

Magnesiocanutite, $\text{NaMnMg}_2[\text{AsO}_4]_2[\text{AsO}_2(\text{OH})_2]$, a new protonated alluaudite-group mineral from the Torrecillas mine, Iquique Province, Chile

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ABSTRACT

The new mineral magnesiocanutite (IMA2016-057), $\text{NaMnMg}_2[\text{AsO}_4]_2[\text{AsO}_2(\text{OH})_2]$, was found at the Torrecillas mine, Iquique Province, Chile, where it occurs as a secondary phase in association with anhydrite, canutite, halite, lavendulan and magnesiokoritnigite. Magnesiocanutite occurs as pale brownish-pink to rose-pink, lozenge-shaped tablets that are often grouped in tightly intergrown aggregates. The crystal forms are $\{110\}$ and $\{102\}$. Crystals are transparent, with vitreous lustre and white to very pale pink streak. The Mohs hardness is $2\frac{1}{2}$, tenacity is brittle, and the fracture is splintery. Crystals exhibit two perfect cleavages: $\{010\}$ and $\{101\}$. The calculated density is 3.957 g/cm^3 . Optically, magnesiocanutite is biaxial (+), with $\alpha = 1.689(2)$, $\beta = 1.700(2)$, $\gamma = 1.730(2)$ (measured in white light); $2V_{\text{meas.}} = 64.3(4)^\circ$; slight dispersion, $r < v$; orientation $Z = \mathbf{b}$; $X \wedge \mathbf{a} = 15^\circ$ in obtuse angle β . The mineral is slowly soluble in dilute HCl at room temperature. Electron-microprobe analyses, provided Na_2O 5.44, CaO 0.26, MgO 8.84, MnO 18.45, CoO 1.47, CuO 2.13, As_2O_5 59.51, $\text{H}_2\text{O}(\text{calc})$ 2.86, total 98.96 wt.%. Magnesiocanutite is monoclinic, $C2/c$, $a = 12.2514(8)$, $b = 12.4980(9)$, $c = 6.8345(5)$ Å, $\beta = 113.167(8)^\circ$, $V = 962.10(13)$ Å³ and $Z = 4$. The eight strongest powder X-ray diffraction lines are [d_{obs} Å(I)(hkl)]: 6.25(42)(020), 3.566(43)(310, $\bar{1}31$), 3.262(96)($\bar{1}12$), 3.120(59)(002, 131, 040, 221), 2.787(93)(400, 022, 041, 330), 2.718(100)(421, 240, 112, 402), 2.641(42)($\bar{1}32$) and 1.5026(43)(multiple). Magnesiocanutite has a protonated alluaudite-type structure ($R_1 = 2.59\%$ for 789 $F_o > 4\sigma F$ reflections) and is the Mg analogue of canutite. Using the results of both the microprobe analyses and structure refinement, the structurally based empirical formula is $\text{Na}(\text{Mn}_{0.78}\text{Mg}_{0.22})_{\Sigma 1.00}(\text{Mg}_{1.04}\text{Mn}_{0.70}\text{Cu}_{0.15}\text{Co}_{0.11})_{\Sigma 2.00}[\text{AsO}_4]_2[\text{AsO}_2(\text{OH})_2]$.

KEYWORDS: magnesiocanutite, new mineral, canutite, arsenate, alluaudite group, crystal structure, Torrecillas mine, Iquique Province, Chile.

Introduction

OVER the last several years, our investigations of the minerals of the Torrecillas mine, a small, long-inactive arsenic mine in the northern Atacama Desert of Chile, have revealed a remarkable

assemblage of rare secondary chlorides, arsenates and arsenites, many of which are new mineral species. To date, the descriptions of eight new minerals have been published: the chloride leverettite (Kampf *et al.*, 2013a); the arsenates magnesiokoritnigite (Kampf *et al.*, 2013b), canutite (Kampf *et al.*, 2014a), chongite (Kampf *et al.*, 2016a), currierite (Kampf *et al.*, 2017a) and juansilvaite (Kampf *et al.*, 2017b); and the arsenites torrecillasite (Kampf *et al.*, 2014b) and gajardoite

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TABLE 1. Comparative data for magnesiocanutite and canutite.

| | Magnesiocanutite | Canutite |
|-----------------------------------|--|--|
| Ideal formula | $\text{NaMnMg}_2[\text{AsO}_4]_2[\text{AsO}_2(\text{OH})_2]$ | $\text{NaMn}_3[\text{AsO}_4]_2[\text{AsO}_2(\text{OH})_2]$ |
| Space group | $C2/c$ | $C2/c$ |
| a (Å) | 12.2514(8) | 12.3132(5) |
| b (Å) | 12.4980(9) | 12.6042(6) |
| c (Å) | 6.8345(5) | 6.8717(5) |
| β (°) | 113.167(8) | 113.500(8) |
| V (Å ³) | 962.10(13) | 978.02(9) |
| Z | 4 | 4 |
| Density calc (g/cm ³) | 3.957 | 4.119 |
| Optical character | biaxial (+) | biaxial (+) |
| α | 1.689(2) | 1.712(3) |
| β | 1.700(2) | 1.725(3) |
| γ | 1.730(2) | 1.756(3) |
| $2V$ (°) | 64.3(4) | 65.6(4) |

(Kampf *et al.*, 2016b). Herein, we describe the ninth new mineral from the Torrecillas mine, the alluaudite-group arsenate magnesio-canutite. Several other potentially new minerals are still under study.

Magnesio-canutite is a member of the alluaudite group and is isostructural with compounds with the protonated alluaudite-type structure, which have the general formula $AM_1M_2[T_1O_4][T_2O_3(\text{OH})]_2$, in which the A cations occupy large sites in channels of a framework composed of M_1 and M_2 octahedra, and T_1 and T_2 tetrahedra. The name magnesio-canutite was chosen to highlight the fact that the mineral is the Mg analogue of canutite, $\text{NaMn}_3[\text{AsO}_4]_2[\text{AsO}_2(\text{OH})_2]$, with Mg rather than Mn dominant in the M_2 site. The cell parameters, densities and optical properties of canutite and magnesio-canutite are compared in Table 1. Powder X-ray diffraction and energy-dispersive spectroscopy (EDS) indicate that other arsenate members of the alluaudite group containing varying contents of Na, Ca, Mg, Mn, Cu and Fe are also present in the Torrecillas secondary assemblages. Based upon the relatively low temperature of the secondary assemblages and the occurrence of the protonated alluaudite-group minerals canutite and magnesio-canutite, it seems probable that other such alluaudite-group phases are also protonated; however, these phases are difficult to define unambiguously because they are generally not suitable for single-crystal study.

The new mineral and the name have been approved by the International Mineralogical Association (IMA2016-057). The description is based upon one

holotype and one cotype specimen that are deposited in the collections of the Natural History Museum of Los Angeles County, 900 Exposition Boulevard, Los Angeles, CA 90007, USA, catalogue numbers 66273 and 66274, respectively.

Occurrence

The new mineral was found at the Torrecillas mine, Salar Grande, Iquique Province, Tarapacá Region, Chile ($\sim 20^\circ 58' 13'' \text{S } 70^\circ 8' 17'' \text{W}$). Torrecillas Hill, on which the Torrecillas mine is located, is composed of four different rock units. The Coastal Range Batholith (mainly gabbros) extends from the seashore to the Pan-American Road along the base of Torrecillas Hill. At the foot of Torrecillas Hill is a small area of contact metamorphic rocks in which garnet crystals occur in metamorphosed shales. Higher on the hill, the rocks are predominantly andesites and porphyritic lavas of the Jurassic La Negra Formation. The Torrecillas deposit, in which the new mineral is found, consists of two main veins rich in secondary arsenic and copper minerals that intersect metamorphosed marine shales and lavas. These mineralized veins are related genetically to the aforementioned andesites and porphyritic lavas of the Jurassic La Negra Formation. More information on the geology and mineralogy of the area is provided by Gutiérrez (1975).

The rare secondary chlorides, arsenates and arsenites have been found at three main sites on the hill: an upper pit measuring ~ 8 m long and



FIG. 1. Intergrown magnesiocanutite crystals. Field of View = 0.45 mm.

3 m deep, a lower pit ~100 m from the upper pit and measuring ~5 m long and 3 m deep, and a mine shaft adjacent to the lower pit and lower on the hill. Magnesiocanutite was found in the upper pit and in the area of the mine shaft by a collecting party consisting of three of the authors (ARK, MD and AAMD) along with Jochen Schlüter and Joe Marty in February 2014.

The new mineral is a secondary phase occurring in association with anhydrite, canutite, halite, lavendulan and magnesiokoritnigite. The

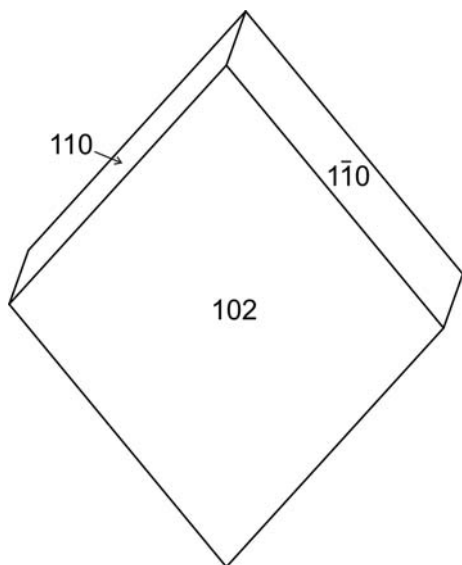


FIG. 2. Crystal drawing of magnesiocanutite, clinographic projection in non-standard orientation, $[20\bar{1}]$ vertical.

secondary assemblages at the Torrecillas deposit are interpreted as having formed from the oxidation of native arsenic and other As-bearing primary phases, followed by later alteration by saline fluids derived from evaporating meteoric water under hyperarid conditions (cf. Cameron *et al.*, 2007).

Physical and optical properties

Magnesiocanutite occurs as pale brownish-pink to rose-pink, lozenge-shaped tablets to ~0.5 mm in maximum dimension. Tablets are often grouped in tightly intergrown aggregates (Fig. 1). The tablets are flattened on $\{102\}$ and slightly elongated on $[20\bar{1}]$; the crystal forms exhibited are $\{110\}$ and $\{102\}$ (Fig. 2). No twinning was observed. Crystals are transparent, with vitreous lustre and white to very pale pink streak. The Mohs hardness is $2\frac{1}{2}$ based upon scratch tests. The tenacity is brittle, and the fracture is splintery as a result of two perfect cleavages, $\{010\}$ and $\{101\}$. Efforts to measure the density by flotation in Clerici solution were unsuccessful because of the difficulty in identifying the most Mg-rich crystal aggregates and the difficulty in observing the aggregates in the solution. The calculated density is 3.957 g/cm^3 based on the empirical formula and 3.764 g/cm^3 based on the ideal formula. The mineral is slowly soluble in dilute HCl at room temperature.

Optically, magnesiocanutite is biaxial (+), with $\alpha = 1.689(2)$, $\beta = 1.700(2)$, $\gamma = 1.730(2)$ determined in white light. The measured $2V$ is $64.3(4)^\circ$ based on extinction data collected on a spindle stage and analysed using *EXCALIBRW* (Gunter *et al.*, 2004). The calculated $2V$ is 63.3° . Slight

TABLE 2. Analytical data (wt.%) for magnesiocanutite.

| Constituent | Mean | Min | Max | SD | Standard |
|--------------------------------|-------|-------|-------|------|-----------|
| Na ₂ O | 5.44 | 5.21 | 5.59 | 0.12 | albite |
| CaO | 0.26 | 0.11 | 0.31 | 0.06 | diopside |
| MgO | 8.84 | 8.17 | 10.35 | 0.69 | diopside |
| MnO | 18.45 | 16.28 | 19.60 | 0.98 | rhodonite |
| CoO | 1.47 | 1.13 | 1.78 | 0.18 | Co metal |
| CuO | 2.13 | 1.20 | 2.75 | 0.51 | Cu metal |
| As ₂ O ₅ | 59.51 | 58.85 | 60.75 | 0.55 | Syn. GaAs |
| H ₂ O* | 2.86 | | | | |
| Total | 98.96 | | | | |

* Calculated on the basis of 3 As, charge balance and 12 O apfu. SD – standard deviation.

TABLE 3. Powder X-ray diffraction data for magnesioacantite.

| I_{obs} | d_{obs} | d_{calc} | I_{calc} | hkl | I_{obs} | d_{obs} | d_{calc} | I_{calc} | hkl |
|------------------|------------------|------------------------------|-------------------|---|------------------|------------------|------------------------------|-------------------|---|
| 4 | 8.37 | 8.3670 | 3 | 1 1 0 | | | 1.7979 | 2 | 6 2 0 |
| 42 | 6.25 | 6.2490 | 49 | 0 2 0 | | | 1.7793 | 3 | $\bar{5}$ 3 3 |
| 21 | 5.59 | 5.6317 | 23 | 2 0 0 | | | 1.7634 | 5 | 1 7 0 |
| 21 | 4.42 | 4.4398 | 14 | 1 1 1 | | | 1.7593 | 2 | $\bar{4}$ 4 3 |
| 33 | 4.13 | 4.1835 | 22 | 2 2 0 | | | 1.7492 | 3 | 1 3 3 |
| 36 | 4.03 | 4.0746 | 39 | $\bar{2}$ 2 1 | 13 | 1.7299 | 1.7257 | 12 | $\bar{1}$ 7 1 |
| 13 | 3.902 | 3.9073 | 17 | 1 3 0 | | | 1.7215 | 3 | $\bar{4}$ 6 1 |
| 10 | 3.776 | 3.7982 | 10 | $\bar{3}$ 1 1 | | | 1.7100 | 2 | 4 2 2 |
| 43 | 3.566 | { 3.5957 3.5419 3.3639 | { 33 15 7 | { 3 1 0 $\bar{1}$ 3 1 $\bar{2}$ 0 2 | 32 | 1.6890 | { 1.6955 1.6805 1.6734 | { 26 18 9 | { $\bar{2}$ 0 4 $\bar{6}$ 4 2 5 5 0 |
| 96 | 3.262 | { 3.2727 3.1417 3.1319 | { 100 20 26 | { $\bar{1}$ 1 2 0 0 2 1 3 1 | 19 | 1.6666 | { 1.6712 1.6608 1.6463 | { 3 2 3 | { 1 7 1 $\bar{1}$ 5 3 $\bar{4}$ 6 2 |
| 59 | 3.120 | { 3.1245 3.0905 | { 13 11 | { 0 4 0 2 2 1 | 6 | 1.6007 | { 1.6091 1.6023 | { 7 2 | { 6 4 0 $\bar{7}$ 3 2 |
| 19 | 2.970 | { 2.9887 2.9620 | { 19 6 | { $\bar{3}$ 1 2 $\bar{2}$ 2 2 | | | { 1.5959 1.5452 | { 6 8 | { 7 1 0 4 4 2 |
| 93 | 2.787 | { 2.8159 2.8069 2.7977 | { 28 4 15 | { 4 0 0 0 2 2 0 4 1 | 23 | 1.5324 | { 1.5444 1.5390 1.5371 | { 7 2 2 | { $\bar{3}$ 7 2 $\bar{6}$ 0 4 $\bar{2}$ 6 3 |
| | | { 2.7890 2.7463 | { 57 9 | { 3 3 0 $\bar{4}$ 2 1 | | | { 1.5316 1.5287 | { 3 13 | { $\bar{1}$ 3 4 $\bar{8}$ 0 2 |
| 100 | 2.718 | { 2.7322 2.6935 2.6871 | { 74 15 30 | { 2 4 0 1 1 2 $\bar{4}$ 0 2 | | | { 1.5274 1.5161 1.5117 | { 3 4 7 | { 2 4 3 0 8 1 $\bar{5}$ 3 4 |
| 42 | 2.641 | 2.6299 | 44 | $\bar{1}$ 3 2 | | | 1.5104 | 4 | $\bar{7}$ 3 3 |
| 18 | 2.578 | 2.5673 | 20 | 4 2 0 | 43 | 1.5026 | 1.5054 | 2 | 2 8 0 |
| | | 2.4686 | 3 | $\bar{4}$ 2 2 | | | 1.5019 | 15 | 3 7 1 |
| | | 2.4028 | 4 | $\bar{5}$ 1 1 | | | 1.5010 | 14 | 7 3 0 |
| 8 | 2.372 | 2.3748 | 11 | 2 0 2 | | | 1.5002 | 2 | $\bar{2}$ 8 1 |
| 3 | 2.296 | 2.3110 | 4 | 3 3 1 | | | 1.4988 | 8 | 1 7 2 |
| 13 | 2.205 | 2.2170 | 17 | 5 1 0 | | | 1.4944 | 2 | $\bar{6}$ 2 4 |
| | | 2.2073 | 2 | $\bar{3}$ 1 3 | | | 1.4902 | 3 | $\bar{2}$ 4 4 |
| | | 2.1853 | 5 | $\bar{4}$ 4 1 | | | 1.4849 | 2 | $\bar{8}$ 2 2 |
| 4 | 2.0652 | 2.0807 | 8 | 3 5 0 | 18 | 1.4809 | 1.4810 | 8 | $\bar{4}$ 4 4 |
| | | 2.0211 | 20 | $\bar{5}$ 3 2 | | | 1.4769 | 2 | 0 6 3 |
| | | 2.0188 | 3 | 3 1 2 | | | 1.4680 | 2 | 1 1 4 |
| 21 | 2.0021 | { 2.0151 2.0121 | { 3 2 | { $\bar{4}$ 2 3 $\bar{1}$ 5 2 | | | 1.4542 | 2 | $\bar{6}$ 6 1 |
| | | 1.9934 | 2 | $\bar{6}$ 0 2 | | | 1.4460 | 4 | 6 4 1 |
| | | 1.9859 | 5 | 0 2 3 | | | 1.4336 | 2 | $\bar{7}$ 1 4 |
| | | 1.9816 | 2 | 5 3 0 | | | 1.4257 | 3 | $\bar{7}$ 5 2 |
| | | 1.9747 | 7 | $\bar{3}$ 3 3 | | | 1.4169 | 2 | $\bar{2}$ 8 2 |
| 26 | 1.9513 | { 1.9422 1.9404 | { 4 22 | { $\bar{2}$ 6 1 $\bar{3}$ 5 2 | 18 | 1.3971 | { 1.4035 1.3992 1.3968 | { 7 5 2 | { 0 4 4 5 7 0 $\bar{1}$ 7 3 |
| 13 | 1.9152 | { 1.9317 1.8991 | { 5 2 | { $\bar{6}$ 2 1 $\bar{6}$ 2 2 | | | 1.3932 | 4 | 1 3 4 |
| | | 1.8772 | 3 | 6 0 0 | | | 1.3919 | 8 | $\bar{3}$ 7 3 |
| | | 1.8580 | 2 | 3 5 1 | | | 1.3912 | 4 | $\bar{4}$ 8 1 |
| | | 1.8522 | 17 | 1 5 2 | 16 | 1.3813 | 1.3888 | 7 | 6 0 2 |
| 34 | 1.8393 | { 1.8405 1.8362 | { 6 20 | { $\bar{2}$ 4 3 3 3 2 | | | 1.3792 | 6 | 2 0 4 |
| | | | | | | | 1.3732 | 4 | $\bar{8}$ 4 2 |

Only calculated lines with intensities of 3 or greater are shown.

$r < v$ dispersion was observed. The optical orientation is $Z = \mathbf{b}$; $X \wedge \mathbf{a} = 15^\circ$ in obtuse angle β . Pleochroism is imperceptible.

Composition

Quantitative analyses (nine points on four crystals) were performed at the University of Utah on a Cameca SX-50 electron microprobe with four wavelength-dispersive spectrometers (WDS) utilizing *Probe for EPMA* software. Analytical conditions were: 15 kV accelerating voltage, 10 nA beam current and a beam diameter of 10 μm . Magnesiocanutite exhibited visible damage under the electron beam. Sodium experienced a time-dependent decrease in intensity under the electron beam, whereas As sustained an increase; both were accounted for by an exponential fit to the intensity

vs. time measurements and extrapolation to zero-time intensity.

No other elements were detected by EDS. Other possible elements were sought by WDS scans, but none were above the detection limits. Raw X-ray intensities were corrected for matrix effects with a ϕ (ρz) algorithm (Pouchou and Pichoir, 1991). Because insufficient material was available for a direct determination of H_2O , the amount of water was calculated on the basis of 3 As atoms per formula unit (apfu), charge balance and 12 O apfu, as determined by the crystal structure analysis (see below). Analytical data are given in Table 2.

The empirical formula is $(\text{Na}_{1.02}\text{Ca}_{0.03}\text{Mn}_{1.51}\text{Mg}_{1.27}\text{Cu}_{0.16}\text{Co}_{0.11})_{\Sigma 4.10}\text{As}_3\text{O}_{12}\text{H}_{1.84}$. The simplified structural formula is $\text{Na}(\text{Mn},\text{Mg})(\text{Mg},\text{Mn})_2[\text{AsO}_4]_2[\text{AsO}_2(\text{OH})_2]$ and the end-member formula is $\text{NaMnMg}_2[\text{AsO}_4]_2[\text{AsO}_2(\text{OH})_2]$, which requires Na_2O 5.68, MgO 14.78, MnO

TABLE 4. Data collection and structure refinement details for magnesiocanutite.

| | |
|--|--|
| Diffractionmeter | Rigaku R-Axis Rapid II |
| X-ray radiation / power | MoK α ($\lambda = 0.71075 \text{ \AA}$)/50 kV, 40 mA |
| Temperature | 298(2) K |
| Structural formula | $\text{Na}(\text{Mn}_{0.748}\text{Mg}_{0.252})_{\Sigma 1.00}(\text{Mg}_{1.114}\text{Mn}_{0.886})_{\Sigma 2.00}[\text{AsO}_4]_2[\text{AsO}_2(\text{OH})_2]$ |
| Space group | $C2/c$ |
| Unit cell dimensions | $a = 12.2514(8) \text{ \AA}$ $b = 12.4980(9) \text{ \AA}$ $c = 6.8345(5) \text{ \AA}$ $\beta = 113.167(8)^\circ$ |
| V | $962.10(13) \text{ \AA}^3$ |
| Z | 4 |
| Density (for above formula) | 3.899 g/cm^3 |
| Absorption coefficient | 12.610 mm^{-1} |
| $F(000)$ | 1061 |
| Crystal size (μm) | $75 \times 45 \times 20$ |
| θ range ($^\circ$) | 3.26 to 25.03 |
| Index ranges | $-14 \leq h \leq 12$, $-14 \leq k \leq 14$, $-8 \leq l \leq 8$ |
| Refls. collected/unique | 4478 / 849; $R_{\text{int}} = 0.051$ |
| Reflections with $F_o > 4\sigma F$ | 789 |
| Completeness to $\theta = 25.03^\circ$ | 99.3% |
| Max. and min. transmission | 0.787 and 0.451 |
| Refinement method | Full-matrix least-squares on F^2 |
| Parameters refined | 93 |
| Goof | 1.110 |
| Final R indices [$F_o > 4\sigma(F)$] | $R_1 = 0.0259$, $wR_2 = 0.0542$ |
| R indices (all data) | $R_1 = 0.0286$, $wR_2 = 0.0556$ |
| Largest diff. peak / hole | $+0.74 / -0.63 \text{ e/\AA}^3$ |

* $R_{\text{int}} = \Sigma |F_o^2 - F_c^2(\text{mean})| / \Sigma [F_o^2]$. Goof = $S = \{\Sigma [w(F_o^2 - F_c^2)^2] / (n-p)\}^{1/2}$. $R_1 = \Sigma |F_o| - |F_c| / \Sigma |F_o|$. $wR_2 = \{\Sigma [w(F_o^2 - F_c^2)^2] / \Sigma [w(F_c^2)^2]\}^{1/2}$; $w = 1 / [\sigma^2(F_o^2) + (aP)^2 + bP]$ where a is 0.0086, b is 5.1983 and P is $[2F_c^2 + \text{Max}(F_o^2, 0)] / 3$.

13.01, As₂O₅ 63.22, H₂O 3.30, total 100 wt.%. The Gladstone-Dale compatibility $1 - (K_p/K_c)$ for the empirical formula is 0.004 in the range of superior compatibility (Mandarino, 2007).

X-ray crystallography and structure refinement

Both powder and single-crystal X-ray diffraction studies were carried out using a Rigaku R-Axis Rapid II curved imaging plate microdiffractometer, with monochromatic MoK α radiation. For the powder diffraction study a Gandolfi-like motion on the ϕ and ω axes was used to randomize the sample and observed d -values and intensities were derived by profile fitting using *JADE 2010* software (Materials Data, Inc.). The powder data presented in Table 3 show good agreement with the pattern calculated from the structure determination. Unit-cell parameters refined from the powder data using *JADE 2010* with whole pattern fitting are $a = 12.183(5)$, $b = 12.506(6)$, $c = 6.802(5)$ Å, $\beta = 112.803(13)^\circ$ and $V = 955.4(9)$ Å³.

The *Rigaku CrystalClear* software package was used for processing the structure data, including the application of an empirical multi-scan absorption correction using *ABSCOR* (Higashi, 2001). The structure was solved using *SIR2011* (Burla *et al.*, 2012). *SHELXL-2013* (Sheldrick, 2015) was used for the refinement of the structure. The location of all non-hydrogen atom sites was straightforward and atom coordinates were then transformed to be consistent with the standardized atom coordinates for alluaudite-group structures (Krivovichev *et al.*, 2013). (Note that the atom coordinates and O atom assignments for canutite were not reported using this standardized approach.) The *A* site was assigned full occupancy by Na. The *M1* and *M2* sites were refined with joint occupancies by Mn and Mg, yielding *M1*: Mn_{0.748}Mg_{0.252} and *M2*: Mn_{0.443}Mg_{0.557}. A difference-Fourier map revealed the location of the H atom corresponding to the O2...O4 hydrogen bond. The data collection and refinement details are given in Table 4, atom coordinates and displacement parameters in Table 5, selected bond distances in Table 6 and a bond valence analysis in Table 7.

Description of the structure

Magnesiocanutite is a member of the alluaudite group and is isostructural with compounds with the protonated alluaudite-type structure, including the

TABLE 5. Atom coordinates and displacement parameters (Å²) for magnesiocanutite.

| | <i>x/a</i> | <i>y/b</i> | <i>z/c</i> | U_{eq} | U^{11} | U^{22} | U^{33} | U^{23} | U^{13} | U^{12} |
|-------|------------|------------|-------------|-------------|------------|------------|------------|--------------|------------|-------------|
| A(Na) | 0 | -0.0103(3) | 1/4 | 0.0458(9) | 0.0219(16) | 0.093(3) | 0.0204(16) | 0 | 0.0057(13) | 0 |
| M1* | 0 | 0.28224(8) | 1/4 | 0.0161(4) | 0.0158(7) | 0.0162(6) | 0.0167(7) | 0 | 0.0068(5) | 0 |
| M2* | 0.29203(8) | 0.65987(7) | 0.37797(14) | 0.0135(4) | 0.0147(6) | 0.0113(5) | 0.0155(6) | 0.0006(3) | 0.0072(4) | 0.0006(3) |
| As1 | 0 | 0.69083(5) | 1/4 | 0.01527(19) | 0.0169(3) | 0.0144(3) | 0.0125(3) | 0 | 0.0036(3) | 0 |
| As2 | 0.21940(4) | 0.88691(3) | 0.11470(6) | 0.01458(16) | 0.0173(3) | 0.0127(3) | 0.0132(3) | -0.00007(15) | 0.0054(2) | 0.00009(17) |
| O1 | 0.4640(3) | 0.7337(2) | 0.5293(4) | 0.0188(7) | 0.0163(15) | 0.0235(16) | 0.0147(15) | -0.0047(12) | 0.0041(13) | -0.0001(13) |
| O2 | 0.1129(3) | 0.6077(2) | 0.2654(5) | 0.0183(7) | 0.0148(15) | 0.0126(15) | 0.0272(18) | 0.0009(12) | 0.0079(13) | 0.0026(12) |
| H | 0.114(5) | 0.517(4) | 0.288(7) | 0.027 | | | | | | |
| O3 | 0.3443(3) | 0.6757(2) | 0.1211(4) | 0.0184(7) | 0.0215(17) | 0.0190(16) | 0.0141(16) | -0.0016(12) | 0.0062(13) | -0.0043(12) |
| O4 | 0.1336(3) | 0.4110(2) | 0.3336(5) | 0.0206(7) | 0.0173(16) | 0.0168(15) | 0.0303(18) | 0.0006(13) | 0.0123(14) | 0.0036(12) |
| O5 | 0.2139(3) | 0.8142(2) | 0.3164(4) | 0.0162(7) | 0.0183(16) | 0.0181(15) | 0.0128(15) | 0.0044(11) | 0.0068(13) | 0.0002(12) |
| O6 | 0.3515(3) | 0.5038(2) | 0.4029(5) | 0.0194(7) | 0.0197(17) | 0.0164(15) | 0.0212(18) | 0.0012(12) | 0.0069(14) | -0.0022(12) |

* Refined occupancies: M1: Mn/Mg = 0.748/0.252(8); M2: Mn/Mg = 0.443/0.557(6).

TABLE 6. Selected bond distances (Å) and angles (°) for magnesiocanutite.

| | | | |
|---------------|----------|----------------|----------------|
| A–O6 (×2) | 2.356(3) | M1–O4 (×2) | 2.204(3) |
| A–O6 (×2) | 2.435(3) | M1–O3 (×2) | 2.206(3) |
| A–O3 (×2) | 2.915(4) | M1–O1 (×2) | 2.206(3) |
| A–O1 (×2) | 3.122(4) | <M1–O> | 2.205 |
| A–O5 (×2) | 3.309(4) | M2–O6 | 2.066(3) |
| <A–O> | 2.827 | M2–O3 | 2.101(3) |
| As1–O1 (×2) | 1.684(3) | M2–O5 | 2.120(3) |
| As1–O2 (×2) | 1.700(3) | M2–O2 | 2.122(3) |
| <As–O> | 1.692 | M2–O5 | 2.143(3) |
| As2–O5 | 1.675(3) | M2–O1 | 2.156(3) |
| As2–O6 | 1.679(3) | <M2–O> | 2.118 |
| As2–O3 | 1.682(3) | | |
| As2–O4 | 1.720(3) | | |
| <As–O> | 1.689 | | |
| Hydrogen bond | | | |
| D–H | $d(D-H)$ | $d(H\cdots A)$ | <DHA |
| O2–H | 1.14(5) | 1.37(6) | 171(6) |
| | | | $d(D\cdots A)$ |
| | | | A |
| | | | O4 |

minerals canutite, $\text{NaMn}_3[\text{AsO}_4]_2[\text{AsO}_2(\text{OH})_2]$ (Kampf *et al.*, 2014a), o'danielite, $\text{NaZn}_3[\text{AsO}_4][\text{AsO}_3(\text{OH})]_2$ (Keller and Hess, 1988) and groatite, $\text{NaCaMn}_2[\text{PO}_4][\text{PO}_3(\text{OH})]_2$ (Cooper *et al.*, 2009). The general formula of these phases is $AM_1M_2[T_1O_4][T_2O_3(\text{OH})]_2$ { $T = \text{P, As}$ }, where the M_1 and M_2 octahedra link by edge-sharing to form staggered chains, T_1 and T_2 tetrahedra cross-link the chains, and A cations occupy large channel sites (Fig. 3).

The structure refinement for magnesiocanutite clearly indicates that Mn and Mg can both occupy the M_1 and M_2 sites, but that there is a distinct

preference of Mn for the M_1 site and Mg for the M_2 site. Although the small amount of Co and Cu indicated in the chemical analysis was not incorporated into the structure refinement, we included these in an analysis of the M_1 and M_2 site occupancies using the program *OccQP* (Wright *et al.*, 2001). This program uses quadratic equations in a constrained least-squares formulation to optimize occupancy assignments based upon site scattering, chemical composition, charge balance, bond valence and cation–anion bond lengths. Assuming full occupancies and using the chemical analytical values for Mn, Mg, Co and Cu, this analysis indicated the M_1 site to

TABLE 7. Bond-valence analysis for magnesiocanutite. Values are expressed in valence units.

| | O1 | O2 | O3 | O4 | O5 | O6 | Σ_c |
|------------|----------|----------|----------|----------|--------------|----------------------|------------|
| A(Na) | 0.03 ×2→ | | 0.05 ×2→ | | 0.02 ×2→ | 0.22 ×2→ 0.18 ×2→ | 1.00 |
| M1 | 0.31 ×2→ | | 0.31 ×2→ | 0.31 ×2→ | | | 1.86 |
| M2 | 0.32 | 0.36 | 0.38 | | 0.36 0.33 | 0.41 | 2.16 |
| As1 | 1.25 ×2→ | 1.20 ×2→ | | | | | 4.90 |
| As2 | | | 1.26 | 1.13 | 1.29 | 1.27 | 4.95 |
| Σ_a | 1.91 | 1.56 | 2.00 | 1.44 | 2.00 | 2.08 | |

Multiplicities indicated by ×2→; bond strengths based upon refined Mn/Mg site occupancies; $\text{Na}^+ - \text{O}$ bond-valence parameters are from Wood and Palenik (1999); other bond-valence parameters are from Brown and Altermatt (1985); hydrogen-bond donor-acceptor contributions for $\text{O2}\cdots\text{O4}$ are not included.

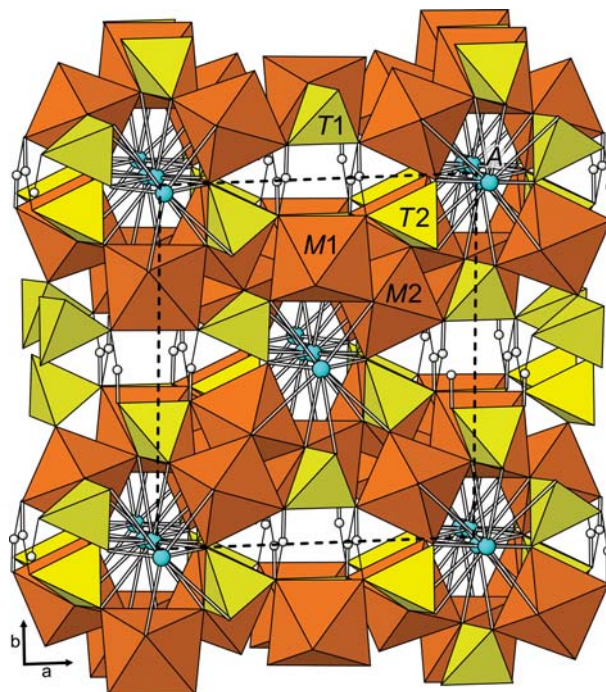


FIG. 3. The crystal structure of magnesiocanutite viewed slightly canted down [001]. Unit cell shown by dashed line. A–O and O–H bonds shown as sticks. Hydrogen bonds shown as thin solid lines.

be occupied by 0.784 Mn and 0.216 Mg and the *M2* site to be occupied by 0.350 Mn, 0.518 Mg, 0.056 Co and 0.076 Cu, providing the structurally based empirical formula $\text{Na}(\text{Mn}_{0.78}\text{Mg}_{0.22})_{\Sigma 1.00}(\text{Mg}_{1.04}\text{Mn}_{0.70}\text{Cu}_{0.15}\text{Co}_{0.11})_{\Sigma 2.00}[\text{AsO}_4]_2[\text{AsO}_2(\text{OH})_2]$.

In the structure of canutite and other protonated alluaudite-type minerals, O4 [part of *T2* (As2) tetrahedron] is an OH group forming a hydrogen bond to O2 [part of *T1* (As1) tetrahedron]. An interesting feature of the magnesiocanutite structure refinement is that the H atom position appears to be closer to O2 than to O4, O2–H = 1.14 and O4–H = 1.37 Å; however, it is worth noting that there is also a residual electron density closer to O2. The very short O2–O4 distance is typical for a symmetrical hydrogen bond, although in this case, the bond is somewhat asymmetrical. The bond-valence sums for O2 and O4 without the H contributions are 1.56 and 1.44 valence units, respectively, consistent with such a hydrogen bond character.

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