



Designation: **D7153 – 15<sup>ε1</sup> D7153 – 22**



**IP 529/15/16**

## Standard Test Method for Freezing Point of Aviation Fuels (Automatic Laser Method)<sup>1,2</sup>

This standard is issued under the fixed designation D7153; 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 ( $\epsilon$ ) indicates an editorial change since the last revision or reapproval.

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<sup>ε1</sup> NOTE—Figures 3, A1.3, and A1.5 were corrected editorially in March 2016.

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### 1. Scope\*

1.1 This test method covers the determination of the temperature below which solid hydrocarbon crystals may form in aviation turbine fuels.

1.2 This test method is designed to cover the temperature range of  $-80^{\circ}\text{C}$  to  $20^{\circ}\text{C}$ ; however, the interlaboratory study mentioned in 12.4 has only demonstrated the test method with fuels having freezing points in the range of  $-60^{\circ}\text{C}$  to  $-42^{\circ}\text{C}$ .

1.3 The values stated in SI units are to be regarded as standard. No other units of measurement are included in this standard.

1.4 *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, health, and environmental practices and to determine the applicability of regulatory limitations prior to use.*

1.5 *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:<sup>3</sup>

**D2386 Test Method for Freezing Point of Aviation Fuels**

**D4057 Practice for Manual Sampling of Petroleum and Petroleum Products**

**D4177 Practice for Automatic Sampling of Petroleum and Petroleum Products**

#### 2.2 Energy Institute Standard:

**IP 16 Determination Freezing Point of Aviation Fuels<sup>4</sup>**

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<sup>1</sup> This test method is under the jurisdiction of ASTM International Committee D02 on Petroleum Products, Liquid Fuels, and Lubricants and is the direct responsibility of ASTM Subcommittee D02.07 on Flow Properties. The technically equivalent standard as referenced is under the jurisdiction of the Energy Institute Subcommittee SC-B-7. Current edition approved April 1, 2015 April 1, 2022. Published April 2015 April 2022. Originally approved in 2005. Last previous edition approved in 2010 as D7153 – 05 (2010); D7153 – 15<sup>ε1</sup>. DOI: 10.1520/D7153-15E01.10.1520/D7153-22.

<sup>2</sup> This test method has been developed through the cooperative effort between ASTM and the Energy Institute, London. ASTM and IP standards were approved by ASTM and EI technical committees as being technically equivalent but that does not imply both standards are identical.

<sup>3</sup> For referenced ASTM standards, visit the ASTM website, [www.astm.org](http://www.astm.org), or contact ASTM Customer Service at [service@astm.org](mailto:service@astm.org). For Annual Book of ASTM Standards volume information, refer to the standard's Document Summary page on the ASTM website.

<sup>4</sup> Annual Book of IP Standards Methods, Vol 1. Available from Energy Institute, 61 New Cavendish St., London, WIG 7AR, U.K.

\*A Summary of Changes section appears at the end of this standard

### 3. Terminology

#### 3.1 Definitions:

3.1.1 *freezing point, n—in aviation fuels*, the fuel temperature at which solid hydrocarbon crystals, formed on cooling, disappear when the temperature of the fuel is allowed to rise under specified conditions of test.

#### 3.2 Definitions of Terms Specific to This Standard:

3.2.1 *automatic laser method, n*—the procedures of automatically cooling a liquid aviation fuel specimen until solid hydrocarbon crystals appear, followed by controlled warming and recording of temperature at which hydrocarbon crystals completely redissolve into the liquid phase.

#### 3.3 Symbols:

*Cd* = the specimen temperature at which the appearance of the first crystals are detected in the specimen by an optical crystal detector under specified conditions of test.

*Co* = the specimen temperature at which the appearance of opacity in the specimen is detected by an optical opacity detector under specified conditions of test.

*Do* = the specimen temperature at which the disappearance of opacity in the specimen is detected by an optical opacity detector under specified conditions of test.

### 4. Summary of Test Method

4.1 A specimen is cooled at a rate of  $10\text{ }^{\circ}\text{C}/\text{min} \pm 5\text{ }^{\circ}\text{C}/\text{min}$  while continuously being illuminated by a laser light source. The specimen is continuously monitored by optical crystal and opacity detectors for the first formation of solid hydrocarbon crystals. Once the hydrocarbon crystals are detected by both sets of optical detectors, the specimen is then warmed at a rate of  $3\text{ }^{\circ}\text{C}/\text{min} \pm 0.5\text{ }^{\circ}\text{C}/\text{min}$ . When initial opacity in the specimen disappears, the specimen is then warmed at a rate of  $12\text{ }^{\circ}\text{C}/\text{min} \pm 1\text{ }^{\circ}\text{C}/\text{min}$ . The specimen temperature at which the last hydrocarbon crystals return to the liquid phase, as detected by the crystal detector, is recorded as the freezing point.

4.2 In certain circumstances, as measured by the apparatus, the specimen is reheated to approximately  $10\text{ }^{\circ}\text{C}$ , then cooled at the rate in 4.1 until hydrocarbon crystals are detected by the crystal detector. The specimen is then warmed at a rate of  $12\text{ }^{\circ}\text{C}/\text{min} \pm 1\text{ }^{\circ}\text{C}/\text{min}$ , until the last hydrocarbon crystals return to the liquid phase. The specimen temperature at which the last hydrocarbon crystals return to the liquid phase, as detected by the crystal detector, is recorded as the freezing point.

### 5. Significance and Use

5.1 The freezing point of an aviation fuel is the lowest temperature at which the fuel remains free of solid hydrocarbon crystals which, if present in the fuel system of the aircraft, can restrict the flow of fuel through filters. The temperature of the fuel in the aircraft tank normally decreases during flight depending on aircraft speed, altitude, and flight duration. The freezing point of the fuel shall always be lower than the minimum operational fuel temperature.

5.2 Petroleum blending operations require precise measurement of the freezing point.

5.3 This test method expresses results to the nearest  $0.1\text{ }^{\circ}\text{C}$ , and it eliminates most of the operator time and judgment required by Test Method **D2386**.

5.4 When a specification requires Test Method **D2386**, do not substitute this test method or any other test method.

### 6. Apparatus

6.1 *Automatic Apparatus*<sup>5</sup>—This apparatus consists of a microprocessor-controlled test cell that is capable of cooling and heating

<sup>5</sup> The sole source of supply of the apparatus known to the committee at this time is ISL model FZP 5G2s series Freezing Point Analyzer, available from PAC - ISL, BP 70285 - VERNON, 14653 CARPIQUET Cedex, France. If you are aware of alternative suppliers, please provide this information to ASTM International Headquarters. Your comments will receive careful consideration at a meeting of the responsible technical committee,<sup>1</sup> which you may attend.