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## Effect of solvent isomers on the gelation properties of tri-aryl amine organogels and their hybrid thermoreversible gels with poly[vinyl chloride]<sup>†</sup>

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The gelation properties of tri-aryl amine (TATA) in two isomers of dichlorobenzene, namely *ortho*-dichlorobenzene (*o*-DCB) and *meta*-dichlorobenzene (*m*-DCB), have been studied by calorimetry for mapping out the temperature–concentration phase diagram. It is shown that both systems behave differently, yet both form molecular compounds. The occurrence of these compounds is further demonstrated by neutron diffraction in *o*-DCB. Depending on the isomer used, X-ray investigations highlight a difference in the position of the diffraction peaks. The morphology observed by scanning electron microscopy reveals fibrillar systems in both cases, yet the fibrils' cross-sections are far larger in *m*-DCB with respect to *o*-DCB. The body of these results is discussed in light of current opinions on the role of the solvent. Ternary gels obtained with PVC have also been investigated by means of the same experimental approach. Calorimetry investigations and X-ray diffraction data show no significant difference for the behavior of the TATA moiety. The morphology studied by scanning electron microscopy allows one to distinguish clearly the TATA fibrils and the PVC network. This again confirms the possibility of preparing functional materials with an organogelator and a polymer through the making of hybrid gels.

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## Introduction

The phenomenon of organogelation is a topical subject as it raises many questions as to the mechanisms responsible for the formation of a three-dimensional network that can retain up to 99% of solvent.<sup>1–5</sup> Also, their potential applications in various domains arise continuously growing interest.<sup>6,7</sup> The term “network” is important as it conveys the notion of a morphology made up with elongated structures such as fibrils.<sup>4</sup>

The possibility of predicting whether a given molecule is to produce a gel in a given solvent<sup>8–13</sup> stands among the different concerns frequently expressed. As suggested by Guenet,<sup>4</sup> a potential organogelator must be a “chimera type” molecule, namely a molecule bearing different types of interactions such as hydrogen bonding, van der Waals interaction,  $\pi$ – $\pi$  interaction, and the like. This can favour the much faster growth of one of the crystalline faces, which generates fibrils instead of

3-D crystals. For instance, in the case of OPV molecules the gel fibrillar structures are produced by the fastest growth of the face involving the  $\pi$ – $\pi$  interaction.<sup>4,7</sup>

The other player in the gelation process is the solvent. Recently, the Hansen parameter has been used abundantly in an attempt to predict whether a gel will form or not in a given solvent.<sup>8–13</sup> Although this parameter can indicate whether a solvent is not too good so as to allow structure formation, nor too bad thus preventing solubilization, it does not give indications on the gel thermodynamic properties nor on its morphology. An example is given by triarylamine tris-amide (TATA) organogels obtained in two solvents of very close Hansen parameters,<sup>13</sup> namely bromobenzene (20.3) and tetrachloroethane (19.8),<sup>8</sup> which possess melting temperatures some 40–60 °C apart.<sup>14,15</sup> Also, the occurrence of TATA molecular compounds has recently been demonstrated in these solvents.<sup>16</sup> The Hansen parameter cannot predict the formation of molecular compounds between the solvent and the organogelator, something reminiscent of the case of polymer thermoreversible gels where the Flory's parameter is also not relevant. Indeed, poor solvents, such as *trans*-decahydronaphthalene, unexpectedly produce molecular compounds with syndiotactic polystyrene and isotactic polystyrene.<sup>17</sup> In fact, the shape of the

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solvent can be a decisive factor, as particularly revealed first in the case of polyethylene oxide compounds that form with *para*-dihalogenobenzene<sup>18,19</sup> but not with *meta*-dihalogenobenzene nor *ortho*-dihalogenobenzene.

Clearly, the organogelation phenomenon is more complex than anticipated, and therefore requires a deeper investigation into the role of the solvent.<sup>20,21</sup> For this purpose, we report here on the effect of the shape of the solvent on the gelation of triarylamine tris-amide (TATA) in two isomers of dichlorobenzene, namely *ortho*-dichlorobenzene and *meta*-dichlorobenzene. It is shown that these solvents of the same Hansen solubility parameter produce organogels with differing thermodynamic properties, molecular structures and morphologies. The case of hybrid materials made up with PVC thermoreversible gels and TATA organogels in these isomers is also tackled.

## Experimental section

### Materials

The fibrillar organogels were obtained from solutions of *tri-aryl tri-amines* designated as TATA in what follows (see Fig. 1). The synthesis and properties of these molecules are extensively described in ref. 22–25.

Of particular interest are the conductivity properties of its organogels, especially in chlorinated solvents.<sup>23,24</sup> Also, it has been shown that small amounts of TATA to PVC thermoreversible gels increase drastically the elastic modulus beyond Voigt's upper limit.<sup>14</sup>

The poly[vinyl chloride], PVC, used for the preparation of the hybrid systems was purchased from Sigma-Aldrich and was used without further purification. From SEC in THF at 25 °C (universal calibration) the weight average molecular weight was found to be  $M_w = 7.9 \times 10^4 \text{ g mol}^{-1}$  with a polydispersity index  $M_w/M_n = 1.87$ . The fractions of tetrads were found to be sss = 41% ssi, iss, sis, ... = 39% and iii = 19% as determined from <sup>13</sup>C-NMR.

*Ortho*-Dichlorobenzene (*o*-DCB), *meta*-dichlorobenzene (*m*-DCB) were purchased from Sigma-Aldrich (purity grade 99.5%), and were used as-received. Deuterated *ortho*-dichlorobenzene

(*o*-DCB<sub>D</sub>) was purchased from Eurisotop, and also used as-received.

The preparation of the binary TATA/DCB gels and of the hybrid networks (PVC/TATA/DCB) consists in preparing mixtures at the desired concentration and composition, and then heating them up to 120–160 °C until clear, homogeneous solutions are obtained. Gels are produced by quenching these solutions at low temperature (0 °C down to –20 °C).

TATA concentration was varied from  $0.5 \times 10^{-2} \text{ g cm}^{-3}$  (or  $0.004 \text{ g g}^{-1}$ ) to  $1.04 \times 10^{-2} \text{ g cm}^{-3}$  (or  $0.065 \text{ g g}^{-1}$ ). Three PVC concentrations have been studied:  $4.82 \times 10^{-2} \text{ g cm}^{-3}$  (or  $3.0 \times 10^{-2}$  in  $\text{g g}^{-1}$ ), designated as PVC5%,  $9.3 \times 10^{-2} \text{ g cm}^{-3}$  (or  $5.8 \times 10^{-2}$  in  $\text{g g}^{-1}$ ) designated as PVC10%, and  $0.135 \text{ g cm}^{-3}$  (or 0.10 in  $\text{g g}^{-1}$ ) designated as PVC15%.

**Differential scanning calorimetry.** The gel formation, the gel melting temperatures, and the corresponding enthalpies of the TATA-C11 moiety were determined by means of a DSC 8500 from PerkinElmer. Three heating and cooling rates were systematically used, namely 5 °C min<sup>-1</sup>, 10 °C min<sup>-1</sup> and 15 °C min<sup>-1</sup> within a temperature range from –20 °C to 160 °C.

About 30–40 mg of gel were introduced into stainless steel pans that were hermetically sealed by means of an *o*-ring. A first run was systematically performed for erasing the sample history, as well as to suppress parasitic mechanical effects at the gel melting. The weight of the sample was systematically checked after the different cycles in order to evaluate any solvent loss.

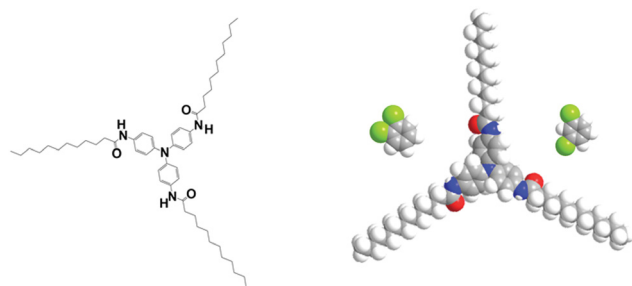
Although PVC gel formation arises from the crystallization of the longest syndiotactic sequences, the thermal behaviour of its gels is not detectable by DSC, an observation already reported in several papers.<sup>26,27</sup> Only the first run gives the melting range, while the enthalpy is highly overestimated. Consequently, PVC gel formation was determined by the tube tilting. This further implies that the first order transition observed by DCS on the hybrid systems arise from the only TATA moiety.

### Scanning electron microscopy

A piece of a gel from PVC/DCB or PVC/TATA/DCB was placed onto a cryo-holder and quenched by a plunge into a nitrogen slush. After transfer of the sample into the Quorum PT3010 chamber attached to the microscope, the sample was coated by sputter deposition with a thin Pt layer, and then fractured with a razor blade. Subsequent etching was performed at  $T = -70 \text{ °C}$  so as to expose the details of the morphology. Samples were eventually transferred in the FEG-cryoSEM (Hitachi SU8010). Images were taken with the SE-in lens detector at 1 keV and at  $T = -150 \text{ °C}$ .

### X-Ray diffraction

X-Ray diffraction experiments were performed using a diffractometer developed by the Institut Charles Sadron Différix platform. The instrument operates with a pinhole collimator and a hybrid photon counting detector (HPC-Dectrics Pilatus<sup>®</sup>3 R 300 K). The monochromatic beam ( $\lambda = 0.154 \text{ nm}$ ) is obtained by reflection of the primary beam produced by means of an X-ray



**Fig. 1** left: Chemical structure of the tris-amide triarylamine C11 in its completely extended conformation with the phenyl group tilting.<sup>22</sup> This molecule can be inscribed in a cylinder of radius  $r \approx 2.1 \text{ nm}$ . Right space-filling model of the same molecule together with the two conformers of dichlorobenzene at the same scale; left, *ortho*-dichlorobenzene (*o*DCB) and right, *meta*-dichlorobenzene (*m*-DCB).



generator from Rigaku (Micromax 007HF) onto a confocal mirror with multilayer coating (Confocal Max-Flux™ Optic, Rigaku). The size of the beam incident on the sample is about 800 μm. The distance between the sample and the detector is set in such a way as to access scattering vectors  $q = 4\pi\sin(\theta/2)/\lambda$  ( $\theta$  = diffraction angle) ranging from  $q = 0.4$  to  $10 \text{ nm}^{-1}$ . Data reduction is performed following the standard procedure for isotropic scattering. Calibration of the scattering vector range is performed with a silver behenate sample. Sample holders consist of cells of 1 mm thick with calibrated mica windows. Typical counting times are about 10 mins.

### Neutron diffraction

Neutron diffraction experiments were carried out at Institut Laue-Langevin on the D16 camera.<sup>28</sup> This camera is equipped with a helium detector of  $320 \times 320 \text{ mm}^2$  with  $1 \times 1 \text{ mm}^2$  pixels. The detector can rotate so as to access larger  $q$  values. In these experiments three angles were used:  $0^\circ$  for the transmission measurements, and  $10.8^\circ$  and  $27^\circ$  for the determination of the diffracted intensities. The neutron wavelength is selected by reflection by a focussing pyrolytic graphite monochromator. Experiments reported herein were obtained with a neutron wavelength of  $\lambda = 0.4454 \text{ nm}$ , providing the following  $q$ -range with the two-angle set-up:  $0.6 \text{ nm}^{-1} \leq q \leq 9 \text{ nm}^{-1}$  (further details are available at <https://www.ill.eu/users/instruments/instruments-list/d16/characteristics>). The sample holder temperature can be controlled with an external bath. Four temperatures were used:  $T = 20^\circ \text{C}$ ,  $T = 50^\circ \text{C}$ ,  $T = 70^\circ \text{C}$ , and  $T = 85^\circ \text{C}$ .

The samples were prepared in closed quartz cells of 2 mm thickness purchased from B. Thuet (Blodelsheim, France). The required amounts of TATA and PVC powders are first introduced, and then mixed by stirring vigorously prior to the addition of the solvent. The mixture is finally heated up to  $160^\circ \text{C}$ . This procedure allows one to prepare more easily homogeneous solutions within a few minutes. Gels were formed by cooling the solutions in a fridge kept at  $4^\circ \text{C}$ . Gel samples were systematically prepared in the same way with hydrogenous *o*-dichlorobenzene and perdeuterated *o*-dichlorobenzene for testing the existence of TATA/*o*-dichlorobenzene molecular compounds. These experiments were restricted to *o*-DCB as deuterated *m*-DCB was not available.

## Results and discussion

### (1) Thermodynamic properties of tri-aryl-triamine binary gels

The thermodynamic properties have been studied by DSC. Typical thermograms for TATA/*o*-DCB and TATA/*m*-DCB are displayed in Fig. 2. These thermograms differ significantly with the DCB isomer.

In the case of *m*-DCB, a maximum of three endotherms is observed while up to five endotherms are seen with *o*-DCB. From the endotherms the temperature-concentration phase diagrams have been mapped out as shown in Fig. 2. They reveal in *m*-DCB that the endotherm at  $T = 50 \pm 1^\circ \text{C}$  is a

non-variant event. The endotherm at  $T = 109 \pm 1^\circ \text{C}$  is also a non-variant event up to  $X_{\text{TATA}} = 0.036$ , and then increases. As to the terminal endotherm, it increases monotonously with TATA concentration.

In the case of *o*-DCB the endotherms at  $T = 16 \pm 1^\circ \text{C}$ ,  $T = 37 \pm 1^\circ \text{C}$  and  $T = 61 \pm 1^\circ \text{C}$  represent non-variant events, while the terminal endotherm, which corresponds to the gel macroscopic melting, increases with TATA concentration as also observed with *m*-DCB.

The endotherm at  $T = 85^\circ \text{C}$  is a non-variant event up to  $X_{\text{TATA}} = 0.022$ , and then also increases beyond this concentration. It is worth noting that the terminal gel melting is some  $10^\circ \text{C}$  higher in *m*-DCB than in *o*-DCB. As is customary, the dotted lines in both phase diagrams as well as the italics stand for possible extensions and for plausible phases that fulfil Gibbs phase rules.<sup>29</sup>

In the case of TATA/*o*-DCB systems, the melting enthalpies  $\Delta H$  related to the non-variant events are about the same irrespective of the transition, but are relatively low with respect to those of the terminal melting  $\Delta H_f$ , which are about five times larger (see Fig. S1, ESI†). Extrapolation of  $\Delta H_f$  to  $X_{\text{TATA}} = 1$  yields  $\Delta H_f = 41 \text{ J g}^{-1}$ , a value about 25% lower than that measured in the solid state.<sup>15</sup> This would be consistent with a molecular compound rather than a solid phase. Note that these enthalpies are not heating-rate dependent in both systems, which emphasizes the absence of any noticeable kinetic effects. We further observe that the terminal melting enthalpies are larger in *m*-DCB than in *o*-DCB (see Fig. S1, ESI†).

The shapes of the phase diagrams suggest that the non-variant events correspond to the transformations of molecular compounds.<sup>19,29</sup> This assumption will receive further confirmation in what follows. In *m*-DCB, compound  $C_1$  transforms into compound  $C_2$  at  $T = 51.5^\circ \text{C}$ , which is likely to be related to a change of the solvation degree, or in other words to a change of the compound stoichiometry. The same comments apply to TATA/*o*-DCB gels with the occurrence of 4 molecular compounds. The following sequence takes place  $C_1 \Rightarrow C_2$  at  $T = 16 \pm 1^\circ \text{C}$ ,  $C_2 \Rightarrow C_3$  at  $T = 37 \pm 1^\circ \text{C}$  and  $C_3 \Rightarrow C_4$  at  $T = 61 \pm 1^\circ \text{C}$ . The stoichiometries of the different compounds cannot be inferred from the present data. In view of the size of the solvent molecules with respect to the space between the aliphatic arms, the number of DCB molecules per TATA moiety is likely to be high. There might be two types of solvent molecules, those tightly bound close to the TATA core, and those loosely bound located at the edge.

In both cases, the last non-variant event at  $T = 106^\circ \text{C}$  for TATA/*m*-DCB gels and  $T = 84^\circ \text{C}$  for TATA/*o*-DCB gels may be assigned to a *metatectic transition*<sup>18</sup> corresponding to the phases  $P_\alpha$  and  $P_\beta$  (see Fig. S2, ESI†). The metatectic concentration where three phases are supposed to coexist, namely  $P_\alpha$  and  $P_\beta$ , and  $C_4$ , is about  $C_{\text{TATA}} \approx 0.025 \text{ g g}^{-1}$  in *o*-DCB and  $C_{\text{TATA}} \approx 0.036 \text{ g g}^{-1}$  in *m*-DCB. Assuming metatectic transitions at  $85^\circ \text{C}$  for TATA/*o*-DCB and at  $105^\circ \text{C}$  for TATA/*m*-DCB relies on the existence of several crystalline forms of TATA in the solid state (see Fig. S3, ESI†) so that  $P_\alpha$  and  $P_\beta$  could correspond to the recently reported structure II and III,<sup>15</sup> respectively. Yet, at this stage, we cannot completely disregard their possible molecular compound status.



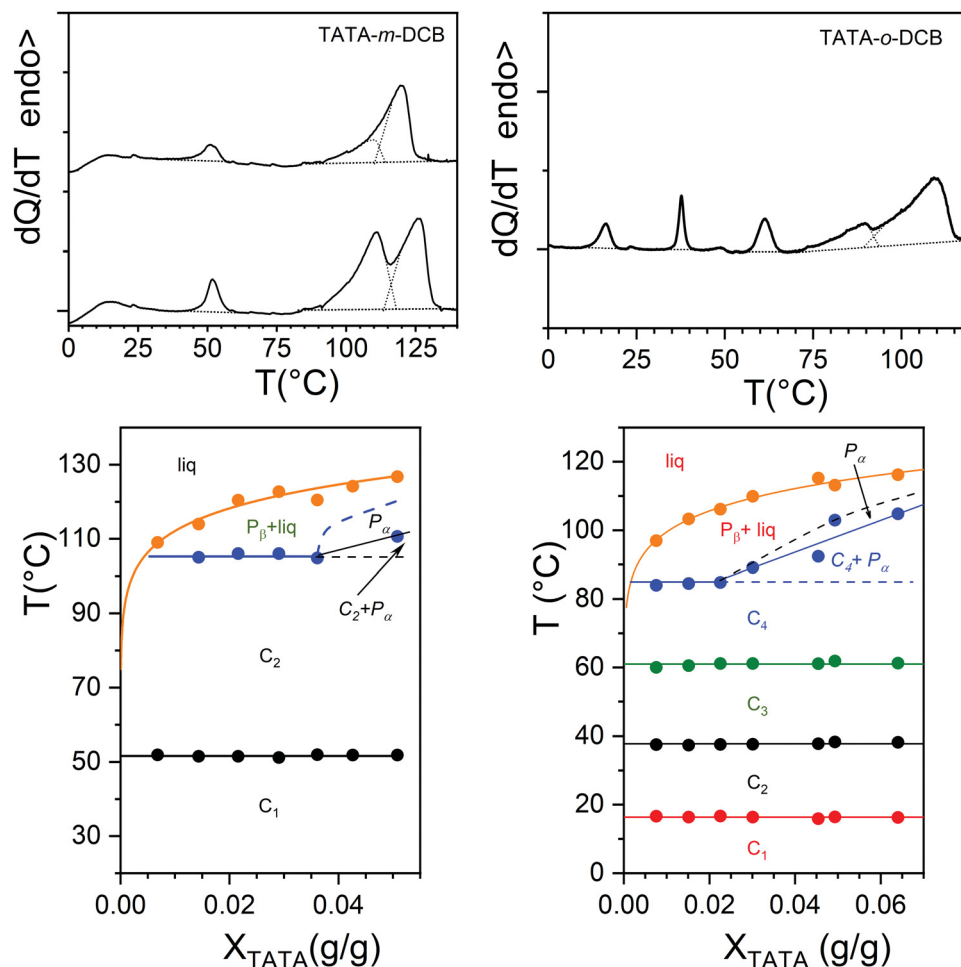


Fig. 2 (a) Top left, typical DSC thermograms for TATA/*m*-dichlorobenzene gels; (b) top right, typical DSC thermograms for TATA/*o*-dichlorobenzene gels; (c) bottom left, temperature–concentration phase diagram for TATA/*m*-dichlorobenzene gels, the orange line is a guide for the eyes; (d) bottom right, temperature–concentration phase diagram for TATA/*o*-dichlorobenzene gels, the orange line is a guide for the eyes.

## (2) Molecular structure of tri-aryl-triamine binary gels

The molecular structure of TATA/*o*-DCB gels has been chiefly investigated by neutron diffraction as this technique is well-suited to the case of molecular compounds.<sup>30–34</sup> Complementary X-ray diffraction investigations have also been performed.

The intensity diffracted by any kind of radiation for a binary system is written:

$$I(q) = \bar{A}_i^2(q)S_i(q) + \bar{A}_j^2(q)S_j(q) + 2\bar{A}_i(q)\bar{A}_j(q)S_{ij}(q) \quad (1)$$

where  $A_i$ ,  $A_j$  and  $S_i(q)$ ,  $S_j(q)$  are the scattering amplitudes and the structure factors of species  $i$  and  $j$ , respectively, while  $S_{ij}(q)$  is the cross-term related to the intermolecular terms. For TATA/*o*-DCB gels, one may contemplate two possible cases: either the solvent is totally excluded from the fibrils, a case designated as a *solid phase*, or the TATA molecules and the solvent co-crystallize, thus forming a *molecular compound*.

If the system forms a solid phase, then  $I(q)$  reduces to:

$$I(q) = \bar{A}_{\text{TATA}}^2(q)S_{\text{TATA}}(q) + \bar{A}_{\text{oDCB}}^2(q)S_{\text{oDCB}}(q). \quad (2)$$

with obvious meaning as to the subscripts.

This implies that altering the value of  $\bar{A}_{\text{oDCB}}$  is not to modify the diffraction pattern of the crystalline TATA fibrils. In other words, the relative intensity of the peaks with respect to one another will not change as  $S_{\text{oDCB}}(q)$  is chiefly featureless except for a broad maximum.

Conversely, if a molecular compound is formed, then the cross-term in relation (1) cannot be discarded any longer. As a result, if one can modify the solvent contrast,  $\bar{A}_{\text{solv}}$  the diffraction pattern will be altered.<sup>31–33</sup> Testing the occurrence of molecular compounds can be achieved by means of neutron diffraction by comparing the diffraction pattern obtained in hydrogenous and in deuterated solvents as the scattering length of hydrogen ( $a_{\text{H}} = -0.375 \times 10^{-12}$  cm) differs drastically from that of deuterium ( $a_{\text{D}} = 0.670 \times 10^{-12}$  cm).<sup>34</sup> The neutron diffraction peak intensity can therefore vary significantly with the solvent labelling. In this paper, only systems in *o*-DCB have been considered since deuterated *m*-DCB was not commercially available (see Table ST1, ESI†).

The neutron diffraction patterns for TATA/*o*-DCB<sub>H</sub> and TATA/*o*-DCB<sub>D</sub> are given in Fig. 3 together with the X-ray diffraction pattern obtained for TATA/*o*-DCB<sub>H</sub>. For the neutron



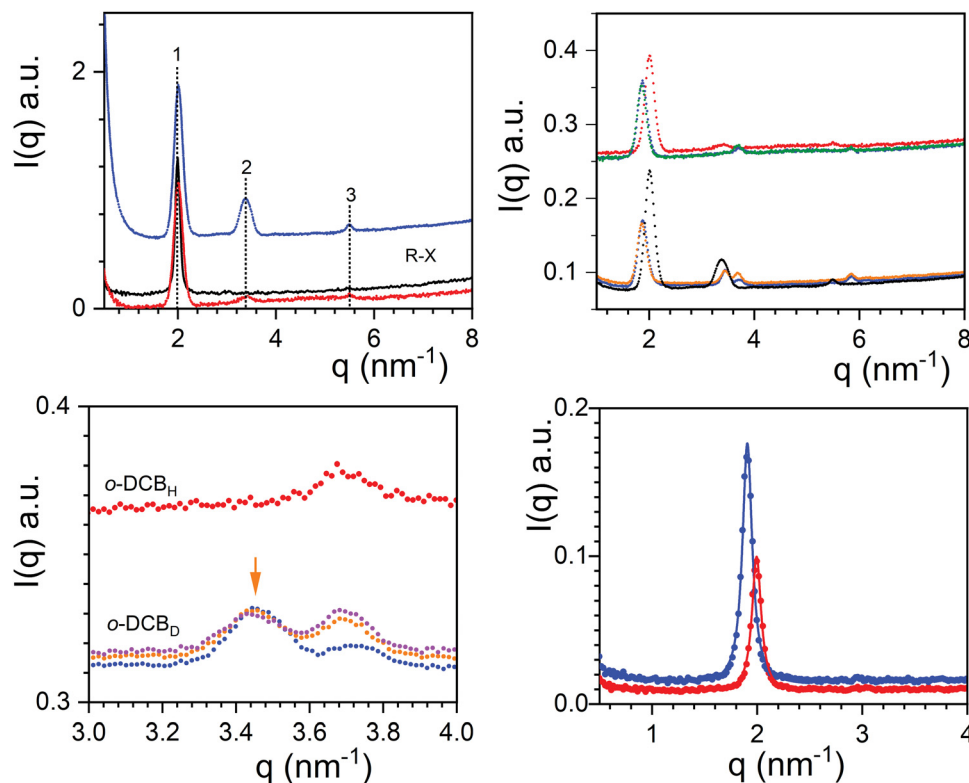


Fig. 3 (a) Upper left, neutron diffraction patterns for TATA/o-DCB<sub>H</sub> (red,  $C_{\text{TATA}} = 4.2 \times 10^{-2} \text{ g cm}^{-3}$ ) and TATA/o-DCB<sub>D</sub> (blue,  $C_{\text{TATA}} = 4.1 \times 10^{-2} \text{ g cm}^{-3}$ ), and the X-ray diffraction pattern (black,  $C_{\text{TATA}} = 4.0 \times 10^{-2} \text{ g cm}^{-3}$ ); (b) upper right, neutron diffraction patterns for: upper curves = TATA/o-DCB<sub>H</sub> ( $C_{\text{TATA}} = 4.2 \times 10^{-2} \text{ g cm}^{-3}$ ) with red = 20 °C, green = 50 °C, lower curves TATA/o-DCB<sub>D</sub> ( $C_{\text{TATA}} = 4.64 \times 10^{-2} \text{ g cm}^{-3}$ ), with black = 20 °C, blue = 50 °C, orange = 70 °C. (c) Lower left, diffraction curve of TATA/o-DCB<sub>D</sub> for different temperatures for the peaks at  $q = 3.439 \text{ nm}^{-1}$  and  $q = 3.675 \text{ nm}^{-1}$ , blue  $T = 50 \text{ °C}$ , orange  $T = 70 \text{ °C}$ , pink  $T = 85 \text{ °C}$ , red TATA/o-DCB<sub>H</sub>  $T = 85 \text{ °C}$ . (d) Lower right, X-ray diffraction patterns in the low  $q$  range of TATA/o-DCB (red) and TATA/*m*-DCB (blue), the solid lines are fits with a Lorentzian function.

diffraction patterns, three peaks are observed at  $q_1 = 2.00 \text{ nm}^{-1}$ ,  $q_2 = 3.38 \text{ nm}^{-1}$ , and  $q_3 = 5.49 \text{ nm}^{-1}$  with intensities that differ as to whether hydrogenous or deuterated *o*-dichlorobenzene is used. The intensity ratio of peaks 1 and 2 is  $I_2/I_1 = 0.315$  for TATA/o-DCB<sub>D</sub> against  $I_2/I_1 = 0.053$  for TATA/o-DCB<sub>H</sub>, which confirms the occurrence of the molecular compound  $C_2$  in agreement with the conclusions drawn from the phase diagram. The X-ray diffraction pattern is nearly identical to the TATA/o-DCB<sub>H</sub> system as it shows only the peak located at  $q_1 = 1.99 \text{ nm}^{-1}$ .

A tentative model of the clathrate type<sup>35,36</sup> is shown in Fig. S6 (ESI<sup>†</sup>) as a basis for discussion. It relies on previous calculations that suggested that TATA helices arrange in rows.<sup>15</sup> At the present resolution level, the helical structure scatters as a hollow solid cylinder. The intensity diffracted by an array of such cylinders is written:<sup>37–39</sup>

$$I(q) \sim \frac{4\pi\mu_L}{q^3 r_{\text{out}}^2} \left[ \frac{A_{\text{in}}}{A_{\text{m}}} J_1(q\gamma r_{\text{out}}) + \frac{A_{\text{out}}}{A_{\text{m}}} \times \{J_1(qr_{\text{out}}) - \gamma J_1(q\gamma r_{\text{out}})\} \right]^2 \times \sum_{j=1}^n \sum_{k=1}^n J_o(qd_{jk}) \quad (3)$$

where  $A_{\text{in}}$ ,  $A_{\text{out}}$ ,  $r_{\text{in}}$ , and  $r_{\text{out}}$  are the scattering amplitudes and

the radii of the inner cylinder and of the outer hollow cylinder, respectively, with  $\gamma = r_{\text{in}}/r_{\text{out}}$ , and  $A_{\text{m}} = \gamma^2 A_{\text{in}} + [1 - \gamma^2]A_{\text{out}}$ ,  $d_{jk}$  is the distance between the axis of cylinders  $j$  and  $k$ ,  $n$  is the number of cylinders, and  $J_1$  and  $J_o$  are Bessel functions of the first type and order 1 and 0, respectively. The first term in eqn (3) is the helix form factor, while the second term stands for the intermolecular interactions.

By considering helices arranged in rows instead of hexagonally assembled, one can reproduce fairly well the peak at  $q_1 = 2.00 \text{ nm}^{-1}$  with a close FWHM value by taking  $r_{\text{out}} = 1.7 \text{ nm}$ ,  $\gamma = 0.41$ ,  $n = 20$ , and the distance between adjacent helices  $d = 3.4 \text{ nm}$  (see Fig. S4, ESI<sup>†</sup>). The value of  $r_{\text{out}}$  which is lower than the value expected for the all-extended structures (2.1 nm), suggests therefore some disorganization of the aliphatic arms.

The additional peaks observed with the deuterated solvent imply an organization of the solvent differing from that of the TATA helices. It is, however, not possible to elaborate further with the present data.

Neutron diffraction patterns for TATA/o-DCB<sub>H</sub> and TATA/o-DCB<sub>D</sub> as a function of temperature are displayed in Fig. 3. As will be further discussed below, the results at  $T = 50 \text{ °C}$ ,  $70 \text{ °C}$  and  $85 \text{ °C}$  are all consistent with the existence of molecular compounds.

A conspicuous change of the diffraction pattern is seen after transforming  $C_2$  into  $C_3$ . Peak  $q_1$  shifts to a lower  $q$  value



(from  $q = 2.00 \text{ nm}^{-1}$  to  $1.872 \text{ nm}^{-1}$ ) while peaks  $q_2$  and peak  $q_3$  shift to higher  $q$  values (from  $3.380 \text{ nm}^{-1}$  to  $3.439 \text{ nm}^{-1}$ , and from  $5.846 \text{ nm}^{-1}$  to  $5.497 \text{ nm}^{-1}$ ). Also, a new peak occurs at  $q = 3.704 \text{ nm}^{-1}$ . The most prominent peak, here  $q_1$ , is generally related to the distance between first neighbours, namely between TATA helices in the present case. Its shift to lower  $q$  values therefore suggests an increase of this distance. This may suggest that the stoichiometry, namely the number of solvent molecules per TATA moiety, increases at the  $C_2 \Rightarrow C_3$  transition instead of decreasing as is usually observed. In other words, the structure would absorb instead of releasing solvent molecules. The slight decrease of turbidity detected at this transition is consistent with this assumption (see Fig. S5, ESI<sup>†</sup>) as raising the compound stoichiometry is likely to bring the refractive index of the TATA fibrils closer to that of the surrounding solvent.

The transformations  $C_3 \Rightarrow C_4$ , and  $C_4 \Rightarrow P_x$  do not entail a modification of the positions of the peaks of the diffraction pattern. The changes only occur at the intensity level, particularly the ratio of the intensities of the peak at  $q = 3.439 \text{ nm}^{-1}$  vs. the peak at  $q = 3.704 \text{ nm}^{-1}$ ,  $I(3.704)/I(3.439)$  for the system TATA/*o*-DCB<sub>D</sub> (see Fig. 3c upper right, and Fig. S6, ESI<sup>†</sup>). This effect is possibly related to a change of stoichiometry that occurs without alteration of the molecular arrangement.

Comparison of the diffraction patterns obtained at  $85^\circ\text{C}$  for *o*-DCB<sub>H</sub> and *o*-DCB<sub>D</sub> is consistent with the existence of a compound. Yet, it seems difficult to conclude that phase  $P_\beta$  is a molecular compound as the presence of compound C4 cannot be excluded. Due to technical constraints we could not increase the temperature above  $85^\circ\text{C}$ , which would have been the only way to settle this issue.

Finally, X-ray diffraction patterns show a slight difference in peak positions between gels in *o*-DCB and *m*-DCB (see Fig. 3). The intense peak  $q_1$  stands at a slightly lower  $q$  for *m*-DCB ( $q = 1.905 \text{ nm}^{-1}$ ) than for *o*-DCB ( $q = 1.99 \text{ nm}^{-1}$ ). This therefore suggests a slightly different packing in each isomer, which is in line with the discrepancies observed in the temperature-concentration phase diagram.

Note that in all the diffraction patterns shown here, the full-width at half maximum (FWHM) of the first peak is about 0.1–0.12 which suggests a rather low degree of disorganization as the correlation length is about  $2\pi/\text{FWHM} \approx 60\text{--}50 \text{ nm}$ .

### (3) Morphology of tri-aryl-triamine binary gels

Morphologies obtained by means of scanning electron microscopy are shown in Fig. 4. Again, as already reported for the thermodynamic properties, the solvent isomer has a marked impact on the microscopic framework of TATA/DCB gels. TATA/*m*-DCB gels consist of a jumble of flat, chipped laths of width somewhere between  $0.5\text{--}3 \mu\text{m}$  (Fig. 4a). Most of the laths are not connected nor branched to one another. TATA/*o*-DCB gels are made up with needles of cross-section somewhere between  $0.06\text{--}0.4 \mu\text{m}$ , that, too, display no interconnections (Fig. 4b). Therefore, gels in DCB possess a similar network architecture, yet with sizes differing by one order of magnitude depending upon the isomer used.

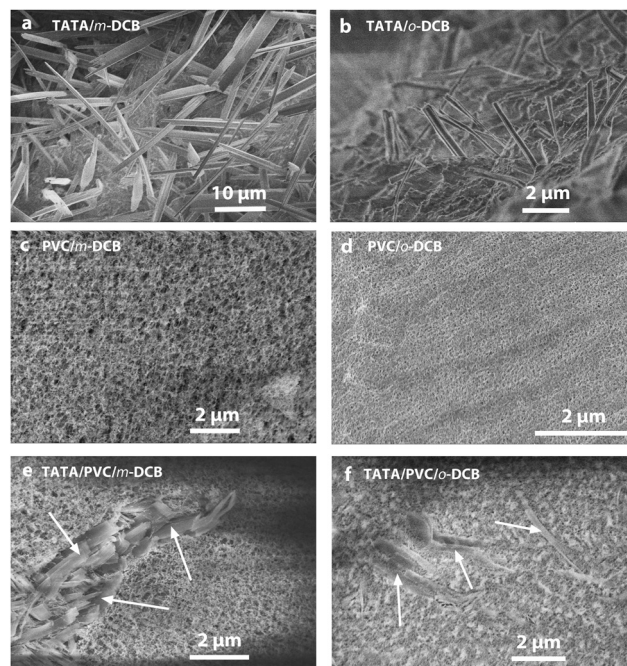


Fig. 4 Scanning electron micrographs: (a) TATA/*m*-DCB  $C_{\text{TATA}} = 0.01 \text{ g cm}^{-3}$ ; (b) TATA/*o*-DCB  $C_{\text{TATA}} = 0.01 \text{ g cm}^{-3}$ ; (c) PVC/*m*-DCB  $C_{\text{PVC}} = 0.05 \text{ g cm}^{-3}$ ; (d) PVC/*o*-DCB  $C_{\text{PVC}} = 0.05 \text{ g cm}^{-3}$ ; (e) TATA/PVC/*m*-DCB  $C_{\text{TATA}} = 0.01 \text{ g cm}^{-3}$ ,  $C_{\text{PVC}} = 0.05 \text{ g cm}^{-3}$ , arrows indicate the TATA fibrils; (f) TATA/PVC/*o*-DCB  $C_{\text{TATA}} = 0.01 \text{ g cm}^{-3}$ ,  $C_{\text{PVC}} = 0.05 \text{ g cm}^{-3}$ , arrows indicate the TATA fibrils.

In principle, TATA helices are supposed to be aligned along the length of the laths/needles.<sup>23</sup> The smaller width observed in *o*-DCB with respect to *m*-DCB may be related to their disparate molecular compounds, whose aggregation of their solvated helices could be rapidly arrested by accumulation of packing defects. Another possibility could be that helix growth is much faster than the helix-helix aggregation, in which case the material is rapidly consumed thus limiting the lateral dimensions.

### (4) The ternary gels PVC/TATA/DCB

As was already described in previous papers, the gelation temperature  $T_{\text{gel}}$  of PVC gels cannot be straightforwardly determined by DSC. The tilting tube method has been used in this study for accessing this parameter.<sup>15</sup> As can be seen in Fig. 5 upper left, the gelation temperature depends significantly upon the DCB isomer, about  $15\text{--}20^\circ\text{C}$  higher in *m*-DCB than in *o*-DCB. The PVC gel morphology is also affected by the DCB isomer as can be seen in Fig. 5c and d, but appears basically as a network of connected fibrils. In *m*-DCB, the mesh size is in the submicron range (about  $0.1\text{--}0.2 \mu\text{m}$ ) while the fibril cross-section is a few tens of nm (around  $40 \text{ nm}$ ). In *o*-DCB, the mesh size is smaller (about  $0.1 \mu\text{m}$ ) while the fibril cross-section looks about the same at the current resolution.

The TATA gel formation temperature is substantially higher in both isomers than that of the PVC gels for all the investigated range of PVC compositions (Fig. 5 upper right). This implies that the TATA network is likely to form before the PVC's during



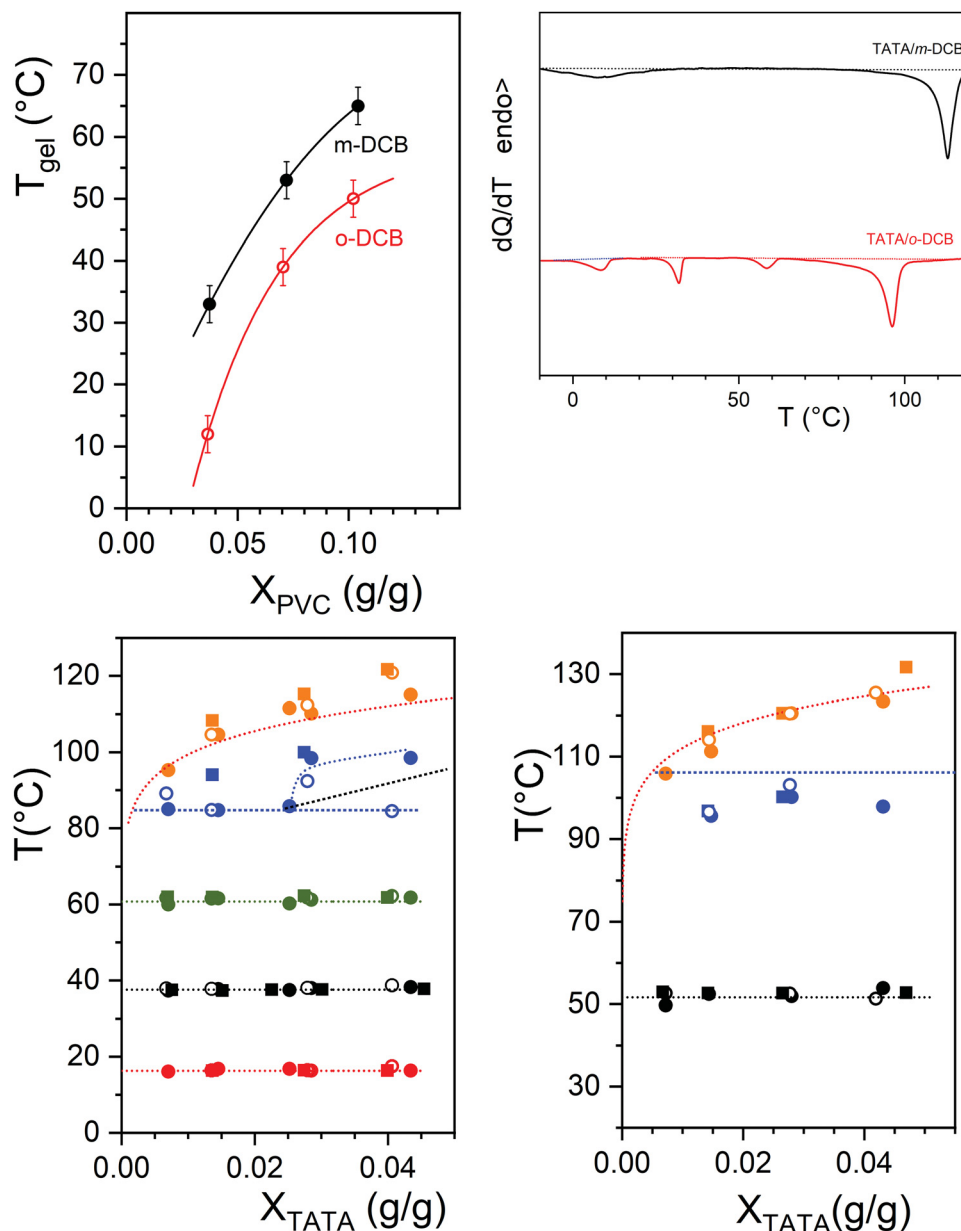


Fig. 5 (a) Upper left, formation temperatures of PVC gels (5%, 10% and 15%) in *m*-DCB (black) and in *o*-DCB (red); (b) upper right, typical DSC formation exotherms for TATA in *m*-DCB (black) and in *o*-DCB (red); (c) lower left, temperature concentration phase diagrams for TATA/PVC/*o*-DCB systems, the dotted lines stand for the TATA/*o*-DCB binary system, (●, ●, ●, ●, ●) = TATA in PVC5%, (○, ○, ○, ○, ○) = TATA in PVC10%, (■, ■, ■, ■, ■) = TATA in PVC15%; (d) lower right, temperature concentration phase diagrams for TATA/PVC/*m*-DCB systems. The dotted lines stand for the TATA/*m*-DCB binary system, (●, ●, ●) = TATA in PVC5%, (○, ○, ○) = TATA in PVC10%, (■, ■, ■) = TATA in PVC15%.

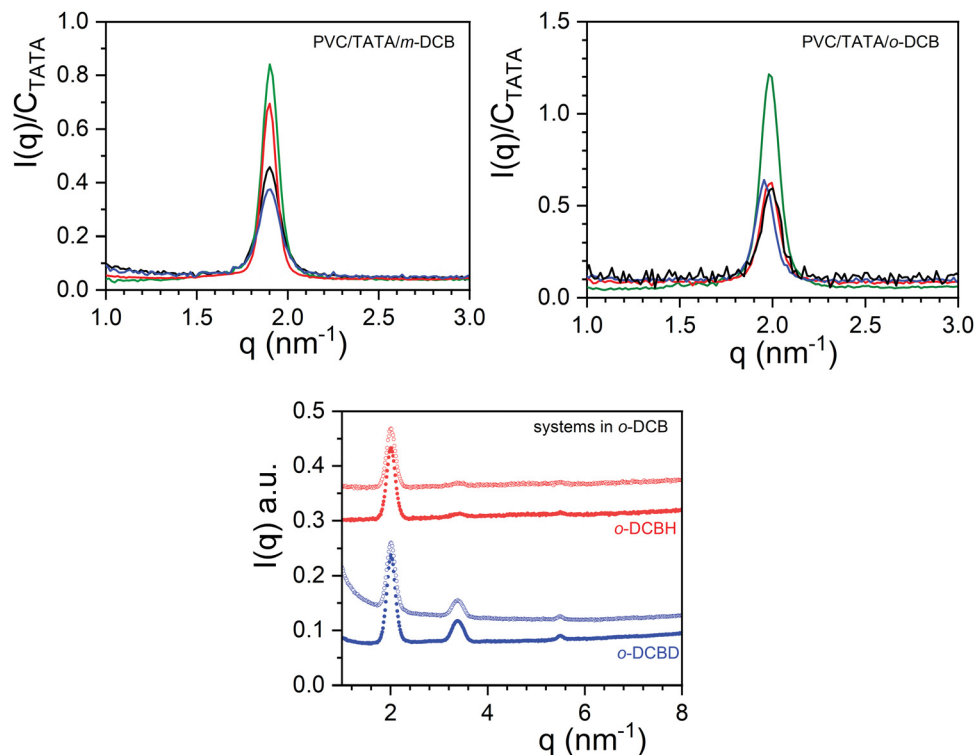
the cooling process of the ternary systems. The morphology of the hybrid systems shown in micrographs 5e and 5f confirm this statement as the TATA fibrils can be distinctly discriminated from the PVC network. Clearly, the PVC gel grows within the TATA network.

The temperature–concentration phase diagrams determined for the different PVC compositions are displayed in Fig. 5 where the dotted lines stand for the transition determined for TATA/DCB binary gels. As a rule, the temperatures of the first-order thermal events associated with the TATA moiety are virtually unaltered by the PVC chains. Yet, the metatectic line is not so

well-defined for the TATA/PVC/*m*-DCB system, and appears to be shifted downwards by about 10 °C.

While the temperature–concentration phase diagrams of the ternary systems are virtually unchanged with respect to the TATA binary gels, the associated enthalpies turn out to be lower and their values are highly scattered (see Fig. S7, ESI†). This implies that a fraction of the TATA molecules does not gel. Since the TATA network forms first, it is surmised that these TATA molecules establish a privileged interaction with PVC chains that prevents them from aggregating.





**Fig. 6** (a) Upper left, X-ray diffraction patterns in the low  $q$  range for TATA/PVC/ $m$ -DCB, (green) = TATA/ $m$ -DCB, (red) = PVC5%, (black) = PVC10%, (blue) = PVC15%.  $C_{TATA} = 0.04$  g cm<sup>-3</sup>. Curves are rescaled by the TATA concentration. (b) Upper right, X-ray diffraction patterns in the low  $q$  range for TATA/PVC/ $o$ -DCB, (green) = TATA/ $o$ -DCB (red) = PVC5%, (black) = PVC10%, (blue) = PVC15%.  $C_{TATA} = 0.041$  g cm<sup>-3</sup>. Curves are rescaled by the TATA concentration. (c) Lower, neutron diffraction pattern in  $o$ -DCBH (upper red curves) and  $o$ -DCBD<sub>D</sub> (lower blue curves) for TATA/ $o$ -DCB (●), and for TATA/PVC10%/o-DCB (○).

The existence of isolated TATA molecules is further confirmed by the X-ray diffraction patterns obtained on the hybrid gels at room temperature (see Fig. 6). While there is no change as to the position of the peaks for the different PVC compositions, the intensities differ significantly. For PVC/TATA/ $o$ -DCB systems the intensity is about half the value of that in the binary gels independent of the PVC composition. Conversely, the intensity decreases with increasing PVC composition for PVC/TATA/ $m$ -DCB systems.

As expected in view of the temperature–concentration phase diagrams, molecular compound  $C_2$  is still present in the ternary PVC/TATA/ $o$ -DCB hybrid gels as ascertained by neutron diffraction data in Fig. 6. Peaks remain at the same positions, and the intensity ratios are the same. Only an upturn is seen for the PVC/TATA/ $o$ -DCBD<sub>D</sub> sample due to the PVC coherent scattering.

## Concluding remarks

Results presented herein show the significant effect of the solvent isomer on the thermodynamic properties, the morphology, and to a lesser extent the molecular structure of TATA organogels. Molecular compounds are, however, formed for both solvents with different stoichiometries. These results highlight the complexity of the phenomenon of organogelation.

In the present case, the Hansen parameters are the same for  $o$ -DCB and  $m$ -DCB<sup>13</sup> ( $h = 20.5$ ), and yet the thermodynamic properties differ considerably. Clearly, the Hansen solubility parameter chiefly indicates whether structure formation is possible or not. Yet, it cannot predict the morphology, particularly whether structures that comply to the definition of a network are generated. For instance, when spherulites are formed<sup>40</sup> the gel status can be legitimately questioned. Note that relying on routine rheological experiments where  $G' > G''$  can be deceiving.<sup>41,42</sup> Indeed, the rheological definition of a true gel states that an elastic modulus at 0 frequency (or infinite time) should be observed. Recent observation by Collin *et al.*<sup>42</sup> for organogels, but also former investigations on polymer thermoreversible gels<sup>41</sup> report that this condition is not always fulfilled.

This investigation therefore opens up new questions such as resolving the structure of the solvent in the TATA molecular compounds. One may wonder whether there is simply a solvent shell around the 20<sub>1</sub> TATA helix reported earlier,<sup>23</sup> or a new helical form that can accommodate the solvent molecules through intercalation, has to be considered.

It is here further observed that the TATA molecules keep their properties when imbedded in a PVC matrix. This again confirms previous results<sup>14,15</sup> that show the possibility of preparing functional materials with an organogelator and a polymer by making hybrid gels.





Finally, the recent reports on other systems suggest that the occurrence of molecular compounds with other self-assembling systems may be a more general trend.<sup>43,44</sup>

## Conflicts of interest

The authors declare no conflict of interest.

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## References

- 1 P. Terech and R. G. Weiss, Low molecular mass gelators of organic liquids and the properties of their gels, *Chem. Rev.*, 1997, **97**, 3133–3159.
- 2 In *Molecular Gels: Materials with Self-Assembled Fibrillar Networks*, ed. P. Terech and R. G. Weiss, Springer Verlag, 2006.
- 3 R. G. Weiss, The past, present and future of molecular gels. What is the status of the field, and where is it going?, *J. Am. Chem. Soc.*, 2014, **136**, 7519–7530.
- 4 J. M. Guenet, *Organogels: thermodynamics, structure, solvent role and properties*, Springer International Publishing, N.Y., 2016. According to the dictionary: a network is “a large system of lines, tubes, wires, etc. that cross one another or are connected with one another”.
- 5 In *Molecular Gels, Structure and Dynamics*, ed., R. G. Weiss, Monograph in Supramolecular Chemistry, Royal Society of Chemistry, London, 2018.
- 6 X. L. Liu and J. L. Li, *Soft Fibrillar Materials: Fabrication and Applications*, Wiley-VCH, 2013.
- 7 S. S. Babu, V. K. Praveen and A. Ajayaghosh, Functional  $\pi$ -Gelators and Their Applications, *Chem. Rev.*, 2014, **114**, 1973–2129.
- 8 C. M. Hansen, *Hansen Solubility Parameter, Users' Handbook*, CRC Press, London, 2nd edn, 2007; The three Hansen parameters are:  $\delta_d = 18$ ,  $\delta_p = 4.4$ ,  $\delta_H = 4.2$  for tetrachloroethane,  $\delta_d = 19.2$ ,  $\delta_p = 5.5$ ,  $\delta_H = 4.1$  for bromobenzene. For the solvents used here,  $\delta_d = 19.2$ ,  $\delta_p = 6.3$ ,  $\delta_H = 3.3$  for *o*-DCB, and  $\delta_d = 19.2$ ,  $\delta_p = 6.3$ ,  $\delta_H = 3.3$  for *m*-DCB, the resulting *h* is 20.5.
- 9 R. F. Fedors, A method for estimating both the solubility parameters and the molar volume of liquids, *Polym. Eng. Sci.*, 1974, **14**, 147.
- 10 M. Raynal and L. Bouteiller, Organogel formation rationalized by Hansen solubility parameters, *Chem. Commun.*, 2011, **47**, 8271.
- 11 N. Yan, Z. Xu, K. K. Diehn, S. R. Raghavan, Y. Fang and R. G. Weiss, How do liquid mixtures solubilize insoluble gelators? Self-assembly properties of pyrenyl-glucono gelators in tetrahydrofuran-water mixtures, *J. Am. Chem. Soc.*, 2013, **135**, 8989.
- 12 J. Bonnet, G. Suissa, M. Raynal and L. Bouteiller, Organogel formation rationalized by Hansen solubility parameters: dos and don'ts, *Soft Matter*, 2014, **10**, 3154.
- 13 C. M. Hansen, *Hansen Solubility Parameters: A User's Handbook*, CRC Press, Boca Raton, 2nd edn, 2007.
- 14 B. Kiflemariam, D. Collin, O. Gavata, A. Carvalho, E. Moulin, N. Giuseppone and J. M. Guenet, Hybrid materials from tri-aryl amine organogelators and poly[vinyl chloride] networks, *Polymer*, 2020, **207**, 122814.
- 15 P. Talebpour, B. Heinrich, O. Gavata, A. Carvalho, E. Moulin, N. Giuseppone and J. M. Guenet, Modulation of the Molecular Structure of Tri-aryl Amine Fibrils in Hybrid Poly[vinyl chloride] Gel/Organogel Systems, *Macromolecules*, 2021, **54**, 8104.
- 16 J.-M. Guenet, B. Demé, O. Gavata, E. Moulin and N. Giuseppone, Evidence by neutron diffraction of molecular compounds in triarylamine tris-amide organogels and in their hybrid thermoreversible gels with PVC, *Soft Matter*, 2022, **18**, 2851.
- 17 J. M. Guenet, *Polymer\_solvent molecular compounds*, Elsevier, London, 2008.
- 18 J. J. Point and C. Coutelier, Linear high polymers as host in intercalates. Introduction and example, *J. Polym. Sci., Part B: Polym. Phys.*, 1985, **23**, 231.
- 19 J. J. Point, C. Coutelier and D. Villers, Intercalates with linear polymer host. Part 4. Structure of p-C<sub>6</sub>H<sub>4</sub>XY-poly(oxyethylene) intercalates from unit-cell determination and energy minimization, *J. Phys. Chem.*, 1986, **90**, 3277.
- 20 L. Feng and K. A. Cavicchi, Investigation of the relationships between the thermodynamic phase behavior and gelation behavior of a series of tripodal trisamide compounds, *Soft Matter*, 2012, **8**, 6483.
- 21 J. M. Guenet, Physical Aspects of Organogelation: A Point of View, *Gels*, 2021, **7**, 65.
- 22 E. Moulin, F. Niess, M. Maaloum, E. Buhler, L. Nyrkova and N. Giuseppone, The hierarchical self-assembly of charge nanocarriers: a highly cooperative process promoted by visible light, *Angew. Chem., Int. Ed.*, 2010, **49**, 6974.
- 23 J. J. Armao IV, M. Maaloum, T. Ellis, G. Fuks, M. Rawiso, E. Moulin and N. Giuseppone, Healable supramolecular polymers as organic metals, *J. Am. Chem. Soc.*, 2014, **136**, 11382.
- 24 L. Nyrkova, E. Moulin, J. J. Armao IV, M. Maaloum, B. Heinrich, M. Rawiso, F. Niess, J. J. Cid, N. Jouault, E. Buhler, A. Semenov and N. Giuseppone, Supramolecular self-assembly and radical kinetics in conducting self-replicating nanowires, *ACS Nano*, 2014, **8**, 10111.
- 25 E. Moulin, J. J. Armao IV and N. Giuseppone, Triaryl amine-based supramolecular polymers: structure, dynamics, and functions, *Acc. Chem. Res.*, 2019, **52**, 975.
- 26 Y. C. Yang and P. H. Geil, Morphology and properties of PVC/solvent gels, *J. Macromol. Sci., Part B: Phys.*, 1983, **22**, 463.



- 27 P. H. Mutin and J. M. Guenet, Physical gels from PVC: ageing and solvent effect on thermal behavior, swelling and compression modulus, *Macromolecules*, 1989, **22**, 843.
- 28 V. Cristiglio, B. Giroud, L. Didier and B. Demé, D16 is back to business: more neutrons, more space, more fun, *J. Neutron Res.*, 2015, **26**, 23.
- 29 See for instance: A. Reisman, *Phase Equilibria: Basic Principles, Applications, experimental Techniques*, Acad. Press, London, 1970.
- 30 R. E. Rundle, C. G. Schull and E. O. Wollan, The crystal structure of thorium and zirconium dihydrides by X-ray and neutron diffraction, *Acta Crystallogr.*, 1952, **5**, 22.
- 31 J. J. Point, P. Damman and J. M. Guenet, Neutron diffraction study of poly(ethylene oxide) p dihalogenobenzene crystalline complexes, *Polym. Commun.*, 1991, **32**, 477.
- 32 C. Daniel, A. Menelle, A. Brulet and J. M. Guenet, Thermo-reversible Gelation of Syndiotactic Polystyrene in Toluene and Chloroform, *Polymer*, 1997, **38**, 4193.
- 33 F. Kaneko, N. Seto, S. Sato, A. Radulescu, M. M. Schiavone, J. Allgaier and K. Ute, Development of a Simultaneous SANS/FTIR Measuring System, *Chem. Lett.*, 2015, **44**, 497.
- 34 G. E. Bacon, Coherent neutron-scattering amplitudes, *Acta Crystallogr., Sect. A: Cryst. Phys., Diffr., Theor. Gen. Crystallogr.*, 1972, **A28**, 357.
- 35 H. Tadokoro, Y. Chatani, T. Yoshihara, S. Tahara and S. Murahashi, Structural studies on polyethers,  $[-(\text{CH}_2)_m-\text{O}-]_n$ . II. Molecular structure of polyethylene oxide, *Makromol. Chem.*, 1964, **73**, 109.
- 36 A. Chenite and F. Brisse, Structure and conformation of poly(ethylene oxide), PEO, in the trigonal form of the PEO-urea complex at 173 K, *Macromolecules*, 1991, **24**, 2221.
- 37 O. A. Pringle and P. W. Schmidt, Small-Angle X-Ray Scattering from Helical Macromolecules, *J. Appl. Crystallogr.*, 1971, **4**, 290.
- 38 G. Oster and D. P. Riley, Scattering from Cylindrically Symmetric Systems, *Acta Cryst.*, 1952, **5**, 272.
- 39 P. Mittelbach and G. Porod, Zur Röntgenkleinwinkelstreuung verdünnter kolloiden Systeme, *Acta Phys. Austriaca*, 1961, **14**, 185.
- 40 M. Lescanne, A. Colin, O. Mondain-Monval, F. Fages and J. L. Pozzo, Structural Aspects of the Gelation Process Observed with Low Molecular Mass Organogelators, *Langmuir*, 2003, **19**, 2013.
- 41 J. M. Guenet and G. B. McKenna, The concentration dependence of the compression modulus of iPS/cis decalin gels, *J. Polym. Sci., Polym. Phys. Ed.*, 1986, **24**, 2499.
- 42 D. Collin, R. Covis, F. Allix, B. Jamart-Grégoire and P. Martinoty, Jamming transition in solutions containing organogelator molecules of amino-acid type: rheological and calorimetry experiments, *Soft Matter*, 2013, **9**, 2947.
- 43 K. K. Kartha, S. S. Babu, S. Srinivasan and A. Ajayaghosh, Attogram Sensing of Trinitrotoluene with a Self-Assembled Molecular Gelator, *J. Am. Chem. Soc.*, 2012, **134**, 4834.
- 44 G. Ghosh, A. Chakraborty, P. Pal, B. Jana and S. Ghosh, Direct Participation of Solvent Molecules in the Formation of Supramolecular Polymers, *Chem. – Eur. J.*, 2022, DOI: [10.1002/chem.202201082](https://doi.org/10.1002/chem.202201082).

