Isomeric trimesogens exhibiting modulated nematic mesophases†

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Modulated nematic phases, in which the constituent molecules possess some form of superstructure in addition to the average orientational organization and lack positional order present in classical nematic phases, are of current fascination in liquid crystal research. Most of the recent focus has been on the twist-bend nematic phase. This phase exhibits phenomena that are of fundamental importance to science; the spontaneous formation of a helical pitch of nanometre scale in a fluid and the spontaneous breaking of mirror symmetry leading to the emergence of chiral domains in an achiral system. In this communication we report on the synthesis of two liquid-crystalline trimesogens whose structures are an extension of the known dimeric material CB6OCB. One material (A63) exhibits a nematic phase with a twist-bend modulation (N_{TB} phase), whereas a structurally isomeric material (B63) exhibits an unusual nematic like mesophase which we believe is a candidate for being a new form of modulated nematic. Whereas the properties of dimeric LCs are somewhat indifferent to chemical composition, the stark difference in behaviour between the two materials herein shows that oligomers may exhibit an unexpected sensitivity to molecular structure.

Experimental

In our previous attempt to synthesis A63, we were unable to cleave the benzyl ether without accompanying nitrile reduction,40 in the present work we circumvented this issue by moving to the use of a tert butyldimethylsilyl (TBDMS) protective group. The phenol group of the intermediate i1, synthesis reported in,40 was protected with a TBDMS group and i2. Mitsunobu etherification with i4 gave i5; cleavage of the TBDMS group was achieved using TBAF. Etherification of i6 with i7 afforded the trimesogen A63.

The trimesogen B63, an isomer of A63, was prepared via Mitsunobu etherification of i7 and i8 as shown in Scheme 2. Synthetic details, chemical characterisation and details of instrumentation used are given in the ESI† to this communication; both trimesogenic materials were obtained in high purity (99.4% for A63 and 99.7% for B63) as evidenced by HPLC.
Results and discussion

The transition temperatures of both $A_{63}$ and $B_{63}$ were determined using a combination of polarized optical microscopy (POM), differential scanning calorimetry (DSC) and small angle X-ray scattering (SAXS). Transition temperatures and associated enthalpies of transition for each material are presented in Table 1.

The trimesogen $A_{63}$ can be envisaged as an extension of the well-known liquid-crystalline dimer CB6OCB (Cr 102 N$_{TB}$ 110.5 N 154.2 Iso); it is unsurprising therefore that this material mirrors the behaviour of the parent compound and exhibits nematic and twist-bend phases, although we note that the trimesogen ($A_{63}$) has significantly higher melting points and transition temperatures than the parent (CB6OCB), while the associated enthalpy associated with the N-N$_{TB}$ phase transition is an order of magnitude larger in $A_{63}$. The identification of the nematic phase was trivial based on its characteristic schlieren texture (Fig. 2a), high fluidity, Brownian motion, and diffuse scattering of X-rays at small angles (Fig. 2d and SI-7†). With further cooling a transition into the twist-bend phase occurred; regions exhibiting a schlieren texture in the nematic phase exhibited a “blocky” texture (Fig. 2b), planar aligned regions transformed into the rope texture (Fig. 2c) although homeotropic regions remained optically extinct. When the rope texture was annealed for 5 minutes at a constant temperature of 125 °C we observed striations forming at an angle of ~30° to the ‘length’ of the rope; the possibility for the formation of fibre-like structures in the N$_{TB}$ has been noted previously$^{43}$ and recently demonstrated to have a “fish-bone” like structure from SEM imaging of photopolymerised twist-bend dimers.$^{43}$

When studied by SAXS the N$_{TB}$ phase of $A_{63}$ exhibited a diffuse scattering at small angles (Fig. 2e). The small angle scattering peak had a $d$-spacing of 14.1 Å which was considered as a first approximation to be roughly 1/3 of the molecular length. DFT (B3LYP/6-31G(d)) calculations show that if the two outer mesogenic units are trans to each other then the molecular length is 43.5 Å compared with a value of 39 Å in the comparable cis geometry, however this neglects the flexibility inherent in such systems and is best considered as an approximation. The $d$-spacing values in both phases, taken in conjunction with the calculated molecular lengths, indicate the sub-units of the trimesogen behave essentially as free entities in both phases rather than the trimesogen behaving as one large ‘rod-like’ molecule.

As mentioned briefly, the use of a single conformer neglects the inherent flexibility afforded by the hexyloxy spacer, present in both materials. Using fully relaxed scans (at the DFT (B3LYP/6-31G(d)) level in Gaussian 09 revision e01) we studied a 1/2 fragment of both trimers. We allow each dihedral shown in Fig. 3a to rotate in 10° steps, giving plots of conformer energy as a function of each individual angle. For the 2nd dihedral we observe the gauche states to be lower in energy than the trans form ($\Delta E$, 0.6 kJ mol$^{-1}$), for the other dihedrals the trans form is lower than gauche by 2–6 kJ mol$^{-1}$. While calculation of the full conformer distribution for both materials is too computationally expensive to be feasible, even using the RIS approximation and semi-empirical methods, the present study illustrates the flexibility inherent to these systems.

Given the similarities in molecular structure it had been anticipated that $B_{63}$ would exhibit similar transition temperatures and phase behaviour to $A_{63}$. $B_{63}$ exhibits lower transition temperatures than $A_{63}$, with identification of the nematic phase being as described previously. Upon cooling into the next phase (N$_X$) a series of unusual optical textures were observed, which we
Radial integration of each SAXS pattern gave scattered intensity as a function of $q$; fitting the raw scattering data with a Lorentzian function gave the $d$-spacings of the small- (001) and wide- (100) angle peaks (Fig. 4f), which we find to be temperature invariant. The small-angle scattering peak has a $d$-spacing of $\sim 14$ Å in both phases; we do not observe any scattering events at smaller angles up to the limit of our experimental setup at $\sim 100$ Å. We obtain molecular lengths at the DFT (B3LYP/6-31G(d)) level; as with $A6_3$, we find that if the two outer mesogenic units are trans to each other the molecular length is 42.7 Å compared with a value of 38.5 Å in the comparable cis geometry, i.e. the $d$-spacing of the small angle peak corresponds to $\sim 1/3$ of the molecular length. Although this neglects the flexibility of $B6_3$, but the present results indicate that, as with $A6_3$, the trimesogen does not behave as one large ‘rod-like’ molecule. The correlation lengths were determined from the FWHM of each peak; whereas the correlation length parallel to the director decreases slightly at the N-NX phase transition the perpendicular correlation length increases marginally, however we find the magnitude of both correlation lengths is consistent with a nematic-like phase. Lastly, we plot the maximum intensity of both small- and wide-angle peaks as a function of temperature, we observe that there is a decrease in the intensity small-angle scattering at the N-NX transition however there is a corresponding increase in the intensity wide-angle scattering.

Good quality planar alignment was achieved for both compounds when filled into cells with a buffered antiparallel alignment layer (purchased from Instec thickness $\sim 5$ μm). The

describe as follows. On an untreated glass slide we observed that regions exhibiting a schlieren texture initially take on a patchy appearance at the phase transition into the NX phase before then forming a nondescript, sand-like texture (Fig. 4a–c). We performed SAXS on $B6_3$, collecting frames on cooling in 1.2 °C intervals from the clearing point until the sample crystallised at $\sim 103$ °C. The nematic nature of both phases, hinted at by the diffuse small angle peak in SAXS (2D patterns in Fig. 4d and e, data in Fig. 4f–h). Both the onset temperature and the enthalpy associated with the transition, was confirmed by the diffuse small angle peak in SAXS (2D patterns in Fig. 4d and e, data in Fig. 4f–h). Both the onset temperature and the enthalpy associated with the transition were found to be invariant with the heat/cool rate employed in DSC studies – unlike for the dimer CB11CB where the NTB-NX transition appears almost second order and resolves into a small first order peak with varying scan rate,\(^{45}\) although this phenomenon is not repeated by either material in this present work.

Fig. 2 Characterisation of $A6_3$: photomicrographs (×100, crossed polarisers) of (a) the nematic phase of $A6_3$ at 135 °C; (b) the same region as a cooled into the twist-bend phase at 128 °C; (c) the rope texture of the twist-bend phase at 128 °C; (d) 2D SAXS pattern for the nematic phase of $A6_3$ at 145 °C; (e) 2D SAXS pattern for the NTB phase of $A6_3$ at 125 °C – note due to a tendency for sample crystallisation the exposure time in the NTB phase was shorter (30 seconds vs. 300 seconds) leading to a poorer signal-to-noise ratio.

### Table 1 Transition temperatures and associated enthalpies of the isomeric trimesogens $A6_3$ and $B6_3$ as determined using DSC at a heat/cool rate of 10 °C. Note, presented values are the average of 3 cycles

<table>
<thead>
<tr>
<th>Cpd</th>
<th>X</th>
<th>Y</th>
<th>MP</th>
<th>$N_{X}$</th>
<th>$N_{NTB}$</th>
<th>$N_{Iso}$</th>
</tr>
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<tbody>
<tr>
<td>A6_3</td>
<td>-CH$_2$-</td>
<td>-O-</td>
<td>7 °C</td>
<td>140.9</td>
<td>133.3</td>
<td>170.9</td>
</tr>
<tr>
<td>$\Delta H$ [kJ mol$^{-1}$]</td>
<td>30.0</td>
<td>30.0</td>
<td>30.0</td>
<td></td>
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$B6_3$ | -O- | -CH$_2$- | 118.4 | 116.4 | 156.7 |
| $\Delta H$ [kJ mol$^{-1}$] | 16.7 | 1.4 | 2.0 |
Fig. 3  The dihedrals shown in (a) were allowed to rotate in 10 °C steps using fully relaxed scans at the B3LYP/6-31G(d) level of DFT as implemented in Gaussian G09 revision e.01, the plots (b–g) show the energy of as a function of each dihedral relative to that of the lowest energy conformer; (b) dihedral 1; (c) dihedral 2; (d) dihedral 3; (e) dihedral 4; (f) dihedral 5; (g) dihedral 6.

Fig. 4  Characterisation of B62: photomicrographs (×100, crossed polarisers) of (a) the nematic phase at 145 °C; (b) the transition into the N_x phase at 118.5 °C; (c) the N_x phase at 116 °C; (d) 2D SAXS pattern of the nematic phase at 137 °C; (e) 2D SAXS pattern of the N_x phase at 111 °C; (f) plot of the d-spacing of the 001 (small angle) and 100 (wide angle) peaks in SAXS as a function of reduced temperature; (g) plot of the correlation length parallel (ζ_∥) and perpendicular (ζ_⊥) to the director as a function of reduced temperature; (h) plot of the intensity (arbitrary units) of the small- and wide-peaks during SAXS as a function of reduced temperature. In plots (f–h) the dashed line corresponds to the N–N_x phase transition.
behaviour of A63 was in keeping with our studies on CB11CB in the same confined geometry; the well aligned planar texture in the nematic phase (Fig. 5a) gives way to domains of differing handedness at the N_{TB} transition (Fig. 5b), with further cooling giving a rope-like texture perpendicular to the rubbing direction of the cell (Fig. 5c). This behaviour is typical for the N_{TB} phase. The isomeric trimesogen B63 also afforded a well aligned planar texture in the nematic phase, however with further cooling a blocky texture emerged (Fig. 5d) which slowly transformed into a parabolic texture (Fig. 5e). Some regions formed a quasi-planar texture featuring regions with different handed domains (Fig. 5f); this is reminiscent of the twist-bend phase, however, this texture in the twist-bend phase gives way to the rope texture (Fig. 5b and c) yet in the N_{X} phase of B63 we did not observe a fully-fledged rope texture. Whereas the optical texture of the N_{TB} phase changes significantly with cooling, the changes in the optical texture of the N_{X} phase exhibited by B63 are less pronounced.

We next subjected a sample of B63 to slow cooling across the N–N_{X} phase transition in order to obtain superior optical textures, with a view to ruling out this phase as simply being a twist-bend nematic. The typical schlieren texture of the nematic phase (Fig. 6a) is made up of disclinations, following the transition into the N_{X} phase these lines become broken and fractal (Fig. 6b). At the N–N_{TB} phase transition planar aligned regions adopt a blocky texture (for example Fig. 2b), whereas at the N–N_{X} a curved, fractal texture is obtained (Fig. 6b).

Additionally the N_{TB} phase exhibits focal-conic defects, whereas none have been observed in the optical textures of the N_{X} phase of B63. With further cooling there is a significant change in the birefringence of the sample, with long curved domains appearing (Fig. 6c). The sample could easily be sheared to give a fairly uniform homeotropic texture (Fig. 6d).

Uniform homeotropic alignment of both the nematic and N_{X} phases of B63 was realised using glass slides treated with piranha solution (30% hydrogen peroxide and 95% sulphuric acid in a 1 : 3 ratio); subsequent conoscopy demonstrates the N_{X} phase to be optically uniaxial (Fig. 6e) and, a pronounced change in the birefringence of the sample, with long curved domains appearing (Fig. 6c). The sample could easily be sheared to give a fairly uniform homeotropic texture (Fig. 6d).

Conoscopy gives crucial information about the local structure of the N_{X} phase; firstly as this phase is optically uniaxial it is implied that the modulation period (P_{mod}) is much shorter than the wavelength of visible light (P_{mod} ≪ λ). As the phase is optically positive the molecules must be tilted away from the director by less than the magic angle. Although the twist-bend phase would satisfy both of these observations, as well as the diffuse nematic-like SAXS patterns, the optical textures observed for B63 are sufficiently distinct from those of A63 and other twist-bend materials to suggest this is not the case.

If the N_{X} phase has a structure with modulation of electron density (e.g. SmA, splay-bend nematic) Bragg scattering should be observed; as this has not been observed it raises two
possibilities for the structure of the Nₓ phase exhibited by B₆₃: the modulation period is large enough that the corresponding Bragg scattering peak (and its harmonics) occur at larger d-spacings than those we can observe using our SAXS setup (~100 Å). Alternatively, if there is no modulation of electron density associated with the Nₓ phase (similar to that observed for the Nₜₜₖ) then non-resonant X-ray experiments are blind to the modulation period. We note the possibility that the structure may have no modulation whatsoever; we consider this unlikely given the changes in optical textures. The change in SAXS peak intensity as well as the associated enthalpy of transition of the two materials are comparable, though not identical, for both materials and to previously reported trimesogens exhibiting the twist-bend phase.³⁴,³⁶

In the original work on the CBₙCB materials the phase now identified as being Nₜₜₖ was often identified as being smectic, presumably because of its smectic-like optical textures resulting from its periodic structure. We note that the trimesogen shown in Fig. 7, described 20 years ago by both Zuev and Imrie in separate publications, is reported as exhibiting nematic and smectic phases. Given the similarity of the molecular structure of this material to A₆₃ and B₆₃ a thorough re-examination of such materials may be warranted.

Conclusions

Two trimesogenic liquid-crystalline materials – both isomers of one another – have been prepared and characterised. The method used to prepare these materials should finally allow access to chemically stable oligomeric materials of any given length, and polydispersity approaching unity. One material (A₆₃) exhibits nematic and twist-bend mesophases, however its counterpart material (B₆₃) exhibits a nematic-like mesophase.
structure for which we have not been able to provide a definitive identity. Studies to date point to a modulated nematic phase, with a small modulation period, and although it is possible that the observed ‘N’ phase may be simply a twist-bend nematic, the potential immiscibility of the two materials as well as the remarkably different optical textures suggest this is not the case and thus a new liquid crystal phase may be a possibility that warrants further study on this material, its analogues and higher oligomeric homologues.

**Conflicts of interest**

There are no conflicts of interest to declare.

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**References**