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# Exploring short strong hydrogen bonds engineered in organic acid molecular crystals for temperature dependent proton migration behaviour using single crystal synchrotron X-ray diffraction (SCSXR D)<sup>†</sup>

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Seven multi-component molecular crystals containing O–H $\cdots$ O/O<sup>+</sup>–H $\cdots$ O<sup>–</sup> and N<sup>+</sup>–H $\cdots$ O<sup>–</sup> short strong hydrogen bonds (SSHBs) have been engineered by combining substituted organic acids with hydrogen bond acceptor molecules *N,N*-dimethylurea and isonicotinamide. In these materials, the shortest of the SSHBs are formed in the *N,N*-dimethylurea set for the *ortho/para* nitro-substituted organic acids whilst a twisted molecular approach favours the shorter SSHBs N<sup>+</sup>–H $\cdots$ O<sup>–</sup> in the isonicotinamide set. Temperature dependent proton migration behaviour has been explored in these systems using single crystal synchrotron X-ray diffraction (SCSXR D). By using a protocol which considers a combination of structural information when assessing the hydrogen atom (H-atom) behaviour, including refined H-atom positions alongside heavy atom geometry and Fourier difference maps, temperature dependent proton migration is indicated in two complexes (2: *N,N*-dimethylurea 2,4-dinitrobenzoic acid 1:1 and 5: isonicotinamide phthalic acid 2:1). We also implement Hirshfeld atom refinement for further confidence in this observation; this highlights the importance of having corroborating trends when applying the SCSXR D technique in these studies. Further insights into the SSHB donor–acceptor distance limit for temperature dependent proton migration are also revealed. For the O–H $\cdots$ O/O<sup>+</sup>–H $\cdots$ O<sup>–</sup> SSHBs, the systems here support the previously proposed maximum limit of 2.45 Å whilst for the charge assisted N<sup>+</sup>–H $\cdots$ O<sup>–</sup> SSHBs, a limit in the region of 2.55 Å may be suggested.

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## Introduction

Short strong hydrogen bonds (SSHBs) are those with very short donor–acceptor distances, typically less than 2.5 Å for O–H $\cdots$ O HBs<sup>1,2</sup> and less than 2.6 Å for N–H $\cdots$ O HBs.<sup>3</sup> These interactions may be partly covalent in character<sup>4,5</sup> and have a potential energy surface (PES) which approaches the shape of

a symmetric single well potential with a diminished barrier to hydrogen atom (H-atom) motion.<sup>6</sup> Proton transfer, in the form of migration, may occur in the presence of this ‘low’ barrier, involving a gradual shift/movement in H-atom position across the HB; this effect often occurs under an external stimulus including the application of an electric field,<sup>7,8</sup> or changes in pressure<sup>9</sup> or temperature.<sup>10</sup>

Proton migration across SSHBs can lead to a number of potentially useful properties in solid-state materials including ferroelectric<sup>11</sup> or colour changing<sup>12</sup> behaviour. Its study can reveal insight into the occurrence of such properties and have implications in functional materials design. A temperature dependency of this behaviour has been observed across SSHBs in a number of solid-state organic materials. These include across O–H $\cdots$ O SSHBs in substituted urea:organic acid molecular complexes<sup>10,13</sup> and across charge assisted N<sup>+</sup>–H $\cdots$ O<sup>–</sup> SSHBs in N-heterocycle:organic acid molecular complexes.<sup>14–16</sup> For the urea:organic acid systems, the SSHB donor–acceptor distance is found to ‘tune’ the migration

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<sup>†</sup> Electronic supplementary information (ESI) available: Crystallographic data; selected molecular geometry tables; selected hydrogen bond tables/plots; selected Fourier difference electron density maps. CCDC 1921985–1922038. For ESI and crystallographic data in CIF or other electronic format see DOI: 10.1039/c9ce00925f



behaviour; an O–H...O donor–acceptor distance of 2.45 Å has been proposed as a limit for migration, above which proton migration behaviour would not be expected.<sup>10,13</sup> No such limit has been proposed for charge assisted N<sup>+</sup>–H...O<sup>−</sup> SSHBs. However, temperature dependent proton migration is reported only for molecular complexes where the donor–acceptor distance is less than 2.55 Å.<sup>14–16</sup>

Solid-state proton transfer processes across HBs, including temperature dependent proton migration, are typically followed using neutron diffraction; a method which allows the unambiguous determination of H-atom positions and, often important in such studies, also reveals H-atom anisotropic displacement parameters (ADPs).<sup>17</sup> Other experimental methods such as <sup>14</sup>NQR (nuclear quadrupole resonance) spectroscopy,<sup>18</sup> X-ray photoelectron spectroscopy<sup>19</sup> and solid-state NMR<sup>20,21</sup> combined with computational methods<sup>22,23</sup> can offer alternatives to diffraction methods in determining H-atom positions. Recently, X-ray diffraction has been shown to be useful in such studies, notwithstanding the fact that the precision of the determination of H-atom parameters is lower.<sup>24–27</sup> Hirshfeld atom refinement (HAR)<sup>27</sup> can contribute to overcoming such deficiencies allowing the precise and accurate location of H-atoms, comparable to neutron determinations,<sup>26,28</sup> from X-ray diffraction data.

The present work explores SSHBs engineered in multi-component organic acid : *N,N*-dimethylurea/isonicotinamide molecular crystals for temperature dependent proton migration behaviour. These systems have arisen from a wider study into the SSHB propensity of organic co-formers.<sup>29</sup> *N,N*-Dimethylurea and isonicotinamide molecules readily recognise carboxylic acid groups to form robust heterodimeric hydrogen bonded motifs<sup>30,31</sup> across which migration behaviour has been observed previously.<sup>10,13–16</sup> The selected organic acid co-formers (Fig. 1) in this study include *ortho* or nitro-substituted benzoic acids targeted for their increased acidity<sup>32</sup> with the potential to form HBs with a charge-assisted component that can impart additional strength to the interaction,<sup>33</sup> increasing the likelihood of exhibiting migration behaviour.

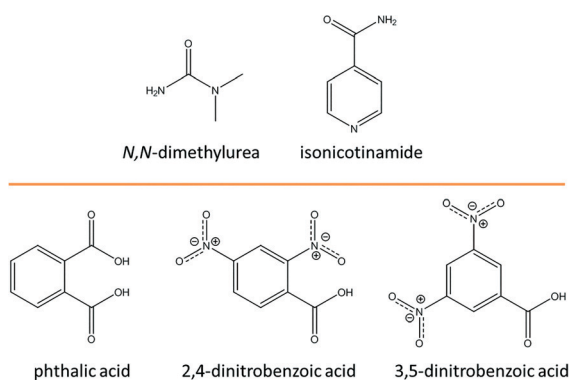


Fig. 1 The *N,N*-dimethylurea/isonicotinamide components and organic acid co-formers.

Single crystal synchrotron X-ray diffraction (SCSXR) has been implemented here to explore the temperature dependent proton migration behaviour of the SSHBs in the prepared systems; nearly all of these form with SSHB donor–acceptor distances approaching the region where this behaviour has been previously reported.<sup>13,15,16</sup> This technique is shown to be valid in identifying such behaviour by using a protocol which assesses potential migration behaviour across a number of parameters and which additionally implements HAR. This combined approach allows greater confidence in any behaviour observed. The use of SCSXR in this context is advantageous because the high intensity of the synchrotron X-rays allows for rapid data collection enabling multiple measurements under different conditions or of different complexes to be carried out faster than a single laboratory XRD or neutron experiment.

## Experimental

### Complex preparation

All single crystal samples were prepared by evaporation from solution. Single crystals of 1 (*N,N*-dimethylurea phthalic acid 2:1) and 5 (isonicotinamide phthalic acid 2:1) were grown from a 2:1 stoichiometric ratio of the two components, respectively, whilst 2 (*N,N*-dimethylurea 2,4-dinitrobenzoic acid 1:1), 3 (*N,N*-dimethylurea 3,5-dinitrobenzoic acid 1:1), 4 (*N,N*-dimethylurea 3,5-dinitrobenzoic acid 2:2), 6 (isonicotinamide 2,4-dinitrobenzoic acid 1:1), 7 (isonicotinamide 2,4-dinitrobenzoic acid 2:2) and 8 (isonicotinamide 3,5-dinitrobenzoic acid 1:1) were grown from a 1:1 stoichiometric ratio of the two components. Crystallisations were carried out in methanol solvent for 2 (at 40 °C), 5 (30 °C), 7 (at room temperature) or in ethanol for 1, 3, 6 (at room temperature) and 8 (at 35 °C).

### Data collection and treatment

SCSXR data were collected from 2, 4, 6 and 8 on beamline I19 (EH1) at Diamond Light Source, U.K. (NR18193-1) at  $\lambda = 0.6889$  Å using a Fluid Film Devices Ltd diffractometer equipped with a PILATUS 2M detector. The sample temperature was controlled (between 100 and 350 K) using an Oxford Cryosystems Cryostream Plus. Data were collected using the in-house General Data Acquisition (GDA) software and processed using in-house *xia2* software.<sup>34</sup> For 8, data were additionally scaled using the Bruker software SADABS.<sup>35</sup> SCSXR data were collected from 1, 3, 5 and 7 on beamline 11.3.1 at the Advanced Light Source, U.S.A. using Si (111) monochromated radiation at  $\lambda = 0.7749$  Å (1) and  $\lambda = 0.8856$  Å (3, 5 and 7). SCSXR data were collected from 1 using a Bruker AXS D8 three-circle diffractometer equipped with a Bruker AXS APEX2 CCD detector. SCSXR data were collected from 3, 5 and 7 using a Bruker AXS D8 three-circle diffractometer equipped with a Bruker AXS PHOTON 100 CMOS detector. The sample temperature was controlled for collections between 100 and 350 K using an Oxford Cryosystems Cryostream Plus. Data were collected and processed using



Bruker AXS APEX2<sup>36</sup> and Bruker software SADABS<sup>35</sup> versions 2014/2 and 2014/15.

### Structure solution and refinement details

All 100 K structures of 1–8 were solved using SHELXS-2018<sup>37</sup> and refined using SHELXL-2018<sup>38</sup> within the WinGX package.<sup>39</sup> For successive temperatures, the 100 K structure was used as the starting point for the SHELXL refinements. H-Atoms were nearly all located from Fourier difference maps and their positions and isotropic thermal parameters allowed to refine freely. Exceptions to this are in 1–4 where disorder is indicated in the *N,N*-dimethylurea methyl groups at elevated temperatures and so the methyl H-atoms were placed geometrically using HFIX 137 at these temperatures. A SHEL command was used for each system set to the maximum diffraction resolution of the highest temperature data set (cut-off criteria where  $I/\sigma \geq 1-2$ ,  $R_{\text{int}} \leq 50\%$ ) allowing comparability of multi-temperature data sets. Crystallographic data for 1–8 are given in ESI† Tables S1 to S8. HAR was performed for systems 2, 5 and 7 that exhibited the most interesting proton migration behaviour from the SHELXL refinements. Data were first merged in Bruker software XPREF<sup>40</sup> followed by crystal structure refinement using SHELXL. On finalising the structure, a LIST 4 FVAR<sup>2</sup> command was used and the output  $F_{\text{obs}}$  structure factors and finalised CIF were used as the input to HAR. The programme HAR (16.05.31 v. c4e213d) was run on the Linux platform (Linux-4.13.0-41-generic) on the Diamond Light Source (U.K.) cluster using the selected refinement options: shelx-f2 (reflection file  $F^2$  format), dispersion details for X-ray wavelength, basis-set using def2-TZVP level of theory, anisotropic or isotropic refinement of H-atom ADPs (depending on refinement stability) and the use of a restricted Hartree–Fock self-consistent field. The crystal fragment used was the asymmetric unit. The refinements took between 5 to 48 hours depending on complexity of asymmetric unit. The refinement was deemed to be successful when run to convergence with values of  $\chi^2 < 10$  and maximum shift values refined to 3 decimal places. HAR refinement data are included in the ESI† Tables S2, S5 and S7.

## Results and discussion

### SSHB formation and proton transfer

$\text{O}-\text{H}\cdots\text{O}/\text{O}^+-\text{H}\cdots\text{O}^-$  SSHBs. The crystallisation of *N,N*-dimethylurea with the organic acid co-formers generates molecular crystals 1 to 4. These include a 2:1 system of *N,N*-dimethylurea and phthalic acid (1) respectively, a 1:1 adduct of *N,N*-dimethylurea and 2,4-dinitrobenzoic acid (2), a 1:1 adduct of *N,N*-dimethylurea 3,5-dinitrobenzoic acid (3) and a 2:2 system of *N,N*-dimethylurea and 3,5-dinitrobenzoic acid (4). 3 and 4 are non-polymorphic forms crystallising with the same component stoichiometry (1:1) but differing values of  $Z'$  (3  $Z' = 1$  versus 4  $Z' = 2$ ). In the crystal structures of 1–4, molecular association of the two components occurs *via* a carboxylic acid:amide  $R_2^2(8)$  HB synthon (Fig. 2) in which the

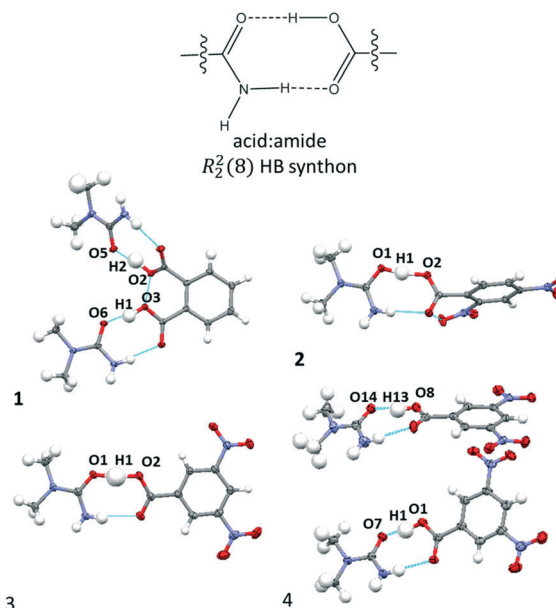


Fig. 2 The carboxylic acid:amide  $R_2^2(8)$  hydrogen bonded synthons formed in the crystal structures of 1–4.

$\text{O}-\text{H}\cdots\text{O}$  HB dominates synthon formation.<sup>41</sup> This synthon is commonly formed between substituted urea and organic acid components;<sup>42,43</sup> its frequency of formation is consistent with complex stoichiometry, as noted by Alhalaweh *et al.* (2010).<sup>44</sup>

The donor–acceptor distances of the  $\text{O}-\text{H}\cdots\text{O}$  HBs in 1–4 at 100 K (Table 1) are below 2.5 Å and can therefore be classed as short strong hydrogen bonds.<sup>45</sup> In most cases, the donor–acceptor distances are within the range of the previously proposed limit for temperature dependent proton migration across  $\text{O}-\text{H}\cdots\text{O}$  SSHBs ( $\sim 2.45$  Å)<sup>13</sup> and are good candidates for multi-temperature studies. The shortest HBs are formed in 2 and 4 having donor–acceptor distances in the region of 2.44 Å, just below the proposed migration limit. The complementary  $\text{N}-\text{H}\cdots\text{O}$  HB in the  $R_2^2(8)$  HB synthon mirrors this interaction and tends to be shorter if the  $\text{O}-\text{H}\cdots\text{O}$  HB is shorter.

The  $\text{O}-\text{H}\cdots\text{O}$  SSHBs form with varying degrees of proton transfer in the 100 K structures. In 1 and 4, no proton transfer is evident as the carboxylic acid proton resides on the acid molecule; 1 and 4 are therefore neutral complexes at 100 K and considered co-crystals.<sup>46</sup> For 2 and 3, the  $\text{O}-\text{H}\cdots\text{O}$  SSHBs are charge assisted ( $\text{O}^+-\text{H}\cdots\text{O}^-$ ) as a result of the H-atom being fully (2) or partially (3) transferred between the organic acid and *N,N*-dimethylurea molecule. 2 is therefore a salt at this temperature<sup>46</sup> whilst the refined H-atom positions at 100 K for 3 indicate a centralised position (the  $d_{\text{O}-\text{H}}$  1.23(7) Å and  $d_{\text{H}\cdots\text{O}}$  1.26(7) Å distances are not significantly different given the esds) meaning it is an intermediate between a salt and co-crystal.<sup>47</sup>

Generally, where the interaction has a charge assisted component ( $\text{O}^+-\text{H}\cdots\text{O}^-$ ), as in 2 and 3, the donor–acceptor distances are shorter. This can be explained where the charges on the donor–acceptor atoms significantly increase



**Table 1** Hydrogen bond parameters of the carboxylic acid :amide  $R_2^2(8)$  HB synthon in 1–4: D is the HB donor atom and A is the HB acceptor atom

| System | Components   | Interaction         | $dD-H$ (Å) | $H\cdots A$ (Å) | $dD\cdots A$ (Å) | $\angle DHA$ (°) |
|--------|--|---------------------|------------|-----------------|------------------|------------------|
| 1      | 2 : 1 <i>N,N</i> -dimethylurea phthalic acid           | O2–H2 $\cdots$ O5   | 1.01(3)    | 1.46(3)         | 2.4520(16)       | 167(2)           |
|        |  | N2–H14 $\cdots$ O1  | 0.95(2)    | 2.03(2)         | 2.9743(19)       | 170.1(18)        |
|        |  | O3–H1 $\cdots$ O6   | 1.09(3)    | 1.44(3)         | 2.4925(16)       | 160(3)           |
|        |  | N4–H21 $\cdots$ O4  | 0.93(2)    | 2.06(2)         | 2.9778(19)       | 174(2)           |
| 2      | 1 : 1 <i>N,N</i> -dimethylurea 2,4-dinitrobenzoic acid | O1–H1 $\cdots$ O2   | 1.10(2)    | 1.36(2)         | 2.4435(10)       | 168.6(18)        |
|        |  | N1–H2 $\cdots$ O3   | 0.884(16)  | 2.030(16)       | 2.9100(12)       | 173.1(14)        |
| 3      | 1 : 1 <i>N,N</i> -dimethylurea 3,5-dinitrobenzoic acid | O1–H1 $\cdots$ O2   | 1.22(7)    | 1.26(7)         | 2.469(3)         | 172(6)           |
|        |  | N4–H12 $\cdots$ O3  | 0.88(4)    | 2.11(5)         | 2.964(3)         | 162(3)           |
| 4      | 2 : 2 <i>N,N</i> -dimethylurea 3,5-dinitrobenzoic acid | O1–H1 $\cdots$ O7   | 1.06(2)    | 1.39(2)         | 2.4445(12)       | 170(2)           |
|        |  | N3–H5 $\cdots$ O2   | 0.890(19)  | 1.99(2)         | 2.8795(14)       | 173.9(17)        |
|        |  | O8–H13 $\cdots$ O14 | 1.03(2)    | 1.45(2)         | 2.4640(11)       | 167(2)           |
|        |  | N7–H17 $\cdots$ O9  | 0.905(18)  | 2.041(18)       | 2.9272(14)       | 166.0(16)        |

the strength of the interaction<sup>33</sup> by reinforcing its electrostatic character;<sup>48,49</sup> leading to the shorter donor–acceptor distances. Furthermore, where the charge assisted component of the interaction is enhanced by the organic acid having substituents that are either electron withdrawing<sup>50</sup> (2–4) or in the *ortho/para* position (2),<sup>32,51,52</sup> the donor–acceptor distances are shorter; these substituents increase the acidity of the carboxylic acid group and impart a greater charge assisted component to the SSHB. These observations are useful in targeting short O–H $\cdots$ O SSHBs in the future. The shorter O–H $\cdots$ O donor–acceptor distances also form where the DHA angle approaches *ca.* 170° suggesting this may be an optimum angle to facilitate SSHB formation for this set of materials.

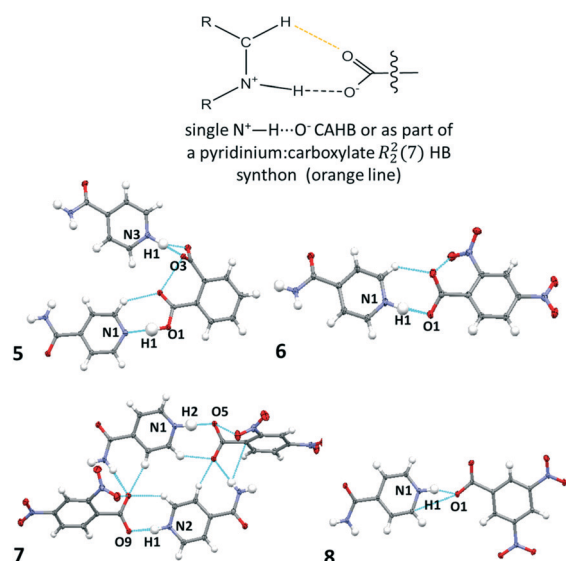
**$N^+H\cdots O^-$  SSHBs.** The crystallisation of isonicotinamide with the organic acid co-formers generates four molecular crystals, 5–8. These include a 2 : 1 system of isonicotinamide and phthalic acid respectively (5), a 1 : 1 system of isonicotinamide and 2,4-dinitrobenzoic acid (6), a 2 : 2 system of isonicotinamide and 2,4-dinitrobenzoic acid (7) and a 1 : 1 system of isonicotinamide and 3,5-dinitrobenzoic acid (8). 6 and 7 are non-polymorphic forms crystallising with the same component stoichiometry (1 : 1) but differing values of  $Z'$  (6  $Z' = 1$  versus 7  $Z' = 2$ ). The crystal structure of 8 has been reported previously and is included for comparison with this related set of structures.<sup>53</sup> In the crystal structures of 5–8, molecular association of the two components occurs *via* charge assisted  $N^+H\cdots O^-$  HBs. For 6 and 7, the  $N^+H\cdots O^-$  interaction is assisted by a weak C–H $\cdots$ O HB to give a pyridinium:carboxylate  $R_2^2(7)$  hydrogen bond synthon (Fig. 3). This synthon is commonly reported in pyridine carboxylic acid molecular complexes<sup>54</sup> and can contribute to favourable mechanical properties or plasticity in materials.<sup>41</sup> Two symmetry independent pyridinium:carboxylate  $R_2^2(7)$  HB synthons form in 7 consistent with complex stoichiometry (2 : 2).

The refined H-atom positions indicate that 5–8 are all salts at 100 K; an acidic proton has been transferred to the isonicotinamide molecule in each case. In 5, both phthalic acid carboxylic acid groups form HBs to a different isonicotinamide molecule; only one of these interactions is

charge assisted, the other is a neutral O–H $\cdots$ N HB which is moderate in strength ( $d_{N\cdots O}$  2.696(1) Å). This interaction is assisted by a weak C–H $\cdots$ O hydrogen bond and is a pyridine:carboxylic acid  $R_2^2(7)$  HB synthon.

The charge assisted  $N^+H\cdots O^-$  HBs in 5–8 have donor–acceptor distances (Table 2) equivalent to or shorter than 2.6 Å and are therefore classed as strong HBs. For all but 6, these distances are close to the region where temperature dependent proton migration across  $N^+H\cdots O^-$  SSHBs has previously been observed, near 2.55 Å;<sup>3,14–16</sup> the shortest is formed in 8. The shorter  $N^+H\cdots O^-$  HBs are formed as a discrete link  $D_1^1(3)$  between isonicotinamide and organic acid co-former, as in 5 and 8. In this scenario, the carboxylate group is significantly twisted out of co-planarity of the isonicotinamide N-heterocyclic ring (CONC torsions: 5 –85.41(9)°, 8 29.3(2)°). This arrangement may allow a closer approach of the carboxylate group to the protonated isonicotinamide molecule.

Beyond the charge assisted  $N^+H\cdots O^-$  interactions, the isonicotinamide molecules (protonated and neutral) associate



**Fig. 3** The charge assisted  $N^+H\cdots O^-$  HBs in 5–8 formed singularly (5 and 8) or as part of a pyridinium:carboxylate  $R_2^2(7)$  hydrogen bond synthon (6, 7).



**Table 2** Hydrogen bond parameters of the interactions connecting isonicotinamide and acid co-formers in 5–8: D is the HB donor atom and A is the HB acceptor atom

| System | Components                                    | Interaction | D–H (Å)   | H···A (Å) | dD...A (Å) | <DHA (°) |
|--------|---|-------------|-----------|-----------|------------|----------|
| 5      | 2 : 1 isonicotinamide phthalic acid           | N3–H3···O3  | 1.102(18) | 1.449(18) | 2.5499(10) | 176.7(2) |
|        |   | O1–H1···N1  | 1.04(2)   | 1.66(2)   | 2.6955(10) | 179(2)   |
| 6      | 1 : 1 isonicotinamide 2,4-dinitrobenzoic acid | N1–H1···O1  | 1.046(18) | 1.572(18) | 2.6000(8)  | 166.2(2) |
| 7      | 2 : 2 isonicotinamide 2,4-dinitrobenzoic acid | N2–H1···O9  | 0.985(18) | 1.660(19) | 2.6347(12) | 169.5(2) |
|        |   | N1–H2···O5  | 1.05(2)   | 1.50(2)   | 2.5542(12) | 178(2)   |
| 8      | 1 : 1 isonicotinamide 3,5-dinitrobenzoic acid | N1–H1···O1  | 1.084(18) | 1.484(18) | 2.5350(13) | 161.4(2) |

via centrosymmetric amide:amide  $R_2^2(8)$  HB dimers formed of N–H···O HBs. Common structural features are found in these dimers where the charge assisted  $N^+–H···O^-$  HBs are shorter; the amide:amide HB distances are longer (ESI† Table S9,  $d_{D...A}$  IN:IN) whilst the isonicotinamide amide group is increasingly out of co-planarity of the N-heterocycle ring (ESI† Table S9, <OCCC). It may be that these features work to facilitate or compensate for the formation of the shorter charge assisted  $N^+–H···O^-$  HBs in these types of materials.

In terms of crystal engineering approaches, there is no clear distinction between one acid co-former over the other, in terms of best choice to access the shorter of the charge assisted  $N^+–H···O^-$  HBs. However, it is clear that from this set, the combination of isonicotinamide with either *ortho*-substituted or dinitrobenzoic acids is a good route to forming shorter hydrogen bonding interactions. As is the case for the O–H···O SSHBs, this may be a result of the increased acidity of the carboxylic acid group<sup>32,50–52</sup> and thus an enhanced charge-assisted component of the HB<sup>55,56</sup> for these types (nitro) or positions (*ortho*) of substituent group.

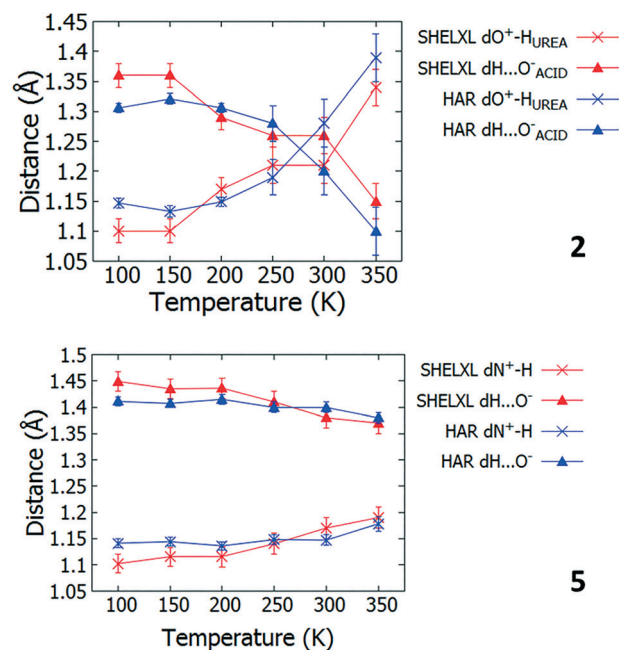
### Multi temperature measurements reveal migratory behaviour in 2 and 5

Multi temperature SCSXRD measurements were performed on 1 to 8 between 100 and 350 K to explore the SSHBs for temperature dependent proton migration behaviour. The high throughput nature of the SCSXRD technique was exploited in collecting a significant number of data points for each system whilst attenuating the synchrotron beam to ensure the sample continued to diffract across the whole temperature range. Following these measurements, migratory behaviour is indicated in 2 and 5 only.

For systems 2 and 5, in both the SHELXL<sup>57</sup> and HAR<sup>27</sup> crystal structures, a lengthening of the D–H distance and a shortening of the H···A distance occurs in the SSHB as a function of temperature (Fig. 4, ESI† Tables S10 and S11) suggestive of a migrating H-atom. In 2, the H-atom migrates away from the urea group up to 250 K where it occupies a more central position. Between 300 and 350 K, almost complete transfer to the organic acid is indicated. In 5, the H-atom migrates away from the HB donor group ( $N^+$ ) to a more central position in the charge assisted  $N^+–H···O^-$  SSHB as a function of temperature. The D–H/H···A distances change gradually as a function of temperature and in a consistent manner for 2 (up to 300 K)

and 5 (up to 350 K) whilst at the extremes of the migration temperature range they are appreciably different (2  $d_{O-H}$  1.10(2) Å at 100 K vs. 1.34(3) Å at 350 K, 5  $d_{N^+-H}$  1.102(18) Å at 100 K vs. 1.19(2) Å at 350 K). The fact there is agreement between the SHELXL<sup>57</sup> and HAR<sup>27</sup> crystal structures is significant and adds further confidence to the observation of the migration behaviour. The reliability of the SHELXL<sup>57</sup> refinements is aided by low values for the standard uncertainties on the D–H and H···A distances (of 0.02–0.03 Å) over the temperature range. On implementing the HAR<sup>27</sup> method, the precision of the H-atom positions is improved at low temperature, as is typically seen;<sup>26</sup> the observed migration trend is also more consistent. As is usually observed between X-rays and neutrons,<sup>24</sup> the changes in H-atom position observed as a function of temperature between 100 and 350 K are generally larger from the SHELXL<sup>57</sup> refinements for 2 and 5 (0.24(4) and 0.09(3) Å, respectively) than they are from the HAR<sup>27</sup> refinements (2, 0.18(1) and 5, 0.038(12) Å), giving values closest to those found by neutron diffraction.

The extent of temperature dependent proton migration occurring in 2 is similar to that in 1 : 1 adduct urea phosphoric



**Fig. 4** The D–H and H···A distances in 2 and 5 as refined using SHELXL<sup>57</sup> and HAR<sup>27</sup> over the 100 to 350 K temperature range.



acid,<sup>10</sup> where the H-atom migrates across the charge assisted  $O^+-H\cdots O^-$  hydrogen bond away from the protonated donor (urea) to a central position at 300 K. For 5, there are similarities in the nature of the migration occurring with 4,4-bipyridine 1,2,3,4-tetracarboxylic acid<sup>16</sup> and pyridine-3,5-dicarboxylic acid<sup>16</sup> where the H-atom moves away from the nitrogen donor towards the centre of the  $N^+-H\cdots O^-$  CAHB as a function of temperature. In contrast to 5, in these literature systems, the proton movement occurs to a far greater extent, located closer to the organic acid at room temperature.

Over the migration temperature range, the donor-acceptor distances of the SSHBs of interest are essentially invariant in both 2 and 5 (Fig. 5). This indicates that the observed H-atom migration behaviours are not simply a result of a lengthening SSHB due to lattice expansion as a function of temperature. In 2, the most significant change is between 300 and 350 K and may explain the slight inconsistency in the  $O^+-H$  and  $H\cdots O^-$  distances in this region that suggest a sudden large jump in H-atom position, as can be seen in Fig. 4.

Direct visualisation of the H-atom electron density in Fourier difference electron density maps can yield additional insight into proton transfer behaviour.<sup>58</sup> These maps are a valuable tool, giving a clearer and independent picture from X-ray diffraction data,<sup>24</sup> as they are not dependent on the refined parameters of the H atom involved in the SSHB, which is removed in the calculation of the map. Two dimensional Fourier difference electron density maps (FMAPs) have been generated as a function of temperature for SSHBs  $O1^+-H1\cdots O5^-$  (2) (Fig. 6) and  $N3^+-H3\cdots O3^-$  (5) (Fig. 7) from the SCSXRD data using the LIST 3 command in the SHELX<sup>37</sup> instruction file. These maps have been generated with the H-atom of interest removed from the model such that its density is visible (contours in the red to white region) when visualised using the MAPVIEW option in WinGX.<sup>39</sup> In the 100 K FMAPs for 2 and 5, the H-atom density is partially smeared along the HB, asymmetric in shape with a tail end that points towards the acceptor atom; this is characteristic of a H-atom in a short strong interaction.<sup>13</sup> In 2, this smearing increases with increasing temperature up to 250 K where it becomes more

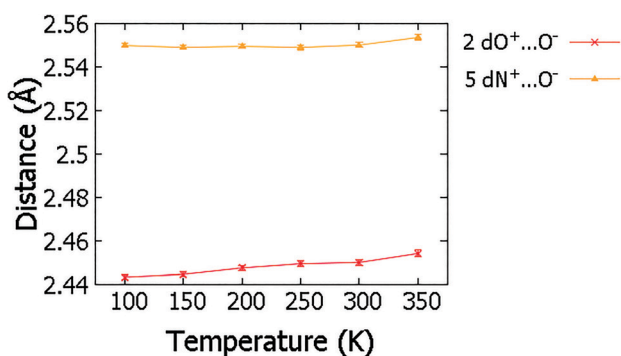


Fig. 5 The SSHB donor-acceptor distance in complex 2 and 5 obtained from refinement in SHELXL<sup>57</sup> over the 100 to 350 K temperature range.

elongated in shape. At 300 K, the peak maxima has moved to a central position and the H-atom density is evenly smeared across the centre of the SSHB, indicating a more centrally located H-atom in agreement with the refined H-atom positions at this temperature. At 350 K, the H-atom density peak maxima is located closer to the acceptor  $O2^+$  and the extended tail points more towards the opposite end of the SSHB, indicating its transfer. This switch in asymmetry of the H-atom density as a function of temperature is similar to that seen in the FMAPs of an equivalent temperature dependent proton migration material, the 1:1 adduct of *N,N*-dimethylurea oxalic acid.<sup>59</sup> Similarly in 5, ramping the temperature to 250 K causes the H-atom density to change shape and become more elongated along the SSHB. At 300 K the H-atom density peak maxima broadens suggesting increased thermal motion that could correspond to its movement towards the  $O3^-$  acceptor atom. At 350 K, the H-atom density is more symmetrically smeared along the interaction and the peak maximum occupies a more central position. It is less clear in this case (5) whether the changes occurring in the H-atom density as a function of temperature are consistent with proton migration or simply an effect of increased thermal motion of the H-atom with increasing temperature of the system;<sup>13</sup> the changes are less significant than seen for 2.

The generated FMAPs for 2 and 5 are of good quality in each case featuring low and consistent noise over the temperature range aiding the determination of the H-atom behaviour. In general, there is a slight difference in the FMAPs in the refined H-atom position and the peak maximum. This is likely due to the SSHB H-atom being refined with an isotropic thermal parameter which, characteristically in an SSHB, has a more anisotropic shape; this leads to an apparent shift of the atom away from its true position during the refinement.<sup>24</sup> The difference is, however, small and consistent over the temperature range meaning the refined H-atom positions are made more reliable for deducing proton migration trends.

For 2 and 5, changes in molecular geometries of the heavy atom groups forming the SSHBs as a function of temperature provide corroborating evidence for a migrating H-atom (Fig. 8, ESI† Tables S12 and S13). In both 2 and 5, the bond distances of the protonated groups ( $C=O_{UREA}^+$  and  $C-N_{IN}^+$ ) forming the SSHBs undergo some shortening as a function of temperature indicating a slight increase in their bond orders and supporting the migration of the H-atom away from this end of the HB. At 350 K, the bond distance for the  $C=O_{UREA}^+$  group in 2 is longer than might be expected for a fully deprotonated *N,N*-dimethylurea (WIFKEB02 1.245(2) Å at 273 K).<sup>60</sup> This post migration distance is, however, consistent with related *N,N*-dimethylurea systems exhibiting similar migration behaviour.<sup>13</sup> At the opposite end of the SSHB, the  $C-O^-$  bond distances for both 2 and 5 also shorten as a function of temperature. This is less consistent with migration behaviour; an increase in this bond length would be expected where the H-atom migrates towards this group. This effect is, however, very small and could be caused by a redistribution



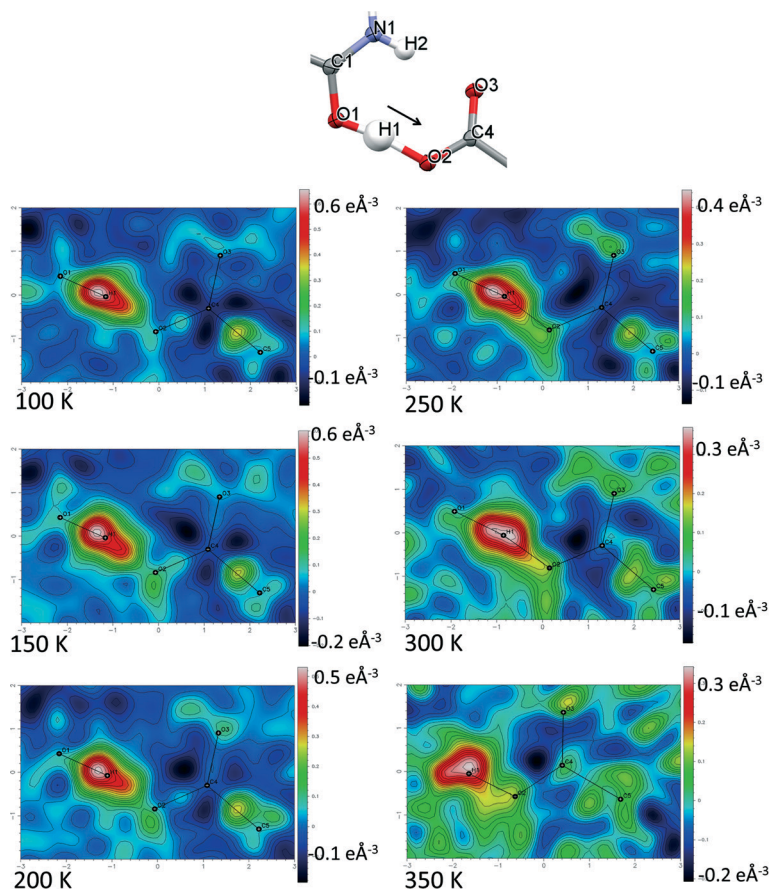


Fig. 6 Fourier difference electron density maps showing the electron density associated with the H-atom in the  $\text{O}^+-\text{H}\cdots\text{O}^-$  SSHB in 2 over the 100 to 350 K temperature range. The 350 K FMAP is missing atom O1 where it was not possible to generate a 2D plane containing all three atoms of the SSHB.

of charge in the carboxylate group as a function of temperature.

### Inconclusive or absent behaviour

Temperature dependent proton migration behaviour in contrast cannot be confirmed in 1, 3, 4, 6–8. For 7 and 8, the behaviour of the SSHB H-atom appears on the edge of migratory behaviour. Changes in the  $\text{N}^+-\text{H}$  and  $\text{H}\cdots\text{O}^-$  distances as a function of temperature correspond to only a slight migration of the H-atom towards the centre of the  $\text{N}^+-\text{H}\cdots\text{O}^-$  SSHBs (Fig. 9, ESI† Tables S14 and S15). Investigating these migrations in more detail in 7, it is clear that these changes are within error in the SHELXL<sup>57</sup> refinement. Also, the agreement between the SHELXL<sup>57</sup> and HAR<sup>27</sup> methods is not good; this is likely due to the HAR<sup>27</sup> method affording longer D–H distances closer in value to those found from neutron diffraction which are typically longer probing nuclear position rather than electron density.<sup>24</sup> A significant lengthening of the SSHB donor–acceptor distance also occurs for 7 (Fig. 10) such that any changes in the SSHB H-atom are more likely attributed to HB expansion. The supporting FMAPs in the region of the SSHBs show inconsistent behaviour (Fig. 11) meaning it is not possible to confidently identify migration

behaviour in this material. They do not support the changes in the  $\text{N}^+-\text{H}$  and  $\text{H}\cdots\text{O}^-$  distances as a function of temperature, instead showing no identifiable trend. An elongation of the hydrogen atom electron density occurs along the hydrogen bond but this does not increase consistently across the temperature range. From 250 K, this density appears as two peaks suggesting that proton disorder is more likely to be occurring and by 350 K, the H-atom peak is becoming lost in the increased thermal motion of the entire system; its signal cannot be deciphered from background noise. In 8, the SSHB donor–acceptor distances in contrast show minor variations as a function of temperature (Fig. 10) meaning any observed migratory behaviour would be ruled out from being attributed to lattice expansions. However, in the FMAPs (Fig. 12) the H-atom density remains close to the  $\text{N}^+$  donor atom indicating a static H-atom over the temperature range. Only significant structural changes in the HB groups (Fig. 13, ESI† Tables S16 and S17) that would support migratory behaviour as a function of temperature occur in 7; shortening of the isonicotinamide HB donor group C–N bond distances occur as a function of temperature, similar to those in 5. System 7 would benefit from additional experimental studies to fully elucidate the proton transfer behaviour; it has however not been possible to grow further single crystals of 7 suitable for



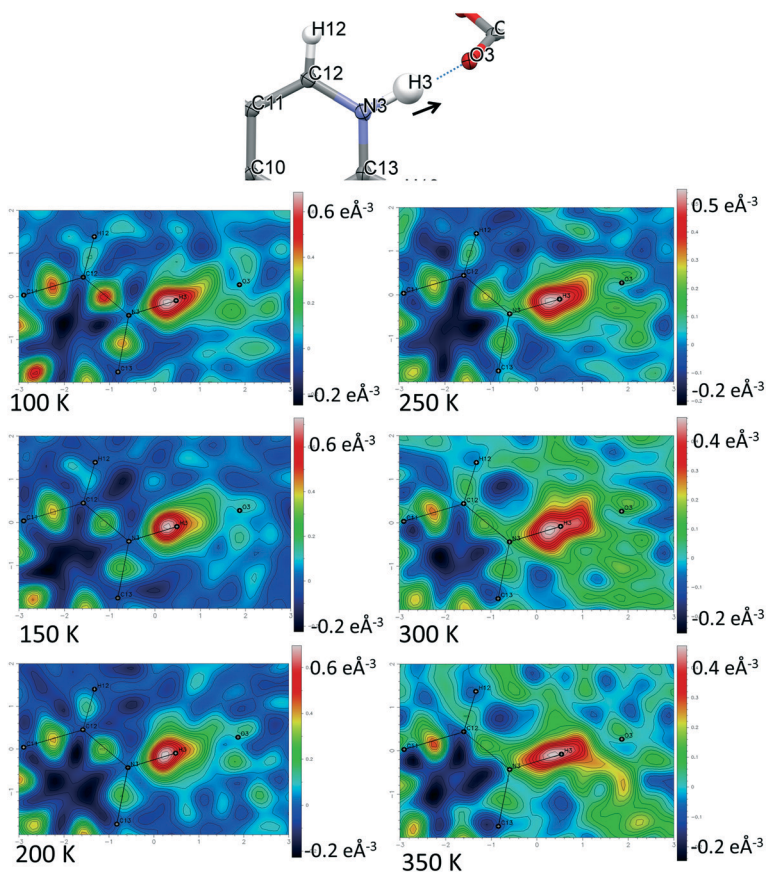


Fig. 7 Fourier difference electron density maps showing the electron density associated with the H-atom in the  $\text{N}^+\text{-H}\cdots\text{O}^-$  SSHB in **5** over the 100 to 350 K temperature range.

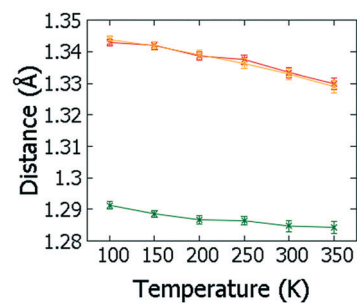
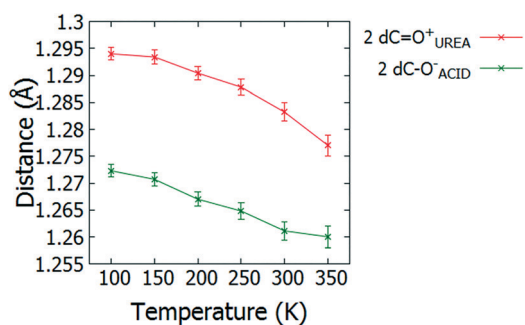


Fig. 8 Bond distances of the groups forming the SSHBs in **2** (top) and **5** (bottom) over the 100 to 350 K temperature range.

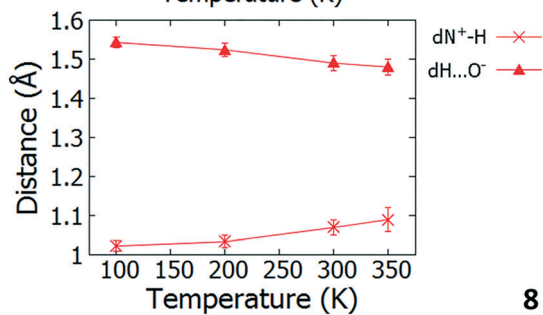
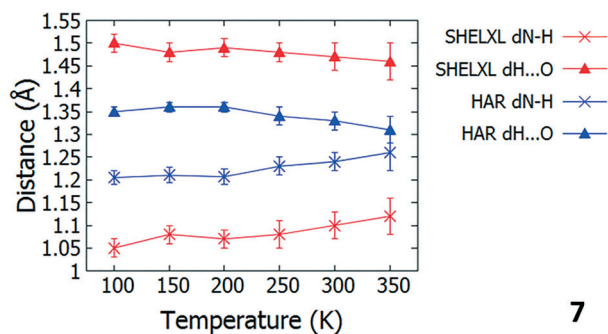


Fig. 9 Bond distances of the groups forming the SSHBs in **7** (top) and **8** (bottom) over the 100 to 350 K temperature range.





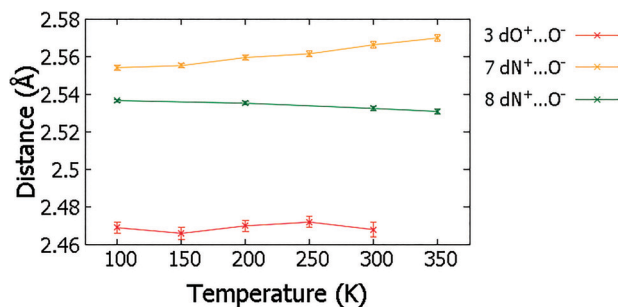


Fig. 10 The SSHB donor-acceptor distance in complex 3, 7 and 8 over the 100 to 350 K temperature range.

diffraction measurements since these SCSXRD measurements were undertaken, only the 1:1 system 6 is currently accessible.

The migration behaviour for 3 is unclear when considering the refined H-atom positions (ESI† Fig. S1, Table S18) and FMAPs (ESI† Fig. S2) over the 100 to 300 K temperature range. The H-atom position appears to fluctuate as a function of temperature whilst the determination of behaviour is made difficult by the significant noise present in the H-atom positions and FMAPs. The SSHB donor-acceptor distance also fluctuates significantly (Fig. 10) with temperature, meaning any migration behaviour is more likely to be attributed to

an expanding HB. This system could benefit from multi-temperature neutron diffraction measurements (requiring the growth of larger single crystals) as there may be interesting proton disorder behaviour above 100 K; though this suggestion is based on FMAPs with high levels of noise and is not suggested in the acid C=O/C-O bond lengths. In 1, 4 and 6 the evidence points to systems where temperature dependent proton migration behaviour is absent. Little variation in the D-H and H...A distances occur as a function of temperature (ESI† Table S19–S21, Fig. S1) and they are not significantly different at the temperature extremes. The FMAPs corroborate a static H-atom whose density remains close to the SSHB donor atom as a function of temperature (ESI† Fig. S3–S5).

### Insights into temperature dependent proton migration across SSHBs

The study of SSHBs in molecular crystals for temperature dependent proton migration using the SCSXRD technique adds to an understanding of the migration process in the solid-state. The high throughput nature of the SCSXRD technique means the number of systems explored for this migration behaviour can be greatly increased. In particular, this study has identified two new temperature dependent proton migration

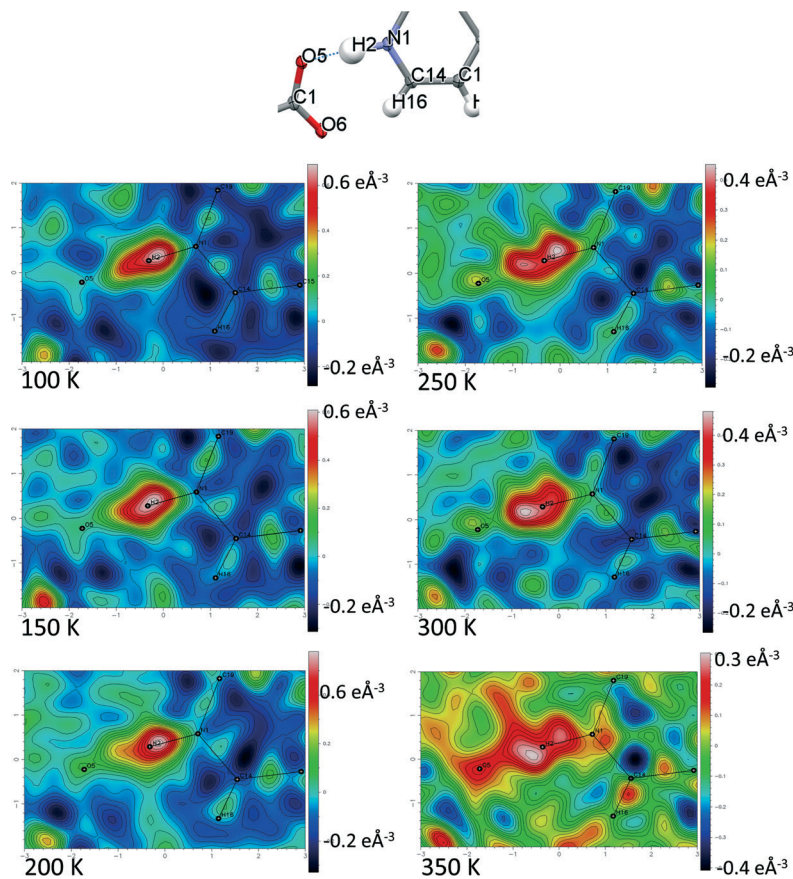


Fig. 11 Fourier difference electron density maps showing the electron density associated with the H-atom in the N<sup>+</sup>-H...O<sup>-</sup> SSHB in 7 over the 100 to 350 K temperature range.



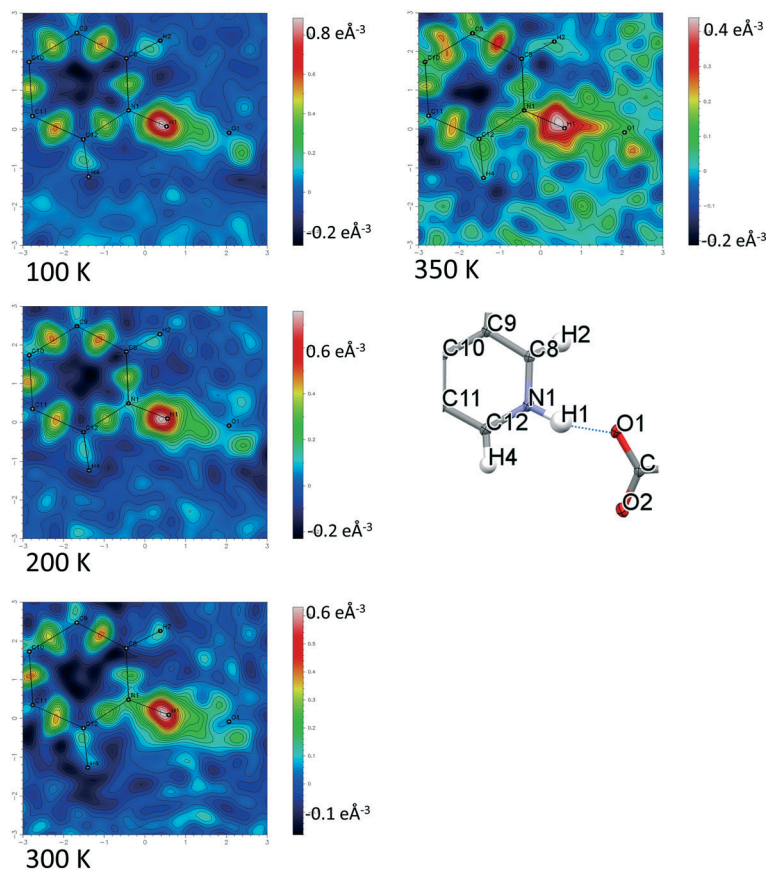


Fig. 12 Fourier difference electron density maps showing the electron density associated with the H-atom in the  $\text{N}^+\text{-H}\cdots\text{O}^-$  SSHB in **8** over the 100 to 350 K temperature range.

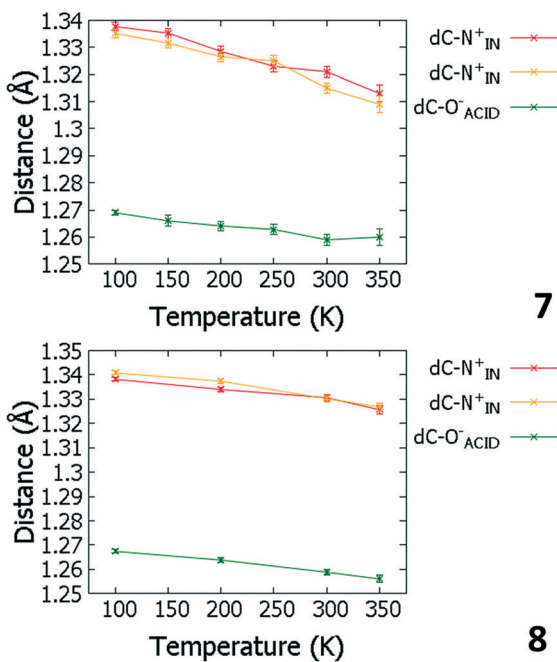


Fig. 13 Bond distances of the groups forming the SSHBs in **7** (top) and **8** (bottom) over the 100 to 350 K temperature range.

materials, **2** and **5**, two in which this behaviour is less clear (**7** and **8**) and four (**1**, **3**, **4** and **6**) in which this behaviour is absent.

A comparison with proton migration systems found previously<sup>3,10,13–16,61</sup> reveals insights related to SSHB donor–acceptor distance. For  $\text{O-H}\cdots\text{O}$  SSHBs, in substituted urea: organic acid systems, a limiting donor–acceptor distance of 2.45 Å has been proposed,<sup>13</sup> above which temperature-dependent proton migration is not expected. The related multi-component molecular crystals studied here containing  $\text{O-H}\cdots\text{O}/\text{O}^+\text{-H}\cdots\text{O}^-$  SSHBs are consistent with this proposal;<sup>59</sup> proton migration is not indicated in any of the molecular complexes with  $\text{O}\cdots\text{O}$  donor–acceptor distances above 2.45 Å, only in **2** where the  $\text{O}\cdots\text{O}$  distance is 2.441(1) Å at 100 K. For  $\text{N}^+\text{-H}\cdots\text{O}^-$  SSHBs in related N-heterocycle:organic acid molecular complexes, temperature dependent proton migration has not been identified in systems with donor–acceptor distances longer than 2.55 Å, having been reported previously in isoniazid 4-aminosalicylic acid ( $d_{\text{N}\cdots\text{O}}$  2.548(2) Å),<sup>14</sup> pyridine-3,5-dicarboxylic acid ( $d_{\text{N}\cdots\text{O}}$  2.523(2) Å)<sup>15</sup> and 4,4'-bipyridine benzene-1,2,4,5-tetracarboxylic acid ( $d_{\text{N}\cdots\text{O}}$  2.5220(17) Å).<sup>16</sup> **5** has the longest  $\text{N}^+\text{-H}\cdots\text{O}^-$  hydrogen bond across which temperature dependent proton migration is suggested ( $d_{\text{N}\cdots\text{O}}$  2.551(1) Å). It is not possible to confidently identify the presence of temperature dependent proton migration in **7**, with a



slightly longer SSHB donor–acceptor distance ( $d_{N\cdots O}$  2.555(1) Å). It may be, for  $N^+-H\cdots O^-$  SSHBs, that the donor–acceptor limiting distance is in the region of 2.55 Å. This is, however, a preliminary assessment and more cases are needed before a firm conclusion about  $N^+-H\cdots O^-$  donor–acceptor distances can be made. It is interesting that for 8, despite the very short HB donor–acceptor distance of 2.5350(13) Å at 100 K, temperature dependent migration behaviour appeared to be absent. This may be related to the SSHB angle of this interaction which has reduced directionality (*ca.* 161°) compared to in systems 5 and 7 (*ca.* 180°). It may be that enhanced directionality in charge assisted  $N^+-H\cdots O^-$  HBs favour migratory behaviour; pyridine-3,5-dicarboxylic acid (CSD refcode DINICA12),<sup>15</sup> isoniazid 4-aminosalicylic acid (URUDER)<sup>14</sup> and 4,4'-bipyridine benzene-1,2,4,5-tetracarboxylic acid (WISNAN)<sup>62</sup> all have HB angles greater than *ca.* 168°.

In the complexes reported here, the donor–acceptor distance is therefore shown to ‘tune’ the occurrence of temperature dependent proton migration across SSHBs, occurring in some instances but not in others. This is in part known where donor–acceptor distances affect the shape of the HB PES<sup>6</sup> and can be correlated to HB strength and PES symmetry.<sup>63</sup> A shorter donor–acceptor distance favours an increasingly symmetric HB PES with a low barrier to proton transfer; in this situation, migration behaviour is more likely.

## Conclusions

In this work we have engineered seven multi-component molecular crystals containing SSHBs by combining isonicotinamide and *N,N*-dimethylurea components with organic acid co-formers. Analysis of the SSHBs across the systems revealed that the strategy of selecting nitro-substituted organic acids favoured the formation of shorter hydrogen bonds in general whilst for the *N,N*-dimethylurea set, the presence of *ortho/para* substituents was additionally favourable. These findings contribute to the predictable design of these short interactions in the solid-state. If donor–acceptor distances in SSHBs can be predictably tuned, this would be a step forward in accessing proton migration materials more predictably in future.

SCSXRDX has been used to investigate the SSHBs for temperature dependent proton migration behaviour in eight multi component molecular crystals. This technique has potential in identifying such behaviour by implementing a protocol which considers the refined H-atom positions alongside those from Hirshfeld atom refinement in conjunction with other information including Fourier difference maps (FMAPs) and selected heavy atom structural parameters. Following this protocol, two new temperature dependent proton migration materials have been identified (2 and 5) in which migration of the H-atom occurs from the donor atom across the SSHB either towards the centre (5) or to the opposite side of the HB (2). The evaluation of such behaviour is aided by the availability of high quality, multi-condition data using the SCSXRDX technique. These findings contribute to the develop-

ment of the SCSXRDX method to allow it to be used routinely and with more confidence in future studies of proton transfer behaviour. Further insight into the dependence of temperature dependent proton migration on SSHB donor–acceptor distances is also revealed which can help to identify potential proton migration materials in future.

## Conflicts of interest

There are no conflicts of interest to declare.

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## References

- 1 L. H. R. Dos Santos, B. L. Rodrigues, Y. M. Idemori and N. G. Fernandes, *J. Mol. Struct.*, 2012, **1014**, 102–109.
- 2 P. Gilli, V. Bertolasi, V. Ferretti and G. Gilli, *J. Am. Chem. Soc.*, 1994, **116**, 909–915.
- 3 T. Steiner, I. Majerz and C. C. Wilson, *Angew. Chem., Int. Ed.*, 2001, **40**, 2651–2654.
- 4 M. Schmidtman, M. J. Gutmann, D. S. Middlemiss and C. C. Wilson, *CrystEngComm*, 2007, **9**, 743–745.
- 5 M. V. Vener, E. O. Levina, A. A. Astakhov and V. G. Tsirelson, *Chem. Phys. Lett.*, 2015, **638**, 233–236.
- 6 C. L. Perrin, *Acc. Chem. Res.*, 2010, **43**, 1550–1557.
- 7 I. A. Abronin, I. A. Koval'chuk and V. P. Sakun, *Russ. J. Phys. Chem. B*, 2016, **10**, 357–359.
- 8 Z.-S. Yao, K. Yamamoto, H.-L. Cai, K. Takahashi and O. Sato, *J. Am. Chem. Soc.*, 2016, **138**, 12005–12008.
- 9 D. M. S. Martins, D. S. Middlemiss, C. R. Pulham, C. C. Wilson, M. T. Weller, P. F. Henry, N. Shankland, K. Shankland, W. G. Marshall, R. M. Ibberson, K. Knight, S. Moggach, M. Brunelli and C. A. Morrison, *J. Am. Chem. Soc.*, 2009, **131**, 3884–3893.
- 10 C. C. Wilson, K. Shankland and N. Shankland, *Z. Kristallogr.*, 2001, **216**, 303–306.
- 11 K. Lee, B. Kolb, T. Thonhauser, D. Vanderbilt and D. C. Langreth, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2012, **86**, 104102.
- 12 M. T. Reetz, S. Höger and K. Harms, *Angew. Chem., Int. Ed. Engl.*, 1994, **33**, 181–183.



- 13 A. O. F. Jones, M.-H. Lemeë-Cailleau, D. M. S. Martins, G. J. McIntyre, I. D. H. Oswald, C. R. Pulham, C. K. Spanswick, L. H. Thomas and C. C. Wilson, *Phys. Chem. Chem. Phys.*, 2012, **14**, 13273–13283.
- 14 P. Grobelny, A. Mukherjee and G. R. Desiraju, *CrystEngComm*, 2011, **13**, 4358–4364.
- 15 J. A. Cowan, J. A. K. Howard, G. J. McIntyre, S. M. F. Lo and I. D. Williams, *Acta Crystallogr., Sect. B: Struct. Sci.*, 2005, **61**, 724–730.
- 16 J. A. Cowan, J. A. K. Howard, G. J. McIntyre, S. M.-F. Lo and I. D. Williams, *Acta Crystallogr., Sect. B: Struct. Sci.*, 2003, **59**, 794–801.
- 17 L. H. Thomas, N. Blagden, M. J. Gutmann, A. A. Kallay, A. Parkin, C. C. Seaton and C. C. Wilson, *Cryst. Growth Des.*, 2010, **10**, 2770–2774.
- 18 A. Gregorović, T. Apih, V. Žagar and J. Seliger, *Phys. Chem. Chem. Phys.*, 2019, **21**, 306–313.
- 19 J. S. Stevens, S. J. Byard, C. C. Seaton, G. Sadiq, R. J. Davey and S. L. M. Schroeder, *Phys. Chem. Chem. Phys.*, 2014, **16**, 1150–1160.
- 20 L. Rajput, M. Banik, J. R. Yarava, S. Joseph, M. K. Pandey, Y. Nishiyama and G. R. Desiraju, *IUCrJ*, 2017, **4**, 466–475.
- 21 P. Cerreia Vioglio, M. R. Chierotti and R. Gobetto, *Adv. Drug Delivery Rev.*, 2017, **117**, 86–110.
- 22 R. K. Harris, P. Y. Ghi, R. B. Hammond, C.-Y. Ma and K. J. Roberts, *Chem. Commun.*, 2003, 2834–2835.
- 23 M. Siskos, M. Choudhary and I. Gerathanassis, *Molecules*, 2017, **22**, 415.
- 24 A. Parkin, S. M. Harte, A. E. Goeta and C. C. Wilson, *New J. Chem.*, 2004, **28**, 718–721.
- 25 A. O. F. Jones, N. Blagden, G. J. McIntyre, A. Parkin, C. C. Seaton, L. H. Thomas and C. C. Wilson, *Cryst. Growth Des.*, 2013, **13**, 497–509.
- 26 M. Wońska, S. Grabowsky, P. M. Dominiak, K. Woźniak and D. Jayatilaka, *Sci. Adv.*, 2016, **2**, e1600192.
- 27 S. C. Capelli, H.-B. Burgi, B. Dittrich, S. Grabowsky and D. Jayatilaka, *IUCrJ*, 2014, **1**, 361–379.
- 28 M. Fugel, D. Jayatilaka, E. Hupf, J. Overgaard, V. R. Hathwar, P. Macchi, M. J. Turner, J. A. K. Howard, O. V. Dolomanov, H. Puschmann, B. B. Iversen, H.-B. Burgi and S. Grabowsky, *IUCrJ*, 2018, **5**, 32–44.
- 29 L. K. Saunders, *PhD Thesis*, University of Bath, U.K., 2016.
- 30 D. Braga and F. Grepioni, *Making Crystals by Design: Methods, Techniques and Applications*, Wiley-VCH, Weinheim, 2007.
- 31 M. Schmidtman, L. J. Farrugia, D. S. Middlemiss, M. J. Gutmann, G. J. McIntyre and C. C. Wilson, *J. Phys. Chem. A*, 2009, **113**, 13985–13997.
- 32 J. D. Hepworth, D. R. Waring and M. J. Waring, *Aromatic Chemistry*, Royal Society of Chemistry, Cambridge, 2002.
- 33 A. J. L. Jesus and J. S. Redinha, *J. Phys. Chem. A*, 2011, **115**, 14069–14077.
- 34 G. Winter, *J. Appl. Crystallogr.*, 2010, **43**, 186–190.
- 35 SADABS, Bruker AXS Inc., Madison, Wisconsin, USA.
- 36 Apex2, Bruker AXS Inc., Madison, Wisconsin, U.S.A.
- 37 G. Sheldrick, *Acta Crystallogr., Sect. A: Found. Crystallogr.*, 2008, **64**, 112–122.
- 38 P. Muller, R. Herbst-Irmer, A. L. Spek, T. R. Schneider and M. R. Sawaya, *Crystal Structure Refinement: A Crystallographer's Guide to SHELXL*, Oxford University Press, Oxford, 2006.
- 39 L. Farrugia, *J. Appl. Crystallogr.*, 2012, **45**, 849–854.
- 40 XPREP, Bruker AXS Inc., Madison, Wisconsin, USA.
- 41 S. Saha and G. R. Desiraju, *J. Am. Chem. Soc.*, 2018, **140**, 6361–6373.
- 42 A. O. F. Jones, C. K. Leech, G. J. McIntyre, C. C. Wilson and L. H. Thomas, *CrystEngComm*, 2014, **16**, 8177–8184.
- 43 L. K. Saunders, H. Nowell, P. R. Raithby and C. C. Wilson, *CrystEngComm*, 2016, **18**, 5916–5929.
- 44 A. Alhalaweh, S. George, D. Boström and S. P. Velaga, *Cryst. Growth Des.*, 2010, **10**, 4847–4855.
- 45 G. A. Jeffrey, *An Introduction to Hydrogen Bonding*, Oxford University Press, New York, 1997.
- 46 C. B. Aakeröy, M. E. Fasulo and J. Desper, *Mol. Pharmaceutics*, 2007, **4**, 317–322.
- 47 A. R. Eberlin, M. D. Eddleston and C. S. Frampton, *Acta Crystallogr., Sect. C: Cryst. Struct. Commun.*, 2013, **69**, 1260–1266.
- 48 C. B. Aakeroy and K. R. Seddon, *Chem. Soc. Rev.*, 1993, **22**, 397–407.
- 49 M. D. Ward, *Struct. Bonding*, 2009, **132**, 1–24.
- 50 M. A. Hernandez and A. Rathinavelu, *Basic Pharmacology Understanding Drug Actions and Reactions*, CRC Press, Boca Raton, 2006.
- 51 P. M. Dewick, *Essentials of Organic Chemistry: For Students of Pharmacy, Medicinal Chemistry and Biological Chemistry*, John Wiley & Sons, Ltd, Chichester, 2006.
- 52 T. L. Lemke, *Review of Organic Functional Groups: Introduction to Medicinal Organic Chemistry*, Lippincott Williams & Wilkins, Baltimore, 2003.
- 53 S. Tothadi and G. R. Desiraju, *Philos. Trans. R. Soc., A*, 2012, **370**, 2900–2915.
- 54 N. Shan, A. D. Bond and W. Jones, *Cryst. Eng.*, 2002, **5**, 9–24.
- 55 L. H. Thomas, A. O. F. Jones, A. A. Kallay, G. J. McIntyre and C. C. Wilson, *Cryst. Growth Des.*, 2016, **16**, 2112–2122.
- 56 S. J. Grabowski, *Annu. Rep. Prog. Chem., Sect. C: Phys. Chem.*, 2006, **102**, 131–165.
- 57 G. Sheldrick, *Acta Crystallogr., Sect. C: Struct. Chem.*, 2015, **71**, 3–8.
- 58 L. H. Thomas, A. J. Florence and C. C. Wilson, *New J. Chem.*, 2009, **33**, 2486–2490.
- 59 A. O. F. Jones, *PhD Thesis*, University of Bath, 2012.
- 60 G. Song, F.-Y. Bai, N. Xing, C. Chen, H. Shan and Y.-H. Xing, *Chem. Res. Chin. Univ.*, 2013, **29**, 408–413.
- 61 S. J. Ford, G. J. McIntyre, M. R. Johnson and I. R. Evans, *CrystEngComm*, 2013, **15**, 7576–7582.
- 62 A. J. Lough, P. S. Wheatley, G. Ferguson and C. Glidewell, *Acta Crystallogr., Sect. B: Struct. Sci.*, 2000, **56**, 261–272.
- 63 M. E. Tuckerman, D. Marx, M. L. Klein and M. Parrinello, *Science*, 1997, **275**, 817–820.

