

Supporting Information

Star-Shaped Triphenylene Discotic Liquid Crystalline Oligomers and their Hydrogen-Bonded Supramolecular Complexes with Simple Acids

Ke-Qing Zhao, Xiao-Yan Bai, Bo Xiao, Yue Gao, Ping Hu, Bi-Qin Wang, Qing-Dao Zeng, Chen Wang, Benoît Heinrich, Bertrand Donnio

Table of Content

1 ^1H NMR spectra: Figures S1-S20 (1a-d ; 2a-d ; 3 , 3a ; 4a-d ; 5a-f)	pages	2-11
2 ^{13}C NMR spectra: Figures S21-S24 (4a-d)	pages	12-13
3 IR: Figures S25-S33 (4d , hexanoic acid, 5a , 5b ; benzoic acid, 5c , 5d ; 4-hexoxybenzoic acid, 5e)	pages	14-18
4 DSC: Figures S34-S39 (1b , 1d ; 2b , 2d ; 4c , 4d)	page	19
5 SAXS: Figures S40-S42 (5a-f)	page	20
6 SAXS in isotropic liquid: Figure S43 (4a,c,d)	page	21

¹H NMR

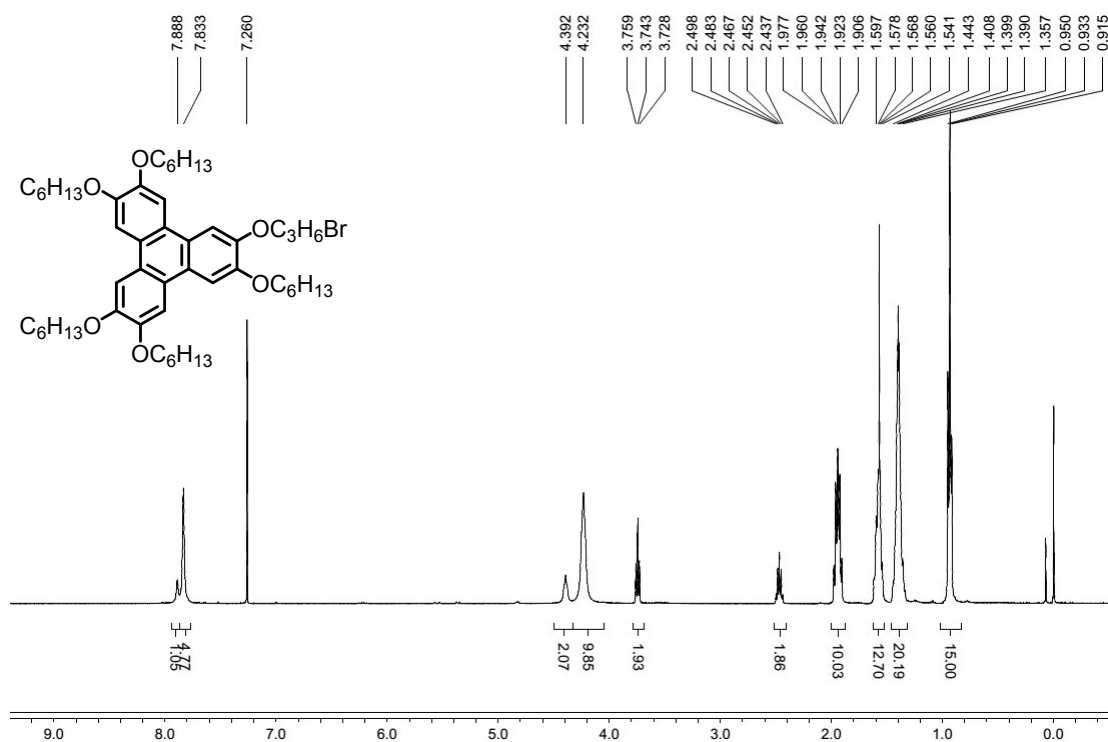


Figure S1: ¹H NMR (CDCl₃, 400 MHz) spectrum of **1a**

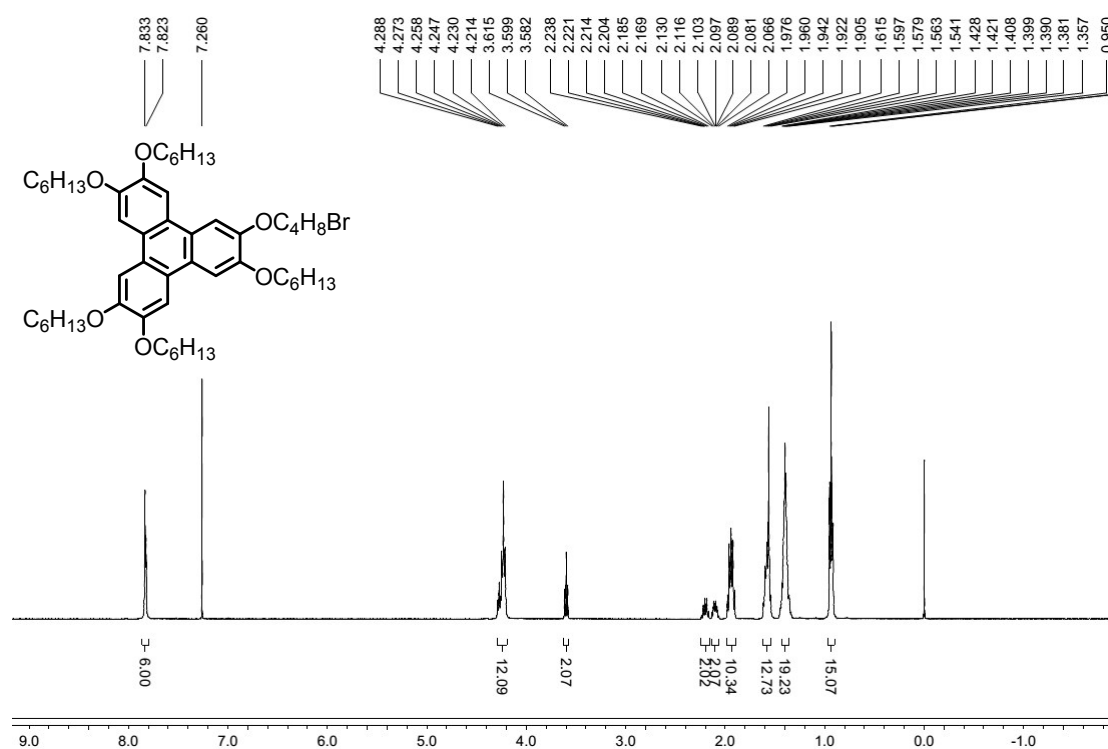


Figure S2: ¹H NMR (CDCl₃, 400 MHz) spectrum of **1b**

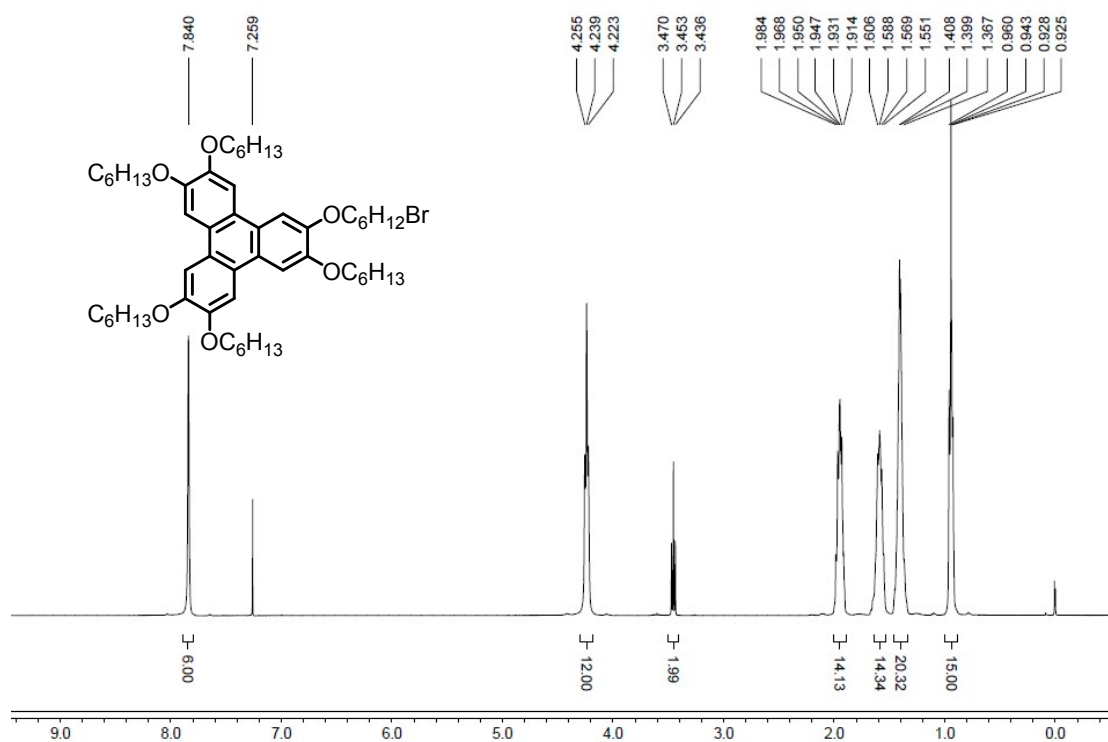


Figure S3: ^1H NMR (CDCl_3 , 400 MHz) spectrum of **1c**

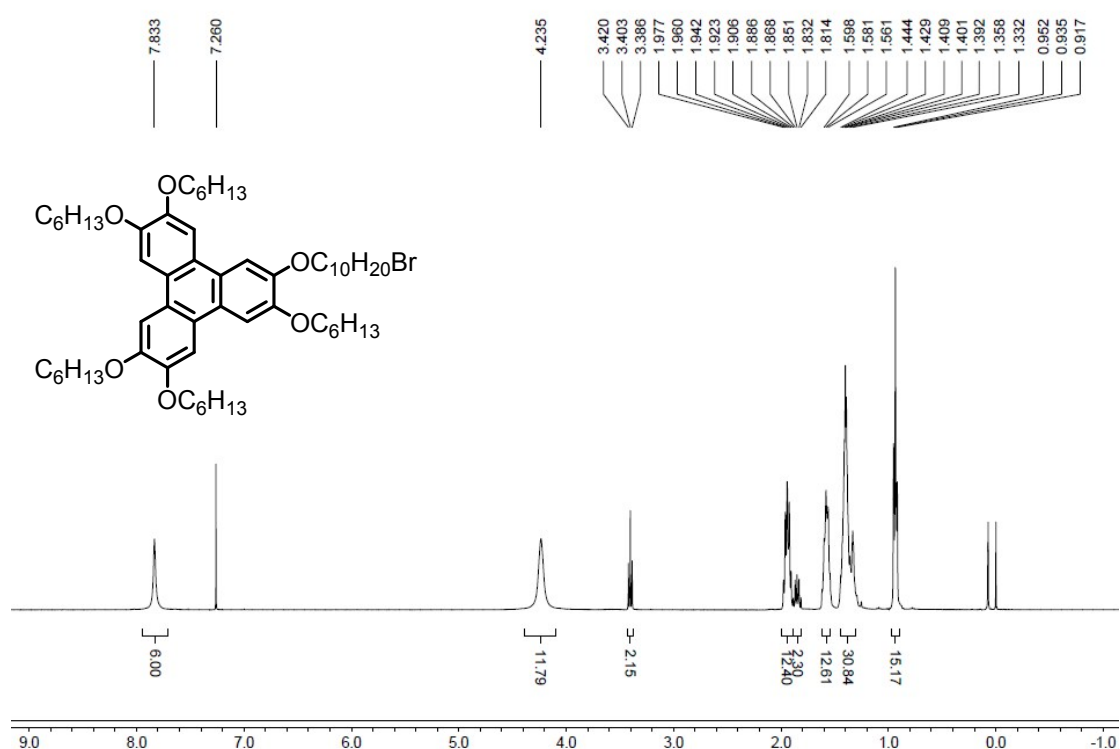


Figure S4: ^1H NMR (CDCl_3 , 400 MHz) spectrum of **1d**

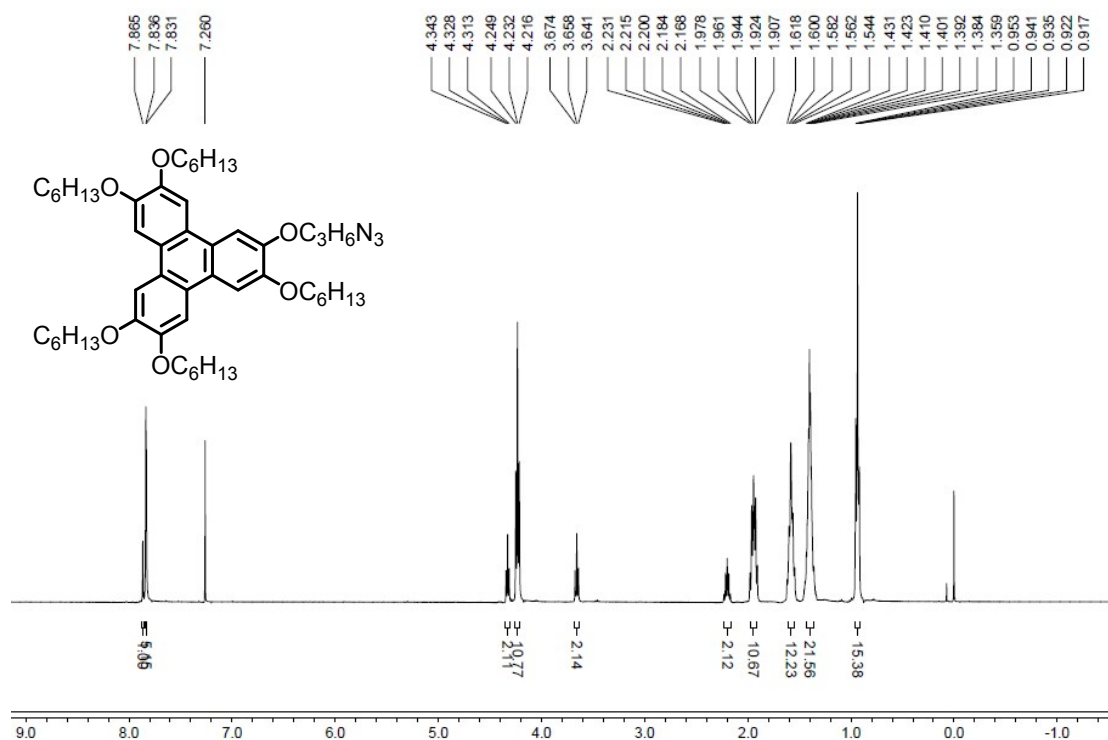


Figure S5: ^1H NMR (CDCl₃, 400 MHz) spectrum of **2a**

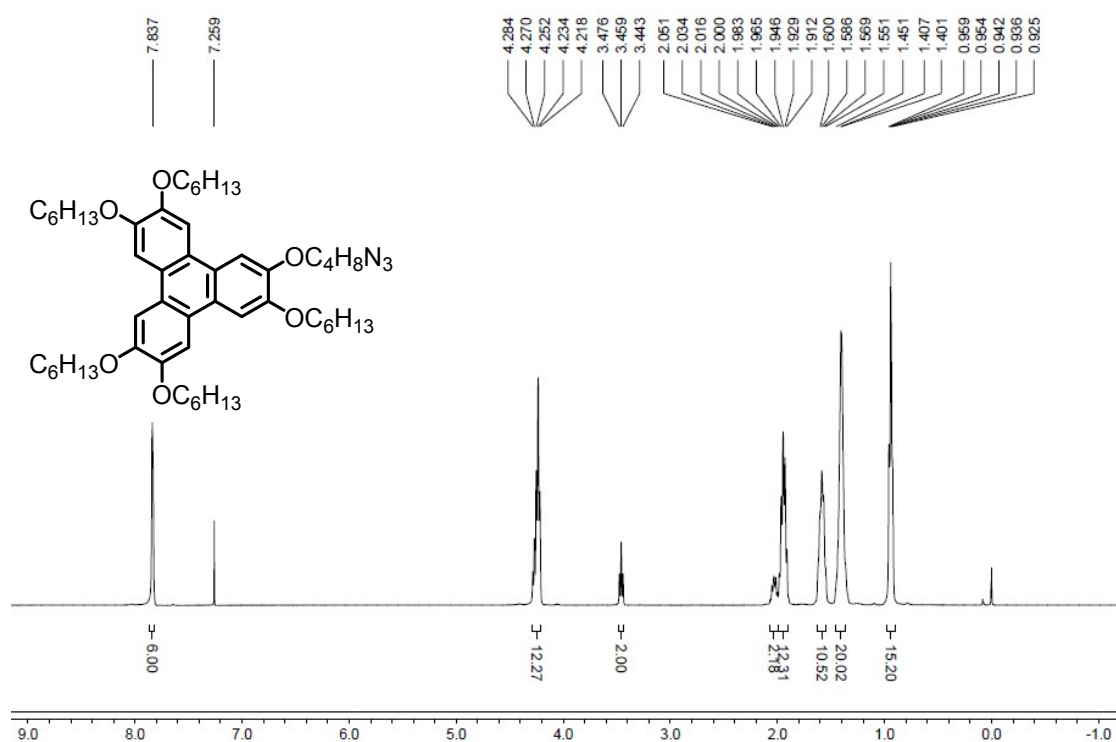


Figure S6: ^1H NMR (CDCl₃, 400 MHz) spectrum of **2b**

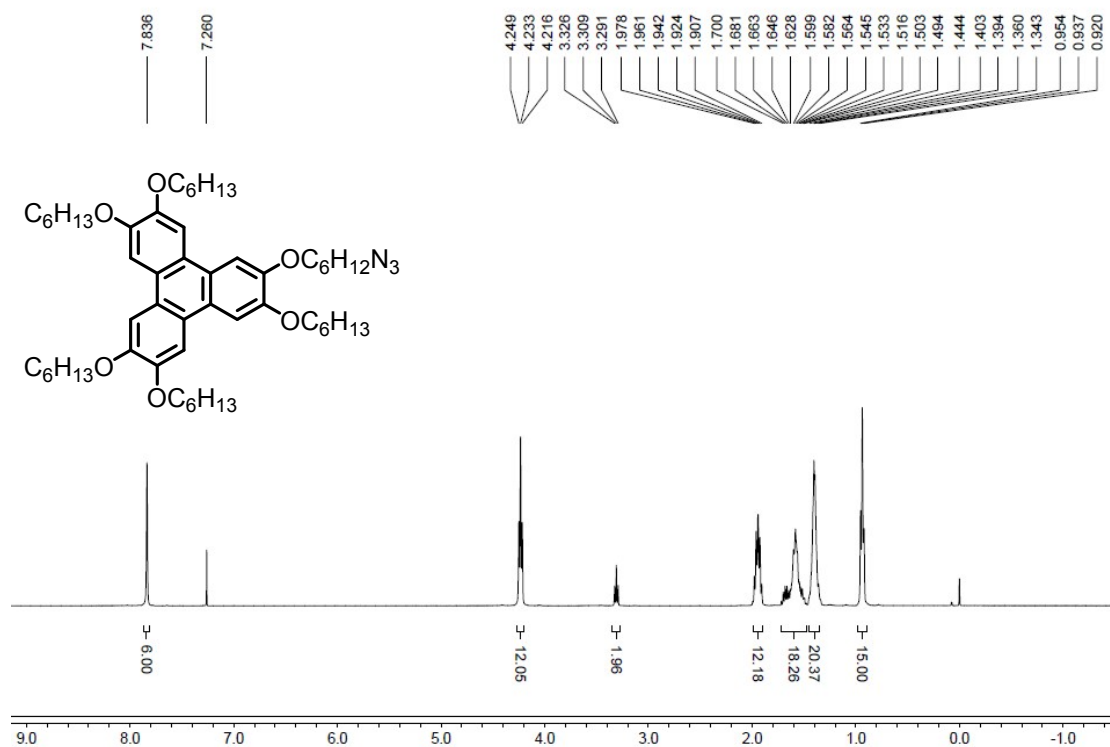


Figure S7: ^1H NMR (CDCl₃, 400 MHz) spectrum of **2c**

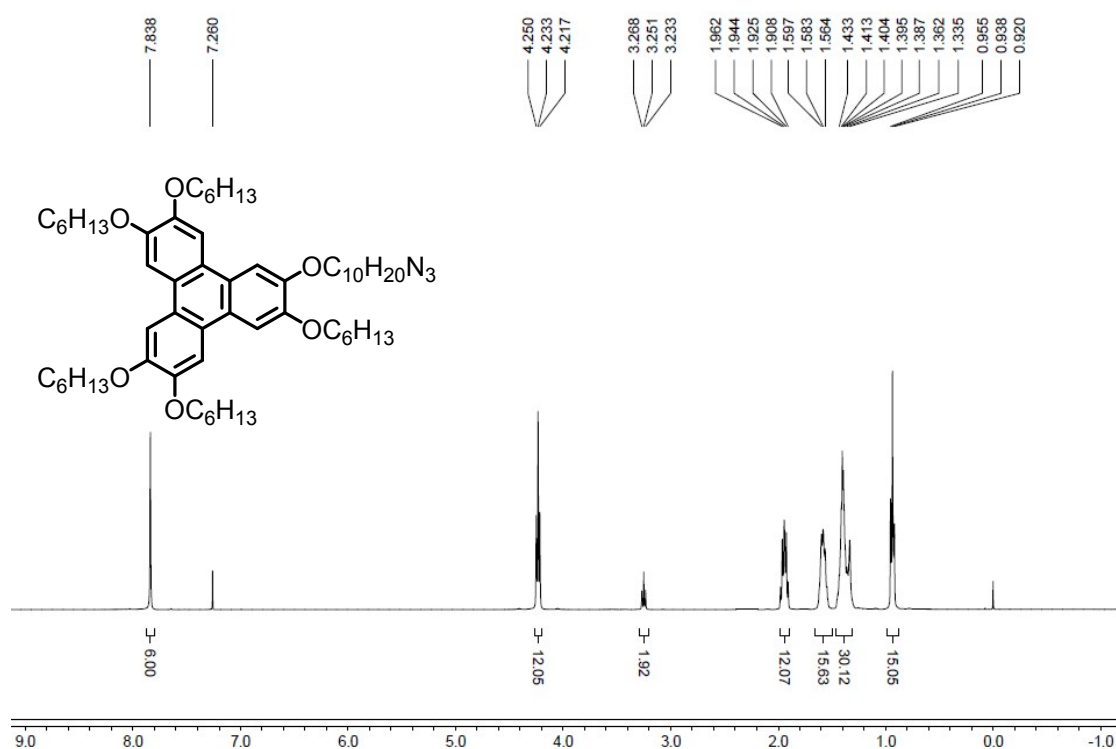


Figure S8: ^1H NMR (CDCl₃, 400 MHz) spectrum of **2d**

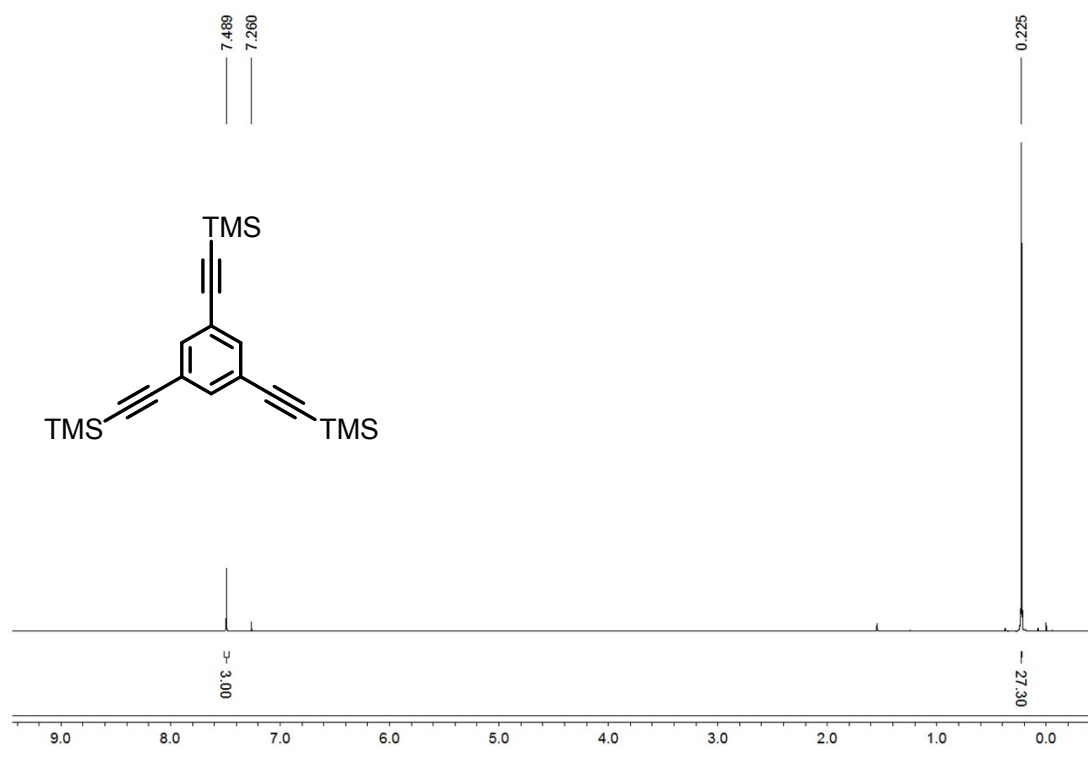


Figure S9: ^1H NMR (CDCl₃, 400 MHz) spectrum of **3**

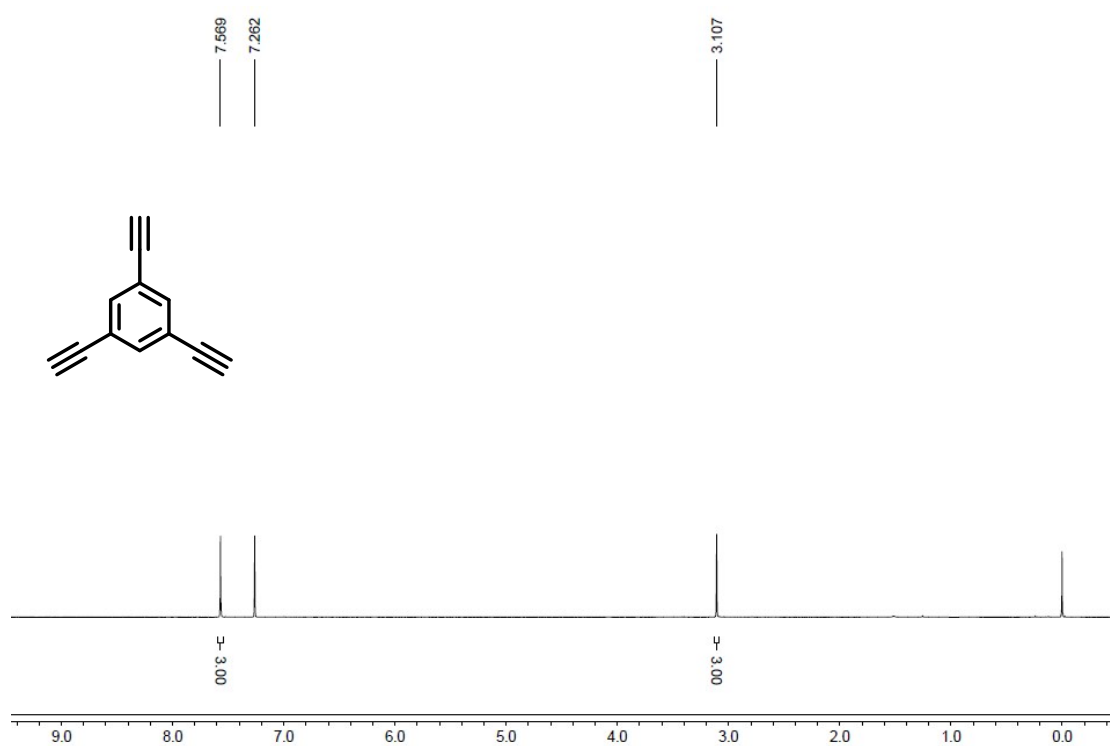


Figure S10: ^1H NMR (CDCl₃, 400 MHz) spectrum of **3a**

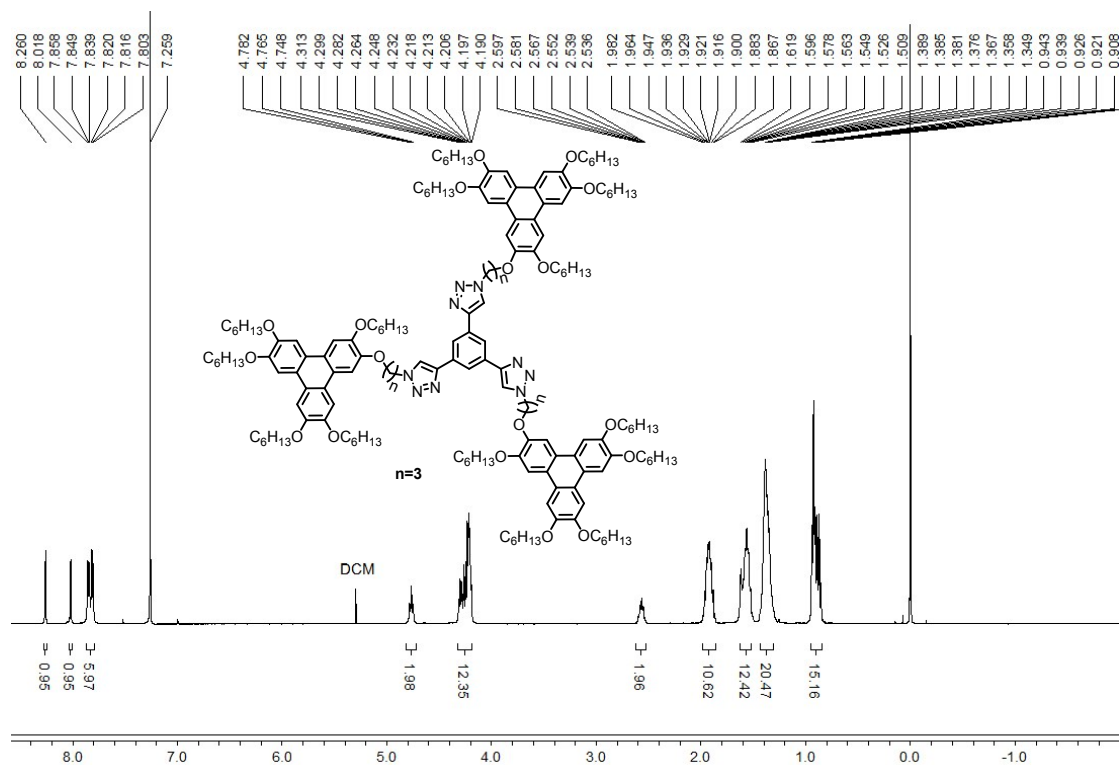


Figure S11: ^1H NMR (CDCl_3 , 400 MHz) spectrum of **4a**, $n = 3$

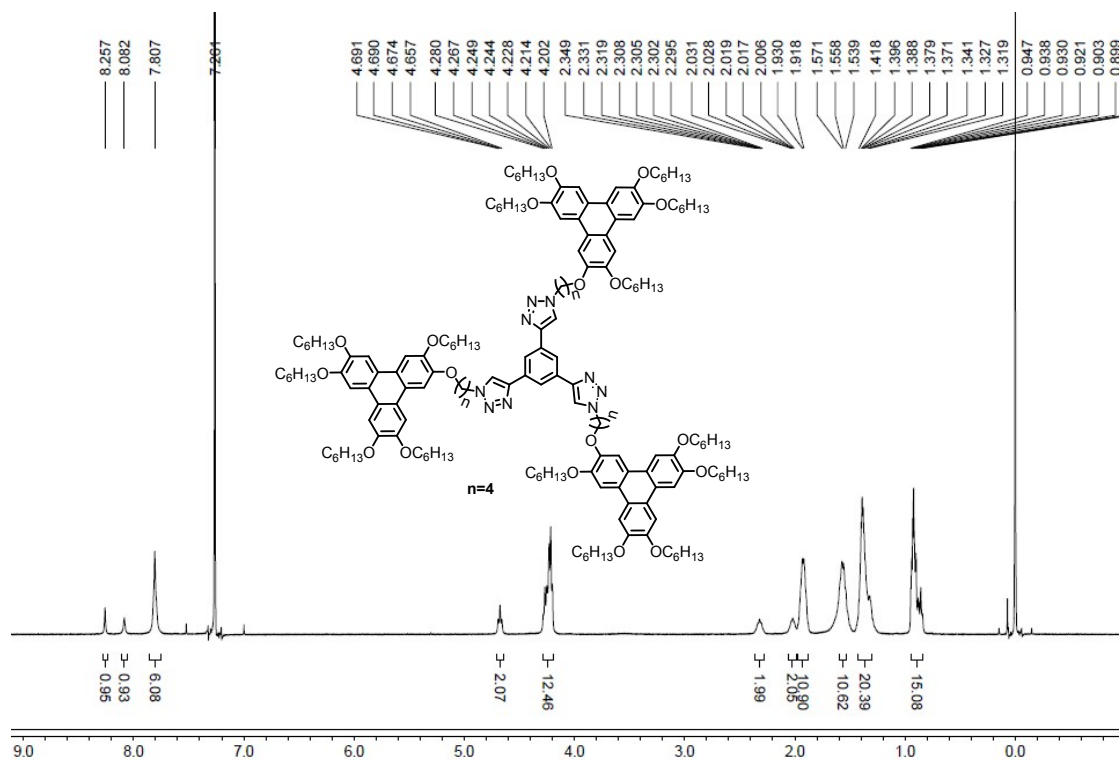


Figure S12: ^1H NMR (CDCl_3 , 400 MHz) spectrum of **4b**, $n = 4$

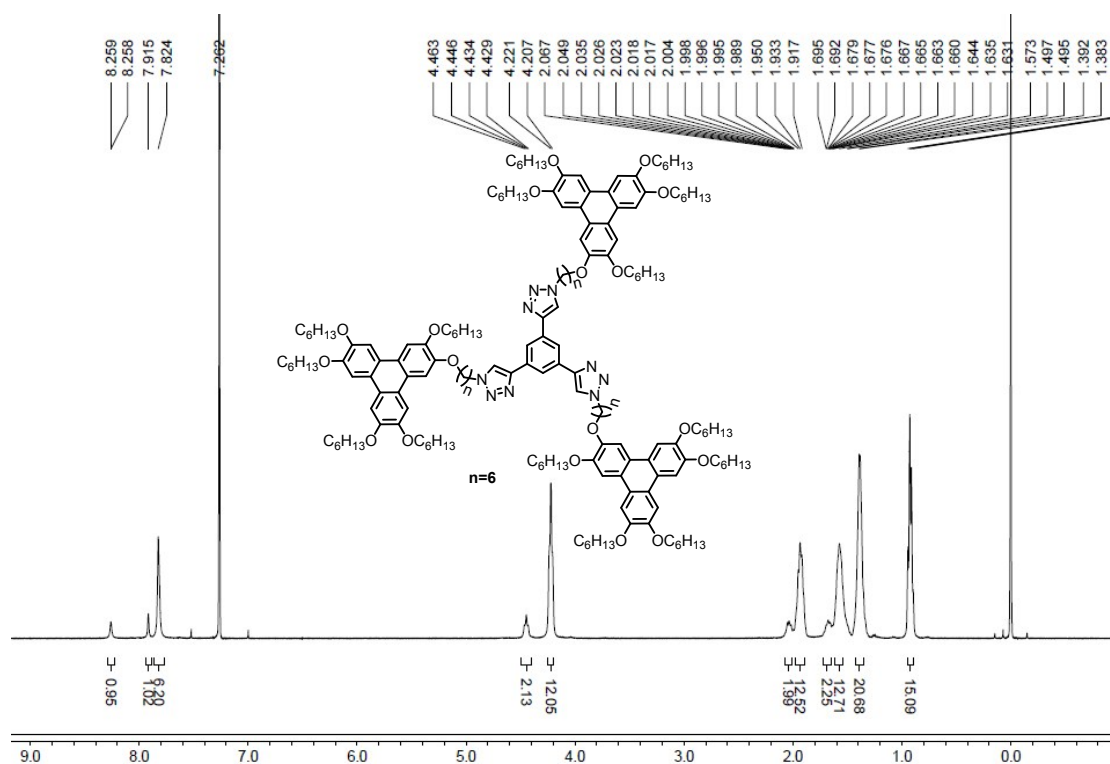


Figure S13: ¹H NMR (CDCl₃, 400 MHz) spectrum of **4c**, n = 6

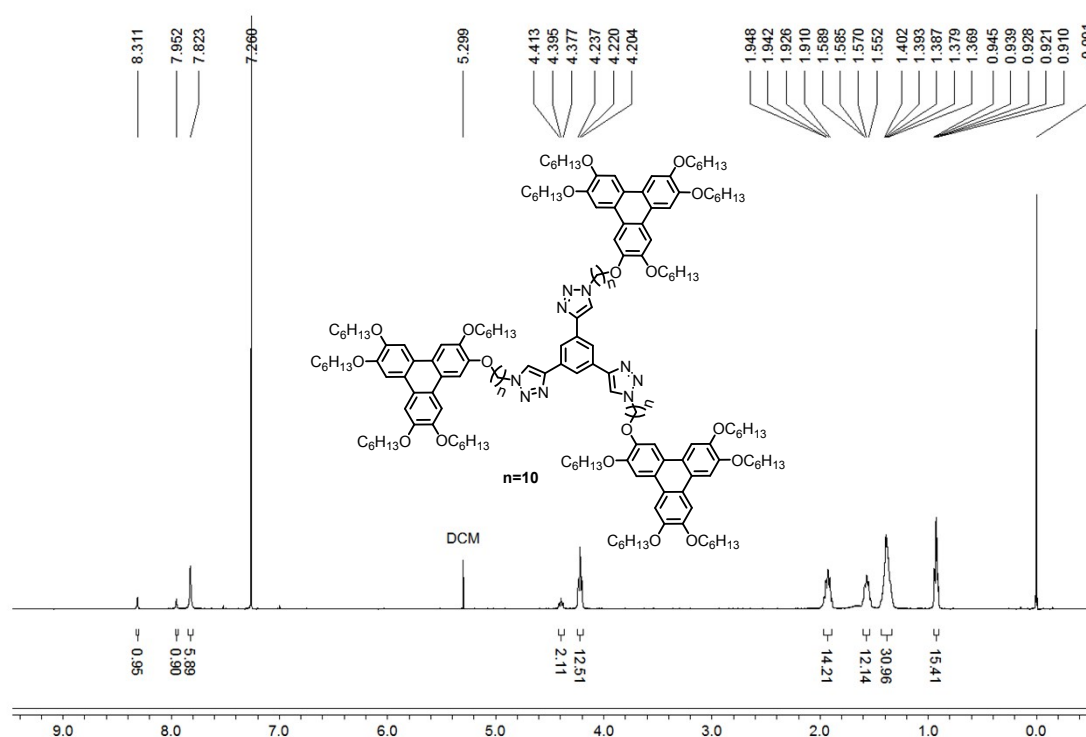


Figure S14: ¹H NMR (CDCl₃, 400 MHz) spectrum of **4d**, n = 10

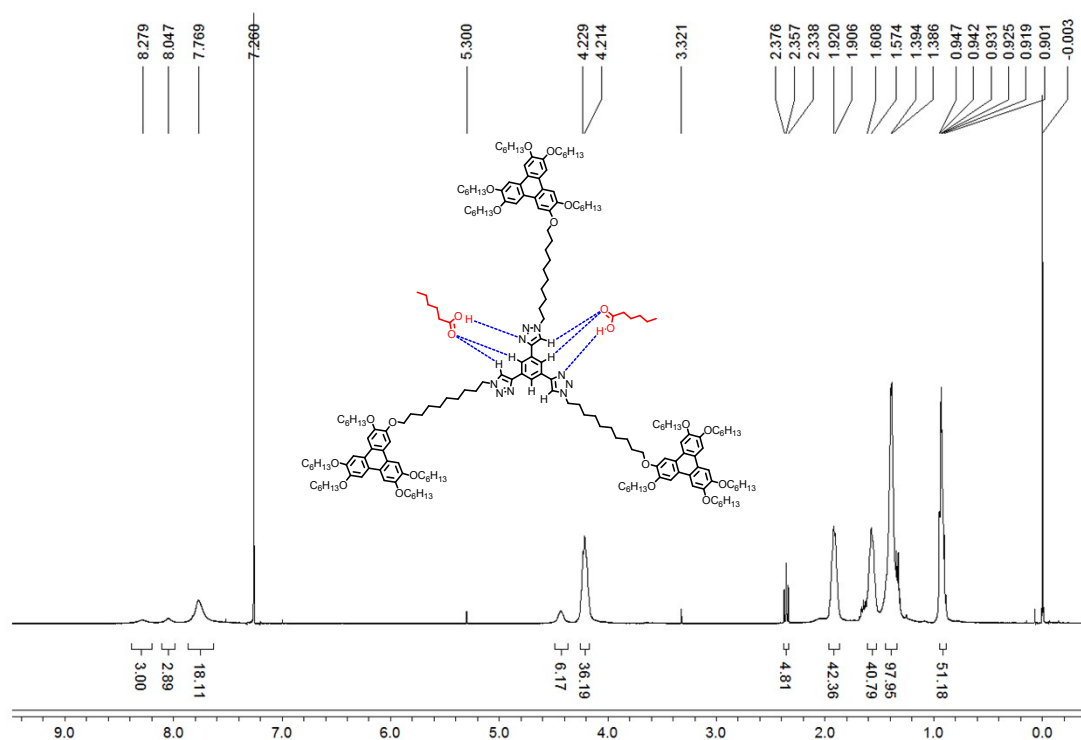


Figure S15: ^1H NMR (CDCl_3 , 400 MHz) spectrum of **5a**

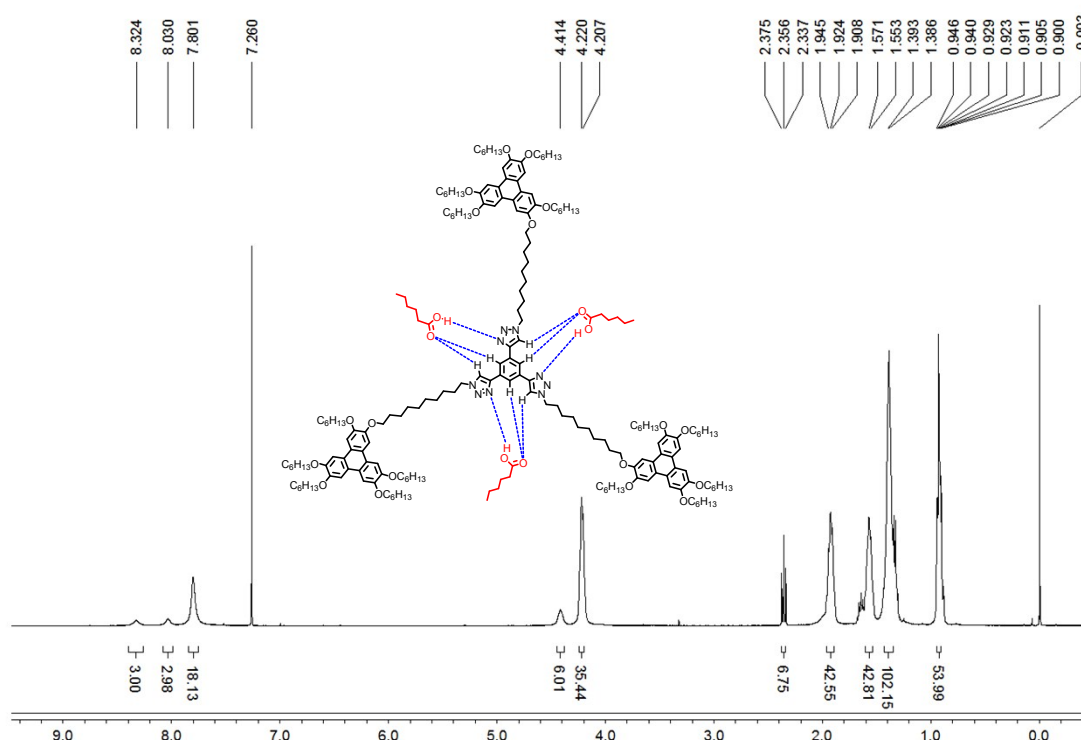


Figure S16: ^1H NMR (CDCl_3 , 400 MHz) spectrum of **5b**

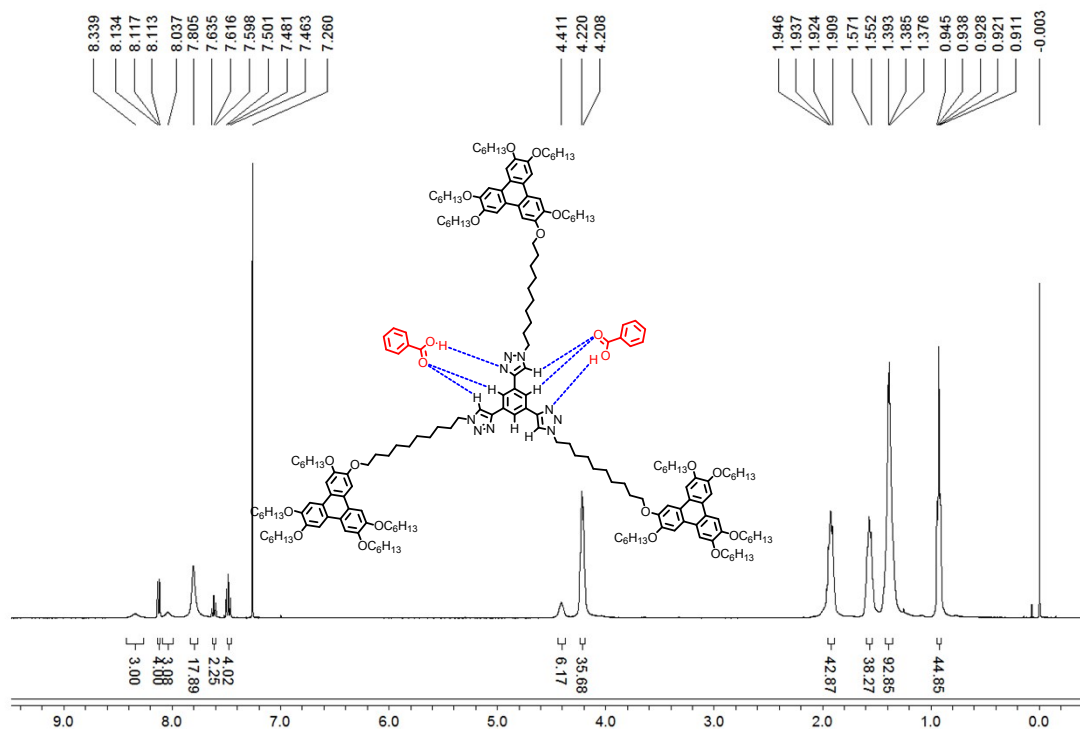


Figure S17: ^1H NMR (CDCl_3 , 400 MHz) spectrum of **5c**

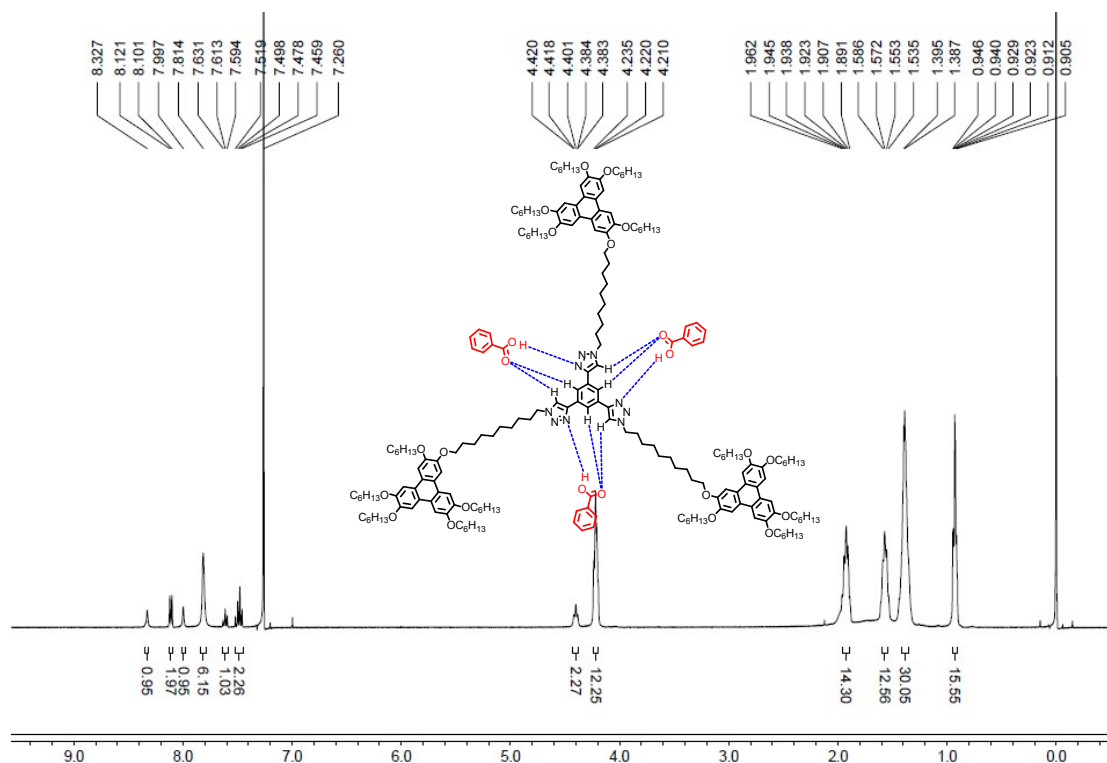


Figure S18: ^1H NMR (CDCl_3 , 400 MHz) spectrum of **5d**

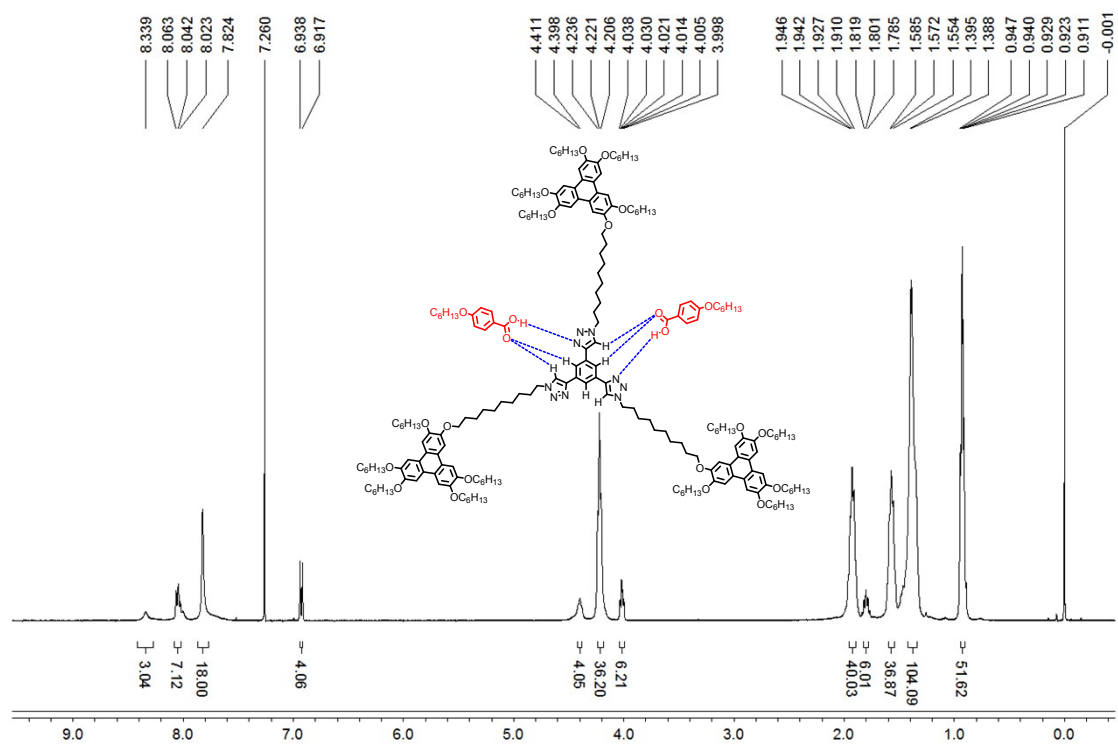


Figure S19: ¹H NMR (CDCl₃, 400 MHz) spectrum of **5e**

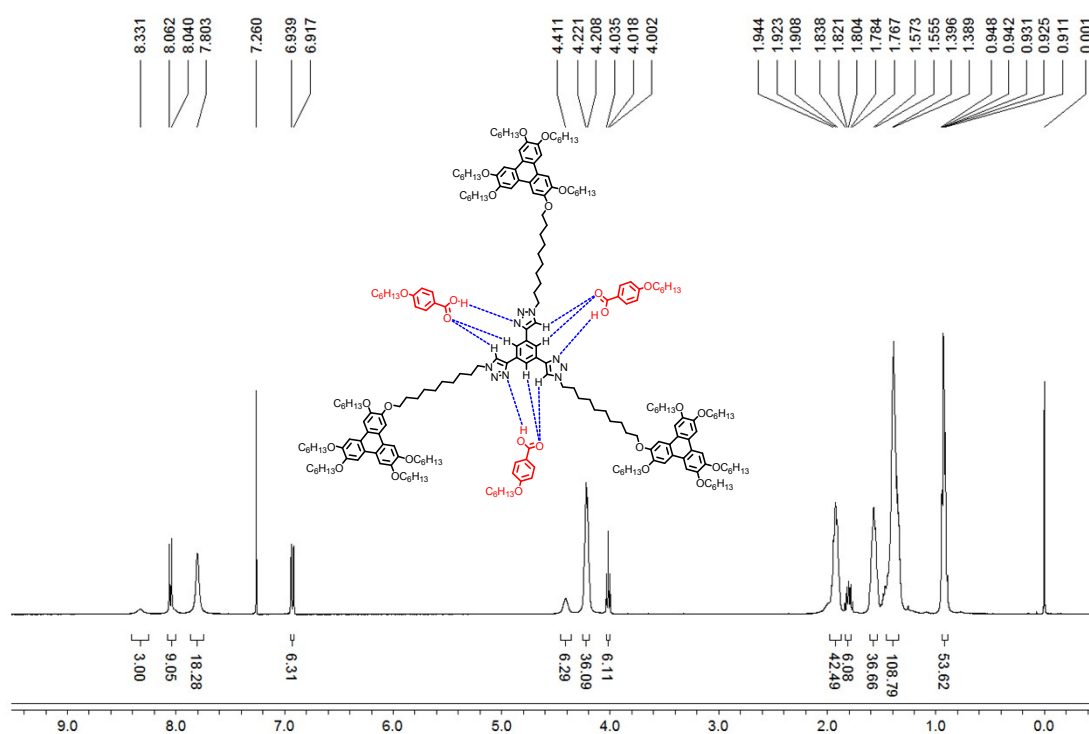


Figure S20: ¹H NMR (CDCl₃, 400 MHz) spectrum of **5f**

2. ^{13}C NMR

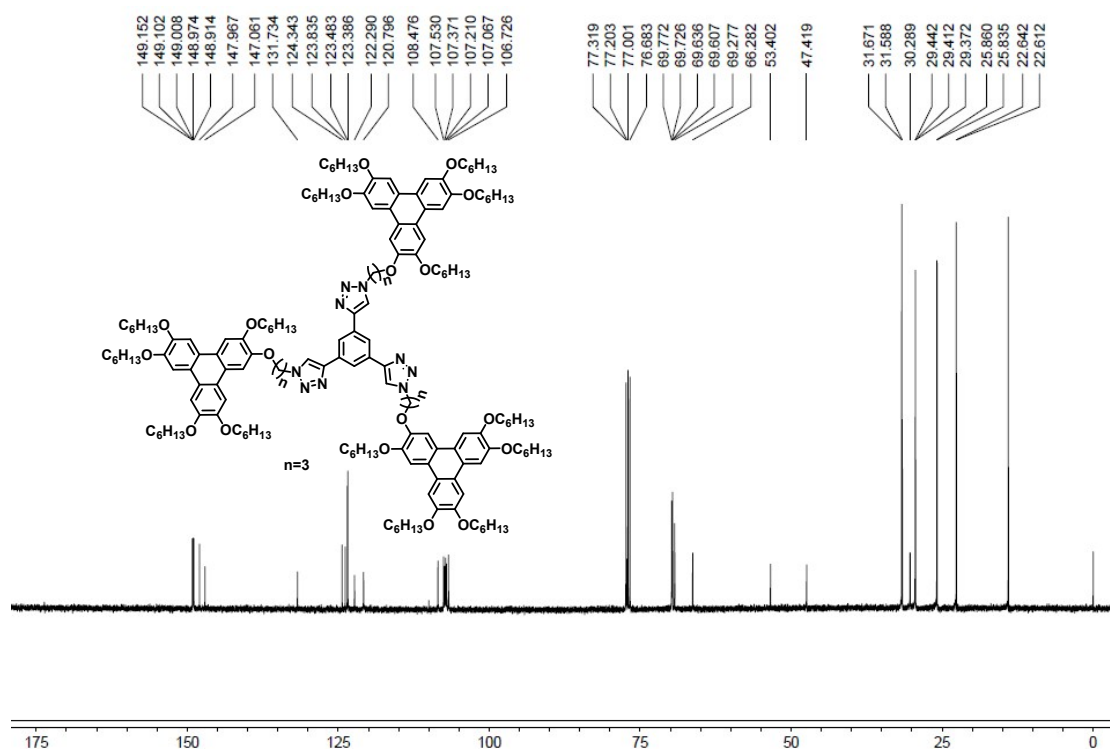


Figure S21: ^{13}C NMR (CDCl_3 , 100 MHz) spectrum of **4a**, $n=3$

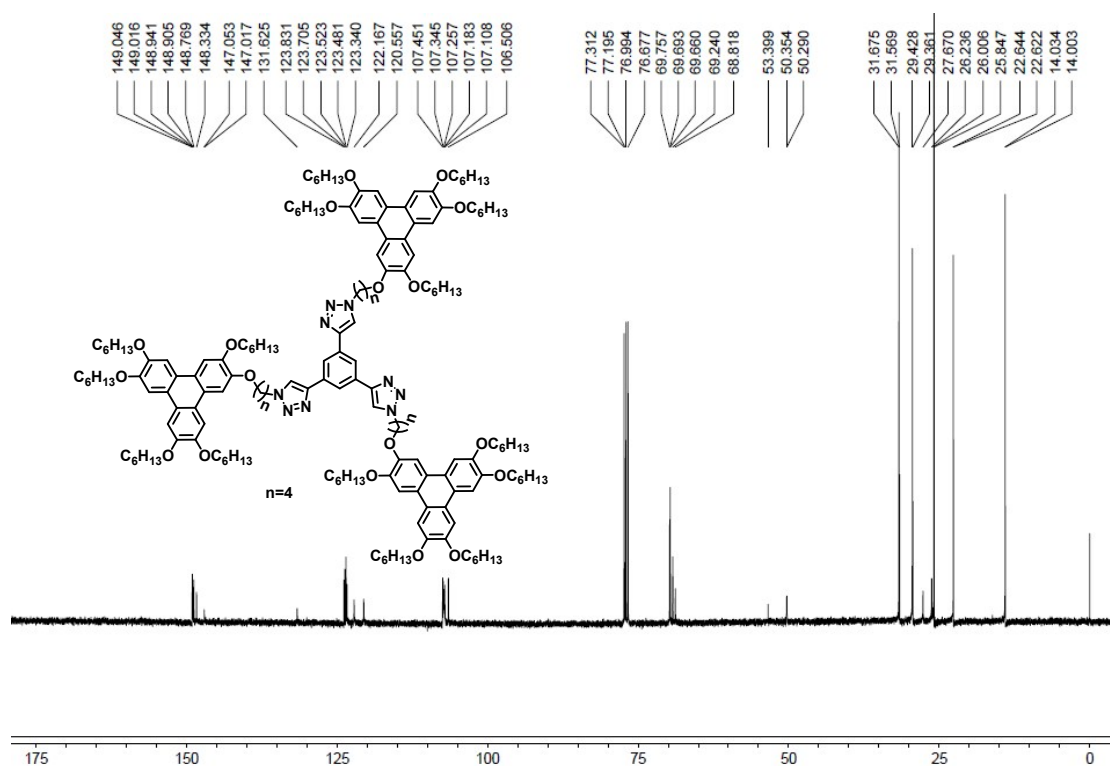


Figure S22: ^{13}C NMR (CDCl_3 , 100 MHz) spectrum of **4b**, $n=4$

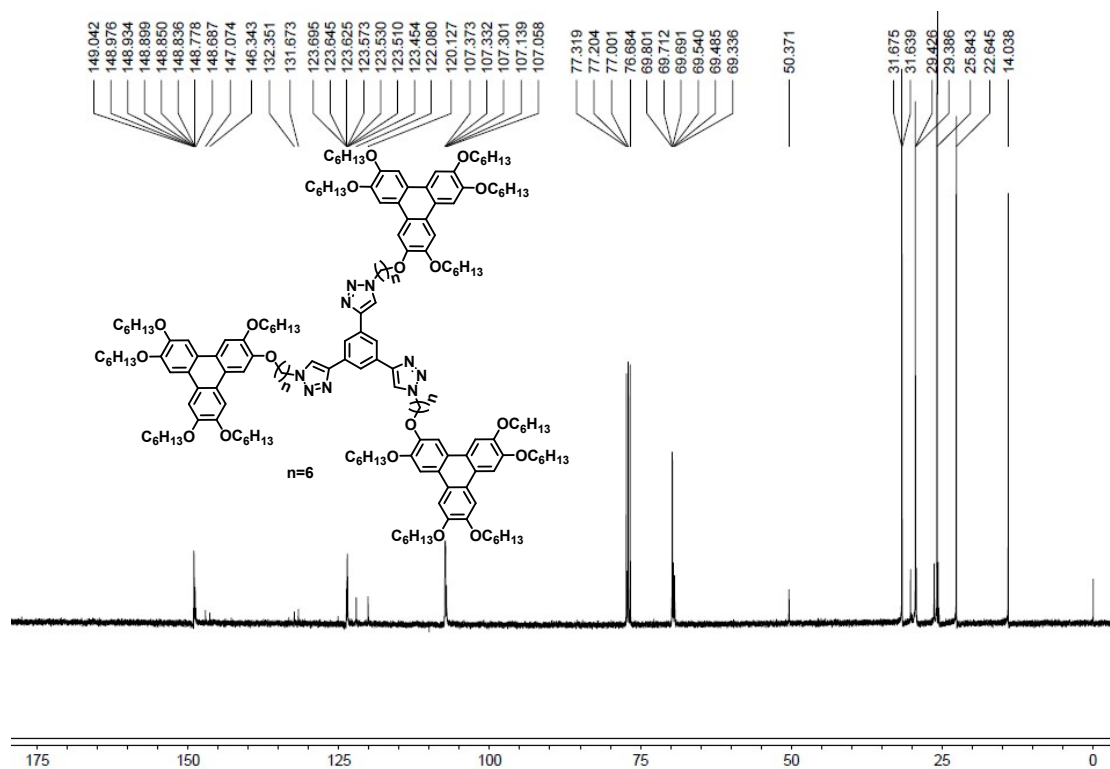


Figure S23: ^{13}C NMR (CDCl_3 , 100 MHz) spectrum of **4c**, $n=6$

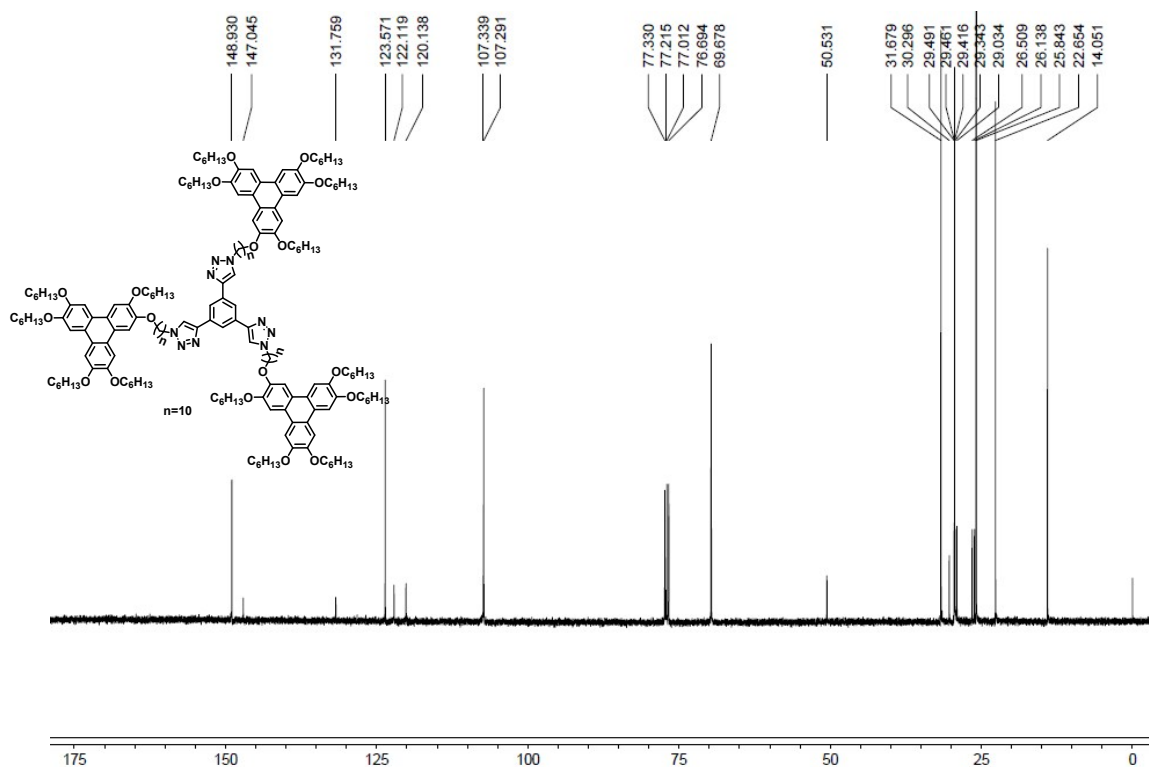


Figure S24: ^{13}C NMR (CDCl_3 , 100 MHz) spectrum of **4d**, $n=10$

3. IR

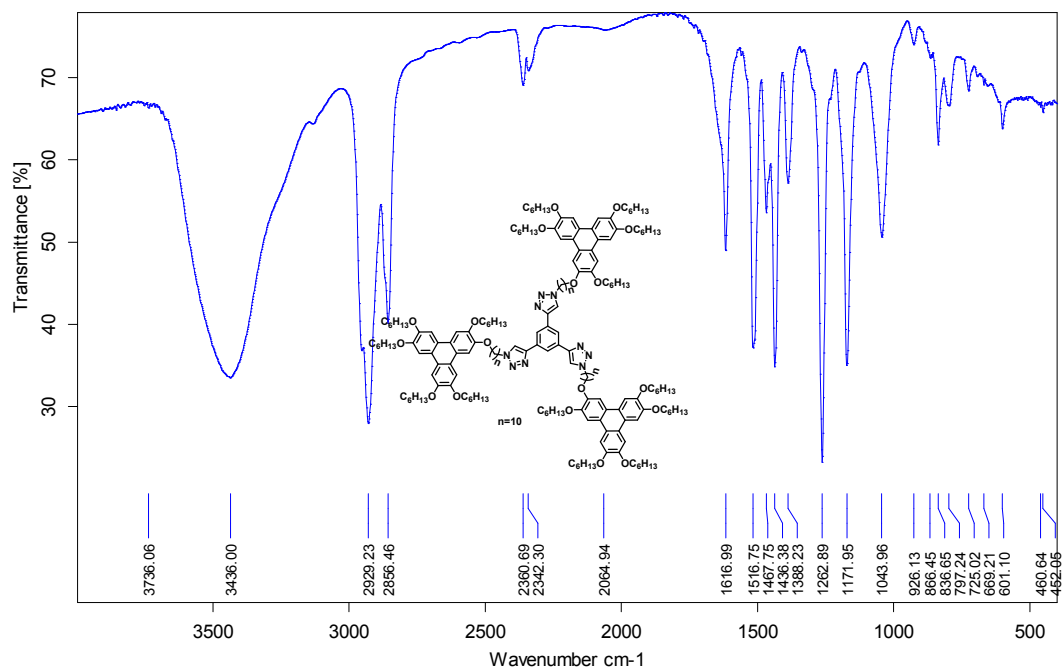


Figure S25: IR spectrum of **4d**, $n = 10$

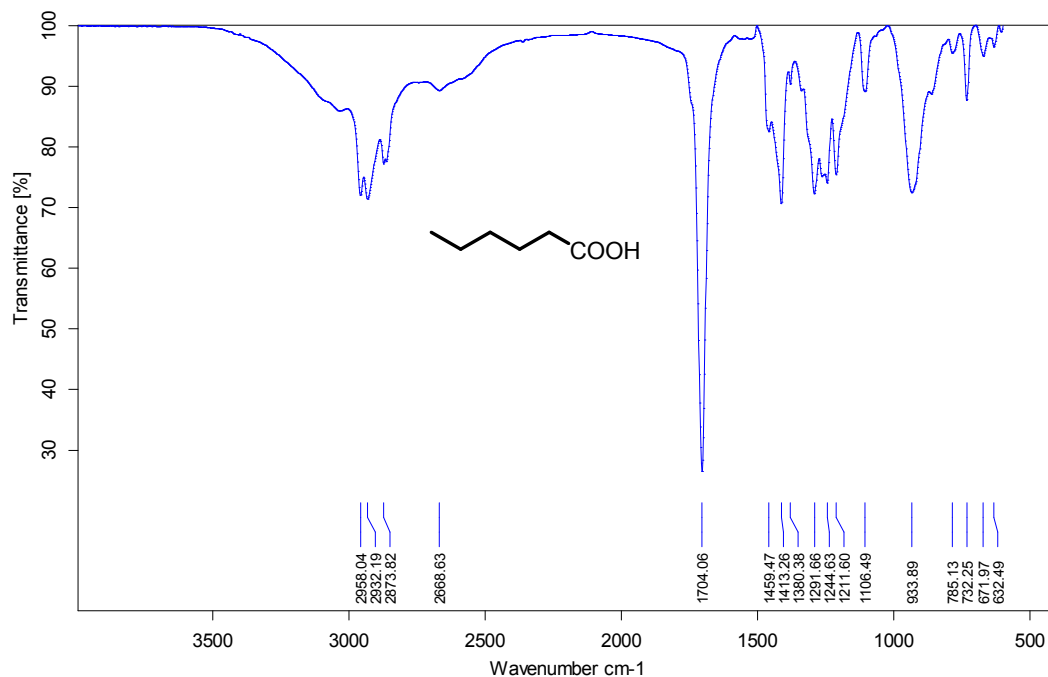


Figure S26: IR spectrum of hexanoic acid

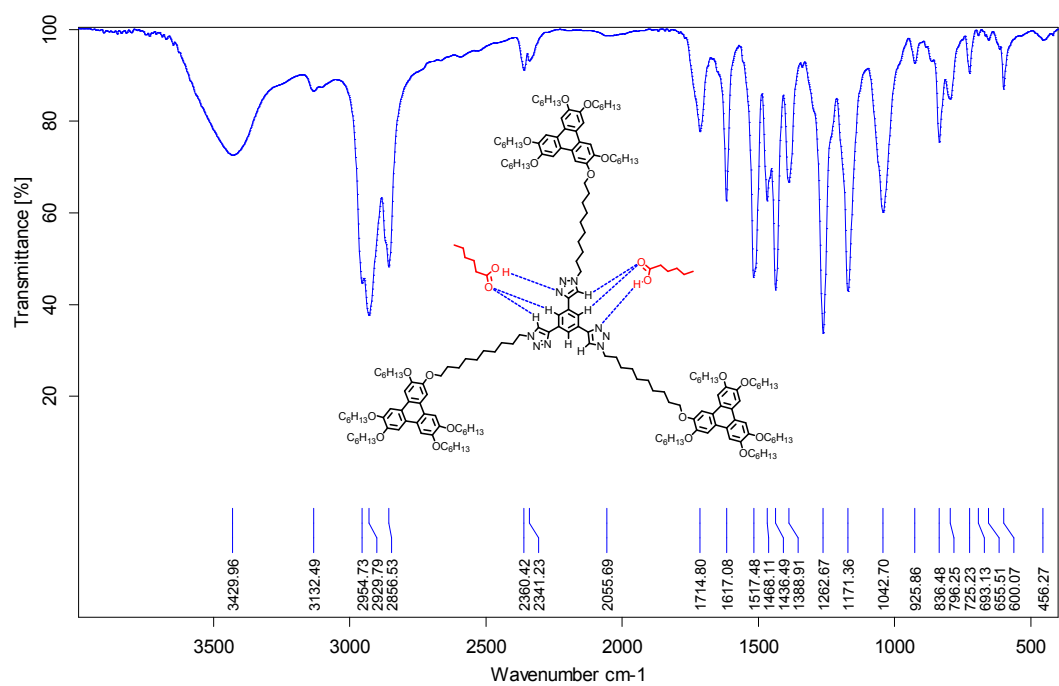


Figure S27: IR spectrum of **5a**

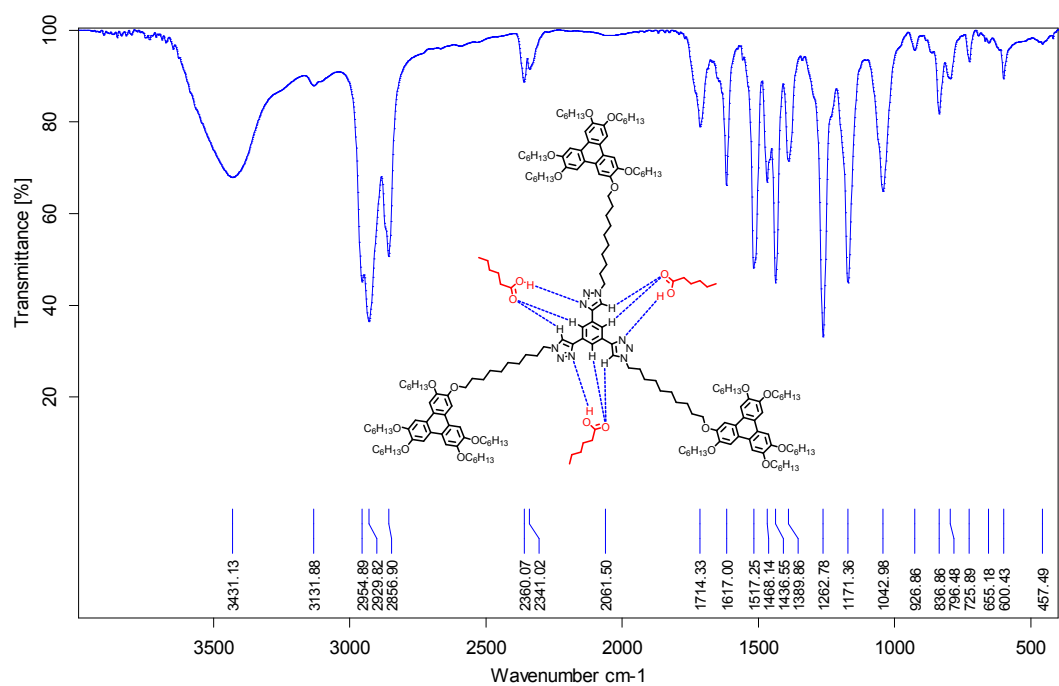


Figure S28: IR spectrum of **5b**

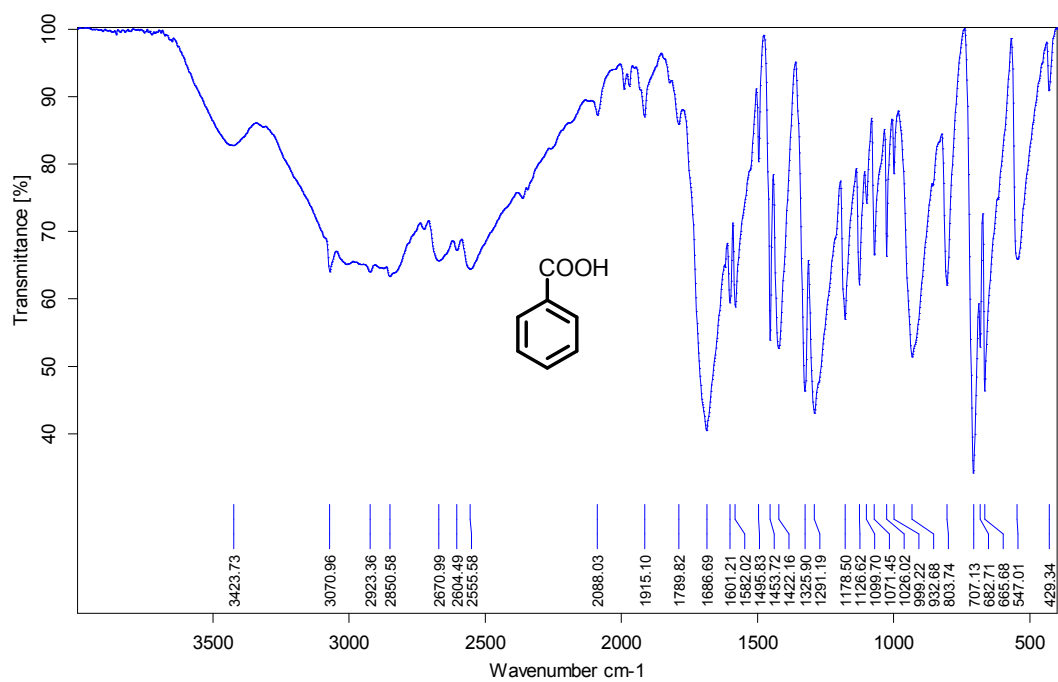


Figure S29: IR spectrum of benzoic acid

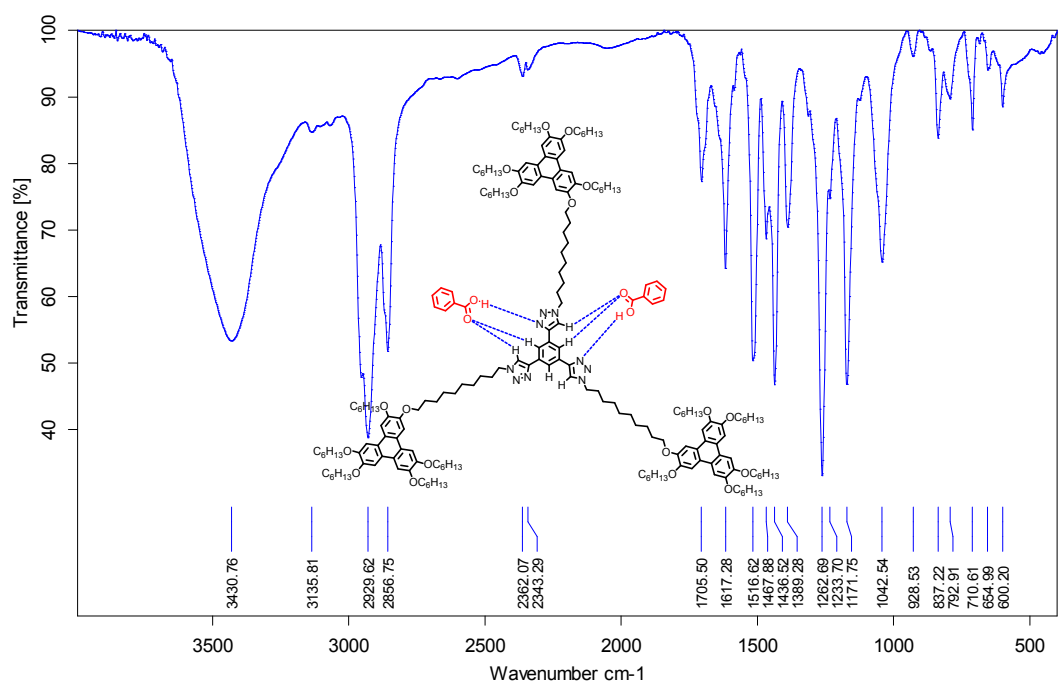


Figure S30: IR spectrum of **5c**

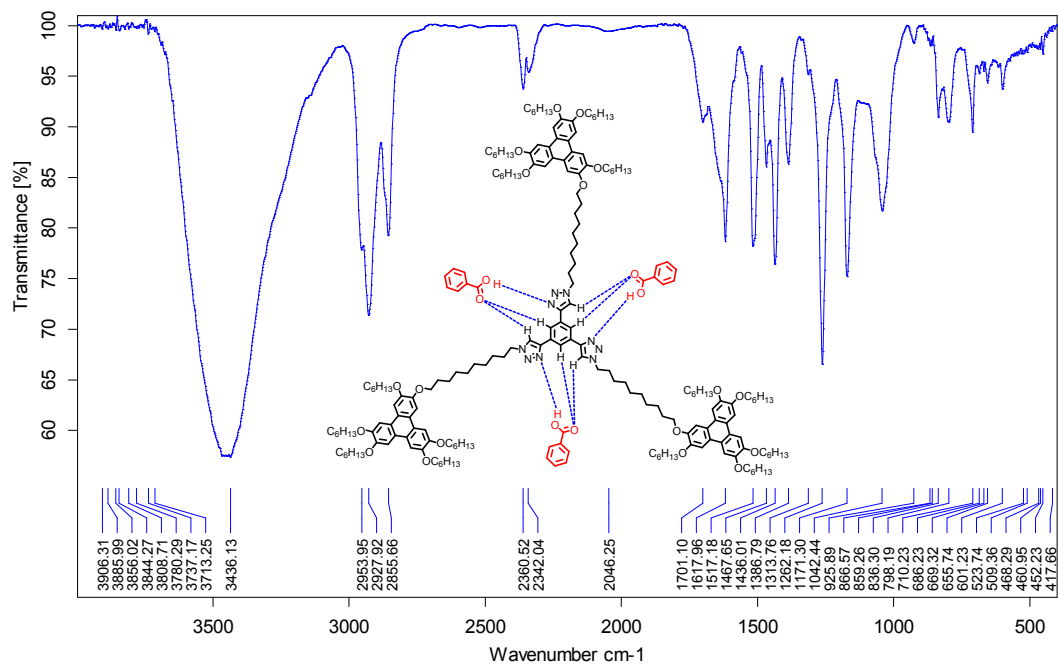


Figure S31: IR spectrum of **5d**

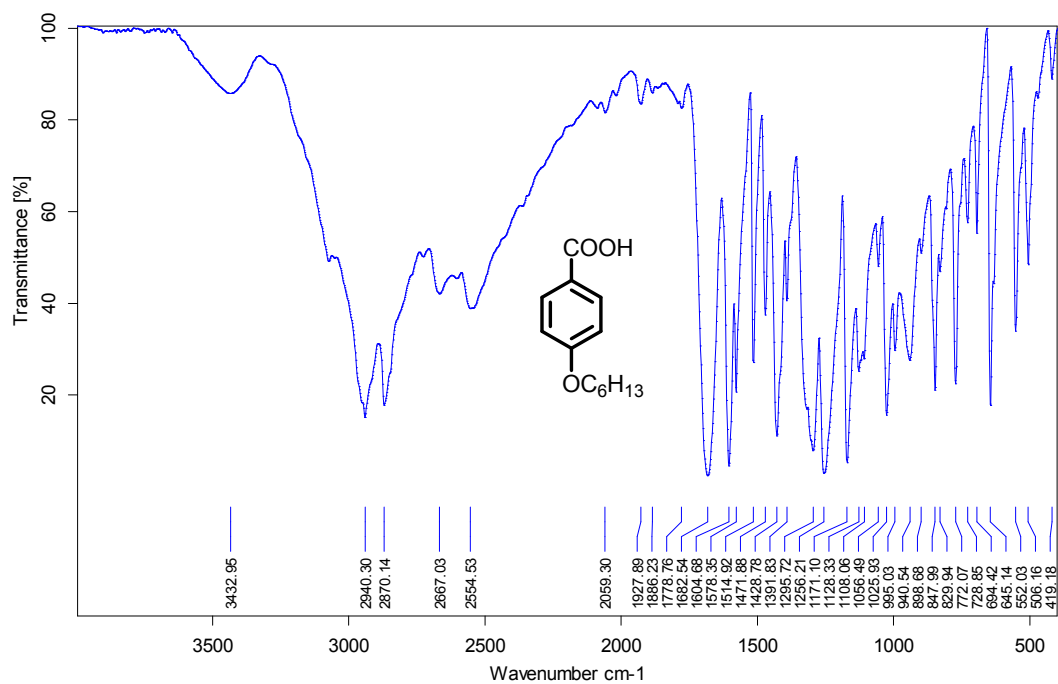


Figure S32: IR spectrum of 4-hexyloxybenzoic acid

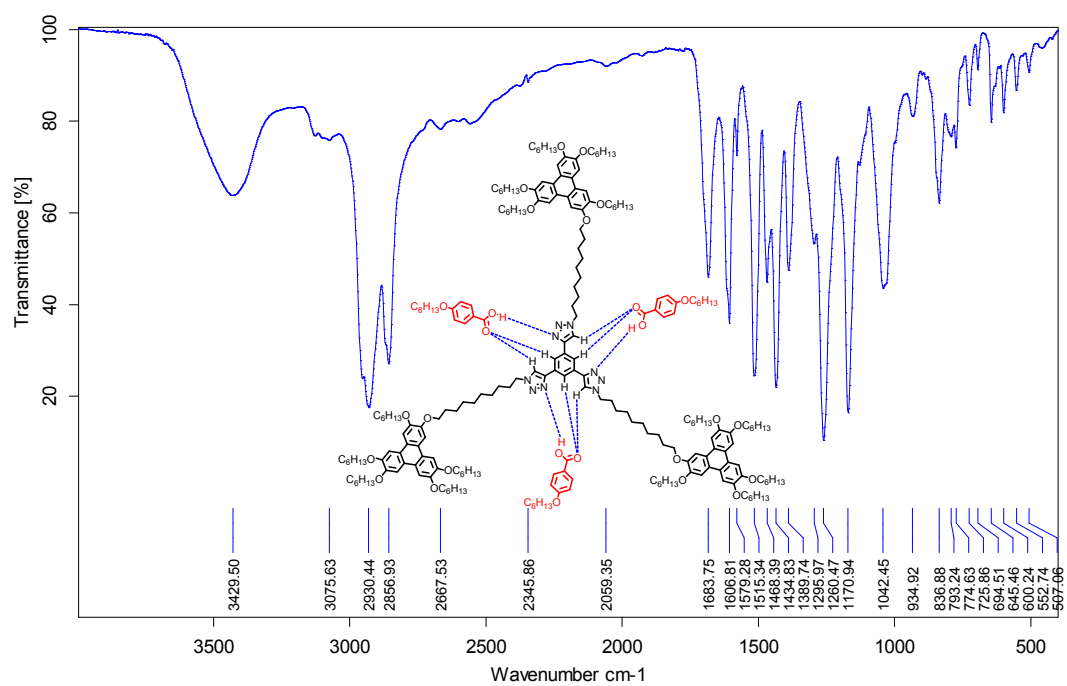


Figure S33: IR spectrum of **5f**

4. DSC

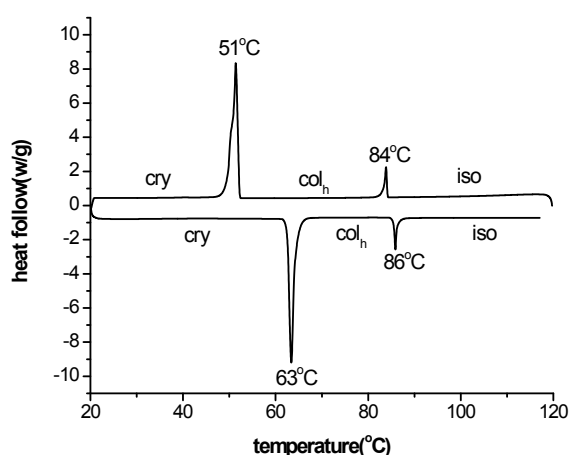


Figure S34: DSC traces of **1b**

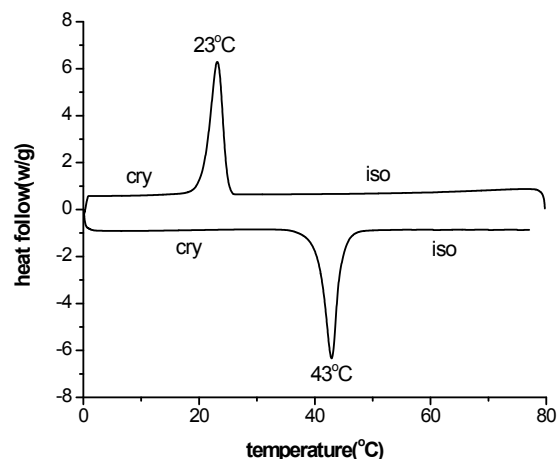


Figure S35: DSC traces of **1d**

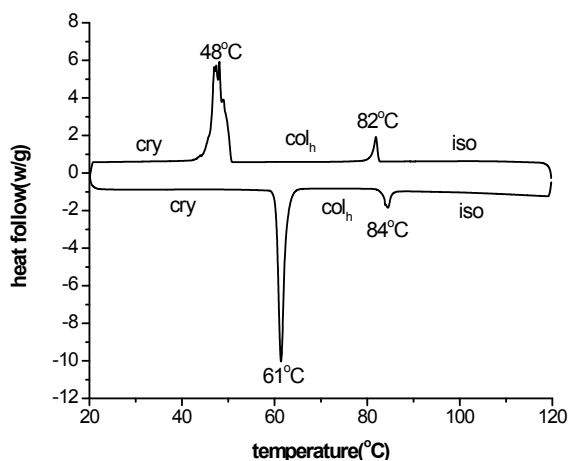


Figure S36: DSC traces of **2b**

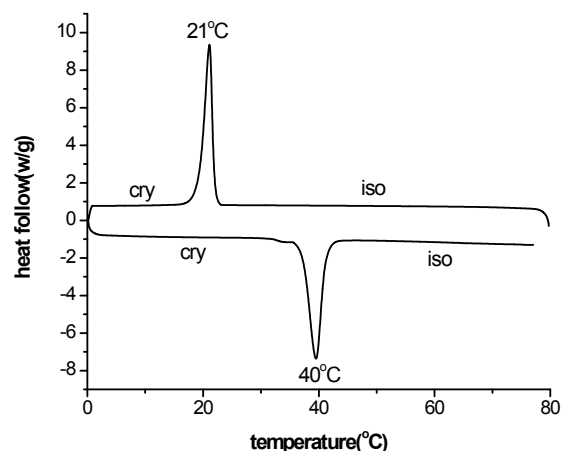


Figure S37: DSC traces of **2d**

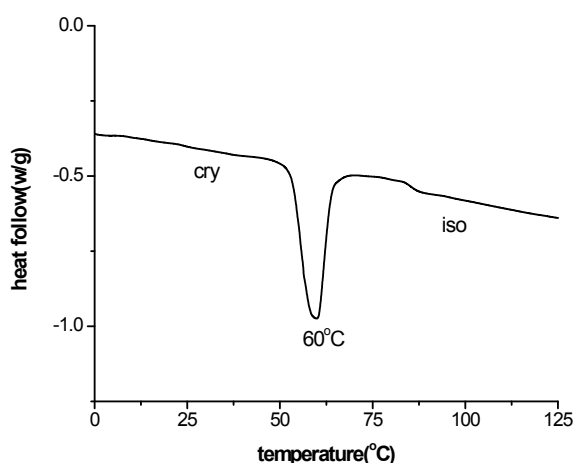


Figure S38: DSC traces of **4c**, n = 6

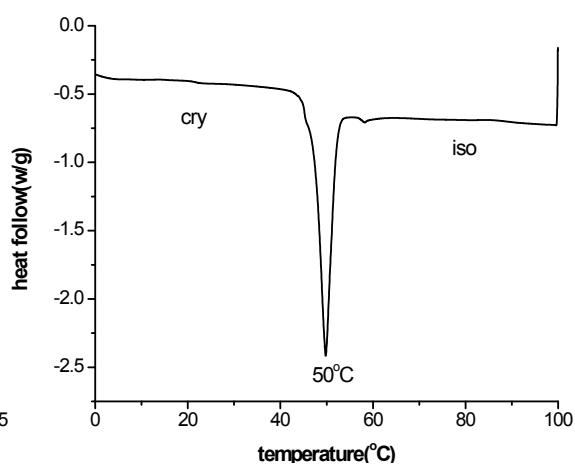


Figure S39: DSC traces of **4d**, n = 10

5. SAXS

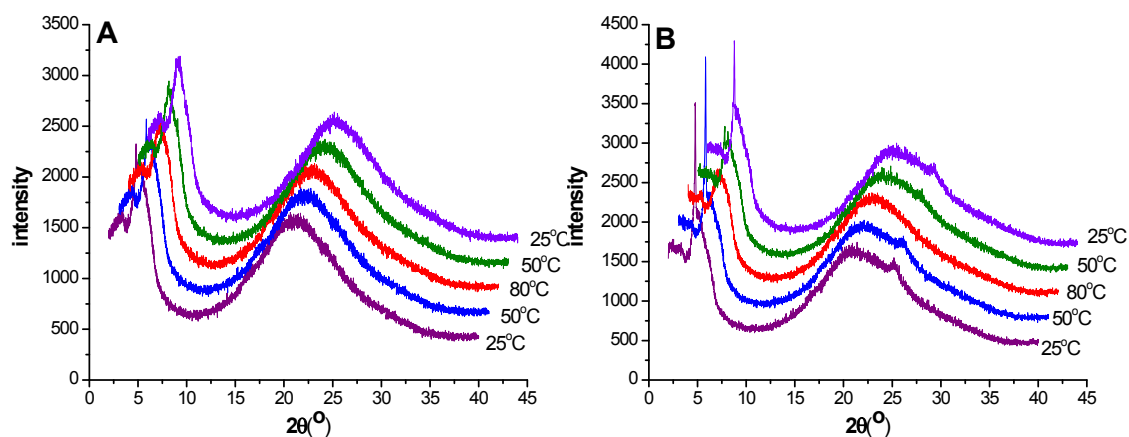


Figure S40: SAXS patterns of at variable temperatures of **5a** (A) and **5b** (B).

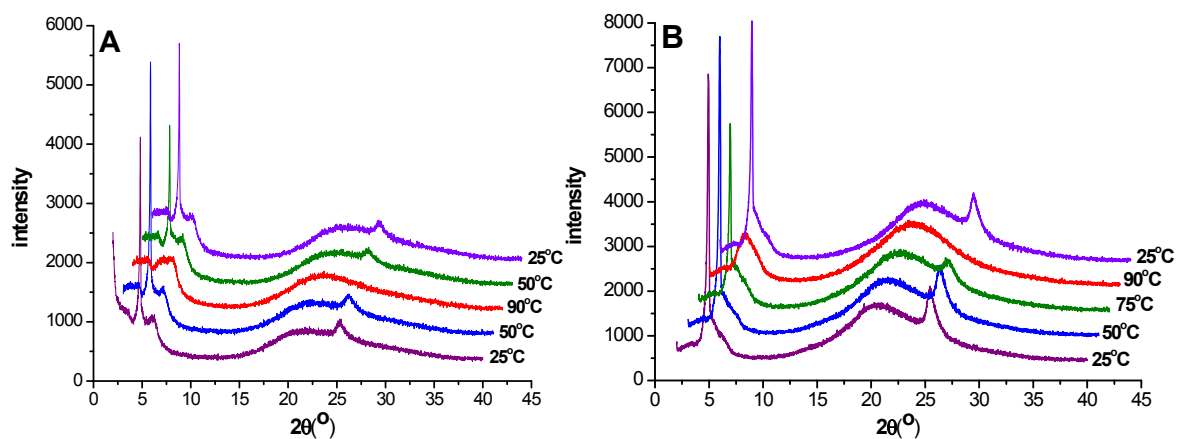


Figure S41: SAXS patterns of at variable temperatures of **5c** (A) and **5d** (B).

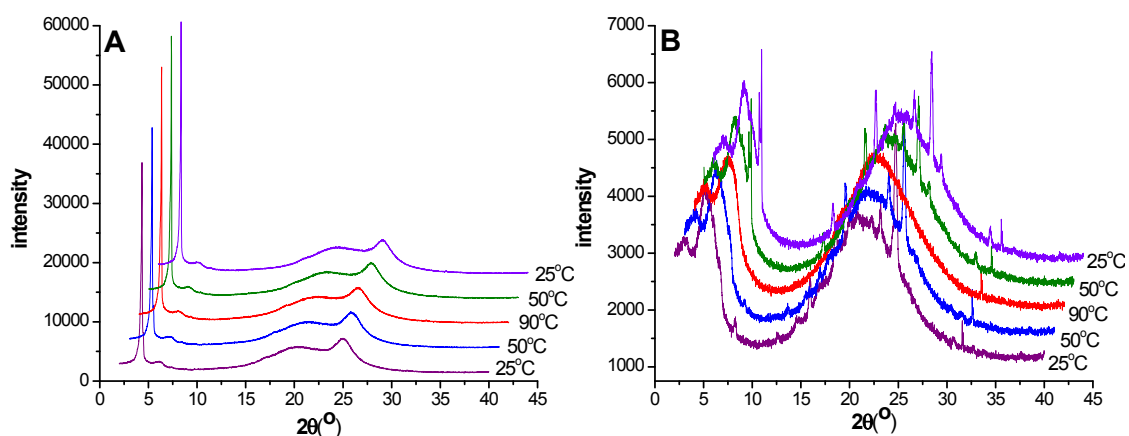


Figure S42: SAXS patterns of at variable temperatures of **5e** (A) and **5f** (B).

6. SAXS in isotropic liquid

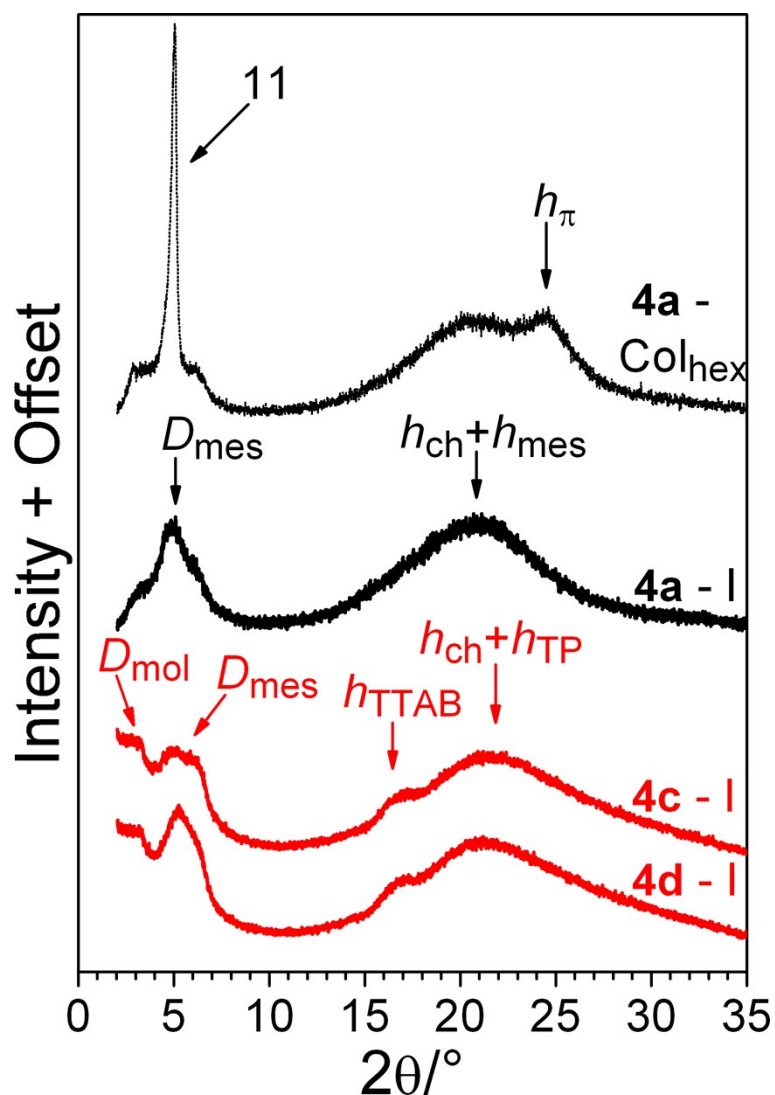


Figure S43: SAXS patterns in the isotropic state of **4a** at 130°C, **4c** at 75°C and **4d** at 70°C, compared to the room-temperature hexagonal columnar phase of **4a**; (11) and h_π are respectively the most intense reflection of the columnar lattice and the scattering signal from π -stacked triphenylene units; D_{mes} , h_{mes} and h_{ch} come from piling and lateral distances between mesogens, and from lateral distances between chains; h_{TTAB} and h_{TP} correspond to piling distances between tris-triazolyl-benzene units (TTAB) and loosely piled triphenylene units (TP); D_{mol} stands for the lateral distance between piled molecules, thus between TTAB separated by TP.