

**Chemical and structural characterization of UICC crocidolite fibres from Koegas Mine, Northern Cape (South Africa)**Alessandro Pacella<sup>1,\*</sup>, Giovanni B. Andreozzi<sup>1</sup>, Luca Nodari<sup>2</sup>, Paolo Ballirano<sup>1</sup><sup>1</sup> Department of Earth Sciences, Sapienza University of Rome, Piazzale Aldo Moro 5, I-00185 Rome, Italy<sup>2</sup> Institute of Condensed Matter Chemistry and Technologies for Energy (ICMATE), National Research Council of Italy (CNR), Corso Stati Uniti 4, 35127 Padova, Italy**ARTICLE INFO**

Submitted: August 2019

Accepted: October 2019

Available on line: November 2019

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DOI: 10.2451/2019PM910

How to cite this article:

Pacella A. et al. (2019)

Period. Mineral. 88, 297-306

**ABSTRACT**

In the present work we report the full structural and spectroscopic characterization of an UICC crocidolite standard sample, that is a fibrous riebeckite from Koegas Mine, Northern Cape (South Africa). The chemical composition was obtained by SEM-EDS and cation site partition was retrieved by complementing chemical, Mössbauer and X-ray powder diffraction data. Cell parameters, fractional coordinates, and site scattering for  $M(1)$ ,  $M(2)$ ,  $M(3)$ ,  $M(4)$  were refined using the Rietveld method.

The UICC crocidolite standard sample shows the chemical formula  $^A\text{Na}_{0.026}^B\text{Na}_{2.000}^C(\text{Fe}^{2+}_{2.212}\text{Fe}^{3+}_{2.041}\text{Mg}_{0.747})_{\Sigma=5.000}^T(\text{Si}_{7.950}\text{Al}_{0.022})_{\Sigma=7.972}\text{O}_{22}^{\text{O}(3)}(\text{OH})_2$ , very close to that of the end-member riebeckite  $^A\text{Na}_{2.000}^B\text{Na}_{2.000}^C(\text{Fe}^{2+}_3\text{Fe}^{3+}_2)_{\Sigma=5}^T\text{Si}_8\text{O}_{22}^{\text{O}(3)}(\text{OH})_2$ , the major difference being the  $\text{Fe}^{3+}/\text{Fe}_{\text{tot}}$  ratio of 0.48, slightly higher than the ideal ratio of 0.40. Cation site distribution of the UICC crocidolite standard sample is:  $[\text{Mg}_{0.50}\text{Fe}^{3+}_{0.31}\text{Fe}^{2+}_{1.19}]_{\Sigma=2.00}$  at  $M(1)$ ;  $[\text{Mg}_{0.10}\text{Fe}^{3+}_{1.60}\text{Fe}^{2+}_{0.30}]_{\Sigma=2.00}$  at  $M(2)$ ;  $[\text{Mg}_{0.15}\text{Fe}^{3+}_{0.13}\text{Fe}^{2+}_{0.72}]_{\Sigma=1.00}$  at  $M(3)$ . Refined cell parameters are:  $a=9.73516(19)$  Å,  $b=18.0453(3)$  Å,  $c=5.32895(9)$  Å,  $\beta=103.5159(12)^\circ$ ,  $V=910.23(3)$  Å<sup>3</sup>. Quantitative Phase Analysis indicates that 93.7(3) wt% of the sample consists of crocidolite, the remaining being distributed among five different phases including magnetite and quartz.

Keywords: asbestos; UICC crocidolite; Scanning Electron Microscopy with Energy Dispersive System (SEM-EDS); Mössbauer Spectroscopy; X-ray Powder Diffraction (XRPD), Rietveld method; surface reactivity.

**INTRODUCTION**

Asbestos is a group of six naturally-occurring silicate including one fibrous serpentine (chrysotile) and five fibrous amphiboles (anthophyllite, tremolite, actinolite, riebeckite, and grunerite, with the last two commercially known as crocidolite and amosite, respectively). Asbestos has been widely used in the past in a number of industrial settings due to its high tensile strength, flexibility, electrical and thermal resistance, but today is banned in several countries, due to health risks posed by their inhalation (IARC, 2012). Indeed, despite an extraordinary

research effort, the mechanism through which asbestos fibres may give rise to disease is not yet completely clear. Toxicological studies show that interactions between fibrous material and biological environment are strongly dependent on both the morphology and the crystal chemistry of mineral fibres (Stanton et al., 1981; Bonneau et al., 1986; Fubini, 1993, 1996; Gilmour et al., 1997). In particular, both the presence and structural coordination of surface iron are considered key factors in the toxicity of asbestos (Fubini and Mollo, 1995; Martra et al., 2003; Favero-Longo et al., 2005; Gazzano et al., 2005). Besides,

the recent work of Andreozzi et al. (2017) showed that the amphibole asbestos (crocidolite and tremolite) fibre surface may undergo chemical modifications when in contact with fluids at physiological pH of 7.4, with consequent modification of the Fe topochemistry resulting in the modulation of the fibre reactivity. In particular, they showed that the radical production is dependent on specific surface Fe sites, occupied by either  $\text{Fe}^{2+}$  or  $\text{Fe}^{3+}$ , with unsaturated coordinative valences. Furthermore, Andreozzi et al. (2017) highlighted a different behaviour of crocidolite with respect to tremolite in both dissolution and Fe speciation kinetics. Iron-rich crocidolite exhibited the fastest dissolution leading to a pronounced transfer of bulk Fe to the surface, with consequent formation of an armoring of Fe-rich nanoparticles. Iron-poor tremolite, on the contrary, showed a slow dissolution process with partial amorphisation of external layers and absence of armoring.

Although crocidolite (i.e., fibrous riebeckite) is the most studied type of asbestos from the toxicological point of view, only very few structural data are available in the scientific literature. The first structural refinement of a fibrous riebeckite from Bolivia was done by Whittaker (1949), indeed representing the first example of a crystal structure refinement of a fibrous amphibole. According to its formula,  $(\text{Na}_{1.38}\text{K}_{0.13}\text{Ca}_{0.17}\text{Mg}_{0.25})(\text{Mg}_{2.81}\text{Fe}^{3+}_{1.66}\text{Fe}^{2+}_{0.48}\text{Al}_{0.05})[\text{Si}_{7.94}\text{Al}_{0.06}\text{O}_{22}](\text{OH})_2$ , the investigated sample may be classified as a magnesio-riebeckite. The refined structure is very similar to that later reported by Hawthorne (1978) for fluoro-riebeckite and by Hawthorne et al. (2008) for several magnesio-riebeckite samples. Nevertheless, the UICC (Union for International Cancer Control) crocidolite standard is a riebeckite from South Africa that commonly contain higher Fe contents (Hodgson, 1979; Miyano and Beukes, 1997; Pacella et al., 2014). Bloise et al. (2016) investigated the thermal behavior of UICC crocidolite from Koegas Mine using thermo-gravimetry coupled with differential scanning calorimetry. Results obtained showed that the fibers decompose at around 850 °C with the formation of magnetite, cristobalite and acmite. Interestingly, despite the thermal treatment and the following change of the amphibole structure, the sample still preserves the fibrous habit (pseudomorphosis). Quantitative data on the percentage of the actual asbestos and the mineral species present as impurities in an UICC crocidolite standard sample are reported by Kohyama et al. (1986), showing a content of >99 wt% crocidolite and <1 wt% quartz. More recently, Della Ventura et al. (2018) indicated, in addition to quartz, the presence of very minor amounts of hematite, magnetite, talc, and lizardite for the same sample. Gualtieri et al. (2013) investigated the structural and microstructural changes of an UICC crocidolite standard sample after

injecting in mice peritoneum. Results indicated a partial fibre degradation of the crystalline structure following the incubation in the biological matrix. Rietveld refinement data highlighted intra-structure cation mobilization with Fe and Na migrating from site  $M(2)$  to site  $M(1)$  and from site  $A(m)$  to site  $M(4)$ , respectively.

To clarify crocidolite structural details and define its cation distribution scheme, in the present work we carried out a full structural and spectroscopic characterization of an UICC crocidolite standard sample from Koegas Mine, Northern Cape (South Africa). Cation site partition, especially of  $\text{Fe}^{2+}$  and  $\text{Fe}^{3+}$  ionic species, was obtained from SEM-EDS chemical data integrated by  $^{57}\text{Fe}$  Mössbauer spectroscopy and X-ray powder diffraction data.

## EXPERIMENTAL

### Scanning Electron Microscopy with Energy Dispersive System (SEM-EDS)

The micro-chemical characterization was performed using a Quanta 400 SEM (FEI, Hillsboro, Oregon, USA) equipped with an EDX Genesis EDS system. Operating conditions were: 15 kV accelerating voltage, 11 mm working distance, 0° tilt angle. SEM analysis was done on the flattened part of fibre bundles (6 analytical points) in order to avoid the effects due to particle geometry on the ability to obtain quantitative EDS data (Pacella et al., 2016; Newbury and Ritchie, 2013; Paoletti et al., 2011, 2008). In addition, analyses on dispersed fibers were performed and results confirmed the homogeneity of the fibre chemical composition (e.g. Ca absence). Table 1 shows the average chemical composition and crystal-chemical formula normalized on the basis of 24 O. Cations are reported in atoms per formula unit (apfu) and were assigned following Hawthorne et al. (2012). In addition, the morphology of the fibres was investigated by Field-Emission SEM Gemini 500.

### $^{57}\text{Fe}$ Mössbauer Spectroscopy

About 10 mg of UICC crocidolite were gently ground in an agate mortar with acetone and mixed with a powdered acrylic resin to avoid (or reduce) preferred orientations. The absorber was within the limits for the thin absorber thickness described by Long et al. (1983). Data were collected at room temperature and calibrated against both the 4-lines and 6-lines  $\alpha$ -Fe spectra, using a conventional spectrometer system operated in constant acceleration mode, with a  $^{57}\text{Co}$  source of nominal strength of 1850 MBq (50 mCi) in rhodium matrix, and recorded in a multichannel analyzer using 512 channels. After velocity calibration against a spectrum of high-purity  $\alpha$ -iron foil (25  $\mu\text{m}$  thick), the raw data were folded to 256 channels. The spectrum was fitted using Recoil 1.04 fitting program

Table 1. Chemical composition of the UICC crocidolite standard sample from SEM-EDX data

Oxides	Wt%		Cations on basis of 24 (O,OH,F)	
SiO <sub>2</sub>	52.37(19)	T	Si	7.950(18)
Al <sub>2</sub> O <sub>3</sub>	0.12(18)		Al	0.022(32)
Fe <sub>2</sub> O <sub>3</sub>	17.87(32)		total	<b>7.972(14)</b>
FeO	17.42(31)	C	Fe <sup>3+</sup>	2.041(39)
MgO	3.30(35)		Fe <sup>2+</sup>	2.212(43)
CaO	-		Mg	0.747(77)
Na <sub>2</sub> O	6.88(27)		total	<b>5.000(5)</b>
H <sub>2</sub> O	2.04	B	Na	2.026(75)
Total	100.00		total	<b>2.026(75)</b>

Note: Standard deviations are reported in brackets.

(Lagarec and Rancourt, 1988). Pure Lorentian line shape were fitted, and results were satisfactory. After a number of trials, a model based on five Lorentzian doublets, with three doublets for Fe<sup>2+</sup> and two doublets for Fe<sup>3+</sup>, was chosen (Table 2). An alternative fitting strategy based on quadrupole splitting distributions was tested, and results were in agreement with those obtained using the Lorentzian approach (reduced  $\chi^2$  between 1.0 and 1.5 and reasonable parameters, in line with the specific literature). The extended spectrum (6-lines calibration) revealed the presence of minor magnetic contribution to the absorption that was fitted with two magnetic sextets. Uncertainties were calculated using the covariance matrix and the errors were estimated to be approximately  $\pm 3\%$  for both Fe<sup>2+</sup> and Fe<sup>3+</sup> absorption areas.

#### X-ray Powder Diffraction (XRPD)

X-ray powder diffraction data were collected on a Bruker AXS D8 Advance operating in  $\theta/\theta$  transmission mode. The powder was charged in a 0.5 mm diameter borosilicate glass capillary that was aligned on a

standard goniometer stage. The instrument is fitted with an incident beam focussing Göbel mirror and a PSD VÅntec-1. Measurement was performed, using CuK $\alpha$ , in the 5-145° 2 $\theta$  angular range, 0.022° 2 $\theta$  step size and 10 s counting time. Data were evaluated by the Rietveld method using Topas V4.2 (Bruker AXS, 2009) applying the Fundamental Parameters Approach FPA (Cheary and Coelho, 1992). Owing to the strong correlations with site occupancies the structure of crocidolite was refined keeping fixed all displacement parameters to reference data, following the same approach used by Andreozzi et al. (2009). No restraints on bond distances and angles were imposed. Site scattering (*s.s.*) at *M*(1), *M*(2), *M*(3) and *M*(4) was optimized. Very low electron density was detected at A-type sites in keeping with the very small cationic excess at *M*(4). For stabilizing the refinement, a single *A* site was used to model this behaviour. Careful scrutiny of the pattern revealed the occurrence of minor magnetite, calcite, quartz, siderite and minnesotaite. In particular, minnesotaite (Fe<sup>2+</sup>,Mg)<sub>3</sub>[Si<sub>4</sub>O<sub>10</sub>](OH)<sub>2</sub> is a fairly common phyllosilicate pertaining to the pyrophyllite-talc group (Guggenheim and Eggleton, 1986) that has been reported as an accessory phase at the Koegas-Westerberg area (Cilliers et al., 1961). For the accessory phases, only scale factors, cell parameters and peak shapes were refined.

Peak broadening followed a Lorentzian (size) and a Gaussian (strain) behaviour (Delhez et al., 1993). The  $\epsilon_0$  microstrain (lattice strain), defined as  $\beta_i = 4\epsilon_0 \tan \theta$ , and the volume-weighted mean column height  $L_{vol}$ , defined as  $\beta_i = l/L_{vol} \cos \theta$  were extracted from the integral breadths  $\beta_i$  of the individual reflections (Ballirano and Sadun, 2009). Subsequently, an attempt to model a size-dependent anisotropic broadening of the reflections of crocidolite was carried out using the ellipsoid-model of Katerinopoulou et al. (2012) describing the diffraction-vector dependent broadening of diffraction maxima. In the monoclinic symmetry, the shape ellipsoid parameters  $b_{ij}$  are constrained as  $b_{11} \neq b_{22} \neq b_{33}$ ;  $b_{12} = b_{23} = 0$ . An improvement

Table 2. Room temperature <sup>57</sup>Mössbauer parameters of the UICC crocidolite standard sample.

UICC crocidolite	$\delta$	$\Delta E_Q$	$\Gamma$	Assignment	Area (%)	Fe <sup>3+</sup> /Fe <sub>Tot</sub> (raw)	Fe <sup>3+</sup> /Fe <sub>Tot</sub> (corr)
	1.14	2.88	0.32	<sup>V</sup> Fe <sup>2+</sup>	25	0.53	0.48
	1.14	2.39	0.36	<sup>V</sup> Fe <sup>2+</sup>	15		
	1.08	2.23	0.76	<sup>V</sup> Fe <sup>2+</sup>	7		
	0.40	0.44	0.28	<sup>V</sup> Fe <sup>3+</sup>	23		
	0.33	0.00	1.50	<sup>V</sup> Fe <sup>3+</sup>	30		

Note: Fit with Lorentzian doublets; temperature=298 K;  $\delta$ =centre shift (with respect to an  $\alpha$ -iron foil);  $\Delta E_Q$ =quadrupole splitting;  $\Gamma$ =full width at half maximum; Area=absorption area. Errors were estimated at about  $\pm 2$  mm/s for  $\delta$ ,  $\Delta E_Q$  and  $\Gamma$ , and no less than  $\pm 3\%$  absolute for doublet areas.

of the fit was observed as well as a reduction of standard deviations. Absorption was modelled following the approach of Sabine et al. (1998) for a cylindrical sample. Preferred orientation of crocidolite was modelled by means of spherical harmonics (nine refinable parameters up to the 4<sup>th</sup> order), resulting in a minor improvement of the fit, coherently with extremely small parameters, as expected for a capillary mount.

The Quantitative Phase Analysis (QPA) results are reported in Table 3, relevant parameters of the refinement, including statistical indicators, are listed in Table 4. Rietveld plots are shown in Figure 1. QPA follows the check list indicated in Gualtieri et al. (2019). Table 5 reports the relevant bond distances of crocidolite. The

Table 3. Quantitative Phase Analysis of the UICC crocidolite standard sample.

Phases	wt%	Reference structural data
Crocidolite	93.67(19)	Hawthorne (1978)
Magnetite	1.85(6)	Wechsler et al. (1984)
Quartz	1.48(4)	Le Page and Donnay (1976)
Calcite	1.30(9)	Ballirano (2011)
Siderite	1.12(9)	Effenberger et al. (1981)
Minnesotaite	0.57(6)	Guggenheim and Eggleton (1986)

corresponding CIF file of UICC crocidolite is available for downloading at the web site of the journal.

Table 4. Microstructural parameters of the UICC crocidolite standard sample (see text) and agreement factors (as defined in Young, 1993) of the Rietveld refinement. Isotropic and anisotropic refers to refinements carried out with an isotropic/anisotropic model of peak broadening (see text for details).

	isotropic	anisotropic
$L_{vol}$ (nm)	63.9(15)	-
$b_{11}$	-	4.37(14)
$b_{22}$	-	0.56(4)
$b_{33}$	-	9.7(5)
$b_{13}$	-	3.57(19)
$\epsilon_o$	0.040(5)	0.0714(19)
$R_{Bragg}$ (%)	0.86	0.70
$R_{wp}$ (%)	2.74	2.69
$R_p$ (%)	1.66	1.66
GoF	4.41	4.32
DWd	1.49	1.56

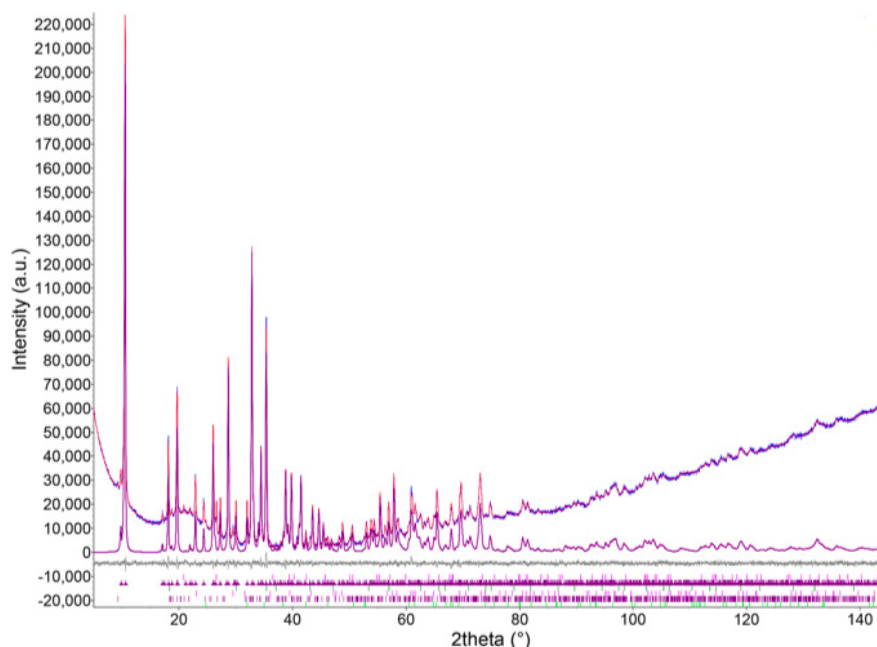


Figure 1. Rietveld plots of UICC crocidolite sample. Blue dotted line: experimental pattern; red continuous line: calculated; grey continuous line: difference plot; red continuous line calculated crocidolite contribution; vertical bars: position of the calculated Bragg reflections of, from above to below: quartz, crocidolite, magnetite, calcite, minnesotaite and siderite.

Table 5. Relevant bond distances (in Å) of the UICC crocidolite standard sample.

<i>T</i> (1)	-O(7)	1.626(5)	<i>T</i> (2)	-O(4)	1.589(8)
	-O(6)	1.629(6)		-O(5)	1.612(8)
	-O(1)	1.632(10)		-O(2)	1.640(9)
	-O(5)	1.635(9)		-O(6)	1.672(8)
< <i>T</i> (1)-O>		1.630	< <i>T</i> (2)-O>		1.628
<i>M</i> (1)	-O(3) x2	2.083(7)	<i>M</i> (2)	-O(4) x2	1.912(7)
	-O(1) x2	2.085(9)		-O(2) x2	2.066(8)
	-O(2) x2	2.126(7)		-O(1) x2	2.112(7)
< <i>M</i> (1)-O>		2.098	< <i>M</i> (2)-O>		2.030
<i>M</i> (3)	-O(1) x4	2.116(7)	<i>M</i> (4)	-O(4) x2	2.359(9)
	-O(3) x2	2.129(12)		-O(2) x2	2.364(8)
< <i>M</i> (3)-O>		2.120		-O(6) x2	2.513(8)
				-O(5) x2	2.900(8)
<i>A</i>	-O(7) x2	2.540(10)	< <i>M</i> (4)-O>		2.534
	-O(5) x4	2.887(7)			
	-O(6) x4	3.296(6)			
< <i>A</i> -O>		2.981			

## RESULTS AND DISCUSSION

The fibres of UICC crocidolite standard sample here examined appear straight, rigid and very tiny even when observed at high magnification through FE-SEM images. In addition, high-resolution images show that the micrometric fibres may split into smaller fibrils with diameter of ca. 0.1 µm (Figure 2).

The SEM-EDS analyses revealed substantial chemical homogeneity of the fibres (Table 1). Combining chemical data with Mössbauer results the following crystal-chemical formula was retrieved:  ${}^A\text{Na}_{0.026}\text{Na}_{2.000}\text{C}(\text{Fe}^{2+}_{2.212}\text{Fe}^{3+}_{2.041}\text{Mg}_{0.747})_{\Sigma=5.000}\text{T}(\text{Si}_{7.950}\text{Al}_{0.022})_{\Sigma=7.972}\text{O}_{22}\text{O}^{(3)}(\text{OH})_2$ , fairly close to the end-member riebeckite, ideally  ${}^A\text{Na}_2\text{C}(\text{Fe}^{2+}_3\text{Fe}^{3+}_2)_{\Sigma=5}\text{T}\text{Si}_8\text{O}_{22}\text{O}^{(3)}(\text{OH})_2$ .

The chemical composition of the investigated sample is closely comparable to those obtained on two UICC crocidolite standard samples coming from the same locality and previously studied by Pacella et al. (2014) [ ${}^A\text{B}(\text{K}_{0.01}\text{Na}_{1.64}\text{Ca}_{0.14}\text{Mg}_{0.16})_{\Sigma=1.95}\text{C}(\text{Fe}^{2+}_{2.13}\text{Fe}^{3+}_{2.30}\text{Mg}_{0.55}\text{Mn}_{0.01}\text{Ti}_{0.01})_{\Sigma=5.00}(\text{Si}_{7.82}\text{Al}_{0.02})_{\Sigma=7.84}\text{O}_{22}(\text{OH})_{2.1}$ ] and by Della Ventura et al. (2018) [ ${}^A(\text{K}_{0.01}\text{Na}_{0.03})_{\Sigma=0.04}\text{B}(\text{Ca}_{0.04}\text{Na}_{1.96})_{\Sigma=2.00}\text{C}(\text{Fe}^{2+}_{2.22}\text{Fe}^{3+}_{2.15}\text{Mg}_{0.52}\text{Mn}_{0.01})_{\Sigma=4.90}(\text{Si}_{7.92}\text{Al}_{0.02})_{\Sigma=7.94}\text{O}_{22}(\text{OH})_{2.2}$ ], using Mössbauer data coupled with ICP-OES and SEM/EDS analysis, respectively. In

detail, the main differences between the UICC crocidolite sample investigated in this work and the other two samples are the absence of Ca and a slightly lower  $\text{Fe}_{\text{tot}}$  content.

The room temperature  ${}^{57}\text{Fe}$  Mössbauer spectrum of the investigated UICC crocidolite sample calibrated against the 4-lines  $\alpha$ -Fe spectrum shows two main absorptions between -1 and 3 mm/s (Figure 3a). The crocidolite spectrum is well comparable with literature spectra collected on UICC crocidolite (Stroink et al., 1980) and on riebeckite (Susta et al., 2018). The presence of very weak absorption in the external part of the spectrum raised the suspect of a magnetic contribution from Fe oxides/hydroxides, possibly due to accessory phases and/or surface alteration. The crocidolite extended spectrum (calibrated against the 6-lines  $\alpha$ -Fe spectrum) confirmed the presence of a magnetic Fe oxide that was identified as magnetite, with two sextets characterized by centre shift  $\delta=0.23$  and 0.66 mm/s, magnetic field  $H=48.87$  and 46.07 Tesla, and absorption area  $\sim 3$  and 7%  $\text{Fe}_{\text{Tot}}$ , respectively (Figure 3b). The absorption contribution of the two magnetic sextets accounts for  $\sim 10\%$  of the total Fe, which allows quantifying the magnetite as  $\sim 3$  wt% of the UICC crocidolite standard sample. Apart from the magnetite contribution, the experimental Mössbauer absorption



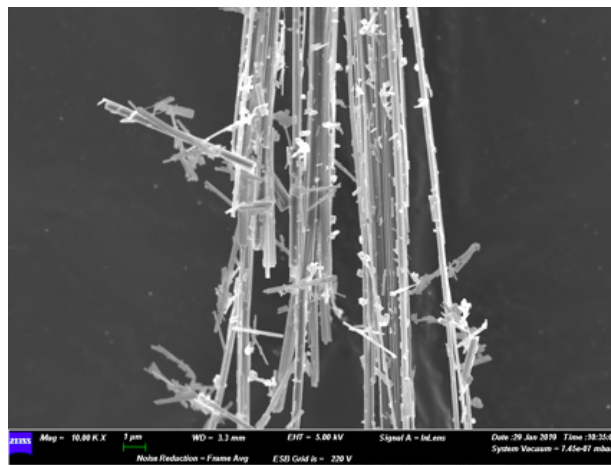
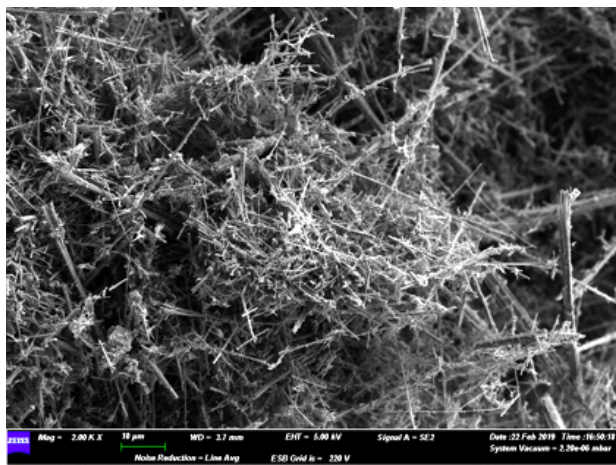


Figure 2. FE-SEM images of UICC crocidolite fibres at increasing magnification clearly showing single fibrils coming from the splitting of micrometric fibres.

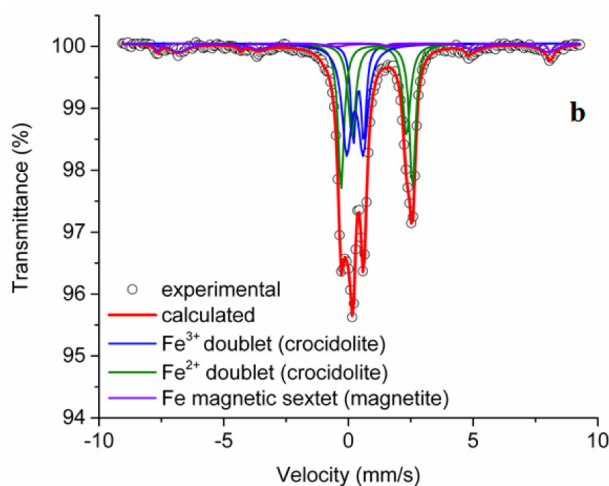
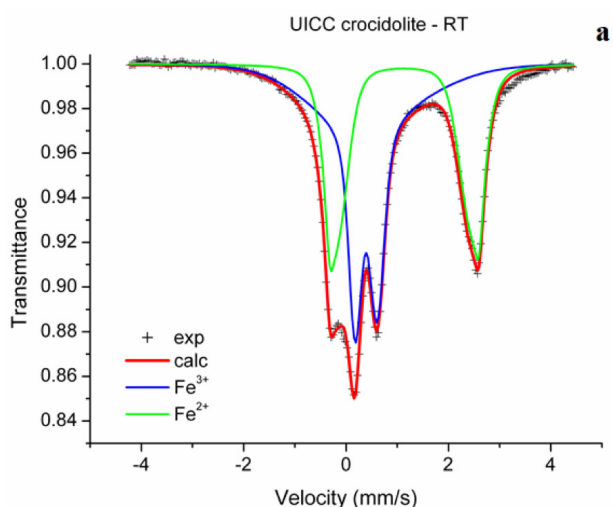


Figure 3. Room temperature  $^{57}\text{Fe}$  Mössbauer spectrum of the UICC crocidolite sample: a) spectrum calibrated against the 4-lines  $\alpha$ -Fe spectrum (fit with Quadrupole Splitting Distribution); b) extended spectrum, calibrated against the 6-lines  $\alpha$ -Fe spectrum, showing the magnetite contribution (fit with pure Lorentzian lines).

spectrum of crocidolite was reproduced by the convolution of five doublets representing  $\text{Fe}^{2+}$  and  $\text{Fe}^{3+}$  ionic species in octahedral coordination (Table 2). In particular,  $\text{Fe}^{2+}$  contribution was modelled by three doublets with almost constant centre shift ( $\delta$  values close to 1.1 mm/s) and clearly distinguished quadrupole splitting ( $\Delta E_Q$ ) values: 2.9, 2.4 and 2.2 mm/s. These data are in agreement with the existing literature and are interpreted as representing  $\text{Fe}^{2+}$  at octahedrally-coordinated sites with different next-nearest neighbour coordination environments. The contribution of  $\text{Fe}^{3+}$  was modelled by two components with comparable  $\delta$  of 0.3-0.4 mm/s and distinguished by  $\Delta E_Q$  of 0 and 0.4 mm/s. The  $\text{Fe}^{3+}/\text{Fe}_{\text{Tot}}$  ratio was quantified on the basis of spectral area measurement (and reported

in the “raw” column in Table 3), later corrected for the temperature effect (“corr” column in Table 3) applying the empirical correction factor of Dyar et al. (1993). The corrected  $\text{Fe}^{3+}/\text{Fe}_{\text{Tot}}$  ratio obtained for UICC crocidolite is 0.48, which suggests that this sample is slightly more oxidized than the end-member riebeckite (ideal ratio of 0.40).

Refined UICC crocidolite cell parameters are  $a=9.73516(19)$  Å,  $b=18.0453(3)$  Å,  $c=5.32895(9)$  Å,  $\beta=103.5159(12)^\circ$ ,  $V=910.23(3)$  Å<sup>3</sup>. The QPA indicates that 93.7(3) wt.% of the sample consists of crocidolite, the remaining being distributed among five different phases, the relatively most abundant of which is the magnetite, quantified at ~2 wt% (Table 3). Results here obtained are

only in partial agreement with the findings of Kohyama et al. (1986), who reported, for the same sample, a content of >99 wt% crocidolite and <1 wt% quartz.

Structural details of the UICC crocidolite standard sample here examined are compared with those reported by Gualtieri et al. (2013) for the same material and Susta et al. (2018) for riebeckite in Table 6. The results of the present refinement are very close to those of Susta et al. (2018) as the mean bond distances differ of <0.01 Å with the only exception of <M(1)-O> whose difference is of 0.020 Å, likely due to slightly different chemical composition of the two samples. Larger differences are observed with respect to data reported by Gualtieri et al. (2013), in particular with respect to the <M(3)-O> bond distance (that was reported to be 2.217 Å, a value anomalously large). The observed discrepancy may be attributed, as pointed out by the authors, to the significant

contribution to the diffraction pattern of the components used for sample preparation, which affected the accuracy of the structural data.

The refined site scattering (*s.s.*) at C group sites (120.2(8) e<sup>-</sup>) is in excellent agreement with that from combined chemical and Mössbauer data (119(3) e<sup>-</sup>), as reported in Table 7. Following the approach of Vignaroli et al. (2014), an indirect Fe<sup>2+</sup>/Fe<sup>3+</sup> partition was performed by comparing the <r<sup>M</sup>> mean cationic radii calculated from both the refined <M(1,2,3)-O> bond distances and from the proposed site partition. The <r<sup>M</sup>> were calculated following the conventional assignment to the C group sites (Hawthorne, 1981) and using the cation ionic radii taken from Shannon (1976) by considering an <sup>III</sup>O radius of 1.36 Å (Table 7). The result of this process produces a Fe<sup>2+</sup>/Fe<sup>3+</sup>=1.07, which is in very good agreement with 1.08(4) obtained combining chemical and Mössbauer

Table 6. Comparison of mean bond distances (in Å) of the UICC crocidolite standard sample with reference data.

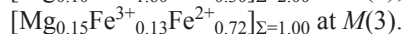
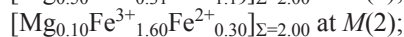
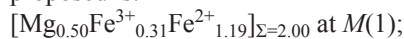
Mean bond distances	UICC crocidolite (present work)	Gualtieri et al. (2013)	Susta et al. (2018)*
<T(1)-O>	1.630	1.650	1.625
<T(2)-O>	1.628	1.631	1.632
<M(1)-O>	2.098	2.086	2.118
<M(2)-O>	2.030	2.068	2.031
<M(3)-O>	2.120	2.217	2.123
<M(4)-O>	2.534	2.498	2.541

Note: \*Data of Susta et al. (2018) refer to riebeckite.

Table 7. Crystal chemistry of the UICC crocidolite standard sample. Left side: experimental site scattering (exp. *s.s.*) at C sites from Rietveld refinement, Mg-Fe partition from *s.s.*, Fe<sup>2+</sup>-Fe<sup>3+</sup> partition from <r<sup>M</sup>>. Right side: calculated site scattering (calc. *s.s.*) from chemical formula, Fe<sup>2+</sup>-Fe<sup>3+</sup> partition from combined SEM-EDX and Mössbauer data.

Site	exp. <i>s.s.</i> (e <sup>-</sup> )	Mg-Fe partition from <i>s.s.</i>	Fe <sup>2+</sup> -Fe <sup>3+</sup> partition from <r <sup>M</sup> >	calc. <i>s.s.</i> (e <sup>-</sup> )	Fe <sup>2+</sup> -Fe <sup>3+</sup> partition from SEM and Mössbauer data
T					[Si <sub>7.95(2)</sub> Al <sub>0.02(3)</sub> ]
C					
M(1)	45.4(2)	Mg <sub>0.47</sub> Fe <sub>1.53(2)</sub>	[Mg <sub>0.47</sub> Fe <sup>3+</sup> <sub>0.41</sub> Fe <sup>2+</sup> <sub>1.12</sub> ]		[Mg <sub>0.50</sub> Fe <sup>3+</sup> <sub>0.31</sub> Fe <sup>2+</sup> <sub>1.19</sub> ]
M(2)	50.6(3)	Mg <sub>0.10</sub> Fe <sub>1.90(1)</sub>	[Mg <sub>0.10</sub> Fe <sup>3+</sup> <sub>1.58</sub> Fe <sup>2+</sup> <sub>0.32</sub> ]		[Mg <sub>0.10</sub> Fe <sup>3+</sup> <sub>1.60</sub> Fe <sup>2+</sup> <sub>0.30</sub> ]
M(3)	24.2(3)	Mg <sub>0.13</sub> Fe <sub>0.87(1)</sub>	[Mg <sub>0.13</sub> Fe <sup>3+</sup> <sub>0.09</sub> Fe <sup>2+</sup> <sub>0.78</sub> ]		[Mg <sub>0.13</sub> Fe <sup>3+</sup> <sub>0.13</sub> Fe <sup>2+</sup> <sub>0.72</sub> ]
<b>S<sub>M(1)+M(2)+M(3)</sub></b>	<b>120.2(8)</b>	<b>Mg<sub>0.70</sub>Fe<sub>4.30(4)</sub></b>	<b>[Mg<sub>0.70</sub>Fe<sup>3+</sup><sub>2.08</sub>Fe<sup>2+</sup><sub>2.22</sub>]</b>	<b>119(3)</b>	<b>[Mg<sub>0.75(8)</sub>Fe<sup>3+</sup><sub>2.04(4)</sub>Fe<sup>2+</sup><sub>2.21(4)</sub>]</b>
Fe <sup>2+</sup> /Fe <sup>3+</sup>			1.07		1.08(4)
B					
M(4)	20.8(2)	Na <sub>1.89(1)</sub>	Na <sub>1.89</sub>		Na <sub>2.00(7)</sub>
Am	0.6(3)	Na <sub>0.05(2)</sub>	Na <sub>0.05(2)</sub>		Na <sub>0.03(7)</sub>

data. According to this procedure, the UICC crocidolite sample allocates  $\text{Fe}^{3+}$  with the following site-preference sequence  $M(2) \gg M(1) > M(3)$  and  $\text{Fe}^{2+}$  with the sequence  $M(3) > M(1) \gg M(2)$ . As a result, the final partition here proposed is:



Comparison between measured and calculated  $\langle r^M \rangle$  mean cationic radii shows an excellent agreement, with discrepancies ranging within one standard deviation of  $M$ -O bond distances.

## CONCLUSIONS

This work reports an accurate crystal-chemical characterization of the UICC crocidolite standard sample coming from Koegas Mine, Northern Cape (South Africa) obtained using a multi-analytical chemical, spectroscopic and diffractometric approach. The chemical formula and the cation distribution, obtained on the basis of a combination of all data, showed that the investigated sample is very close to the riebeckite end member.

Although the UICC crocidolite standard reference has been extensively used for toxicological investigations, an accurate, full crystal structural characterization of the fibres was still missing. Our QPA revealed the presence of about 7 wt% of accessory phases, much more than those observed by Kohyama et al. (1986) and Della Ventura et al. (2018), highlighting the occurrence of non-negligible contents of impurities in the UICC crocidolite standard samples. Notably, the presence of quartz, a mineral of well known toxic potential, and Fe-bearing phases such as magnetite and minor siderite and minnesotaite, is a relevant aspect to be taken in due consideration in case of reactivity and toxicity studies on UICC crocidolite fibres.

## ACKNOWLEDGEMENTS

Francesco Turci of the Department of Chemistry, University of Turin, Italy, is thanked for providing the crocidolite standard sample. Many thanks to Marco Albano for technical assistance with the SEM-EDS analyses at CNR-IGAG, UOS of Rome, Italy, and Lorenzo Arrizza of the Microscopy Centre, University of L'Aquila, Italy, for fibre FE-SEM images.

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