

Synthesis and Characterization of a Oligomeric Conjugated Metal-Containing Poly(*p*-phenylenevinylene) Analogue

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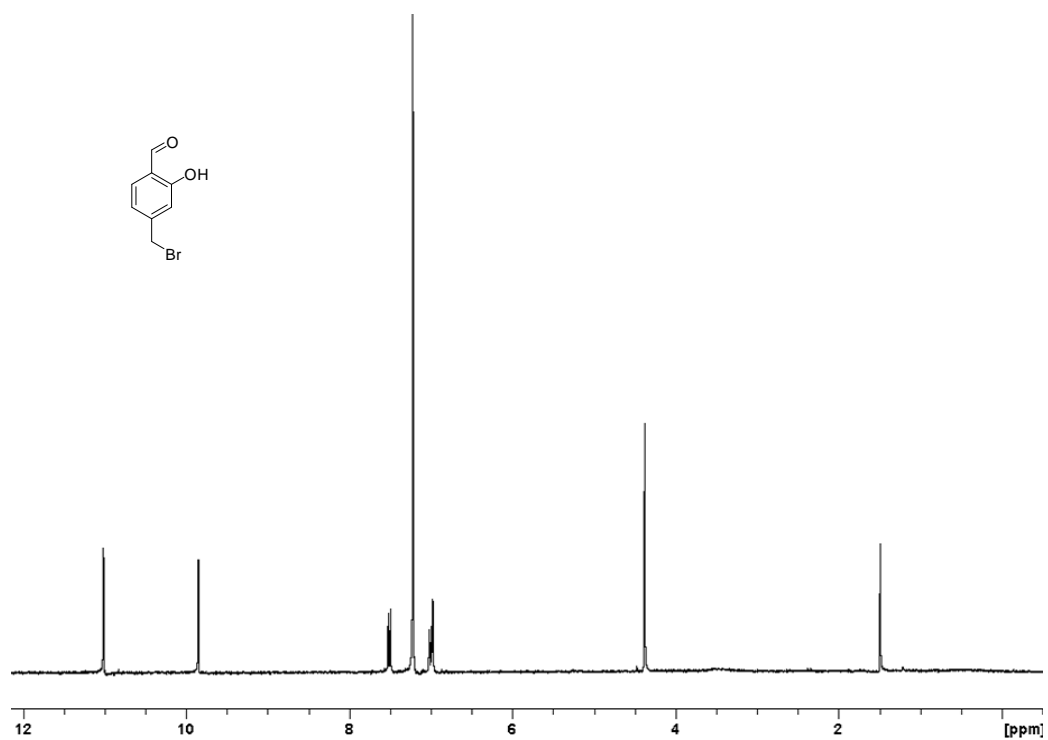


Figure S1. ¹H NMR spectrum of **6** (300 MHz, CDCl₃, 298 K).

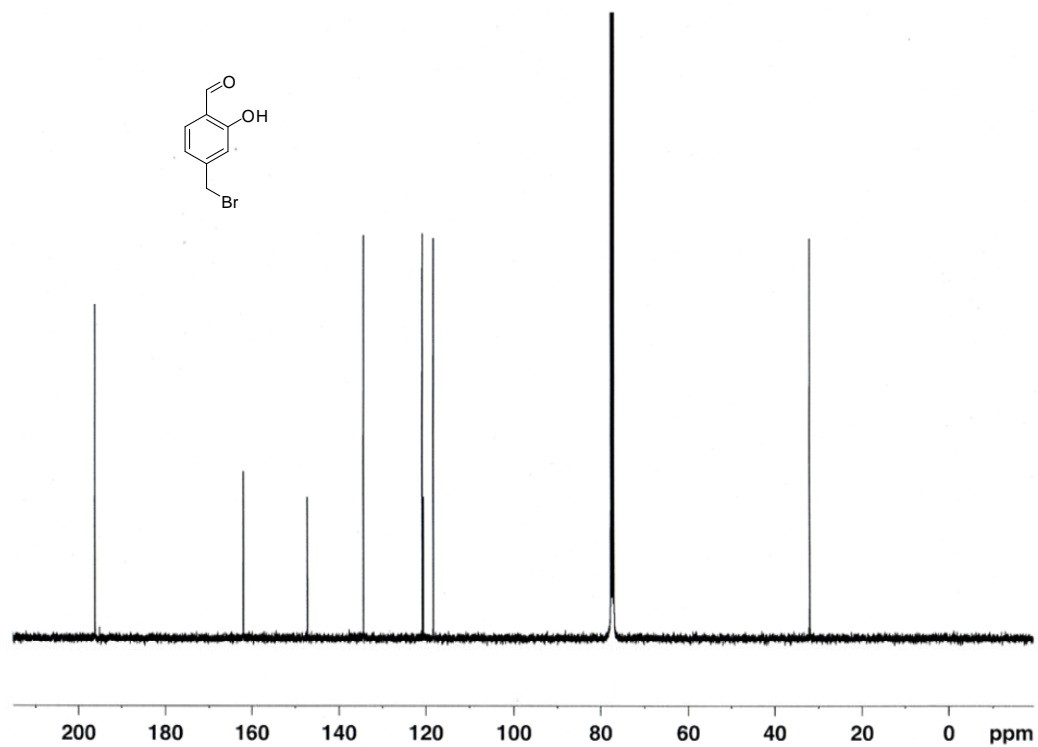


Figure S2. ¹³C NMR spectrum of **6** (100.6 MHz, CDCl₃, 298 K).

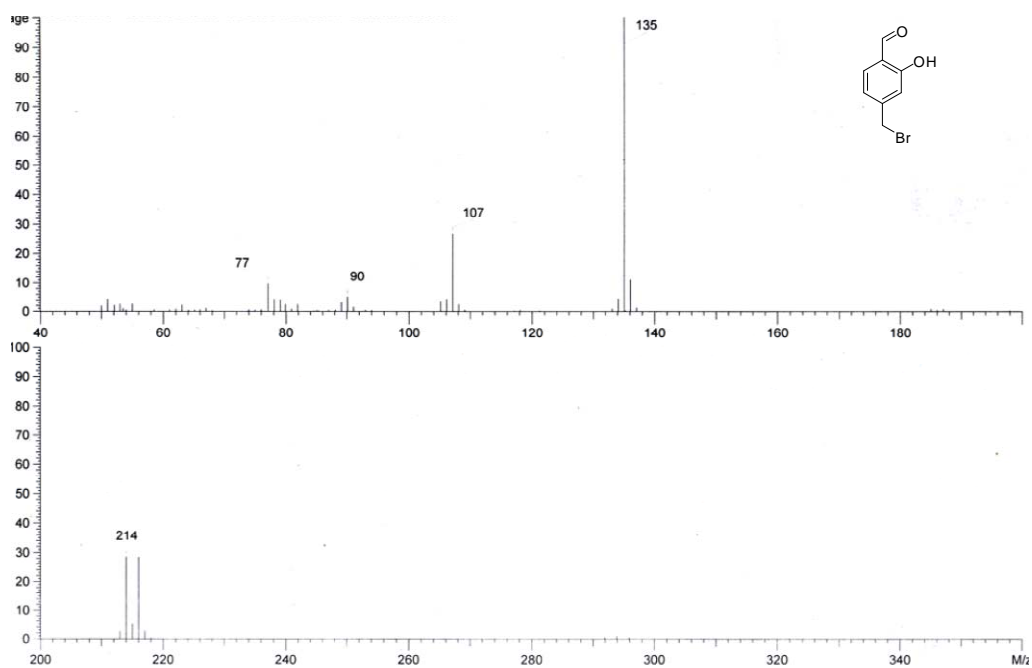


Figure S3. EI-MS spectrum of **6**

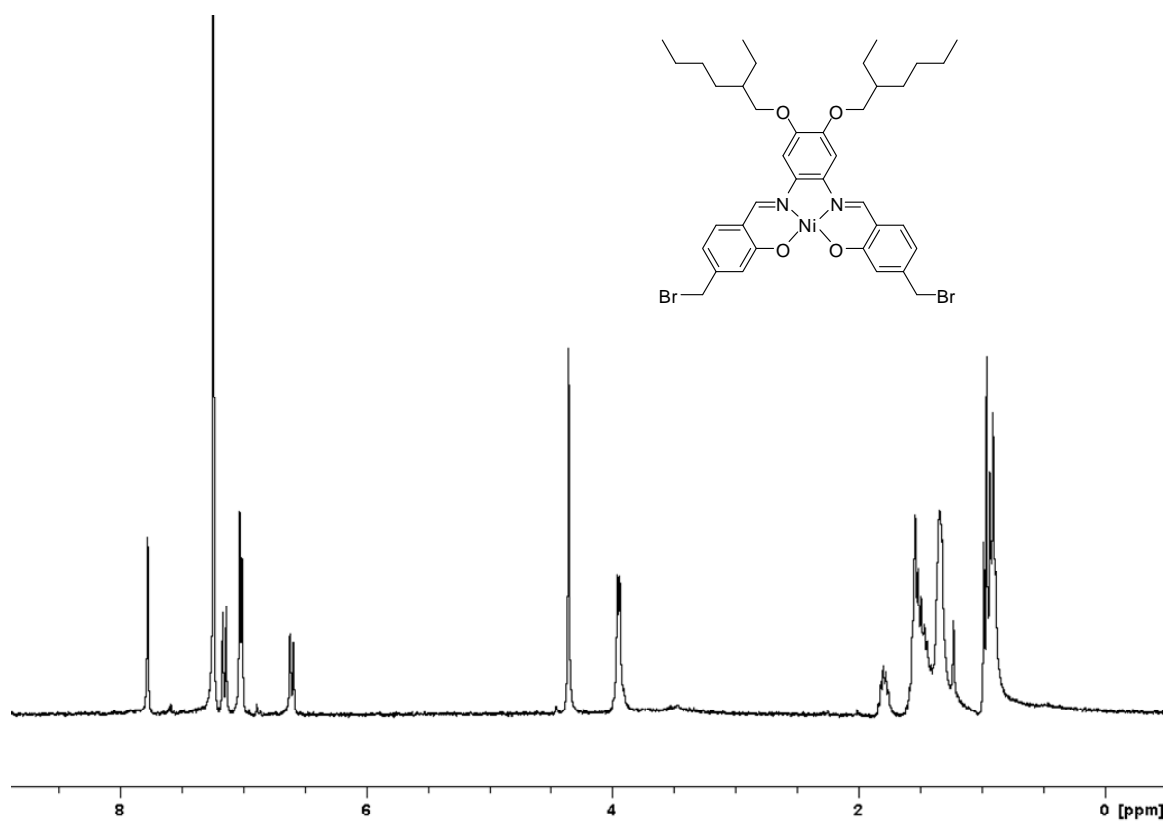


Figure S4. ¹H NMR spectrum of **2** (300 MHz, CDCl₃ (with trace of THF), 298 K)

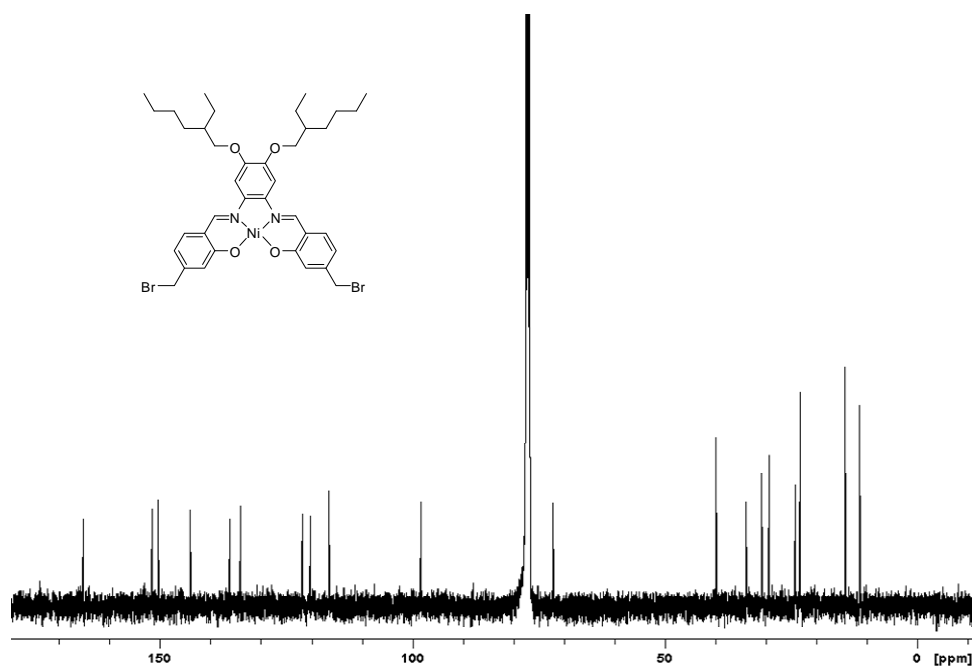


Figure S5. ^{13}C NMR spectrum of **2** (100.6 MHz, CDCl_3 , 298 K).

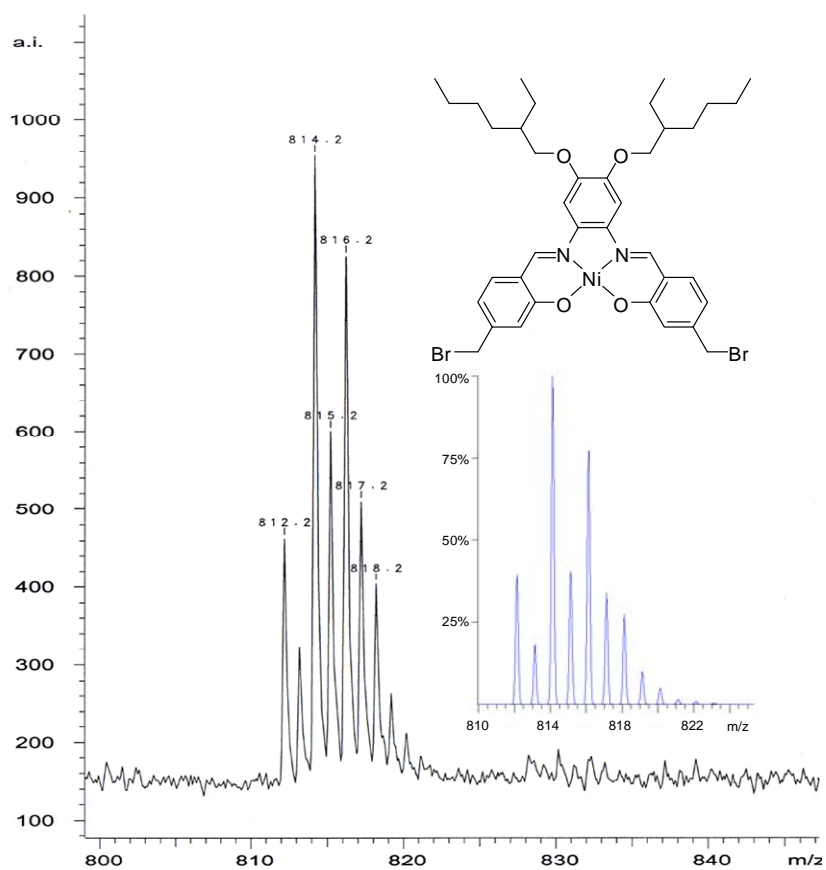


Figure S6. MALDI-TOF mass spectrum of **2**. Inset: Simulated isotope distribution for $[2]^+$

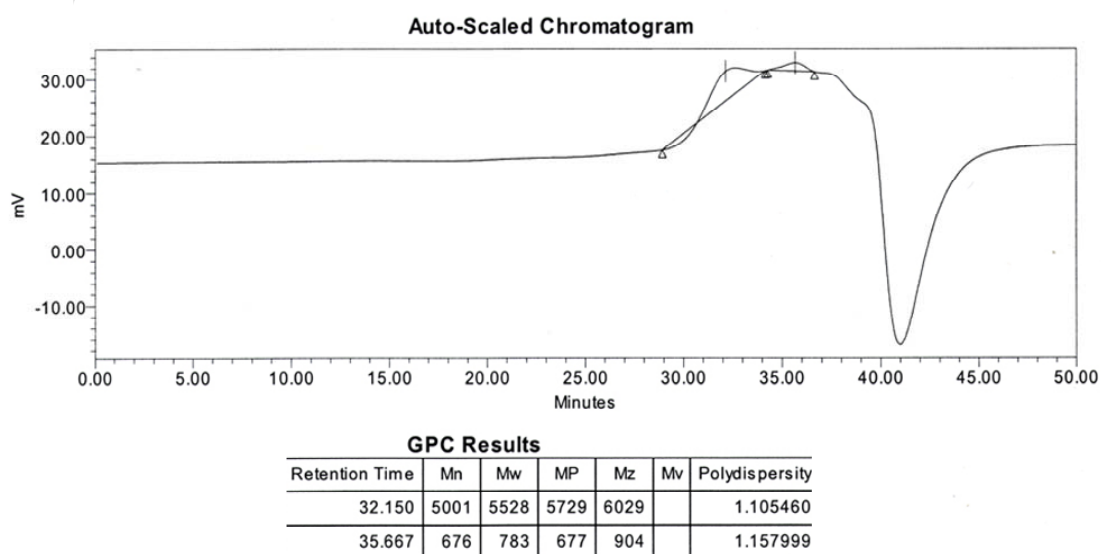


Figure S7. GPC of polymer **8**

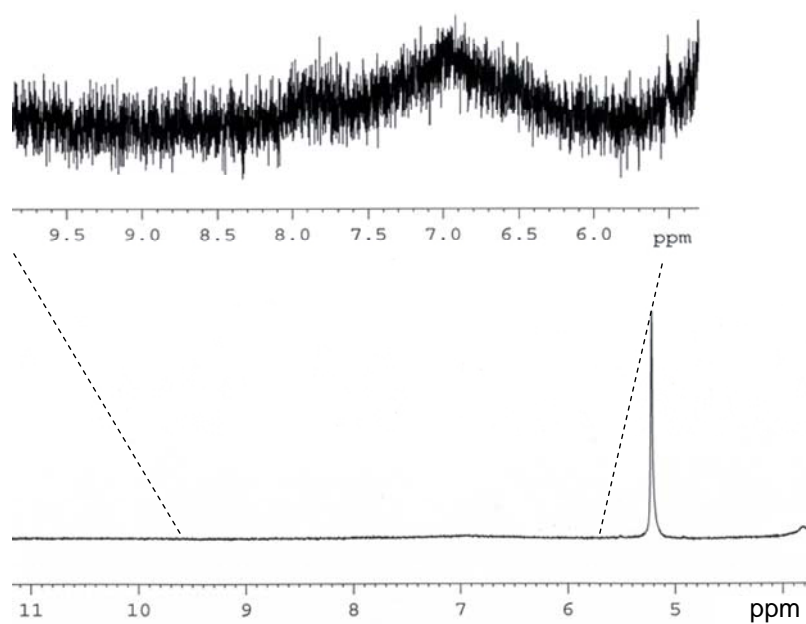


Figure S8. Partial of ^1H NMR spectrum of **8** (400 MHz, CD_2Cl_2 (with a trace of $\text{THF-}d_8$), 298K).

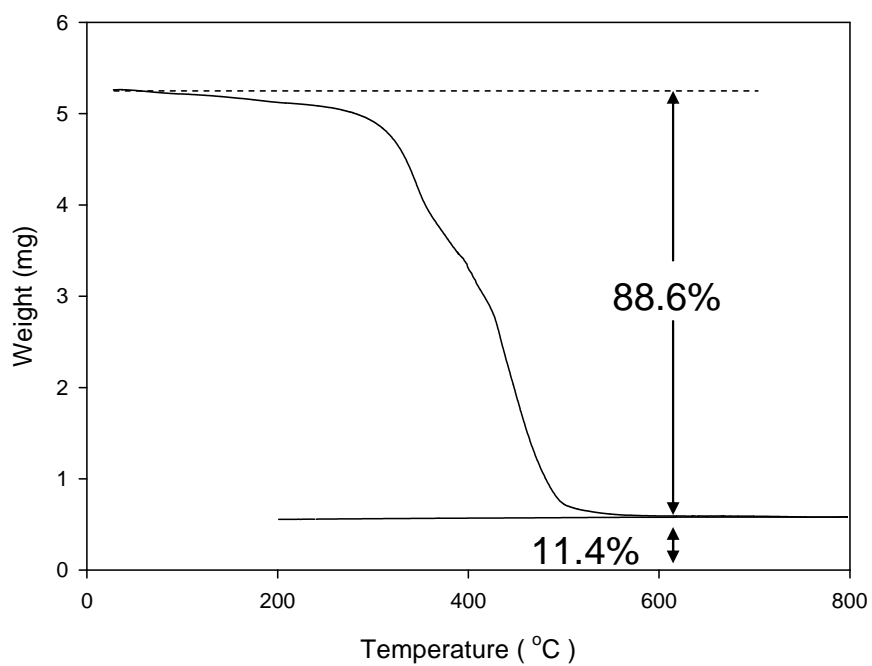


Figure S9. TGA of polymer **8**

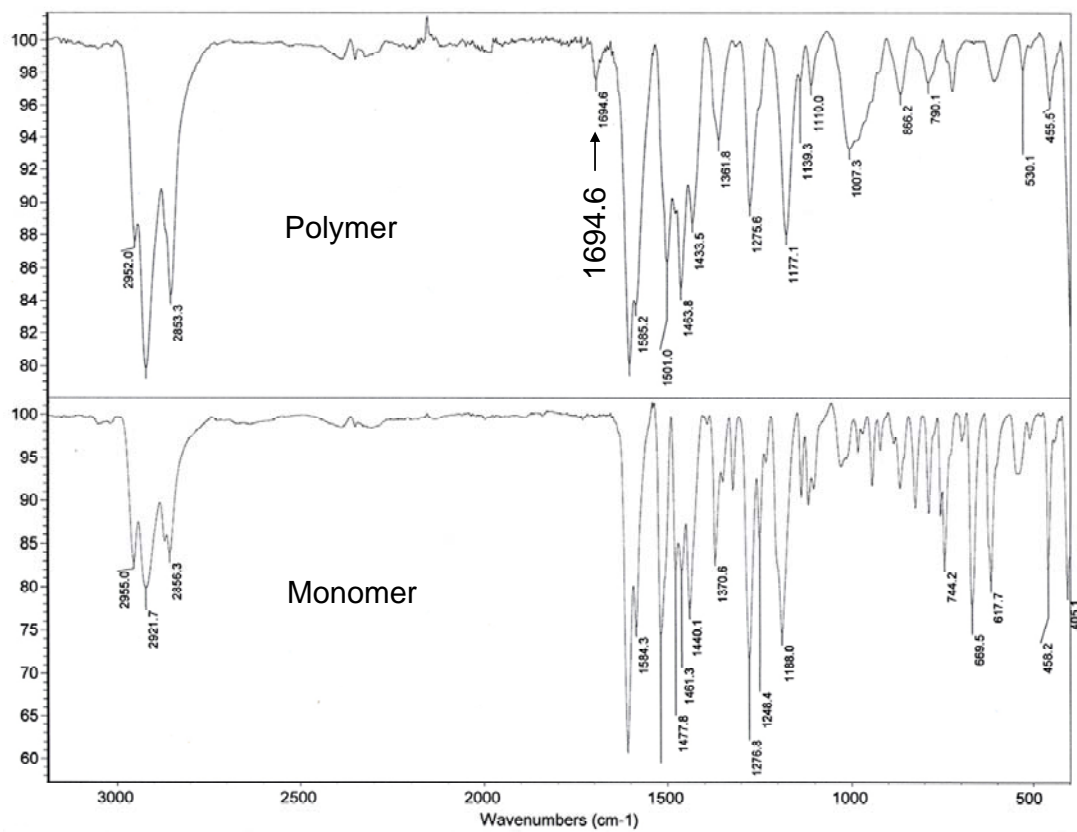


Figure S10. IR spectra of polymer **8** and monomer **2**

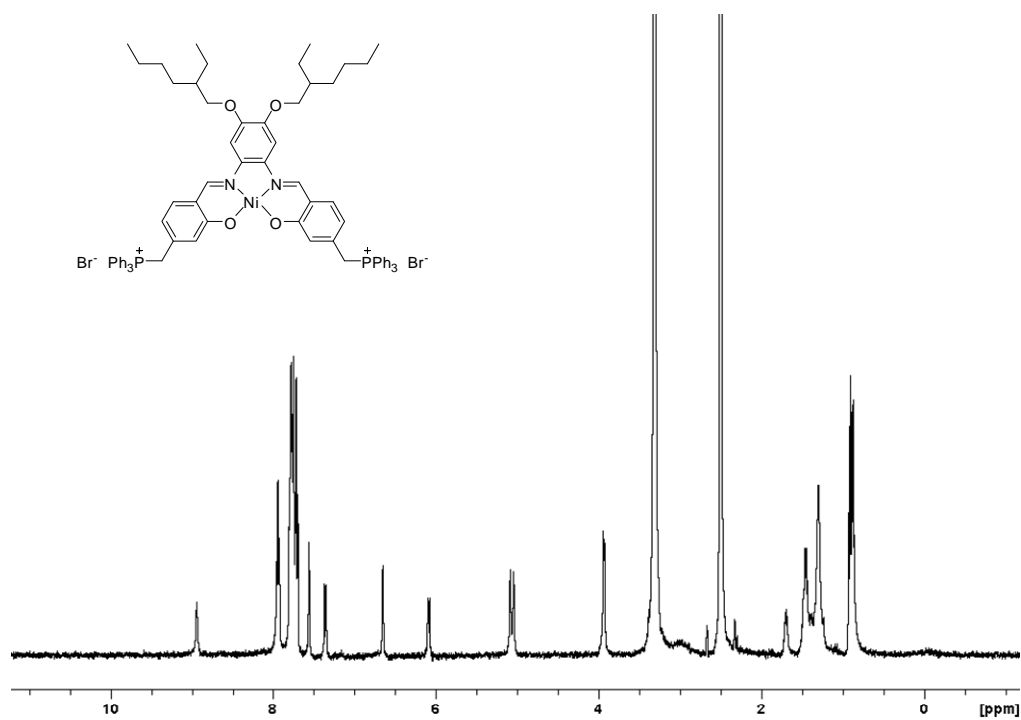


Figure 11. ¹H NMR spectrum of **9** (400 MHz, DMSO-*d*₆, 298 K).

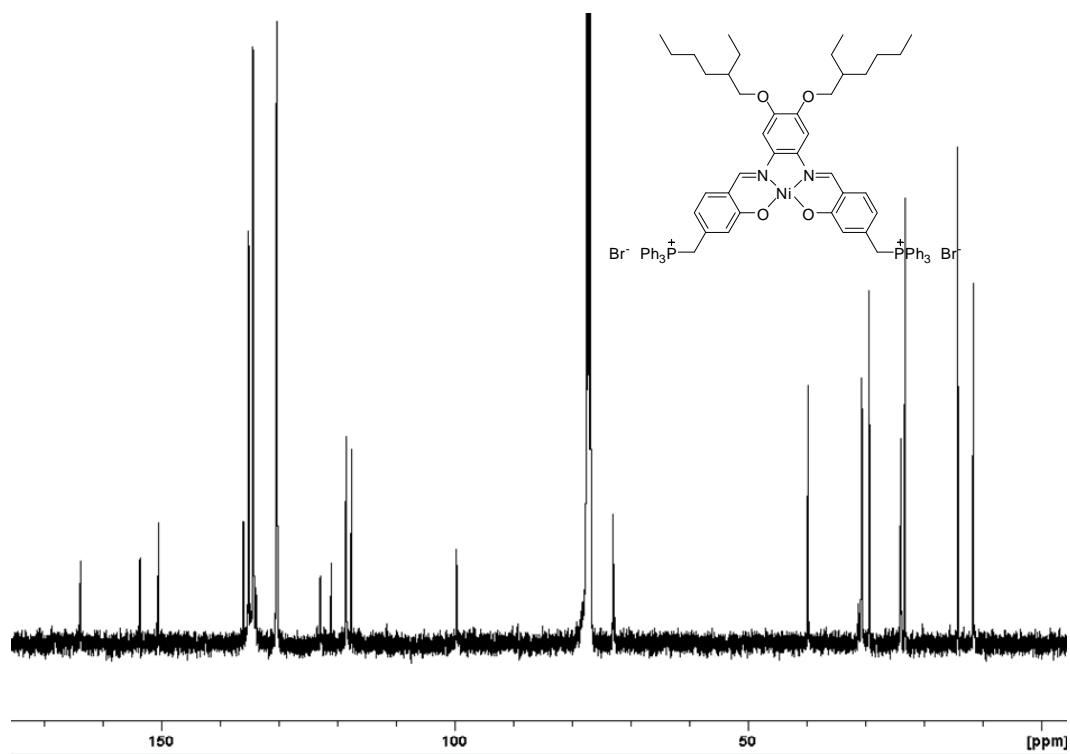


Figure S12. ¹³C NMR spectrum of **9** (100.6 MHz, CDCl₃, 298 K).

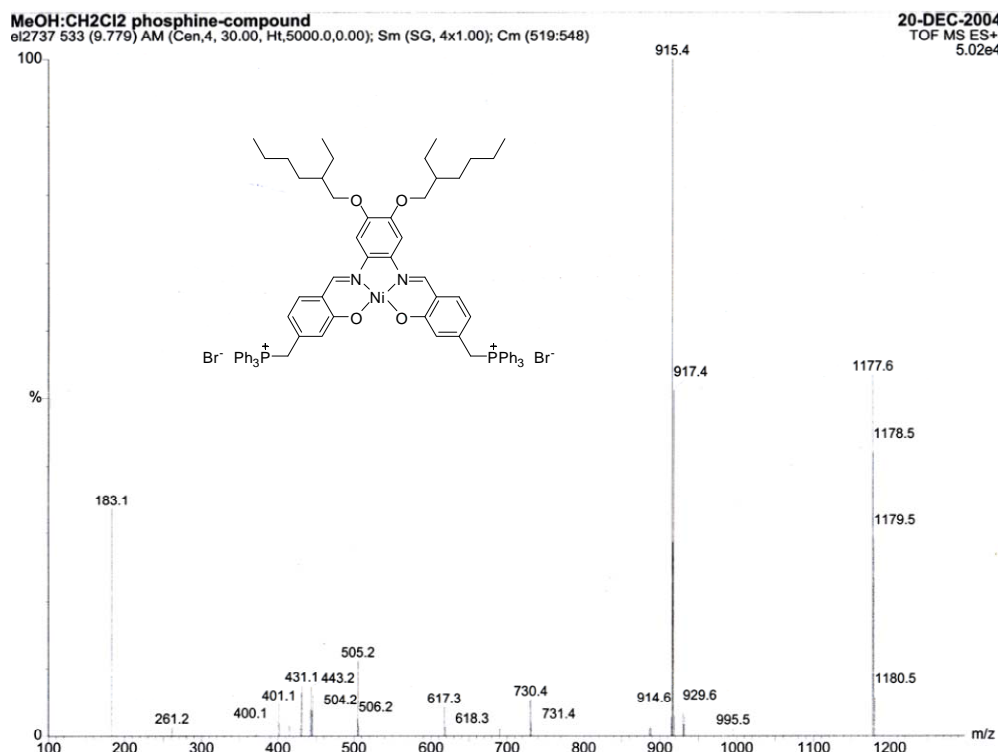


Figure S13. ESI-MS of 9.

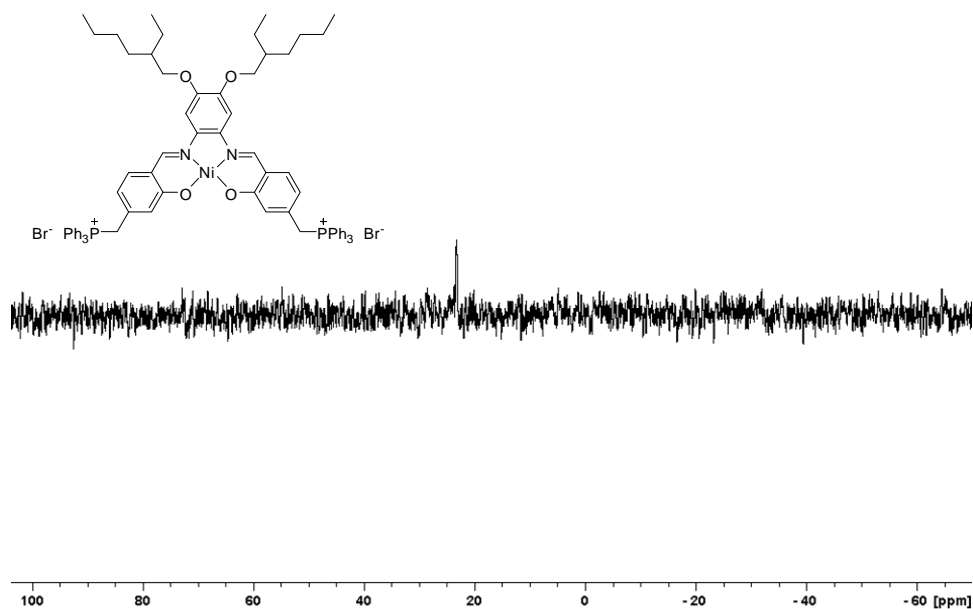


Figure S14. ³¹P NMR spectrum of 9 (121.4 MHz, DMSO-*d*₆, 298 K, 8% H₃PO₄ aqueous solution as external reference).

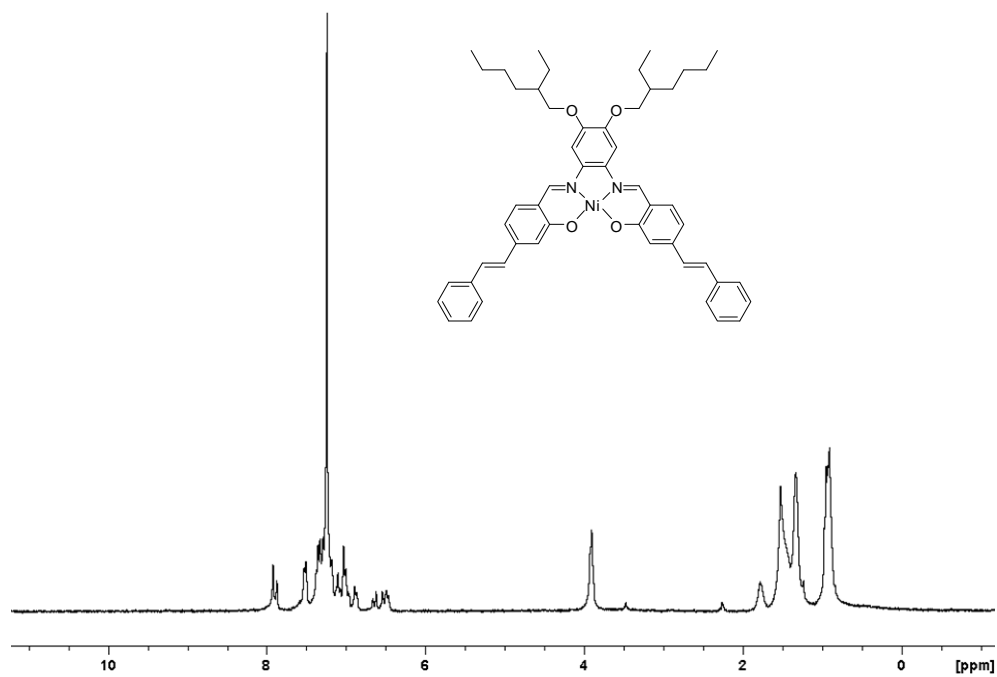


Figure S15. ¹H NMR spectrum of **10** (400 MHz, CDCl₃, 298 K).

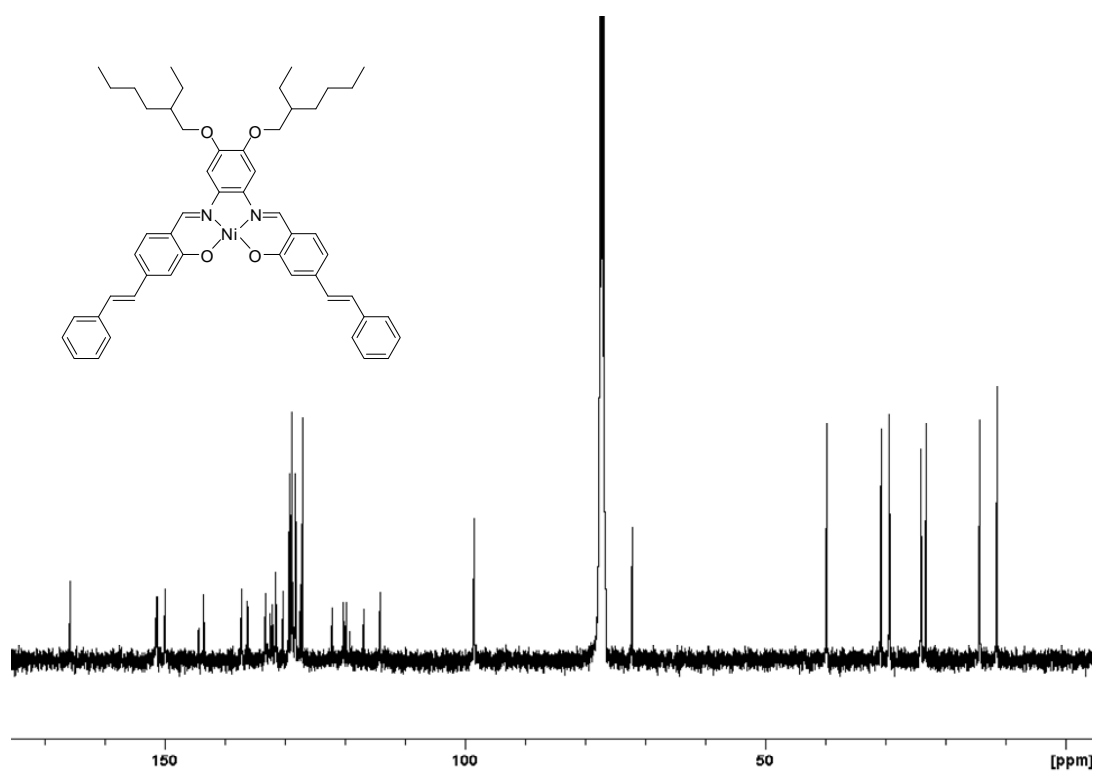


Figure S16. ¹³C NMR spectrum of **10** (100.6 MHz, DMSO-*d*₆, 298 K).

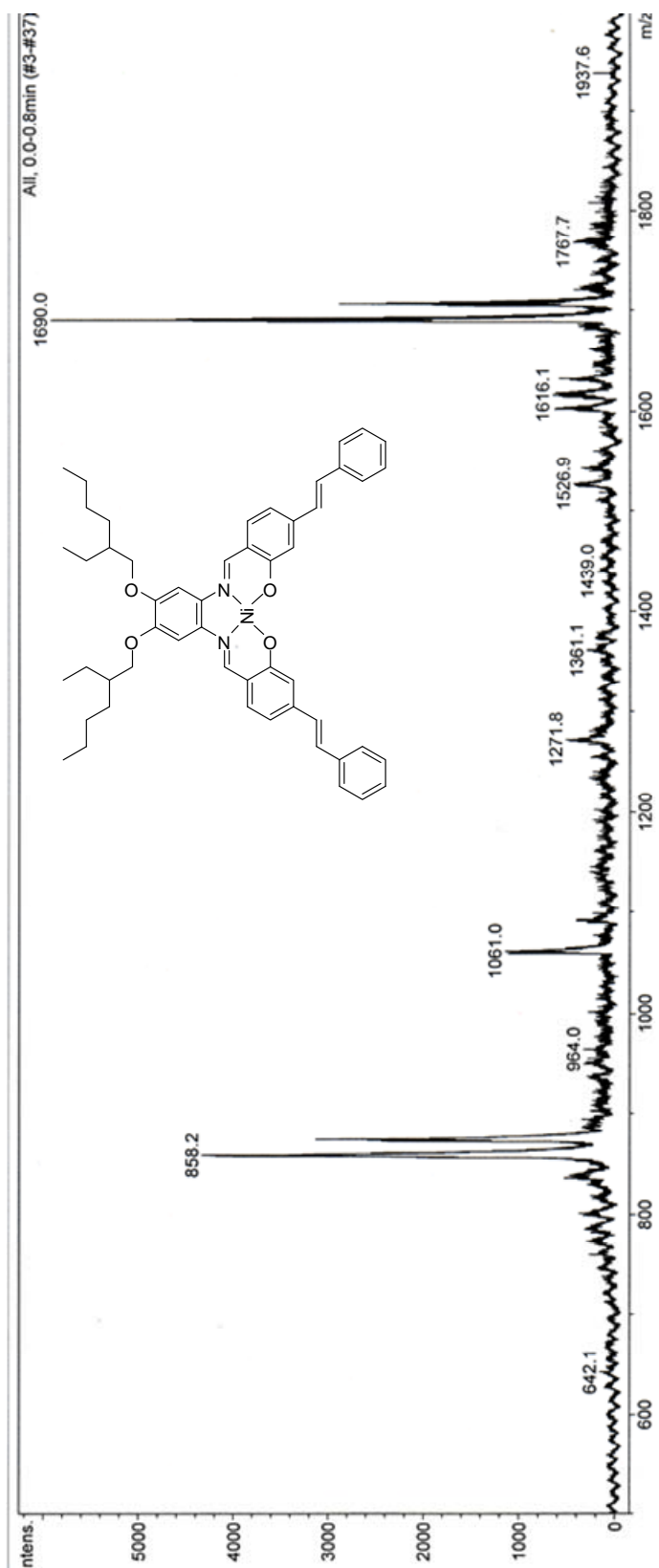


Figure S17. ESI-MS spectrum of **10**.