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#### CERTIFICATE OF ANALYSIS FOR

# **COPPER-GOLD STANDARD**

## **OREAS 50P**

## SUMMARY STATISTICS

Recommended Values, 95% Confidence and Tolerance Intervals

Constituent	Recommended value	95% Confidence interval		Toleranc 1-α=0.99	e interval 9, ρ=0.95
		Low	High	Low	High
Gold, Au (ppm)	0.727	0.706	0.748	0.723	0.731
Copper, Cu (%)	0.691	0.681	0.701	0.681	0.701

Prepared by: Ore Research & Exploration Pty Ltd April 2004

## INTRODUCTION

OREAS reference materials (RMs) are intended to provide a low cost method of evaluating and improving the quality of precious and base metal analysis of geological samples. To the analyst they provide an effective means of calibrating analytical equipment, assessing new techniques and routinely monitoring in-house procedures. To the explorationist they provide an important control in analytical data sets related to exploration from the grass roots level through to prospect evaluation. To the mine geologist they provide a valuable tool in grade control and QA/QC management programs. Following the implementation of new processing technology Ore Research & Exploration now produces gold RMs exhibiting a level of homogeneity previously unattainable. In certain instances RMs produced from a single source are sufficiently homogeneous to produce a relatively coarse-grained form designed to simulate drill chip samples. These have a grain size of minus 3mm and are designated with a "C" suffix to the RM identification number. These standards are packaged in 0.5-1kg units following homogenisation and are intended for submission to analytical laboratories in subsample sizes of as little as 250g. They offer the added advantages of providing a check on both sample preparation and analytical procedures while acting as a blind standard to the assay laboratory. The more conventional pulped standards have a grain size of minus 20 to minus 75 microns and a higher degree of homogeneity. These standards are distinguished by a "P" suffix to the standard identification number. In line with ISO recommendations successive batch numbers are now designated by the lower case suffixes "a", "b", "c", "d", etc.

### SOURCE MATERIALS

Reference material OREAS 50P is one of four porphyry copper-gold standards prepared from ore samples from the Northparkes Mine, central western New South Wales, Australia.

Mineralisation in the region is hosted by a sequence of late Ordovician to Early Silurian volcanics, intrusives and sediments that occur within the Bogan Gate Synclinorial Zone of the Lachlan Fold Belt. The western portion of this zone is dominated by volcanics and host to the Goonumbla porphyry copper-gold deposits. The Late Ordovician Goonumbla Volcanics host the Northparkes deposits and are interpreted to have erupted from shallow water to partly emergent volcanic centres. They exhibit a broad range in composition from shoshonite through to latite to trachyte.

Coeval sub-volcanic quartz monzonite porphyries (and attendant mineralisation) have intruded the volcanics. They are generally small, sub-vertical, pipe-like intrusives. Typically the mineralised porphyries contain plagioclase and quartz phenocrysts in a matrix of finegrained potassium feldspar and quartz with minor biotite and hornblende.

Copper-gold mineralisation occurs as stockwork quartz veins and disseminations associated with potassic alteration. This alteration is intimately associated spatially and temporally with the small finger-like quartz monzonite porphyries that intrude the Goonumbla Volcanics. Sulphides are zoned laterally from the centres of mineralisation. The central portions are bornite-rich with minor chalcopyrite, zoning outward through equal concentrations of bornite and chalcopyrite, to an outermost chalcopyrite-rich zone. Pyrite increases outward at the expense of bornite.

## **COMMINUTION AND HOMOGENISATION PROCEDURES**

The material was prepared in the following manner:

- a) drying;
- b) crushing and screening;
- c) preliminary homogenisation;
- d) milling to minus 20 microns;
- e) final homogenisation;
- f) bagging into 20kg sublots.

## **ANALYSIS OF OREAS 50P**

The indicative major and trace element composition of OREAS 50P is given in Table 1. The constituents are the means of duplicate XRF analyses determined using a borate fusion method at the University of Melbourne, Victoria, Australia, and are uncertified values.

Table 1. Indicative major and trace element composition of reference material OREAS 50P;  $SiO_2$  to Total in weight percent (Total includes traces); rest in parts per million.

Constituent	Concentration	Constituent	Concentration
SiO <sub>2</sub>	55.64	Ba	714
TiO <sub>2</sub>	0.74	CI	169
$AI_2O_3$	17.46	Со	25
Fe <sub>2</sub> O <sub>3</sub>	7.41	Cr	32
MnO	0.12	Ga	22
MgO	3.43	Nb	1
CaO	2.22	Nd	16
Na <sub>2</sub> O	5.59	Ni	34
K <sub>2</sub> O	3.38	Rb	53
$P_2O_5$	0.43	Sc	22
SO <sub>3</sub>	0.21	Sr	693
LOI	2.63	Th	17
Total	100.12	V	252
		Y	19
		Zn	61
		Zr	70

Fifteen commercial laboratories participated in the certification program for gold and copper and are listed in the section headed Participating Laboratories. To maintain anonymity laboratories have been randomly assigned a number code 1 through 15. Their results together with uncorrected means, medians, one sigma standard deviations, relative standard deviations and percent deviation of lab means from the corrected mean of means (PDM<sup>3</sup>) are presented in Tables 2 to 4. The parameter PDM<sup>3</sup> is a measure of laboratory accuracy while the relative standard deviation is an effective measure of analytical precision where homogeneity of the test material has been confirmed. The analytical

methods employed by each laboratory are given in the table captions. With the exception of Becquerel, six 110g samples were submitted to each laboratory for analysis. These samples were duplicate scoop splits from three separate 1kg test units taken during the bagging stage. This two-stage nested design for the interlaboratory program was amenable to analysis of variance (ANOVA) treatment and enabled a comparative assessment of within- and between-unit homogeneity. The twenty-six INAA samples, on which much of the homogeneity evaluation is based, were also taken at regular intervals throughout the bagging stage and are considered representative of the entire batch.

Gold was determined in six replicate assays using a fire assay technique (40-50g charge with new pots) with flame AAS or ICPOES finish at thirteen laboratories (Table 2), while Becquerel determined gold in twenty-six samples via instrumental neutron activation analysis (INAA) using 0.5gm analytical subsample weights (Table 3). Copper was determined via four acid (HF-HNO<sub>3</sub>-HCIO<sub>4</sub>-HCI) digest with ICPOES or AAS finish (Table 4).

Table 2.Analytical results for gold in standard OREAS 50P (FA\*AAS - fire assay / atomic<br/>absorption spectrometry; FA\*OES - fire assay / inductively coupled plasma optical<br/>emission spectrometry; INAA - instrumental neutron activation analysis; Std.Dev.<br/>and Rel.Std.Dev. are one sigma values; PDM³ is percent deviation of lab mean from<br/>corrected mean of means; outliers in bold; values in parts per billion.

Replicate	Lab 1	Lab 2	Lab 3	Lab 4	Lab 5	Lab 6	Lab 7
Number	FA*AAS	FA*AAS	FA*AAS	FA*AAS	FA*OES	FA*AAS	FA*AAS
1	740	720	695	700	785	585	789
2	740	780	684	730	780	630	672
3	740	730	705	720	776	655	697
4	730	730	700	720	778	660	706
5	720	750	700	720	753	650	704
6	730	740	709	700	774	665	700
Mean	733	742	699	715	774	641	711
Std. Dev.	8	21	9	12	11	30	40
Rel.Std.Dev.	1.11%	2.88%	1.25%	1.71%	1.43%	4.67%	5.62%
PDM <sup>3</sup>	0.81%	1.95%	-3.94%	-1.71%	6.44%	-11.9%	-2.22%

Table 2.	Continued.						
Replicate	Lab 8	Lab 9	Lab 10	Lab 11	Lab 12	Lab 13	Lab 14
Number	FA*AAS	FA*AAS	FA*AAS	FA*AAS	FA*AAS	FA*OES	INAA
1	764	700	710	733	695	717	
2	785	734	710	733	681	702	
3	792	679	770	735	704	747	Refer to
4	694	706	780	740	708	745	Table 3
5	775	725	740	743	692	728	
6	786	710	770	746	684	734	
Mean	766	709	747	738	694	729	772
Std. Dev.	37	19	31	6	11	17	15
Rel.Std.Dev.	4.78%	2.73%	4.21%	0.75%	1.54%	2.36%	2.00%
PDM <sup>3</sup>	5.30%	-2.54%	3.65%	1.64%	-4.63%	0.51%	6.13%

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Table 3. Analytical results for gold in standard OREAS 50P via	3
instrumental neutron activation analysis using a 0.5g analytica	
subsample weight (abbreviations as in Table 2; values in parts	3
per billion).	

Replicate	Lab 14	
Number	INAA	
1	783	
2	757	
3	797	
4	781	
5	769	
6	791	
7	753	
8	761	
9	782	
10	788	
11	790	
12	756	
13	771	
14	784	
15	773	
16	778	
17	757	
18	748	
19	796	
20	778	
21	791	
22	754	
23	770	
24	758	
25	760	
26	749	
Mean	772	
Std. Dev.	15	
Rel.Std.Dev.	2.00%	

Table 4. Analytical results for copper in standard OREAS 50P (4AD\*OES - four acid digest / inductively coupled plasma optical emission spectrometry; 4AD\*AAS - four acid digest / atomic absorption spectrometry, other abbreviations as in Table 2; values in parts per million).

Replicate	Lab 1	Lab 2	Lab 3	Lab 4	Lab 5	Lab 6	Lab 7
Number	4AD*OES	4AD*OES	4AD*OES	4AD*AAS	4AD*OES	4AD*OES	4AD*OES
1	6530	6030	6930	6880	6960	7000	6860
2	6660	6270	7130	6840	6840	6850	6940
3	6570	6070	7050	6970	6800	6900	7050
4	6670	5890	7310	6530	6800	6850	6850
5	6740	5940	6970	6730	6880	6900	7010
6	6590	6070	6970	7110	6990	6800	6910
Mean	6627	6045	7060	6843	6878	6883	6937
Std. Dev.	77	132	142	200	81	68	80
Rel.Std.Dev.	1.16%	2.18%	2.01%	2.92%	1.18%	0.99%	1.16%
PDM <sup>3</sup>	-4.11%	-12.52%	2.16%	-0.97%	-0.46%	-0.39%	0.38%

Table 4.	Continued.						
Replicate	Lab 8	Lab 9	Lab 10	Lab 11	Lab 12	Lab 13	Lab 15
Number	4AD*AAS	4AD*AAS	4AD*OES	4AD*AAS	4AD*AAS	4AD*OES	4AD*OES
1	7200	7100	6901	6700	6840	6940	6800
2	7300	7000	6963	6720	6770	6930	6940
3	7200	6900	6957	6760	6860	6940	6880
4	7600	6900	6961	6660	6810	6970	6930
5	7500	6800	6962	6670	6800	6950	7020
6	7200	6900	6987	6680	6790	6970	6980
Mean	7333	6933	6955	6698	6812	6950	6925
Std. Dev.	175	103	29	37	33	17	77
Rel.Std.Dev.	2.39%	1.49%	0.41%	0.55%	0.49%	0.24%	1.12%
PDM <sup>3</sup>	6.12%	0.33%	0.65%	-3.07%	-1.43%	0.21%	0.57%

# STATISTICAL EVALUATION OF ANALYTICAL DATA FOR OREAS 50P

#### **Recommended Value and Confidence Limits**

The certified value is the mean of means of accepted replicate values of accepted participating laboratories computed according to the formulae

$$\overline{x}_i = \frac{1}{n_i} \sum_{j=1}^{n_i} x_{ij}$$

$$\ddot{x} = \frac{l}{p} \sum_{i=1}^{p} \bar{x}_i$$

where

 $x_{ij}$  is the jth result reported by laboratory i; p is the number of participating laboratories;  $n_i$  is the number of results reported by laboratory i;  $\overline{x_i}$  is the mean for laboratory i;  $\ddot{x}$  is the mean of means.

The confidence limits were obtained by calculation of the variance of the consensus value (mean of means) and reference to Student's-*t* distribution with degrees of freedom (*p*-1).

$$\hat{V}(\ddot{x}) = \frac{1}{p(p-1)} \sum_{i=1}^{p} (\bar{x}_i - \ddot{x})^2$$
Confidence limits =  $\ddot{x} \pm t_{1-x/2} (p-1) (\hat{V}(\ddot{x}))^{1/2}$ 

where  $t_{1-x/2}(p-1)$  is the 1-x/2 fractile of the t-distribution with (p-1) degrees of freedom.

The distribution of the values are assumed to be symmetrical about the mean in the calculation of the confidence limits.

The test for rejection of individual outliers from each laboratory data set was based on *z* scores (rejected if  $|z_i| > 2.5$ ) computed from the robust estimators of location and scale, *T* and *S*, respectively, according to the formulae

$$S = 1.483 \text{ median } / x_j - \text{median } (x_i) /$$
$$z_i = \frac{x_i - T}{S}$$

where

*T* is the median value in a data set; *S* is the median of all absolute deviations from the sample median multiplied by 1.483, a correction factor to make the estimator consistent with the usual parameter of a normal distribution.

Individual outliers and, more rarely, laboratory means deemed to be outlying are shown in bold italics and have been omitted in the determination of recommended values.

Constituent	Recommended value	95% Confidence interva	
		Low	High
Gold, Au (ppm)	0.727	0.706	0.748
Copper, Cu (wt. %)	0.691	0.681	0.701

Table 5. Recommended values and 95% confidence intervals for OREAS 50P.

#### Statement of Homogeneity

The standard deviation of each laboratory data set includes error due to both the imprecision of the analytical method employed and to possible inhomogeneity of the material analysed. The standard deviation of the pooled individual analyses of all participating laboratories includes error due to the imprecision of each analytical method, to possible inhomogeneity of the material analysed and, in particular, to deficiencies in accuracy of each analytical method. In determining tolerance intervals for copper that component of error attributable to measurement inaccuracy was eliminated by transformation of the individual results of each data set to a common mean (the uncorrected grand mean) according to the formula

$$x'_{ij} = x_{ij} - \overline{x}_i + \frac{\sum_{i=1}^p \sum_{j=1}^{n_i} x_{ij}}{\sum_{i=1}^p n_i}$$

where

 $x_{ij}$  is the jth raw result reported by laboratory i;  $x'_{ij}$  is the jth transformed result reported by laboratory i;  $n_i$  is the number of results reported by laboratory i; p is the number of participating laboratories;  $\overline{x}_i$  is the raw mean for laboratory i.

The homogeneity of each constituent was determined from tables of factors for two-sided tolerance limits for normal distributions (ISO 3207) in which

Lower limit is  $\ddot{x} - k'_2(n, p, l - \alpha)s''_g$ Upper limit is  $\ddot{x} + k'_2(n, p, l - \alpha)s''_g$ 

where

*n* is the number of results;  $1-\alpha$  is the confidence level; *p* is the proportion of results expected within the tolerance limits;  $k'_2$  is the factor for two-sided tolerance limits (*m*,  $\alpha$  unknown);  $s''_{\pi}$  is the corrected grand *s* tan dard deviation.

The meaning of these tolerance limits may be illustrated for copper, where 99% of the time at least 95% of subsamples will have concentrations lying between 0.681% and 0.701%. 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 (IS0 Guide 35).

The corrected grand standard deviation,  $s_g$ , used to compute the tolerance intervals is the weighted means of standard deviations of all data sets for a particular constituent according to the formula

$$s''_{g} = \frac{\sum_{i=1}^{p} (s_{i}(1 - \frac{s_{i}}{s'_{g}}))}{\sum_{i=1}^{p} (1 - \frac{s_{i}}{s'_{g}})}$$

where

 $1 - \left(\frac{s_i}{2s'_{\varphi}}\right)$  is the weighting factor for laboratory *i*;

 $s^{\prime}_{g}$  is the grand standard deviation computed from the transformed (i.e. means -

adjusted) results

according to the formula

$$s'_{g} = \left[\frac{\sum_{i=j}^{p} \sum_{j=i}^{n_{i}} (x'_{ij} - \overline{x}'_{i})^{2}}{\sum_{i=l}^{p} n_{i} - l}\right]^{1/2}$$

where  $\bar{x}'_i$  is the transformed mean for laboratorty *i* 

The weighting factors were applied to compensate for the considerable variation in analytical precision amongst participating laboratories. Hence, weighting factors for each data set have been constructed so as to be inversely proportional to the standard deviation of that data set. It should be noted that estimates of tolerance by this method are considered conservative as a significant proportion of the observed variance, even in those laboratories exhibiting the best analytical precision, can presumably be attributed to measurement error.

For gold a more simplified procedure was used in the determination of homogeneity. This entailed using the high precision INAA data alone, obtained on an analytical subsample weight of 0.5gm (compared to 40-50gm for the fire assay method). By employing a sufficiently reduced subsample weight in a series of determinations by the same method, analytical error becomes negligible in comparison to subsampling error. The corresponding standard deviation at a 50gm subsample weight can then be determined from the observed standard deviation of the 0.5gm data using the known relationship between the two parameters (Kleeman, 1967). The homogeneity of gold was then determined from tables of factors for two-sided tolerance limits for normal distributions. The high level of repeatability indicated by the low coefficients of variation in Table 2 and the 0.5gm Becquerel data) is consistent with the very narrow calculated tolerance interval and is confirmation of the excellent homogeneity of gold in OREAS 50P.

Constituent	Recommended value	Tolerance interval 1-α=0.99, ρ=0.95	
		Low	High
Gold, Au (ppm)	0.727	0.723	0.731
Copper, Cu (wt. %)	0.691	0.681	0.701

Table 6. Recommended values and tolerance limits for OREAS 50P.

No outliers were removed from the INAA results prior to the calculation of tolerance intervals for gold, although for copper, outliers were removed prior to the calculation of  $s_{q'}$ 

and a weighting factor of zero was applied to those data sets where  $s_l/2s_g' > 1$  (i.e. where the weighting factor 1-  $s_l/2s_g' < 0$ ).

# PARTICIPATING LABORATORIES

Acme Analytical Laboratories, Vancouver, BC, Canada Actlabs Pacific Pty Ltd, Redcliffe, WA, Australia ALS Chemex, Santiago, Chile ALS Chemex, Sparks, Nevada, USA Amdel Laboratories, Thebarton, SA, Australia Amdel Laboratories, Wangara, WA, Australia Becquerel Laboratories Inc, Lucas Heights, NSW, Australia Cantech Laboratories Inc, Calgary, Canada Cone Geochemical, Lakewood, Colorado, USA Genalysis Laboratory Services, Maddington, WA, Australia McPhar Geoservices (Phil.) Inc., Makati, Philippines SGS, Welshpool, WA, Australia SGS, Garbutt, QLD, Australia Ultra Trace, Canning Vale, WA, Australia Intertek Testing Services, Jakarta, Indonesia

## PREPARER AND SUPPLIER OF THE REFERENCE MATERIAL

The copper-gold ore reference material, OREAS 50P has been prepared and certified and is supplied by:

Ore Research & Exploration Pty Ltd 6 – 8 Gatwick Road Bayswater North VIC 3153 AUSTRALIA

Telephone	(03) 9729 0333	International	+613-9729 0333
Facsimile	(03) 9729 4777	International	+613-9729 4777
Email	info@ore.com.au	Web	www.ore.com.au

It is available in unit sizes of 60g laminated foil packets.

### INTENDED USE

OREAS 50P is a reference material intended for the following:

- i) for the calibration of instruments used in the determination of the concentration of gold and copper;
- ii) for the verification of analytical methods for gold and copper;
- iii) for the preparation of secondary reference materials of similar composition;
- iv) as an arbitration sample for commercial transactions.

## STABILITY AND STORAGE INSTRUCTIONS

OREAS 50P has been prepared from a sulphide-poor mineralised quartz monzonite porphyry sample. The robust foil laminate film used to package it is an effective barrier to oxygen and moisture and the sealed CRM is considered to have long-term stability under normal storage conditions.

## INSTRUCTIONS FOR THE CORRECT USE OF THE REFERENCE MATERIAL

The recommended values for OREAS 50P refers to the concentration levels of gold and copper after removal of hygroscopic moisture by drying in air to constant mass at 105<sup>°</sup> C. In its packaged state a hygroscopic moisture content of 1.11% has been established. If the reference material is not dried by the user prior to analysis, the recommended values should be corrected to the moisture-bearing basis.

## 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.

# **CERTIFYING OFFICER:** Dr Paul Hamlyn

### REFERENCES

Ingamells, C. O. and Switzer, P. (1973), Talanta 20, 547-568. ISO Guide 35 (1985), Certification of reference materials - General and statistical principals. ISO Guide 3207 (1975), Statistical interpretation of data - Determination of a statistical tolerance interval.

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