

## **CERTIFICATE OF ANALYSIS FOR**

# Carlin Ore (Leeville Mine, Nevada, USA) CERTIFIED REFERENCE MATERIAL OREAS 277

Table 1. Certified Values and Performance Gates for OREAS 277.

Constituent	Certified	A	Absolute \$	Standard	Deviation	าร	Relative Standard Deviations			5% window	
Constituent	Value	1SD	2SD Low	2SD High	3SD Low	3SD High	1RSD	2RSD	3RSD	Low	High
Pb Fire Assay											
Au, ppm	3.39	0.120	3.15	3.63	3.03	3.75	3.54%	7.08%	10.62%	3.22	3.56
Au, opt	0.099	0.004	0.092	0.106	0.088	0.109	3.54%	7.08%	10.62%	0.094	0.104
Infrared Combustion	Infrared Combustion										
C, wt.%	1.86	0.051	1.75	1.96	1.70	2.01	2.76%	5.52%	8.29%	1.76	1.95
C-(Carbonate), wt.%	1.71	0.040	1.63	1.79	1.59	1.83	2.35%	4.70%	7.05%	1.62	1.79
S, wt.%	0.619	0.043	0.532	0.706	0.489	0.749	7.01%	14.03%	21.04%	0.588	0.650
Aqua Regia Digestio	on										
Ag, ppm	0.093	0.011	0.072	0.115	0.061	0.126	11.67%	23.34%	35.02%	0.089	0.098
Al, wt.%	0.923	0.049	0.825	1.020	0.776	1.069	5.30%	10.60%	15.90%	0.876	0.969
As, ppm	467	23	421	512	399	534	4.86%	9.71%	14.57%	443	490
B, ppm	< 10	IND	IND	IND	IND	IND	IND	IND	IND	IND	IND

SI unit equivalents: ppm (parts per million)  $\equiv$  mg/kg  $\equiv$   $\mu$ g/g  $\equiv$  0.0001 wt.%  $\equiv$  1000 ppb (parts per billion).

1opt (troy ounce per short ton)  $\equiv$  34.2857ppm.

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.



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#### Table 1 continued.

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Constituent	Certified	P	Absolute \$	Standard	Deviation	าร	Relative	Standard D	eviations	5% w	indow
Constituent	Value	1SD	2SD Low	2SD High	3SD Low	3SD High	1RSD	2RSD	3RSD	Low	High
Aqua Regia Digestic	on continu	ued									
Be, ppm	0.33	0.05	0.24	0.43	0.20	0.47	13.79%	27.57%	41.36%	0.32	0.35
Bi, ppm	0.20	0.016	0.17	0.24	0.16	0.25	7.59%	15.18%	22.77%	0.19	0.21
Ca, wt.%	4.50	0.186	4.13	4.87	3.94	5.06	4.12%	8.25%	12.37%	4.27	4.72
Cd, ppm	0.36	0.029	0.30	0.42	0.27	0.45	8.06%	16.12%	24.18%	0.34	0.38
Ce, ppm	28.2	1.64	25.0	31.5	23.3	33.2	5.80%	11.60%	17.40%	26.8	29.7
Co, ppm	5.99	0.372	5.24	6.73	4.87	7.10	6.22%	12.44%	18.66%	5.69	6.29
Cr, ppm	27.8	1.53	24.8	30.9	23.3	32.4	5.48%	10.96%	16.45%	26.4	29.2
Cs, ppm	2.05	0.125	1.81	2.30	1.68	2.43	6.07%	12.13%	18.20%	1.95	2.16
Cu, ppm	25.5	1.74	22.1	29.0	20.3	30.8	6.82%	13.64%	20.45%	24.3	26.8
Dy, ppm	1.51	0.048	1.41	1.60	1.36	1.65	3.20%	6.39%	9.59%	1.43	1.58
Er, ppm	0.66	0.049	0.56	0.76	0.51	0.81	7.43%	14.86%	22.29%	0.63	0.69
Eu, ppm	0.55	0.07	0.40	0.69	0.33	0.76	13.03%	26.06%	39.09%	0.52	0.57
Fe, wt.%	1.80	0.082	1.64	1.97	1.56	2.05	4.53%	9.06%	13.59%	1.71	1.90
Ga, ppm	2.92	0.223	2.47	3.37	2.25	3.59	7.65%	15.30%	22.95%	2.78	3.07
Gd, ppm	2.18	0.169	1.84	2.52	1.68	2.69	7.75%	15.49%	23.24%	2.07	2.29
Ge, ppm	0.054	0.006	0.042	0.066	0.036	0.073	11.39%	22.78%	34.17%	0.051	0.057
Hf, ppm	0.47	0.06	0.35	0.58	0.30	0.64	12.19%	24.38%	36.57%	0.44	0.49
Hg, ppm	4.45	0.433	3.59	5.32	3.15	5.75	9.73%	19.45%	29.18%	4.23	4.68
Ho, ppm	0.26	0.015	0.23	0.29	0.21	0.30	5.74%	11.48%	17.22%	0.24	0.27
In, ppm	0.022	0.002	0.018	0.026	0.017	0.027	8.08%	16.15%	24.23%	0.021	0.023
K, wt.%	0.373	0.023	0.327	0.419	0.304	0.443	6.21%	12.42%	18.63%	0.354	0.392
La, ppm	15.2	0.79	13.6	16.7	12.8	17.5	5.21%	10.43%	15.64%	14.4	15.9
Li, ppm	13.7	1.35	11.0	16.4	9.7	17.7	9.83%	19.67%	29.50%	13.0	14.4
Lu, ppm	0.079	0.005	0.069	0.089	0.064	0.094	6.37%	12.75%	19.12%	0.075	0.083
Mg, wt.%	1.27	0.057	1.16	1.39	1.10	1.44	4.51%	9.03%	13.54%	1.21	1.33
Mn, wt.%	0.027	0.001	0.024	0.030	0.022	0.031	5.44%	10.87%	16.31%	0.025	0.028
Mo, ppm	4.51	0.253	4.00	5.02	3.75	5.27	5.62%	11.24%	16.86%	4.28	4.73
Na, wt.%	0.017	0.005	0.006	0.028	0.001	0.034	31.57%	63.14%	94.72%	0.016	0.018
Nb, ppm	0.077	0.017	0.043	0.110	0.026	0.127	21.98%	43.96%	65.94%	0.073	0.080
Nd, ppm	13.8	0.74	12.3	15.3	11.6	16.0	5.32%	10.64%	15.96%	13.1	14.5
Ni, ppm	20.1	1.33	17.5	22.8	16.2	24.1	6.58%	13.16%	19.74%	19.1	21.1
P, wt.%	0.050	0.002	0.046	0.054	0.045	0.056	3.69%	7.37%	11.06%	0.048	0.053
Pb, ppm	7.25	0.366	6.52	7.98	6.15	8.35	5.05%	10.09%	15.14%	6.89	7.61
Pd, ppb	< 10	IND	IND	IND	IND	IND	IND	IND	IND	IND	IND
Pr, ppm	3.52	0.237	3.04	3.99	2.81	4.23	6.74%	13.48%	20.22%	3.34	3.70
Pt, ppb	< 5	IND	IND	IND	IND	IND	IND	IND	IND	IND	IND
Rb, ppm	26.8	2.06	22.7	30.9	20.6	33.0	7.69%	15.38%	23.06%	25.4	28.1
Re, ppm	0.003	0.001	0.002	0.004	0.001	0.004	19.01%	38.01%	57.02%	0.003	0.003
S, wt.%	0.641	0.038	0.566	0.716	0.528	0.753	5.85%	11.70%	17.56%	0.609	0.673
Sb, ppm	30.4	3.9	22.5	38.3	18.6	42.3	12.97%	25.94%	38.91%	28.9	32.0
	2.60	0.148	2.31	2.90		3.05	5.69%	11.37%	17.06%	2.47	2.73
Sc, ppm					2.16						
Se, ppm	0.70	0.17	0.36	1.04	0.19	1.21	24.49%	48.98%	73.46%	0.66	0.73

SI unit equivalents: ppm (parts per million)  $\equiv$  mg/kg  $\equiv$   $\mu$ g/g  $\equiv$  0.0001 wt.%  $\equiv$  1000 ppb (parts per billion).

Note 1: intervals may appear asymmetric due to rounding.

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## Table 1 continued.

0	Certified	P	Absolute \$	Standard	Deviation	าร	Relative	Standard D	eviations	5% w	indow
Constituent	Value	1SD	2SD Low	2SD High	3SD Low	3SD High	1RSD	2RSD	3RSD	Low	High
Aqua Regia Digestion continued											
Sm, ppm	2.57	0.167	2.24	2.90	2.07	3.07	6.49%	12.97%	19.46%	2.44	2.70
Sn, ppm	0.93	0.10	0.73	1.13	0.64	1.23	10.57%	21.15%	31.72%	0.88	0.98
Sr, ppm	56	2.8	50	61	47	64	5.10%	10.20%	15.29%	53	58
Ta, ppm	< 0.01	IND	IND	IND	IND	IND	IND	IND	IND	IND	IND
Tb, ppm	0.29	0.029	0.23	0.35	0.21	0.38	9.97%	19.94%	29.91%	0.28	0.31
Te, ppm	0.13	0.03	0.08	0.18	0.05	0.21	19.92%	39.84%	59.75%	0.12	0.14
Th, ppm	5.16	0.377	4.41	5.92	4.03	6.29	7.29%	14.59%	21.88%	4.90	5.42
Ti, wt.%	0.035	0.002	0.031	0.039	0.028	0.041	6.17%	12.34%	18.52%	0.033	0.037
TI, ppm	6.88	0.514	5.85	7.91	5.34	8.42	7.47%	14.94%	22.40%	6.53	7.22
Tm, ppm	0.090	0.008	0.074	0.105	0.067	0.113	8.63%	17.26%	25.90%	0.085	0.094
U, ppm	1.52	0.120	1.28	1.76	1.16	1.88	7.87%	15.74%	23.61%	1.44	1.60
V, ppm	31.5	2.92	25.6	37.3	22.7	40.2	9.27%	18.54%	27.80%	29.9	33.0
W, ppm	2.10	0.33	1.43	2.77	1.10	3.10	15.86%	31.71%	47.57%	2.00	2.21
Y, ppm	6.86	0.405	6.05	7.67	5.65	8.08	5.91%	11.81%	17.72%	6.52	7.21
Yb, ppm	0.56	0.045	0.47	0.65	0.43	0.70	8.01%	16.03%	24.04%	0.54	0.59
Zn, ppm	53	3.4	46	60	43	63	6.51%	13.02%	19.53%	50	55
Zr, ppm	15.2	1.52	12.2	18.2	10.6	19.7	9.99%	19.99%	29.98%	14.4	16.0

SI unit equivalents: ppm (parts per million)  $\equiv$  mg/kg  $\equiv$   $\mu$ g/g  $\equiv$  0.0001 wt.%  $\equiv$  1000 ppb (parts per billion).

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.

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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 'Intended Use' should be read carefully.

OREAS 277 is one of a suite of three Carlin ore CRMs available: OREAS 277: 3.39ppm (0.1opt); OREAS 278: 4.99ppm (0.15opt) and OREAS 279: 6.54ppm (0.19opt). OREAS 277 contains 67 certified values including C and S speciation and full ICP-OES and MS suites by aqua regia digestion.

Furthermore, 105 indicative values are provided including Au by cyanide leach, Au by aqua regia digestion and major and trace element composition (see Table 2 below). Au by the aforementioned leach methods was uncertifiable due to a lack of interlaboratory consensus. The refractory nature of the gold ore and presence of carbon caused a wide variation in results. However, it is important to note that a high level of repeatability (precision) was displayed within each laboratory's data set for both cyanide leach and aqua regia digestion.

Tabulated results of all elements together with uncorrected means, medians, standard deviations, relative standard deviations and per cent deviation of lab means from the corrected mean of means (PDM³) are presented in the detailed certification data for this CRM (OREAS 277-DataPack.3.0.201218\_140109.xlsx).

# **SOURCE MATERIAL**

OREAS 277 was prepared from a blend of high grade gold-bearing ore and barren sediments (shale, quartz and limestone). The ore was sourced from the Leeville Mine, located near the western crest of the Tuscarora Mountains, about 20 miles northwest of Carlin in Nevada, USA. Leeville is an underground high-grade refractory gold deposit located on the Carlin Trend. Gold mineralisation is hosted within decalcified and weakly to moderately silicified rocks composed of 60 to 70% quartz, 10 to 30% dolomite, 5 to 12% kaolinite, 2 to 4% illite and 2 to 4% pyrite (Jackson et al, 1997).

## **PERFORMANCE GATES**

Table 1 above shows intervals calculated for two and three standard deviations. As a guide these intervals may be regarded as warning or rejection for multiple 2SD outliers, or rejection for individual 3SD outliers in QC monitoring, although their precise application should be at the discretion of the QC manager concerned (also see 'Intended Use' section below). Westgard Rules extend the basics of single-rule QC monitoring using multi-rules

(for more information visit www.westgard.com/mltirule.htm). A second method utilises a 5% window calculated directly from the certified value.

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

i.e. Certified Value ± 10% ± 2DL (adapted from Govett, 1983).

Table 2. Indicative Values for OREAS 277.

Pb Fire Assay			T able 2	. Illuicative va	1003 101	OILLAG ETT	•		
Ag         ppm         0.267         Pd         ppb         0.453         Pt         ppb         0.402           Cyanide Leach           Au         ppm         0.818         Image: Cyanide of the pipe of	Constituent	Unit	Value	Constituent	Unit	Value	Constituent	Unit	Value
Au	Pb Fire Assay					•			
Au	Ag	ppm	0.267	Pd	ppb	0.453	Pt	ppb	0.402
Infrared Combustion	Cyanide Leach								
C-(non.Carbonate)         wt.%         0.157         C-(Organic)         wt.%         0.070         S-(Sulphide)         wt.%         0.416           C-(Graphite)         wt.%         0.097         S-(Sulphate)         wt.%         0.187         wt.%         0.416           Aqua Regia Digestion           Au         ppm         1.74         Ir         ppb         0.930         Si         wt.%         0.051           Ba         ppm         584         Rh         ppb         3.38         wt.%         0.051           X-ray Photon Assay           Au         ppm         3.41         Ir         ppb         3.38         wt.%         0.051           Borate Fusion XRF           Al2O3         wt.%         5.39         MgO         wt.%         2.45         S         wt.%         0.629           CaO         wt.%         6.61         MnO         wt.%         0.030         SiO2         wt.%         71.89           Fe <sub>2</sub> O3         wt.%         2.72         Na <sub>2</sub> O         wt.%         0.430         TiO2         wt.%         0.360           K <sub>2</sub> O         wt.%         0.113         wt.% <td< td=""><td>Au</td><td>ppm</td><td>0.818</td><td></td><td></td><td></td><td></td><td></td><td></td></td<>	Au	ppm	0.818						
C-(Graphite)         wt.%         0.097         S-(Sulphate)         wt.%         0.187         Aqua Regia Digestion           Au         ppm         1.74         Ir         ppb         0.930         Si         wt.%         0.051           Ba         ppm         584         Rh         ppb         3.38         Si         wt.%         0.051           X-ray Photon Assay           Au         ppm         3.41         Si         Si         wt.%         0.629           Borate Fusion XRF           Al <sub>2</sub> O <sub>3</sub> wt.%         5.39         MgO         wt.%         2.45         Si         wt.%         0.629           CaO         wt.%         6.61         MnO         wt.%         0.030         SiO <sub>2</sub> wt.%         7.199           Fe <sub>2</sub> O <sub>3</sub> wt.%         2.72         Na <sub>2</sub> O         wt.%         0.430         TiO <sub>2</sub> wt.%         0.360           K <sub>2</sub> O         wt.%         7.19         Wt.%         0.113         Wt.%         0.360           Thermogravimetry           Lol1000         wt.%         7.19         Wt.%         0.133         Sm         ppm         3.29	Infrared Combustic	on							
Aqua Regia Digestion           Au         ppm         1.74         Ir         ppb         0.930         Si         wt.%         0.051           Ba         ppm         584         Rh         ppb         3.38  .	C-(non.Carbonate)	wt.%	0.157	C-(Organic)	wt.%	0.070	S-(Sulphide)	wt.%	0.416
Au         ppm         1.74         Ir         ppb         0.930         Si         wt.%         0.051           Ba         ppm         584         Rh         ppb         3.38         —	C-(Graphite)	wt.%	0.097	S-(Sulphate)	wt.%	0.187			
Name	Aqua Regia Digest	ion							
Au	Au	ppm	1.74	lr	ppb	0.930	Si	wt.%	0.051
Au         ppm         3.41         Borate Fusion XRF           Al₂O₃         wt.%         5.39         MgO         wt.%         2.45         S         wt.%         0.629           CaO         wt.%         6.61         MnO         wt.%         0.030         SiO₂         wt.%         71.89           Fe₂O₃         wt.%         2.72         Na₂O         wt.%         0.430         TiO₂         wt.%         0.360           K₂O         wt.%         1.10         P₂O₅         wt.%         0.113         TiO₂         wt.%         0.360           Thermogravimetry           LOI¹000         wt.%         7.19         Image: Single singl	Ва	ppm	584	Rh	ppb	3.38			
May	X-ray Photon Assa	y							
Al <sub>2</sub> O <sub>3</sub> wt.%         5.39         MgO         wt.%         2.45         S         wt.%         0.629           CaO         wt.%         6.61         MnO         wt.%         0.030         SiO <sub>2</sub> wt.%         71.89           Fe <sub>2</sub> O <sub>3</sub> wt.%         2.72         Na <sub>2</sub> O         wt.%         0.430         TiO <sub>2</sub> wt.%         0.360           K <sub>2</sub> O         wt.%         1.10         P <sub>2</sub> O <sub>5</sub> wt.%         0.113         TiO <sub>2</sub> wt.%         0.360           Thermogravimetry           LOI <sup>1000</sup> wt.%         7.19         Image: Colspan="6">Trick         Wt.%         0.113         Image: Colspan="6">Trick         Vt.%         0.360           Thermogravimetry         Trick         Trick <td>Au</td> <td>ppm</td> <td>3.41</td> <td></td> <td></td> <td></td> <td></td> <td></td> <td></td>	Au	ppm	3.41						
CaO         wt.%         6.61         MnO         wt.%         0.030         SiO2         wt.%         71.89           Fe2O3         wt.%         2.72         Na2O         wt.%         0.430         TiO2         wt.%         0.360           K2O         wt.%         1.10         P2O5         wt.%         0.113         TiO2         wt.%         0.360           Thermogravimetry           LOI <sup>1000</sup> wt.%         7.19         Image: Calcal Science Sc	Borate Fusion XRF	•							
Fe <sub>2</sub> O <sub>3</sub> wt.%         2.72         Na <sub>2</sub> O         wt.%         0.430         TiO <sub>2</sub> wt.%         0.360           K <sub>2</sub> O         wt.%         1.10         P <sub>2</sub> O <sub>5</sub> wt.%         0.113         TiO <sub>2</sub> wt.%         0.360           Thermogravimetry           LOI <sup>1000</sup> wt.%         7.19         Image: Colspan="6">Trian in the colspan="6">Tr	$Al_2O_3$	wt.%	5.39	MgO	wt.%	2.45	S	wt.%	0.629
K2O         wt.%         1.10         P2Os         wt.%         0.113         Image: colspan="8">Image: colspan="8">Image: colspan="8">Image: colspan="8">Image: colspan="8">Wt.%         1.10         P2Os         wt.%         0.113         Image: colspan="8">Image: colspan="8">I	CaO	wt.%	6.61	MnO	wt.%	0.030	SiO <sub>2</sub>	wt.%	71.89
Thermogravimetry   LOI <sup>1000</sup>   wt.%   7.19	Fe <sub>2</sub> O <sub>3</sub>	wt.%	2.72	Na₂O	wt.%	0.430	TiO <sub>2</sub>	wt.%	0.360
LOI <sup>1000</sup> wt.%         7.19         Image: Control of the control of	K <sub>2</sub> O	wt.%	1.10	P <sub>2</sub> O <sub>5</sub>	wt.%	0.113			
Laser Ablation ICP-MS           Ag         ppm         0.100         Hf         ppm         3.33         Sm         ppm         3.29           As         ppm         389         Ho         ppm         0.50         Sn         ppm         2.30           Ba         ppm         995         In         ppm         < 0.05	Thermogravimetry								
Ag         ppm         0.100         Hf         ppm         3.33         Sm         ppm         3.29           As         ppm         389         Ho         ppm         0.50         Sn         ppm         2.30           Ba         ppm         995         In         ppm         < 0.05	LOI <sup>1000</sup>	wt.%	7.19						
As ppm 389 Ho ppm 0.50 Sn ppm 2.30  Ba ppm 995 In ppm <0.05 Sr ppm 66  Be ppm 1.10 La ppm 19.2 Ta ppm 0.49  Bi ppm 0.20 Lu ppm 0.20 Tb ppm 0.45  Cd ppm 0.40 Mn wt.% 0.029 Te ppm <0.53  Ce ppm 35.1 Mo ppm 4.30 Th ppm 6.53  Co ppm 6.40 Nb ppm 6.21 Ti wt.% 0.220  Cr ppm 48.5 Nd ppm 16.6 Tl ppm 9.60  Cs ppm 3.53 Ni ppm 23.0 Tm ppm 0.21	Laser Ablation ICP	-MS							
Ba         ppm         995         In         ppm         < 0.05         Sr         ppm         66           Be         ppm         1.10         La         ppm         19.2         Ta         ppm         0.49           Bi         ppm         0.20         Lu         ppm         0.20         Tb         ppm         0.45           Cd         ppm         0.40         Mn         wt.%         0.029         Te         ppm         < 0.2	Ag	ppm	0.100	Hf	ppm	3.33	Sm	ppm	3.29
Be         ppm         1.10         La         ppm         19.2         Ta         ppm         0.49           Bi         ppm         0.20         Lu         ppm         0.20         Tb         ppm         0.45           Cd         ppm         0.40         Mn         wt.%         0.029         Te         ppm         < 0.2	As	ppm	389	Но	ppm	0.50	Sn	ppm	2.30
Bi         ppm         0.20         Lu         ppm         0.20         Tb         ppm         0.45           Cd         ppm         0.40         Mn         wt.%         0.029         Te         ppm         < 0.2	Ва	ppm	995	In	ppm	< 0.05	Sr	ppm	66
Cd         ppm         0.40         Mn         wt.%         0.029         Te         ppm         < 0.2           Ce         ppm         35.1         Mo         ppm         4.30         Th         ppm         6.53           Co         ppm         6.40         Nb         ppm         6.21         Ti         wt.%         0.220           Cr         ppm         48.5         Nd         ppm         16.6         Tl         ppm         9.60           Cs         ppm         3.53         Ni         ppm         23.0         Tm         ppm         0.21	Be	ppm	1.10	La	ppm	19.2	Та	ppm	0.49
Ce         ppm         35.1         Mo         ppm         4.30         Th         ppm         6.53           Co         ppm         6.40         Nb         ppm         6.21         Ti         wt.%         0.220           Cr         ppm         48.5         Nd         ppm         16.6         Tl         ppm         9.60           Cs         ppm         3.53         Ni         ppm         23.0         Tm         ppm         0.21	Bi	ppm	0.20	Lu	ppm	0.20	Tb	ppm	0.45
Co         ppm         6.40         Nb         ppm         6.21         Ti         wt.%         0.220           Cr         ppm         48.5         Nd         ppm         16.6         Tl         ppm         9.60           Cs         ppm         3.53         Ni         ppm         23.0         Tm         ppm         0.21	Cd	ppm	0.40	Mn	wt.%	0.029	Te	ppm	< 0.2
Cr         ppm         48.5         Nd         ppm         16.6         TI         ppm         9.60           Cs         ppm         3.53         Ni         ppm         23.0         Tm         ppm         0.21	Ce	ppm	35.1	Мо	ppm	4.30	Th	ppm	6.53
Cs ppm 3.53 Ni ppm 23.0 Tm ppm 0.21	Co	ppm	6.40	Nb	ppm	6.21	Ti	wt.%	0.220
	Cr	ppm	48.5	Nd	ppm	16.6	TI	ppm	9.60
Cu ppm 25.0 Pb ppm 8.00 U ppm 2.65	Cs	ppm	3.53	Ni	ppm	23.0	Tm	ppm	0.21
	Cu	ppm	25.0	Pb	ppm	8.00	U	ppm	2.65

SI unit equivalents: ppm (parts per million)  $\equiv$  mg/kg  $\equiv$   $\mu$ g/g  $\equiv$  0.0001 wt.%  $\equiv$  1000 ppb (parts per billion). Note: the number of significant figures reported is not a reflection of the level of certainty of stated values. They are instead an artefact of ORE's in-house CRM-specific LIMS.

#### Table 2 continued.

Constituent	Unit	Value	Constituent	Unit	Value	Constituent	Unit	Value	
Laser Ablation ICP	Laser Ablation ICP-MS continued								
Dy	ppm	2.54	Pr	ppm	4.45	V	ppm	73	
Er	ppm	1.41	Rb	ppm	52	W	ppm	13.5	
Eu	ppm	0.64	Re	ppm	< 0.01	Υ	ppm	14.4	
Ga	ppm	7.20	Sb	ppm	36.2	Yb	ppm	1.32	
Gd	ppm	2.94	Sc	ppm	6.05	Zn	ppm	65	
Ge	ppm	0.93	Se	ppm	< 5	Zr	ppm	120	
4-Acid Digestion	4-Acid Digestion								
Al	wt.%	2.60	Cu	ppm	46.5	Pb	ppm	10.9	
As	ppm	465	Fe	wt.%	1.75	Re	ppm	13.2	
Ва	ppm	901	K	wt.%	0.860	Sb	ppm	40.7	
Be	ppm	1.00	Mg	wt.%	1.38	Se	ppm	9.20	
Bi	ppm	< 13	Mn	wt.%	0.028	Sn	ppm	2.42	
Ca	wt.%	4.76	Мо	ppm	12.6	Te	ppm	< 6	
Cd	ppm	2.00	Na	wt.%	0.305	Ti	wt.%	0.182	
Со	ppm	7.50	Ni	ppm	18.0	TI	ppm	< 8	
Cr	ppm	25.2	Р	wt.%	0.046	Zn	ppm	52	

SI unit equivalents: ppm (parts per million)  $\equiv$  mg/kg  $\equiv$   $\mu$ g/g  $\equiv$  0.0001 wt.%  $\equiv$  1000 ppb (parts per billion). Note: the number of significant figures reported is not a reflection of the level of certainty of stated values. They are instead an artefact of ORE's in-house CRM-specific LIMS.

## COMMINUTION AND HOMOGENISATION PROCEDURES

The material constituting OREAS 277 was prepared in the following manner:

- Drying to constant mass at 105°C;
- Crushing and milling of the barren sediments to 98% minus 75 microns;
- Crushing and milling of the ore material to 100% minus 30 microns;
- Blending in appropriate proportions to achieve the desired grade;
- Packaging in 60g units sealed in laminated foil pouches and 500g units in plastic jars.

## PHYSICAL PROPERTIES

OREAS 277 was tested at ORE Research & Exploration Pty Ltd's onsite facility for various physical properties. Table 3 presents these findings that should be used for informational purposes only.

Table 3. Physical properties of OREAS 277.

Bulk Density (g/L)	Moisture%	Munsell Notation <sup>‡</sup>	Munsell Color <sup>‡</sup>
735	0.70	N7	Light Gray

<sup>&</sup>lt;sup>‡</sup>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

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

- Gold by fire assay (15-50g charge weight) with AAS (31 laboratories), ICP-OES (8 laboratories) or ICP-MS (2 laboratories) finish;
- Gold by cyanide leach A variety of cyanide leach methods were undertaken by the participating laboratories including the use of LeachWELL tablets, alkaline added sodium cyanide solution as well as sodium cyanide liquor with LeachWELL powder. The sample weights included: 5g (1 laboratory by AAS finish), 10g (1 laboratory by ICP-OES finish), 30g (8 laboratories by AAS, 1 laboratory by ICP-OES and 1 laboratory ICP-MS finish), 50g (1 laboratory by AAS and 2 laboratory by ICP-MS finish) and 200g (5 laboratories by AAS, 2 laboratories by ICP-OES and 2 laboratories by ICP-MS finish);
- Aqua regia digestion for full ICP-OES and ICP-MS elemental suites (up to 34 laboratories depending on the element);
- Total Carbon (29 laboratories) and Total Sulphur (30 laboratories) by infrared combustion furnace;
- Carbonate Carbon and Non-Carbonate Carbon by a range of methodologies including HCl or HClO<sub>4</sub> leach followed by infrared combustion furnace or coulometry, or via difference: Total C minus Non-Carbonate C (for Carbonate C determination) or, Total C minus Carbonate C (for Non-Carbonate C determination);
- Graphitic Carbon by HCl digestion to remove carbonates followed by roasting to remove organic carbon, the residue is then determined by infrared combustion furnace;
- Sulphate Sulphur by HCl leach followed by gravimetric (6 laboratories) or ICP-OES (1 laboratory) finish or roasting (at 500-550°C) followed by infrared combustion furnace (7 laboratories);
- Sulphide Sulphur by difference (Total S minus Sulphate Sulphur).

Not all forms of Carbon and Sulphur were able to be certified. Other than the dominant forms (Total Carbon, Carbonate Carbon and Total Sulphur), the others forms (Non-Carbonate Carbon, Graphitic Carbon, Organic Carbon, Sulphate Sulphur and Sulphide Sulphur) displayed insufficient interlaboratory consensus and are shown in Table 2 as indicative values only.

To confirm homogeneity, gold by instrumental neutron activation analysis (INAA) was undertaken on 20 x 85mg subsamples by the Australian Nuclear Science and Technology Organisation (ANSTO) located in Lucas Heights, NSW, Australia (see Table 5 in the 'Homogeneity Evaluation' section below).

For the round robin program twenty 3.0kg test units were taken at predetermined intervals during the bagging stage, immediately following homogenisation and are considered representative of the entire prepared batch. Six 100-350g pulp samples were submitted to each laboratory for analysis (the weight provided depended on whether the laboratory was anticipated to undertake assays by gold cyanide leach). The samples received by each laboratory were obtained by taking two samples from each of three separate 3.0kg 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.

Table 1 provides performance gate intervals for the 67 certified values based on their pooled 1SD's. Table 2 shows 105 indicative values including gold by Chrysos Corporation's Photon Assay technique and major and trace element characterisation by Bureau Veritas Perth using the following methodologies:

- Major oxides by lithium borate fusion with X-ray fluorescence;
- LOI at 1000°C by thermogravimetric analyser;
- Trace element characterisation by laser ablation with ICP-MS finish.

Table 2 also presents twenty additional indicative values (Al to Zn) by 4-acid digestion with ICP-OES/MS finish performed by one of the participating laboratories. Table 3 provides some indicative physical properties and Table 4 presents 95% confidence and tolerance limits. Gold homogeneity (via INAA) is shown in Table 5 and is also demonstrated by a nested ANOVA program using fire assay (see 'nested ANOVA' section).

Results are also presented in scatter plots for gold by fire assay, total carbon by infrared combustion furnace and total sulphur by infrared combustion furnace (Figures 1 to 3, respectively) together with ±3SD (magenta) and ±5% (yellow) control lines and certified value (green line). Accepted individual results are coloured blue and individual and dataset outliers are identified in red and violet, respectively.

## STATISTICAL ANALYSIS

**Standard Deviation** intervals (see Table 1) provide an indication of a level of performance that might reasonably be expected from a laboratory being monitored by this CRM in a QA/QC program. They take into account errors attributable to measurement uncertainty and CRM variability. For an effective CRM the contribution of the latter should be negligible in comparison to measurement errors. The Standard Deviation values include all sources of measurement uncertainty: between-lab variance, within-run variance (precision errors) and CRM variability.

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

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

Certified Values, Standard Deviations, Confidence Limits and Tolerance Limits (Table 4) have been determined for each analyte following removal of individual, laboratory dataset (batch) and 3SD outliers (single iteration).

For individual outliers within a laboratory batch the z-score test is used in combination with a second method that determines the per cent deviation of the individual value from the

batch median. Outliers in general are selected on the basis of z-scores > 2.5 and with per cent deviations (i) > 3 and (ii) more than three times the average absolute per cent deviation for the batch. In certain instances statistician's prerogative has been employed in discriminating outliers.

Each laboratory data set mean is tested for outlying status based on z-score discrimination and rejected if > 2.5. After individual and laboratory data set (batch) outliers have been eliminated a non-iterative 3 standard deviation filter is applied, with those values lying outside this window also relegated to outlying status.

**Certified Values** are the means of accepted laboratory means after outlier filtering. The INAA data (see Table 5) is omitted from determination of the certified value for Au and is used solely for the calculation of Tolerance Limits and homogeneity evaluation of OREAS 277 (see 'Homogeneity Evaluation' section below).

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

**Indicative (uncertified) values** (Table 2) are provided for the major and trace elements determined by borate fusion XRF ( $Al_2O_3$  to  $TiO_2$ ), laser ablation with ICP-MS (Ag to Zr) and LOI at  $1000^{\circ}$ C and are the means of duplicate assays from Bureau Veritas, Perth. Au by Chrysos Corporation's Photon Assay technique is also shown. Additional indicative values by other analytical methods are present where the number of laboratories reporting a particular analyte is insufficient (< 5) to support certification or where inter-laboratory consensus is poor.

Table 4. 95% Confidence & Tolerance Limits for OREAS 277.

Ormstitusest	Certified	95% Confid	ence Limits	95% Tolerance Limits				
Constituent	Value	Low	High	Low	High			
Pb Fire Assay								
Au, Gold (ppm)	3.39	3.36	3.43	3.38*	3.40*			
Au, Gold (opt)	0.099	0.098	0.100	0.099*	0.099*			
Infrared Combustion (wt.%)								
C, Carbon	1.86	1.84	1.88	1.82	1.89			
C-(Carbonate)	1.71	1.69	1.73	1.69	1.73			
S, Sulphur	0.619	0.603	0.635	0.606	0.632			
Aqua Regia Digestion								
Ag, Silver (ppm)	0.093	0.089	0.098	0.082	0.105			
Al, Aluminium (wt.%)	0.923	0.904	0.941	0.899	0.946			
As, Arsenic (ppm)	467	459	475	457	476			
B, Boron (ppm)	< 10	IND	IND	IND	IND			
Be, Beryllium (ppm)	0.33	0.31	0.36	0.32	0.35			
Bi, Bismuth (ppm)	0.20	0.20	0.21	0.19	0.22			

SI unit equivalents: ppm (parts per million)  $\equiv$  mg/kg  $\equiv$   $\mu$ g/g  $\equiv$  0.0001 wt.%  $\equiv$  1000 ppb (parts per billion).

Note: intervals may appear asymmetric due to rounding.



<sup>\*</sup>Gold Tolerance Limits for typical 30g fire assay are determined from 20 x 85mg INAA results and the Sampling Constant (Ingamells & Switzer, 1973).

# Table 4 continued.

	Certified	4 continued.	lence Limits	95% Toler	ance Limits
Constituent	Value	Low	High	Low	High
Aqua Regia Digestion continued		2011	ı ııgıı	2011	i iigii
Ca, Calcium (wt.%)	4.50	4.43	4.57	4.43	4.57
Cd, Cadmium (ppm)	0.36	0.35	0.37	0.33	0.38
Ce, Cerium (ppm)	28.2	27.5	29.0	27.5	29.0
Co, Cobalt (ppm)	5.99	5.83	6.14	5.78	6.19
Cr, Chromium (ppm)	27.8	27.3	28.4	26.8	28.8
Cs, Caesium (ppm)	2.05	2.00	2.11	1.99	2.12
Cu, Copper (ppm)	25.5	24.9	26.2	24.3	26.8
Dy, Dysprosium (ppm)	1.51	1.47	1.54	1.43	1.58
Er, Erbium (ppm)	0.66	0.61	0.71	0.62	0.70
Eu, Europium (ppm)	0.55	0.61	0.62	0.62	0.70
	1.80		1.83		
Fe, Iron (wt.%)		1.78		1.77	1.84
Ga, Gallium (ppm)	2.92	2.82	3.02	2.80	3.04
Gd, Gadolinium (ppm)	2.18	2.03	2.34	2.08	2.28
Ge, Germanium (ppm)	0.054	0.048	0.060	IND	IND
Hf, Hafnium (ppm)	0.47	0.44	0.49	0.44	0.49
Hg, Mercury (ppm)	4.45	4.25	4.66	4.30	4.61
Ho, Holmium (ppm)	0.26	0.24	0.27	0.24	0.27
In, Indium (ppm)	0.022	0.021	0.023	0.019	0.025
La, Lanthanum (ppm)	15.2	14.8	15.5	14.8	15.5
Li, Lithium (ppm)	13.7	13.1	14.3	13.1	14.2
Lu, Lutetium (ppm)	0.079	0.075	0.082	IND	IND
Mg, Magnesium (wt.%)	1.27	1.25	1.29	1.25	1.30
Mn, Manganese (wt.%)	0.027	0.026	0.027	0.026	0.027
Mo, Molybdenum (ppm)	4.51	4.41	4.61	4.34	4.68
Na, Sodium (wt.%)	0.017	0.015	0.019	0.017	0.018
Nb, Niobium (ppm)	0.077	0.066	0.088	IND	IND
Nd, Neodymium (ppm)	13.8	13.2	14.5	13.2	14.5
Ni, Nickel (ppm)	20.1	19.7	20.6	19.5	20.7
P, Phosphorus (wt.%)	0.050	0.049	0.051	0.049	0.051
Pb, Lead (ppm)	7.25	7.08	7.42	6.99	7.51
Pd, Palladium (ppb)	< 10	IND	IND	IND	IND
Pr, Praseodymium (ppm)	3.52	3.32	3.72	3.35	3.69
Pt, Platinum (ppb)	< 5	IND	IND	IND	IND
Rb, Rubidium (ppm)	26.8	25.8	27.7	26.1	27.4
Re, Rhenium (ppm)	0.003	0.003	0.003	IND	IND
S, Sulphur (wt.%)	0.641	0.626	0.655	0.628	0.654
Sb, Antimony (ppm)	30.4	29.0	31.9	29.1	31.7
Sc, Scandium (ppm)	2.60	2.55	2.66	2.48	2.72

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

Note: intervals may appear asymmetric due to rounding.



Table 4 continued.

O a matitus mat	Certified	95% Confid	dence Limits	95% Tolei	ance Limits
Constituent	Value	Low	High	Low	High
Aqua Regia Digestion continue	d		•		•
Se, Selenium (ppm)	0.70	0.60	0.79	IND	IND
Sm, Samarium (ppm)	2.57	2.40	2.74	2.49	2.65
Sn, Tin (ppm)	0.93	0.89	0.98	0.87	0.99
Sr, Strontium (ppm)	56	55	57	54	57
Ta, Tantalum (ppm)	< 0.01	IND	IND	IND	IND
Tb, Terbium (ppm)	0.29	0.27	0.31	0.28	0.31
Te, Tellurium (ppm)	0.13	0.12	0.14	0.10	0.16
Th, Thorium (ppm)	5.16	4.99	5.33	5.00	5.32
Ti, Titanium (wt.%)	0.035	0.034	0.036	0.034	0.036
Tl, Thallium (ppm)	6.88	6.66	7.09	6.64	7.11
Tm, Thulium (ppm)	0.090	0.082	0.098	IND	IND
U, Uranium (ppm)	1.52	1.47	1.57	1.48	1.56
V, Vanadium (ppm)	31.5	30.4	32.6	30.5	32.5
W, Tungsten (ppm)	2.10	1.95	2.25	2.03	2.17
Y, Yttrium (ppm)	6.86	6.69	7.04	6.68	7.04
Yb, Ytterbium (ppm)	0.56	0.51	0.61	0.53	0.60
Zn, Zinc (ppm)	53	52	54	51	54
Zr, Zirconium (ppm)	15.2	14.5	15.9	14.7	15.7
W, Tungsten (ppm)	2.10	1.95	2.25	2.03	2.17
Y, Yttrium (ppm)	6.86	6.69	7.04	6.68	7.04
Yb, Ytterbium (ppm)	0.56	0.51	0.61	0.53	0.60
Zn, Zinc (ppm)	53	52	54	51	54
Zr, Zirconium (ppm)	15.2	14.5	15.9	14.7	15.7

SI unit equivalents: ppm (parts per million)  $\equiv$  mg/kg  $\equiv$   $\mu$ g/g  $\equiv$  0.0001 wt.%  $\equiv$  1000 ppb (parts per billion).

Note: intervals may appear asymmetric due to rounding.

#### **Homogeneity Evaluation**

For analytes other than gold the tolerance limits (ISO 16269:2014) shown in Table 4 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 arsenic by aqua-regia digestion, where 99% of the time  $(1-\alpha=0.99)$  at least 95% of subsamples ( $\rho=0.95$ ) will have concentrations lying between 457 and 476 ppm. Put more precisely, this means that if the same number of subsamples were taken and analysed in the same manner repeatedly, 99% of the tolerance intervals so constructed would cover at least 95% of the total population, and 1% of the tolerance intervals would cover less than 95% of the total population (ISO Guide 35). **Please note that tolerance limits pertain to the homogeneity of the CRM only and should not be used as control limits for laboratory performance.** 

Table 5 below shows the gold INAA data determined on 20 x 85mg subsamples of OREAS 277. An equivalent scaled version of the results is also provided to demonstrate the level of repeatability that would be achieved if 30g fire assay determinations were undertaken

without the normal measurement error associated with this methodology. The homogeneity of gold has been determined by INAA using the reduced analytical subsample method which utilises the known relationship between standard deviation and analytical subsample weight (Ingamells and Switzer, 1973). In this approach the sample aliquot is substantially reduced to a point where most of the variability in replicate assays should be due to inhomogeneity of the reference material (i.e. sampling error) and measurement error becomes negligible. In this instance a subsample weight of 85 milligrams was employed and the 1RSD of 0.06% was calculated for a 30g fire assay sample (1.18% at 85mg weights) and confirms the high level of gold homogeneity in OREAS 277.

Table 5. Neutron Activation Analysis of Au (in ppm) on 20 x 85mg subsamples and showing the equivalent results scaled to a 30g sample mass typical of fire assay determination.

Replicate	Au	Au
No	85mg actual	30g equivalent*
1	3.448	3.456
2	3.510	3.460
3	3.484	3.458
4	3.399	3.454
5	3.467	3.457
6	3.488	3.459
7	3.435	3.456
8	3.419	3.455
9	3.485	3.458
10	3.472	3.458
11	3.461	3.457
12	3.493	3.459
13	3.371	3.452
14	3.430	3.456
15	3.458	3.457
16	3.520	3.460
17	3.398	3.454
18	3.488	3.459
19	3.495	3.459
20	3.419	3.455
Mean	3.457	3.457
Median	3.464	3.457
Std Dev.	0.041	0.002
Rel.Std.Dev.	1.184%	0.063%

\*Results calculated for a 30g equivalent sample mass using the formula: 
$$x^{30g \ Eq} = \frac{(x^{INAA} - \bar{X}) \times RSD@30g}{RSD@85mg} + \bar{X}$$
 where  $x^{30g \ Eq} =$  equivalent result calculated for a 30g sample mass  $(x^{INAA}) =$  raw INAA result at 85mg  $\bar{X} =$  mean of 85mg INAA results

The homogeneity of OREAS 277 has also been evaluated in a **nested ANOVA** of the round robin program. Each of the forty-two round robin laboratories received six samples per CRM and these samples were made up of paired samples from three different, non-adjacent sampling intervals. The purpose of the ANOVA evaluation is to test that no statistically significant difference exists in the variance between units to that of the

variance within units. This allows an assessment of homogeneity across the entire prepared batch of OREAS 277. The test was performed using the following parameters:

- Gold fire assay 246 samples (41 laboratories each providing analyses on 3 pairs of samples);
- Null Hypothesis, H<sub>0</sub>: Between-unit variance is no greater than within-unit variance (reject H<sub>0</sub> if *p*-value < 0.05);
- Alternative Hypothesis, H<sub>1</sub>: Between-unit variance is greater than within-unit variance.

P-values are a measure of probability where values less than 0.05 indicate a greater than 95% probability that the observed differences in within-unit and between-unit variances are real. The datasets were filtered for both individual and laboratory data set (batch) outliers prior to the calculation of the p-value. This process derived a p-value of 0.91 which is an insignificant result and the Null Hypothesis is therefore retained. Additionally, none of the other certified values showed significant p-values. Please note that only results for constituents present in concentrations well above the detection levels (i.e. >20 x Lower Limit of Detection) for the various methods undertaken were considered for the objective of evaluating homogeneity.

It is important to note that ANOVA is not an absolute measure of homogeneity. Rather, it establishes whether or not the analytes are distributed in a similar manner throughout the packaging run of OREAS 277 and whether the variance between two subsamples from the same unit is statistically distinguishable from the variance of two subsamples taken from any two separate units. A reference material therefore can possess poor absolute homogeneity yet still pass a relative homogeneity (ANOVA) test if the within-unit heterogeneity is large and similar across all units.

Based on the statistical analysis of the results of the inter-laboratory certification program it can be concluded that OREAS 277 is fit-for-purpose as a certified reference material (see 'Intended Use' below).

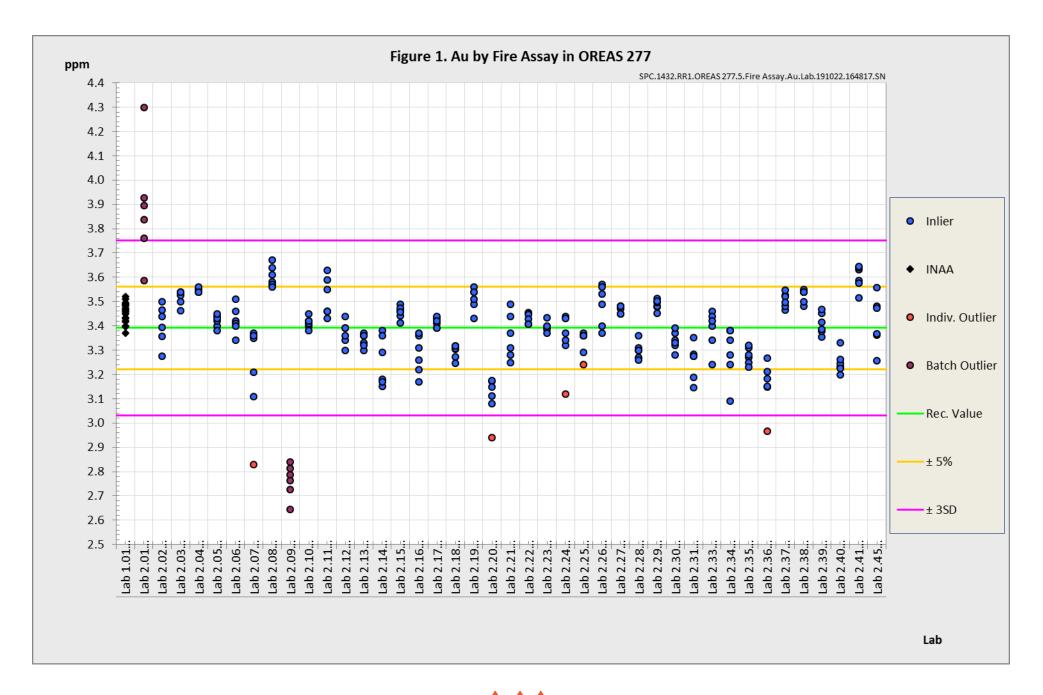
# PARTICIPATING LABORATORIES

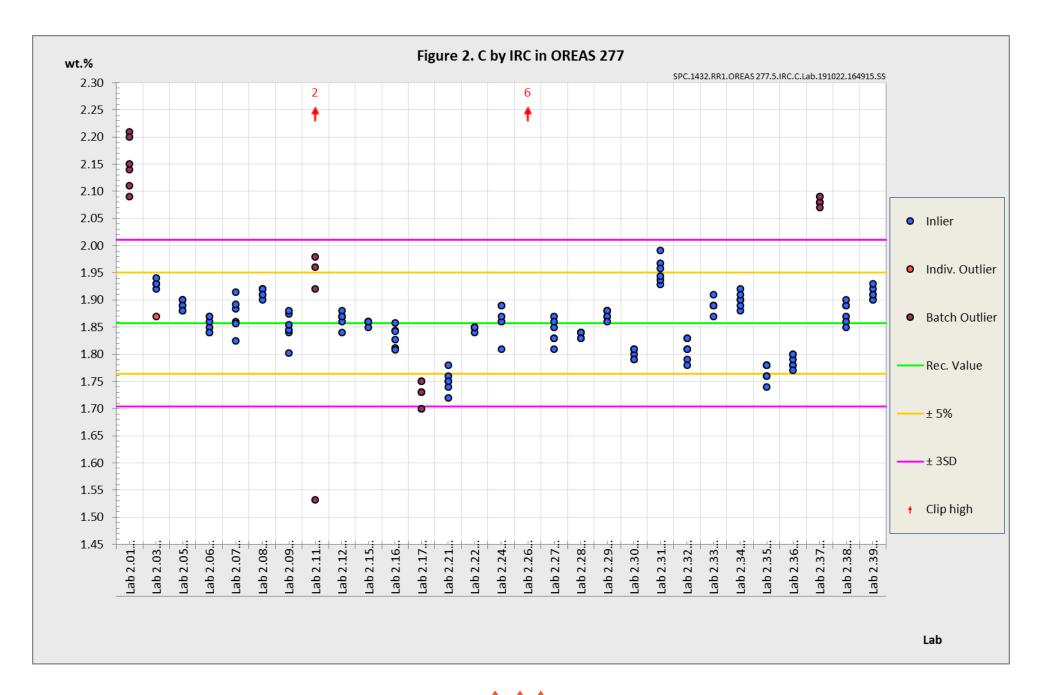
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- 2. AGAT Laboratories, Mississauga, Ontario, Canada
- 3. Alex Stewart International, Mendoza, Argentina
- 4. ALS, Lima, Peru
- 5. ALS, Loughrea, Galway, Ireland
- 6. ALS, Perth, WA, Australia
- 7. ALS, Reno, Nevada, USA
- 8. ALS, Vancouver, BC, Canada
- 9. American Assay Laboratories, Sparks, Nevada, USA
- 10. ANSTO, Lucas Heights, NSW, Australia
- 11. Bureau Veritas Commodities Canada Ltd, Vancouver, BC, Canada
- 12. Bureau Veritas Geoanalytical, Adelaide, SA, Australia
- 13. Bureau Veritas Geoanalytical, Perth, WA, Australia

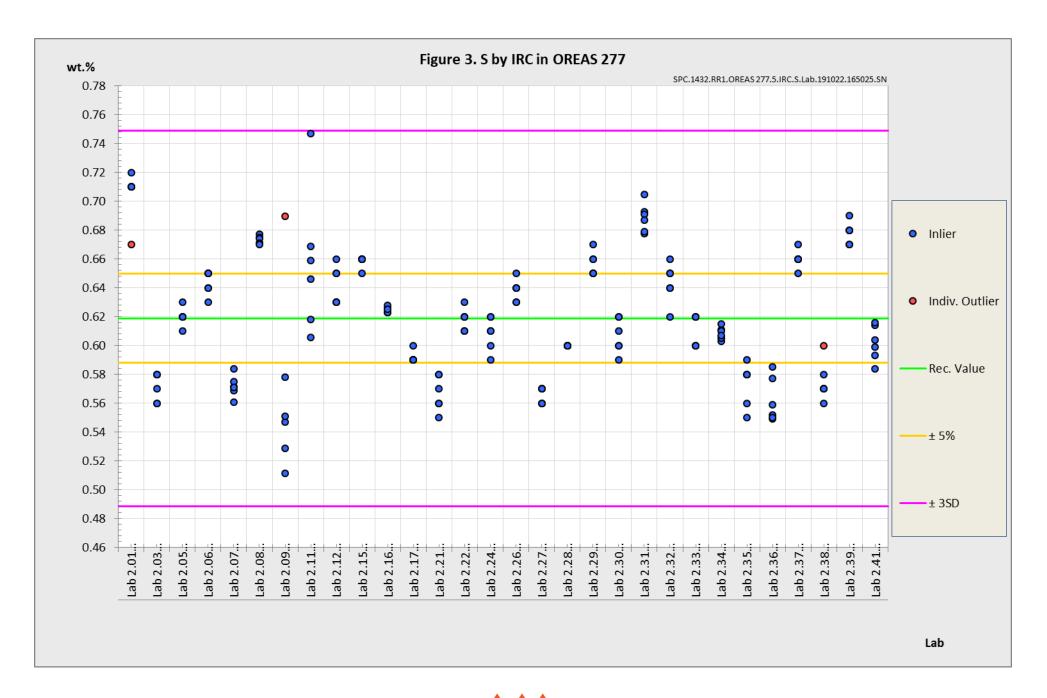


- 14. CERTIMIN, Lima, Peru
- 15. Chrysos Corporation, Perth, WA, Australia
- 16. Couer Mining Rochester Laboratory, Lovelock, Nevada, USA
- 17. Gekko Assay Labs, Ballarat, VIC, Australia
- 18. Information and Research Center, Kara-Balta, Chuy Region, Kyrgyzstan
- 19. Inspectorate (BV), Lima, Peru
- 20. Inspectorate America Corporation (BV), Sparks, Nevada, USA
- 21. Intertek Genalysis, Perth, WA, Australia
- 22. Intertek Tarkwa, Tarkwa, Ghana
- 23. Intertek Testing Services, Townsville, QLD, Australia
- 24. Intertek Testing Services Philippines, Cupang, Muntinlupa, Philippines
- 25. MinAnalytical Services, Kalgoorlie, WA, Australia
- 26. MinAnalytical Services, Perth, WA, Australia
- 27. Nagrom, Perth, WA, Australia
- 28. Nevada Gold Mines Assay Lab, Carlin, Nevada, USA
- 29. On Site Laboratory Services, Bendigo, VIC, Australia
- 30. Ontario Geological Survey, Sudbury, Ontario, Canada
- 31. PT Geoservices Ltd, Cikarang, Jakarta Raya, Indonesia
- 32. PT Intertek Utama Services, Jakarta Timur, DKI Jakarta, Indonesia
- 33. PT SGS Indo Assay Laboratories, Jakarta, Indonesia
- 34. Reminex Centre de Recherche, Marrakesh, Marrakesh-Safi, Morocco
- 35. Rio Tinto Kennecott Copper Central Laboratory, Kennecott, Utah, USA
- 36. Saskatchewan Research Council, Saskatoon, Saskatchewan, Canada
- 37. SGS, Randfontein, Gauteng, South Africa
- 38. SGS Australia Mineral Services, Kalgoorlie, WA, Australia
- 39. SGS Canada Inc., Vancouver, BC, Canada
- 40. SGS del Peru, Lima, Peru
- 41. SGS Lakefield Research Ltd, Lakefield, Ontario, Canada
- 42. SGS Tarkwa, Tarkwa, Western Region, Ghana
- 43. Shiva Analyticals Ltd, Bangalore North, Karnataka, India
- 44. Skyline Assayers & Laboratories, Tucson, Arizona, USA
- 45. UIS Analytical Services, Centurion, South Africa

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







## PREPARER AND SUPPLIER

Certified reference material OREAS 277 was prepared, certified and supplied by:



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## METROLOGICAL TRACEABILITY

The analytical samples were selected in a manner representative of the entire batch of the prepared CRM. This 'representivity' was maintained in each submitted laboratory sample batch and ensures the user that the data is traceable from sample selection through to the analytical results that underlie the consensus values. Each analytical data set has been validated by its assayer through the inclusion of internal reference materials and QC checks during analysis.

The laboratories were chosen on the basis of their competence (from past performance in inter-laboratory programs undertaken by ORE Pty Ltd) for a particular analytical method, analyte or analyte suite and sample matrix. Most of these laboratories have and maintain ISO 17025 accreditation. The certified values presented in this report are calculated from the means of accepted data following robust statistical treatment, as detailed in this report.

Guide ISO/TR 16476:2016, section 5.3.1 describes metrological traceability in reference materials as it pertains to the transformation of the measurand. In this section it states, "Although the determination of the property value itself can be made traceable to appropriate units through, for example, calibration of the measurement equipment used, steps like the transformation of the sample from one physical (chemical) state to another cannot. Such transformations may only be compared with a reference (when available), or among themselves. For some transformations, reference methods have been defined and may be used in certification projects to evaluate the uncertainty associated with such a transformation. In other cases, only a comparison among different laboratories using the same method is possible. In this case, certification takes place on the basis of agreement among independent measurement results (see ISO Guide 35:2006, Clause 10)."

#### COMMUTABILITY

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

## **INTENDED USE**

OREAS 277 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 277 may be used to calibrate the entire procedure by producing a pure substance CRM transformed into a calibration solution.

OREAS 277 is intended for the following uses:

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

# 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. Aqua regia is a partial empirical digest and differences in recoveries for various analytes are commonplace. These are caused by variations in the digest conditions which can include the ratio of nitric to hydrochloric acids, acid strength, temperatures, leach times and secondary digestions. Recoveries for sulphide-hosted base metal sulphides approach total values, however other analytes, in particular the lithophile elements, show greater sensitivity to method parameters. This can result in lack of consensus in an inter-laboratory certification program for these elements.

The approach applied here is to report certified values in those instances where reasonable agreement exists amongst a majority of participating laboratories. The results from 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.

# 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 inter-laboratory 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.

## STABILITY AND STORAGE INSTRUCTIONS

OREAS 277 has been prepared from primary gold ore blended with barren sediments. It is low in reactive sulphide (0.62 wt.% S) and in its unopened state and under normal conditions of storage has a shelf life beyond ten years. Its stability will be monitored at regular intervals and purchasers notified if any changes are observed.

## INSTRUCTIONS FOR CORRECT USE

The certified values for OREAS 277 refer to the concentration level in its packaged state. It should not be dried prior to weighing and analysis.

# HANDLING INSTRUCTIONS

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

# **LEGAL NOTICE**

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

# **DOCUMENT HISTORY**

Revision No.	Date	Changes applied
2	18 <sup>th</sup> Dec, 2020	Status of specific forms of C and S changed from Certified to Indicative.
1	25 <sup>th</sup> Aug 2020	Corrected indicative values for XRF, laser ablation and LOI¹000°C.
0	23 <sup>rd</sup> Oct 2019	First publication.

# QMS CERTIFICATION

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.





## **CERTIFYING OFFICER**

18<sup>th</sup> December, 2020

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



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