# **Short Communications**

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Acta Cryst. (1963). 16, 834

The neutron coherent scattering length for magnesium. By T. M. SABINE and J. D. BROWNE, Australian Atomic Energy Commission, Research Establishment, Lucas Heights, N.S.W., Australia

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The coherent scattering length of magnesium was given as  $0.52 \pm 0.01 \times 10^{-12}$  cm by Bacon (1952) from measurements with MgAl<sub>2</sub>O<sub>4</sub> and MgO. However, the value given in *International Tables for X-ray Crystallography* (1962) is  $0.54 \times 10^{-12}$  cm. This figure was a compromise (Bacon, private communication), taking account also of measurements of the total and incoherent cross-section of magnesium. In connection with work in progress in this laboratory neutron powder intensities have been measured for MgO. The coherent scattering length for Mg deduced from these data, using the accepted value of  $0.577 \times 10^{-12}$  cm for oxygen, is  $0.516 \pm 0.006 \times 10^{-12}$  cm, which confirms Bacon's original lower figure.

## References

BACON, G. E. (1952). Acta Cryst. 5, 684.
International Tables for X-ray Crystallography. (1962).
Vol. III, p. 229. Birmingham: Kynoch Press.

#### Acta Cryst. (1963). 16, 834

X-ray evidence of plutonium(III) oxalate decahydrate. By D. M. CHACKRABURTTY, Radiochemistry and Isotope Division, Atomic Energy Establishment, Trombay, Bombay-73, India

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A bluish-white precipitate has been obtained by adding excess of 0.06M oxalic acid in plutonium(III) solution, prepared by adding 1 ml of 10% hydroxylamine hydrochloride solution to 30 mg of plutonium(IV) in 0.5Mnitric acid. This precipitate has been studied by means of the thermogravimetric balance in air and in an argon atmosphere and it has been suggested that the first peak in the thermogravimetric curve has a molecular formula corresponding to  $Pu_2(C_2O_4)_3.10H_2O$  (Dawson, 1952,

Table	1.	X-ray	diffraction	data	of
	Р	u.(C.O	). 10 H.O		

~ `	$\alpha_{2}(\bigcirc_{2}\bigcirc_{4})_{3}$	01220
d	Ι	hkl
6·669 Å	s+	011, 111
5.132	mw	$200, 10\overline{2}$
4·974	mw +	$21\overline{1}$
4·681	m +	020, 111
4·567	vw	002, 210
3.523	w	20 <b>3</b> , 102
<b>3</b> ∙090	vw	003
2.966	w +	$40\overline{2}, 031$
2.918	vw	013
2.778	vw	$12\overline{3}, 320, 202$
2.720	vw	41 <del>1</del> , 131
2.617	vw	$103, 23\overline{2}$
2.340	vw	$040, 23\overline{3}$
2.271	vw	330
2.242	vw	014, 420
$2 \cdot 171$	vw	141
$2 \cdot 107$	vw	$52\overline{2}, 43\overline{1}, 24\overline{2}$
2.044	vw	104
1.957	vw	$24\overline{3}$
1.903	vww	$34\overline{3}$

Rest too weak

Regnaut, Faugeras, Brut, Helou & Redon, 1958; Rao, Subramaniam & Welch, 1962).

X-ray diffraction data of the compound, given in Table 1, were obtained from patterns taken in a 19 cm Unicam camera of Bradley–Jay type with Cu  $K\alpha$  radiation. The crystal system is monoclinic and unit-cell dimensions are:

 $a = 11.84 \pm 0.005, b = 9.40 \pm 0.004,$  $c = 10.66 \pm 0.005 \text{ Å}; \beta = 120^{\circ} 13';$ 

these are similar to the unit-cell dimensions of lanthanum oxalate decahydrate obtained from single-crystal studies by Victor & McCrone (1952). It therefore appears that  $Pu_2(C_2O_4)_3$ .  $10H_2O$  is isostructural with  $La_2(C_2O_4)_3$ .  $10H_2O$ . The probable space group is  $P2_1/m$ . Only  $PuO_2$  lines were detected when plutonium oxalate was heated to 900 °C.

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