

Dicaesium magnesium bis(dihydrogen phosphate(V)) dihydrate

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Key indicators: single-crystal X-ray study; $T = 173$ K; mean $\sigma(\text{P}-\text{O}) = 0.006$ Å; R factor = 0.048; wR factor = 0.125; data-to-parameter ratio = 12.3.

The title compound, $\text{Cs}_2\text{Mg}(\text{H}_2\text{P}_2\text{O}_7)_2 \cdot 2\text{H}_2\text{O}$, is isostructural with the related known isoformular phosphates. The crystal framework consists of corner-sharing MgO_6 and $\text{H}_2\text{P}_2\text{O}_7$ polyhedra, leading to tunnels parallel to the b -axis direction in which Cs^+ ions are located. The $\text{H}_2\text{P}_2\text{O}_7$ unit shows a bent eclipsed conformation. The Mg^{2+} ion lies on an inversion center. The water molecules form hydrogen bonds to O atoms of two different dihydrogenphosphate ions, which are further hydrogen bonded to symmetry-equivalent dihydrogenphosphate ions.

Related literature

For isostructural phosphates, see: Capitelli *et al.* (2004), $(\text{NH}_4)_2\text{Mn}(\text{H}_2\text{P}_2\text{O}_7)_2 \cdot 2\text{H}_2\text{O}$; Essehli *et al.* (2005a), $(\text{NH}_4)_2\text{Zn}(\text{H}_2\text{P}_2\text{O}_7)_2 \cdot 2\text{H}_2\text{O}$; Essehli *et al.* (2005b), $(\text{NH}_4)_2\text{Ni}(\text{H}_2\text{P}_2\text{O}_7)_2 \cdot 2\text{H}_2\text{O}$; Essehli *et al.* (2005c), $(\text{NH}_4)_2\text{Co}(\text{H}_2\text{P}_2\text{O}_7)_2 \cdot 2\text{H}_2\text{O}$; Tahiri *et al.* (2004), $\text{K}_2\text{Ni}(\text{H}_2\text{P}_2\text{O}_7)_2 \cdot 2\text{H}_2\text{O}$; Tahiri *et al.* (2003), $\text{K}_2\text{Zn}(\text{H}_2\text{P}_2\text{O}_7)_2 \cdot 2\text{H}_2\text{O}$; Harcharras *et al.* (2003), $\text{K}_2\text{Mg}(\text{H}_2\text{P}_2\text{O}_7)_2 \cdot 2\text{H}_2\text{O}$. For the biological activity of inorganic acidic diphosphates containing $\text{HP}_2\text{O}_7^{3-}$ or $\text{H}_2\text{P}_2\text{O}_7^{2-}$ anions, see: Andreeva *et al.* (2001).

Experimental

Crystal data

$\text{Cs}_2\text{Mg}(\text{H}_2\text{P}_2\text{O}_7)_2 \cdot 2\text{H}_2\text{O}$ $\alpha = 83.776$ (16) $^\circ$
 $M_r = 678.07$ $\beta = 68.558$ (15) $^\circ$
 Triclinic, $P\bar{1}$ $\gamma = 87.850$ (17) $^\circ$
 $a = 7.0935$ (15) Å $V = 392.87$ (14) Å³
 $b = 7.4606$ (15) Å $Z = 1$
 $c = 8.0230$ (15) Å Mo $K\alpha$ radiation

$\mu = 5.17$ mm⁻¹
 $T = 173$ (2) K

0.19 × 0.15 × 0.08 mm

Data collection

Stoe IPDSII two-circle diffractometer 3316 measured reflections
 1420 independent reflections
 Absorption correction: multi-scan (MULABS; Spek, 2003; Blessing, 1995) 1245 reflections with $I > 2\sigma(I)$
 $R_{\text{int}} = 0.082$
 $T_{\text{min}} = 0.440$, $T_{\text{max}} = 0.683$

Refinement

$R[F^2 > 2\sigma(F^2)] = 0.048$ H atoms treated by a mixture of independent and constrained refinement
 $wR(F^2) = 0.125$
 $S = 1.02$
 1420 reflections
 115 parameters
 3 restraints
 $\Delta\rho_{\text{max}} = 2.00$ e Å⁻³
 $\Delta\rho_{\text{min}} = -2.73$ e Å⁻³

Table 1

Hydrogen-bond geometry (Å, °).

$D-H \cdots A$	$D-H$	$H \cdots A$	$D \cdots A$	$D-H \cdots A$
$\text{O1W}-\text{H1WA} \cdots \text{O1}^{\text{i}}$	0.839 (10)	2.01 (4)	2.804 (8)	158 (9)
$\text{O1W}-\text{H1WB} \cdots \text{O6}^{\text{ii}}$	0.839 (10)	1.97 (3)	2.778 (9)	162 (9)
$\text{O3}-\text{H3} \cdots \text{O6}^{\text{ii}}$	0.84	1.72	2.551 (8)	172
$\text{O7}-\text{H7} \cdots \text{O1}^{\text{iii}}$	0.84	1.71	2.518 (8)	159

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

Data collection: *X-AREA* (Stoe & Cie, 2001); cell refinement: *X-AREA*; data reduction: *X-AREA*; program(s) used to solve structure: *SHELXS97* (Sheldrick, 2008); program(s) used to refine structure: *SHELXL97* (Sheldrick, 2008); molecular graphics: *XP* in *SHELXTL-Plus* (Sheldrick, 2008); software used to prepare material for publication: *SHELXL97*.

Supplementary data and figures for this paper are available from the IUCr electronic archives (Reference: BR2088).

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supplementary materials

Acta Cryst. (2009). E65, i3 [doi:10.1107/S1600536808042384]

Dicaesium magnesium bis(dihydrogen phosphate(V)) dihydrate

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Comment

Inorganic acidic diphosphates containing HP_2O_7 or $\text{H}_2\text{P}_2\text{O}_7$ hold important biochemical activities, such as inhibitors of human immunodeficiency enzymes as reported by Andreeva *et al.* (2001). In the framework of our systematic research on these phosphates, we report on the new compound $\text{Cs}_2\text{Mg}(\text{H}_2\text{P}_2\text{O}_7)_2 \cdot 2\text{H}_2\text{O}$. Detailed studies on structure determinations of such phosphates are available in related crystallography literature.

The crystal packing of $\text{Cs}_2\text{Mg}(\text{H}_2\text{P}_2\text{O}_7)_2 \cdot 2\text{H}_2\text{O}$ is a 3D network made upon edges sharing $[\text{MgO}_6]$ octahedra and dihydrogendiphosphate $[\text{H}_2\text{P}_2\text{O}_7]$. These delimitate tunnels along *b* direction, where Cs^+ ions are located. A projection onto *ac*-plan is depicted on Fig. 1.

Mg^{2+} cation sites are on inversion center. It is coordinated by four O atoms from two bidendate $[\text{H}_2\text{P}_2\text{O}_7]$ groups and two remaining O atoms from water molecule (Fig. 2).

$\text{H}_2\text{P}_2\text{O}_7$ shows bent eclipsed conformation. Distances and angles in $[\text{MgO}_6]$ and $[\text{H}_2\text{P}_2\text{O}_7]$ are as usual as in related phosphates structures. The $[\text{MgO}_6]$ are isolated in the structure, with an Mg-Mg distance over 7 Å.

Experimental

Crystals of $\text{Cs}_2\text{Mn}(\text{H}_2\text{P}_2\text{O}_7)_2 \cdot 2\text{H}_2\text{O}$ were grown at room temperature by slow evaporation from water-ethanol (80/20) of aqueous solution containing a stoichiometric the mixture : $\text{MgCl}_2 \cdot 6\text{H}_2\text{O}$ (0.231 mg, 1 mmol), Cs_2CO_3 (0.24 mg, 1 mmol), and $\text{K}_4\text{P}_2\text{O}_7$ (0.5 mg, 1 mmol). The solution was stirred for two hours at leaved to stand at room temperature. Crystals suitable for X-ray analysis were formed after few days.

Refinement

All H atoms were located in a difference map. The water H atoms were refined with the O-H bonds restrained to 0.84 (1) Å and the H...H distances restrained to 1.4 (1) Å and with fixed individual displacement parameters [$U(\text{H}) = 1.2 U_{\text{eq}}(\text{O})$]. The H atoms of the hydroxyl groups bonded to P were refined using a riding model with O-H = 0.84 Å, $U(\text{H}) = 1.2 U_{\text{eq}}(\text{O})$ and P-O-H = 109.5 °.

Figures

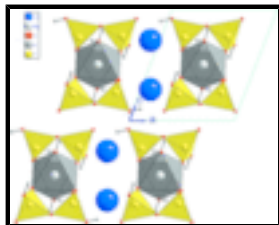


Fig. 1. Crystal structure of $\text{Cs}_2\text{Mn}(\text{H}_2\text{P}_2\text{O}_7)_2 \cdot 2\text{H}_2\text{O}$ viewed along b direction.

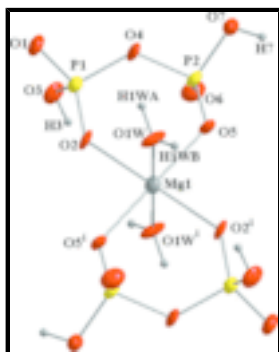


Fig. 2. Mg coordination in $\text{Cs}_2\text{Mn}(\text{H}_2\text{P}_2\text{O}_7)_2 \cdot 2\text{H}_2\text{O}$. Displacement ellipsoids are drawn at the 50% probability level. [Symmetry code: (i) $-x, -y, -z$.

Dicaesium magnesium bis(dihydrogen phosphate) dihydrate

Crystal data

$\text{Cs}_2\text{Mg}(\text{H}_2\text{P}_2\text{O}_7)_2 \cdot 2\text{H}_2\text{O}$

$M_r = 678.07$

Triclinic, $P\bar{1}$

Hall symbol: $-P\ 1$

$a = 7.0935(15)\ \text{\AA}$

$b = 7.4606(15)\ \text{\AA}$

$c = 8.0230(15)\ \text{\AA}$

$\alpha = 83.776(16)^\circ$

$\beta = 68.558(15)^\circ$

$\gamma = 87.850(17)^\circ$

$V = 392.87(14)\ \text{\AA}^3$

$Z = 1$

$F_{000} = 318$

$D_x = 2.866\ \text{Mg m}^{-3}$

Mo $K\alpha$ radiation

$\lambda = 0.71073\ \text{\AA}$

Cell parameters from 3316 reflections

$\theta = 3.7\text{--}25.5^\circ$

$\mu = 5.17\ \text{mm}^{-1}$

$T = 173(2)\ \text{K}$

Plate, colourless

$0.19 \times 0.15 \times 0.08\ \text{mm}$

Data collection

Stoe IPDSII two-circle diffractometer

Radiation source: fine-focus sealed tube

Monochromator: graphite

$T = 173(2)\ \text{K}$

ω scans

Absorption correction: multi-scan (MULABS; Spek, 2003; Blessing, 1995)

$T_{\min} = 0.440$, $T_{\max} = 0.683$

1420 independent reflections

1245 reflections with $I > 2\sigma(I)$

$R_{\text{int}} = 0.082$

$\theta_{\text{max}} = 25.3^\circ$

$\theta_{\text{min}} = 3.7^\circ$

$h = -8 \rightarrow 8$

$k = -8 \rightarrow 8$

3316 measured reflections

$l = -9 \rightarrow 9$

Refinement

Refinement on F^2

Secondary atom site location: difference Fourier map

Least-squares matrix: full

Hydrogen site location: inferred from neighbouring sites

$R[F^2 > 2\sigma(F^2)] = 0.048$

H atoms treated by a mixture of independent and constrained refinement

$wR(F^2) = 0.125$

$$w = 1/[\sigma^2(F_o^2) + (0.084P)^2]$$

where $P = (F_o^2 + 2F_c^2)/3$

$S = 1.02$

$(\Delta/\sigma)_{\max} < 0.001$

1420 reflections

$\Delta\rho_{\max} = 2.00 \text{ e } \text{\AA}^{-3}$

115 parameters

$\Delta\rho_{\min} = -2.73 \text{ e } \text{\AA}^{-3}$

3 restraints

Extinction correction: SHELXL97 (Sheldrick, 2008),

$$F_c^* = kF_c[1 + 0.001 \times F_c^2 \lambda^3 / \sin(2\theta)]^{-1/4}$$

Primary atom site location: structure-invariant direct methods

Extinction coefficient: 0.011 (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 (\AA^2)

	x	y	z	$U_{\text{iso}}^*/U_{\text{eq}}$
Cs1	0.09044 (7)	0.70386 (7)	0.27841 (6)	0.0186 (3)
Mg1	0.5000	0.5000	0.5000	0.0137 (8)
O1W	0.4211 (9)	0.2941 (8)	0.7146 (7)	0.0195 (14)
H1WA	0.413 (14)	0.319 (13)	0.817 (6)	0.023*
H1WB	0.355 (13)	0.201 (8)	0.719 (12)	0.023*
P1	0.3736 (3)	0.2528 (3)	0.2388 (3)	0.0129 (5)
P2	0.7759 (3)	0.1985 (3)	0.2518 (3)	0.0130 (5)
O1	0.3041 (8)	0.3164 (9)	0.0855 (8)	0.0190 (13)
O2	0.3344 (8)	0.3767 (8)	0.3829 (7)	0.0151 (12)
O3	0.2791 (9)	0.0628 (8)	0.3238 (8)	0.0181 (13)
H3	0.2828	0.0417	0.4276	0.022*
O4	0.6144 (8)	0.2214 (8)	0.1485 (7)	0.0162 (12)
O5	0.7572 (8)	0.3505 (8)	0.3636 (7)	0.0155 (12)
O6	0.7434 (9)	0.0105 (8)	0.3514 (8)	0.0216 (14)

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O7	0.9737 (9)	0.2074 (9)	0.0826 (8)	0.0201 (14)
H7	1.0655	0.2565	0.1046	0.024*

Atomic displacement parameters (\AA^2)

	U^{11}	U^{22}	U^{33}	U^{12}	U^{13}	U^{23}
Cs1	0.0193 (4)	0.0231 (4)	0.0164 (4)	0.0023 (2)	-0.0101 (2)	-0.0027 (2)
Mg1	0.0124 (18)	0.018 (2)	0.0124 (18)	0.0022 (15)	-0.0069 (15)	-0.0011 (15)
O1W	0.030 (4)	0.022 (3)	0.010 (3)	-0.003 (3)	-0.012 (3)	-0.002 (2)
P1	0.0121 (10)	0.0171 (11)	0.0121 (10)	0.0017 (8)	-0.0073 (8)	-0.0031 (8)
P2	0.0131 (10)	0.0190 (11)	0.0104 (9)	0.0015 (8)	-0.0080 (8)	-0.0035 (8)
O1	0.016 (3)	0.029 (3)	0.017 (3)	-0.001 (2)	-0.012 (2)	-0.002 (3)
O2	0.013 (3)	0.021 (3)	0.017 (3)	0.000 (2)	-0.012 (2)	-0.006 (2)
O3	0.022 (3)	0.017 (3)	0.021 (3)	-0.005 (2)	-0.012 (3)	-0.005 (2)
O4	0.013 (3)	0.027 (3)	0.014 (3)	0.003 (2)	-0.011 (2)	-0.005 (2)
O5	0.014 (3)	0.025 (3)	0.012 (3)	0.002 (2)	-0.008 (2)	-0.009 (2)
O6	0.027 (3)	0.022 (3)	0.017 (3)	-0.002 (3)	-0.010 (3)	-0.001 (2)
O7	0.012 (3)	0.037 (4)	0.014 (3)	-0.001 (3)	-0.006 (2)	-0.008 (3)

Geometric parameters (\AA , $^\circ$)

Cs1—O7 ⁱ	3.092 (6)	O1W—Cs1 ^{iv}	3.608 (6)
Cs1—O3 ⁱⁱ	3.151 (6)	O1W—H1WA	0.839 (10)
Cs1—O2	3.155 (6)	O1W—H1WB	0.839 (10)
Cs1—O6 ⁱⁱⁱ	3.233 (7)	P1—O2	1.498 (6)
Cs1—O2 ^{iv}	3.259 (6)	P1—O1	1.510 (6)
Cs1—O4 ⁱ	3.295 (5)	P1—O3	1.566 (6)
Cs1—O5 ^v	3.401 (5)	P1—O4	1.612 (6)
Cs1—O5 ^{vi}	3.450 (6)	P1—Cs1 ^{iv}	4.100 (2)
Cs1—O1	3.461 (6)	P2—O5	1.493 (6)
Cs1—O1W ^v	3.486 (6)	P2—O6	1.518 (6)
Cs1—O1W ^{iv}	3.608 (6)	P2—O7	1.553 (6)
Cs1—P1	3.836 (2)	P2—O4	1.637 (5)
Mg1—O2	2.046 (5)	P2—Cs1 ⁱ	3.995 (2)
Mg1—O2 ^v	2.046 (5)	O2—Cs1 ^{iv}	3.259 (6)
Mg1—O5 ^v	2.103 (6)	O3—Cs1 ^{viii}	3.151 (6)
Mg1—O5	2.103 (6)	O3—H3	0.8400
Mg1—O1W ^v	2.103 (5)	O4—Cs1 ⁱ	3.295 (5)
Mg1—O1W	2.103 (5)	O5—Cs1 ^v	3.401 (5)
Mg1—Cs1 ^v	4.0953 (9)	O5—Cs1 ^{vii}	3.450 (6)
Mg1—Cs1 ^{vii}	4.1789 (11)	O6—Cs1 ^{ix}	3.233 (6)
Mg1—Cs1 ^{iv}	4.1790 (11)	O7—Cs1 ⁱ	3.092 (6)
O1W—Cs1 ^v	3.486 (6)	O7—H7	0.8400
O7 ⁱ —Cs1—O3 ⁱⁱ	103.03 (16)	O5—Mg1—Cs1 ^v	56.01 (15)
O7 ⁱ —Cs1—O2	127.15 (16)	O1W ^v —Mg1—Cs1 ^v	121.65 (16)

O3 ⁱⁱ —Cs1—O2	108.02 (15)	O1W—Mg1—Cs1 ^v	58.35 (16)
O7 ⁱ —Cs1—O6 ⁱⁱⁱ	74.51 (15)	O2—Mg1—Cs1	48.97 (16)
O3 ⁱⁱ —Cs1—O6 ⁱⁱⁱ	72.09 (16)	O2 ^v —Mg1—Cs1	131.03 (16)
O2—Cs1—O6 ⁱⁱⁱ	155.89 (14)	O5 ^v —Mg1—Cs1	56.01 (15)
O7 ⁱ —Cs1—O2 ^{iv}	112.39 (14)	O5—Mg1—Cs1	123.99 (15)
O3 ⁱⁱ —Cs1—O2 ^{iv}	108.69 (15)	O1W ^v —Mg1—Cs1	58.35 (16)
O2—Cs1—O2 ^{iv}	96.77 (14)	O1W—Mg1—Cs1	121.65 (16)
O6 ⁱⁱⁱ —Cs1—O2 ^{iv}	61.80 (15)	Cs1 ^v —Mg1—Cs1	180.000 (7)
O7 ⁱ —Cs1—O4 ⁱ	44.14 (14)	O2—Mg1—Cs1 ^{vii}	130.16 (15)
O3 ⁱⁱ —Cs1—O4 ⁱ	84.93 (15)	O2 ^v —Mg1—Cs1 ^{vii}	49.84 (16)
O2—Cs1—O4 ⁱ	97.36 (14)	O5 ^v —Mg1—Cs1 ^{vii}	124.62 (16)
O6 ⁱⁱⁱ —Cs1—O4 ⁱ	106.59 (14)	O5—Mg1—Cs1 ^{vii}	55.38 (16)
O2 ^{iv} —Cs1—O4 ⁱ	156.16 (13)	O1W ^v —Mg1—Cs1 ^{vii}	59.70 (16)
O7 ⁱ —Cs1—O5 ^v	169.36 (14)	O1W—Mg1—Cs1 ^{vii}	120.30 (16)
O3 ⁱⁱ —Cs1—O5 ^v	68.88 (15)	Cs1 ^v —Mg1—Cs1 ^{vii}	61.976 (19)
O2—Cs1—O5 ^v	52.76 (15)	Cs1—Mg1—Cs1 ^{vii}	118.02 (2)
O6 ⁱⁱⁱ —Cs1—O5 ^v	108.24 (14)	O2—Mg1—Cs1 ^{iv}	49.84 (16)
O2 ^{iv} —Cs1—O5 ^v	77.41 (13)	O2 ^v —Mg1—Cs1 ^{iv}	130.16 (16)
O4 ⁱ —Cs1—O5 ^v	126.34 (13)	O5 ^v —Mg1—Cs1 ^{iv}	55.38 (16)
O7 ⁱ —Cs1—O5 ^{vi}	86.75 (14)	O5—Mg1—Cs1 ^{iv}	124.62 (16)
O3 ⁱⁱ —Cs1—O5 ^{vi}	160.22 (15)	O1W ^v —Mg1—Cs1 ^{iv}	120.30 (16)
O2—Cs1—O5 ^{vi}	78.07 (14)	O1W—Mg1—Cs1 ^{iv}	59.70 (16)
O6 ⁱⁱⁱ —Cs1—O5 ^{vi}	94.58 (15)	Cs1 ^v —Mg1—Cs1 ^{iv}	118.024 (19)
O2 ^{iv} —Cs1—O5 ^{vi}	51.54 (14)	Cs1—Mg1—Cs1 ^{iv}	61.98 (2)
O4 ⁱ —Cs1—O5 ^{vi}	113.40 (13)	Cs1 ^{vii} —Mg1—Cs1 ^{iv}	180.0
O5 ^v —Cs1—O5 ^{vi}	103.09 (12)	Mg1—O1W—Cs1 ^v	90.75 (19)
O7 ⁱ —Cs1—O1	82.39 (15)	Mg1—O1W—Cs1 ^{iv}	90.08 (18)
O3 ⁱⁱ —Cs1—O1	132.65 (15)	Cs1 ^v —O1W—Cs1 ^{iv}	178.20 (18)
O2—Cs1—O1	45.25 (13)	Mg1—O1W—H1WA	119 (7)
O6 ⁱⁱⁱ —Cs1—O1	150.16 (14)	Cs1 ^v —O1W—H1WA	72 (7)
O2 ^{iv} —Cs1—O1	112.32 (15)	Cs1 ^{iv} —O1W—H1WA	106 (7)
O4 ⁱ —Cs1—O1	66.40 (15)	Mg1—O1W—H1WB	124 (6)
O5 ^v —Cs1—O1	97.95 (14)	Cs1 ^v —O1W—H1WB	125 (7)
O5 ^{vi} —Cs1—O1	65.07 (13)	Cs1 ^{iv} —O1W—H1WB	55 (7)
O7 ⁱ —Cs1—O1W ^v	119.58 (14)	H1WA—O1W—H1WB	113 (8)
O3 ⁱⁱ —Cs1—O1W ^v	59.65 (15)	O2—P1—O1	116.7 (4)
O2—Cs1—O1W ^v	51.99 (14)	O2—P1—O3	110.1 (3)
O6 ⁱⁱⁱ —Cs1—O1W ^v	131.44 (15)	O1—P1—O3	108.9 (3)
O2 ^{iv} —Cs1—O1W ^v	128.02 (13)	O2—P1—O4	108.7 (3)
O4 ⁱ —Cs1—O1W ^v	75.64 (13)	O1—P1—O4	106.0 (3)
O5 ^v —Cs1—O1W ^v	50.71 (13)	O3—P1—O4	106.0 (3)

supplementary materials

O5 ^{vi} —Cs1—O1W ^v	130.02 (14)	O2—P1—Cs1	52.5 (2)
O1—Cs1—O1W ^v	76.60 (14)	O1—P1—Cs1	64.3 (3)
O7 ⁱ —Cs1—O1W ^{iv}	61.98 (13)	O3—P1—Cs1	126.4 (2)
O3 ⁱⁱ —Cs1—O1W ^{iv}	119.49 (15)	O4—P1—Cs1	127.4 (2)
O2—Cs1—O1W ^{iv}	128.11 (13)	O2—P1—Cs1 ^{iv}	46.6 (2)
O6 ⁱⁱⁱ —Cs1—O1W ^{iv}	47.53 (15)	O1—P1—Cs1 ^{iv}	110.0 (2)
O2 ^{iv} —Cs1—O1W ^{iv}	50.41 (12)	O3—P1—Cs1 ^{iv}	69.8 (2)
O4 ⁱ —Cs1—O1W ^{iv}	105.98 (12)	O4—P1—Cs1 ^{iv}	143.1 (2)
O5 ^v —Cs1—O1W ^{iv}	127.65 (13)	Cs1—P1—Cs1 ^{iv}	64.84 (4)
O5 ^{vi} —Cs1—O1W ^{iv}	50.22 (14)	O5—P2—O6	116.1 (3)
O1—Cs1—O1W ^{iv}	104.72 (14)	O5—P2—O7	112.9 (3)
O1W ^v —Cs1—O1W ^{iv}	178.20 (18)	O6—P2—O7	110.9 (4)
O7 ⁱ —Cs1—P1	105.41 (12)	O5—P2—O4	110.8 (3)
O3 ⁱⁱ —Cs1—P1	122.63 (12)	O6—P2—O4	106.4 (3)
O2—Cs1—P1	22.12 (11)	O7—P2—O4	98.0 (3)
O6 ⁱⁱⁱ —Cs1—P1	164.02 (12)	O5—P2—Cs1 ⁱ	120.3 (2)
O2 ^{iv} —Cs1—P1	104.76 (11)	O6—P2—Cs1 ⁱ	123.5 (3)
O4 ⁱ —Cs1—P1	82.42 (11)	O4—P2—Cs1 ⁱ	53.5 (2)
O5 ^v —Cs1—P1	74.87 (11)	P1—O1—Cs1	92.5 (3)
O5 ^{vi} —Cs1—P1	69.55 (10)	P1—O2—Mg1	137.3 (4)
O1—Cs1—P1	23.15 (9)	P1—O2—Cs1	105.4 (3)
O1W ^v —Cs1—P1	63.00 (11)	Mg1—O2—Cs1	101.8 (2)
O1W ^{iv} —Cs1—P1	117.80 (10)	P1—O2—Cs1 ^{iv}	113.8 (3)
O2—Mg1—O2 ^v	179.999 (1)	Mg1—O2—Cs1 ^{iv}	101.5 (2)
O2—Mg1—O5 ^v	89.5 (2)	Cs1—O2—Cs1 ^{iv}	83.23 (14)
O2 ^v —Mg1—O5 ^v	90.5 (2)	P1—O3—Cs1 ^{viii}	148.3 (3)
O2—Mg1—O5	90.5 (2)	P1—O3—H3	109.5
O2 ^v —Mg1—O5	89.5 (2)	Cs1 ^{viii} —O3—H3	102.2
O5 ^v —Mg1—O5	179.999 (1)	P1—O4—P2	126.8 (4)
O2—Mg1—O1W ^v	89.7 (2)	P1—O4—Cs1 ⁱ	128.3 (3)
O2 ^v —Mg1—O1W ^v	90.3 (2)	P2—O4—Cs1 ⁱ	103.0 (2)
O5 ^v —Mg1—O1W ^v	89.1 (2)	P2—O5—Mg1	129.5 (3)
O5—Mg1—O1W ^v	90.9 (2)	P2—O5—Cs1 ^v	120.1 (3)
O2—Mg1—O1W	90.3 (2)	Mg1—O5—Cs1 ^v	93.14 (17)
O2 ^v —Mg1—O1W	89.7 (2)	P2—O5—Cs1 ^{vii}	127.4 (3)
O5 ^v —Mg1—O1W	90.9 (2)	Mg1—O5—Cs1 ^{vii}	94.5 (2)
O5—Mg1—O1W	89.1 (2)	Cs1 ^v —O5—Cs1 ^{vii}	76.91 (12)
O1W ^v —Mg1—O1W	180.0 (3)	P2—O6—Cs1 ^{ix}	123.7 (3)
O2—Mg1—Cs1 ^v	131.03 (16)	P2—O7—Cs1 ⁱ	114.5 (3)
O2 ^v —Mg1—Cs1 ^v	48.97 (16)	P2—O7—H7	109.5
O5 ^v —Mg1—Cs1 ^v	123.99 (15)	Cs1 ⁱ —O7—H7	122.0

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

Hydrogen-bond geometry (\AA , $^\circ$)

$D-H\cdots A$	$D-H$	$H\cdots A$	$D\cdots A$	$D-H\cdots A$
O1W—H1WA \cdots O1 ^x	0.839 (10)	2.01 (4)	2.804 (8)	158 (9)
O1W—H1WB \cdots O6 ^{xi}	0.839 (10)	1.97 (3)	2.778 (9)	162 (9)
O3—H3 \cdots O6 ^{xi}	0.84	1.72	2.551 (8)	172
O7—H7 \cdots O1 ^{vii}	0.84	1.71	2.518 (8)	159

Symmetry codes: (x) $x, y, z+1$; (xi) $-x+1, -y, -z+1$; (vii) $x+1, y, z$.

Fig. 1

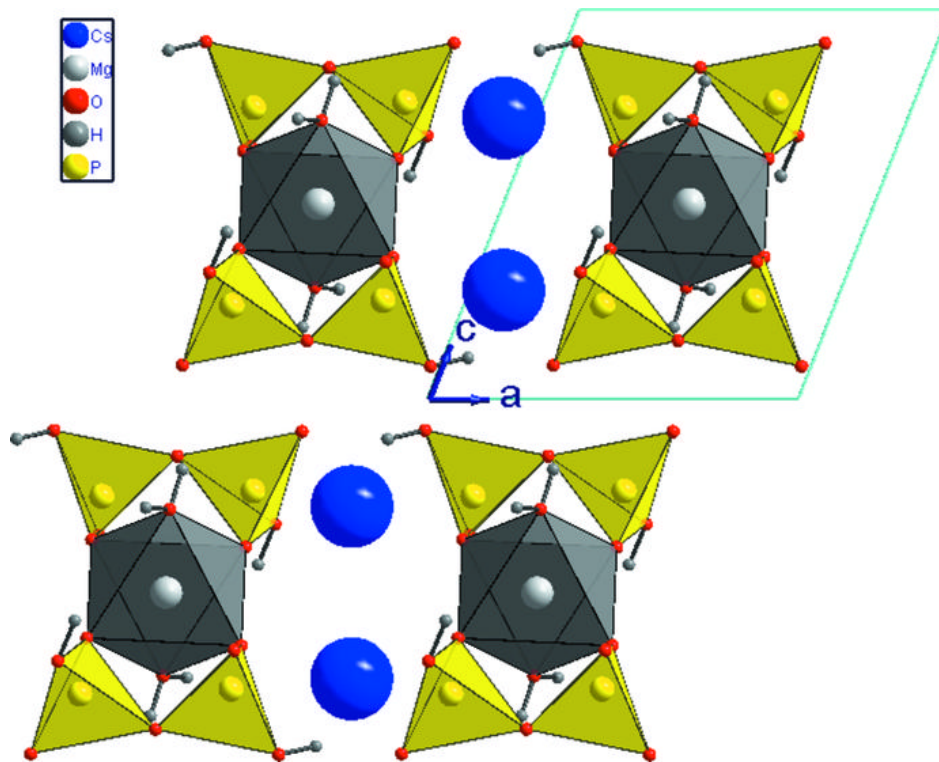


Fig. 2

