

5,15-Bis(3,5-di-*tert*-butylphenyl)-10,20-bis(phenylethynyl)porphyrin

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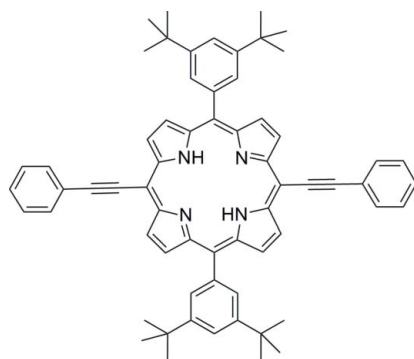
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Key indicators: single-crystal X-ray study; $T = 213$ K; mean $\sigma(\text{C}-\text{C}) = 0.008$ Å; disorder in main residue; R factor = 0.107; wR factor = 0.315; data-to-parameter ratio = 11.4.

In the centrosymmetric title compound, $\text{C}_{64}\text{H}_{62}\text{N}_4$, the two phenylethynyl groups lie at diagonal *meso* positions. The 24-membered porphyrin has in-plane distortion with respect to the mean plane of the macrocycle and two intra-ring bifurcated $\text{N}-\text{H}\cdots(\text{N},\text{N})$ hydrogen bonds occur. The dihedral angles between the phenyl rings in the phenylethynyl group and the 3,5-bis(*tert*-butyl)phenyl group with respect to the mean plane of the porphyrin are 17.2 (2) and 59.2 (3)°. The *tert*-butyl groups are disordered over two sets of sites in a 0.661 (13):0.339 (13) ratio.

Related literature

For background to porphyrin structures and electronic properties, see: Anderson *et al.* (1994, 1998); Fujita *et al.* (1995); Henari *et al.* (1997); Huuskonen *et al.* (1998); LeCours *et al.* (1996); Screen *et al.* (2002); Seo *et al.* (2008); Silvers & Tulinsky (1967).



Experimental

Crystal data

$\text{C}_{64}\text{H}_{62}\text{N}_4$
 $M_r = 887.18$

Triclinic, $P\bar{1}$
 $a = 9.9598$ (19) Å

$b = 10.496$ (2) Å
 $c = 13.925$ (3) Å
 $\alpha = 86.236$ (4)°
 $\beta = 80.266$ (4)°
 $\gamma = 82.765$ (4)°
 $V = 1421.8$ (5) Å³

$Z = 1$
Mo $K\alpha$ radiation
 $\mu = 0.06$ mm⁻¹
 $T = 213$ K
 $0.45 \times 0.10 \times 0.05$ mm

Data collection

Bruker SMART CCD
diffractometer
Absorption correction: multi-scan
(*SADABS*; Bruker, 2003)
 $T_{\min} = 0.974$, $T_{\max} = 0.997$

6144 measured reflections
4136 independent reflections
2068 reflections with $I > 2\sigma(I)$
 $R_{\text{int}} = 0.101$
 $\theta_{\text{max}} = 23.5^\circ$

Refinement

$R[F^2 > 2\sigma(F^2)] = 0.107$
 $wR(F^2) = 0.315$
 $S = 0.99$
4136 reflections
363 parameters

193 restraints
H-atom parameters constrained
 $\Delta\rho_{\text{max}} = 1.13$ e Å⁻³
 $\Delta\rho_{\text{min}} = -0.30$ e Å⁻³

Table 1

Hydrogen-bond geometry (Å, °).

$D-\text{H}\cdots A$	$D-\text{H}$	$\text{H}\cdots A$	$D\cdots A$	$D-\text{H}\cdots A$
$\text{N1}-\text{H1A}\cdots\text{N2}$	0.87	2.44	2.972 (6)	120
$\text{N1}-\text{H1A}\cdots\text{N2}^i$	0.87	2.35	2.891 (5)	121

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

Data collection: *SMART* (Bruker, 2003); cell refinement: *SAINT* (Bruker, 2003); data reduction: *SAINT*; program(s) used to solve structure: *SHELXS97* (Sheldrick, 2008); program(s) used to refine structure: *SHELXL97* (Sheldrick, 2008); molecular graphics: *SHELXTL* (Sheldrick, 2008); software used to prepare material for publication: *SHELXTL* and *PLATON* (Spek, 2009).

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Supplementary data and figures for this paper are available from the IUCr electronic archives (Reference: HB5134).

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supplementary materials

Acta Cryst. (2009). E65, o3004–o3005 [doi:10.1107/S1600536809045954]

5,15-Bis(3,5-di-*tert*-butylphenyl)-10,20-bis(phenylethynyl)porphyrin

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Comment

The electronic and steric tunings through surrogating various substituents on the *meso*- and beta-carbons play crucial role in the synthesis of diverse porphyrin systems. In this respect, *meso*-ethynyl porphyrins have attracted the attention due to their wide utilization for the development of conjugated electronic (LeCours *et al.*, 1996; Anderson *et al.*, 1994; Henari *et al.*, 1997; Screen *et al.*, 2002) and light harvesting materials. As an extension of our research on the conjugated photoelectronic materials (Seo *et al.*, 2008), the title compound (I) was prepared and its crystal structure determined.

The molecular structure of C₆₄H₆₂N₄ (I) is shown in Figure 1. The structure shows symmetric molecular system due to presence of inversion center (Ci) in the core of porphyrin macrocycle. On assuming all four nitrogen atoms of all pyrrolic groups in a plane, it was observed that the *meso*-carbons were deviated with ± 0.072 Å from the least-squares plane and find analogy with earlier reported compound 5,15-bis(3,5-di-*tert*-butylphenyl)-10,20-bis(trimethylsilylethynyl)porphyrin (Huuskonen *et al.*, 1998). The structural analysis of the porphyrin macrocycle reveals that the plane of phenyl rings in the phenylethynyl groups is slightly twisted with the dihedral angle of 17.17° with respect to the least-squares plane of the porphyrin, in contrast the plane of aryl rings associated with 3,5-bis(*tert*-butyl)phenyl groups (59.19°) (Figure 2). The dihedral angle associated with phenylethynyl groups (17.17°) is also much smaller with respect to phenyl planes slanting (61–63°) in tetraphenylporphyrin (Silvers *et al.*, 1967). The comparison of this result with the tetraphenylporphyrin, it was observed that on increasing the conjugation with phenyl groups leads to release of steric strain. The distances between the nitrogen atoms (N1—N2 = 2.972, N1—N2' = 2.891, N1—N1' = 4.167, N2—N2' = 4.126 Å) involve in the formation of basal parallelogram based core along with the C—C (triple bond) (1.165 Å) bond lengths and C(*meso*)-C(α)-C(β) angles (178.07°) involving ethynyl groups show the analogy with previous reports (Fujita *et al.*, 1995; Huuskonen *et al.*, 1998). In addition, the distances involving the diagonal *meso*-carbon atoms (C5—C5' = 6.815, C10—C10' = 6.994 Å) also differ to each other and correlate with the previous work (Huuskonen *et al.*, 1998). The shortest intermolecular distances for the pi-pi and pi-H interactions are not observed less than 6.085 and 3.84 Å, respectively and imply the steric strain between aryl groups of adjacent porphyrin prevent the strong interactions in between adjacent molecular system (Anderson *et al.*, 1998).

Experimental

The title compound was prepared from the corresponding dipyrromethane and phenylpropargyl aldehyde as follows. BF₃.OEt₂ (25 ML, 0.2 mmol) was added to a solution of *meso*-(3,5-di-*tert*-butylphenyl) dipyrromethane (0.669 g, 2.0 mmol) and phenylpropargyl aldehyde (245 ML, 2.0 mmol) in dry CH₂Cl₂ (200 ml). The reaction mixture was stirred for 10 min at room temperature.

2,3-Dichloro-5,6-dicyano-*p*-benzoquinone (340 mg, 0.15 mmol) was added and further stirred for 30 min. After evaporation of solvent to dryness, the title compound was separated by column chromatography (SiO₂, CH₂Cl₂:*n*-hexane = 1:4). Recrystallization from a CH₂Cl₂/CH₃CN solution afforded purple crystalline solid. Yield: 124 mg (14%). ¹H NMR (200 MHz, CDCl₃): Δ 9.70 (d, 4H), 8.89 (d, 4H), 8.06 (s, 4H), 8.00 (t, 2H), 7.82 (s, 2H), 1.55 (s, 36H), -1.92 (br, 2H). UV-vis (CH₂Cl₂): λ_{\max} (log E) 443 (6.04), 554 (4.38), 594(4.92), 623 (4.25), 681 (4.61) nm.

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Purple needles of (I) were grown by slow diffusion of CH₃CN to a CH₂Cl₂ solution of the title compound.

Refinement

The carbon atoms C26—C28 and C30—C32 and their attached H atoms are disordered over two sets of sites in a 66.1:33.1 ratio with total site occupancy of 1.00 for each one of them. The contributions of the mostly disordered solvent molecules were removed from the diffraction data using the SQUEEZE routine of *PLATON* software (Spek, 2009), and then final refinements were carried out. All the non-hydrogen atoms were refined anisotropically, and hydrogen atoms were placed in their geometrically ideal positions.

Figures

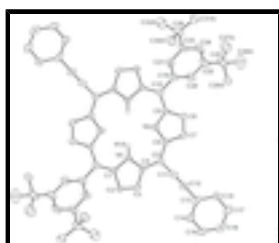


Fig. 1. The molecular structure of (I) showing 30% probability displacement ellipsoids. Hydrogen atoms (except to N—H) have been omitted for the clarity.



Fig. 2. A view of molecular structure of (I) showing the tilted planes of the different aryl groups.

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Crystal data

C₆₄H₆₂N₄

M_r = 887.18

Triclinic, *P* $\bar{1}$

Hall symbol: -P 1

a = 9.9598 (19) Å

b = 10.496 (2) Å

c = 13.925 (3) Å

α = 86.236 (4)°

β = 80.266 (4)°

γ = 82.765 (4)°

V = 1421.8 (5) Å³

Z = 1

*F*₀₀₀ = 474

D_x = 1.036 Mg m⁻³

Mo *K*α radiation, λ = 0.71073 Å

Cell parameters from 4136 reflections

θ = 2.0–23.5°

μ = 0.06 mm⁻¹

T = 213 K

Needle, purple

0.45 × 0.10 × 0.05 mm

Data collection

Bruker SMART CCD
diffractometer

Radiation source: fine-focus sealed tube

Monochromator: graphite

T = 213 K

4136 independent reflections

2068 reflections with *I* > 2σ(*I*)

*R*_{int} = 0.101

θ_{max} = 23.5°

ω scans $\theta_{\min} = 2.0^\circ$
 Absorption correction: multi-scan $h = -8 \rightarrow 11$
 (SADABS; Bruker, 2003)
 $T_{\min} = 0.974$, $T_{\max} = 0.997$ $k = -11 \rightarrow 11$
 6144 measured reflections $l = -11 \rightarrow 15$

Refinement

Refinement on F^2 Secondary atom site location: difference Fourier map
 Least-squares matrix: full Hydrogen site location: inferred from neighbouring sites
 $R[F^2 > 2\sigma(F^2)] = 0.107$ H-atom parameters constrained
 $wR(F^2) = 0.315$ $w = 1/[\sigma^2(F_o^2) + (0.1906P)^2]$
 where $P = (F_o^2 + 2F_c^2)/3$
 $S = 0.99$ $(\Delta/\sigma)_{\max} < 0.001$
 4136 reflections $\Delta\rho_{\max} = 1.13 \text{ e } \text{\AA}^{-3}$
 363 parameters $\Delta\rho_{\min} = -0.30 \text{ e } \text{\AA}^{-3}$
 193 restraints Extinction correction: none
 Primary atom site location: structure-invariant direct methods

Fractional atomic coordinates and isotropic or equivalent isotropic displacement parameters (\AA^2)

	<i>x</i>	<i>y</i>	<i>z</i>	$U_{\text{iso}}^*/U_{\text{eq}}$	Occ. (<1)
N1	0.4997 (4)	0.8413 (3)	0.5981 (3)	0.0445 (11)	
H1A	0.4940	0.9116	0.5616	0.053*	
N2	0.3805 (4)	0.9196 (3)	0.4184 (3)	0.0481 (11)	
C1	0.5695 (5)	0.8204 (4)	0.6753 (3)	0.0429 (13)	
C2	0.5537 (6)	0.6929 (5)	0.7151 (4)	0.0529 (14)	
H2	0.5907	0.6530	0.7689	0.063*	
C3	0.4771 (6)	0.6413 (5)	0.6622 (4)	0.0600 (16)	
H3	0.4511	0.5578	0.6718	0.072*	
C4	0.4409 (5)	0.7342 (4)	0.5880 (4)	0.0449 (13)	
C5	0.3646 (5)	0.7155 (4)	0.5160 (4)	0.0482 (13)	
C6	0.3377 (5)	0.8019 (5)	0.4380 (4)	0.0468 (13)	
C7	0.2576 (6)	0.7764 (5)	0.3646 (4)	0.0577 (15)	
H7	0.2154	0.7020	0.3608	0.069*	
C8	0.2563 (6)	0.8818 (5)	0.3030 (4)	0.0564 (15)	
H8	0.2137	0.8929	0.2472	0.068*	
C9	0.3299 (5)	0.9725 (4)	0.3367 (3)	0.0409 (12)	
C10	0.3527 (5)	1.0946 (5)	0.2919 (3)	0.0470 (13)	
C11	0.3101 (6)	0.5928 (5)	0.5201 (4)	0.0557 (14)	
C12	0.2674 (6)	0.4937 (5)	0.5260 (4)	0.0606 (16)	
C13	0.2183 (6)	0.3674 (5)	0.5331 (4)	0.0584 (15)	
C14	0.2721 (6)	0.2699 (5)	0.5915 (4)	0.0605 (15)	
H14	0.3410	0.2849	0.6265	0.073*	
C15	0.1152 (6)	0.3433 (5)	0.4811 (4)	0.0626 (16)	
H15	0.0781	0.4072	0.4393	0.075*	

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C16	0.2265 (7)	0.1512 (5)	0.5990 (5)	0.0712 (18)	
H16	0.2655	0.0850	0.6381	0.085*	
C17	0.0704 (7)	0.2243 (6)	0.4930 (4)	0.0760 (19)	
H17	-0.0007	0.2083	0.4605	0.091*	
C18	0.1249 (8)	0.1287 (6)	0.5501 (5)	0.0757 (19)	
H18	0.0929	0.0475	0.5558	0.091*	
C19	0.2853 (6)	1.1365 (5)	0.2057 (4)	0.0511 (14)	
C20	0.1439 (6)	1.1470 (5)	0.2137 (4)	0.0636 (16)	
H20	0.0914	1.1283	0.2745	0.076*	
C21	0.3627 (6)	1.1657 (4)	0.1160 (4)	0.0541 (14)	
H21	0.4586	1.1596	0.1104	0.065*	
C22	0.0761 (6)	1.1849 (6)	0.1337 (4)	0.0704 (17)	
C23	0.3015 (6)	1.2037 (5)	0.0341 (4)	0.0612 (15)	
C24	0.1597 (6)	1.2114 (6)	0.0466 (4)	0.0714 (18)	
H24	0.1172	1.2366	-0.0081	0.086*	
C25	-0.0782 (7)	1.2010 (8)	0.1463 (5)	0.090 (2)	
C26A	-0.1491 (10)	1.1396 (11)	0.2354 (9)	0.077 (3)	0.661 (13)
H26A	-0.1029	1.0541	0.2459	0.115*	0.661 (13)
H26B	-0.2433	1.1337	0.2283	0.115*	0.661 (13)
H26C	-0.1479	1.1907	0.2908	0.115*	0.661 (13)
C27A	-0.1397 (14)	1.3240 (12)	0.1232 (13)	0.118 (5)	0.661 (13)
H27A	-0.2385	1.3240	0.1324	0.178*	0.661 (13)
H27B	-0.1063	1.3481	0.0558	0.178*	0.661 (13)
H27C	-0.1171	1.3852	0.1655	0.178*	0.661 (13)
C28A	-0.1218 (15)	1.1113 (16)	0.0638 (12)	0.147 (6)	0.661 (13)
H28A	-0.0845	1.0225	0.0739	0.221*	0.661 (13)
H28B	-0.0852	1.1424	-0.0013	0.221*	0.661 (13)
H28C	-0.2209	1.1172	0.0711	0.221*	0.661 (13)
C26B	-0.125 (4)	1.110 (3)	0.176 (3)	0.156 (15)	0.339 (13)
H26D	-0.2235	1.1246	0.1789	0.234*	0.339 (13)
H26E	-0.1041	1.0889	0.2416	0.234*	0.339 (13)
H26F	-0.0867	1.0400	0.1344	0.234*	0.339 (13)
C27B	-0.121 (3)	1.3362 (18)	0.2041 (19)	0.109 (9)	0.339 (13)
H27D	-0.0753	1.4043	0.1668	0.164*	0.339 (13)
H27E	-0.0933	1.3249	0.2679	0.164*	0.339 (13)
H27F	-0.2195	1.3590	0.2115	0.164*	0.339 (13)
C28B	-0.117 (3)	1.263 (3)	0.048 (2)	0.153 (12)	0.339 (13)
H28D	-0.0526	1.3227	0.0203	0.229*	0.339 (13)
H28E	-0.2091	1.3088	0.0598	0.229*	0.339 (13)
H28F	-0.1147	1.1962	0.0025	0.229*	0.339 (13)
C29	0.3836 (7)	1.2398 (6)	-0.0623 (4)	0.0763 (18)	
C30A	0.495 (6)	1.131 (5)	-0.084 (4)	0.094 (9)	0.121 (8)
H30A	0.5050	1.0805	-0.0243	0.141*	0.121 (8)
H30B	0.5801	1.1652	-0.1102	0.141*	0.121 (8)
H30C	0.4709	1.0776	-0.1311	0.141*	0.121 (8)
C31A	0.294 (5)	1.274 (6)	-0.138 (4)	0.089 (10)	0.121 (8)
H31A	0.3509	1.2927	-0.2001	0.134*	0.121 (8)
H31B	0.2299	1.3483	-0.1193	0.134*	0.121 (8)
H31C	0.2446	1.2018	-0.1446	0.134*	0.121 (8)

C32A	0.445 (7)	1.358 (5)	-0.035 (5)	0.110 (10)	0.121 (8)
H32A	0.5169	1.3299	0.0039	0.165*	0.121 (8)
H32B	0.3740	1.4150	0.0026	0.165*	0.121 (8)
H32C	0.4841	1.4041	-0.0938	0.165*	0.121 (8)
C30B	0.3606 (9)	1.1477 (8)	-0.1410 (5)	0.092 (3)	0.879 (8)
H30D	0.4134	1.1703	-0.2035	0.138*	0.879 (8)
H30E	0.2640	1.1563	-0.1463	0.138*	0.879 (8)
H30F	0.3903	1.0595	-0.1217	0.138*	0.879 (8)
C31B	0.3428 (10)	1.3783 (8)	-0.0936 (6)	0.109 (3)	0.879 (8)
H31D	0.3979	1.3990	-0.1557	0.163*	0.879 (8)
H31E	0.3579	1.4346	-0.0449	0.163*	0.879 (8)
H31F	0.2466	1.3899	-0.1003	0.163*	0.879 (8)
C32B	0.5416 (8)	1.2160 (9)	-0.0616 (5)	0.086 (2)	0.879 (8)
H32D	0.5902	1.2415	-0.1249	0.130*	0.879 (8)
H32E	0.5680	1.1255	-0.0475	0.130*	0.879 (8)
H32F	0.5645	1.2664	-0.0120	0.130*	0.879 (8)

Atomic displacement parameters (\AA^2)

	U^{11}	U^{22}	U^{33}	U^{12}	U^{13}	U^{23}
N1	0.048 (3)	0.031 (2)	0.050 (3)	-0.0016 (18)	-0.004 (2)	0.0123 (18)
N2	0.053 (3)	0.039 (2)	0.050 (3)	-0.0027 (19)	-0.002 (2)	0.0033 (19)
C1	0.043 (3)	0.033 (3)	0.046 (3)	0.006 (2)	-0.001 (2)	0.012 (2)
C2	0.057 (4)	0.043 (3)	0.059 (3)	-0.008 (3)	-0.014 (3)	0.014 (3)
C3	0.058 (4)	0.047 (3)	0.071 (4)	-0.009 (3)	-0.006 (3)	0.026 (3)
C4	0.050 (3)	0.032 (3)	0.048 (3)	-0.005 (2)	0.005 (3)	0.006 (2)
C5	0.049 (3)	0.041 (3)	0.051 (3)	-0.007 (2)	-0.002 (3)	0.006 (2)
C6	0.035 (3)	0.050 (3)	0.051 (3)	-0.001 (2)	0.001 (2)	-0.002 (2)
C7	0.061 (4)	0.055 (3)	0.059 (4)	-0.021 (3)	-0.006 (3)	0.004 (3)
C8	0.061 (4)	0.066 (4)	0.043 (3)	-0.014 (3)	-0.006 (3)	0.000 (3)
C9	0.035 (3)	0.039 (3)	0.046 (3)	0.000 (2)	-0.004 (2)	0.001 (2)
C10	0.043 (3)	0.050 (3)	0.043 (3)	0.003 (2)	0.000 (2)	-0.001 (2)
C11	0.059 (4)	0.046 (3)	0.060 (4)	-0.014 (3)	-0.004 (3)	0.011 (3)
C12	0.071 (4)	0.046 (3)	0.062 (4)	-0.012 (3)	-0.001 (3)	0.008 (3)
C13	0.056 (4)	0.043 (3)	0.071 (4)	-0.009 (3)	0.007 (3)	-0.008 (3)
C14	0.055 (4)	0.049 (3)	0.075 (4)	-0.007 (3)	-0.004 (3)	-0.001 (3)
C15	0.071 (4)	0.054 (3)	0.059 (4)	-0.006 (3)	-0.004 (3)	0.000 (3)
C16	0.065 (4)	0.044 (3)	0.093 (5)	0.001 (3)	0.009 (4)	0.011 (3)
C17	0.090 (5)	0.084 (5)	0.058 (4)	-0.038 (4)	0.000 (3)	-0.014 (3)
C18	0.080 (5)	0.055 (4)	0.085 (5)	-0.020 (3)	0.016 (4)	-0.008 (4)
C19	0.053 (4)	0.052 (3)	0.046 (3)	0.003 (3)	-0.008 (3)	-0.005 (2)
C20	0.049 (4)	0.076 (4)	0.057 (4)	0.011 (3)	0.004 (3)	-0.001 (3)
C21	0.054 (4)	0.053 (3)	0.049 (3)	0.007 (3)	-0.002 (3)	-0.003 (2)
C22	0.057 (4)	0.086 (4)	0.061 (4)	0.020 (3)	-0.010 (3)	0.000 (3)
C23	0.062 (4)	0.062 (3)	0.057 (4)	0.011 (3)	-0.018 (3)	0.004 (3)
C24	0.063 (5)	0.095 (4)	0.050 (4)	0.024 (3)	-0.015 (3)	0.000 (3)
C25	0.054 (5)	0.121 (6)	0.086 (5)	0.029 (4)	-0.020 (4)	0.009 (4)
C26A	0.038 (5)	0.093 (6)	0.096 (7)	-0.002 (4)	-0.014 (5)	0.018 (5)

supplementary materials

C27A	0.097 (8)	0.117 (8)	0.135 (9)	0.002 (6)	-0.020 (7)	0.011 (7)
C28A	0.109 (9)	0.171 (10)	0.167 (10)	-0.024 (7)	-0.031 (7)	-0.014 (8)
C26B	0.149 (17)	0.162 (17)	0.157 (17)	-0.013 (10)	-0.031 (10)	0.002 (10)
C27B	0.099 (11)	0.111 (11)	0.114 (12)	-0.006 (8)	-0.015 (9)	0.000 (9)
C28B	0.147 (15)	0.165 (15)	0.148 (14)	-0.006 (9)	-0.037 (9)	-0.006 (9)
C29	0.081 (4)	0.089 (4)	0.048 (3)	0.016 (3)	-0.002 (3)	0.004 (3)
C30A	0.089 (12)	0.096 (11)	0.092 (13)	0.003 (8)	-0.010 (9)	0.002 (9)
C31A	0.087 (12)	0.095 (14)	0.085 (12)	-0.007 (8)	-0.015 (9)	0.005 (9)
C32A	0.112 (15)	0.112 (12)	0.106 (15)	-0.011 (9)	-0.021 (10)	-0.012 (10)
C30B	0.083 (5)	0.133 (6)	0.052 (4)	0.007 (4)	0.000 (3)	-0.010 (4)
C31B	0.114 (6)	0.108 (5)	0.089 (5)	-0.002 (4)	0.002 (5)	0.039 (4)
C32B	0.083 (5)	0.122 (6)	0.045 (4)	0.001 (4)	0.003 (3)	0.009 (4)

Geometric parameters (Å, °)

N1—C4	1.357 (6)	C29—C30A	1.49 (4)
N1—C1	1.367 (6)	C29—C31A	1.49 (4)
N2—C6	1.354 (6)	C29—C31B	1.515 (9)
N2—C9	1.378 (6)	C29—C30B	1.570 (10)
C1—C10 ⁱ	1.397 (7)	C29—C32B	1.563 (10)
C1—C2	1.431 (6)	C29—C32A	1.55 (5)
C2—C3	1.329 (7)	N1—H1A	0.869 (3)
C3—C4	1.433 (7)	C2—H2	0.941 (6)
C4—C5	1.393 (7)	C3—H3	0.940 (5)
C5—C6	1.407 (7)	C8—H8	0.940 (6)
C5—C11	1.454 (7)	C14—H14	0.940 (6)
C6—C7	1.455 (7)	C15—H15	0.940 (5)
C7—C8	1.355 (7)	C16—H16	0.940 (6)
C8—C9	1.420 (7)	C17—H17	0.940 (7)
C9—C10	1.415 (6)	C18—H18	0.941 (7)
C10—C1 ⁱ	1.397 (7)	C20—H20	0.940 (5)
C10—C19	1.488 (7)	C21—H21	0.939 (6)
C11—C12	1.164 (7)	C24—H24	0.940 (6)
C12—C13	1.462 (7)	C26A—H26A	0.97 (1)
C13—C14	1.377 (8)	C26A—H26B	0.97 (1)
C13—C15	1.409 (8)	C26A—H26C	0.97 (1)
C14—C16	1.371 (7)	C27A—H27A	0.97 (1)
C15—C17	1.369 (8)	C27A—H27B	0.97 (2)
C16—C18	1.364 (9)	C27A—H27C	0.97 (2)
C17—C18	1.355 (9)	C28A—H28A	0.97 (2)
C19—C20	1.385 (7)	C28A—H28B	0.97 (2)
C19—C21	1.390 (7)	C28A—H28C	0.97 (1)
C20—C22	1.408 (8)	C30A—H30A	0.97 (5)
C21—C23	1.395 (7)	C30A—H30B	0.97 (6)
C22—C24	1.383 (8)	C30A—H30C	0.97 (6)
C22—C25	1.506 (9)	C31A—H31A	0.97 (5)
C23—C24	1.386 (8)	C31A—H31B	0.96 (5)
C23—C29	1.502 (8)	C31A—H31C	0.97 (6)
C25—C26B	1.14 (3)	C32A—H32A	0.97 (7)

C25—C27A	1.400 (13)	C32A—H32B	0.97 (6)
C25—C26A	1.473 (12)	C32A—H32C	0.97 (6)
C25—C28B	1.57 (3)	N1—H1A	0.869 (3)
C25—C27B	1.651 (16)	C2—H2	0.941 (6)
C25—C28A	1.676 (16)		
C4—N1—C1	108.7 (4)	C28B—C25—C28A	59.0 (12)
C6—N2—C9	107.9 (4)	C27B—C25—C28A	143.9 (12)
N1—C1—C10 ⁱ	126.7 (4)	C30A—C29—C31A	116 (3)
N1—C1—C2	107.9 (5)	C30A—C29—C23	106 (2)
C10 ⁱ —C1—C2	125.4 (5)	C31A—C29—C23	112 (2)
C3—C2—C1	107.4 (5)	C30A—C29—C31B	142 (2)
C2—C3—C4	108.6 (5)	C31A—C29—C31B	58 (2)
N1—C4—C5	126.4 (4)	C23—C29—C31B	111.3 (5)
N1—C4—C3	107.4 (5)	C30A—C29—C30B	65 (2)
C5—C4—C3	126.1 (5)	C31A—C29—C30B	54 (2)
C4—C5—C6	126.8 (4)	C23—C29—C30B	109.0 (6)
C4—C5—C11	116.8 (4)	C31B—C29—C30B	110.0 (6)
C6—C5—C11	116.4 (5)	C30A—C29—C32B	44 (2)
N2—C6—C5	126.7 (5)	C31A—C29—C32B	136 (2)
N2—C6—C7	109.1 (4)	C23—C29—C32B	112.4 (5)
C5—C6—C7	124.2 (5)	C31B—C29—C32B	110.1 (7)
C8—C7—C6	105.9 (5)	C30B—C29—C32B	103.9 (6)
C7—C8—C9	108.7 (5)	C30A—C29—C32A	110 (3)
N2—C9—C8	108.4 (4)	C31A—C29—C32A	112 (3)
N2—C9—C10	125.5 (5)	C23—C29—C32A	101 (2)
C8—C9—C10	126.0 (5)	C31B—C29—C32A	54 (2)
C1 ⁱ —C10—C9	124.6 (5)	C30B—C29—C32A	150 (2)
C1 ⁱ —C10—C19	117.9 (4)	C32B—C29—C32A	66 (2)
C9—C10—C19	117.4 (5)	C25—C26A—H26A	110 (1)
C12—C11—C5	178.0 (6)	C25—C26A—H26B	109 (1)
C11—C12—C13	178.2 (6)	C25—C26A—H26C	109 (1)
C14—C13—C15	118.9 (5)	H26A—C26A—H26B	109 (1)
C14—C13—C12	120.2 (6)	H26A—C26A—H26C	109 (1)
C15—C13—C12	120.9 (5)	H26B—C26A—H26C	109 (1)
C16—C14—C13	120.7 (6)	C25—C27A—H27A	109 (1)
C17—C15—C13	118.2 (6)	C25—C27A—H27B	110 (1)
C18—C16—C14	120.3 (6)	C25—C27A—H27C	109 (1)
C18—C17—C15	122.4 (7)	H27A—C27A—H27B	109 (1)
C17—C18—C16	119.4 (6)	H27A—C27A—H27C	109 (1)
C20—C19—C21	118.9 (5)	H27B—C27A—H27C	109 (1)
C20—C19—C10	120.3 (5)	C25—C28A—H28A	109 (1)
C21—C19—C10	120.8 (5)	C25—C28A—H28B	109 (1)
C19—C20—C22	122.2 (5)	C25—C28A—H28C	109 (1)
C23—C21—C19	121.6 (5)	H28A—C28A—H28B	109 (2)
C24—C22—C20	115.8 (6)	H28A—C28A—H28C	110 (2)
C24—C22—C25	123.6 (6)	H28B—C28A—H28C	109 (2)
C20—C22—C25	120.6 (5)	C23—C29—C30A	106 (2)
C24—C23—C21	116.7 (5)	C23—C29—C31A	111 (2)

supplementary materials

C24—C23—C29	121.1 (5)	C23—C29—C32A	101 (2)
C21—C23—C29	122.1 (6)	C30A—C29—C31A	116 (3)
C22—C24—C23	124.8 (6)	C30A—C29—C32A	110 (3)
C26B—C25—C27A	131 (2)	C31A—C29—C32A	112 (3)
C26B—C25—C26A	36.7 (19)	C29—C30A—H30A	109 (5)
C27A—C25—C26A	113.9 (9)	C29—C30A—H30B	109 (5)
C26B—C25—C22	113 (2)	C29—C30A—H30C	109 (5)
C27A—C25—C22	115.8 (9)	H30A—C30A—H30B	110 (5)
C26A—C25—C22	115.9 (6)	H30A—C30A—H30C	110 (5)
C26B—C25—C28B	117 (2)	H30B—C30A—H30C	110 (5)
C27A—C25—C28B	48.9 (12)	C29—C31A—H31A	109 (5)
C26A—C25—C28B	136.7 (14)	C29—C31A—H31B	110 (5)
C22—C25—C28B	106.5 (13)	C29—C31A—H31C	109 (5)
C26B—C25—C27B	120 (2)	H31A—C31A—H31B	110 (5)
C27A—C25—C27B	45.0 (9)	H31A—C31A—H31C	109 (5)
C26A—C25—C27B	85.5 (11)	H31B—C31A—H31C	110 (5)
C22—C25—C27B	102.9 (11)	C29—C32A—H32A	110 (5)
C28B—C25—C27B	93.6 (14)	C29—C32A—H32B	110 (5)
C26B—C25—C28A	64 (2)	C29—C32A—H32C	110 (5)
C27A—C25—C28A	102.6 (10)	H32A—C32A—H32B	109 (6)
C26A—C25—C28A	98.7 (9)	H32A—C32A—H32C	109 (6)
C22—C25—C28A	107.1 (7)	H32B—C32A—H32C	109 (6)

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

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

$D-H\cdots A$	$D-H$	$H\cdots A$	$D\cdots A$	$D-H\cdots A$
N1—H1A \cdots N2	0.87	2.44	2.972 (6)	120
N1—H1A \cdots N2 ⁱ	0.87	2.35	2.891 (5)	121

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

Fig. 1

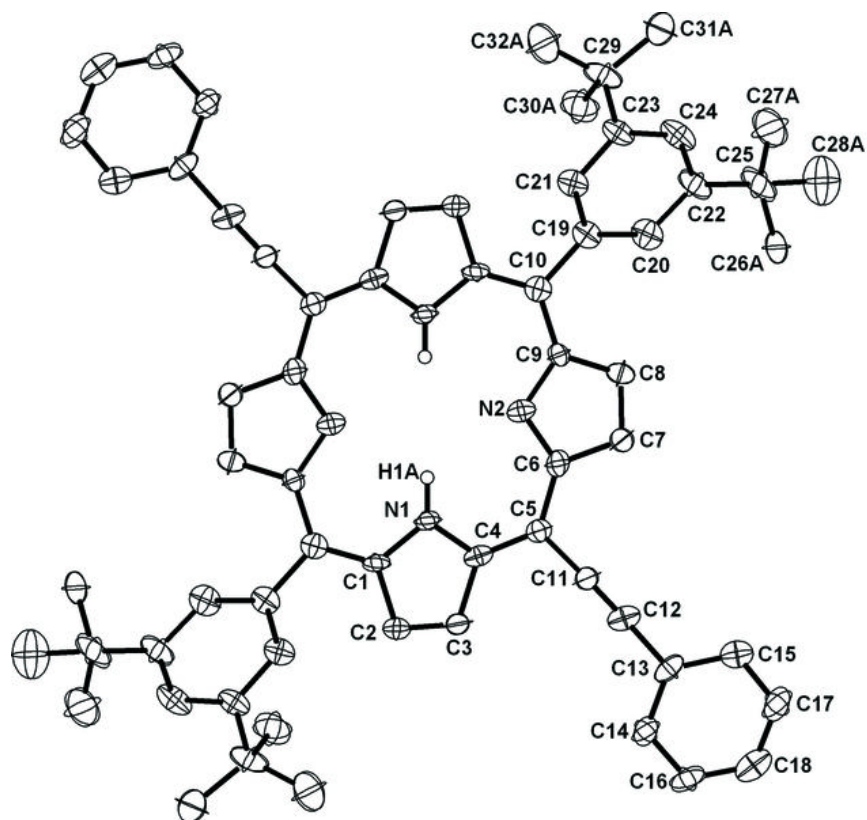


Fig. 2

