

Di- μ -chlorido-bis{[2-(morpholino-methyl)phenyl- κ^2 C¹,N]palladium(II)}

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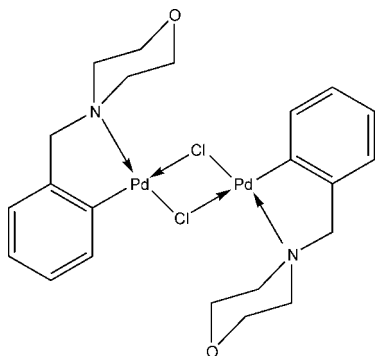
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Key indicators: single-crystal X-ray study; $T = 297$ K; mean $\sigma(\text{C}-\text{C}) = 0.005$ Å; R factor = 0.029; wR factor = 0.059; data-to-parameter ratio = 17.2.

The title compound, $[\text{Pd}_2(\text{C}_{11}\text{H}_{14}\text{NO})_2\text{Cl}_2]$, has a dimeric structure with Cl atoms bridging the two Pd atoms, one half of the molecule being generated by symmetry due to the crystallographic inversion centre located in the middle of the perfectly planar Pd_2Cl_2 ring. The five-membered ring adopts an envelope conformation, while the morpholino group has a chair conformation. The geometry around the metal centres is distorted square-planar, as a result of a strong intramolecular $\text{N}\rightarrow\text{Pd}$ coordination *trans* to a $\text{Pd}-\text{Cl}$ bond. In the crystal structure, the dimeric structure is strengthened by intermolecular $\text{C}-\text{H}\cdots\text{Cl}$ hydrogen bonds. $\text{C}-\text{H}\cdots\text{C}_{\text{phenyl}}$ interactions link the dimers into a columnar supramolecular array along the a axis; the dimers are further connected by $\text{C}-\text{H}\cdots\text{Ph}$ interactions into a three-dimensional supramolecular arrangement.

Related literature

For related literature, see: Copolovici *et al.* (2007, 2008); Crispini *et al.* (1992); Fuchita *et al.* (1999); Mahalakshmi *et al.* (2003); Mentis *et al.* (1997, 2004, 2005); Phadnis *et al.* (2002, 2003); Emsley (1994); IUPAC (1979).



Experimental

Crystal data

$[\text{Pd}_2(\text{C}_{11}\text{H}_{14}\text{NO})_2\text{Cl}_2]$
 $M_r = 636.20$
 Monoclinic, $P2_1/c$
 $a = 8.1234$ (6) Å
 $b = 16.4437$ (13) Å
 $c = 8.8298$ (7) Å
 $\beta = 101.9570$ (10)°

$V = 1153.88$ (15) Å³
 $Z = 2$
 Mo $K\alpha$ radiation
 $\mu = 1.81$ mm⁻¹
 $T = 297$ (2) K
 $0.22 \times 0.17 \times 0.17$ mm

Data collection

Bruker SMART APEX CCD
 area-detector diffractometer
 Absorption correction: multi-scan
 (*SADABS*; Bruker, 2000)
 $T_{\text{min}} = 0.677$, $T_{\text{max}} = 0.733$

9118 measured reflections
 2354 independent reflections
 2245 reflections with $I > 2\sigma(I)$
 $R_{\text{int}} = 0.026$

Refinement

$R[F^2 > 2\sigma(F^2)] = 0.028$
 $wR(F^2) = 0.059$
 $S = 1.15$
 2354 reflections

137 parameters
 H-atom parameters constrained
 $\Delta\rho_{\text{max}} = 0.49$ e Å⁻³
 $\Delta\rho_{\text{min}} = -0.43$ e Å⁻³

Table 1

Selected geometric parameters (Å, °).

Cl1—Pd1	2.3232 (8)	N1—Pd1	2.118 (2)
Cl1—Pd1 ⁱ	2.4815 (8)	Cl1—Pd1	1.971 (3)
Cl1—Pd1—Cl1 ⁱ	82.66 (3)	Cl1—Pd1—N1	81.69 (11)
N1—Pd1—Cl1	174.34 (7)	Pd1—Cl1—Pd1 ⁱ	97.34 (3)
N1—Pd1—Cl1 ⁱ	102.76 (6)	C7—N1—Pd1	104.01 (17)
C1—Pd1—Cl1	92.88 (9)	C8—N1—Pd1	117.35 (18)
C1—Pd1—Cl1 ⁱ	175.53 (9)	C11—N1—Pd1	111.92 (17)

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

Table 2

Hydrogen-bond geometry (Å, °).

C_g is the centroid of the C1—C6 ring.

$D-\text{H}\cdots A$	$D-\text{H}$	$\text{H}\cdots A$	$D\cdots A$	$D-\text{H}\cdots A$
C9—H9B ⁱ ⋯Cl1 ⁱ	0.97	2.49	3.370 (3)	150
C7—H7B ⁱⁱ ⋯C1 ⁱⁱⁱ	0.97	2.67	3.588 (4)	159
C11—H11A ⁱⁱⁱ ⋯C _g ⁱⁱⁱ	0.97	2.93	3.829 (3)	154

Symmetry codes: (i) $-x + 1, -y + 1, -z$; (ii) $-x, -y + 1, -z$; (iii) $x, -y + \frac{3}{2}, z + \frac{1}{2}$.

Data collection: *SMART* (Bruker, 2000); cell refinement: *SAINT-Plus* (Bruker, 2001); data reduction: *SAINT-Plus*; program(s) used to solve structure: *SHELXS97* (Sheldrick, 2008); program(s) used to refine structure: *SHELXL97* (Sheldrick, 2008); molecular graphics: *DIAMOND* (Brandenburg & Putz, 2006); software used to prepare material for publication: *pubCIF* (Westrip, 2008).

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Supplementary data and figures for this paper are available from the IUCr electronic archives (Reference: HK2489).

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supporting information

Acta Cryst. (2008). E64, m1057–m1058 [doi:10.1107/S1600536808022575]

Di- μ -chlorido-bis[2-(morpholinomethyl)phenyl- κ^2C^1,N]palladium(II)}

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S1. Comment

There are scarce data reported upon the reactivity of triorganoantimony(III) compounds toward palladium(II) chloride or corresponding complexes. The compounds of the type $[PdCl_2(SbR_3)_2]$, $R = iPr$ (Phadnis *et al.*, 2002), *ortho*-tolyl (Mentes & Fawcett, 2005), were obtained by reaction of $[PdCl_2(COD)]$, COD = cycloocta-1,5-diene, with the corresponding stibines. The reaction of $[PdCl_2(PhCN)_2]$ with SbR_3 ($R = 2$ -thienyl) resulted in a black insoluble powder (possible Pd metal) due to a decomposition process (Mahalakshmi *et al.*, 2003), whilst the reaction of $[PdCl_2(PhCN)_2]$ with tri(allyl)-stibine and tris(2-methylallyl)stibine afforded the transmetalation products $[Pd_2(\mu-Cl)_2R_2]$, $R =$ allyl, 2-methylallyl, and the fate of the stibine residue is unknown (Phadnis *et al.*, 2003). Reaction of $SbPh_3$ with $[PdCl_2(COD)]$ or $PdCl_2$ afforded *trans*- $[PdCl(\sigma-Ph)(SbPh_3)_2]$ due to the easy cleavage of $Sb-C$ bond in $SbPh_3$, whilst reaction of Na_2PdCl_4 with $SbPh_3$ gave *cis*- $[PdCl_2(SbPh_3)_2]$ contaminated with small amount of chloro σ -phenyl complex (Mentes *et al.*, 1997). Related to our interest on the synthesis and chemical properties of organoantimony(III) compounds, we performed the reaction between $Sb[C_6H_4CH_2\{N(CH_2CH_2O)_2\}_2]_3$ and $[PdCl_2(MeCN)_2]$ in acetonitrile/chloroform mixture, at room temperature.

The title compound has a dimeric structure, with two palladacycles bridged through the chlorine atoms, resulting in a perfectly planar Pd_2Cl_2 core. One half of the molecule is generated by symmetry owing the crystallographic inversion centre located in the middle of the Pd_2Cl_2 ring. The cycle is distorted from the ideal square as reflected by the differences in the $Pd-Cl$ bond lengths and the $Cl1-Pd1-Cl1^i$ and $Pd1-Cl1-Pd1^i$ bond angles (Table 1) [symmetry code: (i) = 1 - x, 1 - y, -z].

The N atom from the pendant arm coordinates the metal centre resulting in a square planar $(C,N)PdCl_2$ core, in which the distortion is mainly due to the PdC_3N ring constraint. The two equivalent organic ligands from the dimer are in a *trans* arrangement with respect to the $Pd \cdots Pd$ axis (Fig. 1). The $Pd1-Cl1$ [1.971 (2) Å] bond is smaller than the sum of the corresponding covalent radii, its magnitude being similar to those found for related compounds for which partially $Pd-C$ multiple bond was assumed (Crispini *et al.*, 1992, Fuchita *et al.*, 1999, Mentes *et al.*, 2004).

An almost ideal *chair* conformation was observed for the morpholinyl groups with torsion angles [$C8-N1-C11-C10 = -52.6$ (3)° and $C10-O1-C9-C8 = 58.4$ (3)°] similar with those found in 4-benzylmorpholin-4-ium chloride (Copolovici *et al.*, 2007) and in tris[2-(morpholin-4-ylmethyl)phenyl- κ^2C^1,N]antimony(III) (Copolovici *et al.*, 2008).

As a result of the intramolecular coordination of the N1 atom from the organic ligand to Pd1 atom, a nonplanar five-membered ring is formed, with nitrogen atom lying out of the $Pd1/C1/C2/C7$ best plane [0.666 (2) Å]. The dihedral angle between the $Pd1/N1/C7$ and $Pd1/C1/C2/C7$ planes is 39.2 (1)°. This induces planar chirality (with the aromatic ring and the nitrogen atom as chiral plane and pilot atom, respectively, IUPAC, 1979). In the crystal of the title compound, the dimer contains both R_N and S_N^i isomers.

In the crystal structure, the dimer is strengthened by hydrogen-bond type interactions (Table 2) involving the Cl from one molecular unit and the methylenic proton of the morpholinyl group from the other molecular unit from the same dimer [the sum of van der Waals radii of the corresponding atoms $\Sigma r_{vdw}(Cl,H) = 3.00$ Å; Emsley, 1994] (Fig. 2).

Intermolecular $C-H\cdots C_{\text{phenyl}}$ interactions (Table 2) between a methylene hydrogen from the pendant arm and a carbon atom from the aromatic ring of another dimer connect the molecular units in a columnar arrangement along the a axis. Furthermore, these arrays are interlinked by intermolecular $C-H\cdots Ph$ interactions (Table 2) in a three-dimensional supramolecular arrangement in the crystal structure (Fig.3).

S2. Experimental

For the preparation of the title compound, $PdCl_2$ (0.1 g, 0.56 mmol) in acetonitrile (40 ml) was refluxed for 3 h, and then allowed to cool to room temperature. A solution of $Sb[C_6H_4CH_2\{N(CH_2CH_2)_2O\}-2]_3$ (0.369 g, 0.56 mmol) in $CHCl_3$ (35 ml) was added and the mixture was stirred at room temperature for 30 h, under an N_2 atmosphere. The reaction mixture was filtered off and the solvents were removed under vacuum. The yellow oil obtained was triturated with hexane (2 x 20 ml) and then washed with petroleum ether (2 x 10 ml) to give a yellow solid. Yellow crystals suitable for X-ray diffraction studies were obtained by slow diffusion of hexane into a solution of CH_2Cl_2 of the title compound (1:1 v/v ratio) (yield: 0.217 g, 47%). Anal. Found: C, 41.38; H, 4.63; N, 4.73. Calc. for $C_{22}H_{28}Cl_2N_2O_2Pd_2$ (636.22): C, 41.53; H, 4.44; N, 4.40%. FT-IR (KBr, cm^{-1}): C-H stretch aromatic: 3038, 3035, 2958, 2891; 2857, 1436, 1114, 1069, 868, 745.

S3. Refinement

H atoms were positioned geometrically, with C-H = 0.93 and 0.97 Å for aromatic and methylene H, respectively, and constrained to ride on their parent atoms with $U_{\text{iso}}(H) = 1.2U_{\text{eq}}(C)$,

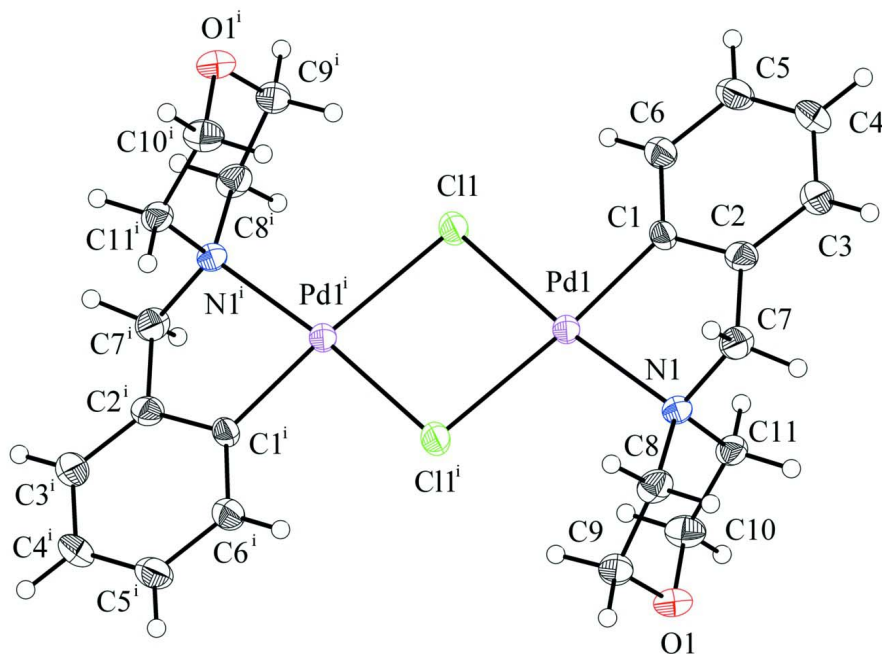
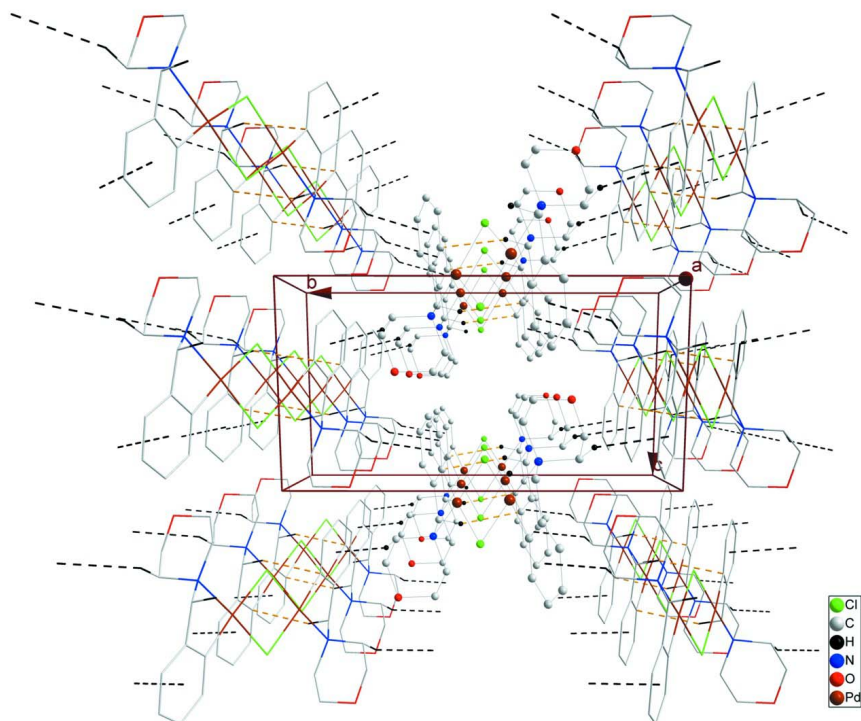


Figure 1

The molecular structure of the title molecule, with the atom-numbering scheme. Displacement ellipsoids are drawn at the 30% probability level [symmetry code: (i) $1 - x, 1 - y, -z$].

**Figure 3**

A packing diagram of the title compound, showing the supramolecular arrangement.

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Crystal data

[Pd₂(C₁₁H₁₄NO)₂Cl₂]

$M_r = 636.20$

Monoclinic, $P2_1/c$

Hall symbol: -P 2ybc

$a = 8.1234$ (6) Å

$b = 16.4437$ (13) Å

$c = 8.8298$ (7) Å

$\beta = 101.957$ (1)°

$V = 1153.88$ (15) Å³

$Z = 2$

$F(000) = 632$

$D_x = 1.831$ Mg m⁻³

Mo $K\alpha$ radiation, $\lambda = 0.71073$ Å

Cell parameters from 4334 reflections

$\theta = 2.5$ – 27.3 °

$\mu = 1.81$ mm⁻¹

$T = 297$ K

Block, yellow

$0.22 \times 0.17 \times 0.17$ mm

Data collection

Bruker SMART APEX CCD area-detector
diffractometer

Radiation source: fine-focus sealed tube

Graphite monochromator

φ and ω scans

Absorption correction: multi-scan

(*SADABS*; Bruker, 2000)

$T_{\min} = 0.677$, $T_{\max} = 0.733$

9118 measured reflections

2354 independent reflections

2245 reflections with $I > 2\sigma(I)$

$R_{\text{int}} = 0.026$

$\theta_{\text{max}} = 26.4$ °, $\theta_{\text{min}} = 2.5$ °

$h = -10 \rightarrow 10$

$k = -20 \rightarrow 20$

$l = -11 \rightarrow 11$

Refinement

Refinement on F^2
 Least-squares matrix: full
 $R[F^2 > 2\sigma(F^2)] = 0.028$
 $wR(F^2) = 0.059$
 $S = 1.15$
 2354 reflections
 137 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
 $w = 1/[\sigma^2(F_o^2) + (0.0166P)^2 + 1.0911P]$
 where $P = (F_o^2 + 2F_c^2)/3$
 $(\Delta/\sigma)_{\max} = 0.002$
 $\Delta\rho_{\max} = 0.49 \text{ e } \text{Å}^{-3}$
 $\Delta\rho_{\min} = -0.43 \text{ e } \text{Å}^{-3}$

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 $F^2 > 2\sigma(F^2)$ 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 (Å^2)

	<i>x</i>	<i>y</i>	<i>z</i>	$U_{\text{iso}}^*/U_{\text{eq}}$
Pd1	0.32057 (3)	0.559622 (13)	0.02382 (2)	0.03252 (8)
Cl1	0.41182 (11)	0.49501 (6)	-0.17788 (9)	0.0591 (3)
O1	0.4523 (3)	0.69670 (15)	0.4417 (3)	0.0524 (6)
N1	0.2135 (3)	0.61896 (14)	0.1932 (3)	0.0311 (5)
C1	0.1119 (3)	0.59843 (17)	-0.1126 (3)	0.0332 (6)
C2	-0.0122 (4)	0.62200 (19)	-0.0349 (4)	0.0382 (7)
C3	-0.1640 (4)	0.6534 (2)	-0.1153 (4)	0.0564 (10)
H3	-0.2460	0.6700	-0.0624	0.068*
C4	-0.1921 (4)	0.6598 (2)	-0.2752 (4)	0.0578 (10)
H4	-0.2932	0.6811	-0.3296	0.069*
C5	-0.0721 (4)	0.6349 (2)	-0.3534 (4)	0.0494 (8)
H5	-0.0930	0.6382	-0.4608	0.059*
C6	0.0798 (4)	0.60484 (19)	-0.2732 (4)	0.0401 (7)
H6	0.1614	0.5888	-0.3270	0.048*
C7	0.0286 (4)	0.6089 (2)	0.1360 (4)	0.0426 (7)
H7A	-0.0318	0.6480	0.1864	0.051*
H7B	-0.0055	0.5547	0.1601	0.051*
C8	0.2571 (4)	0.5867 (2)	0.3546 (3)	0.0437 (8)
H8A	0.2506	0.5278	0.3510	0.052*
H8B	0.1743	0.6059	0.4112	0.052*
C9	0.4285 (4)	0.6111 (2)	0.4406 (4)	0.0493 (8)
H9A	0.4448	0.5917	0.5464	0.059*
H9B	0.5125	0.5854	0.3929	0.059*
C10	0.4272 (4)	0.7249 (2)	0.2865 (4)	0.0497 (8)
H10A	0.5065	0.6983	0.2344	0.060*

H10B	0.4479	0.7830	0.2862	0.060*
C11	0.2517 (4)	0.70759 (18)	0.2014 (3)	0.0378 (7)
H11A	0.1730	0.7351	0.2529	0.045*
H11B	0.2365	0.7292	0.0972	0.045*

Atomic displacement parameters (Å²)

	U^{11}	U^{22}	U^{33}	U^{12}	U^{13}	U^{23}
Pd1	0.03461 (14)	0.03664 (14)	0.02611 (12)	0.00850 (9)	0.00582 (9)	0.00045 (9)
C11	0.0574 (5)	0.0891 (7)	0.0284 (4)	0.0394 (5)	0.0034 (4)	-0.0061 (4)
O1	0.0565 (14)	0.0596 (15)	0.0356 (12)	0.0015 (12)	-0.0028 (10)	-0.0127 (11)
N1	0.0336 (12)	0.0330 (12)	0.0270 (12)	-0.0006 (10)	0.0071 (10)	-0.0009 (10)
C1	0.0313 (14)	0.0282 (14)	0.0368 (16)	0.0048 (12)	-0.0006 (12)	0.0000 (12)
C2	0.0324 (15)	0.0417 (17)	0.0391 (17)	-0.0050 (13)	0.0039 (13)	-0.0066 (13)
C3	0.0319 (17)	0.077 (3)	0.057 (2)	0.0086 (17)	0.0022 (16)	-0.0190 (19)
C4	0.0371 (18)	0.069 (2)	0.057 (2)	0.0090 (17)	-0.0122 (16)	-0.0084 (19)
C5	0.0468 (19)	0.056 (2)	0.0398 (18)	0.0014 (16)	-0.0038 (15)	0.0025 (16)
C6	0.0390 (16)	0.0432 (17)	0.0365 (16)	0.0032 (14)	0.0043 (13)	0.0025 (14)
C7	0.0344 (16)	0.0509 (19)	0.0446 (18)	-0.0057 (14)	0.0131 (14)	-0.0087 (15)
C8	0.060 (2)	0.0444 (18)	0.0291 (16)	0.0036 (16)	0.0144 (15)	0.0038 (13)
C9	0.059 (2)	0.059 (2)	0.0274 (16)	0.0164 (17)	0.0032 (15)	0.0011 (15)
C10	0.050 (2)	0.050 (2)	0.0455 (19)	-0.0111 (16)	0.0020 (15)	-0.0035 (16)
C11	0.0464 (17)	0.0309 (15)	0.0343 (16)	0.0022 (13)	0.0042 (13)	-0.0015 (12)

Geometric parameters (Å, °)

Pd1—C11 ⁱ	2.4815 (8)	C7—H7A	0.9700
C11—Pd1	2.3232 (8)	C7—H7B	0.9700
C11—Pd1 ⁱ	2.4815 (8)	C8—N1	1.493 (4)
N1—Pd1	2.118 (2)	C8—C9	1.496 (5)
C1—C2	1.387 (4)	C8—H8A	0.9700
C1—C6	1.392 (4)	C8—H8B	0.9700
C1—Pd1	1.971 (3)	C9—O1	1.421 (4)
C2—C3	1.389 (4)	C9—H9A	0.9700
C2—C7	1.492 (4)	C9—H9B	0.9700
C3—C4	1.387 (5)	C10—O1	1.421 (4)
C3—H3	0.9300	C10—C11	1.495 (4)
C4—C5	1.369 (5)	C10—H10A	0.9700
C4—H4	0.9300	C10—H10B	0.9700
C5—C6	1.381 (4)	C11—N1	1.489 (4)
C5—H5	0.9300	C11—H11A	0.9700
C6—H6	0.9300	C11—H11B	0.9700
C7—N1	1.492 (4)		
C11—Pd1—C11 ⁱ	82.66 (3)	C1—C6—H6	119.7
N1—Pd1—C11	174.34 (7)	N1—C7—C2	108.9 (2)
N1—Pd1—C11 ⁱ	102.76 (6)	N1—C7—H7A	109.9
C1—Pd1—C11	92.88 (9)	C2—C7—H7A	109.9

C1—Pd1—C11 ⁱ	175.53 (9)	N1—C7—H7B	109.9
C1—Pd1—N1	81.69 (11)	C2—C7—H7B	109.9
Pd1—C11—Pd1 ⁱ	97.34 (3)	H7A—C7—H7B	108.3
C10—O1—C9	108.9 (2)	N1—C8—C9	113.6 (3)
C7—N1—C8	107.9 (2)	N1—C8—H8A	108.8
C7—N1—Pd1	104.01 (17)	C9—C8—H8A	108.8
C8—N1—Pd1	117.35 (18)	N1—C8—H8B	108.8
C11—N1—Pd1	111.92 (17)	C9—C8—H8B	108.8
C11—N1—C7	108.0 (2)	H8A—C8—H8B	107.7
C11—N1—C8	107.2 (2)	O1—C9—C8	112.4 (3)
C2—C1—C6	118.8 (3)	O1—C9—H9A	109.1
C2—C1—Pd1	114.2 (2)	C8—C9—H9A	109.1
C6—C1—Pd1	127.1 (2)	O1—C9—H9B	109.1
C1—C2—C3	120.7 (3)	C8—C9—H9B	109.1
C1—C2—C7	115.4 (3)	H9A—C9—H9B	107.9
C3—C2—C7	123.9 (3)	O1—C10—C11	110.8 (3)
C4—C3—C2	119.4 (3)	O1—C10—H10A	109.5
C4—C3—H3	120.3	C11—C10—H10A	109.5
C2—C3—H3	120.3	O1—C10—H10B	109.5
C5—C4—C3	120.4 (3)	C11—C10—H10B	109.5
C5—C4—H4	119.8	H10A—C10—H10B	108.1
C3—C4—H4	119.8	N1—C11—C10	112.2 (3)
C4—C5—C6	120.2 (3)	N1—C11—H11A	109.2
C4—C5—H5	119.9	C10—C11—H11A	109.2
C6—C5—H5	119.9	N1—C11—H11B	109.2
C5—C6—C1	120.5 (3)	C10—C11—H11B	109.2
C5—C6—H6	119.7	H11A—C11—H11B	107.9
Pd1 ⁱ —C11—Pd1—C1	-179.66 (9)	C1—C2—C7—N1	32.1 (4)
Pd1 ⁱ —C11—Pd1—C11 ⁱ	0.0	C3—C2—C7—N1	-149.6 (3)
C11—N1—Pd1—C1	-84.42 (19)	N1—C8—C9—O1	54.4 (4)
C7—N1—Pd1—C1	31.98 (19)	O1—C10—C11—N1	-61.0 (4)
C8—N1—Pd1—C1	151.0 (2)	C10—C11—N1—C7	168.6 (3)
C11—N1—Pd1—C11 ⁱ	96.05 (18)	C10—C11—N1—C8	52.6 (3)
C7—N1—Pd1—C11 ⁱ	-147.55 (17)	C10—C11—N1—Pd1	-77.4 (3)
C8—N1—Pd1—C11 ⁱ	-28.5 (2)	C2—C7—N1—C11	77.7 (3)
C6—C1—C2—C3	-1.6 (5)	C2—C7—N1—C8	-166.7 (3)
Pd1—C1—C2—C3	177.7 (3)	C2—C7—N1—Pd1	-41.4 (3)
C6—C1—C2—C7	176.8 (3)	C9—C8—N1—C11	-49.5 (3)
Pd1—C1—C2—C7	-3.9 (3)	C9—C8—N1—C7	-165.6 (3)
C1—C2—C3—C4	1.1 (5)	C9—C8—N1—Pd1	77.4 (3)
C7—C2—C3—C4	-177.1 (3)	C11—C10—O1—C9	61.6 (4)
C2—C3—C4—C5	0.4 (6)	C8—C9—O1—C10	-58.5 (4)
C3—C4—C5—C6	-1.4 (6)	C2—C1—Pd1—N1	-16.4 (2)
C4—C5—C6—C1	0.9 (5)	C6—C1—Pd1—N1	162.9 (3)

C2—C1—C6—C5	0.6 (5)	C2—C1—Pd1—C11	162.0 (2)
Pd1—C1—C6—C5	-178.6 (2)	C6—C1—Pd1—C11	-18.7 (3)

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

Hydrogen-bond geometry (Å, °)

<i>D—H...A</i>	<i>D—H</i>	<i>H...A</i>	<i>D...A</i>	<i>D—H...A</i>
C9—H9B...C11 ⁱ	0.97	2.49	3.370 (3)	150
C7—H7B...C1 ⁱⁱ	0.97	2.67	3.588 (4)	159
C11—H11A...Cg ⁱⁱⁱ	0.97	2.93	3.829 (3)	154

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