

**CERTIFICATE OF ANALYSIS FOR**

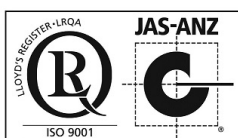
**CERTIFIED REFERENCE MATERIAL**

**OREAS 753b**

**(Pegmatite Li Ore, Western Australia)**



Accredited for compliance with ISO 17034



COA-1862-OREA753b-R1  
BUP-70-10-01 Ver:2.0

28-Oct-2024

**Table 1. Certified Values, Uncertainty & Tolerance Intervals for multi-elements by 4-acid digestion in OREAS 753b.**

Constituent	Certified Value <sup>†</sup>	95 % Expanded Uncertainty		95 % Tolerance Limits	
		Low	High	Low	High
<b>4-Acid Digestion</b>					
Ag, Silver (ppm)	0.273	0.240	0.306	0.251	0.295
Al, Aluminium (wt.%)	6.01	5.78	6.25	5.86	6.16
As, Arsenic (ppm)	4.75	4.46	5.03	4.37	5.12
Ba, Barium (ppm)	130	126	135	127	134
Be, Beryllium (ppm)	11.7	11.3	12.1	11.4	12.0
Bi, Bismuth (ppm)	0.72	0.67	0.77	0.68	0.76
Ca, Calcium (wt.%)	0.228	0.218	0.237	0.221	0.235
Cd, Cadmium (ppm)	0.14	0.11	0.17	IND	IND
Ce, Cerium (ppm)	7.92	7.08	8.76	7.44	8.41
Co, Cobalt (ppm)	1.38	1.30	1.47	1.34	1.43
Cr, Chromium (ppm)	17.0	14.8	19.1	16.0	17.9
Cs, Caesium (ppm)	170	163	176	166	173
Cu, Copper (ppm)	18.0	17.3	18.8	17.4	18.7
Dy, Dysprosium (ppm)	0.46	0.40	0.52	0.44	0.48
Er, Erbium (ppm)	0.21	0.17	0.24	0.19	0.22
Eu, Europium (ppm)	0.15	0.13	0.17	IND	IND
Fe, Iron (wt.%)	0.615	0.596	0.633	0.599	0.631
Ga, Gallium (ppm)	46.5	44.6	48.4	45.3	47.7
Gd, Gadolinium (ppm)	0.71	0.65	0.78	0.65	0.78
Ge, Germanium (ppm)	0.077	0.042	0.112	IND	IND
Hf, Hafnium (ppm)	1.86	1.77	1.96	1.79	1.94
Ho, Holmium (ppm)	0.076	0.056	0.096	IND	IND
K, Potassium (wt.%)	0.938	0.913	0.964	0.920	0.957
La, Lanthanum (ppm)	3.80	3.32	4.28	3.49	4.11
Li, Lithium (wt.%)	1.03	1.01	1.06	1.02	1.05
Li <sub>2</sub> O, Lithium oxide (wt.%)	2.23	2.18	2.27	2.20	2.25
Lu, Lutetium (ppm)	0.025	0.015	0.035	IND	IND
Mg, Magnesium (wt.%)	0.120	0.114	0.127	0.118	0.123
Mn, Manganese (wt.%)	0.133	0.129	0.137	0.130	0.136
Mo, Molybdenum (ppm)	3.36	3.21	3.52	3.24	3.48
Na, Sodium (wt.%)	0.412	0.401	0.424	0.403	0.422
Nb, Niobium (ppm)	102	98	106	99	105
Nd, Neodymium (ppm)	3.94	3.57	4.32	3.65	4.24
Ni, Nickel (ppm)	8.32	7.91	8.73	8.04	8.60
P, Phosphorus (wt.%)	0.049	0.047	0.051	0.048	0.050

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

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

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

**Table 1 continued.**

Constituent	Certified Value <sup>†</sup>	95 % Expanded Uncertainty		95 % Tolerance Limits	
		Low	High	Low	High
<b>4-Acid Digestion continued</b>					
Pb, Lead (ppm)	4.64	4.41	4.86	4.42	4.86
Pr, Praseodymium (ppm)	1.07	0.97	1.18	0.96	1.19
Rb, Rubidium (ppm)	1215	1171	1259	1191	1239
Re, Rhenium (ppm)	< 0.002	IND	IND	IND	IND
S, Sulphur (wt.%)	0.025	0.019	0.031	0.024	0.026
Sb, Antimony (ppm)	0.64	0.60	0.68	0.61	0.67
Sc, Scandium (ppm)	1.03	0.91	1.14	0.93	1.13
Sm, Samarium (ppm)	0.82	0.75	0.88	0.71	0.92
Sn, Tin (ppm)	109	104	115	106	113
Sr, Strontium (ppm)	38.3	36.6	39.9	37.1	39.4
Ta, Tantalum (ppm)	283	267	298	275	291
Tb, Terbium (ppm)	0.10	0.09	0.11	IND	IND
Te, Tellurium (ppm)	0.071	0.041	0.101	IND	IND
Th, Thorium (ppm)	1.92	1.72	2.12	1.80	2.04
Ti, Titanium (wt.%)	0.052	0.050	0.054	0.051	0.054
Tl, Thallium (ppm)	8.73	8.37	9.08	8.52	8.94
U, Uranium (ppm)	1.86	1.66	2.07	1.78	1.95
V, Vanadium (ppm)	8.11	7.65	8.57	7.73	8.50
W, Tungsten (ppm)	3.37	3.03	3.70	3.09	3.65
Y, Yttrium (ppm)	1.97	1.81	2.13	1.84	2.10
Yb, Ytterbium (ppm)	0.18	0.16	0.20	IND	IND
Zn, Zinc (ppm)	169	164	175	166	173
Zr, Zirconium (ppm)	16.2	14.9	17.6	15.4	17.1

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

Note: intervals may appear asymmetric due to rounding.

<sup>†</sup>The 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 Values, Uncertainty & Tolerance Intervals for other measurands in OREAS 753b.**

Constituent	Certified Value	95 % Expanded Uncertainty		95 % Tolerance Limits	
		Low	High	Low	High
<b>Borate / Peroxide Fusion ICP</b>					
Al, Aluminium (wt.%)	6.30	6.17	6.43	6.19	6.40
B, Boron (ppm)	80	65	95	65	95
Ba, Barium (ppm)	131	125	137	127	136
Be, Beryllium (ppm)	12.0	10.9	13.0	IND	IND
Bi, Bismuth (ppm)	0.75	0.56	0.93	IND	IND
Ca, Calcium (wt.%)	0.224	0.199	0.248	0.207	0.240
Ce, Cerium (ppm)	8.89	8.17	9.60	8.49	9.29
Co, Cobalt (ppm)	1.58	0.96	2.20	IND	IND
Cs, Caesium (ppm)	168	162	173	165	170
Cu, Copper (ppm)	20.9	18.3	23.5	IND	IND
Dy, Dysprosium (ppm)	0.69	0.57	0.80	0.61	0.76
Er, Erbium (ppm)	0.37	0.30	0.43	0.31	0.43
Eu, Europium (ppm)	0.16	0.14	0.19	IND	IND
Fe, Iron (wt.%)	0.624	0.600	0.647	0.607	0.641
Ga, Gallium (ppm)	48.5	45.6	51.4	46.9	50.1
Gd, Gadolinium (ppm)	0.82	0.63	1.00	0.72	0.91
Ge, Germanium (ppm)	2.66	2.10	3.23	IND	IND
Ho, Holmium (ppm)	0.13	0.10	0.16	IND	IND
In, Indium (ppm)	< 0.2	IND	IND	IND	IND
K, Potassium (wt.%)	0.966	0.927	1.005	0.933	0.999
La, Lanthanum (ppm)	4.55	4.18	4.91	4.21	4.88
Li, Lithium (wt.%)	1.04	1.02	1.07	1.02	1.06
Li <sub>2</sub> O, Lithium oxide (wt.%)	2.24	2.19	2.30	2.20	2.29
Lu, Lutetium (ppm)	0.066	0.034	0.098	IND	IND
Mg, Magnesium (wt.%)	0.128	0.120	0.136	0.124	0.131
Mn, Manganese (wt.%)	0.135	0.129	0.140	0.131	0.139
Mo, Molybdenum (ppm)	3.50	2.80	4.20	IND	IND
Nb, Niobium (ppm)	102	98	105	99	104
Nd, Neodymium (ppm)	4.02	3.56	4.48	3.69	4.35
P, Phosphorus (wt.%)	0.047	0.045	0.050	0.045	0.049
Pr, Praseodymium (ppm)	1.08	0.96	1.20	0.98	1.19
Rb, Rubidium (ppm)	1226	1165	1288	1205	1247
Re, Rhenium (ppm)	< 0.1	IND	IND	IND	IND
Sb, Antimony (ppm)	0.67	0.51	0.83	IND	IND
Si, Silicon (wt.%)	38.08	37.00	39.16	37.50	38.66
Sm, Samarium (ppm)	0.84	0.69	0.99	0.76	0.92
Sn, Tin (ppm)	162	155	168	156	167

SI unit equivalents: ppm (parts per million;  $1 \times 10^{-6}$ )  $\equiv$  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
<b>Borate / Peroxide Fusion ICP continued</b>					
Sr, Strontium (ppm)	47.4	42.9	51.8	44.9	49.8
Ta, Tantalum (ppm)	295	280	310	288	302
Tb, Terbium (ppm)	0.12	0.10	0.14	IND	IND
Th, Thorium (ppm)	2.08	1.89	2.27	1.79	2.36
Ti, Titanium (wt.%)	0.052	0.048	0.056	IND	IND
Tl, Thallium (ppm)	8.86	8.36	9.35	8.46	9.25
U, Uranium (ppm)	2.07	1.85	2.29	1.86	2.29
V, Vanadium (ppm)	10.5	8.6	12.5	IND	IND
W, Tungsten (ppm)	3.81	2.79	4.84	IND	IND
Y, Yttrium (ppm)	3.46	2.98	3.95	3.04	3.88
Yb, Ytterbium (ppm)	0.34	0.23	0.45	IND	IND
Zn, Zinc (ppm)	164	152	177	159	169
<b>Borate Fusion XRF</b>					
Al <sub>2</sub> O <sub>3</sub> , Aluminium(III) oxide (wt.%)	12.04	11.89	12.18	11.95	12.12
BaO, Barium oxide (ppm)	150	105	195	IND	IND
CaO, Calcium oxide (wt.%)	0.323	0.316	0.330	0.315	0.331
Cr <sub>2</sub> O <sub>3</sub> , Chromium(III) oxide (ppm)	< 100	IND	IND	IND	IND
Fe <sub>2</sub> O <sub>3</sub> , Iron(III) oxide (wt.%)	0.899	0.884	0.914	0.883	0.915
K <sub>2</sub> O, Potassium oxide (wt.%)	1.13	1.12	1.15	1.12	1.15
MgO, Magnesium oxide (wt.%)	0.214	0.202	0.225	0.206	0.222
MnO, Manganese oxide (wt.%)	0.178	0.171	0.184	0.172	0.183
Na <sub>2</sub> O, Sodium oxide (wt.%)	0.559	0.538	0.580	0.546	0.572
Nb, Niobium (ppm)	93	71	116	IND	IND
P <sub>2</sub> O <sub>5</sub> , Phosphorus(V) oxide (wt.%)	0.110	0.104	0.115	0.106	0.113
Rb, Rubidium (ppm)	1200	1122	1278	1130	1270
SiO <sub>2</sub> , Silicon dioxide (wt.%)	81.48	80.95	82.01	81.10	81.87
Sn, Tin (ppm)	151	103	199	IND	IND
SO <sub>3</sub> , Sulphur trioxide (wt.%)	0.057	0.046	0.067	0.050	0.064
TiO <sub>2</sub> , Titanium dioxide (wt.%)	0.087	0.079	0.095	0.084	0.090
V <sub>2</sub> O <sub>5</sub> , Vanadium(V) oxide (ppm)	< 100	IND	IND	IND	IND
Zn, Zinc (ppm)	170	146	194	IND	IND
<b>Thermogravimetry</b>					
LOI <sup>1000</sup> , Loss On Ignition @1000°C (wt.%)	0.872	0.786	0.958	0.830	0.914

SI unit equivalents: ppm (parts per million;  $1 \times 10^{-6}$ )  $\equiv$  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 753b.**

Constituent	Unit	Value	Constituent	Unit	Value	Constituent	Unit	Value
<b>4-Acid Digestion</b>								
B	ppm	4.00	In	ppm	0.023	Tm	ppm	0.031
Hg	ppm	< 1	Se	ppm	0.75			
<b>Peroxide Fusion ICP</b>								
Ag	ppm	< 10	Hg	ppm	< 5	Sc	ppm	< 5
As	ppm	4.03	Na	wt.%	0.448	Se	ppm	< 10
Cd	ppm	< 10	Ni	ppm	17.3	Te	ppm	< 1
Cr	ppm	39.2	Pb	ppm	5.89	Tm	ppm	0.047
Hf	ppm	2.61	S	wt.%	0.028	Zr	ppm	38.8
<b>Borate Fusion XRF</b>								
Ag	ppm	0.201	Hf	ppm	< 80	Sm	ppm	0.87
As	ppm	< 100	Ho	ppm	0.12	SrO	ppm	85
Be	ppm	12.2	In	ppm	< 0.1	Ta	ppm	263
Bi	ppm	< 100	La	ppm	34.5	Tb	ppm	0.13
Ce	ppm	8.58	Lu	ppm	0.050	Te	ppm	< 0.1
Co	ppm	< 100	Mo	ppm	< 50	Th	ppm	21.1
Cs	ppm	183	Nd	ppm	3.82	Tl	ppm	8.23
CuO	ppm	39.4	NiO	ppm	26.1	Tm	ppm	0.052
Dy	ppm	0.64	Pb	ppm	11.4	U	ppm	7.05
Er	ppm	0.36	Pr	ppm	1.06	W	ppm	< 10
Eu	ppm	0.17	Re	ppm	< 0.1	Y	ppm	42.0
Ga	ppm	45.6	Sb	ppm	< 50	Yb	ppm	0.35
Gd	ppm	0.81	Sc	ppm	2.67	Zr	ppm	63
H <sub>2</sub> O-	wt.%	0.180	Se	ppm	0.92			
<b>Aqua Regia Digestion</b>								
Cs	ppm	170						

SI unit equivalents: ppm (parts per million;  $1 \times 10^{-6}$ )  $\equiv$  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.

## TABLE OF CONTENTS

INTRODUCTION.....	8
SOURCE MATERIAL .....	8
COMMINUTION AND HOMOGENISATION PROCEDURES .....	8
PHYSICAL PROPERTIES.....	9
ANALYTICAL PROGRAM.....	9
STATISTICAL ANALYSIS .....	9
Certified Values and their uncertainty intervals .....	9
Indicative (uncertified) values .....	10
Homogeneity Evaluation.....	10
PERFORMANCE GATES .....	11
PARTICIPATING LABORATORIES .....	15
PREPARER AND SUPPLIER .....	15
METROLOGICAL TRACEABILITY .....	18
COMMUTABILITY.....	18
INTENDED USE.....	19
MINIMUM SAMPLE SIZE.....	19
PERIOD OF VALIDITY & STORAGE INSTRUCTIONS .....	19
INSTRUCTIONS FOR HANDLING & CORRECT USE .....	20
LEGAL NOTICE .....	20
DOCUMENT HISTORY.....	21
CERTIFYING OFFICER.....	21
QMS CERTIFICATION.....	21
REFERENCES.....	21

## LIST OF TABLES

Table 1. Certified Values, Uncertainty & Tolerance Intervals for multi-elements by 4-acid digestion in OREAS 753b.....	2
Table 2. Certified Values, Uncertainty & Tolerance Intervals for other measurands in OREAS 753b.....	4
Table 3. Indicative Values for OREAS 753b.....	6
Table 4. Physical properties of OREAS 753b.....	9
Table 5. Performance Gates for OREAS 753b.....	11

## LIST OF FIGURES

Figure 1. Li <sub>2</sub> O by 4-Acid digestion in OREAS 753b.....	16
Figure 2. Li <sub>2</sub> O by borate / peroxide fusion ICP in OREAS 753b.....	17

## 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 (PDM<sup>3</sup>) are presented in the detailed certification data for this CRM (**OREAS 753b-DataPack.1.2.241028\_151052.xlsx**). Results are also presented in scatter plots for Li<sub>2</sub>O (wt.%) by 4-acid digestion and borate/peroxide fusion with ICP in Figures 1 and 2 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 753b was prepared from a blend of spodumene concentrate derived from the processing of lithium pegmatite ores sourced from the Greenbushes area of southwest Western Australia, Londonderry lithium-pegmatite ore (containing elevated levels of lithium, rubidium, caesium, tin and tantalum), barren granodiorite and quartz. The barren granodiorite was sourced from the mafic, S-Type, Late Devonian Bulla Granodiorite complex located in northern Melbourne, Australia.

## COMMINUTION AND HOMOGENISATION PROCEDURES

The material constituting OREAS 753b was prepared in the following manner:

- Drying of ore and barren materials to constant mass at 105 °C;
- Crushing and milling of the barren materials to > 98 % minus 75 µm;
- Multi-stage milling of ore and spodumene concentrate materials to 100 % minus 30 µm;
- Check analysis of ore and spodumene concentrate for contained Li concentration;
- Blending the ore, concentrate and barren materials in appropriate proportions to achieve the desired Li grade;



- Homogenisation using OREAS' novel processing technologies;
- Packaging in 10g units in laminated foil pouches and 500 g units in plastic wide-mouth jars.

## PHYSICAL PROPERTIES

OREAS 753b 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 753b.**

Bulk Density (kg/m <sup>3</sup> )	Moisture (wt.%)	Munsell Notation <sup>‡</sup>	Munsell Color <sup>‡</sup>
764	0.43	N8	Very Light Gray

<sup>‡</sup>The Munsell Rock Color Chart helps geologists and archaeologists 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-eight commercial analytical laboratories participated in the program to certify the elements reported in Table 1 and 2. The following methods were employed:

- Lithium borate or sodium peroxide fusion with full suite ICP-OES and ICP-MS elemental packages (up to 24 laboratories depending on the element);
- 4-acid (HNO<sub>3</sub>-HF-HClO<sub>4</sub>-HCl) digestion with full suite ICP-OES and ICP-MS elemental packages (up to 26 laboratories depending on the element);
- Lithium borate fusion whole rock analysis package by X-ray fluorescence (up to 20 laboratories depending on the element);
- Thermogravimetry: Loss on Ignition (LOI) at 1000 °C (8 laboratories used a thermogravimetric analyser, 8 laboratories included LOI with their fusion package and 5 laboratories used a conventional muffle furnace).

For the round robin program twelve 1 kg 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 15 g scoop splits from each of three separate 1 kg 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) 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 lithium ( $\text{Li}_2\text{O}$ ) by 4-acid digestion with ICP, where 99 % of the time ( $1-\alpha=0.99$ ) at least 95 % of subsamples ( $p=0.95$ ) will have concentrations lying between 2.20 and 2.25 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 753b 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](http://www.westgard.com/mltirule.htm)). A second method utilises a 5 % window calculated directly from the certified value.

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

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

**Table 5. Performance Gates for OREAS 753b.**

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
<b>4-Acid Digestion</b>											
Ag, ppm	0.273	0.042	0.188	0.358	0.146	0.400	15.56%	31.11%	46.67%	0.259	0.287
Al, wt. %	6.01	0.360	5.29	6.73	4.93	7.09	5.98%	11.97%	17.95%	5.71	6.31
As, ppm	4.75	0.359	4.03	5.46	3.67	5.82	7.57%	15.15%	22.72%	4.51	4.98
Ba, ppm	130	6	119	142	113	148	4.47%	8.94%	13.41%	124	137
Be, ppm	11.7	0.60	10.5	12.9	9.9	13.5	5.13%	10.27%	15.40%	11.1	12.3
Bi, ppm	0.72	0.048	0.62	0.82	0.58	0.86	6.66%	13.32%	19.98%	0.68	0.76
Ca, wt. %	0.228	0.009	0.209	0.246	0.200	0.255	3.98%	7.96%	11.95%	0.216	0.239
Cd, ppm	0.14	0.02	0.09	0.18	0.07	0.21	17.09%	34.17%	51.26%	0.13	0.14
Ce, ppm	7.92	1.48	4.97	10.88	3.49	12.36	18.65%	37.31%	55.96%	7.53	8.32
Co, ppm	1.38	0.073	1.24	1.53	1.16	1.60	5.26%	10.53%	15.79%	1.31	1.45
Cr, ppm	17.0	3.7	9.5	24.4	5.8	28.1	21.89%	43.78%	65.68%	16.1	17.8
Cs, ppm	170	9	151	188	141	198	5.53%	11.05%	16.58%	161	178
Cu, ppm	18.0	0.87	16.3	19.8	15.4	20.6	4.80%	9.60%	14.41%	17.1	18.9
Dy, ppm	0.46	0.05	0.36	0.56	0.31	0.61	10.55%	21.11%	31.66%	0.44	0.48
Er, ppm	0.21	0.03	0.14	0.28	0.10	0.31	16.78%	33.57%	50.35%	0.20	0.22
Eu, ppm	0.15	0.015	0.12	0.18	0.11	0.20	9.59%	19.17%	28.76%	0.14	0.16
Fe, wt. %	0.615	0.027	0.562	0.668	0.535	0.694	4.32%	8.64%	12.97%	0.584	0.646
Ga, ppm	46.5	2.94	40.6	52.4	37.7	55.3	6.32%	12.65%	18.97%	44.2	48.8
Gd, ppm	0.71	0.037	0.64	0.79	0.60	0.82	5.24%	10.48%	15.72%	0.68	0.75
Ge, ppm	0.077	0.020	0.037	0.117	0.017	0.137	25.97%	51.94%	77.91%	0.073	0.081
Hf, ppm	1.86	0.101	1.66	2.06	1.56	2.16	5.41%	10.83%	16.24%	1.77	1.95
Ho, ppm	0.076	0.011	0.054	0.099	0.043	0.110	14.71%	29.43%	44.14%	0.072	0.080

SI unit equivalents: ppm (parts per million;  $1 \times 10^{-6}$ )  $\equiv$  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
<b>4-Acid Digestion continued</b>											
K, wt.%	0.938	0.040	0.857	1.019	0.817	1.060	4.32%	8.63%	12.95%	0.891	0.985
La, ppm	3.80	0.71	2.37	5.23	1.66	5.94	18.78%	37.55%	56.33%	3.61	3.99
Li, wt.%	1.03	0.026	0.98	1.09	0.96	1.11	2.48%	4.95%	7.43%	0.98	1.09
Li <sub>2</sub> O, wt.%	2.23	0.055	2.12	2.34	2.06	2.39	2.48%	4.95%	7.43%	2.12	2.34
Lu, ppm	0.025	0.006	0.014	0.036	0.008	0.042	22.58%	45.15%	67.73%	0.024	0.026
Mg, wt.%	0.120	0.007	0.106	0.135	0.099	0.142	6.00%	12.00%	18.01%	0.114	0.126
Mn, wt.%	0.133	0.005	0.124	0.142	0.119	0.146	3.42%	6.85%	10.27%	0.126	0.139
Mo, ppm	3.36	0.184	3.00	3.73	2.81	3.91	5.46%	10.92%	16.39%	3.19	3.53
Na, wt.%	0.412	0.015	0.383	0.441	0.369	0.456	3.53%	7.05%	10.58%	0.392	0.433
Nb, ppm	102	5	92	113	86	118	5.15%	10.30%	15.45%	97	107
Nd, ppm	3.94	0.246	3.45	4.44	3.21	4.68	6.23%	12.47%	18.70%	3.75	4.14
Ni, ppm	8.32	0.438	7.44	9.19	7.00	9.63	5.26%	10.53%	15.79%	7.90	8.73
P, wt.%	0.049	0.003	0.044	0.054	0.041	0.057	5.26%	10.52%	15.77%	0.047	0.052
Pb, ppm	4.64	0.255	4.13	5.15	3.87	5.40	5.49%	10.98%	16.47%	4.40	4.87
Pr, ppm	1.07	0.073	0.93	1.22	0.86	1.29	6.81%	13.61%	20.42%	1.02	1.13
Rb, ppm	1215	73	1068	1362	995	1436	6.05%	12.09%	18.14%	1155	1276
Re, ppm	< 0.002	IND	IND	IND	IND	IND	IND	IND	IND	IND	IND
S, wt.%	0.025	0.004	0.017	0.033	0.013	0.037	16.26%	32.51%	48.77%	0.024	0.026
Sb, ppm	0.64	0.042	0.56	0.73	0.51	0.77	6.60%	13.21%	19.81%	0.61	0.67
Sc, ppm	1.03	0.14	0.76	1.30	0.62	1.43	13.18%	26.35%	39.53%	0.98	1.08
Sm, ppm	0.82	0.057	0.70	0.93	0.65	0.99	6.93%	13.86%	20.79%	0.78	0.86
Sn, ppm	109	6	97	122	90	129	5.89%	11.78%	17.67%	104	115
Sr, ppm	38.3	2.33	33.6	42.9	31.2	45.3	6.10%	12.21%	18.31%	36.3	40.2
Ta, ppm	283	27	228	338	201	365	9.69%	19.38%	29.07%	269	297
Tb, ppm	0.10	0.01	0.07	0.13	0.06	0.14	14.41%	28.83%	43.24%	0.10	0.11
Te, ppm	0.071	0.015	0.042	0.101	0.027	0.116	20.78%	41.55%	62.33%	0.068	0.075
Th, ppm	1.92	0.30	1.33	2.52	1.03	2.81	15.46%	30.92%	46.39%	1.83	2.02
Ti, wt.%	0.052	0.003	0.047	0.058	0.044	0.060	5.36%	10.72%	16.07%	0.049	0.055
Tl, ppm	8.73	0.395	7.94	9.52	7.54	9.91	4.53%	9.06%	13.59%	8.29	9.16
U, ppm	1.86	0.23	1.39	2.33	1.16	2.57	12.57%	25.14%	37.72%	1.77	1.96
V, ppm	8.11	0.427	7.26	8.97	6.83	9.39	5.26%	10.52%	15.78%	7.71	8.52
W, ppm	3.37	0.36	2.64	4.09	2.28	4.45	10.73%	21.45%	32.18%	3.20	3.53
Y, ppm	1.97	0.23	1.50	2.44	1.27	2.67	11.90%	23.80%	35.70%	1.87	2.07
Yb, ppm	0.18	0.03	0.11	0.24	0.08	0.28	18.32%	36.65%	54.97%	0.17	0.19
Zn, ppm	169	9	151	188	142	197	5.33%	10.66%	15.99%	161	178
Zr, ppm	16.2	1.59	13.1	19.4	11.5	21.0	9.79%	19.58%	29.36%	15.4	17.0
<b>Borate / Peroxide Fusion ICP</b>											
Al, wt.%	6.30	0.112	6.07	6.52	5.96	6.63	1.78%	3.57%	5.35%	5.98	6.61
B, ppm	80	14	53	108	39	121	17.14%	34.29%	51.43%	76	84
Ba, ppm	131	7	117	145	110	153	5.35%	10.70%	16.05%	125	138
Be, ppm	12.0	1.04	9.9	14.1	8.9	15.1	8.69%	17.38%	26.07%	11.4	12.6
Bi, ppm	0.75	0.11	0.53	0.96	0.42	1.07	14.36%	28.72%	43.08%	0.71	0.78
Ca, wt.%	0.224	0.026	0.172	0.275	0.147	0.301	11.48%	22.97%	34.45%	0.212	0.235

SI unit equivalents: ppm (parts per million;  $1 \times 10^{-6}$ )  $\equiv$  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
<b>Borate / Peroxide Fusion ICP continued</b>											
Ce, ppm	8.89	0.94	7.01	10.77	6.07	11.70	10.57%	21.13%	31.70%	8.44	9.33
Co, ppm	1.58	0.38	0.81	2.35	0.43	2.73	24.31%	48.62%	72.93%	1.50	1.66
Cs, ppm	168	5	157	178	152	183	3.18%	6.36%	9.54%	159	176
Cu, ppm	20.9	2.6	15.7	26.1	13.1	28.7	12.48%	24.95%	37.43%	19.8	21.9
Dy, ppm	0.69	0.07	0.54	0.83	0.47	0.90	10.37%	20.75%	31.12%	0.65	0.72
Er, ppm	0.37	0.05	0.26	0.47	0.21	0.52	13.88%	27.76%	41.64%	0.35	0.38
Eu, ppm	0.16	0.02	0.12	0.21	0.10	0.23	13.97%	27.94%	41.91%	0.16	0.17
Fe, wt. %	0.624	0.022	0.579	0.668	0.557	0.691	3.58%	7.17%	10.75%	0.593	0.655
Ga, ppm	48.5	2.13	44.2	52.8	42.1	54.9	4.39%	8.79%	13.18%	46.1	50.9
Gd, ppm	0.82	0.10	0.62	1.01	0.52	1.11	11.96%	23.92%	35.88%	0.77	0.86
Ge, ppm	2.66	0.45	1.76	3.57	1.30	4.03	17.02%	34.05%	51.07%	2.53	2.80
Ho, ppm	0.13	0.02	0.09	0.17	0.07	0.19	15.97%	31.94%	47.91%	0.12	0.14
In, ppm	< 0.2	IND	IND	IND	IND	IND	IND	IND	IND	IND	IND
K, wt. %	0.966	0.053	0.861	1.071	0.808	1.124	5.46%	10.92%	16.37%	0.918	1.014
La, ppm	4.55	0.440	3.67	5.43	3.23	5.87	9.68%	19.36%	29.04%	4.32	4.77
Li, wt. %	1.04	0.034	0.98	1.11	0.94	1.14	3.23%	6.45%	9.68%	0.99	1.09
Li <sub>2</sub> O, wt. %	2.24	0.072	2.10	2.39	2.03	2.46	3.23%	6.45%	9.68%	2.13	2.36
Lu, ppm	0.066	0.017	0.031	0.101	0.014	0.118	26.40%	52.80%	79.21%	0.063	0.069
Mg, wt. %	0.128	0.007	0.114	0.142	0.107	0.148	5.40%	10.80%	16.20%	0.121	0.134
Mn, wt. %	0.135	0.005	0.125	0.145	0.119	0.150	3.80%	7.59%	11.39%	0.128	0.142
Mo, ppm	3.50	0.69	2.13	4.87	1.44	5.56	19.62%	39.24%	58.86%	3.32	3.67
Nb, ppm	102	3	95	108	91	112	3.32%	6.63%	9.95%	96	107
Nd, ppm	4.02	0.49	3.03	5.01	2.54	5.50	12.26%	24.53%	36.79%	3.82	4.22
P, wt. %	0.047	0.004	0.040	0.054	0.036	0.058	7.76%	15.53%	23.29%	0.045	0.049
Pr, ppm	1.08	0.13	0.82	1.35	0.69	1.48	12.15%	24.30%	36.45%	1.03	1.14
Rb, ppm	1226	46	1134	1319	1087	1366	3.78%	7.56%	11.34%	1165	1288
Re, ppm	< 0.1	IND	IND	IND	IND	IND	IND	IND	IND	IND	IND
Sb, ppm	0.67	0.10	0.47	0.86	0.38	0.96	14.45%	28.91%	43.36%	0.63	0.70
Si, wt. %	38.08	1.276	35.53	40.63	34.25	41.91	3.35%	6.70%	10.05%	36.17	39.98
Sm, ppm	0.84	0.11	0.61	1.07	0.50	1.18	13.46%	26.93%	40.39%	0.80	0.88
Sn, ppm	162	11	139	184	128	195	6.84%	13.67%	20.51%	153	170
Sr, ppm	47.4	6.2	35.0	59.8	28.8	66.0	13.09%	26.17%	39.26%	45.0	49.7
Ta, ppm	295	22	251	340	229	362	7.52%	15.04%	22.56%	280	310
Tb, ppm	0.12	0.02	0.09	0.16	0.08	0.17	13.07%	26.14%	39.21%	0.12	0.13
Th, ppm	2.08	0.21	1.66	2.50	1.45	2.71	10.03%	20.06%	30.09%	1.98	2.18
Ti, wt. %	0.052	0.003	0.045	0.058	0.042	0.061	6.10%	12.20%	18.30%	0.049	0.054
Tl, ppm	8.86	0.448	7.96	9.75	7.51	10.20	5.06%	10.12%	15.18%	8.41	9.30
U, ppm	2.07	0.21	1.65	2.50	1.43	2.71	10.28%	20.56%	30.84%	1.97	2.18
V, ppm	10.5	2.2	6.2	14.9	4.0	17.0	20.57%	41.13%	61.70%	10.0	11.1
W, ppm	3.81	0.64	2.53	5.10	1.89	5.74	16.79%	33.59%	50.38%	3.62	4.01
Y, ppm	3.46	0.46	2.55	4.37	2.09	4.83	13.18%	26.35%	39.53%	3.29	3.63
Yb, ppm	0.34	0.06	0.22	0.46	0.16	0.52	17.83%	35.66%	53.49%	0.32	0.35
Zn, ppm	164	14	137	192	123	206	8.38%	16.75%	25.13%	156	172

SI unit equivalents: ppm (parts per million;  $1 \times 10^{-6}$ )  $\equiv$  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
<b>Borate Fusion XRF</b>											
Al <sub>2</sub> O <sub>3</sub> , wt. %	12.04	0.203	11.63	12.44	11.43	12.64	1.68%	3.37%	5.05%	11.43	12.64
BaO, ppm	150	43	64	236	21	279	28.58%	57.17%	85.75%	143	158
CaO, wt. %	0.323	0.007	0.308	0.338	0.301	0.345	2.30%	4.61%	6.91%	0.307	0.339
Cr <sub>2</sub> O <sub>3</sub> , ppm	< 100	IND	IND	IND	IND	IND	IND	IND	IND	IND	IND
Fe <sub>2</sub> O <sub>3</sub> , wt. %	0.899	0.022	0.854	0.943	0.832	0.966	2.48%	4.95%	7.43%	0.854	0.944
K <sub>2</sub> O, wt. %	1.13	0.022	1.09	1.18	1.07	1.20	1.94%	3.88%	5.82%	1.08	1.19
MgO, wt. %	0.214	0.018	0.177	0.250	0.159	0.268	8.51%	17.01%	25.52%	0.203	0.224
MnO, wt. %	0.178	0.006	0.165	0.190	0.159	0.196	3.43%	6.86%	10.29%	0.169	0.187
Na <sub>2</sub> O, wt. %	0.559	0.029	0.502	0.617	0.473	0.645	5.15%	10.29%	15.44%	0.531	0.587
Nb, ppm	93	23	46	140	23	163	25.08%	50.17%	75.25%	89	98
P <sub>2</sub> O <sub>5</sub> , wt. %	0.110	0.006	0.098	0.121	0.092	0.127	5.26%	10.51%	15.77%	0.104	0.115
Rb, ppm	1200	53	1095	1305	1042	1358	4.39%	8.77%	13.16%	1140	1260
SiO <sub>2</sub> , wt. %	81.48	0.903	79.68	83.29	78.77	84.19	1.11%	2.22%	3.33%	77.41	85.56
Sn, ppm	151	37	77	225	39	262	24.64%	49.28%	73.92%	143	158
SO <sub>3</sub> , wt. %	0.057	0.014	0.030	0.084	0.016	0.097	23.72%	47.44%	71.16%	0.054	0.060
TiO <sub>2</sub> , wt. %	0.087	0.006	0.074	0.100	0.067	0.106	7.44%	14.87%	22.31%	0.082	0.091
V <sub>2</sub> O <sub>5</sub> , ppm	< 100	IND	IND	IND	IND	IND	IND	IND	IND	IND	IND
Zn, ppm	170	17	136	204	119	221	10.03%	20.07%	30.10%	161	178
<b>Thermogravimetry</b>											
LOI <sup>1000</sup> , wt. %	0.872	0.083	0.705	1.038	0.622	1.122	9.55%	19.10%	28.65%	0.828	0.915

SI unit equivalents: ppm (parts per million;  $1 \times 10^{-6}$ )  $\equiv$  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.

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4. ALS, Johannesburg, South Africa
5. ALS, Lima, Peru
6. ALS, Loughrea, Galway, Ireland
7. ALS, Malaga, WA, Australia
8. ALS, Vancouver, BC, Canada
9. ARGETEST Mineral Processing, Ankara, Central Anatolia, Turkey
10. Bureau Veritas Commodities Canada Ltd, Vancouver, BC, Canada
11. CERTIMIN, Lima, Peru
12. CRS Laboratories Oy, Kempele, Northern Ostrobothnia, Finland
13. Inspectorate (BV), Lima, Peru
14. Intertek, Cupang, Muntinlupa, Philippines
15. Intertek, Perth, WA, Australia
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***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.***

## PREPARER AND SUPPLIER

Certified reference material OREAS 753b is prepared, certified and supplied by:



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Figure 1. Li<sub>2</sub>O by 4-Acid Digestion in OREAS 753b

SPC.1862.RR1.OREAS 753b.1.4-Acid.Li2O.Lab.240722.234714.SN

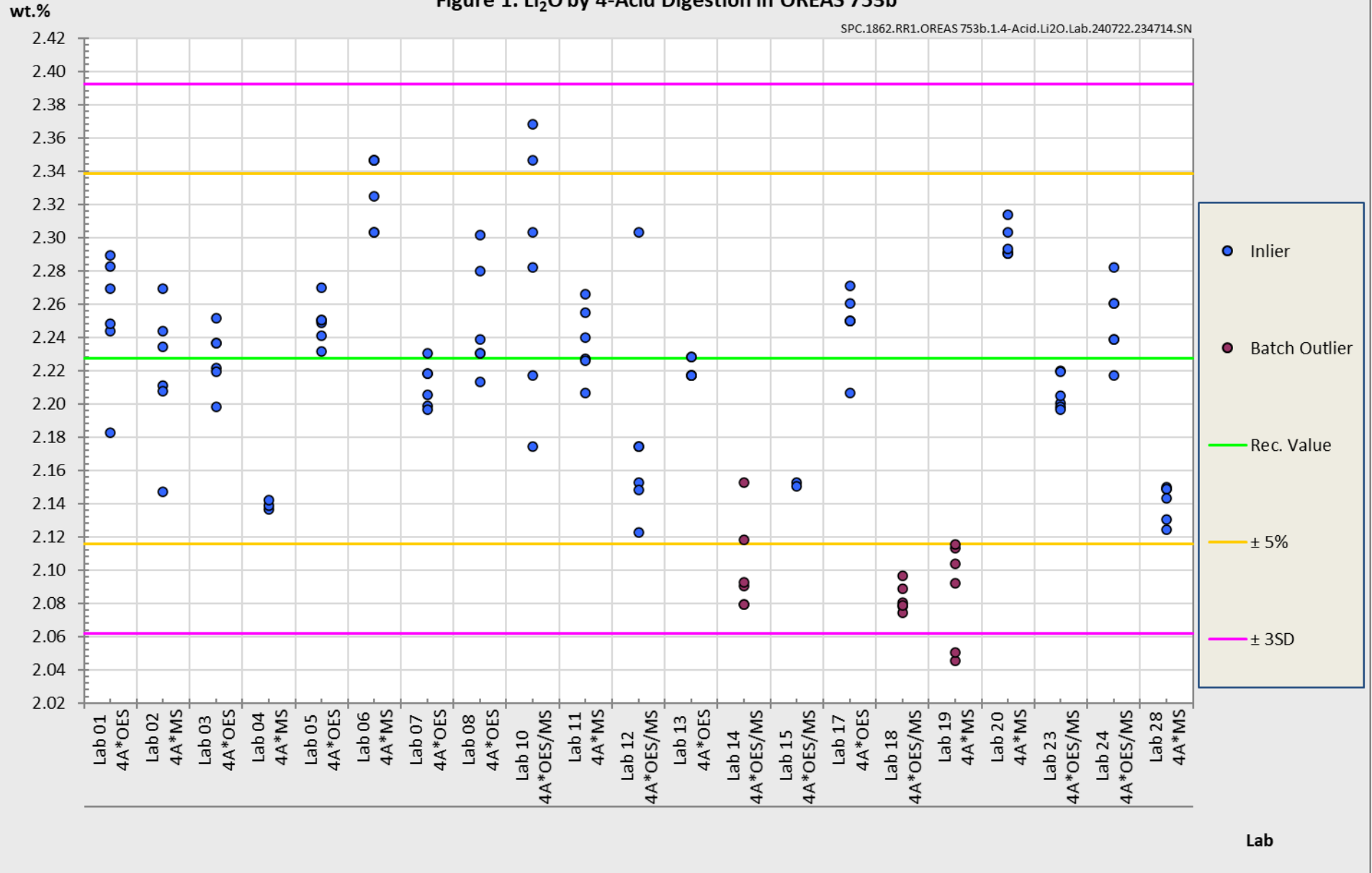
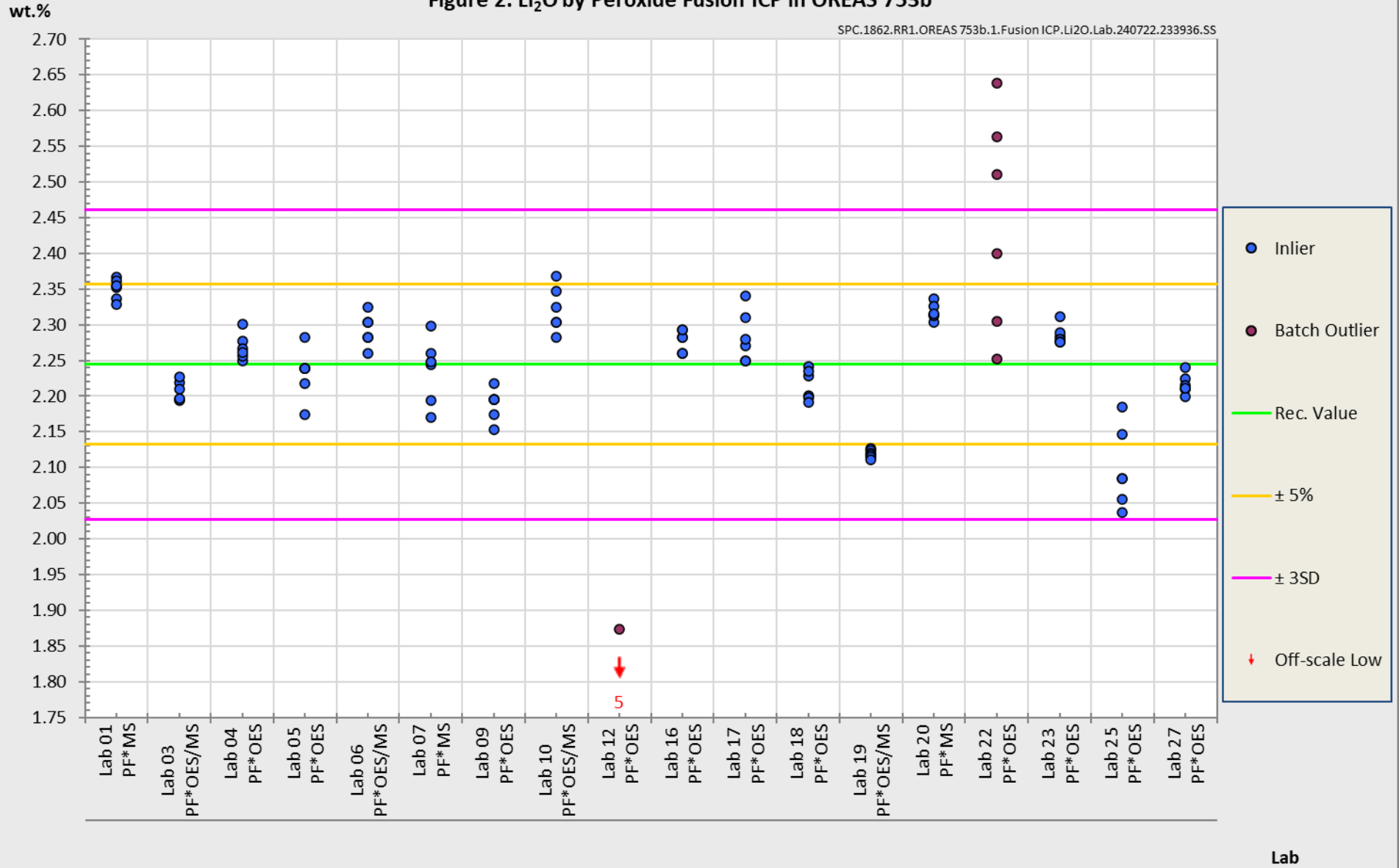




Figure 2. Li<sub>2</sub>O by Peroxide Fusion ICP in OREAS 753b

SPC.1862.RR1.OREAS 753b.1.Fusion ICP.Li2O.Lab.240722.233936.SS



## 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 753b 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 753b may be used to calibrate the entire procedure by producing a pure substance CRM transformed into a calibration solution.

OREAS 753b 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 / Sodium peroxide fusion with ICP-OES and/or MS finish:  $\geq 0.2$  g;
- Borate fusion with X-ray fluorescence finish:  $\geq 0.2$  g;
- Loss on Ignition (LOI) at 1000 °C:  $\geq 1$  g;
- Multi-elements by 4-acid digestion with ICP-OES and/or MS finish:  $\geq 0.25$  g.

## PERIOD OF VALIDITY & STORAGE INSTRUCTIONS

The certification of OREAS 753b remains valid, within the specified measurement uncertainties, until at least February 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

OREAS 753b is packaged in single-use laminated foil sachets. Following analysis, it is the manufacturer's expectation that any remaining material is discarded. It is the user's responsibility to prevent contamination and avoid prolonged exposure of the sample to the atmosphere prior to analysis.

## **Repeat-use packaging (e.g., 500 g plastic jars)**

After taking a subsample, users should replace the lid of the jar promptly and securely to prevent accidental spills and airborne contamination. OREAS 753b contains a non-hygroscopic\* matrix with an indicative value for moisture provided to enable users to check for changes to stored material by determining moisture in the user's laboratory and comparing the result to the value in Table 4 in this certificate.

The stability of the CRM in regard to oxidation from the breakdown of sulphide minerals to sulphates is negligible given its low sulphur concentration (~0.03 wt.% S).

\*A non-hygroscopic matrix means exposure to atmospheres significantly different, in terms of temperature and humidity, from the climate during manufacturing should have negligible impact on the precision of results. Hygroscopic moisture is the amount of adsorbed moisture (weakly held H<sub>2</sub>O- molecules on the surface of exposed material) following exposure to the local atmosphere. Usually, equilibration of material to the local atmosphere will only occur if the material is spread into a thin (~2mm thick) layer and left exposed for a period of 2 hours.

## **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.

## **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

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## DOCUMENT HISTORY

Revision No.	Date	Changes applied
1	28 <sup>th</sup> October, 2024	Changed Fe <sub>2</sub> O <sub>3</sub> to Fe for 4-Acid Digestion and Borate / Peroxide Fusion ICP data in Table 1, 2 & 5.
0	9 <sup>th</sup> September, 2024	First publication.

## CERTIFYING OFFICER



28<sup>th</sup> October, 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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