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2-[(*E*)-2-(4-Chlorophenyl)ethenyl]-1-methylpyridinium iodide monohydrate¹

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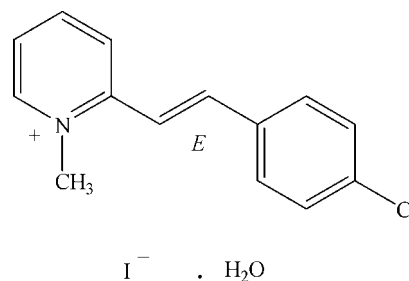
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Key indicators: single-crystal X-ray study; $T = 100$ K; mean $\sigma(\text{C}-\text{C}) = 0.005$ Å; R factor = 0.038; wR factor = 0.104; data-to-parameter ratio = 25.9.

In the title compound, $\text{C}_{14}\text{H}_{13}\text{ClN}^+\cdot\text{I}^-\cdot\text{H}_2\text{O}$, the cation is nearly planar and exists in an *E* configuration; the dihedral angle between the pyridinium and benzene rings is 0.98 (17)°. The cations stack in an anti-parallel manner along the *a* axis through two π - π interactions between the pyridinium and benzene rings [centroid-centroid distances 3.569 (2) and 3.6818 (13) Å, respectively]. The cation, anion and water molecule are linked into a chain along the *a* axis by weak $\text{C}-\text{H}\cdots\text{O}$ and $\text{C}-\text{H}\cdots\text{I}$ interactions together with $\text{O}-\text{H}\cdots\text{I}$ hydrogen bonds and the chains are further connected into a three-dimensional network.

Related literature

For bond-length data, see: Allen *et al.* (1987). For related structures, see, for example: Chantrapromma *et al.* (2007*a,b,c*). For background on non-linear optical properties, see, for example: Lakshmanaperumal *et al.* (2004); Marder *et al.* (1994); Qiu *et al.* (2007); Williams (1984); Zhai *et al.* (1999); Zhan *et al.* (2006).



Experimental

Crystal data

$\text{C}_{14}\text{H}_{13}\text{ClN}^+\cdot\text{I}^-\cdot\text{H}_2\text{O}$

$M_r = 375.62$

Monoclinic, $P2_1/c$

$a = 7.0876$ (1) Å

$b = 9.8096$ (2) Å

$c = 21.0940$ (4) Å

$\beta = 95.147$ (1)°

$V = 1460.68$ (5) Å³

$Z = 4$

Mo $K\alpha$ radiation

$\mu = 2.36$ mm⁻¹

$T = 100.0$ (1) K

$0.28 \times 0.17 \times 0.07$ mm

Data collection

Bruker SMART APEXII CCD

area-detector diffractometer

Absorption correction: multi-scan

(*SADABS*; Bruker, 2005)

$T_{\min} = 0.560$, $T_{\max} = 0.845$

18928 measured reflections

4241 independent reflections

3486 reflections with $I > 2\sigma(I)$

$R_{\text{int}} = 0.038$

Refinement

$R[F^2 > 2\sigma(F^2)] = 0.037$

$wR(F^2) = 0.104$

$S = 1.13$

4241 reflections

164 parameters

H-atom parameters constrained

$\Delta\rho_{\text{max}} = 2.13$ e Å⁻³

$\Delta\rho_{\text{min}} = -0.79$ e Å⁻³

Table 1

Hydrogen-bond geometry (Å, °).

$D-\text{H}\cdots A$	$D-\text{H}$	$\text{H}\cdots A$	$D\cdots A$	$D-\text{H}\cdots A$
$\text{O1W}-\text{H1W1}\cdots\text{I1}$	0.86	2.87	3.592 (4)	143
$\text{O1W}-\text{H2W1}\cdots\text{I1}^i$	0.85	2.87	3.567 (4)	141
$\text{C14}-\text{H14A}\cdots\text{O1W}$	0.96	2.50	3.202 (5)	130
$\text{C14}-\text{H14D}\cdots\text{O1W}^{ii}$	0.96	2.56	3.460 (5)	157
$\text{C1}-\text{H1A}\cdots\text{I1}^{iii}$	0.93	3.20	3.830 (4)	127
$\text{C2}-\text{H2A}\cdots\text{I1}^{iv}$	0.93	3.17	3.825 (4)	129
$\text{C3}-\text{H3A}\cdots\text{I1}^{iv}$	0.93	3.21	3.840 (4)	127

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

Data collection: *APEX2* (Bruker, 2005); cell refinement: *APEX2*; data reduction: *SAINT* (Bruker, 2005); program(s) used to solve structure: *SHELXTL* (Sheldrick, 2008); program(s) used to refine structure: *SHELXTL*; molecular graphics: *SHELXTL*; software used to prepare material for publication: *SHELXTL* and *PLATON* (Spek, 2003).

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Supplementary data and figures for this paper are available from the IUCr electronic archives (Reference: IS2329).

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supplementary materials

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2-[(*E*)-2-(4-Chlorophenyl)ethenyl]-1-methylpyridinium iodide monohydrate

K. Chanawanno, S. Chantrapromma and H.-K. Fun

Comment

In the last two decades, many efforts were focused on the discovery of new organic materials which exhibit large nonlinear optical (NLO) properties and would have applications in the fields of optoelectronics and photonics (Lakshmanaperumal *et al.*, 2004; Marder *et al.*, 1994; Qiu *et al.*, 2007; Zhai *et al.*, 1999; Zhan *et al.*, 2006). In order to obtain second-order NLO single crystals, the main requirements should be the choice of molecules with large hyperpolarizability (β) and the alignment of these molecules with optimal orientation into a noncentrosymmetric space group in the crystal (Williams, 1984). Among the known organic NLO materials, ionic chromophores are of great interest because they exhibit large first hyperpolarizabilities (β) and have high melting points and hardness of their crystals. At the molecular level, a generally popular approach towards NLO materials is to design and synthesize compounds with extended conjugated π -systems with donor and acceptor groups because such compounds are likely to exhibit large values of molecular hyperpolarizability (β) and to possess polarization. Styryl pyridinium derivatives are considered to be good conjugated π -systems. In continuation of our on-going research on nonlinear optical materials (Chantrapromma *et al.*, 2007*a,b,c*), the title compound, (I), was synthesized and the X-ray structure analysis was carried out in order to obtain detailed information about the molecular packing. However, compound (I) crystallizes in monoclinic space group $P2_1/c$ and doesn't exhibit second-order nonlinear optic properties.

The asymmetric unit of the title compound consists of $C_{14}H_{13}ClN^+$ cation, I^- anion and one water molecule (Fig. 1). The conformation of the cation is essentially planar as indicated by the dihedral angle between the pyridinium (N1/C1—C5) and the benzene (C8—C13) rings, being $0.98(17)^\circ$. The mean plane through C5/C6/C7/C8 plane makes dihedral angles of $6.1(4)^\circ$ and $6.4(4)^\circ$ with pyridinium and benzene rings, respectively. The cation exists in the *E* configuration and the torsion angle C5—C6—C7—C8 = $-179.2(3)^\circ$. The bond distances and angles in (I) have normal values (Allen *et al.*, 1987) and comparable with closely related structures (Chantrapromma *et al.*, 2007*a,b,c*).

The packing of the molecule down the *c* axis (Fig. 2), showing that the cation is linked with water molecule by weak C—H \cdots O interactions (Table 1) and linked with I^- anions by weak C—H \cdots I interactions (Table 1) whereas the I^- anion is linked with water molecule by O—H \cdots I hydrogen bonds, forming one-dimensional chains along the *a* axis. These chains are further connected into a three-dimensional network (Fig. 2). $\pi\cdots\pi$ interactions involving pyridinium and benzene rings were also observed with $Cg_1\cdots Cg_2$ distances of $3.662(2)$ Å (symmetry code; $1 - x, -y, 1 - z$) and $3.569(2)$ Å (symmetry code; $2 - x, -y, 1 - z$); Cg_1 and Cg_2 are the centroids of the N1/C1—C5 pyridinium and C8—C13 benzene rings, respectively. The crystal is stabilized by O—H \cdots I hydrogen bond, weak C—H \cdots O and C—H \cdots I interactions.

Experimental

The title compound was prepared by mixing solutions of 1,2-dimethylpyridinium iodide, 4-chlorobenzaldehyde and piperidine (1:1:1 molar ratio) in methanol. The resulting solution was refluxed for 12 hr under a nitrogen atmosphere. The solid which formed was filtered and washed with chloroform. Orange plate-like single-crystal suitable for X-ray diffraction

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analysis was obtained by recrystallization from methanol by slow evaporation of the solvent at ambient temperature after several days, Mp. 492–493 K.

Refinement

All H atoms were placed in calculated positions (O—H = 0.85–0.86 and C—H = 0.93–0.96 Å) and refined as riding, with $U_{\text{iso}}(\text{H}) = 1.2U_{\text{eq}}(\text{C})$ or $1.5U_{\text{eq}}(\text{O, methyl C})$. A rotating group model was used for the methyl groups. The highest residual electron density peak is located at 0.76 Å from I1 and the deepest hole is located at 0.59 Å from I1.

Figures

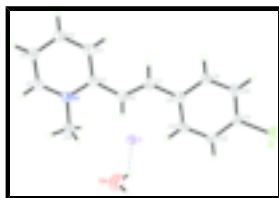


Fig. 1. The title compound showing 50% probability displacement ellipsoids and the atom-numbering scheme. The O—H...I hydrogen bond was drawn as dashed line.

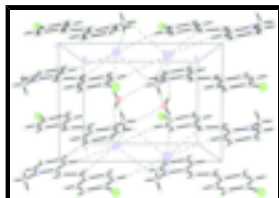


Fig. 2. The packing diagram of the title structure viewed approximately along the *c* axis. Hydrogen bonds were drawn as dashed lines.

2-[(*E*)-2-(4-Chlorophenyl)ethenyl]-1-methylpyridinium iodide monohydrate

Crystal data

$\text{C}_{14}\text{H}_{13}\text{ClN}^+\text{I}^-\text{H}_2\text{O}$

$M_r = 375.62$

Monoclinic, $P2_1/c$

Hall symbol: -P 2ybc

$a = 7.0876$ (1) Å

$b = 9.8096$ (2) Å

$c = 21.0940$ (4) Å

$\beta = 95.147$ (1)°

$V = 1460.68$ (5) Å³

$Z = 4$

$F_{000} = 736$

$D_x = 1.708$ Mg m⁻³

Melting point: 492-493 K

Mo $K\alpha$ radiation

$\lambda = 0.71073$ Å

Cell parameters from 4241 reflections

$\theta = 1.9$ – 30.0°

$\mu = 2.36$ mm⁻¹

$T = 100.0$ (1) K

Plate, orange

$0.28 \times 0.17 \times 0.07$ mm

Data collection

Bruker SMART APEXII CCD area-detector diffractometer

Radiation source: fine-focus sealed tube

Monochromator: graphite

Detector resolution: 8.33 pixels mm⁻¹

4241 independent reflections

3486 reflections with $I > 2\sigma(I)$

$R_{\text{int}} = 0.038$

$\theta_{\text{max}} = 30.0^\circ$

$T = 100.0(1)$ K $\theta_{\min} = 1.9^\circ$
 ω scans $h = -9 \rightarrow 9$
 Absorption correction: multi-scan (SADABS; Bruker, 2005) $k = -13 \rightarrow 12$
 $T_{\min} = 0.561$, $T_{\max} = 0.845$ $l = -26 \rightarrow 29$
 18928 measured reflections

Refinement

Refinement on F^2 Secondary atom site location: difference Fourier map
 Least-squares matrix: full Hydrogen site location: inferred from neighbouring sites
 $R[F^2 > 2\sigma(F^2)] = 0.037$ H-atom parameters constrained
 $wR(F^2) = 0.104$ $w = 1/[\sigma^2(F_o^2) + (0.0483P)^2 + 1.1431P]$
 where $P = (F_o^2 + 2F_c^2)/3$
 $S = 1.13$ $(\Delta/\sigma)_{\max} = 0.001$
 4241 reflections $\Delta\rho_{\max} = 2.13 \text{ e } \text{\AA}^{-3}$
 164 parameters $\Delta\rho_{\min} = -0.79 \text{ e } \text{\AA}^{-3}$
 Primary atom site location: structure-invariant direct methods Extinction correction: none

Special details

Experimental. The data was collected with the Oxford Cyrosystem Cobra low-temperature attachment.

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 (\AA^2)

	x	y	z	$U_{\text{iso}}^*/U_{\text{eq}}$
I1	0.96440 (3)	0.30947 (2)	0.374487 (10)	0.02517 (9)
Cl1	0.65015 (14)	-0.33018 (11)	0.25791 (5)	0.0349 (2)
N1	0.7517 (4)	0.2927 (3)	0.58820 (14)	0.0228 (6)
O1W	0.4685 (5)	0.3289 (4)	0.39663 (18)	0.0589 (10)
H1W1	0.5617	0.2949	0.3782	0.088*
H2W1	0.3691	0.2881	0.3808	0.088*
C1	0.7754 (5)	0.3535 (4)	0.64637 (17)	0.0254 (7)
H1A	0.7521	0.4464	0.6498	0.030*
C2	0.8329 (5)	0.2805 (4)	0.69966 (18)	0.0277 (8)
H2A	0.8473	0.3228	0.7393	0.033*
C3	0.8698 (5)	0.1412 (4)	0.69383 (17)	0.0262 (7)

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H3A	0.9115	0.0900	0.7294	0.031*
C4	0.8440 (5)	0.0817 (4)	0.63559 (17)	0.0264 (7)
H4A	0.8674	-0.0111	0.6318	0.032*
C5	0.7828 (5)	0.1569 (4)	0.58078 (17)	0.0221 (7)
C6	0.7492 (5)	0.0969 (4)	0.51828 (17)	0.0256 (7)
H6A	0.7187	0.1548	0.4839	0.031*
C7	0.7592 (5)	-0.0353 (4)	0.50694 (17)	0.0268 (7)
H7A	0.7883	-0.0912	0.5421	0.032*
C8	0.7293 (5)	-0.1035 (4)	0.44465 (16)	0.0235 (7)
C9	0.7663 (5)	-0.2435 (4)	0.44164 (18)	0.0266 (7)
H9A	0.8075	-0.2903	0.4786	0.032*
C10	0.7425 (5)	-0.3135 (4)	0.38414 (19)	0.0271 (8)
H10A	0.7680	-0.4063	0.3824	0.033*
C11	0.6808 (5)	-0.2432 (4)	0.33021 (17)	0.0238 (7)
C12	0.6421 (5)	-0.1050 (4)	0.33059 (17)	0.0258 (7)
H12A	0.6005	-0.0596	0.2932	0.031*
C13	0.6668 (5)	-0.0354 (4)	0.38847 (17)	0.0250 (7)
H13A	0.6414	0.0575	0.3896	0.030*
C14	0.6940 (5)	0.3805 (4)	0.53292 (18)	0.0298 (8)
H14D	0.6802	0.4727	0.5470	0.045*
H14A	0.5754	0.3489	0.5126	0.045*
H14B	0.7889	0.3769	0.5032	0.045*

Atomic displacement parameters (\AA^2)

	U^{11}	U^{22}	U^{33}	U^{12}	U^{13}	U^{23}
II	0.02744 (14)	0.02228 (14)	0.02596 (14)	0.00264 (9)	0.00324 (9)	0.00111 (9)
Cl1	0.0368 (5)	0.0385 (6)	0.0293 (5)	-0.0051 (4)	0.0027 (4)	-0.0126 (4)
N1	0.0206 (13)	0.0229 (16)	0.0249 (15)	0.0010 (11)	0.0027 (11)	0.0046 (12)
O1W	0.0347 (17)	0.083 (3)	0.058 (2)	0.0104 (16)	-0.0028 (15)	-0.028 (2)
C1	0.0246 (17)	0.0238 (18)	0.0278 (18)	0.0003 (14)	0.0028 (14)	0.0000 (15)
C2	0.0255 (17)	0.032 (2)	0.0261 (18)	-0.0004 (14)	0.0028 (14)	-0.0020 (15)
C3	0.0209 (16)	0.031 (2)	0.0263 (17)	0.0017 (14)	-0.0021 (13)	0.0030 (15)
C4	0.0218 (16)	0.0267 (19)	0.0307 (18)	-0.0010 (14)	0.0017 (14)	0.0034 (15)
C5	0.0175 (15)	0.0230 (18)	0.0265 (17)	-0.0009 (12)	0.0050 (12)	-0.0007 (14)
C6	0.0265 (17)	0.0245 (19)	0.0258 (17)	0.0002 (14)	0.0018 (14)	0.0001 (14)
C7	0.0272 (17)	0.0261 (19)	0.0268 (18)	0.0021 (14)	0.0002 (14)	-0.0019 (15)
C8	0.0227 (16)	0.0244 (19)	0.0236 (16)	-0.0012 (13)	0.0027 (13)	-0.0021 (14)
C9	0.0288 (18)	0.0231 (19)	0.0272 (18)	0.0000 (14)	-0.0009 (14)	0.0014 (15)
C10	0.0266 (17)	0.0195 (18)	0.035 (2)	-0.0016 (14)	0.0038 (15)	0.0009 (15)
C11	0.0203 (16)	0.0253 (19)	0.0257 (17)	-0.0029 (13)	0.0025 (13)	-0.0064 (14)
C12	0.0233 (16)	0.027 (2)	0.0267 (17)	0.0026 (14)	-0.0011 (13)	0.0029 (14)
C13	0.0246 (16)	0.0172 (17)	0.0329 (19)	0.0022 (13)	0.0013 (14)	0.0009 (14)
C14	0.037 (2)	0.025 (2)	0.0281 (18)	0.0052 (15)	0.0056 (15)	0.0032 (15)

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

Cl1—C11	1.744 (4)	C6—H6A	0.9300
N1—C1	1.361 (5)	C7—C8	1.473 (5)

N1—C5	1.361 (5)	C7—H7A	0.9300
N1—C14	1.478 (5)	C8—C13	1.397 (5)
O1W—H1W1	0.8628	C8—C9	1.400 (5)
O1W—H2W1	0.8523	C9—C10	1.391 (5)
C1—C2	1.364 (5)	C9—H9A	0.9300
C1—H1A	0.9300	C10—C11	1.368 (5)
C2—C3	1.399 (6)	C10—H10A	0.9300
C2—H2A	0.9300	C11—C12	1.383 (5)
C3—C4	1.358 (5)	C12—C13	1.396 (5)
C3—H3A	0.9300	C12—H12A	0.9300
C4—C5	1.407 (5)	C13—H13A	0.9300
C4—H4A	0.9300	C14—H14D	0.9600
C5—C6	1.444 (5)	C14—H14A	0.9600
C6—C7	1.322 (5)	C14—H14B	0.9600
C1—N1—C5	121.6 (3)	C13—C8—C9	118.5 (3)
C1—N1—C14	117.3 (3)	C13—C8—C7	123.3 (3)
C5—N1—C14	121.0 (3)	C9—C8—C7	118.2 (3)
H1W1—O1W—H2W1	106.3	C10—C9—C8	121.0 (3)
N1—C1—C2	121.1 (4)	C10—C9—H9A	119.5
N1—C1—H1A	119.4	C8—C9—H9A	119.5
C2—C1—H1A	119.4	C11—C10—C9	118.8 (3)
C1—C2—C3	119.0 (4)	C11—C10—H10A	120.6
C1—C2—H2A	120.5	C9—C10—H10A	120.6
C3—C2—H2A	120.5	C10—C11—C12	122.5 (3)
C4—C3—C2	119.2 (3)	C10—C11—C11	119.1 (3)
C4—C3—H3A	120.4	C12—C11—C11	118.4 (3)
C2—C3—H3A	120.4	C11—C12—C13	118.4 (3)
C3—C4—C5	121.7 (4)	C11—C12—H12A	120.8
C3—C4—H4A	119.2	C13—C12—H12A	120.8
C5—C4—H4A	119.2	C12—C13—C8	120.9 (3)
N1—C5—C4	117.4 (3)	C12—C13—H13A	119.6
N1—C5—C6	119.3 (3)	C8—C13—H13A	119.6
C4—C5—C6	123.4 (3)	N1—C14—H14D	109.5
C7—C6—C5	123.9 (3)	N1—C14—H14A	109.5
C7—C6—H6A	118.0	H14D—C14—H14A	109.5
C5—C6—H6A	118.0	N1—C14—H14B	109.5
C6—C7—C8	127.0 (4)	H14D—C14—H14B	109.5
C6—C7—H7A	116.5	H14A—C14—H14B	109.5
C8—C7—H7A	116.5		
C5—N1—C1—C2	0.5 (5)	C5—C6—C7—C8	-179.2 (3)
C14—N1—C1—C2	-178.4 (3)	C6—C7—C8—C13	-6.4 (6)
N1—C1—C2—C3	0.7 (5)	C6—C7—C8—C9	173.2 (4)
C1—C2—C3—C4	-1.3 (5)	C13—C8—C9—C10	0.3 (5)
C2—C3—C4—C5	0.6 (5)	C7—C8—C9—C10	-179.4 (3)
C1—N1—C5—C4	-1.1 (5)	C8—C9—C10—C11	-0.4 (5)
C14—N1—C5—C4	177.7 (3)	C9—C10—C11—C12	0.3 (5)
C1—N1—C5—C6	177.9 (3)	C9—C10—C11—C11	-179.9 (3)
C14—N1—C5—C6	-3.2 (5)	C10—C11—C12—C13	-0.2 (5)

supplementary materials

C3—C4—C5—N1	0.6 (5)	C11—C11—C12—C13	-179.9 (3)
C3—C4—C5—C6	-178.4 (3)	C11—C12—C13—C8	0.0 (5)
N1—C5—C6—C7	-173.9 (3)	C9—C8—C13—C12	-0.1 (5)
C4—C5—C6—C7	5.1 (6)	C7—C8—C13—C12	179.6 (3)

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

$D-H\cdots A$	$D-H$	$H\cdots A$	$D\cdots A$	$D-H\cdots A$
O1W—H1W1 \cdots I1	0.86	2.87	3.592 (4)	143
O1W—H2W1 \cdots I1 ⁱ	0.85	2.87	3.567 (4)	141
C14—H14A \cdots O1W	0.96	2.50	3.202 (5)	130
C14—H14D \cdots O1W ⁱⁱ	0.96	2.56	3.460 (5)	157
C1—H1A \cdots I1 ⁱⁱⁱ	0.93	3.20	3.830 (4)	127
C2—H2A \cdots I1 ^{iv}	0.93	3.18	3.825 (4)	129
C3—H3A \cdots I1 ^{iv}	0.93	3.21	3.840 (4)	127

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

Fig. 1

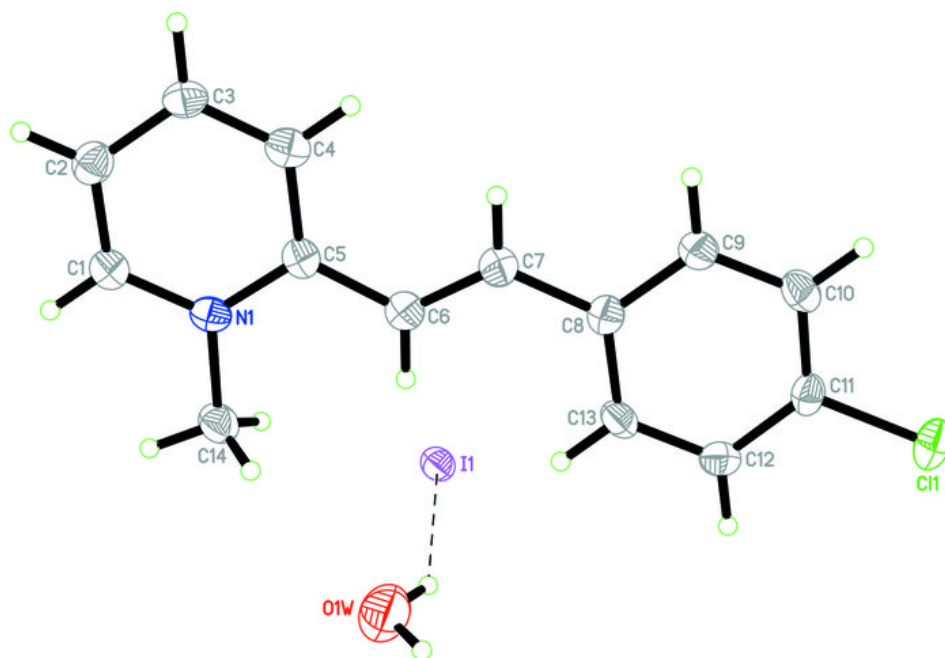


Fig. 2

