

# Potassium *N*-chloro-*o*-toluene-sulfonamidate monohydrate

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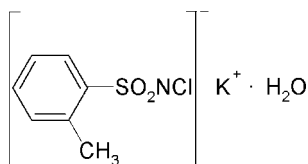
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 Key indicators: single-crystal X-ray study;  $T = 293$  K; mean  $\sigma(\text{C}-\text{C}) = 0.005$  Å;  $R$  factor = 0.044;  $wR$  factor = 0.123; data-to-parameter ratio = 16.0.

In the crystal structure of the title compound,  $\text{K}^+ \cdot \text{C}_7\text{H}_7\text{ClNO}_2\text{S}^- \cdot \text{H}_2\text{O}$ , the  $\text{K}^+$  ion is heptacoordinated by two O atoms from water molecules, four sulfonyl O atoms and the Cl atom of the anion. The S–N distance of 1.584 (3) Å is consistent with an S–N double bond. In the crystal, anions are connected by  $\text{K}^+$  cations into layers parallel to the *ab* plane. The water molecules are coordinated to the  $\text{K}^+$  cations and are additionally linked by intermolecular O–H...Cl and O–H...N hydrogen bonding.

## Related literature

For our studies of the effect of substituents on the structures of *N*-haloarylsulfonamides, see: Gowda *et al.* (2009, 2011*a,b*); and on the oxidative strengths of *N*-halolarylsulfonamides, see: Gowda & Kumar (2003); Usha & Gowda (2006). For similar structures, see: George *et al.* (2000); Olmstead & Power (1986). For the preparation of the title compound, see: Jyothi & Gowda (2004).



## Experimental

### Crystal data

 $\text{K}^+ \cdot \text{C}_7\text{H}_7\text{ClNO}_2\text{S}^- \cdot \text{H}_2\text{O}$   
 $M_r = 261.76$ 

 Orthorhombic, *Pbca*  
 $a = 11.4968$  (9) Å

 $b = 6.7990$  (5) Å  
 $c = 26.883$  (2) Å  
 $V = 2101.4$  (3) Å<sup>3</sup>  
 $Z = 8$ 

 Mo  $K\alpha$  radiation  
 $\mu = 0.94$  mm<sup>-1</sup>  
 $T = 293$  K  
 $0.42 \times 0.40 \times 0.30$  mm

### Data collection

 Oxford Diffraction Xcalibur diffractometer with a Sapphire CCD detector  
 Absorption correction: multi-scan (*CrysAlis RED*; Oxford)

 Diffraction, 2009)  
 $T_{\min} = 0.694$ ,  $T_{\max} = 0.766$   
 4396 measured reflections  
 2150 independent reflections  
 1992 reflections with  $I > 2\sigma(I)$   
 $R_{\text{int}} = 0.017$ 

### Refinement

 $R[F^2 > 2\sigma(F^2)] = 0.044$   
 $wR(F^2) = 0.123$   
 $S = 0.92$   
 2150 reflections  
 134 parameters  
 4 restraints

 H atoms treated by a mixture of independent and constrained refinement  
 $\Delta\rho_{\text{max}} = 0.42$  e Å<sup>-3</sup>  
 $\Delta\rho_{\text{min}} = -0.59$  e Å<sup>-3</sup>
**Table 1**

Hydrogen-bond geometry (Å, °).

<i>D</i> —H... <i>A</i>	<i>D</i> —H	H... <i>A</i>	<i>D</i> ... <i>A</i>	<i>D</i> —H... <i>A</i>
O3—H31...Cl1 <sup>i</sup>	0.85 (1)	2.74 (2)	3.568 (3)	166 (4)
O3—H32...N1 <sup>ii</sup>	0.85 (1)	2.08 (1)	2.909 (4)	167 (3)

 Symmetry codes: (i)  $x + \frac{1}{2}, -y + \frac{5}{2}, -z$ ; (ii)  $-x + 1, -y + 2, -z$ .

Data collection: *CrysAlis CCD* (Oxford Diffraction, 2009); cell refinement: *CrysAlis RED* (Oxford Diffraction, 2009); data reduction: *CrysAlis RED*; program(s) used to solve structure: *SHELXS97* (Sheldrick, 2008); program(s) used to refine structure: *SHELXL97* (Sheldrick, 2008); molecular graphics: *PLATON* (Spek, 2009); 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: NC2234).

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## supporting information

*Acta Cryst.* (2011). E67, m961 [doi:10.1107/S1600536811023555]

## Potassium *N*-chloro-*o*-toluenesulfonamide monohydrate

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### S1. Comment

The crystal structure of the title compound (I) was determined as a part of a project to explore the substituent effects and the effect of replacing sodium ions by potassium ions on the solid state structures of *N*-halo-arylsulfonamides (Gowda & Kumar, 2003; Usha & Gowda, 2006, Gowda *et al.*, 2009, 2011*a,b*). The structure resembles those of potassium *N*, 2-dichloro-benzenesulfonamide sesquihydrate (II)(Gowda *et al.*, 2011*a*), potassium *N*-bromo, *o*-toluenesulfonamide sesquihydrate (Gowda *et al.*, 2011*b*) and sodium *N*-chloro, *o*-toluenesulfonamide sesquihydrate (IV) (Gowda *et al.*, 2009) and other sodium *N*-chloro-aryl- sulfonamides (George *et al.*, 2000; Olmstead & Power, 1986).

In the crystal structure of the title compound the K<sup>+</sup> ion is hepta coordinated by two O atoms from water molecules, four sulfonyl O atoms and one Cl atom of the *N*-chloro,*o*-toluenesulfonamide anions (Fig. 1). This coordination geometry is different from that in II and III, in which the potassium cations are hepta coordinated by three O atoms from water molecules and by four sulfonyl O atoms and and in III, in which the cations are octahedral coordinated.

The S—N distance of 1.584 (3)Å is consistent with an S—N double bond and is in agreement with the observed values of 1.582 (2)Å in II, 1.577 (5)Å in III and 1.590 (2) Å in IV.

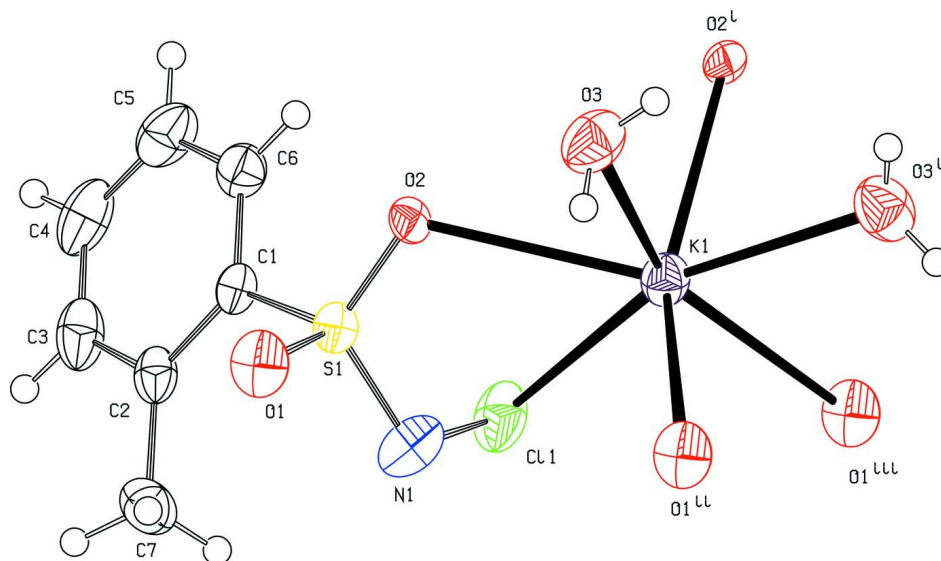
The crystal structure comprises sheets parallel to the *ab* plane (Fig. 2). The molecular packing is additionally stabilized by O—H···Cl and O—H···N hydrogen bonds (Table 1).

### S2. Experimental

The title compound was prepared by the method similar to that reported in literature (Jyothi & Gowda, 2004). *o*-Toluenesulfonamide (2 g) was dissolved in hot aqueous solution (70° C) of 5 M KOH (40 ml). The resulting solution was filtered and Chlorine gas was passed through the clear solution of *o*-toluenesulfonamide in KOH to obtain the title compound. It was filtered under suction, quickly washed with a minimum quantity of ice cold water and dried. The purity of the compound was checked by determining its melting point (155° C). Colourless prisms of the compound were obtained from its aqueous solution at room temperature.

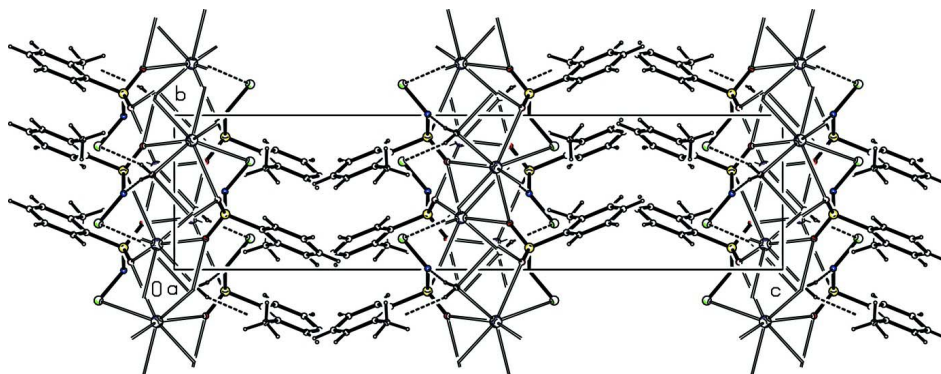
### S3. Refinement

The water H atoms were located in difference map and were refined with O—H distance restrained to 0.85 (2) Å and H—H distance restrained to 1.365 Å. The other H atoms were positioned with idealized geometry using a riding model with the aromatic C—H = 0.93 Å and methyl C—H = 0.96 Å. All H atoms were refined with isotropic displacement parameters (set to 1.2 times of the  $U_{eq}$  of the parent atom).



**Figure 1**

Molecular structure of the title compound showing the coordination geometry of the  $K^+$  cations with labelling and displacement ellipsoids drawn at the 50% probability level. Symmetry codes: (i)  $-x + 3/2, y + 1/2, z$ ; (ii)  $-x + 1, -y + 2, -z$ ; (iii)  $x, y + 1, z$ .



**Figure 2**

Crystal structure of the title compound with view in the direction of the  $a$  axis and hydrogen bonding drawn as dashed lines.

### Potassium *N*-chloro-*o*-toluenesulfonamidate monohydrate

#### Crystal data

$K^+ \cdot C_7H_7ClNO_2S^- \cdot H_2O$

$M_r = 261.76$

Orthorhombic,  $Pbca$

Hall symbol:  $-P 2ac 2ab$

$a = 11.4968 (9) \text{ \AA}$

$b = 6.7990 (5) \text{ \AA}$

$c = 26.883 (2) \text{ \AA}$

$V = 2101.4 (3) \text{ \AA}^3$

$Z = 8$

$F(000) = 1072$

$D_x = 1.655 \text{ Mg m}^{-3}$

Mo  $K\alpha$  radiation,  $\lambda = 0.71073 \text{ \AA}$

Cell parameters from 2737 reflections

$\theta = 3.0\text{--}27.7^\circ$

$\mu = 0.94 \text{ mm}^{-1}$

$T = 293 \text{ K}$

Prism, colourless

$0.42 \times 0.40 \times 0.30 \text{ mm}$

*Data collection*

Oxford Diffraction Xcalibur  
diffractometer with a Sapphire CCD detector  
Radiation source: fine-focus sealed tube  
Graphite monochromator  
Rotation method data acquisition using  $\omega$  scans  
Absorption correction: multi-scan  
(*CrysAlis RED*; Oxford Diffraction, 2009)  
 $T_{\min} = 0.694$ ,  $T_{\max} = 0.766$

4396 measured reflections  
2150 independent reflections  
1992 reflections with  $I > 2\sigma(I)$   
 $R_{\text{int}} = 0.017$   
 $\theta_{\max} = 26.4^\circ$ ,  $\theta_{\min} = 3.5^\circ$   
 $h = -2 \rightarrow 14$   
 $k = -8 \rightarrow 5$   
 $l = -33 \rightarrow 14$

*Refinement*

Refinement on  $F^2$   
Least-squares matrix: full  
 $R[F^2 > 2\sigma(F^2)] = 0.044$   
 $wR(F^2) = 0.123$   
 $S = 0.92$   
2150 reflections  
134 parameters  
4 restraints  
Primary atom site location: structure-invariant  
direct methods

Secondary atom site location: difference Fourier  
map  
Hydrogen site location: inferred from  
neighbouring sites  
H atoms treated by a mixture of independent  
and constrained refinement  
 $w = 1/[\sigma^2(F_o^2) + (0.0717P)^2 + 5.301P]$   
where  $P = (F_o^2 + 2F_c^2)/3$   
 $(\Delta/\sigma)_{\max} = 0.001$   
 $\Delta\rho_{\max} = 0.42 \text{ e } \text{\AA}^{-3}$   
 $\Delta\rho_{\min} = -0.59 \text{ e } \text{\AA}^{-3}$

*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>	$U_{\text{iso}}^*/U_{\text{eq}}$
K1	0.63387 (6)	1.33711 (10)	0.02856 (2)	0.0327 (2)
Cl1	0.47408 (9)	1.20077 (12)	0.12455 (3)	0.0481 (3)
S1	0.55243 (6)	0.85908 (10)	0.08371 (2)	0.0260 (2)
O1	0.5175 (2)	0.6996 (3)	0.05138 (8)	0.0412 (5)
N1	0.4458 (2)	1.0067 (4)	0.08324 (10)	0.0368 (6)
O3	0.7703 (2)	1.1144 (4)	-0.03605 (9)	0.0494 (6)
H31	0.817 (3)	1.179 (6)	-0.0541 (12)	0.059*
H32	0.713 (2)	1.084 (7)	-0.0541 (12)	0.059*
O2	0.66241 (18)	0.9505 (3)	0.07163 (8)	0.0367 (5)
C1	0.5708 (2)	0.7601 (4)	0.14466 (10)	0.0251 (5)
C2	0.4782 (3)	0.6704 (4)	0.17010 (11)	0.0305 (6)
C3	0.5030 (3)	0.5932 (5)	0.21709 (12)	0.0426 (8)
H3	0.4441	0.5301	0.2346	0.051*
C4	0.6116 (4)	0.6073 (5)	0.23828 (12)	0.0492 (9)
H4	0.6249	0.5548	0.2697	0.059*

C5	0.7001 (3)	0.6985 (5)	0.21318 (13)	0.0466 (8)
H5	0.7732	0.7096	0.2277	0.056*
C6	0.6803 (3)	0.7742 (5)	0.16600 (11)	0.0348 (6)
H6	0.7406	0.8345	0.1487	0.042*
C7	0.3566 (3)	0.6577 (5)	0.15053 (14)	0.0427 (8)
H7A	0.3582	0.6144	0.1165	0.051*
H7B	0.3205	0.7849	0.1524	0.051*
H7C	0.3129	0.5657	0.1701	0.051*

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

	$U^{11}$	$U^{22}$	$U^{33}$	$U^{12}$	$U^{13}$	$U^{23}$
K1	0.0370 (4)	0.0316 (4)	0.0294 (3)	−0.0002 (3)	−0.0028 (2)	0.0026 (3)
Cl1	0.0630 (6)	0.0315 (4)	0.0497 (5)	0.0118 (4)	0.0137 (4)	−0.0022 (3)
S1	0.0319 (4)	0.0261 (4)	0.0199 (3)	0.0016 (3)	0.0016 (2)	0.0001 (2)
O1	0.0564 (14)	0.0378 (12)	0.0293 (11)	−0.0012 (11)	−0.0026 (10)	−0.0105 (10)
N1	0.0396 (14)	0.0365 (14)	0.0344 (13)	0.0095 (11)	−0.0050 (10)	0.0007 (11)
O3	0.0535 (15)	0.0560 (16)	0.0387 (12)	0.0129 (13)	0.0017 (11)	0.0097 (12)
O2	0.0377 (10)	0.0383 (12)	0.0340 (11)	−0.0015 (8)	0.0096 (9)	0.0056 (9)
C1	0.0325 (13)	0.0201 (12)	0.0228 (12)	0.0029 (11)	−0.0005 (10)	−0.0009 (10)
C2	0.0406 (15)	0.0216 (13)	0.0293 (14)	0.0025 (11)	0.0092 (12)	−0.0011 (11)
C3	0.062 (2)	0.0311 (16)	0.0344 (16)	0.0036 (15)	0.0161 (15)	0.0062 (13)
C4	0.079 (3)	0.0414 (18)	0.0274 (15)	0.0164 (18)	−0.0037 (16)	0.0082 (14)
C5	0.057 (2)	0.0455 (19)	0.0377 (17)	0.0096 (16)	−0.0175 (15)	0.0012 (15)
C6	0.0377 (15)	0.0324 (15)	0.0343 (15)	0.0002 (13)	−0.0037 (12)	0.0013 (12)
C7	0.0375 (16)	0.0408 (18)	0.0499 (19)	−0.0067 (14)	0.0107 (14)	−0.0013 (15)

*Geometric parameters (Å, °)*

K1—O2 <sup>i</sup>	2.724 (2)	O3—H31	0.848 (10)
K1—O1 <sup>ii</sup>	2.777 (2)	O3—H32	0.847 (10)
K1—O3	2.788 (3)	O2—K1 <sup>v</sup>	2.724 (2)
K1—O3 <sup>i</sup>	2.790 (3)	C1—C6	1.386 (4)
K1—O1 <sup>iii</sup>	2.870 (2)	C1—C2	1.405 (4)
K1—O2	2.891 (2)	C2—C3	1.397 (4)
K1—Cl1	3.3006 (11)	C2—C7	1.497 (5)
K1—N1	3.447 (3)	C3—C4	1.375 (6)
K1—H31	3.25 (4)	C3—H3	0.9300
K1—H32	2.95 (4)	C4—C5	1.369 (6)
Cl1—N1	1.755 (3)	C4—H4	0.9300
S1—O1	1.446 (2)	C5—C6	1.388 (4)
S1—O2	1.446 (2)	C5—H5	0.9300
S1—N1	1.584 (3)	C6—H6	0.9300
S1—C1	1.784 (3)	C7—H7A	0.9600
O1—K1 <sup>ii</sup>	2.777 (2)	C7—H7B	0.9600
O1—K1 <sup>iv</sup>	2.870 (2)	C7—H7C	0.9600
O3—K1 <sup>v</sup>	2.790 (3)		

O2 <sup>i</sup> —K1—O1 <sup>ii</sup>	153.30 (7)	O1—S1—O2	115.46 (14)
O2 <sup>i</sup> —K1—O3	86.25 (8)	O1—S1—N1	104.81 (15)
O1 <sup>ii</sup> —K1—O3	79.74 (8)	O2—S1—N1	113.75 (14)
O2 <sup>i</sup> —K1—O3 <sup>i</sup>	74.57 (7)	O1—S1—C1	107.58 (13)
O1 <sup>ii</sup> —K1—O3 <sup>i</sup>	80.01 (7)	O2—S1—C1	105.34 (13)
O3—K1—O3 <sup>i</sup>	75.94 (5)	N1—S1—C1	109.75 (13)
O2 <sup>i</sup> —K1—O1 <sup>iii</sup>	93.83 (7)	S1—O1—K1 <sup>ii</sup>	134.92 (14)
O1 <sup>ii</sup> —K1—O1 <sup>iii</sup>	87.16 (6)	S1—O1—K1 <sup>iv</sup>	130.02 (14)
O3—K1—O1 <sup>iii</sup>	149.53 (7)	K1 <sup>ii</sup> —O1—K1 <sup>iv</sup>	92.84 (6)
O3 <sup>i</sup> —K1—O1 <sup>iii</sup>	74.75 (8)	S1—N1—C11	109.14 (15)
O2 <sup>i</sup> —K1—O2	89.39 (6)	S1—N1—K1	86.03 (11)
O1 <sup>ii</sup> —K1—O2	107.42 (7)	C11—N1—K1	70.34 (10)
O3—K1—O2	72.04 (7)	K1—O3—K1 <sup>v</sup>	101.62 (8)
O3 <sup>i</sup> —K1—O2	144.97 (8)	K1—O3—H31	116 (3)
O1 <sup>iii</sup> —K1—O2	138.41 (7)	K1 <sup>v</sup> —O3—H31	117 (3)
O2 <sup>i</sup> —K1—C11	103.04 (5)	K1—O3—H32	93 (3)
O1 <sup>ii</sup> —K1—C11	103.35 (6)	K1 <sup>v</sup> —O3—H32	120 (3)
O3—K1—C11	130.37 (6)	H31—O3—H32	107.3 (16)
O3 <sup>i</sup> —K1—C11	153.67 (7)	S1—O2—K1 <sup>v</sup>	136.54 (13)
O1 <sup>iii</sup> —K1—C11	79.32 (5)	S1—O2—K1	112.42 (12)
O2—K1—C11	59.65 (4)	K1 <sup>v</sup> —O2—K1	100.64 (7)
O2 <sup>i</sup> —K1—N1	122.85 (7)	C6—C1—C2	121.1 (3)
O1 <sup>ii</sup> —K1—N1	83.01 (7)	C6—C1—S1	117.5 (2)
O3—K1—N1	105.34 (8)	C2—C1—S1	121.4 (2)
O3 <sup>i</sup> —K1—N1	162.46 (7)	C3—C2—C1	116.6 (3)
O1 <sup>iii</sup> —K1—N1	100.15 (7)	C3—C2—C7	119.1 (3)
O2—K1—N1	46.20 (6)	C1—C2—C7	124.2 (3)
C11—K1—N1	30.05 (5)	C4—C3—C2	122.2 (3)
O2 <sup>i</sup> —K1—H31	80.0 (6)	C4—C3—H3	118.9
O1 <sup>ii</sup> —K1—H31	81.2 (6)	C2—C3—H3	118.9
O3—K1—H31	13.6 (5)	C3—C4—C5	120.1 (3)
O3 <sup>i</sup> —K1—H31	62.7 (6)	C3—C4—H4	119.9
O1 <sup>iii</sup> —K1—H31	137.2 (6)	C5—C4—H4	119.9
O2—K1—H31	84.2 (6)	C4—C5—C6	119.8 (3)
C11—K1—H31	143.4 (6)	C4—C5—H5	120.1
N1—K1—H31	118.9 (5)	C6—C5—H5	120.1
O2 <sup>i</sup> —K1—H32	102.8 (3)	C5—C6—C1	120.1 (3)
O1 <sup>ii</sup> —K1—H32	63.8 (5)	C5—C6—H6	120.0
O3—K1—H32	16.6 (3)	C1—C6—H6	120.0
O3 <sup>i</sup> —K1—H32	78.7 (8)	C2—C7—H7A	109.5
O1 <sup>iii</sup> —K1—H32	143.6 (7)	C2—C7—H7B	109.5
O2—K1—H32	74.7 (8)	H7A—C7—H7B	109.5
C11—K1—H32	126.5 (7)	C2—C7—H7C	109.5
N1—K1—H32	97.6 (6)	H7A—C7—H7C	109.5
H31—K1—H32	24.8 (3)	H7B—C7—H7C	109.5
N1—C11—K1	79.61 (10)		
O2 <sup>i</sup> —K1—C11—N1	-135.35 (10)	N1—K1—O3—K1 <sup>v</sup>	-57.22 (10)

O1 <sup>ii</sup> —K1—C11—N1	48.70 (10)	O1—S1—O2—K1 <sup>v</sup>	-29.4 (2)
O3—K1—C11—N1	-39.09 (12)	N1—S1—O2—K1 <sup>v</sup>	-150.65 (16)
O3 <sup>i</sup> —K1—C11—N1	143.14 (15)	C1—S1—O2—K1 <sup>v</sup>	89.11 (19)
O1 <sup>iii</sup> —K1—C11—N1	133.16 (10)	O1—S1—O2—K1	107.12 (14)
O2—K1—C11—N1	-53.88 (10)	N1—S1—O2—K1	-14.13 (17)
O2—S1—O1—K1 <sup>ii</sup>	-99.9 (2)	C1—S1—O2—K1	-134.36 (11)
N1—S1—O1—K1 <sup>ii</sup>	26.1 (2)	O2 <sup>i</sup> —K1—O2—S1	147.79 (8)
C1—S1—O1—K1 <sup>ii</sup>	142.86 (17)	O1 <sup>ii</sup> —K1—O2—S1	-53.29 (14)
O2—S1—O1—K1 <sup>iv</sup>	58.3 (2)	O3—K1—O2—S1	-125.95 (14)
N1—S1—O1—K1 <sup>iv</sup>	-175.69 (16)	O3 <sup>i</sup> —K1—O2—S1	-150.81 (12)
C1—S1—O1—K1 <sup>iv</sup>	-58.93 (19)	O1 <sup>iii</sup> —K1—O2—S1	52.71 (17)
O1—S1—N1—C11	176.31 (15)	C11—K1—O2—S1	42.26 (10)
O2—S1—N1—C11	-56.67 (19)	N1—K1—O2—S1	8.18 (10)
C1—S1—N1—C11	61.05 (19)	O2 <sup>i</sup> —K1—O2—K1 <sup>v</sup>	-60.99 (11)
O1—S1—N1—K1	-116.09 (11)	O1 <sup>ii</sup> —K1—O2—K1 <sup>v</sup>	97.93 (8)
O2—S1—N1—K1	10.93 (13)	O3—K1—O2—K1 <sup>v</sup>	25.27 (8)
C1—S1—N1—K1	128.65 (10)	O3 <sup>i</sup> —K1—O2—K1 <sup>v</sup>	0.41 (16)
K1—C11—N1—S1	78.36 (14)	O1 <sup>iii</sup> —K1—O2—K1 <sup>v</sup>	-156.08 (8)
O2 <sup>i</sup> —K1—N1—S1	-57.37 (13)	C11—K1—O2—K1 <sup>v</sup>	-166.52 (9)
O1 <sup>ii</sup> —K1—N1—S1	115.47 (11)	N1—K1—O2—K1 <sup>v</sup>	159.39 (12)
O3—K1—N1—S1	38.17 (12)	O1—S1—C1—C6	119.2 (2)
O3 <sup>i</sup> —K1—N1—S1	130.0 (2)	O2—S1—C1—C6	-4.5 (3)
O1 <sup>iii</sup> —K1—N1—S1	-158.69 (10)	N1—S1—C1—C6	-127.3 (2)
O2—K1—N1—S1	-6.91 (9)	O1—S1—C1—C2	-60.5 (3)
C11—K1—N1—S1	-111.95 (14)	O2—S1—C1—C2	175.8 (2)
O2 <sup>i</sup> —K1—N1—C11	54.58 (11)	N1—S1—C1—C2	53.0 (3)
O1 <sup>ii</sup> —K1—N1—C11	-132.58 (10)	C6—C1—C2—C3	-1.4 (4)
O3—K1—N1—C11	150.12 (9)	S1—C1—C2—C3	178.3 (2)
O3 <sup>i</sup> —K1—N1—C11	-118.0 (3)	C6—C1—C2—C7	177.1 (3)
O1 <sup>iii</sup> —K1—N1—C11	-46.74 (10)	S1—C1—C2—C7	-3.2 (4)
O2—K1—N1—C11	105.04 (11)	C1—C2—C3—C4	1.5 (5)
O2 <sup>i</sup> —K1—O3—K1 <sup>v</sup>	65.85 (9)	C7—C2—C3—C4	-177.1 (3)
O1 <sup>ii</sup> —K1—O3—K1 <sup>v</sup>	-136.97 (10)	C2—C3—C4—C5	-0.3 (5)
O3 <sup>i</sup> —K1—O3—K1 <sup>v</sup>	140.87 (13)	C3—C4—C5—C6	-0.9 (5)
O1 <sup>iii</sup> —K1—O3—K1 <sup>v</sup>	157.04 (11)	C4—C5—C6—C1	1.0 (5)
O2—K1—O3—K1 <sup>v</sup>	-24.72 (8)	C2—C1—C6—C5	0.2 (5)
C11—K1—O3—K1 <sup>v</sup>	-38.10 (13)	S1—C1—C6—C5	-179.5 (3)

Symmetry codes: (i)  $-x+3/2, y+1/2, z$ ; (ii)  $-x+1, -y+2, -z$ ; (iii)  $x, y+1, z$ ; (iv)  $x, y-1, z$ ; (v)  $-x+3/2, y-1/2, z$ .

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

$D-H\cdots A$	$D-H$	$H\cdots A$	$D\cdots A$	$D-H\cdots A$
O3—H31 $\cdots$ C11 <sup>vi</sup>	0.85 (1)	2.74 (2)	3.568 (3)	166 (4)
O3—H32 $\cdots$ N1 <sup>ii</sup>	0.85 (1)	2.08 (1)	2.909 (4)	167 (3)

Symmetry codes: (ii)  $-x+1, -y+2, -z$ ; (vi)  $x+1/2, -y+5/2, -z$ .