

Ralphcannonite, $\text{AgZn}_2\text{TlAs}_2\text{S}_6$, a new mineral of the routhierite isotypic series from Lengenbach, Binn Valley, Switzerland

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ABSTRACT

The new mineral species ralphcannonite, $\text{AgZn}_2\text{TlAs}_2\text{S}_6$, was discovered in the Lengenbach quarry, Binn Valley, Wallis, Switzerland. It occurs as metallic black equant, isometric to prismatic crystals, up to 50 μm , associated with dufrénoysite, hatchite, realgar and baryte. Minimum and maximum reflectance data for COM wavelengths in air are [λ (nm): R (%)]: 471.1: 25.8/27.1; 548.3: 25.2/26.6; 586.6: 24.6/25.8; 652.3: 23.9/24.8. Electron microprobe analyses give (wt.%): Cu 2.01(6), Ag 8.50(16), Zn 10.94(20), Fe 3.25(8), Hg 7.92(12), Tl 24.58(26), As 18.36(19), Sb 0.17(4), S 24.03(21), total 99.76(71). On the basis of 12 atoms per formula unit, the chemical formula of ralphcannonite is $\text{Ag}_{0.63(2)}\text{Cu}_{0.25(2)}\text{Zn}_{1.35(5)}\text{Fe}_{0.47(1)}\text{Hg}_{0.32(2)}\text{Tl}_{0.97(3)}[\text{As}_{1.97(6)}\text{Sb}_{0.01(1)}]_{\Sigma 1.98(5)}\text{S}_{6.03(8)}$. The new mineral is tetragonal, space group $I42m$, with $a = 9.861(2)$, $c = 11.125(3)$ Å, $V = 1081.8(4)$ Å³, $Z = 4$. The main diffraction lines of the calculated powder diagram are [d (in Å), intensity, hkl]: 4.100, 85, 211; 3.471, 40, 103; 2.954, 100, 222; 2.465, 24, 400; 2.460, 39, 303. The crystal structure of ralphcannonite has been refined by X-ray single-crystal data to a final $R_1 = 0.030$, on the basis of 140 observed reflections [$F_o > 4\sigma(F_o)$]. It shows a three dimensional framework of (Ag,Zn)-centred tetrahedra (1 $M1$ + 2 $M2$), with channels parallel to $[001]$ hosting TlS_6 and (As,Sb) S_3 disymmetric polyhedra. Ralphcannonite is derived from its isotype routhierite $^{M1}\text{Cu}^{M2}\text{Hg}_2\text{TlAs}_2\text{S}_6$ through the double heterovalent substitution $^{M1}\text{Cu}^+ + ^{M2}\text{Hg}^{2+} \rightarrow ^{M1}\text{Zn}^{2+} + ^{M2}\text{Ag}^+$. This substitution obeys a steric constraint, with Ag^+ , the largest cation relative to Zn^{2+} and Cu^+ , entering the largest $M2$ site, as observed in arsiccioite. The ideal crystal-chemical formula of ralphcannonite is $^{M1}\text{Zn}^{M2}(\text{Zn}_{0.5}\text{Ag}_{0.5})_2\text{TlAs}_2\text{S}_6$.

KEYWORDS: ralphcannonite, new mineral species, sulfosalt, thallium, zinc, silver, arsenic, crystal structure, Lengenbach, Binn Valley, Switzerland.

Introduction

AMONG the world-class localities for the study of sulfosalt assemblages, Lengenbach occupies an outstanding position, owing to the large number of sulfosalt species (more than 30 species as of

November 2014) with it as their type locality and to the complex and intriguing geochemistry, with an original combination of numerous cations (Pb, Tl, Ag, Cu, Zn, Cd, Sn, As and Sb). In particular, Lengenbach can be considered the most important locality world-wide for the study of the crystal-chemistry of thallium sulfosalts. Indeed, 19 thallium sulfosalts have been identified from the Lengenbach dolostone so far; among these, 15 were first described from this locality (Table 1).

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Following the sulfosalt systematics given by Mořlo *et al.* (2008), thallium sulfosalts from Lengenbach can be classified in the following groups (the very recently approved mineral spaltiite is not considered because its crystallographic data have not been published yet):

- (1) sulfosalts with an atom ratio of cation/chalcogen = 1: the hatchite isotypes (hatchite, wallisite) and the two members of the weissbergite homeotypic pair, i.e. weissbergite and lorándite;
- (2) lead sulfosalts based on large 2D fragments of the PbS/SnS archetype, represented by members of the sartorite homologous series (philrothite) and related compounds (dalnegroite, parapierrroite);
- (3) specific Tl sulfosalts (structures with SnS layers), particularly represented by members of the hutchinsonite merotypic series (bernardite, hutchinsonite, edenharterite, jentschite, imhofite). Other minerals belonging to this type are sicherite, gabrielite and erniggllite. Possibly, the recently described mineral raberite could be classified within this group;
- (4) sulfosalts with an excess of small monovalent cations (Ag,Cu) relative to (As, Sb, Bi), represented by the member of the routhierite isotypic series stalderite.

This latter mineral, as well as the members of the routhierite isotypic series, deserve a careful study, owing to their variable crystal chemistry, possibly reflecting the geochemistry of the crystallizing medium, as described for the routhierite-arsiccioite pair by Biagioni *et al.* (2014b). Preliminary energy dispersive spectrometry (EDS) chemical analyses performed on specimens of stalderite showed the occurrence of Ag- and Fe-rich varieties. In particular, a specimen examined in 2003 proved to be an Ag-rich Fe-bearing stalderite. In 2013, new EDS chemical analyses confirmed its high Ag content, suggesting the need for a more accurate characterization.

Crystallographic studies and chemical analyses allowed us to describe this phase as the new mineral ralphcannonite. The mineral and its name have been approved by the Commission on New Minerals, Nomenclature and Classification of the International Mineralogical Association (IMA 2014-077, Bindi *et al.*, 2015). The holotype specimen of ralphcannonite is deposited in the mineralogical collection of the Museo di Storia Naturale, Università degli Studi di Firenze, Via La Pira 4, Florence, Italy, under catalog number 3145/I. The name honours Ralph Cannon (b. 1956) for his contribution to the knowledge of the mineralogy of the Lengenbach quarry. Ralph Cannon

TABLE 1. Thallium sulfosalts from Lengenbach. Sulfosalts that have their type locality at Lengenbach are in bold.

Mineral	Chemical formula	References
Bernardite	TlAs ₅ S ₈	Hofmann <i>et al.</i> (1993)
Dalnegroite	(Tl ₂ PbSb)(Sb ₃ As ₆)S ₁₇	Nestola <i>et al.</i> (2009)
Edenharterite	TlPbAs ₃ S ₆	Graeser and Schwander (1992)
Erniggllite	SnTl ₂ As ₂ S ₆	Graeser <i>et al.</i> (1992)
Gabrielite	Cu ₂ AgTl ₂ As ₃ S ₇	Graeser <i>et al.</i> (2007)
Hatchite	AgTlPbAs ₂ S ₅	Solly and Smith (1912)
Hutchinsonite	TlPbAs ₅ S ₉	Solly (1905)
Imhofite	Tl _{5,8} As _{15,4} S ₂₆	Burri <i>et al.</i> (1965)
Jentschite	TlPbAs ₂ SbS ₆	Graeser and Edenharter (1997)
Lorándite	TlAsS ₂	Graeser (1967)
Parapierrroite	TlSb ₅ S ₈	Nestola (unp. data, 2014)
Philrothite	TlAs ₃ S ₅	Bindi <i>et al.</i> (2014)
Raberite	Ag ₄ Tl ₅ As ₆ SbS ₁₅	Bindi <i>et al.</i> (2012)
Ralphcannonite	AgZn ₂ TlAs ₂ S ₆	This work
Sicherite	Ag ₂ Tl(As,Sb) ₃ S ₆	Graeser <i>et al.</i> (2001)
Spaltiite	Cu ₂ Tl ₂ As ₂ S ₅	Graeser <i>et al.</i> (2014)
Stalderite	CuZn ₂ TlAs ₂ S ₆	Graeser <i>et al.</i> (1995)
Wallisite	CuTlPbAs ₂ S ₅	Nowacki (1965)
Weissbergite	TlSbS ₂	Roth <i>et al.</i> (2014)

started working in the Lengenbach quarry in 1996. In 2003, when the Lengenbach Research Association (FGL, *Forschungsgemeinschaft Lengenbach*) was founded, he was appointed as Technical Head for specimen extraction, an activity he has been carrying out ever since with great dedication and intuition. He has also analysed several specimens from this locality and has published many papers about Lengenbach mineralogy.

The aim of this paper is the description of the new mineral species ralphcannonite and its relationships with the other members of the routhierite isotypic series.

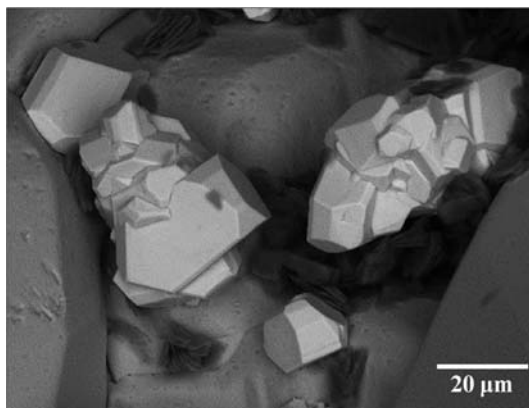


FIG. 1. Ralphcannonite, equant crystals on realgar.

Occurrence and mineral description

The Lengenbach quarry (latitude $46^{\circ}21'54''\text{N}$, longitude $8^{\circ}13'15''\text{E}$) exploits a Triassic dolostone overlying the gneiss basement at the northern front of the Monte Leone Nappe, in the Penninic Domain of the Alps. These rocks have been metamorphosed up to upper greenschist–lower amphibolite facies conditions. The dolostone is 240 m thick at Lengenbach; mineralization occurs in the uppermost part of the sequence, close to the contact with the overlying Jurassic to Lower Cretaceous *Bündnerschiefer*. Hofmann and Knihl (1996) recognized four major types of mineralization: (1) stratiform layers of pyrite with minor galena, sphalerite and xenomorphic sulfosalts; (2) massive to interstitial sulfosalt accumulations; (3) discordant sulfosalt and sulfide veins; and (4) idiomorphic crystals within druses and open fissures. Ralphcannonite was discovered in the type (2) mineralization, corresponding to the so-called ‘Zone 1’ of Graeser *et al.* (2008). The specimen studied was originally found in 1989 by the *Arbeitsgemeinschaft Lengenbach* (Lengenbach Working Association). Then, via the Natural History Museum in Bern, the specimen found its way into the private collection of one of the authors (TR).

Physical and optical properties

Ralphcannonite was observed as euhedral crystals; on the basis of scanning electron microscopy images (Fig. 1) and the crystal forms reported by Graeser *et al.* (1995) for stalderite, the dominant forms could be the prism $\{110\}$ and the basal pynacoid $\{001\}$, with the bipyramids $\{101\}$ and $\{111\}$, as well as the prism $\{100\}$ as accessory forms. Ralphcannonite is brittle, with irregular fracture; the streak is black, the

lustre is metallic. Micro-indentation measurements carried out with a VHN load of 30 g give a mean value of 120 kg mm^{-2} (range: 116–128; 3 measurements) corresponding to a Mohs hardness of $\sim 2\text{--}2\frac{1}{2}$.

In plane-polarized incident light, ralphcannonite is greyish in colour. Under crossed polars, it is very weakly anisotropic, with greyish to light blue rotation tints. Internal reflections are very weak. There is no optical evidence of growth zonation.

Reflectance measurements were performed in air by means of a MPM-200 Zeiss microphotometer equipped with a MSP-20 system processor on a Zeiss Axioplan ore microscope. The filament temperature was $\sim 3350 \text{ K}$. An interference filter was adjusted, in turn, to select four wavelengths (approximating those recommended by the Commission on Ore Mineralogy of the IMA) for measurement (471.1, 548.3, 586.6 and 652.3 nm). Readings were taken for specimen and standard (SiC) maintained under the same focus conditions. The diameter of the circular measuring area was 0.1 mm. Reflectance percentages for $R_{\min}$ and $R_{\max}$ are 25.8, 27.1 (471.1 nm), 25.2, 26.6 (548.3 nm), 24.6, 25.8 (586.6 nm) and 23.9, 24.8 (652.3 nm), respectively.

Owing to the small amount of pure material available, the density was not measured; the calculated density, based on the empirical formula (see below), is 4.927 g cm^{-3} .

In the specimen studied, ralphcannonite is associated with hatchite, dufrénoysite, realgar and baryte.

Chemical analysis

A preliminary chemical analysis using EDS performed on the crystal fragment used for the

TABLE 2. Microprobe analyses of ralphcannonite: chemical composition as wt.% and number of atoms on the basis of 12 atoms per formula unit (a.p.f.u.).

Element	wt.%	range	e.s.d.
Cu	2.01	1.78–2.22	0.06
Ag	8.50	8.19–8.80	0.16
Zn	10.94	10.25–11.33	0.20
Fe	3.25	3.10–3.34	0.08
Hg	7.92	7.10–8.50	0.12
Tl	24.58	23.87–25.26	0.26
As	18.36	17.58–18.96	0.19
Sb	0.17	0.09–0.25	0.04
S	24.03	23.61–24.66	0.21
Total	99.76	99.25–100.88	0.71
	a.p.f.u.	range	e.s.d.
Cu	0.255	0.226–0.279	0.022
Ag	0.634	0.606–0.663	0.021
Zn	1.346	1.262–1.400	0.054
Fe	0.468	0.449–0.482	0.013
Hg	0.318	0.286–0.344	0.022
Tl	0.968	0.940–1.005	0.028
As	1.972	1.908–2.037	0.057
Sb	0.011	0.006–0.016	0.004
S	6.030	5.949–6.141	0.078
Ev	0.1	–3.7–3.0	2.5

Valence equilibrium: Ev (%) = $[\Sigma(\text{val}+) - \Sigma(\text{val}-)] \times 100 / \Sigma(\text{val}-)$.

single-crystal X-ray diffraction did not indicate the presence of elements ($Z > 9$) other than Fe, Cu, Zn, As, Ag, Hg, Tl and S.

The same fragment was then analysed with a JEOL 8200 electron microprobe (wavelength dispersive mode, accelerating voltage 15 kV, beam current 20 nA, beam size 1 μm). Counting times are 20 s for peak and 15 s for background. The following standards (element, emission line) were used: Ag metal (Ag $L\alpha$), Cu metal (Cu $K\alpha$), Zn metal (Zn $K\alpha$), Fe metal (Fe $K\alpha$), cinnabar (Hg $M\alpha$), TlAsS₂ (Tl $M\alpha$, S $K\alpha$), As metal (As $L\alpha$) and Sb metal (Sb $L\alpha$). Cadmium and Se were sought but were below the detection limit. The crystal fragment was found to be homogeneous within analytical error. Chemical data are given in Table 2.

On the basis of 12 atoms per formula unit (a.p.f.u.), the chemical formula of ralphcannonite corresponds to $\text{Ag}_{0.63(2)}\text{Cu}_{0.25(2)}\text{Zn}_{1.35(5)}\text{Fe}_{0.47(1)}\text{Hg}_{0.32(2)}\text{Tl}_{0.97(3)}[\text{As}_{1.97(6)}\text{Sb}_{0.01(1)}]\Sigma_{1.98(5)}\text{S}_{6.03(8)}$.

Crystallography

For the X-ray single-crystal study, the intensity data were collected using an Oxford Diffraction Xcalibur 3 diffractometer equipped with a Sapphire 2 CCD area detector, with Mo $K\alpha$ radiation. The detector-to-crystal distance was 60 mm and 288 frames were collected using ω scan mode, in 0.5° slices, with an exposure time of 45 s per frame. Intensity integration and standard Lorentz-polarization corrections were performed with the *CrysAlis RED* (Oxford Diffraction, 2006) software package. The program *ABSPACK* in *CrysAlis RED* (Oxford Diffraction, 2006) was used for the absorption correction. The statistical tests on the distribution of $|E|$ values ($|E^2 - 1| = 0.808$) and the systematic absences suggested the space group $I42m$. The refined cell parameters are $a = 9.861(2)$, $c = 11.125(3)$ Å, $V = 1081.8(4)$ Å³.

The crystal structure of ralphcannonite was refined starting from the atomic coordinates given by Bindi (2008) for routhierite, using *Shelxl-97* (Sheldrick, 2008). Scattering curves for neutral atoms were taken from the *International Tables for Crystallography* (Wilson, 1992). Crystal data and details of intensity data collection and refinement are reported in Table 3. Owing to the relatively small number of observed reflections, only an isotropic model was refined for ralphcannonite. After several cycles of refinement, the R_1 converged to 0.030, confirming the validity of the structural model. We noticed that the goodness-of-fit of the structure refinement is very low (i.e. 0.289), a feature probably related to the low-quality diffraction data set.

Four independent cation sites occur in the crystal structure of ralphcannonite, labelled as Tl, M1, M2 and As. The M1 and M2 sites correspond to Cu and Hg sites in the crystal structure of routhierite (Bindi, 2008; Biagioni *et al.*, 2014a). The occupancies of these four independent cation sites were refined using the following scattering curves: Tl site: Tl vs. $\square$; M1 site: Cu vs. $\square$; M2 site: Hg vs. $\square$; As site: As vs. $\square$. The Tl, M1 and As sites were found to be fully occupied and their site occupancies were fixed to 1. At the M2 site, the refined site-scattering value was 43.9 electrons, compatible with a mixed (Ag,Zn,Fe,Hg) occupancy. Final atom coordinates and isotropic displacement parameters are given in Table 4, whereas selected bond distances are reported in Table 5. The refined and calculated site scattering (in electrons per formula unit, e.p.f.u.) on the basis of the proposed site populations are given in Table 6. Iron has been considered as divalent, in

TABLE 3. Crystal and experimental data for ralphcannonite.

Crystal data	
Crystal size (mm^3)	0.035 x 0.045 x 0.055
Cell setting, space group	Tetragonal, $I42m$
a , c ($\text{\AA}$)	9.861(2), 11.125(3)
V ($\text{\AA}^3$)	1081.8(4)
Z	4
Data collection and refinement	
Radiation, wavelength ($\text{\AA}$)	$\text{MoK}\alpha$, $\lambda = 0.71073$
Temperature (K)	293
$2\theta_{\text{max}}$ ($^\circ$)	56.05
Measured reflections	7722
Unique reflections	633
Reflections with $F_o > 4\sigma(F_o)$	140
R_{int}	0.0408
$R\sigma$	0.0498
Range of h , k , l	$-7 \leq h \leq 8$, $0 \leq k \leq 12$, $0 \leq l \leq 13$
$R [F_o > 4\sigma(F_o)]$	0.0299
R (all data)	0.0420
wR (on F^2)	0.1048
Number of least-squares parameters	20
Maximum and minimum residual peak ($\text{e \AA}^{-3}$)	0.93 (at 1.94 $\text{\AA}$ from S2); -1.86 (at 1.19 $\text{\AA}$ from T1)

order to achieve electrostatic neutrality; this is consistent with the observation of Makovicky *et al.* (1990) that the presence of Zn and Hg favours Fe^{2+} with respect to Fe^{3+} . Finally, Table 7 shows the results of the bond-valence calculations.

Owing to the very small size and amount of crystals available, a powder X-ray diffraction pattern was not collected. Table 8 reports the calculated powder X-ray diffraction data of ralphcannonite on the basis of the crystallographic data collected through single-crystal X-ray diffraction.

Crystal structure description

The crystal structure of ralphcannonite is isotypic with those of routhierite, stalderite and arsiccioite (Graeser *et al.*, 1995; Bindi, 2008; Biagioni *et al.*, 2014*a,b*), showing a framework formed by two independent MeS_4 tetrahedra sharing corners, hosting channels parallel to $[001]$. These channels contain TlS_6 and $(\text{As,Sb})\text{S}_3$ polyhedra, sharing corners and edges with the tetrahedron framework (Fig. 2).

TABLE 4. Atomic coordinates and isotropic displacement parameters ($\text{\AA}^2$) for ralphcannonite.

Site	Wyckoff site	x/a	y/b	z/c	U_{iso}
T1	$4e$	0	0	0.3524(2)	0.0413(7)
M1	$4d$	0	$\frac{1}{2}$	$\frac{1}{4}$	0.0334(14)
M2	$8f$	0.2228(2)	$\frac{1}{2}$	$\frac{1}{2}$	0.0405(11)
As	$8i$	0.2611(4)	0.2611(4)	0.2587(6)	0.0400(12)
S1	$16j$	0.0943(6)	0.3308(7)	0.3743(7)	0.037(2)
S2	$8i$	0.1257(7)	0.1257(7)	0.1295(9)	0.040(3)

TABLE 5. Selected bond distances (in Å) in ralphcannonite.

Tl	–S2	$3.037(10) \times 2$	M1	–S1	$2.358(7) \times 4$
	–S1	$3.401(7) \times 4$			
	average	3.280			
As	–S1	$2.198(8) \times 2$	M2	–S1	$2.417(6) \times 2$
	–S2	$2.374(11)$		–S2	$2.519(7) \times 2$
	average	2.257		average	2.468

As stated above, four independent cation sites occur. The two independent tetrahedral sites, *M1* and *M2*, have average bond distances of 2.358 and 2.468 Å, respectively. The $\langle M1-S \rangle$ bond distance agrees closely with that observed by Graeser *et al.* (1995) for the Cu site of staldelite, i.e. 2.357(4) Å, whereas the *M2* site of ralphcannonite is enlarged with respect to the corresponding Zn site of staldelite, having an average bond distance of 2.406 Å. This is in agreement with the replacement of the small Zn (and possibly Cu) atoms by the larger Ag at the *M2* site.

The As site forms a trigonal pyramid with three S atoms, with an average bond distance of 2.257 Å. The refinement of the site occupancy points to the full occupancy of this site by arsenic, in agreement with chemical data showing only a negligible amount of Sb (0.01 a.p.f.u.). Actually the AsS_3 geometry is distorted, with two relatively short As–S1 bond distances (2.198 Å) and a longer one (2.378 Å).

The coordination of the Tl site can be described as an orthorhombic pyramid with a split apex, as in the other members of the routhierite isotypic series. On the other site of the pyramidal split apex, a relatively short Tl–Tl distance occurs, i.e. 3.284(4) Å. This value agrees perfectly with that reported for staldelite (3.28 Å) by Graeser *et al.* (1995) and is in the range of Tl–Tl distances observed in other members of the routhierite group, i.e. 3.33 Å and 3.47 Å for routhierite from Monte Arsiccio and Jas Roux, respectively (Bindi, 2008; Biagioni *et al.*, 2014a) and values ranging between 3.25 and 3.51 Å for the split Tl site in arsiccioite (Biagioni *et al.*, 2014b). As reported by previous authors, such short Tl–Tl distances most likely indicate some type of Tl–Tl interaction.

 TABLE 6. Refined site-scattering values (e.p.f.u.), assigned site population (a.p.f.u.) and comparison between observed and calculated bond distances (in Å) at *M1*, *M2* and As sites in ralphcannonite.

Site	XRD	EPMA	Site population	$\langle Me-S \rangle_{obs}$	$\langle Me-S \rangle_{calc}$
<i>M1</i>	29.0	29.8	$[Zn_{0.751}Cu_{0.249}]^*$	2.358	2.352
<i>M2</i>	43.8	42.4	$Ag_{0.317}Zn_{0.290}Fe_{0.234}Hg_{0.159}$	2.468	2.499
As	33.0	33.1	$[As_{0.995}Sb_{0.005}]^*$	2.257	2.261

XRD – X-ray diffraction; EPMA – electron probe microanalysis.

*Values normalized to 1

TABLE 7. Bond-valence sums (BVS, in valence units) calculated using the parameters given by Brese and O'Keeffe (1991).

	<i>M1</i>	<i>M2</i>	Tl	As	Σ anions
S1	0.43^{x4}	0.51^{x2}	0.12^{x4}	1.17^{x2}	2.23
S2		$^{2x}0.39^{x2}$	0.34^{x2}	0.74	1.86
Σ cations	1.72	1.80	1.16	3.08	
Theoretical	1.75	1.68	1.00	3.00	

Note: In mixed sites, the bond-valence contribution of each cation has been weighted according to its occupancy (see Table 6). Left and right superscripts indicate the number of bonds involving cations and anions, respectively.

TABLE 8. Calculated powder X-ray diffraction data for ralphcannonite. Intensity and d_{hkl} were calculated using the software *Powdercell 2.3* (Kraus and Nolze, 1996) on the basis of the structural model given in Table 4; only reflections with $I_{\text{calc}} > 5$ are listed. The five strongest reflections are given in bold.

I_{calc}	d_{calc}	$h\ k\ l$	I_{calc}	d_{calc}	$h\ k\ l$
8	6.973	1 1 0	5	2.324	3 3 0
9	4.930	2 0 0	8	2.201	3 2 3
85	4.100	2 1 1	14	2.076	3 1 4
12	3.486	2 2 0	10	2.010	4 1 3
40	3.471	1 0 3	10	1.845	4 0 4
17	3.118	3 1 0	15	1.807	5 2 1
100	2.954	2 2 2	13	1.743	4 4 0
20	2.656	3 2 1	9	1.741	4 3 3
17	2.583	1 1 4	9	1.728	4 2 4
24	2.465	4 0 0	6	1.604	6 1 1
39	2.460	3 0 3	6	1.594	3 1 6
11	2.422	2 0 4	7	1.588	5 1 4
6	2.338	4 1 1	11	1.501	6 2 2

Bond-valence balance

Bond-valence sums (BVS), according to Brese and O'Keeffe (1991), are reported in Table 7, on the basis of the site population given in Table 6.

With respect to the BVS of routhierite and arsiccioite, where differences up to 25% between observed and calculated BVS have been reported (Biagioni *et al.*, 2014a,b), only minor deviations from the expected values (up to 7% for the $M2$ site) have been observed, confirming the validity of the proposed site population. A good agreement between observed and calculated BVS also occurs for the As site. The BVS excess for the Tl atom is similar to that reported in arsiccioite and routhierite (Biagioni *et al.*, 2014a,b) and can be related to the overestimation of the bond parameter tabulated by Brese and O'Keeffe (1991) for the pair (Tl,S), i.e. 2.63 Å, as suggested by Biagioni *et al.* (2014b). Using a value of $R_{\text{Tl,S}} = 2.55$ Å, a BVS of 0.95 valence units (v.u.) can be calculated for the Tl site.

Crystal chemistry

Structural formula

The crystal-chemical formula of ralphcannonite can be written as $^{M1}(\text{Zn}_{0.75}\text{Cu}_{0.25})^{M2}(\text{Ag}_{0.32}\text{Zn}_{0.29}\text{Fe}_{0.23}\text{Hg}_{0.16})_2\text{TlAs}_2\text{S}_6$, with the relative error of the valence equilibrium $E_v = +0.9$.

Graeser *et al.* (1995) proposed that in staldierite the small $M1$ site is occupied by Cu^+ , whereas the divalent cations (Zn^{2+} and minor Fe^{2+} and Hg^{2+}) are hosted at the larger $M2$ site. On the contrary, the results of the crystal structure study suggest that $M1$ is preferentially occupied by Zn^{2+} , whereas $M2$ hosts the larger cations Ag^+ and Hg^{2+} , replaced by Zn^{2+} and Fe^{2+} . As reported by Biagioni *et al.* (2014b) for arsiccioite, the cation distribution among the $M1$ and $M2$ sites is guided by the steric constraint that prevails over the valence state. Therefore, the largest cation, Ag^+ is hosted at the large $M2$ site, whereas Zn^{2+} , the smallest cation, preferentially occupies the $M1$ site.

Similarly to the ideal formula of arsiccioite, $^{M1}\text{Hg}^{M2}(\text{Ag}_{0.5}\text{Hg}_{0.5})_2\text{TlAs}_2\text{S}_6$, the formula of ralphcannonite could be written as $^{M1}\text{Zn}^{M2}(\text{Ag}_{0.5}\text{Zn}_{0.5})_2\text{TlAs}_2\text{S}_6$. The Ag content of ralphcannonite is actually a little lower than the ideal one, and its deficit is probably balanced through the heterovalent substitution $^{M1}\text{Zn}^{2+} + ^{M2}\text{Ag}^+ \rightarrow ^{M1}\text{Cu}^+ + ^{M2}(\text{Zn,Fe,Hg})^{2+}$.

Nomenclature of the routhierite isotypic series

The general crystal-chemical formula of the members of the routhierite isotypic series can be written as $(\text{Me}_{1-2x}^{+}\text{Me}_{2x}^{2+})(\text{Me}_{1-x}^{2+}\text{Me}_x^{+})_2\text{Tl}(\text{As}_{1-y}\text{Sb}_y)_2\text{S}_6$, with $0 \leq x \leq 0.5$ and $y < 0.5$. The cation Me^+ is represented by Cu^+ and Ag^+ , whereas bivalent cations Me^{2+} are Zn^{2+} , Hg^{2+} and Fe^{2+} . Minor Cd^{2+} (up to 0.02 a.p.f.u.) has been reported in arsiccioite (Biagioni *et al.*, 2014b).

The crystal-structure studies of arsiccioite and ralphcannonite indicate that Ag prefers the largest $M2$ site; this observation can also be extended to Ag-bearing routhierite (e.g. specimen from Monte Arsiccio – Biagioni *et al.*, 2014a,b). On the contrary, the smaller cations (Zn^{2+} , Cu^+) are preferentially hosted at the $M1$ site.

Graeser *et al.* (1995), in their study of staldierite, proposed the site population $M1 = \text{Cu}_{1.00}$ and $M2 = \text{Zn}_{0.55}\text{Fe}_{0.25}\text{Hg}_{0.20}$; however, as pointed out by Biagioni *et al.* (2014b), a site population with $M1 = \text{Zn}_{1.00}$ and $M2 = \text{Cu}_{0.50}\text{Fe}_{0.25}\text{Hg}_{0.20}\text{Zn}_{0.05}$ cannot be excluded, resulting in very similar site-scattering values, BVS and calculated average bond distances. Consequently, one could also verify a different cation distribution for ralphcannonite, with Zn completely hosted at the $M1$ site and a mixed (Ag,Cu,Fe,Hg,Zn) occupancy at the $M2$ site. In this case, the idealized site population should be $M1 = \text{Zn}_{1.00}$ and $M2 = \text{Ag}_{0.32}\text{Cu}_{0.13}\text{Fe}_{0.23}\text{Hg}_{0.16}\text{Zn}_{0.16}$. Assuming such a

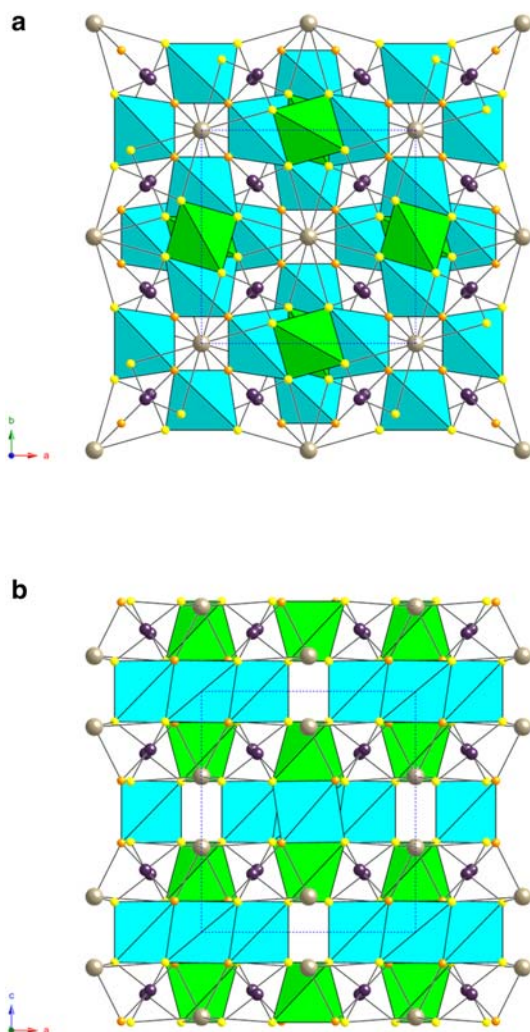


FIG. 2. Crystal structure of ralphcannonite, as seen down [001] (a) and [010] (b). Polyhedra: green: $M1$ tetrahedra; light blue: $M2$ tetrahedra. Circles: grey: Tl site; violet: As site; yellow: S1 site; orange: S2 site.

distribution, the BVS, average bond distances and calculated site scattering are similar to those proposed in Table 6, suggesting that the actual site occupancy, by using conventional X-ray diffraction, cannot be known, owing to the similar values of the scattering factors for Cu, Zn and also Fe. It is noteworthy that the site population proposed in Table 6 would correspond to the ideal formula $\text{Zn}(\text{Ag}_{0.5}\text{Zn}_{0.5})_2\text{TlAs}_2\text{S}_6$, whereas in the alternative cation distribution the formula should be written as $\text{Zn}(\text{Ag}_{0.5}\text{Fe}_{0.5})_2\text{TlAs}_2\text{S}_6$, with Fe^{2+} being the dominant bivalent cation at $M2$.

Consequently, the same chemistry could correspond to two different crystal-chemical formulae. Taking into account this issue, in our opinion, the classification of the members of the routhierite isotypic series should be based on the chemical data only, taking into account the combination of dominant Me^+ and Me^{2+} cations. In this way the following species can be defined (Fig. 3): (1) routhierite – $\text{Me}^+ = \text{Cu}^+$, $\text{Me}^{2+} = \text{Hg}^{2+}$; (2) stalderite – $\text{Me}^+ = \text{Cu}^+$, $\text{Me}^{2+} = \text{Zn}^{2+}$; (3) arsiccioite – $\text{Me}^+ = \text{Ag}^+$, $\text{Me}^{2+} = \text{Hg}^{2+}$; (4) ralphcannonite – $\text{Me}^+ = \text{Ag}^+$, $\text{Me}^{2+} = \text{Zn}^{2+}$, being the general chemical formula $\text{Me}^+\text{Me}_2^{2+}\text{TlAs}_2\text{S}_6$.

Relationships between the crystal chemistry of 'routhierites' and the ore geochemistry

The members of the routhierite isotypic series are very rare minerals. Well-characterized specimens of routhierite have only been reported so far from Jas Roux, France and the Monte Arsiccio mine, Italy (Johan *et al.*, 1974; Biagioni *et al.*, 2014a). The other three phases have only been reported from their type localities, Monte Arsiccio mine (arsiccioite – Biagioni *et al.*, 2014b) and the Lengenbach quarry (stalderite – Graeser *et al.*, 1995; ralphcannonite – this work).

These minerals can have a complex chemistry, being characterized by the occurrence of Tl, As, Cu, Ag, Hg, Zn and minor Fe and Sb. In particular, the chemical variability of the tetrahedral cations (Cu, Ag, Hg, Zn and Fe) seems to be closely related to the ore geochemistry, as discussed by Biagioni *et al.* (2014b) for the pair routhierite-arsiccioite from Monte Arsiccio. This locality, as well as Jas Roux, are characterized by a relatively large abundance of Hg, as evidenced by the occurrence of the Hg members of the routhierite isotypic series, as well as other Hg phases, i.e. aktashite, $\text{Cu}_6\text{Hg}_3\text{As}_4\text{S}_{12}$ and laffittite, AgHgAsS_3 (Favreau *et al.*, 2011; Biagioni *et al.*, 2014c).

The ore geochemistry at the Lengenbach quarry, on the contrary, seems to be Hg-depleted. In fact, the Zn-dominant (sometimes Fe-enriched) members of the routhierite isotypic series occur there, stalderite and ralphcannonite. In addition, the Zn isotype of aktashite, nowackiite, $\text{Cu}_6\text{Zn}_3\text{As}_4\text{S}_{12}$, has been identified (Marumo and Burri, 1965). The variable Ag/Cu ratio within the crystallizing medium controls the crystallization of the stalderite-ralphcannonite pair, as well as other isotypic compounds described from the Lengenbach quarry, e.g. hatchite, $\text{AgTlPbAs}_2\text{S}_5$ and wallisite, $\text{CuTlPbAs}_2\text{S}_5$.

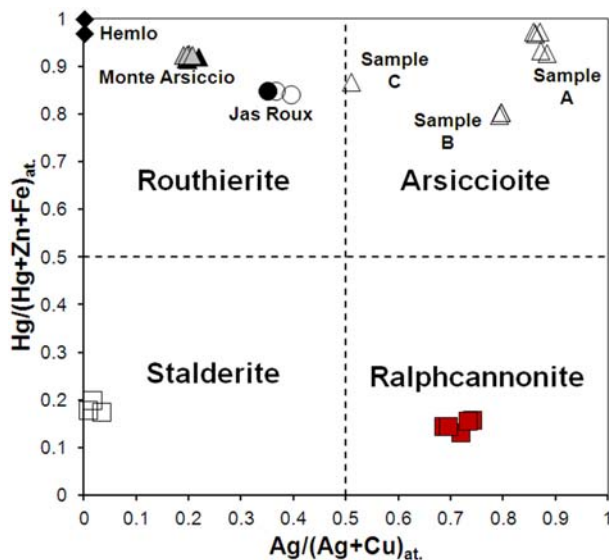


FIG. 3. $\text{Ag}/(\text{Ag} + \text{Cu})$ versus $\text{Hg}/(\text{Hg} + \text{Zn} + \text{Fe})$ atomic ratios in the routhierite isotypic series. Open triangles: arsiccioite from Monte Arsiccio mine (Biagioni *et al.*, 2014b). Filled triangles: routhierite from Monte Arsiccio mine (black, Biagioni *et al.*, 2014b; grey, Biagioni *et al.*, 2014a). Open circles: routhierite from Jas Roux (Johan *et al.*, 1974). Filled circles: routhierite from Jas Roux (Bindi, 2008). Open squares: stalderite from Lengenbach (Graeser *et al.*, 1995). Filled squares (red): ralphcannonite (this work). Filled lozenges: 'Sb-routhierite' from Hemlo (Harris, 1989).

Conclusion

The description of ralphcannonite increases the knowledge about the crystal chemistry of the routhierite isotypic series, bringing new data to the systematics of thallium sulfosalts. The unexpected cation distribution described by Biagioni *et al.* (2014b) in arsiccioite has been confirmed, with Ag, the largest cation, preferentially hosted at the largest $M2$ site: the priority of the steric effect over the valence state in the members of the routhierite isotypic series is thus confirmed.

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