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# Thionated PDI supramolecular polymers: controlling aggregation mechanisms, morphology and function†

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Controlling the self-assembly of functional organic molecules provides opportunities to optimise the interlinked properties and structures of these materials for specific applications. Perylene diimide (PDI) derivative, **PDI1**, was used as a model monomer to investigate the effects of heteroatom substitution in supramolecular polymer systems. In addition to the previously reported optoelectronic effects (including shifted absorbance and quenched fluorescence), thionation was found to have a significant impact on the self-assembly behaviour of **PDI1**. The thionated analogues of this species displayed a diverse range of self-assembled morphologies, rationalised through changes in intermolecular interactions. Furthermore, thionation results in a transition from highly cooperative to isodesmic assembly mechanism. Such mechanistic control should also be applicable to many other supramolecular systems based on either hydrogen-bonding or  $\pi$ -stacking interactions, and presents an important route for the regulation of dynamic polymer properties. Finally, thionation was found to result in a substantial increase in conductivity recorded within analogous supramolecular polymer structures.

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## Introduction

In the field of supramolecular polymers (SMPs), *i.e.*, polymers connected by non-covalent, directional interactions between their constituent monomers,<sup>1</sup> significant research efforts are focused on gaining control over the assembly behavior. Several promising routes are being explored to obtain this desired control, including chemical modification<sup>2–4</sup> or altering the external environment, such as solvent or temperature.<sup>3,5–7</sup> One of the most attractive building blocks for such assemblies are perylene diimides (PDIs), also called perylene bisimides (PBIs). PDIs possess a range of useful optoelectronic properties, including high fluorescence quantum yields, high chemical and thermal stability, and N-type semiconductivity. The change and control of the assembly behaviour can have a significant influence on the properties of the resulting SMPs, especially the optoelectronic properties, depending on whether the monomers undergo H- or J-aggregation.<sup>3</sup>

The formation of one-dimensional self-assembled structures from PDI derivatives has been the subject of much research, owing largely to their fascinating array of potential applications.<sup>8</sup>

The unique properties of PDIs are well-suited to utilisation in the fields of electronics, photocatalysis, bioimaging and in thermoelectric devices, amongst others, as shown by Zhu *et al.* and others in the field.<sup>9–13</sup> More recently, the ability of PDIs to readily form anionic species has been demonstrated to offer excellent material properties, for example as thermoelectric materials and as photosensitizers for hydrogen evolution reactions (HERs).<sup>14,15</sup> The alignment of chromophores achieved in one-dimensional assemblies further enhances the desirable functionalities displayed by these materials, for example, through  $\pi$ -stacking-mediated exciton diffusion.<sup>8</sup>

In order to tune the properties, self-assembly and therefore applications, the versatility of the PDI motif has been exploited by selective functionalisation in a number of positions, for example by the laboratories of Würthner and Zhu.<sup>3,4,14–25</sup> Imide substitution is the most facile route for functionalisation, and may be used to alter the aggregation behaviour and solubility of PDI derivatives. Such changes have minimal effect on the PDI optoelectronic properties, owing to nodes in the frontier molecular orbitals at the imide-nitrogen position.<sup>10,26</sup> Conversely, substitution in the bay positions has strong substituent-dependent effects on the optoelectronic properties of the PDI chromophore.<sup>20</sup> Targeted changes at the bay positions allow tailoring of these properties for particular applications. The resultant core distortion from this substitution pattern, however, has a significant impact on the self-assembly

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behaviour, in many cases inhibiting the formation of well-defined nanostructures.<sup>27</sup>

One area that has received comparatively little attention is functionalization within the core PDI moiety. Heteroatom substitution has been applied to a number of organic semiconductors, and has proven to be a versatile strategy to optimize their properties, including solubility, film morphology, and charge carrier mobility.<sup>28–31</sup> This approach has only recently been utilised for both PDI and related naphthalene diimide (NDI) systems.<sup>32</sup>

Oxygen-to-sulphur exchange has a number of reported effects on the optoelectronic properties of these chromophores. Firstly, absorbance is bathochromically shifted significantly, resulting in a distinctive colour change. PDIs typically fluoresce very strongly with quantum yields close to unity.<sup>33–35</sup> Thionation has been shown to result in complete quenching of this fluorescence, instead forming longer-lived triplet excited states leading to phosphorescence.<sup>36</sup> These species may be desirable for their increased rate of charge separation in organic photovoltaic (OPV) devices *via* singlet fission, or for generating singlet oxygen for therapeutic applications.<sup>37–39</sup>

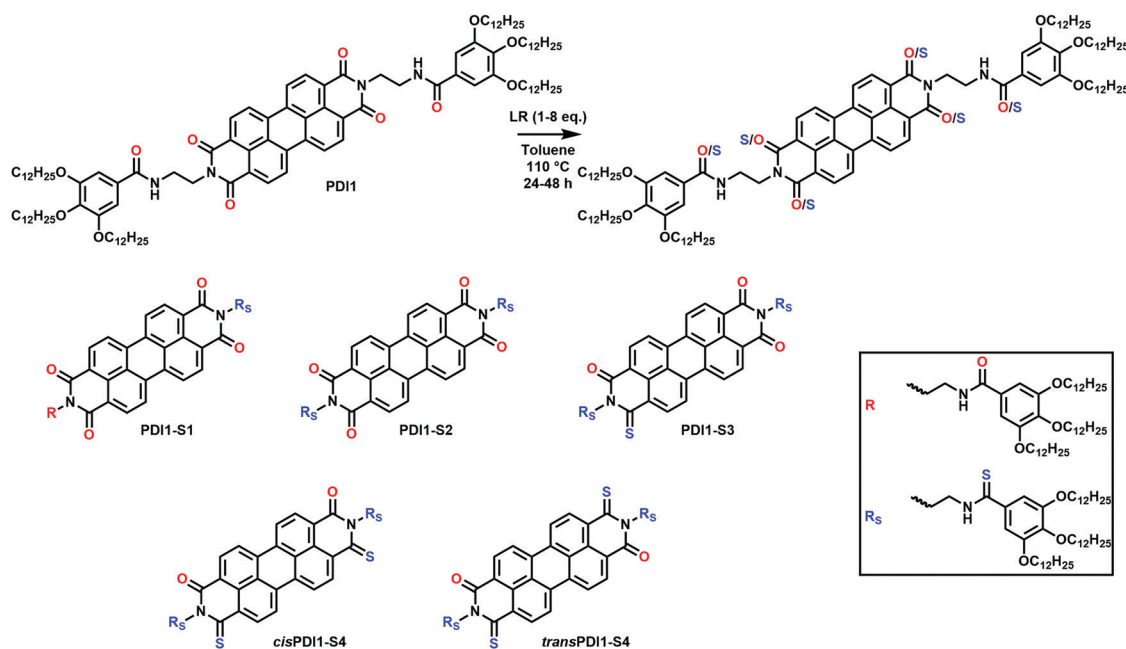
Despite the promising properties displayed by these thionated materials, a very limited number of studies have been conducted on their ability to form self-assembled structures. Differences in film-forming behaviour and solid-state absorbance spectra, however, suggest thionation has a number of effects and is worthy of further investigation. We herein present a comprehensive study on the effects of thionation on the self-assembly behaviour of a previously reported PDI derivative,<sup>5,40</sup> **PDI1**, shown in Scheme 1. We highlight the opportunities that thionation provides for fine control over the self-assembly behaviour of this known PDI derivative.

## Results and discussion

### Synthesis and characterisation

Thionation of **PDI1** was achieved by reaction of the parent PDI with Lawesson's reagent (LR) in anhydrous toluene, as described in Scheme 1.<sup>36</sup> This reaction generated a mixture of thionated products, with between one and six substitutions found in crude reaction mixtures. As demonstrated in Table S1 (ESI<sup>†</sup>), alterations to either the molar ratio of LR to **PDI1** or the reaction length could be used to favour products with differing levels of thionation.

Despite the more complex side chains employed in this study (when compared with previous studies on PDI thionation), separation of each product was successfully achieved with conventional chromatography using a combination of preparative TLC and silica gel column chromatography. By this method, products containing between one and four sulphur atoms were isolated, as shown in Scheme 1, and termed **PDI1-S1** to **PDI1-S4**, respectively. Interestingly, when four substitutions occur, two distinct regioisomers are observed, denoted *cis*-**PDI1-S4** and *trans*-**PDI1-S4**, depending on the relative positions of the two core sulphur atoms. An additional product substituted with five sulphur atoms was also isolated, however yield was insufficient to allow extensive characterisation. Notably, two products were not obtained after purification: the third *geminal* S4-substituted isomer, and the fully-thionated S6-substituted product. Findings in previous studies suggest the former is energetically unfavoured, and likely only present in trace amounts unless favoured by steric blocking of one imide group.<sup>36,41,42</sup> Peaks consistent with trace amounts of an S6-substituted product were observed in mass spectra of the crude reaction mixtures upon thionation of **PDI1**;



**Scheme 1** General conditions used for the thionation of **PDI-1** with Lawesson's reagent (LR). Chemical structures of products isolated from thionation with between one and four oxygen-to-sulphur substitutions.



however, despite extensive efforts, significant quantities of this product could not be isolated. All thionated products obtained in significant amounts were fully characterised by  $^1\text{H}$  and  $^{13}\text{C}$  NMR spectroscopy (Fig. S1–S10, ESI $^\dagger$ ) and MALDI-ToF mass spectrometry (Fig. S12–S16, ESI $^\dagger$ ). To confirm the identities of each regioisomer of **PDI1-S4**, additional  $^1\text{H}$ - $^1\text{H}$  ROESY NMR analysis was conducted on these two compounds and is shown in Fig. S11 (ESI $^\dagger$ ). In agreement with previous reports on thionated PDI systems,<sup>36,43</sup> the *trans* isomer displayed additional through-space coupling interactions between the bridgehead protons, confirming their close proximity in this compound. Comparable peaks were not observed for *cis***PDI1-S4**.

The complete absence of products exhibiting thionation of only the imide oxygen atoms indicates a remarkable preference for thionation of the side chain amide group compared with the imide positions.

### Optoelectronic properties

UV/Vis and fluorescence spectroscopy were used to investigate the optoelectronic properties of **PDI1** and its thionated analogues. Spectra, shown in Fig. 1a, were recorded in chloroform (at a concentration of  $1 \times 10^{-5}$  M) to minimise aggregation.

The absorbance spectrum of **PDI1** displays a vibronic progression typical of PDI materials, with a maximum absorbance

at 529 nm, and progressively weaker peaks at shorter wavelengths. **PDI1-S1** and **PDI1-S2** exhibit spectra almost identical to the parent PDI, with absorbance maxima at 530 and 531 nm, respectively. The absence of significant changes is consistent with substitution occurring only within the side chains, which does not affect the optoelectronic properties of the core PDI chromophore. Conversely, products containing 3 or more sulphur atoms displayed successive bathochromic shifts of their absorbance spectra, dependent on the number of thionated imide positions: **PDI1-S3** displayed a maximum absorbance at 580 nm, whereas both S4-substituted PDIs exhibited identical maxima at 622 nm. This behaviour is consistent with previous reports on NDI and PDI thionation and is attributed to a concurrent increase in energy of the HOMO and decrease in that of the LUMO, and an increase in the electron affinity associated with increasing degree of thionation.<sup>32,36,44</sup>

The typically strong fluorescence of PDI materials is completely quenched in all derivatives exhibiting any degree of core thionation, as shown in Fig. 1b. This remarkably strong quenching effect has been attributed to a reordering of electronic states upon thionation, such that inter-system crossing to a triplet excited state is facilitated.<sup>36</sup> Conversely, thionation of only the side chain amide groups resulted in a significant, though not complete, quenching as shown in the spectra of **PDI1-S1** and **PDI1-S2**. This more gradual quenching, evidenced by a successive decrease in fluorescence intensity upon thionation of neither, one or both amide groups, is likely to occur due to more conventional mechanisms, such as the heavy atom effect.<sup>45,46</sup> This effect is likely to be further exacerbated by the well-documented intramolecular hydrogen-bonding interactions between amide and imide groups within **PDI1**, resulting in a closer proximity of the heavier sulphur atoms to the PDI chromophore.<sup>5</sup>

### Self-assembled morphology

To explore differences in behaviour between **PDI1** and its thionated analogues, the solution self-assembly of all obtained derivatives was investigated in methylcyclohexane (MCH). This solvent has previously been demonstrated to strongly induce aggregation in **PDI1** and therefore serves as an ideal medium to highlight morphological differences.<sup>5</sup> TEM images of the typical morphologies shown by each derivative in this solvent are shown in Fig. 2.

As reported elsewhere, at the dilute concentrations studied here ( $5 \times 10^{-5}$  M), **PDI1** (Fig. 2a) forms networks of interlinked nanofibers, likely driven predominantly by a combination of hydrogen-bonding and  $\pi$ -stacking interactions.<sup>5,40</sup> Individual nanofibers were highly elongated, often multiple micrometres in length. Widths ranged from unimolecular strands of approximately 3 nm to ribbon-like structures more than 500 nm in width. The significant proportion of narrow supra-molecular strands suggests they are relatively stable, with aggregation into larger bundles occurring only due to the high density of nanofibers present.

The assembly of **PDI1-S1** was studied by a similar procedure, with typical morphologies observed shown in Fig. 2b. This compound also forms fibrous structures but with clear

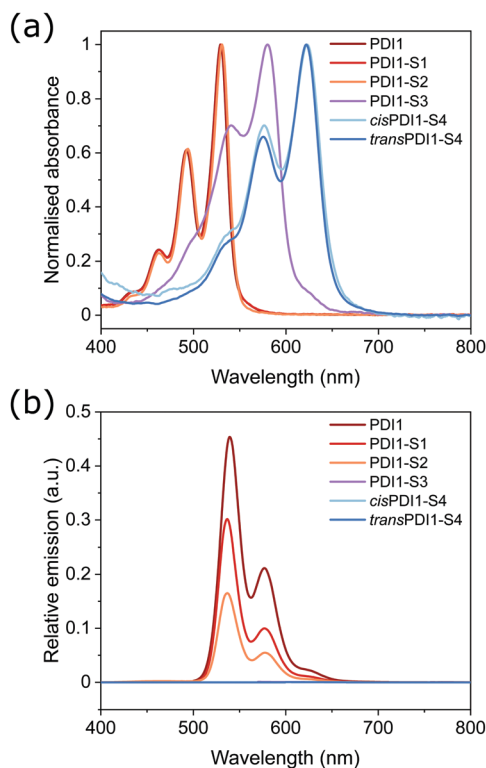


Fig. 1 UV/Vis absorbance spectra (a), and fluorescence spectra (b), of **PDI1**, **PDI1-S1**, **PDI1-S2**, **PDI1-S3**, **PDI1-S4**, *cis***PDI1-S4**, *trans***PDI1-S4**, and **PDI1-S5**. All spectra recorded at a concentration of  $10^{-5}$  M in chloroform. Excitation wavelengths were chosen based on the maxima of the corresponding absorbance; 530 nm for **PDI1**, **PDI1-S1**, and **PDI1-S2**, 580 nm for **PDI1-S3**, 622 nm for *cis***PDI1-S4** and *trans***PDI1-S4**, and 670 nm for **PDI1-S5**.



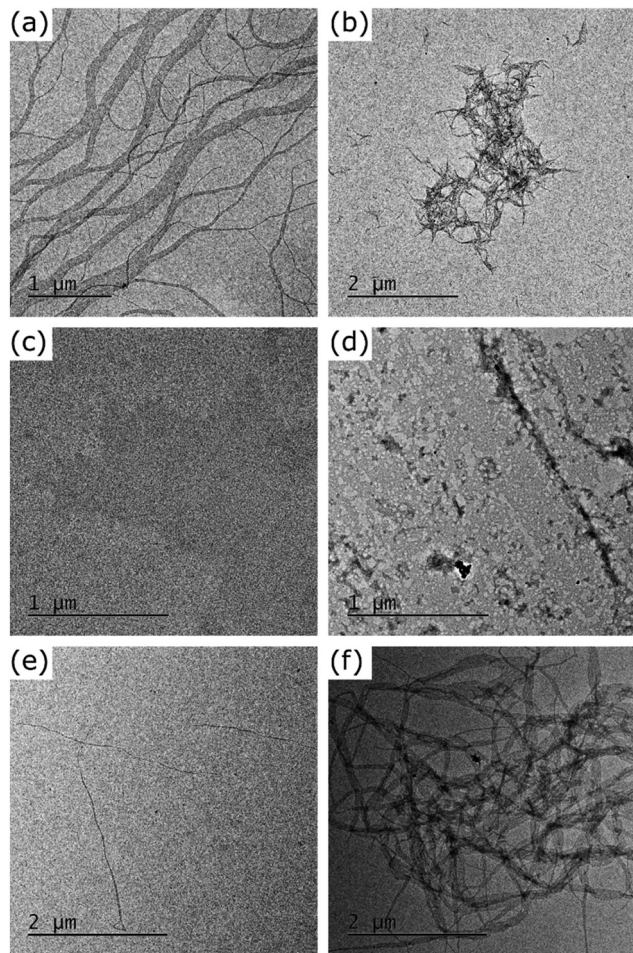


Fig. 2 TEM images of the morphologies formed by the self-assembly of **PDI1** (a), **PDI1-S1** (b), **PDI1-S2** (c), **PDI1-S3** (d), **cisPDI1-S4** (e), and **transPDI1-S4** (f), in MCH at a concentration of  $5 \times 10^{-5}$  M.

differences to those formed by the parent **PDI1**. Most notably, nanofiber lengths are significantly shorter, with typical lengths between 100 and 300 nm, and lengths greater than 500 nm uncommon.

Further increases to the degree of thionation have large effects on the self-assembly behaviour of **PDI1** derivatives. As shown in Fig. 2c, **PDI1-S2** displays no ordered nanostructures up to concentrations of  $5 \times 10^{-5}$  M in MCH. Instead, regions of increased electron density are visible in the TEM images of these samples. Despite the absence of ordered structures, the visibility of these regions by TEM without staining does not rule out self-assembly. **PDI1-S3** displays similar behaviour, with amorphous films, shown in Fig. 2d, present across the majority of the substrate. In contrast to **PDI1-S2**, however, these morphologies exhibit higher electron densities, and possess occasional regions of greater anisotropy.

Unlike lesser thionated derivatives, both S4-substituted isomers formed well-defined SMPs similar to those displayed by the parent **PDI1** (shown in Fig. 2e and f). Interestingly, significant differences between the morphologies formed by each regioisomer were evident. **cisPDI1-S4** assembles into

individual, well-dispersed supramolecular fibers. Nanofiber lengths ranged from hundreds of nanometres to multiple micrometres. The width of these fibers was predominantly between 2 and 20 nm, with a mean of 8.5 nm (width distribution shown in Fig. S17, ESI<sup>†</sup>). Conversely, nanofibers formed by the self-assembly of **transPDI1-S4**, although similar in length, possessed significantly greater widths. Typical widths of these ribbon-like structures were between 30 and 200 nm, though a limited number of thinner fibers and sections were also present, with a mean value of 70.4 nm (width distribution shown in Fig. S18, ESI<sup>†</sup>). Also notable are differences in the subsequent aggregation of the assembled nanofibers. Fibers formed by **transPDI1-S4** showed a strong tendency to aggregate and were almost exclusively present in large clusters, or in extended fibrous networks. This behaviour is in stark contrast to that of **cisPDI1-S4** nanofibers, which were dispersed (as single fibers) across the surface of the grid.

These morphological differences can be rationalised by considering changes in the relative strengths of the non-covalent interactions that lead to SMP formation. As discussed previously, the self-assembly of **PDI1** is driven by aromatic stacking interactions of the PDI core complemented by hydrogen-bonding interactions between adjacent amide groups in both side chains. Computational studies on model thioamide systems have previously revealed a decrease in hydrogen bond acceptor ability and an increase in hydrogen bond donor ability compared with the corresponding oxoamide.<sup>47</sup> For an amide group within a supramolecular polymer, acting as both donor and acceptor of hydrogen bonds, an overall decrease in interaction energy of approximately  $1.25 \text{ kcal mol}^{-1}$  upon thionation can be expected. Similar studies were carried out by Tilley *et al.* to assess the effect of imide thionation on the strength of stacking interactions between a pair of model PDI derivatives.<sup>41</sup> Thionation was found to increase the interaction energy, with each successive oxygen-sulphur substitution leading to a comparable average increase of  $1.38 \text{ kcal mol}^{-1}$ .

Based on these effects, a progressive decrease in interaction energy is expected upon the initial thionation of one or both side chain amides (in **PDI1-S1** and **PDI1-S2**, respectively) compared with **PDI1**. The shorter nanofibers and absence of anisotropic structures shown by these two derivatives are in good agreement with this decrease in interaction strength. Subsequent core thionation leads to a successive increase in interaction strength between adjacent unimers. Notably, both S4-substituted PDI derivatives are expected to display similar interaction energies to the parent **PDI1**. Morphological studies corroborate this behaviour, with all three systems displaying similar nanofibrous structures. The expected changes in interaction energy upon thionation are summarised graphically in Fig. S19 (ESI<sup>†</sup>).

The apparent similarities in the morphologies formed by **PDI1**, **cisPDI1-S4** and **transPDI1-S4** prompted further investigation of these species by AFM, with typical structures observed shown in Fig. 3. On carbon-coated mica substrates, AFM revealed comparable nanofiber structures to those observed on the carbon-coated TEM grids used. Apparent increases in widths compared with TEM images are likely to be a result of either slight differences in



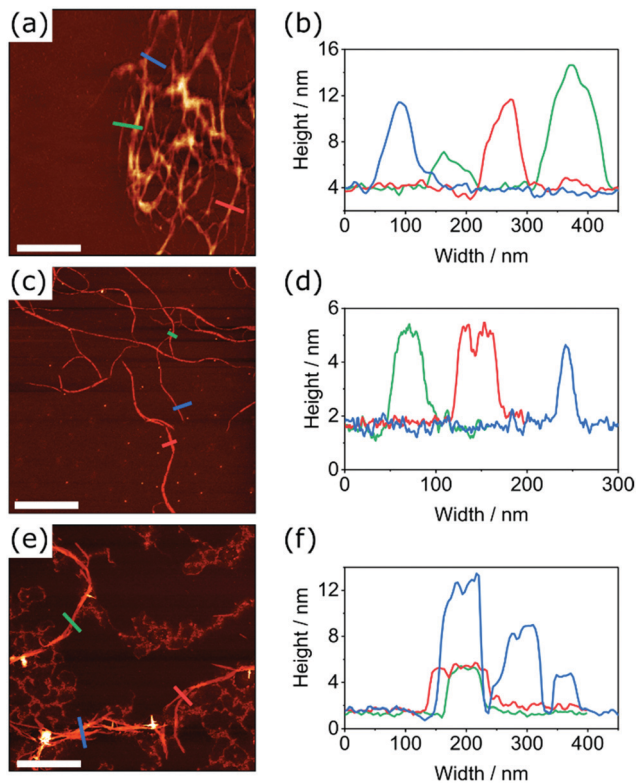


Fig. 3 AFM height images of the morphologies formed by the self-assembly of **PDI1** (a) *cisPDI1-S4* (c), and *transPDI1-S4* (e) in MCH at a concentration of  $5 \times 10^{-5}$  M. Height profiles along each corresponding coloured line, (b, d and f). Scale bars = 1  $\mu$ m.

sample preparation methodology, or peak broadening artefacts. All three PDI derivatives studied formed nanofibers with similar minimum heights of between 2.5–3.5 nm, likely indicative of supramolecular strands of unimolecular thickness.

In agreement with previous reports, **PDI1** nanofibers were also found to display regions of greater width and height which likely arise due to lateral aggregation of individual supramolecular strands.<sup>5,40</sup> Nanofibers of *transPDI1-S4* showed similar behaviour with distinct heights of 2.5–3.5, 6–7 or 10–11 nm observed across the majority of structures. This incremental increase in height likely indicates overlapping of multiple fibers of unimolecular thickness. In contrast to the other two species, *cisPDI1-S4* nanofibers showed exclusively unimolecular heights of 2.5–3.5 nm, regardless of width or length.

The unexpectedly dissimilar self-assembly behaviour of regioisomers *cisPDI1-S4* and *transPDI1-S4* may be attributed to the effects of heteroatom substitution on the aforementioned lateral aggregation of nanofibers. Relatively strong interactions between divalent sulphur atoms and aromatic systems are well-documented, most commonly in the context of proteins.<sup>48–51</sup> Non-covalent interactions between Group 16 elements, often referred to as chalcogen bonding, are increasingly utilised in a range of supramolecular systems and may display strengths comparable with conventional hydrogen bonds.<sup>52,53</sup> The strength of such interactions increases considerably from oxygen to sulphur and other larger chalcogen atoms.<sup>54</sup>

We postulate that both aforementioned interactions contribute to the aggregation of individual co-facially-stacked supramolecular strands into larger bundled structures. Additional S–S and S– $\pi$  interactions should encourage this type of lateral aggregation on the sulphur-substituted edges of thionated **PDI1** derivatives. In *transPDI1-S4*, sulphur atoms are positioned on opposite edges of the PDI motif and are therefore always exposed, leading to the formation of wider structures with a tendency to aggregate further into fibrous networks. For *cisPDI1-S4*, both sulphur atoms are positioned on the same edge of the PDI core. Upon lateral aggregation of individual strands, both sulphur atoms may be positioned internally, inhibiting further bundling of these structures. Such behaviour supports the remarkable preference of this derivative for nanofibers less than 10 nm in width, compared to those with widths almost an order of magnitude greater formed by *transPDI1-S4*.

### Mechanism of self-assembly

Unlike their covalent counterparts, the formation mechanisms of SMPs have significant effects on the properties of the resultant materials.<sup>55</sup> Two predominant mechanisms exist; isodesmic, where the binding constant is independent of polymer length, and cooperative, where two distinct phases are present, characterised by different binding constants.<sup>56</sup> Changes in the degree of aggregation as a function of temperature allow for the most reliable identification of polymerisation mechanism.<sup>57</sup> In this study, we therefore employed variable-temperature UV-Vis spectroscopy measurements to investigate the effects of thionation on the mechanism of self-assembly.

Changes in the absorbance profiles of **PDI1** and *transPDI1-S4* in MCH upon heating from 293 to 373 K are shown in Fig. 4. A concentration of  $5 \times 10^{-6}$  M was selected, such that near complete aggregation was observed at the lowest temperature, and a completely unimeric sample obtained at the highest temperature. Changes in absorbance for the remaining thionated derivatives are shown in Fig. S20 (ESI<sup>†</sup>). Prior to heating, all investigated materials displayed similar differences to the corresponding unimeric spectra shown in Fig. 1;  $\lambda_{\max}$  for each was shifted hypsochromically (by approximately 30–70 nm with respect to the corresponding unimer), and a new bathochromically shifted shoulder was apparent (at a wavelength 20–30 nm longer than the  $\lambda_{\max}$  of the corresponding unimer). These changes are consistent with H-aggregation, with the partially allowed J-band resulting from rotational displacement of PDI cores.<sup>58</sup>

Upon heating, spectral changes associated with disaggregation were observed for all samples. The change in absorbance at the  $\lambda_{\max}$  of the corresponding unimeric spectra was used to determine the degree of aggregation at each temperature for each sample using eqn (S1) (ESI<sup>†</sup>). Degree of aggregation was plotted as a function of temperature for each sample in Fig. 5. Data for each compound were fitted to either the cooperative model proposed by Smulders *et al.*,<sup>57,59</sup> or to a sigmoidal curve (indicative of an isodesmic mechanism<sup>60</sup>) depending on which gave the lowest  $R^2$  parameter. Samples best modelled by a cooperative mechanism



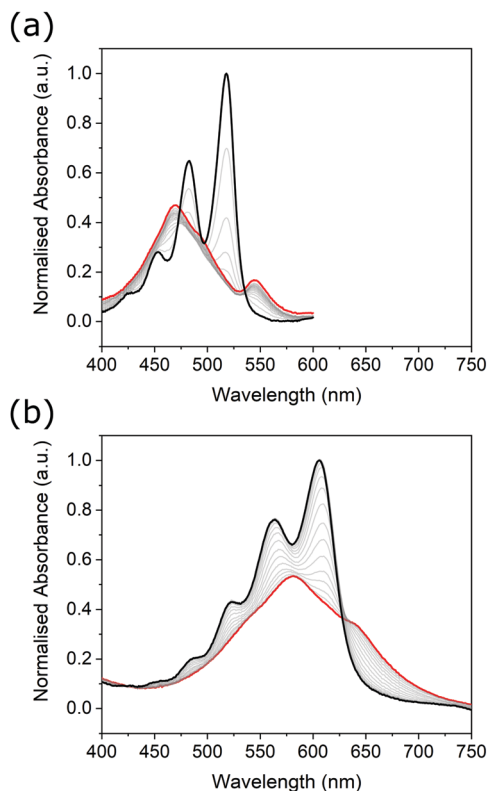


Fig. 4 Changes in the UV/Vis absorbance spectra of **PDI1** (a), and **trans-PDI1-S4** (b), upon heating from 293 K (red profiles) to 373 K (black profiles) in 5 K intervals. Spectra recorded at a concentration of  $5 \times 10^{-6}$  M in MCH.

were initially fitted by eqn (S2) (ESI<sup>†</sup>) at temperatures below nucleation, and eqn (S3) (ESI<sup>†</sup>) at higher temperatures.

In agreement with previous reports,<sup>5</sup> **PDI1** displays a highly cooperative mechanism, characterised by a sharp transition between nucleation and elongation. Specifically, a very steep elongation curve is observed, with near complete aggregation persisting at a relatively high temperature of approximately 340 K. For the initial two sulphur substitutions (*i.e.*, **PDI1-S1** and **PDI1-S2**), the cooperative assembly mechanism of **PDI1** is maintained. A clear transition between nucleation and elongation is visible, depicted by a change of symbol (black squares to red circles) in Fig. 5b and c. Successive thionation for these two derivatives, however, resulted in a significant reduction in the steepness of the elongation curve. Both derivatives therefore show incomplete aggregation, even at the lowest temperature studied.

Thermodynamic parameters extracted directly from the fitting function reveal a decrease in the magnitude of  $\Delta H_e$ , the enthalpic change during elongation, from  $-137.8$  kJ mol<sup>-1</sup> for **PDI1** to  $-45.1$  and  $-22.6$  kJ mol<sup>-1</sup> for **PDI1-S1** and **PDI1-S2**, respectively. Eqn (S4) and (S5) (ESI<sup>†</sup>) were used to estimate the degree of polymerisation. At 293 K, a degree of polymerisation  $> 10\,000$  was determined for **PDI1**, alongside values of 390 for **PDI1-S1** and 290 for **PDI1-S2**. Full thermodynamic parameters for cooperatively fitted systems are given in Table S2 (ESI<sup>†</sup>).

For systems best modelled by an isodesmic mechanism, the equilibrium constant,  $K_e$ , was estimated using eqn (S6) and (S7) (ESI<sup>†</sup>) at each temperature value. Thermodynamic parameters

for polymerisation were then determined *via* Van't Hoff plots, as shown in Fig. S21 (ESI<sup>†</sup>), and described by eqn (S8) and (S9) (ESI<sup>†</sup>). Compared with **PDI1-S2**, the magnitude of  $\Delta H$  increased considerably for **PDI1-S3** to a value of  $-78.7$  kJ mol<sup>-1</sup>. Further increases in magnitude were observed for **cisPDI1-S4** and **transPDI1-S4**, with values of  $-91.8$  and  $-103.5$  kJ mol<sup>-1</sup>, respectively. The degree of polymerisation at 293 K for each sample was estimated using eqn (S6) (ESI<sup>†</sup>). **PDI1-S3** displayed a degree of polymerisation of 3.5, **cisPDI1-S4** a value of 3.2, and **transPDI1-S4** a value of 7.0. Full thermodynamic parameters derived for compounds that polymerised *via* isodesmic mechanisms are given in Table S3 (ESI<sup>†</sup>).

The overall trends in the enthalpic changes determined here are in good agreement with those predicted based on differences in intermolecular interactions upon thionation. The enthalpy loss for the polymerisation of **PDI1** decreases significantly upon the initial thionation of side chain amide groups, indicative of a decrease in the strength of hydrogen-bonding interactions. Further thionation of the core imide groups then leads to enhancement of stacking interactions, reflected by a successive increase in the magnitude of  $\Delta H$  from S2- to S4-substituted **PDI1** analogues. The more negative value of  $\Delta H$  displayed by the *trans* S4-substituted isomer compared to the *cis* isomer further supports the formation of additional energetically favourable S-S or S- $\pi$  interactions, leading to lateral aggregation as described above. Changes in the estimated degree of polymerisation followed a similar pattern for **PDI1**, **PDI1-S1**, and **PDI1-S2**; however, the remaining three compounds investigated all displayed low values between of 3 and 7. Such behaviour is typical of isodesmic systems, where a significant proportion of the material may remain in a unimeric or oligomeric state.

Overall, a clear transition from a highly cooperative polymerisation process to one that proceeds isodesmically is evident upon successive oxygen-to-sulphur substitutions. This behaviour may also be rationalised by changes in the relative strengths of noncovalent interactions. A common source of cooperativity in SMP formation is the electronic influence in hydrogen-bonding interactions along a polymer chain.<sup>56</sup> Conversely, supramolecular polymers formed predominantly by  $\pi$ -stacking interactions typically display isodesmic formation mechanisms.<sup>56</sup> As previously discussed, amide thionation reduces overall hydrogen-bonding ability, whereas thionation of PDI imide positions increases their tendency to  $\pi$ -stack. Thionation in either position would therefore be expected to cause a shift from a binding mode with a large hydrogen-bonding contribution to one based largely on aromatic stacking. This change is consistent with the gradual transition of observed mechanism from cooperative to isodesmic upon thionation.

### Electrostatic and conductivity measurements

The semiconductivity displayed by PDI-based materials is important in a range of applications. To assess if changes in conductivity were exhibited within individual SMPs upon thionation, we performed both electrostatic force microscopy (EFM) and PeakForce tunnelling AFM (PF-TUNA) on self-assembled



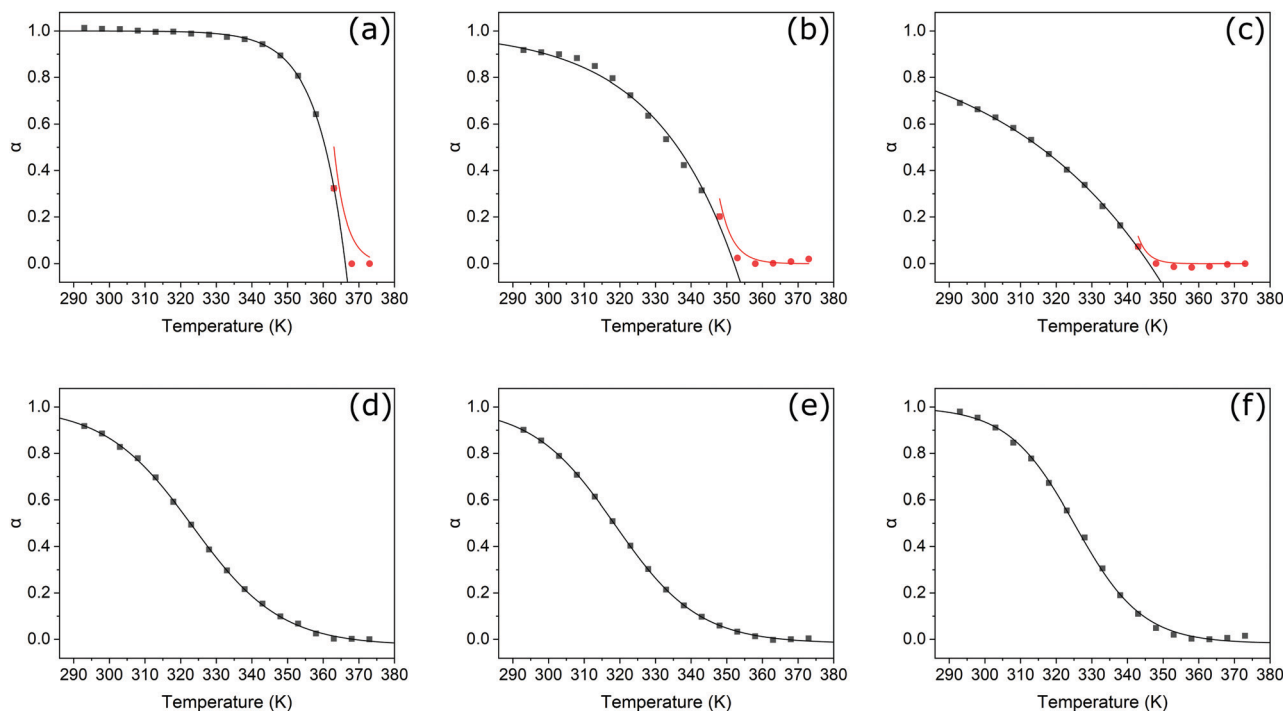


Fig. 5 Changes in degree of aggregation,  $\alpha$ , as a function of temperature in K, and fitting with either cooperative or isodesmic models, for **PDI1** (a), **PDI1-S1** (b), **PDI1-S2** (c), **PDI1-S3** (d), **cisPDI1-S4** (e), and **transPDI1-S4** (f). For species best fitted with a cooperative mechanism, data are separated into the nucleation phase (red circles), and elongation phase (black squares). Fitting curves for each regime are shown in the corresponding colour.

samples of **PDI1** and **cisPDI1-S4**. Comparison of electronic properties of these two systems (*i.e.*, control **PDI1** and thionated **cisPDI1-S4**) is possible as the fiber-like morphology of these compounds upon self-assembly is very similar and the non-contact lift mode and  $< 1$  nN force regime of EFM and PF-TUNA, respectively, ensure minimal disruption to the molecular organisation within the self-assembled architectures.

When a conductive cantilever moves through an electrostatic field a force is induced dependent on the magnitude of the field. If oscillating at its resonance frequency, the induced force will alter the resonance creating a phase shift in the cantilever's motion. The concurrent measurements of the topography and electrostatic interaction (detailed methodology given in ESI<sup>†</sup>) for **PDI1** and **cisPDI1-S4** are shown in Fig. 6a and b, respectively, as 3-dimensional surfaces overlaid with the induced phase shifts as blue colour maps. Data are shown for short sections of nanofibers with comparable heights of 3.5–4.5 nm for each compound. Measured with the same conductive cantilever, qualitative comparison of the magnitude of the electrostatic field is possible when the induced phase shifts are normalised to the carbon background. Phase profiles, extracted orthogonally to nanofiber long-axes are presented in Fig. 6c, demonstrating a greater contrast in phase between nanofiber and substrate, and so greater electrostatic field, for thionated **cisPDI1-S4** compared with the parent **PDI1**.

The currents measured through thionated **cisPDI1-S4** and the parent **PDI1** by PF-TUNA under 500 mV potential are presented in Fig. 6d. While the potential was kept low to prevent possible damage to the delicate assemblies in the

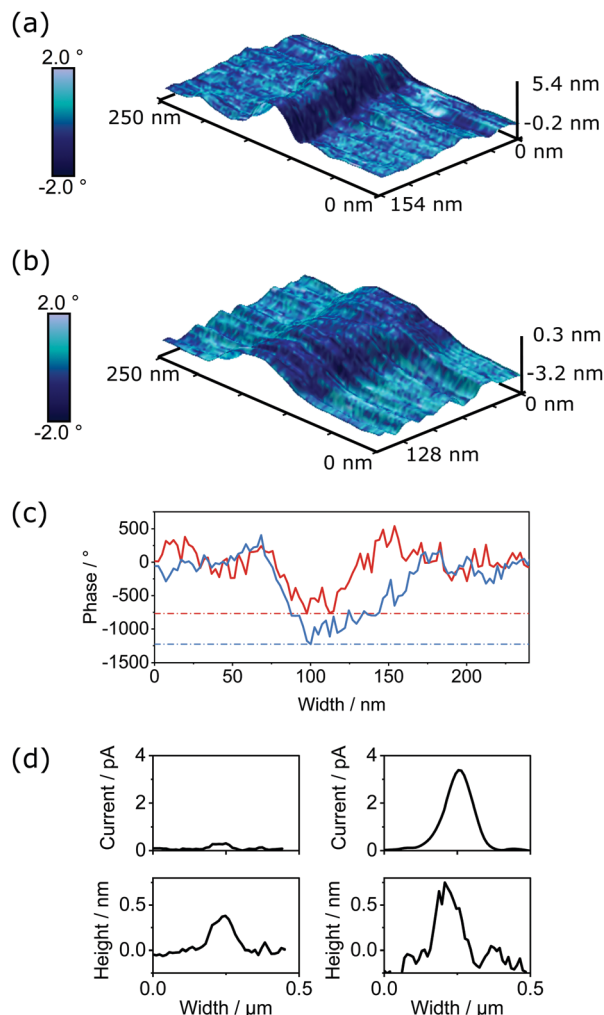
confined geometry beneath the cantilever tip it can clearly be observed that for a comparable height of fiber **cisPDI1-S4** exhibits a greater degree of conductivity than that of parent **PDI1**. In light of the difference in electrostatic potential displayed by the two comparable supramolecular structures, it appears reasonable to infer that the greater level of conductivity observed in **cisPDI1-S4** by PF-TUNA occurs upon the thionation processes described here.

PF-TUNA measurements were also carried out on a sample of **transPDI1-S4** nanofibers. Peak current values observed for this compound, and the two other species measured by this method are compared in Fig. S22 (ESI<sup>†</sup>). For **transPDI1-S4**, we measured peak current values between those observed for **PDI1** and **cisPDI1-S4**. It should, however, be noted that the current values obtained are dependent on many factors including the morphology of the sample. We have chosen to focus our comparison on **PDI1** and **cisPDI1-S4** as these compounds display similar self-assembled morphologies, meaning conductivity differences are likely a direct result of the oxygen-to-sulphur substitution. The notably different morphology displayed by **transPDI1-S4**, described above, could have a substantial influence on the peak current recorded for this compound.

## Conclusions

In this work, we present a comprehensive investigation into the effects of thionation on PDI-based SMPs. Thionation provides a facile method by which to tune the optoelectronic properties of





**Fig. 6** 3D micrographs showing combined topographical (vertical axis) and phase data (colourmap) obtained by EFM measurements on self-assembled nanofibers of **PDI1** (a), and **cisPDI1-S4** (b). (c) Corresponding phase profiles extracted from EFM data for **PDI1** (solid red line) and **cisPDI1-S4** (solid blue line). Data are normalised such that the mean background phase is  $0^\circ$  for both samples. Dashed lines of the corresponding colour indicate the greatest phase contrast compared with the substrate for each sample. (d) Height and current profiles extracted from PF-TUNA measurements for samples of **PDI1** (left) and **cisPDI1-S4** (right) nanofibers.

these materials by simple stoichiometric adjustments. Substitution within side-chain amide groups largely retains the optoelectronic properties of the parent **PDI1**, whereas core substitution results in significant changes. Namely, fluorescence is quenched, and absorbance is bathochromically shifted by approximately 50 nm for each successive substitution. Such readily adjustable optoelectronic properties could facilitate the tuning of supramolecular constructs for specific applications, for example to better match the solar spectrum in photovoltaic devices.<sup>61</sup> Moreover, the formation of the longer-lived triplet excited states responsible for the fluorescence quenching may allow enhancement of exciton transport distances within self-assembled structures.<sup>36</sup>

Profound changes in the self-assembly behaviour of these materials were also observed. Thionation has significant effects on the morphology of the self-assembled structures and the ability of each compound to form linear SMPs. Differences in behaviour are rationalised by considering changes in binding energy upon thionation, and the introduction of additional S-S and S- $\pi$  interactions. Mechanistic investigations reveal that thionation evokes a transition from a highly cooperative to an isodesmic assembly model. Such mechanistic control should also be applicable to many other supramolecular systems based on either hydrogen-bonding or  $\pi$ -stacking interactions, and presents an important tool for the future design and regulation of dynamic polymer properties.<sup>62</sup>

Overall, the approach described provides a facile route to simultaneously exert control over the mechanism, morphology and functionality of self-assembled **PDI** materials. Thionated **PDI**s already show great promise in OFET devices,<sup>41</sup> and further optimisation of their self-assembly, such as the work described here, should facilitate their application in the diverse range of fields for which one-dimensional **PDI**-based materials are well suited.<sup>8</sup>

## Experimental

### Synthesis

**Synthesis of PDI-1.** **PDI1** was synthesized using an adapted method based on that reported by Ogi *et al.*<sup>5,40</sup>

### General procedure for thionation of **PDI1**<sup>36</sup>

**PDI1** was dried into a flask with Lawesson's reagent (1–8 eq.) and placed under a nitrogen atmosphere. Toluene (anhydrous, 2 mL) was added and the mixture heated to reflux for 24–48 hours. The mixture was cooled to room temperature, the solvent removed under reduced pressure, and the product purified by preparative TLC (silica, chloroform:toluene), followed by silica gel microcolumn chromatography (chloroform:hexane) to afford the product. Please see ESI† for detailed methodologies, conditions (p4–6) and associated spectra (Fig. S1–S16, ESI†).

### Self-assembly of **PDI1** and thionated analogues

**PDI1** and its thionated analogues were each dispersed in degassed MCH at concentrations of  $5 \times 10^{-5}$  or  $5 \times 10^{-6}$  M under nitrogen atmosphere. Samples were heated briefly to  $90^\circ\text{C}$  until dissolved, allowed to cool to room temperature, then aged for at least 24 hours prior to microscopy studies or spectroscopic analysis. Please see ESI† for detailed sample preparation and analysis methodologies (p2–3, p23–24).

All relevant underlying data are provided as ESI† accompanying this paper.

## Author contributions

Henry E. Symons: conceptualization, formal analysis, investigation, methodology, validation, visualization, writing – original draft; Maximilian J. L. Hagemann: investigation, validation, writing – review





and editing; Robert L. Harniman: investigation, methodology, resources, writing – review and editing; Charl F. J. Faul: conceptualization, project administration, resources, supervision, writing – review and editing.

## Conflicts of interest

There are no conflicts to declare.

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