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## Structure Reports

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**(1*S*,5*R*)-1-(4-Fluorophenyl)-3-azonia-bicyclo[3.1.0]hexane chloride**

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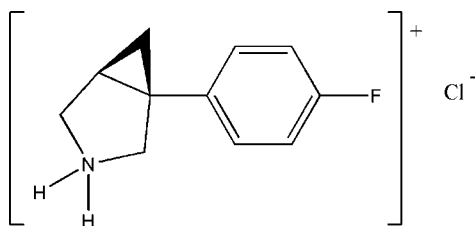
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Key indicators: single-crystal X-ray study;  $T = 296$  K; mean  $\sigma(\text{C}-\text{C}) = 0.002$  Å;  $R$  factor = 0.025;  $wR$  factor = 0.066; data-to-parameter ratio = 17.2.

The absolute structure of the title compound,  $\text{C}_{11}\text{H}_{13}\text{FN}^+\cdot\text{Cl}^-$ , has been determined. The five-membered ring has an envelope conformation with the N atom at the flap position. In the crystal structure, the  $\text{Cl}^-$  anion links with the organic cation *via*  $\text{N}-\text{H}\cdots\text{Cl}$  hydrogen bonding.

## Related literature

For related structures, see: McArdle *et al.* (2004).

## Experimental

## Crystal data

 $\text{C}_{11}\text{H}_{13}\text{FN}^+\cdot\text{Cl}^-$   
 $M_r = 213.67$ 

 Orthorhombic,  $P2_12_12_1$   
 $a = 6.9146$  (10) Å

 $b = 7.8048$  (11) Å  
 $c = 19.448$  (3) Å  
 $V = 1049.6$  (3) Å<sup>3</sup>  
 $Z = 4$ 

 Mo  $K\alpha$  radiation  
 $\mu = 0.34$  mm<sup>-1</sup>  
 $T = 296$  K  
 $0.50 \times 0.36 \times 0.25$  mm

## Data collection

 Bruker APEXII CCD diffractometer  
 Absorption correction: multi-scan (SADABS; Sheldrick, 1996)  
 $T_{\min} = 0.766$ ,  $T_{\max} = 0.919$ 

 6726 measured reflections  
 2292 independent reflections  
 2244 reflections with  $I > 2\sigma(I)$   
 $R_{\text{int}} = 0.028$ 

## Refinement

 $R[F^2 > 2\sigma(F^2)] = 0.025$   
 $wR(F^2) = 0.066$   
 $S = 1.05$   
 2292 reflections  
 133 parameters  
 H atoms treated by a mixture of independent and constrained refinement

 $\Delta\rho_{\text{max}} = 0.38$  e Å<sup>-3</sup>  
 $\Delta\rho_{\text{min}} = -0.17$  e Å<sup>-3</sup>  
 Absolute structure: Flack (1983), 934 Friedel pairs  
 Flack parameter:  $-0.03$  (5)

Table 1

Hydrogen-bond geometry (Å, °).

$D-\text{H}\cdots A$	$D-\text{H}$	$\text{H}\cdots A$	$D\cdots A$	$D-\text{H}\cdots A$
$\text{N1}-\text{H1}\cdots\text{Cl1}$	0.939 (18)	2.146 (18)	3.0837 (13)	176.1 (15)
$\text{N1}-\text{H2}\cdots\text{Cl1}^i$	0.899 (17)	2.275 (17)	3.0907 (13)	150.8 (14)

Symmetry code: (i)  $x - \frac{1}{2}, -y + \frac{3}{2}, -z$ .

Data collection: APEX2 (Bruker, 2007); cell refinement: SAINT-Plus (Bruker, 2007); data reduction: SAINT-Plus; 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.

Supplementary data and figures for this paper are available from the IUCr electronic archives (Reference: XU2479).

## References

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 Flack, H. D. (1983). *Acta Cryst.* **A39**, 876–881.  
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 Sheldrick, G. M. (2008). *Acta Cryst.* **A64**, 112–122.

**supplementary materials**

*Acta Cryst.* (2009). E65, o677 [ doi:10.1107/S1600536809006953 ]

## (1*S*,5*R*)-1-(4-Fluorophenyl)-3-azoniabicyclo[3.1.0]hexane chloride

C. H. Görbitz, T. Hansen and K. Vestli

### Comment

The title compound was prepared as a potential triple neurotransmitter reuptake inhibitor. Details will be published elsewhere. The molecular structure is shown in Fig. 1. The five-membered ring has an envelope conformation with N1 located 0.454 (2) Å above the plane constituted by C1, C2, C4 and C5, on the same side as C3, giving the six-membered ring N1—C1—C2—C3—C4—C5 a distinct boat conformation.

Two different  $P2_1/c$  polymorphs, with  $Z' = 1$  and 4, respectively, were obtained for the racemate of bicifadine hydrochloride (McArdle *et al.*, 2004), which has a methyl group rather than a F atom in the phenyl ortho position. Ring puckering remains unchanged, but phenyl rotations vary; C5—C4—C6—C11 is thus 59.81 (19)° for the title compound, but 83.9° for polymorph 1 of bicifadine and between -10.1 and -30.8° for polymorph 2 (1*S*,5*R*-enantiomers).

### Experimental

Block-shaped crystals were prepared from an acetonitrile solution by slow evaporation at room temperature.

### Refinement

Positional parameters were refined for the two H atoms bonded to N. Other H atoms were positioned with idealized geometry and fixed C—H = 0.93 (aromatic), 0.97 (methylene) or 0.98 Å (methine).  $U_{\text{iso}}(\text{H})$  values were  $1.2U_{\text{eq}}(\text{C,N})$ .

### Figures

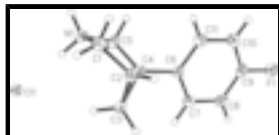


Fig. 1. The asymmetric unit of (I). Displacement ellipsoids are shown at the 50% probability level and H atoms are shown as spheres of arbitrary size.

## (1*S*,5*R*)-1-(4-Fluorophenyl)-3-azoniabicyclo[3.1.0]hexane chloride

### Crystal data

$\text{C}_{11}\text{H}_{13}\text{FN}^+\cdot\text{Cl}^-$

$M_r = 213.67$

Orthorhombic,  $P2_12_12_1$

$a = 6.9146$  (10) Å

$b = 7.8048$  (11) Å

$c = 19.448$  (3) Å

$D_x = 1.352$  Mg m<sup>-3</sup>

Mo  $K\alpha$  radiation

$\lambda = 0.71073$  Å

Cell parameters from 4938 reflections

$\theta = 2.1$ – $27.1^\circ$

$\mu = 0.34$  mm<sup>-1</sup>

$T = 296$  K

# supplementary materials

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$$V = 1049.6 (3) \text{ \AA}^3$$

$$Z = 4$$

$$F_{000} = 448$$

Block, colourless

$$0.50 \times 0.36 \times 0.25 \text{ mm}$$

## Data collection

Bruker APEXII CCD  
diffractometer

2292 independent reflections

Radiation source: fine-focus sealed tube

2244 reflections with  $I > 2\sigma(I)$

Monochromator: graphite

$$R_{\text{int}} = 0.028$$

Detector resolution: 8.3 pixels  $\text{mm}^{-1}$

$$\theta_{\text{max}} = 27.1^\circ$$

$$T = 296 \text{ K}$$

$$\theta_{\text{min}} = 2.1^\circ$$

sets of exposures each taken over  $0.5^\circ$   $\omega$  rotation  
scans

$$h = -8 \rightarrow 4$$

Absorption correction: multi-scan  
(SADABS; Sheldrick, 1996)

$$k = -9 \rightarrow 10$$

$$T_{\text{min}} = 0.766, T_{\text{max}} = 0.919$$

$$l = -24 \rightarrow 24$$

6726 measured reflections

## Refinement

Refinement on  $F^2$

Hydrogen site location: inferred from neighbouring  
sites

Least-squares matrix: full

H atoms treated by a mixture of  
independent and constrained refinement

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

$$w = 1/[\sigma^2(F_o^2) + (0.036P)^2 + 0.1918P]$$

$$wR(F^2) = 0.066$$

$$\text{where } P = (F_o^2 + 2F_c^2)/3$$

$$S = 1.05$$

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

2292 reflections

$$\Delta\rho_{\text{max}} = 0.38 \text{ e \AA}^{-3}$$

133 parameters

$$\Delta\rho_{\text{min}} = -0.17 \text{ e \AA}^{-3}$$

Primary atom site location: structure-invariant direct  
methods

Extinction correction: none

Secondary atom site location: difference Fourier map

Absolute structure: Flack (1983), 934 Friedel pairs

Flack parameter:  $-0.03 (5)$

## Special details

**Experimental.** Crystallized from acetonitrile solution

**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.** Data were collected by measuring three sets of exposures with the detector set at  $2\theta = 29^\circ$ , crystal-to-detector distance 6.00 cm. Refinement of  $F^2$  against ALL reflections.

*Fractional atomic coordinates and isotropic or equivalent isotropic displacement parameters ( $\text{\AA}^2$ )*

x

y

z

$U_{\text{iso}}^*/U_{\text{eq}}$

C11	0.90409 (5)	0.57733 (4)	0.024884 (16)	0.01896 (9)
F1	-0.08580 (14)	0.45054 (11)	0.37784 (4)	0.0280 (2)
C6	0.2562 (2)	0.43164 (19)	0.20420 (7)	0.0184 (3)
C5	0.3941 (2)	0.60246 (17)	0.10328 (6)	0.0185 (3)
H51	0.2695	0.6595	0.1020	0.022*
H52	0.4871	0.6774	0.1254	0.022*
N1	0.45923 (17)	0.55635 (15)	0.03221 (6)	0.0168 (2)
H1	0.595 (3)	0.557 (2)	0.0300 (8)	0.020*
H2	0.422 (3)	0.640 (2)	0.0035 (8)	0.020*
C4	0.3797 (2)	0.43318 (18)	0.14105 (6)	0.0172 (3)
C7	0.3340 (2)	0.3976 (2)	0.26874 (7)	0.0225 (3)
H71	0.4644	0.3701	0.2726	0.027*
C11	0.0603 (2)	0.4731 (2)	0.19938 (7)	0.0235 (3)
H111	0.0067	0.4972	0.1566	0.028*
C9	0.0279 (2)	0.44386 (19)	0.32031 (7)	0.0217 (3)
C1	0.3826 (2)	0.38066 (16)	0.01663 (7)	0.0190 (3)
H11	0.4678	0.3197	-0.0145	0.023*
H12	0.2545	0.3868	-0.0036	0.023*
C8	0.2196 (2)	0.4041 (2)	0.32777 (7)	0.0248 (3)
H81	0.2720	0.3821	0.3709	0.030*
C2	0.3764 (2)	0.29489 (18)	0.08635 (7)	0.0206 (3)
H21	0.2921	0.1953	0.0929	0.025*
C3	0.5512 (2)	0.31501 (19)	0.13212 (7)	0.0224 (3)
H31	0.6677	0.3624	0.1119	0.027*
H32	0.5730	0.2285	0.1671	0.027*
C10	-0.0560 (2)	0.4790 (2)	0.25780 (8)	0.0255 (3)
H101	-0.1867	0.5059	0.2546	0.031*

Atomic displacement parameters ( $\text{\AA}^2$ )

	$U^{11}$	$U^{22}$	$U^{33}$	$U^{12}$	$U^{13}$	$U^{23}$
C11	0.01835 (16)	0.01641 (15)	0.02212 (15)	-0.00080 (13)	-0.00166 (13)	0.00305 (12)
F1	0.0339 (5)	0.0273 (5)	0.0227 (4)	0.0066 (4)	0.0116 (4)	0.0024 (3)
C6	0.0212 (7)	0.0151 (6)	0.0189 (6)	-0.0006 (6)	0.0024 (5)	0.0012 (5)
C5	0.0235 (7)	0.0150 (6)	0.0170 (6)	-0.0025 (6)	0.0013 (6)	0.0003 (5)
N1	0.0187 (6)	0.0147 (5)	0.0169 (5)	0.0003 (5)	-0.0001 (4)	0.0017 (4)
C4	0.0182 (6)	0.0161 (6)	0.0172 (6)	0.0000 (6)	0.0002 (5)	0.0013 (5)
C7	0.0215 (7)	0.0241 (7)	0.0217 (6)	0.0035 (6)	0.0007 (5)	0.0037 (6)
C11	0.0230 (8)	0.0292 (7)	0.0184 (6)	-0.0007 (6)	-0.0016 (5)	0.0019 (5)
C9	0.0292 (7)	0.0160 (7)	0.0198 (6)	0.0007 (6)	0.0086 (6)	0.0005 (5)
C1	0.0209 (7)	0.0162 (6)	0.0198 (6)	-0.0026 (5)	0.0019 (6)	-0.0017 (5)
C8	0.0318 (8)	0.0253 (7)	0.0174 (6)	0.0046 (7)	0.0003 (6)	0.0049 (6)
C2	0.0250 (7)	0.0137 (6)	0.0232 (6)	-0.0006 (6)	0.0044 (6)	0.0004 (5)
C3	0.0227 (7)	0.0230 (7)	0.0216 (7)	0.0055 (6)	0.0024 (5)	0.0066 (5)
C10	0.0204 (7)	0.0302 (8)	0.0259 (7)	0.0024 (6)	0.0025 (6)	0.0016 (6)

Geometric parameters ( $\text{\AA}$ ,  $^\circ$ )

F1—C9	1.3683 (15)	C7—H71	0.9300
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## supplementary materials

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C6—C7	1.3911 (19)	C11—C10	1.393 (2)
C6—C11	1.396 (2)	C11—H111	0.9300
C6—C4	1.4959 (18)	C9—C8	1.369 (2)
C5—N1	1.4977 (16)	C9—C10	1.375 (2)
C5—C4	1.5149 (18)	C1—C2	1.5128 (18)
C5—H51	0.9700	C1—H11	0.9700
C5—H52	0.9700	C1—H12	0.9700
N1—C1	1.5011 (16)	C8—H81	0.9300
N1—H1	0.939 (18)	C2—C3	1.509 (2)
N1—H2	0.899 (17)	C2—H21	0.9800
C4—C3	1.512 (2)	C3—H31	0.9700
C4—C2	1.5156 (19)	C3—H32	0.9700
C7—C8	1.395 (2)	C10—H101	0.9300
C7—C6—C11	118.70 (12)	F1—C9—C8	118.57 (13)
C7—C6—C4	121.43 (13)	F1—C9—C10	118.25 (13)
C11—C6—C4	119.80 (12)	C8—C9—C10	123.18 (13)
N1—C5—C4	104.93 (10)	N1—C1—C2	103.48 (11)
N1—C5—H51	110.8	N1—C1—H11	111.1
C4—C5—H51	110.8	C2—C1—H11	111.1
N1—C5—H52	110.8	N1—C1—H12	111.1
C4—C5—H52	110.8	C2—C1—H12	111.1
H51—C5—H52	108.8	H11—C1—H12	109.0
C5—N1—C1	107.42 (10)	C9—C8—C7	118.03 (13)
C5—N1—H1	109.9 (9)	C9—C8—H81	121.0
C1—N1—H1	110.4 (10)	C7—C8—H81	121.0
C5—N1—H2	108.2 (11)	C3—C2—C1	117.40 (13)
C1—N1—H2	116.1 (11)	C3—C2—C4	59.99 (9)
H1—N1—H2	104.7 (16)	C1—C2—C4	108.28 (11)
C3—C4—C2	59.78 (10)	C3—C2—H21	118.9
C3—C4—C5	115.15 (12)	C1—C2—H21	118.9
C3—C4—C6	122.49 (12)	C4—C2—H21	118.9
C6—C4—C5	116.26 (12)	C2—C3—C4	60.23 (9)
C6—C4—C2	124.19 (12)	C2—C3—H31	117.7
C5—C4—C2	106.36 (10)	C4—C3—H31	117.7
C6—C7—C8	121.09 (14)	C2—C3—H32	117.7
C6—C7—H71	119.5	C4—C3—H32	117.7
C8—C7—H71	119.5	H31—C3—H32	114.9
C10—C11—C6	120.85 (13)	C9—C10—C11	118.15 (14)
C10—C11—H111	119.6	C9—C10—H101	120.9
C6—C11—H111	119.6	C11—C10—H101	120.9
C1—C2—C4—C5	-1.68 (16)	C5—N1—C1—C2	-30.53 (14)
N1—C1—C2—C4	19.46 (15)	F1—C9—C8—C7	179.97 (13)
N1—C5—C4—C2	-16.93 (15)	C10—C9—C8—C7	-0.6 (3)
C5—C4—C6—C11	59.81 (19)	C6—C7—C8—C9	0.4 (2)
C4—C5—N1—C1	29.91 (15)	N1—C1—C2—C3	-45.58 (15)
C7—C6—C4—C3	34.1 (2)	C6—C4—C2—C3	-110.84 (15)
C11—C6—C4—C3	-149.04 (14)	C5—C4—C2—C3	109.97 (13)
C7—C6—C4—C2	107.32 (17)	C6—C4—C2—C1	137.51 (14)

C11—C6—C4—C2	-75.82 (19)	C3—C4—C2—C1	-111.65 (14)
C7—C6—C4—C5	-117.04 (15)	C1—C2—C3—C4	96.23 (13)
N1—C5—C4—C6	-159.86 (12)	C6—C4—C3—C2	113.59 (15)
N1—C5—C4—C3	46.86 (15)	C5—C4—C3—C2	-94.97 (12)
C11—C6—C7—C8	0.2 (2)	F1—C9—C10—C11	179.61 (13)
C4—C6—C7—C8	177.04 (14)	C8—C9—C10—C11	0.2 (2)
C7—C6—C11—C10	-0.6 (2)	C6—C11—C10—C9	0.4 (2)
C4—C6—C11—C10	-177.53 (14)		

*Hydrogen-bond geometry* ( $\text{\AA}$ ,  $^\circ$ )

<i>D</i> —H $\cdots$ <i>A</i>	<i>D</i> —H	H $\cdots$ <i>A</i>	<i>D</i> $\cdots$ <i>A</i>	<i>D</i> —H $\cdots$ <i>A</i>
N1—H1 $\cdots$ C11	0.939 (18)	2.146 (18)	3.0837 (13)	176.1 (15)
N1—H2 $\cdots$ C11 <sup>i</sup>	0.899 (17)	2.275 (17)	3.0907 (13)	150.8 (14)

Symmetry codes: (i)  $x-1/2, -y+3/2, -z$ .

Fig. 1

