Electronic Supplementary Material (ESI) for RSC Advances.

SUPPORTING INFORMATION

Three component synthesis of 2-oxindole *via* sequential Michael addition, intramolecular cyclization and aromatization

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and Nattamai Bhuvanesh

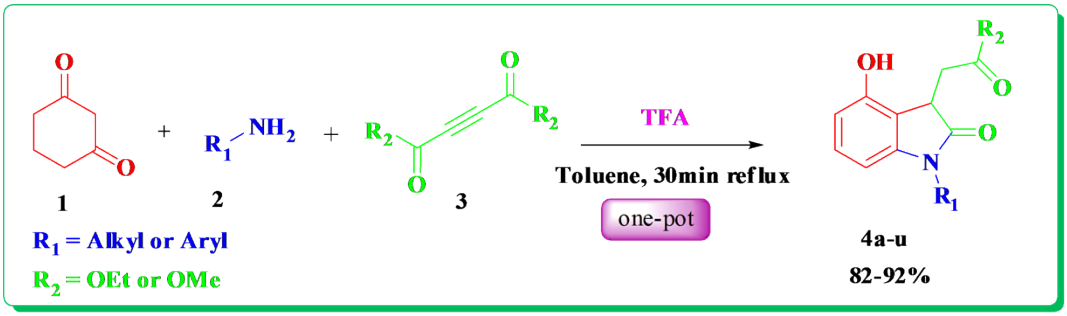

Table of Contents

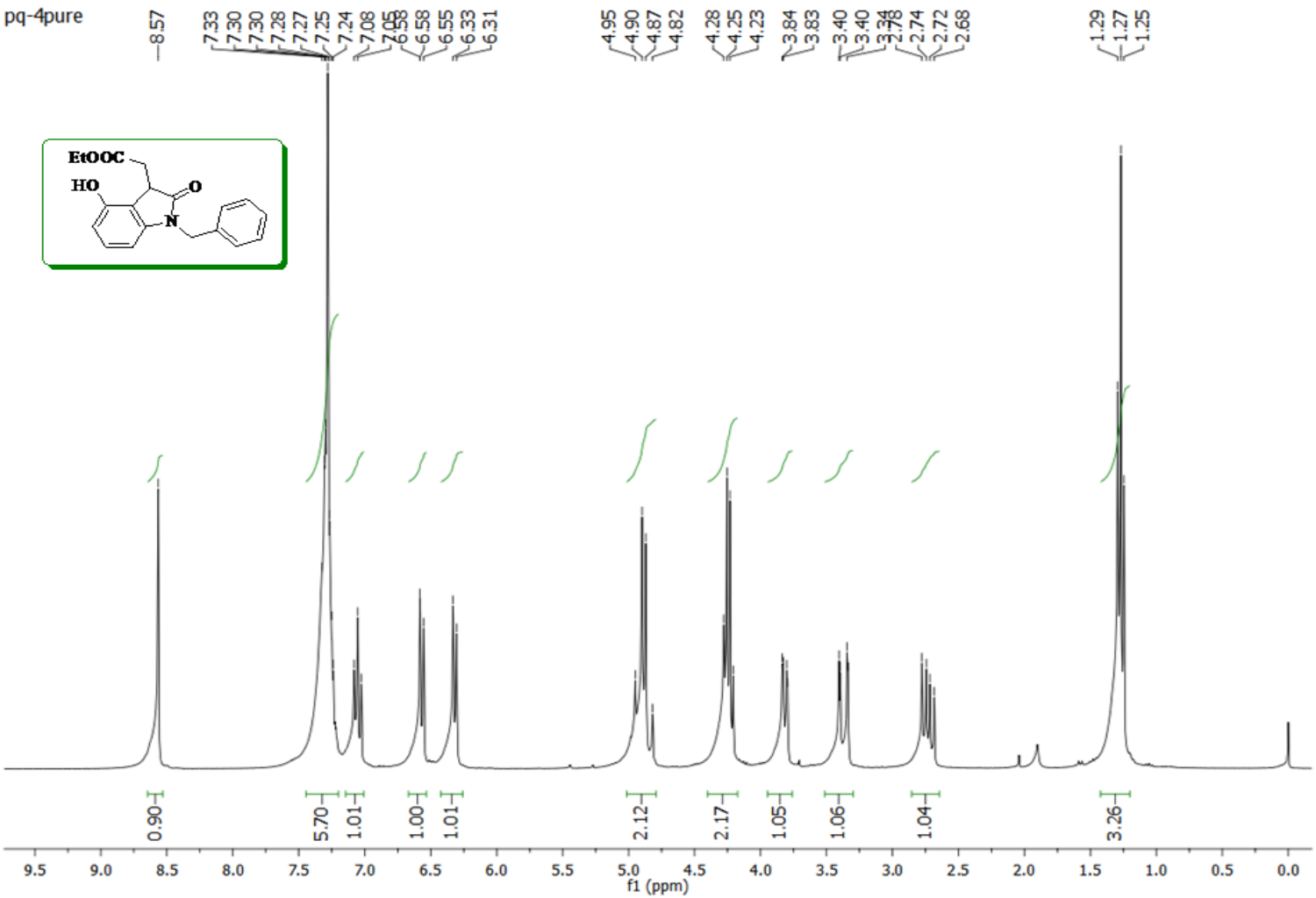

Figure S1. ^{1}H NMR spectrum of **4a**

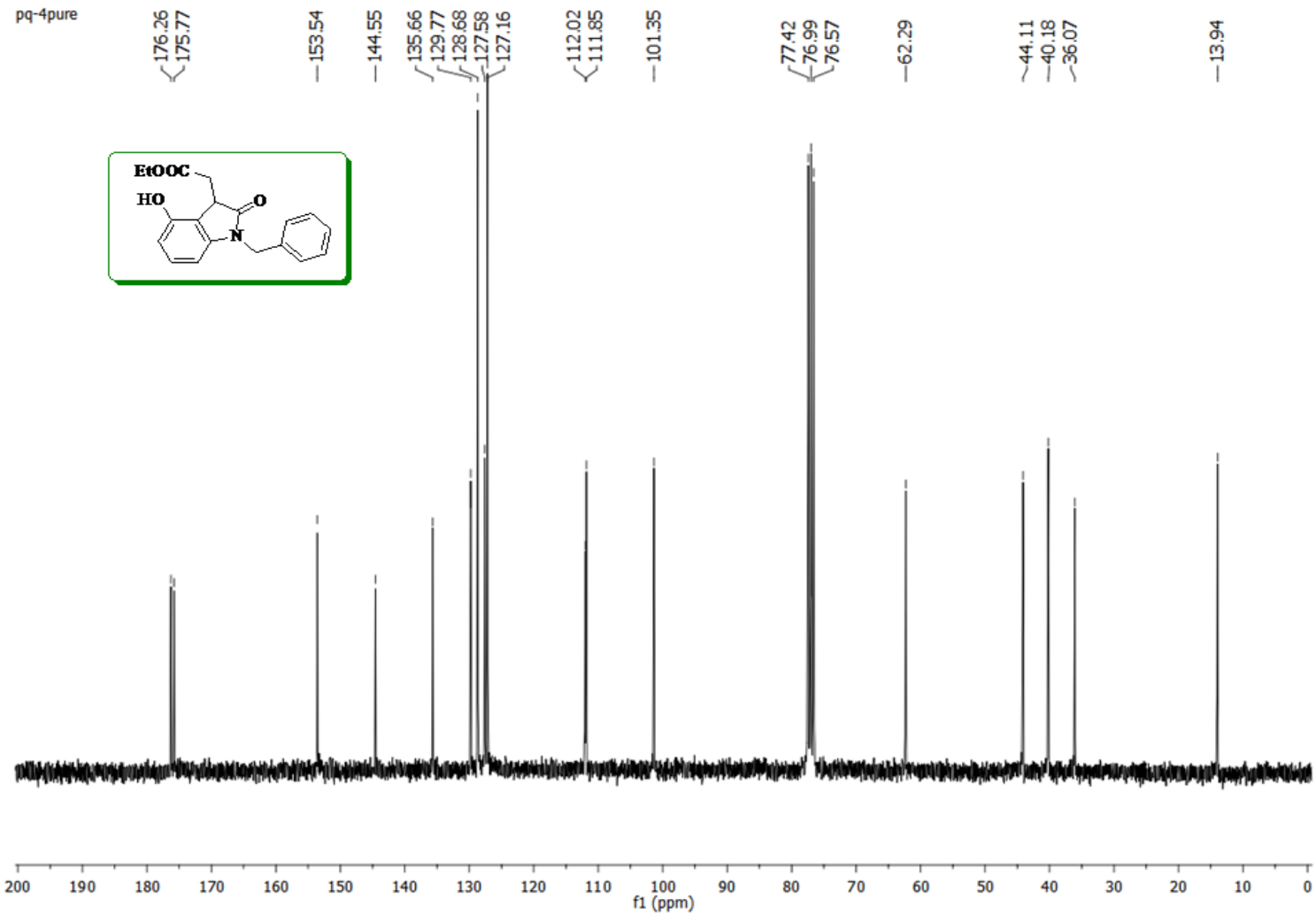

Figure S2. ^{13}C NMR spectrum of **4a**

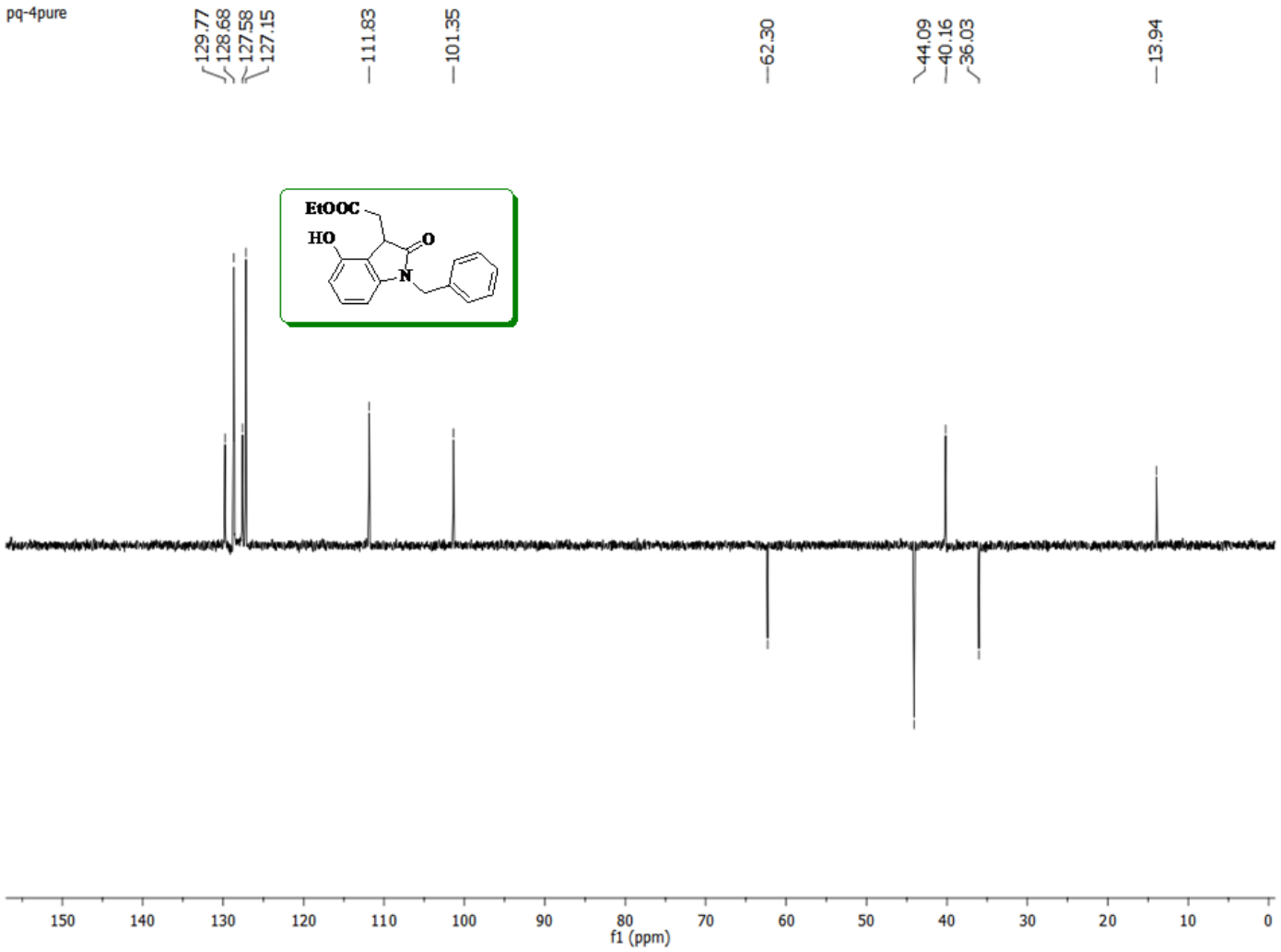

Figure S4. DEPT-135 spectrum of **4a**

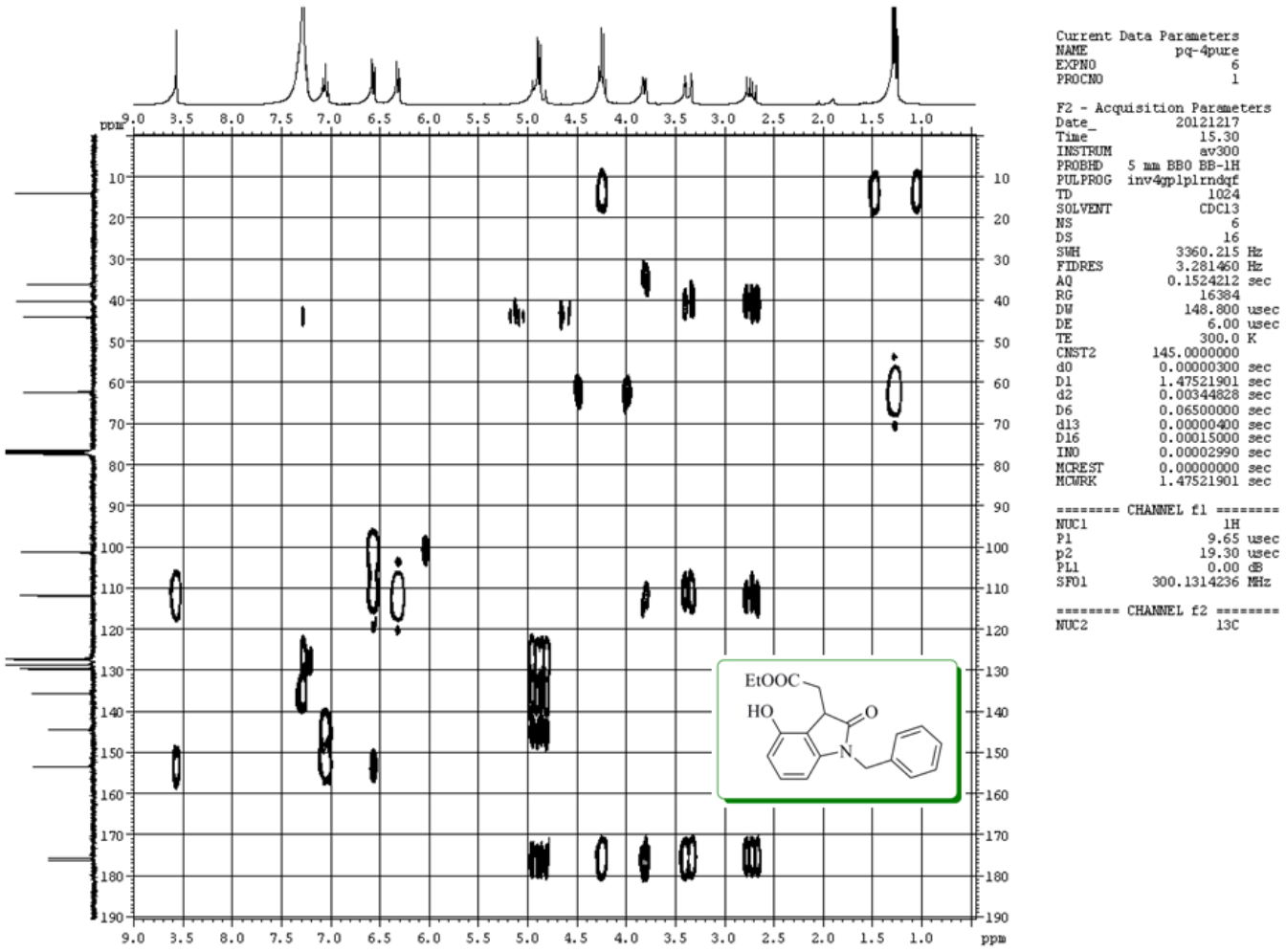

Figure S4. HMBC spectrum of **4a**

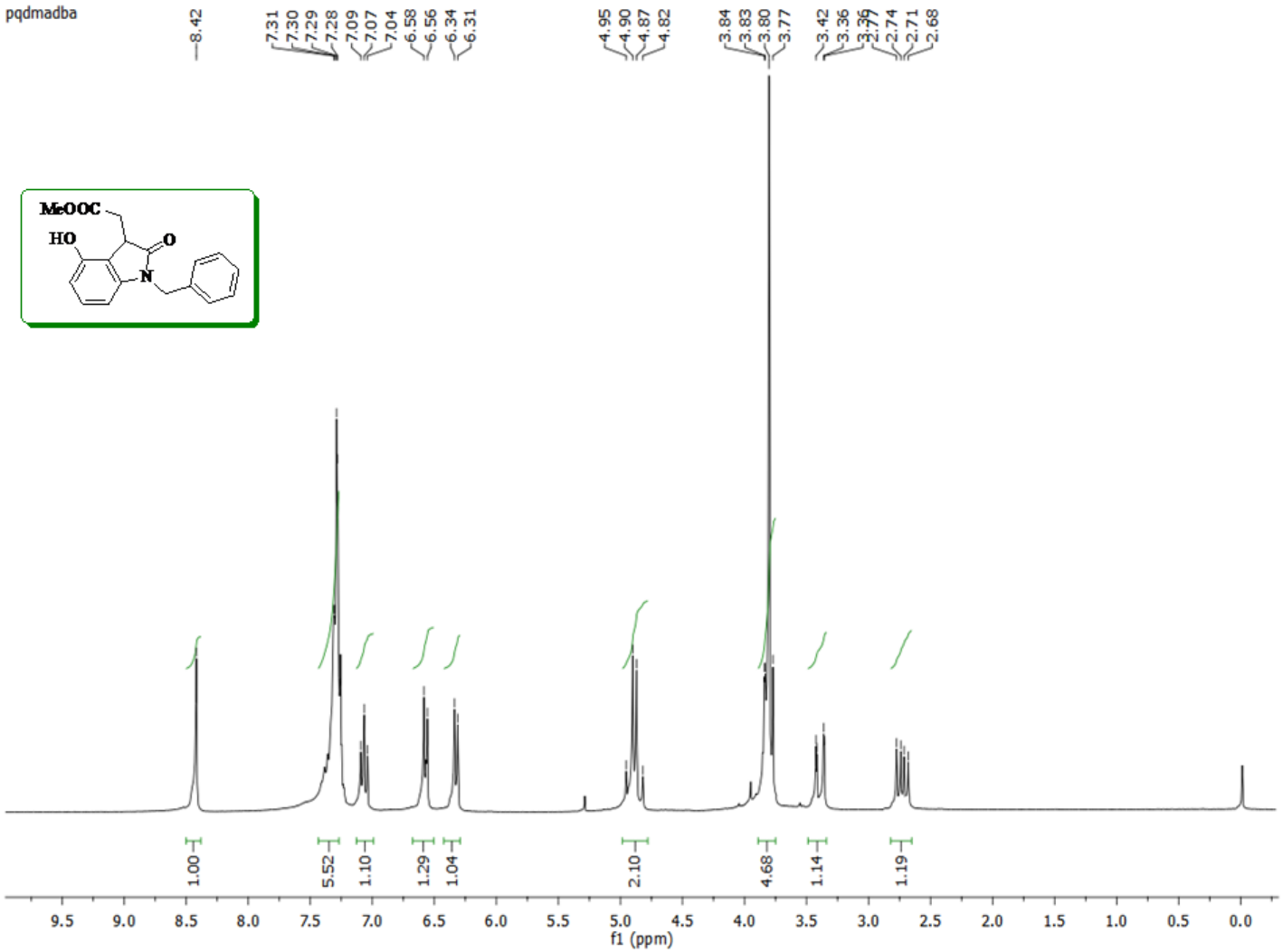

Figure S5. ^{1}H NMR spectrum of **4b**

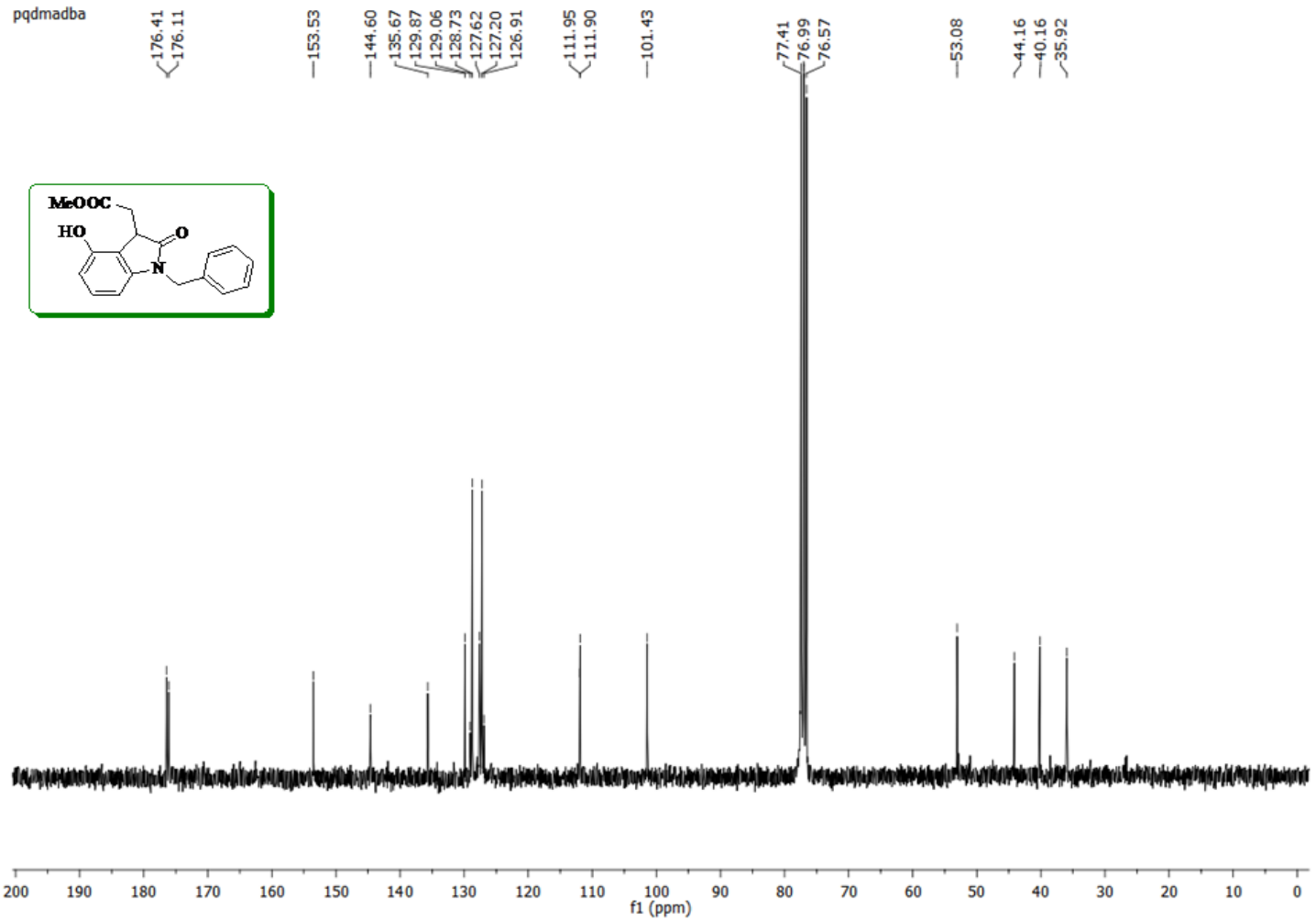

Figure S6. ^{13}C NMR spectrum of **4b**

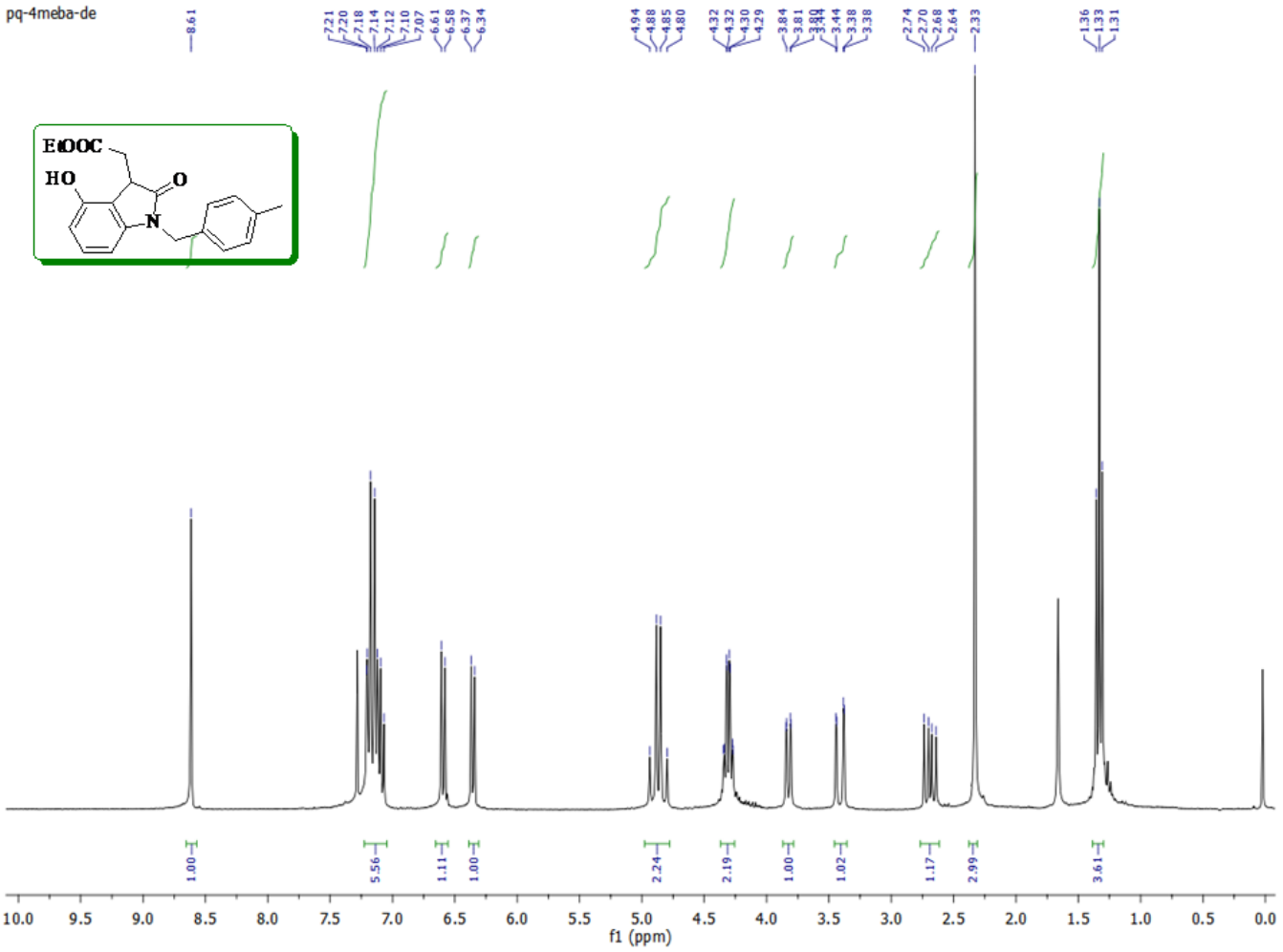

Figure S7. ^{1}H NMR spectrum of **4c**

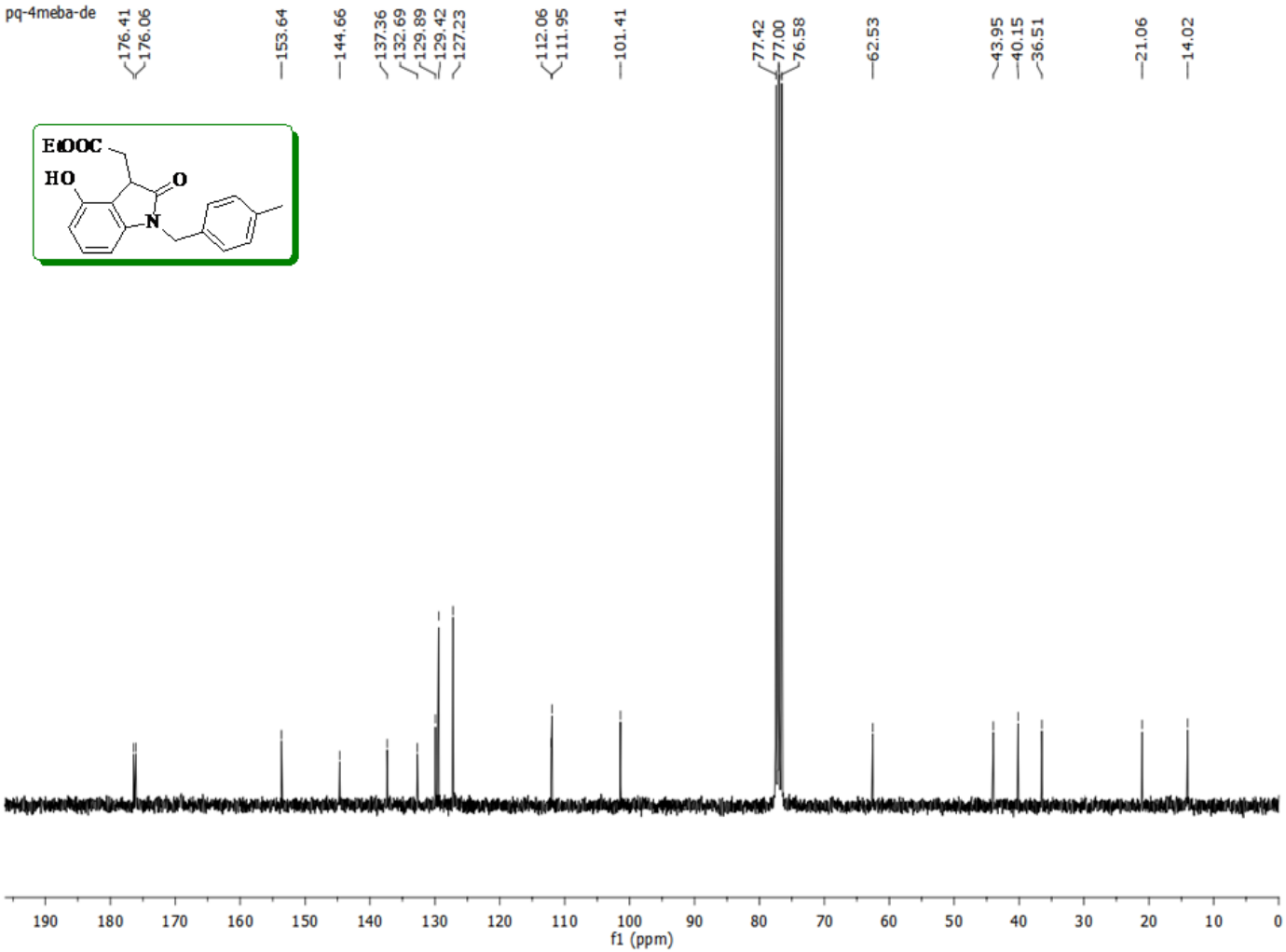

Figure S8. ^{13}C NMR spectrum of **4c**

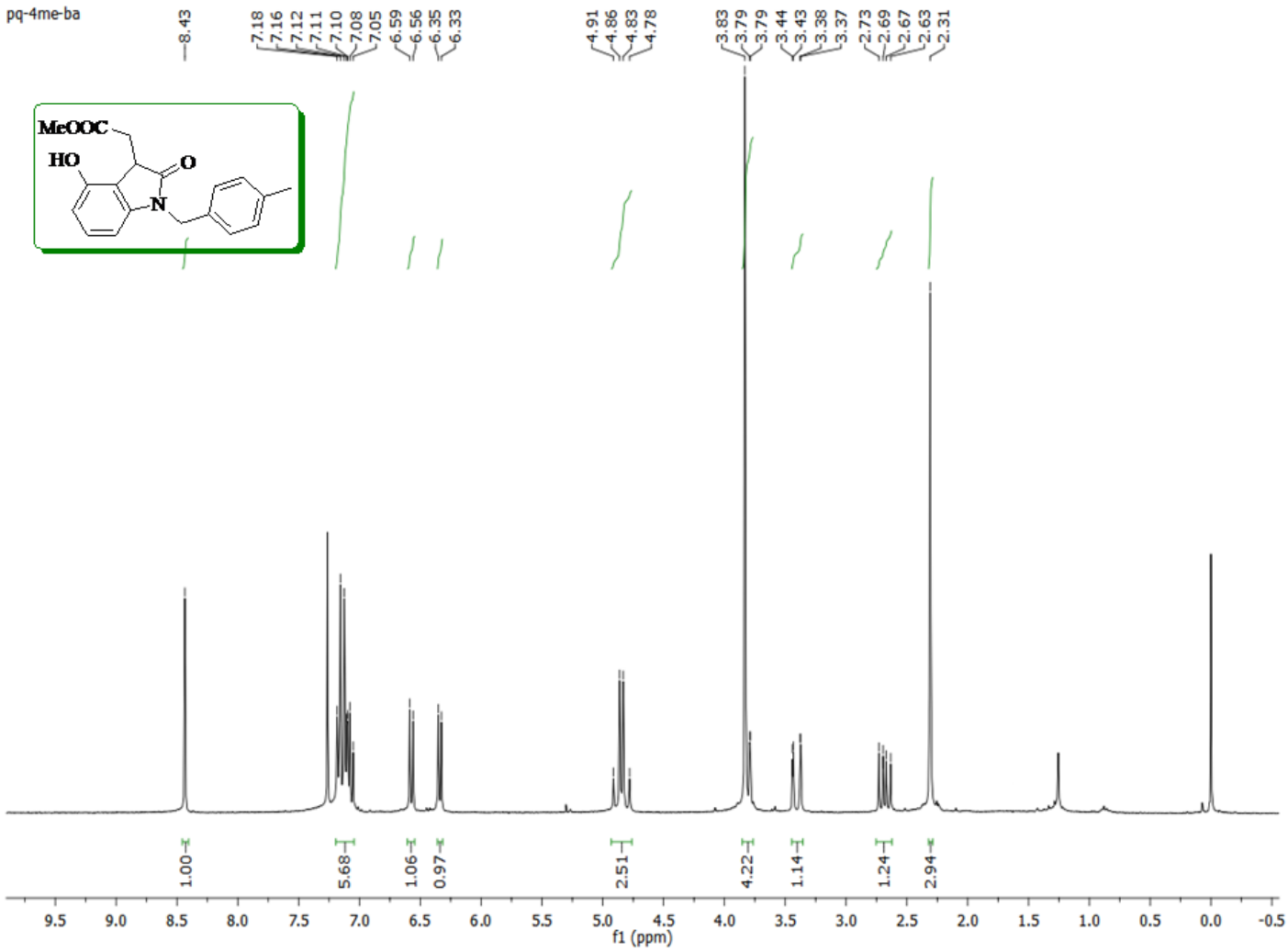

Figure S9. ^{1}H NMR spectrum of **4d**

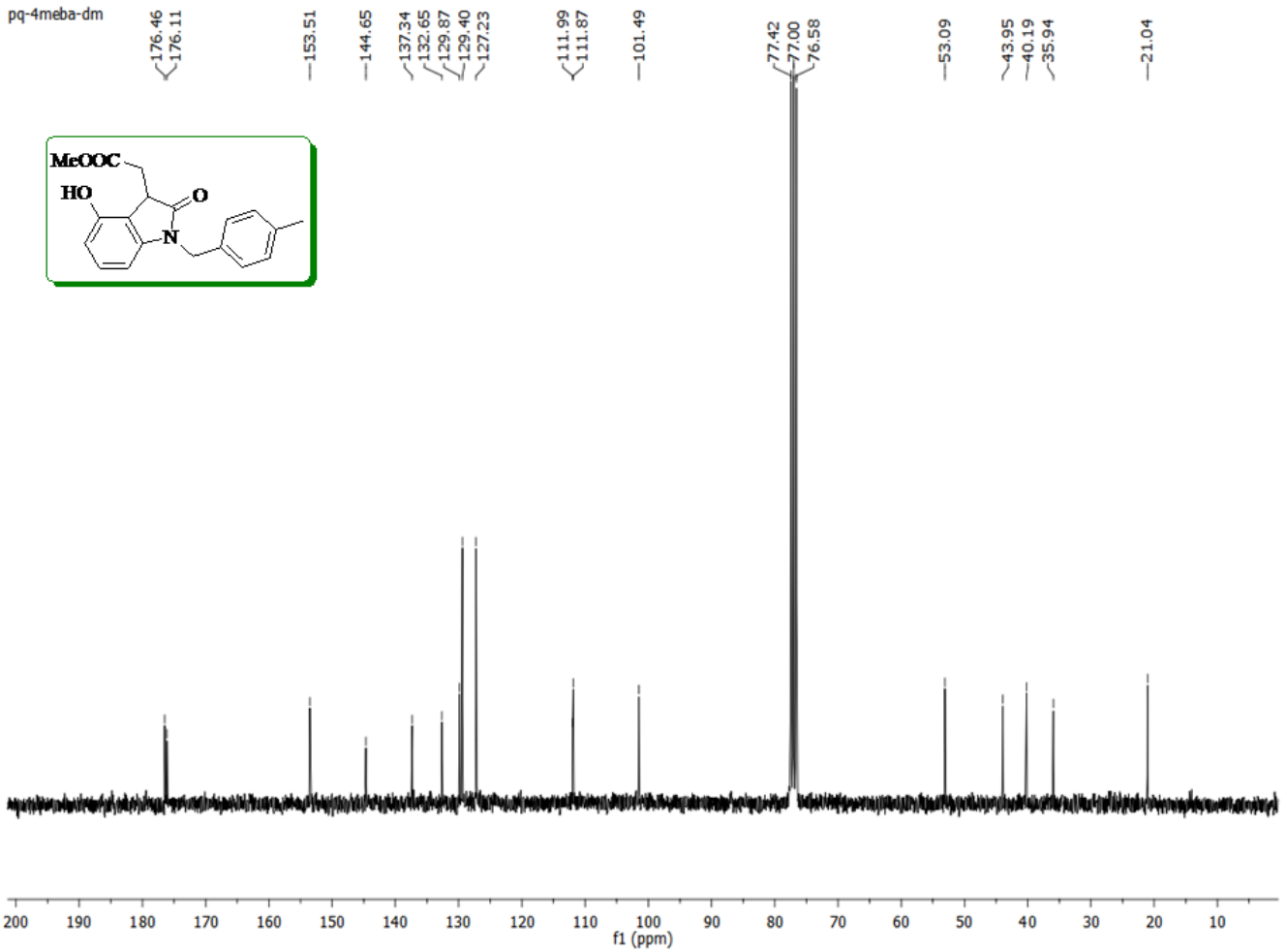

Figure S10. ^{13}C NMR spectrum of **4d**

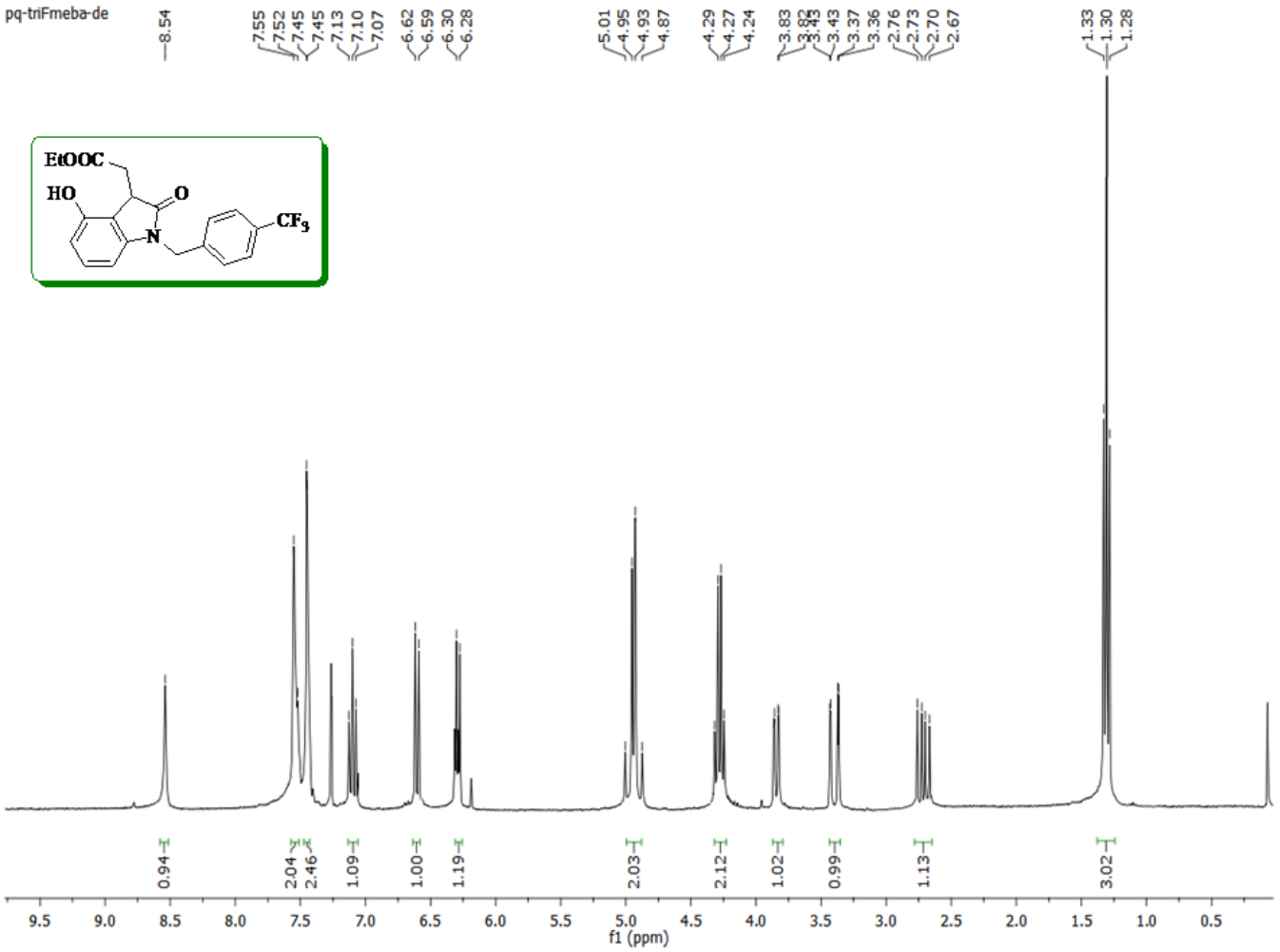

Figure S11. ^{1}H NMR spectrum of **4e**

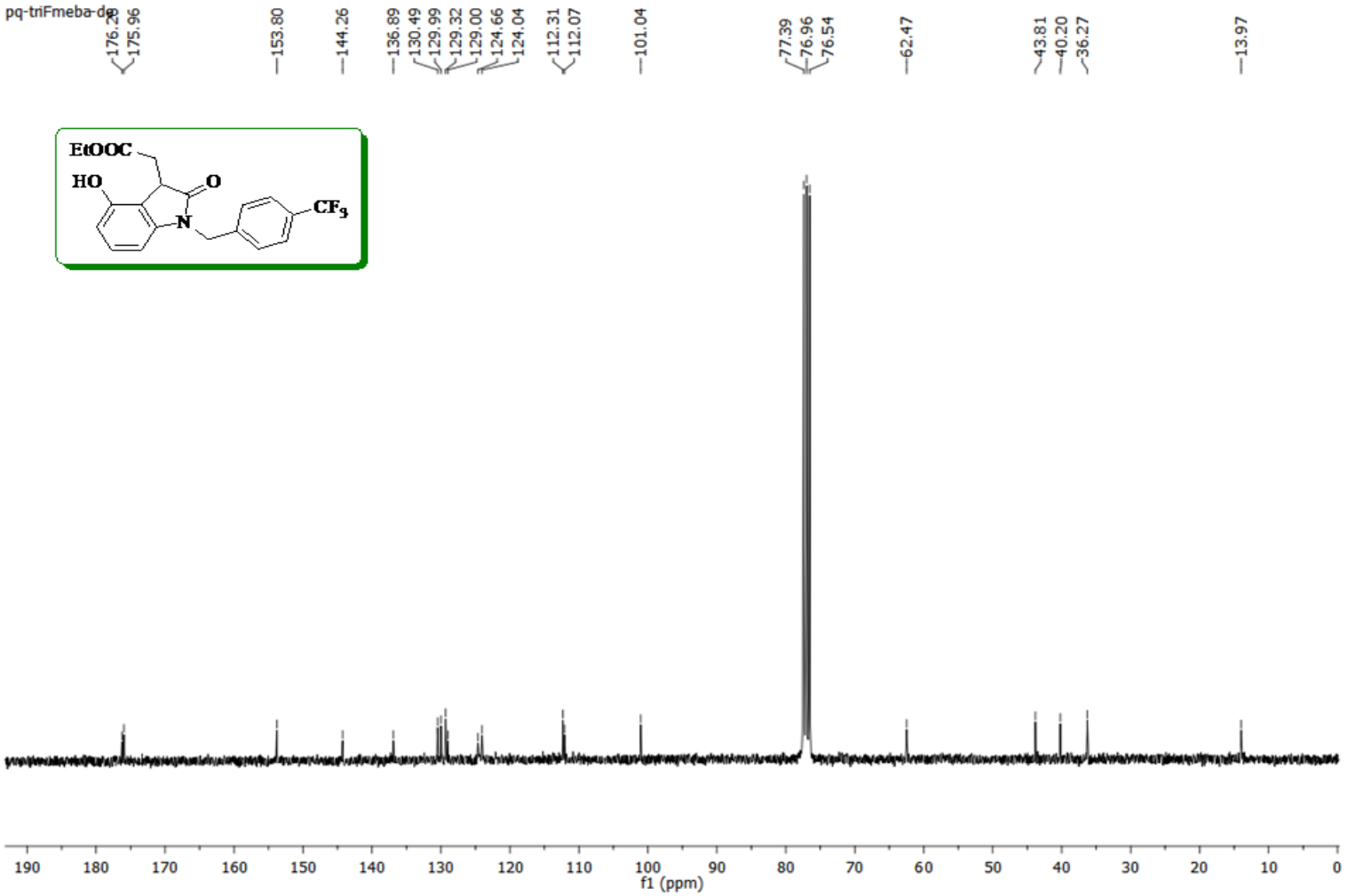

Figure S12. ^{13}C NMR spectrum of **4e**

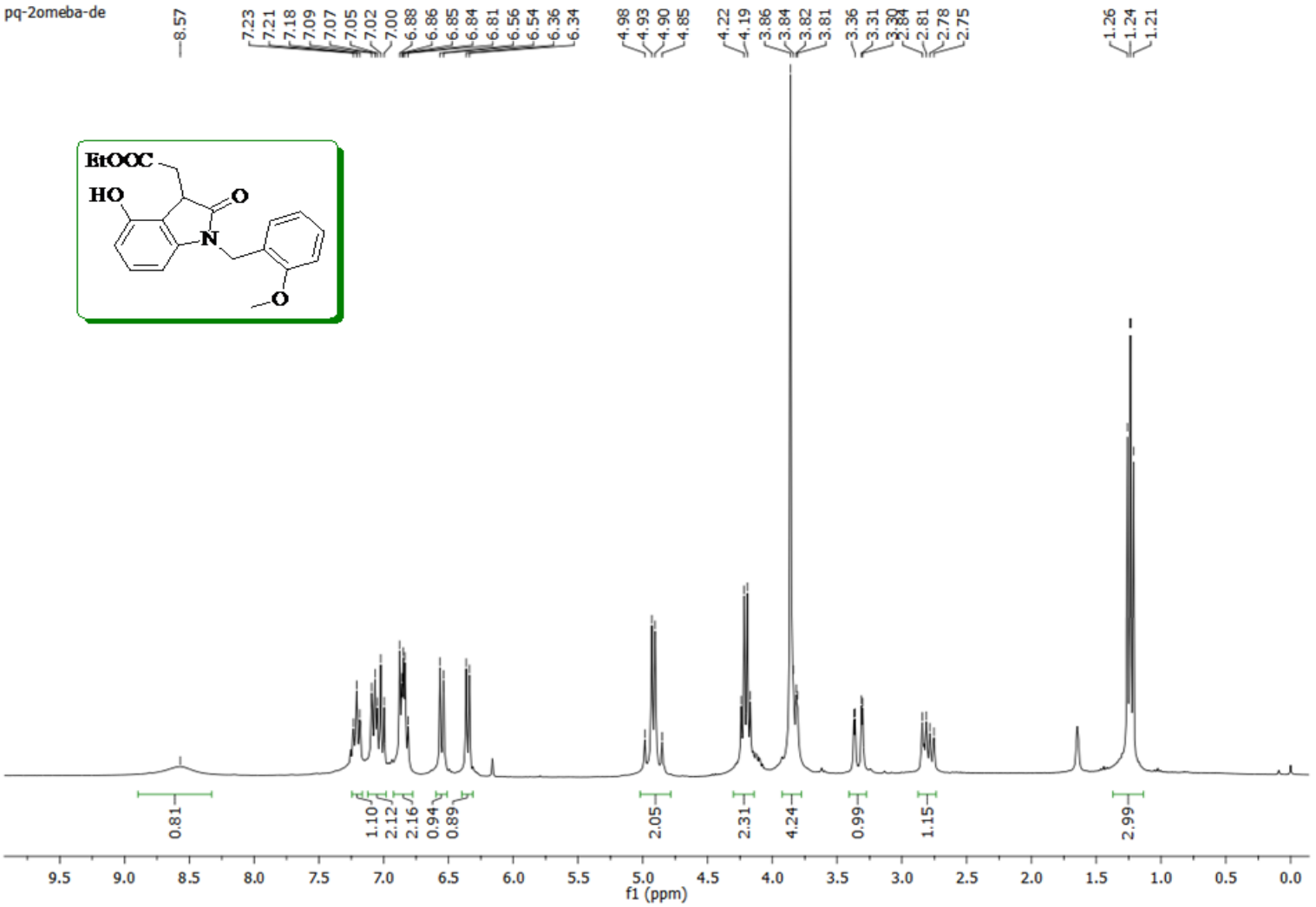

Figure S13. ^{1}H NMR spectrum of **4f**

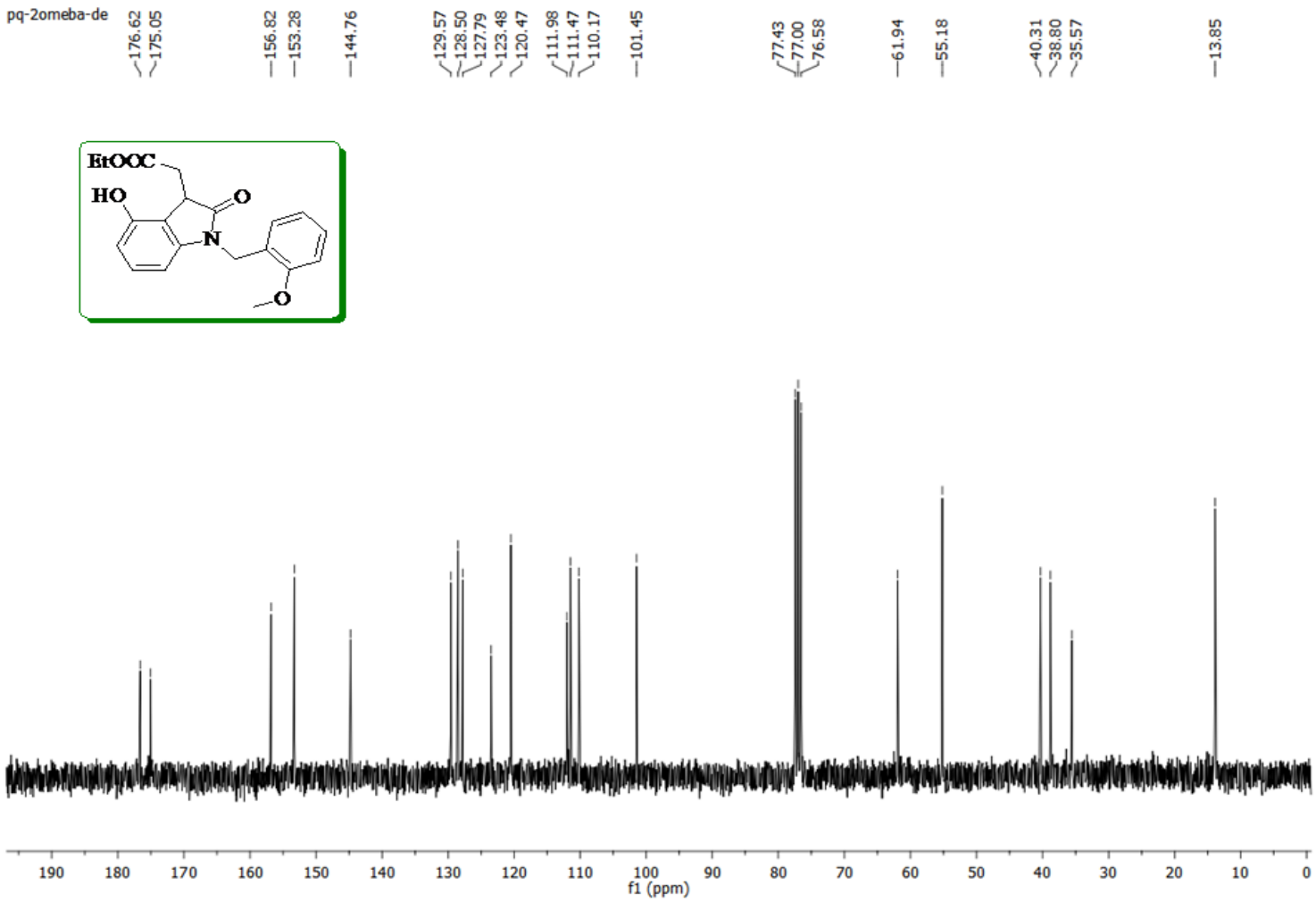

Figure S14. ^{13}C NMR spectrum of **4f**

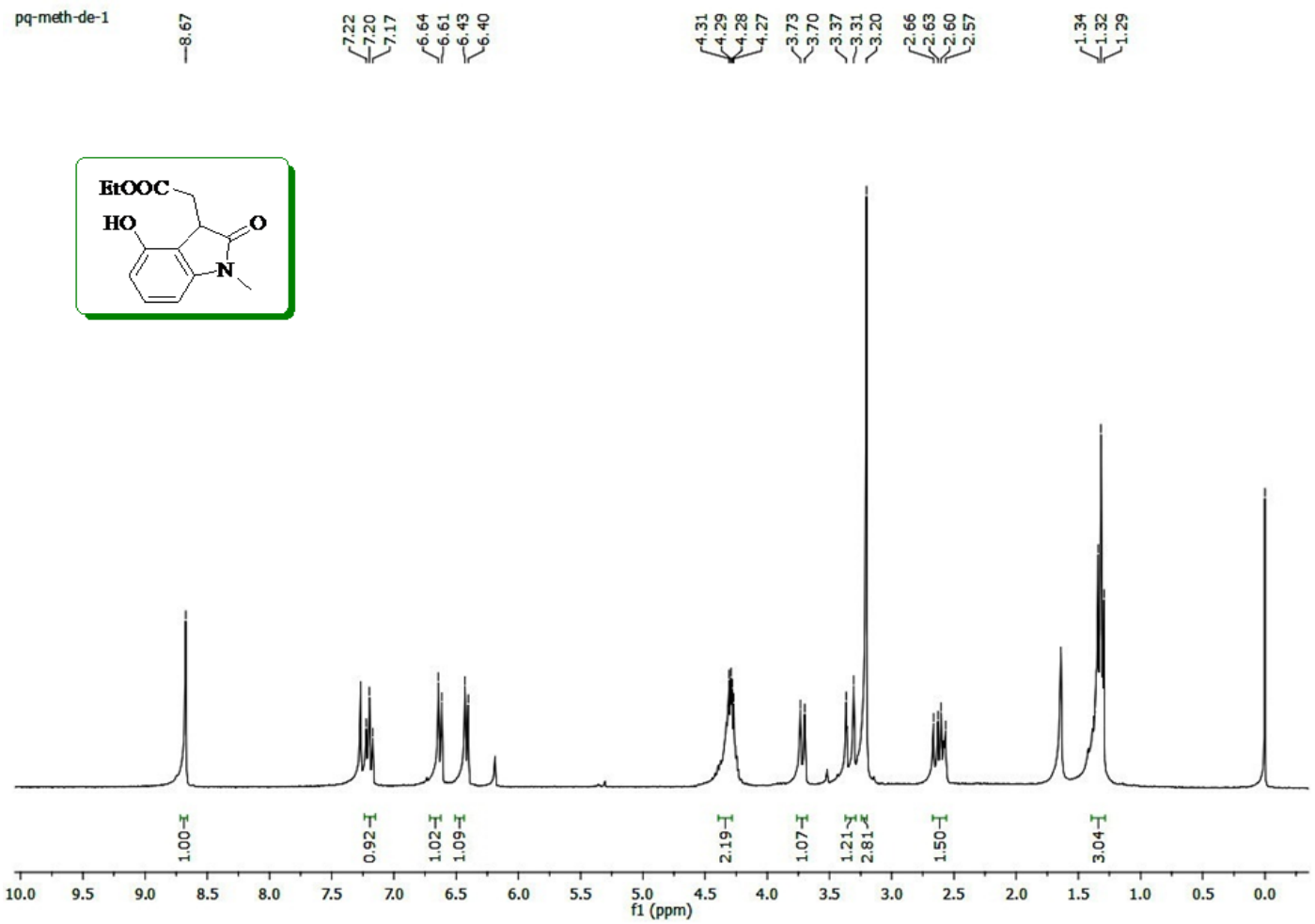

Figure S15. ^{1}H NMR spectrum of **4g**

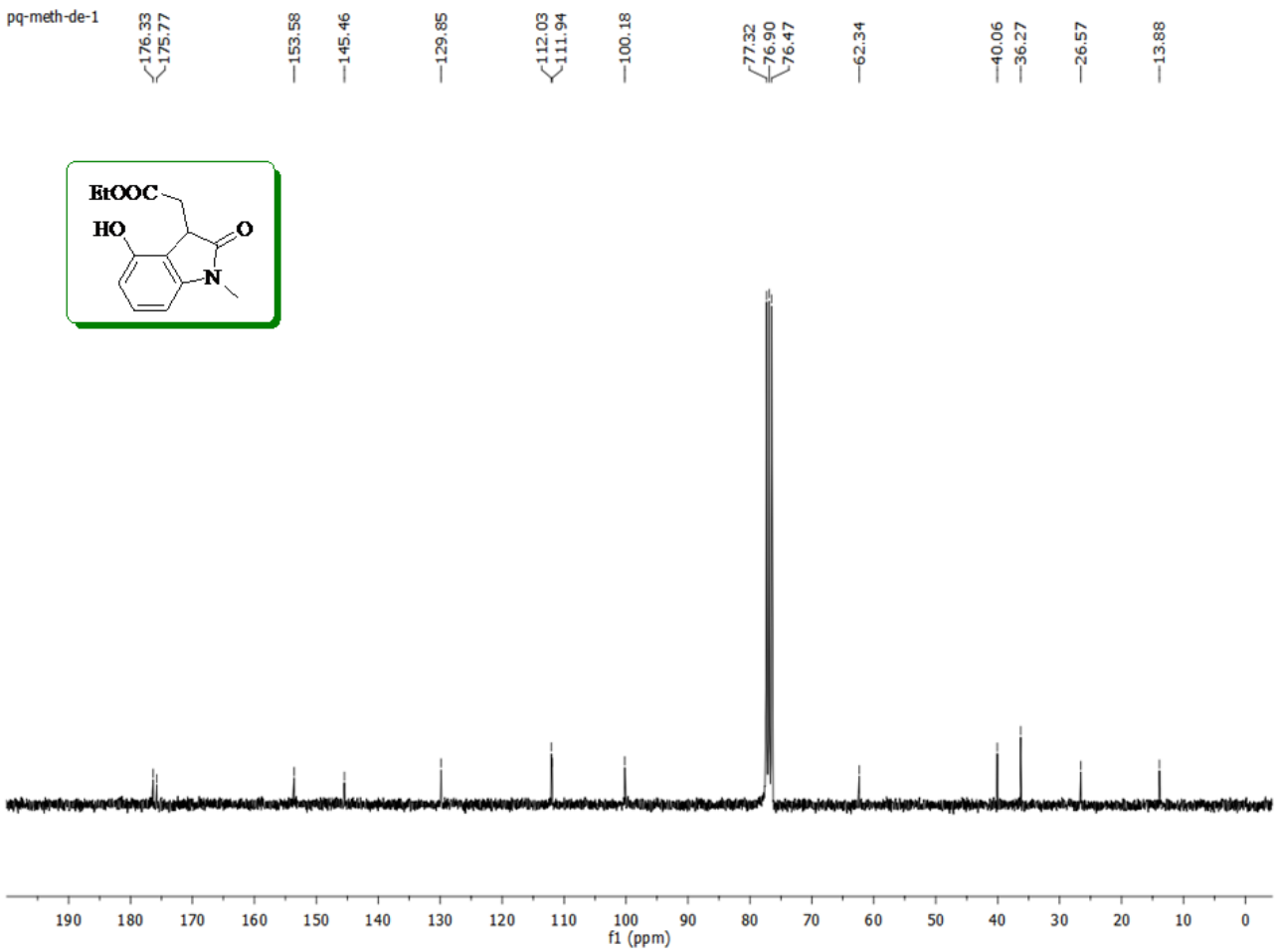

Figure S16. ^{13}C NMR spectrum of **4g**

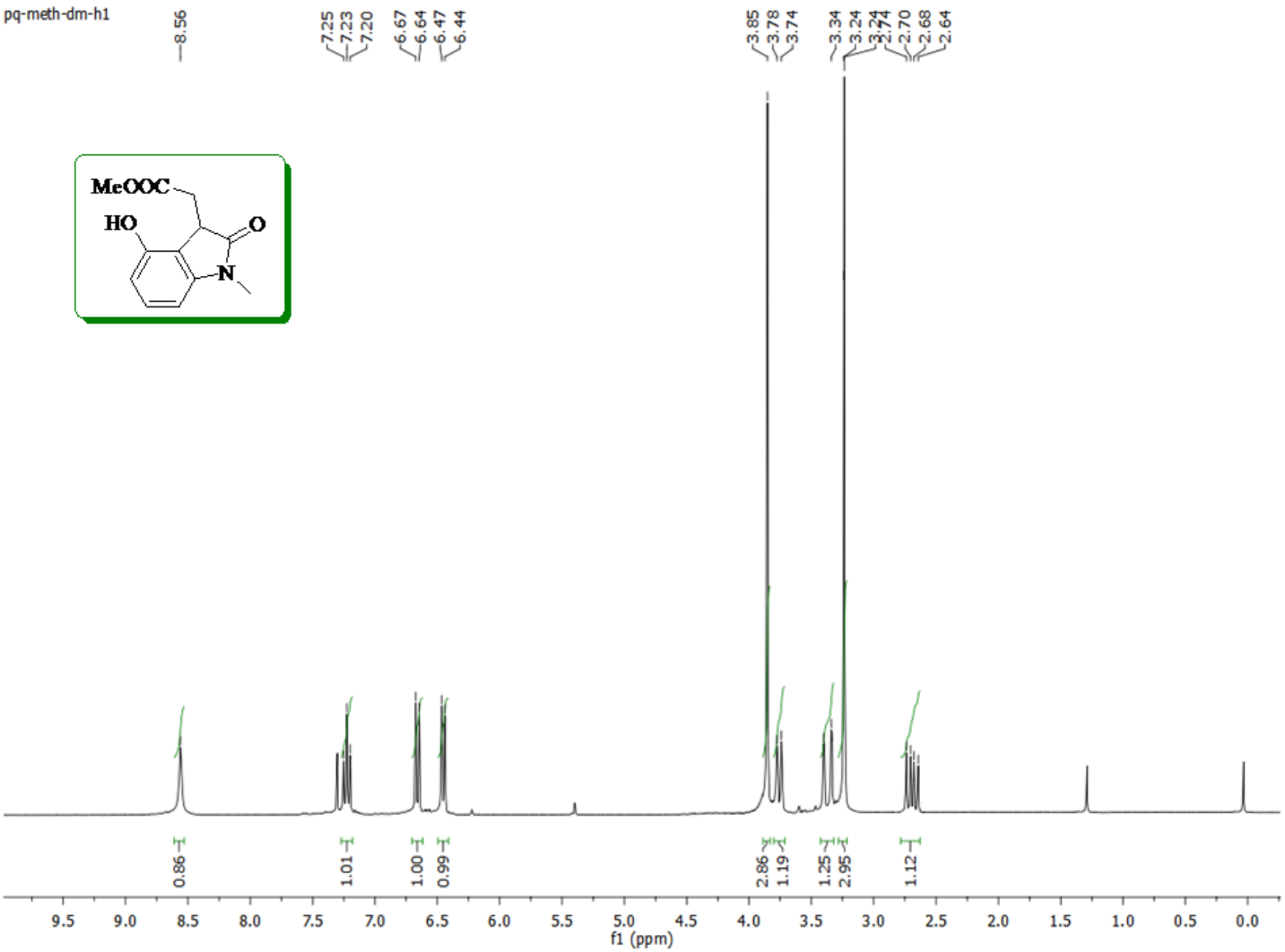

Figure S17. ^{1}H NMR spectrum of **4h**

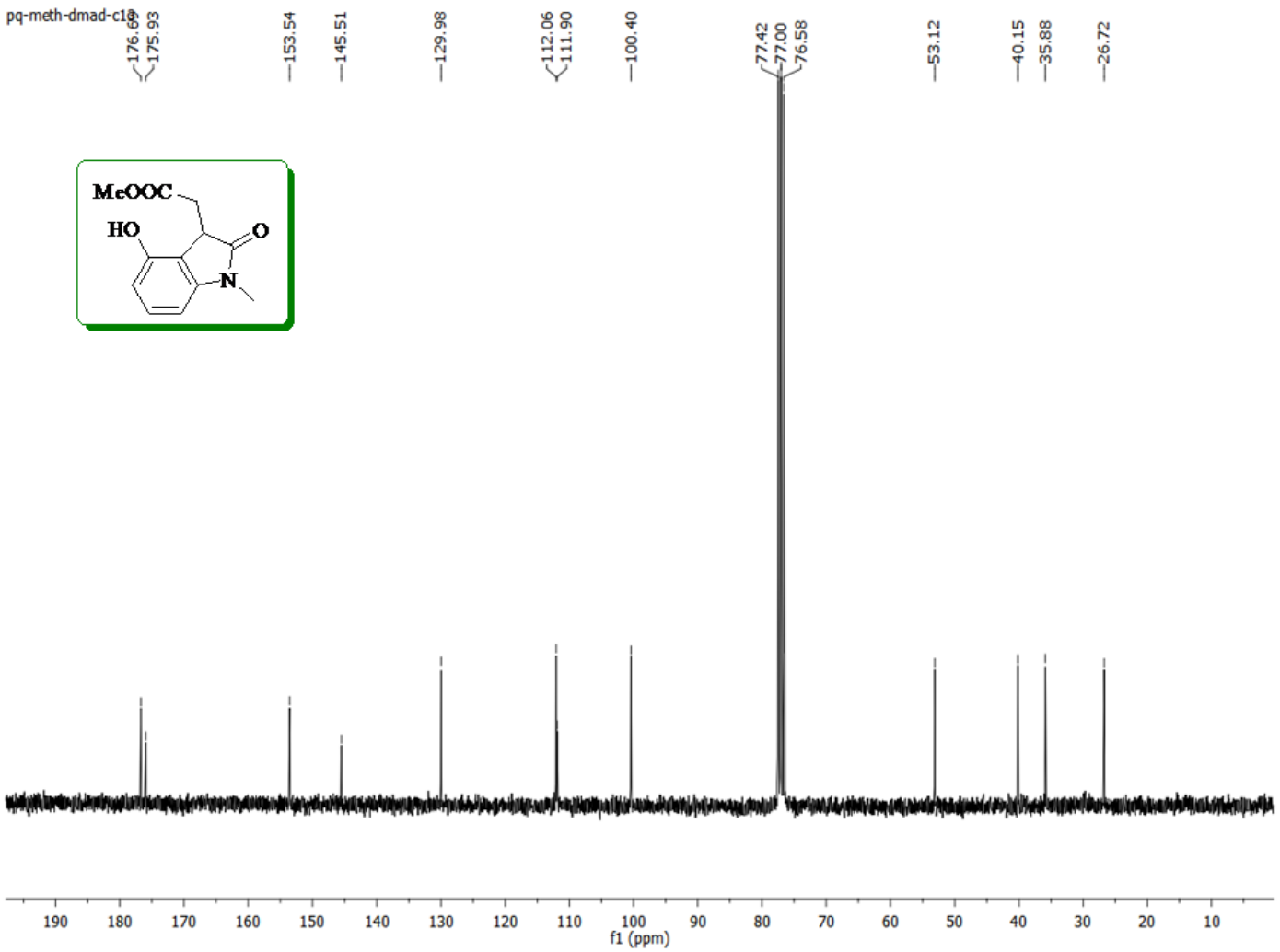

Figure S18. ^{13}C NMR spectrum of **4h**

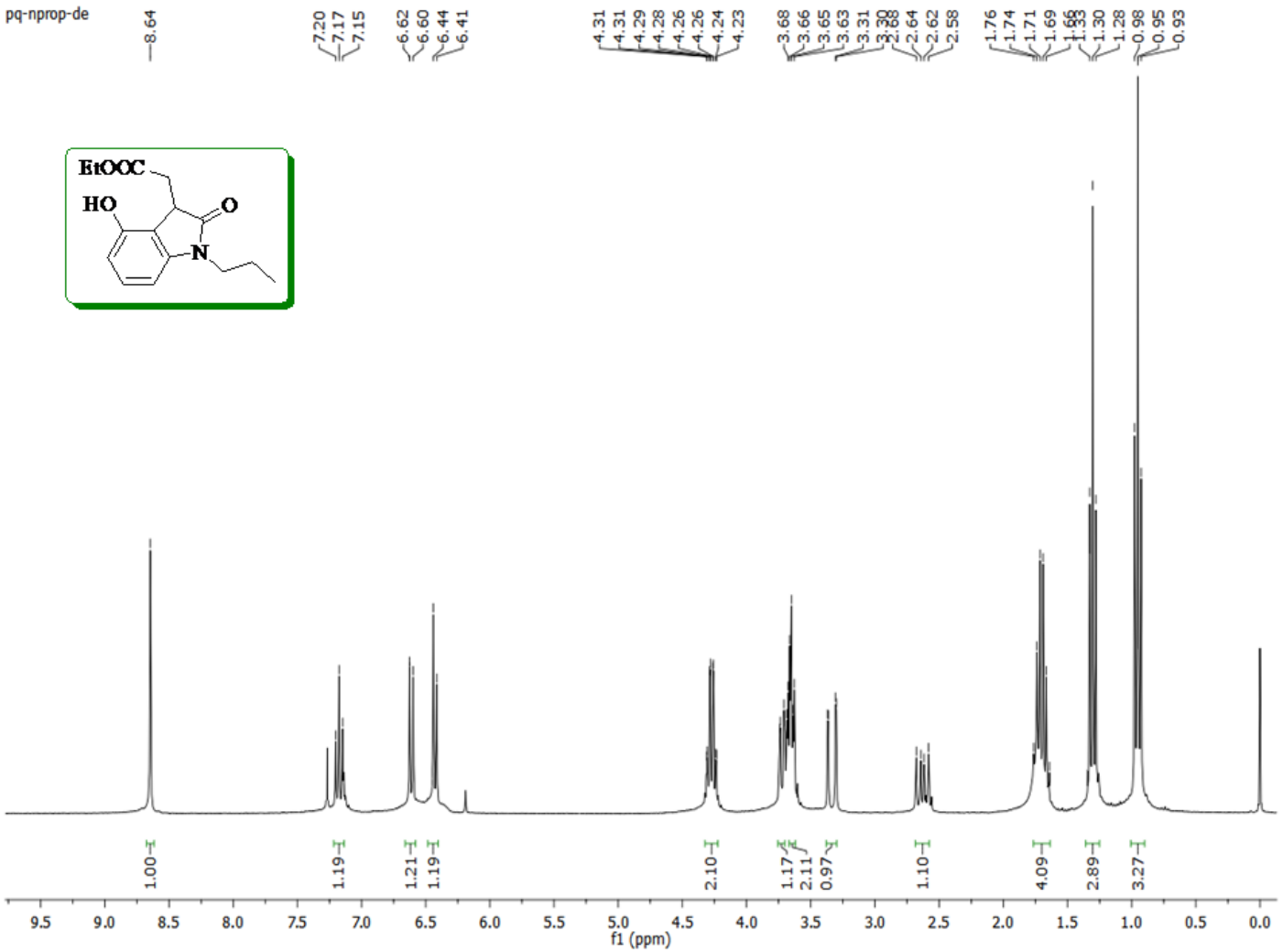

Figure S19. ^{1}H NMR spectrum of **4i**

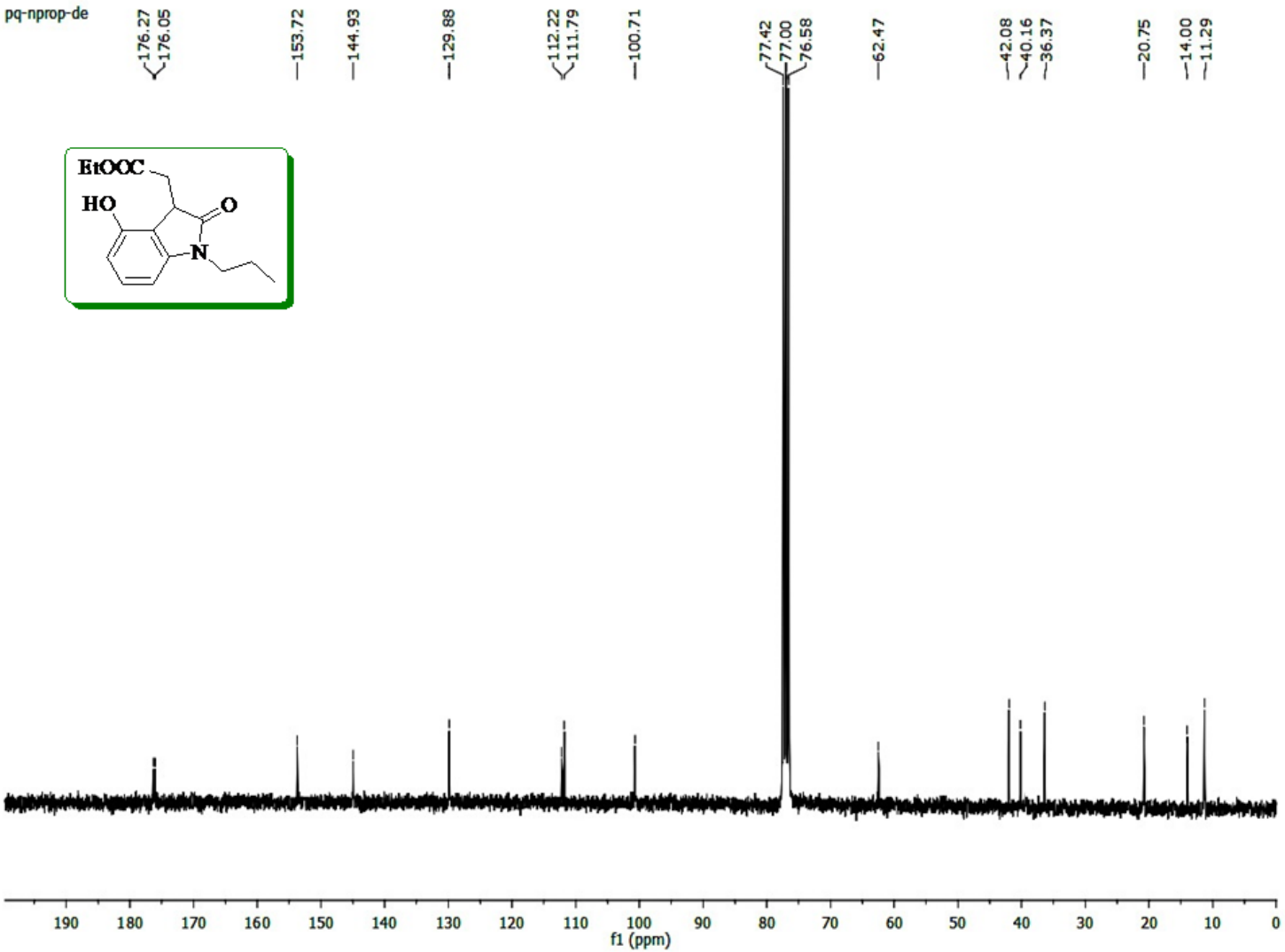

Figure S20. ^{13}C NMR spectrum of **4i**

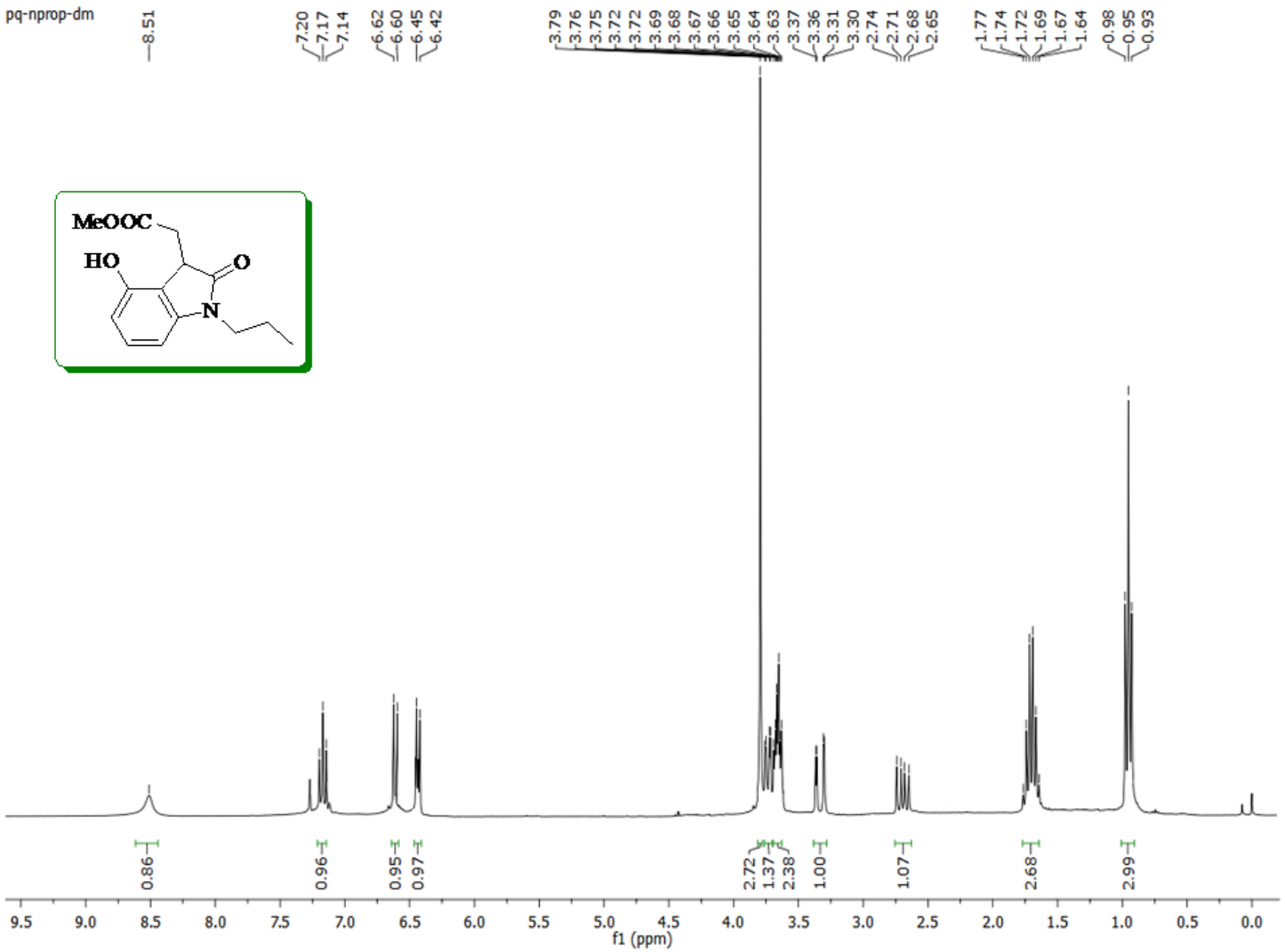

Figure S21. ^{1}H NMR spectrum of **4j**

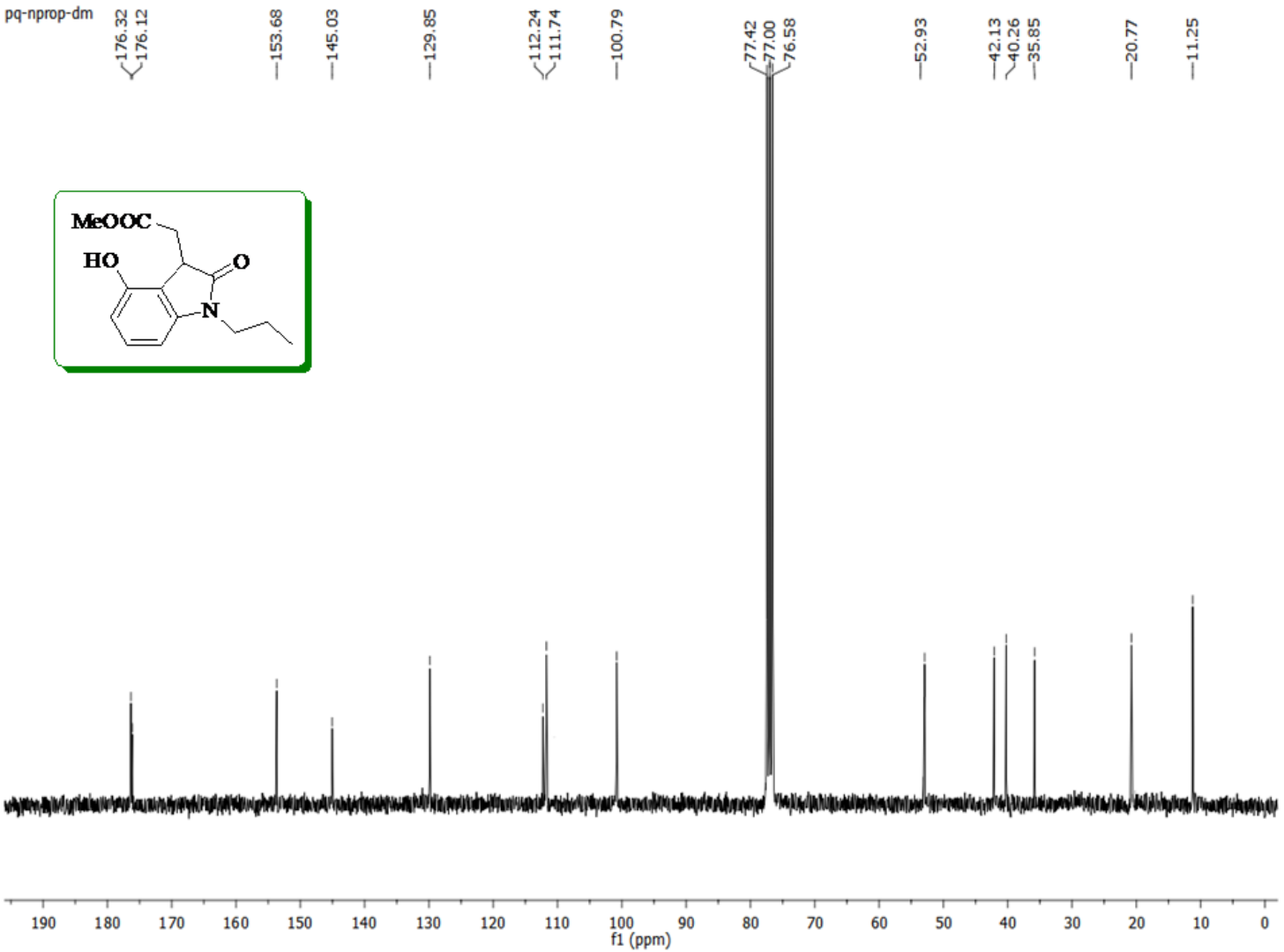

Figure S22. ^{13}C NMR spectrum of **4j**

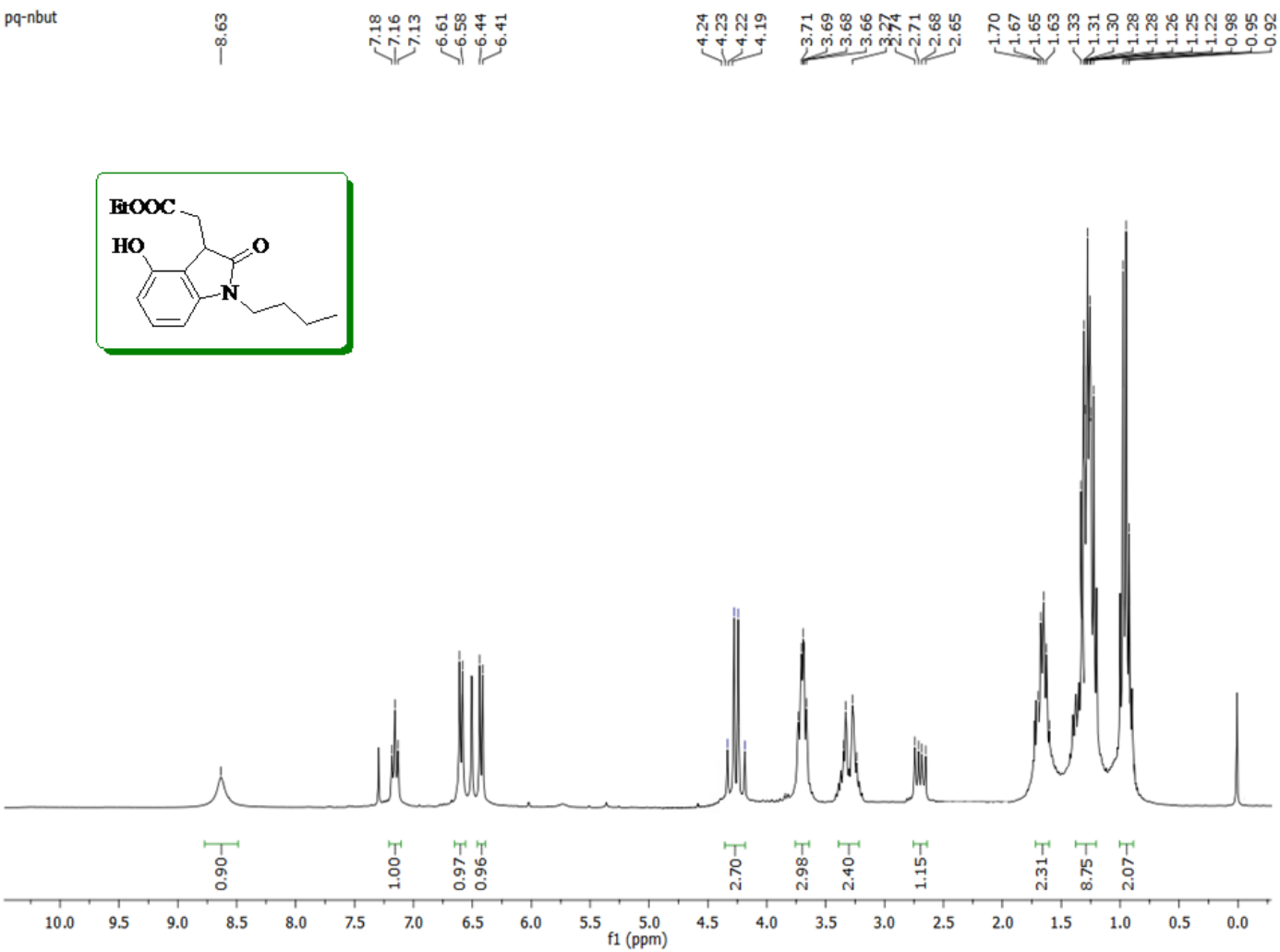

Figure S23. ^{1}H NMR spectrum of **4k**

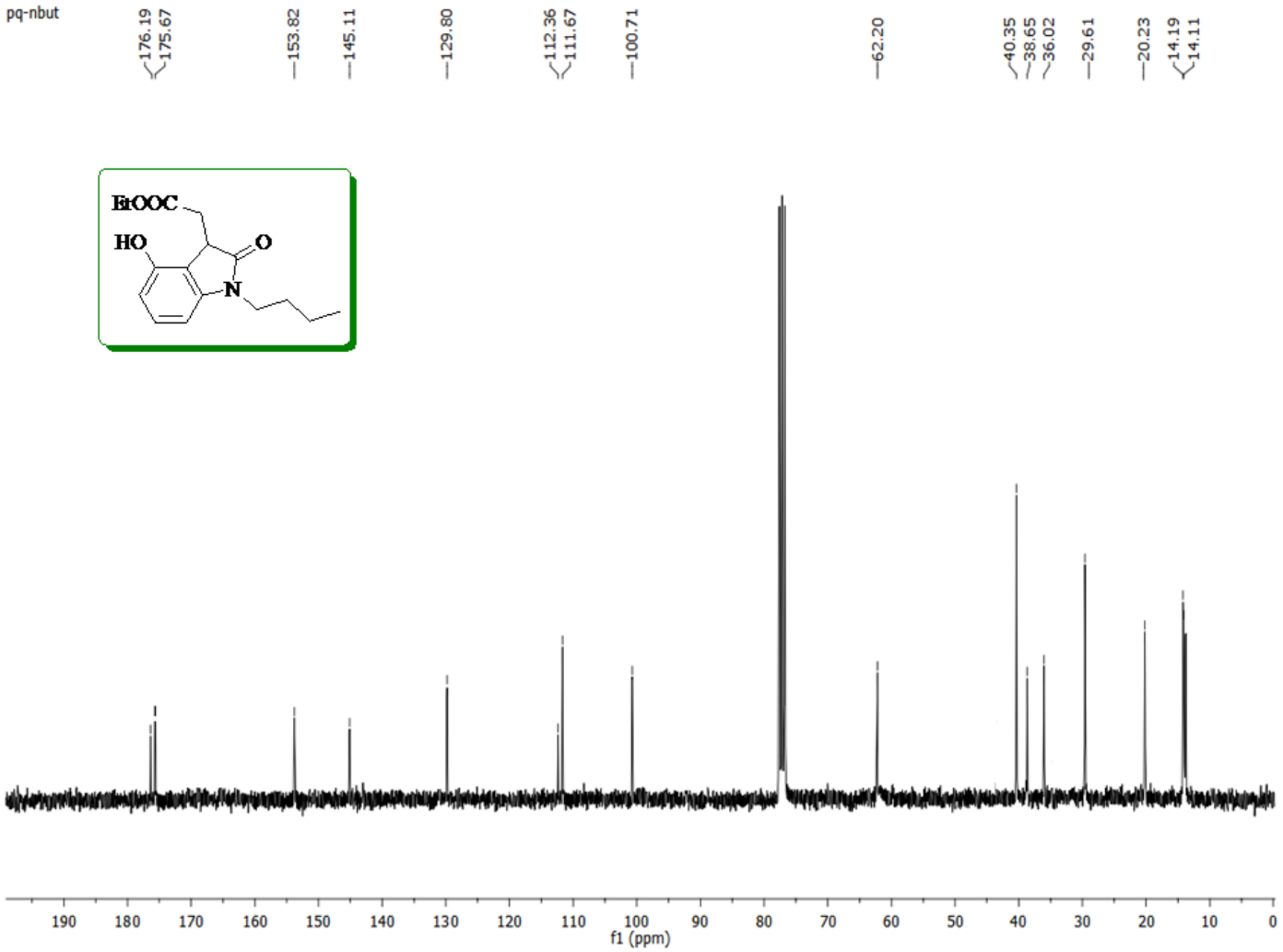

Figure S24. ^{13}C NMR spectrum of **4k**

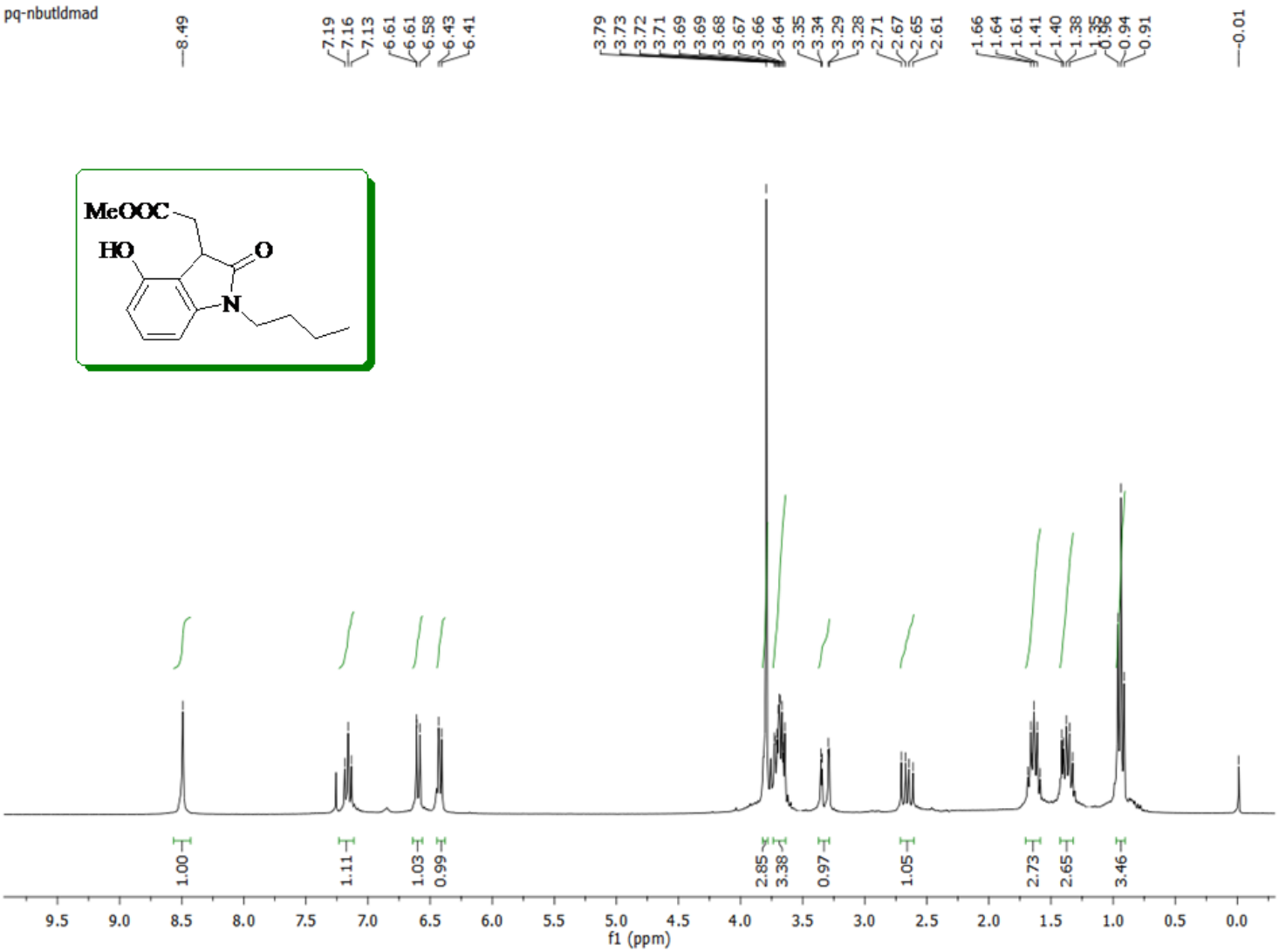

Figure S25. ^{1}H NMR spectrum of **4l**

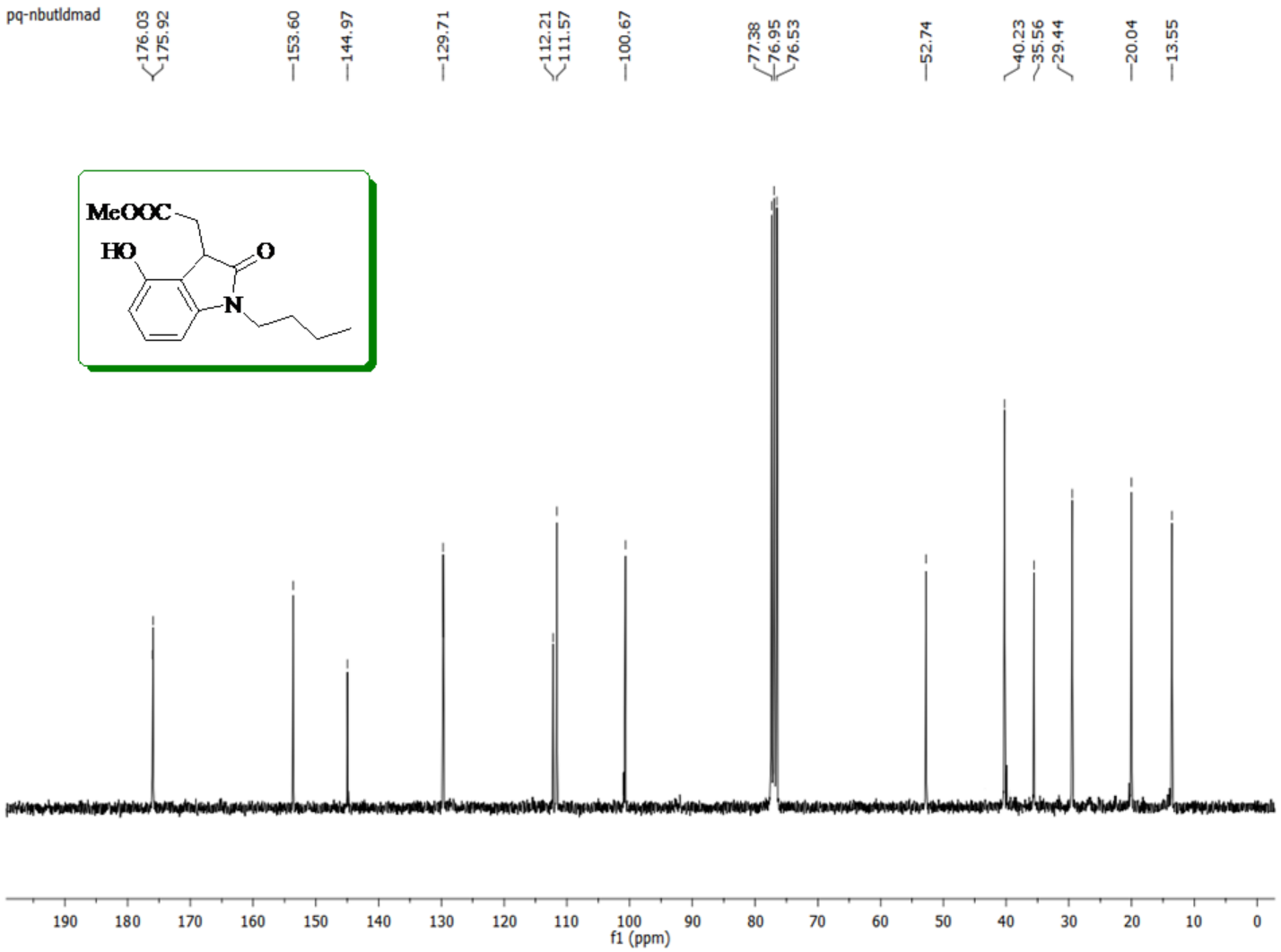

Figure S26. ^{13}C NMR spectrum of **4l**

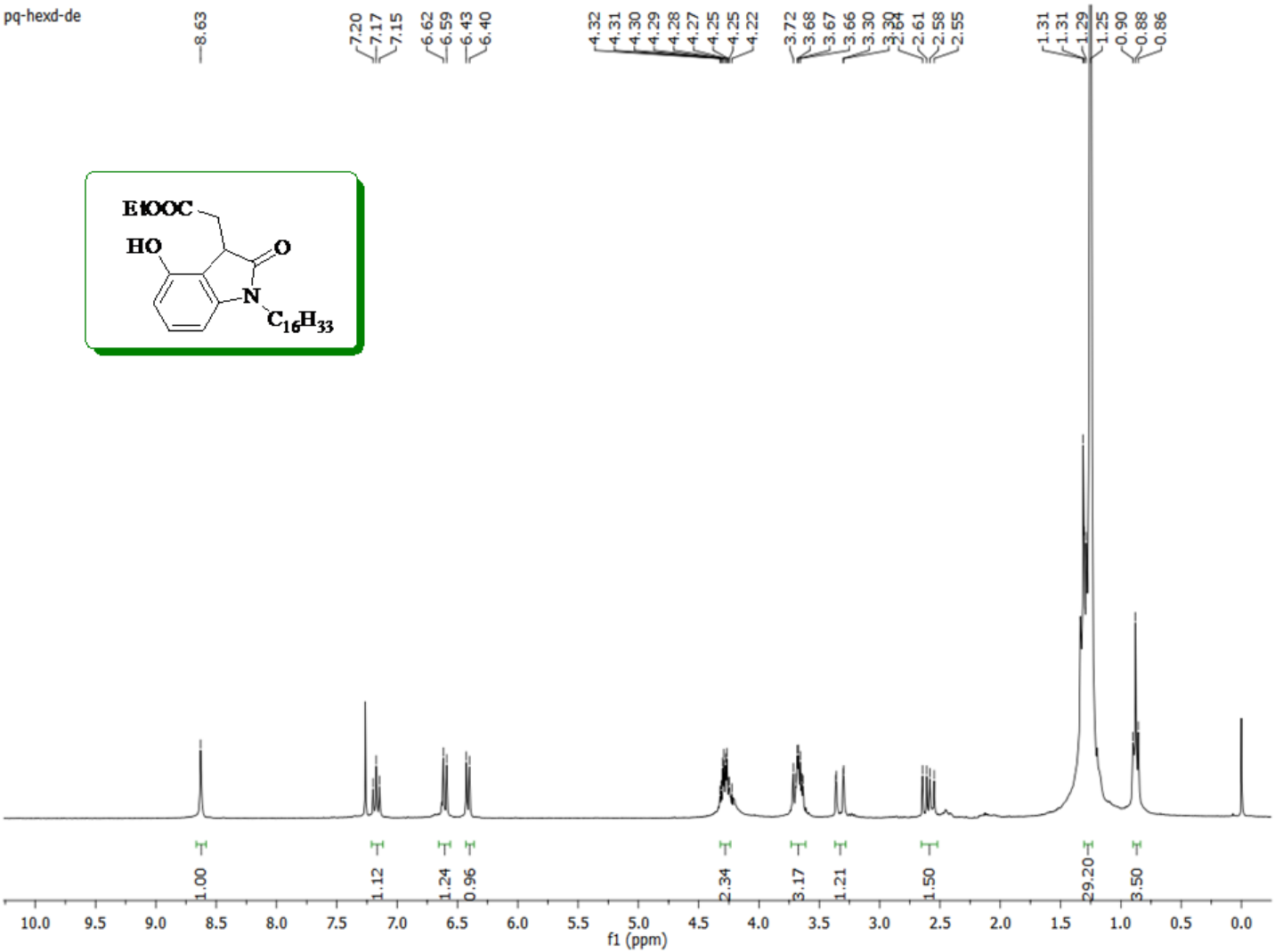

Figure S27. ^{1}H NMR spectrum of **4m**

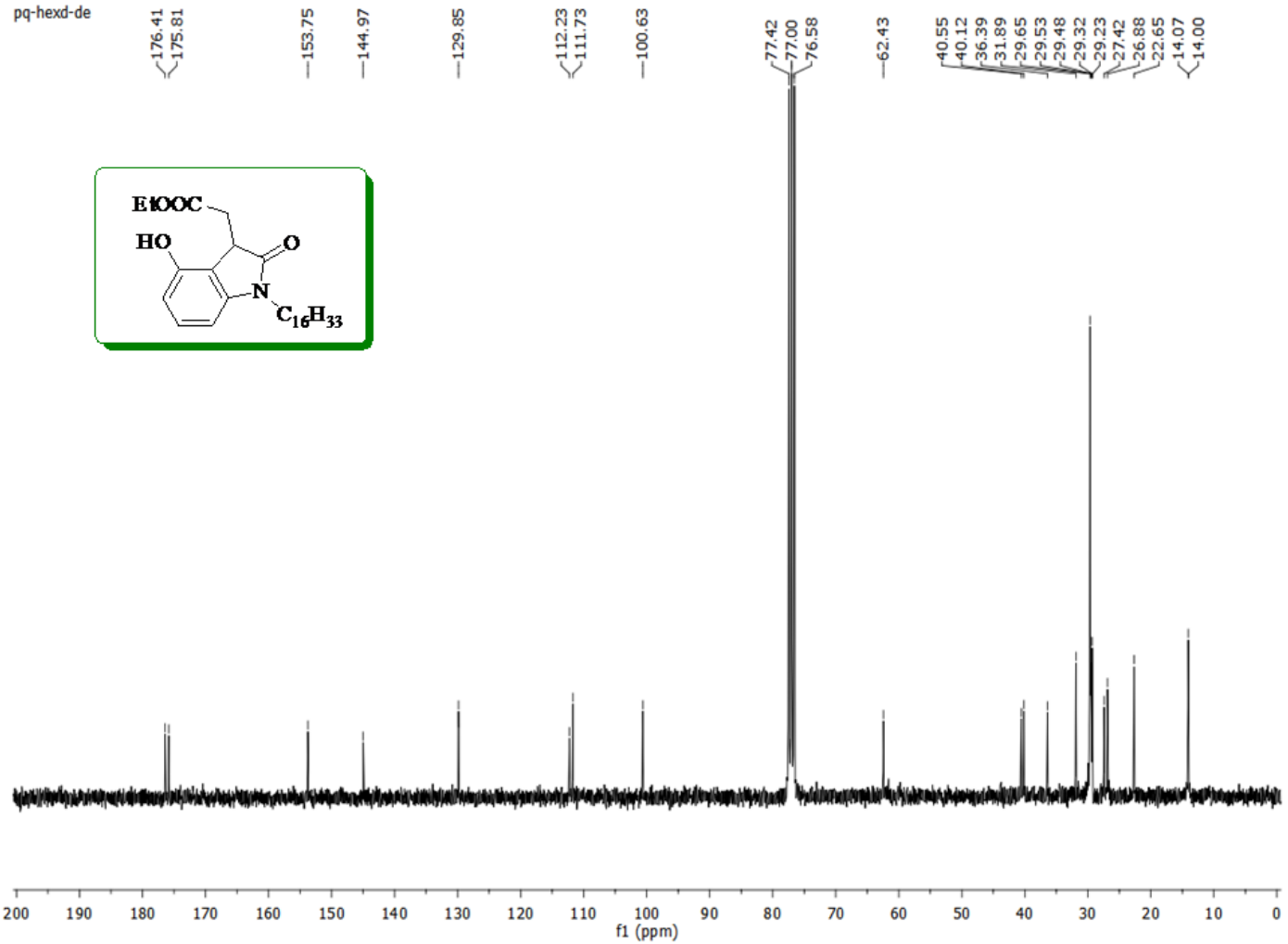

Figure S28. ^{13}C NMR spectrum of **4m**

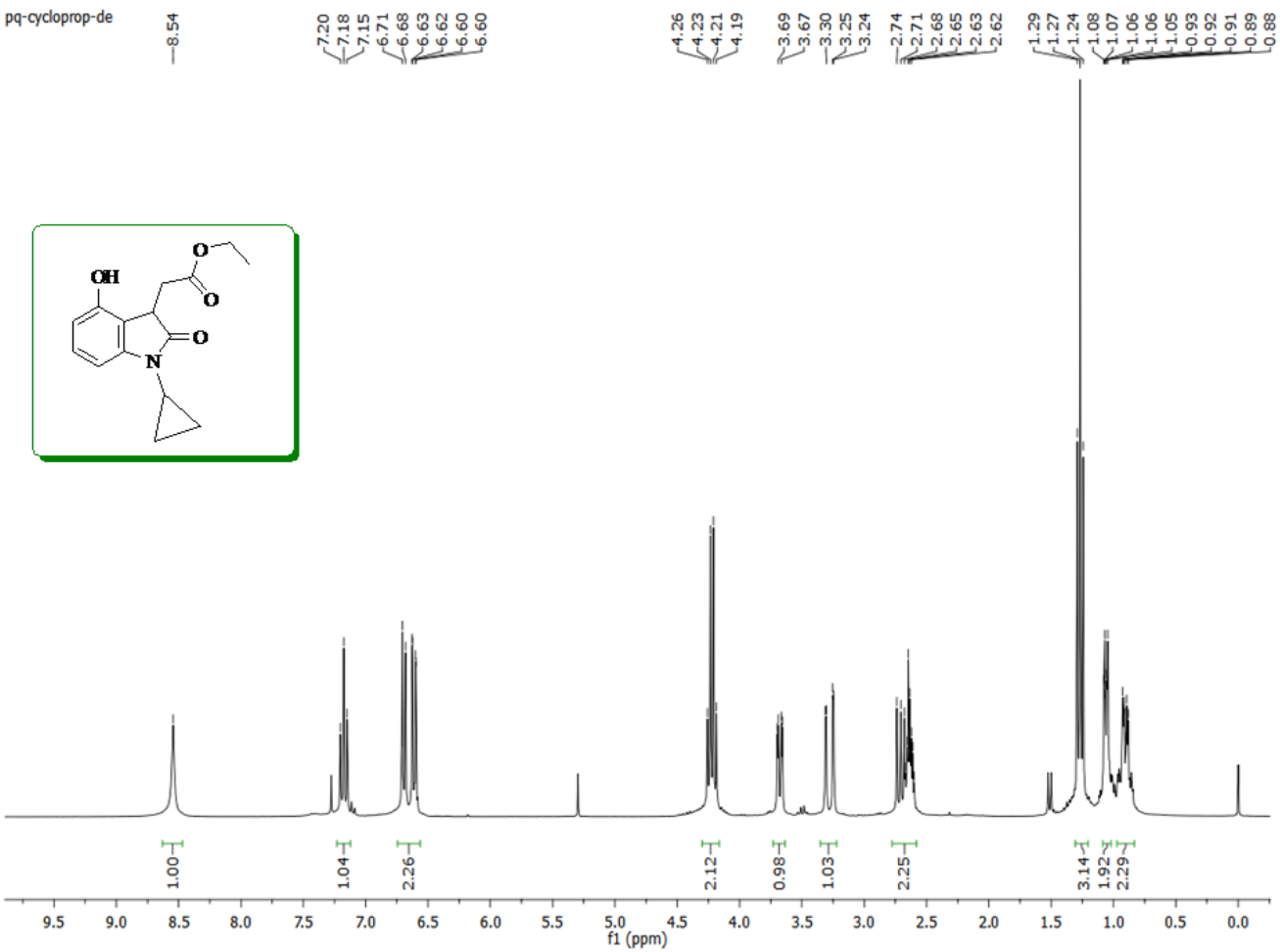

Figure S29. ^{1}H NMR spectrum of **4n**

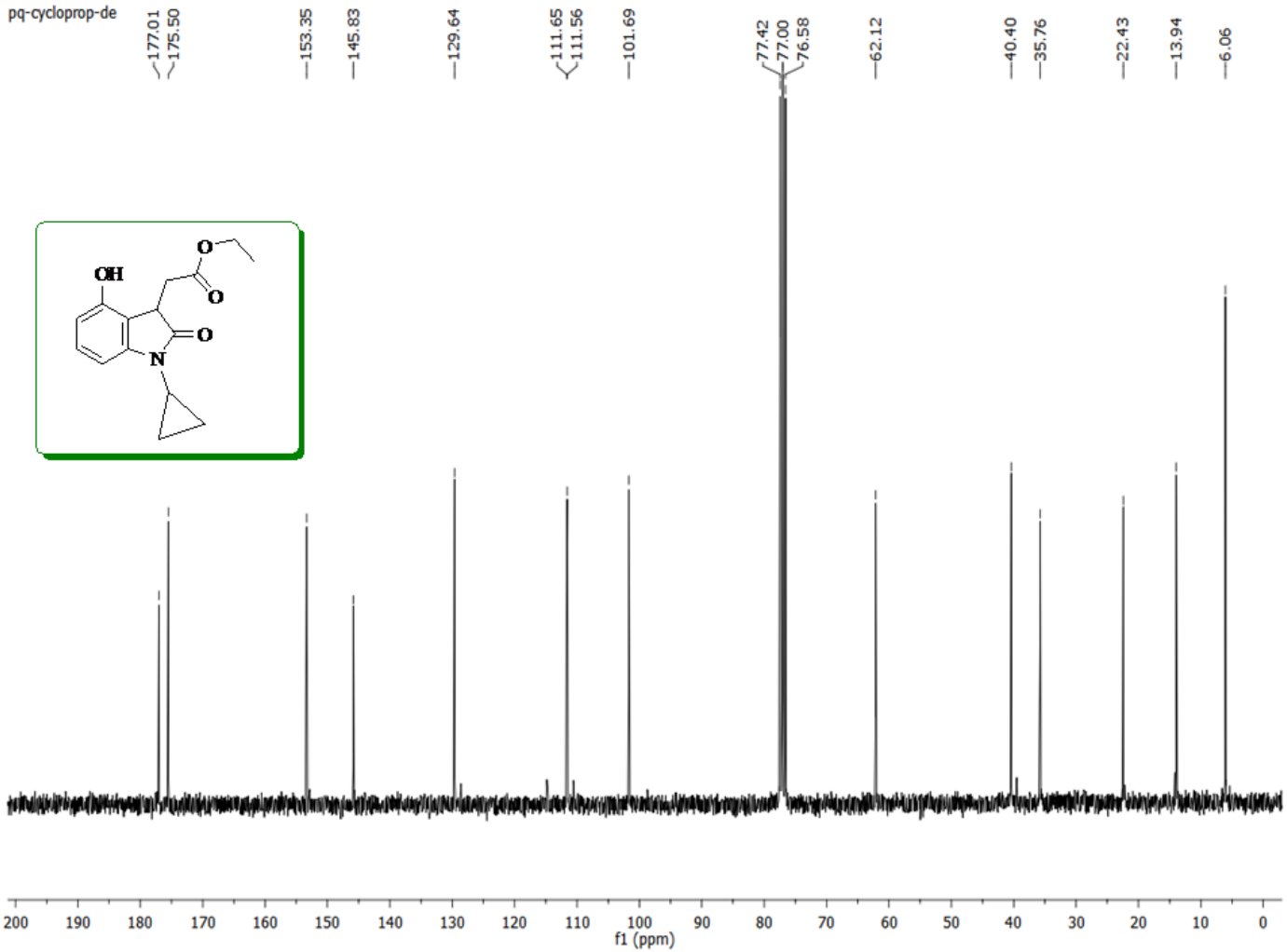

Figure S30. ^{13}C NMR spectrum of **4n**

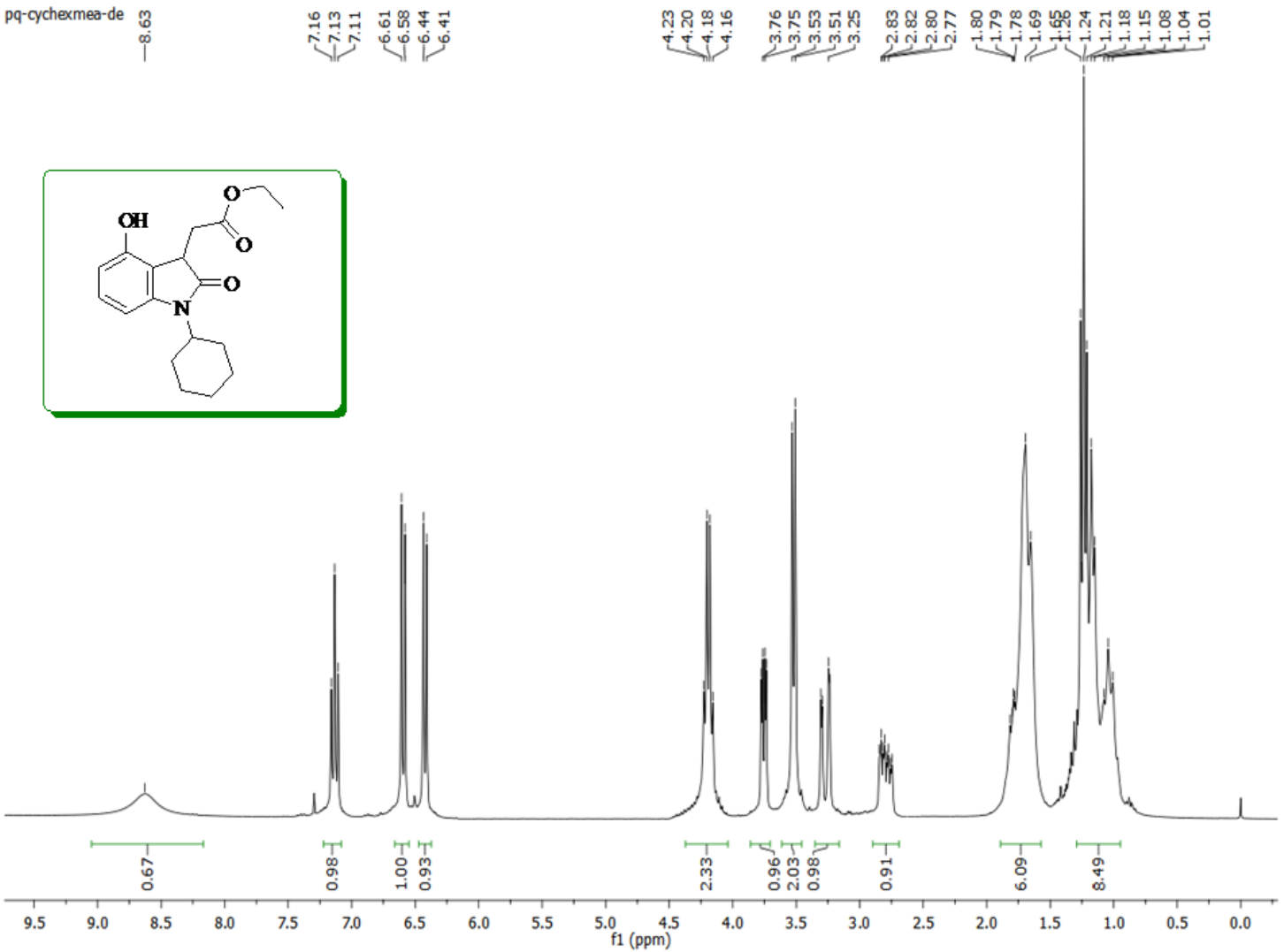

Figure S31. ^{1}H NMR spectrum of **4o**

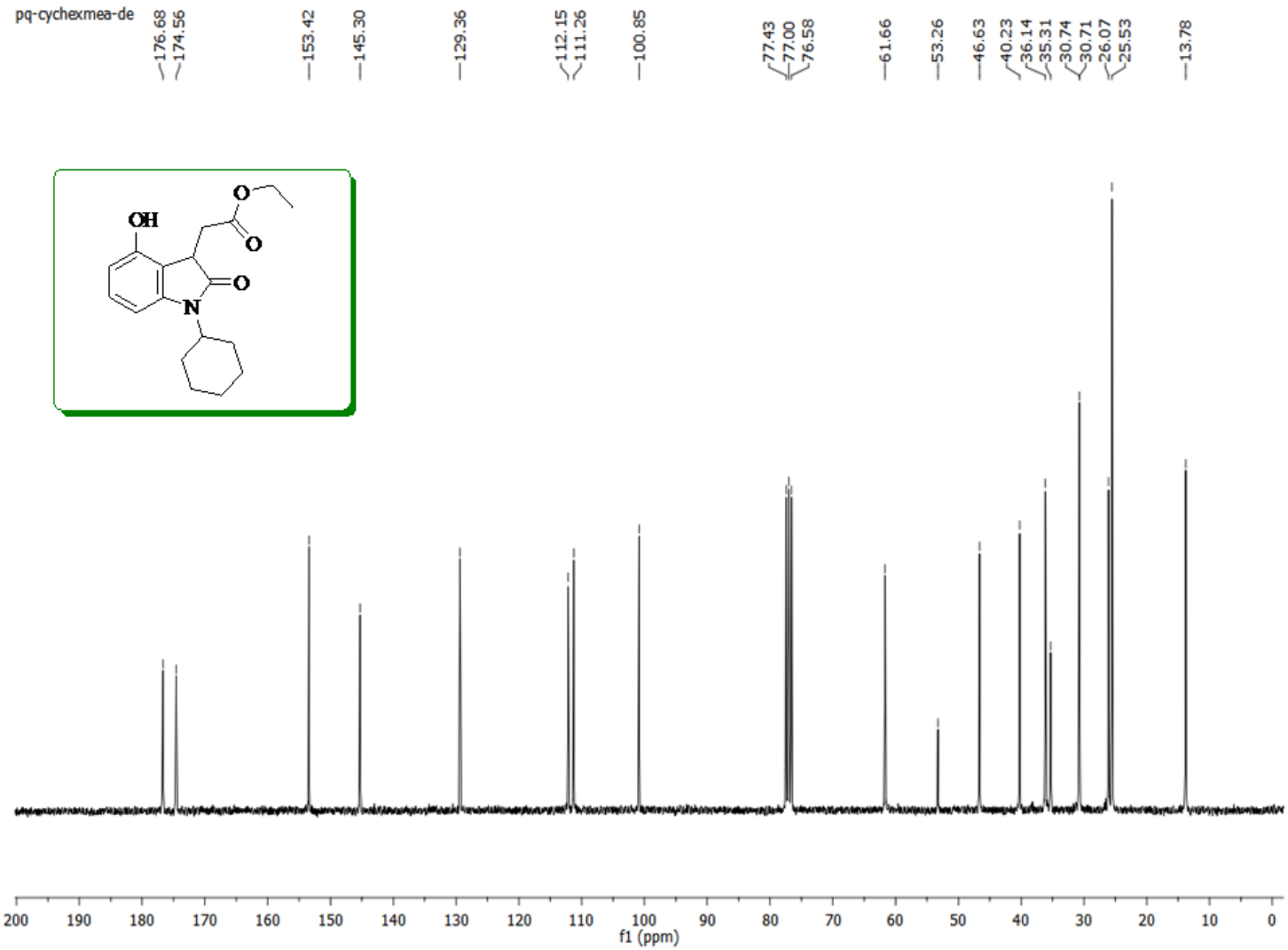

Figure S32. ^{13}C NMR spectrum of **4o**

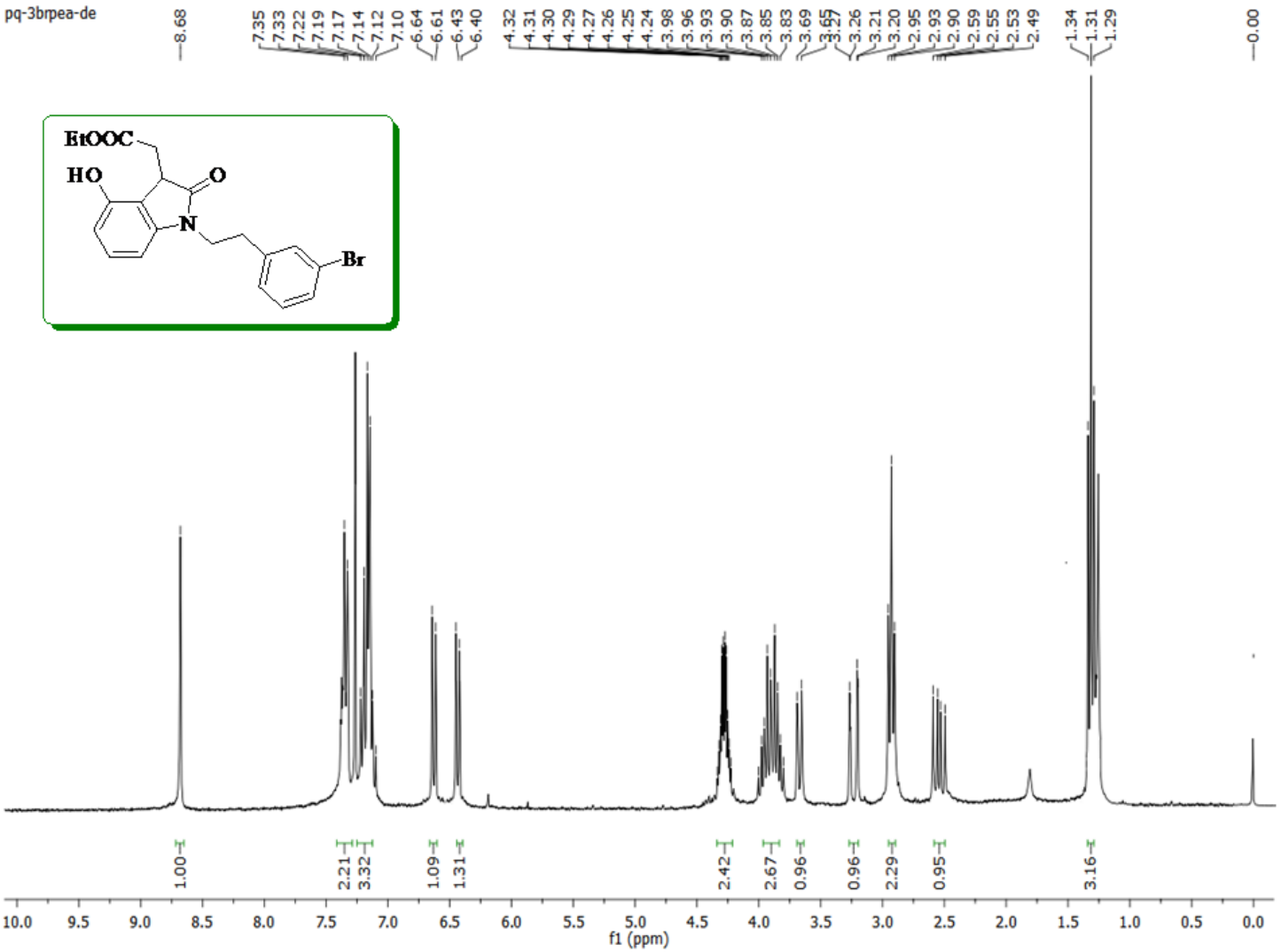

Figure S33. ^{1}H NMR spectrum of **4p**

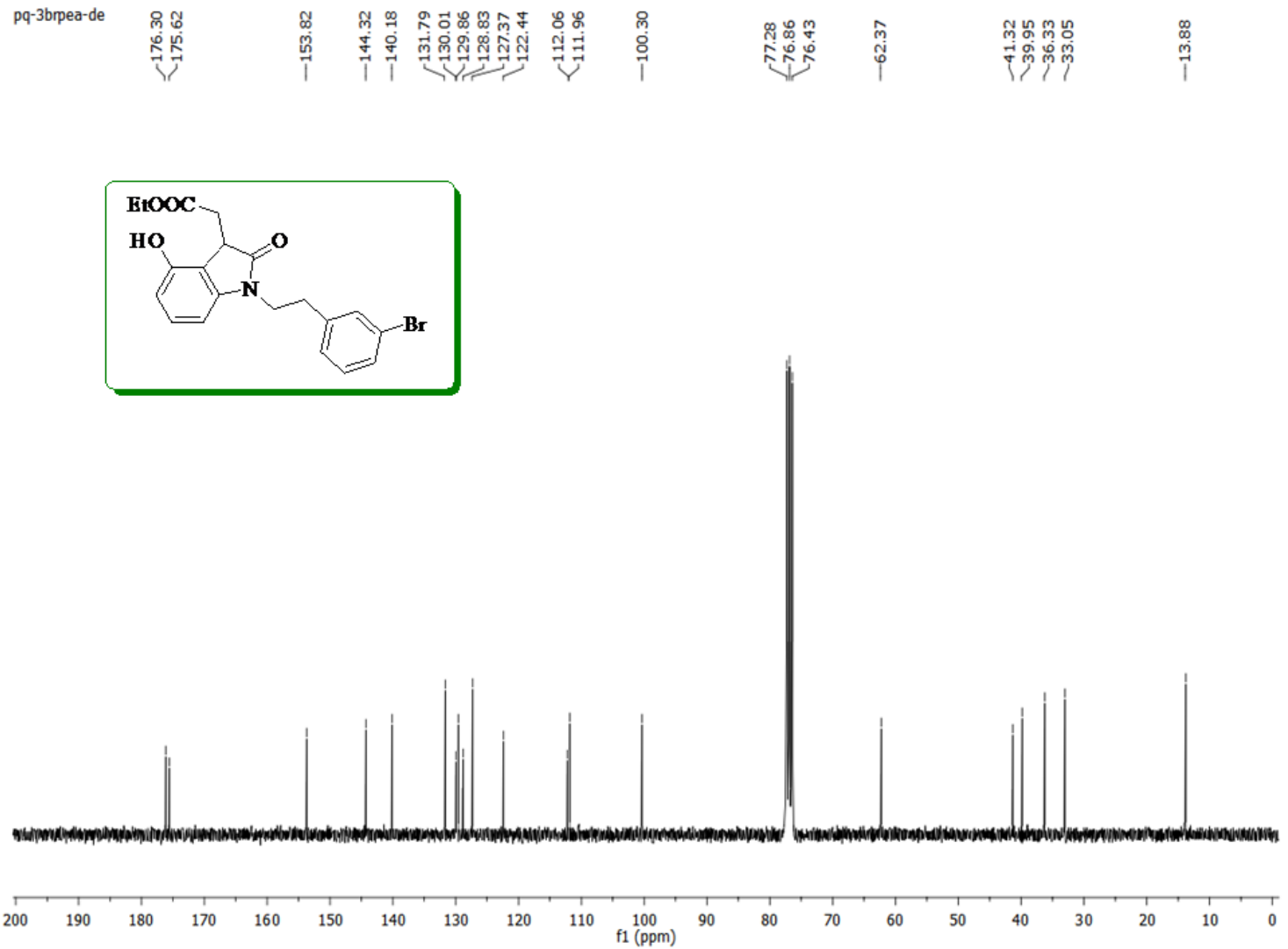

Figure S34. ^{13}C NMR spectrum of **4p**

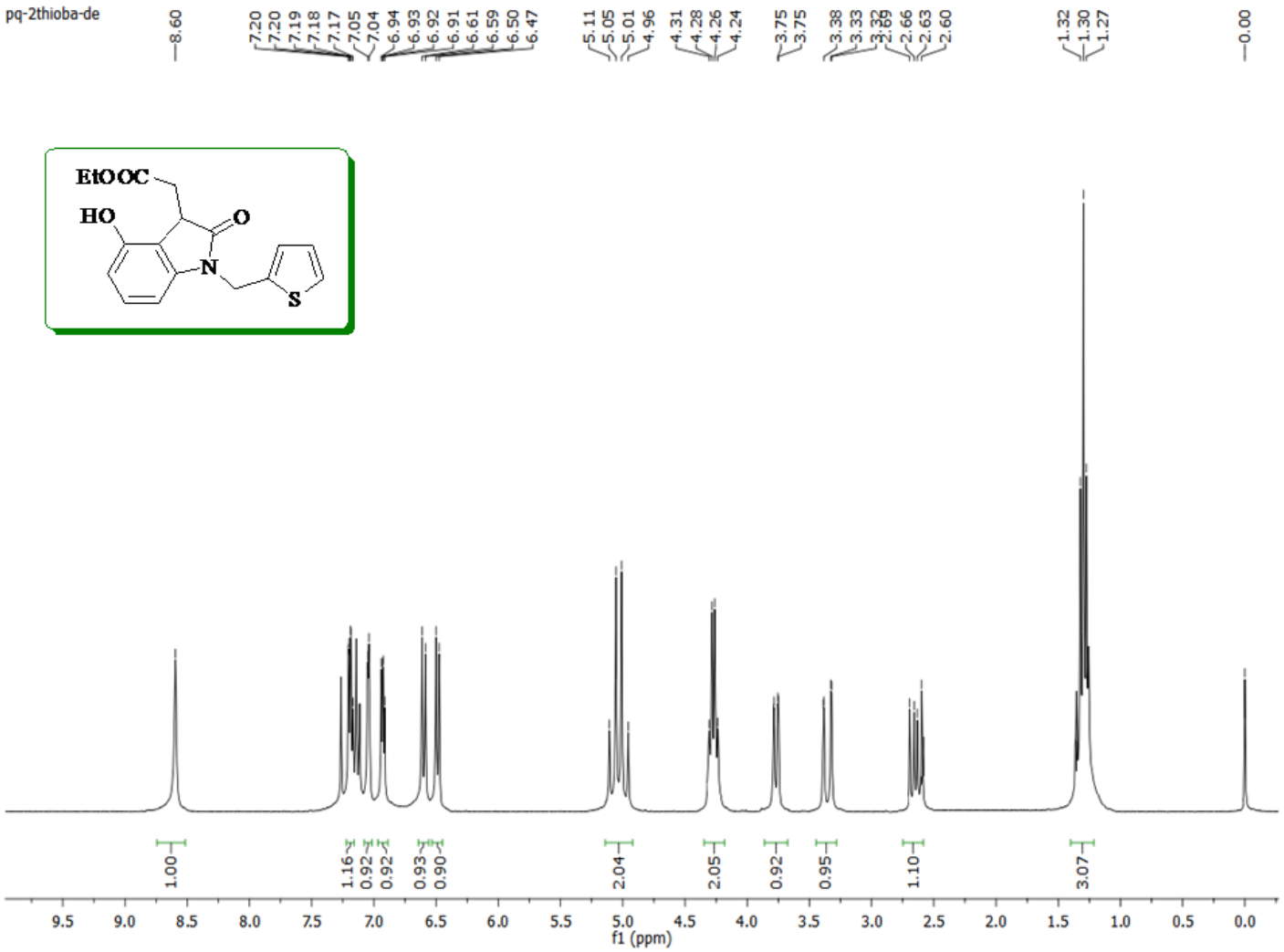

Figure S35. ^{1}H NMR spectrum of **4q**

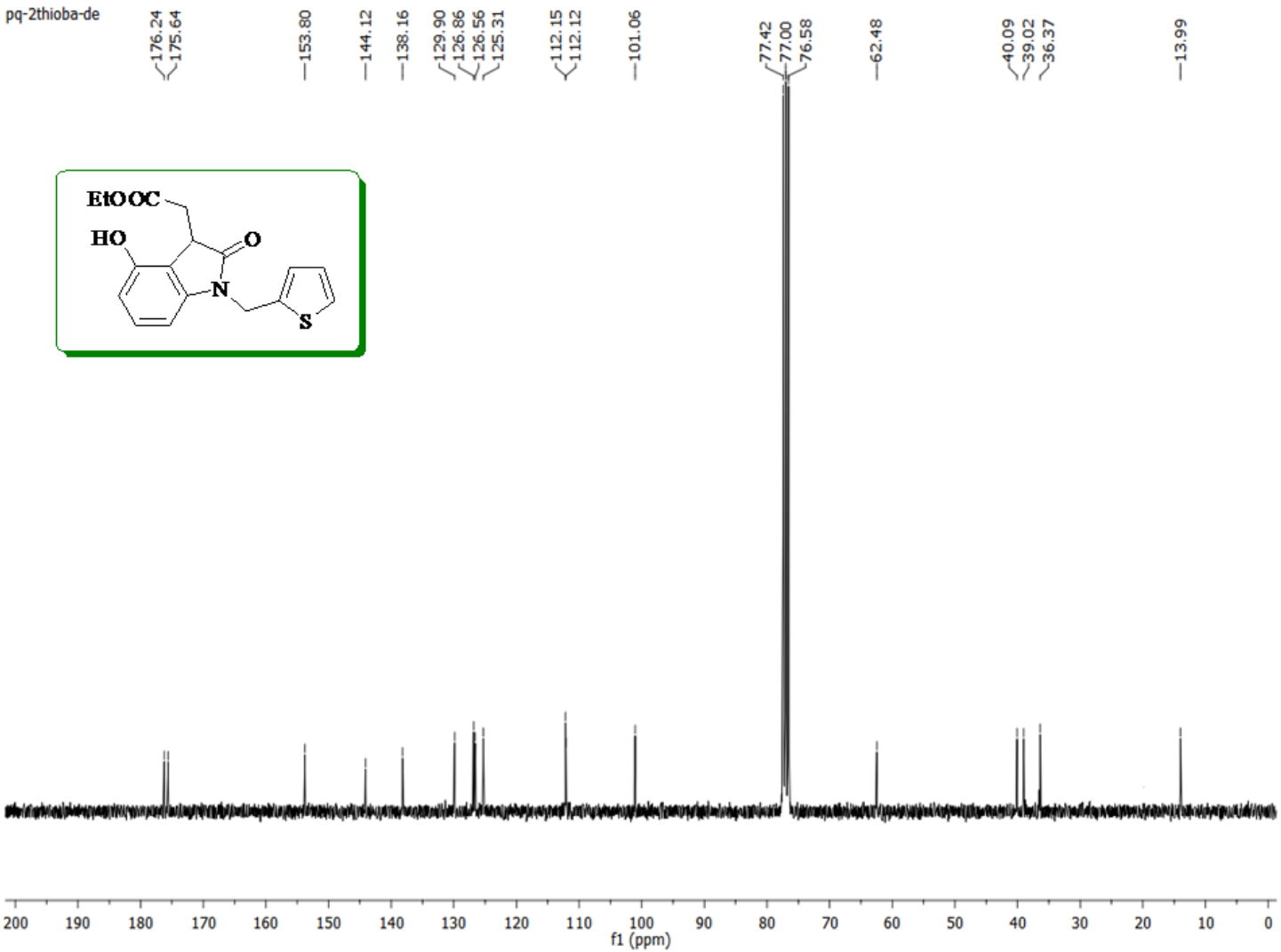

Figure S36. ^{13}C NMR spectrum of **4q**

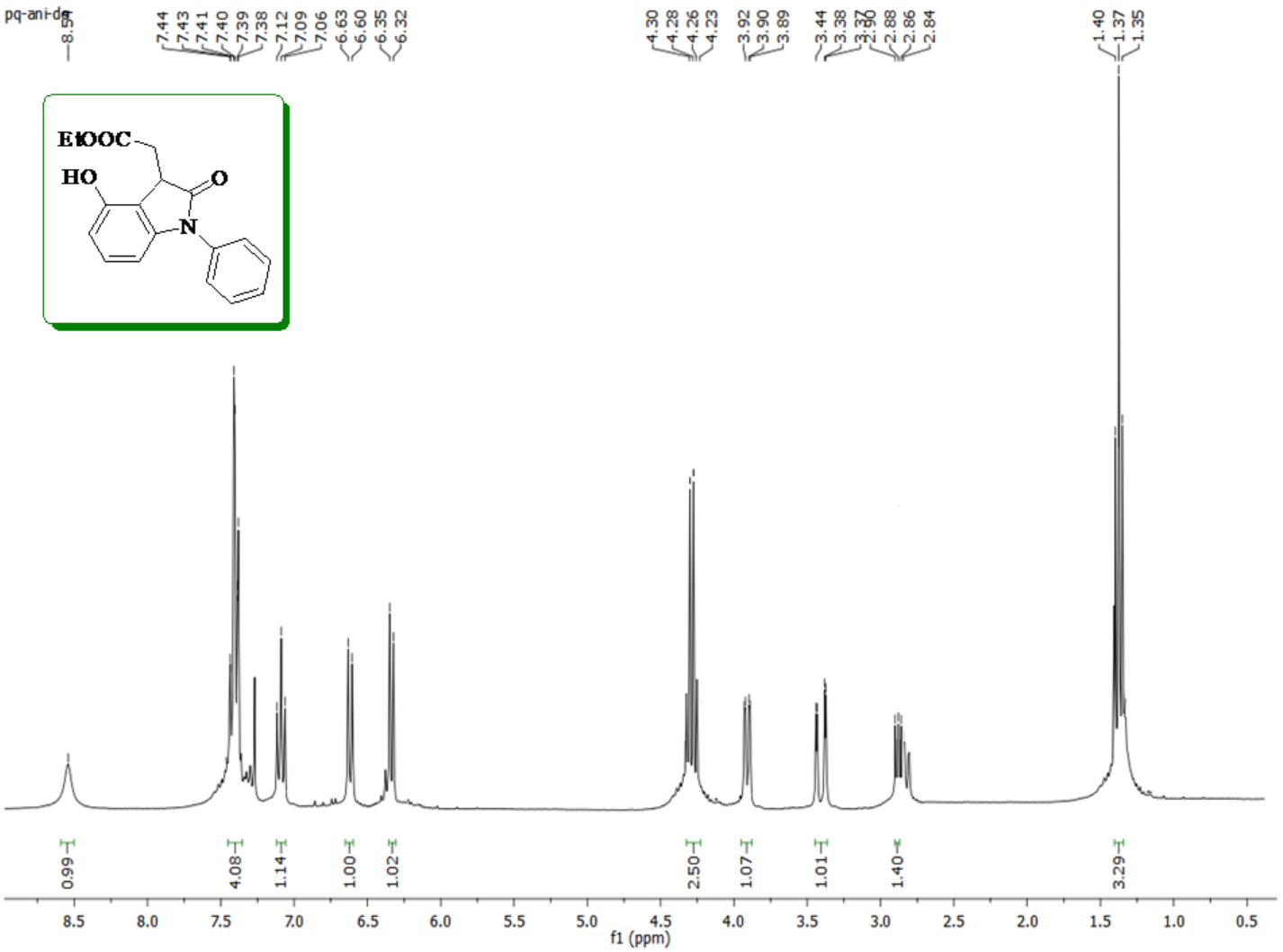

Figure S37. ^{1}H NMR spectrum of **4r**

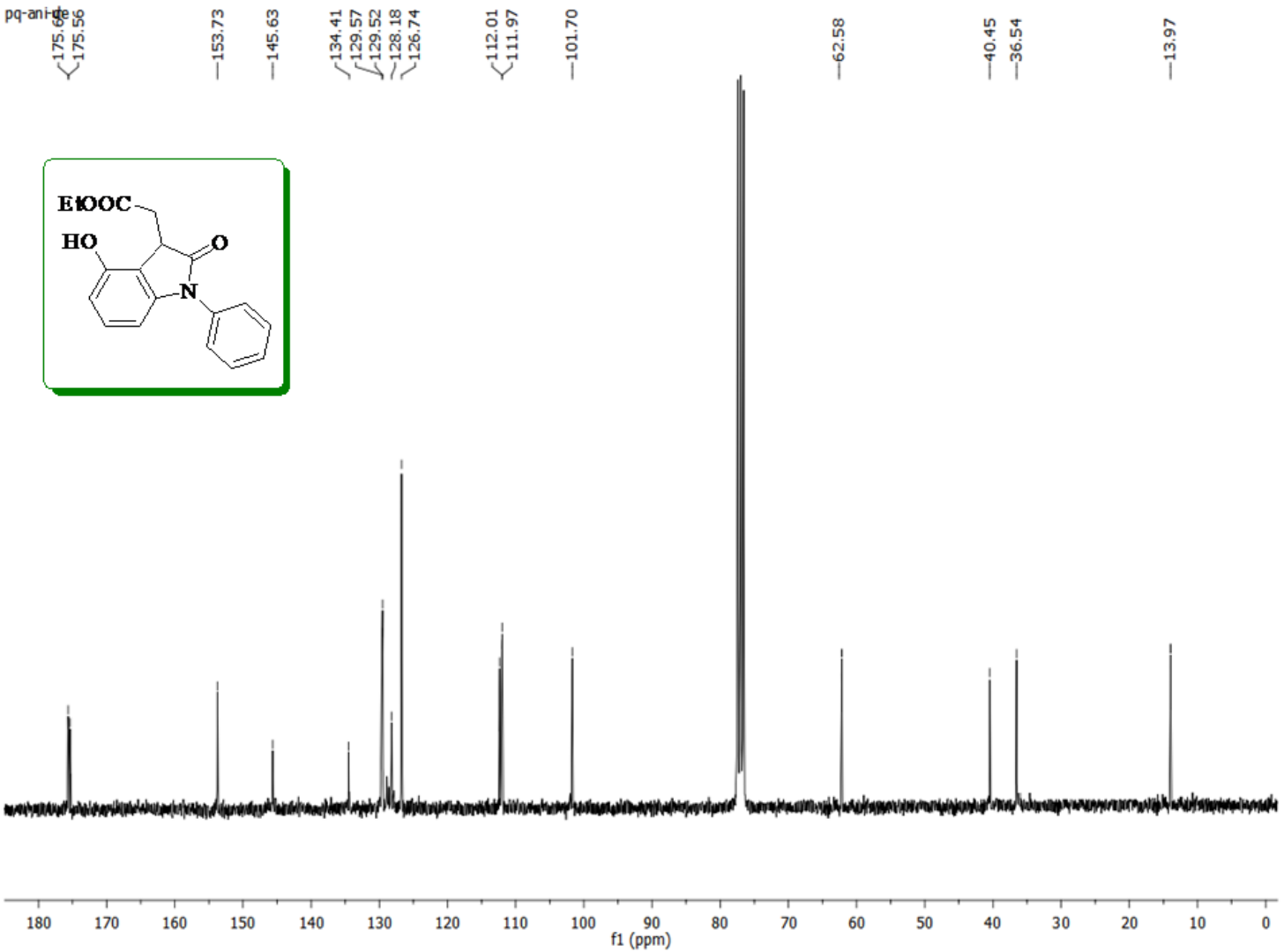

Figure S38. ^{13}C NMR spectrum of **4r**

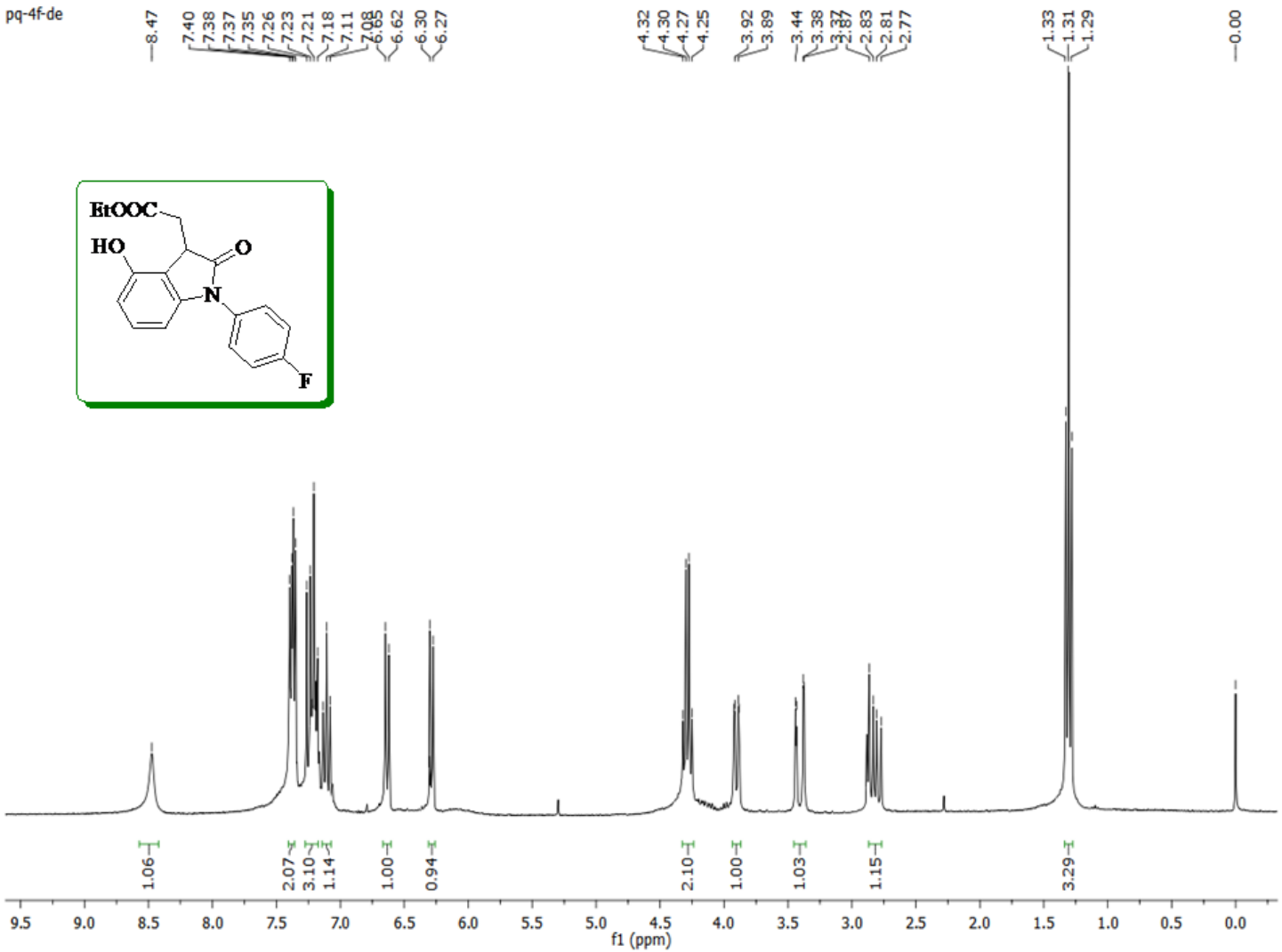

Figure S39. ^{1}H NMR spectrum of **4s**

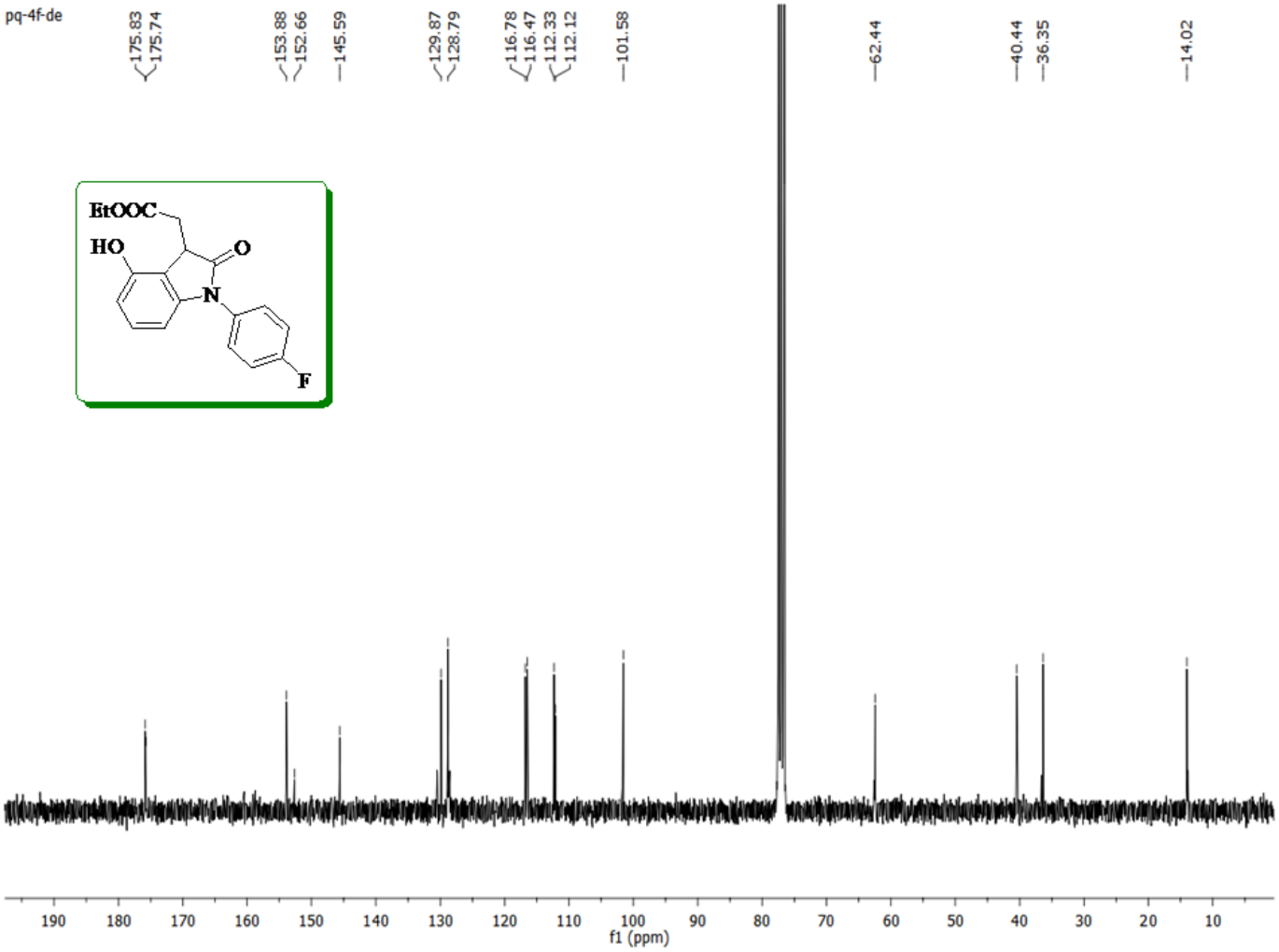

Figure S40. ^{13}C NMR spectrum of **4s**

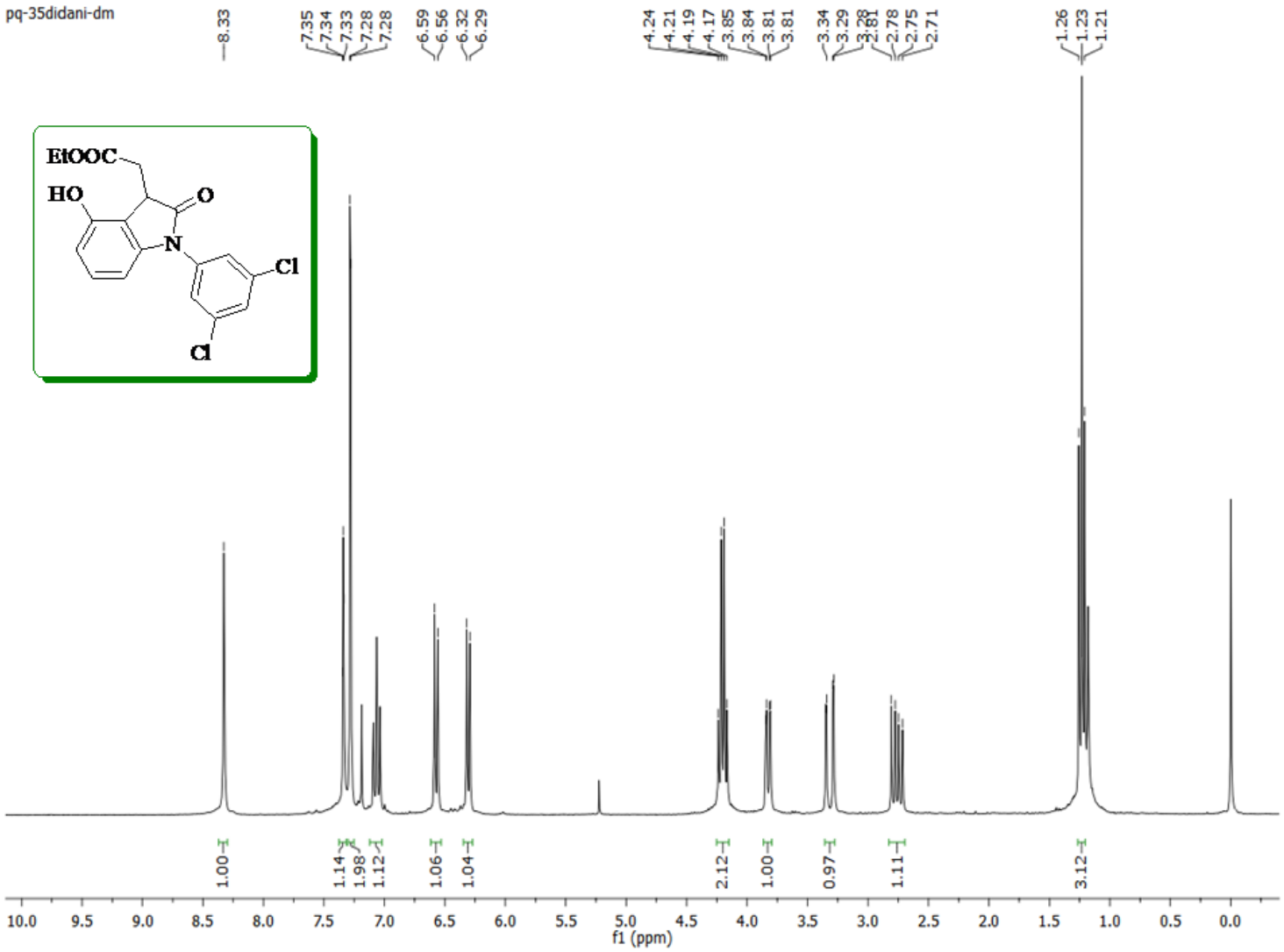

Figure S41. ^{1}H NMR spectrum of **4t**

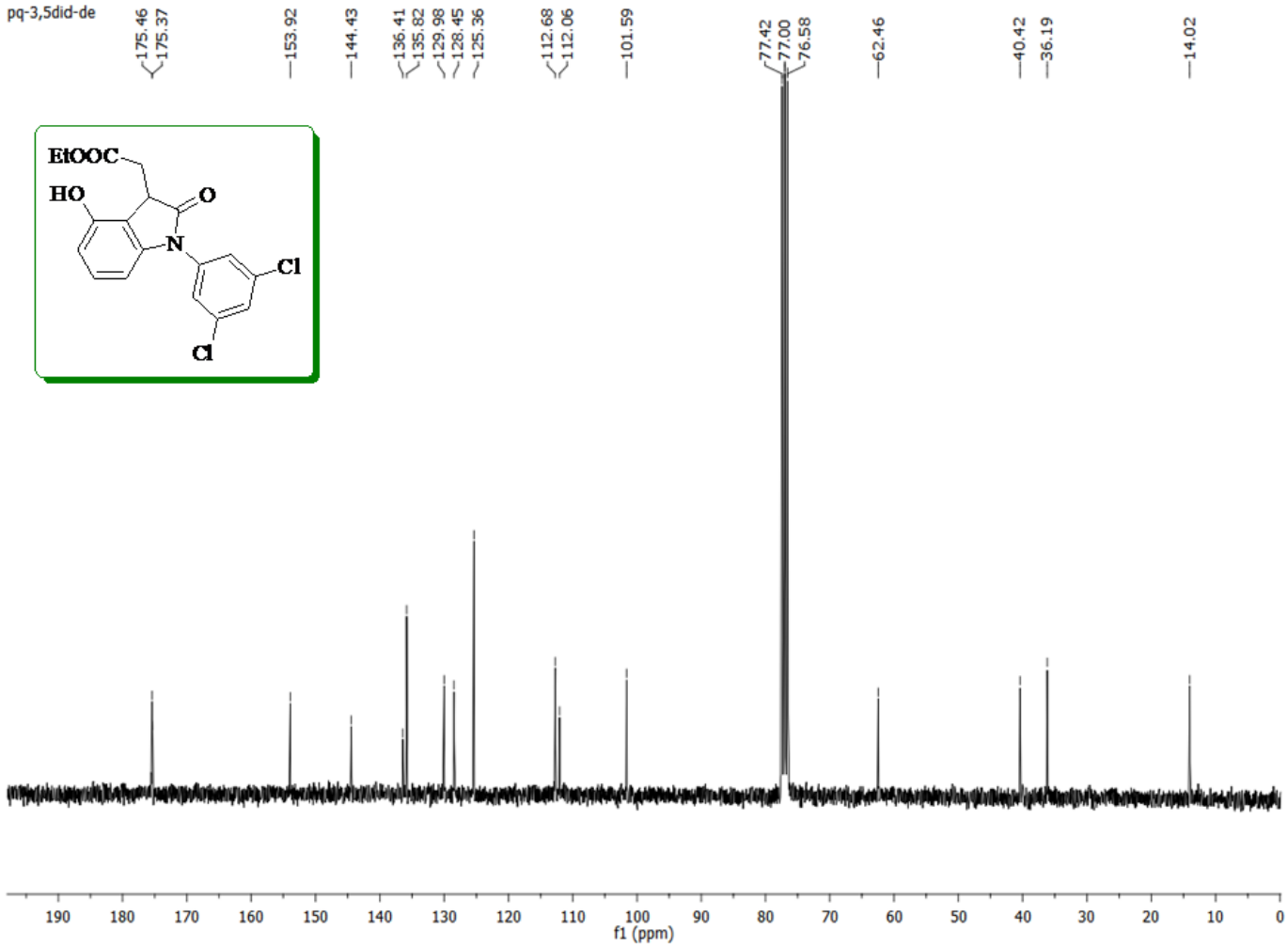

Figure S42. ^{13}C NMR spectrum of **4t**

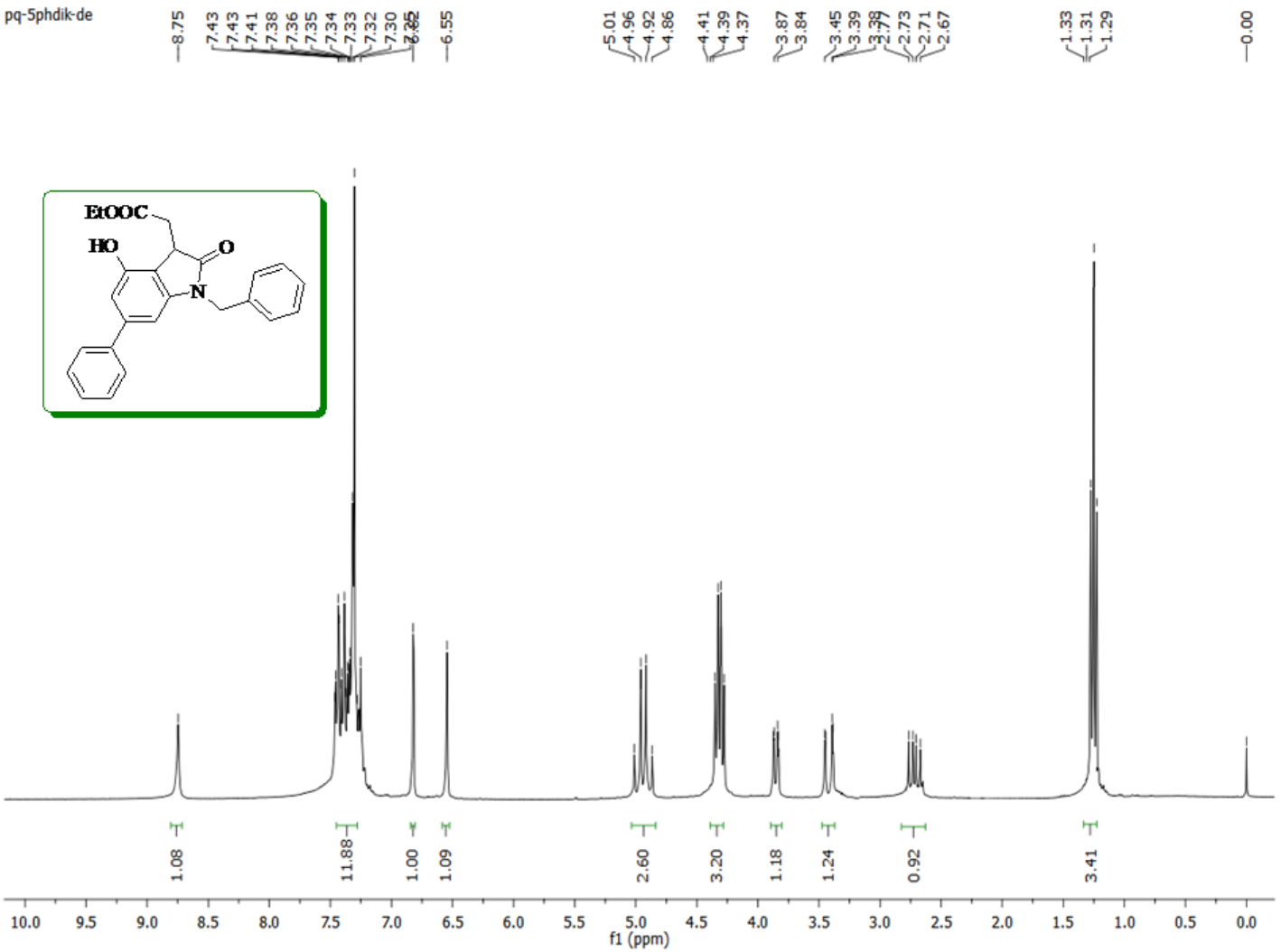

Figure S43. ^{1}H NMR spectrum of **4u**

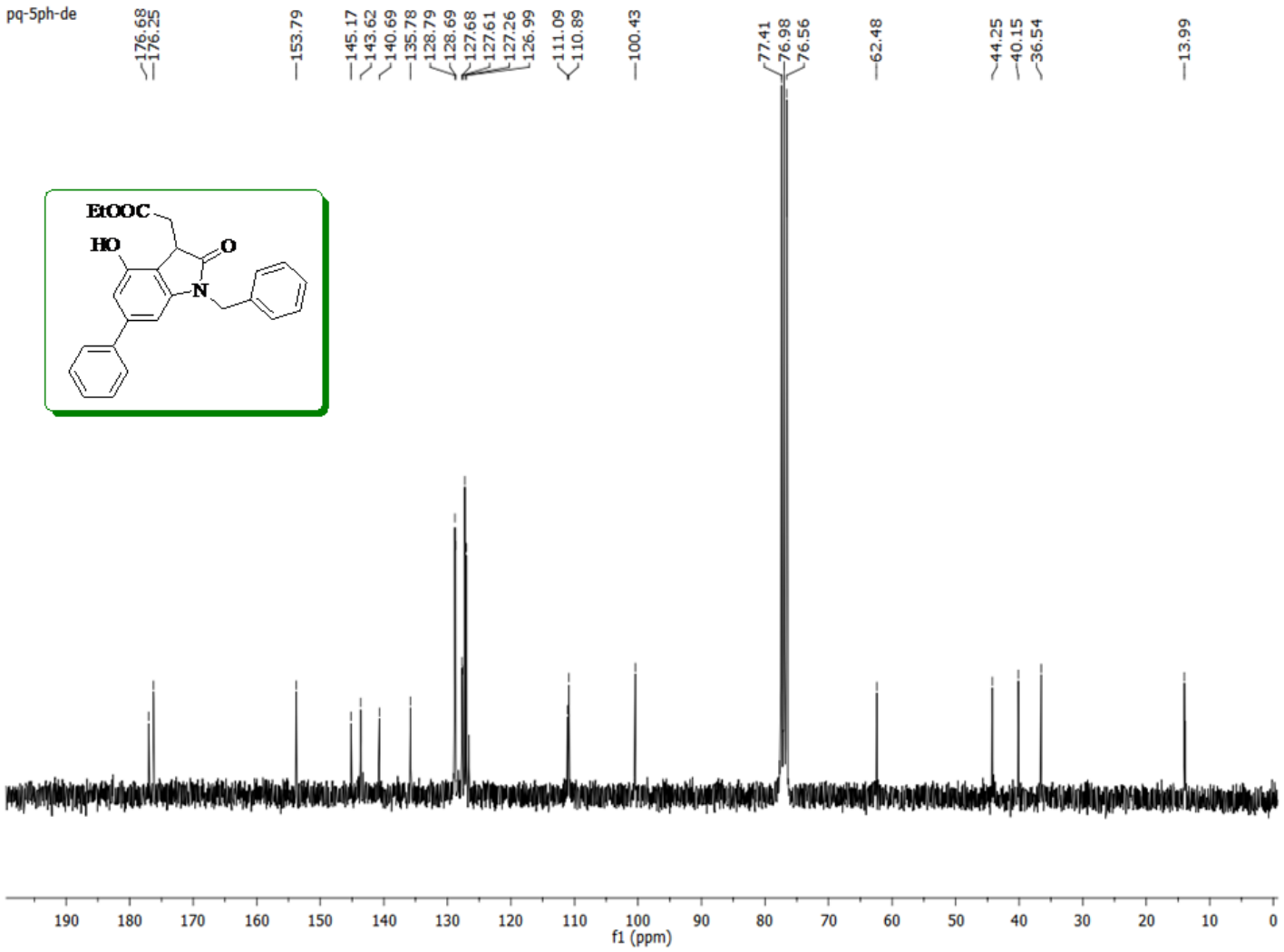

Figure S44. ^{13}C NMR spectrum of **4u**

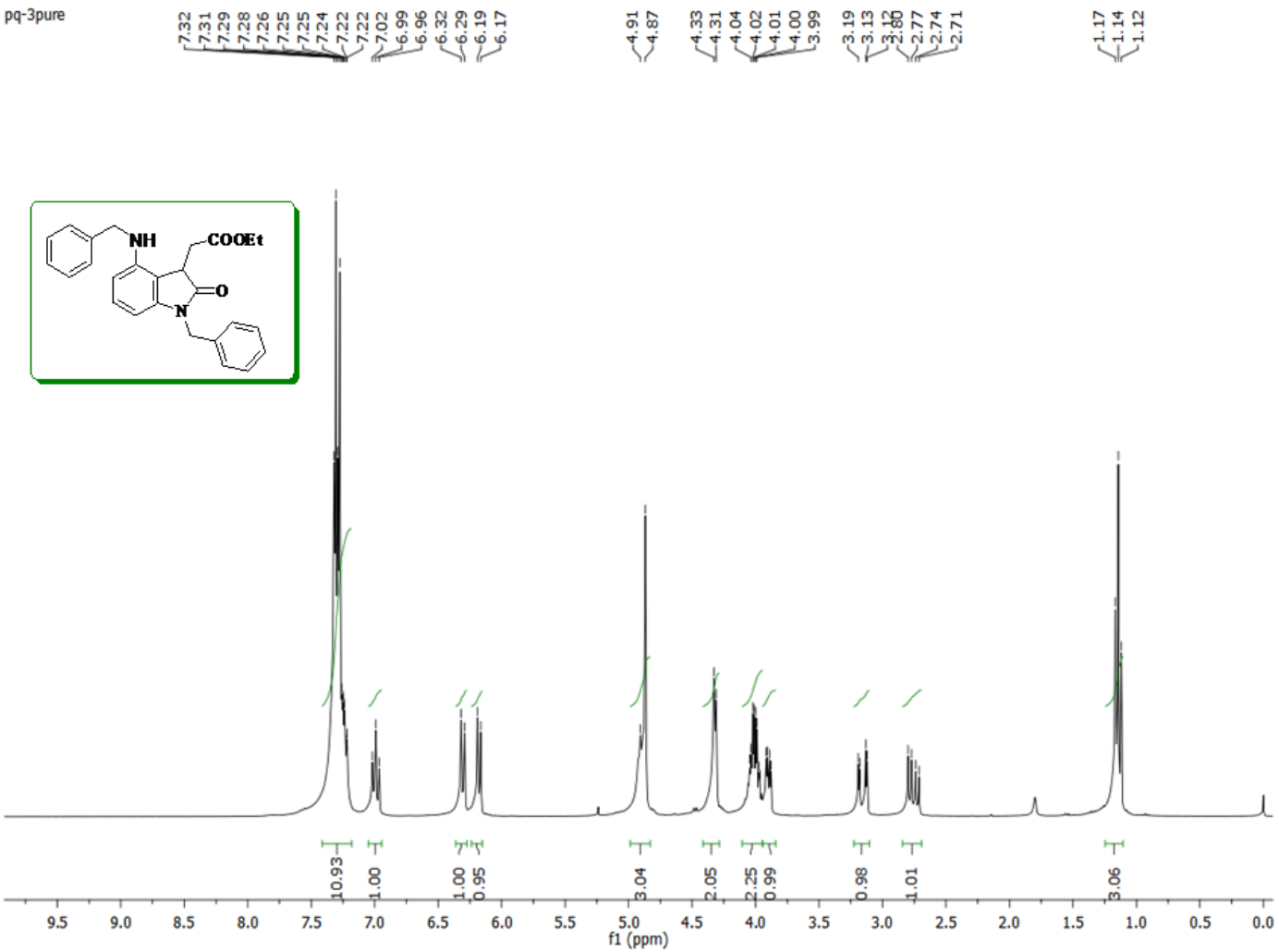

Figure S45. ^{1}H NMR spectrum of **5**

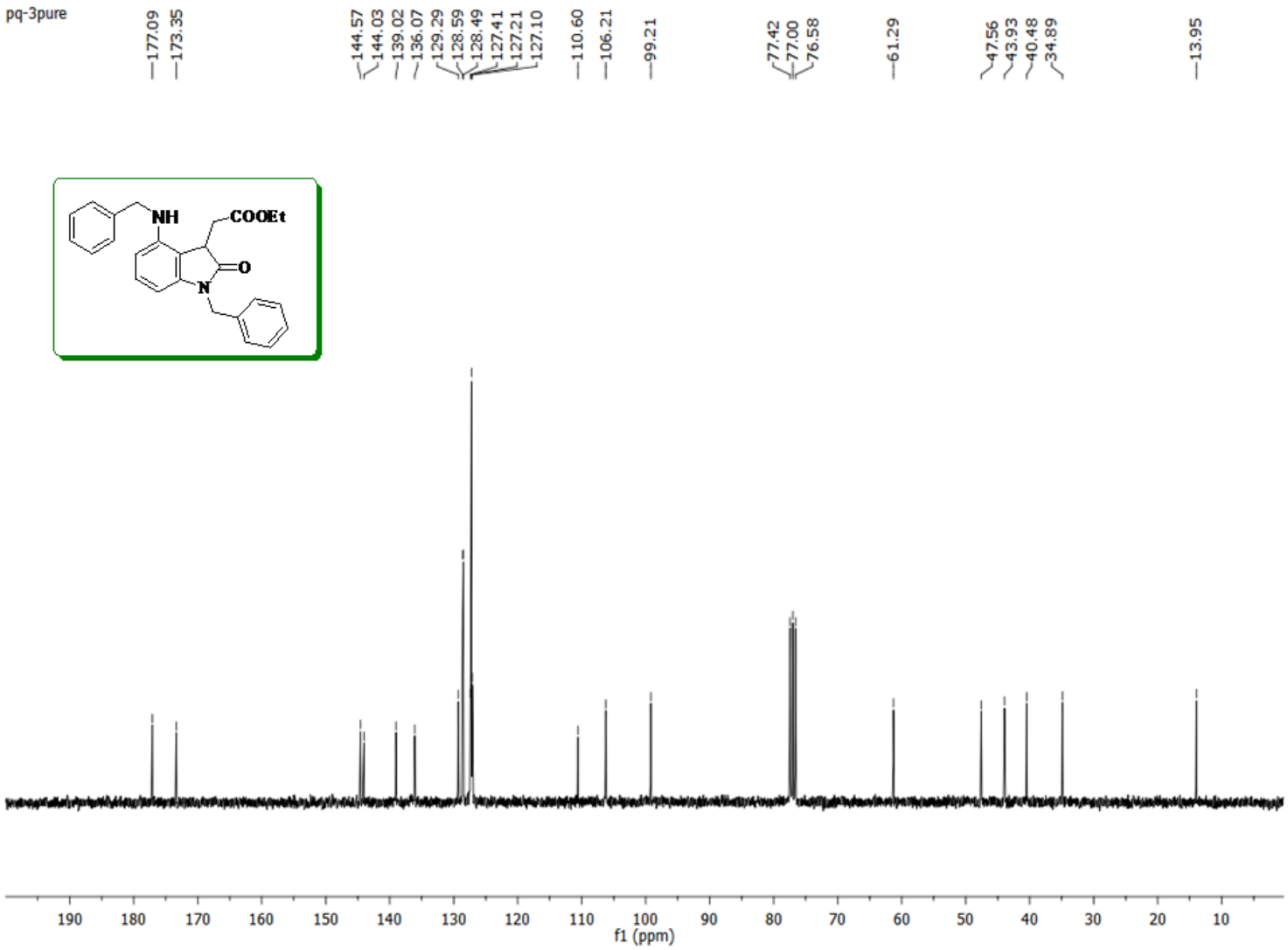

Figure S46. ^{13}C NMR spectrum of **5**

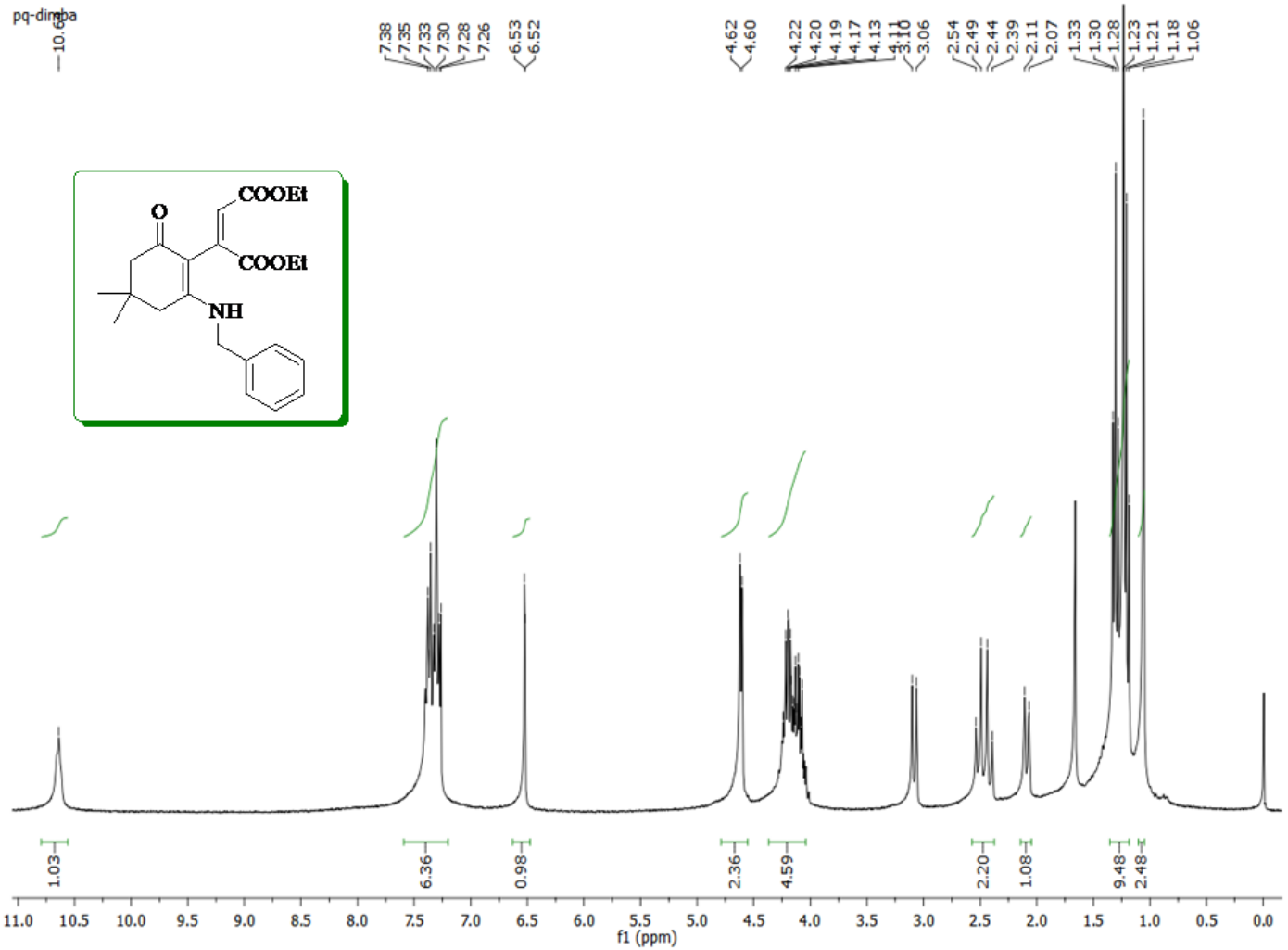

Figure S47. ^{1}H NMR spectrum of **8a**

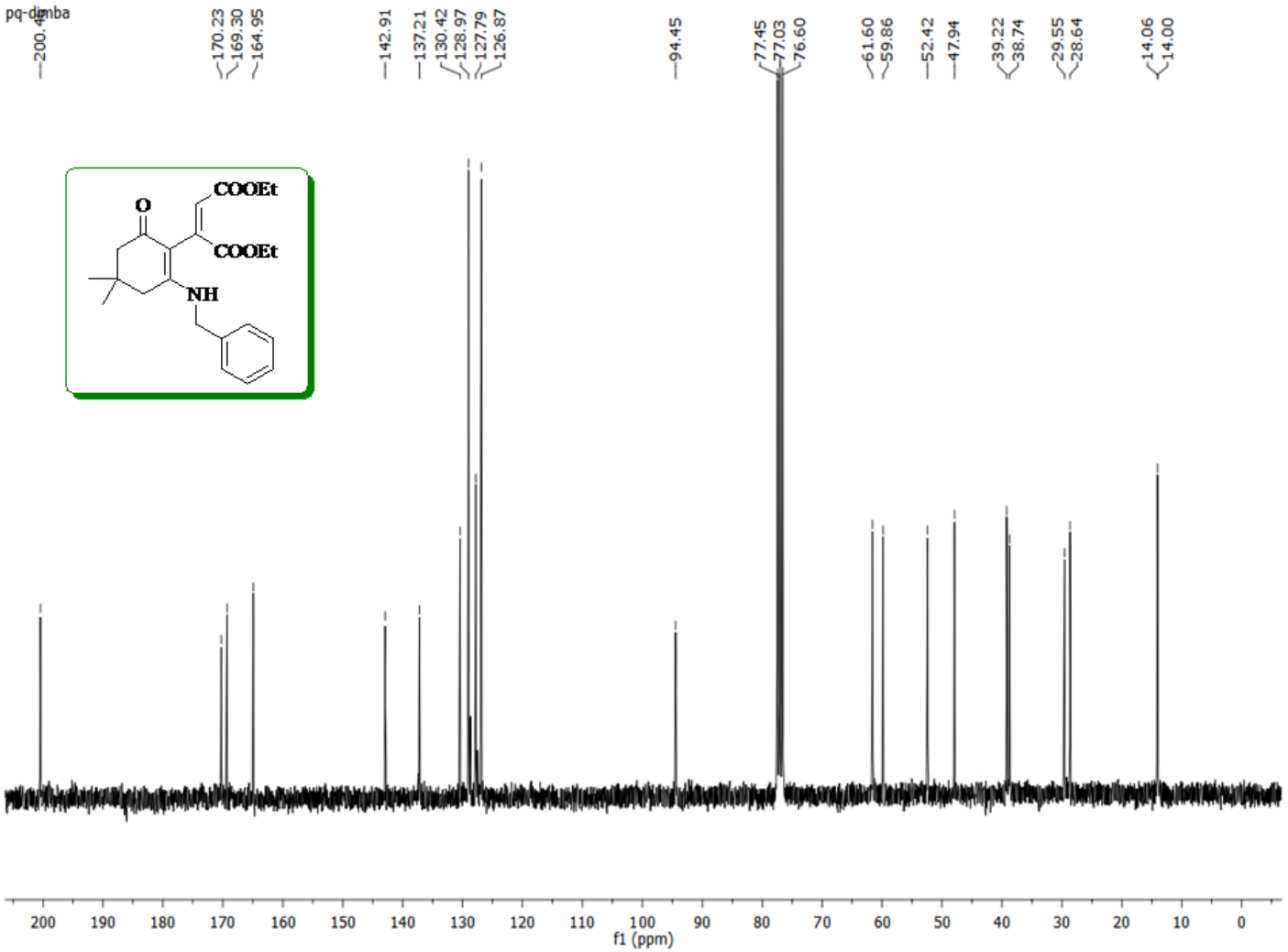

Figure S48. ^{13}C NMR spectrum of **8a**

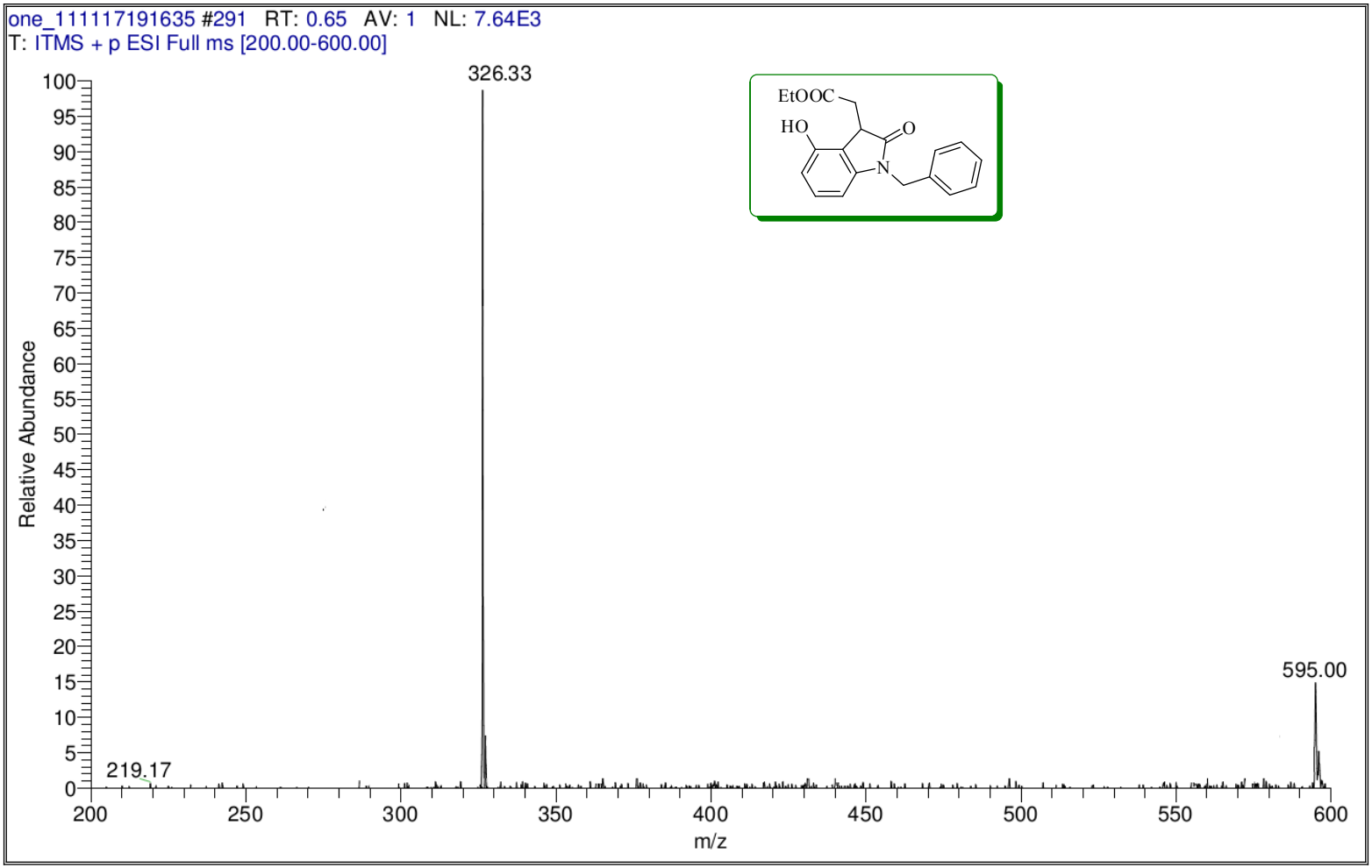

Figure S49. Mass spectrum of **4a**

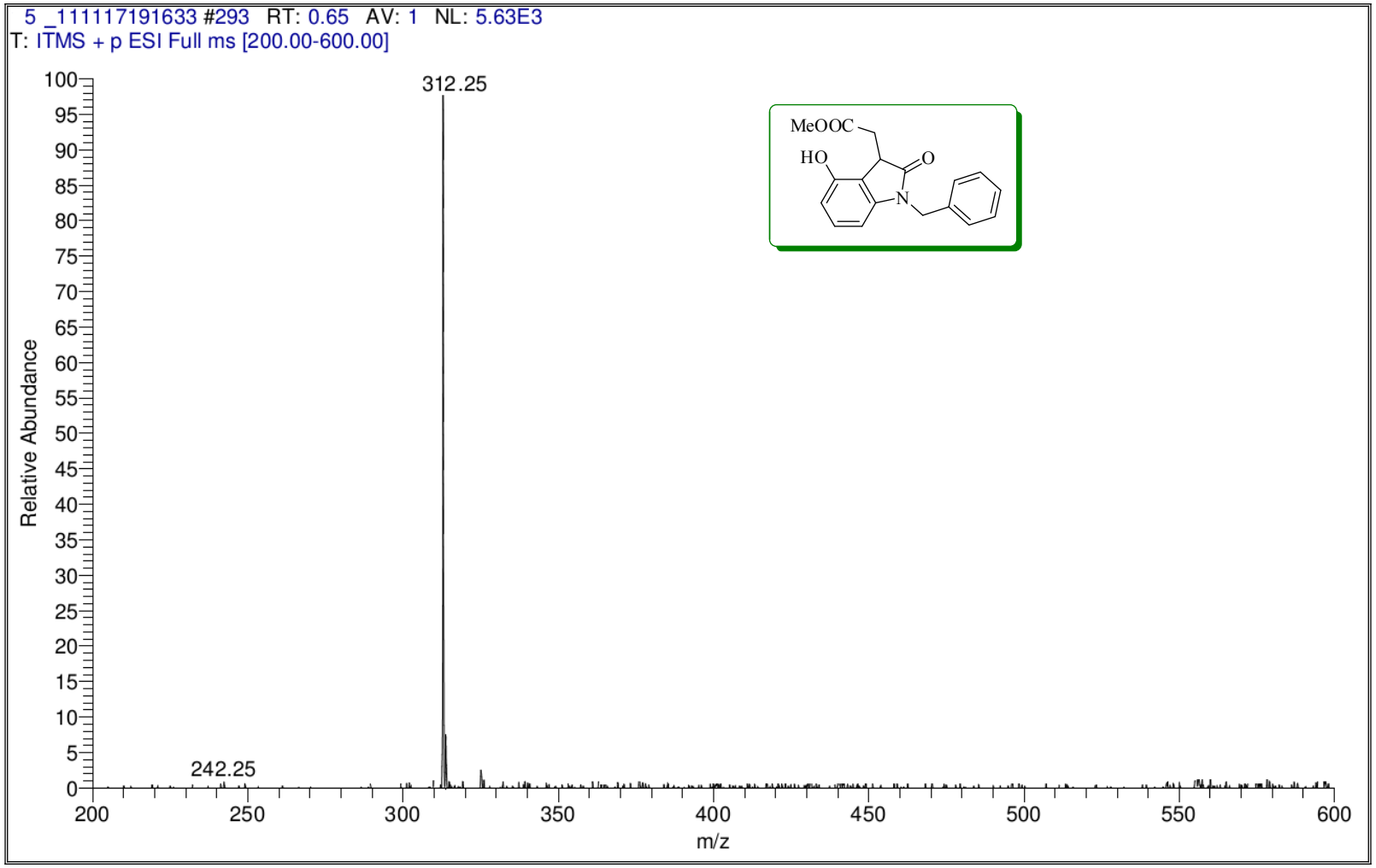

Figure S50. Mass spectrum of **4b**

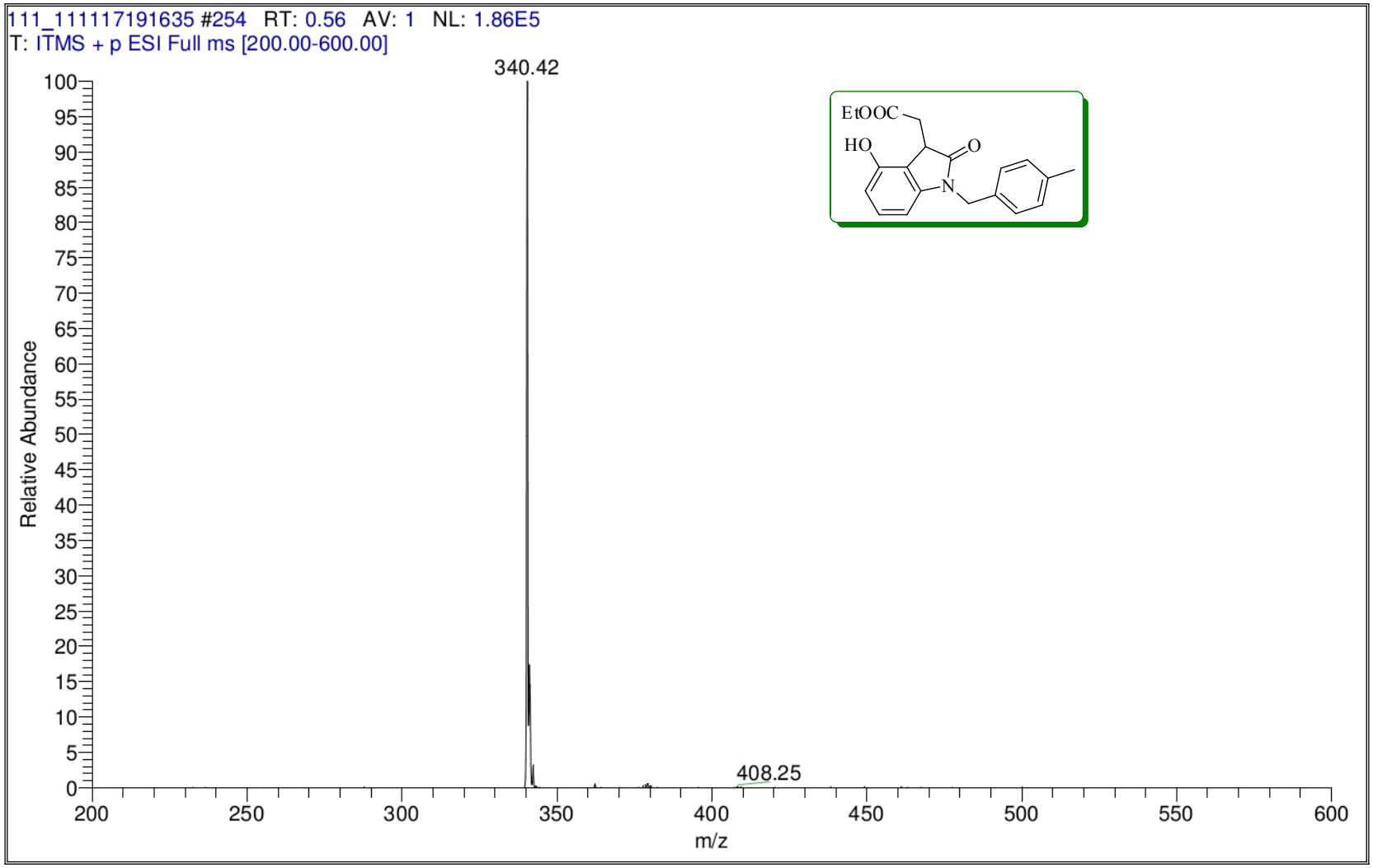

Figure S51. Mass spectrum of **4c**

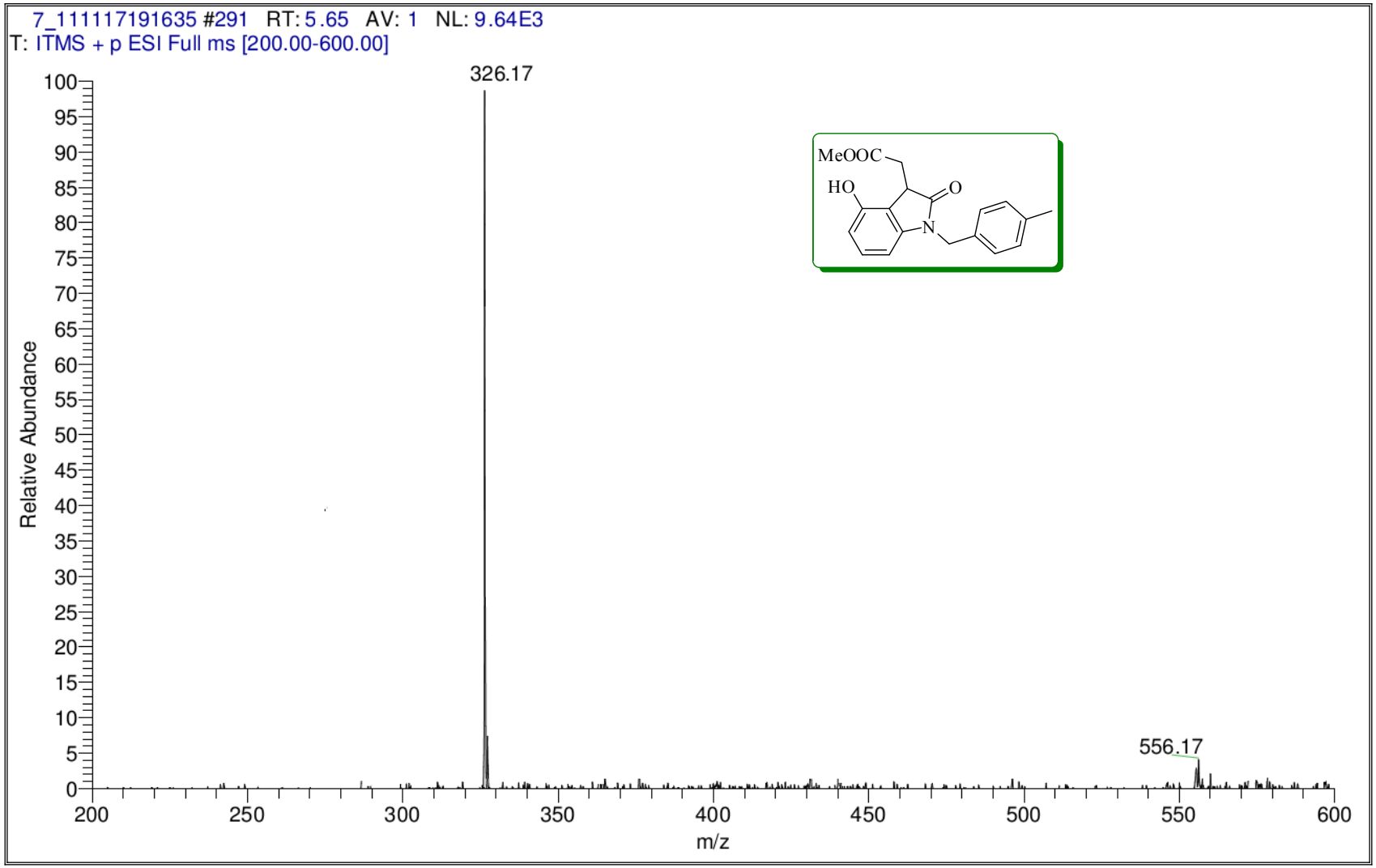

Figure S52. Mass spectrum of **4d**

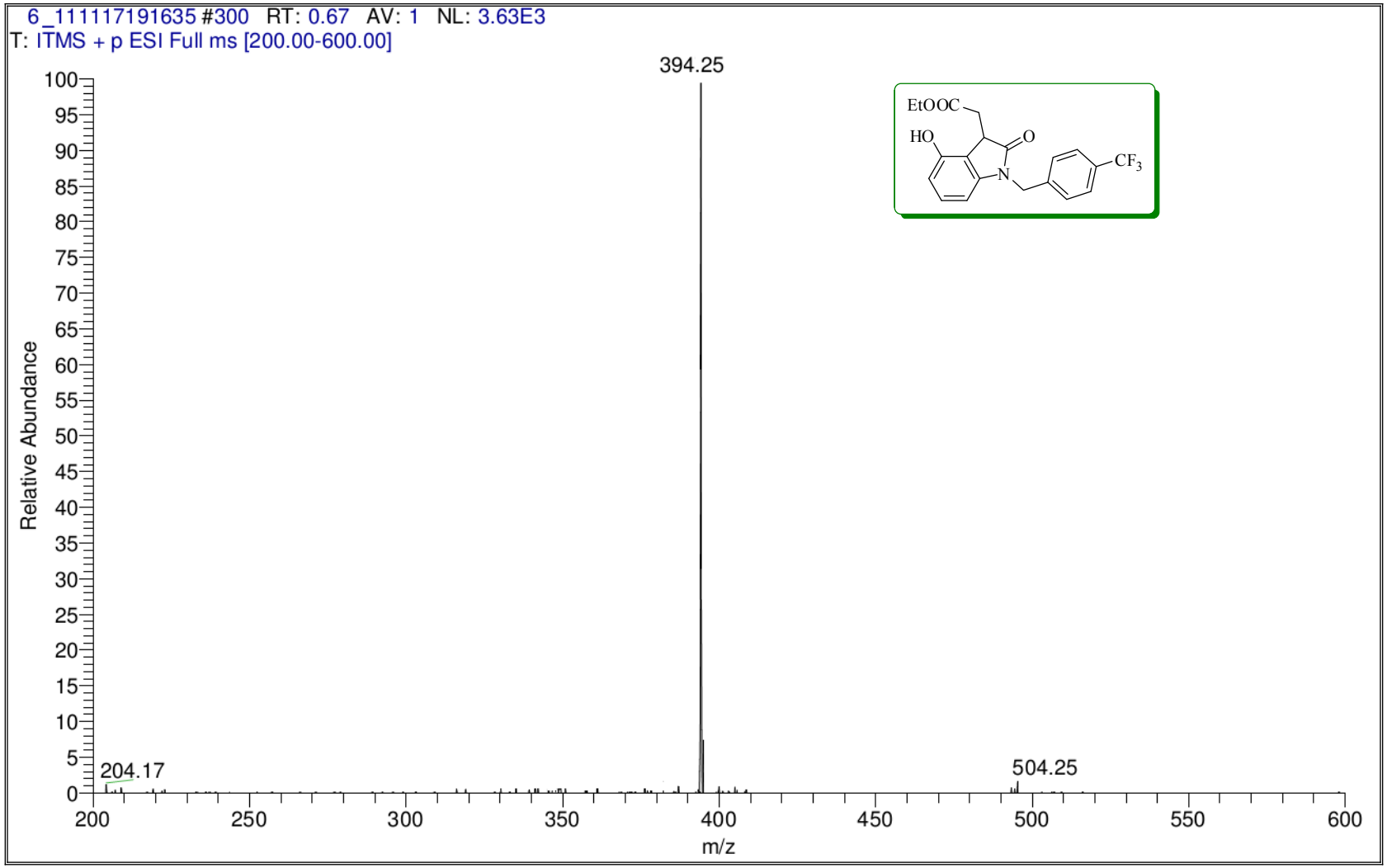

Figure S53. Mass spectrum of **4e**

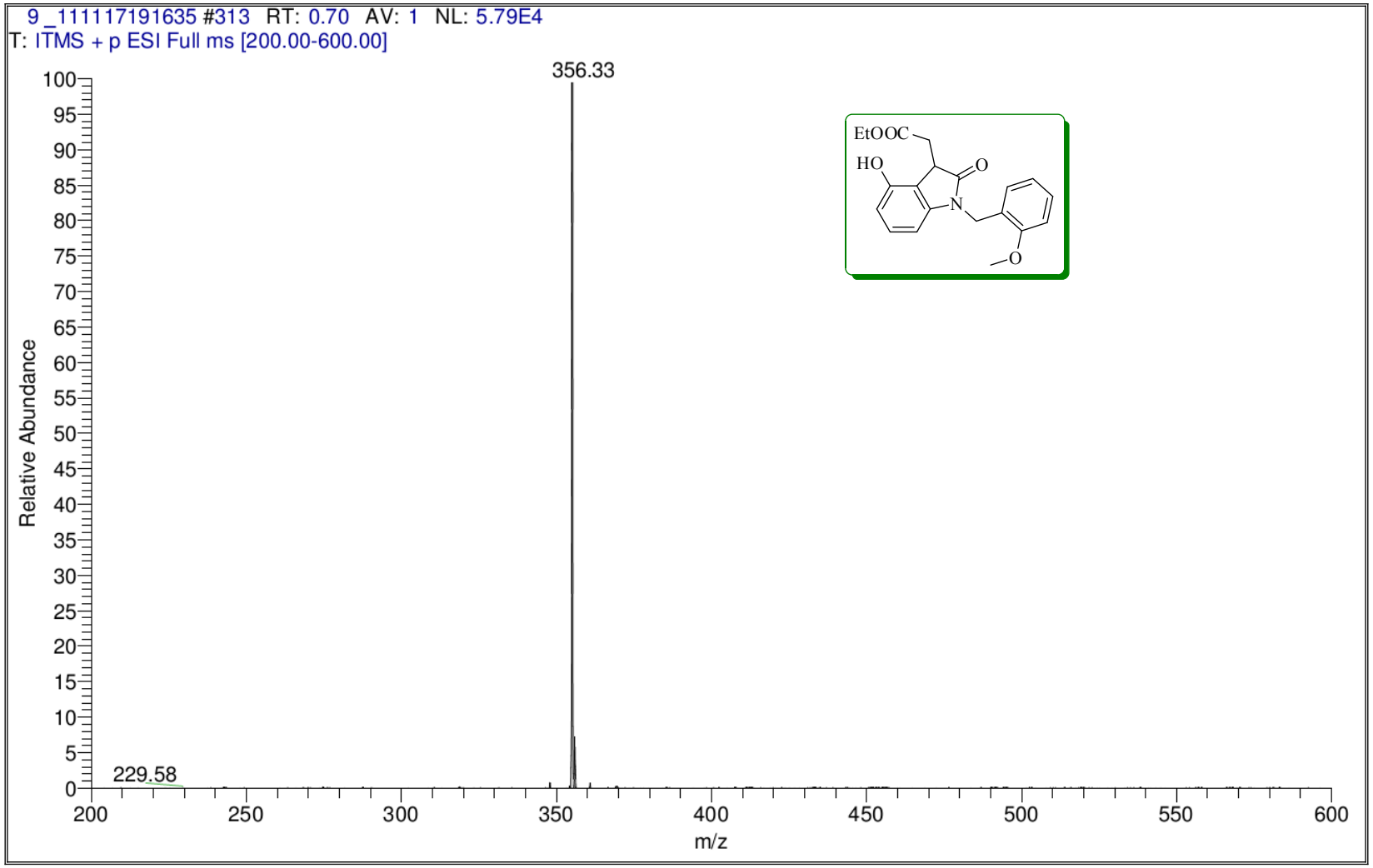

Figure S54. Mass spectrum of **4f**

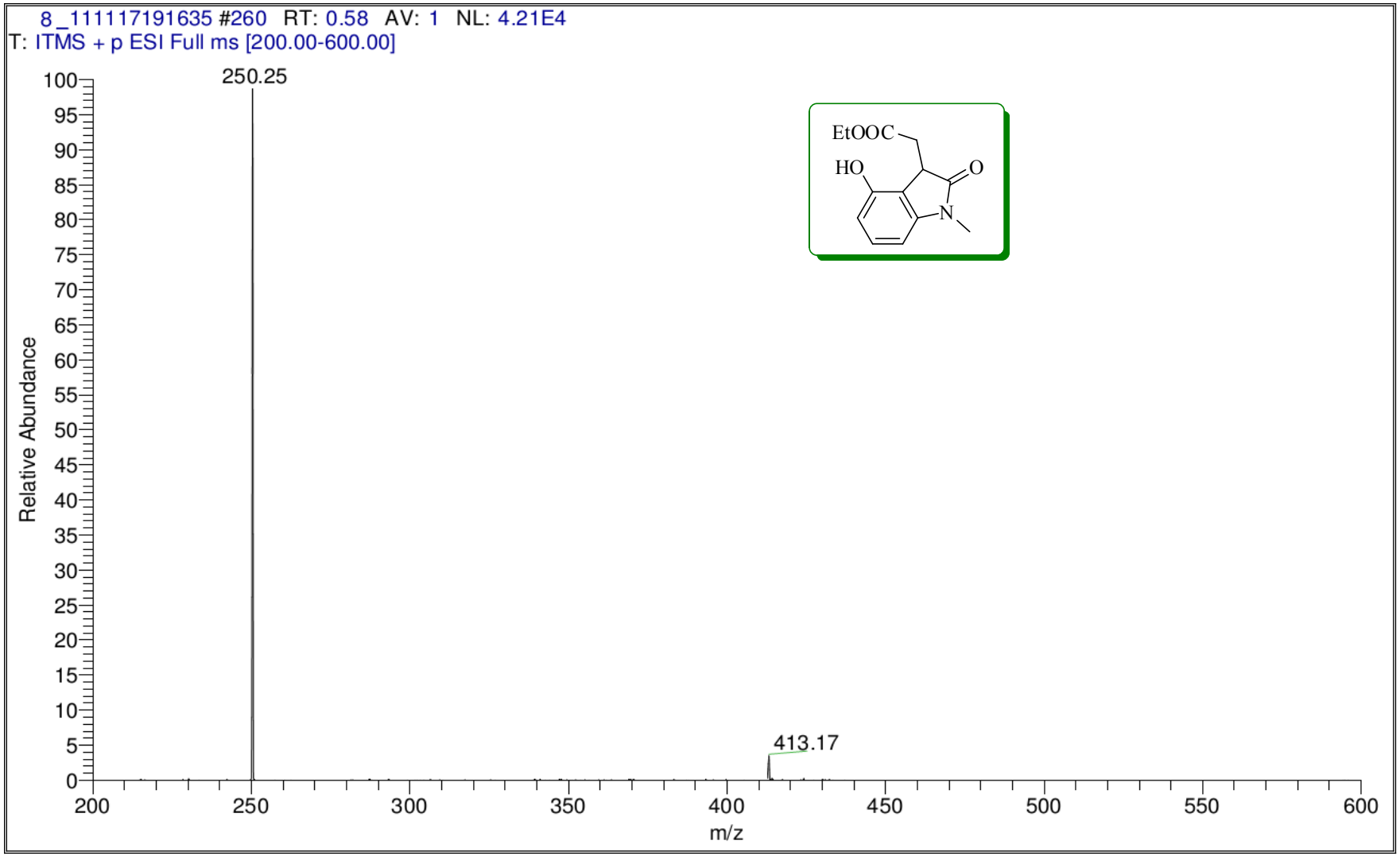

Figure S55. Mass spectrum of **4g**

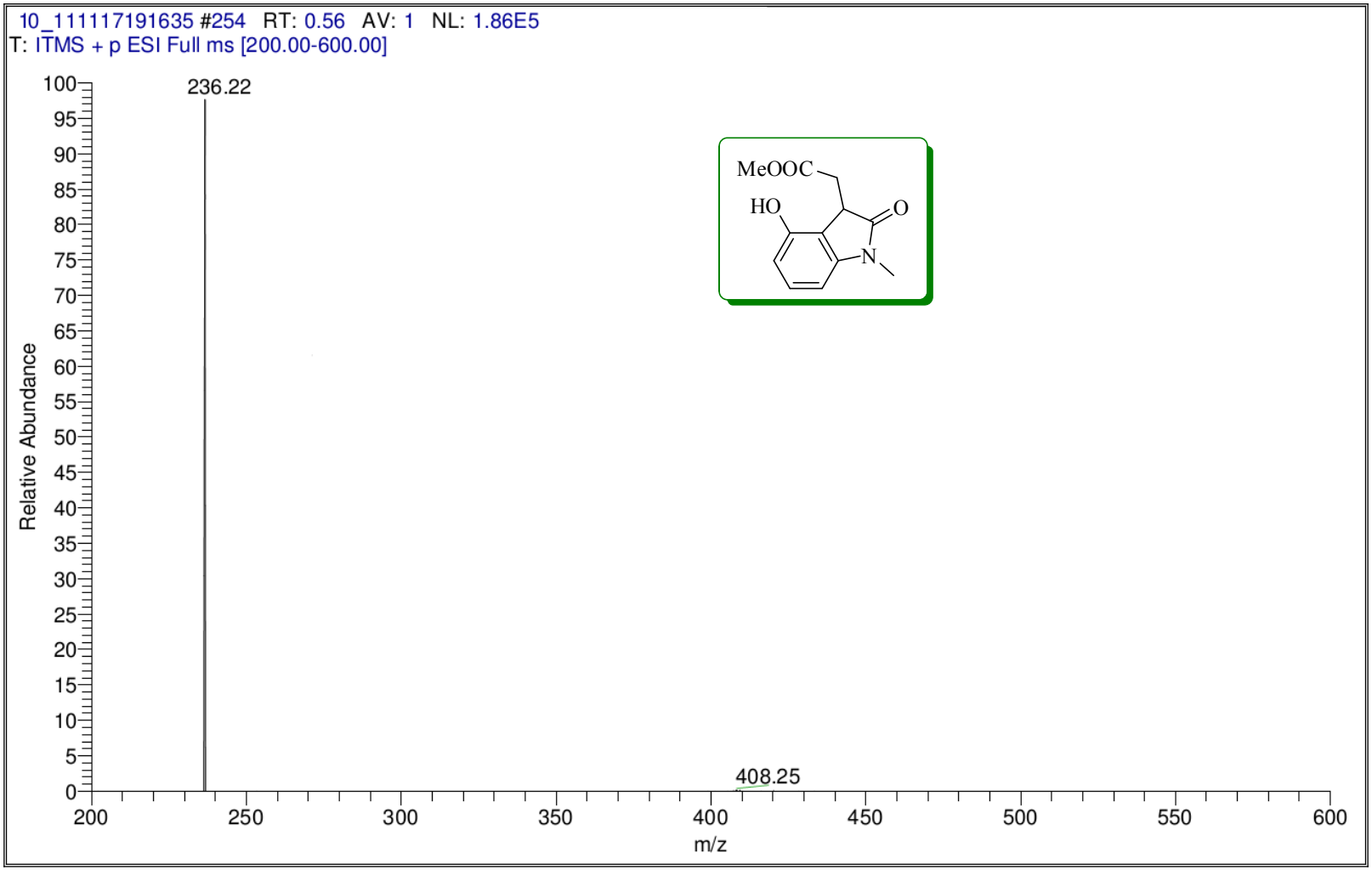

Figure S56. Mass spectrum of **4h**

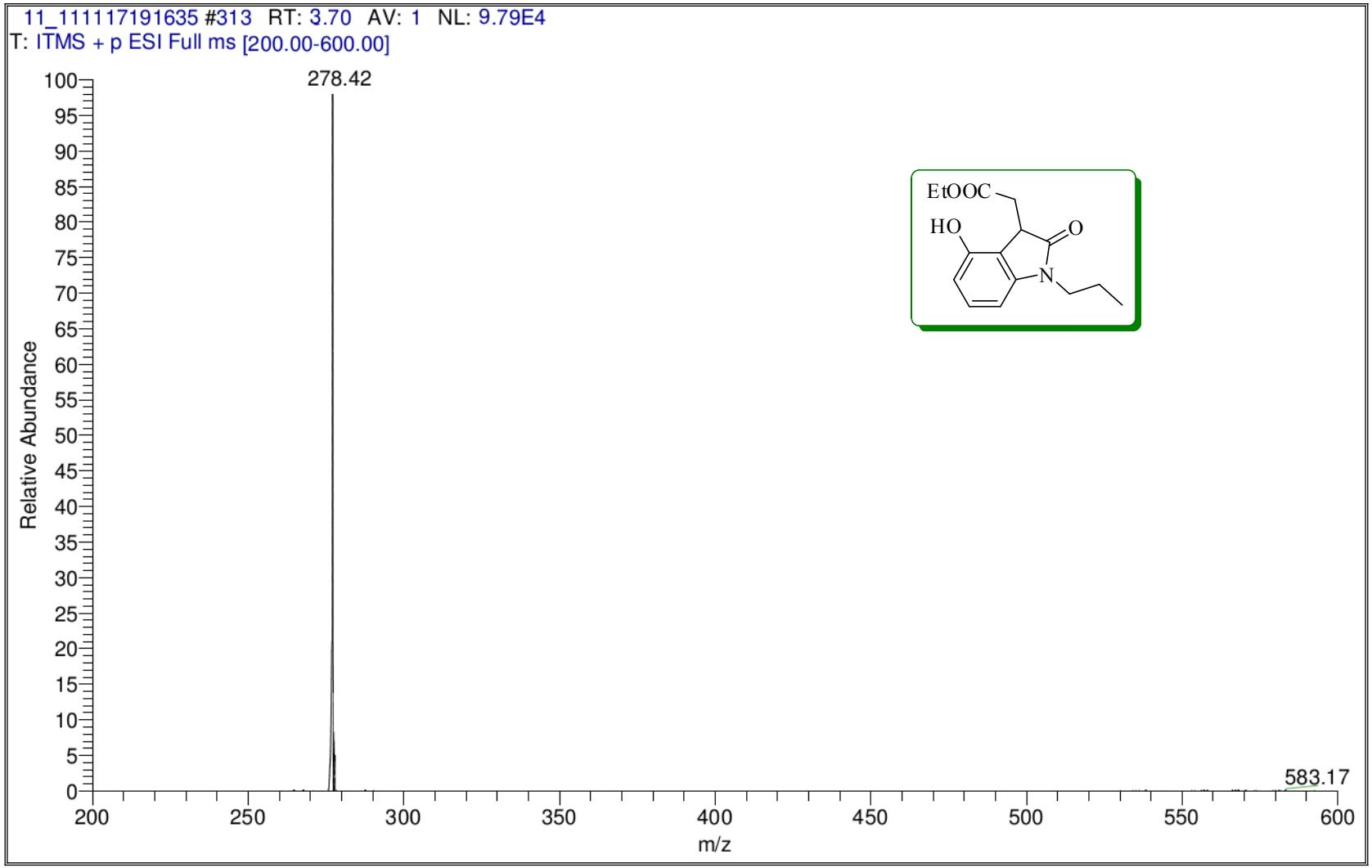

Figure S57. Mass spectrum of **4i**

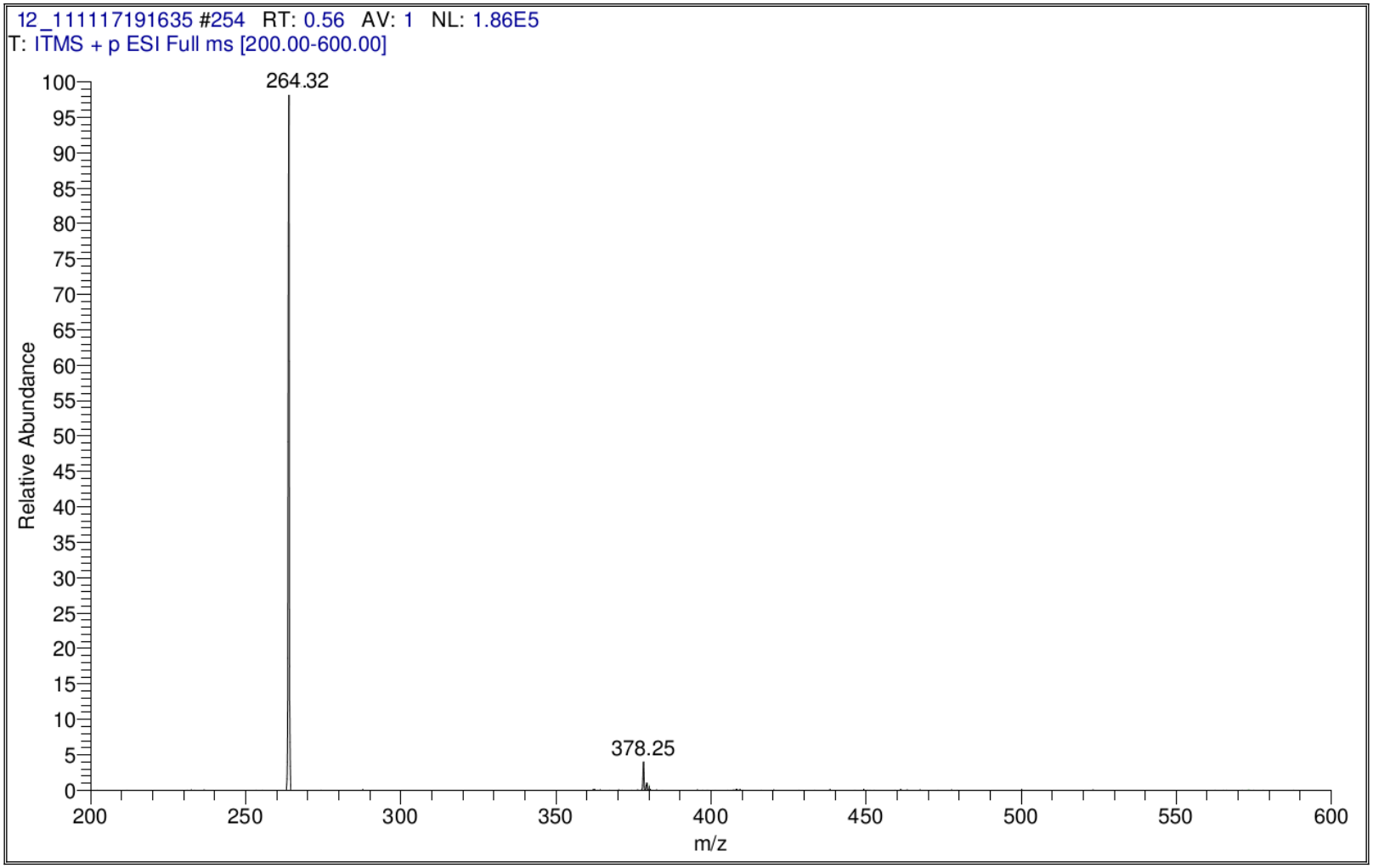

Figure S58. Mass spectrum of **4j**

Selected Crystallographic Data for 4a

Empirical formula	C19 H19 N O4	
Formula weight	325.35	
Temperature	110(2) K	
Wavelength	1.54178 Å	
Crystal system	Triclinic	
Space group	P-1	
Unit cell dimensions	a = 8.4369(10) Å	α = 75.851(6)°.
	b = 12.6589(13) Å	β = 89.973(7)°.
	c = 15.5346(16) Å	γ = 83.469(7)°.
Volume	1597.7(3) $Å^3$	
Z	4	
Density (calculated)	1.353 Mg/m^3	
Absorption coefficient	0.779 mm^{-1}	
F(000)	688	
Crystal size	0.08 x 0.07 x 0.03 mm^3	
Theta range for data collection	2.93 to 60.00°.	
Index ranges	-9<=h<=9, -13<=k<=14, 0<=l<=17	
Reflections collected	6568	
Independent reflections	6577 [R(int) = 0.0000]	
Completeness to theta = 60.00°	88.6 %	
Absorption correction	Semi-empirical from equivalents	
Max. and min. transmission	0.9770 and 0.9403	
Refinement method	Full-matrix least-squares on F^2	
Data / restraints / parameters	6577 / 0 / 438	
Goodness-of-fit on F^2	1.073	
Final R indices [I>2sigma(I)]	R1 = 0.0571, wR2 = 0.1642	
R indices (all data)	R1 = 0.0825, wR2 = 0.1867	
Largest diff. peak and hole	0.354 and -0.357 $e.Å^{-3}$	