Electronic Supplementary Material (ESI) for Organic & Biomolecular Chemistry.

Electronic Supplementary Information

$SnCl_4$-Mediated One-Pot Synthesis of 2,4,5-Trisubstituted Thiazoles from Nitro-Substituted Donor-Acceptor Cyclopropanes and Thioamides

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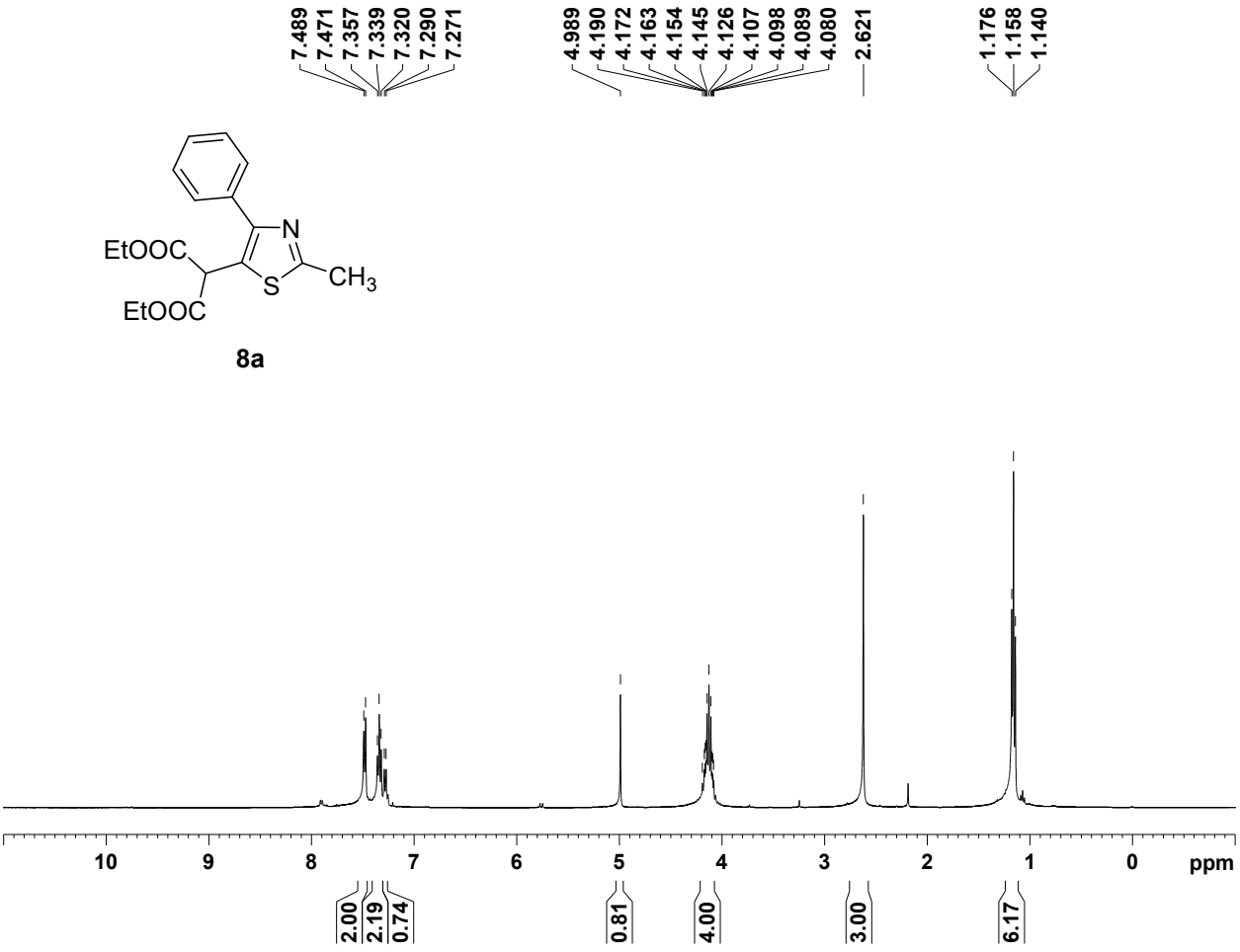

Figure 1. ^{1}H NMR (400 MHz, $CDCl_3$) spectrum of **8a**

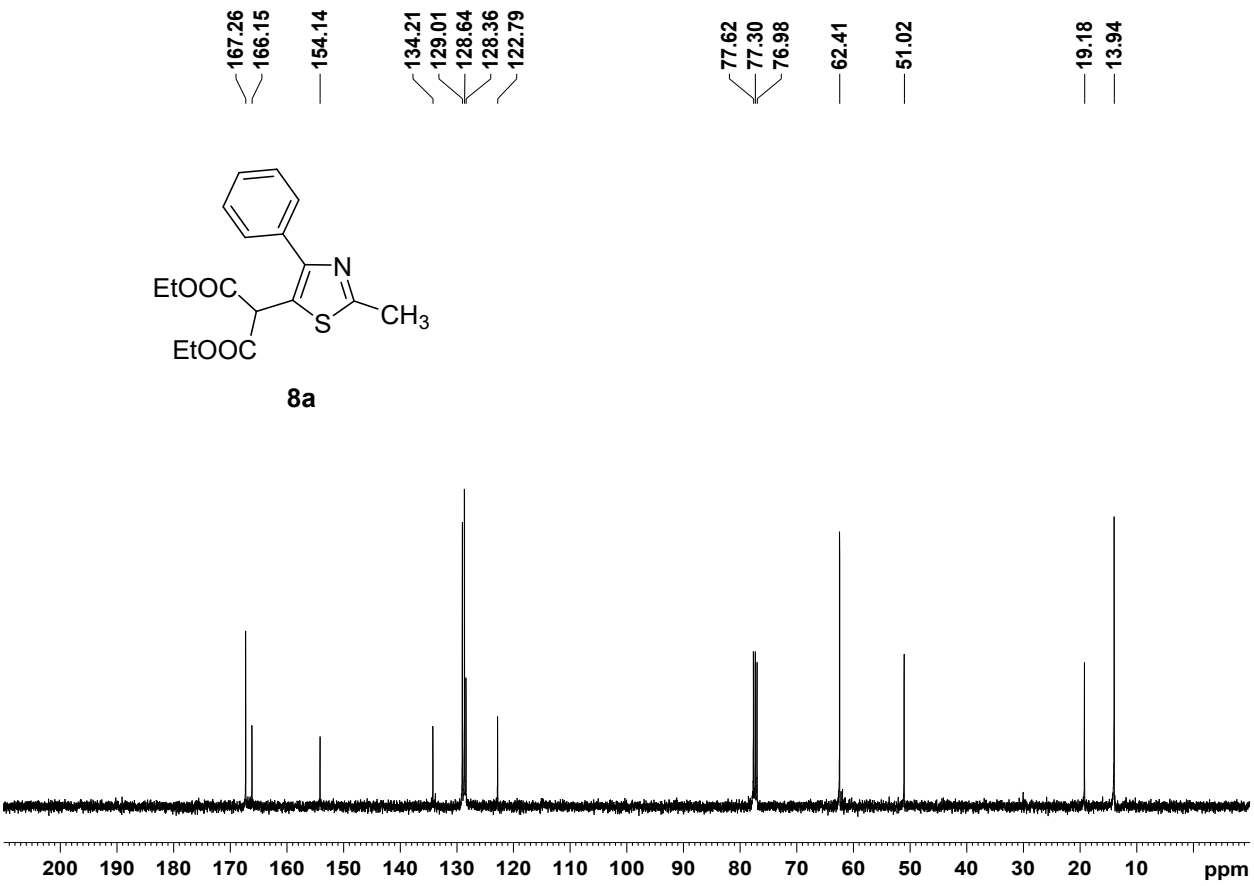

Figure 2. ^{13}C NMR (100 MHz, $CDCl_3$) spectrum of **8a**

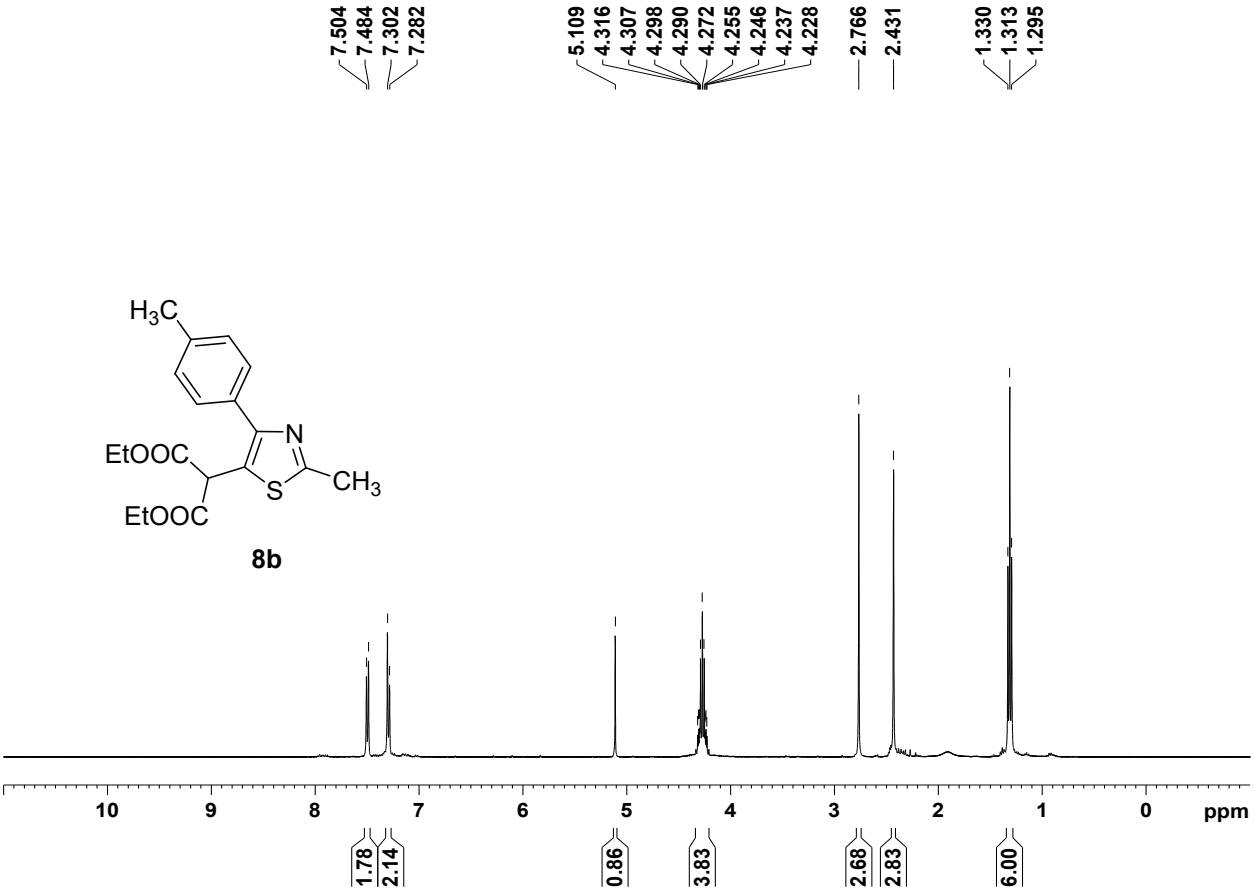

Figure 3. ^{1}H NMR (400 MHz, $CDCl_3$) spectrum of **8b**

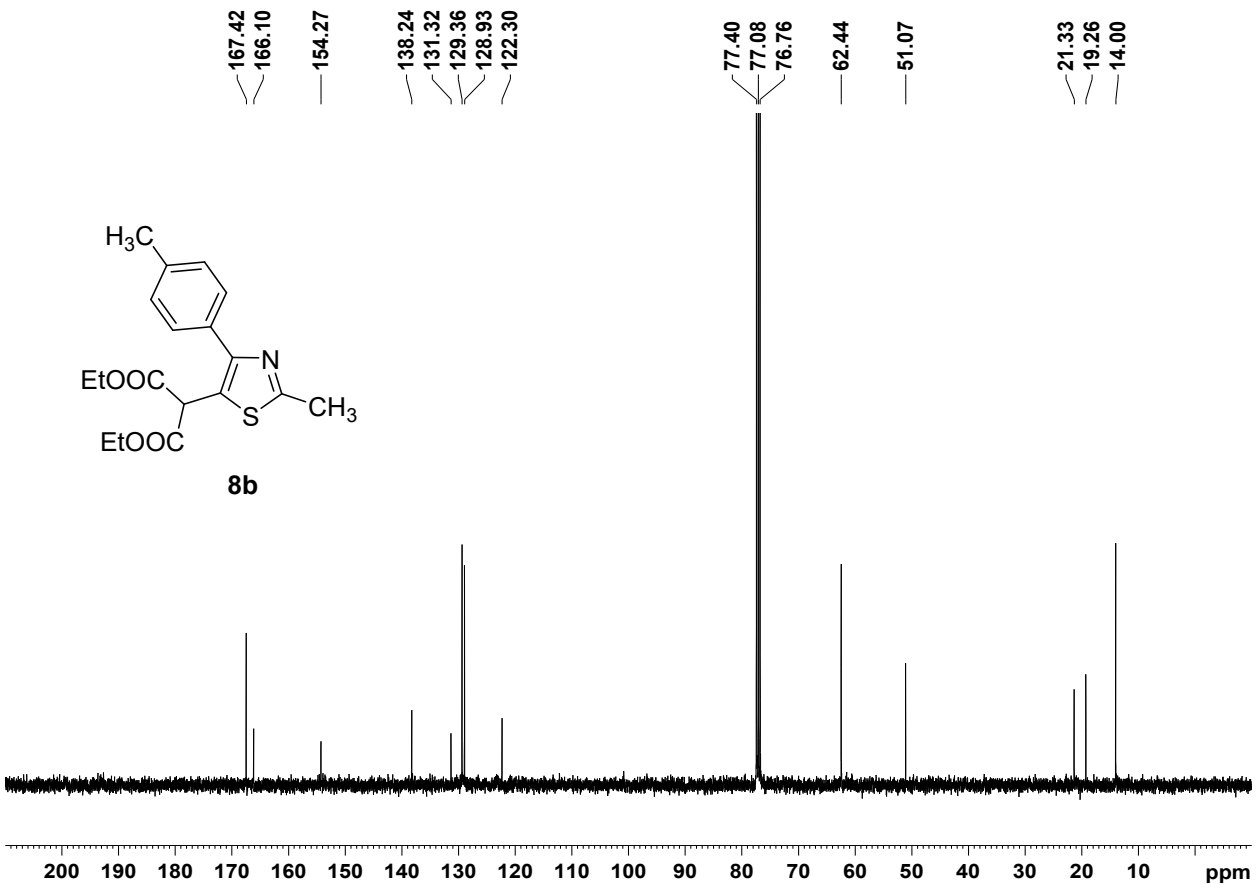

Figure 4. ^{13}C NMR (100 MHz, $CDCl_3$) spectrum of **8b**

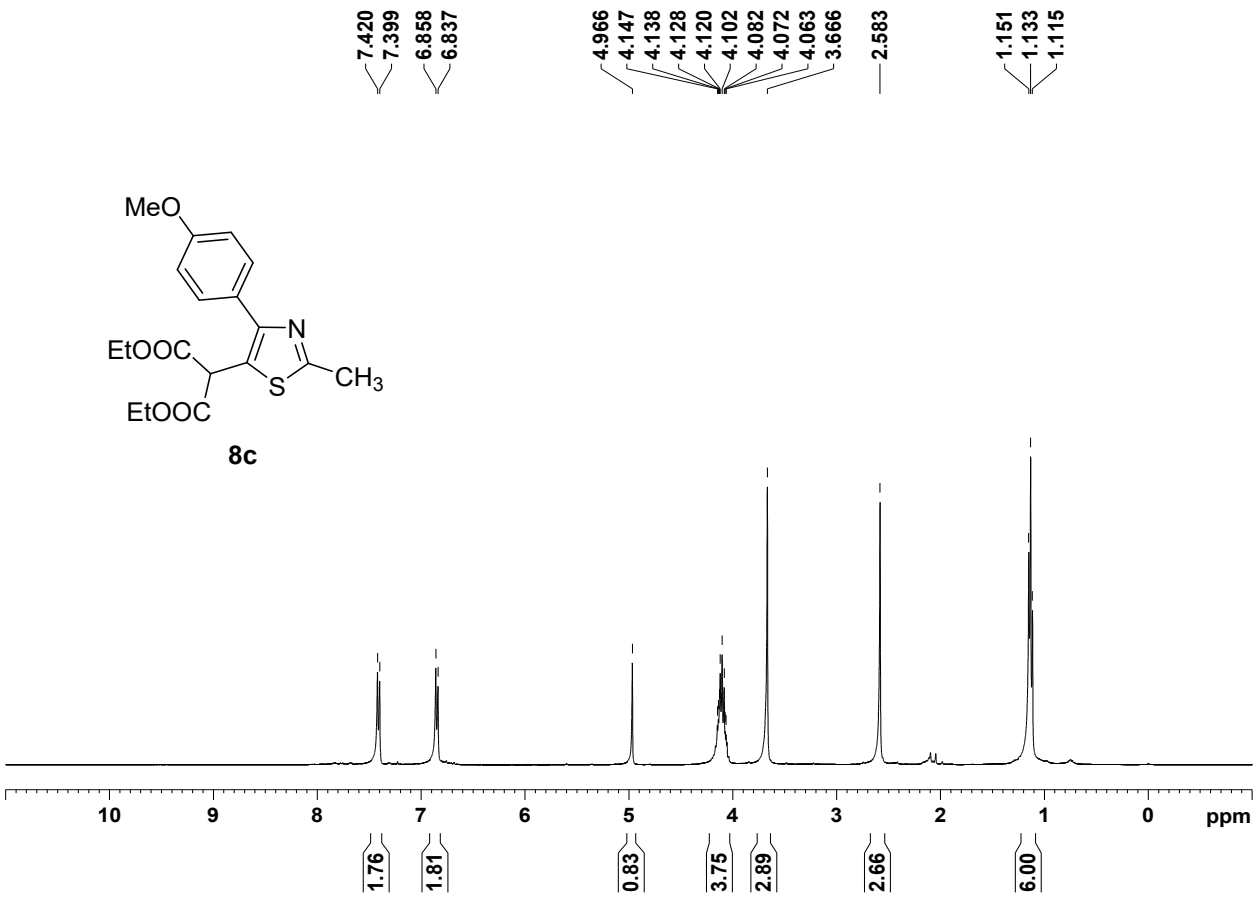

Figure 5. ^{1}H NMR (400 MHz, $CDCl_3$) spectrum of **8c**

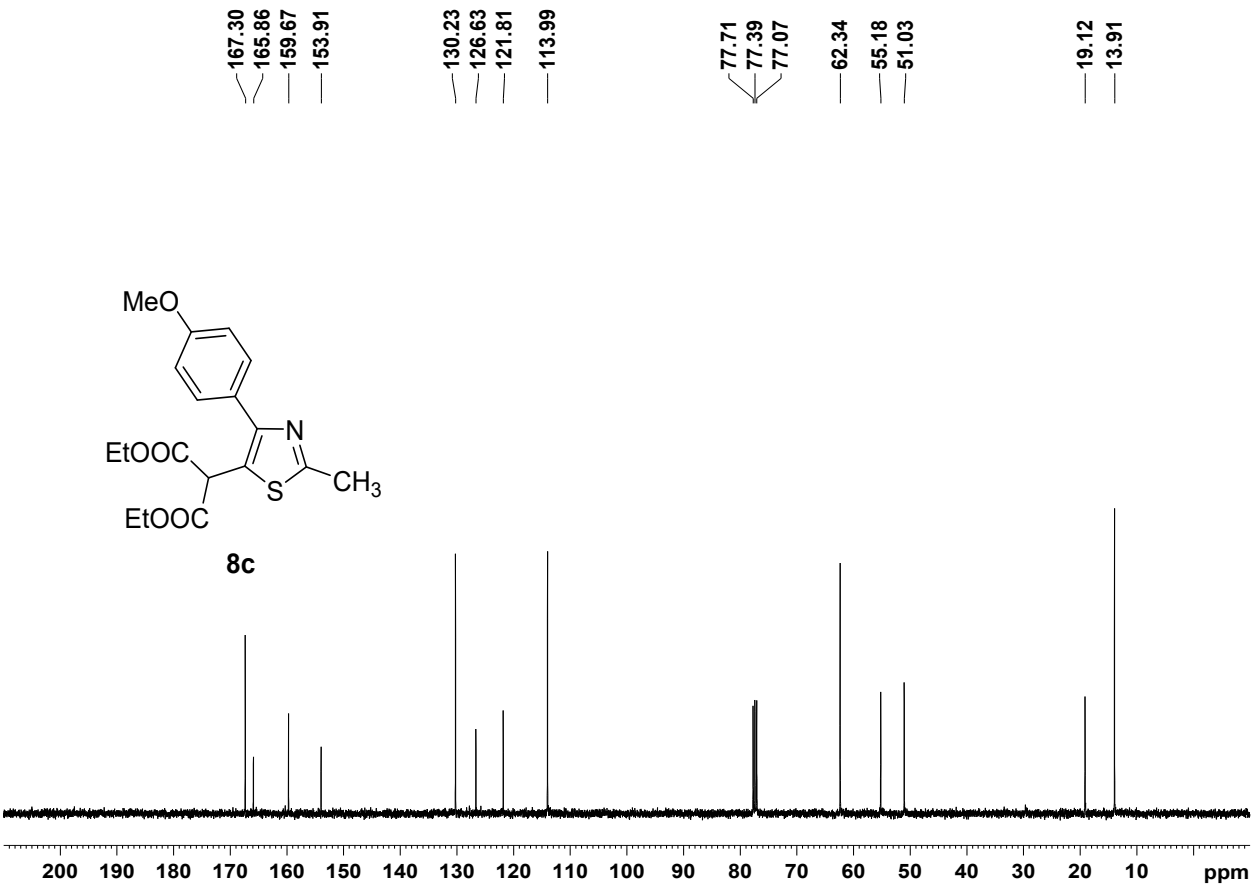

Figure 6. ^{13}C NMR (100 MHz, $CDCl_3$) spectrum of **8c**

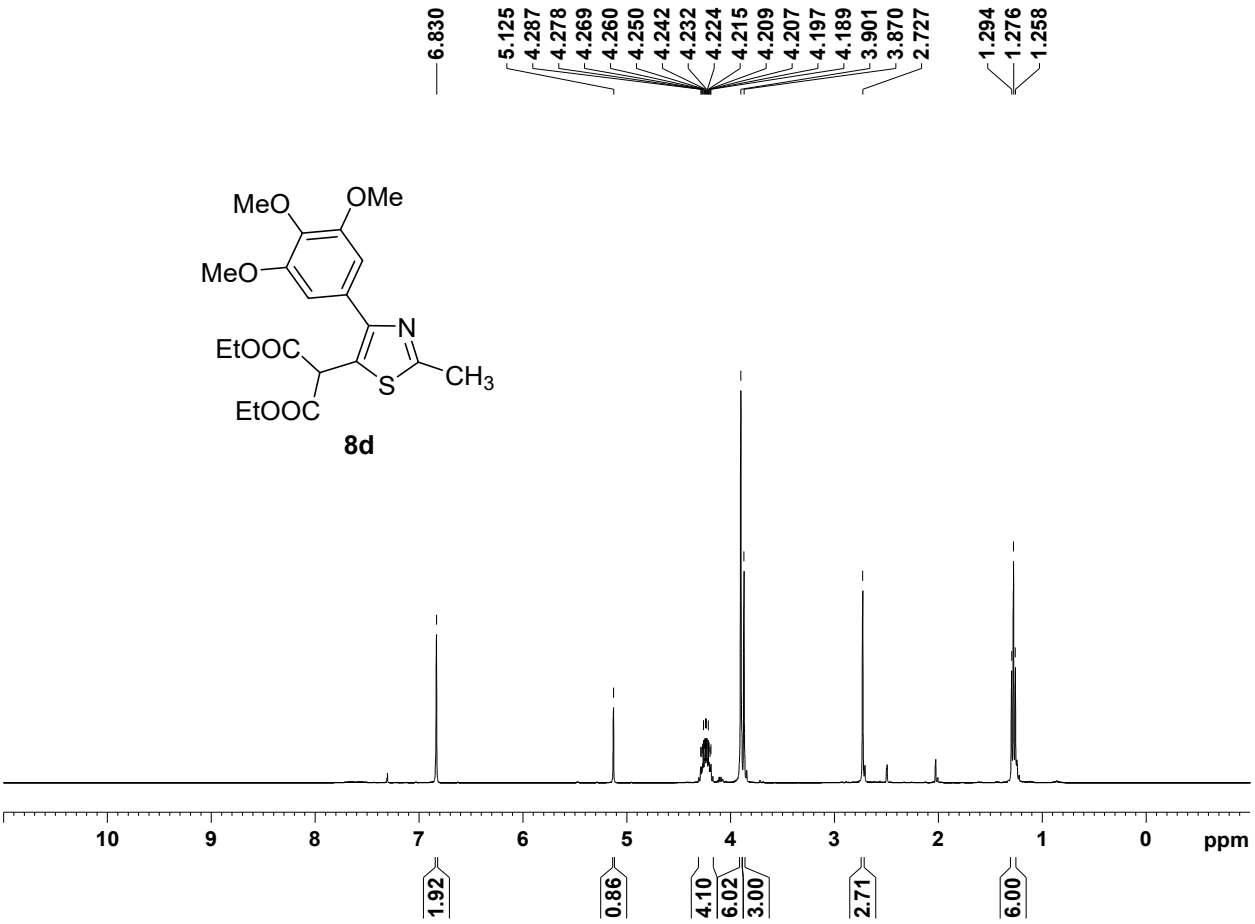

Figure 7. ^{1}H NMR (400 MHz, $CDCl_3$) spectrum of **8d**

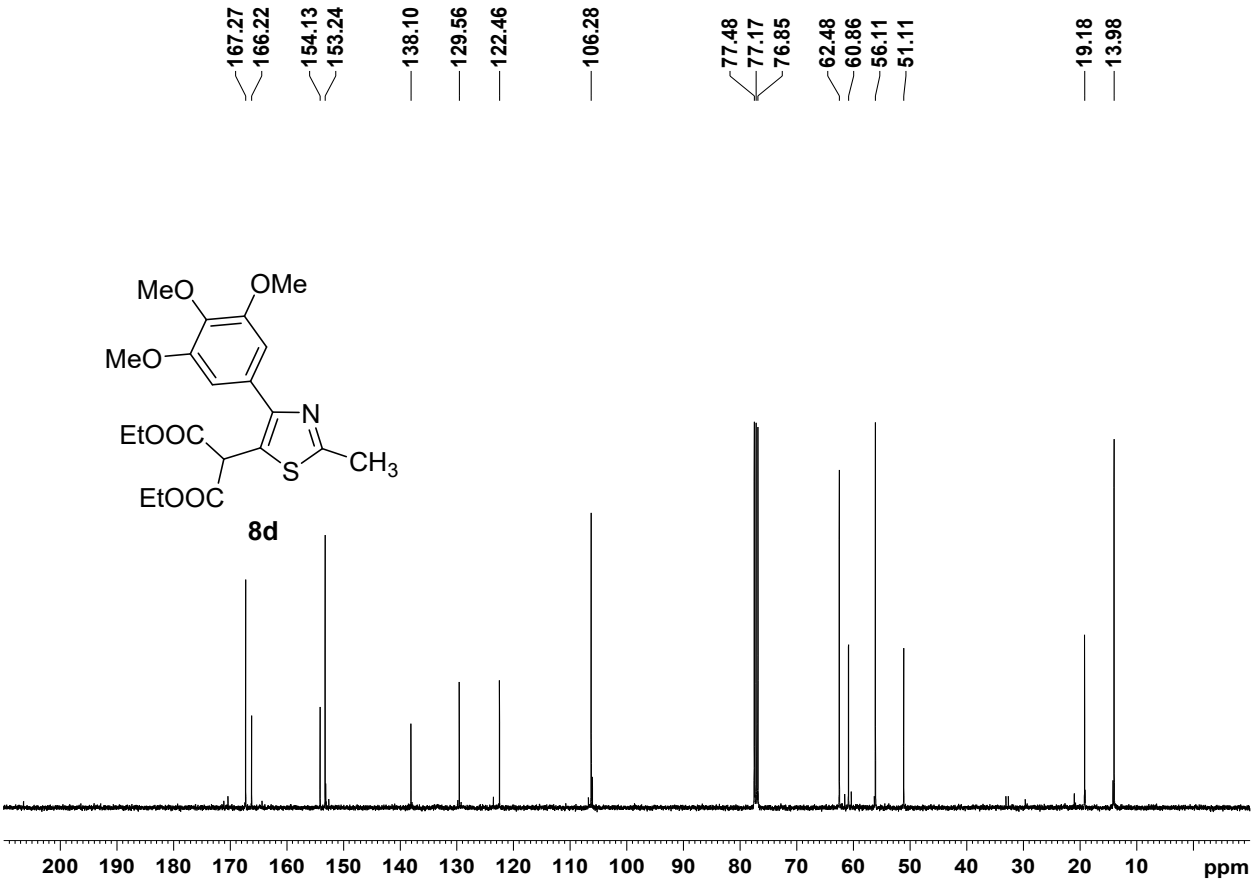

Figure 8. ^{13}C NMR (100 MHz, $CDCl_3$) spectrum of **8d**

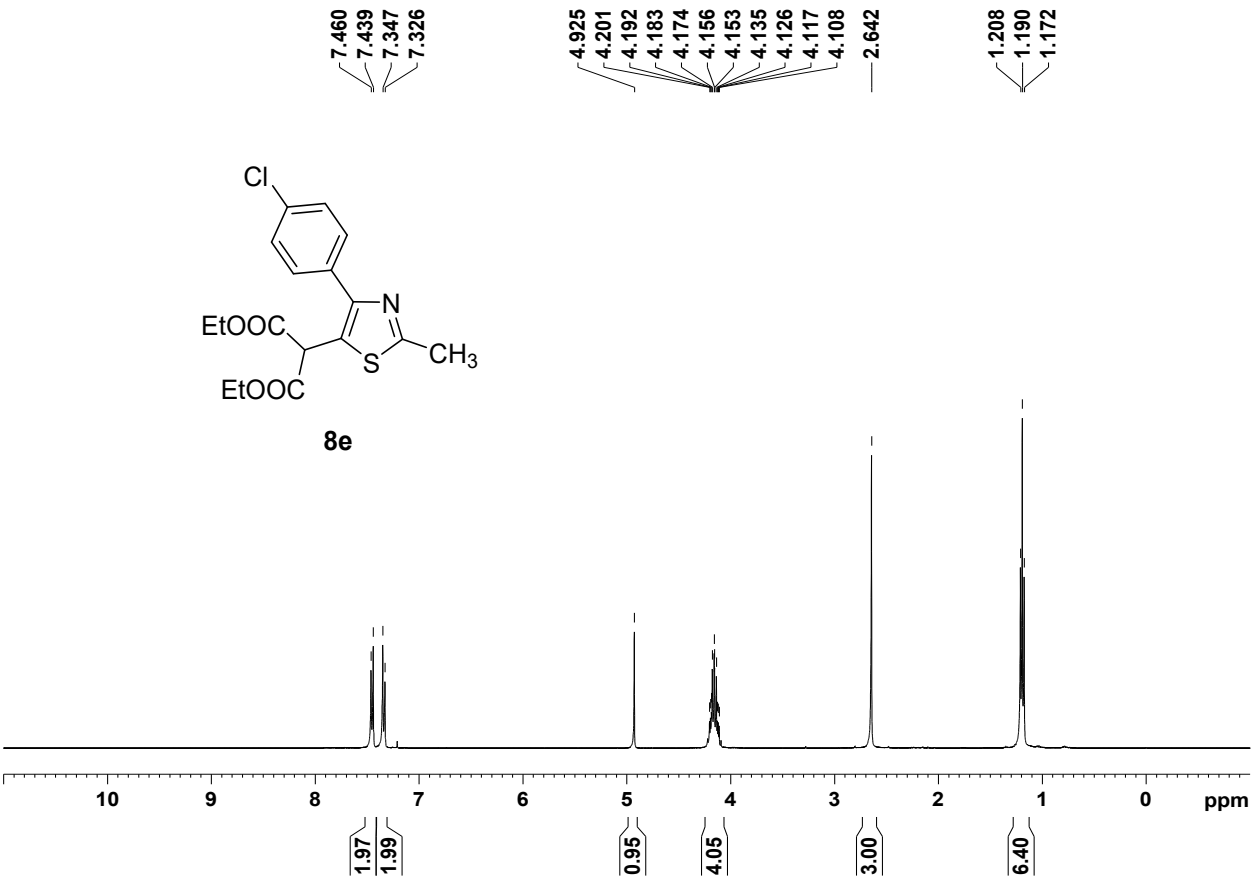

Figure 9. ^{1}H NMR (400 MHz, $CDCl_3$) spectrum of **8e**

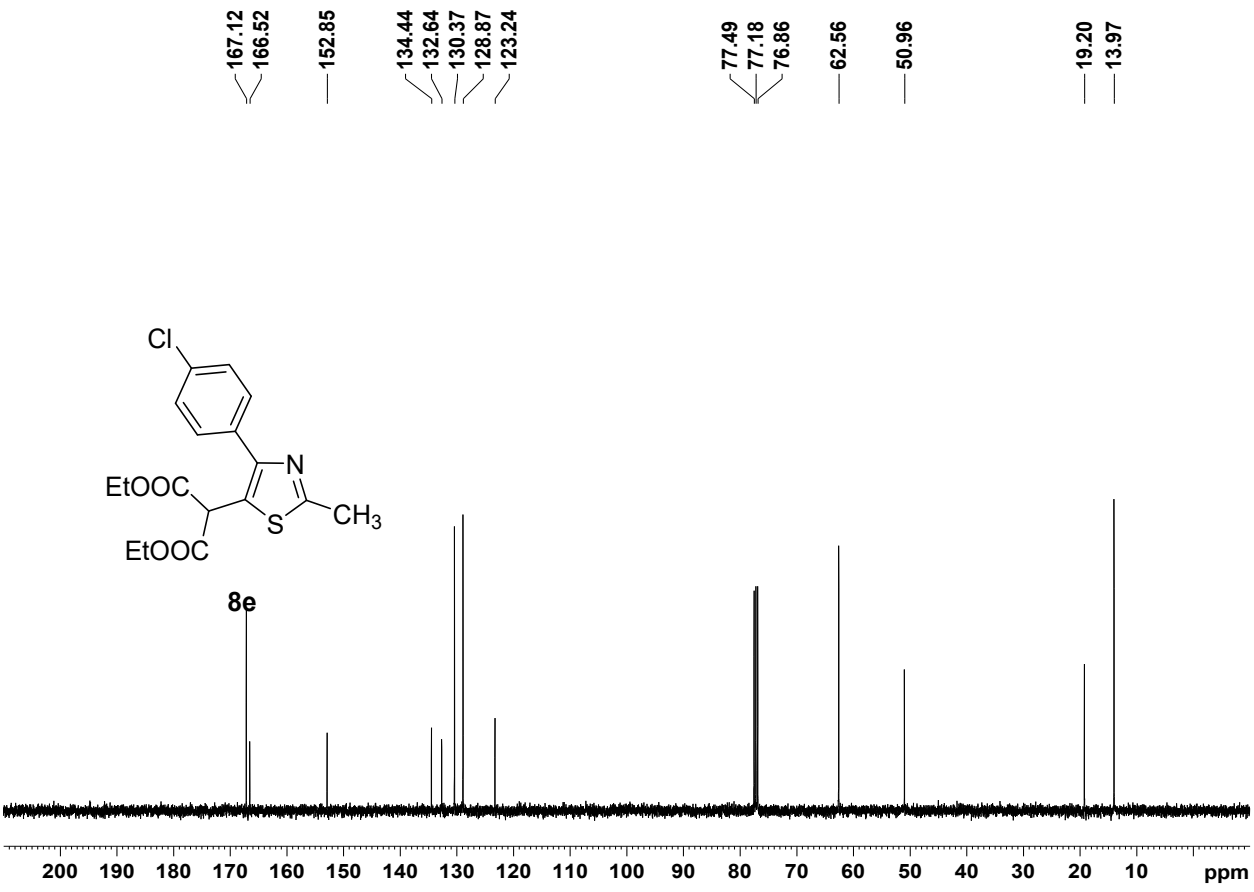

Figure 10. ^{13}C NMR (100 MHz, $CDCl_3$) spectrum of **8e**

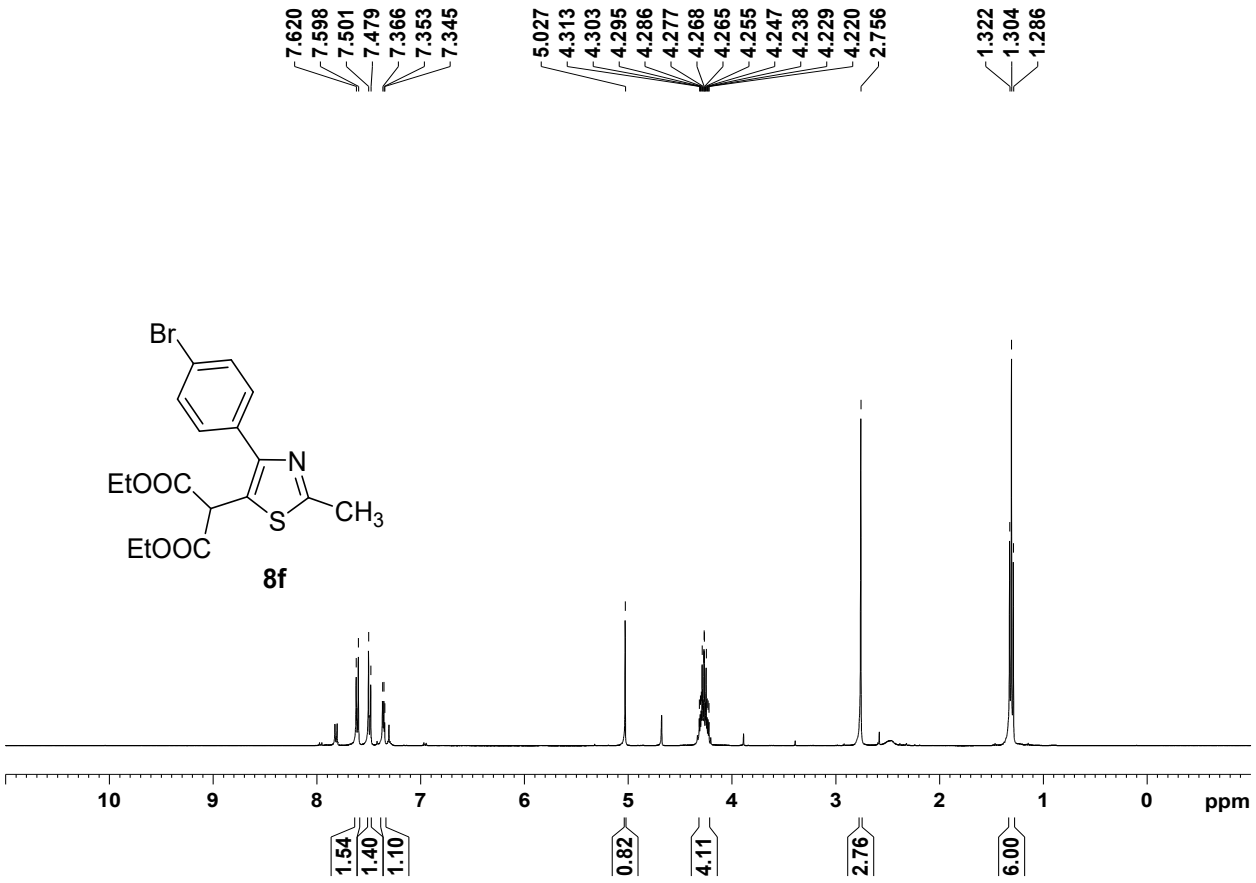

Figure 11. ^{1}H NMR (400 MHz, $CDCl_3$) spectrum of **8f**

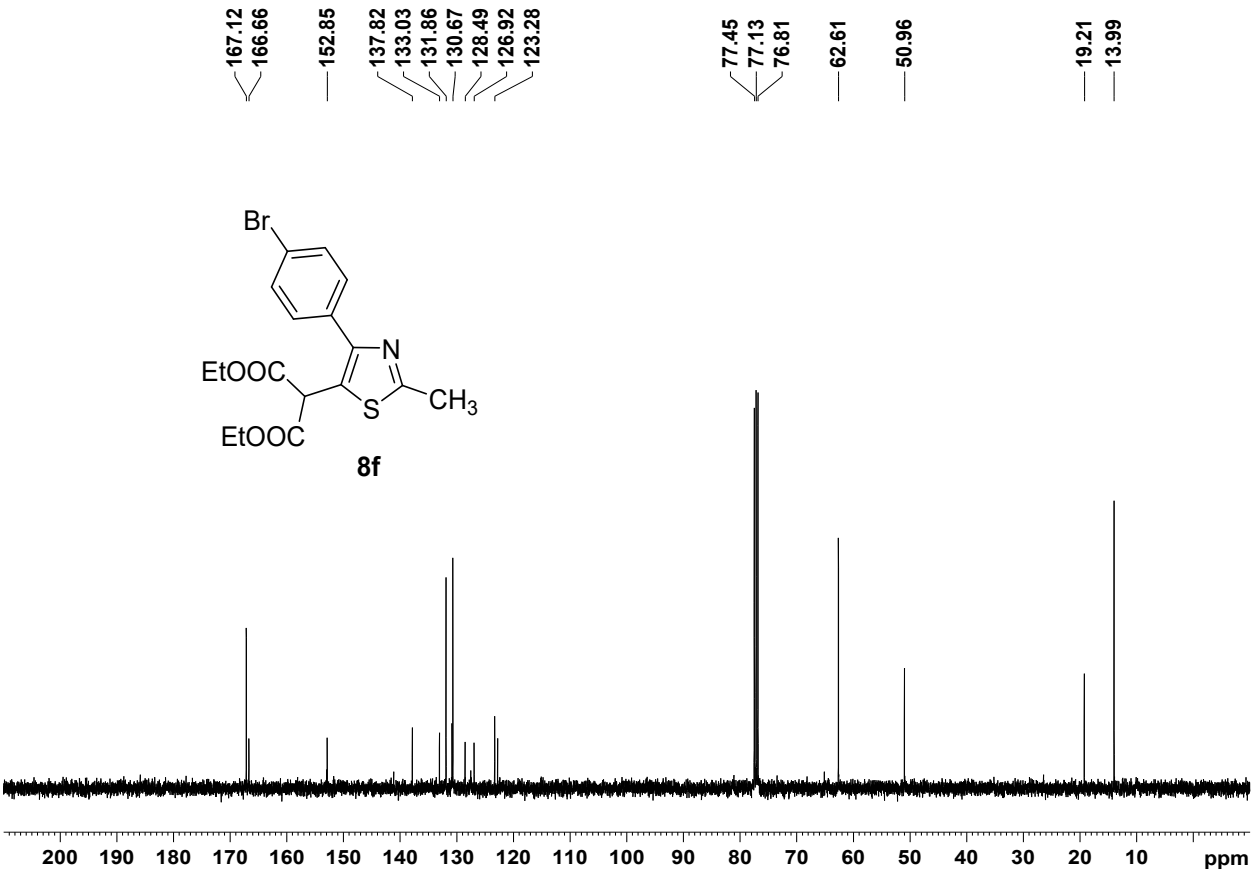

Figure 12. ^{13}C NMR (100 MHz, $CDCl_3$) spectrum of **8f**

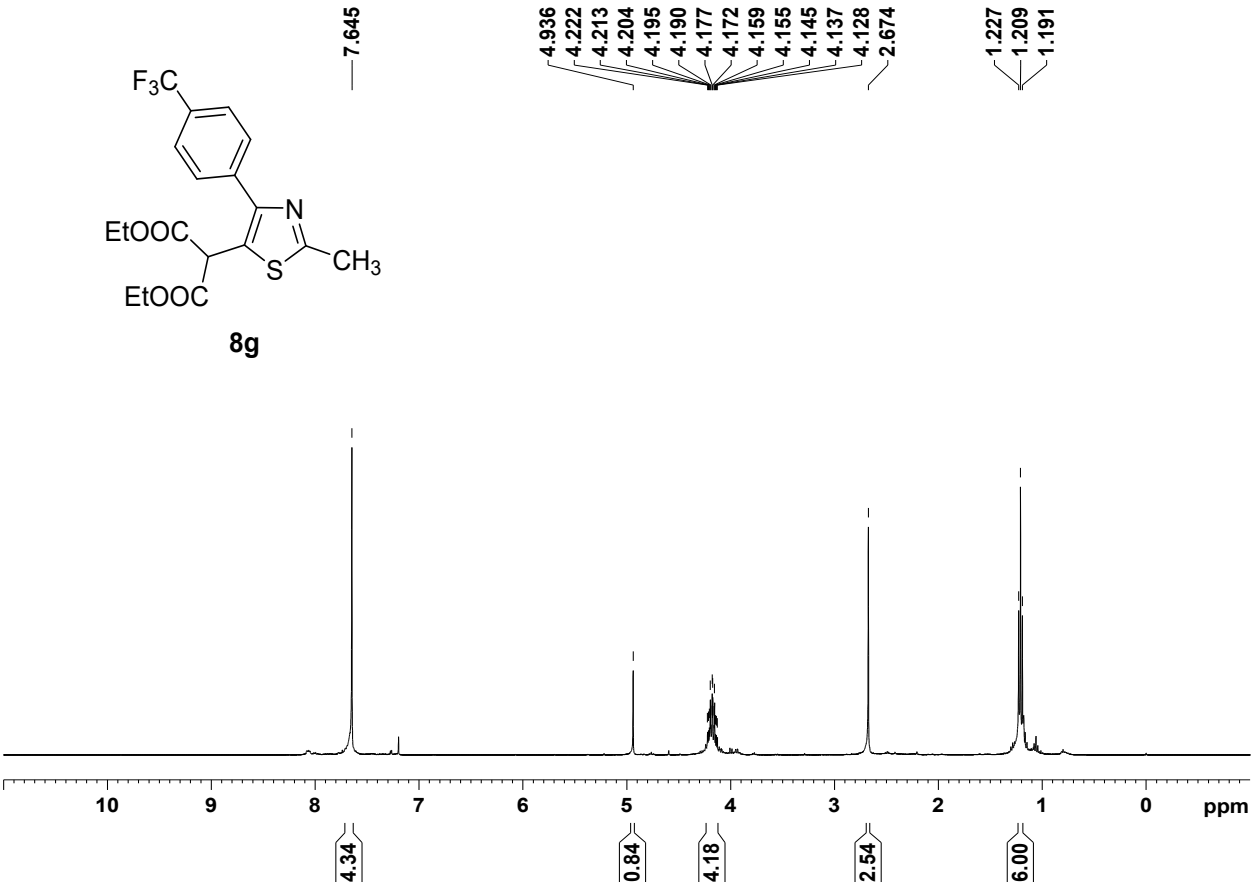

Figure 13. ^{1}H NMR (400 MHz, $CDCl_3$) spectrum of **8g**

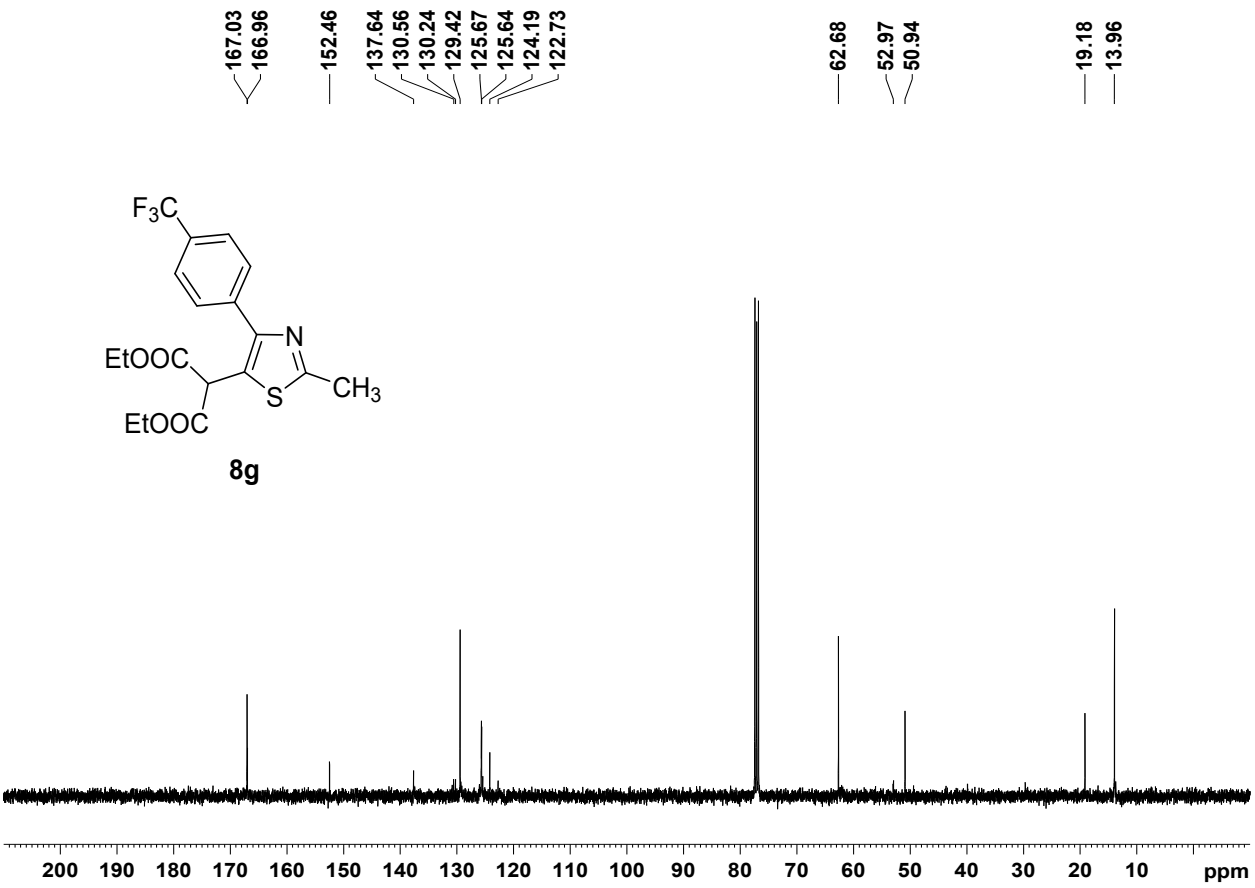

Figure 14. ^{13}C NMR (100 MHz, $CDCl_3$) spectrum of **8g**

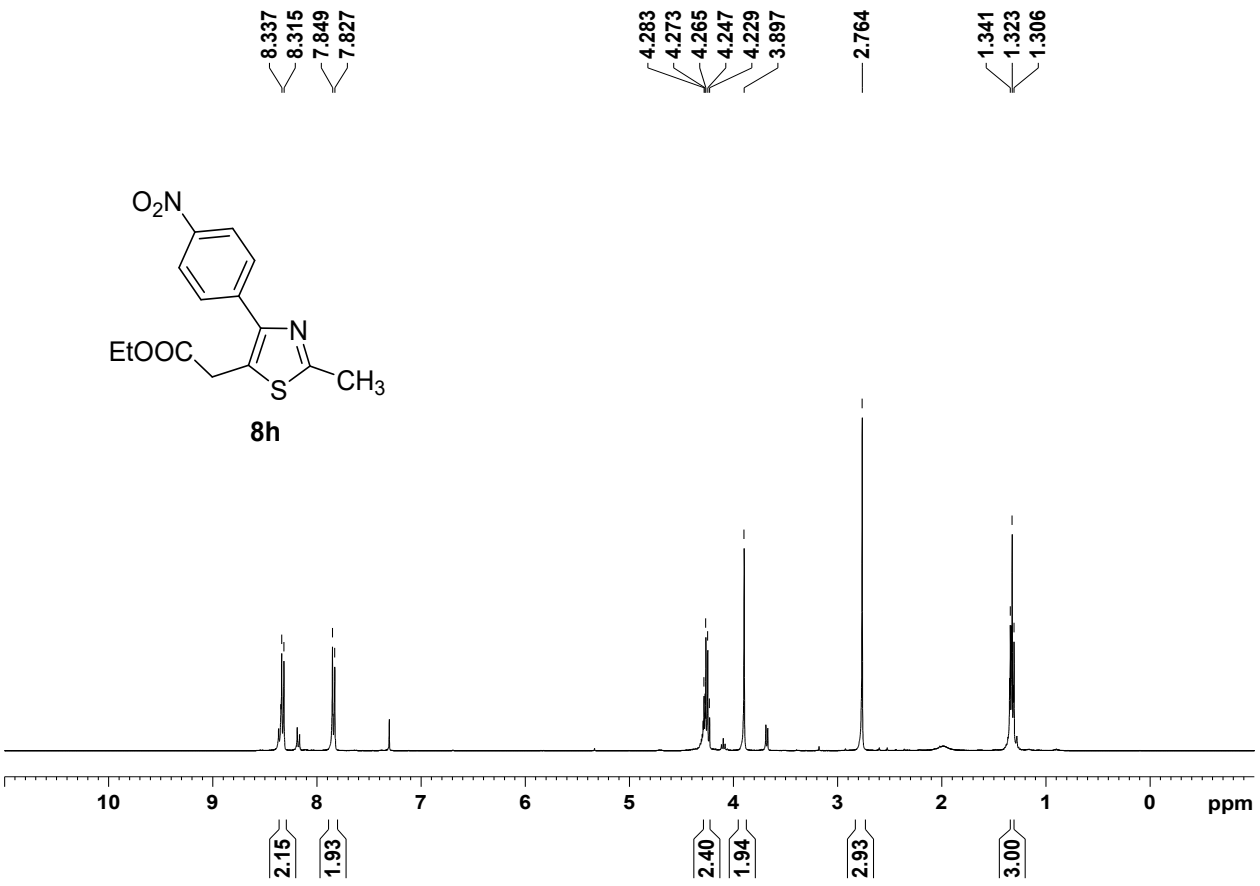

Figure 15. ^{1}H NMR (400 MHz, $CDCl_3$) spectrum of **8h**

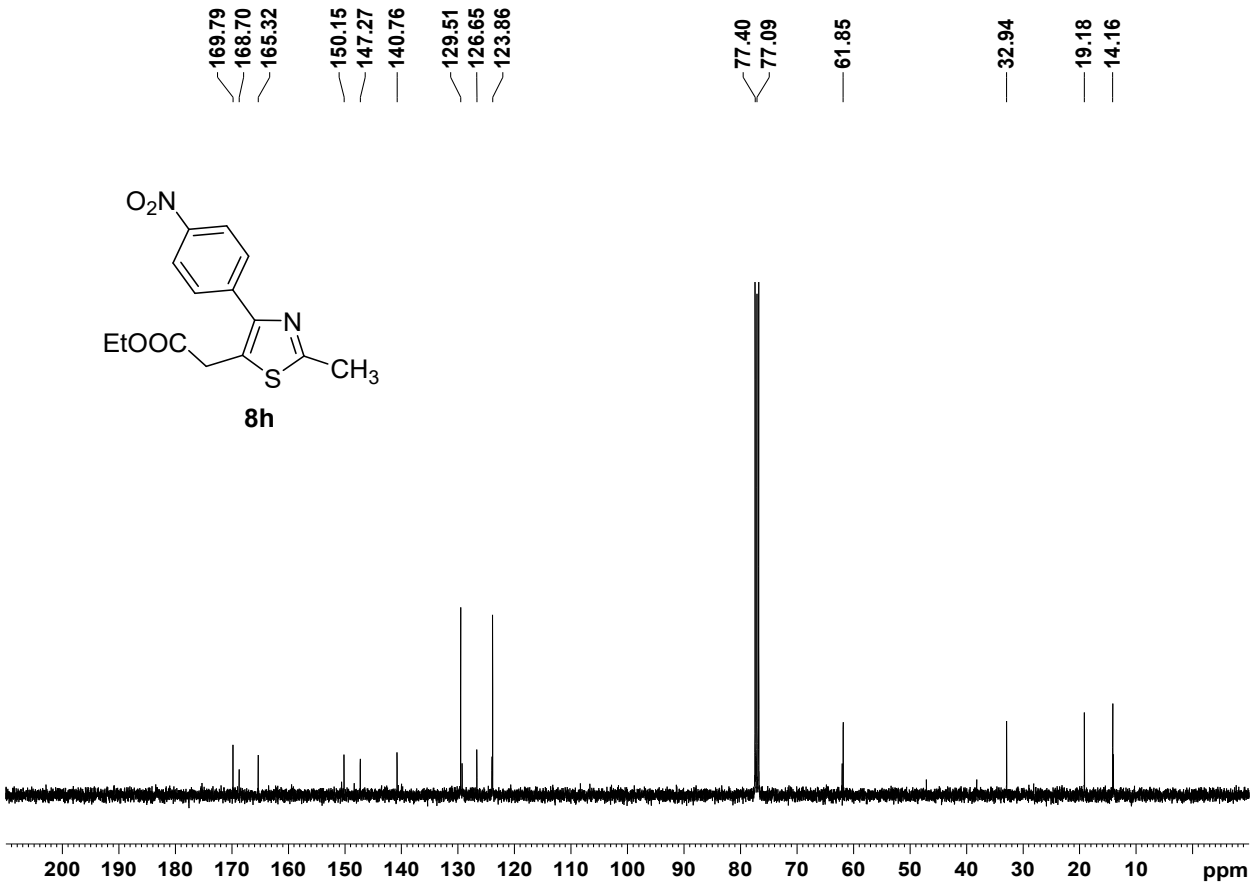

Figure 16. ^{13}C NMR (100 MHz, $CDCl_3$) spectrum of **8h**

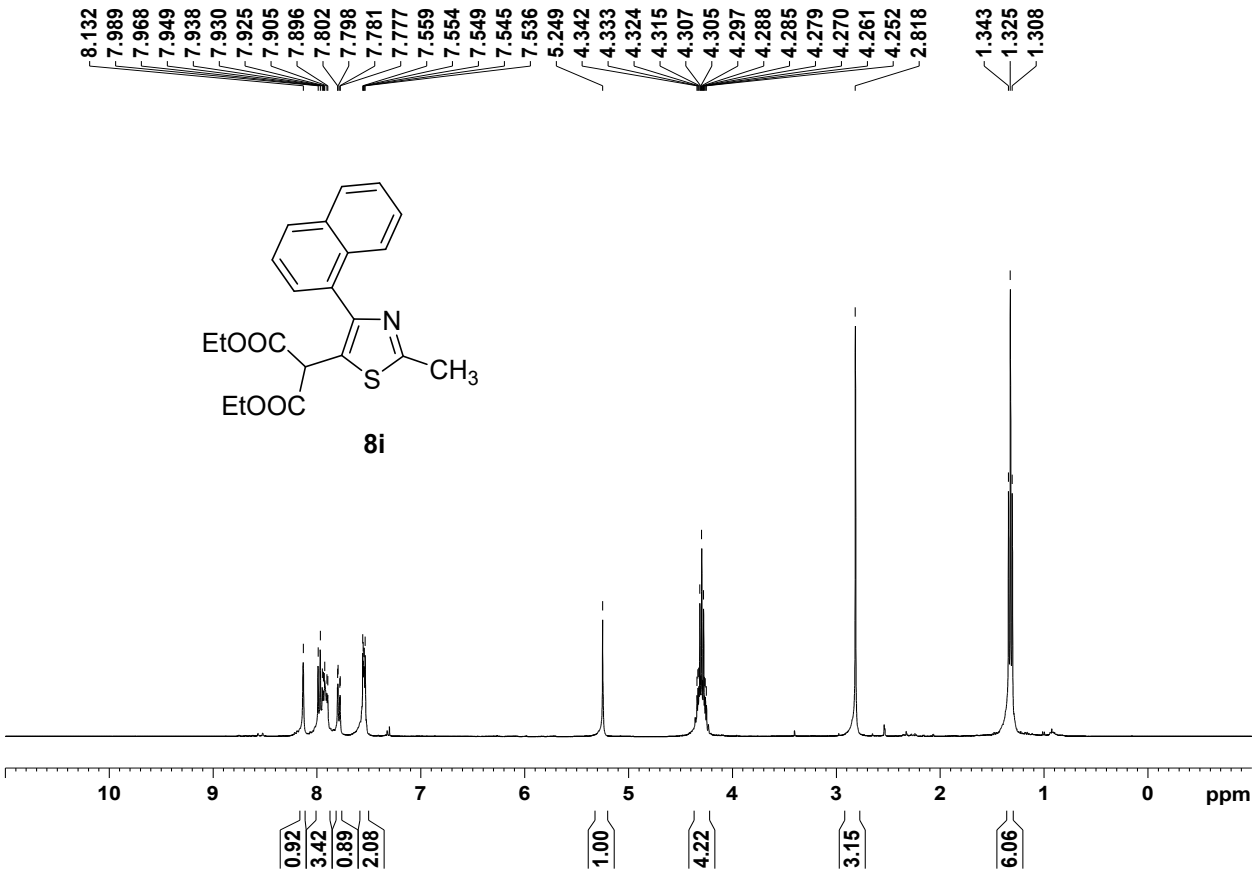

Figure 17. ^{1}H NMR (400 MHz, $CDCl_3$) spectrum of **8i**

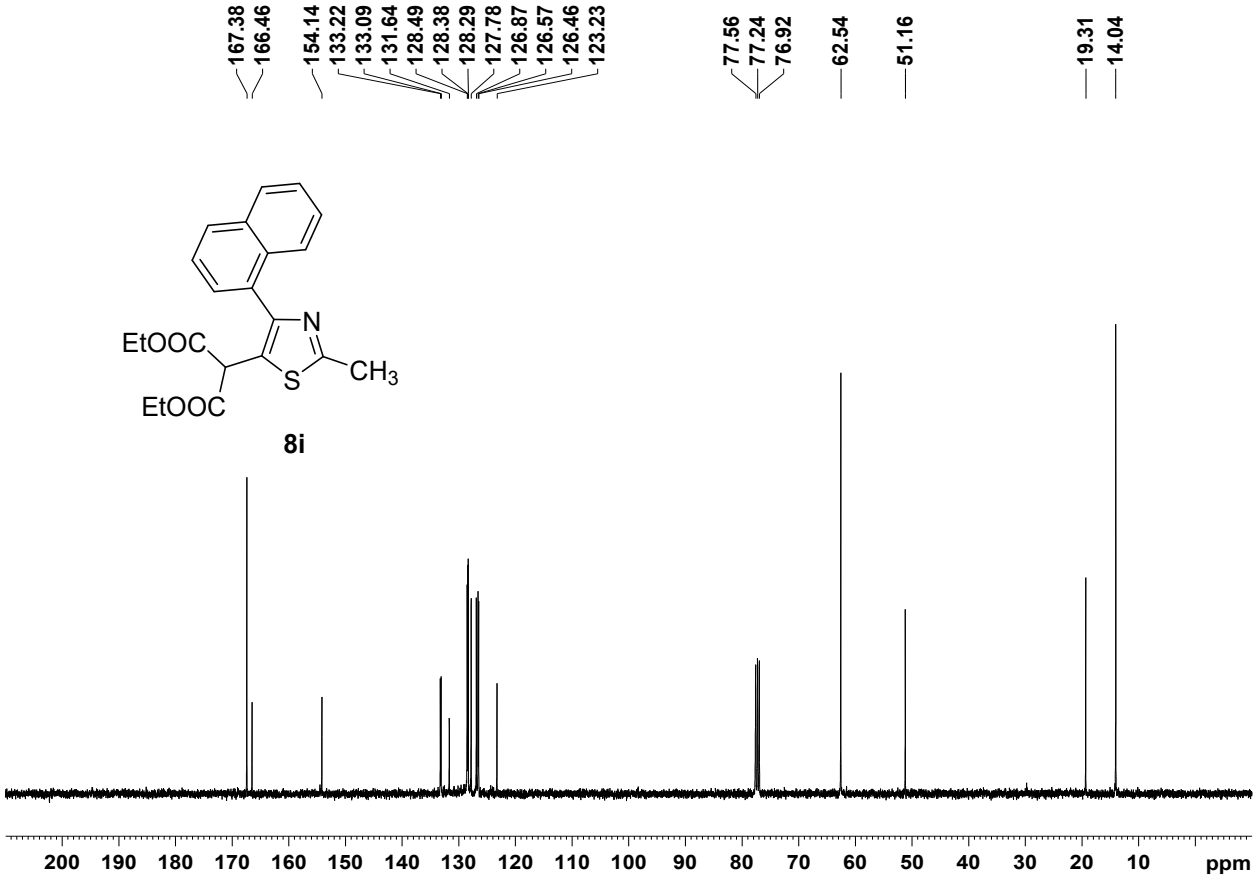

Figure 18. ^{13}C NMR (100 MHz, $CDCl_3$) spectrum of **8i**

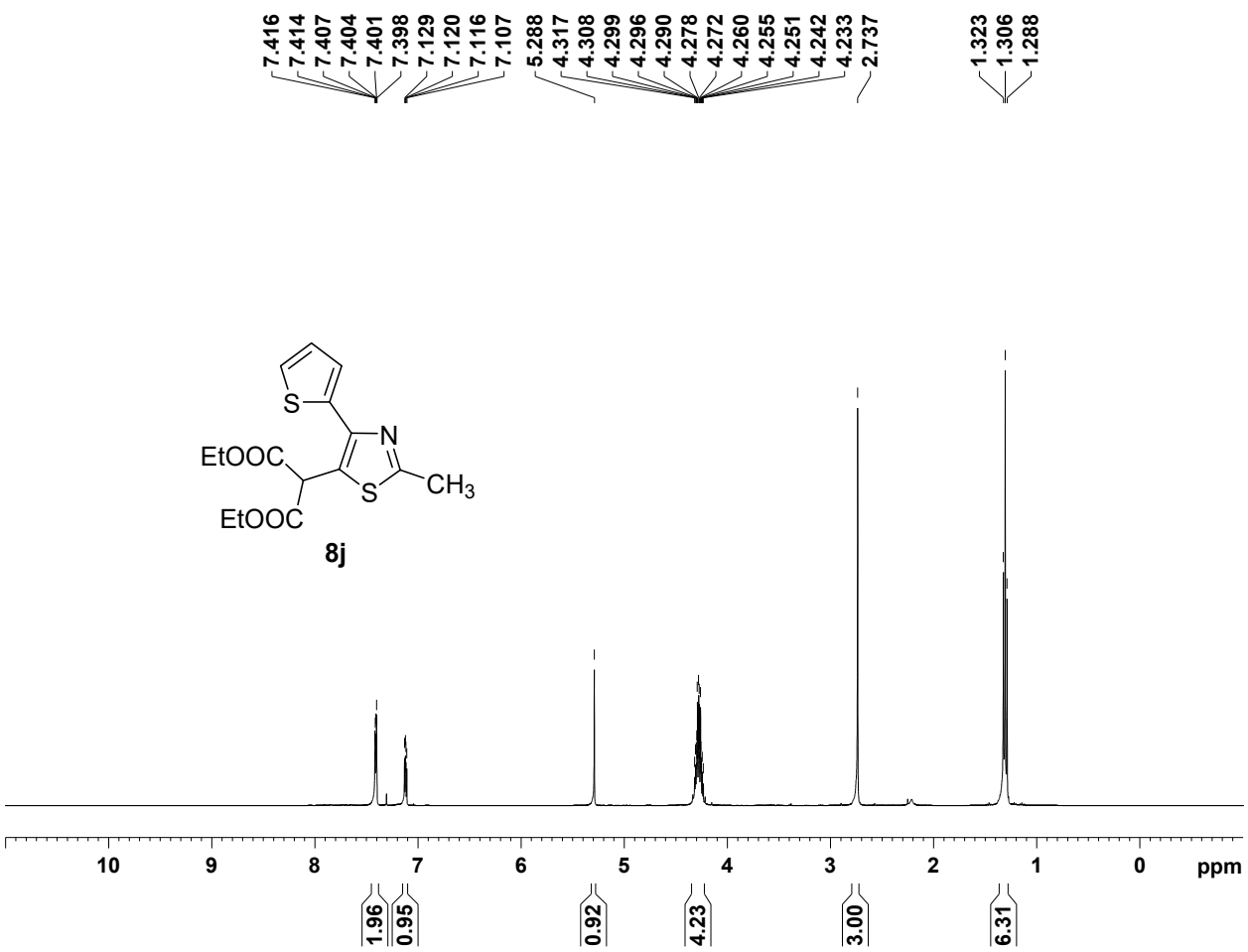

Figure 19. ^{1}H NMR (400 MHz, $CDCl_3$) spectrum of **8j**

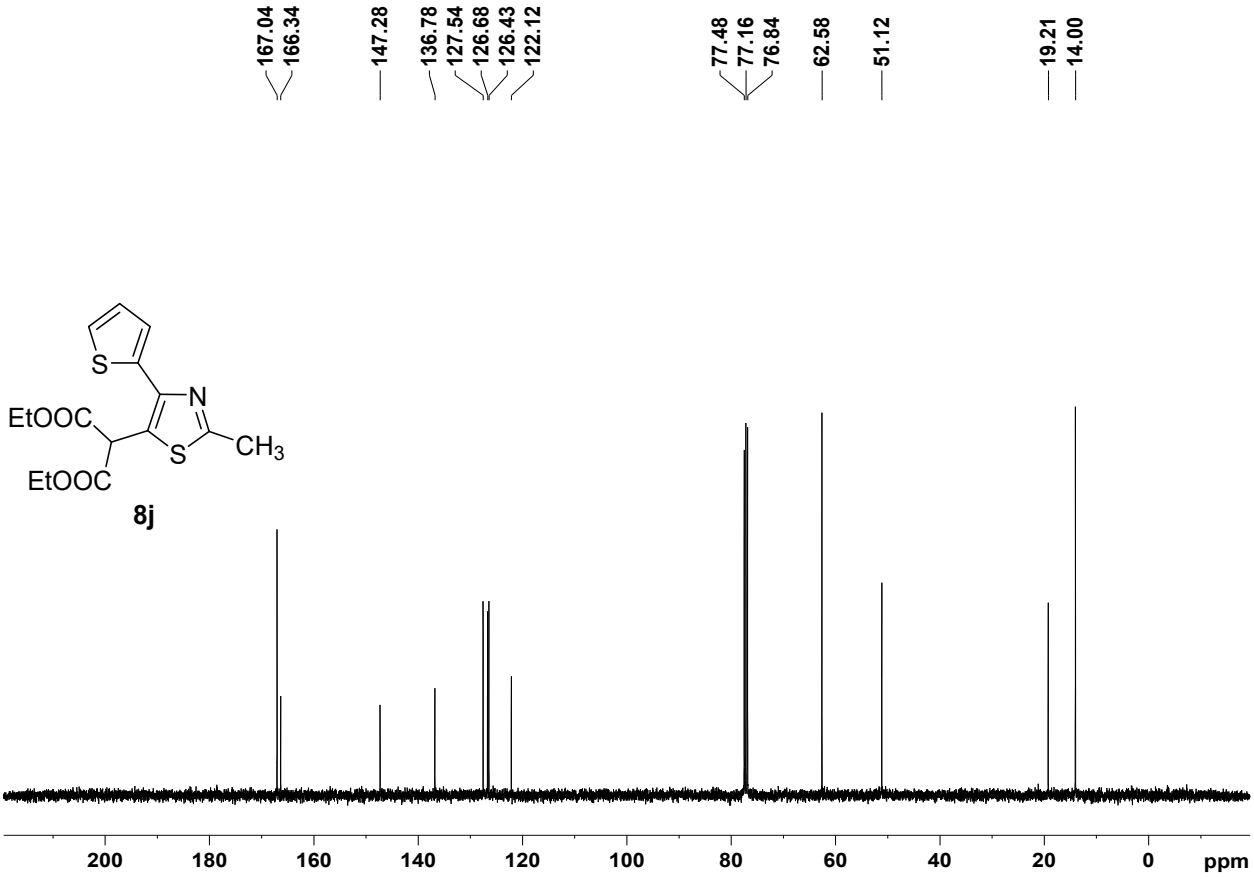

Figure 20. ^{13}C NMR (100 MHz, $CDCl_3$) spectrum of **8j**

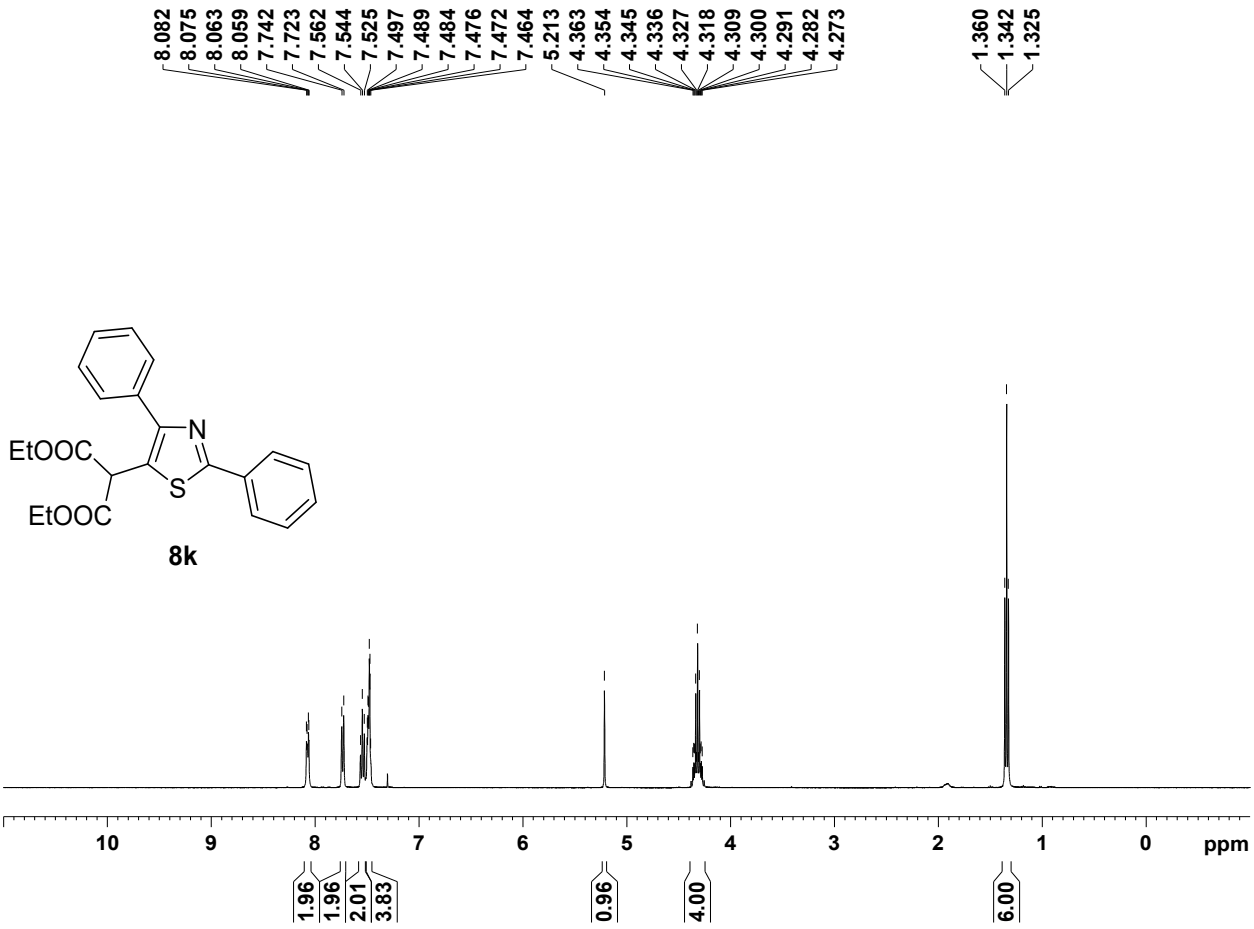

Figure 21. ^{1}H NMR (400 MHz, $CDCl_3$) spectrum of **8k**

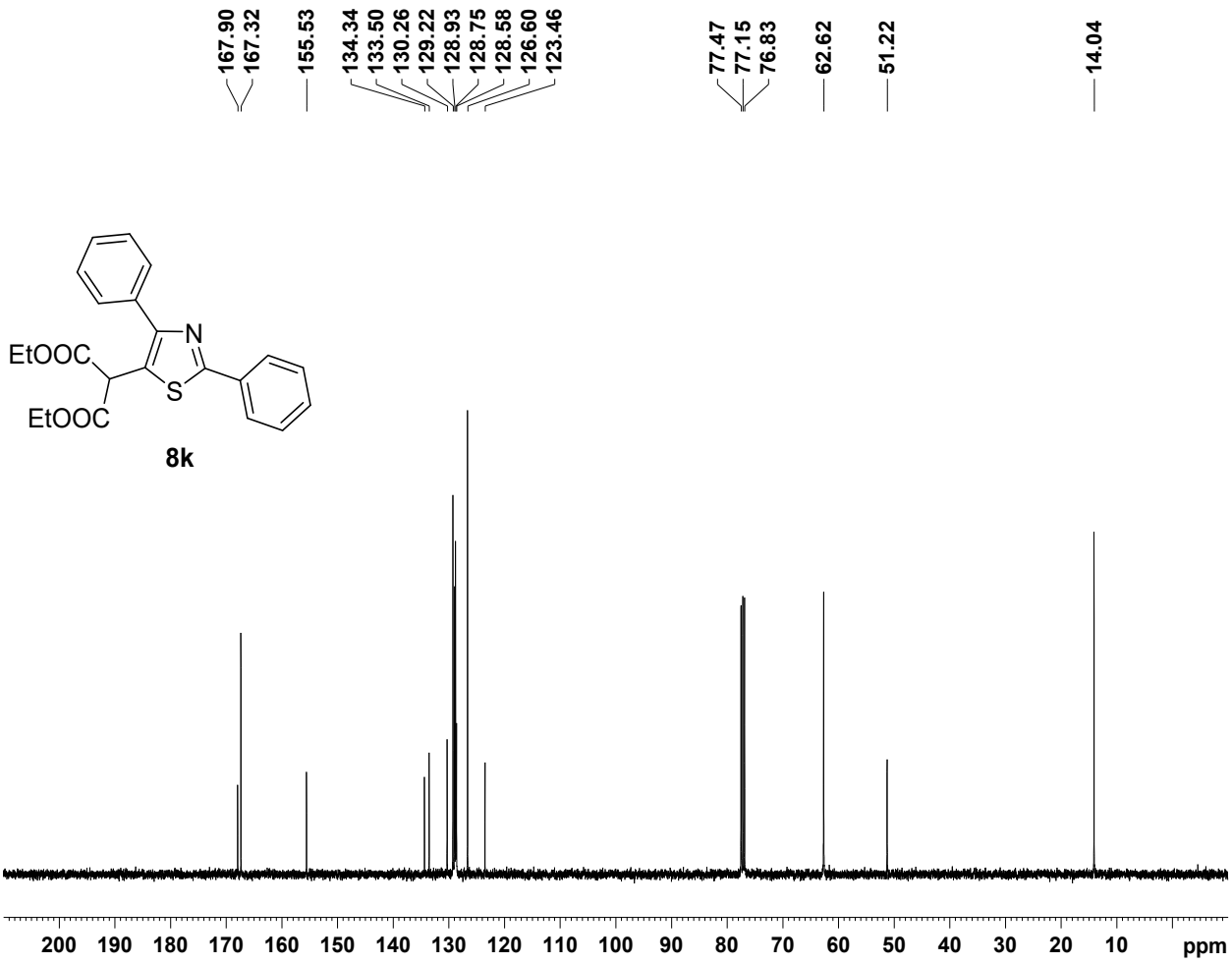

Figure 22. ^{13}C NMR (100 MHz, $CDCl_3$) spectrum of **8k**

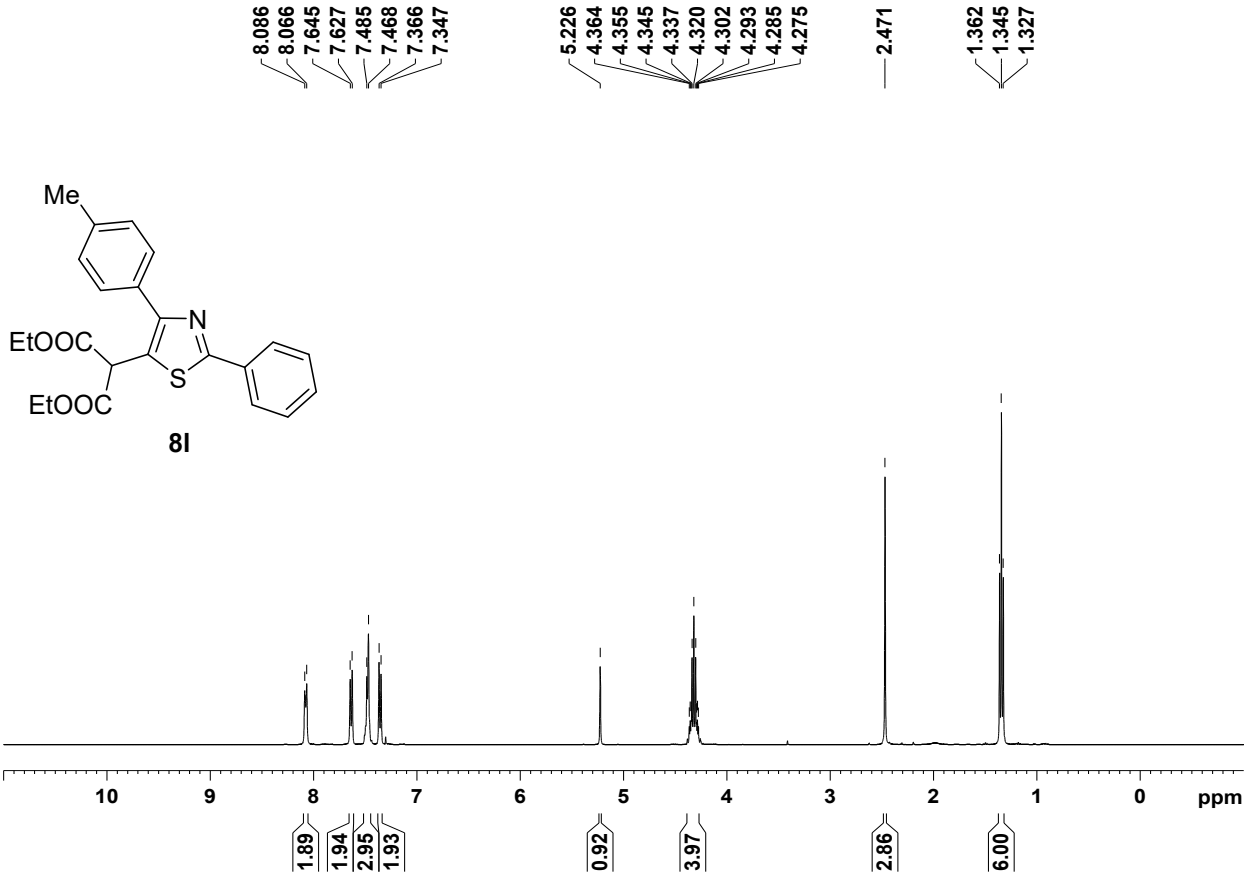

Figure 23. ^{1}H NMR (400 MHz, $CDCl_3$) spectrum of **8l**

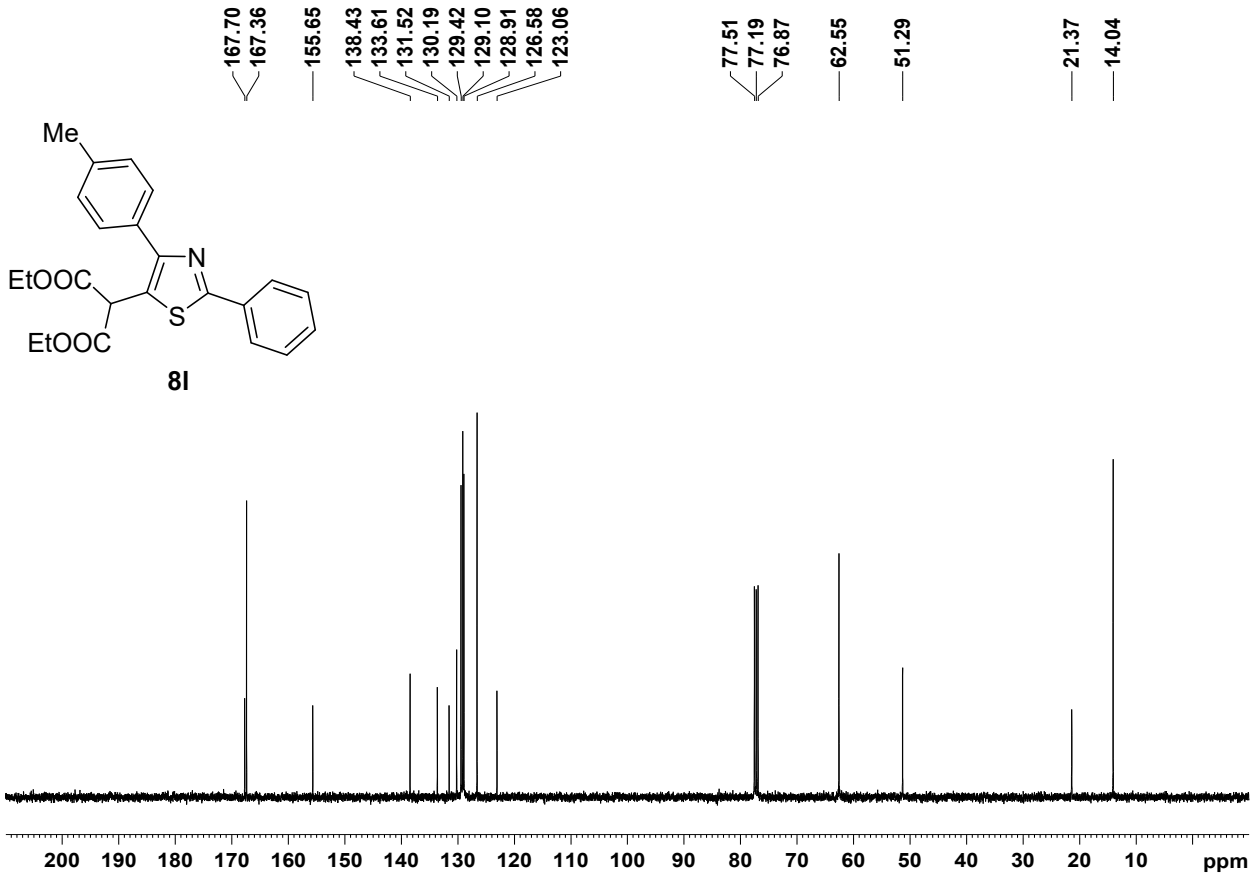

Figure 24. ^{13}C NMR (100 MHz, $CDCl_3$) spectrum of **8l**

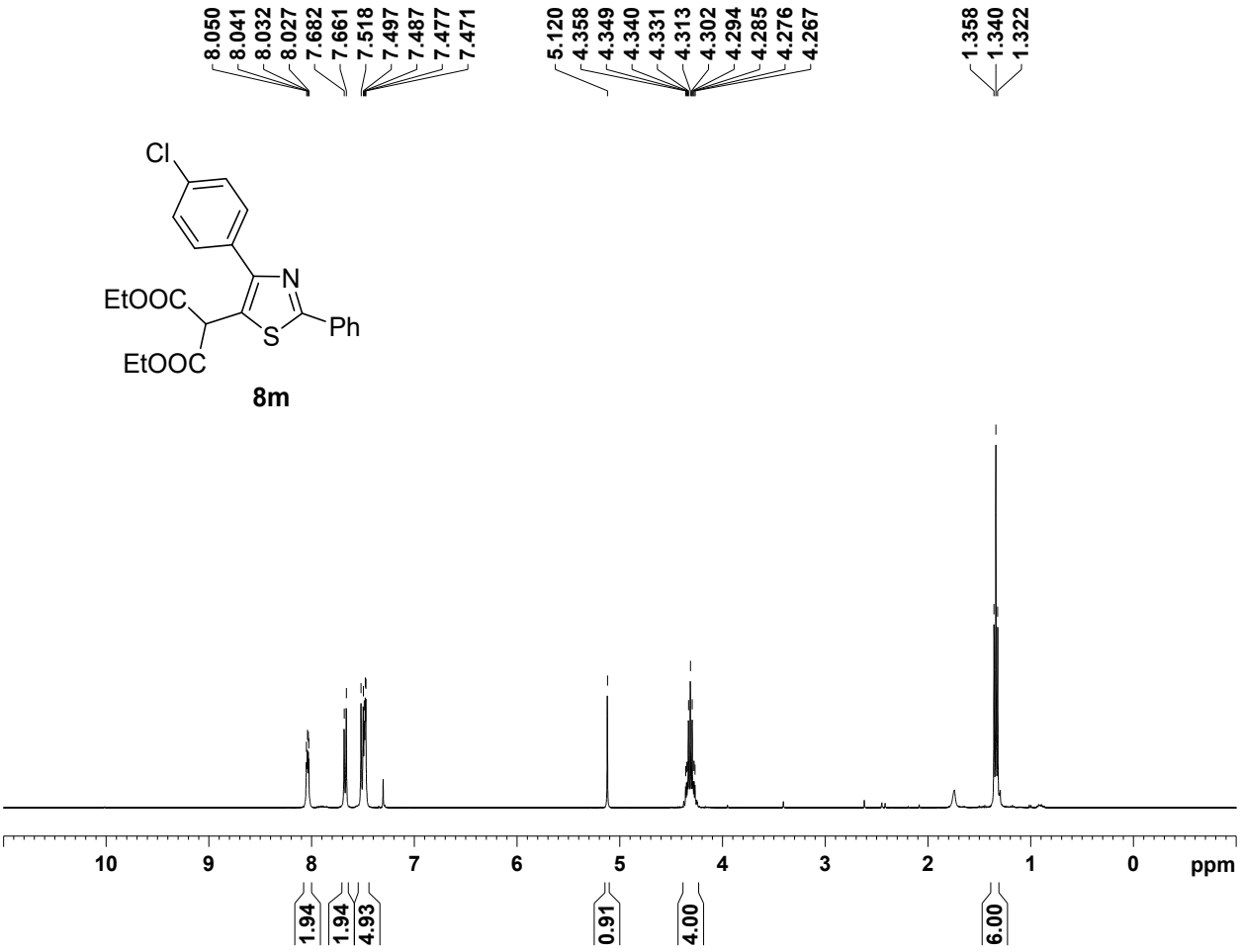

Figure 25. ^{1}H NMR (400 MHz, $CDCl_3$) spectrum of **8m**

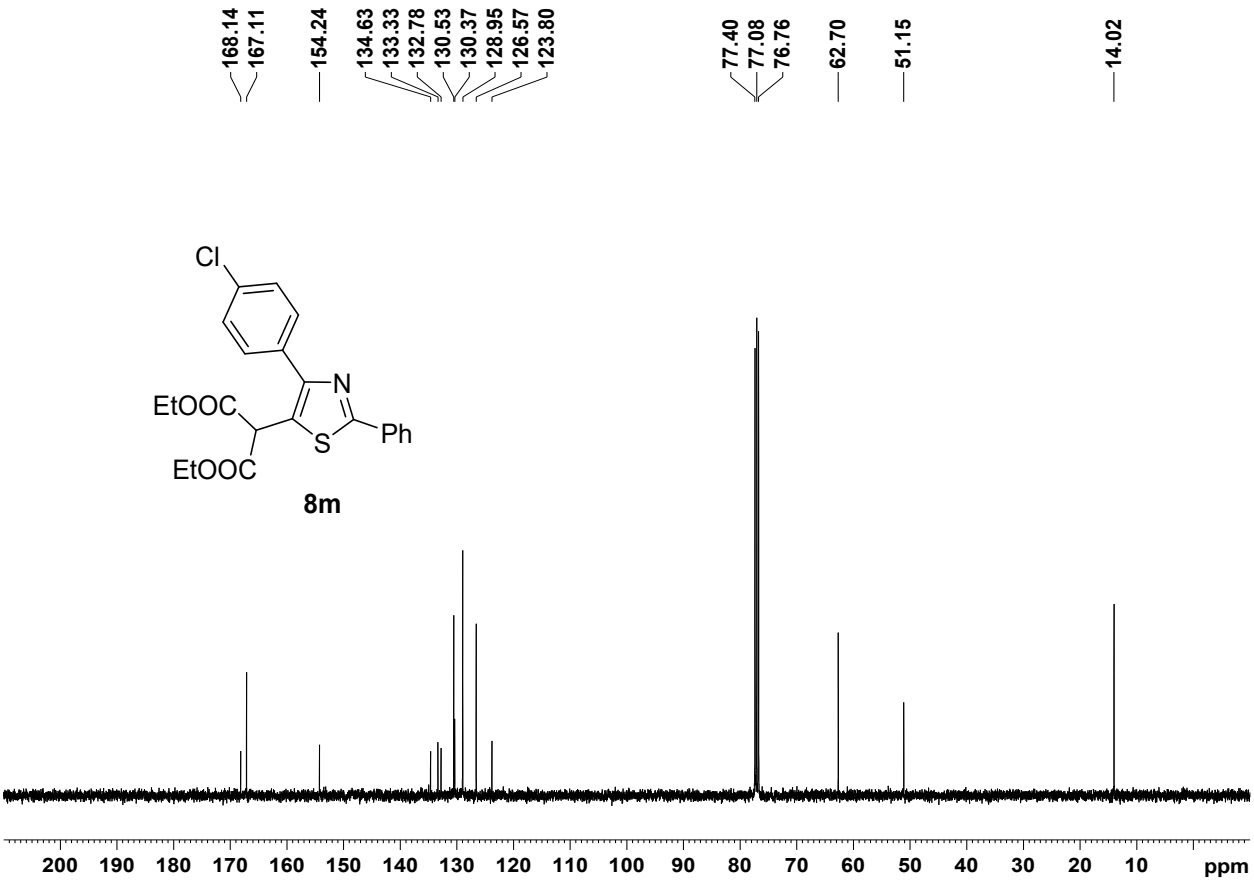

Figure 26. ^{13}C NMR (100 MHz, $CDCl_3$) spectrum of **8m**

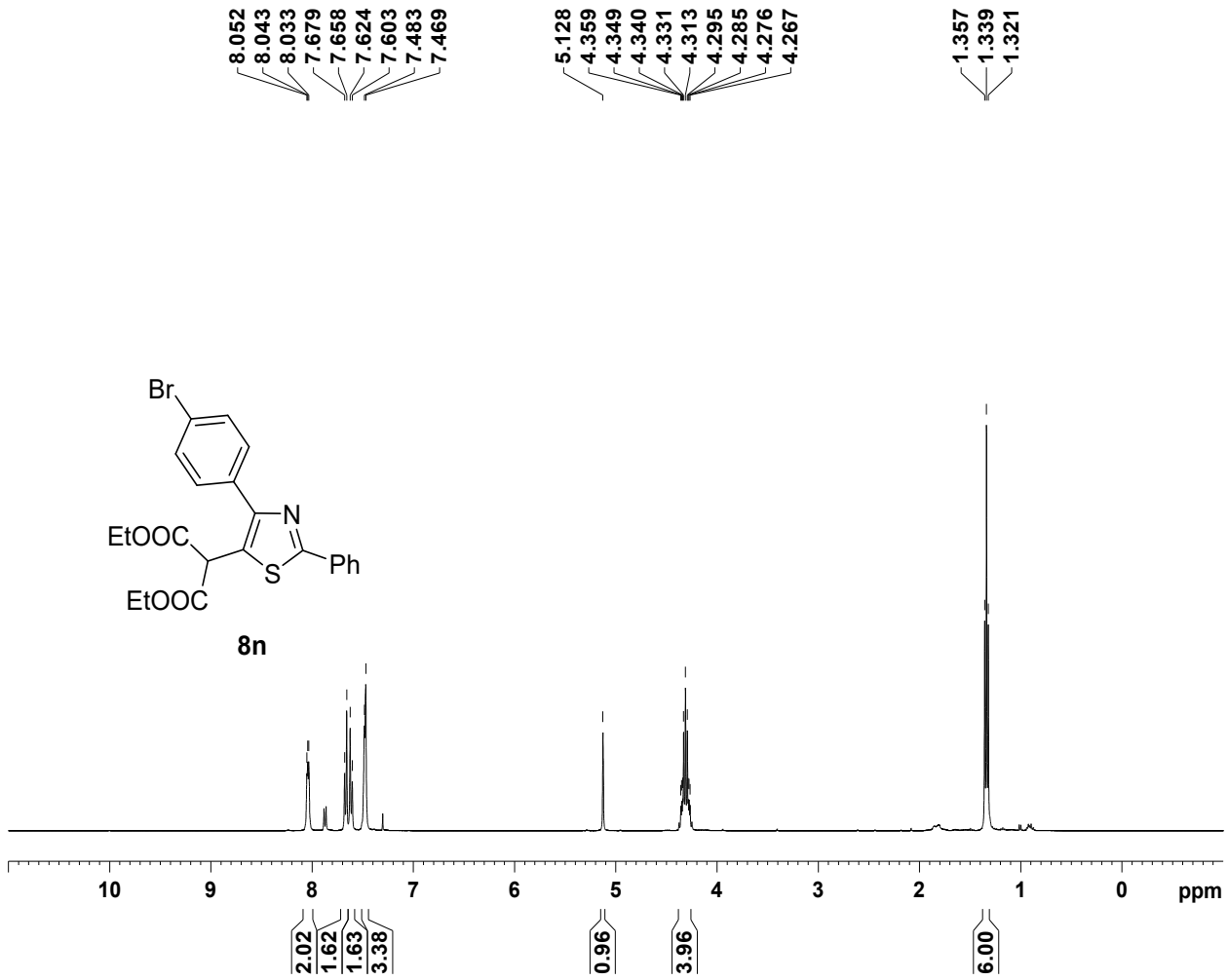

Figure 27. ^{1}H NMR (400 MHz, $CDCl_3$) spectrum of **8n**

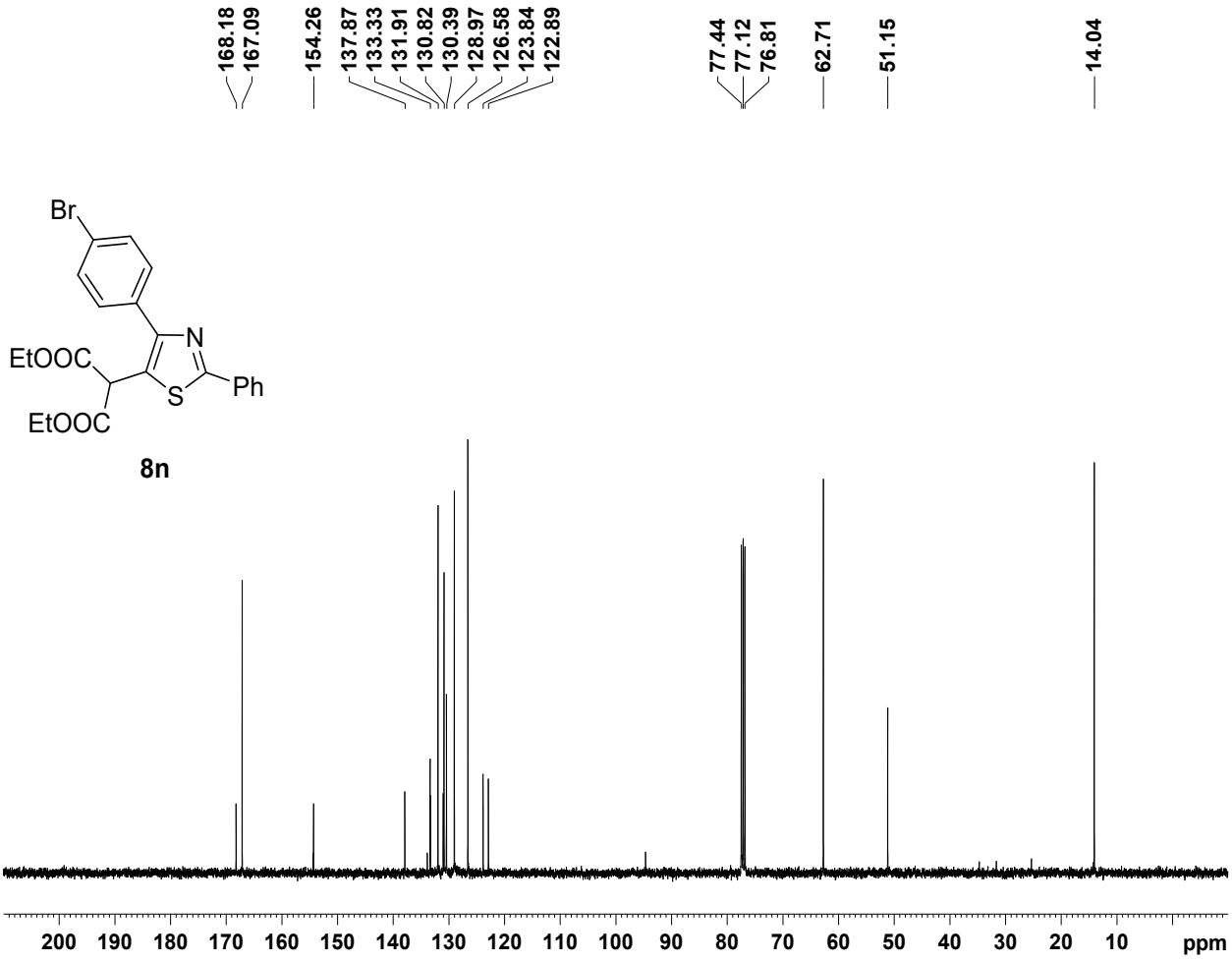

Figure 28. ^{13}C NMR (100 MHz, $CDCl_3$) spectrum of **8n**

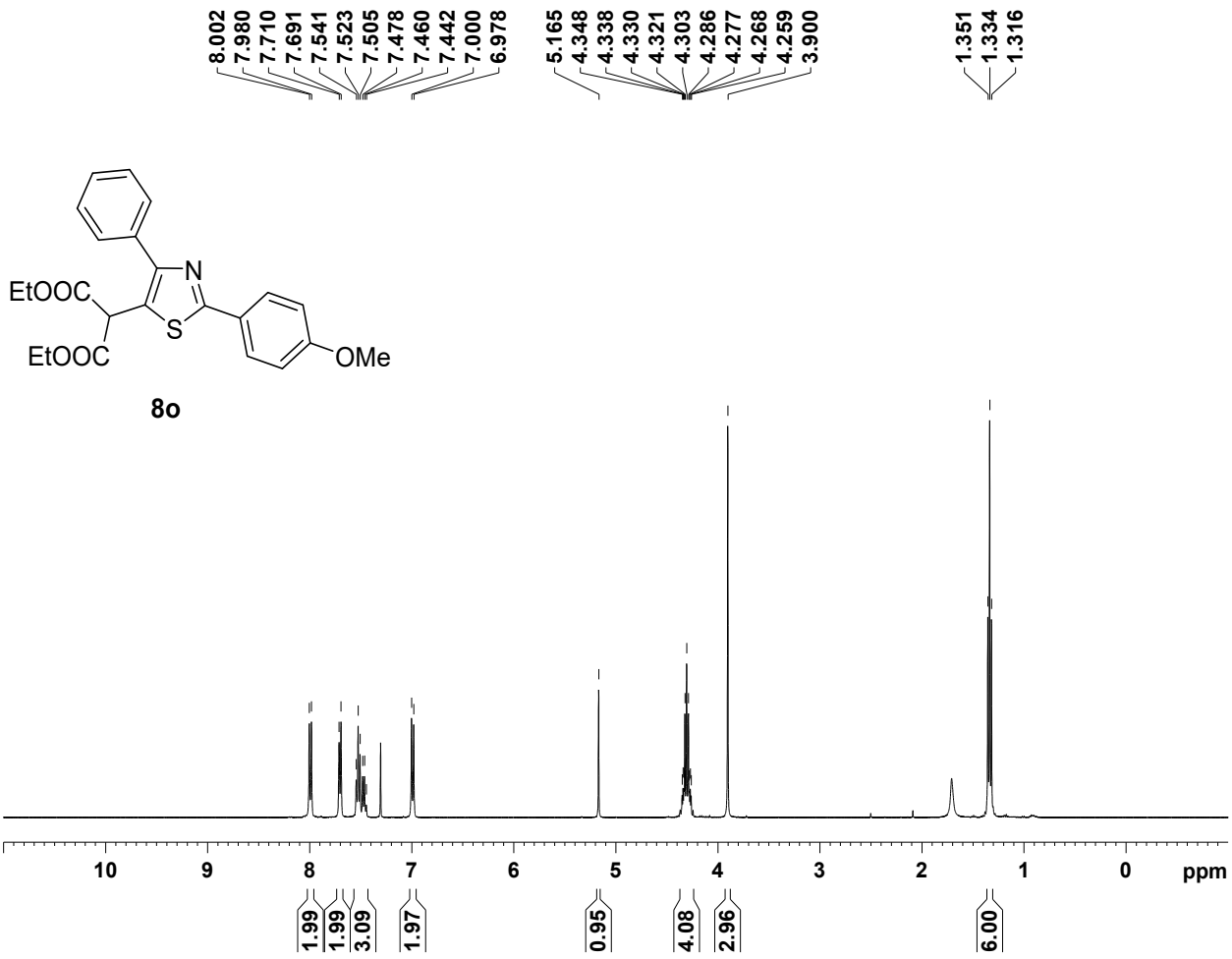

Figure 29. ^{1}H NMR (400 MHz, $CDCl_3$) spectrum of **8o**

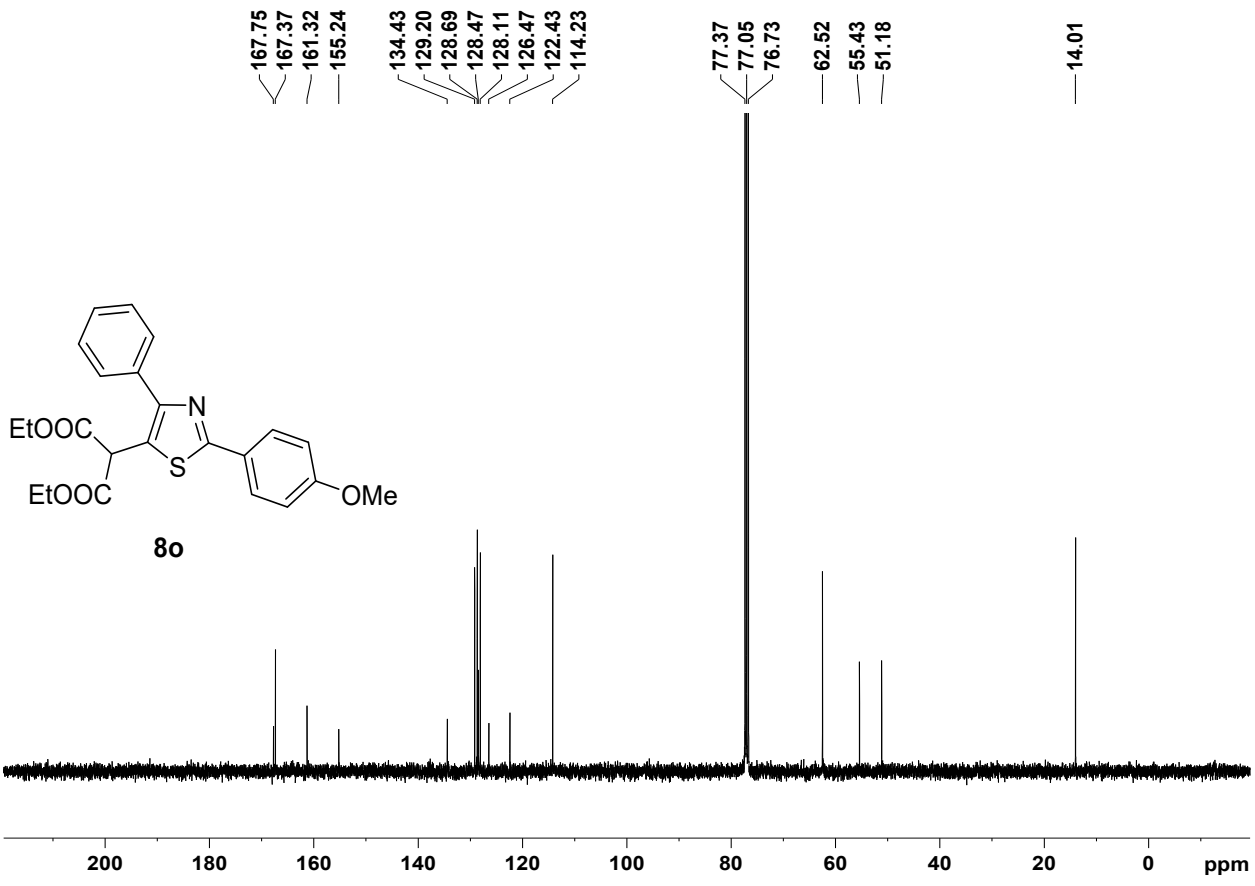

Figure 30. ^{13}C NMR (100 MHz, $CDCl_3$) spectrum of **8o**

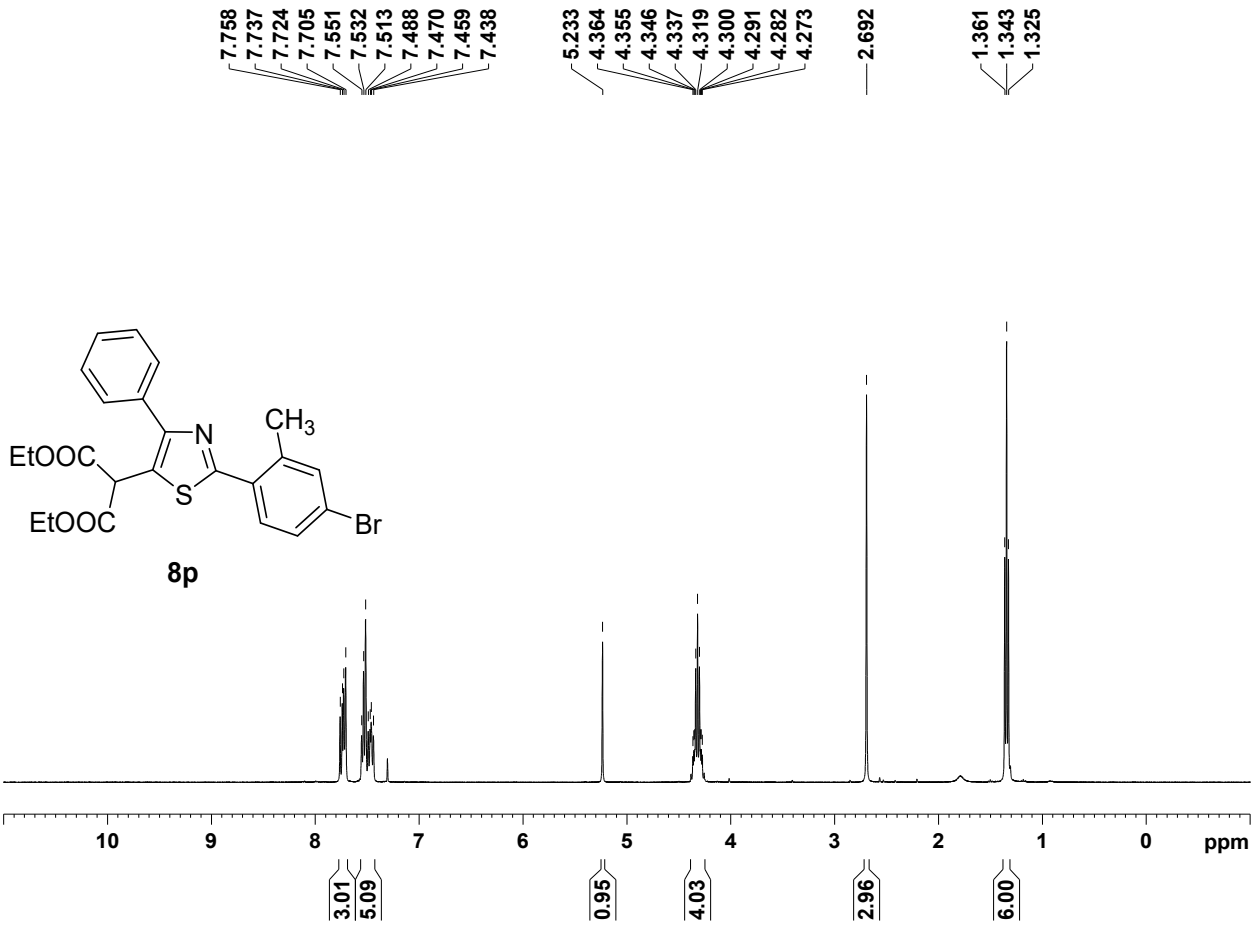

Figure 31. ^{1}H NMR (400 MHz, $CDCl_3$) spectrum of **8p**

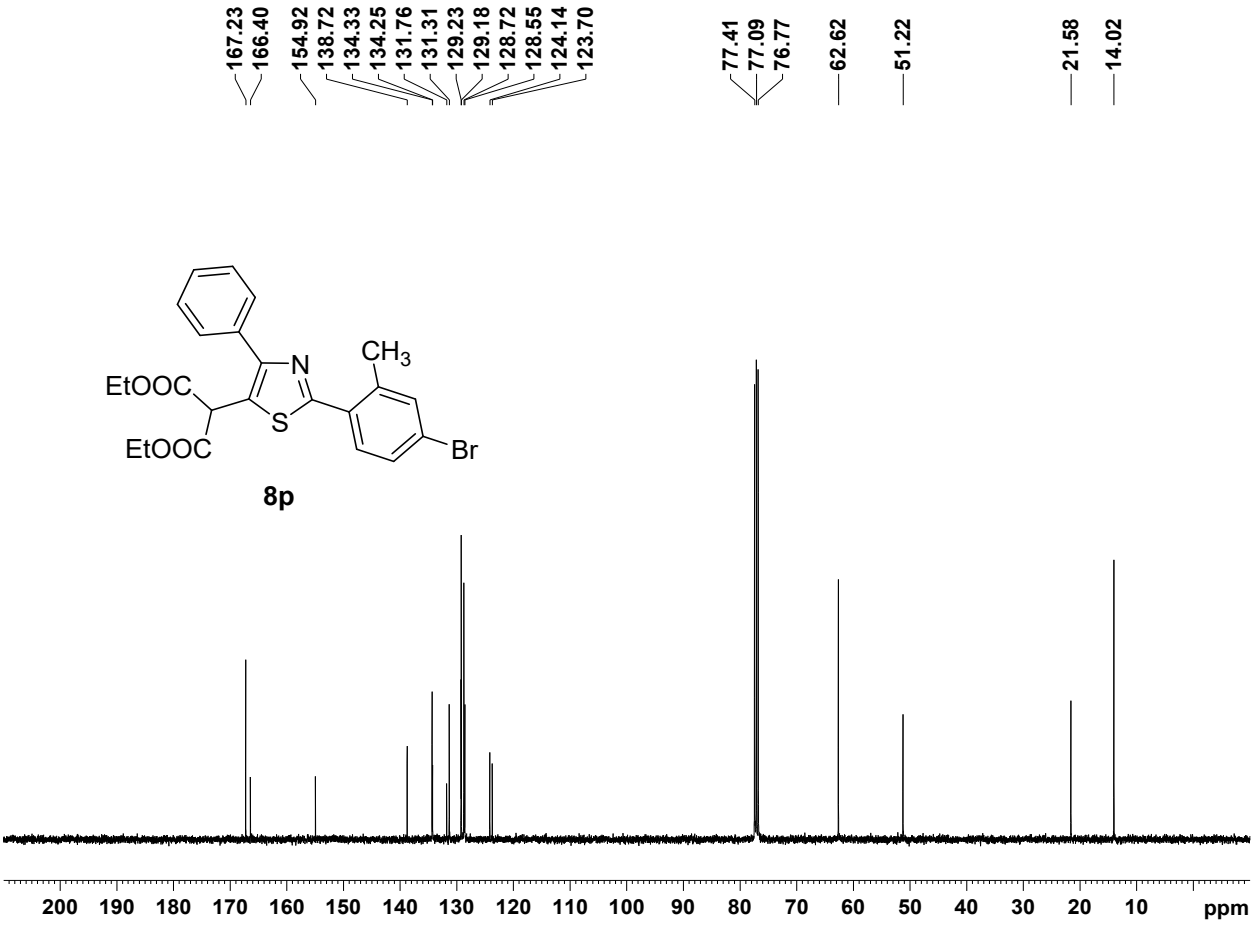

Figure 32. ^{13}C NMR (100 MHz, $CDCl_3$) spectrum of **8p**

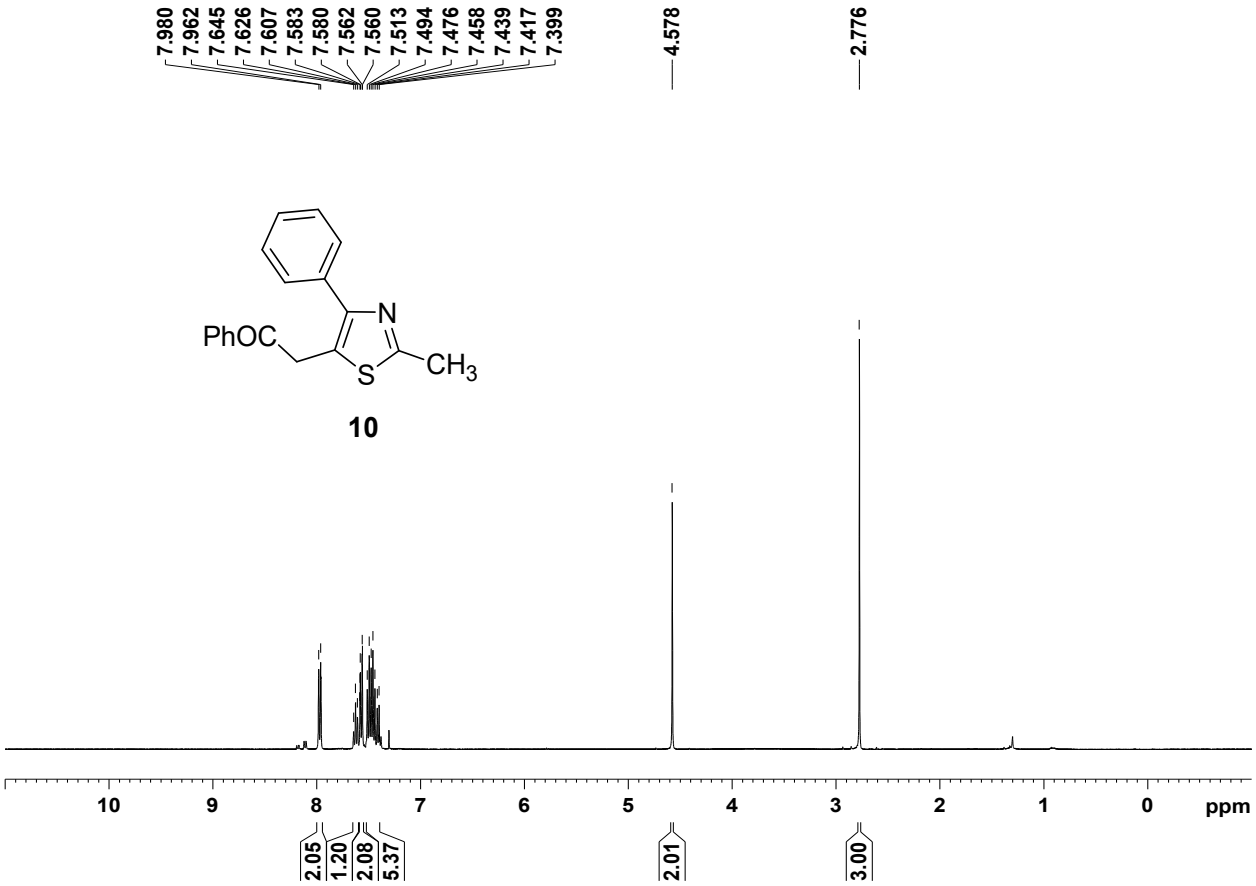

Figure 33. ^{1}H NMR (400 MHz, $CDCl_3$) spectrum of **10**

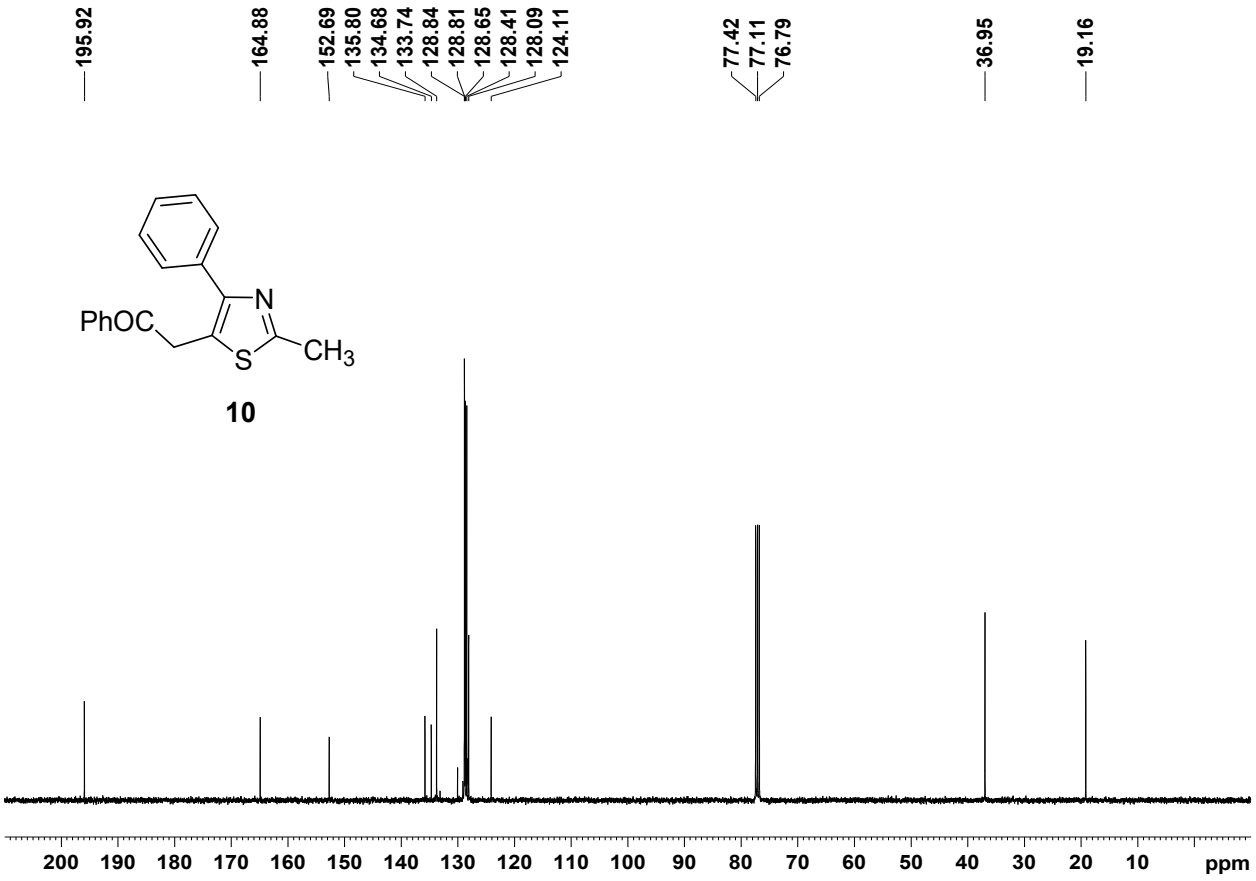

Figure 34. ^{13}C NMR (100 MHz, $CDCl_3$) spectrum of **10**

B. X-ray structure of 8o

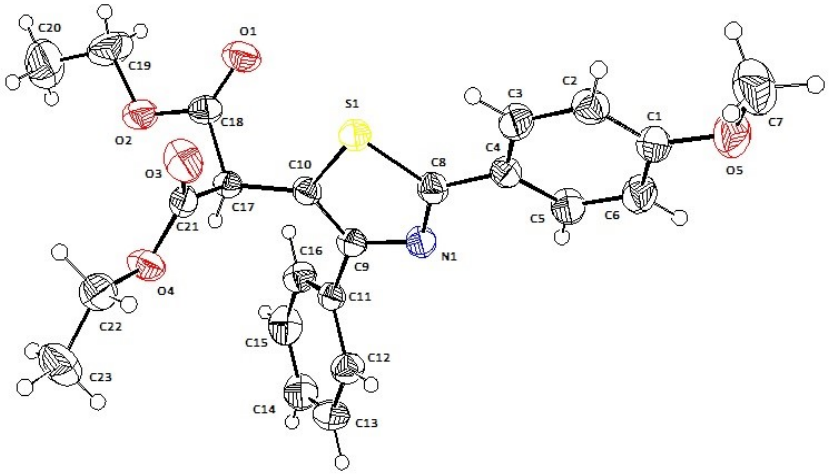

Figure 1. ORTEP plot of the crystal structure of **8o** (at 40% probability level)

Crystal structure determination: Single crystals suitable for X-ray studies were grown by recrystallization of **8o** from DCM/chloroform (1:9). X-ray data were collected on a CCD diffractometer using graphite-monochromated Mo-Kα radiation.

Table 1. Selected crystal parameters and refinement metrics for **8o**

CCDC Number for **8o**	2201951
Chemical formula	$C_{23}H_{23}NO_5S$
Formula weight	425.4990
Crystal system	Monoclinic
Space group	P 2_1/c
a (Å)	9.7261 (9)
b (Å)	26.408 (3)
c (Å)	8.5437 (7)
α (°)	90
β (°)	91.509 (3)
γ (°)	90
Volume (Å3)	2193.7 (2)
Z	4
Reflections collected	3859
R, wR2	0.0924, 0.2053
Goodness–of–fit on F2	1.152