

CERTIFICATE OF ANALYSIS FOR

OREAS 554b

Copper-Cobalt Ore (Kinsevere Mine, Katanga Province, Democratic Republic of the Congo)





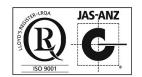


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Table 1. Certified Values, Uncertainty & Tolerance Intervals for elements by 4-Acid Digestion and Aqua Regia Digestion in OREAS 554b.

•	Certified		ed Uncertainty	95% Tolerance Limits		
Constituent	Value [†]	Low	High	Low	High	
4-Acid Digestion						
Ag, Silver (ppm)	0.135	0.103	0.168	0.123	0.148	
Al, Aluminium (wt.%)	6.47	6.26	6.67	6.34	6.59	
As, Arsenic (ppm)	75	73	78	72	78	
Ba, Barium (ppm)	586	543	630	564	609	
Be, Beryllium (ppm)	4.17	3.99	4.36	4.06	4.28	
Bi, Bismuth (ppm)	1.29	1.22	1.35	1.25	1.32	
Ca, Calcium (wt.%)	0.114	0.106	0.122	0.111	0.117	
Cd, Cadmium (ppm)	0.38	0.34	0.42	0.36	0.41	
Ce, Cerium (ppm)	134	129	138	130	138	
Co, Cobalt (wt.%)	0.788	0.766	0.810	0.777	0.799	
Cr, Chromium (ppm)	93	89	98	90	97	
Cs, Caesium (ppm)	2.44	2.34	2.55	2.36	2.53	
Cu, Copper (wt.%)	3.70	3.62	3.77	3.63	3.76	
Dy, Dysprosium (ppm)	4.48	3.96	5.00	4.26	4.70	
Er, Erbium (ppm)	2.69	2.31	3.06	2.52	2.86	
Eu, Europium (ppm)	0.97	0.91	1.03	0.92	1.02	
Fe, Iron (wt.%)	3.66	3.59	3.73	3.59	3.72	
Ga, Gallium (ppm)	26.5	25.4	27.6	25.8	27.3	
Gd, Gadolinium (ppm)	5.08	4.41	5.74	4.86	5.29	
Ge, Germanium (ppm)	0.18	0.14	0.23	0.17	0.20	
Hf, Hafnium (ppm)	4.75	4.55	4.94	4.56	4.94	
Ho, Holmium (ppm)	0.88	0.79	0.98	0.84	0.93	
In, Indium (ppm)	2.08	2.00	2.15	2.01	2.14	
K, Potassium (wt.%)	2.57	2.50	2.63	2.52	2.62	
La, Lanthanum (ppm)	64	61	67	62	66	
Li, Lithium (ppm)	52	50	53	51	53	
Lu, Lutetium (ppm)	0.41	0.39	0.42	0.38	0.44	
Mg, Magnesium (wt.%)	2.46	2.39	2.54	2.42	2.51	
Mn, Manganese (wt.%)	0.065	0.063	0.066	0.063	0.066	
Mo, Molybdenum (ppm)	14.7	14.0	15.4	14.3	15.2	
Na, Sodium (wt.%)	0.063	0.058	0.068	0.061	0.065	
Nb, Niobium (ppm)	15.0	13.3	16.6	14.3	15.6	
Nd, Neodymium (ppm)	58	54	62	56	60	
Ni, Nickel (ppm)	151	147	155	148	154	
P, Phosphorus (wt.%)	0.033	0.032	0.035	0.032	0.034	
Pb, Lead (ppm)	15.9	15.1	16.6	15.3	16.5	

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.



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

Table 1 Continued.										
Constituent	Certified	•	ed Uncertainty	95% Tolerance Limits						
	Value [†]	Low	High	Low	High					
4-Acid Digestion continue			T							
Pr, Praseodymium (ppm)	15.5	14.3	16.6	14.9	16.0					
Rb, Rubidium (ppm)	106	101	112	103	110					
Re, Rhenium (ppm)	0.024	0.021	0.027	0.022	0.026					
S, Sulphur (wt.%)	3.57	3.48	3.65	3.49	3.64					
Sb, Antimony (ppm)	3.79	3.58	3.99	3.61	3.97					
Sc, Scandium (ppm)	12.4	11.8	12.9	12.0	12.7					
Se, Selenium (ppm)	5.99	5.40	6.59	5.66	6.33					
Sm, Samarium (ppm)	8.29	7.87	8.70	7.96	8.62					
Sn, Tin (ppm)	3.21	3.05	3.37	3.07	3.35					
Sr, Strontium (ppm)	66	64	68	64	67					
Ta, Tantalum (ppm)	1.05	0.92	1.18	0.98	1.11					
Tb, Terbium (ppm)	0.77	0.64	0.91	0.73	0.82					
Te, Tellurium (ppm)	0.078	0.058	0.098	IND	IND					
Th, Thorium (ppm)	15.4	14.6	16.3	14.9	15.9					
Ti, Titanium (wt.%)	0.275	0.241	0.310	0.266	0.285					
TI, Thallium (ppm)	0.98	0.93	1.03	0.95	1.02					
Tm, Thulium (ppm)	0.39	0.33	0.44	0.36	0.41					
U, Uranium (ppm)	8.26	7.86	8.66	8.00	8.51					
V, Vanadium (ppm)	316	307	324	309	323					
W, Tungsten (ppm)	2.63	2.49	2.78	2.49	2.78					
Y, Yttrium (ppm)	22.8	21.6	24.0	22.2	23.4					
Yb, Ytterbium (ppm)	2.70	2.44	2.96	2.58	2.81					
Zn, Zinc (ppm)	31.6	30.3	33.0	30.2	33.0					
Zr, Zirconium (ppm)	165	161	170	162	169					
Aqua Regia Digestion										
Ag, Silver (ppm)	0.088	0.076	0.101	IND	IND					
Al, Aluminium (wt.%)	0.868	0.779	0.956	0.835	0.901					
As, Arsenic (ppm)	72	70	75	71	74					
Au, Gold (ppm)	< 0.02	IND	IND	IND	IND					
B, Boron (ppm)	< 10	IND	IND	IND	IND					
Ba, Barium (ppm)	81	71	91	78	84					
Be, Beryllium (ppm)	1.84	1.71	1.96	1.80	1.87					
Bi, Bismuth (ppm)	1.19	1.14	1.24	1.15	1.22					
Ca, Calcium (wt.%)	0.108	0.103	0.113	0.105	0.111					
Cd, Cadmium (ppm)	0.38	0.35	0.42	0.35	0.41					
Ce, Cerium (ppm)	46.3	42.2	50.3	45.0	47.5					

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; 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).



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

Table 1 Continued.									
Constituent	Certified	95% Expande	ed Uncertainty	95% Tolerance Limits					
Consuluent	Value [†]	Low	High	Low	High				
Aqua Regia Digestion cor	ntinued								
Co, Cobalt (wt.%)	0.759	0.733	0.785	0.748	0.770				
Cr, Chromium (ppm)	33.3	31.0	35.5	31.9	34.6				
Cs, Caesium (ppm)	0.43	0.36	0.50	0.41	0.45				
Cu, Copper (wt.%)	3.74	3.68	3.79	3.68	3.79				
Dy, Dysprosium (ppm)	1.10	0.86	1.33	1.05	1.15				
Er, Erbium (ppm)	0.46	0.33	0.58	0.43	0.49				
Eu, Europium (ppm)	0.43	0.29	0.56	0.41	0.44				
Fe, Iron (wt.%)	3.51	3.40	3.63	3.45	3.58				
Ga, Gallium (ppm)	7.13	6.48	7.78	6.91	7.35				
Gd, Gadolinium (ppm)	2.31	2.06	2.56	2.11	2.50				
Ge, Germanium (ppm)	0.093	0.060	0.127	IND	IND				
Hf, Hafnium (ppm)	0.33	0.30	0.36	0.31	0.35				
Hg, Mercury (ppm)	0.054	0.039	0.070	IND	IND				
Ho, Holmium (ppm)	0.18	0.14	0.22	IND	IND				
In, Indium (ppm)	1.47	1.41	1.53	1.44	1.51				
K, Potassium (wt.%)	0.203	0.186	0.220	0.193	0.214				
La, Lanthanum (ppm)	20.9	19.4	22.5	20.2	21.7				
Li, Lithium (ppm)	27.7	24.7	30.7	26.7	28.7				
Mg, Magnesium (wt.%)	1.88	1.77	1.98	1.83	1.92				
Mn, Manganese (wt.%)	0.064	0.062	0.066	0.063	0.065				
Mo, Molybdenum (ppm)	14.0	13.4	14.5	13.6	14.3				
Na, Sodium (wt.%)	0.010	0.009	0.012	0.010	0.011				
Nb, Niobium (ppm)	0.083	0.066	0.100	IND	IND				
Nd, Neodymium (ppm)	21.1	15.0	27.3	20.4	21.9				
Ni, Nickel (ppm)	147	142	152	144	150				
P, Phosphorus (wt.%)	0.020	0.020	0.021	0.019	0.021				
Pb, Lead (ppm)	11.8	11.0	12.6	11.5	12.1				
Pr, Praseodymium (ppm)	5.46	3.85	7.07	5.24	5.67				
Rb, Rubidium (ppm)	8.01	7.20	8.82	7.68	8.34				
Re, Rhenium (ppm)	0.021	0.019	0.024	0.019	0.023				
S, Sulphur (wt.%)	3.48	3.38	3.58	3.42	3.54				
Sb, Antimony (ppm)	2.22	2.02	2.42	2.12	2.31				
Sc, Scandium (ppm)	2.91	2.71	3.11	2.75	3.07				
Se, Selenium (ppm)	5.95	5.35	6.56	5.61	6.29				
Sm, Samarium (ppm)	3.72	2.88	4.56	3.53	3.91				
Sn, Tin (ppm)	1.31	1.21	1.41	1.27	1.35				

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



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

Constituent	Certified	95% Expande	d Uncertainty	95% Tolerance Limits						
Constituent	Value [†]	Low	High	Low	High					
Aqua Regia Digestion continued										
Sr, Strontium (ppm)	15.8	13.8	17.8	15.1	16.5					
Ta, Tantalum (ppm)	< 0.01	IND	IND	IND	IND					
Tb, Terbium (ppm)	0.26	0.21	0.30	0.24	0.28					
Te, Tellurium (ppm)	0.057	0.047	0.068	IND	IND					
Th, Thorium (ppm)	6.93	6.33	7.54	6.70	7.17					
U, Uranium (ppm)	3.22	3.04	3.39	3.12	3.31					
V, Vanadium (ppm)	52	48	56	50	54					
W, Tungsten (ppm)	0.91	0.83	1.00	0.86	0.97					
Y, Yttrium (ppm)	4.37	4.04	4.70	4.24	4.50					
Yb, Ytterbium (ppm)	0.37	0.25	0.48	0.33	0.41					
Zn, Zinc (ppm)	28.2	27.1	29.3	27.2	29.2					
Zr, Zirconium (ppm)	10.4	9.7	11.2	10.1	10.8					

SI unit equivalents: ppm (parts per million; 1×10^{-6}) \equiv mg/kg. Note: intervals may appear asymmetric due to rounding. [†]This operationally defined measurand meets the requirements of ISO 17034 and all participating laboratories comply with the requirements of ISO 17025.

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. Certified Value, Uncertainty & Tolerance Intervals for other measurands in OREAS 554b.

Constituent	Certified	95% Expand	ed Uncertainty	95% Tolerance Limits		
Constituent	Value	Low	High	Low	High	
Borate Fusion XRF						
Al ₂ O ₃ , Aluminium(III) oxide (wt.%)	12.54	12.40	12.68	12.41	12.68	
BaO, Barium oxide (ppm)	707	630	784	682	732	
CaO, Calcium oxide (wt.%)	0.158	0.151	0.165	IND	IND	
Co, Cobalt (wt.%)	0.794	0.767	0.820	0.783	0.805	
Cu, Copper (wt.%)	3.67	3.60	3.75	3.64	3.71	
Fe ₂ O ₃ , Iron(III) oxide (wt.%)	5.30	5.22	5.39	5.26	5.35	
K₂O, Potassium oxide (wt.%)	3.13	3.06	3.20	3.10	3.16	
MgO, Magnesium oxide (wt.%)	4.15	4.10	4.21	4.10	4.20	
MnO, Manganese oxide (wt.%)	0.085	0.082	0.088	0.083	0.087	
Ni, Nickel (ppm)	159	148	170	IND	IND	
P ₂ O ₅ , Phosphorus(V) oxide (wt.%)	0.079	0.075	0.084	0.077	0.082	
SiO ₂ , Silicon dioxide (wt.%)	56.28	55.88	56.68	55.90	56.66	
SO ₃ , Sulphur trioxide (wt.%)	8.91	8.56	9.26	8.81	9.01	
Sr, Strontium (ppm)	66	52	79	IND	IND	
TiO ₂ , Titanium dioxide (wt.%)	0.747	0.729	0.766	0.731	0.764	
V ₂ O ₅ , Vanadium(V) oxide (ppm)	558	490	626	519	598	
Zr, Zirconium (ppm)	207	197	217	IND	IND	

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

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

Note: intervals may appear asymmetric due to rounding; IND = indeterminate (due to limited reading resolution of the methods employed).



Table 2 continued.									
Constituent	Certified		ed Uncertainty						
	Value	Low	High	Low	High				
Thermogravimetry		1			I				
LOI ¹⁰⁰⁰ , Loss On Ignition @1000°C (wt.%)	10.98	10.78	11.19	10.89	11.08				
Infrared Combustion									
C, Carbon (wt.%)	3.77	3.71	3.83	3.73	3.81				
S, Sulphur (wt.%)	3.60	3.54	3.65	3.55	3.64				
Sulphuric Acid 5% Leach									
Co, Cobalt (wt.%)	0.269	0.235	0.302	0.262	0.276				
Cu, Copper (wt.%)	0.948	0.916	0.981	0.935	0.962				
Peroxide Fusion ICP									
Al, Aluminium (wt.%)	6.54	6.43	6.65	6.41	6.67				
As, Arsenic (ppm)	75	68	83	73	78				
B, Boron (ppm)	209	177	242	197	222				
Ba, Barium (ppm)	650	623	676	633	666				
Be, Beryllium (ppm)	4.79	4.27	5.30	IND	IND				
Bi, Bismuth (ppm)	1.41	1.29	1.53	IND	IND				
Ca, Calcium (wt.%)	0.115	0.093	0.138	0.103	0.128				
Ce, Cerium (ppm)	148	139	156	141	154				
Co, Cobalt (wt.%)	0.795	0.774	0.815	0.780	0.809				
Cr, Chromium (ppm)	112	101	123	107	117				
Cs, Caesium (ppm)	2.62	2.29	2.96	2.46	2.79				
Cu, Copper (wt.%)	3.73	3.66	3.79	3.64	3.81				
Dy, Dysprosium (ppm)	6.54	6.01	7.08	6.11	6.97				
Er, Erbium (ppm)	3.73	3.39	4.07	3.44	4.02				
Eu, Europium (ppm)	1.10	1.02	1.18	1.05	1.15				
Fe, Iron (wt.%)	3.69	3.60	3.78	3.64	3.75				
Ga, Gallium (ppm)	27.0	25.4	28.7	25.3	28.7				
Gd, Gadolinium (ppm)	7.06	6.26	7.87	6.64	7.49				
Ge, Germanium (ppm)	2.28	1.63	2.93	2.00	2.56				
Ho, Holmium (ppm)	1.28	1.15	1.41	1.20	1.36				
In, Indium (ppm)	2.04	1.89	2.19	1.88	2.21				
K, Potassium (wt.%)	2.58	2.50	2.65	2.51	2.64				
La, Lanthanum (ppm)	73	70	76	70	76				
Li, Lithium (ppm)	54	50	57	52	55				
Lu, Lutetium (ppm)	0.55	0.42	0.67	0.49	0.60				
Mg, Magnesium (wt.%)	2.50	2.44	2.56	2.45	2.55				
Mn, Manganese (wt.%)	0.067	0.064	0.070	0.066	0.069				
Mo, Molybdenum (ppm)	15.0	13.7	16.3	14.4	15.7				
Nb, Niobium (ppm)	22.9	20.5	25.2	21.3	24.4				
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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; IND = indeterminate (due to limited reading resolution of the methods employed).



0 + i + +	Certified	95% Expande	ed Uncertainty	95% Tolerance Limits		
Constituent	Value	Low	High	Low	High	
Peroxide Fusion ICP cont	inued					
Nd, Neodymium (ppm)	61	56	66	58	64	
Ni, Nickel (ppm)	157	147	167	151	163	
P, Phosphorus (wt.%)	0.032	0.024	0.041	IND	IND	
Pb, Lead (ppm)	21.0	17.4	24.6	17.9	24.1	
Pr, Praseodymium (ppm)	17.0	15.7	18.4	16.1	18.0	
Rb, Rubidium (ppm)	107	101	113	103	111	
S, Sulphur (wt.%)	3.60	3.51	3.70	3.53	3.68	
Sb, Antimony (ppm)	3.96	3.49	4.44	IND	IND	
Sc, Scandium (ppm)	11.4	10.1	12.6	IND	IND	
Si, Silicon (wt.%)	26.68	26.11	27.25	26.13	27.22	
Sm, Samarium (ppm)	8.83	8.10	9.56	8.11	9.56	
Sn, Tin (ppm)	3.10	2.41	3.78	IND	IND	
Sr, Strontium (ppm)	73	68	77	70	76	
Ta, Tantalum (ppm)	1.77	1.45	2.09	IND	IND	
Tb, Terbium (ppm)	1.14	1.00	1.27	1.05	1.22	
Th, Thorium (ppm)	15.6	14.9	16.2	15.0	16.1	
Ti, Titanium (wt.%)	0.436	0.422	0.450	0.426	0.445	
Tl, Thallium (ppm)	1.03	0.88	1.18	IND	IND	
Tm, Thulium (ppm)	0.53	0.49	0.56	0.49	0.56	
U, Uranium (ppm)	8.62	8.00	9.24	8.23	9.01	
V, Vanadium (ppm)	333	318	348	325	340	
W, Tungsten (ppm)	3.10	1.89	4.31	IND	IND	
Y, Yttrium (ppm)	34.8	33.1	36.5	33.1	36.5	
Yb, Ytterbium (ppm)	3.51	3.14	3.88	3.19	3.83	
Zn, Zinc (ppm)	33.5	28.2	38.8	31.5	35.6	

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

Table 3. Indicative Values for OREAS 554b.

		Tabi	e 3. Indicative	Values	IOI OILAG 3	J-D.		
Constituent	Unit	Value	Constituent	Unit	Value	Constituent	Unit	Value
Borate Fusion	on XRF	continued						
Ag	ppm	0.040	HfO ₂	ppm	312	Sc	ppm	13.6
As	ppm	78	Hg	ppm	< 100	Se	ppm	2.33
Be	ppm	2.17	Но	ppm	1.18	Sm	ppm	7.74
Bi	ppm	< 100	In	ppm	2.38	Sn	ppm	57
Cd	ppm	< 100	La	ppm	73	Та	ppm	1.90
Се	ppm	115	Lu	ppm	0.48	Tb	ppm	1.09
Cl	ppm	171	Мо	ppm	< 50	Te	ppm	11.9
Cr ₂ O ₃	ppm	152	Na ₂ O	wt.%	0.102	Th	ppm	15.3
Cs	ppm	2.63	Nb	ppm	28.8	TI	ppm	0.79
Dy	ppm	5.78	Nd	ppm	54	Tm	ppm	0.50
Er	ppm	3.41	Pb	ppm	< 100	U	ppm	8.27
Eu	ppm	0.94	Pr	ppm	14.8	W	ppm	7.22
Ga	ppm	26.8	Rb	ppm	88	Y	ppm	34.7
Gd	ppm	6.13	Re	ppm	18.9	Yb	ppm	3.28
Ge	ppm	< 100	Sb	ppm	< 50	Zn	ppm	43.9
Peroxide Fu)						
Ag	ppm	0.386	Hg	ppm	< 0.1	Se	ppm	10.3
Cd	ppm	< 10	Na	wt.%	0.057	Te	ppm	< 2
Hf	ppm	5.79	Re	ppm	0.052	Zr	ppm	214
Ag	ppm	0.386	Hg	ppm	< 0.1	Se	ppm	10.3
4-Acid Diges			<u> </u>					
Hg	ppm	< 1	Pt	ppb	9.33			
Aqua Regia		on		''				
Ir	ppm	< 0.003	Pt	ppb	5.81	Tm	ppm	0.065
Lu	ppm	0.066	Ti	wt.%	0.002			
Pd	ppb	< 10	TI	ppm	0.33			
3-Acid Diges								
Ag	ppm	0.407	Gd	ppm	5.67	S	wt.%	3.40
Al ₂ O ₃	wt.%	12.37	Hf	ppm	5.32	Sc	ppm	13.0
Ва	ppm	635	Но	ppm	0.88	Sm	ppm	7.88
Be	ppm	5.30	K ₂ O	wt.%	3.15	Sn	ppm	3.18
Bi	ppm	1.82	La	ppm	72	Sr	ppm	69
CaO	wt.%	0.152	Li	ppm	54	Та	ppm	0.32
Cd	ppm	0.50	MgO	wt.%	4.15	Tb	ppm	0.72
Ce	ppm	130	MnO	wt.%	0.081	Th	ppm	17.4
Co	wt.%	0.858	Мо	ppm	16.8	TiO ₂	wt.%	0.739
Cr	ppm	103	Na ₂ O	wt.%	0.082	U	ppm	9.04
Cs	ppm	2.43	Nb	ppm	20.8	V	ppm	332
Cu	wt.%	3.97	Nd	ppm	59	W	ppm	3.53
Dy	ppm	4.74	Ni	ppm	181	Y	ppm	26.9
Er	ppm	2.77	P ₂ O ₅	wt.%	< 0.002	Yb	ppm	2.75
Eu	ppm	1.11	Pb	ppm	19.0	Zn	ppm	73
Fe ₂ O ₃	wt.%	5.19	Pr	ppm	15.8	Zr	ppm	179
			· 1 v 10-9\ = ug/			l .		l .

SI unit equivalents: ppb (parts per billion; 1 x 10^{-9}) $\equiv \mu g/kg$; ppm (parts per million; 1 x 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 and 2 provides the certified values and their associated 95% expanded uncertainty and tolerance intervals, Table 3 shows indicative values, Table 4 provides some indicative physical properties, Table 5 provides indicative mineralogy based on semi-quantitative XRD analysis and Table 6 presents the performance gate intervals for all certified values. 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 554b-DataPack.1.0.240216_184819.xlsx).

Results are also presented in scatter plots for Co and Cu by multiple operationally defined methods including borate fusion with XRF finish, peroxide fusion with ICP-OES/MS finish, 4-acid digestion with ICP-OES/MS finish (and/or AAS finish) and aqua regia digestion with ICP-OES/MS finish (and/or AAS finish) in Figures 1 to 8 respectively, together with ±3SD (magenta) and ±5% (yellow) control lines and certified value (green line). Accepted individual results are coloured blue and individual and dataset outliers are identified in red and violet, respectively.

SOURCE MATERIAL

OREAS 554b was prepared from copper-cobalt sulphide ore samples sourced from MMG's Kinsevere Mine blended with barren black slate and a minor addition of cobalt concentrate. 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 554b was prepared in the following manner:

- Drying of ore and barren materials to constant mass at 105°C;
- Drying of cobalt concentrate to constant mass at 95°C;
- Crushing and milling of the ore materials to 100% minus 30 microns;
- Crushing and milling of the barren materials to >98% minus 75 microns;



- Deagglomeration of cobalt concentrate and screening at 150 microns;
- Check analysis of ores and concentrate for contained Co and Cu concentrations;
- Blending ores, Co concentrate and barren materials in appropriate proportions to achieve the desired grades;
- Homogenisation using OREAS' novel processing technologies;
- Packaging in 10g units sealed under nitrogen in laminated foil pouches.

PHYSICAL PROPERTIES

OREAS 554b was tested at ORE Research & Exploration Pty Ltd's onsite facility for various physical properties. Table 4 presents these findings that should be used for informational purposes only.

Table 4. Physical properties of OREAS 554b.

Bulk Density (kg/m³)	Bulk Density (kg/m³) Moisture (wt.%)		Munsell Color‡
730	0.84	N2	GrayishBlack

[‡]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.

MINERALOGY

The semi-quantitative XRD results shown in Table 5 below have been normalised to 100 % and represent the relative proportion of crystalline material. Totals greater or less than 100 % are due to rounding errors. Some amorphous material might be present. A trace amount of plagioclase appears to be present. The quantification of the kandite group is performed on a largely overlapped pattern and carriers high than usual uncertainty. A trace amount of arsenopyrite, malachite and talc may also be present.

Table 5. Indicative mineralogy of OREAS 554b based on semi-quantitative XRD analysis.

Mineral / Mineral Group	% (mass ratio)
Chlorite	3
Kandite group	2
Annite - biotite - phlogopite	4
Muscovite	17
K-feldspar and/or rutile	0
Tourmaline	< 1
Quartz	42
Dolomite - ankerite	1
Pyrite	0
Chalcopyrite	6
Linnaeite group	1
Magnetite and/or antlerite	1
Goethite	< 1
Malachite	0
Magnesite	22
Rutile	1

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ANALYTICAL PROGRAM

Thirty analytical laboratories participated in the program to characterise the elements reported in Tables 1 and 2. The following elements and methods were undertaken:

- Lithium borate fusion whole rock analysis package by X-ray fluorescence (up to 20 laboratories depending on the element; except for one laboratory that used pressed powder pellet with XRF);
- Loss on Ignition (LOI) at 1000° C (14 laboratories used a thermogravimetric analyser. 6 laboratories included LOI with their fusion package and 4 laboratories used a conventional muffle furnace);
- Infrared combustion furnace/CS analyser to determine C (25 laboratories) and S (26 laboratories);
- Co (9 laboratories) and Cu (19 laboratories) by *5% sulphuric acid leach with ICP-OES or AAS finish:
- Full ICP-OES and ICP-MS elemental suites by sodium peroxide fusion (up to 22 laboratories depending on the element);
- Full ICP-OES and ICP-MS elemental suites by 4-acid (HCI-HNO₃-HF-HClO₄) digestion (up to 27 laboratories depending on the element);
- Full ICP-OES and ICP-MS elemental suites by aqua regia digestion (up to 27 laboratories depending on the element).

For the round robin program, ten 800g test units were taken at predetermined intervals immediately following homogenisation and are considered representative of the entire prepared batch. The six samples received by each laboratory were obtained by taking two 30g scoop splits from each of three separate 800g 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.

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^{*}See 'Appendix' for specified methodology.

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]. 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.

Standard Deviation intervals (see Table 5) provide an indication of a level of performance that might reasonably be expected from a laboratory being monitored by this CRM in a QA/QC program. They take into account errors attributable to measurement uncertainty and CRM variability. For an effective CRM the contribution of the latter should be negligible in comparison to measurement errors. The Standard Deviation values include all sources of measurement uncertainty: between-lab variance, within-run variance (precision errors) and CRM variability.

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

The SD for each analyte's certified value is calculated from the same filtered data set used to determine the certified value, i.e., after removal of all individual, lab dataset (batch) and 3SD outliers (single iteration). These outliers can only be removed after the absolute homogeneity of the CRM has been independently established, i.e., the outliers must be confidently deemed to be analytical rather than arising from inhomogeneity of the CRM. The standard deviation is then calculated for each analyte from the pooled accepted analyses generated from the certification program.

Homogeneity Evaluation

The tolerance limits (ISO 16269:2014) shown in Tables 1 and 2 were determined using an analysis of precision errors method and are considered a conservative estimate of true homogeneity. The meaning of tolerance limits may be illustrated for copper by fusion with XRF, where 99% of the time (1- α =0.99) at least 95% of subsamples (ρ =0.95) will have concentrations lying between 3.64 and 3.71 wt.%. Put more precisely, this means that if the same number of subsamples were taken and analysed in the same manner repeatedly, 99% of the tolerance intervals so constructed would cover at least 95% of the total population, and 1% of the tolerance intervals would cover less than 95% of the total population (ISO Guide 35). *Please note that tolerance limits pertain to the homogeneity of the CRM only and should not be used as control limits for laboratory performance.*

The homogeneity of OREAS 554b has also been evaluated in an ANOVA study for all certified analytes present in concentrations well above detection levels (i.e. >20 x Lower Limit of Detection) for the various methods undertaken. This study tests the null hypothesis that no statistically significant difference exists between the between-unit variance and the within-unit variance (i.e. *p*-values <0.05 indicate rejection of the null hypothesis). Of the 195 certified values, no failures were observed indicating no evidence to reject the null hypothesis.

Based on the statistical analysis of ANOVA and the results of the interlaboratory certification program, it can be concluded that OREAS 554b is fit-for-purpose as a certified reference material (see 'Intended Use' below).

PERFORMANCE GATES

The standard deviations (SD's) intervals reported in Table 6 provide an indication of a level of performance that might reasonably be expected from a laboratory being monitored by this CRM in a QA/QC program. They take into account errors attributable to measurement uncertainty and CRM variability. For an effective CRM the contribution of the latter should be negligible in comparison to measurement errors. The Standard Deviation values include all sources of measurement uncertainty: between-lab variance, within-run variance (precision errors) and CRM variability.

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

The standard deviation is then calculated for each analyte from the pooled accepted analyses generated from the certification program.

Table 6 below shows intervals calculated for two and three standard deviations. As a guide these intervals may be regarded as warning or rejection for multiple 2SD outliers, or rejection for individual 3SD outliers in QC monitoring, although their precise application should be at the discretion of the QC manager concerned (also see 'Intended Use' section below). Westgard Rules extend the basics of single-rule QC monitoring using multi-rules (for more information visit www.westgard.com/mltirule.htm). A second method utilises a 5% window calculated directly from the certified value. Standard deviation is also shown in relative percent for one, two and three relative standard deviations (1RSD, 2RSD and 3RSD) to facilitate an appreciation of the magnitude of these numbers and a comparison with the 5% window. Caution should be exercised when concentration levels approach lower limits of detection of the analytical methods employed as performance gates calculated from standard deviations tend to be excessively wide whereas those determined by the 5% method are too narrow. One approach used at commercial laboratories is to set the acceptance criteria at twice the detection level (DL) ± 10%.

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

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Table 6. Performance Gates for OREAS 554b.

Table 6. Performance Gates for OREAS 5540.											
Constituent	Certified		Absolute				Relative	Standard D	eviations	5% w	indow
Constituent	Value	1SD	2SD Low	2SD High	3SD Low	3SD High	1RSD	2RSD	3RSD	Low	High
Borate Fusion	XRF		LOW	riigii	LOW	riigii					
Al ₂ O ₃ , wt.%	12.54	0.196	12.15	12.94	11.96	13.13	1.56%	3.13%	4.69%	11.92	13.17
BaO, ppm	707	93	522	893	429	985	13.11%	26.21%	39.32%	672	743
CaO, wt.%	0.158	0.005	0.148	0.168	0.143	0.173	3.10%	6.21%	9.31%	0.150	0.166
Co, wt.%	0.794	0.041	0.711	0.876	0.670	0.918	5.21%	10.41%	15.62%	0.754	0.833
Cu, wt.%	3.67	0.110	3.45	3.89	3.34	4.00	3.01%	6.01%	9.02%	3.49	3.86
Fe ₂ O ₃ , wt.%	5.30	0.125	5.05	5.55	4.93	5.68	2.36%	4.72%	7.08%	5.04	5.57
K ₂ O, wt.%	3.13	0.121	2.89	3.37	2.77	3.49	3.85%	7.70%	11.55%	2.97	3.29
MgO, wt.%	4.15	0.078	4.00	4.31	3.92	4.38	1.87%	3.74%	5.62%	3.94	4.36
MnO, wt.%	0.085	0.006	0.074	0.096	0.068	0.101	6.48%	12.97%	19.45%	0.081	0.089
Ni, ppm	159	17	124	194	106	211	11.00%	22.01%	33.01%	151	167
P ₂ O ₅ , wt.%	0.079	0.006	0.067	0.092	0.061	0.098	7.74%	15.49%	23.23%	0.075	0.083
SiO ₂ , wt.%	56.28	0.485	55.31	57.25	54.83	57.74	0.86%	1.72%	2.58%	53.47	59.10
SO ₃ , wt.%	8.91	0.430	8.05	9.77	7.62	10.20	4.83%	9.66%	14.49%	8.47	9.36
Sr, ppm	66	15	35	96	20	111	23.00%	46.00%	68.99%	62	69
TiO ₂ , wt.%	0.747	0.024	0.700	0.795	0.676	0.819	3.19%	6.38%	9.58%	0.710	0.785
V ₂ O ₅ , ppm	558	88	383	733	296	821	15.68%	31.36%	47.04%	530	586
Zr, ppm	207	8	191	224	182	232	3.98%	7.97%	11.95%	197	217
Thermogravin						-					
LOI ¹⁰⁰⁰ , wt.%	10.98	0.352	10.28	11.69	9.93	12.04	3.20%	6.41%	9.61%	10.44	11.53
Infrared Comb											
C, wt.%	3.77	0.114	3.54	4.00	3.43	4.11	3.02%	6.03%	9.05%	3.58	3.96
S, wt.%	3.60	0.076	3.44	3.75	3.37	3.82	2.11%	4.22%	6.32%	3.42	3.78
Sulphuric Acid											
Co, wt.%	0.269	0.025	0.219	0.319	0.194	0.344	9.32%	18.64%	27.97%	0.255	0.282
Cu, wt.%	0.948	0.048	0.853	1.044	0.806	1.091	5.01%	10.02%	15.04%	0.901	0.996
Peroxide Fusi	on ICP										
Al, wt.%	6.54	0.149	6.24	6.84	6.09	6.99	2.28%	4.56%	6.84%	6.21	6.87
As, ppm	75	7.0	62	89	55	96	9.24%	18.48%	27.71%	72	79
B, ppm	209	35	140	279	105	314	16.66%	33.32%	49.99%	199	220
Ba, ppm	650	37	575	725	537	762	5.76%	11.52%	17.28%	617	682
Be, ppm	4.79	0.334	4.12	5.46	3.78	5.79	6.98%	13.96%	20.94%	4.55	5.03
Bi, ppm	1.41	0.084	1.24	1.57	1.15	1.66	5.96%	11.92%	17.88%	1.34	1.48
Ca, wt.%	0.115	0.027	0.062	0.169	0.035	0.196	23.31%	46.62%	69.93%	0.110	0.121
Ce, ppm	148	4	141	155	137	158	2.38%	4.75%	7.13%	140	155
Co, wt.%	0.795	0.022	0.751	0.839	0.729	0.861	2.77%	5.54%	8.31%	0.755	0.834
Cr, ppm	112	22	69	155	47	177	19.39%	38.78%	58.16%	106	118
Cs, ppm	2.62	0.159	2.31	2.94	2.15	3.10	6.04%	12.09%	18.13%	2.49	2.76
Cu, wt.%	3.73	0.092	3.54	3.91	3.45	4.00	2.47%	4.94%	7.42%	3.54	3.91
Dy, ppm	6.54	0.390	5.76	7.32	5.37	7.71	5.97%	11.94%	17.90%	6.21	6.87
Er, ppm	3.73	0.174	3.38	4.08	3.21	4.26	4.67%	9.35%	14.02%	3.55	3.92
Eu, ppm	1.10	0.082	0.93	1.26	0.85	1.35	7.49%	14.97%	22.46%	1.04	1.15
Fe, wt.%	3.69	0.135	3.42	3.96	3.29	4.10	3.66%	7.31%	10.97%	3.51	3.88
Ga, ppm	27.0	1.52	24.0	30.1	22.5	31.6	5.62%	11.25%	16.87%	25.7	28.4
Gd, ppm	7.06	0.450	6.16	7.96	5.72	8.41	6.36%	12.73%	19.09%	6.71	7.42
SI unit equivale										i .	1

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.

						itinuea.					
Canatituant	Certified			Standard			Relative	Standard D	eviations	5% w	indow
Constituent	Value	1SD	2SD Low	2SD High	3SD Low	3SD High	1RSD	2RSD	3RSD	Low	High
Peroxide Fus	Peroxide Fusion ICP continued										
Ge, ppm	2.28	0.32	1.64	2.92	1.32	3.24	14.05%	28.09%	42.14%	2.17	2.40
Ho, ppm	1.28	0.058	1.17	1.40	1.11	1.45	4.51%	9.02%	13.53%	1.22	1.35
In, ppm	2.04	0.090	1.86	2.22	1.77	2.31	4.38%	8.77%	13.15%	1.94	2.15
K, wt.%	2.58	0.106	2.37	2.79	2.26	2.89	4.10%	8.19%	12.29%	2.45	2.71
La, ppm	73	3.8	66	81	62	84	5.18%	10.35%	15.53%	69	77
Li, ppm	54	4.4	45	62	40	67	8.27%	16.54%	24.80%	51	56
Lu, ppm	0.55	0.09	0.37	0.72	0.28	0.81	16.32%	32.63%	48.95%	0.52	0.57
Mg, wt.%	2.50	0.099	2.30	2.70	2.20	2.79	3.95%	7.91%	11.86%	2.37	2.62
Mn, wt.%	0.067	0.003	0.061	0.073	0.058	0.076	4.60%	9.19%	13.79%	0.064	0.070
Mo, ppm	15.0	1.02	13.0	17.1	11.9	18.1	6.82%	13.63%	20.45%	14.3	15.8
Nb, ppm	22.9	1.78	19.3	26.4	17.5	28.2	7.79%	15.57%	23.36%	21.7	24.0
Nd, ppm	61	4.2	53	69	48	74	6.90%	13.80%	20.70%	58	64
Ni, ppm	157	14	130	184	116	198	8.61%	17.23%	25.84%	149	165
P, wt.%	0.032	0.005	0.023	0.042	0.018	0.047	14.41%	28.82%	43.23%	0.031	0.034
Pb, ppm	21.0	3.6	13.9	28.1	10.4	31.7	16.91%	33.82%	50.73%	20.0	22.1
Pr, ppm	17.0	1.36	14.3	19.7	13.0	21.1	7.97%	15.94%	23.92%	16.2	17.9
Rb, ppm	107	6	94	120	88	126	5.88%	11.76%	17.64%	102	112
S, wt.%	3.60	0.120	3.36	3.84	3.24	3.96	3.33%	6.66%	9.98%	3.42	3.78
Sb, ppm	3.96	0.292	3.38	4.55	3.09	4.84	7.38%	14.76%	22.14%	3.76	4.16
Sc, ppm	11.4	0.92	9.5	13.2	8.6	14.1	8.09%	16.17%	24.26%	10.8	11.9
Si, wt.%	26.68	0.548	25.58	27.77	25.04	28.32	2.05%	4.11%	6.16%	25.35	28.01
Sm, ppm	8.83	0.587	7.66	10.00	7.07	10.59	6.64%	13.28%	19.93%	8.39	9.27
Sn, ppm	3.10	0.58	1.93	4.26	1.35	4.85	18.84%	37.68%	56.52%	2.94	3.25
Sr, ppm	73	7	58	88	50	95	10.24%	20.49%	30.73%	69	76
Ta, ppm	1.77	0.118	1.53	2.00	1.41	2.12	6.65%	13.31%	19.96%	1.68	1.86
Tb, ppm	1.14	0.070	1.00	1.28	0.93	1.35	6.18%	12.37%	18.55%	1.08	1.19
Th, ppm	15.6	0.38	14.8	16.3	14.4	16.7	2.47%	4.94%	7.41%	14.8	16.3
Ti, wt.%	0.436	0.013	0.411	0.461	0.398	0.474	2.89%	5.78%	8.66%	0.414	0.458
TI, ppm	1.03	0.10	0.82	1.24	0.71	1.34	10.19%	20.37%	30.56%	0.98	1.08
Tm, ppm	0.53	0.030	0.47	0.59	0.44	0.62	5.75%	11.50%	17.25%	0.50	0.55
U, ppm	8.62	0.604	7.41	9.82	6.81	10.43	7.00%	14.01%	21.01%	8.19	9.05
V, ppm	333	22	288	378	266	400	6.69%	13.39%	20.08%	316	350
W, ppm	3.10	0.92	1.27	4.93	0.35	5.85	29.56%	59.13%	88.69%	2.94	3.25
Y, ppm	34.8	1.92	30.9	38.6	29.0	40.5	5.52%	11.04%	16.57%	33.0	36.5
Yb, ppm	3.51	0.273	2.97	4.06	2.69	4.33	7.77%	15.53%	23.30%	3.34	3.69
Zn, ppm	33.5	4.7	24.2	42.9	19.5	47.5	13.94%	27.87%	41.81%	31.8	35.2
4-Acid Digest	1	0.029	0.000	0.101	0.052	0.210	20 520/	41.049/	G1 EE0/	0.120	0.142
Ag, ppm	0.135	0.028	0.080	0.191	0.052	0.219	20.52%	41.04%	61.55%	0.129	0.142 6.70
Al, wt.%	6.47 75	0.240 3.3	5.99 69	6.95 82	5.75 65	7.18 85	3.70% 4.37%	7.41% 8.74%	11.11% 13.11%	6.14 72	6.79 79
As, ppm Ba, ppm	586	5.3 58	471	702	413	760	9.84%	19.68%	29.52%	557	616
Be, ppm	4.17	0.281	3.61	4.73	3.33	5.01	6.74%	13.48%	29.32%	3.96	4.38
Bi, ppm	1.29	0.261	1.13	1.44	1.05	1.52	6.12%	12.23%	18.35%	1.22	1.35
Ca, wt.%	0.114	0.019	0.094	0.134	0.085	0.144	8.63%	17.25%	25.88%	0.108	0.120
SI unit equivale	I .			l .	l .	l .	l .		l .	0.700	0.120

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; IND = indeterminate.



rable 6 continued.											
Constituent	Certified			Standard			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	4-Acid Digestion continued										
Cd, ppm	0.38	0.035	0.31	0.45	0.28	0.49	9.16%	18.31%	27.47%	0.36	0.40
Ce, ppm	134	6	122	146	116	152	4.50%	9.00%	13.49%	127	141
Co, wt.%	0.788	0.036	0.716	0.860	0.680	0.896	4.57%	9.15%	13.72%	0.749	0.827
Cr, ppm	93	8.8	76	111	67	120	9.44%	18.89%	28.33%	89	98
Cs, ppm	2.44	0.106	2.23	2.66	2.13	2.76	4.33%	8.66%	12.99%	2.32	2.57
Cu, wt.%	3.70	0.121	3.46	3.94	3.34	4.06	3.27%	6.54%	9.81%	3.51	3.88
Dy, ppm	4.48	0.67	3.14	5.82	2.48	6.48	14.92%	29.83%	44.75%	4.26	4.70
Er, ppm	2.69	0.41	1.87	3.51	1.46	3.92	15.21%	30.43%	45.64%	2.55	2.82
Eu, ppm	0.97	0.059	0.85	1.09	0.79	1.15	6.05%	12.10%	18.15%	0.92	1.02
Fe, wt.%	3.66	0.131	3.40	3.92	3.27	4.05	3.59%	7.18%	10.77%	3.48	3.84
Ga, ppm	26.5	1.18	24.2	28.9	23.0	30.1	4.46%	8.91%	13.37%	25.2	27.9
Gd, ppm	5.08	0.80	3.47	6.69	2.66	7.49	15.86%	31.71%	47.57%	4.82	5.33
Ge, ppm	0.18	0.05	0.09	0.28	0.04	0.33	26.61%	53.22%	79.83%	0.17	0.19
Hf, ppm	4.75	0.178	4.39	5.10	4.21	5.28	3.75%	7.49%	11.24%	4.51	4.98
Ho, ppm	0.88	0.080	0.72	1.05	0.64	1.13	9.07%	18.13%	27.20%	0.84	0.93
In, ppm	2.08	0.097	1.88	2.27	1.79	2.37	4.66%	9.32%	13.98%	1.97	2.18
K, wt.%	2.57	0.092	2.38	2.75	2.29	2.84	3.57%	7.13%	10.70%	2.44	2.70
La, ppm	64	4.1	56	72	52	76	6.43%	12.85%	19.28%	61	67
Li, ppm	52	2.4	47	57	45	59	4.59%	9.18%	13.76%	49	55
Lu, ppm	0.41	0.015	0.38	0.44	0.36	0.45	3.68%	7.36%	11.03%	0.39	0.43
Mg, wt.%	2.46	0.100	2.26	2.66	2.16	2.76	4.04%	8.09%	12.13%	2.34	2.59
Mn, wt.%	0.065	0.003	0.059	0.070	0.057	0.072	4.05%	8.11%	12.16%	0.061	0.068
Mo, ppm	14.7	0.93	12.9	16.6	11.9	17.5	6.32%	12.64%	18.95%	14.0	15.4
Na, wt.%	0.063	0.011	0.042	0.084	0.031	0.095	16.92%	33.84%	50.76%	0.060	0.066
Nb, ppm	15.0	2.5	9.9	20.0	7.4	22.5	16.75%	33.50%	50.25%	14.2	15.7
Nd, ppm	58	4.4	49	67	45	71	7.53%	15.07%	22.60%	55	61
Ni, ppm	151	8	135	167	128	175	5.21%	10.43%	15.64%	144	159
P, wt.%	0.033	0.002	0.028	0.038	0.026	0.040	7.09%	14.17%	21.26%	0.031	0.035
Pb, ppm	15.9	0.75	14.4	17.4	13.6	18.1	4.75%	9.50%	14.26%	15.1	16.7
Pr, ppm	15.5	1.6	12.2	18.7	10.6	20.3	10.46%	20.91%	31.37%	14.7	16.2
Rb, ppm	106	8	90	123	82	131	7.62%	15.24%	22.86%	101	112
Re, ppm	0.024	0.003	0.018	0.030	0.016	0.032	11.62%	23.25%	34.87%	0.023	0.025
S, wt.%	3.57	0.134	3.30	3.83	3.17	3.97	3.75%	7.51%	11.26%	3.39	3.75
Sb, ppm	3.79	0.309	3.17	4.41	2.86	4.71	8.16%	16.33%	24.49%	3.60	3.98
Sc, ppm	12.4	0.68	11.0	13.7	10.3	14.4	5.49%	10.97%	16.46%	11.7	13.0
Se, ppm	5.99	0.527	4.94	7.05	4.41	7.57	8.79%	17.57%	26.36%	5.69	6.29
Sm, ppm	8.29	0.484	7.32	9.26	6.83	9.74	5.84%	11.68%	17.52%	7.87	8.70
Sn, ppm	3.21	0.214	2.79	3.64	2.57	3.85	6.65%	13.30%	19.95%	3.05	3.37
Sr, ppm	66	3.6	59	73	55	77	5.46%	10.92%	16.38%	63	69
Ta, ppm	1.05	0.21	0.62	1.47	0.40	1.69	20.45%	40.90%	61.36%	0.99	1.10
Tb, ppm	0.77	0.12	0.54	1.01	0.42	1.13	15.28%	30.55%	45.83%	0.73	0.81
Te, ppm	0.078	0.021	0.036	0.120	0.015	0.141	27.03%	54.05%	81.08%	0.074	0.082
Th, ppm	15.4	0.95	13.5	17.3	12.6	18.3	6.16%	12.33%	18.49%	14.6	16.2
Ti, wt.%	0.275	0.073	0.130	0.421	0.057	0.493	26.38%	52.75%	79.13%	0.262	0.289

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.



Table 6 Continued.											
0 " 1	Certified		Absolute Standard Deviations Relative Standard Devia				eviations	5% w	indow		
Constituent	Value	1SD	2SD Low	2SD	3SD	3SD High	1RSD	2RSD	3RSD	Low	High
4-Acid Digest	ion continue	ed	LOW	High	Low	High					
Tm, ppm	0.39	0.06	0.27	0.50	0.21	0.56	14.87%	29.74%	44.61%	0.37	0.41
U, ppm	8.26	0.581	7.10	9.42	6.52	10.00	7.03%	14.07%	21.10%	7.85	8.67
V, ppm	316	15	286	345	272	359	4.62%	9.25%	13.87%	300	331
W, ppm	2.63	0.187	2.26	3.01	2.07	3.19	7.11%	14.21%	21.32%	2.50	2.76
Y, ppm	22.8	1.14	20.5	25.1	19.4	26.2	5.00%	10.00%	15.00%	21.7	23.9
Yb, ppm	2.70	0.35	2.00	3.40	1.65	3.75	12.99%	25.98%	38.97%	2.56	2.83
Zn, ppm	31.6	1.77	28.1	35.2	26.3	36.9	5.60%	11.20%	16.80%	30.0	33.2
Zr, ppm	165	5	155	176	149	182	3.25%	6.50%	9.76%	157	174
Aqua Regia D	igestion										
Ag, ppm	0.088	0.011	0.067	0.110	0.057	0.120	11.93%	23.85%	35.78%	0.084	0.093
Al, wt.%	0.868	0.185	0.498	1.238	0.313	1.423	21.31%	42.62%	63.93%	0.824	0.911
As, ppm	72	3.8	65	80	61	84	5.24%	10.49%	15.73%	69	76
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	81	14	53	109	38	123	17.50%	34.99%	52.49%	77	85
Be, ppm	1.84	0.22	1.39	2.28	1.17	2.50	12.07%	24.13%	36.20%	1.75	1.93
Bi, ppm	1.19	0.065	1.06	1.32	0.99	1.38	5.50%	11.00%	16.50%	1.13	1.25
Ca, wt.%	0.108	0.004	0.099	0.117	0.094	0.121	4.15%	8.29%	12.44%	0.102	0.113
Cd, ppm	0.38	0.028	0.33	0.44	0.30	0.46	7.24%	14.47%	21.71%	0.36	0.40
Ce, ppm	46.3	7.6	31.0	61.5	23.3	69.2	16.51%	33.03%	49.54%	43.9	48.6
Co, wt.%	0.759	0.054	0.650	0.868	0.596	0.922	7.17%	14.34%	21.51%	0.721	0.797
Cr, ppm	33.3	4.2	24.9	41.6	20.8	45.7	12.48%	24.97%	37.45%	31.6	34.9
Cs, ppm	0.43	0.12	0.18	0.68	0.06	0.81	28.73%	57.45%	86.18%	0.41	0.46
Cu, wt.%	3.74	0.087	3.56	3.91	3.47	4.00	2.33%	4.66%	7.00%	3.55	3.92
Dy, ppm	1.10	0.19	0.71	1.48	0.52	1.67	17.55%	35.09%	52.64%	1.04	1.15
Er, ppm	0.46	0.09	0.28	0.63	0.20	0.72	19.05%	38.11%	57.16%	0.44	0.48
Eu, ppm	0.43	0.10	0.23	0.62	0.13	0.72	22.91%	45.81%	68.72%	0.41	0.45
Fe, wt.%	3.51	0.177	3.16	3.87	2.98	4.05	5.03%	10.06%	15.10%	3.34	3.69
Ga, ppm	7.13	1.22	4.70	9.57	3.48	10.79	17.08%	34.15%	51.23%	6.78	7.49
Gd, ppm	2.31	0.159	1.99	2.63	1.83	2.79	6.90%	13.80%	20.70%	2.19	2.42
Ge, ppm	0.093	0.027	0.040	0.147	0.013	0.174	28.84%	57.67%	86.51%	0.089	0.098
Hf, ppm	0.33	0.05	0.24	0.42	0.19	0.47	13.84%	27.68%	41.52%	0.31	0.35
Hg, ppm	0.054	0.010	0.035	0.073	0.025	0.083	17.64%	35.29%	52.93%	0.051	0.057
Ho, ppm	0.18	0.03	0.12	0.24	0.09	0.27	16.53%	33.06%	49.59%	0.17	0.19
In, ppm	1.47	0.062	1.35	1.60	1.29	1.66	4.21%	8.42%	12.62%	1.40	1.55
K, wt.%	0.203	0.030	0.144	0.262	0.115	0.292	14.54%	29.09%	43.63%	0.193	0.213
La, ppm	20.9	3.2	14.6	27.3	11.4	30.4	15.11%	30.23%	45.34%	19.9	22.0
Li, ppm	27.7	6.1	15.5	39.9	9.3	46.1	22.10%	44.20%	66.31%	26.3	29.1
Mg, wt.%	1.88	0.20	1.48	2.27	1.29	2.47	10.50%	21.01%	31.51%	1.78	1.97
Mn, wt.%	0.064	0.003	0.057	0.070	0.054	0.073	5.03%	10.06%	15.09%	0.061	0.067
Mo, ppm	14.0	0.69	12.6	15.3	11.9	16.0	4.97%	9.94%	14.92%	13.3	14.7
Na, wt.%	0.010	0.002	0.006	0.014	0.005	0.016	18.99%	37.98%	56.98%	0.010	0.011
Nb, ppm	0.083	0.016	0.051	0.115	0.035	0.131	19.11%	38.21%	57.32%	0.079	0.087
Nd, ppm	21.1	5.4	10.3	32.0	4.8	37.5	25.72%	51.45%	77.17%	20.1	22.2

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; IND = indeterminate.



	Certified		Absolute	Standard	Deviations	3	Relative	Standard D	eviations	5% w	5% window	
Constituent	Value	1SD	2SD Low	2SD High	3SD Low	3SD High	1RSD	2RSD	3RSD	Low	High	
Aqua Regia D	Aqua Regia Digestion continued											
Ni, ppm	147	7	133	161	126	168	4.76%	9.53%	14.29%	140	154	
P, wt.%	0.020	0.001	0.018	0.023	0.017	0.024	5.16%	10.31%	15.47%	0.019	0.021	
Pb, ppm	11.8	1.7	8.5	15.1	6.9	16.8	14.01%	28.01%	42.02%	11.2	12.4	
Pr, ppm	5.46	1.42	2.62	8.30	1.20	9.72	26.01%	52.03%	78.04%	5.18	5.73	
Rb, ppm	8.01	1.29	5.44	10.58	4.15	11.87	16.06%	32.12%	48.19%	7.61	8.41	
Re, ppm	0.021	0.002	0.017	0.026	0.015	0.028	10.32%	20.65%	30.97%	0.020	0.023	
S, wt.%	3.48	0.144	3.19	3.77	3.05	3.91	4.14%	8.28%	12.43%	3.31	3.66	
Sb, ppm	2.22	0.30	1.61	2.82	1.31	3.12	13.57%	27.13%	40.70%	2.10	2.33	
Sc, ppm	2.91	0.32	2.27	3.54	1.96	3.86	10.88%	21.76%	32.65%	2.76	3.05	
Se, ppm	5.95	0.550	4.85	7.05	4.30	7.60	9.25%	18.49%	27.74%	5.65	6.25	
Sm, ppm	3.72	0.59	2.54	4.90	1.94	5.50	15.91%	31.83%	47.74%	3.53	3.91	
Sn, ppm	1.31	0.126	1.06	1.56	0.93	1.69	9.63%	19.27%	28.90%	1.24	1.38	
Sr, ppm	15.8	3.7	8.4	23.2	4.7	27.0	23.42%	46.85%	70.27%	15.0	16.6	
Ta, ppm	< 0.01	IND	IND	IND	IND	IND	IND	IND	IND	IND	IND	
Tb, ppm	0.26	0.03	0.19	0.32	0.16	0.35	12.21%	24.41%	36.62%	0.24	0.27	
Te, ppm	0.057	0.012	0.034	0.081	0.022	0.093	20.48%	40.97%	61.45%	0.054	0.060	
Th, ppm	6.93	1.06	4.82	9.05	3.77	10.10	15.22%	30.44%	45.66%	6.59	7.28	
U, ppm	3.22	0.259	2.70	3.73	2.44	3.99	8.04%	16.09%	24.13%	3.05	3.38	
V, ppm	52	7	38	66	32	73	13.09%	26.19%	39.28%	49	55	
W, ppm	0.91	0.11	0.70	1.13	0.59	1.23	11.73%	23.45%	35.18%	0.87	0.96	
Y, ppm	4.37	0.56	3.25	5.49	2.69	6.04	12.78%	25.56%	38.34%	4.15	4.59	
Yb, ppm	0.37	0.08	0.20	0.54	0.12	0.62	22.73%	45.46%	68.19%	0.35	0.39	
Zn, ppm	28.2	1.61	25.0	31.4	23.4	33.0	5.71%	11.42%	17.13%	26.8	29.6	
Zr, ppm	10.4	1.5	7.4	13.5	5.9	15.0	14.57%	29.13%	43.70%	9.9	10.9	

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

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



PARTICIPATING LABORATORIES

- 1. Actlabs, Ancaster, Ontario, Canada
- 2. African Natural Resources & Mines Ltd, Suleja, Niger State, Nigeria
- 3. AGAT Laboratories, Calgary, Alberta, Canada
- 4. ALS, Brisbane, QLD, Australia
- 5. ALS, Lima, Peru
- 6. ALS, Loughrea, Galway, Ireland
- ALS, Malaga, WA, Australia
- 8. ALS, Vancouver, BC, Canada
- 9. American Assay Laboratories, Sparks, Nevada, USA
- 10. ARGETEST Mineral Processing, Ankara, Central Anatolia, Turkey
- 11. Bureau Veritas Commodities Canada Ltd, Vancouver, BC, Canada
- 12. CERTIMIN, Lima, Peru
- 13. ESAN Istanbul, Istanbul, Turkey
- 14. Inspectorate (BV), Lima, Peru
- 15. Intertek Genalysis, Perth, WA, Australia
- 16. Intertek Testing Services, Townsville, QLD, Australia
- 17. Intertek Testing Services Philippines, Cupang, Muntinlupa, Philippines
- 18. Labwest Minerals Analysis, Perth, WA, Australia
- 19. MSALABS, Vancouver, BC, Canada
- 20. Ontario Geological Survey, Sudbury, Ontario, Canada
- 21. PT Geoservices Ltd, Cikarang, Jakarta Raya, Indonesia
- 22. PT Intertek Utama Services, Jakarta Timur, DKI Jakarta, Indonesia
- 23. Saskatchewan Research Council, Saskatoon, Saskatchewan, Canada
- 24. SGS, Randfontein, Gauteng, South Africa
- 25. SGS del Peru, Lima, Peru
- 26. Shiva Analyticals Ltd, Bangalore North, Karnataka, India
- 27. Skyline Assayers & Laboratories, Tucson, Arizona, USA
- 28. 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

Certified reference material OREAS 554b is prepared, certified and supplied by:



ORE Research & Exploration Pty Ltd 37A Hosie Street

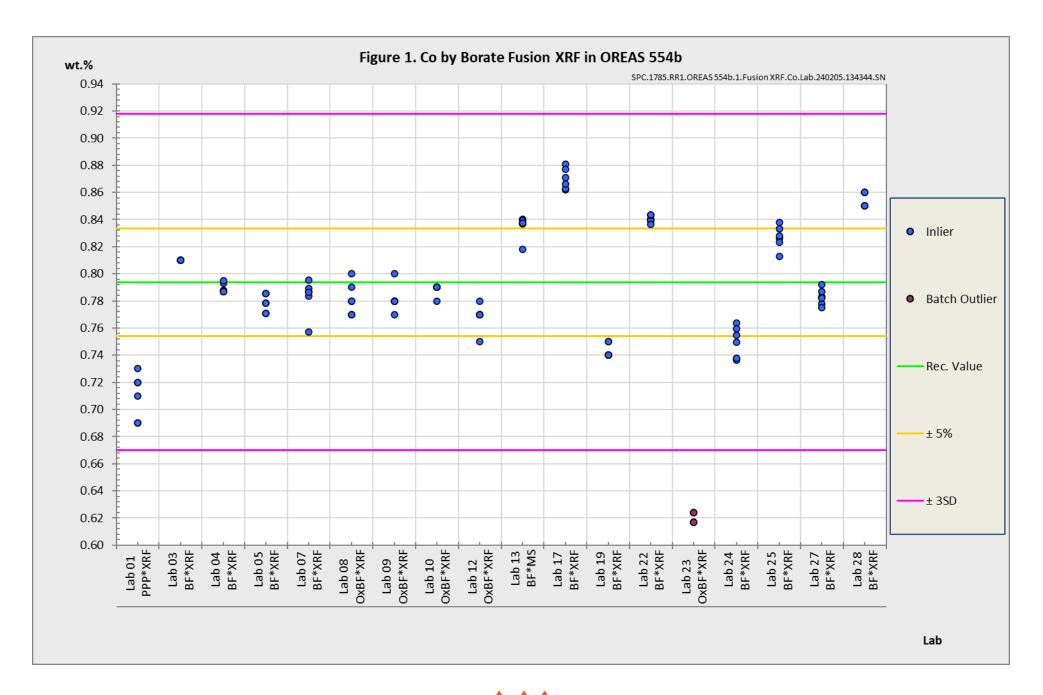
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AUSTRALIA

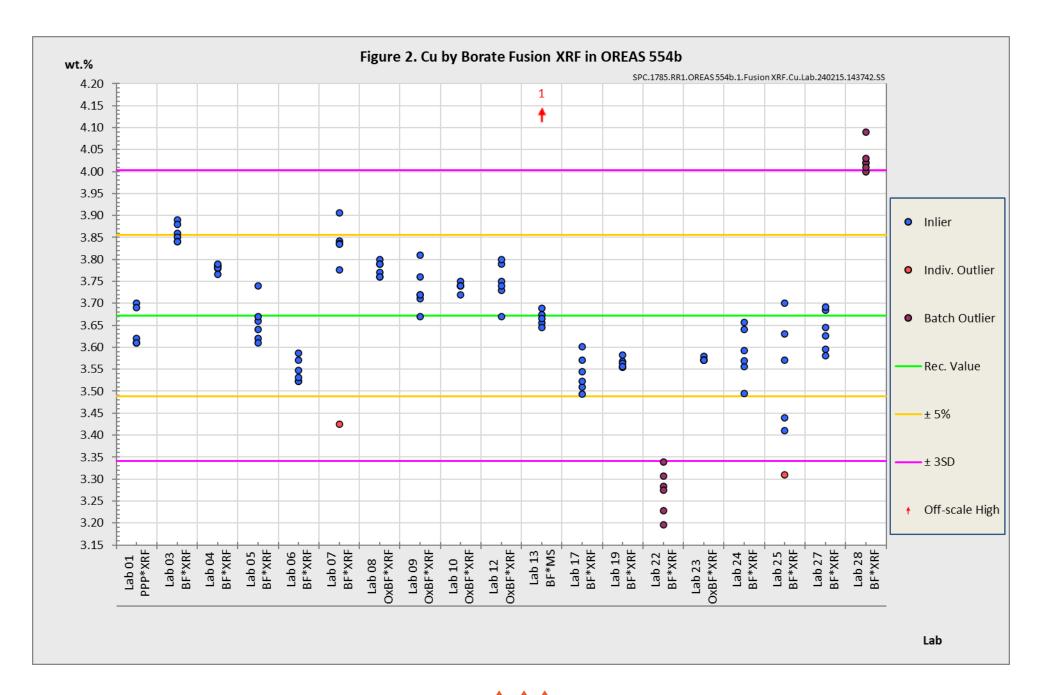
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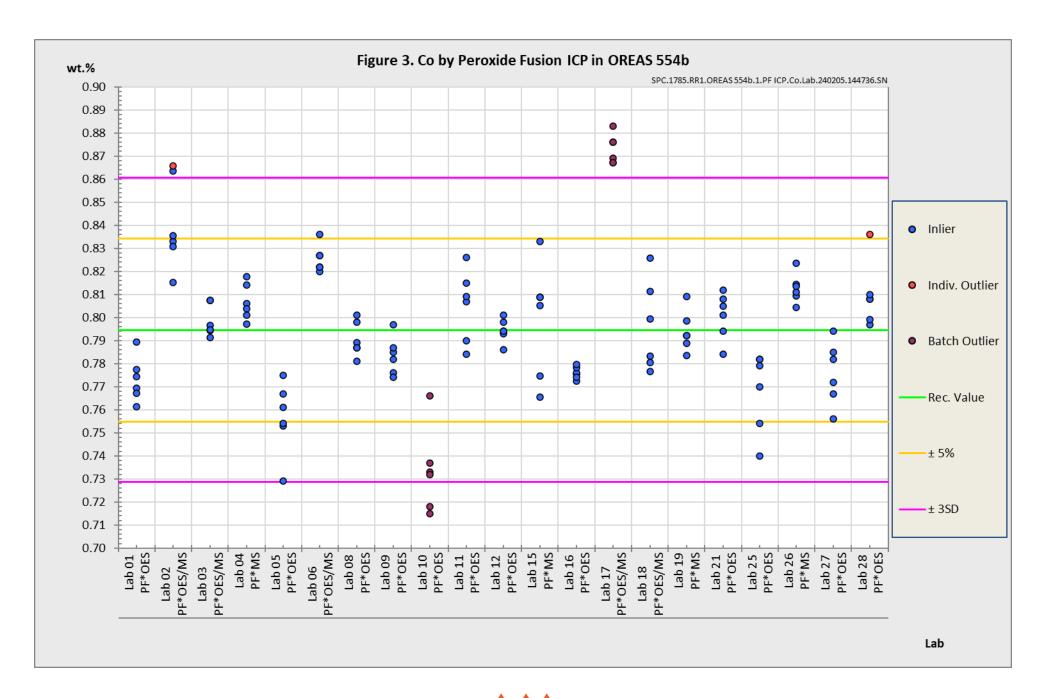
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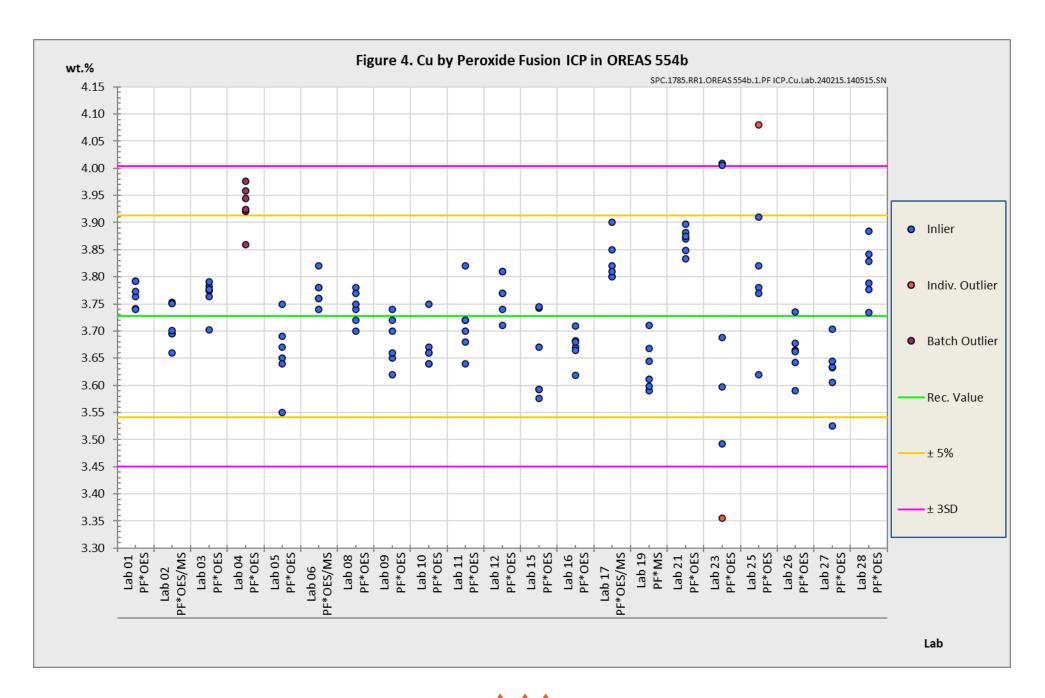
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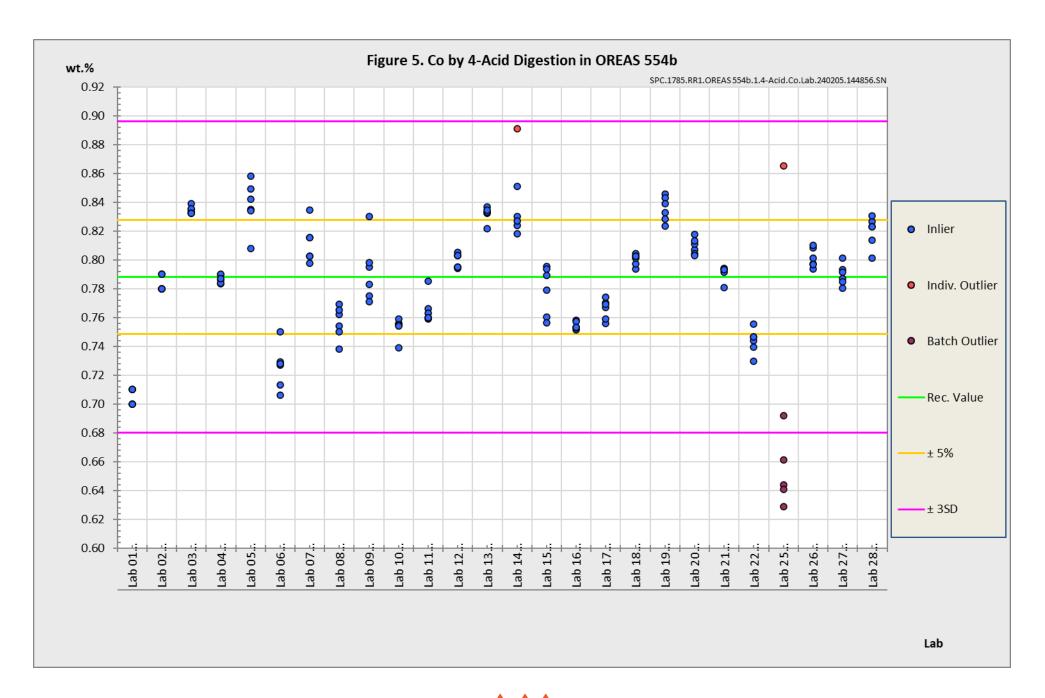


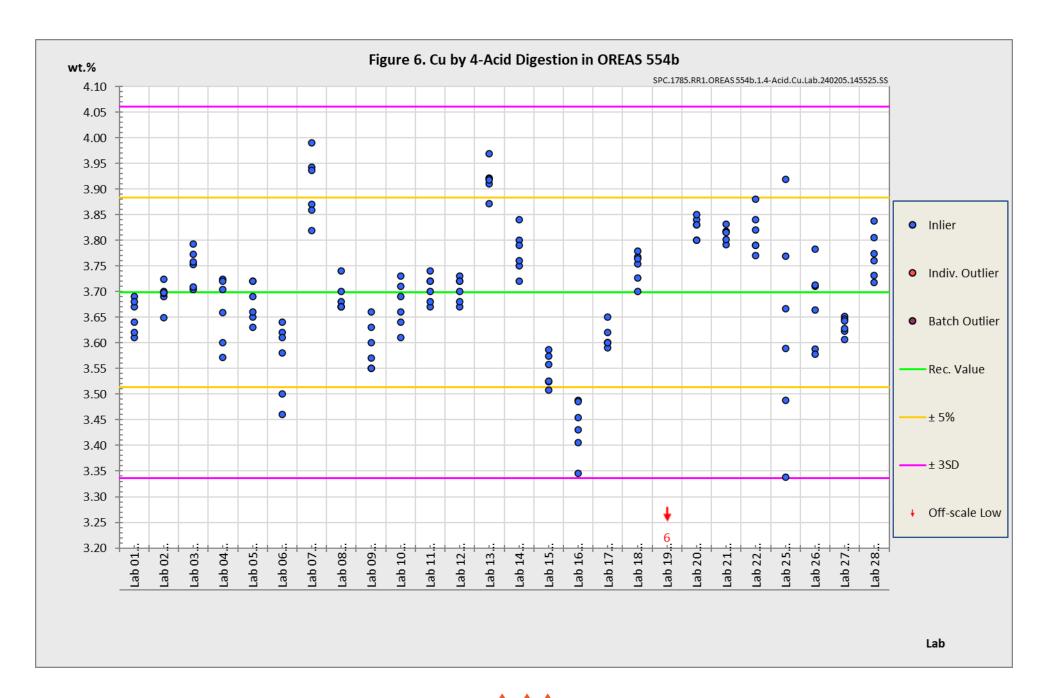


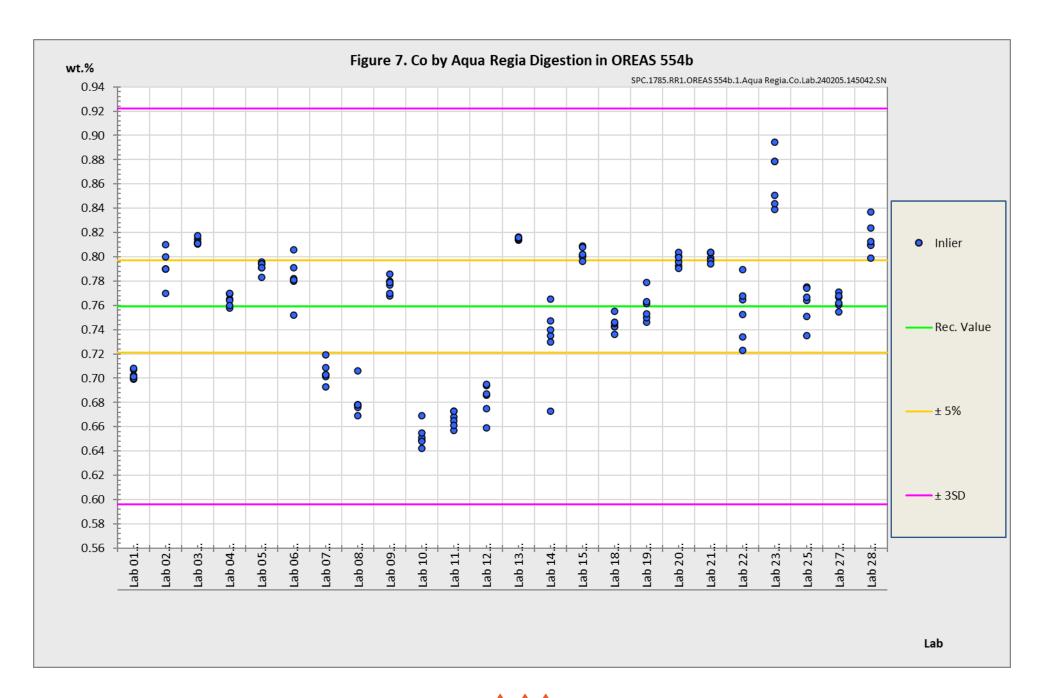


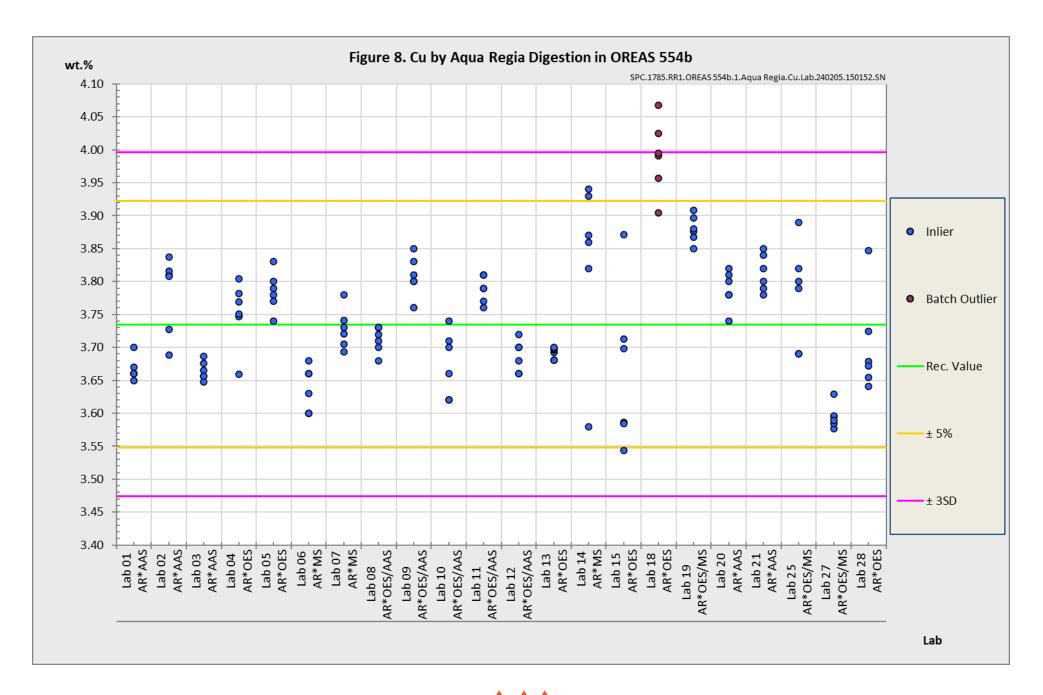












METROLOGICAL TRACEABILITY

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)). In line with popular use, all data within tables in this certificate are expressed as the mass fraction in either weight percent (wt.%) or parts per million (ppm).

The analytical samples sent to participating laboratories were selected in a manner to be representative of the entire prepared batch of CRM. This 'representivity' was maintained in each submitted laboratory sample batch and ensures the user that the data is traceable from sample selection through to the analytical results. The systematic sampling method was chosen due to the low risk of overlooking repetitive effects or trends in the batch due to the way the CRM was processed. In line with ISO 17025 [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 and aqua regia digestion methods (Table 1). The other operationally defined measurands characterised in this certificate (Table 2) are derived from data procured mostly from ISO 17025 accredited laboratories. The certified values presented in this report are calculated from the means of accepted data following robust technical and statistical analysis as detailed in this report.

Guide ISO/TR 16476:2016, section 5.3.1 describes metrological traceability in reference materials as it pertains to the transformation of the measurand. In this section it states, "Although the determination of the property value itself can be made traceable to appropriate units through, for example, calibration of the measurement equipment used, steps like the transformation of the sample from one physical (chemical) state to another cannot. Such transformations may only be compared with a reference (when available), or among themselves. For some transformations, reference methods have been defined and may be used in certification projects to evaluate the uncertainty associated with such a transformation. In other cases, only a comparison among different laboratories using the same procedure is possible. In this case, it is impossible to demonstrate absence of method bias; therefore, the result is an operationally defined measurand (ISO Guide 35:2017, 9.2.4c)." Certification takes place on the basis of agreement among operationally defined, independent measurement results.

COMMUTABILITY

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

INTENDED USE

OREAS 554b 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 554b may be used to calibrate the entire procedure by producing a pure substance CRM transformed into a calibration solution.

OREAS 554b 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:

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

PERIOD OF VALIDITY & STORAGE INSTRUCTIONS

The certification of OREAS 554b remains valid, within the specified measurement uncertainties, until September 2033, provided the CRM is handled and stored in accordance with the instructions given below. This certification is nullified if the CRM is any way changed or contaminated.

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

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



Single-use sachets (e.g., 10g units)

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, well ventilated area at temperatures between -5° 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 6 are intended only to be used as a preliminary guide as to what a laboratory may be able to achieve. Over a period of time monitoring your own laboratory's data for this CRM, SD's should be calculated directly from your own laboratory's process. This will enable you to establish more specific performance gates that are fit for purpose for your application as well as the ability to monitor bias. If your long-term trend analysis shows an average value that is within the 95% expanded uncertainty then generally there is no cause for concern in regard to bias.

For use with the aqua regia digestion method

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

The approach applied here is to report certified values in those instances where reasonable agreement exists amongst a majority of participating laboratories. The results of specific laboratories may differ significantly from the certified values, but will, nonetheless, be valid

and reproducible in the context of the specifics of the aqua regia method in use. Users of this reference material should, therefore, be mindful of this limitation when applying the certified values in a quality control program.

LEGAL NOTICE

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

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DOCUMENT HISTORY

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0	6 th March, 2024	First publication.

CERTIFYING OFFICER

Sign

6th March, 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 the two sulphuric acid leaches, specific methodologies were detailed for the participating laboratories to follow:

5% sulphuric acid leach

- 1. Weigh $0.500 \pm 0.002g$ of sample pulp into a clean 250 ml flask.
- 2. Add to the flask 0.5g 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.
- 7. The solution is now ready to be analysed by ICP or AAS.