



Standard Specification for Pyrolysis Liquid Biofuel¹

This standard is issued under the fixed designation D7544; 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 grades of pyrolysis liquid biofuel produced from biomass intended for use in various types of fuel-burning equipment under various climatic and operating conditions. These grades are described as follows:

1.1.1 Grade G is intended for use in industrial burners equipped to handle the pyrolysis liquid biofuels meeting the requirements listed for Grade G in [Table 1](#). The pyrolysis liquid biofuel listed under Grade G in [Table 1](#) is not intended for use in residential heaters, small commercial boilers, engines, or marine applications.

1.1.2 Grade D is intended for use in commercial/industrial burners requiring lower solids and ash content and which are equipped to handle the pyrolysis liquid biofuels meeting the requirements listed for Grade D in [Table 1](#). The pyrolysis liquid biofuel listed under Grade D in [Table 1](#) is not intended for use in residential heaters, engines, or marine applications not modified to handle these types of fuels.

NOTE 1—For information on the significance of the physical, chemical, and performance properties identified in this specification, see [Appendix X1](#).

1.2 This specification is for use in contracts for the purchase of pyrolysis liquid biofuel and for guidance of consumers of this type of fuel.

1.3 Nothing in this specification should preclude observance of national or local regulations, which may be more restrictive.

NOTE 2—The generation and dissipation of static electricity may create problems in the handling of pyrolysis liquid biofuel. For more information on the subject, see [Guide D4865](#).

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

1.4.1 *Exception*—BTU units are included for information only in [3.1.5.1](#).

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.

2. Referenced Documents

2.1 *ASTM Standards*:²

[D93 Test Methods for Flash Point by Pensky-Martens Closed Cup Tester](#)

[D97 Test Method for Pour Point of Petroleum Products](#)

[D240 Test Method for Heat of Combustion of Liquid Hydrocarbon Fuels by Bomb Calorimeter](#)

[D396 Specification for Fuel Oils](#)

[D445 Test Method for Kinematic Viscosity of Transparent and Opaque Liquids \(and Calculation of Dynamic Viscosity\)](#)

[D482 Test Method for Ash from Petroleum Products](#)

[D4052 Test Method for Density, Relative Density, and API Gravity of Liquids by Digital Density Meter](#)

[D4294 Test Method for Sulfur in Petroleum and Petroleum Products by Energy Dispersive X-ray Fluorescence Spectrometry](#)

[D4865 Guide for Generation and Dissipation of Static Electricity in Petroleum Fuel Systems](#)

[D5854 Practice for Mixing and Handling of Liquid Samples of Petroleum and Petroleum Products](#)

[D6469 Guide for Microbial Contamination in Fuels and Fuel Systems](#)

[D7579 Test Method for Pyrolysis Solids Content in Pyrolysis Liquids by Filtration of Solids in Methanol](#)

[E70 Test Method for pH of Aqueous Solutions With the Glass Electrode](#)

[E203 Test Method for Water Using Volumetric Karl Fischer Titration](#)

3. Terminology

3.1 *Definitions of Terms Specific to This Standard:*

3.1.1 *bulk fuel, n*—fuel in the storage facility in quantities over 190 L.

¹ This specification is under the jurisdiction of ASTM Committee D02 on Petroleum Products and Lubricants and is the direct responsibility of Subcommittee D02.E0 on Burner, Diesel, Non-Aviation Gas Turbine, and Marine Fuels.

Current edition approved June 1, 2012. Published September 2012. Originally approved in 2009. Last previous edition approved in 2010 as D7544-10. DOI:10.1520/D7544-12.

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

TABLE 1 Detailed Requirements for Pyrolysis Liquid Biofuels

| Property | Test Method | Grade G | Grade D |
|--|-------------------|---------|---------|
| Gross Heat of Combustion, MJ/kg, min | D240 | 15 | 15 |
| Water Content, % mass, max | E203 | 30 | 30 |
| Pyrolysis Solids Content, % mass, max | D7579 | 2.5 | 0.25 |
| Kinematic Viscosity at 40°C, mm ² /s, max | D445 ^A | 125 | 125 |
| Density at 20°C, kg/dm ³ | D4052 | 1.1–1.3 | 1.1–1.3 |
| Sulfur Content, % mass, max | D4294 | 0.05 | 0.05 |
| Ash Content, % mass, max | D482 | 0.25 | 0.15 |
| pH. | E70 | Report | Report |
| Flash Point, °C, min | D93, Procedure B | 45 | 45 |
| Pour Point, °C, max | D97 | –9 | –9 |

^A Without filtering.

3.1.2 *char, n*—fine carbonaceous powder that is separated from the vapors of biomass during pyrolysis.

3.1.2.1 *Discussion*—Pyrolysis liquid biofuel contains uniformly suspended char.

3.1.3 *commercial burner, n*—device which produces heat for commercial use through the combustion of liquid fuels.

3.1.3.1 *Discussion*—Commercial burners are typically designed for processes that provide direct heating. Commercial boilers or heaters—small to medium indirect heating units which transfer thermal energy to water or other fluids or gases for use in heating in commercial settings, power generation and in manufacturing processes. These boilers can be classified as small or medium commercial boilers with a heat input of between 10.5 to 105 GJ/h (10 to 100 × 106 BTU/h).

3.1.4 *fuel degradation products, n*—those materials that are formed in fuel during extended storage or exposure to high temperatures.

3.1.4.1 *Discussion*—During storage, reactive organic compounds in pyrolysis liquid can act together to form larger molecules (fuel degradation products), which can become insoluble or increase the fuel viscosity, or both.

3.1.5 *industrial burner, n*—device which produces heat for industrial use through the combustion of liquid fuels.

3.1.5.1 *Discussion*—Industrial burners are typically designed for one of two applications: (1) industrial furnaces—integral components of manufacturing processes that provide direct heating; for example, in aggregate, cement, lime, or phosphate kilns; coke ovens; or blast, smelting, melting, refining, or drying ovens and (2) industrial boilers—large indirect heating units which transfer thermal energy to water or other fluids or gases for use in heating in industrial settings, power generation and in manufacturing processes. These boilers can be classified as utility/large industrial boilers with a heat input greater than 105 GJ/h (100 × 106 BTU/h) or small industrial boilers with a heat input of between 10.5 to 105 GJ/h (10 to 100 × 106 BTU/h).

3.1.6 *long-term storage*—storage of fuel for longer than 3 months after it is received by the user.

3.1.7 *medium-term storage*—storage of fuel for up to 3 months after it is received by the user.

3.1.7.1 *Discussion*—It is recommended that fuel be consumed within 6 months of receipt.

3.1.8 *pyrolysis, n*—chemical decomposition of organic materials by heating in the absence of oxygen.

3.1.9 *pyrolysis liquid biofuel, n*—liquid product from the pyrolysis of biomass.

3.1.9.1 *Discussion*—Pyrolysis liquid biofuel is comprised of a complex mixture of the decomposition products of renewable resources such as ligno-cellulosic biomass including highly oxygenated organic compounds. It is produced from the pyrolysis of biomass, followed by the rapid condensation of its vapors.

3.1.10 *pyrolysis solids, n*—solid particles contained within the pyrolysis liquid biofuel.

3.1.10.1 *Discussion*—Pyrolysis solids is comprised of ash and char.

4. General Requirements

4.1 The pyrolysis liquid biofuel specified in this specification shall remain uniform in medium-term storage and not separate by gravity into layers.

NOTE 3—Long-term storage or equipment down time can necessitate circulation of pyrolysis liquid biofuel in-tank to prevent such separation. The buyer and seller should agree on any requirements for long-term storage. If minor separation occurs during medium-term storage, mild agitation or product circulation should reverse such separation.

5. Detailed Requirements

5.1 The various grades of pyrolysis liquid biofuel shall conform to the detailed requirements shown in **Table 1**.

5.2 The properties selected for limitation are those that are believed to be of the greatest significance in obtaining acceptable performance of the burner.

6. Sampling

6.1 Review all intended test methods prior to sampling to understand the importance and effects of sampling technique, proper containers, and special handling required for each test method. See **Table 2**.

6.2 As indicated in **4.1**, during medium-term storage, pyrolysis liquid biofuel shall remain uniform and not separate into layers. Note, however, that separation may occur during long-term storage. Therefore, samples should be well mixed when transferring from the primary sampling process or container, or both, to another container or analytical apparatus, or both. Sampling from an active circulation loop or a well mixed or agitated tank is preferred. Refer to Practice **D5854** for more guidance on mixing and handling samples.

TABLE 2 Typical Sampling Procedures for Containers

| Type of Container | Procedure |
|---|---------------------------------------|
| Storage tanks that are well-mixed by circulation or agitation | Automatic or Manual Pipeline Sampling |
| Tank cars, tank trucks, or ship tanks | Dipper Sampling or All-Level Sampling |

6.3 *Sample Size*—A minimum of 1 L is recommended.

7. Test Methods

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

- 7.1.1 *Gross Heat of Combustion*—Test Method **D240**.
- 7.1.2 *Water Content*—Test Method **E203**.
- 7.1.3 *Pyrolysis Solids Content*—See Test Method **D7579**.
- 7.1.4 *Kinematic Viscosity*—Test Method **D445**.
- 7.1.5 *Density*—Test Method **D4052**.

- 7.1.6 *Sulfur*—Test Method **D4294**.
- 7.1.7 *Ash Content*—Test Method **D482**.
- 7.1.8 *pH*—Test Method **E70**.
- 7.1.9 *Flash Point*—Test Method **D93**.
- 7.1.10 *Pour Point*—Test Method **D97**.

8. Keywords

8.1 pyrolysis liquid biofuel; bio-oil; pyrolysis oil; pyoil; py-oil; bio-crude-oil; bio-fuel-oil; wood liquids; burner fuel; renewable energy; alternative energy; fuel oils; furnace oils

APPENDIXES

(Nonmandatory Information)

X1. SIGNIFICANCE OF ASTM SPECIFIED PROPERTIES FOR PYROLYSIS LIQUID BIOFUEL

X1.1 Heat of Combustion

X1.1.1 Heat of combustion is a measure of the energy content of pyrolysis liquid. As a reference, pyrolysis liquid has approximately half the heat of combustion on a volumetric basis compared to #2, **D396** fuel oil. The heat of combustion is an important basis for quantifying its monetary value and for equipment selection or design, or both.

X1.2 Water Content

X1.2.1 Higher water content leads to lower overall viscosity and heat of combustion of pyrolysis liquid.

X1.2.2 Water content reduces the flame temperature of pyrolysis liquid, which contributes to lower nitrogen dioxide emissions during combustion.

X1.2.3 Water content higher than the limit, can cause phase separation, leading to a non-homogenous mixture.

X1.3 Char Content

X1.3.1 Pyrolysis liquid biofuel contains uniformly suspended char. Pyrolysis liquid can be produced to various char concentrations. Increasing char content may also increase the ash content, viscosity and pour point of pyrolysis liquid biofuel and affect the pyrolysis liquid biofuel handling, atomization and storage stability in a negative manner.

X1.4 Viscosity

X1.4.1 The viscosity of a pyrolysis liquid is a measure of its resistance to flow. It is of major importance so that adequate preheating facilities can be provided to permit the pyrolysis liquid to be pumped to the burner and to provide good atomization. However, it is equally important that the maximum viscosity under the existing conditions be such that the pyrolysis liquid can be pumped satisfactorily from the storage tank to the preheater.

X1.5 Density

X1.5.1 Density alone is of little significance as an indication of the burning characteristics of pyrolysis liquid. However, when used in conjunction with other properties, it is of value in mass-volume relationships.

X1.6 Sulfur Content

X1.6.1 Knowledge of the sulfur content is important for evaluating the Sulfur Oxides emissions from combusting pyrolysis liquid.

X1.7 Ash Content

X1.7.1 Ash is the amount of noncombustible material in pyrolysis liquid biofuel. Ash-forming materials may be present in pyrolysis liquid biofuel in two forms, solid particles or water soluble metallic compounds, or both. Depending on the size and type (sand, char, etc.) of the solid ash particles, these particles can contribute to wear of burner pumps and valves, and decrease fuel efficiency by fouling heat exchange surfaces. The soluble metallic compounds have little or no effect on wear or plugging, but they can contain elements that produce corrosion and deposits on boiler heating surfaces. Excessive amounts of ash also may necessitate particulate collection equipment for compliance with national or local air emission regulations.

X1.8 pH

X1.8.1 Due to its low pH, materials which are in contact with pyrolysis liquid shall be selected for suitability and corrosion resistance.

X1.9 Flash Point

X1.9.1 The flash point of a fuel is an indication of the maximum temperature at which it can be stored and handled without serious fire hazard.

X1.10 Pour Point

X1.10.1 The pour point of pyrolysis liquid is an indication of the lowest temperature at which the pyrolysis liquid is capable of flowing under very low forces. The pour point is

prescribed in accordance with the conditions of storage and use. Higher pour point fuels are permissible where heated storage and adequate piping facilities are provided.

X2. LONG-TERM STORAGE OF PYROLYSIS LIQUID

X2.1 Scope

X2.1.1 This appendix provides guidance for consumers of pyrolysis liquid who may wish to store quantities of the fuel for extended periods.

X2.1.2 Pyrolysis liquid has adequate stability properties to withstand normal storage and use without formation of troublesome amounts of insoluble degradation products.

X2.1.3 The following suggested practices are general in nature and should not be considered substitutes for any requirements imposed by the warranty of the fuel equipment manufacturer or by federal, state or local government regulations. Further guidance in developing a fuel management system could be provided by fuel suppliers.

X2.2 Terminology (See Section 3)

X2.3 Fuel Selection

X2.3.1 The stability properties of pyrolysis liquid are not fully understood and appear to depend on the biomass feedstock sources, the pyrolysis reaction conditions, the pyrolysis solids content of the pyrolysis liquid, whether additives are present and storage conditions.

X2.3.2 The composition and stability properties of pyrolysis liquid produced at specific production plants can be different. A special requirement of the user, such as long-term storage, should be discussed with the supplier.

X2.4 Fuel Storage Conditions

X2.4.1 Store at a minimum temperature of 10°C to maintain adequate fluidity, but not higher than 30°C for long-term storage. High storage temperatures accelerate fuel degradation. Recommended storage temperature is 15 to 20°C.

X2.4.2 Store in containers built of corrosion resistant steel and materials such as AISI 304, AISI 316, PTFE (polytetrafluoroethylene), PP (polypropylene), HDPE (high density polyethylene) and PVC (polyvinylchloride).

X2.4.3 Long-term storage in an agitated or circulated container is recommended to maintain homogeneity.

X2.4.4 Limit exposure to air to prevent oxidation.

X2.5 Fuel Additives

X2.5.1 Fuel additives are not required in pyrolysis liquid biofuel.

X2.5.2 Polar solvent additives, in particular methanol and ethanol, can be used to improve the long-term storage of pyrolysis liquid. For maximum benefits, additives should be added directly following production. The addition of a polar solvent to the pyrolysis liquid can significantly lower the flash point and appropriate precautions shall be taken.

X2.5.3 Hydrocarbon fuel is not miscible in pyrolysis liquid biofuel.

X2.6 Tests for Fuel Quality

X2.6.1 Work has started on a stability index that correlates the long-term storage stability of pyrolysis liquid against accelerated aging tests at increased temperatures.

X2.7 Fuel Monitoring

X2.7.1 Stored fuel should be periodically sampled and its quality assessed. Measurement and comparison of the stored fuel's viscosity by Test Method **D445** can be used to assess fuel degradation.

X2.8 Use of Degraded Fuels

X2.8.1 Fuels that have undergone mild-to-moderate degradation can often be consumed in a normal way, depending on the fuel system requirements. Filters and other cleanup equipment may require special attention and increased maintenance. Burner nozzle or injector fouling may occur more rapidly.

X2.8.2 If for example, the fuel is too thick for a particular application, mixing in a polar solvent to the pyrolysis liquid prior to use is known to reduce its viscosity and with vigorous mixing will dissolve some insolubles back into the fuel. The addition of a polar solvent to the pyrolysis liquid can significantly lower the flash point and appropriate precautions shall be taken.

X3. HAZARDS

X3.1 The biofuels defined by this specification may have a lower pH and gross heat of combustion than **D396** specification fuels. The materials of construction and equipment sizing of the industrial burner and its fuel handling system shall be compatible with the properties of the biofuels defined by this specification.

X4. MICROBIAL CONTAMINATION

X4.1 Uncontrolled microbial contamination in fuel systems can cause or contribute to a variety of problems including increased corrosivity, and decreased stability, filterability, and caloric value. Microbial processes in fuel systems can also cause or contribute to system damage.

X4.2 Because the microbes contributing to the aforementioned problems may not be necessarily present in the fuel itself, no microbial quality criterion for fuels is recommended.

However, it is important that personnel responsible for fuel quality understand how uncontrolled microbial contamination can affect fuel quality.

X4.3 Guide **D6469** provides personnel with limited microbiological background an understanding of the symptoms, occurrences, and consequences of chronic microbial contamination. Guide **D6469** also suggests means for detecting and controlling microbial contamination in fuels and fuel systems.

SUMMARY OF CHANGES

Committee D02 has identified the location of selected changes to this standard since the last issue (D7544–10) that may impact the use of this standard.

(1) Added **1.1.1**, **1.1.2**, and **3.1.3**.

(2) Revised **3.1.8**, **5.1**, and **Table 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 ASTM website (www.astm.org/COPYRIGHT).