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Pentalanthanum zinc diplumbide, La₅Zn_{1-x}Pb_{2+x} ($x \simeq 0.6$)

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 Key indicators: single-crystal X-ray study; $T = 153$ K; mean $\sigma(\text{Pb-La}) = 0.001$ Å; disorder in main residue; R factor = 0.024; wR factor = 0.058; data-to-parameter ratio = 19.6.

The title non-stoichiometric pentalanthanum zinc diplumbide, La₅Zn_{1-x}Pb_{2+x} ($x \simeq 0.6$), was prepared from the elements in an evacuated silica ampoule. It adopts the Nb₅Sn₂Si-type structure (space group *I4/mcm*, Pearson symbol *tI32*), a ternary ordered superstructure of the W₅Si₃ type. Among the four independent crystallographic positions, three are fully occupied by La (Wyckoff *16k*), La (*4b*), and Pb (*8h*) and one is occupied by a statistical mixture [occupancy ratio 0.394 (12): 0.606 (12)] of Zn and Pb (*4a*). The structure is constructed by face-sharing 10-vertex polyhedra around the unmixed Pb sites. These fragments enclose channels of *trans*-face-sharing tetragonal antiprisms occupied by the disordered Zn and Pb sites.

Related literature

For general background to electronic structure calculations, see: Andersen *et al.* (1986); Becke & Edgecombe (1990); Dronskowski & Blöchl (1993); Lange (1999); Nowak *et al.* (1991). For related structures, see: Aronsson (1955); Oshchapovsky *et al.* (2011a,b, 2012a,b); Stetskiv *et al.* (2012). For isotopic structures, see: Horyn & Lukaszewich (1970).

Experimental

Crystal data

La₅Zn_{0.394}Pb_{2.606} $M_r = 1259.40$ Tetragonal, *I4/mcm* $a = 12.7630$ (18) Å $c = 6.3680$ (13) Å $V = 1037.3$ (3) Å³ $Z = 4$ Mo $K\alpha$ radiation $\mu = 63.04$ mm⁻¹ $T = 153$ K

0.04 × 0.02 × 0.01 mm

Data collection

Bruker P4 CCD diffractometer

Absorption correction: multi-scan

(XSCANS; Bruker, 1999) $T_{\min} =$ 0.332, $T_{\max} = 0.525$

3933 measured reflections

365 independent reflections

352 reflections with $I > 2\sigma(I)$ $R_{\text{int}} = 0.122$

Refinement

 $R[F^2 > 2\sigma(F^2)] = 0.024$ $wR(F^2) = 0.058$ $S = 1.25$

352 reflections

18 parameters

 $\Delta\rho_{\text{max}} = 2.55$ e Å⁻³ $\Delta\rho_{\text{min}} = -1.27$ e Å⁻³
Table 1

 Bond lengths, negative iCOHP^a values and distance contractions in the La₅Zn_{1-x}Pb_{2+x} compound.

Bond	Length (δ in Å)	-iCOHP ^a (eV)	Contraction ^b (%) $\frac{(r_1+r_2)-\delta}{r_1+r_2} \times 100\%$
La1—La1	3.611	0.59	3.4
La1—La1	3.825	0.45	-2.3
La1—La1	3.997	0.40	-6.9
La1—La2	4.088	0.40	-9.3
La1—La1	4.193	0.32	-12.1
La2I—La2	3.184	1.26	14.9
La1—Zn4	3.365	0.26	-5.2
La2—Pb3	3.319	1.10	7.0
La1—Pb3	3.339	1.09	6.5
La1—Pb3	3.484	0.93	2.4
La1—Pb3	3.681	0.72	-3.1
Zn4—Zn4	3.184	-0.49	-19.7

Notes: (a) integrated Crystal Orbital Hamiltonian Population (see Dronskowski & Blöchl, 1993); calculated negative iCOHP values enable qualitative estimation of energies of two-center bonds; (b) based on metallic radii of elements $R(\text{La}) = 1.86$ Å $R(\text{Zn}) = 1.33$ Å and $R(\text{Pb}) = 1.70$ Å.

Data collection: *XSCANS* (Bruker, 1999); cell refinement: *XSCANS*; data reduction: *XSCANS*; program(s) used to solve structure: *SHELXS97* (Sheldrick, 2008); program(s) used to refine structure: *SHELXL97* (Sheldrick, 2008); molecular graphics: *DIAMOND* (Brandenburg, 2006) and *VESTA* (Momma & Izumi, 2008); software used to prepare material for publication: *pubCIF* (Westrip, 2010).

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Supplementary data and figures for this paper are available from the IUCr electronic archives (Reference: HB7160).

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Pentalanthanum zinc diplumbide, $\text{La}_5\text{Zn}_{1-x}\text{Pb}_{2+x}$ ($x \approx 0.6$)**Igor Oshchapovsky, Volodymyr Pavlyuk, Grygoriy Dmytriv and Bernd Harbrecht****S1. Comment**

The title compound, (I), adopts a $\text{Nb}_5\text{Sn}_2\text{Si}$ (Horyn & Lukaszewich, 1970) type structure (space group $I4/mcm$, Pearson symbol $tI32$), a ternary ordered superstructure of the W_5Si_3 type (Aronsson, 1955). A relevant detail of the structure of (I) is presented in Fig. 1.

The coordination polyhedron of the La1 atoms is a 15-vertices polyhedron (CN=15). The nearest neighbours of La2 atoms form 14-vertices polyhedra with CN=14 (see Fig. 1a). The 10-vertices polyhedra around Pb3 can be seen as the major building block of the structure. The face-sharing 10-vertices polyhedra form a three-dimensional framework encasing channels of face-sharing tetragonal antiprisms which in turn accommodate Zn4 and Pb4 with C.N. 10 in a statistical way (see Fig. 1 b, 1c).

This work continues the investigation of $\{\text{La}, \text{Tb}\}\text{-Zn}\text{-}\{\text{Sn}, \text{Pb}\}$ ternary systems. In previous articles the bonding interactions in the compounds LaZn_4 (Oshchapovsky *et al.*, 2012a), LaZn_5 (Oshchapovsky *et al.*, 2012b), $\text{LaZn}_{12.37}$ (Oshchapovsky *et al.*, 2011b) and $\text{TbLi}_{1-x}\text{Zn}_x\text{Sn}_2$ (Stetskiv *et al.*, 2012) were estimated on the basis of electronic structure calculations using the TB-LMTO-ASA program package (Andersen *et al.*, 1986; Nowak *et al.*, 1991) with the additional implementation of two algorithms, the electron localization function (ELF) (Becke & Edgecombe, 1990) and – for quantifying nearest-neighbour bonding interactions – crystal orbital Hamilton population (COHP) (Dronskowski & Blöchl, 1993) analyses. Results arising from these analyses were put in relation to selected crystallographic data of $\text{La}_5\text{Zn}_2\text{Sn}$ (Oshchapovsky *et al.*, 2011a), namely a differential electronic density map obtained from diffraction data.

In all above mentioned compounds bonding interactions evolve similarly. Atoms of the rare-earth elements usually form metallic bonds with a certain ionic component as they donate part of their electrons to other atoms. The atoms of d- and p-metals, namely zinc and tin, accept these electrons used for covalent bonds among each other. This outcome can be easily explained by the higher electronegativity of zinc and tin compared with rare-earth and alkaline metals (Lange, 1999). Judged on the basis of integrated COHP, (iCOHP) values, $R\text{-}\{\text{Zn}, \text{Sn}\}$ bonds are significantly weaker than $\{\text{Zn}, \text{Sn}\}\text{-}\{\text{Zn}, \text{Sn}\}$ bonds. Latter mostly show significant bond length contraction compared to the next contacts in the elemental metals. The differential electronic density map of the structure of $\text{La}_5\text{Zn}_2\text{Sn}$, obtained from experimental diffraction data, indicate donation of electrons to zinc and tin atoms in the similar way as in other compounds, for which electronic structure calculations were performed.

The results of electronic structure calculations for (I) are given in Fig. 2 and 3. As an algorithm for calculating electronic structures for substances with mixed-occupied sites is not implemented in TB-LMTO-ASA package, a fictitious, crystallographically ordered phase was introduced by assuming the statistically occupied Zn/Pb site solely occupied by Zn without change of other crystallographic data.

Similar to $R\text{-Zn}$ binary and $R\text{-Zn-Sn}$ ternary compounds, large values of the electron localization function (ELF) around Pb3 and Zn4 (originally mixed-occupied) are found. In accordance with similar results for previously cited, structurally related compounds, these findings are interpreted as resulting from donation of electrons from lanthanum

atoms to zinc and lead atoms (see Fig. 2a,b).

The values of ELF around Pb3 atoms are significantly larger than around Zn4 atoms, which indicate that bonds between Pb3 and their neighbours should be significantly stronger than bonds between Zn4 atoms and their neighbours. The calculated iCOHP values enable qualitative estimation of energies of two-center bonds and confirm previous assumption. Negative values of iCOHP together with bond lengths and bond contractions are given in Table.

The DOS plot indicates metallic conductivity for the ordered model compound (see Fig. 3a) with small dip at -1 eV relative to Fermi level. Based on a rigid band approximation it can be suggested that these features don't change qualitatively for the title compound. A small dip in the DOS plot near Fermi level could be indirect proof of donation of electrons from lanthanum atoms towards zinc and lead atoms.

The graph negative iCOHP values *versus*. distance contraction shown in Fig. 3 b substantiates the assumption that the polyhedra around Pb3 are indeed the building blocks of the structure if seen as condensed into the given three-dimensional framework by sharing faces. Apparently, this interpretation is not only useful from a crystallographic point of view but also supported by the hierarchy of bonding interactions in the solid: La—Pb bonds are stronger and show larger distance contractions than most La—La bonds. An exception is the particularly strong La^{2IX}—La^{2X} bond enhancing the stability of the framework by connecting adjacent building blocks.

S2. Experimental

A small irregular grey single-crystal of the title compound of suitable quality was isolated from an alloy with composition La₇ZnPb₂ prepared in the course of a systematic investigation of the ternary La—Zn—Pb system. The preparation process was according to that described for La₅Zn₂Sn (Oshchapovsky *et al.*, 2011a).

The sample was prepared by melting pieces of pure metals (99.9% La, 99.999% Zn, 99.99% Pb) in an evacuated silica ampoule in a resistance furnace with subsequent annealing at 600°C for 30 days. Final phase analysis revealed the presence of the title compound in samples with composition La₇ZnPb₂ together with other lanthanum-rich ternary alloys. The latter samples, however, had not reached completely the liquid state, since the use of silica ampoules constraints the maximum temperature to 900°C. This limitation may hamper complete equilibration of the sample.

S3. Refinement

The fractional site occupancies of Pb4 and Zn4 were constrained to sum to unity.

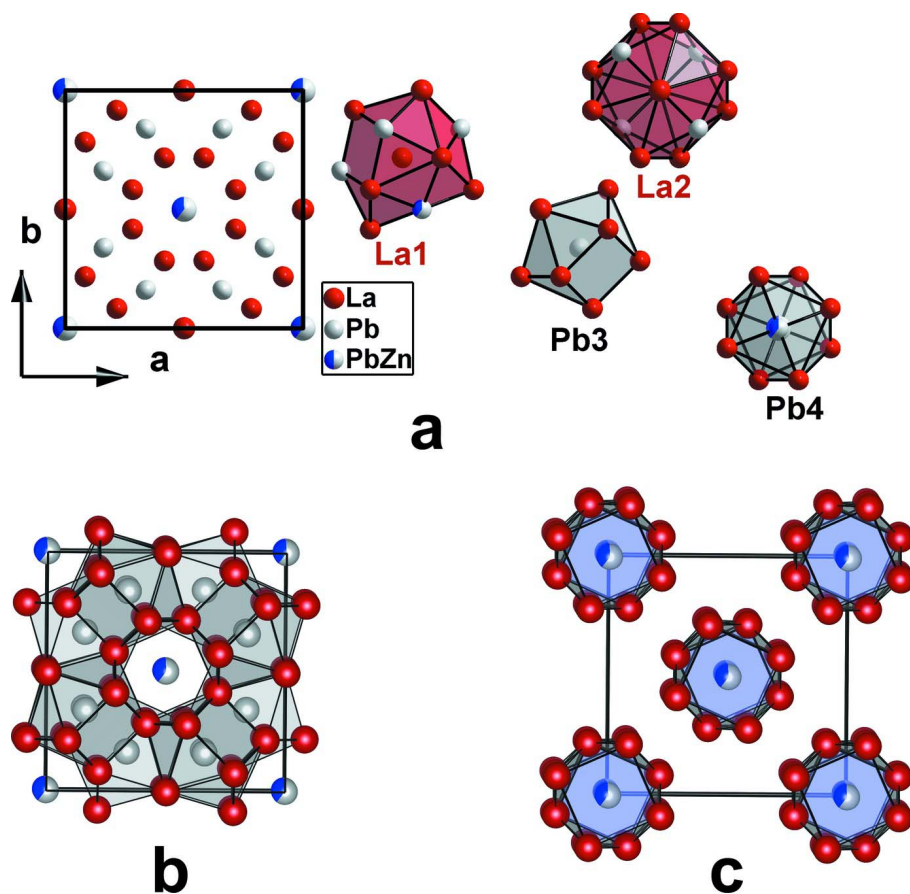


Figure 1

Details of the crystal structure of $\text{La}_5\text{Zn}_{1-x}\text{Pb}_{2+x}$: Projection of the unit cell and coordination polyhedra (a); building blocks (b); channels in the structure (c).

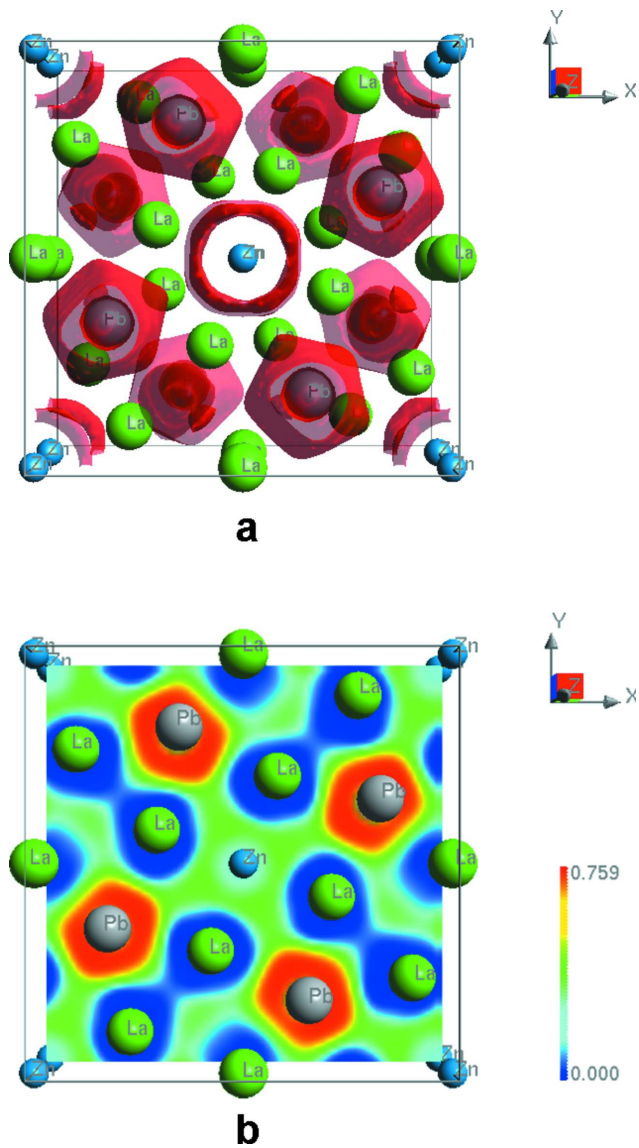


Figure 2

Results of ELF calculations for the $\text{La}_5\text{Zn}_{1-x}\text{Pb}_{2+x}$ compound: Isosurfaces drawn at 0.45 level (a); section [001] drawn at $z=1/2$ (b).

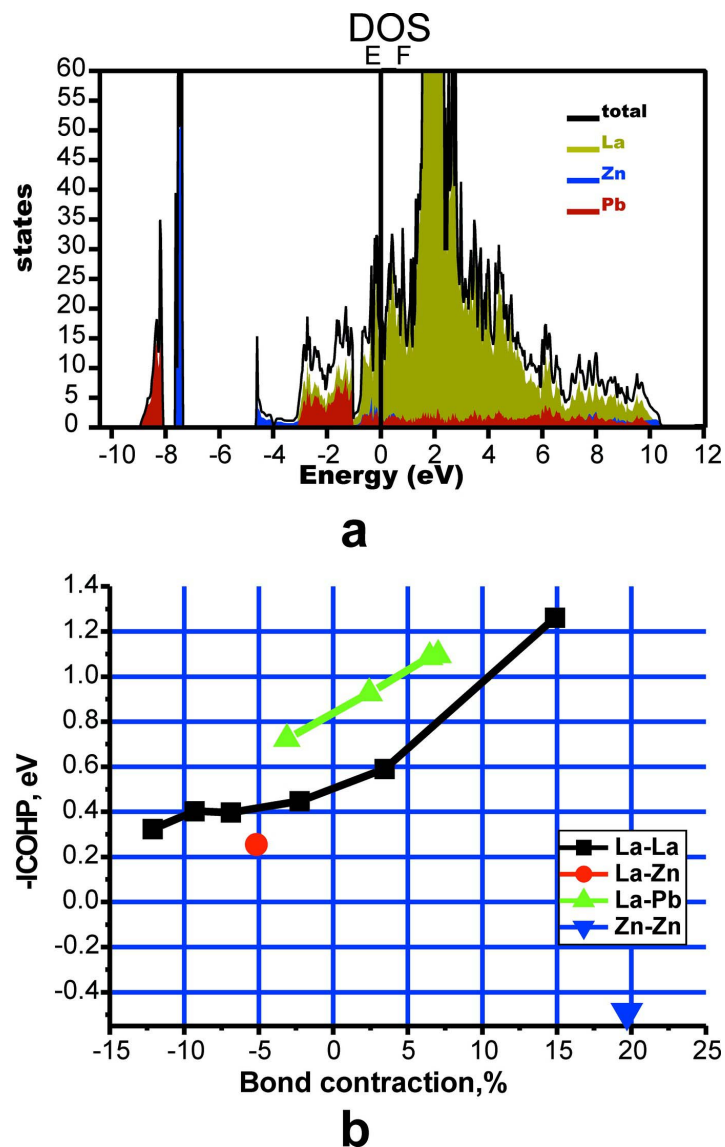


Figure 3

Density of states plot (a); Comparison of bond contractions and -iCOHP values in the title compound (b).

Pentalanthanum zinc diplumbide

Crystal data

$\text{La}_5\text{Pb}_{2.606}\text{Zn}_{0.394}$

$M_r = 1259.40$

Tetragonal, $I4/mcm$

Hall symbol: $-I\ 4\ 2c$

$a = 12.7630\ (18)\ \text{\AA}$

$c = 6.3680\ (13)\ \text{\AA}$

$V = 1037.3\ (3)\ \text{\AA}^3$

$Z = 4$

$F(000) = 2041.6$

$D_x = 8.068\ \text{Mg m}^{-3}$

Mo $K\alpha$ radiation, $\lambda = 0.71073\ \text{\AA}$

Cell parameters from 3933 reflections

$\theta = 4.5\text{--}27.9^\circ$

$\mu = 63.04\ \text{mm}^{-1}$

$T = 153\ \text{K}$

Irregularly shaped, metallic grey

$0.04 \times 0.02 \times 0.01\ \text{mm}$

Data collection

Bruker P4 CCD diffractometer	3933 measured reflections
Radiation source: fine-focus sealed tube	365 independent reflections
Graphite monochromator	352 reflections with $I > 2\sigma(I)$
ω scans	$R_{\text{int}} = 0.122$
Absorption correction: multi-scan (XSCANS; Bruker, 1999)	$\theta_{\text{max}} = 27.9^\circ$, $\theta_{\text{min}} = 4.5^\circ$
$T_{\text{min}} = 0.332$, $T_{\text{max}} = 0.525$	$h = -16 \rightarrow 16$
	$k = -15 \rightarrow 16$
	$l = -8 \rightarrow 8$

Refinement

Refinement on F^2	Secondary atom site location: difference Fourier map
Least-squares matrix: full	$w = 1/[\sigma^2(F_o^2) + 23.7905P]$
$R[F^2 > 2\sigma(F^2)] = 0.024$	where $P = (F_o^2 + 2F_c^2)/3$
$wR(F^2) = 0.058$	$(\Delta/\sigma)_{\text{max}} = 0.012$
$S = 1.25$	$\Delta\rho_{\text{max}} = 2.55 \text{ e } \text{\AA}^{-3}$
352 reflections	$\Delta\rho_{\text{min}} = -1.27 \text{ e } \text{\AA}^{-3}$
18 parameters	Extinction correction: SHELXL97 (Sheldrick, 2008), $F_c^* = kFc[1 + 0.001x Fc^2 \lambda^3 / \sin(2\theta)]^{-1/4}$
0 restraints	Extinction coefficient: 3948 (389)
Primary atom site location: structure-invariant direct methods	

Special details

Geometry. All e.s.d.'s (except the e.s.d. in the dihedral angle between two l.s. planes) are estimated using the full covariance matrix. The cell e.s.d.'s are taken into account individually in the estimation of e.s.d.'s in distances, angles and torsion angles; correlations between e.s.d.'s in cell parameters are only used when they are defined by crystal symmetry. An approximate (isotropic) treatment of cell e.s.d.'s is used for estimating e.s.d.'s involving l.s. planes.

Refinement. Refinement of F^2 against ALL reflections. The weighted R -factor wR and goodness of fit S are based on F^2 , conventional R -factors R are based on F , with F set to zero for negative F^2 . The threshold expression of $F^2 > \sigma(F^2)$ is used only for calculating R -factors(gt) etc. and is not relevant to the choice of reflections for refinement. R -factors based on F^2 are statistically about twice as large as those based on F , and R -factors based on ALL data will be even larger.

Fractional atomic coordinates and isotropic or equivalent isotropic displacement parameters (\AA^2)

	x	y	z	$U_{\text{iso}}^*/U_{\text{eq}}$	Occ. (<1)
La1	0.08312 (9)	0.21698 (9)	0.0000	0.0229 (4)	
La2	0.0000	0.5000	0.2500	0.0188 (6)	
Pb3	0.16124 (5)	0.66124 (5)	0.0000	0.0187 (4)	
Pb4	0.0000	0.0000	0.2500	0.0231 (9)	0.606 (12)
Zn4	0.0000	0.0000	0.2500	0.0231 (9)	0.394 (12)

Atomic displacement parameters (\AA^2)

	U^{11}	U^{22}	U^{33}	U^{12}	U^{13}	U^{23}
La1	0.0150 (6)	0.0150 (6)	0.0386 (9)	0.0005 (4)	0.000	0.000
La2	0.0171 (6)	0.0171 (6)	0.0222 (12)	0.000	0.000	0.000
Pb3	0.0137 (4)	0.0137 (4)	0.0287 (7)	-0.0013 (3)	0.000	0.000
Pb4	0.0142 (8)	0.0142 (8)	0.0410 (16)	0.000	0.000	0.000
Zn4	0.0142 (8)	0.0142 (8)	0.0410 (16)	0.000	0.000	0.000

Geometric parameters (Å, °)

La1—Pb3 ⁱ	3.3394 (14)	La2—La1 ⁱⁱⁱ	4.0875 (12)
La1—Zn4 ⁱⁱ	3.3658 (11)	Pb3—La2 ⁱⁱⁱ	3.3173 (9)
La1—Pb4 ⁱⁱ	3.3658 (11)	Pb3—La1 ^{xiv}	3.3394 (14)
La1—Pb4	3.3658 (11)	Pb3—La1 ^{xv}	3.3394 (14)
La1—Pb3 ⁱⁱⁱ	3.4846 (12)	Pb3—La1 ⁱⁱⁱ	3.4846 (12)
La1—La1 ^{iv}	3.608 (2)	Pb3—La1 ^{iv}	3.4846 (12)
La1—Pb3 ^v	3.6807 (8)	Pb3—La1 ^{xii}	3.6807 (8)
La1—Pb3 ^{vi}	3.6807 (8)	Pb3—La1 ^{xvi}	3.6807 (8)
La1—La1 ^{vii}	3.8261 (13)	Pb3—La1 ^{xi}	3.6807 (8)
La1—La1 ^{viii}	3.8261 (13)	Pb3—La1 ^{xvii}	3.6807 (8)
La1—La1 ^{ix}	3.9969 (14)	Pb4—Zn4 ^{xviii}	3.1840 (7)
La2—La2 ^x	3.1840 (7)	Pb4—Zn4 ⁱⁱ	3.1840 (6)
La2—La2 ⁱⁱⁱ	3.1840 (6)	Pb4—Pb4 ^{xviii}	3.1840 (7)
La2—Pb3 ⁱⁱⁱ	3.3173 (9)	Pb4—Pb4 ⁱⁱ	3.1840 (6)
La2—Pb3	3.3173 (9)	Pb4—La1 ^{xix}	3.3658 (11)
La2—Pb3 ^v	3.3173 (9)	Pb4—La1 ⁱⁱ	3.3658 (11)
La2—Pb3 ^{xi}	3.3173 (9)	Pb4—La1 ^{viii}	3.3658 (11)
La2—La1 ^{xii}	4.0875 (12)	Pb4—La1 ^{xx}	3.3658 (11)
La2—La1 ^{xi}	4.0875 (12)	Pb4—La1 ^{xxi}	3.3658 (11)
La2—La1 ^{xiii}	4.0875 (12)	Pb4—La1 ^{xxii}	3.3658 (11)
La2—La1 ^v	4.0875 (12)	Pb4—La1 ^{xxiii}	3.3658 (11)
Pb3 ⁱ —La1—Zn4 ⁱⁱ	97.62 (3)	La1 ^{xii} —La2—La1	149.92 (3)
Pb3 ⁱ —La1—Pb4 ⁱⁱ	97.62 (3)	La1 ^{xi} —La2—La1	98.725 (6)
Zn4 ⁱⁱ —La1—Pb4 ⁱⁱ	0.0	La1 ^{xiii} —La2—La1	107.88 (3)
Pb3 ⁱ —La1—Pb4	97.62 (3)	La1 ^v —La2—La1	98.725 (6)
Zn4 ⁱⁱ —La1—Pb4	56.46 (2)	La1 ⁱⁱⁱ —La2—La1	134.156 (16)
Pb4 ⁱⁱ —La1—Pb4	56.46 (2)	La2 ⁱⁱⁱ —Pb3—La2	57.36 (2)
Pb3 ⁱ —La1—Pb3 ⁱⁱⁱ	165.81 (4)	La2 ⁱⁱⁱ —Pb3—La1 ^{xiv}	137.584 (15)
Zn4 ⁱⁱ —La1—Pb3 ⁱⁱⁱ	94.87 (3)	La2—Pb3—La1 ^{xiv}	137.584 (15)
Pb4 ⁱⁱ —La1—Pb3 ⁱⁱⁱ	94.87 (3)	La2 ⁱⁱⁱ —Pb3—La1 ^{xv}	137.584 (15)
Pb4—La1—Pb3 ⁱⁱⁱ	94.87 (3)	La2—Pb3—La1 ^{xv}	137.584 (15)
Pb3 ⁱ —La1—La1 ^{iv}	57.30 (2)	La1 ^{xiv} —Pb3—La1 ^{xv}	65.40 (4)
Zn4 ⁱⁱ —La1—La1 ^{iv}	143.576 (15)	La2 ⁱⁱⁱ —Pb3—La1 ⁱⁱⁱ	73.83 (2)
Pb4 ⁱⁱ —La1—La1 ^{iv}	143.576 (15)	La2—Pb3—La1 ⁱⁱⁱ	73.83 (2)
Pb4—La1—La1 ^{iv}	143.576 (15)	La1 ^{xiv} —Pb3—La1 ⁱⁱⁱ	75.81 (4)
Pb3 ⁱⁱⁱ —La1—La1 ^{iv}	108.51 (2)	La1 ^{xv} —Pb3—La1 ⁱⁱⁱ	141.21 (2)
Pb3 ⁱ —La1—Pb3 ^v	79.94 (3)	La2 ⁱⁱⁱ —Pb3—La1 ^{iv}	73.83 (2)
Zn4 ⁱⁱ —La1—Pb3 ^v	147.35 (3)	La2—Pb3—La1 ^{iv}	73.83 (2)
Pb4 ⁱⁱ —La1—Pb3 ^v	147.35 (3)	La1 ^{xiv} —Pb3—La1 ^{iv}	141.21 (2)
Pb4—La1—Pb3 ^v	91.356 (15)	La1 ^{xv} —Pb3—La1 ^{iv}	75.81 (4)
Pb3 ⁱⁱⁱ —La1—Pb3 ^v	93.10 (3)	La1 ⁱⁱⁱ —Pb3—La1 ^{iv}	142.98 (5)
La1 ^{iv} —La1—Pb3 ^v	60.650 (17)	La2 ⁱⁱⁱ —Pb3—La1 ^{xii}	120.60 (3)
Pb3 ⁱ —La1—Pb3 ^{vi}	79.94 (3)	La2—Pb3—La1 ^{xii}	71.26 (2)
Zn4 ⁱⁱ —La1—Pb3 ^{vi}	91.356 (15)	La1 ^{xiv} —Pb3—La1 ^{xii}	69.21 (3)
Pb4 ⁱⁱ —La1—Pb3 ^{vi}	91.356 (15)	La1 ^{xv} —Pb3—La1 ^{xii}	100.06 (3)

Pb4—La1—Pb3 ^{vi}	147.35 (3)	La1 ⁱⁱⁱ —Pb3—La1 ^{xii}	64.48 (2)
Pb3 ⁱⁱⁱ —La1—Pb3 ^{vi}	93.10 (3)	La1 ^{iv} —Pb3—La1 ^{xii}	119.919 (19)
La1 ^{iv} —La1—Pb3 ^{vi}	60.650 (17)	La2 ⁱⁱⁱ —Pb3—La1 ^{xvi}	71.26 (2)
Pb3 ^v —La1—Pb3 ^{vi}	119.78 (3)	La2—Pb3—La1 ^{xvi}	120.60 (3)
Pb3 ⁱ —La1—La1 ^{vii}	122.81 (3)	La1 ^{xiv} —Pb3—La1 ^{xvi}	69.21 (3)
Zn4 ⁱⁱ —La1—La1 ^{vii}	55.363 (14)	La1 ^{xv} —Pb3—La1 ^{xvi}	100.06 (3)
Pb4 ⁱⁱ —La1—La1 ^{vii}	55.363 (14)	La1 ⁱⁱⁱ —Pb3—La1 ^{xvi}	64.48 (2)
Pb4—La1—La1 ^{vii}	102.64 (3)	La1 ^{iv} —Pb3—La1 ^{xvi}	119.919 (19)
Pb3 ⁱⁱⁱ —La1—La1 ^{vii}	60.24 (3)	La1 ^{xii} —Pb3—La1 ^{xvi}	119.78 (3)
La1 ^{iv} —La1—La1 ^{vii}	113.086 (18)	La2 ⁱⁱⁱ —Pb3—La1 ^{xi}	120.60 (3)
Pb3 ^v —La1—La1 ^{vii}	150.47 (3)	La2—Pb3—La1 ^{xi}	71.26 (2)
Pb3 ^{vi} —La1—La1 ^{vii}	55.274 (16)	La1 ^{xiv} —Pb3—La1 ^{xi}	100.06 (3)
Pb3 ⁱ —La1—La1 ^{viii}	122.81 (3)	La1 ^{xv} —Pb3—La1 ^{xi}	69.21 (3)
Zn4 ⁱⁱ —La1—La1 ^{viii}	102.64 (3)	La1 ⁱⁱⁱ —Pb3—La1 ^{xi}	119.919 (19)
Pb4 ⁱⁱ —La1—La1 ^{viii}	102.64 (3)	La1 ^{iv} —Pb3—La1 ^{xi}	64.48 (2)
Pb4—La1—La1 ^{viii}	55.363 (14)	La1 ^{xii} —Pb3—La1 ^{xi}	58.70 (3)
Pb3 ⁱⁱⁱ —La1—La1 ^{viii}	60.24 (3)	La1 ^{xvi} —Pb3—La1 ^{xi}	167.71 (4)
La1 ^{iv} —La1—La1 ^{viii}	113.086 (18)	La2 ⁱⁱⁱ —Pb3—La1 ^{xvii}	71.26 (2)
Pb3 ^v —La1—La1 ^{viii}	55.274 (16)	La2—Pb3—La1 ^{xvii}	120.60 (3)
Pb3 ^{vi} —La1—La1 ^{viii}	150.47 (3)	La1 ^{xiv} —Pb3—La1 ^{xvii}	100.06 (3)
La1 ^{vii} —La1—La1 ^{viii}	112.64 (6)	La1 ^{xv} —Pb3—La1 ^{xvii}	69.21 (3)
Pb3 ⁱ —La1—La1 ^{ix}	59.42 (3)	La1 ⁱⁱⁱ —Pb3—La1 ^{xvii}	119.919 (19)
Zn4 ⁱⁱ —La1—La1 ^{ix}	53.576 (15)	La1 ^{iv} —Pb3—La1 ^{xvii}	64.48 (2)
Pb4 ⁱⁱ —La1—La1 ^{ix}	53.576 (15)	La1 ^{xii} —Pb3—La1 ^{xvii}	167.71 (4)
Pb4—La1—La1 ^{ix}	99.20 (3)	La1 ^{xvi} —Pb3—La1 ^{xvii}	58.70 (3)
Pb3 ⁱⁱⁱ —La1—La1 ^{ix}	124.98 (2)	La1 ^{xi} —Pb3—La1 ^{xvii}	119.78 (3)
La1 ^{iv} —La1—La1 ^{ix}	90.0	Zn4 ^{xviii} —Pb4—Zn4 ⁱⁱ	180.0
Pb3 ^v —La1—La1 ^{ix}	138.92 (3)	Zn4 ^{xviii} —Pb4—Pb4 ^{xviii}	0.0
Pb3 ^{vi} —La1—La1 ^{ix}	51.362 (18)	Zn4 ⁱⁱ —Pb4—Pb4 ^{xviii}	180.0
La1 ^{vii} —La1—La1 ^{ix}	64.79 (2)	Zn4 ^{xviii} —Pb4—Pb4 ⁱⁱ	180.0
La1 ^{viii} —La1—La1 ^{ix}	154.150 (9)	Zn4 ⁱⁱ —Pb4—Pb4 ⁱⁱ	0.0
La2 ^x —La2—La2 ⁱⁱⁱ	180.0	Pb4 ^{xviii} —Pb4—Pb4 ⁱⁱ	180.0
La2 ^x —La2—Pb3 ⁱⁱⁱ	118.679 (10)	Zn4 ^{xviii} —Pb4—La1 ^{xix}	61.771 (11)
La2 ⁱⁱⁱ —La2—Pb3 ⁱⁱⁱ	61.321 (10)	Zn4 ⁱⁱ —Pb4—La1 ^{xix}	118.229 (11)
La2 ^x —La2—Pb3	118.679 (10)	Pb4 ^{xviii} —Pb4—La1 ^{xix}	61.771 (11)
La2 ⁱⁱⁱ —La2—Pb3	61.321 (10)	Pb4 ⁱⁱ —Pb4—La1 ^{xix}	118.229 (11)
Pb3 ⁱⁱⁱ —La2—Pb3	122.64 (2)	Zn4 ^{xviii} —Pb4—La1 ⁱⁱ	118.229 (11)
La2 ^x —La2—Pb3 ^v	61.321 (10)	Zn4 ⁱⁱ —Pb4—La1 ⁱⁱ	61.771 (11)
La2 ⁱⁱⁱ —La2—Pb3 ^v	118.679 (10)	Pb4 ^{xviii} —Pb4—La1 ⁱⁱ	118.229 (11)
Pb3 ⁱⁱⁱ —La2—Pb3 ^v	103.315 (9)	Pb4 ⁱⁱ —Pb4—La1 ⁱⁱ	61.771 (11)
Pb3—La2—Pb3 ^v	103.315 (9)	La1 ^{xix} —Pb4—La1 ⁱⁱ	137.93 (4)
La2 ^x —La2—Pb3 ^{xi}	61.321 (10)	Zn4 ^{xviii} —Pb4—La1 ^{viii}	61.771 (11)
La2 ⁱⁱⁱ —La2—Pb3 ^{xi}	118.679 (10)	Zn4 ⁱⁱ —Pb4—La1 ^{viii}	118.229 (11)
Pb3 ⁱⁱⁱ —La2—Pb3 ^{xi}	103.315 (9)	Pb4 ^{xviii} —Pb4—La1 ^{viii}	61.771 (11)
Pb3—La2—Pb3 ^{xi}	103.315 (9)	Pb4 ⁱⁱ —Pb4—La1 ^{viii}	118.229 (11)
Pb3 ^v —La2—Pb3 ^{xi}	122.64 (2)	La1 ^{xix} —Pb4—La1 ^{viii}	77.072 (9)
La2 ^x —La2—La1 ^{xii}	67.078 (8)	La1 ⁱⁱ —Pb4—La1 ^{viii}	143.26 (4)
La2 ⁱⁱⁱ —La2—La1 ^{xii}	112.922 (8)	Zn4 ^{xviii} —Pb4—La1	118.229 (11)

Pb3 ⁱⁱⁱ —La2—La1 ^{xii}	153.655 (15)	Zn4 ⁱⁱ —Pb4—La1	61.771 (11)
Pb3—La2—La1 ^{xii}	58.513 (15)	Pb4 ^{xviii} —Pb4—La1	118.229 (11)
Pb3 ^v —La2—La1 ^{xii}	101.555 (15)	Pb4 ⁱⁱ —Pb4—La1	61.771 (11)
Pb3 ^{xi} —La2—La1 ^{xii}	54.961 (15)	La1 ^{xix} —Pb4—La1	72.85 (3)
La2 ^x —La2—La1 ^{xi}	67.078 (8)	La1 ⁱⁱ —Pb4—La1	123.54 (2)
La2 ⁱⁱⁱ —La2—La1 ^{xi}	112.922 (8)	La1 ^{viii} —Pb4—La1	69.27 (3)
Pb3 ⁱⁱⁱ —La2—La1 ^{xi}	153.655 (15)	Zn4 ^{xviii} —Pb4—La1 ^{xx}	118.229 (11)
Pb3—La2—La1 ^{xi}	58.513 (15)	Zn4 ⁱⁱ —Pb4—La1 ^{xx}	61.771 (11)
Pb3 ^v —La2—La1 ^{xi}	54.961 (15)	Pb4 ^{xviii} —Pb4—La1 ^{xx}	118.229 (11)
Pb3 ^{xi} —La2—La1 ^{xi}	101.555 (15)	Pb4 ⁱⁱ —Pb4—La1 ^{xx}	61.771 (11)
La1 ^{xii} —La2—La1 ^{xi}	52.38 (3)	La1 ^{xix} —Pb4—La1 ^{xx}	69.27 (3)
La2 ^x —La2—La1 ^{xiii}	112.922 (8)	La1 ⁱⁱ —Pb4—La1 ^{xx}	77.072 (9)
La2 ⁱⁱⁱ —La2—La1 ^{xiii}	67.078 (8)	La1 ^{viii} —Pb4—La1 ^{xx}	137.93 (4)
Pb3 ⁱⁱⁱ —La2—La1 ^{xiii}	54.961 (15)	La1—Pb4—La1 ^{xx}	77.072 (9)
Pb3—La2—La1 ^{xiii}	101.555 (15)	Zn4 ^{xviii} —Pb4—La1 ^{xxi}	118.229 (11)
Pb3 ^v —La2—La1 ^{xiii}	153.655 (15)	Zn4 ⁱⁱ —Pb4—La1 ^{xxi}	61.771 (11)
Pb3 ^{xi} —La2—La1 ^{xiii}	58.513 (15)	Pb4 ^{xviii} —Pb4—La1 ^{xxi}	118.229 (11)
La1 ^{xii} —La2—La1 ^{xiii}	98.725 (6)	Pb4 ⁱⁱ —Pb4—La1 ^{xxi}	61.771 (11)
La1 ^{xi} —La2—La1 ^{xiii}	149.92 (3)	La1 ^{xix} —Pb4—La1 ^{xxi}	143.26 (4)
La2 ^x —La2—La1 ^v	67.078 (8)	La1 ⁱⁱ —Pb4—La1 ^{xxi}	77.072 (9)
La2 ⁱⁱⁱ —La2—La1 ^v	112.922 (8)	La1 ^{viii} —Pb4—La1 ^{xxi}	72.85 (3)
Pb3 ⁱⁱⁱ —La2—La1 ^v	58.513 (15)	La1—Pb4—La1 ^{xxi}	77.072 (10)
Pb3—La2—La1 ^v	153.655 (15)	La1 ^{xx} —Pb4—La1 ^{xxi}	123.54 (2)
Pb3 ^v —La2—La1 ^v	101.555 (15)	Zn4 ^{xviii} —Pb4—La1 ^{xxii}	61.771 (11)
Pb3 ^{xi} —La2—La1 ^v	54.961 (15)	Zn4 ⁱⁱ —Pb4—La1 ^{xxii}	118.229 (11)
La1 ^{xii} —La2—La1 ^v	107.88 (3)	Pb4 ^{xviii} —Pb4—La1 ^{xxii}	61.771 (11)
La1 ^{xi} —La2—La1 ^v	134.156 (16)	Pb4 ⁱⁱ —Pb4—La1 ^{xxii}	118.229 (11)
La1 ^{xiii} —La2—La1 ^v	55.81 (2)	La1 ^{xix} —Pb4—La1 ^{xxii}	123.54 (2)
La2 ^x —La2—La1 ⁱⁱⁱ	112.922 (8)	La1 ⁱⁱ —Pb4—La1 ^{xxii}	72.85 (3)
La2 ⁱⁱⁱ —La2—La1 ⁱⁱⁱ	67.078 (8)	La1 ^{viii} —Pb4—La1 ^{xxii}	77.072 (9)
Pb3 ⁱⁱⁱ —La2—La1 ⁱⁱⁱ	101.555 (15)	La1—Pb4—La1 ^{xxii}	137.93 (4)
Pb3—La2—La1 ⁱⁱⁱ	54.961 (15)	La1 ^{xx} —Pb4—La1 ^{xxii}	143.26 (4)
Pb3 ^v —La2—La1 ⁱⁱⁱ	153.655 (15)	La1 ^{xxi} —Pb4—La1 ^{xxii}	69.27 (3)
Pb3 ^{xi} —La2—La1 ⁱⁱⁱ	58.513 (15)	Zn4 ^{xviii} —Pb4—La1 ^{xxiii}	61.771 (11)
La1 ^{xii} —La2—La1 ⁱⁱⁱ	55.81 (2)	Zn4 ⁱⁱ —Pb4—La1 ^{xxiii}	118.229 (11)
La1 ^{xi} —La2—La1 ⁱⁱⁱ	98.725 (6)	Pb4 ^{xviii} —Pb4—La1 ^{xxiii}	61.771 (11)
La1 ^{xiii} —La2—La1 ⁱⁱⁱ	52.38 (3)	Pb4 ⁱⁱ —Pb4—La1 ^{xxiii}	118.229 (11)
La1 ^v —La2—La1 ⁱⁱⁱ	98.725 (6)	La1 ^{xix} —Pb4—La1 ^{xxiii}	77.072 (9)
La2 ^x —La2—La1	112.922 (8)	La1 ⁱⁱ —Pb4—La1 ^{xxiii}	69.27 (3)
La2 ⁱⁱⁱ —La2—La1	67.078 (8)	La1 ^{viii} —Pb4—La1 ^{xxiii}	123.54 (2)
Pb3 ⁱⁱⁱ —La2—La1	54.961 (15)	La1—Pb4—La1 ^{xxiii}	143.26 (4)
Pb3—La2—La1	101.555 (15)	La1 ^{xx} —Pb4—La1 ^{xxiii}	72.85 (3)
Pb3 ^v —La2—La1	58.513 (15)	La1 ^{xxi} —Pb4—La1 ^{xxiii}	137.93 (4)
Pb3 ^{xi} —La2—La1	153.655 (15)	La1 ^{xxii} —Pb4—La1 ^{xxiii}	77.072 (9)

Symmetry codes: (i) $-y+1, x, z$; (ii) $-x, -y, -z$; (iii) $-x, -y+1, -z$; (iv) $-y+1/2, -x+1/2, z$; (v) $y-1/2, -x+1/2, -z+1/2$; (vi) $y-1/2, -x+1/2, -z-1/2$; (vii) $-x, y, -z-1/2$; (viii) $-x, y, -z+1/2$; (ix) $y, x, -z-1/2$; (x) $-x, -y+1, -z+1$; (xi) $-y+1/2, x+1/2, z+1/2$; (xii) $x, -y+1, z+1/2$; (xiii) $y-1/2, x+1/2, -z$; (xiv) $y, -x+1, -z$; (xv) $-x+1/2, y+1/2, -z$; (xvi) $x, -y+1, z-1/2$; (xvii) $-y+1/2, x+1/2, z-1/2$; (xviii) $-x, -y, -z+1$; (xix) $y, x, -z+1/2$; (xx) $y, -x, -z$; (xxi) $-y, x, z$; (xxii) $-y, -x, z+1/2$; (xxiii) $x, -y, z+1/2$.