

NMNS#12965 before an atomic site became dominant in Zn (i.e. c. 20 wt.% ZnO). The distribution of Zn in these sonolite samples could not have been determined without the linear equation solutions.

#### Acknowledgements

The authors thank Dr F. C. Hawthorne, University of Manitoba, for making available the four-circle diffractometer.

#### References

- Cromer, D. T. and Liberman, D. (1970) Relativistic calculation of anomalous scattering factors for X-rays. *J. Chem. Phys.*, **53**, 1891–8.
- Cromer, D. T. and Mann, J. B. (1968) X-ray scattering factors computed from numerical Hartree-Fock wave functions. *Acta Cryst.*, **A24**, 321–4.
- Dunn, P. J. (1985) Manganese humites and leuco-phoenicitites from Franklin and Sterling Hill, New Jersey: parageneses, compositions, and implications for solid solution limits. *Amer. Mineral.*, **70**, 379–87.
- Francis, C. A. (1985) Crystal structure refinement of magnesian alleghanyite. *Amer. Mineral.*, **70**, 182–5.
- Francis, C. A. and Ribbe, P. R. (1978) Crystal structures of the humite minerals: V. Magnesian manganhumite. *Amer. Mineral.*, **63**, 874–7
- Kato, T., Ito, Y. and Hashimoto, N. (1989) The crystal structures of sonolite and jerrygibbsite. *Neues Jahrb. Mineral., Mh.*, 410–30.
- North, A. C. T., Phillips, D. C. and Mathews, F. S. (1968) A semi-empirical method of absorption correction. *Acta Cryst.*, **A24**, 351–9.
- Ribbe, P. H. (1982) The humite series and Mn-analogs. *Reviews in Mineralogy, Orthosilicates* (Ribbe, P. H., ed.), **5**, 231–74, Mineral. Soc. Amer., Washington, D.C.
- Shannon, R. D. (1976) Revised effective ionic radii and systematic studies of interatomic distances in halides and chalcogenides. *Acta Cryst.*, **A32**, 751–67.
- Sheldrick, G. M. (1990) SHELXTL, a crystallographic computing package, revision 4.1. Siemens Analytical Instruments, Inc., Madison, Wisconsin.

[Manuscript received 3 June 1993;  
revised 3 August 1993]

© Copyright the Mineralogical Society

KEYWORDS: crystal structure, sonolite, New Jersey

*Mineral Sciences Section,  
Canadian Museum of Nature,  
Ottawa,  
Ontario KIP 6P4, Canada.*

JOEL D. GRICE

*Department of Mineral Sciences,  
Smithsonian Institution,  
Washington, D.C. 20560, USA.*

PETE J. DUNN

MINERALOGICAL MAGAZINE, JUNE 1994, VOL. 58, PP. 336–338

## Lead oxychlorides at Elura, western NSW, Australia

SEVERAL different Pb oxychloride minerals are known to form under saline conditions, the most common being laurionite  $[\text{Pb}(\text{OH})\text{Cl}]$  and mendipite  $[\text{Pb}_3\text{O}_2\text{Cl}_2]$ . Several others, penfieldite

$[\text{Pb}_2\text{Cl}_3(\text{OH})]$  and fiedlerite  $[\text{Pb}_3\text{Cl}_4(\text{OH})_2]$  are known mainly as marine corrosion products of ancient Pb objects or slags (Edwards *et al.*, 1992). The rare minerals, blixite  $[\text{Pb}_2\text{Cl}(\text{O},\text{OH})_{2-x}]$  where

TABLE 1. Ideal compositions of lead oxychlorides (wt.%)

Mineral Formula	Laurionite PbCl(OH)	Mendipite Pb <sub>3</sub> O <sub>2</sub> Cl <sub>2</sub>	Blixite <sup>1</sup> Pb <sub>2</sub> Cl(O,OH) <sub>1.7</sub>	Damaraita Pb <sub>4</sub> O <sub>3</sub> Cl <sub>2</sub>	Brilon mineral <sup>2</sup> Pb <sub>4</sub> O <sub>3</sub> (Cl,SO <sub>4</sub> ) <sub>2</sub>
Pb	79.8	85.8	87.0	87.5	86.2
Cl	13.7	9.8	7.7	7.5	5.6
O	6.2	4.4	5.7	5.1	6.8
H	0.4	-	0.1	-	-
S	-	-	-	-	1.4

<sup>1</sup> actual composition determined by Gabrielson *et al.*, 1958

<sup>2</sup> actual composition determined by Rouse and Dunn, 1990

$x \sim 0.3$ ] and damaraita [Pb<sub>4</sub>O<sub>3</sub>Cl<sub>2</sub>] have been recorded only at Langban, Sweden (Gabrielson *et al.*, 1958) and the Kombat Mine, Namibia (Criddle *et al.*, 1990) respectively. This note documents the presence of blixite and mendipite within the weathered profile at the Elura Zn-Pb-Ag deposit, 43 km NE of Cobar and 600km NE of Sydney, New South Wales.

Mineralization at Elura grades 5.8% Pb, 8.4% Zn and 130 g/t Ag and when fresh consists of assemblages of pyrite, sphalerite, pyrrhotite, galena and minor chalcopyrite in a gangue of siderite, quartz, muscovite and chlorite.

Weak weathering at about 130m depth results in the partial destruction of siderite and sphalerite with the intensity increasing until at 100 m only oxidized material is present (Taylor *et al.*, 1984). Supergene sulphides characterized by pyrite, galena, chalcocite, digenite and covellite occur at about 100 m as an interface between the oxidate zone and primary sulphide-bearing assemblages. Although this supergene sulphide zone is only a few metres thick, it exhibits gross zonation from Pb-rich (Pb = 30–50%) to Cu-rich assemblages (Cu > 35%) at its top. As well as the sulphides, anglesite, cerussite, blixite and mendipite are developed in this narrow zone.

Because of the relatively small size of the aggregates of crystals (~50 × 20 μm), the compositions of the Pb oxychlorides and their relationships to other lead minerals at Elura have been studied using an electron microprobe. Although chemical data readily differentiate most of the naturally occurring Pb oxychlorides, blixite [Pb<sub>2</sub>Cl(O,OH)<sub>2-x</sub>] and damaraita [Pb<sub>4</sub>O<sub>3</sub>Cl<sub>2</sub>] have similar proportions of Pb and Cl (Table 1). Thus, whereas the Pb oxychloride with ~9% Cl at Elura is readily identified as mendipite (Table 2), lower Cl content material could be either blixite or damaraita. However, the presence of an X-ray diffraction peak at 2.93Å

(the most intense peak for blixite, JCPDS 12-542) suggests the former. The blixite/mendipite association in the Mendip Hills, England (Symes and Embrey, 1977) is also consistent with a similar association at Elura (see below). Although some S occurs in the Pb oxychlorides at Elura (Table 2), this probably represents contamination from associated anglesite rather than incorporation of SO<sub>4</sub> into Cl sites as in blixite-type occurrences at Brilon, Germany (Rouse and Dunn, 1990). Lower S in mendipite replacing cerussite than in that replacing anglesite (Table 2) is consistent with this interpretation.

In Pb oxychloride-bearing samples from Elura, the following associations are present. Within sample 71072, mendipite replaces both cerussite and anglesite (Fig. 1) but because anglesite extensively replaces cerussite (Fig. 2), the mendi-

TABLE 2. Compositions of lead oxychlorides, Elura (wt.%)

Sample No. No. of analyses	Mendipite		Blixite	
	71072	71072	71073	71073
	6 <sup>1</sup>	1 <sup>2</sup>	4	4
Pb	83.43	83.52	85.80	85.19
Cl	9.24	8.29	8.84	6.20
Cu	0.23	0.18	0.13	0.11
Zn	0.01	0.03	0.04	0.06
Fe <sup>III</sup>	1.07	0.92	0.50	0.19
S	0.28	0.10	0.61	0.22
O <sup>3</sup>	5.28	5.19	5.77	5.63
Total	99.54	98.23	101.69	97.60

<sup>1</sup> mendipite replacing anglesite

<sup>2</sup> mendipite replacing cerussite

<sup>3</sup> determined by stoichiometry

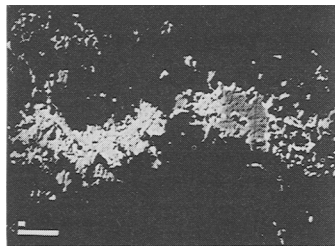


FIG. 1. Mendipite (dark grey) replacing anglesite (pale grey). Sample 71072. Scale bar  $\equiv$  10  $\mu$ m.

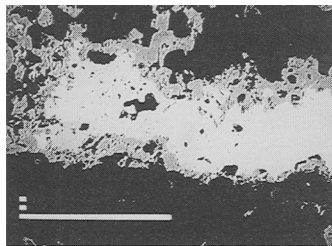


FIG. 2. Anglesite (grey) replacing cerussite (white). Sample 71072. Scale bar  $\equiv$  100  $\mu$ m.

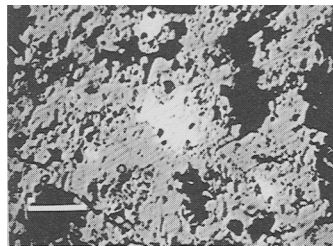


FIG. 3. Blixite/mendipite (white) replaced by anglesite (grey). Sample 71073. Scale bar  $\equiv$  10  $\mu$ m.

pite is probably all formed subsequent to anglesite. However in sample 71073 blixite or blixite/mendipite is replaced by anglesite (Fig. 3) with residual Cu sulphides often present in the Pb oxychlorides. Where blixite/mendipite masses are present, the blixite appears to replace mendipite.

Mendipite formation is restricted to temperatures above 29°C (Edwards *et al.*, 1992) but oxidation of sulphides could easily generate such a temperature at Elura. Therefore, the observed blixite replacement of mendipite could result from a decreasing temperature as well as decreased Cl or increased pH (Edwards *et al.*, 1992). The formation of mendipite also requires low  $\text{CO}_3^{2-}$  and  $\text{SO}_4^{2-}$  environments; hence the replacement of anglesite probably reflects an environment where  $\text{Cl}^-$  has been residually concentrated as  $\text{CO}_3^{2-}$  and  $\text{SO}_4^{2-}$  have been deposited as cerussite and anglesite. Anglesite replacement of mendipite implies exhaustion of  $\text{Cl}^-$  and a new supply of  $\text{SO}_4^{2-}$  being introduced. These features imply either two periods of anglesite formation at Elura or different processes occurring in different micro-environments within the supergene sulphide zone. Either case reflects the operation of complex weathering processes during supergene sulphide formation.

#### Acknowledgements

Ken Kinealy (CSIRO, Division of Exploration Geoscience) assisted in operating the electron microprobe. Discussions with Peter Williams (Univ. Western Sydney) have stimulated interest

in documenting the Pb oxychlorides at Elura. Comments by Peter Williams and Bill Birch on an earlier draft are gratefully acknowledged.

#### References

- Criddle, A. J., Keller, P., Stanley, C. J. and Innes, J. (1990) Damaraitite, a new lead oxychloride mineral from the Kombat mine, Namibia (South West Africa). *Mineral. Mag.*, **54**, 593–8.
- Edwards, R., Gillard, R. D., Williams, P. A. and Pollard, A. M. (1992) Studies of secondary mineral formation in the PbO–H<sub>2</sub>O–HCl system. *Mineral. Mag.*, **56**, 53–65.
- Gabrielson, O., Parwel, A. and Wickman, F. E. (1958) Blixite, a new lead oxyhalide from Langban. *Arkiv. Mineral. Geol. Stockholm*, **2**, 411–5.
- Rouse, R. C. and Dunn, P. J. (1990) A new lead sulfate oxychloride related to the nadorite group from Brilon, Germany. *Neues Jahrb. Mineral., Mh.*, 337–42.
- Symes, R. F. and Embrey, P. G. (1977) Mendipite and other rare oxychloride minerals from the Mendip Hills, Somerset, England. *Mineral. Record*, **8**, 298–303.
- Taylor, G. F., Wilmshurst, J. R., Togashi, Y. and Andrew, A. S. (1984) Geochemical and mineralogical haloes about the Elura Zn–Pb–Ag orebody, western New South Wales. *J. Geochem. Expl.*, **22**, 265–90.

[Manuscript received 20 May 1993]

© Copyright the Mineralogical Society

KEYWORDS: lead oxychlorides, blixite, mendipite, Elura deposit, New South Wales, Australia.

CSIRO Division of Exploration and Mining,  
PO Box 136, North Ryde,  
NSW 2113, Australia

K. M. SCOTT