

Supporting Information

Direct electrosynthesis of a series of novel caffeic acid analogues through a clean and serendipitous domino oxidation/thia-Michael reaction

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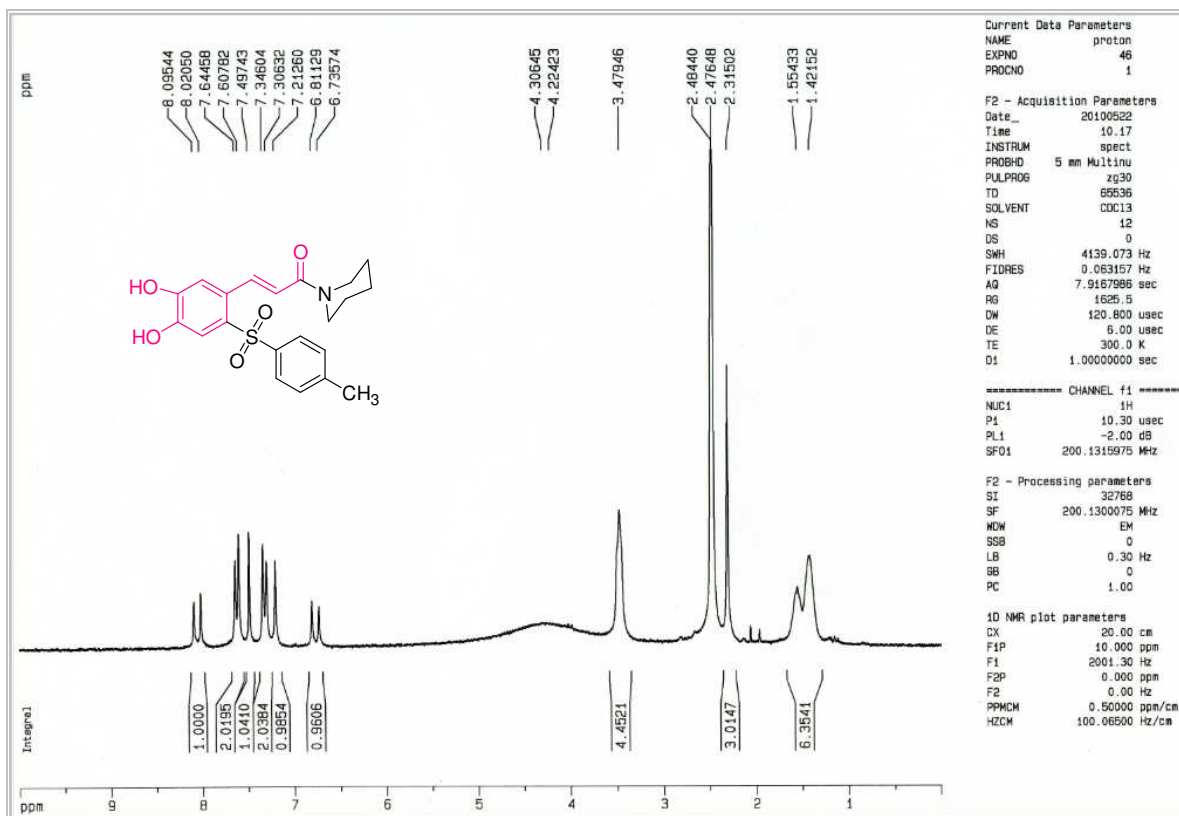


Figure S-1. ¹H NMR spectrum of compound 5a in DMSO-d₆

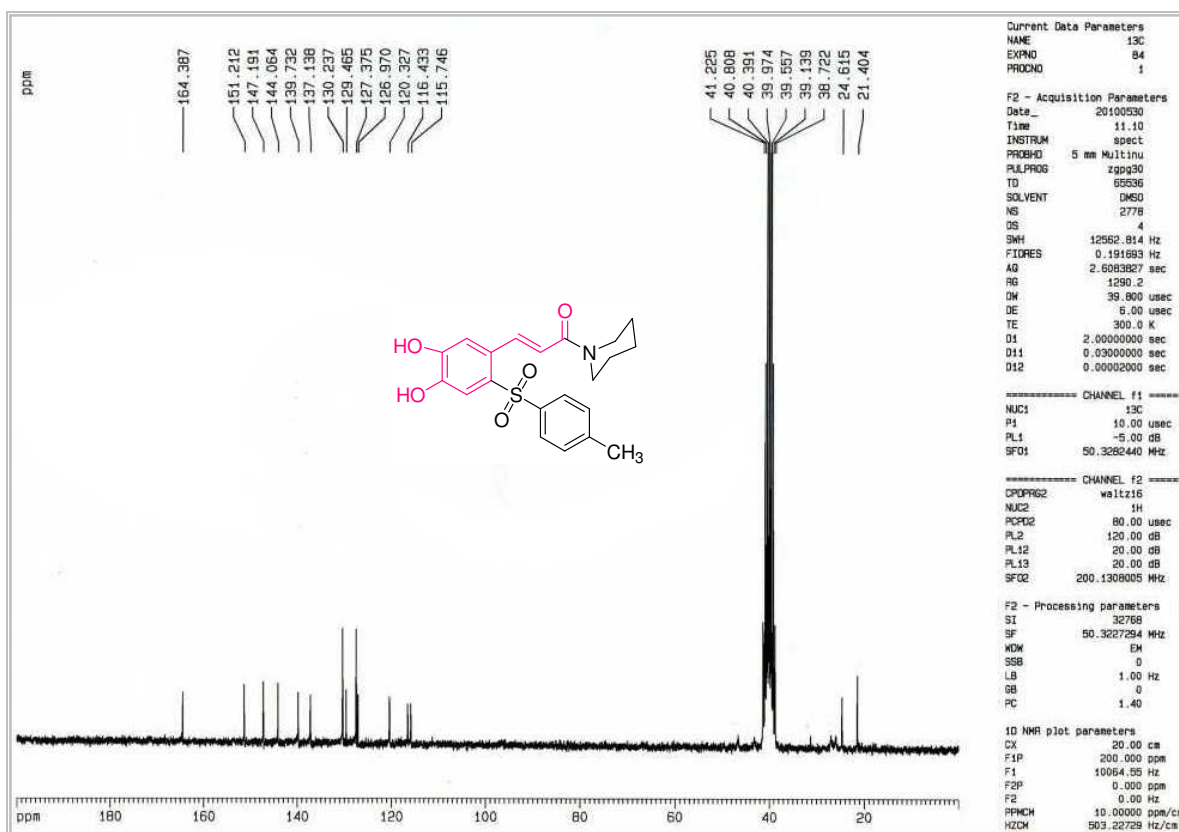


Figure S-2. ¹³C NMR spectrum of compound 5a in DMSO-d₆

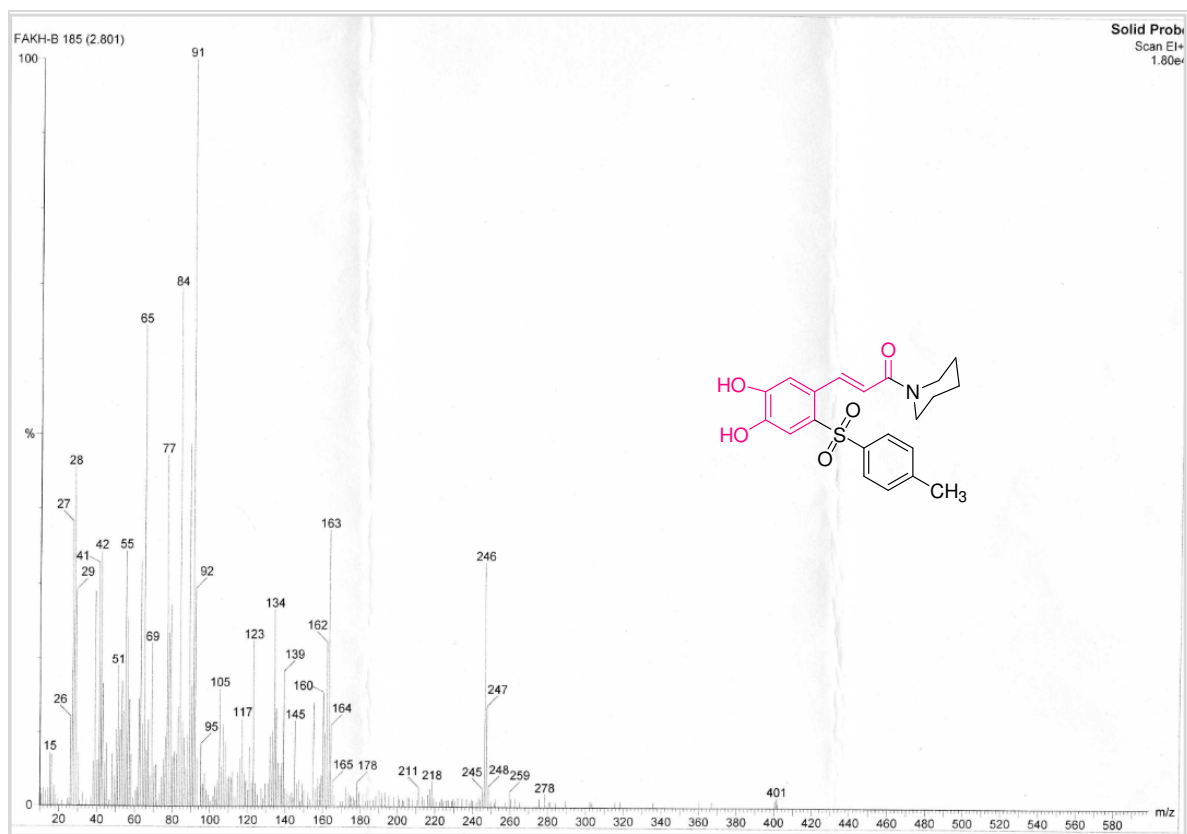


Figure S-3. Mass spectrum of compound **5a**

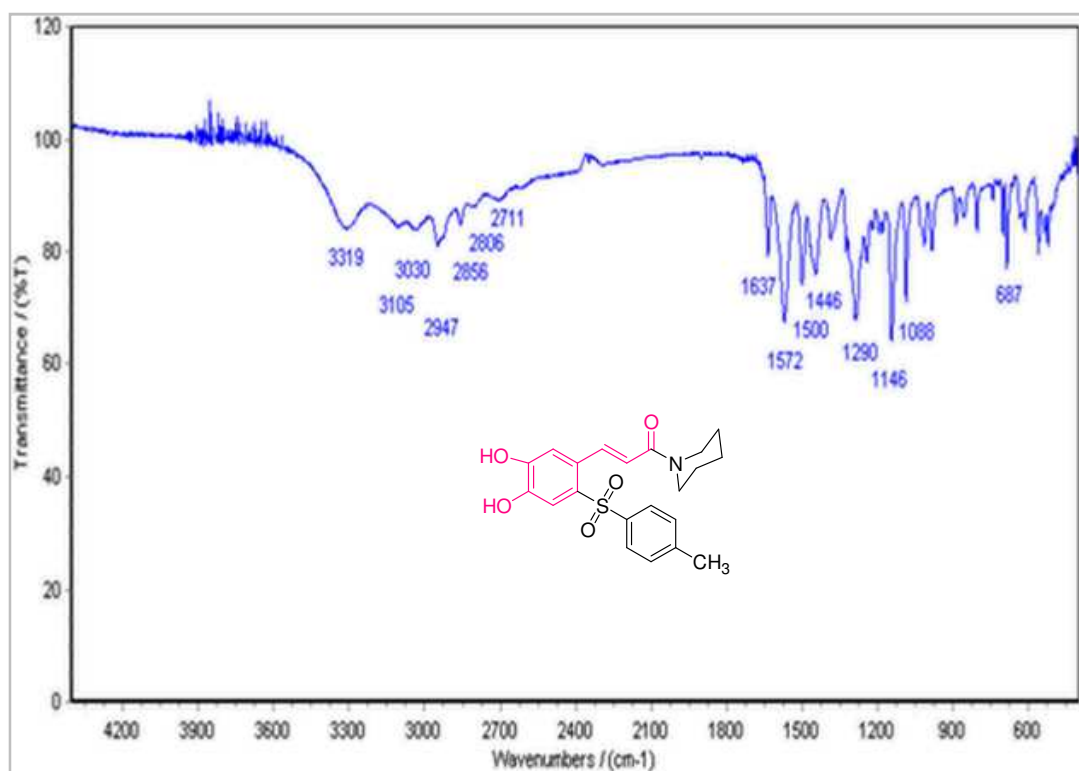


Figure S-4. FT-IR spectrum of compound **5a**

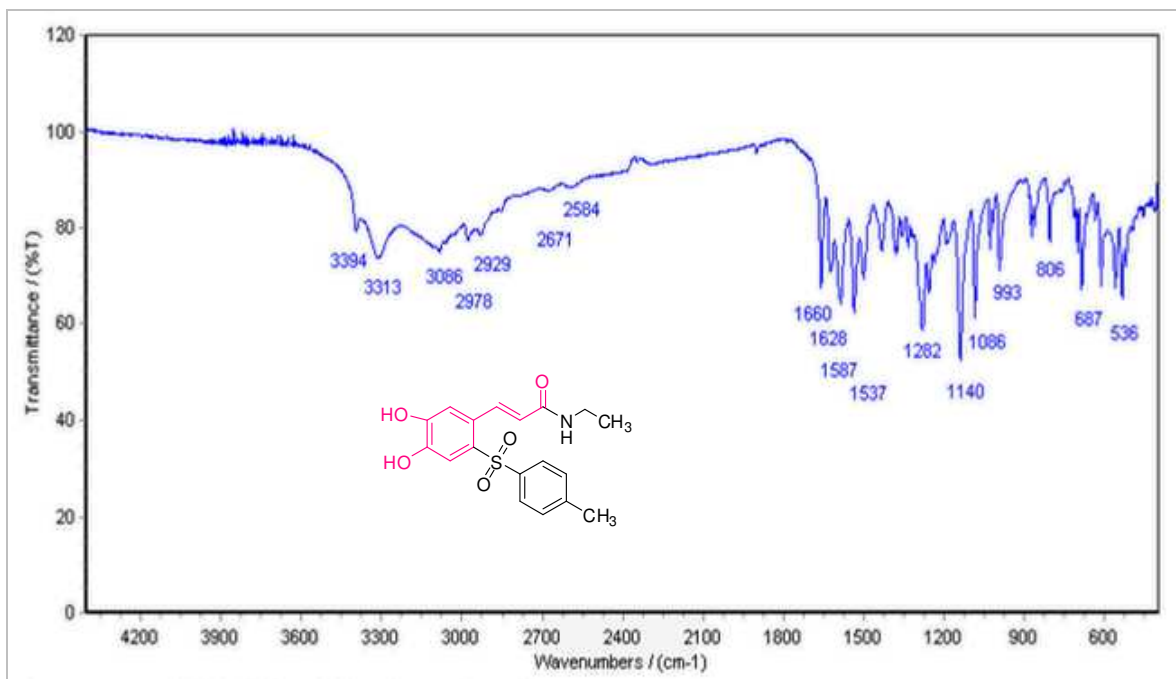


Figure S-7. FT-IR spectrum of compound **5b**

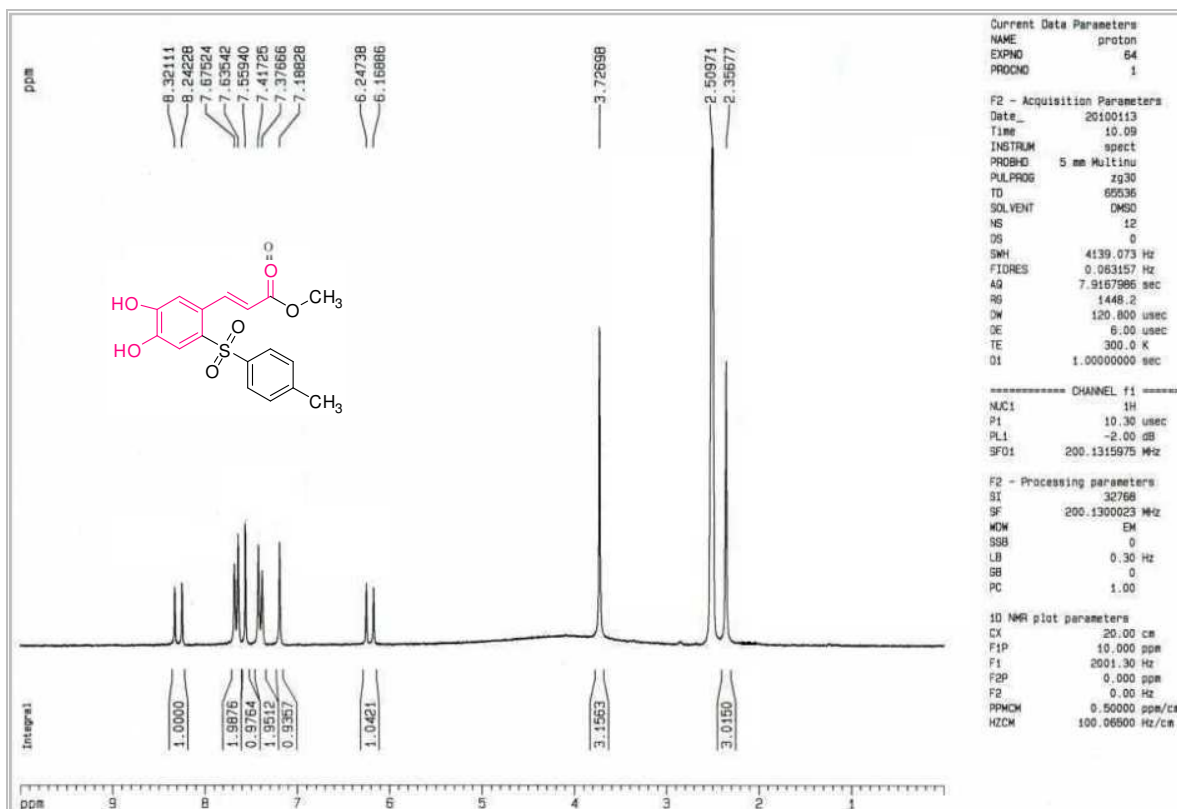


Figure S-8. ^1H NMR spectrum of compound **5c** in DMSO-d_6

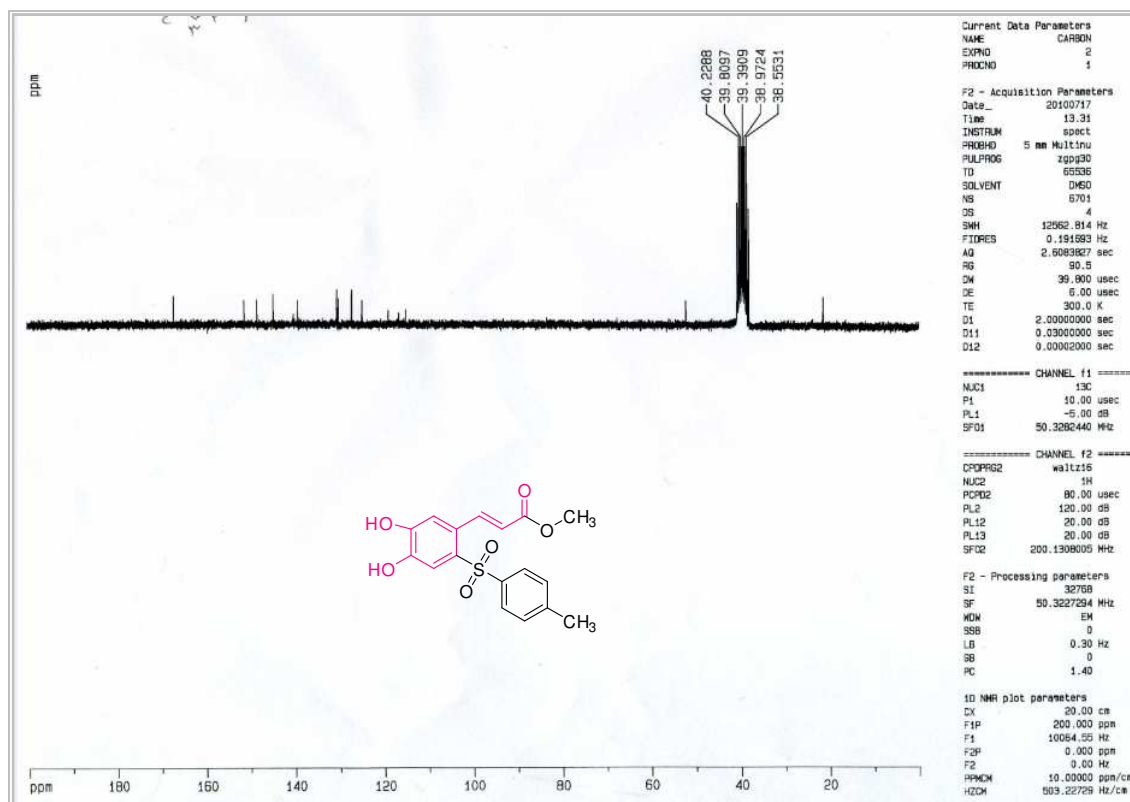


Figure S-9. ^{13}C NMR spectrum of compound **5c** in DMSO-d_6

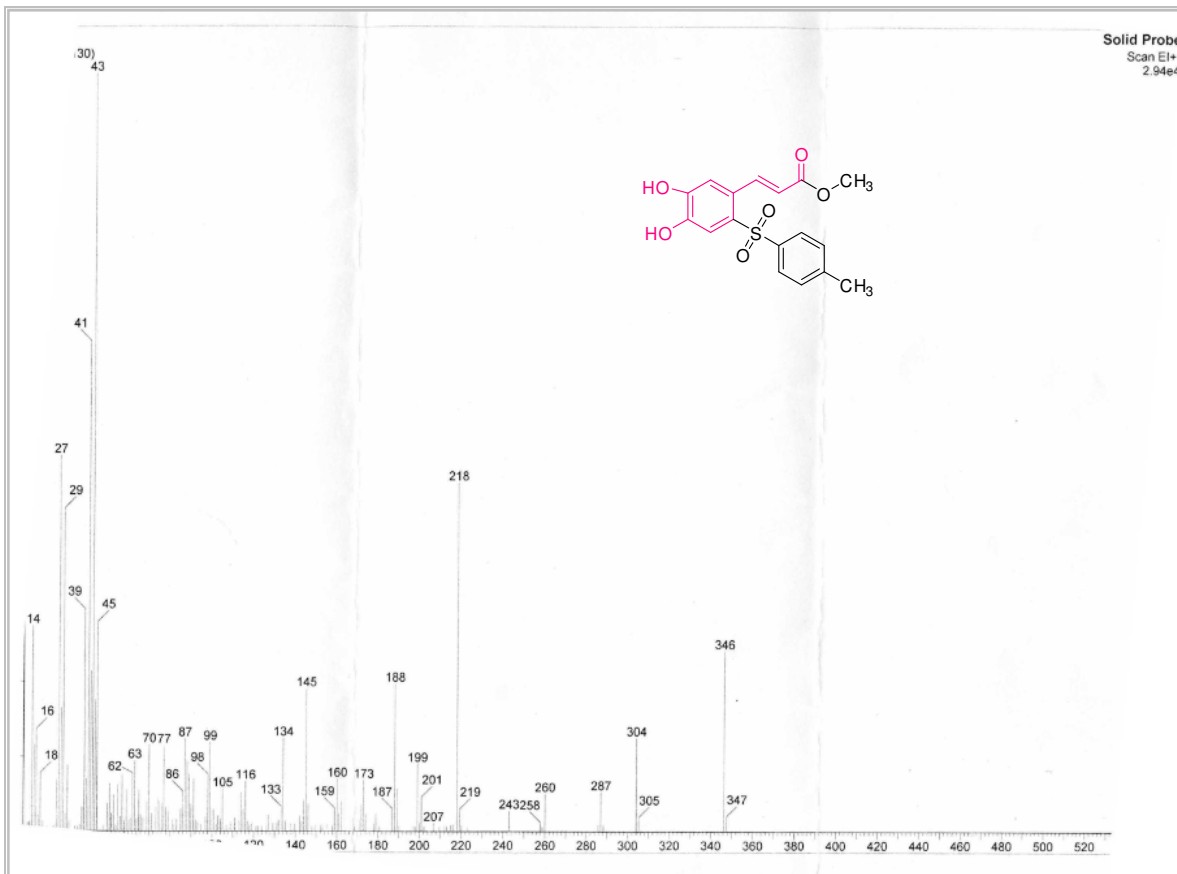


Figure S-10. Mass spectrum of compound 5c

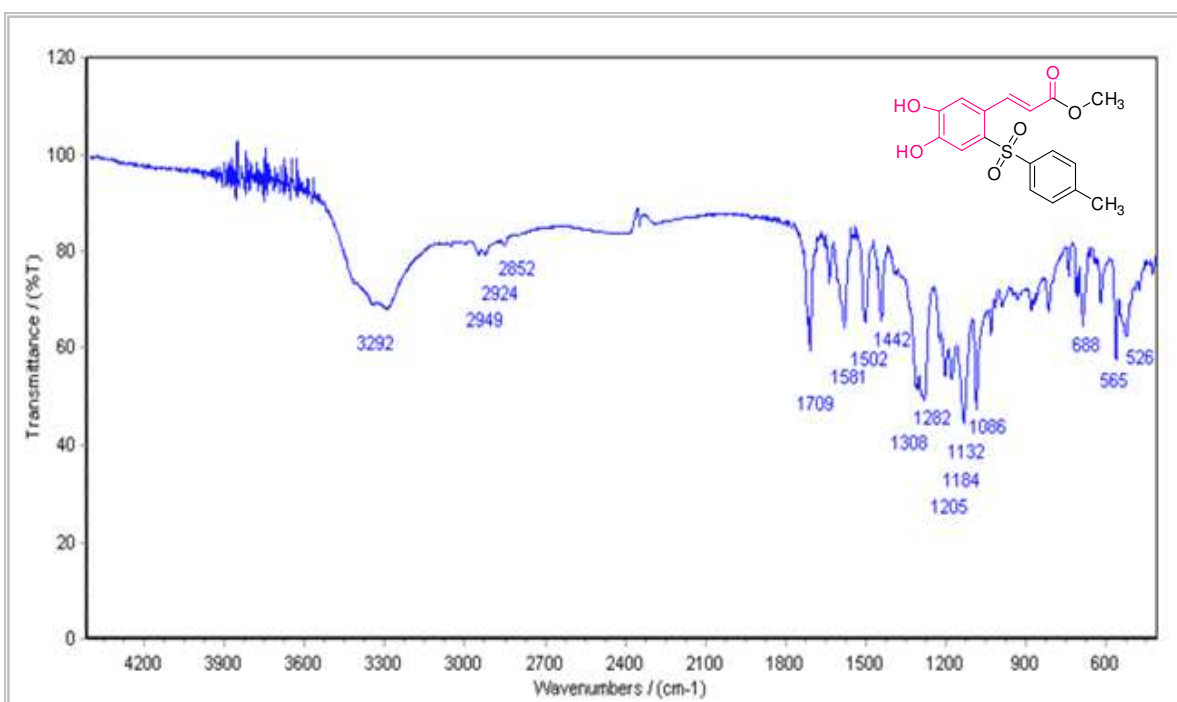


Figure S-11. FT-IR spectrum of compound 5c

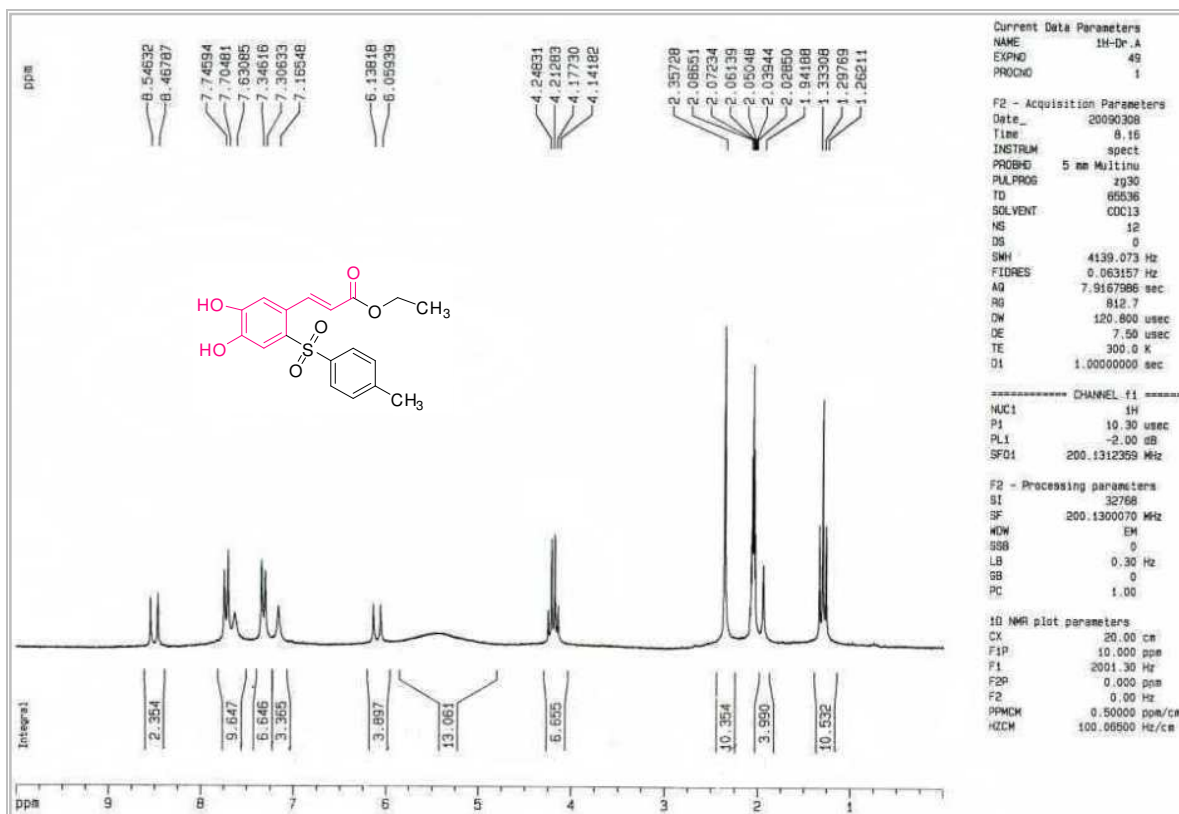


Figure S-12. ¹H NMR spectrum of compound **5d** in acetone-d₆

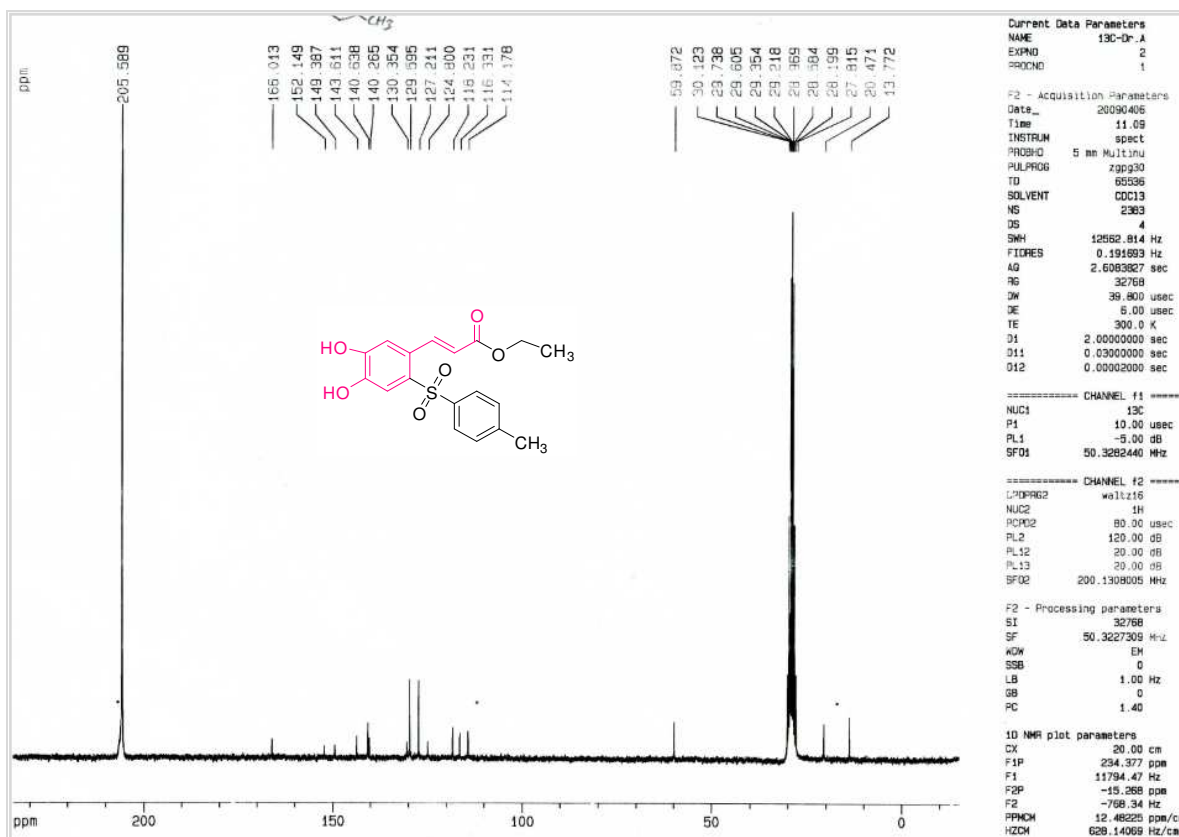


Figure S-13. ¹³C NMR spectrum of compound **5d** in acetone-d₆

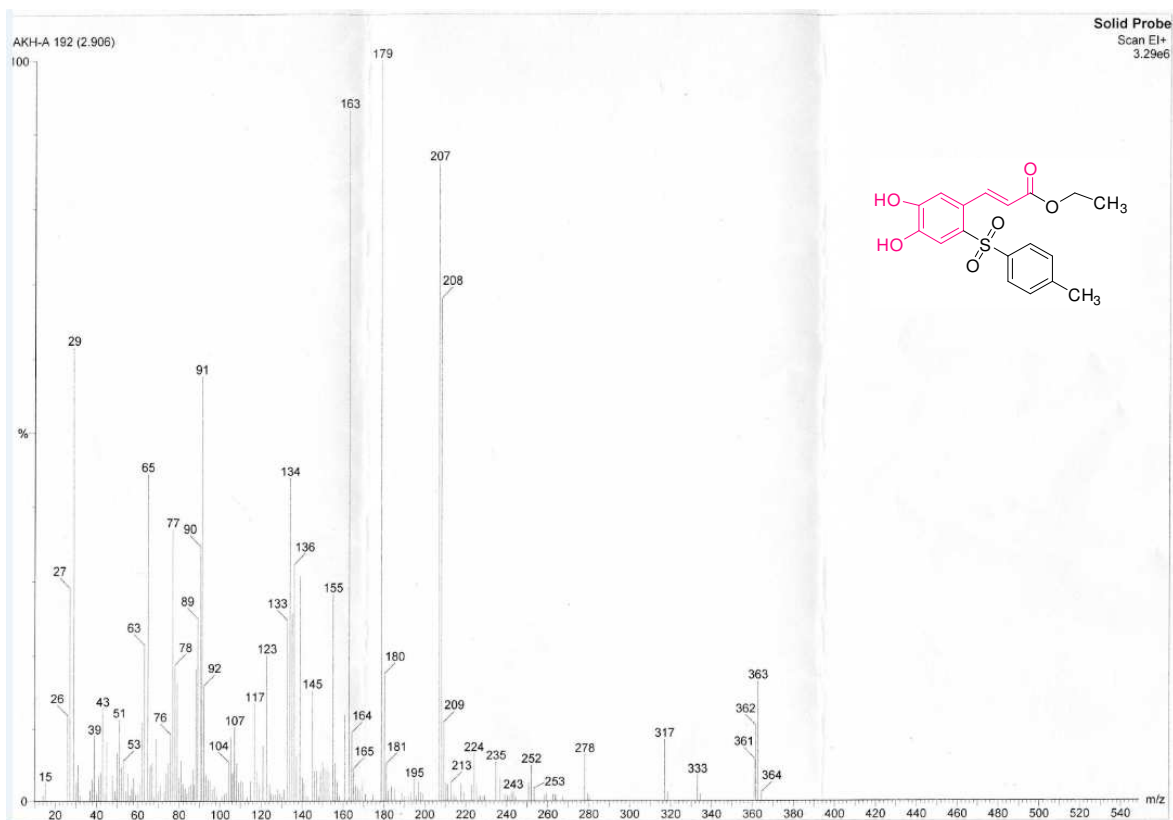


Figure S-14. Mass spectrum of compound **5d**

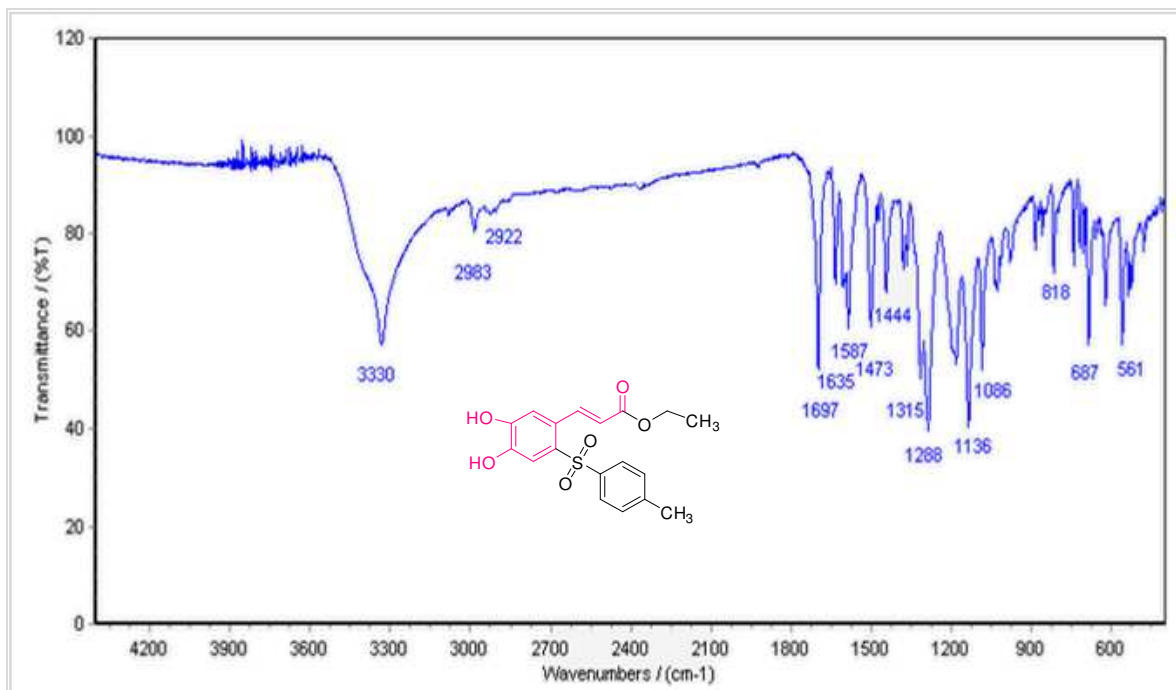


Figure S-15. FT-IR spectrum of compound **5d**

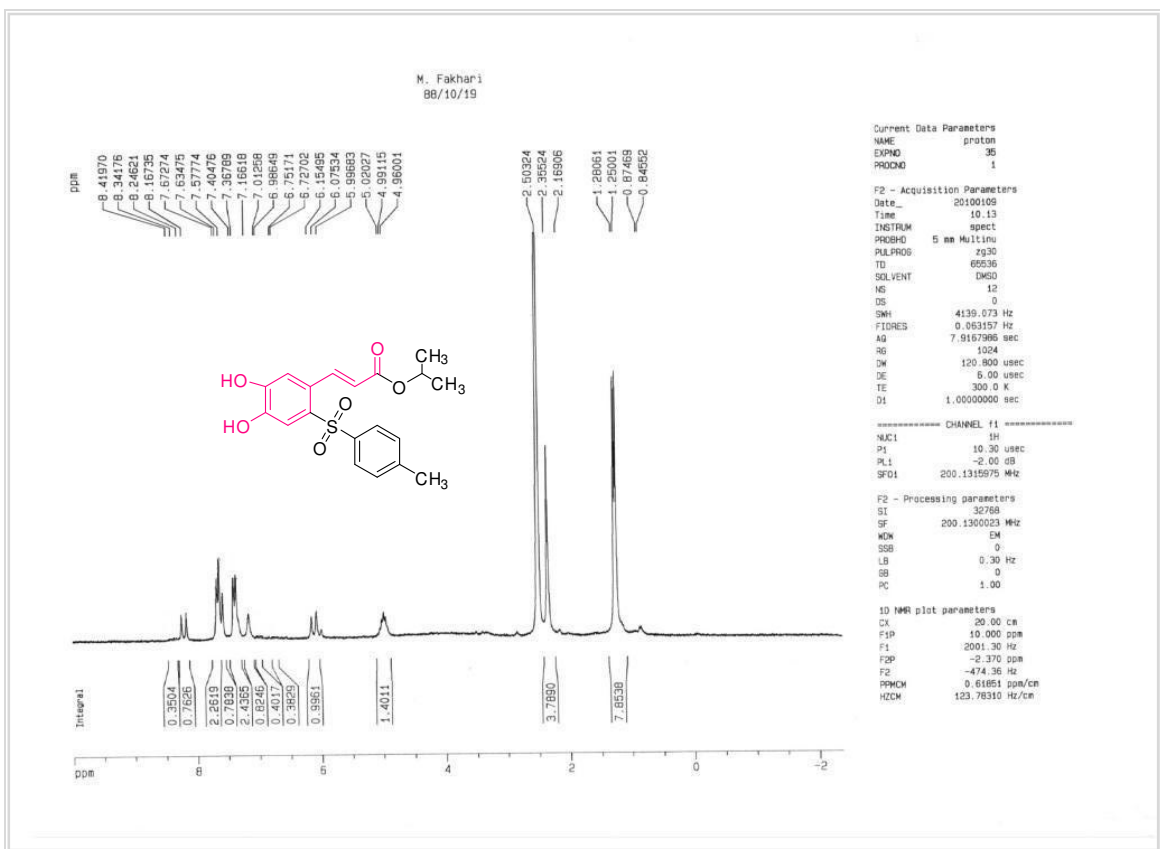


Figure S-16. ¹H NMR spectrum of compound **5e** in DMSO-d₆

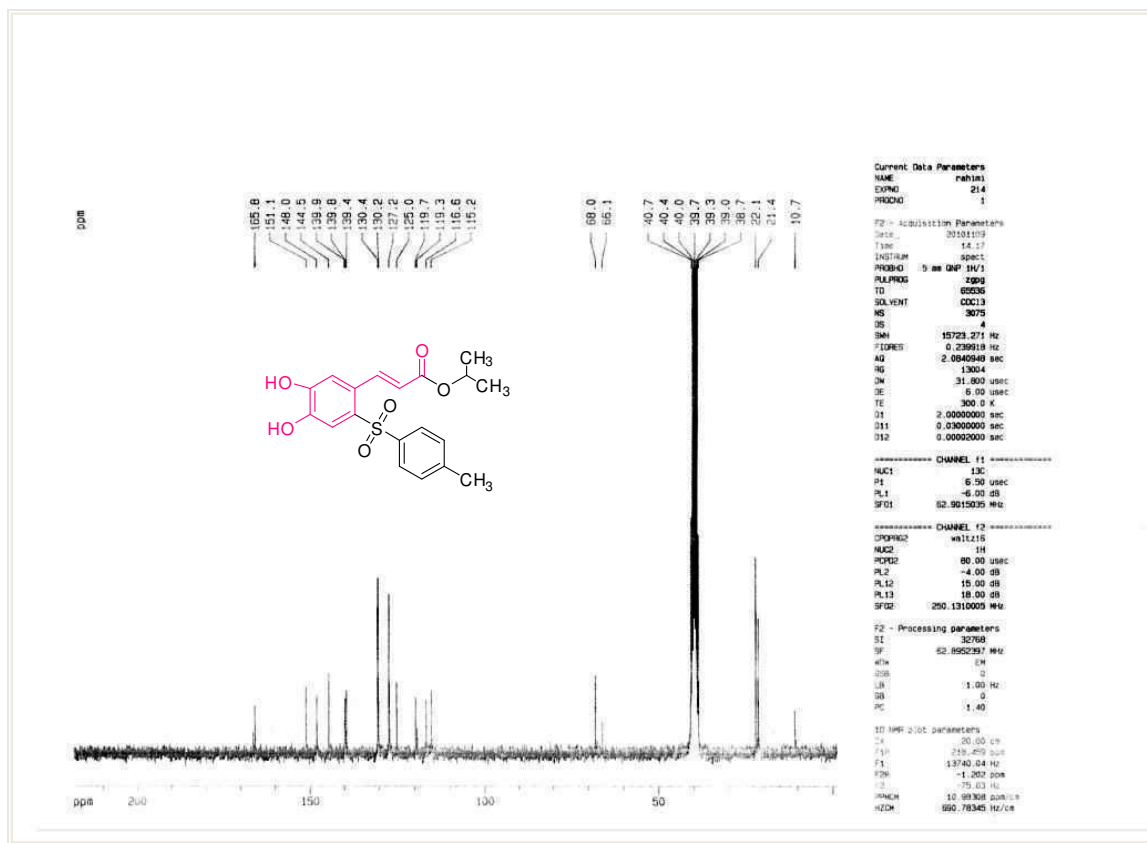


Figure S-17. ¹³C NMR spectrum of compound **5e** in DMSO-d₆

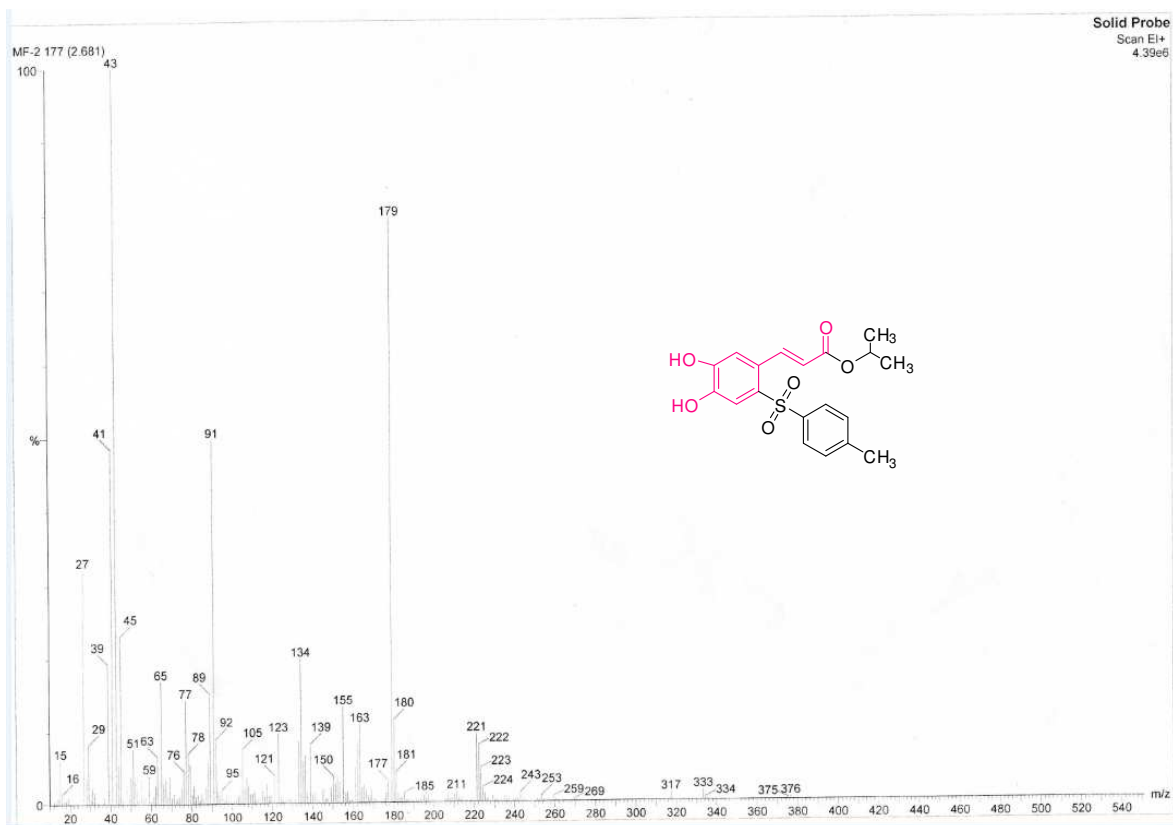


Figure S-18. Mass spectrum of compound 5e

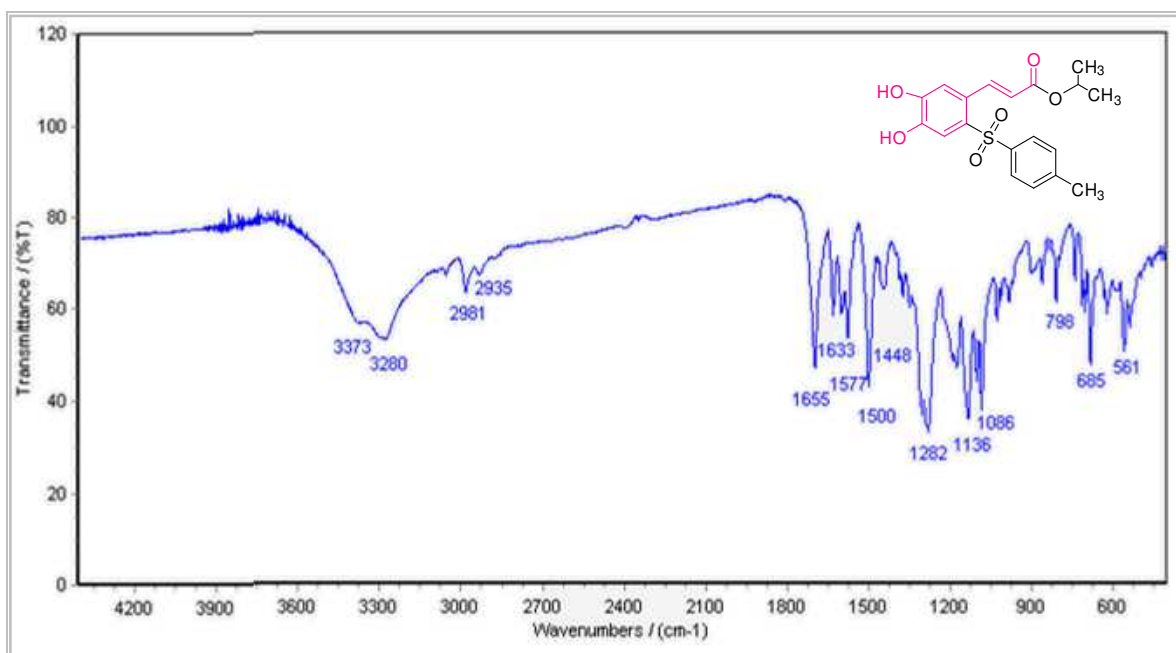


Figure S-19. FT-IR spectrum of compound 5e