

Designation: D565 - 99 (Reapproved 2013)

Standard Test Method for Carbonizable Substances in White Mineral Oil¹

This standard is issued under the fixed designation D565; the number immediately following the designation indicates the year of original adoption or, in the case of revision, the year of last revision. A number in parentheses indicates the year of last reapproval. A superscript epsilon (ε) indicates an editorial change since the last revision or reapproval.

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

1. Scope

- 1.1 This test method covers white mineral oil (Mineral Oil USP and Light Mineral Oil NF) to determine whether it conforms to the standard of quality required for pharmaceutical use as defined by the United States Pharmacopeia and the National Formulary, or the Food and Drug Administration.
- 1.2 The values stated in SI units are to be regarded as standard. No other units of measurement are included in this standard.
- 1.2.1 *Exception*—Dimension requirements for the color comparator in Fig. 1 are in SI and inch-pound units.
- 1.3 This standard does not purport to address all of the safety concerns, if any, associated with its use. It is the responsibility of the user of this standard to establish appropriate safety and health practices and determine the applicability of regulatory limitations prior to use. For specific hazard statements, see Section 6.

2. Referenced Documents

2.1 ASTM Standards:²

D1193 Specification for Reagent Water

2.2 Official Compendia:³

United States Pharmacopeia—Current Edition

Monograph on Mineral Oil

National Formulary—Current Edition

Monograph on Light Mineral Oil

2.3 Government Document:⁴ 21CFR 172.878 Food and Drug Administration Title

3. Summary of Test Method

3.1 The mineral oil is treated with concentrated sulfuric acid (H_2SO_4) under prescribed conditions and the resulting color is compared with a reference standard to determine whether it passes or fails the test.

4. Significance and Use

4.1 This test method is a means for ascertaining whether pharmaceutical mineral oil conforms to the standards of the United States Pharmacopeia, the National Formulary, and the Food and Drug Administration.

5. Apparatus

- 5.1~Test~Tube, as shown in Fig. 1, of heat-resistant glass fitted with a well-ground glass stopper, the stopper and the tube bearing identical and indestructible numbers. The tube shall be $140~\pm~2~$ mm in length and between 14.5~and 15.0~mm in outside diameter, and shall be calibrated at the $5~\pm~0.2~$ mL and $10~\pm~0.2~$ mL liquid levels. The capacity of the tube with stopper inserted shall be between 13.6~and 15.6~mL. A rolled edge can be provided for suspending the tube on the cover of the water bath
- 5.2 Water Bath, suitable for immersing the test tube above the 10 mL line equipped to maintain a temperature of 100 ± 0.5 °C. The bath shall be provided with a cover of any suitable material with holes approximately 16 mm in diameter through which the test tubes can be suspended.
- 5.3 *Color Comparator*, of a suitable type for observing the color of the acid layer in comparison with the reference standard color solution. The size and shape of the comparator are optional, but the size and shape of the apertures shall conform to the dimensions prescribed in Fig. 1.

¹ This test method is under the jurisdiction of ASTM Committee D02 on Petroleum Products, Liquid Fuels, and Lubricantsand is the direct responsibility of Subcommittee D02.06 on Analysis of Lubricants.

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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.

³ Available from U.S. Pharmacopeial Convention, 12601 Twinbrook Parkway, Rockville, MD 20852.

⁴ Available from Standardization Documents Order Desk, DODSSP, Bldg. 4, Section D, 700 Robbins Ave., Philadelphia, PA 19111-5098.