

QOTHO CERTIFIED REFERENCE MATERIAL (QCRM)

QCRM-5-131

COPPER ORE

CERTIFICATE OF ANALYSIS

CERTIFIED VALUES			
ANALYTES	UNITS	CONCENTRATIONS	EXPANDED UNCERTAINTY
Co	%	0.30	±0.02
Cu	%	1.36	±0.02
Mg	%	1.56	±0.11
ASSIGNED VALUES (FOR INFORMATION ONLY)			
ANALYTES	UNITS	CONCENTRATIONS	EXPANDED UNCERTAINTY
Al	%	2.84	±0.29
Cu (Soluble)	%	1.28	±0.08
Fe	%	1.13	±0.07
Mn	%	0.057	±0.005
Ni	%	0.005	±0.003
Si	%	38.8	±1.7
Zn	%	0.008	±0.003

1. Use:

QCRM-5-131 is a certified reference material which is suitable for use as random control samples in routine analytical laboratory quality control, when inserted within a batch of samples and measured in parallel to the unknown. The QCRM can also be used as a control sample in the analysis of samples of a similar type, verification of analytical methods for Copper Ore and as a calibration standard for the calibration of equipment used for analysing similar materials.

2. Origin of Material:

This standard was sponsored by Metalkol RTR Operation (ERG Africa) in the Kolwezi district of the Democratic Republic of Congo.

3. Mineral and Chemical Composition:

The Copper belt forms one of the world's greatest metallogenic provinces containing over a third of the world's cobalt reserves and a tenth of the world's copper reserves. The oxide ore mineralogy is composed predominantly of malachite, pseudo-malachite, minor chrysocolla and heterogenite. Underlying sulphide mineralization comprises chalcopyrite, chalcocite, and bornite. This tail sample is the waste-product from the Metalkol Roan Tailings Reclamation (RTR) plant.

4. Date of Initial Issue:

20 March 2024.

5. Packaging & Handling instructions:

The material was packaged as 100g unit sizes, placed in geo-envelopes, within a vacuum sealed aluminium foil bag. Open the seal of the foil with care and shake or otherwise agitate prior to use. Normal safety precautions for handling fine particulate matter are recommended, such as the use of safety glasses, breathing protection, gloves, and a laboratory coat. Once opened, material must be stored in a cool, dry environment. Results on page 1 is presented on dry basis. Analysis should therefore be done on dry basis, after drying to constant mass, at 105 degrees Celsius.

6. Method of Preparation:

The material was sieved through a 75-micron screen and the oversize was re-milled to ensure 100% passing through the screen. It was then blended, systematically divided, and packaged into 100-gram zip-lock bags. Randomly selected samples, from the bags, were tested in-house via XRF, to confirm homogeneity. Once confirmed and certification completed, the items were placed in geo-envelopes and vacuum sealed in aluminium foil bags.

7. Methods of Analysis used:

- Multi-acid digestion with AAS or ICP-OES finish
- Sodium peroxide fusion with ICP-OES finish
- Powder samples with XRF finish
- Fused beads with XRF finish
- Iron & Copper by multi-acid digestion and potentiometric finish.
- Silica by multi-acid digestion and gravimetric finish.
- Acid soluble copper by weak sulphuric acid leach and AAS finish
- Sulphur by thermal combustion analysis.

8. Analysis required:

An instruction letter was sent to all participants. The analysis required was noted in the instruction letter and reporting template, including but not limited to Ag, Al, As, Au, Ca, Co, Cu, Cu (Soluble), Fe, Mg, Mn, Ni, Pb, S, Si, U and Zn.

9. Participating Laboratories:

No	LABORATORY	COUNTRY
1.	AHK DRC South	Democratic Republic of Congo
2.	AHK Kitwe	Zambia
3.	ALS Kansanshi	Zambia
4.	Ero Brasil Caraiba	Brazil
5.	Intertek Tschudi	Namibia
6.	Kamoto Copper Company	Democratic Republic of Congo
7.	Lubambe Copper Mine	Zambia
8.	Metalkol	Democratic Republic of Congo
9.	Mitra Sk South Africa	South Africa
10.	Mopani Nkana	Zambia
11.	Mutanda Mining	Democratic Republic of Congo
12.	Palabora Mining Company	South Africa
13.	Quality Lab Services Nababeeb	South Africa
14.	Robinson International	Democratic Republic of Congo
15.	UIS Analytical Services	South Africa
16.	Zambia Revenue Authority	Zambia

10. Assay Data:

Data used for Assigning Values and Certification.

LABORATORY	Ag	Al	As	Ca	Co	Cu	Cu (Soluble)	Fe	Mg	Mn	Ni	Pb	S	Si	Zn
UNIT	g/t	%	ppm	%	%	%	%	%	%	%	%	%	%	%	%
LAB001		2.768						1.22						39.076	
LAB002						1.364									
LAB003		2.98		0.212	0.307	1.397		1.234	1.599	0.062				39.151	
LAB004		2.906			0.296	1.355		1.187	1.595	0.054			0.038	38.5	0.007
LAB005					0.291	1.366	1.107	1.156	3.16	0.062		< 0.010	0.041	39.49	
LAB006		2.795			0.289	1.353	1.265	1.083	1.534	0.053	< 0.010				0.007
LAB007							1.3								
LAB008						1.386									
LAB009		2.332		0.363	0.297	1.369	1.194	1.122	1.204	0.052	0.005			21.065	0.011
LAB010				0.487	0.309	1.37		1.115	1.692	0.06	0.006				0.015
LAB011				0.315	0.289	1.393	1.375	0.96	1.506	0.051			0.08		
LAB012						1.29									
LAB013						1.335									
LAB014		2.871		0.355	0.319		1.281	1.164	1.662	0.056	0.005				0.006
LAB015			28.15	0.195	0.288			1.09	1.498	0.058	0.007	0.01	0.066	37.945	0.01
LAB016						1.28									
LAB017						1.43									
LAB018							1.29								
LAB019	2.733	3.143		0.213		1.368	1.37	1.068	1.553	0.055					0.005
LAB020	0.993	4.555	103.95	0.13	0.3	1.355	1.32	1.17	1.58	0.059	0.003	0.004	0.028		0.006

11. Method of Certification:

QM is a SANAS Accredited Proficiency Testing Scheme Provider, No. PTS0012

This material was distributed as test items, in the Qotho Copper PT Round 5 of 2022. The participating laboratories were each given 1 randomly selected sample from the batch, to analyse and report on in duplicate. Some laboratories reported results via more than one analytical method. Obvious blunders were resolved with the laboratories (if any), after which the data was processed using Robust Statistics, through PROLab Plus.

Not all the participating laboratories were accredited. Historical performance in Qotho PT Schemes, as well as an evaluation of the CRM QA/QC data generated by the laboratories, during the analysis of this QRM, were considered, to evaluate the competence of laboratories. Where competence could not be confirmed, the affected data was deselected from the dataset. Certification of analytes was then done, provided that a minimum of 10 qualifying datapoints remained available.

Where analytes could not be certified, estimate concentrations were assigned, using all the data in the dataset.

12. Measurement of Uncertainty:

Standard uncertainty, u_{CRM} , was calculated according to ISO 13528 (equation 6), and it includes the effects of uncertainty due to inhomogeneity, transport, potential instability, and laboratory uncertainty. Because of all the uncertainties under consideration, QM further applies an expanded uncertainty, for certification purposes. The measurement uncertainty of the certified value is therefore calculated as follows: $U_{CRM} = k u_{CRM}$, where k is a coverage factor, which is determined from the Student's t -distribution, based on the degrees of freedom, per analyte.

This presents a certified value, as follows: $x_{CRM} \pm U_{CRM}$.

Measurement uncertainty for Assigned values, are calculated in the same manner.

Laboratories which prefer to use the 95% measurement uncertainty, rather than the expanded uncertainty, all available information relating to measurement uncertainty of the certified/assigned values, are given below:

Analyte	Unit of measure	s (Standard Deviation of Dataset)	ν (Degrees of Freedom)	k (Coverage Factor)	u (Standard Uncertainty)	95% Measurement Uncertainty	Expanded Uncertainty
Al	%	0.286	7	2.365	0.123	± 0.25	± 0.29
Co	%	0.011	9	2.262	0.006	± 0.01	± 0.02
Cu	%	0.033	14	2.145	0.01	± 0.02	± 0.02
Cu (Soluble)	%	0.088	8	2.306	0.034	± 0.07	± 0.08
Fe	%	0.093	11	2.201	0.031	± 0.06	± 0.07
Mg	%	0.125	10	2.228	0.047	± 0.09	± 0.11
Mn	%	0.005	10	2.228	0.002	± 0.004	± 0.005
Ni	%	0.002	4	2.776	0.001	± 0.002	± 0.003
Si	%	1.223	5	2.571	0.634	± 1.3	± 1.7
Zn	%	0.003	7	2.365	0.001	± 0.002	± 0.003

13. Metrological Traceability:

The values quoted herein are based on the consensus values derived from statistical analysis of the data from an inter laboratory measurement program. Traceability to SI units is via the accredited laboratories, as ISO 17025 requires laboratories to use CRM's traceable to the SI units, during the calibration of their equipment. Not all laboratories were accredited.

Fortunately, most laboratories reported on the QA/QC CRMs used during the analysis of this QRM and reported the values obtained during the sample run. Evaluation of their QA/QC performance serves as further evidence of metrological traceability.

Equivalence tests were performed on all analytes to determine whether the metrologically traceable data and those for which traceability evidence was not available, could be treated as equal (at a level of significance of $\alpha = 0.05$). Where equivalent, all the data was used. Where not equivalent, only the metrologically traceable data was considered.

14. Minimum sample size:

The recommended minimum sample size for the use of this material is as per the participants method validation criteria.

15. Period of validity:

The certified values are valid for this product, while still sealed in its original packaging, for a minimum period of 5 years from date of Initial Certification. Stability monitoring of inventory will be done at regular intervals. Any concerns regarding potential instability of the material, will immediately be communicated to all consumers.

16. Legal:

This certificate and the reference material described in it were prepared with due care and attention. The requirements of ISO Guide 31, ISO/IEC 17043 and ISO 17034 were followed in the preparation of this reference material and certificate of analysis.

Qotho Minerals, however, accepts no liability for any decisions or actions taken following the use of the reference material. The company has a complaints procedure, which will be made available upon request, should there be any dissatisfaction with either the product or the COA.

Certifying & Technical Signatories	
<i>Dr Hannelie de Beer (Pr. Sci. Nat.)</i>	<i>Takudzwa Tsapayi (Pr. Sci. Nat.)</i>
20 March 2024	

This Certificate of Analysis (CoA) has been electronically signed using an Advanced Electronic Signature (AES) in terms of the Electronic Communications and Transactions Act No. 15, 2002 (ECT Act). Any amendments to the CoA can be detected by reference to the Signature Panel displayed in the electronic pdf version of the CoA.

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