

CERTIFICATE OF ANALYSIS FOR

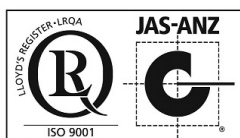
CERTIFIED REFERENCE MATERIAL

OREAS 553b

Copper-Cobalt Ore (Democratic Republic of the Congo)



Accredited for compliance with ISO 17034



COA-1904-OREA553b-R0
BUP-70-10-01 Ver:2.0

12-Nov-2024

Table 1. Certified Values, Uncertainty & Tolerance Intervals for multi-elements by 4-acid digestion in OREAS 553b.

Constituent	Certified Value [†]	95% Expanded Uncertainty		95 % Tolerance Limits	
		Low	High	Low	High
4-Acid Digestion					
Ag, Silver (ppm)	0.103	0.085	0.121	IND	IND
Al, Aluminium (wt.%)	6.41	6.25	6.56	6.29	6.53
As, Arsenic (ppm)	54	51	57	52	55
Ba, Barium (ppm)	262	251	273	255	268
Be, Beryllium (ppm)	4.71	4.56	4.86	4.58	4.83
Bi, Bismuth (ppm)	1.70	1.62	1.78	1.65	1.75
Ca, Calcium (wt.%)	0.313	0.304	0.322	0.305	0.320
Cd, Cadmium (ppm)	0.30	0.27	0.32	0.27	0.32
Ce, Cerium (ppm)	144	135	154	140	149
Co, Cobalt (wt.%)	0.587	0.577	0.597	0.581	0.593
Cr, Chromium (ppm)	91	85	96	88	94
Cs, Caesium (ppm)	2.08	1.98	2.18	2.00	2.16
Cu, Copper (wt.%)	3.60	3.54	3.66	3.57	3.64
Dy, Dysprosium (ppm)	4.44	3.98	4.91	4.24	4.64
Er, Erbium (ppm)	2.85	2.54	3.16	2.73	2.96
Eu, Europium (ppm)	0.81	0.68	0.94	0.77	0.85
Fe, Iron (wt.%)	3.18	3.12	3.25	3.13	3.23
Ga, Gallium (ppm)	27.9	26.7	29.1	27.2	28.6
Gd, Gadolinium (ppm)	5.01	4.31	5.70	4.86	5.15
Hf, Hafnium (ppm)	5.46	5.19	5.73	5.30	5.62
Ho, Holmium (ppm)	0.96	0.82	1.11	0.93	1.00
In, Indium (ppm)	2.14	2.06	2.21	2.08	2.20
K, Potassium (wt.%)	2.55	2.49	2.61	2.50	2.61
La, Lanthanum (ppm)	68	63	74	66	70
Li, Lithium (ppm)	63	61	64	61	64
Lu, Lutetium (ppm)	0.42	0.36	0.48	0.40	0.44
Mg, Magnesium (wt.%)	3.58	3.49	3.67	3.50	3.65
Mn, Manganese (wt.%)	0.013	0.013	0.014	0.013	0.014
Mo, Molybdenum (ppm)	15.1	14.5	15.7	14.7	15.4
Na, Sodium (wt.%)	0.045	0.042	0.048	0.043	0.047
Nb, Niobium (ppm)	17.4	15.4	19.4	16.6	18.2
Nd, Neodymium (ppm)	63	56	70	61	64
Ni, Nickel (ppm)	124	122	127	123	126
P, Phosphorus (wt.%)	0.035	0.034	0.036	0.034	0.036
Pb, Lead (ppm)	11.2	10.5	12.0	10.8	11.7
Pr, Praseodymium (ppm)	16.9	15.2	18.5	16.4	17.3

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

Note: intervals may appear asymmetric due to rounding.

Table 1 continued.

Constituent	Certified Value [†]	95 % Expanded Uncertainty		95 % Tolerance Limits	
		Low	High	Low	High
4-Acid Digestion continued					
Rb, Rubidium (ppm)	96	90	101	92	99
Re, Rhenium (ppm)	0.027	0.024	0.031	0.024	0.031
S, Sulphur (wt.%)	3.34	3.27	3.40	3.29	3.39
Sb, Antimony (ppm)	3.01	2.79	3.24	2.84	3.18
Sc, Scandium (ppm)	12.5	11.9	13.2	12.2	12.8
Se, Selenium (ppm)	5.73	4.96	6.50	5.16	6.30
Sm, Samarium (ppm)	8.69	7.77	9.61	8.42	8.96
Sn, Tin (ppm)	2.81	2.66	2.96	2.65	2.97
Sr, Strontium (ppm)	70	67	73	68	72
Ta, Tantalum (ppm)	1.26	1.09	1.43	1.20	1.33
Tb, Terbium (ppm)	0.74	0.66	0.83	0.69	0.79
Te, Tellurium (ppm)	0.063	0.049	0.078	IND	IND
Th, Thorium (ppm)	15.5	14.8	16.1	14.8	16.1
Ti, Titanium (wt.%)	0.258	0.223	0.293	0.250	0.266
Tl, Thallium (ppm)	0.67	0.63	0.70	0.64	0.70
Tm, Thulium (ppm)	0.42	0.33	0.51	0.39	0.46
U, Uranium (ppm)	8.60	8.26	8.95	8.30	8.90
V, Vanadium (ppm)	348	339	358	341	356
W, Tungsten (ppm)	2.52	2.32	2.72	2.37	2.67
Y, Yttrium (ppm)	23.8	22.5	25.0	23.1	24.5
Yb, Ytterbium (ppm)	2.84	2.53	3.15	2.73	2.95
Zn, Zinc (ppm)	16.1	15.1	17.2	15.5	16.8
Zr, Zirconium (ppm)	188	181	195	183	192

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).

Table 2. Certified Values, Uncertainty & Tolerance Intervals for other measurands in OREAS 553b.

Constituent	Certified Value	95 % Expanded Uncertainty		95 % Tolerance Limits	
		Low	High	Low	High
Sulphuric Acid 5% Leach					
Co, Cobalt (wt.%)	0.127	0.119	0.135	0.124	0.130
Cu, Copper (wt.%)	1.07	1.02	1.11	1.04	1.09
Infrared Combustion					
C, Carbon (wt.%)	4.51	4.43	4.59	4.47	4.55
S, Sulphur (wt.%)	3.40	3.35	3.45	3.37	3.44
Borate Fusion XRF					
Al ₂ O ₃ , Aluminium(III) oxide (wt.%)	12.51	12.32	12.69	12.37	12.64
BaO, Barium oxide (ppm)	303	241	365	IND	IND
CaO, Calcium oxide (wt.%)	0.435	0.423	0.447	0.421	0.449
Co, Cobalt (wt.%)	0.587	0.572	0.601	0.577	0.597
Cu, Copper (wt.%)	3.62	3.56	3.69	3.57	3.67
Fe ₂ O ₃ , Iron(III) oxide (wt.%)	4.60	4.53	4.67	4.55	4.65
K ₂ O, Potassium oxide (wt.%)	3.04	2.99	3.10	3.01	3.08
MgO, Magnesium oxide (wt.%)	5.94	5.84	6.04	5.86	6.02
MnO, Manganese oxide (wt.%)	0.017	0.014	0.019	IND	IND
Ni, Nickel (ppm)	125	87	164	IND	IND
P ₂ O ₅ , Phosphorus(V) oxide (wt.%)	0.083	0.074	0.092	0.079	0.087
SiO ₂ , Silicon dioxide (wt.%)	53.72	53.17	54.28	53.28	54.16
SO ₃ , Sulphur trioxide (wt.%)	8.59	8.37	8.82	8.44	8.75
SrO, Strontium oxide (ppm)	96	75	116	IND	IND
TiO ₂ , Titanium dioxide (wt.%)	0.791	0.775	0.808	0.774	0.809
V ₂ O ₅ , Vanadium(V) oxide (ppm)	614	528	700	IND	IND
Zr, Zirconium (ppm)	240	187	294	IND	IND
Thermogravimetry					
LOI ¹⁰⁰⁰ , Loss On Ignition @1000°C (wt.%)	12.72	12.53	12.92	12.64	12.81
Peroxide Fusion ICP					
Al, Aluminium (wt.%)	6.47	6.31	6.63	6.36	6.57
As, Arsenic (ppm)	53	47	59	50	56
B, Boron (ppm)	217	196	238	210	224
Ba, Barium (ppm)	264	254	274	257	271
Be, Beryllium (ppm)	5.42	4.62	6.22	IND	IND
Bi, Bismuth (ppm)	1.77	1.60	1.94	IND	IND
Ca, Calcium (wt.%)	0.307	0.280	0.334	0.292	0.321
Ce, Cerium (ppm)	152	144	159	147	156
Co, Cobalt (wt.%)	0.581	0.565	0.597	0.574	0.589
Cr, Chromium (ppm)	110	97	124	102	119
Cs, Caesium (ppm)	2.13	2.01	2.25	2.02	2.24

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).

Table 2 continued.

Constituent	Certified Value	95 % Expanded Uncertainty		95 % Tolerance Limits	
		Low	High	Low	High
Peroxide Fusion ICP continued					
Cu, Copper (wt.%)	3.61	3.50	3.71	3.55	3.66
Dy, Dysprosium (ppm)	7.03	6.61	7.45	6.66	7.40
Er, Erbium (ppm)	3.73	3.39	4.08	3.50	3.97
Eu, Europium (ppm)	0.91	0.82	1.00	0.82	1.00
Fe, Iron (wt.%)	3.17	3.08	3.25	3.12	3.22
Ga, Gallium (ppm)	27.7	26.3	29.0	26.4	28.9
Gd, Gadolinium (ppm)	6.87	6.45	7.29	6.44	7.31
Ge, Germanium (ppm)	2.24	1.35	3.12	IND	IND
Ho, Holmium (ppm)	1.35	1.21	1.49	1.18	1.53
In, Indium (ppm)	2.22	2.03	2.40	2.07	2.37
K, Potassium (wt.%)	2.58	2.51	2.65	2.52	2.65
La, Lanthanum (ppm)	77	73	81	74	80
Li, Lithium (ppm)	65	60	69	63	67
Lu, Lutetium (ppm)	0.56	0.47	0.65	0.50	0.61
Mg, Magnesium (wt.%)	3.57	3.48	3.65	3.48	3.66
Mn, Manganese (wt.%)	0.014	0.013	0.015	0.014	0.014
Mo, Molybdenum (ppm)	16.0	14.5	17.4	IND	IND
Nb, Niobium (ppm)	25.6	23.8	27.3	24.0	27.2
Nd, Neodymium (ppm)	65	62	67	62	67
Ni, Nickel (ppm)	126	116	136	119	132
P, Phosphorus (wt.%)	0.037	0.027	0.048	IND	IND
Pb, Lead (ppm)	32.8	21.9	43.8	IND	IND
Pr, Praseodymium (ppm)	18.1	17.3	19.0	17.4	18.8
Rb, Rubidium (ppm)	96	93	100	93	99
Re, Rhenium (ppm)	< 0.1	IND	IND	IND	IND
S, Sulphur (wt.%)	3.29	3.18	3.41	3.22	3.36
Sb, Antimony (ppm)	3.35	2.93	3.77	2.99	3.71
Sc, Scandium (ppm)	12.2	11.2	13.3	IND	IND
Si, Silicon (wt.%)	25.17	24.57	25.78	24.74	25.60
Sm, Samarium (ppm)	8.90	8.21	9.59	8.52	9.28
Sr, Strontium (ppm)	70	66	74	67	72
Ta, Tantalum (ppm)	2.01	1.66	2.36	1.77	2.25
Tb, Terbium (ppm)	1.18	1.10	1.27	1.10	1.27
Th, Thorium (ppm)	16.0	15.3	16.6	15.4	16.5
Ti, Titanium (wt.%)	0.461	0.447	0.475	0.448	0.474
Tl, Thallium (ppm)	0.73	0.61	0.84	IND	IND
Tm, Thulium (ppm)	0.57	0.51	0.63	0.52	0.62

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 2 continued.

Constituent	Certified Value	95 % Expanded Uncertainty		95 % Tolerance Limits	
		Low	High	Low	High
Peroxide Fusion ICP continued					
U, Uranium (ppm)	8.70	8.28	9.12	8.42	8.98
V, Vanadium (ppm)	365	352	378	358	371
W, Tungsten (ppm)	3.18	2.50	3.86	IND	IND
Y, Yttrium (ppm)	35.6	34.1	37.1	33.8	37.4
Yb, Ytterbium (ppm)	3.63	3.26	4.00	3.43	3.84
Aqua Regia Digestion					
Ag, Silver (ppm)	0.077	0.064	0.089	IND	IND
Al, Aluminium (wt.%)	0.940	0.869	1.010	0.903	0.976
As, Arsenic (ppm)	51	49	53	50	53
Au, Gold (ppm)	< 0.02	IND	IND	IND	IND
B, Boron (ppm)	< 10	IND	IND	IND	IND
Ba, Barium (ppm)	49.9	46.5	53.3	47.1	52.7
Be, Beryllium (ppm)	2.16	2.01	2.30	2.09	2.22
Bi, Bismuth (ppm)	1.58	1.51	1.65	1.54	1.62
Ca, Calcium (wt.%)	0.300	0.289	0.311	0.292	0.308
Cd, Cadmium (ppm)	0.30	0.27	0.32	0.28	0.32
Ce, Cerium (ppm)	42.9	39.4	46.3	41.2	44.5
Co, Cobalt (wt.%)	0.548	0.521	0.574	0.535	0.560
Cr, Chromium (ppm)	33.9	32.1	35.8	32.7	35.1
Cs, Caesium (ppm)	0.29	0.25	0.34	0.28	0.31
Cu, Copper (wt.%)	3.60	3.56	3.65	3.57	3.64
Dy, Dysprosium (ppm)	0.80	0.68	0.92	0.78	0.82
Er, Erbium (ppm)	0.36	0.33	0.40	0.34	0.38
Eu, Europium (ppm)	0.31	0.27	0.36	0.30	0.33
Fe, Iron (wt.%)	2.97	2.89	3.05	2.91	3.02
Ga, Gallium (ppm)	7.67	7.01	8.34	7.35	8.00
Gd, Gadolinium (ppm)	1.74	1.48	1.99	1.68	1.79
Hf, Hafnium (ppm)	0.30	0.28	0.32	0.28	0.32
Hg, Mercury (ppm)	0.050	0.037	0.064	IND	IND
Ho, Holmium (ppm)	0.14	0.12	0.16	0.13	0.14
In, Indium (ppm)	1.54	1.46	1.63	1.46	1.62
K, Potassium (wt.%)	0.190	0.180	0.200	0.180	0.200
La, Lanthanum (ppm)	19.0	17.5	20.4	18.3	19.7
Li, Lithium (ppm)	35.5	32.2	38.8	34.2	36.8
Lu, Lutetium (ppm)	0.041	0.037	0.045	0.036	0.046
Mg, Magnesium (wt.%)	2.85	2.73	2.97	2.78	2.91
Mn, Manganese (wt.%)	0.013	0.012	0.013	0.012	0.013

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 2 continued.

Constituent	Certified Value	95 % Expanded Uncertainty		95 % Tolerance Limits	
		Low	High	Low	High
Aqua Regia Digestion continued					
Mo, Molybdenum (ppm)	14.2	13.7	14.7	13.8	14.5
Na, Sodium (wt.%)	0.010	0.009	0.011	IND	IND
Nd, Neodymium (ppm)	20.2	15.6	24.7	19.6	20.7
Ni, Nickel (ppm)	115	111	119	113	118
P, Phosphorus (wt.%)	0.021	0.020	0.021	0.020	0.021
Pb, Lead (ppm)	8.42	7.99	8.85	8.10	8.74
Pr, Praseodymium (ppm)	5.28	4.22	6.34	5.11	5.45
Rb, Rubidium (ppm)	6.29	5.66	6.93	5.96	6.63
Re, Rhenium (ppm)	0.026	0.023	0.029	0.024	0.028
S, Sulphur (wt.%)	3.26	3.18	3.35	3.21	3.32
Sb, Antimony (ppm)	1.86	1.71	2.00	1.76	1.95
Sc, Scandium (ppm)	3.30	3.08	3.52	3.08	3.52
Se, Selenium (ppm)	5.65	5.18	6.12	5.22	6.09
Sm, Samarium (ppm)	3.20	2.67	3.73	3.08	3.32
Sn, Tin (ppm)	1.28	1.19	1.37	1.21	1.35
Sr, Strontium (ppm)	16.4	15.1	17.8	15.7	17.1
Ta, Tantalum (ppm)	< 0.01	IND	IND	IND	IND
Tb, Terbium (ppm)	0.19	0.17	0.21	0.18	0.20
Te, Tellurium (ppm)	0.048	0.033	0.062	IND	IND
Th, Thorium (ppm)	6.18	5.80	6.55	5.97	6.39
U, Uranium (ppm)	3.01	2.86	3.15	2.92	3.09
V, Vanadium (ppm)	56	51	60	53	58
W, Tungsten (ppm)	0.96	0.88	1.05	0.91	1.02
Y, Yttrium (ppm)	3.10	2.88	3.32	2.96	3.23
Yb, Ytterbium (ppm)	0.28	0.22	0.34	0.26	0.31
Zn, Zinc (ppm)	13.0	12.0	14.0	11.8	14.2
Zr, Zirconium (ppm)	9.39	8.63	10.15	8.97	9.81

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 553b.

Constituent	Unit	Value	Constituent	Unit	Value	Constituent	Unit	Value
4-Acid Digestion								
B	ppm	20.0	Hg	ppm	0.058			
Ge	ppm	0.27	Pt	ppb	7.33			
Borate Fusion XRF								
Ag	ppm	0.032	Hf	ppm	42.5	Se	ppm	3.83
As	ppm	97	Ho	ppm	1.38	Sm	ppm	8.75
Be	ppm	5.00	In	ppm	2.20	Sn	ppm	33.0
Bi	ppm	< 100	La	ppm	81	Ta	ppm	69
Cd	ppm	< 10	Lu	ppm	0.55	Tb	ppm	1.20
Ce	ppm	151	Mo	ppm	24.4	Te	ppm	0.075
Cl	ppm	413	Na ₂ O	wt. %	0.074	Th	ppm	15.7
Cr ₂ O ₃	ppm	143	Nb	ppm	44.6	Tl	ppm	0.57
Cs	ppm	2.05	Nd	ppm	63	Tm	ppm	0.55
Dy	ppm	6.81	Pb	ppm	< 50	U	ppm	46.7
Er	ppm	4.08	Pr	ppm	18.0	W	ppm	4.85
Eu	ppm	0.94	Rb	ppm	67	Y	ppm	49.0
F	ppm	< 5000	Re	ppm	< 0.1	Yb	ppm	3.75
Ga	ppm	27.3	Sb	ppm	30.4	Zn	ppm	< 50
Gd	ppm	7.01	Sc	ppm	14.7			
Peroxide Fusion ICP continued								
Ag	ppm	1.56	Na	wt. %	0.067	Zn	ppm	27.2
Cd	ppm	0.33	Se	ppm	9.75	Zr	ppm	241
Hf	ppm	7.97	Sn	ppm	2.86			
Aqua Regia Digestion								
Ge	ppm	0.12	Pt	ppb	3.00	Tm	ppm	0.059
Nb	ppm	0.086	Ti	wt. %	0.001			
Pd	ppb	< 10	Tl	ppm	0.23			
3-Acid Digestion (no HF)								
As	ppm	< 10	Ni	ppm	102	V	ppm	102
Co	wt. %	0.564	Pb	ppm	157	Zn	ppm	103
Cu	wt. %	3.60	S	wt. %	3.32			

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 553b-DataPack.1.0.240828_101056.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 $\pm 3SD$ (magenta) and $\pm 5\%$ (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 553b 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, carrollite, 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 553b 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 553b 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 553b.

Bulk Density (kg/m ³)	Moisture (wt.%)	Munsell Notation [‡]	Munsell Color [‡]
707	0.61	N3	Dark Gray

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

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 ($1-\alpha=0.99$) at least 95 % of subsamples ($p=0.95$) will have concentrations lying between 3.57 wt. % and 3.64 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 553b 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) ± 10 %.

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

Table 5. Performance Gates for OREAS 553b.

Constituent	Certified Value	Absolute Standard Deviations					Relative Standard Deviations			5 % window	
		1SD	2SD Low	2SD High	3SD Low	3SD High	1RSD	2RSD	3RSD	Low	High
4-Acid Digestion											
Ag, ppm	0.103	0.012	0.078	0.127	0.066	0.140	11.93%	23.86%	35.78%	0.098	0.108
Al, wt. %	6.41	0.205	6.00	6.82	5.79	7.02	3.20%	6.40%	9.61%	6.09	6.73
As, ppm	54	2.9	48	59	45	62	5.44%	10.88%	16.32%	51	56
Ba, ppm	262	14	235	289	221	302	5.17%	10.34%	15.51%	249	275
Be, ppm	4.71	0.187	4.33	5.08	4.14	5.27	3.98%	7.96%	11.94%	4.47	4.94
Bi, ppm	1.70	0.081	1.54	1.86	1.46	1.94	4.74%	9.47%	14.21%	1.61	1.78
Ca, wt. %	0.313	0.011	0.291	0.334	0.281	0.345	3.44%	6.88%	10.32%	0.297	0.328
Cd, ppm	0.30	0.016	0.26	0.33	0.25	0.34	5.41%	10.82%	16.24%	0.28	0.31
Ce, ppm	144	15	115	173	100	188	10.11%	20.22%	30.33%	137	151
Co, wt. %	0.587	0.012	0.563	0.612	0.550	0.624	2.09%	4.17%	6.26%	0.558	0.616
Cr, ppm	91	10	70	111	60	122	11.39%	22.79%	34.18%	86	95
Cs, ppm	2.08	0.106	1.87	2.29	1.76	2.40	5.09%	10.17%	15.26%	1.98	2.19

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.

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

Table 5 continued.

Constituent	Certified Value	Absolute Standard Deviations					Relative Standard Deviations			5 % window	
		1SD	2SD Low	2SD High	3SD Low	3SD High	1RSD	2RSD	3RSD	Low	High
4-Acid Digestion continued											
Cu, wt. %	3.60	0.062	3.48	3.73	3.42	3.79	1.71%	3.42%	5.13%	3.42	3.78
Dy, ppm	4.44	0.318	3.81	5.08	3.49	5.40	7.16%	14.32%	21.48%	4.22	4.66
Er, ppm	2.85	0.281	2.29	3.41	2.01	3.69	9.87%	19.73%	29.60%	2.71	2.99
Eu, ppm	0.81	0.11	0.59	1.03	0.47	1.14	13.79%	27.58%	41.37%	0.77	0.85
Fe, wt. %	3.18	0.078	3.03	3.34	2.95	3.42	2.44%	4.87%	7.31%	3.02	3.34
Ga, ppm	27.9	1.24	25.4	30.4	24.2	31.6	4.45%	8.89%	13.34%	26.5	29.3
Gd, ppm	5.01	0.63	3.75	6.26	3.12	6.89	12.55%	25.10%	37.65%	4.76	5.26
Hf, ppm	5.46	0.210	5.04	5.88	4.83	6.09	3.85%	7.69%	11.54%	5.19	5.73
Ho, ppm	0.96	0.10	0.77	1.16	0.67	1.26	10.16%	20.31%	30.47%	0.91	1.01
In, ppm	2.14	0.089	1.96	2.32	1.87	2.41	4.17%	8.34%	12.51%	2.03	2.25
K, wt. %	2.55	0.086	2.38	2.73	2.30	2.81	3.38%	6.76%	10.14%	2.43	2.68
La, ppm	68	10	49	88	39	98	14.32%	28.63%	42.95%	65	72
Li, ppm	63	2.3	58	67	56	70	3.69%	7.37%	11.06%	60	66
Lu, ppm	0.42	0.05	0.31	0.52	0.26	0.57	12.45%	24.89%	37.34%	0.40	0.44
Mg, wt. %	3.58	0.131	3.31	3.84	3.18	3.97	3.67%	7.34%	11.01%	3.40	3.76
Mn, wt. %	0.013	0.000	0.013	0.014	0.013	0.014	2.10%	4.19%	6.29%	0.013	0.014
Mo, ppm	15.1	0.67	13.7	16.4	13.1	17.1	4.42%	8.84%	13.25%	14.3	15.8
Na, wt. %	0.045	0.006	0.033	0.056	0.028	0.062	12.81%	25.61%	38.42%	0.043	0.047
Nb, ppm	17.4	3.0	11.3	23.4	8.3	26.5	17.41%	34.82%	52.23%	16.5	18.3
Nd, ppm	63	7	49	76	42	83	10.79%	21.58%	32.36%	60	66
Ni, ppm	124	3	118	131	114	135	2.77%	5.55%	8.32%	118	131
P, wt. %	0.035	0.002	0.031	0.039	0.029	0.041	5.49%	10.97%	16.46%	0.033	0.037
Pb, ppm	11.2	1.06	9.1	13.4	8.0	14.4	9.47%	18.95%	28.42%	10.7	11.8
Pr, ppm	16.9	1.20	14.4	19.3	13.2	20.5	7.14%	14.27%	21.41%	16.0	17.7
Rb, ppm	96	5.0	86	106	81	111	5.24%	10.49%	15.73%	91	101
Re, ppm	0.027	0.002	0.023	0.032	0.021	0.034	8.02%	16.04%	24.07%	0.026	0.029
S, wt. %	3.34	0.075	3.19	3.49	3.11	3.56	2.25%	4.50%	6.75%	3.17	3.50
Sb, ppm	3.01	0.34	2.33	3.70	1.99	4.04	11.37%	22.73%	34.10%	2.86	3.16
Sc, ppm	12.5	0.75	11.0	14.0	10.3	14.7	5.97%	11.94%	17.90%	11.9	13.1
Se, ppm	5.73	0.528	4.67	6.78	4.15	7.31	9.21%	18.42%	27.64%	5.44	6.02
Sm, ppm	8.69	0.89	6.90	10.47	6.00	11.37	10.30%	20.60%	30.90%	8.25	9.12
Sn, ppm	2.81	0.159	2.50	3.13	2.34	3.29	5.64%	11.27%	16.91%	2.67	2.95
Sr, ppm	70	2.8	64	76	62	79	4.06%	8.11%	12.17%	67	74
Ta, ppm	1.26	0.20	0.86	1.67	0.65	1.88	16.18%	32.36%	48.55%	1.20	1.33
Tb, ppm	0.74	0.09	0.56	0.92	0.48	1.01	11.95%	23.89%	35.84%	0.70	0.78
Te, ppm	0.063	0.011	0.042	0.085	0.031	0.095	16.98%	33.96%	50.94%	0.060	0.066
Th, ppm	15.5	0.62	14.2	16.7	13.6	17.3	4.00%	8.00%	11.99%	14.7	16.2
Ti, wt. %	0.258	0.071	0.117	0.399	0.046	0.470	27.35%	54.70%	82.05%	0.245	0.271
Tl, ppm	0.67	0.048	0.57	0.76	0.53	0.81	7.12%	14.24%	21.36%	0.64	0.70
Tm, ppm	0.42	0.08	0.27	0.57	0.19	0.65	18.03%	36.05%	54.08%	0.40	0.44
U, ppm	8.60	0.421	7.76	9.44	7.34	9.86	4.89%	9.78%	14.68%	8.17	9.03
V, ppm	348	14	321	376	307	389	3.92%	7.85%	11.77%	331	366
W, ppm	2.52	0.215	2.09	2.95	1.88	3.16	8.53%	17.06%	25.59%	2.39	2.65

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.

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

Table 5 continued.

Constituent	Certified Value	Absolute Standard Deviations					Relative Standard Deviations			5 % window	
		1SD	2SD Low	2SD High	3SD Low	3SD High	1RSD	2RSD	3RSD	Low	High
4-Acid Digestion continued											
Y, ppm	23.8	0.97	21.8	25.7	20.9	26.7	4.06%	8.13%	12.19%	22.6	25.0
Yb, ppm	2.84	0.41	2.03	3.65	1.62	4.06	14.33%	28.66%	42.99%	2.70	2.98
Zn, ppm	16.1	1.21	13.7	18.6	12.5	19.8	7.49%	14.97%	22.46%	15.3	17.0
Zr, ppm	188	10	168	208	158	218	5.30%	10.61%	15.91%	178	197
Sulphuric Acid 5% Leach											
Co, wt. %	0.127	0.008	0.111	0.143	0.103	0.151	6.22%	12.43%	18.65%	0.121	0.133
Cu, wt. %	1.07	0.079	0.91	1.22	0.83	1.30	7.37%	14.75%	22.12%	1.01	1.12
Infrared Combustion											
C, wt. %	4.51	0.138	4.23	4.79	4.09	4.92	3.07%	6.14%	9.21%	4.28	4.73
S, wt. %	3.40	0.085	3.23	3.57	3.15	3.66	2.49%	4.97%	7.46%	3.23	3.57
Borate Fusion XRF											
Al ₂ O ₃ , wt. %	12.51	0.197	12.11	12.90	11.91	13.10	1.57%	3.15%	4.72%	11.88	13.13
BaO, ppm	303	45	214	393	169	437	14.72%	29.44%	44.16%	288	319
CaO, wt. %	0.435	0.015	0.406	0.464	0.391	0.479	3.38%	6.75%	10.13%	0.413	0.457
Co, wt. %	0.587	0.017	0.553	0.620	0.536	0.637	2.86%	5.72%	8.58%	0.557	0.616
Cu, wt. %	3.62	0.069	3.48	3.76	3.41	3.83	1.91%	3.83%	5.74%	3.44	3.80
Fe ₂ O ₃ , wt. %	4.60	0.074	4.45	4.75	4.38	4.82	1.62%	3.24%	4.86%	4.37	4.83
K ₂ O, wt. %	3.04	0.091	2.86	3.22	2.77	3.32	2.99%	5.98%	8.96%	2.89	3.20
MgO, wt. %	5.94	0.127	5.69	6.19	5.56	6.32	2.13%	4.27%	6.40%	5.64	6.24
MnO, wt. %	0.017	0.004	0.009	0.025	0.005	0.029	23.83%	47.66%	71.49%	0.016	0.017
Ni, ppm	125	21	83	168	62	189	16.93%	33.86%	50.79%	119	132
P ₂ O ₅ , wt. %	0.083	0.007	0.069	0.097	0.062	0.104	8.48%	16.95%	25.43%	0.079	0.087
SiO ₂ , wt. %	53.72	0.586	52.55	54.89	51.97	55.48	1.09%	2.18%	3.27%	51.04	56.41
SO ₃ , wt. %	8.59	0.215	8.17	9.02	7.95	9.24	2.50%	4.99%	7.49%	8.16	9.02
SrO, ppm	96	21	53	138	32	159	22.11%	44.21%	66.32%	91	100
TiO ₂ , wt. %	0.791	0.015	0.761	0.822	0.746	0.837	1.91%	3.81%	5.72%	0.752	0.831
V ₂ O ₅ , ppm	614	77	460	768	384	845	12.52%	25.04%	37.55%	583	645
Zr, ppm	240	48	145	336	97	384	19.87%	39.75%	59.62%	228	252
Thermogravimetry											
LOI ¹⁰⁰⁰ , wt. %	12.72	0.344	12.03	13.41	11.69	13.76	2.71%	5.41%	8.12%	12.09	13.36
Peroxide Fusion ICP											
Al, wt. %	6.47	0.156	6.15	6.78	6.00	6.93	2.41%	4.82%	7.23%	6.14	6.79
As, ppm	53	6	40	66	34	72	12.07%	24.13%	36.20%	50	56
B, ppm	217	15	187	247	172	262	6.94%	13.89%	20.83%	206	228
Ba, ppm	264	10	244	284	234	294	3.79%	7.59%	11.38%	251	277
Be, ppm	5.42	0.70	4.03	6.81	3.33	7.51	12.84%	25.68%	38.52%	5.15	5.69
Bi, ppm	1.77	0.114	1.54	2.00	1.43	2.11	6.41%	12.83%	19.24%	1.68	1.86
Ca, wt. %	0.307	0.037	0.232	0.381	0.195	0.419	12.15%	24.30%	36.46%	0.291	0.322
Ce, ppm	152	5	142	162	137	166	3.27%	6.54%	9.81%	144	159
Co, wt. %	0.581	0.023	0.535	0.628	0.511	0.651	4.01%	8.02%	12.03%	0.552	0.610
Cr, ppm	110	16	78	143	61	160	14.85%	29.69%	44.54%	105	116
Cs, ppm	2.13	0.141	1.85	2.42	1.71	2.56	6.60%	13.21%	19.81%	2.03	2.24
Cu, wt. %	3.61	0.110	3.39	3.83	3.28	3.94	3.06%	6.12%	9.18%	3.43	3.79

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.

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

Table 5 continued.

Constituent	Certified Value	Absolute Standard Deviations					Relative Standard Deviations			5 % window	
		1SD	2SD Low	2SD High	3SD Low	3SD High	1RSD	2RSD	3RSD	Low	High
Peroxide Fusion ICP continued											
Dy, ppm	7.03	0.304	6.42	7.64	6.12	7.94	4.32%	8.64%	12.96%	6.68	7.38
Er, ppm	3.73	0.329	3.07	4.39	2.74	4.72	8.82%	17.65%	26.47%	3.55	3.92
Eu, ppm	0.91	0.09	0.72	1.10	0.63	1.19	10.39%	20.78%	31.17%	0.86	0.96
Fe, wt. %	3.17	0.092	2.98	3.35	2.89	3.44	2.89%	5.79%	8.68%	3.01	3.33
Ga, ppm	27.7	1.09	25.5	29.8	24.4	30.9	3.92%	7.85%	11.77%	26.3	29.0
Gd, ppm	6.87	0.419	6.03	7.71	5.61	8.13	6.09%	12.19%	18.28%	6.53	7.21
Ge, ppm	2.24	0.67	0.89	3.59	0.22	4.26	30.11%	60.22%	90.33%	2.13	2.35
Ho, ppm	1.35	0.107	1.14	1.57	1.03	1.67	7.95%	15.89%	23.84%	1.28	1.42
In, ppm	2.22	0.166	1.88	2.55	1.72	2.72	7.51%	15.01%	22.52%	2.11	2.33
K, wt. %	2.58	0.094	2.39	2.77	2.30	2.87	3.65%	7.30%	10.94%	2.45	2.71
La, ppm	77	4.1	69	85	65	89	5.33%	10.65%	15.98%	73	81
Li, ppm	65	4.6	56	74	51	79	7.06%	14.11%	21.17%	62	68
Lu, ppm	0.56	0.050	0.46	0.66	0.41	0.71	9.01%	18.02%	27.04%	0.53	0.59
Mg, wt. %	3.57	0.087	3.39	3.74	3.31	3.83	2.43%	4.85%	7.28%	3.39	3.75
Mn, wt. %	0.014	0.002	0.011	0.017	0.009	0.019	11.80%	23.61%	35.41%	0.013	0.015
Mo, ppm	16.0	1.47	13.0	18.9	11.6	20.4	9.22%	18.45%	27.67%	15.2	16.8
Nb, ppm	25.6	1.58	22.4	28.7	20.8	30.3	6.19%	12.39%	18.58%	24.3	26.8
Nd, ppm	65	2.1	60	69	58	71	3.24%	6.48%	9.72%	61	68
Ni, ppm	126	11	103	149	91	160	9.13%	18.25%	27.38%	119	132
P, wt. %	0.037	0.008	0.022	0.053	0.014	0.060	20.48%	40.95%	61.43%	0.035	0.039
Pb, ppm	32.8	19.5	0.0	71.9	0.0	91.4	59.53%	119.1%	178.6%	31.2	34.5
Pr, ppm	18.1	0.51	17.1	19.1	16.6	19.6	2.79%	5.58%	8.36%	17.2	19.0
Rb, ppm	96	4.1	88	104	84	108	4.22%	8.44%	12.66%	91	101
Re, ppm	< 0.1	IND	IND	IND	IND	IND	IND	IND	IND	IND	IND
S, wt. %	3.29	0.188	2.92	3.67	2.73	3.86	5.70%	11.41%	17.11%	3.13	3.46
Sb, ppm	3.35	0.274	2.80	3.90	2.53	4.17	8.17%	16.34%	24.51%	3.18	3.52
Sc, ppm	12.2	0.99	10.3	14.2	9.3	15.2	8.10%	16.20%	24.30%	11.6	12.9
Si, wt. %	25.17	0.789	23.59	26.75	22.80	27.54	3.13%	6.27%	9.40%	23.91	26.43
Sm, ppm	8.90	0.663	7.58	10.23	6.91	10.89	7.45%	14.89%	22.34%	8.46	9.35
Sr, ppm	70	3.6	62	77	59	81	5.22%	10.44%	15.66%	66	73
Ta, ppm	2.01	0.30	1.42	2.60	1.12	2.90	14.74%	29.48%	44.23%	1.91	2.11
Tb, ppm	1.18	0.068	1.05	1.32	0.98	1.39	5.71%	11.42%	17.13%	1.12	1.24
Th, ppm	16.0	0.39	15.2	16.7	14.8	17.1	2.46%	4.93%	7.39%	15.2	16.8
Ti, wt. %	0.461	0.017	0.427	0.495	0.410	0.512	3.70%	7.41%	11.11%	0.438	0.484
Tl, ppm	0.73	0.065	0.60	0.86	0.53	0.92	8.90%	17.79%	26.69%	0.69	0.77
Tm, ppm	0.57	0.052	0.47	0.67	0.41	0.73	9.08%	18.15%	27.23%	0.54	0.60
U, ppm	8.70	0.411	7.88	9.52	7.47	9.93	4.73%	9.45%	14.18%	8.26	9.13
V, ppm	365	16	333	397	317	413	4.37%	8.74%	13.12%	347	383
W, ppm	3.18	0.49	2.19	4.17	1.69	4.66	15.59%	31.17%	46.76%	3.02	3.33
Y, ppm	35.6	1.89	31.8	39.4	29.9	41.3	5.31%	10.63%	15.94%	33.8	37.4
Yb, ppm	3.63	0.348	2.94	4.33	2.59	4.68	9.58%	19.17%	28.75%	3.45	3.82

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

IND = indeterminate.

Note 1: intervals may appear asymmetric due to rounding.

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

Table 5 continued.

Constituent	Certified Value	Absolute Standard Deviations					Relative Standard Deviations			5 % window	
		1SD	2SD Low	2SD High	3SD Low	3SD High	1RSD	2RSD	3RSD	Low	High
Aqua Regia Digestion											
Ag, ppm	0.077	0.008	0.060	0.094	0.051	0.102	11.02%	22.03%	33.05%	0.073	0.081
Al, wt. %	0.940	0.135	0.670	1.209	0.535	1.344	14.34%	28.69%	43.03%	0.893	0.987
As, ppm	51	2.5	46	56	44	59	4.92%	9.84%	14.76%	49	54
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	49.9	6.1	37.8	62.0	31.7	68.1	12.15%	24.30%	36.45%	47.4	52.4
Be, ppm	2.16	0.189	1.78	2.53	1.59	2.72	8.76%	17.52%	26.28%	2.05	2.26
Bi, ppm	1.58	0.111	1.36	1.80	1.25	1.91	7.02%	14.03%	21.05%	1.50	1.66
Ca, wt. %	0.300	0.013	0.274	0.327	0.260	0.340	4.43%	8.87%	13.30%	0.285	0.315
Cd, ppm	0.30	0.023	0.25	0.35	0.23	0.37	7.86%	15.73%	23.59%	0.28	0.31
Ce, ppm	42.9	6.2	30.4	55.3	24.2	61.6	14.55%	29.09%	43.64%	40.7	45.0
Co, wt. %	0.548	0.053	0.441	0.654	0.388	0.708	9.74%	19.47%	29.21%	0.520	0.575
Cr, ppm	33.9	3.31	27.3	40.5	24.0	43.9	9.77%	19.54%	29.31%	32.2	35.6
Cs, ppm	0.29	0.08	0.14	0.45	0.06	0.53	27.07%	54.13%	81.20%	0.28	0.31
Cu, wt. %	3.60	0.077	3.45	3.76	3.37	3.84	2.15%	4.30%	6.44%	3.42	3.78
Dy, ppm	0.80	0.08	0.63	0.97	0.55	1.05	10.48%	20.96%	31.44%	0.76	0.84
Er, ppm	0.36	0.030	0.30	0.42	0.27	0.45	8.16%	16.33%	24.49%	0.35	0.38
Eu, ppm	0.31	0.04	0.24	0.39	0.21	0.42	11.24%	22.48%	33.72%	0.30	0.33
Fe, wt. %	2.97	0.111	2.75	3.19	2.63	3.30	3.75%	7.50%	11.25%	2.82	3.12
Ga, ppm	7.67	1.29	5.09	10.26	3.79	11.56	16.86%	33.72%	50.58%	7.29	8.06
Gd, ppm	1.74	0.23	1.28	2.19	1.06	2.41	13.02%	26.04%	39.07%	1.65	1.82
Hf, ppm	0.30	0.024	0.25	0.35	0.23	0.37	7.82%	15.65%	23.47%	0.29	0.32
Hg, ppm	0.050	0.012	0.026	0.075	0.013	0.087	24.48%	48.96%	73.45%	0.048	0.053
Ho, ppm	0.14	0.03	0.09	0.19	0.06	0.22	18.87%	37.74%	56.61%	0.13	0.15
In, ppm	1.54	0.108	1.33	1.76	1.22	1.87	7.02%	14.04%	21.06%	1.47	1.62
K, wt. %	0.190	0.017	0.157	0.224	0.140	0.240	8.81%	17.63%	26.44%	0.181	0.200
La, ppm	19.0	2.4	14.1	23.8	11.7	26.2	12.75%	25.50%	38.25%	18.0	19.9
Li, ppm	35.5	6.3	22.9	48.0	16.7	54.3	17.67%	35.34%	53.01%	33.7	37.2
Lu, ppm	0.041	0.002	0.036	0.046	0.034	0.049	6.09%	12.18%	18.26%	0.039	0.043
Mg, wt. %	2.85	0.209	2.43	3.27	2.22	3.47	7.32%	14.65%	21.97%	2.71	2.99
Mn, wt. %	0.013	0.001	0.011	0.014	0.010	0.015	6.10%	12.20%	18.30%	0.012	0.013
Mo, ppm	14.2	0.61	13.0	15.4	12.4	16.0	4.29%	8.58%	12.87%	13.5	14.9
Na, wt. %	0.010	0.001	0.008	0.011	0.008	0.012	7.47%	14.93%	22.40%	0.009	0.010
Nd, ppm	20.2	3.7	12.7	27.6	9.0	31.3	18.50%	36.99%	55.49%	19.1	21.2
Ni, ppm	115	4	107	124	102	129	3.82%	7.64%	11.46%	110	121
P, wt. %	0.021	0.001	0.018	0.023	0.016	0.025	6.75%	13.49%	20.24%	0.020	0.022
Pb, ppm	8.42	0.837	6.75	10.09	5.91	10.93	9.94%	19.89%	29.83%	8.00	8.84
Pr, ppm	5.28	0.89	3.50	7.06	2.61	7.95	16.88%	33.76%	50.64%	5.01	5.54
Rb, ppm	6.29	1.14	4.01	8.57	2.87	9.72	18.12%	36.23%	54.35%	5.98	6.61
Re, ppm	0.026	0.001	0.023	0.029	0.022	0.030	5.63%	11.27%	16.90%	0.025	0.027
S, wt. %	3.26	0.103	3.06	3.47	2.95	3.57	3.17%	6.34%	9.51%	3.10	3.43
Sb, ppm	1.86	0.23	1.40	2.31	1.17	2.54	12.27%	24.54%	36.82%	1.76	1.95
Sc, ppm	3.30	0.36	2.58	4.02	2.22	4.38	10.91%	21.81%	32.72%	3.13	3.46

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.

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

Table 5 continued.

Constituent	Certified Value	Absolute Standard Deviations					Relative Standard Deviations			5 % window	
		1SD	2SD Low	2SD High	3SD Low	3SD High	1RSD	2RSD	3RSD	Low	High
Aqua Regia Digestion continued											
Se, ppm	5.65	0.367	4.92	6.39	4.55	6.75	6.49%	12.98%	19.47%	5.37	5.94
Sm, ppm	3.20	0.46	2.27	4.12	1.81	4.58	14.47%	28.94%	43.41%	3.04	3.36
Sn, ppm	1.28	0.070	1.14	1.42	1.07	1.49	5.46%	10.93%	16.39%	1.21	1.34
Sr, ppm	16.4	2.5	11.5	21.3	9.0	23.8	15.05%	30.09%	45.14%	15.6	17.2
Ta, ppm	< 0.01	IND	IND	IND	IND	IND	IND	IND	IND	IND	IND
Tb, ppm	0.19	0.015	0.16	0.22	0.15	0.24	7.85%	15.71%	23.56%	0.18	0.20
Te, ppm	0.048	0.012	0.024	0.071	0.013	0.083	24.42%	48.84%	73.26%	0.045	0.050
Th, ppm	6.18	0.66	4.85	7.51	4.18	8.17	10.76%	21.52%	32.28%	5.87	6.49
U, ppm	3.01	0.226	2.55	3.46	2.33	3.68	7.51%	15.01%	22.52%	2.85	3.16
V, ppm	56	9	39	73	30	82	15.38%	30.76%	46.14%	53	59
W, ppm	0.96	0.067	0.83	1.10	0.76	1.17	6.93%	13.86%	20.79%	0.92	1.01
Y, ppm	3.10	0.38	2.34	3.85	1.97	4.22	12.14%	24.28%	36.42%	2.94	3.25
Yb, ppm	0.28	0.04	0.20	0.36	0.17	0.40	13.69%	27.38%	41.07%	0.27	0.30
Zn, ppm	13.0	1.23	10.5	15.5	9.3	16.7	9.47%	18.94%	28.42%	12.4	13.7
Zr, ppm	9.39	1.35	6.69	12.10	5.33	13.45	14.40%	28.80%	43.21%	8.92	9.86

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.

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. ARGTEST 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 does not correspond with the Lab ID numbering on the scatter plots below.

Figure 1. Co by 4-Acid Digestion in OREAS 553b

SPC.1904.RR1.OREAS 553b.1.4-Acid.Co.Lab.240826.094634.SS

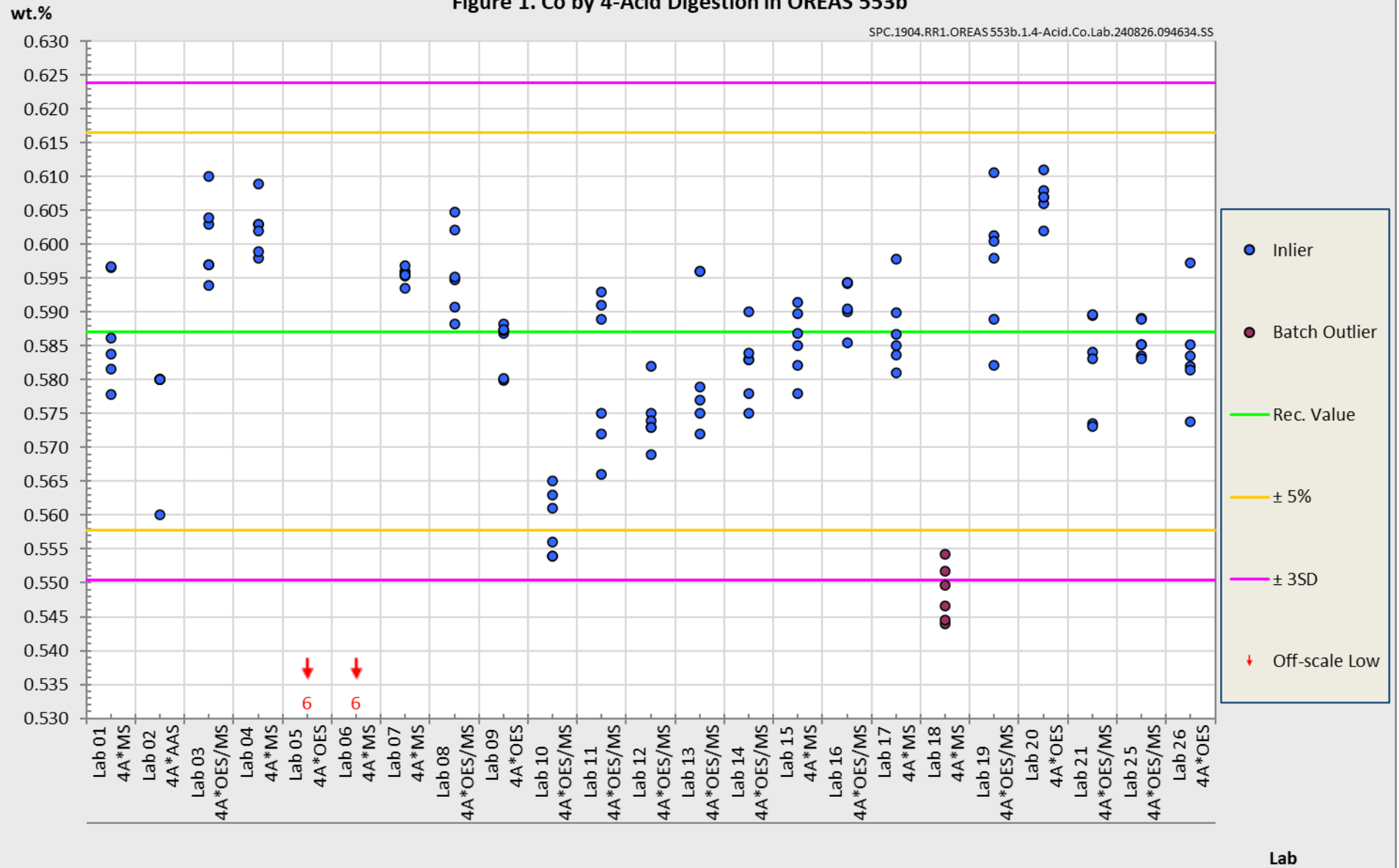


Figure 2. Cu by 4-Acid Digestion in OREAS 553b

SPC.1904.RR1.OREAS 553b.1.4-Acid.Cu.Lab.240826.095127.SN

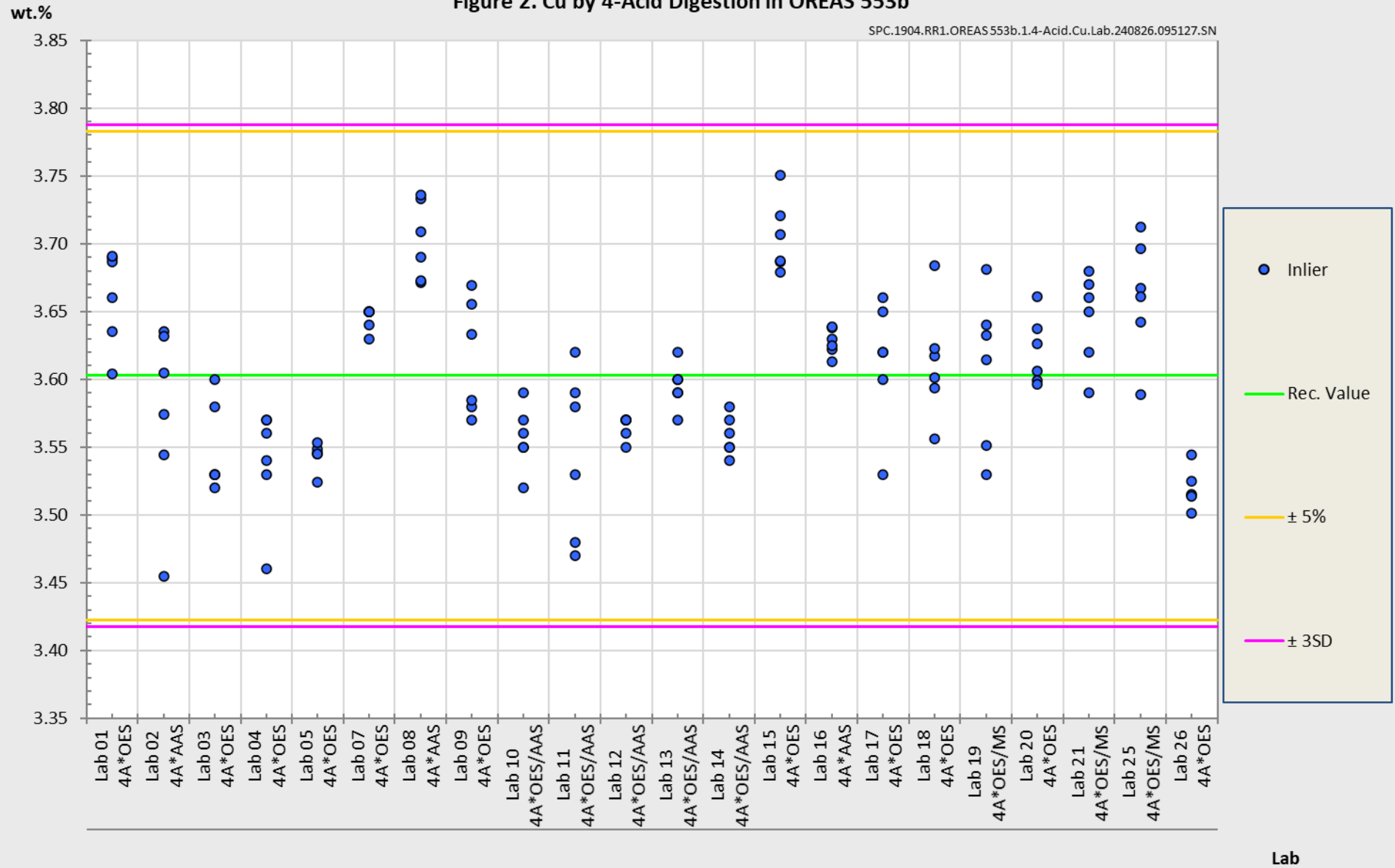


Figure 3. Co by Borate Fusion XRF in OREAS 553b

SPC.1904.RR1.OREAS 553b.1.Fusion XRF.Co.Lab.240828.010850.SS

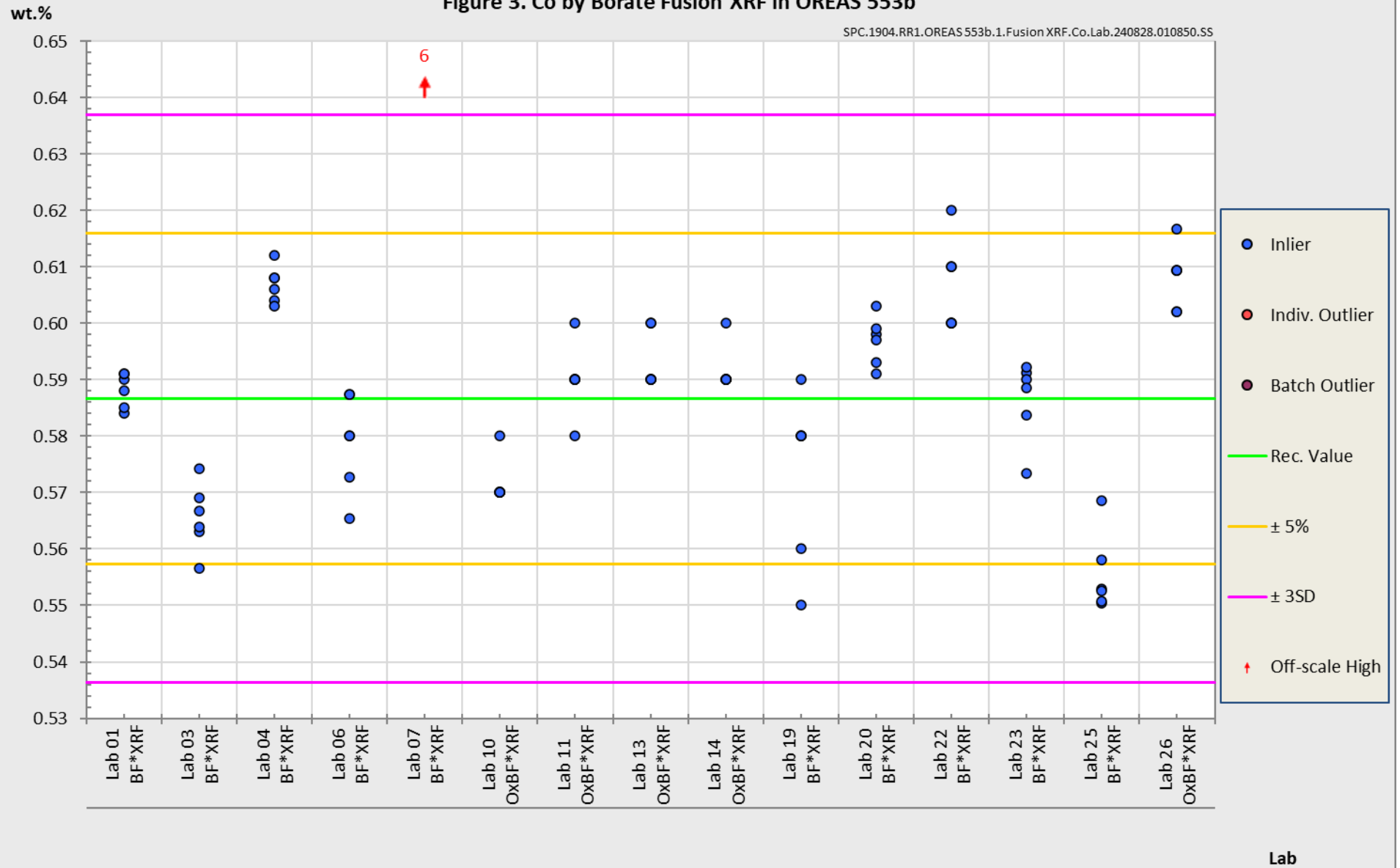


Figure 4. Cu by Borate Fusion XRF in OREAS 553b

SPC.1904.RR1.OREAS 553b.1.Fusion XRF.Cu.Lab.240919.214205.SS

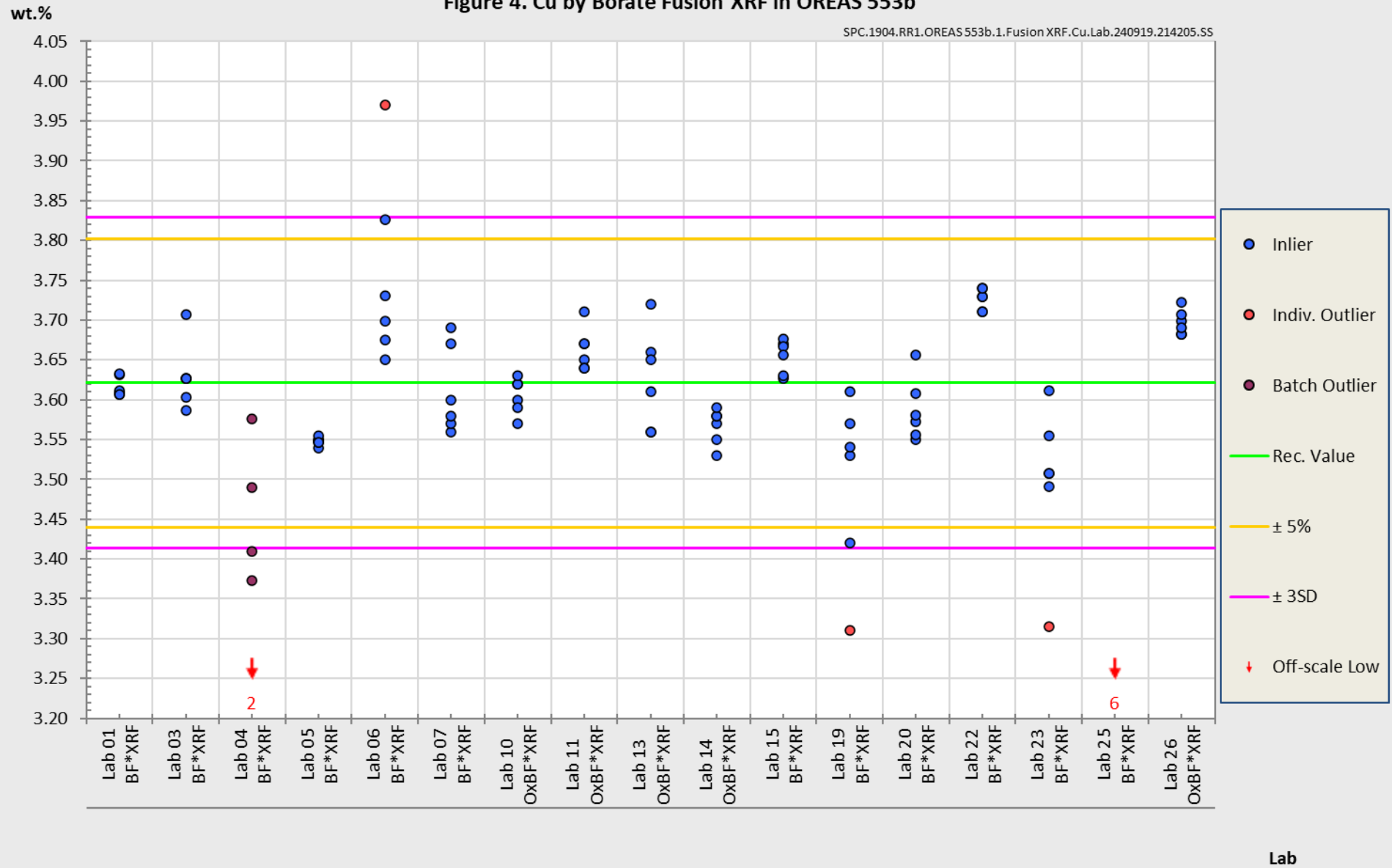


Figure 5. Co by Peroxide Fusion ICP in OREAS 553b

SPC.1904.RR1.OREAS 553b.1.PF ICP.Co.Lab.240826.095012.SS

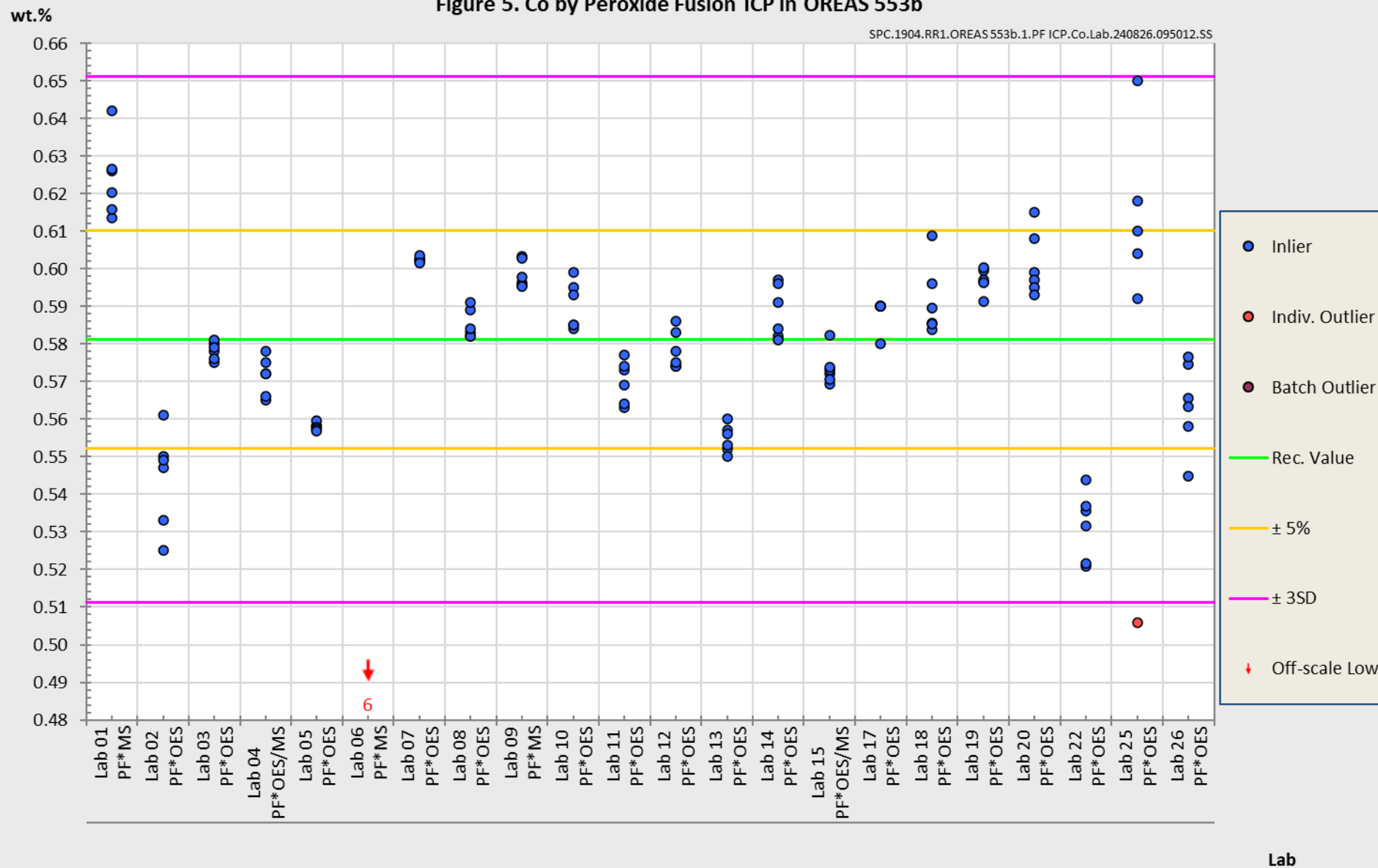
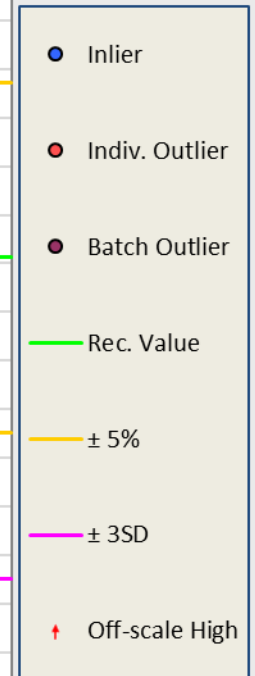
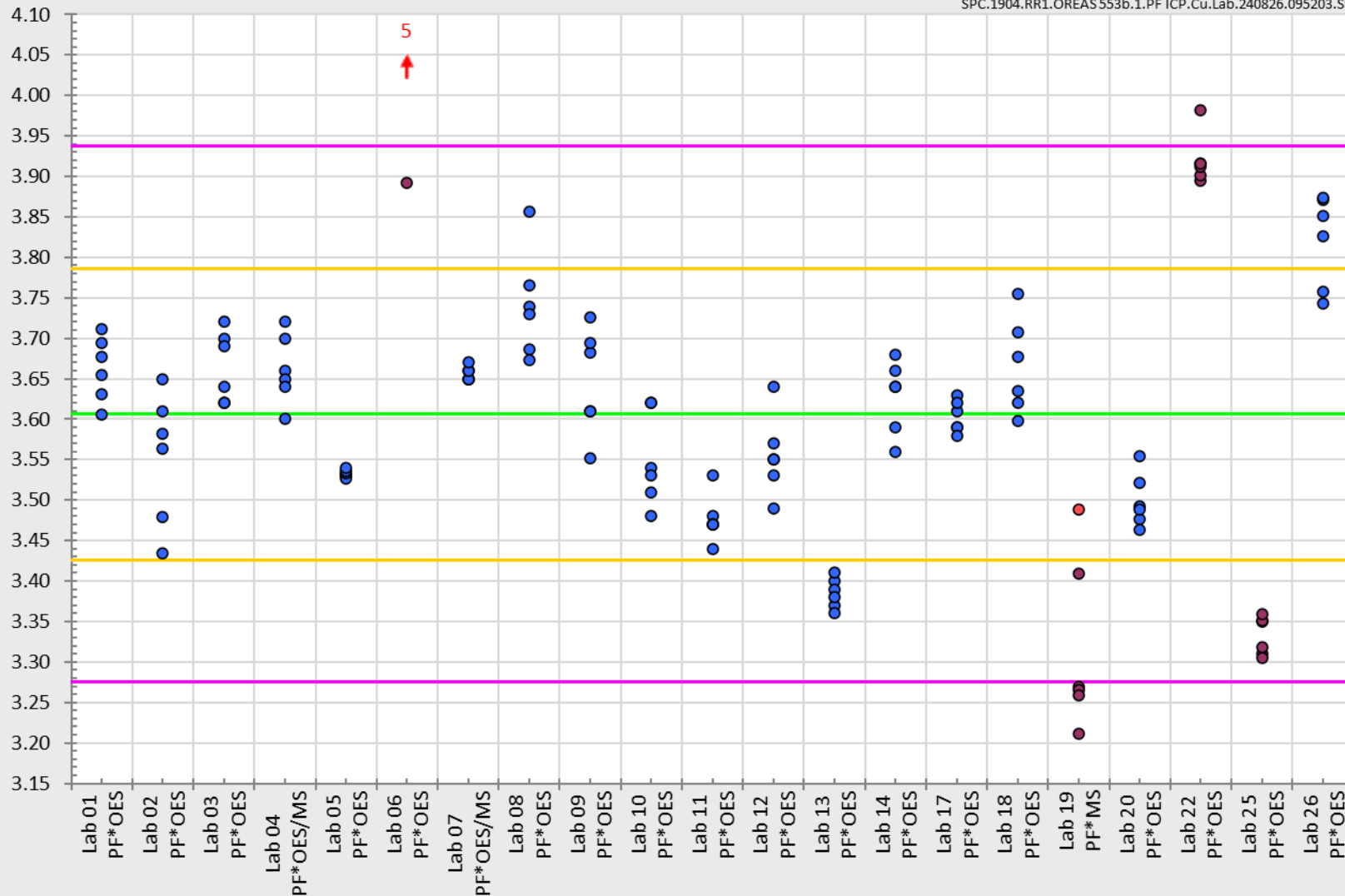


Figure 6. Cu by Peroxide Fusion ICP in OREAS 553b

SPC.1904.RR1.OREAS 553b.1.PF ICP.Cu.Lab.240826.095203.SS

wt. %



Lab

Figure 7. Co by Aqua Regia Digestion in OREAS 553b

SPC.1904.RR1.OREAS 553b.1.Aqua Regia.Co.Lab.240828.011100.SN

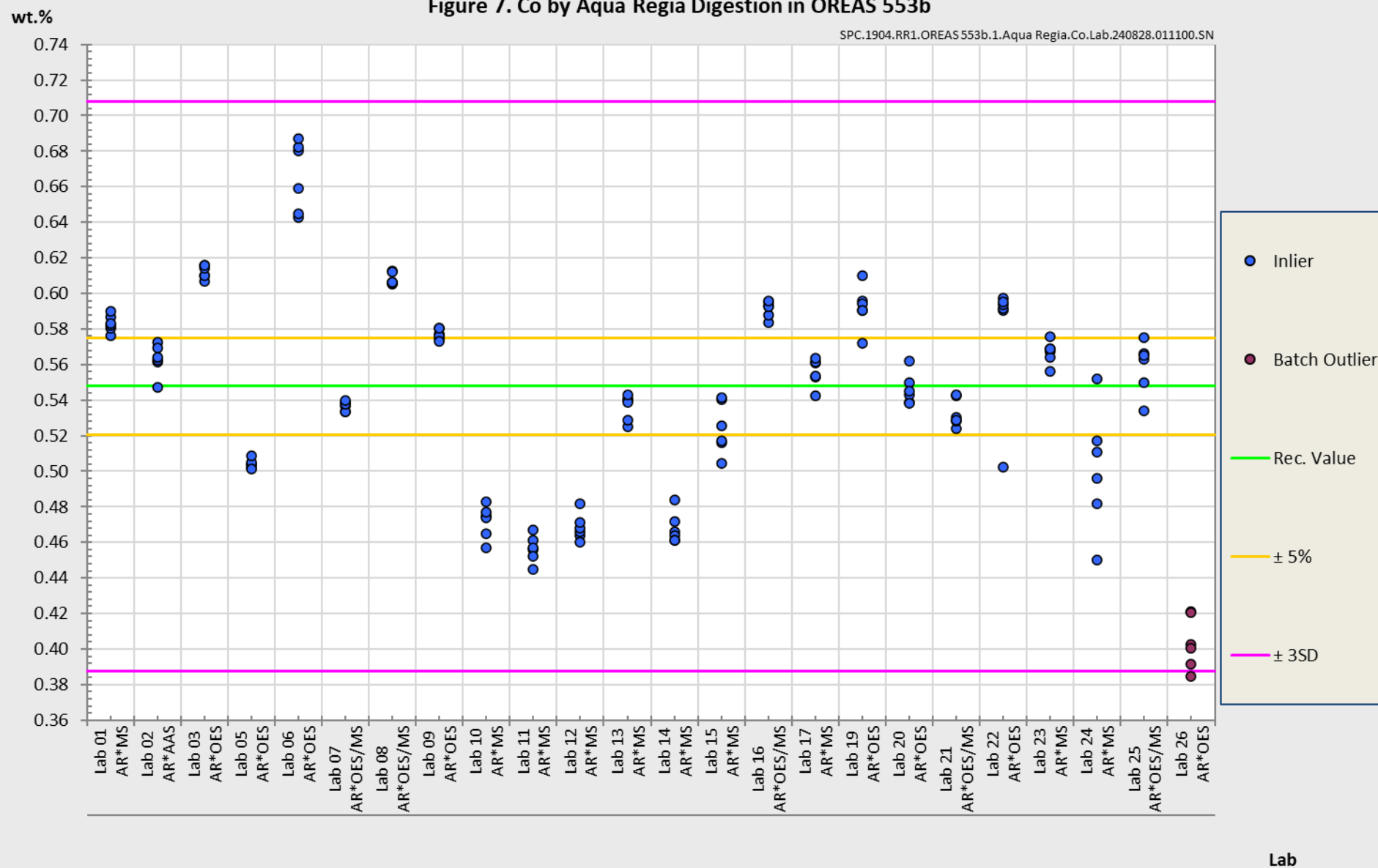
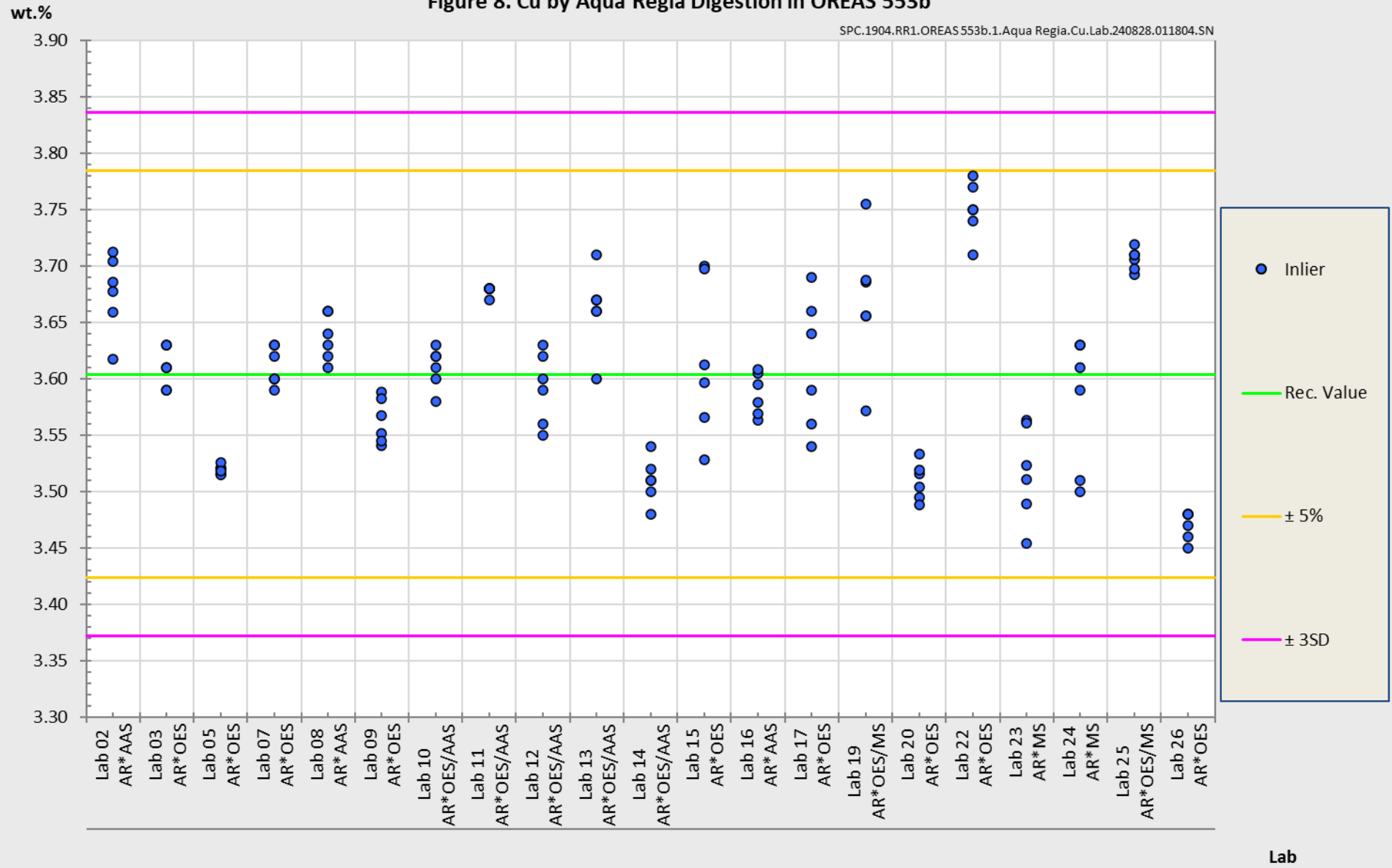


Figure 8. Cu by Aqua Regia Digestion in OREAS 553b

SPC.1904.RR1.OREAS 553b.1.Aqua Regia.Cu.Lab.240828.011804.SN



PREPARER AND SUPPLIER

Certified reference material OREAS 553b is prepared, certified and supplied by:



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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 553b 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 553b may be used to calibrate the entire procedure by producing a pure substance CRM transformed into a calibration solution.

OREAS 553b 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: ≥ 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 553b 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, well-ventilated 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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0	12 th November, 2024	First publication.

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