an analytichem company

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CERTIFICATE OF ANALYSIS FOR

CERTIFIED REFERENCE MATERIAL

OREAS 606b

High Sulphidation Epithermal Au-Cu-Ag Ore

(Mt Carlton, Queensland, Australia)

Table 1. Certified Value, Uncertainty & Tolerance Intervals for Au by FA in OREAS 606b.

Constituent	Certified Value†	95% Expanded Uncertainty		95% Tolerance Limits	
		Low	High	Low	High
Pb Fire Assay					
Au, Gold (ppm)	0.386	0.380	0.391	0.384*	0.387*

SI unit equivalents: ppm (parts per million; 1 x 10^{-6}) ≡ mg/kg. Note: intervals may appear asymmetric due to rounding.

†The operationally defined measurand meets the requirements of ISO 17034 and all participating laboratories comply with the requirements of ISO 17025.

*Gold Tolerance Limits for typical 30g fire assay are determined from 20 x 85mg INAA results and the Sampling Constant (Ingamells & Switzer, 1973).

Table 2. Certified Value, Uncertainty & Tolerance Intervals for other measurands in OREAS 606b.

Constituent	Certified Value	95% Expanded Uncertainty		95% Tolerance Limits	
		Low	High	Low	High
Aqua Regia Digestion (sample weights 10-50g)					
Au, Gold (ppm)	0.375	0.362	0.388	0.373*	0.377*
Infrared Combustion					
S, Sulphur (wt.%)	0.460	0.447	0.473	0.446	0.474
4-Acid Digestion					
Ag, Silver (ppm)	1.53	1.44	1.62	1.46	1.59
Al, Aluminium (wt.%)	7.54	7.29	7.78	7.39	7.68
As, Arsenic (ppm)	89	85	93	86	92
Ba, Barium (ppm)	2892	2797	2987	2821	2963
Be, Beryllium (ppm)	2.98	2.81	3.15	2.88	3.08
Bi, Bismuth (ppm)	4.83	4.60	5.06	4.72	4.94
Ca, Calcium (wt.%)	1.10	1.07	1.13	1.07	1.13
Cd, Cadmium (ppm)	1.24	1.13	1.35	1.16	1.32
Ce, Cerium (ppm)	89	85	93	87	91
Co, Cobalt (ppm)	2.25	2.13	2.38	2.13	2.37
Cr, Chromium (ppm)	14.7	12.9	16.5	13.2	16.2
Cs, Caesium (ppm)	6.11	5.79	6.44	5.98	6.25
Cu, Copper (wt.%)	0.033	0.032	0.034	0.033	0.034
Dy, Dysprosium (ppm)	3.55	3.38	3.73	3.38	3.73
Er, Erbium (ppm)	1.01	0.93	1.08	0.96	1.05
Eu, Europium (ppm)	1.40	1.26	1.53	1.36	1.44
Fe, Iron (wt.%)	2.33	2.26	2.40	2.27	2.39
Ga, Gallium (ppm)	23.0	22.0	24.0	22.5	23.6
Gd, Gadolinium (ppm)	5.60	5.09	6.12	5.43	5.78
Ge, Germanium (ppm)	0.19	0.13	0.26	0.17	0.22
Hf, Hafnium (ppm)	6.48	6.17	6.80	6.25	6.71
Ho, Holmium (ppm)	0.48	0.44	0.53	0.45	0.51
In, Indium (ppm)	0.14	0.13	0.15	0.13	0.15

SI unit equivalents: ppm (parts per million; 1 x 10^{-6}) ≡ mg/kg; wt.% (weight per cent) ≡ % (mass fraction).

*Gold Tolerance Limits for typical 15-50g aqua regia digestion methods are determined from 20 x 85mg INAA results and the Sampling Constant (Ingamells & Switzer, 1973).

Note: intervals may appear asymmetric due to rounding.

Table 2 continued.

Constituent	Certified Value	95% Expanded Uncertainty		95% Tolerance Limits	
		Low	High	Low	High
4-Acid Digestion continued					
K, Potassium (wt.%)	3.02	2.92	3.11	2.94	3.09
La, Lanthanum (ppm)	45.0	42.8	47.2	43.6	46.4
Li, Lithium (ppm)	32.1	30.8	33.4	31.3	32.9
Lu, Lutetium (ppm)	0.074	0.058	0.091	IND	IND
Mg, Magnesium (wt.%)	0.127	0.120	0.133	0.124	0.129
Mn, Manganese (wt.%)	0.028	0.027	0.029	0.028	0.029
Mo, Molybdenum (ppm)	4.38	4.12	4.65	4.21	4.56
Na, Sodium (wt.%)	2.67	2.59	2.76	2.61	2.74
Nb, Niobium (ppm)	18.5	17.7	19.3	18.0	19.0
Nd, Neodymium (ppm)	37.4	35.0	39.9	36.3	38.5
Ni, Nickel (ppm)	3.56	3.13	3.99	3.29	3.83
P, Phosphorus (wt.%)	0.030	0.029	0.031	0.029	0.031
Pb, Lead (ppm)	157	151	163	153	161
Pr, Praseodymium (ppm)	10.1	9.4	10.7	9.8	10.4
Rb, Rubidium (ppm)	139	133	145	136	142
Re, Rhenium (ppm)	< 0.002	IND	IND	IND	IND
S, Sulphur (wt.%)	0.468	0.450	0.486	0.457	0.479
Sb, Antimony (ppm)	6.38	6.04	6.71	6.07	6.68
Sc, Scandium (ppm)	4.26	3.91	4.61	4.13	4.39
Se, Selenium (ppm)	1.80	1.11	2.48	1.67	1.93
Sm, Samarium (ppm)	7.27	6.89	7.65	7.08	7.46
Sn, Tin (ppm)	3.72	3.54	3.89	3.59	3.84
Sr, Strontium (ppm)	200	192	207	195	204
Ta, Tantalum (ppm)	1.36	1.29	1.44	1.32	1.41
Tb, Terbium (ppm)	0.73	0.66	0.80	0.71	0.75
Te, Tellurium (ppm)	0.84	0.73	0.94	0.73	0.94
Th, Thorium (ppm)	14.8	14.1	15.5	14.4	15.2
Ti, Titanium (wt.%)	0.116	0.113	0.120	0.114	0.119
Tl, Thallium (ppm)	0.89	0.85	0.92	0.86	0.91
Tm, Thulium (ppm)	0.10	0.10	0.11	IND	IND
U, Uranium (ppm)	5.57	5.28	5.85	5.39	5.74
V, Vanadium (ppm)	4.97	4.48	5.45	4.65	5.29
W, Tungsten (ppm)	1.85	1.71	1.99	1.73	1.98
Y, Yttrium (ppm)	14.8	14.2	15.5	14.4	15.3
Yb, Ytterbium (ppm)	0.61	0.54	0.68	0.58	0.64
Zn, Zinc (ppm)	316	305	327	308	324
Zr, Zirconium (ppm)	243	232	254	237	249

SI unit equivalents: ppm (parts per million; 1 x 10^{-6}) ≡ mg/kg; wt.% (weight per cent) ≡ % (mass fraction).

Note: intervals may appear asymmetric due to rounding;

IND = indeterminate (due to limited reading resolution of the methods employed. For practical purposes the 95% Expanded Uncertainty can be set between zero and a two times multiple of the upper bound/non-detect limit value).

Table 2 continued.

Constituent	Certified Value	95% Expanded Uncertainty		95% Tolerance Limits	
		Low	High	Low	High
Aqua Regia Digestion					
Ag, Silver (ppm)	1.49	1.40	1.57	1.25	1.72
Al, Aluminium (wt.%)	0.804	0.772	0.835	0.781	0.827
As, Arsenic (ppm)	86	83	89	84	88
B, Boron (ppm)	< 10	IND	IND	IND	IND
Ba, Barium (ppm)	212	198	225	204	219
Be, Beryllium (ppm)	0.51	0.47	0.55	0.49	0.54
Bi, Bismuth (ppm)	4.96	4.73	5.19	4.73	5.19
Ca, Calcium (wt.%)	0.632	0.614	0.650	0.619	0.646
Cd, Cadmium (ppm)	1.19	1.11	1.28	1.13	1.26
Ce, Cerium (ppm)	56	53	59	55	58
Co, Cobalt (ppm)	2.02	1.90	2.14	1.92	2.12
Cr, Chromium (ppm)	16.1	15.1	17.2	15.2	17.1
Cs, Caesium (ppm)	1.48	1.36	1.59	1.42	1.53
Cu, Copper (wt.%)	0.034	0.033	0.035	0.033	0.034
Dy, Dysprosium (ppm)	2.08	1.88	2.29	1.99	2.18
Er, Erbium (ppm)	0.51	0.44	0.58	0.47	0.55
Eu, Europium (ppm)	0.70	0.60	0.79	0.67	0.72
Fe, Iron (wt.%)	2.00	1.94	2.06	1.97	2.03
Ga, Gallium (ppm)	5.90	5.59	6.21	5.70	6.10
Gd, Gadolinium (ppm)	3.82	3.55	4.08	3.66	3.97
Ge, Germanium (ppm)	0.12	0.09	0.16	IND	IND
Hf, Hafnium (ppm)	1.22	1.14	1.29	1.16	1.27
Hg, Mercury (ppm)	0.048	0.033	0.064	IND	IND
Ho, Holmium (ppm)	0.25	0.22	0.27	0.23	0.26
In, Indium (ppm)	0.10	0.10	0.11	0.10	0.11
K, Potassium (wt.%)	0.235	0.220	0.250	0.224	0.246
La, Lanthanum (ppm)	27.9	26.4	29.4	27.2	28.6
Li, Lithium (ppm)	19.9	19.0	20.7	19.3	20.4
Lu, Lutetium (ppm)	0.028	0.019	0.036	IND	IND
Mg, Magnesium (wt.%)	0.088	0.087	0.090	0.087	0.090
Mn, Manganese (wt.%)	0.025	0.024	0.026	0.025	0.026
Mo, Molybdenum (ppm)	4.00	3.70	4.30	3.82	4.18
Na, Sodium (wt.%)	0.075	0.069	0.082	0.070	0.081
Nb, Niobium (ppm)	0.90	0.74	1.06	0.81	1.00
Nd, Neodymium (ppm)	25.5	22.8	28.2	24.8	26.2
Ni, Nickel (ppm)	3.41	3.10	3.72	3.12	3.70
P, Phosphorus (wt.%)	0.022	0.021	0.023	0.021	0.022

SI unit equivalents: ppm (parts per million; 1 x 10^{-6}) ≡ mg/kg; wt.% (weight per cent) ≡ % (mass fraction).

Note: intervals may appear asymmetric due to rounding;

IND = indeterminate (due to limited reading resolution of the methods employed. For practical purposes the 95% Expanded Uncertainty can be set between zero and a two times multiple of the upper bound/non-detect limit value).

Table 2 continued.

Constituent	Certified Value	95% Expanded Uncertainty		95% Tolerance Limits	
		Low	High	Low	High
Aqua Regia Digestion continued					
Pb, Lead (ppm)	128	123	132	125	130
Pr, Praseodymium (ppm)	6.49	5.83	7.14	6.11	6.86
Rb, Rubidium (ppm)	13.3	12.5	14.2	12.8	13.8
Re, Rhenium (ppm)	< 0.05	IND	IND	IND	IND
S, Sulphur (wt.%)	0.251	0.241	0.261	0.244	0.258
Sb, Antimony (ppm)	4.84	4.50	5.18	4.61	5.07
Sc, Scandium (ppm)	1.52	1.31	1.72	1.42	1.61
Se, Selenium (ppm)	1.28	1.06	1.51	IND	IND
Sm, Samarium (ppm)	4.62	4.07	5.17	4.36	4.87
Sn, Tin (ppm)	1.15	1.03	1.27	1.09	1.20
Sr, Strontium (ppm)	24.8	23.8	25.9	24.1	25.5
Ta, Tantalum (ppm)	< 0.01	IND	IND	IND	IND
Tb, Terbium (ppm)	0.46	0.41	0.52	0.44	0.49
Te, Tellurium (ppm)	0.80	0.71	0.89	0.75	0.85
Th, Thorium (ppm)	10.8	10.3	11.4	10.5	11.2
Ti, Titanium (wt.%)	0.021	0.019	0.023	0.020	0.022
Tl, Thallium (ppm)	0.25	0.23	0.27	0.24	0.26
U, Uranium (ppm)	2.76	2.56	2.96	2.67	2.85
V, Vanadium (ppm)	1.97	1.88	2.06	1.86	2.07
W, Tungsten (ppm)	0.52	0.47	0.57	0.48	0.56
Y, Yttrium (ppm)	7.75	7.35	8.16	7.50	8.00
Yb, Ytterbium (ppm)	0.23	0.19	0.27	0.21	0.25
Zn, Zinc (ppm)	282	274	291	276	289
Zr, Zirconium (ppm)	43.0	39.8	46.2	41.5	44.4

SI unit equivalents: ppm (parts per million; 1 x 10^{-6}) ≡ mg/kg; wt.% (weight per cent) ≡ % (mass fraction).
Note: intervals may appear asymmetric due to rounding;

IND = indeterminate (due to limited reading resolution of the methods employed. For practical purposes the 95% Expanded Uncertainty can be set between zero and a two times multiple of the upper bound/non-detect limit value).

Table 3. Indicative Values for OREAS 606b.

Constituent	Unit	Value	Constituent	Unit	Value	Constituent	Unit	Value
Infrared Combustion								
C	wt.%	*0.169*	C-(Inorganic)	wt.%	*0.127*	C-(Organic)	wt.%	*0.043*
Alkaline Leach								
S-(Sulphate)	wt.%	*0.245*	S-(Sulphide)	wt.%	*0.208*			
Acid Leach								
S-(Sulphate)	wt.%	*0.158*						
4-Acid Digestion								
B	ppm	*14.2*	Hg	ppm	*0.023*			
Aqua Regia Digestion								
Pd	ppb	*18.3*	Pt	ppb	*< 5*	Tm	ppm	*0.047*
Borate Fusion XRF								
Al_2O_3	wt.%	*14.86*	Fe_2O_3	wt.%	*3.41*	S	wt.%	*0.473*
As	ppm	*80*	K_2O	wt.%	*3.69*	SiO_2	wt.%	*69.24*
BaO	ppm	*3439*	MgO	wt.%	*0.240*	Sn	ppm	*55*
CaO	wt.%	*1.54*	MnO	wt.%	*0.034*	Sr	ppm	*200*
Cl	ppm	*20.0*	Na_2O	wt.%	*3.67*	TiO_2	wt.%	*0.203*
Co	ppm	*< 10*	Ni	ppm	*< 10*	V_2O_5	ppm	*17.9*
Cr_2O_3	ppm	*36.5*	P_2O_5	wt.%	*0.066*	Zn	ppm	*320*
Cu	wt.%	*0.036*	Pb	ppm	*180*	Zr	ppm	*270*
Thermogravimetry								
LOI^{1000}	wt.%	*1.98*						
Laser Ablation ICP-MS								
Ag	ppm	*2.15*	Hf	ppm	*7.67*	Sm	ppm	*7.33*
As	ppm	*101*	Ho	ppm	*0.52*	Sn	ppm	*4.50*
Ba	ppm	*2890*	In	ppm	*0.13*	Sr	ppm	*195*
Be	ppm	*3.30*	La	ppm	*44.9*	Ta	ppm	*1.40*
Bi	ppm	*6.15*	Lu	ppm	*0.070*	Tb	ppm	*0.75*
Cd	ppm	*1.60*	Mn	wt.%	*0.029*	Te	ppm	*1.00*
Ce	ppm	*87*	Mo	ppm	*4.30*	Th	ppm	*14.3*
Co	ppm	*2.80*	Nb	ppm	*18.3*	Ti	wt.%	*0.119*
Cr	ppm	*18.0*	Nd	ppm	*38.4*	Tl	ppm	*1.20*
Cs	ppm	*6.16*	Ni	ppm	*5.00*	Tm	ppm	*0.12*
Cu	wt.%	*0.035*	Pb	ppm	*189*	U	ppm	*5.76*
Dy	ppm	*3.48*	Pr	ppm	*10.3*	V	ppm	*6.15*
Er	ppm	*0.99*	Rb	ppm	*137*	W	ppm	*1.50*
Eu	ppm	*1.25*	Re	ppm	*0.008*	Y	ppm	*14.5*
Ga	ppm	*23.3*	Sb	ppm	*7.70*	Yb	ppm	*0.58*
Gd	ppm	*5.61*	Sc	ppm	*3.75*	Zn	ppm	*373*
Ge	ppm	*1.53*	Se	ppm	*< 5*	Zr	ppm	*275*
3-Acid Digestion (no HF)								
Ag	ppm	*1.79*	Gd	ppm	*5.60*	S	wt.%	*0.468*
Al_2O_3	wt.%	*14.28*	Hf	ppm	*6.38*	Sc	ppm	*4.05*
Ba	ppm	*2797*	Ho	ppm	*0.43*	Sm	ppm	*6.97*
Be	ppm	*3.00*	K_2O	wt.%	*3.56*	Sn	ppm	*3.72*

SI unit equivalents: ppb (parts per billion; 1 x 10^{-9}) ≡ µg/kg; ppm (parts per million; 1 x 10^{-6}) ≡ mg/kg; wt.% (weight per cent) ≡ % (mass fraction).

Note: the number of significant figures reported is not a reflection of the level of certainty of stated values. They are instead an artefact of ORE's in-house CRM-specific LIMS.

Table 3 continued.

Constituent	Unit	Value	Constituent	Unit	Value	Constituent	Unit	Value
3-Acid Digestion (no HF) continued								
Bi	ppm	*4.90*	La	ppm	*45.3*	Sr	ppm	*203*
CaO	wt.%	*1.55*	Li	ppm	*31.5*	Ta	ppm	*1.31*
Cd	ppm	*1.25*	MgO	wt.%	*0.213*	Tb	ppm	*0.68*
Ce	ppm	*83*	MnO	wt.%	*0.035*	Th	ppm	*13.2*
Co	ppm	*2.14*	Mo	ppm	*4.40*	TiO_2	wt.%	*0.188*
Cr	ppm	*14.7*	Na_2O	wt.%	*3.57*	U	ppm	*5.54*
Cs	ppm	*6.00*	Nb	ppm	*17.6*	V	ppm	*4.98*
Cu	wt.%	*0.033*	Nd	ppm	*36.6*	W	ppm	*2.02*
Dy	ppm	*3.43*	Ni	ppm	*4.12*	Y	ppm	*13.9*
Er	ppm	*0.97*	P_2O_5	wt.%	*0.071*	Yb	ppm	*0.56*
Eu	ppm	*1.83*	Pb	ppm	*156*	Zn	ppm	*324*
Fe_2O_3	wt.%	*3.16*	Pr	ppm	*10.0*	Zr	ppm	*287*
Ga	ppm	*21.4*	Rb	ppm	*134*			

SI unit equivalents: ppm (parts per million; 1 x 10^{-6}) ≡ mg/kg; wt.% (weight per cent) ≡ % (mass fraction).

Note: the number of significant figures reported is not a reflection of the level of certainty of stated values. They are instead an artefact of ORE’s in-house CRM-specific LIMS.

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INTRODUCTION

OREAS reference materials are intended to provide a low-cost method of evaluating and improving the quality of analysis of geological samples. To the geologist they provide a means of implementing quality control in analytical data sets generated in exploration from the grass roots level through to prospect evaluation, and in grade control at mining operations. To the analyst they provide an effective means of calibrating analytical equipment, assessing new techniques and routinely monitoring in-house procedures. OREAS reference materials enable users to successfully achieve process control of these tasks because the observed variance from repeated analysis has its origin almost exclusively in the analytical process rather than the reference material itself. In evaluating laboratory performance with this CRM, the section headed ‘Instructions for correct use’ should be read carefully.

Table 1 (all laboratories accredited to ISO 17025) and Table 2 (most laboratories accredited to ISO 17025) provide the certified values and their associated 95% expanded uncertainty and tolerance intervals, Table 3 shows indicative values including major and trace element characterisation, Table 4 provides some indicative physical properties and Table 5 provides indicative mineralogy based on semi-quantitative XRD analysis. Table 6 shows gold homogeneity via INAA with a nested ANOVA (see 'Homogeneity Evaluation' section), while Table 7 presents the performance gate intervals for all certified values.

Tabulated results of all analytes together with uncorrected means, medians, standard deviations, relative standard deviations and per cent deviation of lab means from the corrected mean of means (PDM[3]) are presented in the detailed certification data for this CRM (**OREAS 606b-DataPack.1.0.240517_170242.xlsx**). Results are also presented in scatter plots for gold by fire assay, copper by 4-acid digestion and silver by 4-acid digestion (Figures 1 to 3, respectively) together with ±3SD (magenta) and ±5% (yellow) control lines and certified value (green line). Accepted individual results are coloured blue and individual and dataset outliers are identified in red and violet, respectively.

SOURCE MATERIAL

OREAS 606b was prepared from a blend of gold-copper-silver ores from Evolution Mining’s Mount Carlton Operation in Queensland, Australia and argillic rhyodacite sourced from a quarry east of Melbourne, Australia. The mineralisation assemblage at Mount Carlton consists of pyrite, enargite/tennantite, tetrahedrite, digenite, covellite, sphalerite, galena, alunite, dickite, kaolinite and vuggy silica, hosted in advanced argillic altered rhyodacite containing sulphur-salts.

COMMINUTION AND HOMOGENISATION PROCEDURES

The material constituting OREAS 606b was prepared in the following manner:

- Drying of ore materials to constant mass at 85° C;
- Drying of barren rhyodacite to constant mass at 105° C;
- Crushing and milling of ore materials to 100% minus 30 microns;
- Crushing and milling of barren rhyodacite to 98% minus 75 microns;
- Blending in appropriate proportions to achieve the desired grades;
- Homogenisation using OREAS’ novel processing technologies;

Packaging in 10g and 60g units in laminated foil pouches and 500g units in plastic jars.

PHYSICAL PROPERTIES

OREAS 606b was tested at ORE Research & Exploration Pty Ltd's onsite facility for various physical properties. Table 4 presents these findings that should be used for informational purposes only.

Table 4. Physical properties of OREAS 606b.

Bulk Density (kg/m^3)	Moisture (wt.%)	Munsell Notation‡	Munsell Color‡
795	0.20	N8	Very Light Gray

‡The Munsell Rock Color Chart helps geologists and archeologists communicate with colour more effectively by cross-referencing ISCC-NBS colour names with unique Munsell alpha-numeric colour notations for rock colour samples.

MINERALOGY

The semi-quantitative XRD results shown in Table 5 below were undertaken by ALS Metallurgy in Balcatta, Western Australia. The results have been normalised to 100 per cent and represent the relative proportion of crystalline material. Totals greater or less than 100 per cent are due to rounding errors. The Kandite group appears to be mainly kaolinite and dickite. Chalcopyrite and calcite are reported together due to their overlapping patterns. Samples might contain some illite which is reported under muscovite. Some amorphous material might be present in the samples.

Table 5. Indicative mineralogy of OREAS 606b based on semi-quantitative XRD analysis.

Mineral / Mineral Group	% (mass ratio)
Kandite group	2
Chlorite	5
Annite - biotite - phlogopite	1
Muscovite	4
Plagioclase	18
K-feldspar	9
Quartz	54
Magnetite	3
Pyrite	1
Chalcopyrite and/or calcite	1
Alunite	3

ANALYTICAL PROGRAM

Twenty-eight commercial analytical laboratories participated in the program to certify the elements reported in Table 1. The following methods were employed:

- Gold by fire assay using a 25-50g charge weight with AAS finish (22 laboratories) and ICP-OES (6 laboratories);
- Gold by aqua regia digestion using a 10-50g sample mass with ICP-MS finish (12 laboratories), AAS (10 laboratories) finish and ICP-OES (1 laboratory);
- Sulphur by infrared combustion furnace (25 laboratories);

- Full ICP-OES and MS elemental suites by 4-acid digestion (up to 26 laboratories depending on the element);
- Full ICP-OES and MS elemental suites by aqua regia digestion (up to 26 laboratories depending on the element).

The following Sulphur "species" were also requested from the laboratories offering this methodology:

- Sulphate S by Na_2CO_3 leach of sulphates, precipitation as barium sulphate with gravimetric finish (or by difference using the Total S value minus the Sulphide S);
- Sulphide S by Na_2CO_3 leach of sulphates followed by infrared combustion furnace (or by difference using the Total S value minus the Sulphate S).

In addition, instrumental neutron activation analysis (INAA) of Au on 20 x 85mg subsamples was undertaken at ANSTO, Lucas Heights to confirm homogeneity (see Table 6).

Table 3 above shows indicative values including major and trace element characterisation based on two samples analysed at Bureau Veritas in Perth, Western Australia which includes:

- Major oxides by lithium borate fusion with X-ray fluorescence;
- LOI at 1000°C by thermogravimetric analyser;
- Trace elements by laser ablation (on the fused bead) with ICP-MS finish.

Table 3 also includes indicative values for the Sulphur "species" and other elements where interlaboratory consensus or the quantity of data was insufficient for certification.

For the round robin program twelve 3kg test units (lots) were taken at predetermined intervals during the bagging stage, immediately following homogenisation and are considered representative of the entire prepared batch. Six 100g pulp samples were submitted to each laboratory for analysis and were selected systematically to maximise representation. For example, from the 12 sampling lot intervals, the six samples a laboratory may receive could be from the odd lots, or the even lots. I.e., 1, 3, 5, 7, 9 and 11 or 2, 4, 6, 8, 10 and 12). All samples lots within a laboratory sample set were randomised prior to assigning sample numbers.

The 20 individual INAA results upon which much of the homogeneity evaluation is based, included paired 10g samples taken from 10 different sampling lot intervals. This format enabled a nested ANOVA treatment of the INAA results to evaluate homogeneity (see 'Homogeneity Evaluation' section below).

STATISTICAL ANALYSIS

Certified Values and their uncertainty intervals (Tables 1 and 2) have been determined for each analyte following removal of individual, laboratory dataset (batch) and 3SD outliers (single iteration).

For individual outliers within a laboratory batch the z-score test is used in combination with a second method that determines the per cent deviation of the individual value from the batch median. Outliers in general are selected on the basis of z-scores > 2.5 and with per cent deviations (i) > 3 and (ii) more than three times the average absolute per cent deviation for the batch. Each laboratory data set mean is tested for outlying status based on z-score discrimination and rejected if > 2.5. After individual and laboratory data set (batch) outliers have been eliminated a non-iterative 3 standard deviation filter is applied, with those values lying outside this window also relegated to outlying status. However, while statistics are taken into account, the exercise of a statistician's prerogative plays a significant role in identifying outliers.

Certified Values are the means of accepted laboratory means after outlier filtering and are the present best estimate of the true value. The INAA data (see Table 6) is omitted from determination of the certified value for Au and is used solely for the calculation of Tolerance Limits and homogeneity evaluation.

95% Expanded Uncertainty provides a 95% probability that the true value of the analyte under consideration lies between the upper and lower limits and is calculated according to the method outlined in ISO 98-3:2008 [6, 16]. All known or suspected sources of bias have been investigated or taken into account.

Indicative (uncertified) values (Table 3) are present where the number of laboratories reporting a particular analyte is insufficient (< 5) to support certification or where interlaboratory consensus is poor. This data is intended for ‘informational purposes’ only.

Homogeneity Evaluation

The tolerance limits (ISO 16269:2014) shown in Tables 1 and 2 were determined using an analysis of precision errors method and are considered a conservative estimate of true homogeneity. The meaning of tolerance limits may be illustrated for copper by 4-acid digestion, where 99% of the time (1-α=0.99) at least 95% of subsamples (ρ=0.95) will have concentrations lying between 0.033 and 0.034 wt.%. Put more precisely, this means that if the same number of subsamples were taken and analysed in the same manner repeatedly, 99% of the tolerance intervals so constructed would cover at least 95% of the total population, and 1% of the tolerance intervals would cover less than 95% of the total population (ISO Guide 35). ***Please note that tolerance limits pertain to the homogeneity of the CRM only and should not be used as control limits for laboratory performance.***

The homogeneity of gold has been determined by INAA at ANSTO using the reduced analytical subsample method which utilises the known relationship between standard deviation and analytical subsample weight (Ingamells and Switzer, 1973 [2]). In this approach the sample aliquot is substantially reduced to a point where most of the variability in replicate assays should be due to inhomogeneity of the reference material and measurement error becomes negligible. Table 6 below shows the gold INAA data determined on 20 x 85mg subsamples of OREAS 606b. An equivalent scaled version of the results is also provided to demonstrate an appreciation of what this data means if 30g fire assays were undertaken without the normal measurement error associated with this methodology. In this instance, the 1RSD of 0.15% calculated for a 30g fire assay sample (2.90% at 85mg weights) confirms the high level of gold homogeneity in OREAS 606b.

Table 6. Neutron Activation Analysis of Au (in ppm) on 20 x 85mg subsamples and showing the equivalent results scaled to a 30g sample mass typical of fire assay determination.

Replicate No	Au 85mg actual	Au 30g equivalent*
1	0.412	0.408
2	0.413	0.408
3	0.397	0.407
4	0.398	0.407
5	0.403	0.408
6	0.403	0.408
7	0.394	0.407
8	0.431	0.409
9	0.397	0.407
10	0.428	0.409
11	0.401	0.407
12	0.400	0.407
13	0.411	0.408
14	0.410	0.408
15	0.413	0.408
16	0.415	0.408
17	0.405	0.408
18	0.395	0.407
19	0.398	0.407
20	0.433	0.409
Mean	0.408	0.408
Median	0.404	0.408
Std Dev.	0.012	0.001
Rel.Std.Dev.	**2.90%**	**0.15%**

*Results calculated for a 30g equivalent sample mass using the formula: $x^{30g\ Eq} = \frac{(x^{INAA} - \bar{X}) \times RSD@30g}{RSD@85mg} + \bar{X}$

where $x^{30g\ Eq}$ = equivalent result calculated for a 30g sample mass
(x^{INAA}) = raw INAA result at 85mg
$\bar{X}$ = mean of 85mg INAA results

The homogeneity of OREAS 606b has also been evaluated in an Analysis of Variance (**ANOVA**) of the INAA data. The 20 samples were comprised of paired samples from each of 10 different sampling lot intervals (representative of the prepared batch) and were randomised prior to assigning sample numbers. The duplicate samples enabled an ANOVA by comparison of within- and between-unit variances across the 10 pairs. The purpose of the ANOVA is to test that no statistically significant difference exists in the variance between units to that of the variance within units. This allows an assessment of homogeneity across the entire prepared batch of OREAS 606b. The test was performed using the following parameters:

- Gold INAA – 20 results (1 laboratory providing duplicate analyses on 10 samples where each sample can be viewed as a 'unit');
- Null Hypothesis, H_0: Between-unit variance is no greater than within-unit variance (reject H_0 if p-value < 0.05);
- Alternative Hypothesis, H_1: Between-unit variance is greater than within-unit variance.

The data was not filtered for outliers prior to the calculation of the *p*-value. This process derived a *p*-value of 0.50, a statistically insignificant result so the Null Hypothesis is accepted.

It is important to note that ANOVA is not an absolute measure of homogeneity. Rather, it establishes whether or not the analytes are distributed in a similar manner throughout the packaging run of OREAS 606b and whether the variance between two subsamples from the same unit is statistically distinguishable from the variance of two subsamples taken from any two separate units. A reference material therefore can possess poor absolute homogeneity yet still pass a relative homogeneity (ANOVA) test if the within-unit heterogeneity is large and similar across all units. Based on the statistical analysis of ANOVA and the results of the interlaboratory certification program, it can be concluded that OREAS 606b is fit-for-purpose as a certified reference material (see 'Intended Use' below).

PERFORMANCE GATES

The standard deviations (SD's) intervals reported in Table 7 provide an indication of a level of performance that might reasonably be expected from a laboratory being monitored by this CRM in a QA/QC program. They take into account errors attributable to measurement uncertainty and CRM variability. For an effective CRM the contribution of the latter should be negligible in comparison to measurement errors. The Standard Deviation values include all sources of measurement uncertainty: between-lab variance, within-run variance (precision errors) and CRM variability.

In the application of SD's in monitoring performance it is important to note that not all laboratories function at the same level of proficiency and that different methods in use at a particular laboratory have differing levels of precision. Each laboratory has its own inherent SD (for a specific concentration level and analyte-method pair) based on the analytical process and this SD is not directly related to the round robin program (see 'Intended Use' section for more detail). The SD for each analyte's certified value is calculated from the same filtered data set used to determine the certified value, i.e., after removal of all individual, lab dataset (batch) and 3SD outliers (single iteration). These outliers can only be removed after the absolute homogeneity of the CRM has been independently established, i.e., the outliers must be confidently deemed to be analytical rather than arising from inhomogeneity of the CRM. ***The standard deviation is then calculated for each analyte from the pooled accepted analyses generated from the certification program.***

Table 7 below shows intervals calculated for two and three standard deviations. As a guide these intervals may be regarded as warning or rejection for multiple 2SD outliers, or rejection for individual 3SD outliers in QC monitoring, although their precise application should be at the discretion of the QC manager concerned (also see 'Intended Use' section below). Westgard Rules extend the basics of single-rule QC monitoring using multi-rules (for more information visit www.westgard.com/mltirule.htm). A second method utilises a 5% window calculated directly from the certified value.

Standard deviation is also shown in relative percent for one, two and three relative standard deviations (1RSD, 2RSD and 3RSD) to facilitate an appreciation of the magnitude of these numbers and a comparison with the 5% window. Caution should be exercised when concentration levels approach lower limits of detection of the analytical methods employed as performance gates calculated from standard deviations tend to be excessively wide whereas those determined by the 5% method are too narrow. One approach used at commercial laboratories is to set the acceptance criteria at twice the detection level (DL) ± 10%.

i.e., Certified Value ± 10% ± 2DL [1].

Table 7. Performance Gates for OREAS 606b.

Constituent	Certified Value	Absolute Standard Deviations					Relative Standard Deviations			5% window	
		1SD	2SD Low	2SD High	3SD Low	3SD High	1RSD	2RSD	3RSD	Low	High
Pb Fire Assay											
Au, ppm	0.386	0.015	0.356	0.416	0.341	0.431	3.89%	7.78%	11.68%	0.366	0.405
Aqua Regia Digestion (sample weights 10-50g)											
Au, ppm	0.375	0.032	0.310	0.440	0.278	0.472	8.62%	17.24%	25.86%	0.356	0.394
Infrared Combustion											
S, wt.%	0.460	0.018	0.425	0.495	0.407	0.513	3.83%	7.65%	11.48%	0.437	0.483
4-Acid Digestion											
Ag, ppm	1.53	0.088	1.35	1.70	1.26	1.79	5.76%	11.53%	17.29%	1.45	1.60
Al, wt.%	7.54	0.307	6.92	8.15	6.62	8.46	4.07%	8.14%	12.21%	7.16	7.91
As, ppm	89	5.1	79	100	74	105	5.70%	11.40%	17.09%	85	94
Ba, ppm	2892	128	2637	3147	2509	3274	4.41%	8.82%	13.23%	2747	3036
Be, ppm	2.98	0.156	2.67	3.29	2.51	3.45	5.23%	10.47%	15.70%	2.83	3.13
Bi, ppm	4.83	0.243	4.34	5.31	4.10	5.56	5.03%	10.07%	15.10%	4.59	5.07
Ca, wt.%	1.10	0.031	1.04	1.16	1.01	1.19	2.80%	5.60%	8.39%	1.05	1.16
Cd, ppm	1.24	0.098	1.05	1.44	0.95	1.53	7.87%	15.73%	23.60%	1.18	1.30
Ce, ppm	89	5.3	78	99	73	105	5.98%	11.96%	17.94%	84	93
Co, ppm	2.25	0.171	1.91	2.60	1.74	2.77	7.61%	15.22%	22.83%	2.14	2.37
Cr, ppm	14.7	1.7	11.4	18.0	9.7	19.7	11.26%	22.52%	33.78%	14.0	15.5
Cs, ppm	6.11	0.406	5.30	6.92	4.89	7.33	6.65%	13.30%	19.95%	5.81	6.42
Cu, wt.%	0.033	0.001	0.031	0.036	0.029	0.038	4.26%	8.53%	12.79%	0.032	0.035
Dy, ppm	3.55	0.118	3.32	3.79	3.20	3.91	3.31%	6.62%	9.93%	3.38	3.73
Er, ppm	1.01	0.070	0.87	1.15	0.80	1.22	6.91%	13.81%	20.72%	0.96	1.06
Eu, ppm	1.40	0.137	1.12	1.67	0.99	1.81	9.77%	19.54%	29.31%	1.33	1.47
Fe, wt.%	2.33	0.086	2.15	2.50	2.07	2.59	3.71%	7.42%	11.13%	2.21	2.44
Ga, ppm	23.0	1.41	20.2	25.9	18.8	27.3	6.10%	12.20%	18.30%	21.9	24.2
Gd, ppm	5.60	0.405	4.79	6.41	4.39	6.82	7.22%	14.45%	21.67%	5.32	5.88
Ge, ppm	0.19	0.08	0.03	0.35	0.00	0.43	41.07%	82.14%	123.2%	0.18	0.20
Hf, ppm	6.48	0.284	5.91	7.05	5.63	7.33	4.38%	8.77%	13.15%	6.16	6.81
Ho, ppm	0.48	0.043	0.40	0.57	0.35	0.61	8.89%	17.78%	26.66%	0.46	0.51
In, ppm	0.14	0.011	0.12	0.16	0.11	0.17	7.66%	15.32%	22.97%	0.14	0.15
K, wt.%	3.02	0.154	2.71	3.33	2.55	3.48	5.12%	10.24%	15.36%	2.87	3.17
La, ppm	45.0	2.95	39.1	50.9	36.1	53.8	6.56%	13.12%	19.68%	42.7	47.2
Li, ppm	32.1	1.45	29.2	35.0	27.8	36.5	4.51%	9.01%	13.52%	30.5	33.7
Lu, ppm	0.074	0.015	0.044	0.104	0.029	0.119	20.30%	40.59%	60.89%	0.070	0.078
Mg, wt.%	0.127	0.006	0.114	0.139	0.108	0.146	5.03%	10.07%	15.10%	0.120	0.133
Mn, wt.%	0.028	0.001	0.026	0.031	0.024	0.032	4.47%	8.94%	13.40%	0.027	0.029
Mo, ppm	4.38	0.182	4.02	4.75	3.84	4.93	4.15%	8.30%	12.45%	4.16	4.60
Na, wt.%	2.67	0.125	2.42	2.92	2.30	3.05	4.68%	9.36%	14.04%	2.54	2.81
Nb, ppm	18.5	0.99	16.5	20.5	15.5	21.5	5.34%	10.67%	16.01%	17.6	19.4
Nd, ppm	37.4	3.22	31.0	43.9	27.8	47.1	8.62%	17.23%	25.85%	35.6	39.3
Ni, ppm	3.56	0.289	2.98	4.14	2.69	4.43	8.11%	16.22%	24.33%	3.38	3.74
P, wt.%	0.030	0.001	0.028	0.032	0.027	0.033	3.30%	6.59%	9.89%	0.028	0.031
Pb, ppm	157	7	142	172	135	179	4.75%	9.49%	14.24%	149	165
Pr, ppm	10.1	0.64	8.8	11.3	8.2	12.0	6.33%	12.65%	18.98%	9.6	10.6

SI unit equivalents: ppm (parts per million; 1 x 10^{-6}) ≡ mg/kg; wt.% (weight per cent) ≡ % (mass fraction).

Note 1: intervals may appear asymmetric due to rounding.

Note 2: the number of decimal places quoted does not imply accuracy of the certified value to this level but are given to minimise rounding errors when calculating 2SD and 3SD windows.

Table 7 continued.

Constituent	Certified Value	Absolute Standard Deviations					Relative Standard Deviations			5% window	
		1SD	2SD Low	2SD High	3SD Low	3SD High	1RSD	2RSD	3RSD	Low	High
4-Acid Digestion continued											
Rb, ppm	139	10	120	159	110	168	7.03%	14.06%	21.09%	132	146
Re, ppm	< 0.002	IND	IND	IND	IND	IND	IND	IND	IND	IND	IND
S, wt.%	0.468	0.020	0.428	0.509	0.408	0.529	4.30%	8.61%	12.91%	0.445	0.492
Sb, ppm	6.38	0.398	5.58	7.17	5.18	7.57	6.24%	12.48%	18.72%	6.06	6.70
Sc, ppm	4.26	0.43	3.39	5.13	2.96	5.56	10.17%	20.34%	30.51%	4.05	4.47
Se, ppm	1.80	0.68	0.44	3.16	0.00	3.83	37.76%	75.51%	113.3%	1.71	1.89
Sm, ppm	7.27	0.434	6.40	8.13	5.97	8.57	5.97%	11.93%	17.90%	6.90	7.63
Sn, ppm	3.72	0.163	3.39	4.04	3.23	4.20	4.38%	8.75%	13.13%	3.53	3.90
Sr, ppm	200	10	180	219	170	229	5.00%	9.99%	14.99%	190	210
Ta, ppm	1.36	0.087	1.19	1.54	1.10	1.63	6.38%	12.76%	19.14%	1.30	1.43
Tb, ppm	0.73	0.050	0.63	0.83	0.58	0.88	6.80%	13.59%	20.39%	0.69	0.77
Te, ppm	0.84	0.075	0.69	0.98	0.61	1.06	8.94%	17.88%	26.81%	0.79	0.88
Th, ppm	14.8	0.71	13.4	16.2	12.7	16.9	4.83%	9.65%	14.48%	14.1	15.5
Ti, wt.%	0.116	0.004	0.108	0.124	0.104	0.128	3.54%	7.07%	10.61%	0.110	0.122
Tl, ppm	0.89	0.043	0.80	0.97	0.76	1.02	4.90%	9.79%	14.69%	0.84	0.93
Tm, ppm	0.10	0.007	0.09	0.12	0.08	0.13	6.83%	13.67%	20.50%	0.10	0.11
U, ppm	5.57	0.291	4.98	6.15	4.69	6.44	5.23%	10.46%	15.70%	5.29	5.84
V, ppm	4.97	0.61	3.75	6.18	3.14	6.79	12.27%	24.54%	36.81%	4.72	5.21
W, ppm	1.85	0.149	1.55	2.15	1.40	2.30	8.05%	16.10%	24.14%	1.76	1.94
Y, ppm	14.8	0.71	13.4	16.3	12.7	17.0	4.76%	9.53%	14.29%	14.1	15.6
Yb, ppm	0.61	0.06	0.48	0.74	0.42	0.80	10.46%	20.92%	31.38%	0.58	0.64
Zn, ppm	316	16	285	347	269	362	4.91%	9.82%	14.73%	300	332
Aqua Regia Digestion											
Ag, ppm	1.49	0.113	1.26	1.71	1.15	1.82	7.60%	15.20%	22.80%	1.41	1.56
Al, wt.%	0.804	0.049	0.705	0.902	0.656	0.951	6.11%	12.23%	18.34%	0.763	0.844
As, ppm	86	4.5	77	95	72	99	5.30%	10.60%	15.90%	82	90
B, ppm	< 10	IND	IND	IND	IND	IND	IND	IND	IND	IND	IND
Ba, ppm	212	20	173	251	153	271	9.26%	18.52%	27.78%	201	222
Be, ppm	0.51	0.042	0.43	0.60	0.39	0.64	8.12%	16.25%	24.37%	0.49	0.54
Bi, ppm	4.96	0.326	4.31	5.61	3.98	5.94	6.58%	13.15%	19.73%	4.71	5.21
Ca, wt.%	0.632	0.027	0.579	0.685	0.552	0.712	4.20%	8.41%	12.61%	0.601	0.664
Cd, ppm	1.19	0.069	1.06	1.33	0.99	1.40	5.77%	11.54%	17.31%	1.13	1.25
Ce, ppm	56	4.0	48	64	44	68	7.06%	14.13%	21.19%	53	59
Co, ppm	2.02	0.062	1.90	2.15	1.84	2.21	3.06%	6.11%	9.17%	1.92	2.12
Cr, ppm	16.1	0.83	14.5	17.8	13.6	18.6	5.17%	10.34%	15.51%	15.3	16.9
Cs, ppm	1.48	0.15	1.17	1.79	1.02	1.94	10.40%	20.80%	31.19%	1.40	1.55
Cu, wt.%	0.034	0.001	0.031	0.036	0.030	0.037	3.56%	7.13%	10.69%	0.032	0.035
Dy, ppm	2.08	0.164	1.76	2.41	1.59	2.58	7.89%	15.77%	23.66%	1.98	2.19
Er, ppm	0.51	0.07	0.38	0.64	0.32	0.71	12.73%	25.47%	38.20%	0.49	0.54
Eu, ppm	0.70	0.060	0.58	0.82	0.52	0.88	8.65%	17.30%	25.95%	0.66	0.73
Fe, wt.%	2.00	0.066	1.87	2.13	1.80	2.20	3.29%	6.57%	9.86%	1.90	2.10
Ga, ppm	5.90	0.432	5.04	6.76	4.60	7.19	7.32%	14.64%	21.95%	5.60	6.19
Gd, ppm	3.82	0.236	3.35	4.29	3.11	4.53	6.17%	12.34%	18.52%	3.63	4.01

SI unit equivalents: ppm (parts per million; 1 x 10^{-6}) ≡ mg/kg; wt.% (weight per cent) ≡ % (mass fraction).

Note 1: intervals may appear asymmetric due to rounding; IND = indeterminate.

Note 2: the number of decimal places quoted does not imply accuracy of the certified value to this level but are given to minimise rounding errors when calculating 2SD and 3SD windows.

Table 7 continued.

Constituent	Certified Value	Absolute Standard Deviations					Relative Standard Deviations			5% window	
		1SD	2SD Low	2SD High	3SD Low	3SD High	1RSD	2RSD	3RSD	Low	High
Aqua Regia Digestion continued											
Ge, ppm	0.12	0.04	0.05	0.20	0.01	0.24	30.85%	61.70%	92.56%	0.12	0.13
Hf, ppm	1.22	0.068	1.08	1.35	1.01	1.42	5.60%	11.20%	16.80%	1.16	1.28
Hg, ppm	0.048	0.009	0.030	0.066	0.021	0.075	18.57%	37.14%	55.71%	0.046	0.051
Ho, ppm	0.25	0.017	0.21	0.28	0.19	0.30	7.09%	14.18%	21.28%	0.23	0.26
In, ppm	0.10	0.006	0.09	0.12	0.09	0.12	5.85%	11.70%	17.54%	0.10	0.11
K, wt.%	0.235	0.018	0.199	0.271	0.181	0.289	7.64%	15.29%	22.93%	0.224	0.247
La, ppm	27.9	2.07	23.8	32.0	21.7	34.1	7.44%	14.87%	22.31%	26.5	29.3
Li, ppm	19.9	1.26	17.3	22.4	16.1	23.6	6.34%	12.67%	19.01%	18.9	20.9
Lu, ppm	0.028	0.005	0.018	0.038	0.013	0.043	18.06%	36.11%	54.17%	0.026	0.029
Mg, wt.%	0.088	0.003	0.082	0.095	0.078	0.099	3.85%	7.69%	11.54%	0.084	0.093
Mn, wt.%	0.025	0.001	0.023	0.027	0.022	0.028	3.72%	7.44%	11.16%	0.024	0.026
Mo, ppm	4.00	0.302	3.39	4.60	3.09	4.90	7.55%	15.10%	22.65%	3.80	4.20
Na, wt.%	0.075	0.006	0.063	0.088	0.057	0.094	8.03%	16.05%	24.08%	0.072	0.079
Nb, ppm	0.90	0.19	0.51	1.29	0.32	1.48	21.52%	43.04%	64.57%	0.86	0.95
Nd, ppm	25.5	2.26	21.0	30.1	18.8	32.3	8.85%	17.70%	26.55%	24.3	26.8
Ni, ppm	3.41	0.244	2.92	3.90	2.68	4.14	7.15%	14.30%	21.45%	3.24	3.58
P, wt.%	0.022	0.001	0.020	0.023	0.019	0.024	3.83%	7.67%	11.50%	0.021	0.023
Pb, ppm	128	4	119	136	115	140	3.16%	6.31%	9.47%	121	134
Pr, ppm	6.49	0.290	5.91	7.07	5.61	7.36	4.48%	8.95%	13.43%	6.16	6.81
Rb, ppm	13.3	1.26	10.8	15.9	9.5	17.1	9.46%	18.92%	28.38%	12.7	14.0
Re, ppm	< 0.05	IND	IND	IND	IND	IND	IND	IND	IND	IND	IND
S, wt.%	0.251	0.015	0.222	0.281	0.207	0.296	5.91%	11.82%	17.73%	0.239	0.264
Sb, ppm	4.84	0.49	3.86	5.82	3.38	6.30	10.08%	20.16%	30.24%	4.60	5.08
Sc, ppm	1.52	0.17	1.17	1.87	0.99	2.04	11.50%	22.99%	34.49%	1.44	1.59
Se, ppm	1.28	0.16	0.96	1.61	0.80	1.77	12.50%	24.99%	37.49%	1.22	1.35
Sm, ppm	4.62	0.63	3.37	5.87	2.74	6.49	13.55%	27.10%	40.65%	4.39	4.85
Sn, ppm	1.15	0.13	0.89	1.40	0.77	1.53	11.08%	22.15%	33.23%	1.09	1.20
Sr, ppm	24.8	1.31	22.2	27.5	20.9	28.8	5.27%	10.54%	15.81%	23.6	26.1
Ta, ppm	< 0.01	IND	IND	IND	IND	IND	IND	IND	IND	IND	IND
Tb, ppm	0.46	0.029	0.40	0.52	0.37	0.55	6.25%	12.49%	18.74%	0.44	0.48
Te, ppm	0.80	0.067	0.67	0.93	0.60	1.00	8.36%	16.71%	25.07%	0.76	0.84
Th, ppm	10.8	0.88	9.0	12.6	8.2	13.5	8.16%	16.31%	24.47%	10.3	11.3
Ti, wt.%	0.021	0.002	0.017	0.025	0.014	0.028	10.39%	20.79%	31.18%	0.020	0.022
Tl, ppm	0.25	0.019	0.21	0.29	0.20	0.31	7.44%	14.88%	22.32%	0.24	0.26
U, ppm	2.76	0.203	2.35	3.17	2.15	3.37	7.35%	14.70%	22.05%	2.62	2.90
V, ppm	1.97	0.130	1.71	2.23	1.58	2.36	6.59%	13.18%	19.76%	1.87	2.07
W, ppm	0.52	0.06	0.41	0.63	0.35	0.69	10.73%	21.46%	32.19%	0.50	0.55
Y, ppm	7.75	0.546	6.66	8.84	6.11	9.39	7.05%	14.09%	21.14%	7.36	8.14
Yb, ppm	0.23	0.03	0.18	0.28	0.15	0.30	11.18%	22.35%	33.53%	0.22	0.24
Zn, ppm	282	13	255	309	242	322	4.74%	9.48%	14.23%	268	296
Zr, ppm	43.0	5.5	32.0	53.9	26.5	59.4	12.75%	25.50%	38.25%	40.8	45.1

SI unit equivalents: ppm (parts per million; 1 x 10^{-6}) ≡ mg/kg; wt.% (weight per cent) ≡ % (mass fraction).

Note 1: intervals may appear asymmetric due to rounding; IND = indeterminate.

Note 2: the number of decimal places quoted does not imply accuracy of the certified value to this level but are given to minimise rounding errors when calculating 2SD and 3SD windows.

PARTICIPATING LABORATORIES

1. Actlabs, Ancaster, Ontario, Canada
2. AGAT Laboratories, Calgary, Alberta, Canada
3. Alex Stewart International, Mendoza, Argentina
4. ALS, Johannesburg, South Africa
5. ALS, Lima, Peru
6. ALS, Loughrea, Galway, Ireland
7. ALS, Malaga, WA, Australia
8. ALS, Vancouver, BC, Canada
9. American Assay Laboratories, Sparks, Nevada, USA
10. ANSTO, Lucas Heights, NSW, Australia
11. ARGETEST Mineral Processing, Ankara, Central Anatolia, Turkey
12. Bureau Veritas Commodities Canada Ltd, Vancouver, BC, Canada
13. Bureau Veritas Geoanalytical, Perth, WA, Australia
14. CERTIMIN, Lima, Peru
15. ESAN Istanbul, Istanbul, Turkey
16. Inspectorate (BV), Lima, Peru
17. Intertek, Cupang, Muntinlupa, Philippines
18. Intertek Genalysis, Perth, WA, Australia
19. Intertek Minerals Ltd, Tarkwa, Western Region, Ghana
20. Intertek Testing Services, Townsville, QLD, Australia
21. Koza Gold (Ovacik Gold Mine), Bergama, Izmir, Turkey
22. Paragon Geochemical Laboratories, Sparks, Nevada, USA
23. PT Geoservices Ltd, Cikarang, Jakarta Raya, Indonesia
24. PT Intertek Utama Services, Jakarta Timur, DKI Jakarta, Indonesia
25. Reminex Centre de Recherche, Marrakesh, Marrakesh-Safi, Morocco
26. Saskatchewan Research Council, Saskatoon, Saskatchewan, Canada
27. SGS Australia Mineral Services, Perth, WA, Australia
28. SGS Mineral Services, Townsville, QLD, Australia
29. Shiva Analyticals Ltd, Bangalore North, Karnataka, India
30. Stewart Assay & Environmental Laboratories LLC, Kara-Balta, Chüy, Kyrgyzstan

Please note: To preserve anonymity, the above numbered alphabetical list of participating laboratories does not correspond with the Lab ID numbering on the scatter plots below.

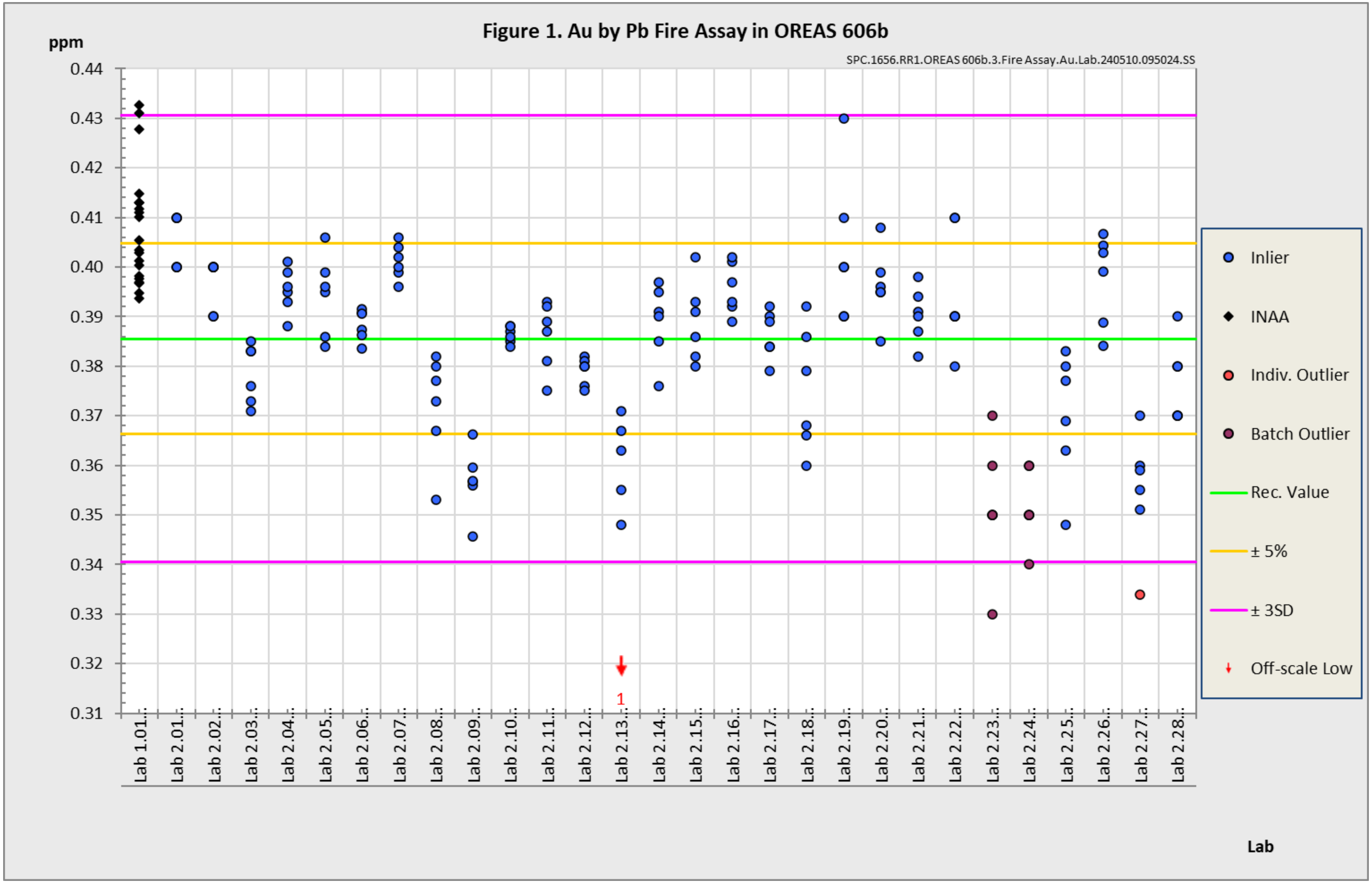
Figure 1. Au by Pb Fire Assay in OREAS 606b
SPC.1656.RR1.OREAS 606b.3.Fire Assay.Au.Lab.240510.095024.SS
ppm
0.44
0.43
0.42
0.41
0.40
0.39
0.38
0.37
0.36
0.35
0.34
0.33
0.32
0.31
Lab 1.01...
Lab 2.01...
Lab 2.02...
Lab 2.03...
Lab 2.04...
Lab 2.05...
Lab 2.06...
Lab 2.07...
Lab 2.08...
Lab 2.09...
Lab 2.10...
Lab 2.11...
Lab 2.12...
Lab 2.13...
Lab 2.14...
Lab 2.15...
Lab 2.16...
Lab 2.17...
Lab 2.18...
Lab 2.19...
Lab 2.20...
Lab 2.21...
Lab 2.22...
Lab 2.23...
Lab 2.24...
Lab 2.25...
Lab 2.26...
Lab 2.27...
Lab 2.28...
1
Lab
Inlier
INAA
Indiv. Outlier
Batch Outlier
Rec. Value
± 5%
± 3SD
Off-scale Low

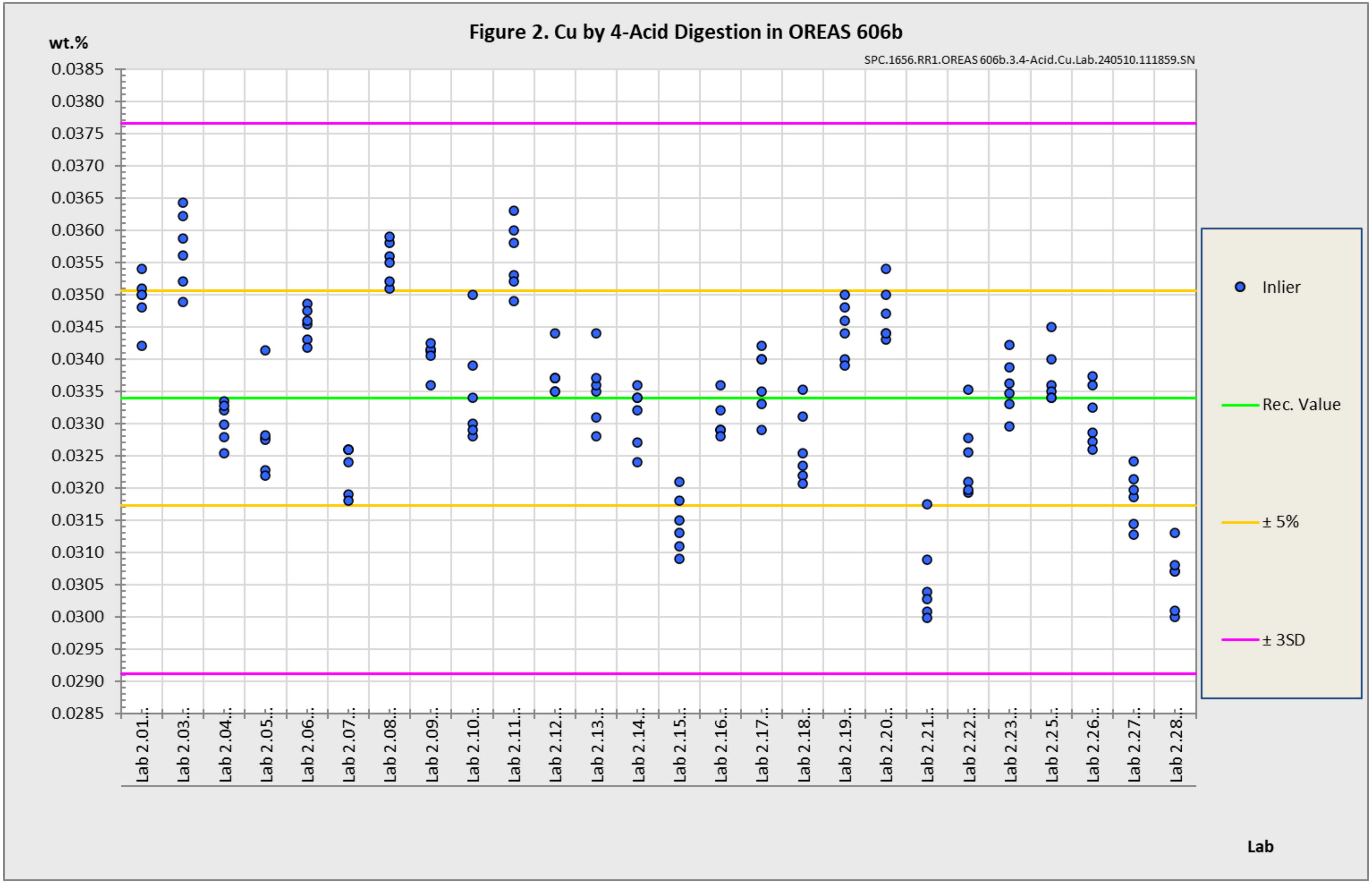
Figure 2. Cu by 4-Acid Digestion in OREAS 606b
wt.%
SPC.1656.RR1.OREAS 606b.3.4-Acid.Cu.Lab.240510.111859.SN
0.0385
0.0380
0.0375
0.0370
0.0365
0.0360
0.0355
0.0350
0.0345
0.0340
0.0335
0.0330
0.0325
0.0320
0.0315
0.0310
0.0305
0.0300
0.0295
0.0290
0.0285
Lab 2.01...
Lab 2.03...
Lab 2.04...
Lab 2.05...
Lab 2.06...
Lab 2.07...
Lab 2.08...
Lab 2.09...
Lab 2.10...
Lab 2.11...
Lab 2.12...
Lab 2.13...
Lab 2.14...
Lab 2.15...
Lab 2.16...
Lab 2.17...
Lab 2.18...
Lab 2.19...
Lab 2.20...
Lab 2.21...
Lab 2.22...
Lab 2.23...
Lab 2.25...
Lab 2.26...
Lab 2.27...
Lab 2.28...
Inlier
Rec. Value
± 5%
± 3SD
Lab

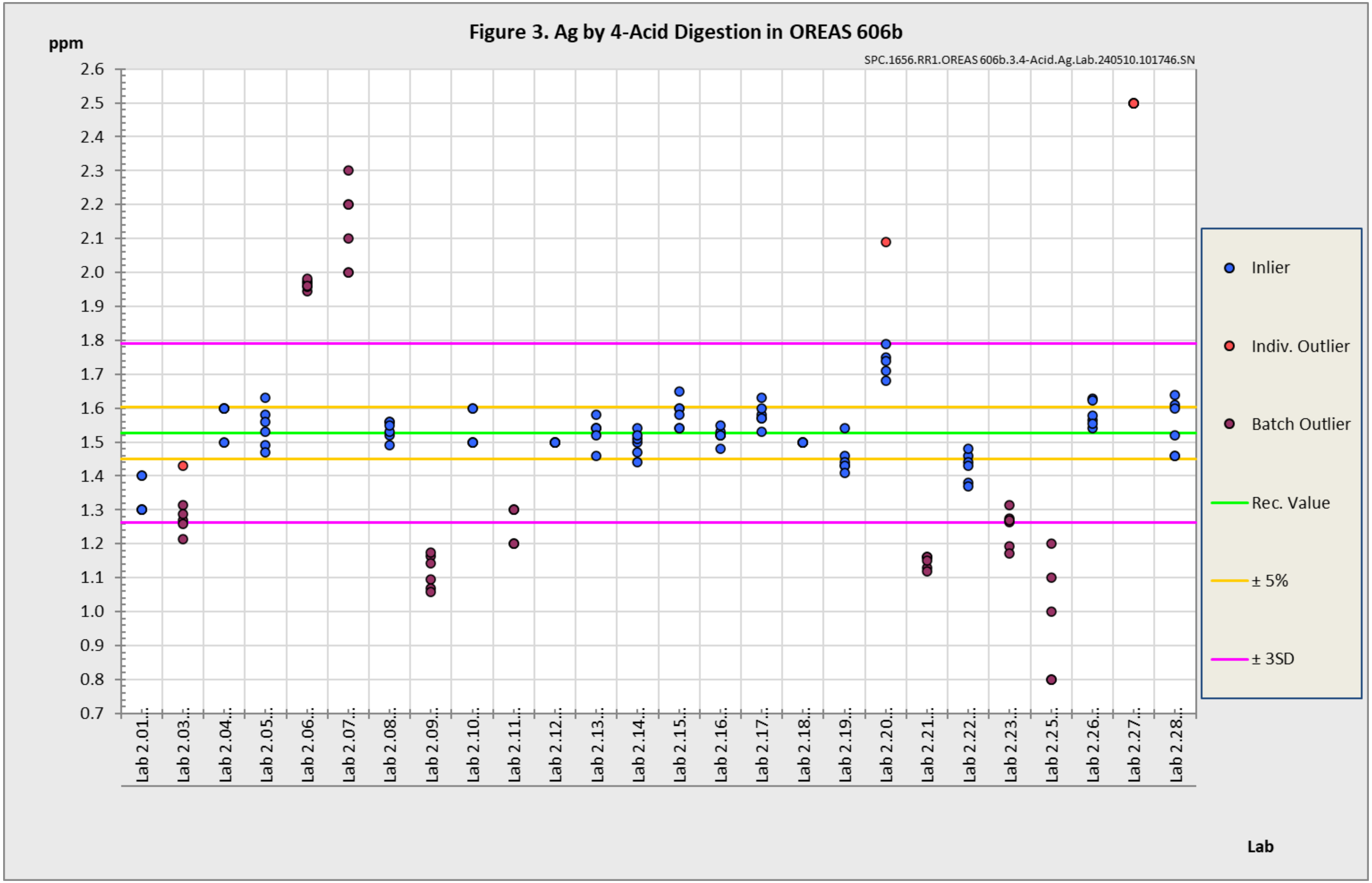
Figure 3. Ag by 4-Acid Digestion in OREAS 606b
SPC.1656.RR1.OREAS 606b.3.4-Acid.Ag.Lab.240510.101746.SN
ppm
2.6
2.5
2.4
2.3
2.2
2.1
2.0
1.9
1.8
1.7
1.6
1.5
1.4
1.3
1.2
1.1
1.0
0.9
0.8
0.7
Lab 2.01...
Lab 2.03...
Lab 2.04...
Lab 2.05...
Lab 2.06...
Lab 2.07...
Lab 2.08...
Lab 2.09...
Lab 2.10...
Lab 2.11...
Lab 2.12...
Lab 2.13...
Lab 2.14...
Lab 2.15...
Lab 2.16...
Lab 2.17...
Lab 2.18...
Lab 2.19...
Lab 2.20...
Lab 2.21...
Lab 2.22...
Lab 2.23...
Lab 2.25...
Lab 2.26...
Lab 2.27...
Lab 2.28...
Lab
Inlier
Indiv. Outlier
Batch Outlier
Rec. Value
± 5%
± 3SD

PREPARER AND SUPPLIER

Certified reference material OREAS 606b is prepared, certified and supplied by:

ORE Research & Exploration Pty Ltd	Tel:	+613-9729 0333
37A Hosie Street	Web:	www.oreas.com
Bayswater North VIC 3153	Email:	info@ore.com.au
AUSTRALIA		

METROLOGICAL TRACEABILITY

The interlaboratory results that underpin the certified values are metrologically traceable to the international measurement scale (SI) of mass (either as a % mass fraction or as milligrams per kilogram (mg/kg)). In line with popular use, all data within tables in this certificate are expressed as the mass fraction in either weight percent (wt.%) or parts per million (ppm).

The analytical samples sent to participating laboratories were selected in a manner to be representative of the entire prepared batch of CRM. This ‘representivity’ was maintained in each submitted laboratory sample batch and ensures the user that the data is traceable from sample selection through to the analytical results. The systematic sampling method was chosen due to the low risk of overlooking repetitive effects or trends in the batch due to the way the CRM was processed. In line with ISO 17025 [9], each analytical data set received from the participating laboratories has been validated by its assayer through the inclusion of internal reference materials and QC checks during and post analysis.

The participating laboratories were chosen on the basis of their competence (from past performance in interlaboratory programs undertaken by ORE Pty Ltd) for a particular analytical method, analyte or analyte suite and sample matrix. These laboratories are accredited to ISO 17025 for Au by fire assay (Table 1). The other operationally defined measurands characterised in this certificate (Table 2) are derived from data procured mostly from ISO 17025 accredited laboratories. The certified values presented in this report are calculated from the means of accepted data following robust technical and statistical analysis as detailed in this report.

Guide ISO/TR 16476:2016, section 5.3.1 describes metrological traceability in reference materials as it pertains to the transformation of the measurand. In this section it states, *“Although the determination of the property value itself can be made traceable to appropriate units through, for example, calibration of the measurement equipment used, steps like the transformation of the sample from one physical (chemical) state to another cannot. Such transformations may only be compared with a reference (when available), or among themselves. For some transformations, reference methods have been defined and may be used in certification projects to evaluate the uncertainty associated with such a transformation.* ***In other cases, only a comparison among different laboratories using the same procedure is possible. In this case, it is impossible to demonstrate absence of method bias; therefore, the result is an operationally defined measurand*** *(ISO Guide 33405:2024-05, 9.2.4c).”* Certification takes place on the basis of agreement among operationally defined, independent measurement results.

COMMUTABILITY

The measurements of the results that underlie the certified values contained in this report were undertaken by methods involving pre-treatment (fusion/digestion) of the sample. This served to reduce the sample to a simple and well understood form permitting calibration using simple solutions of the CRM. Due to these methods being well understood and highly effective, commutability is not an issue for this CRM. All OREAS CRMs are sourced from natural ore minerals meaning they will display similar behaviour as routine ‘field’ samples in the relevant measurement process. Care should be taken to ensure ‘matrix matching’ as close as practically achievable. The matrix and mineralisation style of the CRM is described in the ‘Source Material’ section and users should select appropriate CRMs matching these attributes to the field samples being analysed.

INTENDED USE

OREAS 606b is intended to cover all activities needed to produce a measurement result. This includes extraction, possible separation steps and the actual measurement process (the signal producing step). OREAS 606b may be used to calibrate the entire procedure by producing a pure substance CRM transformed into a calibration solution.

OREAS 606b is intended for the following uses:

- For the monitoring of laboratory performance in the analysis of analytes reported in Tables 1 and 2 in geological samples;
- For the verification of analytical methods for analytes reported in Tables 1 and 2;
- For the calibration of instruments used in the determination of the concentration of analytes reported in Tables 1 and 2. When a value provided in this certificate is used to calibrate a measurement process, the uncertainty associated with that value should be appropriately propagated into the user’s uncertainty calculation. Users can determine an approximation of the standard uncertainty by calculating one fourth of the width of the Expanded Uncertainty interval given in this certificate (Expanded Uncertainty intervals are provided in Tables 1 and 2).

MINIMUM SAMPLE SIZE

To relate analytical determinations to the values in this certificate, the minimum mass of sample used should match the typical mass that the laboratories used in the interlaboratory (round robin) certification program. This means that different minimum sample masses should be used depending on the operationally defined methodology as follows:

- Gold by fire assay: ≥15g;
- Gold by aqua regia digestion: ≥10g;
- Total S by infrared combustion furnace/CS analyser: ≥0.1g;
- Multi-elements by 4-acid digestion with ICP-OES and/or MS finish: ≥0.25g;
- Multi-elements by aqua regia digestion with ICP-OES and/or MS finish: ≥0.5g;
- Sulphate Sulphur via various leaching methods: ≥0.1g;
- Sulphide Sulphur via various leaching methods: ≥0.1g.

PERIOD OF VALIDITY & STORAGE INSTRUCTIONS

The certification of OREAS 606b remains valid, within the specified measurement uncertainties, until December 2033, provided the CRM is handled and stored in accordance with the instructions given below. This certification is nullified if the CRM is any way changed or contaminated.

Store in a clean and cool dry place away from direct sunlight.

Long-term stability will be monitored at appropriate intervals and purchasers notified if any changes are observed. The period of validity may well be indefinite and will be reassessed prior to expiry with the aim of extending the validity if possible.

Single-use sachets

OREAS 606b contains a very low level of reactive sulphide (0.46 wt.% S) and is packaged in single-use laminated foil sachets. Following analysis, it is the manufacturer's expectation that any remaining material is discarded. It is the user's responsibility to prevent contamination and avoid prolonged exposure of the sample to the atmosphere prior to analysis.

INSTRUCTIONS FOR HANDLING & CORRECT USE

Pre-homogenisation of the CRM prior to subsampling and analysis is not necessary as there is no particle segregation under transport [13].

Fine powders pose a risk to eyes and lungs and therefore standard precautions including the use of safety glasses and dust masks are advised.

QC monitoring using multiples of the Standard Deviation (SD)

In the application of SD's in monitoring performance it is important to note that not all laboratories function at the same level of proficiency and that different methods in use at a particular laboratory have differing levels of precision. Each laboratory has its own inherent SD (for a specific concentration level and analyte-method pair) based on the analytical process and this SD is not directly related to the round robin program.

The majority of data generated in the round robin program was produced by a selection of world class laboratories. The SD's thus generated are more constrained than those that would be produced across a randomly selected group of laboratories. To produce more generally achievable SD's the 'pooled' SD's provided in this report include interlaboratory bias. This 'one size fits all' approach may require revision at the discretion of the QC manager concerned following careful scrutiny of QC control charts.

The performance gates shown in Table 7 are intended only to be used as a preliminary guide as to what a laboratory may be able to achieve. Over a period of time monitoring your own laboratory's data for this CRM, SD's should be calculated directly from your own laboratory's process. This will enable you to establish more specific performance gates that are fit for purpose for your application as well as the ability to monitor bias. If your long-term trend analysis shows an average value that is within the 95% expanded uncertainty then generally there is no cause for concern in regard to bias.

For use with the aqua regia digestion method

It is important to note that in the analytical industry there is no standardisation of the aqua regia digestion process. This method is a partial empirical digest and differences in recoveries for various analytes are commonplace. These are caused by variations in the digest conditions and can include the ratio of nitric to hydrochloric acids, acid strength, temperatures, leach times and secondary digestions. Recoveries for sulphide-hosted base metal sulphides approach total values, however, other analytes, in particular the lithophile elements, show greater sensitivity to method parameters. This can result in lack of consensus in an inter-laboratory certification program for these elements.

The approach applied here is to report certified values in those instances where reasonable agreement exists amongst a majority of participating laboratories. The results of specific laboratories may differ significantly from the certified values, but will, nonetheless, be valid and reproducible in the context of the specifics of the aqua regia method in use. Users of this reference material should, therefore, be mindful of this limitation when applying the certified values in a quality control program.

LEGAL NOTICE

Ore Research & Exploration Pty Ltd has prepared and statistically evaluated the property values of this reference material to the best of its ability. The Purchaser by receipt hereof releases and indemnifies Ore Research & Exploration Pty Ltd from and against all liability and costs arising from the use of this material and information.

DOCUMENT HISTORY

Revision No.	Date	Changes applied
0	19th June, 2024	First publication.

CERTIFYING OFFICER

19th June, 2024

Craig Hamlyn (B.Sc. Hons - Geology), Technical Manager - ORE P/L

QMS CERTIFICATION

ORE Pty Ltd is accredited for compliance with ISO 17034:2016.

ORE Pty Ltd is ISO 9001:2015 certified by Lloyd's Register Quality Assurance Ltd for its quality management system including development, manufacturing, certification and supply of CRMs.

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[6] ISO Guide 98-3:2008. Guide to the expression of uncertainty in measurement (GUM:1995).

[7] ISO 16269:2014. Statistical interpretation of data – Part 6: Determination of statistical tolerance intervals.

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