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Dichlorido(6,6′-dimethyl-2,2′-bipyridine-κ^2N,N')cobalt(II)

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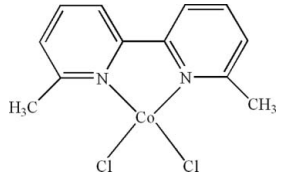
Key indicators: single-crystal X-ray study; T = 298 K; mean σ(C–C) = 0.008 Å; R factor = 0.064; wR factor = 0.226; data-to-parameter ratio = 23.4.

In the title compound, $[CoCl_2(C_{12}H_{12}N_2)]$, the Co^{II} atom is four-coordinated in a distorted tetrahedral geometry by two N atoms from a 6,6′-dimethyl-2,2′-bipyridine ligand and two terminal Cl atoms. Intermolecular C—H···Cl hydrogen bonds and π–π stacking interactions between the pyridine rings [centroid–centroid distances = 3.788 (1) and 3.957 (1) Å] are present in the crystal structure.

Related literature

For related structures, see: Akbarzadeh Torbati *et al.* (2010); Alizadeh *et al.* (2010); Alizadeh, Kalateh, Ebadi *et al.* (2009); Alizadeh, Kalateh, Khoshtarkib *et al.* (2009); Alizadeh, Khoshtarkib *et al.* (2009); Baker *et al.* (1988); Itoh *et al.* (2005); Kou *et al.* (2008); Onggo *et al.* (2005).



Experimental

Crystal data

$[CoCl_2(C_{12}H_{12}N_2)]$
M_r = 314.07
Monoclinic, $P2_1/c$
a = 7.6292 (14) Å
b = 9.8034 (14) Å
c = 17.980 (4) Å
β = 93.990 (15)°
V = 1341.5 (4) Å^3
Z = 4
Mo $K\alpha$ radiation
μ = 1.66 mm^{-1}
T = 298 K
0.50 × 0.19 × 0.13 mm

Data collection

Bruker APEX CCD diffractometer
Absorption correction: multi-scan (*SADABS*; Bruker, 2001)
T_{min} = 0.690, T_{max} = 0.810
10258 measured reflections
3609 independent reflections
2642 reflections with $I > 2\sigma(I)$
R_{int} = 0.100

Refinement

$R[F^2 > 2\sigma(F^2)]$ = 0.064
$wR(F^2)$ = 0.226
S = 1.13
3609 reflections
154 parameters
H-atom parameters constrained
$\Delta\rho_{max}$ = 0.47 e Å^{-3}
$\Delta\rho_{min}$ = −0.92 e Å^{-3}

Table 1
Selected bond lengths (Å).

Co1—N1	2.042 (3)	Co1—Cl1	2.2193 (13)
Co1—N2	2.053 (3)	Co1—Cl2	2.2269 (13)

Table 2
Hydrogen-bond geometry (Å, °).

D—H···A	D—H	H···A	D···A	D—H···A
C5—H5···Cl1^{i}	0.93	2.82	3.565 (7)	138

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

Data collection: *SMART* (Bruker, 2007); cell refinement: *SAINT* (Bruker, 2007); data reduction: *SAINT*; program(s) used to solve structure: *SHELXTL* (Sheldrick, 2008); program(s) used to refine structure: *SHELXTL*; molecular graphics: *ORTEP-3* (Farrugia, 1997); software used to prepare material for publication: *WinGX* (Farrugia, 1999).

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Supplementary data and figures for this paper are available from the IUCr electronic archives (Reference: HY2353).

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supplementary materials

Acta Cryst. (2010). E**66**, m1284 [**doi:10.1107/S1600536810036846**]

Dichlorido(6,6'-dimethyl-2,2'-bipyridine-κ^2N,N')cobalt(II)

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Comment

6,6'-Dimethyl-2,2'-bipyridine (6,6'-dmbipy) is a good bidentate ligand, and numerous complexes with 6,6'-dmbipy have been prepared, such as that of zinc (Alizadeh, Kalateh, Ebadi *et al.*, 2009; Alizadeh, Kalateh, Khoshtarkib *et al.*, 2009; Alizadeh, Khoshtarkib *et al.*, 2009), copper (Itoh *et al.*, 2005), nickel (Kou *et al.*, 2008), cadmium (Alizadeh *et al.*, 2010) and ruthenium (Onggo *et al.*, 2005). We report here the synthesis and crystal structure of the title compound.

In the title compound (Fig. 1), the Co^{II} atom is four-coordinated in a distorted tetrahedral geometry by two N atoms from a 6,6'-dmbipy ligand and two terminal Cl atoms. The Co—N and Co—Cl bond lengths and angles (Table 1) are within normal range as observed in $[Co(6,6'\text{-dmbpy})Cl_2].1/2(C_6H_6)$ (Baker *et al.*, 1988) and $[Co(dmphen)Cl_2]$ (dmphen = 2,9-dimethyl-1,10-phenanthroline) (Akbarzadeh Torbati *et al.*, 2010) .

In the crystal structure, intermolecular C—H···Cl hydrogen bonds (Table 2) and π–π contacts (Fig. 2) between the pyridine rings, $Cg1\cdots Cg2^{i}$ and $Cg1\cdots Cg2^{ii}$ [symmetry codes: (i) 1-x, 1-y, 1-z; (ii) -x, 1-y, 1-z. *Cg*1 and *Cg*2 are the centroids of the N1, C2–C6 ring and N2, C7–C11 ring], stabilize the structure, with centroid–centroid distances of 3.788 (1) and 3.957 (1) Å.

Experimental

For the preparation of the title compound, a solution of 6,6'-dmbipy (0.25 g, 1.34 mmol) in methanol (15 ml) was added to a solution of $CoCl_2.6H_2O$ (0.37 g, 1.34 mmol) in acetonitrile (15 ml) and the resulting blue solution was stirred for 15 min at 313 K. This solution was left to evaporate slowly at room temperature. After one week, blue block crystals of the title compound were isolated (yield: 0.32 g, 76%).

Refinement

All H atoms were positioned geometrically and refined as riding atoms, with C—H = 0.93 (aromatic) and 0.96 (methyl) Å and with $U_{iso}(H) = 1.2U_{eq}(C)$.

Figures

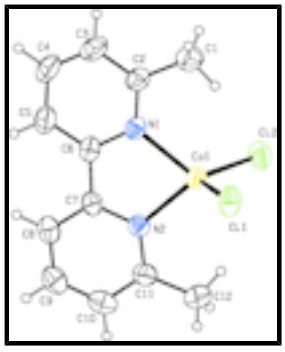

Fig. 1. The molecular structure of the title compound. Displacement ellipsoids are drawn at the 30% probability level.

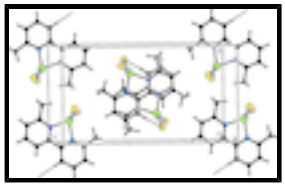

Fig. 2. Crystal packing diagram for the title compound. Dashed lines denote hydrogen bonds.

Dichlorido(6,6'-dimethyl-2,2'-bipyridine-κ^2N,N')cobalt(II)

Crystal data

$[CoCl_2(C_{12}H_{12}N_2)]$	$F(000) = 636$
$M_r = 314.07$	$D_x = 1.555$ Mg m^{-3}
Monoclinic, $P2_1/c$	Mo $K\alpha$ radiation, $\lambda = 0.71073$ Å
Hall symbol: -P 2ybc	Cell parameters from 1009 reflections
$a = 7.6292$ (14) Å	$\theta = 2.3$–$29.3°$
$b = 9.8034$ (14) Å	$\mu = 1.66$ mm^{-1}
$c = 17.980$ (4) Å	$T = 298$ K
$\beta = 93.990$ (15)°	Block, blue
$V = 1341.5$ (4) Å^3	0.50 × 0.19 × 0.13 mm
$Z = 4$	

Data collection

Bruker APEX CCD diffractometer	3609 independent reflections
Radiation source: fine-focus sealed tube	2642 reflections with $I > 2\sigma(I)$
graphite	$R_{int} = 0.100$
φ and ω scans	$\theta_{max} = 29.3°$, $\theta_{min} = 2.3°$
Absorption correction: multi-scan (*SADABS*; Bruker, 2001)	$h = -10 \rightarrow 10$
$T_{min} = 0.690$, $T_{max} = 0.810$	$k = -13 \rightarrow 11$
10258 measured reflections	$l = -24 \rightarrow 24$

Refinement

Refinement on F^2	Primary atom site location: structure-invariant direct methods
Least-squares matrix: full	Secondary atom site location: difference Fourier map
$R[F^2 > 2\sigma(F^2)] = 0.064$	Hydrogen site location: inferred from neighbouring sites
$wR(F^2) = 0.226$	H-atom parameters constrained
$S = 1.13$	$w = 1/[\sigma^2(F_o^2) + (0.1251P)^2 + 0.225P]$ where $P = (F_o^2 + 2F_c^2)/3$
3609 reflections	$(\Delta/\sigma)_{max} < 0.001$
154 parameters	$\Delta\rho_{max} = 0.47$ e Å^{-3}
0 restraints	$\Delta\rho_{min} = -0.92$ e Å^{-3}

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

	x	*y*	*z*	U_{iso}*/U_{eq}
C1	0.3248 (9)	0.4581 (6)	0.2579 (3)	0.0900 (16)
H1A	0.4369	0.4668	0.2373	0.108*
H1B	0.2338	0.4866	0.2217	0.108*
H1C	0.3061	0.3646	0.2712	0.108*
C2	0.3209 (6)	0.5449 (5)	0.3254 (3)	0.0654 (10)
C3	0.3451 (8)	0.6844 (6)	0.3227 (4)	0.0832 (15)
H3	0.3656	0.7264	0.2777	0.100*
C4	0.3390 (8)	0.7594 (5)	0.3849 (4)	0.0857 (16)
H4	0.3592	0.8529	0.3835	0.103*
C5	0.3022 (7)	0.6964 (5)	0.4517 (4)	0.0778 (14)
H5	0.2945	0.7475	0.4950	0.093*
C6	0.2773 (5)	0.5568 (4)	0.4525 (2)	0.0572 (9)
C7	0.2356 (5)	0.4790 (4)	0.5193 (2)	0.0544 (8)
C8	0.2233 (6)	0.5387 (6)	0.5874 (3)	0.0719 (12)
H8	0.2462	0.6312	0.5940	0.086*
C9	0.1762 (7)	0.4593 (7)	0.6459 (3)	0.0807 (14)
H9	0.1637	0.4985	0.6924	0.097*
C10	0.1481 (7)	0.3233 (7)	0.6357 (3)	0.0765 (13)
H10	0.1156	0.2693	0.6749	0.092*
C11	0.1681 (7)	0.2654 (5)	0.5660 (3)	0.0661 (10)
C12	0.1486 (11)	0.1176 (6)	0.5507 (4)	0.100 (2)
H12A	0.0319	0.0892	0.5603	0.120*
H12B	0.2324	0.0678	0.5824	0.120*
H12C	0.1688	0.1000	0.4995	0.120*
N1	0.2875 (4)	0.4836 (3)	0.38883 (19)	0.0537 (7)
N2	0.2096 (4)	0.3440 (4)	0.50901 (18)	0.0541 (7)
Co1	0.24226 (7)	0.28007 (5)	0.40212 (3)	0.0549 (2)
Cl1	0.47387 (17)	0.14706 (13)	0.38988 (8)	0.0811 (4)
Cl2	0.01041 (17)	0.19582 (14)	0.33528 (8)	0.0769 (4)

Atomic displacement parameters (Å^2)

	U^{11}	U^{22}	U^{33}	U^{12}	U^{13}	U^{23}
C1	0.124 (4)	0.079 (3)	0.071 (3)	−0.003 (3)	0.029 (3)	0.007 (3)
C2	0.068 (2)	0.058 (2)	0.071 (3)	−0.0068 (19)	0.0113 (19)	0.0074 (19)
C3	0.089 (3)	0.064 (3)	0.097 (4)	−0.021 (3)	0.006 (3)	0.021 (3)
C4	0.096 (4)	0.049 (2)	0.111 (5)	−0.018 (2)	0.003 (3)	0.005 (3)
C5	0.084 (3)	0.058 (2)	0.090 (3)	−0.010 (2)	−0.006 (3)	−0.013 (2)
C6	0.0507 (17)	0.0491 (19)	0.072 (2)	0.0005 (15)	0.0021 (16)	−0.0091 (17)
C7	0.0490 (17)	0.059 (2)	0.0559 (19)	0.0040 (15)	0.0048 (14)	−0.0108 (16)
C8	0.075 (3)	0.074 (3)	0.068 (3)	0.002 (2)	0.010 (2)	−0.019 (2)
C9	0.082 (3)	0.095 (4)	0.066 (3)	0.009 (3)	0.011 (2)	−0.018 (3)
C10	0.073 (3)	0.099 (4)	0.058 (2)	0.006 (3)	0.009 (2)	0.010 (2)
C11	0.073 (3)	0.069 (3)	0.057 (2)	0.001 (2)	0.0055 (18)	0.0050 (19)
C12	0.154 (6)	0.071 (3)	0.077 (3)	−0.019 (4)	0.017 (4)	0.014 (3)
N1	0.0525 (15)	0.0479 (16)	0.0617 (17)	−0.0034 (13)	0.0120 (13)	−0.0002 (13)
N2	0.0572 (16)	0.0543 (17)	0.0512 (16)	0.0003 (14)	0.0072 (13)	−0.0035 (13)
Co1	0.0631 (4)	0.0453 (3)	0.0569 (4)	−0.0020 (2)	0.0095 (2)	−0.0053 (2)
Cl1	0.0798 (7)	0.0659 (7)	0.0979 (9)	0.0142 (6)	0.0080 (6)	−0.0239 (6)
Cl2	0.0753 (7)	0.0790 (7)	0.0759 (7)	−0.0148 (6)	0.0013 (5)	−0.0134 (6)

Geometric parameters (Å, °)

C1—C2	1.483 (7)	C8—C9	1.377 (8)
C1—H1A	0.9600	C8—H8	0.9300
C1—H1B	0.9600	C9—C10	1.361 (9)
C1—H1C	0.9600	C9—H9	0.9300
C2—N1	1.330 (5)	C10—C11	1.393 (7)
C2—C3	1.382 (7)	C10—H10	0.9300
C3—C4	1.342 (9)	C11—N2	1.338 (6)
C3—H3	0.9300	C11—C12	1.480 (8)
C4—C5	1.397 (9)	C12—H12A	0.9600
C4—H4	0.9300	C12—H12B	0.9600
C5—C6	1.382 (6)	C12—H12C	0.9600
C5—H5	0.9300	Co1—N1	2.042 (3)
C6—N1	1.358 (5)	Co1—N2	2.053 (3)
C6—C7	1.477 (6)	Co1—Cl1	2.2193 (13)
C7—N2	1.349 (5)	Co1—Cl2	2.2269 (13)
C7—C8	1.365 (5)		
C2—C1—H1A	109.5	C10—C9—C8	119.8 (5)
C2—C1—H1B	109.5	C10—C9—H9	120.1
H1A—C1—H1B	109.5	C8—C9—H9	120.1
C2—C1—H1C	109.5	C9—C10—C11	119.7 (5)
H1A—C1—H1C	109.5	C9—C10—H10	120.2
H1B—C1—H1C	109.5	C11—C10—H10	120.2
N1—C2—C3	120.7 (5)	N2—C11—C10	120.0 (5)
N1—C2—C1	117.3 (4)	N2—C11—C12	116.6 (4)

C3—C2—C1	121.9 (5)	C10—C11—C12	123.4 (5)
C4—C3—C2	120.0 (5)	C11—C12—H12A	109.5
C4—C3—H3	120.0	C11—C12—H12B	109.5
C2—C3—H3	120.0	H12A—C12—H12B	109.5
C3—C4—C5	119.7 (5)	C11—C12—H12C	109.5
C3—C4—H4	120.1	H12A—C12—H12C	109.5
C5—C4—H4	120.1	H12B—C12—H12C	109.5
C6—C5—C4	118.8 (5)	C2—N1—C6	120.7 (4)
C6—C5—H5	120.6	C2—N1—Co1	125.8 (3)
C4—C5—H5	120.6	C6—N1—Co1	113.5 (3)
N1—C6—C5	120.0 (5)	C11—N2—C7	120.0 (4)
N1—C6—C7	116.2 (3)	C11—N2—Co1	126.3 (3)
C5—C6—C7	123.9 (4)	C7—N2—Co1	113.7 (3)
N2—C7—C8	121.6 (4)	N1—Co1—N2	81.00 (14)
N2—C7—C6	115.7 (3)	N1—Co1—Cl1	114.87 (10)
C8—C7—C6	122.7 (4)	N2—Co1—Cl1	114.93 (10)
C7—C8—C9	118.8 (5)	N1—Co1—Cl2	115.70 (10)
C7—C8—H8	120.6	N2—Co1—Cl2	118.35 (10)
C9—C8—H8	120.6	Cl1—Co1—Cl2	109.67 (5)
N1—C2—C3—C4	1.6 (8)	C5—C6—N1—Co1	−179.0 (3)
C1—C2—C3—C4	179.4 (6)	C7—C6—N1—Co1	−0.2 (4)
C2—C3—C4—C5	−2.2 (9)	C10—C11—N2—C7	1.3 (7)
C3—C4—C5—C6	1.7 (9)	C12—C11—N2—C7	−177.7 (5)
C4—C5—C6—N1	−0.5 (7)	C10—C11—N2—Co1	−179.2 (4)
C4—C5—C6—C7	−179.2 (5)	C12—C11—N2—Co1	1.7 (7)
N1—C6—C7—N2	−1.0 (5)	C8—C7—N2—C11	1.0 (6)
C5—C6—C7—N2	177.7 (4)	C6—C7—N2—C11	−178.9 (4)
N1—C6—C7—C8	179.2 (4)	C8—C7—N2—Co1	−178.5 (3)
C5—C6—C7—C8	−2.1 (6)	C6—C7—N2—Co1	1.6 (4)
N2—C7—C8—C9	−2.6 (7)	C2—N1—Co1—N2	−178.0 (3)
C6—C7—C8—C9	177.2 (4)	C6—N1—Co1—N2	0.8 (3)
C7—C8—C9—C10	1.9 (8)	C2—N1—Co1—Cl1	68.6 (3)
C8—C9—C10—C11	0.4 (8)	C6—N1—Co1—Cl1	−112.6 (2)
C9—C10—C11—N2	−2.0 (8)	C2—N1—Co1—Cl2	−60.8 (3)
C9—C10—C11—C12	177.0 (6)	C6—N1—Co1—Cl2	118.0 (2)
C3—C2—N1—C6	−0.4 (7)	C11—N2—Co1—N1	179.2 (4)
C1—C2—N1—C6	−178.3 (4)	C7—N2—Co1—N1	−1.4 (3)
C3—C2—N1—Co1	178.3 (4)	C11—N2—Co1—Cl1	−67.5 (4)
C1—C2—N1—Co1	0.4 (6)	C7—N2—Co1—Cl1	111.9 (3)
C5—C6—N1—C2	−0.1 (6)	C11—N2—Co1—Cl2	64.7 (4)
C7—C6—N1—C2	178.7 (3)	C7—N2—Co1—Cl2	−115.8 (2)

Hydrogen-bond geometry (Å, °)

D—H···*A*	*D*—H	H···*A*	*D*···*A*	*D*—H···*A*
C5—H5···Cl1^{i}	0.93	2.82	3.565 (7)	138

Symmetry codes: (i) −*x*+1, −*y*+1, −*z*+1.

Fig. 1

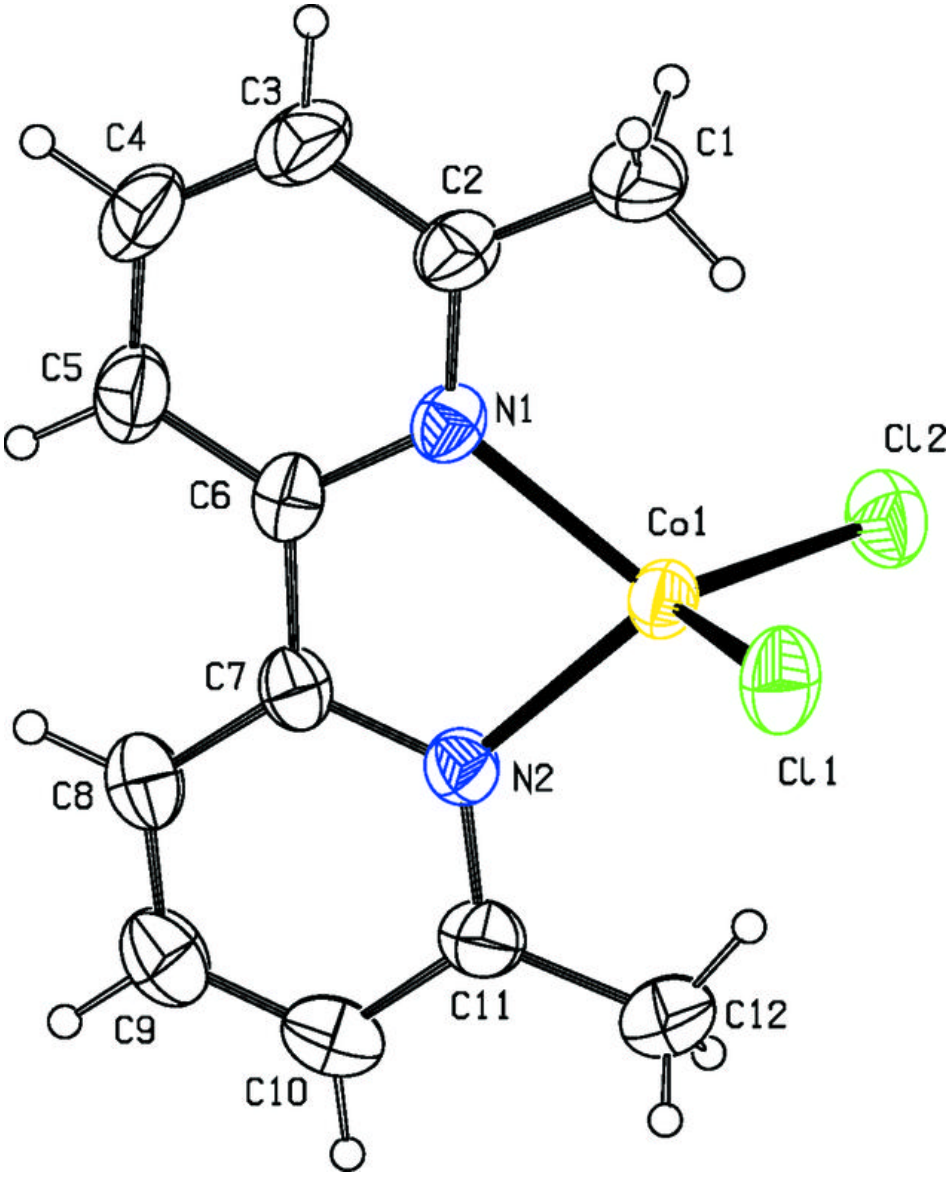

Fig. 2