

New Flavone from the Leaves of *Neea theifera* (Nyctaginaceae)

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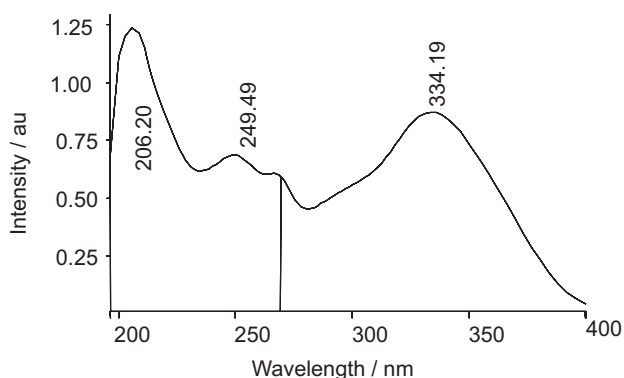


Figure S1. UV spectrum of (1) (MeOH).

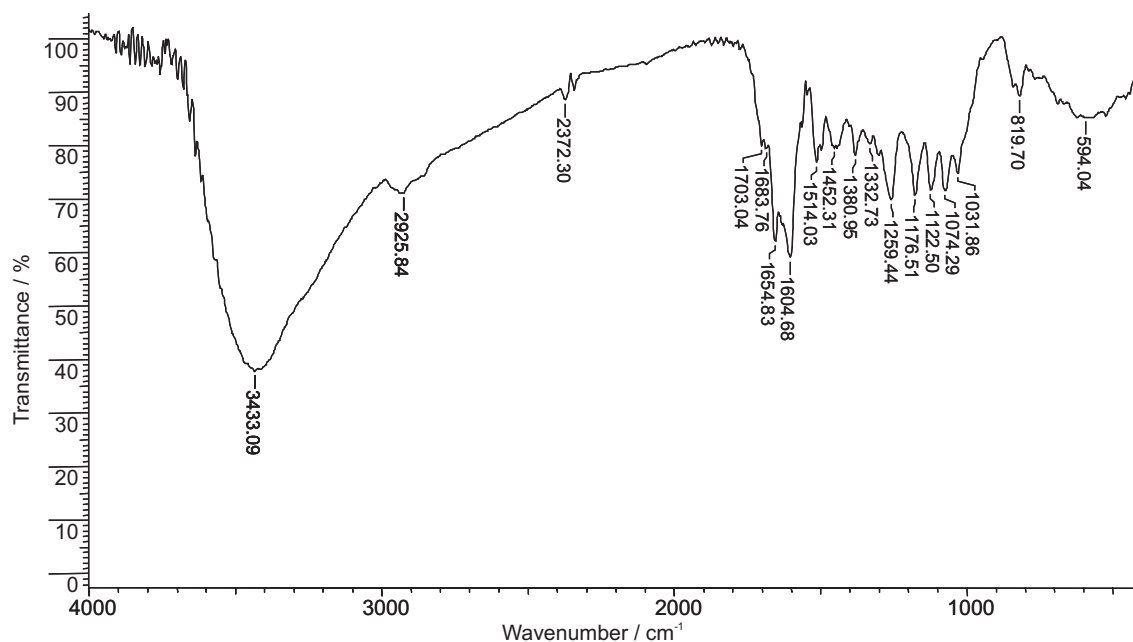
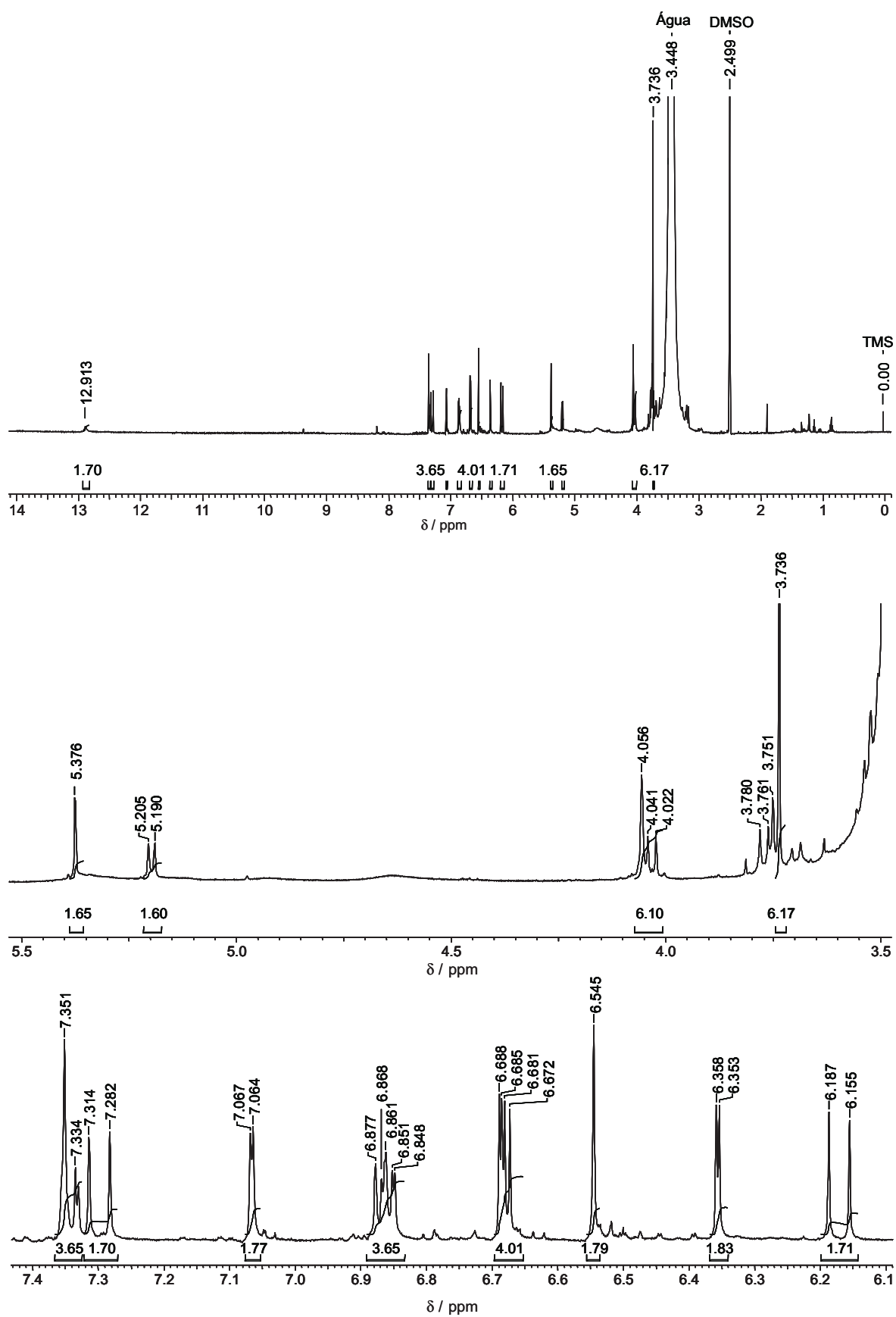


Figure S2. IR spectrum of (1) (KBr disk).

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Figure S3. ^1H NMR spectrum of (1) ($\text{DMSO}-d_6$, 11.7 T, TMS, ppm).

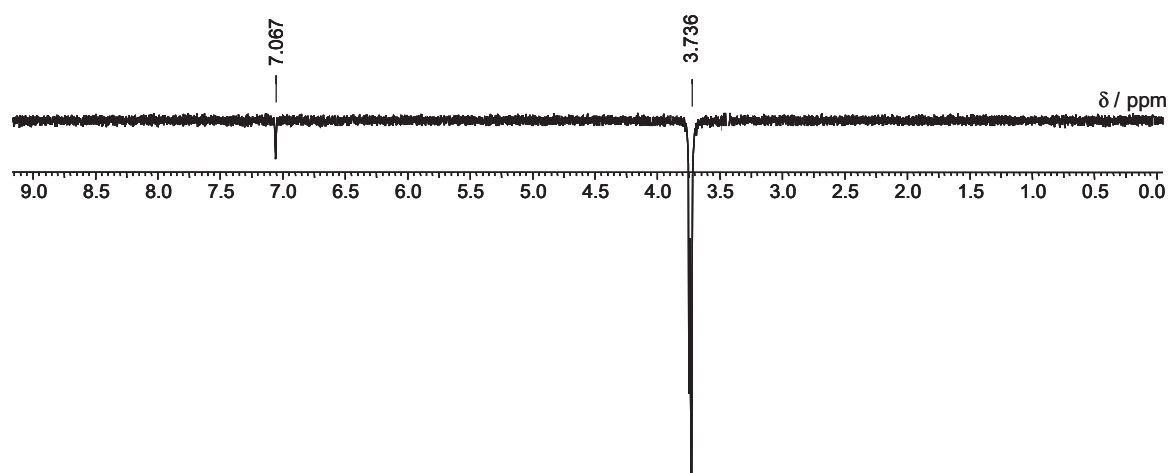


Figure S4. ¹H NMR 1D-NOESY of (1) (DMSO-*d*₆, 11.7 T, ppm).

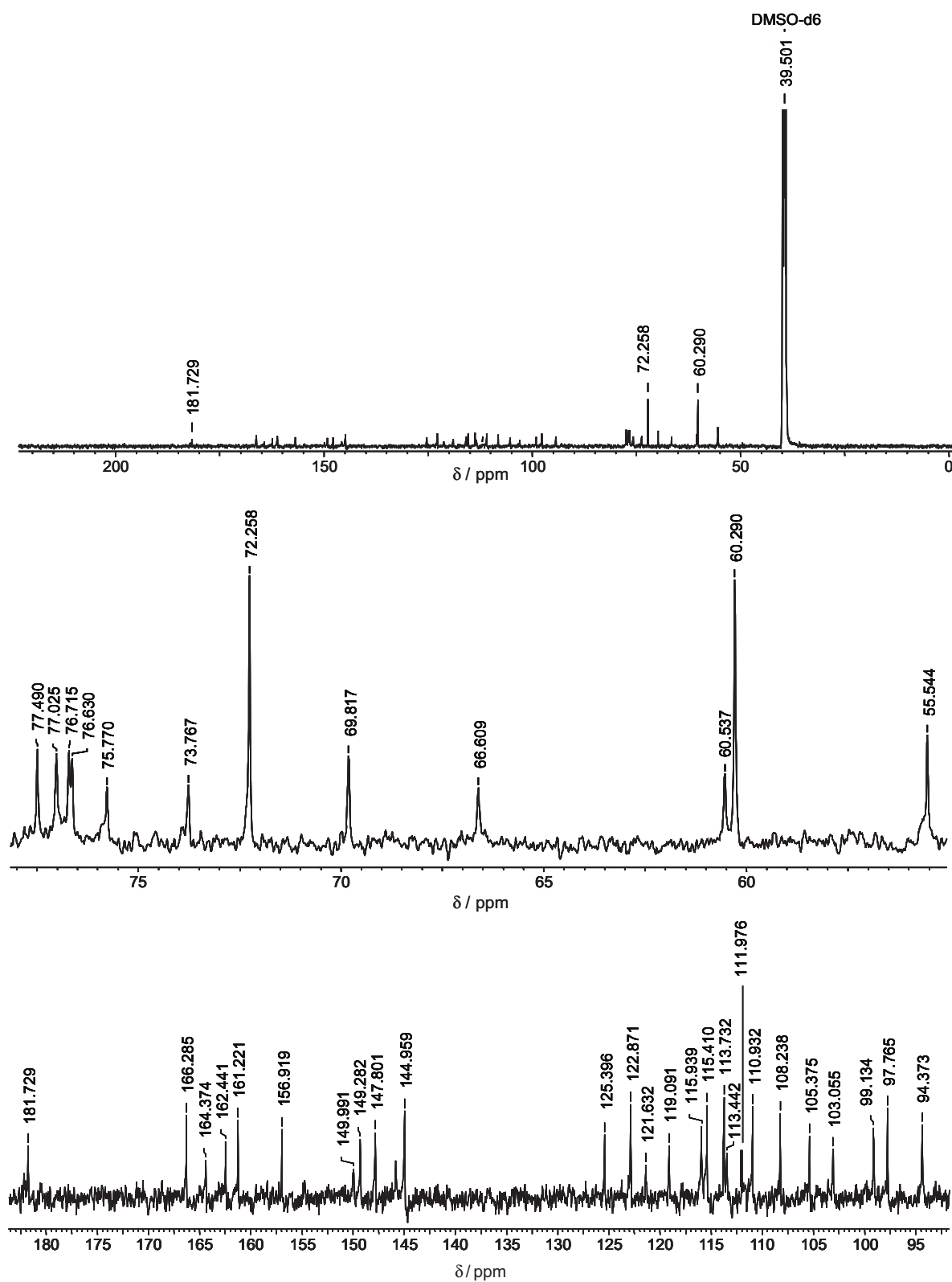


Figure S5. ^{13}C NMR spectrum of (I) ($\text{DMSO}-d_6$, 11.7 T, ppm).

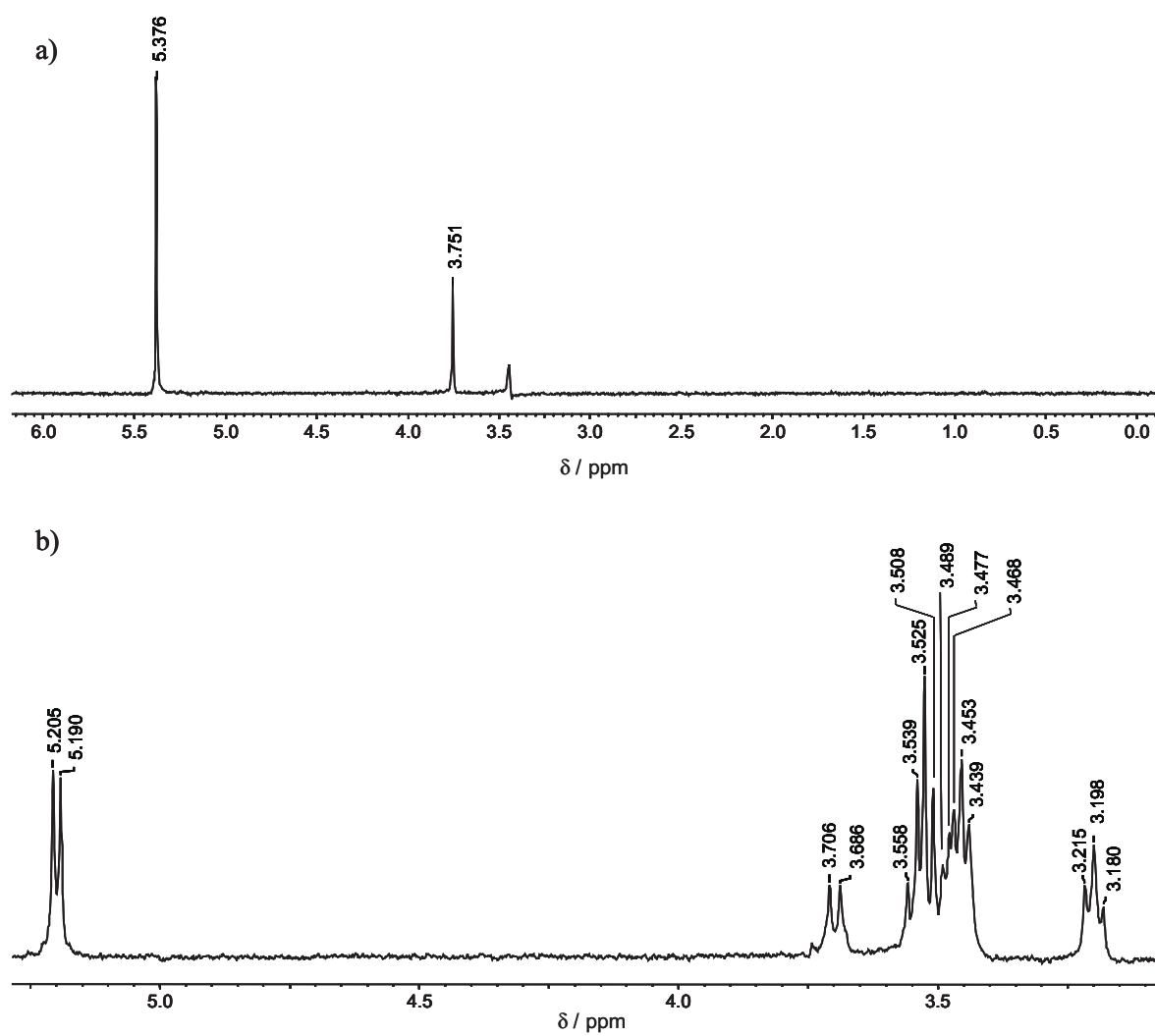


Figure S6. TOCSY spectrum of (I) (DMSO- d_6 , 11.7 T, ppm): a) Apiose; b) Glucose.

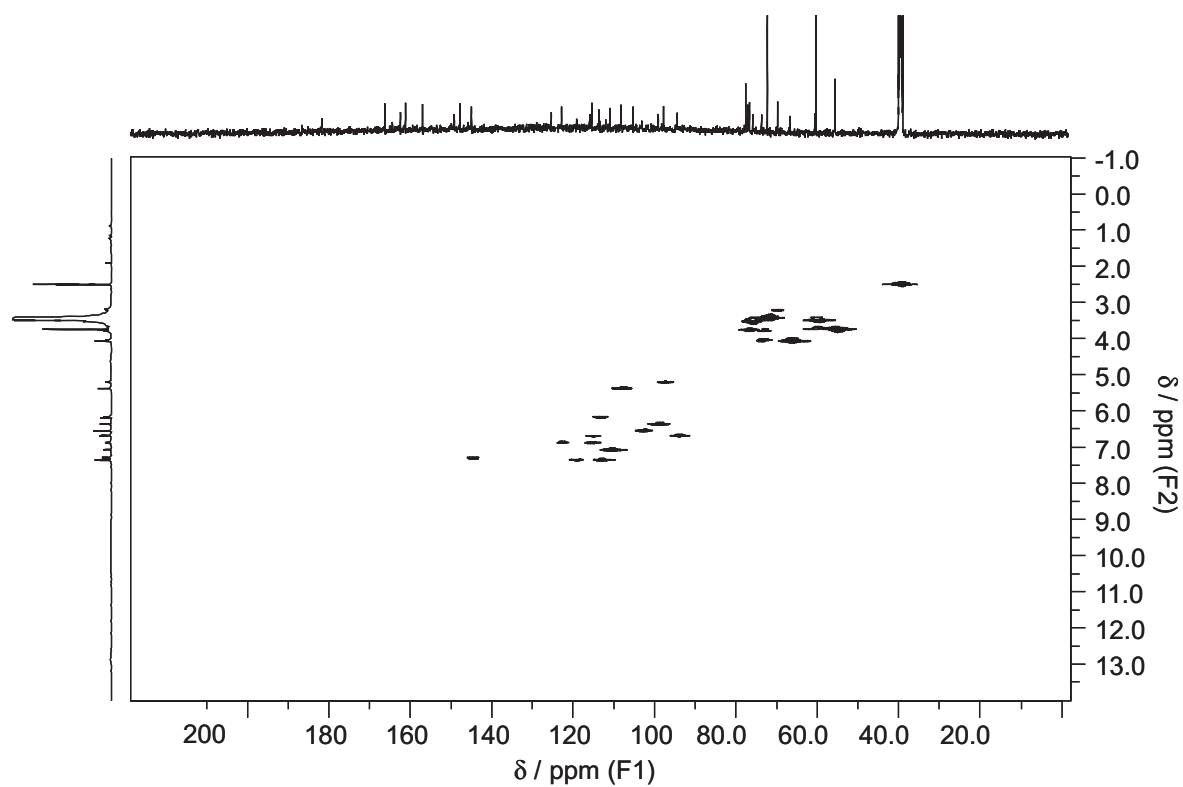


Figure S7. gHMBC spectrum of (1) (DMSO- d_6 , 11.7 T, TMS, ppm).

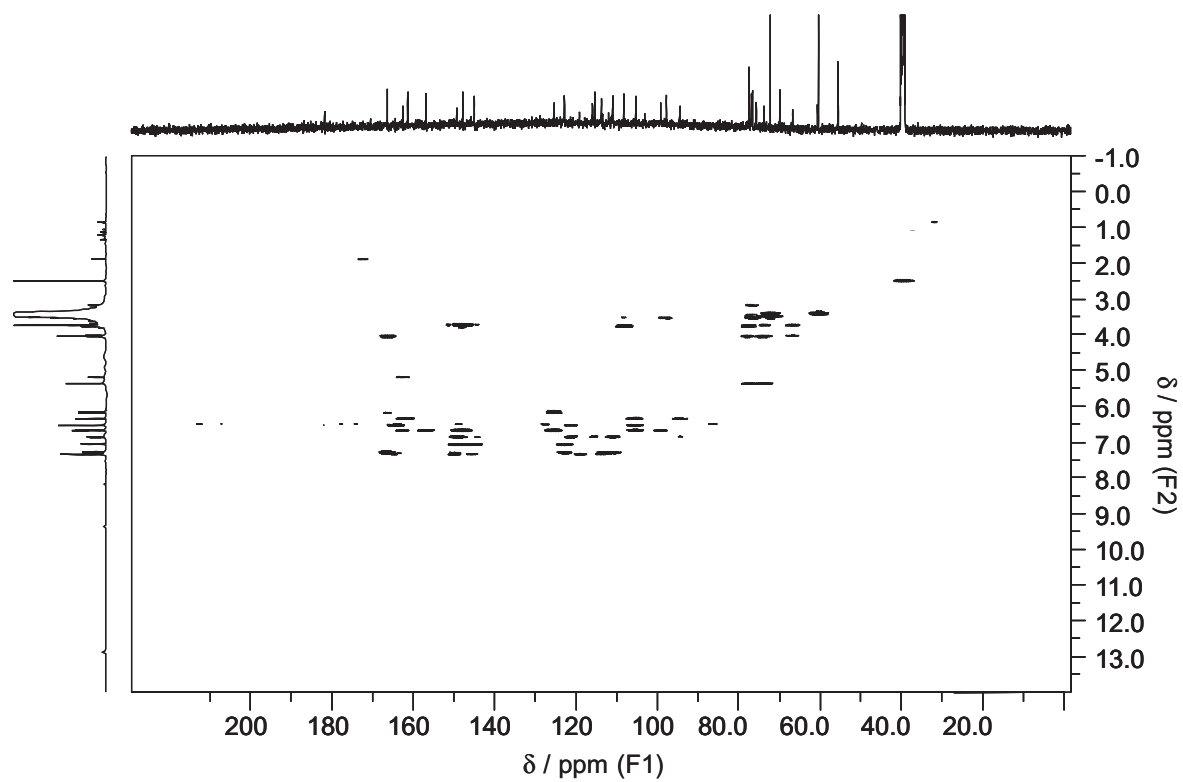


Figure S8. gHMBC spectrum of (1) (DMSO- d_6 , 11.7 T, TMS, ppm).

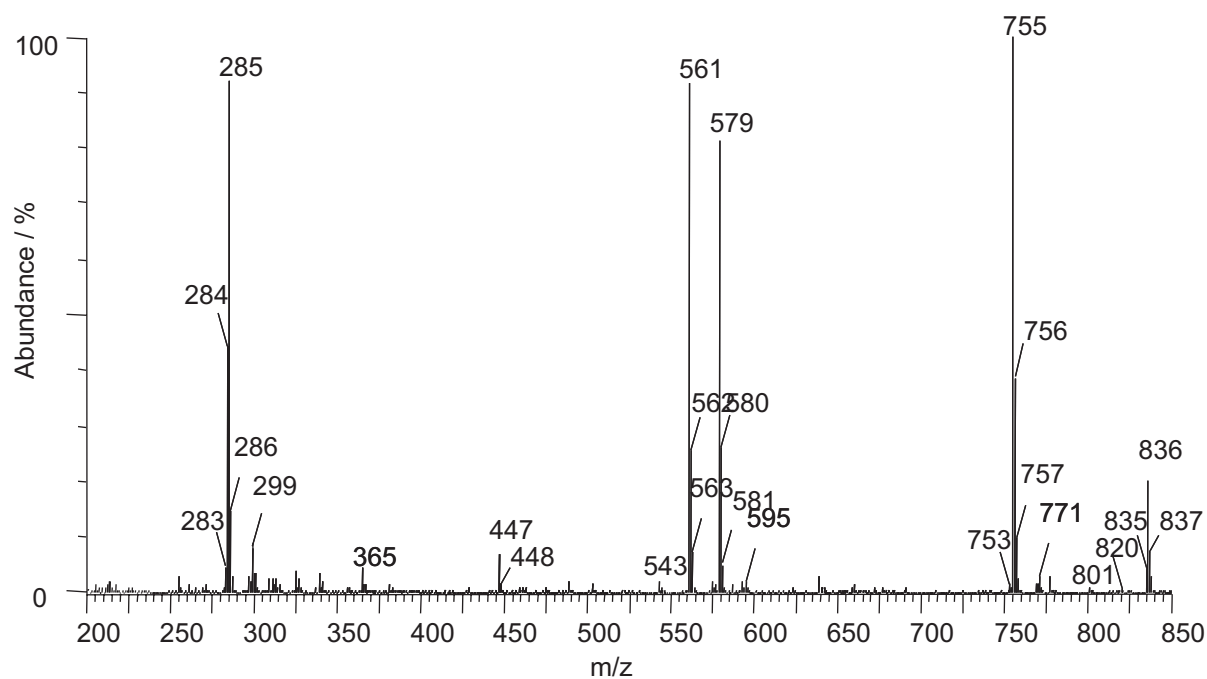


Figure S9. ESI-MS spectrum of (1) (negative mode, 70 V).

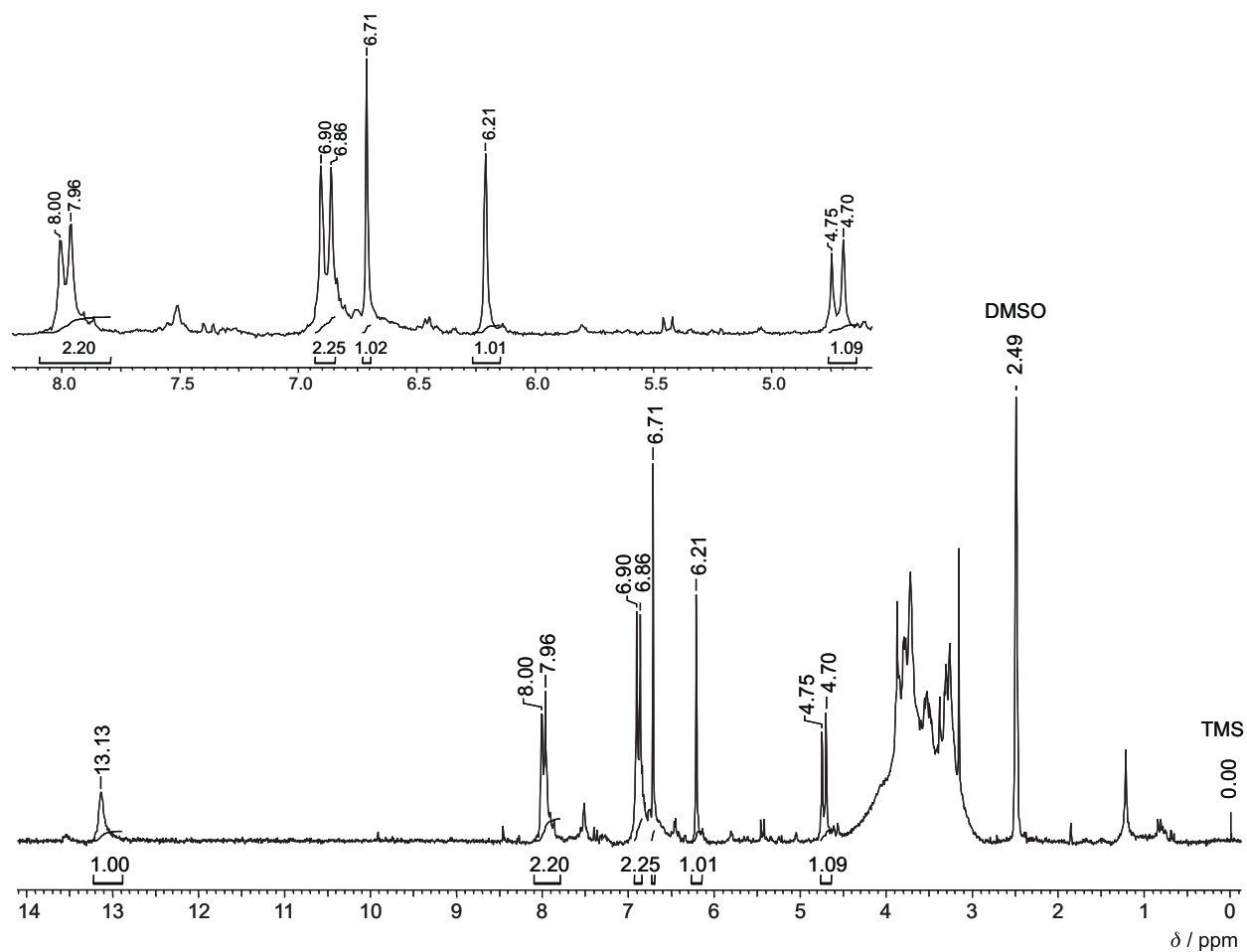


Figure S10. ^1H NMR spectrum of vitexin ($\text{DMSO}-d_6$, 4.7 T, TMS, ppm).

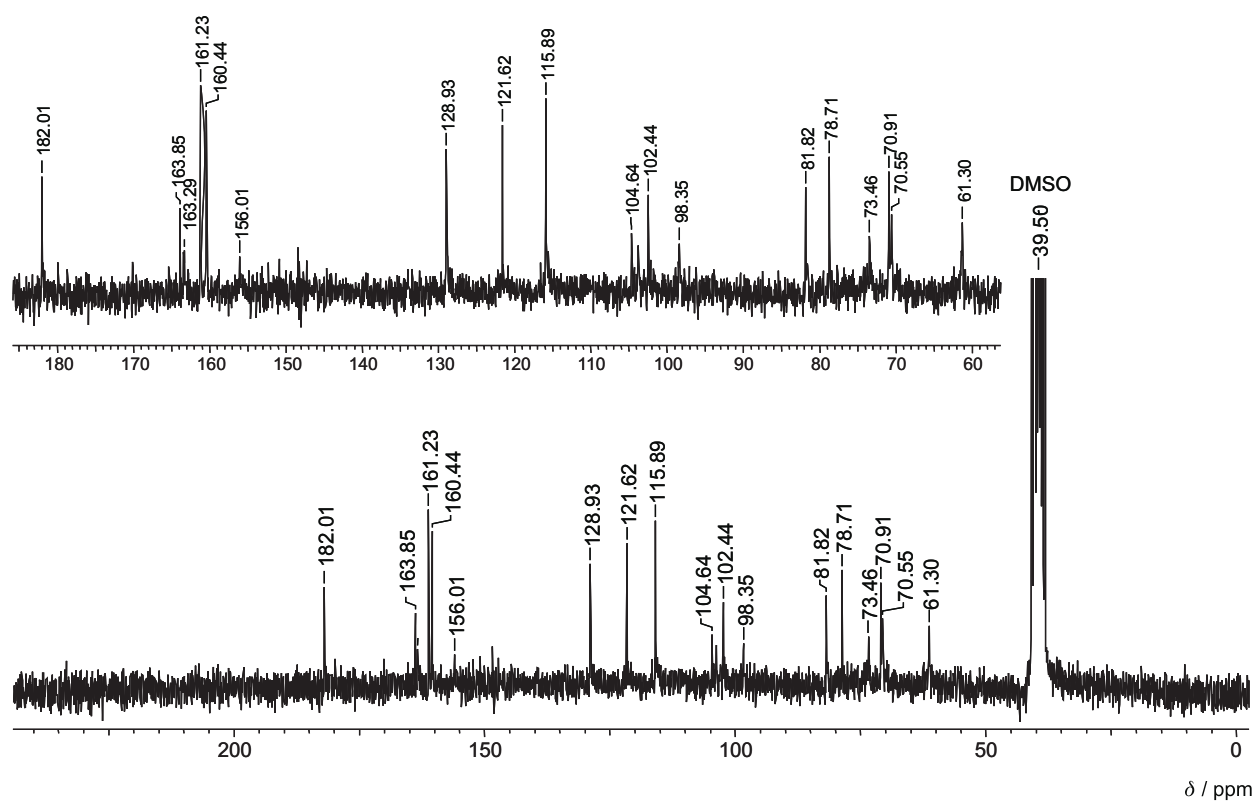


Figure S11. ^{13}C NMR spectrum of vitexin (DMSO-d_6 , 4.7 T, ppm).

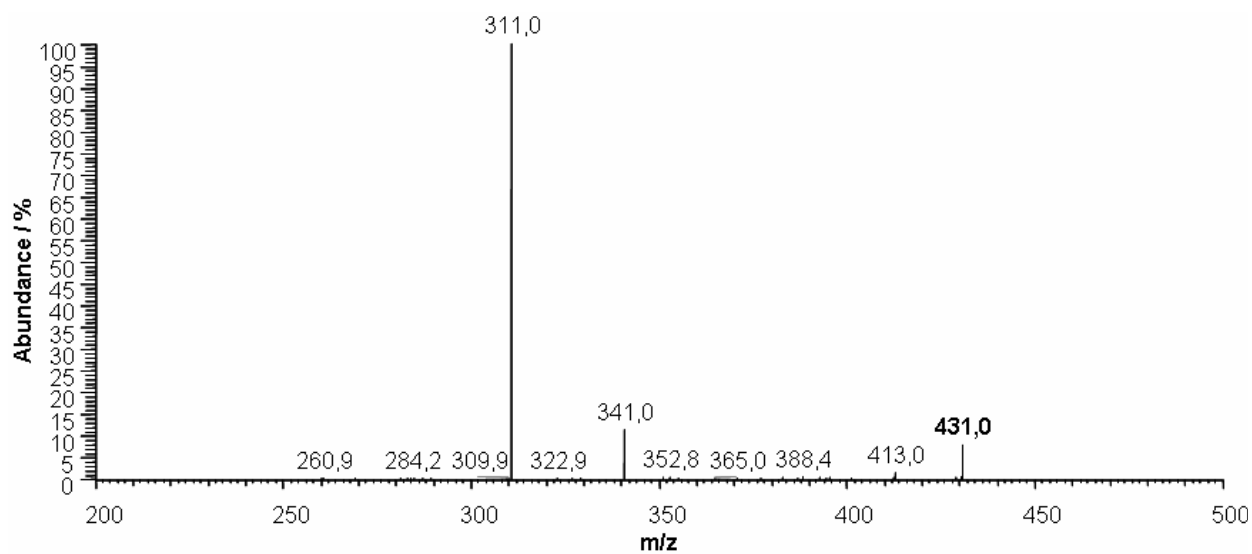


Figure S12. ESI-MS spectrum of vitexin (negative mode, 70 V).

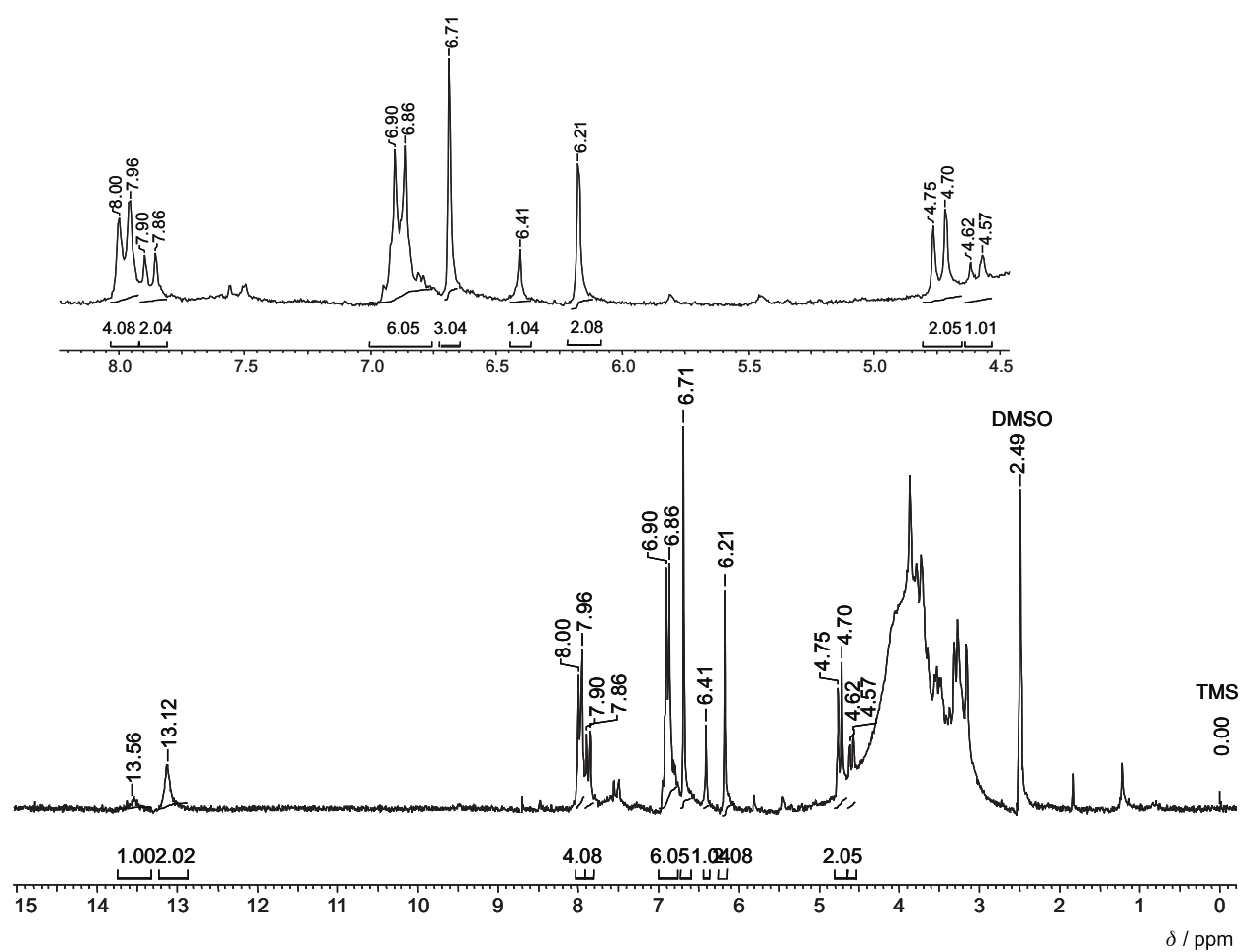


Figure S13. ^1H NMR spectrum of the mixture of isovitexin and vitexin ($\text{DMSO}-d_6$, 4.7 T, TMS, ppm).

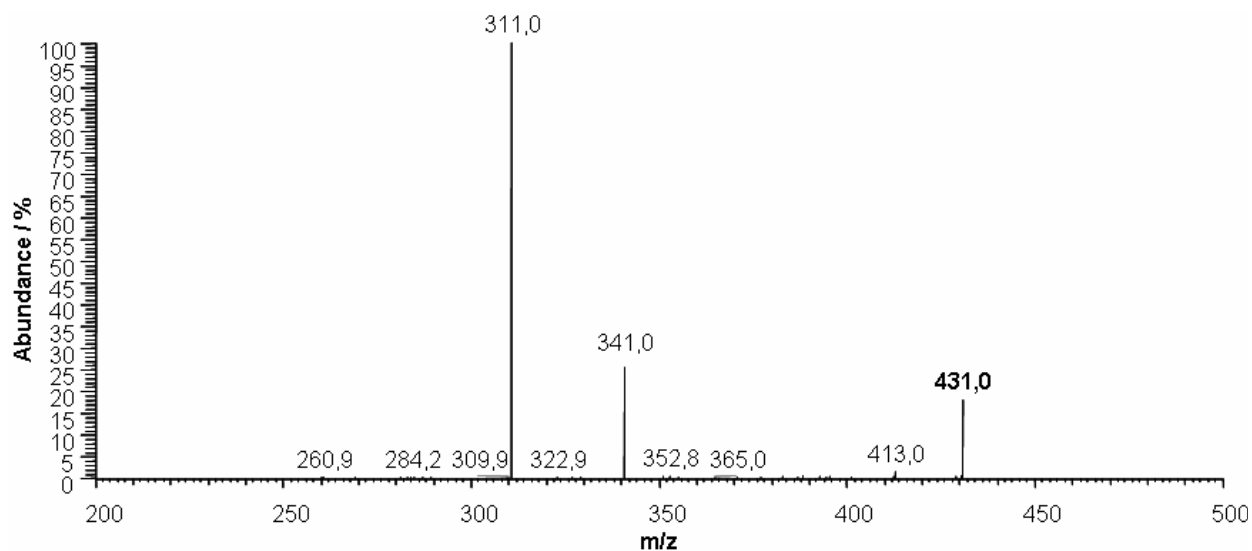


Figure S14. ESI-MS spectrum of the mixture of isovitexin and vitexin (negative mode, 70 V).

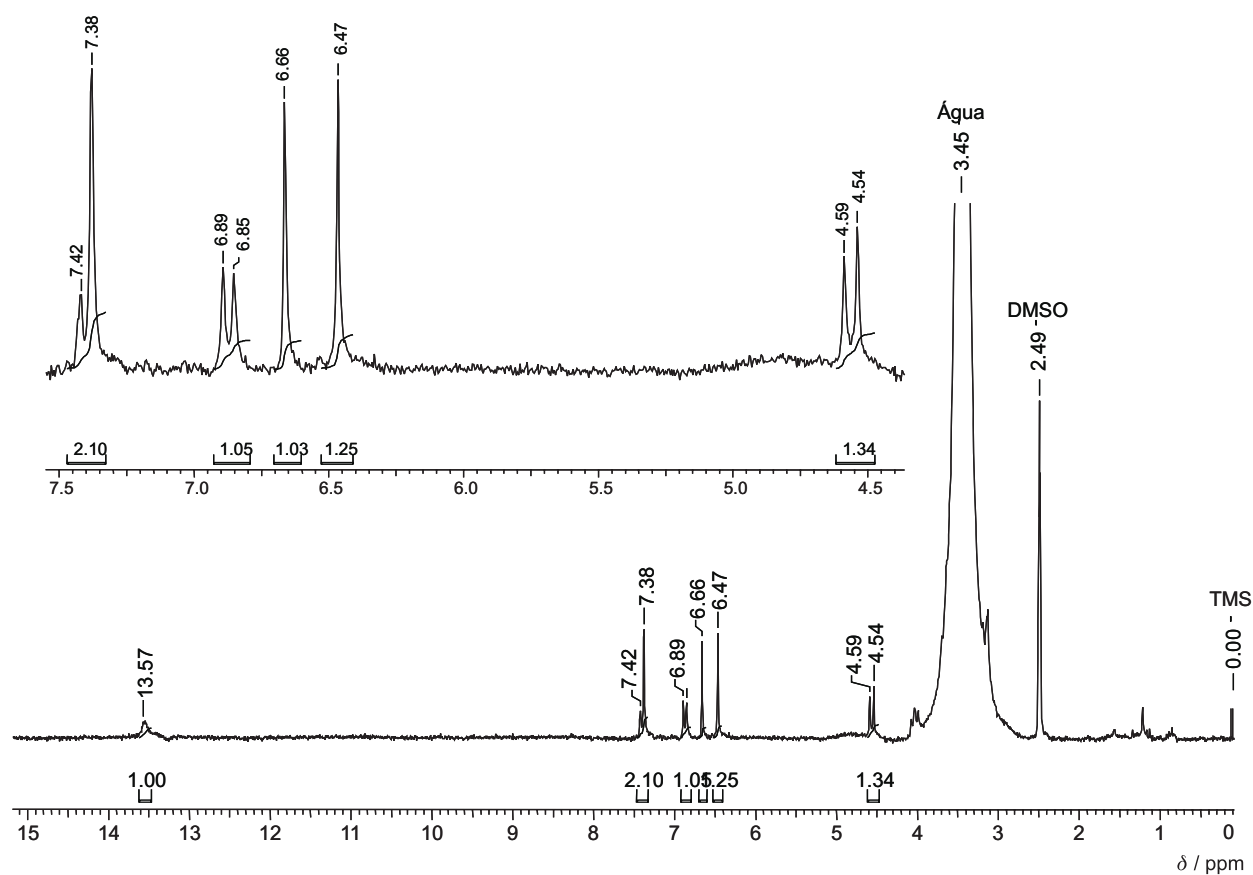


Figure S15. ¹H NMR spectrum of isoorientin (DMSO-*d*₆, 4.7 T, TMS, ppm).

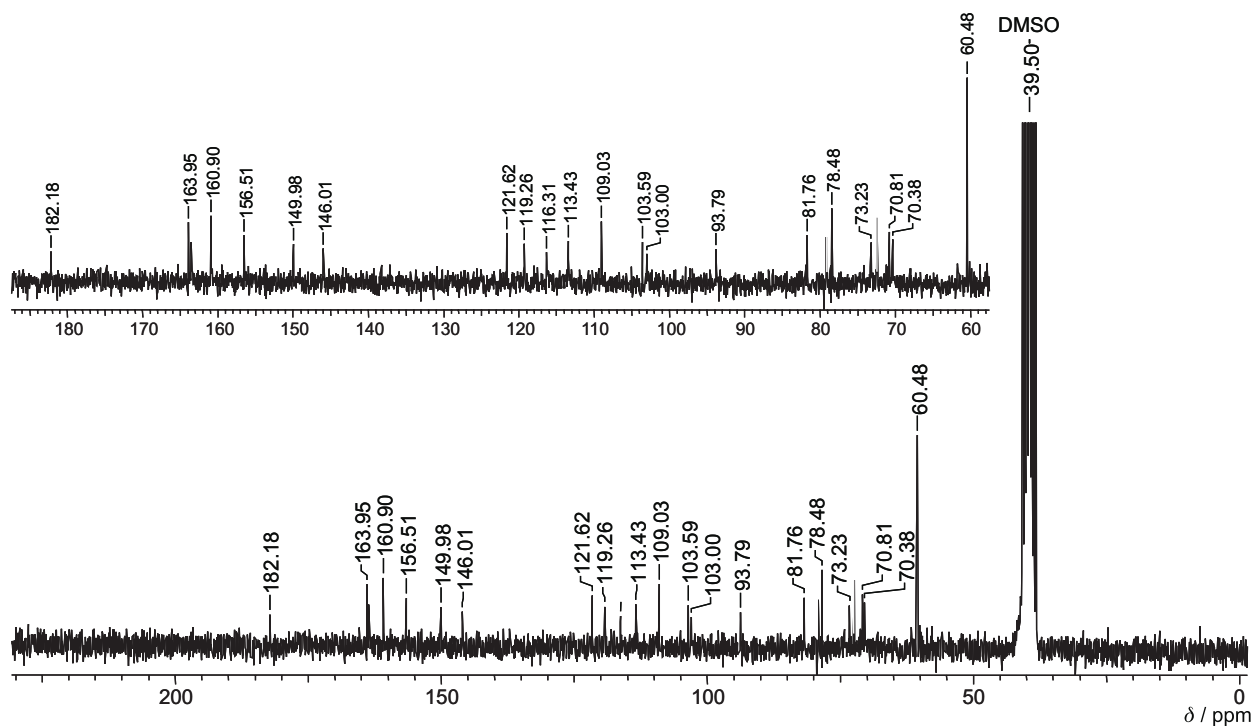


Figure S16. ¹³C NMR spectrum of isoorientin (DMSO-*d*₆, 4.7 T, ppm).

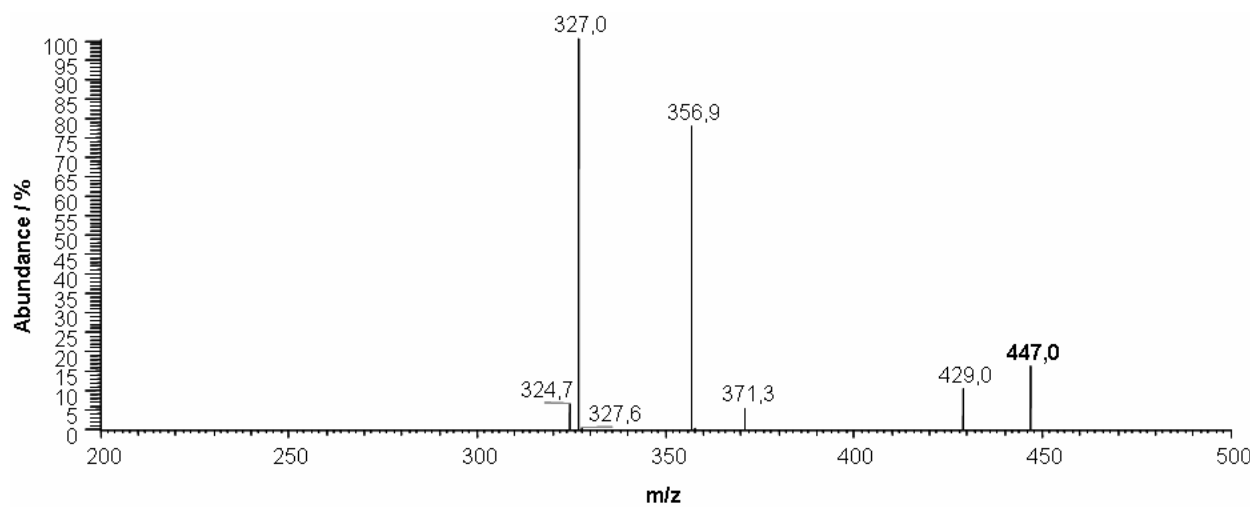


Figure S17. ESI-MS spectrum of isoorientin (negative mode, 70 V).

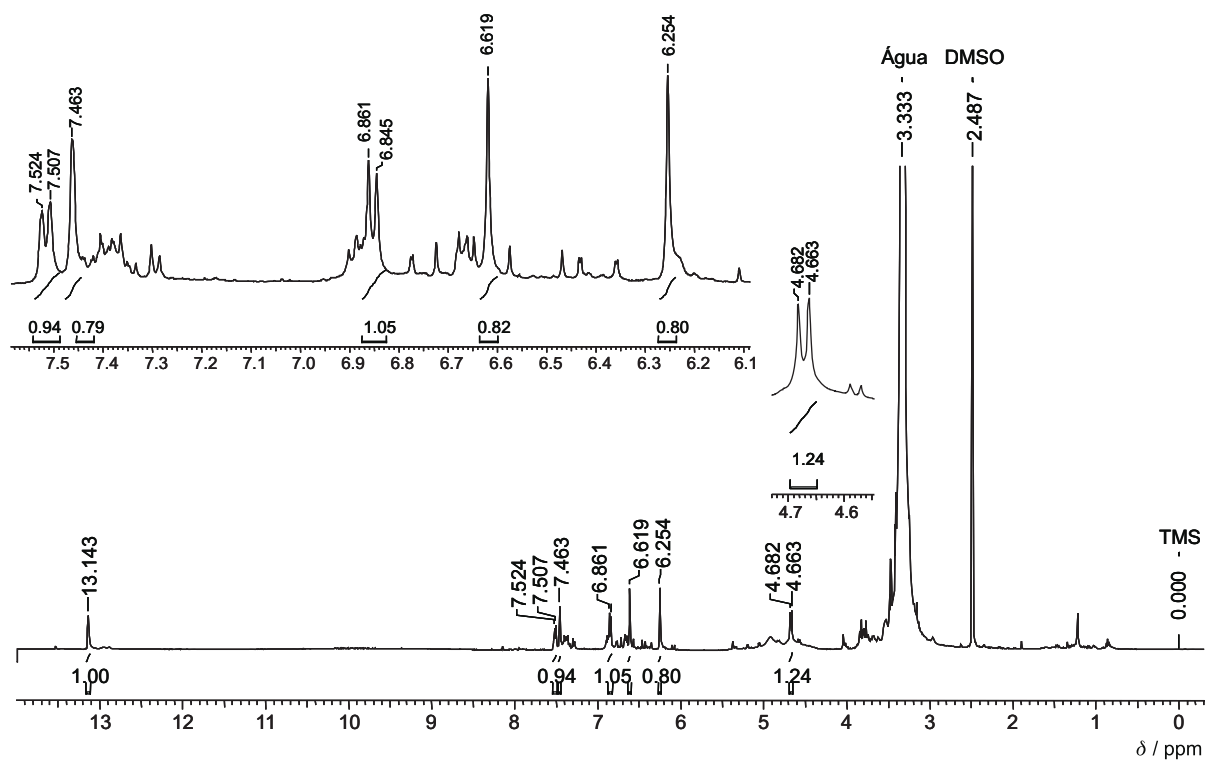


Figure S18. ^1H NMR spectrum of orientin ($\text{DMSO}-d_6$, 4.7 T, TMS, ppm).

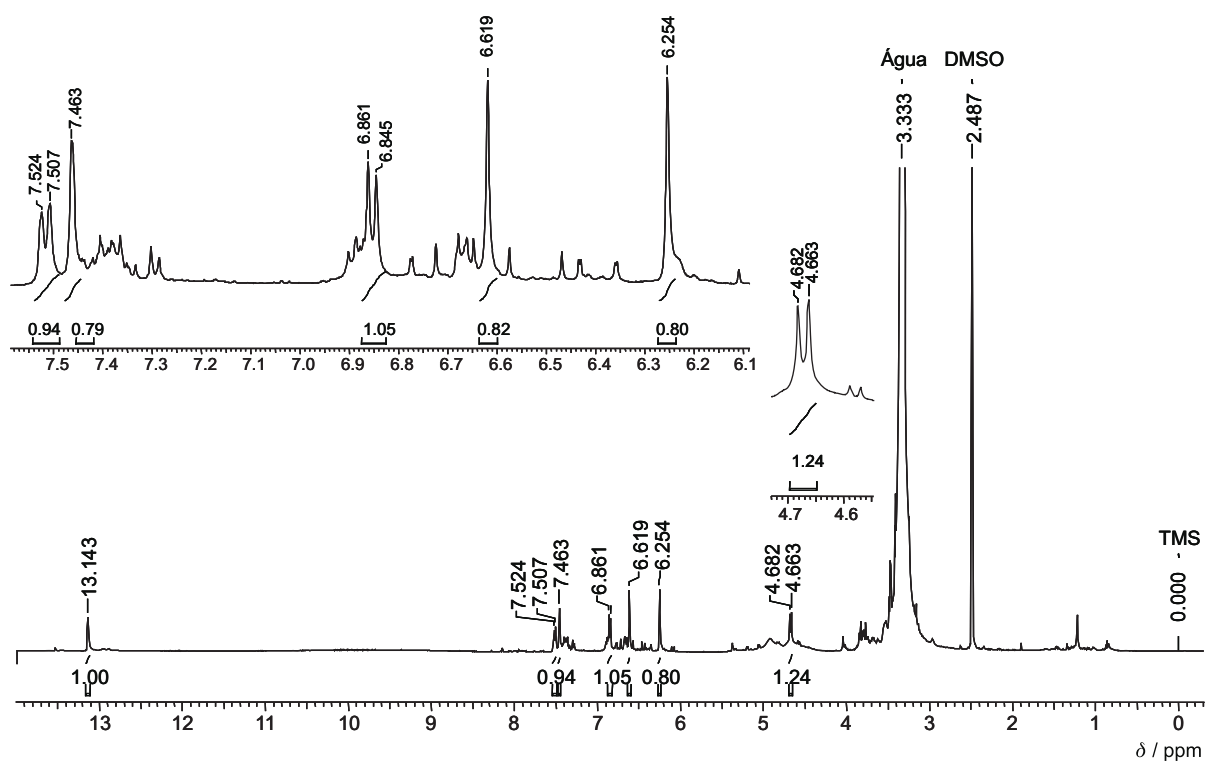


Figure S19. ¹³C NMR spectrum of orientin (DMSO-*d*₆, 4.7 T, ppm).

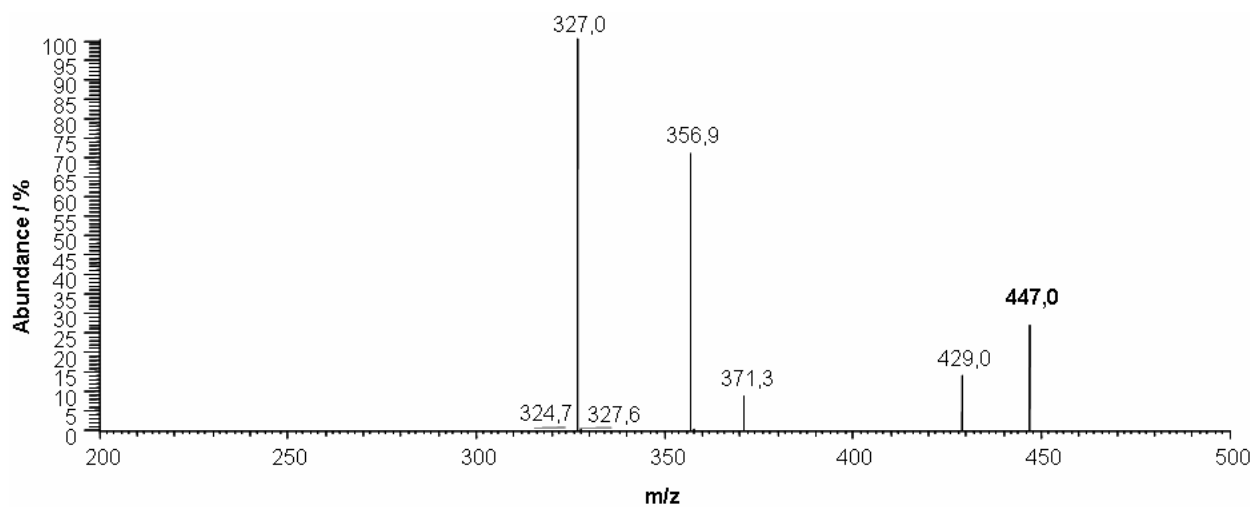


Figure S20. ESI-MS spectrum of orientin (negative mode, 70 V).

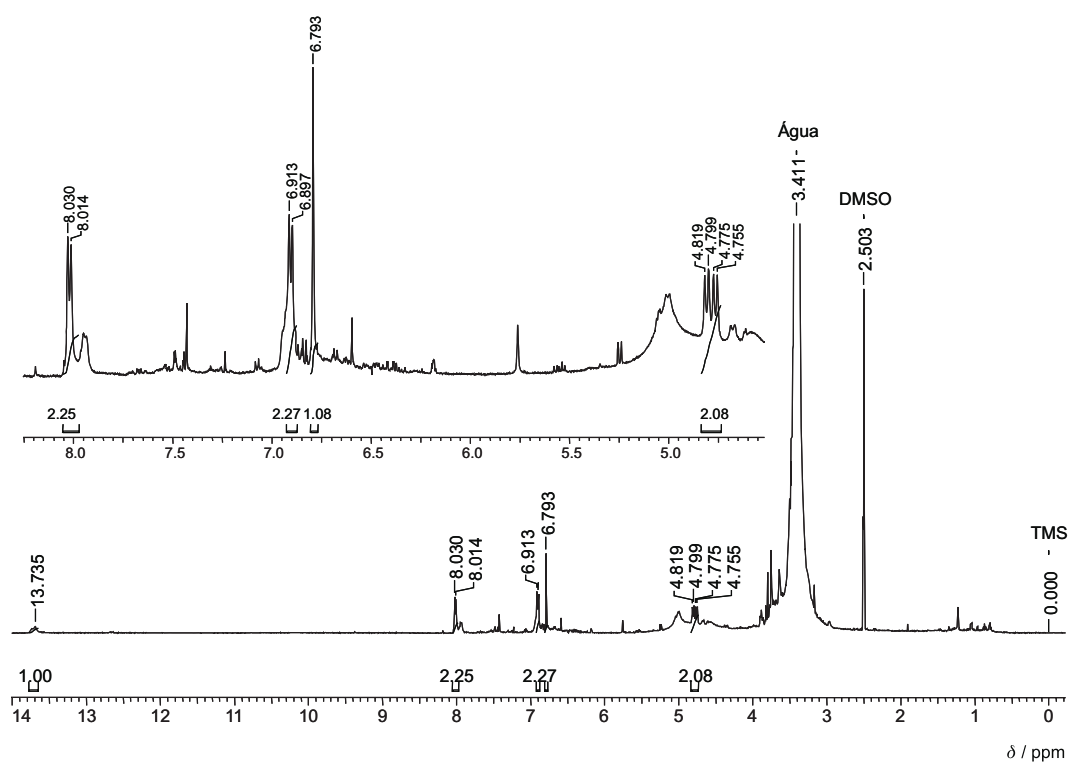


Figure S21. ^1H NMR spectrum of vicenin-2 ($\text{DMSO}-d_6$, 11.7 T, TMS, ppm).

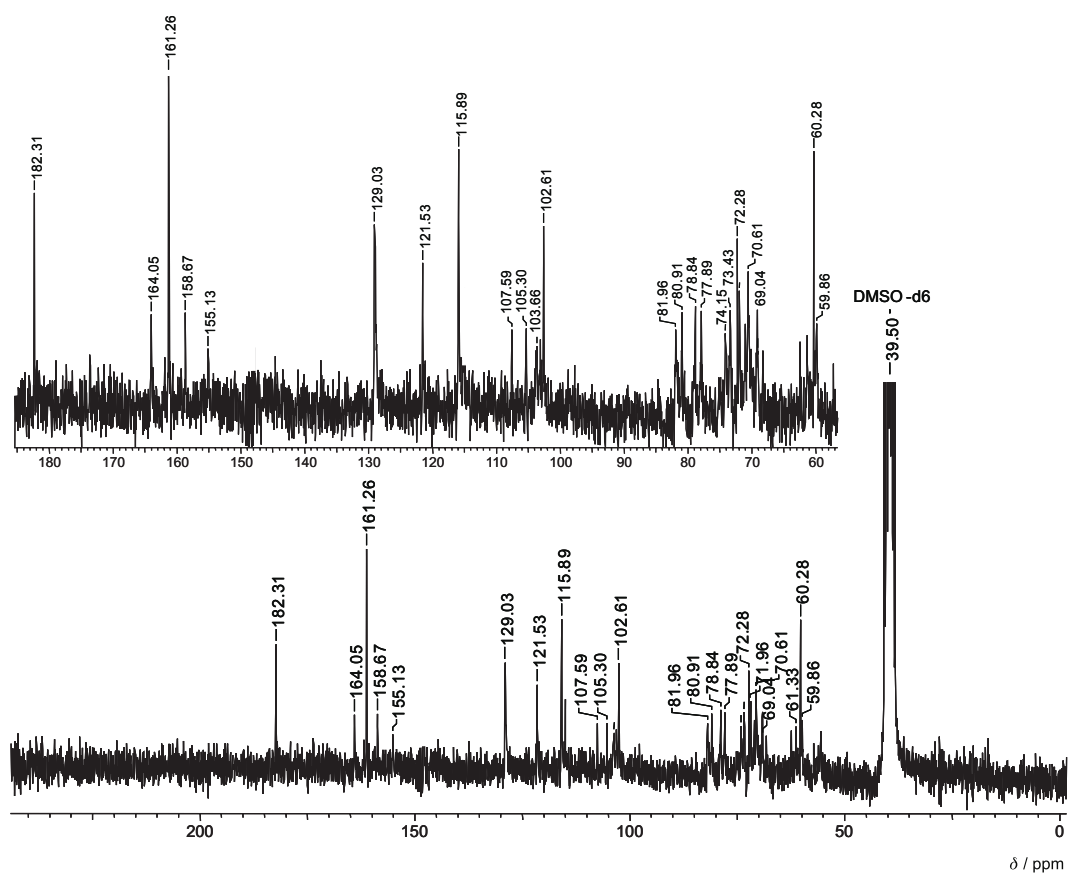


Figure S22. ^{13}C NMR spectrum of vicenin-2 ($\text{DMSO}-d_6$, 11.7 T, TMS, ppm).

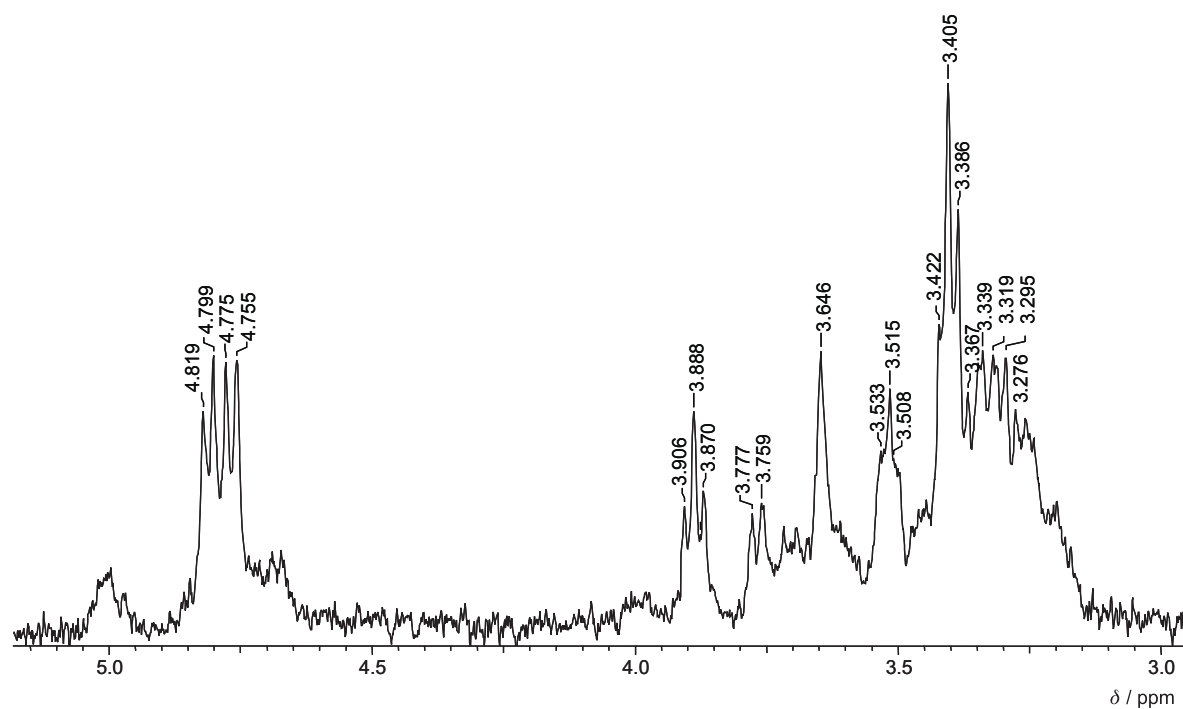


Figure S23. TOCSY spectrum of vicenin-2 (DMSO- d_6 , 11.7 T, ppm, irradiation of δ 4.809).

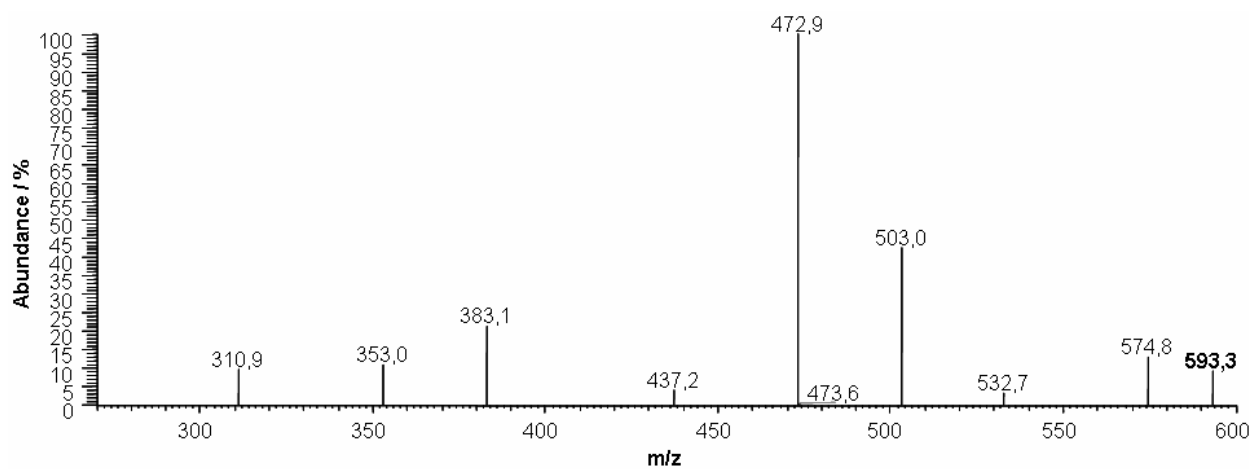


Figure S24. ESI-MS spectrum of vicenin-2 (negative mode, 70 V).

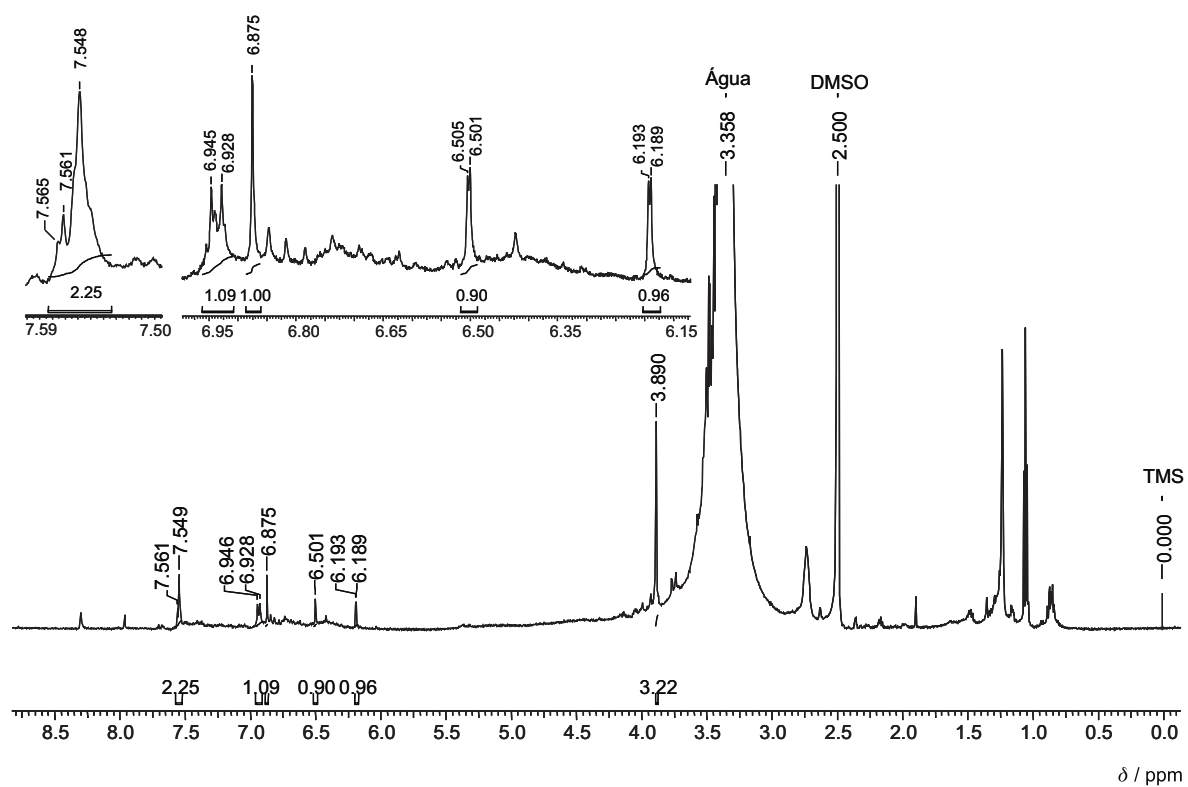


Figure S25. ¹H NMR spectrum of chrysoeriol (DMSO-*d*₆, 11.7 T, ppm).

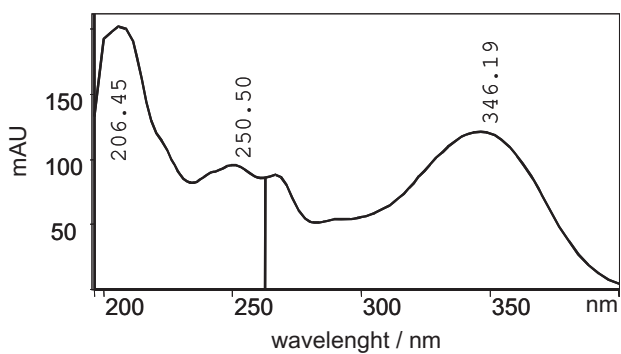


Figure S26. UV spectrum of chrysoeriol (MeOH).

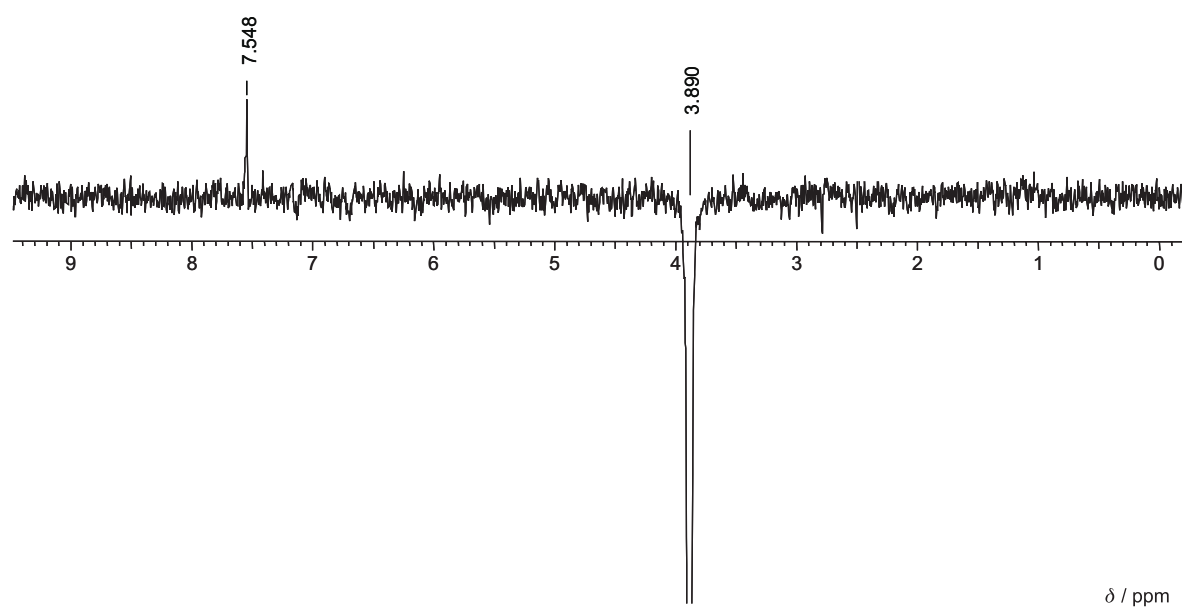


Figure S27. ¹H NMR 1D-NOESY of chrysoeriol (DMSO-*d*₆, 11.7 T, ppm, irradiation of δ 3.890).

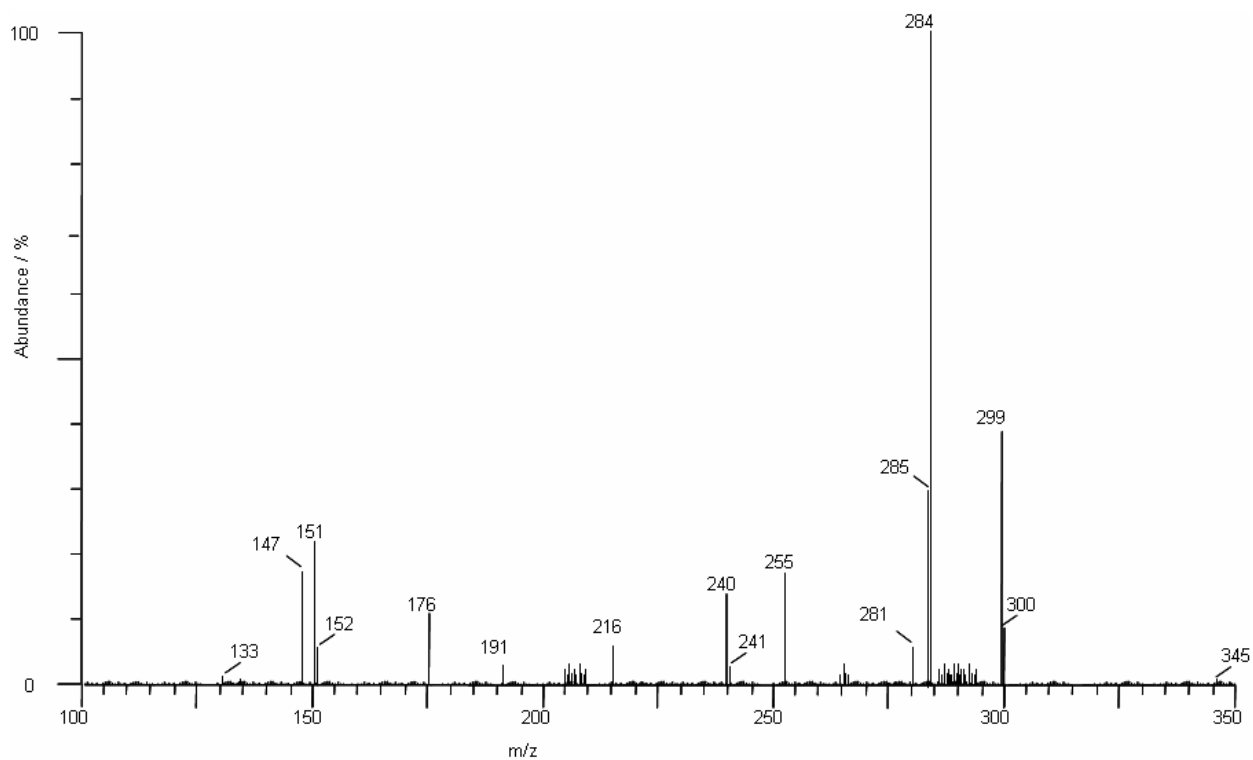


Figure S28. ESI-MS spectrum of chrysoeriol (negative mode, 70 V).

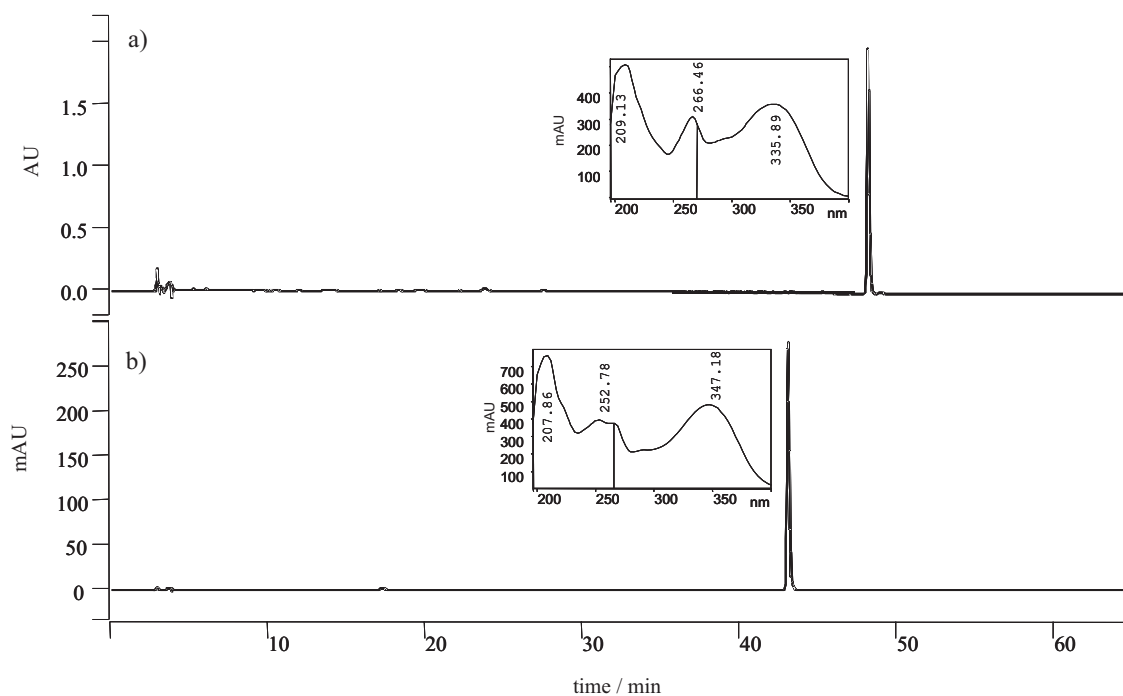


Figure S29. Chromatograms of the co-injections of a) apigenin isolated + apigenin Sigma® (1:1 m/m) and b) luteolin isolated + luteolin Sigma® (1:1 m/m) (C18, 250 × 4.60 mm i. d. × 5 µm). A binary gradient elution system with solvent A (0.05% TFA in H₂O) and solvent B (0.05% TFA in CH₃CN) was used, with linear gradient starting from 68:32 (A:B) at 20 min and then changed to 75:35 (A:B) for 40 min. The flow-rate was 1.0 mL min⁻¹. λ 254 nm.