Electronic Supplementary Information

Hierarchical Superstructures from A Star-shaped Molecule Consisted of A Cyclic Oligosiloxane with Cyanobiphenyl Moieties

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Characterization data

Chemical structures and purities of SiLC and its intermediates were confirmed by thin layer chromatography (TLC), ¹H NMR, and ¹³C NMR in deuterated chloroform (CDCl₃). All the chemical compounds synthesized in this research were purified by column chromatography using the silica gel. Chemical shifts were quoted in part per million (ppm) with a reference of tetramethylsilane (TMS). Chemical structure of compound 4-cyano-4'-hex-5-en-1-oxy-biphenyl (1) was confirmed by disappearance of -C-OH proton peak corresponding to the 4'-hydroxy-4-biphenyl-carbonitrile and appearance of -CH=CH₂ of olefin group (5.03 ppm, 2H). Chemical structure of SiLC (2) was identified by increased integration of total protons and a new peak as -Si-CH₃ (0.08 ppm, 12H) corresponding to the 2,4,6,8-tetramethylcyclotetrasiloxane. Carbon atoms of SiLC molecules were identified by both solution (Fig. S3) and solid state (Fig. 5) ¹³C NMR and their peak position of carbon atoms were also well matched. The carbon atoms in cyanobiphenyl groups appear between 100 ppm and 170 ppm, while carbons in the alkyl chains are chemically shifted between 100 ppm.



Fig. S1 ¹H NMR spectrum of compound 1



Fig. S2 ¹H NMR spectrum of compound $\bf 2$



Fig. S3 ¹³C NMR spectrum of compound 2