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Bis[4-(dimethylamino)pyridinium] octaaquachloridolanthanum(III) tetrachloride trihydrate

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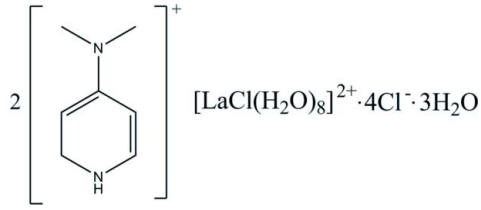
Key indicators: single-crystal X-ray study; T = 180 K; mean σ(C–C) = 0.004 Å; R factor = 0.023; wR factor = 0.056; data-to-parameter ratio = 22.7.

The title organic–inorganic salt, $(C_7H_{11}N_2)_2[LaCl(H_2O)_8]Cl_4\cdot 3H_2O$, consists of two 4-(dimethylamino)pyridinium and one $[La(H_2O)_8Cl]^{2+}$ cations, four chloride anions and three solvent water molecules. In the crystal, the various units are connected by N—H···Cl, O—H···Cl, O—H···O and N—H···O hydrogen bonds, forming a network of alternating organic and inorganic layers. The 4-(dimethylamino)pyridinium cations stack along the c axis, while the inorganic layers lie parallel to the ac plane. The chloride anions are located between these entities, forming hydrogen bonds with the NH atom of the pyridinium ions and the water molecules. There are also C—H···Cl hydrogen bonds present involving one of the 4-(dimethylamino)pyridinium cations, resulting in the formation of a three-dimensional supramolecular architecture.

Related literature

For common applications of organic–inorganic hybrid materials, see: Cui *et al.* (2000); Lacroix *et al.* (1994); Chakravarthy & Guloy (1997). For the crystal structures of compounds involving 4-(dimethylamino)pyridinium, see: Chao *et al.* (1977); Mayr-Stein & Bolte (2000); Lo & Ng (2008, 2009); Koon *et al.* (2009). For hydrogen-bond motifs, see: Bernstein *et al.* (1995).



Experimental

Crystal data

$(C_7H_{11}N_2)_2[LaCl(H_2O)_8]Cl_4\cdot 3H_2O$
M_r = 760.69
Triclinic, $P\overline{1}$
a = 9.6741 (4) Å
b = 12.6695 (7) Å
c = 14.3601 (7) Å
α = 68.354 (5)°
β = 75.273 (4)°
γ = 84.264 (4)°
V = 1582.16 (15) Å^3
Z = 2
Mo $K\alpha$ radiation
μ = 1.82 mm^{-1}
T = 180 K
0.43 × 0.28 × 0.08 mm

Data collection

Oxford Xcalibur Sapphire1 diffractometer
Absorption correction: multi-scan (*CrysAlis PRO*; Agilent, 2011)
T_{min} = 0.548, T_{max} = 0.864
33782 measured reflections
7144 independent reflections
6518 reflections with $I > 2\sigma(I)$
R_{int} = 0.038

Refinement

$R[F^2 > 2\sigma(F^2)]$ = 0.023
$wR(F^2)$ = 0.056
S = 1.11
7144 reflections
320 parameters
H-atom parameters constrained
$\Delta\rho_{max}$ = 0.76 e Å^{-3}
$\Delta\rho_{min}$ = −1.03 e Å^{-3}

Table 1
Hydrogen-bond geometry (Å, °).

D—H···A	D—H	H···A	D···A	D—H···A
N2—H2A···Cl5^{i}	0.86	2.71	3.314 (3)	129
N2—H2A···O1W^{ii}	0.86	2.24	2.909 (3)	134
N4—H4A···Cl4^{iii}	0.86	2.51	3.213 (2)	140
N4—H4A···Cl4^{iv}	0.86	2.77	3.418 (3)	133
O1—H11···Cl2^{v}	0.79	2.46	3.2316 (19)	165
O1—H12···O1W^{vi}	0.78	2.01	2.784 (2)	167
O2W—H12W···Cl3^{iii}	0.85	2.38	3.222 (2)	169
O3W—H13W···Cl2^{vii}	0.85	2.51	3.293 (2)	153
O2—H21···Cl3^{viii}	0.85	2.32	3.1537 (17)	166
O1W—H21W···Cl1	0.85	2.31	3.1538 (19)	173
O2—H22···Cl4	0.85	2.27	3.1023 (17)	168
O2W—H22W···Cl2	0.85	2.37	3.213 (2)	172
O3W—H23W···Cl2^{v}	0.85	2.34	3.193 (2)	176
O3—H31···Cl4^{viii}	0.84	2.34	3.1471 (17)	160
O3—H32···Cl1^{vi}	0.84	2.36	3.1413 (16)	157
O4—H41···Cl3^{viii}	0.84	2.31	3.1459 (17)	173
O4—H42···O2W	0.85	1.95	2.791 (3)	178
O5—H51···Cl1	0.84	2.38	3.1707 (19)	158
O5—H52···Cl2	0.85	2.43	3.241 (2)	160
O6—H61···Cl4	0.84	2.35	3.1708 (17)	164
O6—H62···Cl1	0.85	2.32	3.1250 (16)	158
O7—H71···Cl5^{v}	0.84	2.46	3.1402 (17)	139
O7—H72···Cl1	0.84	2.41	3.2287 (18)	162
O8—H81···O3W	0.85	1.94	2.786 (3)	172
O8—H82···Cl3	0.84	2.48	3.2711 (19)	156
C11—H11A···Cl3^{iii}	0.93	2.80	3.612 (3)	147
C14—H14B···Cl1	0.96	2.75	3.639 (4)	154

Symmetry codes: (i) $x-1, y+1, z$; (ii) $-x+1, -y+2, -z+1$; (iii) $x, y-1, z$; (iv) $-x+1, -y, -z+2$; (v) $-x+2, -y+1, -z+1$; (vi) $x+1, y, z$; (vii) $x, y+1, z$; (viii) $-x+2, -y+1, -z+2$.

Data collection: *CrysAlis PRO* (Agilent, 2011); cell refinement: *CrysAlis PRO*; data reduction: *CrysAlis PRO*; program(s) used to solve structure: *SIR92* (Altomare *et al.*, 1993); program(s) used to refine structure: *SHELXL97* (Sheldrick, 2008); molecular graphics: *ORTEPIII* (Burnett & Johnson, 1996) and *ORTEP-3 for Windows* (Farrugia, 2012); software used to prepare material for publication: *SHELXL97*.

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Supplementary data and figures for this paper are available from the IUCr electronic archives (Reference: SU2504).

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supporting information

Acta Cryst. (2012). E**68**, m1321–m1322 [doi:10.1107/S1600536812040901]

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S1. Comment

Organic–inorganic hybrid compounds are of great interest because of their special magnetic (Cui *et al.*, 2000), electronic (Lacroix *et al.*, 1994) and optoelectronic properties (Chakravarthy & Guloy, 1997). It is expected that the packing interactions that govern the crystal organization will be influenced by the features of the cations and anions, which in turn will affect specific properties of the solids. The supramolecular networks become especially interesting when the cation and anion can participate in hydrogen-bonding. As part of a study of the effect of cations and anions on the crystal structures of organic–inorganic compounds, we report herein on the crystal structure of the title compound. This type of hybrid material generally exhibits a structure consisting of alternating organic–inorganic layers, characterized by isolated anions as found with other compounds involving 4-(dimethylamino)pyridinium (Chao *et al.*, 1977; Mayr-Stein & Bolte, 2000; Lo and Ng, 2008, 2009; Koon *et al.*, 2009).

The title structure contains three cations, one inorganic $[La(H_2O)_8Cl]^{2+}$ cation and two independent monoprotonated 4-(dimethylamino)pyridinium cations, four chloride anions and three water molecules (Fig. 1). Atom La1 is coordinated by eight water molecules with distances ranging from 2.510 (1) to 2.588 (2) Å, and by one chloride ion with La1—Cl5 = 2.8829 (6) Å. The overall structure consists of layers stacked along the c axis. The chloride anions are located between the organic entities forming hydrogen bonds with the NH atoms of the pyridinium ions and the water molecules (Table 1).

Each Cl^- anion accepts hydrogen bonds which can be divided into two groups. The first group involves hydrogen bonds linking $Cl4^-$ with two organic cations via the pyridinium N4—H4A H atom (Table 1), generating centrosymmetric $R_2^2(4)$ motifs (Bernstein *et al.*, 1995) along the c axis at y = 1/2. The second 4-(dimethylamino)pyridinium molecule is linked to one $[La(H_2O)_8Cl]^{2+}$ cation through an intermolecular N2—H2A···$Cl5^i$ hydrogen bond [symmetry code: (i) *x* - 1, *y* + 1, *z*] which can be described by the graph-set motif D(3). The second type of hydrogen bond, in which the Cl^- anion is the acceptor, is a linkage between the water molecules (free and coordinated) and the Cl^- anion. The inorganic $[La(H_2O)_8Cl]^{2+}$ cations are indirectly linked via Cl^- anions through intermolecular O—H···Cl and O—H···O hydrogen bonds generating cycles $R_2^2(8)$ and $R_6^2(12)$, which connect cationic and anionic entities (Fig. 2 and Table 1).

In the 4-(dimethylamino)pyridinium cations the N—C bond linking the dimethylamino substituent to the pyridinium ring is characteristically short [1.321 (3) and 1.324 (3)Å]. The dimethylamino group lies close to the plane of the pyridinium ring with a dihedral angle, between the pyridinium and the dimethylamine plane (C/N/C atoms), of 3.5 (3) and 2.0 (3)°.

On the structural level, the atomic arrangement of this material consists of a network of alternating organic–inorganic layers. The chloride anions are located between these entities forming hydrogen bonds with the NH atoms of the pyridinium ions and the water molecules. There are also C—H···Cl interactions present (Table 1) involving one of the 4-(dimethylamino)pyridinium cations, which results in the formation of a three-dimensional supramolecular architecture.

S2. Experimental

4-(Dimethylamino)pyridine (1 mmol, 0.08g) and hydrochloric acid (1M) were added slowly to a solution of $LaCl_3.6H_2O$ (1mmol, 0.08g). The mixture was refluxed at 353 K for about 1 h and then cooled to room temperature. Slow evaporation of the solvent at room temperature lead to the formation of colourless plate-like crystals of the title compound.

S3. Refinement

The H atoms of the coordinated water molecules were located in difference Fourier syntheses and were initially refined using distance restraints: O—H = 0.85 (2) Å, and H···H = 1.40 (2) Å, with U_{iso}(H) = 1.5U_{eq}(O). In the final cycles of refinement they were constrained to ride on their parent O atoms. The N-bound H atoms were located in a difference Fourier map but like the C-bound H atoms they were included in calculated positions and treated as riding atoms: N—H = 0.86 Å, C—H = 0.93 and 0.96 Å for CH and CH_3 H atoms, respectively, with U_{iso}(H) = 1.5U_{eq}(C) for the methyl groups and = 1.2U_{eq}(N,C) for the other H atoms.

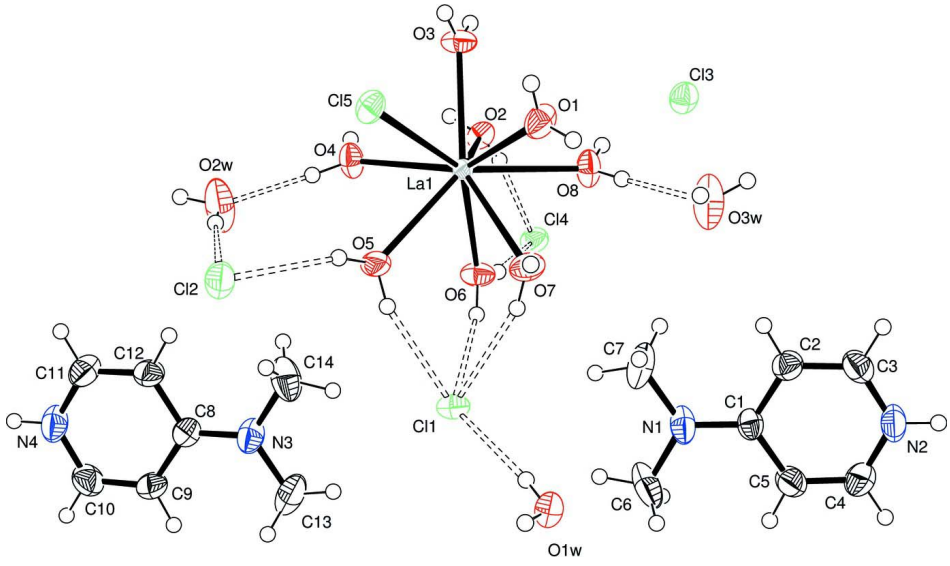

Figure 1

The molecular structure of the title compound, showing the atom-numbering. Displacement ellipsoids are drawn at the 50% probability level. The O—H···Cl hydrogen bonds are shown as double dashed lines.

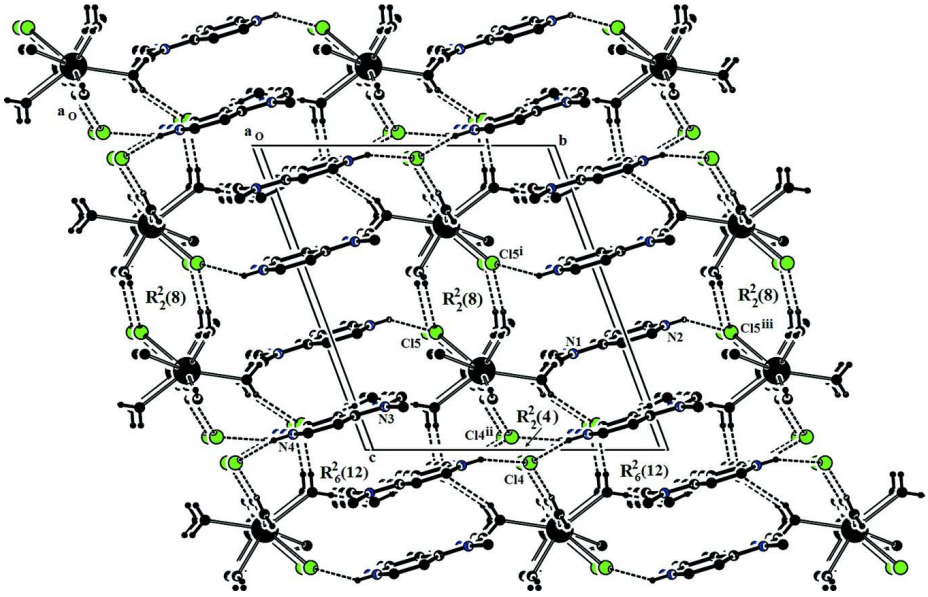

Figure 2

A view along the a axis of the three-dimensional hydrogen-bonded network of the title compound, showing the aggregation of the hydrogen-bonding motifs, $R_2^2(4)$, $R_2^2(8)$ and $R_6^2(12)$. Hydrogen bonds are drawn as dashed lines. [symmetry codes: (i) -x + 2, -y + 1, -z + 1; (ii) -x + 2, -y + 1, -z + 2; (iii) x - 1, y + 1, z].

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Crystal data

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Triclinic, $P\overline{1}$
Hall symbol: -P 1
a = 9.6741 (4) Å
b = 12.6695 (7) Å
c = 14.3601 (7) Å
α = 68.354 (5)°
β = 75.273 (4)°
γ = 84.264 (4)°
V = 1582.16 (15) Å^3
Z = 2
$F(000)$ = 772
D_x = 1.597 Mg m^{-3}
Mo $K\alpha$ radiation, λ = 0.71073 Å
Cell parameters from 22004 reflections
θ = 3.0–28.3°
μ = 1.82 mm^{-1}
T = 180 K
Plate, colourless
0.43 × 0.28 × 0.08 mm

Data collection

Oxford Xcalibur Sapphire1 diffractometer
Radiation source: Enhance (Mo) X-ray Source
Graphite monochromator
Detector resolution: 8.2632 pixels mm^{-1}
ω scans
Absorption correction: multi-scan (*CrysAlis PRO*; Agilent, 2011)
T_{min} = 0.548, T_{max} = 0.864
33782 measured reflections
7144 independent reflections
6518 reflections with $I > 2\sigma(I)$
R_{int} = 0.038
θ_{max} = 27.5°, θ_{min} = 3.0°
h = −12→12
k = −16→16
l = −18→18

Refinement

Refinement on F^2
Least-squares matrix: full
$R[F^2 > 2\sigma(F^2)] = 0.023$
$wR(F^2) = 0.056$
$S = 1.11$
7144 reflections
320 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
$w = 1/[\sigma^2(F_o^2) + (0.0257P)^2 + 0.4439P]$ where $P = (F_o^2 + 2F_c^2)/3$
$(\Delta/\sigma)_{max} = 0.003$
$\Delta\rho_{max} = 0.76$ e Å^{-3}
$\Delta\rho_{min} = -1.03$ e Å^{-3}

Special details

Geometry. Bond distances, angles etc. have been calculated using the rounded fractional coordinates. All su's are estimated from the variances of the (full) variance-covariance matrix. The cell esds are taken into account in the estimation of distances, angles and torsion angles

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 (Å²)

	x	*y*	*z*	U_{iso}*/U_{eq}
La1	0.98356 (1)	0.46700 (1)	0.74019 (1)	0.0192 (1)
Cl5	1.19118 (6)	0.36620 (5)	0.61364 (4)	0.0342 (2)
O1	1.12672 (17)	0.62452 (14)	0.58572 (13)	0.0428 (6)
O2	0.95865 (16)	0.45114 (13)	0.92689 (11)	0.0297 (5)
O3	1.21742 (15)	0.45676 (15)	0.78883 (12)	0.0346 (5)
O4	0.97883 (18)	0.26017 (13)	0.86085 (12)	0.0361 (5)
O5	0.85068 (17)	0.34195 (15)	0.68566 (14)	0.0436 (6)
O6	0.71719 (15)	0.46052 (14)	0.83325 (11)	0.0343 (5)
O7	0.83661 (17)	0.59127 (14)	0.61058 (12)	0.0374 (5)
O8	0.93133 (19)	0.65745 (14)	0.76941 (12)	0.0383 (6)
N1	0.4241 (2)	0.81063 (18)	0.68388 (17)	0.0442 (7)
N2	0.4146 (3)	1.15628 (18)	0.59093 (18)	0.0461 (8)
C1	0.4218 (2)	0.9227 (2)	0.65305 (18)	0.0322 (7)
C2	0.5477 (3)	0.9873 (2)	0.62318 (19)	0.0370 (8)
C3	0.5401 (3)	1.1014 (2)	0.5933 (2)	0.0416 (8)
C4	0.2929 (3)	1.0995 (2)	0.6179 (2)	0.0479 (9)
C5	0.2927 (3)	0.9862 (2)	0.6480 (2)	0.0436 (9)
C6	0.2940 (4)	0.7466 (3)	0.7103 (3)	0.0670 (11)
C7	0.5568 (4)	0.7456 (2)	0.6892 (3)	0.0618 (11)
N3	0.4436 (2)	0.07909 (18)	0.85825 (17)	0.0432 (7)
N4	0.4840 (2)	−0.26628 (19)	0.94889 (18)	0.0473 (8)
C8	0.4561 (2)	−0.0328 (2)	0.88707 (17)	0.0334 (7)
C9	0.3378 (2)	−0.1043 (2)	0.9155 (2)	0.0395 (8)
C10	0.3548 (3)	−0.2174 (2)	0.9445 (2)	0.0465 (9)
C11	0.5998 (3)	−0.2022 (2)	0.9229 (2)	0.0444 (8)
C12	0.5908 (2)	−0.0884 (2)	0.89126 (19)	0.0378 (8)

C13	0.3050 (3)	0.1370 (3)	0.8590 (3)	0.0613 (11)
C14	0.5691 (4)	0.1508 (3)	0.8259 (3)	0.0609 (11)
Cl1	0.54269 (5)	0.45268 (5)	0.68027 (4)	0.0344 (2)
Cl2	0.91549 (7)	0.13843 (5)	0.59452 (5)	0.0433 (2)
Cl3	0.97602 (7)	0.78867 (5)	0.91759 (4)	0.0388 (2)
Cl4	0.65967 (6)	0.50633 (5)	1.04257 (4)	0.0340 (2)
O1W	0.42022 (18)	0.63535 (14)	0.50183 (13)	0.0391 (5)
O2W	0.9449 (3)	0.06121 (17)	0.82838 (17)	0.0733 (9)
O3W	0.9016 (3)	0.86973 (17)	0.62175 (16)	0.0676 (8)
H11	1.09960	0.68050	0.54710	0.0640*
H12	1.20950	0.61670	0.56780	0.0640*
H21	0.99110	0.39140	0.96650	0.0450*
H22	0.88380	0.47160	0.96220	0.0450*
H31	1.22900	0.46190	0.84310	0.0520*
H32	1.29400	0.44680	0.74990	0.0520*
H41	0.99590	0.24250	0.91960	0.0540*
H42	0.96630	0.19990	0.85180	0.0540*
H51	0.76890	0.35990	0.67430	0.0650*
H52	0.88930	0.29500	0.65740	0.0650*
H61	0.68810	0.48120	0.88450	0.0510*
H62	0.65100	0.46360	0.80330	0.0510*
H71	0.87490	0.61120	0.54710	0.0560*
H72	0.75130	0.57010	0.62470	0.0560*
H81	0.92450	0.71900	0.72020	0.0570*
H82	0.95110	0.67060	0.81790	0.0570*
H2	0.63590	0.95080	0.62430	0.0440*
H2A	0.41240	1.22910	0.57180	0.0550*
H3	0.62340	1.14270	0.57390	0.0500*
H4	0.20700	1.13940	0.61560	0.0570*
H5	0.20670	0.94860	0.66590	0.0520*
H6A	0.22750	0.76040	0.76690	0.1010*
H6B	0.31660	0.66700	0.72960	0.1010*
H6C	0.25220	0.77020	0.65180	0.1010*
H7A	0.61200	0.75610	0.62060	0.0920*
H7B	0.53530	0.66650	0.72600	0.0920*
H7C	0.61060	0.77130	0.72440	0.0920*
H4A	0.49240	−0.33900	0.96850	0.0570*
H9	0.24700	−0.07270	0.91420	0.0470*
H10	0.27570	−0.26280	0.96190	0.0560*
H11A	0.68830	−0.23740	0.92680	0.0530*
H12A	0.67300	−0.04580	0.87200	0.0450*
H13A	0.25120	0.10850	0.82580	0.0920*
H13B	0.31940	0.21710	0.82290	0.0920*
H13C	0.25350	0.12350	0.92890	0.0920*
H14A	0.61670	0.13040	0.88130	0.0920*
H14B	0.53970	0.22900	0.80830	0.0920*
H14C	0.63330	0.14010	0.76690	0.0920*
H11W	0.43610	0.61070	0.45270	0.0590*

H21W	0.45310	0.59110	0.55210	0.0590*
H12W	0.96180	−0.00920	0.85550	0.1100*
H22W	0.94090	0.07510	0.76670	0.1100*
H13W	0.92930	0.93010	0.62480	0.1010*
H23W	0.94670	0.86620	0.56370	0.1010*

Atomic displacement parameters (Å²)

	U^{11}	U^{22}	U^{33}	U^{12}	U^{13}	U^{23}
La1	0.0199 (1)	0.0190 (1)	0.0202 (1)	0.0010 (1)	−0.0061 (1)	−0.0082 (1)
Cl5	0.0366 (3)	0.0363 (3)	0.0326 (3)	0.0084 (2)	−0.0054 (2)	−0.0195 (3)
O1	0.0333 (9)	0.0315 (10)	0.0442 (10)	0.0023 (7)	0.0006 (7)	0.0014 (8)
O2	0.0320 (8)	0.0342 (9)	0.0248 (7)	0.0070 (7)	−0.0090 (6)	−0.0129 (7)
O3	0.0220 (7)	0.0563 (11)	0.0345 (8)	0.0010 (7)	−0.0080 (6)	−0.0258 (8)
O4	0.0578 (11)	0.0232 (9)	0.0292 (8)	−0.0007 (7)	−0.0161 (7)	−0.0077 (7)
O5	0.0318 (8)	0.0492 (11)	0.0738 (13)	0.0131 (8)	−0.0257 (8)	−0.0438 (10)
O6	0.0251 (7)	0.0533 (11)	0.0307 (8)	0.0001 (7)	−0.0078 (6)	−0.0214 (8)
O7	0.0343 (8)	0.0452 (11)	0.0312 (8)	0.0027 (7)	−0.0141 (7)	−0.0083 (8)
O8	0.0621 (11)	0.0244 (9)	0.0325 (9)	0.0036 (8)	−0.0162 (8)	−0.0124 (7)
N1	0.0510 (13)	0.0267 (12)	0.0516 (13)	−0.0011 (10)	−0.0022 (10)	−0.0167 (10)
N2	0.0626 (15)	0.0242 (11)	0.0532 (14)	0.0006 (10)	−0.0205 (11)	−0.0114 (10)
C1	0.0363 (12)	0.0279 (12)	0.0336 (12)	0.0008 (10)	−0.0043 (9)	−0.0153 (10)
C2	0.0324 (12)	0.0374 (14)	0.0396 (13)	0.0019 (10)	−0.0075 (10)	−0.0130 (11)
C3	0.0449 (14)	0.0366 (15)	0.0413 (14)	−0.0113 (12)	−0.0094 (11)	−0.0093 (12)
C4	0.0444 (15)	0.0405 (16)	0.0643 (18)	0.0127 (12)	−0.0178 (13)	−0.0249 (14)
C5	0.0326 (12)	0.0411 (16)	0.0624 (17)	−0.0019 (11)	−0.0070 (12)	−0.0267 (14)
C6	0.078 (2)	0.0392 (18)	0.082 (2)	−0.0244 (16)	−0.0016 (18)	−0.0250 (17)
C7	0.080 (2)	0.0334 (16)	0.068 (2)	0.0189 (15)	−0.0171 (17)	−0.0187 (15)
N3	0.0473 (12)	0.0294 (12)	0.0466 (12)	0.0040 (10)	−0.0076 (10)	−0.0096 (10)
N4	0.0484 (13)	0.0284 (12)	0.0572 (14)	0.0009 (10)	−0.0025 (11)	−0.0133 (11)
C8	0.0348 (12)	0.0335 (13)	0.0300 (11)	0.0025 (10)	−0.0070 (9)	−0.0102 (10)
C9	0.0262 (11)	0.0425 (15)	0.0491 (15)	0.0028 (10)	−0.0077 (10)	−0.0170 (12)
C10	0.0385 (14)	0.0438 (17)	0.0563 (17)	−0.0104 (12)	−0.0038 (12)	−0.0193 (14)
C11	0.0366 (13)	0.0414 (16)	0.0491 (15)	0.0077 (11)	−0.0074 (11)	−0.0130 (13)
C12	0.0267 (11)	0.0382 (15)	0.0451 (14)	−0.0026 (10)	−0.0058 (10)	−0.0123 (12)
C13	0.072 (2)	0.0458 (18)	0.067 (2)	0.0261 (16)	−0.0248 (17)	−0.0222 (16)
C14	0.076 (2)	0.0347 (17)	0.0626 (19)	−0.0163 (15)	−0.0038 (16)	−0.0109 (15)
Cl1	0.0247 (2)	0.0445 (3)	0.0366 (3)	0.0009 (2)	−0.0099 (2)	−0.0158 (3)
Cl2	0.0529 (4)	0.0348 (3)	0.0459 (3)	0.0063 (3)	−0.0157 (3)	−0.0177 (3)
Cl3	0.0485 (3)	0.0358 (3)	0.0342 (3)	0.0040 (3)	−0.0174 (3)	−0.0110 (3)
Cl4	0.0317 (3)	0.0403 (3)	0.0372 (3)	0.0066 (2)	−0.0128 (2)	−0.0208 (3)
O1W	0.0455 (10)	0.0304 (9)	0.0404 (9)	0.0044 (7)	−0.0106 (8)	−0.0125 (8)
O2W	0.138 (2)	0.0342 (12)	0.0592 (13)	0.0006 (13)	−0.0433 (14)	−0.0167 (10)
O3W	0.1061 (18)	0.0352 (12)	0.0511 (12)	0.0020 (11)	−0.0040 (12)	−0.0139 (10)

Geometric parameters (Å, º)

La1—Cl5	2.8829 (6)	O1W—H11W	0.8500
La1—O1	2.5585 (17)	O1W—H21W	0.8500
La1—O2	2.5632 (15)	N4—H4A	0.8600
La1—O3	2.5101 (15)	O2W—H22W	0.8500
La1—O4	2.5505 (17)	O2W—H12W	0.8500
La1—O5	2.5710 (19)	O3W—H13W	0.8500
La1—O6	2.5775 (15)	O3W—H23W	0.8500
La1—O7	2.5885 (17)	C1—C2	1.418 (4)
La1—O8	2.5786 (19)	C1—C5	1.419 (4)
O1—H11	0.7900	C2—C3	1.348 (4)
O1—H12	0.7800	C4—C5	1.338 (4)
O2—H21	0.8500	C2—H2	0.9300
O2—H22	0.8500	C3—H3	0.9300
O3—H31	0.8400	C4—H4	0.9300
O3—H32	0.8400	C5—H5	0.9300
O4—H41	0.8400	C6—H6B	0.9600
O4—H42	0.8500	C6—H6C	0.9600
O5—H51	0.8400	C6—H6A	0.9600
O5—H52	0.8500	C7—H7C	0.9600
O6—H61	0.8400	C7—H7A	0.9600
O6—H62	0.8500	C7—H7B	0.9600
O7—H71	0.8400	C8—C9	1.412 (3)
O7—H72	0.8400	C8—C12	1.422 (3)
O8—H81	0.8500	C9—C10	1.342 (4)
O8—H82	0.8400	C11—C12	1.343 (4)
N1—C1	1.322 (4)	C9—H9	0.9300
N1—C6	1.456 (5)	C10—H10	0.9300
N1—C7	1.458 (4)	C11—H11A	0.9300
N2—C3	1.339 (4)	C12—H12A	0.9300
N2—C4	1.336 (4)	C13—H13B	0.9600
N2—H2A	0.8600	C13—H13A	0.9600
N3—C8	1.324 (4)	C13—H13C	0.9600
N3—C14	1.462 (5)	C14—H14C	0.9600
N3—C13	1.462 (4)	C14—H14A	0.9600
N4—C10	1.344 (4)	C14—H14B	0.9600
N4—C11	1.341 (4)		

Cl5—La1—O1	70.83 (4)	C4—N2—H2A	120.00
Cl5—La1—O2	130.08 (4)	C8—N3—C14	121.1 (2)
Cl5—La1—O3	72.34 (4)	C13—N3—C14	116.7 (3)
Cl5—La1—O4	78.77 (4)	C8—N3—C13	122.3 (2)
Cl5—La1—O5	71.63 (4)	C10—N4—C11	120.3 (3)
Cl5—La1—O6	142.42 (4)	H11W—O1W—H21W	112.00
Cl5—La1—O7	101.27 (4)	C11—N4—H4A	120.00
Cl5—La1—O8	139.80 (4)	C10—N4—H4A	120.00
O1—La1—O2	122.67 (5)	H12W—O2W—H22W	108.00

O1—La1—O3	77.88 (5)	H13W—O3W—H23W	107.00
O1—La1—O4	146.68 (6)	C2—C1—C5	115.6 (2)
O1—La1—O5	112.45 (6)	N1—C1—C2	122.3 (2)
O1—La1—O6	130.10 (5)	N1—C1—C5	122.0 (2)
O1—La1—O7	65.71 (5)	C1—C2—C3	120.3 (3)
O1—La1—O8	70.66 (6)	N2—C3—C2	121.1 (3)
O2—La1—O3	65.87 (5)	N2—C4—C5	121.1 (3)
O2—La1—O4	68.56 (5)	C1—C5—C4	120.9 (3)
O2—La1—O5	124.62 (6)	C3—C2—H2	120.00
O2—La1—O6	70.04 (5)	C1—C2—H2	120.00
O2—La1—O7	128.53 (5)	C2—C3—H3	119.00
O2—La1—O8	65.90 (5)	N2—C3—H3	119.00
O3—La1—O4	80.24 (6)	N2—C4—H4	119.00
O3—La1—O5	135.98 (6)	C5—C4—H4	119.00
O3—La1—O6	135.89 (5)	C4—C5—H5	119.00
O3—La1—O7	142.78 (6)	C1—C5—H5	120.00
O3—La1—O8	88.74 (6)	H6A—C6—H6C	109.00
O4—La1—O5	68.67 (6)	N1—C6—H6B	109.00
O4—La1—O6	82.76 (6)	N1—C6—H6A	109.00
O4—La1—O7	135.63 (6)	H6A—C6—H6B	109.00
O4—La1—O8	133.68 (5)	N1—C6—H6C	109.00
O5—La1—O6	71.22 (6)	H6B—C6—H6C	109.00
O5—La1—O7	69.44 (6)	H7B—C7—H7C	109.00
O5—La1—O8	135.28 (6)	N1—C7—H7A	109.00
O6—La1—O7	70.55 (5)	N1—C7—H7B	109.00
O6—La1—O8	74.49 (6)	H7A—C7—H7B	109.00
O7—La1—O8	72.67 (6)	H7A—C7—H7C	109.00
La1—O1—H11	130.00	N1—C7—H7C	110.00
La1—O1—H12	119.00	N3—C8—C9	122.6 (2)
H11—O1—H12	111.00	N3—C8—C12	121.6 (2)
La1—O2—H21	117.00	C9—C8—C12	115.8 (2)
La1—O2—H22	123.00	C8—C9—C10	120.8 (2)
H21—O2—H22	108.00	N4—C10—C9	121.3 (3)
La1—O3—H31	126.00	N4—C11—C12	121.5 (3)
La1—O3—H32	121.00	C8—C12—C11	120.3 (2)
H31—O3—H32	113.00	C10—C9—H9	120.00
La1—O4—H41	121.00	C8—C9—H9	120.00
La1—O4—H42	131.00	N4—C10—H10	119.00
H41—O4—H42	109.00	C9—C10—H10	119.00
La1—O5—H51	121.00	N4—C11—H11A	119.00
La1—O5—H52	126.00	C12—C11—H11A	119.00
H51—O5—H52	110.00	C11—C12—H12A	120.00
La1—O6—H61	122.00	C8—C12—H12A	120.00
La1—O6—H62	122.00	N3—C13—H13B	109.00
H61—O6—H62	112.00	H13A—C13—H13C	109.00
La1—O7—H71	119.00	N3—C13—H13C	109.00
La1—O7—H72	114.00	H13A—C13—H13B	110.00
H71—O7—H72	112.00	N3—C13—H13A	110.00

La1—O8—H81	121.00	H13B—C13—H13C	109.00
La1—O8—H82	124.00	N3—C14—H14C	110.00
H81—O8—H82	111.00	H14A—C14—H14C	109.00
C1—N1—C6	121.2 (2)	H14B—C14—H14C	109.00
C1—N1—C7	121.9 (2)	H14A—C14—H14B	109.00
C6—N1—C7	116.8 (3)	N3—C14—H14A	109.00
C3—N2—C4	121.0 (2)	N3—C14—H14B	109.00
C3—N2—H2A	119.00		

C6—N1—C1—C2	−177.6 (3)	C5—C1—C2—C3	0.5 (4)
C7—N1—C1—C2	−0.1 (4)	N1—C1—C2—C3	−179.3 (2)
C6—N1—C1—C5	2.6 (4)	C2—C1—C5—C4	−0.8 (4)
C7—N1—C1—C5	−179.9 (3)	N1—C1—C5—C4	179.1 (2)
C3—N2—C4—C5	0.0 (4)	C1—C2—C3—N2	−0.1 (4)
C4—N2—C3—C2	−0.2 (4)	N2—C4—C5—C1	0.5 (4)
C13—N3—C8—C9	3.4 (4)	C12—C8—C9—C10	0.1 (4)
C13—N3—C8—C12	−176.6 (3)	N3—C8—C9—C10	−179.9 (2)
C14—N3—C8—C12	2.2 (4)	N3—C8—C12—C11	178.6 (2)
C14—N3—C8—C9	−177.9 (3)	C9—C8—C12—C11	−1.4 (3)
C11—N4—C10—C9	−0.8 (4)	C8—C9—C10—N4	1.0 (4)
C10—N4—C11—C12	−0.5 (4)	N4—C11—C12—C8	1.6 (4)

Hydrogen-bond geometry (Å, º)

D—H···*A*	*D*—H	H···*A*	*D*···*A*	*D*—H···*A*
N2—H2*A*···Cl5^{i}	0.86	2.71	3.314 (3)	129
N2—H2*A*···O1*W*^{ii}	0.86	2.24	2.909 (3)	134
N4—H4*A*···Cl4^{iii}	0.86	2.51	3.213 (2)	140
N4—H4*A*···Cl4^{iv}	0.86	2.77	3.418 (3)	133
O1—H11···Cl2^{v}	0.79	2.46	3.2316 (19)	165
O1—H12···O1*W*^{vi}	0.78	2.01	2.784 (2)	167
O2*W*—H12*W*···Cl3^{iii}	0.85	2.38	3.222 (2)	169
O3*W*—H13*W*···Cl2^{vii}	0.85	2.51	3.293 (2)	153
O2—H21···Cl3^{viii}	0.85	2.32	3.1537 (17)	166
O1*W*—H21*W*···Cl1	0.85	2.31	3.1538 (19)	173
O2—H22···Cl4	0.85	2.27	3.1023 (17)	168
O2*W*—H22*W*···Cl2	0.85	2.37	3.213 (2)	172
O3*W*—H23*W*···Cl2^{v}	0.85	2.34	3.193 (2)	176
O3—H31···Cl4^{viii}	0.84	2.34	3.1471 (17)	160
O3—H32···Cl1^{vi}	0.84	2.36	3.1413 (16)	157
O4—H41···Cl3^{viii}	0.84	2.31	3.1459 (17)	173
O4—H42···O2*W*	0.85	1.95	2.791 (3)	178
O5—H51···Cl1	0.84	2.38	3.1707 (19)	158
O5—H52···Cl2	0.85	2.43	3.241 (2)	160
O6—H61···Cl4	0.84	2.35	3.1708 (17)	164
O6—H62···Cl1	0.85	2.32	3.1250 (16)	158
O7—H71···Cl5^{v}	0.84	2.46	3.1402 (17)	139
O7—H72···Cl1	0.84	2.41	3.2287 (18)	162

O8—H81···O3*W*	0.85	1.94	2.786 (3)	172
O8—H82···Cl3	0.84	2.48	3.2711 (19)	156
C11—H11*A*···Cl3^{iii}	0.93	2.80	3.612 (3)	147
C14—H14*B*···Cl1	0.96	2.75	3.639 (4)	154

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