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CERTIFICATE OF ANALYSIS FOR

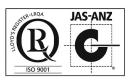
CERTIFIED REFERENCE MATERIAL

OREAS 551b

Copper-Cobalt Ore (Democratic Republic of the Congo)



Accredited for compliance with ISO 17034



COA-1904-OREA551b-R0 BUP-70-10-01 Ver:2.0

Table 1. Certified Values, Uncertainty & Tolerance Intervals for multi-elements by 4-acid digestion in OREAS 551b.

	1					
Constituent	Certified	95 % Expanded Uncertainty		95 % Tolerance Limits		
	Value [†]	Low	High	Low	High	
4-Acid Digestion	1	-			1	
Ag, Silver (ppm)	0.302	0.226	0.379	0.230	0.374	
Al, Aluminium (wt.%)	6.18	5.99	6.37	6.08	6.28	
As, Arsenic (ppm)	50	48	53	48	52	
Ba, Barium (ppm)	328	317	340	319	338	
Be, Beryllium (ppm)	4.14	3.95	4.32	4.03	4.24	
Bi, Bismuth (ppm)	2.64	2.53	2.75	2.56	2.72	
Ca, Calcium (wt.%)	0.835	0.811	0.859	0.823	0.846	
Cd, Cadmium (ppm)	0.31	0.28	0.34	0.28	0.34	
Ce, Cerium (ppm)	145	137	153	140	149	
Co, Cobalt (wt.%)	0.285	0.278	0.292	0.282	0.288	
Cr, Chromium (ppm)	79	73	86	76	83	
Cs, Caesium (ppm)	1.89	1.78	2.01	1.80	1.98	
Cu, Copper (wt.%)	2.77	2.71	2.83	2.74	2.80	
Dy, Dysprosium (ppm)	4.46	3.96	4.95	4.31	4.60	
Er, Erbium (ppm)	2.72	2.39	3.06	2.61	2.84	
Eu, Europium (ppm)	0.80	0.73	0.87	0.76	0.84	
Fe, Iron (wt.%)	2.89	2.80	2.98	2.84	2.94	
Ga, Gallium (ppm)	25.9	25.0	26.8	25.1	26.6	
Gd, Gadolinium (ppm)	5.14	4.47	5.81	4.97	5.31	
Hf, Hafnium (ppm)	4.79	4.56	5.03	4.55	5.03	
Ho, Holmium (ppm)	0.93	0.82	1.04	0.88	0.98	
In, Indium (ppm)	1.58	1.52	1.64	1.53	1.63	
K, Potassium (wt.%)	2.17	2.11	2.23	2.12	2.22	
La, Lanthanum (ppm)	77	73	81	73	80	
Li, Lithium (ppm)	81	78	83	79	83	
Lu, Lutetium (ppm)	0.39	0.34	0.43	0.36	0.41	
Mg, Magnesium (wt.%)	4.31	4.16	4.46	4.24	4.38	
Mo, Molybdenum (ppm)	10.8	10.2	11.3	10.4	11.1	
Na, Sodium (wt.%)	0.045	0.042	0.049	0.043	0.047	
Nb, Niobium (ppm)	15.4	13.1	17.7	14.5	16.3	
Nd, Neodymium (ppm)	61	55	67	59	63	
Ni, Nickel (ppm)	100	97	103	98	102	
P, Phosphorus (wt.%)	0.050	0.048	0.052	0.048	0.052	
Pb, Lead (ppm)	11.8	11.0	12.5	11.3	12.2	
Pr, Praseodymium (ppm)	17.0	15.9	18.1	16.4	17.6	
Rb, Rubidium (ppm)	82	79	86	79	85	

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

[†]The operationally defined measurand meets the requirements of ISO 17034 [9] and all participating laboratories comply with the requirements of ISO 17025 [8].

Note: intervals may appear asymmetric due to rounding.

Table 1 continued.					
Constituent	Certified	95 % Expand	ed Uncertainty	95 % Toler	ance Limits
Constituent	Value [†]	Low	High	Low	High
4-Acid Digestion contin	ued				
Re, Rhenium (ppm)	0.022	0.020	0.025	0.020	0.025
S, Sulphur (wt.%)	2.09	2.02	2.16	2.05	2.12
Sb, Antimony (ppm)	2.62	2.37	2.87	2.39	2.85
Sc, Scandium (ppm)	12.0	11.5	12.6	11.7	12.4
Se, Selenium (ppm)	4.96	4.44	5.48	4.50	5.42
Sm, Samarium (ppm)	8.33	7.56	9.10	7.91	8.76
Sn, Tin (ppm)	2.41	2.25	2.57	2.28	2.55
Sr, Strontium (ppm)	78	75	81	76	80
Ta, Tantalum (ppm)	1.13	0.94	1.32	1.05	1.21
Tb, Terbium (ppm)	0.77	0.70	0.84	0.74	0.80
Te, Tellurium (ppm)	0.066	0.041	0.091	IND	IND
Th, Thorium (ppm)	14.3	13.5	15.1	13.6	15.1
Ti, Titanium (wt.%)	0.242	0.208	0.276	0.233	0.250
TI, Thallium (ppm)	0.58	0.54	0.61	0.54	0.62
Tm, Thulium (ppm)	0.40	0.30	0.50	0.38	0.42
U, Uranium (ppm)	10.7	10.2	11.1	10.3	11.0
V, Vanadium (ppm)	256	248	264	250	262
W, Tungsten (ppm)	3.54	3.09	3.99	3.09	3.99
Y, Yttrium (ppm)	22.4	21.5	23.2	21.8	22.9
Yb, Ytterbium (ppm)	2.63	2.37	2.89	2.50	2.76
Zn, Zinc (ppm)	22.5	20.9	24.0	21.3	23.6
Zr, Zirconium (ppm)	171	164	178	167	174

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

[†]The operationally defined measurand meets the requirements of ISO 17034 [9] and all participating laboratories comply with the requirements of ISO 17025 [8].

IND = indeterminate (due to limited reading resolution of the methods employed).

Constituent	Certified	95 % Expande	ed Uncertainty	95 % Toler	ance Limits
Constituent	Value	Low	High	Low	High
Sulphuric Acid 5% Leach					
Co, Cobalt (wt.%)	0.077	0.073	0.082	0.075	0.079
Cu, Copper (wt.%)	1.12	1.07	1.16	1.09	1.15
Infrared Combustion					
C, Carbon (wt.%)	3.76	3.68	3.85	3.72	3.80
S, Sulphur (wt.%)	2.10	2.06	2.13	2.07	2.13
Borate Fusion XRF					1
Al ₂ O ₃ , Aluminium(III) oxide (wt.%)	11.93	11.71	12.15	11.82	12.04
BaO, Barium oxide (ppm)	372	278	466	IND	IND
CaO, Calcium oxide (wt.%)	1.17	1.15	1.19	1.15	1.19
Co, Cobalt (wt.%)	0.289	0.280	0.299	0.282	0.297
Cu, Copper (wt.%)	2.78	2.73	2.83	2.73	2.82
Fe ₂ O ₃ , Iron(III) oxide (wt.%)	4.19	4.14	4.25	4.15	4.24
K ₂ O, Potassium oxide (wt.%)	2.60	2.56	2.65	2.57	2.64
MgO, Magnesium oxide (wt.%)	7.16	7.08	7.25	7.09	7.23
MnO, Manganese oxide (wt.%)	0.024	0.020	0.028	0.022	0.027
Ni, Nickel (ppm)	101	47	154	IND	IND
P ₂ O ₅ , Phosphorus(V) oxide (wt.%)	0.119	0.108	0.129	0.114	0.123
SiO ₂ , Silicon dioxide (wt.%)	55.38	54.90	55.85	55.05	55.71
SO ₃ , Sulphur trioxide (wt.%)	5.27	5.11	5.43	5.19	5.35
SrO, Strontium oxide (ppm)	98	79	118	IND	IND
TiO ₂ , Titanium dioxide (wt.%)	0.739	0.716	0.762	0.722	0.757
V ₂ O ₅ , Vanadium(V) oxide (ppm)	452	350	553	IND	IND
Zr, Zirconium (ppm)	211	122	301	IND	IND
Thermogravimetry					I
LOI ¹⁰⁰⁰ , Loss On Ignition @1000°C (wt.%)	11.84	11.57	12.11	11.70	11.97
Peroxide Fusion ICP					1
Al, Aluminium (wt.%)	6.19	6.02	6.36	6.07	6.32
As, Arsenic (ppm)	49.9	45.6	54.3	48.7	51.2
B, Boron (ppm)	209	185	233	198	220
Ba, Barium (ppm)	337	326	348	324	350
Be, Beryllium (ppm)	4.85	4.34	5.37	4.36	5.34
Bi, Bismuth (ppm)	2.81	2.62	3.00	2.67	2.95
Ca, Calcium (wt.%)	0.834	0.783	0.886	0.795	0.873
Ce, Cerium (ppm)	155	149	162	149	161
Co, Cobalt (wt.%)	0.289	0.280	0.298	0.284	0.295
Cr, Chromium (ppm)	107	95	120	98	116
		+	l .		1

Table 2. Certified Values, Uncertainty & Tolerance Intervals for other measurands in OREAS 551b.

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).

		2 continued.	od Upcortainty	05 % Tolo	rance Limits
Constituent	Certified Value	-	ed Uncertainty		
Peroxide Fusion ICP continue		Low	High	Low	High
Cu, Copper (wt.%)	2.75	2.66	2.84	2.70	2.81
Dy, Dysprosium (ppm)	6.50	5.99	7.01	6.11	6.88
Er, Erbium (ppm)	3.49	3.17	3.80	3.26	3.71
Eu, Europium (ppm)	0.85	0.73	0.97	0.79	0.91
Fe, Iron (wt.%)	2.89	2.82	2.96	2.84	2.94
Ga, Gallium (ppm)	25.8	24.2	27.3	24.9	26.7
Gd, Gadolinium (ppm)	6.58	6.10	7.05	6.14	7.01
Ge, Germanium (ppm)	2.14	1.79	2.50	IND	IND
Ho, Holmium (ppm)	1.29	1.17	1.41	1.19	1.38
In, Indium (ppm)	1.64	1.50	1.77	IND	IND
K, Potassium (wt.%)	2.21	2.12	2.30	2.13	2.30
La, Lanthanum (ppm)	80	77	83	78	83
Li, Lithium (ppm)	81	76	86	78	84
Lu, Lutetium (ppm)	0.53	0.46	0.60	0.48	0.58
Mg, Magnesium (wt.%)	4.30	4.19	4.42	4.21	4.39
Mn, Manganese (wt.%)	0.020	0.019	0.021	0.019	0.021
Mo, Molybdenum (ppm)	11.1	9.4	12.7	IND	IND
Nb, Niobium (ppm)	24.1	22.2	26.0	22.6	25.6
Nd, Neodymium (ppm)	64	61	67	62	67
Ni, Nickel (ppm)	103	90	115	98	107
P, Phosphorus (wt.%)	0.052	0.038	0.067	IND	IND
Pb, Lead (ppm)	13.2	10.4	15.9	IND	IND
Pr, Praseodymium (ppm)	17.9	16.7	19.1	17.0	18.8
Rb, Rubidium (ppm)	82	78	86	79	84
Re, Rhenium (ppm)	< 0.1	IND	IND	IND	IND
S, Sulphur (wt.%)	2.09	2.01	2.16	2.02	2.15
Sb, Antimony (ppm)	2.03	2.01	3.15	2.02	3.08
Sc, Scandium (ppm)	11.5	10.5	12.5	IND	IND
Si, Silicon (wt.%)	26.08	25.26	26.91	25.70	26.46
Sm, Samarium (ppm)	8.92	8.47	9.37	8.48	9.36
Sr, Strontium (ppm)	78	73	84	75	82
Ta, Tantalum (ppm)	1.82	1.56	2.07	1.56	2.07
Tb, Terbium (ppm)	1.82	1.00	1.19	1.01	1.18
Th, Thorium (ppm)	1.10	14.0	1.19	14.2	1.18
			+ +		0.450
Ti, Titanium (wt.%)	0.435	0.420	0.450	0.419	
TI, Thallium (ppm)	0.62	0.52	0.72	IND	IND
Tm, Thulium (ppm)	0.53	0.47	0.60	0.49	0.58

SI unit equivalents: ppm (parts per million; 1×10^{-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).

	Certified	Certified 95 % Expanded Ur		95 % Tole	rance Limits
Constituent	Value	Low	High	Low	High
Peroxide Fusion ICP continue	d				L
U, Uranium (ppm)	10.8	10.2	11.4	10.4	11.2
V, Vanadium (ppm)	274	259	289	266	283
W, Tungsten (ppm)	4.09	3.03	5.14	IND	IND
Y, Yttrium (ppm)	32.8	31.6	34.0	31.2	34.3
Yb, Ytterbium (ppm)	3.42	3.12	3.73	3.23	3.62
Aqua Regia Digestion		·	· · ·		
Ag, Silver (ppm)	0.256	0.171	0.342	0.215	0.298
Al, Aluminium (wt.%)	1.31	1.23	1.39	1.26	1.36
As, Arsenic (ppm)	48.2	46.1	50.3	46.6	49.8
Au, Gold (ppm)	< 0.02	IND	IND	IND	IND
B, Boron (ppm)	< 10	IND	IND	IND	IND
Ba, Barium (ppm)	59	53	65	56	62
Be, Beryllium (ppm)	1.99	1.88	2.10	1.94	2.04
Bi, Bismuth (ppm)	2.47	2.36	2.57	2.39	2.54
Ca, Calcium (wt.%)	0.813	0.784	0.842	0.799	0.827
Cd, Cadmium (ppm)	0.31	0.28	0.33	0.29	0.33
Ce, Cerium (ppm)	41.1	37.7	44.5	39.2	43.0
Co, Cobalt (wt.%)	0.271	0.257	0.285	0.265	0.277
Cr, Chromium (ppm)	32.3	30.7	33.9	30.9	33.7
Cs, Caesium (ppm)	0.32	0.28	0.36	0.31	0.33
Cu, Copper (wt.%)	2.73	2.68	2.77	2.70	2.75
Dy, Dysprosium (ppm)	0.97	0.87	1.06	0.91	1.02
Er, Erbium (ppm)	0.44	0.41	0.47	0.43	0.45
Eu, Europium (ppm)	0.29	0.25	0.32	0.27	0.30
Fe, Iron (wt.%)	2.66	2.57	2.75	2.62	2.70
Ga, Gallium (ppm)	8.54	7.93	9.15	8.15	8.93
Gd, Gadolinium (ppm)	1.78	1.53	2.04	1.71	1.86
Hf, Hafnium (ppm)	0.32	0.29	0.34	0.30	0.33
Hg, Mercury (ppm)	0.058	0.044	0.071	IND	IND
Ho, Holmium (ppm)	0.17	0.14	0.20	0.16	0.18
In, Indium (ppm)	1.05	0.99	1.10	1.00	1.10
K, Potassium (wt.%)	0.194	0.182	0.205	0.181	0.206
La, Lanthanum (ppm)	18.7	17.3	20.1	18.0	19.4
Li, Lithium (ppm)	54	50	59	52	56
Lu, Lutetium (ppm)	0.052	0.045	0.059	0.040	0.064
Mg, Magnesium (wt.%)	3.57	3.44	3.69	3.50	3.64
Mn, Manganese (wt.%)	0.018	0.018	0.019	0.018	0.019

SI unit equivalents: ppm (parts per million; 1×10^{-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	95 % Expande	ed Uncertainty	95 % Tole	rance Limits	
Constituent	Value	Low	High	Low	High	
Aqua Regia Digestion continued	-				-	
Mo, Molybdenum (ppm)	10.1	9.7	10.5	9.8	10.4	
Na, Sodium (wt.%)	0.010	0.008	0.011	IND	IND	
Nd, Neodymium (ppm)	18.1	14.7	21.5	17.7	18.4	
Ni, Nickel (ppm)	93	89	97	91	96	
P, Phosphorus (wt.%)	0.036	0.034	0.037	0.035	0.037	
Pb, Lead (ppm)	8.29	7.85	8.73	8.04	8.54	
Pr, Praseodymium (ppm)	4.81	3.94	5.68	4.69	4.92	
Rb, Rubidium (ppm)	6.59	5.90	7.27	6.26	6.92	
Re, Rhenium (ppm)	0.021	0.018	0.023	0.017	0.025	
S, Sulphur (wt.%)	2.06	2.00	2.12	2.03	2.09	
Sb, Antimony (ppm)	1.62	1.46	1.79	1.52	1.72	
Sc, Scandium (ppm)	4.06	3.85	4.27	3.88	4.24	
Se, Selenium (ppm)	4.93	4.55	5.32	4.66	5.21	
Sm, Samarium (ppm)	2.86	2.45	3.27	2.78	2.94	
Sn, Tin (ppm)	1.09	1.00	1.18	1.03	1.15	
Sr, Strontium (ppm)	23.1	21.7	24.5	22.1	24.1	
Ta, Tantalum (ppm)	< 0.01	IND	IND	IND	IND	
Tb, Terbium (ppm)	0.20	0.19	0.22	0.20	0.21	
Te, Tellurium (ppm)	0.047	0.031	0.063	IND	IND	
Th, Thorium (ppm)	5.43	4.99	5.86	5.22	5.63	
U, Uranium (ppm)	4.99	4.76	5.23	4.86	5.12	
V, Vanadium (ppm)	56	52	59	54	58	
W, Tungsten (ppm)	1.59	1.29	1.88	1.38	1.79	
Y, Yttrium (ppm)	3.86	3.62	4.10	3.73	3.99	
Yb, Ytterbium (ppm)	0.36	0.29	0.42	0.34	0.38	
Zn, Zinc (ppm)	19.9	18.8	21.1	19.1	20.7	
Zr, Zirconium (ppm)	10.4	9.9	11.0	10.0	10.9	

Table 2 contin 4

SI unit equivalents: ppm (parts per million; 1×10^{-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)

			_					
Constituent	Unit	Value	Constituent	Unit	Value	Constituent	Unit	Value
4-Acid Digest	tion						· · · · · ·	
В	ppm	21.0	Hg	ppm	0.15			
Ge	ppm	0.25	Pt	ppb	7.50			
Borate Fusio	n XRF							
Ag	ppm	0.031	Hf	ppm	35.0	Se	ppm	3.33
As	ppm	90	Но	ppm	1.23	Sm	ppm	8.12
Be	ppm	4.33	In	ppm	1.57	Sn	ppm	< 50
Bi	ppm	< 100	La	ppm	84	Та	ppm	62
Cd	ppm	< 10	Lu	ppm	0.48	Tb	ppm	1.07
Ce	ppm	141	Мо	ppm	19.4	Те	ppm	0.13
CI	ppm	362	Na ₂ O	wt.%	0.068	Th	ppm	14.7
Cr ₂ O ₃	ppm	119	Nb	ppm	38.7	TI	ppm	0.50
Cs	ppm	1.78	Nd	ppm	62	Tm	ppm	0.50
Dy	ppm	6.52	Pb	ppm	46.7	U	ppm	62
Er	ppm	3.66	Pr	ppm	18.0	W	ppm	< 10
Eu	ppm	0.80	Rb	ppm	73	Y	ppm	43.6
F	ppm	< 5000	Re	ppm	< 0.1	Yb	ppm	3.62
Ga	ppm	25.0	Sb	ppm	< 50	Zn	ppm	< 50
Gd	ppm	6.34	Sc	ppm	13.5			
Peroxide Fus	ion ICP	continued						
Ag	ppm	1.09	Na	wt.%	0.059	Zn	ppm	23.3
Cd	ppm	0.41	Se	ppm	9.75	Zr	ppm	217
Hf	ppm	7.06	Sn	ppm	2.80			
Aqua Regia D	Digestio	n						
Ge	ppm	0.094	Pt	ppb	2.50	Tm	ppm	0.059
Nb	ppm	0.057	Ti	wt.%	0.002			
Pd	ppb	< 10	TI	ppm	0.23			
3-Acid Digest	tion (no	HF)						
As	ppm	< 10	Ni	ppm	97	V	ppm	87
Со	wt.%	0.278	Pb	ppm	135	Zn	ppm	82
Cu	wt.%	2.75	S	wt.%	2.09			

Table 3. Indicative Values for OREAS 551b.

SI unit equivalents: ppm (parts per million; 1×10^{-6}) \equiv mg/kg; wt.% (weight per cent) \equiv % (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 (generated from data supplied by laboratories all accredited to ISO 17025 for 4-acid digestion) and Table 2 (generated from data supplied by laboratories mostly 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 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 (PDM³) are presented in the detailed certification data for this CRM (**OREAS 551b-DataPack.1.0.240828_100502.xlsx**).

Results are also presented in scatter plots for Co and Cu by multiple operationally defined methods including 4-acid digestion with ICP-OES/MS finish (and/or AAS finish), borate fusion with XRF finish, peroxide fusion with ICP-OES/MS finish, and aqua regia digestion with ICP-OES/MS finish (and/or AAS finish) in Figures 1 to 8 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.

SOURCE MATERIAL

OREAS 551b was prepared from copper-cobalt sulphide ore samples sourced from MMG's Kinsevere Mine blended with barren black slate. The Kinsevere Mine is located in the Haut-Katanga province about 30 km from Lubumbashi in the south-east of the Democratic Republic of the Congo (DRC). The hypogene mineralisation at Kinsevere occurs as stratabound, veins and breccias consisting of mainly chalcopyrite, carrolite, bornite and occasionally pyrite and chalcocite, hosted within the Mine series carbonaceous shales, siltstones, and dolomites of the Roan Group belonging to the Katangan Supergroup stratigraphy.

COMMINUTION AND HOMOGENISATION PROCEDURES

The material constituting OREAS 551b was prepared in the following manner:

• Drying the ores and barren black slate to constant mass at 105 °C;

- Multi-stage milling of ores and barren black slate to achieve a particle size distribution of > 99.5 % passing 75 μm;
- Preliminary homogenisation of ore source materials;
- Representative sampling and check assaying of ore source materials;
- Blending the ores and barren black slate in appropriate proportions to achieve target grades;
- Homogenisation using OREAS' novel processing technologies;
- Packaging in 10 g units sealed in laminated foil pouches and 500 g units in plastic jars.

PHYSICAL PROPERTIES

OREAS 551b 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. Phy	sical properties	of OREAS 551b.
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Bulk Density (kg/m ³)	Moisture (wt.%)	Munsell Notation [‡]	Munsell Color [‡]
644	0.88	N3	Dark Gray

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

ANALYTICAL PROGRAM

Twenty-six commercial analytical laboratories participated in the program to certify the elements reported in Table 1 and 2. The following methods were employed:

- 4-acid (HNO₃-HF-HClO₄-HCl) digestion with full suite ICP-OES and ICP-MS elemental packages (up to 23 laboratories depending on the element);
- Aqua regia digestion for full elemental suite ICP-OES and ICP-MS (up to 26 laboratories depending on the element).
- Lithium borate fusion whole rock analysis package by X-ray fluorescence (up to 17 laboratories depending on the element);
- Thermogravimetry: Loss on Ignition (LOI) at 1000 °C (14 laboratories used a thermogravimetric analyser, 4 laboratories used a conventional muffle furnace and 3 laboratories included LOI with their fusion package);
- C and S by infrared combustion furnace/CS analyser (23 laboratories);
- Cu and Co by 5 % sulphuric acid leach* with ICP or AAS finish (up to 20 laboratories);
- Lithium borate or sodium peroxide fusion with full suite ICP-OES and ICP-MS elemental packages (up to 22 laboratories depending on the element).

*See 'Appendix' for specified methodology.

For the round robin program twelve 800 g test units were taken at predetermined intervals during the bagging stage, immediately following homogenisation and are considered representative of the entire prepared batch. The six samples received by each laboratory were obtained by taking two 30 g scoop splits from each of three separate 800 g test units. This format enabled a nested ANOVA treatment of the results to evaluate homogeneity, i.e.

to ascertain whether between-unit variance is greater than within-unit variance (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.

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 [5, 15]. 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.

Standard Deviation intervals (see Table 5, 'Performance Gates') 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.*

Homogeneity Evaluation

The tolerance limits (ISO 16269:2014) [6] 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 2.74 wt. % and 2.80 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. *Please note that tolerance limits pertain to the homogeneity of the CRM only and should not be used as control limits for laboratory performance.*

Based on the statistical analysis of the results of the interlaboratory certification program, it can be concluded that OREAS 551b is fit-for-purpose as a certified reference material (see 'Intended Use' below).

PERFORMANCE GATES

Table 5 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) \pm 10 %.

		1		enonina			JREAS 5	515.			
Ormatiturent	Certified		Absolute	Standard	Deviations	6	Relative	Standard D	eviations	5 % w	rindow
Constituent	Value	1SD	2SD Low	2SD High	3SD Low	3SD High	1RSD	2RSD	3RSD	Low	High
4-Acid Digest	ion										
Ag, ppm	0.302	0.057	0.188	0.417	0.131	0.474	18.94%	37.87%	56.81%	0.287	0.317
Al, wt.%	6.18	0.201	5.78	6.58	5.58	6.78	3.25%	6.51%	9.76%	5.87	6.49
As, ppm	50	2.1	46	55	44	57	4.27%	8.54%	12.81%	48	53
Ba, ppm	328	17	294	363	277	380	5.24%	10.48%	15.73%	312	345
Be, ppm	4.14	0.161	3.82	4.46	3.65	4.62	3.89%	7.79%	11.68%	3.93	4.34
Bi, ppm	2.64	0.131	2.38	2.90	2.25	3.03	4.97%	9.93%	14.90%	2.51	2.77
Ca, wt.%	0.835	0.017	0.801	0.869	0.784	0.886	2.04%	4.09%	6.13%	0.793	0.876
Cd, ppm	0.31	0.024	0.26	0.36	0.24	0.38	7.60%	15.19%	22.79%	0.29	0.33
Ce, ppm	145	12	121	168	109	180	8.20%	16.40%	24.60%	137	152
Co, wt.%	0.285	0.006	0.272	0.298	0.266	0.304	2.25%	4.50%	6.75%	0.271	0.299
Cr, ppm	79	8	63	96	54	105	10.58%	21.17%	31.75%	75	83
Cs, ppm	1.89	0.086	1.72	2.07	1.63	2.15	4.56%	9.12%	13.67%	1.80	1.99

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

Table 5. Performance Gates for OREAS 551b

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

Note 1: intervals may appear asymmetric due to rounding.

		Absolute Standard Deviations					Relative	Standard D	eviations	5 % window	
Constituent	Certified Value		-	1	1	I.	Ttelative			J 70 W	
	Value	1SD	2SD Low	2SD High	3SD Low	3SD High	1RSD	2RSD	3RSD	Low	High
4-Acid Digest	1	1	1							1	1
Cu, wt.%	2.77	0.048	2.67	2.86	2.62	2.91	1.73%	3.47%	5.20%	2.63	2.91
Dy, ppm	4.46	0.52	3.41	5.51	2.88	6.03	11.76%	23.53%	35.29%	4.23	4.68
Er, ppm	2.72	0.33	2.06	3.39	1.73	3.72	12.15%	24.30%	36.45%	2.59	2.86
Eu, ppm	0.80	0.047	0.71	0.90	0.66	0.94	5.82%	11.64%	17.45%	0.76	0.84
Fe, wt.%	2.89	0.079	2.73	3.05	2.66	3.13	2.72%	5.44%	8.17%	2.75	3.04
Ga, ppm	25.9	1.25	23.4	28.4	22.1	29.6	4.85%	9.70%	14.56%	24.6	27.2
Gd, ppm	5.14	0.68	3.78	6.50	3.10	7.18	13.25%	26.49%	39.74%	4.88	5.40
Hf, ppm	4.79	0.242	4.31	5.27	4.07	5.52	5.04%	10.08%	15.12%	4.55	5.03
Ho, ppm	0.93	0.081	0.77	1.09	0.69	1.17	8.69%	17.38%	26.08%	0.88	0.98
In, ppm	1.58	0.073	1.43	1.72	1.36	1.80	4.62%	9.23%	13.85%	1.50	1.66
K, wt.%	2.17	0.066	2.04	2.30	1.97	2.37	3.03%	6.06%	9.09%	2.06	2.28
La, ppm	77	5.1	67	87	62	92	6.67%	13.34%	20.01%	73	81
Li, ppm	81	2.4	76	86	74	88	2.98%	5.96%	8.93%	77	85
Lu, ppm	0.39	0.05	0.29	0.49	0.24	0.54	12.87%	25.75%	38.62%	0.37	0.41
Mg, wt.%	4.31	0.159	3.99	4.63	3.83	4.79	3.69%	7.39%	11.08%	4.10	4.53
Mn, wt.%	0.019	0.001	0.018	0.021	0.018	0.021	3.18%	6.35%	9.53%	0.018	0.020
Mo, ppm	10.8	0.50	9.8	11.8	9.3	12.3	4.65%	9.30%	13.95%	10.2	11.3
Na, wt.%	0.045	0.006	0.032	0.058	0.026	0.065	14.28%	28.57%	42.85%	0.043	0.048
Nb, ppm	15.4	3.0	9.5	21.4	6.5	24.3	19.28%	38.56%	57.85%	14.6	16.2
Nd, ppm	61	4.7	52	70	47	75	7.71%	15.41%	23.12%	58	64
Ni, ppm	100	2.8	94	105	92	108	2.77%	5.54%	8.32%	95	105
P, wt.%	0.050	0.003	0.045	0.055	0.042	0.058	5.21%	10.42%	15.63%	0.048	0.053
Pb, ppm	11.8	1.07	9.6	13.9	8.6	15.0	9.06%	18.13%	27.19%	11.2	12.3
Pr, ppm	17.0	0.95	15.1	18.9	14.1	19.8	5.60%	11.21%	16.81%	16.1	17.8
Rb, ppm	82	3.6	75	90	71	93	4.43%	8.87%	13.30%	78	86
Re, ppm	0.022	0.002	0.019	0.026	0.017	0.028	8.55%	17.09%	25.64%	0.021	0.023
S, wt.%	2.09	0.090	1.91	2.27	1.82	2.36	4.30%	8.60%	12.90%	1.98	2.19
Sb, ppm	2.62 12.0	0.32	1.98 10.7	3.27	1.65	3.60	12.35%	24.70%	37.05%	2.49	2.76
Sc, ppm	4.96	0.65		13.3	10.1	14.0	5.39% 7.20%	10.78% 14.40%	16.17% 21.60%	11.4	12.6
Se, ppm		0.357	4.25	5.68	3.89	6.03 10.13				4.71	5.21
Sm, ppm	8.33 2.41	0.600 0.168	7.13 2.08	9.53 2.75	6.53 1.91	2.92	7.20% 6.96%	14.41% 13.92%	21.61% 20.88%	7.92 2.29	8.75 2.53
Sn, ppm Sr, ppm	78	2.5	73	83	71	86	3.20%	6.39%	9.59%	74	82
Ta, ppm	1.13	0.18	0.77	1.49	0.58	1.68	16.11%	32.23%	48.34%	1.07	1.19
Tb, ppm	0.77	0.18	0.64	0.90	0.58	0.96	8.28%	16.56%	24.84%	0.73	0.81
Te, ppm	0.066	0.004	0.04	0.90	0.029	0.90	18.48%	36.96%	55.44%	0.73	0.069
Th, ppm	14.3	0.62	13.1	15.6	12.5	16.2	4.31%	8.62%	12.93%	13.6	15.1
Ti, wt.%	0.242	0.02	0.112	0.371	0.047	0.436	26.80%	53.59%	80.39%	0.230	0.254
TI, wt. %	0.242	0.003	0.112	0.65	0.047	0.430	5.91%	11.81%	17.72%	0.230	0.254
Tm, ppm	0.38	0.034	0.23	0.03	0.48	0.66	21.41%	42.82%	64.23%	0.38	0.42
U, ppm	10.7	0.03	9.7	11.7	9.2	12.2	4.76%	9.52%	14.27%	10.1	11.2
											269
											3.72
V, ppm W, ppm	256 3.54	7 0.47	243 2.61	269 4.47	236 2.14	276 4.94	2.63% 13.14%	5.25% 26.29%	7.88% 39.43%	243 3.36	

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

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

	Table 5 continued.										
Constituent	Certified		Absolute	Standard	Deviations	5	Relative	Standard D	eviations	5 % w	vindow
Constituent	Value	1SD	2SD Low	2SD High	3SD Low	3SD High	1RSD	2RSD	3RSD	Low	High
4-Acid Digest	ion continue	ed									
Y, ppm	22.4	0.99	20.4	24.3	19.4	25.3	4.44%	8.87%	13.31%	21.2	23.5
Yb, ppm	2.63	0.37	1.90	3.36	1.53	3.73	13.94%	27.88%	41.82%	2.50	2.76
Zn, ppm	22.5	1.38	19.7	25.2	18.3	26.6	6.14%	12.29%	18.43%	21.3	23.6
Zr, ppm	171	9	152	190	142	199	5.53%	11.06%	16.59%	162	179
Sulphuric Aci	d 5% Leach										
Co, wt.%	0.077	0.005	0.068	0.087	0.063	0.092	6.14%	12.27%	18.41%	0.074	0.081
Cu, wt.%	1.12	0.083	0.95	1.28	0.87	1.37	7.39%	14.79%	22.18%	1.06	1.18
Infrared Comb	oustion										
C, wt.%	3.76	0.163	3.44	4.09	3.28	4.25	4.32%	8.65%	12.97%	3.58	3.95
S, wt.%	2.10	0.063	1.97	2.22	1.91	2.28	2.98%	5.97%	8.95%	1.99	2.20
Borate Fusior	n XRF										
Al ₂ O ₃ , wt.%	11.93	0.186	11.56	12.30	11.37	12.49	1.56%	3.12%	4.68%	11.33	12.53
BaO, ppm	372	62	248	496	186	558	16.66%	33.32%	49.99%	353	391
CaO, wt.%	1.17	0.022	1.12	1.21	1.10	1.23	1.91%	3.81%	5.72%	1.11	1.23
Co, wt.%	0.289	0.009	0.271	0.307	0.262	0.316	3.11%	6.23%	9.34%	0.275	0.304
Cu, wt.%	2.78	0.045	2.69	2.87	2.64	2.91	1.63%	3.25%	4.88%	2.64	2.92
Fe ₂ O ₃ , wt.%	4.19	0.048	4.10	4.29	4.05	4.34	1.15%	2.30%	3.45%	3.98	4.40
K ₂ O, wt.%	2.60	0.057	2.49	2.72	2.43	2.77	2.20%	4.40%	6.59%	2.47	2.73
MgO, wt.%	7.16	0.143	6.88	7.45	6.73	7.59	1.99%	3.99%	5.98%	6.80	7.52
MnO, wt.%	0.024	0.003	0.017	0.031	0.014	0.035	14.19%	28.37%	42.56%	0.023	0.025
Ni, ppm	101	24	53	149	29	173	23.82%	47.64%	71.46%	96	106
P ₂ O ₅ , wt.%	0.119	0.008	0.102	0.135	0.093	0.144	7.13%	14.26%	21.40%	0.113	0.124
SiO ₂ , wt.%	55.38	0.455	54.47	56.29	54.01	56.74	0.82%	1.64%	2.47%	52.61	58.15
SO ₃ , wt.%	5.27	0.140	4.99	5.55	4.85	5.69	2.66%	5.33%	7.99%	5.01	5.54
SrO, ppm	98	19	60	137	40	157	19.77%	39.54%	59.31%	93	103
TiO ₂ , wt.%	0.739	0.021	0.697	0.781	0.677	0.802	2.82%	5.64%	8.46%	0.702	0.776
V ₂ O ₅ , ppm	452	94	263	641	168	735	20.91%	41.82%	62.72%	429	474
Zr, ppm	211	56	100	323	44	379	26.42%	52.85%	79.27%	201	222
Thermogravin	-			1	[[[[[
LOI ¹⁰⁰⁰ , wt.%	11.84	0.465	10.91	12.77	10.44	13.24	3.93%	7.86%	11.79%	11.25	12.43
Peroxide Fusi											
Al, wt.%	6.19	0.167	5.86	6.53	5.69	6.70	2.70%	5.41%	8.11%	5.88	6.50
As, ppm	49.9	4.50	40.9	58.9	36.4	63.4	9.02%	18.04%	27.06%	47.4	52.4
B, ppm	209	16	178	240	162	256	7.52%	15.04%	22.56%	199	219
Ba, ppm	337	11	314	359	303	370	3.32%	6.63%	9.95%	320	354
Be, ppm	4.85	0.52	3.82	5.88	3.30	6.40	10.63%	21.26%	31.89%	4.61	5.09
Bi, ppm	2.81	0.163	2.48	3.13	2.32	3.30	5.79%	11.57%	17.36%	2.67	2.95
Ca, wt.%	0.834	0.068	0.699	0.970	0.631	1.038	8.13%	16.25%	24.38%	0.793	0.876
Ce, ppm	155	5	145	165	140	170	3.28%	6.56%	9.83%	147	163
Co, wt.%	0.289	0.008	0.274	0.305	0.266	0.312	2.69%	5.38%	8.07%	0.275	0.304
Cr, ppm	107	21	64	150	43	172	20.05%	40.11%	60.16%	102	113
Cs, ppm	1.88	0.19	1.50	2.26	1.31	2.45	10.16%	20.32%	30.48%	1.78	1.97
Cu, wt.%	2.75	0.081	2.59	2.92	2.51	3.00	2.95%	5.90%	8.85%	2.62	2.89

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

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

Ormetiturent	Certified		Absolute	Standard	Deviation	S	Relative	Standard D	eviations	ns 5 % window		
Constituent	Value	1SD	2SD Low	2SD High	3SD Low	3SD High	1RSD	2RSD	3RSD	Low	High	
Peroxide Fus	ion ICP cont	tinued										
Dy, ppm	6.50	0.261	5.98	7.02	5.71	7.28	4.02%	8.03%	12.05%	6.17	6.82	
Er, ppm	3.49	0.37	2.74	4.23	2.37	4.61	10.70%	21.41%	32.11%	3.31	3.66	
Eu, ppm	0.85	0.09	0.67	1.03	0.58	1.12	10.47%	20.95%	31.42%	0.81	0.89	
Fe, wt.%	2.89	0.094	2.70	3.08	2.61	3.17	3.24%	6.48%	9.73%	2.75	3.03	
Ga, ppm	25.8	1.35	23.1	28.5	21.7	29.8	5.25%	10.50%	15.74%	24.5	27.1	
Gd, ppm	6.58	0.639	5.30	7.86	4.66	8.49	9.72%	19.45%	29.17%	6.25	6.91	
Ge, ppm	2.14	0.39	1.36	2.93	0.97	3.32	18.26%	36.51%	54.77%	2.04	2.25	
Ho, ppm	1.29	0.075	1.14	1.44	1.06	1.51	5.79%	11.58%	17.38%	1.22	1.35	
In, ppm	1.64	0.128	1.38	1.89	1.25	2.02	7.82%	15.63%	23.45%	1.56	1.72	
K, wt.%	2.21	0.116	1.98	2.45	1.87	2.56	5.23%	10.47%	15.70%	2.10	2.32	
La, ppm	80	3.8	73	88	69	91	4.69%	9.39%	14.08%	76	84	
Li, ppm	81	4.5	72	90	68	94	5.49%	10.98%	16.48%	77	85	
Lu, ppm	0.53	0.046	0.44	0.62	0.40	0.67	8.56%	17.11%	25.67%	0.51	0.56	
Mg, wt.%	4.30	0.160	3.98	4.62	3.82	4.78	3.73%	7.46%	11.19%	4.09	4.52	
Mn, wt.%	0.020	0.002	0.016	0.024	0.014	0.027	10.81%	21.61%	32.42%	0.019	0.021	
Mo, ppm	11.1	1.4	8.2	13.9	6.8	15.4	12.94%	25.88%	38.82%	10.5	11.6	
Nb, ppm	24.1	1.85	20.4	27.8	18.5	29.6	7.70%	15.41%	23.11%	22.9	25.3	
Nd, ppm	64	3.9	57	72	53	76	6.06%	12.12%	18.18%	61	68	
Ni, ppm	103	13	77	129	63	142	12.77%	25.53%	38.30%	98	108	
P, wt.%	0.052	0.008	0.036	0.069	0.028	0.077	15.32%	30.64%	45.96%	0.050	0.055	
Pb, ppm	13.2	1.7	9.8	16.6	8.1	18.3	12.92%	25.85%	38.77%	12.5	13.8	
Pr, ppm	17.9	1.25	15.4	20.4	14.1	21.7	7.01%	14.02%	21.04%	17.0	18.8	
Rb, ppm	82	3.3	75	88	72	91	4.01%	8.02%	12.03%	78	86	
Re, ppm	< 0.1	IND	IND	IND	IND	IND	IND	IND	IND	IND	IND	
S, wt.%	2.09	0.115	1.86	2.32	1.74	2.43	5.53%	11.05%	16.58%	1.98	2.19	
Sb, ppm	2.67	0.225	2.22	3.12	2.00	3.35	8.42%	16.85%	25.27%	2.54	2.80	
Sc, ppm	11.5	0.90	9.7	13.3	8.8	14.2	7.87%	15.74%	23.61%	10.9	12.1	
Si, wt.%	26.08	0.818	24.45	27.72	23.63	28.54	3.14%	6.27%	9.41%	24.78	27.39	
Sm, ppm	8.92	0.514	7.89	9.95	7.38	10.46	5.76%	11.52%	17.29%	8.47	9.37	
Sr, ppm	78	5.0	68	88	63	93	6.34%	12.67%	19.01%	74	82	
Ta, ppm	1.82	0.25	1.31	2.33	1.05	2.58	14.03%	28.07%	42.10%	1.73	1.91	
Tb, ppm	1.10	0.099	0.90	1.29	0.80	1.39	9.08%	18.16%	27.24%	1.04	1.15	
Th, ppm	14.6	0.73	13.2	16.1	12.5	16.8	4.96%	9.91%	14.87%	13.9	15.4	
Ti, wt.%	0.435	0.017	0.401	0.469	0.384	0.486	3.92%	7.85%	11.77%	0.413	0.457	
TI, ppm	0.62	0.08	0.47	0.77	0.39	0.85	12.39%	24.78%	37.17%	0.59	0.65	
Tm, ppm	0.53	0.048	0.44	0.63	0.39	0.68	9.03%	18.06%	27.08%	0.50	0.56	
U, ppm	10.8	0.71	9.4	12.2	8.7	12.9	6.60%	13.20%	19.80%	10.3	11.3	
V, ppm	274	14	247	302	233	316	5.04%	10.08%	15.12%	261	288	
W, ppm	4.09	0.69	2.70	5.47	2.00	6.17	16.98%	33.96%	50.94%	3.88	4.29	
Y, ppm	32.8	1.45	29.9	35.7	28.4	37.1	4.43%	8.86%	13.29%	31.2	34.4	
Yb, ppm	3.42	0.36	2.71	4.13	2.35	4.49	10.40%	20.79%	31.19%	3.25	3.59	

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

IND = indeterminate.

Note 1: intervals may appear asymmetric due to rounding.

	Certified		Absolute	Standard	Deviations	S	Relative Standard Deviations			5 % window	
Constituent	Value	1SD	2SD Low	2SD High	3SD Low	3SD High	1RSD	2RSD	3RSD	Low	High
Aqua Regia D	igestion										
Ag, ppm	0.256	0.074	0.109	0.403	0.036	0.477	28.71%	57.41%	86.12%	0.244	0.269
AI, wt.%	1.31	0.13	1.04	1.58	0.91	1.72	10.23%	20.46%	30.69%	1.25	1.38
As, ppm	48.2	2.81	42.6	53.8	39.8	56.6	5.84%	11.68%	17.52%	45.8	50.6
Au, ppm	< 0.02	IND	IND	IND	IND	IND	IND	IND	IND	IND	IND
B, ppm	< 10	IND	IND	IND	IND	IND	IND	IND	IND	IND	IND
Ba, ppm	59	8	43	75	35	83	13.35%	26.71%	40.06%	56	62
Be, ppm	1.99	0.169	1.65	2.33	1.48	2.50	8.51%	17.03%	25.54%	1.89	2.09
Bi, ppm	2.47	0.152	2.16	2.77	2.01	2.92	6.17%	12.33%	18.50%	2.34	2.59
Ca, wt.%	0.813	0.036	0.741	0.885	0.704	0.921	4.45%	8.89%	13.34%	0.772	0.854
Cd, ppm	0.31	0.022	0.26	0.35	0.24	0.37	7.08%	14.16%	21.24%	0.29	0.32
Ce, ppm	41.1	5.9	29.3	52.8	23.5	58.7	14.31%	28.61%	42.92%	39.0	43.1
Co, wt.%	0.271	0.027	0.216	0.326	0.189	0.353	10.12%	20.23%	30.35%	0.258	0.285
Cr, ppm	32.3	2.57	27.2	37.4	24.6	40.0	7.95%	15.90%	23.85%	30.7	33.9
Cs, ppm	0.32	0.07	0.18	0.46	0.11	0.53	21.78%	43.56%	65.34%	0.30	0.33
Cu, wt.%	2.73	0.072	2.58	2.87	2.51	2.94	2.64%	5.29%	7.93%	2.59	2.86
Dy, ppm	0.97	0.079	0.81	1.12	0.73	1.20	8.22%	16.43%	24.65%	0.92	1.01
Er, ppm	0.44	0.033	0.37	0.50	0.34	0.54	7.43%	14.85%	22.28%	0.42	0.46
Eu, ppm	0.29	0.026	0.24	0.34	0.21	0.36	9.00%	18.01%	27.01%	0.27	0.30
Fe, wt.%	2.66	0.106	2.45	2.87	2.35	2.98	3.97%	7.95%	11.92%	2.53	2.80
Ga, ppm	8.54	0.92	6.70	10.38	5.77	11.31	10.80%	21.59%	32.39%	8.11	8.97
Gd, ppm	1.78	0.23	1.32	2.25	1.09	2.48	13.03%	26.05%	39.08%	1.70	1.87
Hf, ppm	0.32	0.04	0.24	0.39	0.20	0.43	11.73%	23.46%	35.19%	0.30	0.33
Hg, ppm	0.058	0.011	0.035	0.080	0.024	0.092	19.40%	38.79%	58.19%	0.055	0.061
Ho, ppm	0.17	0.02	0.13	0.20	0.12	0.22	10.50%	21.00%	31.50%	0.16	0.18
In, ppm	1.05	0.059	0.93	1.17	0.87	1.22	5.61%	11.22%	16.83%	1.00	1.10
K, wt.%	0.194	0.019	0.155	0.233	0.135	0.252	10.03%	20.07%	30.10%	0.184	0.203
La, ppm	18.7	2.3	14.2	23.2	11.9	25.5	12.13%	24.26%	36.39%	17.8	19.6
Li, ppm	54	9	36	72	27	81	16.48%	32.95%	49.43%	51	57
Lu, ppm	0.052	0.004	0.044	0.060	0.040	0.064	7.69%	15.37%	23.06%	0.050	0.055
Mg, wt.%	3.57	0.217	3.13	4.00	2.92	4.22	6.07%	12.15%	18.22%	3.39	3.75
Mn, wt.%	0.018	0.001	0.017	0.020	0.016	0.021	4.94%	9.88%	14.82%	0.017	0.019
Mo, ppm	10.1	0.46	9.2	11.0	8.7	11.5	4.55%	9.10%	13.65%	9.6	10.6
Na, wt.%	0.010	0.002	0.006	0.013	0.005	0.014	16.53%	33.06%	49.59%	0.009	0.010
Nd, ppm	18.1	3.0	12.0	24.1	9.0	27.1	16.68%	33.37%	50.05%	17.2	19.0
Ni, ppm	93	5.5	82	104	77	110	5.92%	11.84%	17.76%	89	98
P, wt.%	0.036	0.003	0.030	0.041	0.028	0.044	7.70%	15.39%	23.09%	0.034	0.038
Pb, ppm	8.29	0.725	6.84	9.74	6.11	10.46	8.74%	17.49%	26.23%	7.88	8.70
Pr, ppm	4.81	0.76	3.30	6.32	2.54	7.07	15.71%	31.42%	47.13%	4.57	5.05
Rb, ppm	6.59	1.09	4.40	8.77	3.31	9.86	16.57%	33.14%	49.71%	6.26	6.92
Re, ppm	0.021	0.002	0.018	0.024	0.016	0.026	7.92%	15.84%	23.76%	0.020	0.022
S, wt.%	2.06	0.081	1.90	2.22	1.82	2.30	3.94%	7.89%	11.83%	1.96	2.16
Sb, ppm	1.62	0.23	1.17	2.08	0.94	2.30	14.02%	28.04%	42.06%	1.54	1.70
Sc, ppm	4.06	0.336	3.39	4.73	3.05	5.07	8.27%	16.54%	24.81%	3.86	4.26

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

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

Ormetiturent	Certified		Absolute	Standard	Deviations	6	Relative	Standard D	eviations	5 % w	vindow
Constituent	Value	1SD	2SD Low	2SD High	3SD Low	3SD High	1RSD	2RSD	3RSD	Low	High
Aqua Regia D	igestion co	ntinued									
Se, ppm	4.93	0.386	4.16	5.71	3.77	6.09	7.83%	15.67%	23.50%	4.69	5.18
Sm, ppm	2.86	0.36	2.14	3.58	1.77	3.94	12.63%	25.26%	37.88%	2.71	3.00
Sn, ppm	1.09	0.071	0.95	1.23	0.88	1.30	6.52%	13.05%	19.57%	1.04	1.15
Sr, ppm	23.1	2.6	17.8	28.4	15.2	31.0	11.40%	22.79%	34.19%	21.9	24.3
Ta, ppm	< 0.01	IND	IND	IND	IND	IND	IND	IND	IND	IND	IND
Tb, ppm	0.20	0.019	0.16	0.24	0.15	0.26	9.51%	19.02%	28.52%	0.19	0.21
Te, ppm	0.047	0.013	0.021	0.072	0.009	0.084	27.16%	54.32%	81.47%	0.044	0.049
Th, ppm	5.43	0.59	4.25	6.61	3.66	7.20	10.88%	21.76%	32.64%	5.16	5.70
U, ppm	4.99	0.334	4.32	5.66	3.99	5.99	6.70%	13.39%	20.09%	4.74	5.24
V, ppm	56	7	42	69	36	76	11.96%	23.93%	35.89%	53	59
W, ppm	1.59	0.29	1.01	2.16	0.73	2.44	18.03%	36.06%	54.10%	1.51	1.67
Y, ppm	3.86	0.42	3.02	4.70	2.60	5.13	10.92%	21.84%	32.75%	3.67	4.05
Yb, ppm	0.36	0.04	0.28	0.44	0.24	0.47	11.21%	22.42%	33.62%	0.34	0.37
Zn, ppm	19.9	1.44	17.0	22.8	15.6	24.2	7.23%	14.45%	21.68%	18.9	20.9
Zr, ppm	10.4	0.99	8.5	12.4	7.5	13.4	9.46%	18.93%	28.39%	9.9	11.0

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

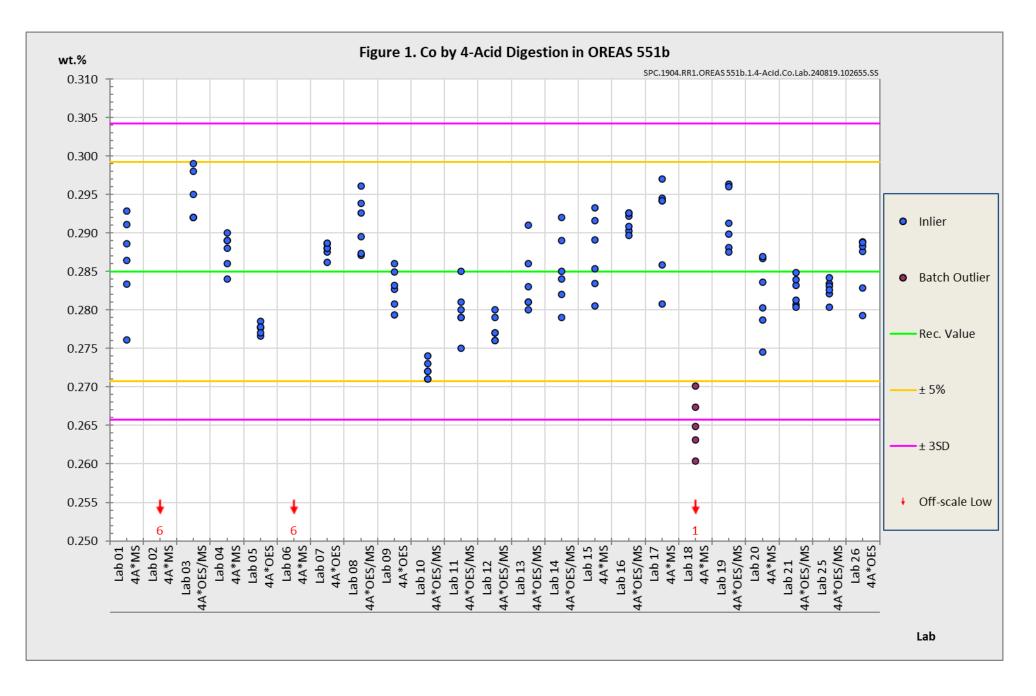
Note 1: intervals may appear asymmetric due to rounding; IND = indeterminate.

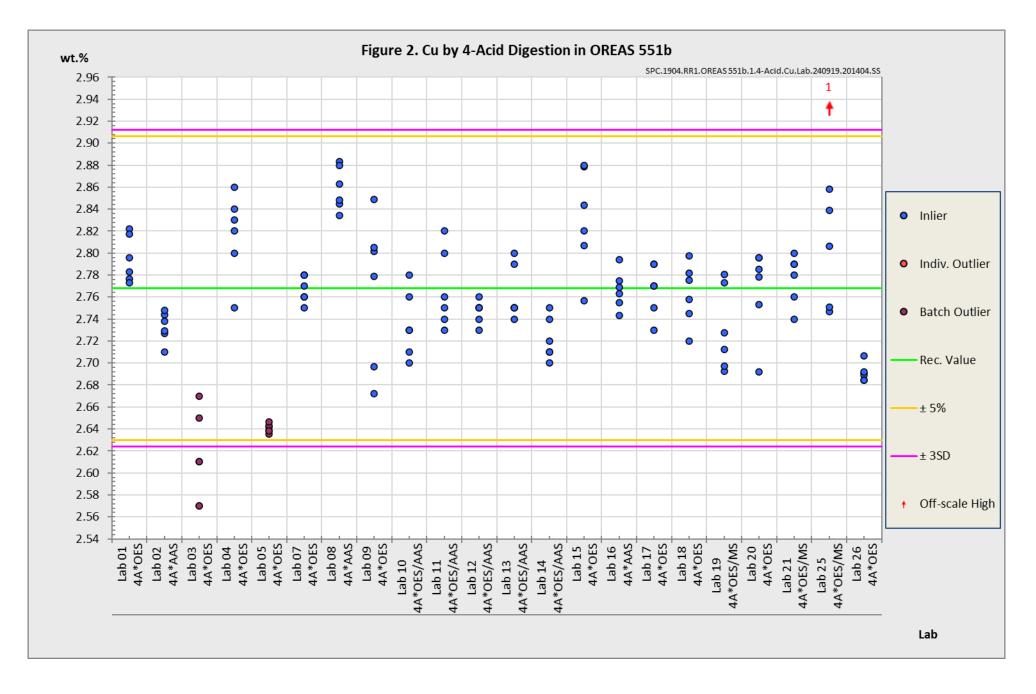
Note 2: the number of decimal places quoted does not imply accuracy of the certified value to this level but are given to minimise rounding errors when calculating 2SD and 3SD windows.

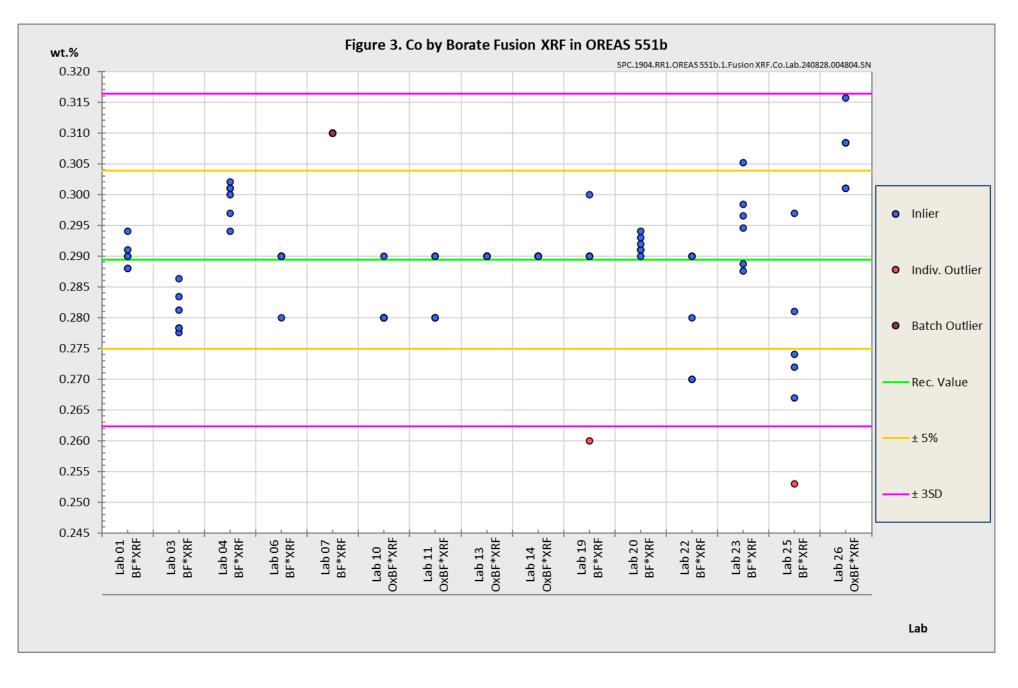
PARTICIPATING LABORATORIES

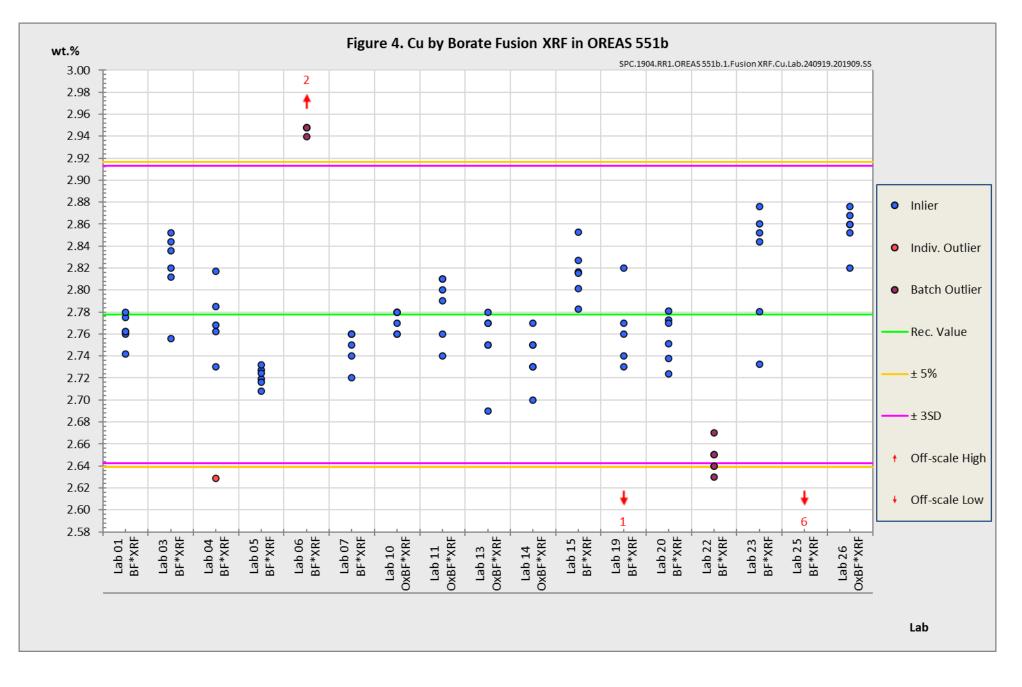
- 1. Actlabs, Ancaster, Ontario, Canada
- 2. ALS, Brisbane, QLD, Australia
- 3. ALS, Lima, Peru
- 4. ALS, Loughrea, Galway, Ireland
- 5. ALS, Malaga, WA, Australia
- 6. ALS, Vancouver, BC, Canada
- 7. American Assay Laboratories, Sparks, Nevada, USA
- 8. ARGETEST Mineral Processing, Ankara, Central Anatolia, Turkey
- 9. Bureau Veritas Commodities Canada Ltd, Vancouver, BC, Canada
- 10. CERTIMIN, Lima, Peru
- 11. ESAN Istanbul, Istanbul, Turkey
- 12. Inspectorate (BV), Lima, Peru
- 13. Intertek, Cupang, Muntinlupa, Philippines
- 14. Intertek, Perth, WA, Australia
- 15. Intertek, Townsville, QLD, Australia
- 16. Labwest Minerals Analysis, Perth, WA, Australia
- 17. PT Geoservices Ltd, Cikarang, Jakarta Raya, Indonesia
- 18. PT Intertek Utama Services, Jakarta Timur, DKI Jakarta, Indonesia
- 19. Saskatchewan Research Council, Saskatoon, Saskatchewan, Canada
- 20. SGS, Randfontein, Gauteng, South Africa
- 21. SGS Australia Mineral Services, Perth, WA, Australia
- 22. SGS Canada Inc., Vancouver, BC, Canada
- 23. SGS del Peru, Lima, Peru
- 24. Shiva Analyticals Ltd, Bangalore North, Karnataka, India
- 25. Skyline Assayers & Laboratories, Tucson, Arizona, USA
- 26. Stewart Assay & Environmental Laboratories LLC, Kara-Balta, Chüy, Kyrgyzstan

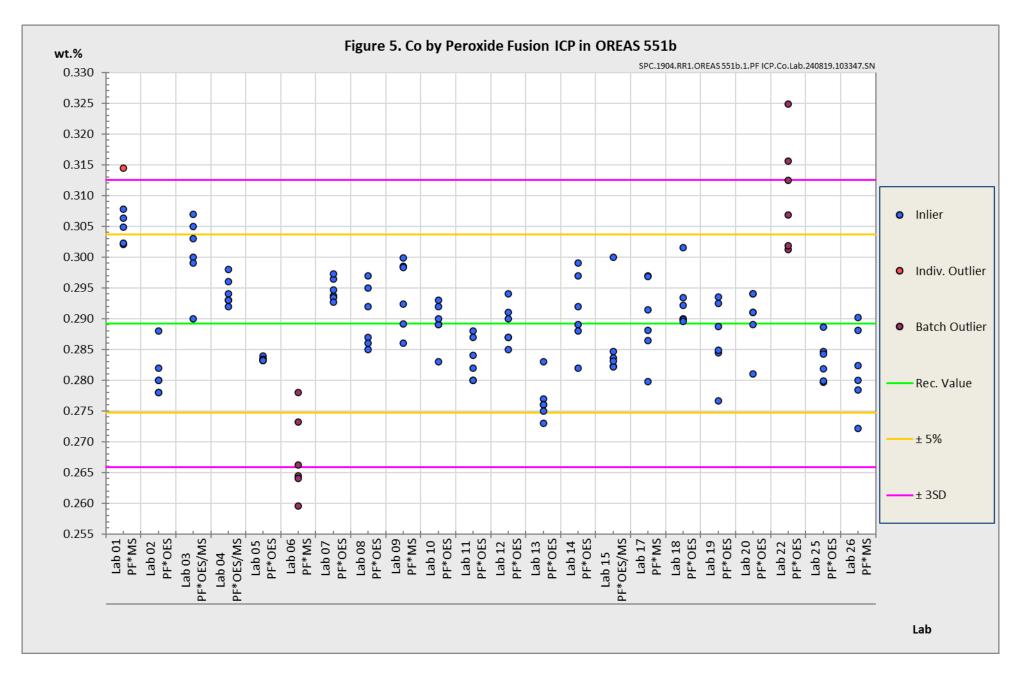
Please note: To preserve anonymity, the above numbered alphabetical list of participating laboratories <u>does not</u> correspond with the Lab ID numbering on the scatter plots below.

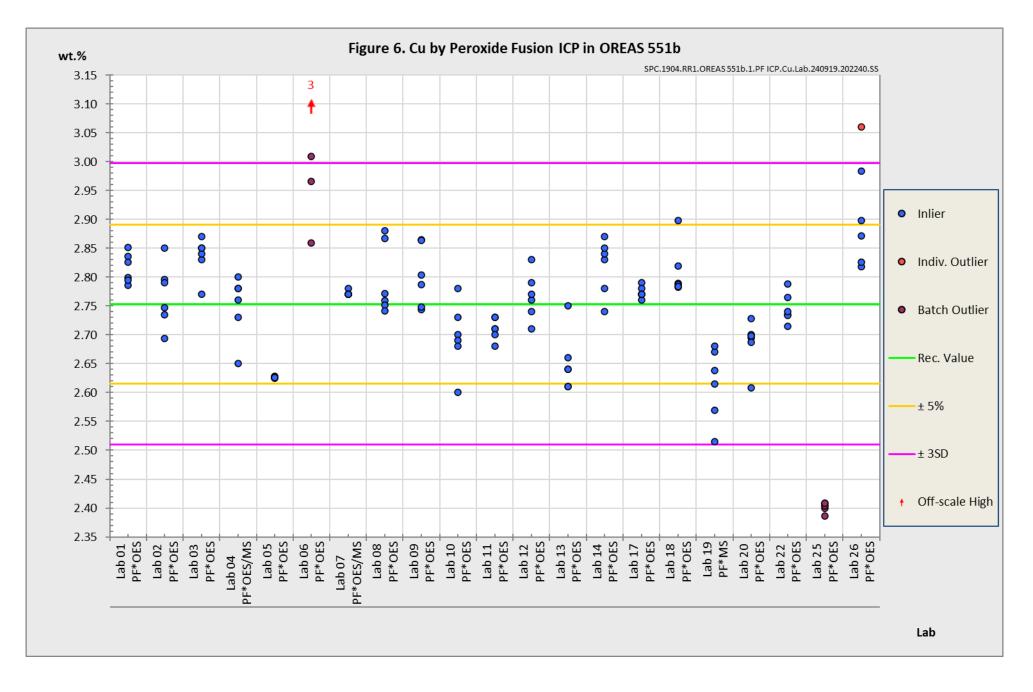


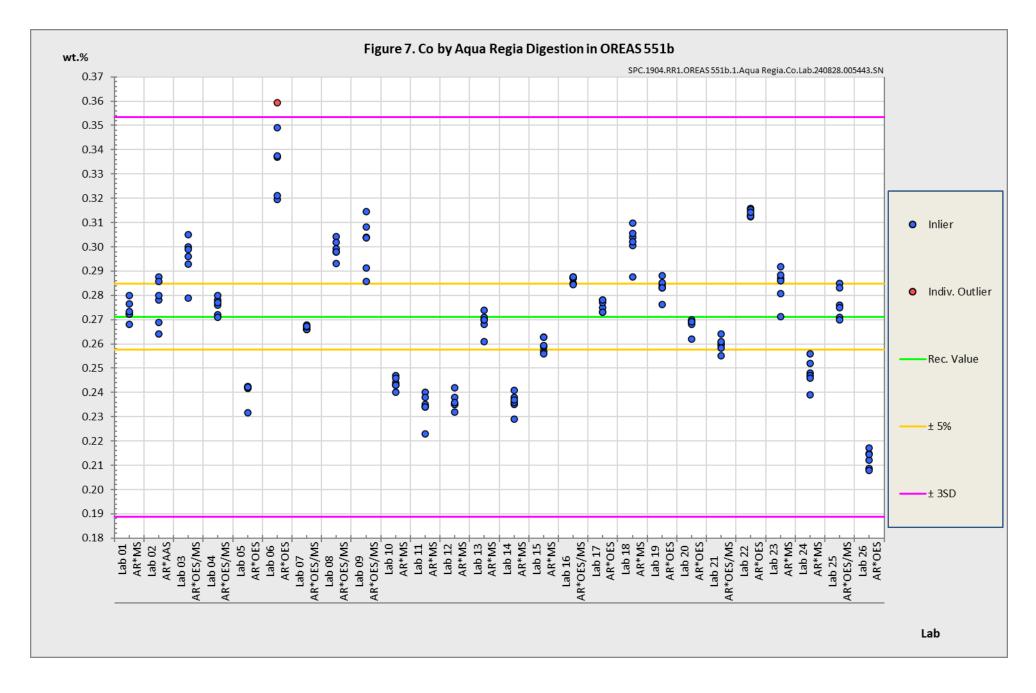


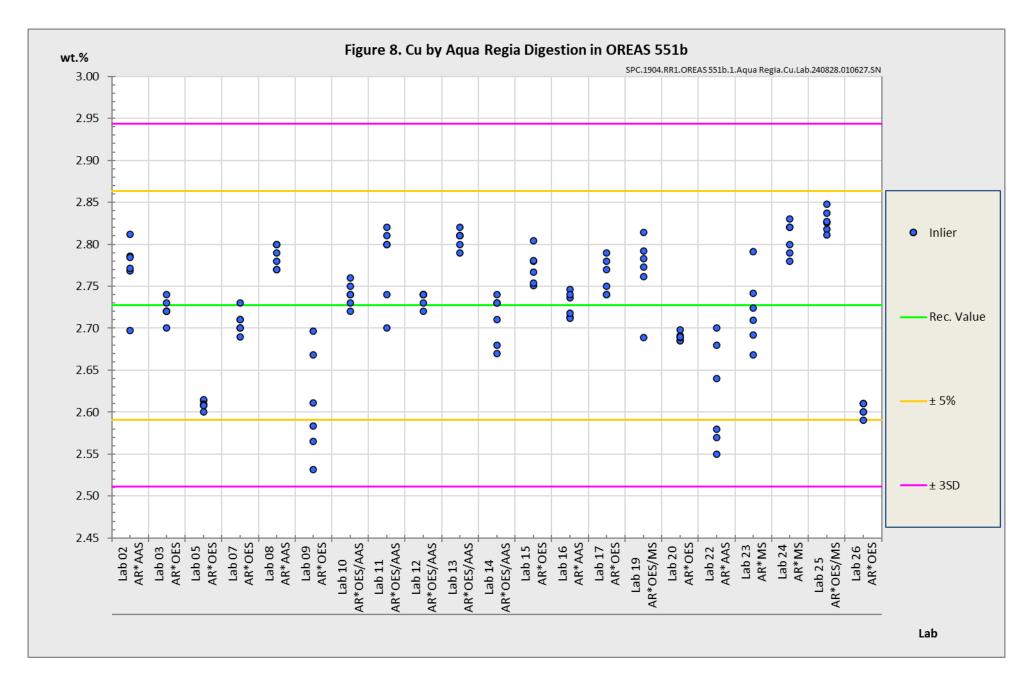












PREPARER AND SUPPLIER

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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)) [14]. 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 representativeness 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 [8], 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 4-acid digestion (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 [7], 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) [4]." 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 551b 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 551b may be used to calibrate the entire procedure by producing a pure substance CRM transformed into a calibration solution.

OREAS 551b 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:

- 4-acid digestion with ICP-OES and/or MS finish: ≥ 0.25 g;
- Cu and Co by 5 % sulphuric acid leach with ICP or AAS finish: 0.5 g;
- C and S by infrared combustion furnace/CS analyser: ≥ 0.1 g;
- Lithium borate fusion with X-ray fluorescence finish: ≥ 0.2 g;
- Loss on Ignition (LOI) at 1000 °C: \geq 1 g;
- Sodium peroxide fusion with ICP-OES and/or MS finish: ≥ 0.2 g;
- Aqua regia digestion with ICP-OES and/or MS finish: ≥ 0.5 g.



PERIOD OF VALIDITY & STORAGE INSTRUCTIONS

The certification of OREAS 551b remains valid, within the specified measurement uncertainties, until at least March 2039, 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

Following analysis of the CRM subsample it is the manufacturers' expectation that any remaining material is discarded. The stability of the material after opening the sachet is not within the scope of proper use. However, if opened sachets are resealed after opening, then under ordinary* storage conditions the CRM will have a shelf-life beyond ten years.

*ordinary storage conditions: means storage not in direct sunlight in a dry, clean, wellventilated area at temperatures between -5 °C and 50 °C.

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 [12].

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 5 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.

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CERTIFYING OFFICER

12th November, 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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APPENDIX

For Cu and Co by sulphuric acid leach, a specific methodology was detailed for the participating laboratories to follow:

5 % sulphuric acid leach

- 1. Weigh 0.500 ± 0.002 g of sample pulp into a clean 250 mL flask.
- 2. Add to the flask 0.5 g of Sodium Sulphite (AR Grade).
- 3. Add 50 mL of approximately 55 g/L Sulphuric acid solution (prepared from a 98 % pure concentrated sulphuric acid).
- 4. Put the cap on the flask and start automatic shaking. Leave the sample on continuous shaking to leach for four (4) hours.
- 5. Remove the cap and add 25 mL of hydrochloric acid.
- 6. Dilute to a final volume of 250 mL with distilled/deionised water up to the mark and mix again by inverting at least 10 times. Allow the solution to settle for 30 minutes.

The solution is now ready to be analysed by ICP or AAS

