

Support Information

Article

One-Pot Synthesis of Stable Poly([c2]Daisy-chain Rotaxane) with Pseudo-stopper via Metathesis Reaction and Thiol-ene Reaction

Risako Kamoto, Kenjiro Onimura and Kazuhiro Yamabuki *

Graduate School of Sciences and Technology for Innovation, Yamaguchi University, 2-16-1 Tokiwadai, Ube, Yamaguchi 755-8611, Japan; b014vfv@yamaguchi-u.ac.jp (R.K.); onimura@yamaguchi-u.ac.jp (K.O.)

* Correspondence: yamabuki@yamaguchi-u.ac.jp

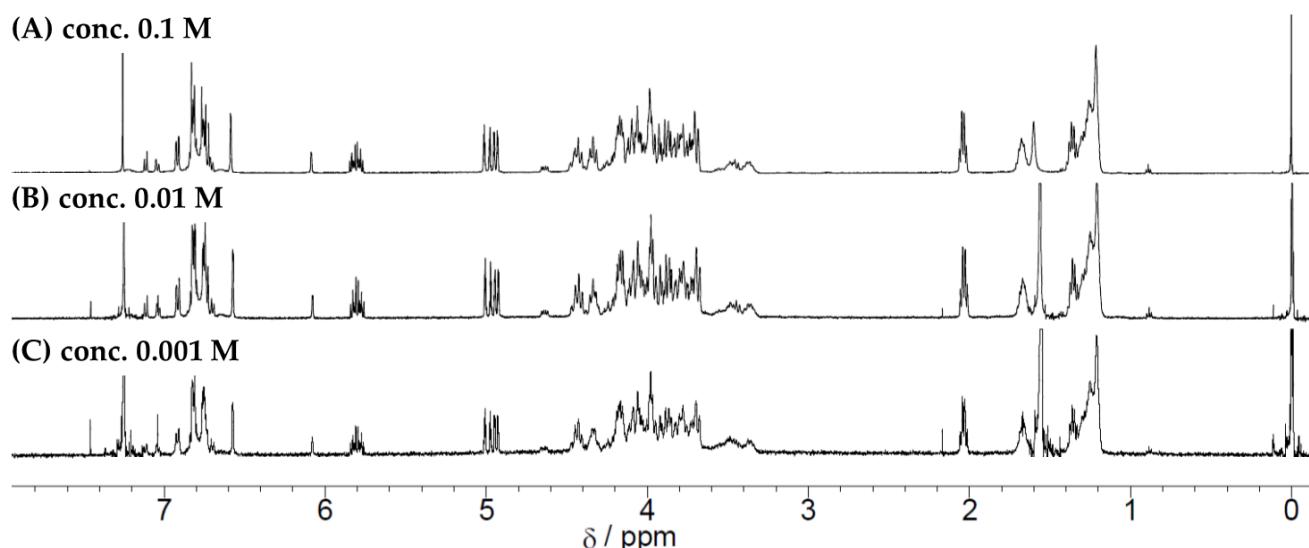


Figure S1. ^1H NMR spectra of H-G monomer at (A) conc. 0.1 M, (B) 0.01, and (C) 0.001 M in CDCl_3 .

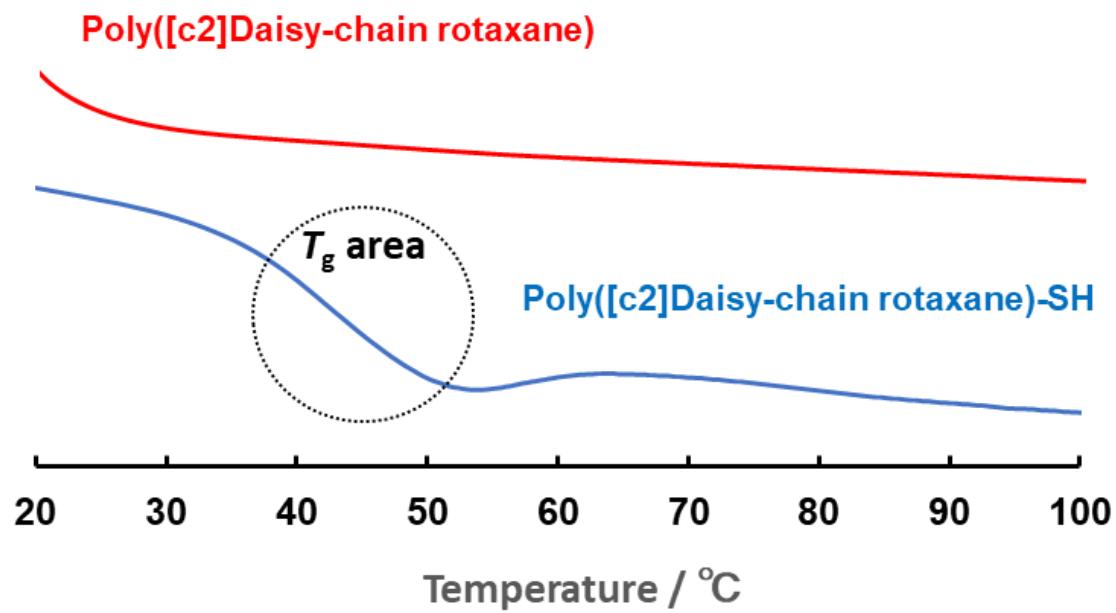


Figure S2. DSC charts of poly([c2]daisy-chain rotaxane) (red line) and poly([c2]daisy-chain rotaxane)-SH (blue line).

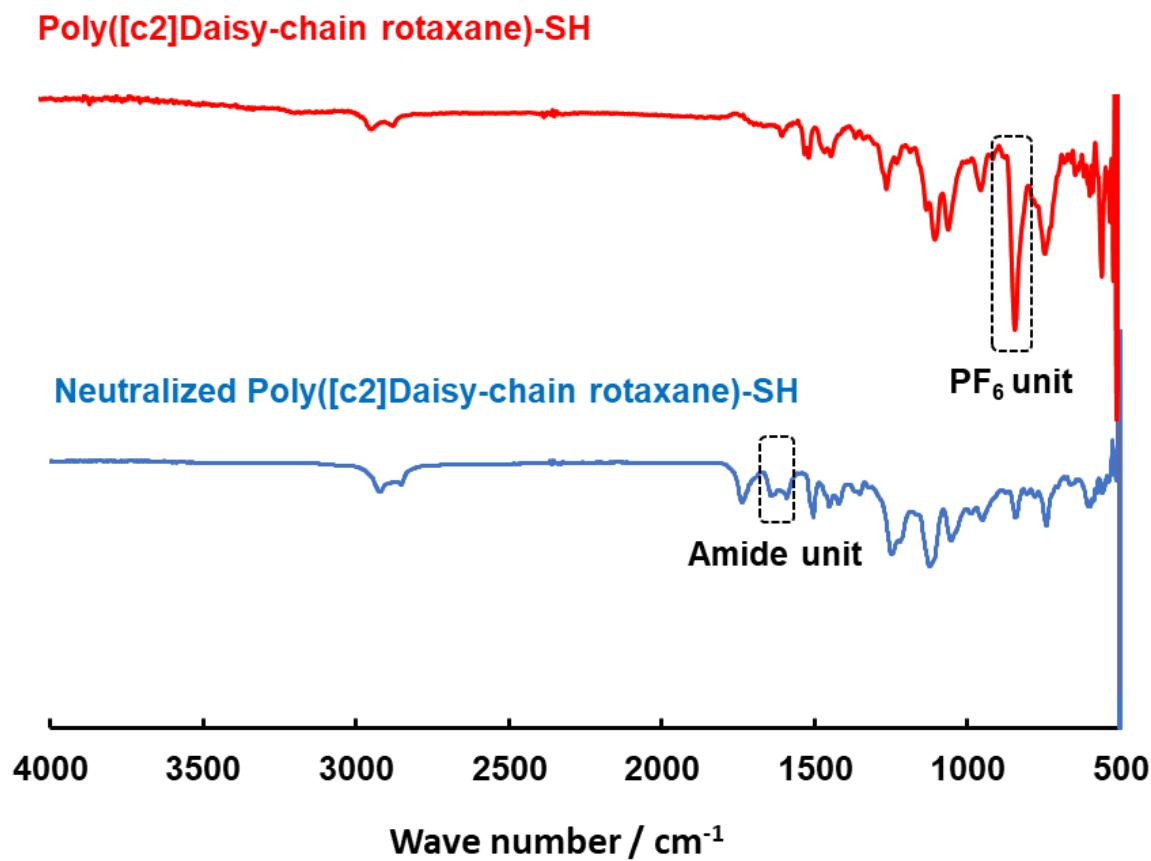


Figure S3. IR spectra of poly([c2]daisy-chain rotaxane)-SH (red line) and neutralized poly([c2]daisy-chain rotaxane)-SH (blue line).

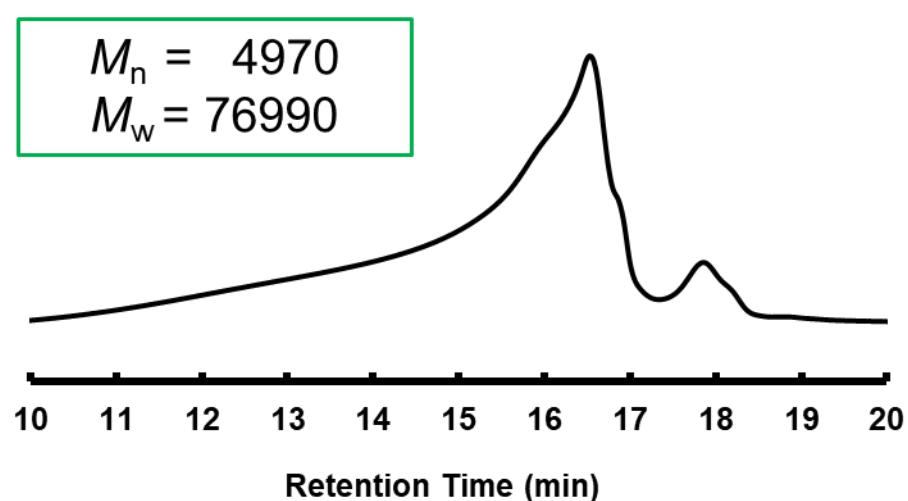


Figure S4. GPC chart of neutralized poly([c2]Daisy-chain rotaxane-SH (eluent: CHCl₃).

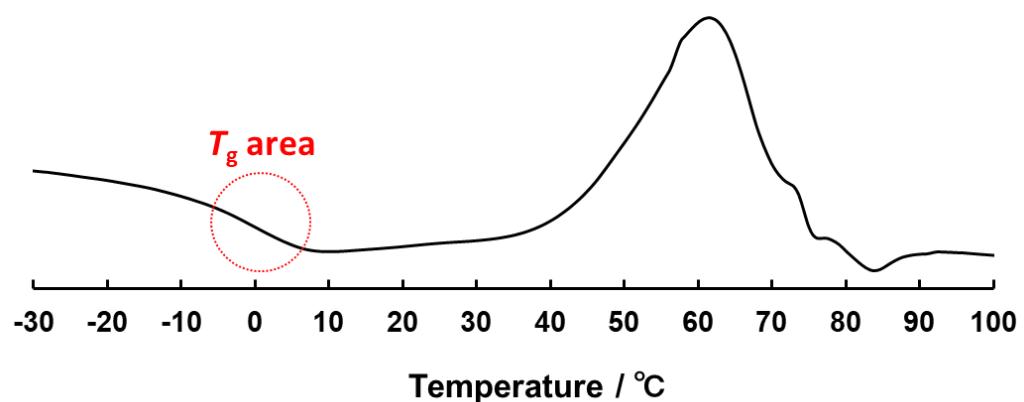


Figure S5. DSC charts of neutralized poly([c2]Daisy-chain rotaxane).

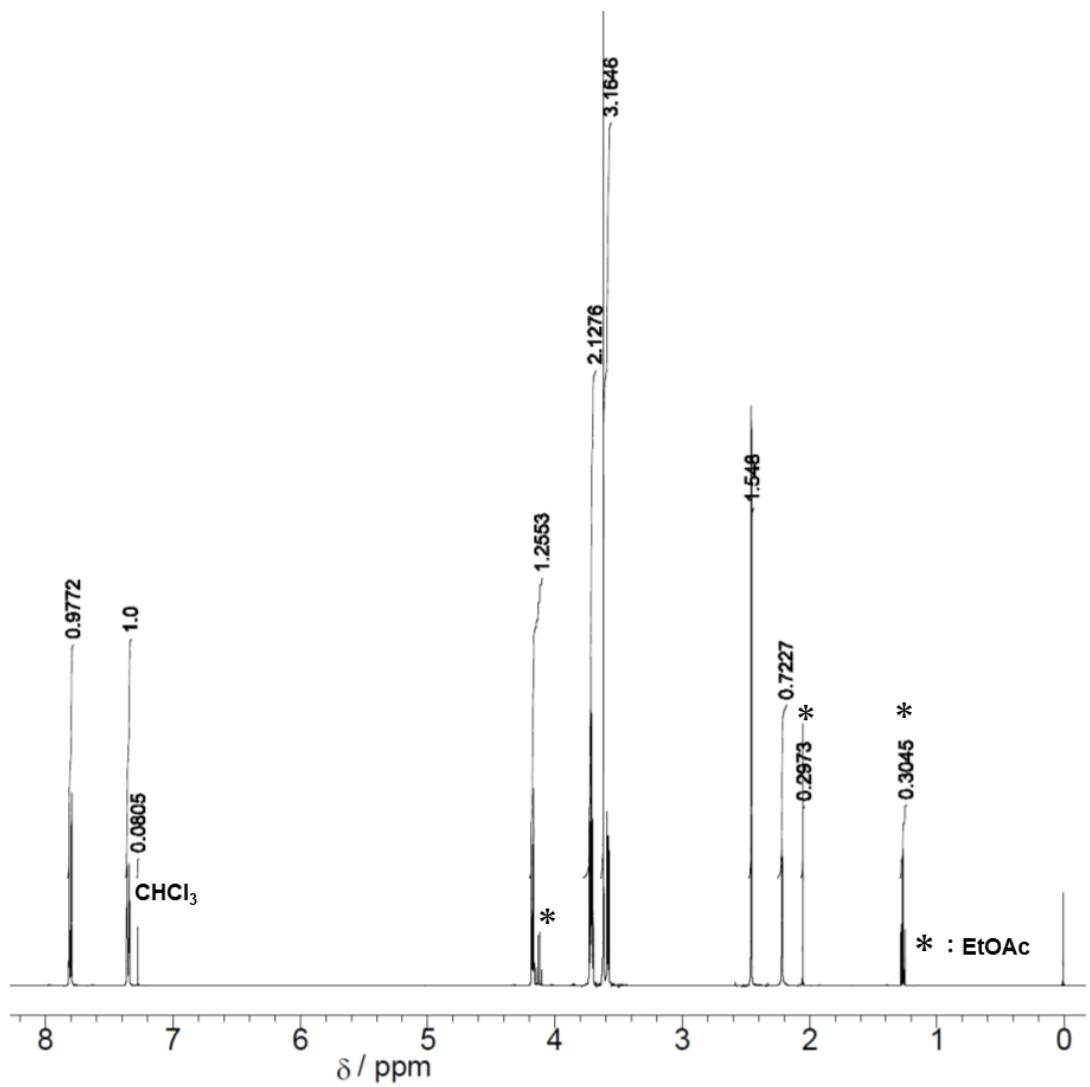


Figure S6. ^1H NMR spectra of compound **1** in CDCl_3 .

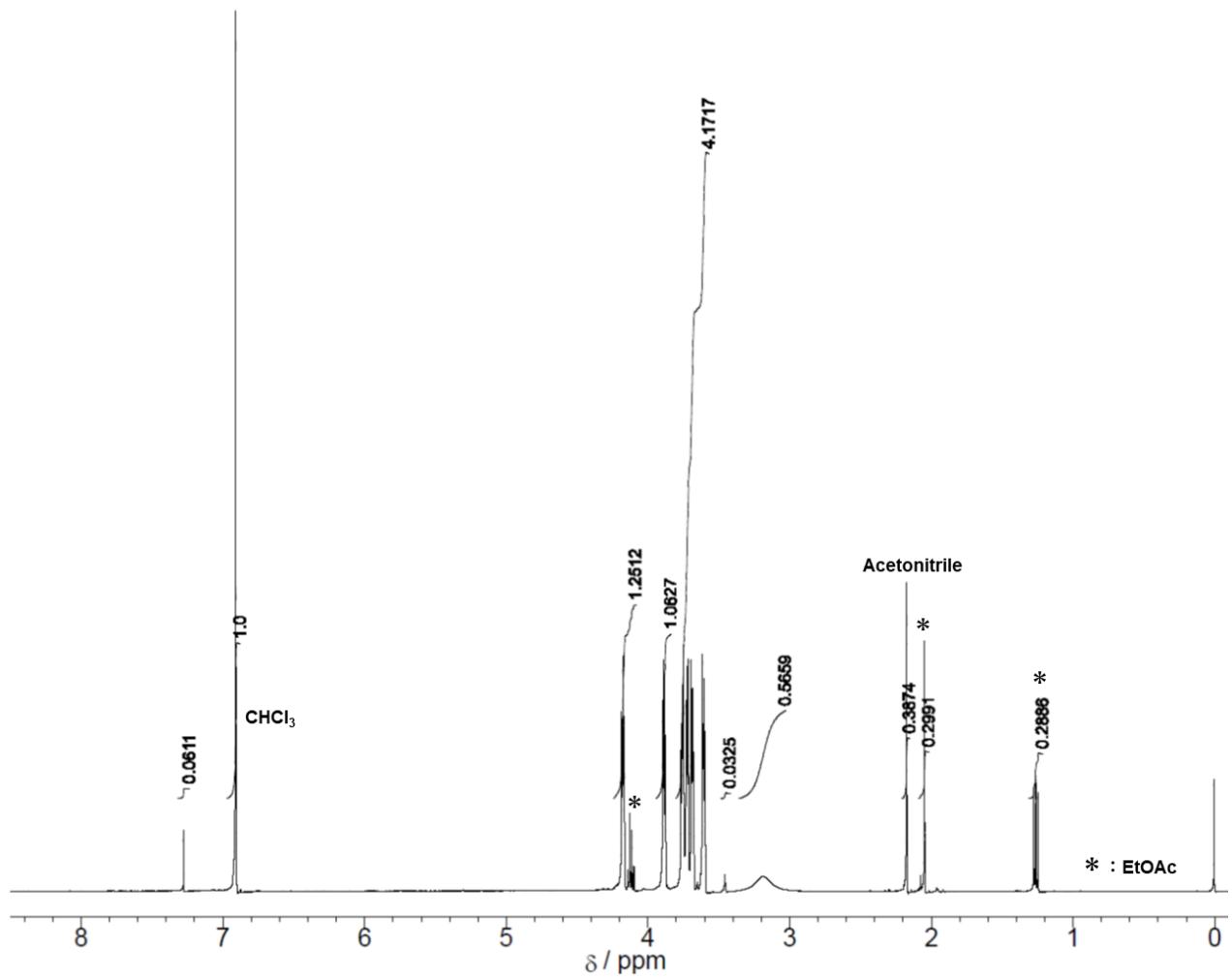


Figure S7. ^1H NMR spectra of compound 2 in CDCl_3 .

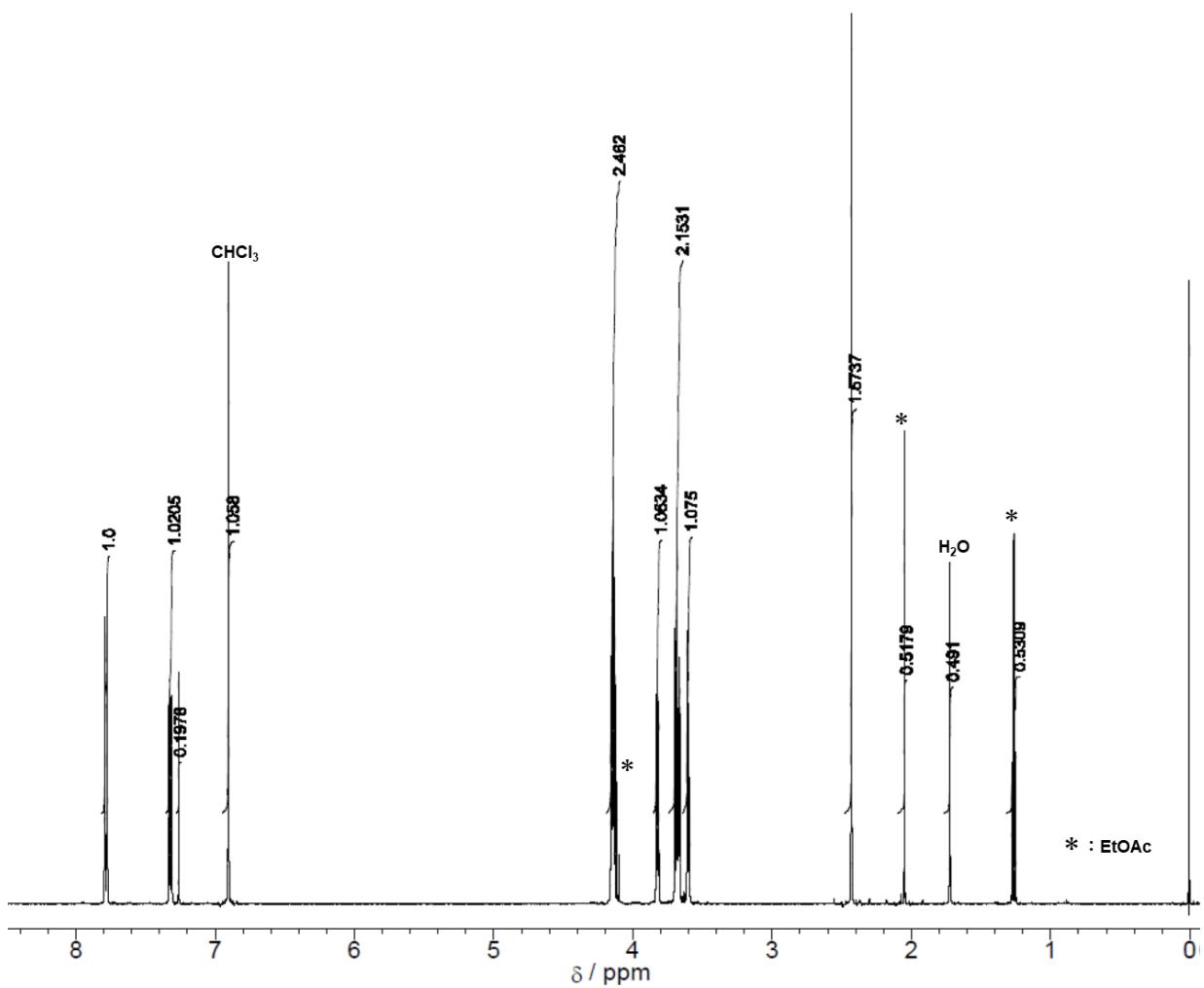


Figure S8. ^1H NMR spectra of compound 3 in CDCl_3 .

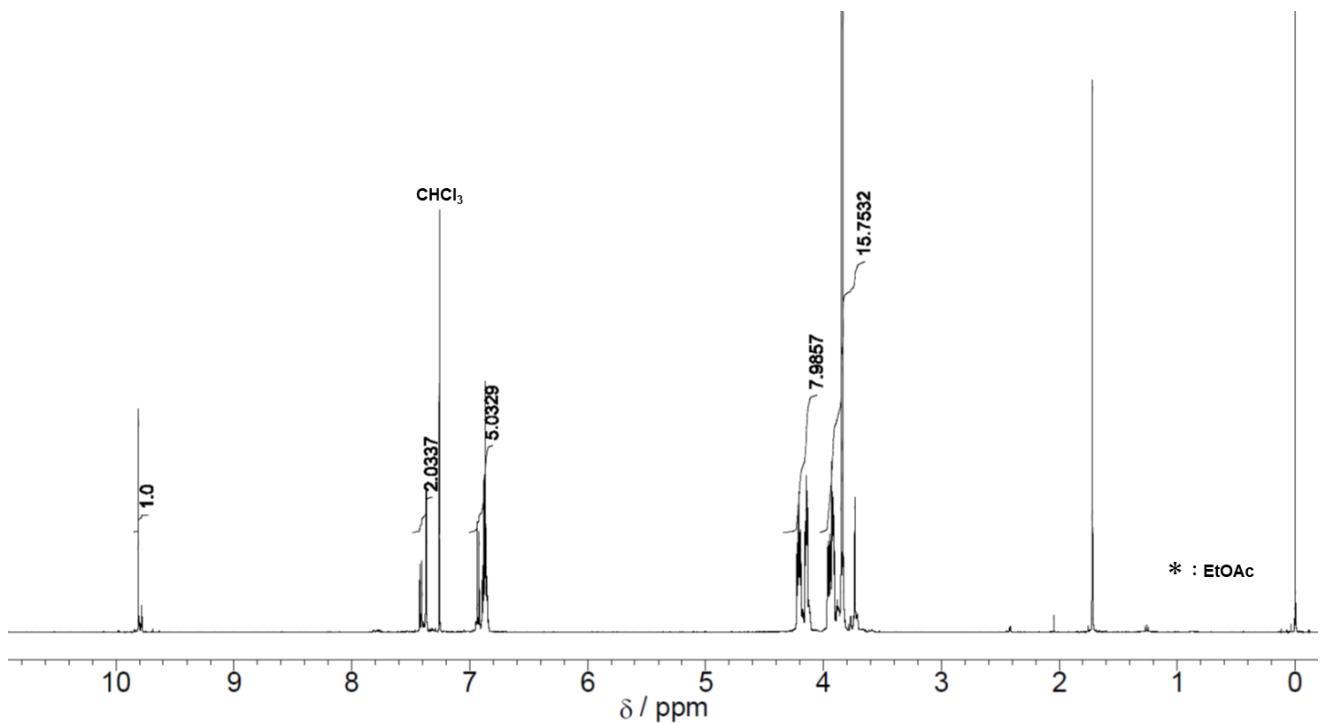


Figure S9. ^1H NMR spectra of compound 4 in CDCl_3 .

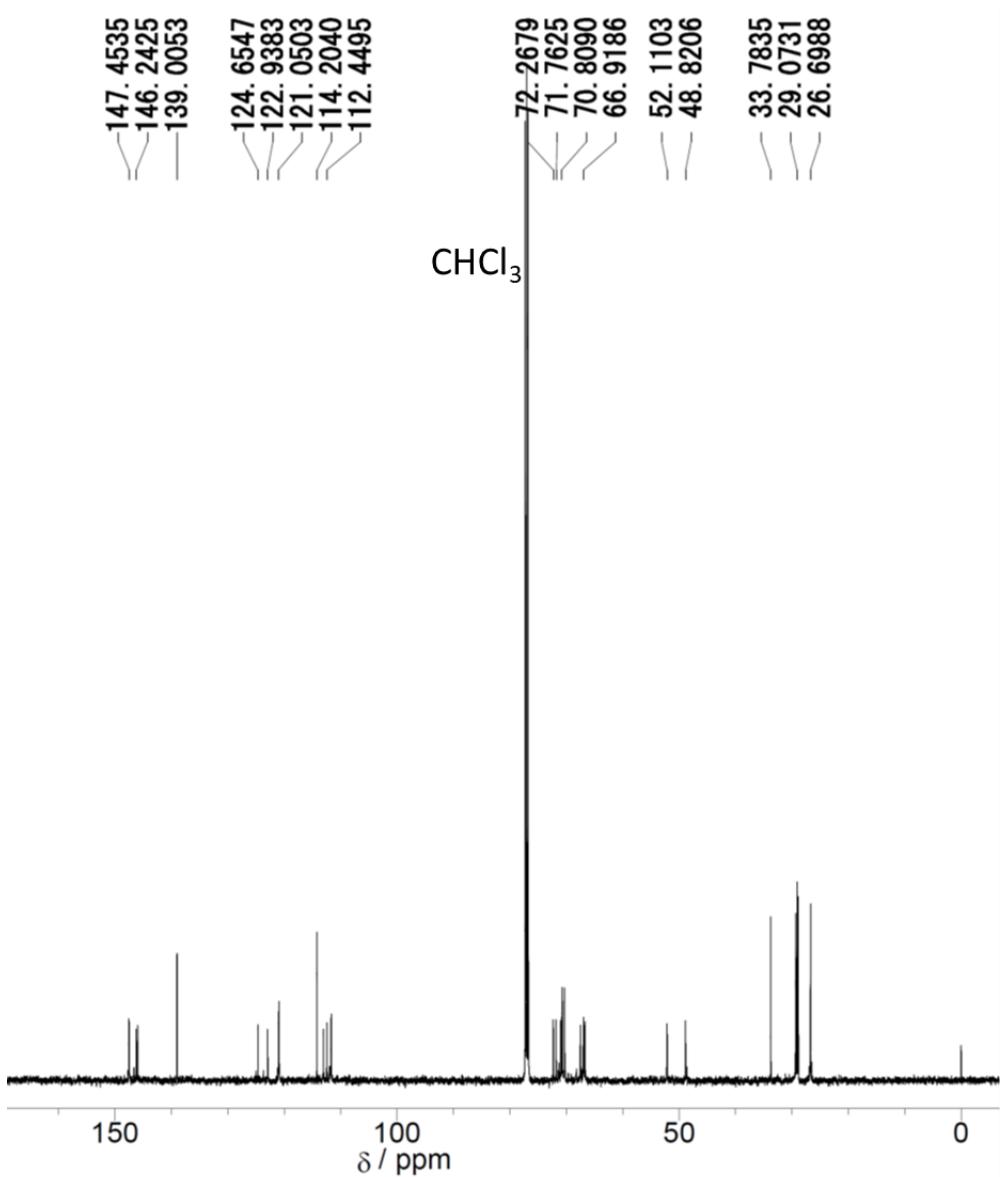


Figure S10. ^{13}C NMR spectra of H-G monomer in CDCl_3 .