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Electronic Supplementary Information

Side-chain Polysiloxane Liquid Crystalline Elastomers from Nonmesogenic Components[†]

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Synthesis of 1-Phenyl-2(4-(undec-10-en-1-yloxy) phenyl)diazene, monomer (M)



SchemeS1 Synthetic pathway followed to prepare olefinic terminated monomer M



Fig.S1 FT - IR spectrum of azo monomer M



Fig.S2 ¹H NMR spectrum of azo monomer M



Fig.S3 DSC thermogram of azo monomer (M) recorded at a heating and cooling rate of

10°C/min





Scheme S2 Synthetic route to prepare vinyl terminated 11UB cross-linking agent, C1



Fig.S4 FT-IR spectrum of cross-linker C1







Fig.S6 DSC thermogram of cross-linker (C1) recorded at a heating and cooling rate of 10°C/min

Synthesis of biphenyl cross-linker C₂



Scheme S3 Synthetic route for cross-linker C2



Fig.S7 FT-IR spectrum of biphenyl cross-linker C₂







Fig.S9 DSC thermogram of cross-linker (C_2) recorded at a heating and cooling rate of 10°C/min



Fig. S10 An optimised molecular model for the cross-linker C_2 in a stretched *all trans* alkyl chain conformation (from ACD chemlab-3D viewer).



Fig. S11 UV-Visible absorption spectra of azo monomer in toluene (a) by shining UV light at 254 nm; *trans* to *cis* isomerisation, (b) by shining visible light(>540);*cis* to *trans* isomerisation.



Fig. S12 The peak absorbance of monomer (348nm) with respect to exposure time for (a) *trans* to *cis* isomerisation and (b) *cis* to *trans* isomerisation.

Video S1: Bending of the LCE film upon UV irradiation (254nm)

Video S2: Unbending of the LCE film upon visible irradiation (>540 nm)