



# National Institute of Standards & Technology

## Certificate of Analysis

### Standard Reference Material<sup>®</sup> 3164

#### Spectrometric Standard Solution

#### Uranium

#### Batch Code 590811

This Standard Reference Material (SRM) is intended for use in atomic absorption spectrometry, inductively coupled plasma spectrometry, spectrophotometry, or any other analytical technique that requires aqueous standard solutions for calibrating instruments. SRM 3164 is a single element solution prepared gravimetrically to contain a nominal 10 mg/mL of uranium with an approximate nitric acid volume fraction of 10 %. The certified value ( $Y$ ) is based on the mass of high-purity oxide dissolved and diluted to known volume. The value has been adjusted upward by 0.1 % relative, based on estimated transpiration losses of solvent through the container walls of 0.2 % relative per year. The density of the solution at 22 °C is 1.078 g/mL.

Metal	Concentration (mg/mL)	Source Purity, %	Acid Volume Fraction Approximate
Uranium	10.01 ± 0.03	NBL CRM 129, U <sub>3</sub> O <sub>8</sub> (99.968)*	HNO <sub>3</sub> , 10 %

\*The total impurities in NBLCRM 129 were determined by spectrochemical analysis and are estimated to be less than 50 mg/kg. The CRM was obtained from the U.S. Department of Energy New Brunswick Laboratory (NBL), Argonne, IL. It is a limited quantity radioactive material that is exempt from radioactive labeling requirements under 49CFR section 173.421. The massic activity of SRM 3164 is less than 500 Bq/g. To determine proper handling, storage, and disposal of this SRM refer to appropriate state and federal regulations.

The uncertainty in the certified value is calculated as

$$U = (2u_c + 0.001Y) \text{ mg/mL}$$

where  $u_c$  the "combined uncertainty" calculated according to the ISO Guide [1,2]. The value  $u_c$  is intended to represent, at the level of one standard deviation, the combined effect of uncertainty components associated with volumetric and gravimetric factors, as well as the purity of the starting material. The additional quantity, 0.001 $Y$ , is an allowance for transpiration of the solution through the container walls, which is estimated to be ± 0.1 % of the certified value during the one-year period of validity of the certification.

The combined uncertainty consists of a Type A component associated with replicate weighings of the uranium oxide and Type B components due to uncertainty in the material purity, material handling, and dilution.

SRM 3164 was prepared by T.A. Butler of the NIST Analytical Chemistry Division. Inductively coupled plasma spectrometric analysis was performed by L.J. Wood of the NIST Analytical Chemistry Division.

The technical and support aspects involved in the preparation, certification, and issuance of the SRM were coordinated through the Standard Reference Materials Program by B.S. MacDonald.

Gaithersburg, MD 20899  
February 7, 1996  
(Revision of certificate dated 12-12-95)

Thomas E. Gills, Chief  
Standard Reference Materials Program

## Procedures for Use

**Stability:** This certification is valid for one year from the shipping date, provided the SRM solution is kept tightly capped and stored under normal laboratory conditions. NIST will monitor the stability of representative solutions from the SRM lot, and if any changes occur that invalidate this certification, NIST will notify purchasers.

**Preparation of Working Standard Solutions:** All solutions should be brought to  $22\text{ }^{\circ}\text{C} \pm 1\text{ }^{\circ}\text{C}$  before use and all glass or plastic surfaces coming into contact with the standard must have been previously cleaned. A working standard solution can be prepared from the SRM solution by serial dilution. Dilutions should be made with certified volumetric class A flasks and 5 mL or 10 mL class A pipets. All volumetric transfers of solutions should be performed using a proven analytical technique. Each dilution should be acidified with an appropriate high-purity acid and diluted to calibrated volume using high-purity water. The stability of the working standard solution will depend on the final acid concentration; therefore, care should be exercised to ensure that the final acid concentration of the dilution closely approximates that of the SRM. The analyst should prepare daily working solutions from 100  $\mu\text{g}/\text{mL}$  dilutions of the original SRM solution.

## NOTICE AND WARNING TO USERS

For some instrumental techniques, small differences in acid type and concentration between the SRM and sample may lead to erroneous results. Therefore, the same acid mixture as is listed on this SRM certificate should be used in making appropriate dilutions and working standards.

## REFERENCES

- [1] *Guide to the Expression of Uncertainty in Measurement*, ISBN 92-67-10188-9, 1st Ed. ISO, Geneva, Switzerland, (1993).
- [2] Taylor, B.N. and Kuyatt, C.E., "Guidelines for Evaluating and Expressing the Uncertainty of NIST Measurement Results", NIST Technical Note 1297, U.S. Government Printing Office, Washington, D.C., (1994).