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Cyclohexane-1,2-diammonium bis(pyridine-2-carboxylate)

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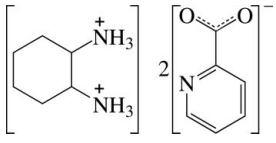
Key indicators: single-crystal X-ray study; T = 293 K; mean σ(C–C) = 0.005 Å; R factor = 0.064; wR factor = 0.171; data-to-parameter ratio = 16.3.

In the dication of the title salt, $C_6H_{16}N_2^{2+}\cdot 2C_6H_4NO_2^-$, the two ammonium groups are in the equatorial positions of the chair-shaped cyclohexyl ring. In the crystal, the cations and anions are linked by N—H···O and N—H···N hydrogen bonds, forming a layer network parallel to the *ac* plane. Weak π–π interactions between adjacent pyridine rings with a centroid–centroid distance of 3.589 (2) Å are also present.

Related literature

For the syntheses and structures of cyclohexane-1,2-diammonium compounds, see: Lin & Lii (1998); Lin & Wang (2000). For the crystal structures of pyridine-2-carboxylates, see: Kim & Ha (2009*a*,*b*,*c*).



Experimental

Crystal data

$C_6H_{16}N_2^{2+}\cdot 2C_6H_4NO_2^-$
M_r = 360.41
Monoclinic, $P2_1/n$
a = 9.2942 (11) Å
b = 20.329 (2) Å
c = 10.2189 (11) Å
β = 101.775 (3)°
V = 1890.1 (4) Å^3
Z = 4
Mo $K\alpha$ radiation
μ = 0.09 mm^{-1}
T = 293 K
0.20 × 0.10 × 0.10 mm

Data collection

Bruker SMART 1000 CCD diffractometer
Absorption correction: none
11005 measured reflections
3854 independent reflections
1741 reflections with $I > 2\sigma(I)$
R_{int} = 0.064

Refinement

$R[F^2 > 2\sigma(F^2)]$ = 0.064
$wR(F^2)$ = 0.171
S = 0.98
3854 reflections
237 parameters
H-atom parameters constrained
$\Delta\rho_{max}$ = 0.18 e Å^{-3}
$\Delta\rho_{min}$ = −0.17 e Å^{-3}

Table 1
Hydrogen-bond geometry (Å, °).

D—H···A	D—H	H···A	D···A	D—H···A
N1—H1A···O3	0.86	1.89	2.749 (3)	175
N1—H1B···O2	0.86	1.92	2.743 (3)	160
N1—H1C···O3^{i}	0.86	2.10	2.790 (3)	137
N1—H1C···N4^{i}	0.86	2.49	3.271 (4)	152
N2—H2A···O1^{ii}	0.86	2.09	2.828 (3)	144
N2—H2A···N3^{ii}	0.86	2.53	3.254 (4)	142
N2—H2B···O1	0.86	2.02	2.807 (3)	152
N2—H2C···O4^{iii}	0.86	1.88	2.734 (3)	171

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

Data collection: *SMART* (Bruker, 2000); cell refinement: *SAINT* (Bruker, 2000); data reduction: *SAINT*; program(s) used to solve structure: *SHELXS97* (Sheldrick, 2008); program(s) used to refine structure: *SHELXL97* (Sheldrick, 2008); molecular graphics: *ORTEP-3* (Farrugia, 1997) and *PLATON* (Spek, 2009); software used to prepare material for publication: *SHELXL97*.

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Supplementary data and figures for this paper are available from the IUCr electronic archives (Reference: NG2649).

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supplementary materials

Acta Cryst. (2009). E**65**, o2571 [**doi:10.1107/S1600536809038689**]

Cyclohexane-1,2-diammonium bis(pyridine-2-carboxylate)

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Comment

The title compound, $C_6H_{16}N_2^{2+}.2C_6H_4NO_2^-$, consists of a doubly protonated cyclohexane-1,2-diammonium dication and two pyridine-2-carboxylate anions (Fig. 1). The dication has two chiral carbon atoms (C1 and C2), and is one of four possible stereoisomers. Both chiral atoms have *R* configuration. The cyclohexane ring of the dication adopts a strain-free chair conformation. The C—C—C bond angles lie in the range of 109.3 (3)°–111.6 (3)°, close to the ideal tetrahedral angle, and all neighboring C—H bonds are staggered. The diammonium groups within the dication are on opposite faces of the cyclohexane ring, that is, *trans* with respect to each other, and therefore the dication exists in the diequatorial conformation. The N1—C1—C2—N2 torsion angle of -59.0 (3)° displays the *gauche* conformation for the four atoms and there is a *gauche* interaction between the two NH_3^+ groups. The carboxylate groups of the anions appear to be delocalized on the basis of the C—O bond lengths [C—O: 1.239 (3)–1.255 (3) Å]. In the crystal structure, the component ions interact by means of many intermolecular N—H···O and N—H···N hydrogen bonds to form a two-dimensional network parallel to the *ac* plane (Table 1 and Fig. 2). There may also be intermolecular π-π interactions between adjacent pyridine rings, with a centroid-centroid distance of 3.589 (2) Å.

Experimental

A solution of a mixture of *cis* and *trans* isomers of 1,2-diaminocyclohexane (0.202 g, 1.769 mmol) and pyridine-2-carboxylic acid (0.294 g, 2.388 mmol) in H_2O (10 ml) was stirred for 3 h at 60 °C. The solvent was removed under vacuum and the residue was washed with ether/acetone/$CHCl_3$, to give a white powder (0.112 g). Crystals suitable for X-ray analysis were obtained by slow evaporation from a CH_3CN solution.

Refinement

H atoms were positioned geometrically and allowed to ride on their respective parent atoms [C—H = 0.98 (CH), 0.97 (CH_2) or 0.93 (aromatic) Å and N—H = 0.86 Å and U_{iso}(H) = 1.2U_{eq}(C, N)].

Figures

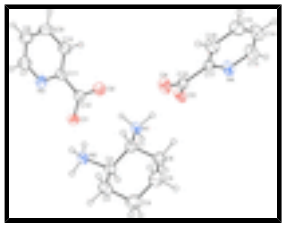

Fig. 1. The structure of the title compound, with displacement ellipsoids drawn at the 30% probability level for non-H atoms.

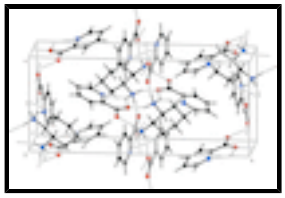

Fig. 2. View of the unit-cell contents of the title compound. Hydrogen-bond interactions are drawn with dashed lines.

Cyclohexane-1,2-diammonium bis(pyridine-2-carboxylate)

Crystal data

$C_6H_{16}N_2^{2+}\cdot 2C_6H_4NO_2^-$	F_{000} = 768
M_r = 360.41	D_x = 1.267 Mg m^{-3}
Monoclinic, $P2_1/n$	Mo $K\alpha$ radiation, λ = 0.71073 Å
Hall symbol: -P 2yn	Cell parameters from 784 reflections
a = 9.2942 (11) Å	θ = 2.3–17.2º
b = 20.329 (2) Å	μ = 0.09 mm^{-1}
c = 10.2189 (11) Å	T = 293 K
β = 101.775 (3)º	Rod, colorless
V = 1890.1 (4) Å^3	0.20 × 0.10 × 0.10 mm
Z = 4	

Data collection

Bruker SMART 1000 CCD diffractometer	1741 reflections with $I > 2\sigma(I)$
Radiation source: fine-focus sealed tube	R_{int} = 0.064
Monochromator: graphite	θ_{max} = 26.4º
T = 293 K	θ_{min} = 2.3º
φ and ω scans	h = −11→11
Absorption correction: None	k = −25→25
11005 measured reflections	l = −12→7
3854 independent reflections	

Refinement

Refinement on F^2	Secondary atom site location: difference Fourier map
Least-squares matrix: full	Hydrogen site location: inferred from neighbouring sites
$R[F^2 > 2\sigma(F^2)]$ = 0.064	H-atom parameters constrained
$wR(F^2)$ = 0.171	$w = 1/[\sigma^2(F_o^2) + (0.0581P)^2]$ where $P = (F_o^2 + 2F_c^2)/3$
S = 0.98	$(\Delta/\sigma)_{max} < 0.001$
3854 reflections	$\Delta\rho_{max}$ = 0.18 e Å^{-3}
237 parameters	$\Delta\rho_{min}$ = −0.16 e Å^{-3}
Primary atom site location: structure-invariant direct methods	Extinction correction: none

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 $F^2 > \sigma(F^2)$ 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)

	x	*y*	*z*	U_{iso}*/U_{eq}
O1	0.5813 (2)	0.56693 (10)	0.5900 (2)	0.0580 (6)
O2	0.6448 (3)	0.60901 (12)	0.7932 (2)	0.0874 (9)
O3	0.9341 (2)	0.53730 (12)	1.1294 (2)	0.0629 (7)
O4	0.7329 (2)	0.53120 (12)	1.2114 (2)	0.0705 (7)
N1	0.7776 (2)	0.49308 (12)	0.8877 (2)	0.0497 (7)
H1A	0.8252	0.5093	0.9617	0.060*
H1B	0.7247	0.5233	0.8425	0.060*
H1C	0.8387	0.4780	0.8423	0.060*
N2	0.5142 (2)	0.44461 (12)	0.6901 (2)	0.0515 (7)
H2A	0.4887	0.4232	0.6164	0.062*
H2B	0.5633	0.4790	0.6768	0.062*
H2C	0.4370	0.4563	0.7182	0.062*
N3	0.4687 (3)	0.68504 (14)	0.4928 (3)	0.0660 (8)
N4	1.1006 (3)	0.56390 (14)	1.3686 (3)	0.0606 (8)
C1	0.6809 (3)	0.43883 (15)	0.9164 (3)	0.0469 (8)
H1	0.6048	0.4573	0.9593	0.056*
C2	0.6076 (3)	0.40179 (15)	0.7922 (3)	0.0494 (8)
H2	0.6844	0.3820	0.7519	0.059*
C3	0.5124 (4)	0.34650 (16)	0.8301 (3)	0.0646 (10)
H3A	0.4353	0.3652	0.8698	0.078*
H3B	0.4664	0.3227	0.7503	0.078*
C4	0.6042 (4)	0.29907 (18)	0.9290 (4)	0.0789 (11)
H4A	0.5408	0.2657	0.9550	0.095*
H4B	0.6754	0.2772	0.8865	0.095*
C5	0.6839 (5)	0.33547 (18)	1.0523 (3)	0.0802 (12)
H5A	0.6128	0.3529	1.1007	0.096*
H5B	0.7474	0.3052	1.1108	0.096*
C6	0.7750 (4)	0.39129 (16)	1.0134 (3)	0.0625 (10)
H6A	0.8516	0.3733	0.9722	0.075*
H6B	0.8219	0.4150	1.0931	0.075*
C7	0.4020 (5)	0.74176 (19)	0.4510 (4)	0.0801 (12)
H7	0.3641	0.7465	0.3600	0.096*
C8	0.3859 (5)	0.79253 (18)	0.5318 (4)	0.0832 (12)

H8	0.3372	0.8307	0.4977	0.100*
C9	0.4430 (5)	0.78617 (18)	0.6647 (4)	0.0863 (13)
H9	0.4344	0.8203	0.7231	0.104*
C10	0.5136 (4)	0.72897 (17)	0.7119 (3)	0.0668 (10)
H10	0.5546	0.7241	0.8024	0.080*
C11	0.5224 (3)	0.67897 (15)	0.6232 (3)	0.0493 (8)
C12	0.5894 (3)	0.61329 (16)	0.6723 (4)	0.0542 (9)
C13	1.1815 (4)	0.58402 (19)	1.4852 (4)	0.0769 (11)
H13	1.2833	0.5832	1.4958	0.092*
C14	1.1234 (4)	0.60559 (18)	1.5892 (4)	0.0715 (11)
H14	1.1846	0.6196	1.6678	0.086*
C15	0.9757 (5)	0.60653 (18)	1.5775 (4)	0.0723 (11)
H15	0.9338	0.6212	1.6474	0.087*
C16	0.8879 (4)	0.58488 (17)	1.4573 (3)	0.0646 (10)
H16	0.7861	0.5843	1.4463	0.078*
C17	0.9540 (3)	0.56461 (14)	1.3564 (3)	0.0452 (8)
C18	0.8658 (4)	0.54258 (15)	1.2225 (3)	0.0496 (8)

Atomic displacement parameters (Å^2)

	U^{11}	U^{22}	U^{33}	U^{12}	U^{13}	U^{23}
O1	0.0663 (15)	0.0528 (14)	0.0526 (14)	0.0046 (11)	0.0071 (11)	−0.0075 (11)
O2	0.126 (2)	0.0742 (18)	0.0482 (16)	0.0264 (16)	−0.0142 (15)	−0.0071 (12)
O3	0.0525 (14)	0.0915 (18)	0.0459 (14)	−0.0065 (12)	0.0128 (11)	−0.0127 (12)
O4	0.0427 (14)	0.102 (2)	0.0660 (17)	−0.0010 (13)	0.0095 (12)	−0.0170 (13)
N1	0.0466 (16)	0.0592 (17)	0.0403 (15)	0.0014 (13)	0.0018 (12)	−0.0019 (12)
N2	0.0448 (15)	0.0642 (18)	0.0444 (16)	−0.0042 (13)	0.0061 (12)	−0.0097 (13)
N3	0.077 (2)	0.062 (2)	0.053 (2)	0.0113 (16)	−0.0005 (15)	−0.0010 (14)
N4	0.0504 (18)	0.078 (2)	0.0512 (18)	−0.0100 (15)	0.0052 (14)	−0.0030 (14)
C1	0.0436 (18)	0.055 (2)	0.0424 (19)	−0.0030 (16)	0.0109 (15)	0.0000 (15)
C2	0.0467 (18)	0.057 (2)	0.0450 (19)	0.0038 (16)	0.0096 (15)	−0.0014 (15)
C3	0.069 (2)	0.065 (2)	0.062 (2)	−0.014 (2)	0.0177 (19)	−0.0052 (18)
C4	0.105 (3)	0.061 (2)	0.072 (3)	−0.006 (2)	0.021 (2)	0.007 (2)
C5	0.111 (3)	0.069 (3)	0.061 (3)	0.002 (2)	0.018 (2)	0.013 (2)
C6	0.073 (2)	0.062 (2)	0.050 (2)	0.0062 (19)	0.0045 (17)	0.0067 (17)
C7	0.104 (3)	0.070 (3)	0.058 (3)	0.012 (2)	−0.002 (2)	0.001 (2)
C8	0.107 (3)	0.056 (3)	0.083 (3)	0.012 (2)	0.011 (3)	0.004 (2)
C9	0.132 (4)	0.048 (2)	0.081 (3)	0.005 (2)	0.025 (3)	−0.016 (2)
C10	0.086 (3)	0.054 (2)	0.058 (2)	−0.002 (2)	0.0076 (19)	−0.0057 (18)
C11	0.0482 (19)	0.050 (2)	0.050 (2)	−0.0064 (16)	0.0087 (16)	−0.0029 (16)
C12	0.052 (2)	0.057 (2)	0.052 (2)	−0.0021 (17)	0.0063 (17)	−0.0047 (18)
C13	0.059 (2)	0.106 (3)	0.062 (3)	−0.019 (2)	0.004 (2)	−0.004 (2)
C14	0.079 (3)	0.088 (3)	0.045 (2)	−0.022 (2)	0.006 (2)	−0.0053 (19)
C15	0.083 (3)	0.083 (3)	0.056 (3)	0.001 (2)	0.028 (2)	−0.0101 (19)
C16	0.054 (2)	0.087 (3)	0.056 (2)	−0.0048 (19)	0.0173 (19)	−0.0143 (19)
C17	0.046 (2)	0.0454 (19)	0.043 (2)	−0.0001 (15)	0.0074 (15)	0.0047 (14)
C18	0.044 (2)	0.053 (2)	0.051 (2)	0.0054 (16)	0.0081 (17)	−0.0018 (16)

Geometric parameters (Å, °)

O1—C12	1.255 (3)	C4—H4A	0.9700
O2—C12	1.241 (3)	C4—H4B	0.9700
O3—C18	1.251 (3)	C5—C6	1.517 (5)
O4—C18	1.239 (3)	C5—H5A	0.9700
N1—C1	1.489 (3)	C5—H5B	0.9700
N1—H1A	0.8600	C6—H6A	0.9700
N1—H1B	0.8600	C6—H6B	0.9700
N1—H1C	0.8600	C7—C8	1.349 (5)
N2—C2	1.493 (3)	C7—H7	0.9300
N2—H2A	0.8600	C8—C9	1.359 (5)
N2—H2B	0.8600	C8—H8	0.9300
N2—H2C	0.8600	C9—C10	1.374 (5)
N3—C11	1.330 (4)	C9—H9	0.9300
N3—C7	1.337 (4)	C10—C11	1.376 (4)
N4—C13	1.337 (4)	C10—H10	0.9300
N4—C17	1.343 (4)	C11—C12	1.515 (4)
C1—C2	1.513 (4)	C13—C14	1.359 (5)
C1—C6	1.526 (4)	C13—H13	0.9300
C1—H1	0.9800	C14—C15	1.353 (5)
C2—C3	1.528 (4)	C14—H14	0.9300
C2—H2	0.9800	C15—C16	1.400 (4)
C3—C4	1.525 (4)	C15—H15	0.9300
C3—H3A	0.9700	C16—C17	1.368 (4)
C3—H3B	0.9700	C16—H16	0.9300
C4—C5	1.517 (5)	C17—C18	1.512 (4)
C1—N1—H1A	109.5	H5A—C5—H5B	108.1
C1—N1—H1B	109.5	C5—C6—C1	111.6 (3)
H1A—N1—H1B	109.5	C5—C6—H6A	109.3
C1—N1—H1C	109.5	C1—C6—H6A	109.3
H1A—N1—H1C	109.5	C5—C6—H6B	109.3
H1B—N1—H1C	109.5	C1—C6—H6B	109.3
C2—N2—H2A	109.5	H6A—C6—H6B	108.0
C2—N2—H2B	109.5	N3—C7—C8	124.5 (4)
H2A—N2—H2B	109.5	N3—C7—H7	117.8
C2—N2—H2C	109.5	C8—C7—H7	117.8
H2A—N2—H2C	109.5	C7—C8—C9	118.0 (4)
H2B—N2—H2C	109.5	C7—C8—H8	121.0
C11—N3—C7	117.1 (3)	C9—C8—H8	121.0
C13—N4—C17	117.1 (3)	C8—C9—C10	119.5 (3)
N1—C1—C2	112.9 (2)	C8—C9—H9	120.2
N1—C1—C6	108.0 (2)	C10—C9—H9	120.2
C2—C1—C6	109.3 (3)	C9—C10—C11	118.8 (3)
N1—C1—H1	108.9	C9—C10—H10	120.6
C2—C1—H1	108.9	C11—C10—H10	120.6
C6—C1—H1	108.9	N3—C11—C10	122.1 (3)
N2—C2—C1	113.2 (2)	N3—C11—C12	117.3 (3)

N2—C2—C3	108.8 (2)	C10—C11—C12	120.6 (3)
C1—C2—C3	109.8 (2)	O2—C12—O1	124.8 (3)
N2—C2—H2	108.3	O2—C12—C11	116.8 (3)
C1—C2—H2	108.3	O1—C12—C11	118.4 (3)
C3—C2—H2	108.3	N4—C13—C14	123.7 (4)
C4—C3—C2	111.0 (3)	N4—C13—H13	118.2
C4—C3—H3A	109.4	C14—C13—H13	118.2
C2—C3—H3A	109.4	C15—C14—C13	119.6 (3)
C4—C3—H3B	109.4	C15—C14—H14	120.2
C2—C3—H3B	109.4	C13—C14—H14	120.2
H3A—C3—H3B	108.0	C14—C15—C16	118.2 (3)
C5—C4—C3	110.7 (3)	C14—C15—H15	120.9
C5—C4—H4A	109.5	C16—C15—H15	120.9
C3—C4—H4A	109.5	C17—C16—C15	119.0 (3)
C5—C4—H4B	109.5	C17—C16—H16	120.5
C3—C4—H4B	109.5	C15—C16—H16	120.5
H4A—C4—H4B	108.1	N4—C17—C16	122.4 (3)
C6—C5—C4	110.6 (3)	N4—C17—C18	115.7 (3)
C6—C5—H5A	109.5	C16—C17—C18	121.9 (3)
C4—C5—H5A	109.5	O4—C18—O3	124.4 (3)
C6—C5—H5B	109.5	O4—C18—C17	119.0 (3)
C4—C5—H5B	109.5	O3—C18—C17	116.6 (3)
N1—C1—C2—N2	−59.0 (3)	C9—C10—C11—C12	−175.5 (3)
C6—C1—C2—N2	−179.2 (2)	N3—C11—C12—O2	177.0 (3)
N1—C1—C2—C3	179.2 (2)	C10—C11—C12—O2	−5.3 (5)
C6—C1—C2—C3	59.0 (3)	N3—C11—C12—O1	−5.1 (4)
N2—C2—C3—C4	177.0 (3)	C10—C11—C12—O1	172.6 (3)
C1—C2—C3—C4	−58.6 (4)	C17—N4—C13—C14	−0.9 (5)
C2—C3—C4—C5	56.3 (4)	N4—C13—C14—C15	0.9 (6)
C3—C4—C5—C6	−54.9 (4)	C13—C14—C15—C16	0.1 (6)
C4—C5—C6—C1	56.8 (4)	C14—C15—C16—C17	−0.9 (5)
N1—C1—C6—C5	177.9 (3)	C13—N4—C17—C16	0.1 (5)
C2—C1—C6—C5	−58.9 (4)	C13—N4—C17—C18	179.1 (3)
C11—N3—C7—C8	−0.1 (6)	C15—C16—C17—N4	0.8 (5)
N3—C7—C8—C9	1.0 (7)	C15—C16—C17—C18	−178.1 (3)
C7—C8—C9—C10	−0.4 (6)	N4—C17—C18—O4	167.7 (3)
C8—C9—C10—C11	−1.0 (6)	C16—C17—C18—O4	−13.2 (5)
C7—N3—C11—C10	−1.5 (5)	N4—C17—C18—O3	−12.5 (4)
C7—N3—C11—C12	176.2 (3)	C16—C17—C18—O3	166.5 (3)
C9—C10—C11—N3	2.0 (5)		

Hydrogen-bond geometry (Å, °)

D—H···*A*	*D*—H	H···*A*	*D*···*A*	*D*—H···*A*
N1—H1A···O3	0.86	1.89	2.749 (3)	175
N1—H1B···O2	0.86	1.92	2.743 (3)	160
N1—H1C···O3^{i}	0.86	2.10	2.790 (3)	137
N1—H1C···N4^{i}	0.86	2.49	3.271 (4)	152

N2—H2A···O1^{ii}	0.86	2.09	2.828 (3)	144
N2—H2A···N3^{ii}	0.86	2.53	3.254 (4)	142
N2—H2B···O1	0.86	2.02	2.807 (3)	152
N2—H2C···O4^{iii}	0.86	1.88	2.734 (3)	171

Symmetry codes: (i) −*x*+2, −*y*+1, −*z*+2; (ii) −*x*+1, −*y*+1, −*z*+1; (iii) −*x*+1, −*y*+1, −*z*+2.

Fig. 1

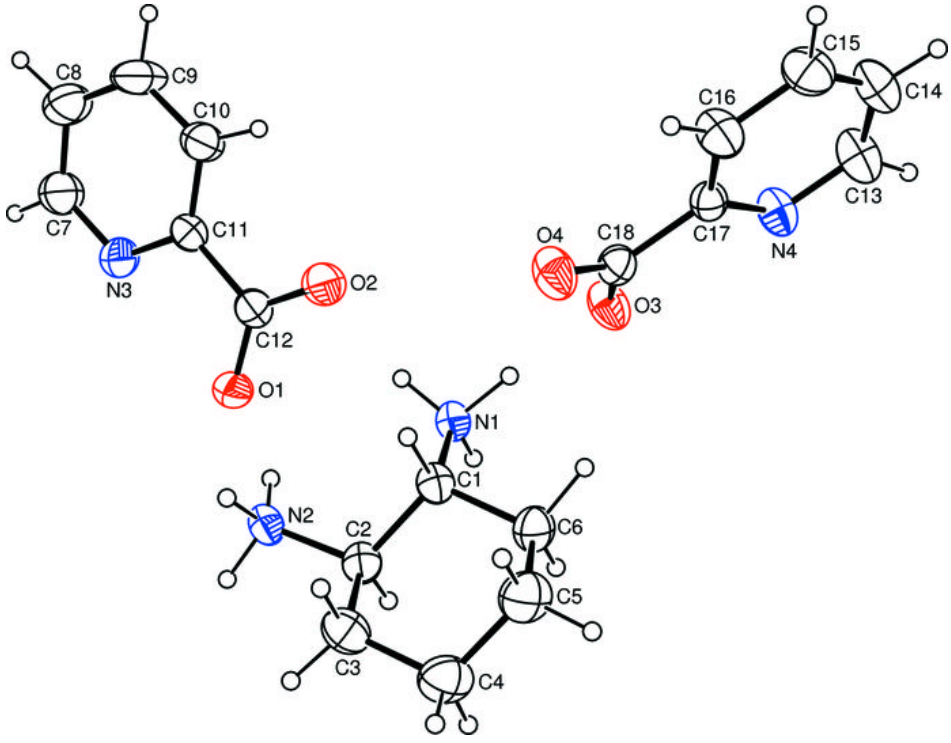
C8
C9
C10
C7
C11
N3
O2
C12
O1
C15
C16
C14
C13
O4
C18
C17
N4
O3
N1
C1
N2
C2
C6
C5
C3
C4

Fig. 2