

SUPPORTING INFORMATION

CONCISE SYNTHESIS OF (±)-CEPHALOTAXINE

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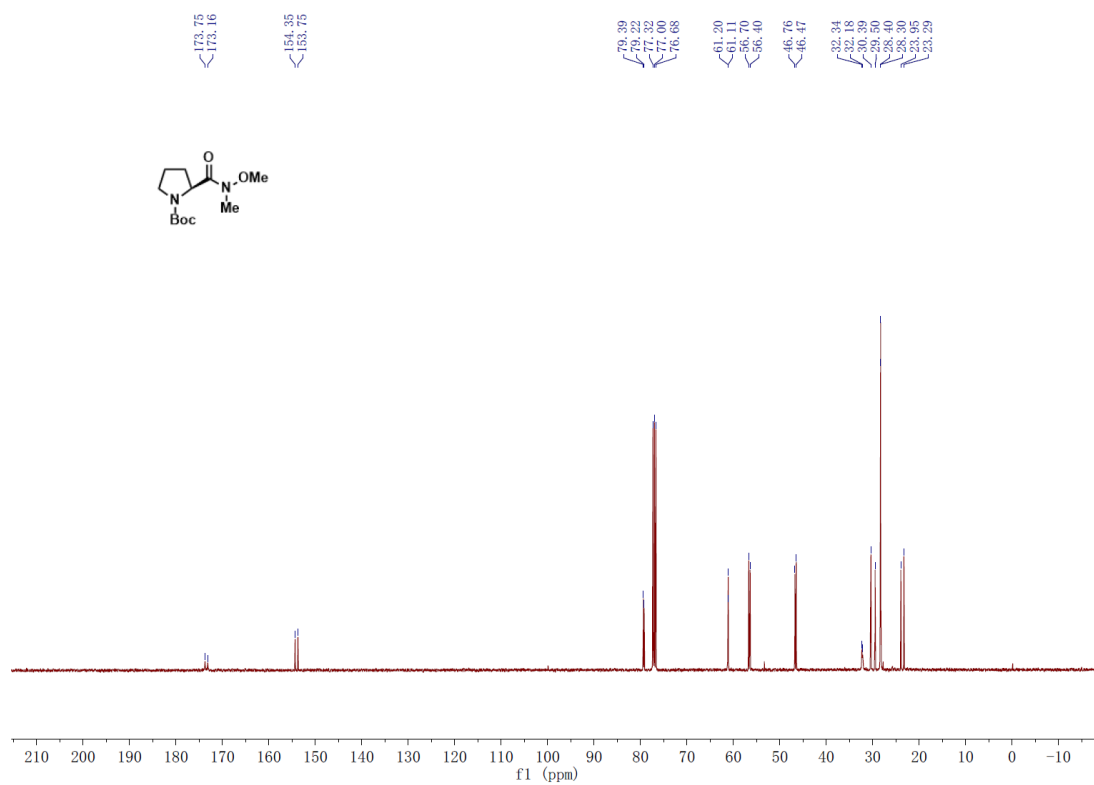
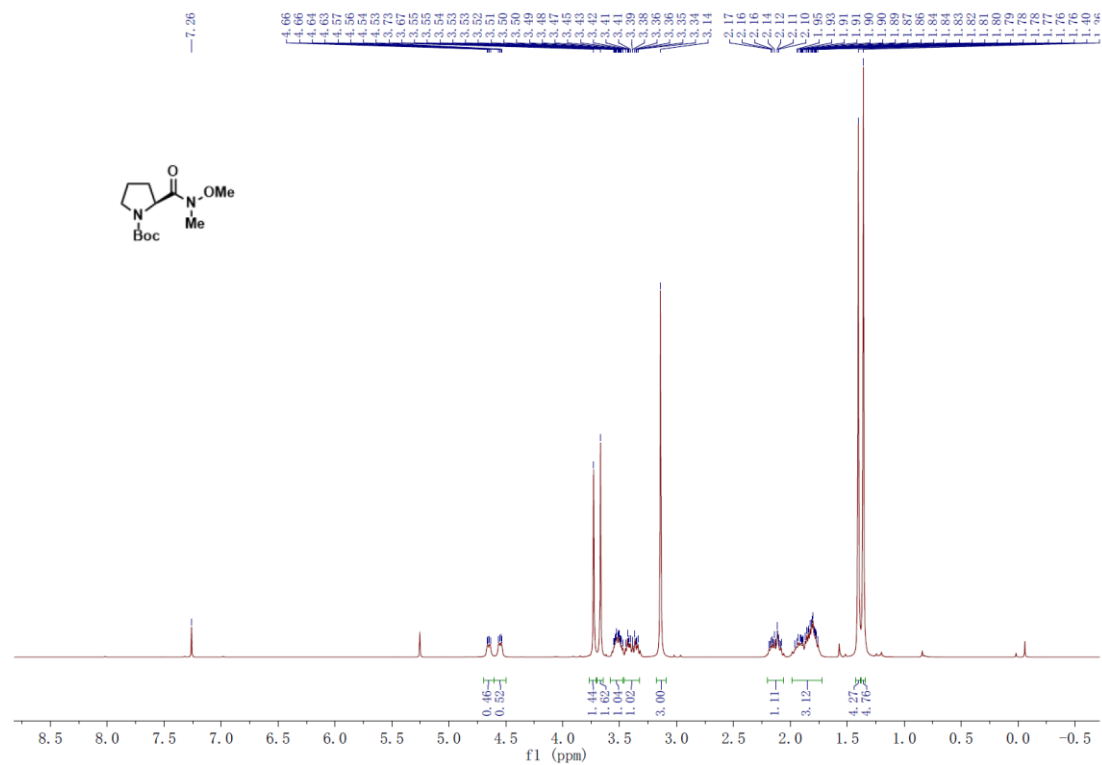
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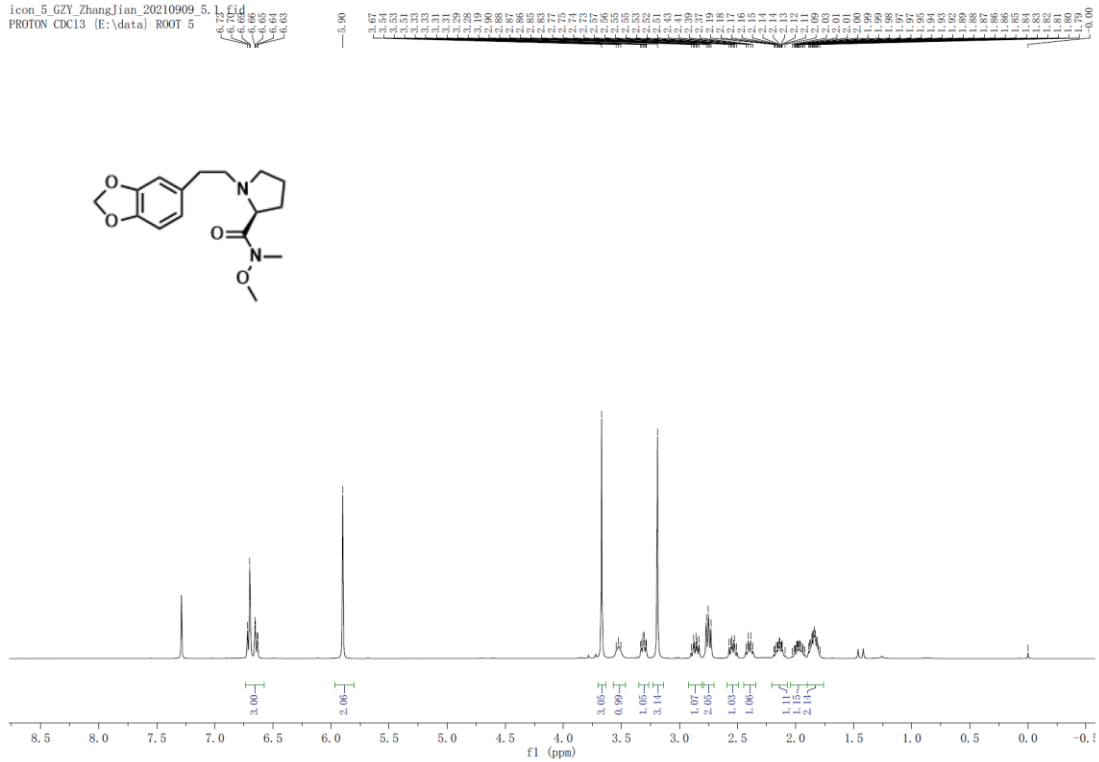
I. General procedure

All solvents were redistilled prior to use and anhydrous solvents were treated using standard techniques. All reactions were monitored by thin-layer chromatography using high silica gel GF pre-coated plates (0.25 mm). IR spectra were recorded on a Thermo Nicolet 6700 FT-IR Spectrometer using KBr disks and are reported in frequency of absorption (cm^{-1}). HRMS were acquired by a Waters Micromass Q-ToF MicroTM instrument using the ESI technique. ¹H and ¹³C NMR spectra were recorded on a Bruker AM-400 spectrometer with TMS as an internal standard and CDCl₃ as solvent. Flash column chromatography was conducted on silica gel (200-300 mesh).

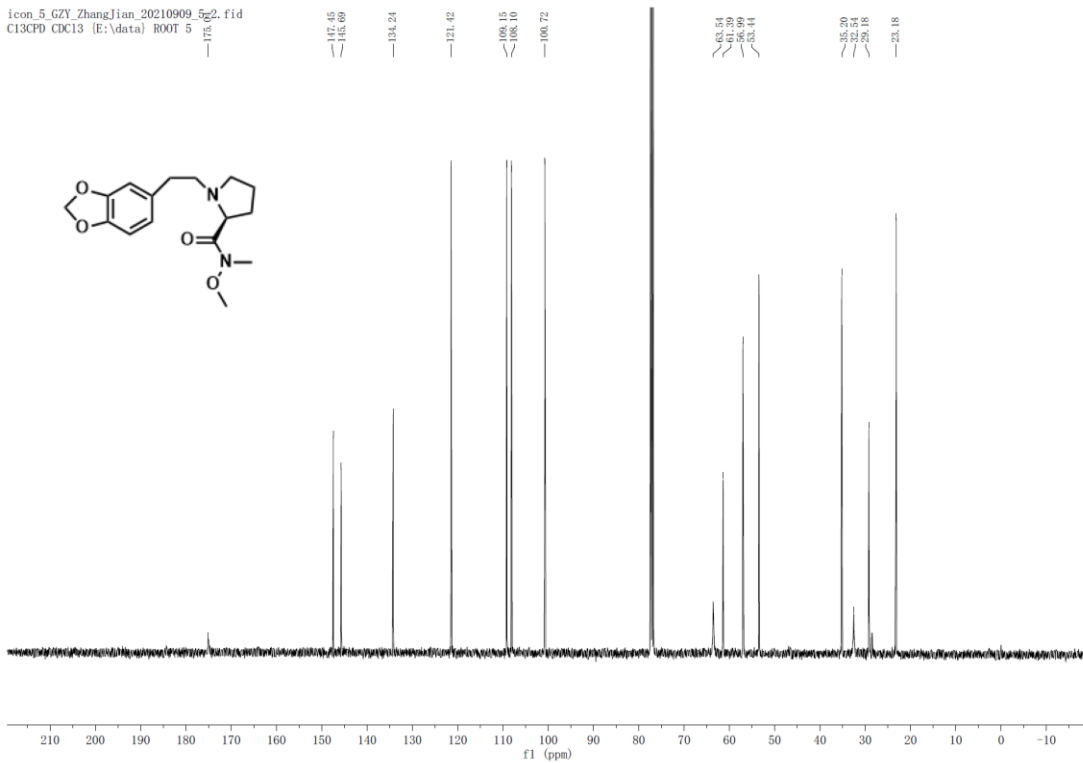
II. ¹H and ¹³C NMR Spectra of Synthesized Compounds

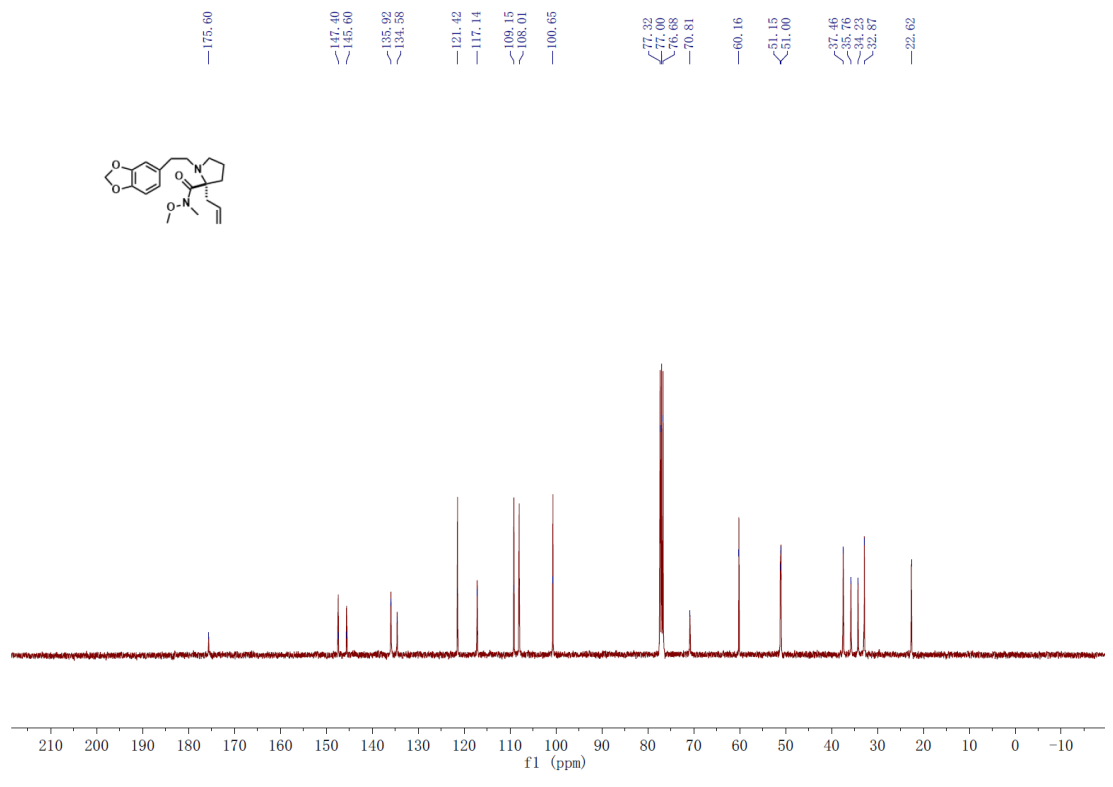
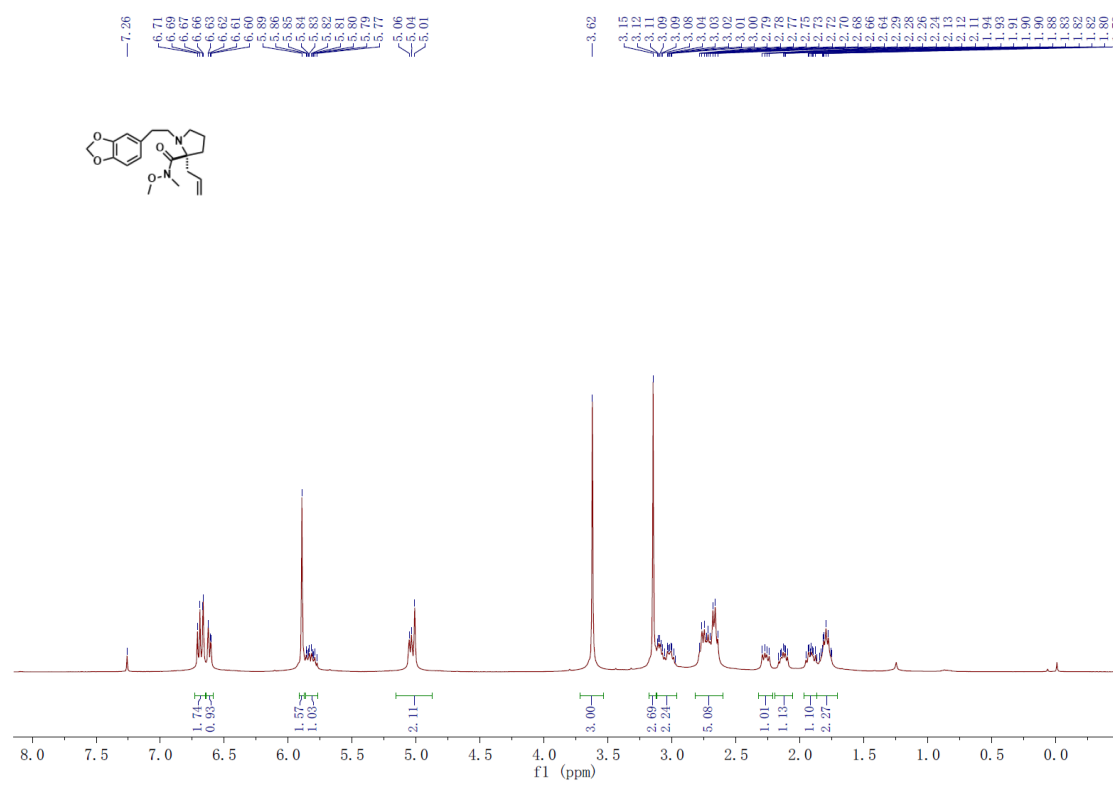


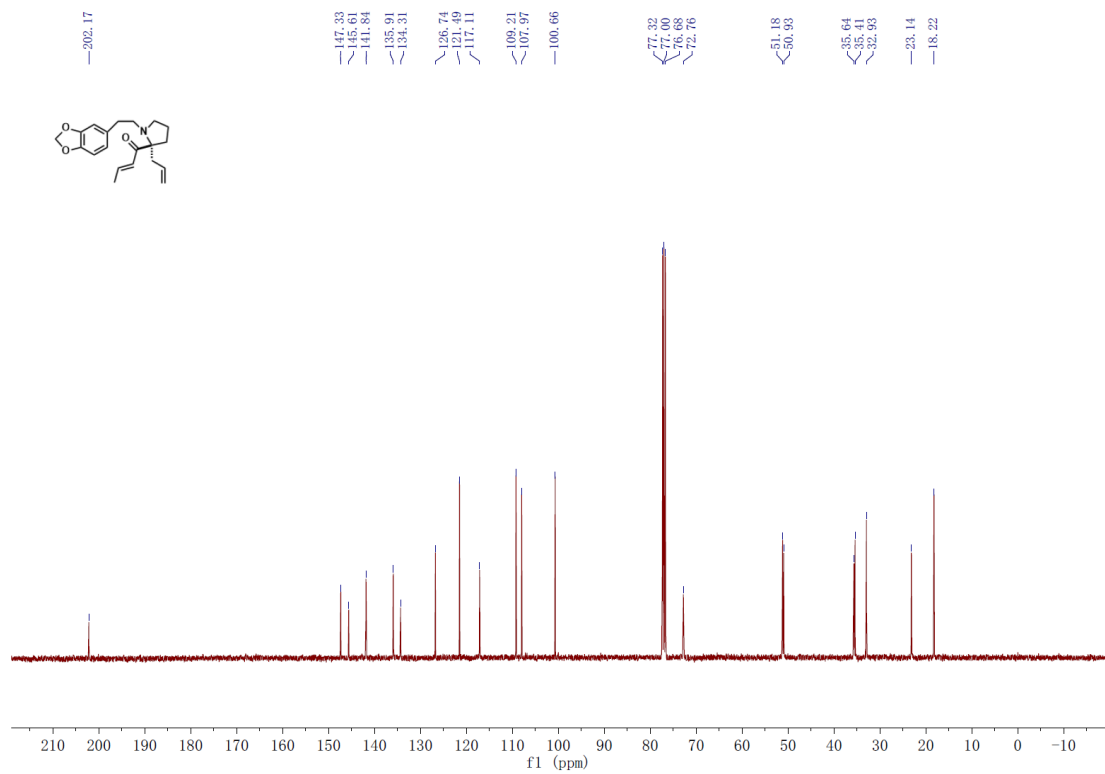
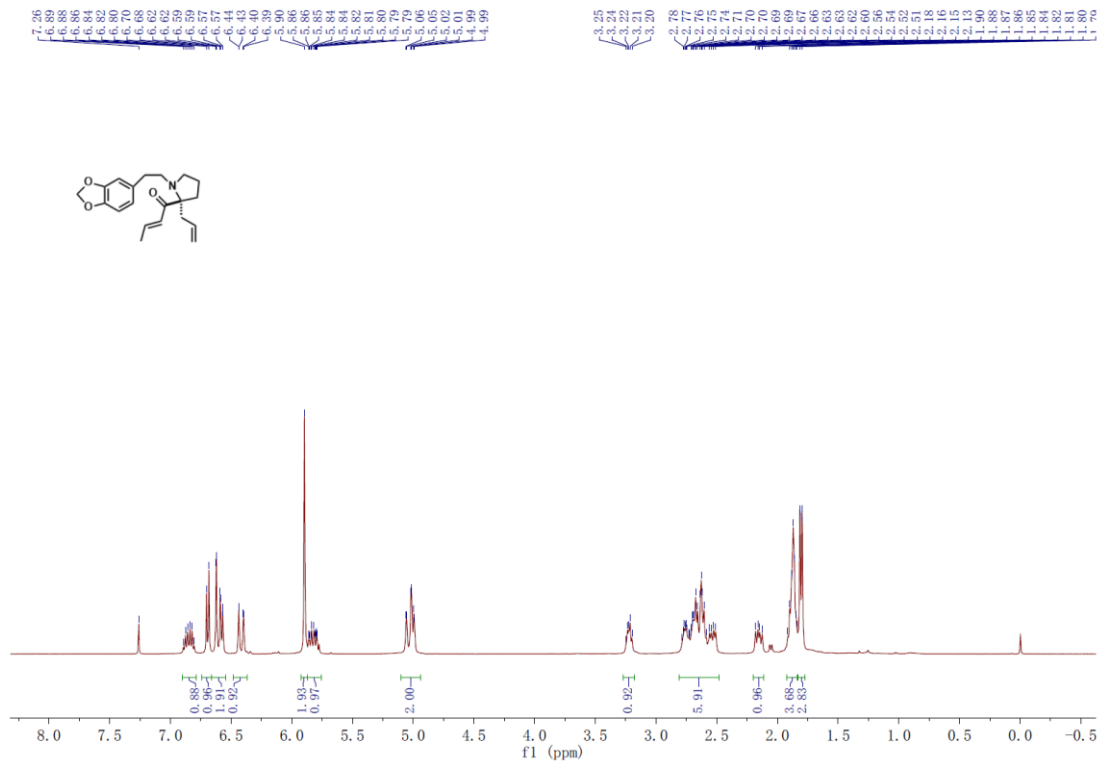
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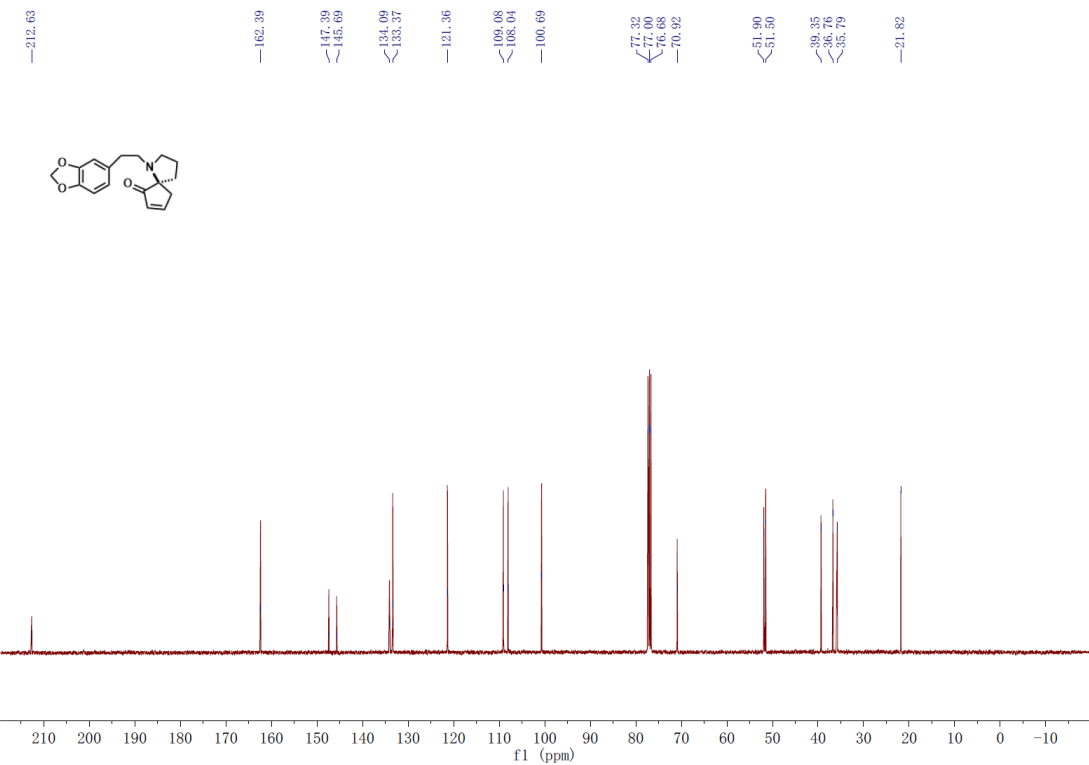
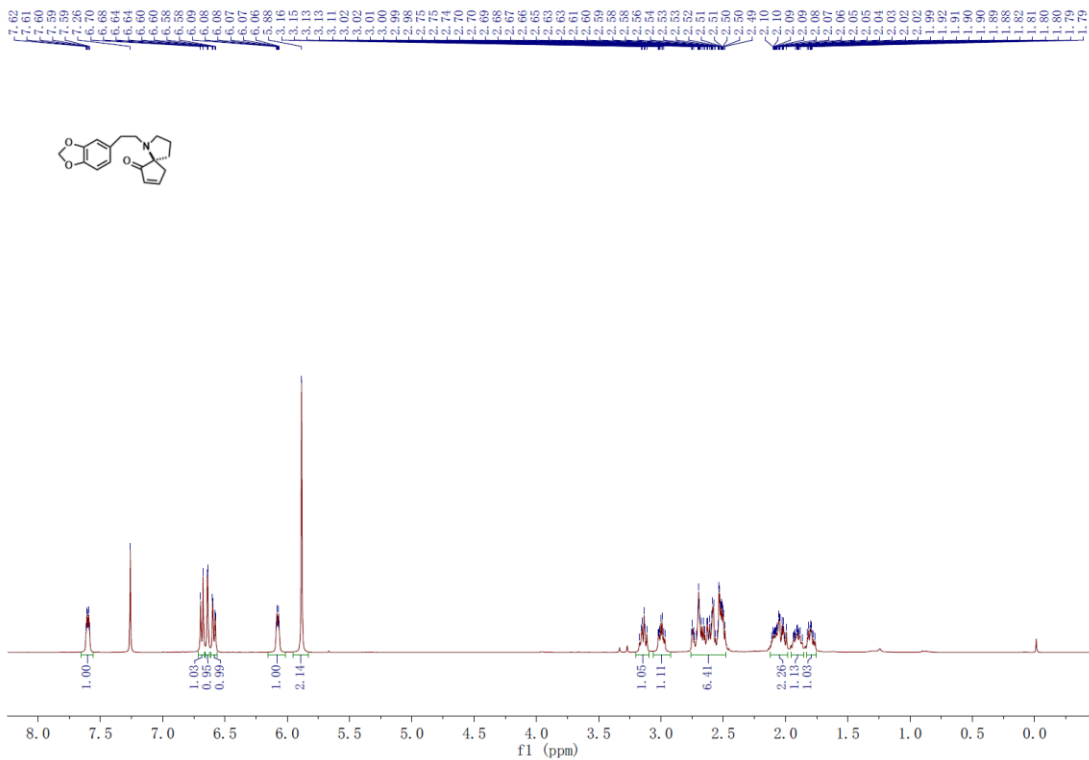


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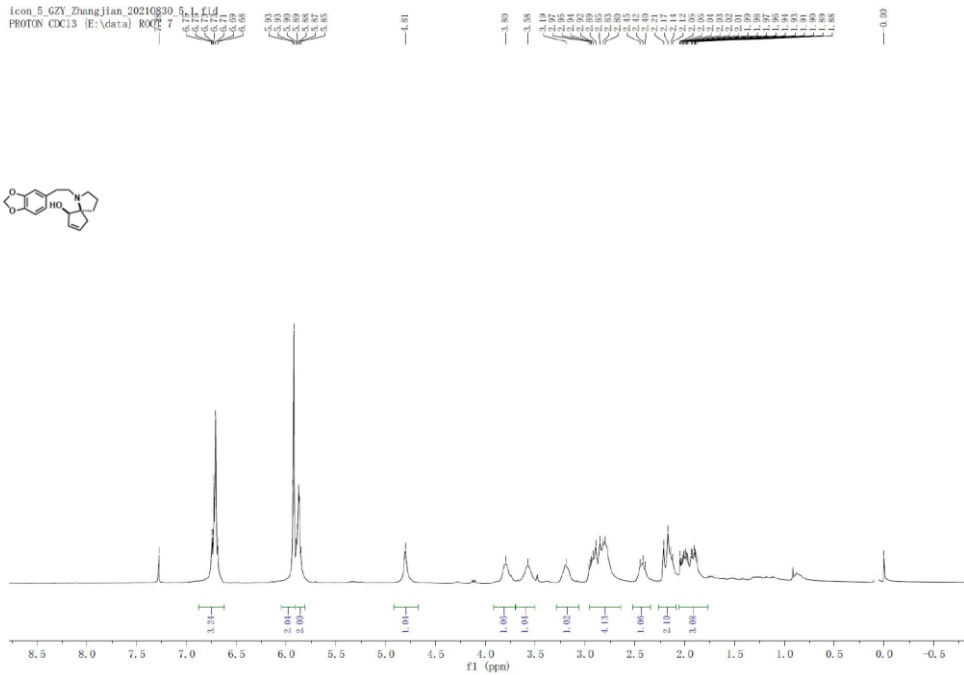








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