[Bis(2-pyridyl)amine-*N*,*N*'](nitrato-*O*,*O*')cobalt(II) nitrate. Corrigendum

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The chemical name and formula in the paper by Castillo, Luque, De la Pinta & Román [*Acta Cryst.* (2001), E**57**, m384–m386] is corrected.

In the paper by Castillo, Luque, De la Pinta & Román (2001), the ligand reported as nitrate should be carbonate and the oxidation state of the cobalt metal atom should be Co[III] rather than Co[II], thus making the correct chemical composition $[Co(CO_3)(C_{10}H_9N_3)_2]NO_3$ and the correct chemical name '[Bis(2-pyridyl)amine-$\kappa^2 N$,*N*'](carbonato-$\kappa^2 O$,*O*')cobalt(III) nitrate'.

Experimental

Crystal data

$[Co(CO_3)(C_{10}H_9N_3)_2]NO_3$	V = 2165.9 (8) Å^3
M_r = 523.35	Z = 4
Monoclinic, $P2_1/n$	Mo $K\alpha$ radiation
a = 17.191 (3) Å	μ = 0.85 mm^{-1}
b = 7.3080 (10) Å	T = 293 K
c = 17.843 (5) Å	$0.42 \times 0.20 \times 0.08$ mm
β = 104.94 (3)°	

Data collection

Stoe IPDS diffractometer	14084 measured reflections
Absorption correction: numerical	4037 independent reflections
(Stoe & Cie, 1998)	2598 reflections with $I > 2\sigma(I)$
T_{min} = 0.815, T_{max} = 0.934	R_{int} = 0.048

Refinement

$R[F^2 > 2\sigma(F^2)]$ = 0.031	316 parameters
$wR(F^2)$ = 0.076	H-atom parameters constrained
S = 0.82	$\Delta\rho_{max}$ = 0.37 e Å^{-3}
4037 reflections	$\Delta\rho_{min}$ = −0.31 e Å^{-3}

Data collection, cell refinement and data reduction: *IPDS Software* (Stoe & Cie, 1998); program(s) used to solve structure: *SIR92* (Altomare *et al.*, 1993); program(s) used to refine structure: *SHELXL93* (Sheldrick, 1993).

Supplementary data and figures for this paper are available from the IUCr electronic archives (Reference: BT9068).

References

Altomare, A., Cascarano, G., Giacovazzo, C. & Guagliardi, A. (1993). *J. Appl. Cryst.* **26**, 343–350.

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