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# A role of intermolecular interaction modulating thermal diffusivity in organosuperelastic and organoferroelastic cocrystals†

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Although the finding of superelasticity and ferroelasticity in organic crystals has been serendipitous, an increasing number of organic crystals with such deformation properties have been witnessed. Understanding the structure–property relationship can aid in the rational selection of intermolecular interactions to design organic crystals with desired superelastic or ferroelastic properties. In this study, we investigated the mechanical deformation in two cocrystals, prepared with the parent compound, 1,4-diodotetrafluorobenzene with two cofomers, 1,2-bis(4-pyridyl)ethane and pyrene. The parent compound and cofomers were chosen to introduce distinct weak interactions such as halogen bonds and C–H...F, and  $\pi$ ... $\pi$  interactions in the crystal structure. The two cocrystals exhibited different mechanical deformations, superelasticity, and ferroelasticity, respectively. The single-crystal X-ray diffraction and energy framework analysis of the crystal structure of the cocrystals revealed that both deformations were caused by mechanical twinning. Interestingly, a difference in the extent of deformation was observed, modulated by a combination of strong and weak intermolecular interactions in the superelastic cocrystal, and only weak interaction in the ferroelastic one. In this comparison, the superelastic cocrystal exhibited higher thermal diffusivity than the ferroelastic cocrystal, indicating the presence of symmetrical and relatively robust intermolecular interactions in the superelastic cocrystal.

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## Introduction

Over the last decade, distinct mechanical deformation behavior in organic molecular crystals has been reported, thereby making them a prospective candidate as next-generation structural materials for actuators,<sup>1</sup> multi-stimuli responsive sensors,<sup>2</sup> optical waveguides,<sup>3</sup> and soft robotics.<sup>4</sup> Organic molecular crystals have been found to exhibit several mechanical deformation behaviors such as elastic and plastic bending,<sup>5–7</sup> light-induced bending,<sup>8,9</sup> superelasticity,<sup>10–12</sup>

ferroelasticity,<sup>13,14</sup> superplasticity,<sup>15</sup> shape memory effect,<sup>16,17</sup> and many more. Superelastic and ferroelastic organic molecular crystals stand out as excellent and superior candidates in materials science due to their reversibility and diffusionless transformation behavior. Although both superelastic and ferroelastic deformations exhibit diffusionless transformation, the reversibility manner of the shape restoration differs. In superelastic deformation, the shape recovery is simply attained by removal of the applied force,<sup>10</sup> whereas in ferroelastic deformation, the shape change due to spontaneous strain is regained by application of force in the opposite direction to that of the deformation direction.<sup>13</sup> Digging into the mechanism, superelastic deformation is enabled by martensitic transformation,<sup>10</sup> whereas ferroelastic deformation is manifested by twinning deformation.<sup>18</sup> Recently, twinning has been found to be the common cause for both superelastic and ferroelastic deformation in organic crystals.<sup>11,19</sup>

While many organic molecular crystals exhibiting superelastic and ferroelastic deformation have been reported in recent years, a rational approach for designing and obtaining the desired superelastic and ferroelastic deformation is lacking.<sup>10–13,20</sup> For crystal structure designing with novel properties, a detailed understanding of various strong and weak molecular interactions is essential to modify the strength and geometry of intermolecular interactions. A thorough analysis

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showed that different crystal structures with polar space groups and weak intermolecular interactions such as weak hydrogen bonding, halogen bonding, and van der Waals interactions contribute to superelastic or ferroelastic deformation.<sup>20–22</sup> The influence of weak intermolecular interactions involving halogen groups such as an aromatic C–F group,<sup>23–26</sup> and halogen–halogen<sup>27,28</sup> and  $\pi$  stacking interactions<sup>29–31</sup> on directing the crystal packing in the presence or absence of strong hydrogen bonds has been extensively studied, and can serve as an excellent tool for crystal designing. Li *et al.* suggested that such crystals could be designed by considering the orientational change of molecules caused by twinning in superelastic and ferroelastic deformation.<sup>32</sup> They prepared a cocrystal composed of 3,5-dimethylbenzoic acid and 4,4'-bipyridine bonded by an acid-pyridine hetero-synthon and resembling an organic trimer, which demonstrated a bidirectional superelastic deformation through a 90° rotation of the aromatic plane of 4,4'-bipyridine.<sup>32</sup> In our previous report about the shape memory effect of 1,2-bis(4pyridyl)ethane dodecafluorosuberate,<sup>17</sup> we realized that the introduction of weak dispersive interactions like C–F $\cdots\pi$ , and C–H $\cdots$ F in the crystal structure changed the deformation behavior from biased ferroelastic<sup>33</sup> to the shape memory effect (interconversion of superelastic to ferroelastic). In this case, this weak dispersion interaction combined with strong hydrogen bonding N<sup>+</sup>–H $\cdots$ O<sup>–</sup> contributed to the shape memory effect with an increased critical forward stress value. Thalladi *et al.* mentioned that C–H $\cdots$ F interactions could be exploited to construct desirable organic crystals as compared to well-known C–H $\cdots$ O and C–H $\cdots$ N interactions owing to their ability to stabilize the crystal structures.<sup>34</sup> Additionally, Jain *et al.* demonstrated the role of halogen bonds in synthesizing higher-order cocrystals and further established that halogen bonds can exhibit analogous behavior to hydrogen bonds.<sup>35</sup> Mukherjee and Desiraju previously indicated that the strength of halogen bonds can manifest in elastic or plastic bending in organic crystals.<sup>36</sup> Thus, halogen bond-mediated/directed synthesis of a supramolecular assembly could be a possible way to design organic molecular crystals with desired superelastic and ferroelastic deformation. Variations in the strength of halogen bonds with excellent directionality<sup>37</sup> and other weak dispersive interactions<sup>38</sup> can manipulate the spontaneous strain generation required for ferroelastic deformation and spontaneous shape recovery in a superelastic deformation.

In addition to developing the designing principles for superelastic and ferroelastic deformation, studying the anisotropic thermal transport properties of these materials such as thermal diffusivity would be insightful owing to the range of thermal management applications of these crystals.<sup>39,40</sup> Previous studies have reported that the anisotropic mechanical properties of soft materials in response to high frequency can be correlated with thermal diffusivity.<sup>41,42</sup> Studying the anisotropic thermal properties in ferroelastic and superelastic organic crystals would be beneficial to understand the manner of heat transport with respect to their structural order. Furthermore, the reversible phase transition properties of these crystals in response to mechanical stress can be utilized as thermal switches.<sup>43,44</sup> The micro-level miniaturization of the integrated

devices demands efficient flexibility and thermal management, which can be fulfilled by the controlled deformation behavior of these crystals and used as wearable devices.<sup>45–47</sup> It has been demonstrated that inorganic ferroelastic materials can be utilized for thermal management applications.<sup>44,48–50</sup> Sieradzki *et al.* studied the changes in the thermal conductivity of Li<sub>2</sub>-TiGeO<sub>5</sub> ceramics in the ferroelastic phase transition or domain formation.<sup>50</sup> Wang *et al.* reported the difference in thermal conductivity values of layered PdSe<sub>2</sub> before and after ferroelastic phase transition by uniaxial compression and demonstrated that the high ratio of the switchable thermal conductivity value between the two phases could be utilized for flexibility and thermal management.<sup>44</sup> However, there is only one study concerning the measurement of anisotropic heat transport properties in a superelastic organic crystal due to the limited approaches/techniques to measure the thermal diffusivity of organic crystals.<sup>40</sup>

Keeping these considerations in mind, we attempted to prepare single crystals of two cocrystals consisting of a weak halogen bond donor and acceptor, where 1,4-diiodotetrafluorobenzene served as the halogen donor. It produced cocrystals with the halogen acceptor 1,2-bis(4-pyridyl)ethane (cocrystal 1) and pyrene (cocrystal 2), respectively. Different deformation behaviors, *i.e.*, superelastic and ferroelastic deformation, were observed in 1 and 2, respectively. The anisotropic thermal diffusivity values of the two cocrystals were measured by typical microscale temperature-wave analysis ( $\mu$ -TWA). A difference in the combination of relatively strong halogen interactions (C–I $\cdots$ N) and weak interactions (C–H $\cdots$ F and  $\pi\cdots\pi$ ) in the crystal structure of the two cocrystals could be responsible for the different mechanical deformations and thermal diffusivity behaviors.

## Results and discussion

### Single crystal synthesis and characterization

For cocrystal preparation, a halogen bond donor, 1,4-diiodotetrafluorobenzene was selected as the parent compound. In previous studies, 1,4-diiodotetrafluorobenzene has been utilized to prepare supramolecular structures owing to its ability to form a range of interactions, including C–I $\cdots$ N, C–H $\cdots$ F, and  $\pi\cdots\pi$ .<sup>51</sup> Such weak C–H $\cdots$ F, C–F $\cdots\pi$  interactions can form crystal packing even in the absence of strong hydrogen bonds.<sup>52</sup> Furthermore, the C–I $\cdots$ N interactions have been demonstrated to be strong and directional to facilitate the self-assembly of a 1D infinite network, and other interactions such as C–H $\cdots$ F and  $\pi\cdots\pi$  enable the formation of a 3D network.<sup>53</sup> In addition, Corradi *et al.* noted that the halogen bonding of 1,4-diiodotetrafluorobenzene was significantly accountable for the supramolecular structure formation with 1,2-bis(4-pyridyl)ethane and hydroquinone compared to hydrogen bonding.<sup>51</sup> In order to manipulate the weak interactions in the crystal structure, two cofomers of 1,2-bis(4-pyridyl)ethane and pyrene were selected on the basis of halogen bond acceptor strength. 1,2-Bis(4-pyridyl)ethane acts as a strong halogen acceptor due to the electron rich nitrogen atom, while pyrene acts as a weak



halogen acceptor due to the absence of an electron rich group in its structure.

The colorless block shape single crystals of cocrystal **1** were obtained by slow evaporation of a dichloromethane solution of 1,4-diiodotetrafluorobenzene and 1,2-bis(4-pyridyl)ethane mixed at a molar ratio of 1 : 1.<sup>54</sup> The single crystal of cocrystal **1** crystallizes in a triclinic crystal system with space group  $P\bar{1}$ . The molecules formed an infinite 1D chain *via* the C–I $\cdots$ N (4.898(4) Å, 175.6°) halogen bonds with  $d(I1\cdots N1) = 2.795(3)$  Å in an ABAB manner along the *a* axis. Furthermore, weak C–H $\cdots$ F hydrogen bonds connect the nearby 1D chains to create a three-dimensional crystal structure along the *bc* plane (Fig. 1 and Table S1, ESI†).

On the other hand, the colorless block-shaped single crystals of cocrystal **2** were obtained by slow evaporation of an ethanolic solution of 1,4-diiodotetrafluorobenzene and pyrene mixed in a molar ratio of 1 : 1.<sup>55</sup> The single crystals of **2** crystallize in

a monoclinic crystal system with space group  $P2_1/c$ . The single crystal of cocrystal **2** formed a column-like structure *via*  $\pi$ – $\pi$  stacking in an A $\cdots$ B $\cdots$ A $\cdots$ B $\cdots$  face-to-face manner along the *a* axis with an interplanar distance of 3.450 Å. Furthermore, the adjacent columns are linked together by C–I $\cdots$ I–C (3.812(6) Å) interactions along the *b* axis (Fig. 1 and Table S1, ESI†). The shortest C–H $\cdots$ F interactions (2.56 Å, 145.05°) were found along the *c* axis, stabilizing the crystal structure of **2**. The structures of cocrystals **1** and **2** along with their deformation behaviors are illustrated in Fig. 1.

### Microscopic observation

For studying the shear deformation behavior at a macroscopic level, mechanical shear stress on both the single crystals of **1** and **2** was applied independently at room temperature under a polarized microscope on the (011) plane and (001) plane,



Fig. 1 (a) Molecular structures and formation of cocrystals **1** and **2**, (b) illustration of the difference in the mechanical deformation of interest, and (c) crystal structures of (i) cocrystal **1** along the *bc* plane and (ii) cocrystal **2** along the *a* axis.



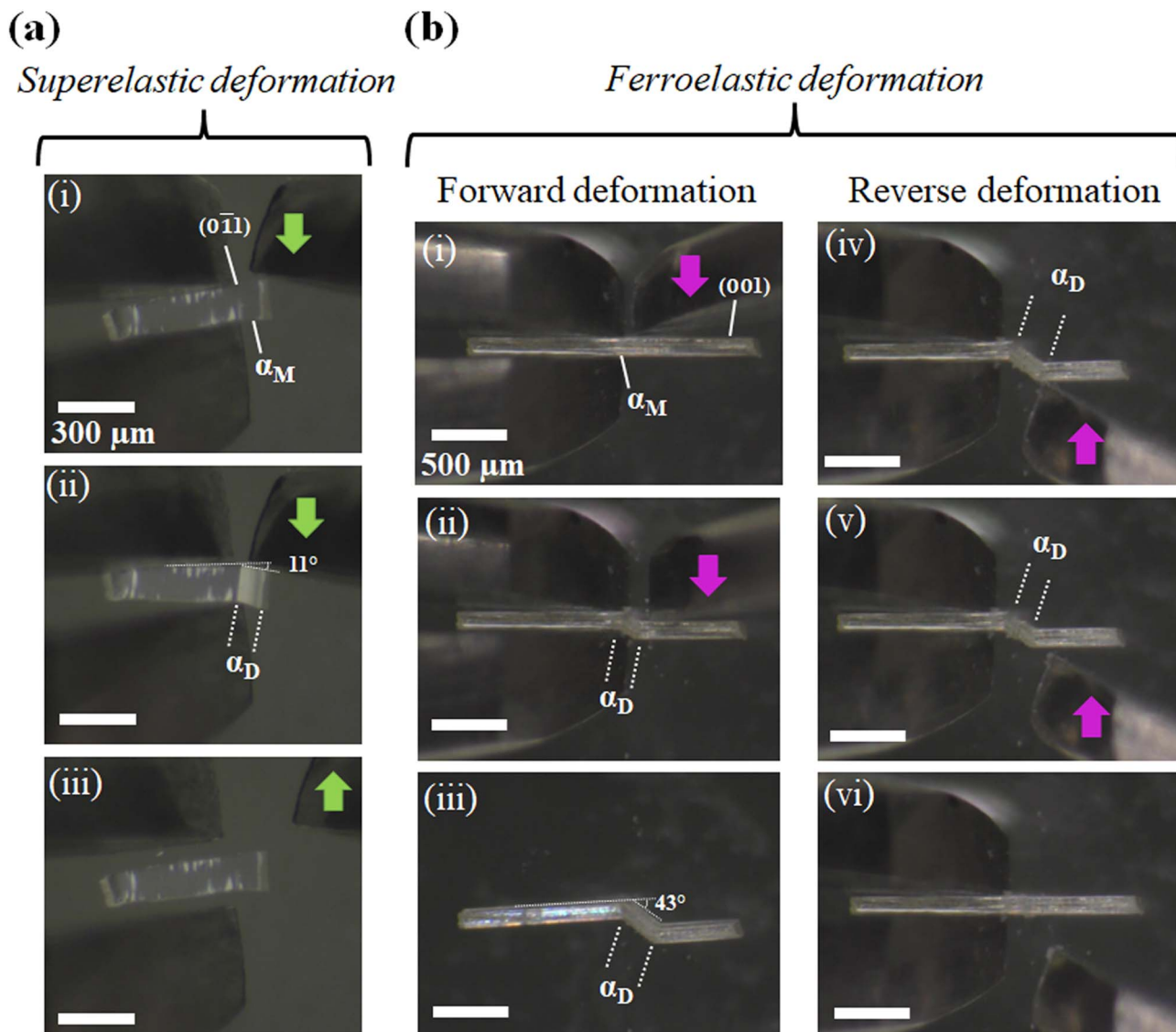


Fig. 2 Microscope observation with tweezers at room temperature: (a) superelastic deformation of cocystal 1 and (b) ferroelastic deformation of cocystal 2.

respectively (Fig. 1(b)). An application of shear stress on the (011) plane of single crystals of 1 caused a superelastic deformation (Fig. 2(a)). The superelastic deformation was initiated by the appearance of a stress-induced twinning domain ( $\alpha_D$ ) generated from the mother crystal ( $\alpha_M$ ) with a distinct boundary line at a bending angle of  $11^\circ$ . The deformed crystal was recovered to its original shape after the applied force was removed. On the other hand, single crystals of 2 showed a ferroelastic deformation on the application of shear stress on the (001) plane (Fig. 2(b)). A bending angle of  $43^\circ$  was demonstrated with the growth of daughter domains ( $\alpha_D$ ) from the mother domain ( $\alpha_M$ ) in the single crystal of 2, and a spontaneous strain was accumulated as residual strain until the removal of the force. Furthermore, on application of reverse force on the (001) plane, the daughter domain returned to its initial orientation, confirming the ferroelasticity.

### Mechanical deformation analysis

The stress–strain relationship between cocystals 1 and 2 was investigated by using a custom-designed setup: the shear force was applied with a jig to the specific crystal plane of the crystal fixed to a glass plate with epoxy glue. In a single crystal specimen of 1, the  $\alpha_D$  domain started to grow from the  $\alpha_M$  domain at an applied stress of 0.27 MPa on the (011) plane. At a critical stress of 0.27 MPa, the stress-induced twinning phase transformation ( $\alpha_D$ ) propagated, and the stress–strain curves displayed a characteristic plateau resembling a superelastic deformation (Fig. 3(a)). The original shape was regained by decreasing the stress in depicting the typical superelastic stress–strain hysteresis loop, validating the superelastic deformation. On the other hand, with the gradual increase of stress on the (001) plane of the single crystal of 2, the  $\alpha_D$  domain was generated at a coercive stress of 0.58 MPa, which was fairly



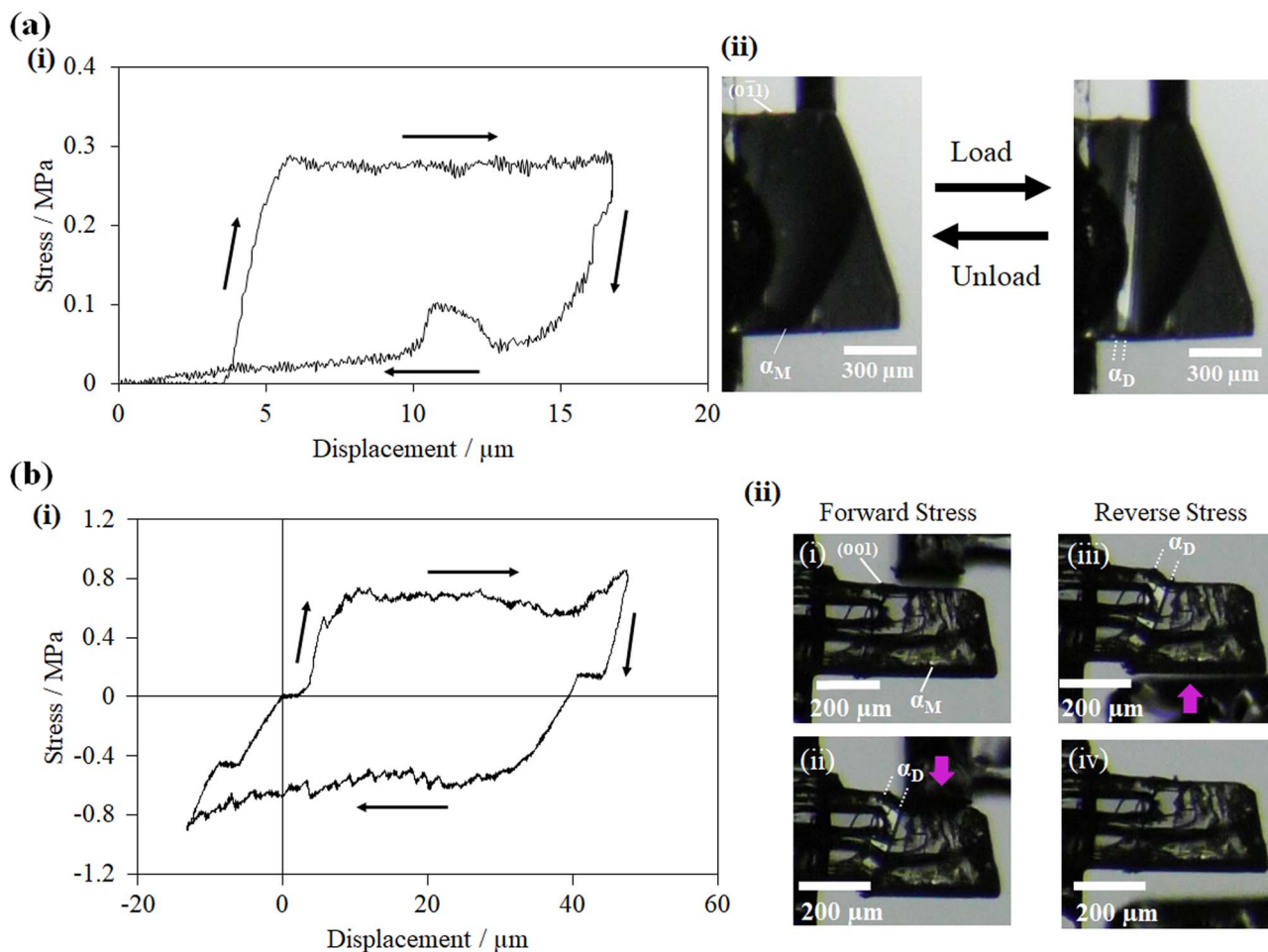


Fig. 3 Shear-strain test: (a) the superelastic hysteresis loop of the cocrystal of 1 (i) and snapshots during measurement (ii) and (b) the ferroelastic hysteresis loop of the cocrystal of 2 (i) and snapshots (ii).

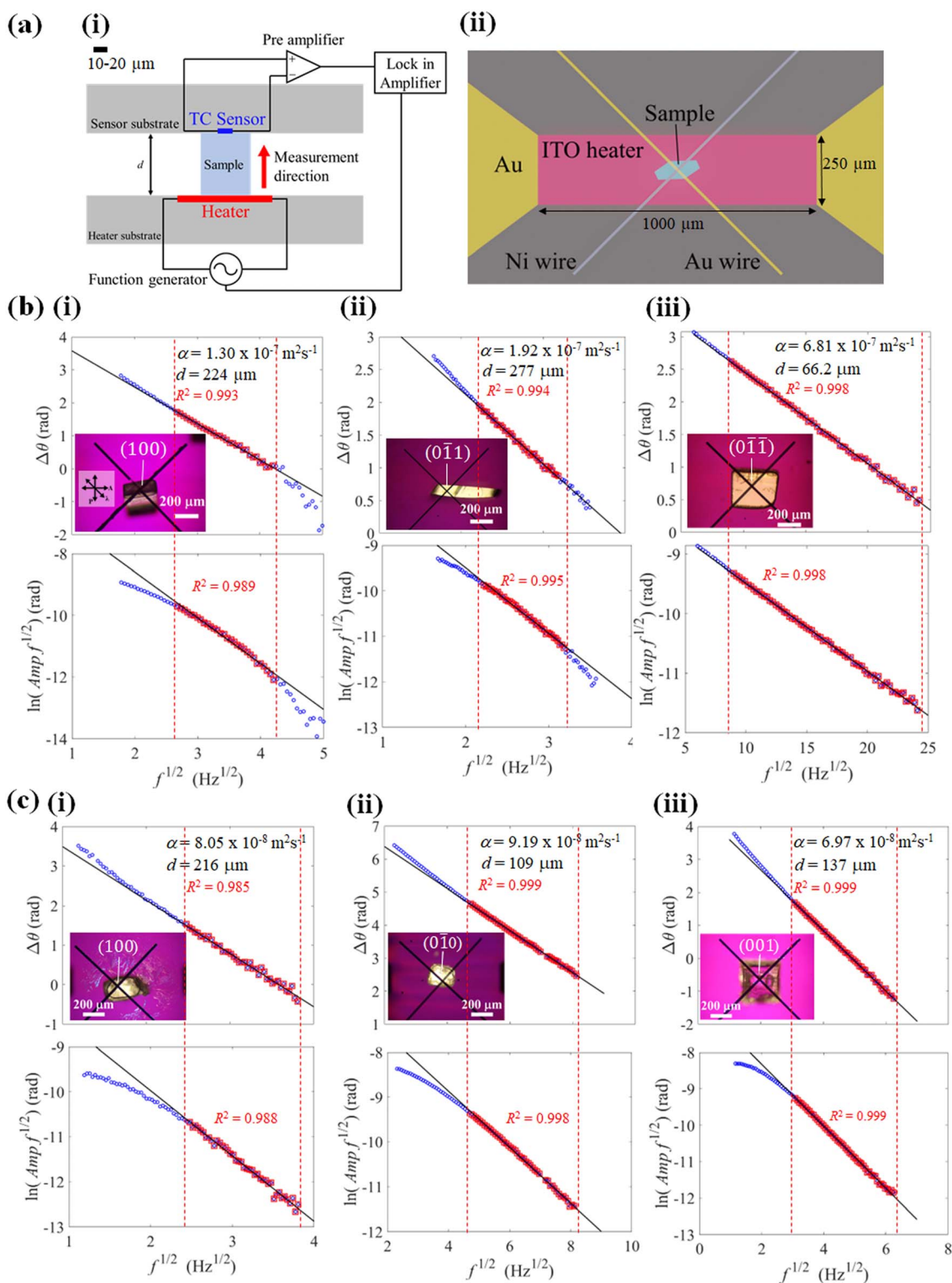
steady. Furthermore, by applying a shear stress of 0.57 MPa from the opposite direction, the  $\alpha_D$  domain was reverted to the  $\alpha_M$  domain with the corresponding release of the spontaneous strain. The hysteresis loop was established in the shear displacement graph, which is the confirmatory indicator of ferroelastic deformation (Fig. 3(b)). The observed stress strengths are similar to those of organosuperelastic/organoferroelastic crystals previously reported.<sup>13–19</sup>

### Thermal diffusivity studies

Thermal diffusivity is a specific property of materials related to the rate of thermal energy transfer through it. Determination of the thermal diffusivity value of the crystals would aid in understanding the heat transfer properties and their applicability in thermal management. In addition, it can also assist in understanding the changes in the thermal properties due to the structure, phase transitions, defects, *etc.* The anisotropic thermal diffusivity measurement of single crystals 1 and 2 were performed by  $\mu$ -TWA measurements<sup>40</sup> along its longer, wider, and thinner axis. The frequency change in the phase delay of the temperature wave was obtained for the three crystallographic axes of the crystals (Fig. 4, S1, S2, and Tables S2, S3, ESI†).

From the measured data, the slope of the high-frequency region, which was decided according to the linearity of both phase and amplitude data, was used to determine the thermal diffusivity. In single crystals of 1, the measured thermal diffusivity was found to be the highest  $6.30 \times 10^{-7} \text{ m}^2 \text{ s}^{-1}$  along the  $[0\bar{1}\bar{1}]$  direction followed by,  $2.04 \times 10^{-7} \text{ m}^2 \text{ s}^{-1}$  along the  $[0\bar{1}1]$  direction, and  $1.12 \times 10^{-7} \text{ m}^2 \text{ s}^{-1}$  along the  $[100]$  direction (Fig. 5, S1, and Table S2, ESI†). In contrast, the thermal diffusivity of the ferroelastic crystal was similar along the  $[0\bar{1}0]$  direction and  $[100]$  directions ( $0.93 \times 10^{-7} \text{ m}^2 \text{ s}^{-1}$ ). However, a lower thermal diffusivity was observed along the  $[001]$  direction ( $0.69 \times 10^{-7} \text{ m}^2 \text{ s}^{-1}$ ) which is the shear stress deformation direction of cocrystal 2 (Fig. 5, S2, and Table S3, ESI†). The highest obtained thermal diffusivity along the  $[0\bar{1}0]$  direction in the ferroelastic crystal was almost 7-fold lower compared to that of superelastic crystal along the  $[0\bar{1}\bar{1}]$  direction. In addition, the thermal diffusivity value of both the crystals was low in the direction of applied shear stress along  $[0\bar{1}1]$  of 1 and  $[001]$  of 2 compared to their orthogonal directions. Furthermore, the thermal diffusivity values in the daughter domains of cocrystals 1 and 2 were found to increase by 17% ( $7.38 \times 10^{-7} \text{ m}^2 \text{ s}^{-1}$ ) and 7% ( $0.99 \times 10^{-7} \text{ m}^2 \text{ s}^{-1}$ ) along the  $[0\bar{1}0]$  and  $[0\bar{1}\bar{1}]$  directions of 1





**Fig. 4** The frequency dependency of the phase delay measured for cocrystals 1 and 2 (a) schematic depiction of the experimental setup from the (i) cross-sectional view and (ii) top-view. The synchronized function generator and the lock-in amplifier determined the periodic heating and detection of the periodic temperature response from the sample as a phase delay. (b) (i), (ii), and (iii) cocrystals of 1 placed along its longer, wider and thinner axes corresponding to the [100], [011] and [011] directions, respectively, and (c) (i), (ii), and (iii) cocrystals of 2 placed along its longer, wider and thinner axes corresponding to the [100], [010], and [001] directions, respectively. The inset shows the optical image of each sample under a crossed-Nicols with a sensitive plate (530 nm wave plate) taken by using a polarized microscope (top view).





Fig. 5 The obtained thermal diffusivity values of cocrystals 1 (light green colored) and 2 (purple colored) with respect to the longer, wider, and thinner axes, respectively. Arrows indicate the direction of shear stress deformations.

and 2, respectively (Fig. S3, Table S4, ESI<sup>†</sup>). The switchable thermal diffusivity values of the mother and daughter domains indicate the potential for a thermal switch application of these crystals.

### Crystallographic studies

The underlying mechanism behind the different deformation behaviors of cocrystals 1 and 2 were investigated by single-crystal XRD analyses of the deformed crystals of 1 and 2. The cocrystal of 1 crystallizes in a triclinic crystal system with the

space group  $P\bar{1}$ . On the other hand, the cocrystal of 2 crystallizes in a monoclinic crystal system with space group  $P2_1/c$ . Both the crystal structures were similar to the previously reported one.<sup>51,55</sup> XRD investigations of 1 and 2 using face indexing revealed that the shear-induced twinning deformation initiated the development of the daughter crystal domain ( $\alpha_D$ ) from the mother crystal domain ( $\alpha_M$ ) upon application of shear stress on the (011) and (001) planes, respectively (Fig. 6, 7, S4 and Table S1, ESI<sup>†</sup>). According to crystallographic investigations, the twinning interfaces of cocrystals 1 and 2 are located at  $(100)_{\alpha_M}/(100)_{\alpha_D}$  and  $(10\bar{1})_{\alpha_M}/(\bar{1}01)_{\alpha_D}$ , respectively, which are perpendicular to the stacked column. A 180° rotation about the rotation axis was associated with the mother domain and daughter domains. This relationship is attributed to the polar  $P\bar{1}$  and  $P2_1/c$  point group symmetry, which caused the molecules to tilt by 41.81° of the 1,4-diiodotetrafluorobenzene molecule and 7.81° of 1,2-bis(4-pyridyl)ethane molecule in the case of cocrystal 1, and 57.48° of the 1,4-diiodotetrafluorobenzene molecule and 63.48° of the pyrene molecule in the case of cocrystal 2, instead of the actual 180° rotation (Fig. S4, ESI<sup>†</sup>). 1,4-diiodotetrafluorobenzene interacted through C–I...N and C–H...F interactions with the nearby molecules in crystal structure 1, where it is surrounded by four 1,2-bis(4-pyridyl)ethane molecules in a rhomboid configuration with sides of 10.401 Å and 7.379 Å (Fig. 6(b)(ii)). 1,4-diiodotetrafluorobenzene was rotated by 41.81° in relation to its mother domain configuration due to the twinning orientational change. As a result, the crystal structure is balanced and has a configuration analogous to a rhomboid

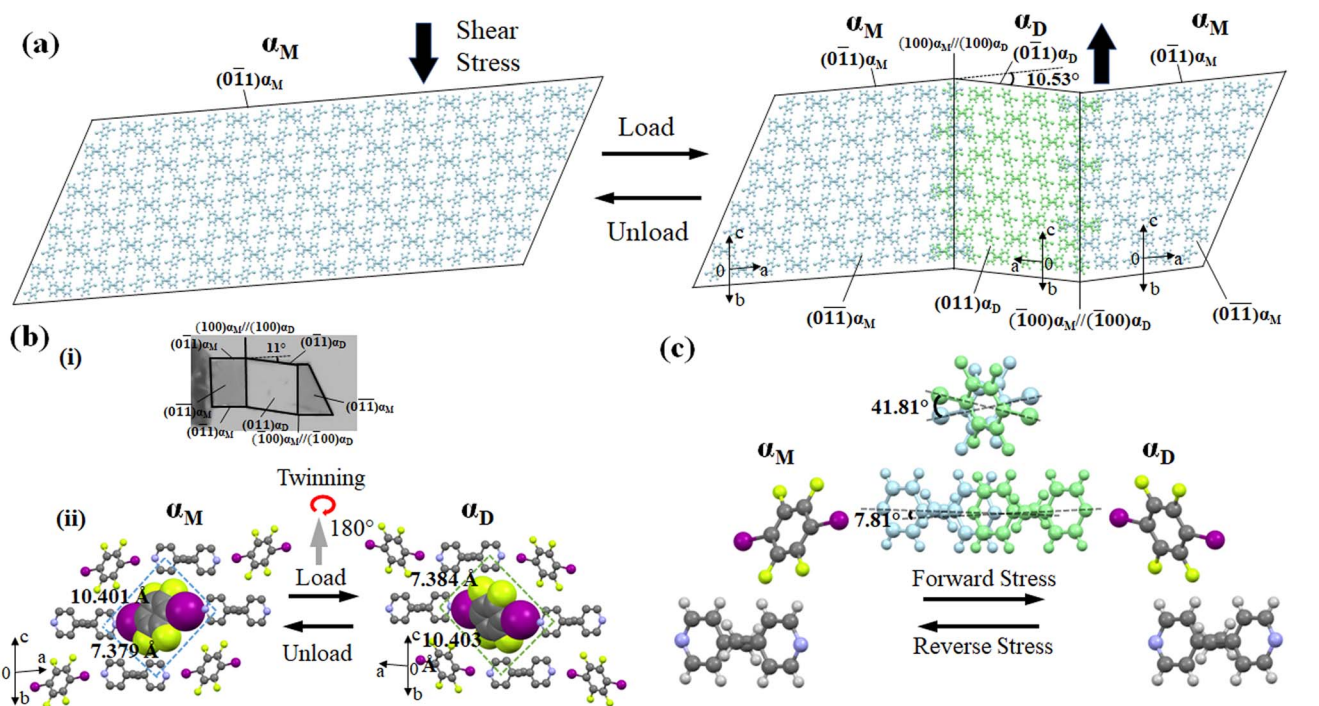


Fig. 6 (a) Estimated connecting manners of the  $\alpha_M$  domain and  $\alpha_D$  domain based on the X-ray diffraction measurements of single crystal 1. (b) Face indexing of the superelastic bent crystal used for SCXRD (i) and estimated twinning manners of the  $\alpha_D$  domain based on the  $\alpha_M$  domain (ii). (c) Estimated molecular movements at the interface at  $(100)_{\alpha_M}/(100)_{\alpha_D}$  (light blue and light green colored molecules indicate the  $\alpha_M$  and  $\alpha_D$  domains, respectively).



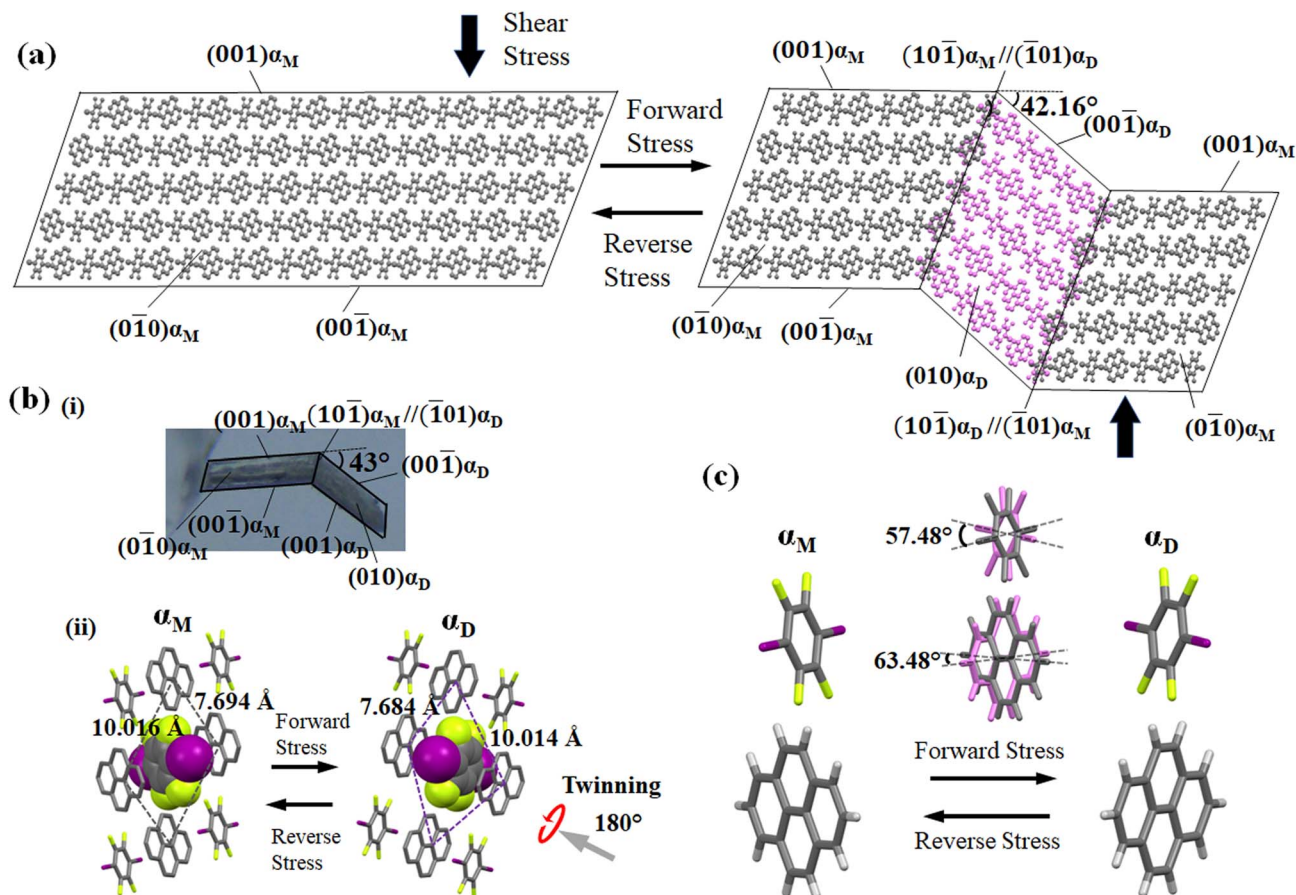


Fig. 7 (a) Estimated connecting manners of the  $\alpha_M$  domain and  $\alpha_D$  domain based on the X-ray diffraction measurements of single crystal 2. (b) (i) Face indexing of the ferroelastic bent crystal used for SCXRD. (ii) Estimated twinning manners of the  $\alpha_D$  domain based on the  $\alpha_M$  domain. (c) Estimated molecular movements at the interface at  $(10\bar{1})\alpha_M // (\bar{1}01)\alpha_D$ . Light gray and light purple colored molecules indicate the  $\alpha_M$  and  $\alpha_D$  domains, respectively.

arrangement rotated by  $90^\circ$  in the daughter phase with sides of  $7.384 \text{ \AA}$  and  $10.403 \text{ \AA}$ . Similar observations were seen for cocrystal 2, which also resembled a tunable rhomboid structure with sides of  $10.016 \text{ \AA}$  and  $7.694 \text{ \AA}$  in the mother domain and  $7.684 \text{ \AA}$  and  $10.014 \text{ \AA}$  in the daughter domain (Fig. 7(b)(ii)).

Although mechanical twinning manifested superelastic and ferroelastic deformations in both crystals, the deformation produced a different bending angle and dissipated energy ( $E_d$ ). A bending angle of  $10.53^\circ$  between  $\alpha_M$  and  $\alpha_D$  can be anticipated based on crystallographic results in cocrystal 1, and this angle aligned well with the  $11^\circ$  bending angle that was measured by optical microscopy. However, the bending angle was higher by  $32^\circ$  in the case of cocrystal 2 compared to cocrystal 1, which was estimated to be  $42.16^\circ$  by crystallographic analysis and  $43^\circ$  by optical microscopy. Due to the molecular movement by such a large angle, significant coercive stress was observed during the daughter domain development in ferroelastic deformation. Furthermore, the lower tilt angle of molecules in the case of molecule 1 could be due to the presence of relatively strong and ordered C-I $\cdots$ N ( $4.898(4) \text{ \AA}$ ,  $175.6^\circ$ ) halogen interactions, and the stable twisted chair form of bipyridine moieties which would stabilize the crystal during the application of stress.

In cocrystal 1, the strong C-I $\cdots$ N halogen bond is clarified by the shorter I-N ( $2.795(3) \text{ \AA}$ ) distance compared to other bipyridine-based halogen bonded complexes, relating to the two distantly placed twisted chair forms of the bipyridine rings in the structure. The strong C-I $\cdots$ N halogen-bonded interactions and weak C-H $\cdots$ F ( $3.528(3) \text{ \AA}$ ,  $141.98^\circ$  (F1 $\cdots$ H6a ( $2.71 \text{ \AA}$ ))) hydrogen-bonded interactions in cocrystal 1 can make its crystal packing robust compared to cocrystal 2 (Fig. 8(a)).

In cocrystal 2, C-I/I-C ( $d(I-I) 3.812(6) \text{ \AA}$ ) interactions hold the parallel-placed columns together. The connectivity between columns appears to be strengthened by C-I $\cdots$  $\pi$  ( $3.468(6) \text{ \AA}$ ) halogen bonding and C-H $\cdots$ F ( $3.367(5) \text{ \AA}$ ,  $145.05^\circ$ ) interactions  $d(F1\cdots H19) = 2.56 \text{ \AA}$  to maintain the 3D crystal structure (Fig. 8(b)).

### Correlation of anisotropic structural arrangement with mechanical behavior and thermal diffusivity

Since cocrystals 1 and 2 have a diversity of contacts, including halogen,  $\pi\cdots\pi$ , and C-H $\cdots$ F interactions, it can be anticipated that the forces retaining them are diverse in strength. Hirshfeld surface analysis was utilized to evaluate the relative efficacy of the competing weak hydrogen bonding,  $\pi\cdots\pi$ , and halogen-







Fig. 8 Crystal packing with the 3D topology of energy frameworks of cocrystal 1 (i) and 2 (ii). Red and blue colored bidirectional arrows indicate the highest and lowest thermal diffusivity direction of cocrystals 1 and 2, respectively. Red and green colored tubes indicate the electrostatic and dispersion energy, respectively.

bonding interactions (Fig. S5, ESI<sup>†</sup>). Short interactions were discovered to be distinct in the mother phase of cocrystals 1 and 2. In cocrystal 1, the F $\cdots$ H (34.6%) interaction was found to be the highest interaction supporting the three-dimensional structure, followed by I $\cdots$ H (9.8%), C $\cdots$ I (9.1%), and I $\cdots$ N (4.7%) interactions. Nevertheless, the contributions of all interactions changed in cocrystal 2. However, F $\cdots$ H (23.4%) interaction was also seen to be the highest in cocrystal 2 followed by I $\cdots$ H (15.3%) and C $\cdots$ C (11.8%) interactions. It can be inferred that the halogen and weak hydrogen bond interactions play a significant role in determining the crystal structure of 1. On the other hand, weak hydrogen bonds and  $\pi\cdots\pi$  interactions play a crucial role in maintaining the crystal structure of 2. Furthermore, energy framework analysis suggests that the C–I $\cdots$ N halogen interaction in cocrystal 1 is a strong, electrostatically dominant interaction with an energy of  $-21.1\text{ kJ mol}^{-1}$  that holds the structure together with weak C–H $\cdots$ F interactions (Fig. 8(a) and S6, ESI<sup>†</sup>). In contrast, weak electrostatic C–I/C–I interactions were found to be accompanied by the presence of the dispersion-dominated C–H $\cdots$ F and  $\pi\cdots\pi$  interactions in cocrystal 2, which have a  $-8.6\text{ kJ mol}^{-1}$  energy along the *ac* plane (Fig. 8 and S7, ESI<sup>†</sup>). The comparatively stronger interactions such as C–I $\cdots$ N, C–I, and C–H $\cdots$ F in cocrystal 1 could have promoted the superelastic deformation rather than ferroelastic deformation. Furthermore, the C–I $\cdots$ N halogen interaction tapes that make 2D sheets with the help of C–H $\cdots$ F interactions grow along the  $[0\bar{1}\bar{1}]$  direction in a symmetrical stack in cocrystal 1. This could be the reason for the highest

thermal diffusivity along the  $[0\bar{1}\bar{1}]$  direction where the robust, strong, and electrostatically dominated C–I $\cdots$ N halogen bonds determine the thermal diffusivity value. In contrast, dispersion dominated C–I/C–I and  $\pi\cdots\pi$  interactions contained in cocrystal 2 resulted in its lower thermal diffusivity than that of cocrystal 1.

## Conclusions

In conclusion, we presented two cocrystals consisting of (1,2-bis(4-pyridyl)ethane and 1,4-diodotetrafluorobenzene), and (pyrene and 1,4-diodotetrafluorobenzene), which are denoted as 1 and 2, respectively. Cocrystals 1 and 2 demonstrated superelastic and ferroelastic deformations, respectively. From the crystal structure analysis, the underlying cause of the deformation was found to be mechanical twinning. The difference in the deformation behavior was discussed through the distinct combination of strong halogen bonds, such as C–I $\cdots$ N and C $\cdots$ I, and weak interactions, such as C–H $\cdots$ F and  $\pi\cdots\pi$  interactions in the cocrystal structures. The thermal diffusivity measurements on the ferroelastic and superelastic crystals showed the axial-dependent anisotropic thermal diffusivity of the cocrystals. A 7-fold difference in thermal diffusivity was observed in superelastic crystal 1, the highest in the orthogonal plane to the shear-induced direction, whereas in crystal 2 the axial dependence was less pronounced. This study provides insights into the interplay with the variation of inter-molecular interactions in organosuperelastic and organoferroelastic



behavior and the way for further design and regulation of functional organic solids with thermo-mechanical properties.

## Data availability

All experimental, thermal diffusivity, and crystallographic data are available in the ESI.†

## Author contributions

S. R. carried out the experiments, analyzed the results, and composed the manuscript. R. M., M. R., and J. M. conducted thermal diffusivity investigations. J. M. and S. T. organized the project and edited the manuscript.

## Conflicts of interest

The authors declare no competing financial interest.

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