

# Self-assembly and mesophase formation in a non-ionic chromonic liquid crystal system

Walker, Martin; Masters, Andrew J.; Wilson, Mark R.

DOI:

[10.1039/c4cp03092c](https://doi.org/10.1039/c4cp03092c)

License:

Creative Commons: Attribution (CC BY)

*Document Version*

Publisher's PDF, also known as Version of record

*Citation for published version (Harvard):*

Walker, M, Masters, AJ & Wilson, MR 2014, 'Self-assembly and mesophase formation in a non-ionic chromonic liquid crystal system: insights from dissipative particle dynamics simulations', *Physical Chemistry Chemical Physics*, vol. 16, no. 42, pp. 23074-23081. <https://doi.org/10.1039/c4cp03092c>

[Link to publication on Research at Birmingham portal](#)

## General rights

Unless a licence is specified above, all rights (including copyright and moral rights) in this document are retained by the authors and/or the copyright holders. The express permission of the copyright holder must be obtained for any use of this material other than for purposes permitted by law.

- Users may freely distribute the URL that is used to identify this publication.
- Users may download and/or print one copy of the publication from the University of Birmingham research portal for the purpose of private study or non-commercial research.
- User may use extracts from the document in line with the concept of 'fair dealing' under the Copyright, Designs and Patents Act 1988 (?)
- Users may not further distribute the material nor use it for the purposes of commercial gain.

Where a licence is displayed above, please note the terms and conditions of the licence govern your use of this document.

When citing, please reference the published version.

## Take down policy

While the University of Birmingham exercises care and attention in making items available there are rare occasions when an item has been uploaded in error or has been deemed to be commercially or otherwise sensitive.

If you believe that this is the case for this document, please contact [UBIRA@lists.bham.ac.uk](mailto:UBIRA@lists.bham.ac.uk) providing details and we will remove access to the work immediately and investigate.



Cite this: *Phys. Chem. Chem. Phys.*,  
2014, 16, 23074

# Self-assembly and mesophase formation in a non-ionic chromonic liquid crystal system: insights from dissipative particle dynamics simulations

Martin Walker,<sup>a</sup> Andrew J. Masters<sup>b</sup> and Mark R. Wilson<sup>\*a</sup>

Results are presented from a dissipative particle dynamics (DPD) simulation of a model non-ionic chromonic system, TP6EO2M, composed of a poly(ethylene glycol) functionalised aromatic (triphenylene) core. The simulations demonstrate self-assembly of chromonic molecules to form single molecule stacks in solution at low concentrations, the formation of a nematic mesophase at higher concentrations and a columnar phase in the more concentrated regime. The simulation model used allows very large system sizes, of many thousands of particles, to be studied. This provides, for the first time, an opportunity to study chromonic phase behaviour by simulation without severe restrictions imposed by system size. In the low concentration limit, the simulations demonstrate approximate isodesmic association from which a binding energy can be obtained, allowing the simulations to be tuned to reproduce the behaviour of the real experimental system.

Received 14th July 2014,  
Accepted 4th September 2014

DOI: 10.1039/c4cp03092c

www.rsc.org/pccp

## 1 Introduction

Chromonic liquid crystals are a sub-class of the lyotropic liquid crystal family.<sup>1,2</sup> The molecular features that drive the formation of chromonic phases are a polyaromatic core, with polar or solubilising groups at the periphery. In water, although in principle other dispersing media could be used, chromonic molecules self-assemble to form stacks with single<sup>3–5</sup> or multiple-molecule cross-sections.<sup>6–8</sup> Upon aggregation, aromatic groups are aligned into a stack (or column), limiting unfavourable interactions with the solvent.<sup>3</sup> At higher concentrations, when aggregates are sufficiently long, mesophase formation occurs from entropy-driven alignment of the long anisotropic columns, resulting in a nematic (N) phase composed of stacks with a size distribution that is dependent upon temperature and concentration. At still higher concentrations a chromonic M phase can form, with a regular two-dimensional packing of columns on a hexagonal lattice.<sup>2</sup> There is considerable current interest in chromonics because of applications in areas such as thin film fabrication,<sup>9</sup> real-time microbial sensors,<sup>10,11</sup> and controllable side-by-side or end-to-end assembly of gold nanorods.<sup>12</sup>

Chromonic mesogens tend to have a strong enthalpic contribution towards the self-assembly process. They aggregate

in a reversible manner, with aggregation occurring in the absence of a critical micelle concentration, and often at extremely low concentrations.<sup>13</sup> The favourable enthalpic interactions are usually attributed to  $\pi$ - $\pi$  stacking of the aromatic cores, leading to the growth of linear aggregates. Since aggregation is a one-dimensional process (for simple single molecule stacks) each molecule only interacts strongly with its two neighbours, above and below in the stack. Molecules at the end of the stack interact strongly with only one neighbouring molecule. This configuration implies an isodesmic binding model; that is the binding environment does not change as the aggregates grow in size.<sup>14</sup> Adding a molecule to the stack increases the stack size, and replaces the end of the stack with another identical to its predecessor. This is distinctly different from micellar growth. As a micelle grows the curvature of the interface of the micelle changes, leading to a different binding environment for each additional molecule.

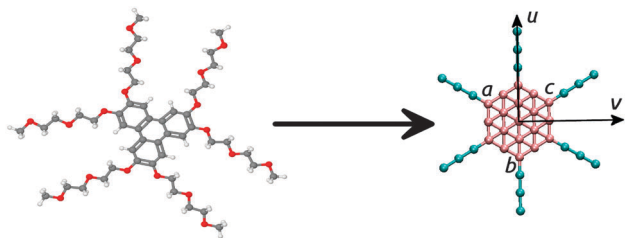
Most identified chromonics fall under the banner of ionic chromonics, with a large chromonic ion that self-assembles, which is responsible for the shape of aggregates in solution, and a dissociated solvated counter ion. Here, counter ions are well-solvated and appear to be only partially associated with the aggregate species.<sup>3</sup> Far less studied are non-ionic chromonics, which tend to have far larger aromatic cores with a number of solubilising groups attached that stabilise the molecule in solution.<sup>15,16</sup>

This study considers the non-ionic chromonic 2,3,6,7,10,11-hexa-(1,4,7-trioxaethyl)-triphenylene, TP6EO2M (Fig. 1). TP6EO2M is a neutral molecule with  $D_{3h}$  symmetry, and is composed of a triphenylene core with six ethylene oxide arms (each ethylene oxide

<sup>a</sup> Department of Chemistry, Durham University Science Laboratories, South Road, Durham, DH1 3LE, UK. E-mail: mark.wilson@durham.ac.uk;  
Tel: +44 (0)191 384 1158

<sup>b</sup> School of Chemical Engineering and Analytical Science, University of Manchester, Manchester, M13 9PL, UK. E-mail: andrew.masters@manchester.ac.uk;  
Tel: +44 (0)161 2754679





**Fig. 1** Molecular structure of a TP6EO2M molecule (left) and the structure of the dissipative particle dynamics representation used in this work (right) for  $L = 3$  chain beads. Beads marked *a*, *b* and *c* are used to apply an aligning field as described in Section 2.1. The vertical and horizontal arrows show the parallel and perpendicular axes respectively.

oligomer unit is terminated with a methyl group). Previous studies<sup>17,18</sup> have shown that TP6EO2M in water has a stable nematic phase, which extends from 14 wt% to 51.2 wt%, depending on temperature, with a hexagonal columnar phase at higher concentrations. An isodesmic binding model has been suggested for the formation of TP6EO2M aggregates.<sup>19</sup>

Simulation studies have provided some useful insights into chromonic behaviour.<sup>14,19,20</sup> In recent work Chami and Wilson<sup>3</sup> have provided one of the first detailed atomistic studies, looking at the structure and dynamics of molecules within a chromonic column in water and the structure of a (pre-assembled) N phase. However, for a molecule as large as TP6EO2M such studies are limited, by computational expense, to a relatively small number of molecules in water; and cannot be used to study bulk phase behaviour or to look at aggregation as a function of concentration.

In the current work, we use the dissipative particle dynamics (DPD)<sup>21</sup> technique to study a model for TP6EO2M at a coarse-grained level. We show how altering the ethylene oxide arm length has subtle effects on local packing, and greatly impacts on the phase behaviour of the system in water. Additionally we show how DPD parameters can be tuned to fit binding energies and reproduce experimental phase behaviour.

## 2 Computational methods

### 2.1 DPD model

DPD has been successfully applied to simulate a number of soft matter systems: block co-polymers,<sup>22–24</sup> colloidal suspensions,<sup>25</sup> lipid bilayers,<sup>26,27</sup> nanoparticles<sup>28,29</sup> and liquid crystalline phases<sup>30,31</sup> including those formed from complex polyphilic molecules.<sup>32–35</sup> Molecular systems are readily coarse-grained to a DPD representation by replacing multi-atom chemical groups with simple, single-site microphase segregating groups (beads). The beads then interact through space with three simple forces acting along the inter-bead vector: a conservative force ( $F_C$ ), a dissipative force ( $F_D$ ) and a random force ( $F_R$ ).

$$F_C = a_{ij}(1 - r_{ij}/r_{cut}) \quad \text{for } |r_{ij}| < r_{cut}$$

or

$$F_C = 0 \quad \text{for } |r_{ij}| \geq r_{cut} \quad (1)$$

where  $a_{ij}$  represents the maximum repulsive force between particles  $i$  and  $j$ , and  $r_{cut}$  is the interaction cutoff, which, as standard, is taken to a single unit of length.

$$F_D = -\gamma\omega^2 r_{ij}(r_{ij} \cdot v_{ij}) \quad \text{for } |r_{ij}| < r_{cut}$$

$$F_D = 0 \quad \text{for } |r_{ij}| \geq r_{cut}; \quad (2)$$

$$F_R = \frac{\sigma\omega r_{ij}\theta}{\sqrt{dt}} \quad \text{for } |r_{ij}| < r_{cut} \quad (3)$$

$$F_R = 0 \quad \text{for } |r_{ij}| \geq r_{cut}$$

where the relationship between  $\sigma$  and  $\gamma$  ( $\sigma^2 = 2\gamma k_B T^*$ ) effectively controls the temperature. How rapidly the system equilibrates with respect to temperature can be controlled with the friction coefficient  $\omega$ , and  $\theta$  is a random normally-distributed variable ( $0 < \theta < 1$ ). Further details of the technique have been reviewed in the literature.<sup>36</sup>

In principle, non-ionic chromonic molecules can be represented by two types of chemical group: an unsaturated aromatic-like group that can be used to represent the core and a hydrophilic group to represent peripheral solubilising groups. In the case of TP6EO2M, in order to properly capture molecular structure (and hence self-assembly behaviour) it is necessary to include additional bonding and angle parameters to accurately represent the molecular shape. The coarse-graining scheme used in this work is shown in Fig. 1. We used a rigid disc-like core, composed of 19 aromatic type beads arranged hexagonally in 2 dimensions, surrounded with six flexible arms (see Fig. 1) each represented by  $L$  beads. We note that while the  $D_{3h}$  symmetry of the atomistic representation has been increased to  $D_{6h}$  within the coarse-grained model, this (38 bead) model is still able to capture the key interactions of TP6EO2M responsible for self-assembly and mesophase formation.

We have also carried out test calculations of a more fine-grained representation of TP6EO2M with  $D_{3h}$  symmetry. However, here a larger number of beads is required to represent the reduced,  $D_{3h}$ , symmetry of the core. This, combined with additional intramolecular interaction terms to maintain geometry, necessitates the use of a smaller DPD time-step. Hence, as the self-assembly behaviour was very similar in the two models in initial tests, we have concentrated on the more coarse-grained representation in the work presented here.

The usual reduced DPD units were employed throughout. Distances were scaled by unit length,  $l = 1$ , we used beads of size  $r_{cut}/l = 1$  with a reduced mass of  $m = 1$ . Energies were scaled by unit energy,  $\epsilon = 1$  and temperature was measured in units of  $T^* = k_B T/\epsilon$ . Our simplified TP6EO2M core maintained rigidity with harmonic springs binding beads at an ideal distance of  $r/l = 0$  using the repulsive DPD potential to provide the repulsive component of the “bonding” potential (and hence a finite bond length). Harmonic angle potentials were used to link any three linear adjacent beads (in Fig. 1) with an ideal angle of  $180^\circ$ . The force constants for these two interactions were set to  $\kappa_{bond}/(\epsilon l^{-2}) = 20$ ,  $\kappa_{angle}/(\epsilon \text{ rad}^2) = 20$  respectively. The ethylene oxide units are represented by simple beads bonded with the



same harmonic potentials described for the core, no angle potentials were imposed on the ethylene oxide units.

The properties of the two types of coarse-grained groups aim to mimic the properties of the aromatic-core and the ethylene oxide chains within TP6EO2M, and so, in the first instance the DPD parameters,  $a_{ij}$  were selected to encourage microphase segregation, allowing the ethylene oxide arm and water to freely mix but, keeping the aromatic core separate. The corresponding parameter set was  $a_{ij}/\epsilon = 25$  when  $i = j$ , or  $i = \text{water}$  and  $j = \text{ethylene oxide}$ , and  $a_{ij}/\epsilon = 45$  for all other interactions.

As the coarse-graining scheme used is relatively crude, we initially varied the ethylene oxide chain length  $L$  from zero to 5 units. Treating  $L$  in this way, for fixed values of  $a_{ij}$  allow us to obtain a chain length that best matches experimental<sup>15</sup> data for TP6EO2M.

1000 molecules of TP6EO2M were placed at random positions and orientations within a cubic simulation box. The required concentration was then obtained by adding an appropriate number of water beads in random positions. To explore the influence of arm length, simulations were performed on molecules with an arm length  $L = 0-5$ , over the range of concentrations 25 wt%, 50 wt%, 75 wt% and 100 wt% (no water). To study binding energies additional longer simulations at 10 wt% were also performed to minimise the influence of highly fluctuating larger aggregates. Given the uniform mass of DPD beads used in these simulations, the weight percent can be mapped in a 1:1 manner with the volume fraction.

Equations of motion were solved using the Velocity Verlet integration algorithm, using the DPD module of the DL\_MESO<sup>37,38</sup> program (DL\_MESO\_DPD) in the constant-NVT (canonical) ensemble. In each case, simulations were started from a random configuration, which was quenched to the desired temperature ( $T^* = 1$ ). More complicated cooling procedures were found to be unnecessary, since the unbound water content allowed for sufficient system flexibility to equilibrate under any set of conditions without the equilibrium configuration depending on thermal history (or nucleation events). We found that 1 000 000 steps with a time step of  $\tau/(l(m/\epsilon)^{1/2}) = 0.02$  was sufficient to ensure a well equilibrated phase in most cases. The parameter slowest to equilibrate was the cluster size/distribution, simulations where this distribution was found unequilibrated were continued for a further 1 000 000 equilibrated steps. The DPD parameter for noise amplitude was taken to be  $\sigma = 3.6710$ . All simulations were performed at a particle site density  $\rho/l^3 = 3.0$ .

Simulations of ordered phases (nematic, columnar and crystal phases) started from a random arrangement of molecules often failed to equilibrate, resulting in linear aggregates with no long range positional, or orientational order. These aggregates were often locked into position by periodic boundary conditions, and could not dynamically reorient due to packing restraints of adjacent aggregates. To aid the equilibration of ordered phases we investigated the influence of a pseudo magnetic field, applied to orient the discs in a single direction. Once a phase formed, the field was removed and a further 100 000 equilibration time steps were performed to study a field free simulation. To impose a field we first calculated the normal vector to the plane defined by three

of the corner aromatic beads,  $a$ ,  $b$  and  $c$  (Fig. 1). The field acted to align this vector to the  $z$ -axis of the simulation cell. The energy due to the field is:

$$|E_{\text{mag}}| = 0.5k_{\text{field}}(\theta - \theta_{\text{eqm}}) \quad (4)$$

where  $\theta = \cos^{-1}((\mathbf{r}_{ab} \times \mathbf{r}_{bc}) \cdot \mathbf{D})$ ,  $\theta_{\text{eqm}} = 0$ , with a field director  $\mathbf{D} = (001)$ . A field strength of  $k_{\text{field}}/(\epsilon/l^{-1} \text{ rad}^{-2}) = 2.0$  was selected as sufficiently large to align the short molecular axis but sufficiently small to minimise the effects of the field on phase boundaries. The presence of the field did not influence phase stability for any of the systems in water. However, without a field it was difficult for long columns formed from self-assembly of an initial random configuration of mesogens to fully align because of the possibility of long aggregates interacting with themselves across periodic boundary conditions. For thermotropics the field helped stabilize mesophases for two systems, as described below.

## 2.2 Analysis of simulations

**Cluster analysis.** Cluster analysis was carried out following the method of Stoddard.<sup>39</sup> Using the central bead of the core as a reference site, any molecule that fell within a radius of 2.5 bead lengths ( $l$ ) was considered to be part of the same aggregate. A value of  $2.5l$  was chosen as it correctly identified the number of molecules within chromonic columns (checked visually); picking up the interactions of molecules that aligned parallel in a shifted or eclipsed configuration, while molecules with an orthogonal arrangement could not achieve this distance of approach without substantial molecular distortion. Cluster calculations sampled an equilibrated simulation 100 times over 10 000 000 time steps.

**Pair distribution functions.** Two dimensional pair distribution functions,  $g(u,v)$ , were calculated using a molecular frame normal to the system director (see Fig. 1). To allow for changes along the system director, each molecule was analysed applying a cutoff distance of  $2.5l$  along the director axis. An additional cutoff of half the simulation box dimension was applied normal to the system director. The parallel axis was defined as the unit vector along the direction from the centre of the core to the core bead where an ethylene oxide arm attaches. The perpendicular axis was implicitly defined as normal to the parallel axis and the molecular short axis (Fig. 1). For comparison we also tested a second version of  $g(u,v)$ , where the parallel vector was defined from the centre of the core to the bead at the end of a selected ethylene oxide arm. No discernible difference was found between the two  $g(u,v)$  definitions.

## 3 Results and discussion

### 3.1 Arm length $L = 0$

An arm length of zero at 100 wt% corresponds to a simulation of a thermotropic polyaromatic discotic liquid crystal. The polyaromatic mesogen shows no clear ordering upon cooling, short linear aggregates form, but fail to equilibrate on any reasonable timescale. To aid in equilibration a weak field ( $k_{\text{field}}/(\epsilon/l^{-1} \text{ rad}^{-2}) = 2.0$ ) was used to align the molecular short axis,



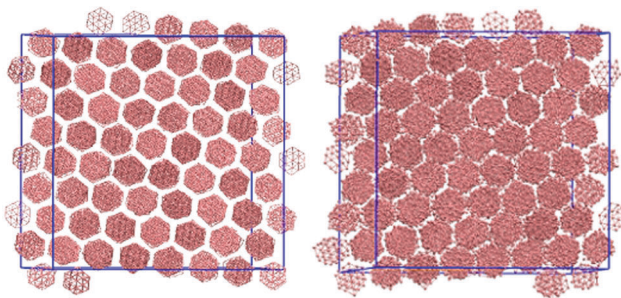


Fig. 2 Crystal columnar (left) and columnar (right) phases of the zero arm ( $L = 0$ ) model. Transitions occurred at  $T^* = 0.7$  (crystal–columnar),  $T^* = 1.5$  (columnar–nematic),  $T^* = 1.9$  (nematic–isotropic) in the presence of a stabilising field.

inducing a series of transitions: isotropic–nematic ( $T^* = 1.9$ ), nematic–columnar ( $T^* = 1.5$ ), columnar–crystalline ( $T^* = 0.7$ ). For the columnar phase (Fig. 2) each molecule has the freedom to rotate giving each column a circular cross section, whereas in the crystalline phase the molecular rotations are frozen such that each column maintains a hexagonal cross-section. The transition between the two highly ordered phases can easily be distinguished through analysis of the  $g(u,v)$  plots (Fig. 3).  $g(u,v)$  for the crystal phase which show long range in-plane orientational order, whereas the columnar mesophase shows hexagonal packing of columns but exhibits in-plane molecular rotations. This leads to a loss of the sharp peaks within the rings that arise from the hexagonal tiling. At higher temperature, the columnar phase melts to give a discotic nematic, with no hexagonal order.

Removal of the field from each equilibrated phase resulted in loss of both the nematic and columnar phases, only the crystal and isotropic phases remained stable with a single transition crystal–isotropic phase transition at  $T^* = 0.8$ .

Since there are no ethylene oxide arms to aid solvation in water, upon addition of water, one would expect any ordered phase to swell, to accommodate minimum amounts of water, then phase separate to form a two phase region. This does

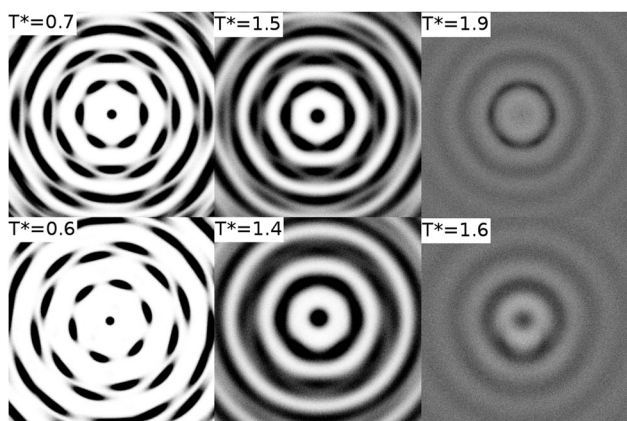


Fig. 3 Two dimensional  $g(u,v)$  for zero arm ( $L = 0$ ) molecule (top) and single arm ( $L = 1$ ) molecule (bottom). The plots show crystal (left), columnar (middle) and nematic (right) phases in the presence of a stabilising field.

indeed occur and is seen at the next concentration increment (75 wt%). A biphasic region in the phase diagram becomes evident with the two phases corresponding to a region of pure water and a region of essentially pure chromonic material.

### 3.2 Arm length $L = 1$

Adding six ethylene oxide arms each with a length of a single bead, reduces phase transition temperatures slightly but does not dramatically change the thermotropic phase behaviour. Once again no ordering was observed on cooling from an isotropic liquid. However, for the solventless system, in an aligning field, transitions occur at  $T^* = 1.6$  (isotropic–nematic),  $T^* = 1.4$  (nematic–columnar),  $T^* = 0.6$  (columnar–crystalline). In comparison to the  $L = 0$  system the only notable differences are seen in a slight increase in the column spacing in the crystal and columnar phases, and a change in the in-plane ordering within the crystal. To accommodate the ethylene oxide groups the columns are spaced further apart. Additionally, since a single bead arm has very little conformational freedom, it fills space without distorting the molecule. Consequently, the columns are reorientated by approximately 20 degrees relative to the  $L = 0$  crystal (as seen in Fig. 3). In the columnar phase increases in the spacing between columns leads to more diffuse peaks in  $g(u,v)$  and less orientational order relative to the  $L = 0$  model.

Removal of the field only affected the thermotropic nematic phase stability. Both the crystal and columnar phases were stable with and without an aligning field; with the columnar phase melting directly to an isotropic phase on increasing temperature.

The  $L = 1$  system displays further phase changes upon the addition of water. The ethylene oxide arms add a favourable interaction between the molecule and water, stabilising the inclusion of water within a single phase. No two phase regions were observed, even down to a 25 wt% concentration. The columns clearly interact less with each other as the amount of solvent increases (see Fig. 4). At 75 wt% the 2d-orientational correlation of the pure melt has been lost. The ring pattern seen

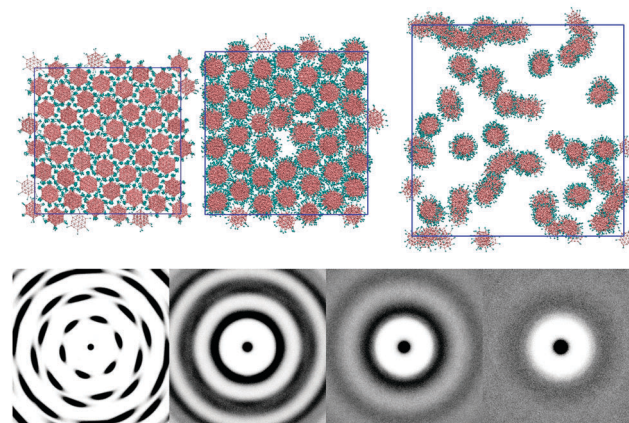


Fig. 4 Top: configurations of the  $L = 1$  system at 100 wt%, 75 wt% and 25 wt% (left middle and right, respectively) at  $T^* = 0.6$ ; brown sites show aromatic groups, blue sites show ethylene oxide segments and water is not shown for clarity. Bottom:  $g(u,v)$  for four concentrations 100 wt%, 75 wt%, 50 wt% and 25 wt% (left to right).



in  $g(u,v)$  is commensurate with a chromonic M phase (hexagonally arranged columns). At 50 wt% (N phase) the hexagonal arrangement of columns has disappeared. However, a ring structure persists, indicating that there is a weak column-column interaction. At 25 wt% (nematic-isotropic phase transition) all structure in the pair distribution function is lost apart from an exclusion zone around each core unit that no other columns can enter; indicating the columns randomly fill space.

### 3.3 Arm length $L = 2-3$

Increasing the arm length to two beads per arm adds a soft, distortable corona around each molecule. 2d-orientational order between columns is now lost for the thermotropic case: even at low temperatures ( $T^* = 0.5$ ) each molecule within a column has freedom to rotate around the molecular primary axis. At low temperatures we see therefore only a columnar phase (see Fig. 5). This phase can accommodate water by swelling to give both M (at 75 wt%) and N phases. The system shows a nematic phase at 50 wt% and 25 wt% but orientational ordering of columns is lost at 10 wt% giving an isotropic arrangement of aggregates.

Molecules with  $L = 3$  arms exhibit a very similar trend to the  $L = 2$  system. The exception occurs at 50 wt% concentration. Here, the phase is nematic but shows local hexagonal arrangement of columns for thin slices parallel to the director (see Fig. 6: top); suggesting that 50 wt% is on the cusp of a hexagonal-nematic columnar transition. The  $g(u,v)$  (see Fig. 6: bottom) shows that in the thermotropic melt there is no in-plane orientational order within a column but the ring pattern and intensity corresponds to a hexagonal columnar phase. At 50 wt% the water volume reduces the packing forces between columns, allowing columns a greater configurational freedom and resulting in a pattern in which the columns appear to form strings in the 2D projections (Fig. 5 and 6). However, each column is fully solvated by water and remains independent of adjacent columns. The M phase is evident at 75 wt% and, as before, at 25 wt% the pair distribution function shows no positional order in the cross section normal to the columnar director.

### 3.4 Arm length $L = 4-5$

Arm lengths of  $L = 4$  and  $L = 5$  show similar behaviour to the  $L = 3$  system, with the exception that at 50 wt% an M phase is seen. There are two competing factors responsible for the stabilisation of ordered phases at lower concentrations: as the

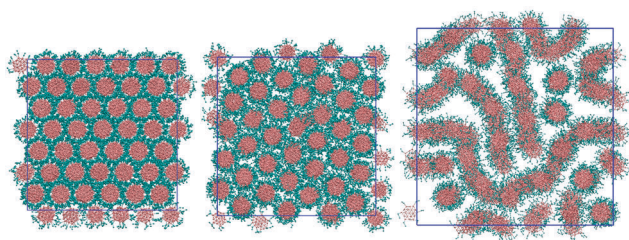


Fig. 5 Configurations of the  $L = 2$  model at 100 wt% (left), 75 wt% (middle) and 50 wt% (right).  $T^* = 1.0$ .

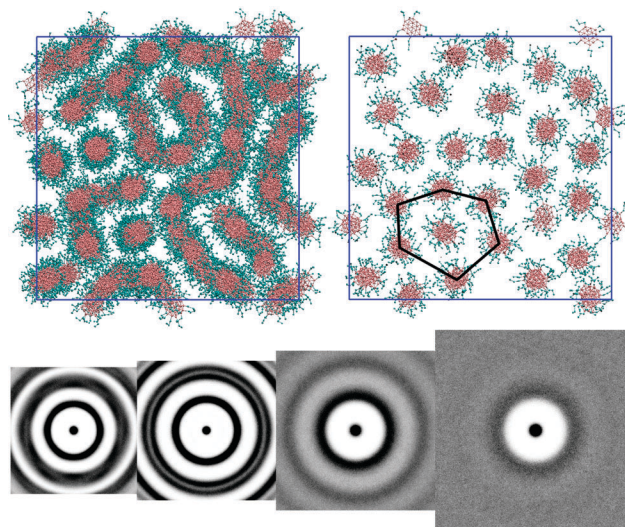


Fig. 6 Top (left): a configuration of the  $L = 3$  model at 50 wt%. Top (right): a slab of the same concentration,  $3.6l$  thick, with an irregular hexagonal overlay (right). Bottom:  $g(u,v)$  plots for  $L = 3$  model at four concentrations 100 wt%, 75 wt%, 50 wt% and 25 wt% (left to right).  $T^* = 1.0$ .

arms increase in length the fraction of rigid core present in the system decreases, acting to destabilise the phases relative to water content, this is counteracted by the longer arms allowing a higher capacity for swelling of a phase, and thereby incorporating more solvent.

### 3.5 Aggregation at low concentrations

To test the self-assembly properties of our model against experimental behaviour and to show how DPD parameters can be fine-tuned for chromonic systems, an additional series of lower concentration calculations were performed. Simulations with arm length of 3 beads ( $L = 3$ ) were selected for this experiment, since the behaviour of the  $L = 3$  model best represents our target experimental system TP6EO2M.<sup>15</sup> Simulations were performed at a concentration of 10 wt%, in the isotropic phase. At this concentration columns are sufficiently short to remove the possibility of periodically infinite columns and to reduce the frequency of any poorly sampled, larger aggregates.

The standard isodesmic model assumes that the equilibrium constant for binding an additional molecule is the same across each  $n$ -mer for each further binding event. This leads to the production of an exponential distribution of monomers in aggregates. In Fig. 7, we show the distribution of cluster sizes for different temperatures. At higher temperatures ( $T^* \geq 1.5$ ) monomers dominate, while at lower temperatures ( $T^* < 1.1$ ) larger aggregates dominate. In between there is a regime where the number of dimers exceeds the number of bare monomers.

The lower part of Fig. 7 shows the probability of finding molecules with zero bound faces *i.e.* monomers, molecules with one face bound representing either dimers or molecules at the end of an aggregate, and aggregates molecules with two bound faces. At high temperatures ( $T^* = 1.5$ )  $\sim 30\%$  of the molecules are monomers, with  $\sim 65\%$  end aggregate molecules and  $\sim 5\%$  mid aggregate molecules. As the temperature decreases we



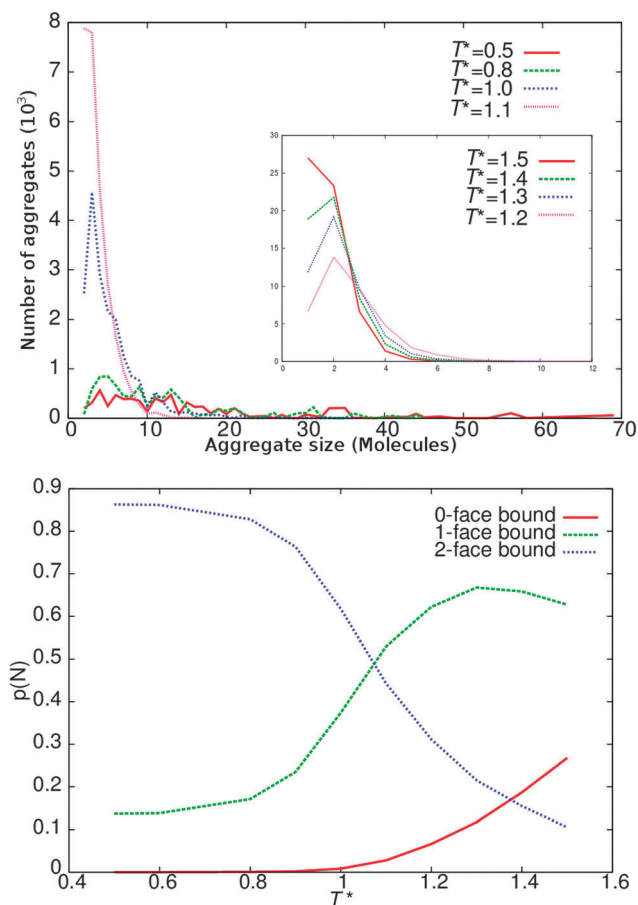


Fig. 7 Top: distribution of clusters of different sizes for  $T^* = 0.5, 0.8, 1.0, 1.1$ . Top (insert): distribution of clusters of different sizes for  $T^* = 1.5, 1.4, 1.3, 1.2$ . Bottom: probability of molecules having zero, one or two bound faces.

initially observe an increase in both forms of bound molecules and an associated decrease in monomers; indicating that more aggregates form and aggregates also increase in size. As the temperature is reduced further the monomer numbers continue to decrease while the number of molecules in a mid aggregate configuration increases. However, the number of end aggregate molecules decreases also; indicating (for fixed number of molecules) that fewer but larger aggregates have formed. At lower temperatures still ( $T^* = 0.8$ ) there are effectively no

monomers remaining and only a small number of end aggregate molecules.

It is interesting to compare the results seen here with the Monte Carlo simulations of a simple disc model for a chromonic system by Edwards *et al.*<sup>19</sup> They note that a modification to the usual isodesmic association model is required to explain their results, whereby the equilibrium constant becomes concentration dependent. However, the results in the current study are more consistent with a different type of modification to isodesmic association, in which there is more than one binding event. In our case initial formation of dimers is favoured over addition of further monomers. The latter appears also to be seen in some experimental systems. For sunset yellow, Chami and Wilson<sup>3</sup> have attempted to quantify this by calculating free energies of binding to aggregate stacks within an atomistic simulation model. In their work a small preference is seen ( $\sim 2 \text{ kJ mol}^{-1}$ ) for aggregation of two monomers to form a dimer, in comparison to addition of a monomer to a larger (dimer or above) aggregate. In the DPD model used here, all interactions are repulsive, consequently a small preference arises naturally for dimer formation due to the absence of next nearest neighbour repulsions (particularly arising from the flexible chains).

To quantify this further we define equilibrium constants for the binding process

$$K_n = \frac{[C_n]M^0}{[C_{n-1}][C_1]}, \quad (5)$$

where  $[C_n]$  is the concentration of aggregates of size  $n$  and  $M^0$  is a standard state of 1 M. Examination of the equilibrium constants in Table 1 show that there is indeed a small difference in the monomer to dimer equilibrium constant compared to the  $(n - 1)$ mer to  $n$ -mer binding; thus explaining the lower than expected monomer concentrations of Fig. 7. To a lesser extent we also note the formation of a trimer is also slightly favoured over higher aggregates.

For a standard constant- $NpT$  simulation binding enthalpies can be extracted directly from the slope of a Van't Hoff plot. Our DPD simulations are in the constant- $NVT$  ensemble but given

$$\left(\frac{\partial \ln K}{\partial T}\right)_p = \left(\frac{\partial \ln K}{\partial T}\right)_V - \left(\frac{\partial \ln K}{\partial p}\right)_T \left(\frac{\partial p}{\partial T}\right)_V \quad (6)$$

Table 1 Variation of simulation pressure and equilibrium constants for formation of dimers and higher aggregates as a function of temperature.  $n$ -mer equilibrium constants and standard errors are calculated over  $7n$ -mers for  $T^* = 1.5$  and respectively for 7, 8, 10, 15, 18 and 19 aggregates for the additional temperatures given

$T^*$	$p^*$	$\frac{[C_2]M^0}{[C_1][C_1]}/10^{-5}$	$\frac{[C_3]M^0}{[C_2][C_1]}/10^{-5}$	$\frac{[C_4]M^0}{[C_3][C_1]}/10^{-5}$	$\frac{[C_5]M^0}{[C_4][C_1]}/10^{-5}$	$\frac{[C_n]M^0}{[C_{n-1}][C_1]}/10^{-5}$
1.5	26.80	3.2	1.0	0.77	0.76	$0.91 \pm 0.1$
1.4	26.44	6.1	2.0	1.4	1.4	$1.5 \pm 0.1$
1.3	26.08	13.7	4.2	3.0	2.6	$2.9 \pm 0.27$
1.2	25.57	30.9	10.3	7.5	5.8	$7.2 \pm 0.98$
1.1	25.54	100.8	35.4	21.0	21.1	$22.1 \pm 2.6$
1.0	25.04	379.5	220.9	77.4	91.9	$103.8 \pm 14$
0.9	24.71	3771.4	688.8	282.1	540.5	$485.0 \pm 47$



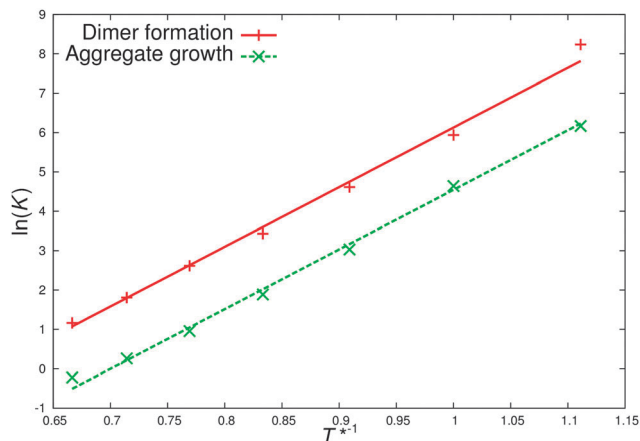


Fig. 8 Van't Hoff plots for dimer formation and aggregate growth. The slopes lead to an almost identical association enthalpy of  $-15.2RT$  for dimer and aggregate growth but with differing entropies,  $-9.0R$  and  $-10.6R$ .

and  $(\partial \ln K / \partial p)_T$  is small, and additionally our system size is relatively large, we can neglect the second term on the right hand side of eqn (6), writing

$$\left(\frac{\partial \ln K}{\partial T}\right)_p \approx \left(\frac{\partial \ln K}{\partial T}\right)_v \approx \frac{\Delta H^\circ}{RT^2}, \quad (7)$$

or alternatively

$$\left(\frac{\partial \ln K}{\partial(1/T)}\right)_v \approx \frac{-\Delta H^\circ}{R}, \quad (8)$$

Fig. 8 shows the Van't Hoff plots for dimer formation and growth of aggregates. Within errors the slopes lead to almost identical values for  $\Delta H^\circ$  of  $-15.2RT$ , indicating, for this model, that the less favourable association to form aggregates (in comparison to dimers) arises from entropic factors; most likely due to neighbour interactions in the aggregate restricting conformational freedom.

The association enthalpy in our model is easily tuned by the DPD repulsion parameter  $a_{ij}$ . This means that it should, in future work, be possible to tune the association behaviour of models such as this by reference to small atomistic simulations in dilute solution.

## 4 Conclusions

A simple DPD model has been developed to study chromonic self-assembly and phase formation in the non-ionic chromonic TP6EO2M. The computational speed of the model allows the phase diagram to be probed across a range of concentrations from pure mesogen to dilute solution. Depending on concentration and the length of the hydrophilic chain, we see the formation of a crystal phase, a columnar phase, and a stable chromonic nematic phase; in addition to association to give an isotropic arrangement of chromonic aggregates at low concentrations.

In dilute solution, association behaviour is approximately idodesmic, leading to an approximate exponential distribution

of monomers in aggregates. However, we note a small deviation from idodesmic behaviour in the case of dimers. Here, as in some experimental systems, entropic factors lead to the association of two monomers being slightly more favourable than aggregate growth. This probably arises because chains belonging to a molecule in the middle of a stack are restricted by the presence of two neighbours, whereas chains belonging to end molecules are only restricted by a neighbour on one side.

## Acknowledgements

The authors would like to acknowledge the support of the Engineering and Physical Sciences Research Council (EPSRC) in the UK through grants EP/J004413/1 and EP/J004707/1.

## References

- 1 J. Lydon, *Curr. Opin. Colloid Interface Sci.*, 2004, **8**, 480–490.
- 2 J. Lydon, *J. Mater. Chem.*, 2010, **20**, 10071–10099.
- 3 F. Chami and M. R. Wilson, *J. Am. Chem. Soc.*, 2010, **132**, 7794–7802.
- 4 D. J. Edwards, J. W. Jones, O. Lozman, A. P. Ormerod, M. Sinyureva and G. J. T. Tiddy, *J. Phys. Chem. B*, 2008, **112**, 14628–14636.
- 5 V. R. Horowitz, L. A. Janowitz, A. L. Modic, P. A. Heiney and P. J. Collings, *Phys. Rev. E: Stat., Nonlinear, Soft Matter Phys.*, 2005, **72**, 041710.
- 6 M. R. Tomasik and P. J. Collings, *J. Phys. Chem. B*, 2008, **112**, 9883–9889.
- 7 A. J. Dickinson, N. D. LaRacune, C. B. McKitterick and P. J. Collings, *Mol. Cryst. Liq. Cryst.*, 2009, **509**, 9[751]–20[762].
- 8 D. Perahia, E. J. Wachtel and Z. Luz, *Liq. Cryst.*, 1991, **9**, 479–492.
- 9 K. V. Kaznatcheev, P. Dudin, O. D. Lavrentovich and A. P. Hitchcock, *Phys. Rev. E: Stat., Nonlinear, Soft Matter Phys.*, 2007, **76**, 061703.
- 10 S. V. Shiyonovskii, O. D. Lavrentovich, T. Schneider, T. Ishikawa, I. I. Smalyukh, C. J. Woolverton, G. D. Niehaus and K. J. Doane, *Mol. Cryst. Liq. Cryst.*, 2005, **434**, 587–598.
- 11 S. V. Shiyonovskii, T. Schneider, I. I. Smalyukh, T. Ishikawa, G. D. Niehaus, K. J. Doane, C. J. Woolverton and O. D. Lavrentovich, *Phys. Rev. E: Stat., Nonlinear, Soft Matter Phys.*, 2005, **71**, 020702.
- 12 H. S. Park, A. Agarwal, N. A. Kotov and O. D. Lavrentovich, *Langmuir*, 2008, **24**, 13833–13837.
- 13 E. A. Mills, M. H. Regan, V. Stanic and P. J. Collings, *J. Phys. Chem. B*, 2012, **116**, 13506–13515.
- 14 P. K. Maiti, Y. Lansac, M. A. Glaser and N. A. Clark, *Liq. Cryst.*, 2002, **29**, 619–626.
- 15 Z. H. Al-Lawati, B. Alkhairalla, J. P. Bramble, J. R. Henderson, R. J. Bushby and S. D. Evans, *J. Phys. Chem. C*, 2012, **116**, 12627–12635.
- 16 R. E. Hughes, S. P. Hart, D. A. Smith, B. Movaghar, R. J. Bushby and N. Boden, *J. Phys. Chem. B*, 2002, **106**, 6638–6645.





- 17 N. Boden, R. J. Bushby, L. Ferris, C. Hardy and F. Sixl, *Liq. Cryst.*, 1986, **1**, 109–125.
- 18 R. Hentschke, P. J. B. Edwards, N. Boden and R. J. Bushby, *Macromol. Symp.*, 1994, **81**, 361–367.
- 19 R. G. Edwards, J. R. Henderson and R. L. Pinning, *Mol. Phys.*, 1995, **86**, 567–598.
- 20 P. Prybytak, W. J. Frith and D. J. Cleaver, *Interface Focus*, 2012, **2**, 651–657.
- 21 P. J. Hoogerbrugge and J. M. V. A. Koelman, *EPL*, 1992, **19**, 155.
- 22 R. D. Groot, T. J. Madden and D. J. Tildesley, *J. Chem. Phys.*, 1999, **110**, 9739–9749.
- 23 A. A. Gavrilov, Y. V. Kudryavtsev and A. V. Chertovich, *J. Chem. Phys.*, 2013, **139**, 224901.
- 24 L. He, Z. Chen, R. Zhang, L. Zhang and Z. Jiang, *J. Chem. Phys.*, 2013, **138**, 094907.
- 25 E. S. Boek, P. V. Coveney, H. N. W. Lekkerkerker and P. van der Schoot, *Phys. Rev. E: Stat. Phys., Plasmas, Fluids, Relat. Interdiscip. Top.*, 1997, **55**, 3124–3133.
- 26 S. Yamamoto, Y. Maruyama and S. Hyodo, *J. Chem. Phys.*, 2002, **116**, 5842–5849.
- 27 J. C. Shillcock and R. Lipowsky, *J. Chem. Phys.*, 2002, **117**, 5048–5061.
- 28 H. Xu, Q. Zhang, H. Zhang, B. Zhang and C. Yin, *J. Theor. Comput. Chem.*, 2013, **12**, 1250111.
- 29 X. Chen, F. Tian, X. Zhang and W. Wang, *Soft Matter*, 2013, **9**, 7592–7600.
- 30 A. AlSunaidi, W. K. Den Otter and J. H. R. Clarke, *Philos. Trans. R. Soc., A*, 2004, **362**, 1773–1781.
- 31 A. AlSunaidi, W. K. den Otter and J. H. R. Clarke, *J. Chem. Phys.*, 2013, **138**, 154904.
- 32 M. Bates and M. Walker, *Soft Matter*, 2009, **5**, 346–353.
- 33 M. A. Bates and M. Walker, *Phys. Chem. Chem. Phys.*, 2009, **11**, 1893–1900.
- 34 M. A. Bates and M. Walker, *Mol. Cryst. Liq. Cryst.*, 2010, **525**, 204–211.
- 35 M. A. Bates and M. Walker, *Liq. Cryst.*, 2011, **38**, 1749–1757.
- 36 R. D. Groot and P. B. Warren, *J. Chem. Phys.*, 1997, **107**, 4423–4435.
- 37 M. A. Seaton, R. L. Anderson, S. Metz and W. Smith, *Mol. Simul.*, 2013, **39**, 796–821.
- 38 DL\_MESO is a mesoscale simulation package written by R. Qin, W. Smith and M. A. Seaton and has been obtained from STFC's Daresbury Laboratory via the website [http://www.ccp5.ac.uk/DL\\_MESO](http://www.ccp5.ac.uk/DL_MESO).
- 39 S. D. Stoddard, *J. Comput. Phys.*, 1978, **27**, 291–293.

