

## Hexa- $\mu$ -chlorido-hexachlorido( $\eta^6$ -hexamethylbenzene)trialuminium(III)-lanthanum(III) benzene solvate

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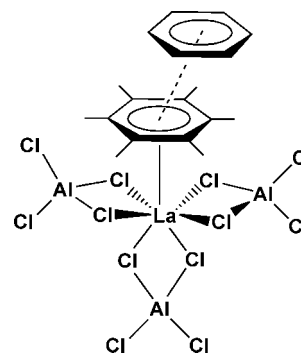
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Key indicators: single-crystal X-ray study;  $T = 173$  K; mean  $\sigma(\text{C}-\text{C}) = 0.005$  Å; disorder in main residue;  $R$  factor = 0.025;  $wR$  factor = 0.063; data-to-parameter ratio = 22.5.

In the title compound,  $[\text{Al}_3\text{LaCl}_{12}(\text{C}_{12}\text{H}_{18})]\cdot\text{C}_6\text{H}_6$ , all molecules are located on a mirror plane. Three chloridoaluminate groups and a hexamethylbenzene molecule are bound to the central lanthanum(III) ion, forming a distorted pentagonal bipyramid with the  $\eta^6$ -coordinated arene located at the apical position. The hexamethylbenzene ligand disordered between two orientations in a 1:1 ratio is also involved in parallel-slipped  $\pi$ - $\pi$  stacking intermolecular interactions with a benzene solvent molecule [centroid-centroid distance 3.612 (4) Å].

### Related literature

For the previously characterized lanthanum chloroaluminate and chlorogallate complexes, see: Filatov *et al.* (2008). For a recent review of other lanthanide chloroaluminate complexes, see: Bochkarev (2002). For complexes of lanthanide chlorogallates with polycyclic aromatic systems, see: Gorlov *et al.* (2008).



### Experimental

#### Crystal data

$[\text{Al}_3\text{LaCl}_{12}(\text{C}_{12}\text{H}_{18})]\cdot\text{C}_6\text{H}_6$	$V = 3404.9$ (3) Å <sup>3</sup>
$M_r = 885.62$	$Z = 4$
Orthorhombic, $Pnma$	Mo- $K\alpha$ radiation
$a = 12.2127$ (6) Å	$\mu = 2.28$ mm <sup>-1</sup>
$b = 16.4205$ (8) Å	$T = 173$ K
$c = 16.9790$ (8) Å	$0.22 \times 0.20 \times 0.16$ mm

#### Data collection

Bruker SMART APEX CCD area-detector diffractometer	28535 measured reflections
Absorption correction: multi-scan (SADABS; Bruker, 2003)	4250 independent reflections
$T_{\min} = 0.613$ , $T_{\max} = 0.697$	3911 reflections with $I > 2\sigma(I)$
	$R_{\text{int}} = 0.018$

#### Refinement

$R[F^2 > 2\sigma(F^2)] = 0.025$	189 parameters
$wR(F^2) = 0.063$	H-atom parameters not refined
$S = 1.05$	$\Delta\rho_{\max} = 0.89$ e Å <sup>-3</sup>
4250 reflections	$\Delta\rho_{\min} = -0.99$ e Å <sup>-3</sup>

Table 1

Selected bond lengths (Å).

La1—C1	2.945 (3)	La1—Cl4	2.9298 (6)
La1—C2	2.965 (2)	La1—Cl5	2.9083 (7)
La1—C3	2.957 (3)	La1—Cl6	2.9097 (7)
La1—C4	2.941 (4)	La1—Cg1	2.613 (3)
La1—Cl3	2.9128 (5)		

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: SHELXTL (Sheldrick, 2008) and ORTEP-3 for Windows (Farrugia, 1997); software used to prepare material for publication: SHELXTL and publCIF (Westrip, 2009).

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Supplementary data and figures for this paper are available from the IUCr electronic archives (Reference: CV2516).

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## supporting information

*Acta Cryst.* (2009). E65, m286–m287 [doi:10.1107/S1600536809004899]

## Hexa- $\mu$ -chlorido-hexachlorido( $\eta^6$ -hexamethylbenzene)-trialuminium(III)lanthanum(III) benzene solvate

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### S1. Comment

We have recently reported X-ray structural characterization of the first two lanthanum(III) chloroaluminate complexes with neutral arenes,  $[\text{La}(\eta^6\text{-C}_6\text{H}_5\text{Me})(\text{AlCl}_4)_3]$  and  $[\text{La}(\eta^6\text{-C}_6\text{Me}_6)(\text{AlCl}_4)_3]$ , as well as of the first lanthanum(III) chlorogallate complex,  $[\text{La}(\eta^6\text{-C}_6\text{Me}_6)(\text{GaCl}_4)_3]$  (Filatov *et al.*, 2008). The  $[\text{La}(\eta^6\text{-C}_6\text{Me}_6)(\text{AlCl}_4)_3] \cdot 0.5\text{C}_6\text{H}_6$  complex crystallizes in the monoclinic  $P2_1/c$  space group with the  $\beta$  angle being close to  $90^\circ$  ( $\beta = 90.27^\circ$ ). We later found that under slightly different experimental conditions, namely at a higher temperature (285 *versus* 273 K), the lanthanum complex with hexamethylbenzene,  $[\text{La}(\eta^6\text{-C}_6\text{Me}_6)(\text{AlCl}_4)_3] \cdot \text{C}_6\text{H}_6$  (I), is crystallized.

The molecular structure of (I) is comprised of the three chloroaluminate groups and a hexamethylbenzene molecule bound to the central lanthanum(III) ion (Fig.1). The coordination polyhedron is a distorted pentagonal bipyramid with the  $\eta^6$ -arene located at the apical position. The La–C bond distances span from 2.941 (4) to 2.965 (2) Å with a La–centroid distance being 2.613 (3) Å. These distances are comparable to those found in the previously reported complex  $[\text{La}(\eta^6\text{-C}_6\text{Me}_6)(\text{AlCl}_4)_3] \cdot 0.5\text{C}_6\text{H}_6$  (II) [La–C 2.927 (7)–3.035 (7) Å; La–centroid 2.633 (7) Å].

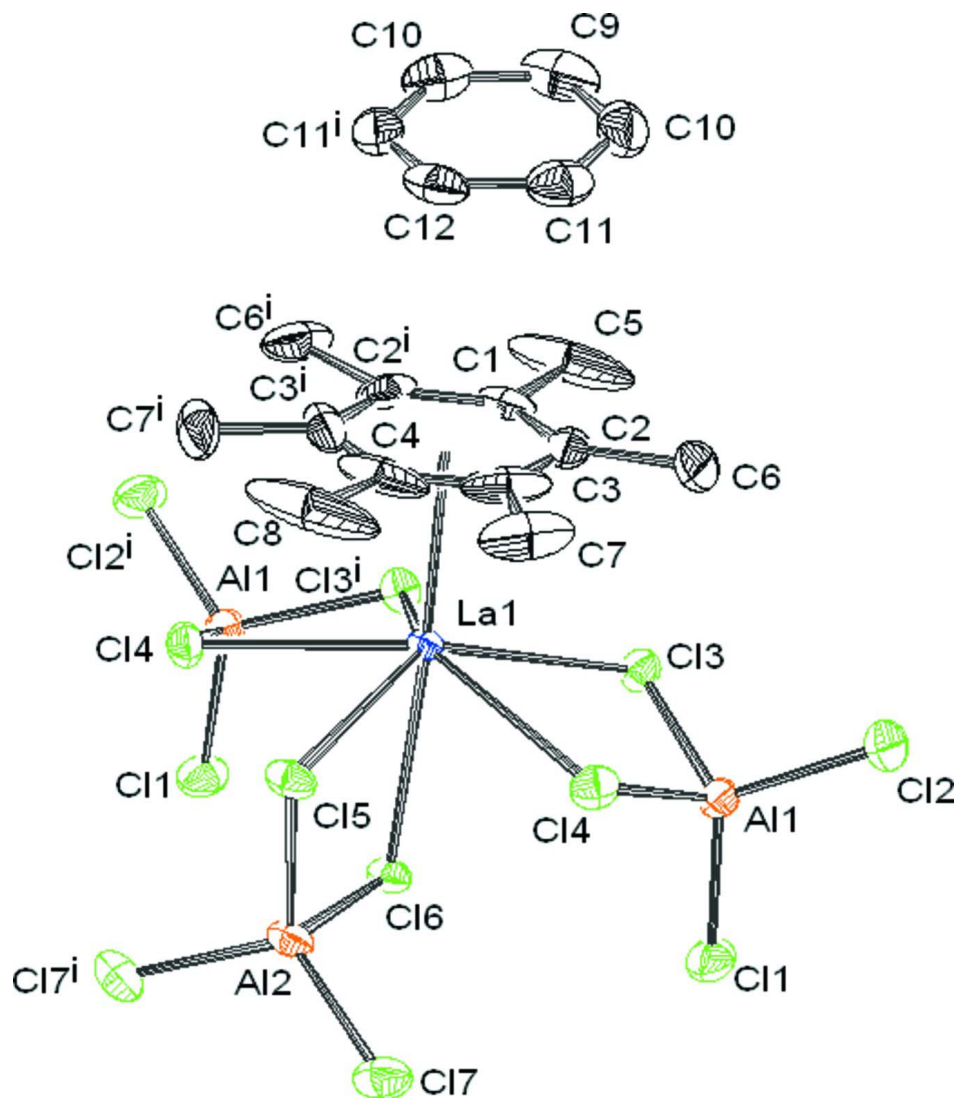
In (I), coordinated hexamethylbenzene is engaged into  $\pi$ - $\pi$  stacking interactions with a solvent benzene molecule. The intercentroid distance between their 6-membered rings is 3.612 (4) Å. The two ring planes are not parallel and the dihedral angle is  $12.7^\circ$  (Fig.2). In the above hemisolvate (II), on the contrary, both benzene faces are involved in  $\pi$ - $\pi$  stacking interactions as benzene is sandwiched between two hexamethylbenzene molecules. The distance between the centroids of the hexamethylbenzene and benzene rings (3.688 (4) Å) is noticeably longer than that found in (I).

### S2. Experimental

$\text{LaCl}_3$  (100 mg, 0.41 mmol),  $\text{AlCl}_3$  (163 mg, 1.22 mmol), hexamethylbenzene (66 mg, 0.41 mmol) and an excess of aluminium foil were placed into a Schlenk flask inside the glove box. Benzene (10 ml) was added to the flask and the mixture was refluxed for two hours. The  $\text{LaCl}_3$ ,  $\text{AlCl}_3$ , and hexamethylbenzene dissolved completely to give a yellow solution. The solution was filtered while hot through a pad of Celite and then kept at  $12^\circ\text{C}$  under argon for 2 days to afford a yellow crystalline material. Yield: 240 mg (65%). IR data ( $\text{cm}^{-1}$ ): 3091 (w), 3071 (w), 3036 (w), 1598 (m), 1531 (w), 1478 (m), 1423 (s), 1382 (m), 1332 (m), 1272 (m), 1180 (w), 1076 (w), 983 (w), 824 (w), 677 (s).

### S3. Refinement

All C–H atoms were refined using the riding model approximation, with C–H = 0.95–0.98 Å [ $U_{\text{iso}}(\text{H}) = 1.2$  or  $1.5U_{\text{eq}}(\text{C})$ ]. All other atoms were refined anisotropically. Large anisotropy of the carbon atoms of hexamethylbenzene suggests the presence of disorder. It was modeled over two rotational orientations in a 1:1 ratio. The C5 and C8 carbon atoms lie on a mirror plane and are constrained to have identical anisotropic displacement parameters (EADP command in the *SHELXL* realm).



**Figure 1**

A view of the molecular structure of (I), along with the atom numbering scheme [symmetry code: (i)  $x, -y + 1/2, z$ ]. Displacement ellipsoids are drawn at the 30% probability level. Hydrogen atoms and disordered parts are omitted for clarity.

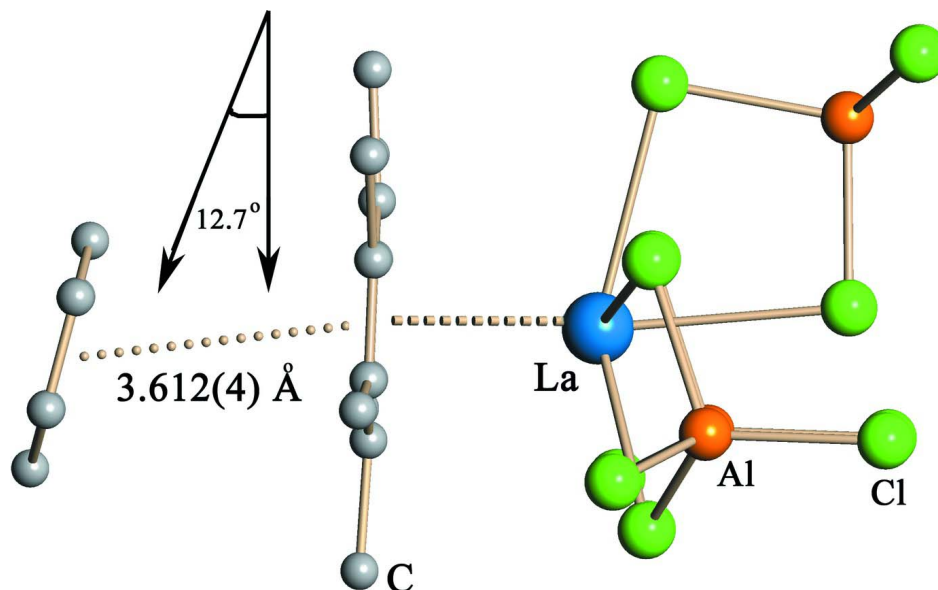


Figure 2

A view of the molecular structure of (I) showing  $\pi$ - $\pi$  stacking interactions between coordinated  $C_6Me_6$  and benzene rings.

### Hexa- $\mu$ -chlorido-hexachlorido( $\eta^6$ -hexamethylbenzene)trialuminium(III)lanthanum(III) benzene solvate

#### Crystal data

$[Al_3LaCl_{12}(C_{12}H_{18})] \cdot C_6H_6$

$M_r = 885.62$

Orthorhombic,  $Pnma$

Hall symbol: -P 2ac 2n

$a = 12.2127$  (6) Å

$b = 16.4205$  (8) Å

$c = 16.9790$  (8) Å

$V = 3404.9$  (3) Å<sup>3</sup>

$Z = 4$

$F(000) = 1728$

$D_x = 1.728$  Mg m<sup>-3</sup>

Mo  $K\alpha$  radiation,  $\lambda = 0.71073$  Å

Cell parameters from 7019 reflections

$\theta = 2.5$ – $28.2^\circ$

$\mu = 2.28$  mm<sup>-1</sup>

$T = 173$  K

Block, yellow

$0.22 \times 0.20 \times 0.16$  mm

#### Data collection

Bruker SMART APEX CCD area-detector diffractometer

Radiation source: fine-focus sealed tube

Graphite monochromator

$0.30^\circ$   $\omega$  scans

Absorption correction: multi-scan

(*SADABS*; Bruker, 2003)

$T_{\min} = 0.613$ ,  $T_{\max} = 0.697$

28535 measured reflections

4250 independent reflections

3911 reflections with  $I > 2\sigma(I)$

$R_{\text{int}} = 0.018$

$\theta_{\max} = 28.3^\circ$ ,  $\theta_{\min} = 2.1^\circ$

$h = -15 \rightarrow 16$

$k = -21 \rightarrow 21$

$l = -22 \rightarrow 22$

#### Refinement

Refinement on  $F^2$

Least-squares matrix: full

$R[F^2 > 2\sigma(F^2)] = 0.025$

$wR(F^2) = 0.063$

$S = 1.05$

4250 reflections

189 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 not refined

$$w = 1/[\sigma^2(F_o^2) + (0.0304P)^2 + 3.1621P]$$

where  $P = (F_o^2 + 2F_c^2)/3$   
 $(\Delta/\sigma)_{\max} < 0.001$

$$\Delta\rho_{\max} = 0.89 \text{ e } \text{\AA}^{-3}$$

$$\Delta\rho_{\min} = -0.99 \text{ e } \text{\AA}^{-3}$$

### 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 ( $\text{\AA}^2$ )

	<i>x</i>	<i>y</i>	<i>z</i>	$U_{\text{iso}}^*/U_{\text{eq}}$	Occ. (<1)
La1	0.106393 (12)	0.2500	0.425620 (9)	0.02758 (6)	
Al1	0.09537 (6)	0.04272 (4)	0.32343 (5)	0.04146 (16)	
Al2	-0.20200 (8)	0.2500	0.41914 (6)	0.0403 (2)	
Cl1	-0.02765 (6)	0.03298 (4)	0.23836 (5)	0.06224 (19)	
Cl2	0.18966 (7)	-0.06204 (4)	0.34109 (6)	0.0684 (2)	
Cl3	0.19872 (5)	0.14875 (3)	0.30139 (3)	0.04206 (13)	
Cl4	0.02586 (5)	0.08226 (4)	0.43571 (3)	0.04489 (14)	
Cl5	-0.09121 (6)	0.2500	0.52121 (4)	0.04250 (18)	
Cl6	-0.08346 (6)	0.2500	0.32208 (4)	0.03682 (16)	
Cl7	-0.29273 (6)	0.14181 (5)	0.41927 (4)	0.05841 (18)	
C1	0.3384 (3)	0.2500	0.4729 (2)	0.0515 (10)	
C2	0.3003 (3)	0.17691 (17)	0.50333 (19)	0.0609 (8)	
C3	0.2212 (3)	0.1777 (3)	0.5621 (2)	0.0823 (13)	
C4	0.1820 (3)	0.2500	0.5901 (2)	0.096 (3)	
C5	0.4271 (4)	0.2500	0.4107 (3)	0.194 (5)	
H5A	0.4789	0.2943	0.4214	0.291*	0.50
H5B	0.3940	0.2580	0.3587	0.291*	0.50
H5C	0.4659	0.1978	0.4118	0.291*	0.50
C6	0.3258 (8)	0.0879 (6)	0.4928 (7)	0.087 (3)	0.50
H6A	0.3781	0.0706	0.5332	0.130*	0.50
H6B	0.3577	0.0791	0.4405	0.130*	0.50
H6C	0.2583	0.0561	0.4977	0.130*	0.50
C7	0.1582 (8)	0.1206 (6)	0.6166 (5)	0.093 (3)	0.50
H7A	0.1812	0.1299	0.6711	0.139*	0.50
H7B	0.1735	0.0640	0.6019	0.139*	0.50
H7C	0.0796	0.1312	0.6117	0.139*	0.50
C6X	0.3773 (8)	0.1077 (7)	0.4617 (6)	0.084 (4)	0.50
H6X1	0.4398	0.0956	0.4959	0.126*	0.50
H6X2	0.4039	0.1281	0.4109	0.126*	0.50
H6X3	0.3344	0.0581	0.4533	0.126*	0.50
C7X	0.2146 (9)	0.0787 (5)	0.5824 (6)	0.092 (3)	0.50
H7X1	0.1425	0.0659	0.6045	0.137*	0.50

H7X2	0.2716	0.0646	0.6206	0.137*	0.50
H7X3	0.2257	0.0473	0.5339	0.137*	0.50
C8	0.1026 (4)	0.2500	0.6586 (3)	0.194 (5)	
H8A	0.0691	0.1960	0.6636	0.291*	0.50
H8B	0.0454	0.2907	0.6494	0.291*	0.50
H8C	0.1420	0.2633	0.7072	0.291*	0.50
C9	0.5948 (4)	0.2500	0.6119 (3)	0.088 (2)	
H9	0.6600	0.2500	0.5814	0.106*	
C10	0.5464 (3)	0.1771 (2)	0.6345 (2)	0.0750 (10)	
H10	0.5779	0.1265	0.6193	0.090*	
C11	0.4547 (3)	0.1790 (2)	0.6780 (2)	0.0679 (9)	
H11	0.4212	0.1293	0.6936	0.082*	
C12	0.4094 (3)	0.2500	0.7000 (3)	0.0646 (12)	
H12	0.3449	0.2500	0.7312	0.077*	

*Atomic displacement parameters (Å<sup>2</sup>)*

	$U^{11}$	$U^{22}$	$U^{33}$	$U^{12}$	$U^{13}$	$U^{23}$
La1	0.02175 (8)	0.03570 (10)	0.02528 (9)	0.000	0.00028 (5)	0.000
Al1	0.0428 (4)	0.0342 (3)	0.0475 (4)	-0.0003 (3)	-0.0093 (3)	-0.0026 (3)
Al2	0.0247 (4)	0.0577 (6)	0.0384 (5)	0.000	0.0008 (4)	0.000
Cl1	0.0664 (4)	0.0525 (4)	0.0678 (4)	0.0015 (3)	-0.0311 (3)	-0.0082 (3)
Cl2	0.0666 (4)	0.0438 (3)	0.0947 (6)	0.0136 (3)	-0.0232 (4)	-0.0036 (4)
Cl3	0.0434 (3)	0.0424 (3)	0.0404 (3)	-0.0030 (2)	0.0099 (2)	-0.0068 (2)
Cl4	0.0460 (3)	0.0431 (3)	0.0456 (3)	-0.0101 (2)	0.0013 (2)	0.0080 (2)
Cl5	0.0288 (3)	0.0687 (5)	0.0300 (3)	0.000	0.0031 (3)	0.000
Cl6	0.0277 (3)	0.0530 (4)	0.0298 (3)	0.000	-0.0025 (3)	0.000
Cl7	0.0418 (3)	0.0720 (5)	0.0615 (4)	-0.0171 (3)	0.0028 (3)	-0.0021 (3)
C1	0.0227 (14)	0.099 (3)	0.0325 (16)	0.000	-0.0035 (12)	0.000
C2	0.0654 (17)	0.0486 (14)	0.0688 (18)	0.0173 (13)	-0.0438 (16)	-0.0151 (13)
C3	0.071 (2)	0.114 (3)	0.0617 (19)	-0.055 (2)	-0.0416 (18)	0.055 (2)
C4	0.0269 (19)	0.237 (9)	0.0259 (18)	0.000	-0.0009 (14)	0.000
C5	0.0328 (17)	0.507 (15)	0.0414 (19)	0.000	0.0080 (14)	0.000
C6	0.082 (7)	0.060 (5)	0.118 (9)	0.034 (5)	-0.054 (6)	-0.025 (5)
C7	0.099 (7)	0.110 (7)	0.070 (5)	-0.061 (6)	-0.042 (5)	0.058 (5)
C6X	0.075 (6)	0.089 (8)	0.087 (7)	0.044 (6)	-0.037 (5)	-0.040 (6)
C7X	0.111 (8)	0.067 (5)	0.097 (7)	-0.032 (5)	-0.053 (6)	0.048 (5)
C8	0.0328 (17)	0.507 (15)	0.0414 (19)	0.000	0.0080 (14)	0.000
C9	0.052 (3)	0.172 (7)	0.041 (2)	0.000	0.0042 (19)	0.000
C10	0.081 (2)	0.080 (2)	0.0634 (19)	0.029 (2)	-0.0217 (18)	-0.0169 (17)
C11	0.070 (2)	0.069 (2)	0.0653 (19)	-0.0142 (17)	-0.0277 (16)	0.0189 (16)
C12	0.039 (2)	0.105 (4)	0.049 (2)	0.000	-0.0136 (17)	0.000

*Geometric parameters (Å, °)*

La1—C1	2.945 (3)	C3—C7X	1.663 (8)
La1—C2	2.965 (2)	C4—C3 <sup>i</sup>	1.366 (5)
La1—C3	2.957 (3)	C4—C8	1.514 (6)

La1—C4	2.941 (4)	C5—H5A	0.9800
La1—C13	2.9128 (5)	C5—H5B	0.9800
La1—C14	2.9298 (6)	C5—H5C	0.9800
La1—C15	2.9083 (7)	C8—H8A	0.9800
La1—C16	2.9097 (7)	C8—H8B	0.9800
La1—Cg1	2.613 (3)	C8—H8C	0.9800
Cg1—Cg2	3.612 (4)	C6—H6A	0.9800
La1—C13 <sup>i</sup>	2.9128 (5)	C6—H6B	0.9800
La1—C14 <sup>i</sup>	2.9298 (6)	C6—H6C	0.9800
La1—C3 <sup>i</sup>	2.957 (3)	C7—H7A	0.9800
La1—C2 <sup>i</sup>	2.965 (2)	C7—H7B	0.9800
A11—C11	2.0902 (10)	C7—H7C	0.9800
A11—C12	2.0918 (10)	C6X—H6X1	0.9800
A11—C13	2.1828 (9)	C6X—H6X2	0.9800
A11—C14	2.1855 (10)	C6X—H6X3	0.9800
A12—C17	2.0937 (9)	C7X—H7X1	0.9800
A12—C17 <sup>i</sup>	2.0937 (9)	C7X—H7X2	0.9800
A12—C16	2.1936 (12)	C7X—H7X3	0.9800
A12—C15	2.1987 (12)	C9—C10	1.389 (5)
C1—C2 <sup>i</sup>	1.387 (4)	C9—C10 <sup>i</sup>	1.389 (5)
C1—C2	1.387 (4)	C9—H9	0.9500
C1—C5	1.513 (6)	C10—C11	1.343 (5)
C2—C3	1.389 (5)	C10—H10	0.9500
C2—C6	1.505 (10)	C11—C12	1.344 (4)
C2—C6X	1.635 (10)	C11—H11	0.9500
C3—C4	1.366 (5)	C12—C11 <sup>i</sup>	1.344 (4)
C3—C7	1.525 (7)	C12—H12	0.9500
C15—La1—C16	71.09 (2)	C17—A12—C15	108.96 (4)
C15—La1—C13	136.497 (15)	C17 <sup>i</sup> —A12—C15	108.96 (4)
C16—La1—C13	82.586 (17)	C16—A12—C15	100.72 (5)
C15—La1—C13 <sup>i</sup>	136.497 (14)	A11—C13—La1	96.16 (3)
C16—La1—C13 <sup>i</sup>	82.586 (17)	A11—C14—La1	95.61 (3)
C13—La1—C13 <sup>i</sup>	69.61 (2)	A12—C15—La1	94.06 (4)
C15—La1—C14	71.868 (13)	A12—C16—La1	94.13 (4)
C16—La1—C14	76.576 (13)	C2 <sup>i</sup> —C1—C2	119.9 (4)
C13—La1—C14	68.636 (17)	C2 <sup>i</sup> —C1—C5	119.99 (18)
C13 <sup>i</sup> —La1—C14	135.147 (17)	C2—C1—C5	119.99 (18)
C15—La1—C14 <sup>i</sup>	71.868 (13)	C2 <sup>i</sup> —C1—La1	77.23 (17)
C16—La1—C14 <sup>i</sup>	76.576 (13)	C2—C1—La1	77.23 (17)
C13—La1—C14 <sup>i</sup>	135.147 (17)	C5—C1—La1	119.9 (3)
C13 <sup>i</sup> —La1—C14 <sup>i</sup>	68.636 (17)	C1—C2—C3	119.5 (3)
C14—La1—C14 <sup>i</sup>	140.15 (3)	C1—C2—C6	136.6 (6)
C15—La1—C4	74.37 (8)	C3—C2—C6	103.8 (6)
C16—La1—C4	145.46 (8)	C1—C2—La1	75.63 (16)
C13—La1—C4	124.47 (6)	C3—C2—La1	76.11 (16)
C13 <sup>i</sup> —La1—C4	124.47 (6)	C6—C2—La1	120.4 (4)
C14—La1—C4	92.86 (3)	C6X—C2—La1	123.2 (4)



Cl4 <sup>i</sup> —La1—C4	92.86 (3)	C4—C3—C2	120.1 (3)
Cl5—La1—C1	130.25 (7)	C4—C3—C7	98.4 (6)
Cl6—La1—C1	158.66 (7)	C2—C3—C7	141.3 (6)
Cl3—La1—C1	79.93 (6)	C4—C3—La1	76.0 (2)
Cl3 <sup>i</sup> —La1—C1	79.93 (6)	C2—C3—La1	76.76 (15)
Cl4—La1—C1	107.879 (16)	C7—C3—La1	118.9 (3)
Cl4 <sup>i</sup> —La1—C1	107.879 (17)	C3 <sup>i</sup> —C4—C3	120.8 (4)
C4—La1—C1	55.88 (10)	C3 <sup>i</sup> —C4—C8	119.5 (2)
Cl5—La1—C3 <sup>i</sup>	87.49 (8)	C3—C4—C8	119.5 (2)
Cl6—La1—C3 <sup>i</sup>	148.34 (6)	C3 <sup>i</sup> —C4—La1	77.2 (2)
Cl3—La1—C3 <sup>i</sup>	127.81 (6)	C3—C4—La1	77.2 (2)
Cl3 <sup>i</sup> —La1—C3 <sup>i</sup>	98.90 (10)	C8—C4—La1	121.9 (3)
Cl4—La1—C3 <sup>i</sup>	119.39 (10)	C1—C5—H5A	109.5
Cl4 <sup>i</sup> —La1—C3 <sup>i</sup>	74.68 (7)	C1—C5—H5B	109.5
C4—La1—C3 <sup>i</sup>	26.77 (10)	H5A—C5—H5B	109.5
C1—La1—C3 <sup>i</sup>	47.94 (8)	C1—C5—H5C	109.5
Cl5—La1—C3	87.49 (8)	H5A—C5—H5C	109.5
Cl6—La1—C3	148.34 (6)	H5B—C5—H5C	109.5
Cl3—La1—C3	98.90 (10)	C4—C8—H8A	109.5
Cl3 <sup>i</sup> —La1—C3	127.81 (6)	C4—C8—H8B	109.5
Cl4—La1—C3	74.68 (7)	H8A—C8—H8B	109.5
Cl4 <sup>i</sup> —La1—C3	119.39 (10)	C4—C8—H8C	109.5
C4—La1—C3	26.77 (10)	H8A—C8—H8C	109.5
C1—La1—C3	47.94 (8)	H8B—C8—H8C	109.5
C3 <sup>i</sup> —La1—C3	47.35 (17)	C2—C6—H6A	109.5
Cl5—La1—C2 <sup>i</sup>	114.49 (8)	C2—C6—H6B	109.5
Cl6—La1—C2 <sup>i</sup>	154.88 (5)	C2—C6—H6C	109.5
Cl3—La1—C2 <sup>i</sup>	104.12 (8)	C3—C7—H7A	109.5
Cl3 <sup>i</sup> —La1—C2 <sup>i</sup>	77.41 (6)	C3—C7—H7B	109.5
Cl4—La1—C2 <sup>i</sup>	128.51 (5)	C3—C7—H7C	109.5
Cl4 <sup>i</sup> —La1—C2 <sup>i</sup>	82.04 (6)	C2—C6X—H6X1	109.5
C4—La1—C2 <sup>i</sup>	47.68 (9)	C2—C6X—H6X2	109.5
C1—La1—C2 <sup>i</sup>	27.14 (7)	H6X1—C6X—H6X2	109.5
C3 <sup>i</sup> —La1—C2 <sup>i</sup>	27.13 (10)	C2—C6X—H6X3	109.5
C3—La1—C2 <sup>i</sup>	55.61 (8)	H6X1—C6X—H6X3	109.5
Cl5—La1—C2	114.49 (8)	H6X2—C6X—H6X3	109.5
Cl6—La1—C2	154.88 (5)	C3—C7X—H7X1	109.5
Cl3—La1—C2	77.41 (6)	C3—C7X—H7X2	109.5
Cl3 <sup>i</sup> —La1—C2	104.12 (8)	H7X1—C7X—H7X2	109.5
Cl4—La1—C2	82.04 (6)	C3—C7X—H7X3	109.5
Cl4 <sup>i</sup> —La1—C2	128.51 (5)	H7X1—C7X—H7X3	109.5
C4—La1—C2	47.68 (9)	H7X2—C7X—H7X3	109.5
C1—La1—C2	27.14 (7)	C10—C9—C10 <sup>i</sup>	119.1 (5)
C3 <sup>i</sup> —La1—C2	55.61 (8)	C10—C9—H9	120.5
C3—La1—C2	27.13 (10)	C10 <sup>i</sup> —C9—H9	120.5
C2 <sup>i</sup> —La1—C2	47.76 (11)	C11—C10—C9	119.2 (4)
Cl1—Al1—Cl2	115.58 (4)	C11—C10—H10	120.4
Cl1—Al1—Cl3	110.99 (4)	C9—C10—H10	120.4

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C12—A11—C13	111.24 (4)	C10—C11—C12	121.1 (4)
C11—A11—C14	110.26 (4)	C10—C11—H11	119.5
C12—A11—C14	109.45 (5)	C12—C11—H11	119.5
C13—A11—C14	97.89 (3)	C11 <sup>i</sup> —C12—C11	120.5 (5)
C17—A12—C17 <sup>i</sup>	116.10 (6)	C11 <sup>i</sup> —C12—H12	119.8
C17—A12—C16	110.49 (4)	C11—C12—H12	119.8
C17 <sup>i</sup> —A12—C16	110.49 (4)		

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Symmetry code: (i)  $x, -y+1/2, z$ .