



Designation: ~~D4481 – 10~~ **D4481 – 10 (Reapproved 2015)**

Standard Test Method for Total Nickel in Fresh Alumina-Base Catalysts¹

This standard is issued under the fixed designation D4481; 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 test method covers the determination of nickel in fresh alumina-base catalysts and has been tested at nickel concentrations from 2.5 to 60 weight %, expressed as nickel oxide (NiO).

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

1.3 *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 its use.*

2. Referenced Documents

2.1 *ASTM Standards:*²

[D1193 Specification for Reagent Water](#)

[D7442 Practice for Sample Preparation of Fluid Catalytic Cracking Catalysts and Zeolites for Elemental Analysis by Inductively Coupled Plasma Atomic Emission Spectroscopy](#)

[E105 Practice for Probability Sampling of Materials](#)

[E177 Practice for Use of the Terms Precision and Bias in ASTM Test Methods](#)

[E456 Terminology Relating to Quality and Statistics](#)

[E691 Practice for Conducting an Interlaboratory Study to Determine the Precision of a Test Method](#)

3. Summary of Test Method

3.1 The test specimen (as received) is treated with concentrated hydrochloric acid to solubilize the nickel. If necessary, nickel is recovered from any insoluble residue by potassium pyrosulfate fusion, after hydrofluoric-sulfuric acid treatment to remove silica. Ammonium citrate is added to complex the aluminum and buffer the solution. Nickel is precipitated as nickel dimethylglyoxime, Ni (C₄H₇O₂N₂)₂, at a weakly alkaline pH. The precipitate is washed and weighed as Ni (C₄H₇O₂N₂)₂ after drying at 120°C.

3.2 A separate test specimen is taken to determine loss on ignition (LOI) at 550°C. The value is used to calculate the nickel as percent nickel oxide (NiO) on a 550°C dry basis.

4. Significance and Use

4.1 This test method sets forth a procedure by which catalyst samples can be compared either on an interlaboratory or intralaboratory basis. It is anticipated that catalyst producers and users will find this method of value.

5. Interferences

5.1 Cobalt, molybdenum, and aluminum do not interfere. Interferences by elements that precipitate as hydroxides, such as iron, chromium, aluminum, lead, tin, manganese, titanium, and zirconium, are avoided by the addition of ammonium citrate before making the solutions ammoniacal. Copper, present in the 2 to 10 % range, tends to be co-precipitated with the nickel dimethylglyoxime. The only other metal ions precipitated by dimethylglyoxime are palladium, gold, and bismuth.

6. Apparatus

6.1 *Beakers*, 600-mL, 150-mL.

6.2 *Hotplate*.

¹ This test method is under the jurisdiction of Committee [D32](#) on Catalysts and is the direct responsibility of Subcommittee [D32.03](#) on Chemical Composition. Current edition approved April 1, 2010/Dec. 1, 2015. Published May 2010/December 2015. Originally approved in 1985. Last previous edition approved in 2004/2010 as D4481-04-10. DOI: [10.1520/D4481-10](#); [10.1520/D4481-10R15](#).

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.

- 6.3 *Furnace*, electric muffle. Calibrated and capable of maintaining temperatures of $550 \pm 25^{\circ}\text{C}$, and $950 \pm 25^{\circ}\text{C}$.
- 6.4 *Platinum Dishes*, 100-mL³ capacity.
- 6.5 *Mortar and Pestle*, agate, or equivalent mechanical grinder.
- 6.6 *Crucibles*, sintered-glass, 30-mL, medium porosity frit.
- 6.7 *Fiberglass Filter*, 3.2 cm.
- 6.8 *Drying Oven* capable of maintaining a temperature of 120°C .
- 6.9 *Vacuum Filtering Flask*, 500-mL.
- 6.10 *Filter Holder and Filter Disk*, Millipore 0.65 μm -47-mm diameter.
- 6.11 *pH Paper* to detect a value of 9.
- 6.12 *Screen*, 250- μm openings, 60-mesh.
- 6.13 *Analytical Balance*, capable of weighing to nearest 0.1 mg.
- 6.14 *Ashless Filter Pulp*.
- 6.15 *Desiccator*.

7. Reagents

7.1 *Purity of Reagents*—Reagent grade chemicals shall be used in all tests. Unless otherwise indicated, it is intended that all reagents shall conform to the specifications of the Committee on Analytical Reagents of the American Chemical Society, where such specifications are available.³ Other grades may be used, provided it is first ascertained that the reagent is of sufficiently high purity to permit its use without lessening the accuracy of the determination.

7.2 *Purity of Water*—Unless otherwise indicated, references to water shall be understood to mean type IV reagent water as defined in Specification **D1193**.

7.3 *Ammonium Citrate*, dibasic crystal, $(\text{NH}_4)_2\text{HC}_6\text{H}_5\text{O}_7$.

7.4 *Ammonium Hydroxide*, 10 and 25 % solutions.

NOTE 1—Refer to Section 8 on Hazards in Practice **D7442** for hazards associated with handling of acids.

7.5 *Hydrochloric Acid*, concentrated 38 %.

7.6 *Hydrofluoric Acid*, concentrated 48 %.

7.7 *Methyl Red Indicator (o-carboxy benzene azodimethylaniline)*—Dissolve 0.02 g in 60-mL reagent grade ethanol. Add 40.0 mL of distilled water.

7.8 *4A Molecular Sieve*, activated for 2 h at 275°C .

7.9 *Potassium Hydrogen Sulfate*, fused power-acid-flux grade.

7.10 *Sodium Dimethylglyoxime*, 8-Hydrate, crystal, $\text{CH}_3\text{C}:\text{NONaC}:\text{NONaCH}_3 \cdot 8\text{H}_2\text{O}$. Prepare a 1 weight % per volume solution in water.

7.11 *Sulfuric Acid*, 10 and 50 % solutions.

NOTE 2—The sodium salt of dimethylglyoxime is recommended, rather than the organic reagent, dimethylglyoxime because the salt is water soluble. Large excesses added to precipitate the nickel do no harm as the excess dimethylglyoxime is eliminated during the filtration and washing part of the procedure. The organic reagent, dimethylglyoxime, is not very soluble in water and is often added as a 1 % solution in ethanol. When added in this way, some of the excess reagent may contaminate the precipitated nickel complex. For this reason an aqueous solution of the sodium salt is preferred. If the alcohol solution of dimethylglyoxime is to be used, add an amount so that the alcohol content of the sample solution does not exceed 50 % or some of the precipitated nickel dimethylglyoximate may dissolve.

8. Sampling

8.1 The selection of a representative analytical sample from the bulk material is outside the scope of the present test method. It is presumed that parties using this test method for comparison purposes will have agreed on the selection of an analytical sample. If a sampling procedure is desired, Practice **E105** is recommended.

8.2 Grind the sample to pass through a 60-mesh screen.

³ *Reagent Chemicals, American Chemical Society Specifications*, American Chemical Society, Washington, DC. For Suggestions on the testing of reagents not listed by the American Chemical Society, see *Annual Standards for Laboratory Chemicals*, BDH Ltd., Poole, Dorset, U.K., and the *United States Pharmacopeia and National Formulary*, U.S. Pharmacopeial Convention, Inc. (USPC), Rockville, MD.