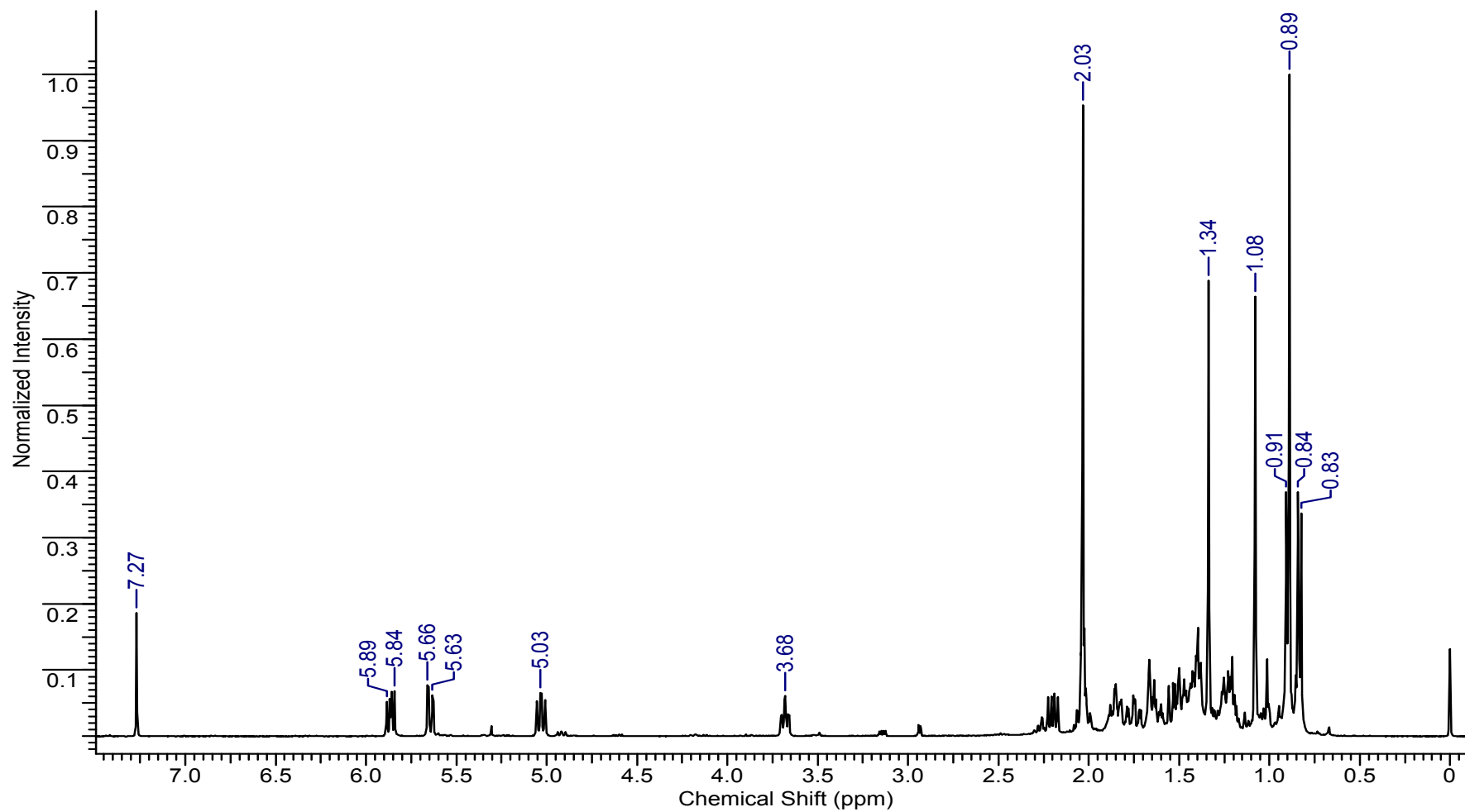


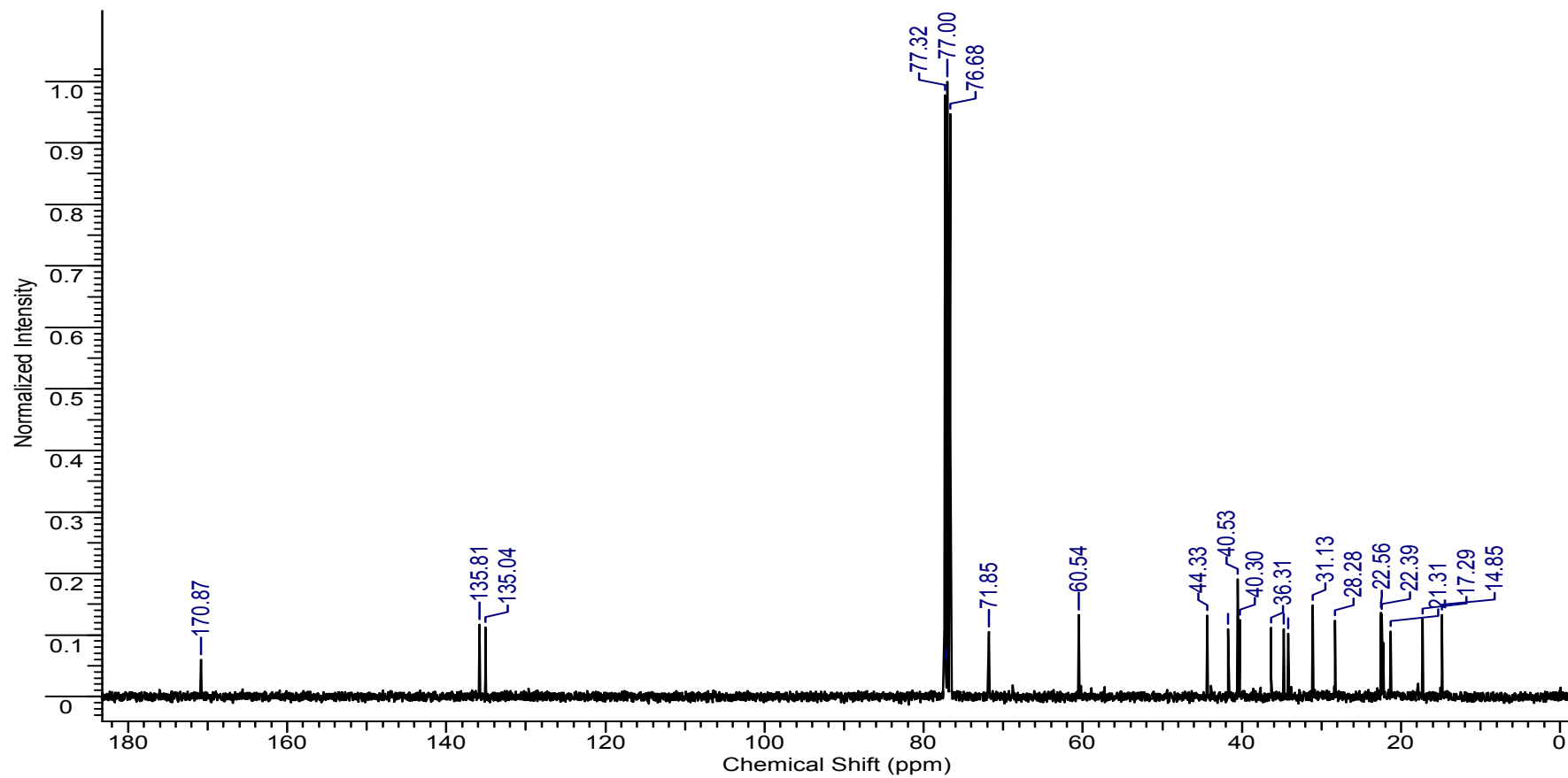
Electronic supplementary Information

**An unusual mulinane diterpenoid from the Chilean
plant *Azorella trifurcata* (Gaertn) Pers.**

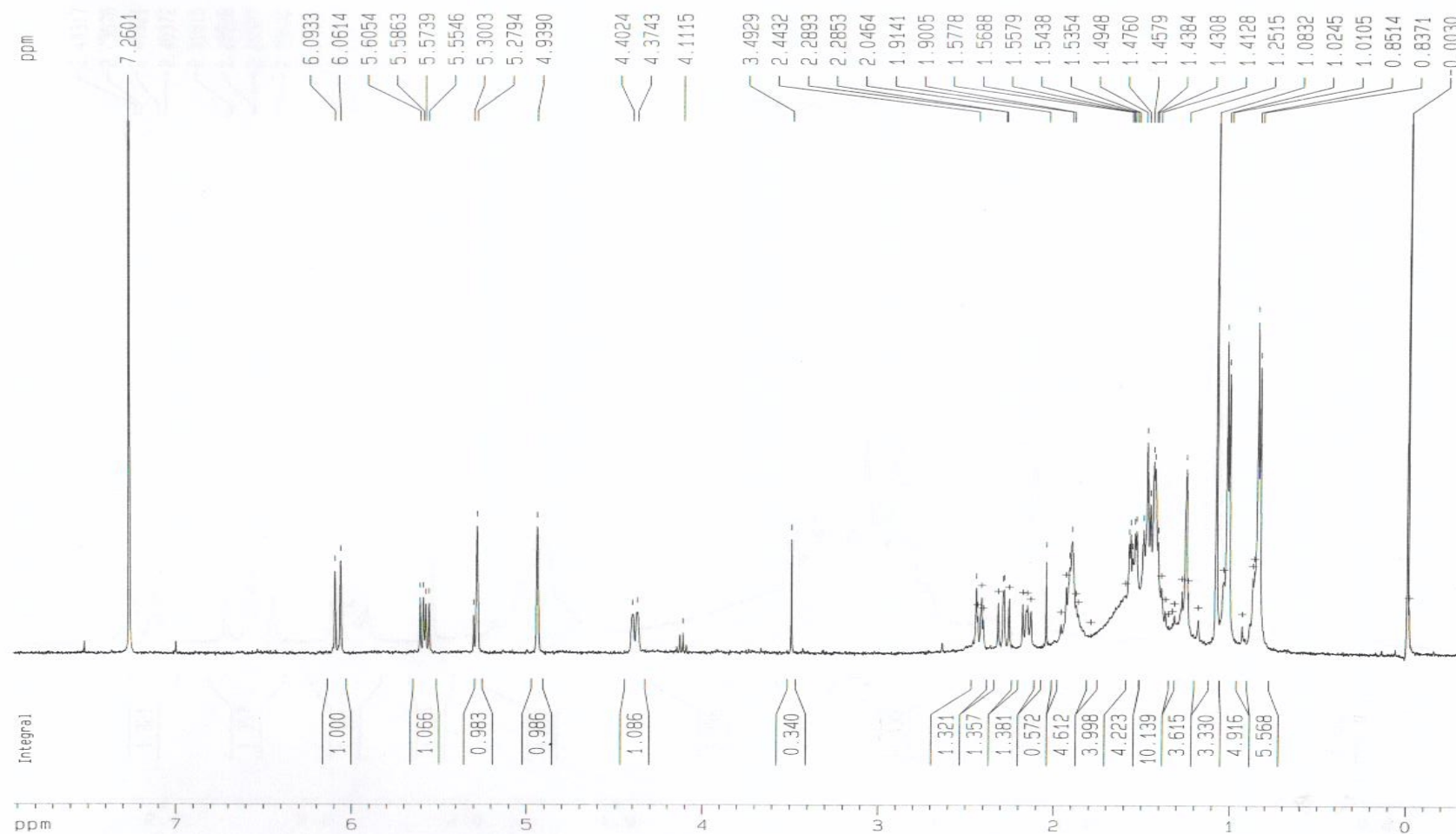
**Carlos Areche,^{*a} Beatriz Sepulveda,^b Aurelio San Martin,^a Olimpo
Garcia-Beltrán,^{a,c} Mario Simirgiotis,^d Alvaro Cañete,^e**



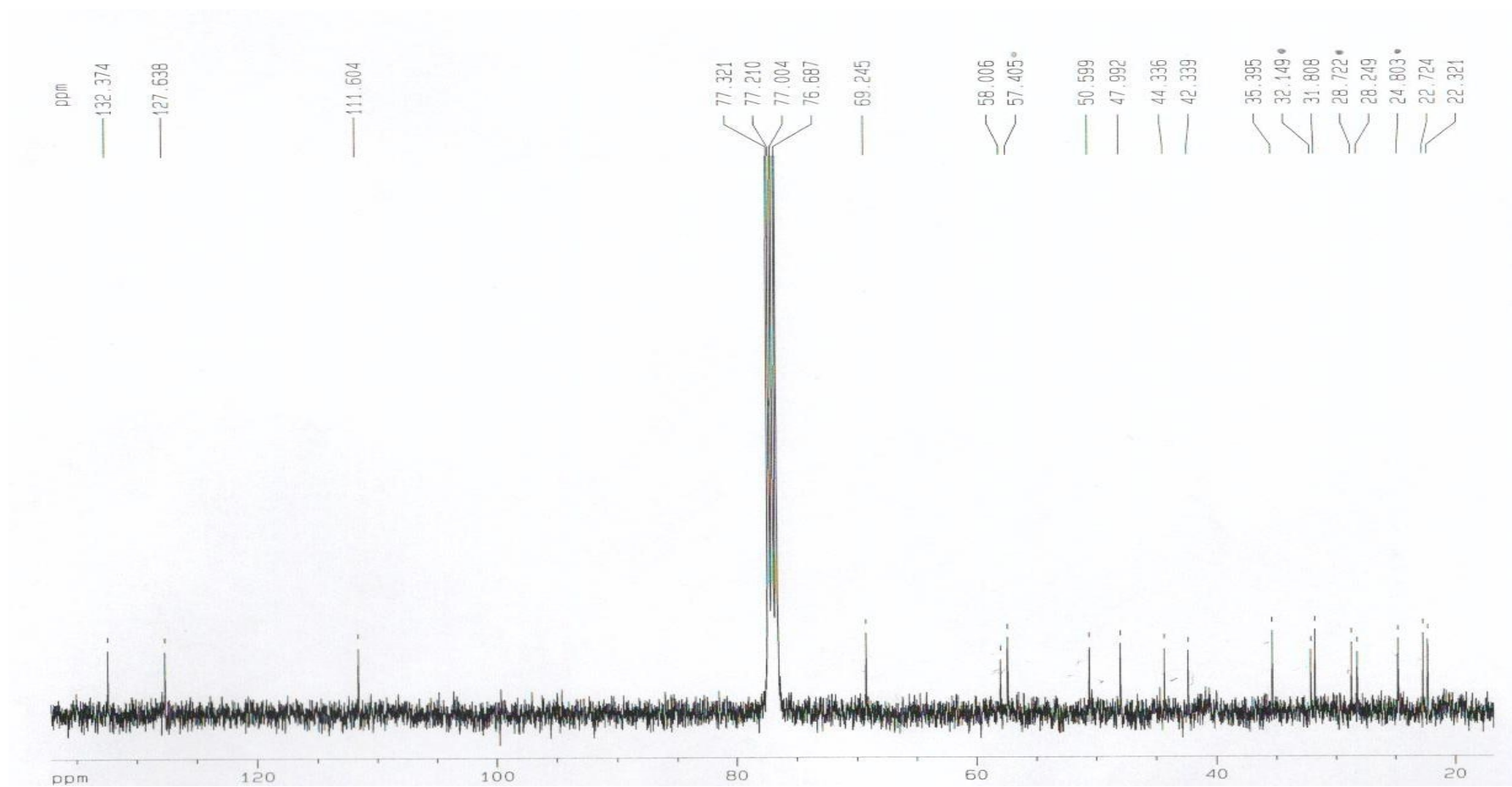
1. ^1H NMR spectrum of 7 α -acetoxy-9-epi-13 β -hydroxymulinane **4** in CDCl_3 .



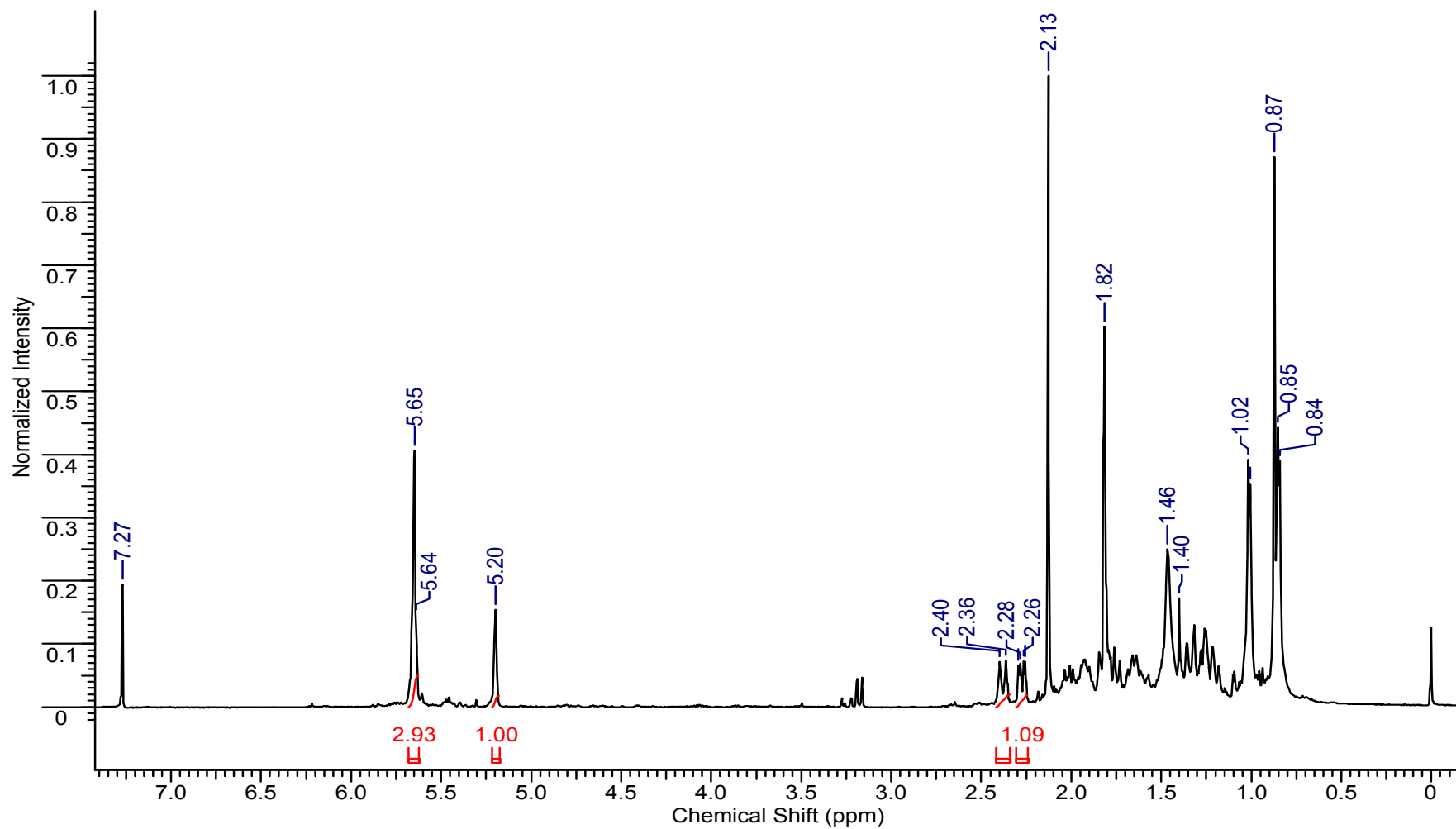
2. ¹³C NMR spectrum of 7 α -acetoxy-9-epi-13 β -hydroxymulinane **4** in CDCl₃.



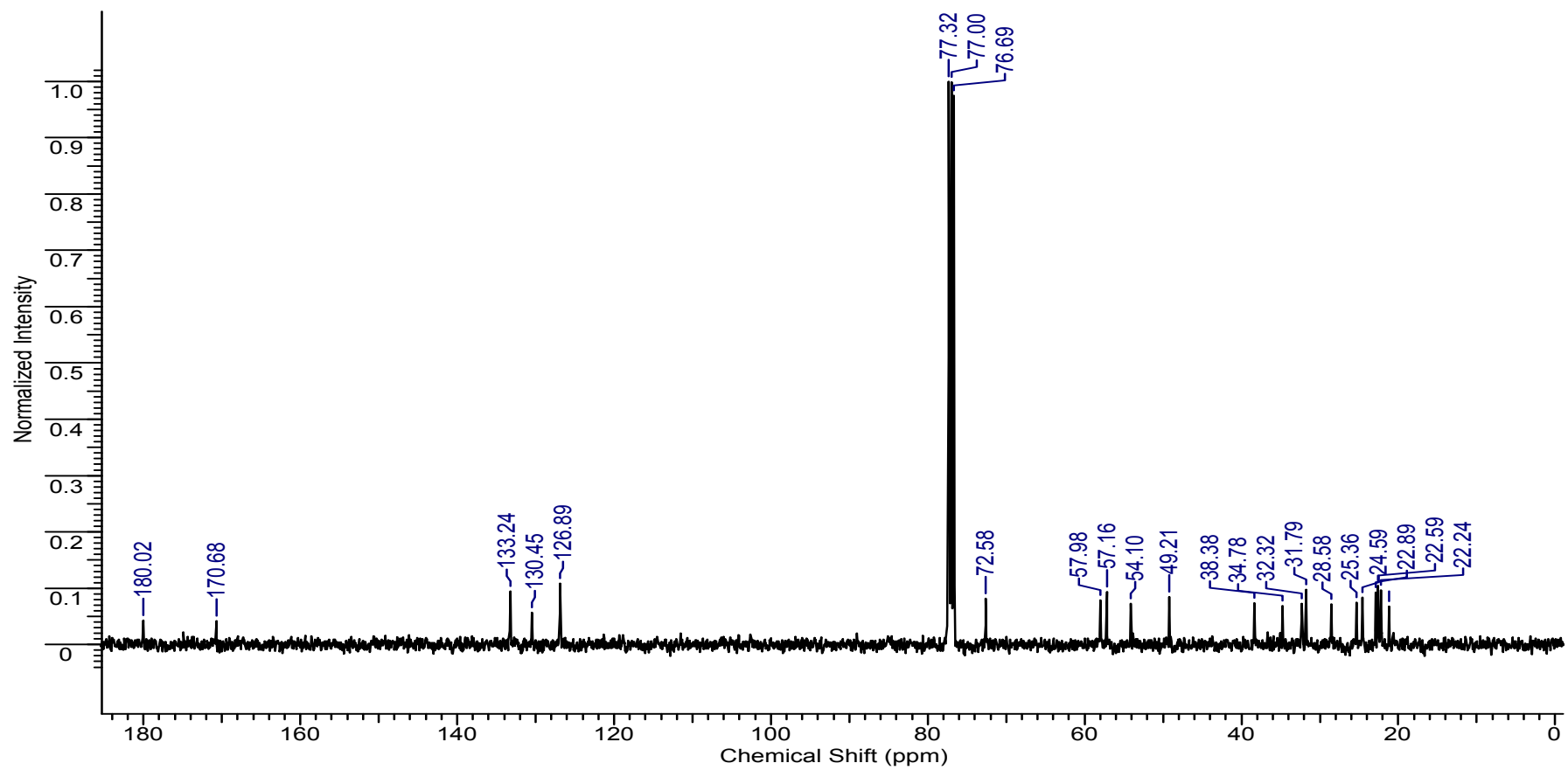
3. ¹H NMR spectrum of 14 α -hydroxymulin-11,13(16)-dien-20-oic acid **5** in CDCl₃.



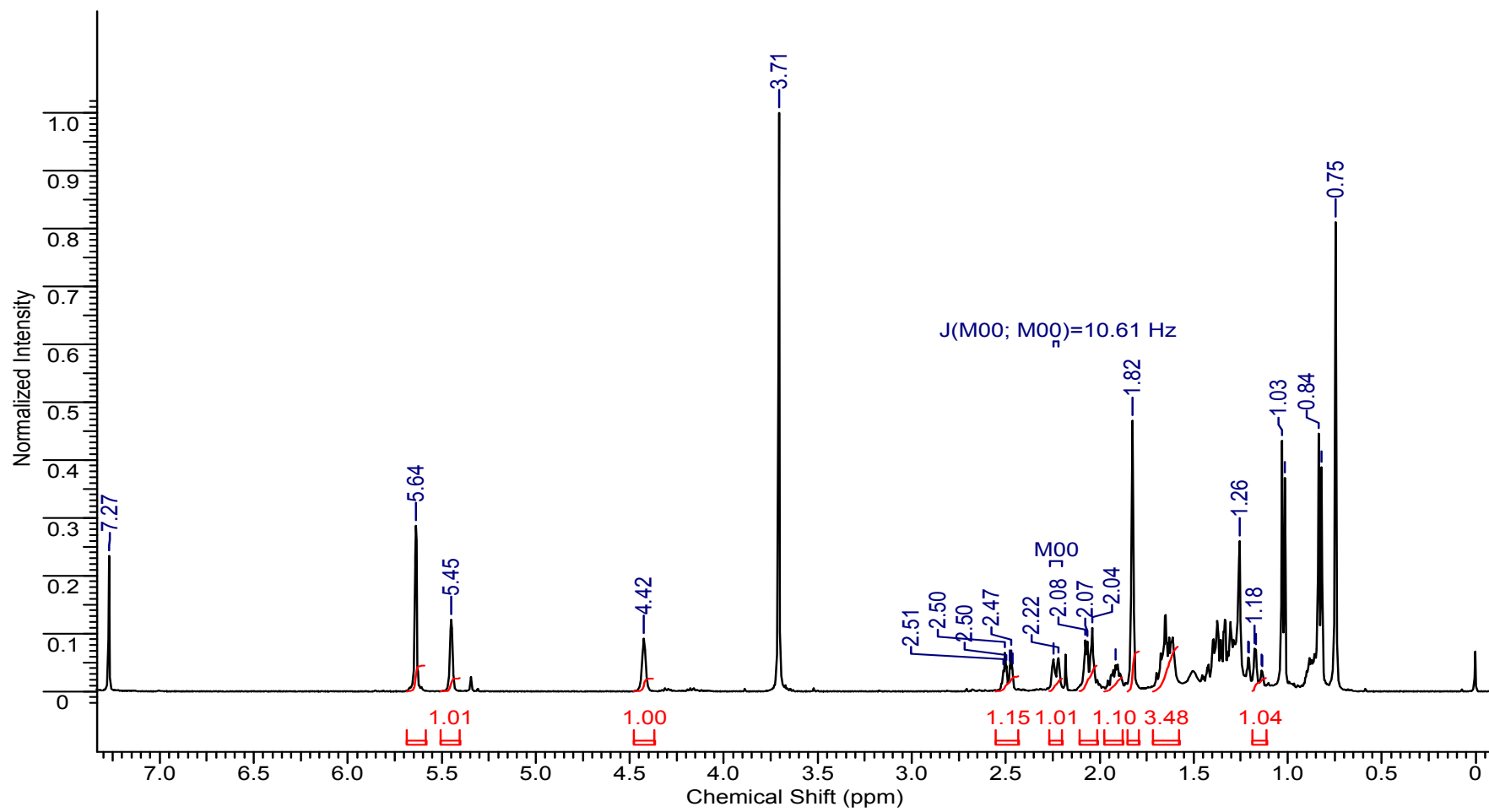
4. ¹³C NMR spectrum of 14 α -hydroxymulin-11,13(16)-dien-20-oic acid **5** in CDCl₃.



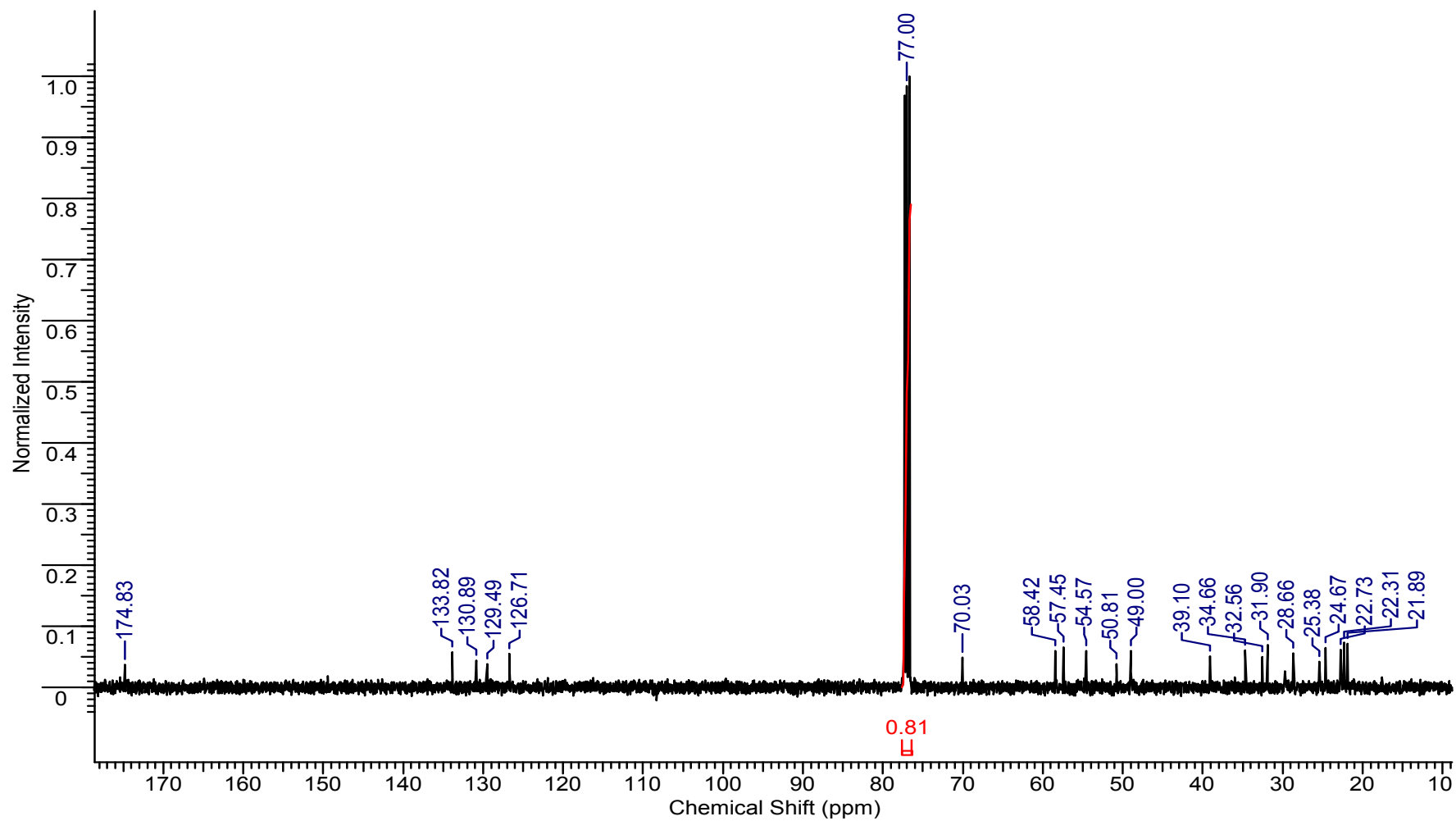
5. ^1H NMR spectrum of 15 α -acetoxymulin-11,13-dien-20-oic acid **8** in CDCl_3 .



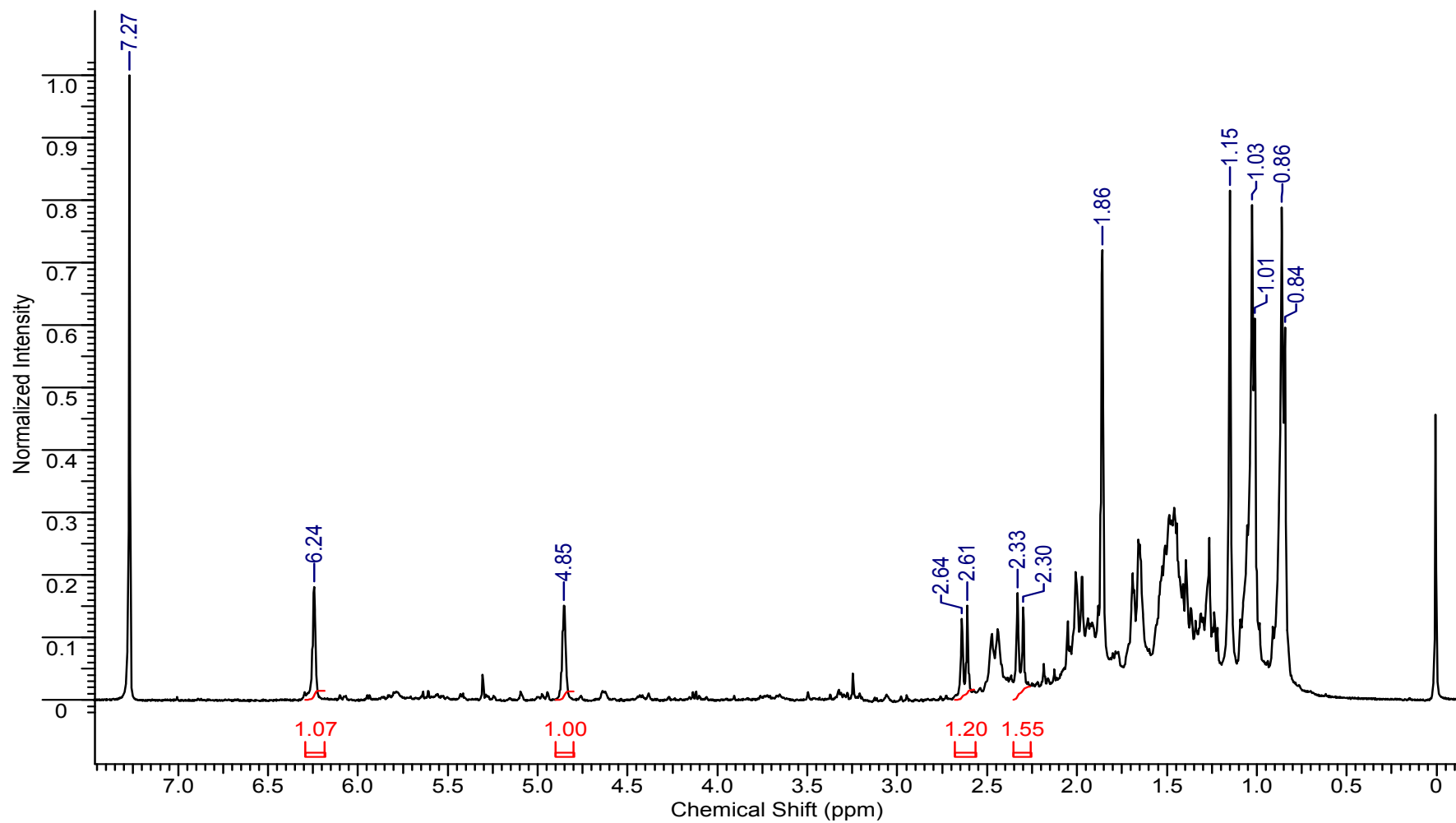
6. ¹³C NMR spectrum of 15 α -acetoxymulin-11,13-dien-20-oic acid **8** in CDCl₃.



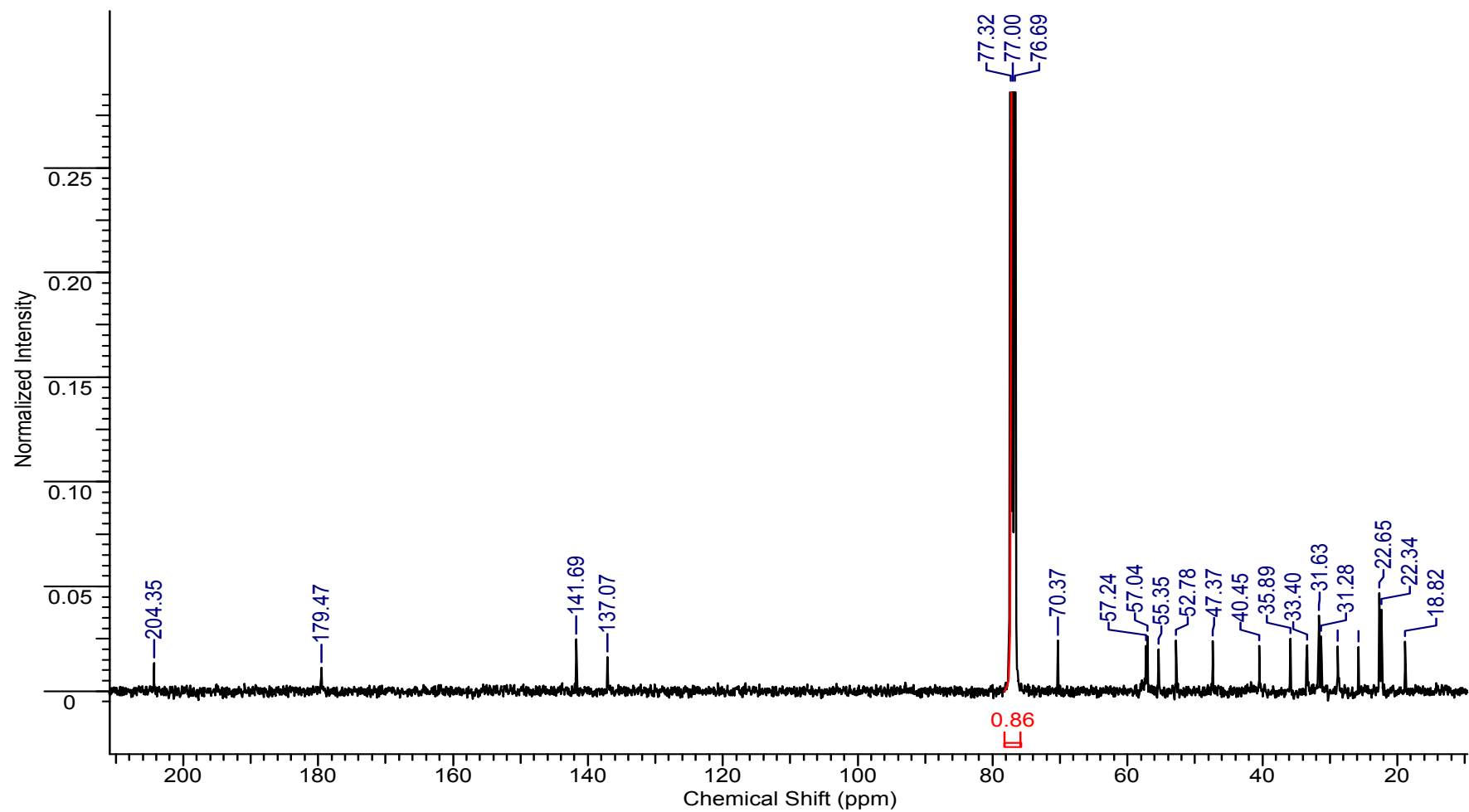
7. ^1H NMR spectrum of 15 α -hydroxymulin-11,13-dien-20-oic acid methyl ester **8a** in CDCl_3



8. ¹³C NMR spectrum of 15 α -hydroxymulin-11,13-dien-20-oic acid methyl ester **8a** in CDCl₃.



9. ^1H NMR spectrum of 11 α -hydroxymulin-12-en-14-one-20-oic acid **9** in CDCl_3 .



10. ¹³C NMR spectrum of 11 α -hydroxymulin-12-en-14-one-20-oic acid **9** in CDCl₃.

Preparation of **8a**

In a flask containing a stirred solution of 10 mg of the compound **8** in Et₂O (5 mL) at 10°C, ethereal diazomethane (0.5 mL) was added dropwise. The resulting solution was stirred for 3h. Then, the solution was concentrated in vacuo and monitored by TLC to give quantitatively the methyl ester. To a solution of 10 mg of this methyl ester in MeOH (5mL), K₂CO₃ (catalytic amount) was added. The resulting solution was stirred for 12h. After usual work-up, purification by CC on silica gel using an *n*-hexane/EtOAc 10:1 to 4:6 gradient gave compound **8a** (8mg).