



Certificate of Analysis

Standard Reference Material[®] 2062

TiAl(NbW) Alloy

This Standard Reference Material (SRM) is intended for use in X-ray fluorescence analysis of titanium-aluminum (Ti-Al) aerospace alloys although it may be a useful reference for other methods of analysis. A unit of SRM 2062 is a flat, polished disk approximately 2.4 cm diameter and 2 mm thick.

Certified Values: A NIST-certified value is a value for which NIST has the highest confidence in its accuracy, in that all known or suspected sources of bias have been investigated or accounted for by NIST. A certified value is the present best estimate of the true value based on the results of analyses performed at NIST. The uncertainty listed with the value is an expanded uncertainty ($k = 2$, 95 % confidence interval [1]) calculated in accordance with the ISO and NIST Guides [2].

Certified values for titanium (Ti), aluminum (Al), niobium (Nb), and tungsten (W), expressed as mass fractions [2], are provided in Table 1. The results are expressed as the certified value \pm an expanded uncertainty. The certified values are the unweighted averages of the concentrations determined by wavelength dispersive X-ray fluorescence spectrometry (WD-XRF) and inductively-coupled plasma optical emission spectrometry (ICP-OES) [3]. The expanded uncertainties for Ti, Al, Nb, and W include the uncertainties attributable to specimen heterogeneity of these elements in addition to the uncertainties from the two quantitative methods.

Testing of the specimen to specimen heterogeneity was performed with WD-XRF and as part of the wavelength dispersive electron probe microanalysis (WD-EPMA) of these specimens. Value assignment categories are based on the definition of terms and modes used at NIST for chemical reference materials [4].

Table 1. Certified Values with Expanded Uncertainties for SRM 2062

Element	Mass Fraction (%)
Titanium	53.92 \pm 0.34
Aluminum	30.31 \pm 0.31
Niobium	10.78 \pm 0.10
Tungsten	4.38 \pm 0.11

Expiration of Certification: The certification of this SRM is valid until **01 January 2029**, within the uncertainty specified, provided the SRM is handled and stored in accordance with the instructions given in this certificate (see "Instructions for Use"). However, the certification may be nullified if the SRM is damaged or contaminated.

Coordination of the technical measurements for certification was accomplished under the direction of R.B. Marinenko of the NIST Surface and Microanalysis Division.

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Certificate Issue Date: 16 May 2005

Robert L. Watters, Jr., Chief
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Heterogeneity and microheterogeneity testing was performed by R.B. Marinenko and J. Kessler of the NIST Surface and Microanalysis Science Division and by R. Herrera-Basurto, Guest Researcher from Centro Nacional de Metrologia, Querétaro, Mexico.

WD-XRF quantitative characterization as well as specimen to specimen heterogeneity testing for certification of this SRM were performed at NIST by J.R. Sieber and A. F. Marlow of the NIST Analytical Chemistry Division. ICP-OES quantitative characterization for certification were performed by L.L. Yu and T.A. Butler, also of the NIST Analytical Chemistry Division.

Statistical consultation for this SRM was provided by S.D. Leigh of the NIST Statistical Engineering Division.

The support aspects involved in the issuance of this SRM were coordinated through the NIST Standard Reference Materials Program by B.S. MacDonald of the Measurement Services Division.

Information Values: An information value is considered to be a value that will be of interest and use to the SRM user, but insufficient information is available to assess the uncertainty associated with the value. Information values for oxygen, and nitrogen are provided in Table 2. The oxygen and nitrogen values are based on inert gas fusion (IGF) analysis.

Table 2. Information Values for SRM 2062

Element	Mass Fraction (%)
Oxygen	0.232
Nitrogen	0.004

Stability: This material is considered to be stable during the period of certification. NIST will monitor this SRM over the period of its certification. If substantive technical changes occur that affect the certification before the expiration of this certificate, NIST will notify the purchaser. Registration (see attached sheet) will facilitate notification.

PREPARATION, HETEROGENEITY, AND ANALYSIS¹

Material Preparation: The material for SRM 2062 was fabricated at Wright-Patterson Air Force Base (Dayton, OH). It was extruded in the form of a 2.4 cm diameter rod that was annealed at 1370 °C to produce only the α -phase of the alloy. The rod was cut into 23 disks, 2 mm thick, that were marked on the unpolished side with a number and letter to identify their original locations in the rod. These disks were polished at NIST on one side and cleaned for EPMA heterogeneity testing. Upon completion of the heterogeneity tests, six of the disks were sliced into 2 mm × 2 mm × 2 mm cubes with a diamond saw. Most of the cubes were retained for use as SRM 2061 for microanalysis while a few were used for quantitative analyses by WD-XRF, ICP-OES, and IGF. The remaining disks were retained for use as SRM 2062.

Heterogeneity: Each of the 23 disks were tested with WD-EPMA for heterogeneity using three independent sampling strategies:

1. 40-point traverses using 2 μ m steps – 2 traverses normal to one another per specimen
2. 20 random point samplings per specimen
3. Nested design sampling using 3 duplicate readings on 4 randomly selected points per specimen to determine the variances for the between specimen and the between points heterogeneities [5].

The total uncertainties determined from test No. 3 are included in the certified uncertainties assigned to each element. In addition, 15 disks were tested for between specimen heterogeneity with WD-XRF.

¹Certain commercial equipment, instruments, or materials are identified in this certificate in order to adequately specify the experimental procedure. Such identification does not imply recommendation or endorsement by the National Institute of Standards and Technology, nor does it imply that the materials or equipment identified are necessarily the best available for the purpose.

Analysis: For WD-XRF analysis [5], acid digestion of single cubes of the alloy (6 total, approx. 40 mg each) was followed by borate fusion with a pre-fused flux (50:50 mix of $\text{Li}_2\text{B}_4\text{O}_7$ and LiBO_2). Calibration was accomplished using synthetic calibrants designed to match the fused unknowns and prepared from high-purity compounds. The $\text{K L}_{2,3}$ ($\text{K}\alpha$) X-ray lines for Al, Ti, and Nb and the $\text{L}_{3\text{-}M_{4,5}}$ ($\text{L}\alpha$) lines for W were used in the analysis.

For ICP-OES analysis [5], single cubes of the alloy were digested in HCl and HF. Manganese was used as an internal standard. For each element, calibration was accomplished with six matrix-matched standards and a linear calibration model. The analytes Al, Ti, Nb, W, and the internal standard, Mn, were measured side-on with respect to the plasma at 394.401 nm, 337.279 nm, 269.706 nm, 239.708 nm, and 260.568 nm, respectively. Analyses were done on two different days and averaged.

Oxygen and nitrogen were analyzed by Luvak, Inc. (Boylston, MA) using IGF. Two of the 2 mm × 2 mm × 2 mm cubes were used for duplicate analyses of both elements.

INSTRUCTIONS FOR USE²

Storage: The alloy should be stored in a closed container such as a plastic box, in a cool, dry location, preferably a desiccator, to avoid dust or moisture contamination.

Use: To relate analytical determinations to the values on this certificate of analysis, the polished side of the specimen must be used in XRF analysis. Avoid analysis of the region within 1 mm of the edge of the specimen. Handle the specimen with clean, powder-free gloves to avoid contamination with body oils and any kind of foreign matter. Avoid contact of any kind with the polished surface once it is clean and dry. If repolishing is required, particles of the polishing material may adhere or become imbedded in the surface. Most can be removed with Scotch[®] Removable Magic[®] Tape or similar transparent (“invisible”) tape followed by thorough rinsing with heptane to ensure removal of any remaining tape adhesive.

Minimum Sample Quantity: For X-ray fluorescence spectrometry, practically any beam size may be used. The material is suitable for measurement by micro X-ray fluorescence spectrometry in which small X-ray beams are employed. For beams smaller than about 1 mm in diameter and based on the microheterogeneity assessment, the sampling of at least 10 different randomly selected points is recommended for Al, Ti, and Nb. A 20-point sampling is recommended for W.

REFERENCES

- [1] Hahn, G.J.; Meeker, W.Q.; *Statistical Intervals: A Guide for Practitioners*; John Wiley & Sons, Inc.: New York (1991).
- [2] ISO; *Guide to the Expression of Uncertainty in Measurement*; ISBN 92-67-10188-9, 1st ed.; International Organization for Standardization: Geneva, Switzerland (1993); see also Taylor, B.N.; Kuyatt, C.E.; *Guidelines for Evaluating and Expressing the Uncertainty of NIST Measurement Results*; NIST Technical Note 1297; U.S. Government Printing Office: Washington, DC (1994); available at <http://physics.nist.gov/Pubs/>.
- [3] Levenson, M.S.; Banks, D.L.; Eberhardt, K.R.; Gill, L.M.; Guthrie, W.F.; Liu, H-K.; Vangel, M.G.; Yen, J.H.; Zhang, N.F.; *An Approach to Combining Results from Multiple Methods Motivated by the ISO GUM*; J. Res. Natl. Inst. Stand. Technol.; Vol. 105, pp. 571–579 (2000).
- [4] May, W.E.; Parris, R.M.; Beck II, C.M.; Fassett, J.D.; Greenberg, R.R.; Guenther, F.R.; Kramer, G.W.; Wise, S.A.; Gills, T.E.; Colbert, J.C.; Gettings, R.J.; MacDonald, B.S.; *Definitions of Terms and Modes Used at NIST for Value-Assignment of Reference Materials for Chemical Measurements*; NIST Special Publication 260–136, p.16; U.S. Government Printing Office: Washington, DC; (2000).
- [5] Sieber, J.R.; Yu, L.L.; Marlow, A.F.; Butler, T.A.; *Traceability and Uncertainty in Alloy Analysis by Borate Fusion and XRF*; X-Ray Spectrom., Vol. 34, pp. 153-159 (2004).

Users of this SRM should ensure that the certificate in their possession is current. This can be accomplished by contacting the SRM Program at: telephone (301) 975-6776; fax (301) 926-4751, email srminfo@nist.gov; or via the Internet at <http://www.nist.gov/srm>.

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