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## Structure Reports

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**(1*S*,2*R*,3*R*)-(–)-2-Hydroxy-3-morpholino-3-phenyl-*N*-(1'-phenylethyl)propionamide**

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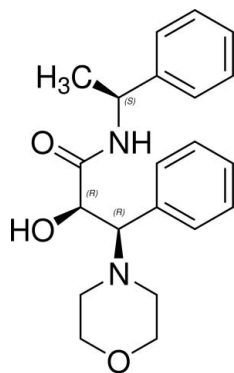
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Key indicators: single-crystal X-ray study;  $T = 293$  K; mean  $\sigma(\text{C}-\text{C}) = 0.007$  Å;  $R$  factor = 0.054;  $wR$  factor = 0.163; data-to-parameter ratio = 11.8.

In the title compound,  $\text{C}_{21}\text{H}_{26}\text{N}_2\text{O}_3$ , the morpholine ring has a chair conformation and the dihedral angle between the two phenyl rings is  $59.0(3)^\circ$ . The crystal packing is stabilized by intermolecular  $\text{O}-\text{H}\cdots\text{O}$  hydrogen bonds, generating a ribbon structure along the  $a$  axis. An intramolecular  $\text{N}-\text{H}\cdots\text{O}$  contact is also present.

## Related literature

For general background, see: Barbaro *et al.* (1992); Szymanski *et al.* (2006); Sheppard *et al.* (2004); Chen *et al.* (1996); Concellón *et al.* (2003*a,b*); Martín *et al.* (2004). For related structures, see: Romero *et al.* (2005*a,b*). For ring conformation analysis, see: Cremer & Pople (1975).



## Experimental

## Crystal data

 $\text{C}_{21}\text{H}_{26}\text{N}_2\text{O}_3$  $M_r = 354.44$ Orthorhombic,  $P2_12_12_1$  $a = 6.0010(17)$  Å $b = 15.659(3)$  Å $c = 20.746(4)$  Å $V = 1949.5(8)$  Å<sup>3</sup> $Z = 4$ Mo  $K\alpha$  radiation $\mu = 0.08$  mm<sup>-1</sup> $T = 293$  K $0.72 \times 0.28 \times 0.16$  mm

## Data collection

Bruker P4 diffractometer

Absorption correction: none

3964 measured reflections

2959 independent reflections

1116 reflections with  $I > 2\sigma(I)$  $R_{\text{int}} = 0.055$ 

3 standard reflections

every 97 reflections

intensity decay: 3%

## Refinement

 $R[F^2 > 2\sigma(F^2)] = 0.054$  $wR(F^2) = 0.163$  $S = 0.89$ 

2959 reflections

251 parameters

H atoms treated by a mixture of independent and constrained refinement

 $\Delta\rho_{\text{max}} = 0.24$  e Å<sup>-3</sup> $\Delta\rho_{\text{min}} = -0.19$  e Å<sup>-3</sup>

Table 1

Hydrogen-bond geometry (Å, °).

$D-H\cdots A$	$D-H$	$H\cdots A$	$D\cdots A$	$D-H\cdots A$
$\text{N1}-\text{H1N}\cdots\text{O1}$	1.00 (7)	1.92 (7)	2.569 (5)	120 (5)
$\text{O1}-\text{H1O}\cdots\text{O2}^1$	0.98 (8)	1.80 (8)	2.753 (4)	163 (7)

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

Data collection: XSCANS (Siemens, 1994); cell refinement: XSCANS; data reduction: XSCANS; program(s) used to solve structure: SIR2004 (Burla *et al.*, 2005); program(s) used to refine structure: SHELXL97 (Sheldrick, 2008); molecular graphics: ORTEP-3 for Windows (Farrugia, 1997); software used to prepare material for publication: WinGX (Farrugia, 1999).

We acknowledge financial support under scholarship DMAS No. 169011 and CONACyT project No. 83185. Special thanks go to Dr Marcos Flores (USAI-FQ-UNAM) for useful comments.

Supplementary data and figures for this paper are available from the IUCr electronic archives (Reference: IS2398).

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**supplementary materials**

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**(1'S,2R,3R)-(-)-2-Hydroxy-3-morpholino-3-phenyl-N-(1'-phenylethyl)propionamide**

**A. Mendoza, D. M. Aparicio, J. L. Terán, D. Gnecco and J. R. Juárez**

**Comment**

The stereoselective synthesis of  $\alpha$ -hydroxyamides (Barbaro *et al.*, 1992; Szymanski *et al.*, 2006) is an important area in asymmetric synthesis because these kinds of intermediates exhibit a great value as synthetic building block in the synthesis of pharmaceutical, agriculture and medicinal compounds (Sheppard *et al.*, 2004; Chen *et al.*, 1996). Numerous methods have been development through the last 10 years. One of the most relevant methodologies is the opening reaction of epoxyamides (Concellón *et al.*, 2003*a,b*; Martín *et al.*, 2004). In this way, we show herein the regiospecific ring opening reaction of epoxyamide derived from (*S*)-(-)-phenyl ethyl amine (Scheme 1). Related structures including (*S*)-(-)-phenyl ethyl amine with different substituents has been previously reported (Romero *et al.*, 2005*a,b*).

In the present paper, we report the structure of title compound, (I), which shows a single asymmetric unit with two aromatic rings. The morpholine ring shows an almost perfect chair conformation with a puckering parameters (Cremer & Pople, 1975) (O3—C2—C1—N2—C4—C3)  $Q = 0.572$  (4) Å,  $\theta_2 = 3.6^\circ$  (4),  $\varphi_2 = 343^\circ$  (10),  $q_2 = 0.026$  (4) Å and  $q_3 = 0.571$  (4) Å. In the crystal structure, the molecules are linked by O—H...O hydrogen bonds (Table 1), generating a ribbon structure along the *a* axis.

**Experimental**

The compound (I) was obtained by *trans*-diastereoisomeric mixture of epoxyamide in anhydrous EtOH. Then, 1.1 equivalents of morpholine were added. The reaction was stirred over night. Finally, the resultant amino alcohol mixture was separate by column chromatography (AcOEt: Petroleum Ether). The absolute configuration of (1'*S*, 2*R*, 3*R*)-(-)-2-hydroxy-3-morpholin-4-yl-3-phenyl-*N*-(1'-phenyl-ethyl)-propionamide was established by the structure determination of a (*S*)-(-)-phenyl ethyl amine of known absolute configuration of starting material [C8 in compound (I)]. The  $H^1$  NMR experiment shows only one diastereomeric compound.

**Refinement**

H atoms bonded to N and O atoms and all optically relevant were located in a difference map and refined as riding on their parent atoms with  $U_{iso}(H) = 1.2U_{eq}$  for N and C—H and  $U_{iso}(H) = 1.5U_{eq}$  for O atoms. H atoms linked to C atoms were placed in geometrical idealized positions and refined as riding on their parent atoms, with C—H = 0.93–0.98 Å and with  $U_{iso}(H) = 1.2U_{eq}(C)$  or  $1.5U_{eq}(\text{methyl C})$ . In the absence of significant anomalous scattering effects, Friedel pairs were merged and the absolute configuration was assigned on the base of synthetic procedure.

## Figures

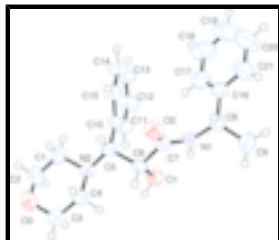


Fig. 1. The molecular structure of the title compound, with atom labels and 50% probability displacement ellipsoids for non-H atoms.

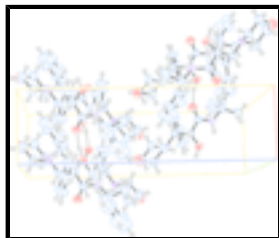


Fig. 2. The packing of the title compound, showing molecules connected by O—H...O hydrogen bonds (dashed lines).

### (1'S,2R,3R)-(-)-2-Hydroxy-3-morpholino-3-phenyl- N-(1'-phenylethyl)propionamide

#### Crystal data

$C_{21}H_{26}N_2O_3$

$M_r = 354.44$

Orthorhombic,  $P2_12_12_1$

Hall symbol: P 2ac 2ab

$a = 6.0010 (17) \text{ \AA}$

$b = 15.659 (3) \text{ \AA}$

$c = 20.746 (4) \text{ \AA}$

$V = 1949.5 (8) \text{ \AA}^3$

$Z = 4$

$F_{000} = 760$

$D_x = 1.208 \text{ Mg m}^{-3}$

Melting point: 431 K

Mo  $K\alpha$  radiation

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

Cell parameters from 30 reflections

$\theta = 3.9\text{--}23.9^\circ$

$\mu = 0.08 \text{ mm}^{-1}$

$T = 293 \text{ K}$

Prism, colorless

$0.72 \times 0.28 \times 0.16 \text{ mm}$

#### Data collection

Bruker P4  
diffractometer

Radiation source: fine-focus sealed tube

Monochromator: graphite

$T = 293 \text{ K}$

$2\theta/\omega$  scans

Absorption correction: none

3964 measured reflections

2959 independent reflections

1116 reflections with  $I > 2\sigma(I)$

$R_{\text{int}} = 0.055$

$\theta_{\text{max}} = 29.0^\circ$

$\theta_{\text{min}} = 1.6^\circ$

$h = -1 \rightarrow 8$

$k = -1 \rightarrow 21$

$l = -1 \rightarrow 28$

3 standard reflections

every 97 reflections

intensity decay: 3%

*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
$R[F^2 > 2\sigma(F^2)] = 0.054$	$w = 1/[\sigma^2(F_o^2) + (0.0711P)^2]$ where $P = (F_o^2 + 2F_c^2)/3$
$wR(F^2) = 0.163$	$(\Delta/\sigma)_{\max} < 0.001$
$S = 0.89$	$\Delta\rho_{\max} = 0.24 \text{ e } \text{\AA}^{-3}$
2959 reflections	$\Delta\rho_{\min} = -0.19 \text{ e } \text{\AA}^{-3}$
251 parameters	Extinction correction: SHELXL97 (Sheldrick, 2008)
Primary atom site location: structure-invariant direct methods	Extinction coefficient: 0.008 (2)
Secondary atom site location: difference Fourier map	

*Special details*

**Geometry.** All s.u.'s (except the s.u. in the dihedral angle between two l.s. planes) are estimated using the full covariance matrix. The cell s.u.'s are taken into account individually in the estimation of s.u.'s in distances, angles and torsion angles; correlations between s.u.'s in cell parameters are only used when they are defined by crystal symmetry. An approximate (isotropic) treatment of cell s.u.'s is used for estimating s.u.'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  $F^2 > 2\sigma(F^2)$  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>	$U_{\text{iso}}^*/U_{\text{eq}}$
O1	-0.1563 (5)	0.92034 (19)	0.75127 (13)	0.0567 (8)
N2	-0.1559 (5)	0.90373 (19)	0.60796 (14)	0.0464 (8)
O2	0.4127 (5)	0.96834 (18)	0.72379 (13)	0.0598 (8)
N1	0.2201 (6)	0.9177 (2)	0.81024 (16)	0.0516 (9)
C10	0.0914 (6)	0.8018 (2)	0.66365 (17)	0.0447 (9)
C7	0.2354 (6)	0.9439 (2)	0.74924 (19)	0.0445 (9)
O3	-0.3564 (6)	0.9556 (2)	0.48787 (14)	0.0770 (10)
C5	0.0430 (7)	0.8947 (2)	0.64872 (18)	0.0449 (9)
C6	0.0197 (6)	0.9474 (2)	0.71116 (18)	0.0434 (9)
C4	-0.2007 (7)	0.9933 (2)	0.5920 (2)	0.0554 (11)
H4A	-0.0697	1.0183	0.5721	0.066*
H4B	-0.2322	1.0248	0.6312	0.066*
C8	0.3993 (7)	0.9216 (3)	0.8565 (2)	0.0552 (11)
C15	0.3024 (8)	0.7691 (2)	0.65361 (19)	0.0535 (10)
H15	0.4145	0.8045	0.6380	0.064*
C16	0.5180 (7)	0.8364 (3)	0.86521 (19)	0.0558 (11)

## supplementary materials

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C11	-0.0702 (7)	0.7484 (3)	0.6885 (2)	0.0610 (11)
H11	-0.2125	0.7695	0.6964	0.073*
C17	0.4526 (9)	0.7626 (3)	0.8349 (2)	0.0703 (13)
H17	0.3292	0.7635	0.8078	0.084*
C3	-0.3969 (8)	1.0009 (3)	0.5463 (2)	0.0676 (13)
H3A	-0.5294	0.9782	0.5669	0.081*
H3B	-0.4235	1.0607	0.5367	0.081*
C1	-0.1284 (8)	0.8570 (3)	0.54712 (19)	0.0637 (12)
H1A	-0.1097	0.7967	0.5564	0.076*
H1B	0.0055	0.8768	0.5257	0.076*
C2	-0.3229 (9)	0.8683 (3)	0.5030 (2)	0.0765 (15)
H2A	-0.2973	0.8365	0.4635	0.092*
H2B	-0.4559	0.8457	0.5233	0.092*
C13	0.1855 (9)	0.6310 (3)	0.6898 (2)	0.0699 (14)
H13	0.2158	0.5736	0.6973	0.084*
C14	0.3492 (8)	0.6836 (3)	0.6666 (2)	0.0638 (12)
H14	0.4918	0.6623	0.6594	0.077*
C21	0.7048 (9)	0.8339 (4)	0.9047 (3)	0.0896 (17)
H21	0.7533	0.8835	0.9250	0.107*
C9	0.3070 (9)	0.9542 (4)	0.9206 (2)	0.0938 (18)
H9A	0.2331	1.0078	0.9138	0.141*
H9B	0.4271	0.9618	0.9506	0.141*
H9C	0.2027	0.9135	0.9375	0.141*
C12	-0.0219 (9)	0.6627 (3)	0.7018 (3)	0.0742 (14)
H12	-0.1316	0.6274	0.7189	0.089*
C18	0.5674 (12)	0.6867 (3)	0.8440 (3)	0.0958 (18)
H18	0.5207	0.6370	0.8236	0.115*
C19	0.7512 (13)	0.6857 (5)	0.8837 (4)	0.119 (2)
H19	0.8297	0.6351	0.8899	0.143*
C20	0.8184 (12)	0.7586 (5)	0.9141 (3)	0.122 (2)
H20	0.9418	0.7574	0.9412	0.147*
H1O	-0.307 (14)	0.929 (5)	0.735 (4)	0.184*
H1N	0.059 (12)	0.906 (4)	0.819 (3)	0.147*
H5	0.180 (12)	0.915 (4)	0.627 (3)	0.147*
H6	-0.004 (11)	1.009 (4)	0.697 (3)	0.147*
H8	0.535 (12)	0.960 (4)	0.839 (3)	0.147*

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

	$U^{11}$	$U^{22}$	$U^{33}$	$U^{12}$	$U^{13}$	$U^{23}$
O1	0.0370 (14)	0.0800 (19)	0.0531 (15)	0.0034 (16)	0.0045 (14)	0.0008 (15)
N2	0.0476 (19)	0.0465 (17)	0.0450 (16)	0.0034 (17)	-0.0023 (16)	0.0024 (14)
O2	0.0429 (17)	0.0697 (18)	0.0668 (18)	-0.0070 (15)	0.0011 (15)	0.0061 (15)
N1	0.0388 (18)	0.065 (2)	0.0507 (19)	0.0006 (18)	-0.0067 (17)	-0.0002 (17)
C10	0.041 (2)	0.045 (2)	0.049 (2)	0.003 (2)	0.0017 (19)	0.0029 (17)
C7	0.038 (2)	0.044 (2)	0.051 (2)	0.0032 (18)	-0.001 (2)	-0.0001 (19)
O3	0.091 (3)	0.086 (2)	0.0543 (17)	0.013 (2)	-0.0085 (19)	0.0103 (16)
C5	0.041 (2)	0.046 (2)	0.048 (2)	0.0057 (19)	0.0005 (19)	0.0005 (17)

C6	0.034 (2)	0.047 (2)	0.049 (2)	0.0020 (19)	0.0026 (18)	0.0012 (18)
C4	0.058 (3)	0.050 (2)	0.058 (2)	0.007 (2)	-0.009 (2)	0.008 (2)
C8	0.047 (2)	0.064 (3)	0.055 (2)	0.008 (2)	-0.011 (2)	-0.009 (2)
C15	0.050 (2)	0.052 (2)	0.058 (2)	0.002 (2)	0.005 (2)	-0.001 (2)
C16	0.048 (3)	0.067 (3)	0.053 (2)	0.005 (2)	0.002 (2)	0.003 (2)
C11	0.048 (3)	0.050 (2)	0.085 (3)	0.002 (2)	0.008 (3)	0.009 (2)
C17	0.069 (3)	0.068 (3)	0.074 (3)	0.007 (3)	0.004 (3)	0.008 (3)
C3	0.073 (3)	0.072 (3)	0.058 (2)	0.016 (3)	-0.003 (3)	0.005 (2)
C1	0.080 (3)	0.061 (2)	0.051 (2)	0.007 (3)	-0.003 (3)	-0.010 (2)
C2	0.083 (4)	0.088 (3)	0.058 (3)	0.004 (3)	-0.016 (3)	-0.010 (3)
C13	0.072 (3)	0.047 (2)	0.090 (3)	0.006 (3)	-0.002 (3)	0.008 (2)
C14	0.057 (3)	0.051 (2)	0.084 (3)	0.016 (2)	0.004 (3)	-0.004 (2)
C21	0.075 (4)	0.098 (4)	0.096 (4)	0.018 (4)	-0.030 (3)	0.003 (3)
C9	0.080 (3)	0.134 (5)	0.068 (3)	0.034 (4)	-0.018 (3)	-0.042 (3)
C12	0.063 (3)	0.052 (3)	0.107 (4)	-0.003 (2)	0.003 (3)	0.018 (3)
C18	0.108 (5)	0.063 (3)	0.116 (4)	0.014 (4)	0.016 (4)	0.011 (3)
C19	0.114 (6)	0.094 (5)	0.149 (6)	0.045 (5)	-0.001 (5)	0.025 (5)
C20	0.104 (5)	0.125 (5)	0.137 (6)	0.035 (5)	-0.042 (5)	0.025 (5)

*Geometric parameters (Å, °)*

O1—C6	1.410 (5)	C16—C21	1.389 (6)
O1—H10	0.98 (8)	C11—C12	1.401 (6)
N2—C4	1.466 (5)	C11—H11	0.9300
N2—C1	1.468 (5)	C17—C18	1.387 (7)
N2—C5	1.470 (5)	C17—H17	0.9300
O2—C7	1.248 (4)	C3—H3A	0.9700
N1—C7	1.333 (5)	C3—H3B	0.9700
N1—C8	1.443 (5)	C1—C2	1.494 (7)
N1—H1N	1.00 (7)	C1—H1A	0.9700
C10—C11	1.380 (5)	C1—H1B	0.9700
C10—C15	1.382 (6)	C2—H2A	0.9700
C10—C5	1.514 (5)	C2—H2B	0.9700
C7—C6	1.517 (5)	C13—C12	1.363 (7)
O3—C2	1.416 (5)	C13—C14	1.370 (6)
O3—C3	1.426 (5)	C13—H13	0.9300
C5—C6	1.542 (5)	C14—H14	0.9300
C5—H5	0.99 (7)	C21—C20	1.376 (8)
C6—H6	1.02 (6)	C21—H21	0.9300
C4—C3	1.516 (6)	C9—H9A	0.9600
C4—H4A	0.9700	C9—H9B	0.9600
C4—H4B	0.9700	C9—H9C	0.9600
C8—C16	1.523 (6)	C12—H12	0.9300
C8—C9	1.528 (6)	C18—C19	1.376 (9)
C8—H8	1.08 (7)	C18—H18	0.9300
C15—C14	1.393 (5)	C19—C20	1.365 (9)
C15—H15	0.9300	C19—H19	0.9300
C16—C17	1.373 (6)	C20—H20	0.9300
C6—O1—H10	116 (4)	C16—C17—H17	119.5

## supplementary materials

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C4—N2—C1	107.6 (3)	C18—C17—H17	119.5
C4—N2—C5	111.8 (3)	O3—C3—C4	111.1 (4)
C1—N2—C5	110.8 (3)	O3—C3—H3A	109.4
C7—N1—C8	124.5 (4)	C4—C3—H3A	109.4
C7—N1—H1N	107 (4)	O3—C3—H3B	109.4
C8—N1—H1N	127 (4)	C4—C3—H3B	109.4
C11—C10—C15	118.4 (4)	H3A—C3—H3B	108.0
C11—C10—C5	121.5 (4)	N2—C1—C2	112.3 (4)
C15—C10—C5	120.1 (4)	N2—C1—H1A	109.1
O2—C7—N1	123.7 (4)	C2—C1—H1A	109.1
O2—C7—C6	119.7 (3)	N2—C1—H1B	109.1
N1—C7—C6	116.5 (3)	C2—C1—H1B	109.1
C2—O3—C3	108.5 (3)	H1A—C1—H1B	107.9
N2—C5—C10	111.5 (3)	O3—C2—C1	111.2 (4)
N2—C5—C6	111.0 (3)	O3—C2—H2A	109.4
C10—C5—C6	111.1 (3)	C1—C2—H2A	109.4
N2—C5—H5	112 (4)	O3—C2—H2B	109.4
C10—C5—H5	104 (4)	C1—C2—H2B	109.4
C6—C5—H5	107 (4)	H2A—C2—H2B	108.0
O1—C6—C7	108.7 (3)	C12—C13—C14	120.0 (4)
O1—C6—C5	113.8 (3)	C12—C13—H13	120.0
C7—C6—C5	109.9 (3)	C14—C13—H13	120.0
O1—C6—H6	110 (4)	C13—C14—C15	120.1 (4)
C7—C6—H6	108 (4)	C13—C14—H14	120.0
C5—C6—H6	106 (4)	C15—C14—H14	120.0
N2—C4—C3	111.0 (3)	C20—C21—C16	120.5 (6)
N2—C4—H4A	109.4	C20—C21—H21	119.8
C3—C4—H4A	109.4	C16—C21—H21	119.8
N2—C4—H4B	109.4	C8—C9—H9A	109.5
C3—C4—H4B	109.4	C8—C9—H9B	109.5
H4A—C4—H4B	108.0	H9A—C9—H9B	109.5
N1—C8—C16	113.0 (3)	C8—C9—H9C	109.5
N1—C8—C9	108.8 (3)	H9A—C9—H9C	109.5
C16—C8—C9	111.1 (4)	H9B—C9—H9C	109.5
N1—C8—H8	112 (3)	C13—C12—C11	120.1 (5)
C16—C8—H8	100 (3)	C13—C12—H12	119.9
C9—C8—H8	112 (3)	C11—C12—H12	119.9
C10—C15—C14	120.8 (4)	C19—C18—C17	119.4 (6)
C10—C15—H15	119.6	C19—C18—H18	120.3
C14—C15—H15	119.6	C17—C18—H18	120.3
C17—C16—C21	118.5 (5)	C20—C19—C18	120.2 (6)
C17—C16—C8	123.3 (4)	C20—C19—H19	119.9
C21—C16—C8	118.1 (4)	C18—C19—H19	119.9
C10—C11—C12	120.6 (4)	C19—C20—C21	120.4 (6)
C10—C11—H11	119.7	C19—C20—H20	119.8
C12—C11—H11	119.7	C21—C20—H20	119.8
C16—C17—C18	121.1 (5)		
C8—N1—C7—O2	6.1 (6)	N1—C8—C16—C17	-3.5 (6)
C8—N1—C7—C6	-170.8 (3)	C9—C8—C16—C17	119.1 (5)



C4—N2—C5—C10	-178.6 (3)	N1—C8—C16—C21	175.0 (4)
C1—N2—C5—C10	-58.5 (4)	C9—C8—C16—C21	-62.5 (5)
C4—N2—C5—C6	57.0 (4)	C15—C10—C11—C12	-1.3 (6)
C1—N2—C5—C6	177.1 (3)	C5—C10—C11—C12	179.9 (4)
C11—C10—C5—N2	-51.9 (5)	C21—C16—C17—C18	1.0 (7)
C15—C10—C5—N2	129.3 (4)	C8—C16—C17—C18	179.5 (4)
C11—C10—C5—C6	72.4 (5)	C2—O3—C3—C4	60.3 (5)
C15—C10—C5—C6	-106.4 (4)	N2—C4—C3—O3	-59.3 (5)
O2—C7—C6—O1	-177.3 (3)	C4—N2—C1—C2	-54.6 (5)
N1—C7—C6—O1	-0.2 (4)	C5—N2—C1—C2	-177.1 (4)
O2—C7—C6—C5	57.6 (5)	C3—O3—C2—C1	-59.9 (5)
N1—C7—C6—C5	-125.4 (3)	N2—C1—C2—O3	59.0 (5)
N2—C5—C6—O1	63.3 (4)	C12—C13—C14—C15	-1.7 (7)
C10—C5—C6—O1	-61.3 (4)	C10—C15—C14—C13	-0.3 (7)
N2—C5—C6—C7	-174.5 (3)	C17—C16—C21—C20	-1.3 (8)
C10—C5—C6—C7	60.9 (4)	C8—C16—C21—C20	-179.8 (6)
C1—N2—C4—C3	54.3 (5)	C14—C13—C12—C11	2.1 (8)
C5—N2—C4—C3	176.2 (3)	C10—C11—C12—C13	-0.6 (7)
C7—N1—C8—C16	-99.7 (5)	C16—C17—C18—C19	-0.6 (8)
C7—N1—C8—C9	136.4 (4)	C17—C18—C19—C20	0.4 (10)
C11—C10—C15—C14	1.7 (6)	C18—C19—C20—C21	-0.7 (11)
C5—C10—C15—C14	-179.4 (4)	C16—C21—C20—C19	1.1 (10)

*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>
N1—H1N···O1	1.00 (7)	1.92 (7)	2.569 (5)	120 (5)
O1—H1O···O2 <sup>i</sup>	0.98 (8)	1.80 (8)	2.753 (4)	163 (7)

Symmetry codes: (i)  $x-1, y, z$ .

Fig. 1

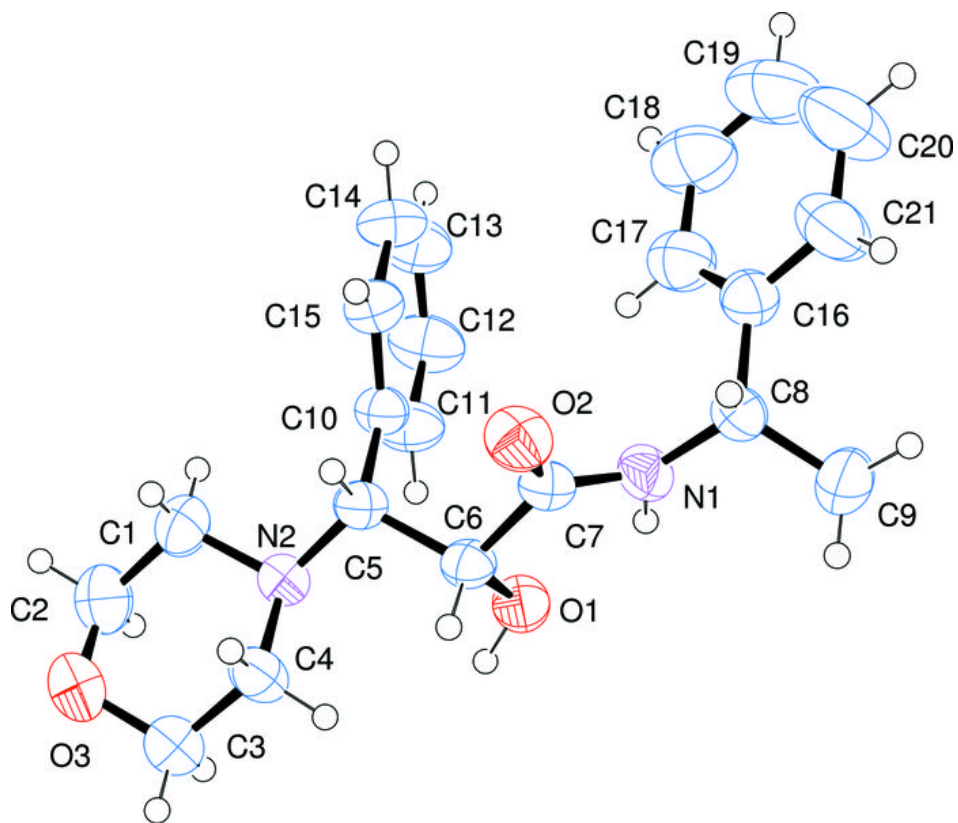


Fig. 2

