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CERTIFICATE OF ANALYSIS

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1.0 ACCREDITATION / REGISTRATION

INORGANIC VENTURES is accredited to ISO Guide 34, "General Requirements for the Competence of Reference Material Producers" and ISO/IEC 17025, "General Requirements for the Competence of Testing and Calibration Laboratories". Inorganic Ventures is also an ISO 9001 registered manufacturer (SAI Global File Number 010105).



2.0 PRODUCT DESCRIPTION

Product Code: Single Analyte Mass Spec Solution

Catalog Number: MSSN-10PPM

Lot Number: J2-SN02082

Matrix: 5% (v/v) HNO3

tr. HF

Value / Analyte(s): 10 μg/mL ea:

Sn

Starting Material: Sn shot
Starting Material Lot#: 1744

Starting Material Lot#: 1744

Starting Material Purity: 99.9998%

3.0 CERTIFIED VALUES AND UNCERTAINTIES

Certified Value: $9.990 \pm 0.070 \,\mu\text{g/mL}$

Certified Density: 1.024 g/mL (measured at 20 ± 1 °C)

Assay Information:

 ANALYTE
 METHOD
 NIST SRM#
 SRM LOT#

 Sn
 Calculated
 See Sec. 4.2

 Sn
 ICP Assay
 3161a
 070330

The following equations are used in the calculation of the certified value and the uncertainty. Reported uncertainties represent expanded uncertainties expressed at approximately the 95% confidence level using a coverage factor of k = 2.

Characterization of CRM/RM by Two Methods

Certified Value, $X_{CRM/RM}$, where two methods of characterization are used is the weighted mean of the two results:

 $\mathsf{X}_{\mathsf{CRM/RM}} = [(\mathsf{w}_{\mathsf{a}})\;(\mathsf{X}_{\mathsf{a}}) + (\mathsf{w}_{\mathsf{b}})\;(\mathsf{X}_{\mathsf{b}})]$

 $\mathbf{X_a}$ = mean of Assay Method A with standard uncertainty $\mathbf{u_{char}}$ a

X_b = mean of Assay Method B with standard uncertainty u_{char b}

w_a and w_b = the weighting factors for each method calculated using the inverse square of the variance:

 $\mathbf{w_a} = (1/u_{\text{char a}})^2 / ((1/u_{\text{char a}})^2 + (1/u_{\text{char b}})^2))$ $\mathbf{w_b} = (1/u_{\text{char b}})^2 / ((1/u_{\text{char a}})^2 + (1/u_{\text{char b}})^2))$

CRM/RM Expanded Uncertainty (±) = $U_{CRM/RM} = k (u_{char}^2 + u_{bb}^2 + u_{lts}^2 + u_{sts}^2)^{1/2}$

k = coverage factor = 2 in all cases at Inorganic Ventures

 $\mathbf{u_{bb}}$ = bottle to bottle homogeneity standard uncertainty

u_{lts} = long term stability standard uncertainty (storage)

u_{sts} = short term stability standard uncertainty (transportation)

Characterization of CRM/RM by One Method

Certified Value, X_{CRM/RM}, where one method of characterization is used is the mean of individual results:

X_{CRM/RM} = mean of Assay Method A with standard uncertainty u_{char a}

CRM/RM Expanded Uncertainty (±) = $U_{CRM/RM} = k (u^2_{char} a + u^2_{bb} + u^2_{lts} + u^2_{sts})^{1/2}$

k = coverage factor = 2 in all cases at Inorganic Ventures

u_{char a} = square root of the sum of the squares of the errors from characterization which include instrumental measurement, density, NIST SRM uncertainty, weighing, and volume

u_{bb} = bottle to bottle homogeneity standard uncertainty

u_{lts} = long term stability standard uncertainty (storage)

u_{sts} = short term stability standard uncertainty (transportation)

4.0 TRACEABILITY TO NIST

- This product is traceable to NIST via an unbroken chain of comparisons. The uncertainties for each certified value are reported, taking into account the SRM/RM uncertainty error and the measurement, weighing and volume dilution errors. In rare cases where no NIST SRM/RM are available, the term 'in-house std.' is specified.

4.1 Thermometer Calibration

- All thermometers are NIST traceable through thermometers that are calibrated by an accredited calibration laboratory.

4.2 Balance Calibration

- All analytical balances are calibrated by an accredited calibration laboratory and procedure. The weights used for testing are annually compared to master weights and are traceable to NIST.

4.3 Glassware Calibration

- An in-house procedure is used to calibrate all Class A glassware used in the manufacturing and quality control of CRM/RMs.

5.0 TRACE METALLIC IMPURITIES (TMI) DETERMINED BY ICP-MS AND ICP-OES (µg/mL)

CRM/RMs are tested for trace metallic impurities by Axial ICP-OES and ICP-MS. The result from the most sensitive method for each element, is reported below. Solutions tested by ICP-MS were analyzed in an ULPA-Filtered Clean Room. An ULPA-Filter is 99.9985% efficient for the removal of particles down to $0.3 \, \mu m$.

М	Ag	<	0.005803	М	Eu	<	0.008705	0	Na		0.000003	0	Se	<	0.050000	0	Zn	<	0.000200
0	Αl		0.000010	0	Fe		0.000001	M	Nb	<	0.001450	0	Si	<	0.003400	М	Zr	<	0.014509
0	As	<	0.005000	0	Ga	<	0.001600	M	Nd	<	0.005803	M	Sm	<	0.002901				
M	Au	<	0.008705	M	Gd	<	0.002901	0	Ni	<	0.010000	s	Sn	<					
0	В	<	0.012000	0	Ge	<	0.004000	n	Os	<		M	Sr	<	0.001450				
0	Ва	<	0.000700	M	Hf	<	0.005803	0	Р	<	0.005000	M	Та	<	0.020312				
M	Ве	<	0.001450	0	Hg	<	0.015000	0	Pb	<	0.010000	M	Tb	<	0.000870				
M	Bi	<	0.001160	M	Но	<	0.001450	M	Pd	<	0.014509	0	Те	<	0.050000				
0	Ca			0	In	<	0.002000	M	Pr	<	0.000870	M	Th	<	0.002901				
0	Cd	<	0.005000	M	lr	<	0.014509	M	Pt	<	0.005803	M	Ti	<	0.145091				
0	Се	<	0.010000	0	K		0.000004	M	Rb	<	0.002901	M	TI	<	0.002901				
0	Co	<	0.002000	M	La	<	0.001450	M	Re	<	0.002901	M	Tm	<	0.001160				
0	Cr	<	0.001000	0	Li	<	0.000020	M	Rh	<	0.002901	M	U	<	0.005803				
M	Cs	<	0.000870	M	Lu	<	0.001160	M	Ru	<	0.005803	0	V	<	0.001200				
M	Cu	<	0.017410	0	Mg		0.000001	n	S	<		M	W	<	0.029018				
M	Dy	<	0.017410	M	Mn	<	0.011607	0	Sb	<	0.010000	M	Υ	<	0.116072				
M	Er	<	0.014509	M	Мо	<	0.005803	M	Sc	<	0.029018	M	Yb	<	0.002901				

M - Checked by ICP-MS O - Checked by ICP-OES i - Spectral Interference

n - Not Checked For s - Solution Standard Element

6.0 INTENDED USE

- For the calibration of analytical instruments and validation of analytical methods as appropriate.

7.0 INSTRUCTIONS FOR THE CORRECT USE OF THIS REFERENCE MATERIAL

7.1 Storage and Handling Recommendations

- Keep cap tightly sealed when not in use. Store and use at $20 \pm 4^{\circ}$ C. Do not pipette from the container. Do not return removed aliquots to container.

Atomic Weight; Valence; Coordination Number; Chemical Form in Solution - 118.71 +4 4,5, 6,7,8 Sn(OH)xFv2-

Chemical Compatibility -Soluble in HCl and dilute HF / HNO3. Avoid neutral to basic media. Unstable at ppm levels with metals that would pull F- away. (i.e. Do not mix with Alkaline or Rare Earths or high levels of transition elements unless they are fluorinated.) Stable with most inorganic anions provided it is in the chemical form shown above.

Stability - 2-100 ppb levels stable (alone or mixed with all other metals that are at comparable levels) as the Sn(OH)xFy2- for 1 year in 1% HNO3 / LDPE container. 1-10,000 ppm single element solutions as the Sn(OH)xFy2- chemically stable for years in 2-5% HNO3 / trace HF in a LDPE container.

Sn Containing Samples (Preparation and Solution) - Metal (Soluble in HF / HNO3 or HCl); Oxides - SnO (soluble in HCl), SnO2 -very resistant to all acids including HF(Fusion with equal parts of Na2CO3 and S. It is then soluble in water or dilute acids as the thiostannate.); Alloys (Treat first 0.1 g with 10 mL conc. H2SO4 to boiling until the alloy disintegrates and nearly all of the sulfuric acid is expelled. Then add 100 mL O2 free water and 50 mL of conc HCl or transfer to a plastic container and add 1 mL HF in either case warming gently to bring about solution.); Organic Matrices (Volatility and precipitation of the insoluble stannic oxide are problems. Consultation of the literature should be made for individual matrices / Sn compounds.)

Atomic Spectroscopic Information (ICP-OES D.L.s are given as radial/axial view):

Technique/Line	Estimated D.L.	Order	Interferences (underlined indicates severe)
ICP-MS 120 amu	5 ppt	N/A	120Te, 104Ru16O,
			104Pd16O
ICP-OES 189.989 nm	0.03 / 0.003 μg/mL	1	
ICP-OES 242.949 nm	0.1 / 0.01 μg/mL	1	W, Mo, Rh ,Ta, Co

HF Note: This standard should not be prepared or stored in glass.

8.0 HAZARDOUS INFORMATION

- Please refer to the Safety Data Sheet for information regarding this CRM/RM.

9.0 HOMOGENEITY

- This solution was mixed according to an in-house procedure and is guaranteed to be homogeneous. Homogeneity data indicate that the end user should take a minimum sample size of 0.2 mL to assure homogeneity.

10.0 QUALITY STANDARD DOCUMENTATION

10.1 10CFR50 Appendix B - Nuclear Regulatory Commission

- Domestic Licensing of Production and Utilization Facilities

10.2 10CFR21 - Nuclear Regulatory Commission

- Reporting defects and Non-Compliance

10.3 ISO 9001 Quality Management System Registration

- SAI Global File Number 010105

10.4 ISO/IEC Guide 17025 "General Requirements for the Competence of Testing and Calibration Laboratories"

- Chemical Testing - Accredited / A2LA Certificate Number 883.01

10.5 ISO/IEC Guide 34 "General Requirements for the Competence of Reference Material Producers"

- Reference Material Producer - Accredited / A2LA Certificate Number 883.02

11.0 CERTIFICATION, EXPIRATION AND PERIOD OF VALIDITY

11.1 Certification Issue Date

August 17, 2015

11.2 Expiration Date

11.3 Period of Validity

- The certification is valid within the measurement uncertainty specified provided the CRM/RM is handled and stored in accordance with instructions given in Sec 7.0 and used prior to the date given in Sec 11.2. This certification is nullified if the CRM/RM is damaged, contaminated, or otherwise modified.

12.0 NAMES AND SIGNATURES OF CERTIFYING OFFICERS

Certificate Prepared By:

James King Jr Product Documentation Supervisor

Certificate Approved By:

Michael Booth QC Supervisor

Certifying Officer:

Paul Gaines PhD., Senior Technical Director Michael 2 Booth

Paul R Laines