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Acanthoic Acid and other Constituents from the Stem of *Annona amazonica* (Annonaceae)

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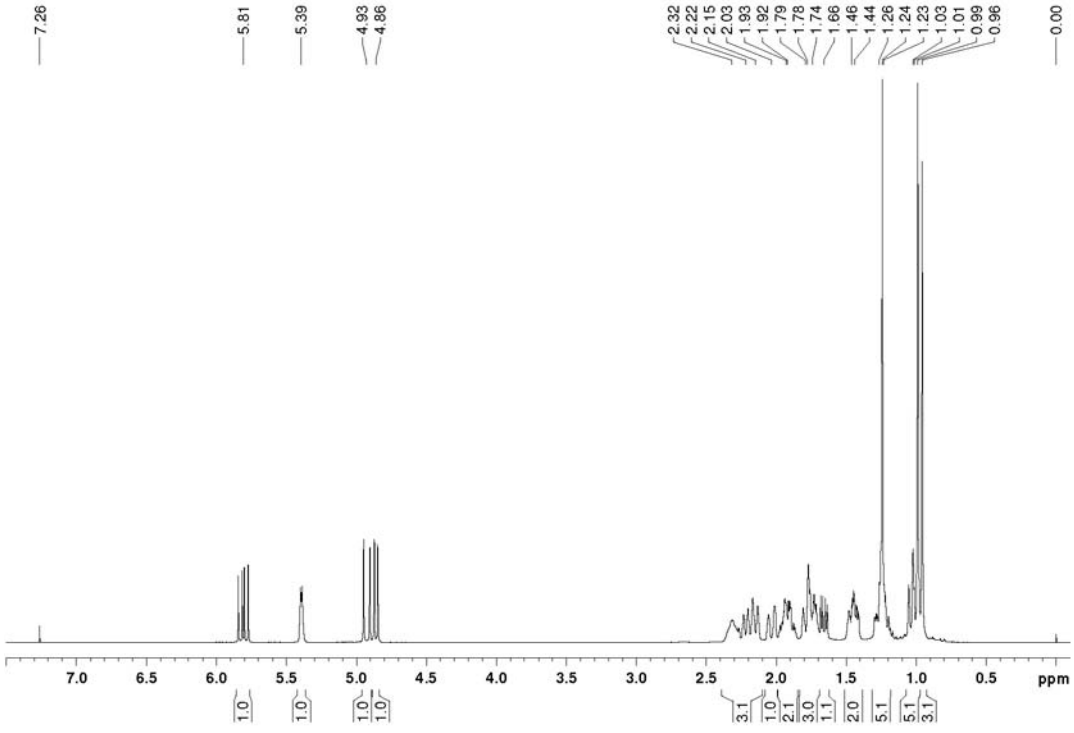

Figure S1. ^{1}H NMR spectrum of acanthoic acid in $CDCl_3$ at 400 MHz.

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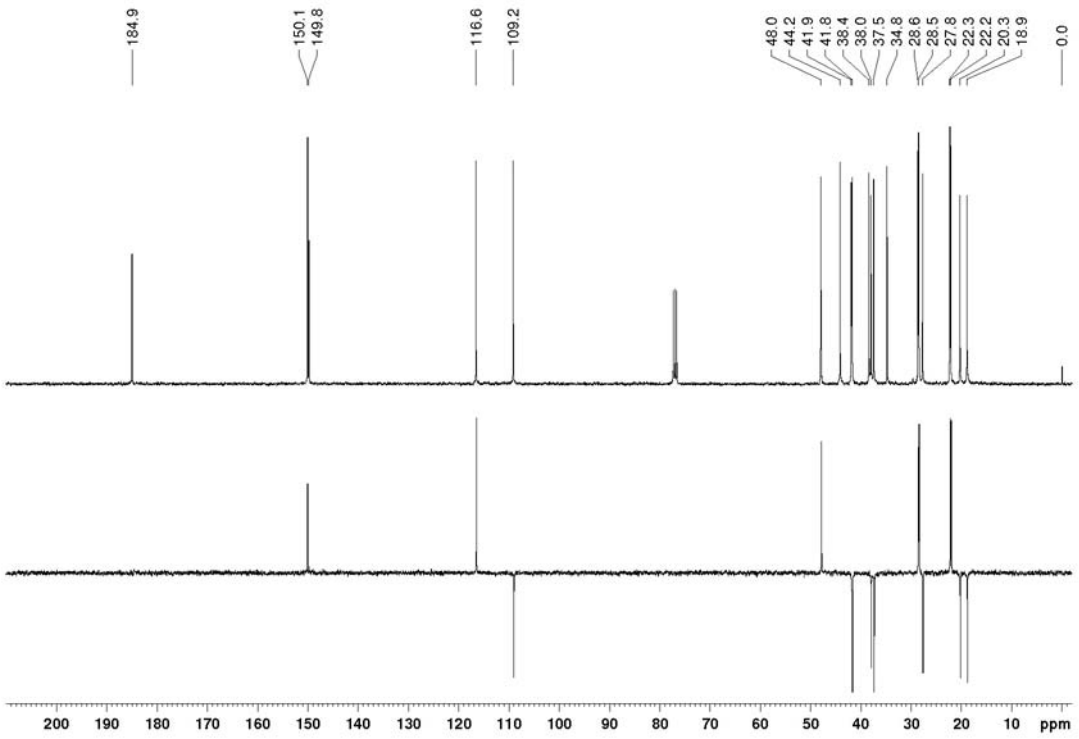

Figure S2. $^{13}C\{^{1}H\}$ and DEPT 135 NMR spectra of acanthoic acid in $CDCl_3$ at 100 MHz.

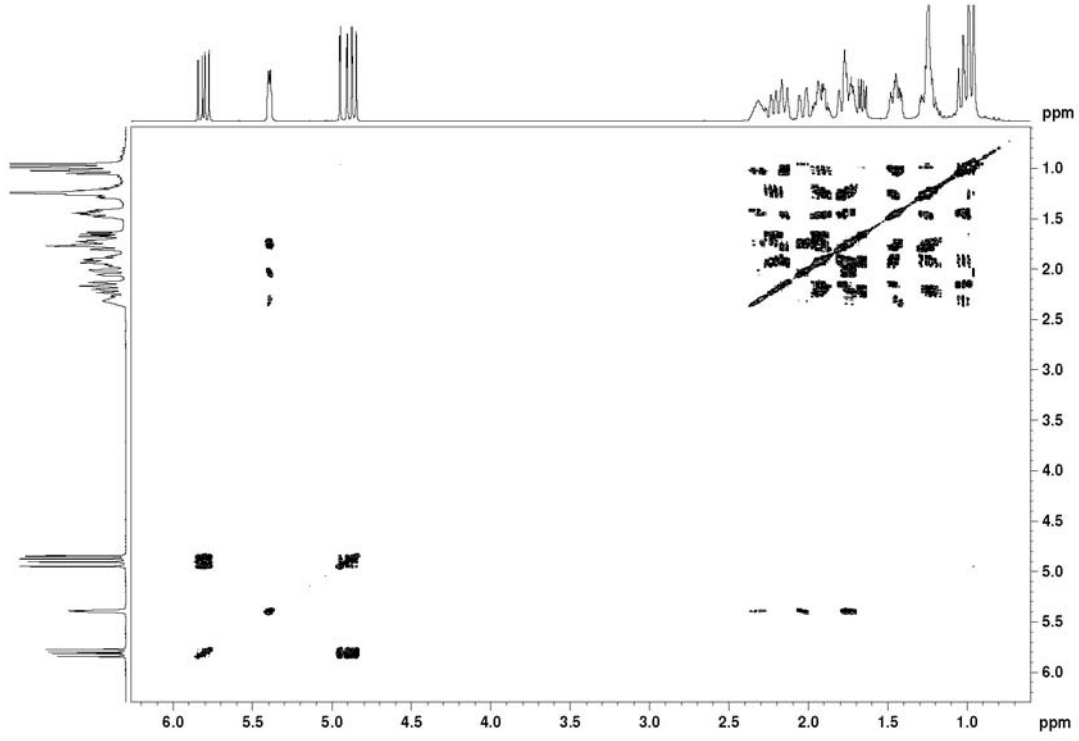

Figure S3. ^{1}H-^{1}H correlation map from the COSY NMR experiment on acanthoic acid in $CDCl_3$ at 400 MHz.

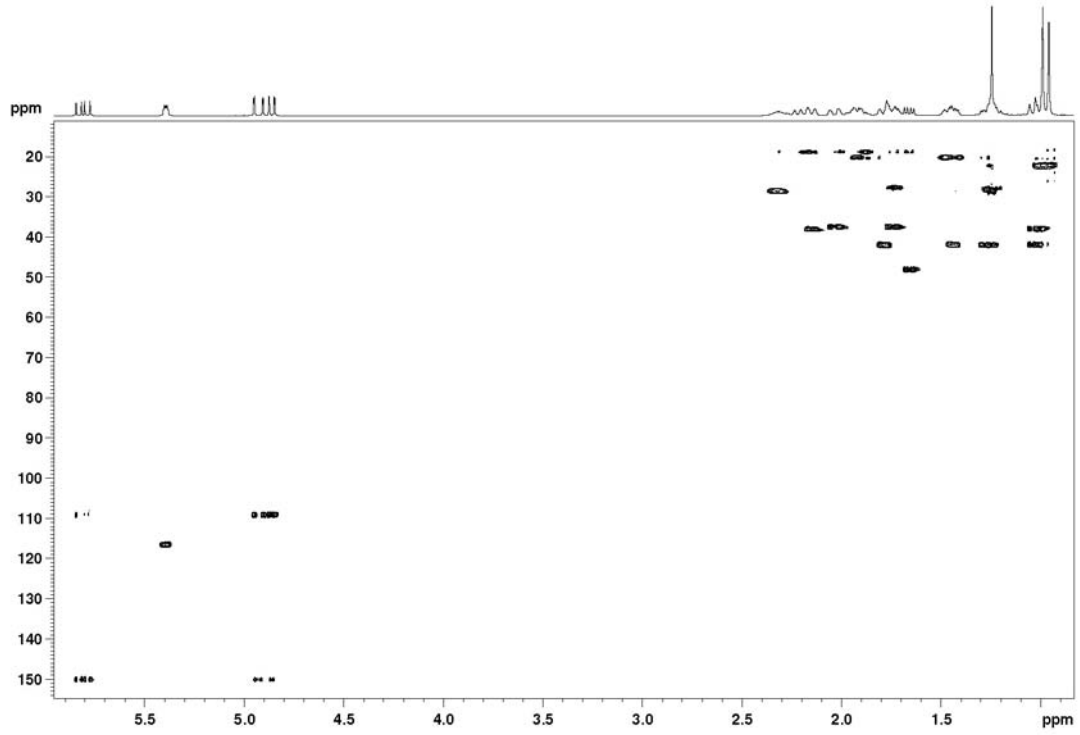

Figure S4. ^{1}H-^{13}C one-bond correlation map from the HSQC NMR experiment on acanthoic acid in $CDCl_3$ at 400 and 100 MHz.

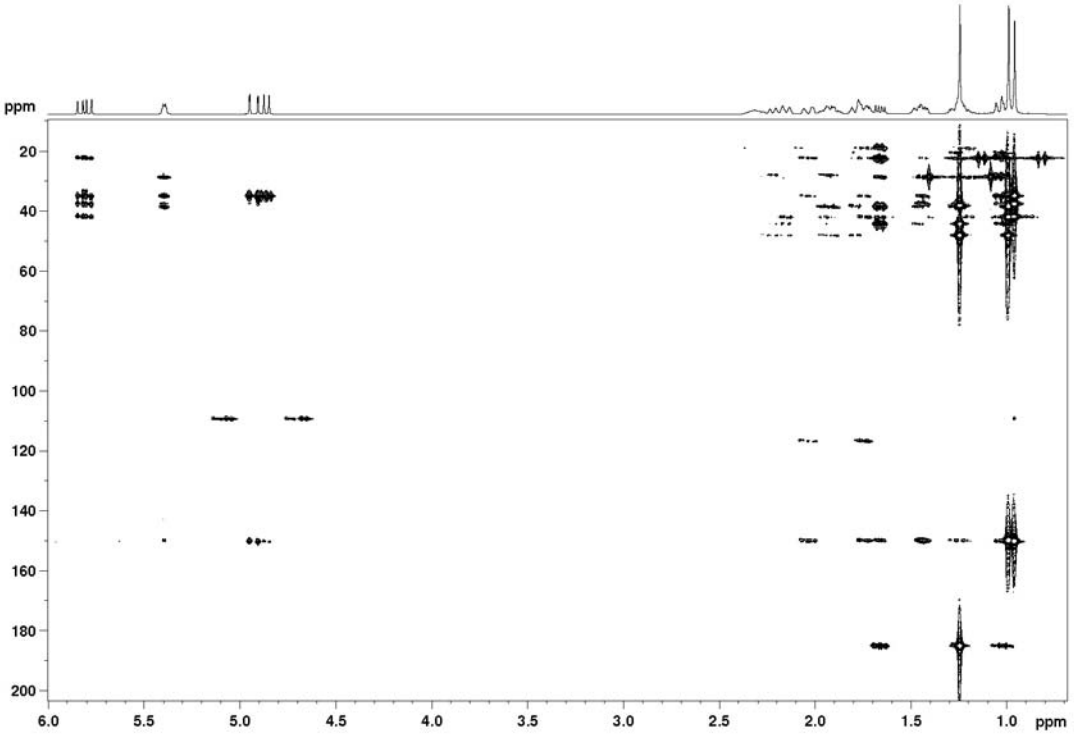

Figure S5. ^{1}H-^{13}C long-range correlation map from the HMBC NMR experiment on acanthoic acid in $CDCl_3$ at 400 and 100 MHz.

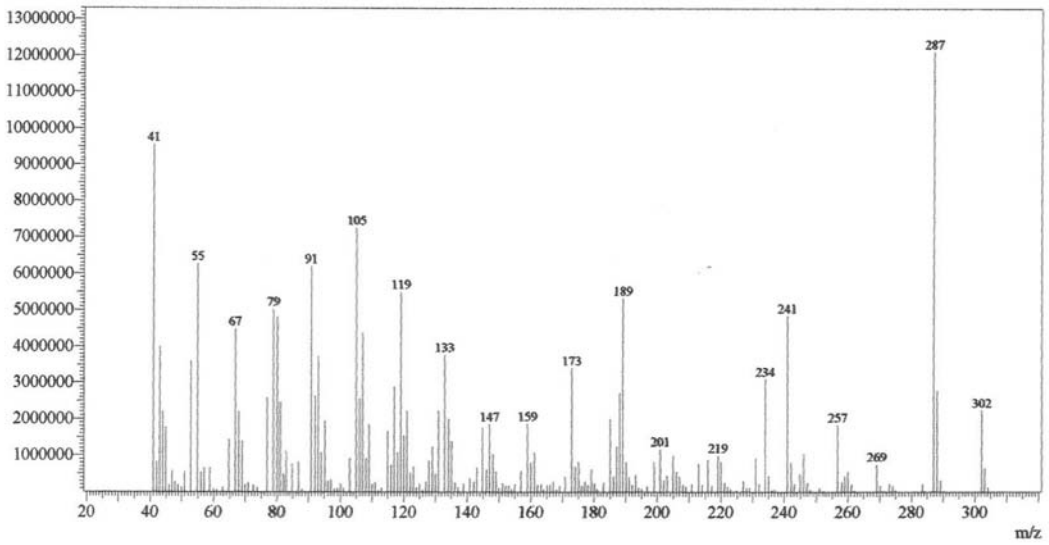

Figure S6. EI-MS of acanthoic acid.

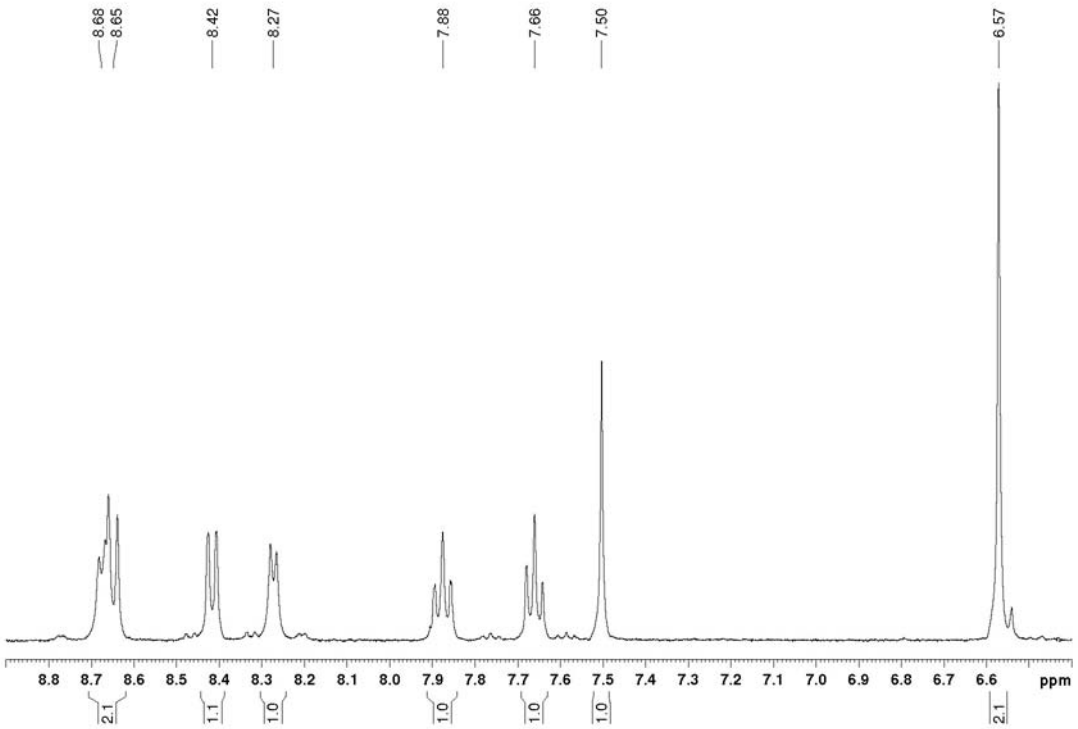

Figure S7. ^{1}H NMR spectrum of liriodenine in MeOD-d_4 at 400 MHz.

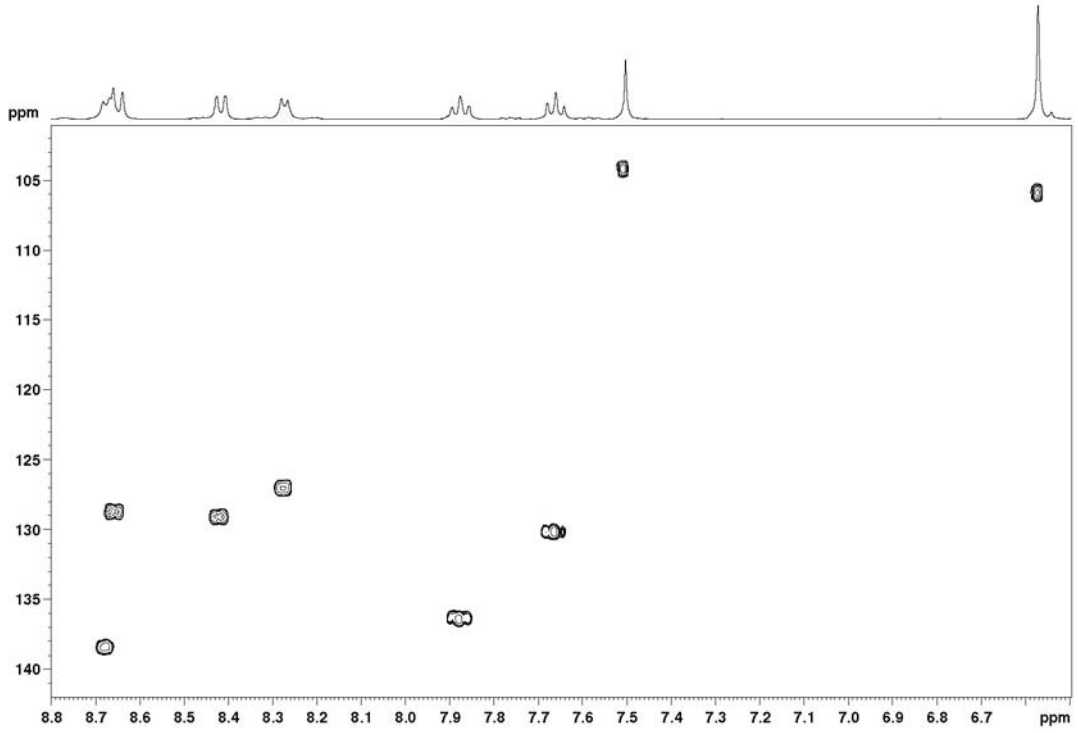

Figure S8. ^{1}H-^{13}C one-bond correlation map from the HSQC NMR experiment on liriodenine in MeOD-d_4 at 400 and 100 MHz.

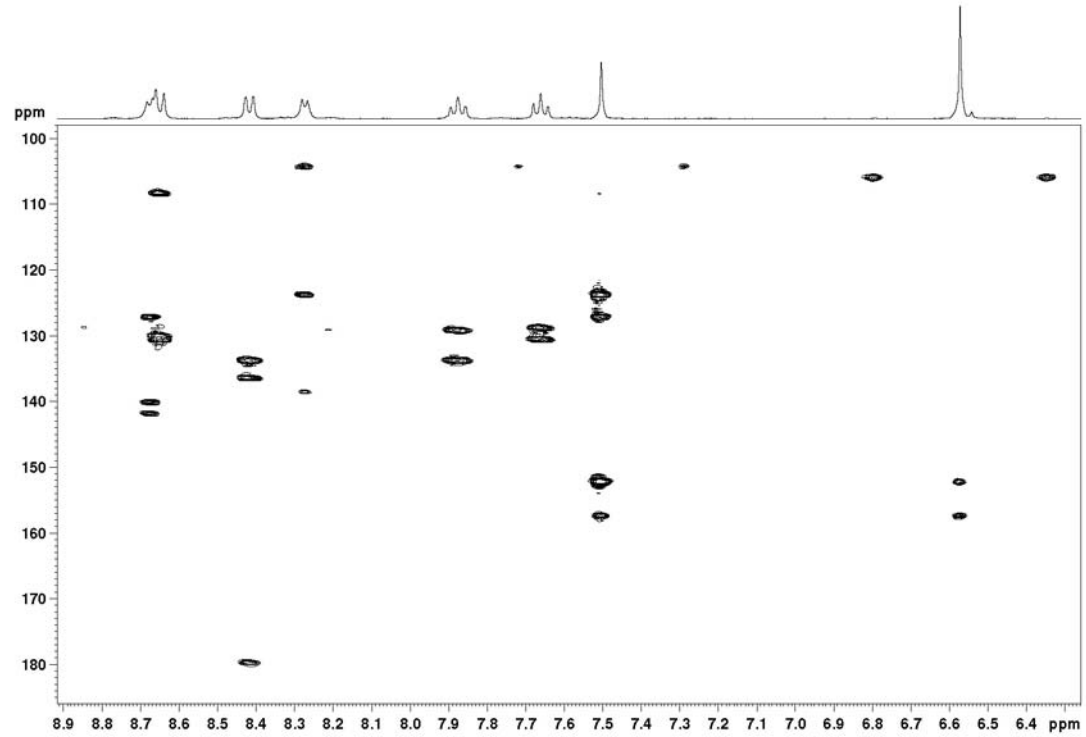

Figure S9. ^{1}H-^{13}C long-range correlation map from the HMBC NMR experiment on liriodenine in MeOD-d_4 at 400 and 100 MHz.

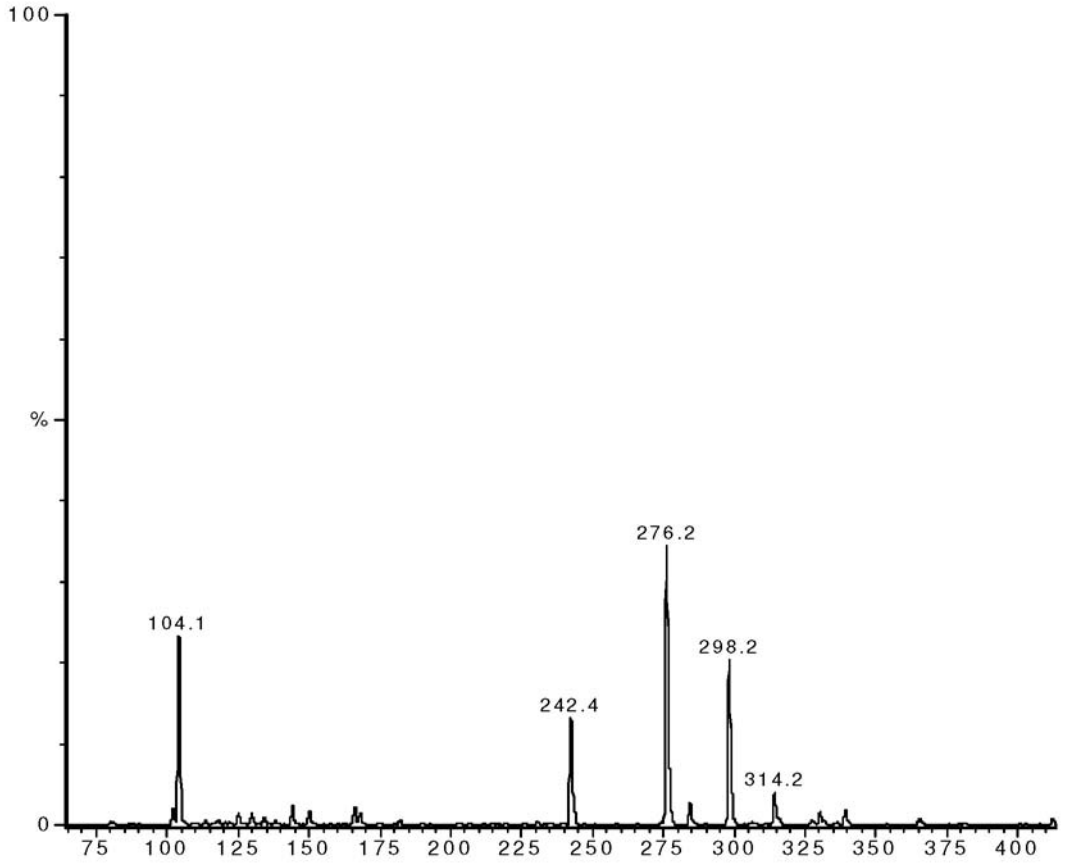

Figure S10. ESI-MS of liriodenine in MeOH, positive mode.

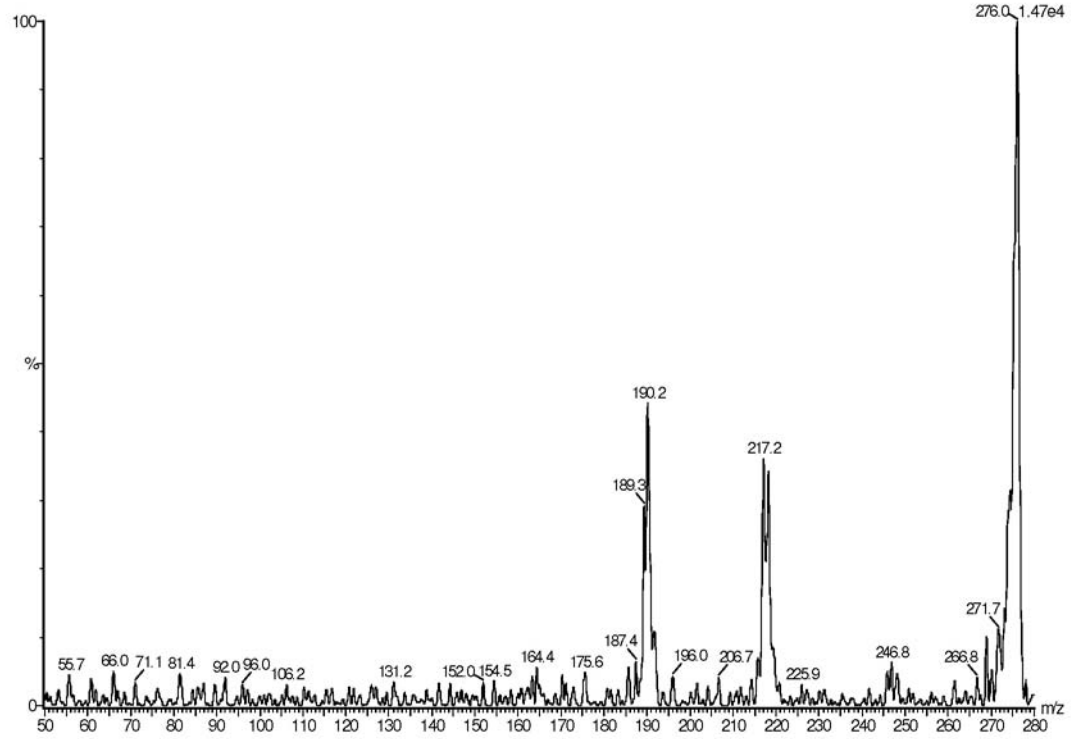

Figure S11. ESI-MS/MS of liriodenine in MeOH, positive mode at 50 eV.

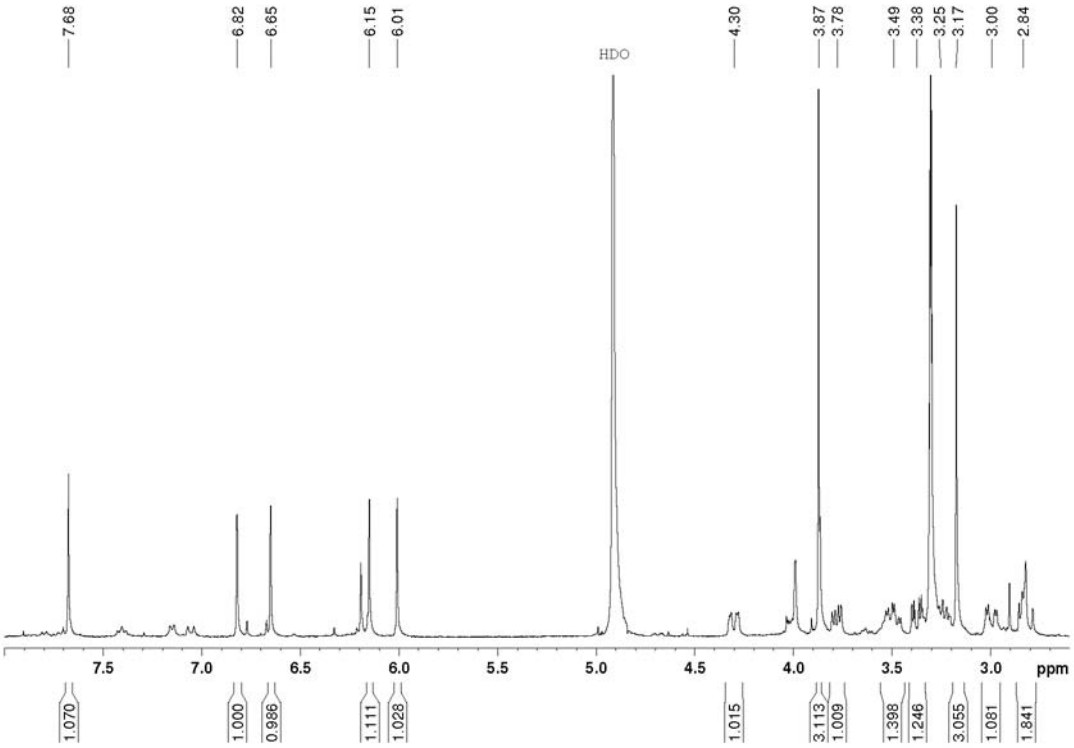

Figure S12. ^{1}H NMR spectrum of cassythicine in MeOD-d_4 at 400 MHz.

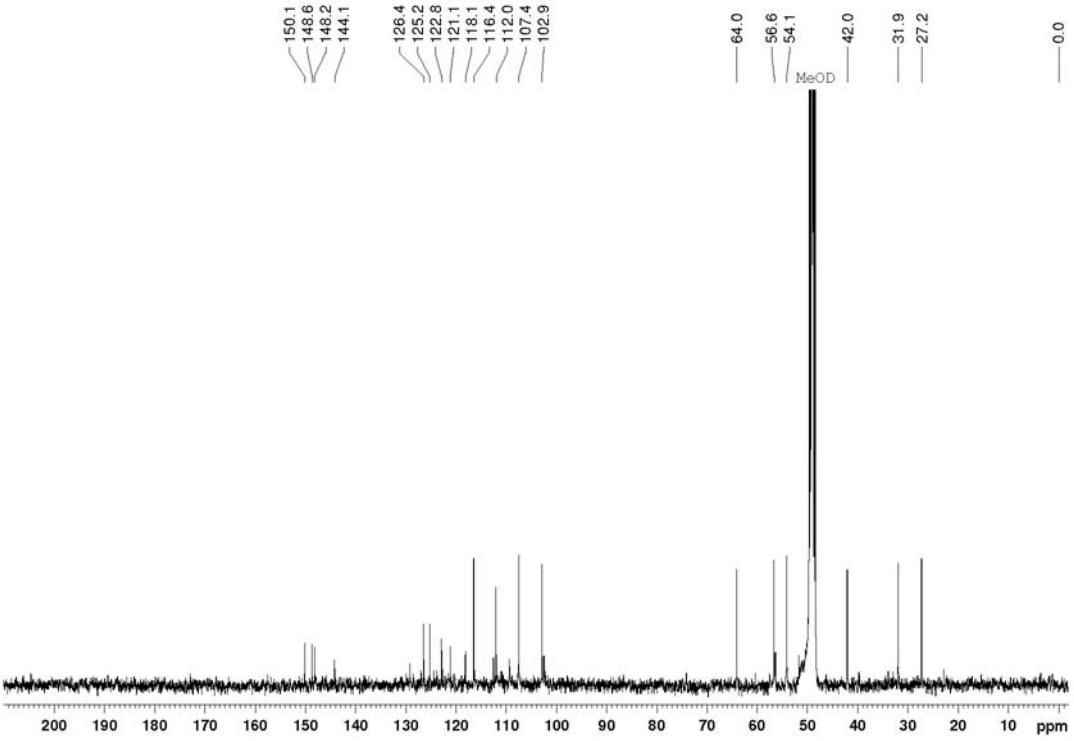

Figure S13. ^{13}C{^{1}H} NMR spectrum of cassythicine in MeOD-d_4 at 100 MHz.

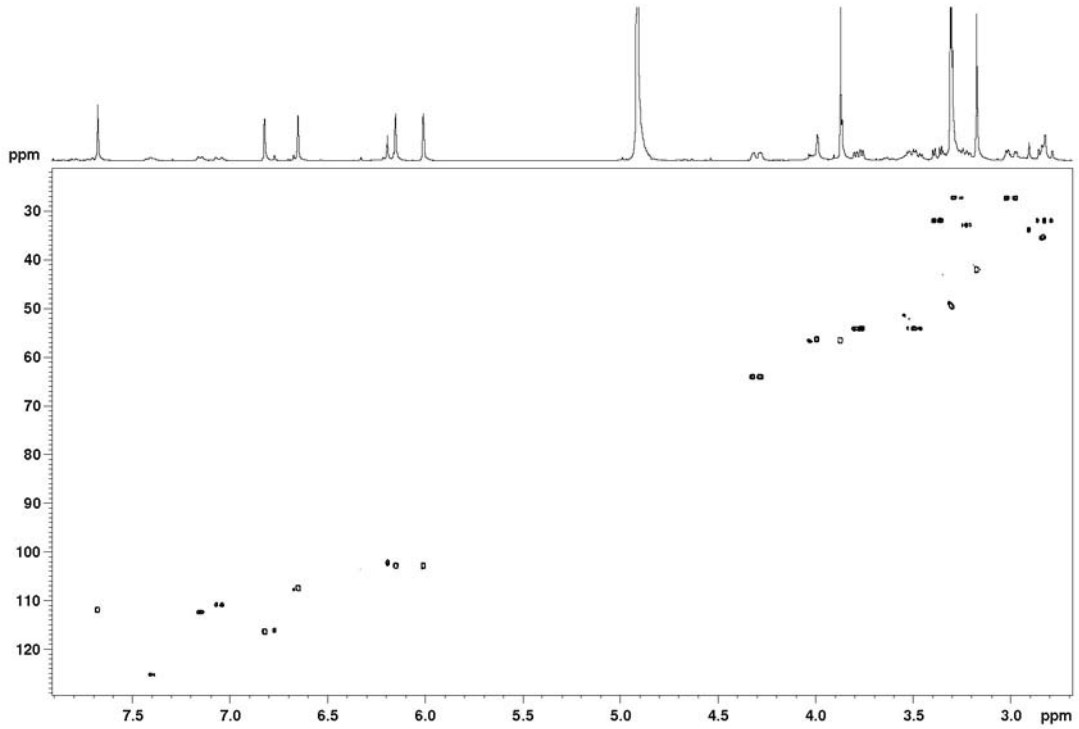

Figure S14. ^{1}H-^{13}C one-bond correlation map from the HSQC NMR experiment on cassythicine in MeOD-d_4 at 400 and 100 MHz.

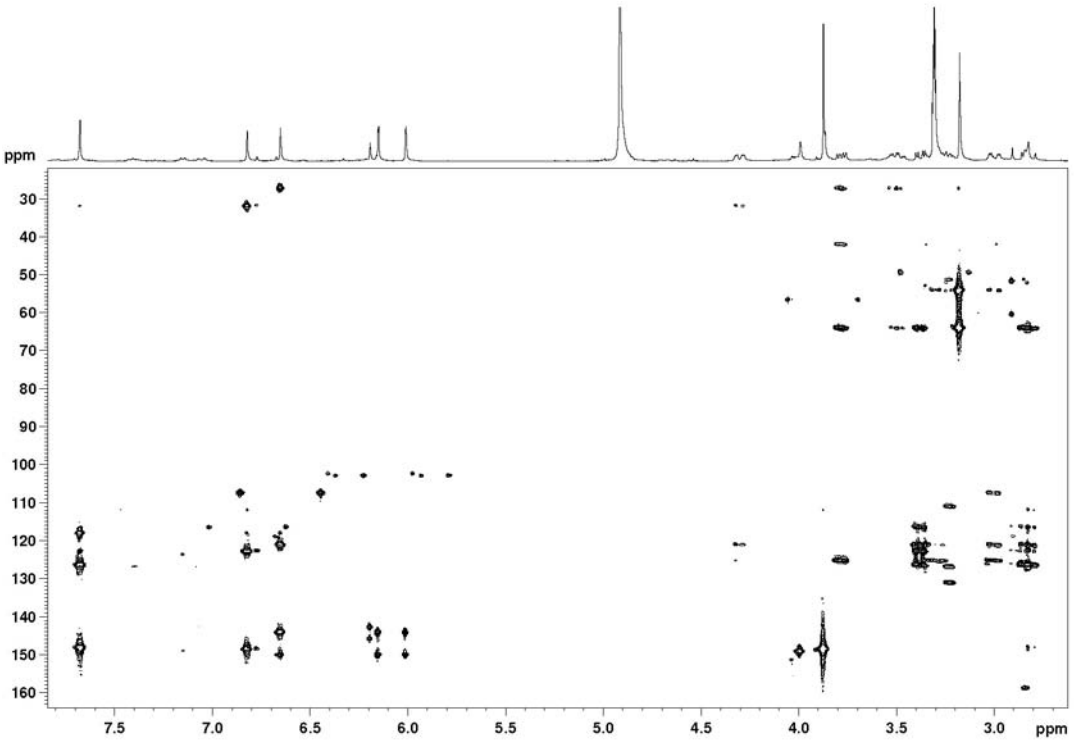

Figure S15. ^{1}H-^{13}C long-range correlation map from the HMBC NMR experiment on cassythicine in MeOD-d_4 at 400 and 100 MHz.

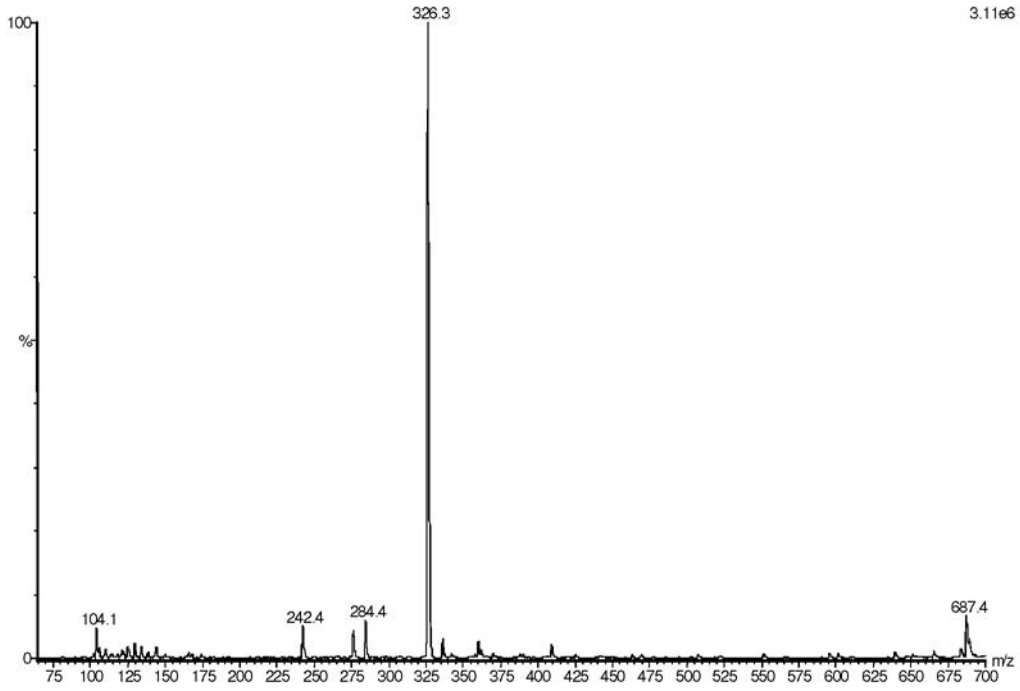

Figure S16. ESI-MS of cassythicine in MeOH, positive mode.

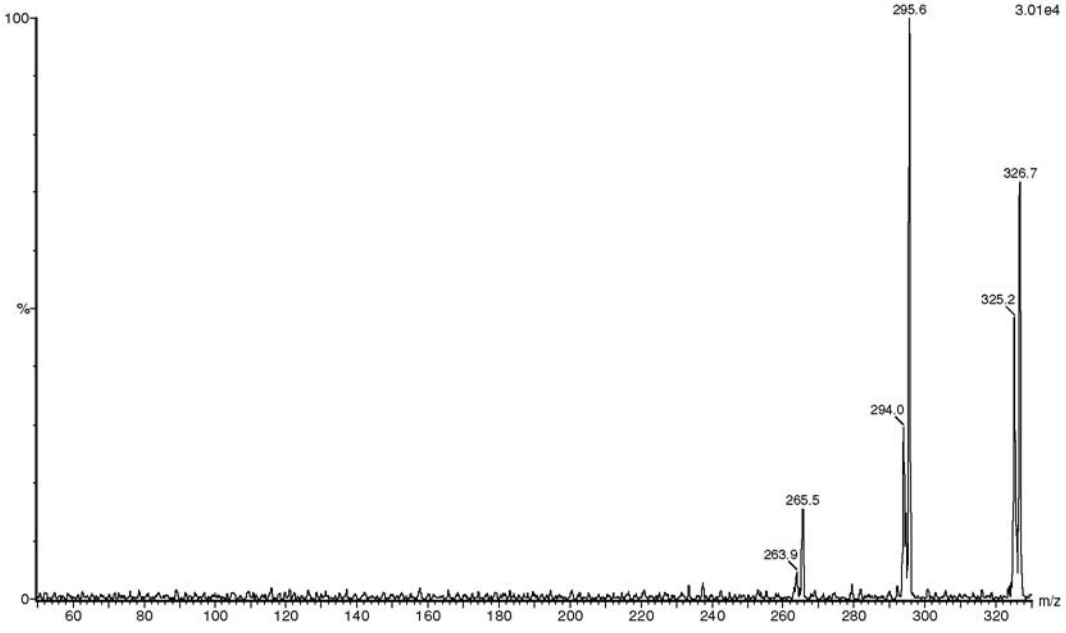

Figure S17. ESI-MS/MS of cassythicine in MeOH, positive mode at 50 eV.

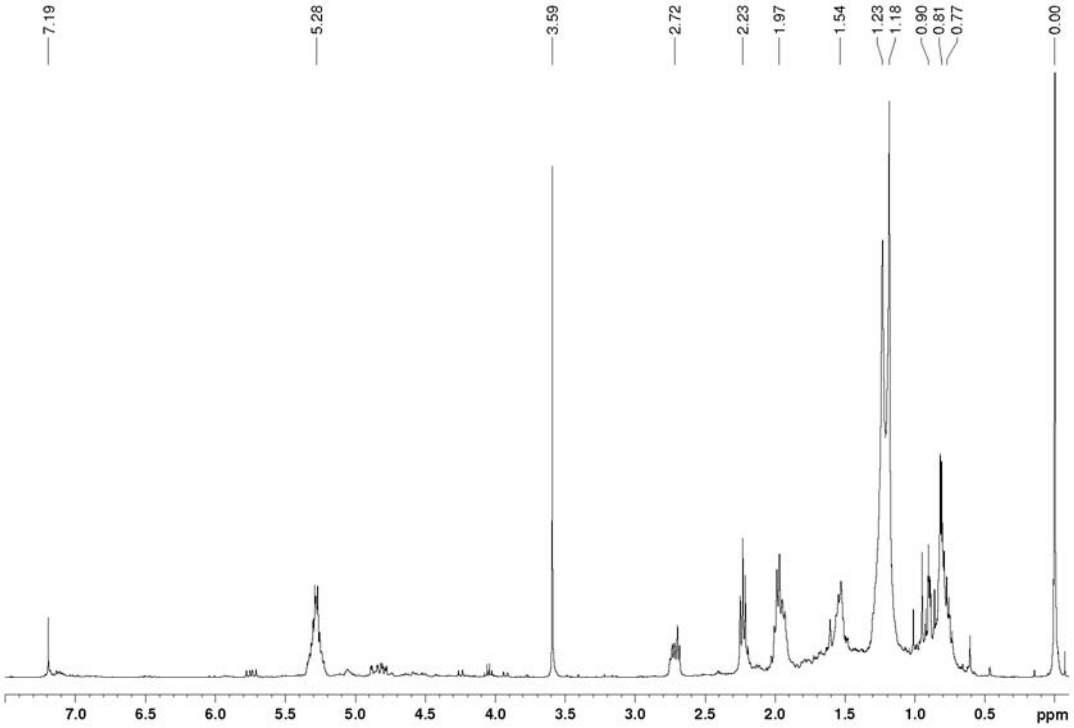

Figure S18. ^{1}H NMR spectrum of the mixture of three methyl esters of the fatty acids, oleic, linoleic, and linolenic in $CDCl_3$ at 400 MHz.

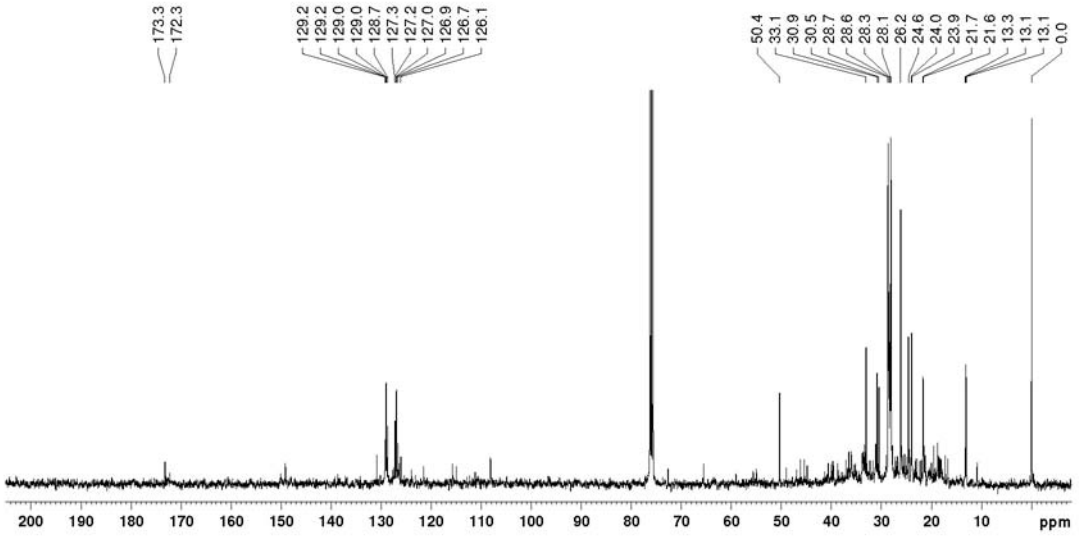

Figure S19. $^{13}C\{^{1}H\}$ spectrum of the mixture of three methyl esters of the fatty acids, oleic, linoleic, and linolenic in $CDCl_3$ at 100 MHz.

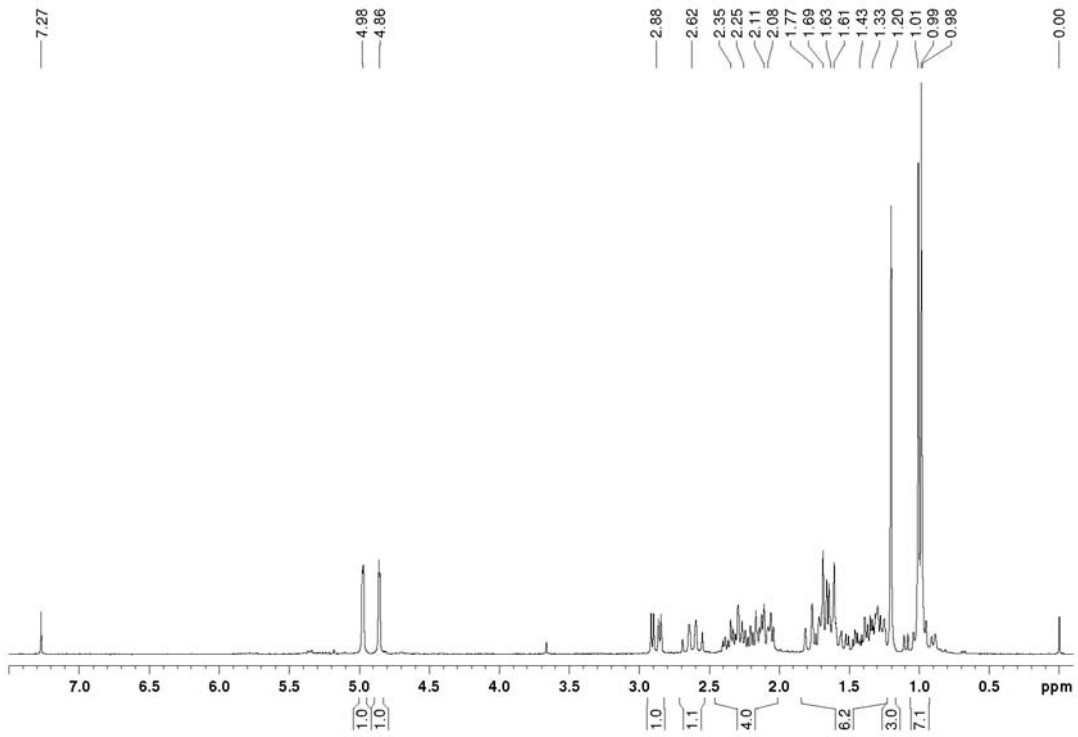

Figure S20. ^{1}H NMR spectrum of caryophyllene oxide in $CDCl_3$ at 200 MHz.

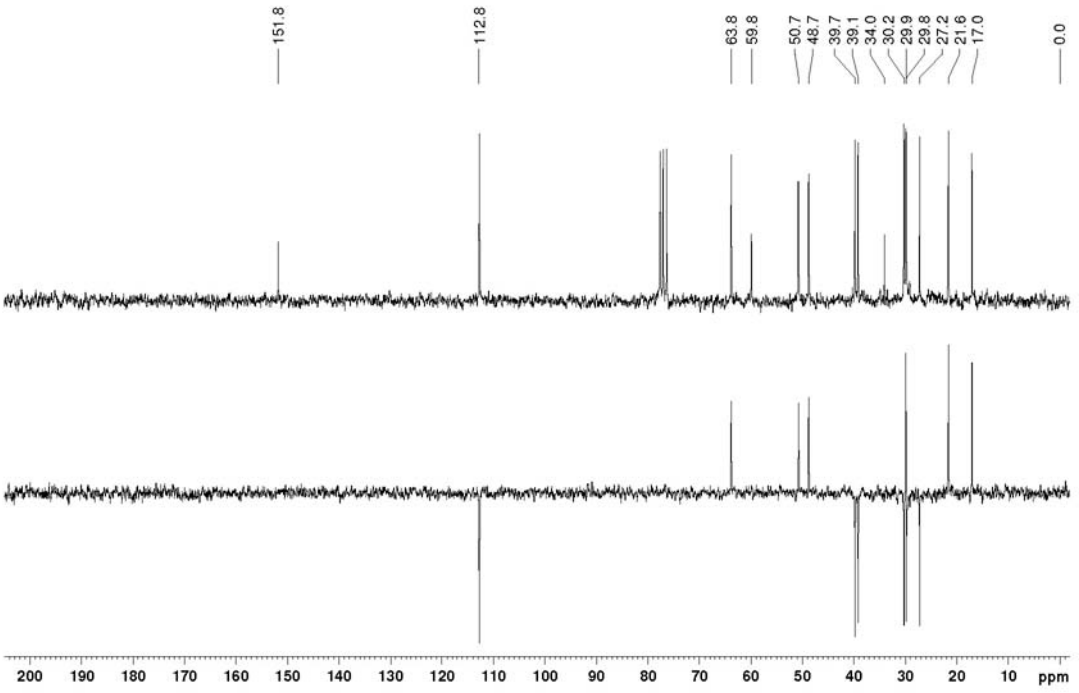

Figure S21. $^{13}C\{^{1}H\}$ and DEPT 135 NMR spectra of caryophyllene oxide in $CDCl_3$ at 50 MHz.

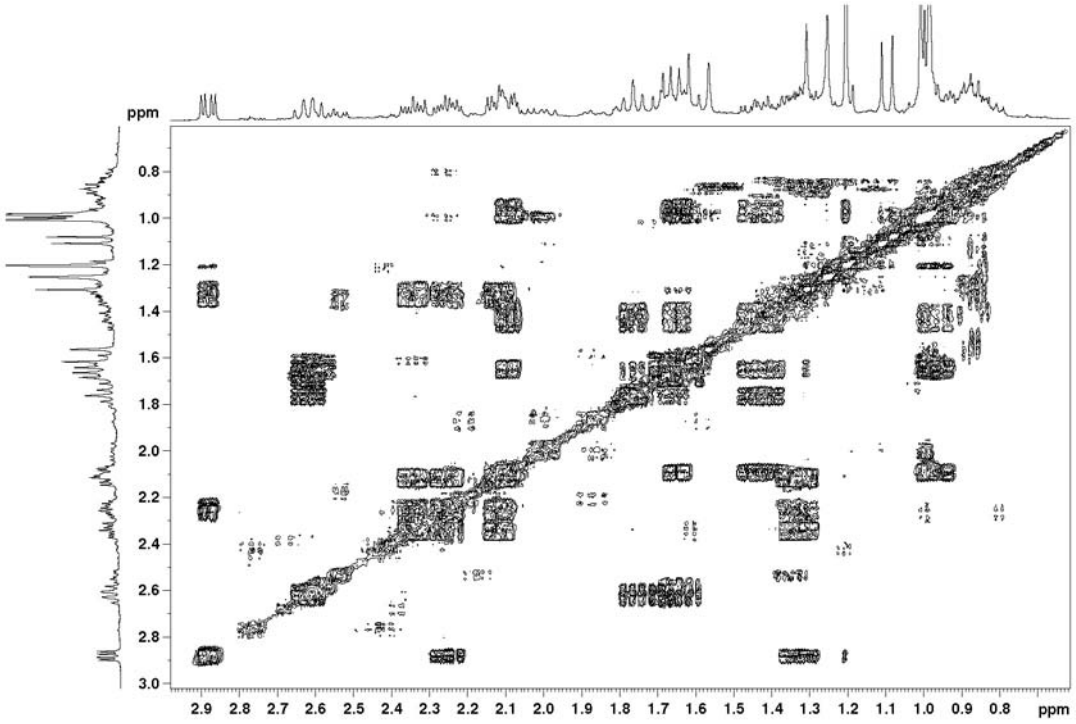

Figure S22. ^{1}H-^{1}H correlation map from COSY NMR experiment of caryophyllene oxide in $CDCl_3$ at 400 MHz.

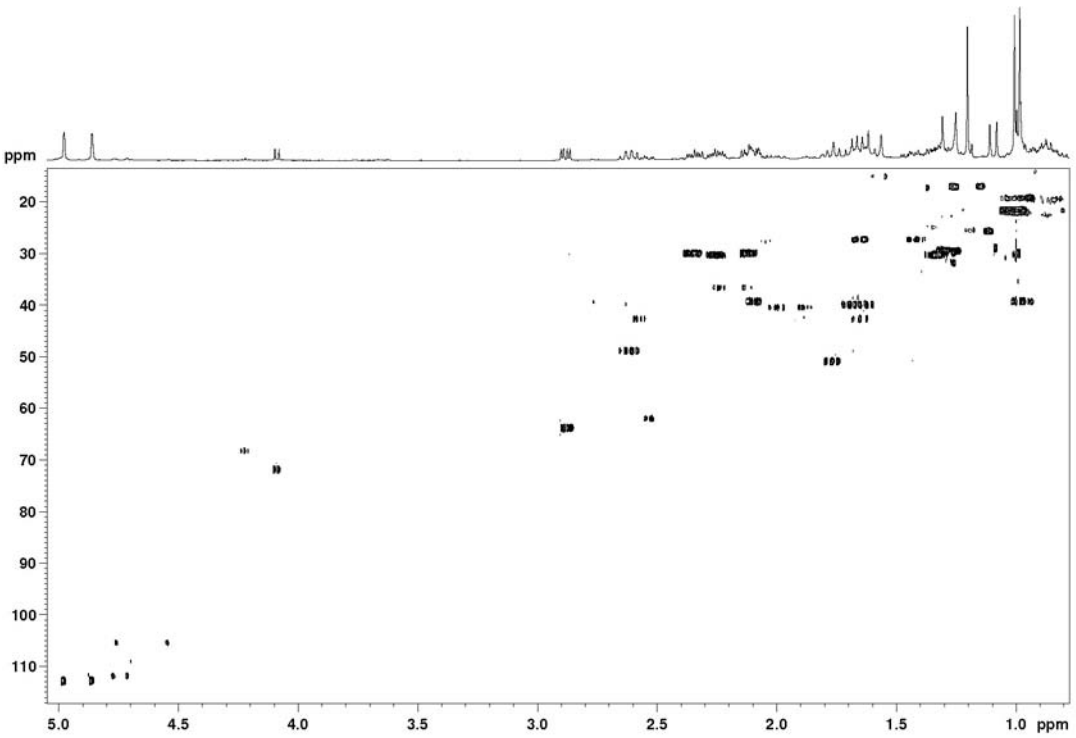

Figure S23. ^{1}H-^{13}C one-bond correlation map from the HSQC NMR experiment on caryophyllene oxide in $CDCl_3$ at 400 and 100 MHz.

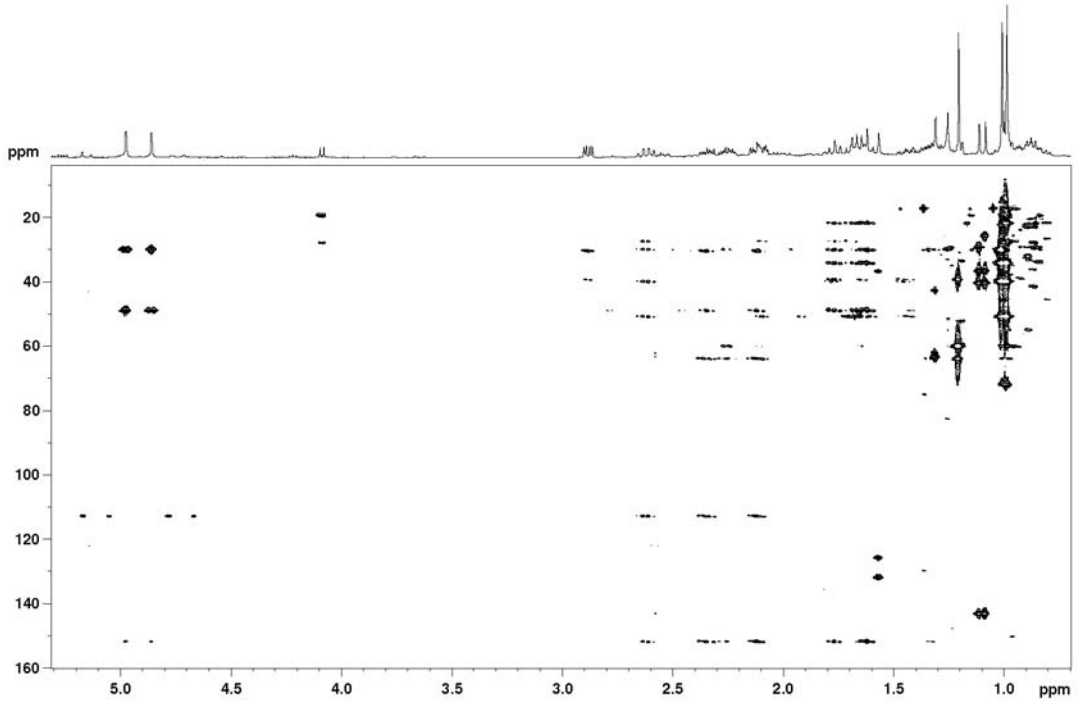

Figure S24. ^{1}H-^{13}C long-range correlation map from the HMBC NMR experiment on caryophyllene oxide in $CDCl_3$ at 400 and 100 MHz.

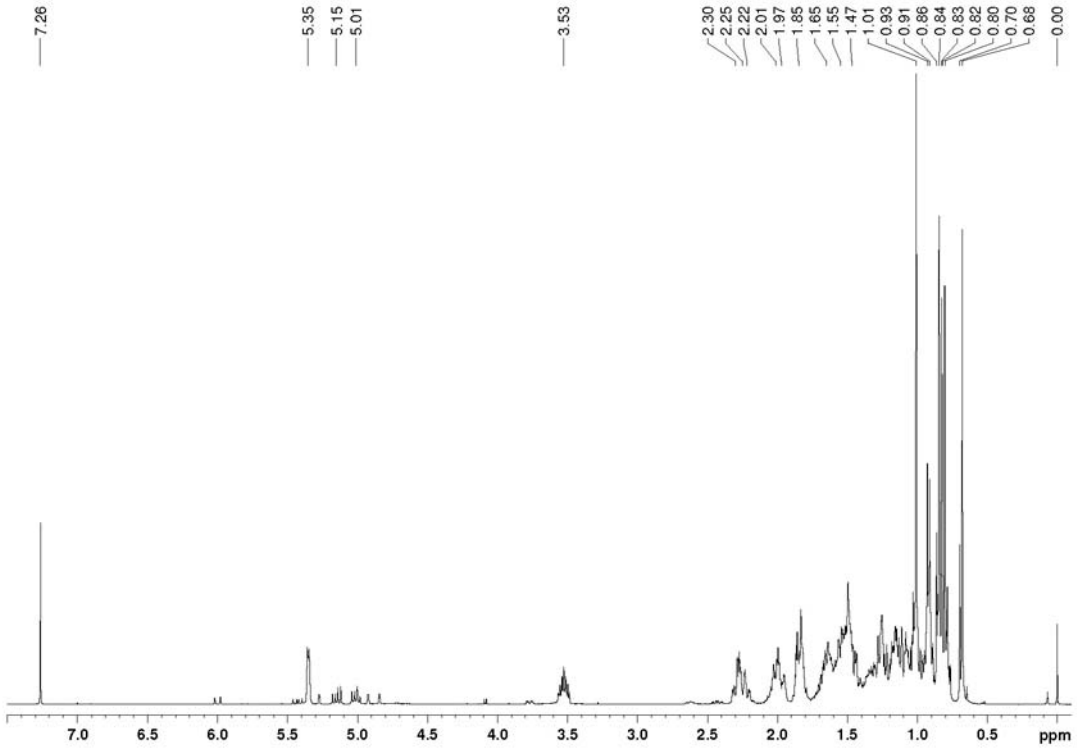

Figure S25. ^{1}H NMR spectrum of β-sitosterol and stigmasterol mixture in $CDCl_3$ at 400 MHz.

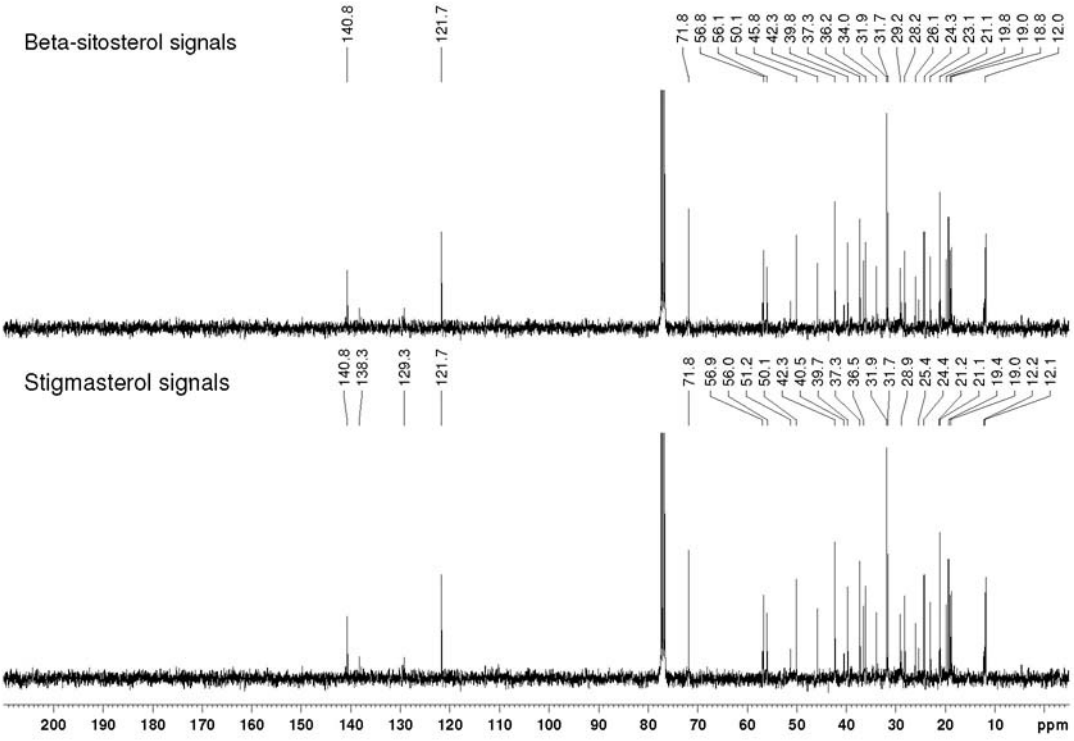

Figure S26. $^{13}C\{^{1}H\}$ NMR spectrum of the β-sitosterol and stigmasterol mixture in $CDCl_3$ at 100 MHz.

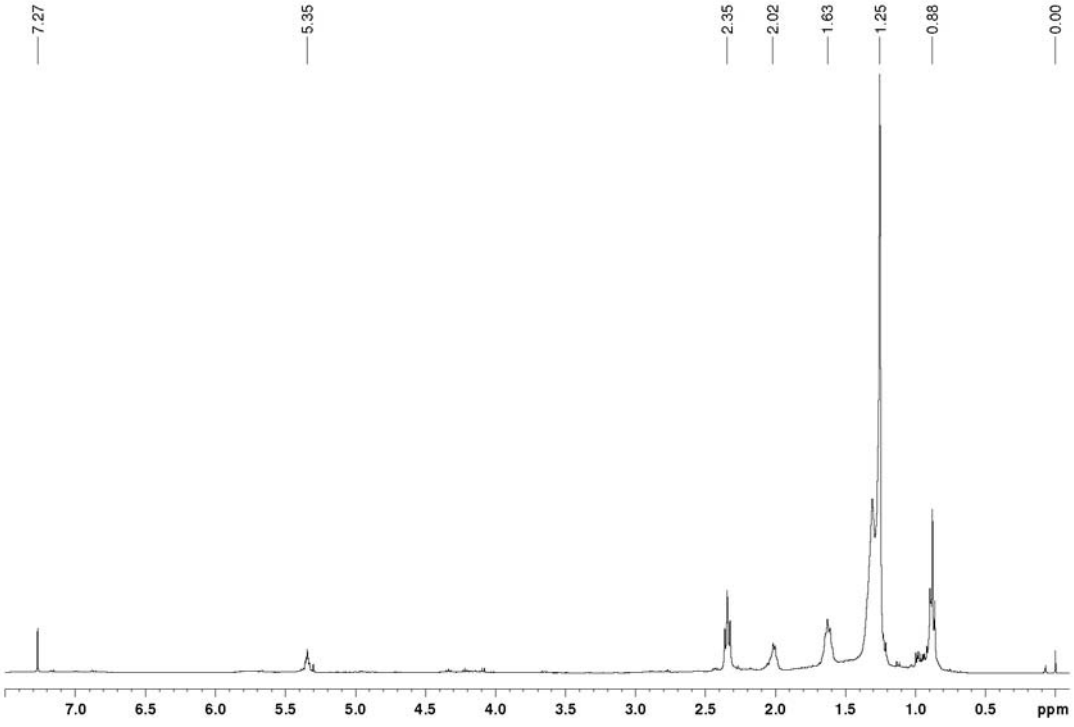

Figure S27. ^{1}H NMR spectrum of oleic acid in $CDCl_3$ at 400 MHz.

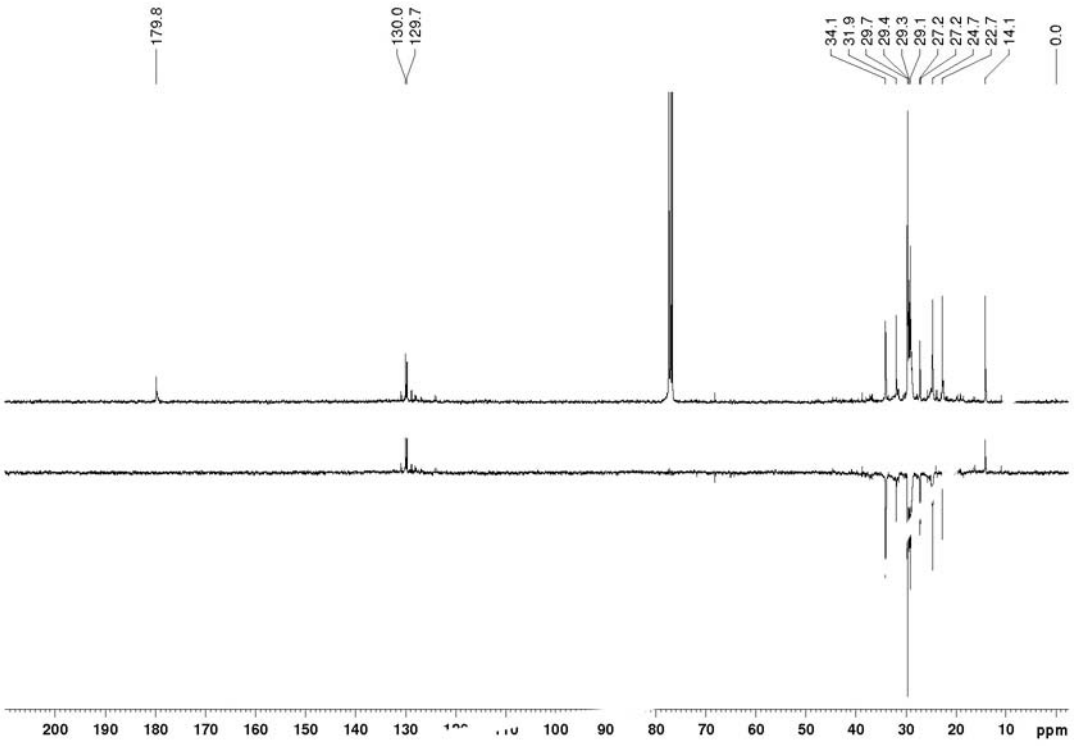

Figure S28. $^{13}C\{^{1}H\}$ and DEPT 135 NMR spectra of oleic acid in $CDCl_3$ at 100 MHz.