

**Saskatchewan Ministry of Energy and Resources  
Minister's Order 169/22**

**Approval of *Directive PNG017: Measurement Requirements for Oil and Gas Operations*  
Section 17, *The Oil and Gas Conservation Act***

In accordance with clause 17(1)(o) of *The Oil and Gas Conservation Act* (the Act) I hereby do order that:

- i) MRO 59/20, dated March 25, 2020 is rescinded;
- ii) the requirements set out substantially in the form of the attached Schedule A apply to the measurement and reporting requirements of wells and facilities licensed in accordance with the Act; and
- iii) the document shown in Schedule A be known as "*Directive PNG017: Measurement Requirements for Oil and Gas Operations*" or "*Directive PNG017*";

Dated at Regina.

Hordenchuk, Sharla ER Digitally signed by  
Hordenchuk, Sharla ER  
Date: 2022.08.15  
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Per: Minister of Energy and Resources  
Sharla Hordenchuk  
Assistant Deputy Minister  
Energy Regulation Division

## SCHEDULE A

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# Measurement Requirements for Oil and Gas Operations

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Directive PNG017

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August 2022

Revision 5.0

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Governing Legislation:

Act: *The Oil and Gas Conservation Act*

Regulation: *The Oil and Gas Conservation Regulations, 2012*

Order: 169/22

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### Record of Change

Version	Date	Description
1.0	May 13, 2015	Approved Initial Draft
1.1	June 22, 2015	Updated Paper Battery requirements in section 12.2.1.2 and corrected several references-draft
1.2	November 17, 2015	Updated version to coincide with AER revisions and adoption of MARP in Saskatchewan-draft
2.0	April 1, 2016	Approved Initial Version (authorized by Minister's Order 47-16)
2.1	August 1, 2017	See What's New Section for changes and all changes are highlighted in red
3.0	December 2019	See What's New Section for changes
4.0	September 2020	See What's New Section for changes
5.0	August 2022	See What's New Section for changes.

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## Introduction

### Purpose

*Directive PNG017: Measurement Requirements for Oil and Gas Operations* (Directive PNG017) consolidates, clarifies, and updates the requirements of the Ministry of Energy and Resources (ER) with respect to measurement points used for accounting and reporting purposes, as well as those measurement points required for upstream petroleum facilities and some downstream pipeline operations under existing regulations. The term measurement as used in this Directive means measurement, accounting, and reporting. While measurement is the determination of a volume, accounting and reporting are integral components of measurement in that after a fluid volume is measured, mathematical procedures (accounting) may have to be employed to arrive at the desired volume to be reported.

### Interpretation

Directive PNG017 includes measurement and reporting requirements that are applicable to oil and gas operations including other industries that are regulated by the Energy Regulation Division of ER. Wells and facilities operated in Saskatchewan must comply with the requirement.

### What's New in This Edition

- Removal of Alberta and British Columbia measurement requirements from various sections in the Directive.
- Restructuring of some sections of the Directive and the relocation of some requirements to different Sections of the Directive.
- Minimizing subjective, vague and inconsistent terminology in favor of objective, clear, and consistent terminology.

[Section 4.3.1.7](#) (amended) Ultrasonic Meters: Reduction of requirements for gas volume metering as Industry has demonstrated that season temperature and pressure factors can be utilized over live temperature and pressure measurement and the uncertainty requirements are still able to be met.

[Section 4.3.6](#) (amended) Volume Calculations: Reduction of requirements for gas volume metering as Industry has demonstrated that season temperature and pressure factors can be utilized over live temperature and pressure measurement and the uncertainty requirements are still able to be met.

[Section 4.3.6.2](#) (amended/new?) Physical Properties of Natural Gas Components: A new requirement for licensees as they must use the most updated physical properties for natural gas when determining gas volumes.

[Section 6.5.1](#) (amended) Crude Oil Single-Well Batteries (facility subtype 311), Crude Oil Multi-well Group Batteries (facility subtype 321), Crude Oil Multi-well Swab Group Battery (314), and Crude Oil Multi-well Swab Paper Battery (facility subtype 316): Clarification on how soon the Gas Oil Ratio (GOR) test must be done for a new non-heavy oil wells.

[Section 6.5.1.1](#) (new) Exception for GOR Testing Frequency: Reduction of requirements for frequency of GOR factor determination for non-heavy oil wells when specific qualifying criteria are met.

[Section 6.5.3](#) (amended) Methods for Determining Factors Used in Estimating Gas Volumes using a GOR Factor for Non-Heavy Oil: Clarifications on what needs to be included in the determination of the GOR factor for heavy and non-heavy oil wells.

[Section 12.2.3](#) (amended) Method for Determining Gas Oil Ratio Factor and GIS: Clarifications on what needs to be included in the determination of the GOR factor for heavy and non-heavy oil wells.

[Section 12.3.5](#) (amended) Water/Steam Primary and Secondary Measurement: Reduction of requirements for water and steam for thermal heavy oil facilities where the primary measurement method must be metered and the secondary measurement can be estimated.

[Section 12.3.8](#) (new requirement) Well Production Measurement: New requirement which will be a more stringent single point measurement uncertainty for group measurement point for a metered volume at a thermal heavy oil battery to match with the single point measurement uncertainty at an individual well.

### **Intent of this Directive**

This Directive specifies:

1. what volumes must be measured and how;
2. what, where, and how volumes may be estimated;
3. if accounting procedures must be performed on the determined volumes and what they are;
4. what data must be kept for audit purposes; and
5. what resultant volumes must be reported to ER.

The licensee must comply with all requirements set out in this Directive.

In this Directive, the term “must” indicates a requirement that must be followed. In some situations, a requirement may be subject to exceptions if specific conditions are met.

The term “should” indicates a recommendation that will not be subject to enforcement. However, ER may direct the licensee in writing to implement changes to improve measurement accuracy, and this direction will become a condition of operation for that facility or facilities.

The Directive does not include instructions on how the volumes must be reported to ER which are included in other ER documents, such as Saskatchewan’s *Directive PNG032: Volumetric, Valuation and Infrastructure Reporting in Petrinex*, but it does include some information on requirements regarding facility subtype, status and code in accordance with those documents.

If requirements in previously-issued ER documents (interim directives, informational letters, guides, etc.) conflict with the requirements in this Directive, the requirements in this Directive replace the prior requirements. Over time, it is intended that all relevant superseded requirements will be rescinded.

## Definitions

Many terms used in this Directive are defined in the Glossary (Appendix 2). However, many critically important definitions are also included within the applicable sections.

## Enforcement

This Directive replaces and supersedes a number of ER documents as identified in Appendix 1. This Directive currently has ER authority in Saskatchewan effective April 1, 2016. ER enforcement of the new requirements, including audits and inspections, that are intended to ensure compliance with the new oil and gas measurement requirements will be applied in accordance with the implementation schedule outlined below. Enforcement actions will be applied according to Section 13.6 of *Directive PNG076: Enhanced Production Audit Program*. As a result of the implementation of Directive PNG017, ER will be rescinding all Measurement Exemptions approved in Saskatchewan before April 1, 2016.

Industry must make continuous progress with respect to compliance for measurement and reporting by April 1, 2021, and ER may require licensees to demonstrate their progress throughout the implementation schedule. Licensees must meet the following implementation schedule:

1. Requirements must be 25 per cent implemented by April 1, 2017.
2. Requirements must be 50 per cent implemented by April 1, 2018.
3. Requirements must be 75 per cent implemented by April 1, 2019.
4. Requirements must be 100 per cent implemented by April 1, 2021.

All licensees are expected to be fully compliant with the ER requirements prior to the implementation of Directive PNG017.

Any facilities licensed after April 1, 2016 must be designed and operated in full compliance with Directive PNG017. For facilities licensed prior to April 1, 2016, licensees are expected to comply with the 5-year implementation schedule.

The licensee may apply for a site-specific measurement exemption through the IRIS generic application process if all the necessary documentation associated with an application is submitted and there is significant evidence to support the exemption (refer to Section 5 of Directive PNG017).

# 1 Standards of Accuracy

## 1.1 Introduction

With regard to accuracy, it is assumed an exact or true value exists for any variable that is valid for the conditions existing at the moment the result is determined. Determining the true value without doubt cannot be done, due to the limitations of measuring equipment and procedures and the possibility of human error. Typically, the closer one wants to approach the true value, the more expense and effort has to be expended.

Measurement in an oil and gas industry context, the principal measurement technologies and procedures are:

- a. Meters for determining flow volumes.
- b. Calculated volumes using a proration formula based on test volumes.
- c. Estimates of volumes based on production facility and product characteristics.
- d. Scales for samples and vehicles.
- e. Gauge boards for tanks.
- f. Gauges for temperature and pressure.

ER has established standards of accuracy for gas and liquid measurement that take into account potential impacts to royalty, equity, reservoir engineering, declining production rates, aging equipment, environment, public safety, accuracy and completeness. The current standards are stated as maximum uncertainty of monthly volume and/or single point measurement uncertainty. The uncertainties are to be applied as “plus/minus” e.g.,  $\pm 5.0$  per cent. Measurement at delivery/sales points must meet the highest accuracy standards because volumes determined at these points can have a direct impact on royalty determination. Other measurement points that play a role in the overall accounting process are subject to less stringent accuracy standards to accommodate physical limitations and/or economics.

The specific standards of accuracy are listed in Section 1.7. Licensees must ensure that their measurement equipment and measurement procedures meet the requirements that are listed in this directive or those that are approved by ER. It is a joint responsibility of the operator of record and licensee to ensure proper reporting as per this Directive and *Directive PNG032: Volumetrics, Valuation, Infrastructure in Petrinex*. The Operator of Record is required to provide the licensee with any documentation or information that may be requested by the licensee in order to demonstrate to ER that the requirements in this directive are being met. If an inspection of a measurement device or of procedure reveals unsatisfactory conditions that preclude the prescribed standards of accuracy from being met ER will direct the licensee to implement changes to improve measurement accuracy, and this direction will become a condition of operation for the facility or facilities.

## 1.2 Applicability and Use of Uncertainties

ER used the uncertainty levels contained in this section to develop many of the requirements for equipment and/or procedures relating to measurement, accounting, and reporting for various aspects of oil and gas production and processing operations, which are explained in detail in other sections. If those requirements are being met and consideration has been made regarding the potential impacts to royalty, equity, reservoir engineering, environment, public safety,

accuracy and completeness, ER considers a licensee to be in compliance without the need to demonstrate compliance with the applicable uncertainty requirements contained in this section.

In some scenarios a licensee may deviate from the minimum requirements for equipment and/or procedures that are stated in this Directive. Refer to Section 5: Site-specific Deviation from Base Requirements.

### **1.3 Maximum Uncertainty of Monthly Volume**

ER requires production data to be reported on a calendar month basis. Maximum uncertainty of monthly volume relates to the limits applicable to equipment and/or procedures used to determine the total monthly volume. Total monthly volumes may result from a single month-long measurement, but more often result from a combination of individual measurements and/or estimations. For example, consider a well in an oil proration battery to which a maximum uncertainty of monthly volume would apply:

1. First, the well is tested, and the oil test rate is used to estimate the well's production for the period until the next test is conducted.
2. The well's total estimated oil production for the month is combined with the month's estimated oil production for the other wells in the battery to arrive at the total estimated monthly oil production for the battery.
3. The total actual monthly oil production for the battery is determined based on measured deliveries out of the battery and inventory change.
4. A proration factor is determined by dividing the actual battery production by the estimated battery production.
5. The proration factor is multiplied by the well's estimated production to determine the well's actual monthly production.

### **1.4 Single Point Measurement Uncertainty**

Single point measurement uncertainty relates to the limits applicable to equipment and/or procedures used to determine a single-phase specific volume at a single measurement point. The oil volume determined during a 24-hour well test conducted on a well in a proration battery is an example of a specific volume determination to which a single point measurement uncertainty limit would apply.

### **1.5 Confidence Level**

The stated uncertainties are not absolute limits. The confidence level, which indicates the probability that true values will be within the stated range, is 95.0 per cent. This implies that there is a 95.0 per cent probability, 19 chances in 20 that the true value will be within the stated range.

### **1.6 Determination of Uncertainties**

The uncertainties referred to relate to the accuracies associated with measurement devices, device calibration, sample gathering and analysis, variable operating conditions, etc. These uncertainties are for single-phase specific volume determination points of specific fluids (oil, gas, or water) or for combinations of two or more such points. These uncertainties do not relate to comparisons of two or more measurement points, such as comparison of inlet volumes to outlet

volumes. Such comparisons are typically expressed as proration factors, allocation factors, or metering differences.

The uncertainties are relevant to equipment at the time of installation. No uncertainty adjustment is required to account for the effects of multiphase fluids, wear, sludge or scale buildup, etc. as it is accepted that such conditions would constitute a bias error to be monitored and accounted for through the use of proration factors, allocation factors, or metering differences.

The methods to be used for determining and combining uncertainties are found in the latest edition of the American Petroleum Institute (API) *Manual of Petroleum Measurement Standards* (MPMS), [Chapter 13](#): Statistical Aspects of Measuring and Sampling or the latest edition of the International Standard Organization (ISO) *Standard 5168: Measurement of Fluid Flow— Estimation of Uncertainty of a Flow-rate Measurement*.

### 1.6.1 Example Calculation

Determination of single point measurement uncertainty for well oil at a proration battery using root sum square methodology:

Individual uncertainties from historical research:

For oil/emulsion measurement:

Oil meter uncertainty = 0.5% (typical manufacturer's specification)

Meter proving uncertainty = 1.5%

Sediment and water (S&W) determination uncertainty = 0.5%

Combined uncertainty =  $\sqrt{[(0.5)^2 + (1.5)^2 + (0.5)^2]}$   
= 1.66% (rounded to 2.0%)

For gas measurement;

Primary element – gas meter uncertainty = 1.0%

Secondary element – (pulse counter or transducer, etc.) uncertainty = 0.5%

Secondary element calibration uncertainty = 0.5%

Tertiary element – (flow calculation, Electronic Flow Measurement (EFM), etc.)  
uncertainty = 0.2%

Gas sampling and analysis uncertainty = 1.5%

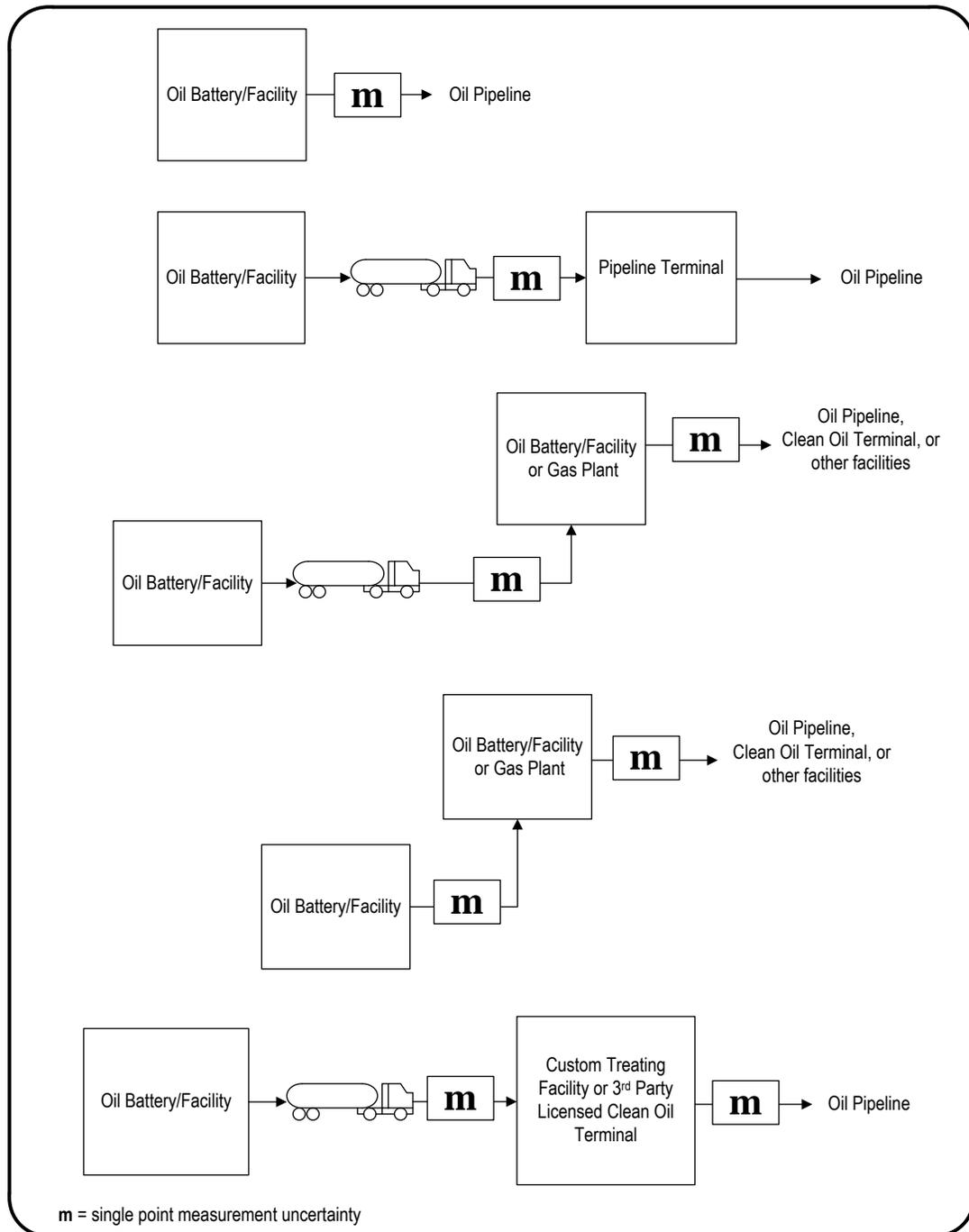
Combined uncertainty =  $\sqrt{[1.0)^2 + (0.5)^2 + (0.5)^2 + (0.2)^2 + (1.5)^2]}$   
= 1.95% (rounded to 2.0%)

### 1.6.2 Oil Systems

#### 1.6.2.1 Oil Systems - Total Battery/Facility Oil

Delivery point measurement, including single-well batteries.

Figure 1.1 Total battery/facility oil - delivery point measurement



Maximum uncertainty of monthly volume = N/A

The uncertainty of the monthly volume will vary, depending upon the number of individual measurements that are combined to yield the total monthly volume.

Single point measurement uncertainty:

Delivery point measures > 100.0 m<sup>3</sup>/day = ±0.5%

Delivery point measures ≤ 100.0 m<sup>3</sup>/day = ±1.0%

The royalty trigger point for oil is at the wellhead. Thus, delivery point measurements are required at the following locations:

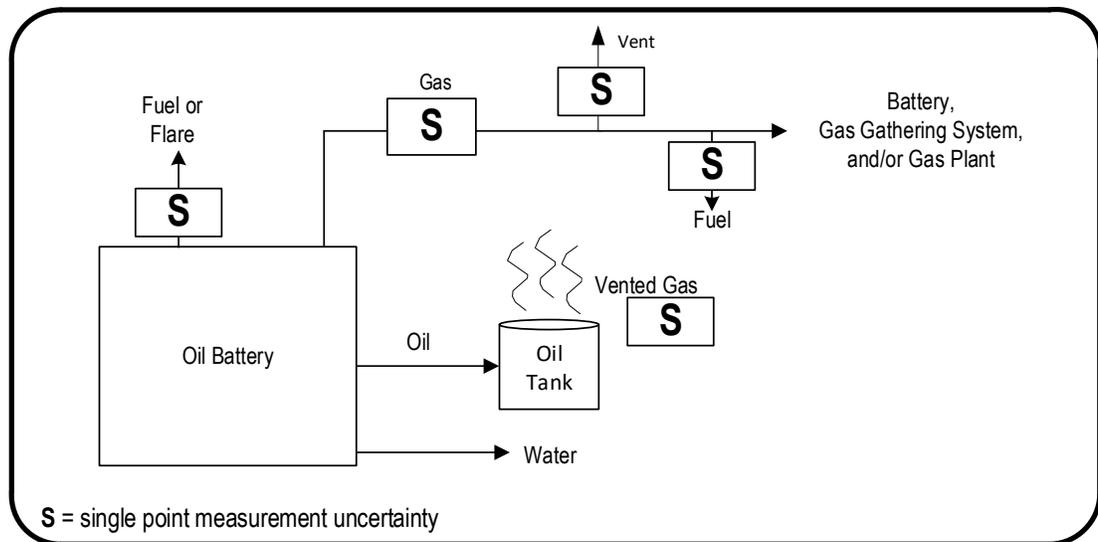
1. Facility dispositions
2. Trucked-in receipts
3. Pipeline receipts
4. Railcar receipts
5. Sales
6. LACT

Excluded: Test points and group points if they are not used for accounting or inventory.

### 1.6.2.2 Oil Systems - Total Battery Gas

Includes produced gas that is vented, flared, or used as fuel, including single-well batteries. Also referred to as associated gas, as it is the gas produced in association with oil production at oil wells.

**Figure 1.2 Total battery gas**



Single point measurement uncertainty:

>  $16.9 \times 10^3 \text{m}^3/\text{day} = \pm 3.0\%$

>  $0.5 \times 10^3 \text{m}^3/\text{day}$  and  $\leq 16.9 \times 10^3 \text{m}^3/\text{day} = \pm 3.0\%$

$\leq 0.5 \times 10^3 \text{m}^3/\text{day} = \pm 10.0\%$

Maximum uncertainty of monthly volume (**M**)

>  $16.9 \times 10^3 \text{m}^3/\text{day} = \pm 5.0\%$

>  $0.5 \times 10^3 \text{m}^3/\text{day}$  and  $\leq 16.9 \times 10^3 \text{m}^3/\text{day} = \pm 10.0\%$

$\leq 0.5 \times 10^3 \text{m}^3/\text{day} = \pm 20.0\%$

Note that **M** is dependent upon combined deliveries, fuel, and vented gas measurement.

The maximum uncertainty of total monthly battery gas volumes allows for reduced emphasis on accuracy as gas production rate declines. For gas rates that are  $\leq 0.5 \text{ } 10^3 \text{ m}^3/\text{day}$ , the gas volumes may be determined by using estimates. Therefore, the maximum uncertainty of monthly volume is set at  $\pm 20.0$  per cent. If gas rates are  $> 0.5 \text{ } 10^3 \text{ m}^3/\text{day}$ , the gas must be metered. However, a component of the total monthly gas volume may include estimates for low volumes of fuel, vented, or flare gas that may add to the monthly uncertainty. At the highest gas production rates, it is expected the use of estimates will be minimal or at least have a minor impact on the accuracy of the total monthly gas volume, thereby resulting in the  $\pm 5.0$  per cent maximum uncertainty of monthly volume.

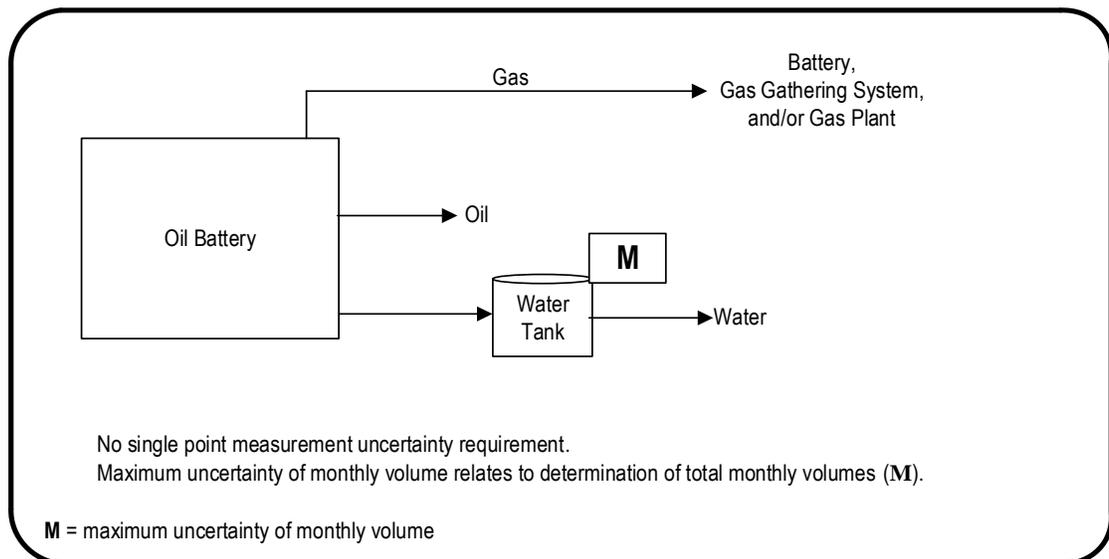
The equipment and/or procedures used to determine the metered gas volumes when metering is required must be capable of meeting a  $\pm 3.0$  per cent single point measurement uncertainty. Due to the difficulty associated with metering very low gas rates, procedures used in determining gas-oil ratios or other factors to be used in estimating gas volumes where rates are  $\leq 0.5 \text{ } 10^3 \text{ m}^3/\text{day}$  must meet a  $\pm 10.0$  per cent single point measurement uncertainty.

These uncertainties do not apply to gas produced in association with heavy oil with a density  $\geq 920.0 \text{ kg/m}^3$  at  $15.0^\circ\text{C}$ .

### 1.6.2.3 Oil Systems - Total Battery Water

Includes single-well batteries.

**Figure 1.3 Total battery water**



Maximum uncertainty of monthly volume:

$> 50.0 \text{ m}^3/\text{month} = \pm 5.0\%$

$\leq 50.0 \text{ m}^3/\text{month} = \pm 20.0\%$

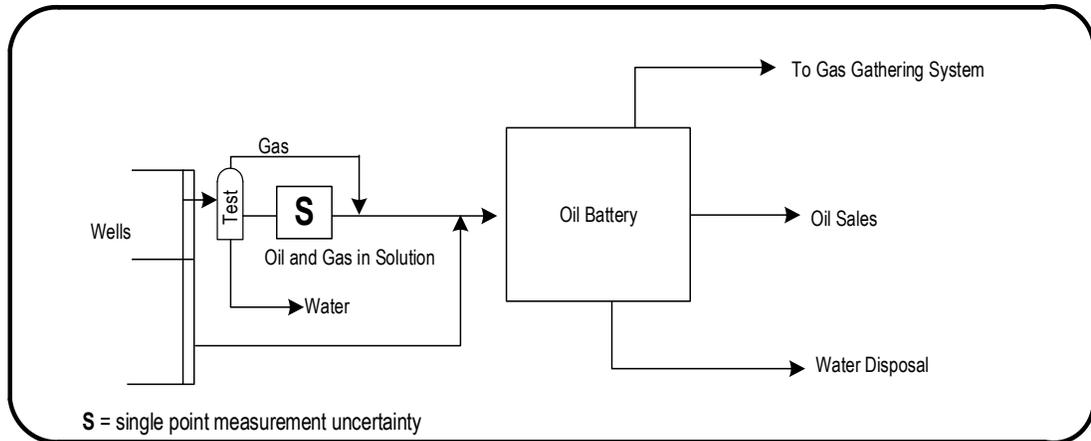
Single point measurement uncertainty = N/A

Total battery water may be determined by metering or estimation, depending on production rates, so no basic requirement has been set for single point measurement uncertainty.

Total battery water production volumes  $\leq 50.0 \text{ m}^3/\text{month}$  may be determined by estimation. Therefore, the maximum uncertainty of monthly volume is set at  $\pm 20.0$  per cent. If the total battery water production volumes are  $> 50.0 \text{ m}^3/\text{month}$ , the water must be separated from the oil and measured. Therefore, the maximum uncertainty of monthly volume is set at  $\pm 5.0$  per cent.

#### 1.6.2.4 Oil Systems - Well Oil - Proration Battery

Figure 1.4 Well oil – proration battery



Single point measurement uncertainty:

All classes =  $\pm 2.0\%$

Maximum uncertainty of monthly volume:

Class 1 (high)  $> 30.0 \text{ m}^3/\text{day} = \pm 5.0\%$

Class 2 (medium)  $> 6.0 \text{ m}^3/\text{day}$  and  $\leq 30.0 \text{ m}^3/\text{day} = \pm 10.0\%$

Class 3 (low)  $> 2.0 \text{ m}^3/\text{day}$  and  $\leq 6.0 \text{ m}^3/\text{day} = \pm 20.0\%$

Class 4 (stripper)  $\leq 2.0 \text{ m}^3/\text{day} = \pm 40.0\%$

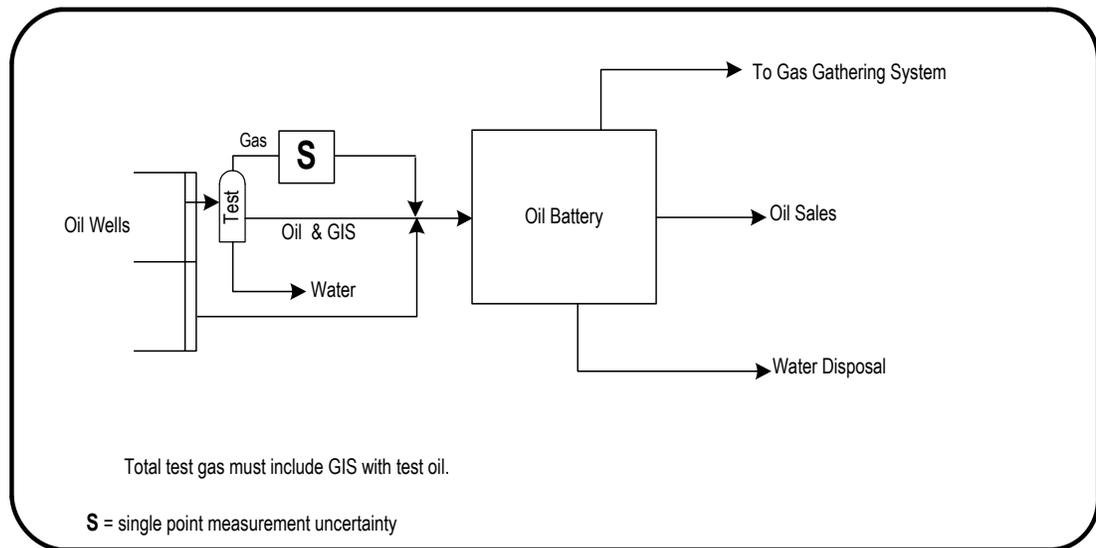
Maximum uncertainty is dependent upon oil and gas test volumes and the number of days the test is used for estimating production, plus correction by a proration factor.

The maximum uncertainty of monthly well oil production volumes for light and medium density oil wells in proration batteries has been developed to allow for reduced emphasis on accuracy as oil production rates decline. Rather than being determined by continuous measurement, monthly well oil production volumes are estimated from well tests and corrected by the use of proration factors to result in actual volumes. Lower rate wells are allowed reduced testing frequencies, which, coupled with the fact that wells may exhibit erratic production rates between tests, results in less certainty that the reported monthly oil production volume will be accurate.

#### 1.6.2.5 Oil Systems - Well Gas - Proration Battery

Also referred to as associated gas, as it is the gas produced in association with oil production at oil wells.

Figure 1.5 Well gas – proration battery



Single point measurement uncertainty:

> 16.9 10<sup>3</sup>m<sup>3</sup>/day = ±3.0%

> 0.5 10<sup>3</sup>m<sup>3</sup>/day and ≤ 16.9 10<sup>3</sup>m<sup>3</sup>/day = ±3.0%

≤ 0.5 10<sup>3</sup>m<sup>3</sup>/day = ±10.0%

Maximum uncertainty of monthly volume:

> 16.9 10<sup>3</sup>m<sup>3</sup>/day = ±5.0%

> 0.5 10<sup>3</sup>m<sup>3</sup>/day and ≤ 16.9 10<sup>3</sup>m<sup>3</sup>/day = ±10.0%

≤ 0.5 10<sup>3</sup>m<sup>3</sup>/day = ±20.0%

Maximum uncertainty is dependent upon oil and gas test volumes and the number of days the test is used for estimating production, plus correction by a proration factor.

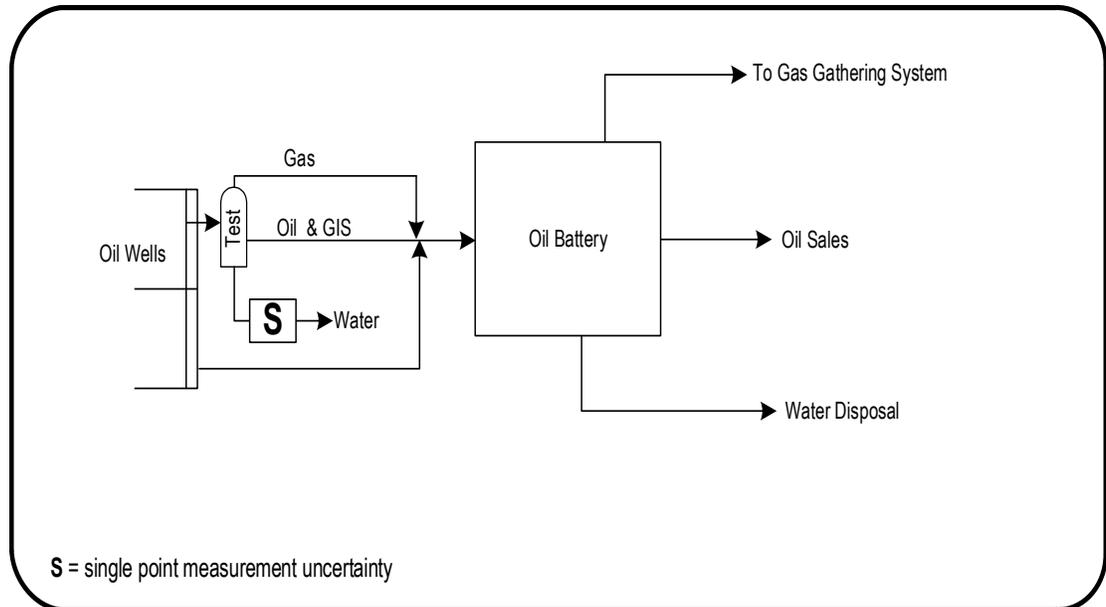
The maximum uncertainty of monthly oil well gas volumes has been developed to allow for reduced emphasis on accuracy as gas production rates decline. Rather than being determined by continuous metering, monthly oil well gas production volumes are estimated from well tests and corrected by the use of proration factors to result in actual volumes. Low gas production rates are typically associated with wells that are allowed reduced testing frequencies, which, coupled with the fact that wells may exhibit erratic production rates between tests, results in less certainty that the reported monthly gas production volume will be accurate.

For gas rates that are ≤ 0.5 10<sup>3</sup>m<sup>3</sup>/day, the well test gas volume may be determined by using estimates. Therefore, the maximum uncertainty of monthly volume is set at ±20.0 per cent. If gas rates are > 0.5 10<sup>3</sup>m<sup>3</sup>/day, the test gas must be metered. However, a component of a well's total test gas volume may include estimates for gas in solution dissolved in the test oil volume, which may add to the monthly uncertainty. At the highest gas production rates, it is expected that the use of estimates will be minimal or at least have a minor impact on the accuracy of the total monthly gas volume, thereby resulting in the ±5.0 per cent maximum uncertainty of monthly volume.

The equipment and/or procedures used to determine the measured test gas volumes if measurement is required must be capable of meeting a  $\pm 3.0$  per cent single point measurement uncertainty. Due to the difficulty associated with measuring very low gas rates, procedures used in determining gas-oil ratios or other factors to be used in estimating gas volumes, if rates are  $\leq 0.5 \text{ } 10^3 \text{ m}^3/\text{day}$ , must meet a  $\pm 10.0$  per cent single point measurement uncertainty.

### 1.6.2.6 Oil Systems - Well Water - Proration Battery

Figure 1.6 Well water – proration battery



Single point measurement uncertainty =  $\pm 10.0\%$

Maximum uncertainty of monthly volume = N/A

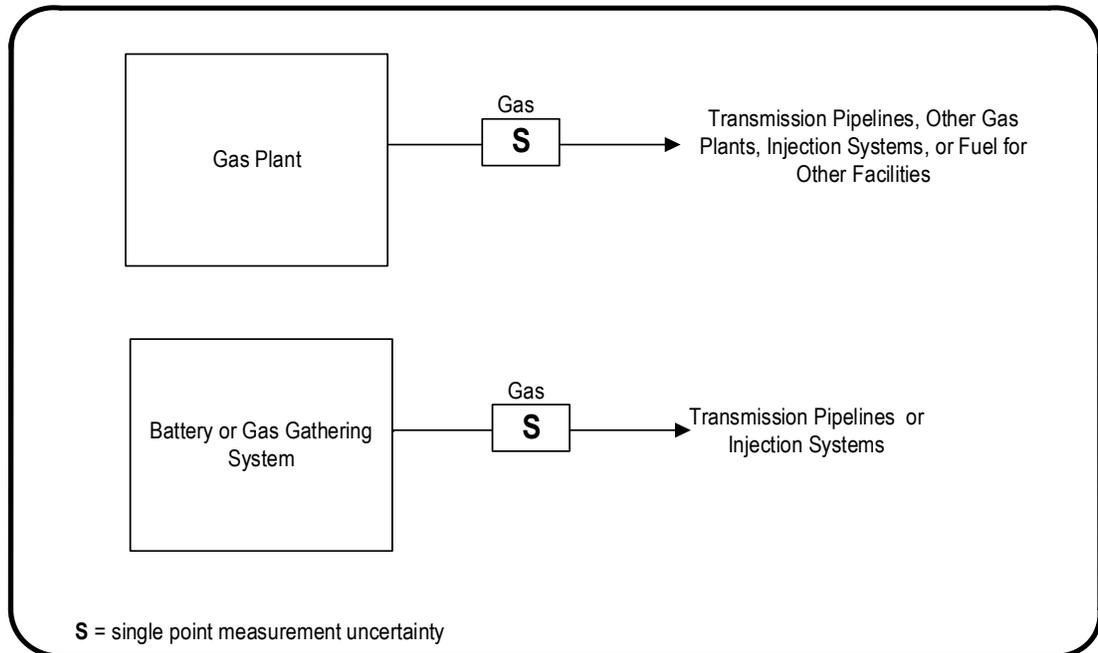
The uncertainty of the monthly volume will vary, depending upon the method used to determine test water rates and the frequency of well tests.

Rather than being determined by continuous measurement, monthly oil well water production volumes are estimated from well tests and corrected by the use of proration factors to result in actual volumes. The water rates determined during the well tests may be inferred from determining the water content of emulsion samples, and in some scenarios, estimates may be used to determine water rates. Therefore, the single point measurement uncertainty is set at  $\pm 10.0$  per cent.

### 1.6.3 Gas Systems

#### 1.6.3.1 Gas Systems - Gas Deliveries – Sales Gas

Figure 1.7 Gas deliveries – sales gas



Single point measurement uncertainty =  $\pm 2.0\%$

Maximum uncertainty of monthly volume = N/A

The total monthly volume may result from a single month-long measurement, making the uncertainty of the monthly volume equivalent to the single point measurement uncertainty.

Since the delivery point is often a custody transfer point, a stringent requirement is set for the single point measurement uncertainty.

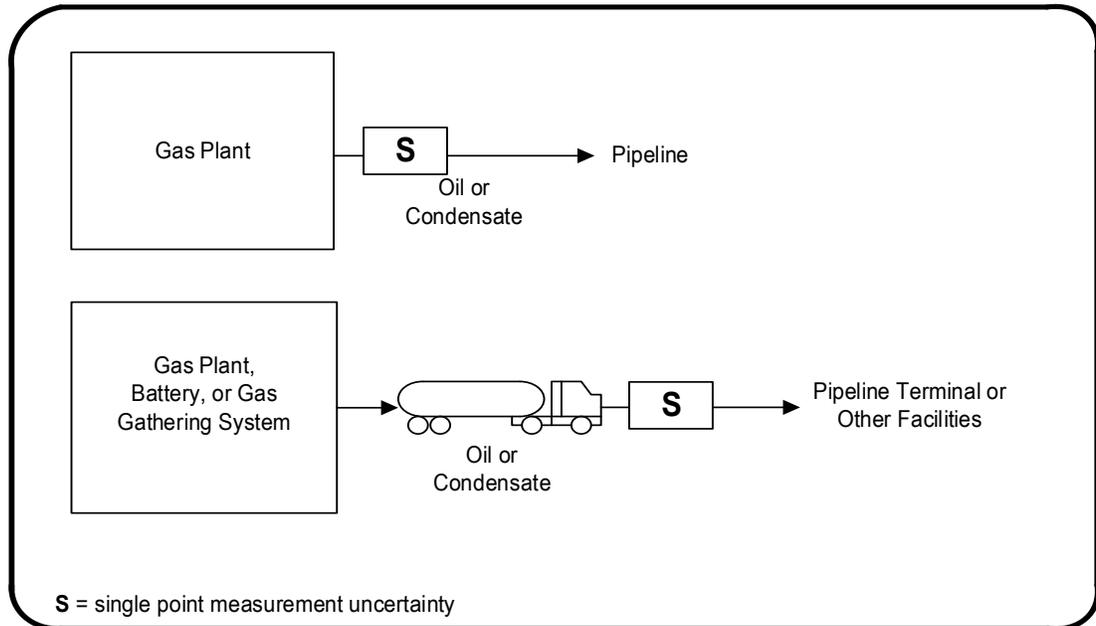
In some scenarios, this type of gas may be delivered to other plants for further processing or to injection facilities. Thus, delivery point measurements are required at the following locations:

1. Gas plant dispositions
2. Sales to downstream
3. Purchase from downstream facilities
4. Cross-border and cross-jurisdiction
5. Gas delivered from one upstream facility to another that is not tied to the same system for FUEL, such as from a gas battery to an oil battery
6. Condensate disposition to an oil facility or for sales

Excluded: Return fuel to the original source facility after the gas has been sweetened.

### 1.6.3.2 Gas Systems - Hydrocarbon Liquid Deliveries

Figure 1.8 Hydrocarbon liquid deliveries



Single point measurement uncertainty:

Delivery point measures  $> 100.0 \text{ m}^3/\text{day} = \pm 0.5\%$

Delivery point measures  $\leq 100.0 \text{ m}^3/\text{day} = \pm 1.0\%$

Maximum uncertainty of monthly volume = N/A

The uncertainty of the monthly volume will vary, depending upon the number of individual measurements that are combined to yield the total monthly volume.

The term delivery point measurement for hydrocarbon liquids refers to the point at which the hydrocarbon liquid production from a battery or facility is measured. Where clean hydrocarbon liquids are delivered directly into a pipeline system via a Lease Automatic Custody Transfer Unit (LACT) measurement or trucked to a pipeline terminal, it can also be referred to as the custody transfer point. The delivery point terminology is from the perspective of the producing battery or facility, but the receiving facility (pipeline, terminal, custom treating facility, other battery, etc.) may refer to this point as its receipt point. The hydrocarbon liquid volume determined at the delivery point is used in all subsequent transactions involving that liquid.

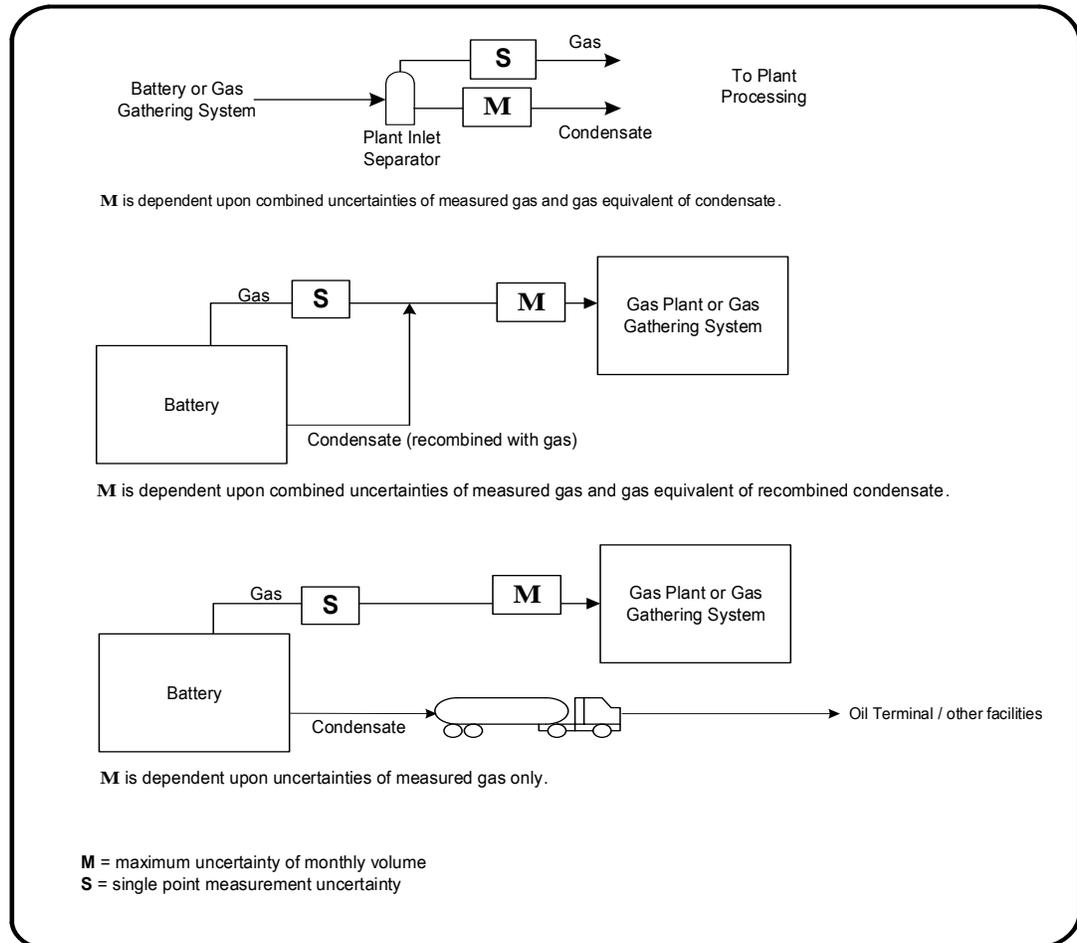
Hydrocarbon liquids delivered out of a gas system at the well, battery, or plant inlet levels are typically condensate, and in some scenarios, they may be considered to be oil. The hydrocarbon liquids delivered out of a gas plant may be pentanes, butane, propane, ethane, or a mixture of various components.

The measurement equipment and/or procedures must be capable of determining the hydrocarbon liquid volume within the stated limits.

For facilities where the hydrocarbon liquid delivery volumes total  $\leq 100.0 \text{ m}^3/\text{day}$ , the single point measurement uncertainty has been increased to allow for the economical handling of hydrocarbon liquids when minimal volumes would not justify the added expense for improved measurement equipment and/or procedures.

### 1.6.3.3 Gas Systems - Plant Inlet or Total Battery/Group Gas

Figure 1.9 Plant inlet or total battery/group gas



Maximum uncertainty of monthly volume =  $\pm 5.0\%$

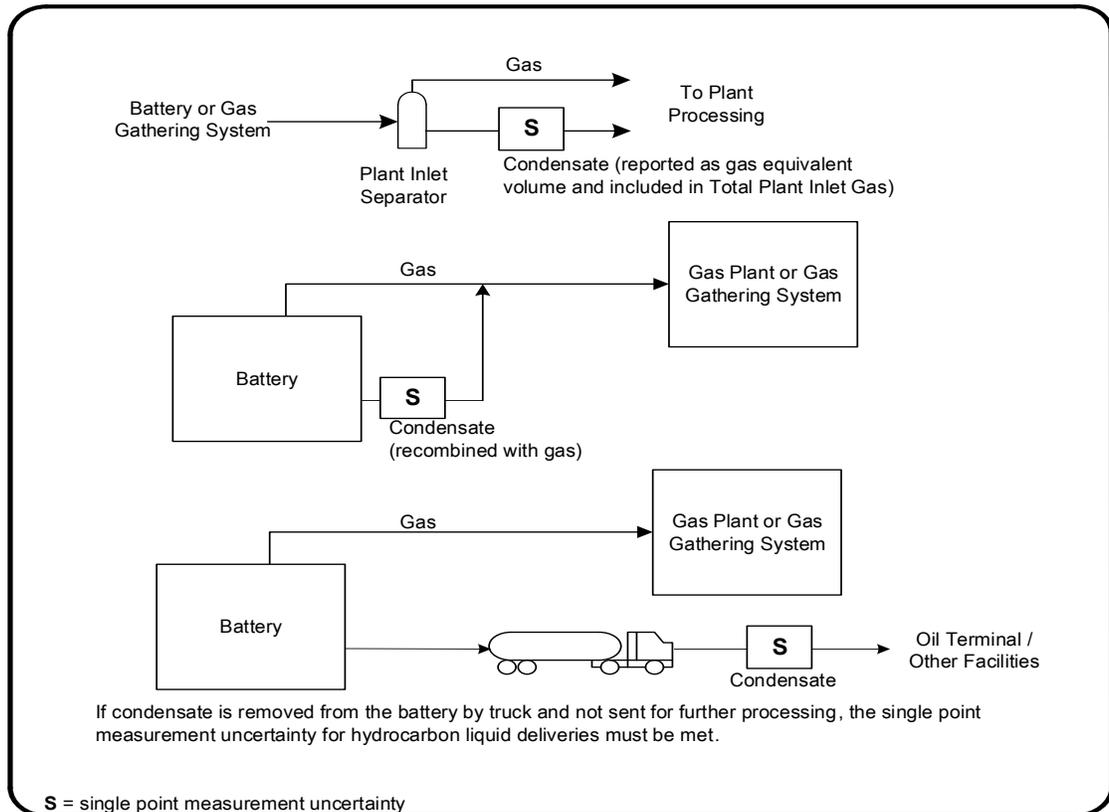
Single point measurement uncertainty =  $\pm 3.0\%$

Plant inlet gas or total battery/group gas is typically unprocessed gas that may vary in composition and may contain entrained liquids. The total reported gas volume could result from combining several measured volumes from various points and may also include the calculated gas equivalent volume of entrained hydrocarbon liquids, typically condensate. The maximum uncertainty of monthly volume is set at  $\pm 5.0$  per cent to allow for the uncertainties associated with measuring gas under these conditions.

The equipment and/or procedures used to determine the measured gas volumes must be capable of meeting a  $\pm 3.0$  per cent single point measurement uncertainty.

## 1.6.3.4 Gas Systems - Plant Inlet or Total Battery Group Condensate - Recombined

Figure 1.10 Plant inlet or total battery/group condensate - recombined



Single point measurement uncertainty =  $\pm 2.0\%$

Maximum uncertainty of monthly volume = N/A

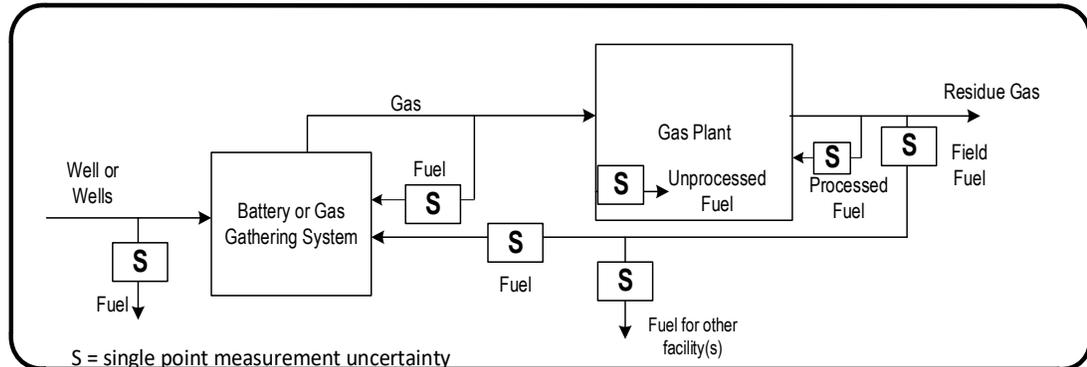
The condensate volume is included in the total gas volume for reporting purposes and is therefore covered by the maximum uncertainty of monthly volume for the plant inlet or total battery/group gas.

Plant inlet condensate is typically separated from the inlet stream and sent through the plant for further processing. For reporting purposes, the gas equivalent of the plant inlet condensate is included in the total plant inlet gas volume. If total battery/group condensate upstream of the plant inlet is separated and measured prior to being recombined with the gas production, the condensate is converted to a gas equivalent volume and included in the gas production volume. In either scenario, the condensate single point measurement uncertainty is set at  $\pm 2.0$  per cent for the liquid volume determination.

Note that if plant inlet or total battery/group condensate is separated and delivered out of the system at that point, the condensate measurement is subject to the single point measurement uncertainties stipulated for hydrocarbon liquid deliveries stated in [Section 1.6.3.2](#).

### 1.6.3.5 Gas Systems - Fuel Gas

Figure 1.11 Fuel gas



Single point measurement uncertainty:

$> 0.5 \text{ } 10^3\text{m}^3/\text{day} = \pm 3.0\%$

$\leq 0.5 \text{ } 10^3\text{m}^3/\text{day} = \pm 10.0\%$

Maximum uncertainty of monthly volume:

$> 0.5 \text{ } 10^3\text{m}^3/\text{day} = \pm 5.0\%$

$\leq 0.5 \text{ } 10^3\text{m}^3/\text{day} = \pm 20.0\%$

Note that maximum uncertainty is dependent upon combined uncertainties of various fuel sources at each reporting facility.

The maximum uncertainty of monthly fuel gas volumes allows for reduced emphasis on accuracy as gas flow rates decline.

For all upstream oil and gas facilities, if the annual average fuel gas rate is  $\leq 0.5 \text{ } 10^3\text{m}^3/\text{day}$  on a per-site basis, the gas volume may be determined by using estimates. Therefore, the maximum uncertainty of the monthly volume is set at  $\pm 20.0$  per cent. If the annual average fuel gas rates are  $> 0.5 \text{ } 10^3\text{m}^3/\text{day}$  on any site, the gas must be metered, but since the gas being used as fuel may be unprocessed gas and part of the total fuel gas volume may include some estimated volumes up to  $0.5 \text{ } 10^3\text{m}^3/\text{day}$ , the maximum uncertainty of the monthly volume is set at  $\pm 5.0$  per cent to allow for the uncertainties associated with measuring gas under those conditions. See Section 4.2 for more detail.

Gas used for pneumatic devices that is vented or flared must be reported as vented or flared, respectively.

For facilities licensed prior to January 1, 2020 or for non-licensed facilities built before January 1, 2020, the volume of gas emitted by pneumatic devices may be estimated and then subtracted from the metered fuel gas volume in the case where the metered fuel gas provides the pneumatic gas supply.

For facilities licensed after January 1, 2020 or for non-licensed facilities built after January 1, 2020, the facility must be built so that the metered fuel gas does not include gas emitted by pneumatic devices.

Gas used for pilot, purge, sweep, blanket and makeup gas must be reported as flared.

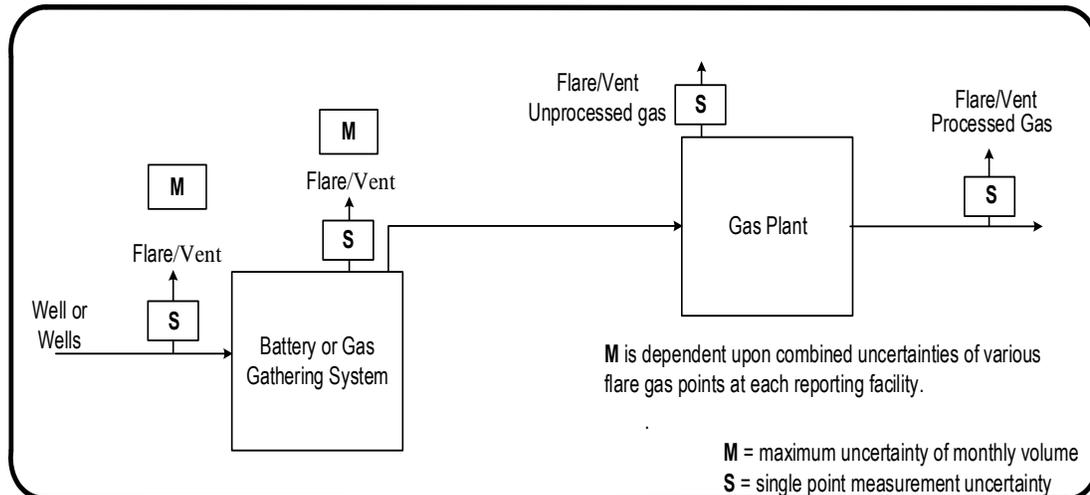
For facilities licensed prior to January 1, 2020 or for non-licensed facilities built before January 1, 2020, the volume of gas used as pilot, purge, sweep, blanket, and makeup gas may be estimated and then subtracted from metered fuel gas in the case where metered fuel gas also provides the pilot, purge, sweep, blanket, and makeup gas supply. The volume that is subtracted from fuel gas does not contribute to the allowance of  $0.5 \times 10^3 \text{m}^3/\text{day}$  that may be estimated for flare gas.

For facilities licensed after January 1, 2020 or for non-licensed facilities built after January 1, 2020, the facility must be built so that the metered fuel gas does not include pilot, purge, sweep, blanket, and makeup gas supply.

The equipment and/or procedures used to determine the measured gas volumes if metering is required must be capable of meeting a  $\pm 3.0$  per cent single point measurement uncertainty. Due to the difficulty associated with measuring very low gas rates, procedures used in determining gas-oil ratios or other factors to be used in estimating gas volumes, if rates are  $\leq 0.5 \times 10^3 \text{m}^3/\text{day}$ , must meet a  $\pm 10.0$  per cent single point measurement uncertainty.

### 1.6.3.6 Gas Systems - Flare and Vent Gas

Figure 1.12 Flare and vent gas



Maximum uncertainty of monthly volume =  $\pm 20.0\%$

Single point measurement uncertainty =  $\pm 5.0\%$

Flare gas may be clean processed gas or it may be unprocessed gas, depending on the point in the system from which gas is being flared. Continuous and intermittent flared and vent volumes at all oil or gas production or processing facilities, including thermal *in situ* facilities but excluding non-thermal heavy oil and bitumen facilities, where annual average total flared and vented volumes per facility are  $> 0.5 \times 10^3 \text{m}^3/\text{day}$  excluding pilot, purge, or dilution gas must be metered.

Gas used for pilot, purge, sweep, blanket and makeup gas must be reported as flared.

For facilities licensed prior to January 1, 2020 or for non-licensed facilities built before January 1, 2020, the volume of gas used as pilot, purge, sweep, blanket, and makeup gas may be estimated and then subtracted from metered fuel gas in the case where metered fuel gas also provides the pilot, purge, sweep and blanket gas supply. The volume that is subtracted from fuel gas does not contribute to the allowance of  $0.5 \times 10^3 \text{m}^3/\text{day}$  that may be estimated for flare gas.

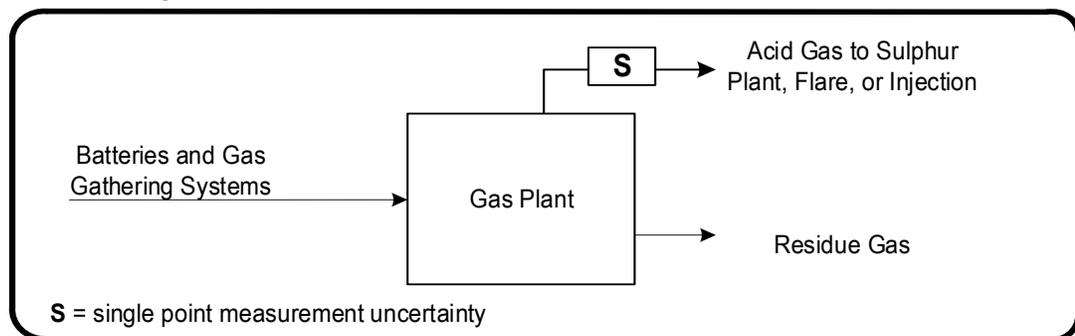
For facilities licensed after January 1, 2020 or for non-licensed facilities built after January 1, 2020, the facility must be built so that the metered fuel gas does not include pilot, purge, sweep, blanket, and makeup gas supply.

Uncombusted gas released to the atmosphere, including fugitive emissions, must be reported as vent gas using the methodologies in *Guideline PNG035: Estimating Venting and Fugitive Emissions*. When a fugitive emission is discovered the licensee must estimate and report the amount of gas released from the time of discovery until the fugitive emission is eliminated. If, at a facility, all gas that is received or produced is vented including casing gas, then no fugitive emissions need to be reported. All documentation relating to the fuel, flare and vent including fugitive emission must be kept for ER to review.

Sites requiring flare or vent gas metering may estimate up to  $0.5 \times 10^3 \text{m}^3/\text{day}$ . Any continuous and intermittent flare and vent volumes at non-thermal heavy crude oil or bitumen facilities that are  $> 2.0 \times 10^3 \text{m}^3/\text{day}$  must be metered. Sites requiring flare or vent gas metering may estimate up to  $2.0 \times 10^3 \text{m}^3/\text{day}$ . Flare lines usually operate in a shut-in condition and may be required to accommodate partial or full volumes of gas production during flaring conditions. In some scenarios if flaring is infrequent and no measurement equipment is in place, flare volumes must be estimated such as flaring at SW Saskatchewan gas wells in a proration battery where there is no on-site measurement equipment. Therefore, the maximum uncertainty of the monthly volume is set at  $\pm 20.0$  per cent, to allow for the erratic conditions associated with flare measurement.

### 1.6.3.7 Gas Systems - Acid Gas

Figure 1.13 Acid gas



Single point measurement uncertainty =  $\pm 10.0\%$  for low pressure acid gas before compression, and =  $\pm 3.0\%$  after compression.

Maximum uncertainty of monthly volume = N/A

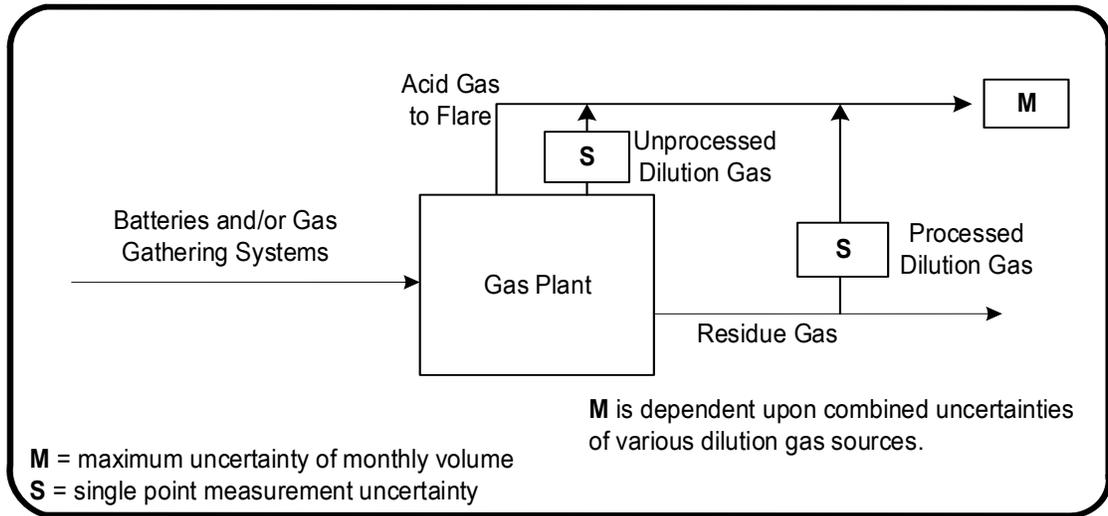
The total monthly volume may result from a single month-long measurement, making the uncertainty of the monthly volume equivalent to the single point measurement uncertainty.

Acid gas usually contains a great deal of water vapour and has other conditions associated with it, such as very low pressure that affects measurement accuracy. Therefore, the single point measurement uncertainty is set at  $\pm 10.0$  per cent.

See Section **Error! Reference source not found.**6.3 for details.

### 1.6.3.8 Gas Systems - Dilution Gas

Figure 1.14 Dilution gas



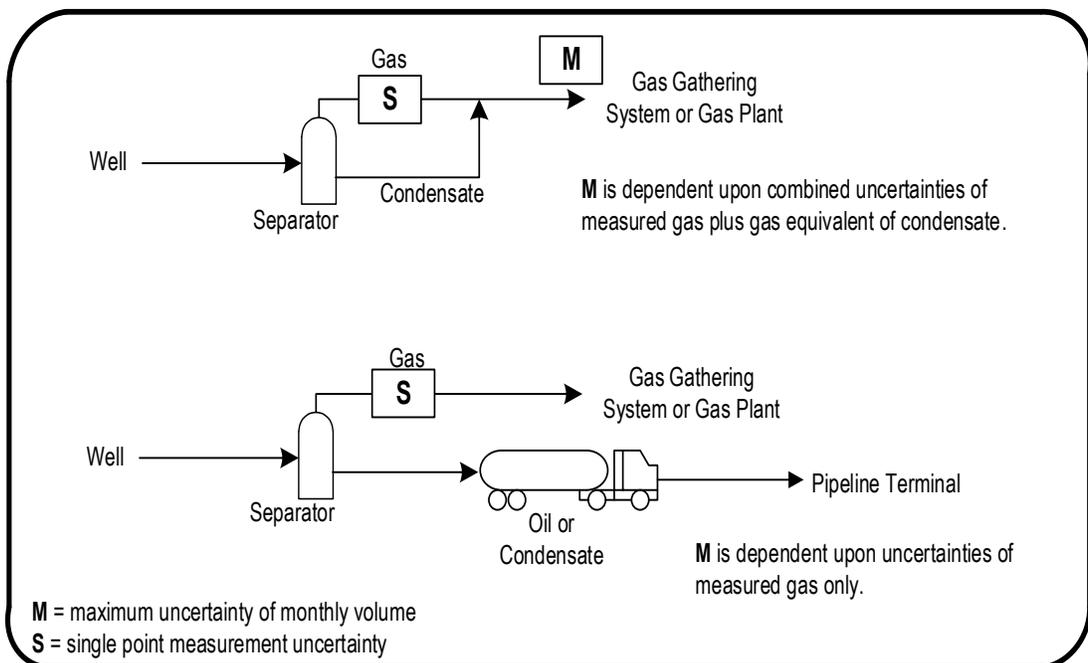
Single point measurement uncertainty =  $\pm 3.0\%$

Maximum uncertainty of monthly volume =  $\pm 5.0\%$

Dilution gas is gas used to provide adequate heating value for incineration or flaring acid gas. Since it must be metered, it is subject to the same uncertainties as stated in Section 1.6.3.5 for fuel gas that must be determined by metering.

### 1.6.3.9 Gas Systems – Well Gas - Well Site Separation

Figure 1.15 Well gas – well site separation



Single point measurement uncertainty =  $\pm 3.0\%$

Maximum uncertainty of monthly volume:

$> 16.9 \text{ } 10^3 \text{ m}^3/\text{day} = \pm 5.0\%$

$\leq 16.9 \text{ } 10^3 \text{ m}^3/\text{day} = \pm 10.0\%$

If production components from gas wells are separated and continuously measured, the maximum uncertainty of monthly well gas volumes allows for reduced emphasis on accuracy as gas production rates decline. Since the separated gas is unprocessed and may still contain entrained liquids at the measurement point and a component of the total reported well gas production may include the calculated gas equivalent volume of the well's condensate production, the maximum uncertainty of monthly volumes also allows for the uncertainties associated with measuring gas under those conditions.

The equipment and/or procedures used to determine the separated measured well gas volumes must be capable of meeting a  $\pm 3.0$  per cent single point measurement uncertainty.

### 1.6.3.10 Gas Systems - Well Gas - Proration Battery

Figure 1.16 Well gas (effluent measurement battery)

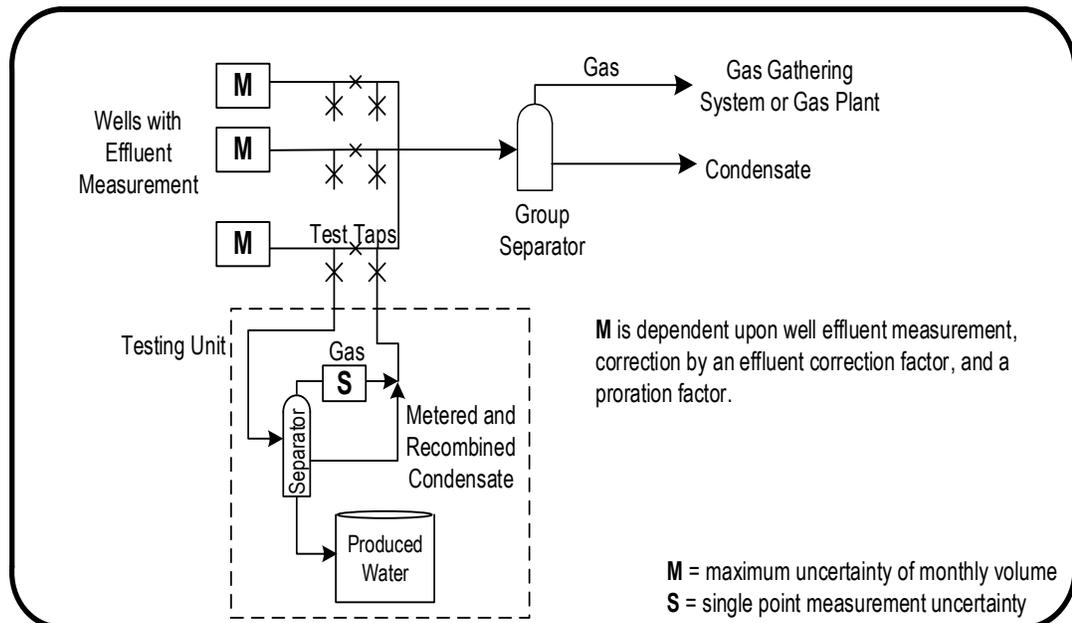
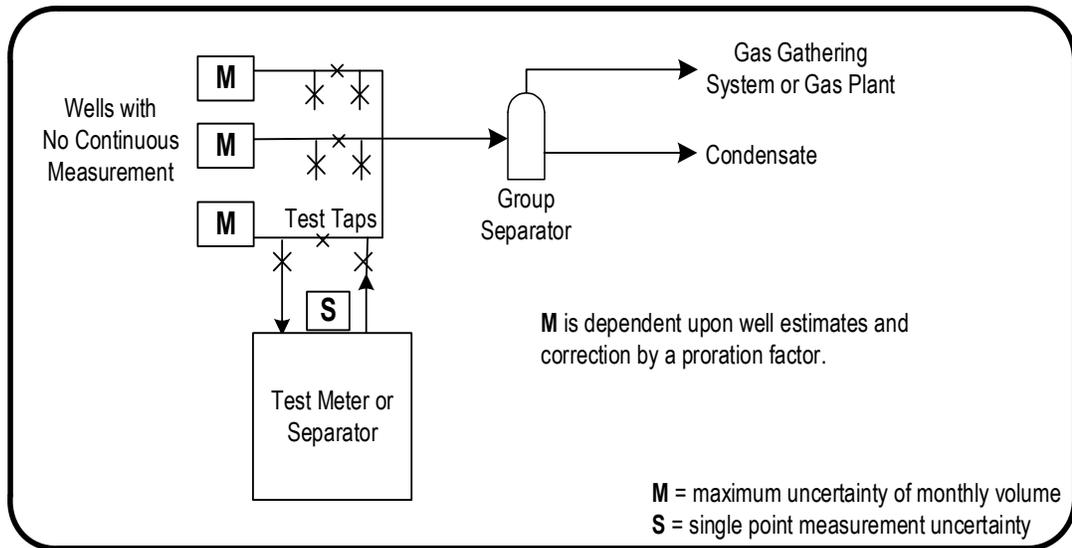


Figure 1.17 Well gas - SW Saskatchewan or other approved multi-well proration battery



Single point measurement uncertainty =  $\pm 3.0\%$

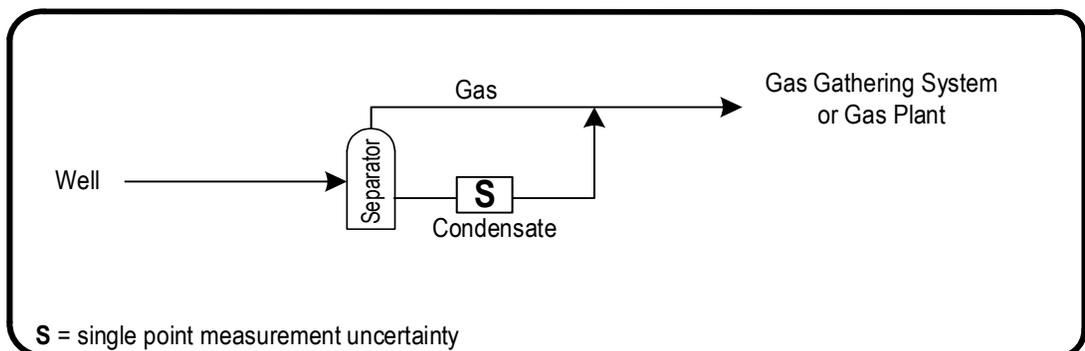
Maximum uncertainty of monthly volume =  $\pm 15.0\%$

If production components from gas wells are not separated and continuously measured, the gas wells are subject to a proration accounting system. There are two types of gas proration batteries. Effluent gas wells have continuous effluent measurement, and the actual production is prorated based on the measurement of group gas and liquid components following separation at a central location. Dry gas wells approved to operate without continuous measurement have the production estimated based on periodic tests, and the actual production is prorated based on the measurement of group volumes at a central location. For both types of proration batteries, the maximum uncertainty of the monthly well gas volume is set at  $\pm 15.0$  per cent to allow for the inaccuracies associated with these types of measurement systems.

The equipment and/or procedures used to determine the measured well test gas volumes downstream of separation during effluent meter correction factor tests or during the periodic dry gas well tests must be capable of meeting a  $\pm 3.0$  per cent single point measurement uncertainty.

### 1.6.3.11 Gas Systems - Well Condensate - Recombined

Figure 1.18 Well condensate - recombined



Single point measurement uncertainty =  $\pm 2.0\%$

Maximum uncertainty of monthly volume = N/A

The gas equivalent of the condensate volume is included in the total well gas volume for reporting purposes and is therefore covered by the monthly uncertainty for the well gas.

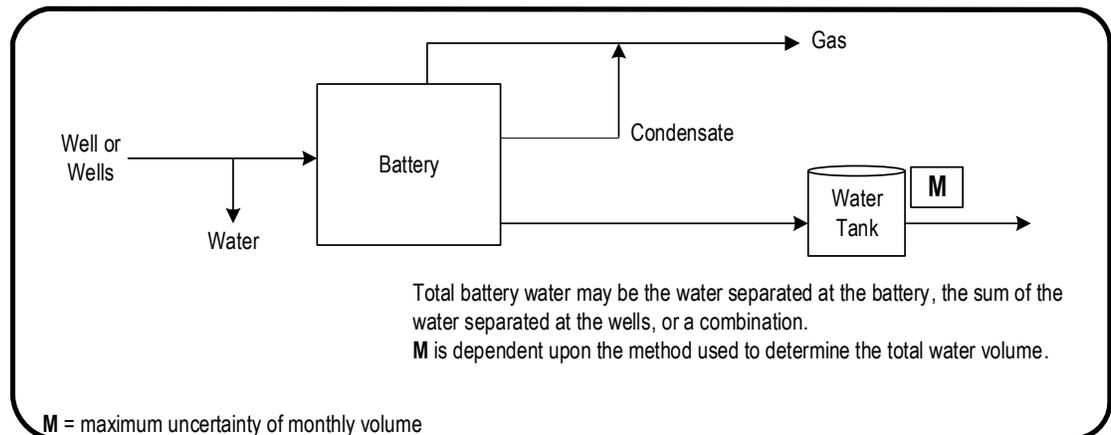
If condensate produced by a gas well is separated and measured at the wellhead prior to being recombined with the gas production, the condensate is mathematically converted to a gas equivalent volume and added to the well gas production volume. In this scenario, the condensate single point measurement uncertainty is set at  $\pm 2.0$  per cent for the liquid volume determination. No requirement has been set for the maximum uncertainty of monthly volume because the gas equivalent of the condensate volume is included in the total well gas volume for reporting purposes.

In the scenario of a gas well subject to effluent measurement, the gas equivalent of the condensate volume is included in the well's total gas production volume. The liquid volume determination, which is done during the effluent meter correction factor test, is subject to a single point measurement uncertainty of  $\pm 2.0$  per cent. No requirement has been set for the maximum uncertainty of monthly volume because the gas equivalent of the condensate volume is included in the total well gas volume for reporting purposes.

Note that if condensate produced by a gas well is separated at the wellhead and delivered out of the system at that point, the condensate is reported as a liquid volume. In this scenario, the condensate measurement is subject to the single point measurement uncertainties stipulated for hydrocarbon liquid deliveries stated in Section 1.6.3.2.

### 1.6.3.12 Gas Systems - Total Battery Water

Figure 1.19 Total battery water



Single point measurement uncertainty = N/A

Maximum uncertainty of monthly volume =  $\pm 5.0\%$

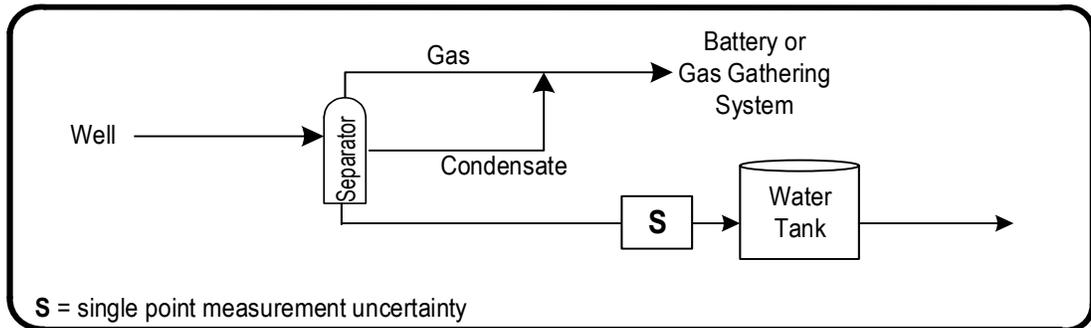
Total battery water may be determined by an individual group measurement, by totaling individual well measurements, or by totaling individual well estimates, so no basic requirement for measurement uncertainty has been set.

Total battery water in a gas system may be collected at a central location where it can be measured prior to disposal, or it may be a summation of individual well estimates or

measurements of water collected at multiple locations and disposed from those sites. The  $\pm 5.0$  per cent maximum uncertainty of monthly volume allows for some leeway in volume determination.

### 1.6.3.13 Gas Systems - Well Water

Figure 1.20 Well water



Single point measurement uncertainty =  $\pm 10.0\%$

Maximum uncertainty of monthly volume = N/A

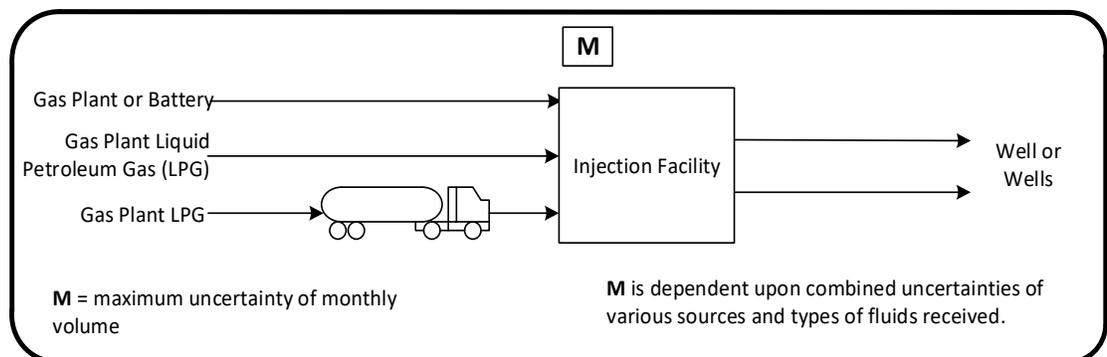
The uncertainty of the monthly volume will vary, depending upon whether produced volumes are subject to individual well measurement, estimation, or proration.

Water production at gas wells may be determined by measurement after separation, or if separators are not used, it may be determined by using water-gas ratios determined from engineering calculations or semiannual tests. To allow for the various methods used to determine production volumes, the single point measurement uncertainty is set at  $\pm 10.0$  per cent.

## 1.6.4 Injection/Disposal Systems

### 1.6.4.1 Injection/Disposal Systems - Total Gas

Figure 1.21 Total gas



Maximum uncertainty of monthly volume =  $\pm 5.0\%$

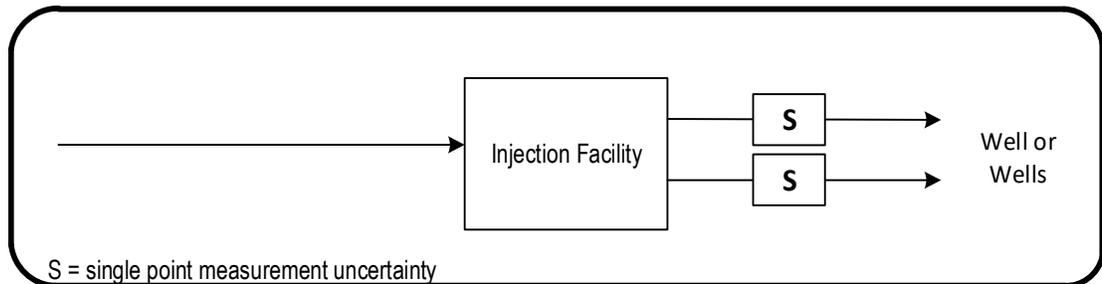
Single point measurement uncertainty = N/A

The single point measurement uncertainty will vary depending on the source and type of fluids received.

Gas used in injection/disposal systems may be clean processed gas or unprocessed gas that may contain entrained liquids, and in some scenarios several sources may make up the total gas volume received by an injection system. This includes gas injected into caverns. The maximum uncertainty of monthly volume is set at  $\pm 5.0$  per cent to allow for the uncertainties associated with measuring gas under those conditions.

#### 1.6.4.2 Injection/Disposal Systems - Well Gas

Figure 1.22 Well gas



Single point measurement uncertainty =  $\pm 3.0\%$

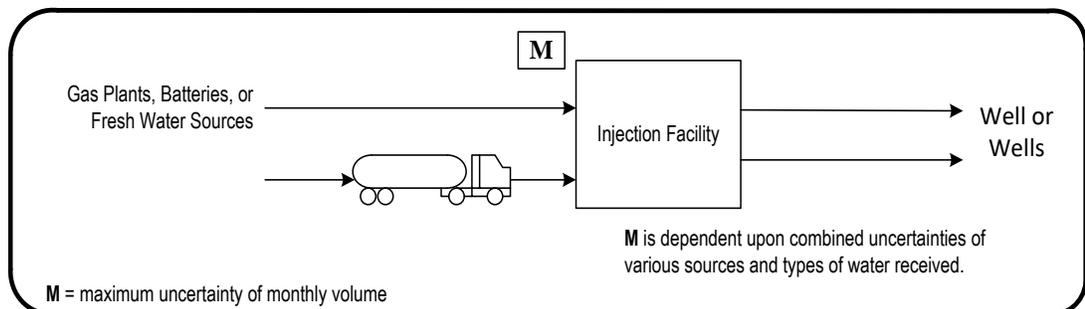
Maximum uncertainty of monthly volume = N/A

The total monthly volume may result from a single month-long measurement, making the uncertainty of the monthly volume equivalent to the single point measurement uncertainty.

The gas injected/disposed into each well must be measured at the injection site and may consist of clean processed gas and/or unprocessed gas that may contain entrained liquids. The equipment and/or procedures used to determine the gas volumes injected/disposed into each well must be capable of meeting a  $\pm 3.0$  per cent single point measurement uncertainty.

#### 1.6.4.3 Injection/Disposal Systems - Total Water

Figure 1.23 Total water



Maximum uncertainty of monthly volume =  $\pm 5.0\%$

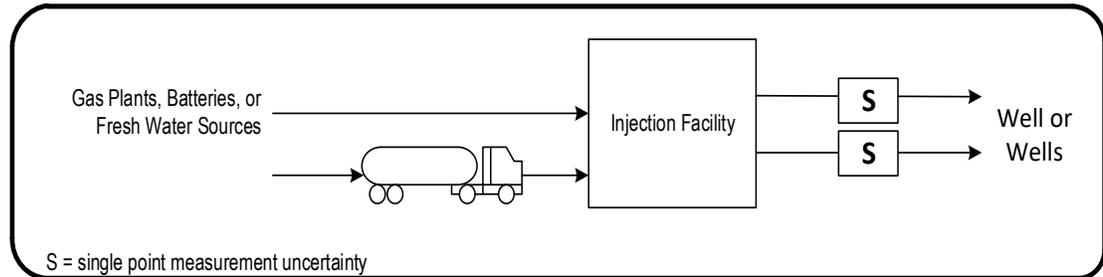
Single point measurement uncertainty = N/A

To be equivalent to the requirements for Oil Systems - Total Battery Water (Section 1.6.2.3) and Gas Systems - Total Battery Water (Section 1.6.3.12).

Water used in injection/disposal systems may be produced water from oil or gas batteries, fresh water from water source wells, or waste water. To be equivalent to the requirements for total oil and gas battery water, the maximum uncertainty of monthly volume is set at  $\pm 5.0$  per cent.

#### 1.6.4.4 Injection/Disposal Systems - Well Water/Steam

Figure 1.24 Well water/steam



Single point measurement uncertainty =  $\pm 5.0\%$

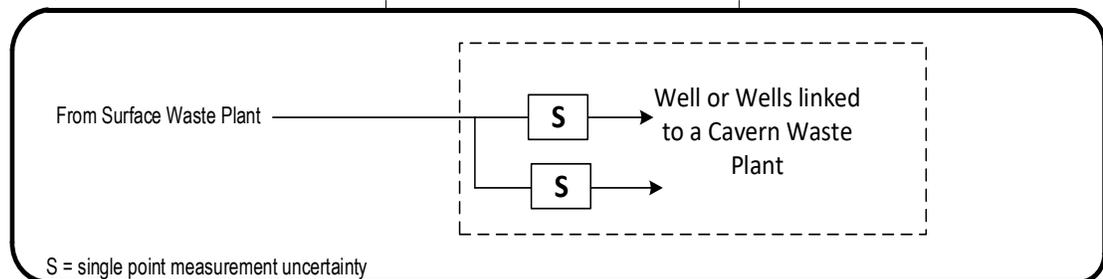
Maximum uncertainty of monthly volume = N/A

The total monthly volume may result from a single month-long measurement, making the uncertainty of the monthly volume equivalent to the single point measurement uncertainty.

The water/steam injected/disposed into each well must be measured at the injection site. The single point measurement uncertainty is set at  $\pm 5.0$  per cent. For water and steam production at a thermal *in situ* facility, the single point measurement uncertainty is set at  $\pm 2.0$  per cent, see Sections 12.3.3 and 12.3.4 for details.

#### 1.6.4.5 Injection/Disposal Systems – Waste injected into Cavern

Figure 1.25 Well waste



Single point measurement uncertainty =  $\pm 5.0\%$

Maximum uncertainty of monthly volume = N/A

The total monthly volume may result from a single month-long measurement, making the uncertainty of the monthly volume equivalent to the single point measurement uncertainty.

The waste into each well must be metered at the injection site. The single point measurement uncertainty is set at  $\pm 5.0$  per cent.

For withdrawals from the cavern the same single point measurement uncertainty is required.

## 1.7 Standards of Accuracy—Summary

### 1.7.1 Oil Systems – excluding heavy oil

		Flow Rate	Maximum uncertainty of monthly volume ± %	Single point measurement uncertainty ± %
(i)	Total battery oil (delivery point measurement)	Delivery point measures > 100.0 m <sup>3</sup> /d	N/A	0.5
		Delivery point measures ≤ 100.0 m <sup>3</sup> /day	N/A	1.0
(ii)	Total battery gas (includes produced gas that is vented, flared, or used as fuel)	> 16.9 10 <sup>3</sup> m <sup>3</sup> /day	5.0	3.0
		> 0.5 10 <sup>3</sup> m <sup>3</sup> /day and ≤ 16.9 10 <sup>3</sup> m <sup>3</sup> /day	10.0	3.0
		≤ 0.5 10 <sup>3</sup> m <sup>3</sup> /day	20.0	10.0
(iii)	Total battery water	> 50.0 m <sup>3</sup> /month	5.0	N/A
		≤ 50.0 m <sup>3</sup> /month	20.0	N/A
(iv)	Well oil (proration battery)	Class 1 (high), > 30.0 m <sup>3</sup> /day	5.0	2.0
		Class 2 (medium), > 6.0 m <sup>3</sup> /day and ≤ 30.0 m <sup>3</sup> /day	10.0	2.0
		Class 3 (low), > 2.0 m <sup>3</sup> /day and ≤ 6.0 m <sup>3</sup> /day	20.0	2.0
		Class 4 (stripper), ≤ 2.0 m <sup>3</sup> /day	40.0	2.0

		Flow Rate	Maximum uncertainty of monthly volume ± %	Single point measurement uncertainty ± %
(v)	Well gas (proration battery)	> 16.9 10 <sup>3</sup> m <sup>3</sup> /day	5.0	3.0
		> 0.5 10 <sup>3</sup> m <sup>3</sup> /day and ≤ 16.9 10 <sup>3</sup> m <sup>3</sup> /day	10.0	3.0
		≤ 0.5 10 <sup>3</sup> m <sup>3</sup> /day	20.0	10.0
(vi)	Well water		N/A	10.0

### 1.7.2 Gas Systems

		Flow Rate	Maximum uncertainty of monthly volume ± %	Single point measurement uncertainty ± %
(i)	Gas deliveries (sales gas)		N/A	2.0
(ii)	Hydrocarbon liquid deliveries	Delivery point measures ≤ 100.0 m <sup>3</sup> /day	N/A	1.0
		Delivery point measures > 100.0 m <sup>3</sup> /day	N/A	0.5
(iii)	Plant inlet or total battery/group gas		5.0	3.0
(iv)	Plant inlet or total battery/group condensate (recombined)		N/A	2.0
(v)	Fuel gas	> 0.5 10 <sup>3</sup> m <sup>3</sup> /day	5.0	3.0

		Flow Rate	Maximum uncertainty of monthly volume ± %	Single point measurement uncertainty ± %
		≤ 0.5 10 <sup>3</sup> m <sup>3</sup> /day	20.0	10.0
(vi)	Flare and vent gas		20.0	5.0
(vii)	Acid gas before compression		N/A	10.0
	Acid gas after compression		N/A	3.0
(viii)	Dilution gas		5.0	3.0
(ix)	Well gas (well site separation)	> 16.9 10 <sup>3</sup> m <sup>3</sup> /day	5.0	3.0
		≤ 16.9 10 <sup>3</sup> m <sup>3</sup> /day	10.0	3.0
(x)	Well gas (proration battery)		15.0	3.0
(xi)	Well condensate (recombined)		N/A	2.0
(xii)	Total battery water		5.0	N/A
(xiii)	Well water		N/A	10.0

### 1.7.3 Injection/Disposal Systems

		Maximum uncertainty of monthly volume ± %	Single point measurement uncertainty ± %
(i)	Total gas	5.0	N/A
(ii)	Well gas	N/A	3.0
(iii)	Total water/waste	5.0	N/A

		Maximum uncertainty of monthly volume ± %	Single point measurement uncertainty ± %
(iv)	Well water/steam/waste	N/A	5.0
	Produced water/steam at thermal <i>in situ</i> facilities	N/A	2.0
(v)	Brine disposal well, see <a href="#">Section 15.2.9</a>	N/A	5.0

#### 1.7.4 Heavy Oil - excluding Thermal In Situ Operations (from Section 12)

		Maximum uncertainty of monthly volume ± %	Single point measurement uncertainty ± %
(i)	Gas	N/A	3.0
(ii)	Liquid products received into a cleaning plant, excluding the effect of S&W and density determination	N/A	1.0
(iii)	Sales oil delivery point from a treatment facility	N/A	0.5
(iv)	Test emulsion meter, excluding the effect of S&W determination	N/A	2.0

#### 1.7.5 Thermal In Situ Operations (from Section 12)

		Maximum uncertainty of monthly volume ± %	Single point measurement uncertainty ± %
(i)	Gas production or injection	N/A	3.0

		Maximum uncertainty of monthly volume ± %	Single point measurement uncertainty ± %
(ii)	Emulsion testing using grouped well metering when the subsurface drainage area has coalesced excluding the effects S&W determination		2.0
(iii)	Emulsion test using metering for individual wells - excluding the effects S&W determination	N/A	2.0
(iv)	Clean heavy oil sales	N/A	0.5
(v)	Wellhead steam injection (CWE)	N/A	5.0
(vi)	All other steam measurement, including steam leaving a steam plant (CWE)	N/A	2.0
(vii)	Liquid solvent injection	N/A	2.0
(viii)	Fresh, brackish, or produced water delivered to or from an injection facility	N/A	2.0
(ix)	Boiler feed water, boiler blowdown	N/A	2.0
(ix)	Water disposal	N/A	2.0

## 1.8 Measurement Schematics

This section presents the requirements for measurement schematics used for measurement, accounting, and reporting of oil and gas facilities. Measurement schematics are required to ensure measurement, accounting, and reporting compliance and are visual tools showing the current physical layout of the facility. Schematics should be regularly reviewed and used by groups such as operations, engineering, and production accounting to ensure a common understanding. For the purpose of this Directive, process flow diagrams (PFD), piping and instrumentation diagrams, and process and instrumentation diagrams (P&ID) are not considered measurement schematics.

### 1.8.1 Measurement Schematics Requirements

The operator of record, the company who reports the monthly production to ER generally via Petrinex, is responsible for creating, confirming, and revising any measurement schematics. The

well licensee and physical operator shall provide all assistance they can. The schematics must be used by operations and production accounting to ensure that the reported volumes are in compliance with ER's reporting and licensing requirements. How the required information is shown on a measurement schematic is up to the operator of record to decide as long as the measurement schematic is clear and comprehensive.

The operator of record must have measurement schematics for all the existing applicable facilities outlined in Directive PNG017 by April 1, 2018.

For new facilities, the measurement schematic must be created before the facility is brought on active (i.e. start of operations of a facility) and must be reviewed and updated on an annual basis. See Section 1.8.2 of Measurement Schematic update requirements.

The measurement schematic can be stored electronically or in hard-copy format. A master copy of the measurement schematic must be retained at a central location and previous versions must be stored for a minimum of 18 months.

**The measurement schematic must include the following:**

1. Facility name, facility licensee name, and operator name if different, facility licence number (optional), surface location of facility (i.e. legal survey location: 16-06-056-02W2)
  - a. For reporting facilities with satellites, the surface location of the satellite facility ID of the satellite, satellite's licence number (optional), and equipment located at that satellite is required.
2. Unique Well Identifier (UWI), surface location, and well licence (optional).
3. Facility boundaries between each reporting facility with associated reporting facility ID (Petrinex's facility ID) and facility subtypes. For larger facilities, an optional field flow diagram may be used to show facility delineation. See Appendix 8 for an example.
4. Flow lines with flow direction and product type (i.e. emulsion, oil, gas, water, etc.) that move fluids in and out of the facility(s) and those that connect the essential process equipment within the facility, including recycle lines and bypasses to measurement equipment. Identify if oil is tied into a gas system.
5. Flow split or diversion points (headers) with their Legal Survey Location if they are not on a well or facility lease site.
6. Process equipment that changes the state or composition of the fluid(s) within the facility, such as separators, treaters, dehydrators, compressors, sweetening and refrigeration units, etc.
7. Measurement points and storage tanks or vessels that are used for estimating, accounting, or reporting purposes, including:
  - a. Type of instrumentation (charts, EFM, or readouts)
  - b. Type of meter(s) if applicable
  - c. Testing or proving taps required by ER
8. Fuel, flare, or vent take-off points – default to estimated if meter not shown.
9. Energy source (gas, propane, electricity) used for equipment if not metered or estimated as part of total site fuel.

10. Permanent flare points

11. Fresh water sources, such as lakes and rivers

Multiple facilities can be shown on a single page and a representative equipment configuration can be referenced on a measurement schematic where several wells or facilities have the exact same measurement and equipment set up. For example, if a crude oil multi-well proration battery has several satellites and each satellite has several well sites and all satellites and well sites are constructed the same, it is acceptable to show the detailed configuration of one satellite and one wellsite and labelling them as being typical satellite and well site configurations; and providing a list of all the satellite facility IDs on that page unless the facility subtype is 363. The reporting facility ID and all applicable well and facility information must still be listed on the measurement schematic and separate list of all the information is not acceptable unless the facility subtypes are a 313, 371, or 381 as per stated below. Also it is not acceptable to have a separate list of equipment which may be at the site.

**Additional information required on the schematic:**

**Wells**

1. A list of all producing, water source, injection/disposal, and shut-in wells.
2. Reporting event for wells with downhole commingled stratigraphic units or zones.
3. Identify mechanical lift, such as plunger lift, pump jack, etc.
4. Suspended wells are optional, if shown, identify them as suspended.
5. Except for:
  - a. Heavy Crude Oil Paper Battery (facility subtype 313): The well list is not required to be on the measurement schematic but must state how many wells are in the battery and must be available upon request by ER.
  - b. Gas Multi-well Proration SW Saskatchewan Batteries (facility subtypes 363): The well list is not required to be on the measurement schematic but must state how many wells there are on each branch coming into the battery location and must be available upon request by ER.
  - c. Gas Test Battery (facility subtype 371) and Drilling and Completing Battery (facility subtype 381): No measurement schematic is required until the well is tied to a production battery and starts producing.

**Process Equipment**

1. Normally closed valves that can change production flow.
2. Identify if compressors are electric or gas drive. If they are gas drive, then the HP or KW rating is required unless fuel gas is measured as part of total fuel within a facility. Some cross-border facilities may be required to measure fuel for some compressors individually.
3. Normally open valves, such as emergency shutdown valves (ESDs), pressure-control valves (PCVs), and block valves, are not required as they can be considered default flow.
4. Pressure safety valves (PSV) are not required.

### **Measurement Points**

1. Identify non-accounting meters if shown.
2. Originating facility ID or Legal Survey Locations for truck-in receipt points is not required.

### **Storage Tanks and Vessels**

1. Include fluid type for these tanks, vessels, and caverns, such as oil, emulsion, condensate, plant product, waste, or water; tank and vessel capacity may be shown on separate document and should be available upon request
2. Identify if the tank or vessel is underground or default to aboveground.
3. Identify optional non-reporting chemical storage or pop tanks if shown.
4. Identify if the tank or vessel is tied into a vapour recovery system (VRU) or flare system with the default being to vented.

### **Measurement, Accounting and Reporting Plan (MARP)**

A MARP for Thermal In Situ Projects the measurement schematic must include these additional items:

1. blowdown lines
2. ponds – volume and fluid type
3. meter ID and sample point ID
4. tank gauge
5. pumps
6. secondary measurement points

## **1.8.2 Measurement Schematic Updates**

Changes affecting reporting must be redlined on the measurement schematic at the field level when they occur and communicated to the production accountant at a date set by the operator of record to facilitate accurate reporting before the Petrinex submission deadline.

1. Physical changes, such as wells, piping, or equipment additions or removal, require a measurement schematic update.
2. Temporary changes within the same reporting period do not require a measurement schematic update.

The master copy of the measurement schematic must be updated annually to reflect any changes or deletions. There must be verification of the revisions or, if no revisions, confirmation of no change. Documentation of the verification may be stored separately from the measurement schematic but must be available on request.

A facility that is reactivated must have an up-to-date measurement schematic within three months of reactivation or after the implementation period.

### 1.8.3 Measurement Schematic Availability

Schematics must be provided by the operator of record to the following external parties upon request:

1. Facility licensee of the subject facility
2. The company that performs the volumetric reporting for the facility and the well licensee of wells within a reporting facility
3. The company that performs the product and residue gas allocations up to the allocation point(s)
4. ER or other Saskatchewan regulators, Indian Oil and Gas Canada (IOGC) and cross border Regulatory bodies.
5. Licensee of receipt/disposition points—all reporting measurement points for the facility only

### 1.9 Facility Delineation Requirements

Delineation of lease sites and geographic areas into reporting facilities is based on the measurement, accounting, and reporting rules described in this Directive and *Directive PNG032: Volumetric, Valuation and Infrastructure Reporting in Petrinex*.

Facility delineation requires accurate information on process flows and measurement points in the field, as well as a sound understanding of ER facility definitions and facility subtypes outlined in the aforementioned directives.

Multiple measurement points and regulatory flexibility can result in more than one way of delineating some facilities. However, the following general guidelines can be used.

1. All gas and liquid received into and delivered from a facility must be continuously or batch measured in a single phase.
2. Wells and the associated equipment are only linked to and reported under batteries (BT), injection facilities (IF), or water source facilities (WT).
  - a. Gas wells are linked to and reported under gas batteries.
  - b. Crude oil wells and heavy crude oil wells are linked to and reported under crude oil batteries and heavy crude oil batteries, respectively.
  - c. Disposal wells are linked to and reported under disposal facilities.
  - d. Injection wells are linked to and reported under injection facilities.
  - e. Source water wells are linked to and reported under water treatment facilities (WT).
3. Measured and prorated wells must not be linked to the same battery and must be reported under separate reporting codes.

## 2 Calibration and Proving

Metering devices all require various types of maintenance to ensure operating conditions meet the uncertainty requirements outlined in Section 1 – Standard of Accuracy. This Section presents the base requirements and exceptions for maintaining metering devices.

The calibration and proving requirements stipulated in this Directive are applicable to measurement devices used to determine volumes for ER-required accounting and reporting purposes. These requirements are not applicable to measurement devices used only for a licensee's internal accounting purposes. The requirements are considered minimums, and a licensee may choose to apply more stringent requirements.

If a licensee wishes to deviate from these requirements or exceptions other than applying more stringent requirements, see Section 5: Site-Specific Deviation from Base Requirements to determine if the deviation requires submission of an application to and approval by ER.

### 2.1 Frequency

The accuracy of measurement devices may deviate over time, due to wear, changes in operating conditions, changes in ambient conditions, etc. Generally, the more important the accuracy of a measurement device is, the more frequently it must be calibrated or proved. Example: For an annual frequency, if the last calibration was performed in May 2006, the licensee has to perform another calibration by the end of June 2007 (end of the calendar quarter).

#### 2.1.1 Frequency Exceptions

1. If the use or operation of a measurement device requiring monthly, bimonthly or quarterly calibration/proving is suspended for at least seven consecutive days, the scheduled calibration/proving may be delayed by the number of days the device was not in service. Documentation of the amount of time the device was not in service must be kept and made available to ER on request. If this exception is being applied, the licensee must attach a tag to the meter indicating that the exception is in effect and the next scheduled calibration/proving date. This exception is not applicable to measurement devices subject to calibration/proving frequencies that are semiannual or longer.
2. If a liquid meter is removed from service for bench proving but is put on the shelf and not returned to service, the countdown to the next required bench proving does not start until the meter is returned to service. The licensee must attach a tag to the meter indicating the installation date, but leaving the original proving tag intact.
3. ER may request that calibration/proving of a meter be done at any time or may shorten or extend the due date for scheduled calibration/proving, depending on the specific circumstances at a measurement point.

### 2.2 Accuracy of Provers and Calibration Instruments

Provers and other instruments used for calibration of measurement devices must be tested for accuracy prior to first being used or immediately following any repairs (prior to being put back into service) or alterations being conducted on them, and periodically, in accordance with the following:

1. Portable provers must be calibrated every two years using measurement standards that are traceable to the standards listed in Section 2.2.1.
2. Stationary provers must be calibrated every four years using measurement standards that are traceable to the standards listed in Section 2.2.1.
3. Calibration instruments, such as manometers, thermometers, pressure gauges, deadweight testers, electronic testers, etc., must be tested for accuracy every two years against instruments having accuracy traceable to the standards listed in Section 2.2.1.
4. Master meters must be proved quarterly using a calibrated prover. The fluid used to prove the master meter must have properties similar to the fluids measured by the meters it will be used to prove. The master meter must be proved at flow rates that are comparable to the conditions it will be used for.
5. The measurement uncertainty of the proving or calibrating device must be equal to or better than the uncertainty of the device being proved or calibrated.

### **2.2.1 Proving and Calibration Procedure Standards**

The procedures to be followed for these accuracy tests must be designed to provide consistent and repeatable results and must take into consideration the actual operational conditions the device will encounter. The calibration and proving procedures must be in accordance with the following standards:

1. Procedures specified by Measurement Canada (An Agency of Industry Canada),
2. Procedures described in the API Manual of Petroleum Measurement Standards,
3. The device manufacturer's recommended procedures, or
4. Other applicable procedures accepted by an appropriate industry technical standards association.

Records of the foregoing accuracy tests must be kept for a minimum of three years following the expiry of the applicable test and provided to ER on request.

## **2.3 Gas Meters**

### **2.3.1 Gas Meter Calibration Requirements**

The term gas meter is broadly used to describe all of the equipment or devices that are collectively used to arrive at an indication of a gas volume. Typically, various values, such as differential pressure, static pressure and temperature, must be determined and used to calculate a gas volume. Depending on the specific gas meter, each of those values may be determined by individual devices or equipment.

Calibration of gas meter elements requires the instrumentation to be subjected to various actual pressures, temperatures, and other values that are concurrently subjected to the calibration equipment. If the meter element or end device does not indicate the same value as the calibration equipment, adjustments must be made to the meter element and/or end device.

Some meter equipment technologies may require alternative equipment and procedures for calibration, which is acceptable provided that the equipment and procedures are capable of confirming that the meter elements are functioning properly and are sensing and transmitting accurate data to the end device.

Orifice meters are commonly used to measure gas volumes. Gas orifice meters themselves (the meter run and orifice plate-holding device) do not require calibration/proving. However, the associated meter elements and the end devices to which they are connected must be calibrated, as described in Section 2.3.5.

If devices other than orifice meters are used to measure gas, the associated meter elements and the end devices to which they are connected must be calibrated at the same frequency as orifice meters. The required procedures must be designed to provide consistent and repeatable results and must take into consideration the actual operational conditions the device will encounter.

### **2.3.2 Gas Meter Calibration Frequency**

The frequency of meter element calibration and end devices must be:

1. Within the first calendar month of operation of a new meter
2. By the end of the calendar month following installation, after service or repairs have been made to the meter
3. Semiannually thereafter if the meter is used in a gas plant or for sales/delivery point, see Section 1.7.2 for details
4. Annually for all other meters

See Section 2.3.4 for the exceptions that extend calibration frequency.

### **2.3.3 Gas Meter Internal Inspection**

A key contributor to meter accuracy is the condition of the internal components of the gas meter. Examples of internal components are orifice plates, vortex shedder bars, and turbine rotors. The procedure to inspect internal components is:

1. The internal component must be removed from service,
2. It must be inspected,
3. It must be replaced or repaired if found to be damaged, and
4. It can then be placed back in service.

This procedure must be in accordance with the following:

1. The required frequency for inspection of the gas meter primary element is semiannually for gas plant accounting meters and sales/delivery point meters and annually for all other gas meters.
2. Whenever possible, the inspection should be done at the same time as the calibration of the meter elements and end device, but to accommodate operational constraints the inspection may be conducted at any time, provided that the frequency requirement is met.
3. Inspections must be done in accordance with procedures specified by the API, the American Gas Association (AGA), other relevant standards organizations, other applicable industry-accepted procedures, or the device manufacturer's recommended procedures, whichever are most applicable and appropriate.
4. A tag or label must be attached to the meter or end device that identifies the meter serial number, the date of the internal inspection, and any other relevant details.

5. A detailed record of the inspection documenting the condition of the internal components found and any repairs or changes made to the internal components must be kept for at least one year and provided to ER on request.

### 2.3.4 Gas Meter Calibration and Proving Exceptions

1. If the “as found” calibration check of the gas meter confirms that the accuracy of all readings or outputs are within  $\pm 0.25$  per cent of full scale, with the exception of  $\pm 1.0^{\circ}\text{C}$  for the temperature element, no adjustment of the instrumentation is required.
2. If meter elements and end devices have been found to not require adjustment for three consecutive calibrations, as indicated in item 1 above, the minimum time between routine calibrations may be doubled. A tag must be attached to the meter indicating that this exception is being applied and the date of the next scheduled calibration. The records of the calibrations that qualify the meter for this exception must be kept for at least one year and made available to ER on request.
3. If redundant gas meters are installed for a measurement point or redundant meter elements and/or end devices are installed on a single gas meter, the minimum time between routine calibration of the meter elements and end devices may be doubled, provided that daily volumes from each end device are compared at least monthly and found to be within  $\pm 0.25$  per cent of each other. If the daily volumes are not found to be within  $\pm 0.25$  per cent of each other, immediate calibration of both sets of equipment is required. A tag must be attached to the meter indicating that this exception is being applied and the date of the next scheduled calibration. The records of the monthly comparisons and any calibrations that are done must be kept for at least one year and made available to ER on request.
4. If rotary, turbine, or other types of gas meters with internal moving parts are used to measure gas, such as fuel gas, they must be proved at a frequency of once every seven years following an initial proving prior to installation. The calibration of related meter elements must follow Section 2.3.1. These meters must also be proved immediately following any repairs or alterations being conducted on them. The proving may be done with the meter in service, or the meter may be removed from service and proved in a shop at a pressure that is within the normal operating condition for that meter location unless it can be shown that proving at lower pressure conditions will not change the uncertainty of the meter, such as in the scenario of a rotary meter. A tag or label must be attached to the meter that identifies the meter serial number, the date of the proving, and the meter factor determined by the proving. A detailed report indicating the details of the proving operation must be either left with the meter or readily available for inspection by ER. (If the detailed report is left with the meter, the foregoing requirement relating to the tag or label is considered to be met.)
5. For meters used in effluent measurement that require proving, such as a turbine meter, the proving must be performed by using a gas master meter or other provers in single-phase proving runs. For effluent correction factor (ECF) – water gas ratio (WGR) testing, see Section 7.4.
6. If the internal components of gas meters have been found to be clean and undamaged for three consecutive inspections, the minimum time between

inspections may be doubled. When the internal components are found to be dirty or damaged on any subsequent inspection, the frequency for inspections will revert back to the original requirement.

7. If the inspection of internal components of a gas meter requires the meter to be removed from service and there is no meter bypass installed, it is acceptable to defer a scheduled internal component inspection until the next time the gas meter run is shut down, provided that shutting down and depressuring the gas meter run to remove and inspect the internal components would be very disruptive to operations, require excessive flaring, or cause a safety concern, and:
  - a. Previous internal component inspections have proven to be satisfactory; or
  - b. The meter run is installed in a flow stream where the risk of internal component damage is low, e.g., sales gas, fuel gas; or
  - c. The measurement system at the facility provides sufficient assurance, through volumetric and/or statistical analysis, that internal component damage will be detected in a timely manner.
8. In the scenario of an orifice meter, if the orifice plate is mounted in a quick-change (senior) orifice meter assembly and when attempting to conduct an inspection of the orifice plate the fitting is found to be leaking between the chambers such that the meter run must be shut down and depressured to safely remove the orifice plate, it is acceptable to defer a scheduled orifice plate inspection until the next time the gas meter run is shut down, provided that:
  - a. shutting down and depressuring the gas meter run to remove the orifice plate would be very disruptive to operations, require excessive flaring; or
  - b. the orifice meter assembly is scheduled for repairs to be conducted the next time the gas meter run is shut down to eliminate the cause of the leak and allow future scheduled orifice plate inspections to be conducted; and one of the following must be true:
    - i. Previous orifice plate inspections have proven to be satisfactory;
    - ii. The meter run is installed in a flow stream where the risk of orifice plate damage is low, e.g., sales gas, fuel gas, etc.; or
    - iii. The measurement system at the facility provides sufficient assurance, through volumetric and/or statistical analysis, that orifice plate damage will be detected in a timely manner.
9. Internal metering diagnostics may be used to determine if the structural integrity of the primary element is within acceptable operating parameters and checked at the same required intervals as an internal inspection. Then internal inspection is not required until an alarm or error is generated by the device or as recommended by the manufacturer. The licensee must maintain documentation on the diagnostic capability of the measurement system and make it available to ER on request. An initial baseline diagnostic profile must be performed and documented during the commissioning process.
10. Single phase in-line proving of the gas meter may be used to determine if the primary element/meter element is within acceptable operating parameters and

proved at the same required intervals as an internal inspection. Then internal inspection is not required until the uncertainty limits are exceeded.

Should the primary element inspections be deferred in accordance with any of the foregoing exceptions, the licensee must be able to demonstrate to ER, on request, that the situation meets the conditions identified. If these exceptions are being used, it must be clearly indicated on a tag or label attached to the meter or end device. Evidence in battery or facility logs that the internal component inspection has been scheduled for the next shutdown must be available for inspection by ER. For the purposes of these exceptions, shutdown means any scheduled discontinuation of flow through the meter that is of sufficient duration to allow the operations needed to remove and inspect the internal component. If an unscheduled shutdown appears that it will allow sufficient time to conduct internal component inspection operations, the licensee should conduct those inspections prior to the conclusion of this unscheduled shutdown.

### 2.3.5 Orifice Meters with Chart Recorder Calibrations

The procedure for orifice meter chart recorder (meter element and end device) calibration must be in accordance with the following:

1. Pen arc, linkage, pressure stops, and spacing must be inspected and if necessary be adjusted.
2. The differential pressure element must be calibrated at zero, full span, and nine ascending/descending points throughout its range. A zero check of the differential under normal operating pressure must be done before and after the calibration.
3. The static pressure element must be calibrated at:
  - a. zero;
  - b. 50.0 per cent of full span; and
  - c. full span.
4. If a temperature element is in place, the temperature element must be calibrated at three temperatures:
  - a. operating temperature;
  - b. one colder temperature; and
  - c. one warmer temperature.
5. If a thermometer is in place and used to determine flowing gas temperature, the thermometer must be checked at two points:
  - a. operating temperature; and
  - b. one other temperature.

If the thermometer is found not to read within  $\pm 1^{\circ}\text{C}$  it must be replaced.

6. If a thermometer or other temperature measuring device is not left in place and is transported by a licensee to determine flowing gas temperatures at multiple sites, the accuracy of that device must be verified at the same frequency and in the same manner as a thermometer left in place, and the record of verification must be readily available for inspection by ER for a period of one year.

7. Subsequent to the meter calibration, a tag or label must be attached to the meter or end device that identifies the meter serial number, orifice plate size and the date of the calibration.
8. A detailed report indicating the tests conducted on the meter during the calibration and the conditions “as found” and “as left” must be either left with the meter or end device or readily available for inspection by ER. If the detailed report is left with the meter, the foregoing requirement relating to the tag or label is considered to be met.

### 2.3.6 EFM Meter Calibrations

For gas meters using digital (smart) transmitters connected to a remote terminal unit (RTU) or electronic flow measurement (EFM) at measurement points that do not require delivery point measurement, the transmitter must be verified or calibrated at least every five years in accordance with the following conditions:

1. The digital transmitter is used to measure static pressure, differential pressure or temperature. This exception applies only to digital transmitters and does not apply to analog transmitters. Analog transmitters must be calibrated in accordance with the frequency specified in Section 2.3.2 or to the frequency allowed by other exceptions in Section 2.3.4.
2. For digital transmitters that are currently installed at existing facilities, if the last “as found” calibration check (verification) does not necessitate further calibration in accordance with item #1 in Section 2.3.4 (i.e., the accuracy of all outputs were within  $\pm 0.25$  per cent of full scale), then the next verification or calibration must be conducted in not more than five years. The only exception is where digital transmitters are used at a gas plant inlet that has multiple inlet separators. In this case, the digital transmitters on the inlet separators must be verified or calibrated at minimum biennially. For gas plants with a single inlet separator, the digital transmitters on the inlet separator must be verified or calibrated at least every five years.
3. New digital transmitters and previously used digital transmitters that are installed in a new location must be verified or calibrated at the time of installation and after one year of operation. If the first-year verification or calibration results do not necessitate further calibration in accordance with item #1 in Section 2.3.4 (i.e., the accuracy of all outputs were within  $\pm 0.25$  per cent of full scale) then the next verification or calibration must be in no more than five years. The only exception is where digital transmitters are used at a gas plant inlet that has multiple inlet separators. In that case, the digital transmitters on the inlet separators must be verified or calibrated at minimum biennially. For gas plants with a single inlet separator, the digital transmitters on the inlet separator must be verified or calibrated at least every five years.
4. If calibration of a digital transmitter is required after the first year of operation or as a result of a verification check, then the transmitter must be verified or calibrated in the subsequent year. Digital transmitters must be installed, set up, and verified or calibrated in accordance with the procedures described in the most current version of the Industry Measurement Group’s Intelligent Transmitter Commissioning and Verification industry recommended practice. Note that for new or newly installed

digital transmitters, the differential pressure transmitter must be zero verified and adjusted at static operating pressure during the first-year verification or calibration. If the verification or calibration confirms that the zero reading is within  $\pm 0.25$  per cent, then the differential pressure zero does not need to be verified again for the remainder of the five-year term. If the differential pressure zero reading requires adjustment (i.e., reading is  $> \pm 0.25$  per cent), then it must be verified or calibrated in the subsequent year.

5. If the static operating pressure changes  $> \pm 1750.0$  kPa during the five years with no verification or calibration, then the differential pressure must be zero verified and adjusted at the new static operating pressure within the first month of the pressure change.
6. When verifying or calibrating the analog output signal transmitter, it is the analog output to the RTU or EFM system that must be compared to the reference value. Do not decouple the digital transmitter from the analog output to assess only the digital signal.
7. The output signal from the transmitter must match the received value at the RTU/EFM system.
8. A tag must be attached to the transmitter indicating the date of the last verification or calibration and that the EFM verification or calibration frequency described in this section is being applied. The records of the verifications and calibrations that qualify the meter for the extended verification or calibration frequency must be kept for at least five years and made available to ER on request.

The procedure for calibration of an EFM system must be in accordance with the following:

1. For digital transmitters, as defined above, in Section 2.5.4 (1), the differential pressure element must be calibrated at:
  - a. zero;
  - b. 50.0 per cent of calibrated full span; and
  - c. calibrated full span.
2. For analog transmitters, the differential pressure element must be calibrated at:
  - a. zero;
  - b. 50.0 per cent of full span (ascending);
  - c. full span (ascending); and
  - d. either 80.0 per cent and 20.0 per cent or 75.0 per cent and 25.0 per cent of full span (descending).
3. A zero check of the differential under normal operating pressure must be done before and after the calibration.
4. The static pressure element must be calibrated at:
  - a. zero;

- b. 50.0 per cent of full span; and
  - c. full span
5. If a temperature element is in place, the temperature probe must be verified at two temperatures:
  - a. the normal operating temperature; and
  - b. one colder temperature or one warmer temperature.

The temperature probe must be calibrated or replaced if found not to be within  $\pm 1.0^{\circ}\text{C}$ . Additionally, if an EFM system is used, the temperature probe and transmitter must be verified as a single unit, not decoupled and verified separately, and the indicated value of the transmitter that is sent to the EFM should be compared to the reference value.

6. Subsequent to the meter calibration, a tag or label must be attached to the meter or end device that identifies the meter serial number, orifice plate size and the date of the calibration.
7. A detailed report indicating the tests conducted on the meter or end device during the calibration and the conditions “as found” and “as left” must be either left with the meter or end device or readily available for inspection by ER. If the detailed report is left with the meter or end device, the foregoing requirement relating to the tag or label is considered to be met.
8. If data from the meter or end device are sent to another location(s) for flow calculations via DCS, SCADA, RTU, or other means of communication, the reading of the calibration must be verified at the receipt location of such data to ensure accurate data transmission.

## 2.4 Liquid Meters

Oil and other liquid production and disposition volumes except gas well condensate under certain conditions, see Section 2.6, must always be reported as liquid volumes at  $15.0^{\circ}\text{C}$  and either equilibrium vapour pressure or 101.325 kilopascals (kPa) absolute pressure.

### 2.4.1 Liquid Meter Proving

The frequency and methodology for calibrating the secondary meter element are the same as in Section 2.3. The frequency and methodology for inspecting the primary meter element is the same as in Section 2.3.

Meters used to measure hydrocarbons, water, and emulsions are subject to the following general proving requirements. However, there are additional specific requirements depending on the fluid types, as detailed in Sections 2.4 through 2.8.

1. The design and operation of the entire meter system must meet the requirements of this directive.
2. The design and operation of the meter installation must ensure that the conditions of fluid flow through the meter are within the manufacturer’s recommended operating range.
3. The meter must be installed upstream of a snap acting control/dump valve, if present.

4. The size of the prover taps and operation of the prover must not restrict or alter the normal flow through the meter. Tank-type volumetric or gravimetric provers must be connected downstream of the meter and downstream of a snap acting control/dump valve, but other provers, such as ball provers, pipe provers, or master meters, may be connected either upstream (provided there is no gas breakout) or downstream of the meter and upstream of a snap acting control/dump valve. The location of the proving taps will dictate the proving method(s) that can be used.
5. A new hydrocarbon meter must be proved within the first calendar month of operation or immediately following any repairs being conducted on the meter or any changes to the meter installation. Note that the resultant meter factor must be applied back to the volumes measured after the commencement of operation, repair, or change. A new water meter must be proved within the first three months of operation or immediately following any repairs being conducted on the meter or any changes to the meter installation and no retroactive application of meter factor is required.
6. The meters must be proved according to the frequency in [Table 2.1](#).

**Table 2.1 Meter proving frequency requirements and proving methods**

Application	Meter type	Proving method					Proving frequency
		Pipe/compact/small volume prover	Master meter	Volumetric vessel/tank prover	Bench proving	Calibrate transmitter	
Live oil/condensate (meter at well/battery or test meter)	PD/turbine	A <sup>1</sup>	A	A	A <sup>2</sup>	N/A	Annual <sup>7</sup>
	Vortex/Coriolis	A <sup>3</sup>	A <sup>3</sup>	A <sup>3</sup>	A <sup>2,3</sup>	N/A	Annual <sup>7</sup>
	Differential producer	N/A	N/A	N/A	N/A	A	Annual
Live oil/condensate (gas plant inlet separator or cross border)	PD/turbine	A <sup>3</sup>	A	A	A <sup>2</sup>	N/A	Semi-annual
	Vortex/Coriolis	A <sup>3</sup>	A <sup>3</sup>	A <sup>3</sup>	A <sup>2,3</sup>	N/A	Semi-annual
	Differential producer	N/A	N/A	N/A	N/A	A	Semi-annual
Dead oil, stable HVP liquids, or delivery points <sup>4</sup>	PD/turbine	A	A	A <sup>5</sup>	N/A	N/A	Monthly <sup>6</sup>
	Coriolis/ultrasonic	A	A	A <sup>5</sup>	N/A	N/A	Monthly <sup>6</sup>
Water	PD/turbine	A	A	A	A <sup>2</sup>	N/A	Annual <sup>7</sup>
	Vortex/Coriolis/magnetic/ultrasonic	A <sup>3</sup>	A <sup>3</sup>	A <sup>3</sup>	A <sup>2,3</sup>	NA	Annual <sup>7</sup>
	Differential producer	N/A	N/A	N/A	N/A	A	Annual

<sup>1</sup> A = acceptable method; N/A = not applicable.

<sup>2</sup> See Sections 2.5.1, 2.6.2, and 2.8 for bench proving information.

<sup>3</sup> For meter proving exceptions, see Section 2.4.2.

<sup>4</sup> A delivery point may be emulsion, crude oil, crude bitumen, condensate, LPGs, ethane, or NGLs.

- <sup>5</sup> For live oil/condensate delivery point only.
- <sup>6</sup> If flow is < than 100.0 m<sup>3</sup>/day, quarterly proving is acceptable. For other exceptions, see Section 2.4.2.
- <sup>7</sup> If flow is ≤ 2.0 m<sup>3</sup>/day, biennial proving is acceptable
7. The meter must be proved in line at normal operating conditions unless otherwise exempt by ER.
  8. If a master meter is used for proving, it must have an uncertainty rating equal to or better than the meter it is being used to prove.
  9. Each proving run must consist of a representative volume of the normally metered fluid being directed into the prover or through the master meter.
  10. If a meter is proved after a period of regular operation, an “as found” proving run must be performed prior to conducting any repairs on the meter or replacing the meter.
  11. An acceptable initial proving, also referred to as the first proving of a new or repaired meter, and all subsequent proving must consist of the number of consecutive runs, each with a meter factor (MF) that is within the specified deviation range of the average MF of all applicable runs, as specified in Table 2.2. The resultant meter factor will be the average of all the applicable run meter factors. Proving procedures using more than the specified number of runs are allowed, provided that the licensee can demonstrate that the alternative procedures provide a meter factor of equal or better accuracy.

**Table 2.2 Proving requirements for hydrocarbons, water, and emulsions**

Hydrocarbon meter type	Initial prove: number of required consecutive runs	Subsequent prove: number of required consecutive runs		Maximum MF deviation allowed from average of all applicable runs (%)
		As found MF ≤ ± 0.5% of previous	As found MF > ± 0.5% of previous	
Live oil – field proving (see 2.5)	4	1	4	1.5
Live oil – shop proving (see 2.5)	4	4	4	0.5
Dead oil, condensate at equilibrium, high vapour pressure liquids (see 2.5 & 2.6.1)	3	1	3	0.25
Live condensate – field proving (see 2.6.2)	4	1	4	2
Live condensate – shop proving (see 2.6.2)	4	4	4	0.5
Water– field proving (see 2.8)	4	1	4	1.5
Water– shop proving (see 2.8)	4	4	4	1.5

12. Whenever possible, the inspection of internal components should be done at the same time as the meter end device calibration, but to accommodate operational constraints the inspection may be conducted at any time, provided the frequency requirement is met.
13. A detailed report indicating the type of prover or master meter used, the run details, and the calculations conducted during the proving must be either left with the meter or readily available for inspection by ER. If the detailed report is left with the meter, the requirement stated in point #14 relating to the tag or label is considered to be met. If the proving involved the use of a shrinkage factor instead of de-gassing, a copy of the sample analysis must be attached to the proving report.

14. Subsequent to the meter proving, a tag or label must be attached to the meter that identifies the meter serial number, the date of the proving, the type of prover or master meter used, and the average meter factor. If the meter is connected to an electronic readout, it may be possible to program the meter factor into the software to allow the meter to indicate corrected volumes. If the meter is connected to a manual readout, it is necessary to apply the meter factor to the observed meter readings to result in corrected volumes.
15. LACT meters may use the proving procedure in API-MPMS, Chapter 4: Proving Systems, instead of the procedure in Section 2.4.

## 2.4.2 Liquid Meter Proving Exceptions

1. If a meter used to measure fluids at flowline conditions is a type that uses no internal moving parts, e.g., orifice meter, vortex meter, cone meter, Coriolis meter or ultrasonic meter, it does not require proving, provided that all of the following conditions are met:
  - a. The measurement point does not require delivery point measurement uncertainty (e.g., an oil-well test meter). In some cases, hydrocarbon fluid measurement at separator or flowline conditions requires delivery point measurement uncertainty (e.g., a gas-well producing oil).
  - b. The flow through the meter must be continuous (not intermittent) or the meter must qualify for bench proving or be a Coriolis-type meter with sufficient structural integrity as determined by internal diagnostics (see point c) and maintained within the rates specified by the meter manufacturer as providing accurate measurement. This exception does not apply to master meter proving (see Section 2.2 for requirement). If there is a dump valve as part of the Coriolis or bench-proved measurement system, the dump valve must be checked for leaks at the same proving frequency set out in [Table 2.1](#).
  - c. The design and operation of the entire meter system must be in accordance with the meter manufacturer's specifications.
  - d. The meter element/end device(s) must be calibrated at the frequencies specified in [Section 2.3](#), using procedures specified in [Section 2.3](#), by the API MPMS, the AGA, the device manufacturer, or other applicable industry-accepted procedures, whichever are most appropriate and applicable.
  - e. The internal components of the primary element must be removed from service at the same frequency as indicated in [Table 2.1](#), inspected, replaced or repaired if found to be damaged, and then placed back in service, in accordance with procedures specified by the API, the AGA, other relevant standards organizations, other applicable industry-accepted procedures, or the device manufacturer's recommended procedures, whichever are most applicable and appropriate. Internal metering diagnostics may be used to determine if the structural integrity of the primary element is within acceptable operating parameters and checked at the same required intervals as an internal inspection. Then internal inspection is not required until an alarm or error is generated by the device or as recommended by

- the manufacturer. An initial baseline diagnostic profile must be performed and documented during the commissioning process. The licensee must maintain documentation on the diagnostic capability of the measurement system and make that available to ER on request.
- f. If a meter is to be proved just like one with internal moving parts, no internal inspection is required.
  - g. Whenever possible, the inspection of internal components should be done at the same time as the meter end device calibration, but to accommodate operational constraints the inspection may be conducted at any time, provided the frequency requirement is met.
  - h. A tag or label must be attached to the meter (or end device) that identifies the primary device serial number and the date of the calibration.
  - i. A tag or label must be attached to the meter (or end device) that identifies the primary device serial number, the date of the internal component inspection, and any other relevant details (e.g., the size of the orifice plate installed in the meter).
  - j. A detailed report indicating the tests conducted on the meter during the calibration and the conditions “as found” and “as left” must be either left with the meter (or end device) or readily available for inspection by ER. (If the detailed report is left with the meter, the foregoing requirement relating to the tag or label is considered to be met.)
  - k. A detailed record of the internal component inspection documenting their condition “as found” and any repairs or changes made to them must be either left with the meter (or end device) or readily available for inspection by ER. (If the detailed report is left with the meter or readily available, the foregoing requirement relating to the tag or label is considered to be met.)
2. If the volume of fluid measured by a delivery point or LACT meter is  $\leq 100.0 \text{ m}^3/\text{day}$ , the meter proving frequency may be extended to quarterly. The tag attached to the meter must clearly indicate that the meter measures  $\leq 100.0 \text{ m}^3/\text{day}$  and that the meter is on a quarterly proving frequency. The required proving frequency will revert back to monthly if the meter begins measuring volumes  $> 100.0 \text{ m}^3/\text{day}$ .
  3. For delivery point or LACT meters, if the meter factor has been found to be within  $\pm 0.5$  per cent of the average of the meter factors from the previous three consecutive proves, the meter proving frequency may be extended to quarterly. The tag attached to the meter must clearly indicate that the meter has been found to have consistent meter factors and is on a quarterly proving frequency. The required proving frequency will revert back to monthly whenever the meter factor determined during a proving is found to not be within  $\pm 0.5$  per cent of the average of the meter factors from the previous three consecutive proves.
  4. For delivery point meters that measure trucked-in oil, emulsion, and condensate and that have no moving internal parts (e.g., Coriolis meter, ultrasonic meter, orifice meter, vortex meter, cone meter), the meter may be proved semiannually if the current meter factor is within  $\pm 0.5$  per cent of the average of the previous three consecutive factors. The tag attached to the meter must clearly indicate that the meter has been found to have consistent meter factors and is on a semiannual

proving frequency. The required proving frequency will revert back to monthly whenever the meter factor determined during a proving is not within  $\pm 0.5$  percent of the average of the previous three factors. The meter must requalify for the exception before the proving frequency can again be extended to semiannual. The meter must be proved following repairs to the meter changes to the metering installation.

5. If a meter that required internal inspection is used to measure liquid hydrocarbons and no meter bypass is installed, it is acceptable to defer a scheduled internal component inspection until the next time the liquid meter run is shut down, provided that shutting down and depressuring the meter run to remove and inspect the internal components would be very disruptive to operations or present a safety concern and:
  - a. previous internal component inspections have proven to be satisfactory; or
  - b. the meter run is installed in a flow stream where the risk of internal component damage is low, for example, processed or filtered liquids; or
  - c. the measurement system at the facility provides sufficient assurance, through volumetric and/or statistical analysis, that internal component damage will be detected in a timely manner.

## 2.5 Oil Meters

Live oil and dead oil require distinctly different proving procedures:

1. Live oil - Live oil meters are typically those used to measure volumes of oil or oil/water emulsion produced through test separators, but they also include meters used to measure well or group oil or oil/water emulsions that are delivered to other batteries or facilities by pipeline prior to the pressure being reduced to atmospheric. If oil production is measured prior to being reduced to atmospheric pressure, the proving procedures must allow for the volume reduction that will occur when the gas in solution with the live oil is allowed to evolve upon pressure reduction.
2. Dead oil - Dead oil meters are typically those used for delivery point measurement of clean oil that has been de-gassed to atmospheric pressure. These meters may be found measuring oil being pumped from a battery into a pipeline or measuring oil being pumped from a truck into a pipeline terminal, battery, or other facility. No consideration for gas in solution is required when proving meters used to measure dead oil.

### 2.5.1 Additional Proving Requirements for Live Oil Meters

To account for the shrinkage that will occur at the metering point due to the gas held in solution with live oil, the amount of shrinkage must be determined either by physically de-gassing the prover oil volumes or by calculating the shrinkage based on an analysis of a sample of the live oil or a software simulation. Calculation of shrinkage volumes is most often used to mitigate safety and environmental concerns if the live oil volumes are measured at high pressures or if the live oil contains hydrogen sulphide ( $H_2S$ ).

Additional proving requirements for live oil are as follows:

1. If the proving procedure includes de-gassing the prover to physically reduce the pressure of the hydrocarbons to atmospheric pressure:
  - a. The prover must be a tank-type volumetric or gravimetric prover;
  - b. Each proving run must consist of a representative volume of hydrocarbons or hydrocarbons/water emulsion being directed through the meter and into the prover and the liquid volume then being reduced in pressure to atmospheric pressure. The resultant volume determined by the prover, after application of any required correction factors, is divided by the metered volume to determine the meter factor; and
  - c. The amount of time required to de-gas the prover volume and arrive at a stable atmospheric pressure in the prover will vary, depending on the initial fluid pressure and the fluid characteristics.
2. If the proving procedure uses a shrinkage factor, rather than de-gassing, to adjust the prover volume to atmospheric conditions:
  - a. A shrinkage factor representative of the fluid passing through the meter must be determined and used to adjust the meter volumes to atmospheric conditions.
    - i. The shrinkage factor may either be incorporated into the meter factor or be applied to metered volumes after they are adjusted by the meter factor; and
    - ii. The shrinkage factor must be based upon analysis of a sample of the metered fluid taken at normal operating conditions (see Section 14.3);
  - b. Whenever the operating conditions at the meter experience a change that could affect the shrinkage factor by  $> \pm 5.0$  per cent, a new shrinkage factor must be determined based upon analysis of a sample of the metered fluid taken at the new operating conditions. Consideration must be given to proving the meter at the new operating conditions to determine if the meter factor has been affected; and
  - c. The tag attached to the meter must indicate that a shrinkage factor was used instead of de-gassing the prover and whether the shrinkage factor was incorporated into the meter factor or will be applied separately.
3. When proving a test oil meter, a well that is representative of the battery's average well production characteristics must be directed through the test separator for each of the four runs. If there are wells in the battery with production characteristics that vary significantly from the average, consider determining specific meter factors to be used for each of those wells.
4. In the scenario of a test oil meter, the meter factor must include a correction factor to adjust the metered volume to 15.0°C unless the meter is temperature compensated. Although the actual fluid temperature may vary with ambient temperature, it is acceptable to assume that the temperature observed at the time of proving is reasonably representative of the temperature experienced at the meter until the next proving. This requirement does not apply to meter technologies that do not require correction for temperature.

5. In the scenario of a live oil delivery point meter, the meter factor must not include a correction factor for temperature. The meter must either be temperature compensated or a fluid temperature must be taken daily and the metered volume must be corrected to 15.0°C. This requirement does not apply to meter technologies that do not require correction for temperature.

#### 2.5.1.1 Oil Meter Exceptions

1. In situations where individual well production rates are so low that proving a test oil meter in accordance with the requirements listed in Section 2.5.1 would require more than one hour for an individual proving run, it is acceptable to modify the proving procedures. The following modifications, in order of ER's preference, may be used to reduce proving time:
  - a. Produce several wells through the test separator at one time to increase the volume available for the proving runs.
  - b. If the de-gassing procedure is being used, de-gas the first run only, and then use those data to calculate a shrinkage factor, which can be applied to subsequent runs conducted without de-gassing.
  - c. Use the highest rate well for all proving runs.
  - d. Conduct only three proving runs.

The detailed proving report must clearly indicate if any of the aforementioned modifications was used to prove the meter.

2. A live oil meter may be removed from service and bench proved:
  - a. If the meter is used to measure test volumes of non-heavy oil or emulsion and the average rate of flow of oil in the emulsion streams of all the wells tested through the meter is  $\leq 2.0 \text{ m}^3/\text{day}$  and no well is  $> 4.0 \text{ m}^3/\text{day}$  of oil production in the emulsion stream; or
  - b. If the meter is used to measure test volumes of heavy oil or emulsion (density  $\geq 920.0 \text{ kg/m}^3$ )
3. Bench proving must be conducted in accordance with the following in addition to the general procedure in Section 2.4 where applicable:
  - a. The meter installation must be inspected as follows, and corrective action must be taken when required:
    - i. The flow rate through the meter must be observed to verify that it is within the manufacturer's recommended operating ranges; and
    - ii. The dump valve must not be leaking with no flow registered between dumps.
  - b. The bench proving may be conducted with a volumetric or gravimetric prover or with a master meter, as follows:
    - i. Water is typically used as the proving fluid, but varsol or some other light hydrocarbon fluid may be used for the proving; and
    - ii. Corrections for the temperature and pressure of the proving fluid must be made, where applicable.

If the gas held in solution with the fluid produced through the meter is of sufficient volume to significantly affect the fluid volume indicated by the meter, the shrinkage factor must be determined to correct for the effect of the gas in solution and provide that factor to the meter calibration shop so it may be built into the meter factor.

## 2.6 Condensate Meters

Condensate is subject to two different sets of measurement, accounting, and reporting rules. If condensate volumes are measured and delivered at equilibrium vapour pressure, the volume must be determined and reported as a liquid volume at 15.0°C and equilibrium vapour pressure. If live condensate volumes are measured and delivered at flowline conditions, the volume must be determined at flowline pressure and corrected to 15.0°C, but the volume is reported as a gas equivalent volume at standard conditions (101.325 kPa absolute and 15.0°C).

### 2.6.1 Proving Condensate Meters at Equilibrium Conditions

Meters that measure condensate stored and delivered as a liquid at atmospheric pressure or equilibrium pressure are typically delivery point measurement meters and are therefore subject to the same proving requirements and exceptions applicable to meters used for dead oil measurement (see Sections 2.4 and 2.5).

### 2.6.2 Proving Condensate Meters at Flowline Conditions

When a meter that requires proving is used to measure live condensate at flowline conditions, it must be subjected to the proving requirements in Section 2.4.

#### 2.6.2.1 Condensate Meter at Flowline Conditions Proving Exceptions

A meter used to measure live condensate at flowline conditions may be removed from service and bench proved, in accordance with the following:

1. If the meter is used to measure live condensate production on a continuous or intermittent basis, the rate of flow through the meter must be  $\leq 2.0 \text{ m}^3/\text{day}$  or it must be  $\leq 3.0 \text{ m}^3/\text{day}$  with the gas equivalent volume of the daily condensate volume being  $\leq 3.0$  per cent of the daily gas volume related to the condensate production. If the meter is used on a portable test unit, there is no volume limitation, but consideration should be given to proving the meter in line if significant condensate production is observed during the test.
2. The meter installation must be inspected as follows, and corrective action must be taken where required:
  - a. The flow rate through the meter must be observed to verify that it is within the manufacturer's recommended operating ranges; and
  - b. The dump valve must not be leaking with no flow registered between dumps.

## 2.7 Other Liquid Hydrocarbon Meters

Meters used to measure other high vapour pressure liquid hydrocarbons, such as propane, butane, pentanes plus, gas liquid/liquid petroleum gas (NGL/LPG), etc., are subject to the same proving requirements and exceptions set out in Sections 2.4 and 2.6.1.

## 2.8 Water Meters

If a meter is used to measure water production, injection, or disposal or injection of other water-based fluids, in addition to the requirements in Section 2.4:

1. The meter must be installed and proved within the first three months of operation. Note that the meter factor may be assumed to be 1.0000 until the first proving is conducted.
2. The proving may be conducted in line at field operating conditions, or the meter may be removed from service and bench proved using water as the test fluid.

The proving may be conducted using a volumetric prover, a gravimetric prover, or a master meter.

If a meter is proved after a period of regular operation, an “as found” proving run must be performed prior to conducting any repairs on the meter or replacing the meter. An acceptable proving must consist of four consecutive runs one of which may be the “as found” run, each providing a meter factor within  $\pm 1.5$  per cent of the average of the four factors. The resultant meter factor is the average of the four applicable meter factors. Proving procedures using more than four runs will be allowed, provided that the licensee can demonstrate that the alternative procedures provide a meter factor of equal or better accuracy.

Following the meter proving, a tag or label must be attached to the meter that identifies the meter serial number, the date of the proving, and the average meter factor. If the meter is connected to an electronic readout, it may be possible to program the meter factor into the software to allow the meter to indicate corrected volumes. If the meter is connected to a manual readout, it is necessary to apply the meter factor to the observed meter readings to result in corrected volumes.

A detailed report indicating the type of prover or master meter used, the run details, and the calculations conducted during the proving must be either left with the meter or readily available for inspection by ER. If the detailed report is left with the meter, the foregoing requirement relating to the tag or label is considered to be met.

## 2.9 Product Analyzers

If a product analyzer (water cut analyzer) is used to determine water production, it must be calibrated annually using procedures recommended by the manufacturer.

Following the calibration, a tag or label must be attached to the product analyzer that identifies the analyzer serial number and the date of the calibration. A detailed report indicating the calibration procedure used and the calibration details must be either left with the analyzer or readily available for inspection by ER. If the detailed report is left with the analyzer or readily available, the foregoing requirement relating to the tag or label is considered to be met.

## 2.10 Automatic Tank Gauges

### 2.10.1 Inventory Measurement Calibration

If automatic tank gauge devices are used to indicate fluid levels in tanks for monthly inventory measurement, they must be calibrated on site within the first month of operation and annually thereafter. The calibration procedures must adhere to at least one of following list of procedure standards, as available and applicable:

1. The device manufacturer's recommended procedures;
2. Procedures described in the API Manual of Petroleum Measurement Standards; or
3. Other applicable procedures accepted by an appropriate industry technical standards association.

A record of the calibration must be made available to ER on request.

### 2.10.2 Delivery Point Measurement Calibration

If automatic tank gauge devices are used to indicate fluid levels in tanks for delivery point measurement of oil or oil/water emulsion, such as truck volume receipts at batteries/facilities or batch deliveries into a pipeline, they must be calibrated on site within the first month of operation and monthly thereafter. The calibration procedures must be in accordance with the following list of procedure standards, as available and applicable:

1. The device manufacturer's recommended procedures;
2. Procedures described in the API Manual of Petroleum Measurement Standards; or
3. Other applicable procedures accepted by an appropriate industry technical standards association.

A record of the calibration must be made available to ER on request.

#### 2.10.2.1 Delivery Point Calibration Frequency Exception

Where the accuracy of an automatic tank gauge  $\leq \pm 0.5$  per cent of full scale for three consecutive months, the calibration frequency may be extended from monthly to quarterly. The record of calibration must clearly indicate that the device has been found to demonstrate consistent accuracy and is on a quarterly calibration frequency. The records of the calibrations that qualify the device for this exception must be kept and made available to ER on request. The calibration frequency will revert back to monthly whenever the accuracy is  $> \pm 0.5$  per cent of full scale.

## 2.11 Manual Tank Gauges for Oil Measurement

Tank gauging refers to determining levels in a tank and using those levels to calculate a volume increase or decrease in the tank. The level may be determined by using an automatic tank gauge device or by manually determining the level with a gauge tape. In either scenario, the volume of the tank relative to its height at any given point must be determined. This is referred to as the tank calibration, or tank strapping, and results in the creation of a tank gauge table.

### 2.11.1 Inventory Measurement Calibration

If tank gauging is used only for monthly inventory measurement, specific tank calibration procedures are not required. It is acceptable to use gauge tables provided by the tank manufacturer or, if those are unavailable, generic gauge tables applicable to the tank size/type being used.

### 2.11.2 Delivery Point Measurement Calibration

If tank gauging is used for delivery point measurement of oil or oil/water emulsion, such as truck volume receipts at batteries/facilities or batch deliveries into a pipeline, the specific tanks being used must be calibrated on site within the first month of operation and any time the tank is damaged or altered. The calibration must result in the creation of a gauge table for each tank, which must then be used in conjunction with tank gauge readings to determine volumes. Calibration procedures must be in accordance with applicable methods stipulated in the *API Manual of Petroleum Measurement Standards*.

A record of the calibration must be made available to ER on request.

## 2.12 Weigh Scales

Weigh scales used to measure oil/water emulsion and clean oil receipts at batteries, custom treating plants, pipeline terminals, and other facilities must be approved and inspected prior to use, in accordance with Measurement Canada requirements.

Weigh scales must be tested for accuracy in accordance with the following schedule:

1. Monthly;
2. Immediately (by the end of the calendar month) following any incident in which the scale may have been damaged;
3. Immediately (by the end of the calendar month) following any changes or modifications being made to the scale;
4. and
5. The complete set of procedures set out by Measurement Canada for determining weigh scale accuracy must be used following any damage or modifications and at least annually.

The monthly accuracy tests may be done using the complete set of procedures set out by Measurement Canada or, as a minimum, using the following abbreviated procedure:

1. Zero check: Determine if the scale reads zero with no weight on the scale;
2. Add a 10.0 kg standard weight: Determine if the scale reads 10.0 kg;
3. Remove the 10.0 kg standard weight: Determine if the scale returns to zero;
4. Add a test load consisting of 10 000.0 kg of standard weights or, alternatively, durable object(s) of known weight (minimum 5000.0 kg): Determine if the scale reads the correct weight of the test load (acceptable error is  $\pm 0.2$  per cent of the test load);
5. Add a loaded truck, typical of the loads routinely handled by the scale: Note the total weight of the test load and truck;

6. Remove the test load and note the weight of the truck alone: Determine if the scale reading correctly indicates the removal of the test load (acceptable error is  $\pm 0.2$  per cent of the test load); and
7. Remove the truck: Determine if the scale returns to zero with no weight on the scale.

If as a result of the aforementioned tests the weigh scale is found to not be accurate, it must be calibrated and retested until found to be accurate and then sealed by a heavy-duty scale service company. The service company must then send a written report to Measurement Canada documenting the adjustment and/or repairs.

A detailed record of the accuracy tests and any calibration activities must be kept in close proximity to the weigh scale, retained for at least one year, and made available to ER on request. This record must include the following information:

1. Make, model, serial number, and capacity of the weigh scale and any associated equipment;
2. Date of the accuracy test;
3. Details of the tests performed and the results noted; and
4. Details regarding any alterations or calibration performed on the weigh scale.

#### **2.12.1.1 Weigh Scale Test Frequency Exceptions**

1. If the volume of fluid measured by a weigh scale is  $\leq 100.0 \text{ m}^3/\text{day}$ , the monthly accuracy test frequency may be extended to quarterly. The detailed record of the accuracy tests must clearly indicate that the weigh scale measures  $\leq 100.0 \text{ m}^3/\text{day}$  and that the weigh scale is on a quarterly testing frequency. The required testing frequency will revert back to monthly if the weigh scale begins measuring volumes  $> 100.0 \text{ m}^3/\text{day}$ .
2. If the weigh scale has been found to not require calibration adjustments for three consecutive months, the monthly accuracy test frequency may be extended to quarterly. The required accuracy test frequency will revert back to monthly whenever a quarterly accuracy test determines that the weigh scale requires calibration adjustments.

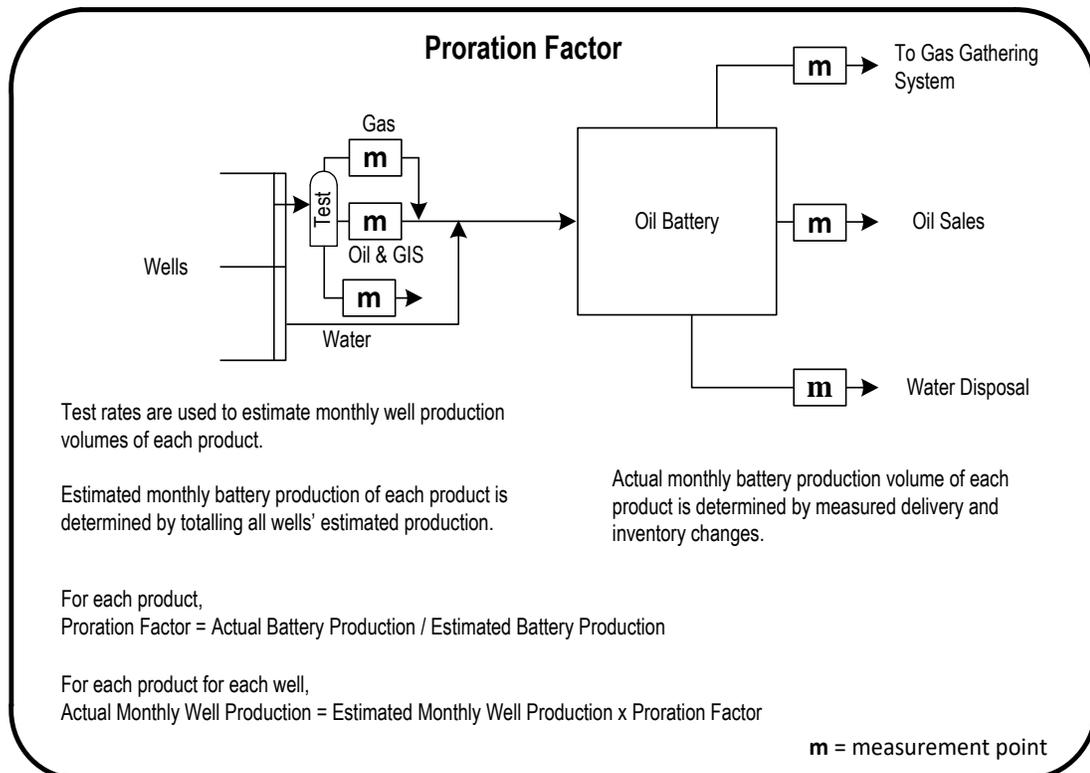
### 3 Proration Factors, Allocation Factors, and Metering Difference

#### 3.1 Proration Factors and Allocation Factors

Proration is an accounting system or procedure where the total actual monthly battery production is equitably distributed among wells in the battery. This system is applicable when the production of wells producing to a battery is commingled before separation and measurement, and each well's monthly production is initially estimated, based on well test data. In this type of system, proration factors are used to correct estimated volumes to actual volumes.

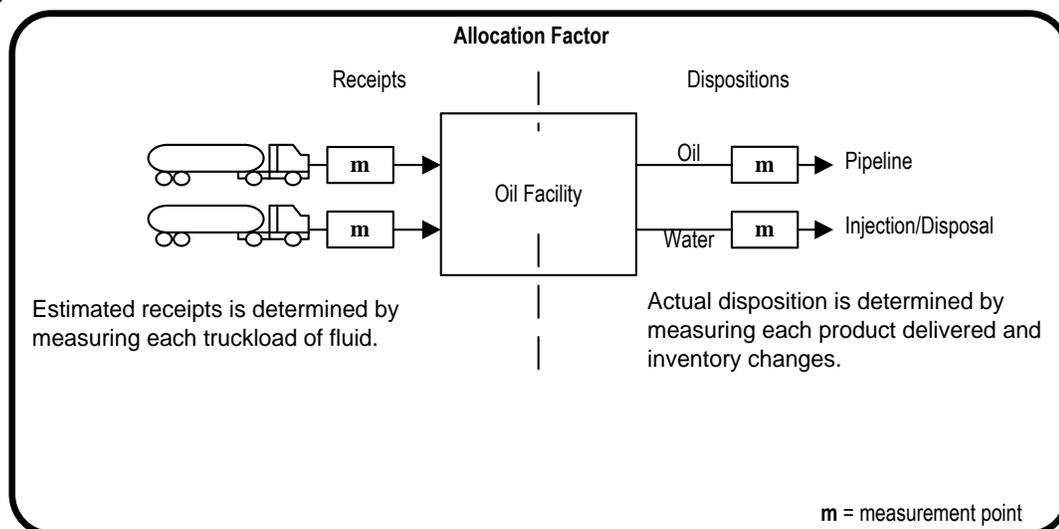
In the scenario of an oil proration battery (Figure 3.1), the oil, gas, and water produced by individual wells are not continuously measured. Instead, the wells are periodically tested to determine the production rates of oil, gas, and water. The rates determined during the well test are used to estimate the well's production for the time period beginning with the well test and continuing until another test is conducted. The estimated monthly production so determined for each well in the battery is totaled to arrive at the battery's total monthly estimated production. The total actual oil, gas, and water production volumes for the battery are determined, and for each fluid, the total actual volume is divided by the total estimated production to yield a proration factor. The proration factor is multiplied by each well's estimated production to yield the well's actual production. Similar accounting procedures are used for gas batteries subject to proration.

Figure 3.1 Proration Factor



An allocation factor is a type of proration factor. It is used at facilities where only fluids received by truck are handled, such as custom treating facilities and third-party operated oil terminals (Figure 3.2). The name of the factor has been chosen to reflect the differences between batteries that receive fluids from wells through flow lines where proration factors are used and facilities that receive fluids from batteries only by truck where allocation factors are used. The purpose of an allocation factor is similar to a proration factor, in that it is used to correct fluid receipt volumes (considered estimates) to actual volumes based on disposition measurements taken at the outlet of the facility and also considering inventory change. The allocation factor is determined by dividing the monthly total actual volume for each fluid by the monthly total estimated volume for each fluid. The total estimated volume of each fluid received from each source is multiplied by the allocation factor for that fluid to yield the actual volume received from that source. Examples of how to calculate and report the proration and allocation factors can be found in *Directive PNG032* and Section 6 of *Directive PNG017*.

**Figure 3.2 Allocation factor**



The allocation factors discussed in this section are not to be confused with the process whereby products delivered out of a gas plant are allocated back to each well in the system, based on individual well production volumes and gas analyses.

Measurement accuracy and uncertainty generally relate to random errors and, as such, are not directly comparable to proration and allocation factors, which generally relate to bias errors. ER's Standards of Accuracy (Section 1) focus on specific measurement points, i.e., inlet or outlet, whereas proration and allocation factors relate to a comparison of inlet (or estimated production) to outlet measurement. It is important to note that the acceptable factor ranges, or targets, for different products may be different due to the products being subjected to different levels of uncertainty. For example, the acceptable factor ranges for oil and water in a non-heavy oil proration battery are different, because while the estimated production volumes of oil and water are determined by the same type of measurement, the outlet volumes of the clean oil and water are not determined by the same type of measurement.

When measurement equipment and procedures conform to all applicable standards, it is assumed that the errors that occur in a series of measurements will be either plus or minus and will cancel each other out to some degree. Where a bias error occurs in a series of measurements, there will be no plus/minus and all of the measurements are assumed to be in

error by the same amount and in the same direction. Proration factors and allocation factors are therefore used to equitably correct all measurements for bias errors.

### 3.1.1 Acceptable Ranges for Proration and Allocation Factors

If measurement and accounting procedures meet applicable requirements, any proration factor or allocation factor should be acceptable, since it is assumed that the factor will correct for a bias error that has occurred. ER requires proration factors and allocation factors to be monitored by licensees and used as a warning flag to identify when the measurement system at a facility is experiencing problems that require investigation.

ER deems the ranges of proration factors and allocation factors indicated in this section to be acceptable ranges. When a factor is found to exceed these limits, the licensee is required to investigate the cause of the factor being outside the acceptable range and document the results of the investigation and the actions taken to correct the situation. Action required by the licensee regarding the investigations into the cause of the proration or allocation factor being outside the acceptable range may include, but is not limited to confirming adherence to:

1. S&W measurement requirements.
2. Related fluid measurement system design, installation and operation requirements.
3. Proving and calibration requirements.
4. The inspections of the primary element of meters used in the proration measurement system.

ER acknowledges that at some facilities, physical limitations or the economics applicable to a particular situation may prohibit the resolution of situations where factors are consistently in excess of the acceptable ranges indicated in this section. In this scenario, the licensee must document the reason(s) that prohibit further action from being taken. This information does not have to be routinely submitted to ER, but must be available to ER on request for audit.

If the cause of a factor being outside these acceptable ranges is determined and the error can be quantified, ER requires the reported volumetric data to be amended, thereby bringing the factor back into line. If the cause is determined and action is taken to correct the situation for future months, but the findings are not quantifiable for past months, amendments are not required to be submitted.

#### 3.1.1.1 Proration Factors

**Table 3.1 Proration factors**

Facility	Oil	Gas	Water
Crude oil battery (facility subtype: 322)	0.95000 – 1.05000	0.90000 – 1.10000	0.90000 – 1.10000
Heavy crude oil battery—primary production and waterflood operations (facility subtype: 327)	0.85000 – 1.15000	no stated expectation due to generally low production volumes	0.85000 – 1.15000

Facility	Oil	Gas	Water
Heavy crude oil battery – thermal recovery operations (facility subtypes: 344)	0.85000 – 1.15000	no stated expectation due to the nature of thermal production	0.85000 – 1.15000
Gas battery – SW Saskatchewan (facility subtypes: 363)		0.80000 – 1.20000	
Gas battery – outside SW Saskatchewan (facility subtype: 364)		0.90000 – 1.10000	0.90000 – 1.10000
Gas battery – effluent measurement (facility subtype: 362)		0.90000 – 1.10000	0.90000 – 1.10000

### 3.1.1.2 Allocation Factors

**Table 3.2 Allocation factors**

Facility	Oil	Gas	Water
Custom Treating facilities (facility subtypes: 611 and 612)	0.95000 – 1.05000		0.90000 – 1.10000
Terminals (facility subtypes 671, 673, 674 and 675)	0.95000 – 1.05000		

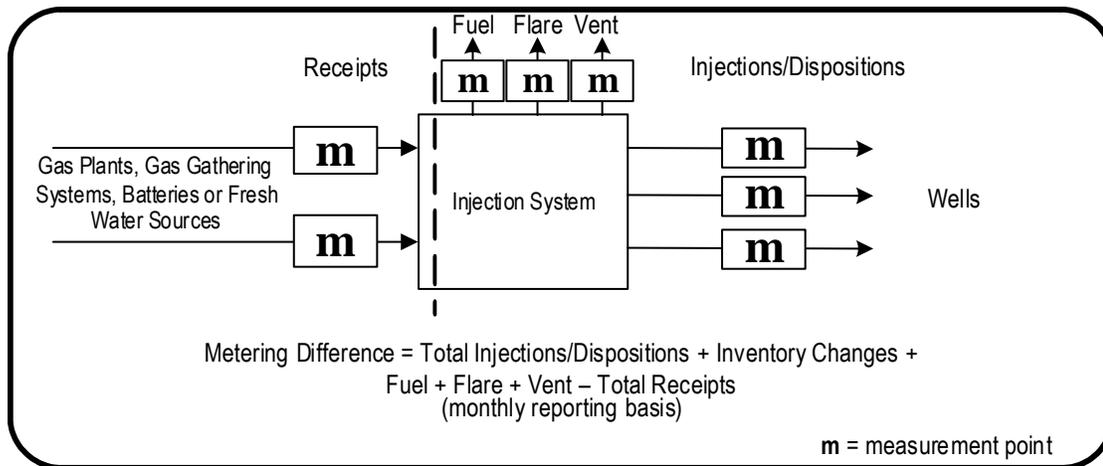
## 3.2 Metering Difference for Fluids other than Oil

For volumetric reporting purposes, a metering difference is used to balance, on a monthly basis, any difference that occurs between the measured inlet/receipt volumes and the measured outlet/disposition volumes at a facility. Metering difference is generally acceptable as an accounting/reporting entity if a difference results from two or more measurements of the same product. Metering differences occur because no two measurement devices provide the exact same volume, due to the uncertainties associated with the devices. However, a more significant cause of metering differences is that the product measured at the inlet to a facility is usually altered by the process within the facility, resulting in a different product or products being measured at the outlet of the facility. It should be noted that metering difference differs from proration and allocation factors in that for facilities where those factors are used, the difference occurs between estimated and actual volumes.

Metering difference may be used as follows:

**Injection/disposal facilities (Figure 3.3)** - Receipts into these facilities are typically measured prior to being split up and delivered to individual wells, where each well's volume is metered prior to injection/disposal.

**Figure 3.3 Injection/disposal facilities**

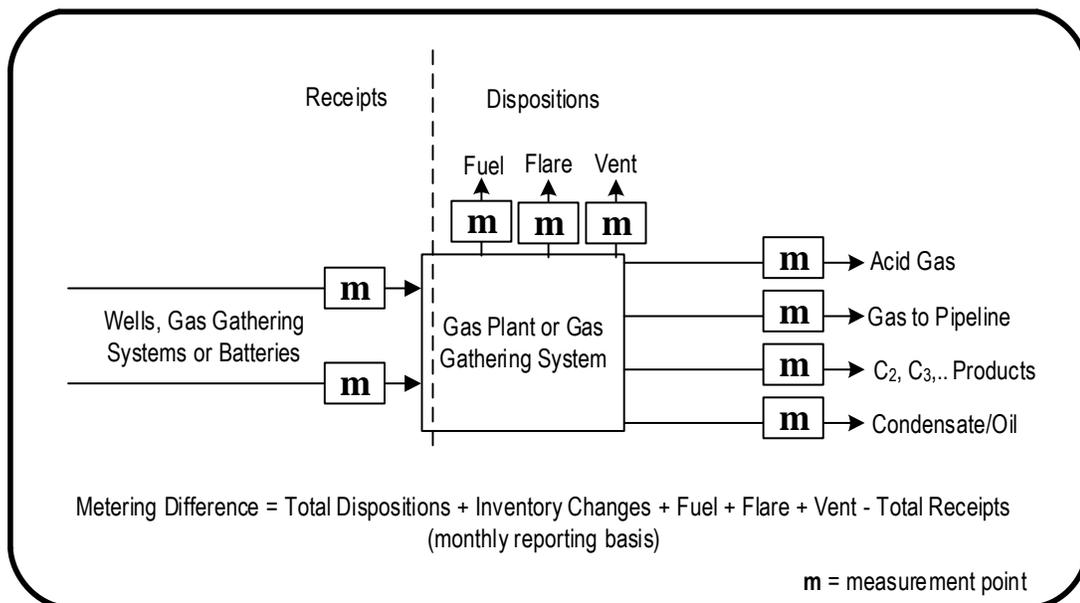


**Batteries** - Metering difference does not apply to any measured or proration battery except gas metering difference for subtype 321 and 326 batteries.

**Gathering systems** (Figure 3.4) - Receipts into these facilities are typically measured prior to being subjected to some sort of limited processing, which may include liquids removal and compression, and the resultant product(s) is measured prior to delivery to a sales point or to a gas plant for further processing.

**Gas plants** (Figure 3.4) - Receipts into these facilities are typically measured prior to being processed into salable products, and those products are measured prior to delivery to a sales point.

**Figure 3.4 Gas plant or gas gathering system**



### 3.2.1 Acceptable Metering Difference Range

If measurement and accounting procedures meet applicable requirements, metering differences up to ±5.0 per cent of the total inlet/receipt volume are deemed to be acceptable. ER requires

the metering difference to be monitored by licensees and used as a warning flag to identify when the measurement system at a facility is experiencing problems that require investigation.

When a metering difference is found to be equal to or greater than  $\pm 5.0$  per cent, the licensee is required to investigate the cause of the metering difference and document the results of the investigation and the actions taken to correct the situation. ER acknowledges that in some facilities, physical limitations and/or the economics applicable to a particular situation may prohibit the resolution of situations where the metering difference is consistently in excess of the range indicated. In such scenarios, the licensee must document the reason(s) that prohibit further action from being taken. This information does not have to be routinely submitted to ER, but must be available to ER on request for audit.

If the cause of an metering difference greater than  $\pm 5.0$  per cent is determined and the error can be quantified, ER requires the incorrectly reported production data to be amended, thereby bringing the metering difference back into an acceptable range. If the cause is determined and action is taken to correct the situation for future months, but the findings are not quantifiable for past months, amendments are not required to be submitted.

## 4 Gas Measurement

This section presents the base requirements and exceptions for gas measurement from any source in the upstream and midstream oil and gas industry that are used for determining volumes for reporting to ER. The term gas includes measurement of hydrocarbon gases as well as non-hydrocarbon gases such as but not limited to hydrogen sulphide, carbon dioxide, helium, hydrogen, and nitrogen.

### 4.1 General Requirements for all Facilities

Unless otherwise stated, the following general gas measurement requirements apply to all Petrinex facility subtypes.

All gas production and injection:

1. must be continuously and accurately metered or
2. where eligible, determined by engineering estimation if the exception conditions described below are met or
3. subject to the conditions of a site-specific ER approval if one has been obtained.

A gas measurement system may deviate from these base requirements if:

1. the conditions in Section 4.4 are met; or
2. the deviation is provided for in Section 1: Standards of Accuracy.

Licensees must report FUEL, FLARE and VENT volumes based on the definitions listed in Directive PNG017 Appendix 2.

Uncombusted gas released to the atmosphere, including fugitive emissions, must be reported as vent gas using the methodologies in *Guideline PNG035: Estimating Venting and Fugitive Emissions*. When a fugitive emission is discovered the licensee must estimate and report the amount of gas released from the time of discovery until the fugitive emission is eliminated. If, at a facility, all gas that is received or produced is vented including casing gas, then no fugitive emissions need to be reported. All documentation relating to the fuel, flare and vent including fugitive emission must be kept for ER to review.

Monthly gas volumes must be reported in units of  $10^3\text{m}^3$  and rounded to 1 decimal place as per *Directive PNG032: Volumetric, Valuation and Infrastructure Reporting in Petrinex*.

Standard conditions for use in calculating and reporting gas volumes are 101.325 kPa absolute and 15.0°C.

1. At all facilities, fuel gas volumes that are  $\leq 0.5 \times 10^3\text{m}^3/\text{day}$ , based on an annual average, may be estimated. If the annual average fuel gas is  $> 0.5 \times 10^3\text{m}^3/\text{day}$ , then the fuel gas must be metered (see Figure 1.11). At facilities where fuel gas metering is required, up to  $0.5 \times 10^3\text{m}^3/\text{day}$  may be estimated. The estimation method must be an acceptable method that is listed in Directive PNG017, *Guideline PNG035: Estimating Venting and Fugitive Emissions* or manufacturer's equipment engineering data recommendations. The licensee must keep the estimation documentation and provide it to ER upon request.

If the site has more than one Petrinex reporting facility, only the fuel for the overall site must be metered; it must then be allocated to and reported for each reporting facility

- provided that the facilities have common working interest ownership and there are no royalty trigger measurement points across the facilities. If the working interest ownership is not common or there are royalty trigger measurement points across the reporting facilities, then any fuel gas volumes crossing reporting facility boundaries must be metered. The only exception is for integrated oilfield waste management facilities (OWMF) with waste plant, custom treater, and injection facilities on the same site, in which case fuel receipt (REC) is to be reported at the waste plant and total OWMF fuel use at the same facility.
2. For non-heavy oil and gas facilities where gas production is  $\leq 0.5 \text{ } 10^3\text{m}^3/\text{day}$ , based on an annual average, may be estimated. If the annual average gas production is  $> 0.5 \text{ } 10^3\text{m}^3/\text{day}$ , then the gas production must be metered (see Figure 1.12). Section 6.5 outlines the acceptable methods for estimating gas production for non-heavy oil facilities. Section 7 outlines the acceptable methods for estimating gas production at gas facilities based on periodic proration testing. For heavy oil facilities where gas production volumes that are  $\leq 2.0 \text{ } 10^3\text{m}^3/\text{day}$ , based on an annual average, may be estimated. If the annual average gas production is  $> 2.0 \text{ } 10^3\text{m}^3/\text{day}$ , then the gas production must be metered. Section 12.2.2 outlines the acceptable methods for estimating gas production. The licensee must keep the estimation documentation and provide it to ER upon request.
  3. For non-heavy oil and gas facilities, flare gas or vent gas volumes that are  $\leq 0.5 \text{ } 10^3\text{m}^3/\text{day}$ , based on an annual average, may be estimated. If the annual average flare gas or vent gas is  $> 0.5 \text{ } 10^3\text{m}^3/\text{day}$ , then the flare gas or vent gas, respectively, must be metered (see Figure 1.12). Facilities requiring flare gas or vent gas metering may estimate up to  $0.5 \text{ } 10^3\text{m}^3/\text{day}$  of each. For heavy oil facilities, flare gas or vent gas volumes that are  $\leq 2.0 \text{ } 10^3\text{m}^3/\text{day}$ , based on an annual average, may be estimated. If the annual average flare gas or vent gas is  $> 2.0 \text{ } 10^3\text{m}^3/\text{day}$ , then the flare gas or vent gas, respectively, must be metered (see Figure 1.12). Facilities requiring flare gas or vent gas metering may estimate up to  $2.0 \text{ } 10^3\text{m}^3/\text{day}$  of each. The estimation method must be an acceptable method that is listed in Directive PNG017, *Guideline PNG035: Estimating Venting and Fugitive Emissions* or manufacturer's equipment engineering data recommendations. The licensee must keep the estimation documentation and provide it to ER upon request.
  4. Gas used for pneumatic devices that is vented or flared must be reported as vented or flared gas, respectively. Licensed and non-licensed facilities must be constructed so that the metered fuel gas does not include pneumatic device gas supply. For facilities that are licensed prior to January 1, 2020 or for non-licensed facilities that were constructed before January 1, 2020, the volume of gas emitted by pneumatic devices may be estimated and then subtracted from the metered fuel gas volume in the case where the metered fuel gas provides the pneumatic device gas supply. The volume that is subtracted from fuel gas does not contribute to the allowance of  $0.5 \text{ } 10^3\text{m}^3/\text{day}$  that may be estimated for flare or vent gas.
  5. Gas used for pilot, purge, sweep, blanket and makeup gas must be reported as flared gas if the gas is combusted. Licensed and non-licensed facilities must be constructed so that the metered fuel gas does not include pilot, purge, sweep, blanket, and makeup gas supply. For facilities that are licensed prior to January 1, 2020 or for non-licensed facilities that were constructed before January 1, 2020, the volume of gas used as pilot, purge, sweep, blanket and makeup gas may be estimated and then subtracted from metered fuel gas in the case where metered fuel gas also provides the pilot, purge and makeup gas supply. The volume

that is subtracted from fuel gas does not contribute to the allowance of 0.5 10<sup>3</sup>m<sup>3</sup>/day that may be estimated for flare gas.

## 4.2 Gas Measurement and Accounting Requirements for Various Facility Subtypes

This section outlines specific requirements for various facility subtypes. General measurement requirements, including meter design, operation, and maintenance requirements are detailed in other sections.

### 4.2.1 Oil Batteries

1. All wells linked to the battery for reporting purposes must be classified as oil wells.
2. Subject to Section 5.5 exception criteria, all wells linked to a multi-well battery for reporting purposes, must be subject to the same type of measurement.
3. Production from gas batteries or other oil batteries may not be connected to an oil proration battery upstream of the oil battery group gas measurement point unless specific criteria are met and/or ER approval of an application is obtained. See Section 5: Site-specific Deviation from Base Requirements, Measurement by Difference.

#### 4.2.1.1 Single-well Battery (facility subtypes: 311 and 325)

1. Gas must be separated from oil, emulsion or water, when present, and metered or estimated separately from the liquid volumes as a single phase.

#### 4.2.1.2 Multi-well Group Battery (facility subtypes: 321 and 326)

1. Each well must have its gas separated from oil or emulsion and metered or estimated as a single phase, similar to a single-well battery.
2. All separation and measurement equipment for the wells in the battery, including the tanks but excluding the wellheads, must share a common surface location.

#### 4.2.1.3 Proration Battery (facility subtypes: 322, 327, and 344)

1. All well production is commingled prior to the total battery gas being separated from oil or emulsion and metered or estimated as a single phase.
2. Individual monthly well gas production is estimated based on well tests and corrected to the reported monthly volume through the use of a proration factor.

#### 4.2.1.4 Crude Oil Multi-well Swab Battery (facility subtypes: 314 and 316)

Monthly gas production from individual crude oil swab wells must be metered or estimated (where applicable) and reported.

### 4.2.2 Gas Batteries

1. All wells linked to the battery for reporting purposes must be classified as gas wells.
2. Gas wells may produce condensate or oil.
3. A mixture of measured and prorated wells (mixed measurement) may be physically connected to the same battery if:

- a. Exception criteria specified in Section 5: Site-specific Deviation from Base Requirements under Measurement by Difference are met, or'
  - b. Site-specific approval has been obtained, and the measured well(s) must have their own separate reporting facility(s) to deliver gas into the proration battery.
4. Well(s) with no phase-separated measurement, including effluent wells, are not allowed to be linked to a Gas Multi-well Group Battery (facility subtype 361).
  5. All gas and recombined liquids from wells linked to a Gas Multiwell Battery (either facility subtypes 361, 362, 363, 364) must be connected by pipeline to a common point.
  6. Gas production from oil wells or oil batteries or from other gas wells or gas batteries must not be connected to a gas proration battery upstream of the gas proration battery group measurement point unless the exception criteria in Section 5: Site-specific Deviation from Base Requirements under Measurement by Difference are met or site-specific approval is obtained.
  7. Well status on Petrinex since gas royalties are based solely on monthly production volumes, only GAS ACTIVE well fluid mode type is used.

#### 4.2.2.1 Single-well Battery (facility subtype: 351)

1. Gas must be separated from water and condensate or oil (if applicable) and continuously measured as a single phase.
2. Condensate produced must be reported as a liquid if it is disposed directly from the battery without further processing. For wells that produce  $\leq 2.0 \text{ m}^3/\text{day}$  of total liquid (i.e., condensate and water) and that direct condensate and water production to lease tanks or to a single emulsion tank, licensees may use the disposition equals production reporting methodology for reporting condensate and water production. This reporting methodology eliminates the requirement to report monthly condensate and water tank inventories. If licensees choose to use this reporting method,
  - a. They must account for existing tank inventories of condensate and water with the initial reporting and
  - b. If the well status is changed to suspended after implementation, the condensate and water tank inventories must be disposed of (i.e., tank emptied) in the reporting month that the well status is changed.

The disposition equals production method of reporting may also be used for water reporting in the case where the separated condensate is recombined with the gas stream and sent to a gathering system and the separated water is directed to a lease tank for disposition.

Refer to Section 12.2.1.1 for a further explanation of the disposition equals production reporting method.

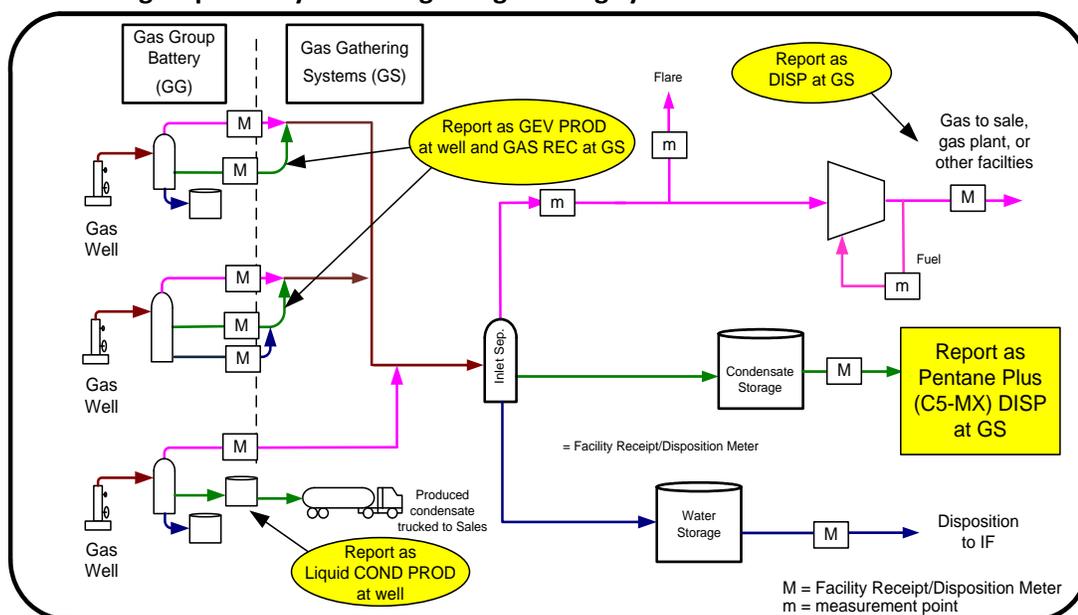
3. Condensate that is recombined with the gas production after separation and measurement must be converted to a gas equivalent volume and added to the measured single-phase gas volume for reporting purposes.
4. Condensate that is trucked from the battery to a gas plant for further processing must be reported as a liquid condensate volume.

- Oil produced in conjunction with the gas must be reported as oil at standard conditions. The gas-in-solution (GIS) with the oil at the point of measurement must be estimated and added to the gas production volume. See Section 6.5.3.

#### 4.2.2.2 Multi-well Group Battery (facility subtypes: 361)

- Each well must have its own separation and measurement equipment, similar to a single-well battery.
- The wells in the group battery may all be identical with regard to the handling of condensate and water, or there may be a mixture of handling methods. The rules for reporting condensate as a gas equivalent or as a liquid are the same as those for single-well gas batteries (see Figure 4.1). The rules for using the disposition equals production reporting methodology for condensate and water are the same as those for single-well gas batteries. See Section 4.2.2.1.
- The volumes measured at each well separator must be used to report the production, PROD, volume on Petrinex. There must not be any proration from any downstream measurement point.

Figure 4.1 Gas group battery delivering to a gathering system



There is no group measurement point requirement for fluids from the gas group wells, but the wells must deliver to a common facility, normally a gas gathering system, with the metering difference reported at the gas gathering system. Hydrocarbon liquids and/or water may be tanked and disposed of by truck and reported as liquid DISP. Recombined hydrocarbon liquids reported as gas equivalent volume and recombined water reported as liquid water must be sent to the same common facility as the gas. Multiple gas groups can deliver to the same gas gathering system.

If the gas gathering system further disposes of the fluids, similar to Figure 4.1, each fluid type (gas, hydrocarbon liquids, water) disposition must be measured and reported. The gas gathering system will also report a metering difference.

See Appendix 9 for schematics of the following scenarios of grouped wells:

Scenario 1 – 1 licensee with 1 reporting entity (1 gas group)

Scenario 2 – 1 to 4 licensees/equity partners with 4 reporting entities (4 single-well batteries) with licensed compressor facility on one well

Scenario 3 – 1 or 2 licensees with 2 reporting entities (1 single-well batteries and 1 gas group)

Scenario 4 – 1 or 2 licensees with 2 reporting entities (gas groups)

Scenario 5 – 1 or 2 licensees with 2 reporting entities (gas groups) with licensed compressor on one well

#### **4.2.2.3 Multi-well Effluent Measurement Battery (facility subtype: 362)**

The definition of and the requirements for facility subtype 362 is found in Section 7. See Section 7 for requirements.

#### **4.2.2.4 Multi-well Proration Battery (facility subtypes: 363 and 364)**

The definition of and the requirements for facility subtypes 363 and 364 are found in Section 7. See Section 7 for requirements.

#### **4.2.3 Gas Gathering System (facility subtypes: 621 and 622)**

A reporting facility consisting of pipelines that move gas and liquids (primarily gas) from one facility to another. A gathering system may also include compressor stations, line heaters, and dehydration equipment located on the system but not associated with any battery, injection facility, gas plant, or other facilities. Inlet measurement usually consists of the battery or facility group measurement point.

Outlet measurement usually consists of the gas plant inlet measurement.

See Section 15.2.1.6 for water reporting requirements.

#### **4.2.4 Gas Plant (facility subtypes: 401, 402, 403, 404, 405, 406)**

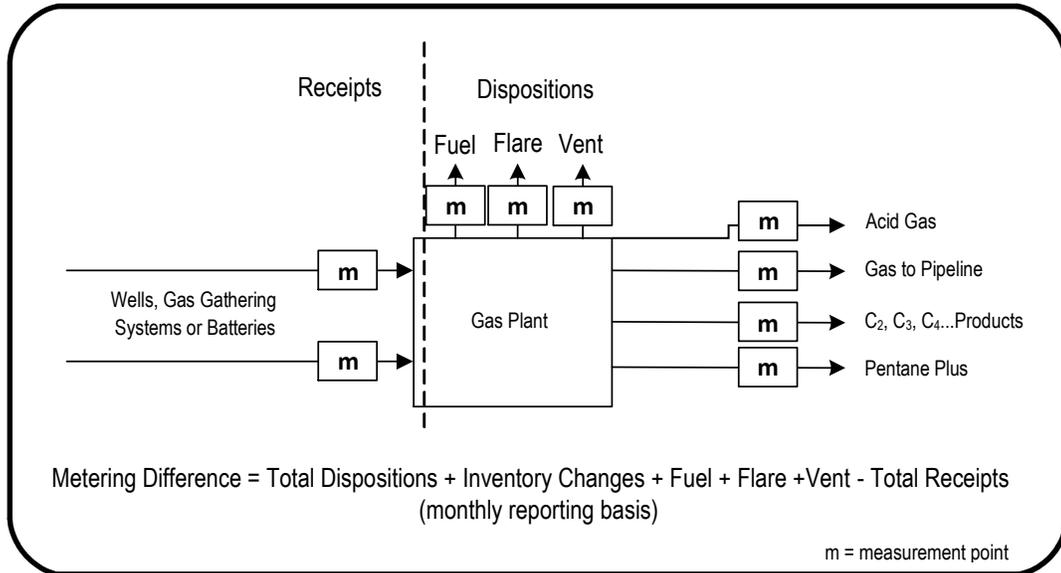
For gas plant facility subtypes definitions see *Directive PNG032: Volumetric, Valuation, and Infrastructure Reporting in Petrinex*.

A gas processing plant (gas plant) is a system or arrangement of equipment used for the extraction of hydrogen sulfide, helium, ethane, natural gas liquids, or other substances from raw gas; does not include a wellhead separator, treater, dehydrator, or production facility that recovers < 2.0 m<sup>3</sup>/day of hydrocarbon liquids without using a liquid extraction process (e.g., refrigeration, desiccant). In addition, does not include an arrangement of equipment that removes small amounts of sulfur (< 0.1 tonne/day) through the use of nonregenerative scavenging chemicals that generate no hydrogen sulfide or sulfur dioxide.

Each plant inlet stream must have inlet separation and continuous measurement for all liquids and gas before commingling with other streams and must be used to report volume on Petrinex for the plant receipt from upstream facilities and for plant balance. However, there are situations where the raw gas has been stripped of its liquid (not recombined downstream) and measured upstream of the plant site. If all streams entering a gas plant on the same gas gathering system are “dry” (the absence of free liquids), the gas plant inlet measurement may consist of the gas gathering system outlet measurement or battery group measurement.

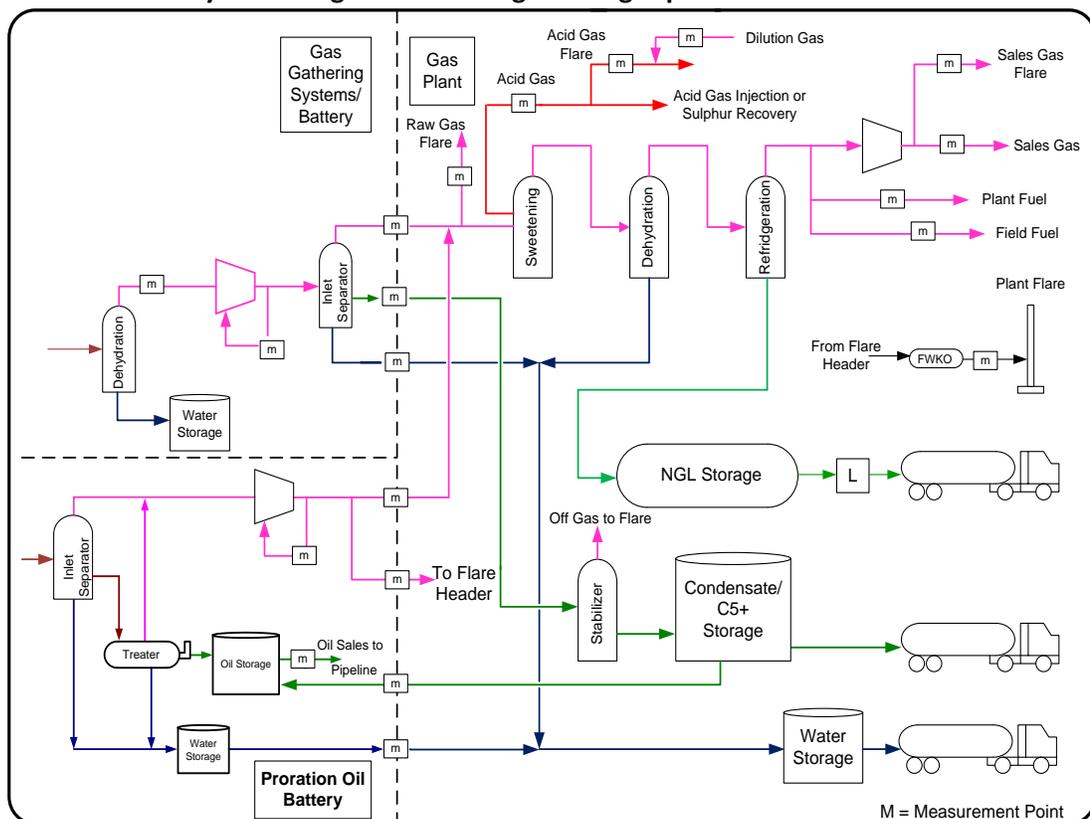
Measurement of all gas dispositions out of the gas plant, such as sales, lease fuel for other facilities, flare and vent gas, acid gas disposition, and any volumes used at the gas plant, is required unless otherwise exempted by ER. Monthly liquid inventory change must be accounted for and reported to Petrinex. (See Figure 4.2.)

**Figure 4.2 Typical gas plant measurement and reporting points**



## Delineation for an Oil Battery Delivering to or Receiving from a Gas Plant on Same Site

Figure 4.3 Oil battery delivering to or receiving from a gas plant



Oil battery gas and water sent to a gas plant for further processing or disposition and gas for flaring must be measured and reported as disposition from the oil battery to the gas plant. The gas plant will report the receipts, total flare, and dispositions.

Gas plant condensate, C<sub>5+</sub>, and/or NGL sent to an oil battery must be measured and reported as disposition to the oil battery. The receipt measurement point at the oil battery is not a royalty trigger point but still requires delivery point measurement.

### 4.2.5 Gas Fractionation Plant (facility subtype: 407)

In addition to the requirements in Section 4.2.4, condensate delivered to a gas fractionation plant must be measured and reported as a liquid receipt in cubic metres.

### 4.2.6 Injection or Disposal Facility (facility subtypes: 501, 503, 504, 505, 506, 507, 510, 511, 512, 514, 516, 517, 518, 519)

Gas must be continuously metered as a single phase. For acid gas injection requirements, see Section 11 - Acid Gas and Sulphur Measurement. For steam injection see Section 12 – Heavy Oil Measurement (Section 12.3).

### 4.2.7 Meter Station (facility subtypes: 631, 632, 633, 634, 640)

Gas must be continuously metered as a single phase.

**4.2.8 Other Facilities (facility subtypes: 204, 207, 208, 210, 211, 212, 213, 214, 371, 381, 671, 673, 674, 675, 676, 701, 702, 703, 904, 905, 906, 907)**

If gas is present, these facilities are required to measure and report the gas. Gas must be measured as a single phase.

**4.3 Base Requirements for Gas Measurement**

**4.3.1 Design and Installation of Gas Measurement Devices**

The design and installation of measurement equipment must be in accordance with the following or as approved by Measurement Canada:

**4.3.1.1 General Design and Installation Requirements**

Measured gas must be corrected to and reported at standard conditions of 101.325 kPa and 15.0°C. Therefore, pressure and temperature measurement devices must be installed in accordance with Section 4.3.3 and 4.3.4.

In some cases, such as the meters listed below, manufacture’s recommended installation is allowed. If manufacture’s recommended meter installation instructions are utilized, the Licensee must provide documentation to ER (if requested) that shows that the meter and the installation meet the uncertainty requirements stipulated in Section 1.

**4.3.1.2 Orifice Meters**

If an orifice meter is used to measure gas, it must be designed and installed according to the applicable AGA Report #3: Orifice Metering of Natural Gas and Other Related Hydrocarbon Fluids (AGA3) listed in Table 4.1.

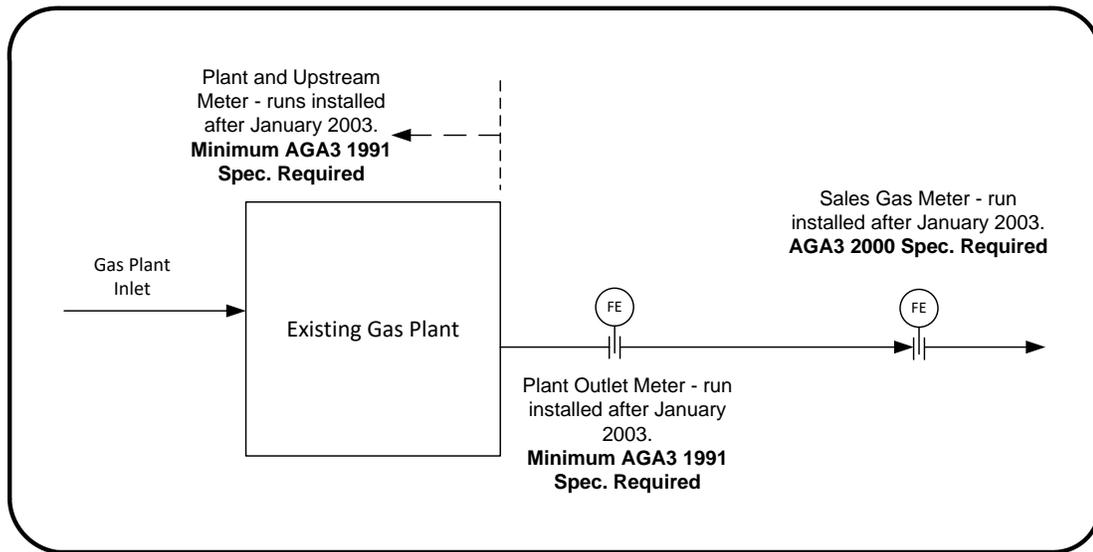
**Table 4.1 Orifice meter design requirement (see detailed explanation in this section)**

<b>Meter run date of manufacture</b>	<b>Applicable AGA3 (API MPMS 14.3, Part 2) version</b>
Before February, 2003	AGA3 1991 or earlier meter run with upstream and/or downstream ID marking – may be reused or relocated for its designed application except to replace a meter where AGA3 2000 spec is required.  Non-AGA meter run or run not marked with upstream or downstream ID – grandfathered for the existing volumetric throughput application, if relocated, it must be refurbished to AGA3 (1985) or later specification but cannot be used for sales/delivery point measurement.
After January, 2003 (except for sales/delivery point meters measuring sales specification processed gas)	February, 1991 or April, 2000, or later
All sales/delivery point meters measuring sales specification	April, 2000 or later

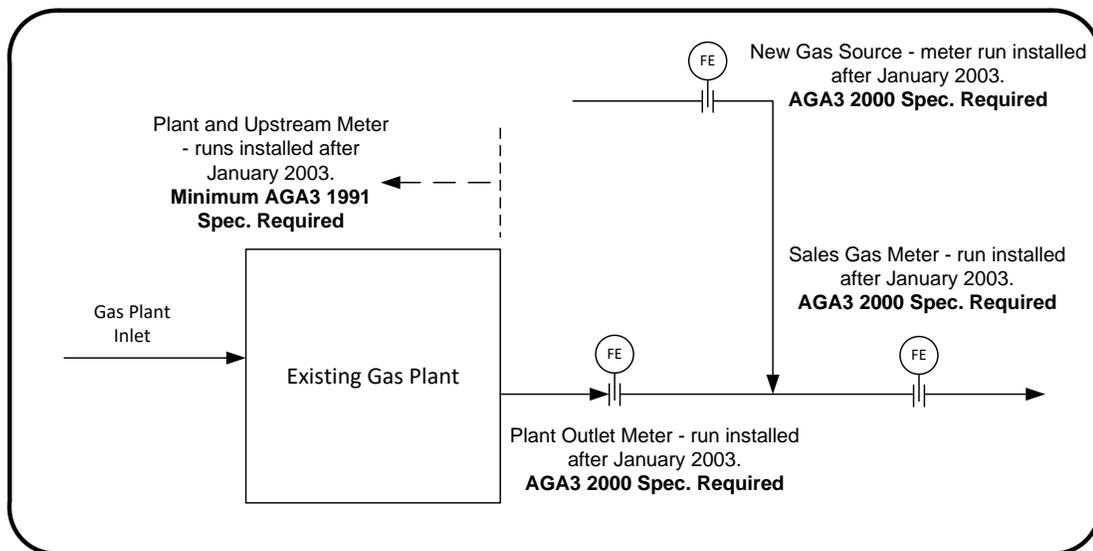
processed gas manufactured after January, 2003	
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When a meter such as a gas plant outlet meter is used to check sales/delivery point measurement and is not normally used to report volumes to ER, it does not require AGA3 April, 2000 specification. However, Licensees are advised that when another gas source ties in to the sales pipeline between the check meter and the sales/delivery point meter, the check meter could become a sales or delivery point meter, and be subject to the requirements of that type of meter as illustrated in Figures 4.4 and 4.5.

**Figure 4.4 Gas plant outlet meter AGA3 1991 specification requirements**



**Figure 4.5 Gas plant outlet meter AGA3 2000 specification requirements**



A permanently marked plate with the following information must be attached to each meter run and maintained in readable condition (not painted over or covered with insulation, etc.) for inspection:

- a. Manufacturer's name
- b. Serial number
- c. Date of manufacture
- d. Average upstream inside diameter (ID) of the meter run at 25.4 mm upstream of the orifice plate in millimetres to one decimal place (or to three decimal places if indicated in inches).
- e. AGA3 version/year (for new runs only after January 31, 2005) and/or AGA3 configuration for runs installed after January, 2003 and not using the April, 2000 specification.

The orifice plate must be permanently marked with the plate bore in millimetres to two decimal places or to three decimal places if indicated in inches, preferably within 6 mm of the outside edge of the plate, to avoid interfering with normal flow if the marking creates a dent or protrusion on the plate surface.

Temperature measurement must be installed according to the applicable AGA3 specifications and the temperature must be determined in accordance with Section 4.3.4.

#### **4.3.1.3 Turbine and Vortex Meters**

Turbine or vortex meters, used to measure gas must be designed and installed according to the provisions of the 1985 or later editions of the AGA Report #7: Measurement of Gas by Turbine Meters (AGA7) or the manufacturer's recommendation and must meet uncertainty requirements stipulated in Section 1.

Temperature measurement is to be installed according to AGA7, i.e., between one and five pipe diameters downstream of the meter or the meter manufacturer's recommendation and the temperature as per Section 4.3.3.

The installation must include instrumentation that allows for continuous pressure, temperature, and compressibility corrections either on site, e.g., electronic correctors, EFM systems or at a later date, e.g., pressure and temperature charts.

#### **4.3.1.4 Rotary Meters**

Rotary metering systems must be designed and installed according to the provisions of the 1992 or later edition of the ANSI B109.3: Rotary Type Gas Displacement Meters or the manufacturer's recommendation and must meet the uncertainty requirements stipulated in Section 1.

#### **4.3.1.5 Diaphragm Meters**

Diaphragm displacement meters must be designed and installed according to the provisions of the 1992 or later edition of the ANSI B109.1: Diaphragm Type Gas Displacement Meters (up to 500 cubic feet/hour capacity), or ANSI B109.2: Diaphragm Type Gas Displacement Meters (over 500 cubic feet/hour capacity), and/or the manufacturer's recommendation and must meet the uncertainty requirements stipulated in Section 1.

#### **4.3.1.6 Venturi or Flow Nozzle Meters**

Venturi or flow nozzle type of meters must be installed according to the provisions of the 1991 or later edition of the ISO Standard 5167: Measurement of fluid flow by means of orifice plates, nozzles and venturi tubes inserted in circular cross-section conduits running full (ISO 5167),

other recognized International Standards or the meter manufacturer's recommendation and must meet the uncertainty requirements stipulated in Section 1.

#### **4.3.1.7 Ultrasonic Meters**

Ultrasonic meters must be designed and installed according to the provisions of the 1998 or later editions of AGA Report No. 9: Measurement of Gas by Multipath Ultrasonic Meters (AGA9), an applicable standard of an appropriate industry technical standards association (ISO, AGA, API) or the meter manufacturer's recommendation and must meet the uncertainty requirements stipulated in Section 1.

Exception:

Ultrasonic metering systems used for flare gas measurement may be installed without live temperature compensation and in accordance with the manufacturer's recommendations. In this scenario the flowing gas temperature must be obtained and updated for volume determination at least once per quarter, the measurement uncertainty requirement for the measurement point stipulated in Section 1 must be met, and the operator must be able to demonstrate to ER, upon request, that the required measurement uncertainty is being met.

#### **4.3.1.8 Coriolis Meters**

Coriolis mass metering systems must be designed and installed according to the provisions of the latest edition of AGA Report No. 11: *Measurement of Natural Gas by Coriolis Meter* and the uncertainty requirements stipulated in Section 1 must be met. External gas sample analysis and density must be used to determine the gas volume at standard conditions (see section 4.3.5).

#### **4.3.1.9 Thermal Mass Meters**

Thermal Mass meters must be designed and installed to applicable standard of an appropriate industry technical standards association or the meter manufacturer's recommendation and must meet the uncertainty requirements stipulated in Section 1. Thermal Mass meters should only be used if:

- a. The density does not change; or
- b. The effect of density change on the volume is within the uncertainty requirements for that application; or
- c. The density can be determined and recorded for flow calculation.

These meters are not recommended for use at gas plant flare stacks unless the above criteria can be met.

#### **4.3.1.10 Other Meters**

If meters other than those listed, for example cones or wedge meters, etc. are used to measure gas, they must be installed according to applicable standard of an appropriate industry technical standards association, the meter manufacturer's recommendation and must meet the uncertainty requirements stipulated in Section 1.

#### **4.3.1.11 Electronic Flow Measurement Systems (EFM)**

See Section 4.3.7.

### 4.3.2 Chart Operations

The chart drive for a circular chart recorder used to measure gas well gas production or group oil battery gas production must not be more than 8 days per cycle unless the exception criteria specified in Section 5: Site-Specific Deviation from Base Requirements are met or ER site-specific approval is obtained. A 24-hour chart drive is required for test gas measured associated with Class 1 and 2 proration oil wells. An 8 day chart drive may be used for test gas measurement associated with Class 3 and 4 proration oil wells. See Section 6.5 Proration Well Testing for more detail on classes of wells. If the mode of operation causes painting on the chart or because of cycling or on/off flows, a 24-hour chart is required for any gas measurement point or EFM must be used.

The Licensee must ensure that:

1. The meter location is properly identified on the chart.
2. The chart is correctly dated.
3. The on and off chart times are recorded on the chart to the nearest quarter hour if not actual.
4. The correct orifice plate and line size are recorded on the chart.
5. The time to the nearest quarter hour of any orifice plate change is indicated on the chart, along with the new orifice plate size.
6. It is noted on the charts if the differential, pressure, or temperature range of the recorder has been changed or if they are different from the ranges printed on the chart
7. The flowing gas temperature is recorded on the chart in accordance with [Table 4.2](#).
8. Proper chart reading instructions are provided when the pen fails to record because of sensing line freezing, clock stoppage, pens out of ink, overlapping traces, or other reasons. Example instructions include:
  - a. drawing in the estimated traces,
  - b. requesting to read as average flow for the missing period, or
  - c. providing an estimate of the differential and static pressure.
9. Any data or traces that require correction must not be covered over or obscured by any means

The Licensee should ensure that:

1. A notation is made on the chart with regard to whether or not the meter is set up for atmospheric pressure for square root charts.
2. The accuracy of the meter clock speed is checked and the chart reader is instructed accordingly of any deviations.
3. The differential pen is zeroed once per chart cycle.
4. Differential pen recordings are at 33.0 per cent or more within the chart range whenever possible.
5. Static pen recordings are at 20.0 per cent or more within the chart range whenever possible.

6. When there is a painted differential band, instructions are provided as to where it should be read. There are various ways to read a painted chart:
  - a. If the differential pen normally records at the top of the painted band but spikes quickly down and up during separator dump cycles, it is reasonable to read the differential near the top of the band or vice versa.
  - b. If the differential pen is in constant up and down motion, it is reasonable to read the differential at the root mean square (RMS) of the band or in a sine wave motion alternating between the top and bottom of the painted area.
7. The pen trace colours conform to the industry-accepted practice (RED for differential, BLUE for static, and GREEN for temperature). However, any colour may be used, provided the colour used is documented.

#### 4.3.2.1 Ministry of Energy and Resources Site-Specific Requests

If an inspection of a measurement device or of procedures reveals unsatisfactory conditions that significantly reduce measurement accuracy, ER will direct that the licensee implement changes to improve measurement accuracy, and this direction will become a condition of operation for that facility or facilities. Examples of unsatisfactory conditions applicable to orifice chart recorders are as follows:

1. Thick pen traces that will cause excessive error when reading the traces.
2. Painting traces.
3. Differential or static pens recording too low on the chart—in scenarios where it is unavoidable because of low flow rate, high shut-in pressure, and equipment or operating pressure range limitations.

#### 4.3.2.2 Chart Reading

The chart integrator/planimeter operator **must** ensure the following:

1. Visible gaps between the integrator/planimeter traces and chart traces are minimized.
2. The counter is read correctly.
3. The integrator is calibrated as per the specified calibration frequency and after each change of pens.
4. The correct integrator or square root planimeter constants are noted.
5. The correct integrator setback is recorded.
6. The correct coefficient, using all of the required factors, is recorded.

#### 4.3.2.3 Digital Chart Reading Technology

Some chart reading technology uses digital scanning technology to scan and store an image of the chart and the use of computer programs to read and interpret the digital image of the chart and the pen traces.

The use of digital technologies to read charts does not require prior approval of ER, but the licensee using any new technology must be able to demonstrate that the following requirements are met:

1. The equipment and/or procedures used to read the chart must not alter or destroy the chart such that it cannot subsequently be read using conventional equipment and/or procedures.
2. The accuracy and repeatability of the new equipment and/or procedures must be equal to or better than conventional equipment and/or procedures.

The following requirements are specific to the use of digital scanning technology for reading charts:

1. The original chart must be retained for at least 12 months, 18 months for gas production associated with heavy oil or crude bitumen, or alternatively the licensee may choose the following procedure for audit trail:
  - a. An original scanned image of the chart, both front and back, must be stored so that it cannot be changed. If the chart back is blank, the back does not need to be scanned provided there is a statement entered in the record to that effect. There must be a method to confirm that a set of front and back scans belong to the same chart if scanned and stored. No alteration or editing of the original scanned image is allowed.
  - b. At least two separate electronic copies of the scanned images must be retained and one copy must be stored off site at a different physical address/location for the applicable required period.

Note that although the ER accepts the electronic submission for audits, other jurisdictions might not. Therefore, the original chart should be kept for other jurisdictional audits.

2. Editing or alterations may only be made to a copy of the original scanned image of the chart. If the edited version is used for accounting purposes, the edited or altered image must be stored for the applicable required period and in the same manner as in item #1 above.
3. An image of the chart showing how the chart pen traces were read or interpreted must be stored for the applicable required period and in the same manner as in item #1 above.
4. If there are any changes or additions to those requirements and recommendations specific to chart scanning, these must be documented and made available for instructing chart analysts. An additional requirement specific to chart scanning is as follows:
  - a. When a differential pen is not zeroed correctly, the zero line must be adjusted to the correct position if it is obvious on the chart such as when the zeroing was out when changing charts but the pen was not adjusted and/or as documented by the licensee. Other situations will require the judgment of the chart analyst and confirmation from the facility licensee. Any zero adjustment must only reposition the zero line and must maintain the entire span of the pen. The distance between the actual zero and the pen trace must not be altered.
5. For ER inspection/audit purposes, the licensee must upon request:
  - a. submit any original paper charts or the scanned original images or make them available for on-line viewing; and
  - b. submit all edited images or make them available for on-line viewing.

Note that the software used to open the scanned images should be readily and freely available on the market. In case there is any specific/proprietary image reader software required to view the scanned and stored chart images, it must also be submitted.

6. Upon request of the licensee, the vendor must demonstrate the accuracy of the scanning and integration technology by performing three consecutive scans, with a rotation of the chart image of about 120.0° before each scan, and integrations of the same chart image. The calculated volumes from each reading must be within  $\pm 0.5$  per cent of the average of the three scans and integrations.
7. ER may check the accuracy of the chart-reading technology and volume calculations by providing charts with known calculated volumes. The volumes determined by the chart reading technology must be within  $\pm 0.5$  per cent of ER's known values.

### 4.3.3 Sensing Line Installation for Differential meters

This section applies to differential meters such as orifice, cone or venturi meters. Note that there are exceptions from these requirements detailed in the next section.

1. Accounting meters using differential pressure sensing devices must be equipped with full port valves at the metering tap on the sensing lines. The valves must be the same size as the sensing lines (12.7 mm [1/2 inch] minimum for meter runs 102.0 mm [4.0 inches] in diameter or larger, and 9.5 mm [3.8 inch] minimum for meter runs less than 102.0 mm). All metering design and installation must ensure that the sensing line diameter does not change from the sensing tap valve to the manifold for deliver point, group point, and sales point measurement.
2. Sharing of metering taps by multiple differential pressure devices is not allowed.
3. A separate set of valve manifolds must be used for each device.
4. Equipment and sensing lines must be suitably winterized to prevent them from freezing.
5. Sensing lines must be self-draining towards the sensing taps to prevent liquid from being trapped in the line if they do not meet the exception criteria for changes in sensing line diameter specified in Section 4.3.2.1.
6. Sensing lines should not exceed 1.0 m in length and should have a slope of not less than 25.4 mm per 300.0 mm from the transmitter to the changer.

#### 4.3.3.1 Exceptions from Requirements for Sensing Line Installation for Differential meters

Grandfathering of existing differential pressure-sensing tap valves installed prior to April 1, 2016 is granted without application unless any of the following situations exist:

1. The metering device is being upgraded, refurbished, and commissioned within a new application or relocated;
2. The metering device does not meet the single point uncertainty limit, as detailed in Section 1: Standards of Accuracy;
3. The metering point is subject to noticeable pulsation effects, such as physical vibration or audible flow noise, or is downstream of a reciprocal compressor on the same site; or
4. The metering point is at a delivery point, group point, sales point, or custody transfer point.

Grandfathering of changes in sensing line diameter from the sensing tap to the manifold, such as drip pots, installed before April 1, 2016, is granted without application unless:

1. The metering device does not meet the single point uncertainty limit, as stipulated in Section 1: Standards of Accuracy;
2. The metering point is subject to noticeable pulsation effects, such as physical vibration or audible flow noise, or is downstream of a reciprocal compressor on the same site;
3. The metering point is at a delivery point, group point, sales point, or custody transfer point; or
4. The fuel measurement point does not have a clean, dry fuel source at a facility, such as a gas plant.

If the current metering installation does not meet the grandfathering requirement, licensees must make any necessary changes required to bring the installation into compliance with this Section by April 1, 2017.

#### 4.3.4 Temperature

The flowing gas temperature must be measured and recorded according to Table 4.2.

**Table 4.2 Temperature reading frequency table for gas measurement**

Minimum temperature reading frequency	Criteria or events
Continuous	Sales/delivery points and/or EFM devices
Daily	$> 16.9 \text{ } 10^3\text{m}^3/\text{day}$
Weekly	$\leq 16.9 \text{ } 10^3\text{m}^3/\text{day}$
Daily	a. Production (proration) volume testing, or b. Nonroutine or emergency flaring and venting

Note that the temperature-measuring element must be installed on the meter run if present or near the meter such that it will be sensing the flowing gas stream temperature. Using the surface temperature of the piping or use a thermowell location where there is normally no flow is not acceptable. A meter equipped with a temperature compensation device is considered to have continuous temperature measurement.

#### 4.3.5 Pressure

Where the pressure at the meter may drop below atmospheric pressure, absolute pressure measurement is required.

#### 4.3.6 Volume Calculations

The gas volume calculations comply if the following requirements are met:

1. If an orifice meter is used to measure gas, the licensee must use the 1985 or later editions of the AGA3 to calculate the gas volumes.
2. If a positive displacement meter or a linear type of meter, such as a turbine, ultrasonic, or vortex meter, is used to measure gas, volumes must be calculated

according to the provisions of the 1985 or later editions of the AGA7. Corrections for static pressure, temperature, and compressibility are required.

Exception:

Meters installed at metering points that do not require delivery point measurement uncertainty and are operating under a fixed pressure setting  $\leq 700$  kPa(g) (i.e., directly downstream of a pressure regulating valve with no process or other equipment installed between the pressure regulating valve and the meter) do not require continuous pressure and temperature compensation. Instead, operating pressure and temperature correction factors may be used for volumetric correction. The meter operating pressure must be obtained and updated annually and the flowing temperature must be obtained and updated quarterly for volume determination.

3. If a venturi or flow nozzle type of meter is used to measure gas, volumes must be calculated according to the provisions of the 1991 or later edition of the ISO 5167 or the meter manufacturer's recommended calculation procedures.
4. If a Coriolis mass meter is used to measure gas, volumes must be calculated from the measured mass flow and the density at standard conditions derived from a representative gas sample analysis, including corrections for compressibility because the flowing density measured by the Coriolis mass meter is of insufficient accuracy in a gas application and must not be used to compute volumes.
5. If meter types other than those listed in the previous points, such as v-cones or wedge meters, are used to measure gas, volumes must be calculated according to the applicable standard of an appropriate industry technical standards association or the meter manufacturer's recommendation.
6. If condensate production from a gas well is required to be reported as a gas equivalent volume, the calculation of the gas equivalent factor must be performed in accordance with the methodologies outlined in Section 8.3. The following are the general requirements:
  - a. The gas equivalent volume (GEV) is to be determined based on the latest condensate sample analysis.
  - b. The gas equivalent volume can be determined using the volume fractions, mole fractions, or mass fractions of the condensate analysis.
  - c. The gas equivalent volume can be determined using all of the individual components in the condensate analysis, or the C<sub>5</sub> and/or heavier components in the sample can be grouped as C<sub>5+</sub>, C<sub>6+</sub>, C<sub>7+</sub> or other heavier component groups. If the heavier components are grouped, the gas equivalent factor for the grouped components must be calculated using the molecular weight and/or relative density of the grouped components.

#### 4.3.6.1 Compressibility Factors Used in Gas Volume Calculations

Produced or injected gas volume measurements must be corrected for pressure, temperature, gas composition, and the compressibility of the natural gas.

Exception: For meters that do not require delivery point measurement uncertainty, correction for compressibility is not required (i.e. a compressibility factor 1.0 can be applied) at operating pressures  $\leq 700.0$  kPa(g).

The AGA8<sup>1</sup> (1992 or later) or Redlich-Kwong with Wichert-Aziz sour gas corrections method should be used for the calculation of the compressibility factors. However, other methods can also be used, provided that the licensee documents the reason for their use. Other methods that could be used are

1. Pitzer et al. with Wichert-Aziz sour gas corrections
2. Dranchuk, Purvis, Robinson with Wichert-Aziz sour gas corrections (Standing and Katz)
3. Dranchuk, Abou-Kassam with Wichert-Aziz sour gas corrections (Starling)
4. Hall, Yarborough with Wichert-Aziz sour gas corrections
5. ISO 20765-2: Natural Gas – Calculation of thermodynamic Properties – Part 2: Single-phase Properties (Gas, Liquid, and Dense Fluid) for Extended Ranges of Application (Groupe Européen de Recherches Gazières, 2008)

ER will also accept the use of methods other than those listed. If others are used, a suitable reference and comparison to the AGA8 (1992) method or to experimental results and the justification for use must be documented and provided to ER for inspection on request.

The AGA8 publication includes several approaches for estimating the properties of natural gas for use in the AGA8 calculation. The full compositional analysis (Detail) method must be used rather than the less accurate partial composition method.

If paper charts are used, the compressibility factor should be calculated at least once for each gas chart cycle. Flow computers and other EFM systems used for gas measurement must calculate and update the compressibility or super compressibility factor at least once for each hourly quantity transaction record and whenever the gas composition is updated. For more information on the frequency of compressibility/supercompressibility factor calculating and updating please refer to API MPMS, chapter 21.1, “Flow Measurement Using Electronic Metering Systems – Electronic Gas Measurement.”

#### **4.3.6.2 Physical Properties of Natural Gas Components**

ER adopts the physical properties contained in the most recent edition of the Gas Processors Suppliers Association (GPSA) *SI Engineering Data Book*<sup>1</sup> or the Gas Processors Association (GPA) 2145<sup>1</sup> publication, whichever is the most current. The licensee must ensure that it is using the up-to-date list. If an EFM system does not have the capability to accept updated physical constants, then the existing set of physical constants may be used; however, that type of EFM system must not be used for measurement of processed gas at measurement points that require delivery point measurement accuracy. For standards, such as AGA8, that have imbedded physical constants different in value from those in GPA 2145 or GPSA *SI Engineering Data Book*<sup>1</sup>, changes to such standards are not required unless they are made by the standards association.

### **4.3.7 Data Verification and Audit Trail**

#### **4.3.7.1 General**

The field data, records, and any calculations or estimations, including EFM calculation or estimations, relating to ER-required production data submitted to Petrinex must be supplied

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<sup>1</sup> See Section 4.6: References for complete bibliographical details for these citations.

upon request by ER. The reported data verification and audit trails must be in accordance with the following:

1. When a bypass around a meter is opened or when, for any reason, gas does not reach the meter or the recording device, a reasonable estimate of the unmeasured volume must be determined, the method used to determine the estimate must be documented, and a record of the event must be made.
2. A record must be maintained that identifies the gas stream being metered, the measurement devices, and all measurements, inputs, times, and events related to the determination of gas volumes. (See [Section 4.3.1 Operations](#) for more detail on orifice chart recorders). If EFM is used, the required data must be collected and retained according to Section 4.4.
3. Test records are any documentation produced in the testing or operation of metering equipment that affects measured volumes. This includes the record containing volume verification and calibration measurements for all secondary and tertiary elements.
4. When a gas metering error is discovered, the licensee of the facility must immediately correct the cause of the error and submit amended monthly production reports to Petrinex to correct all affected gas volumes.
5. All flared and vented gas must be reported as described in *Directive PNG032 – Volumetric, Valuation and Infrastructure Reporting*. Incinerated gas must be reported as FLARE gas if an incinerator is used in place of a flare stack. This does not apply to acid gas streams that are incinerated at a gas plant as part of normal operations; in such cases, the incinerated acid gas must be reported as activity of Flare and product type of ACID GAS.
6. When the fuel usage, flaring, or venting location is within a gas gathering system but is not at a licensed entity:
  - a. it must be reported as an activity associated with the closest licensed facility (e.g., compressor) within the gas gathering system; or
  - b. if there is no applicable licensed facility within the gas gathering system, it must be reported as an activity associated with the gas gathering system itself.
7. Licensees must not prorate or allocate flared and vented volumes that occur at a facility to other upstream facilities and/or well locations.
8. Dilution gas, purge gas, or gas used to maintain a minimum heating value of the flared or incinerated gas must be reported as FLARE. The reported total flare volume must include all of these gas volumes.
9. Production hours for gas wells designed to operate on an on/off cycle basis, such as intermittent timers, pump-off controls, plunger lifts, manual on/off, or pump jacks, that are operating normally and as designed on repeated cycles, and where part of the operation involves shutdown of pump equipment and/or shut-in as part of the repeated cycles, are to be considered on production even when the wells are not flowing. At least one on/off cycle must be completed within a reporting period. Physical well shut-ins that are not part of a repeated cycle and emergency shutdown (ESDs) are considered down time. The operation personnel are required to make a

decision based on the operating environment if the wells are not shut in but may or may not have production.

10. Gas used for instrumentation, pumps, and purging, that is then vented or flared must be reported as vent gas or flare gas, respectively, on a per-site basis.

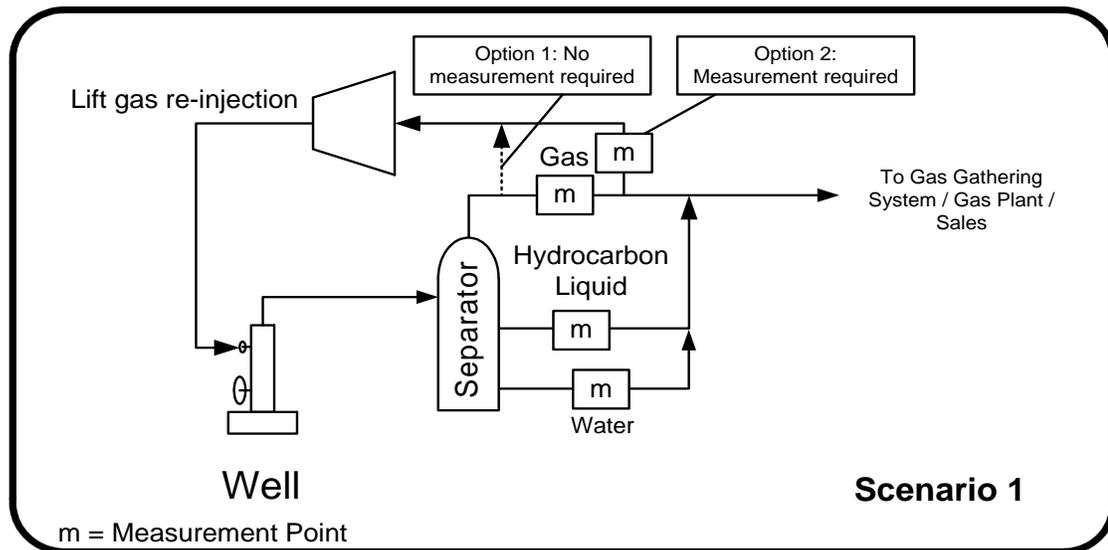
#### 4.3.7.2 Gas Lift Systems for Both Oil and Gas Wells

There are four gas source scenarios, and each one may be subject to different measurement, reporting, and sampling and analysis requirements when gas is injected into the wellbore to assist in lifting the liquids to the surface.

##### Scenario 1

There is no external gas source for the lift gas used given the raw gas is being separated and recirculated continuously at the well with compressor(s). Regular sampling and analysis frequency for the well type applies (see Section 8.4).

Figure 4.6 Lift gas from existing well – scenario 1



**Option 1:** If the lift gas is taken from upstream of the production measurement point, then there is no lift gas reporting requirement.

**Option 2:** If the lift gas is taken from downstream of the production measurement point, then measurement of the lift gas is required and the total well gas production will be the difference between the total measured production volume and the measured lift gas volume.

##### Scenario 2

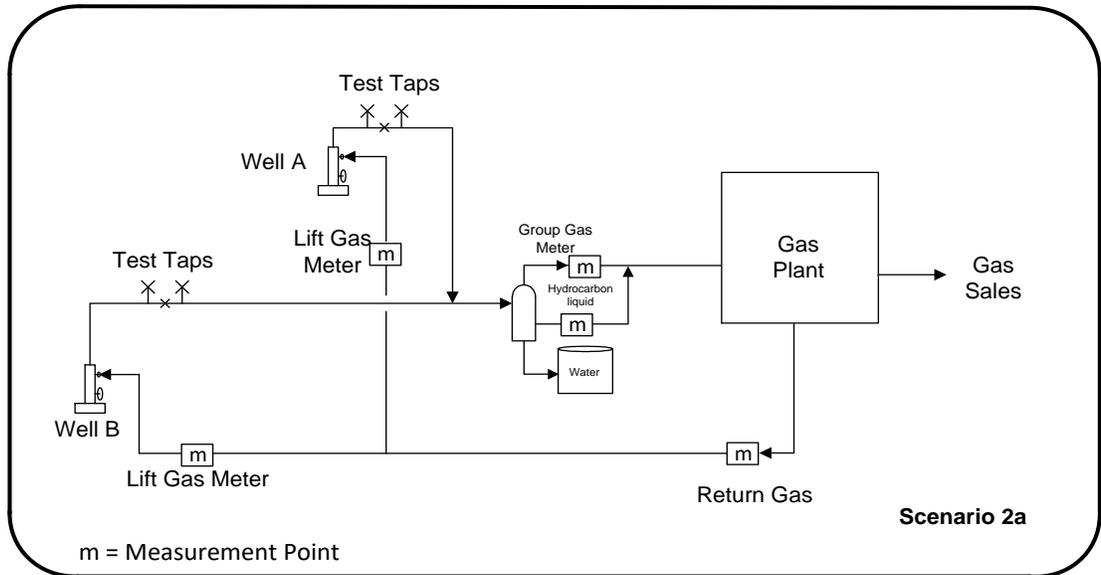
This scenario applies to the lift gas received back from a downstream gas plant/facility that is classified as return gas (no royalty implications).

Measurement is required at the battery level for any gas coming back from the gas plant/facility after sweetening/processing and reported as REC, but such measurement is not delivery point measurement. Part of this return gas could be used for fuel at the well. The lift gas injected into the wellbore must be measured and regular sampling and analysis frequency for the well type applies (see Section 8.4).

There are two possibilities under Scenario 2 (see Figures 4.7 and 4.8).

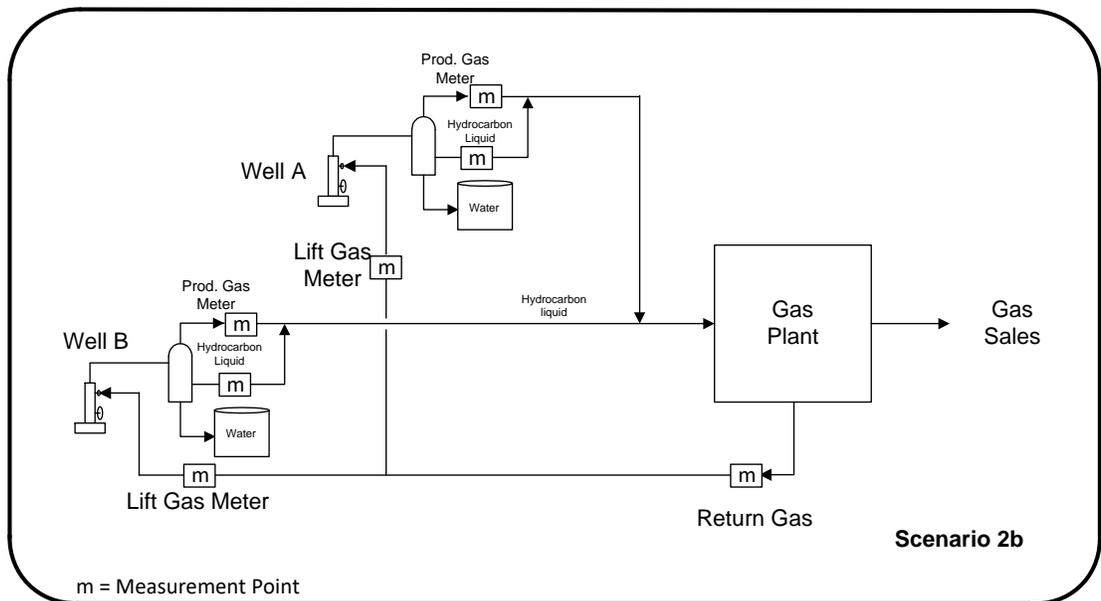
For proration tested wells, the gas lift volume during the test period must be netted off the total test gas production volume to determine the estimated gas production volume for each well.

**Figure 4.7 Lift gas using return gas from plant – scenario 2a**



For continuously measured wells, the gas lift volume must be netted off the total measured gas production volume to determine the actual gas production volume for each well.

**Figure 4.8 Lift gas using return gas from plant – scenario 2b**



**Scenario 3**

This scenario applies to lift gas that comes from royalty exempt sources. The measurement and reporting requirements are the same as Scenario 2.

The gas sampling and analysis frequency for this type of gas lift well is semiannual.

#### 4.4 Electronic Flow Measurement (EFM) for Gas

An EFM system is defined as any flow measurement and related system that collects data and performs flow calculations electronically. If it is part of a Distributed Control System (DCS), Supervisory Control and Data Acquisition system (SCADA) or Programmable Logic Controller system (PLC), only the EFM portion has to meet the requirements in this section.

The following systems are not defined as an EFM:

1. Any meter with an electronic totalizer or pulse counter that does not perform flow calculations with or without built-in temperature compensation.
2. A Remote Terminal Unit (RTU) that transmits any data other than flow data and does not calculate flow.

##### 4.4.1 Base Requirements for EFM

If an EFM is used to calculate volumes for ER accounting purpose, the licensee must be able to verify that it is performing within ER-allowed deviation limits defined in this section.

When any parameter that affects the flow calculation is changed, such as orifice plate size, meter factor, fluid analysis, or transmitter range, a signoff procedure or an event log must be set up to ensure that the change is made in the EFM system. All data and reports must be retained for a minimum of 12 months.

Hardware and software requirements:

1. The EFM data storage capability must exceed the time period used for data transfer from the EFM system.
2. The EFM system must be provided with the capability to retain data in the event of a power failure, e.g., battery backup, UPS, EPROM.
3. System access must have appropriate levels of security, with the highest level of access restricted to authorized personnel.
4. The EFM system must be set to alarm on out-of-range inputs, such as temperature, pressure, differential pressure (if applicable), flow, low power, and communication failures.
5. Any EFM configuration changes or forced inputs that affect measurement computations must be documented either electronically via audit trails or on paper.

The values calculated from forced data must be identified as such.

##### 4.4.2 EFM Performance Evaluation Test

A performance evaluation test must be completed within two weeks after the EFM is put into service and immediately after any change to the computer program or algorithms that affects the flow calculation on a per software version basis. The evaluation must be documented for ER audit purposes on request. For existing EFM systems, ER encourages licensees to conduct their own performance evaluations. A performance evaluation must be conducted and submitted for ER audit on request. ER considers either one of the following methods acceptable for performance evaluation:

1. Conduct a performance evaluation test on the system by inputting known values of flow parameters into the EFM to verify the volume calculation, coefficient factors,

and other parameters. The first seven test cases included in this section are for gas orifice meters (AGA3 flow calculations), each with different flow conditions and gas properties. Test Case 8 is for the AGA7 flow calculation for positive displacement or linear meters. Other manufacturers' recommended equations can also be used to evaluate the EFM performance. The seven AGA3 test cases could also be used to evaluate any compressibility or super compressibility factors used in other flow calculations using the same gas composition, pressure, and temperature in the calculation as inputs.

2. Evaluate the EFM calculation accuracy with a flow calculation checking program that performs within the allowed deviation limits for all the factors and parameters listed in the test cases. A snapshot of the instantaneous flow parameters and factors, flow rates, and configuration information is to be taken from the EFM and input into the checking program. If the instantaneous EFM flow parameters, factors, and flow rates are not updated simultaneously, multiple snapshots may have to be taken to provide a representative evaluation.

Note that some DCS or other control systems have built-in and/or manual input of pressure and temperature for flow calculations. Since the pressure and temperature are not continuously updated, they are not acceptable for ER accounting and reporting purposes unless ER approval is obtained.

The volumetric flow rate (Q) obtained from a performance evaluation test must agree to within  $\pm 0.25$  per cent of those recorded on the sample test cases or other flow calculation checking programs. If the  $\pm 0.25$  per cent limit is exceeded, the EFM must be subjected to a detailed review of the calculation algorithm to resolve the deviation problem. For gas orifice meters, if no AGA3 factor or parameter outputs are available, the acceptable volumetric gas flow rate limit is lowered to  $\pm 0.15$  per cent.

#### **Test Cases 1 to 7 for Verification of Orifice Meter Gas Flow Calculation Programs**

ER has developed test cases to verify that the EFM system correctly calculates gas flow rates from orifice meters. The seven test cases were calculated on the following basis:

1. They are for flange taps only.
2. The atmospheric pressure is assumed to be 93.08 kPa(a) (13.5 psia).
3. The heaviest carbon component was assumed to be normal heptane.
4. The ideal gas relative density was converted to the real gas relative density.
5. The same static pressure value is used for pressure taps that are located upstream (U/S) or downstream (D/S) of the orifice plate.
6. The AGA3 (1985) results were calculated based on upstream conditions for both upstream and downstream static pressure tap in imperial units and the Y2 factor is also provided for reference. The metric conversion factor for the calculated gas volume is 0.02831685. The compressibility factors were calculated using the Redlich-Kwong (RK) equation with the Wichert-Aziz correction for sour gas.
7. The AGA3 (1990) results were calculated using the Detail AGA8 (1992) compressibility factor calculation and using the upstream expansion factor Y1, as recommended by the AGA3 (1990), Part 1, Section 1.8, even though the pressure tap may be downstream of the orifice plate. The Y2 factor is also provided for reference when applicable.

8. The orifice plate material is assumed to be 316 stainless steel and the meter run to be carbon steel at reference temperature of 20°C, isentropic exponent (k) = 1.3, viscosity = 0.010268 centipoise.
9. The standard conditions of 101.325 kPa[abs] and 15.0°C are used in the calculated temperature base factor ( $F_{tb}$ ) and pressure base factor ( $F_{pb}$ ).

**Test Case 8 for Verification of AGA7 Gas Flow Calculation Programs**

ER has developed a test case to verify that the EFM system correctly calculates gas flow rates using the AGA7 equations. The test case was calculated on the following basis:

1. The heaviest carbon component was assumed to be normal heptane.
2. The compressibility factors were calculated using the Detail AGA8 (1992) or the Redlich-Kwong (RK) equation with the Wichert-Aziz correction for sour gas.

**Table 4.3. Allowable deviation limits for the AGA3 (1985) equation**

AGA3 (1985) factors	Allowed deviation limit from test cases
Y, $F_a$ , $F_r$ , and $F_{tf}$	±0.01%
$F_b$	±0.1%
$F_{gr}$ , $F_{pv}$	±0.2%
Q	±0.25% or ±0.15% without the above factors

**Table 4.4. Allowable deviation limits for the AGA3 (1990) equation**

AGA3 (1990) factors	Allowed deviation limit from test cases
$Y_1$ , and $E_v$	±0.01%
$C_d$ and $Z_b$	±0.1%
$Z_f$	±0.2%
Q	±0.25% or ±0.15% without the above factors

**Table 4.5. Allowable deviation limits for the AGA7 equation**

AGA7 factors	Allowed deviation limit from test cases
$F_{pm}$ (flowing pressure) and $F_{tm}$ (flowing temperature)	±0.1%
S (compressibility)	±0.2%
Q	±0.25% or ±0.15% without the above factors

**TEST CASE 1 (for AGA3 Flow Calculations)**

**Gas Analysis**

N <sub>2</sub>	-	0.0184	iC <sub>4</sub>	-	0.0081
CO <sub>2</sub>	-	0.0000	nC <sub>4</sub>	-	0.0190
H <sub>2</sub> S	-	0.0260	iC <sub>5</sub>	-	0.0038
C <sub>1</sub>	-	0.7068	nC <sub>5</sub>	-	0.0043
C <sub>2</sub>	-	0.1414	C <sub>6</sub>	-	0.0026
C <sub>3</sub>	-	0.0674	C <sub>7</sub>	-	0.0022

Ideal gas relative density - 0.7792

**Meter Data (flange taps)**

Meter run I.D.	-	52.370 mm (2.0618 inches)
Orifice I.D.	-	9.525 mm (0.375 inches)

**Flow Data (24 hr)**

Static pressure	-	2818.09 kPa(a) (408.73 psia)
Differential pressure	-	10.2000 kPa (40.9897 inches H <sub>2</sub> O)
Flowing temperature	-	57.0°C (134.600°F)

**Gas Volume Result**

<b>AGA3 (1985)</b>			<b>AGA3 (1990)</b>		
<b>Factors</b>	<b>U/S Tap</b>	<b>D/S Tap</b>	<b>Factors</b>	<b>U/S Tap</b>	<b>D/S Tap</b>
F <sub>b</sub>	28.4286	28.4286	C <sub>d</sub>	0.5990	0.5990
Y <sub>1</sub>	0.9989	0.9989	Y <sub>1</sub>	0.9989	0.9989
Y <sub>2</sub>	N/A	1.0007	Y <sub>2</sub>	N/A	1.0007
F <sub>tb</sub>	0.9981	0.9981	E <sub>v</sub>	1.0005	1.0005
F <sub>gr</sub>	1.1308	1.1308	Z <sub>b</sub>	0.9959	0.9959
F <sub>a</sub>	1.0012	1.0012	Z <sub>r</sub>	0.9280	0.9277
F <sub>r</sub>	1.0006	1.0006	Q	2.7478	2.7531 10 <sup>3</sup> m <sup>3</sup> /24 hr
F <sub>pb</sub>	1.0023	1.0023			
F <sub>tr</sub>	0.9351	0.9351			
F <sub>pv</sub>	1.0360	1.0361			
C'	31.175	31.179			
Q	2.7422	2.7475 10 <sup>3</sup> m <sup>3</sup> /24 hr			

**TEST CASE 2 (for AGA3 Flow Calculations)**

**Gas Analysis**

N <sub>2</sub>	-	0.0156	iC <sub>4</sub>	-	0.0044
CO <sub>2</sub>	-	0.0216	nC <sub>4</sub>	-	0.0075
H <sub>2</sub> S	-	0.1166	iC <sub>5</sub>	-	0.0028
C <sub>1</sub>	-	0.7334	nC <sub>5</sub>	-	0.0024
C <sub>2</sub>	-	0.0697	C <sub>6</sub>	-	0.0017
C <sub>3</sub>	-	0.0228	C <sub>7</sub>	-	0.0015

Ideal gas relative density - 0.7456

**Meter Data (flange taps)**

Meter run I.D.	-	102.26 mm (4.026 inches)
Orifice I.D.	-	47.625 mm (1.875 inches)

**Flow Data (24 hr)**

Static pressure	-	9100.94 kPa(a) (1319.98 psia)
Differential pressure	-	11.0000 kPa (44.2046 inches H <sub>2</sub> O)
Flowing temperature	-	50.0°C (122.0°F)

**Gas Volume Result**

AGA3 (1985)			AGA3 (1990)		
Factors	U/S Tap	D/S Tap	Factors	U/S Tap	D/S Tap
F <sub>b</sub>	733.697	733.697	C <sub>d</sub>	0.6019	0.6019
Y <sub>1</sub>	0.9996	0.9996	Y <sub>1</sub>	0.9996	0.9996
Y <sub>2</sub>	N/A	1.0002	Y <sub>2</sub>	N/A	1.0003
F <sub>tb</sub>	0.9981	0.9981	E <sub>v</sub>	1.0244	1.0244
F <sub>gr</sub>	1.1564	1.1564	Z <sub>b</sub>	0.9967	0.9967
F <sub>a</sub>	1.0010	1.0010	Z <sub>f</sub>	0.8098	0.8097
F <sub>r</sub>	1.0002	1.0002	Q	146.08	146.18 10 <sup>3</sup> m <sup>3</sup> /24 hr
F <sub>pb</sub>	1.0023	1.0023			
F <sub>if</sub>	0.9452	0.9452			
F <sub>pv</sub>	1.1072	1.1073			
C'	888.905	889.000			
Q	145.93	146.03 10 <sup>3</sup> m <sup>3</sup> /24 hr			

**TEST CASE 3 (for AGA3 Flow Calculations)**

**Gas Analysis**

N <sub>2</sub>	-	0.0500	iC <sub>4</sub>	-	0.0000
CO <sub>2</sub>	-	0.1000	nC <sub>4</sub>	-	0.0000
H <sub>2</sub> S	-	0.2000	iC <sub>5</sub>	-	0.0000
C <sub>1</sub>	-	0.6000	nC <sub>5</sub>	-	0.0000
C <sub>2</sub>	-	0.0500	C <sub>6</sub>	-	0.0000
C <sub>3</sub>	-	0.0000	C <sub>7</sub>	-	0.0000

Ideal gas relative density - 0.8199

**Meter Data (flange taps)**

Meter run I.D.	-	590.55 mm (23.250 inches)
Orifice I.D.	-	304.80 mm (12.000 inches)

**Flow Data (24 hr)**

Static pressure	-	10342.14 kPa(a) (1500.00 psia)
Differential pressure	-	22.1600 kPa (89.0522 inches H <sub>2</sub> O)
Flowing temperature	-	60.0°C (140.0°F)

**Gas Volume Result**

AGA3 (1985)			AGA3 (1990)		
Factors	U/S Tap	D/S Tap	Factors	U/S Tap	D/S Tap
F <sub>b</sub>	30429.66	30429.66	C <sub>d</sub>	0.6029	0.6029
Y <sub>1</sub>	0.9993	0.9993	Y <sub>1</sub>	0.9993	0.9993
Y <sub>2</sub>	N/A	1.0004	Y <sub>2</sub>	N/A	1.0004
F <sub>tb</sub>	0.9981	0.9981	E <sub>v</sub>	1.0375	1.0375
F <sub>gr</sub>	1.1028	1.1028	Z <sub>b</sub>	0.9968	0.9968
F <sub>a</sub>	1.0013	1.0013	Z <sub>r</sub>	0.8216	0.8213
F <sub>r</sub>	1.0001	1.0001	Q	8564.77	8575.48 10 <sup>3</sup> m <sup>3</sup> /24 hr
F <sub>pb</sub>	1.0023	1.0023			
F <sub>if</sub>	0.9309	0.9309			
F <sub>pv</sub>	1.1076	1.1078			
C'	34636.6	34643.21			
Q	8603.19	8614.04 10 <sup>3</sup> m <sup>3</sup> /24 hr			

**TEST CASE 4 (for AGA3 Flow Calculations)**

**Gas Analysis**

N <sub>2</sub>	-	0.0029	iC <sub>4</sub>	-	0.0000
CO <sub>2</sub>	-	0.0258	nC <sub>4</sub>	-	0.0000
H <sub>2</sub> S	-	0.0000	iC <sub>5</sub>	-	0.0000
C <sub>1</sub>	-	0.9709	nC <sub>5</sub>	-	0.0000
C <sub>2</sub>	-	0.0003	C <sub>6</sub>	-	0.0000
C <sub>3</sub>	-	0.0001	C <sub>7</sub>	-	0.0000

Ideal gas relative density - 0.5803

**Meter Data (flange taps)**

Meter run I.D.	-	146.36 mm (5.7622 inches)
Orifice I.D.	-	88.900 mm (3.500 inches)

**Flow Data (24 hr)**

Static pressure	-	9839.99 kPa(a) (1427.17 psia)
Differential pressure	-	6.6130 kPa (26.575 inches H <sub>2</sub> O)
Flowing temperature	-	22.35°C (72.23°F)

**Gas Volume Result**

AGA3 (1985)			AGA3 (1990)		
Factors	U/S Tap	D/S Tap	Factors	U/S Tap	D/S Tap
F <sub>b</sub>	2694.965	2694.97	C <sub>d</sub>	0.6047	0.6047
Y <sub>1</sub>	0.9998	0.9998	Y <sub>1</sub>	0.9998	0.9998
Y <sub>2</sub>	N/A	1.0001	Y <sub>2</sub>	N/A	1.0001
F <sub>tb</sub>	0.9981	0.9981	E <sub>v</sub>	1.0759	1.0759
F <sub>gr</sub>	1.3116	1.3116	Z <sub>b</sub>	0.9980	0.9980
F <sub>a</sub>	1.0001	1.0001	Z <sub>f</sub>	0.8425	0.8425
F <sub>r</sub>	1.0002	1.0002	Q	503.44	503.63 10 <sup>3</sup> m <sup>3</sup> /24 hr
F <sub>pb</sub>	1.0023	1.0023			
F <sub>tf</sub>	0.9884	0.9884			
F <sub>pv</sub>	1.0843	1.0843			
C'	3790.16	3790.31			
Q	501.64	501.82 10 <sup>3</sup> m <sup>3</sup> /24 hr			

**TEST CASE 5 (for AGA3 Flow Calculations)**

**Gas Analysis**

N <sub>2</sub>	-	0.0235	iC <sub>4</sub>	-	0.0088
CO <sub>2</sub>	-	0.0082	nC <sub>4</sub>	-	0.0169
H <sub>2</sub> S	-	0.0021	iC <sub>5</sub>	-	0.0035
C <sub>1</sub>	-	0.7358	nC <sub>5</sub>	-	0.0031
C <sub>2</sub>	-	0.1296	C <sub>6</sub>	-	0.0014
C <sub>3</sub>	-	0.0664	C <sub>7</sub>	-	0.0007

Ideal gas relative density - 0.7555

**Meter Data (flange taps)**

Meter run I.D.	-	154.05 mm (6.0650 inches)
Orifice I.D.	-	95.250 mm (3.750 inches)

**Flow Data (24 hr)**

Static pressure	-	2499.9 kPa(a) (362.58 psia)
Differential pressure	-	75.000 kPa (301.395 inches H <sub>2</sub> O)
Flowing temperature	-	34.0°C (93.2°F)

**Gas Volume Result**

**AGA3 (1985)**

<u>Factors</u>	<u>U/S Tap</u>	<u>D/S Tap</u>
F <sub>b</sub>	3111.24	3111.24
Y <sub>1</sub>	0.9894	0.9897
Y <sub>2</sub>	N/A	1.0044
F <sub>tb</sub>	0.9981	0.9981
F <sub>gr</sub>	1.1485	1.1485
F <sub>a</sub>	1.0005	1.0005
F <sub>r</sub>	1.0001	1.0001
F <sub>pb</sub>	1.0023	1.0023
F <sub>tf</sub>	0.9695	0.9695
F <sub>pv</sub>	1.0382	1.0394
C'	3561.90	3567.34
Q	800.22	813.37 10 <sup>3</sup> m <sup>3</sup> /24 hr

**AGA3 (1990)**

<u>Factors</u>	<u>U/S Tap</u>	<u>D/S Tap</u>
C <sub>d</sub>	0.6042	0.6041
Y <sub>1</sub>	0.9894	0.9897
Y <sub>2</sub>	N/A	1.0044
E <sub>v</sub>	1.0822	1.0822
Z <sub>b</sub>	0.9962	0.9962
Z <sub>f</sub>	0.9240	0.9217
Q	799.83	813.00 10 <sup>3</sup> m <sup>3</sup> /24 hr

**TEST CASE 6 (for AGA3 Flow Calculations)**

**Gas Analysis**

N <sub>2</sub>	-	0.0268	iC <sub>4</sub>	-	0.0123
CO <sub>2</sub>	-	0.0030	nC <sub>4</sub>	-	0.0274
H <sub>2</sub> S	-	0.0000	iC <sub>5</sub>	-	0.0000
C <sub>1</sub>	-	0.6668	nC <sub>5</sub>	-	0.0000
C <sub>2</sub>	-	0.1434	C <sub>6</sub>	-	0.0180
C <sub>3</sub>	-	0.1023	C <sub>7</sub>	-	0.0000

Ideal gas relative density - 0.8377

**Meter Data (flange taps)**

Meter run I.D.	-	52.500 mm (2.0669 inches)
Orifice I.D.	-	19.050 mm (0.750 inches)

**Flow Data (24 hr)**

Static pressure	-	2506.33 kPa(a) (363.50 psia)
Differential pressure	-	17.0500 kPa (68.5171 inches H <sub>2</sub> O)
Flowing temperature	-	7.2°C (44.96°F)

**Gas Volume Result**

AGA3 (1985)			AGA3 (1990)		
Factors	U/S Tap	D/S Tap	Factors	U/S Tap	D/S Tap
F <sub>b</sub>	115.138	115.138	C <sub>d</sub>	0.6005	0.6005
Y <sub>1</sub>	0.9978	0.9978	Y <sub>1</sub>	0.9978	0.9978
Y <sub>2</sub>	N/A	1.0012	Y <sub>2</sub>	N/A	1.0012
F <sub>tb</sub>	0.9981	0.9981	E <sub>v</sub>	1.0088	1.0088
F <sub>gr</sub>	1.0902	1.0902	Z <sub>b</sub>	0.9951	0.9951
F <sub>a</sub>	0.9996	0.9996	Z <sub>f</sub>	0.8588	0.8578
F <sub>r</sub>	1.0003	1.0003	Q	14.687	14.746 10 <sup>3</sup> m <sup>3</sup> /24 hr
F <sub>pb</sub>	1.0023	1.0023			
F <sub>tf</sub>	1.0148	1.0148			
F <sub>pv</sub>	1.0708	1.0714			
C'	136.15	136.22			
Q	14.602	14.660 10 <sup>3</sup> m <sup>3</sup> /24 hr			

**TEST CASE 7 (for AGA3 Flow Calculations)**

**Gas Analysis**

N <sub>2</sub>	-	0.0070	iC <sub>4</sub>	-	0.0062
CO <sub>2</sub>	-	0.0400	nC <sub>4</sub>	-	0.0090
H <sub>2</sub> S	-	0.0000	iC <sub>5</sub>	-	0.0052
C <sub>1</sub>	-	0.8720	nC <sub>5</sub>	-	0.0016
C <sub>2</sub>	-	0.0340	C <sub>6</sub>	-	0.0000
C <sub>3</sub>	-	0.0250	C <sub>7</sub>	-	0.0000

Ideal gas relative density - 0.6714

**Meter Data (flange taps)**

Meter run I.D.	-	52.500 mm (2.0669 inches)
Orifice I.D.	-	12.70 mm (0.50 inches)

**Flow Data (24 hr)**

Static pressure	-	299.92 kPa(a) (43.50 psia)
Differential pressure	-	6.3455 kPa (25.5 inches H <sub>2</sub> O)
Flowing temperature	-	1.67°C (35°F)

**Gas Volume Result**

AGA3 (1985)			AGA3 (1990)		
Factors	U/S Tap	D/S Tap	Factors	U/S Tap	D/S Tap
F <sub>b</sub>	50.523	50.523	C <sub>d</sub>	0.6006	0.6006
Y <sub>1</sub>	0.9933	0.9935	Y <sub>1</sub>	0.9933	0.9934
Y <sub>2</sub>	N/A	1.0039	Y <sub>2</sub>	N/A	1.0039
F <sub>tb</sub>	0.9981	0.9981	E <sub>v</sub>	1.0017	1.0017
F <sub>gr</sub>	1.2190	1.2190	Z <sub>b</sub>	0.9973	0.9973
F <sub>a</sub>	0.9994	0.9994	Z <sub>f</sub>	0.9905	0.9903
F <sub>r</sub>	1.0018	1.0018	Q	1.4335	1.4489 10 <sup>3</sup> m <sup>3</sup> /24 hr
F <sub>pb</sub>	1.0023	1.0023			
F <sub>tf</sub>	1.0250	1.0250			
F <sub>pv</sub>	1.0035	1.0036			
C'	63.013	63.029			
Q	1.4263	1.4416 10 <sup>3</sup> m <sup>3</sup> /24 hr			

**TEST CASE 8 (for AGA7 Flow Calculations)**

**Gas Analysis**

N <sub>2</sub>	-	0.0268	iC <sub>4</sub>	-	0.0123
CO <sub>2</sub>	-	0.0030	nC <sub>4</sub>	-	0.0274
H <sub>2</sub> S	-	0.0000	iC <sub>5</sub>	-	0.0000
C <sub>1</sub>	-	0.6668	nC <sub>5</sub>	-	0.0000
C <sub>2</sub>	-	0.1434	C <sub>6</sub>	-	0.0180
C <sub>3</sub>	-	0.1023	C <sub>7</sub>	-	0.0000

**Flow Data (24 hr)**

Uncorrected volume	-	128.0 10 <sup>3</sup> m <sup>3</sup>
Static pressure	-	2506.33 kPa(a) (363.50 psia)
Flowing temperature	-	7.2°C (44.96°F)

**Gas Volume Result**

**AGA7 (Volumetric Flow)**

**Factors**

F <sub>pm</sub>	24.6784
F <sub>pb</sub>	1.0023
F <sub>tm</sub>	1.0298
F <sub>tb</sub>	0.9981
Using AGA8 compressibility equations,	
S	1.1588
Q	3770.9 10 <sup>3</sup> m <sup>3</sup> /24 hr

Using RK compressibility equations,	
S	1.1467
Q	3731.6 10 <sup>3</sup> m <sup>3</sup> /24 hr

## 4.5 Measurement Reports for EFM Systems

The required information on each report must be stored using electronic media (not necessarily on the EFM) or printed media and can exist individually on different formats or reports and generated on demand for audit, as follows:

1. Daily for daily report required data
2. Monthly for monthly report required data
3. Event and alarm logs at regular intervals before information is overwritten
4. Meter reports generated on request for audit

### 4.5.1 Daily Report

The daily report must include:

1. Meter identification
2. Daily accumulated flow, with indicating flags for estimated flows made by the system or by the operation personnel and alarms that have occurred for over-ranging of end devices
3. Hours on production or hours of flow (specify)
4. Flow data audit trail – include at least one of the following:
  - a. Instantaneous values for flow rate, differential pressure (if applicable), static pressure, and temperature taken at the same time each day, or
  - b. Average daily values for differential pressure (if applicable), static pressure, and temperature, or
  - c. Hourly accumulated flow rate and average hourly values for differential pressure (if applicable), static pressure, and temperature

Existing EFM systems that do not have any of the audit trail capabilities specified in Section 4.5, and cannot develop the capability due to system limitations, should be evaluated for upgrading, especially when new production is tied into the facilities. ER may request upgrades, where audit/inspection results indicate they are warranted.

### 4.5.2 Monthly Report

This report is for the entire system, providing data for each measurement point. It is to contain the following at each measurement point as applicable:

1. Monthly cumulative flow
2. Flags indicating any change made to flow volumes
3. Total hours on production or hours of flow (specify)

### 4.5.3 Meter Report

The meter report details the configuration of each meter and flow calculation information. These values are used as part of the audit trail to confirm that the flow calculation is functioning correctly. Without them there is no way of verifying the accuracy of the system. The meter report must include the following as applicable and be produced on request by ER:

1. Instantaneous Flow Data, including:
  - a. Instantaneous flow rate
  - b. Instantaneous static pressure
  - c. Instantaneous differential pressure
  - d. Instantaneous flowing temperature
  - e. Instantaneous relative density (if live)
  - f. Instantaneous compressibility (if live)
  - g. Instantaneous gas component (if live)
  - h. Optional: instantaneous (AGA3) factors (see the orifice meter test cases in Section 4.4.2 for output information)
2. Current Configuration Information for Differential Meters or Other Types of Meters, whichever are applicable:
  - a. Meter identification
  - b. Date and time
  - c. Contract hour
  - d. Atmospheric pressure
  - e. Pressure base (unless fixed)
  - f. Temperature base (unless fixed)
  - g. Meter tube reference inside diameter
  - h. Orifice plate reference bore size
  - i. Static pressure tap location
  - j. Orifice plate material
  - k. Meter tube material
  - l. Calibrated static pressure range
  - m. Calibrated differential pressure range
  - n. Calibrated temperature range
  - o. High/low differential cutoff
  - p. Relative density (if not live)
  - q. Compressibility (if not live)
  - r. Gas components (if not live)

- s. Meter factor and/or K-Factor
- t. Effluent correction factor

#### 4.5.4 Event Log

This log is used to note and record exceptions and changes to the flow parameter, configuration, programming, and database affecting flow calculations, such as, but not limited to:

1. Orifice size change
2. Transmitter range change
3. Date of gas/liquid analysis update
4. Algorithm changes
5. Meter factor, K-Factor, or effluent correction factor changes
6. Other manual inputs

#### 4.5.5 Alarm Log

The alarm log includes any alarms that may have an effect on the measurement accuracy of the system. The time of each alarm condition and the time of clearing of each alarm must be recorded. Alarms to be reported must include, but are not limited to,

1. Master terminal unit failures
2. Remote terminal unit failures
3. Communication failures
4. Low-power warning
5. High differential pressure (for differential measurement devices)
6. High/low volumetric flow rate (for other types of measurement)
7. Over-ranging of end devices

## 4.6 References

American Gas Association Transmission Measurement Committee Report No. 8 (AGA8), November 1992. *Compressibility and Supercompressibility for Natural Gas and Other Hydrocarbon Gases*.

Dranchuk, P.M., and Abou-Kassam, J.H., "Calculation of Z Factors for Natural Gases Using Equations of State," *The Journal of Canadian Petroleum Technology*, Vol. 14, No. 3, July-September 1975, pp. 34-36.

Dranchuk, P.M., Purvis, R.A., and Robinson, D.B., "Computer Calculation of Natural Gas Compressibility Factors Using the Standing and Katz Correlation," *Institute of Petroleum Technical Series No. 1*, IP 74-008, 1974.

Gas Processors Association, *GPA 2145: "Table of Physical Constants for Hydrocarbons and Other Compounds of Interest to the Natural Gas Industry."*

Gas Processors Suppliers Association, *SI Engineering Data Book*.

Hall, K.R., and Yarborough, L., "A New Equation of State for Z Factor Calculations," *The Oil and Gas Journal*, June 18, 1973, pp. 82-92.

Pitzer, K.S., Lippman, D.Z., Curl, R.F., Huggins, C.M., and Petersen, D.E., "The Volumetric and Thermodynamic Properties of Fluids II. Compressibility Factor, Vapour Pressure and Entropy of Vapourization," *Journal of the American Chemical Society*, Vol. 77, No. 13, July 1955.

Redlich, O., and Kwong, J.N.S., "On the Thermodynamics of Solutions. V. An Equation of State. Fugacities of Gaseous Solutions," *Chemical Review* 44, 1949, pp. 233-244.

Wichert, E., and Aziz, K., "Calculate Z's for Sour Gases," *Hydrocarbon Processing*, Vol. 51, May 1972, pp. 119-122.

Yarborough, L., and Hall, K.R., "How to Solve Equation of State for Z-Factors," *The Oil and Gas Journal*, February 18, 1974, pp. 86-88

## 5 Site-Specific Deviation from Base Requirements

Section 1: Standards of Accuracy states that a licensee may deviate from ER's minimum measurement, accounting, and reporting requirements without specific approval if no royalty, equity, public safety, environmental, or reservoir engineering concerns are associated with the volumes being measured and the licensee is able to demonstrate that the alternative measurement equipment and/or procedures will provide measurement accuracy within the applicable uncertainties.

This section describes situations where a licensee may deviate from the minimum requirements without ER approval, provided that specific criteria are met. Licensees may also apply for approval to deviate from the minimum requirements if the specific criteria are not met; this section indicates what information must be included in such an application. If these exceptions or approvals are in use, ER inspectors and auditors will review the licensee's records for demonstrated compliance with the criteria specified in this section or in the applicable approval.

### 5.1 Site-specific Exceptions

Deviation from base measurement, accounting, and reporting requirements is allowed without submission of an application to ER, provided that all the qualifying criteria listed under the subsequent Exception sections are met.

**Qualifying Criteria** – These criteria detailed in the subsequent sections must be met to qualify for the exception. If the qualifying criteria have been met and the exception is implemented, it may remain in place indefinitely, as long as it does not meet any of the revocation clauses and no physical additions to the facility are made, e.g., new wells or stratigraphic units or zones. If additions or changes are made to the facility, the qualifying criteria must be met for all the wells or stratigraphic units or zones added to the facility for the exception to remain in place.

**Documentation Requirement** – The license must retain all the data and documentation to support the qualifying criteria and the last three testing records (if applicable) for as long as the exception is in effect. ER may revoke an exception when the licensee fails to produce the supporting data or documentation during an audit or inspection. The licensee will have thirty days to meet the applicable base requirements or at ER's discretion, the licensee can negotiate a plan to comply with the exception requirements within an ER-approved timeframe.

### 5.2 Site-specific Applications

If the exception criteria cannot be met or if a specific situation is not covered in this section, the licensee may be allowed to deviate from base measurement, accounting, and reporting requirements upon approval of an application submitted to ER.

A licensee may apply to deviate from base measurement, accounting and reporting requirements site-specific measurement exemption through the IRIS generic application process.

(see <http://www.saskatchewan.ca/business/agriculture-natural-resources-and-industry/oil-and-gas/oil-and-gas-licensing-operations-and-requirements/oil-and-gas-drilling-and-operations/measurement-requirements/apply-for-a-measurement-exemption> )

Site-specific approvals will remain in place indefinitely, unless otherwise stated in approval, including after transfer of the facility to another licensee provided that conditions specified in

the approval are met. If an ER audit or inspection finds that approval conditions are not being met, the approval may be revoked and the licensee may be required to meet applicable base requirements immediately, or other appropriate requirements may be specified.

If a licensee anticipates that proposed changes to the facility may not meet the approval conditions, the licensee may reconfigure the facility to meet base measurement, accounting, and reporting requirements or submit a new application for site-specific approval of deviation from the base requirements. Approval must be in place prior to implementation. Submission of an application does not guarantee that an approval will be granted.

The following information is required for all applications for site-specific deviation from base requirements. Other specific information that may be required is described in the applicable sections that follow.

1. Well and/or facility list, including:
  - a. Reporting-facility ID, facility licence (optional), facility subtype and surface location(s)
  - b. Well information including – well UWI (i.e. unique well identifier), well licence
  - c. Project ID, where applicable
  - d. Respective pool or stratigraphic unit or zone designations and unique identifier for each stratigraphic unit or zone
  - e. Indication as to unit or non-unit operation, if applicable
  - f. Mineral ownership type (i.e. Crown/Freehold)
  - g. Ownership and royalty equity issues, if any

If there are multiple owners each owner must be notified through the Public Notice found in *Directive PNG009: Public Notice Requirements*. With the submission of the application please provide a copy of the notice. If the owner does object then the licensee is required to notify the Ministry of Energy and Resources.
  - h. Up-to-date measurement schematic(s) for the existing system(s) and the proposed new gas or oil source(s), including all tie-in locations, if applicable
  - i. A detailed description of the new proposed operating procedure
  - j. A detailed description including the production accounting formulas for how the volumetrics will be determined
2. Justifications for deviation from measurement requirements, e.g., economics, minimal impact on measurement accuracy.

### 5.3 Chart Cycles Extended Beyond the Required Time Period

Chart cycle is the time required for a circular chart to complete one 360.0° revolution. An extension of the required chart cycle time may be applicable under the following scenarios:

1. The gas well orifice meter chart cycle is greater than eight days;
2. The single-well oil battery orifice gas meter chart cycle is greater than 24.0 hours; or

3. The Class 3 and 4 oil well test orifice gas meter chart cycle is greater than eight days.

Mixing of wells with EFM systems and wells using extended cycle paper charts within the same battery is allowed without approval from ER.

Group, sales, or delivery point meters and Class 1 and 2 oil well test gas meters do not qualify for exception from chart cycle requirements, and approvals for extension of the chart cycle for those meters will not normally be granted.

### 5.3.1 Exception

Orifice meter gas chart cycles may be extended without ER site-specific approval if all the qualifying criteria in Section 5.3.1.1 are met.

#### 5.3.1.1 Qualifying Criteria

Qualifying criteria that must be met includes the following:

1. For a gas multi-well battery, all wells in the battery are gas wells. A single-well battery does not qualify for this exception on its own; the entire group battery or gas gathering system must be considered.
2. For an oil battery, all wells in the battery are oil wells, and each well is linked to either an oil single-well battery or to an oil multi-well group battery, where each well has its own separation and measurement equipment.
3. All wells are subjected to the same type of measurement (all well production is separated and all separated streams are measured, or all well production is subject to effluent measurement) and the same chart cycle.
4. All wells flowing to the battery:
  - a. have common working interest ownership, and if there is no common ownership, written notification has been provided to all working interest participants and no objections have been received;
  - b. have either common Crown or common Freehold royalties. If there are no common Crown or common Freehold royalties and only Freehold royalties are involved, written notification has been given to all Freehold royalty owners. If the owner does object then the licensee is required to notify the Ministry of Energy and Resources.
5. The monthly average volumetric gas flow rate for each well is  $\leq 16.9 \times 10^3 \text{m}^3/\text{day}$  including the gas equivalent of condensate in the case of gas well measurement.
6. The differential pen records at  $\geq 33.0$  per cent within the chart range, and the static pressure pen should record at  $\geq 20.0$  per cent within the chart range. Painted traces must not exceed 4.0 per cent of the differential pressure or static pressure range. Painting occurs when there are quick up and down movements of the pen and there is no visible separation between the up and down traces for a period of time.
7. Temperature must be recorded at a minimum of once per week and if that is not possible, then continuous temperature measurement (temperature pen) is required.
8. The wells that are within the same battery as the extended chart cycle wells and are designed for and/or operate on on/off flows, e.g., plunger lifts, pump-off controls,

intermittent timers, must be measured using EFM. In addition, an extended chart cycle with EFM is allowed.

Exemption: wells producing gas at a rate  $\leq 3.0 \times 10^3 \text{m}^3/\text{day}$  do not have to meet qualifying criteria 6 to 8 to qualify for extended chart cycles; however, all other criteria must be met.

### 5.3.1.2 Revocation of Exception

If any of the following scenarios exists or occurs, the exception is revoked:

1. Gas from an oil battery is delivered to a gas battery.
2. There is mixed measurement within the battery other than with EFM.
3. The oil well is not linked to either an oil single-well battery or to an oil multi-well group battery where each well has its own separation and measurement equipment.
4. The working interest participants for any well flowing to the battery have changed and a new working interest participant objects to the exception.
5. Any well within the battery exceeds the  $16.9 \times 10^3 \text{m}^3/\text{day}$  monthly average gas production rate for three months including gas equivalent of condensate for gas wells.
6. Painted traces for any well exceeded 4.0 per cent of the differential pressure range or the static pressure range.
7. A new well with on/off flows is added to an effluent measurement battery or one or more of the existing wells has been modified to operate on on/off flows but EFM is not used.

Base measurement requirements must be reinstated if the exception is revoked due to any of the scenarios stated in Section 5.3.1.2.

### 5.3.2 Applications

The following information must be submitted with an application to extend orifice meter gas chart cycles if the criteria in Section 5.3.1.1 are not met:

1. All of the information listed in Section 5.2.
2. If there is no common ownership or no common Crown or common Freehold royalty, documentation to address royalty and equity issues demonstrating that written notification was given to all Freehold mineral owners and working interest participants, with no resulting objection received.
3. A discussion of the impact on measurement accuracy of intermingling base chart cycles and extended chart cycles in a common battery and how it may relate to concerns about working interest equity and/or royalty considerations.
4. A minimum of two current, consecutive, representative gas charts. Additionally, the licensee has the option to run the charts on the proposed chart cycle to gather test data for submission and then revert back to the required chart cycle after a maximum test period of 31 days. The original copies of any such charts created must be submitted with the application. The trial run must be clearly identified on the charts.

### 5.3.3 Considerations for Site-specific Approval

1. Differential and static pressures are stable, with essentially uninterrupted flow:
  - a. On/off flow as designed, including plunger lifts, pump-off controls, intermittent timers, etc., that cause painting or spiking, do not normally qualify for chart cycle extension.
  - b. The effects of painting are minimized. The amount of painting that is acceptable is decided case by case.
  - c. The differential pen should record at  $\geq 33.0$  per cent within the chart range and the static pressure pen should record at  $\geq 20.0$  per cent within the chart range.
2. There are minimal equity and royalty concerns.
3. Reservoir engineering concerns: the concern for well measurement accuracy declines, from a reservoir perspective, as the pool depletes. The applicant must provide its assessment/opinion, but ER has to decide on a case-by-case basis if the concerns are relevant.
4. All gas meters producing into the same group measurement point use the same chart cycle, so that they are subject to the same type of error.

### 5.4 Gas Proration Outside SW Saskatchewan Shallow Gas Stratigraphic Units

For wells outside the boundary of and/or producing from stratigraphic units other than those approved for the SW Saskatchewan Shallow Gas Stratigraphic Units (see Section 7.2), it may be acceptable to use a proration system for gas well production instead of having measurement for every well. If a proration system is implemented, all wells linked to the battery for reporting purposes must be subject to the proration system.

#### 5.4.1 Exception

Gas wells may be produced without individual well measurement and be physically connected to a proration battery without ER site-specific approval if all the qualifying criteria in Section 5.4.1.1 are met.

##### 5.4.1.1 Qualifying Criteria

1. All wells are classified as gas wells.
2. All wells flowing to the battery:
  - a. have common ownership, and where there is no common ownership, written notification has been provided to all working interest participants and no objections have been received;
  - b. have either common Crown or common Freehold royalties. If there are no common Crown or common Freehold royalties and only Freehold royalties are involved, written notification has been given to all Freehold royalty owners. If the owner does object then the licensee is required to notify ER.

3. The licensee has discussed and addressed reservoir engineering issues with its own reservoir engineering staff or external knowledgeable personnel to ensure minimal reservoir engineering concerns and has documented the results for audit.
4. Total liquid production at each well in the battery is  $\leq 2.0 \text{ m}^3/\text{day}$  based on the monthly average flow rates recorded during the six months prior to conversion. If a group of new wells not previously on production are to be constructed as a proration battery, the qualifying flow rates must be based on production tests conducted under the anticipated operating conditions of the proration battery.
5. The maximum average daily well gas flow rate of all wells in the battery is  $\leq 10.0 \cdot 10^3 \text{ m}^3/\text{day}$  including gas equivalent volume of condensate, with the highest daily well flow rate  $\leq 16.9 \cdot 10^3 \text{ m}^3$  including gas equivalent volume of condensate and except as allowed below in item 8. If an existing battery with measured gas well production is being converted to a proration battery, qualifying flow rates must be based on the monthly average flow rates recorded during the six consecutive months prior to conversion. If a group of new wells not previously on production are to be constructed as a proration battery, the qualifying flow rates must be based on production tests conducted under the anticipated operating conditions of the proration battery.
6. Periodic well tests are conducted under normal operating conditions to determine hourly flow rates that will be used to estimate monthly well production based on monthly well operating hours. The well tests are conducted for a minimum of 12.0 hours, and all gas, condensate, and water volumes are separated and measured during the test. For gas wells with minimal water production of  $\leq 0.01 \text{ m}^3 \text{ water}/10^3 \text{ m}^3 \text{ gas}$  and no condensate or oil, the testing duration must be sufficient to clearly establish stabilized flow rates and unseparated single-phase testing is allowed.
7. Following the commencement of production at the proration battery, all wells are tested within the first month, then again within six months, and then annually after that. New wells added to the battery at some future date must be tested within the first month of production, then again within six months, and then annually after that.
8. For new wells tying into a gas proration battery and that will be producing  $> 16.9 \cdot 10^3 \text{ m}^3/\text{day}$  but that are expected to drop to  $\leq 16.9 \cdot 10^3 \text{ m}^3/\text{day}$  within six months, every well must be tested monthly for the first six months with a separator or until the production rate has stabilized and annually thereafter. If the gas production rate for any of the wells is  $> 16.9 \cdot 10^3 \text{ m}^3/\text{day}$  after six months of production or the liquid production rate is  $> 2.0 \text{ m}^3/\text{day}$ , a separator must be installed to continuously separate and measure the well production, the well must be linked to a different battery reporting facility ID for reporting purposes, and the measurement-by-difference rules in Section 5.5 apply in this scenario.
9. The flow rates established from the well tests are used to determine estimated monthly well production from the date of the test until the date of the next test, except that the test conducted during the first month of production is also used to estimate the wells' production for the producing days prior to the test. The total measured group gas and liquid production are prorated to the wells, based on each well's estimated production, to determine the reported well production.

### 5.4.1.2 Revocation of Exception

If any of the following scenarios exists or occurs, the exception is revoked:

1. An oil well is added to the battery, or one or more of the existing gas wells has been reclassified as an oil well.
2. The maximum average daily flow rate of all wells in the battery for any month is  $> 10.0 \text{ } 10^3 \text{ m}^3/\text{day}$ , or the highest single well flow rate is  $> 16.9 \text{ } 10^3 \text{ m}^3/\text{day}$  except as allowed in Section 5.4.1.1, item 8.
3. Total liquid volume is  $> 2.0 \text{ m}^3/\text{day}$  during a 24.0 hour test period or prorated to 24.0 hours if the test period is not 24.0 hours.
4. A new well has been added to the proration battery with a daily flow rate  $> 16.9 \text{ } 10^3 \text{ m}^3$  except as allowed in Section 5.4.1.1 item 8 or whose additional volume will cause the average daily well gas flow rate of all wells in the battery to be  $> 10.0 \text{ } 10^3 \text{ m}^3/\text{day}$ .
5. Wells within the proration battery or new wells added to the battery were not tested as required.
6. The gas proration methodology in item 10 under Qualifying Criteria in Section 5.4.1.1 was not followed.

Base measurement requirements must be reinstated if the exception is revoked due to any of the scenarios listed above.

### 5.4.2 Applications

The following information must be submitted with an application to use a proration system, instead of individual gas well measurement, to determine gas well production if the criteria in Section 5.4.1.1 are not met:

1. All of the information listed in [Section 5.2](#);
2. A discussion of the stage of depletion for pools involved and the impact of any reduction in well measurement accuracy that may result from gas proration as it relates to reservoir engineering data needs - discussion of this matter by the licensee with its own reservoir engineering staff or knowledgeable external personnel is required and must be addressed in the application;
3. A clear explanation and flow diagram of proposed well and group measurement devices and locations, the proposed accounting and reporting procedures, and the proposed method and frequency of testing;
4. If there is no common ownership or no common Crown or common Freehold royalty, documentation to address royalty and equity issues demonstrating that written notification was given to all Freehold mineral owners and working interest participants, with no resulting objection received.

### 5.4.3 Considerations for Site-specific Approval

1. All wells must be classified as gas wells.
2. There are minimal equity, royalty, and reservoir engineering concerns.
3. All wells should have similar flow rates.

4. Economic considerations: Would implementation of a proration system reduce costs enough to significantly extend operations? Have other options been considered?
5. Total liquid production at each well in the battery should be  $\leq 2.0 \text{ m}^3/\text{day}$  based on the monthly average flow rates recorded during the six months prior to conversion. If a group of new wells not previously on production are to be constructed as a proration battery, the qualifying flow rates must be based on production tests conducted under the anticipated operating conditions of the proration battery.

## 5.5 Measurement by Difference

Measurement by difference (MbD) is defined as any situation where an unmeasured volume is determined by taking the difference between two or more measured volumes. It results in the unmeasured volume absorbing all the measurement uncertainty associated with the measured volumes. In the scenario of a proration battery, either effluent measurement or periodic testing without continuous measurement, new gas or oil source errors may be difficult to detect because the proration testing errors in the original system can hide the new source errors. Despite these concerns, a properly designed and operated measurement system can minimize the risk and attain reasonable accuracy, provided that the measured source gas or oil rates are a small proportion of the total system delivery rates. MbD is not allowed for multi-well group batteries, single-well batteries, or sales points unless a site-specific approval is obtained from ER prior to implementation.

### 5.5.1 Gas MbD

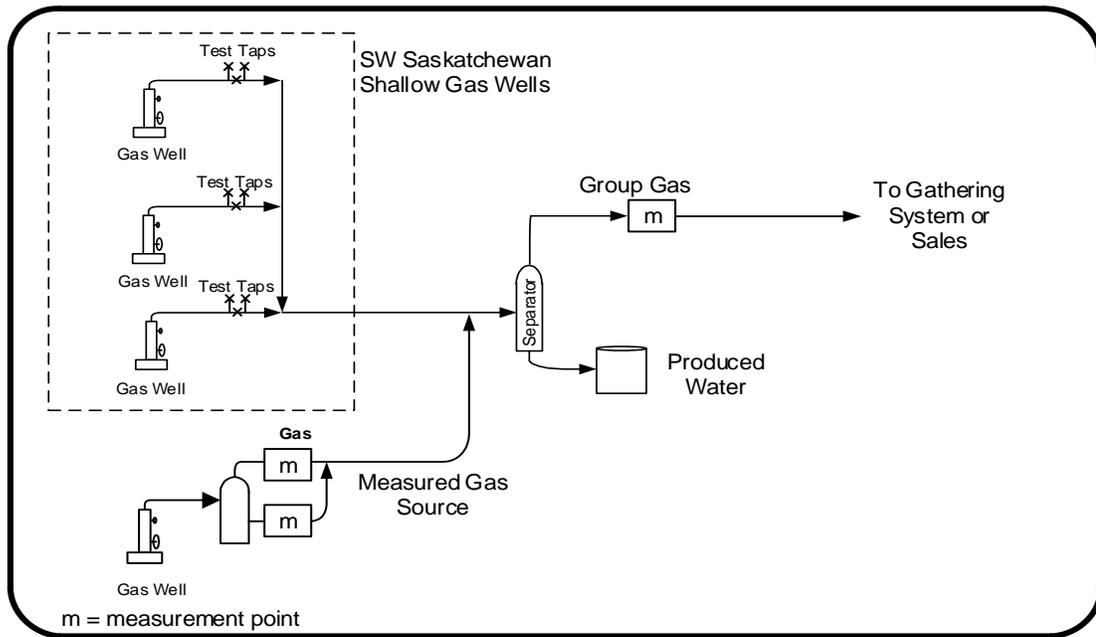
For proration batteries, MbD can include, but is not limited to, the following scenarios.

Note: All schematics are examples only; systems may be configured differently.

#### Scenario 1

Measured gas source(s) other than from the designated SW Saskatchewan Shallow Gas Stratigraphic Units delivering into a Gas Multi-well Proration SW Saskatchewan Battery ([Figure 5.1](#)):

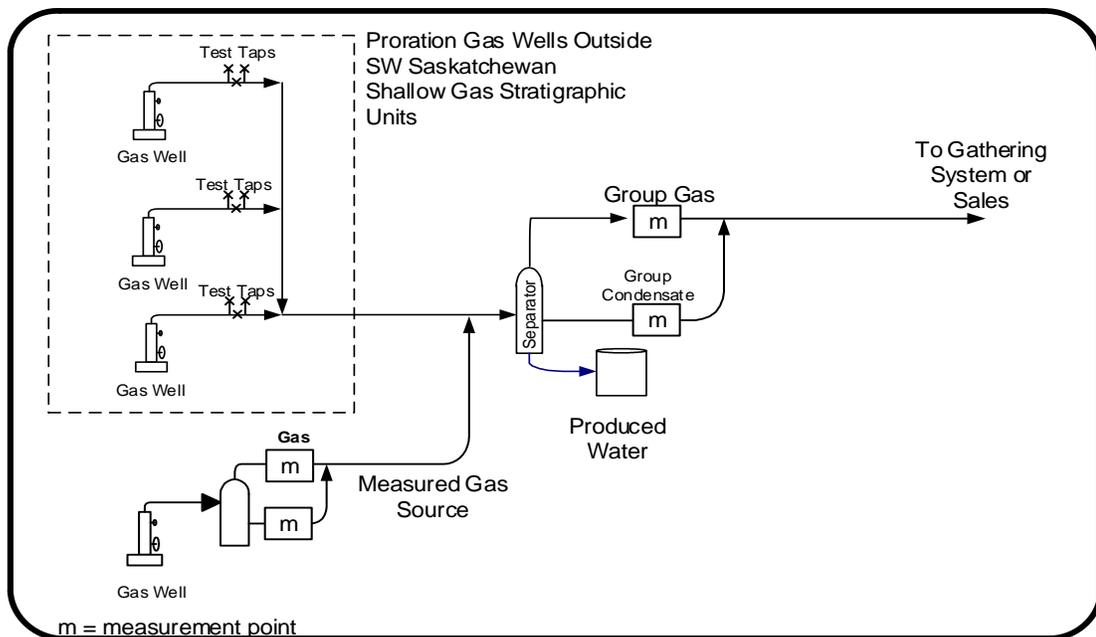
**Figure 5.1 Measured gas source delivering into a gas multi-well proration SW Saskatchewan battery**



**Scenario 2**

Measured gas source(s) delivering into a Gas Multi-well Proration Outside SW Saskatchewan Battery (Figure 5.2):

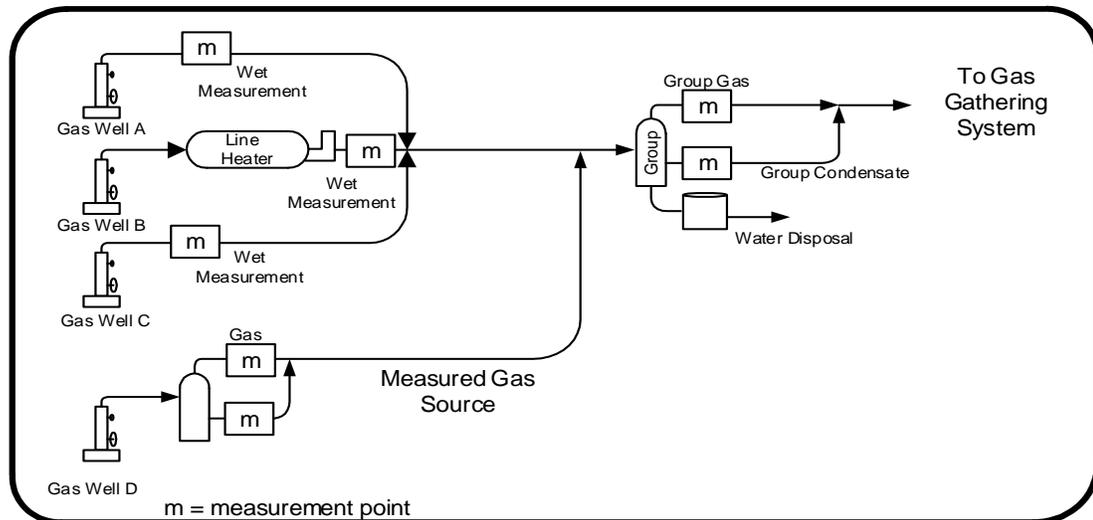
**Figure 5.2 Measured gas source delivering into a gas multi-well proration outside SW Saskatchewan battery**



### Scenario 3

Measured gas source(s) delivering into a gas multi-well effluent measurement battery with battery condensate separated, metered and recombined with battery gas (Figure 5.3):

**Figure 5.3 Measured gas sources delivering into a gas multi-well effluent measurement battery with battery condensate separated, metered, and recombined with battery gas**



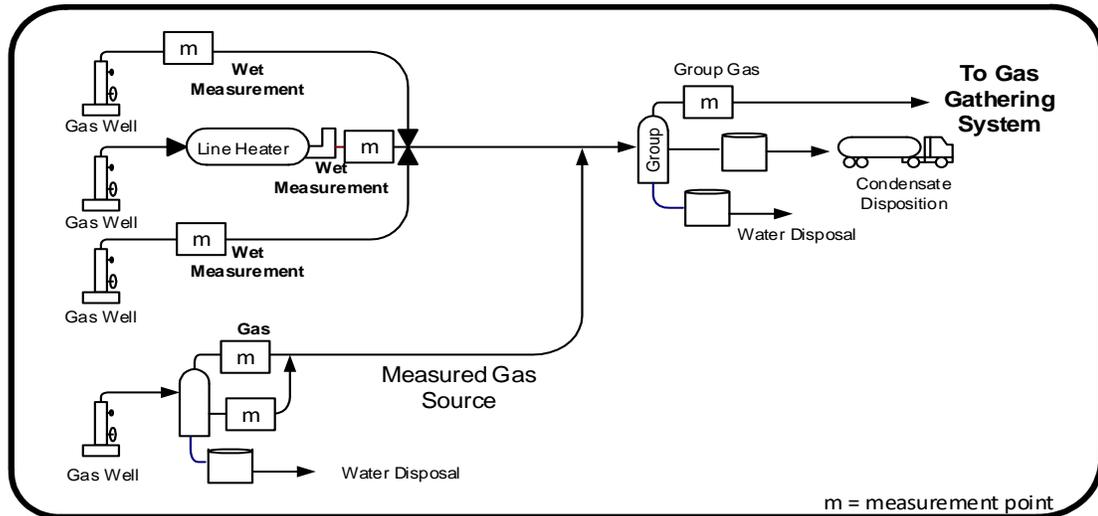
### Scenario 4

Measured gas source(s) delivering into a gas multi-well effluent measurement battery with battery condensate separated and sent to a tank for disposition to sales. (Figure 5.4): Note that this scenario can also occur at gas multi-well proration batteries outside SW Saskatchewan.

In this scenario, the condensate from the measured gas source is reported as a liquid condensate disposition to the effluent battery, rather than being included in the measured gas volume. If this reporting option is used, the following conditions must be adhered to:

1. MbD ratios and qualifying criteria for both gas and oil (condensate) are applicable at the effluent battery (see Section 5.5.3).
2. The condensate meter at the measured gas source must meet delivery point measurement requirements and be proven to base conditions.
3. A live condensate sample and analysis must be obtained at the measured gas source and used to conduct a flash simulation analysis to calculate a GIS at the measured gas source. The liquid condensate disposition from the measured gas source will be the metered condensate and the gas disposition will be the metered gas volume plus the calculated GIS.
4. The effluent battery condensate production will be the battery disposition minus the measured gas source condensate receipt plus change in inventory.

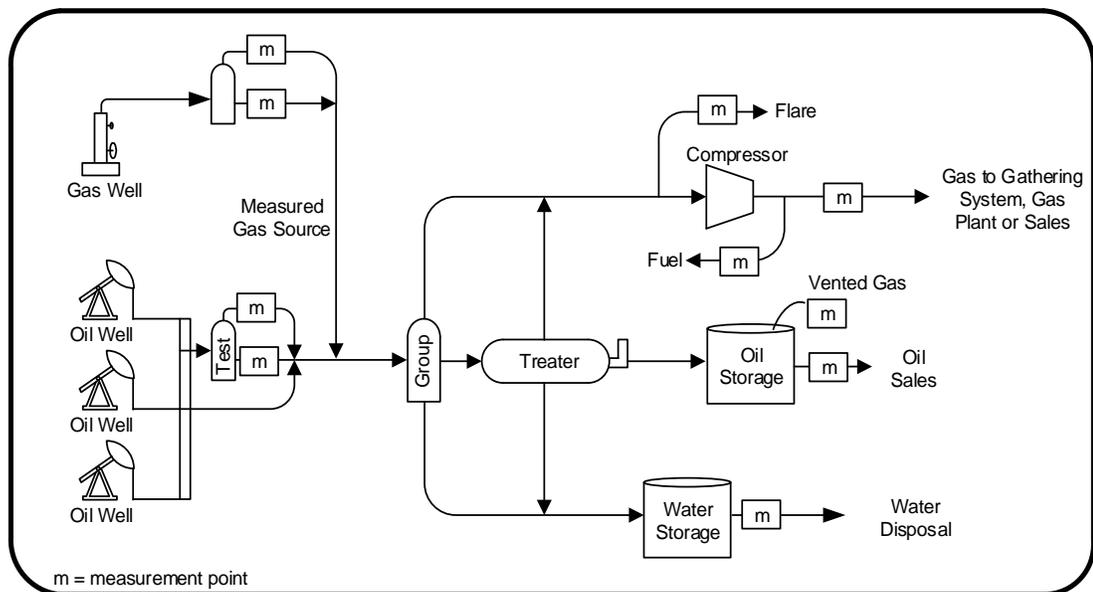
**Figure 5.4 Measured gas sources delivering into a gas multi-well effluent measurement battery with battery condensate separated and sent to a tank for disposition to sales**



**Scenario 5**

Measured gas source(s) delivering into an oil multi-well proration battery (Figure 5.5):

**Figure 5.5 Measured gas sources delivering into a crude oil multi-well proration battery**

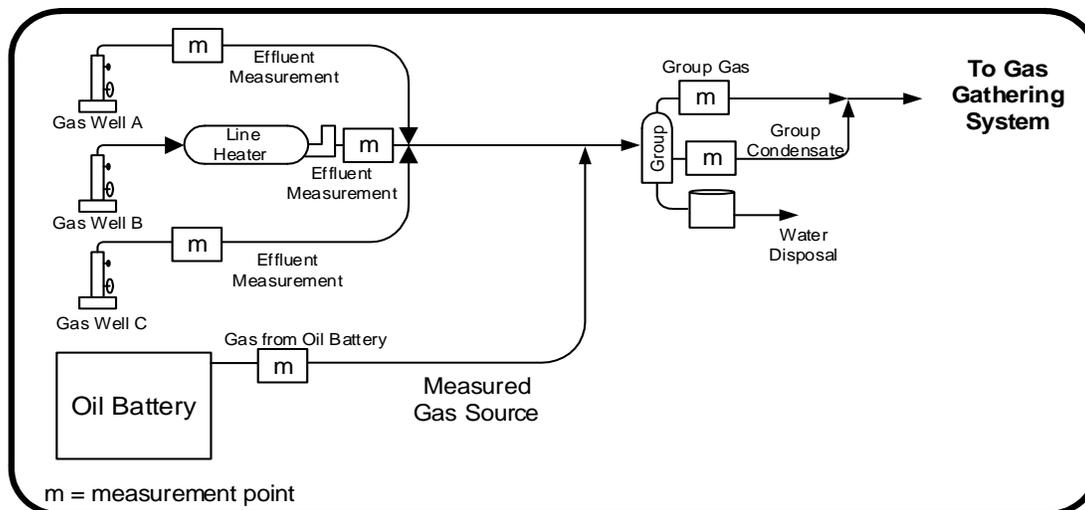


For the measured gas source(s), the applicable condensate metering and reporting option described in Table 5.6 in Section 5.5.5 must be used.

**Scenario 6**

Measured oil facility delivering gas into a gas multi-well proration battery (Figure 5.6):

**Figure 5.6 Measured oil facility gas delivering into a gas multi-well proration battery**



**5.5.1.1 Measured Gas Source into Gas Multi-well Proration Battery**

If any measured gas source will be tied into a gas multi-well proration battery:

1. The gas and liquids from all tied-in gas sources must be separately and continuously metered. If the R ratio in Table 5.1 cannot be met, the licensee may consider the tied-in measured gas wells as continuous or 31 day test and include them as part of the gas proration battery. However, these wells must be tagged as continuous test.
2. The monthly gas volume, including the GEV of condensate where applicable, received from a tied-in measured gas source and any other receipts must be subtracted from the total monthly gas battery disposition gas volume, including GEV of condensate where appropriate, to determine the gas battery monthly gas production volume.
3. Table 5.1 indicates when gas MbD may be acceptable by exception and when submission of an application may be required.

**Table 5.1 Gas MbD by exception and application**

Prorated gas flow rate (excluding all measured gas source)	R*	Application Required
$\leq 0.5 \times 10^3 \text{m}^3/\text{day}$	$< 1.00$	No
$> 0.5 \times 10^3 \text{m}^3/\text{day}$	$\leq 0.35$	No
$> 0.5 \times 10^3 \text{m}^3/\text{day}$	$> 0.35 \text{ and } \leq 0.75$	No**
$> 0.5 \times 10^3 \text{m}^3/\text{day}$	$> 0.75$	Yes

\* Ratio of volume of all tied-in measured gas volumes (including GEV of condensate where applicable) to the total battery gas disposition volume (including fuel, flare, vent volumes, and the GEV of condensate where applicable).

\*\* Must meet additional qualifying criteria, see Section 5.3.1.2

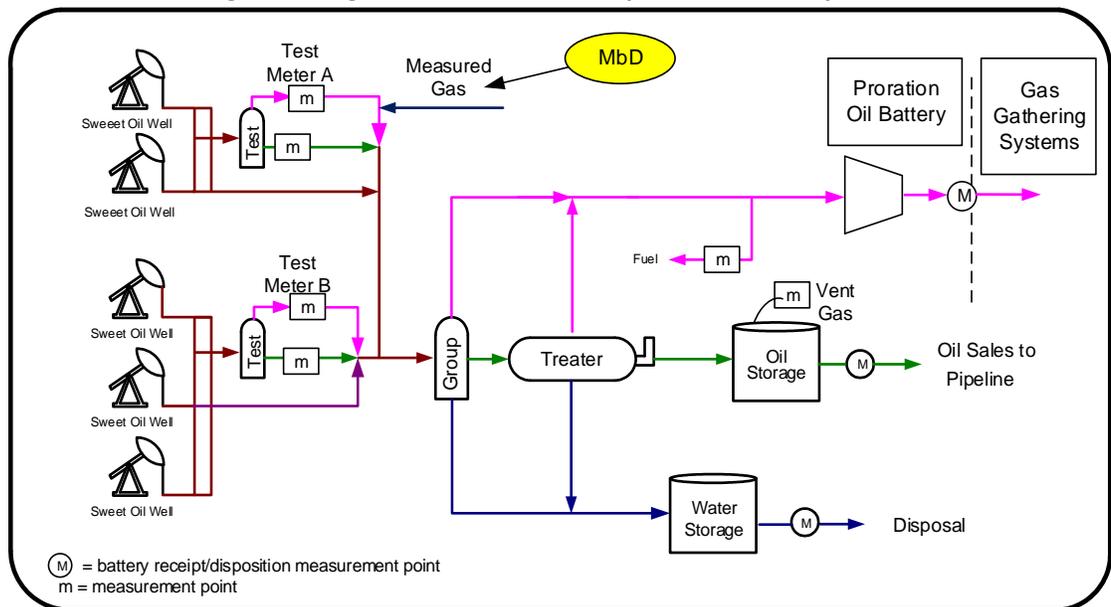
### 5.5.1.2 Measured Gas Source into an Oil Multi-well Proration Battery

If any measured gas source is tied in to an oil multi-well proration battery:

1. The gas and liquids from the tied-in gas source(s) must be separately and continuously metered.
2. The monthly gas volume (including, where applicable, the GEV of the portion of the condensate that will flash into the gas phase at the oil battery) received from a tied-in measured gas source and any other receipts must be subtracted from the total monthly oil battery gas disposition volume to determine the monthly oil battery gas production volume. See Table 5.6 for reporting options.
3. If condensate is received from a tied-in measured gas source, the portion of the monthly condensate volume that will remain in a liquid state at the oil battery must be subtracted from the oil battery total monthly oil disposition (plus/minus inventory changes and minus any other receipts) to determine the monthly oil battery oil production volume. See Table 5.6 for reporting options.

#### Scenario 1

Figure 5.7. Measured gas coming into an oil multi-well proration battery with MbD



#### To calculate reported oil battery gas production:

Total oil battery gas disposition to the gas gathering system is the metered volume after compression, see Figure 5.7. The oil battery gas production is calculated by subtracting the measured gas receipt volumes from the sum total of the oil battery disposition to the gas gathering system, fuel, flare, and vent. The resultant oil battery gas production volume is then prorated to the flowlined oil wells. The amount of measured gas that can be delivered into the oil battery is limited by the MbD ratio in Section 5.5.



**Table 5.2 Gas MbD by exception and application**

Prorated gas flow rate (excluding all measured gas source)	R*	Application Required
$\leq 0.5 \text{ } 10^3\text{m}^3/\text{d}$	$< 1.00$	No
$> 0.5 \text{ } 10^3\text{m}^3/\text{d}$	$\leq 0.35$	No
$> 0.5 \text{ } 10^3\text{m}^3/\text{d}$	$> 0.35 \text{ and } \leq 0.75$	No**
$> 0.5 \text{ } 10^3\text{m}^3/\text{d}$	$> 0.75$	Yes

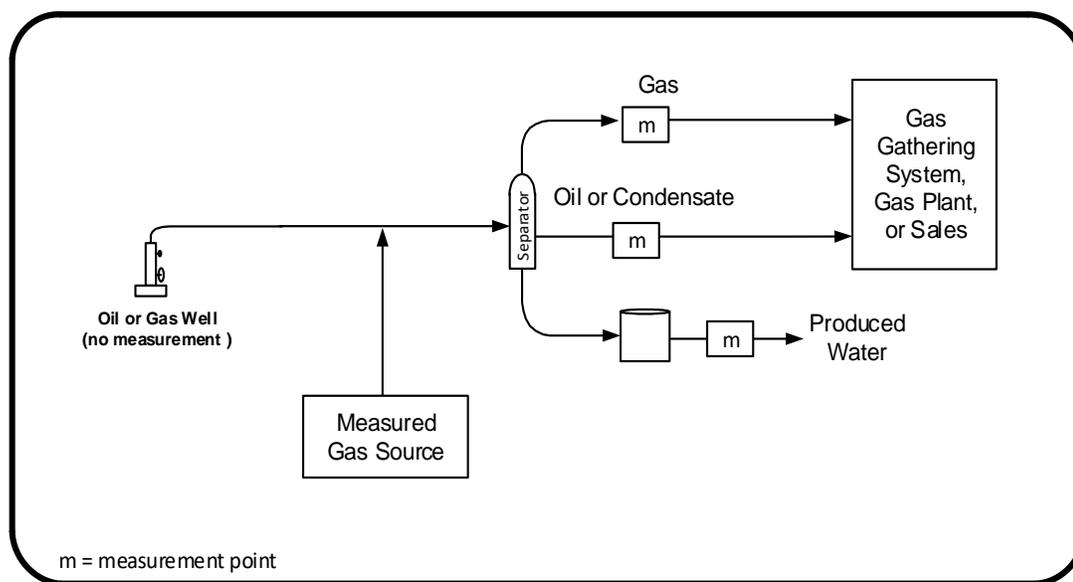
\* Ratio of volume of all tied-in measured gas volumes (including GEV of condensate where applicable) to the total battery gas disposition volume (including fuel, flare, vent volumes, and including GEV of condensate where applicable).

\*\* Must meet additional qualifying criteria, see Section 5.5.3.1.2.

### 5.5.1.3 Measured Gas Source into a Single-Well Battery

Where a measured gas source will be tied into a single-well battery, as shown in [Figure 5.9](#), this situation does not qualify for an exception, and an application must be submitted to and approved by ER prior to implementation.

**Figure 5.9 Measured gas source delivering into a single-well battery**



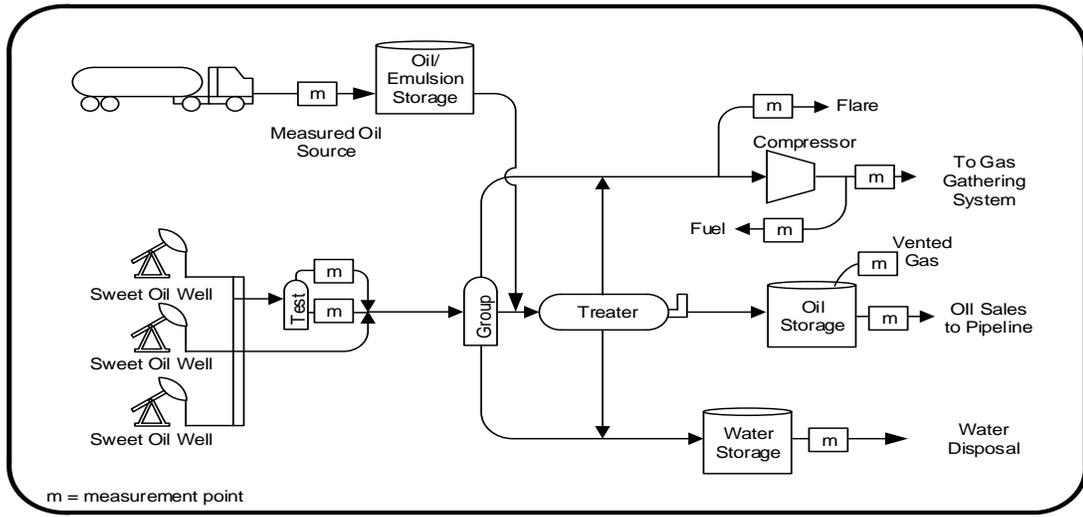
### 5.5.2 Oil MbD

For oil streams, MbD can include but is not limited to the following scenarios.

#### Scenario 1

Measured oil and/or oil-water emulsion from a battery delivering into an oil multi-well proration battery by truck ([Figure 5.10](#)):

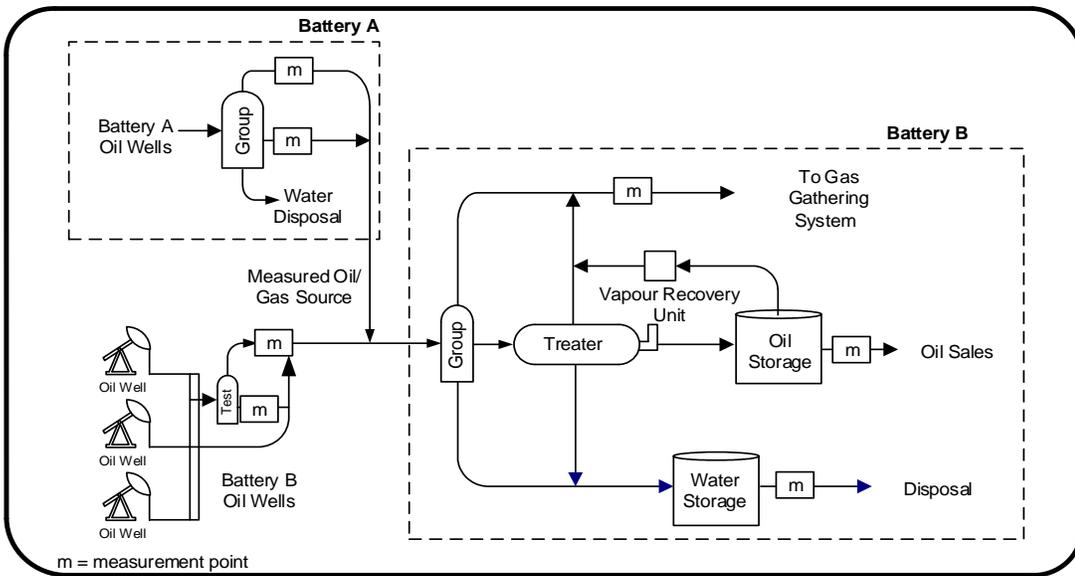
**Figure 5.10 Measured oil or oil emulsion from a battery delivering into an oil multi-well proration battery by truck**



**Scenario 2**

Measured oil and/or oil-water emulsion (and gas if applicable) under pressure from a battery delivering into an oil proration battery by pipeline (Figure 5.11):

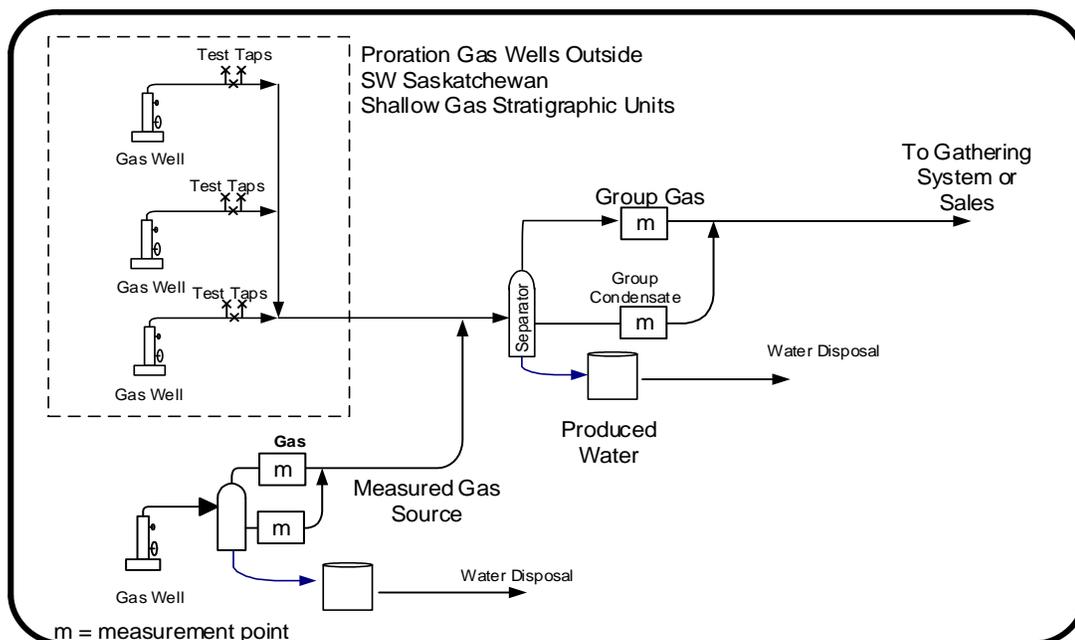
**Figure 5.11 Measured oil or oil emulsion (and gas if applicable) under pressure from a battery delivering into an oil multi-well proration battery by pipeline**



**Scenario 3**

Measured oil and/or oil-water emulsion from a measured gas source delivering into a gas multi-well proration battery or gas plant (Figure 5.12): For specific measurement and reporting information, see Section 5.5.3.1.1 #8.

**Figure 5.12 Measured oil or oil emulsion from a measured gas source delivering into a gas multi-well proration battery or gas plant**



**5.5.2.1 Measured Oil or Oil Emulsion Source into an Oil Multi-well Proration Battery**

If any measured oil and/or oil-water emulsion source is-delivered to an oil proration battery including trucked-in volumes:

1. Measured oil and/or oil-water emulsion delivery/receipt volumes must be determined using equipment and/or procedures that meet delivery point measurement uncertainty requirements. In the scenario of oil-water emulsions, the measurement uncertainty requirements apply to total volume determination only.
2. Measured oil volumes must be determined and reported at standard conditions.
3. The liquids received from the measured oil and/or oil-water emulsion source(s) must be subtracted from the total monthly oil battery oil and water disposition volumes plus/minus inventory changes and minus any other receipts to determine the monthly oil battery oil and water production volumes.
4. [Table 5.3](#) indicates when oil MbD is acceptable by exception and when submission of an application is required.

**Table 5.3 Oil MbD by exception and application**

Measured oil receipt volume	R*	Application Required
≤ 1000.0 m <sup>3</sup> /month	Not applicable	No
> 1000.0 m <sup>3</sup> /month	≤ 0.25	No
> 1000.0 m <sup>3</sup> /month	0.25 < R ≤ 1.00	No**
> 1000.0 m <sup>3</sup> /month	> 1.00	Yes

\* Total measured oil receipt volume divided by the monthly battery oil production

\*\* Must meet additional qualifying criteria, see Section 5.5.3.2.1

5. Consideration should be given to incorporating pipeline measured oil and/or oil-water emulsion source(s) delivered by pipeline as a satellite of the receiving battery, if the battery delivering the oil and/or oil-water emulsion is an oil proration battery, and including it in the receiving battery’s proration system. In that scenario, MbD would be avoided. A pipelined single oil well or oil wells in a multi-well group may also be considered as continuous or 31 day test and included as part of the oil proration battery. However, wells must be tagged as continuous test.

### 5.5.3 Exceptions

MbD is allowed without ER site-specific approval if all of the applicable criteria in this section are met.

If the MbD will involve existing production, initial qualifying flow rates must be based on average calendar daily flow rates (monthly flow rate divided by number of production hours in the month multiplied by 24) recorded during the six months prior to implementation of the MbD. If new measured production is to be connected to a proration battery, the qualifying flow rates must be based on production tests conducted under the anticipated operating conditions.

#### 5.5.3.1 Exceptions for All Measured Gas Streams

For measured gas source(s) from either gas or oil batteries tied into a gas proration battery or an oil proration battery.

##### 5.5.3.1.1 Qualifying Criteria

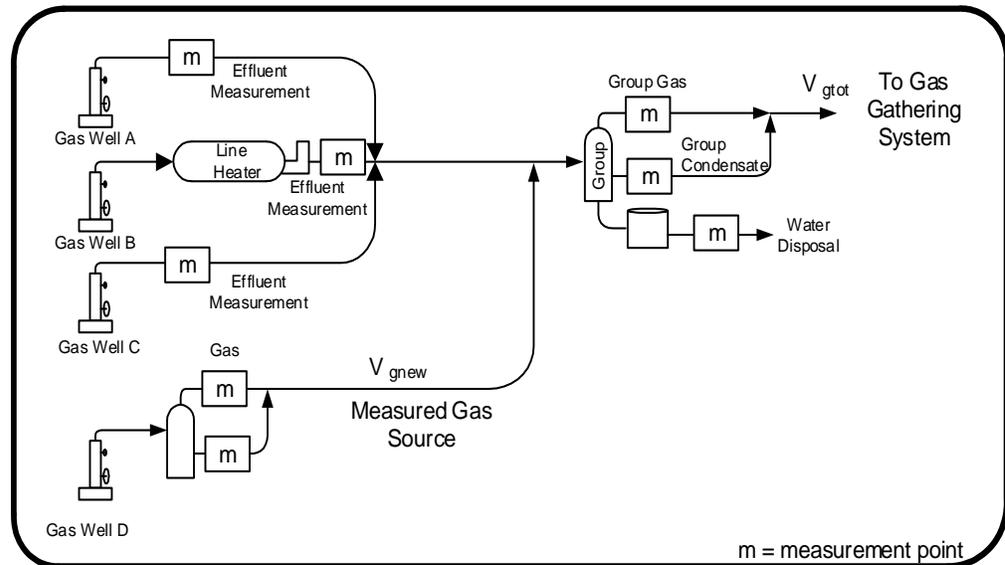
1. Volumetric criteria for measured gas tying into a proration battery

**Table 5.4 Volumetric criteria for measured gas tying into a multi-well proration battery**

Prorated gas flow rate (excluding all measured gas source)	R*
$\leq 0.5 \times 10^3 \text{m}^3/\text{day}$	$< 1.00$
$> 0.5 \times 10^3 \text{m}^3/\text{day}$	$\leq 0.35$
$> 0.5 \times 10^3 \text{m}^3/\text{day}$	$0.35 < R \leq 0.75^{**}$

\*R: Ratio of volume of all tied-in measured gas (including GEV of condensate where applicable) to the total gas disposition volume from the receiving battery (including fuel, flare, vent volumes, and GEV of condensate where applicable).

\*\* Additional qualifying criteria apply, see Section 5.5.3.1.2

**Example:****Figure 5.13 Gas battery example of volumetric criteria for measured gas tying into a multi-well proration battery**

For the gas battery in [Figure 5.13](#),

$V_{gtot} = 100.0 \text{ } 10^3\text{m}^3/\text{day}$  (total of measured gas and GEV of condensate delivered out of the proration battery, including volumes received from Gas Well D)

$V_{gnew} = 30.0 \text{ } 10^3\text{m}^3/\text{day}$  (total of measured gas and GEV of condensate delivered to the proration battery from Gas Well D)

Prorated gas flow rate =  $V_{gtot} - V_{gnew} = 100.0 - 30.0 = 70.0 \text{ } 10^3\text{m}^3/\text{day}$

$R = 30.0/100.0 = 0.3$

Since the prorated flow rate is  $> 0.5 \text{ } 10^3\text{m}^3/\text{day}$  and  $R$  is below 0.35 for the Gas Well D tie-in, it is within the acceptable exception range.

2. All proration wells flowing to the battery:
  - a. have common working interest ownership, and, where there is no common ownership written notification has been provided to all working interest participants. If the owner does object then the licensee is required to notify the Ministry of Energy and Resources;
  - b. have either common Crown or common Freehold royalties. If there are no common Crown or common Freehold royalties and only Freehold royalties are involved, written notification has been given to all Freehold royalty owners. If the owner does object then the licensee is required to notify the Ministry of Energy and Resources.
3. The gas and liquid phases from the tied-in measured gas source(s) are separately and continuously metered.

4. Gas volumes received at a gas proration battery from the tied-in measured gas source(s) include the GEV of the measured condensate volumes if the condensate is recombined with the measured gas volumes from the new tied-in gas source.
5. If the tied-in measured gas source(s) produces condensate and is connected by pipeline to an oil proration battery, the applicable condensate metering and delivery/reporting options described in Table 5.6 in Section 5.5.5 must be used.
6. In the scenario of an oil proration battery or a gas proration battery, the monthly gas volume, including the GEV of condensate where applicable, received from a tied-in measured gas source and any other receipts, is subtracted from the total monthly proration battery gas volume, including the GEV of condensate where applicable, to determine the monthly proration battery gas production volume.
7. In the scenario of an oil proration battery, the monthly liquid condensate volume, where appropriate, received from a tied-in measured gas source, is subtracted from the total monthly oil disposition, plus inventory changes, shrinkage, if applicable, and minus any other receipts, to determine the monthly oil proration battery oil production volume.
8. Oil and/or oil-water emulsion from a tied-in measured gas source may be delivered to a gas proration battery, or gas plant in accordance with the following:
  - a. The oil or oil-water emulsion must be measured with a meter proved to stock tank conditions
  - b. A live oil sample must be taken annually and a multiphase flash liberation or computer simulation must be performed in order to determine the GIS factor of the entrained gas in the oil which must be added to the measured gas volume.
  - c. The oil or oil-water emulsion disposition must be reported as a liquid oil volume and kept whole, as it is reported through the gathering system and gas plant inlet.
  - d. Blending shrinkage requirements in Section 14.3.2 must be adhered to.
  - e. The oil and gas MbD exception qualifying criteria set out in Section 5.5.3 must be adhered to.

**5.5.3.1.2 Additional Qualifying Criteria:  $0.35 < R \leq 0.75$**

1. Single point measurement uncertainty of the measured gas source gas meter and of the prorated battery group gas meter must be  $\leq 2.0$  per cent.
2. EFM must be installed on both the gas and condensate meters at the measured gas source meter(s) and the proration battery group separator.
3. Gas proration factor targets, as set out in Table 3.1 must be maintained.
4. Potential reservoir engineering/management concerns have been considered and determined to be acceptable.

**5.5.3.1.3 Revocation of Exceptions:  $R \leq 0.35$**

If any of the qualifying criteria specified in Section 5.5.3.1.1 is not adhered to, then the exception is revoked and base requirements must be reinstated.

**5.5.3.1.4 Revocation of Exceptions:  $0.35 < R \leq 0.75$**

If any of the qualifying criteria in Section 5.5.3.1.1 or the additional qualifying criteria in Section 5.5.3.1.2 are not adhered to, then the exception is revoked.

However, if the gas proration factor at the proration battery exceeds the proration factor targets as set out in Section 3.1.1, then the licensee must take steps to bring the proration factor back within range within two months after the initial month the proration targets were exceeded. If the gas proration factor cannot be restored to within the target range within two months, the exception is revoked and the licensee must restore the R factor to 0.35 or lower or obtain a site-specific approval.

Base measurement requirements must be reinstated if the exception is revoked.

**5.5.3.2 Exception for Measured Oil Receipts Received by Truck or Pipeline at an Oil Proration Battery**

**5.5.3.2.1 Qualifying Criteria**

1. Volumetric Criteria for Measured Oil Delivered by Truck or Pipeline to an Oil Proration Battery:

**Table 5.5 Volumetric criteria for measured oil delivered to an oil multi-well proration battery**

Measured oil receipt volume	R*
$\leq 1000.0 \text{ m}^3/\text{month}$	Not applicable
$> 1000.0 \text{ m}^3/\text{month}$	$\leq 0.25$
$> 1000.0 \text{ m}^3/\text{month}$	$0.25 < R \leq 1.00^{**}$

\* Total measured oil receipt volume divided by the monthly battery oil production

\*\* Additional qualifying criteria apply, see Section 5.5.3.2.2

2. The monthly oil battery oil and water production volumes are determined by subtracting the monthly measured oil and water receipt volumes from the total monthly oil battery oil and water disposition volumes plus inventory change and minus any other receipts.
3. All wells linked to the oil proration battery (the proration wells):
  - a. have common working interest ownership, and, where there is no common ownership written notification has been provided to all working interest participants. If the owner does object then the licensee is required to notify the Ministry of Energy and Resources;
  - b. have either common Crown or common Freehold royalties. If there are no common Crown or common Freehold royalties and only Freehold royalties are involved, written notification has been given to all Freehold royalty owners., with no resulting objections received. If the owner does object then the licensee is required to notify the Ministry of Energy and Resources.

4. If measured gas from a measured live oil/emulsion production source is also commingled with the production at an oil proration battery (pipelined receipt), the exception criteria for gas MbD must also be met.

#### **5.5.3.2.2 Additional Qualifying Criteria: $0.25 < R \leq 1.00$**

1. Delivery point measurement must be installed at the oil proration battery to meter the measured oil receipts (trucked in or pipelined) and the delivery point measurement uncertainty is  $\leq 0.5$  per cent, irrespective of the daily volume of the metered receipts.
2. Oil (and gas if applicable) proration factor targets, as set out in Table 3.1 must be maintained.
3. Proving requirements and frequency for the delivery point measurement devices must be adhered to.
4. Blending requirements in Section 14.3.2 must be adhered to.
5. Potential reservoir engineering/management risks have been considered and determined to be acceptable.

#### **5.5.3.2.3 Revocation of Exceptions: $R \leq 0.25$**

If any of the qualifying criteria specified in Section 5.5.3.2.1 are not adhered to then the exception is revoked and base measurement requirements must be reinstated.

If the exception is revoked, the licensee must:

1. Deliver all oil receipts  $> 1000.0 \text{ m}^3/\text{month}$  elsewhere;
2. Set up another treater train with separate receipt measurement, tankage, and disposition measurement to process the trucked in or pipelined receipts prior to commingling with the battery production; or
3. Obtain ER site-specific special approval to continue.

#### **5.5.3.2.4 Revocation of Exceptions: $0.25 < R \leq 1.00$**

If any of the qualifying criteria in Section 5.5.3.2.1 or the additional qualifying criteria in Section 5.5.3.2.2 are not adhered to, then the exception is revoked.

However, if the oil (and gas if applicable) proration factor(s) at the proration battery exceeds the proration factor targets as set out in Table 3.1, then the licensee must take steps to bring the proration factors back within range within two months after the initial month the proration targets were exceeded. If the proration factors cannot be restored to within the target range within two months, the exception is revoked and the licensee must restore the R factors to 0.25 or lower for oil and 0.35 or lower for gas or obtain a site-specific approval to continue.

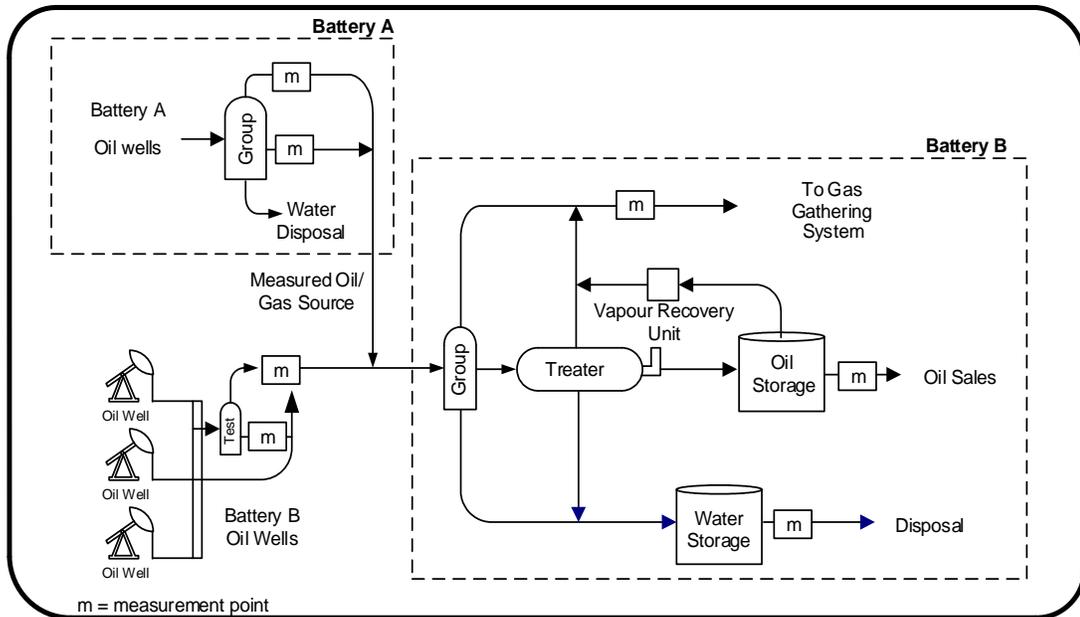
Base measurement requirements must be reinstated if the exception is revoked.

If the exception is revoked the licensee must:

1. Deliver all oil receipts  $> 1000.0 \text{ m}^3/\text{month}$  elsewhere;
2. Set up a treater train with separate receipt measurement, tankage, and disposition measurement to process the trucked in or pipelined receipts prior to commingling with the battery production; or

3. Obtain ER site-specific approval to continue.

**Figure 5.14 Oil battery example of volumetric criteria for measured oil tying into a multi-well proration battery**



Note that with the addition of Battery A production, if the MbD meets all the qualifying criteria and the total oil receipt volume at oil proration Battery B is  $> 100.0 \text{ m}^3/\text{day}$ , the delivery volume must be determined by a measurement device(s) and/or procedures having  $\pm 0.5$  per cent uncertainty, which might require changes in measurement equipment and/or procedures at Battery B.

For this example (Figure 5.14), given the following data:

Battery A oil production volume =  $20.0 \text{ m}^3/\text{d}$

Battery B oil production volume =  $90.0 \text{ m}^3/\text{d}$  before tying in Battery A

Battery A gas production volume =  $15.0 \cdot 10^3 \text{ m}^3/\text{d}$

Battery B gas production volume =  $20.0 \cdot 10^3 \text{ m}^3/\text{d}$  before tying in Battery A

**Step 1:** Calculate the monthly measured oil volume from Battery A delivered to the proration battery (Battery B) and the ratio of the prorated oil production:

Monthly measured oil production volume from Battery A =  $20.0 \text{ m}^3/\text{d} \times 30 \text{ days} = 600.0 \text{ m}^3$

Battery A oil volume as a ratio (R) of Battery B oil production volume =  $20.0 \text{ m}^3/\text{day} / 90.0 \text{ m}^3/\text{day} = 0.22$

**Step 2:** Calculate the R ratio for the commingled gas:

$$R = 15.0 / (15.0 + 20.0) = 0.43$$

Since the oil proration Battery A monthly measured oil volume is below  $1000.0 \text{ m}^3/\text{month}$ , the oil volumetric criteria are met. The gas R ratio is also below 0.75 so an application is not required in this case, provided all prequalifying criteria are met.

#### 5.5.4 Applications

The following information must be submitted with an application for a proration battery to receive measured gas or oil/emulsion sources if the applicable qualifying criteria and additional qualifying criteria in Section 5.5.3.2.2 are not met:

1. All of the information listed in Section 5.2;
2. A discussion of the stage of depletion for pools involved, and the impact of any reduction in well measurement accuracy that may result from Mbd as it relates to reservoir engineering data needs; discussion of this matter by the licensee with its own reservoir engineering staff or knowledgeable external personnel is required and must be addressed in the application;
3. If there is no common ownership or no common Crown or common Freehold royalty, documentation to address royalty and equity issues demonstrating that written notification was given to all Freehold mineral owners and working interest participants, with no resulting objection received.

#### 5.5.5 Considerations for Site-specific Approval

1. There are minimal equity, royalty, and reservoir engineering concerns.
2. Economic considerations, including an assessment of whether implementation of a proration system would reduce costs enough to significantly extend operations, and an assessment of the other options that have been considered.
3. The gas and liquids from the tied-in measured source(s) must be separately and continuously measured.
4. If the tied-in measured gas source(s) produces condensate and it is connected by pipeline to an oil battery, the licensee must choose the applicable condensate delivery/reporting options from [Table 5.6](#):

**Table 5.6 Options for condensate delivery and reporting into an oil multi-well proration battery**

<b>Condensate received at oil battery (from all measured gas sources)</b>	<b>Condensate reporting options</b>
<b><math>\leq 2.0 \text{ m}^3/\text{day}</math> and <math>\leq 5.0\%</math> of total prorated oil production</b>	<ol style="list-style-type: none"> <li>1. Prove the tied in measured gas source condensate meter to live conditions.</li> <li>2. Obtain a live condensate liquid sample and send the sample to a lab for a liquid analysis (to C<sub>30+</sub>).</li> <li>3. Multiply the monthly metered condensate volume by the liquid volume fraction from the analysis to derive the component volumes.</li> <li>4. Report the C<sub>6+</sub> (Hexane plus) as a liquid condensate disposition from the measured gas source to the oil battery.</li> <li>5. Most of the light ends (H<sub>2</sub> to NC<sub>5</sub>) will flash out of the liquid condensate at the oil battery treater. Add the light ends (H<sub>2</sub> to NC<sub>5</sub>) component gas equivalent volumes to the dry flow measured gas component volumes and report this as the total gas disposition from the measured gas source to the oil battery.</li> </ol>
<b><math>&gt; 2.0 \text{ m}^3/\text{day}</math> or <math>&gt; 5.0\%</math> of total prorated oil production</b>	<ol style="list-style-type: none"> <li>1. Prove the tied in measured gas source condensate meter to live conditions.</li> <li>2. Obtain a live condensate liquid sample (to C<sub>30+</sub>) and perform a computer flash simulation to determine how much gas will flash out of the condensate at each production stage, (i.e. separator and treater) at the oil battery. This will allow for a shrinkage factor to be determined.</li> <li>3. Report the condensate stock tank volume derived from the metered condensate volume and the simulation shrinkage factor as a liquid disposition from the measured gas source to the oil battery.</li> <li>4. The flash simulation will also derive the volume and composition of the light ends that will flash out of the condensate at each production stage within the battery. Add the light end (flashed) condensate component gas volumes to the dry flow measured gas component volumes and report this as the total gas disposition from the measured gas source to the oil battery.</li> <li>5. If there are changes to the process (temperature, pressure) at either the measured gas source or oil battery, or if the measured gas source has new richer or leaner wells tied in, a new condensate sample must be obtained and a new computer flash simulation conducted.</li> </ol>

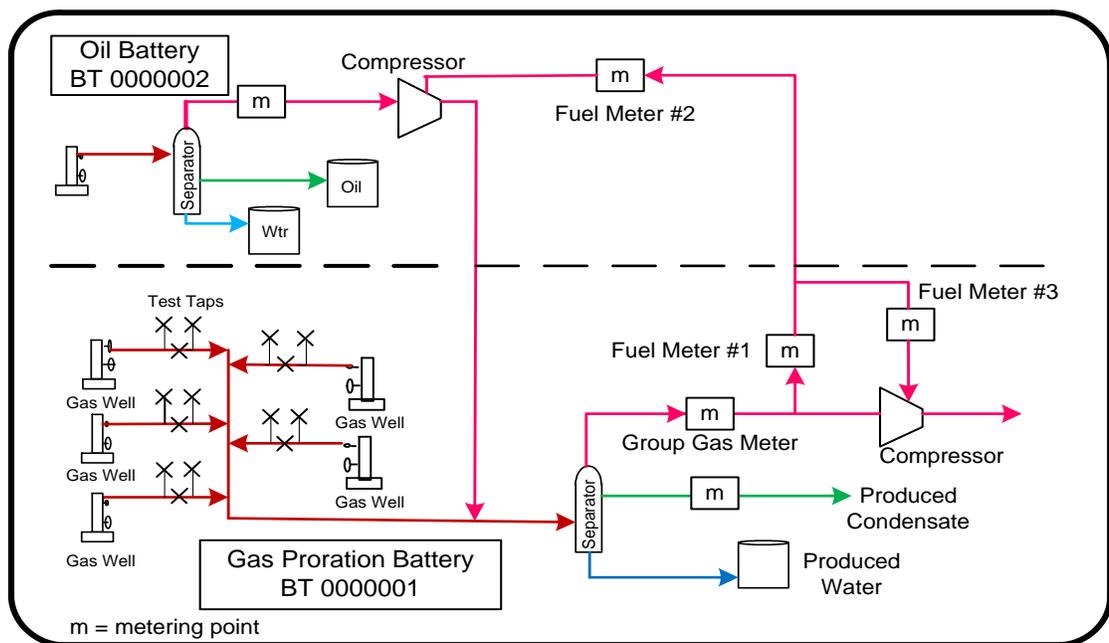
In the scenario of an oil proration battery or a gas proration battery, the monthly gas volume including GEV of condensate where applicable received from a tied-in measured gas source and any other receipts must be subtracted from the total monthly battery gas volume including GEV of condensate where applicable to determine the monthly battery gas production volume.

In the scenario of an oil proration battery, the monthly liquid condensate, oil, or oil-water emulsion volume, where applicable, received from a tied-in measured source must be subtracted from the total monthly oil and/or water disposition plus/minus inventory changes and minus any other receipts to determine the monthly battery oil and/or water production volume.

### 5.5.6 Fuel Gas Mbd

Section 4.3.6.1 (12) describes the requirements for fuel gas measurement and reporting at sites where there may be multiple facility reporting codes and the fuel gas consumption is  $> 0.5 \cdot 10^3 \text{m}^3/\text{day}$ . Situations may occur where fuel gas is metered and consumed at one site and some of the metered fuel gas is then sent to another site (separate geographic location) where it is consumed (see Figure 5.15). Three acceptable fuel gas Mbd scenarios are described below.

Figure 5.15 Fuel gas Mbd scenarios



1. Site fuel gas at BT 0000001 is measured at fuel meter #1. The volume of fuel gas sent to BT 0000002 is  $\leq 0.5 \cdot 10^3 \text{m}^3/\text{day}$ , and the volume of fuel gas consumed at the compressor at BT 0000001 is  $> 0.5 \cdot 10^3 \text{m}^3/\text{day}$ . In this case, fuel gas Mbd is acceptable for the reported fuel gas BT 0000001, and the reported fuel gas at BT 0000001 will equal fuel meter #1 minus fuel meter #2. If the fuel gas sent to BT 0000002 is  $> 0.5 \cdot 10^3 \text{m}^3/\text{day}$  and the fuel gas consumed at the compressor at BT 0000001 is  $\leq 0.5 \cdot 10^3 \text{m}^3/\text{day}$  then fuel gas Mbd is acceptable for the reported fuel

gas at BT 0000002, and the reported fuel gas at BT 0000002 will equal fuel meter #1 minus fuel meter #3.

2. Site fuel gas at BT 0000001 is metered at fuel meter #1. The volume of fuel gas sent to BT 0000002 is  $> 0.5 \text{ } 10^3 \text{ m}^3/\text{day}$ , and the volume of fuel gas consumed at the compressor at BT 0000001 is  $> 0.5 \text{ } 10^3 \text{ m}^3/\text{day}$ . In this case, MbD is acceptable for the fuel gas used at either BT 0000001 or BT 0000002; depending on which site is expected to have the higher reported fuel gas volume. If the fuel gas volume at BT 0000002 will be less than the fuel gas volume at BT 0000001, then fuel gas MbD is acceptable for BT 0000001, and the reported fuel gas at BT 0000001 will equal fuel meter #1 minus fuel meter #2. If the fuel gas volume at BT 0000002 will be less than the fuel gas volume at BT 0000001, then fuel gas MbD is acceptable for BT 0000002, and the reported fuel gas at BT 0000002 will equal fuel meter #1 minus fuel meter #3.
3. Site fuel gas at BT 0000001 is measured at fuel meter #1. The monthly volume of fuel gas sent to BT 0000002 is  $< 0.5 \text{ } 10^3 \text{ m}^3/\text{day}$ , and the monthly volume of fuel gas consumed at the compressor at BT 0000001 is  $< 0.5 \text{ } 10^3 \text{ m}^3/\text{day}$ . In this case, reported fuel gas volumes for BT 0000001 and BT 0000002 may be prorated from the metered monthly fuel gas volume at fuel meter #1 and will be based on each battery's percentage of the total estimated monthly fuel gas volumes at both batteries. For example, reported monthly fuel gas volumes at BT 0000001 = fuel meter #1  $\times$  BT 0000001 estimated fuel  $\div$  (BT 0000001 estimated fuel + BT 0000002 estimated fuel). Battery fuel gas estimates should be based on sound engineering estimates.

## 5.6 Surface Commingling of Multiple Gas Stratigraphic Units or Zones or Wells

If gas wells have been completed in multiple stratigraphic units or zones and those stratigraphic units or zones are segregated in the wellbore and produced separately to surface, or if there are multiple individual gas wells on the same surface location, production from each stratigraphic unit or zone or each well usually has to be measured separately prior to commingling. Where applicable, such stratigraphic units or zones or wells may be commingled at surface prior to the combined production being measured, if the qualifying criteria in Section 5.6.1.1 are met or upon ER approval of an application. Proportionate monthly production volumes must still be determined and reported for each stratigraphic unit or zone or well, in accordance with the applicable criteria and considerations described in Sections 5.6.1.1 and 5.6.3.

The following criteria and considerations do not apply to wells that qualify for the Gas Multi-well Proration SW Saskatchewan Battery procedures if specific stratigraphic units or zones are approved (without application) for commingling in the wellbore.

Commingling of stratigraphic units or zones in the wellbore requires approval from ER.

### 5.6.1 Exceptions

Surface commingling of two gas stratigraphic units or zones in a gas well or separate gas wells on the same surface location prior to measurement is allowed without ER site-specific approval if all the qualifying criteria in Section 5.6.1.1 are met.

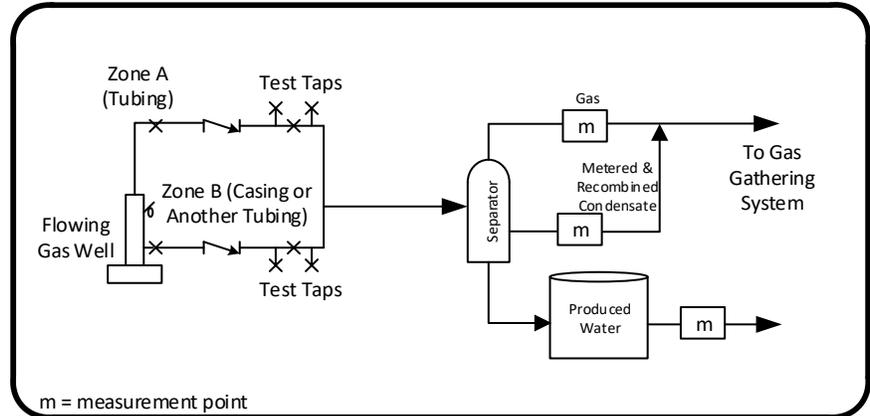
### 5.6.1.1 Qualifying Criteria

1. Both stratigraphic units or zones or wells:
  - a. have common working interest ownership, and where there is no common ownership, written notification has been provided to all working interest participants. If the owner does object then the licensee is required to notify ER;
  - b. have either common Crown or common Freehold royalties. If there are no common Crown or common Freehold royalties and only Freehold royalties are involved, written notification has been given to all Freehold royalty owners. If the owner does object then the licensee is required to notify ER.
2. Monthly average of total liquid production from both stratigraphic units or zones is  $\leq 2.0$  m<sup>3</sup>/day.
3. The combined daily flow rate of both stratigraphic units or zones or wells is  $\leq 16.9$  10<sup>3</sup>m<sup>3</sup>, including GEV of condensate (if recombined). If the stratigraphic units or zones or wells to be commingled will involve existing production, initial qualifying flow rates are based on monthly average flow rates recorded during the six months prior to implementation of the commingling. If new stratigraphic units or zones/wells are to be commingled, the initial qualifying flow rates are based on production tests conducted under the anticipated operating conditions.
4. Shut-in wellhead pressure of the lower pressure stratigraphic zone/well is  $\geq 75.0$  per cent of the shut-in wellhead pressure of the higher-pressure stratigraphic unit or zone.
5. The combined production from both stratigraphic units or zones/wells is measured continuously. Separation before measurement is required for all phases.
6. Check valves are installed on each flow line upstream of the commingling point.
7. Testing requirements:
  - a. Each stratigraphic unit or zone or well must be tested once per month for the first six months after commingling, then annually thereafter, and/or immediately following any significant change to the producing conditions of either stratigraphic unit or zone or well.
  - b. The tests must be conducted for at least 24.0 hours and must involve the separation and measurement of all gas and liquid production.
  - c. If condensate is recombined with the gas production of the commingled stratigraphic units or zones or wells, a sample of the condensate must be taken annually and analyzed and used to determine the factor to be used to determine the GEV.
  - d. The tests for both stratigraphic units or zones or wells must be done consecutively with stabilization periods.
  - e. Any of the three test methods may be used. Methods (1) and (2) are preferred given the testing is conducted under normal flowing conditions and is performed without shutting in stratigraphic units or zones or wells, so minimal stabilization time is required.

**Test Method (1)** Test taps must be installed upstream of the commingling point but downstream of the check valve so that a test

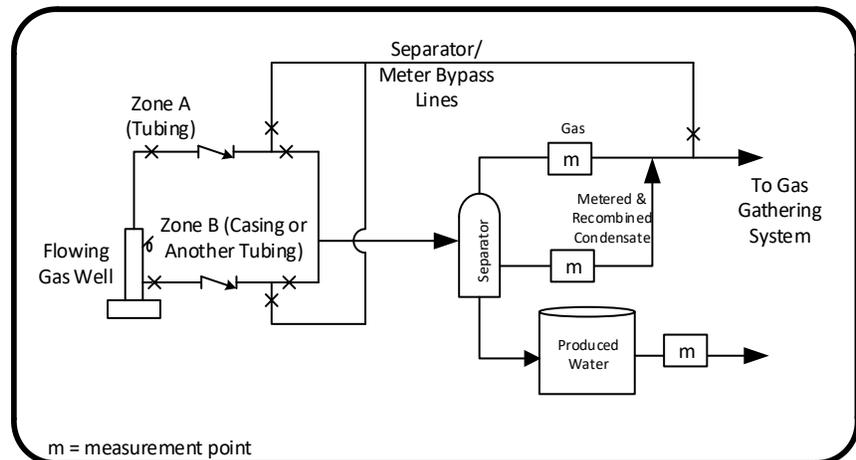
separator unit can be hooked up to test each stratigraphic unit or zone or well individually (Figure 5.16).

**Figure 5.16 Surface comingling of multiple gas zones/wells test method 1**



**Test Method (2)** Install permanent bypasses or taps to hook up temporary bypasses downstream of the check valve so that one stratigraphic unit or zone or well will be bypassing the existing separation and metering equipment while the other stratigraphic unit or zone or well is tested using the existing equipment. Note that the production from the bypassed stratigraphic unit or zone or well must be estimated based on the production test rates (Figure 5.17).

**Figure 5.17 Surface comingling of multiple gas zones/wells test method 2**



**Test Method (3)** Shut in one producing stratigraphic unit or zone at a time and use the existing separation and measurement equipment to test each stratigraphic unit or zone or well individually after stabilization (Figure 5.17).

8. The production rates determined for each stratigraphic unit or zone or well by the periodic tests must be used to estimate the monthly production for each stratigraphic

unit or zone or well from the date they are conducted until the date the next test is conducted. The monthly measured combined production must be prorated to each stratigraphic unit or zone or well based on the estimates, and those prorated volumes must be reported as the monthly production for each stratigraphic unit or zone or well.

### 5.6.1.2 Revocation of Exceptions

If any of the following scenarios exists or occurs, the exception is revoked:

1. The combined production from both stratigraphic units or zones or wells was not measured continuously or there was no separation before measurement.
2. Check valves were not installed on each flow line upstream of the commingling point.
3. Testing requirements in item 7 under Qualifying Criteria in Section 5.6.1.1 were not followed.
4. The gas proration methodology in item 8 under Qualifying Criteria in Section 5.6.1.1 was not followed.

Base measurement requirements must be reinstated if the exception is revoked due to any of these scenarios.

### 5.6.2 Applications

The following information must be submitted with an application to commingle production at surface prior to measurement from multiple stratigraphic units or zones in a gas well or multiple wells on the same surface location if the qualifying criteria in Section 5.6.1.1 are not met:

1. All of the information listed in Section 5.2;
2. Shut-in and proposed operating pressures at the wellhead for all stratigraphic units or zones or wells;
3. Operating pressure for the gathering system at the well's measurement point;
4. Proposed testing procedures to determine the individual stratigraphic unit or zone or well production rates;
5. Proposed accounting procedures for prorating total volumes to the individual stratigraphic units or zones or wells; and
6. All wells flowing to the battery:
  - a. have common working interest ownership, and where there is no common ownership, written notification has been provided to all working interest participants. If the owner does object then the licensee is required to notify ER;
  - b. have either common Crown or common Freehold royalties. If there are no common Crown or common Freehold royalties and only Freehold royalties are involved, written notification has been given to all Freehold royalty owners. If the owner does object then the licensee is required to notify ER.

### 5.6.3 Considerations for Site-specific Approval

1. Generally, there is  $\leq 2.0 \text{ m}^3/\text{day}$  of total liquid production from all stratigraphic units or zones or wells.
2. All stratigraphic units or zones or wells must be classified as gas stratigraphic units or zones or wells.
3. There are minimal equity, royalty, and reservoir engineering concerns.
4. The combined production of all stratigraphic units must be continuously measured. If there are gas and liquid components, they must be separated and measured in a single phase.
5. Check valves must be in place on the flow line upstream of the commingling point.
6. Testing requirements:
  - a. Each stratigraphic unit or zone or well must be tested once per month for the first six months after commingling, then annually after that, and/or immediately following any significant change to the producing conditions of either stratigraphic unit or zone or well.
  - b. The tests must be conducted for at least 24.0 hours in duration and must involve the separation and measurement of all gas and liquid production.
  - c. If condensate is recombined with the gas production of the commingled stratigraphic units or zones or wells, a sample of the condensate must be taken annually and analyzed and used to determine the factor that will be used to determine the GEV.
  - d. The tests for all stratigraphic units or zones or wells must be done consecutively, with stabilization periods.
  - e. Any of the three test methods described in the exceptions in Section 5.6.1.1 may be used, with the consideration that more than two stratigraphic units or zones or wells may be involved. Methods (1) and (2) are preferred, because the testing is conducted under normal flowing conditions without shutting in stratigraphic units or zones or wells, so that minimal stabilization time is required. ER may specify test procedures if specific circumstances warrant them.
7. The production rates determined for each stratigraphic unit or zone or well by the periodic tests must be used to estimate the monthly production for each stratigraphic unit or zone or well from the date they are conducted until the next test is conducted. The monthly measured combined production must be prorated to each stratigraphic unit or zone or well based on the estimates, and those prorated volumes must be reported as the monthly production for each stratigraphic unit or zone or well.

## 6 Non-Heavy Oil Measurement

This section presents the base requirements and exceptions for non-heavy crude oil and emulsion measurements from wells and batteries in the upstream oil and gas industry that are used in determining volumes for reporting to Petrinex. The requirements for crude oil/emulsion volumes transported by truck are detailed in Section 10.

Non-heavy crude oil has the following characteristics:

1. It is a mixture mainly of pentanes and heavier hydrocarbons that may be contaminated with sulphur compounds,
2. It is recovered or is recoverable at a well from an underground reservoir,
3. It is liquid at the conditions under which its volume is measured or estimated, and
4. It has a density of less than 920.0 kg/m<sup>3</sup> at standard conditions.

### 6.1 General Requirements

Crude oil may be found in association with water in an emulsion. In such scenarios, the total liquid volume must be measured, and the relative volumes of oil and water in the emulsion must be determined by obtaining and analyzing a representative sample of the emulsion, by using a product analyzer, or by other means if applicable. Applications for which estimation of water content is appropriate e.g., inventory, are covered in more detail later in this section.

A licensee must measure produced crude oil/emulsion volumes by tank gauging, weigh scale, or meter unless otherwise stated in this Directive. ER will consider an oil measurement system to be in compliance if the requirements detailed in Sections 6.2 and 6.3 are met. ER may stipulate additional or alternative requirements for any specific situation based on a site-specific assessment and will inform licensees in writing of any additional or alternative requirements respecting their facilities.

### 6.2 General Measurement, Accounting, and Reporting Requirements for Various Battery Types

#### 6.2.1 General Accounting Formula

$$\text{Production} = \text{Total disposition} + \text{Closing inventory} - \text{Opening inventory} - \text{Total receipts}$$

#### 6.2.2 Oil Batteries

All wells in the battery must be oil wells.

Liquid production from an oil battery must be measured as an oil, water, or oil/water emulsion volume. This measurement may be performed at the battery site, a truck delivery/receipt point, or a pipeline delivery point. The meter factor obtained from meter proving must be applied to the metered volumes until another prove is conducted.

All wells in a multi-well oil battery must be subject to the same type of measurement: measured or prorated. If there is a mixture of measured and prorated wells delivering production to the proration battery, the measured well(s) must be linked to their own Petrinex reporting facility ID(s) and, the MBD exception criteria in Sections 5.5 must be met or an ER site-specific approval must be obtained.

Production from gas batteries or other oil batteries cannot be connected to an oil proration battery upstream of the oil proration battery group measurement point(s) unless specific MbD exception criteria are met or ER site-specific approval is obtained as per Sections 5.5. For oil delivered to a gas system, see Section 6.2.3.

Any oil well that produces fluids from any stratigraphic unit is considered on production and a battery reporting facility ID is required to report the production on Petrinex even for a test prior to the well being put on permanent production operations, see *Directive PNG032: Volumetric, Valuation and Infrastructure Reporting*.

**6.2.2.1 Crude Oil Single-well Battery (Petrinex facility subtypes: 311)**

Oil/emulsion must be separated from gas and measured.

**6.2.2.2 Crude Oil Multi-well Group Battery (Petrinex facility subtypes: 321)**

Each well must have its own separation and measurement equipment, similar to a single-well battery.

All separation and measurement equipment for the wells in the battery, including the tanks but excluding the wellheads, must share a common surface location.

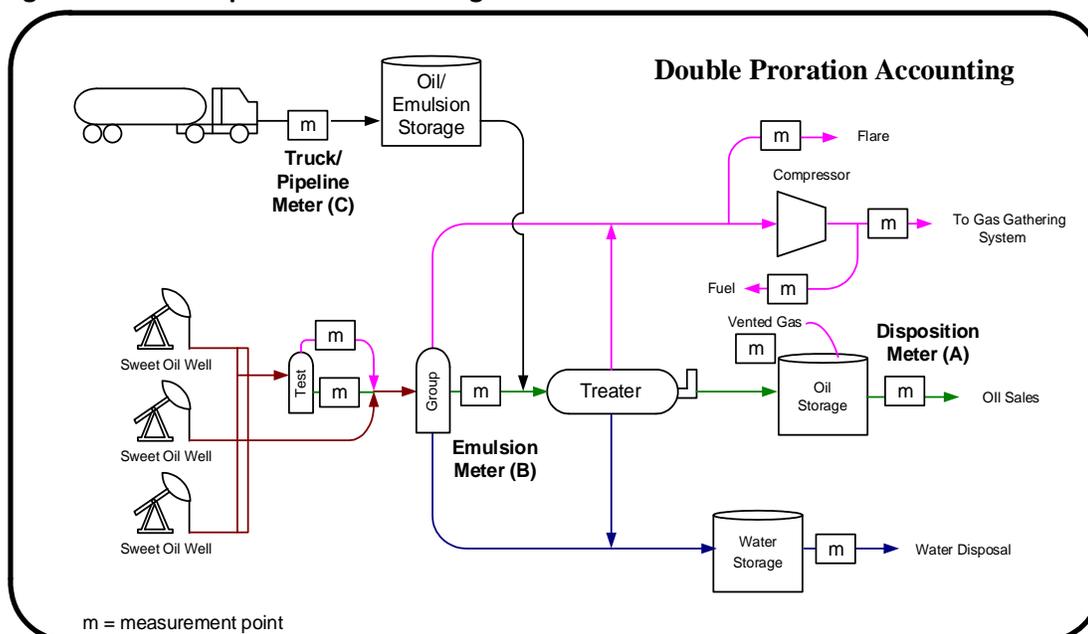
**6.2.2.3 Crude Oil Multi-well Proration Battery (Petrinex facility subtypes: 322)**

All well production is commingled prior to the total battery oil/emulsion being separated from the gas and measured. Individual monthly well oil production is estimated based on periodic well tests and corrected to the reported monthly volume through the use of a proration factor.

Double proration, whereby the proration oil battery disposition volume(s) is prorated to group and receipt measurement points and then further prorated to the wells (see Figure 6.1), is allowed without site-specific approval subject to the following conditions:

1. All prorated oil/emulsion must be measured using measurement systems that meet delivery point measurement uncertainty requirements before commingling with other oil/emulsion receipts. The measurement system must include a meter of sufficient measurement uncertainty and either an online continuous product analyzer or a proportional sampler.
2. The oil/emulsion meter (B) in figure 6.1 must be proved to standard conditions or corrected to standard conditions using a flash liberation analysis.

Figure 6.1 Double proration accounting



Total reported battery oil and water disposition volumes with inventory change must be prorated to the total truck/pipeline volumes measured and the total well emulsion volumes measured (first proration). This proration using PF1 has to be done off-sheet and not reported on Petrinex.

$$PF1 = [\text{meter (A) + INVCL} - \text{INVOP}] \div [\text{meter (B) + meter (C)}]$$

$$\text{Prorated meter (B) volume} = \text{meter (B)} \times PF1$$

$$\text{Prorated individual truck-in and/or pipeline volumes} = \text{meter (C) volumes for each load received} \times PF1$$

$$PF2 = \text{prorated meter (B) volume} \div \text{total estimated production volume}$$

The prorated oil and water volume at the emulsion meter (B) is further prorated using PF2 (second proration) to the tested oil wells. The oil and water proration factors PF2 must then be reported on Petrinex.

### 6.2.3 Gas Batteries Producing Oil

All wells in the battery must be gas wells.

Oil production, receipt, disposition, and inventory volumes must be reported as liquid oil. Oil volumes must not be converted to a gas equivalent volume (GEV) and must not be added to the gas volumes. If oil is recombined with the gas and delivered to a gas plant through a gas gathering system, the oil volume as determined at the battery must be reported as a receipt of OIL by the gas plant. The gas plant must report the oil disposition as appropriate and it is common for gas plants to report oil dispositions as C5-MX or C5-SP.

#### 6.2.3.1 Gas Single-well Battery (Petrinex facility subtype: 351)

Oil/emulsion must be separated from gas and measured.

### 6.2.3.2 Gas Multi-well Group Battery (Petrinex facility subtypes: 361)

Each well must have its own separation and measurement equipment, similar to a single-well battery and its gas production must be connected by pipeline to a common location for further processing.

## 6.3 Base Requirements for Oil Measurement

### 6.3.1 System Design and Installation of Measurement Devices

The system design and installation of oil/emulsion measurement devices must be in accordance with Sections 14.2, 14.3, and 14.7.

EFM systems must be designed and installed according to the requirements in Section 6.8. Any EFM system designed and installed in accordance with API MPMS, Chapter 21.2, is considered to have met the audit trail and reporting requirements, but a performance evaluation is still required in accordance with Section 14.9.2.

### 6.3.2 Volumetric Calculations

Crude oil volume measurements must be determined to a minimum of two decimal places and rounded to one decimal place for monthly reporting. Where there is more than one volume determination within the month at a reporting point, the volumes determined to two decimal places must be totaled prior to the total being rounded to one decimal place for reporting purposes.

#### 6.3.2.1 Temperature Correction Requirements

Temperature measurement used for volume correction must be representative of the actual fluid temperature. Total monthly oil volumes for wells (production) and batteries (production, receipts, dispositions, and delivery point) must be reported in cubic metres at standard temperature of 15.0°C and rounded to the nearest tenth of a cubic metre (0.1 m<sup>3</sup>). Battery or facility opening and closing inventory volumes for monthly reporting must be rounded to the nearest 0.1 m<sup>3</sup> but do not require correction to 15.0°C. The temperature correction (Correction for the effect of Temperature on Liquids [CTL]) factor must be determined in accordance with API MPMS, Chapter 11.1.

In a proration battery, if well test oil volumes are determined by a meter, temperature compensation must be applied using one of the following methods:

1. Apply a composite meter factor that incorporates a CTL factor. To arrive at a composite meter factor, divide the temperature corrected prover volume by the indicated meter volume for each prover run.
2. Apply a CTL factor in real time using an electronic flow measurement system.
3. Apply a CTL factor to the total test volume based on a single temperature measurement taken during the test.

See Section 14.4 for more details.

### **6.3.2.2 Pressure Correction Requirements**

Correction to a 0.0 kPa gauge (atmospheric pressure) must be performed for continuous flow crude oil pipeline measurement where custody transfer measurement is performed. See Section 14.5 for more details.

### **6.3.2.3 Shrinkage Factor**

See Section 14.3 for details.

### **6.3.2.4 General Volume Calculations**

See Section 14.9 for details.

## **6.3.3 Data Verification and Audit Trail**

The field data, records, and any calculations or estimations, including EFM calculations and estimations, relating to ER -required data submitted to Petrinex must be kept for inspection upon request by ER. The reported data verification and audit trails must be in accordance with the following:

1. Test records: any records and documentation produced in the production proration testing of wells that affect measured volumes.
2. Proving records: any records and documentation produced in the proving of meters and calibration of the prover and all peripheral devices if the prover and peripheral devices are owned and operated by the licensee.
3. S&W records: any records and documentation produced in the determination of relative oil/water percentages that affect volumes.
4. Delivery and receipt records: any records and documentation produced in the determination of delivery or receipt volumes.
5. Estimation records: any records and documentation related to the estimation of reported volumes, including estimation methodology, record of event, and approvals.
6. Tank gauging records: any records and documentation produced in the determination of reported volumes.
7. Volume loss records: any records and documentation for volumes lost due to incidents such as theft, spills, and fires.
8. EFM: any records and documentation either electronic, magnetic, or paper form produced in the determination of measured volumes in accordance with the EFM requirements in Section 6.8.

Records of the foregoing must be provided to ER upon request.

## **6.4 Field Operations**

### **6.4.1 Production Hours**

Physical well shut-ins and emergency shutdowns (ESDs) are considered downtime. Other occurrences resulting in downtime include wax or hydrates plugging lines and some other failures. If the well has no oil production but still has gas production, it is considered to be on

production. The operations personnel have to make a determination based on the operating environment in other situations when the wells are not physically shut in but may not have oil and gas production.

Oil wells are considered on production even when the wells are not pumping or flowing in situations where:

1. The wells are operating on an on/off cycle basis, such as intermittent timers, pump-off controls, and plunger lifts;
2. The wells are operating normally and as designed on repeated cycles; and
3. Part of the operation involves shutdown of pump equipment and/or periodic shut-in of the wells as part of the repeated cycle.

#### 6.4.2 Fluid Sampling Requirements for S&W Determination (and Density)

See Section 14.6 for density determination details.

#### 6.4.3 S&W Analysis

Conduct water-cut sampling and analysis for each test.

See Section 14.8 for S&W determination details and Appendix 3 for water-cut procedures.

#### 6.4.4 Proration Well Testing

Proration testing requirements for non-heavy crude oil wells are detailed in Table 6.1.

**Table 6.1 Proration testing requirements for non-heavy crude oil wells**

Class <sup>a</sup>		Oil rate (m <sup>3</sup> /d)	Minimum test frequency	Minimum time between tests <sup>b</sup> (days)	Minimum test duration <sup>c</sup> (hours)
No.	Name				
1	High	> 30.0	3 per month <sup>d</sup>	5.0	12.0
2	Medium	> 6.0 but ≤ 30.0	2 per month <sup>e</sup>	10.0	22.0
3	Low	> 2.0 but ≤ 6.0	1 per month	15.0	22.0
4	Stripper	≤ 2.0	1 every quarter	45.0	22.0

<sup>a</sup> Classification for each well must be determined at least semi-annually based on the average daily oil rate since the last assessment. If a well experiences operational changes that cause a change in the oil rate that could affect the classification, the licensee must immediately change the classification. The average daily oil rate must be based on producing days (not calendar days).

<sup>b</sup> Minimum separation time between tests if minimum number of tests are conducted - the time between tests may be reduced if more than the minimum number of tests are conducted.

<sup>c</sup> Licensees should conduct longer duration tests for wells exhibiting erratic rates to obtain more representative test data.

<sup>d</sup> For Class 1 wells, the minimum test frequency is based on the assumption that the well is on production for the entire calendar month. The test frequency may be reduced to two per month if the well is shut in for at least 10 days within the month and to one per month if the well is shut in for at least 20 days within the month.

<sup>e</sup> For Class 2 wells, the minimum test frequency is based on the assumption that the well is on production for the entire calendar month. The test frequency may be reduced to one per month if the well is shut in for at least 15 days within the month.

#### 6.4.4.1 Well Test Considerations

If there is a change in operating conditions during a test, such as due to a power failure or a change in choke setting, the test must be rejected and a new test must be conducted.

If there is insufficient or lost test data, such as due to meter failure, the test must be rejected and a new test must be conducted.

If there is a significant change in oil, gas, or water for a test, the validity of the test should be questioned and a retest should be considered.

Sufficient purge time must be allowed to ensure that liquids from the previous test are displaced by the new test well liquids.

The pressure difference between the test separator and the group line must not exceed 200.0 kPa.

A well test may be stopped early for operational reasons and still be considered valid. Reasons for the short test must be documented and made available to ER upon request.

#### 6.4.4.2 Common Flow Lines

For common flow lines, a well test must be conducted with all other wells on the common flow line shut. Sufficient purge time must be allowed to ensure that liquids from the common flow line wells are displaced by the test well liquids. Combined (cascade) testing is allowed for common flowlined wells, provided that the conditions in Section 6.4.4.4 are met. However, the combined test must be conducted first, and then the lower gas producing well must be shut in to test the higher gas producing well, allowing sufficient purging and stabilization time.

#### 6.4.4.3 Field Header and Common Flow Line Purging

If a field header is located in the same building as the test separator, the test separator must be purged by allowing at least two liquid dumps to occur prior to starting the well test. The field header must clearly identify which well is tied to the header valves.

If a field header is not located in the same building as the test separator, sufficient purge time must be allowed to ensure that liquids from the previous test are displaced by the new test well liquids.

If two or more wells are tied into a common flow line, only one well must be produced during the well test, and the other well(s) must be shut in. Similar to a field header situation, sufficient purge time must be allowed to ensure that liquids from the previous production condition are displaced by the new test well liquids.

Sufficient purge time must be calculated as follows:

$$\text{Purge time} = \text{test line volume} \div \text{new test well liquid flow rate}$$

**Example:** Calculate the minimum purge time required for the following test line:

Test line dimensions = 1500.0 m length, 88.9 mm OD pipe, 3.2 mm wall thickness

Previous well test flow rates = 5.5 m<sup>3</sup> oil/d, 12.0 m<sup>3</sup> water/d

##### Step 1

$$d = (88.9 - 3.2 \times 2) \div 1000 = 0.0825 \text{ m}$$

$$\text{Test line volume} = (3.142 \times d^2 \times \text{length}) \div 4$$

$$= (3.142 \times (0.0825)^2 \times 1500) \div 4$$

$$= 8.02 \text{ m}^3$$

Where: d = the internal diameter of the flowline

### Step 2

$$\text{Purge time required} = \text{Test line volume (m}^3\text{)} \div \text{Well flow rate (m}^3\text{/hr)}$$

$$\text{Well total liquid flow rate} = (5.5 \text{ m}^3 + 12.0 \text{ m}^3) \div 24.0 \text{ hr}$$

$$= 0.729 \text{ m}^3\text{/hr}$$

$$\text{Purge time required} = 8.02 \text{ m}^3 \div 0.729 \text{ m}^3\text{/hr}$$

$$= 11.0 \text{ hr}$$

Therefore, the minimum purge time required is 11.0 hours.

#### 6.4.4.4 Combined (Cascade) Testing

When a prorated oil well has such low gas production that it cannot properly operate test equipment, a licensee may test two wells simultaneously - combined (cascade) test - through the same test separator. In such scenarios, the following procedure must be followed:

1. Establish oil, gas, and water production volumes for a high gas producing well by testing it individually through the test separator.
2. Conduct a test for both the high gas producing well and a low gas producing well together through the same test separator immediately after testing the high gas producing well, allowing time for stabilization. The testing sequence may be reversed with the testing of the combined wells first.
3. The operating condition of both wells must not be changed. If it is, a new set of tests is required.
4. Total test oil, gas, and water volumes determined for the combined (cascade) test minus the test oil, gas, and water volumes for the high gas producing well will be the test volumes for the low gas producing well.
5. Both wells should have similar S&W percentages. If any of the calculated oil, gas, or water volumes for the low gas producing well are negative, the tests are not valid and both tests must be repeated.

The use of combined (cascade) testing does not require a site-specific approval from ER.

#### Example

Well A = High gas producing

Well B = Low gas producing

**Table 6.3 Test results**

Well	Test date	Oil (m <sup>3</sup> )	Gas (10 <sup>3</sup> m <sup>3</sup> )	Water (m <sup>3</sup> )
------	-----------	-----------------------	---------------------------------------	-------------------------

Well A + B	July 4	80.0	20.0	20.0
Well A	July 5	50.0	19.0	12.0
<b>Well B = (Well A + B) - Well A</b>	<b>July 4</b>	<b>30.0</b>	<b>1.0</b>	<b>8.0</b>

## 6.5 Gas Estimation and Reporting Methods for Non-Heavy Oil Wells

As per Section 4.1, gas streams from non-heavy crude oil wells (with a density  $< 920.0 \text{ kg/m}^3$ ) associated with facility subtypes 311, 314, 316, 321, 322 that is  $\leq 0.5 \text{ } 10^3 \text{ m}^3/\text{day}$ , based on an annual average, the volume may be determined by estimation instead of metering. Gas streams include gas production, well test separator (or dry flow meter) gas, casing gas, flare gas (including incineration), fuel gas, vent gas, and stock tank vapours from produced oil/emulsion tanks and produced water tanks.

Where the gas production rate from a non-heavy crude oil well is  $\leq 0.5 \text{ } 10^3 \text{ m}^3/\text{day}$ , the well gas production may be estimated. The only acceptable estimation method for gas production is using a Gas Oil Ratio (GOR) factor that is applied to the reported monthly oil production. A well's GOR is typically determined by conducting a well test and then dividing the test gas volume by the test oil volume to determine the GOR. The determination of a well's GOR may include some or all of the gas volumes originating from the following sources as applicable:

1. Measured well test separator (or dry flow meter) gas.
2. Measured casing gas.
3. Flare (including incineration), fuel and vent gas volumes.
4. Gas in solution (GIS) determined from a GIS factor, see Section 6.5.3. Although rare, it is possible that a well's GIS gas will be the entire gas volume used in the determination of a well GOR.
5. Stock tank vapours from produced oil/emulsion tanks and produced water tanks.

All applicable well gas production sources must be included in the determination of the well GOR and the number and location of those sources is dependent on the specific operational configuration of the well. A common oversight in the determination of a well's GOR is to not include the well's gas production attributed to the gas that evolves from the oil/emulsion when the oil is depressurized from separator or wellhead pressure to stock tank conditions...the Gas in Solution (GIS). The GIS is typically just one of the gas production sources that must be included in the determination of a well's GOR.

Acceptable methods for determining a GOR factor and the GIS can be found in Section 6.5.3.

In some scenarios a gas volume must be determined, such as the GIS, where the gas is dissolved in an oil volume under pressure, and there is no opportunity to measure the gas volume prior to it being commingled with other gas volumes. In that scenario, the gas volume may be determined by estimation. An example of such a gas volume is the gas held in solution with oil volumes leaving a test separator at an oil proration battery, where the test oil volumes are combined with production from other wells downstream of the test separator. The purpose of estimating the gas in solution is to determine the total gas produced by a well during a production test, since the gas volume measured by the test gas meter will not include the gas that is still in solution with the test oil volume.

A single GIS factor may be determined and used to estimate the gas volume held in solution with the oil stream for each oil stream where the production sources (producing formation) are the same and test separator operating conditions are similar. Additional GIS factors are required for wells in the battery that produce from different formations and where other test separators operate at different pressure and/or temperature conditions. Licensees should also consider determining seasonal GIS factors where ambient temperature differences may affect the factors or when operating conditions change. Acceptable methods for determining the GIS are described in section 6.5.3

### **6.5.1 Crude Oil Single-Well Batteries (facility subtype 311), Crude Oil Multi-well Group Batteries (facility subtype 321), Crude Oil Multi-well Swab Group Battery (314), and Crude Oil Multi-well Swab Paper Battery (facility subtype 316)**

For new non-heavy crude oil wells, a GOR test must be conducted within first 90 days of the well being put on production, and annually thereafter. Licensees must use acceptable estimation methods for gas production until GOR test can be conducted. The GOR tests must be conducted annually to confirm continuing eligibility for gas volume estimation and to update the factors used to estimate gas volumes. GOR tests must also be conducted and factors updated immediately following any operational changes that could cause the factors to change. Licensees should also consider determining seasonal GOR factors if ambient temperature changes.

Acceptable methods for determining a GOR factor and GIS are described in Section 6.5.3.

#### **6.5.1.1 Exception for GOR testing frequency**

For existing non-heavy crude oil single well batteries and non-heavy crude oil multi-well group batteries that are on an annual GOR testing frequency, ER authorizes the extension, without site-specific approval, of the frequency of gas oil ratio (GOR) testing from annually to triennially if all the qualifying criteria below are met. ER may require the licensee of a well to conduct a GOR test that may be witnessed by the Minister; and to submit the results of the test within the time and in the manner specified by the Minister.

##### **Qualifying Criteria**

Qualifying criteria that must be met include the following:

1. The oil well must produce  $\leq 15.0 \text{ m}^3/\text{day}$  of oil on an annual average basis;
2. The oil well must produce  $\leq 0.5 \cdot 10^3 \text{ m}^3/\text{day}$  of gas on an annual average basis;
3. The well must have had two consecutive annual GOR tests conducted subsequent to an initial GOR test, and the initial test can be no older than 36 months. Therefore, for a new well, at least three annual GOR test are required. The two GOR tests must show that the GOR has not changed by more than  $\pm 20.0$  per cent;
4. On an annual basis, the licensee must review and confirm that the well is still eligible for the extended GOR testing frequency and gas estimation based on the oil and gas production rates at the time and the qualifying rates in item 1 and 2 above; and
5. Test records must be kept and made available to ER upon request.

##### **Revocation of Exception**

If any of the following scenarios exists or occurs, the exception is revoked, the well must be GOR tested as soon as is reasonability possible, the GOR testing frequency reverts to annual and the well must requalify for the exception:

1. A licensee fails to provide test records to ER upon request.
2. If any changes are made to the operations of the facility that could affect gas production such as, but not limited to, fracing nearby wells, pressure changes at the well head, new equipment being installed at the facility, or workovers on the well.
3. If a well that qualifies for this exception has a GOR test conducted in the third year and the GOR from that test has changed by more than  $\pm 20.0$  per cent compared to the current GOR, then the next GOR test must be conducted within 12 months and the resulting GOR must not have changed by more than  $\pm 20.0$  per cent to re-qualify the well for triennial testing. This GOR test re-qualification applies to wells that have not undergone operational changes that could have impacted the GOR.

### 6.5.2 Crude Oil Multi-well Proration Battery (facility subtype 322)

For crude oil multi-well proration batteries (facility subtype 322), each well with gas production  $> 0.5 \text{ } 10^3\text{m}^3/\text{day}$  must be proration tested and follow the well proration testing frequency requirements in 6.4.4, unless metered, on a per stream basis. Fuel gas, vent gas or flare gas at a site must be reported at the location which the activity is occurring.

Gas production volumes  $\leq 0.5 \text{ } 10^3\text{m}^3/\text{day}$  may be estimated using one of the following methods:

1. A well-level GOR. A well-level GOR factor must be determined at the frequency listed in Section 6.5.1. Acceptable methods for determining the GOR factor and GIS are described in Section 6.5.3.
2. A battery-level GOR. A battery-level GOR factor (monthly battery gas production  $\div$  monthly battery oil production) must be determined on a monthly basis. To calculate the individual well gas production using the battery-level GOR factor the following conditions must be met:
  - a. All wells using the battery-level GOR must produce  $\leq 0.5 \text{ } 10^3\text{m}^3/\text{day}$  of gas;
  - b. Any well producing  $> 0.5 \text{ } 10^3\text{m}^3/\text{day}$  of gas is not eligible to use the battery-level GOR, and well gas production must be determined using test rates obtained during proration testing;
  - c. Monthly gas and oil volumes from wells not eligible to use the battery-level GOR must be subtracted from the total battery gas and oil production volumes before calculating the battery-level GOR. For gas, the volume to be subtracted would be the total estimated gas determined from proration testing for all the ineligible wells; for oil, the volume would be the total prorated oil production for all the ineligible wells;
  - d. New wells added must produce  $\leq 0.5 \text{ } 10^3\text{m}^3/\text{day}$  of gas for a minimum of six months before being eligible to use the battery-level GOR;
  - e. If there is no common ownership of all the wells in the battery, written notification must have been given to all working interest participants, with no resulting objections; and
  - f. If there is no common Crown or common Freehold royalty and only Freehold royalties are involved in all wells in the battery, written notification must have been

given to all Freehold royalty owners, with no resulting objection received. If there is a mix of Freehold and Crown royalties involved, the licensee must apply to ER for approval if any Freehold royalty owner objects.

### 6.5.3 Methods for Determining Factors Used in Estimating Gas Volumes using a GOR Factor

If gas volumes are determined by estimation using GOR:

1. A 24-hour test must be conducted such that all the applicable gas and oil volumes produced during the test are measured including vented gas (see figure 6.2). The gas volume is to be divided by the oil volume to determine the GOR factor.
2. A sample of oil taken under pressure containing the gas in solution that will be released when the oil pressure is reduced may be submitted to a laboratory where a PVT analysis can be conducted. The analysis must be based on the actual pressure and temperature conditions the oil sample would be subjected to downstream of the sample point. This procedure is used to determine a well's GIS factor. The GIS factor will be calculated based on the volume of gas released from the sample and the volume of oil remaining at the end of the analysis procedure. The volume of GIS gas determined from the GIS factor is then added to the measured test gas volume, and other gas volumes as applicable, to determine the well's total gas production and GOR as described in item 1 above.
3. A sample of oil taken under pressure containing the gas in solution that will be released when the oil pressure is reduced may be submitted to a laboratory where a compositional analysis can be conducted. A computer simulation program may then be used to determine the GIS factor based on the compositional analysis and the actual pressure and temperature conditions the oil sample would be subjected to downstream of the sample point. This is commonly called a flash liberation analysis or FLIB. The volume of GIS gas determined from the GIS factor is then added to the measured test gas volume, and other gas volumes as applicable, to determine the well's total gas production and GOR as described in item 1 above.

In addition to the methods described above to determine the GIS factor, other acceptable methods include:

1. Methods listed under *Guideline PNG035: Estimating Venting and Fugitive Emissions*.
2. A rule of thumb estimate ( $0.0257 \text{ m}^3$  of gas/ $\text{m}^3$  of oil/kPa of pressure drop) may be used as the GIS factor for non-heavy oil production until a more accurate, specific GIS factor is determined. It may be used on a continuous basis, without the need for determining a more accurate GIS factor if well oil production rates are  $\leq 2.0 \text{ m}^3/\text{day}$ .

For each GOR test conducted, the following must be documented and be made available to ER upon request:

- The meter readings, associated data (e.g. temperature and pressure), instantaneous flow rate and cumulative test volume for each time period (e.g. 20.0 minutes) that the measurements were taken for each determination of the value of gas and oil.

- The date, time, and duration of each of those periods. The maximum period duration must be no longer than 1.0 hour.
- The production parameters (e.g. pressure, pump rate) during each of those periods and in the 48.0 hours before each of those periods; and
- A description of the meter(s) and other equipment used during the test. At minimum the meter make, type, and model should be included.
- Any factors used (e.g. GIS) to determine estimated volumes and the associated documentation of the estimation methods

Additional guidance for how to determine GOR and GIS can be found in *Guideline PNG034: Determination and Submission of Gas Oil Ratio*.

### 6.5.3.1 Example Calculations for Estimating Gas Volumes using GOR and GIS Factors

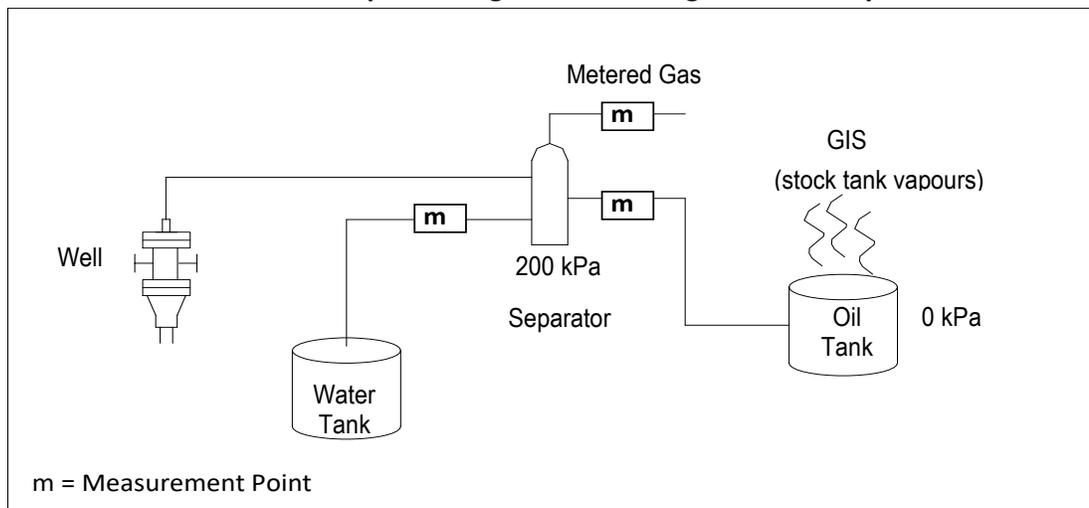
#### **Example 1: Determination of Total Produced Gas for a Single-Well Oil Battery**

Figure 6.2 depicts a single-well battery where a three-phase separator is used to separate oil, gas, and water production from a well. The oil in the separator is under pressure until it is directed to the storage tank, which is at atmospheric pressure (zero kPa gauge). When the oil pressure drops at the tank, the GIS within the oil will be released. The gas leaving the separator in this example is metered, while the GIS released at the tank is estimated using a GIS factor. Total gas production from the well is determined by adding the metered gas and the GIS released at the oil storage tank.

If a single-well battery uses a two-phase separator, the procedure for determining total gas production is the same as for a three-phase separator.

If the gas production rate meets the qualifying criteria for estimation, all production from the well produces directly to a tank without using a separator and there is no free gas in the oil stream, the total gas production may be determined by using only a GIS factor.

**Figure 6.2 Determination of total produced gas for an oil single-well battery**



#### **Sample Calculation: Total Gas Volume at a Single-Well Battery (Figure 6.2)**

Monthly well data (hypothetical) given for this example:

Gas meter volume =  $96.3 \times 10^3 \text{ m}^3$  (from chart readings)

Oil meter volume =  $643.3 \text{ m}^3$  (from meter or tank gauging)

Pressure drop = 200.0 kPa

GIS factor =  $6.37 \text{ m}^3 \text{ gas}/\text{m}^3 \text{ oil}$  or  $0.03185 \text{ m}^3 \text{ gas}/\text{m}^3 \text{ oil}/\text{kPa}$  pressure drop (determined using a method other than the rule of thumb)

**Step 1:** Calculate GIS volume:

$$6.37 \text{ m}^3/\text{m}^3 \times 643.3 \text{ m}^3 = 4097.8 \text{ m}^3 = 4.10 \times 10^3 \text{ m}^3$$

**or**

$$0.03185 \text{ m}^3/\text{m}^3/\text{kPa} \times 643.3 \text{ m}^3 \times 200.0 \text{ kPa} = 4097.8 \text{ m}^3 = 4.10 \times 10^3 \text{ m}^3$$

**Step 2:** Calculate the total battery gas production for the month:

$$96.3 \times 10^3 \text{ m}^3 + 4.1 \times 10^3 \text{ m}^3 = 100.4 \times 10^3 \text{ m}^3$$

Note that total reported battery gas production is to be rounded to one decimal place.

**Example 2: Determination of Total Produced Gas for an Oil Proration Battery**

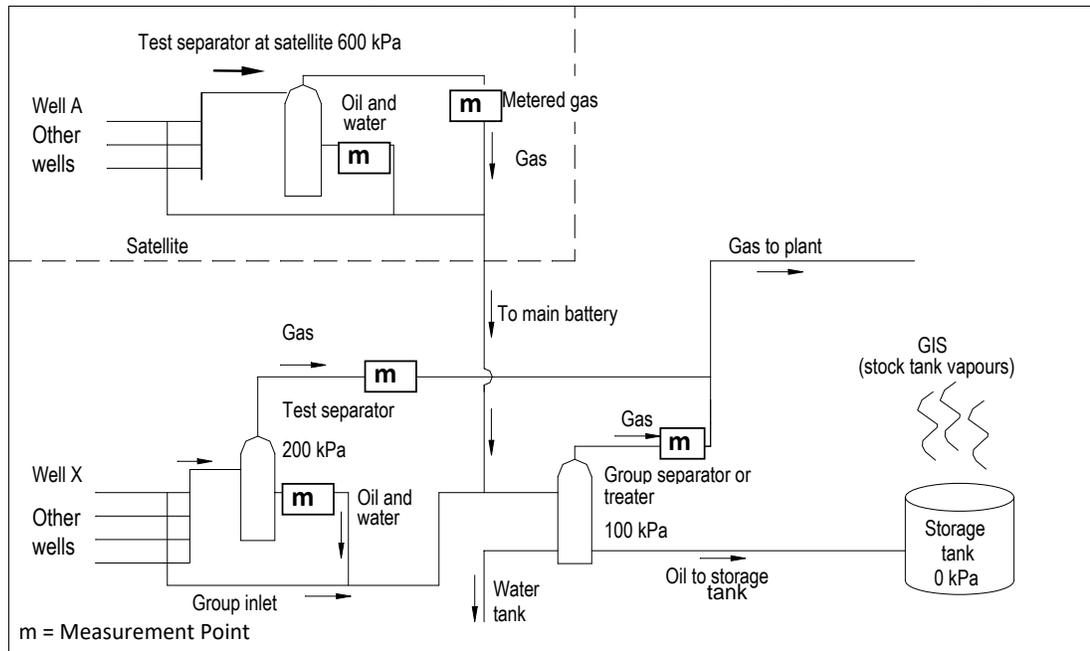
Figure 6.3 depicts a multi-well oil proration battery where production testing of individual wells is done by directing individual well production through a test separator at the main battery site or through a test separator at a satellite facility located away from the main battery site.

In this example, the oil, gas, and water leaving the test separator at the satellite are recombined with the satellite group production and directed to the group separation and measurement equipment at the main battery site. The oil and water leaving the test separator at the main battery site are recombined with the battery group production, but the gas leaving the test separator recombines with the group gas downstream of the group gas measurement point. The oil in the group separator is under pressure until it is directed to the storage tank, which is at atmospheric pressure (zero kPa gauge). When the oil pressure drops at the tank, the GIS with the oil will be released.

The total gas production at the battery will be the sum of all the measured test gas at the battery site, the measured group gas at the battery, and the GIS released at the oil storage tank.

Trucked oil volumes received at the battery must not be included with the total battery oil volume when determining the GIS released at the oil storage tank.

At some facilities a vapour recovery unit (VRU) may be installed to collect any GIS that may be released at the oil storage tank. If the VRU is equipped with a meter or the recovered gas is directed through the group gas meter, a GIS calculation will not be required because the measured VRU gas will either be added to or included in the other measured gas volumes.

**Figure 6.3 Determination of total produced gas for an oil multi-well proration battery****Sample Calculation: Total Gas Production at the Oil Multi-well Proration Battery (Figure 6.3)**

Monthly battery data (hypothetical) given for this example:

Oil production at the proration battery = 745.0 m<sup>3</sup> for the month (from meter and/or tank gauging)

Total test gas measured at the battery site = 30.0 10<sup>3</sup>m<sup>3</sup> (from chart readings)

Measured group gas production = 67.4 10<sup>3</sup>m<sup>3</sup> (from chart readings)

Pressure drop from the group vessel to oil storage tank = 100.0 kPa

GIS factor = 3.99 m<sup>3</sup> gas /m<sup>3</sup> oil or 0.0399 m<sup>3</sup>/m<sup>3</sup>/kPa (determined using a method other than the rule of thumb)

**Step 1:** Calculate the GIS volume:

$$3.99 \text{ m}^3/\text{m}^3 \times 745.0 \text{ m}^3 = 2972.6 \text{ m}^3 = 2.97 \text{ 10}^3\text{m}^3$$

or

$$0.0399 \text{ m}^3/\text{m}^3/\text{kPa} \times 745.0 \text{ m}^3 \times 100.0 \text{ kPa} = 2972.6 \text{ m}^3 = 2.97 \text{ 10}^3\text{ m}^3$$

**Step 2:** Calculate the total produced gas volume for the battery:

$$67.4 \text{ 10}^3\text{m}^3 + 30.0 \text{ 10}^3\text{ m}^3 + 2.97 \text{ 10}^3 \text{ m}^3 = 100.4 \text{ 10}^3 \text{ m}^3$$

Note that total reported battery gas production is to be rounded to one decimal place.

**Example 3: Determination of Individual Well Test Gas for an Oil Multi-well Proration Battery**

Figure 6.3 depicts a multi-well oil proration battery where production testing of individual wells is done by directing individual well production through a test separator at the main battery site or through a test separator at a satellite facility located away from the main battery site. In either scenario, the oil leaving the test separator is under pressure and will be subjected to two stages of pressure drop—one at the group separator and one at the storage tank. The total gas

produced by a well during a test will be the sum of the gas measured as it leaves the test separator and the GIS that will evolve from the test oil volume after leaving the test separator. In the example, the test separators at the battery and satellite operate at significantly different pressures, and the oil leaving the test separator at the satellite will contain more GIS than the oil leaving the test separator at the battery.

**Sample Calculation: Test Gas Production for Wells in the Satellite (Figure 6.3)**

Satellite test data (hypothetical) given for this example for well A:

Metered test oil =  $7.22 \text{ m}^3$  (from oil meter)

Metered test gas =  $1.27 \times 10^3 \text{ m}^3$  (from chart readings)

GIS factor =  $25.62 \text{ m}^3 \text{ gas}/\text{m}^3 \text{ oil}$  or  $0.0427 \text{ m}^3 \text{ gas}/\text{m}^3 \text{ oil}/\text{kPa}$  pressure drop (combined GIS for both stages of pressure drop from test pressure at 600.0 kPa to group pressure at 100.0 kPa to oil storage tank pressure at atmospheric pressure or zero kPa gauge, determined using a method other than the rule of thumb)

**Step 1:** Calculate the GIS volume:

$$0.0427 \text{ m}^3/\text{m}^3/\text{kPa} \times 7.22 \text{ m}^3 \times 600.0 \text{ kPa} = 185.0 \text{ m}^3 = 0.19 \times 10^3 \text{ m}^3$$

**or**

$$25.62 \text{ m}^3/\text{m}^3 \times 7.22 \text{ m}^3 = 185.0 \text{ m}^3 = 0.19 \times 10^3 \text{ m}^3$$

**Step 2:** Calculate the total test gas produced for well A for this test:

$$1.27 \times 10^3 \text{ m}^3 + 0.19 \times 10^3 \text{ m}^3 = 1.46 \times 10^3 \text{ m}^3$$

Note that test gas volumes must be determined to two decimal places (in  $10^3 \text{ m}^3$ ).

**Sample Calculation: Test Gas Production for Wells in the Battery (Figure 6.3)**

Battery test data (hypothetical) given for this example for well X:

Metered test oil =  $3.85 \text{ m}^3$  (from oil meter)

Metered test gas =  $2.33 \times 10^3 \text{ m}^3$  (from chart readings)

GIS factor =  $7.90 \text{ m}^3 \text{ gas}/\text{m}^3 \text{ oil}$  or  $0.0395 \text{ m}^3 \text{ gas}/\text{m}^3 \text{ oil}/\text{kPa}$  pressure drop (combined GIS for both stages of pressure drop from test pressure at 200.0 kPa to group pressure at 100.0 kPa to oil storage tank pressure at atmospheric pressure or zero kPa gauge, determined using a method other than the rule of thumb)

**Step 1:** Calculate the GIS volume:

$$0.0395 \text{ m}^3/\text{m}^3/\text{kPa} \times 3.85 \text{ m}^3 \times 200.0 \text{ kPa} = 30.4 \text{ m}^3 = 0.03 \times 10^3 \text{ m}^3$$

**or**

$$7.90 \text{ m}^3/\text{m}^3 \times 3.85 \text{ m}^3 = 30.4 \text{ m}^3 = 0.03 \times 10^3 \text{ m}^3$$

**Step 2:** Calculate the total test gas produced for well X for this test:

$$2.33 \times 10^3 \text{ m}^3 + 0.03 \times 10^3 \text{ m}^3 = 2.36 \times 10^3 \text{ m}^3$$

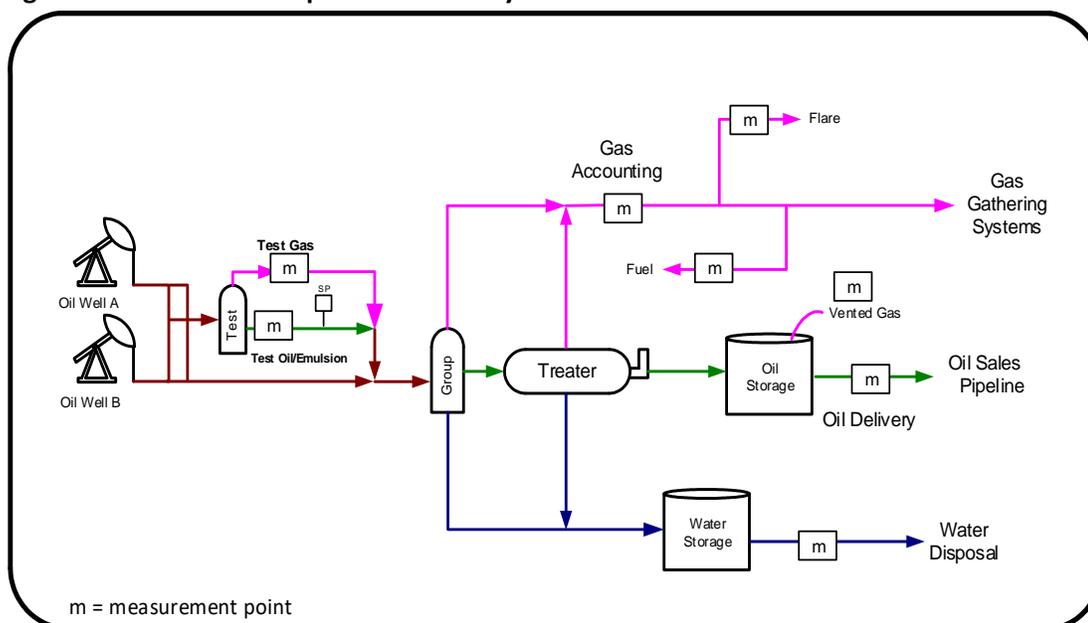
Note that test gas volumes must be determined to two decimal places (in  $10^3 \text{ m}^3$ ).

## 6.6 Oil Multi-well Proration Battery Accounting and Reporting Requirements

Prorated production is an accounting system or procedure in which the total battery production is allocated to wells based on individual well tests. Production from multiple oil wells may be commingled before separation and continuous single-phase measurement of the components (see Figure 6.4). Individual well production must be tested in accordance with Table 6.1 to determine the production rates that will be used to estimate the well's monthly production volume. The estimated monthly well production volume is adjusted using a proration factor and the adjusted volume is the volume reported in Petrinex. The following must be performed (see Section 6.5.1 for details):

1. Test production volumes of gas (in  $10^3\text{m}^3$ ) and oil and water (in  $\text{m}^3$ ) rounded to two decimal places.
2. Record test duration hours to two decimal places with the nearest quarter hour as the minimum resolution.
3. Determine the hourly production rate for each product from the well.
4. Determine the estimated well production by multiplying the hourly rate by the number of hours on production since the last test. This is done repeatedly throughout the month until the entire month's hours on production are accounted for.
5. Determine the prorated production volume by multiplying the estimated well production by the proration factor (the total battery production volume divided by the total estimated battery production volume).

Figure 6.4 Oil multi-well proration battery



The minimum test frequency and duration requirements (see Table 6.1) apply to all non-heavy oil wells under primary production and waterflood operations included in oil proration batteries.

Monitoring the performance of miscible floods and other enhanced oil recovery schemes usually requires testing criteria other than rate alone and therefore testing requirements for miscible flood schemes are set out in each scheme approval.

Licensees must monitor the classification for wells producing to a battery and meet the required testing frequency and duration for each well (see [Table 6.1](#)) unless otherwise approved by ER.

Many low-rate and stripper wells exhibit erratic production rates due to high water-oil ratios or gas-oil ratios, and oversized production lines and test separators can make accurate measurement difficult. Longer test duration can improve test accuracy for many of these wells. To allow licensees the opportunity to conduct longer duration tests, class 3 and 4 wells are allowed to use up to an eight day cycle chart drive for measurement of test gas production volumes.

The use of automatic well testing equipment and procedures with EFM provides licensees the opportunity to conduct tests of shorter durations than specified in [Table 6.1](#). The automated system can monitor the test and use statistical calculation methods to ensure that a representative rate is obtained prior to terminating the test. This practice is acceptable when:

1. the accumulated oil test volume is polled at a frequency of at least once per hour;
2. the criteria for stabilization ensures that the uncertainty for the monthly well oil volume does not exceed half of the maximum uncertainty of monthly volume stipulated in Section 1, Standards of Accuracy; and
3. the computer program is properly documented and the documentation is made available to ER upon request.

The test-to-test method, whereby data from a test are used to estimate production until the next test is conducted, must be used to estimate the production volume from each oil well based on the test rate and the total production hours. This production estimation method and the proration methodology are outlined in Sections 6.6.1 and 6.6.2. A licensee may use its own worksheet format, provided that the required data are retained and made available to ER upon request.

### 6.6.1 Proration Estimated Volume Calculation

Calculate the estimated production of each well from the test data using the sample worksheet below ([Table 6.2](#)).

1. Calculate the test rate/hour for crude oil, gas, and water:

$$\text{Rate per hour} = \frac{\text{test production volume (including GIS volumes for gas)}}{\text{test duration (hr.)}}$$

Enter the test rate per hour rounded to four decimal places.

2. Calculate the hours of production for each test rate during the reporting month. Include only the hours of prorated production:

- a. hours of production from the first day of the month to the start of the first test for the month – data from the last test conducted during the previous month will be used to estimate production until the first test for the month is conducted, and
- b. hours of production from the start of each test conducted during the month up to the start of the next test, or the end of the month, whichever is applicable.

Enter the hours produced rounded to the nearest hour.

3. Calculate the estimated production of oil, gas, and water for the production hours applicable to each test rate:

Estimated production = test rate per hour x hours produced

Enter the estimated production of oil, gas, and water rounded to one decimal place.

4. Calculate the totals for each well:

Add the hours produced that are applicable to each test rate and enter the total.

Add the estimated production of oil, gas, and water, and enter the totals.

Note that if a GOR is used to estimate the well gas production in accordance with Section **Error! Reference source not found.**:

Estimated well gas production = estimated well oil production x GOR

**Table 6.2 Proration estimated volume calculation**

UID	WI 100060100101W400					Test duration <sup>c</sup>	Hourly test rate			Prod	Estimated production		
	Test date		Test oil	Test gas	Test water		Oil	Gas	Water		Oil	Gas	Water
Vessel	dd	mm	m <sup>3</sup>	10 <sup>3</sup> m <sup>3</sup>	m <sup>3</sup>	hours	m <sup>3</sup> /hr	10 <sup>3</sup> m <sup>3</sup> /hr	m <sup>3</sup> /hr	hours	m <sup>3</sup>	10 <sup>3</sup> m <sup>3</sup>	m <sup>3</sup>
Prior mo.	25	6	9.05	1.35	3.53	24.00	0.3771	0.0563	0.1471	96	36.2	5.4	14.1
	5	7	8.85	1.28	3.26	24.00	0.3688	0.0533	0.1358	168	62.0	9.0	22.8
	12	7	9.40	1.51	2.98	24.00	0.3917	0.0629	0.1242	216	84.6	13.6	26.8
	21	7	9.15	1.67	3.65	24.00	0.3813	0.0696	0.1521	264	100.7	18.4	40.2
							<b>Totals</b>			<b>744</b>	<b>283.5</b>	<b>46.4</b>	<b>103.9</b>
UID	WI 100080100101W400					Test duration <sup>c</sup>	Hourly test rate			Prod	Estimated production		
Vessel	Test date		Test oil	Test gas	Test water		Oil	Gas	Water		Oil	Gas	Water
Vessel	dd	mm	m <sup>3</sup>	10 <sup>3</sup> m <sup>3</sup>	m <sup>3</sup>	hours	m <sup>3</sup> /hr	10 <sup>3</sup> m <sup>3</sup> /hr	m <sup>3</sup> /hr	hours	m <sup>3</sup>	10 <sup>3</sup> m <sup>3</sup>	m <sup>3</sup>
Prior mo.	28	6	5.05	0.95	4.15	24.00	0.2104	0.0396	0.1729	48	10.1	1.9	8.3
	3 <sup>a</sup>	7	5.85	1.25	4.50	48.00	0.2406	0.0490	0.1792	336	80.8	16.5	60.2
	4 <sup>a</sup>	7	5.70	1.10	4.10								
	17	7	6.01	1.15	5.00	25.50	0.2357	0.0451	0.1961	168	39.6	7.6	32.9
	24	7	5.40	0.99	4.10	22.75	0.2374	0.0435	0.1802	192	45.6	8.4	34.6
							<b>Totals</b>			<b>744</b>	<b>176.1</b>	<b>34.4</b>	<b>136.0</b>
UID	WI 100160100101W400					Test duration <sup>c</sup>	Hourly test rate			Prod	Estimated production		
Vessel	Test date		Test oil	Test gas	Test water		Oil	Gas	Water		Oil	Gas	Water
Vessel	dd	mm	m <sup>3</sup>	10 <sup>3</sup> m <sup>3</sup>	m <sup>3</sup>	hours	m <sup>3</sup> /hr	10 <sup>3</sup> m <sup>3</sup> /hr	m <sup>3</sup> /hr	hours	m <sup>3</sup>	10 <sup>3</sup> m <sup>3</sup>	m <sup>3</sup>
Prior mo.	1 <sup>b</sup>	7	1.80	1.10	2.20	24.00	0.0750	0.0458	0.0917	24	1.8	1.1	2.2
	2 <sup>b</sup>	7	4.00	2.00	5.00	24.00	0.1667	0.0833	0.2083	120	20.0	10.0	25.0
	7	7	3.95	1.95	4.95	23.00	0.1717	0.0848	0.2152	288	49.4	24.4	62.0
	19	7	4.25	2.05	5.05	26.00	0.1635	0.0788	0.1942	216	35.3	17.0	41.9
	28	7	5.65	2.00	5.50	27.75	0.2036	0.0721	0.1982	96	19.5	6.9	19.0
							<b>Totals</b>			<b>744</b>	<b>126.0</b>	<b>59.4</b>	<b>150.1</b>

Note that test gas volumes must include gas-in-solution (GIS) volumes (see Section **Error! Reference source not found.**).

<sup>a</sup> Tests on July 3 and 4 were comparable and consecutive, e.g., there were no operational changes. Therefore, the results are combined and used as one 48.0-hour test.

<sup>b</sup> Tests on July 1 and 2 were not comparable due to operational changes, e.g., choke/pump speed. Therefore, they are used as separate 24.0-hour tests.

<sup>c</sup> Test duration must be reported to the nearest quarter hour as the minimum resolution (record hours to two decimal places), e.g., 2 hr. and 45 min are entered as 2.75 hr.

## 6.6.2 Calculate Proration Factors and Monthly Production

1. Calculate the total estimated battery production for oil, gas, and water:

Total estimated battery production = sum of all the wells' total estimated production

2. Calculate the total battery production and proration factors for oil, gas, and water:

For oil and water,

Total battery production = total monthly disposition + closing inventory – opening inventory – total receipts

For gas,

Total battery production = total monthly disposition (including fuel, flare, vent) – total receipts

Proration factor = total battery production ÷ total estimated battery production

The proration factors for oil, gas, and water must be rounded to five decimal places.

If a GOR is used to estimate the total battery gas production volume in accordance with Section **Error! Reference source not found.**:

Estimated battery gas production = battery oil production x GOR

Estimated battery gas production = battery gas production

Gas proration factor = 1.00000

3. Calculate each well's monthly prorated production volumes for oil, gas, and water:

Monthly prorated oil volume = well estimated oil production x oil proration factor

Monthly prorated gas volume = well estimated gas production x gas proration factor

Monthly prorated water volume = well estimated water production x water proration factor

4. Check that total well production equals total battery production for oil, gas, and water. If the volumes are not equal due to rounding, minor adjustments to the monthly volumes may be required.

Sum of prorated well production = total battery production

## 6.7 Condensate Receipts at an Oil Battery

If condensate is received by pipeline at an oil battery, the licensee must choose from the applicable condensate reporting options in Table 5.6. The volume of condensate received from an external source that will be reported as a GEV must be subtracted from the total monthly battery gas disposition volume to determine the monthly battery gas production volume.

When condensate is received by truck at an oil battery where a portion of the condensate could flash into the gas phase, the flashed condensate must be reported as a GEV receipt volume and the unflashed condensate must be reported as a liquid condensate receipt.

Note that this may also be applicable to other light hydrocarbons delivered into an oil battery.

## 6.8 Electronic Flow Measurement for Oil Systems

See Section 14.10 for details.

## 6.9 Reporting Requirements and Scenarios for Wells Producing Oil

The following scenarios are the required reporting scenarios for both oil wells and gas wells producing oil. See Section 13 for condensate scenarios.

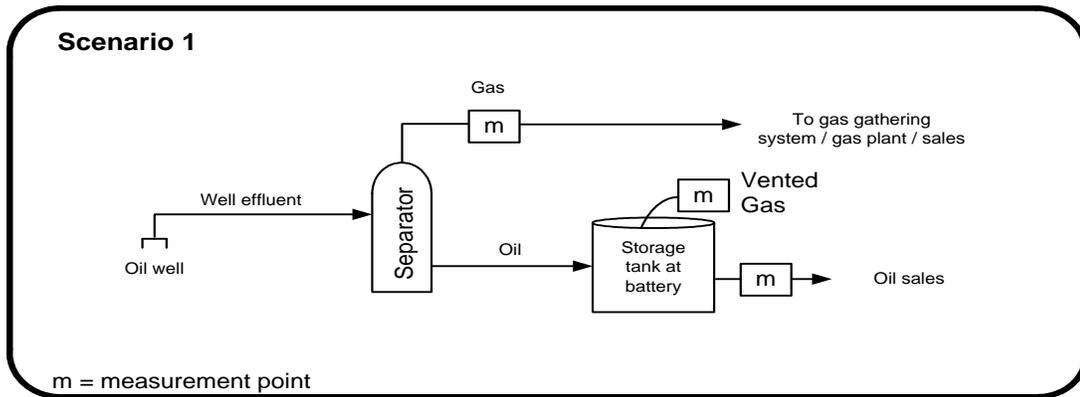
### 6.9.1 Oil Wells

#### Scenario 1

Oil separated from well effluent and sold from battery facilities.

Report as OIL PROD and OIL DISP at the battery in Petrinex.

Figure 6.5 Scenario 1

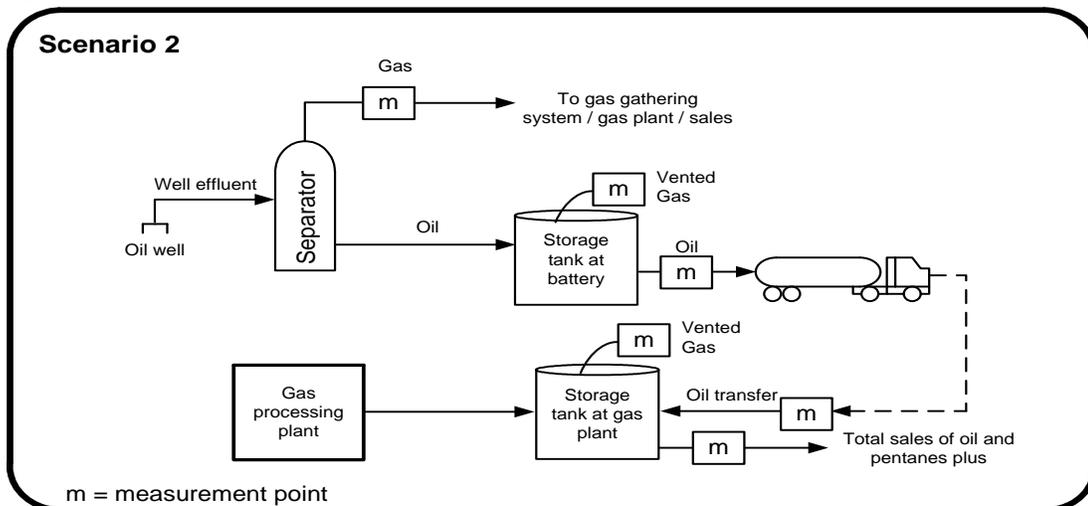


#### Scenario 2

Oil separated from well effluent, measured, and trucked to a tank at the gas plant.

Report as OIL PROD and OIL DISP at battery and OIL REC at the gas plant in Petrinex.

Figure 6.6 Scenario 2



### Scenario 3

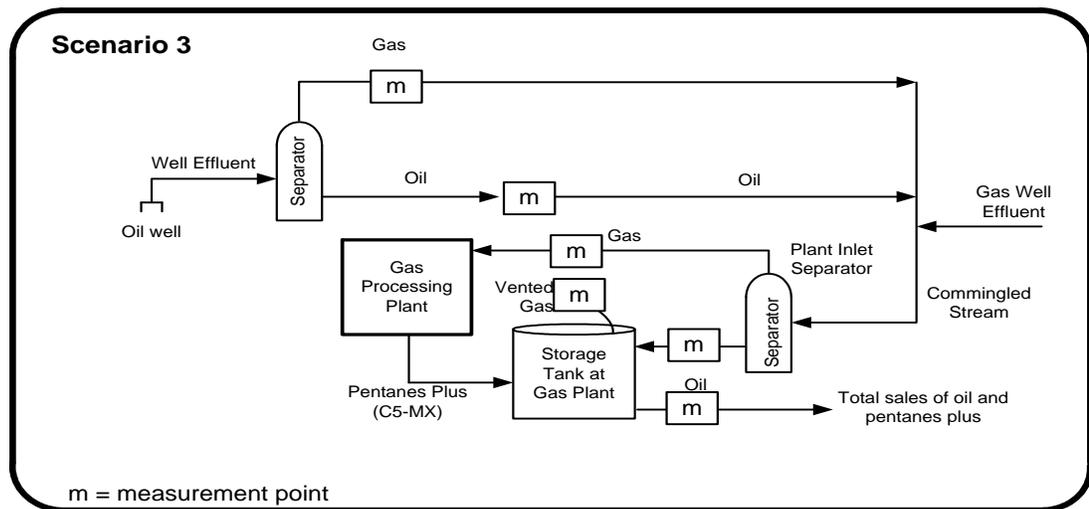
Oil separated from well effluent, measured, commingled with gas, and sent to a gas plant.

Report as OIL PROD and OIL DISP at battery and OIL REC at the gas plant in Petrinex.

Dispositions reported at the gas plant will be the total combined sales of the oil receipt and the plant pentanes plus products.

Note: The total plant inlet volumes reported would normally include the gas equivalent of the inlet condensate, but in this scenario, the inlet condensate volumes used to calculate the total plant inlet must be the net of the oil production received at the plant. The reported plant inlet volumes and the pentanes plus production will be the measured volumes minus the oil receipt.

Figure 6.7 Scenario 3

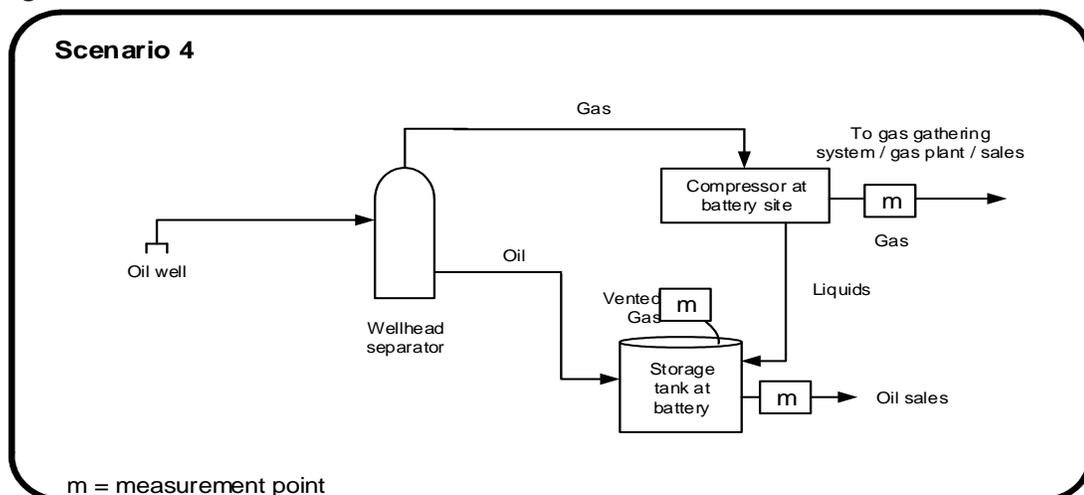


### Scenario 4

Oil separated from well effluent at battery, gas compressed as part of normal battery operations, and additional liquids recovered as a result of compression and commingled with battery oil production.

Report total fluid as OIL PROD and total OIL DISP in Petrinex.

Figure 6.8 Scenario 4

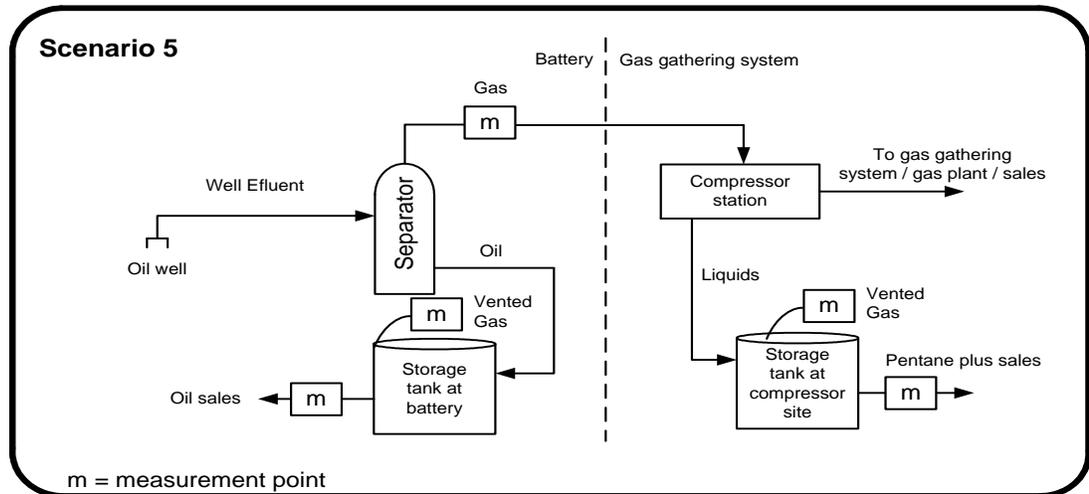


**Scenario 5**

Oil separated from well effluent at battery, gas compressed not as part of normal battery operations, and additional liquids recovered as a result of compression.

Report OIL PROD and OIL DISP at the battery in Petrinex. Hydrocarbon liquids recovered as a result of compression will be reported as pentanes plus (C5-MX) at the gathering system in Petrinex.

**Figure 6.9 Scenario 5**



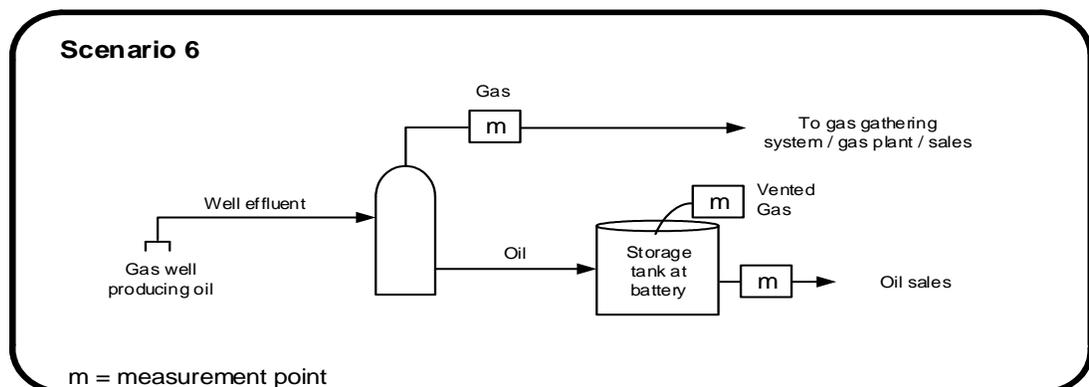
**6.9.2 Gas Well Producing Oil**

**Scenario 6**

Oil separated from well effluent, measured, and sold from battery.

Report as OIL PROD and OIL DISP at the battery in Petrinex.

**Figure 6.10 Scenario 6**



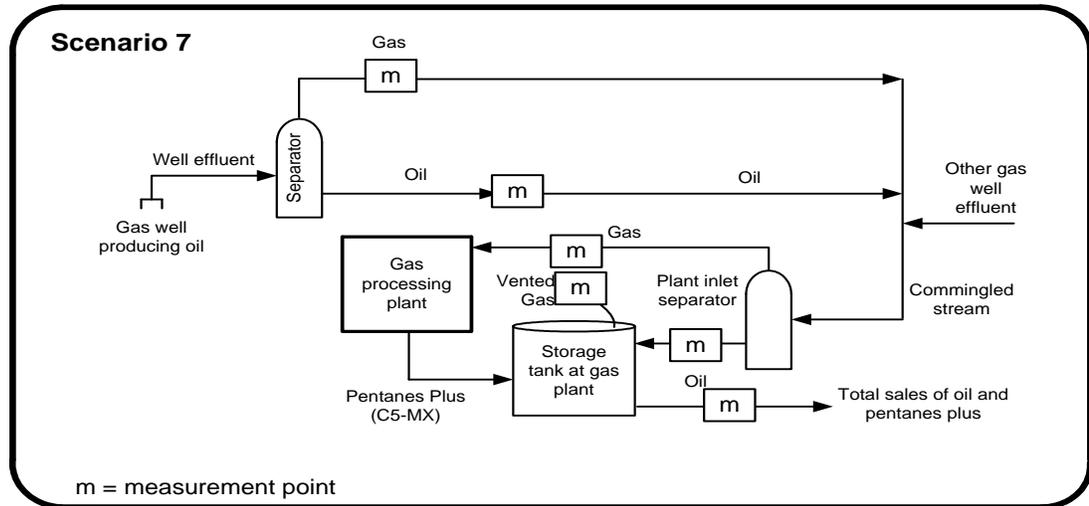
**Scenario 7**

Oil separated from well effluent, measured, commingled with gas, and sent to a gas plant.

Report as OIL PROD and OIL DISP at battery and OIL REC at the gas plant in Petrinex. Dispositions reported at the gas plant will be the total combined sales of this the oil receipt and the gas plant C5-MX products.

Note: The total plant inlet volumes reported would normally include the gas equivalent of the inlet condensate, but in this scenario, the inlet condensate volumes used to calculate the total plant inlet must be the net of the oil production received at to the plant. The reported gas plant inlet volumes and the C5-MX production will be the measured volumes minus the oil receipt.

Figure 6.11 Scenario 7

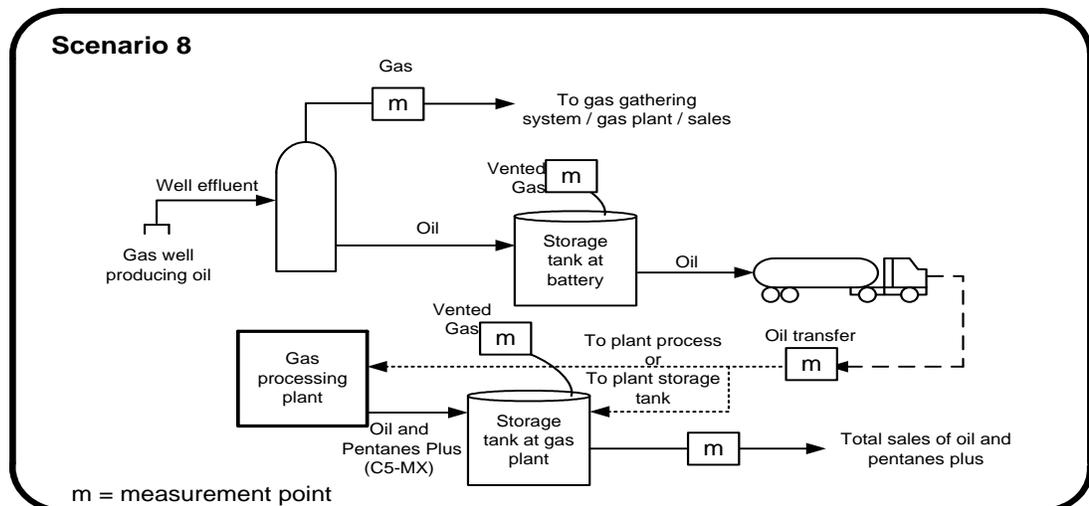


**Scenario 8**

Oil separated from well effluent, measured, and trucked to a gas plant process or a storage tank at the gas plant.

Report as OIL PROD and OIL DISP at battery and OIL REC at the gas plant in Petrinex. Dispositions reported at the gas plant will be the total combined sales receipt and the gas plant C5-MX products.

Figure 6.12 Scenario 8



## 7 Gas Proration Batteries

This section presents the requirements and exceptions relating to measurement, accounting, and reporting for gas proration batteries.

Under certain circumstances gas well licensees have the option of not measuring the gas and/or separated liquids for each well. Instead of the gas and liquids being-separated and measured, they can be prorated. Licensees that decide to install proration measurement systems in accordance with the provisions of this section are accepting higher measurement uncertainty at the wellhead, offset by lower capital and operating costs.

Prorated wells are tested periodically to determine the typical flow rate. The gas and liquids from a number of wells are measured at a group meter, and the volume at the group meter is prorated back to the individual wells based on the most recent test and the hours on production.

The measurement uncertainty realized at individual wells within gas proration batteries is greater than for wells where the gas and liquids are separated and measured. For this reason, licensees should understand the impact of this type of measurement when dealing with partners and third parties.

Prorated wells can be physically tied in to the same system as measured wells but under separate reporting facility IDs. In these scenarios, the measured well production volumes are kept whole, and the difference between the proration battery disposition volume and the measured well volume is prorated to all the proration wells. This is referred to as measurement by difference (MbD), see Section 5.5., increases the measurement uncertainty of the reported prorated well volumes because the measured well volumes are kept whole.

### 7.1 General Requirements

The three types of gas proration batteries include:

1. Gas multi-well proration SW Saskatchewan batteries (Petrinex facility subtype 363),
2. Gas multi-well proration outside SW Saskatchewan batteries (Petrinex facility subtype 364), and
3. Gas multi-well effluent measurement batteries (Petrinex facility subtype 362).

All wells in a gas proration battery must be gas wells and must be connected by flow line to a common group separation and measurement point.

All gas proration batteries require periodic well tests to be conducted to determine production rates, production ratios, and/or ECF that will be used in the determination of monthly estimated well production volumes. Monthly estimated well production volumes are multiplied by proration factors to determine the reported well production volumes. All wells must be tested annually unless otherwise stated in Section 7.

All volumetric calculations must be in  $10^3\text{m}^3$ , to the required decimal places listed in [Table 7.1](#).

**Table 7.1 Required decimal places for volumetric calculations in gas multi-well proration batteries**

Type of calculations	Number of decimals to be calculated to	Number of decimals to be rounded to
Production and estimated production	2	1
Well test gas, GEV of test condensate, test condensate, or test water	3	2
WGR, CGR, and OGR	5	4
Proration factors, ECF	6	5

Test taps must be installed at all proration gas wells. The required test tap locations are specified later in Section 7 for each of the gas multi-well proration battery types.

See Section 8 for sampling and analysis of gas, condensate, and water.

### 7.1.1 Group Measurement

Where delivery point measurement is required, the combined (group) production of all wells in the proration battery must have three-phase separation and be measured as single-phase components. Where delivery point measurement is not required, the group production may be measured using a two-phase separator with an online product analyzer on the liquid leg of the separator provided that:

1. The measurement system design meets the requirements of Section 14 and
2. The condensate and water are recombined with the gas and delivered to a gas gathering system or gas plant for further processing.

If liquid condensate is trucked out of the group separation and measurement point to a gas plant for further processing the condensate must be reported as a liquid condensate disposition from the battery.

Gas wells in any one of the three types of proration batteries must not be commingled with:

1. Measured gas sources or gas from another proration battery prior to group measurement.
2. Gas wells in a different type of gas proration battery, upstream of their respective group measurement points.

Variances from this requirement may be allowed if the Exception criteria in Sections 5.5 and 5.5.1 are met or if site-specific approval has been obtained from ER prior to implementation.

### 7.1.2 Stabilized Flow and Representative Flow

The words stabilized flow and representative flow are used extensively in this section and are defined as:

**Stabilized flow** Indicates a point at which flowing parameters of gas, condensate, or water are producing under normal operating conditions and represent production levels equal to the well's normal average flow rate. Stabilized flow can only be achieved when all testing equipment parameters associated with determining an actual volume has reached equilibrium (i.e., liquid levels in the test separator and

separator pressure and temperature stabilization to normal operating conditions).

**Representative flow**

Used when stabilized flow is not achievable, such as for wells with artificial lift systems and wells with slugging characteristics. The test volumes of gas, condensate, or water must be representative of the well's production capability under normal operating conditions

Wells that use artificial lift systems or characteristically display slug flow are not in stabilized flow and thus must be tested for a minimum duration that completes multiple flow cycles to accurately determine a representative volume of gas, condensate, or water. These representative production volumes are then extrapolated to reflect the wells' production over an extended period of time.

**7.2 Gas Multi-well Proration SW Saskatchewan Batteries (Petrinex facility subtype: 363)**

Gas wells in this type of battery do not require dedicated continuous measurement for each well or a site-specific approval from ER.

Production rates determined during a well test must be used in the proration calculations within 30 days of the test until the next test is conducted.

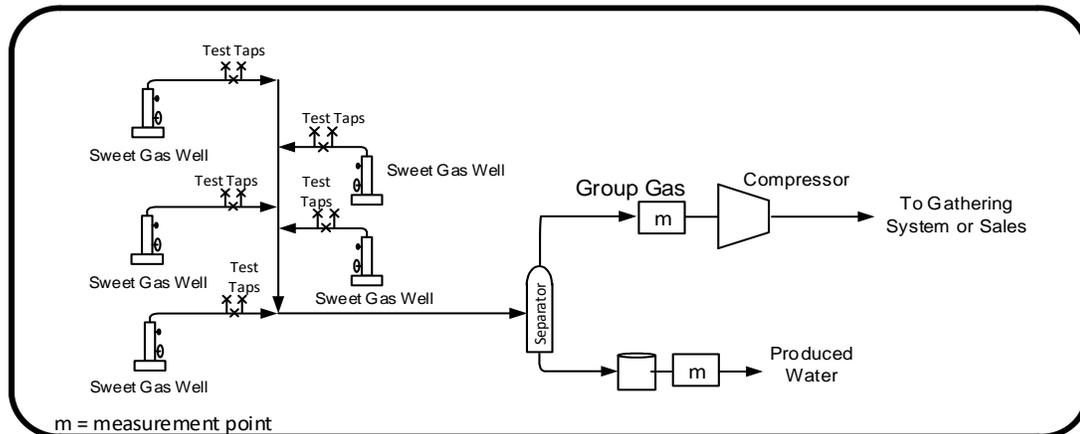
Total battery gas production must be measured and prorated back to the individual wells, based on each well's estimated monthly gas production. Estimated well gas production is based on hourly production rates, determined by periodic well tests and monthly producing hours.

Gas wells that produce from shallow gas stratigraphic units in SW Saskatchewan may be included in these types of batteries. The stratigraphic units include coals and shales from the base of the Glacial Drift to the base of the Upper Cretaceous. The production from two or more of these stratigraphic units without segregation in the wellbore requires approval for commingled production from ER prior to implementation.

**7.2.1 Group Measurement**

Group production is measured by separating the comingled well effluent and measuring each stream in a single phase. At minimum, a two-phase separator is required at the group measurement point because the battery water production must be reported at the battery level. The group measurement point is usually located at the battery site where a compressor is present (see [Figure 7.1](#)).

Figure 7.1 Typical gas multi-well proration SW Saskatchewan battery



### 7.2.2 Size of a Gas Multi-well Proration SW Saskatchewan Battery

There is no limit on the number of flowlined wells that may be in a Gas Multi-well Proration SW Saskatchewan Battery. However, licensees are encouraged to consider the logistics of the battery's operation in determining the size of these batteries, with the key factors being:

1. The ability to conduct representative well tests at the minimum frequency specified in Table 7.2; and
2. The configuration and operating pressures of the battery and flow lines such that all wells can readily flow.

This approach will generally result in the main pipeline system laterals being used to establish a group measurement point.

### 7.2.3 Testing Requirements

Gas production rate tests must be conducted for each well in the battery in accordance with the following requirements:

1. The test must be of sufficient duration to clearly establish a stabilized flow rate.
2. The test must be representative of the well's capability under normal operating conditions.
3. programs and procedures must ensure that all wells are treated equitably within their respective batteries. These types of wells are typically tested by directing flow from the well through a dry flow test meter. However, a test separator system may also be used.
4. New wells must be tested within the first 30 days of production, then again within 12 months, and thereafter according to Table 7.2.

If these requirements cannot be satisfied, the licensee must either reconfigure the system, e.g., redirect some wells to another battery/group measurement point, or test each of the individual wells within the battery once per month.

**Table 7.2 Testing frequency for SW Saskatchewan shallow gas wells**

Minimum rate	Maximum rate	Number of tests	Frequency*
	$\leq 0.5 \text{ } 10^3\text{m}^3/\text{d}$	1	Triennial
$> 0.5 \text{ } 10^3\text{m}^3/\text{d}$	$\leq 5.0 \text{ } 10^3\text{m}^3/\text{d}$	1	Biennial
$> 5.0 \text{ } 10^3\text{m}^3/\text{d}$		1	Annual

\*See Appendix 2 for frequency definition.

## 7.2.4 Production Accounting and Reporting Procedures

### 7.2.4.1 Water Reporting Requirements

The reporting of water production for the wells in Gas Multi-well Proration SW Saskatchewan Batteries is not required. However, all water receipts and dispositions must be reported at the battery level. A receipt from a SKMC for Saskatchewan can be used to balance the disposition at the battery level on Petrinex. If the water is trucked to non-Petrinex reporting facilities or evaporated on site, it must be reported as a disposition to SKMC.

### 7.2.4.2 Gas Production Volume Calculations

The following example assumes there are no measured receipts at the battery upstream of the group measurement point. Monthly gas production volumes are to be calculated as follows:

1. Calculate well gas test rate:

$$\text{Well gas test rate (} 10^3\text{m}^3/\text{hour)} = \text{Well test gas volume (} 10^3\text{m}^3) \div \text{Well test duration (hours)}$$

2. Calculate estimated monthly well gas volume:

$$\text{Estimated monthly well gas volume} = \text{Well gas test rate} \times \text{Monthly total hours of well production}$$

3. Calculate total estimated gas production for the battery:

$$\text{Total battery estimated monthly gas volume} = \text{Sum of all estimated monthly well gas volumes}$$

4. Calculate proration factor for gas:

$$\text{Gas proration factor} = \frac{\text{Total battery measured monthly gas volume}}{\text{Total battery estimated monthly gas volume}}$$

5. Calculate reported monthly (prorated) well gas production:

$$\text{Reported monthly well gas production} = \text{Gas proration factor} \times \text{Estimated monthly well gas volume.}$$

## 7.3 Gas Multi-well Proration Outside SW Saskatchewan Batteries (Petrinex facility subtype 364)

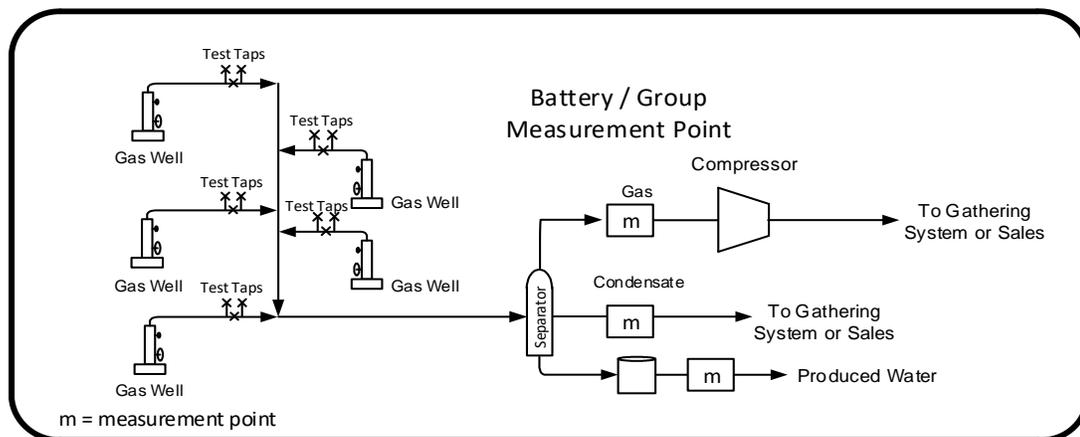
Gas wells in this type of battery do not require dedicated continuous measurement for each well. Production rates, WGRs, and/or CGRs determined during a well test must be used within 30 days in the proration calculations until the next test is conducted.

Total battery gas production must be measured and prorated back to the individual wells based on each well's estimated monthly gas production. Estimated well gas production is based on hourly production rates, determined by well tests, at the frequency listed in Section 7.3.1 and monthly producing hours.

Total battery condensate production must be measured, if present, and it must be prorated back to the individual wells based on each well's CGR from the production tests if the condensate is separated and delivered from the battery. In this scenario the estimated gas production volume at each well will not include the GEV of the condensate. If the condensate is recombined with the gas and delivered to a gathering system or to a gas plant for further processing, the condensate must be reported as a GEV and added to the measured gas production volume and reported on Petrinex.

Total battery water production must be measured and prorated back to the individual wells based on each well's estimated monthly water production. Estimated well water production is based on a WGR, determined by well tests multiplied by the estimated monthly well gas production (see Figure 7.2).

**Figure 7.2 Typical gas multi-well proration outside SW Saskatchewan battery**



If total water production at each well in the battery is  $\leq 0.5 \text{ m}^3/\text{day}$  based on the monthly average flow rates recorded during the six months prior to conversion, water production may be prorated to all wells in the battery based on the estimated gas production at each well. In this scenario, each well's percentage of the total estimated well gas production is multiplied by the battery group measured water production to determine each well's water production. If a group of new wells not previously on production is to be configured as a proration battery, the qualifying flow rates must be based on production tests conducted under the normal operating conditions of the proration battery.

There is no geographical or zonal limitation for this type of proration battery. The exception criteria in Section 5.4 must be met or ER site-specific approval must be obtained prior to the proration battery implementation either at the initial design and installation stage or at a later stage of production when the production rate decreases to a point that continuous measurement is not economical.

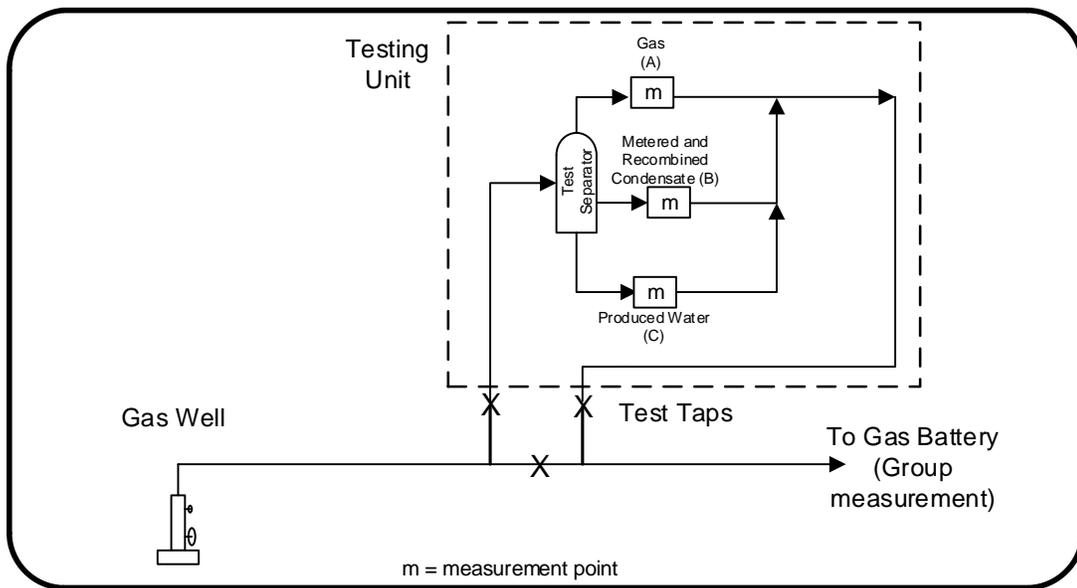
Gas wells producing oil rather than condensate must not be physically tied into a Gas Multi-well Proration Outside SW Saskatchewan battery unless the well oil and gas production volumes are separated and measured prior to commingling with the other wells in the battery and either the Exception criteria in Sections 5.5 and 5.5.1 are met or site-specific approval has been obtained from ER prior to implementation. This is a measurement by difference scenario so a separate Petrinex reporting facility ID must be set up for the gas well producing oil. However, if a gas well classified as producing condensate in a gas Multi-well Proration Outside SW Saskatchewan battery is reclassified by ER as producing oil, the well may remain linked to the battery for reporting purposes provided that the well is equipped with a separator and there is continuous

measurement of the gas, oil, and water or, alternatively, the measurement, accounting, reporting procedures specified in Section 7.3.3 are followed.

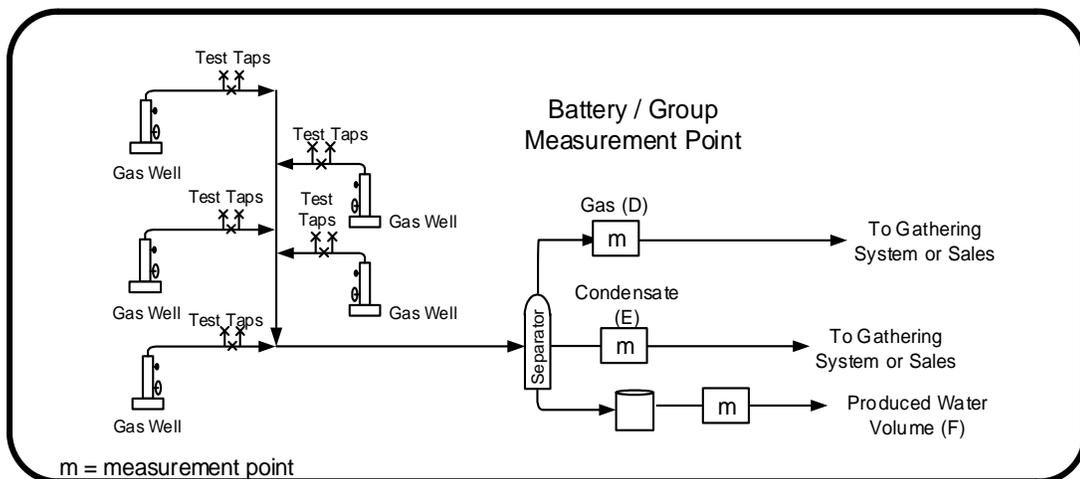
### 7.3.1 Well Testing Requirements

Well testing is typically performed by directing well production through a three-phase portable test separator configured with dedicated meters for gas, condensate, and water. A two-phase separator with a total liquid meter and continuous water cut analyzer is also, acceptable (see Figures 7.3 and 7.4).

**Figure 7.3 Typical testing unit for a gas multi-well proration outside SW Saskatchewan battery**



**Figure 7.4 Typical gas multi-well proration outside SW Saskatchewan battery**



Well tests must be conducted with measurement of all phases as follows:

1. The test must begin only after liquid levels have stabilized.

2. The test duration must be a minimum of 12.0 hours.
3. After the commencement of production at the proration battery, all wells must be tested within the first month, and annually thereafter. New wells added to the battery must be tested within the first month of production, then again within six months, and annually thereafter.
4. Consistent testing procedures must be used for consecutive tests to identify if a change in a well's flow characteristics has occurred.
5. These wells are typically tested by directing flow from the well through a test separator. If the initial testing with a separator shows a liquid-gas ratio (LGR)  $< 0.01 \text{ m}^3 \text{ liquid}/10^3 \text{ m}^3 \text{ gas}$ , other testing methodologies, such as a smaller separator or a dry flow test meter without separation, could be used for the next test. If the total liquid volumes at group measurement point  $> 0.05 \text{ m}^3 \text{ liquid}/10^3 \text{ m}^3 \text{ gas}$  in any month, a test separator must be used to test all the wells within the battery for the next round of testing to determine where the liquid originated.
6. The gas, condensate, and water volumes must be measured.
7. The condensate must be sampled during every test and subjected to a compositional analysis, which is to be used to determine the gas equivalent factor (GEF). The sample may be taken from the condensate leg of a three-phase separator or the liquid leg of a two-phase separator. The water must be removed from the condensate before conducting the analysis.
8. The GEF must be used to convert the liquid condensate volume determined during the test to a GEV, which will be added to the measured test gas volume to determine the total test gas volume if the condensate is recombined with the gas and flowlined to a gas plant or other facility for further processing, see Section 7.3.2.
9. The WGR, CGR, and OGR (if applicable) must be determined by dividing the test water, condensate, and oil volume respectively by the total test gas volume.
10. For orifice meters, the test gas meter must use 24-hour charts for a test period of 72 hours or less, unless electronic flow measurement is used; for testing periods longer than 72 hours, seven day charts may be used, provided that good, readable pen traces are maintained, see Section 4.3.2.

The test frequency may be extended with ER approval.

#### **7.3.1.1 Exception for Gas Multi-well Proration Outside SW Saskatchewan Batteries**

1. New and existing wells producing from shallow gas stratigraphic units in SW Saskatchewan may be tested in accordance with the testing requirements set out in Section 7.2.3.
2. Existing shallow gas wells in batteries located outside the SW Saskatchewan shallow gas stratigraphic unit with a  $\text{LGR} \leq 0.01 \text{ m}^3 \text{ liquid}/10^3 \text{ m}^3 \text{ gas}$  may be tested in accordance with the testing requirements set out in Section 7.2.3.

The stratigraphic units include coals and shales from the base of the Glacial Drift to the base of the Upper Cretaceous. The production from two or more of these stratigraphic units without segregation in the wellbore requires prior approval from ER for commingled production.

### 7.3.2 Production Volume Calculations

The following example assumes there are no measured receipts at the battery upstream of the group measurement point. Monthly production volumes are to be calculated as follows:

1. Calculate well gas test rate, see [Figure 7.3](#):

$$\text{Well gas test rate (10}^3\text{m}^3\text{/hour)} = (\text{Well test gas volume [A]} + \text{GEV of well test condensate [B]}) \div \text{Well test hours}$$

Note: Do not include GEV of [B] if condensate is delivered for sale at the group measurement point or trucked to a gas plant or other facility for further processing.

2. Calculate estimated monthly well gas volume:

$$\text{Estimated monthly well gas volume} = \text{Well gas test rate} \times \text{Monthly total hours of well production}$$

3. Calculate total estimated gas production for the battery:

$$\text{Total battery estimated monthly gas volume} = \text{Sum of all estimated monthly well gas volumes}$$

4. Calculate the well WGR, see [Figure 7.3](#):

$$\text{WGR} = \text{Well test water volume (C)} \div (\text{Well test gas volume [A]} + \text{GEV of well test condensate [B]})$$

Note: Do not include GEV of [B] if condensate is delivered for sale at the group measurement point or trucked to a gas plant or other facility for further processing.

5. Calculate estimated water production for each well:

$$\text{Estimated monthly well water volume} = \text{Estimated monthly well gas volume} \times \text{WGR}$$

6. Calculate total estimated water production for the battery:

$$\text{Total battery estimated monthly water volume} = \text{Sum of all estimated monthly well water volumes}$$

If the condensate is delivered for sale at the group measurement point or trucked to a gas plant or other facility for further processing, calculate the next two items; otherwise go directly to item 9.

7. Calculate the well CGR, see [Figure 7.3](#):

$$\text{CGR} = \text{Well test condensate volume (B)} \div \text{Well test gas volume (A)}$$

8. Calculate estimated condensate production for each well:

$$\text{Estimated monthly well condensate volume} = \text{Estimated monthly well gas volume} \times \text{CGR}$$

9. Calculate total estimated condensate production for the battery:

$$\text{Total battery estimated monthly condensate volume} = \text{Sum of all estimated monthly well condensate volumes}$$

10. Calculate proration factors for gas, condensate (if the condensate is recombined with the gas and flowlined to a gas plant for further processing) and water, see [Figure 7.4](#):

$$\text{Gas Proration Factor (GPF)} = (\text{Total battery measured monthly gas volume [D]} + \text{GEV of total battery condensate [E]}) \div \text{Total battery estimated monthly gas volume}$$

Note: Do not include GEV of [E] if condensate is delivered for sale at the group measurement point or trucked to a gas plant or other facility for further processing.

Water Proration Factor (WPF) = Total battery actual monthly water volume (F) ÷ Total battery estimated monthly water volume

Condensate Proration Factor (CPF) = Total battery measured monthly condensate volume [E] ÷ Total battery estimated monthly condensate volume

11. Calculate reported monthly (prorated) well production:

Reported monthly well gas production = Estimated monthly well gas volume x Gas Proration Factor

Reported monthly well water production = Estimated monthly well water volume x Water Proration Factor

Reported monthly well condensate production = Estimated monthly well condensate volume x Condensate Proration Factor

### 7.3.3 Exception for Gas Wells Producing Oil

If the hydrocarbon liquid that a gas well produces is reclassified by ER from condensate to oil, the well may remain linked, for reporting purposes, in a Gas Multi-well Proration Outside SW Saskatchewan Battery, provided that the well is equipped with a separator and there is continuous measurement of the gas and liquid components or, alternatively, the measurement, accounting, and reporting procedures specified below are followed, (see [Figure 7.5](#)).

Annual Gas Rate–WGR tests must be conducted on the well. An oil-gas ratio (OGR) must also be determined during this test. The WGR, estimated water production, water proration factor, and actual water production are determined in the same manner, see Section 7.3.2.

Units: All gas volumes and GEV are to be in 10<sup>3</sup>m<sup>3</sup> and liquid volumes in m<sup>3</sup>.

1. Calculate well gas test rate, see [Figure 7.6](#):

Well gas test rate = Well test gas volume (A) ÷ Well test hours

2. Calculate estimated monthly well gas volume:

Estimated monthly well gas volume = Well gas test rate x Monthly total hours of well production

3. Calculate the OGR, see [Figure 7.6](#):

OGR = Well test oil volume (B) ÷ Well test gas volume (A)

4. Calculate reported well oil production:

Reported monthly well oil production = Estimated monthly well gas volume x OGR

5. Calculate reported total oil production:

Reported monthly total battery oil production = Sum of all actual monthly well oil volumes

6. At the group measurement point, subtract the oil production volume (item 5) from the total liquid hydrocarbon volume to determine the total battery condensate production. The GEV of the total battery condensate volume, if not delivered for sale or trucked to a gas plant or other facility for further processing, must be added to the measured group gas volume to determine the total battery gas volume, see [Figure 7.5](#):

Total battery condensate volume = Battery total liquid hydrocarbon volume (E) – Reported monthly total battery oil production

7. Calculate total estimated gas production for the battery:

Total battery estimated monthly gas volume = Sum of all estimated monthly well gas volumes

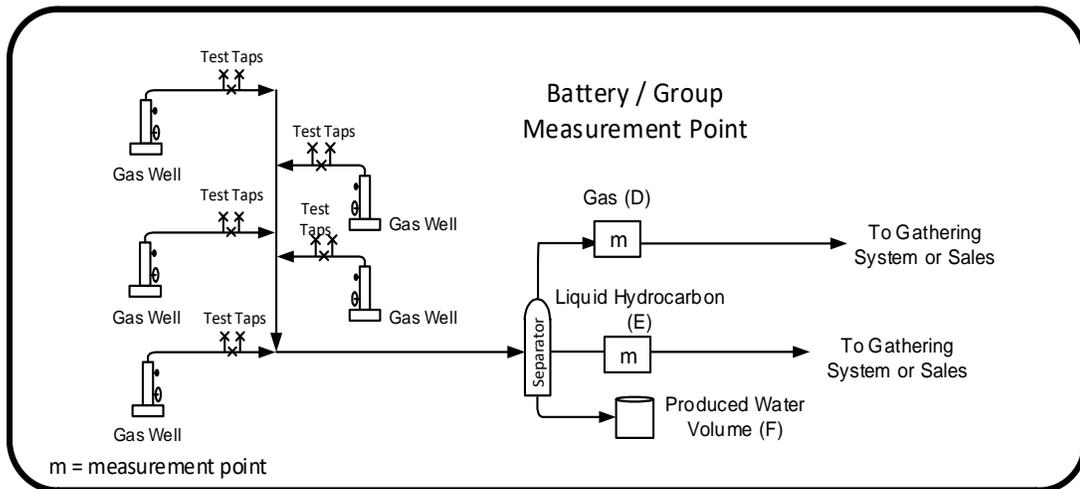
8. Calculate proration factor for gas, see [Figure 7.5](#):

Gas Proration Factor (GPF) = (Total battery measured monthly gas volume [D] + GEV of total battery condensate volume [item 6]) ÷ Total battery estimated monthly gas volume

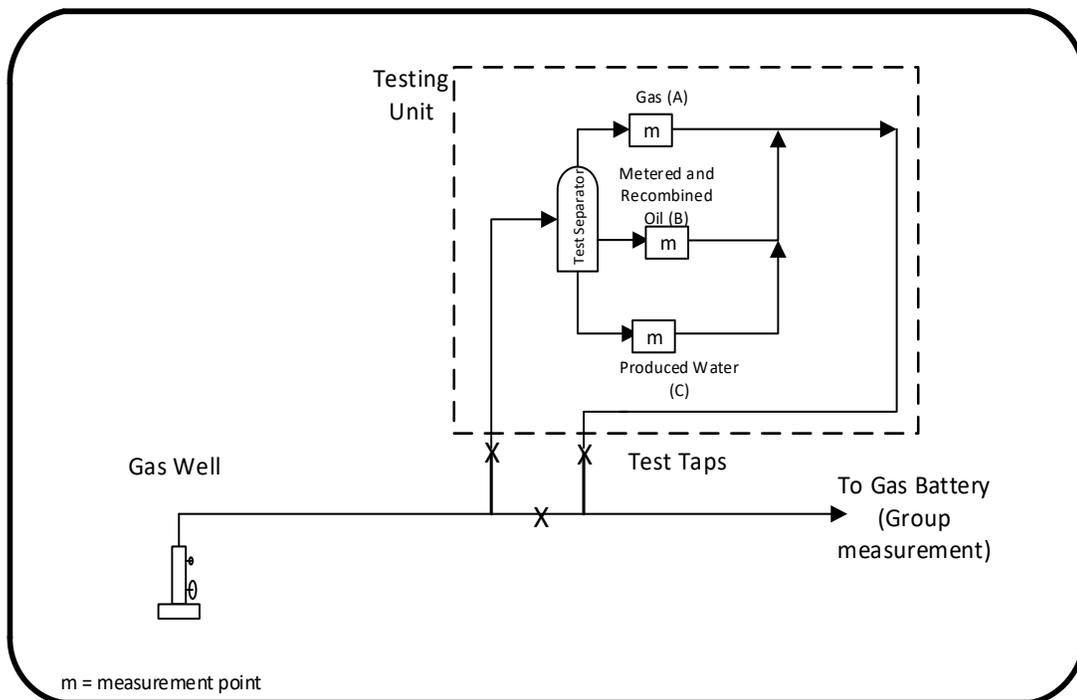
9. Calculate reported monthly (prorated) well gas production:

Reported monthly well gas production = Estimated monthly well gas volume x Gas Proration Factor

**Figure 7.5 Typical gas multi-well proration outside SW Saskatchewan battery**



**Figure 7.6 Typical testing setup**

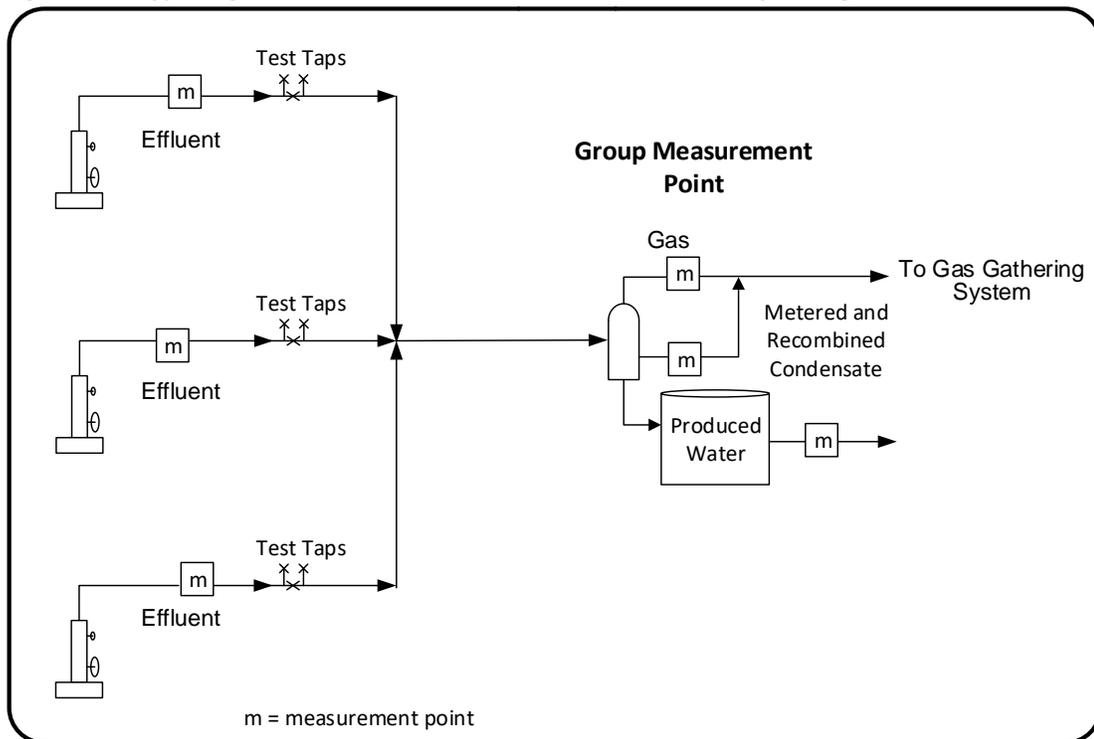


Report the calculated monthly oil production volume as oil produced from the well. Prorate monthly condensate production, if delivered for sale or trucked to a gas plant or other facility for further processing, and water production as in the normal proration battery in Section 7.3.2.

#### 7.4 Gas Multi-well Effluent Measurement Batteries (Petrinex subtype 362)

Gas wells in this type of proration battery have dedicated effluent measurement, whereby total multiphase well fluid passes through a single meter, see Figure 7.7. This type of measurement must be subjected to testing regardless of the type of effluent meter used. For a new completion or recompletion of another stratigraphic unit or zone in an existing well, effluent measurement is not allowed at a certain LGR level, see Section 7.4.1.1 for details.

**Figure 7.7 Typical gas multi-well effluent measurement battery configuration**



Where delivery point measurement is required at the group measurement point, the combined (group) production of all wells in the effluent measurement battery must have three-phase separation and be measured as single-phase components. Where delivery point measurement is not required at the group measurement point, the group production may be measured using “two-phase separation with three-phase measurement”. This means that a two-phase separator with an online product analyzer on the liquid leg of the separator may be used provided that:

1. The measurement system design meets the requirements of Section 14, Figure 14.1
2. The condensate and water are recombined and delivered to a gas gathering system or gas plant for further processing.

The resulting total battery gas production volume (including gas equivalent volume [GEV] of condensate) and total battery water production volume must be prorated back to the wells to determine each well’s reported gas and water production.

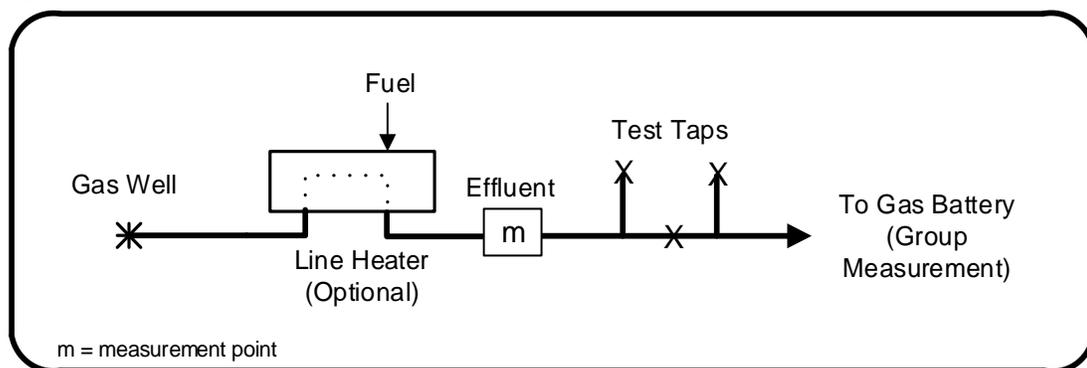
1. If condensate is trucked out of the group separation and measurement point without further processing to a sales point, condensate production must be reported at the wellhead based on the condensate-gas ratio (CGR) from the well test.
2. If liquid condensate is trucked to a gas plant or other facility for further processing the condensate must be reported as a liquid condensate volume.

Production rates, WGR, CGR, and ECF determined during a well test must be used in the estimation/proration calculations within 60 days of the test until the next test is conducted.

Total battery gas production must be measured and prorated back to the individual wells, based on each well's estimated monthly gas production. Estimated well gas production is based on the total volume measured by the effluent meter multiplied by an ECF, see [Figure 7.8](#). The uncertainty of this type of measurement will increase with higher liquid rates, especially under liquid slugging conditions.

[Figure 7.8](#) illustrates a typical gas well effluent metering configuration. Production from the gas well passes through a line heater (optional), where it is heated. This is typically done to vaporize some of the hydrocarbon liquids and heat up the water and the gas in the stream before metering to prevent hydrate formation. For well testing purposes, test taps must be located downstream of the meter within the same pipe run. The line heater, fuel gas tap, and other equipment, if present, must be upstream of the meter or downstream of the test taps to ensure that the test meter is subjected to the same condition as the effluent meter. After measurement, production from the well is commingled with other flowlined effluent measured gas wells in the battery and sent to a group (battery) location, where single-phase (group) measurements of hydrocarbon liquids, gas, and water must be conducted downstream of separation.

**Figure 7.8 Typical gas well effluent metering configuration**



For most wells, the required minimum well testing frequency is annual unless the criteria in [Section 7.4.1.1](#) are met. Total battery water production must be measured and prorated back to the individual wells, based on each well's estimated monthly water production. Estimated well water production is based on a WGR, determined by periodic well tests multiplied by the estimated monthly well gas production. Gas wells that produce oil, rather than condensate, must not be physically tied into a Gas Multi-well Effluent Measurement Battery, unless the well oil and gas production volumes are separated and measured prior to commingling with the

effluent measured wells and either the Exception criteria in Sections 5.5 and 5.5.1 are met or site-specific approval has been obtained from ER prior to implementation. This is a measurement by difference scenario so a separate Petrinex reporting facility ID must set up for the gas well producing oil.

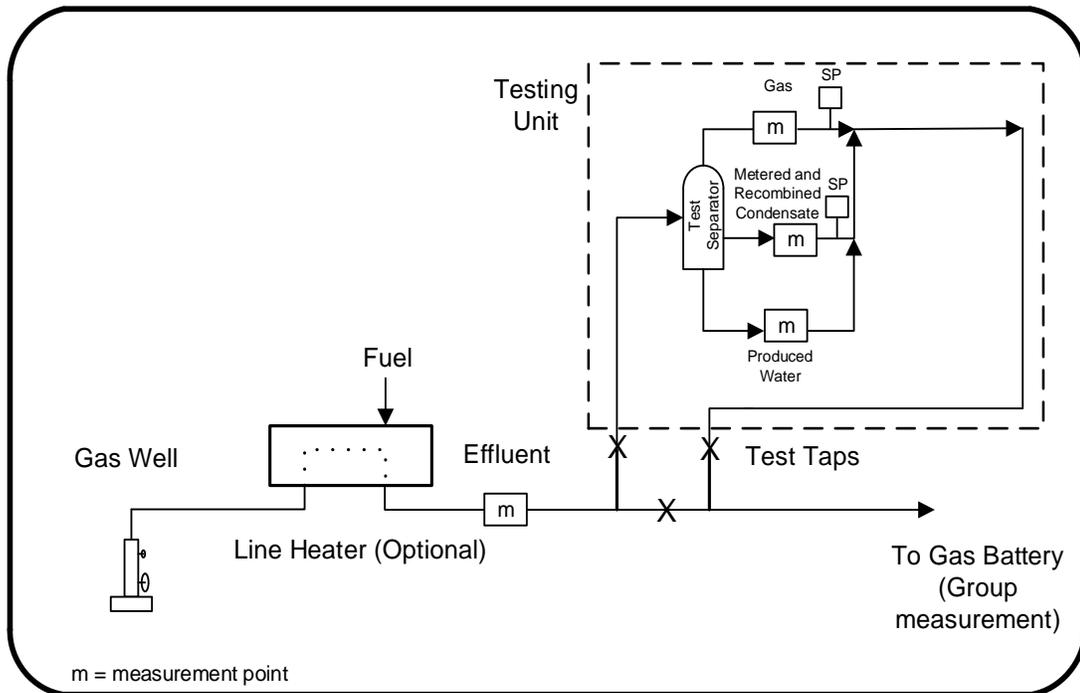
If the hydrocarbon liquid that a gas well produces is reclassified by ER from condensate to oil see Section 7.4.3.

### 7.4.1 Well Testing

Well testing is typically performed by directing well production downstream of the effluent meter and within the same pipe run through a three-phase portable test separator configured with dedicated meters for gas, condensate, and water, see [Figure 7.9](#). Two-phase separator with a total liquid meter and continuous water cut analyzer is also acceptable. The test must be conducted as follows:

1. The test must begin only after liquid levels have stabilized within the test separator.
2. The test duration must be a minimum of 12 hours.
3. All new wells must be tested within the first 30 days of initial production.
4. Consistent testing procedures must be used for consecutive tests to identify if a change in a well's flow characteristics has occurred.
5. The gas, condensate, and water volumes must be measured.
6. The condensate must be sampled during every test and subjected to a compositional analysis, which is to be used to determine the GEF. The sample may be taken from the condensate leg of a three-phase separator or the liquid leg of a two-phase separator. The water must be removed from the condensate before conducting the analysis.
7. The GEF must be used to convert the liquid condensate volume determined during the test to a GEV, which will be added to the measured test gas volume to determine the total test gas volume if the condensate is not delivered for sale at the group measurement point or trucked to a gas plant or other facility for further processing. The ECF can then be determined based on whether the condensate is recombined with the gas, see Section 7.4.2.
8. The WGR must be determined by dividing the test water volume by the sum of the measured test gas volume and the gas equivalent of the measured test condensate volume if the condensate is not delivered for sale at the group measurement point or trucked to a gas plant or other facility for further processing, see Section 7.4.2.
9. For orifice meters, the effluent meter and the test gas meter must use 24-hour charts for a test period of 24 hours or less, unless electronic flow measurement (EFM) is used; for testing periods longer than 24 hours, seven-day charts may be used, provided that good, readable pen traces are maintained, see [Section 4.3.4](#).

Figure 7.9 Typical effluent well measurement configuration with test unit

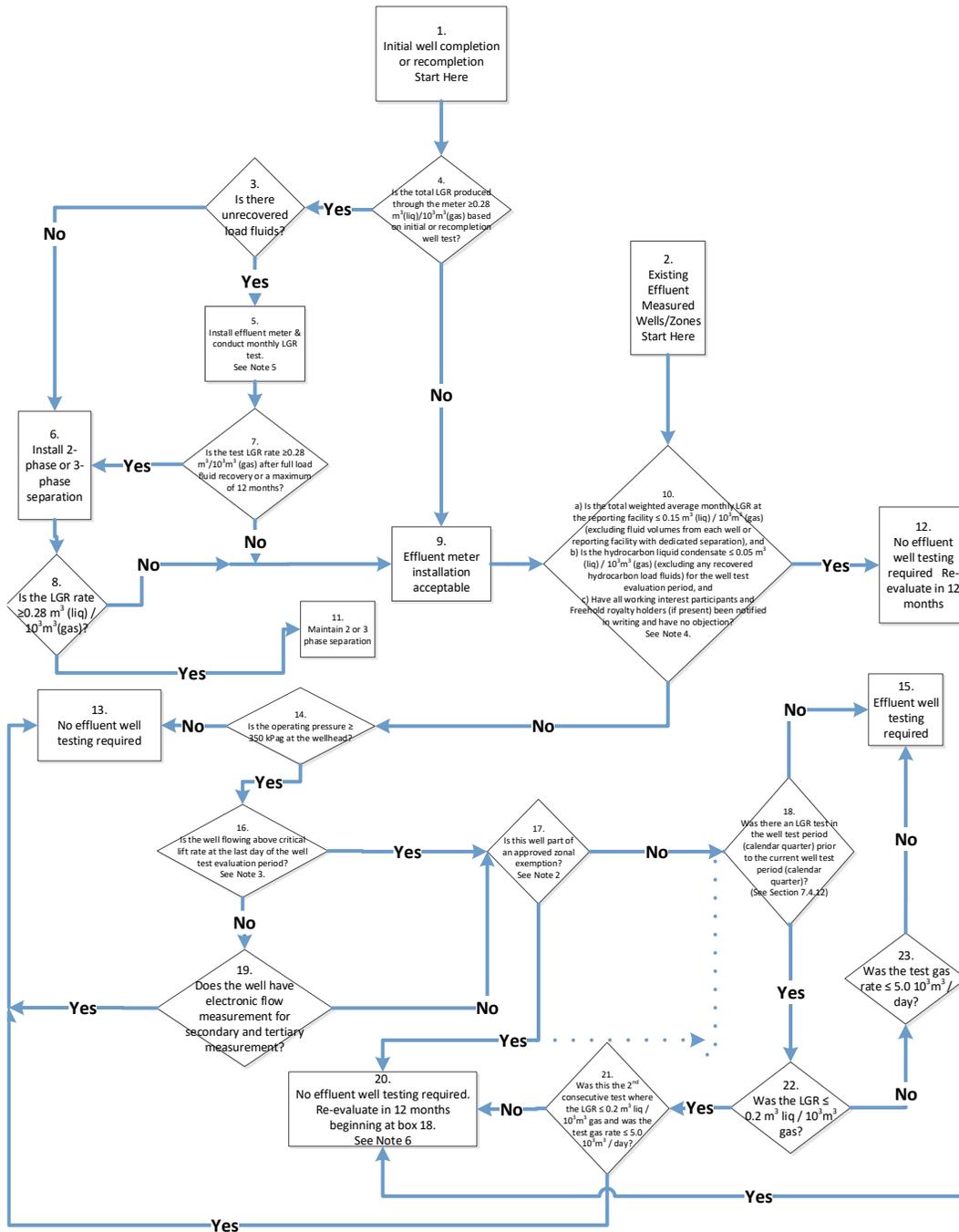


7.4.1.1 Effluent Well Measurement and Testing Decision Tree

The type of measurement and testing frequency for effluent measured wells must follow the decision tree process in [Figure 7.10](#).

Note that the starting point for initial well completion or recompletion is different than for existing effluent measured stratigraphic units or zones or wells.

**Figure 7.10 Effluent well measurement and testing decision tree. (text boxes are numbered from left to right.)**



**Note 1:** Where all wells in a facility are above critical lift and in a deemed exempt stratigraphic unit or zone, if the LGR is > 0.2 m<sup>3</sup> (liq) / 10<sup>3</sup>m<sup>3</sup> (gas) at the respective facility inlet to which the wells flow, the stratigraphic unit or zone is not exempt and the Note 1 path is to be followed.

**Note 2:** ER stratigraphic unit or zonal measurement exemption must be obtained by approval of an application only.

**Note 3:** The Turner Correlation<sup>2</sup> is used to approximate critical lift flow rate. The calculation below produces a value in million standard cubic feet (mmscf) per day. Use a factor of 28.3168 10<sup>3</sup>m<sup>3</sup>/mmscf to convert to metric units. Although there have been further refinements to the Turner Correlation calculation, the formulas below will be applied to determine critical lift flow rate as it relates to the well measurement and testing decision tree. These simplified formulas assume a fixed-gas gravity (G) of 0.6 and fixed-gas temperature (T) of 120.0°F.

$$v_g(\text{Water}) = \frac{5.62(67 - kP)^{0.25}}{(kP)^{0.50}}$$

$$q_g = \frac{3.06Pv_g A}{ZT}$$

$$v_g(\text{Condensate}) = \frac{4.02(45 - kP)^{0.25}}{(kP)^{0.50}}$$

G = gas gravity  
 P = Pressure (absolute) - lb force / square inch  
 T = Temperature (absolute) – degrees Rankine  
 v<sub>g</sub> = Minimum gas velocity required to lift liquids – ft / second  
 Z = Compressibility factor  
 A = Cross sectional area of flow – square feet  
 q<sub>g</sub> = Flow rate – mmscf / day

$$k = \frac{2.693G}{ZT}$$

The following represents a sample Turner Correlation calculation:

Evaluation period: November–October

SCADA daily average tubing pressure, October: 684.6 kPa

Turner Correlation formula: assumes G = 0.6

T = 48.9°C (120°F)

Z = 0.9

Variable	Value	Units	Calculation
G	0.6		
Z	0.9		
T	580	Rankin	[(48.9 x 1.8) + 32] + 460
k	0.003095		(2.693 x 0.6) ÷ (0.9 x 580)
P	114	PSIA	(684.6 kPa ÷ 6.89475) + (101.325 kPa ÷ 6.89475)
A	0.0217	ft <sup>2</sup>	[3.1415 x (1.995 inches ÷ 12) <sup>2</sup> ] ÷ 4 (tubing size = 2 3/8 inches)
q <sub>g</sub>	0.392	mmscf/d	
q <sub>g</sub>	11.10	10 <sup>3</sup> m <sup>3</sup> /d	0.392 mmscf x 28.3168 10 <sup>3</sup> m <sup>3</sup> /mmscf

If both condensate and water are present, use the Turner Correlation for water to evaluate system behavior. The Turner Correlation uses the cross-sectional area of the flow path when calculating liquid lift rates. For example, if the flow path is through the tubing, the minimum gas rate to lift water and condensate is calculated using the inside diameter (ID) of the tubing. When the tubing depth is higher in the wellbore than the midpoint of perforations, the midpoint elevation between the highest and lowest perforations in the casing in a vertical well, the Turner Correlation does not consider the rate required to lift liquids between the midpoint of perforations and the end of the tubing. Ultimately, the liquid lift rate calculations are based on the tubing’s ID or the area of the annulus and not on the casing’s ID unless flow is up the casing only. Midpoint of perforations is the midpoint elevation between the highest and lowest

<sup>2</sup> Turner, R. G., Hubbard, M. G., and Dukler, A. E., 1969, *Analysis and Prediction of Minimum Flow Rate for the Continuous Removal of Liquids from Gas Wells*, JPT 21(11): 1475–1482.

perforations in the casing. The Midpoint of Perforations elevation is used in the Turner Correlation that plays a role in the well testing decision tree found in Section 7.4.1.1.

**Note 4:** Average Monthly LGR/CGR Calculation

Follow [Figure 7.10](#) to determine if a facility exception is appropriate for specific wells that flow to the reporting facility based on the total liquid/condensate volumes versus the total gas volume measured at the group measurement point for the reporting month. Production volumes include not only volumes measured at a group measurement point, but all fluid production volumes used for reporting purposes. This requires accounting for all fluid volumes that are received into or delivered out of the reporting facility for that reporting month.

$$\text{LGR} = \frac{[\text{Total group measured liquids (condensate + water)} + (\text{Disposition} + \text{Inventory change before group measurement}) - \text{Liquid received}]}{[\text{Total group measured gas} + (\text{fuel} + \text{flare} + \text{vent before group measurement}) + \text{Disposition before group gas measurement} - \text{Gas received}]}$$

$$\text{CGR} = \frac{[\text{Total group measured condensate} + (\text{Disposition} + \text{Inventory change before group measurement}) - \text{Condensate received}]}{[\text{Total group measured gas} + (\text{fuel} + \text{flare} + \text{vent before group measurement}) + \text{Disposition before group gas measurement} - \text{Gas received}]}$$

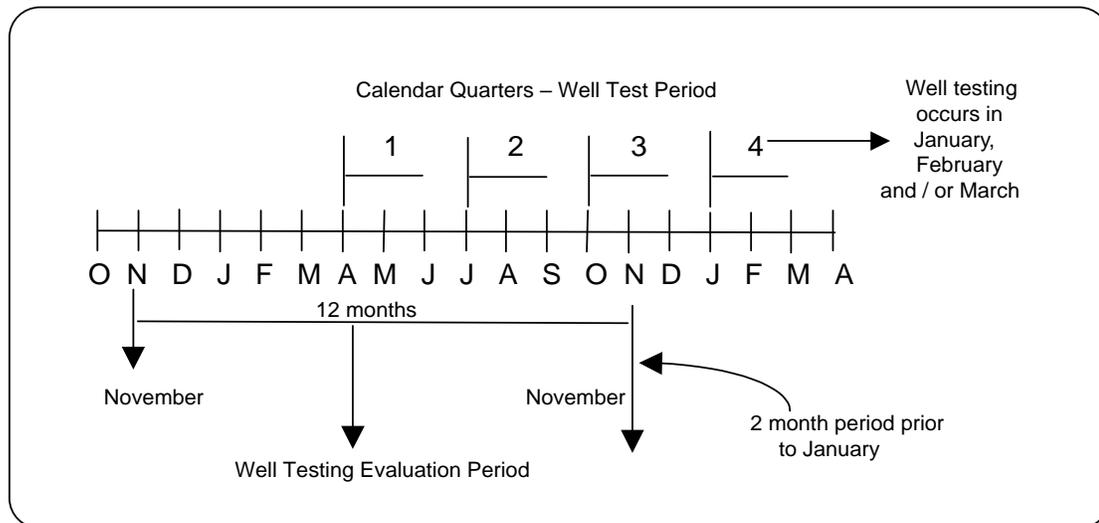
**Note 5:** An initial well test must be conducted within 30 days of the well being put on production and monthly thereafter. The WGR, CGR, and ECF factors from the last test must be used to calculate estimated production until the next test is conducted. Once full fluid recovery is achieved or the 12 month period is passed, whichever comes first, the well must be evaluated according to the decision tree process based on the last well test.

**Note 6:** Wells that require biennial testing must use the ECF, CGR, WGR, and sample analysis from the most current ECF test until the next ECF test results and sample analysis are available.

#### 7.4.1.2 Well Test Evaluation

The well testing evaluation period is based on a cycle of 12 consecutive months that all of the wells in a reporting facility will identically follow. The well test evaluation period must end two months before the planned calendar quarter in which the required well testing will be conducted for a reporting facility. Once the evaluation period is chosen, it remains fixed for a reporting facility. Well testing, when required according to [Figure 7.10](#), must occur once in the fixed calendar quarter. [Figure 7.11](#) provides an illustrated example.

Well and reporting facility data are gathered for the 12 month period identified. The wells and/or the reporting facility are analyzed within the context of the well measurement and testing decision tree. Initializing the design will establish the cycle that is repeated year over year. The licensee is free to choose the well testing calendar quarter based on operations.

**Figure 7.11 Well test evaluation example**

For the purposes of evaluating text box 10 of the well measurement and testing decision tree in [Figure 7.10](#), the reporting facility and the affected wells (i.e. wells without well separation) will be on the same well testing evaluation period. If, however, a reporting facility is not eligible for a reporting facility well testing exception as described in text box 10, the well testing evaluation period can become unique to a well. This means that, for a well that requires testing according to the well measurement and testing decision tree, the well maintains a fixed well testing evaluation period, but the well testing evaluation period may be different for the wells in a reporting facility. Once the well testing period of a calendar quarter is chosen, the licensee must test the well once in the fixed calendar quarter period.

The pressure data, as recorded by the well measurement equipment, will be the monthly average for the last month of the well test evaluation period. If no tubing or casing pressure records are continuously recorded, then the upstream static pressure data from the well's flow meter may be used to approximate the tubing or casing pressure provided that the well's flow meter is located on the same lease site as the wellhead.

#### 7.4.1.3 Record Keeping

The following lists the minimum records required related to well testing and/or the well measurement and testing decision tree where it is applicable:

*General Information:*

1. Licensee
2. Reporting facility – name and surface location
3. Petrinex reporting facility ID
4. Well – unique well identifier (UWI)
5. Production formation – name and/or stratigraphic unit ID

*Well Test Information:*

6. Current well testing date
7. Last well test date
8. Effluent well meter run – internal diameter (mm)

9. Meter run orifice size (mm) (if applicable)
10. Test tap location (relative to effluent meter)
11. Test tap connection – diameter (mm)
12. Last gas sample date
13. Last condensate sample date
14. Test gas average rate ( $10^3 \text{ m}^3/\text{day}$ )
15. Test condensate average rate ( $\text{m}^3/\text{day}$ )
16. Test water average rate ( $\text{m}^3/\text{day}$ )
17. Current WGR ( $\text{m}^3/10^3\text{m}^3$ )
18. Current CGR ( $\text{m}^3/10^3\text{m}^3$ )
19. Current LGR ( $\text{m}^3/10^3\text{m}^3$ )
20. Last WGR ( $\text{m}^3/10^3\text{m}^3$ )
21. Last CGR ( $\text{m}^3/10^3\text{m}^3$ )
22. Last LGR ( $\text{m}^3/10^3\text{m}^3$ )
23. ECF – last value calculated
24. ECF – current value calculated

*Decision Tree Information*

25. Wellhead tubing internal diameter (mm)
26. Wellhead casing internal diameter (mm)
27. Wellhead tubing pressure (KPa)
28. Wellhead casing pressure (KPa)
29. Effluent meter monthly average differential pressure for evaluation period (kPa) – listed by month (optional)
30. Effluent meter monthly average static pressure for evaluation period (KPa) – listed by month
31. Effluent meter monthly average temperature for evaluation period ( $^{\circ}\text{C}$ ) – listed by month (optional)
32. Evaluation period average reporting facility LGR
33. Evaluation period average reporting facility CGR
34. Artificial lift method, i.e., cycling, plunger control
35. Well chart or EFM – model and make
36. Well test evaluation period starting month
37. Well test evaluation period ending month
38. Date well dropped below critical velocity
39. Critical lift calculation for evaluation period

40. Well load fluid volumes for evaluation period
41. Meters used in facility LGR calculations
  - a. Meter tag
  - b. Meter location
  - c. Meter volume
  - d. Meter units ( $10^3\text{m}^3$ , etc.)
42. Well flow volume prior to recompletion (optional)
43. Well recompletion flow volume (optional)

#### 7.4.1.4 Revocation of Exception Allowed by Decision Tree

A testing exception for an effluent gas well may be revoked if certain criteria are not met. Baseline well testing must be conducted if a testing exception is revoked for any of the following reasons:

1. Noncompliance

Potential areas of noncompliance include:

- a. Incorrect exception calculations,
- b. Inadequate record keeping,
- c. Source data for exception calculations cannot be validated, and
- d. Incorrect application/implementation of the well measurement and testing decision tree.

2. A working interest participant or Freehold royalty owner for any flowing well to the reporting facility objects to the exception.

Additionally, if ER has a concern with the activities, operations, production data, or reporting associated with well testing, on notice in writing, ER can partially or fully revoke well testing exceptions and impose, modify, or substitute well testing conditions for any period of time. ER will advise the licensee in writing as to the reason for the revocation, provide a reasonable time period for the licensee to meet the conditions set by ER, and provide an opportunity for the licensee to comment.

#### 7.4.2 Production Volume Calculations

Monthly production volumes are to be calculated as follows (see [Figure 7.12](#)).

Units: All gas volumes and GEV are to be in  $10^3\text{m}^3$  and liquid volumes in  $\text{m}^3$ .

#### 7.4.2.1 Testing-Exempt Battery

For a battery that is exempt from testing, the volumetric calculation is to be based on the following:

$$ECF = 1.00000$$

WGR = Battery-based water-gas ratio

LGR = Battery-based liquid-gas ratio

CGR = Battery-based condensate-gas ratio

If battery condensate volumes are recombined back into the gas stream, the gas equivalent of the recombined liquids will be calculated and added to the measured group gas volume to obtain the total battery gas volume. Condensate liquid volumes will not be prorated to the wells. If battery condensate volumes are tanked and trucked out for sale or to a gas plant for further processing, the condensate liquid volumes will be prorated back to the wells in the battery based on the calculated battery CGR.

Well water production may be determined by either:

1. Calculating the battery water proration factor and then multiplying the well's estimated water production by the battery's water proration factor or
2. Multiplying the wells' percentage of the total estimated gas production by the monthly measured battery water volume. In this case report a battery water proration factor of 1.00000
3. Exception: The licensee may, providing there is no objection from the working interest owners of any well producing into the battery, use the WGR, CGR and ECF from each well's most recent ECF test instead of using the battery-calculated WGR, CGR and ECF of 1.00000. This option may be used as long as the battery qualifies as a test-exempt battery.

#### 7.4.2.2 Testing - Exempt Wells

For batteries with both exempt and nonexempt wells, the volumetric calculation must be based on the following:

$$ECF = 1.00000 \text{ for exempt wells}$$

For the wells that require testing, water production will be prorated to each well based on the well's individual WGR derived from the well tests multiplied by its estimated gas production. For those wells that are test exempt, a battery WGR will be established and applied to all the test-exempt wells after netting off the estimated water production of the tested wells.

If battery condensate volumes are recombined back into the gas stream, the gas equivalent volume of the recombined condensate will be calculated and added to the measured group gas volume (recombined volume and recombined analysis). Condensate liquid volumes will not be prorated to the wells.

If battery condensate volumes are separated and delivered from the battery, the tested wells' estimated condensate production will be calculated based on each well's CGR, derived from the well tests, multiplied by the well's estimated gas production. For those wells that are test exempt, a battery CGR will be established and applied to all the test-exempt wells after netting off the estimated condensate production of the tested wells.

Exception: For test-exempt wells, the licensee may, at its discretion, use the WGR, CGR, and ECF from each well's most recent ECF test instead of using the battery-calculated WGR, CGR, and ECF of 1.00000.

### 7.4.2.3 Testing - Nonexempt Wells

The following example assumes there are no measured receipts at the battery upstream of the group measurement point. If there are no exempt wells in the battery and condensate is delivered for sale at the group measurement point, go directly to item 8. Otherwise, follow items 1 to 7.

1. Calculate the ECF:

$$\text{ECF} = (\text{Well test gas volume [B]} + \text{GEV of well test condensate [C]}) \div \text{Effluent gas volume measured during test (A)}$$

2. Calculate estimated gas production for each well:

$$\text{Estimated monthly well gas volume} = \text{Monthly well effluent volume} \times \text{ECF}$$

3. Calculate the WGR:

$$\text{WGR} = \text{Well test water volume (D)} \div (\text{Well test gas volume [B]} + \text{GEV of well test condensate [C]})$$

4. Calculate estimated water production for each well:

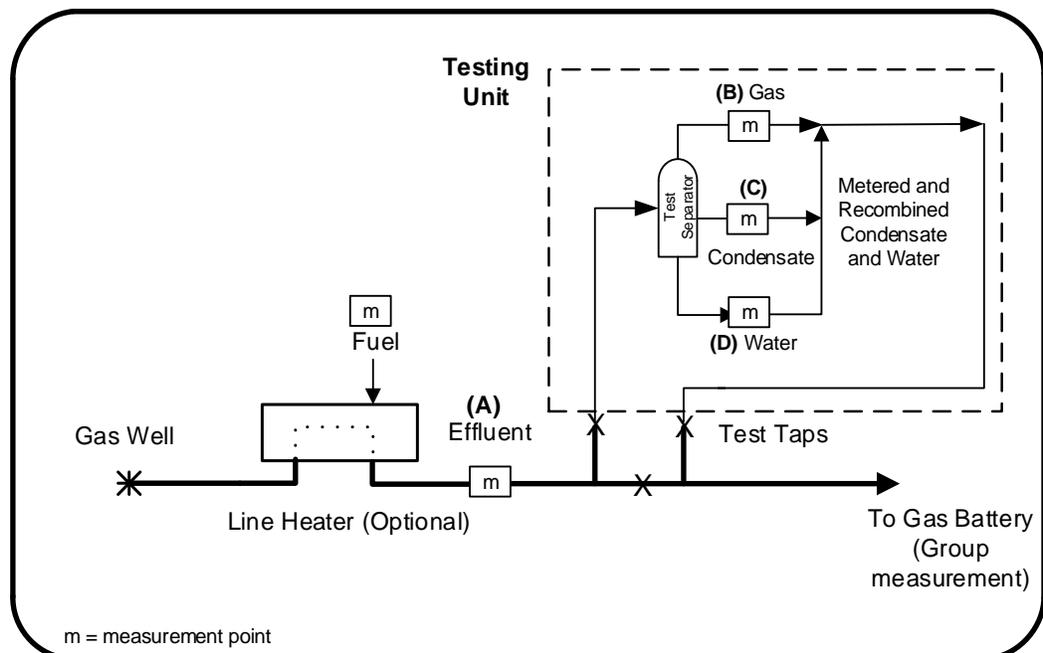
$$\text{Estimated monthly well water volume} = \text{Estimated monthly well gas volume} \times \text{WGR}$$

5. Calculate total battery estimated volumes (gas and water):

$$\text{Total battery estimated monthly gas volume} = \text{Sum of all estimated monthly well gas volumes}$$

$$\text{Total battery estimated monthly water volume} = \text{Sum of all estimated monthly well water volumes}$$

Figure 7.12 Effluent well meter testing configuration with condensate production



6. Calculate proration factors for gas and water:

$$\text{Gas Proration Factor} = (\text{Total battery measured monthly gas volume} + \text{GEV of total battery condensate}) \div \text{Total battery estimated monthly gas volume}$$

$$\text{Water Proration Factor} = \text{Total battery actual monthly water volume} \div \text{Total battery estimated monthly water volume}$$

7. Calculate reported monthly (prorated) well production:

$$\text{Reported monthly well gas production} = \text{Estimated monthly well gas volume} \times \text{Gas Proration Factor}$$

$$\text{Reported monthly well water production} = \text{Estimated monthly well water volume} \times \text{Water Proration Factor}$$

For the battery with condensate delivered for sale at the group measurement point:

8. Calculate the ECF:

$$\text{ECF} = \text{Well test gas volume (B)} \div \text{Effluent gas volume measured during test (A)}$$

9. Calculate the well CGR:

$$\text{CGR} = \text{Well test condensate volume (C)} \div \text{Well test gas volume (B)}$$

10. Calculate the WGR:

$$\text{WGR} = \text{Well test water volume (D)} \div \text{Well test gas volume (B)}$$

11. Calculate estimated gas, condensate, and water production for each well:

$$\text{Estimated monthly well gas volume} = \text{Monthly well effluent volume} \times \text{ECF}$$

$$\text{Estimated monthly well condensate volume} = \text{Estimated monthly well gas volume} \times \text{Condensate Gas Ratio}$$

$$\text{Estimated monthly well water volume} = \text{Estimated monthly well gas volume} \times \text{Water Gas Ratio}$$

12. Calculate total estimated gas, condensate, and water production for the battery:

$$\text{Total battery estimated monthly gas volume} = \text{Sum of all estimated monthly well gas volumes}$$

$$\text{Total battery estimated monthly condensate volume} = \text{Sum of all estimated monthly well condensate volumes}$$

$$\text{Total battery estimated monthly water volume} = \text{Sum of all estimated monthly well water volumes}$$

13. Calculate total battery monthly gas, condensate, and water production:

$$\text{Total battery monthly gas volume} = \text{Total gas disposition} + \text{Flare} + \text{Vent} + \text{Fuel (take off before sales meter)}$$

$$\text{Total battery monthly condensate volume} = \text{Total condensate disposition} + \text{inventory change}$$

$$\text{Total battery monthly water volume} = \text{Total water disposition} + \text{inventory change}$$

14. Calculate proration factors for gas, condensate, and water:

$$\text{Gas Proration Factor (GPF)} = \text{Total battery monthly gas volume} \div \text{Total battery estimated monthly gas volume}$$

$$\text{Condensate Proration Factor (CPF)} = \text{Total battery monthly condensate volume} \div \text{Total battery estimated monthly condensate volume}$$

$$\text{Water Proration Factor (WPF)} = \text{Total battery monthly water volume} \div \text{Total battery estimated monthly water volume}$$

15. Calculate reported monthly (prorated) well production:

$$\text{Reported monthly well gas production} = \text{Estimated monthly well gas volume} \times \text{Gas Proration Factor}$$

Reported monthly well condensate production = Estimated monthly well condensate volume x  
Condensate Proration Factor

Reported monthly well water production = Estimated monthly well water volume x Water  
Proration Factor

### 7.4.3 Sampling and Analysis Requirements

#### 7.4.3.1 Testing Exempted Batteries

For testing exempted batteries, the well sample and analysis used may be either:

1. The sample and analysis obtained from the most recent ECF test or
2. The annual sample and analysis obtained from the group separator, provided that
  - a. There is common ownership in all wells in the battery;
  - b. If there is no common ownership, written notification has been given to all working interest participants, with no resulting objection received; and
  - c. If there is no common Crown or common Freehold royalty and only Freehold royalties are involved, written notification has been given to all Freehold royalty owners, with no resulting objection received. If there is a mix of Freehold and Crown royalty involved, the licensee must apply to ER for approval.
  - d. There are no other receipts into the battery upstream of the group separator.

Regardless of which of the above approaches is used, the licensee may, at its discretion, test and sample any well and use the well sample and analysis to calculate well volume.

#### 7.4.3.2 Testing Exempted Wells

For test-exempt wells in batteries that have tested and test-exempt wells, the well sample and analysis used may be either:

1. The sample and analysis obtained from the most recent ECF test or
2. The annual sample and analysis obtained from the group separator, provided that
  - a. There is common ownership in all wells in the battery;
  - b. If there is no common ownership, written notification has been given to all working interest participants, with no resulting objection received; and
  - c. If there is no common Crown or common Freehold royalty and only Freehold royalties are involved, written notification has been given to all Freehold royalty owners, with no resulting objection received.
  - d. There are no other receipts into the battery upstream of the group separator.

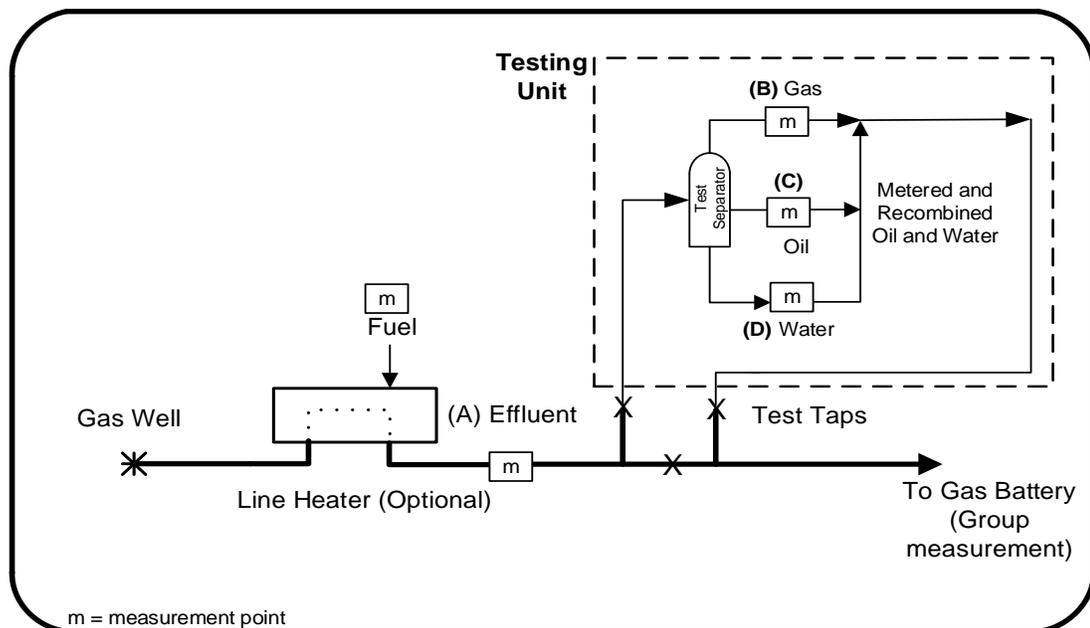
Regardless of which of the above approaches is used, the licensee may, at its discretion, test and sample any test-exempt well and use the well sample and analysis to calculate well volume.

#### 7.4.4 Exception for Gas Wells Producing Oil in Effluent Measurement Battery

If the hydrocarbon liquid that a gas well produces is reclassified by ER from condensate to oil, the well may remain in a Gas Multi-well Effluent Measurement Battery, provided that the well is equipped with a separator and there is continuous measurement of the gas and liquid components or, alternatively, the effluent meter is left in place and the measurement, accounting, and reporting procedures specified below are followed, (see [Figure 7.13](#)).

If the effluent meter is left in place, annual ECF-WGR tests must be conducted on the well. These types of wells do not qualify for the test frequency exceptions or reductions described in Section 7.4. An OGR, to be used for the well oil production calculation, must also be determined during this test. The WGR, estimated water production, water proration factor, and actual water production are determined in the same manner as indicated in Section 7.4.2.

**Figure 7.13 Effluent well configuration for a gas well producing condensate reclassified to a gas well producing oil**



The following example assumes there are no measured receipts at the battery upstream of the group measurement point.

1. Calculate the ECF:

$$\text{ECF} = \text{Well test gas volume (B)} \div \text{Effluent gas volume measured during test (A)}$$

2. Calculate estimated gas production for the well:

$$\text{Estimated monthly well gas volume} = \text{Monthly well effluent volume} \times \text{ECF}$$

3. Calculate the OGR:

$$\text{OGR} = \text{Well test oil volume (C)} \div \text{Well test gas volume (B)}$$

4. Calculate reported well oil production:

$$\text{Reported monthly well oil volume} = \text{Estimated monthly well gas production} \times \text{OGR}$$

5. Calculate the total monthly battery condensate volume:

Total battery condensate volume = Total battery liquid hydrocarbon volume – Total monthly oil volume

6. Report the calculated monthly oil production volume as oil produced from the well. Prorate monthly gas and water production as in Section 7.4.2.

## 8 Gas and Liquid Sampling and Analysis

This section outlines the gas and related liquid sampling and analysis requirements for wells, group measurement points, and other streams associated with the various Petrinex reporting facility subtypes. These requirements add to the requirements in Section 93.1 of *The Oil and Conservation Regulations, 2012*, which continues to apply.

The requirements vary, depending on a number of factors, such as production rate, potential for the composition to change over time, and the end use of the fluid. Where applicable, conditions have been identified under which the sampling and analysis requirements may be altered or eliminated altogether. ER may also consider applications for further requirement alterations or eliminations if the licensee can demonstrate that measurement accuracy would either not be reduced or there would be not negative impacts on royalties, working interest owner equity, environmental protection, public safety or reservoir engineering concerns. In Saskatchewan, all sample analysis reports for both facilities and wells must be submitted to ER in accordance with the specifications listed in *Directive PNG013: Well Data Submission Requirements*.

### 8.1 General

Gas and liquid analyses are required for the determination of gas volumes, conversion of liquid volumes to gas equivalent, and product allocation. The sampling and analysis requirements described in Section 8 pertain only to those areas that affect the calculations and reporting required by ER.

These requirements apply solely to the measurement of hydrocarbon fluids and are not intended to supersede the business requirements that licensees are required to meet regarding product allocations.

If oil is produced from gas wells, as defined by ER, the oil must be reported as liquid oil production and not as a gas equivalent volume (GEV). Therefore, compositional analysis of the oil is not required for that purpose. The oil produced may be combined with the gas and delivered to a gas plant or other facilities for further processing, or the oil could be separated from the gas at the well equipment and directed to tankage, and then for further treatment or sale.

Gas density and composition are integral components of gas volume calculations and plant product allocation calculations. For differential producing meters, such as orifice meters, venturi meters, and flow nozzles, the accuracy of a computed volume and component allocations are very sensitive to the accuracy of the compositional analysis, which is the basis for compressibility factors and density determination. For linear meters, such as ultrasonic and vortex, the compositional analysis is primarily used to determine the compressibility factors.

If liquid condensate produced from a gas well is recombined with the well gas production, the compositional analysis from a condensate sample must be used to determine the GEV of the condensate, which must be added to the well gas volume for reporting purposes. A similar procedure applies to gas gathering systems where liquid condensate is delivered to other facilities. For this reason, the condensate sampling requirements must be at the same frequency as the gas sampling requirements.

If liquid condensate is separated at a well, battery, or gas gathering system and delivered from that point for sale or other disposition, or trucked to a gas plant or other facility for further processing, the condensate must be reported as a liquid volume. Therefore, a compositional

analysis of the condensate is not required for gas equivalent volume determination purposes but may be required for the purposes of the sale.

Sampling and analysis frequencies and updating requirements for the various well and facility subtypes are summarized in Section 8.4. Further details are provided in the sections that follow. These sampling frequencies are the base requirements for gas and related liquid measurement.

## 8.2 Sampling and Analysis Requirements

Except where noted in this Directive, the gas sampling equipment and methodology must follow the requirements set out in API MPMS 14.1 of June 2001, Gas Processors Association (GPA) 2166-05, or other equivalent industry standards.

Except where noted in this Directive, the condensate sampling equipment and methodology must follow the requirements set out in GPA 2174-93, the evacuated cylinder method cited in GPA 2166-05, or in other equivalent industry standards.

Samples and analysis may be obtained by any of the following methods:

1. On-site gas chromatograph (GC)
2. Proportional sampling
3. Spot or grab sampling

Spot or grab samples are acceptable for obtaining gas and liquid analyses once per test or per determination, provided that uncertainty requirements in Section 1, Standards of Accuracy are fulfilled. When the uncertainty requirements cannot be met, licensees must consider more frequent sampling, calculated analyses in Section 8.3.2, proportional samplers, or gas chromatographs.

For example, if the analysis from one-time period to the next is such that the density and/or compressibility changes cause the volume to change by more than the allowable uncertainty, a more frequent analysis is required or an alternative method of obtaining the sample must be used.

The gas and liquid analyses must be updated when operating conditions are significantly altered such as through the addition/removal of compression or line-heating, addition/removal of production sources in a common stream, or wellbore recompletion.

If the gas volumes for all meters in the common stream such as sales, fuel, flare, and injection gases meet the uncertainty guidelines in Section 1, Standards of Accuracy, the licensee may use a single gas analysis for all meters on the common stream.

### 8.2.1 Sampling Procedures

1. Sample points must be located to provide representative samples.
2. Sample probes must not be located within the minimum upstream straight lengths of the meter.
3. Access from grade or platform must be provided for the sample point.
4. If sample transfer tubing is to be used, its length must be minimized.
5. The sample transfer tubing must be oriented to minimize the potential to trap liquids in gas samples and water in condensate samples.

6. A means must be provided to safely purge sample transfer tubing between the sample point and the connection point of the sample cylinder.
7. Sample containers must be clean and meet the pressure, temperature, and materials requirements of the intended service and have the required Regulatory approvals as necessary.
8. The procedures used for sampling, transportation, handling, storage, and analysis must ensure that atmospheric contamination does not occur.

All samples must be analyzed using a gas chromatograph or equivalent to determine the components to a minimum of C<sub>7+</sub> composition except for sales or delivery points where C<sub>6+</sub> composition is acceptable if agreed upon by affected parties. The gas composition analysis must be determined to a minimum of four decimal points as a fraction of 1.0000 or two decimal points as a percentage of 100, and the relative density must be determined to a minimum of three decimal points.

## 8.2.2 Sample Point and Probes

The sample point location and probe installation requirements that follow apply to all measurement points described in this Directive. With the exception of delivery point measurement, current sample point locations and installations that were constructed prior to April 1, 2016 do not have to be upgraded to meet the sample probe requirements but must meet the sample point requirements.

For grandfathered sample point locations as describe above, a sample probe must be installed according to the requirements in this section when an installation is relocated or reused for another well or facility.

### 8.2.2.1 Requirements for Gas Sampling

1. For sampling applications where the gas is at or near its hydrocarbon dew point, a sample probe must be used. This requirement applies to any separator application where hydrocarbon liquids are present.
2. For gas applications where the gas is not near its hydrocarbon dew point a sample probe is recommended.
3. The preferred location for gas sample points is the top of horizontal lines.
4. An optional location for gas sample probes is the side of vertical lines with the probe tip sloping 45.0° downward.
5. Sample probes must be located at least five pipe diameters downstream of any piping disturbances, such as bends, elbows, headers, and tees.
6. The location of the sample point must be such that phase changes due to changes in pressure and/or temperature are minimized. Specifically, for gases at or near their hydrocarbon dew point, sample points must not be located downstream of pressure-reducing components, such as control valves, flow conditioners, and pressure regulators , or long lengths of uninsulated piping or within five pipe diameters downstream of an orifice plate.
7. Sample points may be located downstream of ultrasonic meters that experience minimum pressure drop through the meter unless a flow conditioner is used and the

gas is at or near its hydrocarbon dew point, in which case the sample point must be upstream of the flow conditioner.

8. Insulation and heat tracing must be used to eliminate any cold spots between the sample point and the entry point into the sample container or gas chromatograph where the sample transfer tubing temperature falls below the hydrocarbon dew point, such as at all separator applications.
9. Sample points used to sample blends of two gas streams must have provision for mixing, such as an upstream static mixer, with due consideration to potential phase changes brought about by a pressure drop associated with the mixing device.
10. Orifice meter impulse lines or transmitter manifolds lines must not be used for taking samples.
11. Level gauge connections must not be used for taking samples.

### 8.2.2.2 Requirements for Condensate Sampling

1. With the exception of two-phase separators, a sample probe is recommended.
2. A sample probe must be installed for samples to be used to determine water cut when there is emulsion or a mix of water and hydrocarbon, such as two-phase separators. For such applications, the sampling system design must meet the requirements of API MPMS 8.2 with respect to the use of mixers, sample probe location, and design.
3. The preferred location for condensate sample points is the side of horizontal lines.
4. An optional location for liquid sample points is the side of vertical lines with the probe tip sloping 45.0° downward.
5. The location of the sample point must be such that phase changes due to changes in pressure and/or temperature are minimized. Specifically, sample points must not be located where vapour breakout is likely such as downstream of pressure-reducing components, orifice plates, flow conditioners, turbine, PD or Coriolis mass meters, control valves, and pressure regulators or where the stream temperature has increased.
6. For separator applications, the sample point must be between the separator outlet and the flow/level control valve upstream of the meter, unless a pressure booster pump is used, in which case the sample point must be located between the pump discharge and the meter.
7. Level gauge connections must not be used for taking samples.

### 8.2.3 H<sub>2</sub>S Sampling and Analysis

This section relates to obtaining high pressure samples. Special considerations, such as extra sample(s) or purging, should be taken when obtaining low pressure samples at a boot separator configuration, treater, stabilizer, or at an acid gas facility.

Hydrogen sulphide (H<sub>2</sub>S) is a reactive molecule that presents challenges for sampling and analysis of gas mixtures containing it. H<sub>2</sub>S is easily lost during sampling and analysis, resulting in underreporting of H<sub>2</sub>S concentrations. Factors that affect representative sampling and analysis accuracy through H<sub>2</sub>S loss are:

1. Presence of air, water, or other sulphur-containing molecules
2. Presence of reactive or absorptive sampling container surfaces

3. Presence of a liquid phase, which can absorb H<sub>2</sub>S
4. H<sub>2</sub>S concentration
5. Sample pressure and temperature
6. Analysis method
7. Time lapse between sampling and analysis

The amount of H<sub>2</sub>S lost can be reduced by:

1. Proper sample point selection, which minimizes the presence of contaminants such as air, water, and amines
2. Using clean containers made of materials that minimize H<sub>2</sub>S reactions or absorption
3. Minimizing the time between sampling and analysis

Typical construction materials for cylinders are stainless steel and aluminum. Inert coated cylinders, glass containers, and non-absorptive elastomer bags can be considered to further minimize H<sub>2</sub>S degradation, especially for concentrations of H<sub>2</sub>S less than 5000 ppm when moisture is present.

The choice of analytical technique also affects the amount of H<sub>2</sub>S reported. Instrument-oriented techniques, such as gas chromatography, are typically more precise than chemistry-oriented techniques, such as Tutweiler titrations or stain tubes. However, such instrument-oriented techniques are often impractical for individual well site applications.

Therefore, consideration should be given to analysis technique limitations and sample degradation as they relate to the specific reporting requirements in determining the best approach.

See [Table 8.1](#) for analysis technique comparison.

With the exception of ppm level concentrations of H<sub>2</sub>S in the presence of moisture, a field H<sub>2</sub>S determination along with a laboratory GC analysis are recommended. These analysis techniques provide a degree of redundancy and a check of the field analysis. Above 5.0 per cent H<sub>2</sub>S, the GC value is typically more reliable. Below 5.0 per cent H<sub>2</sub>S, the higher of the two values must be used. Unexpectedly large variances between lab and field H<sub>2</sub>S values must be investigated.

**Table 8.1 H<sub>2</sub>S analysis technique comparison**

Analysis technique	Lower detection limit	Advantages	Limitations
Online GC	500 ppm	Real-time, accuracy Minimal elapsed time	Capital cost, ongoing maintenance
Laboratory GC	500 ppm	Precision, accuracy	Potential degradation during transport that varies with H <sub>2</sub> S concentration
Tutweiler GPA C-1	1500 ppm	On site	Titration apparatus, reagent quality, variability in operator technique, including visual endpoint detection, computations, mercaptan interference
Stain Tubes GPA 2377	1 ppm	On site	Poor precision ( $\pm 25.0\%$ ) Matrix effects as described in manufacturer's specifications

Analysis by gas chromatography is the preferred method at higher H<sub>2</sub>S concentrations.

For H<sub>2</sub>S concentrations between 1500 and 5000 ppm, it is recommended that both stain tube and Tutweiler values be obtained if online GC is not used.

If high accuracy of low-level, below 1500 ppm, H<sub>2</sub>S concentration is required, consideration should be given to using a low-level sulphur-specific detector, such as a GC sulphur chemiluminescence detector. The use of containers that minimize degradation and minimizing the time elapsed between sampling and analysis is also recommended in these situations.

Refer to Appendix 4 for more detail on the analytical methods used in the industry for determining H<sub>2</sub>S concentrations in gas samples.

### 8.2.4 Compositional Analysis of Natural Gas

The two recommended procedures for compositional analysis of natural gas are based on GPA Standard 2286-95: *Tentative Method of Extended Analysis for Natural Gas and Similar Gaseous Mixtures by Temperature Programmed Gas Chromatography* and GPA Standard 2261-00: *Analysis for Natural Gas and Similar Gaseous Mixtures by Gas Chromatography*.

If a thorough molecular weight and density description of the C<sub>7+</sub> fraction is required, analytical methods based on GPA Standard 2286 are more accurate and preferred. Specifically, GPA Standard 2286 makes use of a high-resolution column and flame ionization detector to separate and quantify the heavier C<sub>7+</sub> components, which is then used for calculation purposes. Extended analysis of natural gases is common but has not been fully standardized; therefore, some bias may occur from one laboratory to another.

If the C<sub>7+</sub> properties are well defined or have been agreed upon by all affected parties, analytical methods based on GPA Standard 2261 are suitable. The principal advantage of the precut method specified in GPA Standard 2261 is that all of the C<sub>7+</sub> components can be grouped together into a single sharp chromatograph peak. Grouping of the numerous heavy compounds results in more precise measurement of the combined peak area. The primary disadvantage of GPA Standard 2261 is the lack of information gained with respect to the composition of the C<sub>7+</sub> fraction. Inherently, if the composition of the C<sub>7+</sub> fraction is unknown, some agreed-upon physical properties must be applied for calculation purposes. The GC C<sub>7+</sub> calibration is also affected, which increases the uncertainty of the C<sub>7+</sub> measurement and heating value computation.

If detailed information on C<sub>7+</sub> physical properties is not available, default values can be applied, as in [Table 8.2](#).

**Table 8.2 Recommended default values for C<sub>7+</sub> properties\***

Component Names	Molecular mass grams per mole	Liquid density kg/m <sup>3</sup> at 15°C	Heating value MJ/m <sup>3</sup>
C <sub>7+</sub> , Heptanes plus	95.00	735.0	195.0

\* C<sub>7+</sub> is a pseudo-compound. The values in most scenarios have been found to adequately represent the heavier fraction of natural gas samples.

### 8.3 Gas Equivalent Factor Determination from Condensate

Gas Equivalent Factor (GEF) is the volume of gas,  $10^3\text{m}^3$  at standard conditions that would result from converting  $1.0\text{ m}^3$  of liquid into a gas.

GEF is used when there is a requirement to report the gas equivalent volume (GEV) of condensate and other hydrocarbon liquids to ER.

The GEF of a liquid may be calculated by any one of three methods described in Appendix 5, depending upon the type of component analysis conducted on the liquid such as volume, mole, or mass fractions and the known properties of the liquid.

#### 8.3.1 Engineering Data

Specific constants are used in calculating the GEF. Absolute density of liquids should be used instead of relative density.

The examples in Appendix 5 present the different methodologies used to calculate the GEF. All physical properties are based on GPA Standard 2145-03 (2003 or later) published data.

$$1\text{ kmol} = 23.645\text{ m}^3 @ 101.325\text{ kPa and }15.0^\circ\text{C}.$$

#### 8.3.2 Calculated Compositional Analyses

In some instances, representative sampling of a hydrocarbon stream is not possible or feasible because of economics, and calculation of a fluid composition is required, as described in this section:

**Calculated Well Stream Analysis:** It is not possible to accurately sample multiphase streams, so the composition of a recombined well stream must be determined by calculation. Such an analysis is typically not used for measurement, as it represents a multiphase fluid stream and most gas is measured as single phase. However, some companies use this analysis for calculation of gas volumes from effluent measured wells. Calculated well stream analyses are most commonly used in product allocation calculations.

**Calculated Group Analysis:** It is often difficult to accurately determine the average composition of fluids at a commingled group measurement point, as wells/sources to the group system flow at different rates and the composition is constantly changing. The recommended options for sampling these streams are on-line gas chromatographs or proportional sampling systems. However, if the recommended options are not practical or economical, a flow-weighted calculated analysis may be a viable option.

**Calculated Single Analysis:** Sometimes a single analysis cannot represent the composition for an entire measurement period. In such scenarios, multiple analyses of a single point must be combined to determine the composition for the period. An example of this is a sales gas stream where a proportional sample is taken weekly but a single composition for the month is required.

The principles to be followed for each of these calculated analyses follow.

##### 8.3.2.1 Calculated Well Stream Compositional Analysis

This type of analysis applies to wells only and is meant to represent the hydrocarbon fluid composition produced from a well and/or delivered to a gathering system. In most scenarios, it represents the composition of hydrocarbons being produced from the reservoir. The calculation

is a flow-weighted recombination of the hydrocarbon gas and liquid streams. The accuracy of the flow rates used in this calculation is as important as the gas and liquid composition.

Gas and liquid flow rates from the day of sampling must be used in determining recombined compositions, with the following exceptions:

1. When the daily liquid-to-gas ratio is constant, volumes from an extended period such as multiday or up to monthly may be used.
2. If some of the liquid stream is not recombined in a month such as scenarios where it is produced to a tank, the composition flow volume of the liquids not recombined must be deducted from the initial recombined composition. This is typically performed by recalculating the recombined composition using only the liquid volume that was recombined with the gas during the month.

See the example in Appendix 6.

### **8.3.2.2 Calculated Group Compositional Analysis**

This type of analysis is a flow-weighted representation of the hydrocarbon fluid composition produced from a group of wells or meter points. It is often used at commingled group points such as inlets, compressors and certain process points where it is difficult to obtain representative samples using spot sampling techniques. Ideally, proportional samplers should be employed in such situations. However, when proportional sampling is not practical or possible, a calculated group analysis can be determined based on the volume and composition of the wells/meters that flow to the commingled point. The accuracy of the flow rates used in this calculation is as important as the gas and liquid composition. The flow volumes used for each well/meter should be actual measured volumes for the period that the analysis is being calculated for, typically monthly.

For example, five gas wells producing from different pools with different composition deliver gas to a compressor station where the gas is measured. Accurate spot sampling at the compressor station is difficult due to changing flow rates at the wells. Using spot samples taken at the wells and monthly flow rates, the producer calculates a group analysis for the compressor station meter. Care must be taken when separator liquids are produced that all hydrocarbons are correctly accounted for, regardless of the phase.

See the example in Appendix 6.

### **8.3.2.3 Calculated Single Compositional Analysis**

This type of analysis is a flow-weighted representation of the hydrocarbon fluid composition determined at a single sample point. It is typically used at sample points that have variable compositions and are sampled frequently, such as weekly, using spot or proportional sampling. Ideally, proportional samplers or gas chromatographs should be employed in such situations. When proportional sampling is not practical or possible a calculated single analysis can be determined based on the volume and composition of a group of analyses at the sample point. The accuracy of the flow rates used in this calculation is as important as the gas and liquid composition. The flow volumes used for each sample must be actual measured volumes for the period that the analysis is representative of. For example, a producer takes spot samples of an inlet stream weekly because proportional sampling or on-line sampling is not practical. Using weekly flow rates, the producer calculates a monthly flow-weighted composition of the inlet stream.

See the example in Appendix 6.

## 8.4 Gas and Condensate Sampling and Analysis Requirements

For facilities and wells, Table 8.3 outlines the update frequency of sampling and analysis required for gas and condensate streams. The sampling and analysis of condensate, if applicable, must be performed at the same time as the gas sampling. Configuration examples are shown in Figures 8.1 through 8.17. In each scenario, other similar configurations may also apply.

Gas and condensate sampling and analysis must be conducted on all new wells and new measurement points by the end of the month following the first month of production unless otherwise stated in this Directive. For the time period prior to obtaining an analysis from a new well or measurement point, a representative analysis of gas and condensate may be used for gas volume determination and gas equivalent volume of liquid calculations.

For new wells, where possible, representative analyses must be from a well producing from the same pool with similar separator operating conditions or from sample analyse obtained during well testing while the well is undergoing clean up operations. Analyses obtained during well clean up operations should be carefully reviewed prior to use, as samples are typically taken at different conditions from those of the producing well, and there are often contaminants such as nitrogen or frac fluid in test samples. For other measurement points the representative analysis should be reasonably similar to the expected composition.

All sample and analysis reports must be submitted to ER in accordance with the requirements in *Directive PNG013: Well Data Submission Requirements*.

**Table 8.3 Sampling and analysis frequencies for various types of facilities**

	Type of facility	Gas rate 10 <sup>3</sup> m <sup>3</sup> /d	Sample and analysis type	Sampling point	Frequency
<b>Gas wells/ batteries/ facilities</b>	SW Saskatchewan shallow gas stratigraphic units or zones or areas with minimal water (facility subtype 363) See Section 8.4.1	N/A	Gas only	Group meter	Biennially
	Gas proration outside SW Saskatchewan shallow gas stratigraphic units or zones or areas or effluent measurement battery (facility subtypes 362 and 364) See Sections 8.4.1.1 & 8.4.2	N/A	Gas/condensate	Test meters	At time of testing
		>16.9		Group meter	Annually
		≤16.9			Biennially
	Multi-well group battery or single-well battery with no condensate or oil (facility subtypes 361 and 351) See Section 8.4.3	N/A	Gas only	All meters	First month, then Biennially
	Multi-well group battery or single-well battery with condensate or oil (facility subtypes 361 and 351) See Section 8.4.4	> 16.9	Gas/condensate	Per meter	Annually
≤ 16.9		Gas/condensate	Per meter	Biennially	

	Type of facility		Gas rate 10 <sup>3</sup> m <sup>3</sup> /d	Sample and analysis type	Sampling point	Frequency
	Gas storage schemes, injection (facility subtypes 504, 505, 517, and 519) and withdrawal phase (facility subtypes 351, 361, 362, 363, 364) See Section 8.4.5			Gas	Per injection/production meter	First month, then semi-annually
	Gas cycling schemes See Section 8.4.6	Injection		Gas/condensate	Per injection meter	Per approval or first month, then semiannually (if not in approval)
		Production			Per production meter	Per approval or first month, then semi-annually (if not in approval)
	Gas sales/delivery (all facility subtypes apply) See Section 8.4.7			Gas only	Per meter	Annually
	Gas plants (facility subtypes: 401-407 and 411) See Section 8.4.8.1			Gas/condensate	Per accounting meter	Semiannually
	Gas gathering systems (facility subtypes 611 and 612) See Section 8.4.8.2		> 16.9	Gas/condensate	Per inlet meter	Annually
			≤ 16.9	Gas/condensate	Per inlet meter	Biennially
<b>Non-heavy oil wells/batteries</b>	Single-well/ multi-well group battery (facility subtypes: 311 and 321) See Section 8.4.9	Flared		Gas only	Per meter	Initial <sup>2</sup>
		Conserved	> 16.9	Gas only	Per meter	Annually
			≤ 16.9	Gas only	Per meter	Biennially
	Multi-well proration Battery (facility subtype: 322) for injection phase (facility subtypes: 501, 503, 506, and 510) See Section 8.4.10	Primary production and water flood See Section 8.4.10	> 16.9	Gas only	Per test/group meter per pool	Annually
			≤ 16.9	Gas only	Per test/group meter per pool	Biennially
		Miscible/im miscible flood See Section 8.4.11	Production	Gas only	Per test/group meter	Per approval or quarterly (if not in approval)
	Injection		Per meter	Per approval or monthly (if not in approval)		
<b>Heavy oil<sup>1</sup> batteries</b> See Section 8.4.12	Single-well battery (facility subtypes: 325 and 326)		≤ 2.0	Gas only	Per well	Initial <sup>2</sup>
			> 2.0	Gas only	Per meter	Biennially
	Multi-well proration battery (facility subtypes: 327 and		≤ 2.0	Gas only	Per test/group meter per pool	Initial <sup>2</sup>

	Type of facility	Gas rate 10 <sup>3</sup> m <sup>3</sup> /d	Sample and analysis type	Sampling point	Frequency
	344)	> 2.0	Gas only	Per test/group meter per pool	Biennially
<b>Water source well/battery</b>	Single-well/ multi-well battery (facility subtypes: 906 and 907)		Gas only (if present)	Per well	Initial <sup>2</sup>

1. Heavy oil is crude oil with a density  $\geq 920.0$  kg/m<sup>3</sup> at 15.0 °C
2. Initial - an analysis is required within the first six months of operation only, with no subsequent updates required.

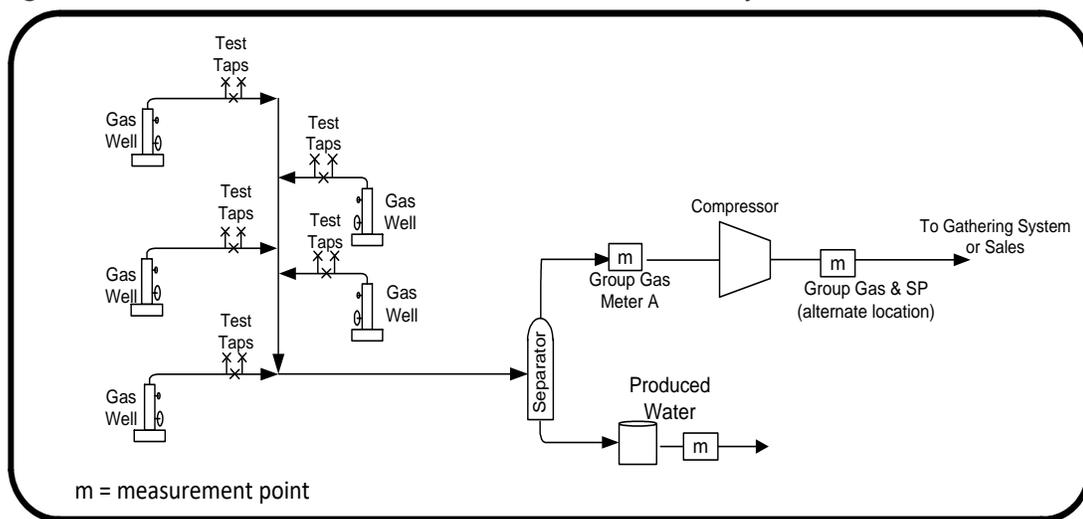
Licensees must ensure that analysis data are used to update volumetric calculations by the end of the month following the receipt of the analysis report. The only exception is for effluent wells, for which the analysis must be updated by the end of the second month following receipt of the analysis report. If sampling and analysis are conducted for other purposes, such as joint venture and allocation agreements, more frequently than required by this Directive, the licensee must use those data to update volumetric calculations.

#### 8.4.1 Gas Multi-well Proration SW Saskatchewan with Minimal Water (Petrinex facility subtypes: 363)

Shallow gas wells are those that produce from shallow gas stratigraphic units or zones, including coals and shales from the bottom of the Glacial Drift to the bottom of the Upper Cretaceous. The production from two or more of these stratigraphic units or zones without segregation in the wellbore requires prior approval from ER for commingled production.

In the configuration of Figure 8.1 of shallow gas wells or CBM wells with water production  $\leq 0.01$  m<sup>3</sup>/10<sup>3</sup>m<sup>3</sup> gas, analyses must be updated biennially at group gas meter A. No sampling and analysis are required at the test meter or well. Density and component analysis data from the group meter sample point may be used for test meter calculations.

Figure 8.1 Gas Multi-well Proration SW Saskatchewan Battery



For shallow gas wells that have been fractured or stimulated using a gaseous medium (e.g., N<sub>2</sub> or CO<sub>2</sub>), gas sampling and analysis frequency must be in accordance with the following:

1. An initial sample and analysis must be obtained within the first month the well is put on production to establish the initial N<sub>2</sub> or CO<sub>2</sub> concentration and other component composition.
2. Where there is adequate analog sample analysis data that is representative of how concentrations of N<sub>2</sub> or CO<sub>2</sub> will decline from month to month in the produced gas, the monthly analog sample analysis data may be used to calculate well volumes in the second to fifth months.
  - a. The analog data set must contain monthly sample analysis data showing how N<sub>2</sub> or CO<sub>2</sub> concentrations decline month over month for up to 12 months and from at least six wells within an eight km radius of the subject well. The analog data set must be a volume-weighted average composition.
3. After being on production for six months, a second sample and analysis must be obtained to confirm that the well gas N<sub>2</sub> or CO<sub>2</sub> concentration is declining as predicted by the analog data set. The sample analysis must be used to re-establish the well gas composition with the analog data set. The monthly analog sample analysis data may be used to calculate well volumes from the seventh to twelfth months.
4. After being on production for one year, the sample and analysis from the group separator may be used to determine the well gas volumes.
5. If analog sample analysis data does not exist as described above, then the wells must be sampled bimonthly until the well gas composition has stabilized; the sample analysis obtained at the group meter may then be used to determine well gas volumes. Composition stabilization means that the mole fraction of N<sub>2</sub> or CO<sub>2</sub> in the total sample analysis is  $\leq 0.05$  or  $\leq 0.02$ , respectively.

If these stratigraphic units or zones are commingled with gas from other outside stratigraphic units or zones, sampling must be conducted biennially on a per pool or stratigraphic unit or zone basis per commingled SW Saskatchewan pools basis from a representative well within the pool.

For exceptions on sampling for these pools or stratigraphic units or zones, see [Section 8.4.1.1](#).

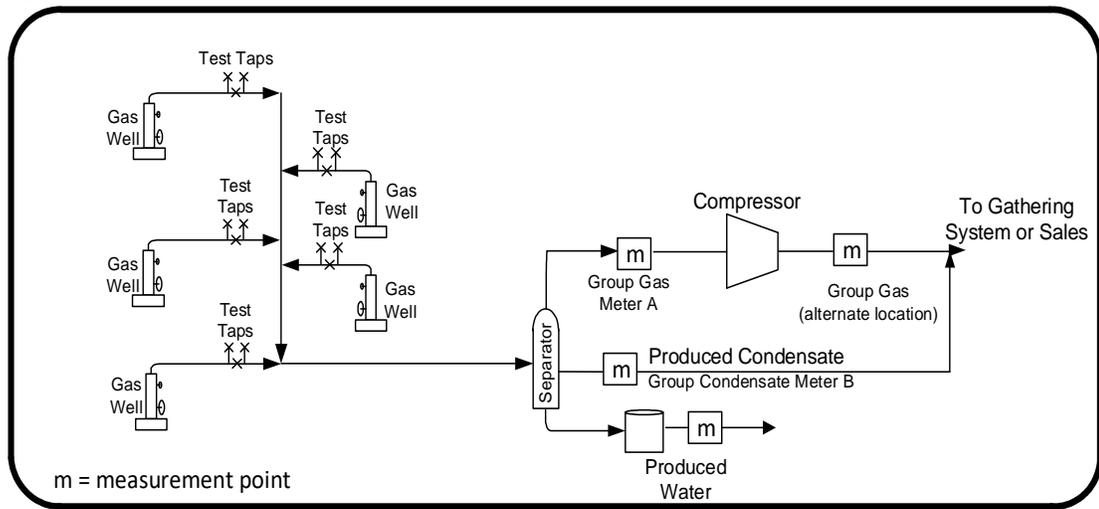
#### **8.4.1.1 Gas Multi-well Proration Outside SW Saskatchewan Batteries (Petrinex facility subtypes: 364)**

In the configuration in [Figure 8.2](#), gas density and composition must be updated

- Annually at group meter A if the daily gas rate is  $> 16.9 \times 10^3 \text{m}^3$ ,
- Biennially at group meter A if the daily gas rate is  $\leq 16.9 \times 10^3 \text{m}^3$ , and
- At each well during a well test.

If condensate at the group measurement point is recombined and delivered to a gas plant, the condensate GEF must be updated

- Annually at group meter B if the daily gas rate is  $> 16.9 \times 10^3 \text{m}^3$ , and
- Biennially at group meter B if the daily gas rate is  $\leq 16.9 \times 10^3 \text{m}^3$ .

**Figure 8.2 Gas Multi-well Proration Outside SW Saskatchewan battery**

#### 8.4.1.2 Exceptions for Sampling Frequency

1. A licensee is not required to update the analyses where three consecutive gas relative density (RD) determinations conducted at the specified determination frequency or, alternatively, no more frequently than once per year are all within  $\pm 1.0$  per cent of the average of the three RD's as shown in Example 8.1. In this situation, there is no need for an application to be submitted to ER. Records and data in support of this exception must be retained by the licensee and made available to ER upon request. Notwithstanding this exception, the licensee must update the gas analyses when changes are made to operating conditions that could affect the gas density by more than  $\pm 1.0$  per cent of the average of the three qualifying RD's.
2. The gas and condensate analyses determined at the group measurement points may be used for the test meters, provided that all wells are from the same pool.
3. For wells producing from multiple pools into a group measurement point, average individual gas and (where applicable) condensate analyses may be used in volume calculations for all the wells in each individual pool (or commingled pool) producing to a test meter provided the following qualifying criteria are met.
  - a. All wells flowing to the group measurement point have common ownership. If there is not common ownership, written notification has been given to all working interest participants, with no resulting objection received.
  - b. All wells flowing to the group measurement point have common Crown or common Freehold royalty. If there is no common Crown or common Freehold royalty and only Freehold royalties are involved, written notification has been given to all Freehold royalty owners, with no resulting objection received. If there is a mix of Freehold and Crown royalties involved, the licensee must apply to ER for approval if any Freehold royalty owner objects after being notified by the licensee.
  - c. The relative densities of the gas and condensate analyses of all common-pool wells (or commingled-pool wells) are within  $\pm 2.0$  per cent of the calculated average gas and condensate relative densities. Initially, the

average relative densities must be calculated based on the most current sample analysis for each well in the pool (or commingled pool).

- i. In each subsequent year, gas and condensate analyses must be obtained from at least four wells in each pool (or commingled pool) or at least 25.0 per cent of the wells in each pool (or commingled pool), whichever is greater. This new data will be used to recalculate average gas and condensate relative densities and, provided that the newly sampled well gas and condensate relative densities variance remains within the  $\pm 2.0$  per cent limit of the average, the exception will remain in effect (see example 8.2). If the well gas and condensate relative densities variance exceeds the  $\pm 2.0$  per cent limit, this exception is revoked. The revocation of the exception remains in place until sampling and analysis of all wells in the pool re-establishes the required RD variance.
- ii. If the pool (or commingled pool) has four or fewer wells flowing to the group measurement point, then, in each subsequent year, gas and condensate analyses must be obtained from one well from each pool (or commingled pool). If the newly sampled well gas and condensate relative densities variance remains within  $\pm 2.0$  per cent of the previous years' relative densities, the exception remains in effect. If the well gas and condensate relative densities variance exceeds the  $\pm 2.0$  per cent limit, this exception is revoked. The revocation of the exception remains in place until sampling and analysis of all wells in the pool re-establishes the required RD variance.
- d. New wells that have been fractured or stimulated with a gaseous medium are not eligible for this exception until their gas and condensate RDs are within the  $\pm 2.0$  per cent variance of the calculated pool average gas and condensate RD.

#### Example 8.1. Meter A RD differences

Sample date	RD	RD Difference from average
June 03, 2000	0.583	- 0.29%
June 09, 2001	0.586	+ 0.22%
June 06, 2002	0.585	+ 0.05%
Average	0.5847	

In this example, meter A would be exempt from the requirement for future updates as the three consecutive RD's are within  $\pm 1.0$  per cent of the average of the three RD's.

#### 8.4.2 Multi-well Effluent Measurement Battery (Petrinex facility subtype: 362)

In the configuration in [Figure 8.3](#) and [Figure 8.4](#), gas analyses, condensate composition, and GEF must be updated at the time of testing each effluent well and as follows:

- Annually at the group gas and condensate meters if the daily group gas rate is  $> 16.9 \times 10^3 \text{m}^3$

- Biennially at the group gas and condensate meters if the daily group gas rate is  $\leq 16.9 \times 10^3 \text{m}^3$

The gas analysis to be used for volumetric calculation at the effluent meter is as follows:

- Option 1: Use the separated gas analysis from the ECF-WGR test; or
- Option 2: Use the recombination of the gas analysis and the condensate analysis from the ECF-WGR test.

Note: All wells within the effluent battery must use the same analysis option.

#### 8.4.2.1 Exceptions for Sampling Frequency

1. A licensee is not required to update analyses at the group gas and condensate metering points if:
  - a. Three consecutive gas RD determinations conducted at the specified determination frequency or no more frequently than once per year are all within  $\pm 1.0$  per cent of the average of the three RD's as shown in Example 8.1; and
  - b. The daily average liquid condensate volume is  $\leq 2.0 \text{ m}^3/\text{day}$  for all reporting months for the previous three years and/or the GEV of the condensate is  $\leq 2.0$  per cent of the recombined total monthly gas volume.

In these situations, there is no need for an application to be submitted to ER. Records and data in support of these exceptions must be retained by the licensee and made available to ER upon request. Notwithstanding these exceptions, the licensee must update the gas analyses when changes are made to operating conditions that could affect the gas RD by more than  $\pm 1.0$  per cent of the average of the three qualifying RD's, and the licensee must update the condensate analyses if the liquid condensate volume or GEV percentage increases beyond the qualifying limits.

2. A licensee is not required to update the well gas and condensate analyses if three consecutive calculated recombined relative density (RD) determinations conducted at the specified determination frequency or, alternatively, no more frequently than once per year are all within  $\pm 1.0$  per cent of the average of the three RDs (see Example 8.1). Specifically, the RDs would be those of the gas plus gas equivalent of the recombined liquid condensate. After the well has fallen below the critical lift velocity as determined using the Turner Correlation calculation described in Section 7.4.1.1 a new gas and condensate analysis must be obtained.

In this situation, there is no need for an application to be submitted to ER. Records and data in support of this exception must be retained by the licensee and made available to ER upon request. Notwithstanding this exception, the licensee must update the gas and condensate analyses when changes are made to producing conditions that could affect the gas density by more than  $\pm 1.0$  per cent of the average of the three qualifying recombined RDs.

Figure 8.3 Gas multi-well effluent measurement battery

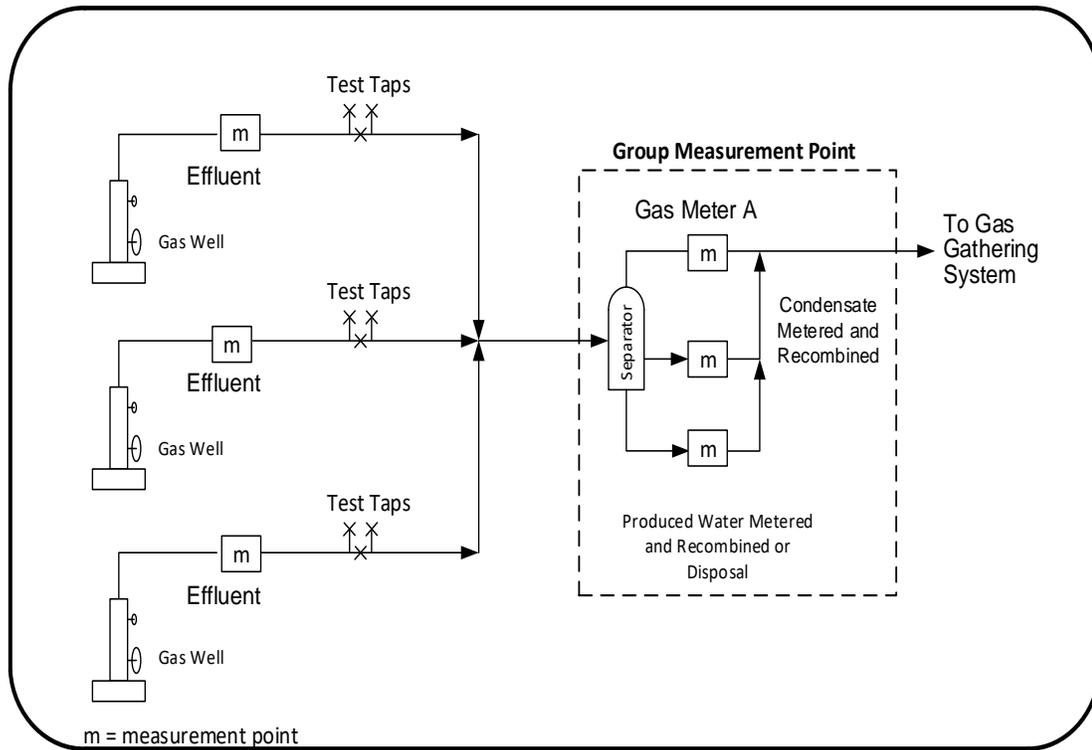
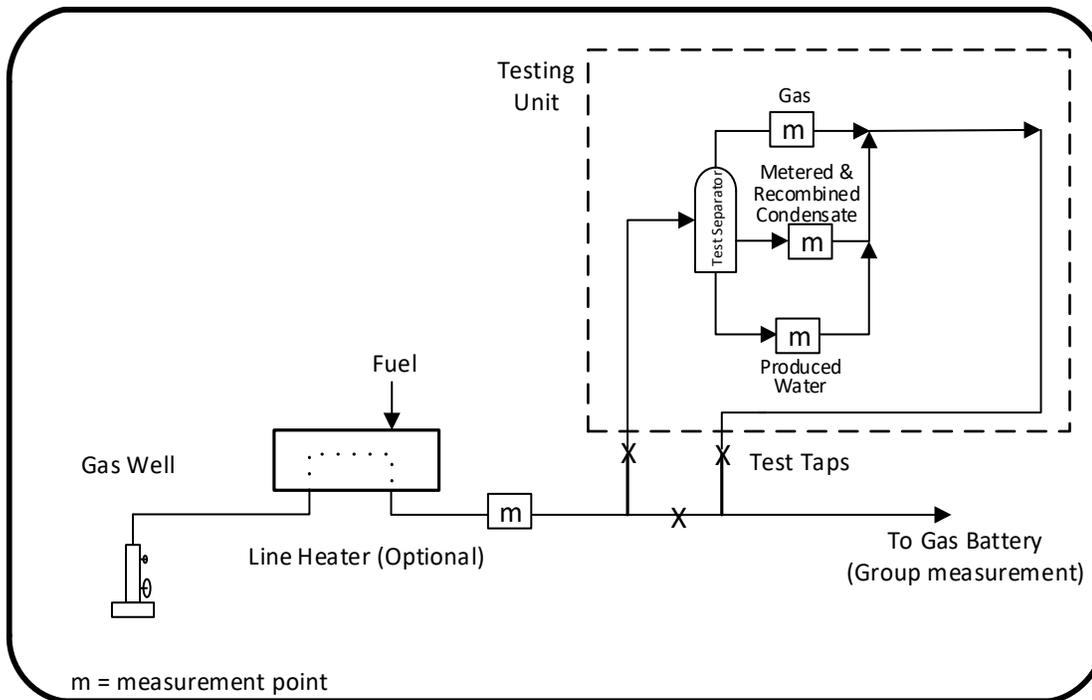


Figure 8.4 Effluent measured well testing configuration



### 8.4.3 Gas Single Well Battery (Petrinex facility subtype: 351) or Gas Multi-well Group Battery (Petrinex facility subtype: 361) and Shallow Gas Wells Producing no Condensate or Oil

For the configuration in Figure 8.5, the gas analysis must be updated within the first year and then biennially at each well gas meter A.

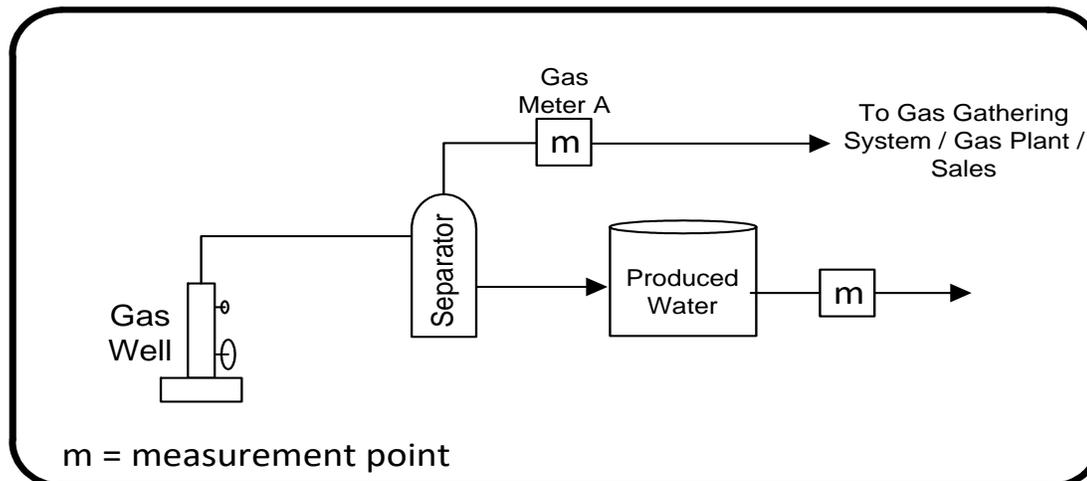
Shallow gas wells are those that produce from the shallow gas stratigraphic units and include coals and shales from the bottom of the Glacial Drift and the bottom of the Upper Cretaceous. The production from two or more of these stratigraphic units without segregation in the wellbore requires prior approval from ER.

For shallow gas wells that are linked to Petrinex facility subtype 351 or 361 and have been fractured or stimulated using a gaseous medium (e.g., N<sub>2</sub> or CO<sub>2</sub>), frequency of gas sampling and analysis must be in accordance with the following:

1. An initial sample and analysis must be obtained in the first month the well is put on production to establish the initial N<sub>2</sub> or CO<sub>2</sub> concentration and other component composition.
2. Where there is adequate analog sample and analysis data that is representative of how concentrations of N<sub>2</sub> or CO<sub>2</sub> will decline from month to month in the produced gas, the monthly analog sample and analysis data may be used to calculate well volumes from the second to fifth months.
  - a. The analog data set must contain monthly sample and analysis data showing how N<sub>2</sub> or CO<sub>2</sub> concentrations decline month over month for up to 12 months and from at least six wells within an eight km radius of the subject well. The analog data set must be a volume-weighted average composition.
3. After being on production for six months, a second sample and analysis must be obtained to confirm that the well gas N<sub>2</sub> or CO<sub>2</sub> concentration is declining as predicted by the analog data set. The sample and analysis must be used to re-establish the well gas composition with the analog data set. In the seventh month and for the duration of the well life cycle, the analog sample and analysis data may be used to determine the well gas volumes.
4. If analog sample and analysis data does not exist as described above, then the well must be sampled bimonthly until the well gas composition has stabilized; after that, no further sampling of the well is required. Composition stabilization means that the mole fraction of N<sub>2</sub> or CO<sub>2</sub> in the total sample analysis is  $\leq 0.05$  or  $\leq 0.02$ , respectively.

For shallow gas wells and coalbed methane wells that have not been fractured or stimulated using a gaseous medium, only a single gas sample and analysis is required through the entire producing life cycle of the well. The operator may determine the timing of the gas sampling, but it must be obtained within the first year of the well being placed on production. A representative sample analysis from an analog well or a calculated average gas composition based on the sample analyses of several analog wells may be used for gas volume determination until the actual well gas sample and analysis are obtained.

**Figure 8.5 Gas single well battery or gas multi-well group battery and shallow gas wells with no condensate or oil production**



#### 8.4.3.1 Exceptions for Analysis Frequency

1. A licensee is not required to update the analyses if three consecutive gas analyses conducted at the specified determination frequency or, alternatively, no more frequently than once per year are all within  $\pm 1.0$  per cent of the average RD of the three analyses as shown in [Example 8.1](#). In this situation, there is no need for an application to be submitted to ER. Records and data in support of this exception must be retained by the licensee and made available to ER upon request. Notwithstanding this exception, the licensee must update the gas analyses when changes are made to operating conditions that could affect the gas analysis by more than  $\pm 1.0$  per cent of the average RD of the three qualifying analysis.
2. A representative analysis for all wells producing to a common gathering system or facility from a common pool can be used if the RD of all common-pool wells are within  $\pm 2.0$  per cent of the average analysis of those wells. Gas analyses must initially be obtained for all the common-pool wells to arrive at the average analysis. Subsequent analyses can be made on 25.0 per cent or at least four wells from the pool (whichever is greater) at the frequency stated in this Directive, provided that the RD variance remains within the  $\pm 2.0$  per cent limit of these wells as shown in [Example 8.2](#). Should the variance exceed this limit, this exception is revoked and biennial analyses must be determined for each measurement point.

#### Example 8.2 Pool RD differences

Consider an 8-well pool producing gas under this configuration:

Well Id	RD	RD Difference from average
11-14	0.602	-0.99%
10-16	0.610	+0.33%
10-21	0.602	-0.99%
9-27	0.616	+1.32%
11-30	0.608	0.00%
6-31	0.616	+1.32%
11-32	0.606	-0.33%

Well Id	RD	RD Difference from average
11-16	0.604	-0.66%
Average	0.608	

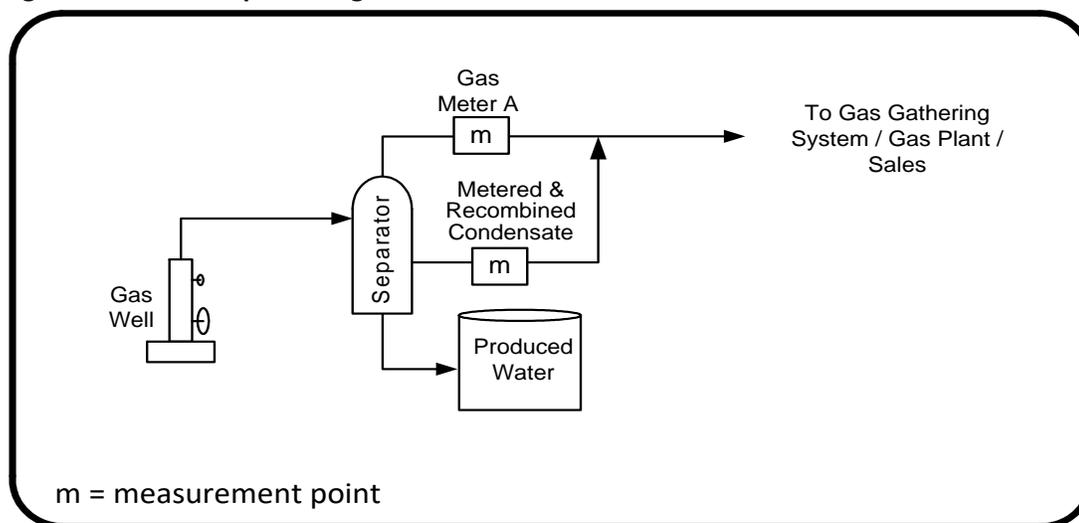
In this scenario, it is acceptable to use the analyses from the well with the RD closest to the average, Well Id 11-30, for all well meters, as all RDs are within  $\pm 2.0$  per cent of the average of all well RDs. The analysis must then be updated biennially for at least four wells from the pool. This exception will remain in place, provided that all four well RD's continue to be within  $\pm 2.0$  per cent of the average of all the updated RD's. When this criteria is not met, sampling and analysis must revert to biennial updates for all wells.

A permanent exception on a pool basis would be available where the updated average RD meets the criterion of Exception 1. Where practical, ER expects the same wells to be used to arrive at the average RD used in pursuit of this exception.

#### 8.4.4 Gas Single Well Battery (Petrinex facility subtype: 351) or Gas Multi-well Group Battery (Petrinex facility subtype: 361) with Condensate or Oil Production

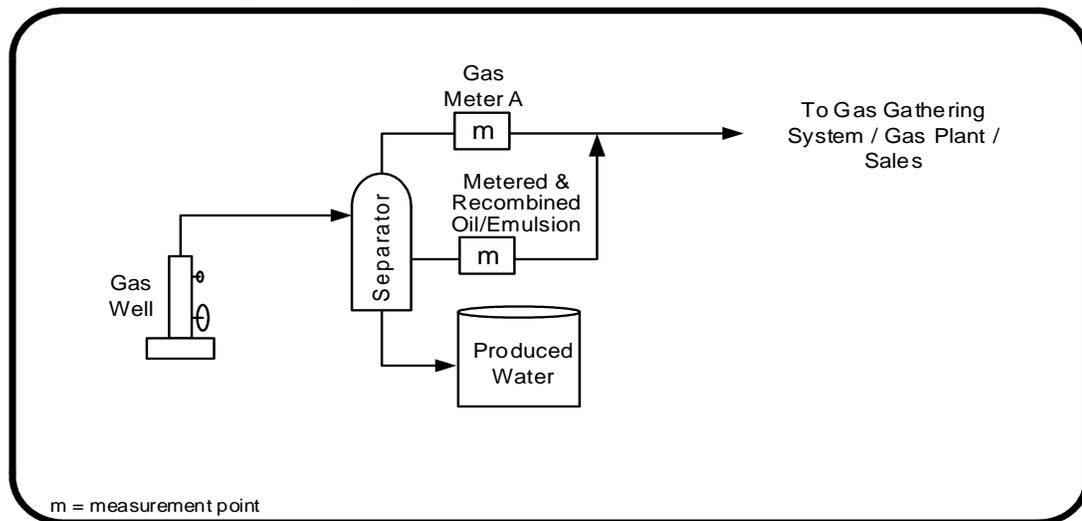
For gas wells producing condensate in [Figure 8.6](#), the frequency of sampling and analysis for gas and condensate depends upon the gas flow rate through gas meter A plus the GEV of condensate. If the flow rate is  $> 16.9 \times 10^3 \text{m}^3/\text{day}$ , the frequency is annual. If the flow rate is  $\leq 16.9 \times 10^3 \text{m}^3/\text{day}$ , the frequency is biennial. The flow rate value is to be based on a rolling 12 month average.

Figure 8.6 Gas well producing condensate



For gas wells producing oil in [Figure 8.7](#), the sampling and analysis of oil/emulsion streams to determine relative oil and water content must conform to the requirements in Sections 6.4.3 and 14.8. The gas sampling frequency is the same as for a gas well producing condensate, except that the total gas flow rate does not include GEV of oil/emulsion.

Figure 8.7 Gas well producing oil



#### 8.4.4.1 Exceptions for Gas Wells with Condensate

1. For gas wells producing condensate, a licensee is not required to update the gas and condensate (if applicable) analyses at the metering points if:
  - a. Three consecutive gas RD determinations conducted at the specified determination frequency or no more frequently than once per year are all within  $\pm 1.0$  per cent of the average of the three RDs as shown in [Example 8.1](#); and
  - b. Daily average liquid condensate volume is  $\leq 2.0 \text{ m}^3/\text{day}$  for all reporting months for the previous three years and/or the GEV of the condensate is  $\leq 2.0$  per cent of the recombined total monthly gas volume.

In these situations, there is no need for an application to be submitted to ER. Records and data in support of these exceptions must be retained by the licensee and made available to ER upon request. Notwithstanding these exceptions, the licensee must update the gas analyses when changes are made to producing conditions that could affect the gas RD by more than  $\pm 1.0$  per cent of the average of the three qualifying RDs, and the licensee must update the condensate analyses if the liquid condensate volume or GEV percentage increases beyond the qualifying limits.

2. A representative analysis for all wells producing to a common gathering system or facility from a common pool may be used if the RD's of all common-pool wells are within  $\pm 2$  per cent of the average RD of those wells. Gas analyses must initially be obtained for all the common-pool wells to arrive at the average RD. Subsequent analyses may be made on 25.0 per cent or at least four wells from the pool, whichever is greater, at the frequency stated in this Directive, provided that the RD variance remains within the  $\pm 2$  per cent limit of these wells as shown in [Example 8.2](#). Should the variance exceed this limit, this exception is revoked and the sampling frequency will revert to the base requirements and analyses must be determined for each measurement point.

### 8.4.5 Underground Gas Storage Facility (Petrinex facility subtype: 505)

For the gas storage configuration shown in [Figure 8.8](#), there are two operational phases to consider:

#### 1. Storage Injection Phase

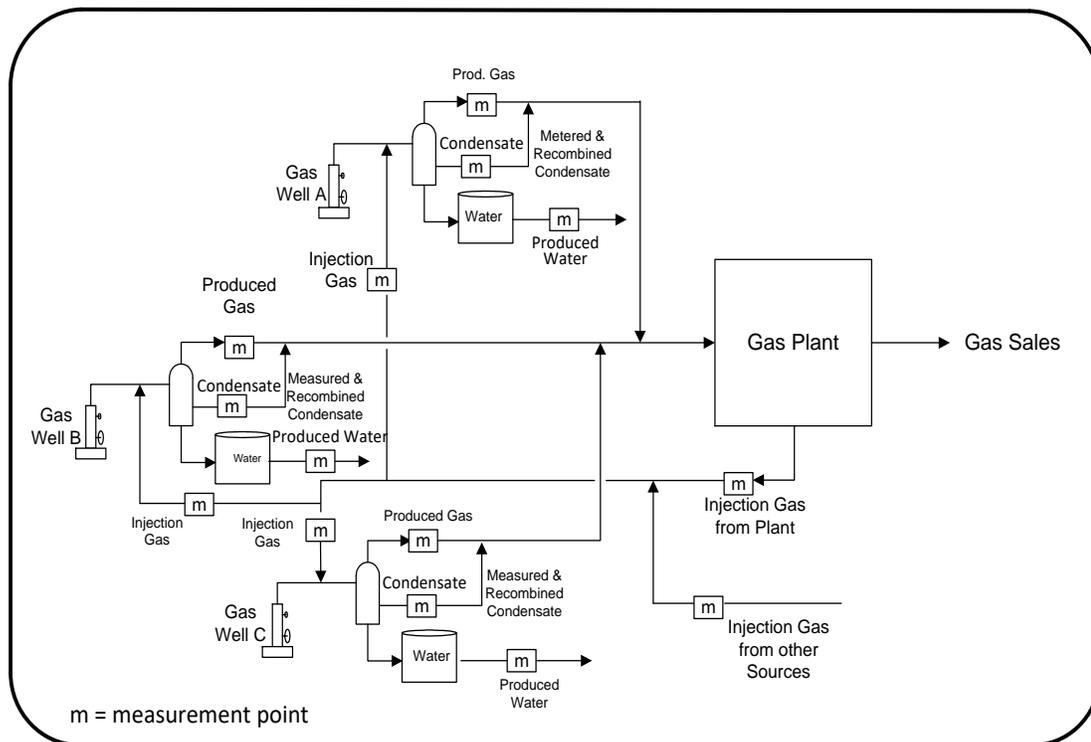
If the injection gas only comes from a single source, an annual common stream sample analysis may be used for all injection meters, and no individual well injection analyses are required.

If there are multiple injection gas sources, sample analysis is required at each source stream and at each well injection measurement point. In this scenario, the minimum analysis frequency for injection meters is semiannual, however, a continuous proportional sampler or a gas chromatograph should be installed to provide more accurate compositions for gas volume calculations. If a continuous proportional sampler or gas chromatograph is installed and used the sample analysis frequency requirement is considered to be met.

#### 2. Storage Recovery Phase

During each recovery phase, analyses must be updated at each gas well's production meters within the first month and semiannually. If a continuous proportional sampler or gas chromatograph is installed and used the sample analysis frequency requirement is considered to be met.

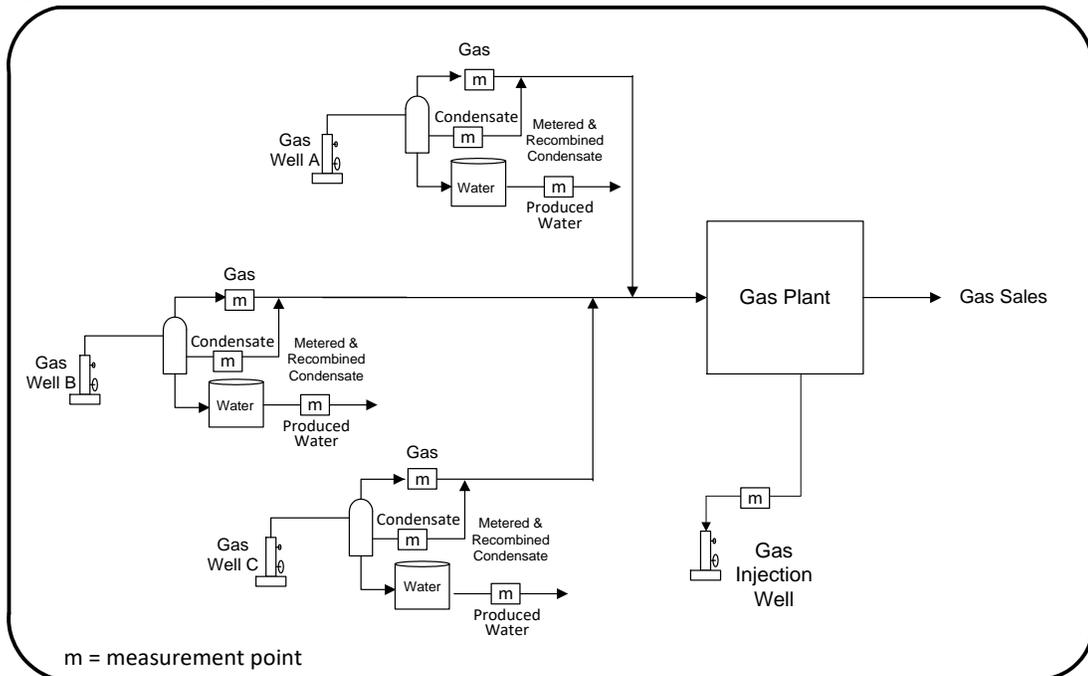
**Figure 8.8 Underground gas storage facility**



### 8.4.6 Gas Cycling Scheme (Petrinex facility subtype: 502)

In the gas cycling scheme configuration shown in Figure 8.9, analyses must be updated at each gas wells production meters, and the injection well meter in accordance with the specific scheme approval. If there are no frequencies specified in the approval, the gas well production meters must have analyses updated semi-annually and the gas injection meter(s) must have analyses updated in accordance with the base requirements such as semiannually for gas plant gas.

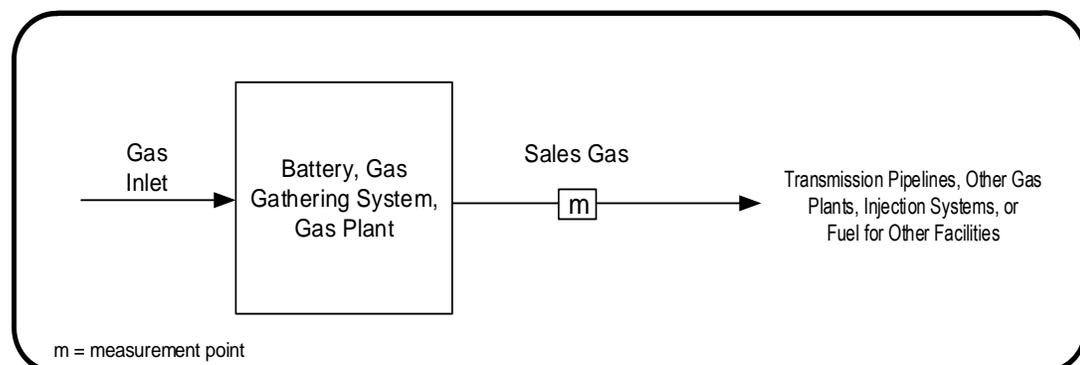
Figure 8.9 Gas cycling scheme



### 8.4.7 Gas Sales/Delivery

In the gas sales/delivery configuration shown in Figure 8.10, gas sales/delivery in this context will typically be clean, processed sales gas that is delivered out of a gas plant or a facility into a transmission pipeline. In some scenarios, this type of gas may be delivered to other plants for further processing or fuel or to injection facilities.

Figure 8.10 Gas sales/delivery



If a meter is used to determine the sales gas/delivery point volume from a battery, gas gathering system, or gas plant, the minimum gas analysis frequency is annual. However, a continuous proportional sampler or a gas chromatograph should be installed to provide more accurate analyses for the gas volume calculation.

#### **8.4.8 Gas Plants (Petrinex facility subtypes: 401 to 407) and Gas Gathering Systems (Petrinex facility subtypes: 621)**

In the configuration shown in [Figure 8.11](#), only one sample point is required for common gas stream volume determination, such as sales gas, which may also be used for fuel, injection, and sales gas flare. Inlet gas sample analysis may be used for inlet gas flare volume determination.

The frequency for sampling and analysis is as follows unless a different frequency has been specified in site-specific approvals, such as gas cycling or miscible/immiscible flood schemes, or for heavy oil well gas production. For gas sales measurement point sampling frequency, see Section 8.4.7.

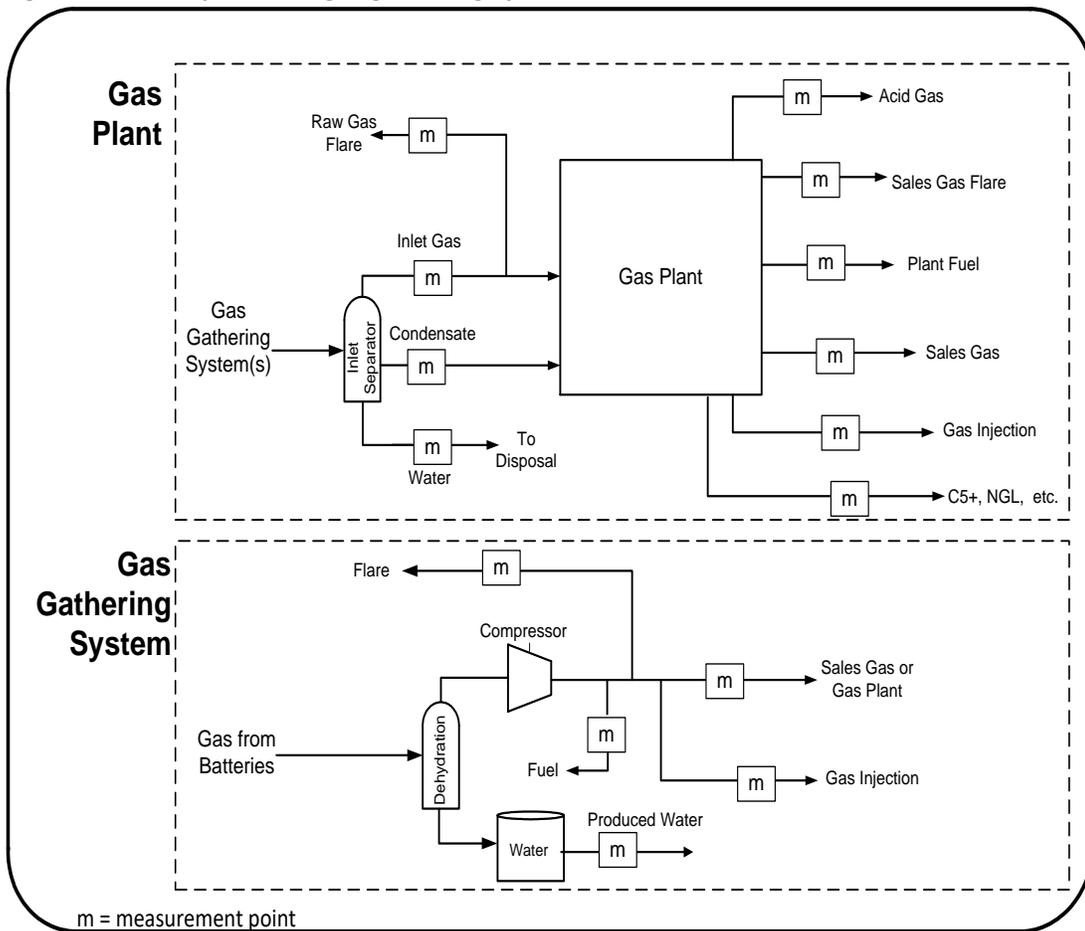
##### **8.4.8.1 Gas Plant**

The minimum frequency for updating analyses at all accounting meters within a gas plant is semiannual. Inlet condensate is reported as a GEV, so condensate analyses are also required. High-vapour pressure liquids, such as C<sub>5</sub>-SP and other NGLs, are to be reported as liquid volumes on Petrinex, which will then perform the GEV calculation automatically using standard factors for plant balancing.

##### **8.4.8.2 Gas Gathering System**

The minimum frequency for updating analyses at all accounting meters within a gas gathering system is annual for flow rates that are  $> 16.9 \times 10^3 \text{m}^3/\text{day}$ . If the flow rate is  $\leq 16.9 \times 10^3 \text{m}^3/\text{day}$ , the frequency is biennial. The flow rate is to be based on a rolling 12 month average. Condensate volumes recombined with gas for delivery to other facilities for processing must be reported as GEV, so analyses are required for updating GEFs. Where condensate is delivered out of a gas gathering system without further processing, it is reported as a liquid volume, but analyses for GEV calculation purposes are required for reporting on Petrinex.

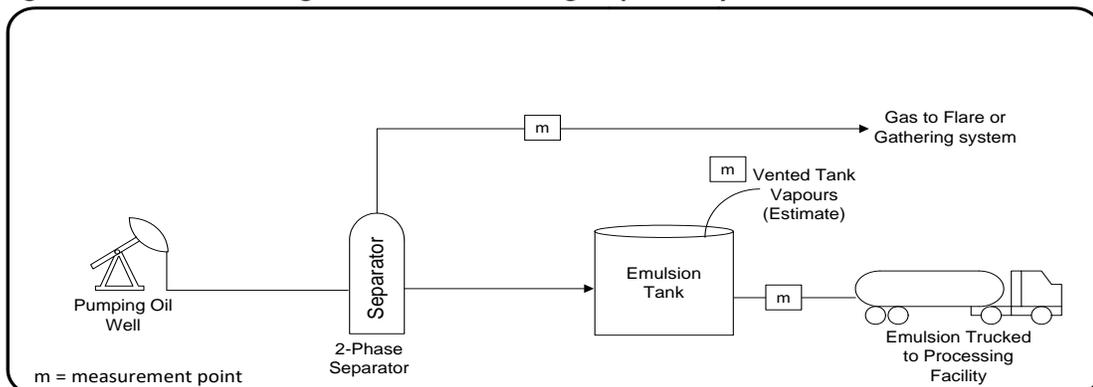
Figure 8.11 Gas plant and gas gathering system



**8.4.9 Crude Oil Single Well Battery (Petrinex facility subtypes: 311) and Crude Oil Multi-well Group Battery (Petrinex subtype: 321)**

In the configuration shown in Figure 8.12, if all associated gas, net of lease fuel, is flared, an initial representative gas analysis is required. If gas is conserved, gas analysis updates are required. If the average flow rate is  $> 16.9 \times 10^3 \text{m}^3/\text{day}$ , the frequency is annual. If the average flow rate is  $\leq 16.9 \times 10^3 \text{m}^3/\text{day}$ , the frequency is biennial.

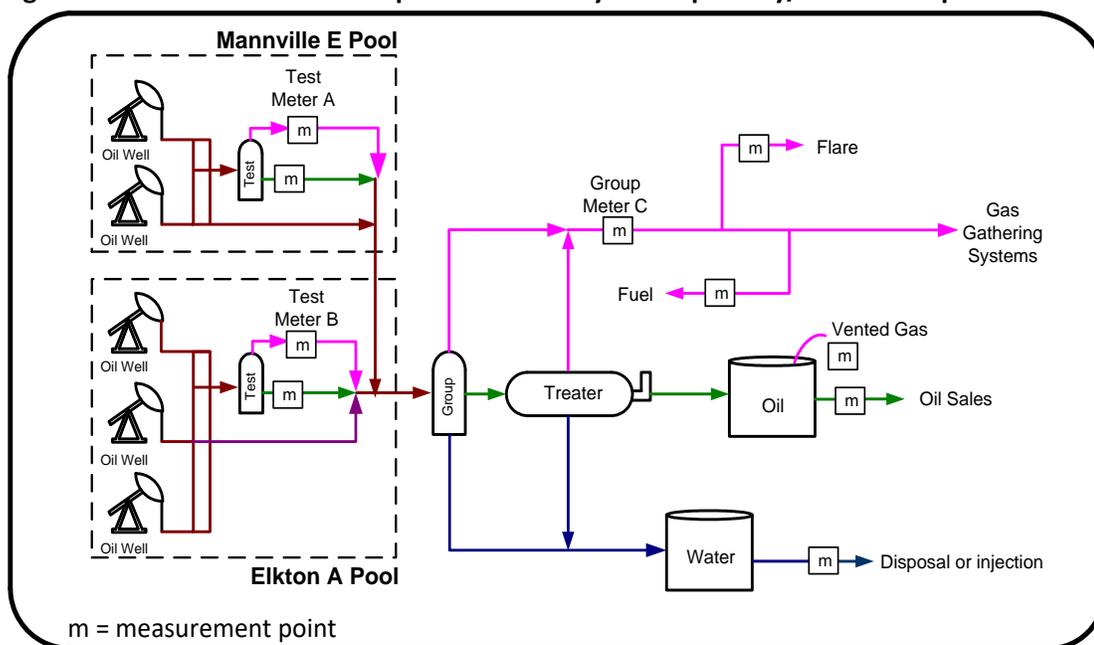
Figure 8.12 Crude oil single-well or multi-well group battery



### 8.4.10 Crude Oil Multi-well Proration Battery (Petrinex facility subtype: 322)

In the configuration shown in Figure 8.13, the gas analyses must be updated at the test meters A and B biennially for test gas rates that are  $\leq 16.9 \text{ } 10^3\text{m}^3/\text{day}$  or annually if the test gas rates are  $> 16.9 \text{ } 10^3\text{m}^3/\text{day}$ .

Figure 8.13 Crude oil multi-well proration battery under primary/waterflood production



It is acceptable to use the gas analysis from a single representative well for all wells within a single pool. If wells from more than one pool are directed through the same test separator, an analysis must be obtained for each pool.

The gas analysis at meter C must be updated annually for gas flow rates that are  $> 16.9 \text{ } 10^3\text{m}^3/\text{day}$  or biennially if the rate through the meter is  $\leq 16.9 \text{ } 10^3\text{m}^3/\text{day}$  based on a rolling 12month average flow rate.

#### Example 8.3 Minimum Gas Analysis Frequency

Consider a five-well proration battery with two wells producing from the Mannville E Pool and three wells producing from the Elkton A Pool. Battery gas production is gathered and conserved.

Pool	Well	Satellite meter	Test gas rate $10^3\text{m}^3/\text{d}$
Mannville E	10-14	Meter A	4.2
Mannville E	10-16	Meter A	6.8
Elkton A	10-21	Meter B	18.0
Elkton A	9-27	Meter B	12.0
Elkton A	10-30	Meter B	6.5
Total rate for Meter C =			47.5

A gas analysis must be established for the Mannville E Pool, as a minimum using either the 10-14 or 10-16 well, and updated biennially at meter A, as the maximum rate through meter A for the Mannville E pool wells is  $\leq 16.9 \text{ } 10^3\text{m}^3/\text{day}$ .

A gas analysis must be determined for the Elkton A Pool at meter B, as a minimum using any one of the three wells, and updated annually, as the maximum rate through meter B for the Elkton A pool wells is  $> 16.9 \times 10^3 \text{m}^3/\text{day}$ .

The gas analysis at meter C must be updated annually, as the flow rate through the meter is  $> 16.9 \times 10^3 \text{m}^3/\text{day}$ .

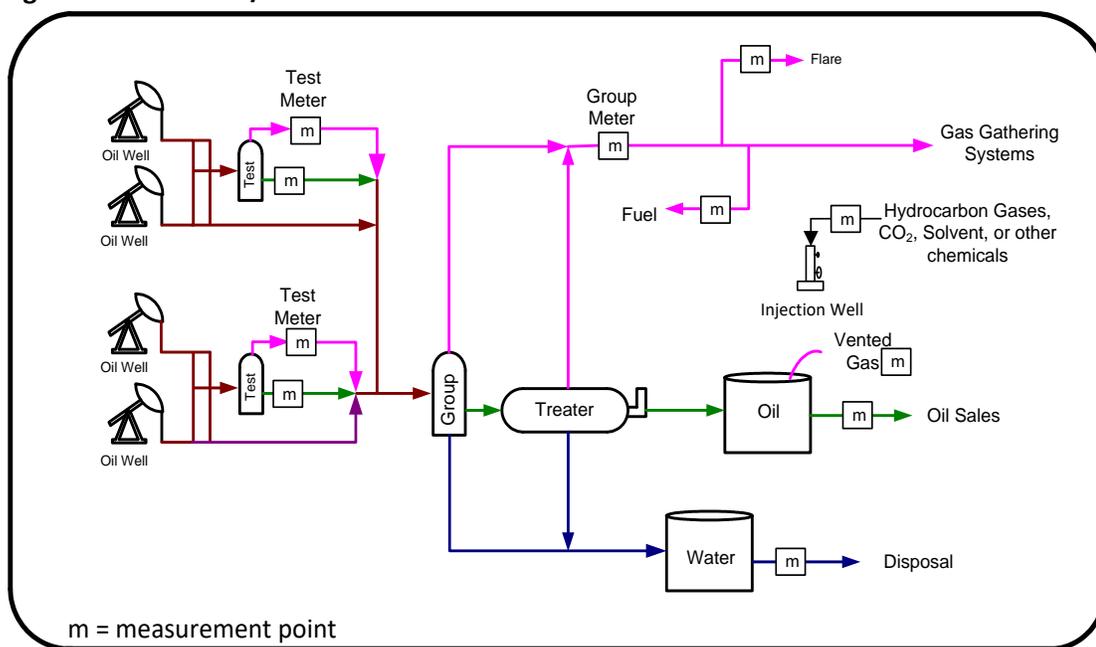
#### 8.4.10.1 Exception for Gas Analysis Frequency

If the total battery gas, net of lease fuel, is flared, an initial pool gas analysis must be determined at meters A and B. Updates of the gas analysis at meter C, at the annual or biennial frequency as determined by the gas flow rate through the meter, is only required if the gas directed through meter C originates from multiple pools. If the gas directed through meter C originates from a single pool, no updates are required subsequent to the initial analysis. However, this exception is revoked as soon as the gas is conserved, and gas analyses must be performed according to the frequencies specified in this section.

#### 8.4.11 Miscible/Immiscible Flood

In the configuration shown in Figure 8.14, analyses must be updated at each test and group meter and the injection well meter in accordance with the specific scheme approval. If there are no frequencies specified in the approval, the test and group meters must have analyses updated quarterly and the injection meter(s) must have analyses updated monthly.

Figure 8.14 Miscible/immiscible flood

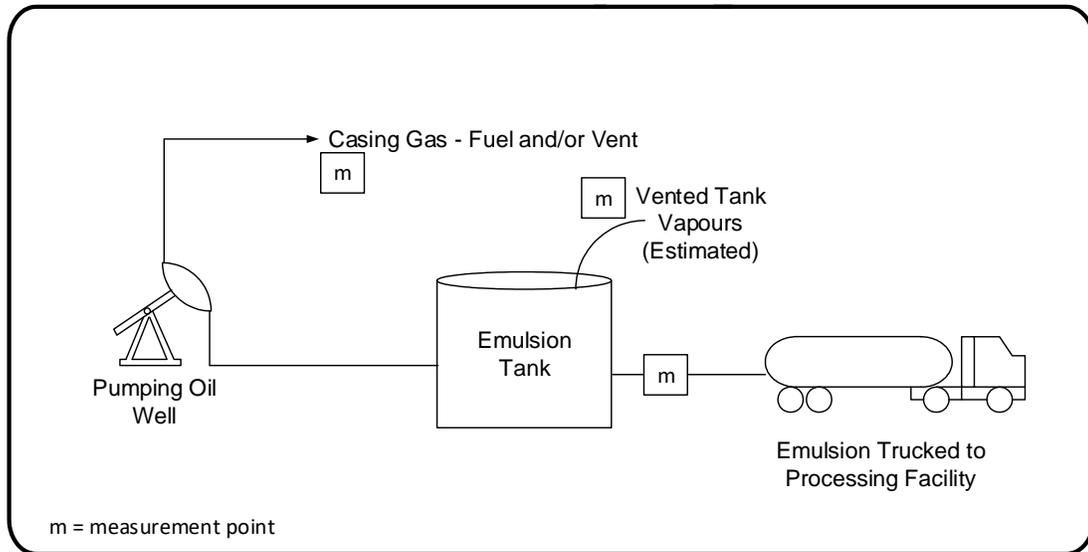


#### 8.4.12 Crude Oil Batteries Producing Heavy Oil (Petrinex facility subtypes: 313, 325, 326, 327, 344)

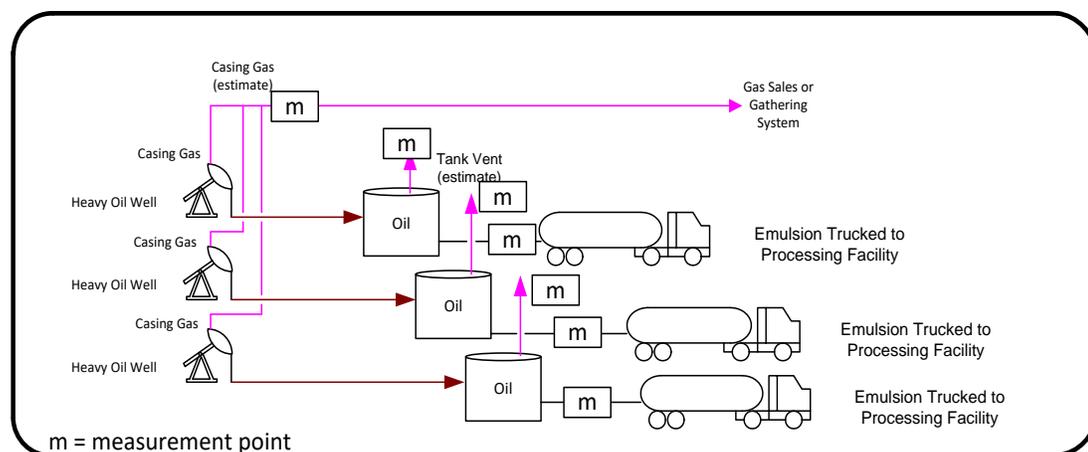
Heavy oil production at a single-well as shown in Figure 8.15 or multi-well group battery as shown in Figure 8.16, or multi-well proration battery in Figure 8.17 typically involves directing all production to a tank without using a separator or gas meter. In these scenarios, gas production may be estimated using a GOR.

If a meter is used to continuously measure gas production and the gas rate is  $\leq 2.0 \times 10^3 \text{ m}^3/\text{day}$ , an initial gas analysis only is required. If the gas production rate is  $> 2.0 \times 10^3 \text{ m}^3/\text{day}$  then an initial gas analysis is required and the analysis must be updated biennially thereafter.

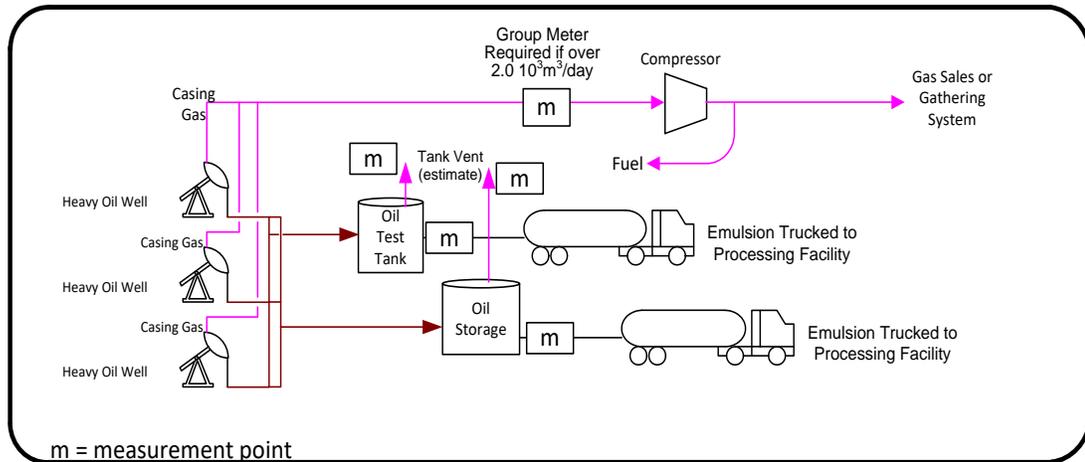
**Figure 8.15 Crude oil single well battery producing heavy oil**



**Figure 8.16 Crude oil multi-well group battery producing heavy oil**



**Figure 8.17 Crude oil multi-well proration battery producing heavy oil**



## 8.5 Oil Sampling and Analysis Requirements

Sampling and analysis must be in accordance with Sections 6, 8, 10, 14 or other equivalent method approved by an appropriate industry standards association.

### 8.5.1 Oil Analysis Requirements for Wells and Facilities

Sampling and analysis of oil/emulsion streams at oil and gas wells and batteries are performed to determine the relative oil and water content of the streams and the density of the oil. Oil/emulsion sampling and analysis are described in Section 6.4.

As per *The Oil and Gas Conservation Regulations, 2012* Section 93.1, the Minister may require an oil sample and analysis to be conducted.

All oil sample analysis reports must be submitted to ER in accordance with the requirements in *Directive PNG013: Well Data Submission Requirements*.

## 9 Cross-Border Measurement

This section presents the measurement requirements for all upstream and midstream oil and gas products crossing a provincial or territorial border.

### 9.1 General Requirements

For those facilities receiving and/or delivering products and waste to another jurisdiction either by truck, rail or pipeline, including pipelines under the National Energy Board (NEB) jurisdiction, each jurisdictional product and waste stream must be separated and measured prior to commingling. The delivery point measurement standards for each jurisdictional authority must be followed, unless site-specific approval from ER and the other jurisdictional authority(ies) has been obtained. All streams must be separated and metered or estimated according to the requirements in this Directive. This can include production, gathering systems, and all fuel, flare, and vent volumes. If the measurement or other equipment requirements for delivery point measurement of hydrocarbon and related fluids from any jurisdiction is different from ER requirements, the higher requirements, such as frequency and accuracy, between the jurisdictions must be followed.

Fuel gas usage at cross-border oil and gas processing facilities must be separately determined and measured if the rate is  $> 0.5 \text{ } 10^3 \text{ m}^3/\text{day}$

for each jurisdiction. If the usage is for production from both jurisdictions, no separate fuel gas metering is required if site-specific approval is obtained from both jurisdictions involved. For example, a compressor used only for gas coming from another jurisdiction into Saskatchewan must be metered separately at the cross-border facility and the fuel gas use for other equipment processing commingled production or the entire facility must be measured with another meter.

### 9.2 Cross-Border Sampling Requirements

Except where otherwise noted, the gas and liquid sampling equipment and methodology must follow the requirements set out in Section 8: Gas and Liquid Sampling and Analysis.

Spot or grab samples are acceptable for obtaining gas and liquid analyses, provided the uncertainty requirements in Section 1 of this Directive are adhered to.

When the uncertainty requirements cannot be met, consider:

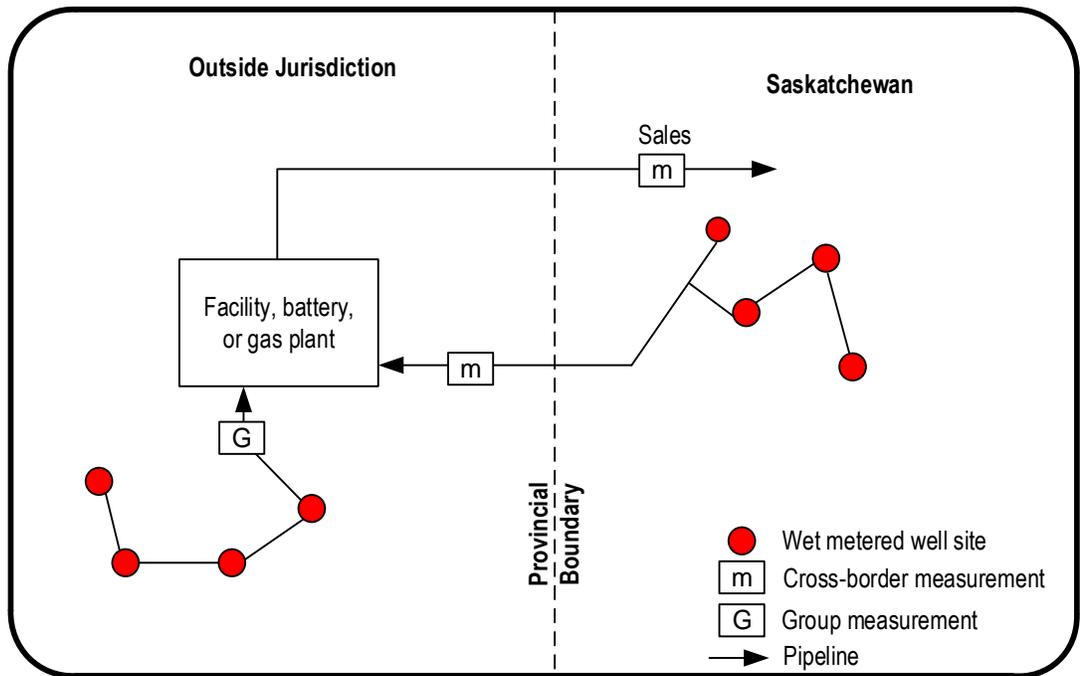
1. More frequent spot sampling for calculated analysis;
2. The use of proportional samplers; or
3. The use of gas chromatographs or other continuous analyzers.

### 9.3 Cross-Border Measurement Points

Figures 9.1 through 9.9 are some of the scenarios to determine if a specific circumstance is considered cross border. Each scenario applies as well if the flow is in the opposite direction. There must be only one cross-border measurement point for each pipeline crossing the provincial boundary unless site-specific approval is obtained from both jurisdictions involved.

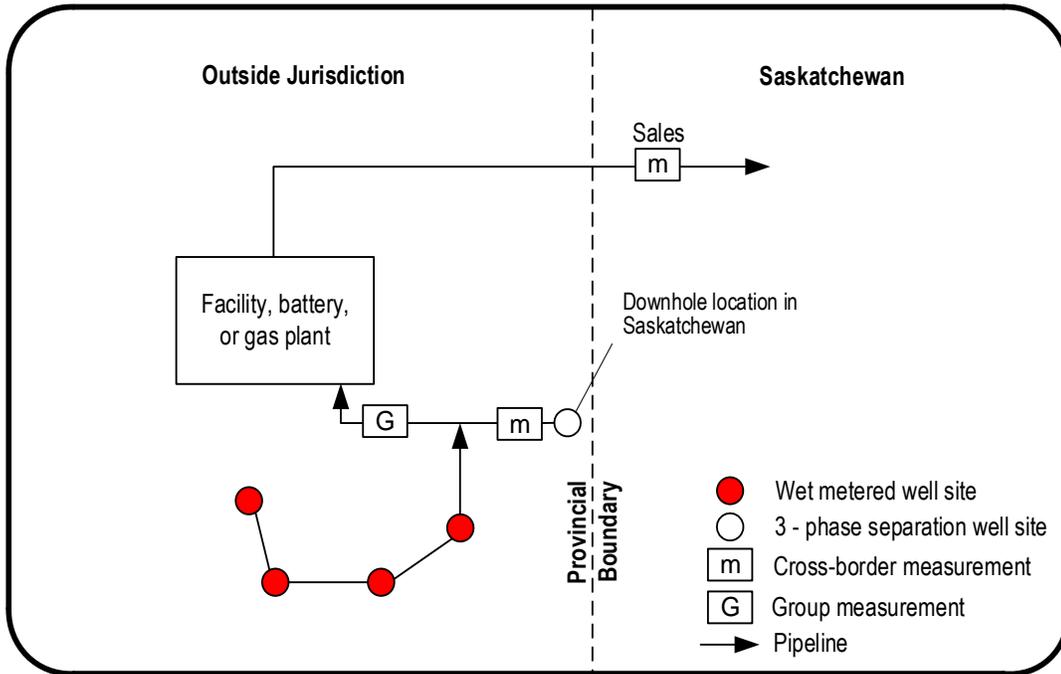
The cross-border measurement point can be on either side of the jurisdictional border before commingling with any fluids from another jurisdiction. Measurement-by-difference rules apply in all situations where there is measured production going into a proration battery.

Figure 9.1 Effluent measured gas wells to an out-of-province location (non-common pool)



For cross-border common pools producing from one or more jurisdictions, if the surface facility is located in one jurisdiction and the well production as defined by the bottom hole location is in another jurisdiction, delivery point measurement of the production is required (see Figure 9.2). The production from this well must be reported as delivered to the other jurisdiction where the surface facility is located.

**Figure 9.2 Gas gathering system of effluent measured gas wells with measured well downhole location in another jurisdiction**



**Figure 9.3 Multiple jurisdictional crossing**

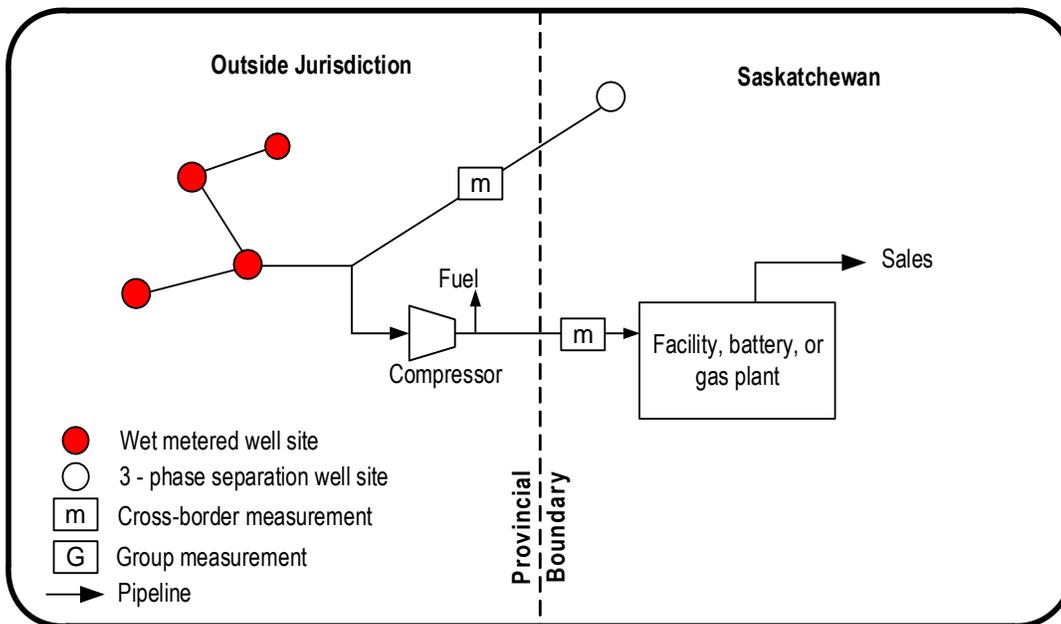


Figure 9.4 Measured gas source from an out-of-province location

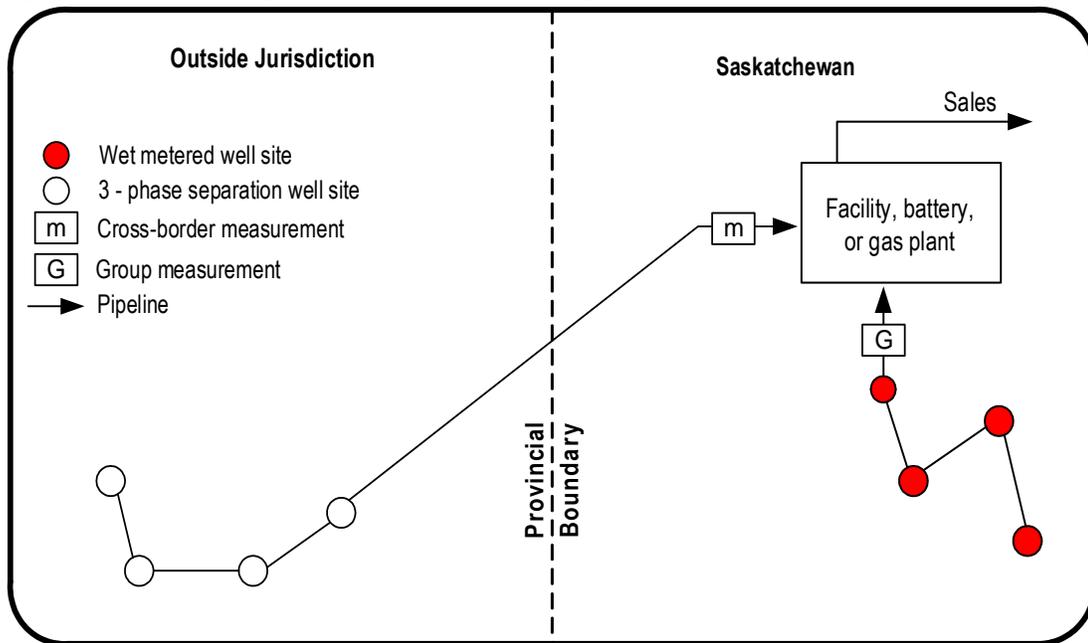


Figure 9.5 Sales gas source from an out-of-province location

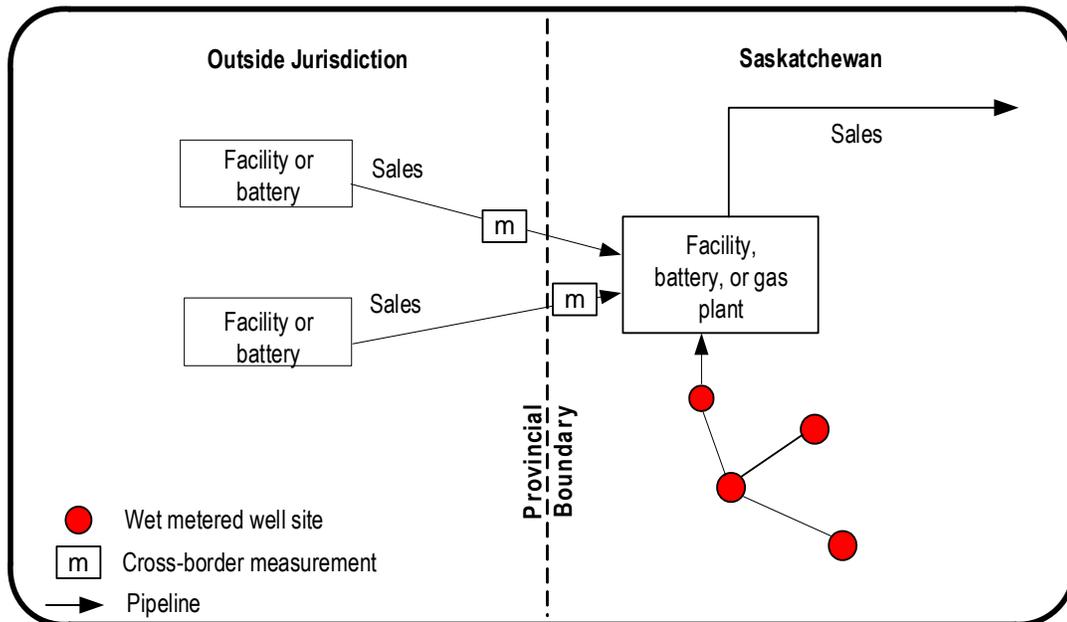
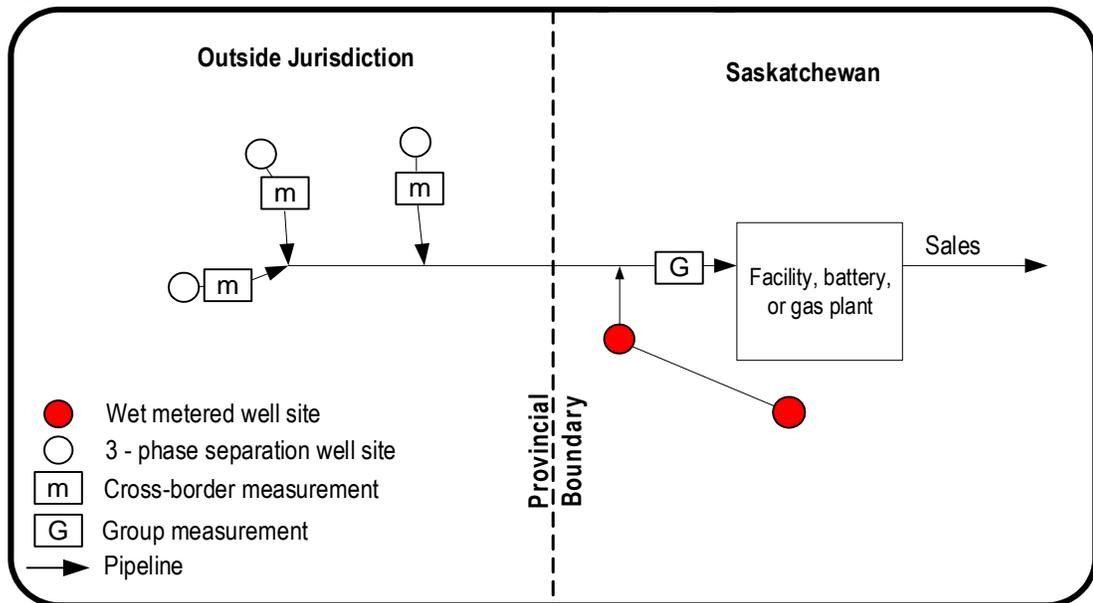


Figure 9.6 Measured sources from an out-of-province location



For Figure 9.6, the three-phase separation wells can be designed to delivery point measurement requirements without another cross-border measurement point.

Measurement-by-difference rules applies in Figures 9.2, 9.5, and 9.6 unless the effluent metered wells have a group measurement point prior to commingling with the measured gas source(s).

Figure 9.7 Sales oil or gas source from an out-of-province location

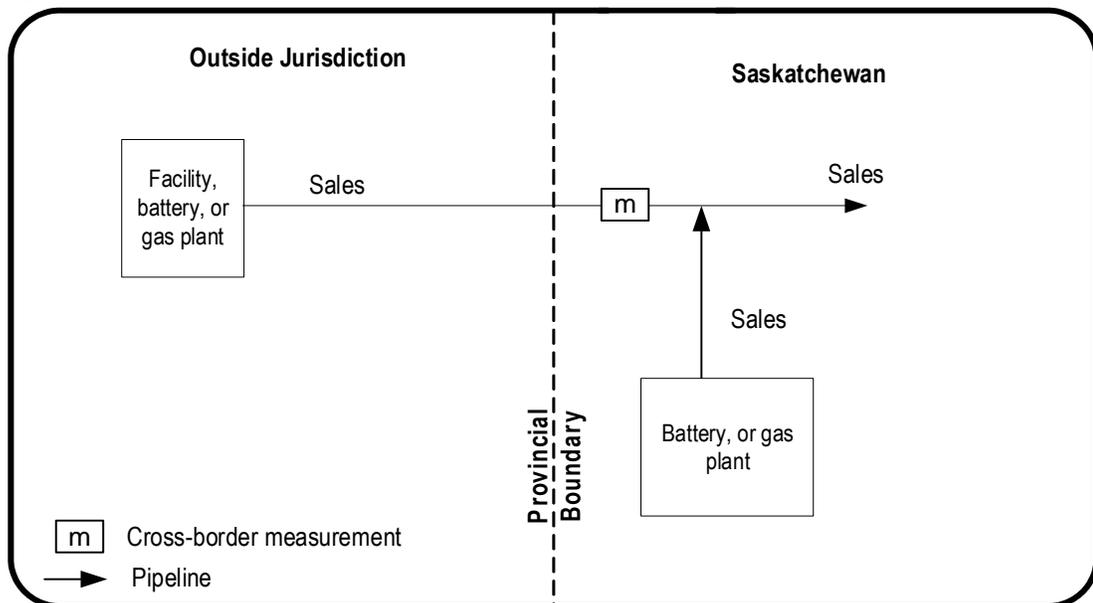


Figure 9.8 Sales oil or gas source from an out-of-province location

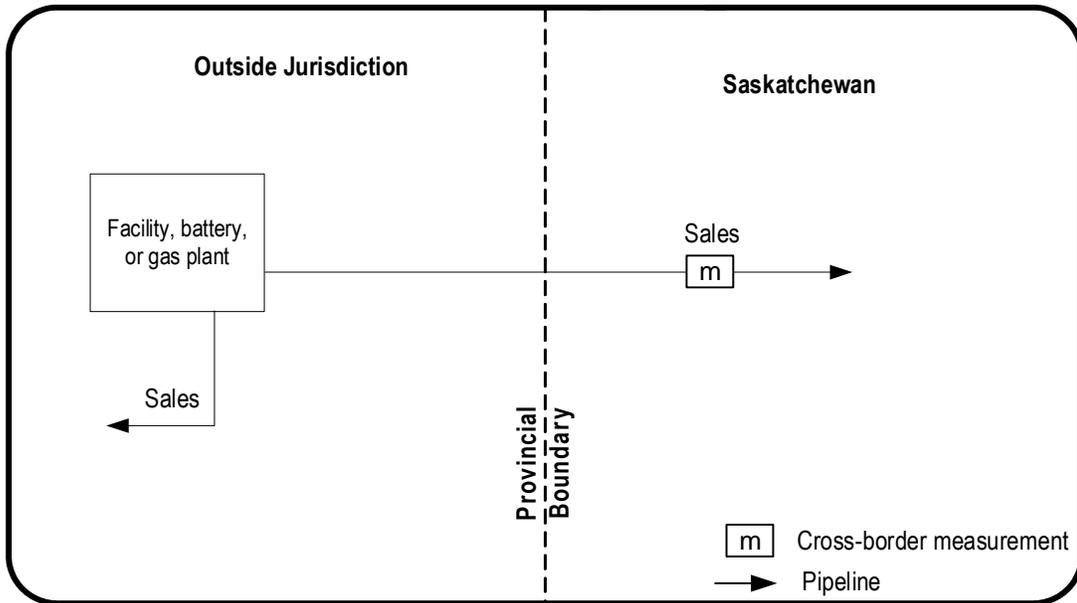
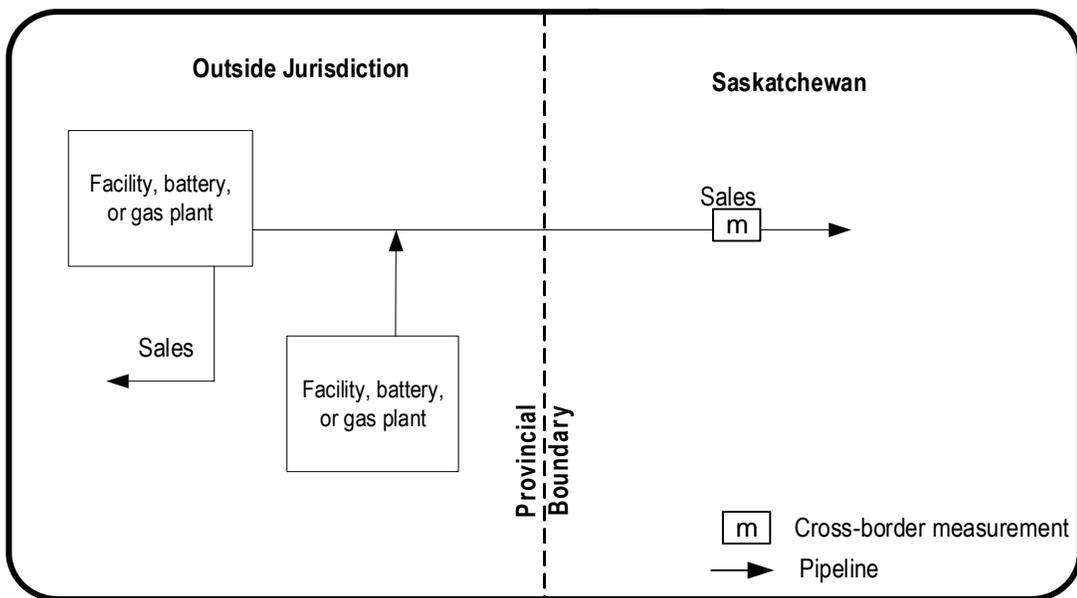


Figure 9.9 Sales oil or gas source from an out-of-province location



## 10 Trucked Liquid Measurement

This section presents the requirements for trucked liquid measurement from oil and gas production facilities to another facility or sales. Applicable liquids include crude oil, condensate, water and NGLs.

### 10.1 General Requirements

Crude oil or condensate may be found in association with water in an emulsion. In such scenarios, the total liquid volume of the trucked load must be measured, and the relative volumes of oil and water in the emulsion must be determined by obtaining and analyzing a representative sample of the emulsion or by using a product analyzer such as a water-cut analyzer or a Coriolis meter's density measurement, where applicable.

A licensee must accurately measure produced liquids/emulsion volumes by using tank gauging, a weigh scale, or a meter, unless otherwise stated in this Directive. The delivery point measurement requirements must be met for all trucked liquids unless the exception conditions in this section are met or a site-specific approval from ER has been obtained as per Section 5.

ER will consider a truck liquid measurement system to be in compliance if the base requirements outlined in Sections 10.1.1, 10.1.2 and 10.1.3 are met. ER may stipulate additional or alternative requirements for any specific situation, based on a site-specific assessment.

All delivery point meters that require delivery point measurement uncertainty must be proved in accordance with Section 2. LACT meters may use the proving procedure in API-MPMS, Chapter 4: Proving Systems, instead of the Section 2 procedure.

#### 10.1.1 Reporting Requirements

Monthly oil, condensate, and water volumes for wells and batteries, for example, (production, receipts, and dispositions) must be reported in m<sup>3</sup> and rounded to the nearest 0.1 m<sup>3</sup>. . Measured volumes must be corrected to 15.0°C and at the greater of 0.0 kPaG or equilibrium vapour pressure at 15.0°C. See Section 6.3.3 for production data verification and audit trail requirements.

For delivery point measurement, hydrocarbon liquid volume must be determined to two decimal places and rounded to one decimal place for monthly reporting. If there is more than one volume determination within the month at a reporting point, the volumes determined to two decimal places must be totaled prior to the total being rounded to one decimal place for reporting purposes.

#### 10.1.2 Temperature Correction Requirements

All delivery point measurement of hydrocarbons and emulsions requires temperature correction to 15.0°C, see Section 6.3.2.1. See Section 14.4 for temperature determination requirements. Composite meter factors are not acceptable for delivery point measurements.

The correction for the effect of temperature on liquids (CTL) factor must be determined in accordance with the API MPMS, Chapter 11.1. LPG must follow the applicable GPA Technical Publication TP-27 or an equivalent applicable procedure accepted by an appropriate industry technical standard association.

### 10.1.3 Pressure Correction Requirements

The correction for the effect of pressure on liquids (CPL) factor must be determined in accordance with API MPMS, Chapter 11, and is required only for LACT applications.

## 10.2 General Trucked Liquid Measurement, Accounting, and Reporting Requirements for Various Facility Types

### 10.2.1 Oil Batteries

For trucked oil/emulsion production into an oil battery, delivery point measurement is required for the total liquid volume. If there is a mixture of trucked-in production and prorated production within the same battery, the exception criteria in Section 5.5.2 must be met or ER site-specific approval must be obtained.

To meet heavy oil trucked production delivery point measurement requirements, the licensee must use an appropriate measurement method based on the fluid characteristics, such as viscosity, temperature, and sand content of the load. Generally, delivery point measurement is performed by using weigh scales or tank gauging (see Section 14.7.5 for tank gauging requirements) with sampling to determine the S&W and/or density. Meters are used only when there are minimal or no solids present in the oil/emulsion, similar to trucking into an oil battery that produces non-heavy oil.

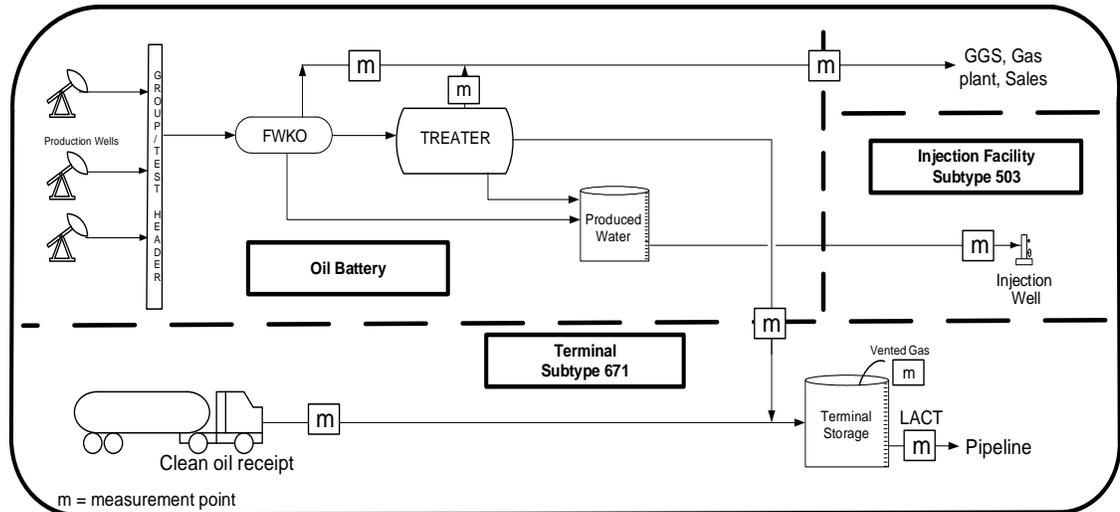
For condensate trucked into an oil battery, delivery point measurement is required for the total liquid volume. The requirements in Section 6.7 must be met.

For any oil battery, the trucked-out liquid is measured at the delivery point located at the receiving facility, and the oil volume determined at the receiving facility must be used as the delivering battery's reported oil disposition. In this scenario, delivery point measurement must be set up at the receipt point only. In the scenario where a facility delivers load oil to wells, delivery point measurement is required at the loading facility. If there are emergencies at the receipt point, the measurement at the originating facility may be used, but only as a temporary solution.

If clean oil from a battery is delivered into an oil pipeline via a LACT unit and that same battery also receives clean trucked oil, condensate, or diluent from other sources a Petrinex terminal (TM) reporting facility ID may be obtained and the clean trucked-in fluid can be reported as being received at the terminal instead of at the battery. Please be aware that reporting receipts at the terminal will trigger oil valuation and possible reduction of trucking charges. In this scenario, the battery oil must also be measured with delivery point measurement before commingling with other fluids at the terminal. The terminal will then deliver the fluid to the pipeline via the LACT unit.

In the scenario described above, if the licensee chooses to not create a terminal and install delivery point measurement for the battery oil being delivered to the terminal, measurement by difference will exist at the battery and the requirements in section 5.5 apply (see Figure 10.1).

Figure 10.1 Oil battery and terminal schematic

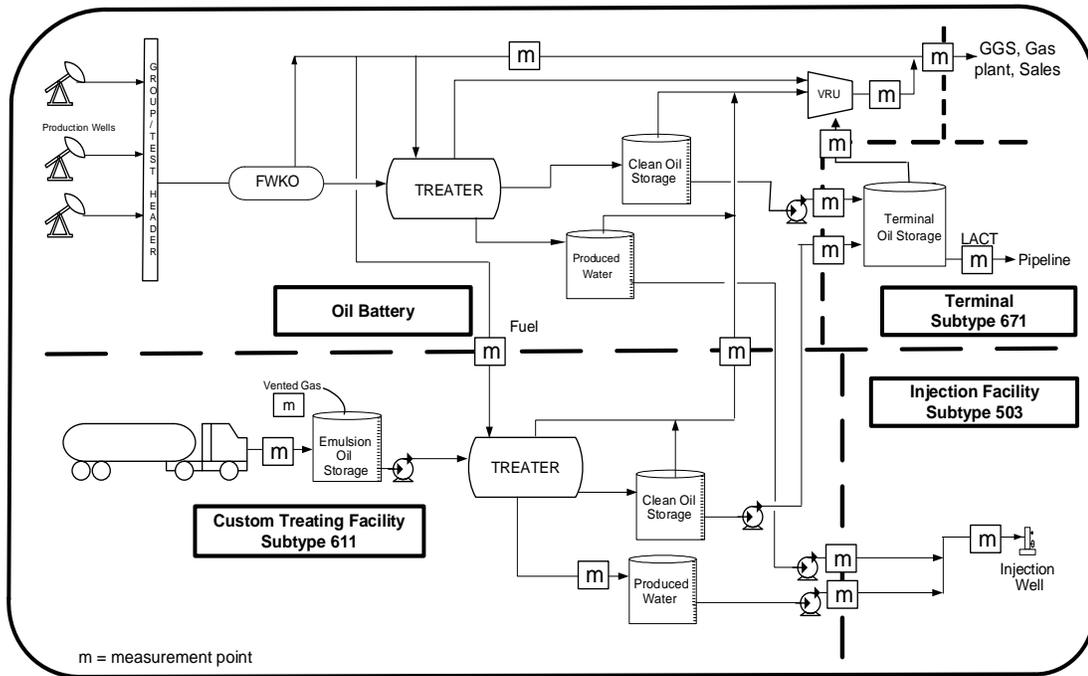


### 10.2.1.1 Custom Treating, Oil Battery, and Terminal Delineation

For reporting purposes, a terminal reporting facility ID may be created in Petrinex when there is more than one source of clean oil going through a LACT meter into an oil pipeline. If there is more than one Petrinex reporting facility on a single site, any oil, water, and gas (except for fuel see Section 4.1) crossing a facility boundary must be measured. If there is blending of hydrocarbon liquids of densities that differ by  $> 40.0 \text{ kg/m}^3$ , such as butane blending with the oil before the LACT, the lighter hydrocarbon used for blending must be received and stored at the terminal and the oil production must be measured before the blending point.

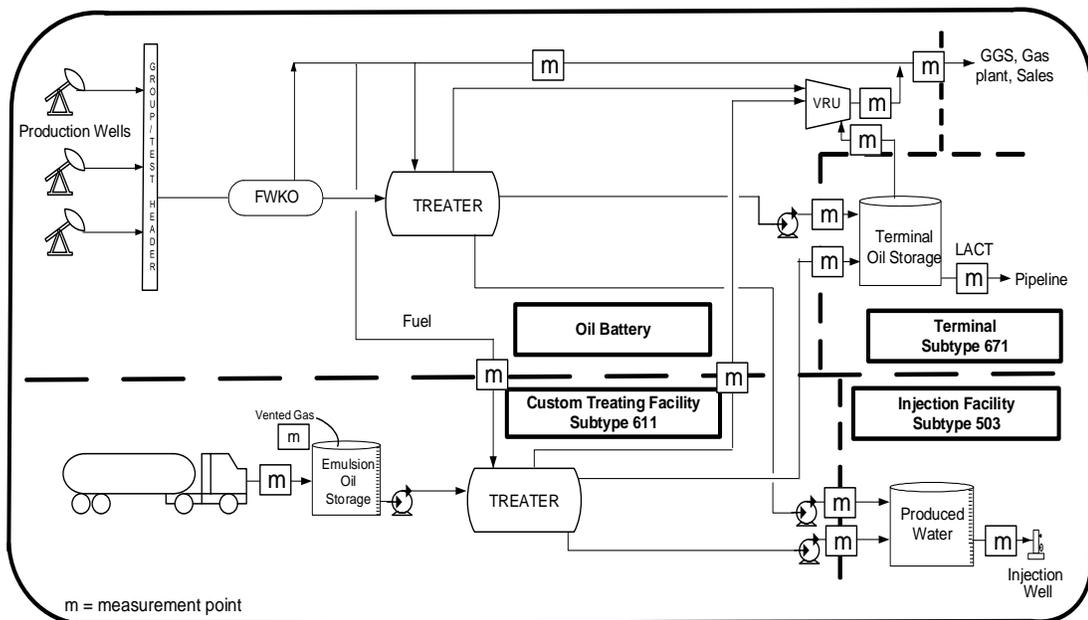
**Scenario 1:** Individual tanks for clean crude oil and produced water for both the crude oil proration battery and the custom treating facility.

Figure 10.2 Custom treating, oil battery, and terminal schematic – scenario 1



**Scenario 2:** Individual metering on treaters for water/oil and a shared tank for clean crude oil and produced water.

Figure 10.3 Custom treating, oil battery, and terminal schematic – scenario 2



The main difference between the two scenarios is that scenario 1 has individual tanks with metering off the tanks, whereas scenario 2 has shared tanks but metering off each treater.

For both scenarios, the transfer of fuel gas from the proration battery to the custom treating facility provides heat for the custom treater and pressure to help dump the treater to storage

tanks. There are also receipt meters for the gas coming back from the custom treater and terminal to the proration battery.

### 10.2.2 Custom Treating Facilities

The measurement requirements are the same as for trucking into a oil battery see Section 10.2.1 and the accounting and reporting procedures must adhere to the requirements in Appendix 4 of *Directive PNG032: Volumetric, Valuation and Infrastructure Reporting in Petrinex*.

### 10.2.3 Terminals

At the terminals that receive either pipelined and/or trucked clean oil, the receipt meter or weigh scale measurement must meet the delivery point measurement requirements.

For facility subtype 671, if the receipt measurement point at the terminal may be considered to be a custody transfer point. That is, there is no proration/allocation from the disposition volumes to the receipt (REC) volumes that are reported to Petrinex. Any measurement beyond this point is considered as downstream operations and not covered in this Directive.

However, if the terminal disposition measurement point is considered the custody transfer point then the licensee must allocate the disposition volume back to the receipts and report the allocation factors in Petrinex then the facility subtype is 673.

### 10.2.4 Gas Plants, Gas Batteries, and Gas Gathering Systems

For gas systems receiving trucked liquid, the measurement requirements are the same as for trucking liquid into a non-heavy oil battery.

### 10.2.5 Water Injection/Disposal Facilities

For water trucked into an injection or disposal facility, delivery point measurement accuracy is not required. See Sections 1.6.4.3 and 1.6.4.4 for facility accuracy requirements.

### 10.2.6 Waste Plant Facilities

A waste plant facility processes volumes of waste generated in the upstream petroleum industry. However, many ER-approved waste plants have an integrated custom treating facility for processing oil/water emulsions extracted from the solids during waste processing. In addition, oil/water emulsions from other batteries may be trucked in and measured independently from the waste oil/water emulsions, and both streams are processed through the same custom treating facility. The total waste oil/emulsion disposition from the waste plant to the custom treater (CT) must have volume and S&W determinations in order to properly allocate the CT clean oil and water disposition volumes back to the trucked in receipts. Therefore, delivery point measurement is required at the receipt point of oil/water emulsions from other batteries and at the total waste oil/emulsion delivery point from the waste plant to the CT.

There may also be injection/disposal facilities that receive other liquids, such as waste streams going into subsurface caverns for disposal. Waste liquids for disposal require the same measurement accuracy as the disposal of produced water. See *Directive PNG032: Volumetric, Valuation and Infrastructure Reporting in Petrinex* for the requirements for waste stream measurement, accounting, and reporting.

### 10.2.6.1 Integrated Waste Processing Facilities

Integrated waste processing facilities include the following facility sub-types:

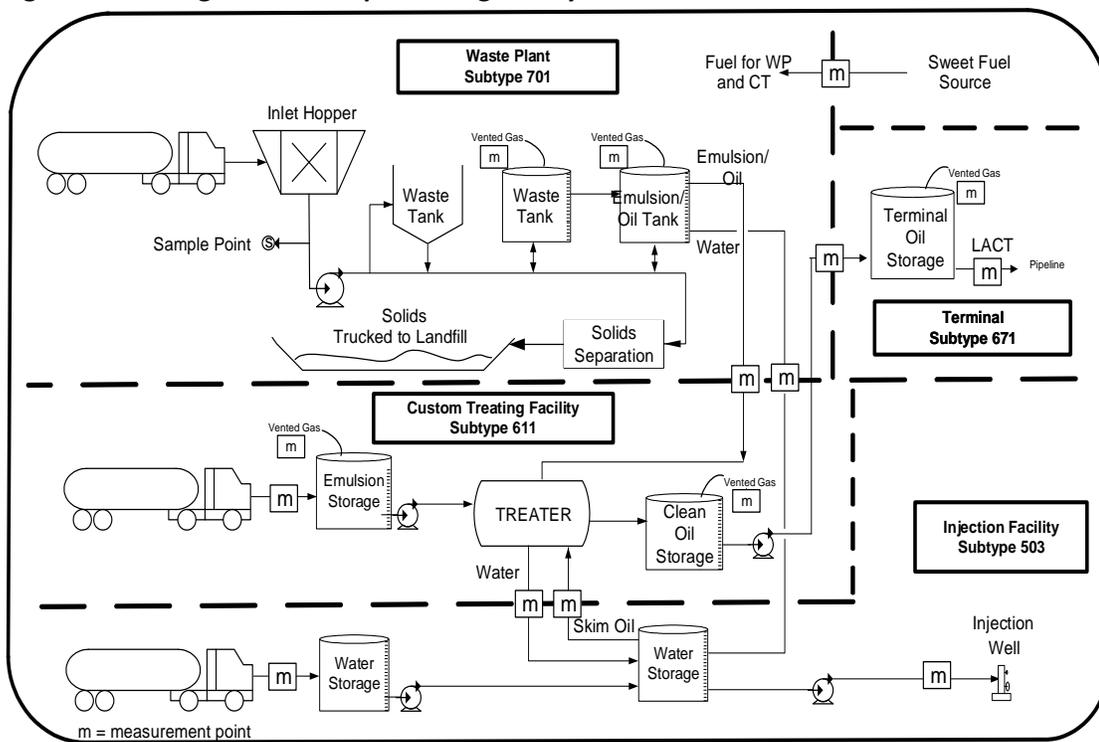
1. Waste Plant ( facility subtype 701)
2. Custom Treater (facility subtype 611)
3. Water Disposal Injection Facility (IF): Disposal (facility subtype 503) and Disposal (Approved as Part of Waste Plant) facility subtype 507.
4. Terminal (TM)

Integrated waste processing facilities are ones with various distinct processing and reporting entities. They are referred to as oilfield waste management facilities (OWMFs), see Figure 10.4.

Any fluids transferred between the different reporting facilities within the integrated site must be measured and reported.

The fuel gas receipt and usage for the site can be reported at the WP. Fuel gas measurement and reporting is not required at the CT or other reporting facilities on the site. See *Directive PNG032: Volumetric, Valuation and Infrastructure Reporting in Petrinex* for more details.

**Figure 10.4 Integrated waste processing facility delineation**



### 10.2.6.2 Waste Caverns

Waste Cavern processing facilities include the following facility sub-types:

1. Surface Waste Plant facility (facility subtypes 701)
2. Cavern Waste Plant facility (facility subtype 702)

Any fluids transferred between the surface waste plant facility and the cavern waste plant facility must be measured and reported. Waste volumes injected and recovered from the

cavern must be metered at each well. The wellhead injection volumes and recovered volumes are not required to be reported at the well level but must be reported as a receipt and disposition at the facility level at the cavern waste plant reporting facility ID. Inventories in the cavern must be reported.

The fuel gas receipt and usage for the site can be reported at the surface waste plant facility. See *Directive PNG032: Volumetric, Valuation and Infrastructure Reporting in Petrinex* for more reporting details.

### 10.3 Design and Installation of Measurement Systems

Delivery point measurement is required for trucked fluid receipts. The gross volume must be measured through a system consisting of inlet tank gauging, inlet meter, or weigh scale. If measurement is by inlet tank gauging, gauge boards must not be used.

Truck ticket estimates, such as volume estimates determined using the truck tank load indicator and used for bill of lading/transportation of dangerous goods purposes, are not acceptable for the purpose of well or facility volume measurement. Truck ticket estimates must not be used for determining volumes unless the requirements in Section 10.3.4 are met.

See Section 14 for liquid measurement design and installation requirements.

#### 10.3.1 Meters

Turbine meters are typically not suitable for viscous fluids and therefore are not recommended for measurement when unloading crude oil.

When metering devices for the purpose of measuring truck delivery/receipt volumes are installed, the following must also be installed:

1. Sample point
2. Air eliminator

For some types of meters and applications, a strainer and a back pressure control system are required. Refer to Figure 14.1 for more information.

Additional requirements for clean oil and pipeline terminals:

1. For mechanical automatic temperature compensators without gravity selection (ATC) or with gravity selection (ATG):
  - a. For new construction or new installations at existing facilities, mechanical ATC and ATG must not be used. For facilities built before April 1, 2016, all existing ATC and ATG are grandfathered at their existing installations and must not be relocated and reused for other installations.
  - b. The difference between actual density and compensation density must be less than 40.0 kg/m<sup>3</sup>.
  - c. Product temperature must be between -10.0°C and +40.0°C excluding LPGs.
  - d. The compensation density, mechanically set density or user-entered density for electronic flow computers, must be a volume weighted average of the expected receipt volumes. When product temperatures exceed +40.0°C, it

may be necessary to reduce the allowable density difference to maintain a  $\pm 0.5$  per cent uncertainty.

- e. Temperature compensation devices must be designed for the actual range of operating temperatures observed. If product temperatures exceed  $+40.0^{\circ}\text{C}$ , it may be necessary to reduce the allowable density difference to maintain a  $\pm 0.5$  per cent uncertainty.

### 10.3.2 Weigh Scales

Weigh scales used for the purpose of delivery point measurement must be verified in accordance with Section 2.12. For sampling points and methods, see Section 8.5. Systems employing weigh scales must also provide for determination of density of oil and water in accordance with one of the following:

1. API MPMS, Chapter 9: Density Determination Using Hydrometer
2. precision laboratory method – ASTM method or
3. on-line densitometer

Also see Section 14.6 for more information regarding the methods that may be used to determine density.

To maintain a measurement uncertainty of  $\pm 0.5$  per cent or less, the net weight of the payload must be  $\geq 40.0$  per cent of the gross vehicle weight and  $\geq 6500.0$  kg. An exemption from this requirement is granted only during seasonal road ban periods when reduced truck loads are mandated by weight restrictions.

### 10.3.3 Exceptions for Truck Measurement Systems

#### 10.3.3.1 Truck-Mounted Level Gauges and Truck-Mounted Meters

Truck gauge level indicators and truck-mounted meters are considered to have met the requirement for low-accuracy measurement with an overall measurement uncertainty of  $\pm 1.0$  per cent or less if the following criteria are met:

1. The battery receives not more than  $100.0 \text{ m}^3$  of trucked-in oil per day.
2. The percentage of trucked-in oil to the battery is  $\leq 10.0$  per cent of the monthly battery oil production volume.
3. The gauges or meters are verified/proved annually and if they are not within  $\pm 1.0$  per cent accuracy they are repaired and recalibrated/reproved.
4. The product temperature is determined to within  $\pm 1.0^{\circ}\text{C}$ , see Section 10.3.2, item 2.
5. The truck gauge levels or meters are initially set by calibrating to a master meter or prover with a demonstrated uncertainty  $\leq 0.2$  per cent.

Additional criteria for truck-mounted level gauges:

1. The stated depth of liquid is within 12.7 mm of a known gauge level marker if used.
2. The depth of liquid is determined while the tank trailer is level to within 150.0 mm over its length.
3. The minimum load on the trailer  $> 65.0$  per cent of full load.

These units can be used for trucked-in delivery point measurement to proration oil batteries.

### 10.3.3.2 Truck Tickets and Lease Tank Gauging

Truck ticket volumes uncorrected for temperature are not acceptable for delivery point measurement of trucked liquid. If the fluid transfers are between unitized facilities or facilities with no equity or royalty concerns, then the temperature correction estimates may be used. The truck ticket must be based on a low-accuracy measurement requirement with an overall uncertainty of  $\pm 1.0$  per cent or less of trucked liquid, such as lease tank gauging at the battery sending the liquid production or truck-mounted meter, for determining inlet volumes at a proration battery if certain situations exist. The S&W per cent and corrected opening and closing readings must be on the ticket or available on a summary sheet for ER audit purposes. An individual truck load must be recorded on its own ticket.

ER may accept low-accuracy measurement with an overall uncertainty of  $\pm 1.0$  per cent or less for trucked liquid production at a proration battery if:

1. Trucked production is temporary, pending battery consolidation within one year or less.
2. Individual well oil volumes being trucked are  $< 2.0 \text{ m}^3/\text{day}$ , see Section 10.3.5.
3. The crude oil volume receipt net of water is  $\leq 10.0$  per cent of the total receiving battery oil production.
4. Truck-mounted meters used for low-accuracy measurement with an overall uncertainty of  $\pm 1.0$  per cent or less are proved in accordance with the requirements in Section 2.

### 10.3.4 Load Fluids

Load fluids, at a minimum, must be measured using devices that meet the requirement for low-accuracy measurement with an overall uncertainty of  $\pm 1.0$  per cent either at the source loading location or at the delivery point.

Reporting of load fluid on Petrinex is limited to oil-based and/or water-based fluid(s) injected during pre-production well stimulation or post-production activities. Only the load fluid product codes OIL, COND, or WATER can be reported. Well drilling fluids must not be reported on Petrinex as load fluids.

Load fluids are reported at the well level except when in an SW Saskatchewan shallow gas battery, since there is no requirement to measure and report water production at this type of well. The load fluid reporting then can be done at the facility level as a receipt (REC) and disposition (DISP). See the *Directive PNG032: Volumetric, Valuation and Infrastructure Reporting in Petrinex* for more reporting procedures.

### 10.3.5 Split Loads

A split load is defined as existing when a truck takes on partial loads from more than one well or battery in a single trip or when load oil is delivered to more than one receipt point or well.

**Requirements:** If the densities of the split load components are different by more than  $40.0 \text{ kg/m}^3$ , blending tables are required to calculate shrinkage. The shrinkage volume is to be prorated back to each battery on a volumetric basis.

**Measurement:** Volume from each well or facility must be measured at the time of loading onto the truck (or off-loading from the truck for load oil) by one of the methods:

1. Gauging the battery lease tank.
2. Gauging the truck tank (not allowed for density difference over 40.0 kg/m<sup>3</sup> for any oils or emulsions).
3. Truck-mounted meter/gauge that meets low-accuracy measurement and is proved at least annually.

Calibrated gauge tables are required for methods 1 and 2.

**Sampling:** Fluid from each single-well oil battery must be sampled to determine the S&W and the oil/water volumes. The truck driver is to collect the samples by taking at least three well-spaced grab samples during the loading period, see Section 14.6 and 8.5.

For load oil, the S&W must be determined at the loading source.

**Records:** The truck tickets must show the individual load volumes, as well as the total volume at delivery (receipt) point, supported by opening and closing gauge or meter readings.

**Accounting:** For battery emulsions, the total load is to be measured and sampled at the receiving location and prorated to each of the wells based on the measured loading volumes and S&W from each of the wells.

For load oil, the initial volume must be measured at the loading source and prorated to each delivery point based on the measured volume delivered to each well.

**Allowed:**

1. Single-well oil battery delivering to other facilities.
2. Gas wells with condensate-water tanks and production less than 2.0 m<sup>3</sup> of total liquids per day.
3. Blending of heavy oil and condensate.
4. Load oil for well servicing only, specifically load oil from a single source only.

**Not Allowed**

1. Multi-well batteries delivering to other facilities other than load oil.
2. Gas wells with production that is > 2.0 m<sup>3</sup> of total liquids per day.

## 10.4 Sampling and Analysis

For trucked-in hydrocarbon and emulsion receipts, a truck thief sampler or a proportional sampler may be used to obtain a sample from the truck tank, see Section 14.6 and 8.5. In some scenarios spot (grab) samples may be used to obtain the sample from the off-load/load line. Automatic sampling methods are preferred. However, manual or tank sampling systems may also be allowed, as described in Sections 10.4.2.

The frequency of sampling or readings must be sufficient to ensure that a representative sample of the entire truck volume is obtained. Consideration must be given to both conditioning the flow stream and locating a probe or sampler. Flow conditioning to ensure turbulent mixing can be achieved through velocity control, piping configurations, or introduction of a mixing element upstream of the sample point. A sample probe is required for truck delivery point measurement sampling unless there is an in-line product analyzer or the sampling is incorporated as part of the measurement system. A mid-pipe probe location must be used for accurate sampling, also see Sections 8.2.1 and 8.2.2.

The licensee must choose the sampling methodology based on emulsion characteristics, stratification, and S&W consistency of each load to obtain a representative sample. API MPMS, Chapter 8.1, Section 8, provides further information on manual sampling procedures.

### 10.4.1 Automatic Sampling

Automatic sampling is typically conducted through the use of proportional samplers. If automatic sampling procedures are used, a manual procedure must also be in place for use when the automatic system is out of service or for intermittent verification of the automatic system reading. For more information, API MPMS, Chapter 8.2, Sections 7 to 15, provide further details on flow conditioning, probe location, and sampling frequency.

Other requirements for automatic sampling:

1. Containers made of suitable material for handling and storage of the sample must be used. Container lids must be vapour tight.
2. All sample containers must be cleaned and dried prior to collection of the next sample.
3. Sample containers must allow adequate room for expansion and content mixing, taking into consideration the temperature of the liquid at the time of filling.
4. The sample containers must be housed in a secured enclosure to prevent any tampering with the sample.
5. Sample lines must be as short as practical and sloped downward to reduce the possibility of plugging up the sample line.

### 10.4.2 Manual Spot (Grab) Sampling

Manual spot (grab) sampling may be acceptable in situations involving a tight emulsion  $\leq 0.5$  per cent S&W in the truck by taking three well-spaced grab samples during the unloading period, see Section 14.6 and 8.5. A single grab sample is not acceptable when there is stratification of S&W within the truck.

The use of manual sampling techniques, such as truck tank thieving, (full height or intermittent) may also be acceptable. However, in the presence of stratification, one unit of height at the

bottom of the truck tank represents a significantly lesser volume than the same unit of height at the midpoint of the truck tank because of the shape of the tank. The resulting S&W from a full-height core sample therefore may not be representative of the entire load. In such cases multiple grab samples must be used.

Lease tank manual sampling is subject to similar stratification limitations excluding the non-uniformity of the tank. These concerns can be reduced by locating any water-emulsion interface and obtaining bottom, middle, and top samples of the emulsion to determine the average water cut of the emulsion. However, lease tank manual sampling requires dedicated tankage for each load received or delivered to avoid mixing of product between deliveries.

Visual estimates or estimates based on changing off-load pump speeds must not be used for free water volume determination.

### 10.4.3 S&W Determination

The licensee must select the most appropriate method for determining the S&W, see Section 14.8 and Appendix 3.

### 10.4.4 Density Determination

Truck load sample density determination at 15.0°C must be conducted at least annually or more frequently if there are changes in the reservoir conditions. Density of the load may be determined by one of the following methods:

1. Truck load samples may be collected from the receiving point and sent to an independent laboratory for analysis to determine density of the liquid hydrocarbon phase and the liquid water phase (if required). The density found in this analysis must be applied to all hydrocarbon liquids coming from the specific facility; or
2. Truck load samples or samples from automatic samplers may be tested for density as outlined in Section 14.6 and 8.5.

In situations where the truck volumes have an S&W that is > 1.0 per cent, density determination at 15.0°C of an emulsion sample is difficult, as there are two different thermal corrections to be applied, one for the water and one for the oil.

There are two options available:

1. The first is to determine the sample density using a precision densitometer that has its measuring cell at 15.0°C. No further corrections are required.
2. The second is to separately predetermine the density at 15.0°C of the water and the oil. When using this option, the emulsion density is calculated by applying the S&W cut to the density of each component. The calculation is

$$\rho_{emulsion} = (\rho_{oil} \times (100 - \%S \& W)) + (\rho_{water} \times \%S \& W)$$

Where:

$\rho_{emulsion}$  is the calculated density of the emulsion at 15.0°C

$\rho_{oil}$  is the density of the oil portion at 15.0°C

$\rho_{water}$  is the density of the water portion at 15.0°C

## **10.5 Volume Determination**

### **10.5.1 Tank Gauging**

Tank gauging procedures are detailed in Section 14.7. The opening and closing levels measured are then converted to volume through the use of gauge tables, supplied by the tank manufacturer, which have been calculated using measurements of the tank. The difference between the closing and opening volumes is the measured volume. If the tank is used for delivery point measurement, the temperature and density of the tank contents must be taken in order to correct the indicated volume to standard conditions before determining the volume difference.

### **10.5.2 Weigh Scales**

The procedure for determining the volume of liquid on a truck using a scale is to weigh the truck before and after loading or unloading and determine the difference to obtain the net weight. The entire load must be weighed at a time. Split weighing, whereby the truck is weighed after unloading a portion of its load to obtain the weight of the unloaded portion, is not permitted unless it is used in cold heavy oil measurement.

To determine the density of the load, an on-line densitometer maybe used or a representative sample must be obtained and the density and temperature measured with a hydrometer and thermometer respectively. The observed density must be corrected to 15.0°C.

The net weight determined during the weighing process divided by the sample density at 15.0°C results in the net volume of the load prior to deductions for S&W.

### **10.5.3 Meters**

Metered volumes must be determined in accordance with Section 14.

## 11 Acid Gas and Sulphur Measurement

This section presents the base requirements and exceptions for acid gas and sulphur measurements at processing plants and injection facilities in the upstream oil and gas industry that are used in determining volumes for reporting to ER.

In a gas processing plant where sour gas is processed, most of the acidic portion of the gas must be removed from the gas stream (sweetening) in order to produce a saleable pipeline-quality gas product. In the process of removing the acidic portion of the sour gas, acid gas, which consists mainly of H<sub>2</sub>S and CO<sub>2</sub>, is generated and must be disposed of in an environmentally and economically acceptable way, such as by elemental sulphur production, acid gas injection, or acid gas flaring.

### 11.1 General Requirements

Sour gas plant inlet gas and acid gas streams must be metered and reported.

The acid gas from the sweetening process is generally saturated with water vapour. This water vapour portion must be subtracted from the saturated acid gas to obtain the dry volume without water vapour. For more information refer to Section 11.2.

### 11.2 Acid Gas Measurement

The quantity of acid gas going to sulphur plants, to compression and injection, or to flaring is generally measured at a low pressure of 50.0 to 110.0 kPag; therefore, the orifice meter, or any other meter, must be appropriately sized and maintained to manufacturer's recommended specifications to achieve accurate measurement.

Acid gas is saturated with water vapour, which represents a significant portion of the total gas measured. The amount of water vapour varies significantly with the temperature of the reflux drum. Therefore, the acid gas meter must have continuous temperature correction (see Section 4.3) to calculate the correct acid gas volume as outlined in this section. The gas density must also include the water content, and the meter coefficient must include a factor to exclude the water vapour content of the gas in the final volume computation for reporting purposes. The accuracy of the gas relative density factor and water content determination must be annually verified to ensure that acid gas measurement uncertainty is within tolerance.

#### 11.2.1 Determining Acid Gas on a Dry Basis

For ideal gases, the total vapour pressure of a system containing several components is the sum of the vapour pressures of the individual components at the temperature of the system.

A component's vapour pressure percentage of the total pressure of a system is equal to the volume percentage of that component in the system. The reflux drum is the vessel in which the acid gas separates from the sweetening solution. The amount of water vapour in the acid gas leaving the reflux drum is a function of the temperature and the absolute pressure in the reflux drum.

### 11.2.1.1 Calculating Acid Gas Flow Rate

The calculation method for the acid gas flow rate is as follows:<sup>3</sup>

- Step 1:** Determine the percentage of water vapour in the acid gas on the basis of the ratio of vapour pressure of water to total pressure in the reflux drum at the reflux drum temperature.
- Step 2:** Convert the acid gas composition from dry basis to wet basis at reflux drum pressure and temperature, and determine the acid gas relative density and compressibility factor on a wet basis at meter pressure and temperature.
- Step 3:** Calculate the acid gas and water vapour flow rate corrected from actual flowing pressure and temperature to standard conditions of 101.325 kPa(a) and 15.0°C.
- Step 4:** The volume calculated in step three contains water vapour in the percentage determined in step one and must be converted to dry basis volume for reporting purposes. An acid gas flow correction factor (CF) has to be applied to correct the acid gas flow from a wet to a dry basis.

$$CF = (100.00 - \% \text{ H}_2\text{O in acid gas}) \div 100$$

$$\text{Dry acid gas flow rate} = CF \times \text{flow rate calculated in Step 3}$$

The H<sub>2</sub>S content of the acid gas is the dry basis acid gas flow times the dry basis percentage of H<sub>2</sub>S in the acid gas divided by 100.

### 11.2.1.2 Calculating Vapour Pressure of Water

The formula for determining the vapour pressure of water<sup>4</sup> is

$$\log P = A - B \div (C + T)$$

where: P = water vapour pressure in mm of mercury

$$A = 8.10765$$

$$B = 1750.280$$

$$C = 235$$

T = temperature of acid gas in reflux drum (°C)

The resulting direct formula for determining the vapour pressure of water in kPa(a) is

$$P_{\text{H}_2\text{O}} = 0.13332 \times 10^{(8.10765 - 1750.280 / (235 + T))}$$

Where: P<sub>H<sub>2</sub>O</sub> = water vapour pressure in kPa(a) at T°C

$$\% \text{ H}_2\text{O in the acid gas} = (100 \% * P_{\text{H}_2\text{O}}) \div (P_{\text{RD}} + P_{\text{atm}})$$

Where: P<sub>RD</sub> = reflux drum pressure, kPag

P<sub>atm</sub> = atmospheric pressure, kPa(a)

<sup>3</sup> Wichert, E., "Water content affects low pressure, acid-gas metering," *Oil & Gas Journal*, January 2, 2006, pp. 44–46.

<sup>4</sup> The vapour pressure of water at a certain temperature can also be obtained from the GPSA *Engineering Data Book*, SI Units version, 12th edition, 2004 or subsequent versions, Figures 24–36.

### 11.2.1.3 Converting Acid Gas Calculation from Dry to Wet Basis

An example acid gas conversion calculation from dry to wet basis with the meter installed upstream of the back-pressure regulator of the reflux drum is provided as follows:

#### A. Reflux drum data

Reflux drum temperature = 40.0°C

Reflux drum pressure = 70.0 kPag

Atmospheric pressure = 95.0 kPa(a)

If the meter is installed upstream of the back-pressure regulator of the reflux drum, the upstream pressure and the temperature of the meter run may be used as the reflux drum pressure and temperature.

#### B. Acid gas components on a dry basis from acid gas analysis:

H<sub>2</sub>S = 65.0%    CO<sub>2</sub> = 33.5%    C<sub>1</sub> = 1.2%    C<sub>2</sub> = 0.3%

#### C. Calculate the percentage of components, including water vapour, on a wet basis:

Percentage of water vapour =  $(100\% \times \text{Vapour pressure of water at } 40.0^\circ\text{C}) \div (\text{Reflux drum gauge pressure} + \text{atmospheric pressure})$

$$P_{\text{H}_2\text{O}} = 0.13332 \times 10^{(8.10765 - 1750.280 / (235 + 40))}$$

$$= 7.377 \text{ kPa(a)}$$

(Note that the vapour pressure of water at 40.0°C is 7.384 kPa(a), from the Saturated Steam Table in the Thermodynamics section of the *GPSA SI Engineering Data Book*, Figures 24-36.)

$$\text{Percentage of water vapour} = 7.377 \div (70.0 + 95.0) \times 100\% = 4.47\%$$

Enter into column 2 (see [Table 11.1](#)) and normalize the gas composition. To determine the component concentration in column 2 (except for H<sub>2</sub>O) multiply the dry basis concentration by  $(100 - 4.47)/100$ .

**Table 11.1 Calculation of relative density (RD) on wet basis**

	<u>Column 1</u>	<u>Column 2</u>	<u>Column 3</u>	<u>Column 4</u>	<u>Column 5</u>
<b>Comp.</b>	<b>Dry basis (%)</b>	<b>Wet basis (%)</b>	<b>Molar mass (kg/kmol)*</b>	<b>(Col. 1 * Col. 3) /100</b>	<b>(Col. 2 * Col. 3) /100</b>
H <sub>2</sub> S	65.00	62.09	34.082	22.153	21.162
CO <sub>2</sub>	33.50	32.00	44.010	14.743	14.083
C <sub>1</sub>	1.20	1.15	16.042	0.193	0.184
C <sub>2</sub>	0.30	0.29	30.069	0.090	0.087
H <sub>2</sub> O	0.00	4.47	18.0153	0.000	0.805
<b>Total</b>	<b>100.00</b>	<b>100.00</b>		<b>37.179</b>	<b>36.321</b>

\* Molar mass of air = 28.9586 kg/kmol (*GPSA Engineering Data Book*, 2004 or later editions, Figure 23-2, or GPA-2145).

From column 4, ideal gas RD, dry basis =  $37.179/28.9586 = 1.284$ .

From column 5, RD wet basis =  $36.321 \div 28.9586 = 1.254$  (this RD is to be used in the flow calculation for acid gas volumes).

#### 11.2.1.4 Difference between the Acid Gas Volume on a Wet Basis and on a Dry Basis

An example calculation is presented to show the difference in the results of the acid gas flow rate and the sulphur content of acid gas using dry versus wet basis metering. The example data for the meter run and assumed conditions are as follows:

1. Orifice meter diameter: 154.051 mm
2. Orifice plate diameter: 76.200 mm
3. Meter upstream pressure: 70.0 kPag
4. Differential pressure: 10.0 kPa
5. Meter temperature: 40.0°C
6. Atmospheric pressure: 95.0 kPa
7. Acid gas composition: as per [Table 11.2](#)

Results with AGA #3 1992 or later method:

1. Flow rate, dry basis without accounting for moisture content =  $33.126 \text{ } 10^3 \text{ m}^3/\text{d}$
2. Sulphur content =  $(33.126 \times 65) \div (100 \times 1.35592) = 29.20 \text{ tonne/day}$
3. Flow rate, wet basis =  $33.499 \text{ } 10^3 \text{ m}^3/\text{d}$ , containing 4.47 percent H<sub>2</sub>O
4. Flow rate, wet basis converted to dry basis =  $(33.499 \times (100 - 4.47)) \div 100$   
=  $32.002 \text{ } 10^3 \text{ m}^3/\text{d}$  dry acid gas equivalent

This volume,  $32.0 \text{ } 10^3 \text{ m}^3$ , is to be reported as Acid Gas on the Petrinex monthly volumetric submission.

An example for percentage difference in acid gas volume between dry and wet basis:

1. Percentage difference in flow rate =  $(33.126 - 32.002) \times 100 \% \div 32.002 = 3.51\%$
2. Sulphur content =  $(32.002 \times 65) \div (100 \times 1.35592) = 28.21 \text{ tonne/day}$
3. Difference in calculated sulphur balance between dry and wet basis metering  
=  $29.20 - 28.21 = 0.99 \text{ tonne/day}$
4. Percentage difference =  $(0.99 \times 100) \div 28.21 = 3.51\%$

Thus, if the moisture content in the metering of the acid gas in this example were ignored, volume determined on a dry basis instead of wet basis, the reported acid gas flow and sulphur content in the acid gas leaving the reflux drum would be 3.51 per cent higher than the correct value.

This method of estimating the water vapour content is valid when the gas is in contact with water in a low-pressure vessel, such as in the reflux drum. The method does not apply to low-pressure gas, such as in a flare line, when the flared gas originates from a high-pressure vessel.

The [Table 11.2](#) summarizes the previous example and also provides the results that are obtained by the 1985 AGA # 3 Report method, using Wichert-Aziz (W-A) compressibility factors.

**Table 11.2 Summary of acid gas volume determination wet vs. dry basis example**

Item	AGA #3, post-1992		AGA #3, 1985, W-A Z factors*	
	Dry basis	Wet basis	Dry basis	Wet basis
Z factor at St'd P and T	0.992 848	0.991 999	0.993 037	0.992 788
Z factor at Meter P, T	0.991 002	0.990 030	0.991 007	0.990 674
Flow rate, 10 <sup>3</sup> m <sup>3</sup> /d	33.126	33.499	33.096	33.481
Corrected to dry gas	-	32.002	-	31.984
% difference	-	3.51	-	3.48
Sulphur flow, tonnes/day	29.20	28.21	29.17	28.19
% difference, tonnes/day	-	3.51	-	3.48

\*Z factors by Wichert-Aziz method, including water content in wet gas.

### 11.2.1.5 Calculation Method of Water Content if Meter Located Downstream of Back-Pressure Valve of Reflux Drum

The water content in the acid gas is a function of the pressure and temperature of the reflux drum. If the acid gas meter is located downstream of the back-pressure regulator of the reflux drum, both the pressure and the temperature of the meter will be somewhat lower than the pressure and temperature of the reflux drum. Under these conditions, it is still necessary to determine the water vapour content of the acid gas stream at the reflux drum pressure and temperature, as shown in the previous example, to correctly calculate the acid gas flow rate.

The reflux drum pressure must be recorded for the correct calculation of the water vapour content of the acid gas. The reflux drum temperature must be used to estimate the water content. However, since the flow data from the meter includes the temperature at the meter run, the reflux drum temperature can be estimated on the basis of the meter temperature, as follows:

$$T_{RD} = ((T_m + 2.28 - (2.28 * P_2)) \div (P_{RD} + P_{atm}))$$

Where:  $T_{RD}$  = reflux drum temperature, °C

$P_2$  = the downstream meter tap pressure, kPa(a)

$P_{RD}$  = reflux drum pressure, kPag

$P_{atm}$  = atmospheric pressure, kPa(a)

$T_m$  = temperature downstream of the orifice plate,

Having estimated the temperature at the reflux drum from the temperature downstream of the orifice plate, the vapour pressure of the water can be determined using the calculations shown in Section 11.2.1.2. The percentage of water vapour in the acid gas can then be determined using the reflux drum pressure, and the same procedure as outlined in the example shown in Section 11.2.1.3 can be used to calculate the acid gas flow rate.

### 11.2.1.6 Effect of Compression and Cooling of Acid Gas

In the situation of acid gas compression and injection, the acid gas flow rate may in some instances be metered after one or more stages of compression and cooling, refer to Section 11.4.5.3. This will remove a sufficient amount of water so that the remaining water vapour in the compressed and cooled acid gas will have little effect on the acid gas metering. In such a situation, it is not necessary to include the effect of water vapour in the metering of the acid gas.

## 11.3 Sulphur Measurement and Pit Volume Determination

### 11.3.1 Sulphur Pit Volume/Tonnage Determination

When pit gauging is used to determine a liquid sulphur volume, the gauging procedures must be conducted in accordance with the following:

1. The licensee must ensure that the gauge/strapping table used to convert the gauge level to a liquid volume is specific for the pit being gauged.
2. Pit gauging must be used for inventory determination only and must not be used for delivery point measurement.
3. All dip sticks and electronic level devices must have a minimum resolution of six mm.
4. It is acceptable to have one reading per determination.
5. The sulphur density at pit temperature is obtained from [Figure 11.1](#).<sup>5</sup>

The general formula for determining the produced sulphur tonnage is as follows:

$$\text{Sulphur tonnage} = \text{Gauge reading} \times \text{CF} \times \text{Sulphur density}$$

where CF = Pit gauge/strapping table conversion factor

### 11.3.2 Sulphur Measurement

For sulphur sales/delivery point measurement using meters, see Section 1.6.3.7 and 14.2 . These meters must be kept at a temperature so that the molten sulphur will not solidify when there is no flow.

For sulphur sales/delivery point measurement using a weigh scale, see Sections 2.12 and 10.3.2.

For daily sulphur production measurement using pit level gauging, two pits are required, one for production and the other for withdrawal using level measurement. The daily sulphur production tonnage must be adjusted by the total monthly disposition at the end of the month.

#### 11.3.2.1 Exception for Sulphur Tonnage Entering a Gas Plant

For daily sulphur production volume determination, if there is only one pit in place in an existing plant and sulphur is being withdrawn without measurement, the licensee may use the measured acid gas volume on a dry basis, provided that there is a continuous acid gas sampling device, such as a gas chromatograph, to calculate the sulphur tonnage entering the sulphur plant. The daily sulphur production can then be calculated using the following formula:

$$\text{Estimated daily sulphur production (t)} = \text{Daily acid gas inlet (t)} - \text{Daily incineration (t)} - \text{Daily flared (t)} - \text{Others if applicable (t)}$$

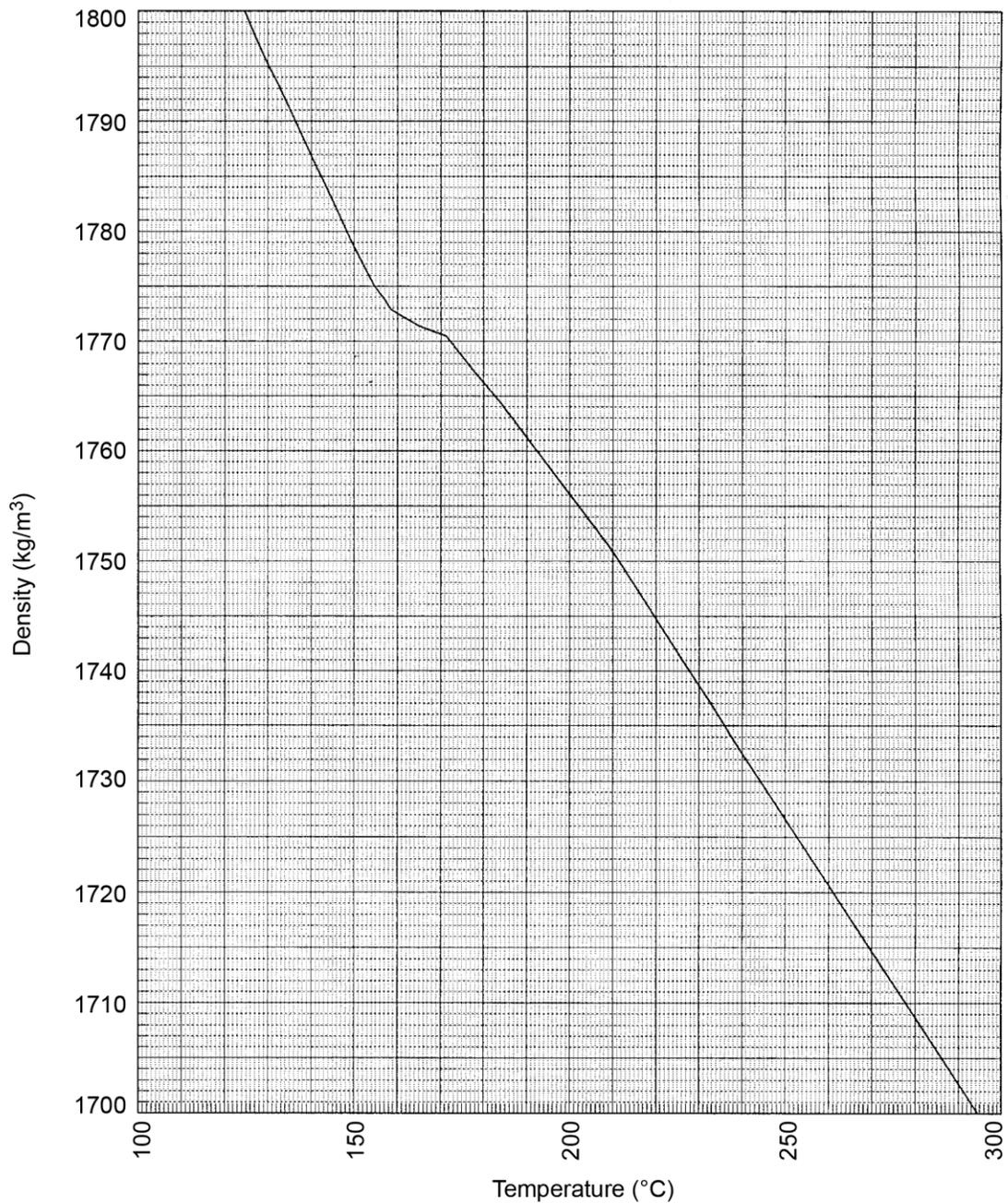
The estimated daily sulphur production tonnage must be adjusted by the total monthly disposition at the end of the month by calculating a proration factor and applying that to all estimated daily production tonnage:

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<sup>5</sup> Tullen, W. N., *The Sulphur Data Book*, New York: McGraw-Hill Book Company, Inc., 1954, p. 17.

Sulphur proration factor ( $S_{pf}$ ) = Total monthly sulphur disposition tonnage (including inventory changes)  $\div$  Total estimated daily sulphur production tonnage

Actual daily sulphur production (t) = Estimated daily sulphur production (t)  $\times$   $S_{pf}$

**Figure 11.1 Liquid sulphur density vs. temperature**

#### 11.4 Overview of Plant Inlet and Outlet Points for H<sub>2</sub>S

Figure 11.2 illustrates the paths by which H<sub>2</sub>S enters the sour gas plant and by which method it can exit from the plant.

Since the flow volumes of the three types of fluid streams out of the inlet separator are required to be measured, it becomes a simple task to account for the amount of H<sub>2</sub>S entering the plant by determining the H<sub>2</sub>S concentration in each stream.

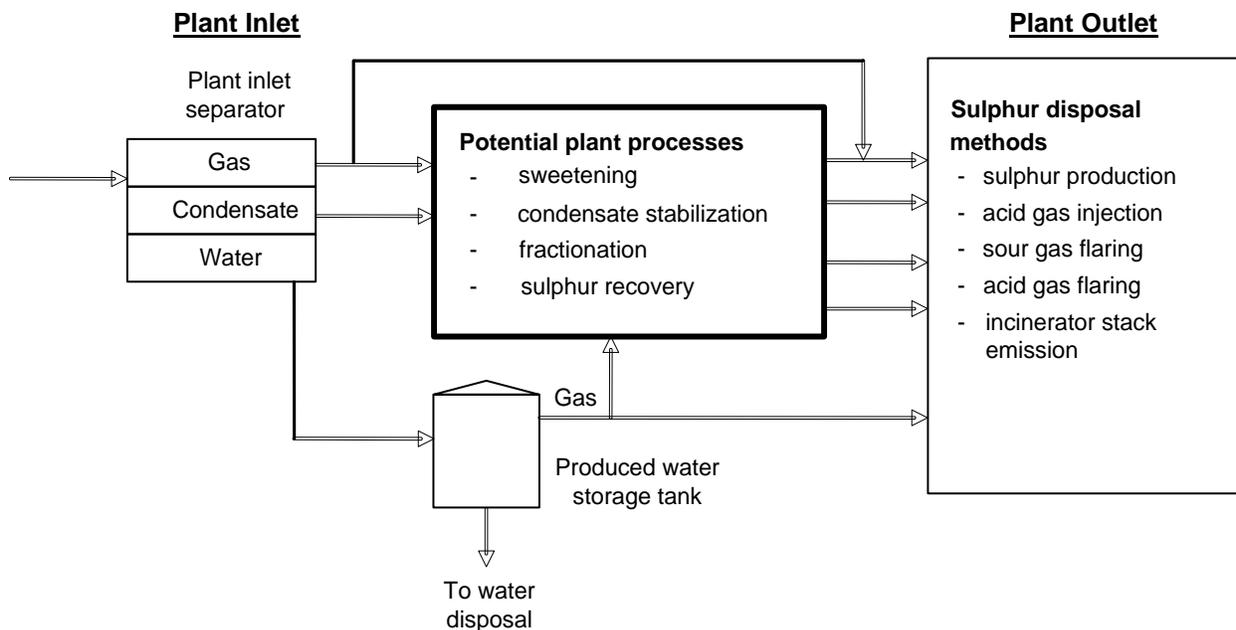
### 11.4.1 Determining H<sub>2</sub>S in Sour Gas

See Section 8.2 for the determination of H<sub>2</sub>S concentration in the inlet gas stream.

### 11.4.2 Determining the Concentration of H<sub>2</sub>S in Condensate

Condensate associated with sour gas will contain some H<sub>2</sub>S. The physical relationship between the concentration of H<sub>2</sub>S in the gas and in the condensate depends on the composition of the gas and the condensate and the pressure and temperature in the plant inlet separator. The concentration of the H<sub>2</sub>S in the condensate is usually determined in a laboratory on condensate samples obtained from the inlet separator.

**Figure 11.2 Sour gas plant process overview**



As long as the gas and condensate entering a sour plant originate from a single pool, the H<sub>2</sub>S concentration in the condensate will likely remain quite stable at the sampled conditions of pressure and temperature. Minor changes in pressure and temperature of the separator will only have a slight influence on the composition of the condensate. If major changes in pressure occur, such as due to installation of plant inlet compression and the resultant lowering of the inlet pressure, new samples must be taken and analyzed.

If the production to the plant occurs from two or more pools with different reservoir fluid compositions, the composition of the condensate will vary. In such scenarios, a vapour/liquid equilibrium correlation between the mole fraction of H<sub>2</sub>S in the sour gas and the condensate can be used to estimate the mole fraction of H<sub>2</sub>S in the condensate based on compositional analysis, computer process simulation, or stabilizer overhead volume and percentage of H<sub>2</sub>S.

### 11.4.3 Determining the Concentration of H<sub>2</sub>S in Inlet Separator Water

The concentration of H<sub>2</sub>S dissolved in the water that enters the plant inlet separator is a function of the separator pressure and temperature, as well as the concentration of H<sub>2</sub>S in the separator sour gas. The amount of H<sub>2</sub>S dissolved in the water can be predicted quite readily with correlations based on compositional analysis or computer process simulation.

Water can be metered ahead of the liquid level control valve on the water outlet line from the separator, or it can be gauged in a low-pressure produced water storage tank. The vapours from the tank are usually swept from the tank to flare with sweet gas. Some plants producing large amounts of sour water have installed a vapour recovery system, by which the tank vapours are conserved and treated in the processing plant.

It is recognized that a portion of the H<sub>2</sub>S in the water of the plant inlet separator will remain in the water when the water is disposed of in a disposal well. This amount of H<sub>2</sub>S is small and will depend on the water temperature in the produced water storage tank at atmospheric pressure, the amount of agitation in the water, and whether sweet gas is used for sweeping the vapours from the tank to flare. The small amount of H<sub>2</sub>S remaining in the water is difficult to estimate and therefore need not be included in the disposal accounting.

### 11.4.4 Calculation Procedure for Estimating the Plant Sulphur Inlet Mass per Day

1. The following streams must be accurately metered:
  - a. sour gas out of the separator, Q, 10<sup>3</sup>m<sup>3</sup>/d
  - b. sour condensate out of the separator, converted to gas equivalent volume, 10<sup>3</sup>m<sup>3</sup>/d
  - c. sour water out of the separator or into the storage tank, m<sup>3</sup>/d.
2. The sulphur content in the sour gas out of the separator can be calculated:  
Sulphur equivalent in sour gas, tonne/day = (Q, 10<sup>3</sup>m<sup>3</sup>/d) x (y) x (1.35592)  
where y is the mole fraction of H<sub>2</sub>S
3. The condensate must be sampled and analyzed semi-annually as a minimum frequency, in accordance with Section 8. When there are continuous gas analyzers and the H<sub>2</sub>S content in the gas stream changes, the sulphur content in the condensate out of the separator can be calculated on the basis of the mole fraction of H<sub>2</sub>S in the separator gas. The following formula should be used. Any alternative methods used must be supported by documentation that it is equivalent to the method outlined in item 1 & 2 and made available to ER upon request.

$$x = y \div K$$

where: x = mole fraction of H<sub>2</sub>S in the separator condensate

y = mole fraction of H<sub>2</sub>S in the sour gas in the plant inlet separator

$$K = A + (B-A) \times (T/66)^{1.2}$$

$$A = -0.7034 (\text{LOG}_{10}(P))^3 + 9.1962 (\text{LOG}_{10}(P))^2 - 39.58 \text{ LOG}_{10}(P) + 56.695$$

$$B = -3.9694 (\text{LOG}_{10}(P))^3 + 46.021 (\text{LOG}_{10}(P))^2 - 178.95 \text{ LOG}_{10}(P) + 234.35$$

T = temperature of the sour gas in the plant inlet separator or metering temperature, °C

P = pressure of the plant inlet separator, kPa(a)

Sulphur equivalent in condensate, tonne/day = (Gas equiv. of condensate, 10<sup>3</sup>m<sup>3</sup>/d) x (x) x (1.35592)

The range of applicability of item 3 for determining x, the mole fraction H<sub>2</sub>S in condensate, is between 700.0 to 9000.0 kPa(a) and 0.0 to 80.0°C.

- The amount of H<sub>2</sub>S dissolved in the water, z (mole fraction), in the plant inlet separator can be estimated by the following formula<sup>6</sup>:

$$z = y \div ((4.53 - 7494.6) \div (P + 758.4 (1.8 T + 32))) \div (P + 4.65 \times y)$$

where all terms are as defined in this section.

Sulphur equivalent in water, t/d, = (1.31) x (water production, m<sup>3</sup>/d) x (z) x (1.35592)

The sum of the results of items 2, 3, and 4 for each sour gas inlet separator is the total sulphur inlet to the plant in tonne/day.

#### 11.4.5 Calculation Procedure for Estimating Plant Sulphur Outlet Mass per Day

There are three sulphur disposal methods approved by ER:

- Sulphur recovery
- Acid gas flaring
- Acid gas compression and injection

Each of these methods is treated separately as far as collecting the disposition data. The plant inlet data are collected identically for the different sulphur disposal methods.

Potential sulphur disposal methods from sour gas plants are:

- Sulphur recovery
- Sour gas flaring or incineration
- Acid gas flaring or incineration
- Sulphur plant incinerator stack emissions

<sup>6</sup> Froning, H. R. Jacoby, R. H., and Richards, W. L., "New K-Data Show Value of Water Wash," Hydrocarbon and Petroleum Refiner, April 1964, pp. 125–130.

5. Sour gas flaring or incineration from the produced water storage tank (> 0.05 tonne/day)

The disposal of the sulphur by any of these methods must be accounted for. This requires measurement of flow rates and knowledge of concentrations of H<sub>2</sub>S in the gas streams.

An important feature of the sulphur measurement on the outlet side is the determination of the H<sub>2</sub>S content of the acid gas out of the reflux drum. This gas stream is fully saturated with water vapour at the operating pressure and temperature of the reflux drum. Depending on what method is used in the determination of the H<sub>2</sub>S content, the results can be on a dry basis or a wet basis. The licensee must determine on which basis the analysis is determined.

The water content of the acid gas out of the reflux drum can be estimated by the procedure in Section 11.2.1.

Any H<sub>2</sub>S determination and any complete analysis of the acid gas stream from the reflux drum presented on a dry basis must be normalized to a wet basis by the inclusion of the water vapour mole fraction. If the H<sub>2</sub>S content in the acid gas is determined on a wet basis, the water vapour content is simply included as calculated previously in Section 11.2.1.3. In any scenario, the wet acid gas composition is to be used in the metering calculations of the acid gas stream at low pressure. This stream is then converted to a dry basis for reporting purposes.

#### 11.4.5.1 Approved Sulphur Recovery Plants

The production of liquid sulphur must be determined by gauging the liquid sulphur level in sulphur production and storage pits or from weigh bills of shipments by truck or sulphur railroad tank cars, plus inventory changes in the pit.

Meters designed for the expected flow conditions and range must be used to measure sweet and sour gas flared if the average flow rate is > 0.5 10<sup>3</sup>m<sup>3</sup>/day on an annual basis. This generally requires a high turndown ratio meter or a combination of a high-range and a low-range meter. A separate acid gas meter designed for the expected flow conditions and range must be used to measure acid gas flared, either continuously or in emergencies, from gas sweetening systems regardless of volume.

The emissions from the sulphur plant emission stack must be monitored for flow rate and SO<sub>2</sub> concentrations.

The emissions of vapours from the produced water storage tank are those that were estimated to be contained in the produced sour water in the plant inlet calculations and must also be reported as flared gas if the volumes are > 0.05 tonnes/day. If the vapours from the water storage tank are recovered through a vapour recovery unit and are injected into a sour plant process stream, they do not form a separate part of the sulphur outlet of the plant but would still be a part of the sulphur inlet.

The total sulphur out of the plant must include:

1. The sum of any liquid production
2. Any amount flared
3. Any amount contained in emissions related to the stack and tank vapours.

The acid gas sent to the sulphur plant is to be reported as shrinkage (SHR), and acid gas flaring at the plant is to be reported as FLARE on Petrinex.

#### 11.4.5.2 Approved Acid Gas Flaring Plants

Plants approved for flaring of acid gas must meter the acid gas leaving the reflux drum of the sweetening process train. The volume calculation procedure must include the effect of the water vapour content in the acid gas at the reflux drum pressure and temperature. The concentration of the H<sub>2</sub>S content of the acid gas stream must be determined at least once per week by a qualified person using the Tutweiler technique or gas chromatography and by applying the calculation procedure to determine the H<sub>2</sub>S concentration in the acid gas. Plants slipping CO<sub>2</sub> into the sales gas, or receiving sour gas from different pools having different H<sub>2</sub>S concentrations in the sour inlet gas, may need to determine the H<sub>2</sub>S concentration in the acid gas more often than once per week. A record of the H<sub>2</sub>S analysis determinations must be maintained for inspection by ER upon request.

Meters designed for the expected flow conditions and range must be used to measure sweet and sour gas flared if the average flow rate is  $> 0.5 \times 10^3 \text{m}^3/\text{day}$  on an annual basis. This generally requires a high turndown ratio meter or a combination of a high-range and a low-range meter.

The emissions of vapours from the produced water storage tank are those that were estimated to be contained in the produced sour water in the plant inlet calculations and must also be reported as flared gas if the volumes are  $> 0.05$  tonnes/day. If the vapours from the water storage tank are recovered through a vapour recovery unit and are injected into a sour plant process stream, they do not form a separate part of the sulphur from the plant.

The sum of the Sulphur, noted in the paragraphs above, must be the sulphur out of the plant. The acid gas flared must be reported on Petrinex as flared acid gas (product type: ACID GAS and activity: FLARE).

#### 11.4.5.3 Approved Acid Gas Injection Plants

Plants approved for injection of acid gas into downhole injection wells must meter the acid gas leaving the reflux drum of the sweetening process train or at some point in the process piping at the plant site. If the gas is metered before the first stage of compression, the volume calculation procedure must include the effect of the water vapour content in the acid gas at the reflux drum pressure and temperature. The concentration of the H<sub>2</sub>S content of the acid gas stream must be checked at least once per week by a qualified person using the Tutweiler technique or gas chromatography and by applying the calculation procedure to determine the H<sub>2</sub>S concentration in the acid gas. Plants slipping CO<sub>2</sub> into the sales gas or receiving sour gas from different pools having different H<sub>2</sub>S concentrations in the sour inlet gas may need to determine the H<sub>2</sub>S concentration in the acid gas more often than once per week. A record of the H<sub>2</sub>S analysis determinations must be maintained for inspection by ER upon request.

Once the acid gas is compressed, it must be measured before injection into each well. If there is more than one injection well, each well must have its own injection measurement at the well site. If sour water is injected together with the acid gas, they must be separately measured before commingling as per the following scenarios.

##### **Scenario 1**

The acid gas meter at plant/facility is before compression. The Injection wellhead meter is required and can be used as the receipt point of the injection facility.

##### **Scenario 2**

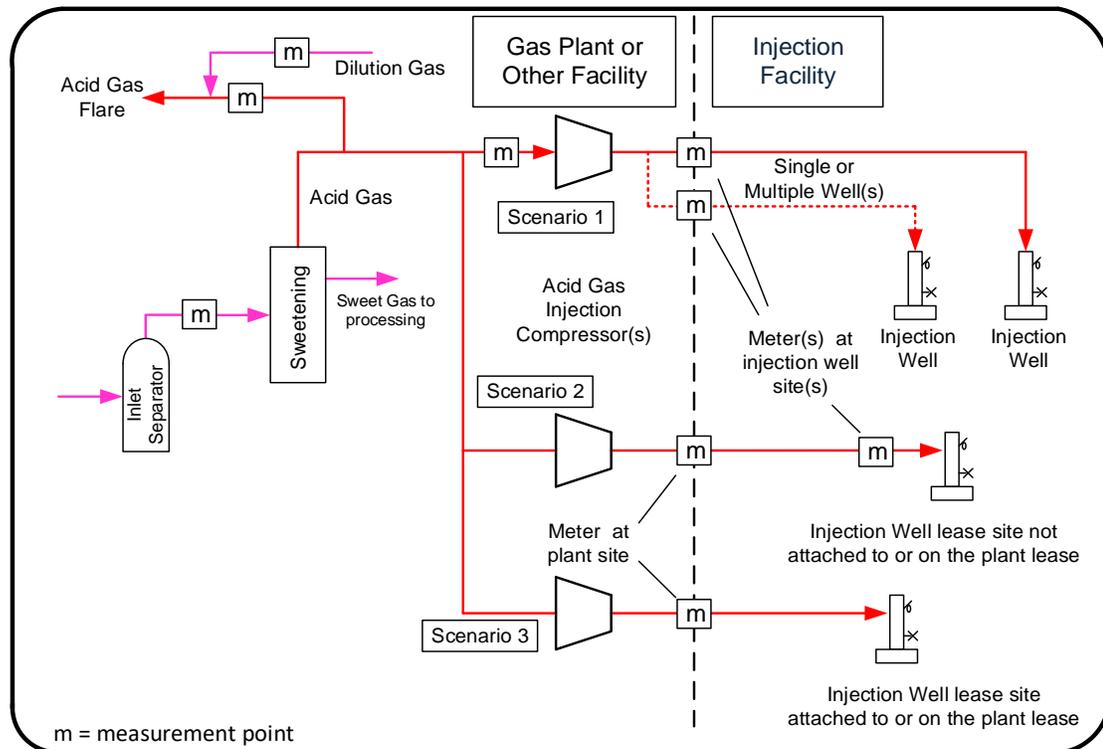
The acid gas meter at plant/facility is after compression and the injection well lease site is not attached to or on the same plant/facility lease. This acid gas meter can be used as

the receipt point of the injection facility and the injection wellhead meter is required. The metering difference is to be reported at the injection facility.

### Scenario 3

The acid gas meter at plant/facility is after compression and the injection well lease site is attached to or on the same plant/facility lease. This acid gas meter can be used as the receipt point of the injection facility and the injection wellhead meter is not required.

**Figure 11.3 Acid gas injection measurement scenarios**



Meters designed for the expected flow conditions and range must be used to measure sweet and sour gas flared if the average flared gas flow rate is  $> 0.5 \times 10^3 \text{ m}^3/\text{day}$  on an annual basis. This generally requires a high turndown ratio meter or a combination of a high-range and a low-range meter. A separate acid gas meter designed for the expected flow conditions and range must be used to measure acid gas flared, either continuously or in emergencies, from gas sweetening systems regardless of volume.

The emissions from the vapours from the produced water storage tank are those that were estimated to be contained in the produced sour water in the plant inlet calculations and must also be reported as flared gas if  $> 0.05 \text{ tonne/day}$ . If the vapours from the water storage tank are recovered through a vapour recovery unit and are injected into a sour plant process stream, they do not form a separate part of the sulphur out of the plant.

The sum of the sulphur noted in the paragraphs above must be the sulphur out of the plant. The acid gas injected is to be reported as a disposition (DISP) of ACID GAS to the injection facility, and acid gas flaring at the plant is to be reported as FLARE of ACID GAS on Petrinex.

## 11.5 Production Data Verification and Audit Trail

The field data, records, any calculations or estimations, and EFM records relating to ER-required production data submitted to Petrinex must be kept for inspection upon request by ER. The records verification and audit trails must be in accordance with the following:

1. Proving/calibration records:
  - a. Any records and documentation produced in the proving/calibration of meters.
  - b. Calibration of the prover and all peripheral devices if the prover and peripheral devices are owned and operated by the licensee.
2. Delivery and receipt records: Any records and documentation produced in the determination of delivery or receipt volumes/tonnage.
3. Estimation records: Any records and documentation related to the estimation of reported volume/tonnage, including estimation methodology, record of event, and approvals.
4. Pit gauging records: Any records and documentation produced in the determination of reported volume/tonnage.
5. Volume/tonnage loss records: Any records and documentation for volumes lost due to incidents such as spills and fires.
6. EFM: Any records and documentation including electronic, magnetic, or paper form produced in the determination of measured volume/tonnage in accordance with the EFM requirements in Section 4.4 for gas and Section 14.9.2 for liquids.

## 12 Heavy Crude Oil Measurement

This section presents the requirements and exceptions for heavy crude oil facilities. This section does not cover heavy crude oil production through mining. The term heavy crude oil is defined as oil with a density  $\geq 920.0$  kg/m<sup>3</sup>.

There are two general categories for the production of heavy oil:

1. primary/secondary which includes cold production and water flood techniques; and
2. *in situ* thermal and solvent schemes.

Blending of heavy oil with lighter hydrocarbon liquids may occur. The lighter hydrocarbon liquids, often called diluent or condensate, are used to reduce the viscosity of the heavy oil and make it easier to process and/or transport.

### 12.1 General Measurement Requirements

In all scenarios, a licensee must measure produced heavy oil volumes, unless otherwise stated in this Directive. ER will consider a heavy oil measurement system to be in compliance if the fuel measurement requirements in Sections 1.6.3.5 and 4.2, the calibration and proving requirements in Section 2, and measurement requirements in Section 14 and in this section are met.

#### 12.1.1 Temperature Correction Requirements

See Section 14.4: Temperature Correction Requirements

##### 12.1.1.1 Exception for Temperature Corrections

If testing heavy oil by test tanks, temperature correction is not required. However, the licensee must be aware that if the temperatures are elevated above standard conditions, the proration factors will be biased by the amount of the temperature correction, approximately 0.07 per cent per degree Celsius (%/°C) at a density of 920.0 kg/m<sup>3</sup>.

#### 12.1.2 Pressure Correction Requirements

See Section 14.5: Pressure Correction Requirements.

#### 12.1.3 EFM Requirements

See Section 14.9.2: Electronic Flow Measurement for Liquid Systems.

#### 12.1.4 Diluent/Condensate Receipts and Blending Shrinkage

Blending shrinkage occurs when two hydrocarbon liquids of different densities are mixed. This mixing results in volumetric discrepancies from the ideal combination, which would yield a volume that would be the sum of the two products. The discrepancy is blending shrinkage, which is the result of smaller molecules of the lighter hydrocarbon filling in the voids or spaces between larger molecules of the heavier hydrocarbon. The result is a combined liquid volume that is less than the sum of the two original volumes. This shrinkage must be determined and properly applied to volumes making up the liquid to ensure proper allocation and reporting.

### **12.1.5 Hydrocarbon Blending and Flashing Shrinkage**

Generally, heavy oil is not significantly affected by shrinkage caused by flashing of light ends. However, there is always potential for flashing shrinkage depending on the actual operating pressure at which the wells are produced. Licensees must evaluate each facility based on its operating characteristics. See Section 14.3 for details.

### **12.1.6 Water Measurement**

Licensees must have proper procedures in place to measure produced water, water receipts, and water injection/disposal to ensure that the information used for reporting is accurate. See Section 1 for measurement accuracy requirements.

As sand may constitute a significant percentage in the S&W content of heavy oil production, licensees must follow Section 14.8 and 8.5 requirements for sampling and analysis to quantify these volumes. The sand volume is included as part of the S&W determination and reported as water.

### **12.1.7 Well Proration Testing**

For heavy oil well testing, licensees must follow the testing frequency requirements in Section 6.4.4. For additional heavy oil well testing requirements refer to Section 12.2.5.

### **12.1.8 Heavy Oil Receipt, Delivery, or Sales**

This includes the delivery or receipt of heavy oil (diluted or non-diluted), diluent, or other hydrocarbon products. For the single point measurement uncertainty of these measurement points see Section 1.7.4.

## **12.2 Primary and Secondary Production**

Well effluent produced from heavy oil wells using natural or other drive mechanisms, such as water-flood wells, flowing wells, or wells with conventional lifting technologies such as pump jacks, progressive cavity pumps, and submersible pumps, is considered primary and secondary production. The well effluent could be flow lined to field test satellites or group production facilities, or the fluids could be produced to tanks at surface and trucked out to a treatment facility. These production wells are subject to the same measurement and reporting requirements as non-heavy oil wells, refer to Section 6, unless otherwise noted in this section.

### **12.2.1 Battery Subtypes**

Battery subtypes are the same as in Section 6.2.2, with the following exceptions:

#### **12.2.1.1 Heavy Crude Oil Single-Well Battery (Petrinex facility subtype: 325) (Disposition = Production)**

Oil and water volumes trucked from lease production tanks must be used to calculate the well production reported on Petrinex. Where a well is producing as a single-well battery (SWB) to a lease tank, the lease tanks can be considered as part of the reservoir and the inventory in the tanks is not reported. This procedure is referred to as the disposition equals production accounting method. This applies to heavy oil production from within an area north of Township

21 and west of Range 4 W3. Oil and water production are only reported when the fluids are trucked from the lease tank to another location.

If a well is on a restricted gas production order or has unrecovered load fluids, reporting of oil and gas production must be done monthly based on inventory change. If there is receipt(s) into the single-well battery, monthly production must be reported based on receipt and inventory volumes and the disposition equals production methodology does not apply.

When using the disposition = production accounting method, it is correct to show hours on production and no production volume if a shipment was not made from a lease tank of a producing well during the reporting period. Conversely, produced fluid removed from a lease tank during a month that a well is shut in is to be shown on the reports as with zero hours of production. Suspended and abandoned heavy oil batteries can report dispositions (and production) up to six months after they become inactive.

#### **12.2.1.2 Heavy Crude Oil Paper Battery (Petrinex facility subtypes: 313)**

Paper batteries are developed to reduce the number of reporting entities for submission to Petrinex. Paper batteries are made up of multiple single heavy oil wells that are grouped, for reporting purposes, into one larger battery. This allows licensees to report production and dispositions from a number of individual wells in one battery, thereby reducing the administrative burden. Paper batteries are treated as multiwell group batteries even though the single wells are not on a common production site.

It is acceptable to move fluids between locations within the same paper battery. These volume movements are not reported to Petrinex but must be managed in a field data capture (FDC) system.

Wells in a paper battery are eligible for reporting using the disposition = production methodology (see Section 12.2.1.1).

As paper batteries are for administrative ease, they must comply with the following criteria:

1. All wells linked to a paper battery must:
  - i. Have an oil density  $\geq 920.0 \text{ kg/m}^3$  and be located north of Township 21 and west of the Third Meridian, in Saskatchewan;
  - ii. Not be connected by flowline to a battery and not be part of an approved Enhanced Oil Recovery project, waterflood project, or any other approved project;
  - iii. Have common working interest ownership and either common crown or common freehold royalties.
  - iv. Receive the same well-head price from the sale of crude oil during the month (i.e. Averaging of gross prices or trucking charges from multiple sales transactions associated with different wells in a paper battery is not be permitted. This means that all oil delivered for sale from any of the single wells linked to the paper battery must receive the same net price at the well-head. This will eliminate averaging of gross prices and trucking charges which creates inequities with respect to the calculation of royalties and taxes on both Crown and Freehold lands.);
  - v. Be within a geographic area no larger than six contiguous townships; and

2. VENT, FLARE, FUEL and EMIS must not be reported at the paper battery level. Any VENT, FLARE, FUEL and EMIS activity must be reported at the location where it physically occurred (i.e. at the well)
3. The licensee must use one of the single-well battery well locations within the paper battery for reporting.

#### **12.2.1.3 Heavy Crude Oil Multi-well Group Battery (Petrinex facility subtypes: 326)**

Each well must have its own separation and measurement equipment similar to a single well battery. The disposition = production accounting method may be used for each well subject to the same limitations as a heavy crude oil single well battery.

All equipment for the wells in the battery must share a common surface location.

If the disposition = production method is not used or cannot be used, inventory at the group battery is to be reported monthly to Petrinex.

#### **12.2.1.4 Heavy Crude Oil Multi-well Proration Battery (Petrinex facility subtypes: 327)**

All well production is commingled prior to the total battery oil/emulsion being separated from the gas and measured. Individual monthly well oil and water production are estimated based on periodic well tests and corrected to the reported monthly volume through the use of a proration factor as stated in Section 6.4.4.

Inventory at the proration battery is reported monthly to Petrinex.

Wells included in a proration battery are not eligible for the disposition = production accounting methodology.

Double proration accounting is permissible, if conditions of Section 6.2.2.3 are met.

### **12.2.2 Gas Estimation and Reporting for Heavy Oil Wells**

As per Sections 4.1, fuel gas volumes at heavy crude oil facility subtypes 313, 325, 326, and 327 that are  $\leq 0.5 \text{ } 10^3 \text{ m}^3/\text{day}$ , based on an annual average, may be determined by estimation instead of metering. If the annual average fuel gas is  $> 0.5 \text{ } 10^3 \text{ m}^3/\text{day}$ , then the fuel gas must be metered. Other gas streams, including gas production, casing gas, flare gas (including incineration), vent gas, and stock tank vapours from produced oil/emulsion tanks and produced water tanks that are  $\leq 2.0 \text{ } 10^3 \text{ m}^3/\text{day}$ , based on an annual average, may be determined by estimation instead of metering. If the annual average gas rate for these streams is  $> 2.0 \text{ } 10^3 \text{ m}^3/\text{day}$ , then the gas must be metered.

When the gas production rate from a heavy crude oil well is  $\leq 2.0 \text{ } 10^3 \text{ m}^3/\text{day}$ , the well gas production may be estimated. The only acceptable estimation method for gas production is using a Gas Oil Ratio (GOR) factor that is applied to the reported monthly oil production. A well's GOR is determined by conducting a well test and then dividing the test gas volume by the test oil volume to determine the GOR. The determination of a well's GOR may include some or all of the gas volumes originating from the following sources as applicable:

- Measured well test separator (or dry flow meter) gas.
- Measured casing gas.
- Flare (including incineration), fuel and vent gas volumes

- Gas in solution (GIS) determined from a GIS factor obtained from a separator or wellhead oil sample under pressure and then sending it to a lab to undergo a PVT analysis or a flash liberation analysis. Although rare, it is possible that a well's GIS gas will be the entire gas volume used in the determination of a well GOR.
- Stock tank vapours from produced oil/emulsion tanks and produced water tanks.

All applicable well gas production sources must be included in the determination of the well's GOR and the number and location of those sources is dependent on the specific operational configuration of the well. A common oversight in the determination of a well's GOR is to not include the well's gas production attributed to the gas that evolves from the oil/emulsion when the oil is depressurized from separator or wellhead pressure to stock tank conditions...the Gas in Solution (GIS). The GIS is typically just one of the gas production sources that must be included in the determination of a well's GOR.

Acceptable methods for determining a GOR factor can be found in Section 12.2.3.

In some scenarios a gas volume must be determined, such as GIS, where the gas is dissolved in an oil volume under pressure, and there is no opportunity to measure the gas volume prior to it being commingled with other gas volumes. In that scenario, the gas volume may be determined by estimation. An example of such a gas volume is the gas held in solution with oil volumes leaving a test separator at an oil proration battery, where the test oil volumes are combined with production from other wells downstream of the test separator. The purpose of estimating the gas in solution is to determine the total gas produced by a well during a production test, since the gas volume measured by the test gas meter will not include the gas that is still in solution with the test oil volume.

A single GIS factor may be determined and used to estimate the gas volume held in solution with the oil stream for each oil stream where the production sources (producing formation) are the same and test separator operating conditions are similar. Additional GIS factors are required for wells in the battery that produce from different formations and where other test separators operate at different pressure and/or temperature conditions. Licensees should also consider determining seasonal GIS factors where ambient temperature differences may significantly affect the factors or when operating conditions change significantly. Acceptable methods for determining the GIS are described in Section 12.2.3.

For single point measurement uncertainty refer to Section 1.7.4.

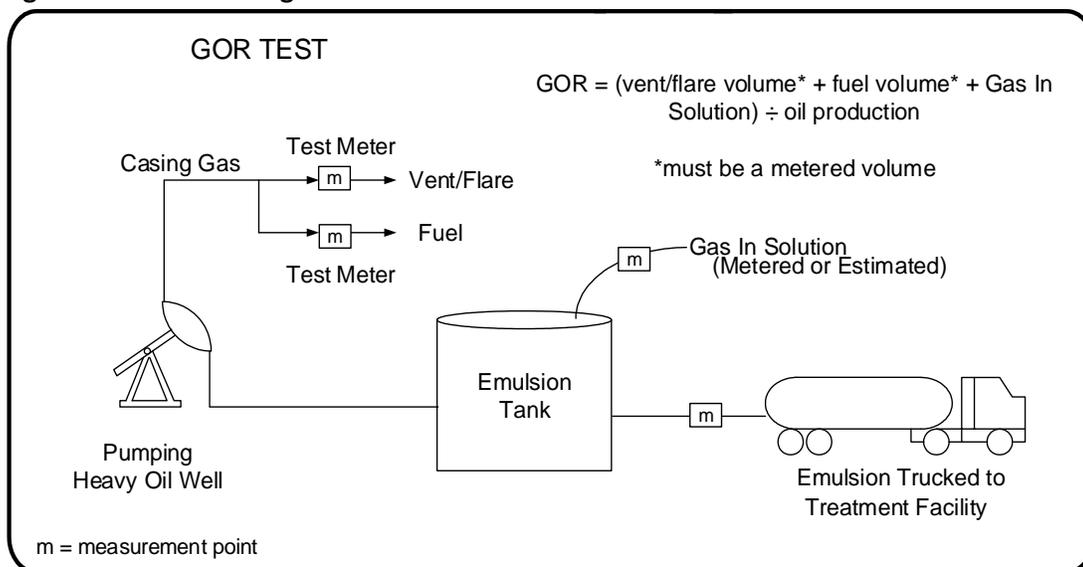
#### **12.2.2.1 Heavy Crude Oil Single-Well Batteries (Petrinex facility subtypes 313 and 325) and Heavy Crude Oil Multi-well Group Batteries (Petrinex facility subtype 326)**

For new heavy crude oil wells, a GOR test should be conducted as soon as is reasonably possible after the well is put on production and well production rates have stabilized, however, notwithstanding the aforementioned, the first GOR test must be conducted within the first six months of the well being put on production. It is acceptable to use the GOR from an analog well producing from the same pool until the first GOR test is conducted. Following the first test, the GOR test frequency must be in accordance with the frequency outlined in Table 12.1.

The procedure used to determine the GOR factor must be in accordance with Section 12.2.3 and the GOR test must be conducted such that all the produced gas, including fuel, flare and vent gas and oil volumes produced during the test are measured (see Figure 12.1). The method to

determine the GOR factor must include the GIS and the method used to determine the GIS must be in accordance with Section 12.2.3.

**Figure 12.1 GOR testing schematic**



GOR test frequencies are detailed in [Table 12.1](#).

**Table 12.1 GOR test frequency requirements**

Gas rate ( $10^3\text{m}^3/\text{d}$ )	Test frequency
$\leq 1.0$	Annually. Due to requirement changes in 2020, licensees who have not already determined a GOR factor as per section 12.2.3 within the 2020 calendar year must: <ul style="list-style-type: none"> <li>determine the GOR factor before the end of the 2020 calendar year</li> <li>use the new GOR factor to report volumes; and</li> <li>the licensee must continue with the GOR factor test frequency onwards.</li> </ul>
$> 1.0$ and $< 2.0$	Semi-annually

**12.2.2.2 Heavy Crude Oil Multi-well Proration Batteries (facility subtype 327)**

For a heavy crude oil multi-well proration battery (facility subtype 327), each well with gas production  $> 2.0 \times 10^3\text{m}^3/\text{day}$  must be well proration tested and follow well proration testing frequency requirements in Section 12.1.7 and Section 12.2.5, unless metered, on a per stream basis. Fuel gas, vent gas or flare gas at a site must be reported at the location at which the activity is occurring.

Gas production volumes  $\leq 2.0 \times 10^3\text{m}^3/\text{day}$  may be estimated using one of the following methods:

1. A well-level GOR at the same initial frequency as a heavy crude oil single well battery and then in accordance with Table 12.1.
2. Battery-level GOR. A battery-level GOR factor (monthly battery gas production  $\div$  monthly battery oil production) must determined on a monthly basis. To calculate the individual

well gas production using the battery-level GOR factor the following conditions must be met:

- a. All wells using the battery-level GOR must produce  $\leq 2.0 \times 10^3 \text{m}^3/\text{day}$  of gas;
- b. Any well producing  $> 2.0 \times 10^3 \text{m}^3/\text{day}$  of gas is not eligible to use the battery-level GOR, and well gas production must be determined using test rates obtained during proration testing;
- c. Monthly gas and oil volumes from wells not eligible to use the battery-level GOR must be subtracted from the total battery gas and oil volumes before calculating the battery-level GOR. For gas, the volume to be subtracted would be the total estimated gas determined from proration testing for all the ineligible wells; for oil, the volume would be the total prorated oil production for all the ineligible wells;
- d. New wells added must produce  $\leq 2.0 \times 10^3 \text{m}^3/\text{day}$  of gas for a minimum of six months before being eligible to use the battery-level GOR;
- e. If there is no common ownership of all the wells in the battery, written notification must have been given to all working interest participants, with no resulting objections; and
- f. If there is no common Crown or common Freehold royalty and only Freehold royalties are involved in all wells in the battery, written notification must have been given to all Freehold royalty owners, with no resulting objection received. If there is a mix of Freehold and Crown royalties involved, the licensee must apply to ER for approval if any Freehold royalty owner objects.

### 12.2.3 Method for Determining Gas Oil Ratio Factor and GIS

Refer to Section 6.5.3 for a description of the procedure and the requirements regarding how to conduct a GOR test and determine GIS. In addition to those requirements, the following additional requirements also apply when conducting a GOR test on a heavy crude oil well:

1. The GOR test duration must be, at minimum, 72.0 hours. Licensees may be exempt from conducting a 72-hour GOR test and instead be allowed to conduct the measurement to determine the gas volume over a continuous period of at least 24-hours, if the qualifying criteria are met.

#### Qualifying Criteria

The gas measurement is taken:

- i. With a continuous measuring device and the variation of flow rate in that continuous period is steady state or a representative flow. Steady state or representative flow rate is such that the average flow rate for any 20-minute period is within  $\pm 5$  per cent of the average flow rate; or
- ii. Using a flow meter (e.g. flow totalizer) with at least one reading taken every 20 minutes within that continuous period. The result must show 95 per cent of the readings as steady state or representative flow as per (i) above.

### Revocation of Exception

If any of the following scenarios exists or occurs, the exception is revoked, the well must be GOR tested as soon as is reasonably possible, the GOR testing duration reverts to 72-hour GOR test and the well must requalify for the exception:

- i. A licensee fails to provide test records to ER upon request which must include the 20 minute readings that prove that the well met the qualifying criteria as well as any additional requested documentation related to the GOR test.
  - ii. If any changes are made to the operations of the facility that could affect gas production such as, but not limited to, fracturing nearby wells, pressure changes at the well head, new equipment being installed at the facility, or workovers on the well.
2. Before the GOR test is conducted there can be no adjustment to the well or operating conditions for at least 48.0 hours before the test that could result in a change to the oil or gas production rates.
  3. Gas sources that must be considered in the determination of the GOR include but are not limited to casing gas, fuel gas, vent gas, flare gas and gas vented from tanks.
  4. The volume of oil produced during the test may be determined using a two or three-phase separator and the liquid hydrocarbon meter must meet the single point measurement uncertainty of  $\pm 2.0$  per cent. The water cut may be obtained using a proportional sampler or an online water cut analyzer as per Section 14.8. Manual sampling (i.e. start of test, middle of test and end of test) for analysis of water content is allowed if the water cut is  $\leq 0.5$  per cent.

If a separator cannot be used to conduct the GOR test and the well effluent is flowlined directly from the wellhead to a production tank then the GOR test oil production volume may be determined by using the monthly well production volume for the same month the gas volume is determined.  $((\text{oil disposition} + \text{tank inventory change}) \div \text{number of hours the well was on production for the month}) \times \text{GOR test duration in hours}$ .

5. The continuous measuring device or flow meter used to determine the GOR Factor must meet the requirements of Section 2.
6. If the licensee is utilizing the production equals disposition methodology and a separator cannot be used, tank inventories must be taken at the start and end of the month for proper determination of the monthly well oil production volume as stated in the formula in point 4. If inventories are not taken for the month for which the GOR test was conducted, the licensee must reconduct the GOR test as soon as is reasonably possible.

The following must be documented and be made available to ER upon request, failure to provide this information will result in ER requiring the licensee to reconduct the GOR test:

- The meter readings, associated data (e.g. temperature and pressure), instantaneous flow rate and cumulative test volume for each time period (e.g. 20 minutes) that the measurements were taken for each determination of the value of gas and oil.
- The date, time, and duration of each of those periods. The maximum period duration must be no longer than 1 hour.

- The production parameters (e.g. pressure, pump rate) during each of those periods and in the 48 hours before each of those periods; and
- A description of the meter(s) and other equipment used during the test. At minimum the meter make, type, and model should be included.
- Any factors used (e.g. GIS) to determine estimated volumes and the associated documentation of the estimation methods
- If a separator cannot be used, then the oil disposition volumes and inventories must be provided that were used in the determination of the oil volume.

#### 12.2.4 Oil and Water Deliveries to a Treatment Facility

Liquid volumes are trucked from single-well batteries, multi-well batteries, and wells within paper batteries to a custom treater or battery. These facilities process oil, water, and sand for disposal from the trucked-in fluids. Typically, received products are measured using weigh scales or inlet meters but tank gauging could also be used, see Section 10.3. The oil and water densities from every well must be updated in accordance with Section 10.4.4 and the S&W of the emulsion delivered to these facilities must be determined on a per-load basis, see Section 14.8.

For single point measurement uncertainties refer to Section 1.

#### 12.2.5 Well Test Measurement with Tank Gauging or Metering

Wells in primary/secondary production of heavy oil must be tested at the same frequency stated in Section 6.4.4 for non-heavy crude oil wells. The tests must be conducted in a consistent manner throughout the month and a test must be conducted when there is a change in well parameters (pump speed, work-over, reactivation, flush-by, etc.) as soon as possible.

A water cut must be determined for each test, and collecting a wellhead sample during a test is acceptable. See Section 14.8.

Temperature correction is not required for well tests using test tanks, see Section 14.4.

Tests may be conducted by using a single isolated tank for the well. [Table 12.2](#) may be used by the licensee to determine the height-to-diameter ratio requirements in accordance with Section 14.7.2, with the exception that the accuracy coefficient in [Table 12.2](#) is a suggested minimum for test fluid volumes.

**Table 12.2 Accuracy coefficients for various measurement types for test tanks**

Level measurement technique	Accuracy coefficient "a"	Maximum level reporting resolution (mm)
Gauge board	1.6	25.0
Manual dip of the tank	0.4	10.0
Electronic (e.g., radar)	0.4	10.0

The accuracy coefficient "a" can be used in the following equation:

$$V \geq a \times d^2 \quad \text{or} \quad d \leq (V \div a)^{0.5}$$

where:

V = test fluid volume, m<sup>3</sup>

a = accuracy coefficient

d = tank diameter, m

On the tank being used for testing, the gauge board float, linkage, and scale must be in good condition. The gauge board markings must be no further than 60.0 mm apart. For gauge measurement on test tanks, one reading of the gauge board is acceptable for the start and end of the test. Where safe work conditions permit, gauge boards should be read at eye level.

If the well emulsion is to be tested using a meter, the meter must be sized to operate within 20.0 per cent to 90.0 per cent of its flow range and installed and operated in accordance with Sections 2.4 and 14.2.

For single point measurement uncertainty refer to Section 1.

### 12.3 Thermal In Situ Battery Operations (Petrinex facility subtypes: 344)

Heavy crude oil enhanced recovery typically requires the injection of steam, sometimes with added solvent or gas.

For thermal production operations, licensees deal with a variety of measurement challenges:

1. Steam injection may be of variable quality at the wellhead, resulting in systematic errors, and may also be present in the vapour state at production facilities, creating further measurement issues.
2. Most thermal projects require the use of a diluent to assist in the separation of water/oil emulsions, so production may not be directly measured at these facilities.
3. Produced fluids are at high temperatures, so all measurements must be temperature corrected except for well testing volumes.
4. At some facilities, highly abrasive fluids are observed as a result of entrained sand, which can damage meters and introduce systematic errors.
5. Injected solvent and gas may be difficult to fully differentiate from produced fluids, especially since the composition of the oil can change across a scheme area and vertically within a stratigraphic unit or zone, and can be further changed due to *in situ* high-temperature reactions.

For these reasons, the measurement plan at thermal operations must be thoroughly considered. *Guideline PNG042: Measurement, Accounting, and Reporting Plan (MARP) Requirements for Thermal Heavy Oil Recovery Projects* is an integral part of developing a thermal project in Saskatchewan. It provides information on the submission requirements of a proposed thermal project measurement and reporting plan. A MARP, approved by ER, must be obtained by the licensee prior to facility licencing.

Unless otherwise stated, all existing thermal *in situ* facilities must meet the requirements set out in this section. For new facilities and expansions to existing facilities, the steam and water measurement requirements are effective immediately.

### 12.3.1 Heavy Crude Oil, Diluent, and Diluted Heavy Crude Oil Delivery Point Measurement

Each licensee is responsible for determining the total heavy crude oil production at its scheme, regardless of whether another party is responsible for operation of the delivery point measurement. The delivery and receipts of diluent, diluted heavy crude oil or heavy crude oil all affect the final determination of production of a scheme. The following are requirements for delivery and receipt points of diluent, diluted heavy crude oil or heavy crude oil:

Receipt point measurement must meet a single point measurement uncertainty of  $\pm 0.5$  per cent and must be proved in accordance with Section 2.4 where applicable.

Density measurement must be taken on the receipt product using one of the following methods:

1. On-line densitometer or Coriolis meter
2. ASTM D4052 - Standard Test Method for Density and Relative Density of Liquids by Digital Density Meter
3. ASTM D5002 - Standard Test Method for Density and Relative Density of Crude Oils by Digital Density Analyzer
4. ASTM D1298/6822 - Standard Test Method for Density and Relative Density or API Gravity of Crude Petroleum and Liquid Petroleum Products by Hydrometer Method

Water content must be determined for delivery/receipt product using the following methods:

1. ASTM D95 - Standard Test Method for Water in Petroleum Products and Bituminous Materials by Distillation
2. ASTM D4006 - Standard Test Method for Water in Crude Oil by Distillation
3. ASTM D4377 - Standard Test Method for Water in Crude Oils by Potentiometric Karl Fischer Titration

Solids content must be determined for delivery/receipt product using the following methods:

1. ASTM D4807 – Standard Test Method for Sediment in Crude Oil by Membrane Filtration; applicable to samples with low mineral content
2. Mineral by Centrifuge Separation; applicable to samples with high mineral content.

For static testing methods, all samples for density, solids, and water must be obtained using proportional sampling and adhere to all requirements in Section 10.4.1.

Trucked in volumes must adhere to sampling requirements in Section 10.4.1 or 10.4.2.

Determination of shrinkage must be consistent with Section 14.3.

Sand volume is to be included as part of the S&W determination and reported as water.

All measurement methods must be conducted in a manner satisfactory to ER. Additional requirements may apply depending on circumstances specific to each project through the MARP approval or site-specific requests.

### 12.3.2 Gas Measurement

As per Sections 4.1, fuel gas volumes at thermal in-situ batteries facility subtypes 344 that are  $\leq 0.5 \text{ } 10^3 \text{ m}^3/\text{day}$ , based on an annual average, may be determined by estimation instead of metering. If the annual average fuel gas is  $> 0.5 \text{ } 10^3 \text{ m}^3/\text{day}$ , then the fuel gas must be metered.

Other gas streams, including gas production, flare gas (including incineration), vent gas, and stock tank vapours from produced oil/emulsion tanks and produced water tanks that are  $\leq 2.0 \times 10^3 \text{m}^3/\text{day}$ , based on an annual average, may be determined by estimation instead of metering. If the annual average gas rate for these streams is  $> 2.0 \times 10^3 \text{m}^3/\text{day}$ , then the gas must be metered.

Gas production measurement from thermal wells can be problematic, particularly at high temperatures where steam and solvent may be present with the gas and the combined flow of gas, steam, and/or solvent would be metered together.

For multi-well proration facilities, the group gas measurement can be used to prorate back to well gas test rates using one of the following methods:

1. If a test separator is used, the total battery gas is prorated back based on the individual well test gas rates after subtracting any steam and lift gas volumes from the produced gas.
2. The individual well GOR may be used in accordance with Section 12.2.2.2.
3. If test gas rates are not consistent and not used or well gas is not measured but total well fluid production and water cut can be determined at the battery, well gas production may be determined using a battery-level GOR. The battery-level GOR and the application to each well can be calculated as follows and will result in a gas proration of 1.00000:

$$\text{Battery GOR} = \frac{\text{Total monthly measured produced gas at battery}}{\text{Total monthly measured produced oil at battery}}$$

$$\text{Well gas volume} = \text{Battery GOR} \times \text{Well prorated (reported) oil volume}$$

The battery-level GOR can only be used if these criteria are met:

- a. There is common working interest ownership of all the wells in the battery;  
and
- b. There are no gas sales or use of the produced gas that would trigger a gas royalty payment.

Total gas flared and total vented gas must be metered (or estimated as applicable, using sound engineering practices) and reported.

Gas can be injected into a reservoir for a variety of different purposes over the life of a well. If gas is injected into a reservoir other than for lift gas, it must be measured and reported. If lift gas is used, the total lift gas volume must be measured and netted off the total produced gas volume before prorating to the wells.

The single point measurement uncertainty for gas produced at a battery or injected at a wellhead is  $\pm 3.0$  per cent.

When injecting gas into the reservoir, licensees must have a methodology in place to determine the resulting production of the injected gas. If a MARP already exists, an update must be submitted for ER approval prior to commencing any gas injection.

All gas measurement devices must be calibrated, proved, or verified annually or as otherwise stated in Section 2.

### 12.3.3 Steam Measurement

Direct measurement of injected steam is challenging. Licensees must contend with steam losses, unmetered utility steam, and changes in steam quality from steam generator to wellhead. Steam must be reported as a cold water equivalent (CWE) at 15.0°C. If it is delivered from another facility for injection, it must be reported on Petrinex as a receipt of the product [STEAM].

The steam injected into each well must be measured on a per-well basis and reported monthly. When measuring wellhead injected steam, the devices used must have a single point measurement uncertainty of  $\pm 5.0$  per cent at the time of installation. Wellhead injection meters with no internal moving parts must have the instrumentation calibrated annually, or before every steam cycle.

At a minimum of once every two steam cycles, licensees that use cyclic steam stimulation (CSS) as a recovery method, must visually inspect the primary element of one wellhead steam injection meter for every five wells on a pad. At a minimum of once per year, steam assisted gravity drainage (SAGD) or other thermal operations must visually inspect the primary element of one wellhead steam injection meter for every five wells on a pad, see Section 12.3.11 for exception. For example, pads that have 6 to 10 wells would require a two well sample set, 11 to 15 wells would require a three well sample set, and so forth. The well(s) chosen as a sample set on a given pad must be representative and not have been selected in the previous five years. If any of the inspected sample set meters are found to be compromised, the licensee must service (clean or replace) all wellhead injection meters included on the pad within 12 months of discovering the compromised meter. Visual evidence of the primary element condition during the meter inspections must be kept on site and made available to ER upon request.

Radiography, optical techniques, or other nonintrusive methods may be used in lieu of physical inspections provided that the images are of sufficient quality to discern any damage to the primary element or the presence of scale. If any of the inspected sample set meters are found to be compromised, the licensee must inspect (non-intrusively) all wellhead injection meters on that pad within one month of discovering the compromised meter. The licensee must service, clean or replace those meters that showed damage, scaling, or yielded inconclusive results within one month. Other verification methods will be considered on a scenario-by-scenario basis.

The wellhead steam volumes reported to Petrinex must be prorated from the total steam volume leaving the steam plant. The monthly facility proration factor and the data used in its calculation must be kept on site and made available to ER upon request. The volume of steam leaving the steam plant may be determined from:

1. Total steam volume leaving the steam plant separator; or
2. Boiler feed water and boiler blowdown.

The steam injected into each well must be measured on a per-well basis and reported monthly. Where steam chambers belonging to a single subsurface drainage area have coalesced for projects using SAGD, licensees may measure steam injected using group measurement for the associated wells in place of individual well measurement. In this scenario, steam injection must continue to be reported to Petrinex for each well using its UWI and it is the licensee's responsibility to justify to ER how individual reported well injection volumes are determined.

A maximum single point measurement uncertainty of  $\pm 2.0$  per cent is required for:

1. Steam entering or leaving the injection facility, excluding wellhead injection, for example, steam transferred to a battery or to or from another injection facility.
2. Steam used for emissions control at the injection facility, e.g., NO<sub>x</sub>, unless the volume is < 2.0 per cent of the injection facility's total water out, in which case it may be estimated using sound engineering practices.
3. Steam leaving a steam plant.

All steam measurement devices must be calibrated/proved/verified on an annual basis or as otherwise stated in this section or in Section 2.3.

#### 12.3.4 Water Measurement

The measurement of water at thermal *in situ* facilities is a key component to evaluating the plant performance and compliance with approvals and regulations. Therefore, a higher degree of accuracy is required than at conventional production operations.

A maximum single point measurement uncertainty of  $\pm 2.0$  per cent is required for the following water streams at injection facilities associated with thermal *in situ* schemes:

1. Fresh and produced water entering or leaving the injection facility;
2. Water injection/disposal;
3. Boiler feed water and boiler blowdown;
4. Water used for emissions control at the injection facility, e.g. NO<sub>x</sub>, unless the volume is < 2.0 per cent of the injection facility's total water OUT, in which case it may be estimated using sound engineering practices;
5. Camp wastewater entering an injection facility, unless the volume is < 2.0 per cent of the injection facility's total water OUT, in which case it may be estimated using sound engineering practices.
6. Freshwater diversion points licensed by Water Security Agency.

Upon application, ER may approve alternative methods for determination of the water stream volumes stated in items 1-6 if it is satisfied that they can be determined with a maximum uncertainty of  $\pm 2.0$  per cent, for example, calculating produced water by subtracting measured fresh and/or brackish water from a measured mixed water stream.

A maximum single point measurement uncertainty of 5.0 per cent is required for all other water uses reported to Petrinex, except if the water streams are small < 2.0 per cent of the total OUT in the facility water balance determination, in which case they may be estimated using sound engineering practices.

All water measurement devices must be calibrated/proved/verified on an annual basis or as otherwise stated in Sections 2.4 and 2.8.

#### 12.3.5 Water/Steam Primary and Secondary Measurement

For produced water entering an injection facility and steam leaving a steam plant, a primary measurement method using direct physical measurement must be in place. A secondary measurement method, which can be physical measurement or estimates based on sound

engineering practices, must also be in place. For example, the primary measurement for steam could be direct high pressure steam measurement or the boiler feed water (BFW) minus boiler blowdown, and the secondary measurement could be BFW times steam quality if the blowdown measurement fails.

Secondary measurement methodologies do not have to meet the calibration, inspection, or proving requirements in this directive.

If both the primary and secondary measurements are on the same pipe run, the operator must use differing measurement technologies to aid in preventing measurement failures from the same mechanism.

If the design of the metering system includes the provision to inspect, conduct maintenance, and repair the primary element without shutting in the stream flow, i.e. meter bypass, and repair of the meter can be completed within two weeks of ceasing primary measurement, then secondary measurement is not required.

### **12.3.6 Solvent Measurement**

The addition of solvent(s) to steam injection can accelerate and improve the recovery of heavy crude oil. Injected substances used as solvents can include, but are not limited to, ethane, propane, butane, and carbon dioxide or some combination of these hydrocarbons.

Individual types of solvents injected into the formation must be measured prior to mixing with steam and/or other solvents. Licensees must comply with the standards of accuracy requirements in Section 1 and the liquid solvent injection requirements stated in Section 12.3.6.

The determination of produced solvent is key to evaluating process performance and economics and determining production volumes. The determination of solvent production is complex and includes, but is not limited to, volumetric measurement, sampling, and compositional analysis.

Consequently, a Guideline PNG042- MARP application must be submitted to address the details of the solvent injection and corresponding production. If a MARP already exists, an update must be submitted and approved by ER prior to commencing any solvent injection.

Where steam chambers belonging to a single subsurface drainage area have coalesced for projects using SAGD, licensees may measure solvent injected on a group measurement basis for the associated wells in place of individual well measurement. Solvent injection must continue to be reported to Petrinex for each well using its UWI and it is the licensee's responsibility to justify to ER how individual reported well injection volumes are determined.

The single point measurement uncertainty for liquid solvent injection is  $\pm 2.0$  per cent.

All injected solvents measured in the gaseous phase are subject to requirements in Sections 12.3.1 and 12.3.2.

All solvent injection measurement devices must be calibrated/proved/verified on an annual basis or as otherwise stated in accordance with Section 2.

### **12.3.7 Production Measurement**

The need for accurate production measurement for heavy crude oil, water, and gas is no different from non-thermal facilities. However, due to the nature of the heavy crude oil in a high temperature production environment, it is impractical to prove meters in the field in a manner used for non-thermal heavy crude oil. Proving trucks cannot handle high temperature

fluids or the high viscosity of cooled heavy crude oil. Therefore, for thermal *in situ* heavy crude oil production, annual verification of measurement devices is sufficient provided that the facility proration factor is within the targets specified in Section 3. If targets are not being met, more frequent verification may be required. For delivery point measurement, sales, or LACT, see Sections 12.3.1 and 14.

Verification can be achieved using:

1. Internal diagnostics of the measurement device if present to check the structural integrity of the primary element;
2. Bench proving; or
3. Other ER -approved methods.

The single point measurement uncertainty requirement for emulsion is  $\pm 2.0$  per cent, excluding the effects of steam and hydrocarbon vapours, as well as the effect of S&W determination.

Determination of S&W from wells faces the same challenges as the production measurement. For this reason, it is necessary that S&W methods take into account the temperature and pressure of the sampled emulsion. All manual S&W samples must be adequately cooled to mitigate the flashing of components. If S&W is determined using an on-line analyzer, provisions to take manual samples must be present to compare against the analyzer if necessary. On-line analyzers must be calibrated annually.

### 12.3.8 Well Production Measurement

Well testing in enhanced thermal operations can only be performed under certain conditions and must be done using a test separation vessel. Licensees using test separation to determine well production must comply with the following:

1. At minimum, each production well must have one valid testing hour for every 40.0 hours the well is in operation.
2. The test duration must be optimized to obtain as many representative production well tests as possible for each month.
3. Where it has been demonstrated that steam chambers have coalesced, licensees may commingle the production of a SAGD producer with an infill production well before testing the wells. Production of oil, gas, and water must continue to be reported to Petrinex for each well and licensees are permitted to use engineering estimates to allocate tested commingled production to individual wells.
4. Sufficient time must be provided between tests to purge the test separator of the emulsion from the previous test.
5. S&W must be determined for each test.
6. Test durations and methods must be reassessed annually to verify that current practices are sufficient to obtain representative data, and this information must be made available to ER upon request.

Notwithstanding the requirements above at certain points in a high-pressure cyclic steam well's production cycle, the process fluids cannot be tested due to high temperatures. In such instances, engineering estimates may be used until tests can be conducted.

In lieu of using a test separation vessel, SAGD wells using mechanical lift may use either of the following methods to determine well production:

1. Continuous unseparated well production measurement on each well (i.e., Coriolis meter with water-cut analyzer or manual sampling)
2. Unseparated well production using periodic tests (i.e., Coriolis meter with water-cut analyzer or manual sampling)

In both cases, the battery production must be prorated back to each well. If the periodic test measurement approach is used, the well-testing criteria described above in this section must be adhered to.

Where steam chambers belonging to a single subsurface drainage area have coalesced for projects using SAGD, licensees may measure well production for the associated wells on a group measurement basis rather than individually. Licensees must continuously monitor S&W for the grouped wells. Production of oil, gas, and water must continue to be reported to Petrinex for each well using its UWI and licensees must be able to justify how they are determining individual well production to ER.

The single point measurement uncertainty for emulsion is  $\pm 2.0$  per cent for individually measured wells, excluding the effects of steam and hydrocarbon vapours, as well as the effect of S&W determination. The single point measurement uncertainty for commingled well metered emulsion is  $\pm 2.0$  per cent, excluding the effects of steam and hydrocarbon vapours, as well as the effect of S&W determination.

Determining well S&W faces the same challenges as measuring production. For this reason, S&W methods must take into account the temperature and pressure of the sampled emulsion. All manual S&W samples must be adequately cooled to mitigate the flashing of components. If S&W is determined using an on-line analyzer, licensees must take manual samples for comparison to the analyzer, if necessary. On-line analyzers must be calibrated annually.

Production measurement methods other than those described above must be approved by ER.

### **12.3.9 Proration Factors**

If the proration factor targets described in Table 3.1 are not being met, ER may require an investigation to determine why. Action required by the licensee may include, but is not limited to:

1. Verifying S&W measurement practices.
2. Verifying related fluid measurement system performance.
3. Proving or calibration of measurement equipment.
4. Inspecting the primary element for meters with no internal moving parts.

### **12.3.10 Uncertainty and Proration Factor Summary**

Refer to Section 1.7 for a summary of uncertainties for heavy oil and to Section 3.1.1.1 for a summary of proration factors for heavy oil.

### 12.3.11 Internal Inspection Exceptions

The inspection requirements stipulated in this exception only apply to measurement devices, including flow meters, used in thermal *in situ* facilities to determine volumes reported to Petrinex. If the internal components of meters used at thermal *in situ* facilities have been found to be clean and undamaged for three consecutive inspections conducted over at least three years, licensees may extend internal inspection frequencies to once every three calendar years. Licensees may not apply this exception if:

1. The meter is used for the measurement or calculation of delivery point measurement, sales, or LACT volumes, including diluent measurement, or
2. The meter is used for flare measurement. For flare meter internal inspection requirements, refer to Section 2.3.4.

Licensees applying the exception criteria must comply with the following:

1. A tag must be attached to the meter indicating that this exception is being applied and the date of the last inspection.
2. If any individual meter fails during a scheduled or nonscheduled calibration or verification or inspection the meter must requalify for the exception.
3. Licensees must keep records of the internal inspections associated with this exception for at least six years and make them available to ER on request.

## 13 Condensate and High Vapour Pressure Liquid Measurement and Reporting

This section presents the requirements and exceptions for condensate measurement and reporting associated with gas well production.

Condensate is a generic term used to describe various types of hydrocarbon liquid products such as: field condensate, condensate separator liquids, condensate stabilized liquids. See Glossary in Appendix 2 for definitions.

High vapour pressure liquids is a generic term for hydrocarbon liquids that are stored and transported under pressure at all times in a liquid phase.

### 13.1 General Measurement and Reporting Requirements

#### 13.1.1 Measurement Requirements

For measurement of condensate and high vapour pressure liquids at all accounting measurement locations within upstream oil and gas facilities, ER will consider the measurement system to be in compliance:

- a. if the measurement requirements in Sections 1.6.3, 4.1, 4.2, the calibration and proving requirements in Section 2, the design and installation of liquid (oil) measurement requirements in Section 14, the sampling and analysis requirements in Section 8, and the trucked liquid requirements in Section 10 are adhered to or
- b. if the measurement requirements of Measurement Canada for high vapour pressure liquids are adhered to.

#### 13.1.2 Reporting Requirements

In Saskatchewan, hydrocarbon liquid with a density  $\leq 780.0 \text{ kg/m}^3$  is classified as condensate and hydrocarbon liquid with a density  $> 780.0 \text{ kg/m}^3$  is classified as crude oil. Condensate production can be reported as a gas or liquid depending on how it is disposed while oil must always be reported as a liquid volume. The general rules are as follows for Petrinex reporting:

1. Oil produced and separated at a gas well or at the group measurement point of gas multi-well proration or effluent measurement batteries, must be reported as liquid OIL irrespective of whether it is measured and recombined with the gas or trucked out for disposition..
2. Condensate produced and separated at a gas well or at the group measurement point of gas multi-well proration or effluent measurement batteries must be converted to a gas equivalent volume and reported as GAS production if it is measured and recombined with the gas and sent to a gathering system or to a gas plant for further processing.
3. Condensate produced and separated at a gas well or at the group measurement point of gas multi-well proration or effluent measurement batteries is considered to be field condensate and must be reported as liquid condensate at the well level if it is trucked out for disposition or trucked to a gas plant for further processing..
4. Condensate separated at a gas gathering system group separation point and stored in a tank for disposition is considered to be pentanes plus and must be reported in Petrinex as a C5-MX volume.

5. Hydrocarbon liquids recovered from processing at a gas processing plant are considered to be by-products of processing and must be reported as specific liquid volumes.
6. The receipt and disposition of load condensate injected, recovered, separated, and stored in a tank at a gas well must be reported as COND or C5-SP volume.
7. Load condensate received from an outside source at any gas well, injected, recovered, separated, recombined with production, and sent to a gas plant must be reported as a receipt of COND or C5-SP at the facility level and a disposition of GAS to the gas plant. A corresponding disposition of COND or C5-SP to SKGE and receipt of GAS from SKGE must also be reported to balance the products at the battery.
8. Load oil received from an outside source at any gas well, injected, recovered, recombined with production, and sent to a gas plant must report an oil receipt and oil disposition at the facility level.

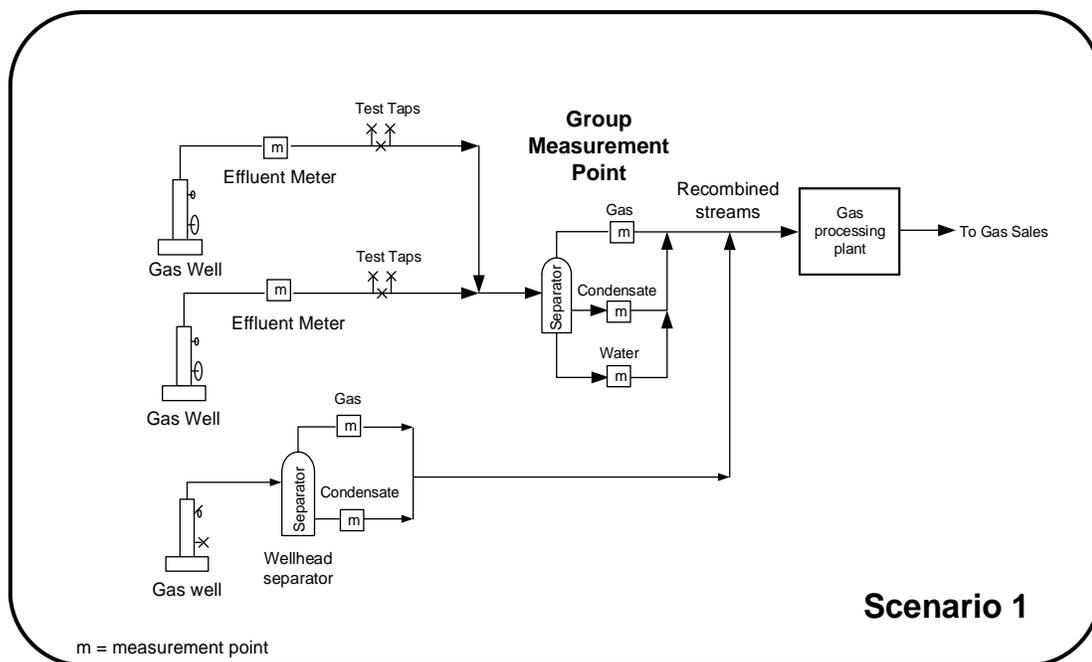
### 13.2 Reporting Scenarios for Gas Wells Producing Condensate

Condensate is associated with gas well production within gas facilities. If there is condensate, the total battery liquid volume must be measured.

#### 13.2.1 Scenario 1 – Condensate that is Effluent Metered and Tested or Proration Tested, or Separated and Recombined

Condensate being wet metered and tested or proration tested only, or separated from other well effluent, metered, combined with gas (single well or multi-well group batteries), and sent to a gas plant for further processing must be converted to its GEV and added to the gas production for reporting purposes.

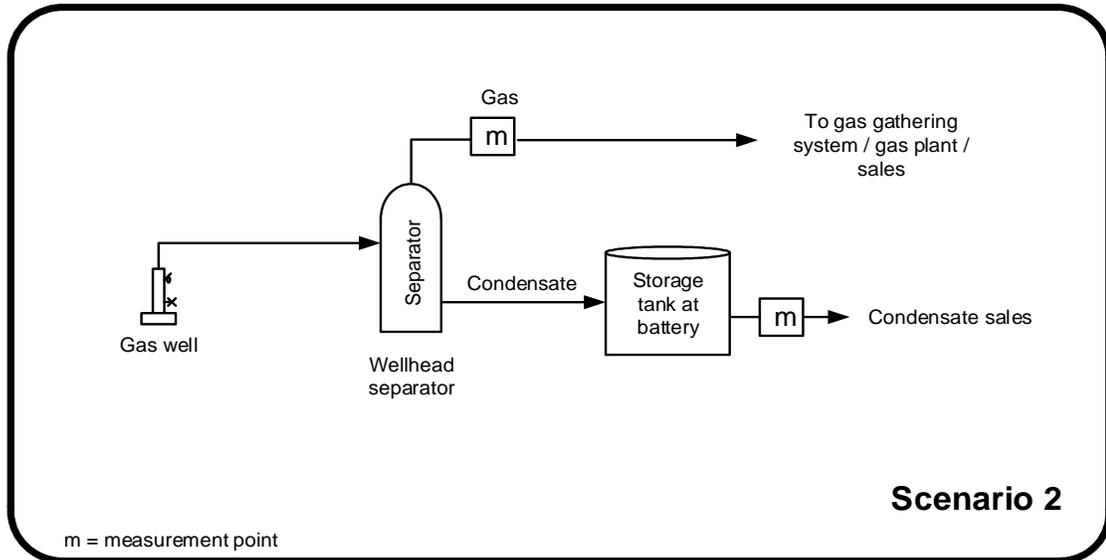
**Figure 13.1 Scenario 1**



### 13.2.2 Scenario 2 – Condensate Production Separated and Disposed of from a Well

Field condensate (COND) must be reported as liquid production at the well if separated from well effluent, measured, and disposed of.

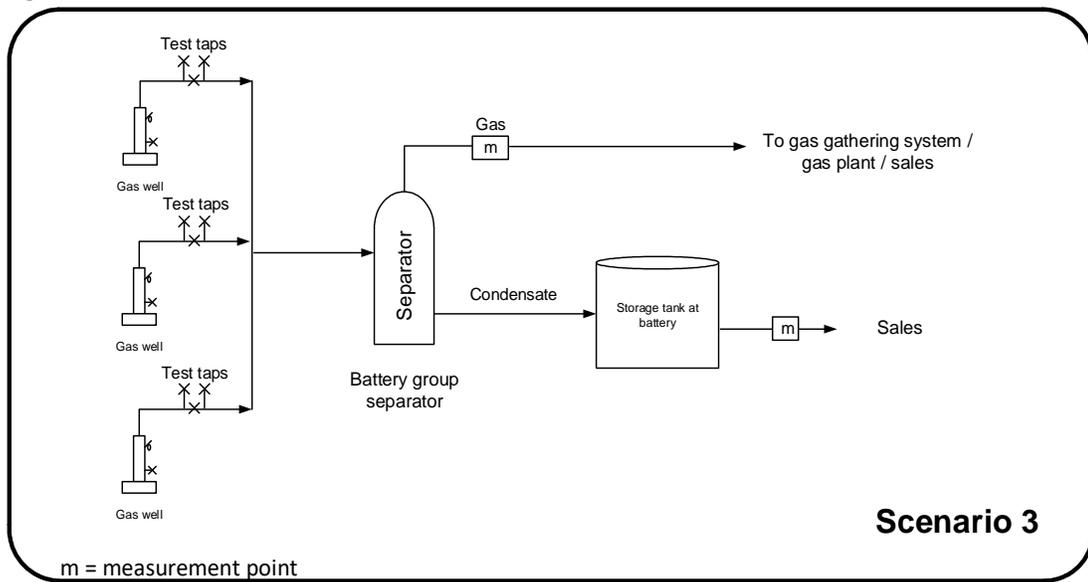
Figure 13.2 Scenario 2



### 13.2.3 Scenario 3 – Condensate Production Separated and Disposed of from a Multi-well Battery

Field condensate (COND) must be reported as a liquid production at the well level if separated from multi-well gas proration or effluent measurement batteries, measured, and disposed of from the battery.

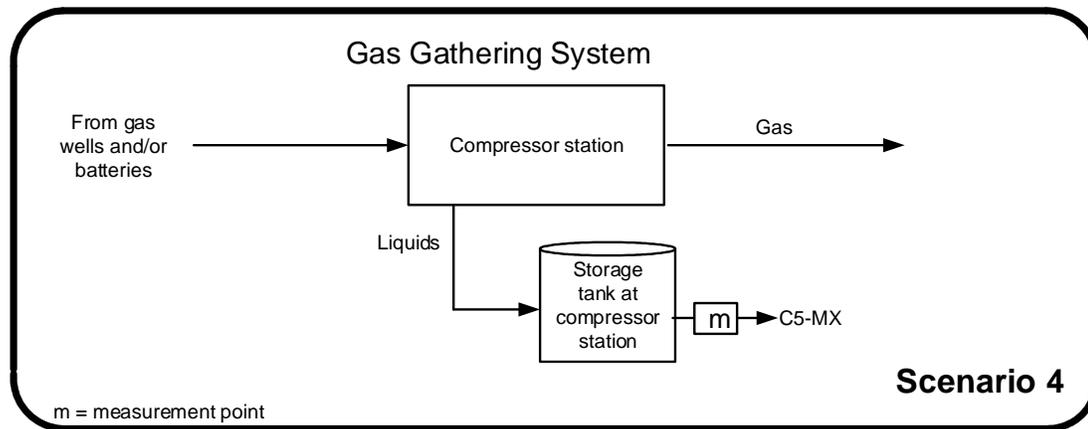
Figure 13.4 Scenario 3



13.2.4 Scenario 4 – Condensate Recovered from Gas Compression

Condensate from one or more batteries recovered as a result of gas compression at a gas gathering system or a gas group battery and disposed of must be reported as C5-MX at the gas gathering system.

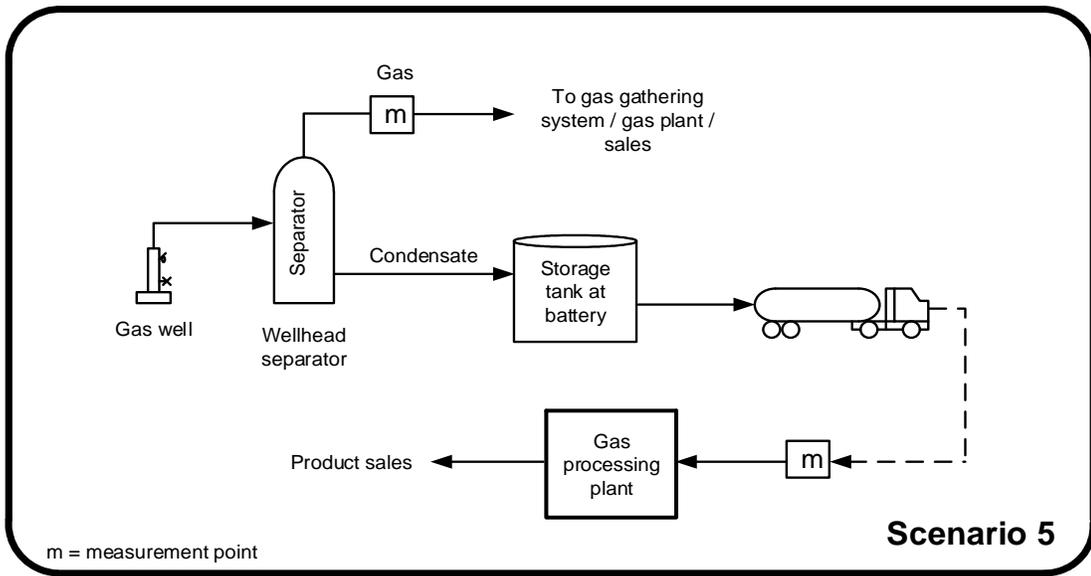
Figure 13.4 Scenario 4



13.2.4.1 Scenario 5 – Condensate Separated and Disposed of by Truck for Processing

Condensate separated from well effluent, measured, and trucked to a gas plant for processing must be reported as COND.

Figure 13.5 Scenario 5



## 14 Hydrocarbon Liquid Measurement

### 14.1 General Requirements

#### 14.1.1 Scope

This section presents the requirements for liquid hydrocarbons measurement related to:

1. crude oil
2. condensate
3. liquefied petroleum gases
  - a. propane
  - b. butane
4. dense phase hydrocarbons
  - a. ethane
  - b. NGLs
5. water

#### 14.1.2 Application of API Measurement Standards

For petroleum liquids, the API MPMS provides requirements for custody transfer measurement of hydrocarbons. For the purposes of this section, the degree of application of MPMS is determined by the level of uncertainty as required in Section 1.

#### 14.1.3 System Design and Installation

The meter system design must meet the overall system uncertainty requirements of Section 1. ER considers a liquid measurement system to be compliant if the requirements in this section are met. Any EFM system designed and installed in accordance with API MPMS, Chapter 21.2, is considered to have met the audit trail and reporting requirements, but a performance evaluation is still required in accordance with Section 14.10 of this Directive.

Liquid measurement systems typically consist of:

1. primary elements, such as a meter;
2. secondary elements, such as temperature and pressure transmitters;
3. in some cases, differential pressure transmitters, level transmitters, and densitometers
4. tertiary elements collectively termed electronic flow measurement (EFM), for example:
  - a. distributed control system (DCS)
  - b. supervisory control and data acquisition system (SCADA)
  - c. flow computers
5. in some cases, mechanical totalizers are used in place of EFM.

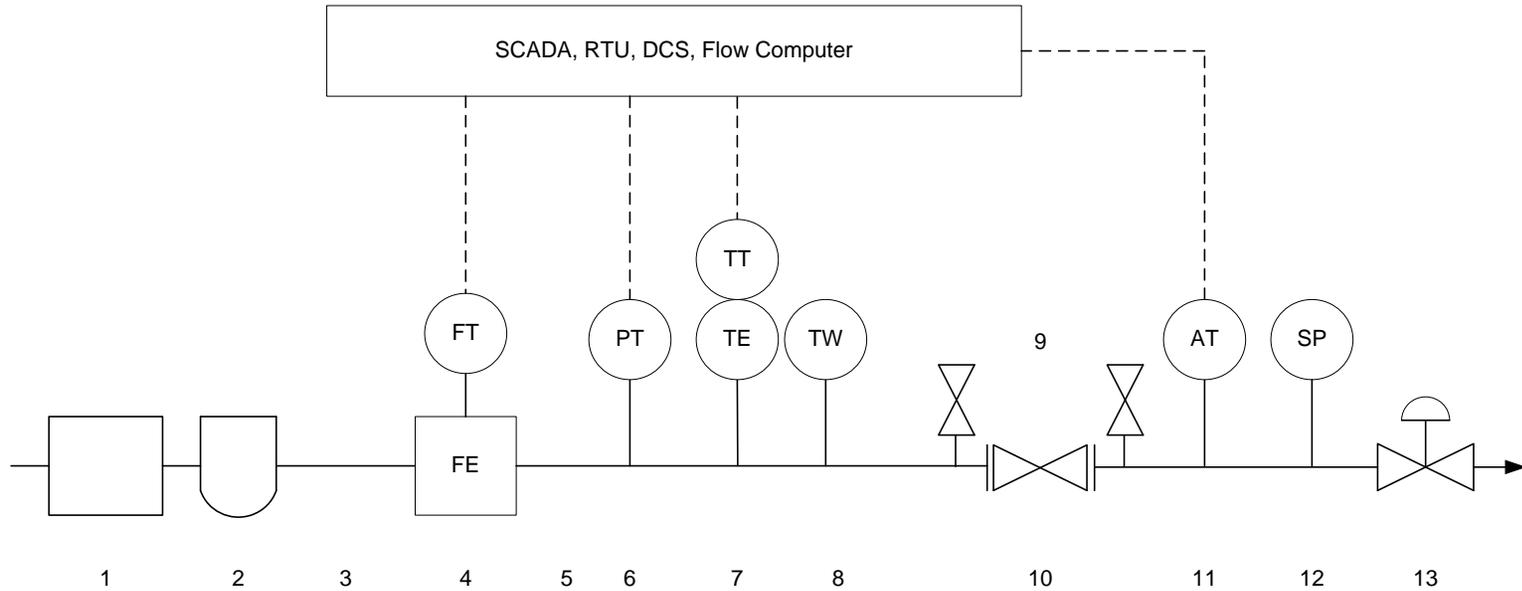
The meter and its associated peripheral equipment, such as strainers and air eliminators where installed, proving valves, and piping must be designed and installed according to applicable

procedures accepted by an appropriate industry technical standards association or the manufacturer's recommendations.

For delivery point applications where in-line proving is to be performed, proving taps and a double block and bleed divert valve must be installed. For positive displacement (PD) and Coriolis meters, proving taps may be upstream or downstream of the meter if a ball prover, pipe prover, or master meter is used. For other types of linear meters or tank provers, the proving taps must be downstream of the meter, refer to Section 2.4 for more detail.

Components of a liquid measurement system are shown in Figure [14.1](#).

Figure 14.1 Typical liquid measurement system



**Components**

- 1) Strainer
- 2) Air eliminator
- 3) Upstream straight lengths
- 4) Meter
- 5) Downstream straight lengths
- 6) Pressure transmitter (if required)
- 7) Temperature transmitter
- 8) Check thermowell
- 9) Prover valves
- 10) Double block and bleed prover divert valve
- 11) Analyzer (e.g., water cut, densitometer)
- 12) Sample point (manual or on-line)
- 13) Flow control valve

**Notes**

- 1) Schematic is generic in nature and therefore all elements may not be required for a specific application or in that order. For example, for water meters, pressure and temperature transmitters for compensation to standard conditions are not normally required.
- 2) Air eliminator is mandatory for truck unloading applications but typically not required for pipeline applications.
- 3) Strainer required for most but not all meter types.
- 4) Upstream and downstream meter straight length requirement varies with meter type and upstream piping disturbances.
- 5) Flow transmitter (FT) may be close coupled to flow sensor (FE) or remote mounted.
- 6) Analyzers are typically water-cut monitors or densitometers.
- 7) Flow control valve may be upstream of prover taps for test separator applications.
- 8) No components such as analyzer fast loops or pressure relief valves should be between meter and prover taps.
- 9) Pressure relief valves should be located to preclude unmeasured fluids via a leaky relief valve.

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## 14.2 Volume Measurement

### 14.2.1 Meter Selection

Appropriate engineering practice is required for selection of meter type and size. Specifically, parameters such as the following must be considered:

1. Process operating conditions such as pressure, temperature, flow rate;
2. Fluid properties such as viscosity, density, contaminants, bubble point;
3. Required accuracy to meet Section 1 uncertainty requirements;
4. Meter pressure drop;
5. Required straight lengths; and
6. Required back pressure.

Parameters known to vary with operating conditions, such as fluid properties such as viscosity and flow rate, must be considered for all operating scenarios like start-up, normal, and upset.

If meters are used for delivery point measurements, electronic temperature compensation is required. For new construction or new installations at existing facilities, mechanical ATC and ATG must not be used. For facilities built before April 1, 2016, all existing ATC and ATG are grandfathered at their existing installations and must not be relocated and reused for other installations.

For meters to be proved using a conventional displacement prover, such as a ball prover or a captive displacer prover such as piston and shaft, pulse outputs are required. For master meter proving, pulse outputs are recommended.

In addition to the meter selection parameters listed above in this section, some upstream applications such as propane sales loading rack at gas plants may also have to meet Measurement Canada requirements.

There are two broad meter types, linear and nonlinear differential producer. The output of linear meters is proportional to flow rate. The output of differential producers is proportional to the flow rate squared. Table 14.1 lists various meter types for volume determination.

**Table 14.1 Meter types**

Linear meters	Nonlinear meters
Positive displacement	Orifice (see Section 4.3.1)
Turbine	Venturi
Vortex	Flow nozzle
Coriolis	Cone
Ultrasonic	Wedge
Magnetic (water or conductive fluids only)	Other differential devices

## 14.3 Shrinkage

For the purpose of this Directive, shrinkage refers to a volume reduction associated with one or more of the following processes:

1. Blending of hydrocarbon streams of varying density such as heavy oil and condensate to reduce the viscosity of the heavy oil for transport by pipeline; and/or

2. Loss of volatile components through vapourization such as flashing, weathering due to a pressure reduction and/or temperature increase or to continued exposure to atmospheric conditions like the conversion of live oil to base conditions.
3. Acid Gas removal, see Section 11

Petrinex-reported shrinkages other than Sections 11 and 14.3 and shrinkage or system loss/gains across facilities or pipeline system are outside the scope of this Directive.

### 14.3.1 Live Oil Shrinkage

Until produced hydrocarbon fluids are stabilized, the oil is normally at its bubble point, also referred to as the equilibrium vapour pressure, condition due to the presence of volatile components. When the oil is discharged to a stock tank at atmospheric condition, the volatile components in the oil evaporate, causing a reduction in liquid volume. When live oils are metered in such equipment as test separators, a shrinkage factor must be applied to correct the measured liquid volume from the metering pressure and temperature to standard conditions. When the meter is proved to standard conditions, the shrinkage factor is incorporated into the meter factor.

### 14.3.2 Hydrocarbon Blending and Flashing Shrinkages

When hydrocarbon molecules of different molecular sizes and intermolecular spacing, also referred to as the density, are mixed, the smaller molecules fill the spaces between the larger molecules. This results in a volume reduction from the arithmetic sum of the volumes of the blend components. The magnitude of this volume reduction is a function of the relative density and volumes of the hydrocarbon blend components. Calculation of shrinkage factors resulting from hydrocarbon blending without flashing must be performed in accordance with API MPMS, Chapter 12.3, or an equivalent procedure accepted by an appropriate industry technical standards association.

In some cases, volume reduction is a combination of the effects of loss of volatile components and intermolecular spacing. For example, blending of condensate or diluent with heavy oil can occur at any point in the production process. The condensate can be introduced in the flow line from the well, at the inlet separator, at the treater, at the storage tank, or at any combination of these ways of introducing the condensate. If condensate is blended with the oil prior to the treater, condensate flashing may also occur.

Blending shrinkage must be determined if the density difference between the hydrocarbon fluids exceeds  $40.0 \text{ kg/m}^3$  and must be reported if the shrinkage volume causes the delivery point volume to shrink by more than 0.1 per cent and more than the  $0.1 \text{ m}^3$  reporting limit on Petrinex. Flashing shrinkage must be determined if the added diluent volume is  $> 2.0 \text{ m}^3/\text{day}$  and/or  $> 5.0$  per cent of total oil production, refer to [Table 5.6](#) for details.

The blending and flashing shrinkage is to be reported as an activity of SHR (which is a type disposition) at the facility for the product type of the diluent volume. The flashing shrinkage is to be convert to a GEV and reported as a REC for the product type of GAS from SK GE.

### 14.3.3 Shrinkage Factor Determination

Live oil shrinkage with entrained gas must be determined by any one of the following techniques:

1. Process simulation software;
2. Manual sampling and laboratory procedure, as referred to in API MPMS, Chapter 20; or
3. Physically de-gassing the prover oil volumes during meter proving of live oils, as referred to in Section 2.5.1.

Calculation of shrinkage volumes or factors is most often used to mitigate safety and environmental concerns if the live oil volumes are measured at high pressures or if the live oil contains H<sub>2</sub>S.

When the manual sampling and laboratory method is used, the shrinkage factor must be based on analysis of a sample of the fluid taken at normal operating conditions. Shrinkage factors must be determined at either a well or battery level. The frequency of shrinkage factor determination must reflect changes in reservoir or operating conditions. Whenever the operating conditions change to a degree that could affect the shrinkage factor, a new shrinkage factor must be determined based upon analysis of a sample of the fluid taken at the new operating conditions.

#### 14.3.4 Shrinkage Factor Application

Shrinkage factors must be applied by being:

1. Incorporated into a meter factor by de-gassing during proving;
2. Incorporated into a meter factor by adjusting the meter factor numerically based on a shrinkage factor determined by process simulation or sampling/analysis; or
3. Applied to metered volumes after they are adjusted by the meter factor (which does not already have the shrinkage factor incorporated into it).

Caution is required to ensure that shrinkage is not applied more than once, such as de-gassing during meter proving and then applying it again as a factor to measured volumes.

#### 14.4 Temperature Measurement

Temperature effects can increase the uncertainty associated with liquid hydrocarbon and water measurements. The magnitude of the effect of temperature measurement errors increases with decreasing hydrocarbon density as illustrated in Table 14.2.

**Table 14.2 Temperature measurement error impact**

Fluid	Approximate error per 1°C temperature measurement error (%)
Propane (510.0 kg/m <sup>3</sup> @15.0°C)	0.29
Butane (600.0 kg/m <sup>3</sup> @15.0°C)	0.18
Condensate (700.0 kg/m <sup>3</sup> @15.0°C)	0.12
Crude oil (820.0 kg/m <sup>3</sup> @15.0°C)	0.09
Crude oil (920.0 kg/m <sup>3</sup> @15.0°C)	0.07
Water	0.02

Therefore, temperature compensation of measured volumes must be provided as required to meet the uncertainty requirements detailed in Section 1 and the requirements of this section.

This applies to delivery point measurement, provers, and others such as LACT's that require temperature compensation for volumetric determination.

Thermowells or direct insertion temperature elements must be used for all temperature measurements. Pipe or meter body skin temperature measurements, such as those used by coriolis meter, are not acceptable unless proven to be within the uncertainty requirements.

Thermowells must be installed in such a manner to be representative of the fluid temperature. Thermowells must not be installed in sections of piping where flow may not be present, for example in dead-ended piping or in a storage tank above the normal liquid level.

With the exception of coriolis or PD meters, thermowells must be installed 5.0 to 10.0 pipe diameters downstream of the meter. For coriolis or PD meters, thermowells must be installed within 10 pipe diameters upstream or downstream of the meter. Valves or pipe restrictions must not be present between the thermowell and the meter's primary element. Meter runs designed for trucked liquid measurement with the existing thermowell(s) within 20.0 diameters of the meter are grandfathered for the existing location and usage. If the meter run is modified or relocated, then these requirements must be met.

Resistance temperature devices (RTD) are the preferred temperature measurement element. Other types of temperature measurement elements, such as thermocouples and thermistors, are acceptable provided that uncertainty requirements are met. Dial thermometers are not acceptable for pipeline-based delivery point measurement.

For pipeline delivery point measurements, two thermowells should be provided, one for measurement and one for verification. For new construction or new installations at existing facilities, mechanical ATC and ATG must not be used. For facilities built before April 1, 2016, all existing ATC and ATG are grandfathered at their existing installations and must not be relocated and reused for other installations

Temperature measurement type, tolerances, and calibration frequency are detailed in [Table 14.3](#).

**Table 14.3 Temperature measurement type, calibration frequency, resolution, and calibration tolerances**

Application	Temperature measurement type <sup>1</sup>	Minimum resolution (°C)	Maximum calibration tolerance (°C)	Verification frequency
Delivery point with meter	Continuous with EFM	0.1	±0.5	Monthly <sup>2</sup>
Well oil (proration battery)	Composite meter factor or continuous with EFM	0.5	±1.0	Annual
Plant inlet or total battery/group condensate (gas gathering system)	Continuous or composite meter factor (see Section 14.9)	0.5	±1.0	Semiannual
Delivery point batch volumes into a pipeline or receipt at a battery/facility using tank gauging	One reading per load	0.1	±0.5	Semiannual

<sup>1</sup> For mechanical ATCs, see Section 10.3.1.

<sup>2</sup> Calibration frequency may be changed to bimonthly if three consecutive verification periods pass without the error exceeding the tolerance.

## 14.5 Pressure Measurement

Pressure compensation of hydrocarbon liquids is required where the meter pressure is above the standard pressure for delivery point measurement unless the meter is proved to standard conditions. The pressure correction referred to as Correction for the effect of Pressure on Liquids (CPL) factor must be determined in accordance with API MPMS, Chapter 11.

Continuous pressure measurements and pressure compensation must be installed where required to meet Section 1 uncertainty requirements.

Pressure transmitters and gauges must be installed in accordance with applicable standards of an appropriate industry technical standards association or manufacturer's recommendations, normally 5.0 to 10.0 pipe diameters downstream of the meter.

## 14.6 Density Determination

Density may be measured manually from a sample or continuously using either a densitometer or a Coriolis meter. Where manual density is used, the manual density value may be derived from a representative grab or composite sample and a laboratory density determination. Whichever method is used, the derivation of the value must be documented and meet the uncertainty requirement.

Continuous density measurements must be provided for mass measurement or if the variability in density is such that use of a fixed density value for temperature compensation would preclude meeting the uncertainty requirements.

On-line densitometers must be installed in accordance with applicable standards of an appropriate industry technical standards association or manufacturer's recommendations, normally 5.0 to 10.0 pipe diameters downstream of linear meters. If a densitometer is used as part of a mass measurement system, for example ethane, NGLs, it must be installed in accordance with API MPMS, Chapter 14.6.

Laboratory density determination may be performed using either the hydrometer methods see API MPMS, Chapter 9 or the precision densitometer method ASTM D4052 for more detail. If practical, densitometer measurements should be made at 15.0°C to preclude the requirement for temperature compensation. If this is not practical for such applications as viscous heavy oil or when using a hydrometer, manual temperature compensation must be provided using the applicable API MPMS table. Refer to Section 14.9 for more detail.

All analyses are required to be submitted to ER as per *Directive PNG013*.

## 14.7 Tank Measurement

Tanks in this section refer to storage tanks that are open to atmosphere, tanks with and without floating roofs, and tanks with blanket gas, as well as bullets and other pressurized storage vessels. The use of tanks open to atmosphere should be limited to liquids with a Reid Vapour Pressure specification of < 103.0 kPa.

Volumetric measurement using storage tanks is based upon a level measurement used in conjunction with a strapping table.

Provided that Section 1 uncertainty tolerances are met, the licensee may use storage tanks for determination of inventory, well test, or delivery point volume measurements. The licensee must ensure that the tank diameter, gauging equipment such as gauge tape or automatic tank

gauge, gauging procedures, and tank strapping table are appropriate for the tank and product being gauged and are capable of achieving the required uncertainty.

Manual gauge boards and automatic tank gauges must be designed, installed, and operated in accordance with manufacturer's specifications and recommendations and must be maintained in good working order.

### 14.7.1 Tank Strapping

Tank strapping tables convert or relate the level to indicated volume.

Depending upon the uncertainty required, tank strapping tables may be prepared using either engineering calculations based upon approximate tank dimensions or via the tank strapping procedures detailed in API MPMS, Chapter 2.

### 14.7.2 Tank Sizing

The relative error of the level measurement is determined by the absolute error of the level measurement relative to the level measured. The level measured or change in level is in turn determined by the diameter of the tank and transaction size. To improve uncertainty, one can measure the level more accurately or increase the level change measured by changing the ratio of tank height to diameter or by increasing the size of the transaction (delivery point) or test volume compared to the overall tank height.

Tank sizing must address the intended use such as delivery point or well test, level measurement technique such as gauge board, hand dip, radar gauge, and well test or transaction volume.

Knowing the transaction or test volume, one can determine tank diameter as follows:

$$d \leq (V \div a)^{1/2}$$

Knowing the tank diameter, one can determine minimum transaction or test volume as follows:

$$V \geq a \times d^2$$

Where:

V = test fluid volume or delivery point batch volume in m<sup>3</sup>

d = tank diameter in metres

a = accuracy coefficient

The accuracy coefficients for non-heavy crude oil applications are:

a = 0.39 for all test fluid volumes

a = 0.39 for delivery point batch volumes ≤ 100.0 m<sup>3</sup>/day

a = 0.92 for delivery point batch volumes > 100.0 m<sup>3</sup>/day

The accuracy coefficients for heavy crude oil applications are:

a = 0.39 for treatment facility receipt batch volumes

a = 0.92 for sales/delivery point batch volume

a = 0.39 for primary heavy oil test fluid volumes

### 14.7.3 Manual Tank Gauging

Manual tank gauging can be accomplished using tank dips or a gauge board.

Gauge boards are acceptable for test tanks and inventory measurements but not for delivery point measurements. See Table 14.4 for marking gradations.

Gauge tapes must have a minimum resolution of 3.0 mm.

**Table 14.4 Gauge board marking gradations**

Gauge board application	Maximum marking separation (mm)
Non-heavy oil testing	25.0
Heavy crude oil testing	60.0
Inventory	150.0

If safe work conditions permit, gauge boards are best read at eye level.

Calibration of gauge boards is not required.

On an annual basis, the licensee must ensure the gauge board is in good working order and the strapping tables are appropriate for the type of measurement and fluid as per API MPMS, Chapter 2.

### 14.7.4 Automatic/Electronic Tank Gauging

Electronic tank gauges must have a minimum resolution of 3.0 mm. One reading of the instrument is acceptable.

Instruments must be calibrated in accordance with the manufacturer's recommendation. See Section 2.10 for frequency requirement.

### 14.7.5 Tank Gauging Applications

#### 14.7.5.1 Inventory Tank Gauging

For monthly inventory measurement gauging, one reading of the gauge tape, gauge board, or automatic tank gauge is acceptable. Levels must be reported to the nearest 75.0 mm.

The tank does not need to be stabilized or isolated for inventory measurements.

#### 14.7.5.2 Test Tank Gauging

For gauge measurement on test tanks, two readings of the gauge board or automatic tank gauge is required, one at the start of the test and one at the end of the test.

Levels are reported to the nearest 10 mm.

#### 14.7.5.3 Delivery Point Measurement

When tank gauging is used to determine an oil/emulsion volume, the gauging procedures must be conducted in accordance with the following:

1. The licensee must ensure that the strapping table has been prepared in accordance with API MPMS, Chapter 2.

2. The licensee must ensure that the tank level is not changing or is stabilized when the gauge readings are taken. This often requires isolating or shutting in the tank before gauging.
3. All gauge tapes and electronic level devices must have a minimum resolution of 3 mm.
4. Manual tank dips are performed in accordance with API MPMS, Chapter 3.1A. For tanks with a nominal capacity > 160.0 m<sup>3</sup>, two consecutive readings within 10.0 mm of each other are required at both the beginning and at the end of the volume transaction. The two readings are averaged. For tanks with a nominal capacity of ≤ 160.0 m<sup>3</sup>, one reading at the beginning and at the end of the volume transaction is acceptable.
5. Automatic tank gauging is performed in accordance with API MPMS, Chapter 3.1B.
6. Temperature measurements are performed in accordance with API MPMS, Chapter 7.
7. Gauge boards must not be used for delivery point measurement.

## **14.8 Sampling and Analysis for S&W and Density Determination**

Sampling and analysis must be in accordance with Sections 6, 8, and 10 or other equivalent method approved by an appropriate industry standards association.

### **14.8.1 Fluid Sampling Requirements for S&W and Density Determination**

S&W determination procedures including the frequency of sampling must be representative of the entire volume transaction as well as the subsequent S&W sample analysis. There are two methods to obtain this measurement: sampling or on-line analysis using a suitable instrument such as water-cut analyzer or product analyzer. Sampling can be categorized by two methods: spot/grab sampling or continuous proportional sampling. It is important that the sample location be carefully selected such that the flowing stream is adequately mixed. This can be achieved by:

1. Installing in-line mixers;
2. Selecting a sampling point that offers the most practical location for collecting a sample that is mixed, such as after valves, elbows, and reducers;
3. Selecting a sampling point that is downstream of a metering point because of the piping elements associated with a meter run; or
4. Collecting samples from a number of different locations, analyzing them, and making a selection based on the location that provides the most consistent and reasonable analysis.

For proration oil testing, grab or spot sampling may be used if the water cut < 10.0 per cent otherwise continuous proportional sampling or the use of a product analyzer is required.

Water-cut analyzers operate on a number of different principles and often are best suited for specific applications. Analyzers must be installed and maintained in accordance with the manufacturer's recommendations.

For a single-well battery or a multi-well group battery trucking emulsion off-site, the volumes are determined at the receiving facilities.

For single-well oil batteries with two-phase or three-phase separators delivering produced oil/emulsions by pipeline to another battery, the sample must be taken at or near the oil/emulsion meter using a continuous proportional sampler. An on-line product analyzer is also acceptable for the determination of water cut. Refer to Section 15.2.2.1 for the exception to these sample taking methods. This is a measurement-by-difference situation at the receiving battery/facility and the oil/emulsion meter at the single well battery must be proved to standard conditions. For more detail on measurement-by-difference refer to Section 5.5.

For an oil battery with emulsion tanks, the oil and water inventory volumes in the emulsion tanks may be determined by one of the following methods:

1. Taking a spot sample, also referred to as a grab sample anywhere between the wellhead or separator and the tank and applying the percentage of sediment and water (percent S&W) to the tank inventory;
2. Using water-indicating paste on the gauge tape to determine the water/oil interface in the tank inventory;
3. Taking the average percent S&W of the total battery production and applying that to the tank inventory;
4. Using the average percent S&W of the trucked out volumes; or
5. Deeming the tank inventory to be entirely oil and making changes/amendments based on delivery volumes.

#### **14.8.2 S&W Determination**

The licensee must select the most appropriate method for determining the percent S&W. There are three static analysis methods of the sampled fluid generally considered acceptable by ER based on the percent S&W:

1. The centrifuge or Karl Fischer method combined with separate method for sediment determination for water cuts  $> 0.0$  per cent and  $\leq 10.0$  per cent;
2. The graduated cylinder method of a larger sample for water cuts  $> 10.0$  per cent and  $\leq 80.0$  per cent and centrifuging the oil emulsion portion; and
3. The graduated cylinder method of a larger sample for water cuts  $> 80.0$  per cent and  $\leq 100.0$  per cent and not centrifuging the oil emulsion portion.

Recommended procedures for these three methods are shown in Appendix 3. Any alternative methods must be supported by testing that shows representative results are achieved and these alternative procedures must be made available to ER upon request.

In some instances, it is possible to use a computer algorithm to determine the oil and water volumes in the emulsion based on the measured densities of the emulsion and the known densities of the oil and water components of the emulsion. The oil and water densities at standard conditions must be based on an analysis of the actual oil and water production being measured and must be corrected for the temperature at which the emulsion density is measured. Temperature correction for produced water density should be calculated in accordance with API MPMS, Chapter 20.1 or some other recognized industry standards organization.

## 14.9 Hydrocarbon Liquid Volume Calculations

Liquid volume measurements must be determined to a minimum of two decimal places and rounded to one decimal place for monthly Petrinex reporting in cubic metres. If there is more than one volume determination within the month at a measurement point, the volumes determined to a minimum of two decimal places must be totaled prior to the total being rounded to one decimal place for Petrinex reporting purposes.

Standard or base conditions for use in calculating and reporting liquid volumes are 15.0°C and 0.0 kPa gauge or the equilibrium vapour pressure at 15.0°C, whichever is higher.

The liquid volume calculations must adhere to the following:

1. Total indicated volume for the transaction period, either daily, weekly, or monthly is measured and recorded. This applies to measurement by meter, weigh scale, or tank gauging.
2. The volumetric meter factor for the flow meter is applied to the total indicated volume.
3. For oil, the percentage of water in the gross volume is determined by measuring the percent S&W of a representative sample or by continuous on-line measurement. The result is a quantified volume of oil and of water.
4. For oil, a shrinkage factor is applied to the volume in order to determine the volume at standard conditions. Some applications may already have the shrinkage factor incorporated into the meter factor.
5. Where required, compensation for the effects of pressure and temperature on the liquid must be applied.
6. Composite meter factors that include temperature correction factors (CTL) must not be used for delivery point measurement. However, they are acceptable for other applications, such as test meters, inlet meters, and water meters, provided that the variability of parameters affecting meter performance such as operating temperature, fluid viscosity, and fluid composition is such that the net effect is within the uncertainty requirements for the application.

### 14.9.1 General Equations for Determining Liquid Volumes at Standard Conditions

#### 14.9.1.1 Linear Meters

##### Indicated Volume

$$IV = \text{closing reading} - \text{opening reading}$$

**or**

$$IV = (\text{closing pulses} - \text{opening pulses}) \div KF$$

##### Gross Standard Volume

$$GSV = IV \times CTL \times CPL \times MF$$

**or**

$$GSV = IV \times CMF$$

**or**

$$GSV = IV \times MF \times DEN_{obs} \div DEN_b$$

or

$$\text{GSV} = \text{Mass} \div \text{DEN}_b$$

### Net Standard Volume

$$\text{CSW} = 1 - (\% \text{S\&W} \div 100)$$

$$\text{NSV} = \text{GSV} \times \text{CSW} \times \text{SF}$$

### Water Cut

$$\text{DEN}_{\text{obs,o}} = \text{DEN}_{b,o} \times \text{CTL}_o$$

$$\text{DEN}_{\text{obs,w}} = \text{DEN}_{b,w} \times \text{CTL}_w$$

$$\text{Water Cut} = (\text{DEN}_{\text{obs,e}} - \text{DEN}_{\text{obs,o}}) \div (\text{DEN}_{\text{obs,w}} - \text{DEN}_{\text{obs,o}})$$

Where:

**CMF – Composite Meter Factor:** A meter factor that includes corrections for the effects of any combination of temperature, pressure, or shrinkage.

**CPL – Correction for the effect of Pressure on Liquid:** Correction for compressibility of liquid at normal operating conditions.

**CTL – Correction for the effect of Temperature on Liquid:** Correction for effect of temperature on liquid at normal operating conditions.

**CTL<sub>o</sub> – Correction for the effect of Temperature on Oil:** Correction for effect of temperature on oil at normal operating conditions.

**CTL<sub>w</sub> – Correction for the effect of Temperature on Water:** Correction for effect of temperature on water at normal operating conditions.

**CSW – Correction for Sediment and Water:** Correction for sediment and water to adjust the gross standard volume of the liquid..

**DEN<sub>b</sub> – Base Density:** Liquid density in kg/m<sup>3</sup> at standard pressure and temperature.

**DEN<sub>b,o</sub> – Base Density – Oil:** Liquid density of oil in kg/m<sup>3</sup> at standard pressure and temperature.

**DEN<sub>b,w</sub> – Base Density – Water:** Liquid density of water in kg/m<sup>3</sup> at standard pressure and temperature.

**DEN<sub>obs</sub> – Observed Density:** Liquid density in kg/m<sup>3</sup> at observed pressure and temperature.

**DEN<sub>obs,o</sub> – Observed Density – Oil:** Oil density in kg/m<sup>3</sup> at observed pressure and temperature.

**DEN<sub>obs,w</sub> – Observed Density – Water:** Water density in kg/m<sup>3</sup> at observed pressure and temperature.

**GSV – Gross Standard Volume:** The volume at standard conditions corrected also for the meter's performance (MF or CMF).

**IV – Indicated Volume:** The change in meter reading that occurs during a receipt or delivery.

**KF – K-Factor:** A term in pulses per unit volume determined during a factory or field proving. The number of pulses generated by a linear meter divided by the k-factor will determine the indicated volume.

**MF – Meter Factor:** A dimensionless term obtained by dividing the volume of a liquid passed through a prover (corrected to standard conditions during proving) by the indicated standard volume (ISVm) as registered by the meter.

**SF - Shrinkage Factor:** a factor reflecting the volume reduction of a liquid associated with one or both of the following two processes:

1. Blending of hydrocarbon streams of varying density such as heavy oil and condensate to reduce the viscosity of the heavy oil for transport by pipeline; and or
2. Loss of volatile components through vapourization such as flashing or weathering due to a pressure reduction and/or temperature increase or to continued exposure to atmospheric conditions such as conversion of live oil to standard conditions.

**NSV – Net Standard Volume:** The gross standard volume corrected for shrinkage and sediment and water.

#### 14.9.1.2 Composite Meter Factors (CMF)

A composite meter factor (CMF) is a meter factor that includes corrections for the effects of any combination of temperature, pressure, or shrinkage.

A CMF may be used:

1. If anticipated changes in pressure and temperature parameters result in uncertainties within those stated in Section 1;
2. For test separators at oil batteries; and
3. For separators at gas wells.

Test separators typically use CMFs to apply temperature correction where an EFM system is not used. The CMF can also include correction for shrinkage. The licensee must ensure that corrections included in CMFs are not being applied elsewhere, such as in a SCADA system or field data capture system.

Note that in separator applications where the hydrocarbon liquid is at its equilibrium vapour pressure, CPL is 1.0 and therefore is not required to be calculated as part of a CMF.

Calculation example for volumetric proving at an oil test separator:

$$CMF_T = IV_P \times CTL_P \div IV_M$$

Where:

$CMF_T$  - CMF that includes correction for the effect of temperature (CTL)

$IV_P$  - Indicated prover volume

$CTL_P$  - CTL calculated using prover temperature during run

$IV_M$  - Indicated meter volume

If the indicated volume of the prover is recorded after de-gassing, the CMF will include correction for shrinkage ( $CMF_{TS}$ ).

#### 14.9.1.3 Orifice Meters

While not as common, orifice meters can be used for liquid measurement. For these applications, either of the following equations must be used.

API MPMS 14.3.1 (AGA-3):

$$Q_b = \frac{Q_m}{\rho_b} = \frac{N_1 C_d E_v Y d^2 \sqrt{\rho_f \Delta P}}{\rho_b}$$

API MPMS 14.8 (Natural Gas Fluids Measurement – Liquefied Petroleum Gas Measurement):

$$Q_b = N_1 C_d E_v Y d^2 \sqrt{\frac{\Delta P}{\rho_f}} (C_{tl} C_{pl})$$

Where:

$N_1$  - Unit conversion factor (0.0000351241 when using SI units listed below)

$C_d$  - Orifice plate coefficient of discharge

$E_v$  - Velocity of approach factor

$Y$  - Expansion factor

$d$  - Orifice plate bore diameter calculated at flowing temperature (mm)

$\Delta P$  - Orifice differential pressure (kPa)

$\rho_f$  - Density of the liquid at flowing conditions (kg/m<sup>3</sup>)

$\rho_b$  - Density of the liquid at standard conditions (kg/m<sup>3</sup>)

$Q_b$  - Volume flow rate at standard conditions (m<sup>3</sup>/sec)

$Q_m$  - Mass (kg)

$C_{tl}$  - Compensation factor for the effect of temperature on liquid

$C_{pl}$  - Compensation factor for the effect of pressure on liquid

For other nonlinear meters, refer to the applicable standard of an appropriate industry technical standards association or manufacturer's documentation for determining volumes at standard conditions.

#### 14.9.1.4 Pressure and Temperature Compensation

CTL and CPL must be calculated in accordance with the current standards in [Table 14.5](#) for the applicable density and temperature range. Calculations for determining CTL or CPL not listed in [Table 14.5](#) are not acceptable.

**Table 14.5 Pressure and temperature compensation standards\***

Standard	Product and density range	Calculation input(s)	Calculation output(s)	Comments
API MPMS 11.1 May 2004	Crude oil, refined products, and lubricating oils 611.16–1163.85 kg/m <sup>3</sup>	Observed density Density @ 15.0°C Flowing temperature Flowing pressure Equilibrium vapour pressure	Density @ 15.0°C CTL CPL VCF	Current
API MPMS 11.2.2M 1986	Hydrocarbon liquid 350.0–637.0 kg/m <sup>3</sup>	Density @ 15.0°C Flowing temperature Flowing pressure Equilibrium vapour pressure	CPL	Current
API MPMS 11.2.4 GPA TP-27 Table 53E September 2007	NGL and LPG 210.0–740.0 kg/m <sup>3</sup>	Observed density Observed temperature	Density @ 15.0°C	Current
API MPMS 11.2.4 GPA TP-27 Table 54E September 2007	NGL and LPG 351.7–687.8 kg/m <sup>3</sup>	Density @ 15.0°C Flowing temperature	CTL	Current

\* Note: The printed API MPMS, Chapter 11.1, Tables 53, 53A, and 53B include correction for the thermal expansion or contraction of a glass hydrometer. Existing computer implementations of these tables may or may not include hydrometer correction.

### 14.9.2 Electronic Flow Measurement for Liquid Systems

An EFM system is any flow measurement and related system that collects data and performs flow calculations electronically. If it is part of a DCS, SCADA, or Programmable Logic Controller system (PLC), only the EFM portion of those systems has to meet the requirements in this section.

The following systems are not defined as an EFM:

1. Any meter with an electronic totalizer or pulse counter that does not perform flow calculations with or without built-in temperature compensation; and
2. A remote terminal unit (RTU) that transmits any data other than flow data and does not calculate flow.

Hardware and software requirements:

1. The EFM system data storage capability must exceed the time period used for data transfer from the EFM system.

2. The EFM system must be provided with the capability to retain data in the event of a power failure. Data retention options include battery backup, uninterrupted power systems, or EPROM.
3. The system must have appropriate levels of access for security, with the highest level of access to the system restricted to authorized personnel.
4. The EFM system must be set to alarm on out-of-range inputs, such as temperature, pressure, differential pressure if it is applicable, flow, low power, or communication failures.
5. Any EFM system configuration changes or forced inputs that affect measurement computations must be documented through either electronic audit trails or paper records.
6. The values calculated from forced data must be identified as such.

### 14.9.3 Performance Evaluation

If an EFM system is used to calculate net liquid volumes, the licensee must be able to verify that it is performing within ER target limits defined in this section.

A performance evaluation test must be completed within two weeks after the EFM system is put into service and immediately after any change to the computer algorithms that affects the flow calculation on a per software version basis, and it must be documented for ER audit upon request. For existing EFM systems, the licensee must conduct a performance evaluation to ensure that they are performing adequately. A performance evaluation must be conducted and submitted for ER audit on request. ER considers either one of the following methods acceptable for performance evaluation.

1. A performance evaluation test conducted on the system by inputting known values of flow parameters into the EFM system to verify the volume calculation and other parameters. The test cases included in this section (Tables 14.6 to 14.9) are for liquid meters each with different flow conditions.

Test cases 1 to 5 for each liquid type are for density correction from flowing temperature to 15.0°C. The hydrometer correction is used to compensate for the glass expansion when a hydrometer is used to measure the density.

Test cases 6 to 10 for each liquid type are for volume correction using CPL and/or CTL factors to correct to standard conditions. Other manufacturer's recommended methodologies can also be used to evaluate the EFM performance, provided that the volumes obtained from a performance evaluation test agree to within  $\pm 0.1$  per cent of those recorded on the sample test cases.

2. Evaluation of the EFM system calculation accuracy with a flow calculation checking program that performs within the target limits for all the factors and parameters listed in the test cases specified in Section 14.9.4. A snapshot of the instantaneous flow parameters and factors, flow rates, and configuration information is to be taken from the EFM system and input into the checking program. If the instantaneous EFM system flow parameters, factors, and flow rates are not updated simultaneously, multiple snapshots may have to be taken to provide a representative evaluation.

The densities in test cases 1 to 5 and 11 to 15, or volumes in test cases 6 to 10 and 16 to 20, obtained from a performance evaluation test must agree to within  $\pm 0.1$  per cent of those recorded on the sample test cases. If the  $\pm 0.1$  per cent limit is exceeded, the EFM system must be subjected to a detailed review of the calculation algorithm to resolve the deviation problem.

#### 14.9.4 Test Cases for Verification of Oil Flow Calculation Programs

These test cases were calculated using the following standards.

**Density @ 15°C CTL/CPL/CTPL:** API MPMS, Chapter 11.1: Temperature and Pressure Volume Correction Factors for Generalized Crude Oils, Refined Products, and Lubricating Oils (May 2004).

**Hydrometer Correction:** API MPMS, Chapter 9.3: Standard Test Method for Density, Relative Density, and API Gravity of Crude Petroleum and Liquid Petroleum Products by Thermohydrometer Method (November 2002).

#### 14.9.5 Test Cases for Verification of NGL and LPG Flow Calculation Programs

These test cases were calculated using the following standards.

**Density @ 15°C:** API MPMS, Chapter 11.2.4 (GPA Technical Publication TP-27): Temperature Correction for the Volume of NGL and LPG, September 2007, Table 53E.

**Hydrometer Correction:** API MPMS, Chapter 9.3: Standard Test Method for Density, Relative Density, and API Gravity of Crude Petroleum and Liquid Petroleum Products by Thermohydrometer Method (November 2002).

**CPL:** API MPMS, Chapter 11.2.2M Compressibility Factors for Hydrocarbons, October 1986.

**CTL:** API MPMS, Chapter 11.2.4 (GPA Technical Publication TP-27): Temperature Correction for the Volume of NGL and LPG, September 2007, Table 54E.

**Table 14.6 Oil density correction test cases—density correction to 15°C**

Test case	Inputs		Outputs	
	Oil density @ observed temp. (kg/m <sup>3</sup> )	Observed temp. (°C)	Oil density corrected to 15°C (kg/m <sup>3</sup> ) with hydrometer correction	Oil density corrected to 15°C (kg/m <sup>3</sup> ) without hydrometer correction
1	875.5	120.0	942.9	945.0
2	693.0	11.4	689.9	689.8
3	644.0	84.45	704.7	705.7
4	625.5	53.05	660.8	661.4
5	779.0	25.0	786.7	786.8

**Table 14.7 Volume correction test cases at atmospheric pressure—volume correction to 15.0°C and 0.0 kPa(g)**

Test case	Inputs				Outputs				
	Metered volume (m <sup>3</sup> )	Density (kg/m <sup>3</sup> ) @ 15.0°C	Observed temp. (°C)	Observed pressure (kPag)	CTL	CPL	CTL corrected volume (m <sup>3</sup> )	CTL & CPL corrected volume (m <sup>3</sup> )	CTL & CPL corrected volume (m <sup>3</sup> ) rounded*
6	60.0	903.5	40.5	700.0	0.98071	1.00050	58.842368	58.871812	58.9
7	15.0	779.0	3.9	400.0	1.01120	1.00034	15.167952	15.173133	15.2
8	100.0	1008.0	89.0	3700.0	0.95472	1.00255	95.472126	95.715578	95.7
9	250.0	875.5	5.0	200.0	1.00799	1.00013	251.998452	252.030396	252.0
10	150.0	640.0	75.0	1000.0	0.90802	1.00365	136.203308	136.700489	136.7

\*The CPL and CTL shown are rounded to five decimal places, but they are not rounded prior to calculating the volumes. Only the final volume is rounded to one decimal place to meet reporting requirements. The corrected volumes are shown to six decimal places for verification purposes.

**Table 14.8 Other liquid hydrocarbon density correction test cases—density correction to 15.0°C**

Test case	Inputs		Outputs	
	Liquid density @ observed temperature and base pressure (kg/m <sup>3</sup> )	Observed temperature (°C)	Liquid density corrected to 15.0°C (kg/m <sup>3</sup> ) with hydrometer correction	Liquid density corrected to 15.0°C (kg/m <sup>3</sup> ) without hydrometer correction
11	525	92.5	614.2	614.9
12	412.5	11.4	404.5	404.5
13	355.5	84.45	506.7	506.9
14	623.5	53.05	658.1	658.7
15	652.5	25	661.3	661.5

**Table 14.9 Volume correction test cases at equilibrium vapour pressure—volume correction to 15.0°C and equilibrium vapour pressure**

Test case	Inputs					Outputs				
	Metered volume (m <sup>3</sup> )	Density (kg/m <sup>3</sup> ) @ 15.0°C and EVP	Observed temp. (°C)	Observed pressure (kPag)	Equilibrium vapour pressure (kPa) @ observed temp.	CTL	CPL	CTL corrected volume (m <sup>3</sup> )	CTL & CPL corrected volume (m <sup>3</sup> )	CTL & CPL corrected volume (m <sup>3</sup> ) rounded*
16	60.0	544.5	40.5	1645.0	738.0	0.93642	1.0054	56.184942	56.488356	56.5
17	15.0	402.0	3.9	1125.0	1125.0	1.05931	1.0000	15.889672	15.889672	15.9
18	100.0	632.0	55.0	348.0	213.0	0.93587	1.0004	93.586521	93.623473	93.6
19	250.0	512.5	5.0	1500.0	494.0	1.02732	1.0041	256.830532	257.880793	257.9
20	150.0	356.5	-14.5	4260.0	1650.0	1.20782	1.0224	181.173148	185.235683	185.2

\*The CPL and CTL shown are rounded as per their respective standards. CPL is rounded to four decimal places and CTL to five decimal places. They are not rounded prior to calculating the volumes. Only the final volume is rounded to one decimal place to meet reporting requirements. The corrected volumes are shown to six decimal places for verification purposes.

## 14.10 Measurement Records

For all metering equipment covered by this section, records must be kept as outlined in the following report types and made available for examination by ER. Licensees are given flexibility in the formatting of these reports. It is not necessary to present the information exactly as outlined.

These records must be maintained for mechanical, electromechanical, or EFM. EFM systems may retain this information automatically. It is advisable to save the records on a regular basis and when metering problems occur, so they are not lost when memory is full or when the EFM system is shut off.

The reports must be recorded using electronic/magnetic (not necessarily on the EFM system), printed, or handwritten media and retained for a minimum of 12 months. They must be produced upon request by ER.

### 14.10.1 The Daily Report

The following information must be recorded on a daily or per test basis for test meters only:

1. Test meter and well identification
2. Test period accumulated flow
3. Hours on production

### 14.10.2 The Monthly Report

The monthly report is for the entire system, providing data for each measurement point. It is to contain the following at each measurement point as applicable:

1. Monthly cumulative flow
2. Indications of any change made to volumes, and supporting documentation
3. Total hours on production for production or test meters only

### 14.10.3 The Event Log

When any parameter that affects the flow calculation is changed, such as meter factor, fluid densities, or transmitter range, a process is required to record the change. In an EFM system this can be accomplished using the event log within the EFM if so equipped. These parameter changes can also be recorded manually on paper or electronic records.

The event log must include such items as:

1. Instrumentation range changes
2. Algorithm changes
3. Meter factor or k-factor changes
4. Orifice plate changes
5. Fixed fluid density changes
6. Other manual inputs

The log must identify the person making the change and the date of the change.

#### 14.10.4 EFM Specific Reports

The following reports are required together with those in 14.11.1 and 14.11.3 where applicable.

##### 14.10.4.1 The Daily Report

The following information must be recorded on a daily basis:

1. Meter identification
2. Daily accumulated flow
3. Hours on production for production or test meters only
4. Production data audit trail—include at least one of the following:
  - a. Instantaneous values for flow rate, operating pressure (if it is applicable) and temperature taken at the same time each day.
  - b. Daily volume and average daily values for operating pressure (if it is applicable) and temperature.
  - c. Hourly accumulated flow rate and average hourly values for operating pressure (if it is applicable) and temperature.

##### 14.10.4.2 The Meter Report

The meter report is primarily required to confirm that the EFM system is operating properly. A meter report is not required when using mechanical or electromechanical systems, where many of these values are fixed. For these mechanical or electromechanical meters, records are required to verify that the various factors used in the calculation are correct.

The meter report details the configuration of each meter and flow calculation information. It must include the required parameters to demonstrate that the net standard volume is being properly computed from the gross indicated volume. The type of EFM system device will determine which of the following are required:

1. Instantaneous flow data
  - a. Gross and net flow rate or gross and net volume calculated over a time period such that the correction factors are not changing
  - b. Operating pressure
  - c. Differential pressure if it is applicable
  - d. Flowing temperature
  - e. Flowing density
  - f. Sediment and water content if an on-line S&W monitor is used
  - g. CTL
  - h. CPL
  - i. CTPL
2. Current configuration information
  - a. Meter identification

- b. Date and time
- c. Standard Pressure
- d. Standard Temperature
- e. Flowing density or density at standard conditions if a fixed density is used
- f. Meter factor and/or k-factor
- g. Shrinkage factor where it is applicable

#### **14.10.4.3 The Alarm Log**

The alarm log includes any alarms that may have an effect on the measurement accuracy of the system. The time of each alarm condition and the time each alarm is cleared must be recorded. The alarm log includes such items as:

1. Master terminal unit failures
2. Remote terminal unit failures
3. Communication failures
4. Low-power warning
5. High/low volumetric flow rate
6. Overranging of end device

## 15 Water Measurement

This section presents the requirements for measurement and reporting of water from oil and gas production, water source, water injection and disposal, waste processing and disposal, storage and disposal cavern, and thermal *in situ* schemes.

All liquid water produced at a well or measured at a group separator is considered production and must be reported to Petrinex. Water that is in the vapour phase under separator conditions must not be reported as production even if it drops out later in a gas gathering system, refer to Section 15.2.1.6 for more details. Reported water volumes must be corrected to 15.0°C when stipulated in this Directive. All sample analyses, test data, and test date records must be submitted to ER upon request.

For the purpose of this Directive, water types are:

1. Water
2. Freshwater
3. Brine (this is only acceptable for potash disposal wells), see Section 15.2.9
4. Water produced related to Geothermal or other minerals extracted from water.

Production of nonsaline water may require a groundwater diversion permit from the Water Security Agency.

### 15.1 Base Requirements for Water Measurement, Volume Calculation, Production Data Verification, and Audit Trail

For general liquid measurement, inventory determination, liquid volume calculation, production data verification, audit trail, and EFM requirements, see Section 14.

For water meter proving method, see Sections 2.4 and 2.78.

The main difference between measurement of water and other hydrocarbon fluids is that the uncertainty limits are generally less stringent for water. See Section 1.7.

### 15.2 Water Measurement and Accounting Requirements for Various Facility Types

#### 15.2.1 Gas Facilities

The three methods for determining gas production for a gas well are separated gas measurement, effluent measurement, and gas proration. The requirements for determining water production vary among these three methods, and there are also some variations within each method.

##### 15.2.1.1 Gas Single-Well Battery (Petrinex facility subtype: 351) or Gas Multi-well Group Battery (Petrinex facility subtypes: 361)

Each gas well must have its' own permanent separation facilities. Water production must be determined by one of the following methods:

1. Metering at the water leg of a three-phase separator;

2. Metering at the water/condensate leg of a two-phase separator and taking a sample of the liquid phase to determine the percent S&W and liquid hydrocarbon. Refer to Section 14.8 for more detail;
3. Directing water production to a tank and delivering by either truck or pipeline for disposition. The monthly production volume is then determined from the delivery volume measured by the receiving facility and tank inventory change by gauging the tank; or

#### **15.2.1.2 Gas Multi-well Proration SW Saskatchewan Battery (Petrinex facility subtype 363)**

Water determination and reporting are not required inside SW Saskatchewan shallow gas areas and stratigraphic units or zones at the well level. They are only required at the battery level. See Section 7.2 for more detail.

#### **15.2.1.3 Gas Multi-well Proration Outside SW Saskatchewan Batteries (Petrinex facility subtype 364)**

Outside SW Saskatchewan shallow gas areas and stratigraphic units or zones that satisfy the proration requirements in Section 5.4, an individual well WGR may be used to determine water production volume. See Section 7.3 for more detail.

#### **15.2.1.4 Gas Multi-well Effluent Measurement Battery (Petrinex facility subtype: 362)**

Gas wells not configured with separation and measurement of each phase at the wellhead and at which all of the multiphase fluid passes through the same meter are subject to effluent or wet gas measurement. The water production volume is calculated based on the WGR obtained from a ECF test. See Section 7.4 for ECF-WGR testing and actual well water production calculation requirements and Section 8.4.2 for sampling and analysis requirements.

#### **15.2.1.5 Low-Pressure Gas Systems**

If all wells at a gas battery operate at a pressure < 350.0 kPa without any mechanical or other external means of water lifting used at any of the wells in the battery, licensees may determine water production using one of the following methods:

1. Battery level WGR, which is determined by metering the battery water production and dividing it by the battery gas production on a monthly basis. The resulting WGR is then applied to all wells within the battery based on the wells' reported gas production to determine each well's reported water production; or
2. Engineering estimates based on the water vapour pressure at the well's flowing conditions, and other methods supported by good engineering practice.

#### **15.2.1.6 Water at Gas Gathering Systems (GS) (Petrinex facility subtypes: 621)**

When water is received or disposed of from a gas gathering system, the volume must be measured and reported in accordance with the following requirements:

1. If there is no known source(s) of water coming into the gas gathering system, all collected water must be reported as water condensation on Petrinex as:

Activity: REC  
Product: WATER  
From/To: SKWC receipt

2. If there is some known source of water, any delivery volume over and above the known source of water delivered into the gas gathering system must be reported on Petrinex as SKWC receipt. This applies when there are multiple single wells, gas group, and/or other facilities tied into the gas gathering system with commingled water.
3. If there is only a single-well battery or one proration battery as the upstream source of the produced water with no water measurement upstream of the disposition location, this location may be designated as the facility group measurement point at the battery, and all water collected can be used for reporting from the battery.
4. The gas gathering system must report receipt of water from any upstream facilities and report disposition if water is going through the gas gathering system to other facilities further downstream.

#### **15.2.1.7 Gas Plants (Petrinex facility subtypes: 401 to 407)**

Gas plant inlet water measurement is required at each inlet separator when the total receipt volume is  $> 50.0 \text{ m}^3/\text{month}$  for the entire plant. If there is only one inlet separator and no other source of water entering the plant, the plant water disposition plus inventory change may be used for inlet water reporting.

#### **15.2.2 Crude Oil Facilities**

The licensee must separate the water from the oil and measure the water if the total water production at a well or battery  $> 50.0 \text{ m}^3$  per month and the water cut is  $> 0.5$  per cent of the total liquid production. The battery water disposition must be measured if  $> 50.0 \text{ m}^3/\text{month}$ . The receiving facility is responsible for measurement and reporting of the water disposition.

The two methods for determining oil/water production for oil wells are separated measurement and proration. The requirements for determining water production vary between these two methods, and there are also some variations within each method. See Section 6.4.

##### **15.2.2.1 Crude Oil Single-Well Battery (Petrinex facility subtype: 311) and Heavy Crude Oil Single-Well Battery (Petrinex facility subtype: 325)**

For a single-well battery trucking its emulsion off site, the water production volume is determined by the battery disposition plus change in inventory and the measurement occurs at the receiving facility. Refer to Section 14.8.1 for more details.

##### **Exception to Single-Well Battery Water Production Volume**

If a single-well battery with a two-phase or three-phase separator is delivering produced oil/emulsions by pipeline to another battery and the water production  $\leq 50.0 \text{ m}^3/\text{month}$ , the water cut may be determined by taking three spot samples at intervals at least one week apart within the month and averaging the results. Refer to Section 14.8.1 for more details on S&W determination.

##### **15.2.2.2 Crude Oil Multi-well Group Battery (Petrinex facility subtype: 321) and Heavy Crude Oil Multi-well Group Battery (Petrinex facility subtype: 326)**

Total water production from a multi-well group oil battery must be determined in the same way as from an oil single-well battery. Refer to [Section 15.2.2.1](#) for more details.

### **15.2.2.3 Crude Oil Multi-well Proration Battery (Petrinex facility subtype: 322) and Heavy Crude Oil Multi-well Proration Battery (Petrinex facility subtype: 327)**

Water production from a multi-well proration oil battery is based on well testing and proration from the battery disposition volumes plus inventory change at month end. (see [Sections 6.4 and 6.6.](#))

### **15.2.2.4 Heavy Crude Oil Paper Battery (Petrinex facility subtypes: 313) and Crude Oil Multi-well Swab Battery (Petrinex facility subtypes: 314 and 316)**

Total water production from a heavy crude oil paper battery or a crude oil multi-well swab battery must be determined in the same way as a heavy crude oil single-well battery. Refer to Section 15.2.2.1 for more details.

## **15.2.3 Water Source Production**

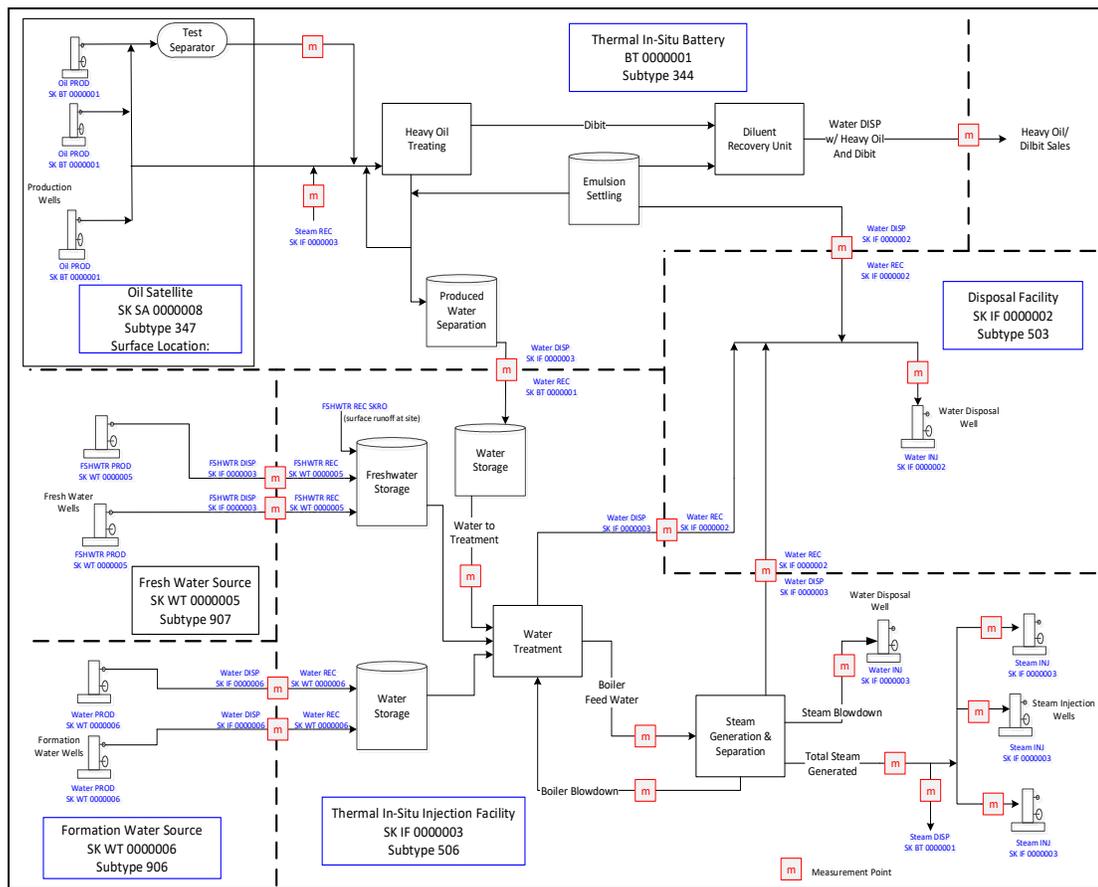
Water produced from water source wells or other sources, such as rivers and lakes, must be continuously measured before commingling with water or fluids from another source. If a source well is producing gas, the related gas production must be separated and measured or estimated and reported.

### **15.2.3.1 Reporting Requirements**

Reporting on Petrinex must be under the SKWS facility subtype 905 for non-ER licensed surface water sources, such as rivers and lakes. If the water is delivered to a thermal *in situ* scheme, the water receipt must be reported at the injection facility and not the battery facility of the thermal *in situ* scheme.

For ER-licensed water source wells, reporting on Petrinex must be under the SKWI code and the applicable facility subtype. If the water is produced from a fresh-water-bearing formation, the water is considered fresh water and reported under the SKWT facility subtype of 907. If the water is produced from a stratigraphic unit that is not fresh-water-bearing, the water is reported under the SKWT facility subtype of 906. If the water production is delivered to a thermal *in situ* scheme, the water receipt from the SKWT facility must be reported at the injection facility and not the battery facility of the thermal *in situ* scheme.

Figure 15.1 Example of water source reporting setup



### 15.2.4 Water Injection and Disposal Facility

Water injected into injection or disposal wells must be continuously metered at each wellhead at the injection site and used for reporting to Petrinex. Water injected into injection or disposal wells constructed prior to September 11, 2012 must be continuously metered into individual wells but may be metered at the injection facility or a field injection header if the meter is not located at the wellhead.

The acid gas injection/disposal measurement scenarios in Section 11.4.5.3 can also be applied to these types of wells, but injection and disposal wells cannot be in the same injection facility except for facility subtype 506. Each injection/disposal well may have its own injection facility reporting code or may be part of another injection facility. If there is more than one facility sending water to the injection facility, each receipt must be measured before commingling.

Skim oil recovered from these facilities must be ≤ 1.0 per cent of the total received volume based on a six month rolling average basis. Licensees may be contacted to explain the origin of the excessive skim oil. If the skim oil is > 1.0 per cent, ER inspector or auditor may direct in writing that the licensee implement changes to improve the skim oil percentage, and these directions will become conditions of operation for that facility. Examples of conditions are as follows:

1. Investigate where the oil originated from.
2. Install delivery point measurement at the receipt point.
3. Inform the source licensee to ship emulsion to a proper treatment facility.
4. Report the oil receipt at the injection facility from the source facility.

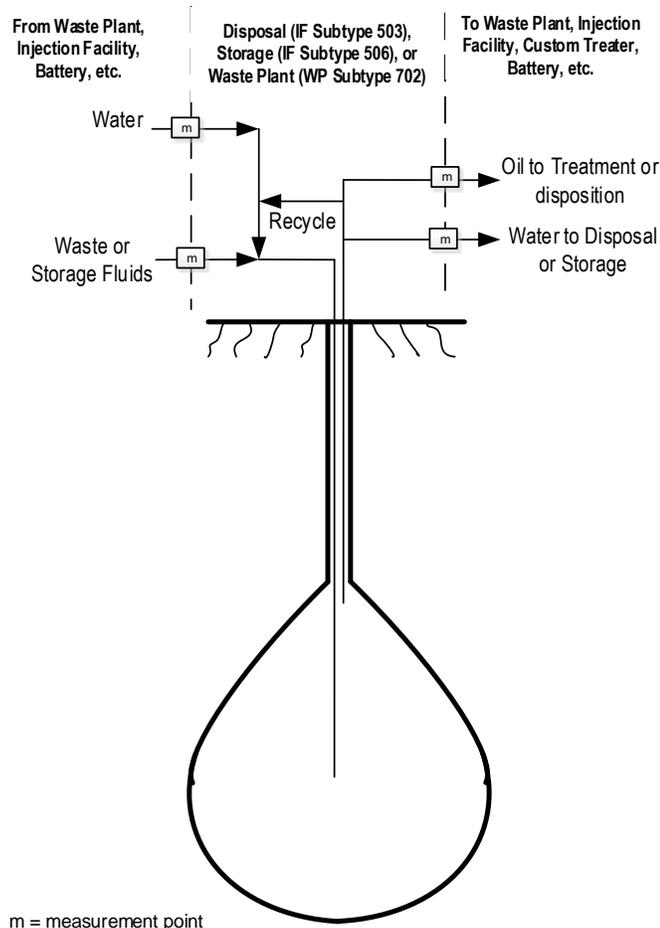
### 15.2.5 Waste Processing and Disposal Facility

All products received at a waste processing facility must be measured, sampled, analyzed, and reported as oil, water, and solids according to the approval conditions. The disposal facility will auto-populate Petrinex with the source of oil and/or water including solids if the fluid is produced from an upstream oil and gas facility, such as tank bottom fluids.

### 15.2.6 Storage and Disposal Cavern

Each storage and disposal cavern must be reported as a separate facility on Petrinex, have its own inlet and outlet measurement system from the cavern washing stage onwards, and maintain separate oil, water, and solids inventory.

**Figure 15.2 Cavern measurement**



### 15.2.7 Thermal In Situ Scheme (Petrinex facility subtype: 506)

Thermal *in situ* schemes use water to generate steam for injection into formations to produce heavy oil. The single point measurement uncertainty requirements for the critical measurement points have been tightened to facilitate water balancing, refer to Sections 12.3.3 through 12.3.4 for more details. The licensee must use the Petrinex subtype code 506 for an injection facility, but a disposal facility can be a facility subtype 503 or part of the facility subtype 506. See [Figure 15.1](#).

All steam volume is to be reported as cold water equivalent volume at 15.0°C.

### 15.2.8 Downhole Water Disposal or Injection

When water is separated from the gas down hole and injected into another stratigraphic unit or zone or formation without coming to surface, the water volume must be measured if it is > 50.0 m<sup>3</sup>/month or estimated if it is ≤ 50.0 m<sup>3</sup>/month.

### 15.2.9 Brine Measurement and Reporting (Petrinex facility subtype: 518)

This section presents the requirements for measurement and reporting of brine injection into potash disposal wells.

The product BRINE is only reported at ER-licensed disposal wells and injection facilities related to the potash industry.

All brine volumes injected into potash disposal wells must be reported to Petrinex. Reported brine volumes must be corrected to 15.0°C.

Brine injected into disposal wells must be continuously metered at each wellhead at the injection site and used for reporting to Petrinex. Brine injected into disposal wells constructed prior to September 11, 2012 must be continuously metered into individual wells but may be metered at the injection facility or a field injection header if the meter is not located at the wellhead.

For general liquid measurement, liquid volume calculation, data verification, audit trail, and EFM requirements, see Section 14.

For brine meter proving method, see [Sections 2.4](#) and [Section 2.8](#).

See Sections 1.7 for uncertainty limits.

### 15.2.10 Load Water Reporting

Current ER load water reporting requirements state that when a well is put on production following load water injection, the water produced from the well must be reported into Petrinex as load water recovery until the entire volume of injected load water is recovered. After all of the load water has been recovered, the water produced from the well must be reported as water production.

In most cases, it is unlikely that all of the injected load water during a well completion operation will be recovered after the well is put on production; some load water will remain in the formation. Licensees may, at their discretion, discontinue reporting load water recovery after a well has been on production for 12 months following load water injection. In this case, the licensee would “zero out” the load water inventory using the load fluid

inventory adjustment—LDINVADJ—activity in Petrinex. Water produced from the well after zeroing out the load water inventory must be reported as water production.

It is the responsibility of the licensee to ensure that fluids such as flow back, load fluid, and produced water are managed properly based on their composition (e.g., sent to an appropriate waste management facility or injection/disposal well, or treated and reused). Information regarding fluid management can be obtained from the Field Services Branch.

### 15.3 Water Gas Ratio (WGR) Testing Methodology

Well testing requirements are as follows:

1. A WGR test is required for gas multi-well proration outside SW Saskatchewan wells.
2. An ECF-WGR test is required for effluent measured wells. Refer to Section 7.4 for more details.
3. New wells must have the required tests conducted within the first 30 days of production and annually thereafter, unless otherwise stated or exempt, refer to Section 5.4 for more detail. If the licensee can demonstrate that the WGR has not stabilized, such as load water has not been fully recovered, multiple tests may have to be conducted over the next few months to determine the stabilized production WGR.
4. Wells that can have operation or production characteristics changed because of such events as workovers or chemical stimulations must have a test conducted within 30 days.

#### 15.3.1 WGR Testing

The WGR test must be conducted using a properly sized three-phase separator with measurement of all phases as follows:

1. The test must begin only after a stabilization period.
2. The test duration must be a minimum of 12.0 hours.
3. If the well is occasionally slugging water, the test duration must be increased to ensure that the test is representative.
4. Consistent testing procedures must be used for consecutive tests to identify when a change in a well's flow characteristics has occurred.
5. The water volume must be measured by collecting it in a suitable container or by using a water meter.
6. The gas and condensate volumes must be measured.
7. The condensate must be sampled during every test and analyzed for the components to determine an updated gas equivalent factor. The sample may be taken from the condensate leg of a three-phase separator or the liquid leg of a two-phase separator. The water must be removed from the condensate before the analysis.
8. The test volumes and date for each test must be recorded.

9. The WGR must be determined by dividing the water volume by the sum of the measured gas volume and the gas equivalent volume (GEV) of the measured condensate (if the condensate is recombined with the gas and sent to a gathering system or gas plant for further processing). For more details refer to Section 7.3.2.
10. If a three-phase separator is not available, alternative equipment, such as a two-phase separator with a total liquid meter and continuous water cut analyzer, is acceptable. Other options that provide equivalent liquid volume accuracy may also be considered on a case-by-case basis by ER.

### 15.3.2 WGR Calculation

WGR calculation for a gas multi-well battery is as follows:

1. Calculate the total monthly gas production volume.

$$\text{Total monthly well gas volume} = \text{Well measured gas volume} + \text{GEV of well measured condensate if recombined.}$$

2. Calculate the WGR for each well.

The WGR is to be calculated to six decimal places and then rounded to five decimal places, as follows:

$$\text{WGR} = \text{Well test water volume} \div (\text{Well test gas volume} + \text{GEV of well test condensate if recombined})$$

3. Calculate reported water production for each well.

Using the WGR, reported water production for the well is calculated as follows:

$$\text{Reported monthly well water volume} = \text{Total monthly well gas volume} \times \text{WGR}$$

For gas multi-well proration batteries, refer to Section 7

## **Appendix 1 Documents Replaced Fully or Partially by Directive PNG017: Measurement Requirements for Oil and Gas Operations**

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### **Saskatchewan**

#### ***Total Replacement (rescinded on date Directive PNG017 becomes effective):***

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Sections 78 through 82 of *The Oil and Gas Conservation Regulations, 2012*

Sections 84 through 87 of *The Oil and Gas Conservation Regulations, 2012*

PNG Guideline 17: Production Testing of Gas Wells

PNG Guideline 23: Monthly Gas Measurement Exemption – Oil Wells

MRO 37/85 - Individual Well Tests in Units – March 11, 1985

All special measurement exemptions granted by Minister's Order for certain designated pools

## Appendix 2 Glossary

The definitions that follow are for the purposes of this Directive only.

**Absolute Density** – The mass per unit volume of a gas or liquid at a specific pressure and temperature. Absolute densities are generally expressed in kg/m<sup>3</sup> at 101.325 kPa(a) and 15.0°C.

**Accuracy** – The ability of a measuring instrument to indicate values closely approximating the true value of the quantity measured.

**Acid Gas** – Gas separated out during the treating of sour gas that contains hydrogen sulphide (H<sub>2</sub>S), totally reduced sulphur compounds, and/or carbon dioxide (CO<sub>2</sub>).

**Allocation Factor** – A factor, that is used to adjust the fluid receipt volumes (considered estimates) to reported volumes based on inventory and disposition measurements at facilities where only fluids received by truck are handled, such as custom treaters or terminals.

**American Petroleum Institute** – Is the source of many measurement standards in the oil and gas industry. (API Manual of Petroleum Measurement Standards).

**Analog Transmitter** – Transmitters that use analog circuitry to convert their sensor output to either 4.0-20.0 milliamps or 1.0-5.0 volts.

**Annually** – Once every four calendar quarters (once a year).

**Annual Average Daily Flow Rate** – A average daily flow rate calculated based on a calendar year. For a new well or facility the daily average flow rate is calculated based on the last 12 months. If the well or facility does not operator for a full calendar year, then the average daily flow rate is calculated by the average flow rate based on the number of days operating. For example: total annual volume divided by number of days the well or facility was operating.

**API** – see American Petroleum Institute.

**Approved** – Written acceptance by ER.

**Associated Gas** – Gas produced in association with oil production at oil wells. Commonly known as solution gas.

**ATC** – see Mechanical Automatic Temperature Compensators without Gravity selection.

**ATG** – see Mechanical Automatic Temperature Compensators with Gravity selection.

**Atmospheric Pressure** – The pressure exerted by the weight of the atmosphere at the point of measurement.

**BA ID** – see Business Associate Identifier.

**Base Conditions** – For Gaseous fluids are 101.325 kPa at 15.0°C and for liquids are 101.325 kPa at 15.0°C. Also known as standard conditions, stock tank conditions or reference conditions.

**Base Pressure** – Atmospheric pressure of 101.325 kPa

**Base Temperature** – 15.0°C

**Battery** - Common storage facilities receiving production from a well or wells and includes equipment for separating the fluid into oil, gas, water and any other substances and for measurement. (Source: *The Oil and Gas Conservation Regulations, 2012*)

**Bias** – Any influence on a result that produces an incorrect approximation of the true value of the variable being measured. Bias is the result of a predictable systematic error.

**Biennially** – Once every eight calendar quarters (once every two years).

**Bimonthly** – Once every two calendar months.

**Boiler Blowdown** – Boilers used in thermal recovery processes typically produce steam with a quality between 75.0 per cent and 80.0 per cent. This results in 20.0 per cent to 25.0 per cent of the boiler feed water not being vapourized. This water stream, which is separated from the steam, leaving the boiler is called boiler blowdown and contains more concentrated total dissolved solids, typically 4.0 to 5.0 times more, than the boiler feed water.

**Brackish Water** – A product type related to cross border reporting for Petrinex volumetrics reporting only. The product Brackish Water (BRKWTR) is only used to accommodate cross border receipts from Alberta and should not be used in any other situations. As defined in Alberta, water from a saline water source that has total dissolved solids  $\geq 4000$  milligrams per litre (mg/L).

**Brine** – A product type related to the potash industry for Petrinex volumetrics reporting only. This product, comprised of water saturated with salt, only reported at disposal wells and injection facilities related to the potash industry.

**Business Associate Identifier** - A unique five-digit identifier assigned to each corporate entity for use in Petrinex.

**Butanes** – A liquid mixture mainly of butanes that ordinarily may contain some propane or pentanes plus. For reporting purposes there are NC4-SP, NC4-MX, IC4-SP, and IC4-MX.

**Calendar Quarter** – January to March, April to June, July to September, October to December.

**Calibration** – The process or procedure of adjusting an instrument, such as a meter, so that its indication or registration is in satisfactory or close agreement with a reference standard. (Source: API MPMS)

**Calibration Standard** – A certified device used in calibration or proving that has a known value traceable to national reference standards maintained by the National Research Council in Ottawa, Ontario.

**CBM** – see **Coalbed Methane**

**CF** – see **Correction Factor**.

**CFM** – see **Composite Meter Factor**.

**CGR** – see **Condensate-Gas Ratio**.

**Clean Oil** – Oil with  $\leq 0.5$  per cent sediment and water.

**Clean Oil Terminal** – Terminals that receive trucked or pipelined clean oil.

**Coalbed Methane** – Natural gas that is found in coals.

**CMF** – see **Composite Meter Factor**.

**Cold Water Equivalent** – Steam volume measurements corrected to a standard temperature of 15.0°C and reported in cubic metres (m<sup>3</sup>).

**Common Crown or Common Freehold Royalty** – When all the wells in a battery are produced:

- 1) under Crown mineral leases and the royalty status is the same for each well;
- 2) under leases granted by one Freehold mineral owner and the Freehold mineral owner receives the same royalty rate for each well; or
- 3) under leases granted by more than one Freehold mineral owner, and the total royalty rate for each owner for each well is the same.

**Common Ownership** – All wells in a battery belong to the same working interest participant, or if there is more than one working interest participant, each working interest participant has the same percentage interest in each well in the battery.

**Composite Meter Factor** – A factor that is calculated by dividing the temperature corrected prover volume by the indicated meter volume for a prover run. The final CMF is often averaged from the results of multiple prover runs. The CMF includes corrections for the effects of any combination of temperature, pressure or shrinkage.

**Compressibility (apparent)** – The arithmetic sum of the actual compressibility of a liquid and the volume change per unit of volume of the confining container caused by a unit change in pressure at a constant temperature.

**Compressibility (liquid)** – The change in volume per unit of volume of a liquid caused by a unit change in pressure at constant temperature.

**Compressibility factor** – The ratio of the actual volume of gas at a given temperature and pressure to the volume of gas when calculated by ideal gas law.

**Compressibility** – The property of a material which permits it to decrease in volume from the ideal state when subjected to an increase in pressure at constant temperature. (Source AGA)

For liquids, see Correction for Pressure and Liquid (CPL)

For gas, compressibility factor “Z” is the deviation from the ideal Boyle’s and Charles’ law behaviour.

**Compressor Station** – An installation of service equipment that receives natural gas from a well, facility, or gathering system prior to delivery to market or other disposition and is intended to maintain or increase the flowing pressure of the gas; includes any equipment for measurement.

**Condensate** – A liquid hydrocarbon product with a density of  $\leq 780.0 \text{ kg/m}^3$  that:

- i. existed in the reservoir in a gaseous phase at original conditions and;
- ii. that is recovered from a gas stream when pressure and temperature are reduced to and not lower than those at base conditions. (Source: *The Oil and Gas Conservation Regulations, 2012*)

**Condensate, Separator Liquids** – Separator liquids are a mixture of hydrocarbon components that remain in a liquid state under the equilibrium pressure and temperature conditions established in a two-phase or three-phase separator. The composition and physical properties of separator liquids are highly variable and are a function of separator inlet fluid composition and separator pressure and temperature conditions.

**Condensate, Stabilized Liquids** – Stabilized liquids are a mixture of hydrocarbon components that remain in the liquid state following single-stage flash evaporation. Stabilized condensate is expected to have low concentrations of light ends ( $C_1$ - $C_4$ ) components. The composition and physical properties of the stabilized condensate is a function of the equilibrium pressure and temperature conditions of the stabilizer and the composition of the stabilizer feed from which it was derived.

**Condensate-Gas Ratio** – Is calculated by dividing the condensate volume by the gas volume of a well test. See [Table 7.1](#). for calculation and rounding requirements.

**Condensate Proration Factor** – Total battery measured daily or monthly condensate volume divided by total battery estimated daily or monthly condensate volume. See Sections 7.3.2 and 7.4.2.

**Continuous Stack Emission Monitor** – A device that measures sulphur stack emissions at a sour gas processing facility.

**Continuous Measurement** – Uninterrupted measurement

**Correction Factor** – A correction factor is any mathematical adjustment made to take into account deviations in volume related to temperature or pressure in either the sample or the measured volume. It must be determined in accordance with API MPMS, Chapter 11.

**Crude Oil** – Crude petroleum oil and any other hydrocarbon, regardless of density, that is or is capable of being produced from a well in liquid form, but does not include condensate. (Source: *The Oil and Gas Conservation Regulations, 2012*)

**Crude Oil Swab Well** – A well without downhole production tubing, mechanical lift and well control equipment. Production occurs through the lifting of well liquids by a device installed on a wireline.

**CSEM** – see **Continuous Stack Emission Monitor**.

**CSS** – see **Cyclic Steam Stimulation**.

**CTL** – see **Correction for the effect of Temperature on Liquids**.

**Custody Transfer Point** – The point where legal and commercial transfer of production physically or is deemed to occur. Every physical custody transfer point is always a delivery point. In Saskatchewan, this custody transfer point is different than the Petrinex custody transfer point which generates a pipeline split.

**Custom Treater** – A system or arrangement of tanks and other surface equipment receiving oil or water emulsion exclusively by truck for separation and treating prior to delivery to market or other disposition.

**CWE** – see **Cold Water Equivalent**.

**Cyclic Steam Stimulation** – A thermal enhanced oil recovery process where alternate cycles of steam injection and oil production are conducted in a reservoir through the same wellbore.

**DCS** – see **Distributed Control System**.

**Dead Oil** – Oil not containing any entrained or formation gas and stabilized to ambient conditions.

**Dead Oil Meters** – Dead oil meters are typically those used for delivery point or custody transfer point measurement of hydrocarbon liquids that have been de-gassed to ambient conditions.

**Deemed Dry Production** – Applies to gas effluent measured wells that qualify for a testing exception based on the effluent testing decision tree. This includes wells that are categorized within a stratigraphic unit or zone-based effluent testing exception where the average liquid gas ratio results of testing are less than  $0.056 \text{ m}^3/10^3\text{m}^3$ .

**Dehydrator** – An apparatus designed and used to remove water from raw gas.

**Delivery Point** – The point at which the production or disposition of hydrocarbon liquid or gas from a facility is measured. Every physical custody transfer point is always a delivery point. A delivery point is not always a custody transfer point. See Section 1 for further information.

**Delivery Point Measurement** – The level of measurement uncertainty required at a delivery point.

**Dew Point** – The temperature at any given pressure at which liquid begins to condense from a gas or vapour. It is specifically applied to the temperature at which water vapour starts to condense from a gas mixture (water dew points) or at which hydrocarbons start to condense (hydrocarbon dew point). (Source AGA Definitions).

**Digital (Smart) Transmitter** – A transmitter that contains a microprocessor used for digital signal processing and calculation purposes. The calculations apply factory characterization of the sensor calibration and dynamic compensation for other process and environmental effects to the sensor output.

**Diluent** – Hydrocarbon liquid such as condensate or C<sub>5</sub>-SP blended with oil to meet pipeline viscosity and density targets.

**Dilution Gas** – Natural gas added to an acid gas stream to ensure adequate heating value during the incineration or flaring of acid gas.

**Distributed Control System** – A control system for a process or plant, wherein control elements are distributed throughout the system. This is in contrast to non-distributed systems, which use a single controller at a central location.

**Dry Gas** – Natural gas without free liquid as per contract conditions consists of little more than methane, producing little condensable heavier hydrocarbon compounds such as propane and butane when brought to the surface.

**Effluent** – Unseparated well production that may contain a mixture of hydrocarbon liquids and gas or water in the stream.

**Effluent Measurement** – The metering of effluent without separation.

**EFM** – see **Electronic Flow Measurement**.

**Electronic Flow Measurement** – Any flow measurement and related system that collects data and performs flow calculations electronically. For more information, reference API MPMS 21.1 and 21.2.

**Emulsion** – A combination of two immiscible liquids, or liquids that do not mix together under normal conditions.

**End Device** – The device or equipment that records the various values used to calculate a volume such as a chart recorder or EFM system. In the scenario of an EFM system, the end device may also perform the calculations necessary to arrive at the measured and corrected gas volume.

**Equilibrium Vapour Pressure (EVP)** – The pressure at which a liquid and its vapour are in equilibrium at a given temperature. When a hydrocarbon liquid has an EVP above the standard pressure (101.325 kPa at 15.0°C), the EVP at 15.0°C is the pressure base.

**Equity** – See common ownership and common royalty.

**Error** – The difference between true and observed values. For more information, see ISO 5168 and ASME MFC 2M.

**Error (random)** – An error that varies in an unpredictable manner when a large number of measurements of the same variable are made under effectively identical conditions.

**Error (spurious)** – A gross error in procedure, for example, human errors or machine malfunctions

**Error (systematic)** – An error that in the course of a number of measurements made under the same conditions on material having the same true value of a variable either remains constant in absolute value and sign or varies in a predictable manner. Systematic errors result in a bias.

**Estimate** – The approximation of a value based on documented and traceable methodologies, calculation, and also based on adequate knowledge of applicable facility processes, metering technology, measurement principles and hydrocarbon and water physical properties.

**Ethane** – A mixture mainly of ethane that ordinarily may contain some methane or propane. Petrinex reporting product types are C<sub>2</sub>-SP (pure ethane) and C<sub>2</sub>-MX (mixture of ethane and propane and other products).

**EVP** – see **Equilibrium Vapour Pressure**.

**Exception** – Circumstances under which if specific qualifying criteria are met measurement devices or procedures are allowed to deviate within specified limits from base measurement requirements.

**Facility** – Any building, structure, installation, equipment or appurtenance that is connected to or associated with the recovery, development, production, storage, handling, processing, treatment or disposal of oil, gas, water, products or other substances, that are produced from or injected into a well, but does not include a pipeline. (Source: *The Oil and Gas Conservation Regulations, 2012*)

**Facility Subtype** – See *Directive PNG032: Volumetric, Valuation and Infrastructure Reporting in Petrinex*

**FDC system** – see **Field Data Capture system**.

**Field Condensate** – Products obtained from natural gas or solution gas before they are delivered to a gathering system. Typically, Field Condensate is a hydrocarbon liquid separated from raw production at a well or a group measurement point, stabilized in a tank, and sold or otherwise disposed of without further processing.

**Field Data Capture System** – A computer system that is used to collect well and facility data about production activities including:

- a. Meter readings and estimates
- b. Production and test hours.
- c. Calculated proration factors and metering differences.
- d. Pressure and temperature readings.
- e. Downtime hours and reasons.

**Flare Gas** - Gas that is combusted in a flare or incinerator at upstream oil and gas operations. Types of gas, if combusted in a flare or incinerator, that must be reported as flare gas include the following:

- Acid gas (routine and non-routine);
- Blanket gas, purge gas, or sweep gas;
- Dilution and make-up gas added to a flare gas stream before flaring or incineration;
- Gas from dehydrator still columns;
- Gas produced during well completions;
- Gas produced during well unloading operations;
- Gas that is flared or incinerated as a result of equipment failures or plant upsets;
- Gas used to operate pneumatic devices (instruments, pumps and compressors starters);
- Pilot gas; and
- Waste gas

**Flow Computer** – A device that calculates and/or compensates the flow or volume based on the meter flow variables. This can be integral to the metering system or completely separate. Examples include, but not limited, are RTU, DCS, or a Net Oil Computer (NOC).

**Flow Meter** – A device used to measure the mass or volumetric flow rate or quantity of a liquid or a gas moving through a pipe.

**Fresh Water** – “fresh-water-bearing formation” means a permanent subsurface water bearing formation with a significant volume of recoverable water that has total dissolved solid concentration < 4 000 milligrams per litre. (Source: *The Oil & Gas Conservation Regulations, 2012*)

**Fuel Gas** – Gas that is combusted and the released energy is used in upstream oil and gas operations. Types of gas that must be reported as fuel gas include gas combusted by the following:

- Catalytic heaters and other building heaters;
- Engines;
- Line heaters;
- Process vessel burners;
- Sulphur recovery unit reaction furnaces; and
- Thermoelectric generators.

**Fugitive Emissions** – Unintentional release of hydrocarbons to the atmosphere.

**Gas** – Natural gas, both before and after it has been subjected to absorption, purification, scrubbing or other treatment or process, and includes all liquid hydrocarbons other than oil and condensate. (Source: *The Oil and Gas Conservation Regulations, 2012*)

**Gas Battery** – A system or arrangement of surface equipment receiving the effluent from one or more gas wells that provides separation, measurement, dehydration, dew point control, compression, or other gas handling functions prior to the delivery to market or other dispositions; does not include gas processing equipment. . Gas batteries include single-well batteries, multi-well group batteries, effluent batteries, and multi-well proration batteries.

**Gas Chromatograph** – An analytical instrument that separates a gas sample into its components and then measures the amount of each separated component. This information is used to determine gas composition for calculating energy content, relative density (specific gravity), compressibility and other related parameters.

**Gas Equivalent Factor** – A factor based on the composition of a hydrocarbon liquid mixture that is used to convert 1.0 m<sup>3</sup> of the hydrocarbon liquid mixture to its equivalent gas volume in 10<sup>3</sup> m<sup>3</sup>. This factor is expressed in terms of 10<sup>3</sup>m<sup>3</sup> gas/ m<sup>3</sup> liquid, is mixture dependent and therefore not a constant for all mixtures.

**Gas Equivalent Volume** – The volume of gas (10<sup>3</sup>m<sup>3</sup>) that would result from converting a known volume of liquid into a gas by applying a Gas Equivalent Factor to the liquid volume.

**Gas Fractionation Plant** – A gas plant that reprocesses natural gas liquids into one or more in-stream components.

**Gas Gathering System** – A reporting entity that may consist of pipelines used to move gas production from oil batteries, gas batteries, or other facilities to another facility, such as a gas plant. This may include compressors, line heaters, dehydrators, measurement, and other equipment.

**Gas in Solution** – Gas dissolved in liquid under pressure.

**Gas Meter** – Broadly used to describe all the equipment or devices that are collectively used to arrive at an indication of a gas volume.

**Gas-Oil Ratio** – The ratio of the number of cubic metres of gas produced from a given source over a given period of time to the number of cubic metres of oil produced from that source over that period.

**Gas Processing Plant (Gas Plant)** – A system or arrangement of equipment used for the extraction of hydrogen sulfide, helium, ethane, natural gas liquids, or other substances from raw gas; does not include a wellhead separator, treater, dehydrator, or production facility that recovers < 2.0 m<sup>3</sup>/day of hydrocarbon liquids without using a liquid extraction process (e.g., refrigeration, desiccant). In addition, does not include an arrangement of equipment that removes small amounts of sulfur (< 0.1 tonne/day) through the use of nonregenerative scavenging chemicals that generate no hydrogen sulfide or sulfur dioxide. (Source: *Directive PNG001: Facility Licence Requirements*)

**Gas Well** – Means:

- (i) a well that is capable of producing gas not associated with oil at the time of production;
- (ii) that part of a well in which the gas-producing stratigraphic unit or zone is successfully segregated from the oil and in which gas is produced separately from the oil;
- (iii) a well from which gas is or is capable of being produced from a reservoir in association with no more than one cubic metre of oil for every 3 500 cubic metres of gas produced from the reservoir; or
- (iv) any other well that may be classified by the Minister pursuant to clause 17(1)(l) of the Act as a gas well for the purposes of the Act and these regulations;

(Source: *The Oil and Gas Conservation Regulations, 2012*)

**GEF** – see **Gas Equivalent Factor**.

**GEV** – see **Gas Equivalent Volume**.

**GIS** – see **Gas In Solution**.

**Good Production Practice** – Production of oil or gas from a well at a rate not governed by an allowable rate of production but limited to what can be produced on the basis of technical parameters without adversely and significantly affecting:

- (i) the ultimate recovery of oil or gas; or
- (ii) the opportunity of other owners to obtain their share of production from the pool;

(Source: *The Oil and Gas Conservation Regulations, 2012*)

**GOR** – see **Gas-Oil Ratio**

**GPP** – see **Good Production Practice**.

**Heavy Oil** – Crude oil having a density  $\geq 920.0$  kg/m<sup>3</sup> at 15.0°C.

**High Vapour Pressure Liquids** – Any hydrocarbon and stabilized hydrocarbon mixture with a Reid Vapour Pressure greater than 14.0 kilopascals (Source: *Oil and Gas Conservation Rules*)

**Hydrocarbon Liquid** – A fluid in the liquid state that primarily consist of one or more of the following: oil, condensate, ethane, propane, butane, pentane, or other heavier hydrocarbon compounds.

**Industry Technical Standards Association** – An industry- and ER -recognized technical association that develops and publishes upstream oil and gas measurement standards and procedures. Organizations include

- the American Petroleum Institute,
- the American National Standards Institute,
- the American Gas Association

- the Gas Processors of America,
- the International Standards Organization, and
- Measurement Canada

**Initial** – As it relates to sample and analysis frequency requirements, an analysis is required within the first six months of operation only, with no subsequent updates required.

**Injection/Disposal Facility** – A system or arrangement of surface equipment associated with the injection or disposal of any substance through one or more wells.

**In Situ Operation** – A scheme or operation ordinarily involving the use of well production operations for the recovery of heavy crude oil from oil sands, or a scheme or operation designated by ER as an *in situ* operation, but does not include a mining operation.

**ISO** – see International Standard Organization

**Internal Inspection** – A visual inspection of the condition of the internal components of a meter including the primary measurement element. The internal components must be removed from service, inspected, replaced or repaired if found to be damaged, and then placed back in service in accordance with this Directive.

**International Standard Organization** – ISO (International Organization for Standardization) is an independent, non-governmental membership organization and the world's largest developer of voluntary International Standards.

**KF** – see **K-Factor**.

**K-Factor** – A term in pulses per unit volume determined during a factory or field proving of a meter. The number of pulses generated by a linear meter divided by the k-factor will determine the indicated volume.

**kPa** – Common multiple units of the pascal are the hectopascal (1 hPa  $\equiv$  100.0 Pa) which is equal to 1.0 mbar, the kilopascal (1.0 kPa  $\equiv$  1000.0 Pa), the megapascal (1 MPa  $\equiv$  1,000,000.0 Pa), and the gigapascal (1 GPa  $\equiv$  1,000,000,000.0 Pa). Standard atmospheric pressure is defined as 101.325 kPa.

**LACT** – see **Lease Automatic Custody Transfer**.

**Lease Automatic Custody Transfer** – An arrangement of equipment that measures the net volume and quality of liquid hydrocarbons. This system provides for the automatic measurement, sampling, and transfer of oil from the lease location into a pipeline. A system of this type is applicable where larger volumes of oil are being produced and must have a pipeline available in which to connect.

**Legal Survey Location** – Is used to identify the location of land parcels in Western Canada. The Legal Survey Locations are based on the Dominion Land Survey (DLS) which is the method used to divide most of Western Canada into one-square-mile (2.6 km<sup>2</sup>) sections for agricultural and other purposes.

**LGR** – see **Liquid to Gas Ratio**.

**Licensee** – The holder of a license according to the records of ER and includes a trustee or receiver-manager of property of a licensee.

**Light Oil** – Oil with a density < 870.0 kg/m<sup>3</sup> at 15.0°C

**Liquefied Petroleum Gas** – LPG consists primarily of propane (C<sub>3</sub>) and butane (C<sub>4</sub>) in a mixture or essentially pure form, with minor components ranging from ethane (C<sub>2</sub>) to normal hexane (C<sub>6</sub>). It is produced either as a by-product of natural gas processing or during refining and processing operations.

**Liquid to Gas Ratio** – A ratio calculated by dividing the total water and/or condensate test volumes by the measured test gas volume.

**Live Oil** – Oil containing mainly pentanes and heavier hydrocarbons that may also contain lighter hydrocarbons with entrained or formation gas under pressure, and is not in a stabilized form. Live oil is commonly measured at the wellhead or facility.

**Live Oil Meters** – Live oil meters are typically those used to measure volumes of oil or oil/water emulsion that are not stabilized.

**Load Fluids** – Any hydrocarbon- or water-based fluids used at any stage in the life of a well including completion, servicing, regular operation, or abandonment. It includes fluids injected into a flowline between a well and the battery to which it produces, e.g., hot oil, dewaxing chemicals.

**Load Oil** – Hydrocarbon-type fluids used as load fluid, including crude oil, condensate, refined oils, and oil-based or oil-soluble chemicals.

**Load Water** – Water-type fluids used as load fluid, including produced, fresh or brackish water and water-based or water-soluble chemicals.

**LPG** – see Liquefied Petroleum Gas.

**Makeup Gas** – Raw or processed gas that is added to another gas stream in order to maintain an adequate heating value during flaring or incineration.

**MARP** – see Measurement, Accounting, and Reporting Plan.

**Master Meter** – A meter of known accuracy that is temporarily connected in series with another meter for the purpose of proving the accuracy of that meter and providing a meter factor.

**Maximum Uncertainty of Monthly Volume** – Relates to the limits applicable to equipment and/or procedures used to determine the total monthly volume.

**MbD** – see Measurement by Difference

**Measured Gas Source(s)** – Single-phase measured gas source(s) downstream of separation and removal of liquids and also includes the gas equivalent volume (GEV) of measured condensate if the condensate is recombined after measurement with the gas downstream of the separator.

**Measured Oil Source(s)** - Oil measured using equipment and/or procedures meeting delivery point measurement requirements and/or uncertainty limits. For emulsion, the delivery point measurement uncertainty limits apply to the total volume determination only.

**Measurement** – A procedure for determining a value for a physical variable. In an oil and gas industry context, the principal measurement technologies and procedures are:

1. Meters for determining flow volumes.
2. Calculated volumes using a proration formula based on test rates.
3. Estimates of volumes based on production facility and product characteristics.
4. Scales for trucked volumes.
5. Automatic gauges and gauge boards for tanks.
6. Instrumentation and gauges for temperature and pressure.

**Measurement, Accounting, and Reporting Plan** – Defined in Saskatchewan *Guideline PNG042: Measurement, Accounting and Reporting Plan (MARP) Requirement for Thermal In-Situ Recovery Projects*

**Measurement by Difference** – Any situation where an unmeasured volume is determined by taking the difference between two or more measured volumes.

**Measurement Canada** – An agency of Industry Canada that is responsible for ensuring businesses and consumers receive fair and accurate measure in financial transactions involving goods and services. The agency develops and administers the laws and requirements governing measurement; evaluates, approves and certifies measuring devices; and investigates complaints of suspected inaccurate measurement.

**Measurement Schematic** – A diagram used to show the actual layout of facilities that traces the normal flow of production from left to right as it moves from wellhead through to sales.

**Measuring Standard** - A device used in calibration or proving that has a known value traceable to national reference standards maintained by the National Research Council in Ottawa, Ontario.

**Medium Oil** – Oil with a density  $\geq 870.0$  and  $< 920.0$  kg/m<sup>3</sup> at 15.0°C

**Meter (noun)** – See Flow Meter.

**Meter (verb)** – To measure using a flow meter.

**Meter Element** – There are three types of meter elements: primary, secondary, and tertiary.

1. Primary element – the internal component(s) of the meter and associated meter run that provides an indication of flow, e.g., orifice plate, shedder bar, or venturi.
2. Secondary element – the part of the meter that senses and records the flow variables, e.g., chart recorder or transmitter.
3. Tertiary element – flow computer that calculates the flow and volume.

**Meter Factor** – A dimensionless number obtained by dividing the volume of gas or liquid passed through a prover (as measured by the prover during proving) by the corresponding meter indicated volume.

**Metering Difference** – Any difference that occurs between the measured inlet (receipt) volumes and the measured outlet (disposition) volumes at a facility.

**Meter Run** – A flow meter installed and calibrated in a section of pipe having adequate upstream and downstream pipe lengths to create a fully developed flow profile.

**Methane** – A mixture mainly of methane that ordinarily may contain some ethane, nitrogen, helium or carbon dioxide.

**MF** – see **Meter Factor**.

**Monthly** – Once per calendar month.

**MPMS** – American Petroleum Institute's **Manual of Petroleum Measurement Standards**.

**Multiphase Fluid** – Unseparated fluid that contains liquids, gases and or solids in a single stream.

**Natural Gas Liquid** – Includes propane, butanes or pentanes plus, or a combination of them, obtained from the processing of raw gas or condensate.

**Net Standard Volume** – The gross standard volume corrected for shrinkage and non-merchantable quantities such as sediment and water.

**NGL** – see **Natural Gas Liquid**.

**Non-Heavy Oil** – Crude oil having a density of  $< 920.0$  kg/m<sup>3</sup> at 15.0°C.

**NSV** – see **Net Standard Volume**.

**OGR** – see **Oil-Gas Ratio**.

**Oil** – Crude petroleum oil and any other hydrocarbon, regardless of density, that is or is capable of being produced from a well in liquid form, but does not include condensate.

(Source: *The Oil and Gas Conservation Regulations, 2012*)

**Oil Battery** – A battery for single or multiple crude oil wells where production may be separated, treated, and measured prior to delivery to sales or other disposition. Oil batteries include single-well batteries, multi-well group batteries, and multi-well proration batteries.

**Oil-Gas Ratio** – Is calculated by dividing the oil volume by the gas volume of a well test. See [Table 7.1](#) for calculation and rounding requirements.

**Oilfield Waste** – Physical waste as that term is ordinarily understood in relation to the activities of the oil and gas industry, but does not include physical waste.

(Source: *The Oil and Gas Conservation Act*)

**Oil Sands** – All sands and rocks that: Contain a highly viscous mixture, composed mainly of hydrocarbons heavier than pentanes, that will not normally flow, in its natural state, to a wellbore; Lie above the top of the Devonian System; and Lie north of Township 73.

(Source: *The Crown Oil and Gas Royalty Regulations, 2012*)

**Oil Well** – Any well capable of producing oil other than a gas well.

(Source: *The Oil and Gas Conservation Regulations, 2012*)

**Operator** – Means: (i) : (i) a person who, as owner, licensee, lessee, sublessee or assignee, has the right to carry on drilling, construction, operation, decommissioning or abandonment of a well or facility and the reclamation of the well or facility site; (ii) in the case of a pipeline, operator as defined in *The Pipeline Regulations, 2000*; (iii) a contractor who on behalf of the person mentioned in subclause (i) or (ii) engages in any of the activities described in that subclause; or (iv) the person designated by the minister as the operator of the well or facility;

(Source: *The Oil and Gas Conservation Regulations, 2012*)

**P&ID** – see **Piping and Instrumentation Diagram**.

**PD Meter** – see **Positive Displacement Meter**.

**Pentanes Plus** – A mixture mainly of pentanes and heavier hydrocarbons that ordinarily may contain some butanes and that is obtained from the processing of raw gas, condensate or crude oil.

**Petrinex** – A centralized web based shared computer system that hosts and disseminates upstream well and facility volumetric data for ER and industry operators of record stakeholders. Each month industry operators of record load volumetric data for the wells and facilities they operate into Petrinex and it is then available to ER and industry operators of record for their specific needs. Petrinex also includes the Enhanced Production Audit Program (EPAP), Oil Valuation, and other functionality. Petrinex was developed and is administered jointly by ER, other Western Regulators, industry operators of record and industry licensees.

**PFD** – see **Process Flow Diagram**.

**Pipeline** – Means:

- (i) a pipe or system of pipes for the transportation of:

- a. Liquid hydrocarbons, including crude oil, multiphase fluids containing hydrocarbons, oil and water emulsions, condensate, liquid petroleum products, natural gas liquids and liquefied petroleum gas;
  - b. Gaseous hydrocarbons, including natural gas, manufactured gas and synthetic gas;
  - c. Water, steam or any other substance where the water, steam or other substance is incidental to or used in the production of crude oil or natural gas; or
  - d. Carbon dioxide;
- (ii) includes any of the following that are incidental to or used in connection with the pipeline:
- a. Tanks, tank batteries, pumps, compressors and racks;
  - b. Storage facilities, loading facilities, terminal facilities, and other similar facilities;
- (Source: *The Pipelines Act, 1998*)

**Piping and Instrumentation Diagram** – A schematic diagram showing piping, equipment, and instrumentation connections within process units.

**Pool** – Means

- i. An underground reservoir that:
  - a) Contains or appears to contain an accumulation of oil or gas; and
  - b) Is separated or appears to be separated from any other reservoir or accumulation in the general structure;
- ii. A portion of an underground reservoir described in subclause (i) that is determined by the minister to be a pool for reasons of development or administration; or
- iii. A group of underground reservoirs described in subclause (i) that is determined by the minister to be a pool for reasons of development or administration;

(Source: *The Oil and Gas Conservation Act*)

**Positive Displacement Meter** – A type of flow meter that measures the volume of fluid by counting repeatedly the filling and discharging of known fixed volumes.

**Primary Measurement** – Where two measurements of a process stream are required, primary measurement is the one that is used on a preferential basis to determine the volume of the process stream.

**Primary Element** – See meter element also known as primary flow element.

**Process and Instrumentation Diagram** – A family of functional one-line diagrams showing hull, mechanical, and electrical systems, such as piping and cable block diagrams.

**Process Flow Diagram** – A diagram commonly used in chemical and process engineering to indicate to the general flow of plant processes and equipment.

**Produced Water** – Water produced in conjunction with oil and natural gas production.

**Propane** – A mixture mainly of propane that ordinarily may contain some ethane and butanes.

**Prorated Production** – Total monthly battery production allocated to wells based on periodic individual well tests and the number of hours during the month that each well was on production.

**Proration** – A procedure in which the total reported monthly battery production is allocated to wells based on periodic individual well tests.

**Proration Battery** – A battery for which well production is determined using proration measurement and accounting procedures.

**Proration Factor** – Ratio of the total reported battery production volume for a fluid divided by the sum of the total estimated well production volumes for that fluid. Separate proration factors are calculated for each fluid (e.g. oil, water and or gas).

**Prover** – A device used to determine the volume of a sample of fluid, to a known standard.

**Prove (proving, proved)** – The procedures or operations whereby a prover volume is compared to an indicated meter volume and both volumes are corrected to applicable pressure and temperature conditions. The prover volume divided by the indicated meter volume yields a meter factor. The meter factor is subsequently applied to indicated meter volumes to determine the adjusted or corrected volume.

**Qualifying Criteria** – Criteria that must be met to qualify for an exception. If the qualifying criteria have been met and the exception is implemented, it may remain in place indefinitely, as long as the exception qualifying criteria continue to be met.

**Quarterly** – Once every calendar quarter. Calendar quarters are:

- a. January – March
- b. April – June
- c. July – September
- d. October – December

**Raw Gas** – Natural gas production from a well before it has been subjected to dehydration, sweetening, or other processes.

**RD** – see **Relative Density**.

**Ministry of Energy and Resources** – Saskatchewan’s Ministry of Energy and Resources (ER) that regulates the upstream oil and gas industry.

**Remote Terminal Unit** – A microprocessor-controlled electronic device that captures and transmits readings from the physical world, such as flow rate, temperature or pressure. Usually, an extension of a SCADA or DCS system; communicates with a host and can operate independently and measure flow parameters, perform calculations, digital control, PID control, etc.

**Relative Density** – Is the ratio of the density (mass of a unit volume) of a substance to the density air or water at standard conditions. Specific gravity is an equivalent term.

**Representative Flow** – Used when stabilized flow is not achievable, such as for wells with artificial lift systems and wells with slugging characteristics. The test volumes of gas, condensate, or water must be representative of the well’s production capability under normal operating conditions.

**Resistance Temperature Devices** – Sensors used to measure temperature by correlating the electrical resistance of the sensor with temperature.

**Return Gas** – Any gas coming back to the battery it was produced from, from a gas plant or other facility after sweetening, or processing.

**Royalty** – For the purpose of this Directive, includes Crown royalties payable pursuant to *The Crown Oil and Gas Royalty Regulations, 2012* and freehold production taxes payable pursuant to *The Freehold Oil and Gas Production Tax Regulations, 2012* and royalties paid to freehold mineral owners. Also see Common Crown and common Freehold Royalty

**RTD** – see **Resistance Temperature Devices**.

**RTU** – see **Remote Terminal Unit**.

**S&W** – see **Sediment and Water**.

**Sales Gas** – A mixture mainly of methane originating from raw gas, if necessary, through the processing of the raw gas for the removal or partial removal of some constituents, and that meets specifications for use as a domestic, commercial, or industrial fuel or as an industrial raw material.

**Satellite** – Surface equipment located between a number of wells and the main battery that is intended to separate and measure the production from each well during the well tests, after which the fluids are recombined and piped to the main battery for separation, treating, measurement, and storage or delivery.

**SCADA** – **Supervisory Control and Data Acquisition System**. A system for gathering and sending coded signals over communication channels to provide control of remote equipment. Typically, the remote equipment is an RTU – see Remote Terminal Unit. See also Electronic Flow Measurement.

**Secondary Element** – See Meter Element.

**Secondary Measurement** – An alternative measurement or calculation method used to determine a volume which is used during repair or downtime of the primary measurement equipment.

**Sediment and Water** – The amount of settled solid and semi-solid components and water in oil.

**Semi-annually** – Once every two calendar quarters.

**Separator** – An unfired apparatus specifically designed and used for separating fluids produced from a well into two or more streams, but does not include a dehydrator.

**Shrinkage** – Refers to a volume reduction associated with one or both of the following two processes:

1. Blending (blending shrinkage) of hydrocarbon streams of varying density such as heavy oil and condensate.
2. Loss of volatile components through vapourization such as flashing (flashing shrinkage) or weathering due to a pressure reduction and/or temperature increase or to continued exposure to atmospheric conditions such as conversion of live oil to base conditions., and/or
3. Acid gas (acid gas shrinkage) converted to elemental sulphur at a sulphur recovery plant

**Shrinkage Factor** – A factor that accounts for a volume reduction due to the blending of liquid hydrocarbons or flashing of gas in solution from a liquid.

**Single Point Measurement Uncertainty** – The uncertainty or accuracy of the equipment and/or procedures used to determine a specific volume at a single measurement point.

**Single-Well Battery** – A licensed well that treats production exclusively from that licensed well.

(Source: *The Oil and Gas Conservation Regulations, 2012*)

**Site** – The area defined by the boundaries of a surface lease for an upstream oil and gas facility or well site and the equipment related to that surface lease.

**Sour Gas** – Sour gas is natural gas (including solution gas) containing more than trace amounts of hydrogen sulfide (H<sub>2</sub>S).

**Southwestern Saskatchewan** – Gas wells located south of Township 28 and West of the Third Meridian in Saskatchewan with completions within the stratigraphic units from the base of the Glacial Drift to the base of the Upper Cretaceous.

**Split Load** – When a truck takes on partial loads from more than one well or battery in a single trip or when load fluids are delivered to more than one receipt point or well.

**Stabilized Flow** – A point at which flowing parameters of gas, condensate, or water are producing under normal operating conditions and represent production level equal to the well's normal average flow rate. See Section 7.1.2 and 7.2.3.

**Steam** – The vapour into which water is changed when heated to its' boiling point. For reporting to Petrinex, the sum of all steam injection volumes of varying quality, reported as a cold water equivalent volume.

**Steam Quality** – The measure of the amount of saturated steam in the vapour phase (mass fraction).

**Stock Tank Vapours** – Gas in a storage tank that has been released from the liquid it was entrained in.

**SWB** – see **Single-Well Battery**

**Sweet Gas** – Natural gas that contains little or no hydrogen sulfide.

**Sweet Oil** – Oil that contains little or no hydrogen sulfide.

**Synthetic Crude Oil** - A mixture, mainly of pentanes and heavier hydrocarbons, that may contain sulphur compounds, that is derived from the processing of heavy oil and that is liquid at the conditions under which its volume is metered or estimated.

**Tertiary Element** – See Meter Element.

**Thermal (Recovery/Production)** – Refers to the use of heat (generally in the form of steam) to lower the viscosity of heavy oil in the reservoir to the point it can be made to flow out of the production well.

**Treater** – A fired apparatus specifically designed and used for separating gas and water from crude oil.

**Triennially** – Once every twelve calendar quarters (once every three years).

**Uncertainty** – The expected range of the true value, given a measurement. It can be expressed as a percentage ( $\pm 1.0\%$ ) or as an absolute measurement ( $\pm 1.0$  Metre). Uncertainty is different from accuracy in that the true value is not known, only the measurement. Also, it is expressed with a specific confidence level (e.g. 95.0%).

**Unique Well Identifier** – Is the standard well identification that was developed for the petroleum industry by the Geoscience Data Committee of the Canadian Petroleum Association (CPA) and has been adopted by ER. It consists of 16 characters, which make up four basic components:

- a. legal survey location
- b. survey system code
- c. location exception code
- d. event sequence code

Together these define the approximate geographical location of the bottom of a drill hole and a specific drilling or producing event at the drill hole.

The unique well identifier, although based on the legal survey position of a well, is primarily for identification rather than location. The location component describes the bottomhole location of the well, not the surface position of the well.

**UWI** – see **Unique Well Identifier**.

**Vent Gas** – Uncombusted gas that is released to the atmosphere at upstream oil and gas operations.

Vent gas includes:

- Blanket gas;
- Facility upsets and emergency shutdown;
- Fugitive emissions;
- Gas from compressor seals, starters, and blowdowns;
- Gas from dehydrator still columns;
- Gas from production tanks, not including methanol and chemical tanks;
- Gas produced during well completions;
- Gas produced during well unloading volumes;
- Gas released during pigging operations;
- Gas used to operate pneumatic devices; and
- Waste gas.

**Verification** – Procedures that establish the accuracy of the “as found” values indicated by a measuring device as compared to the values indicated by a reference standard.

**Water** - For the purpose of this Directive, when the term ‘water’ is used it applies to all types of water such as Water, Fresh Water, and Brine.

**Water Cut** – The ratio of the relative water to oil fractions in a liquid sample or stream. For production determination solids are included in the water volume.

**Water to Gas Ratio** – A ratio calculated by dividing the water volume by the gas volume of a well test. See [Table 7.1](#). for calculation and rounding requirements.

**Well** – means

(i) any opening in the ground made within Saskatchewan from which any oil, gas, oil and gas or other hydrocarbon is, has been or is capable of being produced from a reservoir;

(ii) any opening in the ground that is made for the purpose of:

(A) obtaining water to inject into an underground formation;

(B) injecting any substance into an underground formation;

(C) storing oil, gas or other hydrocarbons underground; or

(D) monitoring reservoir performance and obtaining geological information; or

(iii) any opening in the ground made for informational purposes pursuant to *The Subsurface Mineral Conservation Regulations*;

but does not include seismic shot holes;

(Source: *The Oil and Gas Conservation Regulations, 2012*)

**WGR** – see **Water-Gas Ratio**.

**Working Interest Participant** - A person who owns a legal or beneficial interest in a well or facility pursuant to an agreement that relates to the ownership of the well or facility.

(Source: *The Oil and Gas Conservation Regulations, 2012*)

## Appendix 3 Water-Cut (S&W) Procedures

Water-cut procedures are divided into three categories and described on the following pages. Different procedures are specified for the three categories to improve accuracy and consistency of the S&W determinations. The use of mason jars with measuring tape attached is not acceptable for determining S&W. S&W percentage must be recorded to a minimum of one decimal place.

More information on S&W determination is in API MPMS, Chapter 10.4: Determination of Water and/or Sediment in Crude Oil by the Centrifuge Method (Field Procedure). ER will consider any procedure that meets API MPMS, 10.4 standards to be in compliance with this Directive. It is the responsibility of the licensee to show that its procedure meets the above API standard.

### Category 1: Water-cut $> 0.0\% \leq 10.0\%$

Obtain a representative sample of liquid.

Shake the sample container vigorously to mix it before pouring into the centrifuge tubes.

1. For each of the two tubes, add exactly 50 ml of the sample.
2. For each of the two tubes, add the solvent solution, consisting of premixed solvent and demulsifier to the 100 ml mark.
3. Stopper each tube tightly and invert 10 times.
4. Loosen the stoppers and immerse the tubes in a preheater. Heat the contents to 60.0°C within  $\pm 3.0^\circ\text{C}$ .
5. Stopper each tube tightly and invert 10 times.
6. Place the tubes in the centrifuge machine in a balanced condition and spin for 5 minutes.
7. Immediately after the centrifuge comes to rest, use a thermometer to verify that the sample temperature is within 9.0°C of the test temperature.

If sample temperature is within 9.0°C, go to step 8. If sample temperature is not within 9.0°C, go back to step 4, raise the temperature, and repeat steps 5, 6, and 7.

8. Read and record the volume of water and sediment at the bottom of each tube.
9. Reheat the tubes to the initial spin temperature and return them, without agitation, to the centrifuge machine. Spin for an additional 5.0 minutes. Repeat the procedure until two consecutive, consistent readings are obtained.

For the test to be considered valid, a clear interface must be observed between the oil layer and the separated water. No emulsion should be present immediately above the oil/water interface. A test comprises two tubes of the same sample. Compare the readings of the two tubes. If the difference is greater than one subdivision on the centrifuge tube, the test is invalid and should be repeated.

10. Calculation and reporting:

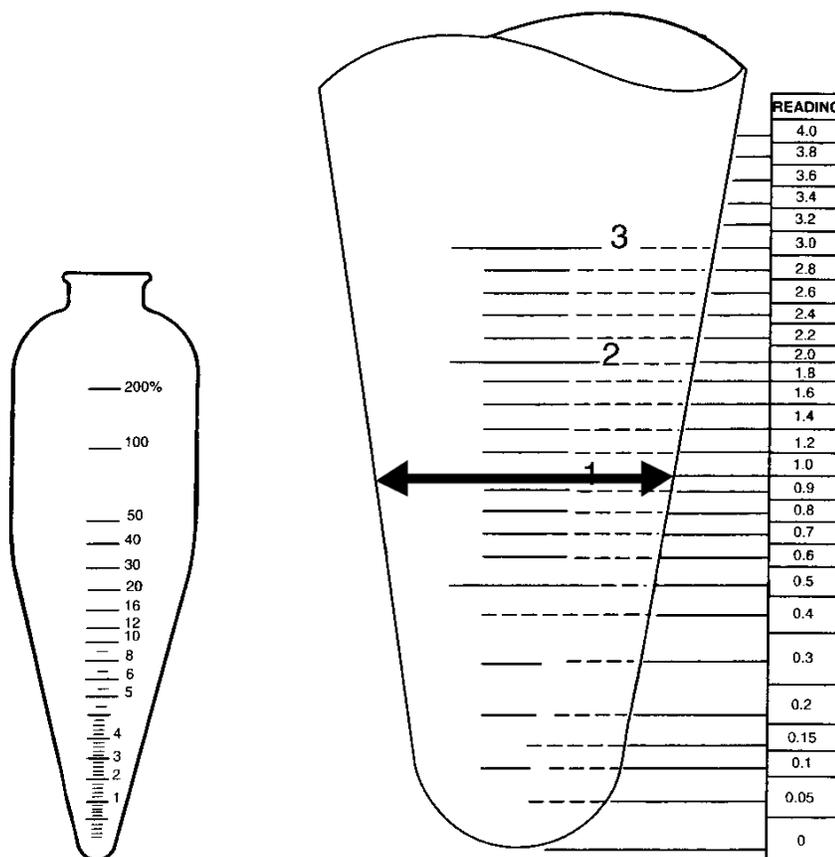
For 200.0 ml tubes: the percentage of water and sediment is the average, to three decimal places, of the values read directly from the two tubes.

For 100 ml tubes: read and record the volume of water and sediment in each tube. Add the readings together and report the sum as the percentage of water and sediment.

**Example 1 (see Figure A4.1)**

100.0 ml centrifuge tubes	200.0 ml centrifuge tubes
If reading from each tube is the same:	
Reading from each tube = 0.50 ml	Reading from each tube = 1.00 ml
Water cut = $(0.50 + 0.50) \div 100 = 1.0\%$	Water cut = $1.00 \div 100 = 1.0\%$
If reading from each tube is not the same:	
Reading from 1 <sup>st</sup> run of each tube = 0.50, 0.60 ml	Reading from 1 <sup>st</sup> run of each tube = 1.00, 1.05 ml
Reading from 2 <sup>nd</sup> run of each tube = 0.50, 0.55 ml	Reading from 2 <sup>nd</sup> run of each tube = 1.00, 1.10 ml
Water cut = $(0.50 + 0.60 + 0.50 + 0.55) \div 2 \div 100 = 1.1\%$	Water cut = $(1.00 + 1.05 + 1.00 + 1.10) \div 4 \div 100 = 1.0\%$

**Figure A4.1. Water-cut > 0.0% ≤ 10.0%**



**Category 2: Water-cut > 10.0% ≤ 80.0%**

Obtain the maximum representative sample of liquid feasible. The representative sample must be a minimum of 800.0 ml.

Transfer the entire sample into an adequately sized graduated cylinder. It may be necessary to wash out the inside of the sample container with a measured volume of solvent to ensure that all of the oil is removed. If this is done, it is necessary to account for the additional amount of solvent added when calculating the water-cut percentage.

Place the graduated cylinder into a heat bath:

1. at or above treater temperature
- or
2. at or above 60.0°C if no treater is involved

until the sample temperature and free water fallout have stabilized. A clear oil/water interface must be visible.

Read and record the total volume, the volume of free water, and the volume of oil/emulsion in the graduated cylinder. Calculate the free water percentage as follows:

$$\text{Percentage of free water} = (\text{Volume of free water} \div \text{Total volume}) \times 100.0\%$$

If solvent and/or demulsifier is added to the sample at any stage of this procedure, it must be accounted for in the calculation as follows:

$$\text{Percentage of free water} = \text{Volume of free water} \div (\text{Total volume} - \text{Volume of solvent/demulsifier}) \times 100.0\%$$

Draw 100.0 ml from the oil/emulsion portion in the graduated cylinder and for each of the two 100.0 ml centrifuge tubes add the oil/emulsion to exactly the 50.0 ml mark. Add solvent to bring the level in the tubes to exactly the 100.0 ml mark. The procedures previously outlined for samples with 0.0 per cent to 10.0 per cent water cut are to be followed, with the exception that the water-cut readings from both tubes are to be added together, even if they are not the same.

Note that if 200.0 ml tubes are to be used, a larger initial sample will be required, and if the water-cut readings from both tubes are not the same, the average of both tubes is to be used as the resultant water cut of the oil/emulsion portion.

From the spinning results, calculate the percentage of water remaining in the oil/emulsion portion as follows:

$$\text{Percentage of water remaining} = \frac{\text{Total oil/emulsion volume in cylinder} \times \text{Water-cut \% of oil/emulsion}}{\text{Total volume}}$$

If solvent and/or demulsifier is added to the sample at any stage of this procedure, it must be accounted for in the calculation as follows:

$$\text{Percentage of water remaining} = \frac{\text{Total oil/emulsion volume in cylinder} \times \text{Water-cut \% of oil/emulsion}}{\text{Total volume} - \text{Volume of solvent/demulsifier}}$$

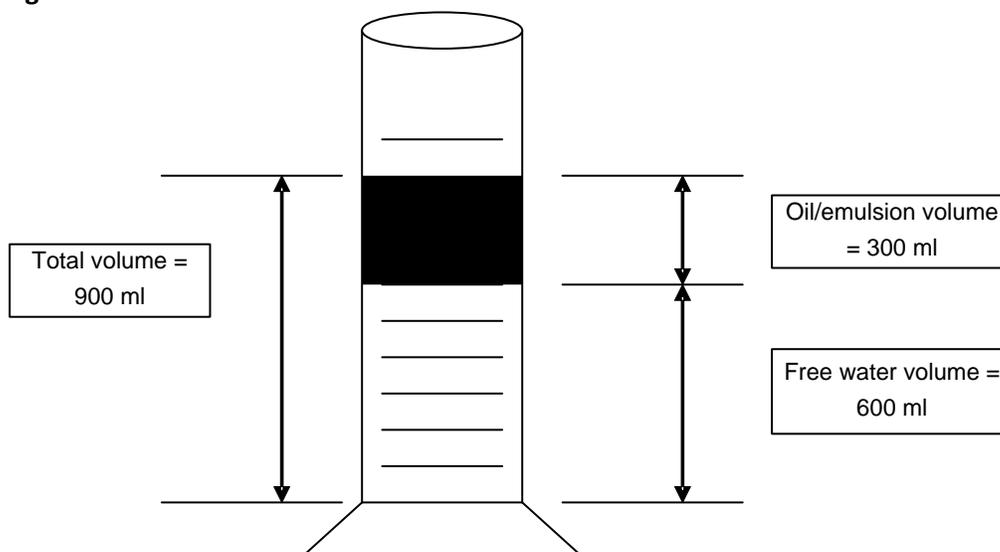
Calculate the total water-cut percentage as follows:

$$\text{Total water-cut \%} = \% \text{ free water} + \% \text{ water remaining}$$

**Example 2 (see Figure A4.2)**

1000.0 ml graduated cylinder
% of free water = $600.0 \text{ ml} \div 900.0 \text{ ml} \times 100\% = 66.7\%$
% of water remaining = $300.0 \text{ ml} \times 10\%^1 \div 900.0 \text{ ml} = 3.3\%$
Total water-cut % = $66.7\% + 3.3\% = 70.0\%$

<sup>1</sup> Water cut of oil portion determined by spinning samples

**Figure A4.2. Water-cut > 10.0% ≤ 80.0%****Category 3: Water-cut > 80.0% ≤ 100.0%**

Obtain the maximum representative sample of liquid feasible. The representative sample must be a minimum of 800.0 ml.

Transfer the entire sample into an adequately sized graduated cylinder. It may be necessary to wash out the inside of the sample container with a measured volume of solvent to ensure that all of the oil is removed. If this is done, it is necessary to account for the additional amount of solvent added when calculating the water-cut percentage.

Place the graduated cylinder into a heat bath

1. at or above treater temperature; or
2. at or above 60.0°C if no treater is involved.

until the sample temperature and free water fallout have stabilized. A clear oil/water interface must be visible. A narrow-necked graduated cylinder should be used to improve accuracy in sample measurement when the water cut is > 90.0 per cent. Refer to [Figure A4.4](#) for an example.

Read and record the total volume and the volume of free water in the graduated cylinder. If no solvent or demulsifier has been added to the sample, calculate the water-cut percentage as follows:

$$\text{Water-cut \%} = \text{Volume of free water} \div \text{Total volume} \times 100.0\%$$

If solvent and/or demulsifier is added to the sample at any stage of this procedure, it must be accounted for in the calculation as follows:

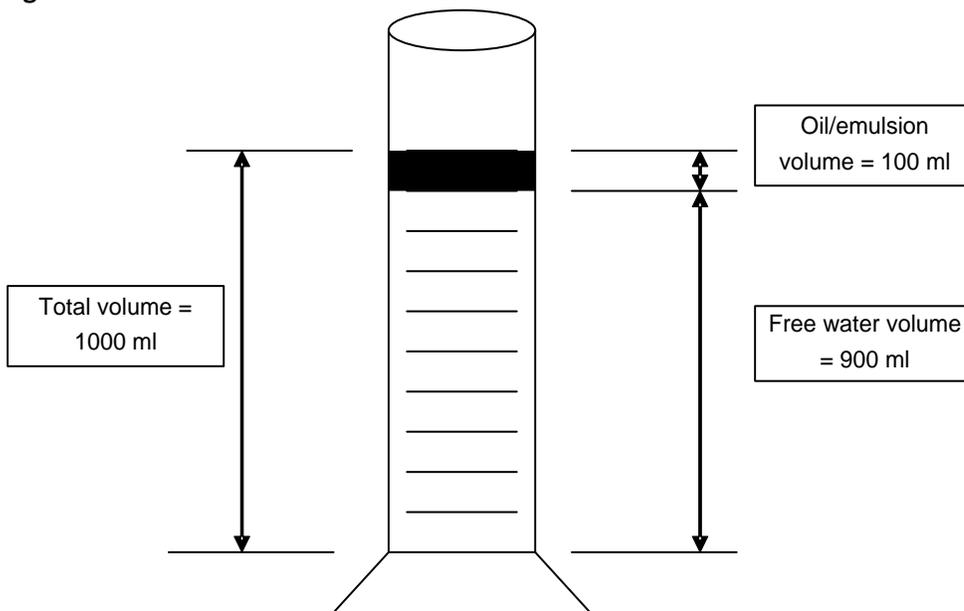
$$\text{Water-cut \%} = \text{Volume of free water} \div (\text{Total volume} - \text{Volume of solvent/demulsifier}) \times 100.0\%$$

The water content of the oil/emulsion portion in the graduated cylinder does not have to be determined, due to the limited amount of the oil/emulsion portion of the sample available at these high water contents. However, if there is enough oil/emulsion volume, the licensee may choose to use the same procedure as that described for the 10.0 per cent to 80.0 per cent S&W, with the option to centrifuge only one sample.

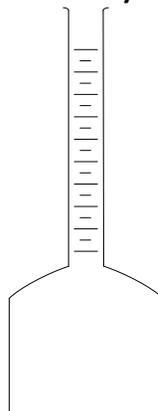
**Example 3 (see Figure A4.3)**

1000.0 ml graduated cylinder
Water-cut % = 900.0 ml ÷ 1000.0 ml x 100%
= 90.0%

**Figure A4.3. Water-cut > 80.0% ≤ 100.0%**



**Figure A4.4. Narrow-necked graduated cylinder**



## Appendix 4 On-site Analytical Techniques for H<sub>2</sub>S Measurement

Refer to Section 11 for related requirements.

On-site measurement of H<sub>2</sub>S in natural gas streams can be accomplished by several different methods. The appropriate method should be selected with an understanding of the benefits and limitations of each method.

### Length of Stain Tubes (GPA Standard 2377-05)

For concentrations below 1500 ppm, the most convenient and economical choice is the use of a length of stain tube. These devices can suffer from some interference, affecting both the precision and the accuracy of the measurements. Nonetheless, for many purposes this technique can provide H<sub>2</sub>S measurements of a suitable quality. The understanding is that the measurement uncertainty is potentially less than the risk of H<sub>2</sub>S degradation if a laboratory method were employed. If the most accurate measurements are required, a second sample can be collected in a suitably inert container and returned to a laboratory for prompt analysis.

### Tutweiler Titration (GPA Standard C-1)

This technique is the method of choice for on-site analysis when the concentration of H<sub>2</sub>S is > 1,500 ppm. The Tutweiler titration can provide accurate measurements of H<sub>2</sub>S using suitably calibrated glassware and chemicals. Operator skill and proper recording of temperatures and atmospheric pressure are also key elements for this technique.

### Instrumental (in-lab) Analytical Techniques for H<sub>2</sub>S Measurement

*Gas Chromatography with Sulphur Selective Detection (ASTM D-5504-01):* Sulphur selective detectors can be coupled with gas chromatographs to achieve a low detection limit for H<sub>2</sub>S and other sulphur compounds, such as mercaptans, sulphides, and disulphides. These instruments are ideal for low concentrations ranging from sub ppm up to several thousand ppm. The sulphur selective detectors are much less susceptible to hydrocarbon interferences and can also identify other sulphur-containing compounds in addition to H<sub>2</sub>S. Suitable sulphur selective detectors are sulphur chemiluminescence detectors (SCD) and pulsed flame photometric detectors (PFPD).

*Gas Chromatography with Thermal Conductivity Detection:* Thermal conductivity detectors can be coupled with gas chromatographs to analyze for intermediate to high levels of H<sub>2</sub>S. H<sub>2</sub>S can be adequately resolved from hydrocarbon components to allow for specific detection. The columns selected for this type of analysis must offer a good balance between high resolution (specificity of H<sub>2</sub>S) and low adsorption of H<sub>2</sub>S. Detection limits for H<sub>2</sub>S levels as low as 300 ppm can be achieved under the right conditions, and the method can also be calibrated for values approaching 100.0 per cent H<sub>2</sub>S. The analytical range for these systems should not exceed the linear range of the column and detector combination. Therefore, acceptable calibration ranges must yield a linear calibration curve (minimum 4 points) with an R<sup>2</sup>-value ≥ 0.99.

## Appendix 5 Gas Equivalent Volume Determination

Liquid Analysis Example (GPA Standard 2145-09)			
Component	Volume Fractions	Mole Fractions	Mass Fractions
N <sub>2</sub>	0.0006	0.0019	0.0008
CO <sub>2</sub>	0.0081	0.0158	0.0109
H <sub>2</sub> S	0	0	0
C <sub>1</sub>	0.0828	0.1617	0.0405
C <sub>2</sub>	0.1117	0.1462	0.0687
C <sub>3</sub>	0.1275	0.1533	0.1056
IC <sub>4</sub>	0.0394	0.0398	0.0362
NC <sub>4</sub>	0.0891	0.0935	0.0849
IC <sub>5</sub>	0.0483	0.0436	0.0492
NC <sub>5</sub>	0.0540	0.0493	0.0556
C <sub>6</sub>	0.0765	0.0614	0.0835
C <sub>7</sub>	0.0880	0.0678	0.1054
C <sub>8</sub>	0.0827	0.0589	0.1032
C <sub>9</sub>	0.0570	0.0368	0.0726
C <sub>10</sub>	0.0363	0.0222	0.0480
C <sub>11</sub>	0.0225	0.0131	0.0305
C <sub>12+</sub>	0.0755	0.0347	0.1044
<b>TOTAL</b>	1.0000	1.0000	1.0000

Properties of C <sub>5+</sub> & C <sub>7+</sub> portion of sample					
	Mol. Fractions	Wt. Fractions	Liq. Vol. Fractions	Mol. Wt. (kg/kmol)	Absolute Density (AD) (kg/m <sup>3</sup> )
C <sub>5+</sub>	0.3878	0.6524	0.5408	107.7	739.33
C <sub>7+</sub>	0.2335	0.4641	0.3620	127.2	785.29

### A5.1 Gas Equivalent Factor by Volume Fraction Calculation

**GEF = Total (Pseudo 10<sup>3</sup>m<sup>3</sup> Gas / m<sup>3</sup> Liquid)**

Condensate Stream: \_\_\_\_\_

Component	Vol. Fraction Liquid Analysis		10 <sup>3</sup> m <sup>3</sup> Gas / m <sup>3</sup> Liquid @ 101.325 kPa & 15.0°C		Pseudo 10 <sup>3</sup> m <sup>3</sup> Gas / m <sup>3</sup> Liquid
N <sub>2</sub>		x	0.68040	=	
CO <sub>2</sub>		x	0.44120	=	
H <sub>2</sub> S		x	0.55460	=	
C <sub>1</sub>		x	0.44217	=	
C <sub>2</sub>		x	0.28140	=	
C <sub>3</sub>		x	0.27213	=	
IC <sub>4</sub>		x	0.22902	=	
NC <sub>4</sub>		x	0.23768	=	
IC <sub>5</sub>		x	0.20485	=	
NC <sub>5</sub>		x	0.20667	=	
C <sub>6</sub>		x	0.18217	=	
C <sub>7</sub>		x	0.16235	=	
C <sub>8</sub>		x	0.14620	=	
C <sub>9</sub>		x	0.13310	=	
C <sub>10</sub>		x	0.12203	=	
				<b>Total =</b>	

**Note: For C<sub>5+</sub>, C<sub>6+</sub>, or C<sub>7+</sub> Sample:**

Properties of C<sub>5+</sub>, C<sub>6+</sub>, or C<sub>7+</sub> sample @ 15.0°C

AD =

Mol. Wt. =

10<sup>3</sup>m<sup>3</sup> Gas / m<sup>3</sup> Liquid = 23.645 (m<sup>3</sup>/kmol) x AD (kg/m<sup>3</sup>) / Mol. Wt. (kg/kmol)

=  Input this factor to the table above for C<sub>5+</sub>, C<sub>6+</sub>, or C<sub>7+</sub>

**GEF = Total (10<sup>3</sup>m<sup>3</sup> Gas / m<sup>3</sup> Liquid)**

=  (10<sup>3</sup>m<sup>3</sup> Gas / m<sup>3</sup> Liquid)

**Example 1 – Gas Equivalent Factor by Volume Fraction Calculation**

GEF = Total (Pseudo 10<sup>3</sup>m<sup>3</sup> Gas / m<sup>3</sup> Liquid)

Condensate Stream: \_\_\_\_\_

Component	Volume Fraction Liquid Analysis		10 <sup>3</sup> m <sup>3</sup> Gas / m <sup>3</sup> Liquid @ 101.325 kPa & 15.0°C		Pseudo 10 <sup>3</sup> m <sup>3</sup> Gas / m <sup>3</sup> Liquid
N <sub>2</sub>	0.0006	x	0.68040	=	0.0004
CO <sub>2</sub>	0.0081	x	0.44120	=	0.0036
H <sub>2</sub> S	0	x	0.55460	=	
C <sub>1</sub>	0.0828	x	0.44217	=	0.0366
C <sub>2</sub>	0.1117	x	0.28140	=	0.0314
C <sub>3</sub>	0.1275	x	0.27213	=	0.0347
IC <sub>4</sub>	0.0394	x	0.22902	=	0.0090
NC <sub>4</sub>	0.0891	x	0.23768	=	0.0212
IC <sub>5</sub>	0.0483	x	0.20485	=	0.0099
NC <sub>5</sub>	0.0540	x	0.20667	=	0.0112
C <sub>6</sub>	0.0765	x	0.18217	=	0.0139
C <sub>7+</sub>	0.3620	x	0.14598	=	0.0528
		x		=	
	1.0000		<b>Total =</b>		0.2247

**Note: For C<sub>7+</sub> Sample:**

Properties of C<sub>7+</sub> sample @ 15.0°C

AD =	785.29
Mol. Wt. =	127.2

10<sup>3</sup>m<sup>3</sup> Gas / m<sup>3</sup> Liquid = 23.645 (m<sup>3</sup>/kmol) x AD (kg/m<sup>3</sup>) / Mol. Wt. (kg/kmol) / 1000 (m<sup>3</sup>/10<sup>3</sup> m<sup>3</sup>)

=  Input this factor to the table above for C<sub>7+</sub>

GEF = Total (Pseudo 10<sup>3</sup>m<sup>3</sup> Gas / m<sup>3</sup> Liquid)

=  (10<sup>3</sup>m<sup>3</sup> Gas / m<sup>3</sup> Liquid)

### A5.2 Gas Equivalent Factor by Mole Fraction Calculation

$$GEF = 23.645 \text{ (m}^3\text{/kmol)} / \text{Total (Pseudo m}^3\text{/kmol)} / 1000 \text{ (m}^3\text{/10}^3\text{ m}^3\text{)}$$

Condensate Stream: \_\_\_\_\_

Component	Mol. Fraction Liquid Analysis		10 <sup>3</sup> m <sup>3</sup> /kmol @ 101.325 kPa & 15.0°C		Pseudo 10 <sup>3</sup> m <sup>3</sup> /kmol
N <sub>2</sub>		x	0.034753	=	
CO <sub>2</sub>		x	0.053590	=	
H <sub>2</sub> S		x	0.042630	=	
C <sub>1</sub>		x	0.053475	=	
C <sub>2</sub>		x	0.084027	=	
C <sub>3</sub>		x	0.086888	=	
IC <sub>4</sub>		x	0.103250	=	
NC <sub>4</sub>		x	0.099482	=	
IC <sub>5</sub>		x	0.115420	=	
NC <sub>5</sub>		x	0.114410	=	
C <sub>6</sub>		x	0.129800	=	
C <sub>7</sub>		x	0.1456640	=	
C <sub>8</sub>		x	0.161730	=	
C <sub>9</sub>		x	0.177650	=	
C <sub>10</sub>		x	0.193770	=	
				<b>Total =</b>	

**Note:**

Properties of C<sub>5+</sub>, C<sub>6+</sub>, or C<sub>7+</sub> Sample @ 15.0°C:

AD =

Mol. Wt. =

**For C<sub>5+</sub>, C<sub>6+</sub>, or C<sub>7+</sub> Sample:**

$$10^3\text{m}^3\text{/kmol} = \text{Mol. Wt.} / \text{AD}$$

=

$$GEF = 23.645 \text{ (m}^3\text{/kmol)} / \text{Total (Pseudo m}^3\text{/kmol)} / 1000 \text{ (m}^3\text{/10}^3\text{ m}^3\text{)}$$

=  (10<sup>3</sup>m<sup>3</sup> Gas / m<sup>3</sup> Liquid)

**Example 2: Gas Equivalent Factor by Mole Fraction Calculation**

GEF = 23.645 (m<sup>3</sup>/kmol) / Total (Pseudo m<sup>3</sup>/kmol) / 1000 (m<sup>3</sup>/10<sup>3</sup> m<sup>3</sup>)

Condensate Stream: \_\_\_\_\_

Component	Mol. Fraction Liquid Analysis		m <sup>3</sup> /kmol @ 101.325 kPa & 15.0°C		Pseudo m <sup>3</sup> /kmol
N <sub>2</sub>	0.0019	x	0.034753	=	0.0001
CO <sub>2</sub>	0.0158	x	0.053590	=	0.0008
H <sub>2</sub> S	0	x	0.042630	=	0
C <sub>1</sub>	0.1617	x	0.053475	=	0.0086
C <sub>2</sub>	0.1462	x	0.084027	=	0.0123
C <sub>3</sub>	0.1533	x	0.086888	=	0.0133
IC <sub>4</sub>	0.0398	x	0.103250	=	0.0041
NC <sub>4</sub>	0.0935	x	0.099482	=	0.0093
IC <sub>5</sub>	0.0436	x	0.115420	=	0.0050
NC <sub>5</sub>	0.0493	x	0.114410	=	0.0056
C <sub>6</sub>	0.0614	x	0.129800	=	0.0080
C <sub>7+</sub>	0.2335	x	0.161980	=	0.0378
		x		=	
		x		=	
	1.0000			<b>Total =</b>	0.1049

**Note:**

Properties of C<sub>7+</sub> sample @ 15.0°C:

AD =	785.29
Mol. Wt. =	127.2

**For C<sub>7+</sub> Fraction:**

m<sup>3</sup>/kmol = Mol. Wt. / AD

=  Input this factor to the table above for C<sub>7+</sub>

$$\text{GEF} = 23.645 \text{ (m}^3\text{/kmol)} / \text{Total (Pseudo m}^3\text{/kmol)} / 1000 \text{ (m}^3\text{/10}^3 \text{ m}^3\text{)}$$

$$= \boxed{0.2254} \text{ (10}^3\text{m}^3 \text{ Gas} / \text{ m}^3 \text{ Liquid)}$$

### A5.3 Gas Equivalent Factor by Mass Fraction Calculation

Step 1. Calculate Pseudo Volume (L) = Mass Fraction ÷ Liquid Density x 1000.0 L/m<sup>3</sup>

Step 2. Calculate Volume Fraction = Component Pseudo Volume ÷ Total Pseudo Volume

Step 3. Calculate Component Pseudo GEF = Volume Fraction x (10<sup>3</sup>m<sup>3</sup> Gas / m<sup>3</sup> Liquid)

Condensate Stream: \_\_\_\_\_

					Step 1	Step 2				Step 3
Component	Mass Fraction Liquid Analysis		Liquid Density (kg/m <sup>3</sup> )		Pseudo Volume (L)	Volume Fraction		10 <sup>3</sup> m <sup>3</sup> Gas /m <sup>3</sup> Liquid @ 101.325 kPa & 15.0°C		Pseudo 10 <sup>3</sup> m <sup>3</sup> Gas / m <sup>3</sup> Liquid
N <sub>2</sub>		÷	806.10	=			x	0.68040	=	
CO <sub>2</sub>		÷	821.20	=			x	0.44120	=	
H <sub>2</sub> S		÷	799.40	=			x	0.55460	=	
C <sub>1</sub>		÷	300.00	=			x	0.44217	=	
C <sub>2</sub>		÷	357.85	=			x	0.28140	=	
C <sub>3</sub>		÷	507.50	=			x	0.27213	=	
IC <sub>4</sub>		÷	562.95	=			x	0.22902	=	
NC <sub>4</sub>		÷	584.25	=			x	0.23768	=	
IC <sub>5</sub>		÷	625.08	=			x	0.20485	=	
NC <sub>5</sub>		÷	630.62	=			x	0.20667	=	
C <sub>6</sub>		÷	663.92	=			x	0.18217	=	
C <sub>7</sub>		÷	688.02	=			x	0.16235	=	
C <sub>8</sub>		÷	706.30	=			x	0.14620	=	
C <sub>9</sub>		÷	721.97	=			x	0.13310	=	
C <sub>10</sub>		÷	734.30	=			x	0.12203	=	
								<b>Total =</b>		

Note: For C<sub>5+</sub>, C<sub>6+</sub>, or C<sub>7+</sub> Sample:

$$\text{Mol. Wt.} = \boxed{\phantom{000000}} \\ \text{AD} = \boxed{\phantom{000000}} \text{ (kg/m}^3\text{)}$$

$$10^3 \text{m}^3 \text{ Gas} / \text{m}^3 \text{ Liquid} = 23.645 \times \text{AD} / \text{Mol. Wt.} / 1000.0 \text{ m}^3 / 10^3 \text{m}^3$$

$$= \text{[Orange Box]} \text{ Input this factor to the table above for } C_{7+}$$

$$\text{GEF} = \text{Total (Pseudo } 10^3 \text{m}^3 \text{ Gas} / \text{m}^3 \text{ Liquid)}$$

$$= \text{[Yellow Box]} (10^3 \text{m}^3 \text{ Gas} / \text{m}^3 \text{ Liquid})$$

### Example 3: Gas Equivalent Factor by Mass Fraction Calculation

Step 1. Calculate Pseudo Volume (L) = Mass Fraction ÷ Liquid Density x 1000.0 L/m<sup>3</sup>

Step 2. Calculate Volume Fraction = Component Pseudo Volume ÷ Total Pseudo Volume

Step 3. Calculate Component Pseudo GEF = Volume Fraction x (10<sup>3</sup>m<sup>3</sup> Gas / m<sup>3</sup> Liquid)

Condensate Stream: \_\_\_\_\_

Component	Mass Fraction Liquid Analysis		Liquid Density (kg/m <sup>3</sup> )		Step 1 Pseudo Volume (L)	Step 2 Volume Fraction		10 <sup>3</sup> m <sup>3</sup> Gas / m <sup>3</sup> Liquid @ 101.325 kPa & 15.0°C		Step 3 Pseudo 10 <sup>3</sup> m <sup>3</sup> Gas / m <sup>3</sup> Liquid
N <sub>2</sub>	0.0008	÷	806.10	=	0.0010	0.0006	x	0.68040	=	0.0004
CO <sub>2</sub>	0.0109	÷	821.20	=	0.0133	0.0081	x	0.44120	=	0.0036
H <sub>2</sub> S	0	÷	799.40	=			x	0.55460	=	
C <sub>1</sub>	0.0405	÷	300.00	=	0.1350	0.0822	x	0.44217	=	0.0363
C <sub>2</sub>	0.0687	÷	357.85	=	0.1920	0.1169	x	0.28140	=	0.0329
C <sub>3</sub>	0.1056	÷	507.50	=	0.2081	0.1267	x	0.27213	=	0.0345
IC <sub>4</sub>	0.0362	÷	562.95	=	0.0643	0.0391	x	0.22902	=	0.0090
NC <sub>4</sub>	0.0849	÷	584.25	=	0.1453	0.0885	x	0.23768	=	0.0210
IC <sub>5</sub>	0.0492	÷	625.08	=	0.0787	0.0479	x	0.20485	=	0.0098
NC <sub>5</sub>	0.0556	÷	630.62	=	0.0882	0.0536	x	0.20667	=	0.0111
C <sub>6</sub>	0.0835	÷	663.92	=	0.1258	0.0766	x	0.18217	=	0.0140
C <sub>7+</sub>	0.4641	÷	785.29	=	0.5910	0.3598	x	0.14584	=	0.0525
	1.0000				1.6427	1.0000	<b>Total =</b>			0.2251

Note: For C<sub>7+</sub> Sample:

$$\begin{aligned} \text{Mol. Wt.} &= 127.2 \\ \text{AD of C}_{7+} \text{ liquid} &= 785.29 \text{ (kg/m}^3\text{)} \end{aligned}$$

$$10^3 \text{m}^3 \text{ Gas} / \text{m}^3 \text{ Liquid} = 23.645 \times \text{AD} / \text{Mol. Wt.} \times 999.10 / 1000.0 \text{ (m}^3/10^3 \text{m}^3\text{)}$$

GEF = Total (Pseudo 10<sup>3</sup>m<sup>3</sup> Gas / m<sup>3</sup> Liquid)

$$= 0.2251 \text{ (10}^3 \text{m}^3 \text{ Gas} / \text{m}^3 \text{ Liquid)}$$

## Appendix 6 Calculated Compositional Analysis Examples

### Calculated Well Stream Compositional Analysis Example

**Step 1:** Collect volumetric and compositional data for both gas and liquid phases.

Gas		Liquid	
Gas Volume ( $10^3\text{m}^3$ )	10000.0	Liquid Volume ( $\text{m}^3$ )	200.0
Composition	Mole %	Composition	Mole %
N <sub>2</sub>	1.00	N <sub>2</sub>	0.00
CO <sub>2</sub>	2.00	CO <sub>2</sub>	1.00
H <sub>2</sub> S	2.40	H <sub>2</sub> S	2.00
C <sub>1</sub>	80.00	C <sub>1</sub>	3.00
C <sub>2</sub>	8.00	C <sub>2</sub>	4.00
C <sub>3</sub>	3.00	C <sub>3</sub>	7.00
IC <sub>4</sub>	1.00	IC <sub>4</sub>	10.00
NC <sub>4</sub>	1.50	NC <sub>4</sub>	15.00
IC <sub>5</sub>	0.20	IC <sub>5</sub>	7.00
NC <sub>5</sub>	0.50	NC <sub>5</sub>	11.00
C <sub>6</sub>	0.30	C <sub>6</sub>	10.00
C <sub>7+</sub>	0.10	C <sub>7+</sub>	30.00
	100.00		100.00

**Step 2:** Convert the condensate liquid volume to a GEV.

- Convert liquid volume to equivalent gas volume using the condensate gas equivalent factor.

**Equation 1:**  $\text{GEV} = \text{Volume of condensate (m}^3) \times \text{GEF (m}^3 \text{ gas per m}^3 \text{ liquid)}$

$$\text{GEV} = 200.0 \text{ (m}^3) \times 220.12 \text{ (m}^3 \text{ gas per m}^3 \text{ liquid)} \div 1000 \text{ (10}^3\text{m}^3/\text{m}^3) \\ = 44.024 \text{ 10}^3\text{m}^3$$

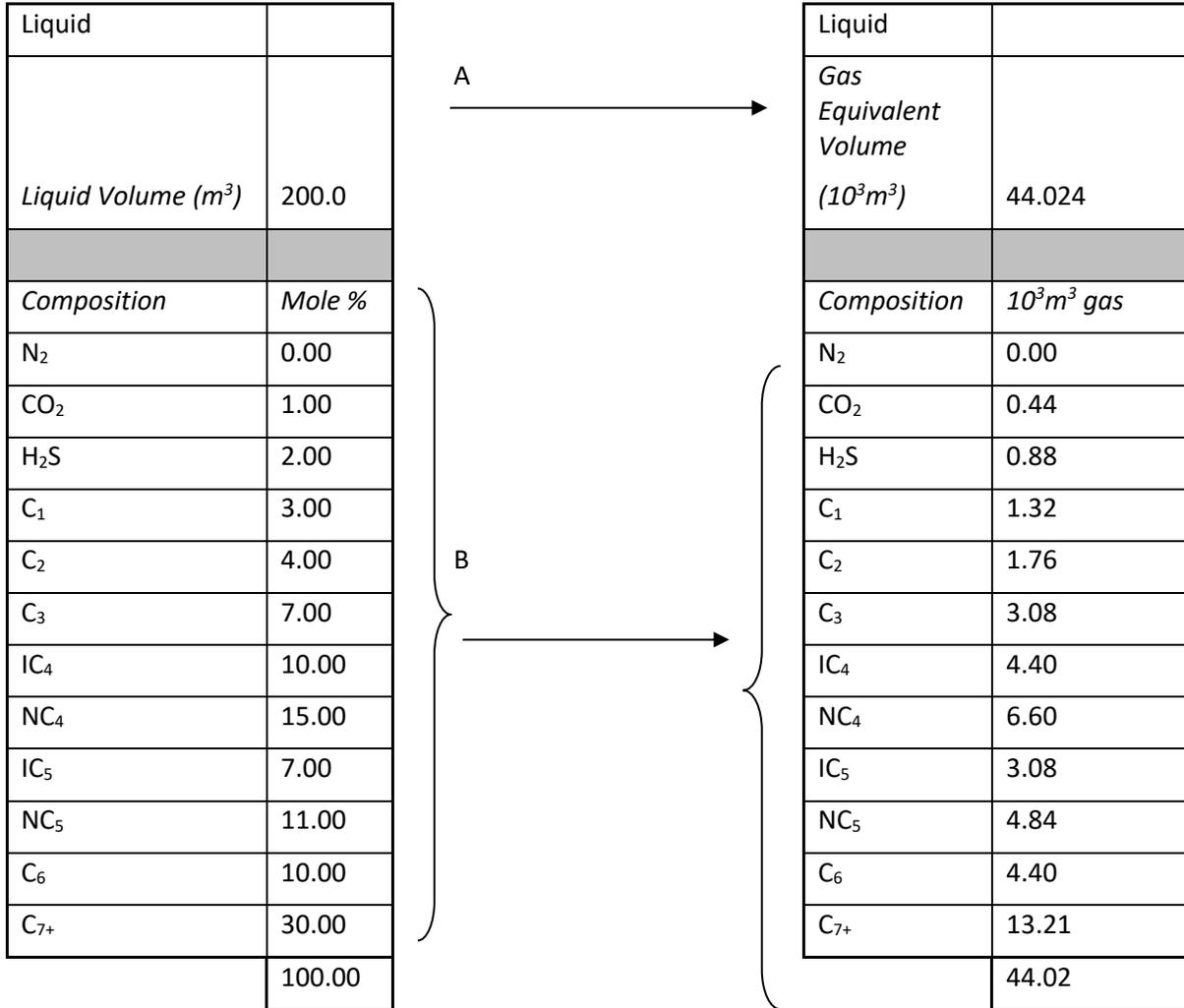
If the gas equivalent factor is not included with the condensate analysis report, it can be calculated.

**Equation 2:**  $\text{GEF} = \text{Absolute Density of Condensate (kg/m}^3 \text{ @ 15.0}^\circ\text{C)} / \text{Molecular weight of the condensate (grams/mole)}$

- Convert the compositional fractions to equivalent gas volumes on a component basis.

**Equation 3:**  $\text{Component gas volume (10}^3\text{m}^3) = [\text{Component mole percent}] \times [\text{GEV}]$

Example: n-pentane equivalent volume:  
 Volume of condensate = 200.0 m<sup>3</sup>  
 Gas Equivalent Factor = 220.12  
 Equivalent n-pentane (NC<sub>5</sub>) gas volume = [11.0%] x [44.024 10<sup>3</sup> m<sup>3</sup>]  
 = 4843.6 m<sup>3</sup>



**Step 3:** Add the gas volumes and liquid gas equivalent volumes and normalize to mole fraction 1.0 or 100.0 per cent.

Gas		Liquid		Recombined Volume		Recombined Composition	
Gas Volume (10 <sup>3</sup> m <sup>3</sup> )	10000.0	Gas Equivalent Volume (10 <sup>3</sup> m <sup>3</sup> )	44.0	Gas Equivalent Volume (10 <sup>3</sup> m <sup>3</sup> )	10044.0	Gas Equivalent Volume (10 <sup>3</sup> m <sup>3</sup> )	10044.0
Composition	10 <sup>3</sup> m <sup>3</sup> gas	Composition	10 <sup>3</sup> m <sup>3</sup> gas	Composition	10 <sup>3</sup> m <sup>3</sup> gas	Composition	Mole %
N <sub>2</sub>	100.0	N <sub>2</sub>	0.00	N <sub>2</sub>	100.0	N <sub>2</sub>	1.00
CO <sub>2</sub>	200.0	CO <sub>2</sub>	0.44	CO <sub>2</sub>	200.4	CO <sub>2</sub>	2.00
H <sub>2</sub> S	240.0	H <sub>2</sub> S	0.88	H <sub>2</sub> S	240.9	H <sub>2</sub> S	2.40
C <sub>1</sub>	8000.0	C <sub>1</sub>	1.32	C <sub>1</sub>	8001.3	C <sub>1</sub>	79.66
C <sub>2</sub>	800.0	C <sub>2</sub>	1.76	C <sub>2</sub>	801.8	C <sub>2</sub>	7.98
C <sub>3</sub>	300.0	C <sub>3</sub>	3.08	C <sub>3</sub>	303.1	C <sub>3</sub>	3.02
IC <sub>4</sub>	100.0	IC <sub>4</sub>	4.40	IC <sub>4</sub>	104.4	IC <sub>4</sub>	1.04
NC <sub>4</sub>	150.0	NC <sub>4</sub>	6.60	NC <sub>4</sub>	156.6	NC <sub>4</sub>	1.56
IC <sub>5</sub>	20.0	IC <sub>5</sub>	3.08	IC <sub>5</sub>	23.1	IC <sub>5</sub>	0.23
NC <sub>5</sub>	50.0	NC <sub>5</sub>	4.84	NC <sub>5</sub>	54.8	NC <sub>5</sub>	0.55
C <sub>6</sub>	30.0	C <sub>6</sub>	4.40	C <sub>6</sub>	34.4	C <sub>6</sub>	0.34
C <sub>7+</sub>	10.0	C <sub>7+</sub>	13.21	C <sub>7+</sub>	23.2	C <sub>7+</sub>	0.23
	10000.0		44.00		10044.0		100.00

### Calculated Group Compositional Analysis Example

**Step 1:**

Collect volumetric and compositional data for both gas and liquid phases for all streams. This information is required for all wells.

**Step 2:**

Mathematically recombine the fluid based on volumetric and compositional data collected in Step 1 for each stream.

**Step 3:**

Add the recombined fluid volumes on a component basis and normalize to 100%.

Stream 1: Gas	
Gas Volume (10 <sup>3</sup> m <sup>3</sup> )	10000
Composition	Mole %
N <sub>2</sub>	1.14
CO <sub>2</sub>	0.16
H <sub>2</sub> S	0.00
C <sub>1</sub>	85.31
C <sub>2</sub>	6.44
C <sub>3</sub>	3.77
IC <sub>4</sub>	0.63
NC <sub>4</sub>	1.32
IC <sub>5</sub>	0.33
NC <sub>5</sub>	0.41
C <sub>6</sub>	0.26
C <sub>7+</sub>	0.23
	100.00

Liquid	
Gas Equivalent Volume (10 <sup>3</sup> m <sup>3</sup> )	800
Composition	Mole %
N <sub>2</sub>	0.12
CO <sub>2</sub>	0.08
H <sub>2</sub> S	0.00
C <sub>1</sub>	22.02
C <sub>2</sub>	6.14
C <sub>3</sub>	8.56
IC <sub>4</sub>	2.62
NC <sub>4</sub>	7.11
IC <sub>5</sub>	3.66
NC <sub>5</sub>	5.73
C <sub>6</sub>	9.73
C <sub>7+</sub>	34.23
	100.00

Recombined Fluid	
Gas Equivalent Volume (10 <sup>3</sup> m <sup>3</sup> )	10800
Composition	Mole %
N <sub>2</sub>	1.06
CO <sub>2</sub>	0.15
H <sub>2</sub> S	0.00
C <sub>1</sub>	80.62
C <sub>2</sub>	6.42
C <sub>3</sub>	4.12
IC <sub>4</sub>	0.78
NC <sub>4</sub>	1.75
IC <sub>5</sub>	0.58
NC <sub>5</sub>	0.80
C <sub>6</sub>	0.96
C <sub>7+</sub>	2.75
	100.00

Stream 2: Gas	
Gas Volume (10 <sup>3</sup> m <sup>3</sup> )	15000
Composition	Mole %
N <sub>2</sub>	1.00
CO <sub>2</sub>	2.00
H <sub>2</sub> S	2.40
C <sub>1</sub>	80.00
C <sub>2</sub>	8.00
C <sub>3</sub>	3.00
IC <sub>4</sub>	1.00
NC <sub>4</sub>	1.50
IC <sub>5</sub>	0.20
NC <sub>5</sub>	0.50
C <sub>6</sub>	0.30
C <sub>7+</sub>	0.10

Liquid	
Gas Equivalent Volume (10 <sup>3</sup> m <sup>3</sup> )	200
Composition	Mole %
N <sub>2</sub>	0.00
CO <sub>2</sub>	1.00
H <sub>2</sub> S	2.00
C <sub>1</sub>	3.00
C <sub>2</sub>	4.00
C <sub>3</sub>	7.00
IC <sub>4</sub>	10.00
NC <sub>4</sub>	15.00
IC <sub>5</sub>	7.00
NC <sub>5</sub>	11.00
C <sub>6</sub>	10.00
C <sub>7+</sub>	30.00

Recombined Fluid	
Gas Equivalent Volume (10 <sup>3</sup> m <sup>3</sup> )	15200
Composition	Mole %
N <sub>2</sub>	0.99
CO <sub>2</sub>	1.99
H <sub>2</sub> S	2.39
C <sub>1</sub>	78.99
C <sub>2</sub>	7.95
C <sub>3</sub>	3.05
IC <sub>4</sub>	1.12
NC <sub>4</sub>	1.68
IC <sub>5</sub>	0.29
NC <sub>5</sub>	0.64
C <sub>6</sub>	0.43
C <sub>7+</sub>	0.49

Total Recombined Fluid	
Gas Equivalent Volume (10 <sup>3</sup> m <sup>3</sup> )	36000
Composition	Mole %
N <sub>2</sub>	0.76
CO <sub>2</sub>	1.44
H <sub>2</sub> S	1.01
C <sub>1</sub>	82.37
C <sub>2</sub>	6.95
C <sub>3</sub>	2.94
IC <sub>4</sub>	0.79
NC <sub>4</sub>	1.37
IC <sub>5</sub>	0.32
NC <sub>5</sub>	0.54
C <sub>6</sub>	0.47
C <sub>7+</sub>	1.04

Stream 3: Gas		Liquid		Recombined Fluid		100.00
Gas Volume (10 <sup>3</sup> m <sup>3</sup> )	10000	Gas Equivalent Volume (10 <sup>3</sup> m <sup>3</sup> )	0	Gas Equivalent Volume (10 <sup>3</sup> m <sup>3</sup> )	10000	
Composition	Mole %	Composition	Mole %	Composition	Mole %	
N <sub>2</sub>	0.10	N <sub>2</sub>	0.00	N <sub>2</sub>	0.10	
CO <sub>2</sub>	2.00	CO <sub>2</sub>	0.00	CO <sub>2</sub>	2.00	
H <sub>2</sub> S	0.00	H <sub>2</sub> S	0.00	H <sub>2</sub> S	0.00	
C <sub>1</sub>	89.40	C <sub>1</sub>	0.00	C <sub>1</sub>	89.40	
C <sub>2</sub>	6.00	C <sub>2</sub>	0.00	C <sub>2</sub>	6.00	
C <sub>3</sub>	1.50	C <sub>3</sub>	0.00	C <sub>3</sub>	1.50	
IC <sub>4</sub>	0.30	IC <sub>4</sub>	0.00	IC <sub>4</sub>	0.30	
NC <sub>4</sub>	0.50	NC <sub>4</sub>	0.00	NC <sub>4</sub>	0.50	
IC <sub>5</sub>	0.08	IC <sub>5</sub>	0.00	IC <sub>5</sub>	0.08	
NC <sub>5</sub>	0.10	NC <sub>5</sub>	0.00	NC <sub>5</sub>	0.10	
C <sub>6</sub>	0.01	C <sub>6</sub>	0.00	C <sub>6</sub>	0.01	
C <sub>7+</sub>	0.01	C <sub>7+</sub>	0.00	C <sub>7+</sub>	0.01	
	100.00		0.00		100.00	

**Calculated Single Compositional Analysis (from Two Samples)**

**Step 1:** Collect spot samples and record the metered volumes associated with each sample.

**Step 2:** Calculate individual component volumes by multiplying the individual component mole fractions or percentage values by the associated metered volumes.

**Example:** Gas Sample #1, Calculation of methane volume

Total Volume = 10,000.0 10<sup>3</sup>m<sup>3</sup>

Methane = 80.00 mole %

Methane Volume = 10,000.0 10<sup>3</sup>m<sup>3</sup> x 0.8000 = 8,000.0 10<sup>3</sup>m<sup>3</sup>

**Step 3:** Normalization: Individual component volumes are summed. The individual component volumes are then be divided into the total to create a normalized (calculated) compositional value.

**Example:** Ethane (C<sub>2</sub>), Calculation of Mole %

Gas Sample #1, C<sub>2</sub> volume: 800.0 10<sup>3</sup>m<sup>3</sup>

Gas Sample #2, C<sub>2</sub> volume: 560.0 10<sup>3</sup>m<sup>3</sup>

Combined, C<sub>2</sub> volume: 1,360.0 10<sup>3</sup>m<sup>3</sup>

Total gas volume: 18,000.0 10<sup>3</sup>m<sup>3</sup>

Calculated C<sub>2</sub> concentration = 1,360.0 10<sup>3</sup>m<sup>3</sup> ÷ 18,000.0 10<sup>3</sup>m<sup>3</sup> = 7.56 mole %

Gas Sample #1			Gas Sample #2			Calculated Single Compositional Analysis		
Gas Volume ( $10^3 m^3$ ) = 10,000.0			Gas Volume ( $10^3 m^3$ ) = 8,000.0			Gas Volume ( $10^3 m^3$ ) = 18,000.0		
Component	Mole %	$10^3 m^3$ gas	Component	Mole %	$10^3 m^3$ gas	Component	<b>Calculated Mole %</b>	$10^3 m^3$ gas
N <sub>2</sub>	1.00	100.0	N <sub>2</sub>	0.60	48.0	N <sub>2</sub>	<b>0.82</b>	148.0
CO <sub>2</sub>	2.00	200.0	CO <sub>2</sub>	2.00	160.0	CO <sub>2</sub>	<b>2.00</b>	360.0
H <sub>2</sub> S	2.40	240.0	H <sub>2</sub> S	1.50	120.0	H <sub>2</sub> S	<b>2.00</b>	360.0
C <sub>1</sub>	80.00	8,000.0	C <sub>1</sub>	83.00	6,640.0	C <sub>1</sub>	<b>81.33</b>	14,640.0
C <sub>2</sub>	8.00	800.0	C <sub>2</sub>	7.00	560.0	C <sub>2</sub>	<b>7.56</b>	1,360.0
C <sub>3</sub>	3.00	300.0	C <sub>3</sub>	2.50	200.0	C <sub>3</sub>	<b>2.78</b>	500.0
IC <sub>4</sub>	1.00	100.0	IC <sub>4</sub>	1.00	80.0	IC <sub>4</sub>	<b>1.00</b>	180.0
NC <sub>4</sub>	1.50	150.0	NC <sub>4</sub>	1.40	112.0	NC <sub>4</sub>	<b>1.46</b>	262.0
IC <sub>5</sub>	0.20	20.0	IC <sub>5</sub>	0.18	14.4	IC <sub>5</sub>	<b>0.19</b>	34.4
NC <sub>5</sub>	0.50	50.0	NC <sub>5</sub>	0.45	36.0	NC <sub>5</sub>	<b>0.48</b>	86.0
C <sub>6</sub>	0.30	30.0	C <sub>6</sub>	0.28	22.4	C <sub>6</sub>	<b>0.29</b>	52.4
C <sub>7+</sub>	0.10	10.0	C <sub>7+</sub>	0.09	7.2	C <sub>7+</sub>	<b>0.10</b>	17.2
	<b>100.00</b>	<b>10,000.0</b>		<b>100.00</b>	<b>8,000.0</b>		<b>100.00</b>	<b>18,000.0</b>

## Appendix 7 Blending Shrinkage Calculation and Reporting Example

### API MPMS, Chapter 12.3, Section 5.3 Equation 3

$$\% \text{ Shrinkage} = 26,900 \times C \times (100 - C)^{0.819} \times (1/dL - 1/dH)^{2.28}$$

Where:

- % Shrinkage = volumetric shrinkage expressed as a % of the total blended mixture volume.
- C = the concentration, in liquid volume %, of the lighter component in the blended mixture.

$$C = \text{Lighter component volume} \div (\text{Heavier component volume} + \text{lighter component volume}) \times 100.$$

- dL = weighted average lighter component density in m<sup>3</sup>/kg.
- dH = weighted average heavier component density in m<sup>3</sup>/kg.
- (1/dL – 1/dH) = inverse density difference of the light (dL) and heavy (dH) components in the blended mixture.

### Example Calculation:

A heavy oil proration battery receives and blends condensate with battery oil production prior to the total blended volume being disposed of. The monthly volumes and densities are shown below.

Condensate received and blended into the heavy crude oil	Volume = 800.0 m <sup>3</sup>
Density = 700.0 kg/m <sup>3</sup>	
Heavy crude oil production	Volume = 2,300.0 m <sup>3</sup> Density = 963.0 kg/m <sup>3</sup>

$$C = (800.0 / (800.0 + 2,300.0)) \times 100 = 25.81 \%$$

$$1/dL = 1/700.0 = 0.00143$$

$$1/dH = 1/963.0 = 0.00104$$

$$\% \text{ Shrinkage} = 26,900 \times C \times (100 - C)^{0.819} \times (1/dL - 1/dH)^{2.28}$$

$$\% \text{ Shrinkage} = 26,900 \times 25.81 \times (100 - 25.81)^{0.819} \times (0.00143 - 0.00104)^{2.28}$$

$$\% \text{ Shrinkage} = 0.3990$$

For blending 2,300.0 m<sup>3</sup> of 963.0 kg/m<sup>3</sup> heavy crude oil and 800.0 m<sup>3</sup> of 700.0 kg/m<sup>3</sup> condensate:

$$\text{Shrinkage Volume} = 0.3993\% \times (800.0 + 2,300.0) = 12.38 \text{ m}^3$$

For Petrinex reporting the shrinkage volume is applied to the lower density fluid so the reporting at the battery would be as follows:

Receipt (REC) of condensate at the battery of 800.0 m<sup>3</sup>

Battery oil production (PROD) of 2,300.0 m<sup>3</sup>

Total production plus receipts = 3,100.0 m<sup>3</sup>

Battery oil disposition (DISP)= 3,087.62 m<sup>3</sup> (report 3,087.6 m<sup>3</sup>)

Condensate shrinkage (SHR) = 12.38 m<sup>3</sup> (report (12.4 m<sup>3</sup>))

Total dispositions = 3,100.0 m<sup>3</sup>

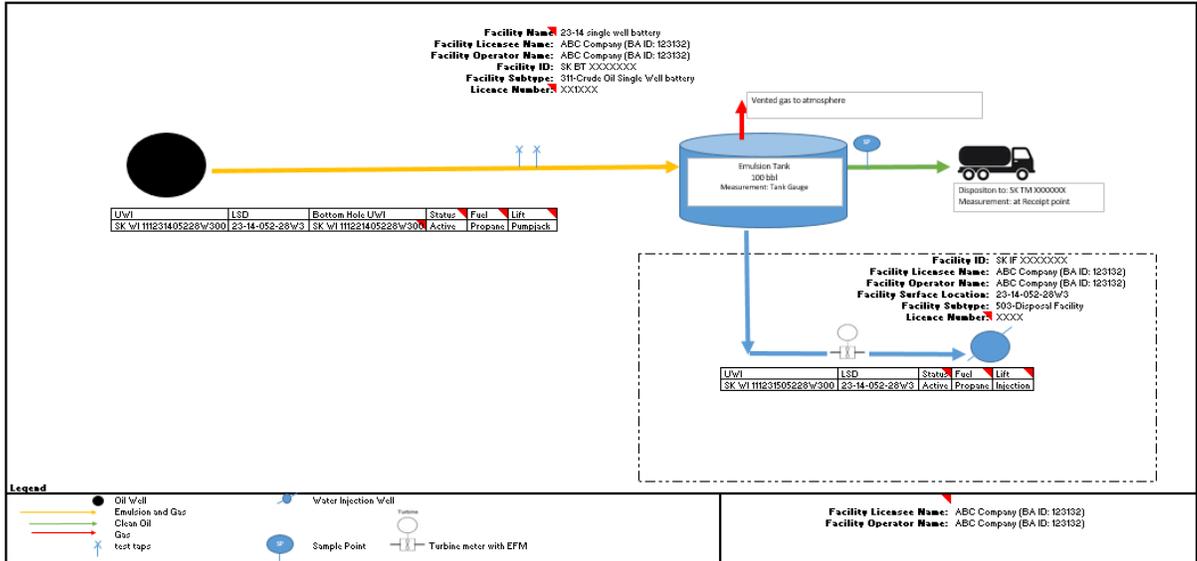
Note that, unlike process shrinkage that is calculated automatically by Petrinex at a gas plant, blending shrinkage has to be calculated “off book” or outside of the Petrinex system and then reported into Petrinex to balance the facility.

This example describes heavy crude oil production being blended with butane at a heavy crude oil multi-well proration battery; however, blending of other hydrocarbon streams at different facility sub-types can also occur.

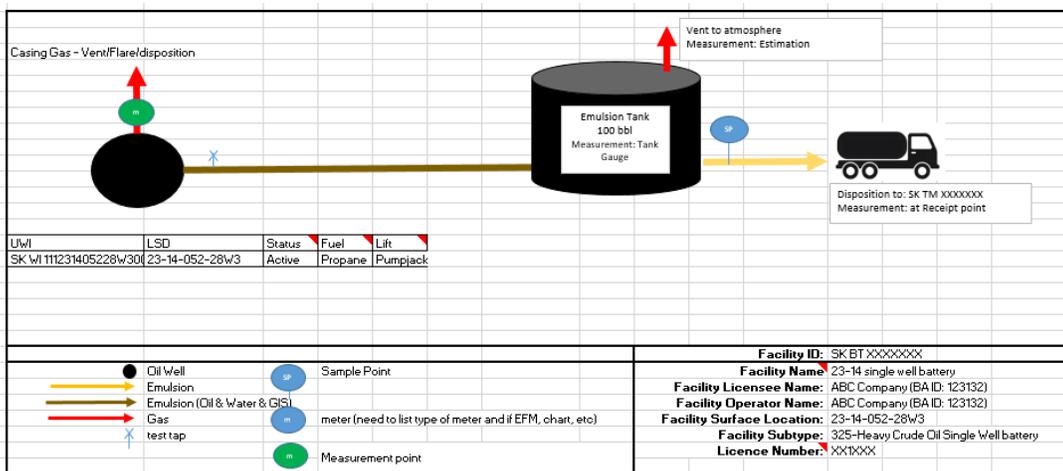
## Appendix 8 Measurement Schematic Example

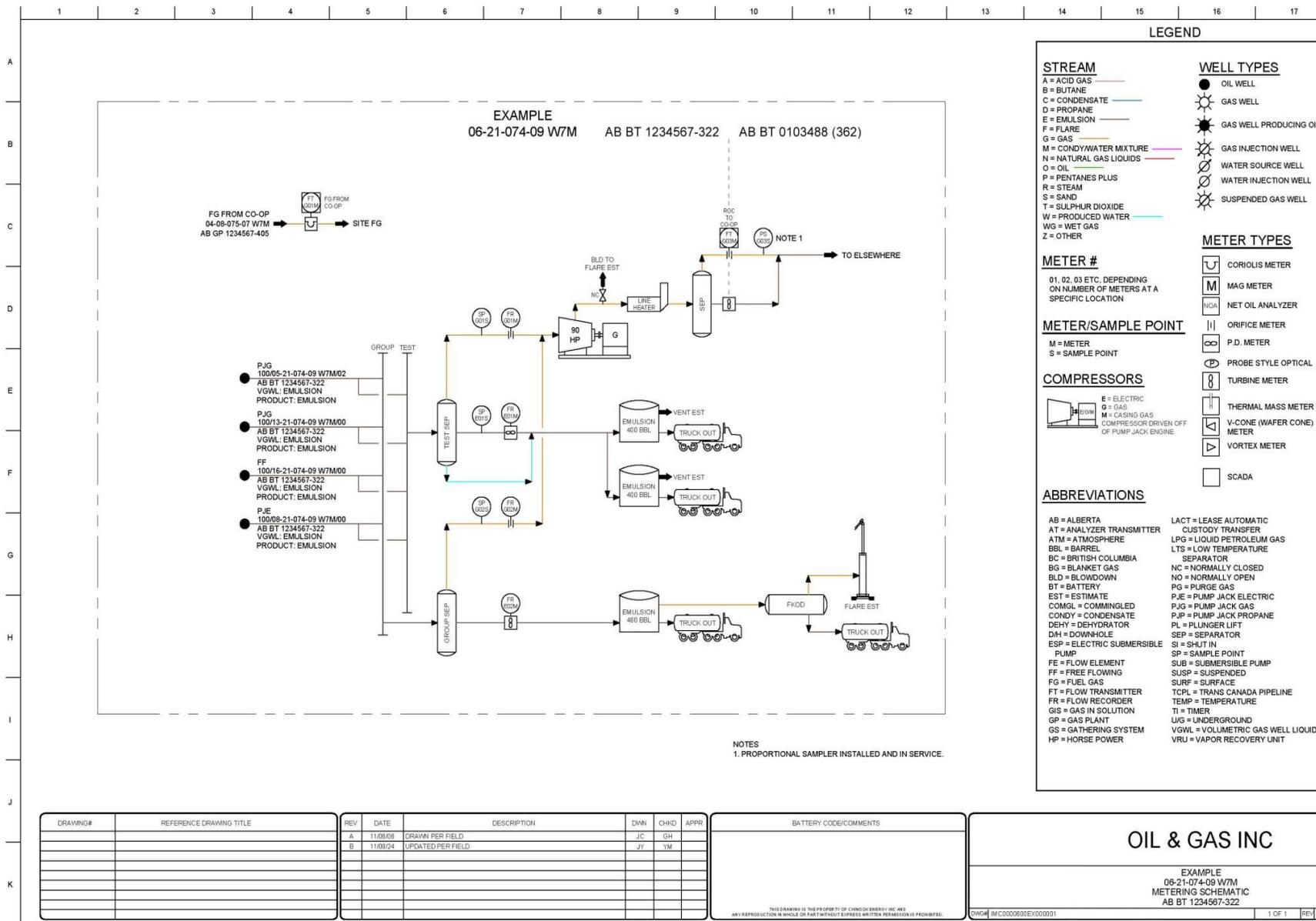
The measurement schematic on the following page is an example of a measurement schematic that is required as described in Section 1.8 and is for information purposes only. A licensee may use other symbols, letters, or words as long as it is clear in the legend or in the schematic what they stand for.

Example of facility subtype 311: Crude Oil Single Well Battery with injection facility



Example of facility subtype 325: Heavy Crude Oil Single Well Battery





**LEGEND**

**STREAM**

- A = ACID GAS
- B = BUTANE
- C = CONDENSATE
- D = PROPANE
- E = EMULSION
- F = FLARE
- G = GAS
- M = CONDY/WATER MIXTURE
- N = NATURAL GAS LIQUIDS
- O = OIL
- P = PENTANES PLUS
- R = STEAM
- S = SAND
- T = SULPHUR DIOXIDE
- W = PRODUCED WATER
- WG = WET GAS
- Z = OTHER

**WELL TYPES**

- OIL WELL
- GAS WELL
- ☀ GAS WELL PRODUCING OIL
- ☀ GAS INJECTION WELL
- ⊕ WATER SOURCE WELL
- ⊕ WATER INJECTION WELL
- ⊕ SUSPENDED GAS WELL

**METER TYPES**

**METER #**

- 01, 02, 03 ETC. DEPENDING ON NUMBER OF METERS AT A SPECIFIC LOCATION

**METER/SAMPLE POINT**

- M = METER
- S = SAMPLE POINT

**COMPRESSORS**

- E = ELECTRIC
- G = GAS
- M = CASING GAS COMPRESSOR DRIVEN OFF OF PUMP JACK ENGINE

**ABBREVIATIONS**

- AB = ALBERTA
- AT = ANALYZER TRANSMITTER
- ATM = ATMOSPHERE
- BBL = BARREL
- BC = BRITISH COLUMBIA
- BG = BLANKET GAS
- BLD = BLOWDOWN
- BT = BATTERY
- EST = ESTIMATE
- CONMGL = COMMINGLED
- CONDY = CONDENSATE
- DEHY = DEHYDRATOR
- DH = DOWNHOLE
- ESP = ELECTRIC SUBMERSIBLE PUMP
- FE = FLOW ELEMENT
- FF = FREE FLOWING
- FG = FUEL GAS
- FT = FLOW TRANSMITTER
- FR = FLOW RECORDER
- GIS = GAS IN SOLUTION
- GP = GAS PLANT
- GS = GATHERING SYSTEM
- HP = HORSE POWER
- LACT = LEASE AUTOMATIC CUSTODY TRANSFER
- LPG = LIQUID PETROLEUM GAS
- LTS = LOW TEMPERATURE SEPARATOR
- NC = NORMALLY CLOSED
- NO = NORMALLY OPEN
- PG = PURGE GAS
- PJE = PUMP JACK ELECTRIC SEPARATOR
- PJG = PUMP JACK GAS
- PJP = PUMP JACK PROPANE
- PL = PLUNGER LIFT
- SEP = SEPARATOR
- SI = SHUT IN
- SP = SAMPLE POINT
- SUB = SUBMERSIBLE PUMP
- SUSP = SUSPENDED SURF = SURFACE
- TCPPL = TRANS CANADA PIPELINE
- TEMP = TEMPERATURE
- TI = TIMER
- UG = UNDERGROUND
- VGWL = VOLUMETRIC GAS WELL LIQUID
- VRU = VAPOR RECOVERY UNIT

DRAWING#	REFERENCE DRAWING TITLE

REV	DATE	DESCRIPTION	DWN	CHKD	APPR
A	11/08/08	DRAWN PER FIELD	JC	GH	
B	11/08/24	UPDATED PER FIELD	JY	YM	

BATTERY CODE/COMMENTS

**OIL & GAS INC**

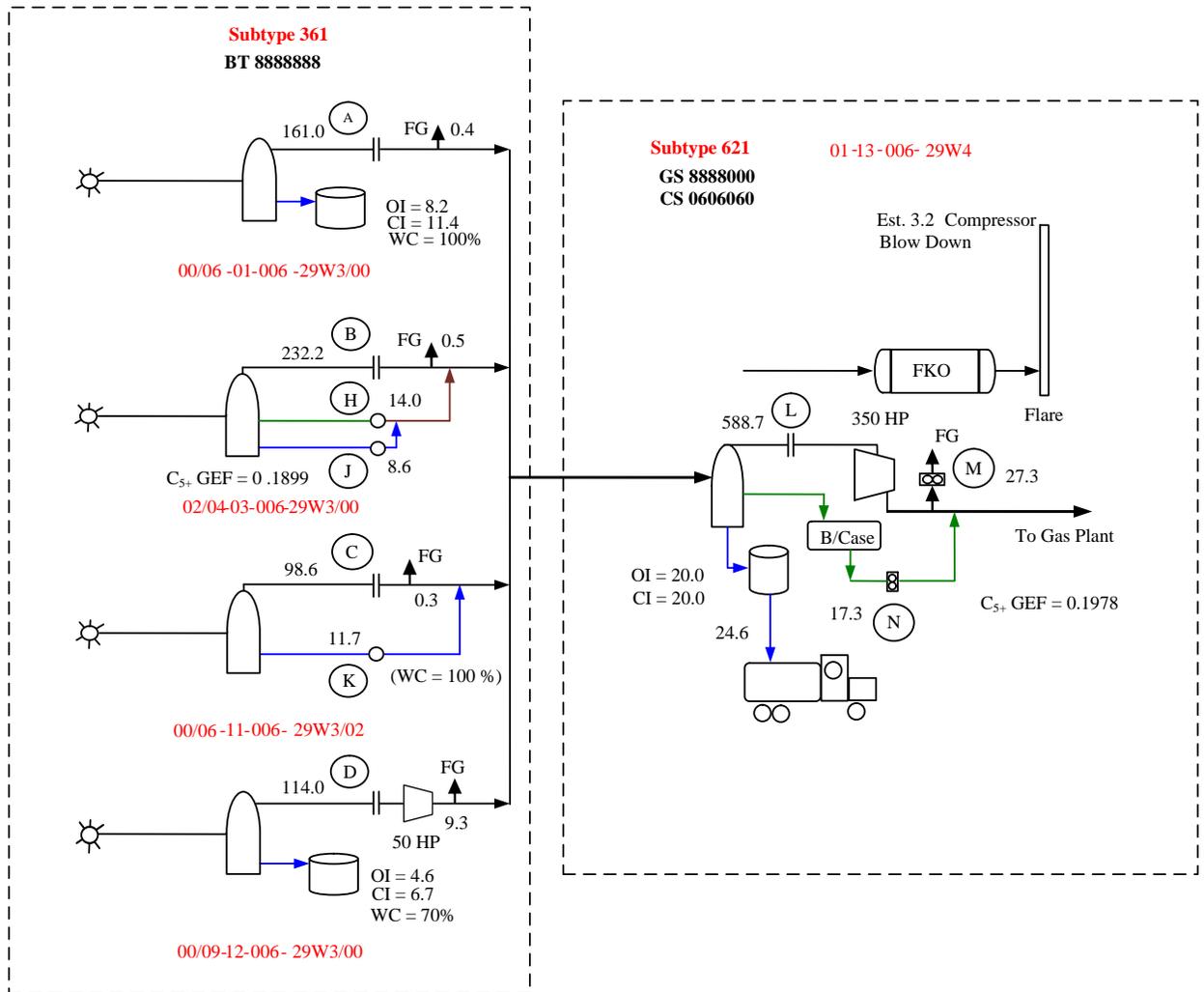
EXAMPLE  
06-21-074-09 W7M  
METERING SCHEMATIC  
AB BT 1234567-322

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1 OF 1 REV 5

## Appendix 9 Gas Group Delineation

Case 1<sup>1,2</sup>

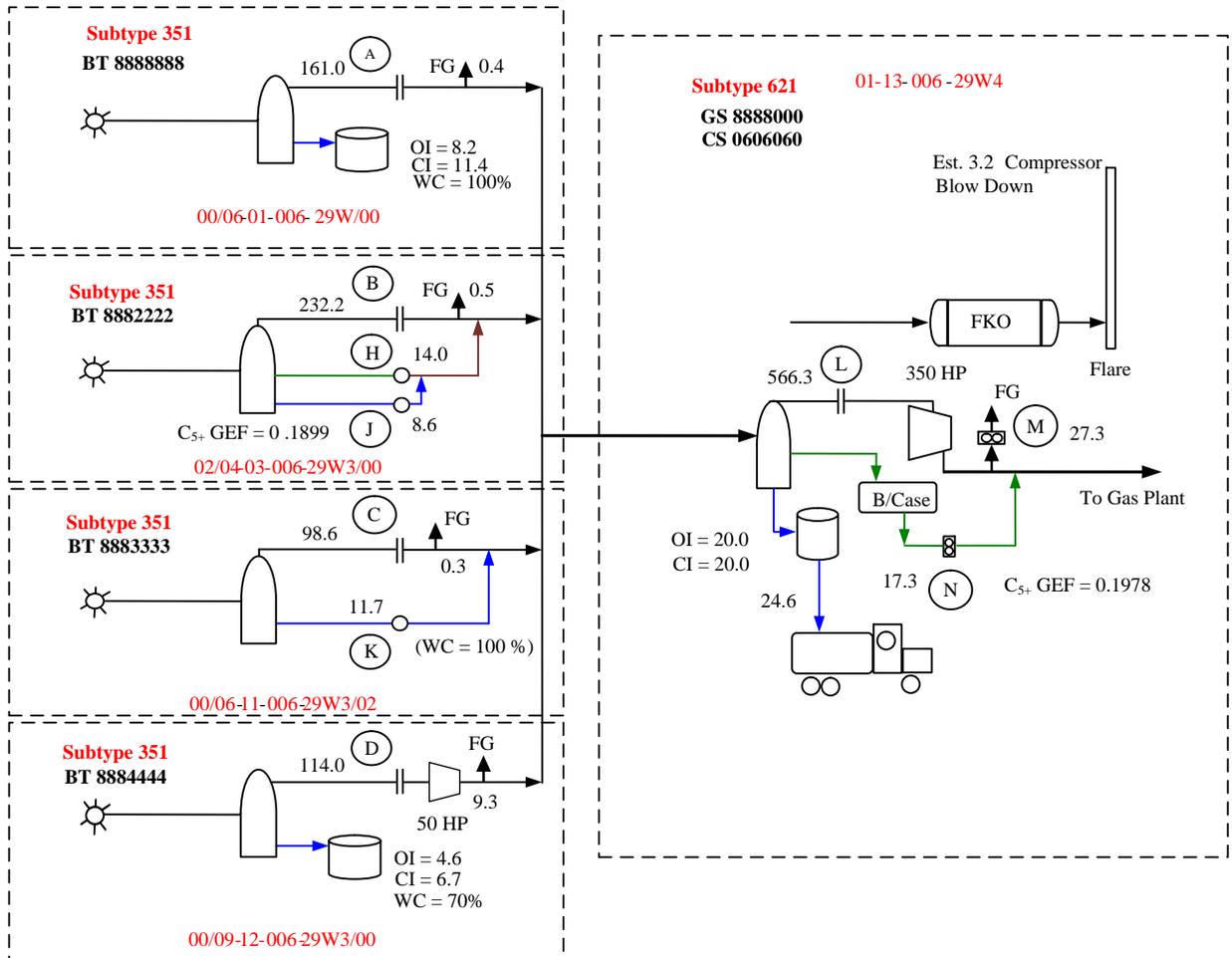


BT 8888888 Gas Production =  $161.0 + 232.2 + (0.1899 \times 14.0) + 98.6 + 114.0 = 608.5$   
 BT 8888888 Gas Delivered =  $608.5 - (0.4 + 0.5 + 0.3 + 9.3) = 598.0$   
 BT 8888888 Water Production =  $3.2 + 8.6 + 11.7 + [(6.7 - 4.6) \times 0.7] = 25.0$   
 BT 8888888 Oil Production =  $[(6.7 - 4.6) \times 0.3] = 0.6$   
 GS 8888000 Receipts = 598.0  
 GS 8888000 Gas Delivered =  $588.7 - 27.3 + (0.1978 \times 17.3) - 3.2 = 561.6$   
 GS 8888000 MD =  $598.0 - 588.7 - (0.1978 \times 17.3) = 5.9$  (1.0%)  
 GS 8888000 Water Receipts =  $8.6 + 11.7 = 20.3$ ; Delivered = 24.6  
 WC Water Receipt =  $24.6 - 20.3 = 4.3$

<sup>1</sup> All wells sweet

<sup>2</sup> All volumes monthly

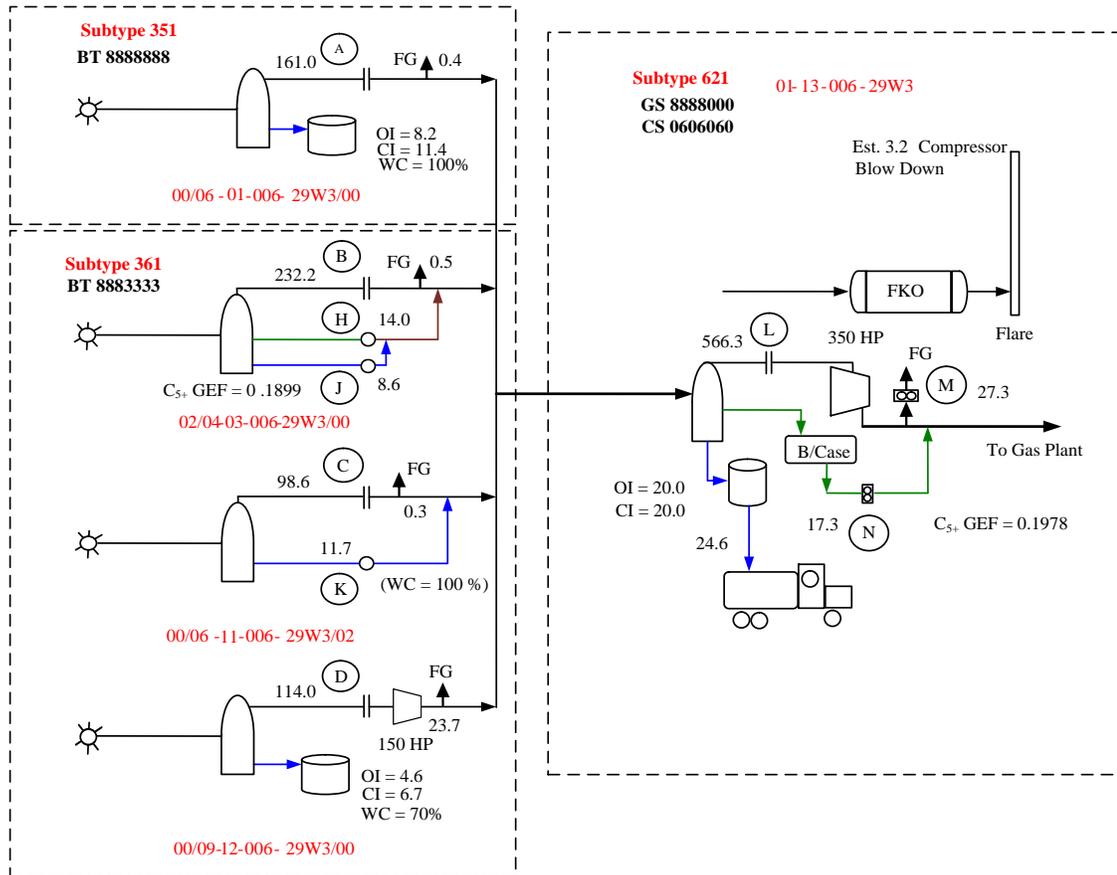
Case 2<sup>1,2</sup>



BT 8888888 Gas Production = 161.0; Delivered = 161.0 – 0.4 = 160.6
BT 8888888 Water Production = 3.2
BT 8882222 Gas Production = 232.2 + (14 x 0.1899) = 234.9; Delivered = 234.9 – 0.5 = 234.4
BT 8882222 Water Production = 8.6
BT 8883333 Gas Production = 98.6; Delivered = 98.6 – 0.3 = 98.3
BT 8883333 Water Production = 11.7
BT 8884444 Gas Production = 114.0; Delivered = 114 – 23.7 = 90.3
BT 8884444 Water Production = (6.7 – 4.6) x 0.7 = 1.5
BT 8884444 Oil Production = (6.7 – 4.6) x 0.3 = 0.6
GS 8888000 Gas Receipts = 160.6 + 234.4 + 98.3 + 90.3 = 583.6
GS 8888000 Gas Delivered = 566.3 – 27.3 + (0.1978 x 17.3) – 3.2 = 539.2
GS 8888000 MD = 583.6 – 569.7 = 13.9 (2.4%)
GS8888000 Water Receipts = 8.6 + 11.7 = 20.3; Delivered = 24.6

<sup>1</sup> All wells sweet  
<sup>2</sup> All volumes monthly

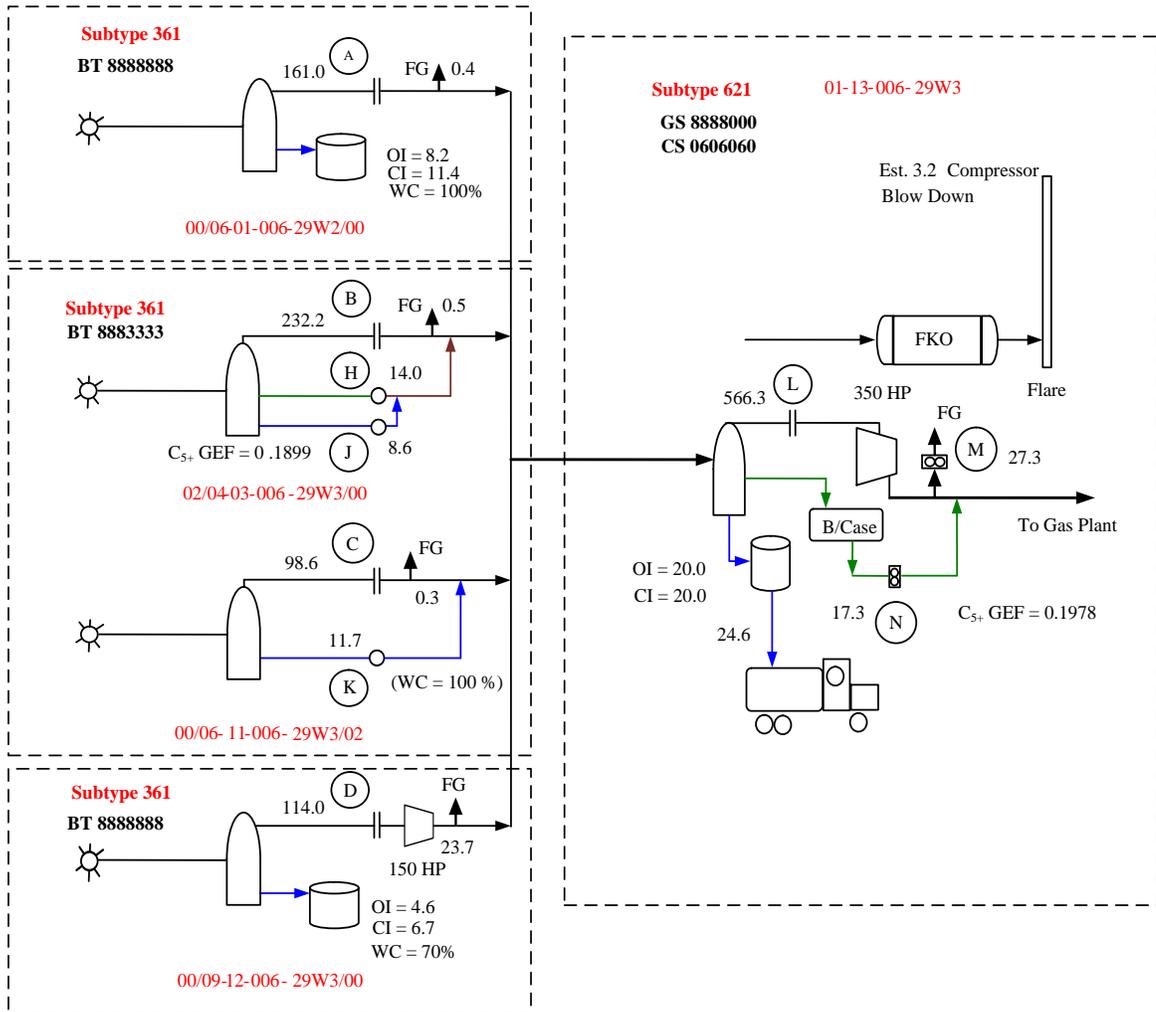
Case 3<sup>1,2</sup>



BT 8888888 Gas Production = 161.0; Delivered = 161.0 – 0.4 = 160.7  
 BT 8888888 Water Production = 3.2  
 BT 8883333 Gas Production = 232.2 + (14 x 0.1899) + 98.6 + 114 = 447.5; Delivered = 447.5 – 24.5 = 423.0  
 BT 8883333 Water Production = 8.6 + 11.7 + [(6.7 – 4.6) x 0.7] = 21.8  
 BT 8883333 Oil Production = (6.7 – 4.6) x 0.3 = 0.6  
 GS 8888000 Receipts = 423.0 + 160.6 = 583.6  
 GS 8888000 Gas Delivered = 566.3 – 27.3 + (0.1978 x 17.3) – 3.2 = 539.2  
 GS 8888000 MD = 583.6 – 569.7 = 13.9 (2.4%)  
 GS 8888000 Water Receipts = 8.6 + 11.7 = 20.3; Delivered = 24.6  
 WC Water Receipt = 24.6 – 20.3 = 4.3

<sup>1</sup> All wells sweet  
<sup>2</sup> All volumes monthly

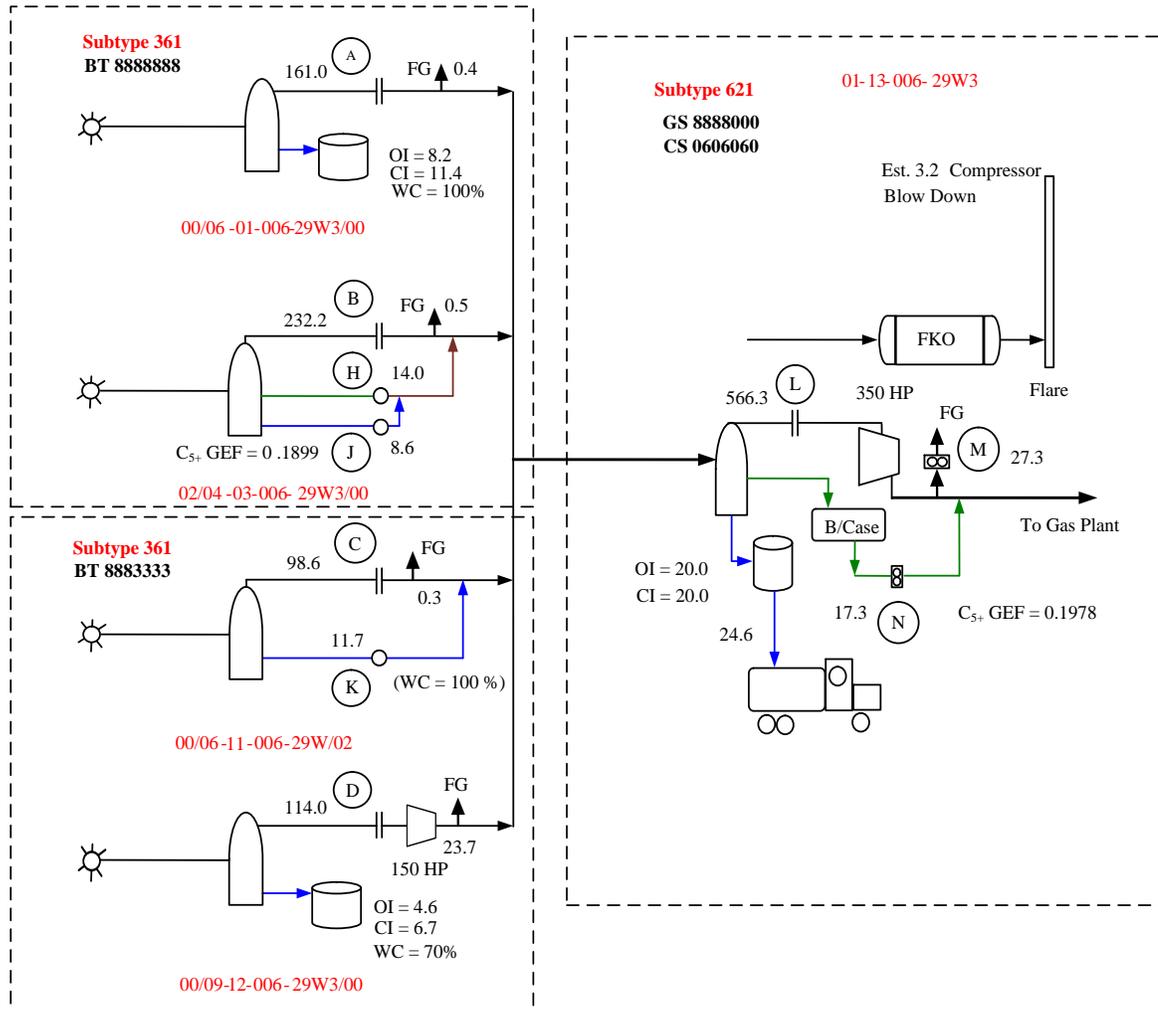
Case 4<sup>1,2</sup>



BT 8888888 Gas Production = 161.0 + 114.0 = 275.0; Delivered = 275 - 0.4 - 23.7 = 250.9  
 BT 8888888 Water Production = (11.4 - 8.2) + [(6.7 - 4.6) x 0.7] = 4.7  
 BT 8888888 Oil Production = (6.7 - 4.6) x 0.3 = 0.6  
 BT 8883333 Gas Production = 232.2 + (14 x 0.1899) + 98.6 = 333.5; Delivered = 333.5 - 0.5 - 0.3 = 332.7  
 BT 8883333 Water Production = 8.6 + 11.7 = 20.3  
 GS 8888000 Receipts = 250.9 + 332.7 = 583.6  
 GS 8888000 Gas Delivered = 566.3 - 27.3 + (0.1978 x 17.3) - 3.2 = 539.2  
 GS 8888000 MD = 583.6 - 569.7 = 13.9 (2.4%)  
 GS 8888000 Water Receipts = 8.6 + 11.7 = 20.3; Delivered = 24.6  
 WC Water Receipt = 24.6 - 20.3 = 4.3

<sup>1</sup> All wells sweet  
<sup>2</sup> All volumes monthly

Case 5<sup>1,2</sup>



BT 8888888 Gas Production =  $161.0 + 232.2 + (14 \times 0.1899) = 395.9$ ; Delivered =  $395.9 - 0.4 - 0.5 = 395.0$   
 BT 8888888 Water Production =  $(11.4 - 8.2) + 8.6 = 11.8$   
 BT 8883333 Gas Production =  $98.6 + 114 = 212.6$ ; Delivered =  $212.6 - 24 = 188.6$   
 BT 8883333 Water Production =  $11.7 + [(6.7 - 4.6) \times 0.7] = 13.2$   
 BT 8883333 Oil Production =  $(6.7 - 4.6) \times 0.3 = 0.6$   
 GS 8888000 Receipts =  $395 + 188.6 = 583.6$   
 GS 8888000 Gas Delivered =  $566.3 - 27.3 + (0.1978 \times 17.3) - 3.2 = 539.2$   
 GS 8888000 MD =  $583.6 - 569.7 = 13.9$  (2.4%)  
 GS 8888000 Water Receipts =  $8.6 + 11.7 = 20.3$ ; Delivered = 24.6  
 WC Water Receipt =  $24.6 - 20.3 = 4.3$

<sup>1</sup> All wells sweet

<sup>2</sup> All volumes monthly