Acta Crystallographica Section E
Structure Reports Online
ISSN 1600-5368

Poly[(μ_4-3-carboxybenzoato-$\kappa^5O^1:O^1,O^{1'}:O^{1'}:O^3$)(quinolin-8-olato-$\kappa^2N,O$)-lead(II)]

Akbar Ghaemi,[a]‡ Zohreh Dadkhah,[a] Seik Weng Ng[b,c] and Edward R. T. Tiekink[b]*

[a]Department of Chemistry, Saveh Branch, Islamic Azad University, Saveh, Iran, [b]Department of Chemistry, University of Malaya, 50603 Kuala Lumpur, Malaysia, and [c]Chemistry Department, Faculty of, Science, King Abdulaziz University, PO Box 80203 Jeddah, Saudi Arabia
Correspondence e-mail: edward.tiekink@gmail.com

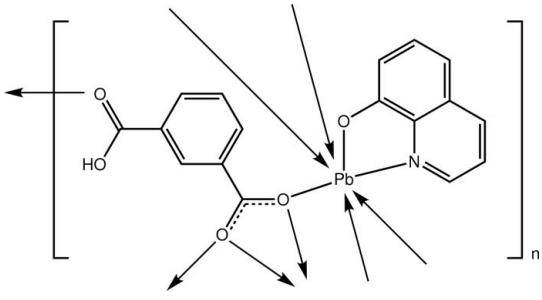
Received 19 December 2011; accepted 21 December 2011

Key indicators: single-crystal X-ray study; T = 100 K; mean σ(C–C) = 0.005 Å; R factor = 0.021; wR factor = 0.049; data-to-parameter ratio = 15.0.

The asymmetric unit of the title complex, $[Pb(C_8H_5O_4)(C_9H_6NO)]_n$, comprises a Pb^{II} cation, a quinolin-8-olate anion and a 3-carboxybenzoate anion. The coordination geometry of the Pb^{II} atom is defined by one N and six O atoms, as well as a stereochemically active lone pair of electrons, and is based on a Ψ-dodecahedron. The quinolin-8-olate is chelating and the 3-carboxybenzoate anion forms bonds to four different Pb^{II} atoms. The benzoate end of the 3-carboxybenzoate ligand chelates one Pb^{II} atom and simultaneously bridges two Pb^{II} atoms on either side, forming a chain along the b axis. The carboxyl end of the 3-carboxybenzoate connects to a neighbouring chain by employing its carbonyl atom to form a bond to a Pb^{II} atom and the hydroxyl group to form a hydrogen bond to a quinolin-8-olate O atom. Thereby, a layer is formed in the bc plane.

Related literature

For background to Pb^{II} mixed quinolate carboxylate structures, see: Shahverdizadeh *et al.* (2008).



Experimental

Crystal data

$[Pb(C_8H_5O_4)(C_9H_6NO)]$
M_r = 516.46
Monoclinic, $P2_1/c$
a = 9.0746 (2) Å
b = 7.0262 (2) Å
c = 22.6919 (6) Å
β = 93.185 (3)°
V = 1444.60 (6) Å^3
Z = 4
Mo $K\alpha$ radiation
μ = 11.71 mm^{-1}
T = 100 K
0.25 × 0.20 × 0.15 mm

Data collection

Agilent SuperNova Dual diffractometer with an Atlas detector
Absorption correction: multi-scan (*CrysAlis PRO*; Agilent, 2010)
T_{min} = 0.158, T_{max} = 0.273
9690 measured reflections
3325 independent reflections
3035 reflections with $I > 2\sigma(I)$
R_{int} = 0.029

Refinement

$R[F^2 > 2\sigma(F^2)]$ = 0.021
$wR(F^2)$ = 0.049
S = 1.01
3325 reflections
221 parameters
1 restraint
H atoms treated by a mixture of independent and constrained refinement
$\Delta\rho_{max}$ = 0.92 e Å^{-3}
$\Delta\rho_{min}$ = −1.27 e Å^{-3}

Table 1
Selected bond lengths (Å).

Pb—O1	2.608 (2)	Pb—O3^{iii}	2.840 (3)
Pb—O1^{i}	2.746 (2)	Pb—O5	2.318 (2)
Pb—O2^{ii}	2.578 (2)	Pb—N1	2.468 (3)
Pb—O2^{i}	2.809 (2)		

Symmetry codes: (i) $-x+1, y-\frac{1}{2}, -z+\frac{1}{2}$; (ii) $x, y-1, z$; (iii) $-x+1, -y+1, -z+1$.

Table 2
Hydrogen-bond geometry (Å, °).

D—H···A	D—H	H···A	D···A	D—H···A
O4—H1···O5^{iii}	0.84 (1)	1.74 (3)	2.539 (4)	158 (6)

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

Data collection: *CrysAlis PRO* (Agilent, 2010); cell refinement: *CrysAlis PRO*; data reduction: *CrysAlis PRO*; program(s) used to solve structure: *SHELXS97* (Sheldrick, 2008); program(s) used to refine structure: *SHELXL97* (Sheldrick, 2008); molecular graphics: *ORTEP-3* (Farrugia, 1997) and *DIAMOND* (Brandenburg, 2006); software used to prepare material for publication: *publCIF* (Westrip, 2010).

We gratefully acknowledge practical support of this study by the Islamic Azad University (Saveh Branch), and thank the University of Malaya for supporting the crystallographic facility.

Supplementary data and figures for this paper are available from the IUCr electronic archives (Reference: QM2046).

‡ Additional correspondence author, e-mail: akbarghaemi@yahoo.com.

References

Agilent (2010). *CrysAlis PRO*. Agilent Technologies, Yarnton, England.
Brandenburg, K. (2006). *DIAMOND*. Crystal Impact GbR, Bonn, Germany.
Farrugia, L. J. (1997). *J. Appl. Cryst.* **30**, 565.
Shahverdizadeh, G. H., Soudi, A. A., Morsali, A. & Retailleau, P. (2008). *Inorg. Chim. Acta*, **361**, 1875–1884.
Sheldrick, G. M. (2008). *Acta Cryst.* A**64**, 112–122.
Westrip, S. P. (2010). *J. Appl. Cryst.* **43**, 920–925.

supplementary materials

Acta Cryst. (2012). E**68**, m97-m98 [**doi:10.1107/S160053681105495X**]

Poly[(μ_4-3-carboxybenzoato-$\kappa^5O^1:O^1,O^{1'}:O^{1'}:O^3$)(quinolin-8-olato-$\kappa^2N,O$)lead(II)]

A. Ghaemi, Z. Dadkhah, S. W. Ng and E. R. T. Tiekink

Comment

Mixed lead(II) complexes of quinolin-8-olate and monofunctional carboxylates have displayed a variety of structural motifs (Shahverdizadeh *et al.*, 2008). In the present report, a 1:1 structure containing quinolin-8-olate and 3-carboxybenzoate is described, (I).

The asymmetric unit of (I) comprises a Pb^{II} cation, a quinolin-8-olate anion and a 3-carboxybenzoate anion, Fig. 1. The coordination geometry of the Pb^{II} atom is defined by a N and six O atoms as well as a stereochemically active lone pair of electrons, and is based on a Ψ-dodecahedron. The quinolin-8-olate anion is chelating, whereas the 3-carboxybenzoate anion is pentadentate, forming bonds to four different Pb^{II} atoms, Table 1. The benzoate group chelates one Pb^{II} atom and each of these O atoms forms a bond to a neighbouring Pb^{II} to form a chain along the *b* axis. Adjacent chains, along the *c* axis, are connected by Pb—*O*(carbonyl) bonds. The hydroxyl group forms a hydrogen bond to the quinolin-8-olate-O atom, Table 2. The result is a layer in the *bc* plane. Layers stack along the *a* axis, Fig. 3, with no specific intermolecular interactions between them.

Experimental

The title complex was obtained by the following method. A methanol solution (10 ml) of 8-hydroxyquinoline (0.145 g, 1 mmol) was added to an aqueous solution (2 ml) of $Pb(NO_3)_2$ (0.331 g, 1 mmol). The mixture was stirred for 10 min. To this solution, was added a DMF solution (5 ml) of isophthalic acid (0.084 g, 0.5 mmol) slowly at room temperature. This mixture was filtered. After keeping the filtrate in air, crystals were formed at the bottom of the vessel upon slow evaporation of the solvents at room temperature. *M*.pt. 558 K (dec.). Yield: 65%.

Refinement

Carbon-bound H-atoms were placed in calculated positions [C—H 0.95 Å, U_{iso}(H) 1.2U_{eq}(C)] and were included in the refinement in the riding model approximation. The acid H-atom was located in a difference Fourier map, and was refined with a distance restraint of O—H 0.84±0.01 Å; its U_{iso} value was refined. The final difference Fourier map had a peak at 0.81 Å from Pb and a hole at 0.90 Å from the same atom.

Figures

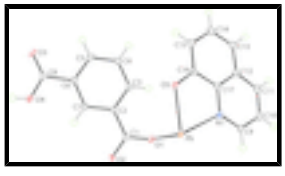


Fig. 1. The asymmetric unit of (I) showing the atom-labelling scheme and displacement ellipsoids at the 50% probability level.

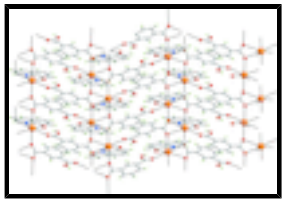


Fig. 2. A view of the layer in the *bc* plane in (I).

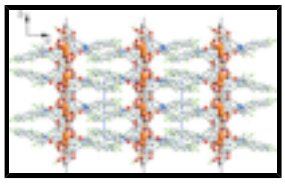


Fig. 3. A view in projection down the *c* axis of the unit-cell contents of (I) highlighting the stacking of layers.

Poly[(μ$_4$-3-carboxybenzoato- κ^5*O*^{1}:*O*^{1},*O*$^{1'}$:*O*$^{1'}$:*O*^{3})(quinolin- 8-olato-κ^2*N*,*O*)lead(II)]

Crystal data

[Pb(C$_8$H$_5$O$_4$)(C$_9$H$_6$NO)]	*F*(000) = 968
M_r = 516.46	D_x = 2.375 Mg m^{-3}
Monoclinic, *P*2$_1$/*c*	Mo *K*α radiation, λ = 0.71073 Å
Hall symbol: -P 2ybc	Cell parameters from 6118 reflections
a = 9.0746 (2) Å	θ = 2.2–27.5°
b = 7.0262 (2) Å	μ = 11.71 mm^{-1}
c = 22.6919 (6) Å	*T* = 100 K
β = 93.185 (3)°	Block, yellow
V = 1444.60 (6) Å^3	0.25 × 0.20 × 0.15 mm
Z = 4	

Data collection

Agilent SuperNova Dual diffractometer with an Atlas detector	3325 independent reflections
Radiation source: SuperNova (Mo) X-ray Source	3035 reflections with $I > 2\sigma(I)$
Mirror	R_{int} = 0.029
Detector resolution: 10.4041 pixels mm^{-1}	θ$_{max}$ = 27.6°, θ$_{min}$ = 2.8°
ω scan	*h* = −11→11
Absorption correction: multi-scan (*CrysAlis PRO*; Agilent, 2010)	*k* = −9→8
T_{min} = 0.158, T_{max} = 0.273	*l* = −29→27
9690 measured reflections	

Refinement

Refinement on F^2	Primary atom site location: structure-invariant direct methods
Least-squares matrix: full	Secondary atom site location: difference Fourier map
$R[F^2 > 2\sigma(F^2)]$ = 0.021	Hydrogen site location: inferred from neighbouring sites

$wR(F^2) = 0.049$	H atoms treated by a mixture of independent and constrained refinement
$S = 1.01$	$w = 1/[\sigma^2(F_o^2) + (0.0237P)^2 + 0.5749P]$ where $P = (F_o^2 + 2F_c^2)/3$
3325 reflections	$(\Delta/\sigma)_{max} = 0.001$
221 parameters	$\Delta\rho_{max} = 0.92$ e Å^{-3}
1 restraint	$\Delta\rho_{min} = -1.27$ e Å^{-3}

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

	x	*y*	*z*	U_{iso}*/U_{eq}
Pb	0.513121 (14)	0.057458 (17)	0.305356 (5)	0.00777 (5)
O1	0.5980 (3)	0.4114 (3)	0.30020 (11)	0.0123 (5)
O2	0.4886 (3)	0.6925 (3)	0.31117 (10)	0.0107 (5)
O3	0.6052 (3)	0.7915 (4)	0.59124 (11)	0.0160 (6)
O4	0.4185 (3)	0.8397 (4)	0.52393 (12)	0.0141 (5)
H1	0.365 (6)	0.890 (8)	0.549 (2)	0.08 (2)*
O5	0.6896 (3)	0.0346 (3)	0.38285 (11)	0.0111 (5)
N1	0.7528 (4)	−0.0485 (4)	0.27073 (13)	0.0104 (6)
C1	0.5716 (4)	0.5570 (5)	0.33079 (16)	0.0099 (7)
C2	0.6368 (4)	0.5693 (4)	0.39260 (16)	0.0084 (7)
C3	0.7672 (4)	0.4692 (5)	0.40797 (16)	0.0112 (7)
H3	0.8141	0.3966	0.3791	0.013*
C4	0.8270 (4)	0.4767 (5)	0.46522 (17)	0.0136 (8)
H4	0.9160	0.4104	0.4754	0.016*
C5	0.7586 (4)	0.5801 (5)	0.50806 (17)	0.0140 (8)
H5	0.8006	0.5848	0.5473	0.017*
C6	0.6275 (4)	0.6774 (5)	0.49317 (15)	0.0111 (7)
C7	0.5672 (4)	0.6744 (5)	0.43551 (15)	0.0100 (7)
H7	0.4794	0.7432	0.4253	0.012*
C8	0.5498 (4)	0.7761 (5)	0.54049 (16)	0.0109 (7)
C9	0.7848 (4)	−0.0880 (5)	0.21594 (16)	0.0116 (8)
H9	0.7081	−0.0799	0.1858	0.014*
C10	0.9255 (4)	−0.1409 (5)	0.19995 (17)	0.0149 (8)
H10	0.9432	−0.1670	0.1599	0.018*
C11	1.0383 (4)	−0.1548 (5)	0.24300 (16)	0.0140 (8)
H11	1.1352	−0.1870	0.2327	0.017*
C12	1.0090 (4)	−0.1206 (5)	0.30261 (16)	0.0117 (8)
C13	1.1178 (4)	−0.1327 (5)	0.34965 (17)	0.0148 (8)
H13	1.2155	−0.1707	0.3422	0.018*
C14	1.0821 (4)	−0.0899 (5)	0.40557 (18)	0.0156 (8)
H14	1.1558	−0.0983	0.4369	0.019*
C15	0.9372 (4)	−0.0329 (5)	0.41800 (18)	0.0144 (8)
H15	0.9155	−0.0037	0.4575	0.017*
C16	0.8271 (4)	−0.0193 (5)	0.37373 (17)	0.0106 (7)
C17	0.8635 (4)	−0.0657 (4)	0.31489 (16)	0.0098 (7)

Atomic displacement parameters (Å^2)

	U^{11}	U^{22}	U^{33}	U^{12}	U^{13}	U^{23}
Pb	0.00854 (8)	0.00841 (7)	0.00647 (8)	0.00005 (5)	0.00123 (5)	−0.00047 (5)
O1	0.0151 (14)	0.0122 (12)	0.0097 (13)	−0.0020 (11)	0.0002 (11)	−0.0010 (10)
O2	0.0123 (13)	0.0092 (12)	0.0107 (13)	0.0024 (10)	0.0018 (10)	−0.0005 (10)
O3	0.0171 (14)	0.0249 (14)	0.0061 (13)	−0.0028 (12)	0.0007 (11)	−0.0049 (11)
O4	0.0134 (14)	0.0197 (13)	0.0093 (13)	0.0005 (12)	0.0022 (11)	−0.0020 (11)
O5	0.0098 (13)	0.0145 (13)	0.0089 (13)	−0.0005 (10)	0.0005 (10)	−0.0008 (10)
N1	0.0118 (16)	0.0112 (15)	0.0080 (16)	−0.0034 (12)	−0.0002 (13)	−0.0007 (11)
C1	0.0112 (18)	0.0116 (17)	0.0072 (18)	−0.0050 (14)	0.0035 (14)	0.0023 (13)
C2	0.0123 (18)	0.0057 (15)	0.0074 (17)	−0.0025 (14)	0.0017 (14)	0.0017 (12)
C3	0.0126 (19)	0.0089 (16)	0.0123 (19)	−0.0005 (15)	0.0036 (15)	−0.0008 (14)
C4	0.0088 (18)	0.0169 (18)	0.015 (2)	0.0027 (15)	−0.0003 (15)	0.0001 (15)
C5	0.013 (2)	0.0189 (18)	0.0094 (19)	−0.0046 (16)	−0.0021 (15)	−0.0007 (15)
C6	0.0154 (19)	0.0104 (16)	0.0080 (18)	−0.0011 (15)	0.0045 (14)	0.0004 (14)
C7	0.0110 (18)	0.0084 (16)	0.0103 (18)	0.0000 (14)	−0.0031 (14)	−0.0012 (13)
C8	0.0120 (18)	0.0099 (16)	0.0108 (18)	−0.0054 (15)	0.0015 (14)	−0.0001 (14)
C9	0.0149 (19)	0.0112 (16)	0.0087 (18)	−0.0008 (15)	−0.0008 (15)	−0.0019 (14)
C10	0.020 (2)	0.0126 (18)	0.0129 (19)	−0.0008 (16)	0.0050 (15)	−0.0023 (14)
C11	0.0153 (19)	0.0098 (16)	0.017 (2)	0.0002 (15)	0.0062 (15)	0.0008 (14)
C12	0.0111 (19)	0.0068 (16)	0.017 (2)	−0.0019 (14)	0.0016 (15)	−0.0027 (14)
C13	0.0078 (18)	0.0150 (17)	0.021 (2)	0.0017 (15)	0.0003 (15)	0.0032 (15)
C14	0.012 (2)	0.0176 (18)	0.017 (2)	−0.0007 (16)	−0.0034 (16)	−0.0005 (15)
C15	0.014 (2)	0.0127 (18)	0.017 (2)	0.0015 (15)	0.0002 (16)	−0.0009 (15)
C16	0.0125 (19)	0.0031 (15)	0.016 (2)	0.0012 (14)	0.0022 (15)	0.0027 (13)
C17	0.0106 (18)	0.0071 (16)	0.0116 (19)	−0.0027 (14)	−0.0006 (15)	−0.0003 (13)

Geometric parameters (Å, °)

Pb—O1	2.608 (2)	C4—C5	1.388 (5)
Pb—O1^{i}	2.746 (2)	C4—H4	0.9500
Pb—O2^{ii}	2.578 (2)	C5—C6	1.397 (5)
Pb—O2^{i}	2.809 (2)	C5—H5	0.9500
Pb—O3^{iii}	2.840 (3)	C6—C7	1.390 (5)
Pb—O5	2.318 (2)	C6—C8	1.489 (5)
Pb—N1	2.468 (3)	C7—H7	0.9500
O1—C1	1.267 (4)	C9—C10	1.397 (5)
O1—Pb^{iv}	2.746 (2)	C9—H9	0.9500
O2—C1	1.278 (4)	C10—C11	1.379 (5)
O2—Pb^{v}	2.578 (2)	C10—H10	0.9500
O3—C8	1.235 (4)	C11—C12	1.414 (5)
O4—C8	1.308 (4)	C11—H11	0.9500
O4—H1	0.840 (10)	C12—C13	1.416 (5)
O5—C16	1.331 (4)	C12—C17	1.418 (5)
N1—C9	1.322 (5)	C13—C14	1.360 (5)

N1—C17	1.385 (4)	C13—H13	0.9500
C1—C2	1.494 (5)	C14—C15	1.418 (6)
C2—C3	1.403 (5)	C14—H14	0.9500
C2—C7	1.400 (5)	C15—C16	1.380 (5)
C3—C4	1.381 (5)	C15—H15	0.9500
C3—H3	0.9500	C16—C17	1.431 (5)
O5—Pb—N1	68.69 (10)	C6—C5—H5	120.2
O5—Pb—O2^{ii}	87.14 (7)	C7—C6—C5	120.4 (3)
N1—Pb—O2^{ii}	78.24 (8)	C7—C6—C8	120.6 (3)
O5—Pb—O1	84.66 (8)	C5—C6—C8	119.0 (3)
N1—Pb—O1	90.33 (9)	C6—C7—C2	119.6 (3)
O2^{ii}—Pb—O1	167.78 (8)	C6—C7—H7	120.2
O5—Pb—O1^{i}	147.23 (8)	C2—C7—H7	120.2
N1—Pb—O1^{i}	84.08 (9)	O3—C8—O4	123.7 (3)
O2^{ii}—Pb—O1^{i}	69.16 (7)	O3—C8—C6	121.9 (3)
O1—Pb—O1^{i}	114.35 (6)	O4—C8—C6	114.4 (3)
O5—Pb—O3^{iii}	71.10 (8)	N1—C9—C10	123.5 (3)
N1—Pb—O3^{iii}	139.12 (8)	N1—C9—H9	118.2
O2^{ii}—Pb—O3^{iii}	106.98 (8)	C10—C9—H9	118.2
O1—Pb—O3^{iii}	78.86 (8)	C11—C10—C9	119.2 (4)
O1^{i}—Pb—O3^{iii}	136.32 (8)	C11—C10—H10	120.4
C1—O1—Pb	132.9 (2)	C9—C10—H10	120.4
C1—O1—Pb^{iv}	96.0 (2)	C10—C11—C12	119.6 (4)
Pb—O1—Pb^{iv}	107.49 (8)	C10—C11—H11	120.2
C1—O2—Pb^{v}	135.0 (2)	C12—C11—H11	120.2
C8—O4—H1	120 (4)	C11—C12—C13	123.4 (4)
C16—O5—Pb	121.1 (2)	C11—C12—C17	117.5 (3)
C9—N1—C17	118.3 (3)	C13—C12—C17	119.1 (3)
C9—N1—Pb	127.4 (2)	C14—C13—C12	119.9 (4)
C17—N1—Pb	114.4 (2)	C14—C13—H13	120.1
O1—C1—O2	122.4 (3)	C12—C13—H13	120.1
O1—C1—C2	118.9 (3)	C13—C14—C15	121.3 (4)
O2—C1—C2	118.6 (3)	C13—C14—H14	119.4
C3—C2—C7	119.9 (3)	C15—C14—H14	119.4
C3—C2—C1	119.3 (3)	C16—C15—C14	121.1 (4)
C7—C2—C1	120.9 (3)	C16—C15—H15	119.5
C4—C3—C2	119.8 (4)	C14—C15—H15	119.5
C4—C3—H3	120.1	O5—C16—C15	123.6 (4)
C2—C3—H3	120.1	O5—C16—C17	118.5 (3)
C3—C4—C5	120.7 (4)	C15—C16—C17	117.9 (3)
C3—C4—H4	119.6	N1—C17—C12	121.9 (3)
C5—C4—H4	119.6	N1—C17—C16	117.4 (3)
C4—C5—C6	119.7 (3)	C12—C17—C16	120.7 (3)
C4—C5—H5	120.2		
O5—Pb—O1—C1	−77.9 (3)	C3—C4—C5—C6	0.2 (6)

N1—Pb—O1—C1	−146.5 (3)	C4—C5—C6—C7	−1.4 (5)
O2^{ii}—Pb—O1—C1	−125.9 (4)	C4—C5—C6—C8	175.7 (3)
O1^{i}—Pb—O1—C1	129.9 (3)	C5—C6—C7—C2	1.5 (5)
O3^{iii}—Pb—O1—C1	−6.2 (3)	C8—C6—C7—C2	−175.5 (3)
O5—Pb—O1—Pb^{iv}	165.82 (11)	C3—C2—C7—C6	−0.5 (5)
N1—Pb—O1—Pb^{iv}	97.27 (11)	C1—C2—C7—C6	177.6 (3)
O2^{ii}—Pb—O1—Pb^{iv}	117.8 (3)	C7—C6—C8—O3	−175.3 (3)
O1^{i}—Pb—O1—Pb^{iv}	13.62 (6)	C5—C6—C8—O3	7.6 (5)
O3^{iii}—Pb—O1—Pb^{iv}	−122.43 (10)	C7—C6—C8—O4	6.3 (5)
N1—Pb—O5—C16	0.4 (2)	C5—C6—C8—O4	−170.8 (3)
O2^{ii}—Pb—O5—C16	78.9 (2)	C17—N1—C9—C10	−1.9 (5)
O1—Pb—O5—C16	−92.1 (2)	Pb—N1—C9—C10	177.7 (3)
O1^{i}—Pb—O5—C16	36.2 (3)	N1—C9—C10—C11	0.4 (5)
O3^{iii}—Pb—O5—C16	−172.1 (2)	C9—C10—C11—C12	1.8 (5)
O5—Pb—N1—C9	−179.4 (3)	C10—C11—C12—C13	179.6 (3)
O2^{ii}—Pb—N1—C9	89.1 (3)	C10—C11—C12—C17	−2.4 (5)
O1—Pb—N1—C9	−95.3 (3)	C11—C12—C13—C14	177.2 (3)
O1^{i}—Pb—N1—C9	19.2 (3)	C17—C12—C13—C14	−0.7 (5)
O3^{iii}—Pb—N1—C9	−168.5 (2)	C12—C13—C14—C15	0.0 (6)
O5—Pb—N1—C17	0.1 (2)	C13—C14—C15—C16	0.1 (6)
O2^{ii}—Pb—N1—C17	−91.4 (2)	Pb—O5—C16—C15	179.1 (3)
O1—Pb—N1—C17	84.3 (2)	Pb—O5—C16—C17	−0.9 (4)
O1^{i}—Pb—N1—C17	−161.3 (2)	C14—C15—C16—O5	−179.5 (3)
O3^{iii}—Pb—N1—C17	11.0 (3)	C14—C15—C16—C17	0.4 (5)
Pb—O1—C1—O2	−108.0 (4)	C9—N1—C17—C12	1.1 (5)
Pb^{iv}—O1—C1—O2	12.7 (4)	Pb—N1—C17—C12	−178.5 (3)
Pb—O1—C1—C2	70.5 (4)	C9—N1—C17—C16	179.0 (3)
Pb^{iv}—O1—C1—C2	−168.8 (3)	Pb—N1—C17—C16	−0.6 (4)
Pb^{v}—O2—C1—O1	−129.2 (3)	C11—C12—C17—N1	1.0 (5)
Pb^{v}—O2—C1—C2	52.3 (5)	C13—C12—C17—N1	179.0 (3)
O1—C1—C2—C3	25.8 (5)	C11—C12—C17—C16	−176.8 (3)
O2—C1—C2—C3	−155.6 (3)	C13—C12—C17—C16	1.2 (5)
O1—C1—C2—C7	−152.3 (3)	O5—C16—C17—N1	1.0 (5)
O2—C1—C2—C7	26.2 (5)	C15—C16—C17—N1	−179.0 (3)
C7—C2—C3—C4	−0.7 (5)	O5—C16—C17—C12	178.9 (3)
C1—C2—C3—C4	−178.9 (3)	C15—C16—C17—C12	−1.1 (5)
C2—C3—C4—C5	0.9 (6)		

Symmetry codes: (i) −*x*+1, *y*−1/2, −*z*+1/2; (ii) *x*, *y*−1, *z*; (iii) −*x*+1, −*y*+1, −*z*+1; (iv) −*x*+1, *y*+1/2, −*z*+1/2; (v) *x*, *y*+1, *z*.

Hydrogen-bond geometry (Å, °)

D—H···*A*	*D*—H	H···*A*	*D*···*A*	*D*—H···*A*
O4—H1···O5^{iii}	0.84 (1)	1.74 (3)	2.539 (4)	158 (6)

Symmetry codes: (iii) −*x*+1, −*y*+1, −*z*+1.

Fig. 1

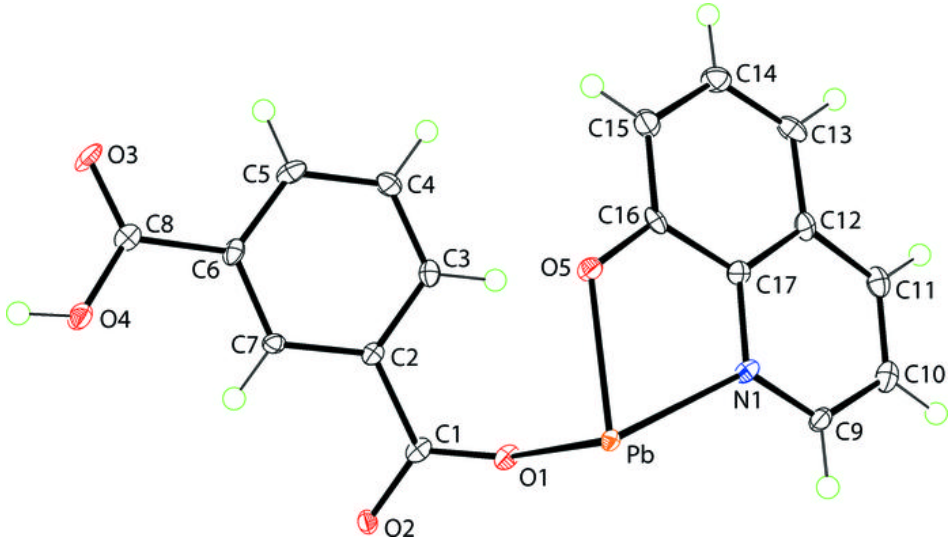

C14
C15
C13
O3
C5
C4
C16
C12
C8
C6
C3
O5
C17
C11
O4
C7
C2
C10
N1
C1
C9
Pb
O1
O2

Fig. 2

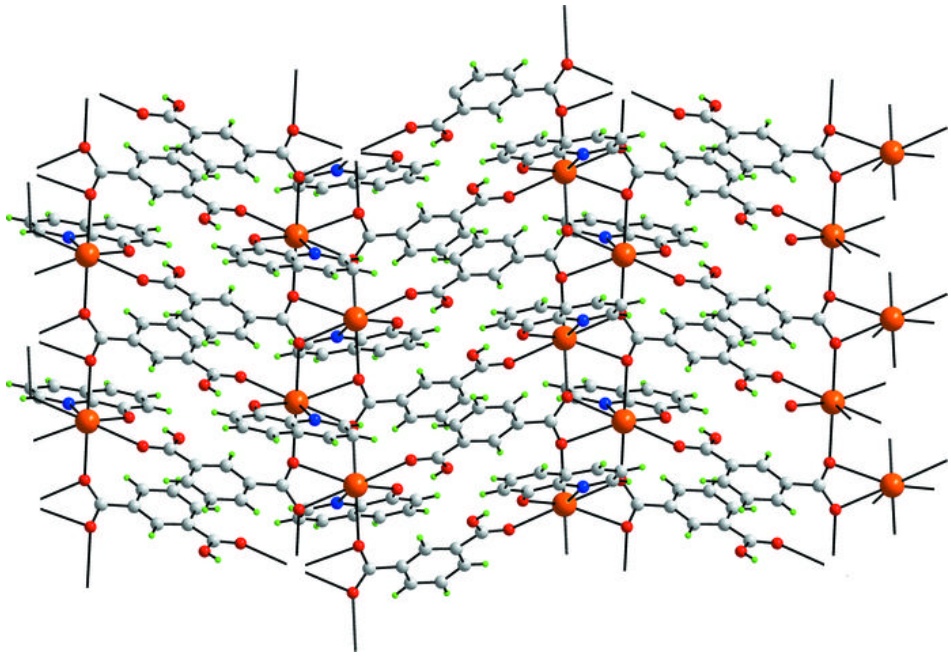


Fig. 3