Table 1. *Final least-squares parameters obtained with weights* $w = (\sigma^2 + 0{\cdot}0005I)^{-1}$

Thermal parameters are the coefficients U_{ij} (in Å²), multiplied by 10^4, in the form:

$$\exp\left[-(U_{11}h^2 + U_{22}k^2 + U_{33}l^2 + 2U_{12}hk + 2U_{13}hl + 2U_{23}kl)\right].$$

	x	y	z	U_{11}	U_{22}	U_{33}	U_{12}	U_{13}	U_{23}
Cl	0·4318 (1)	0·2500	0·1916 (1)	289 (3)	286 (3)	989 (7)	0	13 (3)	0
O(1)	0·3142 (3)	0·2500	0·0664 (4)	540 (13)	522 (13)	1059 (0)	0	−169 (14)	–
O(2)	0·5680 (3)	0·2500	0·1019 (5)	444 (12)	699 (17)	1351 (26)	0	248 (15)	0
O(3)	0·4203 (2)	0·0487 (2)	0·3036 (3)	485 (8)	324 (6)	1171 (15)	12 (6)	−33 (9)	101 (8)
N	0·3191 (2)	0·2500	0·6645 (5)	323 (10)	366 (11)	1058 (23)	3	25 (13)	0

Å, are, within one standard deviation, the same as those of PP (obtained with the positions of the ammonium ions held fixed). The thermal ellipsoids of all atoms in Table 1 are highly anisotropic, with U_{33} two to three times larger than U_{11} or U_{22} throughout. The observed atomic thermal motions of the perchlorate group agree extremely well with values obtained by a rigid-body analysis with the method of Schomaker & Trueblood (1968), with an average difference in U_{ij} of 0·0002 Å². The librational motion of the perchlorate group, as obtained from the rigid-body refinement, is almost isotropic (7·1, 6·9, 6·5° for the amplitudes about principal axes), and causes the apparent Cl–O bond length to be shortened by about 0·02 Å. The translational motion, however, is extremely anisotropic and closely resembles the thermal ellipsoid of the N atom of the ammonium group (0·0291, 0·0281 and 0·0987 Å² for the three principal axes of **T**). It is possible that this relative large translational motion along **c** is correlated with the occurrence of highly temperature-dependent lattice modes observed *via* Raman scattering (Rosasco & Prask, 1975, 1976).

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Acta Cryst. (1976). B**32**, 2920

2-Aminopyridine and 3-aminopyridine. Errata.

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(*Received* 28 *May* 1976; *accepted* 28 *May* 1976)

The figures giving the bond distances and angles in 2-aminopyridine and 3-aminopyridine, by Chao, Schempp & Rosenstein [*Acta Cryst.* (1975), B31, 2922–2924 and 2924–2926], have been transposed. Fig. 1 of the paper on 3-aminopyridine shows the results for 2-aminopyridine and Fig. 2 of the paper on 2-aminopyridine shows those for 3-aminopyridine.

All the relevant information is given in the abstract.

We are grateful to Dr K. D. Holmes for bringing these errors to our attention.

Acta Cryst. (1976). B**32**, 2920

The crystal structure of [1,4-di-(*N*-pyridiniummethyl)benzene]$^{2+}$ (7,7,8,8-tetracyanoquinodimethane)$_4^{2-}$. Erratum.

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(*Received* 1 *June* 1976; *accepted* 4 *June* 1976)

In the previous paper on the title complex [*Acta Cryst.* (1975), B31, 1174–1178] the TCNQ stacking direction was erroneously described as parallel to **b**. It is, in fact, parallel to [110].

In our paper on the crystal structure of 1,4-di-(*N*-pyridiniummethyl)benzene(TCNQ)$_4$ (Ashwell, Wallwork, Baker & Berthier, 1975) we stated that the TCNQ molecules and anions were packed plane-to-plane in columns parallel to **b**. In fact the columns are parallel to [110], *i.e.* in the (001) plane and along the positive diagonal between **a** and **b**. The atomic coordinates, intra- and intermolecular distances and diagrams given in the paper are correct.

Reference

Ashwell, G. J., Wallwork, S. C., Baker, S. R. & Berthier, P. I. C. (1975). *Acta Cryst.* B**31**, 1174–1178.