Standard Practice for Storage and Use of Liquefied Petroleum Gases (LPG) in Sample Cylinders for LPG Test Methods

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1. Scope

1.1 This practice covers information for the storage and use of LPG samples in standard cylinders of the type used in sampling method, Practice D 1265 and floating piston cylinders used in sampling method, Practice D 3700.

1.2 This practice is especially applicable when the LPG sample is used as a quality control (QC) reference material for LPG test methods, such as gas chromatography (GC) analysis (Test Method D 2163) or vapor pressure (Test Method D 6897) that use only a few mL per test, since relatively small portable Department of Transportation (DOT) cylinders (for example, 20 lb common barbecue cylinders) can be used. This practice can be applied to other test methods. However, test methods that require a large amount of sample per test (for example, manual vapor pressure Test Method D 1267) will require QC volumes in excess of 1000 L if stored in standard DOT cylinders or American Society of Mechanical Engineers (ASME) vessels.

2. Referenced Documents

2.1 ASTM Standards:

D 1265 Practice for Sampling Liquefied Petroleum (LP) Gases (Manual Method)2
D 1267 Test Method for Gage Vapor Pressure of Liquefied Petroleum (LP) Gases (LP-Gas Method)2
D 2163 Test Method for Analysis of Liquefied Petroleum (LP) Gases and Propane Concentrates by Gas Chromatography2
D 3700 Practice for Containing Hydrocarbon Fluid Samples Using a Floating Piston Cylinder3
D 6299 Practice for Applying Statistical Quality Assurance Techniques to Evaluate Analytical Measurement System Performance4
D 6897 Test Method for Vapor Pressure of Liquefied Petroleum (LP) (Expansion Method)5

3. Terminology

3.1 Definitions of Terms Specific to This Standard:

3.1.1 floating piston cylinder (FPC)—a high-pressure sample container with a free floating internal piston(s) that effectively divides the container into two or more separate compartments. The sample is contained on one side of the piston (the sample or product side). The chamber on the other side of the piston (the charge or pre-charge side) is maintained at a higher pressure than the vapor pressure of the sample with an inert gas. This allows collection of a sample with no loss of volatile components and no formation of a gaseous phase that may alter the composition of the sample. The cylinder is equipped with a piston follower or indicating rod or other indicating device to show the position of the floating piston.

3.1.2 standard 80 % fill cylinder—a pressure rated cylinder or vessel such as described in Practice D 1265, or conforming to DOT or ASME cylinder standards. These cylinders are not equipped with a floating piston, and have both an equilibrium liquid and vapor phase when used for LPG.

4. Summary of Practice

4.1 This practice provides information for the design and operation of LPG sample storage cylinders taking into account properties of LPG and types of cylinders in common use for storage of LPG.

4.2 This practice provides additional guidelines to Practice D 6299 to determine the minimum volume of LPG sample material required, when used as a QC reference material.

5. Significance and Use

5.1 LPG samples can change composition during storage and use from preferential vaporization of lighter (lower molecular weight) hydrocarbon components, dissolved inert gases (N₂, Ar, He, and so forth) and other dissolved gases/liquids (NH₃, CO₂, H₂S, H₂O, etc.). Careful selection of cylinder type, cylinder volume, and use of inert gas for pressurizing cylinders is required to ensure that composition changes are small enough to maintain the integrity of LPG when used as a QC reference material for various LPG test methods.

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1 This practice is under the jurisdiction of ASTM Committee D02 on Petroleum Products and Lubricants and is the direct responsibility of Subcommittee D02.08 on Volatility.


2 Annual Book of ASTM Standards, Vol 05.01.

3 Annual Book of ASTM Standards, Vol 05.02.

4 Annual Book of ASTM Standards, Vol 05.03.

5 Annual Book of ASTM Standards, Vol 05.04.
5.2 Monitoring of ongoing precision and bias on QC materials using control chart techniques in accordance with Practice D 6299 can be used to establish the need for calibration or maintenance.

6. Reference Materials

6.1 The LPG QC reference material should have a vapor pressure and composition in the range of the samples regularly tested by the equipment. This is particularly important for LPG/natural gas liquid (NGL) mixtures near the critical temperature, as these liquids have large thermal and pressure expansion coefficients.

6.2 LPG QC reference materials should be stored in an environment suitable for long term storage without significant sample degradation for the test(s) being performed.

6.2.1 As an example, evidence of a long term shift or bias in the LPG QC reference material results obtained relative to the established statistical control limits and average value determined for the test initially, may indicate that the composition of the LPG QC reference material has significantly degraded or changed over time. An investigation should be conducted to determine if the long term stability of the QC reference material is the cause for the out-of-control situation.

7. Use of Floating Piston Cylinders for LPG Samples

7.1 Minimum LPG sample volume can be determined in accordance with Practice D 6299.

7.2 Floating piston cylinders (see Fig. 1) are preferred for LPG sample materials for tests involving accurate determination of light gases.

7.3 Excessive inert gas pressure should be avoided for long term storage of vapor pressure QC or calibrant materials in floating piston cylinders. Leakage of inert gas past worn or damaged floating piston seals can cause an increase in dissolved gas concentration and vapor pressure of the QC sample material.

8. Use of Standard 80 % Fill Cylinders for LPG QC Materials

8.1 Common 80 % filled storage tanks or cylinders can be used for LPG QC materials provided that the QC material batch volume is sufficiently large to avoid adverse short term vaporization effects.

8.2 The total initial volume and the minimum unused volume of QC materials stored in standard 80 % fill cylinders must be controlled to ensure that in the short term, composition is constant relative to the precision of the test method.

8.2.1 As liquid is withdrawn from LPG cylinders, a small amount of the remaining liquid must vaporize to replace the volume. This results in a small, but predictable, change in composition and vapor pressure from preferential vaporization of lighter components from the remaining liquid. The composition and vapor pressure changes are known to be approximately linear at low vapor to liquid (V/L) ratios. These changes accelerate and become more significant as the remaining volume of liquid decreases and the cylinder approaches empty. However, if the initial volume is sufficiently large, and the final V/L ratio is limited, the change will occur very slowly over time, and the material is still suitable as a QC. In the short term, the composition is essentially constant relative to the precision of the method.

8.2.2 In the long run, the control limits can be periodically adjusted to compensate for any long-term trend, or the charted response can be compensated for the long-term trend using historical data, or equation of state calculations based on cylinder weight or volume. Consult a statistician for appropriate techniques to develop a prediction model for the long-term trend.

8.2.3 Operation between the 80 % and 20 % fill levels is recommended to satisfy safety requirements and to limit the V/L ratio from 0.25 (1:4) at 80 % liquid filled up to 4 (4:1) at 20 % filled. The cylinder must be re-filled when the liquid level drops below the 20 % level and no further liquid can be withdrawn (see 8.2). This guards against excessive changes in concentration of the remaining QC liquid as would occur with the exponentially increasing vapor/liquid ratio as the liquid volume approaches zero.

8.2.4 The minimum initial QC volume and the maximum number of usable QC runs for the batch volume can be assessed by performing a simple linear regression of the first 20 valid QC results against the observation number and by testing the slope for significance versus zero. Upon a non-significant outcome, continue to perform this regression after every ten additional results until either the slope fails the significance test or the control chart detects a trend. The total number of QC runs cumulated will then constitute the maximum useful runs for the QC batch volume.
NOTE 3—This methodology requires the time between QC results to be long enough such that the long term variation of the test method is observable.

8.3 Common 20 lb or larger DOT approved cylinders (used for home barbecues and mobile applications) equipped with a 20 % liquid level dip tube have been found to be suitable for laboratory GC or instrument vapor pressure (VP) applications that use less than 15 mL per test. The dip tube can be used to establish the 80 % liquid fill by inverting the cylinder and venting liquid using the procedure in Practice D 1265 (see Fig. 2).

8.4 Pressurizing a standard 80 % fill cylinder with an inert gas will result in the inert gas becoming partially soluble in the LPG QC material, which can affect some test results. (Warning—Do not exceed the working temperature or pressure of the storage cylinder.) (Warning—Use re-settable pressure relief valves and not burst disks for laboratory use.)

8.4.1 Common 80 % filled LPG storage cylinders may be pressurized to facilitate liquid transfer and repeatable liquid injections for GC analysis (see Appendix X1). Some GC test methods require specific injection conditions, for example minimum 200 psi above sample vapor pressure, to ensure repeatable liquid injections.

8.4.2 Common 80 % fill QC storage cylinders must not be pressurized with inert gas to facilitate liquid transfer for vapor pressure measurements, as this will affect the result.

8.5 Other vapor tight means of generating sufficient transfer/injection pressure are acceptable, such as magnetically coupled or other sealed cavity pumps.

9. Keywords

9.1 floating piston cylinder; liquefied petroleum gas (LPG); LPG sample storage cylinders; quality control (QC); standard 80% fill cylinder

APPENDIX

(Nonmandatory Information)

X1. INERT GAS PRESSURIZATION WITH STANDARD 80 % FILL CYLINDERS

X1.1 Pressurizing a standard 80 % fill cylinder with an inert gas will result in the inert gas becoming partially soluble in the LPG sample, which can affect some test results.

X1.2 Pressurizing a common 20 lb DOT cylinder to the maximum working pressure of 240 psig will result in approximately 2 mole % nitrogen in the liquid propane, and about 50/50 molar ratio of nitrogen and propane in the equilibrium vapor. The mixture is still at its bubble point, so any increase in temperature or decrease in pressure in sample lines or instrument test cells can still result in formation of vapor.

X1.3 Liquid sample (inject) valves (LSV) are generally slightly above ambient temperature due to proximity to the instrument, and this can cause localized vaporization in the valve and erratic injection volumes. Flushing the valve several times prior to injection provides some local cooling, and it provides for more repeatable liquid injections. In general, the LSV should be kept as close to ambient temperature as practical. This allows the use of lower inert gas pressures or storing the LPG samples at about 5 to 8°C (10 to 15°F) above ambient temperature to obtain repeatable liquid injections.

X1.4 Use of higher inert gas pressures than required to obtain repeatable liquid injections does not limit or control vapor losses in a standard cylinder. Inert gas in a standard cylinder equilibrates with both the liquid and vapor phases, becoming partially dissolved in the liquid. The increase in the total pressure due to inert gas does NOT cause the volatile hydrocarbons to condense or otherwise knock down the hydrocarbon vapor (this is a common misconception). High inert gas pressures cannot compensate for excessive vaporization of the liquid sample. The same errors will be incurred from excessive vapor formation with or without addition of inert gas to the cylinder. The same precautions must be taken to limit vapor losses with or without the use of inert gas to pressurize a standard (non-floating piston) cylinder.

X1.5 Helium is the preferred inert gas for thermal conductivity detector instruments, since it is used as the carrier gas in the GC and will not be detected. Nitrogen will be detected in a thermoconductivity (TC) detector, and it may interfere with the analysis. Nitrogen is not detected and may be used in flame ionization detector (FID) methods, but it may not give as repeatable results as helium at high LSV temperatures due to higher dissolved nitrogen concentration at the same pressure (lower vapor/liquid relative volatility “K” ratio). Heavier inert gases are not recommended.