Hydrogen bonding and proton transfer in cocrystals of 4,4’-bipyridyl and organic acids studied using nuclear quadrupole resonance

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Cocrystals of 4,4’-bipyridyl and several carboxylic acids were grown from the methanol solution of the cocrystal formers. Complete 14N NQR spectra of these cocrystals have been measured using 1H–14N nuclear quadrupole double resonance. The principal values of the quadrupole coupling tensor are calculated from the 14N NQR frequencies. A large variation in the 14N quadrupole coupling constant between 1.3 MHz and 4.7 MHz is observed. A very low 14N quadrupole coupling constant, characteristic for proton transfer O–H···N → O···H–N+, is observed in 4,4’-bipyridyl–oxalic acid (1:1). In 4,4’-bipyridyl–5-chlorosalicylic acid (1:1) the 14N NQR data show the presence of a short, strong N···H···O hydrogen bond. A correlation of the principal values of the 14N quadrupole coupling tensor is observed. The correlation is analyzed in the model, where the deformation of the lone pair electron orbital and the change of the population of the π-electron orbital produce the variation of the 14N quadrupole coupling tensor in the hydrogen bonded 4,4’-bipyridyl. The temperature variation of the 14N quadrupole coupling tensor in 4,4’-bipyridyl–5-chlorosalicylic acid (1:1) is analyzed. Proton displacement within the N···H···O hydrogen bond and the change of the population of the π-electron orbital at the two nitrogen positions in a 4,4’-bipyridyl molecule in the temperature interval between 157 K and 323 K are determined.

Introduction

Crystal engineering is the understanding of intermolecular interactions in the context of crystal packing and the utilization of such understanding in the design of new solids with desired physical, chemical and pharmaceutical properties.1 It is an interplay between the structure and properties of molecules on one hand and those of extended assemblies of molecules on the other. More precisely, one attempts to understand crystal structure and properties in terms of molecular structure and intermolecular interactions.

Cocrystals are often used in crystal engineering. A cocrystal is a nonionic supramolecular complex, which is constructed through several types of interaction, including hydrogen bonding, π–π stacking, and van der Waals forces.2–4 In a cocrystal, homosynthons and heterosynthons generally occur. Their occurrence depends on the molecular architecture and the positions and properties of the functional groups.

Several cocrystals and crystals may exist in more than one crystal structure. These polymorphic forms of a given substance in general differ in the macroscopic properties. Various polymorphs can be obtained by changing the conditions (solvent, temperature, pressure...) during the crystallization. 4,4’-Bipyridyl (BPY, Fig. 1) is a double hydrogen bond acceptor. It is thus a suitable compound to form hydrogen

![Structural formulae of cocrystal formers.](image)
bonded cocrystals with organic acids. Several cocrystals of
4,4′-bipyridyl have already been studied.

The crystal structure of the hydrogen bonded, 1 : 1 molecular
complex of malonic acid with 4,4′-bipyridyl has been deter-
mained. Molecular tape structures are formed where the mole-
cules are bound through strong O–H···N and weak C–H···O
hydrogen bonds. The molecular tapes are arranged in a crossed
fashion.5

BPY and oxalic acid crystallize in a 1 : 1 ratio. The asymmetric
unit consists of one oxalic acid molecule and one BPY molecule
in general positions, together with one half oxalic acid molecule
and one half BPY molecule; the latter two molecules are centro-
symmetric. The molecules are linked in two parallel independent
chains by strong O–H···N hydrogen bonds. In one chain there is
one independent O–H···N hydrogen bond and the molecules lie
on the centres of symmetry. The second chain contains two
independent O–H···N hydrogen bonds and both molecules are
twisted about their central bonds.6

The 1 : 2 adduct of benzene-1,2,4,5-tetracarboxylic acid and
BPY has been studied using single-crystal neutron diffraction.
The structure contains 20 K two k, strong N···O hydrogen bonds:
one O–H···N hydrogen bond and one N···O hydrogen bond. The N–H
distance in the strong, short N···O hydrogen bond changes from 1.207 Å
at 20 K to 1.302 Å 296 K.2 The N–H distance in the short, weak
N···H hydrogen bond is 1.302 Å at 200 K.7

In the 2 : 3 adduct formed between 1,3,5-trihydroxybenzene and
BPY, pairs of molecules of each component are linked by O–H
hydrogen bonds. The phenolic OH group of the second 4HBA molecule forms an O–H···N hydrogen bond to the carboxyl oxygen of the first 4HBA molecule.

In the 2 : 1 cocrystal a 4HBA molecule is sandwiched between
two BPY molecules and forms two O–H···N hydrogen bonds.
The structural roles of the two hydrogen bonded bipyridine
N-atoms are distinct.11

Nuclear quadrupole resonance (NQR) has proven to be a
sensitive technique for the study of molecular electron configu-
ration in solid materials. In hydrogen bonded solids the change
of the electron configuration produced by hydrogen bonding
causes a shift of the NQR frequencies.12 It has recently been
shown how NQR can be used to investigate cocrystals and
crystal polymorphs.13,14

Two studies of cocrystals of 2-amino-4,6-dimethylpyrimidine
and 2,3,5,6-tetramethylypyrazine on one side and several carb-
onyl acids on the other side have already been performed.15,16

In the present paper we present the results of NQR investiga-
tion of cocrystals of BPY and benzoic acid (BA), 5-chlorosalicylic
acid (5CSA), 4-hydroxybenzoic acid (4HBA), oxalic acid (OXA)
and malonic acid (MA). The structural formulae of the molecules
used in the present study are presented in Fig. 1.

In the compounds studied there are two quadrupole atomic
nuclei, 14N and 17O, which are suitable for the investigation
of hydrogen bonds. Due to experimental reasons we decided to
use 14N NQR to characterize the cocrystals and to investigate
the influence of hydrogen bonding on the electron charge
distribution at the positions of the nitrogen atoms and possible
proton transfer in these systems.

14N NQR

The nucleus of the nitrogen isotope 14N has in its ground state
a spin of f = 1 and a nonzero electric quadrupole moment. The
interaction of the nuclear electric quadrupole moment eQ with
the electric field gradient (EFG) tensor Vjk = ∂2V/∂xk∂xj, at
the position of the atomic nucleus, results in three generally
nonequidistant nuclear quadrupole energy levels. The three
resonance (NQR) frequencies υ+ ≥ υ− ≥ υ0 are expressed as

\[ \nu_\pm = \frac{e^2 qQ}{4\hbar} (3 \pm \eta) \quad \nu_0 = \nu_+ - \nu_- = \frac{e^2 qQ}{2\hbar} \eta \]  

Here, e^2 qQ/\hbar is the quadrupole coupling constant (often abbre-
viated as qcc) and η is the asymmetry parameter of the EFG
tensor. They are related to the principal values \( V_{XX}, V_{YY}, \) and \( V_{ZZ} \)
the EFG tensor and the EFG tensor \( \|V_{ZZ}\| > \|V_{YY}\| > \|V_{XX}\| \) as \( e^2 qQ/\hbar = e Q V_{ZZ}/\hbar \)
and \( \eta = (V_{XX} - V_{YY})/V_{ZZ} \). The quadrupole coupling tensor \( q_{ik} \),
which can be determined by NQR or NMR, is equal to
the product of the EFG tensor and the nuclear quadrupole moment
\( e Q \) divided by the Planck constant \( \hbar \),

\[ q_{ik} = e Q V_{ik}/\hbar \]  

It is expressed in frequency units. The present knowledge of nuclear
electric quadrupole moments17 allows us to calculate the EFG
tensor from the quadrupole coupling tensor. The sign of the
largest principal value \( q_{zz} \) of the quadrupole coupling tensor
can in general not be determined by NQR or NMR, so only
the relative signs of the elements of the quadrupole coupling tensor
with respect to \( q_{zz} \) can be determined by these techniques.
Different experimental techniques, as for example microwave
spectroscopy, or quantum chemical calculations are needed to
determine its sign. The absolute value of the largest principal value of the $^{14}$N quadrupole coupling tensor and the asymmetry parameter $\eta$ are calculated from the $^{14}$N NQR frequencies as $|q_{ZZ}| = e^2 qQ/h = 2(\nu_+ - \nu_-)/3$ and $\eta = 2\nu_0/|q_{ZZ}|$. The two smaller principal values of the quadrupole coupling tensor are related to $q_{ZZ}$ and $\eta$ as $q_{YY} = -q_{ZZ}(1 + \eta)/2$ and $q_{XX} = -q_{ZZ}(1 - \eta)/2$.

### Experimental section

The samples of BPY, benzoic acid, 5-chlorosalicylic acid, 4-hydroxybenzoic acid, oxalic acid and malonic acid were purchased at Sigma-Aldrich and used as obtained. The cocrystals were obtained by mixing hot methanol solutions (~50 °C) of cocrystal formers. The solutions were then left at room temperature for a few days until the cocrystals grew. In the case of BPY–OXA (1:1) the solid immediately precipitated from the solution. The obtained solid is not soluble in methanol. We dissolved a part of the obtained solid in hot water, cooled the solution to room temperature and left it at this temperature for several days until the cocrystals grew.

The $^{14}$N NQR frequencies have been measured by $^1$H–$^{14}$N nuclear quadrupole double resonance (NQDR). In the present study, we have used the solid-effect technique and the technique using multiple frequency sweeps and two-frequency irradiation. These techniques are based on magnetic field cycling. The details of the present experimental setup and the measuring procedure were published in a previous paper.

### Results and discussion

The $^1$H–$^{14}$N NQR spectrum of BPY–BA (1:1) is presented in Fig. 2.

We observe two sets of $^{14}$N NQR frequencies ($\nu_0, \nu_-, \nu_+$): (4020 kHz, 3070 kHz, 950 kHz) and (3250 kHz, 2630 kHz, 620 kHz). The frequencies in the first set are close to the $^{14}$N NQR frequencies observed in solid pyridine and may be assigned to the non-hydrogen bonded nitrogen position. The $^{14}$N NQR frequencies in the second set are significantly lower and may be assigned to the hydrogen-bonded nitrogen atom which is hydrogen bonded to carboxylic OH of benzoic acid.

**Table 1** $^{14}$N NQR frequencies, quadrupole coupling constants $e^2 qQ/h$ and asymmetry parameters $\eta$ in cocrystals of BPY and carboxylic acids

<table>
<thead>
<tr>
<th>Substance</th>
<th>$T$ (K)</th>
<th>Nitrogen position</th>
<th>$\nu_0$ (kHz)</th>
<th>$\nu_-$ (kHz)</th>
<th>$\nu_+$ (kHz)</th>
<th>$e^2 qQ/h$ (kHz)</th>
<th>$\eta$</th>
</tr>
</thead>
<tbody>
<tr>
<td>BPY–BA (1:1)</td>
<td>295</td>
<td>N–H–O</td>
<td>3250</td>
<td>2630</td>
<td>620</td>
<td>3920</td>
<td>0.316</td>
</tr>
<tr>
<td></td>
<td>N</td>
<td>N–H–O</td>
<td>4020</td>
<td>3070</td>
<td>950</td>
<td>4727</td>
<td>0.402</td>
</tr>
<tr>
<td></td>
<td>N</td>
<td>N–H–O</td>
<td>2040</td>
<td>1885</td>
<td>155</td>
<td>2617</td>
<td>0.118</td>
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<tr>
<td></td>
<td>N</td>
<td>N–H–O</td>
<td>3567</td>
<td>2979</td>
<td>888</td>
<td>4564</td>
<td>0.389</td>
</tr>
<tr>
<td></td>
<td>295</td>
<td>N–H–O</td>
<td>2010</td>
<td>1866</td>
<td>144</td>
<td>2584</td>
<td>0.111</td>
</tr>
<tr>
<td></td>
<td>N</td>
<td>N–H–O</td>
<td>3582</td>
<td>2982</td>
<td>900</td>
<td>4576</td>
<td>0.393</td>
</tr>
<tr>
<td>BPY–5CSA (1:1)</td>
<td>228</td>
<td>N–H–O</td>
<td>1981</td>
<td>1843</td>
<td>138</td>
<td>2539</td>
<td>0.108</td>
</tr>
<tr>
<td></td>
<td>N</td>
<td>N–H–O</td>
<td>3910</td>
<td>2989</td>
<td>921</td>
<td>4599</td>
<td>0.400</td>
</tr>
<tr>
<td></td>
<td>N</td>
<td>N–H–O</td>
<td>1967</td>
<td>1842</td>
<td>126</td>
<td>2539</td>
<td>0.099</td>
</tr>
<tr>
<td></td>
<td>N</td>
<td>N–H–O</td>
<td>3921</td>
<td>2995</td>
<td>926</td>
<td>4611</td>
<td>0.402</td>
</tr>
<tr>
<td></td>
<td>157</td>
<td>N–H–O</td>
<td>1957</td>
<td>1833</td>
<td>124</td>
<td>2527</td>
<td>0.098</td>
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<tr>
<td></td>
<td>N</td>
<td>N–H–O</td>
<td>3928</td>
<td>2997</td>
<td>931</td>
<td>4617</td>
<td>0.403</td>
</tr>
<tr>
<td>BPY–OXA (1:1)</td>
<td>179</td>
<td>N–H–O</td>
<td>1050</td>
<td>960</td>
<td>90</td>
<td>1340</td>
<td>0.134</td>
</tr>
<tr>
<td>BPY–MA (1:1)</td>
<td>295</td>
<td>N–H–O</td>
<td>2930</td>
<td>2435</td>
<td>495</td>
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<td>N</td>
<td>N–H–O</td>
<td>3275</td>
<td>2670</td>
<td>605</td>
<td>3961</td>
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<tr>
<td></td>
<td>N</td>
<td>N–H–O</td>
<td>3428</td>
<td>2733</td>
<td>695</td>
<td>4107</td>
<td>0.338</td>
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<tr>
<td></td>
<td>N</td>
<td>N–H–O</td>
<td>3950</td>
<td>3000</td>
<td>950</td>
<td>4633</td>
<td>0.410</td>
</tr>
<tr>
<td>BPY–4HBA (2:1)</td>
<td>181</td>
<td>N–H–O</td>
<td>3393</td>
<td>2715</td>
<td>678</td>
<td>4072</td>
<td>0.333</td>
</tr>
<tr>
<td></td>
<td>N–H–O</td>
<td>N–H–O</td>
<td>3410</td>
<td>2728</td>
<td>682</td>
<td>4092</td>
<td>0.333</td>
</tr>
</tbody>
</table>

The possible hydrogen bonding schemes of a BPY molecule are presented in Fig. 3.

Scheme (a) is present in BPY–BA (1:1). Within the experimental resolution of NQDR it may be also concluded that all BPY molecules are crystallographically equivalent.

A similar situation is observed in BPY–5CSA (1:1), where we at room temperature observe two sets of $^{14}$N NQR frequencies: (3882 kHz, 2982 kHz, 900 kHz) and (2010 kHz, 1866 kHz, 144 kHz). The $^{14}$N NQR frequencies in the first set are characteristic for a non-hydrogen bonded nitrogen position. The $^{14}$N NQR frequencies in the second set are rather low. They are in the range of $^{14}$N NQR frequencies observed in pyridine-3,5-dicarboxylic acid, where short, strong N–H–O hydrogen bonds are formed. In this system, by varying temperature, phonon driven proton transfer occurs. At low temperature proton is closer to the hydrogen bond acceptor than to the hydrogen bond donor (N–H–O). Upon increasing temperature proton is continuously displaced from the hydrogen bond acceptor to the hydrogen bond donor (N–H–O → N–H–O). The proton displacement results in a strong temperature variation of the $^{14}$N NQR frequencies. Due to the possibility that a similar effect also occurs in BPY–5CSA (1:1) we...
T relaxation time values are presented in Table 1.

Hydrogen bond donor produces lower $^{14}$N NQR frequencies. According to the crystal structure\textsuperscript{12} we expect to observe four NQR frequencies: (3950 kHz, 3000 kHz, 950 kHz), (3428 kHz, 2733 kHz, 695 kHz), and (3275 kHz, 2670 kHz, 605 kHz). According to the crystal structure\textsuperscript{12} we expect to observe four sets of $^{14}$N NQR frequencies: two from the hydrogen bonded nitrogen positions and two from the non-hydrogen bonded nitrogen positions (two nonequivalent BPY molecules with the hydrogen bond Scheme (a)). We observed only one non-hydrogen bonded position (the first set of NQR frequencies) so we suppose that the splitting of the $^{14}$N NQR frequencies from the two non-hydrogen bonded nitrogen positions is smaller than the experimental resolution of NQDR ($\sim 10$ kHz). In our previous studies we also observed that a lower $pK_a$ of the hydrogen bond donor produces lower $^{14}$N NQR frequencies. We therefore assign the second set of $^{14}$N NQR frequencies to the nitrogen atom which is hydrogen bonded to the phenolic OH group and the third set of $^{14}$N NQR frequencies to the nitrogen atom which is hydrogen bonded to the carboxylic OH group.

In BPY–4HBA (2:1) we performed the measurements at $T = 181$ K due to experimental reasons. Proton spin–lattice relaxation time $T_1$ at the Larmor frequency lower than 1 MHz is too short at room temperature for the present experimental setup to be used. We therefore cooled the sample to a lower temperature to obtain a longer $T_1$. We observed three sets of $^{14}$N NQR frequencies: (3950 kHz, 3000 kHz, 950 kHz), (3428 kHz, 2733 kHz, 695 kHz), and (3275 kHz, 2670 kHz, 605 kHz). According to the crystal structure\textsuperscript{12} we expect to observe four sets of $^{14}$N NQR frequencies: two from the hydrogen bonded nitrogen positions and two from the non-hydrogen bonded nitrogen positions (two nonequivalent BPY molecules with the hydrogen bond Scheme (a)). We observed only one non-hydrogen bonded position (the first set of NQR frequencies) so we suppose that the splitting of the $^{14}$N NQR frequencies from the two non-hydrogen bonded nitrogen positions is smaller than the experimental resolution of NQDR ($\sim 10$ kHz). In our previous studies we also observed that a lower $pK_a$ of the hydrogen bond donor produces lower $^{14}$N NQR frequencies. We therefore assign the second set of $^{14}$N NQR frequencies to the nitrogen atom which is hydrogen bonded to the phenolic OH group and the third set of $^{14}$N NQR frequencies to the nitrogen atom which is hydrogen bonded to the carboxylic OH group.

In BPY–4HBA (2:1) we observe two slightly different sets of $^{14}$N NQR frequencies: (3410 kHz, 2728 kHz, 682 kHz) and (3393 kHz, 2715 kHz, 678 kHz). They are both approximately equal to the $^{14}$N NQR frequencies in BPY–4HBA (2:1) at the nitrogen position which is hydrogen bonded to the phenolic OH group. This situation is expected in polymorphic Form 1 with the hydrogen bonding Scheme (b). The polymorphic Form 2, where we expect two strongly different sets of $^{14}$N NQR frequencies, has not been observed within experimental accuracy. Most probably the mass fraction of this form in the polycrystalline sample is low, lower than approximately 20%.

Two slightly different sets of $^{14}$N NQR frequencies in this compound show that there are two non-equivalent nitrogen positions in the unit cell. This is possible either if the two nitrogen positions in a BPY molecule are distinct or if there are two distinct centrosymmetric BPY molecules in the unit cell.

In BPY–OXA (1:1) we performed the measurements at $T = 181$ K due to experimental reasons. Proton spin–lattice relaxation time $T_1$ at the Larmor frequency lower than 1 MHz is too short at room temperature for the present experimental setup to be used. We therefore cooled the sample to a lower temperature to obtain a longer $T_1$. We observed three sets of $^{14}$N NQR frequencies: (3950 kHz, 3000 kHz, 950 kHz), (3428 kHz, 2733 kHz, 695 kHz), and (3275 kHz, 2670 kHz, 605 kHz). According to the crystal structure\textsuperscript{12} we expect to observe four sets of $^{14}$N NQR frequencies: two from the hydrogen bonded nitrogen positions and two from the non-hydrogen bonded nitrogen positions (two nonequivalent BPY molecules with the hydrogen bond Scheme (a)). We observed only one non-hydrogen bonded position (the first set of NQR frequencies) so we suppose that the splitting of the $^{14}$N NQR frequencies from the two non-hydrogen bonded nitrogen positions is smaller than the experimental resolution of NQDR ($\sim 10$ kHz). In our previous studies we also observed that a lower $pK_a$ of the hydrogen bond donor produces lower $^{14}$N NQR frequencies. We therefore assign the second set of $^{14}$N NQR frequencies to the nitrogen atom which is hydrogen bonded to the phenolic OH group and the third set of $^{14}$N NQR frequencies to the nitrogen atom which is hydrogen bonded to the carboxylic OH group.

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π electron orbital. The two effects are correlated in the case of pyridine and pyrazine.

In substituted and hydrogen-bonded pyrimidine, we analyzed the 14N quadrupole coupling tensor in terms of the population of the π-electron orbital and deformation of the lone pair electron orbital. In the same model we can express the 14N quadrupole coupling tensor q in hydrogen bonded BPY as

\[
q = \begin{pmatrix}
1.434 & 0 & 0 \\
0 & 3.474 & 0 \\
0 & 0 & -4.908
\end{pmatrix} \text{MHz} + x_{LP} \begin{pmatrix}
-0.5 & 0 & 0 \\
0 & -0.5 & 0 \\
0 & 0 & 1
\end{pmatrix}
\]

\[
+ x_\pi \begin{pmatrix}
-0.5 & 0 & 0 \\
0 & 1 & 0 \\
0 & 0 & -0.5
\end{pmatrix}
\]

Here the first tensor contains the principal values of the 14N quadrupole coupling tensor of pyridine in the gas phase, x_{LP} is a parameter related to the deformation of the lone pair electron orbital and x_\pi is proportional to the change of the population of the π-electron orbital with respect to an isolated pyridine molecule. Expressions (2) are obtained when x_{LP} = -1.64x_\pi. The principal values of the quadrupole coupling tensor are then expressed in terms of the parameter x_\pi in the following way:

\[
q_{xx} = 1.434 \text{ MHz} + 0.32x_\pi
\]
\[
q_{yy} = 3.474 \text{ MHz} + 1.82x_\pi
\]
\[
q_{zz} = -4.908 \text{ MHz} - 2.14x_\pi
\]

In the case of a full proton transfer (pyridinium ion) q_{zz} is approximately zero. This corresponds to x_\pi = -2.3 MHz. If we assume that the largest principal value of the axially symmetric contribution of one electron in the nitrogen 2p orbital to the quadrupole coupling tensor is -9 MHz we obtain from the NQR data an increase of the population of the nitrogen π-electron orbital equal to (-2.3 MHz)/(-9 MHz) = 0.25.

The data for the hydrogen bonded nitrogen position in BPY–5CSA are approximately in the center of the correlation diagram. This position indicates the existence of a short, strong N – H – O hydrogen bond. In pyridine-3,5-dicarboxylic acid, where a strong short N – H – O hydrogen bond is formed, proton continuously migrates from the N +–H position at low temperature to the N – H – O position at high temperature. The NQR measurements show a continuous temperature variation of the 14N quadrupole coupling constant $e^2Q/h$. The ratio $\Delta R(N – H)/\Delta e^2Q/h$ is equal to 0.36 Å MHz⁻¹. Here $\Delta R(N – H)$ is the change of the N – H distance and $\Delta e^2Q/h$ is the corresponding change of the 14N quadrupole coupling constant. The 14N NQR frequencies in BPY–5CSA are very close to the 14N NQR frequencies in pyridine-3,5-dicarboxylic acid so we assume that the ratio $\Delta R(N – H)/\Delta e^2Q/h$ is the same in both compounds. Upon increasing the temperature from 157 K to 323 K the 14N quadrupole coupling constant increases for $\Delta e^2Q/h = 90$ kHz, so the nitrogen–proton distance $R(N – H)$ increases for 0.03 Å. The actual increase of $R(N – H)$ may be somewhat larger, because there are two competing effects producing the temperature variation of $e^2Q/h$ at this nitrogen position. Proton displacement towards oxygen increases $e^2Q/h$, whereas molecular thermal motion, which amplitude increases with increasing temperature, produces a decrease of $e^2Q/h$ when temperature increases. If we suppose that the proton displacement dominates the temperature variation of $e^2Q/h$ we may conclude that in BPY–5CSA the proton displacement is in the temperature interval between 157 K and 323 K about four times smaller than in pyridine-3,5-dicarboxylic acid.

To check whether in this compound the proton displacement at the hydrogen bonded nitrogen position influences the electron charge distribution at the non-hydrogen bonded nitrogen position in BPY and to get some more information on thermal motions, we plot the temperature variation of q_{yy} versus q_{zz} for both nitrogen positions. The plot is presented in Fig. 5. The correlation line (expression 2) is also shown in the diagram.

At the hydrogen bonded nitrogen position q_{yy} and q_{yy} increase in magnitude with increasing temperature. The experimental points are located on a line, which is nearly parallel to the correlation line. The distance between the two lines is small. This demonstrates that the temperature variation of the hydrogen bond parameters dominates the temperature variation of the 14N quadrupole coupling tensor at this nitrogen position.

At the non-hydrogen bonded nitrogen position q_{yy} and q_{yy} decrease in magnitude with increasing temperature. The change of $e^2Q/h$ at the non-hydrogen bonded nitrogen position is equal to 60% of the change of $e^2Q/h$ at the hydrogen bonded nitrogen position.
position in the same temperature interval. The line through the experimental points is not parallel to the correlation line. This may be the effect of thermal motions. We may assume that the dominant thermal motion of a BPY molecule is liberation of pyridine rings around the long molecular axis. This motion does not change $q_{zz}$. The principal axis $Z$ is parallel to the long molecular axis. The liberation only reduces the difference between $q_{yy}$ and $q_{xx}$.

We define the molecule-fixed coordinate system $(X, Y, Z)$ along the principal directions of the $^{14}$N quadrupole coupling tensor for a static molecule. In addition we define the crystal-fixed coordinate system $(x, y, z = Z)$ in such a way that the molecular axes $X$ and $Y$ are time average directed along the crystal-fixed axes $x$ and $y$ respectively. The liberation around the $z$ axis produces time variation of $q_{xx}$, $q_{yy}$ and $q_{yx}$. The principal values of the time-averaged quadrupole coupling tensor $(q_i)$, that are observed by NQR, are in the case of small librations equal to

$$q_{zz} = q_{zz}$$

$$q_{yy} = q_{yy} - (q_{yy} - q_{xx})(\phi^2)$$

$$q_{xx} = q_{xx} + (q_{yy} - q_{xx})(\phi^2)$$

(5)

Here $(\phi^2)$ is the mean square libration angle. For the non-hydrogen bonded nitrogen position the difference $q_{yy} - q_{xx}$ is approximately equal to 1.8 MHz, whereas for the hydrogen bonded nitrogen position it is equal to 0.3 MHz. The line through the experimental points at the non-hydrogen bonded nitrogen position becomes parallel to the correlation line when we add the librations in such a way that the following relation holds

$$(q_{yy} - q_{xx})[(\phi^2)_{295 K} - (\phi^2)_{157 K}] = 20 \text{ kHz}$$

(6)

The mean square of the libration angle $(\phi^2)$ is usually proportional to the absolute temperature. If the same holds in the present compound we calculate from expression (6) $(\phi^2)_{295 K}$ as being equal to 0.022 rad$^2$. Its square root is equal to 0.15 rad = 8.5°. Librations with the same amplitude at the hydrogen bonded nitrogen position produce about six times lower effect (0.3 MHz/1.8 MHz = 1/6; 20 kHz/6 MHz = 1/300 kHz) than at the non-hydrogen bonded nitrogen position. This can be the reason why they are not observed at this nitrogen position.

Upon increasing the temperature from 157 K to 323 K the $^{14}$N quadrupole coupling constant at the hydrogen bonded nitrogen position increases for 90 kHz ($\Delta q_{zz} = 90$ kHz), whereas at the non-hydrogen bonded nitrogen position it decreases for 53 kHz ($\Delta q_{zz} = 53$ kHz). According to expression (4) at this temperature change the population of the nitrogen $\pi$-electron orbital at the hydrogen bonded nitrogen positions decreases for 0.005 whereas at the non-hydrogen bonded nitrogen position it increases for 0.003.

In cocrystals of 2,3,5,6-tetramethylpyrazine we also observed a correlation between the $^{14}$N quadrupole coupling constants at the hydrogen bonded and non-hydrogen bonded nitrogen positions. A lower quadrupole coupling constant at the hydrogen bonded nitrogen position results in a higher quadrupole coupling constant at the non-hydrogen bonded nitrogen position. Such a correlation is not observed in cocrystals of BPY.

Conclusions

Cocrystals BPY–BA (1 : 1), BPY–5CSA (1 : 1), BPY–OXA (1 : 1), BPY–MA (1 : 1), BPY–4HBA (2 : 1), and BPY–4HBA (1 : 2) were grown from the methanol solution of the cocrystal formers. The cocrystals BPY–OXA (1 : 1) were also recrystallized from the water solution.

Complete $^{14}$N NQR spectra of these cocrystals have been measured by $^1$H–$^{14}$N nuclear quadrupole double resonance. The principal values of the quadrupole coupling tensor are calculated from the $^{14}$N NQR frequencies. A large variation of $^{14}$N quadrupole coupling constant between 1.3 MHz and 4.7 MHz is observed. At the non-hydrogen bonded nitrogen position in a BPY molecule the $^{14}$N quadrupole coupling constant is around 4.6 MHz. An increasing strength of the O–H–N hydrogen bond reduces the $^{14}$N quadrupole coupling constant. In the case of weak carboxylic acids BA and 4HBA, the $^{14}$N quadrupole coupling constant is reduced to approximately 4 MHz. Polymorphism in BPY–4HBA (1 : 2) has not been observed. A somewhat stronger MA, which forms two equivalent hydrogen bonds with two BPY molecules, reduces the $^{14}$N quadrupole coupling constant to about 3.6 MHz. A still stronger 5CSA reduces $^{14}$N quadrupole coupling constant to about 2.5 MHz. A very low $^{14}$N quadrupole coupling constant, about 1.3 MHz, is observed in BPY–OXA (1 : 1). Such a low $^{14}$N quadrupole coupling constant is characteristic for proton transfer (O–H…N $\rightarrow$ O’–H–N’).

A correlation of the principal values of the $^{14}$N quadrupole coupling tensor is observed in the whole range from the non-interacting pyridine in the gas phase through the presently obtained data to the pyrimidinium ion. The correlation is analyzed in the model, where the deformation of the lone pair electron orbital and the change of the population of the $\pi$-electron orbital produce the variation of the $^{14}$N quadrupole coupling tensor in hydrogen bonded BPY.

The value of the $^{14}$N quadrupole coupling constant at the hydrogen bonded nitrogen position in BPY–5CSA (1 : 1) is in midway between the value observed at the non-hydrogen bonded nitrogen position (~4.6 MHz) and the value observed in the case of the pyrimidinium ion (~1 MHz). Such an intermediate value of the $^{14}$N quadrupole coupling constant is characteristic for a short, strong hydrogen bond. To check for the temperature variation of the proton position within the short, strong hydrogen bond we measured the $^{14}$N NQR frequencies at various temperatures between 157 K and 323 K. The NQR data show that the proton migrates upon increasing the temperature in the direction from the hydrogen bond acceptor to the hydrogen bond donor. The proton displacement is in the temperature range between 157 K and 323 K equal to 0.03 Å.

From the temperature variation of the principal values of the $^{14}$N quadrupole coupling tensor at the two nitrogen positions in
BPY–5CSA (1 : 1) we estimated the mean-square libration angle of the non-hydrogen bonded pyridine ring at room temperature and the change of the population of the $\pi$-electron orbital in the temperature interval between 157 K and 323 K at the two nitrogen positions in a BPY molecule.

References