

Supporting Information

Star Polymerization of Norbornene Derivatives Using Tri-functionalized Blechert's Olefin Metathesis Catalyst

Chulu Zhou^{[a],[b]}, Cuiping Hou^{[a],[c]} and Jianhua Cheng^{*[a],[b]}

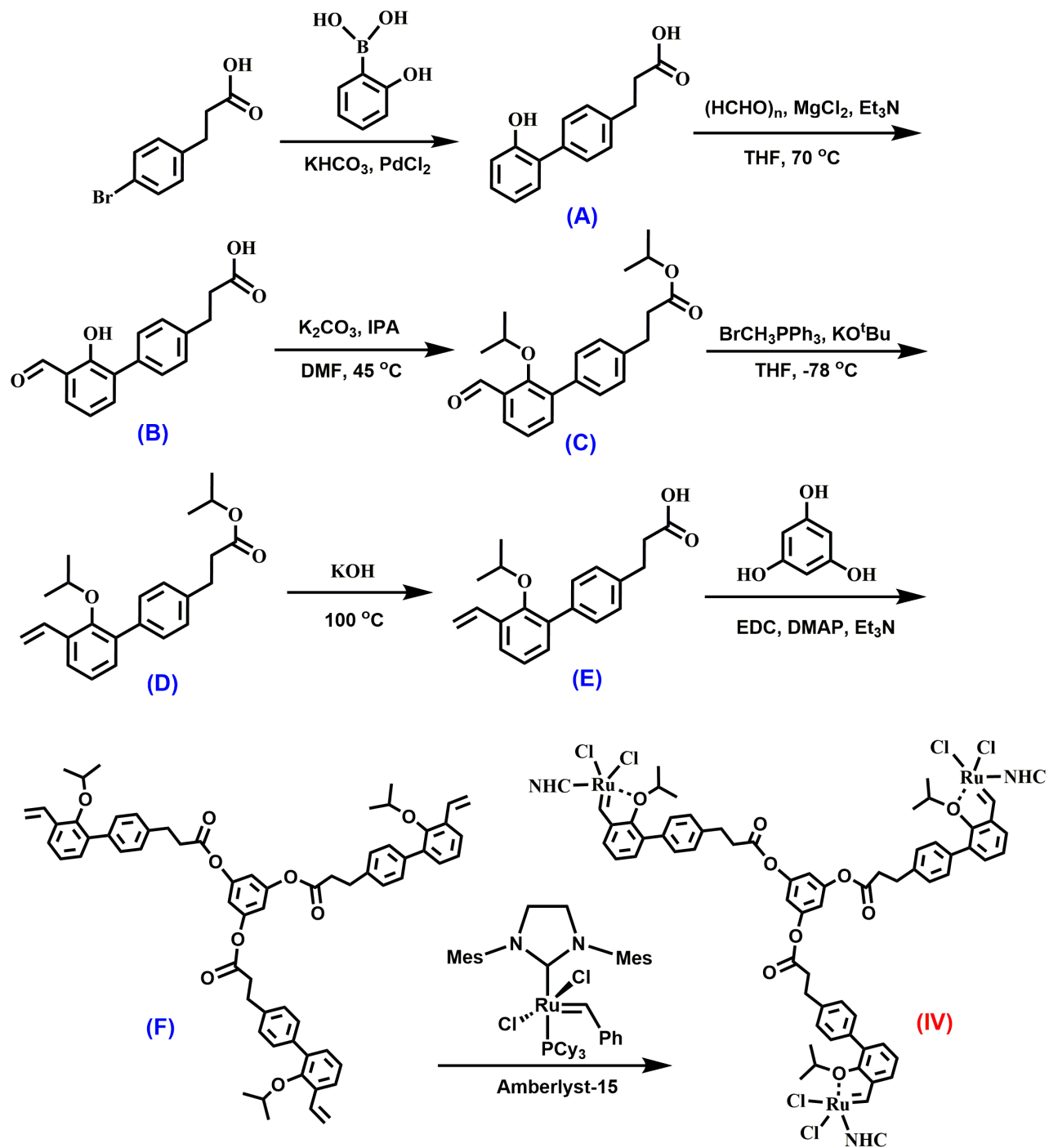
[a] State Key Laboratory of Polymer Physics and Chemistry
Changchun Institute of Applied Chemistry
Chinese Academy of Sciences
No. 5625, Renmin Street, Changchun 130022 (China)
E-Mail: jhcheng@ciac.ac.cn

[b] University of Science and Technology of China
Hefei, Anhui 230029 (China)

[c] University of the Chinese Academy of Sciences,
Changchun Branch, Changchun 130022 (China)

Contents

	Page
1. The route to the synthesis of catalyst IV	S3
2. Selected NMR spectra	S4
3. MALDI-TOF spectra of complex F	S11
4. GPC and NMR characterization of polymers	S12



Scheme S1. The route to the synthesis of star-type trinuclear Ru catalyst (IV).

2. ^1H and $^{13}\text{C}\{^1\text{H}\}$ NMR spectra of complexes A-F and IV

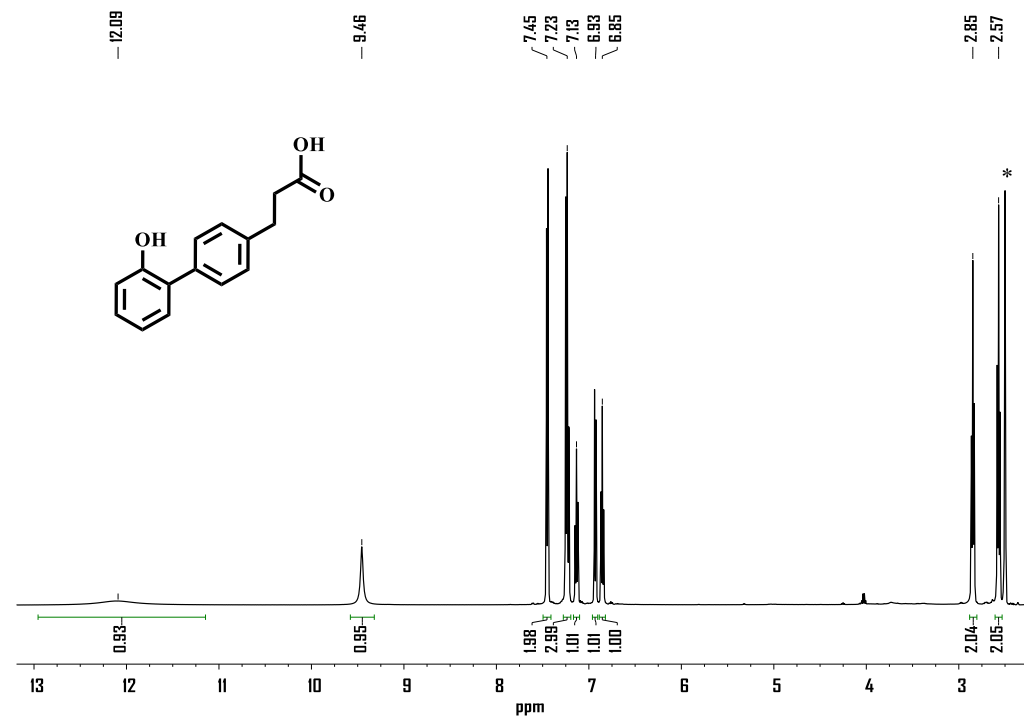


Figure S1. ^1H NMR spectrum (500 MHz) of complex A in $\text{DMSO-}d_6$ (*) at 25 °C.

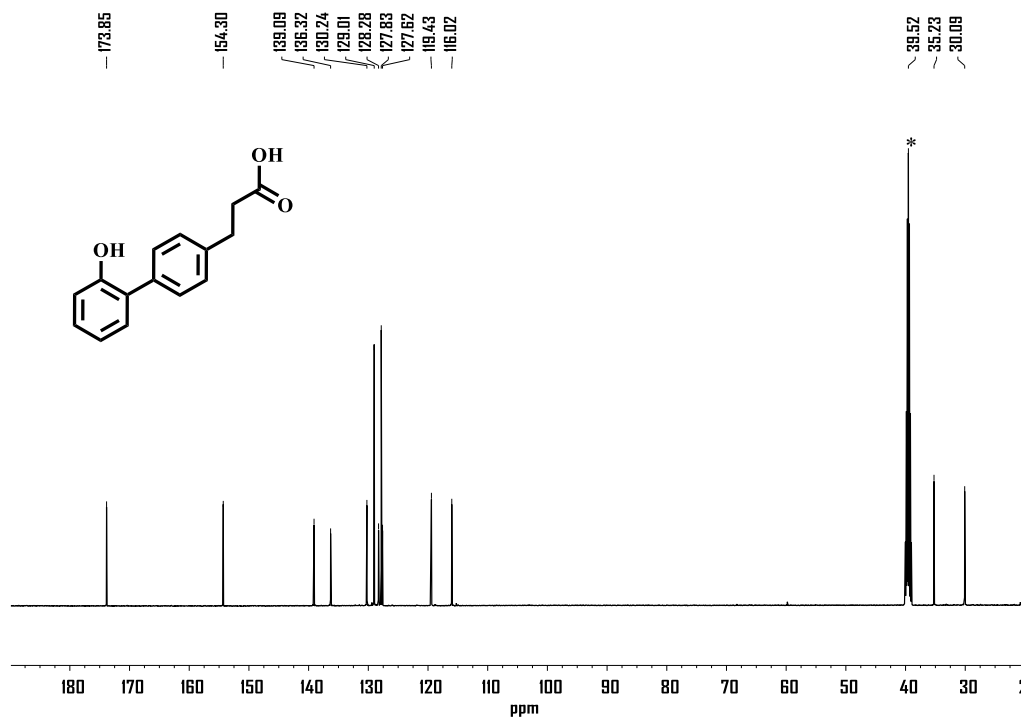


Figure S2. $^{13}\text{C}\{^1\text{H}\}$ NMR spectrum (125 MHz) of complex A in $\text{DMSO-}d_6$ (*) at 25 °C.

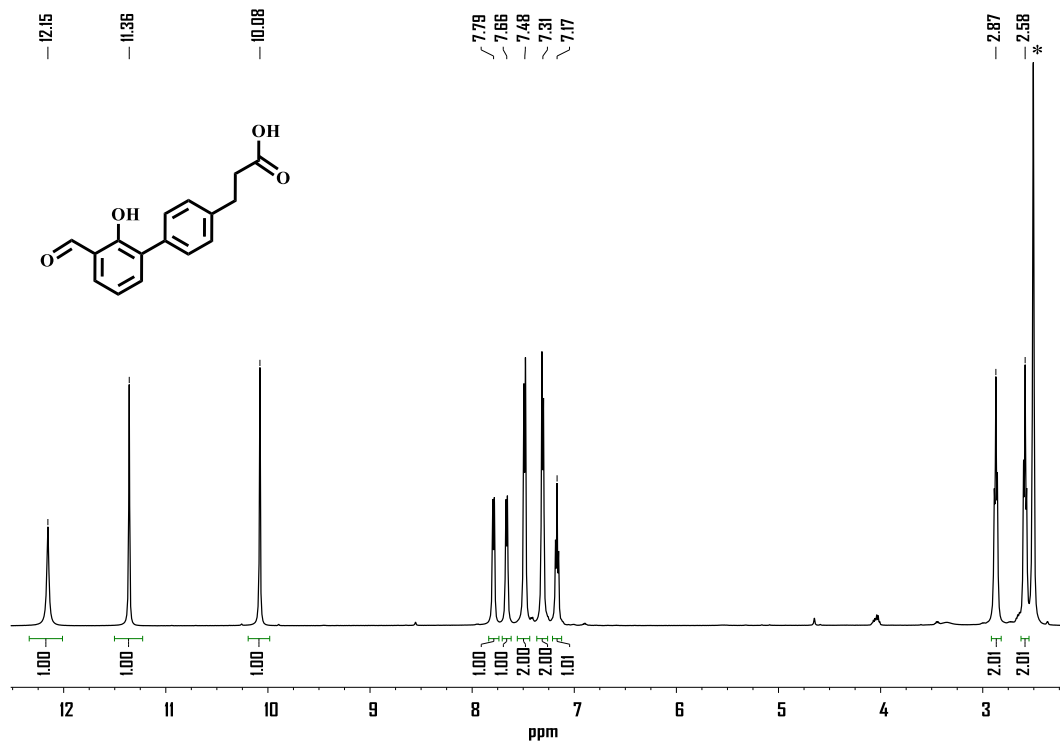


Figure S3. ¹H NMR spectrum (500 MHz) of complex **B** in DMSO-*d*₆(*) at 25 °C.

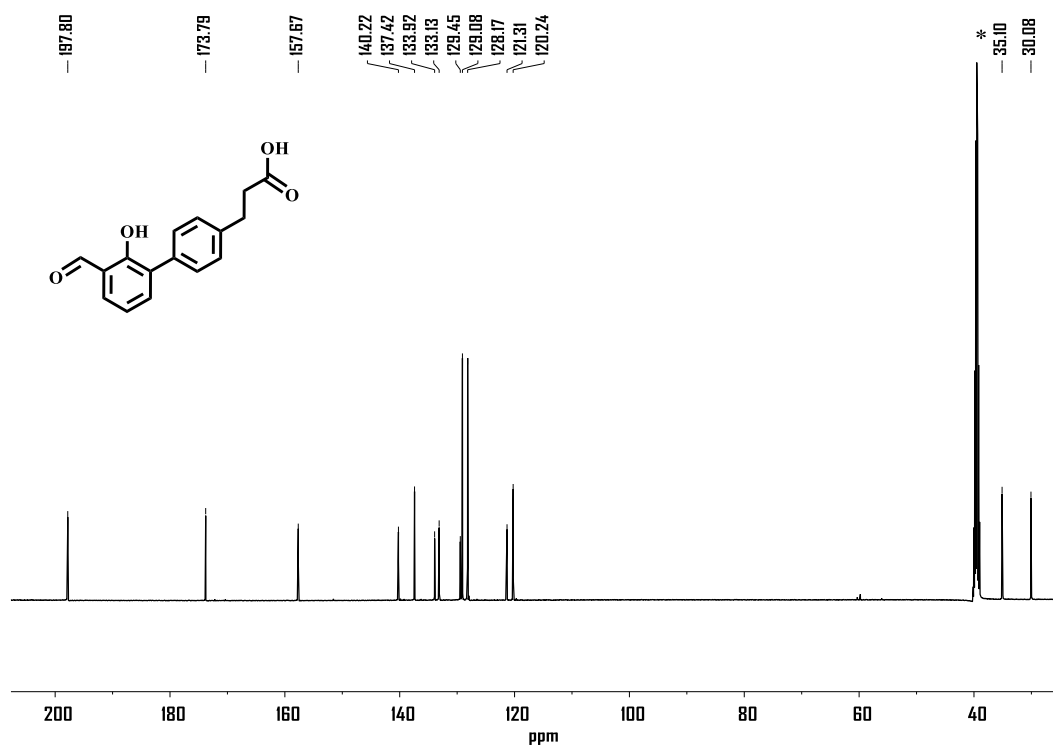


Figure S4. ¹³C{¹H} NMR spectrum (125 MHz) of complex **B** in DMSO-*d*₆(*) at 25 °C.

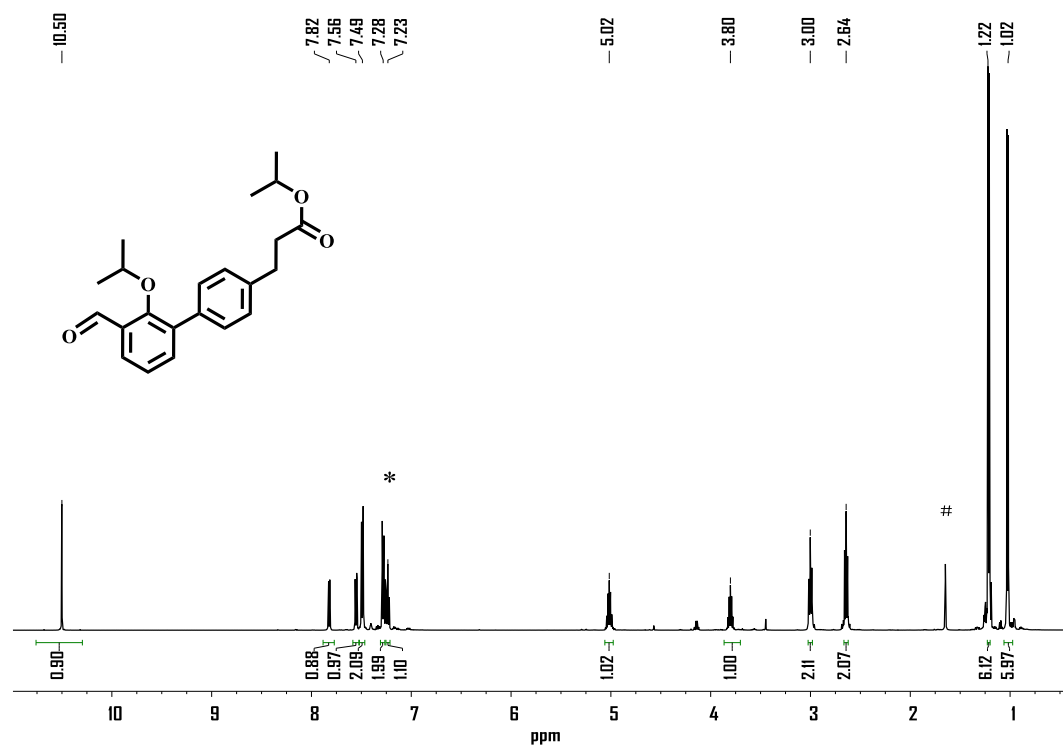


Figure S5. ¹H NMR spectrum (500 MHz) of complex C in CDCl₃ (*) at 25 °C. (#H₂O)

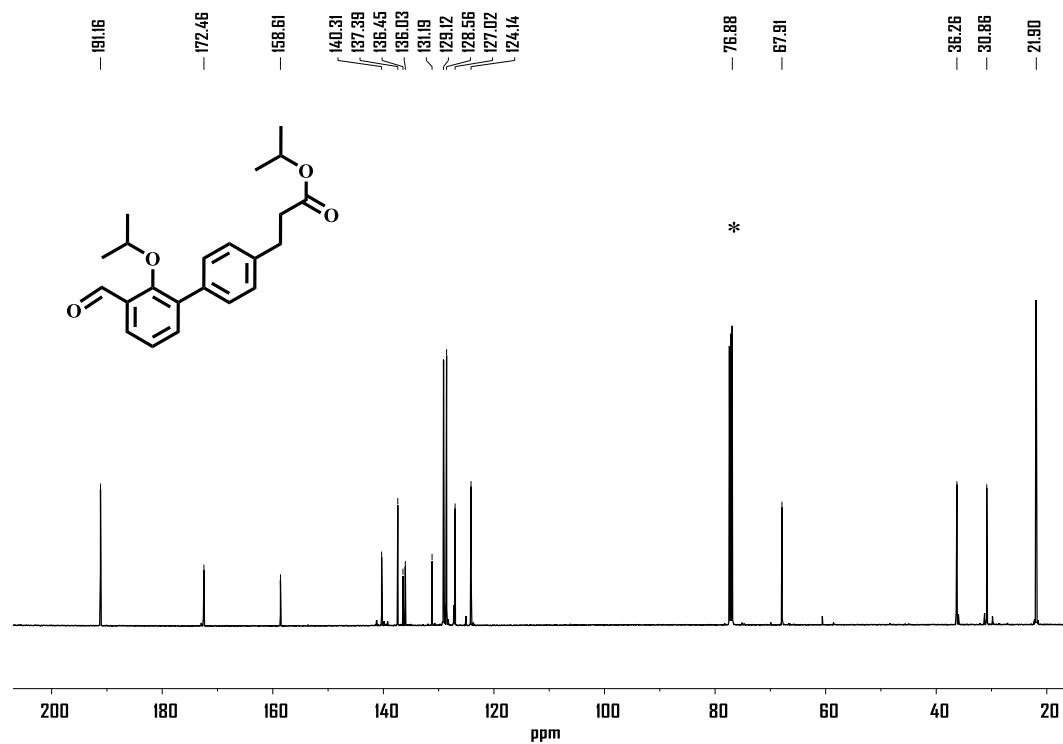


Figure S6. ¹³C{¹H} NMR spectrum (125 MHz) of complex C in CDCl₃ (*) at 25 °C.

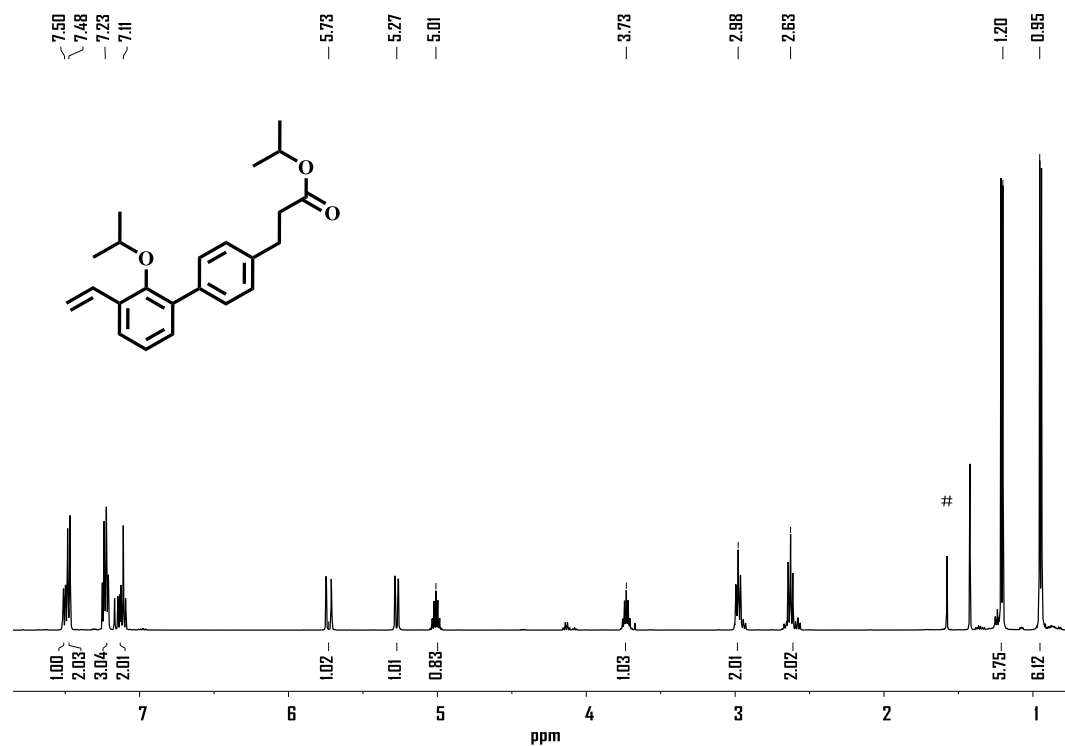


Figure S7. ^1H NMR spectrum (500 MHz) of complex **D** in CDCl_3 (*) at 25 °C. (# H_2O)

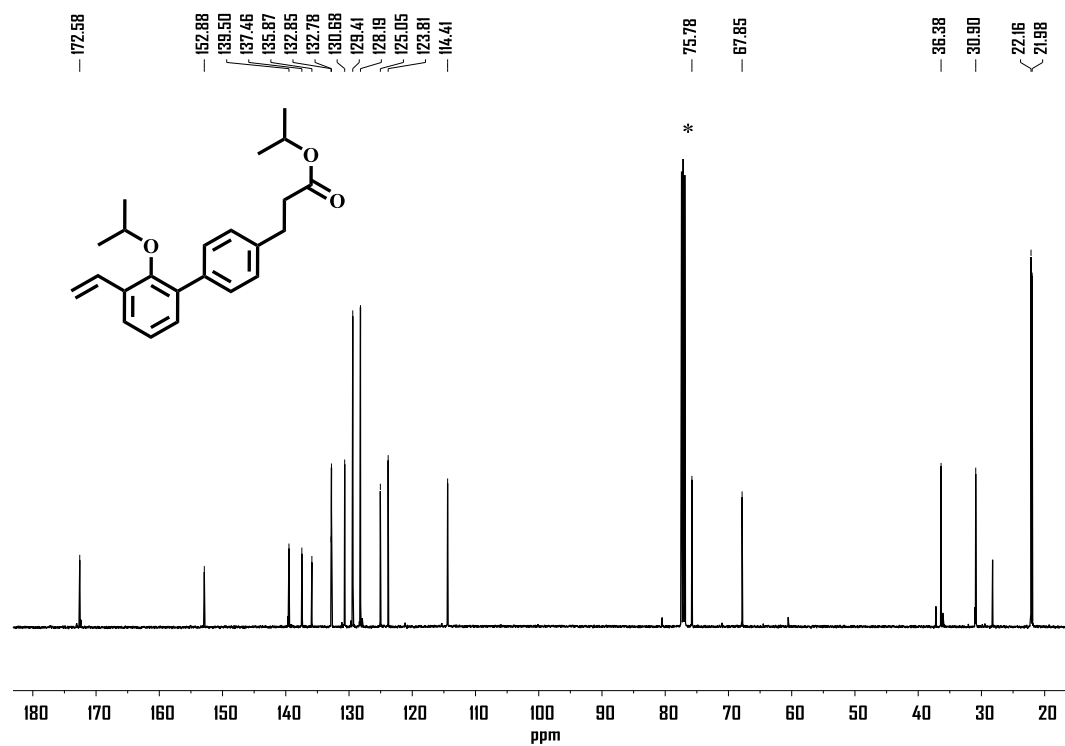


Figure S8. $^{13}\text{C}\{^1\text{H}\}$ NMR spectrum (125 MHz) of complex **D** in CDCl_3 (*) at 25 °C.

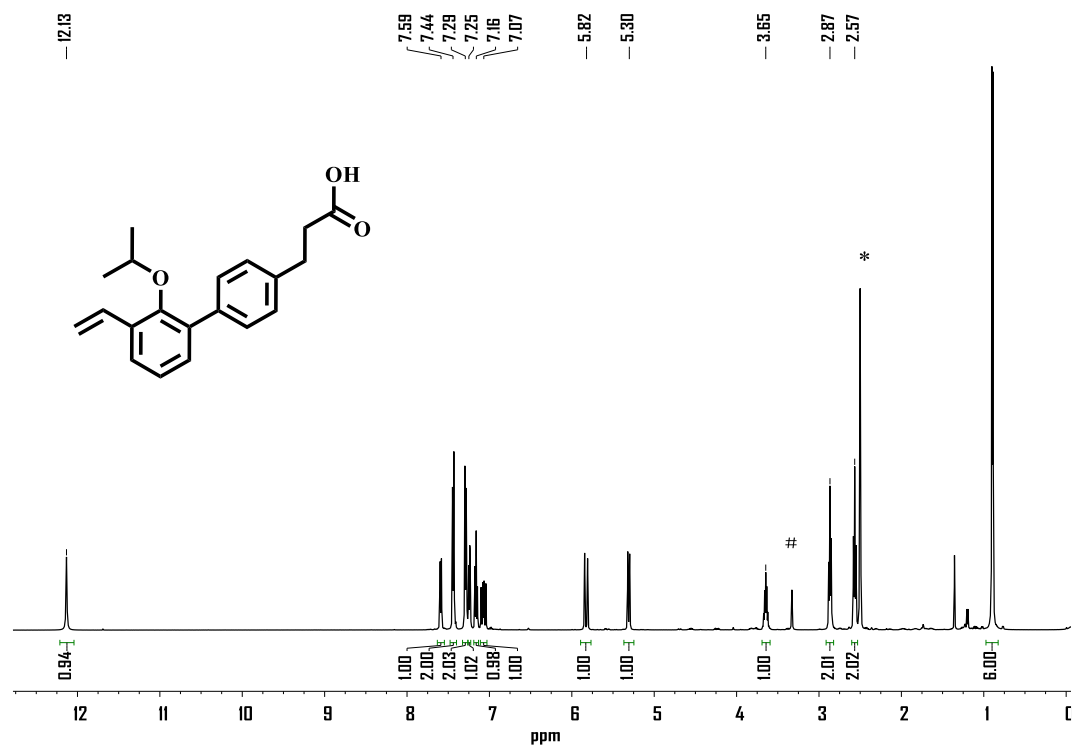


Figure S9. ¹H NMR spectrum (500 MHz) of complex E in DMSO-*d*₆ (*) at 25 °C. (*H₂O)

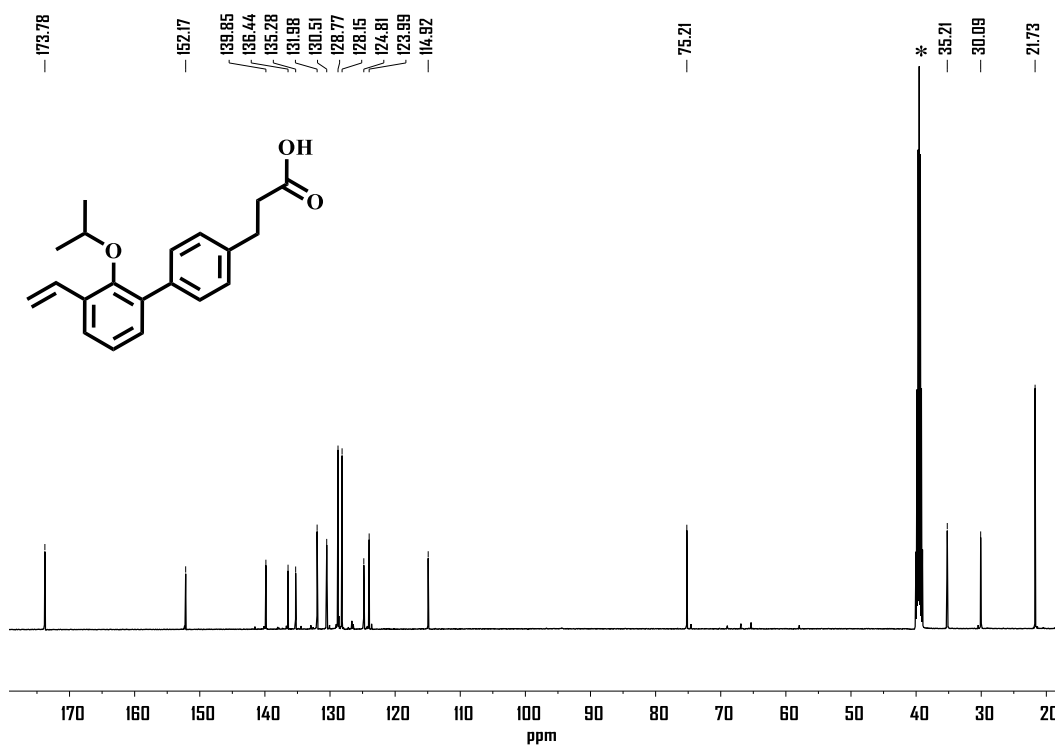


Figure S10. ¹³C{¹H} NMR spectrum (125 MHz) of complex E in DMSO-*d*₆ (*) at 25 °C.

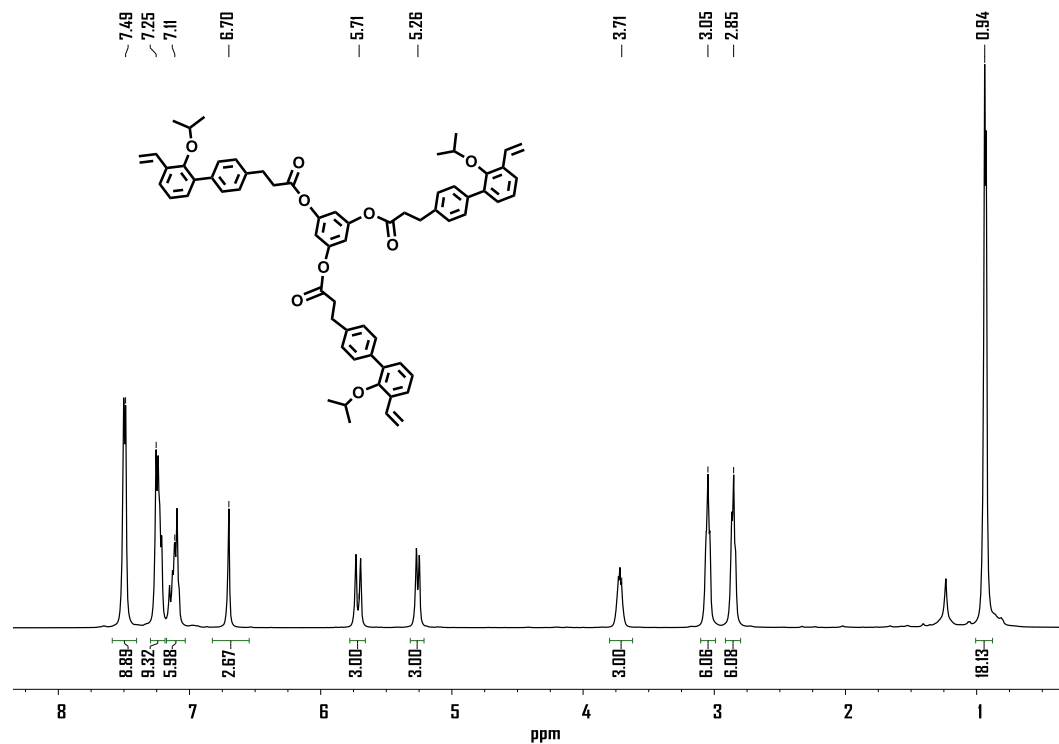


Figure S11. ^1H NMR spectrum (500 MHz) of complex **F** in CDCl_3 (*) at 25 °C.

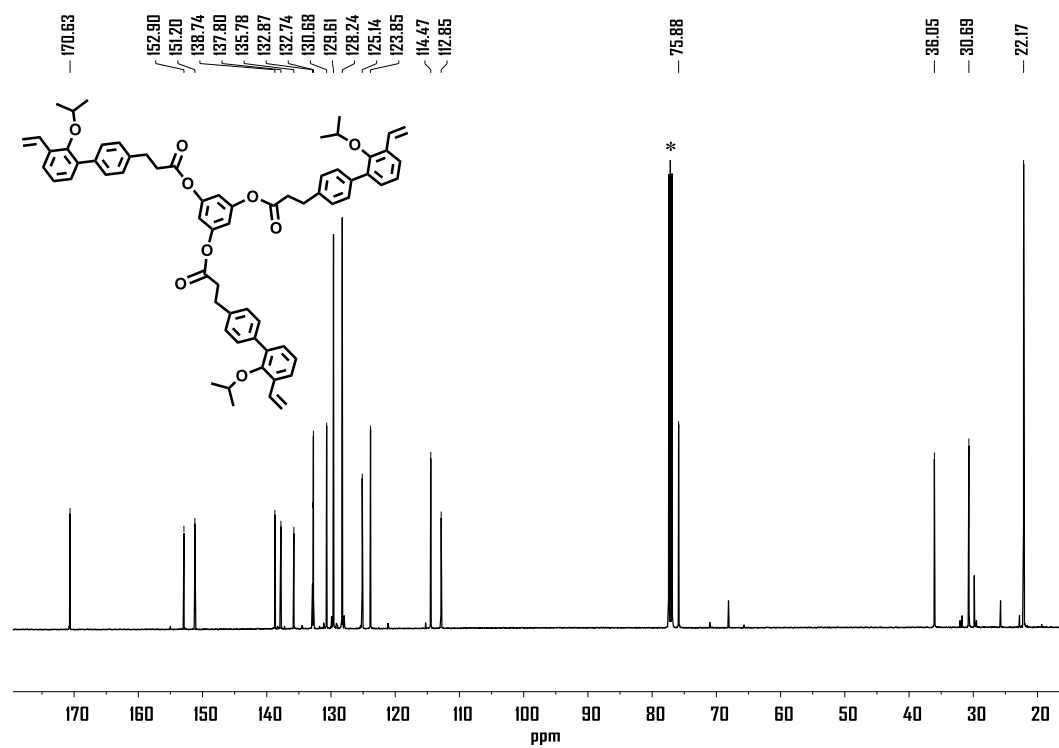


Figure S12. $^{13}\text{C}\{^1\text{H}\}$ NMR spectrum (125 MHz) of complex **F** in CDCl_3 (*) at 25 °C.

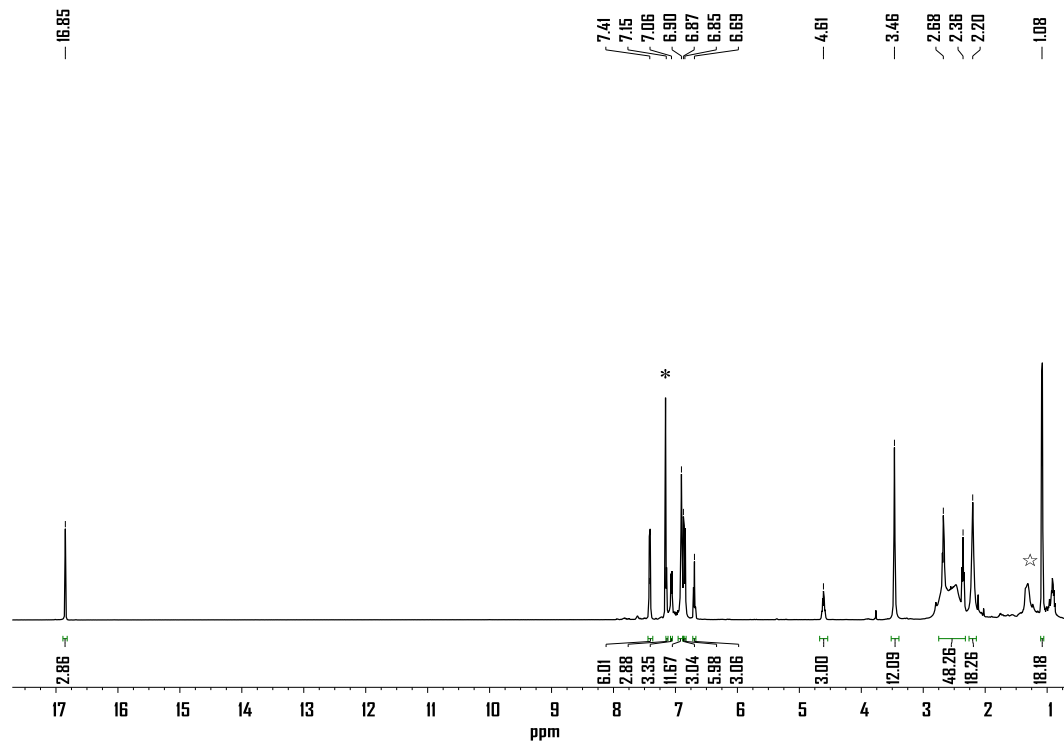


Figure S13. ^1H NMR spectrum (500 MHz) of complex **IV** in C_6D_6 (*) at 25 °C. (* Hexane).

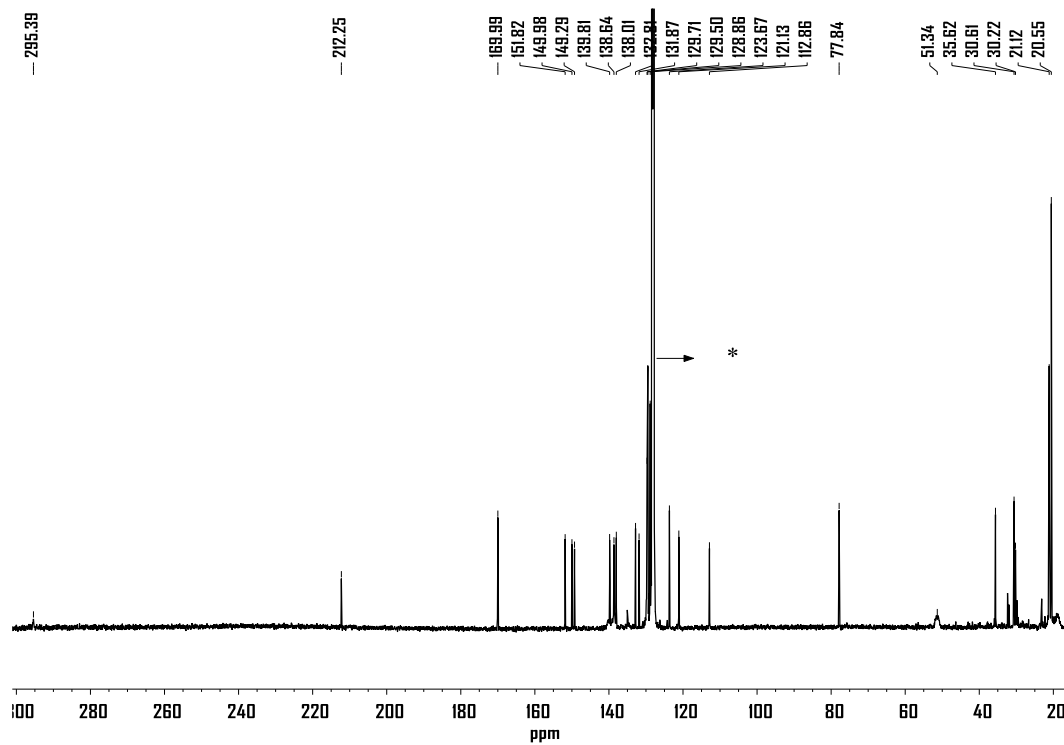


Figure S14. $^{13}\text{C}\{^1\text{H}\}$ NMR spectrum (125 MHz) of complex **IV** in C_6D_6 (*) at 25 °C.

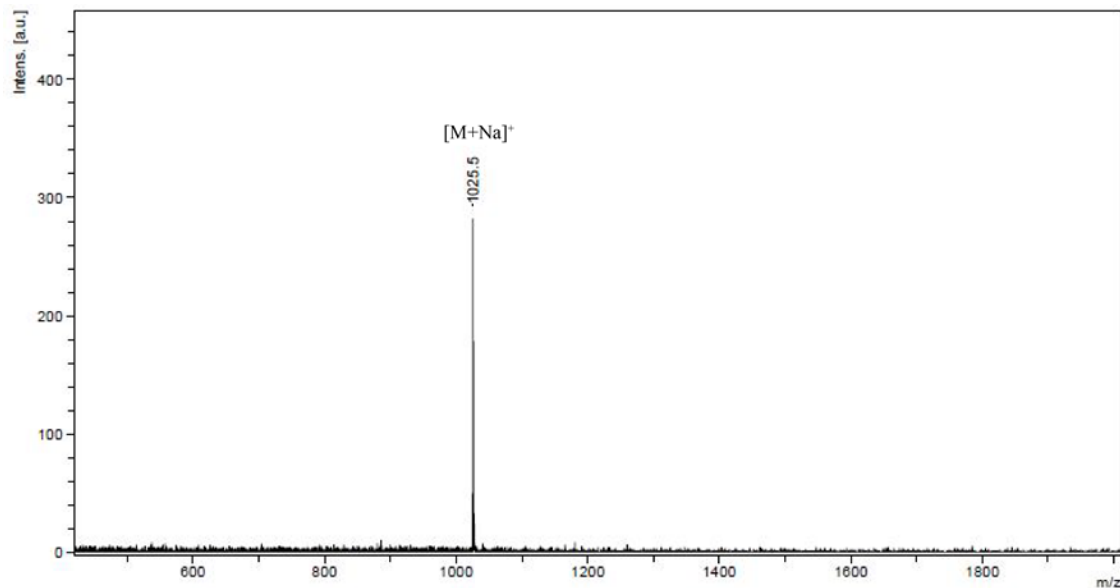


Figure S15. MALDI-TOF of complex F.

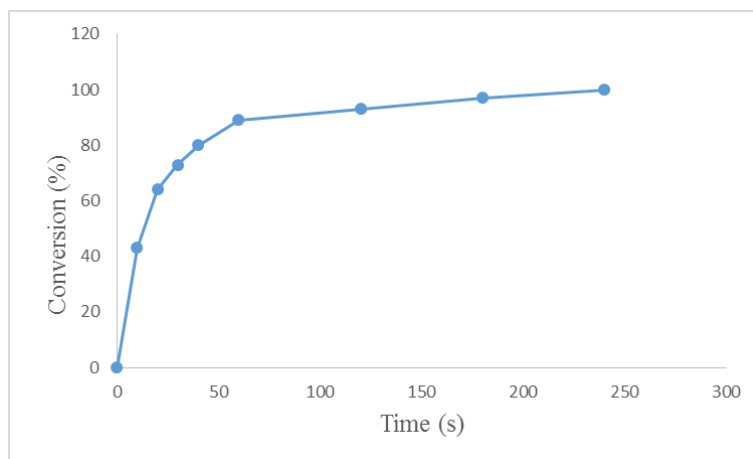


Figure S16. ROMP of monomer 1 in the presence of catalysts III. Conditions: Monomer/catalyst ratio 100:1, 0.3 mM (catalyst concentration) in toluene at room temperature. Conversion was determined by ^1H NMR.

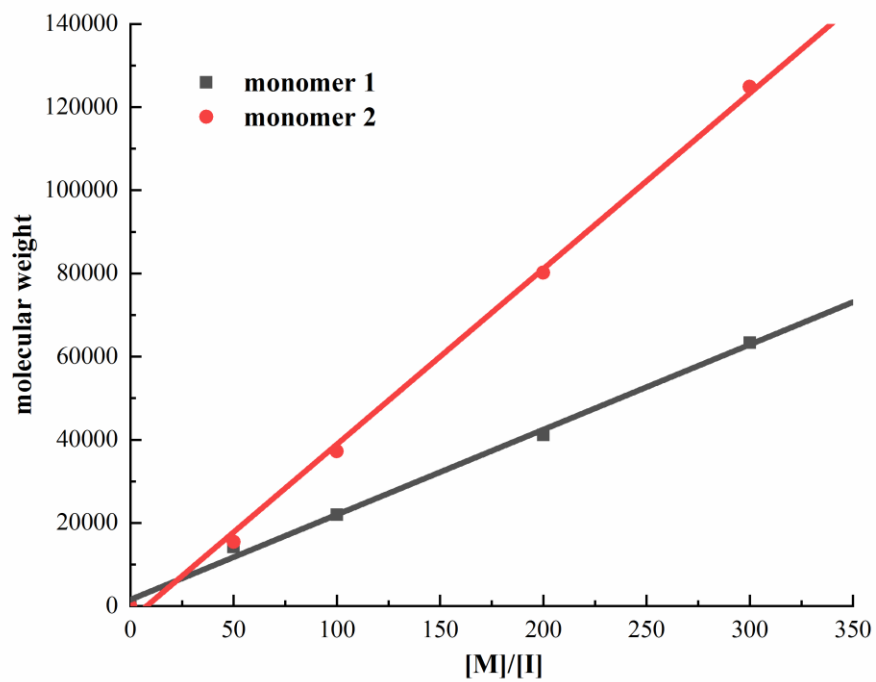


Figure S17. M_n versus M/I for the polymers produced from monomer **1** and **2** by using catalyst III.

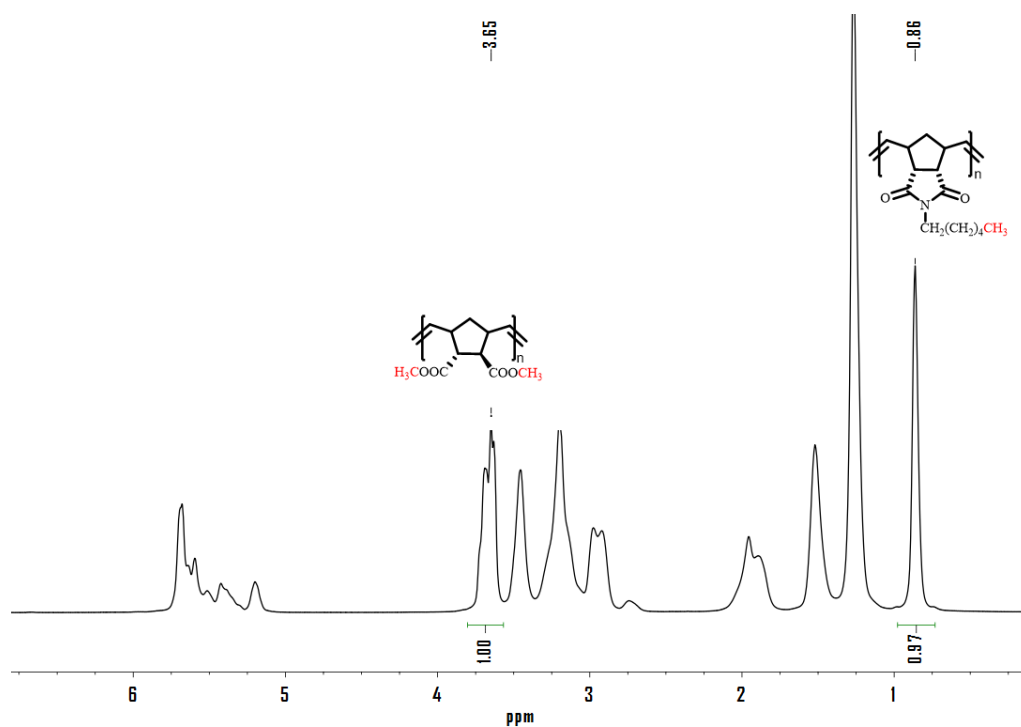


Figure S18. ^1H NMR spectrum (500 MHz) of poly[(**1**)₁₀₀-*b*-(**4**)₂₀₀] in CDCl_3 at 25 °C.

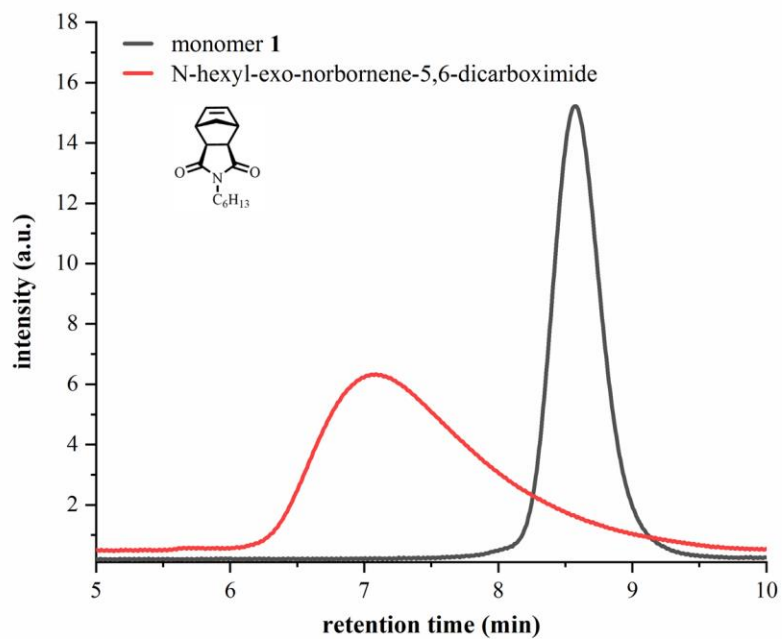


Figure S19. SEC traces of monomer **1** and N-hexyl-exo-norbornene-5, 6-dicarboximide

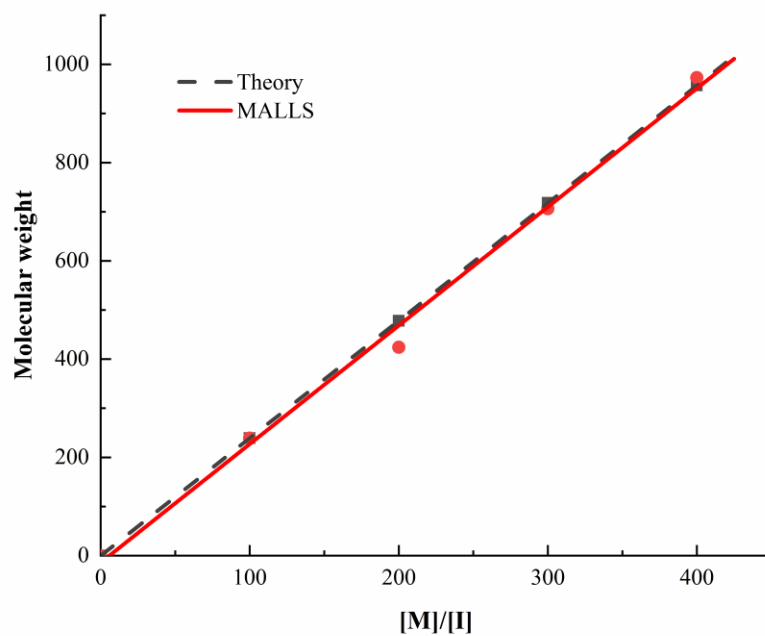


Figure S20. Plots of theoretical molecular weight and absolute molecular weight determined by MALLS-RI detector versus $[M]/[I]$ for dendrimer **5** and using **IV** as catalyst.