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## Structure Reports

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## $\mu$ -Oxalato- $\kappa^4 O^1, O^2:O^1', O^2'$ -bis[aqua-(2,2'-bipyridine- $\kappa N$ )(nitrate- $\kappa^2 O, O')$ -lead(II)]

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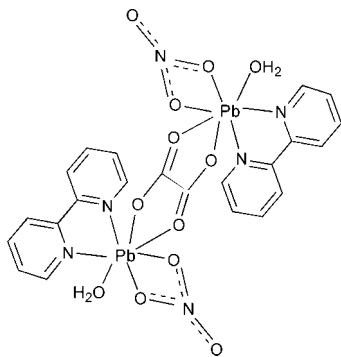
Received 25 July 2012; accepted 22 September 2012

Key indicators: single-crystal X-ray study;  $T = 296$  K; mean  $\sigma(C-C) = 0.011$  Å;  $R$  factor = 0.031;  $wR$  factor = 0.082; data-to-parameter ratio = 12.3.

The title compound,  $[Pb_2(C_2O_4)(NO_3)_2(C_{10}H_8N_2)_2(H_2O)_2]$ , was synthesized hydrothermally. The binuclear complex molecule is centrosymmetric, the inversion centre being located at the mid-point of the oxalate C—C bond. The  $Pb^{II}$  ion is heptacoordinated by the O atom of one water molecule, two oxalate O atoms, two nitrate O atoms and two 2,2'-bipyridine N atoms, forming an irregular coordination environment. Intermolecular O—H...O hydrogen bonds between water molecules and oxalate and nitrate ions result in the formation of layers parallel to (010).  $\pi$ – $\pi$  interactions between pyridine rings in adjacent layers, with centroid–centroid distances of 3.584 (2) Å, stabilize the structural set-up.

### Related literature

For general background to this class of compounds, see: Fan & Zhu (2006); Hamilton *et al.* (2004); Hagrman & Zubieta (2000); Li *et al.* (2002).



### Experimental

#### Crystal data

$[Pb_2(C_2O_4)(NO_3)_2(C_{10}H_8N_2)_2 \cdot (H_2O)_2]$   
 $M_r = 974.82$   
 Monoclinic,  $P2_1/c$   
 $a = 9.5791$  (19) Å  
 $b = 20.6330$  (14) Å  
 $c = 6.7649$  (15) Å  
 $\beta = 91.687$  (1)°  
 $V = 1336.5$  (4) Å<sup>3</sup>  
 $Z = 2$   
 Mo  $K\alpha$  radiation  
 $\mu = 12.66$  mm<sup>-1</sup>  
 $T = 296$  K  
 $0.29 \times 0.28 \times 0.26$  mm

#### Data collection

Bruker SMART CCD diffractometer  
 Absorption correction: multi-scan (SADABS; Bruker, 2003)  
 $T_{min} = 0.120$ ,  $T_{max} = 0.137$   
 7096 measured reflections  
 2336 independent reflections  
 2082 reflections with  $I > 2\sigma(I)$   
 $R_{int} = 0.038$

#### Refinement

$R[F^2 > 2\sigma(F^2)] = 0.031$   
 $wR(F^2) = 0.082$   
 $S = 1.07$   
 2336 reflections  
 190 parameters  
 H-atom parameters constrained  
 $\Delta\rho_{max} = 2.68$  e Å<sup>-3</sup>  
 $\Delta\rho_{min} = -1.19$  e Å<sup>-3</sup>

Table 1

Hydrogen-bond geometry (Å, °).

$D-H\cdots A$	$D-H$	$H\cdots A$	$D\cdots A$	$D-H\cdots A$
$O7-H7A\cdots O5^i$	0.85	2.23	2.875 (7)	133
$O7-H7B\cdots O3^{ii}$	0.85	2.16	2.912 (8)	148

Symmetry codes: (i)  $x, y, z + 1$ ; (ii)  $-x + 2, -y + 1, -z + 2$ .

Data collection: SMART (Bruker, 2003); cell refinement: SAINT (Bruker, 2003); data reduction: SAINT; program(s) used to solve structure: SHELXS97 (Sheldrick, 2008); program(s) used to refine structure: SHELXL97 (Sheldrick, 2008); molecular graphics: DIAMOND (Brandenburg, 2009); software used to prepare material for publication: SHELXTL (Sheldrick, 2008).

This work was supported by the Innovation Project of Guangxi University for Nationalities.

Supplementary data and figures for this paper are available from the IUCr electronic archives (Reference: WM2667).

### References

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## supplementary materials

*Acta Cryst.* (2012). E68, m1310 [doi:10.1107/S1600536812040196]

 **$\mu$ -Oxalato- $\kappa^4O^1, O^2:O^1', O^2'$ -bis[aqua(2,2'-bipyridine- $\kappa N$ )(nitrate- $\kappa^2O, O'$ )lead(II)]**

Gang-Hong Pan, Jin-Niu Tang, Zhong-Jing Huang, Long Li and Chun-Mei Zhang

**Comment**

Complexes containing Pb<sup>II</sup> (Fan & Zhu *et al.*, 2006) are interesting because of the variety of their structures and their potential applications, especially in environmental protection, e.g heavy metal removal. It is known that the introduction of chelating ligands such as 2,2'-bipyridine causes the passivation of metal sites via the N donors of the organic groups and may induce new structural evolution (Hamilton *et al.*, 2004; Hagrman & Zubieta, 2000; Li *et al.*, 2002).

In the title binuclear lead(II) compound, [Pb<sub>2</sub>(C<sub>2</sub>O<sub>4</sub>)(NO<sub>3</sub>)<sub>2</sub>(H<sub>2</sub>O)<sub>2</sub>(C<sub>10</sub>H<sub>8</sub>N<sub>2</sub>)<sub>2</sub>], a centrosymmetric molecule is present, with the centre of symmetry at the mid-point of the C—C oxalate bond (Fig. 1). The Pb<sup>II</sup> ion is hepta-coordinated in an irregular fashion by one water molecule, two nitrate oxygen atoms, two oxalate oxygen atoms, and two nitrogen atoms from 2,2'-bipyridine. The supramolecular assembly in the title compound is completed by O—H...O hydrogen bonds between the coordinating water molecules and oxalate and nitrate O atoms (Table 1), resulting in the formation of layers parallel to (010) (Fig. 2). The structure is further extended by  $\pi$ — $\pi$  stacking interactions between 2,2'-bipyridine molecules of adjacent layers. They overlap with a centroid-to-centroid distance of 3.584 (2) Å.

**Experimental**

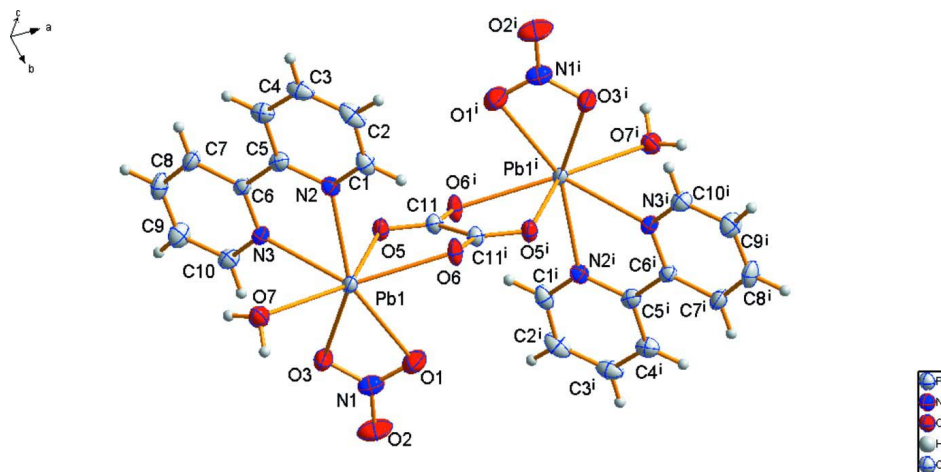
A mixture of oxalic acid (0.0634 g, 0.5 mmol), 2,2'-bipyridine (0.0781 g, 0.5 mmol), Pb(NO<sub>3</sub>)<sub>2</sub> (0.3312 g, 1 mmol), NaOH (0.0400 g, 1 mmol), water (10 ml) and ethanol (5 ml) was placed in a Parr Teflon-lined stainless steel vessel (25 cm<sup>3</sup>). The vessel was then sealed and heated at 403 K for 3 days. Afterwards the mixture was slowly cooled to room temperature, colorless block-shaped crystals of the complex were obtained. Elemental analysis calculated (%<sub>wt</sub>): C 27.11; H 2.07; N 8.62; found (%<sub>wt</sub>): C 27.06; H 2.03; N 8.51.

**Refinement**

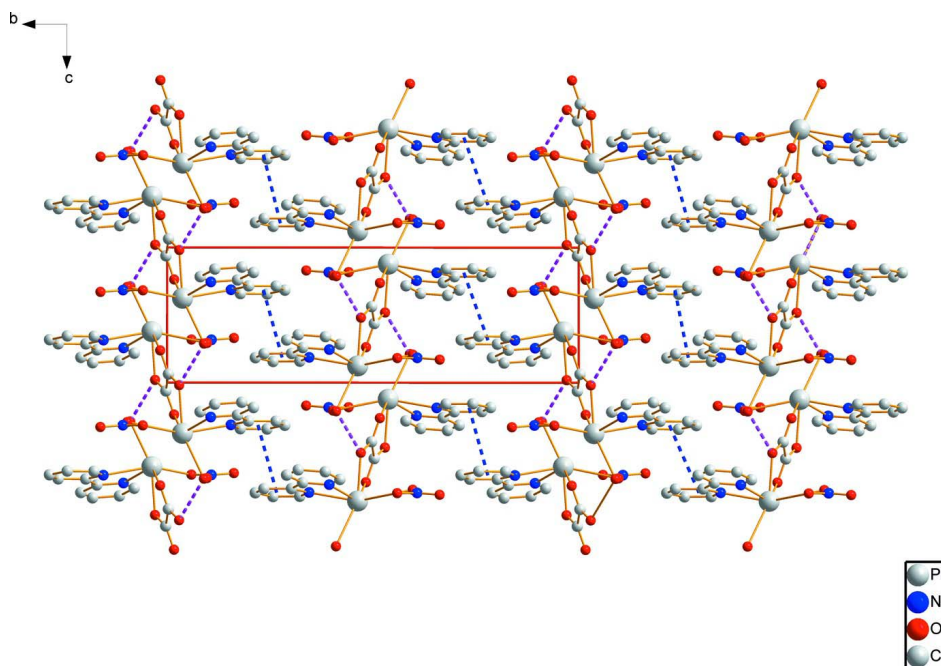
H atoms were positioned geometrically and refined as riding atoms, with C—H = 0.93 Å and  $U_{iso}(H) = 1.2U_{eq}(C)$  for aromatic H atoms. H atoms of the water molecule were located in a difference Fourier map and refined as riding, with O—H = 0.85 Å and with  $U_{iso}(H) = 1.2U_{eq}(O)$ . The maximum remaining electron density is found 0.97 Å from Pb1, and the minimum density 0.91 Å from the same atom.

**Computing details**

Data collection: *SMART* (Bruker, 2003); cell refinement: *SAINTE* (Bruker, 2003); data reduction: *SAINTE* (Bruker, 2003); program(s) used to solve structure: *SHELXS97* (Sheldrick, 2008); program(s) used to refine structure: *SHELXL97* (Sheldrick, 2008); molecular graphics: *DIAMOND* (Brandenburg, 2009); software used to prepare material for publication: *SHELXTL* (Sheldrick, 2008).


**Figure 1**

The coordination environment around Pb<sup>II</sup> in the title compound with the atom-labeling scheme. Displacement ellipsoids for non-hydrogen atoms are drawn at the 30% probability level. [Symmetry code: (i)  $-x+1, -y+1, -z+1$ .]


**Figure 2**

O—H $\cdots$ O hydrogen bonding (purple dashed lines) and  $\pi$ — $\pi$  stacking interactions (blue dashed lines) in the structure of the title compound. All H atoms were omitted for clarity.

**$\mu$ -Oxalato- $\kappa^4 O^1, O^2:O^1', O^2'$ - bis[aqua(2,2'-bipyridine- $\kappa N$ )(nitrate- $\kappa^2 O, O'$ )lead(II)]**

*Crystal data*

[Pb<sub>2</sub>(C<sub>2</sub>O<sub>4</sub>)(NO<sub>3</sub>)<sub>2</sub>(C<sub>10</sub>H<sub>8</sub>N<sub>2</sub>)<sub>2</sub>(H<sub>2</sub>O)<sub>2</sub>]

$M_r = 974.82$

Monoclinic,  $P2_1/c$

Hall symbol:  $-P 2_1/c$

$a = 9.5791(19) \text{ \AA}$

$b = 20.6330(14) \text{ \AA}$

$c = 6.7649(15) \text{ \AA}$

$\beta = 91.687(1)^\circ$

$V = 1336.5(4) \text{ \AA}^3$

$Z = 2$

$F(000) = 908$   
 $D_x = 2.422 \text{ Mg m}^{-3}$   
 Mo  $K\alpha$  radiation,  $\lambda = 0.71073 \text{ \AA}$   
 Cell parameters from 3490 reflections  
 $\theta = 2.3\text{--}28.3^\circ$

$\mu = 12.66 \text{ mm}^{-1}$   
 $T = 296 \text{ K}$   
 Block, colorless  
 $0.29 \times 0.28 \times 0.26 \text{ mm}$

*Data collection*

Bruker SMART CCD  
 diffractometer  
 Radiation source: fine-focus sealed tube  
 Graphite monochromator  
 phi and  $\omega$  scans  
 Absorption correction: multi-scan  
 (SADABS; Bruker, 2003)  
 $T_{\min} = 0.120$ ,  $T_{\max} = 0.137$

7096 measured reflections  
 2336 independent reflections  
 2082 reflections with  $I > 2\sigma(I)$   
 $R_{\text{int}} = 0.038$   
 $\theta_{\text{max}} = 25.0^\circ$ ,  $\theta_{\text{min}} = 2.0^\circ$   
 $h = -11 \rightarrow 10$   
 $k = -24 \rightarrow 24$   
 $l = -7 \rightarrow 8$

*Refinement*

Refinement on  $F^2$   
 Least-squares matrix: full  
 $R[F^2 > 2\sigma(F^2)] = 0.031$   
 $wR(F^2) = 0.082$   
 $S = 1.07$   
 2336 reflections  
 190 parameters  
 0 restraints  
 Primary atom site location: structure-invariant  
 direct methods

Secondary atom site location: difference Fourier  
 map  
 Hydrogen site location: inferred from  
 neighbouring sites  
 H-atom parameters constrained  
 $w = 1/[\sigma^2(F_o^2) + (0.035P)^2 + 5.1328P]$   
 where  $P = (F_o^2 + 2F_c^2)/3$   
 $(\Delta/\sigma)_{\text{max}} = 0.001$   
 $\Delta\rho_{\text{max}} = 2.68 \text{ e \AA}^{-3}$   
 $\Delta\rho_{\text{min}} = -1.19 \text{ e \AA}^{-3}$

*Special details*

**Geometry.** All esds (except the esd in the dihedral angle between two l.s. planes) are estimated using the full covariance matrix. The cell esds are taken into account individually in the estimation of esds in distances, angles and torsion angles; correlations between esds in cell parameters are only used when they are defined by crystal symmetry. An approximate (isotropic) treatment of cell esds is used for estimating esds 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  $F^2 > 2\sigma(F^2)$  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 ( $\text{\AA}^2$ )*

	<i>x</i>	<i>y</i>	<i>z</i>	$U_{\text{iso}}^*/U_{\text{eq}}$
Pb1	0.68880 (3)	0.537173 (12)	0.87652 (4)	0.03036 (13)
N1	0.7954 (9)	0.3925 (3)	0.8164 (10)	0.0504 (18)
N2	0.6271 (6)	0.6548 (3)	0.8129 (9)	0.0345 (14)
N3	0.8863 (6)	0.6061 (3)	0.7555 (8)	0.0317 (13)
O1	0.6656 (8)	0.4032 (3)	0.7926 (12)	0.0678 (18)
O2	0.8406 (9)	0.3377 (3)	0.8302 (12)	0.084 (2)
O3	0.8765 (6)	0.4408 (3)	0.8182 (10)	0.0563 (16)
O5	0.6731 (5)	0.5291 (2)	0.5190 (8)	0.0393 (13)
O6	0.5602 (5)	0.4830 (3)	0.2626 (7)	0.0411 (12)
O7	0.8279 (6)	0.5894 (3)	1.2112 (8)	0.0463 (13)
H7A	0.8289	0.5600	1.2993	0.056*
H7B	0.9123	0.5956	1.1805	0.056*

C1	0.4989 (8)	0.6777 (4)	0.8412 (12)	0.0449 (19)
H1	0.4275	0.6480	0.8607	0.054*
C2	0.4664 (9)	0.7424 (4)	0.8432 (13)	0.052 (2)
H2	0.3755	0.7562	0.8633	0.062*
C3	0.5712 (9)	0.7861 (4)	0.8148 (12)	0.049 (2)
H3	0.5521	0.8303	0.8155	0.058*
C4	0.7051 (9)	0.7647 (4)	0.7851 (12)	0.0408 (18)
H4	0.7772	0.7940	0.7661	0.049*
C5	0.7308 (8)	0.6978 (3)	0.7841 (10)	0.0332 (16)
C6	0.8707 (7)	0.6707 (3)	0.7405 (10)	0.0294 (15)
C7	0.9824 (8)	0.7097 (4)	0.6843 (12)	0.0425 (18)
H7	0.9715	0.7544	0.6737	0.051*
C8	1.1074 (8)	0.6813 (4)	0.6453 (13)	0.051 (2)
H8	1.1828	0.7067	0.6091	0.061*
C9	1.1214 (8)	0.6160 (4)	0.6595 (13)	0.047 (2)
H9	1.2056	0.5961	0.6315	0.057*
C10	1.0100 (8)	0.5801 (4)	0.7156 (11)	0.0384 (17)
H10	1.0206	0.5354	0.7267	0.046*
C11	0.5660 (7)	0.5032 (3)	0.4367 (11)	0.0319 (15)

Atomic displacement parameters ( $\text{\AA}^2$ )

	$U^{11}$	$U^{22}$	$U^{33}$	$U^{12}$	$U^{13}$	$U^{23}$
Pb1	0.02946 (19)	0.03118 (19)	0.03045 (19)	-0.00956 (10)	0.00100 (11)	0.00129 (10)
N1	0.066 (5)	0.037 (4)	0.048 (4)	0.001 (4)	0.006 (3)	0.002 (3)
N2	0.032 (3)	0.036 (3)	0.035 (3)	0.003 (3)	0.002 (3)	-0.002 (3)
N3	0.031 (3)	0.029 (3)	0.035 (3)	-0.007 (2)	0.002 (2)	-0.001 (3)
O1	0.063 (5)	0.044 (4)	0.097 (5)	-0.009 (3)	-0.001 (4)	0.006 (4)
O2	0.119 (7)	0.032 (4)	0.101 (6)	0.018 (4)	0.009 (5)	0.004 (4)
O3	0.043 (3)	0.042 (3)	0.083 (5)	-0.008 (3)	-0.005 (3)	0.006 (3)
O5	0.027 (3)	0.051 (3)	0.040 (3)	-0.011 (2)	0.003 (2)	-0.001 (2)
O6	0.029 (3)	0.065 (3)	0.030 (3)	-0.013 (2)	0.004 (2)	-0.012 (3)
O7	0.040 (3)	0.055 (3)	0.044 (3)	-0.005 (3)	0.003 (2)	0.006 (3)
C1	0.032 (4)	0.062 (6)	0.041 (5)	0.001 (4)	0.005 (3)	-0.001 (4)
C2	0.043 (5)	0.066 (6)	0.047 (5)	0.023 (4)	0.005 (4)	-0.008 (4)
C3	0.065 (6)	0.039 (5)	0.041 (5)	0.017 (4)	-0.007 (4)	-0.005 (4)
C4	0.049 (5)	0.038 (4)	0.035 (4)	0.006 (4)	-0.001 (3)	-0.002 (3)
C5	0.041 (4)	0.030 (4)	0.028 (4)	0.000 (3)	0.001 (3)	-0.005 (3)
C6	0.034 (4)	0.028 (4)	0.026 (4)	-0.005 (3)	0.000 (3)	0.000 (3)
C7	0.042 (4)	0.036 (4)	0.050 (5)	-0.016 (3)	0.004 (3)	0.004 (3)
C8	0.034 (4)	0.061 (6)	0.058 (6)	-0.019 (4)	0.011 (4)	0.001 (4)
C9	0.035 (4)	0.057 (5)	0.051 (5)	0.000 (4)	0.015 (3)	-0.003 (4)
C10	0.037 (4)	0.036 (4)	0.042 (5)	0.000 (3)	0.006 (3)	-0.004 (3)
C11	0.026 (4)	0.033 (4)	0.037 (4)	-0.001 (3)	0.003 (3)	-0.001 (3)

Geometric parameters ( $\text{\AA}$ ,  $^\circ$ )

Pb1—O5	2.425 (5)	C1—C2	1.371 (12)
Pb1—N3	2.522 (5)	C1—H1	0.9300
Pb1—N2	2.532 (6)	C2—C3	1.368 (12)

Pb1—O6 <sup>i</sup>	2.572 (5)	C2—H2	0.9300
Pb1—O3	2.718 (6)	C3—C4	1.377 (11)
Pb1—O7	2.809 (6)	C3—H3	0.9300
N1—O2	1.214 (9)	C4—C5	1.401 (10)
N1—O3	1.263 (9)	C4—H4	0.9300
N1—O1	1.269 (10)	C5—C6	1.490 (10)
N2—C1	1.334 (9)	C6—C7	1.401 (9)
N2—C5	1.351 (9)	C7—C8	1.366 (11)
N3—C10	1.336 (9)	C7—H7	0.9300
N3—C6	1.344 (9)	C8—C9	1.358 (12)
O5—C11	1.270 (9)	C8—H8	0.9300
O6—C11	1.249 (9)	C9—C10	1.362 (11)
O6—Pb1 <sup>i</sup>	2.572 (5)	C9—H9	0.9300
O7—H7A	0.8500	C10—H10	0.9300
O7—H7B	0.8499	C11—C11 <sup>i</sup>	1.554 (13)
O5—Pb1—N3	74.93 (17)	C2—C1—H1	118.1
O5—Pb1—N2	83.62 (18)	C3—C2—C1	118.2 (8)
N3—Pb1—N2	65.04 (19)	C3—C2—H2	120.9
O5—Pb1—O6 <sup>i</sup>	66.07 (16)	C1—C2—H2	120.9
N3—Pb1—O6 <sup>i</sup>	131.97 (18)	C2—C3—C4	120.0 (8)
N2—Pb1—O6 <sup>i</sup>	83.30 (19)	C2—C3—H3	120.0
O5—Pb1—O3	80.03 (19)	C4—C3—H3	120.0
N3—Pb1—O3	81.95 (18)	C3—C4—C5	118.8 (8)
N2—Pb1—O3	146.01 (19)	C3—C4—H4	120.6
O6 <sup>i</sup> —Pb1—O3	116.03 (18)	C5—C4—H4	120.6
O5—Pb1—O7	147.62 (16)	N2—C5—C4	121.1 (7)
N3—Pb1—O7	72.69 (17)	N2—C5—C6	116.8 (6)
N2—Pb1—O7	82.64 (18)	C4—C5—C6	122.0 (7)
O6 <sup>i</sup> —Pb1—O7	140.35 (16)	N3—C6—C7	120.5 (7)
O3—Pb1—O7	95.61 (18)	N3—C6—C5	117.1 (6)
O2—N1—O3	121.0 (8)	C7—C6—C5	122.4 (6)
O2—N1—O1	121.2 (8)	C8—C7—C6	119.1 (7)
O3—N1—O1	117.7 (7)	C8—C7—H7	120.4
C1—N2—C5	118.2 (7)	C6—C7—H7	120.4
C1—N2—Pb1	121.7 (5)	C9—C8—C7	119.8 (7)
C5—N2—Pb1	119.0 (5)	C9—C8—H8	120.1
C10—N3—C6	118.7 (6)	C7—C8—H8	120.1
C10—N3—Pb1	121.2 (5)	C8—C9—C10	118.9 (8)
C6—N3—Pb1	120.0 (4)	C8—C9—H9	120.5
N1—O3—Pb1	99.7 (5)	C10—C9—H9	120.5
C11—O5—Pb1	119.4 (4)	N3—C10—C9	123.0 (7)
C11—O6—Pb1 <sup>i</sup>	114.5 (4)	N3—C10—H10	118.5
Pb1—O7—H7A	106.6	C9—C10—H10	118.5
Pb1—O7—H7B	107.0	O6—C11—O5	124.5 (6)
H7A—O7—H7B	106.7	O6—C11—C11 <sup>i</sup>	118.5 (8)
N2—C1—C2	123.8 (8)	O5—C11—C11 <sup>i</sup>	117.0 (8)
N2—C1—H1	118.1		

Symmetry code: (i)  $-x+1, -y+1, -z+1$ .

Hydrogen-bond geometry (Å, °)

$D-H\cdots A$	$D-H$	$H\cdots A$	$D\cdots A$	$D-H\cdots A$
$O7-H7A\cdots O5^{ii}$	0.85	2.23	2.875 (7)	133
$O7-H7B\cdots O3^{iii}$	0.85	2.16	2.912 (8)	148

Symmetry codes: (ii)  $x, y, z+1$ ; (iii)  $-x+2, -y+1, -z+2$ .