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## Structure Reports

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1-[(Pyrrolidin-1-yl)(*p*-tolyl)methyl]-naphthalen-2-ol

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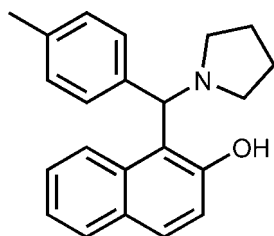
Received 2 September 2008; accepted 5 September 2008

Key indicators: single-crystal X-ray study;  $T = 293$  K; mean  $\sigma(\text{C}-\text{C}) = 0.003$  Å;  
 $R$  factor = 0.069;  $wR$  factor = 0.190; data-to-parameter ratio = 18.7.

In the title compound,  $\text{C}_{22}\text{H}_{23}\text{NO}$ , the dihedral angle between the naphthyl ring system and the benzene ring is  $73.32(6)^\circ$ . An intramolecular  $\text{O}-\text{H}\cdots\text{N}$  hydrogen bond stabilizes the molecular conformation. In the crystal structure, molecules are linked by  $\text{C}-\text{H}\cdots\pi$  interactions, resulting in zigzag chains parallel to the  $[10\bar{1}]$  direction.

## Related literature

For general background on the chemistry of naphthalen-2-ol derivatives, see: Szatmari & Fulop (2004); Zhao & Sun (2005). For puckering and asymmetry parameters, see: Cremer & Pople (1975); Nardelli (1983).



## Experimental

## Crystal data

$\text{C}_{22}\text{H}_{23}\text{NO}$   
 $M_r = 317.41$   
Monoclinic,  $P2_1/n$   
 $a = 10.3467(18)$  Å  
 $b = 16.055(3)$  Å  
 $c = 11.252(2)$  Å  
 $\beta = 106.810(8)^\circ$   
 $V = 1789.2(6)$  Å<sup>3</sup>  
 $Z = 4$   
Mo  $K\alpha$  radiation  
 $\mu = 0.07$  mm<sup>-1</sup>  
 $T = 293(2)$  K  
 $0.25 \times 0.22 \times 0.20$  mm

## Data collection

Rigaku SCXmini diffractometer  
Absorption correction: multi-scan  
(*CrystalClear*; Rigaku, 2005)  
 $T_{\min} = 0.963$ ,  $T_{\max} = 0.989$   
18171 measured reflections  
4086 independent reflections  
2547 reflections with  $I > 2\sigma(I)$   
 $R_{\text{int}} = 0.058$

## Refinement

$R[F^2 > 2\sigma(F^2)] = 0.069$   
 $wR(F^2) = 0.190$   
 $S = 1.06$   
4086 reflections  
219 parameters  
H-atom parameters constrained  
 $\Delta\rho_{\max} = 0.25$  e Å<sup>-3</sup>  
 $\Delta\rho_{\min} = -0.20$  e Å<sup>-3</sup>

Table 1

Hydrogen-bond geometry (Å, °).

$D-H\cdots A$	$D-H$	$H\cdots A$	$D\cdots A$	$D-H\cdots A$
$\text{O1}-\text{H1}\cdots\text{N1}$	0.82	1.88	2.600 (3)	145
$\text{C18}-\text{H18}\cdots\text{Cg1}^i$	0.93	2.66	3.588 (8)	173

Symmetry code: (i)  $x - \frac{1}{2}, -y + \frac{1}{2}, z + \frac{1}{2}$ . Cg1 is the centroid of the C5-C10 benzene ring.

Data collection: *CrystalClear* (Rigaku, 2005); cell refinement: *CrystalClear*; data reduction: *CrystalClear*; program(s) used to solve structure: *SHELXS97* (Sheldrick, 2008); program(s) used to refine structure: *SHELXL97* (Sheldrick, 2008); molecular graphics: *SHELXTL/PC* (Sheldrick, 2008); software used to prepare material for publication: *SHELXTL/PC*.

This work was supported by a start-up grant from Southeast University to HZ.

Supplementary data and figures for this paper are available from the IUCr electronic archives (Reference: RZ2244).

## References

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Zhao, B. & Sun, Y.-X. (2005). *Acta Cryst.* **E61**, m652–m653.

## supporting information

*Acta Cryst.* (2008). E64, o1926 [doi:10.1107/S1600536808028493]

**1-[(Pyrrolidin-1-yl)(*p*-tolyl)methyl]naphthalen-2-ol****Chuanwei Wan and Hong Zhao****S1. Comment**

Compounds derived from naphthalen-2-ol have been of great interest in organic chemistry (Szatmari & Fulop, 2004; Zhao & Sun, 2005). We report here the crystal structure of the title compound (Fig. 1).

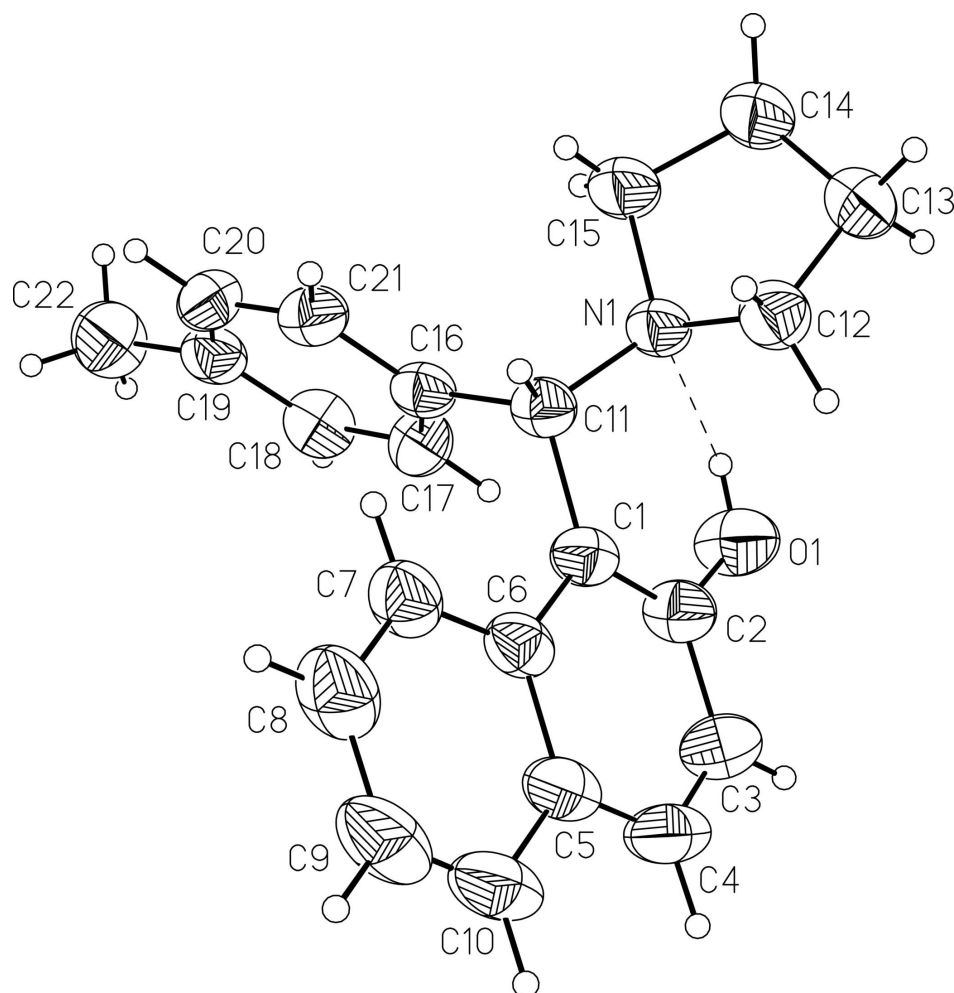
Bond lengths and angles in the title compound have normal values. The dihedral angle between the naphthyl and phenyl rings is 73.32 (6)°. The pyrrolidine ring adopts a twist conformation, as indicated by the puckering parameters ( $q_2 = 0.401$  (2) Å and  $\varphi = 169.2$  (4)°; Cremer & Pople, 1975) and the small value of the displacement asymmetry parameter ( $\Delta C_2(C14) = 0.0301$  (10)°; Nardelli, 1983). The molecular conformation is stabilized by a strong intramolecular O—H $\cdots$ N hydrogen bond (Table 1). In the crystal packing, molecules are linked through C—H $\cdots$  $\pi$  interactions (Table 1) to form zig zag chains running along the [1 0 -1] direction.

**S2. Experimental**

A dry 50 ml flask was charged with benzaldehyde (10 mmol), naphthalen-2-ol (10 mmol), and pyrrolidine (10 mmol). The mixture was stirred at 100°C for 10 h then ethanol (15 ml) was added. After heating under reflux for 30 minutes, the precipitate was filtrated off and washed 3 times with ethanol to give the title compound. Single crystals suitable for X-ray analysis were obtained by slow evaporation of a dichloromethane solution.

**S3. Refinement**

All hydrogen atoms were calculated geometrically, with C—H = 0.93-0.98 Å, O—H = 0.82 Å, and refined as riding with  $U_{iso}(H) = 1.2U_{eq}(C)$  or  $1.2U_{eq}(C, O)$  for methyl and hydroxy hydrogen atoms.

**Figure 1**

The structure of the title compound, showing the atom-numbering scheme. Displacement ellipsoids are drawn at the 50% probability level. The intramolecular O—H $\cdots$ N hydrogen bond is indicated by a dashed line.

### 1-[(Pyrrolidin-1-yl)(*p*-tolyl)methyl]naphthalen-2-ol

#### Crystal data

$C_{22}H_{23}NO$

$M_r = 317.41$

Monoclinic,  $P2_1/n$

Hall symbol:  $-P 2_1n$

$a = 10.3467 (18) \text{ \AA}$

$b = 16.055 (3) \text{ \AA}$

$c = 11.252 (2) \text{ \AA}$

$\beta = 106.810 (8)^\circ$

$V = 1789.2 (6) \text{ \AA}^3$

$Z = 4$

$F(000) = 680$

$D_x = 1.178 \text{ Mg m}^{-3}$

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

Cell parameters from 3280 reflections

$\theta = 2.3\text{--}27.4^\circ$

$\mu = 0.07 \text{ mm}^{-1}$

$T = 293 \text{ K}$

Prism, colourless

$0.25 \times 0.22 \times 0.20 \text{ mm}$

*Data collection*

Rigaku SCXmini  
diffractometer  
Radiation source: fine-focus sealed tube  
Graphite monochromator  
Detector resolution: 13.6612 pixels mm<sup>-1</sup>  
 $\omega$  scans  
Absorption correction: multi-scan  
(*CrystalClear*; Rigaku, 2005)  
 $T_{\min} = 0.963$ ,  $T_{\max} = 0.989$

18171 measured reflections  
4086 independent reflections  
2547 reflections with  $I > 2\sigma(I)$   
 $R_{\text{int}} = 0.058$   
 $\theta_{\max} = 27.4^\circ$ ,  $\theta_{\min} = 2.3^\circ$   
 $h = -13 \rightarrow 13$   
 $k = -20 \rightarrow 20$   
 $l = -14 \rightarrow 14$

*Refinement*

Refinement on  $F^2$   
Least-squares matrix: full  
 $R[F^2 > 2\sigma(F^2)] = 0.069$   
 $wR(F^2) = 0.190$   
 $S = 1.06$   
4086 reflections  
219 parameters  
0 restraints  
Primary atom site location: structure-invariant  
direct methods

Secondary atom site location: difference Fourier  
map  
Hydrogen site location: inferred from  
neighbouring sites  
H-atom parameters constrained  
 $w = 1/[\sigma^2(F_o^2) + (0.0911P)^2 + 0.1709P]$   
where  $P = (F_o^2 + 2F_c^2)/3$   
 $(\Delta/\sigma)_{\max} = 0.005$   
 $\Delta\rho_{\max} = 0.25 \text{ e } \text{Å}^{-3}$   
 $\Delta\rho_{\min} = -0.20 \text{ e } \text{Å}^{-3}$

*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 ( $\text{Å}^2$ )*

	<i>x</i>	<i>y</i>	<i>z</i>	$U_{\text{iso}}^*/U_{\text{eq}}$
C1	0.2106 (2)	0.17738 (14)	0.94453 (19)	0.0419 (5)
C2	0.2318 (2)	0.10314 (14)	1.0087 (2)	0.0484 (6)
C3	0.3641 (2)	0.07349 (17)	1.0645 (2)	0.0593 (7)
H3	0.3762	0.0225	1.1056	0.071*
C4	0.4730 (2)	0.11799 (18)	1.0592 (2)	0.0603 (7)
H4	0.5591	0.0970	1.0959	0.072*
C5	0.4578 (2)	0.19606 (16)	0.9986 (2)	0.0511 (6)
C6	0.3253 (2)	0.22609 (14)	0.9413 (2)	0.0437 (5)
C7	0.3143 (2)	0.30541 (15)	0.8828 (2)	0.0526 (6)
H7	0.2292	0.3276	0.8455	0.063*
C8	0.4263 (3)	0.34974 (18)	0.8801 (3)	0.0680 (8)
H8	0.4161	0.4014	0.8410	0.082*
C9	0.5550 (3)	0.3189 (2)	0.9349 (3)	0.0713 (8)
H9	0.6302	0.3495	0.9317	0.086*
C10	0.5714 (2)	0.2439 (2)	0.9932 (3)	0.0654 (8)
H10	0.6579	0.2236	1.0299	0.078*

C11	0.0689 (2)	0.20844 (12)	0.87783 (19)	0.0396 (5)
H11	0.0738	0.2388	0.8037	0.048*
C12	0.0091 (2)	0.08850 (15)	0.7372 (2)	0.0512 (6)
H12A	0.0886	0.0544	0.7710	0.061*
H12B	0.0248	0.1249	0.6740	0.061*
C13	-0.1145 (2)	0.03501 (16)	0.6848 (2)	0.0603 (7)
H13A	-0.1071	-0.0171	0.7298	0.072*
H13B	-0.1269	0.0231	0.5977	0.072*
C14	-0.2303 (2)	0.08716 (16)	0.7016 (3)	0.0617 (7)
H14A	-0.2831	0.0555	0.7445	0.074*
H14B	-0.2890	0.1048	0.6218	0.074*
C15	-0.1655 (2)	0.16206 (16)	0.7785 (2)	0.0563 (7)
H15A	-0.1704	0.2107	0.7263	0.068*
H15B	-0.2098	0.1745	0.8415	0.068*
C16	0.01458 (19)	0.26802 (13)	0.95666 (19)	0.0395 (5)
C17	0.0117 (2)	0.24824 (14)	1.0757 (2)	0.0477 (5)
H17	0.0454	0.1972	1.1100	0.057*
C18	-0.0407 (2)	0.30347 (15)	1.1441 (2)	0.0528 (6)
H18	-0.0428	0.2884	1.2233	0.063*
C19	-0.0899 (2)	0.38058 (15)	1.0975 (2)	0.0479 (6)
C20	-0.0865 (2)	0.39997 (14)	0.9788 (2)	0.0493 (6)
H20	-0.1189	0.4514	0.9450	0.059*
C21	-0.0361 (2)	0.34482 (14)	0.9096 (2)	0.0458 (5)
H21	-0.0362	0.3595	0.8296	0.055*
C22	-0.1439 (3)	0.44099 (18)	1.1736 (3)	0.0712 (8)
H22A	-0.1290	0.4193	1.2560	0.107*
H22B	-0.2389	0.4488	1.1358	0.107*
H22C	-0.0982	0.4934	1.1776	0.107*
N1	-0.02391 (16)	0.13698 (11)	0.83579 (16)	0.0414 (4)
O1	0.13049 (17)	0.05359 (11)	1.02090 (17)	0.0641 (5)
H1	0.0585	0.0700	0.9744	0.096*

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

	$U^{11}$	$U^{22}$	$U^{33}$	$U^{12}$	$U^{13}$	$U^{23}$
C1	0.0345 (11)	0.0496 (13)	0.0407 (12)	-0.0007 (9)	0.0094 (9)	-0.0015 (9)
C2	0.0431 (12)	0.0535 (14)	0.0463 (13)	-0.0024 (10)	0.0091 (10)	0.0064 (10)
C3	0.0518 (14)	0.0671 (17)	0.0532 (15)	0.0093 (13)	0.0057 (12)	0.0106 (12)
C4	0.0392 (12)	0.0837 (19)	0.0513 (15)	0.0098 (12)	0.0024 (11)	-0.0049 (13)
C5	0.0368 (11)	0.0684 (16)	0.0465 (13)	-0.0041 (11)	0.0097 (10)	-0.0150 (11)
C6	0.0373 (11)	0.0525 (13)	0.0425 (12)	-0.0048 (9)	0.0135 (9)	-0.0096 (10)
C7	0.0464 (13)	0.0510 (14)	0.0629 (16)	-0.0094 (11)	0.0197 (11)	-0.0057 (11)
C8	0.0629 (17)	0.0617 (17)	0.086 (2)	-0.0203 (13)	0.0317 (15)	-0.0078 (14)
C9	0.0527 (16)	0.085 (2)	0.081 (2)	-0.0297 (15)	0.0271 (15)	-0.0208 (16)
C10	0.0368 (12)	0.094 (2)	0.0635 (17)	-0.0096 (13)	0.0123 (12)	-0.0234 (15)
C11	0.0361 (10)	0.0415 (12)	0.0407 (12)	-0.0038 (9)	0.0103 (9)	0.0045 (9)
C12	0.0431 (12)	0.0561 (15)	0.0543 (14)	-0.0004 (10)	0.0143 (11)	-0.0072 (11)
C13	0.0517 (14)	0.0602 (16)	0.0629 (16)	-0.0037 (12)	0.0070 (12)	-0.0137 (12)

C14	0.0407 (12)	0.0626 (16)	0.0748 (18)	-0.0043 (11)	0.0055 (12)	-0.0118 (13)
C15	0.0343 (12)	0.0621 (16)	0.0671 (16)	0.0008 (10)	0.0062 (11)	-0.0113 (12)
C16	0.0313 (10)	0.0451 (12)	0.0409 (12)	-0.0050 (9)	0.0086 (9)	0.0021 (9)
C17	0.0527 (13)	0.0446 (13)	0.0473 (13)	0.0006 (10)	0.0168 (11)	0.0070 (10)
C18	0.0560 (14)	0.0626 (16)	0.0433 (13)	-0.0062 (12)	0.0198 (11)	0.0010 (11)
C19	0.0335 (11)	0.0544 (14)	0.0543 (15)	-0.0033 (10)	0.0105 (10)	-0.0063 (11)
C20	0.0415 (12)	0.0446 (13)	0.0584 (15)	0.0059 (10)	0.0088 (11)	0.0044 (10)
C21	0.0411 (11)	0.0499 (14)	0.0440 (13)	0.0009 (10)	0.0085 (10)	0.0067 (10)
C22	0.0627 (17)	0.077 (2)	0.0782 (19)	0.0021 (14)	0.0275 (15)	-0.0171 (15)
N1	0.0309 (9)	0.0458 (10)	0.0463 (11)	-0.0024 (7)	0.0093 (8)	-0.0028 (8)
O1	0.0508 (10)	0.0634 (12)	0.0733 (13)	-0.0053 (8)	0.0102 (9)	0.0243 (9)

*Geometric parameters (Å, °)*

C1—C2	1.378 (3)	C12—H12B	0.9700
C1—C6	1.430 (3)	C13—C14	1.518 (3)
C1—C11	1.525 (3)	C13—H13A	0.9700
C2—O1	1.354 (3)	C13—H13B	0.9700
C2—C3	1.413 (3)	C14—C15	1.519 (3)
C3—C4	1.351 (4)	C14—H14A	0.9700
C3—H3	0.9300	C14—H14B	0.9700
C4—C5	1.414 (4)	C15—N1	1.476 (3)
C4—H4	0.9300	C15—H15A	0.9700
C5—C10	1.420 (3)	C15—H15B	0.9700
C5—C6	1.420 (3)	C16—C21	1.384 (3)
C6—C7	1.423 (3)	C16—C17	1.385 (3)
C7—C8	1.367 (3)	C17—C18	1.383 (3)
C7—H7	0.9300	C17—H17	0.9300
C8—C9	1.387 (4)	C18—C19	1.383 (3)
C8—H8	0.9300	C18—H18	0.9300
C9—C10	1.357 (4)	C19—C20	1.381 (3)
C9—H9	0.9300	C19—C22	1.504 (3)
C10—H10	0.9300	C20—C21	1.378 (3)
C11—N1	1.483 (2)	C20—H20	0.9300
C11—C16	1.518 (3)	C21—H21	0.9300
C11—H11	0.9800	C22—H22A	0.9600
C12—N1	1.474 (3)	C22—H22B	0.9600
C12—C13	1.511 (3)	C22—H22C	0.9600
C12—H12A	0.9700	O1—H1	0.8200
C2—C1—C6	118.6 (2)	C14—C13—H13A	110.9
C2—C1—C11	121.71 (19)	C12—C13—H13B	110.9
C6—C1—C11	119.71 (19)	C14—C13—H13B	110.9
O1—C2—C1	123.4 (2)	H13A—C13—H13B	108.9
O1—C2—C3	115.8 (2)	C13—C14—C15	105.88 (19)
C1—C2—C3	120.8 (2)	C13—C14—H14A	110.6
C4—C3—C2	121.0 (2)	C15—C14—H14A	110.6
C4—C3—H3	119.5	C13—C14—H14B	110.6

C2—C3—H3	119.5	C15—C14—H14B	110.6
C3—C4—C5	120.8 (2)	H14A—C14—H14B	108.7
C3—C4—H4	119.6	N1—C15—C14	104.57 (18)
C5—C4—H4	119.6	N1—C15—H15A	110.8
C4—C5—C10	121.5 (2)	C14—C15—H15A	110.8
C4—C5—C6	118.6 (2)	N1—C15—H15B	110.8
C10—C5—C6	120.0 (3)	C14—C15—H15B	110.8
C5—C6—C7	116.8 (2)	H15A—C15—H15B	108.9
C5—C6—C1	120.2 (2)	C21—C16—C17	117.6 (2)
C7—C6—C1	123.0 (2)	C21—C16—C11	120.06 (19)
C8—C7—C6	121.4 (2)	C17—C16—C11	122.37 (19)
C8—C7—H7	119.3	C18—C17—C16	120.8 (2)
C6—C7—H7	119.3	C18—C17—H17	119.6
C7—C8—C9	121.0 (3)	C16—C17—H17	119.6
C7—C8—H8	119.5	C19—C18—C17	121.6 (2)
C9—C8—H8	119.5	C19—C18—H18	119.2
C10—C9—C8	120.1 (2)	C17—C18—H18	119.2
C10—C9—H9	120.0	C20—C19—C18	117.3 (2)
C8—C9—H9	120.0	C20—C19—C22	121.5 (2)
C9—C10—C5	120.7 (3)	C18—C19—C22	121.2 (2)
C9—C10—H10	119.6	C21—C20—C19	121.4 (2)
C5—C10—H10	119.6	C21—C20—H20	119.3
N1—C11—C16	111.04 (16)	C19—C20—H20	119.3
N1—C11—C1	110.22 (16)	C20—C21—C16	121.3 (2)
C16—C11—C1	112.58 (17)	C20—C21—H21	119.4
N1—C11—H11	107.6	C16—C21—H21	119.4
C16—C11—H11	107.6	C19—C22—H22A	109.5
C1—C11—H11	107.6	C19—C22—H22B	109.5
N1—C12—C13	103.87 (18)	H22A—C22—H22B	109.5
N1—C12—H12A	111.0	C19—C22—H22C	109.5
C13—C12—H12A	111.0	H22A—C22—H22C	109.5
N1—C12—H12B	111.0	H22B—C22—H22C	109.5
C13—C12—H12B	111.0	C12—N1—C15	103.41 (17)
H12A—C12—H12B	109.0	C12—N1—C11	112.27 (16)
C12—C13—C14	104.3 (2)	C15—N1—C11	113.43 (17)
C12—C13—H13A	110.9	C2—O1—H1	109.5

*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>
O1—H1...N1	0.82	1.88	2.600 (3)	145
C18—H18...Cg1 <sup>i</sup>	0.93	2.66	3.588 (8)	173

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