

CERTIFICATE OF ANALYSIS FOR

PORPHYRY COPPER-GOLD-MOLYBDENUM ORE

(Cadia Valley, New South Wales, Australia)

CERTIFIED REFERENCE MATERIAL OREAS 503d

Constituent	Certified	1SD	95% Confidence Lim		95% Tolerance Limits			
Constituent	Value	130	Low	High	Low	High		
Pb Fire Assay								
Au, Gold (ppm)	0.666	0.015	0.661	0.672	0.663*	0.670*		
Aqua Regia Digestion (sample weights 10-50g)								
Au, Gold (ppm)	0.658	0.025	0.646	0.669	0.654*	0.661*		
4-Acid Digestion								
Ag, Silver (ppm)	1.34	0.066	1.31	1.37	1.27	1.41		
Cu, Copper (wt.%)	0.524	0.010	0.521	0.527	0.514	0.533		
Mo, Molybdenum (ppm)	348	10	344	352	340	356		

Summary Statistics for Key Analytes.

SI unit equivalents: ppm, parts per million \equiv mg/kg \equiv µg/g \equiv 0.0001 wt.% \equiv 1000 ppb, parts per billion.

*Gold Tolerance Limits for typical 30g fire assay and 25g aqua regia digestion methods are determined from 20 x 85mg INAA results and the Sampling Constant (Ingamells & Switzer, 1973).

Note 1: intervals may appear asymmetric due to rounding.



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Constituent	Value	SD	Low	High	Low	High
Pb Fire Assay						
Au, Gold (ppm)	0.666	0.015	0.661	0.672	0.663*	0.670*
Aqua Regia Digestion (sample	ole weights 10	-50g)		1		1
Au, Gold (ppm)	0.658	0.025	0.646	0.669	0.654*	0.661*
4-Acid Digestion				•		1
Ag, Silver (ppm)	1.34	0.066	1.31	1.37	1.27	1.41
Al, Aluminium (wt.%)	7.50	0.281	7.39	7.61	7.35	7.65
As, Arsenic (ppm)	87	5.8	85	89	83	91
Ba, Barium (ppm)	1001	34	987	1015	982	1020
Be, Beryllium (ppm)	2.93	0.226	2.83	3.03	2.81	3.04
Bi, Bismuth (ppm)	1.41	0.081	1.37	1.44	1.34	1.47
Ca, Calcium (wt.%)	2.64	0.107	2.60	2.68	2.60	2.68
Cd, Cadmium (ppm)	0.22	0.04	0.19	0.25	0.20	0.24
Ce, Cerium (ppm)	68	4.6	66	70	66	70
Co, Cobalt (ppm)	16.7	0.77	16.3	17.1	16.2	17.2
Cr, Chromium (ppm)	64	8	60	67	61	67
Cs, Cesium (ppm)	11.2	0.55	10.9	11.4	10.9	11.5
Cu, Copper (wt.%)	0.524	0.010	0.521	0.527	0.514	0.533
Dy, Dysprosium (ppm)	4.63	0.232	4.48	4.79	4.48	4.79
Er, Erbium (ppm)	2.66	0.131	2.60	2.71	2.52	2.79
Eu, Europium (ppm)	1.29	0.16	1.15	1.44	1.18	1.41
Fe, Iron (wt.%)	4.88	0.182	4.80	4.95	4.80	4.95
Ga, Gallium (ppm)	18.5	0.56	18.3	18.7	17.9	19.1
Gd, Gadolinium (ppm)	5.18	0.359	4.92	5.44	4.99	5.37
Hf, Hafnium (ppm)	2.72	0.198	2.63	2.81	2.61	2.83
Ho, Holmium (ppm)	0.92	0.052	0.88	0.96	0.86	0.98
In, Indium (ppm)	0.094	0.009	0.091	0.097	0.085	0.103
K, Potassium (wt.%)	3.13	0.131	3.08	3.18	3.07	3.19
La, Lanthanum (ppm)	34.3	2.19	33.3	35.2	33.0	35.6
Li, Lithium (ppm)	29.3	1.45	28.7	30.0	28.6	30.1
Lu, Lutetium (ppm)	0.36	0.014	0.35	0.37	0.33	0.39
Mg, Magnesium (wt.%)	1.41	0.049	1.39	1.43	1.39	1.44
Mn, Manganese (wt.%)	0.053	0.002	0.052	0.054	0.052	0.054
Mo, Molybdenum (ppm)	348	10	344	352	340	356
Na, Sodium (wt.%)	2.04	0.062	2.01	2.06	2.00	2.08
Nb, Niobium (ppm)	20.1	0.86	19.7	20.4	19.6	20.5

Table 1. Certified Values, SDs, 95% Confidence & Tolerance Limits for OREAS 503d.

*Gold Tolerance Limits for typical 30g fire assay and 25g aqua regia digestion methods are determined from 20 x 85mg INAA results and the Sampling Constant (Ingamells & Switzer, 1973).

Note 1: intervals may appear asymmetric due to rounding.



	Certified	Certified		lence Limits	95% Tolerance Limits	
Constituent	Value	SD	Low	High	Low	High
4-Acid Digestion continued		I	1			
Nd, Neodymium (ppm)	29.5	1.24	28.6	30.4	28.4	30.6
Ni, Nickel (ppm)	33.5	1.68	32.9	34.2	32.1	35.0
P, Phosphorus (wt.%)	0.108	0.004	0.106	0.110	0.106	0.110
Pb, Lead (ppm)	20.6	1.77	19.8	21.4	19.5	21.6
Pr, Praseodymium (ppm)	7.96	0.498	7.51	8.41	7.64	8.29
Rb, Rubidium (ppm)	181	8	177	185	176	186
Re, Rhenium (ppb)	246	13	241	252	235	257
S, Sulphur (wt.%)	0.798	0.027	0.788	0.807	0.777	0.818
Sb, Antimony (ppm)	3.49	0.187	3.40	3.57	3.35	3.62
Sc, Scandium (ppm)	12.5	0.54	12.2	12.7	12.1	12.8
Se, Selenium (ppm)	3.29	0.320	3.03	3.56	2.94	3.65
Sm, Samarium (ppm)	5.85	0.227	5.69	6.01	5.64	6.06
Sn, Tin (ppm)	3.47	0.170	3.39	3.55	3.30	3.64
Sr, Strontium (ppm)	363	13	358	368	355	371
Ta, Tantalum (ppm)	1.46	0.121	1.39	1.52	1.37	1.54
Tb, Terbium (ppm)	0.79	0.031	0.77	0.81	0.76	0.82
Te, Tellurium (ppm)	0.33	0.04	0.30	0.35	0.29	0.36
Th, Thorium (ppm)	17.0	0.81	16.7	17.4	16.5	17.6
Ti, Titanium (wt.%)	0.486	0.020	0.478	0.494	0.477	0.495
TI, Thallium (ppm)	0.92	0.037	0.91	0.94	0.88	0.96
Tm, Thulium (ppm)	0.36	0.04	0.33	0.39	IND	IND
U, Uranium (ppm)	4.63	0.351	4.48	4.78	4.41	4.85
V, Vanadium (ppm)	117	4	116	118	115	119
W, Tungsten (ppm)	3.47	0.286	3.39	3.55	3.18	3.76
Y, Yttrium (ppm)	23.9	1.34	23.4	24.5	23.3	24.6
Yb, Ytterbium (ppm)	2.32	0.129	2.24	2.41	2.25	2.39
Zn, Zinc (ppm)	75	4.4	73	77	73	77
Zr, Zirconium (ppm)	89	7.6	85	92	86	91
Aqua Regia Digestion						
Ag, Silver (ppm)	1.32	0.110	1.27	1.37	1.26	1.38
Al, Aluminium (wt.%)	2.08	0.117	2.03	2.13	2.04	2.12
As, Arsenic (ppm)	87	4.6	85	88	84	90
Ba, Barium (ppm)	361	23	350	372	352	370
Be, Beryllium (ppm)	0.60	0.08	0.56	0.64	0.56	0.64
Bi, Bismuth (ppm)	1.42	0.052	1.39	1.44	1.36	1.48
Ca, Calcium (wt.%)	1.19	0.079	1.16	1.22	1.17	1.21

Note 1: intervals may appear asymmetric due to rounding.



Table 1 continued.								
Constituent	Certified	SD	95% Confid	lence Limits	95% Toler	ance Limits		
Constituent	Value	30	Low	High	Low	High		
Aqua Regia Digestion continued								
Ce, Cerium (ppm)	62	4.3	60	64	61	63		
Co, Cobalt (ppm)	15.9	0.59	15.6	16.2	15.5	16.3		
Cr, Chromium (ppm)	66	3.8	65	68	65	68		
Cs, Cesium (ppm)	9.28	0.516	9.01	9.55	9.06	9.50		
Cu, Copper (wt.%)	0.522	0.011	0.518	0.526	0.513	0.531		
Dy, Dysprosium (ppm)	3.17	0.187	3.01	3.32	3.07	3.27		
Er, Erbium (ppm)	1.70	0.124	1.59	1.81	1.63	1.77		
Eu, Europium (ppm)	0.45	0.029	0.42	0.48	0.43	0.47		
Fe, Iron (wt.%)	4.58	0.175	4.51	4.65	4.51	4.65		
Ga, Gallium (ppm)	8.70	0.542	8.41	8.99	8.33	9.06		
Gd, Gadolinium (ppm)	3.91	0.129	3.79	4.04	3.77	4.06		
Ge, Germanium (ppm)	0.15	0.03	0.12	0.19	IND	IND		
Hf, Hafnium (ppm)	0.48	0.029	0.47	0.50	0.46	0.51		
Ho, Holmium (ppm)	0.60	0.044	0.56	0.64	0.58	0.63		
In, Indium (ppm)	0.074	0.005	0.072	0.075	0.067	0.080		
K, Potassium (wt.%)	1.06	0.038	1.05	1.08	1.05	1.08		
La, Lanthanum (ppm)	30.5	1.63	29.8	31.2	29.8	31.2		
Li, Lithium (ppm)	27.9	3.1	26.5	29.3	27.0	28.8		
Lu, Lutetium (ppm)	0.21	0.008	0.21	0.22	0.21	0.22		
Mg, Magnesium (wt.%)	1.21	0.051	1.19	1.23	1.19	1.23		
Mn, Manganese (wt.%)	0.041	0.001	0.041	0.042	0.040	0.042		
Mo, Molybdenum (ppm)	342	15	337	348	337	348		
Na, Sodium (wt.%)	0.209	0.027	0.197	0.221	0.201	0.217		
Nb, Niobium (ppm)	1.21	0.24	1.06	1.36	1.12	1.30		
Nd, Neodymium (ppm)	25.4	1.33	24.3	26.6	24.8	26.1		
Ni, Nickel (ppm)	31.2	1.36	30.7	31.7	30.3	32.1		
P, Phosphorus (wt.%)	0.105	0.005	0.103	0.108	0.104	0.107		
Pb, Lead (ppm)	8.39	0.602	8.13	8.65	7.97	8.81		
Pr, Praseodymium (ppm)	6.91	0.333	6.63	7.19	6.74	7.08		
Rb, Rubidium (ppm)	120	5	117	122	116	123		
Re, Rhenium (ppb)	239	9	234	243	230	247		
S, Sulphur (wt.%)	0.798	0.037	0.783	0.813	0.781	0.815		
Sb, Antimony (ppm)	2.36	0.28	2.22	2.50	2.28	2.44		
Sc, Scandium (ppm)	7.18	0.524	6.92	7.43	6.95	7.41		
Se, Selenium (ppm)	3.04	0.46	2.71	3.36	2.81	3.26		
Sm, Samarium (ppm)	4.54	0.357	4.23	4.85	4.43	4.65		

SI unit equivalents: ppm, parts per million \equiv mg/kg \equiv µg/g \equiv 0.0001 wt.% \equiv 1000 ppb, parts per billion.

Note 1: intervals may appear asymmetric due to rounding.



Table 1 continued.								
Constituent	Certified	SD	95% Confidence Limits		95% Tolerance Limits			
Constituent	Value	30	Low	High	Low	High		
Aqua Regia Digestion continu	ed							
Sn, Tin (ppm)	2.51	0.136	2.44	2.59	2.42	2.61		
Sr, Strontium (ppm)	88	11	84	93	86	90		
Tb, Terbium (ppm)	0.57	0.035	0.54	0.59	0.55	0.58		
Te, Tellurium (ppm)	0.30	0.03	0.29	0.32	0.28	0.33		
Th, Thorium (ppm)	17.4	0.56	17.1	17.6	16.8	18.0		
Ti, Titanium (wt.%)	0.318	0.019	0.309	0.326	0.311	0.324		
TI, Thallium (ppm)	0.66	0.07	0.63	0.69	0.64	0.68		
Tm, Thulium (ppm)	0.22	0.021	0.20	0.24	0.21	0.23		
U, Uranium (ppm)	4.35	0.283	4.22	4.49	4.19	4.51		
V, Vanadium (ppm)	104	5	102	107	102	107		
W, Tungsten (ppm)	1.91	0.29	1.76	2.06	1.71	2.11		
Y, Yttrium (ppm)	16.9	1.00	16.4	17.3	16.5	17.3		
Yb, Ytterbium (ppm)	1.47	0.065	1.43	1.51	1.41	1.52		
Zn, Zinc (ppm)	67	4.8	65	69	66	69		
Zr, Zirconium (ppm)	12.2	0.89	11.7	12.7	11.9	12.6		

Table 4 as ... -

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.

		Tab	e z. indicative	values		JUJU.		
Constituent	Unit	Value	Constituent	Unit	Value	Constituent	Unit	Value
Pb Fire Assa	ıy				I			
Pd	ppb	9.17	Pt	ppb	< 5			
4-Acid Diges	stion							
Ge	ppm	0.22	Hg	ppm	< 2			
Aqua Regia	Digestio	n						
В	ppm	< 10	Pd	ppb	12.3	Та	ppm	0.011
Cd	ppm	0.22	Pt	ppb	< 5			
Hg	ppm	0.042	Si	wt.%	0.105			
Borate Fusio	on XRF							
Al ₂ O ₃	wt.%	14.64	MgO	wt.%	2.44	SiO ₂	wt.%	61.70
CaO	wt.%	3.63	MnO	wt.%	0.070	SO ₃	wt.%	1.93
Fe ₂ O ₃	wt.%	7.07	Na ₂ O	wt.%	2.75	TiO ₂	wt.%	0.830
K ₂ O	wt.%	3.87	P_2O_5	wt.%	0.250			
Thermogravimetry								
LOI ¹⁰⁰⁰	wt.%	1.83						
Infrared Combustion								
С	wt.%	0.247	S	wt.%	0.767			

Table 2. Indicative Values for OREAS 503d.

SI unit equivalents: ppm, parts per million \equiv mg/kg \equiv µg/g \equiv 0.0001 wt.% \equiv 1000 ppb, parts per billion.

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 2 continued.							
Constituent	Unit	Value	Constituent	Unit	Value	Constituent	Unit	Value
Laser Ablatic	on ICP-M	S						
Ag	ppm	1.45	Hf	ppm	6.45	Sm	ppm	5.95
As	ppm	88	Но	ppm	0.91	Sn	ppm	3.40
Ва	ppm	996	In	ppm	0.10	Sr	ppm	346
Be	ppm	3.10	La	ppm	34.8	Та	ppm	1.42
Bi	ppm	1.51	Lu	ppm	0.36	Tb	ppm	0.77
Cd	ppm	0.25	Mn	wt.%	0.054	Те	ppm	0.40
Ce	ppm	67	Мо	ppm	336	Th	ppm	16.9
Co	ppm	17.6	Nb	ppm	19.9	Ti	wt.%	0.505
Cr	ppm	79	Nd	ppm	30.0	TI	ppm	1.00
Cs	ppm	11.1	Ni	ppm	33.0	Tm	ppm	0.38
Cu	wt.%	0.504	Pb	ppm	21.0	U	ppm	4.82
Dy	ppm	4.46	Pr	ppm	7.97	V	ppm	121
Er	ppm	2.48	Rb	ppm	182	W	ppm	3.25
Eu	ppm	1.22	Re	ppb	340	Y	ppm	24.5
Ga	ppm	18.2	Sb	ppm	3.65	Yb	ppm	2.47
Gd	ppm	4.82	Sc	ppm	12.3	Zn	ppm	75
Ge	ppm	1.45	Se	ppm	< 5	Zr	ppm	240

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.

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.

SOURCE MATERIAL

OREAS 503d was prepared from a blend of porphyry copper-gold ore, barren granodiorite and a minor quantity of Cu-Mo concentrate. The ore was sourced from the Ridgeway underground mine located in the Cadia Valley Operations (CVO) situated in central western New South Wales, Australia. The barren I-type hornblende-bearing granodiorite was sourced from the Late Devonian Lysterfield granodiorite complex located in eastern Melbourne, Australia.

Mineralisation in the CVO region is hosted by a sequence of late Ordovician to Early Silurian volcanics, intrusives and sediments that occur within the Bogan Gate Synclinorial Zone of the Lachlan Fold Belt. The western portion of this zone is dominated by volcanics



and host to the Late Ordovician Goonumbla porphyry copper-gold deposits. These volcanics are interpreted to have erupted from shallow water to partly emergent volcanic centres and show a broad range in composition from shoshonite through to latite to trachyte. Coeval sub-volcanic quartz monzonite porphyries (and attendant mineralisation) have intruded the volcanics. They are generally small, sub-vertical, pipe-like intrusives. Typically the mineralised porphyries contain plagioclase and quartz phenocrysts in a matrix of fine-grained potassium feldspar and quartz with minor biotite and hornblende.

Copper-gold mineralisation occurs as stockwork quartz veins and disseminations associated with potassic alteration. This alteration is intimately associated spatially and temporally with the small finger-like quartz monzonite porphyries that intrude the Goonumbla Volcanics. Sulphides are zoned laterally from the centres of mineralisation. The central portions are bornite-rich with minor chalcopyrite, zoning outward through equal concentrations of bornite and chalcopyrite, to an outermost chalcopyrite-rich zone. Pyrite increases outward at the expense of bornite.

COMMINUTION AND HOMOGENISATION PROCEDURES

The material constituting OREAS 503d was prepared in the following manner:

- Drying of ores and barren granodiorite to constant mass at 105°C;
- Drying of Cu-Mo concentrate to constant mass at 85°C;
- Multi-stage milling of ore and concentrate to 100% minus 30 microns;
- Milling of barren granodiorite to 98% minus 75 microns;
- Combining ore, granodiorite and concentrate in appropriate proportions to achieve target grades;
- Homogenisation;
- Packaging into 10 and 60g units in laminated foil pouches and 500g units in plastic jars.

PHYSICAL PROPERTIES

OREAS 503d was tested at ORE Research & Exploration Pty Ltd's onsite laboratory for various physical properties. Table 3 presents these findings which should be used for informational purposes only.

CRM Name	Bulk Density (g/L)	Moisture%	Munsell Notation [‡]	Munsell Color [‡]				
OREAS 503d	EAS 503d 695.2 0.57		N7	Light Gray				

Table 3. Physical properties of OREAS 503d.

[‡]The Munsell Rock Color Chart helps geologists and archeologists communicate with color more effectively by crossreferencing ISCC-NBS color names with unique Munsell alpha-numeric color notations for rock color samples.



ANALYTICAL PROGRAM

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

- Gold via 15-50g fire assay with AAS finish (14 laboratories) and ICP-OES (12 laboratories) finish;
- Gold via 15-50g aqua regia digestion with ICP-MS finish (11 laboratories), AAS (4 laboratories) and ICP-OES (1 laboratory) finish;
- Aqua regia digestion for full elemental suite ICP-OES/MS and AAS finish (up to 26 laboratories depending on the element);
- 4-Acid digestion for full elemental suite ICP-OES/MS finish (up to 25 laboratories depending on the element);
- Gold by instrumental neutron activation analysis (INAA) on 20 x 85mg subsamples to confirm homogeneity (1 laboratory only: ANSTO, Lucas Heights, Australia);
- Major and trace elements determined by borate fusion XRF (Al₂O₃ to TiO₂), laser ablation with ICP-MS (Ag to Zr), LOI at 1000°C and C+S by infrared combustion furnace (1 laboratory only: Bureau Veritas Geoanalytical, Perth, Australia).

It is important to note that in the analytical industry there is no standardisation of the aqua regia digestion process. Aqua regia is a partial empirical digest and differences in recoveries for various analytes are commonplace. These are caused by variations in the digest conditions which 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.

For the round robin program twenty 1kg test units 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 received by each laboratory were obtained by taking two 100g samples from each of three separate 1kg test units. This format enabled nested ANOVA treatment of the results to evaluate homogeneity, i.e. to ascertain whether between-unit variance is greater than within-unit variance.

Table 1 presents the 119 certified values together with their associated 1SD's, 95% confidence and tolerance limits. Table 2 shows 76 indicative values for major and trace element composition and Table 3 provides indicative physical properties information. Gold homogeneity has been evaluated and confirmed by instrumental neutron activation analysis (INAA) on twenty ~85mg sample portions (see Table 4) and by a nested ANOVA program for both fire assay and aqua regia digestion (see '**nested ANOVA**' section).



Table 5 provides performance gate intervals for the certified values based on their pooled 1SDs. Tabulated results of all elements together with uncorrected means, medians, standard deviations, relative standard deviations and per cent deviation of lab means from the corrected mean of means (PDM³) are presented in the detailed certification data for this CRM (**OREAS 503d DataPack-1.0.190408_171711.xlsx**).

Results are also presented in scatter plots for gold by fire assay, gold by aqua regia digestion, silver by 4-acid digestion, copper by 4-acid digestion and molybdenum by 4-acid digestion (Figures 1 to 5, respectively) together with ± 3 SD (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.

STATISTICAL ANALYSIS

Certified Values, Confidence Limits, Standard Deviations and Tolerance Limits (Table 1) 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. In certain instances statistician's prerogative has been employed in discriminating outliers.

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.

Certified Values are the means of accepted laboratory means after outlier filtering. The INAA data (see Table 4) is omitted from determination of the certified value for Au and is used solely for the calculation of Tolerance Limits and homogeneity evaluation of OREAS 503d.

95% Confidence Limits are inversely proportional to the number of participating laboratories and inter-laboratory agreement. It is a measure of the reliability of the certified value. A 95% confidence interval indicates a 95% probability that the true value of the analyte under consideration lies between the upper and lower limits. *95% Confidence Limits should not be used as control limits for laboratory performance.*

Indicative (uncertified) values are provided in Table 2. Additional indicative values by other analytical methods are present where the number of laboratories reporting a particular analyte is insufficient (< 5) to support certification or where inter-laboratory consensus is poor.

Standard Deviation values (1SDs) are reported in Table 1. They 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.

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.

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 inter-lab 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.

Homogeneity Evaluation

The tolerance limits (ISO 16269:2014) shown in Table 1 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 $(1-\alpha=0.99)$ at least 95% of subsamples ($\rho=0.95$) will have concentrations lying between 0.514 and 0.533 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.*

Table 4 below shows the INAA data determined on 20 x 85mg subsamples of OREAS 503d. An equivalent scaled version of the results is also provided to demonstrate an appreciation of what this data means if 30g fire assay determinations were undertaken without the normal measurement error associated with this methodology.

The homogeneity of gold has been determined by INAA using the reduced analytical subsample method which utilises the known relationship between standard deviation and analytical subsample weight (Ingamells and Switzer, 1973). 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. In this instance a subsample weight of 85 milligrams was employed and the 1RSD of 0.147% was calculated for a 30g fire assay or aqua regia sample (2.75% at 85mg weights). These confirm the high level of gold homogeneity in OREAS 503d.



Replicate	Au	Au
No	85mg actual	30g equivalent*
		- · ·
1	0.662	0.669
2	0.680	0.670
3	0.655	0.668
4	0.698	0.671
5	0.668	0.669
6	0.717	0.672
7	0.681	0.670
8	0.649	0.668
9	0.669	0.669
10	0.678	0.670
11	0.673	0.669
12	0.668	0.669
13	0.680	0.670
14	0.648	0.668
15	0.663	0.669
16	0.657	0.668
17	0.645	0.668
18	0.685	0.670
19	0.660	0.669
20	0.644	0.668
Mean	0.669	0.669
Median	0.668	0.669
Std Dev.	0.018	0.001
Rel.Std.Dev.	2.75%	0.147%

Table 4. Neutron Activation Analysis of Au (in ppm) on 20 x 85mg subsamples showing
the equivalent results scaled to a typical fire assay (30g sample mass) method.

*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

 \overline{X} = mean of 85mg INAA results

The homogeneity of OREAS 503d has also been evaluated in a **nested ANOVA** of the round robin program. Each of the twenty-five round robin laboratories received six samples per CRM and these samples were made up of paired samples from three different, non-adjacent sampling intervals. The purpose of the ANOVA evaluation 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 503d. The test was performed using the following parameters:

- Gold fire assay 156 samples (26 laboratories each providing analyses on 3 pairs of samples);
- Gold aqua regia digestion 102 samples (17 laboratories each providing analyses on 3 pairs of samples);
- Null Hypothesis, H₀: Between-unit variance is no greater than within-unit variance (reject H₀ if *p*-value < 0.05);
- Alternative Hypothesis, H_1 : Between-unit variance is greater than within-unit variance.



P-values are a measure of probability where values less than 0.05 indicate a greater than 95% probability that the observed differences in within-unit and between-unit variances are real. The datasets were filtered for both individual and laboratory data set (batch) outliers prior to the calculation of *p*-values. This process derived *p*-values of 0.999 for Au by fire assay and 0.953 for Au by aqua regia digestion. Both *p*-values are insignificant and the Null Hypothesis is retained. Additionally, none of the other certified values showed significant p-values. Please note that only results for constituents present in concentrations well above the detection levels (i.e. >20 x Lower Limit of Detection) for the various methods undertaken were considered for the objective of evaluating homogeneity

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 503d and whether the variance between two subsamples from the same unit is statistically distinguishable to the variance from two subsamples taken from any two separate units. A reference material therefore, can possess poor absolute homogeneity yet still pass a relative homogeneity test if the within-unit heterogeneity is large and similar across all units.

Based on the statistical analysis of the results of the inter-laboratory certification program it can be concluded that OREAS 503d is fit-for-purpose as a certified reference material (see 'Intended Use' below).

Performance Gates

Table 5 shows 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. 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) \pm 10%.

i.e. Certified Value ± 10% ± 2DL	(adapted from Govett, 1983)
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Constituent	Certified		Absolute	Standard	Deviations Relative			Standard D	eviations	5% w	indow
Constituent	Value	1SD	2SD Low	2SD High	3SD Low	3SD High	1RSD	2RSD	3RSD	Low	High
Pb Fire Assay											
Au, ppm	0.666	0.015	0.636	0.697	0.621	0.712	2.30%	4.59%	6.89%	0.633	0.700
Aqua Regia Digestion (sample weights 10-50g)											
Au, ppm	0.658	0.025	0.609	0.707	0.584	0.732	3.74%	7.48%	11.21%	0.625	0.691

 Table 5. Performance Gates for OREAS 503d.

SI unit equivalents: ppm, parts per million \equiv mg/kg \equiv µg/g \equiv 0.0001 wt.% \equiv 1000 ppb, parts per billion.

Note 1: intervals may appear asymmetric due to rounding.



Constituent	Certified		Absolute	Standard	Deviations	5	Relative	Standard D	5% w	indow	
Constituent	Value	1SD	2SD Low	2SD High	3SD Low	3SD High	1RSD	2RSD	3RSD	Low	High
4-Acid Digest	ion										
Ag, ppm	1.34	0.066	1.21	1.47	1.14	1.54	4.94%	9.87%	14.81%	1.27	1.41
Al, wt.%	7.50	0.281	6.94	8.06	6.66	8.34	3.74%	7.49%	11.23%	7.12	7.87
As, ppm	87	5.8	76	99	70	105	6.69%	13.37%	20.06%	83	92
Ba, ppm	1001	34	932	1070	898	1104	3.44%	6.88%	10.32%	951	1051
Be, ppm	2.93	0.226	2.47	3.38	2.25	3.60	7.72%	15.44%	23.16%	2.78	3.07
Bi, ppm	1.41	0.081	1.24	1.57	1.16	1.65	5.80%	11.59%	17.39%	1.34	1.48
Ca, wt.%	2.64	0.107	2.43	2.85	2.32	2.96	4.06%	8.12%	12.19%	2.51	2.77
Cd, ppm	0.22	0.04	0.13	0.31	0.09	0.35	20.42%	40.84%	61.26%	0.21	0.23
Ce, ppm	68	4.6	59	77	54	82	6.82%	13.64%	20.46%	65	71
Co, ppm	16.7	0.77	15.2	18.3	14.4	19.0	4.58%	9.16%	13.74%	15.9	17.6
Cr, ppm	64	8	47	81	39	89	13.16%	26.31%	39.47%	61	67
Cs, ppm	11.2	0.55	10.1	12.3	9.5	12.8	4.95%	9.90%	14.85%	10.6	11.7
Cu, wt.%	0.524	0.010	0.504	0.543	0.495	0.553	1.86%	3.73%	5.59%	0.498	0.550
Dy, ppm	4.63	0.232	4.17	5.10	3.94	5.33	5.00%	10.00%	15.00%	4.40	4.87
Er, ppm	2.66	0.131	2.40	2.92	2.26	3.05	4.94%	9.87%	14.81%	2.53	2.79
Eu, ppm	1.29	0.16	0.97	1.62	0.81	1.78	12.49%	24.97%	37.46%	1.23	1.36
Fe, wt.%	4.88	0.182	4.51	5.24	4.33	5.42	3.72%	7.45%	11.17%	4.63	5.12
Ga, ppm	18.5	0.56	17.4	19.6	16.8	20.2	3.05%	6.10%	9.15%	17.6	19.4
Gd, ppm	5.18	0.359	4.46	5.90	4.11	6.26	6.93%	13.85%	20.78%	4.92	5.44
Hf, ppm	2.72	0.198	2.32	3.12	2.13	3.32	7.29%	14.59%	21.88%	2.58	2.86
Ho, ppm	0.92	0.052	0.82	1.03	0.76	1.08	5.65%	11.30%	16.96%	0.88	0.97
In, ppm	0.094	0.009	0.075	0.112	0.066	0.121	9.78%	19.56%	29.34%	0.089	0.098
K, wt.%	3.13	0.131	2.87	3.39	2.74	3.52	4.19%	8.38%	12.57%	2.97	3.29
La, ppm	34.3	2.19	29.9	38.7	27.7	40.8	6.39%	12.79%	19.18%	32.6	36.0
Li, ppm	29.3	1.45	26.4	32.2	25.0	33.7	4.95%	9.90%	14.85%	27.9	30.8
Lu, ppm	0.36	0.014	0.33	0.38	0.32	0.40	3.81%	7.62%	11.43%	0.34	0.37
Mg, wt.%	1.41	0.049	1.31	1.51	1.26	1.56	3.48%	6.96%	10.44%	1.34	1.48
Mn, wt.%	0.053	0.002	0.048	0.058	0.046	0.060	4.46%	8.91%	13.37%	0.050	0.056
Mo, ppm	348	10	327	369	317	379	2.97%	5.94%	8.92%	331	366
Na, wt.%	2.04	0.062	1.91	2.16	1.85	2.22	3.04%	6.07%	9.11%	1.94	2.14
Nb, ppm	20.1	0.86	18.3	21.8	17.5	22.6	4.29%	8.59%	12.88%	19.1	21.1
Nd, ppm	29.5	1.24	27.1	32.0	25.8	33.3	4.21%	8.43%	12.64%	28.1	31.0
Ni, ppm	33.5	1.68	30.2	36.9	28.5	38.6	5.01%	10.02%	15.03%	31.9	35.2
P, wt.%	0.108	0.004	0.099	0.117	0.095	0.121	4.03%	8.05%	12.08%	0.103	0.113
Pb, ppm	20.6	1.77	17.0	24.1	15.3	25.9	8.60%	17.19%	25.79%	19.5	21.6
Pr, ppm	7.96	0.498	6.97	8.96	6.47	9.46	6.26%	12.51%	18.77%	7.56	8.36
Rb, ppm	181	8	164	198	156	207	4.69%	9.38%	14.07%	172	190
Re, ppb	246	13	221	272	208	284	5.15%	10.29%	15.44%	234	259
S, wt.%	0.798	0.027	0.744	0.852	0.717	0.879	3.39%	6.77%	10.16%	0.758	0.838
Sb, ppm	3.49	0.187	3.11	3.86	2.92	4.05	5.38%	10.75%	16.13%	3.31	3.66

Table 5 continued.

SI unit equivalents: ppm, parts per million \equiv mg/kg \equiv µg/g \equiv 0.0001 wt.% \equiv 1000 ppb, parts per billion.

Note 1: intervals may appear asymmetric due to rounding.



				IUN							
	Certified		Absolute	Standard	Deviations	5	Relative	Standard D	eviations	5% window	
Constituent	Value	1SD	2SD Low	2SD High	3SD Low	3SD High	1RSD	2RSD	3RSD	Low	High
4-Acid Digest	ion continu	ed						I			
Sc, ppm	12.5	0.54	11.4	13.6	10.8	14.1	4.34%	8.68%	13.02%	11.8	13.1
Se, ppm	3.29	0.320	2.65	3.93	2.33	4.25	9.71%	19.41%	29.12%	3.13	3.46
Sm, ppm	5.85	0.227	5.40	6.30	5.17	6.53	3.88%	7.77%	11.65%	5.56	6.14
Sn, ppm	3.47	0.170	3.13	3.81	2.96	3.98	4.90%	9.80%	14.70%	3.30	3.64
Sr, ppm	363	13	337	389	324	402	3.62%	7.23%	10.85%	345	381
Ta, ppm	1.46	0.121	1.22	1.70	1.10	1.82	8.27%	16.54%	24.81%	1.38	1.53
Tb, ppm	0.79	0.031	0.73	0.85	0.70	0.89	3.98%	7.96%	11.94%	0.75	0.83
Te, ppm	0.33	0.04	0.24	0.41	0.20	0.45	12.96%	25.92%	38.88%	0.31	0.34
Th, ppm	17.0	0.81	15.4	18.7	14.6	19.5	4.73%	9.47%	14.20%	16.2	17.9
Ti, wt.%	0.486	0.020	0.447	0.525	0.427	0.545	4.04%	8.08%	12.13%	0.462	0.510
TI, ppm	0.92	0.037	0.85	1.00	0.81	1.03	3.96%	7.92%	11.87%	0.88	0.97
Tm, ppm	0.36	0.04	0.29	0.44	0.25	0.47	10.15%	20.29%	30.44%	0.34	0.38
U, ppm	4.63	0.351	3.93	5.33	3.58	5.69	7.58%	15.17%	22.75%	4.40	4.86
V, ppm	117	4	110	124	106	128	3.12%	6.25%	9.37%	111	123
W, ppm	3.47	0.286	2.90	4.04	2.61	4.33	8.24%	16.47%	24.71%	3.30	3.64
Y, ppm	23.9	1.34	21.2	26.6	19.9	28.0	5.62%	11.24%	16.86%	22.7	25.1
Yb, ppm	2.32	0.129	2.07	2.58	1.94	2.71	5.54%	11.08%	16.62%	2.21	2.44
Zn, ppm	75	4.4	67	84	62	89	5.78%	11.56%	17.34%	72	79
Zr, ppm	89	7.6	73	104	66	112	8.62%	17.24%	25.86%	84	93
Aqua Regia D	igestion						1	<u> </u>			1
Ag, ppm	1.32	0.110	1.10	1.54	0.99	1.65	8.36%	16.71%	25.07%	1.26	1.39
Al, wt.%	2.08	0.117	1.85	2.31	1.73	2.43	5.60%	11.20%	16.81%	1.98	2.18
As, ppm	87	4.6	78	96	73	100	5.25%	10.51%	15.76%	82	91
Ba, ppm	361	23	314	408	291	432	6.49%	12.98%	19.47%	343	379
Be, ppm	0.60	0.08	0.45	0.75	0.37	0.83	12.86%	25.71%	38.57%	0.57	0.63
Bi, ppm	1.42	0.052	1.31	1.52	1.26	1.57	3.71%	7.41%	11.12%	1.34	1.49
Ca, wt.%	1.19	0.079	1.03	1.35	0.95	1.43	6.68%	13.35%	20.03%	1.13	1.25
Ce, ppm	62	4.3	53	71	49	75	7.00%	14.01%	21.01%	59	65
Co, ppm	15.9	0.59	14.7	17.1	14.1	17.7	3.70%	7.41%	11.11%	15.1	16.7
Cr, ppm	66	3.8	59	74	55	78	5.69%	11.39%	17.08%	63	70
Cs, ppm	9.28	0.516	8.25	10.31	7.73	10.83	5.56%	11.12%	16.67%	8.82	9.74
Cu, wt.%	0.522	0.011	0.499	0.545	0.488	0.556	2.19%	4.39%	6.58%	0.496	0.548
Dy, ppm	3.17	0.187	2.79	3.54	2.61	3.73	5.91%	11.82%	17.73%	3.01	3.33
Er, ppm	1.70	0.124	1.45	1.95	1.33	2.07	7.32%	14.64%	21.97%	1.62	1.79
Eu, ppm	0.45	0.029	0.39	0.51	0.36	0.54	6.44%	12.88%	19.32%	0.43	0.47
Fe, wt.%	4.58	0.175	4.23	4.93	4.05	5.11	3.82%	7.65%	11.47%	4.35	4.81
Ga, ppm	8.70	0.542	7.62	9.78	7.07	10.32	6.23%	12.45%	18.68%	8.26	9.13
Gd, ppm	3.91	0.129	3.66	4.17	3.53	4.30	3.29%	6.57%	9.86%	3.72	4.11
Ge, ppm	0.15	0.03	0.10	0.21	0.07	0.24	18.88%	37.76%	56.65%	0.15	0.16
Hf, ppm	0.48	0.029	0.43	0.54	0.40	0.57	6.09%	12.18%	18.26%	0.46	0.51

Table 5 continued.

SI unit equivalents: ppm, parts per million \equiv mg/kg \equiv µg/g \equiv 0.0001 wt.% \equiv 1000 ppb, parts per billion.

Note 1: intervals may appear asymmetric due to rounding.



			Absolute	Standard			Relative	Standard D	eviations	5% window	
Constituent	Certified Value			2SD		3SD	TClative			570 W	
	Value	1SD	2SD Low	High	3SD Low	High	1RSD	2RSD	3RSD	Low	High
Aqua Regia D	Aqua Regia Digestion continued										
Ho, ppm	0.60	0.044	0.51	0.69	0.47	0.74	7.35%	14.70%	22.06%	0.57	0.63
In, ppm	0.074	0.005	0.063	0.084	0.058	0.090	7.29%	14.58%	21.87%	0.070	0.077
K, wt.%	1.06	0.038	0.99	1.14	0.95	1.18	3.61%	7.23%	10.84%	1.01	1.12
La, ppm	30.5	1.63	27.2	33.7	25.6	35.4	5.33%	10.66%	15.99%	29.0	32.0
Li, ppm	27.9	3.1	21.8	34.0	18.7	37.1	11.00%	22.00%	33.01%	26.5	29.3
Lu, ppm	0.21	0.008	0.20	0.23	0.19	0.24	3.52%	7.04%	10.56%	0.20	0.22
Mg, wt.%	1.21	0.051	1.11	1.31	1.06	1.36	4.24%	8.48%	12.72%	1.15	1.27
Mn, wt.%	0.041	0.001	0.038	0.044	0.037	0.045	3.54%	7.09%	10.63%	0.039	0.043
Mo, ppm	342	15	313	372	298	386	4.29%	8.58%	12.87%	325	360
Na, wt.%	0.209	0.027	0.155	0.263	0.128	0.290	12.94%	25.89%	38.83%	0.199	0.220
Nb, ppm	1.21	0.24	0.73	1.69	0.49	1.93	19.72%	39.44%	59.16%	1.15	1.27
Nd, ppm	25.4	1.33	22.8	28.1	21.4	29.4	5.24%	10.49%	15.73%	24.2	26.7
Ni, ppm	31.2	1.36	28.5	34.0	27.2	35.3	4.35%	8.69%	13.04%	29.7	32.8
P, wt.%	0.105	0.005	0.095	0.116	0.090	0.121	4.97%	9.95%	14.92%	0.100	0.111
Pb, ppm	8.39	0.602	7.19	9.59	6.58	10.19	7.17%	14.34%	21.52%	7.97	8.81
Pr, ppm	6.91	0.333	6.25	7.58	5.91	7.91	4.82%	9.64%	14.46%	6.57	7.26
Rb, ppm	120	5	110	129	105	134	4.04%	8.09%	12.13%	114	126
Re, ppb	239	9	221	256	213	264	3.60%	7.20%	10.81%	227	251
S, wt.%	0.798	0.037	0.724	0.872	0.687	0.909	4.63%	9.25%	13.88%	0.758	0.838
Sb, ppm	2.36	0.28	1.79	2.93	1.51	3.21	12.04%	24.08%	36.11%	2.24	2.48
Sc, ppm	7.18	0.524	6.13	8.23	5.60	8.75	7.31%	14.61%	21.92%	6.82	7.54
Se, ppm	3.04	0.46	2.12	3.96	1.65	4.42	15.18%	30.36%	45.54%	2.89	3.19
Sm, ppm	4.54	0.357	3.82	5.25	3.47	5.61	7.87%	15.74%	23.61%	4.31	4.77
Sn, ppm	2.51	0.136	2.24	2.79	2.11	2.92	5.41%	10.81%	16.22%	2.39	2.64
Sr, ppm	88	11	67	110	56	121	12.18%	24.36%	36.54%	84	93
Tb, ppm	0.57	0.035	0.50	0.64	0.46	0.67	6.23%	12.47%	18.70%	0.54	0.60
Te, ppm	0.30	0.03	0.24	0.37	0.21	0.40	10.71%	21.41%	32.12%	0.29	0.32
Th, ppm	17.4	0.56	16.3	18.5	15.7	19.1	3.21%	6.41%	9.62%	16.5	18.3
Ti, wt.%	0.318	0.019	0.279	0.356	0.260	0.375	6.02%	12.03%	18.05%	0.302	0.333
TI, ppm	0.66	0.07	0.53	0.79	0.46	0.86	10.03%	20.06%	30.10%	0.63	0.69
Tm, ppm	0.22	0.021	0.18	0.26	0.16	0.28	9.56%	19.13%	28.69%	0.21	0.23
U, ppm	4.35	0.283	3.79	4.92	3.51	5.20	6.49%	12.99%	19.48%	4.14	4.57
V, ppm	104	5	94	115	89	120	4.89%	9.78%	14.66%	99	110
W, ppm	1.91	0.29	1.34	2.48	1.05	2.76	14.93%	29.87%	44.80%	1.81	2.00
Y, ppm	16.9	1.00	14.9	18.9	13.9	19.9	5.91%	11.82%	17.72%	16.0	17.7
Yb, ppm	1.47	0.065	1.34	1.60	1.27	1.66	4.45%	8.90%	13.35%	1.39	1.54
Zn, ppm	67	4.8	58	77	53	82	7.12%	14.23%	21.35%	64	71
Zr, ppm	12.2	0.89	10.4	14.0	9.5	14.9	7.32%	14.64%	21.97%	11.6	12.8

Table 5 continued.

SI unit equivalents: ppm, parts per million \equiv mg/kg \equiv µg/g \equiv 0.0001 wt.% \equiv 1000 ppb, parts per billion.

Note 1: intervals may appear asymmetric due to rounding.

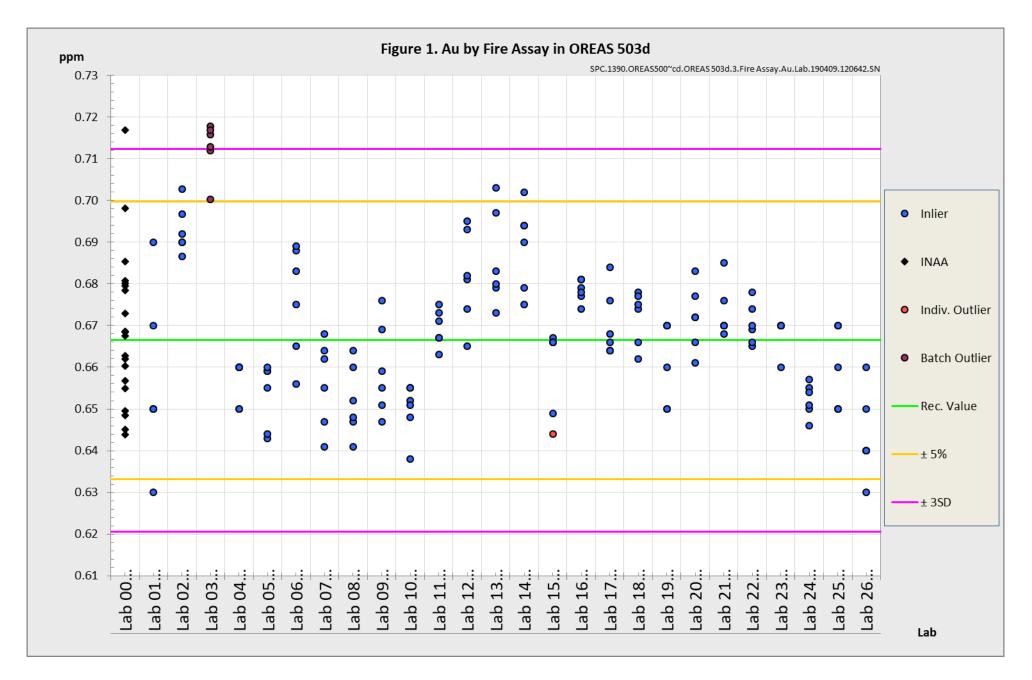


PARTICIPATING LABORATORIES

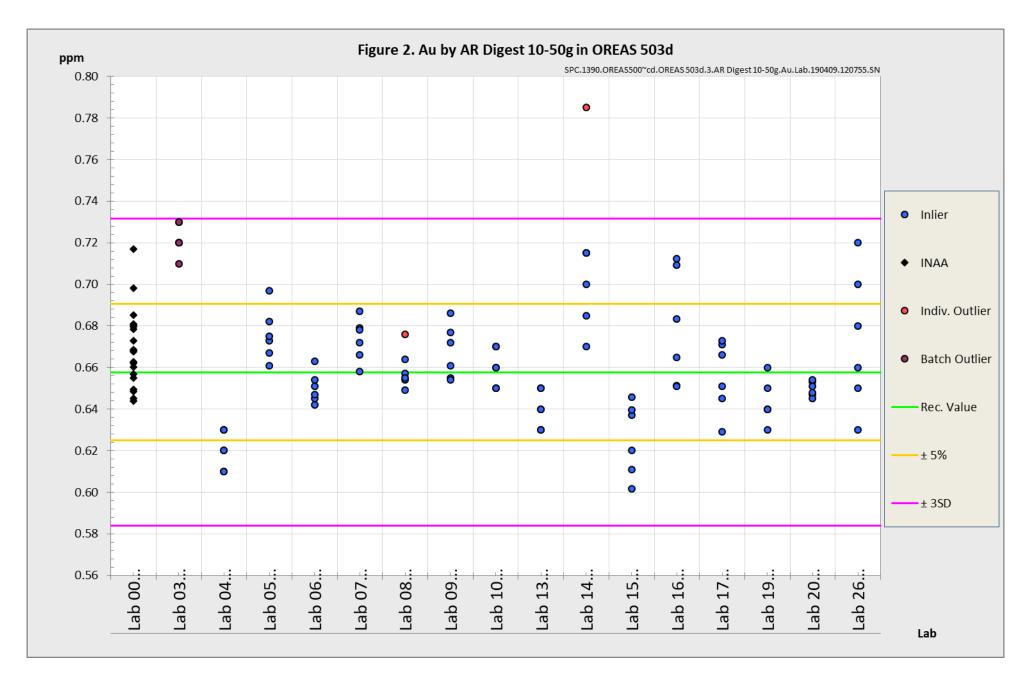
- 1. Alex Stewart International, Mendoza, Argentina
- 2. ALS, Johannesburg, South Africa
- 3. ALS, Lima, Peru
- 4. ALS, Loughrea, Galway, Ireland
- 5. ALS, Perth, WA, Australia
- 6. ALS, Ulaanbaatar, Khan-Uul District, Mongolia
- 7. ALS, Vancouver, BC, Canada
- 8. ANSTO, Lucas Heights, NSW, Australia
- 9. Bureau Veritas Commodities Canada Ltd, Vancouver, BC, Canada
- 10. Bureau Veritas Geoanalytical, Perth, WA, Australia
- 11. CERTIMIN, Lima, Peru
- 12. ESAN Istanbul, Istanbul, Turkey
- 13. Inspectorate (BV), Lima, Peru
- 14. Inspectorate America Corporation (BV), Sparks, Nevada, USA
- 15. Intertek Genalysis, Adelaide, SA, Australia
- 16. Intertek Genalysis, Perth, WA, Australia
- 17. Intertek Testing Services, Townsville, QLD, Australia
- 18. Nagrom, Perth, WA, Australia
- 19. Newcrest Laboratory Services, Orange, NSW, Australia
- 20. PT Geoservices Ltd, Cikarang, Jakarta Raya, Indonesia
- 21. PT Intertek Utama Services, Jakarta Timur, DKI Jakarta, Indonesia
- 22. PT SGS Indo Assay Laboratories, Jakarta, Indonesia
- 23. SGS, Ankara, Anatolia, Turkey
- 24. SGS Canada Inc., Vancouver, BC, Canada
- 25. SGS del Peru, Lima, Peru
- 26. SGS Mongolia, Ulan Bator, Mongolia
- 27. Shiva Analyticals Ltd, Bangalore North, Karnataka, India

Please note: Above numbered alphabetical list of participating laboratories <u>does not</u> reflect the Lab ID numbering on the scatter plots below.

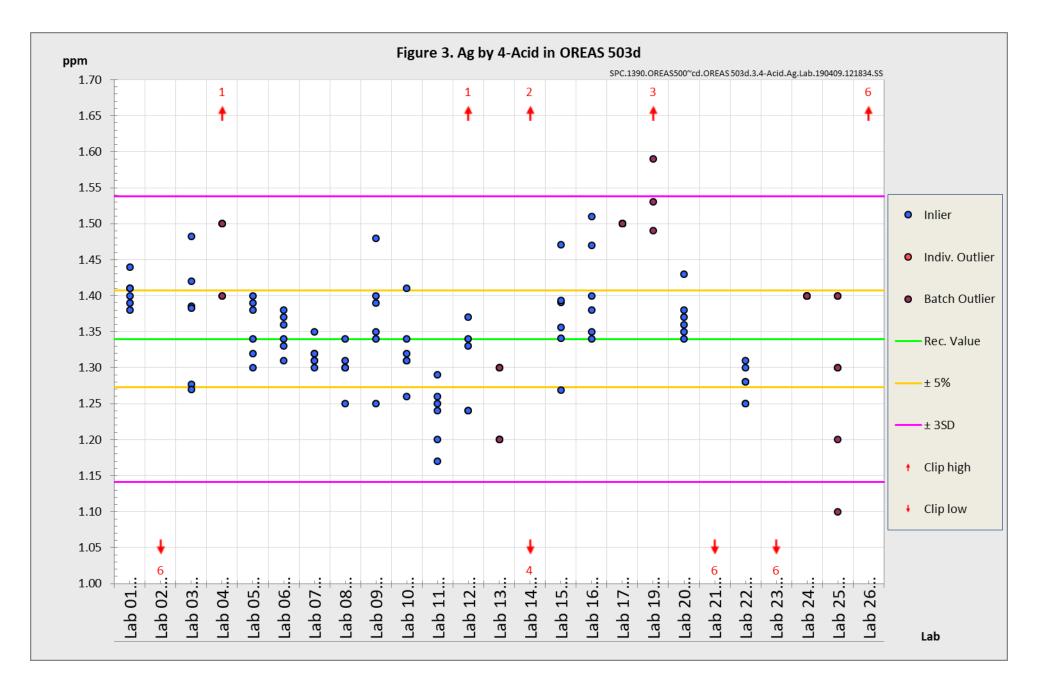




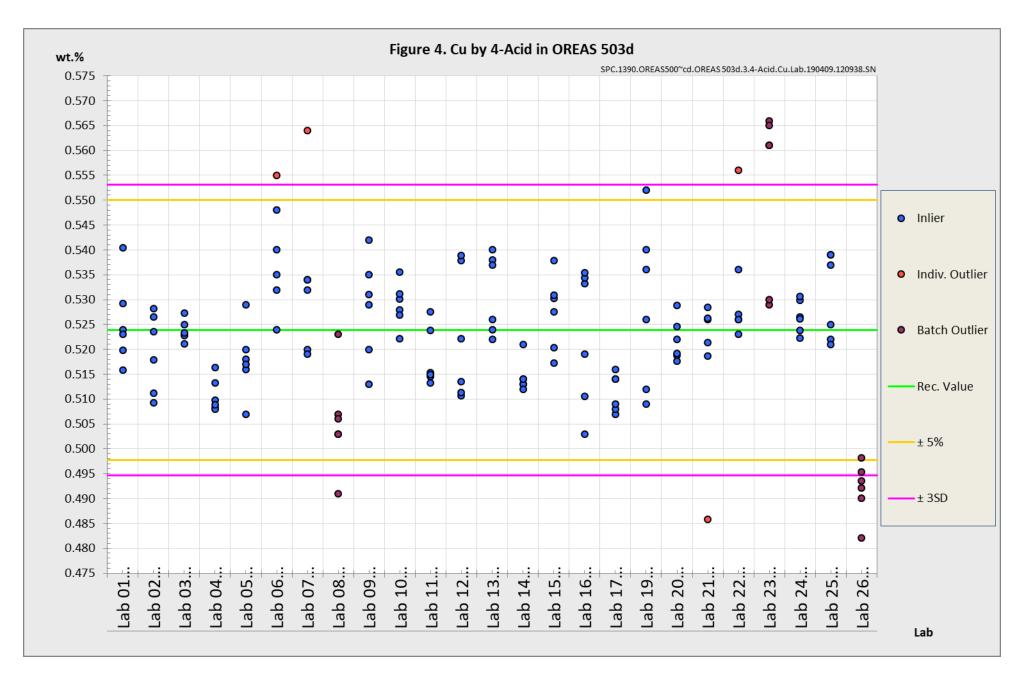




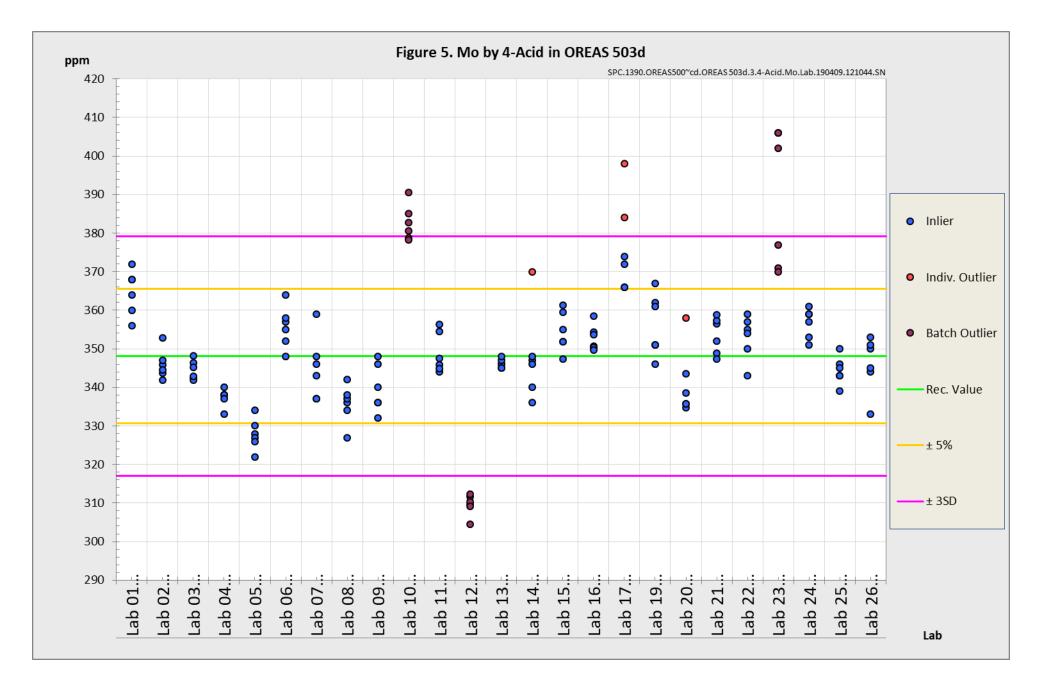














PREPARER AND SUPPLIER

Certified reference material OREAS 503d was prepared, certified and supplied by:



ORE Research & Exploration Pty LtdTel:+613-9729 033337A Hosie StreetFax:+613-9729 8338Bayswater North VIC 3153Web:www.ore.com.auAUSTRALIAEmail:info@ore.com.au

It is available in unit sizes of 10g and 60g (single-use laminated foil pouches) and 500g (plastic jars).

METROLOGICAL TRACEABILITY

The analytical samples were selected in a manner to represent the entire batch of prepared 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 that underlie the consensus values. Each analytical data set has been validated by its assayer through the inclusion of internal reference materials and QC checks during analysis.

The laboratories were chosen on the basis of their competence (from past performance in inter-laboratory programs undertaken by ORE Pty Ltd) for a particular analytical method, analyte or analyte suite, and sample matrix. Most of these laboratories have and maintain ISO 17025 accreditation. The certified values presented in this report are calculated from the means of accepted data following robust statistical treatment 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 method is possible. In this case, certification takes place on the basis of agreement among independent measurement results (see ISO Guide 35:2006, Clause 10)."*

COMMUTABILITY

The measurements of the results that underlie the certified values contained in this report were undertaken by methods involving pre-treatment (digestion/fusion) 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 their field samples.



INTENDED USE

OREAS 503d 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 503d may be used to calibrate the entire procedure by producing a pure substance CRM transformed into a calibration solution.

OREAS 503d is intended for the following uses:

- For the monitoring of laboratory performance in the analysis of analytes reported in Table 1 in geological samples;
- For the verification of analytical methods for analytes reported in Table 1;
- For the calibration of instruments used in the determination of the concentration of analytes reported in Table 1.

STABILITY AND STORAGE INSTRUCTIONS

OREAS 503c has been prepared from a blend of porphyry copper-gold ore, barren granodiorite and a minor quantity of Cu-Mo concentrate. It is low in reactive sulphide (~0.8% S). In its unopened state and under normal conditions of storage it has a shelf life beyond ten years. Its stability will be monitored at regular intervals and purchasers notified if any changes are observed.

INSTRUCTIONS FOR CORRECT USE

The certified values for OREAS 503d refer to the concentration level in its packaged state. It should not be dried prior to weighing and analysis.

HANDLING INSTRUCTIONS

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

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	11 th April 2019	First publication.



QMS ACCREDITED

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



CERTIFYING OFFICER

11th April, 2019

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

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