



ISSN 1600-5368

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Crystal structure of 4,5-bis(3,4,5-trimethoxyphenyl)-2*H*-1,2,3-triazole methanol monosolvate

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Received 17 September 2014; accepted 18 September 2014

Edited by J. Simpson, University of Otago, New Zealand

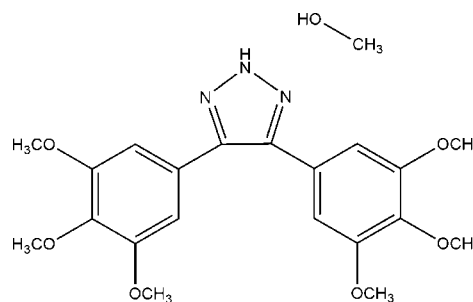
The title compound, $C_{20}H_{23}N_3O_6 \cdot CH_3OH$ was synthesized by [3 + 2] cycloaddition of (*Z*)-2,3-bis(3,4,5-trimethoxyphenyl)acrylonitrile with sodium azide and ammonium chloride in DMF/water. The central nitrogen of the triazole ring is protonated. The dihedral angles between the triazole ring and the 3,4,5-trimethoxyphenyl ring planes are 34.31 (4) and 45.03 (5)°, while that between the 3,4,5-trimethoxyphenyl rings is 51.87 (5)°. In the crystal, the molecules, along with two methanol solvent molecules are linked into an $R_4^4(10)$ centrosymmetric dimer by N—H···O and O—H···N hydrogen bonds.

Keywords: crystal structure; hydrogen bonds; 1,2,3-triazole.

CCDC reference: 1025101

1. Related literature

The synthetic procedure has been described by Madadi *et al.* (2014) and by Penthala *et al.* (2014). For structure-related activity, see: Young & Chaplin (2004); Pettit *et al.* (1995); Hsieh *et al.* (2005); Carr *et al.* (2010); Banimustafa *et al.* (2013); Demchuk *et al.* (2014).



2. Experimental

2.1. Crystal data

$C_{20}H_{23}N_3O_6 \cdot CH_4O$
 $M_r = 433.45$
 Triclinic, $P\bar{1}$
 $a = 10.1458$ (1) Å
 $b = 10.6090$ (1) Å
 $c = 11.0435$ (2) Å
 $\alpha = 89.5708$ (6)°
 $\beta = 72.5903$ (6)°
 $\gamma = 70.7146$ (7)°
 $V = 1065.07$ (2) Å³
 $Z = 2$
 Mo $K\alpha$ radiation
 $\mu = 0.10$ mm⁻¹
 $T = 90$ K
 $0.24 \times 0.22 \times 0.20$ mm

2.2. Data collection

Nonius KappaCCD diffractometer
 Absorption correction: multi-scan (*SADABS*; Sheldrick, 2008*a*)
 $T_{\min} = 0.882$, $T_{\max} = 0.970$
 28813 measured reflections
 4887 independent reflections
 3960 reflections with $I > 2\sigma(I)$
 $R_{\text{int}} = 0.037$

2.3. Refinement

$R[F^2 > 2\sigma(F^2)] = 0.045$
 $wR(F^2) = 0.127$
 $S = 1.05$
 4887 reflections
 291 parameters
 H atoms treated by a mixture of independent and constrained refinement
 $\Delta\rho_{\text{max}} = 0.28$ e Å⁻³
 $\Delta\rho_{\text{min}} = -0.30$ e Å⁻³

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>
N2—H2N···O1M	0.936 (17)	1.797 (18)	2.7303 (15)	174.2 (16)
O1M—H1M···N3 ⁱ	0.84	1.97	2.8101 (15)	177

Symmetry code: (i) $-x + 2, -y + 1, -z + 1$.

Data collection: *COLLECT* (Nonius, 1998); cell refinement: *SCALEPACK* (Otwinowski & Minor, 2006); data reduction: *DENZO-SMN* (Otwinowski & Minor, 2006); program(s) used to solve structure: *SHELXS97* (Sheldrick, 2008*b*); program(s) used to refine structure: *SHELXL2013* (Sheldrick, 2008*b*); molecular graphics: *XP in SHELXTL* (Sheldrick, 2008*b*); software used to prepare material for publication: *SHELX* (Sheldrick, 2008*b*).

Acknowledgements

This investigation was supported NIH/National Cancer Institute (CA140409) United States, and we also thank the Arkansas Research Alliance for financial support.

Supporting information for this paper is available from the IUCr electronic archives (Reference: SJ5427).

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supporting information

Acta Cryst. (2014). E70, o1128–o1129 [doi:10.1107/S1600536814020911]

Crystal structure of 4,5-bis(3,4,5-trimethoxyphenyl)-2*H*-1,2,3-triazole methanol monosolvate

Nikhil Reddy Madadi, Narsimha Reddy Penthala, Shobanbabu Bommagani, Sean Parkin and Peter A. Crooks

S1. Comment

Antimitotic agents are a major class of anticancer drugs that target the microtubules. There are three major binding sites on tubulin, namely the colchicine, taxane and vinca domains. Lately, anti-mitotic agents such as combretastatin A-4 that bind to the colchicine binding site are receiving much attention due to their potent anticancer and antiangiogenic properties. Combretastatin A-4 is a *cis* configured natural product extracted from the South African willow tree *combretum caffrum* (Pettit *et al.*, 1995). Its phosphate prodrug (CA-4P) is currently in phase 3 clinical trials for anaplastic thyroid cancer, and it has successfully arrested tumor growth in a wide spectrum of tumor models (Young and Chaplin, 2004). However, recent studies have reported the chemical instability of CA-4 due to *cis-trans* isomerization to the more thermodynamically stable, but less potent, *trans*-CA-4 isomer (Hsieh *et al.*, 2005). Recently, much research has been conducted to stabilize the *cis* configuration by replacing the ethylene bridge with heterocyclic ring systems, including thiazoles, tetrazoles, imidazoles, pyrazoles, oxazolones, triazoles, and furanones (Carr *et al.*, 2010; Banimustafa *et al.*, 2013; Demchuk *et al.*, 2014). One approach we are exploring is the replacement of the ethylene bridge in the *cis* stilbene scaffold with a triazole moiety to produce geometrically stable triazole analogs of CA4 with improved water solubility.

The title compound structure determination was performed to determine unequivocally the position of the hydrogen atom on the triazole ring system, which cannot be easily determined by NMR spectroscopy, and to obtain detailed information on the structural conformation of the molecule that may be useful in structure-activity relationship (SAR) analysis. The title compound was synthesized in two steps as described by Madadi *et al.*, 2014 and Penthala *et al.*, 2014).

In the first step, (*Z*)-2,3-bis(3,4,5-trimethoxyphenyl)acrylonitrile was synthesized by reacting 2-(3,4,5-trimethoxyphenyl)acetonitrile with 3,4,5-trimethoxybenzaldehyde in 5% NaOMe in methanol to afford the product in 85% yield. In the second step 4,5-bis(3,4,5-trimethoxyphenyl)-2*H*-1,2,3-triazole was synthesized by refluxing a mixture of (*Z*)-2,3-bis(3,4,5-trimethoxyphenyl)acrylonitrile, sodium azide and ammonium chloride in a DMF/water mixture to afford the desired product in 64% yield. The crystal structure of the compound indicates the presence of a 2*H* nitrogen on the triazole ring, *i.e.* protonation of the middle N atom. The molecules, along with two methanol solvent molecules, are linked into an $R^4_4(10)$ centrosymmetric dimer by N—H \cdots O and O—H \cdots N intermolecular hydrogen bonds. The dihedral angles between the triazole ring and the two 3,4,5-trimethoxyphenyl ring planes are 34.31 (4) $^\circ$ and 45.03 (5) $^\circ$, while that between the two 3,4,5-trimethoxyphenyl rings is 51.87 (5) $^\circ$.

S2. Experimental

A mixture of (*Z*)-2,3-bis(3,4,5-trimethoxyphenyl)acrylonitrile, sodium azide and ammonium chloride in a mole ratio of 1:3:3, respectively, was refluxed in 10% aqueous DMF for 5 hrs. The reaction was monitored by TLC. When the (*Z*)-2,3-

bis(3,4,5-trimethoxyphenyl)-acrylonitrile strating material had completely disappeared, cold water was added and the mixture was stirred over 10–15 min, during which the final product precipitated out. The product was purified by flash column chromatography utilizing ethyl acetate/methanol to afford 4,5-bis(3,4,5-trimethoxyphenyl)-2*H*-1,2,3-triazole in 64% yield. Crystallization from methanol afforded a white crystalline product: 4,5-bis(3,4,5-trimethoxyphenyl)-2*H*-1,2,3-triazole methanolate, which was suitable for X-ray crystallographic analysis.

^1H NMR (400 MHz, CDCl_3 -d): δ 3.93 (s, 3H, $-\text{OCH}_3$), 6.87 (d, $J = 8$ Hz, 1H, ArH), 7.05 (d, $J = 12$ Hz, 2H, ArH), 7.42 (s, 2H, ArH), 7.78 (s, 1H, ArH) *p.p.m.* ^{13}C NMR (400 MHz, CDCl_3 -d): δ 55.87, 55.96, 111.05, 127.31, 129.80, 129.89, 130.54, 132.60, 132.86, 149.15, 149.54 *p.p.m.* HRMS (ESI): m/z calcd for $\text{C}_{20}\text{H}_{23}\text{N}_3\text{O}_6$ [$\text{M}-\text{H}$] 350.0463; found 350.0480.

S3. Refinement

H atoms were found in difference Fourier maps. Those bonded to carbon and oxygen were subsequently placed at idealized positions with constrained distances of 0.98 Å (RCH_3), 0.95 Å ($\text{C}_{\text{sp}^2}\text{H}$) and 0.84 Å (OH), while the nitrogen-bound H atom position was refined. $U_{\text{iso}}(\text{H})$ values were set to either $1.2U_{\text{eq}}$ or $1.5U_{\text{eq}}$ (RCH_3 , OH) of the attached atom.

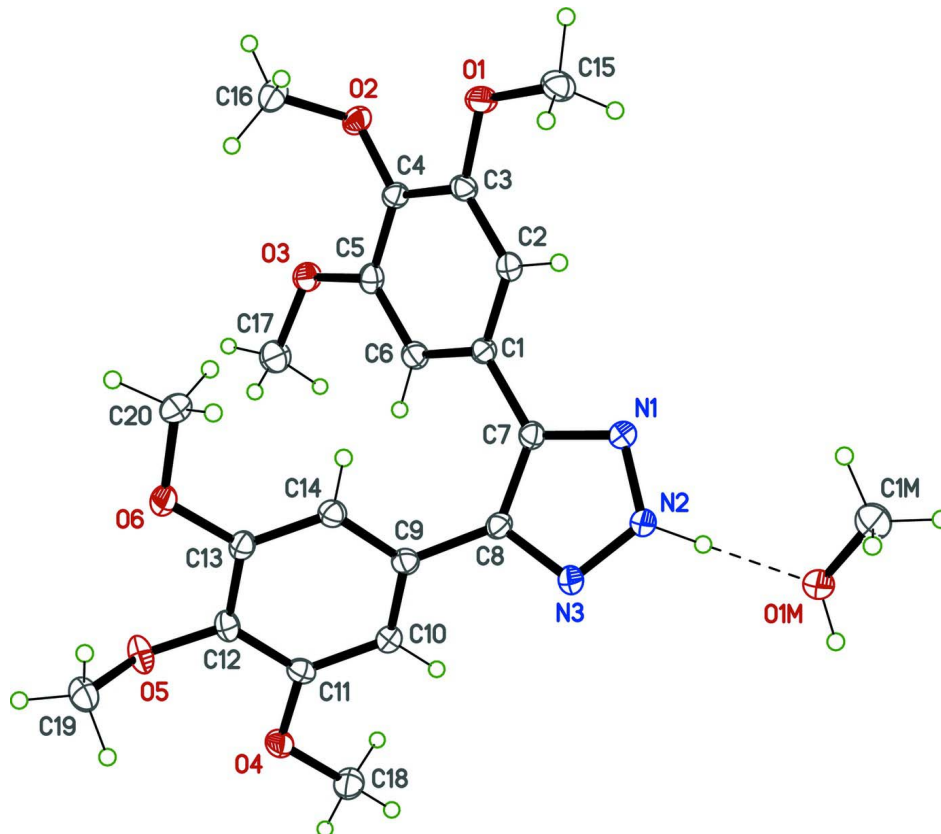


Figure 1

A view of the asymmetric unit of the structure with the atom numbering scheme. Displacement ellipsoids are drawn at the 50% probability level and a hydrogen bond is drawn as a dashed line.

4,5-Bis(3,4,5-trimethoxyphenyl)-2H-1,2,3-triazole methanol monosolvate

Crystal data

$C_{20}H_{23}N_3O_6 \cdot CH_4O$
 $M_r = 433.45$
 Triclinic, $P\bar{1}$
 $a = 10.1458$ (1) Å
 $b = 10.6090$ (1) Å
 $c = 11.0435$ (2) Å
 $\alpha = 89.5708$ (6)°
 $\beta = 72.5903$ (6)°
 $\gamma = 70.7146$ (7)°
 $V = 1065.07$ (2) Å³

$Z = 2$
 $F(000) = 460$
 $D_x = 1.352$ Mg m⁻³
 Mo $K\alpha$ radiation, $\lambda = 0.71073$ Å
 Cell parameters from 11695 reflections
 $\theta = 1.0$ – 27.5 °
 $\mu = 0.10$ mm⁻¹
 $T = 90$ K
 Block, colourless
 $0.24 \times 0.22 \times 0.20$ mm

Data collection

Nonius KappaCCD
 diffractometer
 Radiation source: fine-focus sealed-tube
 Detector resolution: 9.1 pixels mm⁻¹
 φ and ω scans at fixed $\chi = 55$ °
 Absorption correction: multi-scan
 (SADABS; Sheldrick, 2008a)
 $T_{\min} = 0.882$, $T_{\max} = 0.970$

28813 measured reflections
 4887 independent reflections
 3960 reflections with $I > 2\sigma(I)$
 $R_{\text{int}} = 0.037$
 $\theta_{\max} = 27.5$ °, $\theta_{\min} = 1.9$ °
 $h = -13 \rightarrow 13$
 $k = -13 \rightarrow 13$
 $l = -14 \rightarrow 14$

Refinement

Refinement on F^2
 Least-squares matrix: full
 $R[F^2 > 2\sigma(F^2)] = 0.045$
 $wR(F^2) = 0.127$
 $S = 1.05$
 4887 reflections
 291 parameters
 0 restraints
 Primary atom site location: structure-invariant
 direct methods

Secondary atom site location: difference Fourier
 map
 Hydrogen site location: difference Fourier map
 H atoms treated by a mixture of independent
 and constrained refinement
 $w = 1/[\sigma^2(F_o^2) + (0.0731P)^2 + 0.3615P]$
 where $P = (F_o^2 + 2F_c^2)/3$
 $(\Delta/\sigma)_{\max} < 0.001$
 $\Delta\rho_{\max} = 0.28$ e Å⁻³
 $\Delta\rho_{\min} = -0.30$ e Å⁻³

Special details

Experimental. The crystal was mounted with polyisobutene oil on the tip of a fine glass fibre, fastened in a copper mounting pin with electrical solder. It was placed directly into the cold stream of a liquid nitrogen based cryostat. Diffraction data were collected with the crystal at 90 K, which is standard practice in this laboratory for the majority of flash-cooled crystals.

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.

Fractional atomic coordinates and isotropic or equivalent isotropic displacement parameters (Å²)

	<i>x</i>	<i>y</i>	<i>z</i>	$U_{\text{iso}}^*/U_{\text{eq}}$
N1	0.60782 (12)	0.74167 (11)	0.56615 (11)	0.0185 (2)
N2	0.73493 (13)	0.64285 (12)	0.54691 (11)	0.0195 (3)
H2N	0.8264 (19)	0.6486 (17)	0.5010 (16)	0.023*
N3	0.72437 (12)	0.53382 (11)	0.60414 (11)	0.0182 (2)

O1	0.10274 (10)	1.12766 (9)	0.78293 (10)	0.0199 (2)
O2	-0.09566 (10)	1.00928 (10)	0.79884 (9)	0.0203 (2)
O3	-0.01750 (10)	0.74272 (10)	0.76182 (10)	0.0212 (2)
O4	0.55195 (11)	0.11586 (9)	0.76505 (9)	0.0200 (2)
O5	0.35131 (11)	0.22077 (10)	0.98936 (9)	0.0226 (2)
O6	0.26188 (10)	0.48551 (10)	1.06686 (9)	0.0203 (2)
C1	0.34889 (14)	0.77606 (13)	0.68391 (12)	0.0156 (3)
C2	0.30637 (14)	0.91466 (13)	0.71055 (12)	0.0161 (3)
H2	0.3785	0.9560	0.7006	0.019*
C3	0.15713 (14)	0.99221 (13)	0.75193 (12)	0.0160 (3)
C4	0.05096 (14)	0.93156 (13)	0.76591 (12)	0.0162 (3)
C5	0.09440 (14)	0.79251 (14)	0.74196 (12)	0.0171 (3)
C6	0.24333 (14)	0.71422 (14)	0.69882 (13)	0.0177 (3)
H6	0.2728	0.6199	0.6798	0.021*
C7	0.50685 (14)	0.69379 (13)	0.64045 (12)	0.0162 (3)
C8	0.57985 (14)	0.56334 (13)	0.66514 (12)	0.0159 (3)
C9	0.52294 (14)	0.47110 (13)	0.74808 (13)	0.0162 (3)
C10	0.56894 (14)	0.33457 (13)	0.70935 (13)	0.0167 (3)
H10	0.6368	0.2988	0.6270	0.020*
C11	0.51433 (14)	0.25086 (13)	0.79276 (13)	0.0161 (3)
C12	0.41268 (14)	0.30331 (14)	0.91361 (13)	0.0172 (3)
C13	0.36502 (14)	0.44122 (14)	0.94987 (12)	0.0165 (3)
C14	0.42169 (14)	0.52434 (14)	0.86842 (13)	0.0170 (3)
H14	0.3916	0.6173	0.8946	0.020*
C15	0.20801 (16)	1.19411 (14)	0.76486 (15)	0.0231 (3)
H15A	0.2708	1.1587	0.8184	0.035*
H15B	0.1565	1.2907	0.7887	0.035*
H15C	0.2688	1.1785	0.6751	0.035*
C16	-0.17239 (16)	1.02141 (16)	0.93261 (14)	0.0265 (3)
H16A	-0.1634	0.9319	0.9605	0.040*
H16B	-0.2765	1.0745	0.9489	0.040*
H16C	-0.1297	1.0661	0.9799	0.040*
C17	0.02045 (17)	0.60063 (15)	0.75393 (16)	0.0279 (3)
H17A	0.0798	0.5621	0.6663	0.042*
H17B	-0.0694	0.5778	0.7781	0.042*
H17C	0.0771	0.5642	0.8119	0.042*
C18	0.64600 (16)	0.06014 (15)	0.63927 (14)	0.0242 (3)
H18A	0.6015	0.1072	0.5768	0.036*
H18B	0.6593	-0.0354	0.6287	0.036*
H18C	0.7417	0.0703	0.6259	0.036*
C19	0.38996 (17)	0.19362 (16)	1.10416 (14)	0.0247 (3)
H19A	0.4925	0.1338	1.0826	0.037*
H19B	0.3256	0.1506	1.1593	0.037*
H19C	0.3781	0.2780	1.1489	0.037*
C20	0.19984 (16)	0.62778 (14)	1.10034 (13)	0.0220 (3)
H20A	0.2772	0.6621	1.1044	0.033*
H20B	0.1235	0.6476	1.1837	0.033*
H20C	0.1564	0.6709	1.0358	0.033*

O1M	1.00187 (11)	0.66705 (10)	0.42773 (10)	0.0244 (2)
H1M	1.0849	0.6076	0.4149	0.037*
C1M	1.01117 (17)	0.79571 (15)	0.45053 (15)	0.0269 (3)
H1M1	0.9119	0.8626	0.4814	0.040*
H1M2	1.0672	0.8206	0.3710	0.040*
H1M3	1.0608	0.7922	0.5148	0.040*

Atomic displacement parameters (Å²)

	U^{11}	U^{22}	U^{33}	U^{12}	U^{13}	U^{23}
N1	0.0160 (5)	0.0170 (6)	0.0195 (6)	-0.0042 (4)	-0.0029 (4)	0.0023 (5)
N2	0.0155 (5)	0.0163 (6)	0.0228 (6)	-0.0052 (4)	-0.0011 (5)	0.0035 (5)
N3	0.0172 (5)	0.0164 (6)	0.0194 (6)	-0.0070 (4)	-0.0022 (4)	0.0033 (4)
O1	0.0167 (5)	0.0148 (5)	0.0270 (5)	-0.0046 (4)	-0.0063 (4)	0.0008 (4)
O2	0.0127 (4)	0.0229 (5)	0.0217 (5)	-0.0030 (4)	-0.0039 (4)	0.0033 (4)
O3	0.0166 (5)	0.0193 (5)	0.0281 (5)	-0.0088 (4)	-0.0047 (4)	0.0020 (4)
O4	0.0235 (5)	0.0141 (5)	0.0203 (5)	-0.0063 (4)	-0.0039 (4)	0.0011 (4)
O5	0.0322 (6)	0.0247 (5)	0.0171 (5)	-0.0179 (5)	-0.0077 (4)	0.0070 (4)
O6	0.0205 (5)	0.0186 (5)	0.0170 (5)	-0.0062 (4)	0.0003 (4)	0.0006 (4)
C1	0.0150 (6)	0.0161 (6)	0.0139 (6)	-0.0034 (5)	-0.0041 (5)	0.0031 (5)
C2	0.0161 (6)	0.0175 (7)	0.0160 (6)	-0.0074 (5)	-0.0052 (5)	0.0033 (5)
C3	0.0176 (6)	0.0147 (6)	0.0152 (6)	-0.0041 (5)	-0.0062 (5)	0.0030 (5)
C4	0.0138 (6)	0.0180 (7)	0.0156 (6)	-0.0034 (5)	-0.0053 (5)	0.0033 (5)
C5	0.0161 (6)	0.0219 (7)	0.0156 (6)	-0.0087 (5)	-0.0060 (5)	0.0042 (5)
C6	0.0188 (6)	0.0167 (7)	0.0178 (6)	-0.0069 (5)	-0.0054 (5)	0.0025 (5)
C7	0.0169 (6)	0.0164 (7)	0.0144 (6)	-0.0061 (5)	-0.0032 (5)	0.0014 (5)
C8	0.0142 (6)	0.0158 (6)	0.0154 (6)	-0.0043 (5)	-0.0022 (5)	-0.0007 (5)
C9	0.0147 (6)	0.0164 (7)	0.0179 (6)	-0.0050 (5)	-0.0061 (5)	0.0031 (5)
C10	0.0151 (6)	0.0172 (7)	0.0162 (6)	-0.0050 (5)	-0.0033 (5)	0.0012 (5)
C11	0.0167 (6)	0.0138 (6)	0.0195 (6)	-0.0055 (5)	-0.0080 (5)	0.0026 (5)
C12	0.0187 (6)	0.0196 (7)	0.0169 (6)	-0.0106 (5)	-0.0064 (5)	0.0050 (5)
C13	0.0140 (6)	0.0201 (7)	0.0151 (6)	-0.0062 (5)	-0.0040 (5)	0.0022 (5)
C14	0.0168 (6)	0.0153 (6)	0.0186 (6)	-0.0055 (5)	-0.0052 (5)	0.0015 (5)
C15	0.0216 (7)	0.0172 (7)	0.0318 (8)	-0.0079 (6)	-0.0089 (6)	0.0026 (6)
C16	0.0197 (7)	0.0293 (8)	0.0235 (7)	-0.0057 (6)	-0.0001 (6)	0.0031 (6)
C17	0.0246 (7)	0.0220 (8)	0.0373 (9)	-0.0128 (6)	-0.0047 (6)	0.0022 (6)
C18	0.0259 (7)	0.0177 (7)	0.0246 (7)	-0.0079 (6)	-0.0011 (6)	-0.0030 (6)
C19	0.0263 (7)	0.0267 (8)	0.0212 (7)	-0.0093 (6)	-0.0075 (6)	0.0089 (6)
C20	0.0215 (7)	0.0190 (7)	0.0191 (7)	-0.0030 (6)	-0.0017 (5)	-0.0013 (5)
O1M	0.0200 (5)	0.0177 (5)	0.0302 (6)	-0.0046 (4)	-0.0028 (4)	0.0040 (4)
C1M	0.0278 (7)	0.0227 (8)	0.0301 (8)	-0.0100 (6)	-0.0073 (6)	0.0020 (6)

Geometric parameters (Å, °)

N1—N2	1.3243 (16)	C10—C11	1.3970 (18)
N1—C7	1.3473 (16)	C10—H10	0.9500
N2—N3	1.3339 (16)	C11—C12	1.4010 (19)
N2—H2N	0.936 (17)	C12—C13	1.4002 (19)

N3—C8	1.3455 (16)	C13—C14	1.3895 (18)
O1—C3	1.3628 (16)	C14—H14	0.9500
O1—C15	1.4301 (16)	C15—H15A	0.9800
O2—C4	1.3768 (15)	C15—H15B	0.9800
O2—C16	1.4330 (17)	C15—H15C	0.9800
O3—C5	1.3645 (16)	C16—H16A	0.9800
O3—C17	1.4236 (17)	C16—H16B	0.9800
O4—C11	1.3661 (16)	C16—H16C	0.9800
O4—C18	1.4268 (17)	C17—H17A	0.9800
O5—C12	1.3755 (16)	C17—H17B	0.9800
O5—C19	1.4358 (17)	C17—H17C	0.9800
O6—C13	1.3639 (16)	C18—H18A	0.9800
O6—C20	1.4326 (16)	C18—H18B	0.9800
C1—C2	1.3950 (19)	C18—H18C	0.9800
C1—C6	1.4006 (18)	C19—H19A	0.9800
C1—C7	1.4774 (18)	C19—H19B	0.9800
C2—C3	1.3954 (18)	C19—H19C	0.9800
C2—H2	0.9500	C20—H20A	0.9800
C3—C4	1.3982 (19)	C20—H20B	0.9800
C4—C5	1.3953 (19)	C20—H20C	0.9800
C5—C6	1.3950 (18)	O1M—C1M	1.4282 (18)
C6—H6	0.9500	O1M—H1M	0.8400
C7—C8	1.4057 (19)	C1M—H1M1	0.9800
C8—C9	1.4743 (18)	C1M—H1M2	0.9800
C9—C10	1.3936 (19)	C1M—H1M3	0.9800
C9—C14	1.3970 (18)		
N2—N1—C7	104.56 (11)	O6—C13—C12	115.88 (11)
N1—N2—N3	114.49 (11)	C14—C13—C12	120.27 (12)
N1—N2—H2N	124.0 (10)	C13—C14—C9	119.97 (12)
N3—N2—H2N	121.4 (11)	C13—C14—H14	120.0
N2—N3—C8	105.06 (11)	C9—C14—H14	120.0
C3—O1—C15	116.70 (10)	O1—C15—H15A	109.5
C4—O2—C16	113.71 (10)	O1—C15—H15B	109.5
C5—O3—C17	117.30 (11)	H15A—C15—H15B	109.5
C11—O4—C18	116.84 (10)	O1—C15—H15C	109.5
C12—O5—C19	116.01 (11)	H15A—C15—H15C	109.5
C13—O6—C20	116.87 (10)	H15B—C15—H15C	109.5
C2—C1—C6	120.65 (12)	O2—C16—H16A	109.5
C2—C1—C7	119.59 (12)	O2—C16—H16B	109.5
C6—C1—C7	119.77 (12)	H16A—C16—H16B	109.5
C1—C2—C3	119.53 (12)	O2—C16—H16C	109.5
C1—C2—H2	120.2	H16A—C16—H16C	109.5
C3—C2—H2	120.2	H16B—C16—H16C	109.5
O1—C3—C2	124.68 (12)	O3—C17—H17A	109.5
O1—C3—C4	115.06 (11)	O3—C17—H17B	109.5
C2—C3—C4	120.25 (12)	H17A—C17—H17B	109.5
O2—C4—C5	120.10 (11)	O3—C17—H17C	109.5

O2—C4—C3	120.05 (12)	H17A—C17—H17C	109.5
C5—C4—C3	119.81 (12)	H17B—C17—H17C	109.5
O3—C5—C6	124.27 (12)	O4—C18—H18A	109.5
O3—C5—C4	115.33 (11)	O4—C18—H18B	109.5
C6—C5—C4	120.39 (12)	H18A—C18—H18B	109.5
C5—C6—C1	119.32 (12)	O4—C18—H18C	109.5
C5—C6—H6	120.3	H18A—C18—H18C	109.5
C1—C6—H6	120.3	H18B—C18—H18C	109.5
N1—C7—C8	108.49 (11)	O5—C19—H19A	109.5
N1—C7—C1	121.12 (12)	O5—C19—H19B	109.5
C8—C7—C1	130.39 (12)	H19A—C19—H19B	109.5
N3—C8—C7	107.40 (11)	O5—C19—H19C	109.5
N3—C8—C9	121.97 (12)	H19A—C19—H19C	109.5
C7—C8—C9	130.48 (12)	H19B—C19—H19C	109.5
C10—C9—C14	120.46 (12)	O6—C20—H20A	109.5
C10—C9—C8	121.60 (12)	O6—C20—H20B	109.5
C14—C9—C8	117.94 (12)	H20A—C20—H20B	109.5
C9—C10—C11	119.35 (12)	O6—C20—H20C	109.5
C9—C10—H10	120.3	H20A—C20—H20C	109.5
C11—C10—H10	120.3	H20B—C20—H20C	109.5
O4—C11—C10	124.35 (12)	C1M—O1M—H1M	109.5
O4—C11—C12	115.03 (11)	O1M—C1M—H1M1	109.5
C10—C11—C12	120.62 (12)	O1M—C1M—H1M2	109.5
O5—C12—C13	121.12 (12)	H1M1—C1M—H1M2	109.5
O5—C12—C11	119.30 (12)	O1M—C1M—H1M3	109.5
C13—C12—C11	119.29 (12)	H1M1—C1M—H1M3	109.5
O6—C13—C14	123.85 (12)	H1M2—C1M—H1M3	109.5
C7—N1—N2—N3	-0.41 (15)	N2—N3—C8—C9	-175.64 (12)
N1—N2—N3—C8	0.07 (16)	N1—C7—C8—N3	-0.54 (15)
C6—C1—C2—C3	-0.27 (19)	C1—C7—C8—N3	-179.60 (13)
C7—C1—C2—C3	-179.82 (12)	N1—C7—C8—C9	174.91 (13)
C15—O1—C3—C2	3.37 (19)	C1—C7—C8—C9	-4.1 (2)
C15—O1—C3—C4	-177.50 (12)	N3—C8—C9—C10	-47.04 (19)
C1—C2—C3—O1	178.74 (12)	C7—C8—C9—C10	138.07 (15)
C1—C2—C3—C4	-0.36 (19)	N3—C8—C9—C14	132.31 (14)
C16—O2—C4—C5	85.75 (15)	C7—C8—C9—C14	-42.6 (2)
C16—O2—C4—C3	-96.54 (15)	C14—C9—C10—C11	-0.87 (19)
O1—C3—C4—O2	4.99 (18)	C8—C9—C10—C11	178.47 (12)
C2—C3—C4—O2	-175.83 (11)	C18—O4—C11—C10	-4.16 (18)
O1—C3—C4—C5	-177.30 (11)	C18—O4—C11—C12	175.12 (12)
C2—C3—C4—C5	1.9 (2)	C9—C10—C11—O4	-179.85 (12)
C17—O3—C5—C6	8.57 (19)	C9—C10—C11—C12	0.91 (19)
C17—O3—C5—C4	-172.45 (12)	C19—O5—C12—C13	-72.53 (17)
O2—C4—C5—O3	-4.09 (18)	C19—O5—C12—C11	113.70 (14)
C3—C4—C5—O3	178.19 (12)	O4—C11—C12—O5	-4.85 (18)
O2—C4—C5—C6	174.92 (12)	C10—C11—C12—O5	174.46 (11)
C3—C4—C5—C6	-2.79 (19)	O4—C11—C12—C13	-178.74 (11)

O3—C5—C6—C1	-178.92 (12)	C10—C11—C12—C13	0.6 (2)
C4—C5—C6—C1	2.16 (19)	C20—O6—C13—C14	6.21 (18)
C2—C1—C6—C5	-0.6 (2)	C20—O6—C13—C12	-173.47 (11)
C7—C1—C6—C5	178.93 (12)	O5—C12—C13—O6	3.80 (18)
N2—N1—C7—C8	0.56 (15)	C11—C12—C13—O6	177.57 (11)
N2—N1—C7—C1	179.72 (12)	O5—C12—C13—C14	-175.89 (12)
C2—C1—C7—N1	-34.27 (18)	C11—C12—C13—C14	-2.11 (19)
C6—C1—C7—N1	146.18 (13)	O6—C13—C14—C9	-177.50 (12)
C2—C1—C7—C8	144.69 (15)	C12—C13—C14—C9	2.16 (19)
C6—C1—C7—C8	-34.9 (2)	C10—C9—C14—C13	-0.66 (19)
N2—N3—C8—C7	0.28 (15)	C8—C9—C14—C13	179.98 (12)

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>
N2—H2N...O1 <i>M</i>	0.936 (17)	1.797 (18)	2.7303 (15)	174.2 (16)
O1 <i>M</i> —H1 <i>M</i> ...N3 ⁱ	0.84	1.97	2.8101 (15)	177

Symmetry code: (i) $-x+2, -y+1, -z+1$.