



Designation: ~~E126 – 13a~~ E126 – 19

Standard Test Method for Inspection, Calibration, and Verification of ASTM Hydrometers¹

This standard is issued under the fixed designation E126; the number immediately following the designation indicates the year of original adoption or, in the case of revision, the year of last revision. A number in parentheses indicates the year of last reapproval. A superscript epsilon (ϵ) indicates an editorial change since the last revision or reapproval.

This standard has been approved for use by agencies of the U.S. Department of Defense.

1. Scope

1.1 This test method describes the principles, apparatus, and procedures for the inspection, calibration, and verification of ASTM glass hydrometers. This test method is applicable to ASTM hydrometers and may be used for other general hydrometers of the constant-mass, variable-displacement type.

1.2 The values stated in inch-pound units are to be regarded as the standard. The values given in parentheses are mathematical conversions to SI units that are provided for information only and are not considered standard. The metric equivalents of inch-pound units may be approximate.

1.3 *This standard does not purport to address all of the safety concerns, if any, associated with its use. It is the responsibility of the user of this standard to establish appropriate ~~safety~~ safety, health, and ~~health~~ environmental practices and determine the applicability of regulatory limitations prior to use.*

1.4 *This international standard was developed in accordance with internationally recognized principles on standardization established in the Decision on Principles for the Development of International Standards, Guides and Recommendations issued by the World Trade Organization Technical Barriers to Trade (TBT) Committee.*

2. Referenced Documents

2.1 ASTM Standards:²

[D1265 Practice for Sampling Liquefied Petroleum \(LP\) Gases, Manual Method](#)

[D1298 Test Method for Density, Relative Density, or API Gravity of Crude Petroleum and Liquid Petroleum Products by Hydrometer Method](#)

[D1657 Test Method for Density or Relative Density of Light Hydrocarbons by Pressure Hydrometer](#)

[E1 Specification for ASTM Liquid-in-Glass Thermometers](#)

[E77 Test Method for Inspection and Verification of Thermometers](#)

[E100 Specification for ASTM Hydrometers](#)

[E344 Terminology Relating to Thermometry and Hydrometry](#)

[E2251 Specification for Liquid-in-Glass ASTM Thermometers with Low-Hazard Precision Liquids](#)

[E2995 Specification for ASTM Thermohydrometers with Integral Low-Hazard Thermometers](#)

2.2 Other Documents

[ANSI/NCSLI Guide to Expression of Uncertainty in Measurement](#)

3. Terminology

3.1 *Definitions*—The definitions given in Terminology [E344](#) apply.

3.2 *Definitions of Terms Specific to This Standard:*

3.2.1 *API gravity, n*—a relative index of density for petroleum products developed by the American Petroleum Institute. API gravity is defined as:

¹ This test method is under the jurisdiction of ASTM Committee [E20](#) on Temperature Measurement and is the direct responsibility of Subcommittee [E20.05](#) on Liquid-in-Glass Thermometers and Hydrometers.

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² For referenced ASTM standards, visit the ASTM website, [www.astm.org](#), or contact ASTM Customer Service at [service@astm.org](#). For *Annual Book of ASTM Standards* volume information, refer to the standard's Document Summary page on the ASTM website.

$$\text{API Gravity, deg} = [141.5/(\text{rel. density } 60/60 \text{ } ^\circ\text{F})] - 131.5 \quad (1)$$

Values of API gravity are typically expressed in degrees API, that is, 39.60 °API.

3.2.2 *comparator, n*—in this test method, a glass or other transparent cylinder to contain a liquid in which hydrometers may be compared. Examples of suitable comparators are given in [Appendix X1](#).

3.2.3 *density, n*—mass of a unit volume of material.

3.2.3.1 Discussion—

Units of density in hydrometers include kg/l (kilograms per liter), kg/m³ (kilograms per cubic meter), and g/l (grams per liter); each typically expressed as mass per volume at a specified temperature, that is, kg/m³ at 15 °C. As of this writing, only the kg/m³ at 15 °C scale is offered in ASTM hydrometers (see Specification [E100](#)).

3.2.4 *relative density (formerly specific gravity), n*—ratio of the mass of a given volume of material at a stated temperature to the mass of an equal volume of gas-free distilled water at the same or different temperature. Both reference temperatures shall be explicitly stated.

3.2.4.1 Discussion—

Common reference temperatures include 60 °F/60 °F, 20 °C/20 °C, 20 °C/4 °C. The historic term, specific gravity, may still be found.

3.2.5 *specific gravity, n*—historic term, replaced by *relative density*.

3.2.6 *thermo-hydrometer, n*—glass hydrometer having a thermometer combined with a hydrometer in one instrument.

3.2.7 *verification, n*—confirmation, by provision of objective evidence, that the instrument fulfills specified requirements.

3.2.7.1 Discussion—

In this test method, if the hydrometer bears an ASTM designation, the requirements for the maximum scale error and dimensions for the hydrometers given in Specification [E100](#) apply.

3.2.8 Other descriptions of terms relating to thermometers are included in Test Method [E77](#). Any definitions of measurement uncertainty used in this standard are from Terminology [E344](#).

4. Significance and Use

4.1 The purpose of this test method is to establish a common method by which manufacturers, calibration laboratories, and users of hydrometers may inspect, verify, or calibrate them.

4.2 The goal is to provide a standard method that is simple, easily understood, and will produce reliable results.

5. Apparatus

5.1 *Graduated Metal Scales*, of the conventional type, for checking linear dimensions. If more convenient, metal templates on which lines are ruled at suitable distances from reference points corresponding to the maximum and minimum values of the specified dimensions may be used.

5.2 *Micrometers*, of the conventional type, for checking diameters.

5.3 *Polariscope*, for viewing strain patterns in the glass developed during the manufacturing of the hydrometer.

5.4 *Comparators*, for the calibration and verification of hydrometers. Suitable types are described in [Appendix X1](#).

5.5 *Equipment*, for checking the thermometer portion of thermohydrometers as described in Test Method [E77](#).

5.6 *Thermometer(s)*, for use in pressure hydrometer cylinder comparator, ASTM 12C (-20/102 °C, 0.2° divisions), ASTM 12F (-5/215 °F, 0.5° divisions), 136C (-20/60 °C, 0.2° divisions), or ASTM 136F (-5/140 °F, 0.5° divisions) found in ASTM E1 or ASTM S12C (-20/102 °C, 0.2° divisions), or ASTM S12F (-5/215 °F, 0.5° divisions) found in Specification [E2251](#).

6. Reference Standards

6.1 *Standard Hydrometers*—Standard hydrometers shall have similar dimensions and shape to the instruments to be calibrated, (when possible), and shall have dimensions and shapes similar to the instruments to be calibrated.

NOTE 1—The relative density (specific gravity) of liquids used in calibrating hydrometers may be obtained by hydrostatic weighing instead of by the

use of reference standards as described above. Details of the hydrostatic weighing apparatus can be found in the *Dictionary of Applied Physics*³ or *Density of Solids and Liquids*.⁴

6.2 Standards shall be calibrated by either a national metrology body (such as the National Institute of Standards and Technology) or other laboratory competent to calibrate instruments of such precision. The calibration report shall provide traceability to a national metrology body and shall contain a statement of measurement uncertainty. It is desirable that the corrections be stated to one-tenth of a scale division.

6.3 Standards shall be visually inspected every six months or prior to use, whichever is longer.

6.3.1 Visual inspection shall include, but is not limited to, looking for evidence of scratches, etching, scale slippage, deposits on the glass, and discoloration. The presence of any of these defects is an indication that the standard may require re-calibration or replacement.

6.3.2 Experience has shown that the indications of hydrometers may show drift with continued use. A procedure shall be in place to demonstrate continued validity of the calibration results for the standard hydrometer. Such a procedure may include: periodic re-calibration of the standard hydrometers; measurements of hydrometers retained by the testing laboratory for use as check standards; or checks of one standard hydrometer against another.

7. Procedure

7.1 Inspection:

7.1.1 Inspect the hydrometer carefully to be certain there are no cracks, fissures, deep scratches, rough areas, or other obvious damage to the glass. Reject the hydrometer if any of these defects are present.

7.1.2 Using a polariscope, inspect the hydrometer for strain in the glass, especially at the stem/body junction. If the strain appears severe and will compromise the integrity of the hydrometer, reject the instrument. This is particularly important for thermohydrometers. See Test Method E77, 6.1.4 for more details.

7.1.3 Inspect the hydrometer carefully for loose pieces of ballast or other foreign material within the instrument. If present, reject the instrument.

7.1.4 Inspect the paper scale within the hydrometer stem. The paper scale shall be straight and without twist.

7.1.5 Inspect for the presence of a scale slippage indicator. Typically, this is a thin strand of red glass, fused to the inside top of the stem, and terminating at the first major graduation of the hydrometer scale; however, other schemes are permitted, such as etching a line on the glass corresponding to a reference line printed on the scale. If a permitted scale slippage indicator is damaged, incorrectly positioned, or not present, reject the instrument. See Specification E100 for more details.

NOTE 2—Hydrometers that do not carry an ASTM designation may not be required to have a scale slippage indicator. In such cases, a cautionary note on the report would be appropriate.

7.2 Dimensional Inspection:

7.2.1 Check the linear dimensions and diameters for compliance with Specification E100 requirements by comparing the hydrometer with the appropriate device described in 5.1 and 5.2.

7.2.2 Inspect the hydrometers for correctness of graduation spacing. API and Baumé hydrometers are graduated with equal spacing. The interval between graduations of density and relative density (specific gravity) hydrometers is smaller near the bottom of the scale. The proper spacing shall be obtained from the following formula:

$$l = L \times d_2/d \times (d - d_1)/(d_2 - d_1) \quad (2)$$

where:

l = distance from the top line to any line, d , between the top and the bottom,

L = distance between the top and the bottom graduations of the scale,

d_2 = density value, or relative density (specific gravity), of the bottom line, and

d_1 = density value, or relative density (specific gravity), of the top line.

7.2.3 Check the scale of hydrometers graduated to read percent of alcohol by weight or by volume by comparison with the values for master scales given in the *Standard Density and Volumetric Tables*.⁵

7.3 Calibration:

7.3.1 General Considerations:

7.3.1.1 In general, each hydrometer shall be calibrated at a minimum of three calibration points, spaced approximately equally across its range, nominally high, low and mid scale. For example, a hydrometer with a range of 9° to 21° API shall be calibrated at (approximately) 10, 15, and 20 API.

NOTE 3—Certain ASTM hydrometers, notably ASTM 10H and 60H (89/101 °API) and 101H and 310H (0.600 to 0.650 relative density and 500 to

³ *Dictionary of Applied Physics*, MacMillan and Co., London, Vol 3, p. 439.

⁴ "Density of Solids and Liquids," National Institute of Standards and Technology, *Circular No. 487*.

⁵ "Standard Density and Volumetric Tables," National Institute of Standards and Technology, *Circular, No. 19*.

650 kg/m³, respectively) have ranges which cannot be fully calibrated due to fluid limitations. In these cases, the hydrometers may be calibrated at two calibration points.

7.3.1.2 In order that readings shall be uniform and reproducible, the hydrometer must be clean, dry, and at the temperature of the liquid before immersing to take a reading. It is particularly important that the stem be clean so that the liquid will rise uniformly around the stem and merge into an imperceptible film on the stem.

7.3.1.3 *Cleanliness*—The readiness with which proper cleanliness can be obtained depends somewhat on the character of the liquid. Certain liquids, such as mineral oils and strong alcoholic mixtures, adhere to the stem very readily. In such cases, wiping with a lint-free cloth moistened with acetone or alcohol and drying immediately before each reading is usually sufficient. On the other hand, with weak aqueous solutions of sugar, salts, acids, and alcohol, scrupulous cleaning of the stem is required. For such liquids, two methods for preparing instruments for calibration are in common use. In one method, hydrometers are dipped in a mixture of one part concentrated sulfuric acid and two parts fuming sulfuric acid, thoroughly rinsed with water, and dried by wiping with a clean cloth. In the other method, hydrometers are washed with soap and water, dried, and wiped with a cloth moistened with alcohol to remove any residual soap film. The stems can usually be kept clean during the calibration by wiping with a lint-free cloth moistened with alcohol (preferably absolute) and drying before each reading. (**Warning**—EXTREME CAUTION—The cleaning process using concentrated sulfuric acid and fuming sulfuric acid is extremely hazardous. This process should only be carried out in a laboratory setting with appropriate equipment and trained personnel. The hydrometer must be dry before being inserted in the acids. The reaction caused by introducing a wet hydrometer into the acids may splash acids on the operator.)

7.3.1.4 *Influence of Temperature*—For a hydrometer to indicate the density of a specified liquid correctly, it is essential that the liquid be homogenous and uniform in temperature. In comparing two hydrometers having the same standard temperature and made of the same type of glass, the temperature of the liquid need not be considered since the correction required due to variation from the standard temperature is the same for both instruments. But the temperatures of the liquid, the hydrometers, and the surrounding atmosphere shall be nearly equal during the comparison; otherwise, the temperature of the liquid will be changing, causing differences in density. The operator shall allow enough time to achieve this equilibrium. To ensure homogeneity and temperature uniformity in the liquid, thorough mixing is required immediately before making measurements.

NOTE 4—Equipment such as thermometers described in 5.6, or alternative thermometric devices of equal or better accuracy, may be used, if desired.

7.3.1.5 *Influence of Surface Tension*—When a hydrometer is floated in a liquid, a small quantity of the liquid rises about the stem to form a meniscus. This liquid adhering to the stem above the general level of the liquid in which the instrument is floating has the same effect as adding to the mass of the hydrometer, thus increasing the depth of immersion.

7.3.1.6 Because a hydrometer will indicate differently in two liquids having the same density but different surface tensions, and since surface tension is a specific property of liquids, it is necessary to specify the liquid for which a hydrometer is intended. Although hydrometers of equivalent dimensions may be compared, without error, in a liquid differing in surface tension from the specified liquid, the results of comparisons of dissimilar instruments in such a liquid shall be corrected for the effect of the surface tension. [ps://standards.iteh.ai/catalog/standards/sist/480eb705-5a09-45bc-8040-9742e8032454/astm-e126-19](https://standards.iteh.ai/catalog/standards/sist/480eb705-5a09-45bc-8040-9742e8032454/astm-e126-19)

7.3.1.7 In many liquids spontaneous changes in surface tension occur due to the formation of surface films of impurities, which may come from the apparatus, the liquid, or the air. Errors from this cause may be avoided by the use of liquids not subject to such changes. However, if the liquid used is different in surface tension from the specified liquid, a correction is required when dissimilar instruments are compared, as mentioned above. A second method of avoiding these errors is to purify the surface of the calibration liquid by causing an overflow of the liquid before making an observation.

7.3.1.8 The necessity for such special manipulation is confined to the reading of hydrometers in liquids that are subject to surface contamination, such as aqueous solutions or mixtures of acids, alkalis, salts, sugar, and weak alcoholic mixtures. Oils, alcoholic mixtures of strength above ~~40%~~40 % by volume, and other liquids of relatively low surface tension are not, in general, liable to surface contamination sufficient to cause appreciable changes in hydrometer readings.

NOTE 5—For further discussion of surface tension and of meniscus corrections, see Test Method D1298.

7.3.1.9 *Comparison Liquids*—Hydrometers shall be calibrated in liquids similar in density and surface tension to the liquids in which the hydrometer is designed to be used. The liquids in Table 1 are suggested as suitable, although any liquid of proper density and surface tension may be substituted. Intermediate densities may be obtained by mixing.

7.3.1.10 *Ventilation*—Adequate ventilation (fume hood) is desirable with some of the liquids used in calibration. However, care must be taken that the ventilating method does not cause strong air currents or drafts, since such movement of air might influence the calibration.

7.3.1.11 *Taking a Reading*—Observe a point slightly below the plane of the liquid surface and then raise the line of vision until this surface, seen as an ellipse, becomes a straight line. The point where this line cuts the hydrometer scale is the reading of the instrument. Holding a white card behind the comparator just below the liquid level will improve the visibility of the surface. Estimate to one-tenth scale division.

7.4 Performing a Single Substitution Comparison:

7.4.1 Hydrometers for Liquids Having Low Vapor Pressure and All Surface Tensions:

TABLE 1 Liquids Having Suitable Surface Tensions for Comparison Tests

Relative Density, 60/60 °F	Liquid
Hydrometers for Light Liquids Other than Alcohol	
0.5077	pure grade propane ^A
0.5844	pure grade <i>n</i> -butane ^A
0.5967	dimethylpropane
0.6247	isopentane
0.625	petroleum ether
0.6310	<i>n</i> -pentane
0.6540	2,2-dimethylbutane
0.6962	isooctane
0.7504	cyclopentane
0.657 to 1.100	mineral spirits, with a relative density of approximately 0.775 (the relative density of mineral spirits can be lessened with the addition of petroleum ether or increased by the addition of tetrabromoethane)
1.070	3-methylsulfolane
1.100	safrole
Hydrometers for Alcohol	
0.750 to 1.000	ethanol and water
Hydrometers for Heavy Liquids	
1.000 to 2.000	sulfuric acid - water mixtures, or a solution of mineral spirits and tetrabromoethane

^A Because there are at present no standardized hydrometers available in the 0.500 to 0.600 relative density range, pure grade propane and *n*-butane may be used as standardizing liquids. Orders should carry the notation "For ASTM Hydrometer Standard—show relative density on label." The relative density value given on the label will be based on a spectrographic, freezing point, or chromatographic analysis of the material and will be accurate to ±0.0002. In the use of these materials care should be exercised to observe the safety precaution given in Practice [D1265](#).

7.4.1.1 Prepare a worksheet for the hydrometer being calibrated, permitting appropriate spaces each calibration point, the readings of the standard hydrometer, the observed readings of the hydrometer to be calibrated, and all other pertinent details of the calibration.

7.4.1.2 Use of the comparators described in [X1.1](#) may be appropriate. The cylinder shall have sufficient depth so that the hydrometers float freely at least 25 mm (1 in.) above the inside bottom. The cylinder diameter shall be large enough so that there will be at least 12.5 mm (½ in.) between the inner wall and any hydrometer immersed in the cylinder.

7.4.1.3 Clean the standard hydrometer, the hydrometer to be calibrated (DUT), and the comparator thoroughly prior to beginning the comparison. A final wipe, rinse, or immersion of the hydrometers into the comparison fluid is recommended.

7.4.1.4 Introduce enough of the appropriate comparison liquid into the comparator so the hydrometer will float as described in [7.4.1.2](#). Pour the liquid down the side of the comparator to avoid the formation of air bubbles. Adjust the density of the liquid, if necessary, so that its density is within three scale divisions of the nominal calibration point on the hydrometer being calibrated.

NOTE 6—With a liquid mixture, the components with lower surface tension will tend to concentrate at the surface.

NOTE 7—This method has not been evaluated for non-ASTM hydrometers or when the standard hydrometer and test hydrometer are dissimilar. In these cases, it may be beneficial to have a liquid with density within one-half to one scale division of the nominal density being measured.

7.4.1.5 Perform the first single substitution comparison. Stir the liquid, avoiding the formation of air bubbles, and then introduce a standard hydrometer into the cylinder. Slowly immerse the standard hydrometer to slightly beyond the calibration point and allow to float freely. Read the standard hydrometer as directed in [7.3.1.11](#), and record the reading (SR_1) on the worksheet. Remove the standard hydrometer and slowly immerse the DUT to slightly beyond the calibration point, allow it to float freely. Record this reading (R_{1DUT}). Remove the DUT and again immerse the standard hydrometer, read it, and record the reading (SR_2). Average SR_1 and SR_2 and apply the correction from the calibration report to the result. Subtracting the reading of the DUT (R_{1DUT}) from this value yields its correction.

First single substitution:

$$C1_{DUT} = [(SR_1 + SR_2)/2 + C_s] - R1_{DUT}$$

Second single substitution:

$$C2_{DUT} = [(SR_3 + SR_4)/2 + C_s] - R2_{DUT}$$

Third single substitution:

$$C3_{DUT} = [(SR_5 + SR_6)/2 + C_s] - R3_{DUT}$$

$$\text{FINAL CORRECTION}_{DUT} = (C1_{DUT} + C2_{DUT} + C3_{DUT})/3$$

Where:

- SR_1 = first reading of the standard
- SR_2 = second reading of the standard
- SR_3 = third reading of the standard
- SR_4 = fourth reading of the standard

- SR_5 = fifth reading of the standard
 SR_6 = sixth reading of the standard
 C_S = correction to the standard (from its calibration report)
 $R1_{DUT}$ = reading of the DUT in the first single substitution comparison
 $R2_{DUT}$ = reading of the DUT in the second single substitution comparison
 $R3_{DUT}$ = reading of the DUT in the third single substitution comparison
 $C1_{DUT}$ = correction to the DUT determined in the first single substitution comparison
 $C2_{DUT}$ = correction to the DUT determined in the second single substitution comparison
 $C3_{DUT}$ = correction to the DUT determined in the third single substitution comparison

7.4.1.6 Perform the second and third single substitutions (following the process explained in 7.4.1.5) and record all the readings. Once the three comparisons are complete and the readings are recorded, calculate $C1_{DUT}$, $C2_{DUT}$, and $C3_{DUT}$.

NOTE 8—Provided the calibration environment is thermally stable, two or even three DUTs can be read before the second reading of the standard hydrometer is made.

7.4.1.7 $C1_{DUT}$, $C2_{DUT}$ and $C3_{DUT}$ should ideally be in close agreement. One manner to express the correlation of the three values is to calculate their standard deviation. Another manner may be to calculate the range of the values. A third possibility is to estimate the standard deviation from the range of the values by dividing the range by the square root of 3 (1.7321). It is not the intention of this standard to impose a process limit on the dispersion of values; however the user is reminded that the correlation of these values is an important contribution to the uncertainty calculation.

7.4.1.8 Calculate $FINALCORRECTION_{DUT}$, the average of $C1_{DUT}$, $C2_{DUT}$, and $C3_{DUT}$.

Corrected Reading of Standard	Reading of DUT Hydrometer	Correction for DUT Hydrometer
10.08	10.12	-0.04
10.10	10.15	-0.05
10.11	10.16	-0.05

7.4.1.9 Repeat section 7.4 for each calibration point on the DUT.

7.4.1.10 The $FINALCORRECTION_{DUT}$ values determined for a particular DUT at its three calibration points are data that may be presented on a calibration report.

7.4.1.11 *Verification*—if all three $FINALCORRECTION_{DUT}$ values for a particular DUT are equal to or smaller than the permitted scale error values for the DUT provided in Specification E100, then the claim may be made that the DUT complies with the maximum scale error requirements of Specification E100.

7.4.2 *Hydrometers for Liquids Having High Vapor Pressure* **Warning**—This is an extremely hazardous procedure that should only be carried out in a laboratory setting with appropriate equipment, trained personnel, and facilities for dealing with the highly flammable and potentially explosive vapors that are generated during this procedure.

7.4.2.1 A suggested comparator is described in X1.2. <https://standards.iteh.ai/catalog/standards/sist/180cb705-5a09-45bc-8040-9742e8032454/astm-e126-19>

7.4.2.2 Insert the hydrometer to be calibrated and the thermometer into the comparator and attach the cover plate. Connect the source of supply of propane or normal butane to the inlet valve and ascertain that the connections are free from leaks. Open the outlet valve and purge the connections by opening the inlet valve slightly.

7.4.2.3 When the connections have been purged, close the outlet and vent valves and open the inlet valve, permitting the liquid to enter the comparator, until it is completely full. If necessary, the vent valve may be opened slightly to permit complete filling, after which it should be closed.

7.4.2.4 When the comparator has been filled, close the inlet valve and open the outlet valve, permitting the liquid to be withdrawn completely and the pressure inside the comparator to be reduced to that of the ambient atmosphere.

7.4.2.5 Close the outlet valve and open the inlet valve, filling the comparator to a level at which the enclosed hydrometer floats freely. If the vapor pressure is too high to permit adequate filling, cool the comparator (see 7.4.2.6). This is most readily accomplished by repetition of the purging operation. When the comparator is adequately filled, close the inlet valve and examine the apparatus for leaks. If any are detected, withdraw the liquid, correct the leaky condition and repeat the purging and filling operations.

7.4.2.6 Disconnect the comparator from the source of supply of liquid and place it in a water bath maintained at $15.6 \pm 0.3^\circ\text{C}$ ($60.3 \pm 0.5^\circ\text{F}$) until thermal equilibrium has been attained. To accelerate thermal adjustment, occasionally remove the comparator from the water bath, tilt to a horizontal position, rock gently a few times to ensure mixing, and replace in the water bath. Exercise care to prevent damage to the hydrometer.

7.4.2.7 Remove the comparator from the water bath and make the hydrometer readings as quickly as possible, following the procedure outlined in 7.4.1.7 while the hydrometer is floating freely.

7.4.2.8 Withdraw the liquid and repeat the process with a second liquid sample. The two observations should be considered suspect if they differ by more than 0.0005 relative density (specific gravity). The two readings should be averaged in calculating the correction. Take the relative density (specific-gravity) value furnished by the producer of the comparison liquid being used as the true value for purposes of correction calculation.