



Standard Specification for Nuclear Grade Zirconium Oxide Pellets¹

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1. Scope

1.1 This specification applies to pellets of stabilized zirconium oxide used in nuclear reactors.

1.2 The values stated in SI units are to be regarded as the standard. The values given in parentheses are for information only.

2. Referenced Documents

2.1 ASTM Standards:²

C559 Test Method for Bulk Density by Physical Measurements of Manufactured Carbon and Graphite Articles

C859 Terminology Relating to Nuclear Materials

C1065 Specification for Nuclear-Grade Zirconium Oxide Powder

E105 Practice for Probability Sampling of Materials

2.2 ANSI Standard:³

ANSI/ASME NQA-1 Quality Assurance Program Requirements for Nuclear Facility Applications

2.3 U.S. Government Document:⁴

Code of Federal Regulations, Title 10, Part 50—Energy (10 CFR 50) Domestic Licensing of Production and Utilization Facilities

3. Terminology

3.1 Terms shall be defined in accordance with Terminology C859, except for the following:

3.2 *buyer*—the organization issuing the purchase order.

3.3 *pellet*—a fabricated geometric shape of zirconium oxide having a chemical composition as described in Section 4.

3.4 *pellet lot*—the pellets produced from one zirconium oxide powder lot using one set of process parameters. Pellet lot size shall be agreed upon between the seller and the buyer.

3.5 *phase transformation*—the rearrangement of the atomic ordering of a crystalline lattice as a material is cycled through a critical transformation or inversion temperature. The change from one crystalline phase to another may be accompanied by a volume change that could lead to cracks or defects in articles fabricated from such materials.

3.6 *powder lot*—a specified quantity of zirconium-oxide powder with stabilizing additive, blended together such that samples taken in accordance with Section 7 can be considered as representative of the entire quantity.

3.7 *seller*—the zirconium oxide pellet supplier.

3.8 *stabilizing additive*—A material which, when present in sufficient concentration in the subject material exhibiting the phase transformation, produces a stabilized crystalline phase that does not undergo a transformation or inversion at any temperature within the expected fabrication or usage regime of the manufactured pellet. The potentially deleterious volume change is therefore avoided.

4. Chemical Composition

4.1 The starting zirconium oxide powder shall be in accordance with Specification C1065.

4.2 A stabilizing additive shall be used with the zirconium oxide. The recommended stabilizing additive is either calcium oxide (CaO) or yttrium oxide (Y_2O_3). The recommended additive concentration in the case of CaO stabilization is 4 to 8 weight %. In the case of Y_2O_3 stabilization, the recommended additive concentration is 14 to 20 weight %.

4.3 Use analytical chemistry methods as agreed upon between the buyer and seller.

4.4 The impurity concentration excluding the stabilizing additives, shall not exceed 0.5 weight %. Individual element limits are specified in Table 1. The buyer may specify additional limits for any other elements not listed in Table 1.

4.5 The moisture concentration is included in the total hydrogen limit (see Table 1).

5. Physical Requirements

5.1 Physical Dimensions:

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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 American National Standards Institute (ANSI), 25 W. 43rd St., 4th Floor, New York, NY 10036, <http://www.ansi.org>.

⁴ Available from U.S. Government Printing Office Superintendent of Documents, 732 N. Capitol St., NW, Mail Stop: SDE, Washington, DC 20401, <http://www.access.gpo.gov>.

TABLE 1 Impurity Concentration Limits

Element	Maximum Concentration Limit (µg/g pellet)
Hf	200
B	100
Gd	50
Gd + Sm + Eu + Dy	200
Co	100
Si	2000
Fe	1000
Ca ^A	3000
Mg	1200
Al	1500
Ti	100
Th	400
F	30
F + Cl + Br + I	100
H (total hydrogen from all sources)	2

^A This number will be higher if used as a stabilizing addition.

5.1.1 Dimensional requirements shall be in accordance with applicable drawings and purchase order documents.

5.1.2 Pellet dimensions shall be measured to ensure compliance with the buyer's requirements. Sampling plans to meet the acceptance criteria shall be agreed upon between the buyer and the seller to ensure that the pellets represented by the sample are within the required tolerance.

5.2 Density:

5.2.1 Pellet density limits shall be specified by the buyer. The incorporation of a stabilizing additive will introduce a change in the theoretical density of the fabricated pellets and should be taken into account. The method of establishing the theoretical density value shall be agreed upon by the buyer and seller.

5.2.2 The method of density measurement shall be Test Method C559 or an alternative method submitted by the seller for approval by the buyer. Sampling plans to meet the acceptance criteria shall be agreed upon between the buyer and the seller. The method of density measurement and the method of compliance with 5.2.1 shall be approved by the buyer prior to use.

5.3 *Mechanical Properties*—Required mechanical properties and test methods shall be mutually agreed upon between the buyer and the seller. A compression test at 69 MPa (10 000 psi) to ensure pellet integrity is recommended as a suitable test if the pellet is not to be subjected to service temperatures above 1100°C (2012°F). At higher temperatures, a thermal cycling test as given in Appendix X1 followed by a compressive test may be agreed upon between buyer and seller; or alternately, a powder X-ray diffraction pattern may be used to determine that the material has been stabilized in the correct crystalline phase.

5.4 *Visual Appearance*—Visual examination shall be conducted on finished pellets in accordance with Section 7 on Sampling. The seller and the buyer shall agree on visual standards as representing the requirements of 5.4.1, 5.4.2, and 5.4.3. These standards shall be used as acceptance standards for the visual examination of the pellets. In the event of a dispute, the method of defect measurement shall be approved by the buyer prior to use. Maximum permissible defects are defined as follows:

5.4.1 *End Chips*—Pellet end surface shall not be chipped beyond 10 % of the end-face surface area, and no chip shall exceed 1.02 mm (0.040 in.) in depth.

5.4.2 *Circumferential Chips*—Pellet circumferential surfaces shall not be chipped beyond 5 % of the circumferential surface area. No single chip shall exceed a depth of 1.02 mm (0.040 in.).

5.4.3 *Cracks*—Cracks are acceptable, providing the total length of all cracks does not exceed 90 degrees of circumference, and other requirements of the specification are met.

5.4.4 Fissures and other defects shall be evaluated with respect to the criteria of 5.4.1, 5.4.2, and 5.4.3.

6. Cleanliness

6.1 The finished pellets shall be handled in a manner to avoid contamination by grinding fluids and dust, cleaning agents, and organic materials such as plastics and paper used in packaging. Cleaning solutions, if used, shall be free of halides or nonvolatile additives and shall be removed from the pellets prior to sampling and packaging.

7. Sampling

7.1 Sampling plans to meet the acceptance criteria and inspection and measurement procedures that describe the method of compliance with this specification shall be approved by the buyer. The degree of sampling varies with the application and for this reason should be specified on the purchase order. Practice E105 is referenced as a guide.

7.2 Powder and pellet samples shall be taken for quality verification tests, acceptance tests, referee tests and archive samples, as needed.

7.3 Archive samples shall be retained and disposed of in accordance with the buyer's instructions.

8. Inspection and Certification

8.1 The seller shall inspect the material covered by this specification and shall furnish the buyer with certificates of test showing the results of testing and inspection performed on each pellet lot prior to shipment. The seller shall certify that each pellet lot is in compliance with the provisions of this specification.

9. Rejection

9.1 Unless the buyer and seller agree otherwise, rejection and acceptance shall be on a pellet lot basis.

9.2 Pellet lots that fail to conform to the requirements of this specification may be rejected by the buyer. The seller may petition the buyer to waive selected requirements for identified out-of-specification lots. The decision to grant such waiver belongs to the buyer. The seller may also apply any remedy to bring rejected lots into specification providing he can demonstrate to the buyer that such remedy does not impair the function or preclude the certification of the rejected material.

9.3 In the event of disagreement over the results of analysis, samples shall be submitted to a mutually selected referee for resolution.

10. Packaging and Shipping

10.1 The pellets shall be packaged in sealed containers to prevent loss or damage, or both, of material and contamination from airborne or container materials. The exact size and type of packaging shall be as mutually agreed upon between the buyer and seller.

10.2 Each container shall be clearly marked with the following:

10.2.1 Zirconium oxide pellets plus stabilizing additive,

10.2.2 Purchase order number,

10.2.3 Gross, tare, and net weights,

10.2.4 Lot number, and

10.2.5 Name of pellet manufacturer.

11. Quality Assurance

11.1 Quality assurance requirements shall be agreed upon between the buyer and seller when specified in the purchase order. Title 10 CFR, Part 50, Appendix B and ANSI/ASME NQA-1 are referenced as guides.

12. Keywords

12.1 CaO; stabilizer; stabilizing additive; thermal cycling test; Y₂O₃; zirconium oxide

APPENDIX

(Nonmandatory Information)

X1. THERMAL CYCLING TEST

X1.1 The following thermal cycle is recommended as a guide to determine the thermal stability of zirconium-oxide pellets.

X1.1.1 Heat the pellets in air to $1100 \pm 25^{\circ}\text{C}$ ($2012 \pm 77^{\circ}\text{F}$) at a heating rate of at least 100°C/h (212°F/h).

X1.1.2 Hold at $1100 \pm 25^{\circ}\text{C}$ ($2012 \pm 77^{\circ}\text{F}$) for at least 1 h.

X1.1.3 Cool to 500°C (932°F) at a cooling rate of at least 100°C/h (212°F/h).

X1.1.4 Repeat steps **X1.1.1 – X1.1.3** two additional times.

X1.1.5 Cool to room temperature and remove for mechanical testing.

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