



Designation: C1730 – 17 (Reapproved 2022)

Standard Test Method for Particle Size Distribution of Advanced Ceramics by X-Ray Monitoring of Gravity Sedimentation¹

This standard is issued under the fixed designation C1730; 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 particle size distribution of advanced ceramic powders. Experience has shown that this test method is satisfactory for the analysis of silicon carbide, silicon nitride, and zirconium oxide in the size range of 0.1 μm to 50 μm .

1.1.1 However, the relationship between size and sedimentation velocity used in this test method assumes that particles sediment within the laminar flow regime. It is generally accepted that particles sedimenting with a Reynolds number of 0.3 or less will do so under conditions of laminar flow with negligible error. Particle size distribution analysis for particles settling with a larger Reynolds number may be incorrect due to turbulent flow. Some materials covered by this test method may settle in water with a Reynolds number greater than 0.3 if large particles are present. The user of this test method should calculate the Reynolds number of the largest particle expected to be present in order to judge the quality of obtained results. Reynolds number (Re) can be calculated using the following equation:

$$Re = \frac{D^3(\rho - \rho_0)\rho_0 g}{18\eta^2} \quad (1)$$

where:

D = the diameter of the largest particle expected to be present, in cm,

ρ = the particle density, in g/cm^3 ,

ρ_0 = the suspending liquid density, in g/cm^3 ,

g = the acceleration due to gravity, 981 cm/sec^2 , and

η = the suspending liquid viscosity, in poise.

1.1.2 A table of the largest particles that can be analyzed with a suggested maximum Reynolds number of 0.3 or less in water at 35 °C is given for a number of materials in [Table 1](#). A column of the Reynolds number calculated for a 50- μm particle sedimenting in the same liquid system is also given for each material. Larger particles can be analyzed in dispersing media

¹ This test method is under the jurisdiction of ASTM Committee C28 on Advanced Ceramics and is the direct responsibility of Subcommittee C28.03 on Physical Properties and Non-Destructive Evaluation.

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with viscosities greater than that for water. Aqueous solutions of glycerine or sucrose have such higher viscosities.

1.2 The procedure described in this test method may be applied successfully to other ceramic powders in this general size range, provided that appropriate dispersion procedures are developed. It is the responsibility of the user to determine the applicability of this test method to other materials. Note however that some ceramics, such as boron carbide and boron nitride, may not absorb X-rays sufficiently to be characterized by this analysis method.

1.3 The values stated in cgs units are to be regarded as the standard, which is the long-standing industry practice. The values given in parentheses are for information only.

1.4 *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, health, and environmental practices and determine the applicability of regulatory limitations prior to use.* Specific hazard information is given in [Section 8](#).

1.5 *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

C1145 Terminology of Advanced Ceramics

E1617 Practice for Reporting Particle Size Characterization Data

3. Terminology

3.1 For definitions of terms used in this test method, refer to Terminology [C1145](#).

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.

TABLE 1 Maximum Diameter of Ceramic Powders That Can Be Analyzed with Reynolds Number of 0.3 or Less in Water at 35 °C

Particle Composition	Particle Density	Maximum Particle Diameter, μm	Reynolds Number for 50 μm^{A}
Aluminum Nitride	3.26	50.36	0.29
Aluminum Oxide	3.965	46.01	0.39
Cerium Dioxide	7.132	36.13	0.80
Silicon Carbide	3.217	50.68	0.29
Silicon Nitride	3.44	49.09	0.32
Yttrium Oxide	5.01	41.61	0.52
Zirconium Oxide	5.89	38.95	0.63

^A A Reynolds number calculated for 50- μm particles sedimenting in water at 35 °C, with a density of 0.9941 g/cm³ and viscosity of 0.7225 cp. Entries with Reynolds numbers that exceed the recommended upper limit of 0.30 are included to indicate that samples of these materials containing 50- μm particles should not be analyzed using water as a dispersing liquid without addition of a viscosity modifier such as glycerol or sucrose.

4. Summary of Test Method

4.1 A carefully dispersed, homogeneous suspension of the powder is permitted to settle in a cell scanned by a collimated X-ray beam of constant intensity. The net X-ray signal is inversely proportional to the sample concentration in the dispersing medium, and the particle diameter is related to the position of the X-ray beam relative to the top of the cell. Cumulative mass percent versus equivalent spherical diameter is recorded to yield a particle size distribution curve.

5. Significance and Use

5.1 This test method is useful to both suppliers and users of powders, as outlined in 1.1 and 1.2, in determining particle size distribution for product specifications, manufacturing control, development, and research.

5.2 Users should be aware that sample concentrations used in this test method may not be what is considered ideal by some authorities, and that the range of this test method extends into the region where Brownian movement could be a factor in conventional sedimentation. Within the range of this test method, neither the sample concentration nor Brownian movement is believed to be significant. Standard reference materials traceable to national standards, of chemical composition specifically covered by this test method, are available from NIST,³ and perhaps other suppliers.

5.3 Reported particle size measurement is a function of the actual particle dimension and shape factor as well as the particular physical or chemical properties being measured. Caution is required when comparing data from instruments operating on different physical or chemical parameters or with different particle size measurement ranges. Sample acquisition, handling, and preparation can also affect reported particle size results.

5.4 Suppliers and users of data obtained using this test method need to agree upon the suitability of these data to provide specification for and allow performance prediction of the materials analyzed.

6. Apparatus

6.1 *X-Ray Gravitational Sedimentation Particle Size Analyzer*—The analyzer shall utilize extinction of collimated

X-rays to determine particle mass concentration as a homogeneous dispersion of sample sediments under the influence of gravity.⁴

6.2 *Ultrasonic Probe*, consisting of a 200 to 300 W power unit, ultrasonic transducer, and 13-mm (1/2-in.) diameter probe.

6.3 *Balance*, top-loading, accurate to 100 \pm 0.1 g.

6.4 *Stirrer*, magnetic, with 19-mm (3/4-in.) stirrer bar.

7. Reagents and Materials

7.1 *Purity of Reagents*—Reagent-grade chemicals shall be used in all tests. Unless otherwise indicated, it is intended that all reagents conform to the specifications of the Committee on Analytical Reagents of the American Chemical Society where such specifications are available.⁵ 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 *Typical Dispersing Media*—Dissolve 5.0 g of sodium hexametaphosphate [(NaPO₃)₆] in 1000 mL of distilled or deionized water. Alternately, dissolve 5.0 g Darvan C⁶ (ammonium salt of polymethacrylic acid) in 95 mL of distilled or deionized water. This latter solution is typically used after dilution of 0.03 g per 20 mL of distilled or deionized water.

7.3 *pH Adjusters*—Fresh ammonium hydroxide (NH₄OH) at 5 to 10 weight % strength is a common reagent used to raise pH, while fresh nitric acid (HNO₃) at 5 to 10 weight % strength is a common reagent to lower pH.

7.4 *Cleaning Solution*—Prepare a 0.1 % solution by volume of Triton X-100 using distilled or deionized water,⁷ or other suitable laboratory cleaning solution.

⁴ The sole instrument of this type known to the committee at this time is the SediGraph X-ray gravity sedimentation particle size analyzer, available from Micromeritics Instrument Corporation, 4356 Communications Drive, Norcross, GA 30093. If you are aware of alternative suppliers, please provide this information to ASTM International Headquarters. Your comments will receive careful consideration at a meeting of the responsible technical committee,¹ which you may attend.

⁵ *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 *Analar 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.

⁶ Darvan C is a trademarked product of R.T. Vanderbilt Company, Inc., Norwalk, CT 06856-5150.

⁷ Triton X-100 is a trademarked product of Rohm & Haas, Philadelphia, PA and is available from a number of laboratory supply companies.

³ National Institute of Standards and Technology (NIST), 100 Bureau Dr., Stop 2300, Gaithersburg, MD 20899-2300, <http://www.nist.gov>.