



Standard Specification for Hydrocarbon Unleaded Aviation Gasoline¹

This standard is issued under the fixed designation D7547; 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.

1. Scope*

1.1 This specification covers formulating specifications for purchases of aviation gasoline under contract and is intended primarily for use by purchasing agencies.

1.2 Unleaded aviation gasoline defined by this specification is for use in engines and associated aircraft that are specifically approved by the engine and aircraft manufacturers. This fuel is not considered suitable for use in other engines and associated aircraft that are certified to use only leaded aviation gasolines of the same octane grade.

1.3 This specification, unless otherwise provided, prescribes the required properties of unleaded aviation gasoline at the time and place of delivery.

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

1.5 *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.*

1.6 *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

D86 Test Method for Distillation of Petroleum Products and Liquid Fuels at Atmospheric Pressure

- D130 Test Method for Corrosiveness to Copper from Petroleum Products by Copper Strip Test**
- D323 Test Method for Vapor Pressure of Petroleum Products (Reid Method)**
- D873 Test Method for Oxidation Stability of Aviation Fuels (Potential Residue Method)**
- D1094 Test Method for Water Reaction of Aviation Fuels**
- D1298 Test Method for Density, Relative Density, or API Gravity of Crude Petroleum and Liquid Petroleum Products by Hydrometer Method**
- D2386 Test Method for Freezing Point of Aviation Fuels**
- 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**
- D2700 Test Method for Motor Octane Number of Spark-Ignition Engine Fuel**
- D3237 Test Method for Lead in Gasoline by Atomic Absorption Spectroscopy**
- D3338 Test Method for Estimation of Net Heat of Combustion of Aviation Fuels**
- D4052 Test Method for Density, Relative Density, and API Gravity of Liquids by Digital Density Meter**
- D4057 Practice for Manual Sampling of Petroleum and Petroleum Products**
- D4171 Specification for Fuel System Icing Inhibitors**
- D4177 Practice for Automatic Sampling of Petroleum and Petroleum Products**
- 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**
- 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**
- D5006 Test Method for Measurement of Fuel System Icing Inhibitors (Ether Type) in Aviation Fuels**
- D5059 Test Methods for Lead in Gasoline by X-Ray Spectroscopy**
- D5191 Test Method for Vapor Pressure of Petroleum Products (Mini Method)**

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.J0.02 on Spark and Compression Ignition Aviation Engine Fuels.

Current edition approved May 1, 2017. Published July 2017. Originally approved in 2009. Last previous edition approved in 2015 as D7547 – 15^{ε1}. DOI: 10.1520/D7547-17.

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.

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

D6469 Guide for Microbial Contamination in Fuels and Fuel Systems

E29 Practice for Using Significant Digits in Test Data to Determine Conformance with Specifications

3. Terminology

3.1 Definitions:

3.1.1 *unleaded aviation gasoline, n*—gasoline possessing specific properties suitable for fueling aircraft powered by reciprocating spark ignition engines, where lead is not intentionally added for the purpose of enhancing octane performance.

3.1.1.1 *Discussion*—Principal properties include volatility limits, stability, detonation-free performance in the engine for which it is intended, and suitability for low temperature performance.

4. Classification

4.1 Two grades of unleaded aviation gasoline are provided, known as: Grades UL91 and UL94.³

NOTE 1—Grades UL91 and UL94 are based on their octane number as measured by Test Method **D2700** motor method.

5. Materials and Manufacture

5.1 Unleaded aviation gasoline, except as otherwise specified in this specification, shall consist of blends of refined hydrocarbons derived from crude petroleum, natural gasoline, or blends, thereof, with synthetic hydrocarbons or aromatic hydrocarbons, or both.

5.2 *Additives*—These may be added to each grade of unleaded aviation gasoline in the amount and of the composition specified in the following list of approved materials. The quantities and types shall be declared by the manufacturer. Additives added after the point of manufacture shall also be declared.

5.2.1 *Antioxidants*—The following oxidation inhibitors may be added to the gasoline separately, or in combination, in total concentration not to exceed 12 mg of inhibitor (not including weight of solvent) per litre of fuel.

5.2.1.1 2,6-ditertiary butyl-4-methylphenol.

5.2.1.2 2,4-dimethyl-6-tertiary butylphenol.

5.2.1.3 2,6-ditertiary butylphenol.

5.2.1.4 75 % minimum 2,6-ditertiary butylphenol plus 25 % maximum mixed tertiary and tritertiary butylphenols.

5.2.1.5 75 % minimum di- and tri-isopropyl phenols plus 25 % maximum di- and tri-tertiary butylphenols.

5.2.1.6 72 % minimum 2,4-dimethyl-6-tertiary butylphenol plus 28 % maximum monomethyl and dimethyl tertiary butylphenols.

5.2.1.7 N,N'-di-isopropyl-para-phenylenediamine.

5.2.1.8 N,N'-di-secondary-butyl-para-phenylenediamine.

5.2.2 *Fuel System Icing Inhibitor (FSII)*—One of the following may be used:

5.2.2.1 *Isopropyl Alcohol (IPA, propan-2-ol)*, in accordance with the requirements of Specification **D4171** (Type II). May be used in concentrations recommended by the aircraft manufacturer when required by the aircraft owner/operator.

NOTE 2—Addition of isopropyl alcohol (IPA) can reduce knock ratings below minimum specification values (see **X1.2.3**).⁴

5.2.2.2 *Di-Ethylene Glycol Monomethyl Ether (Di-EGME)*, conforming to the requirements of Specification **D4171** (Type III) may be used in concentrations of 0.10 % volume to 0.15 % volume when required by the aircraft owner/operator.

5.2.2.3 Test Method **D5006** may be used to determine the concentration of Di-EGME in aviation fuels.

5.2.3 *Electrical Conductivity Additive*—Stadis 450⁵ in concentrations up to 3 mg/L is permitted. When loss of fuel conductivity necessitates retreatment with electrical conductivity additive, further addition is permissible up to a maximum cumulative level of 5 mg/L of Stadis 450.⁵

5.2.4 *Corrosion Inhibitor Additive*—The following corrosion inhibitors may be added to the gasoline in concentrations not to exceed the maximum allowable concentration (MAC) listed for each additive.

DCI-4A	MAC = 24 g/m ³
DCI-6A	MAC = 15 g/m ³
HITEC 580	MAC = 22.5 g/m ³
NALCO 5403	MAC = 22.5 g/m ³
NALCO 5405	MAC = 11.0 g/m ³
UNICOR J	MAC = 22.5 g/m ³
SPEC-AID 8Q22	MAC = 24.0 g/m ³
TOLAD 351	MAC = 24.0 g/m ³
TOLAD 4410	MAC = 22.5 g/m ³

6. Detailed Requirements

6.1 The unleaded aviation gasoline shall conform to the requirements prescribed in **Table 1**.

6.2 Test results shall not exceed the maximum or be less than the minimum values specified in **Table 1**. No allowance shall be made for the precision of the test methods. To determine the conformance to the specification requirement, a test result may be rounded to the same number of significant figures as in **Table 1** using Practice **E29**. Where multiple determinations are made, the average result, rounded according to Practice **E29**, shall be used.

7. Workmanship, Finish, and Appearance

7.1 The unleaded aviation gasoline specified in this specification shall be free from undissolved water, sediment, and suspended matter. The odor of the fuel shall not be nauseating or irritating. No substances of known dangerous toxicity under usual conditions of handling and use shall be present.

8. Sampling

8.1 Because of the importance of proper sampling procedures in establishing fuel quality, use the appropriate procedures in Practice **D4057** or Practice **D4177**.

³ Supporting data have been filed at ASTM International Headquarters and may be obtained by requesting Research Report RR:D02-1859. Contact ASTM Customer Service at service@astm.org.

⁴ Supporting data have been filed at ASTM International Headquarters and may be obtained by requesting Research Report RR:D02-1526.

⁵ Stadis is a registered trademark marketed by Innospec, Inc., Innospec Manufacturing Park, Oil Sites Road, Ellesmere Port, Cheshire, CH65 4EY, UK.

TABLE 1 Detailed Requirements for Unleaded Aviation Gasolines^A

Property		Grade UL91	Grade UL94	ASTM Test Method ^B
COMBUSTION				
Net heat of combustion, MJ/kg ^C	min	43.5	43.5	D4529 or D3338
Octane Rating				
Knock value, Motor Octane Number ^D	min	91.0	94.0	D2700
COMPOSITION				
Sulfur, mass percent	max	0.05	0.05	D2622
Tetraethyl lead, g Pb/L	max	0.0130	0.0130	D3237 or D5059
Requirements for All Grades				
VOLATILITY				
Vapor pressure, 38 °C, kPa	min		38.0	D323 or D5191 ^E
	max		49.0	
Density at 15 °C, kg/m ³			Report	D1298 or D4052
Distillation, °C				D86
Initial boiling point			Report	
Fuel Evaporated				
10 volume percent at °C	max		75	
40 volume percent at °C	min		75	
50 volume percent at °C	max		105	
90 volume percent at °C	max		135	
Final boiling point	max		170	
Sum of 10 % + 50 % evaporated	tem- peratures	min	135	
Recovery volume percent	min		97	
Residue volume percent	max		1.5	
Loss volume percent	max		1.5	
FLUIDITY				
Freezing point, °C	max		-58 ^F	D2386
CORROSION				
Copper strip, 2 h at 100 °C	max		No. 1	D130
CONTAMINANTS				
Oxidation stability, mg/100 mL (5 h aging) ^G				D873
Potential gum	max		6	
Lead precipitate	max		3	
Water reaction				D1094
Volume change, mL	max		±2	
OTHER				
Electrical conductivity, pS/m	max		450 ^H	D2624

^A For compliance of test results against the requirements of Table 1, see 6.2.

^B The test methods indicated in this table are referred to in Section 10.

^C For all grades use either Eq 1 or Table 1 in Test Method D4529 or Eq 2 in Test Method D3338. Test Method D4809 may be used as an alternative. In case of dispute, Test Method D4809 shall be used.

^D Knock ratings shall be reported to the nearest 0.1 octane/performance number.

^E Test Method D5191 shall be the referee vapor pressure method.

^F If no crystals have appeared on cooling to -58 °C, the freezing point may be reported as less than -58 °C.

^G If mutually agreed upon between the purchaser and the supplier, a 16 h aging gum requirement may be specified instead of the 5 h aging gum test; in such case the gum content shall not exceed 10 mg/100 mL and the visible lead precipitate shall not exceed 4 mg/100 mL. In such fuel the permissible antioxidant shall not exceed 24 mg/L.

^H Applies only when an electrical conductivity additive is used; when a customer specifies fuel containing conductivity additive, the following conductivity limits shall apply under the condition at point of use:

Minimum 50 pS/m

Maximum 450 pS/m.

The supplier shall report the amount of additive added.

8.1.1 Although automatic sampling following Practice D4177 may be useful in certain situations, initial refinery specification compliance testing shall be performed on a sample taken following procedures in Practice D4057.

8.2 A number of unleaded aviation gasoline properties, including copper corrosion, electrical conductivity, and others are very sensitive to trace contamination which can originate

from sample containers. For recommended sample containers, refer to Practice [D4306](#).

9. Report

9.1 The type and number of reports to ensure conformance with the requirements of this specification shall be mutually agreed to by the purchaser and the supplier of the unleaded aviation gasoline.

10. Test Methods

10.1 The requirements enumerated in this specification shall be determined in accordance with the following ASTM test methods:

10.1.1 *Knock Value (MON)*—Test Method [D2700](#).

10.1.2 *Density*—Test Methods [D1298](#) or [D4052](#).

10.1.3 *Distillation*—Test Method [D86](#).

10.1.4 *Vapor Pressure*—Test Methods [D323](#) or [D5191](#).

10.1.5 *Freezing Point*—Test Method [D2386](#).

10.1.6 *Sulfur*—Test Method [D2622](#).

10.1.7 *Net Heat of Combustion*—Test Methods [D4529](#), [D3338](#), or [D4809](#).

10.1.8 *Corrosion (Copper Strip)*—Test Method [D130](#), 2 h test at 100 °C in bomb.

10.1.9 *Potential Gum*—Test Method [D873](#) except that wherever the letter X occurs (referring to oxidation time) insert the number 5, designating the number of hours prescribed in this specification.

10.1.10 *Water Reaction*—Test Method [D1094](#).

10.1.11 *Electrical Conductivity*—Test Method [D2624](#).

10.1.12 *Lead Content*—Test Method [D3237](#) and [D5059](#).

11. Keywords

11.1 Avgas; aviation gasoline; gasoline; unleaded Avgas; unleaded aviation gasoline

APPENDIX

(Nonmandatory Information)

X1. PERFORMANCE CHARACTERISTICS OF UNLEADED AVIATION GASOLINE

X1.1 Introduction

X1.1.1 Unleaded aviation gasoline is a complex mixture of relatively volatile hydrocarbons that vary widely in their physical and chemical properties. The engines and aircraft impose a variety of mechanical, physical, and chemical environments. The properties of unleaded aviation gasoline ([Table X1.1](#)) shall be properly balanced to give satisfactory engine

performance over an extremely wide range of conditions.

X1.1.2 The ASTM requirements summarized in [Table 1](#) are quality limits established on the basis of the broad experience and close cooperation of producers of unleaded aviation gasoline, manufacturers of aircraft engines, and users of both commodities. The values given are intended to define unleaded aviation gasoline suitable for most types of spark-ignition aviation engines; however, certain equipment or conditions of use may require fuels having other characteristics.

X1.1.3 Specifications covering antiknock quality defines the grade of unleaded aviation gasoline. The other requirements either prescribe the proper balance of properties to ensure satisfactory engine performance or limit components of undesirable nature to concentrations so low that they will not have an adverse effect on engine performance.

X1.2 Combustion Characteristics (Antiknock Quality)

X1.2.1 The fuel-air mixture in the cylinder of a spark-ignition engine will, under certain conditions, ignite spontaneously in localized areas instead of progressing from the spark. This may cause a detonation or knock, usually inaudible in aircraft engines. This knock, if permitted to continue for more than brief periods, may result in serious loss of power and damage to, or destruction of, the aircraft engine. When unleaded aviation gasoline is used in other types of aviation engines, for example, in certain turbine engines where specifically permitted by the engine manufacturers, knock or detonation characteristics may not be critical requirements.

X1.2.2 The MON rating is determined in standardized laboratory knock test engines that are operated under prescribed conditions. Results are expressed as octane numbers up to 100. Octane number is defined arbitrarily as the percentage

TABLE X1.1 Performance Characteristics of Unleaded Aviation Gasoline

Performance Characteristics	Test Methods	Sections
Combustion characteristics	knock value (MON)	X1.2
Antiknock quality	isopropyl alcohol	X1.2.3
Fuel metering and aircraft range	density	X1.3.1
Combustion characteristics	net heat of combustion	X1.3.2
Carburetion and fuel vaporization	vapor pressure	X1.4.1
	distillation	X1.4.2
Corrosion of fuel system and engine parts	copper strip corrosion	X1.5.1
	sulfur content	X1.5.2
Fluidity at low temperatures	freezing point	X1.6
Fuel cleanliness, handling, and storage stability	potential gum	X1.7.1
	water reaction	X1.7.3
Miscellaneous	lead content	X1.8.2

of isooctane in that blend of isooctane and n-heptane that the gasoline matches in knock characteristics when compared by the procedure specified. The MON of the gasoline can be used as a guide to the amount of knock-limited power that may be obtained in a full-scale engine under take-off, climb and cruise conditions.

X1.2.3 Since isopropyl alcohol is normally added in the field at the point of use, the operator is cautioned that it may impact octane performance. Depending on octane grade, the addition of the IPA additive may increase or decrease the motor octane rating.

X1.2.4 *Knock Value, MON (Test Method D2700)*—The specification parameter knock value, lists “Motor Octane Number” (MON) as determined by Test Method **D2700**. Historically, aviation lean ratings were determined (from 1941 through 1970) by Test Method D614. An extensive comparison of National Exchange Group data from 1947 through 1964 established that motor octane numbers as determined by Test Methods D357 and D1948 could be converted to equivalent Test Method D614 ratings. A table to convert MON to the corresponding aviation lean rating was included in Test Method **D2700**, which was first issued in 1968 as a revision, consolidation and intended eventual replacement of Test Methods D357 (Withdrawn 1969), D614 (Withdrawn 1970), and D1948 (Withdrawn 1968). Currently unleaded aviation gasoline ratings are determined as MON, Test Method **D2700**.

X1.3 Fuel Metering and Aircraft Range

X1.3.1 *Density*—Density is a property of a fluid and is of significance in metering flow and in mass-volume relationships for most commercial transactions. It is particularly useful in empirical assessments of heating value when used with other parameters such as aniline point or distillation.

X1.3.2 *Net Heat of Combustion*—The net heat of combustion provides a knowledge of the amount of energy obtainable from a given fuel for the performance of useful work, in this instance, power. Aircraft design and operation are dependent upon the availability of a certain predetermined minimum amount of energy as heat. Consequently, a reduction in heat energy below this minimum is accompanied by an increase in fuel consumption with corresponding loss of range. Therefore, a minimum net heat of combustion requirement is incorporated in the specification. The determination of net heat of combustion is time consuming and difficult to conduct accurately. This led to the development and use of the aniline point and density relationship to estimate the heat of combustion of the fuel. This relationship is used along with the sulfur content of the fuel to obtain the net heat of combustion for the purposes of this specification. An alternative calculation, Test Method **D3338**, is based on correlations of aromatics content, density, volatility, and sulfur content. This test method may be preferred at refineries where all these values are normally obtained and the necessity to obtain the aniline point is avoided. The direct measurement method, Test Method **D4809**, is normally used only as a referee method in cases of dispute.

X1.3.3 No great variation in density or heat of combustion occurs in modern unleaded aviation gasolines, since they

depend on hydrocarbon composition that is already closely controlled by other specification properties.

X1.4 Carburetion and Fuel Vaporization

X1.4.1 In many spark-ignition aviation engines, the gasoline is metered in liquid form through the carburetor where it is mixed with air and vaporized before entering the supercharger from which the fuel-air mixture enters the cylinder of the engine. In other types of engines, the fuel may be metered directly into the supercharger, the cylinder, or the combustor. The volatility, the tendency to evaporate or change from a liquid to a gaseous state, is an extremely important characteristic of aviation fuel.

X1.4.2 Gasolines that vaporize too readily may boil in fuel lines or carburetors, particularly as altitude increases, and cause vapor lock with resultant stoppage of fuel flow to the engine. Conversely, fuels that do not completely vaporize may cause engine malfunctioning of other sorts. Therefore, a proper balance of the volatility of the various hydrocarbon components is essential to satisfactory performance of the finished fuel.

X1.4.3 *Vapor Pressure*—The vapor pressure of an unleaded aviation gasoline is the measure of the tendency of the more volatile components to evaporate. Experience has shown that fuels having a Reid vapor pressure no higher than 49 kPa will be free of vapor-locking tendencies under most conditions of aircraft usage. A research report is available.⁶

X1.4.4 *Distillation*—The relative proportions of all the hydrocarbon components of a gasoline are measured in terms of volatility by the range of distillation temperatures. The method is empirical and useful in comparing fuels, but is not intended to separate or identify quantitatively the individual hydrocarbons present in the fuel.

X1.4.4.1 A maximum value is set on the 10 % evaporated point to ensure ease of starting and a reasonable degree of flexibility during the warm-up period. To guard against too high a volatility that might lead to carburetor icing or vapor lock, or both, (also protected against by the vapor pressure test) a minimum value is set for the sum of the 10 and 50 % evaporated points.

X1.4.4.2 A maximum value is specified for the 50 % evaporated temperature to ensure average volatility sufficient to permit adequate evaporation of the fuel in the engine induction system. Insufficient evaporation may lead to loss of power.

X1.4.4.3 A maximum temperature is prescribed for the 90 % evaporated point to prevent too much liquid fuel being delivered to the cylinders, resulting in power loss, and to prevent poor distribution to the various cylinders. Such a condition might lead to excessive leanness in some cylinders with consequent engine roughness, perhaps accompanied by knocking and damage to the engine. Lowered fuel economy and excessive dilution of the lubricating oil may result from too high a 90 % evaporated point.

⁶ Supporting data have been filed at ASTM International Headquarters and may be obtained by requesting Research Report RR:D02-1146.

X1.4.4.4 A minimum value is stipulated for the 40 % evaporated temperature in an effort to control, indirectly, specific gravity and, consequently, carburetor metering characteristics.

X1.4.4.5 A maximum is placed on the final boiling point (end point) which, together with the maximum prescribed for the 90 % evaporated point, is used to prevent incorporation of excessively high boiling components in the fuel that may lead to maldistribution, spark plug fouling, power loss, lowered fuel economy, and lubricating oil dilution.

X1.4.4.6 The stipulation of a minimum recovery and a maximum loss in this specification in conjunction with the vapor pressure requirement is intended to protect against excessive losses by evaporation in storage, handling, and in the aircraft tank. It is also a check on the distillation test technique.

X1.4.4.7 A maximum value is specified for the distillation residue to prevent the inclusion of undesirable high-boiling components essentially impossible to burn in the combustion chamber, the presence of which may reflect the degree of care with which the product is refined or handled. The amount of residue along with the end point temperature can be used as an indication of contamination with high-boiling materials.

X1.5 Corrosion of Fuel System and Engine Parts

X1.5.1 *Copper Strip*—The requirement that gasoline must pass the copper strip corrosion test provides assurance that the product will not corrode the metal parts of fuel systems.

X1.5.2 *Sulfur*—Total sulfur content of aviation fuels is significant because the products of combustion of sulfur can cause corrosive wear of engine parts.

X1.6 Fluidity at Low Temperatures

X1.6.1 A freezing point requirement is specified to preclude solidification of any hydrocarbon components at extremely low temperatures with consequent interference with fuel flow to the engine.

X1.6.2 *Fuel System Icing Inhibitor*—Isopropyl alcohol (IPA), approved in 5.2.2.1, and diethyleneglycol monomethyl ether (Di-EGME), approved in 5.2.2.2, shall be in accordance with the requirements shown in Specification D4171.

X1.7 Fuel Cleanliness, Handling and Storage Stability

X1.7.1 *Potential Gum*—Fuel must be usable after storage for variable periods under a variety of climatic conditions. The potential gum test, which is an accelerated oxidation method, is used to estimate fuel stability in storage and the effectiveness of oxidation inhibitors. If the fuel is to be stored under relatively mild conditions for short periods, an oxidation period of 5 h is generally considered sufficient to indicate if the desired stability has been obtained, whereas a 16 h period is desirable to provide stability assurance for long periods and severe conditions, such as storage in tropical climates.

X1.7.2 *Permissible Oxidation Inhibitors and Oxidation Inhibitor Content*—Antioxidants are used to prevent the formation of gum in fuel during storage. The efficacy of a given inhibitor determined by the apparent oxidation stability of a fuel does not completely establish its suitability for use in an

aircraft engine. Oxidation inhibitors have been found to contribute to excessive induction system deposits; therefore, their acceptability for use must ultimately be determined in the full-scale aircraft engine.

X1.7.2.1 The chemical names of approved inhibitors and the maximum quantities permitted are shown in this specification.

X1.7.3 *Water Reaction*—The water reaction method provides a means of determining the presence of materials readily extractable by water or having a tendency to absorb water. When the fuel consists essentially of hydrocarbon components, there is no measurable change in the volume of the water layer.

X1.7.4 *Electrical Conductivity*—The generation of static electricity can create problems in the handling of unleaded aviation gasolines. Addition of a conductivity improver may be used as an additional precaution to reduce the amount of static electrical charge present during fuel handling. See Guide D4865 for more information.

X1.7.5 *Microbial Contamination*—Uncontrolled microbial contamination in fuel systems may cause or contribute to a variety of problems including corrosion, odor, filter plugging, decreased stability, and deterioration of fuel/water separation characteristics. In addition to system component damage, off-specification fuel can result.

X1.7.6 Guide D6469 provides personnel with limited microbiological background an understanding of the symptoms, occurrence, and consequences of chronic microbial contamination. The guide also suggests means for detection and control. No biocides are approved for unleaded aviation gasoline, therefore engine and airframe manufacturers' guidelines must be followed if they are to be used.

X1.8 Miscellaneous Tests

X1.8.1 *Aromatics Content*—Low boiling aromatics, which are common constituents of unleaded aviation gasolines, are known to affect elastomers to a greater extent than other components in unleaded aviation gasoline. Although Specification D7547 does not include an explicit maximum aromatic limit, other specification limits effectively restrict the aromatic content of unleaded aviation gasolines. Benzene is virtually excluded by the maximum freezing point of -58°C , while other aromatics are limited by the minimum heating value and the maximum distillation end point. Thus, the heating value limits toluene to about 24 %. Xylenes have a slightly higher heating value and, therefore, would permit somewhat higher aromatic concentrations; however, their boiling points (above 138°C) limit their inclusion at levels not higher than 10 %. Total aromatic levels above 25 % in unleaded aviation gasoline are, therefore, extremely unlikely.

X1.8.2 *Lead Content*—A number of analytical test methods are permitted to cover the unintentional presence of lead in unleaded fuel. The intentional addition of lead compounds to unleaded aviation gasoline is not permitted. Industry practice currently limits maximum concentrations of incidental lead to 0.013 g of lead/L (0.05 g/U.S. gal).

X1.9 General

X1.9.1 Further detailed information on the significance of all test methods relevant to unleaded aviation gasoline is provided in Manual MNL 1.⁷

⁷ MNL1, *Significance of Tests for Petroleum Products*, Seventh Edition, Rand, S. J., editor, ASTM International, 2003.

SUMMARY OF CHANGES

Subcommittee D02.J0 has identified the location of selected changes to this standard since the last issue (D7547 – 15^{e1}) that may impact the use of this standard. (Approved May 1, 2017.)

(1) Added new Research Report number in subsection 4.1.

ASTM International takes no position respecting the validity of any patent rights asserted in connection with any item mentioned in this standard. Users of this standard are expressly advised that determination of the validity of any such patent rights, and the risk of infringement of such rights, are entirely their own responsibility.

This standard is subject to revision at any time by the responsible technical committee and must be reviewed every five years and if not revised, either reapproved or withdrawn. Your comments are invited either for revision of this standard or for additional standards and should be addressed to ASTM International Headquarters. Your comments will receive careful consideration at a meeting of the responsible technical committee, which you may attend. If you feel that your comments have not received a fair hearing you should make your views known to the ASTM Committee on Standards, at the address shown below.

This standard is copyrighted by ASTM International, 100 Barr Harbor Drive, PO Box C700, West Conshohocken, PA 19428-2959, United States. Individual reprints (single or multiple copies) of this standard may be obtained by contacting ASTM at the above address or at 610-832-9585 (phone), 610-832-9555 (fax), or service@astm.org (e-mail); or through the ASTM website (www.astm.org). Permission rights to photocopy the standard may also be secured from the Copyright Clearance Center, 222 Rosewood Drive, Danvers, MA 01923, Tel: (978) 646-2600; <http://www.copyright.com/>