

Supplementary Information

Monodisperse oligo(ϵ -caprolactones) with terpenes and alkyl end-groups: synthesis, isolation, characterization, and antibacterial activity

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Table S1 Hydrolytic degradation of initiators (farnesol and 1-pentadecanol), monodisperse monomers (C₁₅F-CL₁ and C₁₅1P-CL₁), and monodisperse dimers (C₁₅F-CL₂ and C₁₅1P-CL₂).

Sample	T_{d1} (°C)	Weight loss (%)	T_{d2} (°C)	Weight loss (%)	T_{d3} (°C)	Weight loss (%)
Farnesol	296	42	368	83	—	—
1-pentadecanol	250	83	—	—	—	—
C ₁₅ F-CL ₁	114	3	386	31	399	70
C ₁₅ 1P-CL ₁	316	54	393	91	—	—
C ₁₅ F-CL ₂	132	2	298	23	410	67
C ₁₅ 1P-CL ₂	369	47	410	80	—	—

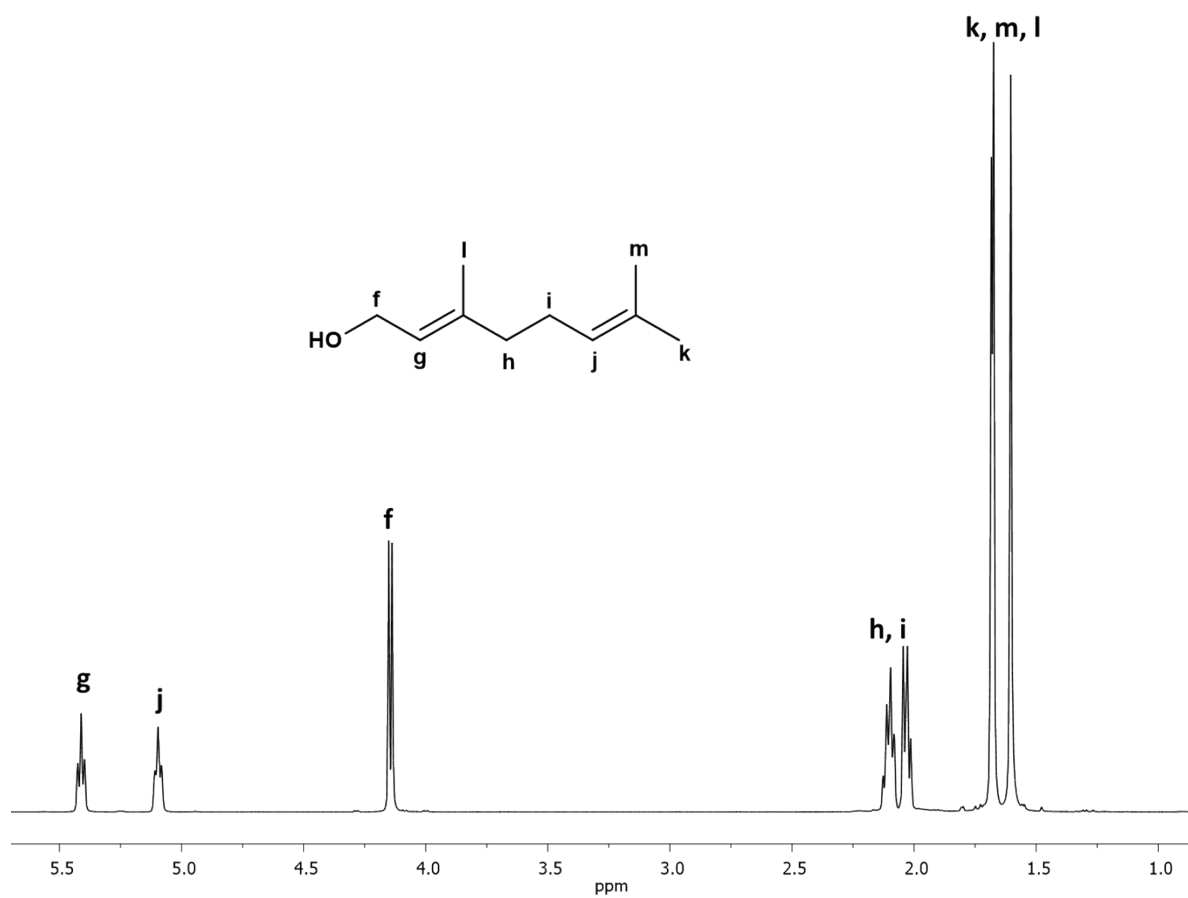


Fig. S1 ^1H NMR (500 MHz) spectrum of geraniol (C_{10}) in CDCl_3 at 40°C .

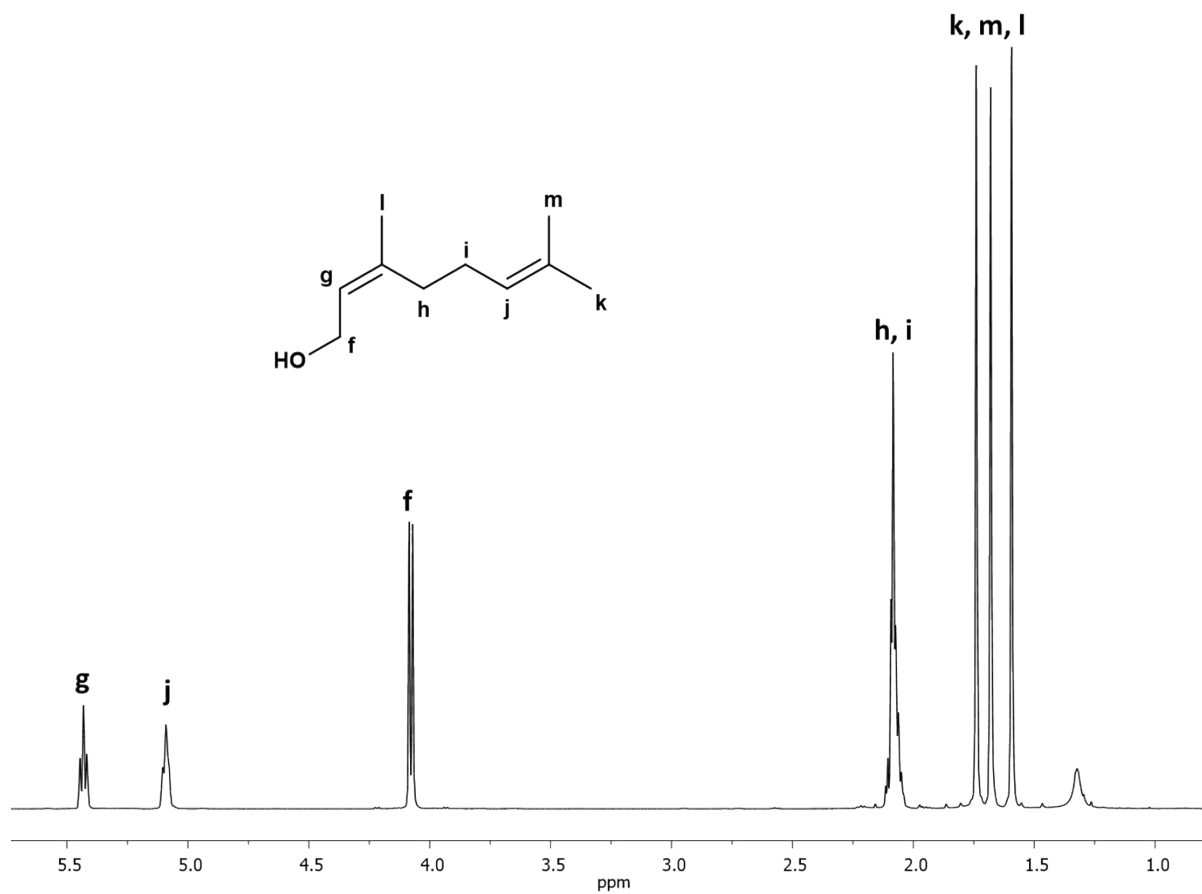


Fig. S2 ^1H NMR (500 MHz) spectrum of nerol (C_{10}) in CDCl_3 at 40°C .

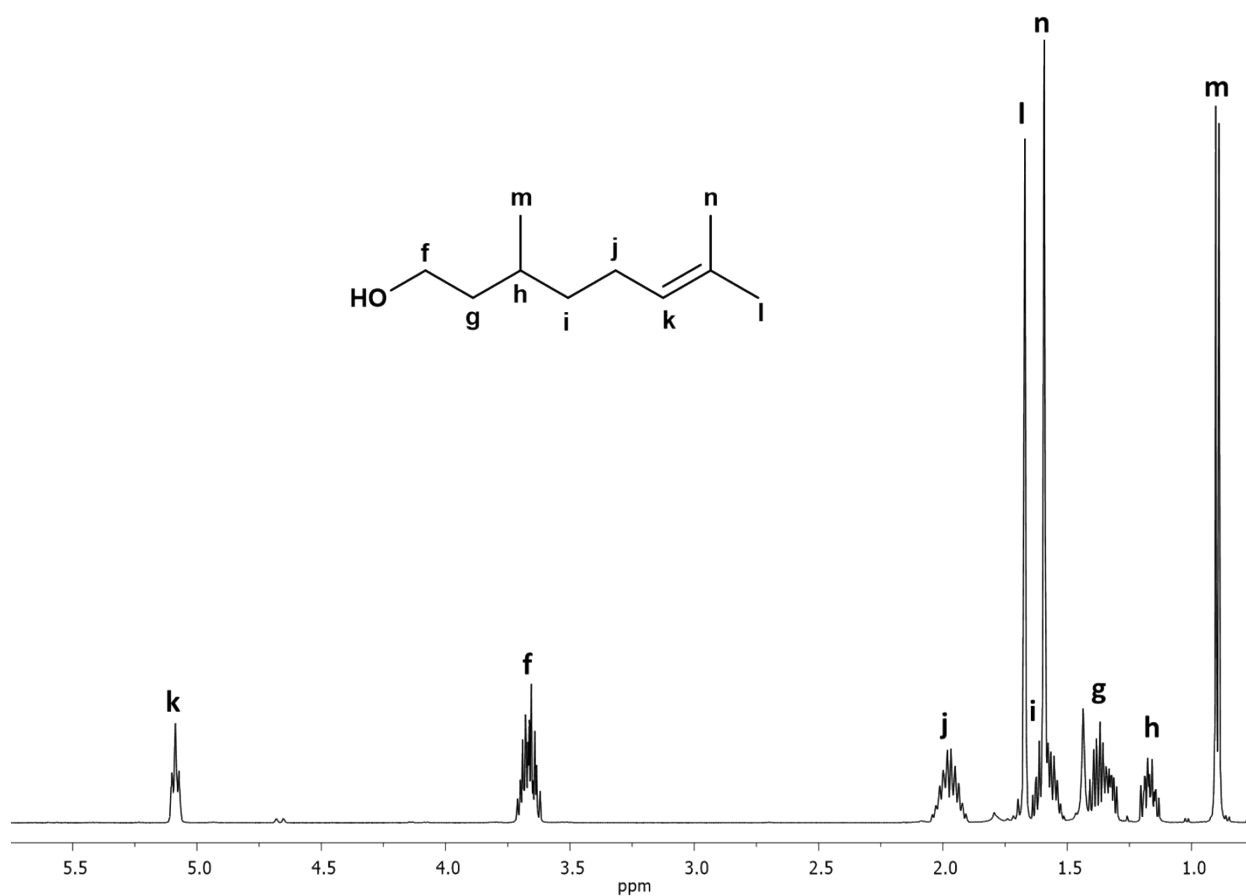


Fig. S3 ^1H NMR (500 MHz) spectrum of β -citronellol (C_{10}) in CDCl_3 at 40°C .

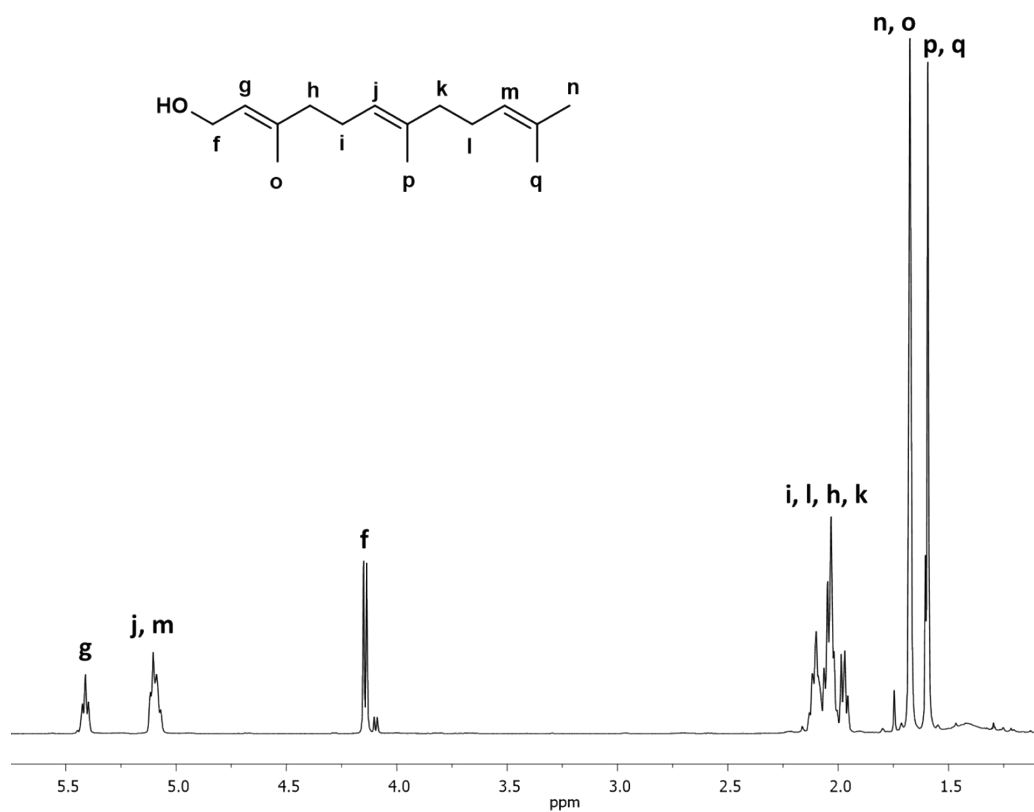


Fig. S4 ¹H NMR (500 MHz) spectrum of farnesol (C₁₅) in CDCl₃ at 40 °C.

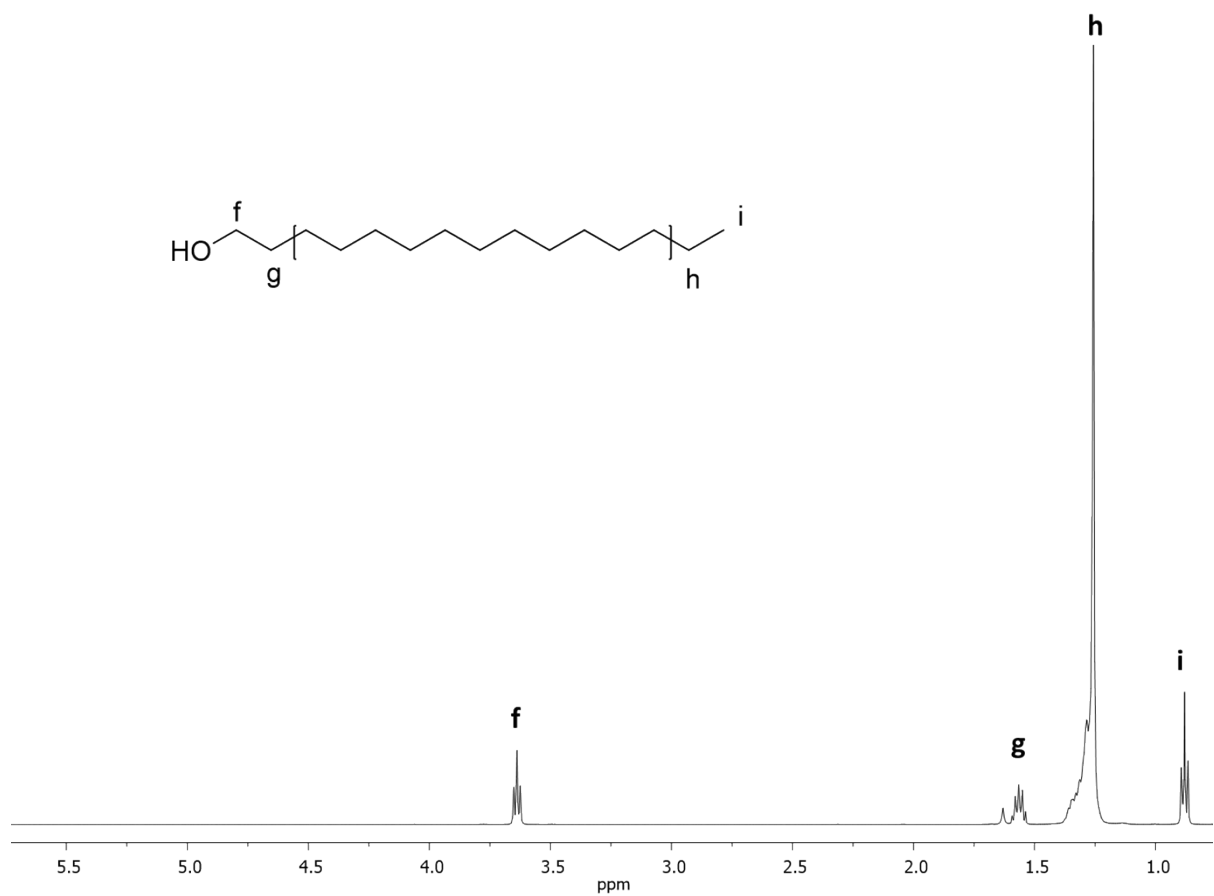


Fig. S5 ¹H NMR (500 MHz) spectrum of 1-pentadecanol (C₁₅) in CDCl₃ at 40 °C.

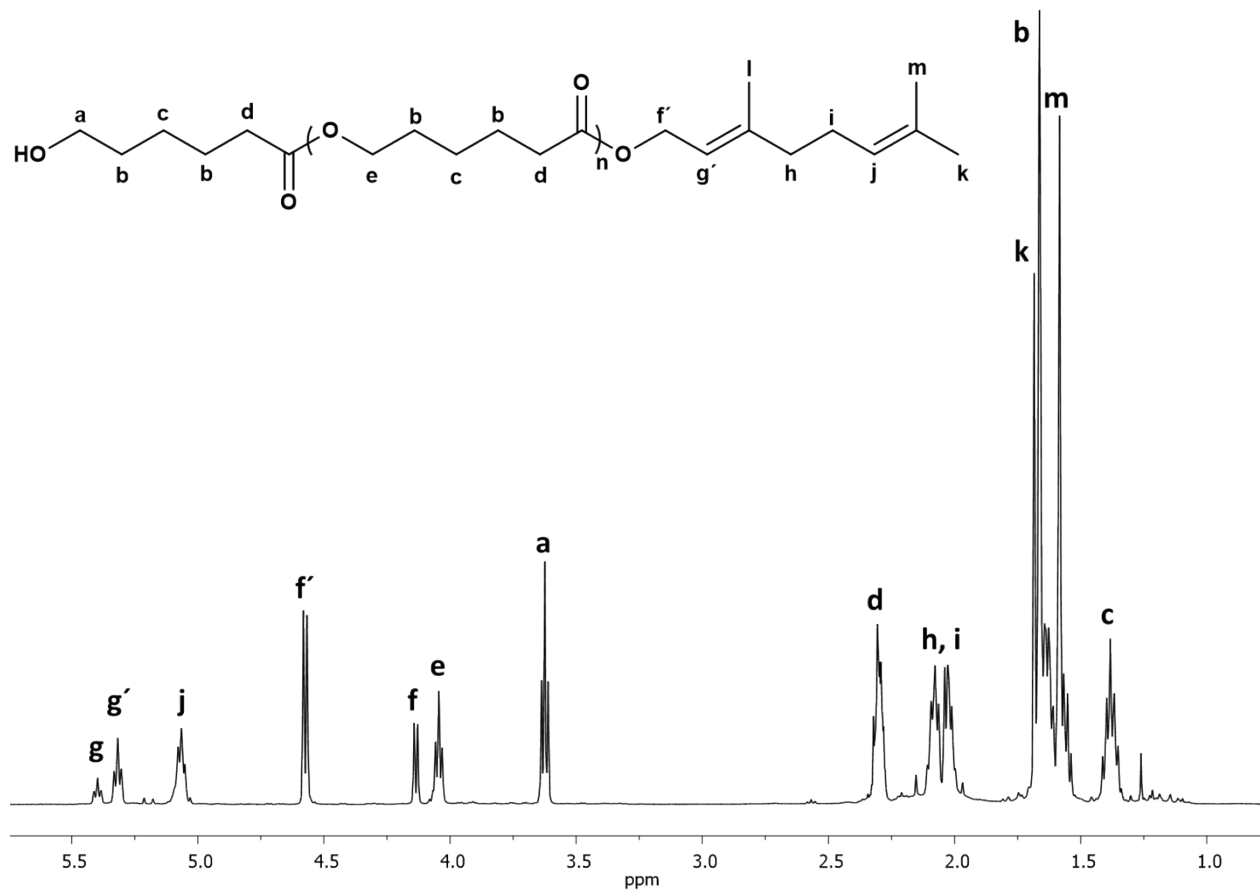


Fig. S6 ¹H NMR (500 MHz) spectrum of oligo(CL) synthesized using geraniol as initiator (C₁₀G-PCL, Table 1) in CDCl₃ at 40 °C.

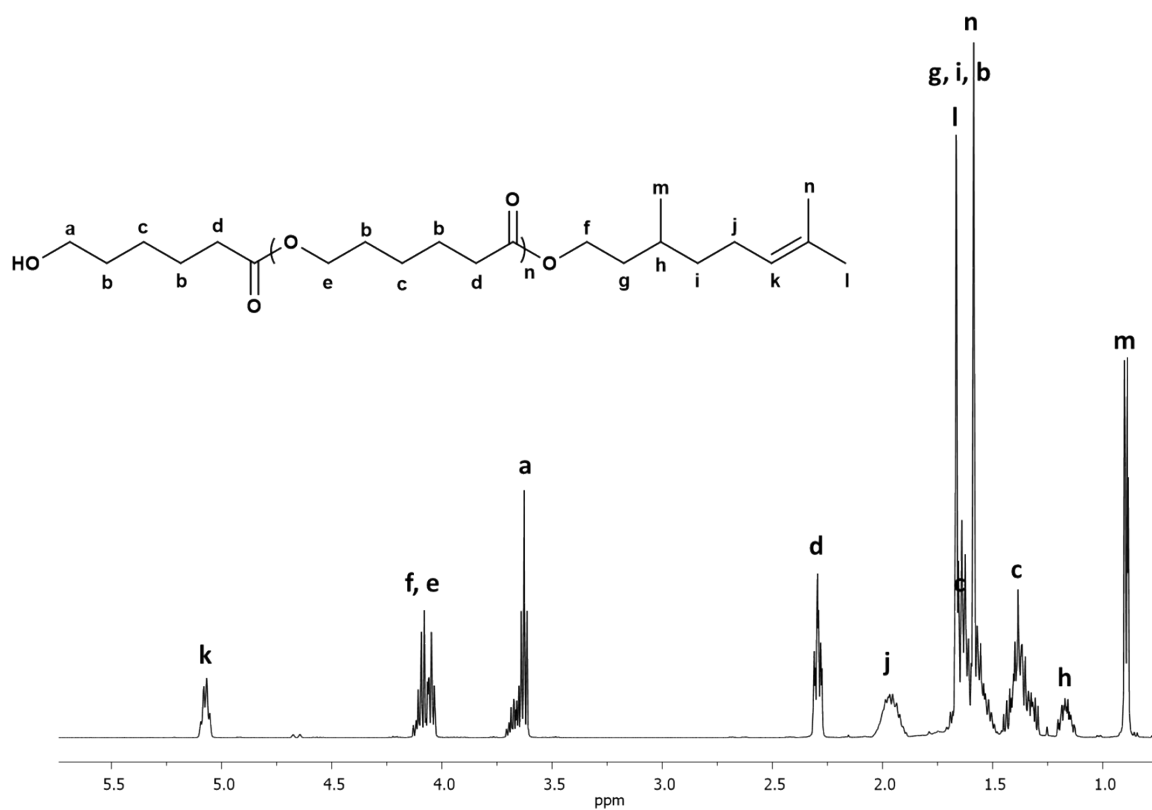


Fig. S7 ¹H NMR (500 MHz) spectrum of oligo(CL) synthesized using β -citronellol as initiator (C₁₀C-PCL, Table 1) in CDCl₃ at 40 °C.

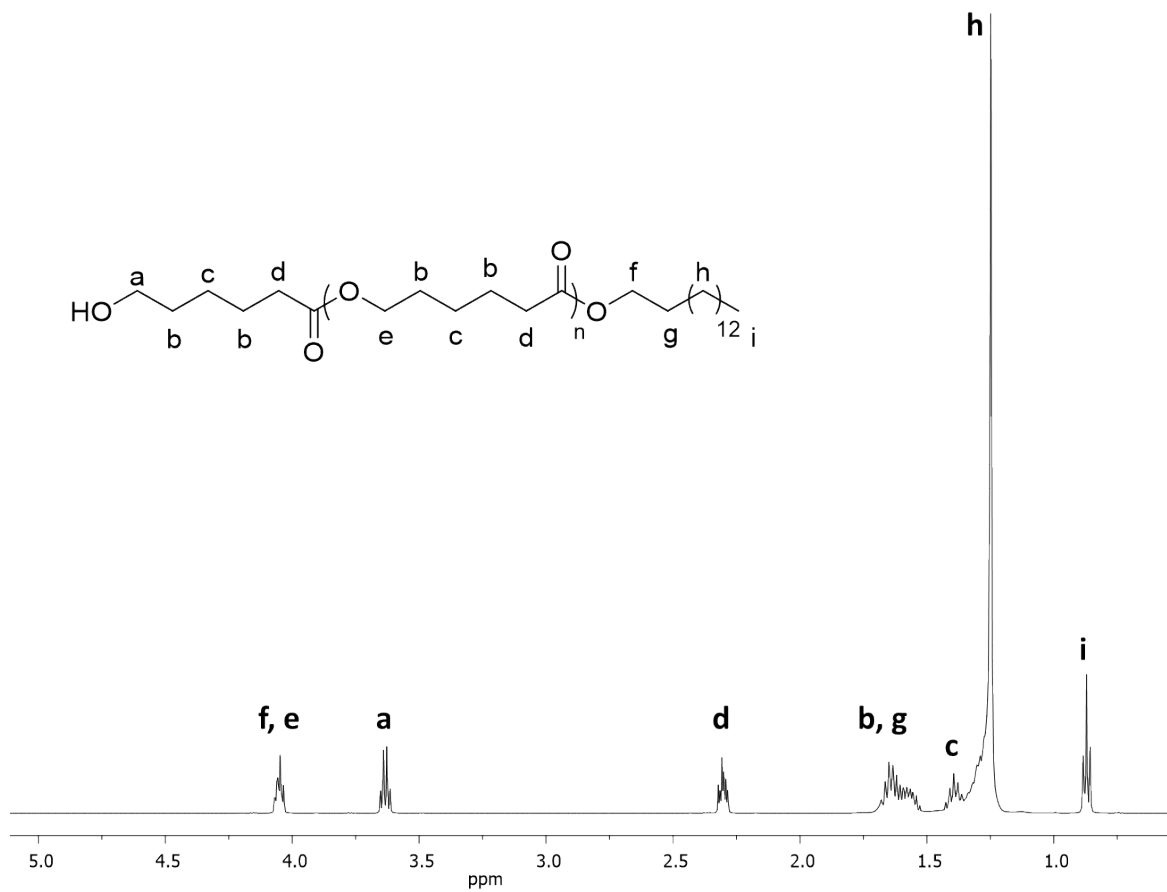


Fig. S8 ¹H NMR (500 MHz) spectrum of oligo(CL) synthesized using 1-pentadecanol as initiator (C₁₅1P-PCL, Table 1) in CDCl₃ at 40 °C.

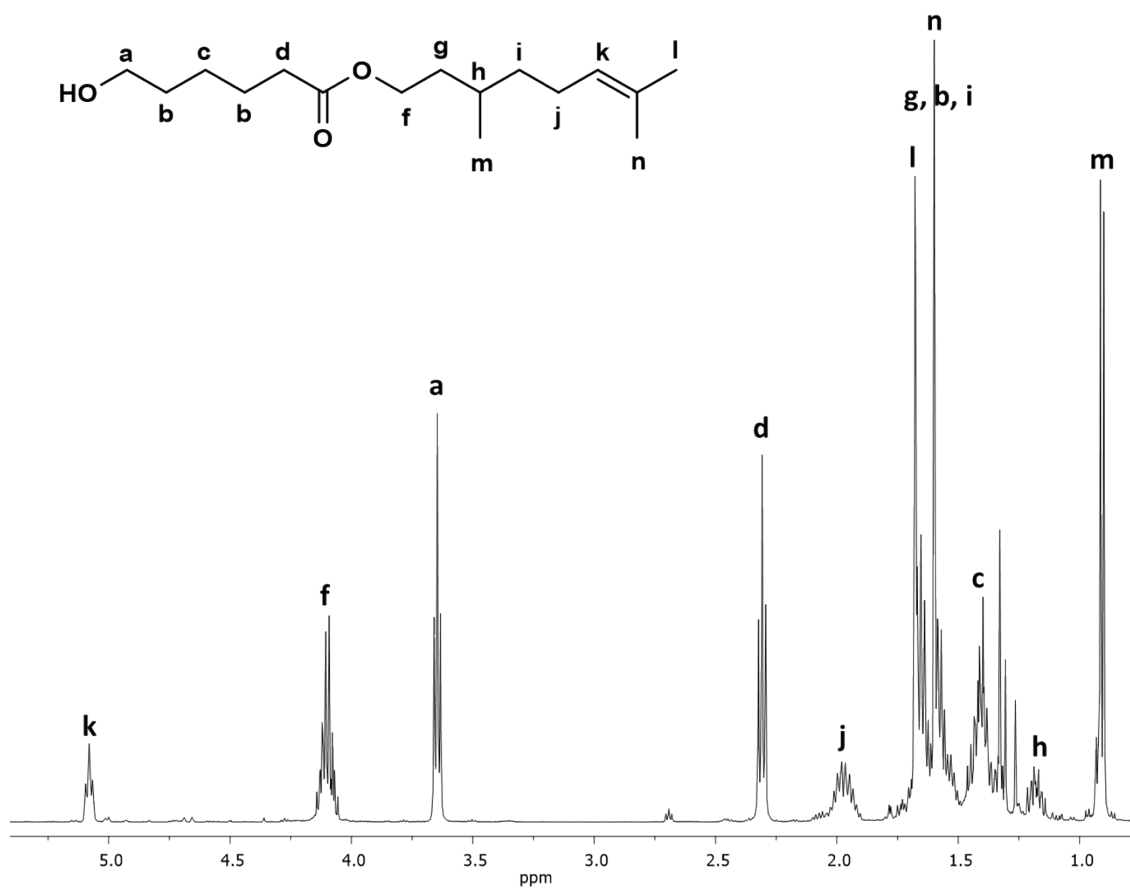


Fig. S9 ¹H NMR (500 MHz) spectrum of a monomer derived from β -citronellol as initiator C₁₀C-CL₁ (monodisperse specie, Table 2) in CDCl₃ isolated by FCC from C₁₀C-PCL (Table 1).

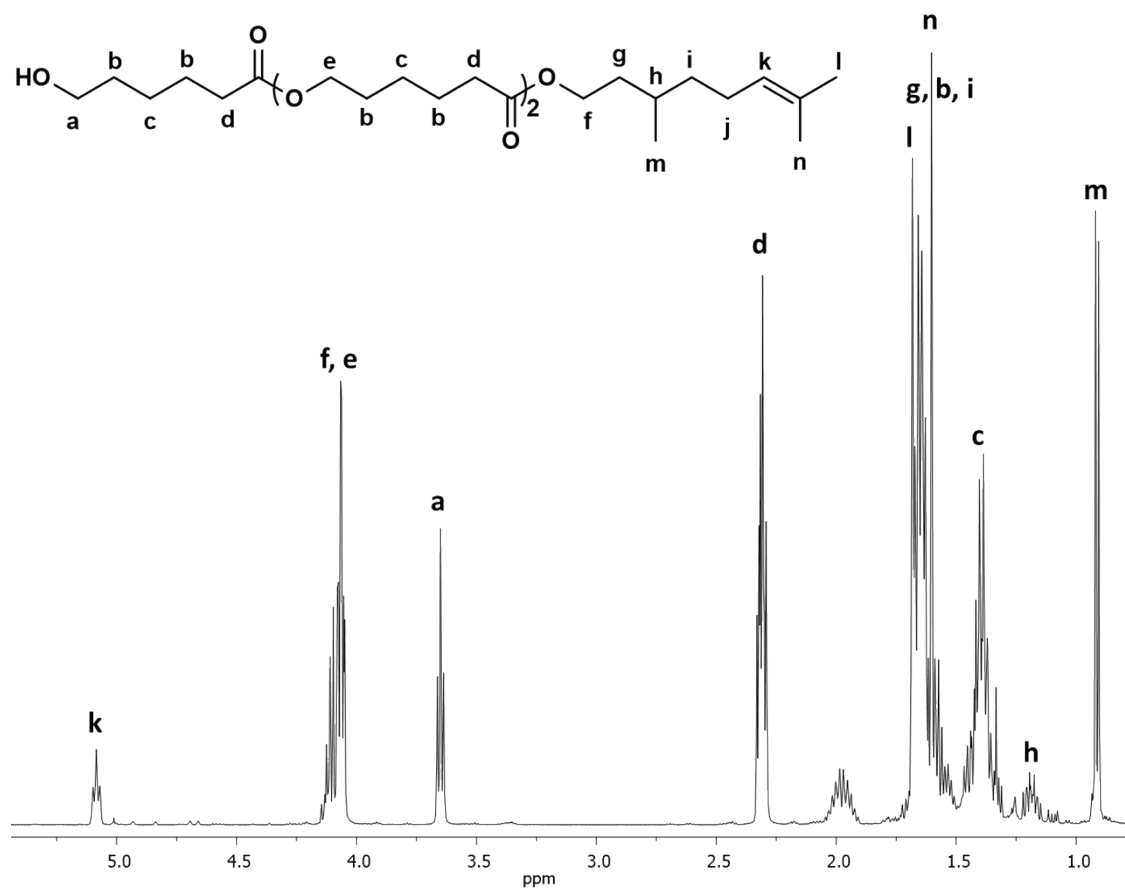


Fig. S10 ^1H NMR (500 MHz) spectrum of a trimer derived from β -citronellol as initiator $\text{C}_{10}\text{C-CL}_3$ (monodisperse specie, Table 2) in CDCl_3 isolated by FCC from $\text{C}_{10}\text{C-PCL}$ (Table 1).

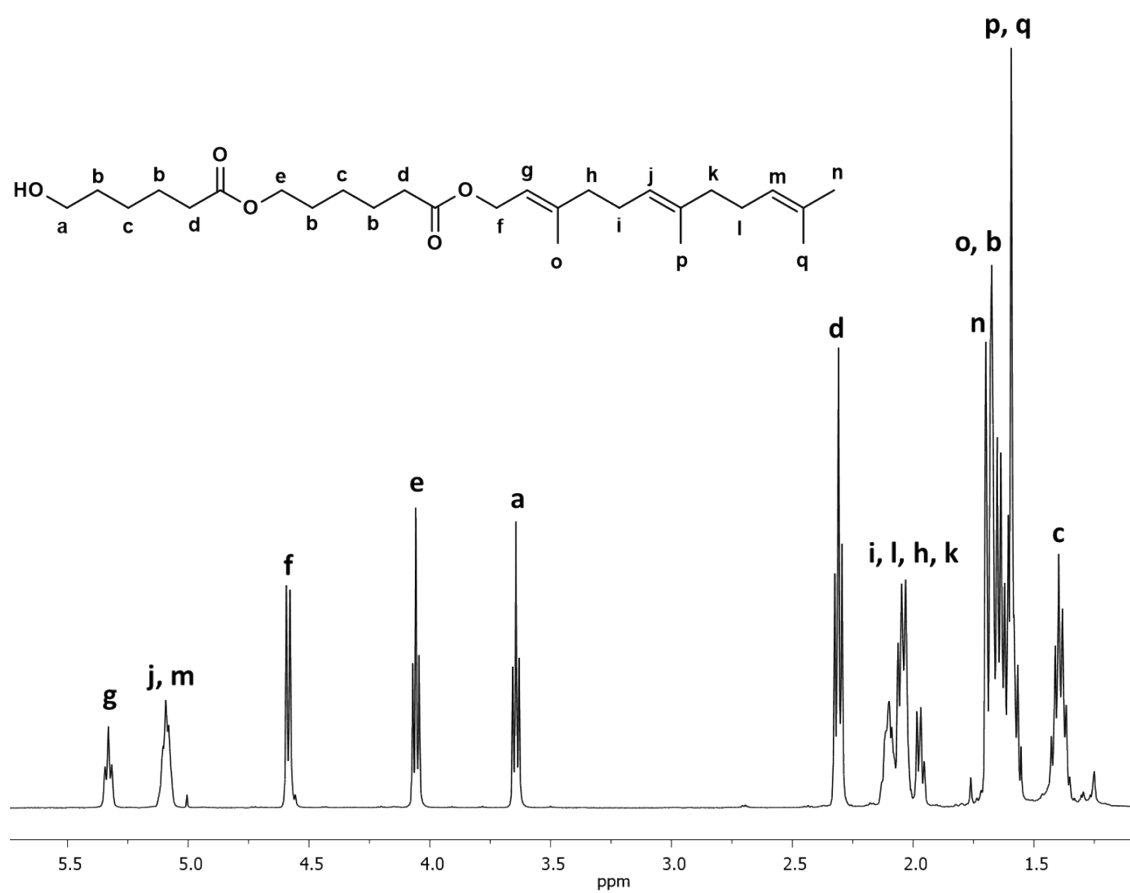


Fig. S11 ¹H NMR (500 MHz) spectrum of a dimer derived from farnesol as initiator C₁₅F-CL₂ (monodisperse specie, Table 2) in CDCl₃ isolated by FCC from C₁₅F-PCL (Table 1).

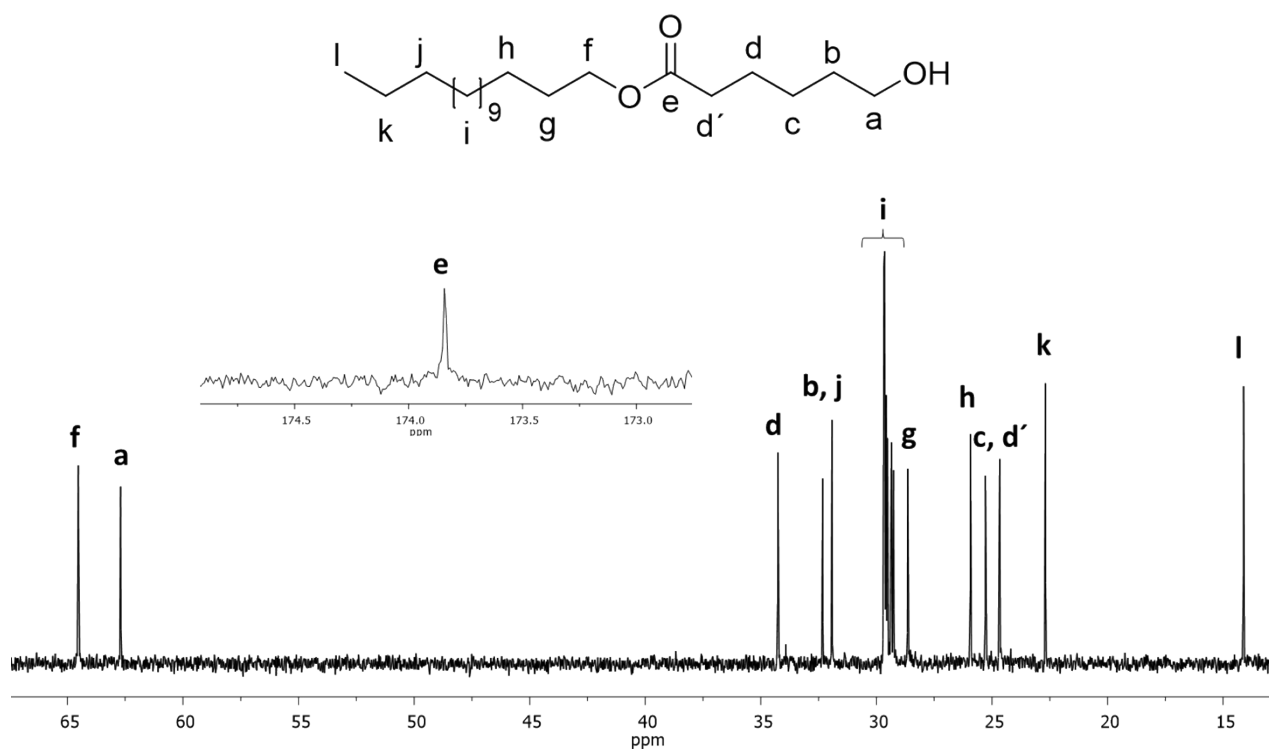


Fig. S12 ^{13}C NMR (500 MHz) spectrum of a monomer derived from 1-pentadecanol as initiator $\text{C}_{15}\text{1P-CL}_1$ (monodisperse specie, Table 2) in CDCl_3 isolated by FCC from $\text{C}_{15}\text{1P-PCL}$ (Table 1).

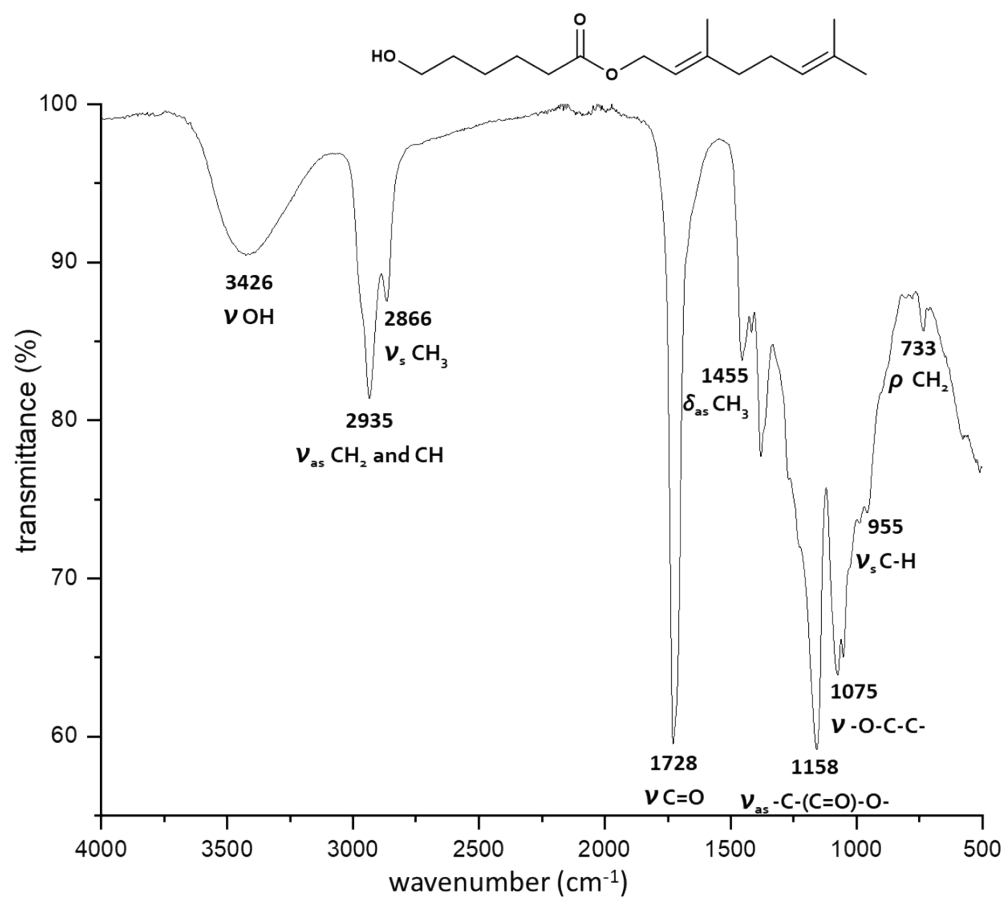


Fig. S13 FT-IR spectrum and assignment of bands from monodisperse monomer $C_{10}G-CL_1$.

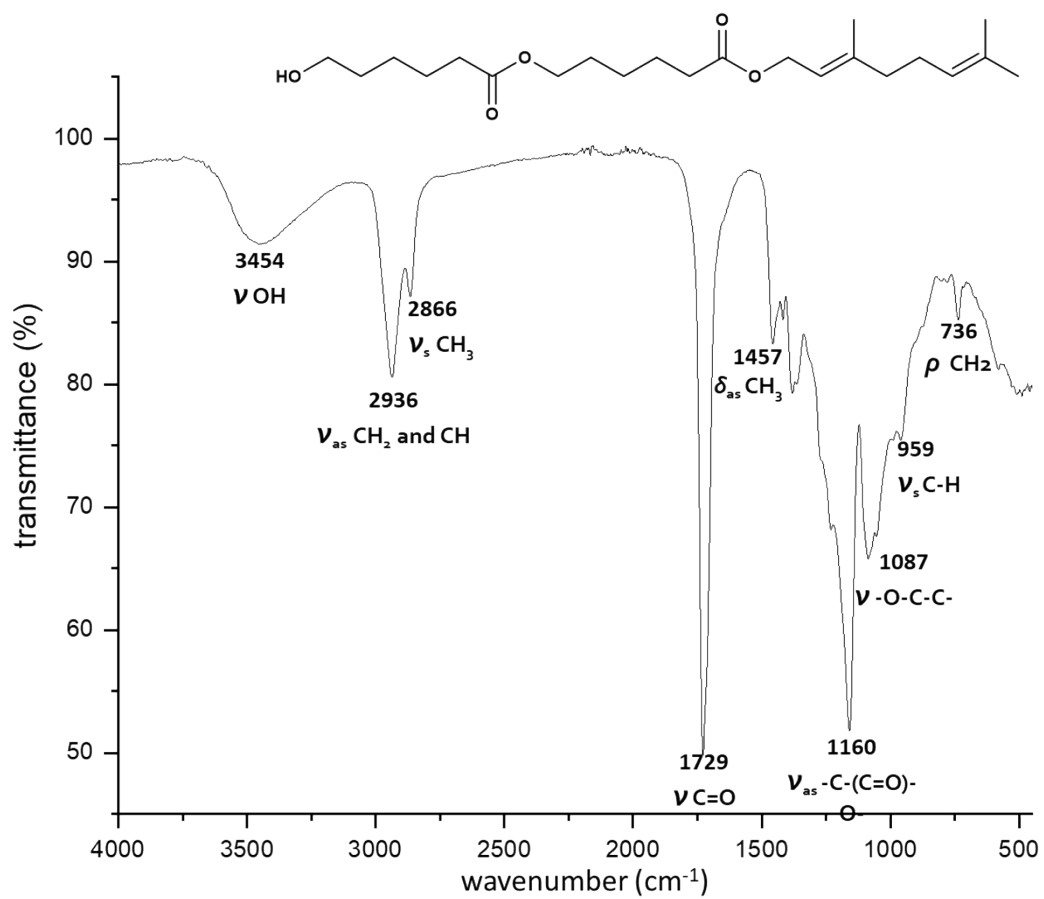


Fig. S14 FT-IR spectrum and assignment of bands from monodisperse dimer C₁₀G-CL₂.

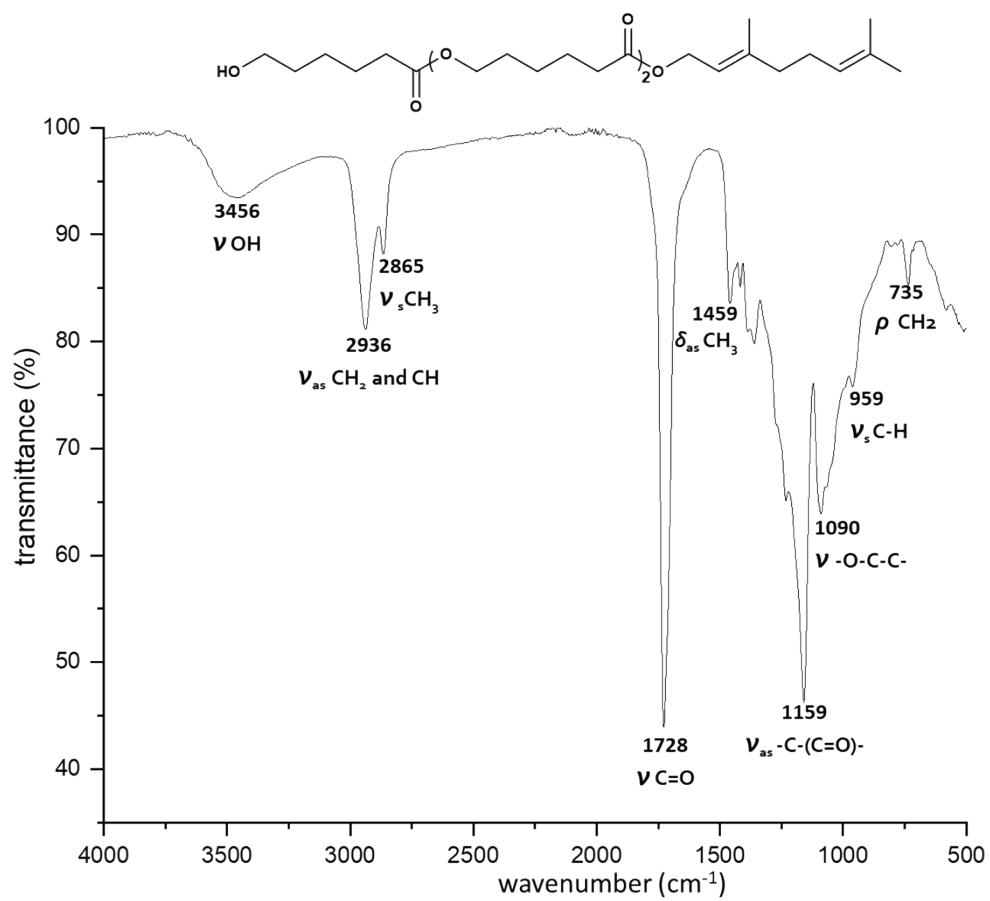


Fig. S15 FT-IR spectrum and assignment of bands from monodisperse trimer C₁₀G-CL₃.

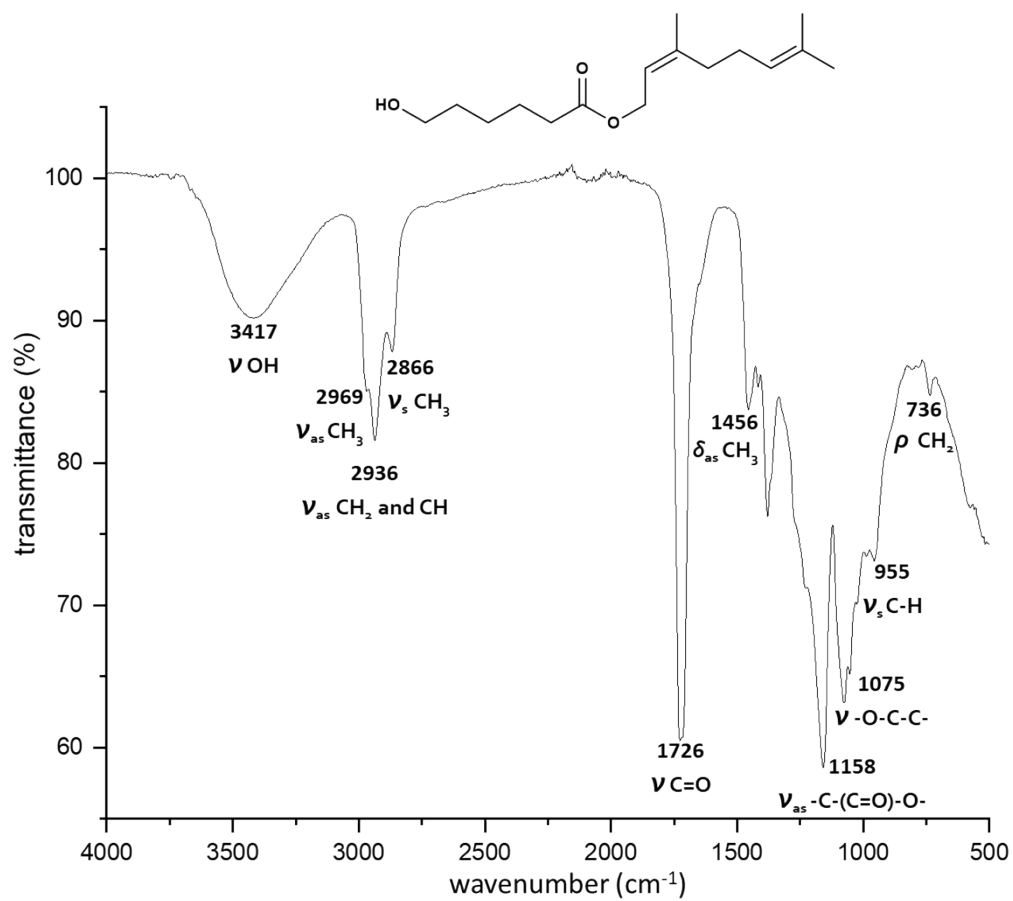


Fig. S16 FT-IR spectrum and assignment of bands from monodisperse monomer C₁₀N-CL₁.

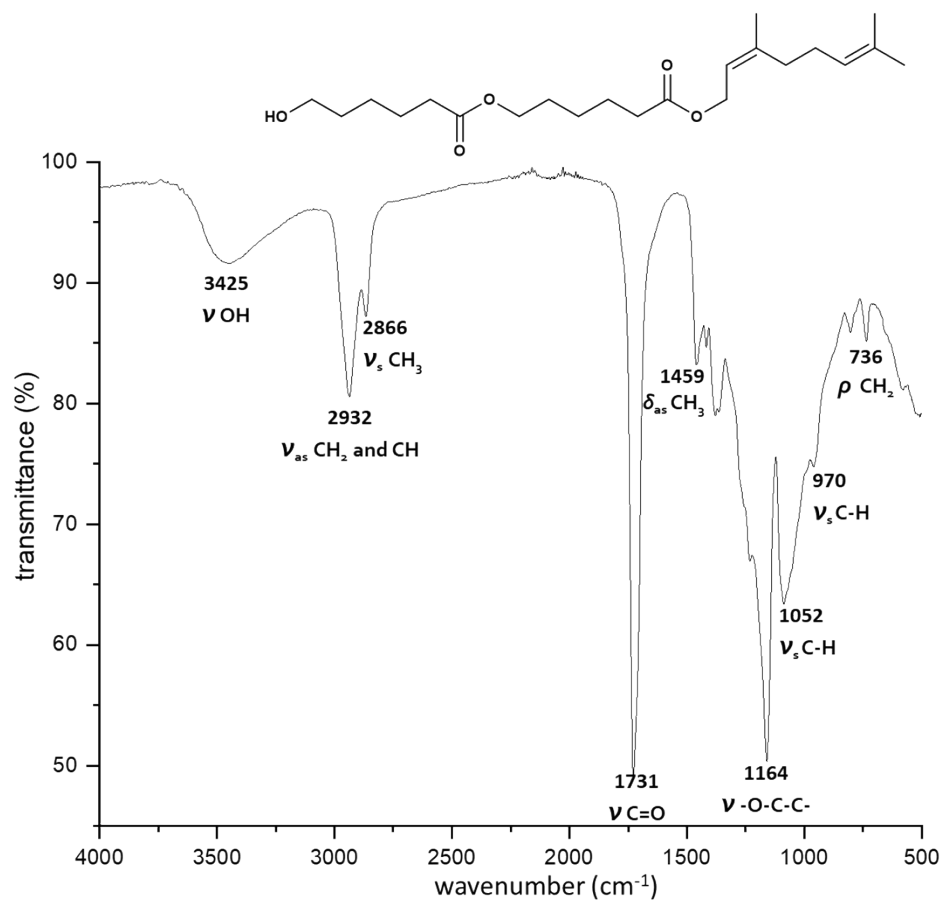


Fig. S17 FT-IR spectrum and assignment of bands from monodisperse dimer C₁₀N-CL₂.

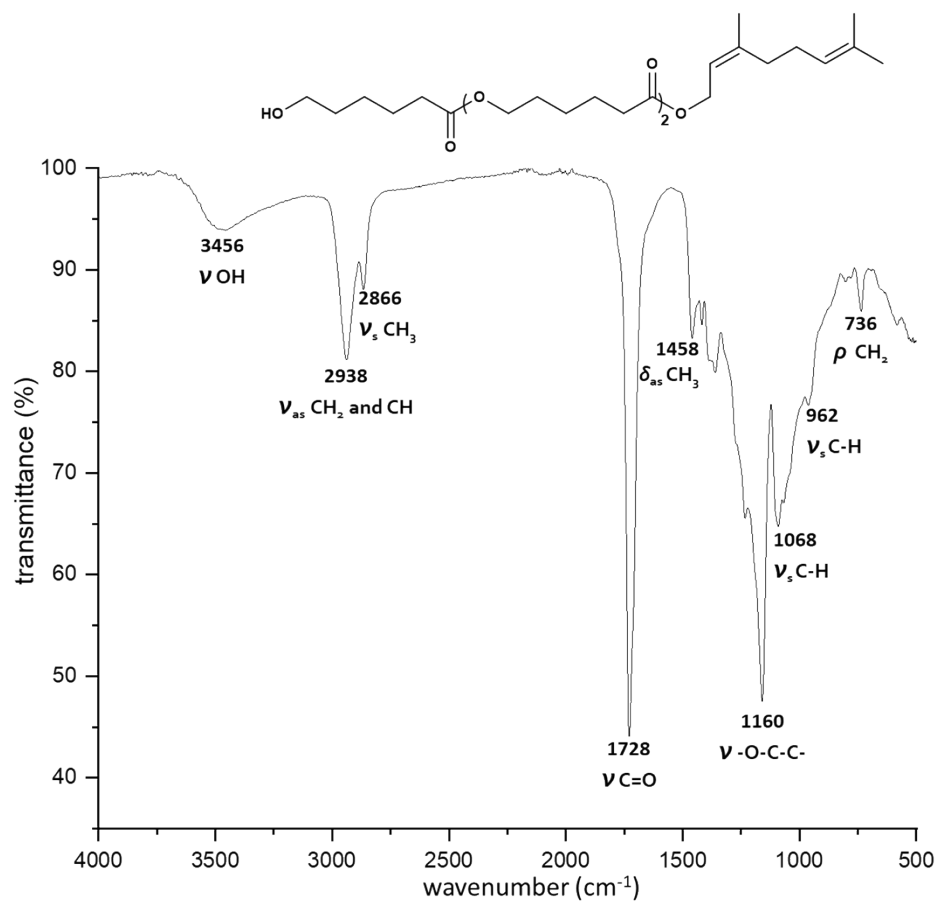


Fig. S18 FT-IR spectrum and assignment of bands from monodisperse trimer $C_{10}N-CL_3$.

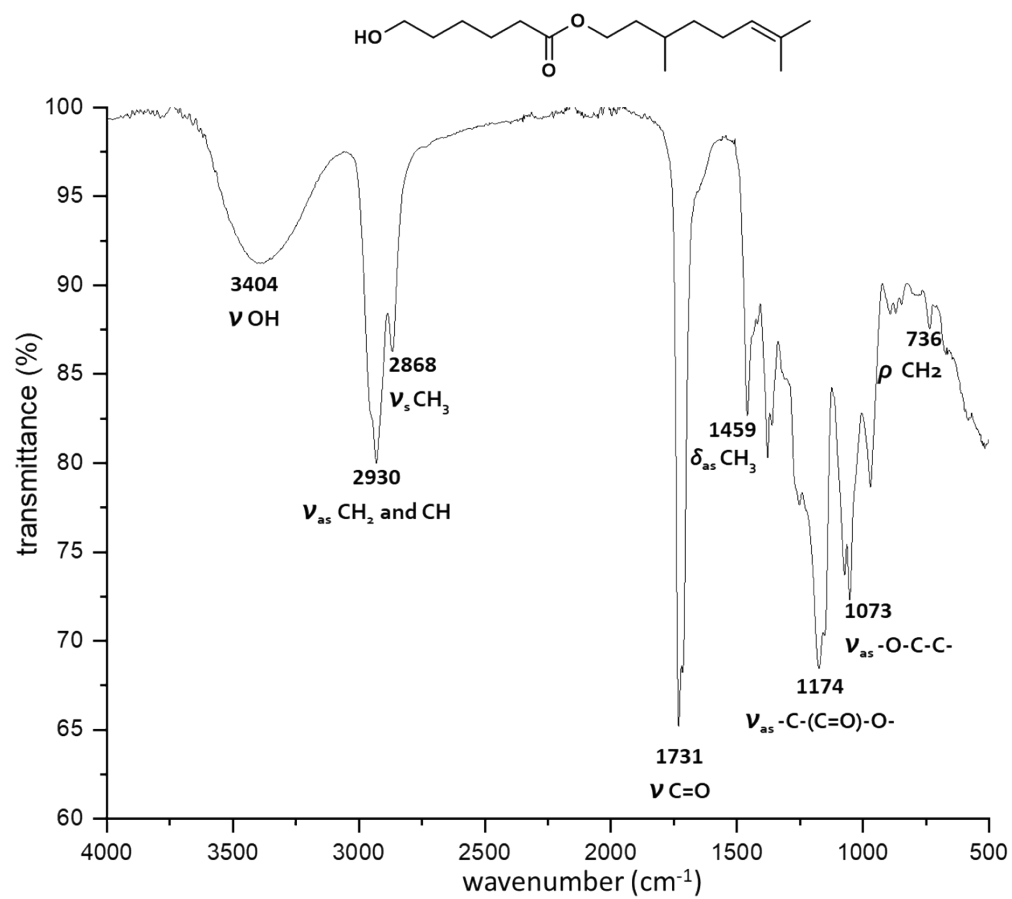


Fig. S19 FT-IR spectrum and assignment of bands from monodisperse monomer C₁₀C-CL₁.

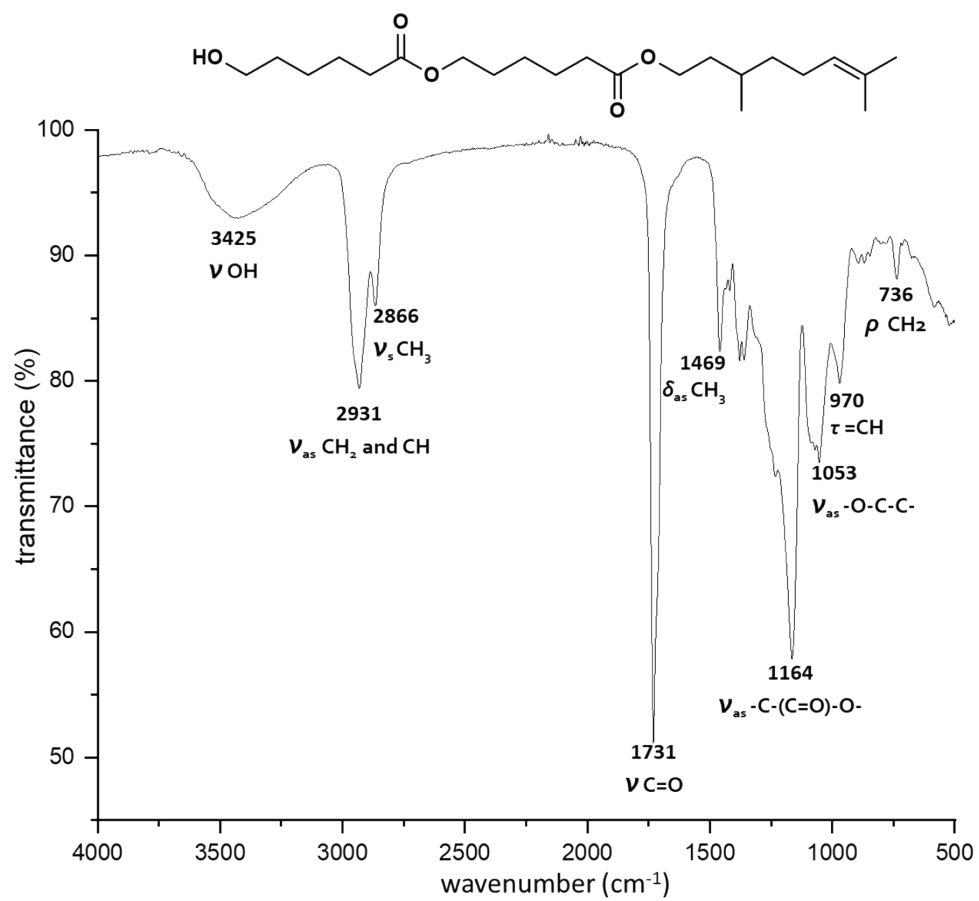


Fig. S20 FT-IR spectrum and assignment of bands from monodisperse dimer C₁₀C-CL₂.

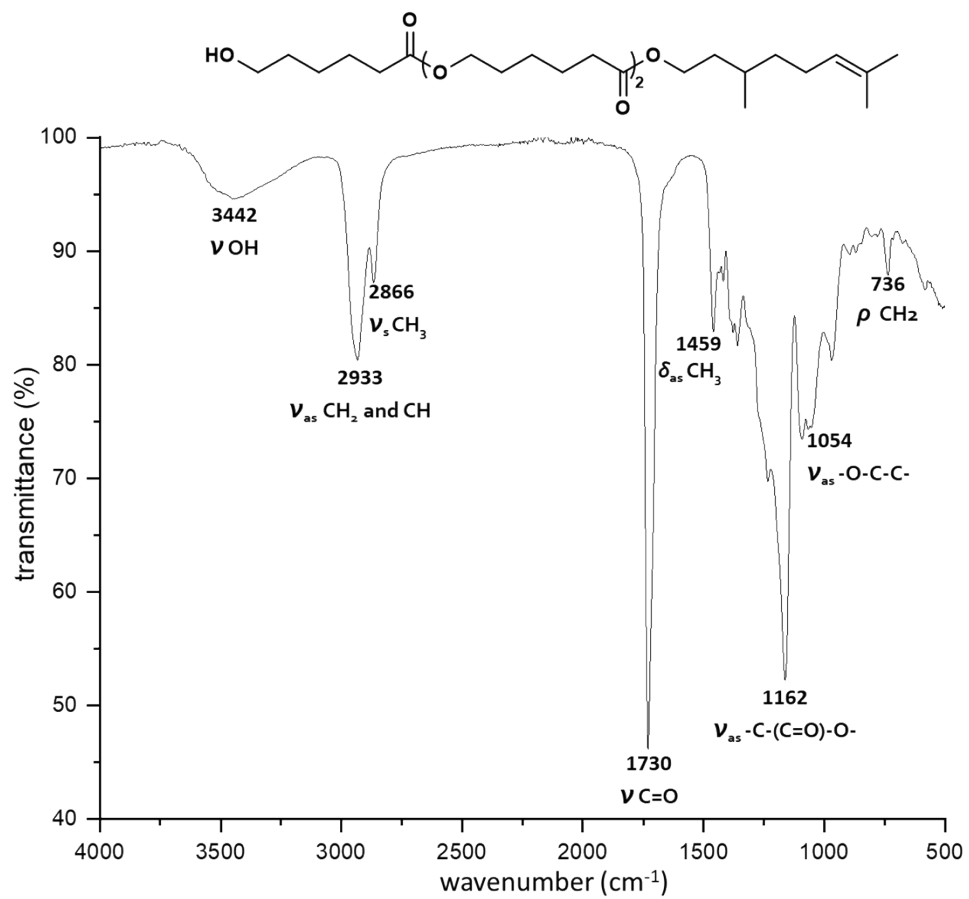
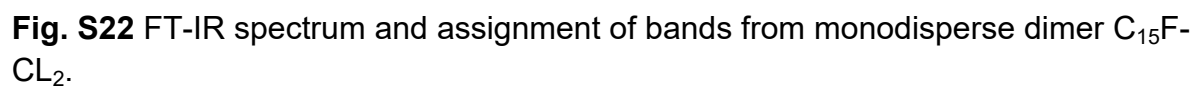
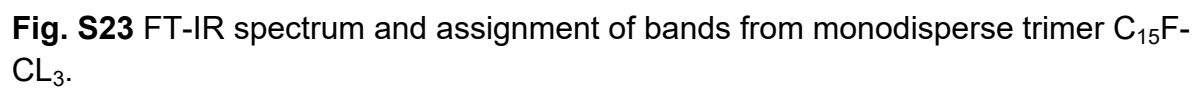


Fig. S21 FT-IR spectrum and assignment of bands from monodisperse trimer $C_{10}C-CL_3$.





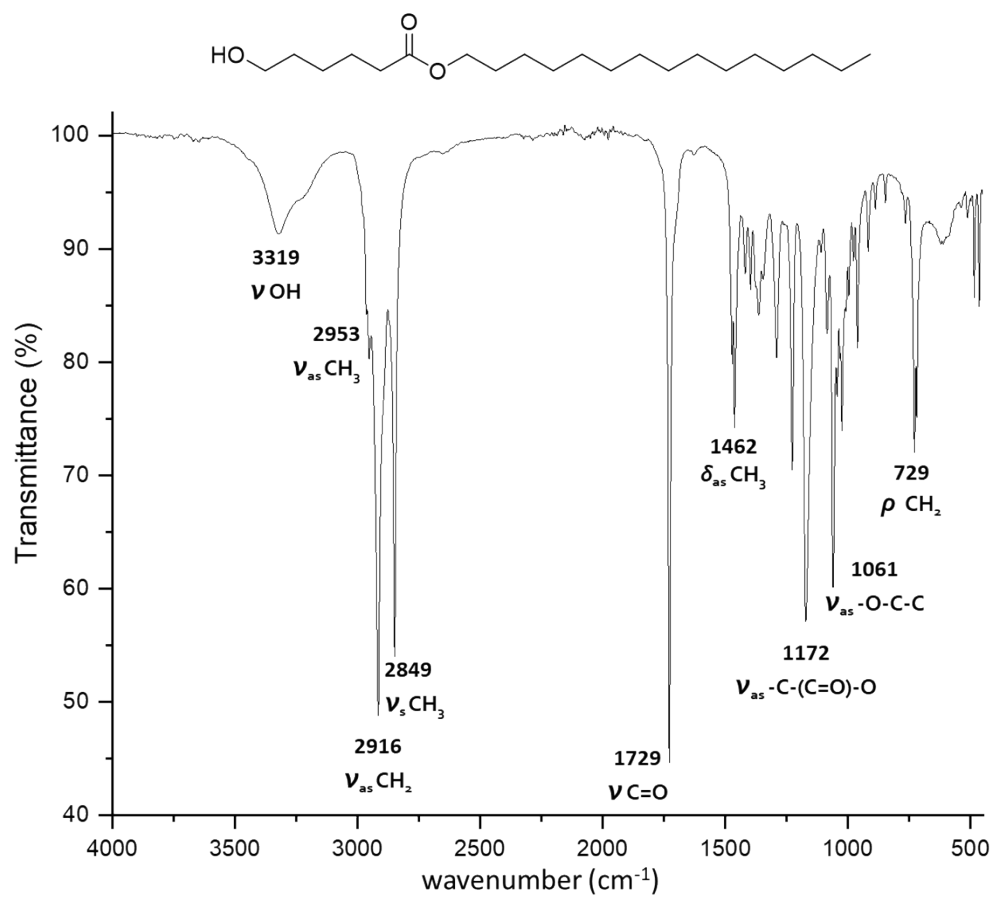
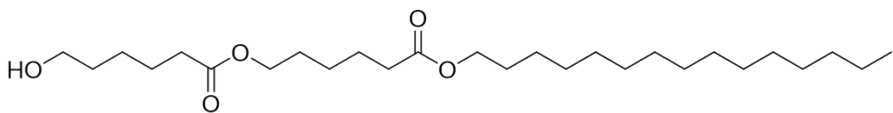


Fig. S24 FT-IR spectrum and assignment of bands from monodisperse monomer C₁₅1P-CL₁.



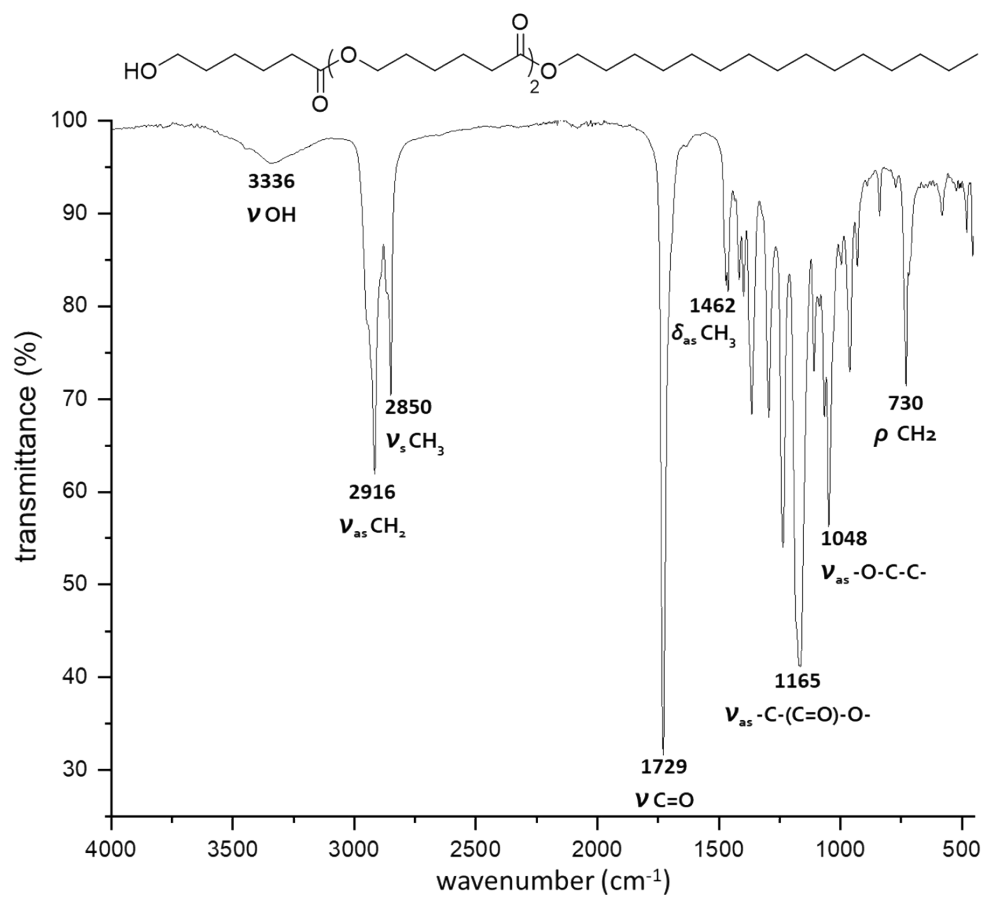


Fig. S26 FT-IR spectrum and assignment of bands from monodisperse trimer $C_{15}1P-CL_3$.

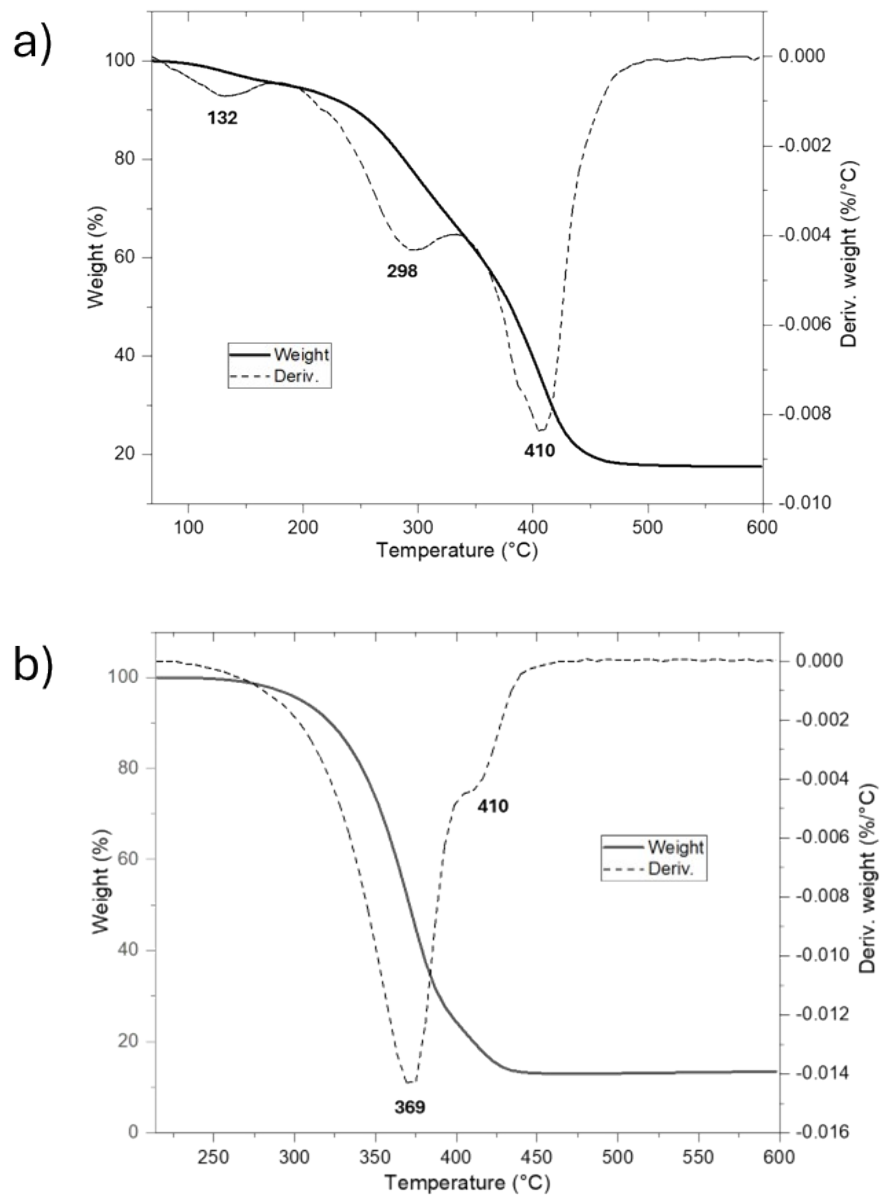


Fig. S27 Thermal degradation (TGA) of a) C₁₅F-CL₂ and b) C₁₅P-CL₂.