Spontaneous mirror symmetry breaking in benzil-based soft crystalline, cubic liquid crystalline and isotropic liquid phases†

Tino Reppe,a Silvio Poppe,a Xiaoqian Cai,b Yu Cao,b Feng Liu,*,b and Carsten Tschierske✉,a

Benzil (diphenylethane-1,2-dione), which is a long known example for an achiral molecule crystallizing in a chiral space group, can also show mirror symmetry breaking in the fluid state if it is suitably functionalized. For some of the new benzil derivatives even three different subsequent mirror symmetry broken soft matter states with a chiral conglomerate structure can be observed. One is an isotropic liquid, the second one a cubic liquid crystal with a complex network structure and the third is a soft crystalline solid. Chirality develops by helical self-assembly combined with dynamic network formation, thus allowing macroscopic chirality synchronization. These achiral molecules, combining a transiently chiral bent core with multiple alkyl chains, provide a unique link between the mirror symmetry breaking phenomena observed for polycatenar and bent-core mesogens. The homogeneously chiral networks are of interest for application as chiral materials, and as templates for chiral recognition, separation and enantioselective catalysis.

Likewise, spontaneous chirality can also emerge during the formation of liquid crystalline (LC) phases and even in liquids due to a dynamic process of chirality synchronization, which is the subject of this contribution.

One of the most prominent examples of achiral molecules crystallizing in a chiral space group is benzil (1, Fig. 1a), whose OA has been more closely studied than for any other organic crystal.⁷ In the chiral crystal structure the molecules assume a helical conformation with a torsion angle of the O=C–C=O bond of about α = 110°. This twisted (skewed) cisoid conformation is also found as a preferred conformation in the gas phase⁸ and in solution.⁹ However, the energy barrier for the rotation around the O=C–C=O bond is relatively low and in solution or in the melted state it leads to optical inactivity. Thus benzil is a typical transiently chiral molecule capable of mirror symmetry breaking in the crystalline state by Viedma ripening.¹¹

Recently, we have observed emergence of OA in bicontinuous cubic (Cub₅₁) LC phases¹² (Fig. 1d) and even in isotropic liquids (Iso[5]) by formation of conglomerates of chiral domains.¹³ This OA is related to the helical nano-scale organization of the molecules in these Cub₅₁ and Iso[5] phases. The question arises whether the transient chirality of the benzil unit can support the development of this spontaneous dissymmetry. Moreover, the twisted conformation leads to a non-linear bent shape of the aromatic core, providing some similarity to bent-core molecules, which are also known for their capability of showing mirror symmetry breaking.¹⁴–¹⁶ In this case besides helix formation the reduction of the phase symmetry to C₂ᵥ, due

Mirror symmetry breaking and development of uniform chirality is considered essential for the development of life. Understanding and controlling the formation of chirality is crucial across different disciplines. In biosystems the chirality of nucleic acids, sugars and amino acids enables the formation of the helical nanostructures of DNA and proteins.¹ Enantioselective synthesis, catalysis and autocatalysis² were successfully developed during recent decades for the synthesis of enantiopure molecules from achiral precursors. This requires expensive chiral reagents or catalysts, synthesized in multistep processes from the chiral pool of nature. However, there is a more economic approach to homochirality, because it can be achieved spontaneously as a result of physical processes such as crystallization,⁴ as for example shown for the Viedma ripening,⁴ where rapidly enantiomerizing racemic mixtures of chiral molecules, or even achiral molecules crystallizing in a chiral space group spontaneously form uniform chiral crystals with optical activity (OA). The combination of generation of chiral centers with deracemization by dynamic crystallization provides a new absolute asymmetric synthesis methodology.⁵

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✉ Institute of Chemistry, Martin Luther University Halle-Wittenberg, Kurt-Mothes-Straße 2, 06120 Halle, Germany. E-mail: carsten.tschierske@chemie.uni-halle.de

a State Key Laboratory for Mechanical Behaviour of Materials, Shaanxi International Research Center for Soft Matter, Xi’an Jiaotong University, Xi’an 710049, P. R. China. E-mail: feng.liu@xjtu.edu.cn

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to the organization of the tilted molecules with their bent direction parallel to the layer planes (see Fig. S16†), contributes to symmetry breaking.14

Besides the well-known case of lyotropic LCs,17 Cub2i phases are also known to be formed in solvent-free (thermotropic) systems by rod-like polyaromatic molecules with two18,19 or multiple alkyl end-chains (polycatenar compounds),20 branched chains (swallow tailed compounds)21 or other bulky end groups.22 In these Cub2i phases the rods form networks and the alkyl chains fill the space between them (Fig. 1c and d). Within the networks the orientation of the rods is perpendicular or slightly tilted with respect to the local network direction and the mismatch of the cross sectional areas of these cores with the cross sectional area of the terminally attached chains leads to the curvature of the aggregates, and simultaneously, to a helical twist along the networks (Fig. 1b).6,12a There are two major types of such thermotropic Cub2i phases, the double gyroid phase (space group Ia3d) and the triple network phase designated as Im3m (Fig. 1c and d).18,21,24 The Ia3d phase involves two helical networks with opposite handedness (red, blue in Fig. 1c) and therefore it is achiral.12a However, in the Im3m phase this degeneracy is broken by the presence of three networks24 and the synchronization of the helix sense between them.12a The chiral space group with the highest symmetry, derived from the achiral Im3m space group would be Ia32,24 but the actual space group was recently identified as I23.25 In this new structural model of the “Im3m” type Cub2i phase there are three networks with all junctions being threefold, but with distances and twist angles between the junctions being slightly different from those in the Ia3d phase (Fig. 1d).

Herein we report the first mirror symmetry broken LC phases formed by a series of suitably designed multi-chain benzil derivatives (Tables 1 and 2). The decyloxy substituted compounds with a different number of chains are labelled as 2–6 according to their order in Table 1. Compounds 3 have three decyloxy chains at one end and either no chain (compound 3/H) or a single n-alkyloxy chain with a variable length at the other end (compounds 3/n); here the number n after the slash indicates the length of this chain (Table 2). The respective compounds were synthesized as described in Scheme S1† and the experimental procedures are given in the ESI.† Analysis of their self-assembly was performed by polarizing optical microscopy (POM), differential scanning calorimetry (DSC) and X-ray diffraction (XRD). As is obvious from Table 1, with the growing number of n-decyloxy chains a transition from lamellar (SmC, 2) via two different types of Cub2i phases (Ia3d, I23[6], 3/10, 4) to hexagonal columnar phases (Colhex, 4, 5) is observed. Only the hexaalkyl substituted compound 6 does not show any LC phase. The SmC and Colhex phases were indicated by their typical birefringent optical textures observed between crossed polarizers (Fig. 2e, h and S2–S5†) and were confirmed by XRD for compounds 2 and 5 (Table S6†). The transition from the isotropic liquid state (Iso) to the Cub2i phases was indicated by a small exotherm in the DSC scans, accompanied by an increase in the viscosity while the phase remained optically isotropic. The cubic phase types are distinguished by optical investigation between polarizers, rotated by a small angle (∼5°) out of the 90° crossed orientation, where only the I23 phase shows a conglomerate of optically active (dark and bright) domains (I23[6] phase), whereas Ia3d is optically inactive (Fig. 2f and g).15 Interestingly, the Cub2i phase type depends on the chain distribution and the chain length (Tables 1 and 2); for the symmetric compound 4 the achiral Ia3d phase occurs as a monotropic (metastable) phase below Colhex, whereas the non-symmetric compound 3/10 has exclusively a broad range of the mirror symmetry broken I23[6] phase. To the best of our knowledge, these are the first benzil derivatives forming cubic LC phases, whereas a few benzil based compounds involving amide groups were found to form exclusively columnar LC phases.26

Here we focus on the homologous series of the non-symmetric tetracatenars 3/n where only 3/H and 3/1 are non-mesomorphic and all the following homologues form Cub2i phases (Table 2). In this series the Cub2i phase type changes...
Table 1: Phase transitions of compounds 2–6 on heating

<table>
<thead>
<tr>
<th>Compd.</th>
<th>A</th>
<th>B</th>
<th>D</th>
<th>E</th>
<th>T°C [ΔH kJ mol⁻¹]</th>
</tr>
</thead>
<tbody>
<tr>
<td>2</td>
<td>H</td>
<td>H</td>
<td>H</td>
<td>H</td>
<td>Cr 149 [27.0] SmCₐ 243 [12.0] Iso</td>
</tr>
<tr>
<td>4</td>
<td>R</td>
<td>H</td>
<td>R</td>
<td>H</td>
<td>Cr 151 [69.5] Cub₃/a23 141 Colhex 143 [12.4] Iso</td>
</tr>
<tr>
<td>5</td>
<td>R</td>
<td>R</td>
<td>R</td>
<td>H</td>
<td>Cr 92 [14.2] Colhex d 105 [7.1] Iso</td>
</tr>
<tr>
<td>6</td>
<td>R</td>
<td>R</td>
<td>R</td>
<td>R</td>
<td>Cr 84 [33.2] Iso</td>
</tr>
</tbody>
</table>

a DSC peak temperatures at 10 K min⁻¹; values in round parantheses indicate monotropic phases, only observed on cooling; abbreviations: Cr = crystalline solid, SmC = tilted lamellar phase, Cub₃/a23 = achiral Cub₃ phase with the la3d space group, Cub₃/a23[8] = mirror symmetry broken Cub₃ phase with the la3d space group; Iso = achiral isotropic liquid; Iso₁[4] = mirror symmetry broken Iso phase; for DSC traces, see Fig. S1, and for textures, see Fig. S2–S5. b d = 3.66 nm. c Enthalpies of both transitions. d ahex = 4.39 nm.

Table 2: Phase transitions, lattice parameters (a_{cub}, nm), number of molecules in the neighbouring rafts (n_{raft}), and twist angles between the rafts (Φ") in the Cub₃ phases of compounds 3/H and 3/n

<table>
<thead>
<tr>
<th>3/n</th>
<th>T°C [ΔH kJ mol⁻¹]</th>
<th>a_{cub}/nm</th>
<th>n_{raft}</th>
<th>Φ&quot;</th>
<th>L_{mol}/nm</th>
</tr>
</thead>
<tbody>
<tr>
<td>3/H</td>
<td>H: Cr 126 [52.2] Iso</td>
<td>—</td>
<td>—</td>
<td>—</td>
<td>4.2</td>
</tr>
<tr>
<td></td>
<td>C: Iso 110 [27.3] Cr</td>
<td>—</td>
<td>—</td>
<td>—</td>
<td>4.3</td>
</tr>
<tr>
<td>3</td>
<td>H: Cr 127 [36.3] Iso</td>
<td>12.9</td>
<td>5.0</td>
<td>6.9</td>
<td>4.4</td>
</tr>
<tr>
<td></td>
<td>C: Iso 115 [36.4] Cr</td>
<td>18.4</td>
<td>4.0</td>
<td>7.6</td>
<td>4.6</td>
</tr>
<tr>
<td>3/2</td>
<td>H: Cr 120 [37.9] la3d(L) 127 [1.8] Iso, 135 [2.8] Iso</td>
<td>18.4</td>
<td>3.8</td>
<td>7.7</td>
<td>4.8</td>
</tr>
<tr>
<td></td>
<td>C: Iso 133 [2.7] Iso, 121 [0.2] Iso₁[4] 110 [0.3] la3d(L) 105 [31.3] Cr</td>
<td>18.4</td>
<td>3.7</td>
<td>7.6</td>
<td>5.2</td>
</tr>
<tr>
<td>3/4</td>
<td>H: Cr 128 [53.5] Iso</td>
<td>11.4</td>
<td>3.3</td>
<td>7.9</td>
<td>5.4</td>
</tr>
<tr>
<td>3/6</td>
<td>H: Cr 118 [64.6] (Cr₁₃[8] 82 [17.3] f23[4] 126 [2.2] Iso, 134 [5.7] Iso</td>
<td>11.5</td>
<td>3.2</td>
<td>7.8</td>
<td>5.8</td>
</tr>
<tr>
<td>3/12</td>
<td>H: Cr 114 [47.0] la3d(L) 135 [3.4] Iso, 139 [6.2] Iso</td>
<td>11.4</td>
<td>3.3</td>
<td>7.9</td>
<td>5.4</td>
</tr>
<tr>
<td></td>
<td>C: Iso 137 [8.9] Iso, 128 [0.3] Iso₁[4] 122 [0.7] la3d(L) 48 [17.9] Cr₁₃ &lt;20 Cr</td>
<td>11.3</td>
<td>3.2</td>
<td>7.8</td>
<td>5.6</td>
</tr>
<tr>
<td>3/16</td>
<td>H: Cr 108 [44.1] la3d(L) 135 [2.7] Iso, 138 [7.2] Iso</td>
<td>11.4</td>
<td>3.3</td>
<td>7.9</td>
<td>5.4</td>
</tr>
<tr>
<td></td>
<td>C: Iso 136 [9.2] Iso, 129 [1.4] Colhex 125 [0.5] la3d(L) 42 [16.9] Cr₁₃ &lt;20 Cr</td>
<td>11.5</td>
<td>3.2</td>
<td>7.8</td>
<td>5.8</td>
</tr>
</tbody>
</table>

a DSC peak temperatures on heating/cooling (H/C) at 10 K min⁻¹; n_{raft} = n_{cell} / (L_{mol} / 0.45) with L_{mol} = 20.68a_{cub}; and L_{mol} = 8.485a_{cub}; Φ(la3d) = 70.5°/ [0.3544a_{cub}/0.45 nm]; Φ(f23) = 90°/ [0.290a_{cub} /0.45 nm]; f_{max} = maximum molecular length as determined with space filling models assuming a molecular bend of 110° and all-trans alkyl chains, see Fig. S15; abbreviations: Iso₁ = achiral cybotactic isotropic liquid (see Fig. 3d), Cr = birefringent crystalline solid; Cr₁₃ = optically isotropic crystalline mesophase; Cr₁₃[8] = mirror symmetry broken Cr₁₃ forming a conglomerate of chiral domains; la3d(L) = short pitch la3d phase with a smaller number of molecules in the unit cell rafts and a larger twist angle Φ than in la3d(L) = long pitch la3d phase with a larger number of molecules in the unit cell rafts and a smaller twist angle Φ; for other abbreviations, see Table 1; chiral phases are shown in bold; because the transitions Iso-Iso₁ are very broad (see Table S1) it is difficult to determine precise enthalpy values; for DSC traces, see Fig. S1 and for XRD data, see Tables S1–S7 and Fig. S7 and S10–S14.
the strongest being indexed either to (211) and (220) of the Ia3d lattice or (321), (400) and (420) reflections of the I23[^1] lattice. The electron density maps (Fig. 2c, d and S9†) reconstructed from these patterns are in agreement with the proposed phase structures shown schematically in Fig. 1c and d. The method of selecting the correct phase combination is described in previous work.12,25,27 The cubic lattice parameter is around 11–13 nm for Ia3d and 18 nm for I23[^1], in line with a double- and triple-network structure, respectively. Remarkably, the Ia3d lattice is the largest for the smallest molecule 3/2 (L_mol = 4.4 nm; c_{cub} = 12.9 nm) and becomes smaller for the Ia3d phases of the larger compounds 3/12–3/16 with much longer chains (L_mol = 5.8–6.3 nm; c_{cub} = 11.3–11.5 nm).28 In line with this, in the Ia3d phase of 3/2 about 5 molecules are arranged side-by-side in the cross section of the networks, whereas for 3/12–3/16 there are only 3.2 molecules (Tables 2 and S7†). The twist between the molecules in the adjacent rafts with a height of 0.45 nm (Φ) can be calculated as ~7° in the long pitch Ia3d(L) phase of 3/2 and ~8° for the short pitch Ia3d(S) phase of compounds 3/12–3/16 (Table 2). The intermediate homologues 3/4–3/10 escape from forming the Ia3d phase by assuming the triple network I23[^1] structure instead. The twist in this network is almost constant 7.6–7.7° and between the angles in the Ia3d(L) and Ia3d(S) phases. The I23[^1] phase obviously allows for these twist angles a better fit of the helical pitch length with junction distances and inter-junction twist angles than in the competing Ia3d lattices.12,25 This phase sequence was also reported for the Cubbi phases of other rod-like molecules18,19,29 and it is observed in the contact regions between the achiral Ia3d(L) and Ia3d(S) phases of 3/2 and 3/12–3/16, respectively, where a concentration gradient develops, and in a certain concentration range the chiral conglomerate of the I23[^1] phase is induced (Fig. 21–k).29,30

In addition to the cubic phases, for compounds 3/2–3/12 a mirror symmetry broken isotropic liquid (Iso1[^1]S) phase occurs as a metastable (monotropic) phase on cooling the achiral Iso phase and replaces a part of the Cubbi phase range (Table 2). The Iso1[^1]S phase disappears at the transition from n = 12 to 14 right after the transition from I23[^1] to Ia3d(S) occurring from n = 10 to 12. Hence, it can be concluded that the local structure in the Iso1[^1]S phase is likely to be related to that of the chiral I23[^1] triple network (Fig. 1d). Remarkably, for 3/2–3/12 the transition Iso–Iso1[^1]S takes place in two separate steps, as indicated by the DSC trace of 3/12 (Fig. 3a). A very broad feature within the isotropic liquid range (with a maximum around 137 °C on cooling) indicates the transition from an ordinary

Fig. 2  (a and b) SAXS diffractions of the I23[^1] (3/4) and the Ia3d(S) (3/16) phases (for the synchrotron source and for numerical data, see Tables S4 and S5† the full indexations of the I23[^1] phase and the WAXS scans are shown in Fig. S7†); (c and d) reconstructed electron density maps of the I23[^1] and Ia3d(S) phase; the aromatic cores are located in the networks formed by the green iso-surfaces, the space between is filled by the alky1 chains; for more details, see the ESI, and Fig. S8† for the individual networks; (e–h) the textures of the (e) SmC phase of 3/10 at 230 °C; (f) the Cubbi/I23[^1] phase of 3/10 at 121 °C and the (g) Cubbi/Ia3d(S) phase of 3/12 at 120 °C, as observed between slightly uncrossed polarizers; (h) Colhax phase of 3/10 at 141 °C, the inset shows the texture with an additional x-plate, indicating that the phase is optically negative, i.e. the orientation of the aromatic cores is perpendicular or only slightly tilted to the column long axis; (i–k) show the induced chiral I23[^1] phase in the contact region between the achiral Ia3d(L) phase of 3/2 (top) and the Ia3d(S) phase of 3/16 (bottom); the orientation of the polarizers is shown by white arrows; the width of the POM images in (e–k) is 200 µm, and in (f), (i) and (k) the contrast is enhanced; for additional textures, see also Fig. S2–S6†
isotropic liquid (Iso) to a kind of cybotactic isotropic liquid with a fluctuating local network structure (Iso)\textsubscript{1},\textsuperscript{31} which is still achiral (Tables 2 and S1\textsuperscript{†}). With further decrease in the temperature the number of linkages between the clusters increases. Mirror symmetry breaking sets in at the next much sharper transition at 128 °C which we attribute to the transition from the cybotactic to a percolated liquid, after crossing a certain critical density of connectivity between the cybotactic clusters,\textsuperscript{31,32} thus leading to a long range transmission of the helical twist, and hence, chirality (Iso)\textsubscript{1}.

In the Iso)\textsubscript{1} range the connections are transient and at the next transition at 122 °C they become permanent with formation of the Ia\textit{3}d phase (or I2\textit{3}[\textgamma] for the shorter homologues, Fig. 3e). In the temperature range of the isotropic liquid phases there is an almost continuous increase in the line width of the small angle XRD scattering (Fig. S1\textsuperscript{†}) and there is also no obvious discontiuity in the viscosity, as indicated by optical microscopy, where all three isotropic liquid phases flow under the influence of gravity. Only at the transition to the cubic phase the material suddenly becomes viscoelastic. The enthalpy of the Iso\textsubscript{1} transition increases with the growing chain length with a distinct jump from \( n = 4 \) to 6 (Table 2), in line with improved rod-chain segregation, supporting the cluster formation.

For the longer homologues 3/14 and 3/16 the Iso)\textsubscript{1} phase is replaced by a Col\textsubscript{hex} phase (Fig. 3b). The growing chain length obviously disfavours the branching, thus leading to predominately linear aggregates which do not form networks, but assume long range order and transform to the achiral Col\textsubscript{hex} phase.

Another interesting feature of compounds 3/6–3/10 is that on further cooling from the Cub\textsubscript{b}/I2\textit{3}[\textgamma] phase a transition to an optically isotropic crystalline mesophase is observed, in which, the conglomerate texture is retained (Col\textsubscript{iso}[\textgamma] phase, see Fig. 4c–f). Because this transition is associated with a change in the XRD pattern and a sharp transition with a significant transition enthalpy of 17–18 kJ mol\textsuperscript{−1} (Table 2 and Fig. 4g), it cannot be a glass transition. The XRD pattern of the Col\textsubscript{iso}[\textgamma] phase is characterized by a relatively broad small angle scattering, with a maximum at \( d = 5.38 \) nm for 3/6 (Fig. 4h) and 5.45 nm for 3/10, approximately corresponding to the lengths of the respective intercalated anti-parallel molecular pairs (5.5 nm, see Fig. S15\textsuperscript{†}). In the wide angle range, there are three broad scatterings with very low intensity (Fig. 4i, S12 and S13\textsuperscript{†}), which can tentatively be attributed to the mean alkyl chain distance (0.44 nm) and the edge-to-edge and face-to-face packing distances of the aromatics (0.56/0.37 nm). This diffraction pattern is similar to those typically recorded for the symmetry broken soft crystalline mesophases of bent-core mesogens, helical nanofilaments (HNFs),\textsuperscript{16,33} helical nano-crystallites (HNCs)\textsuperscript{34} and related helical phases,\textsuperscript{35} which in some respect can be considered as solvent free gels.\textsuperscript{16,36,37} A transition from the LC Cub\textsubscript{b} phase to a soft crystalline network structure, where the poly-aromatic rods and parts of the aliphatic chains assume a crystalline packing, appears likely. Similar to the bent-core mesogens, the directed packing of the twisted and bent 4,4'-diphenylbenzil units (Fig. S15\textsuperscript{†}) is likely to contribute to the development of the helical packing modes in the liquid, LC and especially in the soft crystalline mesophases of compounds 3/n.

For compounds 3/12 and 3/14, forming the achiral Ia\textit{3}d(\textsuperscript{5}) cubic phase instead of I2\textit{3}[\textgamma], the isotropic crystalline phase appears to be achiral (Col\textsubscript{iso}, see Fig. S6\textsuperscript{†}), though the XRD pattern (Fig. S14\textsuperscript{†}) is almost the same as for 3/6 and 3/10. Whether this phase is intrinsically achiral, or the symmetry breaking at the transition from the achiral Ia\textit{3}d(\textsuperscript{5}) to a Col\textsubscript{iso}[\textgamma] phase can only develop locally, as the OA domains are too small to be observable by optical investigation, cannot be decided at present.

In summary, first benzil derivatives forming a series of LC phases, ranging from lamellar via two types of Cub\textsubscript{b} phases to columnar, have been obtained. Mirror symmetry breaking is observed in three of the soft matter phases, the isotropic liquid, the liquid crystalline Cub\textsubscript{b}/I2\textit{3}[\textgamma] phase and in the soft crystalline phase (Col\textsubscript{iso}[\textgamma]). This work contributes to the understanding of the development of mirror symmetry breaking in isotropic liquids as a consequence of network formation and increasing network-connectivity (Fig. 3e). In addition, these molecules with a bend around the central O=C=C=O bond appear to provide the still missing link between the mirror symmetry broken modes of soft self-assembly observed in polycatenars and in bent-core LC systems,\textsuperscript{5,15,16,18,19,19} bent mesogenic dimers,\textsuperscript{16,15,16} and oligomers.\textsuperscript{39} Moreover, as the Iso)\textsubscript{1} and Cub\textsubscript{b}/I2\textit{3}[\textgamma] phases...
can provide giant chirality amplification,\(^{30,31}\) the homogeneously chiral crystalline networks, once developed from these spontaneously mirror symmetry broken soft matter states, could be of interest as chiral templates for chiral recognition and separation, and the emerging field of dynamicenantioselective catalysis.\(^{17,39,40}\)

### Conflicts of interest

There are no conflicts to declare.

### Acknowledgements

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### References


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**Fig. 4** The mirror symmetry broken mesophases of compound 3/6. (a–f) Conglomerates of optically active domains: (a and b) in the liquid $\text{Iso}^{\text{H}}$ phase at 120 °C; (c and d) in the LC $\text{Cub}_{23}^\parallel$ phase at 110 °C and (e and f) in the soft crystalline $\text{Cr}_{\text{iso}}^{\text{a}}$ phase at 60 °C, as observed on cooling between slightly uncrossed polarizers rotated by 5°; (a, c and e) in the counterclockwise and (b, d and f) in the clockwise direction (contrast enhanced). (g) DSC heating and cooling traces (10 K min$^{-1}$); (h) small angle and (i) wide angle XRD pattern in the $\text{Cr}_{\text{iso}}^{\text{a}}$ phase at 50 °C (see also Fig. S12† for the complete diffraction pattern and Fig. S13† for the diffraction pattern of 3/10).
28 This is in contrast to a previous series of rod-like molecules with only two terminal chains, forming a similar sequence la3d–I23[1]–la3d with the growing chain length, but with an increasing d-value. 18a,b,19