

New Journal of Chemistry

Supplementary Information

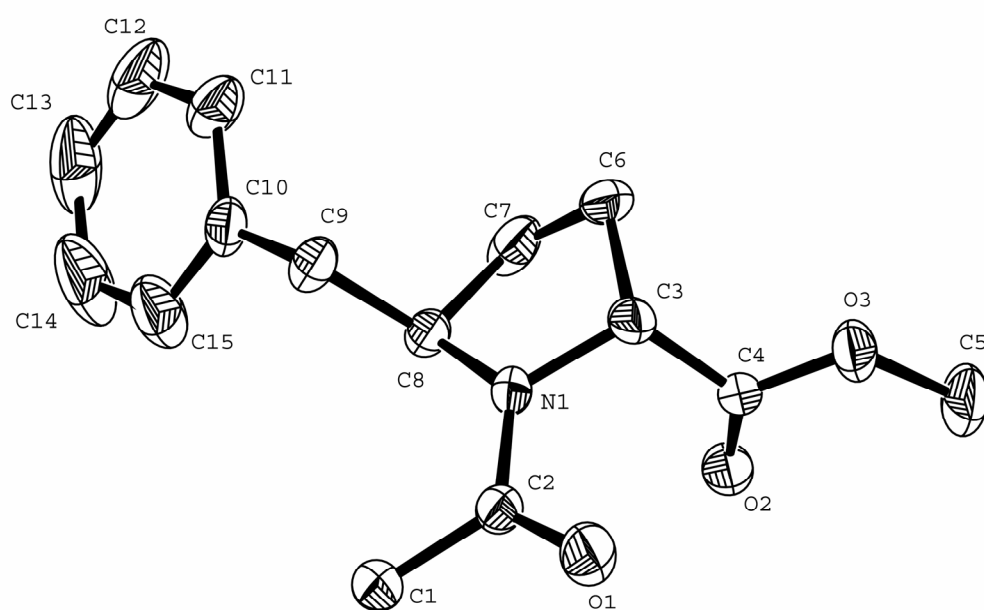
for

Synthesis of enantiomerically pure δ -benzylproline derivatives

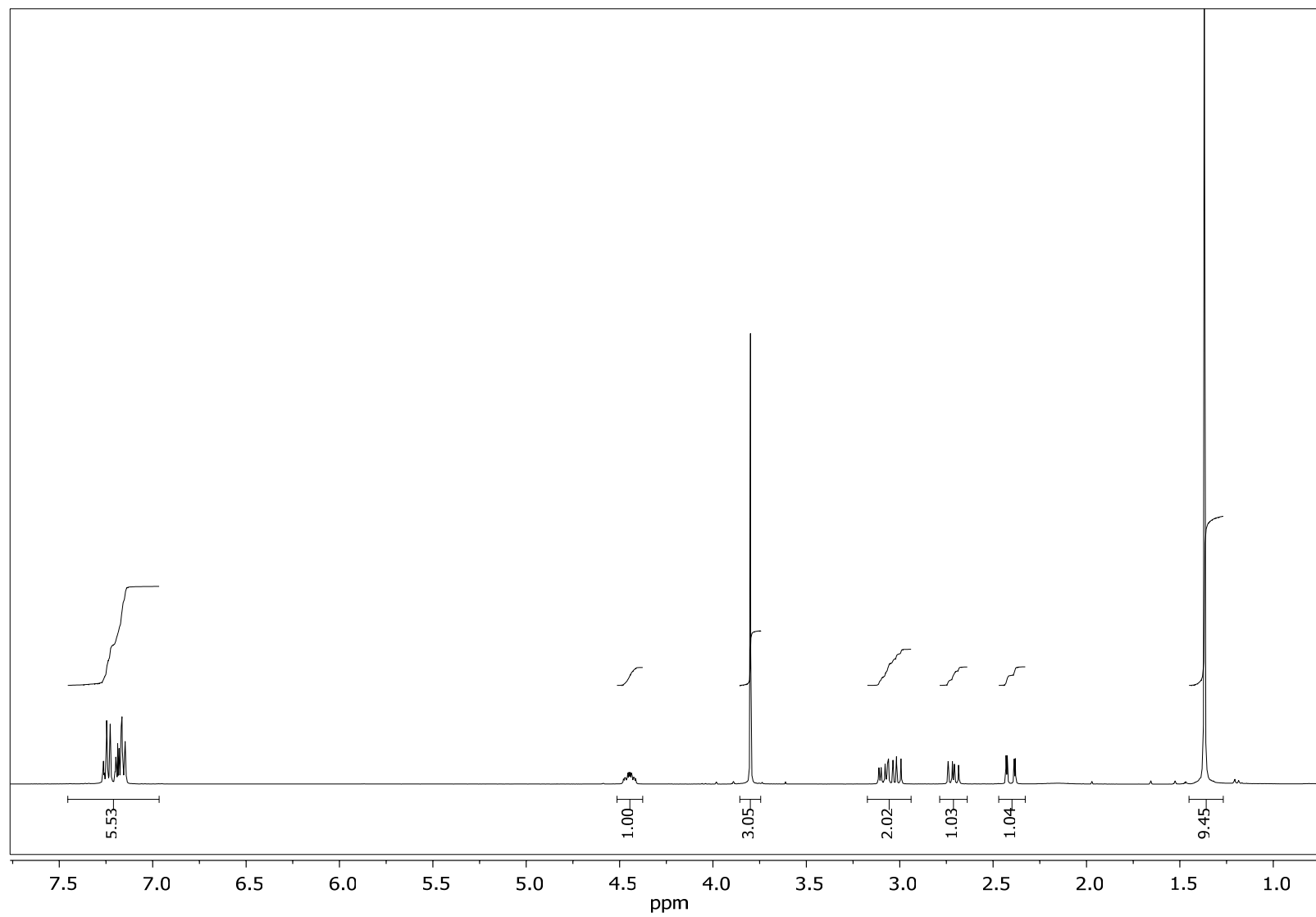
M. Isabel Rodríguez, M. Isabel Calaza, Ana I. Jiménez, and Carlos Cativiela

Contents

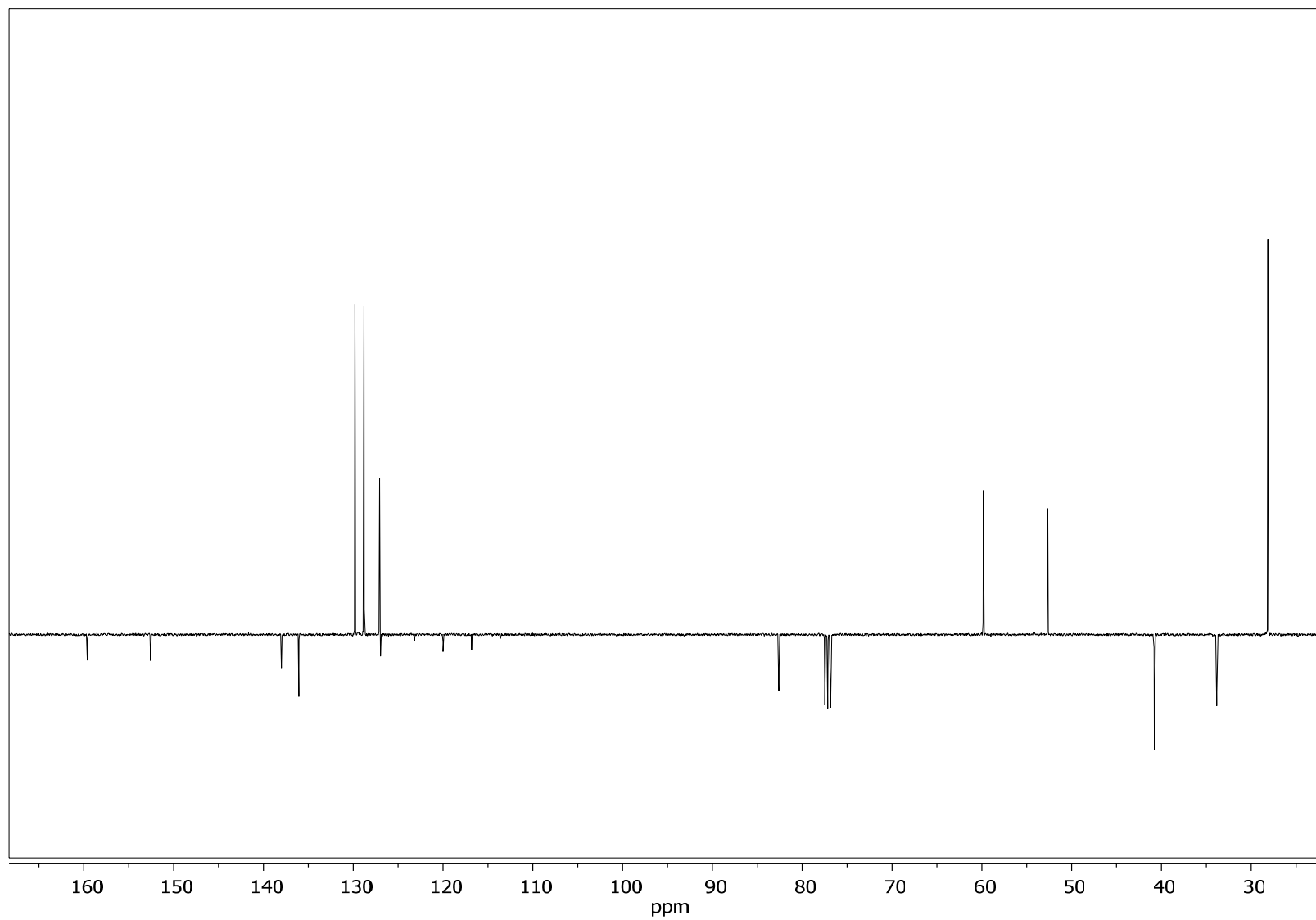
X-ray crystal structure of (2 <i>R</i> ,5 <i>R</i>)- 10 with atom numbering	Page S2
¹ H-NMR and ¹³ C-NMR (APT) spectra of compounds 6–10	Pages S3–S16



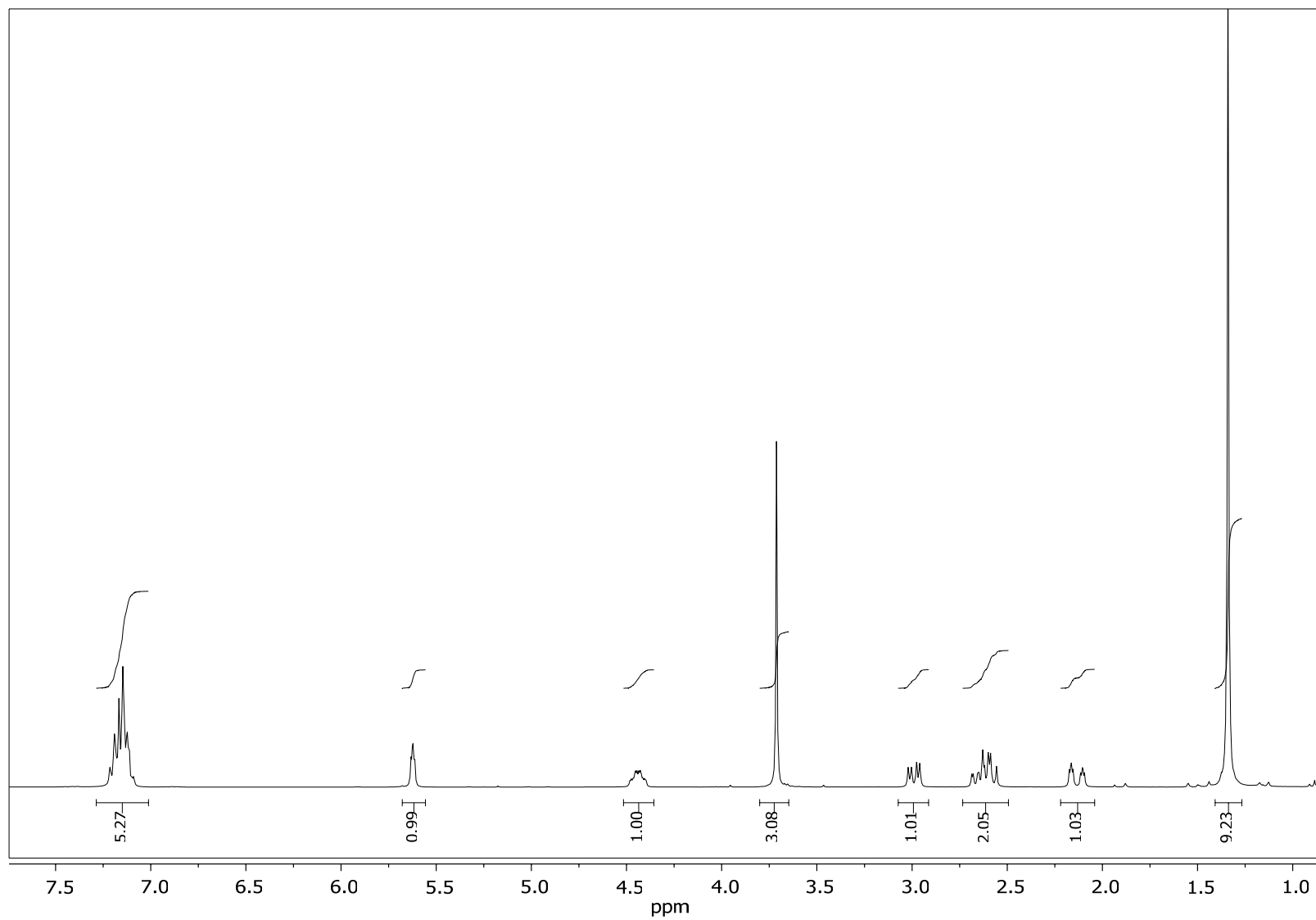
ORTEP diagram for the X-ray crystal structure of (2*R*,5*R*)-**10** with atom numbering. Thermal ellipsoids are shown at 30% probability. Hydrogen atoms have been omitted for clarity.



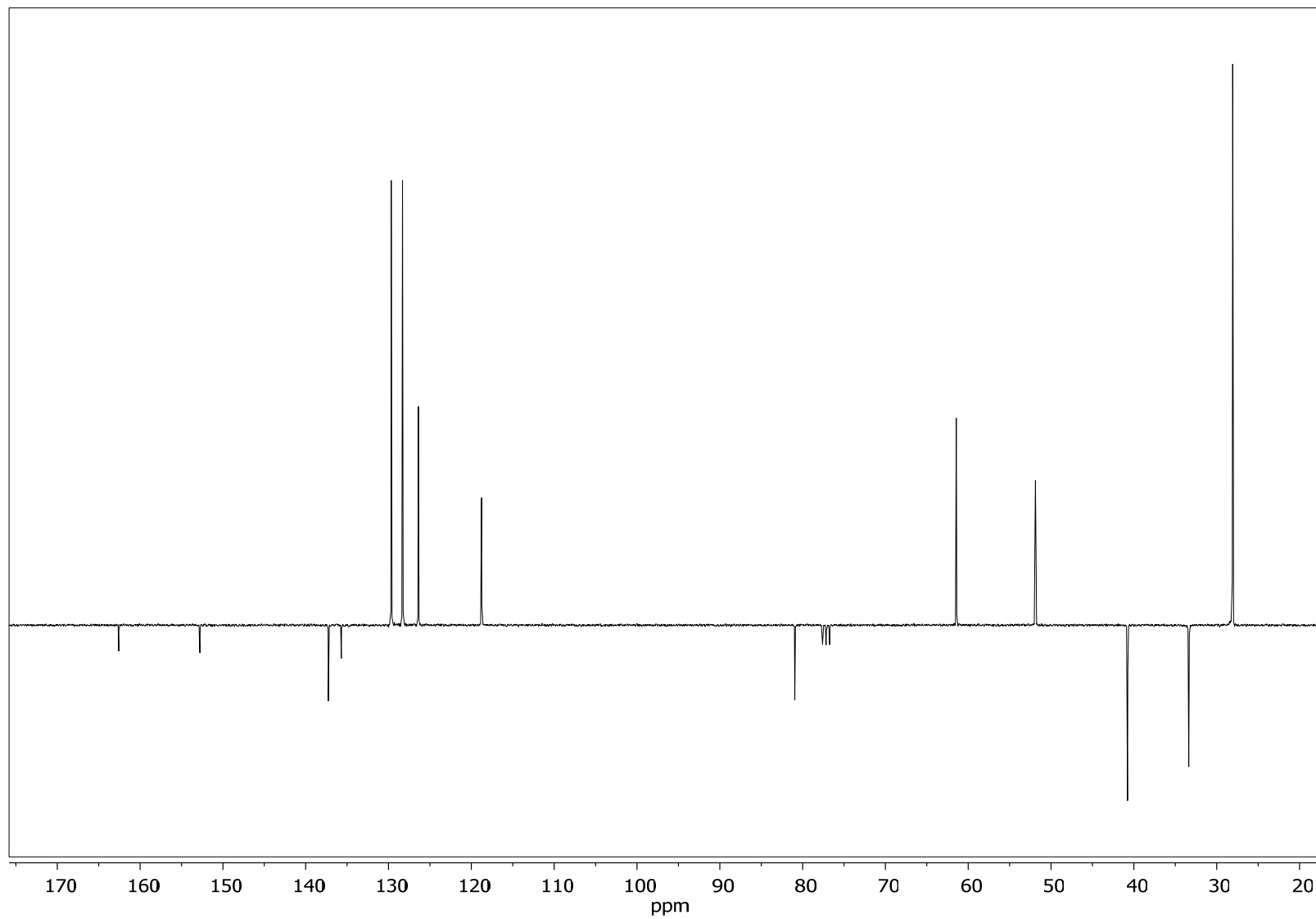
^1H -NMR spectrum of **6** in CDCl_3 (400 MHz, 298K)



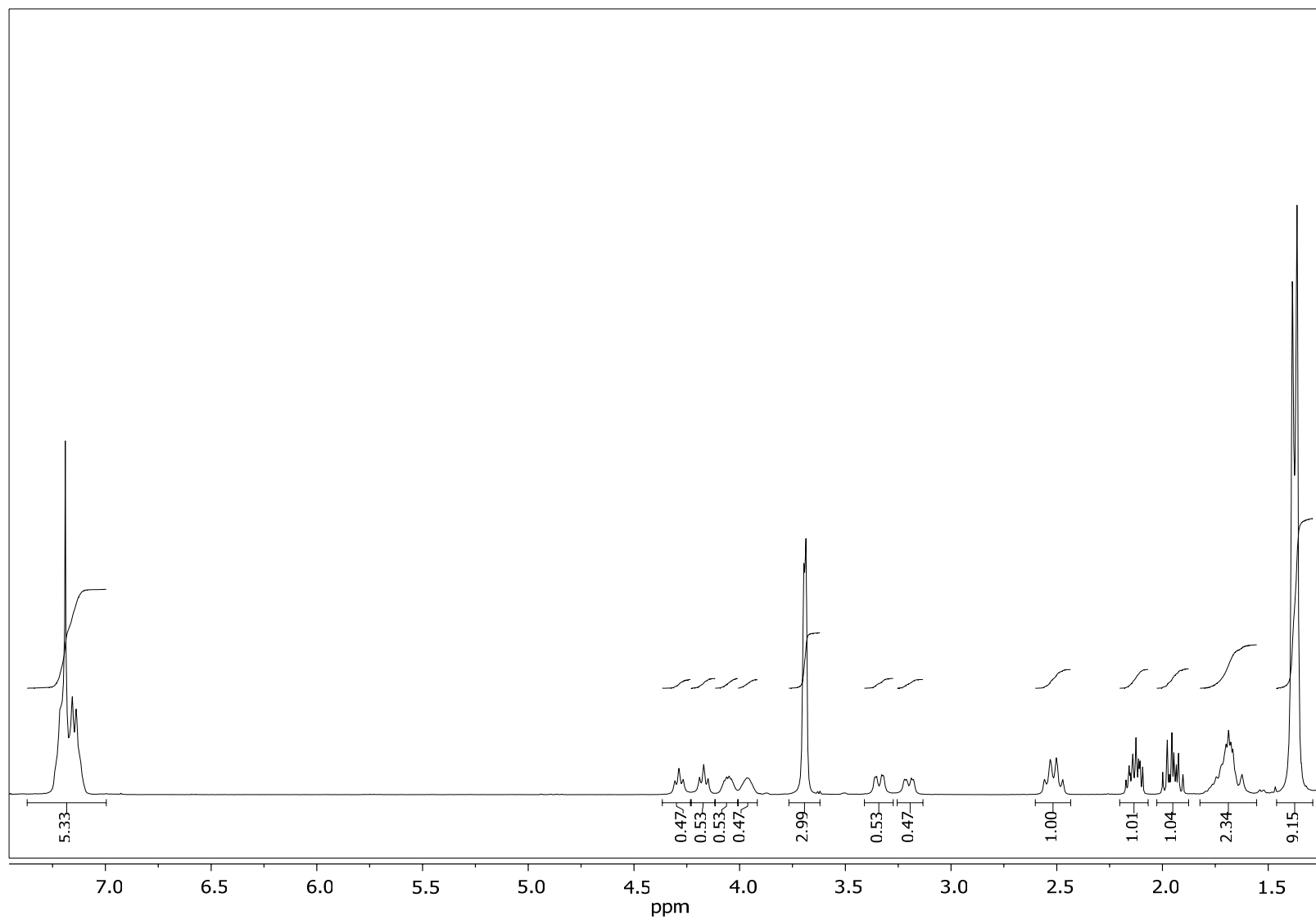
^{13}C -NMR spectrum of **6** in CDCl_3 (100 MHz, 298K)



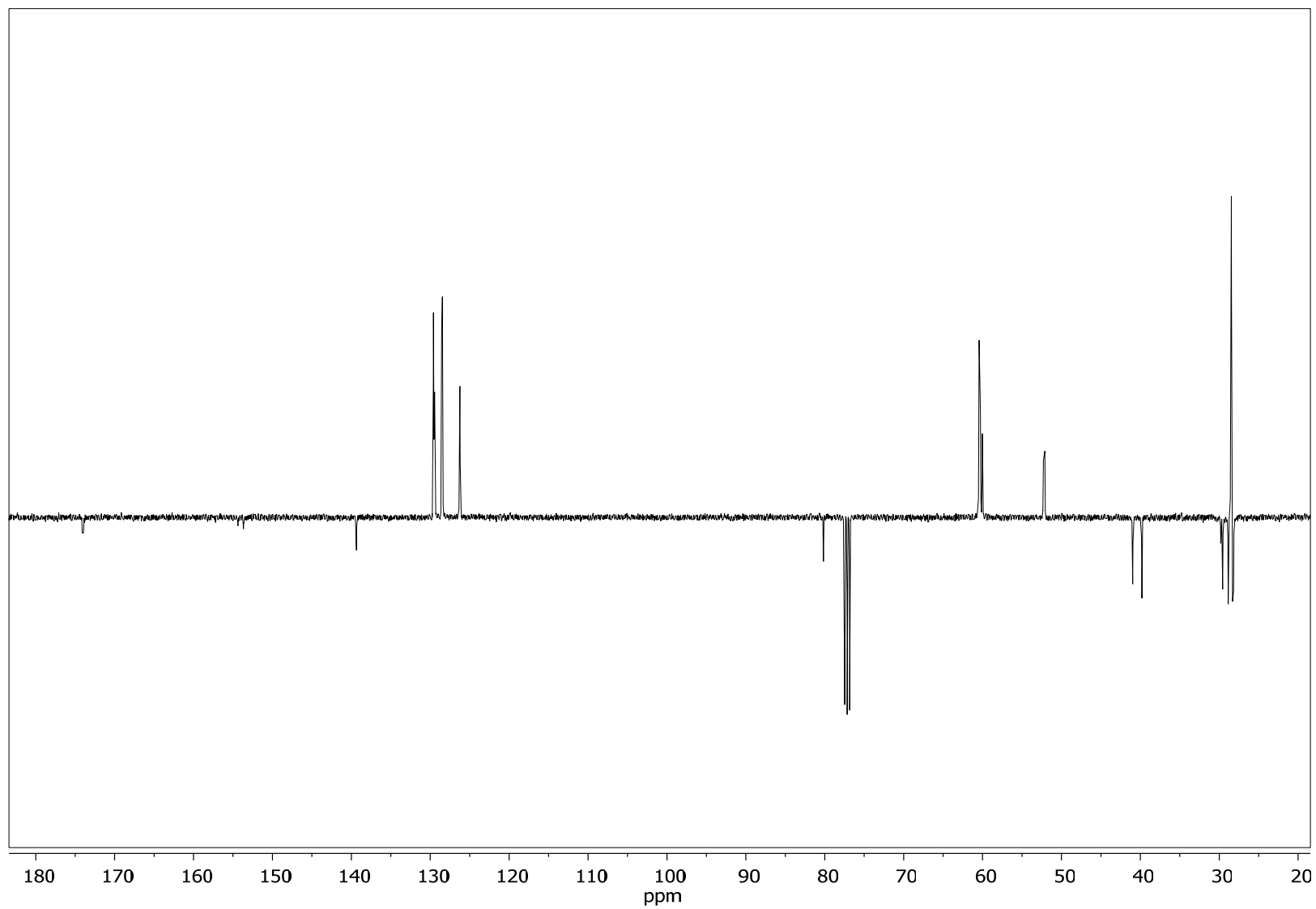
^1H -NMR spectrum of **7** in CDCl_3 (300 MHz, 298K)



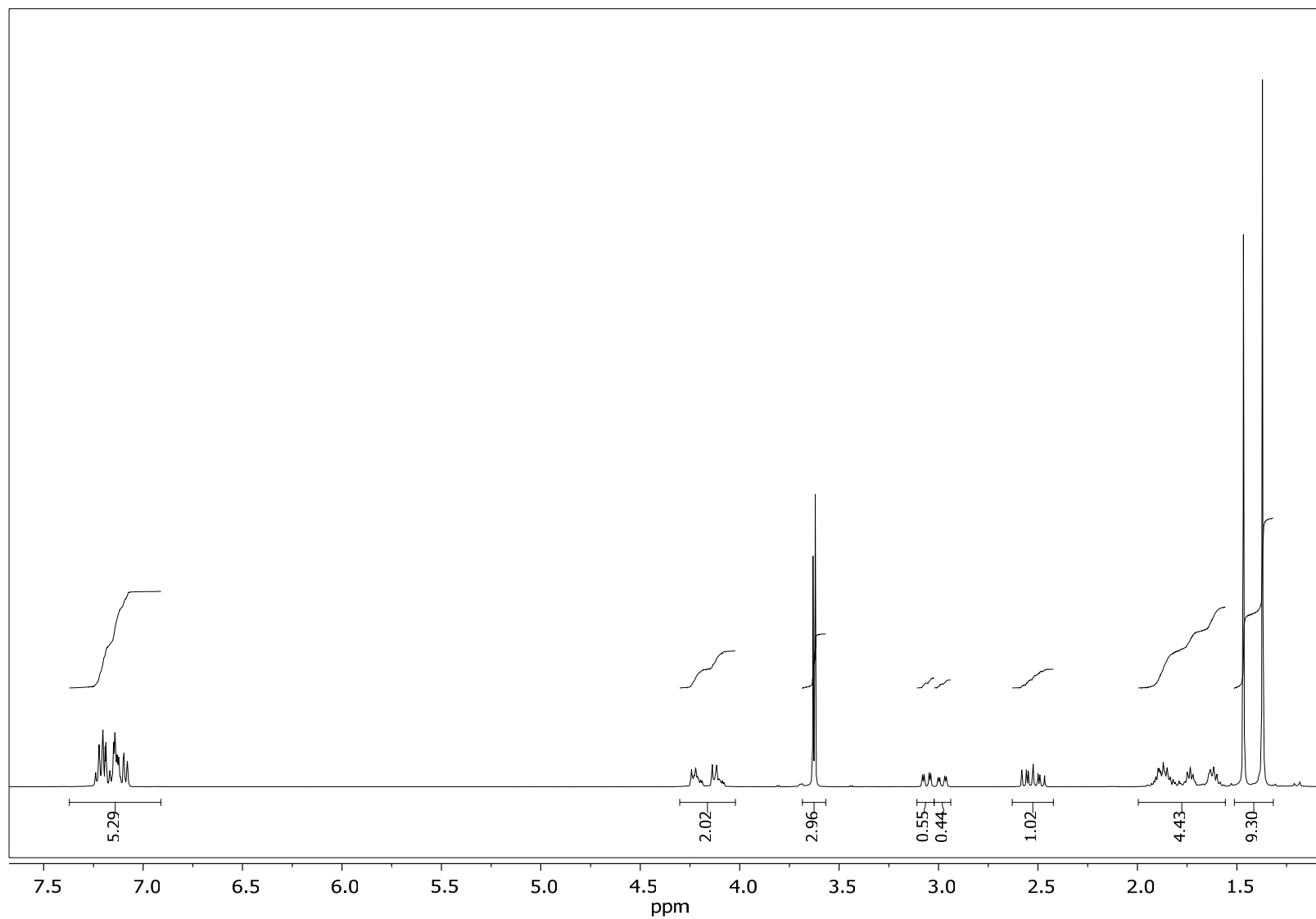
^{13}C -NMR spectrum of **7** in CDCl_3 (75 MHz, 298K)



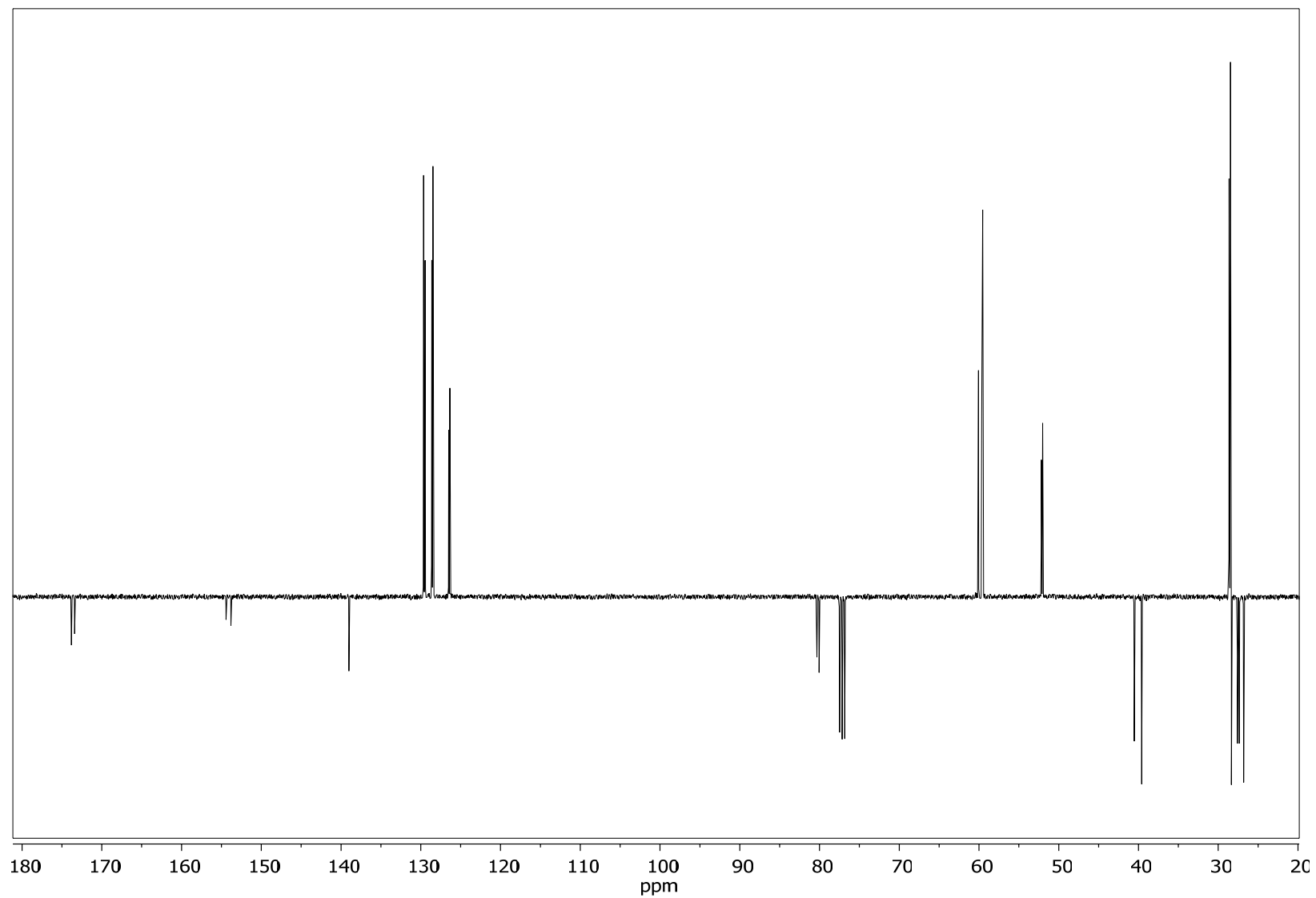
^1H -NMR spectrum of (2S,5R)-8 in CDCl_3 (400 MHz, 298K)



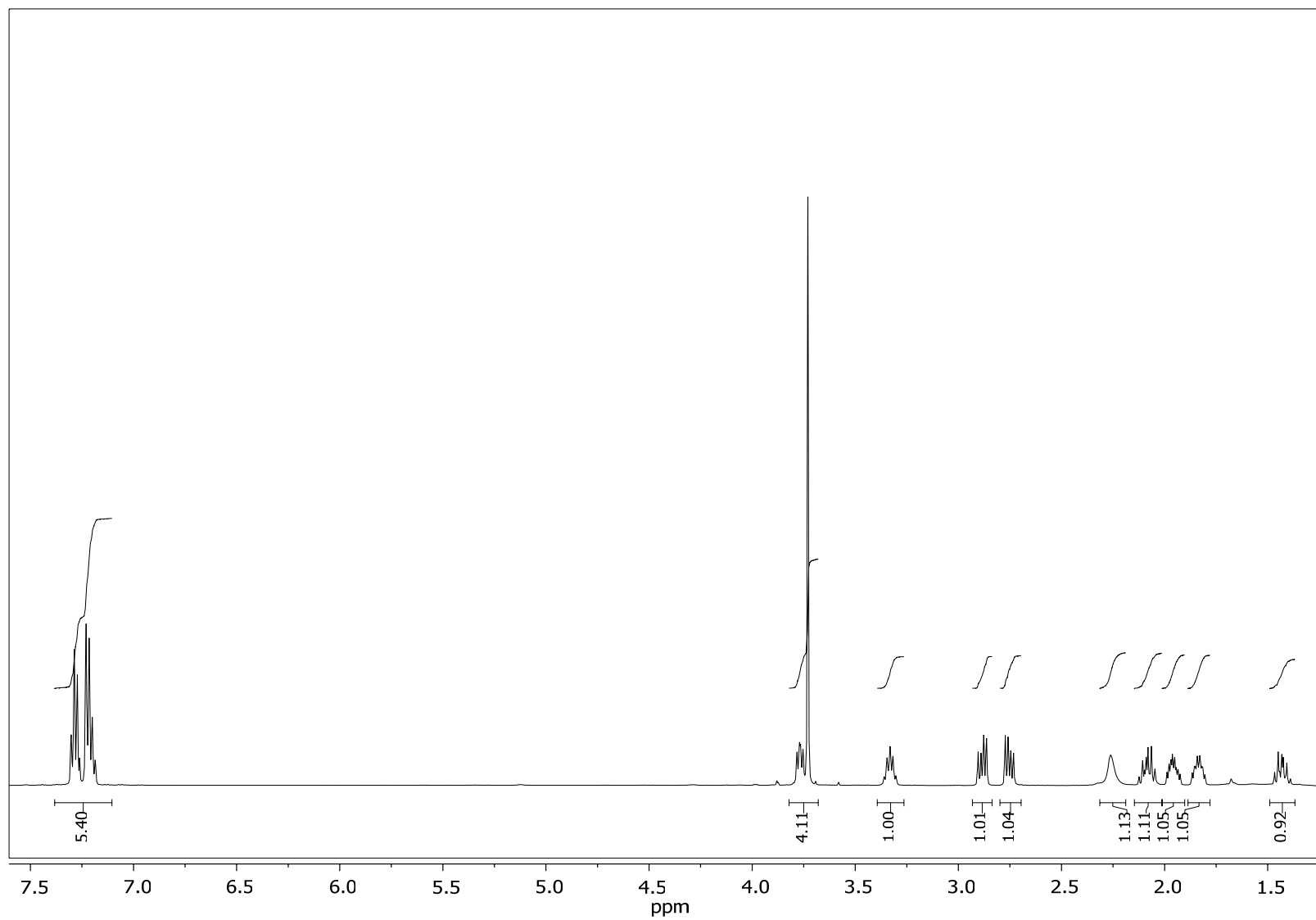
^{13}C -NMR spectrum of (2*S*,5*R*)-**8** in CDCl_3 (100 MHz, 298K)



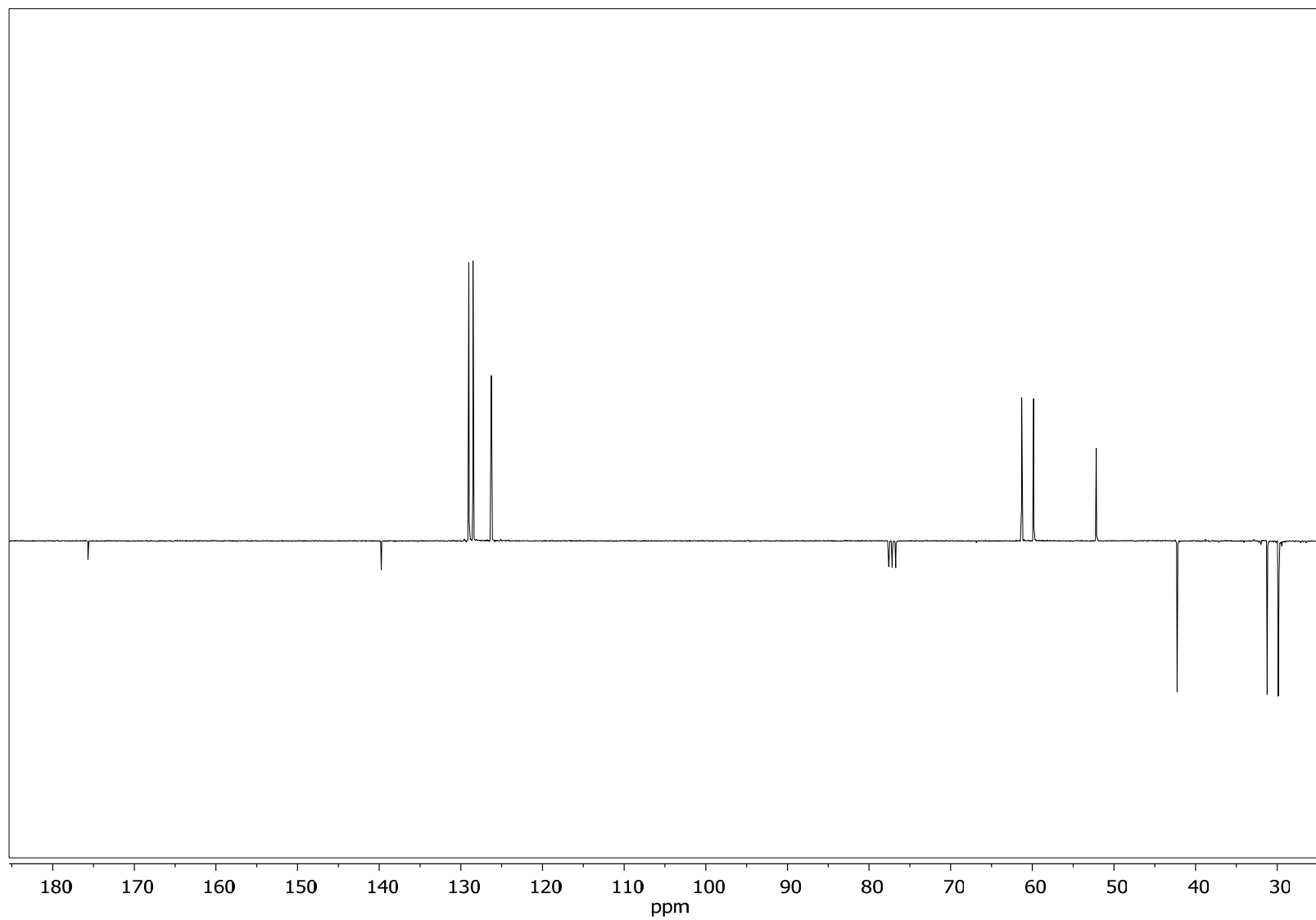
^1H -NMR spectrum of (2*R*,5*R*)-**8** in CDCl_3 (400 MHz, 298K)



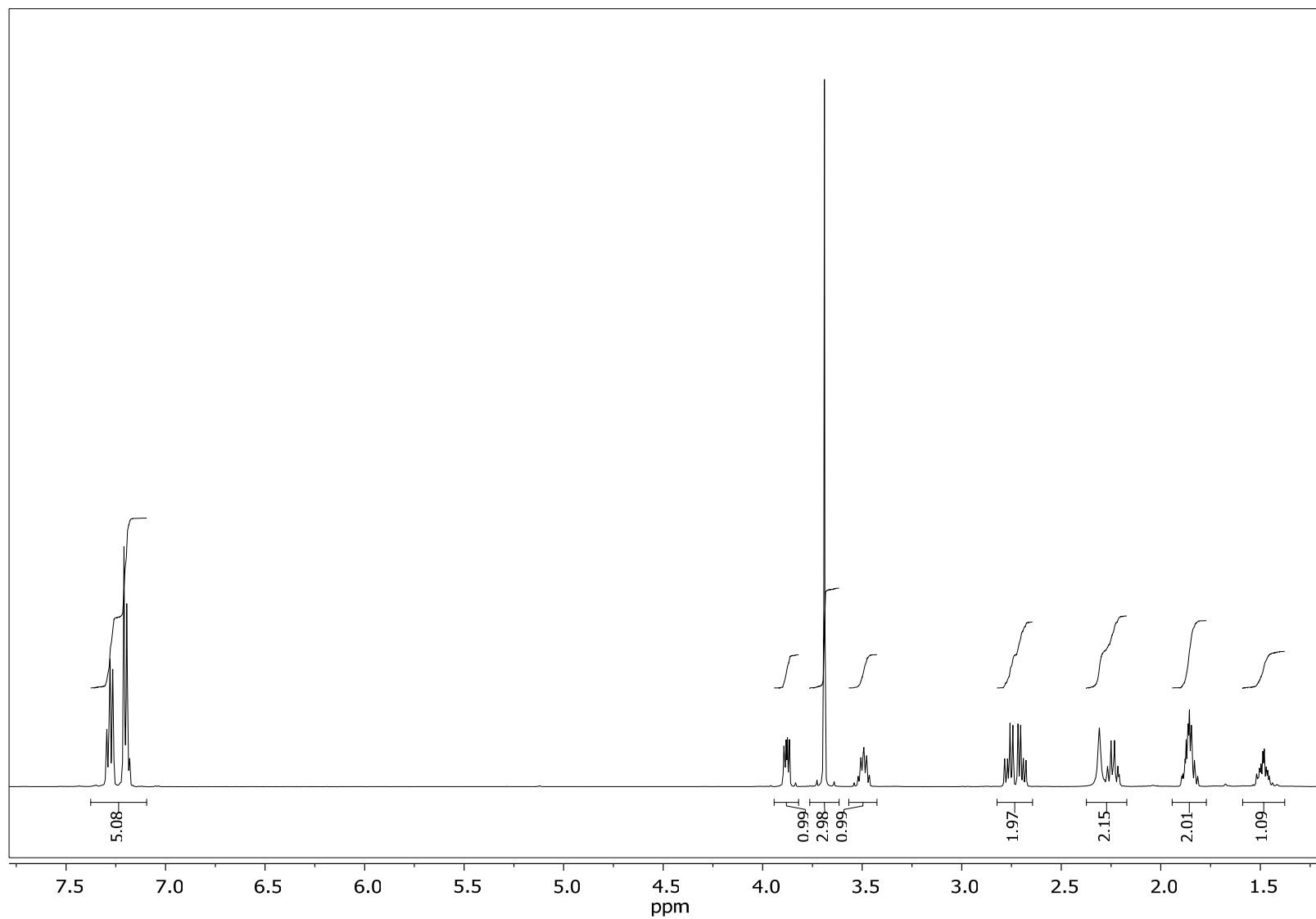
^{13}C -NMR spectrum of (2*R*,5*R*)-**8** in CDCl_3 (100 MHz, 298K)



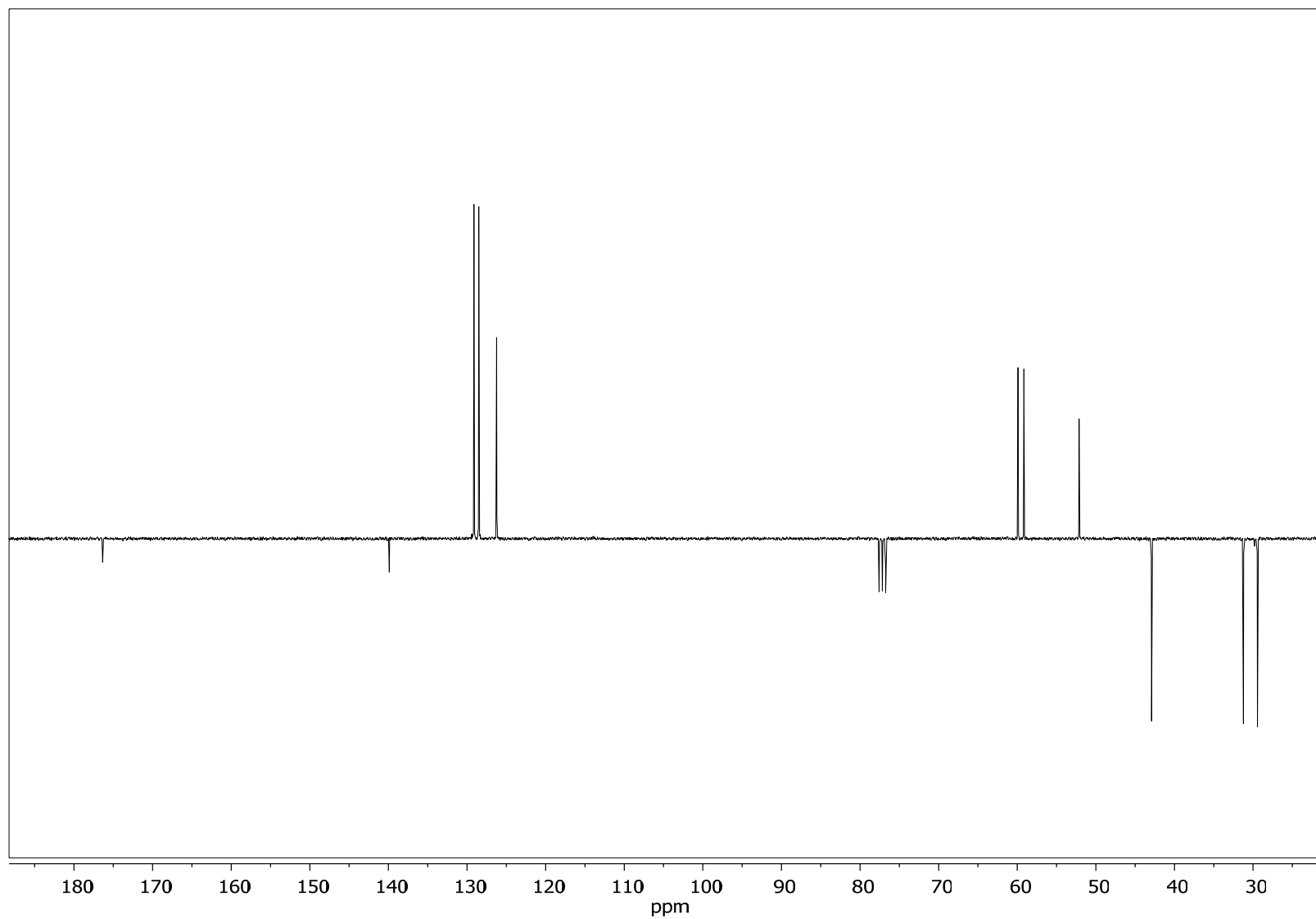
^1H -NMR spectrum of (2S,5R)-9 in CDCl_3 (500 MHz, 298K)



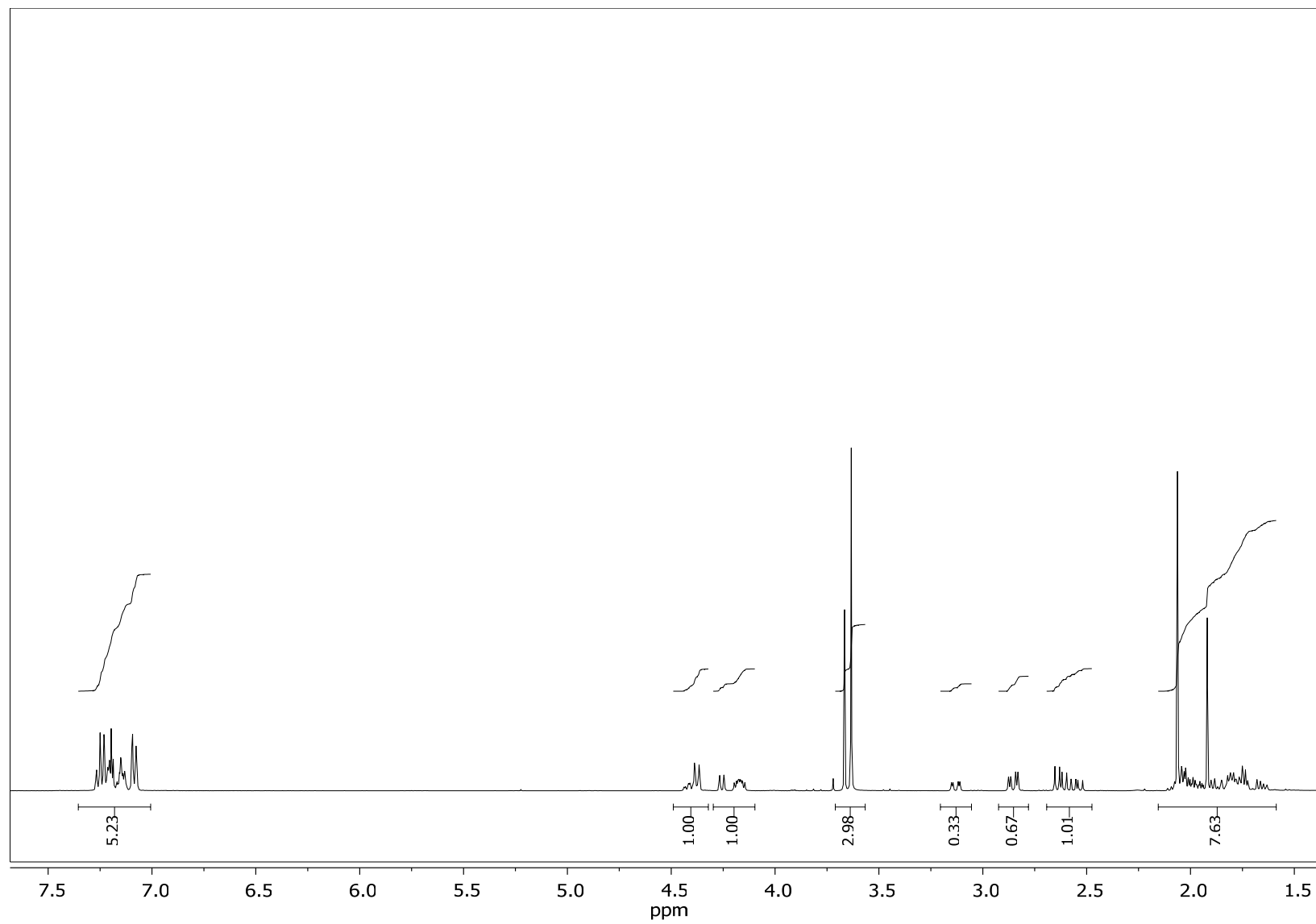
^{13}C -NMR spectrum of (2S,5R)-9 in CDCl_3 (75 MHz, 298K)



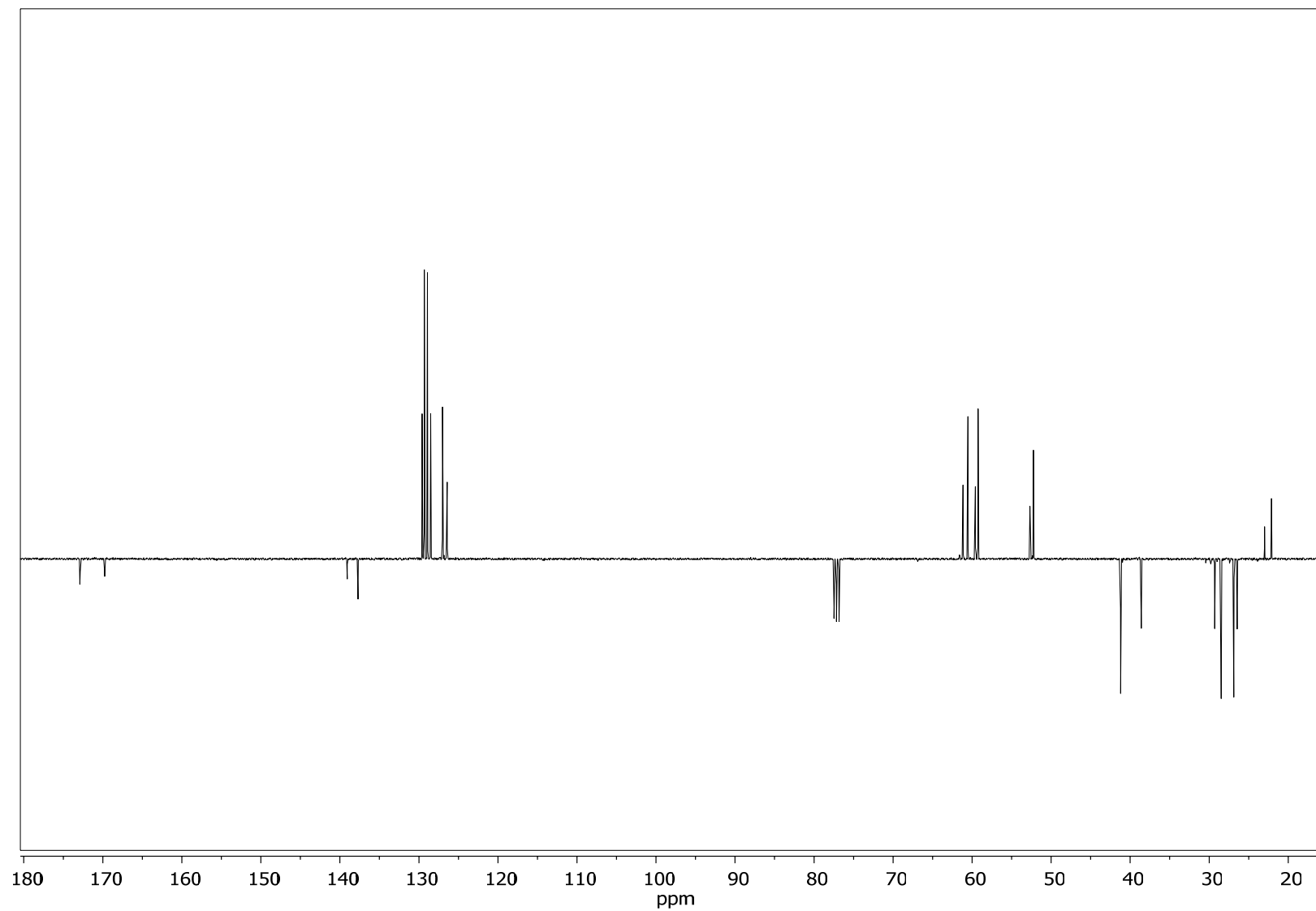
^1H -NMR spectrum of (2*R*,5*R*)-**9** in CDCl_3 (500 MHz, 298K)



^{13}C -NMR spectrum of **(2R,5R)-9** in CDCl_3 (75 MHz, 298K)



^1H -NMR spectrum of (2R,5R)-10 in CDCl_3 (400 MHz, 298K)



^{13}C -NMR spectrum of (2*R*,5*R*)-**10** in CDCl_3 (100 MHz, 298K)