

Designation: D7566 - 15

AnAmerican National Standard

Standard Specification for Aviation Turbine Fuel Containing Synthesized Hydrocarbons¹

This standard is issued under the fixed designation D7566; 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 specification covers the manufacture of aviation turbine fuel that consists of conventional and synthetic blending components.
- 1.2 This specification applies only at the point of batch origination, as follows:
- 1.2.1 Aviation turbine fuel manufactured, certified, and released to all the requirements of Table 1 of this specification (D7566), meets the requirements of Specification D1655 and shall be regarded as Specification D1655 turbine fuel. Duplicate testing is not necessary; the same data may be used for both D7566 and D1655 compliance. Once the fuel is released to this specification (D7566) the unique requirements of this specification are no longer applicable: any recertification shall be done in accordance with Table 1 of Specification D1655.
- 1.2.2 Field blending of synthesized paraffinic kerosine (SPK) blendstocks, as described in Annex A1 (FT SPK), Annex A2 (HEFA SPK), or Annex A3 (SIP) with D1655 fuel (which may on the whole or in part have originated as D7566 fuel) shall be considered batch origination in which case all of the requirements of Table 1 of this specification (D7566) apply and shall be evaluated. Short form conformance test programs commonly used to ensure transportation quality are not sufficient. The fuel shall be regarded as D1655 turbine fuel after certification and release as described in 1.2.1.
- 1.2.3 Once a fuel is redesignated as D1655 aviation turbine fuel, it can be handled in the same fashion as the equivalent refined D1655 aviation turbine fuel.
- 1.3 This specification defines specific types of aviation turbine fuel that contain synthesized hydrocarbons for civil use in the operation and certification of aircraft and describes fuels found satisfactory for the operation of aircraft and engines. The specification is intended to be used as a standard in describing the quality of aviation turbine fuels and synthetic blending

components at the place of manufacture but can be used to

- 1.4 This specification does not define the quality assurance testing and procedures necessary to ensure that fuel in the distribution system continues to comply with this specification after batch certification. Such procedures are defined elsewhere, for example in ICAO 9977, EI/JIG Standard 1530, JIG 1, JIG 2, API 1543, API 1595, and ATA-103.
- 1.5 This specification does not include all fuels satisfactory for aviation turbine engines. Certain equipment or conditions of use may permit a wider, or require a narrower, range of characteristics than is shown by this specification.
- 1.6 While aviation turbine fuels defined by Table 1 of this specification can be used in applications other than aviation turbine engines, requirements for such other applications have not been considered in the development of this specification.
- 1.7 Synthetic blending components, synthetic fuels, and blends of synthetic fuels with conventional petroleum-derived fuels in this specification have been evaluated and approved in accordance with the principles established in Practice D4054.
- 1.8 The values stated in SI units are to be regarded as standard. No other units of measurement are included in this standard.
- 1.9 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 and health practices and determine the applicability of regulatory limitations prior to use.

2. Referenced Documents

2.1 ASTM Standards:²

D56 Test Method for Flash Point by Tag Closed Cup Tester D86 Test Method for Distillation of Petroleum Products at

describe the quality of aviation turbine fuels for contractual transfer at all points in the distribution system.

1.4 This specification does not define the quality assurance

¹ This specification is under the jurisdiction of ASTM Committee D02 on Petroleum Products, Liquid Fuels, and Lubricants and is the direct responsibility of Subcommittee D02.J0.06 on Emerging Turbine Fuels.

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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.

- Atmospheric Pressure
- D93 Test Methods for Flash Point by Pensky-Martens Closed Cup Tester
- D129 Test Method for Sulfur in Petroleum Products (General High Pressure Decomposition Device Method)
- D130 Test Method for Corrosiveness to Copper from Petroleum Products by Copper Strip Test
- D156 Test Method for Saybolt Color of Petroleum Products (Saybolt Chromometer Method)
- D240 Test Method for Heat of Combustion of Liquid Hydrocarbon Fuels by Bomb Calorimeter
- D323 Test Method for Vapor Pressure of Petroleum Products (Reid Method)
- D381 Test Method for Gum Content in Fuels by Jet Evaporation
- D445 Test Method for Kinematic Viscosity of Transparent and Opaque Liquids (and Calculation of Dynamic Viscosity)
- D1266 Test Method for Sulfur in Petroleum Products (Lamp Method)
- D1298 Test Method for Density, Relative Density, or API Gravity of Crude Petroleum and Liquid Petroleum Products by Hydrometer Method
- D1319 Test Method for Hydrocarbon Types in Liquid Petroleum Products by Fluorescent Indicator Adsorption
- D1322 Test Method for Smoke Point of Kerosine and Aviation Turbine Fuel
- D1405 Test Method for Estimation of Net Heat of Combustion of Aviation Fuels
- D1655 Specification for Aviation Turbine Fuels
- D1840 Test Method for Naphthalene Hydrocarbons in Aviation Turbine Fuels by Ultraviolet Spectrophotometry
- D2276 Test Method for Particulate Contaminant in Aviation Fuel by Line Sampling
- D2386 Test Method for Freezing Point of Aviation Fuels
- D2425 Test Method for Hydrocarbon Types in Middle Distillates by Mass Spectrometry
- D2622 Test Method for Sulfur in Petroleum Products by Wavelength Dispersive X-ray Fluorescence Spectrometry
- D2624 Test Methods for Electrical Conductivity of Aviation and Distillate Fuels
- D2710 Test Method for Bromine Index of Petroleum Hydrocarbons by Electrometric Titration
- D2887 Test Method for Boiling Range Distribution of Petroleum Fractions by Gas Chromatography
- D2892 Test Method for Distillation of Crude Petroleum (15-Theoretical Plate Column)
- D3227 Test Method for (Thiol Mercaptan) Sulfur in Gasoline, Kerosine, Aviation Turbine, and Distillate Fuels (Potentiometric Method)
- D3240 Test Method for Undissolved Water In Aviation Turbine Fuels
- D3241 Test Method for Thermal Oxidation Stability of Aviation Turbine Fuels
- D3242 Test Method for Acidity in Aviation Turbine Fuel
- D3338 Test Method for Estimation of Net Heat of Combustion of Aviation Fuels

- D3343 Test Method for Estimation of Hydrogen Content of Aviation Fuels
- D3701 Test Method for Hydrogen Content of Aviation Turbine Fuels by Low Resolution Nuclear Magnetic Resonance Spectrometry
- D3828 Test Methods for Flash Point by Small Scale Closed Cup Tester
- D3948 Test Method for Determining Water Separation Characteristics of Aviation Turbine Fuels by Portable Separometer
- D4052 Test Method for Density, Relative Density, and API Gravity of Liquids by Digital Density Meter
- D4054 Practice for Qualification and Approval of New Aviation Turbine Fuels and Fuel Additives
- D4057 Practice for Manual Sampling of Petroleum and Petroleum Products
- D4171 Specification for Fuel System Icing Inhibitors
- D4176 Test Method for Free Water and Particulate Contamination in Distillate Fuels (Visual Inspection Procedures)
- D4294 Test Method for Sulfur in Petroleum and Petroleum Products by Energy Dispersive X-ray Fluorescence Spectrometry
- D4306 Practice for Aviation Fuel Sample Containers for Tests Affected by Trace Contamination
- D4529 Test Method for Estimation of Net Heat of Combustion of Aviation Fuels
- D4629 Test Method for Trace Nitrogen in Liquid Petroleum Hydrocarbons by Syringe/Inlet Oxidative Combustion and Chemiluminescence Detection
- D4809 Test Method for Heat of Combustion of Liquid Hydrocarbon Fuels by Bomb Calorimeter (Precision Method)
- D4865 Guide for Generation and Dissipation of Static Electricity in Petroleum Fuel Systems
- D4952 Test Method for Qualitative Analysis for Active Sulfur Species in Fuels and Solvents (Doctor Test)
- D4953 Test Method for Vapor Pressure of Gasoline and Gasoline-Oxygenate Blends (Dry Method)
- D5001 Test Method for Measurement of Lubricity of Aviation Turbine Fuels by the Ball-on-Cylinder Lubricity Evaluator (BOCLE)
- D5006 Test Method for Measurement of Fuel System Icing Inhibitors (Ether Type) in Aviation Fuels
- D5190 Test Method for Vapor Pressure of Petroleum Products (Automatic Method) (Withdrawn 2012)³
- D5191 Test Method for Vapor Pressure of Petroleum Products (Mini Method)
- D5291 Test Methods for Instrumental Determination of Carbon, Hydrogen, and Nitrogen in Petroleum Products and Lubricants
- D5452 Test Method for Particulate Contamination in Aviation Fuels by Laboratory Filtration
- D5453 Test Method for Determination of Total Sulfur in Light Hydrocarbons, Spark Ignition Engine Fuel, Diesel Engine Fuel, and Engine Oil by Ultraviolet Fluorescence

³ The last approved version of this historical standard is referenced on www.astm.org.

- D5972 Test Method for Freezing Point of Aviation Fuels (Automatic Phase Transition Method)
- D6045 Test Method for Color of Petroleum Products by the Automatic Tristimulus Method
- D6304 Test Method for Determination of Water in Petroleum Products, Lubricating Oils, and Additives by Coulometric Karl Fischer Titration
- D6379 Test Method for Determination of Aromatic Hydrocarbon Types in Aviation Fuels and Petroleum Distillates—High Performance Liquid Chromatography Method with Refractive Index Detection
- D6469 Guide for Microbial Contamination in Fuels and Fuel Systems
- D6866 Test Methods for Determining the Biobased Content of Solid, Liquid, and Gaseous Samples Using Radiocarbon Analysis
- D7042 Test Method for Dynamic Viscosity and Density of Liquids by Stabinger Viscometer (and the Calculation of Kinematic Viscosity)
- D7153 Test Method for Freezing Point of Aviation Fuels (Automatic Laser Method)
- D7154 Test Method for Freezing Point of Aviation Fuels (Automatic Fiber Optical Method)
- D7359 Test Method for Total Fluorine, Chlorine and Sulfur in Aromatic Hydrocarbons and Their Mixtures by Oxidative Pyrohydrolytic Combustion followed by Ion Chromatography Detection (Combustion Ion Chromatography-CIC)
- E29 Practice for Using Significant Digits in Test Data to Determine Conformance with Specifications
- 2.2 Energy Institute Standards:⁴
- EI/JIG 1530 Quality Assurance Requirements for the Manufacture, Storage and Distribution of Aviation Fuels to Airports
- IP 12 Determination of Specific Energy
- IP 16 Determination of the Freezing Point of Aviation Fuels—Manual Method
- IP 30 Detection of Mercaptans, Hydrogen Sulfide, Elemental Sulfur and Peroxides—Doctor Test Method
- IP 34 Determination of Flash Point—Pensky-Martens Closed Cup Method
- IP 69 Vapour Pressure-Reid Method (St-B-9)
- IP 71, Section 1 Petroleum Products—Transparent and Opaque Liquids—Determination of Kinematic Viscosity and Calculation of Dynamic Viscosity
- IP 123 Petroleum Products—Determination of Distillation Characteristics at Atmospheric Pressure
- IP 154 Petroleum Products—Corrosiveness to Copper— Copper Strip Test
- IP 156 Petroleum Products and Related Materials— Determination of Hydrocarbon Types—Fluorescent Indicator Adsorption Method
- IP 160 Crude Petroleum and Liquid Petroleum Products— Laboratory Determination of Density—Hydrometer Method
- ⁴ Available from Energy Institute, 61 New Cavendish St., London, WIG 7AR, U.K., http://www.energyinst.org.uk.

- IP 170 Determination of Flash Point—Abel Closed-Cup Method
- IP 216 Particulate Contaminant in Aviation Fuel
- IP 225 Determination of Copper in Light Petroleum Distillates—Spectrophotometric Method
- IP 227 Corrosiveness to Silver of Aviation Turbine Fuels— Silver Strip Method
- IP 274 Determination of Electrical Conductivity of Aviation and Distillate Fuels
- IP 299 Determination of Bromine Index—Electrometric Titration Method
- IP 323 Determination of Thermal Oxidation Stability of Gas Turbine Fuels
- IP 336 Petroleum Products—Determination of Sulfur Content—Energy-Dispersive X-ray Fluorescence Spectrometry
- IP 342 Petroleum Products—Determination of Thiol (Mercaptan) Sulfur in Light and Middle Distillate Fuels—Potentiometric Method
- IP 354 Determination of the Acid Number of Aviation Fuels-Colour-Indicator Titration Method
- IP 365 Crude Petroleum and Petroleum Products— Determination of Density—Oscillating U-tube Method
- IP 379 Determination of Organically Bound Trace Nitrogen—Oxidative Combustion and Chemiluminescence Method
- IP 394 Liquid Petroleum Products—Vapour Pressure—Part

 1: Determination of Air Saturated Vapour Pressure
 (ASVP) and Calculated Dry Vapour Pressure Equivalent
 (DVPE)
- IP 406 Petroleum Products—Determination of Boiling Range Distribution by Gas Chromatography
- IP 423 Determination of Particulate Contaminant in Aviation Turbine Fuels by Laboratory Filtration
- IP 435 Determination of the Freezing Point of Aviation Turbine Fuels by the Automatic Phase Transition Method
- IP 436 Determination of Aromatic Hydrocarbon Types in Aviation Fuels and Petroleum Distillates—High Performance Liquid Chromatography Method with Refractive Index Detection
- IP 438 Determination of Water—Coulometric Karl Fischer Titration Method
- IP 475 Petroleum Liquids—Manual Sampling
- IP 523 Determination of Flash Point—Rapid Equilibrium Closed Cup Method
- IP 524 Determination of Flash/No Flash—Rapid Equilibrium Closed Cup Method
- IP 528 Determination for the Freezing Point of Aviation Turbine Fuels—Automatic Fibre Optic Method
- IP 529 Determination of the Freezing Point of Aviation Fuels—Automatic Laser Method
- IP 540 Determination of the Existent Gum Content of Aviation Turbine Fuel—Jet Evaporation Method
- IP 585 Determination of Fatty Acid Methyl Esters (FAME),
 Derived from Bio-diesel Fuel, in Aviation Turbine Fuel—
 GC-MS with Selective Ion Monitoring/Scan Detection
 Method

IP 590 Determination of Fatty Acid Methyl Esters (FAME) in Aviation Turbine Fuel—HPLC Evaporative Light Scattering Detector Method

IP 598 Petroleum Products—Determination of the Smoke Point of Kerosine, Manual and Automated Method

2.3 ANSI Standard:⁵

ANSI 863 Report of Test Results

2.4 API Standards:6

API 1543 Documentation, Monitoring and Laboratory Testing of Aviation Fuel During Shipment from Refinery to Airport

API 1595 Design, Construction, Operation, Maintenance, and Inspection of Aviation Pre-Airfield Storage Terminals⁶

2.5 Joint Inspection Group Standards:⁷

JIG 1 Aviation Fuel Quality Control & Operating Standards for Into-Plane Fuelling Services

JIG 2 Aviation Fuel Quality Control & Operating Standards for Airport Depots & Hydrants⁷

2.6 IATA Guidance:⁸

9680–04 IATA Guidance Material on Microbiological Contamination in Aircraft Fuel Tanks

2.7 UOP Test Methods:9

UOP 389 Trace Metals in Oils by Wet Ash/ICP-AES

2.8 U.S. Department of Defense Specifications: 10

MIL-PRF-25017 Inhibitor, Corrosion/Lubricity Improver, Fuel Soluble

QDS-25017 Qualified Data Set for MIL-PRF-25017 (Inhibitor, Corrosion/Lubricity Improver, Fuel Soluble)

2.9 Other Standards:

ATA-103 Standard for Jet Fuel Quality Control at Airports¹¹
Defence Standard 91-91 Turbine Fuel, Aviation Kerosine
Type, Jet A-1¹²

ICAO 9977 Manual on Civil Aviation Jet Fuel Supply¹³
X001 Test Method for Determination of Farnesane, Saturated Hydrocarbons and Hexahydrofarnesol Content of Synthesized Iso-Paraffins (SIP) Fuels for Blending with Jet Fuel by Gas Chromatography¹⁴

AFRL-RQ-WP-TR-2013-0271 Determination of the Minimum Use Level of Fuel System Icing Inhibitor (FSII) in JP-8 that will Provide Adequate Icing Inhibition and Biostatic Protection for Air Force Aircraft¹⁵

3. General

3.1 This specification, unless otherwise provided, prescribes the required properties of aviation turbine fuel at the time and place of batch origination.

4. Terminology

- 4.1 Definitions:
- 4.1.1 *conventional hydrocarbons, n*—hydrocarbons derived from the following conventional sources: crude oil, natural gas liquid condensates, heavy oil, shale oil, and oil sands.
 - 4.2 Definitions of Terms Specific to This Standard:
- 4.2.1 *batch origination*, *n*—location at which fuel is certified as D7566.
- 4.2.2 *conventional blending component, n*—blending streams derived from conventional hydrocarbons.
- 4.2.3 hydroprocessed, adj—conventional chemical processing in which hydrogen is reacted with organic compounds in the presence of a catalyst to remove impurities such as oxygen, sulfur, nitrogen; to saturate unsaturated hydrocarbons; or to alter the molecular structure of the hydrocarbon molecules.
- 4.2.4 *identified incidental materials, n*—chemicals and compositions that have defined upper content limits in an aviation fuel specification but are not approved additives.
- 4.2.5 synthesized hydrocarbons, n—hydrocarbons derived from alternative sources such as coal, natural gas, biomass, and hydrogenated fats and oils by processes such as gasification, Fischer-Tropsch synthesis, and hydroprocessing.
- 4.2.6 synthetic blending component, n—synthesized hydrocarbons that meet the requirements of Annex A1, Annex A2, or Annex A3.
- 4.2.7 *synthesized iso-paraffins (SIP), n*—synthetic blending component that is comprised essentially of iso-paraffins.
- 4.2.8 synthesized paraffinic kerosine (SPK), n—synthetic blending component that is comprised essentially of isoparaffins, normal paraffins, and cycloparaffins.
- 4.2.8.1 *Discussion*—Trace materials are permitted provided they are components that normally occur in hydroprocessed jet fuel including, but not limited to, trace organics, nitrogen compounds, water, dissolved air, etc.

5. Classification

- 5.1 Two grades of aviation turbine fuels are provided, as follows:
- 5.1.1 *Jet A and Jet A-1*—Relatively high flash point distillates of the kerosine type.
- 5.2 Jet A and Jet A-1 represent two grades of kerosine fuel that differ in freezing point. Other grades would be suitably identified.

⁵ Available from American National Standards Institute (ANSI), 25 W. 43rd St., 4th Floor, New York, NY 10036, http://www.ansi.org.

⁶ Available from American Petroleum Institute (API), 1220 L. St., NW, Washington, DC 20005-4070, http://www.api.org.

⁷ Available from Joint Inspection Group (JIG), http://www.jigonline.com.

⁸ Available from International Air Transport Association (IATA). Head Office: 800 Place Victoria, PO Box 113, Montreal, H4Z 1M1, Quebec, Canada. Executive Office: 33, Route de l'Aeroport, PO Box 416, 1215 Geneva, 15 Airport, Switzerland. www.iata.org.

⁹ Available from ASTM International, www.astm.org, or contact ASTM Customer Service at service@astm.org.

¹⁰ Available from the Standardization Document Order Desk, 700 Robbins, Avenue, Building 4D, Philadelphia PA 19111-5094 (http://assist.daps.dla.mil).

¹¹ Available from Air Transport Association of America, Inc. (ATA) d/b/a Airlines for America, 1301 Pennsylvania Ave. NW, Suite 1100, Washington, D.C. 20004, http://www.airlines.org.

¹² Available from Defence Equipment and Support, UK Defence Standardization, Kentigern House, 65 Brown Street, Glasgow, G2 8EX (http://www.dstan.mod.uk).

¹³ Available from International Civil Aviation Organization (ICAO), 999 University St., Montreal, Quebec H3C 5H7, Canada, http://www.icao.int.

¹⁴ Available from Amyris, http://amyr.is/FarnesaneTestMethod, or contact Amyris at info@amyris.com.

¹⁵ Available from Defense Technical Information Center (DTIC), 8725 John J. Kingman Rd., Ft. Belvoir, VA 22060-6218, http://www.dtic.mil/dtic, accession number ADA595127.

6. Materials and Manufacture

- 6.1 Aviation turbine fuel, except as otherwise defined in this specification, shall consist of the following blends of components or fuels:
- 6.1.1 Conventional blending components or Jet A or Jet A-1 fuel certified to Specification D1655; with up to 50 % by volume of the synthetic blending component defined in Annex A1.
- 6.1.2 Conventional blending components or Jet A or Jet A-1 fuel certified to Specification D1655; with up to 50 % by volume of the synthetic blending component defined in Annex A2.

Note 1—The ability to add 50 % of Annex A1 or Annex A2 blending components (SPK) to Jet A or Jet A-1 is also limited by the physical properties of the fuel with which it is being blended. Practice has shown that density, or aromatic content, or both, of the refined fuel often limit the amount of SPK that can be added to the final blend to less than 50 %.

6.1.3 Conventional blending components or Jet A or Jet A-1 fuel certified to Specification D1655; with up to 10 % by volume of the synthetic blending component defined in Annex A3.

Note 2—The ability to add 10 % of Annex A3 blending components (SIP) to Jet A or Jet A-1 may also be limited by the physical properties of the fuel with which it is being blended. It is possible in extreme cases that viscosity of the refined fuel may limit the amount of SIP that can be added to the final blend to less than 10 %.

- 6.2 Fuels used in certified engines and aircraft are ultimately approved by the certifying authority subsequent to formal submission of evidence to the authority as part of the type certification program for that aircraft and engine model. Additives to be used as supplements to an approved fuel must also be similarly approved on an individual basis (see X1.2.4).
- 6.3 Additives—Only additives approved by the aviation industry (including the aircraft certifying authority) are permitted in the fuel on which an aircraft is operated. The additives approved for use in D7566 jet fuel are shown in Table 1 and Table 2 and may be used within the concentration limits shown in the tables subject to any restrictions described in the table footnotes. ¹⁶
- 6.4 Guidance material is presented in Appendix X3 concerning the need to control processing additives in jet fuel production.
- 6.5 From the point of manufacture to the point of blending to meet this specification, the synthetic blending component shall be handled and transported in the same manner as finished jet fuel in order to maintain product integrity. Appropriate management of change measures shall be used at manufacturing locations, distribution, and storage to maintain product integrity (see Appendix X3).

7. Detailed Requirements

7.1 The aviation turbine fuel shall conform to the requirements prescribed in Table 1 Part 1 and Table 1 Part 2 unless

¹⁶ Supporting data (Guidelines for Approval or Disapproval of Additives) have been filed at ASTM International Headquarters and may be obtained by requesting Research Report RR:D02-1125.

otherwise noted in 7.2, Annex A1, Annex A2, or Annex A3, whichever is applicable.

- 7.2 The fluidity requirement of Part 2 of Table 1 only applies to each batch of fuel containing the synthesized iso-paraffins (SIP) blending component as defined in Annex A3 and blended in accordance with 6.1.3. This requirement does not apply to fuel containing Annex A1 or Annex A2 synthesized components and blended in accordance with 6.1.1 or 6.1.2.
- 7.3 The additional requirements of Part 2 of Table 1 apply only for each batch of fuel intentionally containing a synthetic blending component. The additional requirements of Part 2 of Table 1 are not mandated if conventionally-derived jet fuel is mixed with the residue of a D7566 semi-synthetic aviation turbine fuel in refinery equipment from a previous batch of certified final blended product, for example in a tank heel.
- 7.4 Test results shall not exceed the maximum or be less than the minimum values specified in Table 1, Tables A1.1 and A1.2, Tables A2.1 and A2.2, and Tables A3.1 and A3.2. No allowance shall be made for the precision of the test methods. To determine conformance to the specification requirement, a test result may be rounded to the same number of significant figures as in Table 1, Tables A1.1 and A1.2, Tables A2.1 and A2.2, and Tables A3.1 and A3.2 using Practice E29. Where multiple determinations are made, the average result, rounded in accordance with Practice E29, shall be used.

8. Workmanship, Finish, and Appearance

8.1 The aviation turbine fuel specified in this specification shall be visually free of undissolved water, sediment, and suspended matter. The odor of the fuel shall not be nauseating or irritating. If the fuel has an odor similar to that of "rotten egg," please refer to X1.12.5 for further discussion. No substance of known dangerous toxicity under usual conditions of handling and use shall be present, except as permitted in this specification.

9. Sampling

- 9.1 Because of the importance of proper sampling procedures in establishing fuel quality, use the appropriate procedures in Practice D4057 or IP 475 to obtain a representative sample from the batch of fuel for specification compliance testing. This requirement is met by producing fuel as a discrete batch then testing it for specification compliance. This requirement is not satisfied by averaging online analysis results.
- 9.2 A number of jet fuel properties, including thermal stability, water separation, electrical conductivity, and others, are very sensitive to trace contamination, which can originate from sample containers. For recommended sample containers, refer to Practice D4306.

10. Report

10.1 The type and number of reports to ensure conformance with the requirements of this specification shall be mutually agreed upon by the seller and the purchaser of the aviation turbine fuel.

TABLE 1 Detailed Requirements of Aviation Turbine Fuels Containing Synthesized Hydrocarbons^A

Max		F	Part 1—Basic Requirements	
oidly, both my KOH's promises. When the following requirements all be met. Max	Property		Jet A or Jet A-1	Test Method ^B
Table met - Aconatics, volume percent - Max	COMPOSITION			
Acomatics, volume percent Max Acomatics, volume percent Max 2.6.5 0.3373P1 436 0.2027/P14924 0.30 0.2027/P14924 0.30 0.2027/P14924 0.2087/IP 406° or D86° or IP 123° 0.2087/IP 406° or IP 12			0.10	D3242/IP 354
Acomatics, volume percent Max 26.5 D8379H 4-96 Unif. relocation Times percent Max 0.003 D1266, 02622, D4294, D5453, or IP 336 D1266, D4262, D4294, D5453, D4294,		Max	25	D1319 or IP 156
Max				
Uniful to that image percent Max 0.30 D1266, D2622, D4294, D6453, or IP 336 LOATILITY D2887/IP 406° or D86" or IP 123" Unifoldion remperature (10) Max 205 10 % recovered, temperature (190) report 50 % recovered, temperature (190) Max 300 50 % recovered, temperature (190) Max 300 Final boiling port, resperature Max 300 Sellation residue, percent Max 15 Max 15 D456 or D3828°, IP 170° or IP 523° Benally at 15 °C, kg/m³ Min 36° D56 or D3828°, IP 170° or IP 523° Benally at 15 °C, kg/m³ Max 440 Jet A*1 D58728P-435, D7153VIP 529, D7154VIP 528 Becoatly ~20 °C, mm²/at² Max 427 Jet A*1" D445/P 71, Section 1, D7042* CMBUSTION Min 42.8" D4529, D3338, D4890 or IP 12 to the following requirements shall be metr. Max 3.0 D1840 Q2 Smoke point, mm, and may be precent of the following requirements with in the fill of the following requirements shall be metr. Max 3.0 D130/IP 154 LERIANAL STABLET				
Sellation	Sulfur, total mass percent			
10 % recovered, temperature (T10) 50 % recovered, temperature (T20) 90 % recovered, temperature (T90) 90 % r	VOLATILITY Distillation			D2887/IP 406 ^D or D86 ^E or IP 123 ^E
50 % recovered, temperature (TS0)	Distillation temperature, °C:			
90 % recovered, temperature (T90) Final boiling point, temperature Max	, ,	Max		
Final boling point, temperature				
Isitilation residue, percent				
Isitillation loss, percent and point, "C Min 38" D56 or D3828°, IP 170° or IP 523° ensity at 15 "C, kg/m" Min 38" D56 or D3828°, IP 170° or IP 523° ensity at 15 "C, kg/m" Max -40 Jet A" D8972/IP 435, D7153/IP 529, D7154/IP 528 D2366(IP 16) D445/IP 71, Section 1, D7042° D45/IP 71, Sec	Final boiling point, temperature	Max	300	
Description	Distillation residue, percent	Max	1.5	
LUIDITY	Distillation loss, percent	Max	1.5	
LUIDITY	Flash point, °C	Min	38 ^F	D56 or D3828 ^G , IP 170 ^G or IP 523 ^G
Max	Density at 15 °C, kg/m ³		775 to 840	
D2386/IP 16 D2386/IP 16 D2386/IP 16 D2386/IP 16 D2386/IP 17 Section 1, D7042 ^{-/-} DMBUSTION D445/IP 71, Section 1, D7042 ^{-/-} DMBUSTION D445/IP 71, Section 1, D7042 ^{-/-} D452/IP 27, Section 1, D7042 ^{-/-} D452/IP 28, D452/IP 29, D452/I	LUIDITY			
A	Freezing point, °C	Max	-40 Jet A ^H	D5972/IP 435, D7153/IP 529, D7154/IP 528,
Secosity - 20 °C, mm²/s² Max			_47 let Δ-1 ^H	D2386/IP 16
et heat of combustion, Mu/kg	/iscosity –20 °C, mm²/s/	Max		D445/IP 71, Section 1, D7042 ^J
et heat of combustion, Mu/kg	COMPLICTION			
(1) Smoke point, mm, or Min 18.0 D1322/IP 598 (2) Smoke point, mm, and Min 18.0 D1322/IP 598 (2) Smoke point, mm, and Max 3.0 D1322/IP 598 (2) Smoke point, mm, and Max 3.0 D1322/IP 598 (2) Smoke point, mm, and Max 3.0 D1322/IP 598 (2) Smoke point, mm, and Max 3.0 D1322/IP 598 (2) Smoke point, mm, and Max 3.0 D1322/IP 598 (2) Smoke point, mm, and Max 3.0 D1322/IP 598 (2) Smoke point, mm, and Max 3.0 D1322/IP 598 (2) Smoke point, mm, and Max 3.0 D130/IP 154 (2) Smoke point, mm, and Max 3.0 D130/IP 154 (2) Smoke point, mm, and Max 3.0 D130/IP 154 (2) Smoke point, mm, and Max 3.0 D130/IP 154 (2) Smoke point, mm, and Max 3.0 D130/IP 154 (2) Smoke point, mm, and Max 3.0 D130/IP 154 (2) Smoke point, mm, and Max 3.0 D130/IP 154 (2) Smoke point, mm, and Max 3.0 D130/IP 154 (2) Smoke point, mm, and Max 3.0 D130/IP 154 (2) Smoke point, mm, and Max 3.0 D130/IP 154 (2) Smoke point, mm, and Max 3.0 D130/IP 154 (2) Smoke point, mm, and Max 3.0 D130/IP 154 (2) Smoke point, mm, and Max 3.0 D130/IP 154 (2) Smoke point, mm, and Max 3.0 D130/IP 154 (2) Smoke point, mm, and Max 3.0 D130/IP 156 (2) Smoke point, mm, and Max 3.0 D1319 or IP 156 (2) Smoke point, mm, and Min Min 3.0 D1319 or IP 156 (2) Smoke point, mm, and Min Min 3.0 D1319 or IP 156 (2) Smoke point, mm, and Min Min 3.0 D1319 or IP 156 (2) Smoke point, mm, and Min Min 3.0 D1319 or IP 156 (2) Smoke point, mm, and Min Min 3.0 D1319 or IP 156 (2) Smoke point, mm, and Min Min 3.0 Smoke point, mm, and Min 3.0 Smoke point, mm, and Min 3.0 Smoke point, mm, and Min 3.0 Smoke point, mm, and Min 3.0 Smoke point, mm, and Min 3.0 Smoke point, mm, and Min 3.0 Sm	Net heat of combustion, MJ/kg		42.8 ^K	D4529, D3338, D4809 or IP 12
Naphthalenes, volume, percent			Standards	D4000/ID 500
Naphthalenes, volume, percent				
ORROSION Opper strip, 2 h at 100 °C Max No. 1 D130/IP 154 D3241 [£] /IP 323 [£] HERMAL STABILITY 5 h at control temperature of 260 °C, min Filter pressure drop, mm Hg Max 25 Tube rating: One of the following Interquirements shall be met. Astronomical color deposits (2) Annex A1 VTR, VTR Color Code (2) Annex A2 ITR or Annex A3 ETR, may over area of 2.5 mm² ONTAMINANTS With electrical conductivity additive With electrical conductivity additive Min Base Property Part 2—Extended Requirements Property Jet A or Jet A-1 Test Method® D3948 D1319 or IP 156 Acomatics, volume percent Min ^{P,O} Acomatics, volume percent Min ^{P,O} Min D2887/IP 406 ^O , D86 ^E or IP 123 ^E D5011 With circitry Jubricity, °M mm Max D386 D5001		Min		
Dispersion 2 h at 100 °C	Naphthalenes, volume, percent	Max	3.0	D1840
Dispersion 2 h at 100 °C	CORROCIONI			
### HERMAL STABILITY 5 h at control temperature of 260 °C, min Filter pressure drop, mm Hg		Max	No. 1	D130/IP 154
5 h at control temperature of 260 °C, min Filter pressure drop, mm Hg	55ppor 3mp, 2 m at 100 °C	Docum	ent Preview	D100/II 134
Filter pressure drop, mm Hg Max 25 Tube rating: One of the following prequirements shall be met. Max 85 (1) Annex A1 VTR, VTR Color Code Less than No peacock or abnormal color deposits (2) Annex A2 ITR or Annex A3 ETR, nm avg over area of 2.5 mm² ONTAMINANTS visitent gum, mg/100 mL Max 7 D381, IP 540 IDITIVES See 6.3 Idectrical conductivity additive Min 85 Property Set See 6.3 Idectrical conductivity, pS/m ODE A or Jet A-1 Test Method® ONDITION VOMADICS: One of the following re- vulnements shall be met: Aromatics, volume percent Min No 84 Aromatics, volume percent Min No 85 Aromatics, volume percent Min No 84 Aromatics, volume percent Min No 85 ONDITION VOMADICS: See 6.3 Betrical conductivity additive Min No 98 Property Jet A or Jet A-1 Test Method® ONDITION VOMADICS: One of the following re- vulnements shall be met: Aromatics, volume percent Min No 84 Aromatics, volume percent Min No 85 ONDITION VOMADICS: See 6.3 DE887/IP 408P , D86® or IP 123® ONDITION 150 USBNCITTY USBNCITTY USBNCITTY Min QR Max ONS ONS D5001	THERMAL STABILITY			
Tube rating: One of the following requirements shall be met." (1) Annex A1 VTR, VTR Color Code Less than No peacock or abnormal color deposits (2) Annex A2 ITR or Annex A3 ETR, Max 85 nm avg over area of 2.5 mm² ONTAMINANTS visitent gum, mg/100 mL Max 7 D381, IP 540 D3948 Without electrical conductivity additive Min 85 with electrical conductivity additive Min 70 DDITIVES Part 2—Extended Requirements Property Jet A or Jet A-1 Test Method® OMPOSITION Viromatics: One of the following requirements Min A D6379/IP 436 LAromatics, volume percent Min A D6379/IP 436 OLATILITY DEBOTTOR Min A Max 0.85 D5001	•		25	D3241 ^L /IP 323 ^L
Tube rating: One of the following requirements shall be met. Morphard Standard Stand				
No peacock or abnormal color deposits (2) Annex A2 ITR or Annex A3 ETR, Max 85 nm avg over area of 2.5 mm² ONTAMINANTS sistent gum, mg/100 mL Max 7 D381, IP 540 D3948 Without electrical conductivity additive Min 85 With electrical conductivity additive Min 70 DDITIVES See 6.3 D2624/IP 274 Part 2—Extended Requirements Property Jet A or Jet A-1 Test Method® COMPOSITION				
No peacock or abnormal color deposits (2) Annex A2 ITR or Annex A3 ETR, Max 85 nm avg over area of 2.5 mm² ONTAMINANTS sistent gum, mg/100 mL Max 7 D381, IP 540 D3948 Without electrical conductivity additive Min 85 With electrical conductivity additive Min 70 DDITIVES See 6.3 D2624/IP 274 Part 2—Extended Requirements Property Jet A or Jet A-1 Test Method® COMPOSITION		og/standards/sist/52	23b7a4b-bc82-4862-8eec-dcd	
Abnormal color deposits Abnormal color deposits	(1) Annex A1 VTR, VTR Color Code	Less than		
(2) Annex A2 ITR or Annex A3 ETR, may over area of 2.5 mm² ONTAMINANTS xistent gum, mg/100 mL xistent gum, mg/100			No peacock or	
ONTAMINANTS D381, IP 540 Size of the following re- purperness shall be met:			abnormal color deposits	
ONTAMINANTS xistent gum, mg/100 mL	(2) Annex A2 ITR or Annex A3 ETR,	Max	85	
Max 7 D381, IP 540				
D3948 Without electrical conductivity additive Min 85 With electrical conductivity additive Min 70 No No No No No No No N		Max	7	D381 IP 540
Without electrical conductivity additive Min 85 With electrical conductivity additive Min 70 DDITIVES See 6.3 Petrical conductivity, pS/m D2624/IP 274 Part 2—Extended Requirements Property Jet A or Jet A-1 Test Method [®] COMPOSITION veromatics: One of the following requirements shall be met:			•	
With electrical conductivity additive Min 70 DDITIVES See 6.3 Jectrical conductivity, pS/m D2624/IP 274 Part 2—Extended Requirements Property Jet A or Jet A-1 Test Method ^B COMPOSITION Aromatics: One of the following requirements shall be met: Aromatics, volume percent Min ^{P,Q} 8 D1319 or IP 156 Aromatics, volume percent Min ^{P,Q} 8.4 D6379/IP 436 VOLATILITY D2887/IP 406 ^D , D86 ^E or IP 123 ^E 50-T10, °C Min ^{Q,R} 15 90-T10, °C Min ^{Q,R} 15 UBBRICITY Unbricity, ON mm Max 0.85 D5001		Min	85	
Part 2—Extended Requirements Property Jet A or Jet A-1 Test Method ^B				
Part 2—Extended Requirements Property Jet A or Jet A-1 Test Method ^B	ADDITIVES		See 6.3	
Property Jet A or Jet A-1 Test Method B COMPOSITION Aromatics: One of the following re- puirements shall be met: . Aromatics, volume percent Min P,Q 2. Aromatics, volume percent Min P,Q 3. Aromatics, volume percent Min P,Q 4. Aromatics, volume percent Min P,Q 5. Aromatics, volume percent Min P,Q 6. Aromatics, volume percent Min P,Q 6. To Deserting the following re- puirements shall be met: . Aromatics, volume percent Min P,Q 7. Between the following re- puirements shall be met: . Aromatics, volume percent Min P,Q 7. Between the following re- puirements shall be met: 8. Deserting the following re- puirements shall be met: 8. Deserting the following re- puirements shall be met: 9.	Electrical conductivity, pS/m		0	D2624/IP 274
Property Jet A or Jet A-1 Test Method B COMPOSITION Aromatics: One of the following re- puirements shall be met: . Aromatics, volume percent Min P,Q 2. Aromatics, volume percent Min P,Q 3. Aromatics, volume percent Min P,Q 4. Aromatics, volume percent Min P,Q 5. Aromatics, volume percent Min P,Q 6. Aromatics, volume percent Min P,Q 6. To Deserting the following re- puirements shall be met: . Aromatics, volume percent Min P,Q 7. Between the following re- puirements shall be met: . Aromatics, volume percent Min P,Q 7. Between the following re- puirements shall be met: 8. Deserting the following re- puirements shall be met: 8. Deserting the following re- puirements shall be met: 9.		Do	et O. Estandad Daguiramanta	
COMPOSITION Aromatics: One of the following re- puirements shall be met: . Aromatics, volume percent Min P.Q 2. Aromatics, volume percent Min P.Q 3. Aromatics, volume percent Min P.Q 4. D6379/IP 436 D2887/IP 406 P, D86 or IP 123 or IP 156 D2887/IP 406 P, D86 or IP 123 or IP 156 D2887/IP 406 P, D86 or IP 123 or IP 123 or IP 156 D2887/IP 406 P, D86 or IP 123 or	Property	Pa	· · · · · · · · · · · · · · · · · · ·	Test Method ^B
Aromatics: One of the following re- puirements shall be met: . Aromatics, volume percent Min P,Q 8 D1319 or IP 156 . Aromatics, volume percent Min P,Q VOLATILITY D2887/IP 406P, D86F or IP 123F 50-T10, °C Min Q,R 15 90-T10, °C Min Q,R 15 15 15 15 10-T10, °C Min Q,R 15 15 15 15 15 15 15 15 15 15 15 15 15	COMPOSITION		331.7107 331.77	.co. mounou
. Aromatics, volume percent Min P,Q 8 D1319 or IP 156 2. Aromatics, volume percent Min P,Q 8.4 D6379/IP 436 . Aromatics, volume percent Min P,Q 8.4 D6379/IP 436 . D2887/IP 406 ^D , D86 ^E or IP 123 ^E . T50-T10, °C Min Q,R 15 . T90-T10, °C Min Q,R 40 . UBRICITY . Lubricity, Q,N mm Max 0.85 D5001	Aromatics: One of the following re-			
2. Aromatics, volume percent Min ^{P,Q} 2. Aromatics, volume percent Min ^{P,Q} 3. Description Descr	quirements shall be met:	7.0		
VOLATILITY Described by the control of the control				
Distillation D2887/IP 406 ^D , D86 ^E or IP 123 ^E 50-T10, °C Min ^{Q,R} 15 90-T10, °C Min ^{Q,R} 40 JUBRICITY Ubricity, ^{Q,N} mm Max 0.85 D5001	· •	n ^{P,Q}	8.4	D6379/IP 436
"50-T10, °C Min O,R 15 "90-T10, °C Min O,R 40 UBRICITY .ubricity, O,N mm Max 0.85 D5001	VOLATILITY			
90-T10, °C Min ^{Q,R} 40 .UBRICITY .ubricity, ^{Q,N} mm Max 0.85 D 5001	Distillation			D2887/IP 406^{D} , D86 ^E or IP 123^{E}
190-T10, °C Min Q,R 40 UBRICITY Ubricity, Q,N mm Max 0.85 D5001			15	
.UBRICITY .ubricity, ^{O,N} mm Max 0.85 D5001		n ^{Q,R}		
ubricity, ^{O,N} mm Max 0.85 D5001				
		ax	0.85	D5001
EVIDIT I		ar.	0.00	20001
/iscosity -40 °C, mm ² /s Max 12 D445 ^T /, Section 1 ^T		2.4	10	DA45T / Section 1T



- ^A For compliance of test results against the requirements of Table 1, see 7.3.
- ^B The test methods indicated in this table are referred to in Section 11.
- ^C The mercaptan sulfur determination may be waived if the fuel is considered sweet by the doctor test described in Test Method D4952 or IP 30.
- ^D Distillation property criteria are specified in D86 or IP 123 scale units. D2887/IP 406 results shall be converted to estimated D86 or IP 123 results by application of the correlation in Appendix X5 of D2887 or Annex G of IP 406 for comparison with the specified property criteria. Distillation residue and loss limits provide control of the distillation process during the D86 and IP 123 test methods and do not apply to D2887/IP 406. Distillation residue and loss shall be reported as "not applicable" (N/A) when reporting D2887/IP 406 results.
- ED86 or IP 123 distillation of jet fuel is run at Group 4 conditions, except Group 3 condenser temperature is used.
- F A higher minimum flash point specification may be agreed upon between purchaser and supplier.
- ^G Results obtained by other test methods can be up to 2 °C lower than those obtained by Test Method D56, which is the preferred method. In case of dispute, Test Method D56 will apply.
- HOther freezing points may be agreed upon between supplier and purchaser.
- 1 1 mm 2 /s = 1 cSt.
- 'Test Method D7042 results shall be converted to bias-corrected kinematic viscosity results by the application of the correction described in Test Method D7042, section 15.4.4.
- ^K For all grades use either Eq 1 or Table 1 in Test Method D4529 or Eq 2 in Test Method D3338 or IP 12. Test Method D4809 may be used as an alternative. In case of dispute, Test Method D4809 shall be used.
- ^L D3241/IP 323 Thermal Stability is a critical aviation fuel test, the results of which are used to assess the suitability of jet fuel for aviation operational safety and regulatory compliance. The integrity of D3241/IP 323 testing requires that heater tubes (test coupons) meet the requirements of D3241 Table 2 and give equivalent D3241 results to the heater tubes supplied by the original equipment manufacturer (OEM). A test protocol to demonstrate equivalence of heater tubes from other suppliers is on file at ASTM International Headquarters and can be obtained by requesting Research Report RR:D02-1550. Heater tubes and filter kits, manufactured by the OEM (PAC, 8824 Fallbrook Drive, Houston, TX 77064) were used in the development of the D3241/IP 323 test method. Heater tube and filter kits, manufactured by Falex (Falex Corporation, 1020 Airpark Dr., Sugar Grove, IL, 60554-9585) were demonstrated to give equivalent results (see D3241 for research report references). These historical facts should not be construed as an endorsement or certification by ASTM International.
- ^M Tube deposit ratings shall be measured by D3241 Annex A2 ITR or Annex A3 ETR, when available. If the Annex A2 ITR device reports "N/A" for a tube's volume measurement, the test shall be a failure and the value reported as >85 nm. Visual rating of the heater tube by the method in D3241 Annex A1 is not required when Annex A2 ITR or Annex A3 ETR deposit thickness measurements are reported. In case of dispute between results from visual and metrological methods, the referee shall be considered the Annex A3 ETR method if available, otherwise Annex A2 ITR.
- ^N At point of manufacture.
- O If electrical conductivity additive is used, the conductivity shall not exceed 600 pS/m at the point of use of the fuel. When electrical conductivity additive is specified by the purchaser, the conductivity shall be 50 pS/m to 600 pS/m under the conditions at point of delivery. (1 pS/m = 1 × 10⁻¹² Ω^{-1} m⁻¹)
- ^P Minimum aromatics contents are based on current experience with the approved synthetic fuels and those levels were established from what is typical for refined jet fuel. Research is ongoing on the actual need for aromatics.
- ^Q The minimum aromatics and distillation slope criteria only apply to aviation turbine fuels containing synthesized hydrocarbons produced to this specification and are not applicable to conventional aviation turbine fuels produced to Specification D1655. Some batches of aviation turbine fuels produced to Specification D1655 may not meet the minimum aromatics and distillation slope criteria specified in Table 1 of this specification.
- ¹⁷ These distillation slope limits are based on current experience with the approved synthetic fuels and these values were established from what is typical for refined jet fuel. Research is ongoing on the actual requirements for distillation slope.
- The fluidity requirement applies only to jet fuel containing synthesized iso-paraffins specified in Annex A3 and blended in accordance with 6.1.3. It does not apply to jet fuel containing Annex A1 or Annex A2 synthesized components blended in accordance with 6.1.1 or 6.1.2.
- TD445 or IP 71, Section 1 allows measuring the viscosity at -40 °C, however the precision values were determined down to -20 °C. Data correlating test results at -40 °C for D445 and other related ASTM test methods is provided in Research Report RR:D02-1776, Evaluation of Synthesized Iso-Paraffins produced from Hydroprocessed Fermented Sugars (SIP Fuels), prepared by TOTAL New Energies, Amyris, Inc. and the United States Air Force Research Laboratory (AFRL), Final Version, February 2014. A revision to Test Method D445 to specify measurement precision at -40 °C is in process.

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10.2 A suggested form for reporting inspection data on aviation turbine fuels is given in Appendix X4.

11. Test Methods

Note 3—Where IP test methods are referenced in this standard as alternatives to ASTM test methods, the following nomenclature is used. Where test methods are officially jointed, this is denoted as Dxxxx/IP xxx. Where test methods are technically equivalent or related but not officially jointed, this is denoted as Dxxxx or IP xxx.

- 11.1 Determine the requirements enumerated in this specification in accordance with the following test methods.
- 11.1.1 *Density*—Test Method D1298/IP 160 or D4052 or IP 365.
- 11.1.2 *Distillation*—Test Method D86 or IP 123. For Jet A and Jet A-1, Test Method D2887/IP 406 may be used as an alternate. Results from Test Method D2887/IP 406 shall be reported as estimated D86 or IP 123 results by application of the correlation in Appendix X5 of D2887 or Annex G of IP 406. In case of dispute, Test Method D86 shall be the referee method (see X1.6.1.1).
- 11.1.3 Flash Point—Test Method D56, D3828, IP 170 or IP 523.
- 11.1.4 Freezing Point—Test Method D5972/IP 435, D7153/IP 529, D7154/IP 528, or D2386/IP 16. Any of these test methods may be used to certify and recertify jet fuel.

However, Test Method D2386/IP 16 is the referee method. An interlaboratory study (RR:D02-1572¹⁷) that evaluated the ability of freezing point methods to detect jet fuel contamination by diesel fuel determined that Test Methods D5972/IP 435 and D7153/IP 529 provided significantly more consistent detection of freeze point changes caused by contamination than Test Methods D2386/IP 16 and D7154/IP 528. It is recommended to certify and recertify jet fuel using either Test Method D5972/IP 435 or Test Method D7153/IP 529, or both, on the basis of the reproducibility and cross-contamination detection reported in RR:D02-1572.¹⁷ The cause of freezing point results outside specification limits by automated methods should be investigated, but such results do not disqualify the fuel from aviation use if the results from the referee method (Test Method D2386/IP 16) are within the specification limit.

11.1.5 *Viscosity*—Test Method D445/IP 71, Section 1 or Test Method D7042. Results from Test Method D7042 shall be reported as bias-corrected kinematic viscosity results by application of the correction in Test Method D7042, subsection

¹⁷ Supporting data have been filed at ASTM International Headquarters and may be obtained by requesting Research Report RR:D02-1572. Contact ASTM Customer Service at service@astm.org.

TABLE 2 Detailed Requirements for Additives in Aviation Turbine Fuels

Additive	Dosage	
Fuel Performance Enhancing Additives		
Antioxidants ^{A,B}	24.0 mg/L max ^C	
One of the following:		
2,6 ditertiary-butyl phenol		
2,6 ditertiary-butyl-4-methyl phenol		
2,4 dimethyl-6-tertiary-butyl-phenol		
75 % minimum, 2,6 ditertiary-butyl phenol plus		
25 % maximum mixed tertiary and tritertiary butyl-phenols		
55 % minimum 2,4 dimethyl-6-tertiary-butyl phenol plus		
15 % minimum 2,6 ditertiary-butyl-4-methyl phenol,		
remainder as monomethyl and dimethyl tertiary-butyl phenols		
72 % minimum 2,4 dimethyl-6-tertiary-butyl phenol plus		
28 % maximum monomethyl and dimethyl-tertiary-butyl-phenols		
Metal Deactivator ^A		
N,N-disalicylidene-1,2-propane diamine		
On initial blending	2.0 mg/L max ^{C,D}	
After field reblending cumulative concentration	5.7 mg/L max	
Fuel System Icing Inhibitor ^{E,F}	0.07 % by volume min ^G	
Diethylene Glycol Monomethyl Ether (see Specification D4171 Type III)	0.15 % by volume max	
Fuel Handling and Maintenance Additives		
Electrical Conductivity Improver ^H		
Stadis 450 ⁷		
On initial blending	3 mg/L max	
After field reblending, cumulative concentration	5 mg/L max	
If the additive concentration is unknown at time of retreatment,		
additional concentration is restricted to 2 mg/L max		
Leak Detection Additive	1 mg/kg max	
Tracer A (LDTA-A) ^J		
Teh Standards		
Biocidal Additives-1, 4		
Biobor JF^M		
Kathon FP1.5 ^N		
Kathon FP1.5 ^N Corrosion Inhibitor/Lubricity Improvers ^O S S S S S S S S S S S S S S S S S S S		
One of the following:		
HiTEC 580 Innospec DCI-4A Document Preview	23 mg/L max	
	23 mg/L max	
Nalco 5403	23 mg/L max	

^A The active ingredient of the additive must meet the composition specified.

15.4.4, Relative Bias for jet fuel. In case of dispute, Test Method D445 shall be the referee method.

11.1.6 *Net Heat of Combustion*—Test Method D4529, D3338, D4809, or IP 12.

11.1.7 Corrosion (Copper Strip)—Test Method D130/IP 154.

11.1.8 Total Acidity—Test Method D3242/IP 354.

11.1.9 *Sulfur*—Test Method D1266, D2622, D4294, D5453, or IP 336.

11.1.10 Mercaptan Sulfur—Test Method D3227/IP 342.

11.1.11 Microseparometer—Test Method D3948.

11.1.12 *Existent Gum*—Test Method D381 or IP 540. Test Method D381, using steam jet operating conditions, shall be the referee test method.

^B Supporting data have been filed at ASTM International Headquarters and may be obtained by requesting Research Report RR:D02-1125.

^C Active ingredient (not including weight of solvent).

D If copper contamination is suspected, initial treatment may exceed 2.0 mg/L but cumulative total must be below 5.7 mg/L.775be/astm-d7566-15

^E Quantity must be declared by the fuel supplier and agreed to by the purchaser.

^F DiEGME content can by analyzed by Test Method D5006.

^G The lower FSII concentration limit allowable in Jet Fuel is based on research by the US Air Force as documented in report AFRL-RQ-WP-TR-2013-0271. Some engines and aircraft as certificated require higher minimum concentrations of icing inhibitor than the lower limit in this Jet Fuel specification. When fueling an aircraft, the fuel should be additized to the concentration levels specified in the appropriate engine and aircraft manual

be additized to the concentration levels specified in the appropriate engine and aircraft manual.

He flectrical conductivity improver is used, the conductivity shall not exceed 600 pS/m at the point of use of the fuel. When electrical conductivity additive is specified by the purchaser, the conductivity shall be 50 pS/m to 600 pS/m under the conditions at point of delivery. (1 pS/m = 1 × 10⁻¹² Ω^{-1} m⁻¹)

¹ Stadis 450 is a registered trademark marketed by Innospec Inc., Innospec Manufacturing Park, Oil Sites Road, Ellesmere Port, Cheshire, CH65 4EY, UK.

^JTracer A (LDTA-A) is a registered trademark of Tracer Research Corp., 3755 N. Business Center Dr., Tucson, AZ 85705.

^K Biocidal additives are available for controlled usage. Where such an additive is used in the fuel, the approval status of the additive and associated conditions must be checked for the specific aircraft and engines to be operated.

Lefer to the Aircraft Maintenance Manual (AMM) to determine if either biocide is approved for use and for their appropriate use and dosage.

^M Biobor JF is a registered trademark of Hammonds Technical Services, Inc., 910 Rankin Rd., Houston, TX 77073.

^N KATHON is a trademark of The Dow Chemical Company ("Dow") or an affiliated company of Dow, 2030 Dow Center, Midland, MI 48674. HiTEC 580 is a trademark of Afton Chemical Corp., 500 Spring St., Richmond, VA 23219. Innospec DCI-4A is available from Innospec Inc., Innospec Manufacturing Park, Oil Sites Road, Ellesmere Port, Cheshire, CH65 4EY, UK.

O More information concerning minimum treat rates of corrosion inhibitor/lubricity improver additives is contained in X1.10.2.



- 11.1.13 Thermal Stability—Test Method D3241/IP 323.
- 11.1.14 *Aromatics*—Test Method D1319, IP 156 or D6379/IP 436. Test Method D1319 shall be the referee test method.
 - 11.1.15 Smoke Point—Test Method D1322/IP 598.
 - 11.1.16 Naphthalene Content—Test Method D1840.
- 11.1.17 *Electrical Conductivity*—Test Method D2624 / IP 274.

12. Keywords

12.1 aviation turbine fuel; avtur; Jet A; Jet A-1; jet fuel; synthesized hydrocarbons; synthesized iso-paraffins; synthesized paraffinic kerosine; synthetic blending component; turbine fuel

ANNEXES

(Mandatory Information)

A1. FISCHER-TROPSCH HYDROPROCESSED SYNTHESIZED PARAFFINIC KEROSINE

A1.1 Scope

- A1.1.1 This annex defines hydroprocessed synthesized paraffinic kerosine (SPK) for use as a synthetic blending component in aviation turbine fuels for use in civil aircraft and engines. The specifications in this annex can be used for contractual exchange of synthetic blending components.
- A1.1.2 The synthetic blending components defined in this annex are not satisfactory for aviation turbine engines unless blended with conventional fuel or conventional blending components in accordance with the limitations described in 6.1.1.
- A1.1.3 The values stated in SI units are to be regarded as standard. No other units of measurement are included in this standard.

A1.2 General

A1.2.1 All requirements of the main body of this specification apply except as detailed in this annex.

A1.3 Terminology

- A1.3.1 Definitions of Terms Specific to This Annex:
- A1.3.1.1 Fischer-Tropsch hydroprocessed synthesized paraffinic kerosine (FT-SPK), n—SPK produced from one or more precursors synthesized by Fischer-Tropsch processing.

A1.4 Materials and Manufacture

- A1.4.1 FT-SPK synthetic blending components shall be comprised of hydroprocessed synthesized paraffinic kerosine wholly derived from:
- A1.4.1.1 Paraffins and olefins derived from synthesis gas via the Fischer-Tropsch (FT) process using Iron or Cobalt catalyst.
- A1.4.1.2 Subsequent processing of the product shall include hydrotreating, hydrocracking, or hydroisomerization and is expected to include, but not be limited to, a combination of

other conventional refinery processes such as polymerization, isomerization, and fractionation. 18

A1.5 Detailed Batch Requirements

- A1.5.1 Each batch of synthetic blending component shall conform to the requirements prescribed in Table A1.1.
- A1.5.2 *Test Methods*—Determine the requirements enumerated in this annex in accordance with the following test methods.
- A1.5.2.1 *Density*—Test Method D1298/IP 160, D4052 or IP
- A1.5.2.2 *Distillation*—Test Methods D86 or IP 123, and D2887/IP 406.
- A1.5.2.3 Flash Point—Test Method D56, D3828, IP 170 or IP 523.

A1.5.2.4 Freezing Point—Test Method D5972/IP 435, D7153/IP 529, D7154/IP 528, or D2386/IP 16. Any of these test methods may be used to certify and recertify jet fuel. However, Test Method D2386/IP 16 is the referee method. An interlaboratory study (RR:D02-157217) that evaluated the ability of freezing point methods to detect jet fuel contamination by diesel fuel determined that Test Methods D5972/IP 435 and D7153/IP 529 provided significantly more consistent detection of freeze point changes caused by contamination than Test Methods D2386/IP 16 and D7154/IP 528. It is recommended to certify and recertify jet fuel using either Test Method D5972/IP 435 or Test Method D7153/IP 529, or both, on the basis of the reproducibility and cross-contamination detection reported in RR:D02-1572.¹⁷ The cause of freezing point results outside specification limits by automated methods should be investigated, but such results do not disqualify the fuel from aviation use if the results from the referee method (Test Method D2386/IP 16) are within the specification limit.

¹⁸ Coordinating Research Council (CRC) Report, "Comparative Evaluation of Semi-Synthetic Jet Fuels," September 2008, provides a more detailed description of the composition and performance of FT-SPK blending components that evolved from the evaluation of representative samples of these blending components.



A1.5.2.5 *Total Acidity*—Test Method D3242/IP 354. A1.5.2.6 *Thermal Stability*—Test Method D3241/IP 323.

A1.6 Other Detailed Requirements

A1.6.1 The hydroprocessed SPK blend component shall meet the requirements of Table A1.2. It is not necessary to analyze each batch of hydroprocessed SPK for compliance with Table A1.2 once it is demonstrated that the process scheme is adequately controlled to support the expectation that these requirements are always met. At a minimum, significant changes in production operations shall be cause for recertifying that these limits continue to be met.

A1.6.2 *Test Methods*—Determine the requirements enumerated in this annex in accordance with the following test methods.

A1.6.2.1 Cycloparaffins—Test Method D2425.

A1.6.2.2 Aromatics—Test Method D2425.

A1.6.2.3 Paraffins—Test Method D2425.

A1.6.2.4 Carbon and Hydrogen—Test Method D5291.

A1.6.2.5 Nitrogen—Test Method D4629/IP 379.

A1.6.2.6 Water—Test Method D6304 or IP 438.

A1.6.2.7 *Sulfur*—Test Methods D5453 or D2622. Either of these test methods can be used to certify and recertify jet fuel. However, Test Method D5453 is the referee method.

A1.6.2.8 Metals—Test Method UOP 389.

A1.6.2.9 Halogens—Test Method D7359.

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