

Highly Regioselective Synthesis of 3-Alkenyl-Oxindole Ring-fused 3,3'-disubstituted Oxindoles *via* Direct gamma-substitution of Morita-Baylis-Hillman Carbonates of Isatins with 3-substituted Oxindoles

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Supporting Information

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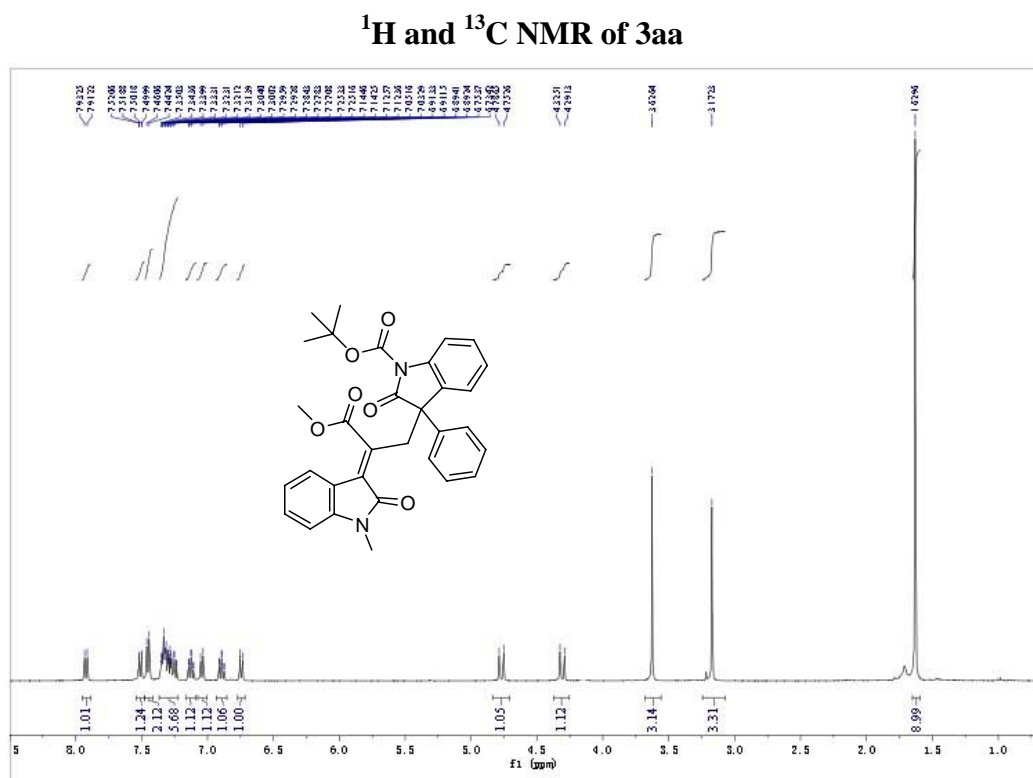
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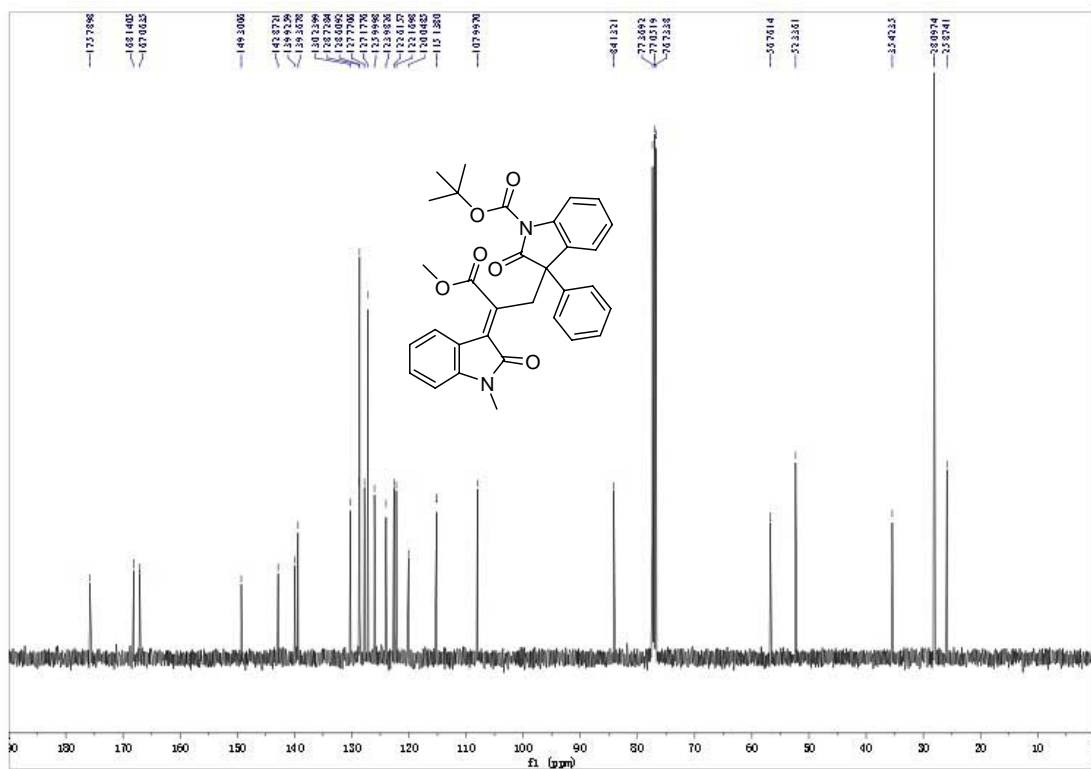
1: General Experimental Information

The ^1H and ^{13}C NMR spectra were recorded on Bruker Avance DMX 400 MHz NMR spectrometers in CDCl_3 and DMSO-d_6 using TMS as internal standard. Chemical shifts were reported as δ values (ppm). Melting points were uncorrected and recorded on an Electrothermal 9100 digital melting point apparatus. High-resolution mass spectra (HRMS-ESI) were obtained on a MicroTM Q-TOF Mass Spectrometer. Crystals suitable for X-ray crystallographic studies were obtained by crystallization from methanol.

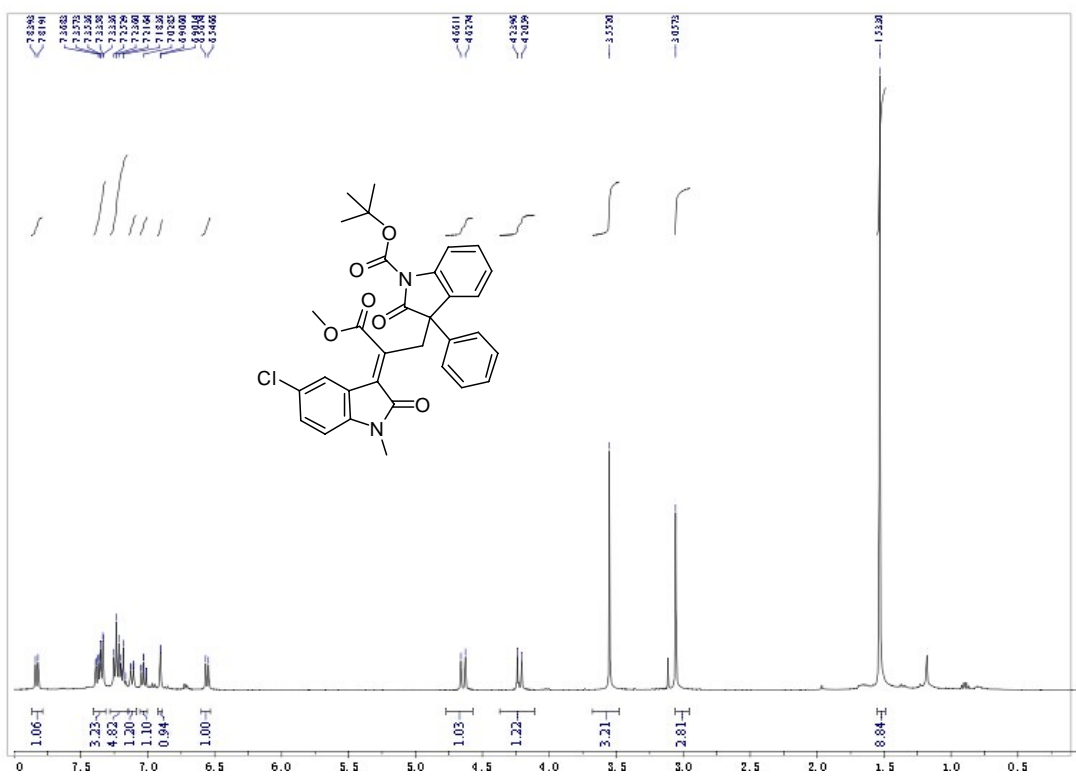
Reagents were purchased from commercial sources and were used as received unless mentioned otherwise. Reactions were monitored by thin layer chromatography using silica gel GF₂₅₄ plates. Column chromatography was performed on silica gel (300-400 mesh).

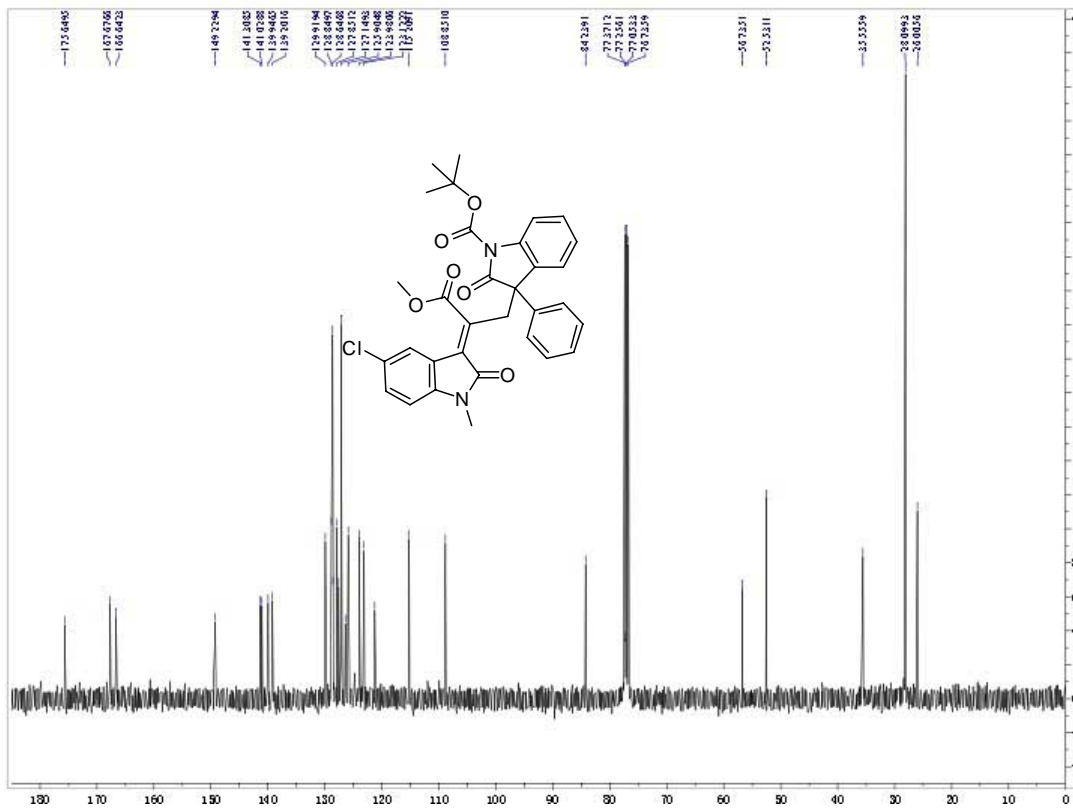
2: The Copies of ^1H NMR and ^{13}C NMR Spectra for Compounds 3



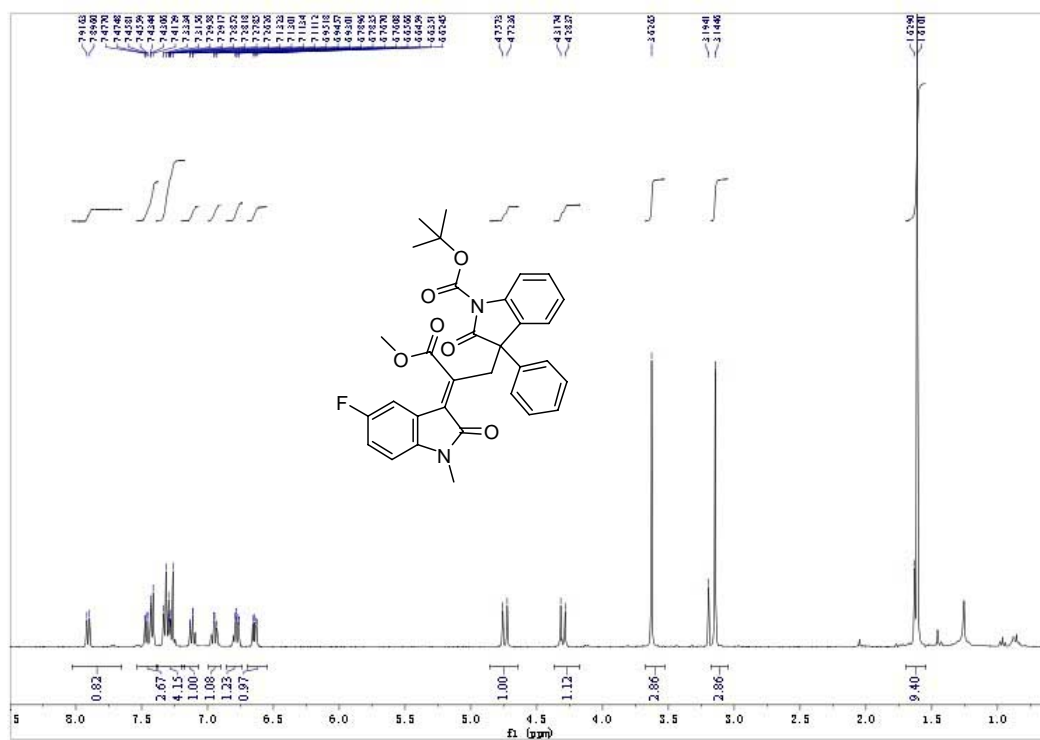


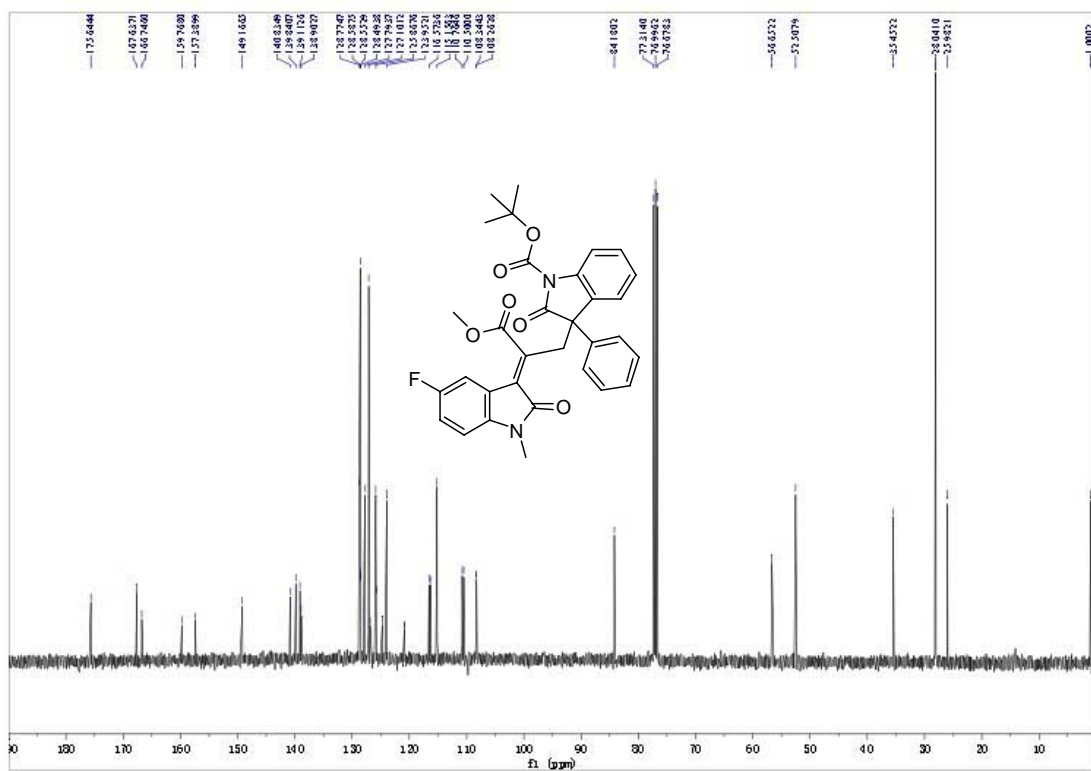
^1H and ^{13}C NMR of 3ab



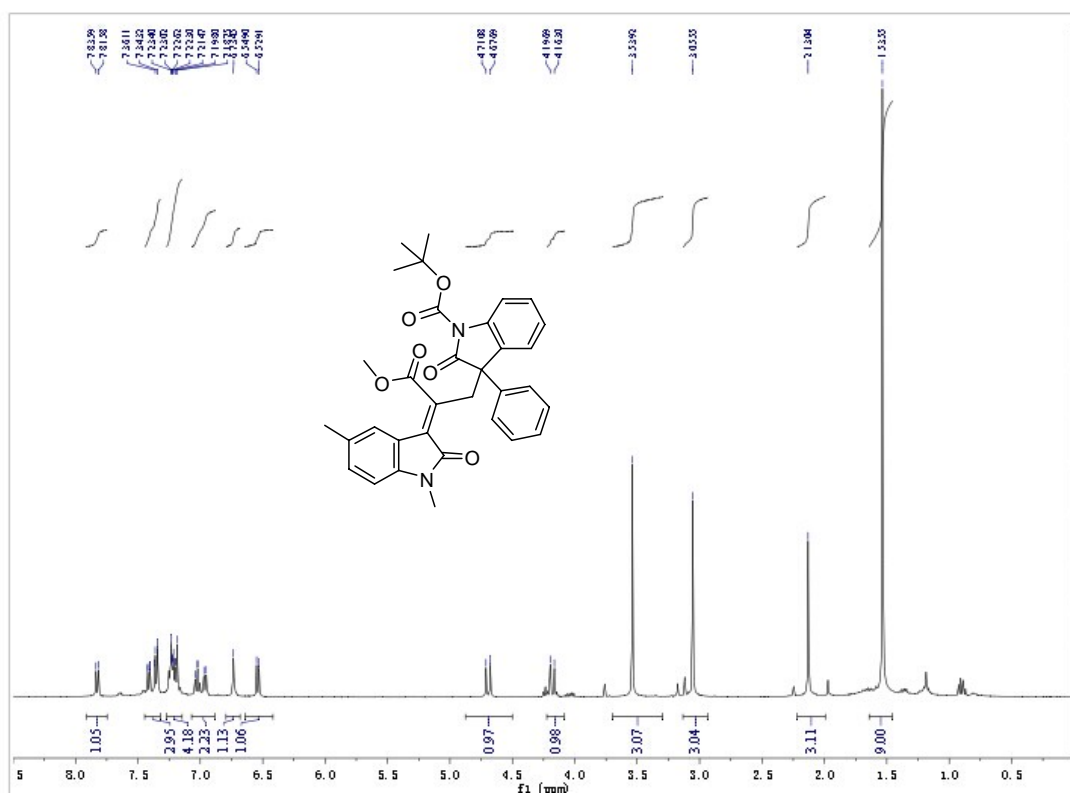


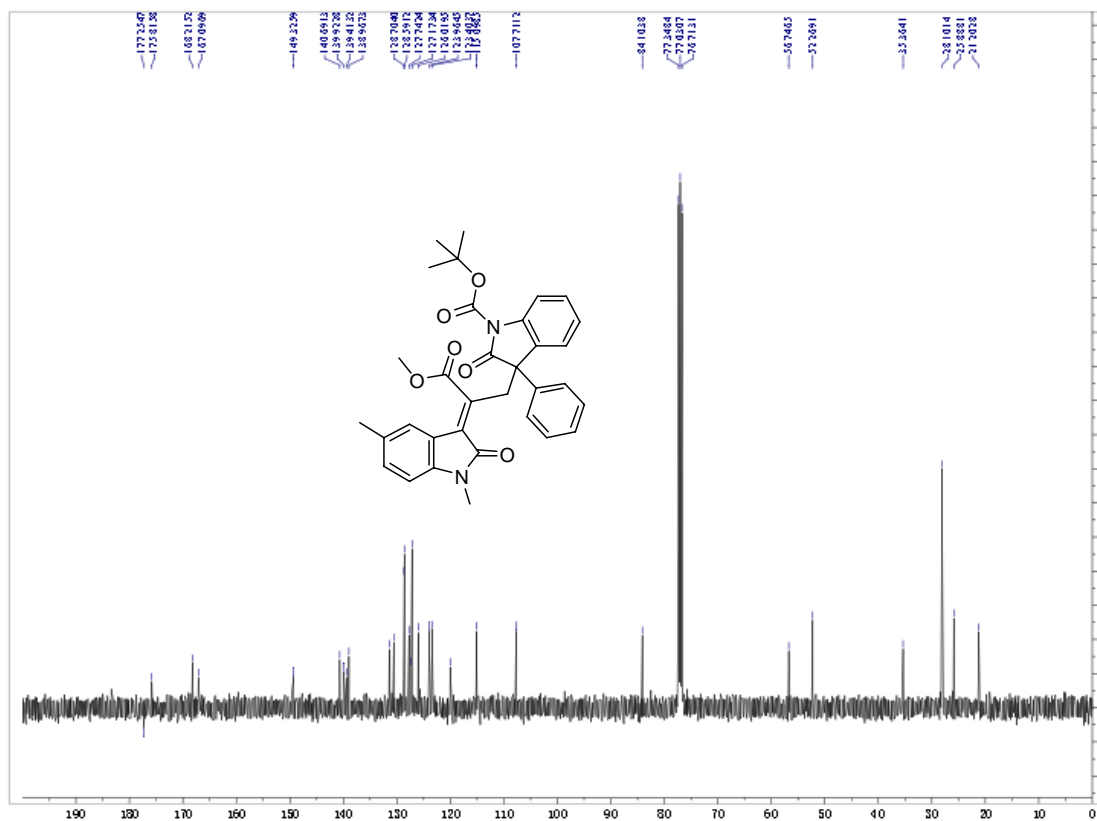
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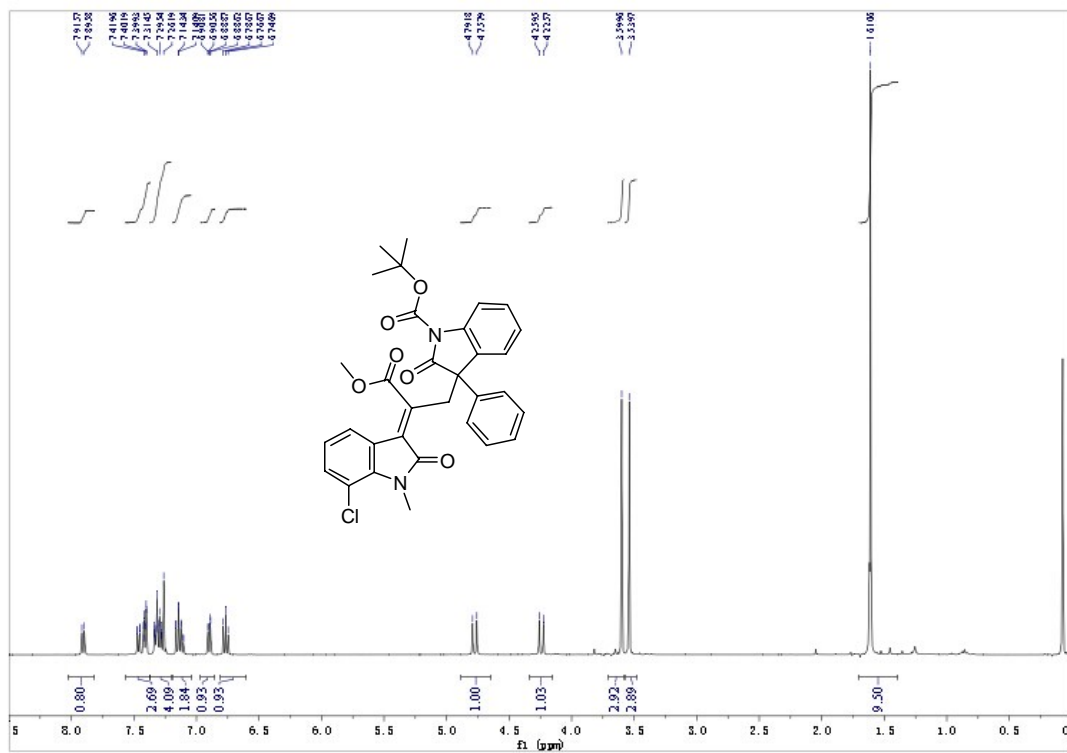


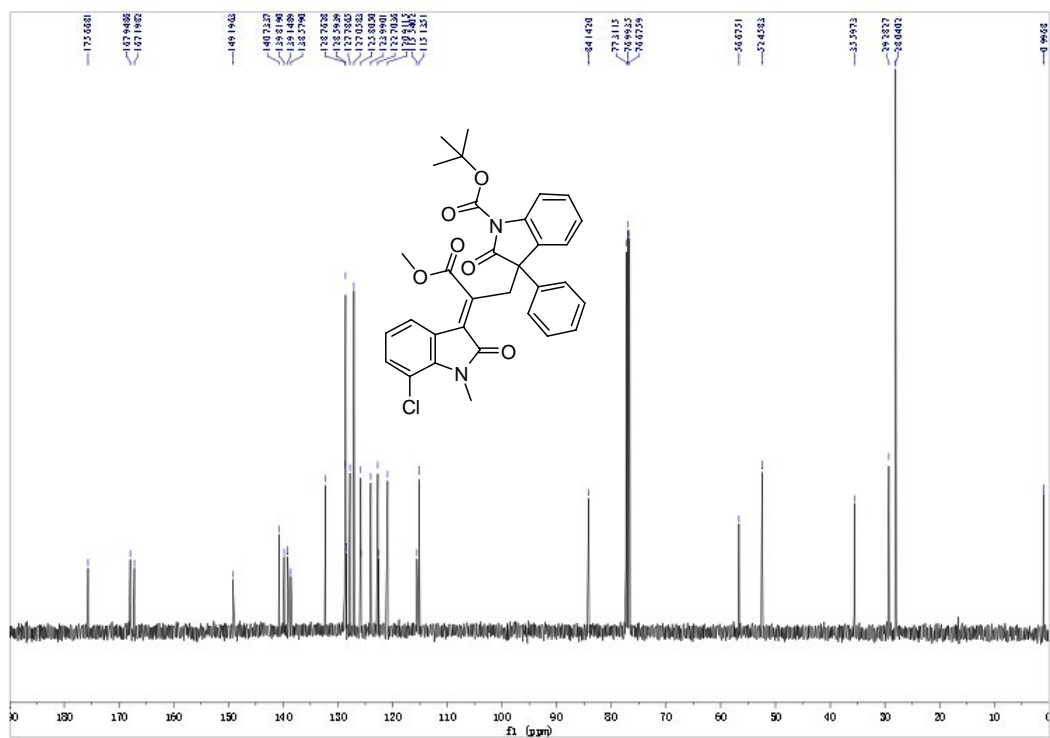
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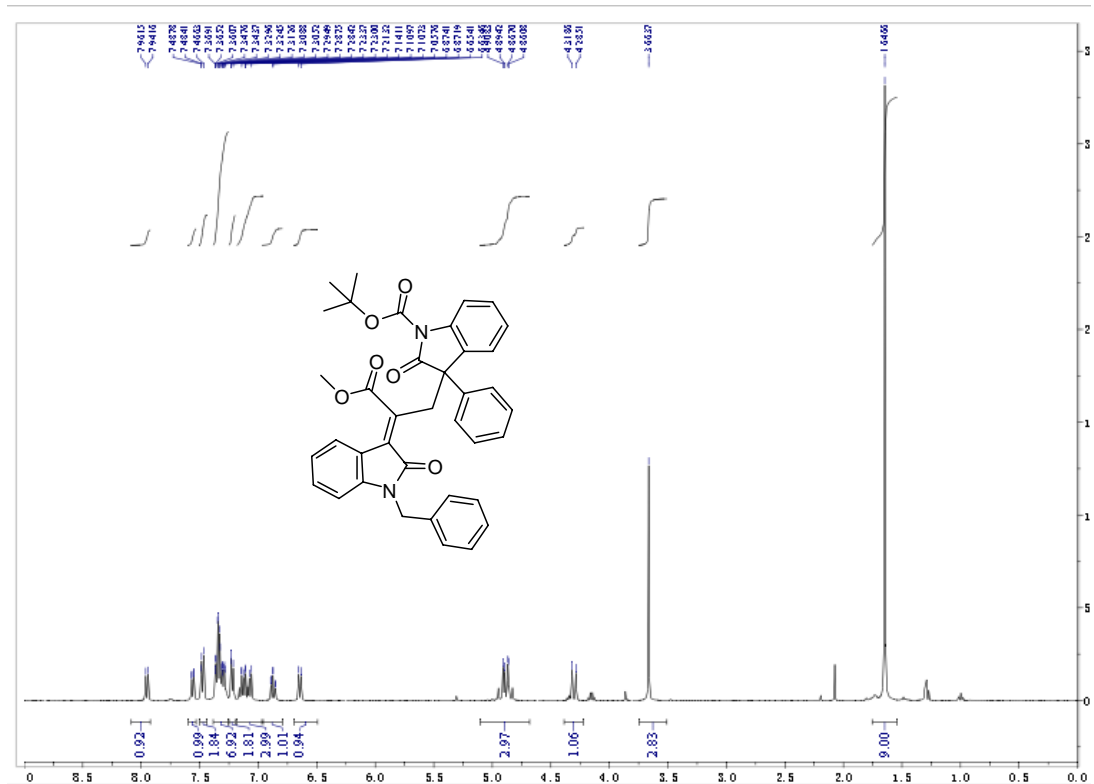


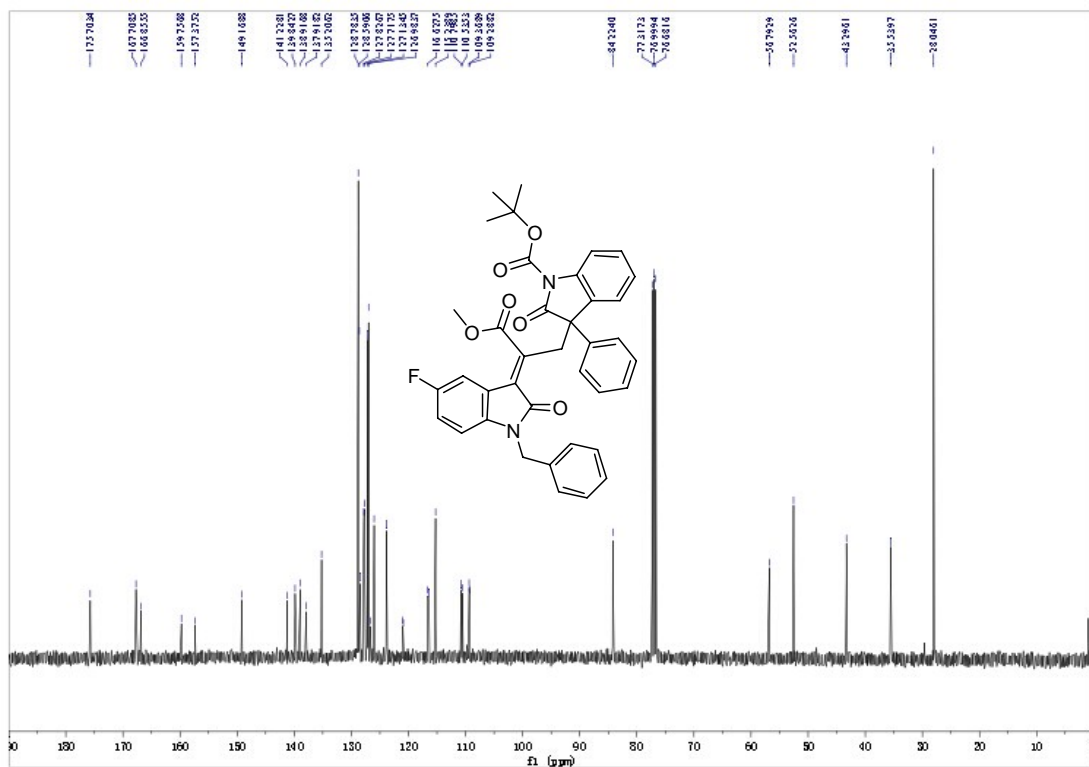
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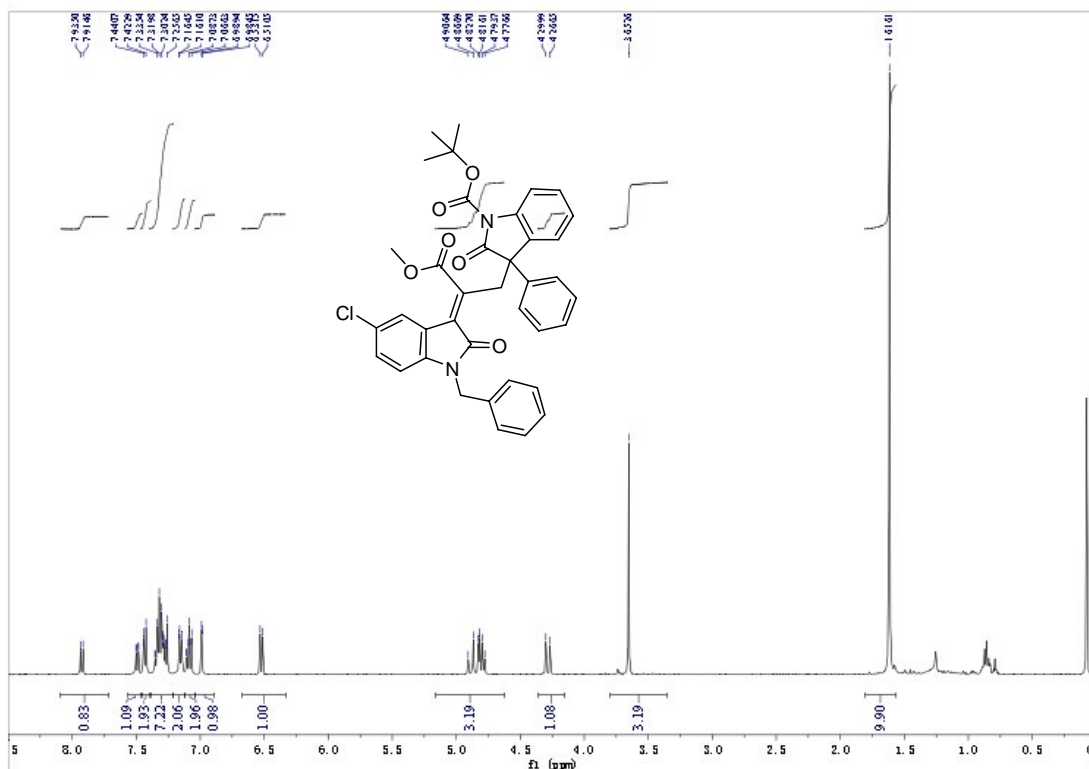


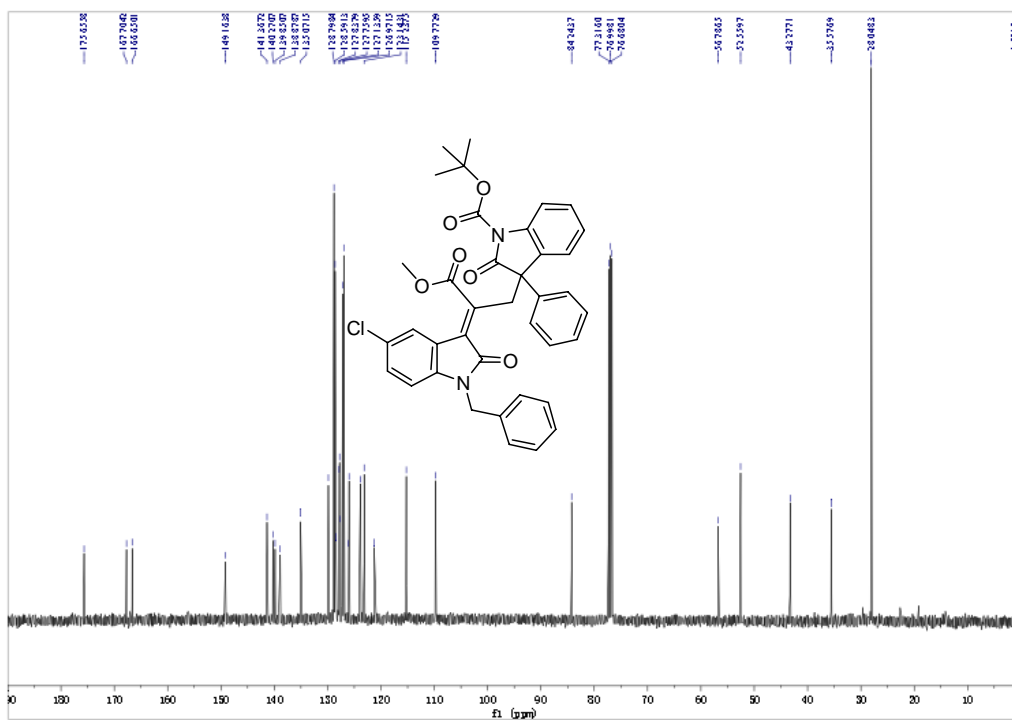
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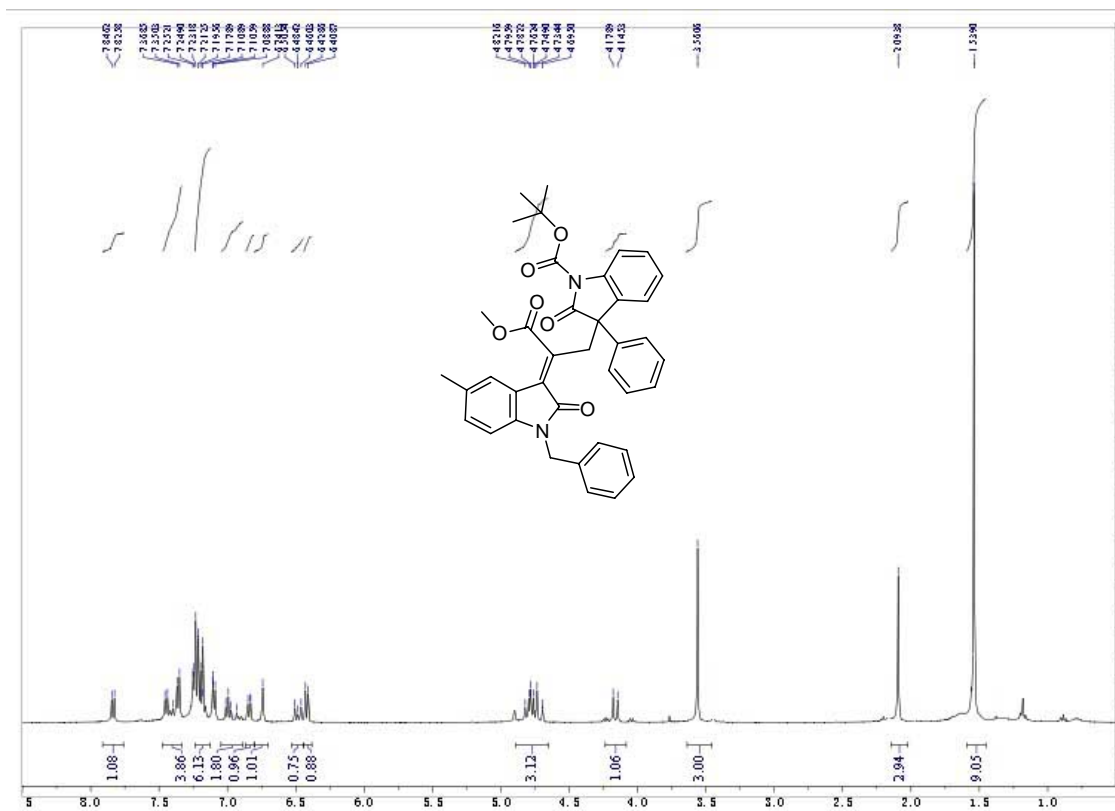


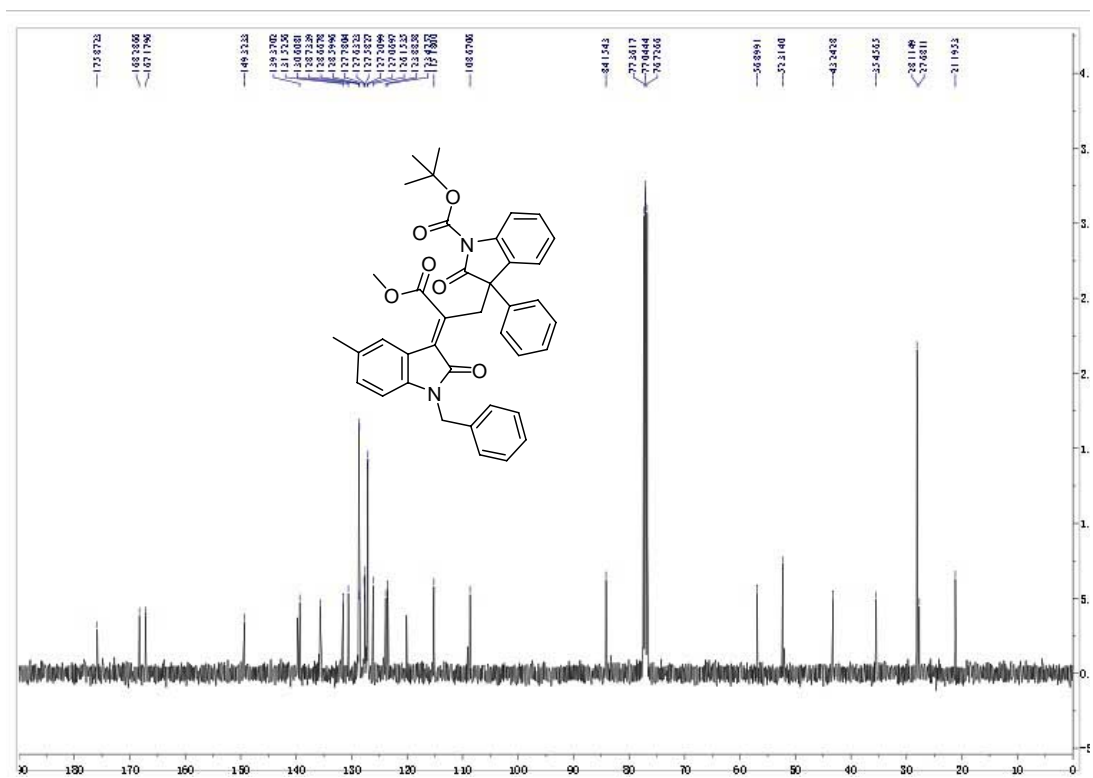
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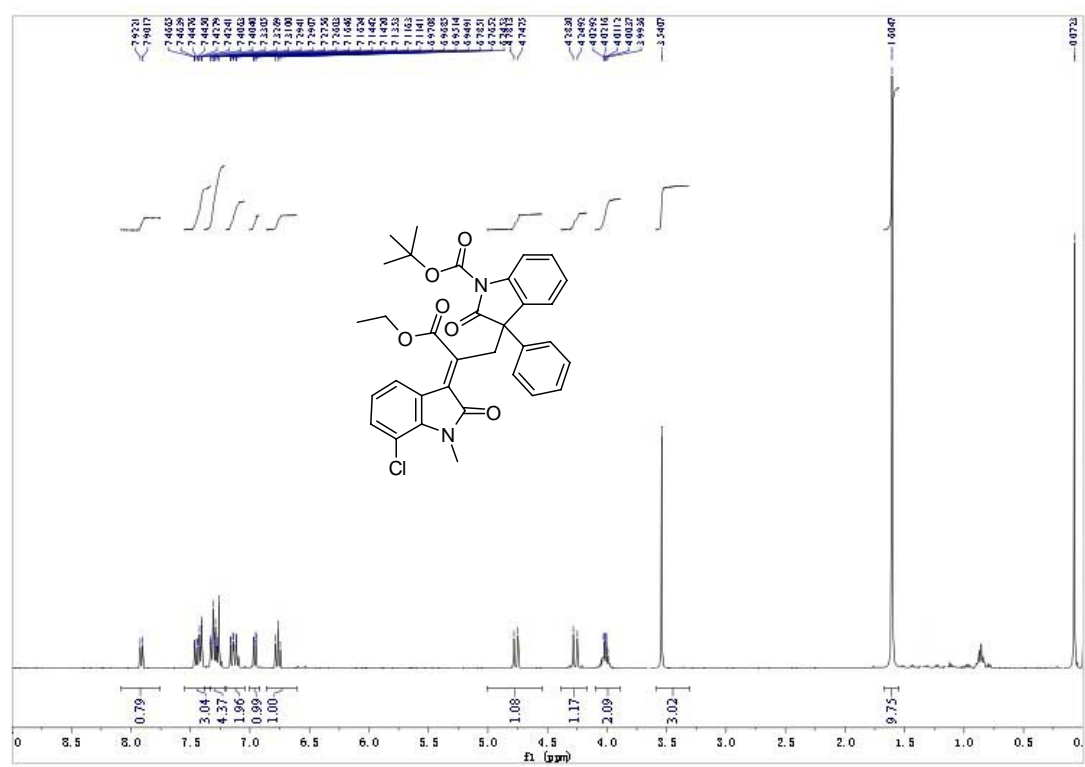


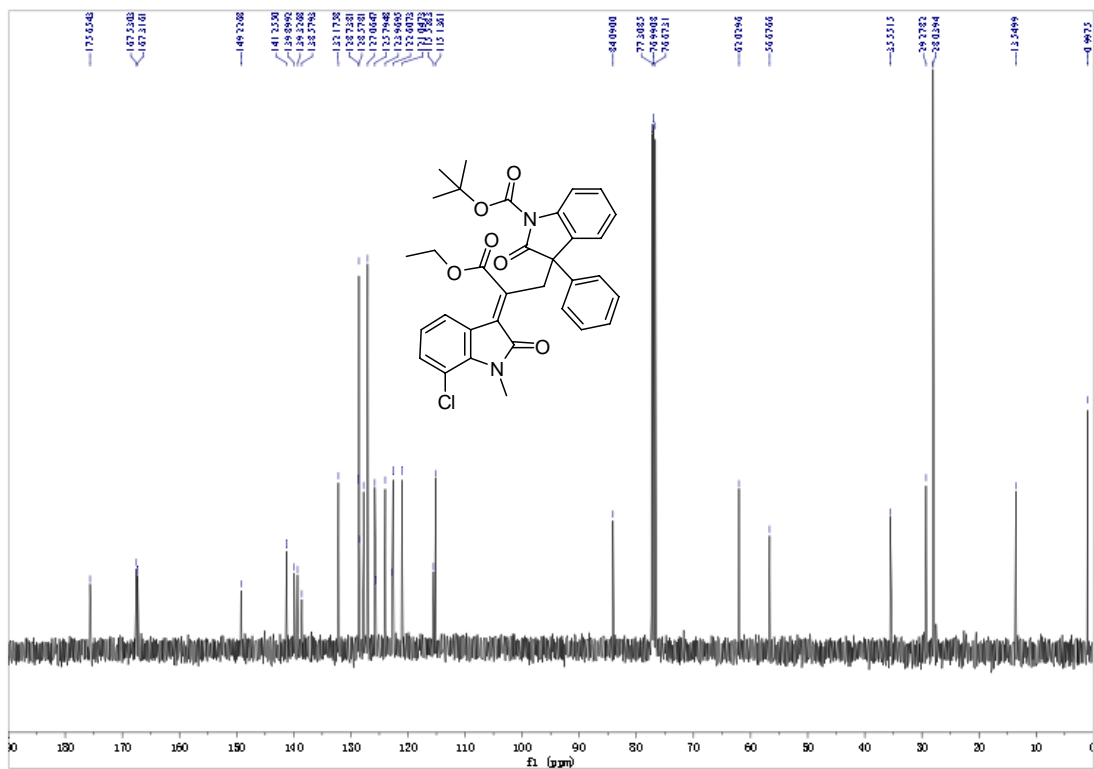
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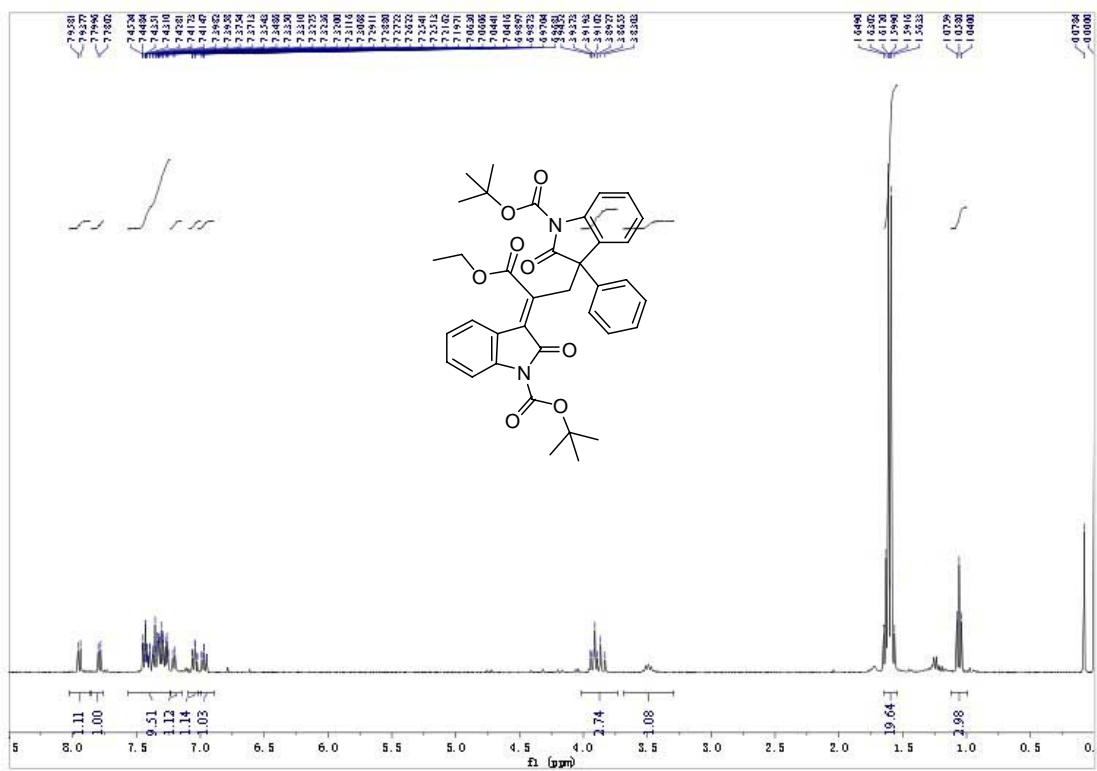


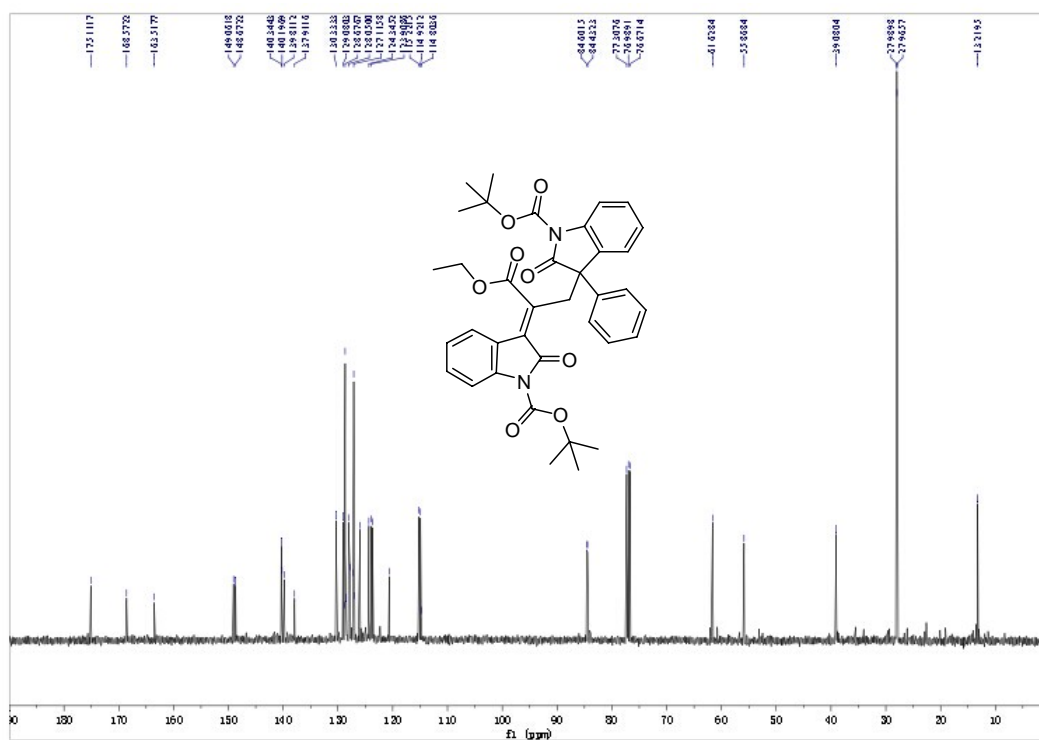
^1H and ^{13}C NMR of 3aj



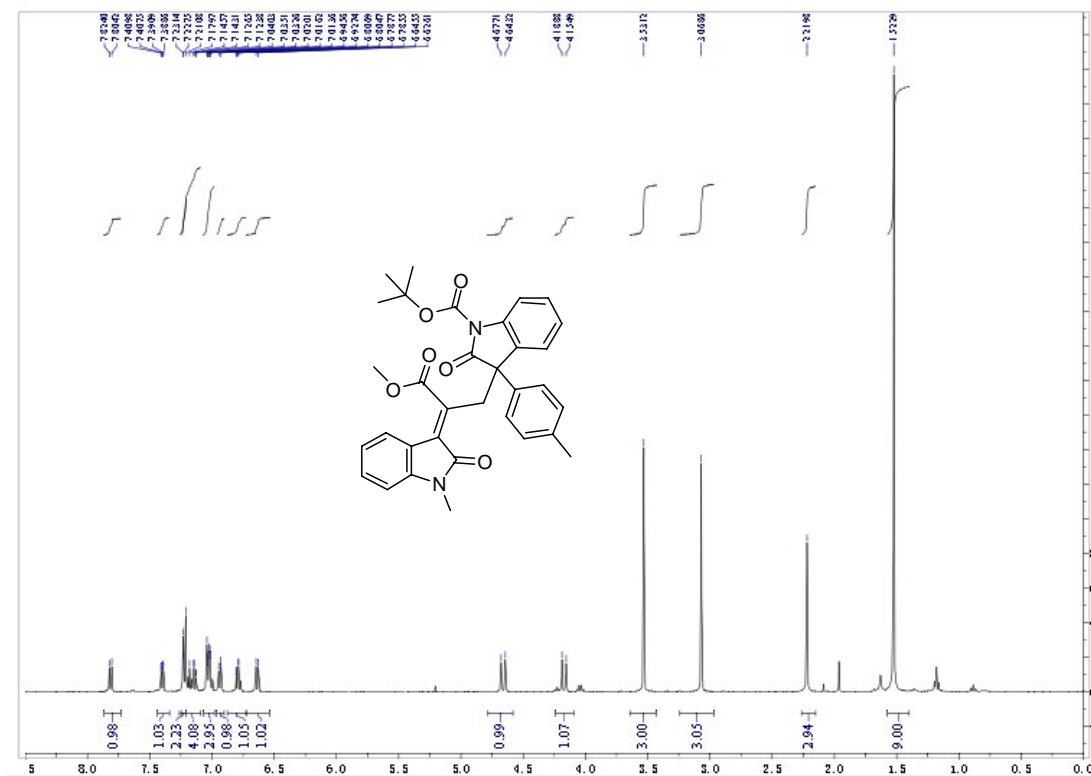


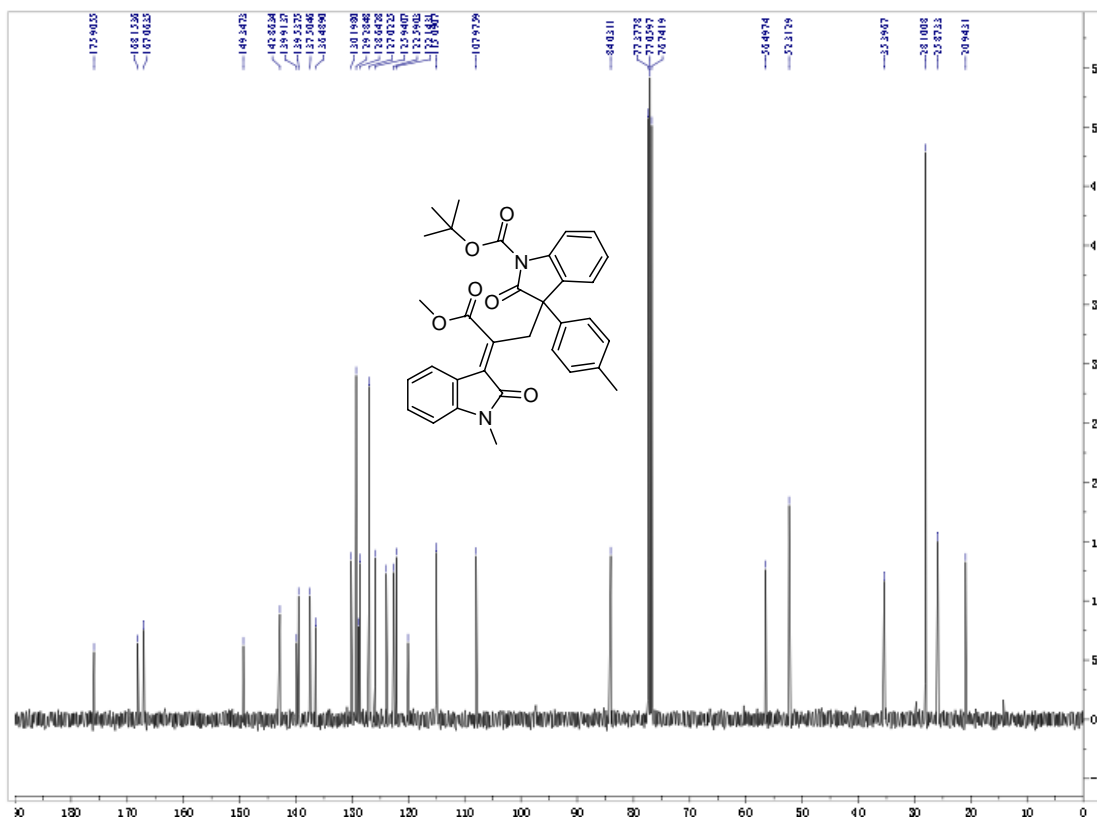
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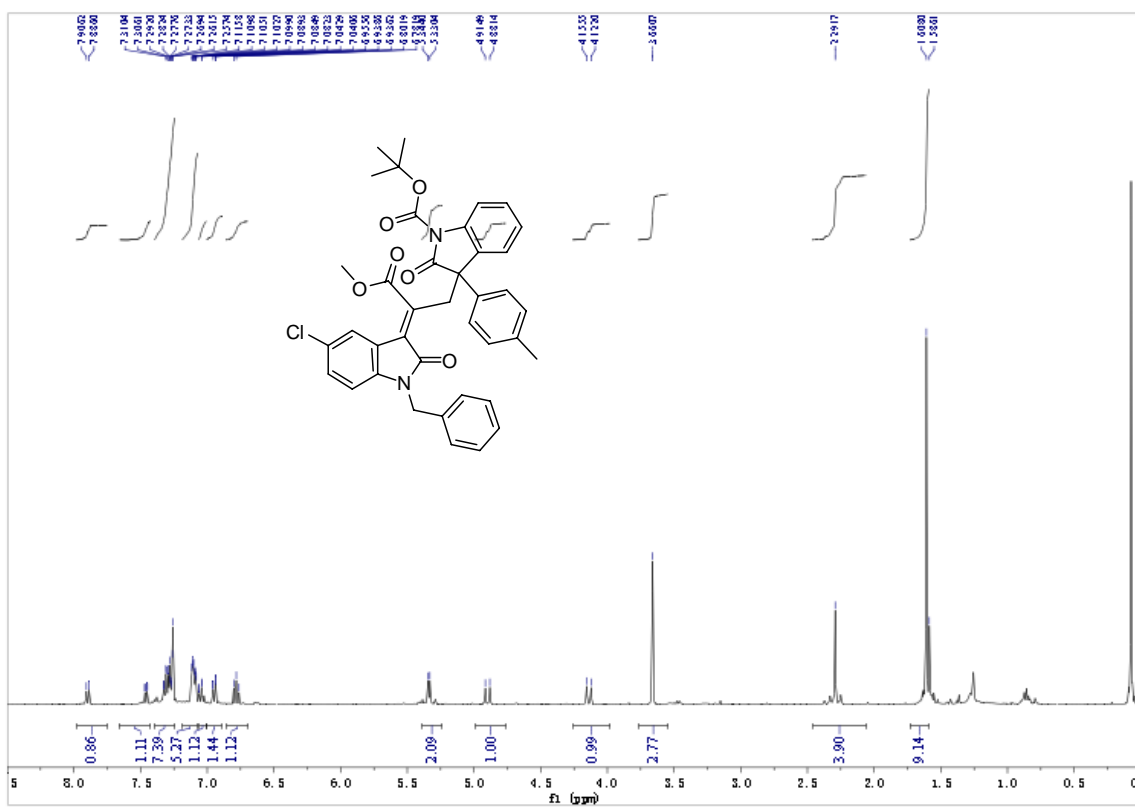


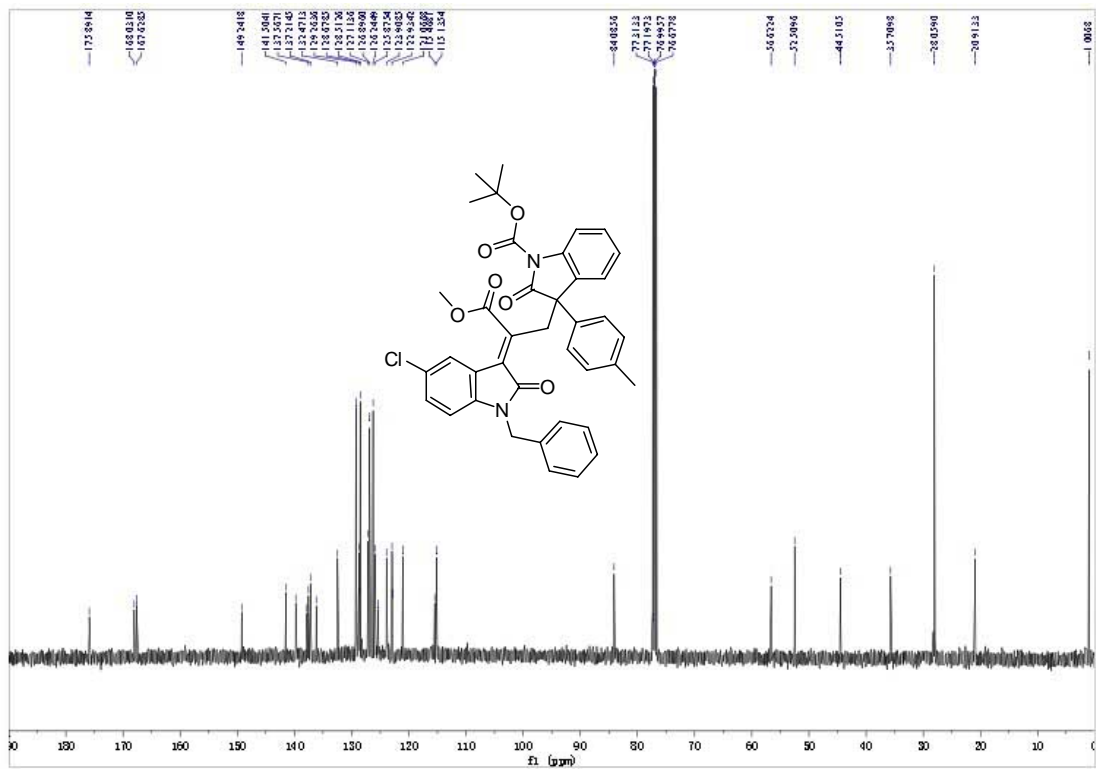
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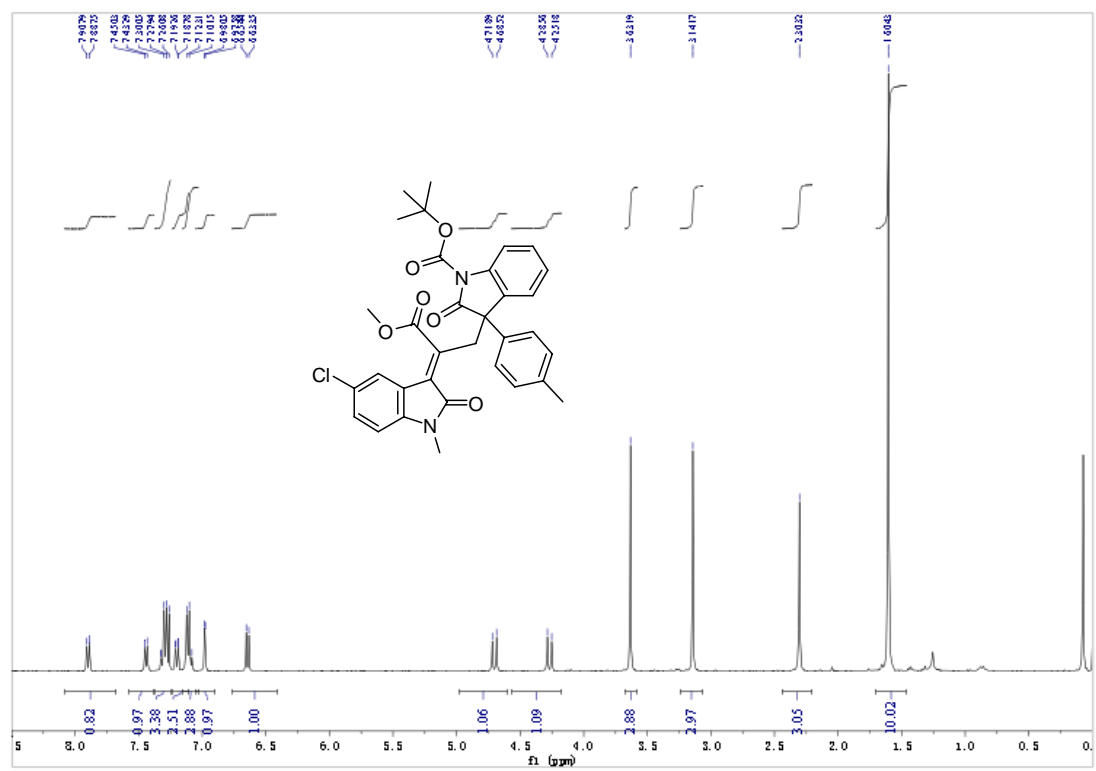


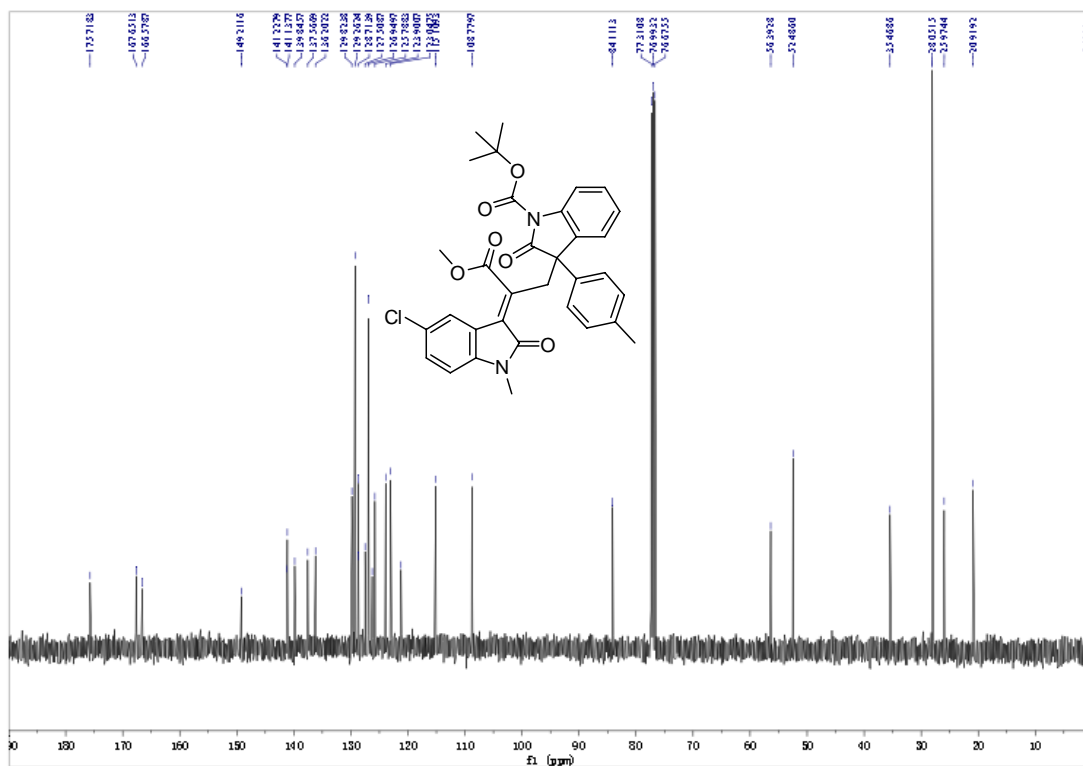
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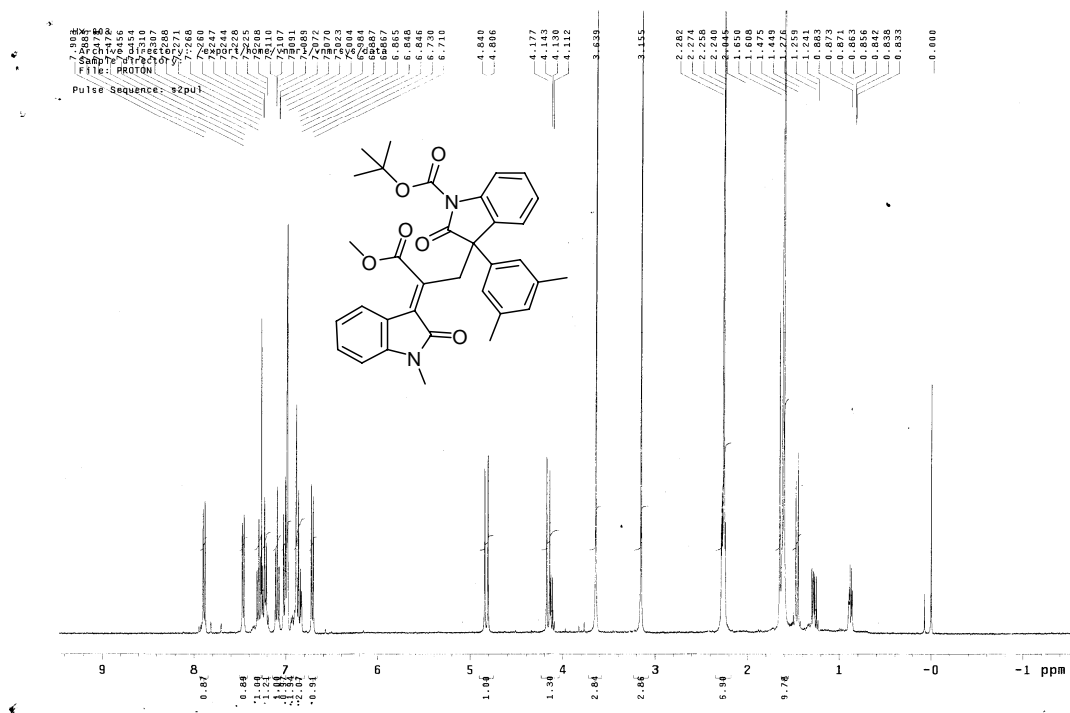


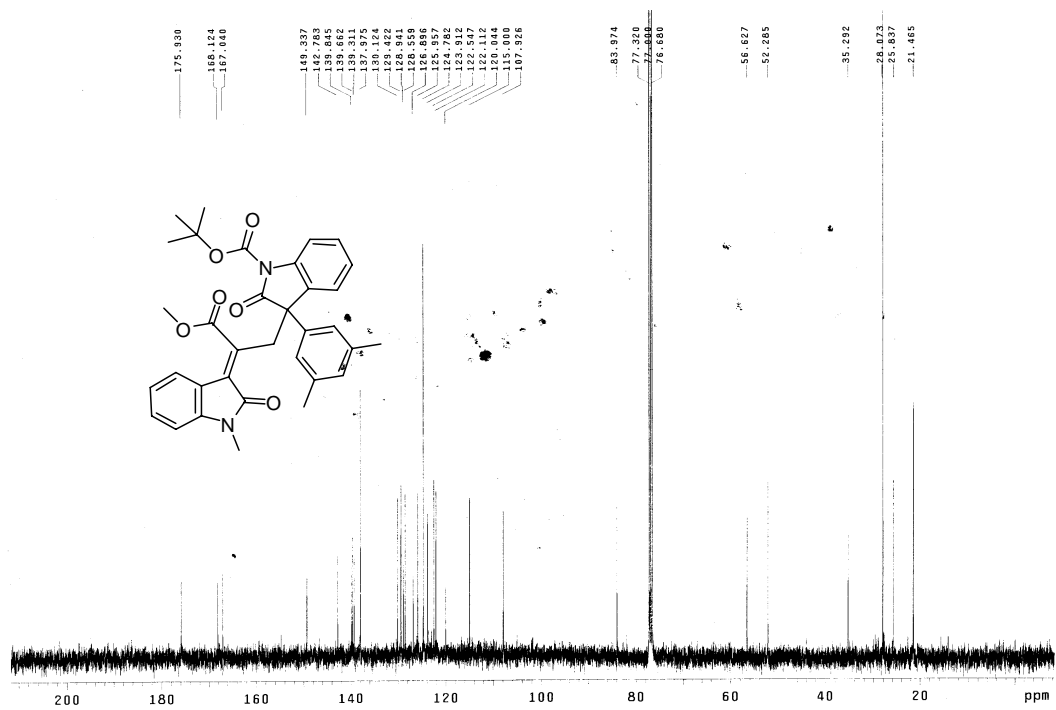
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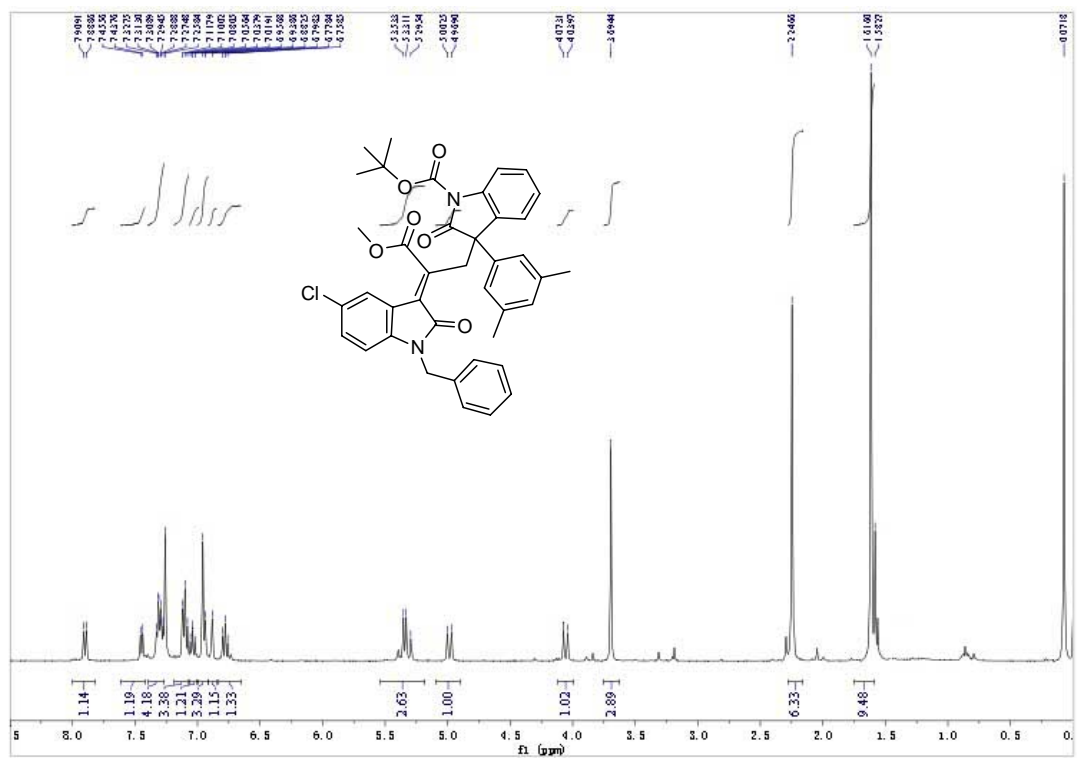


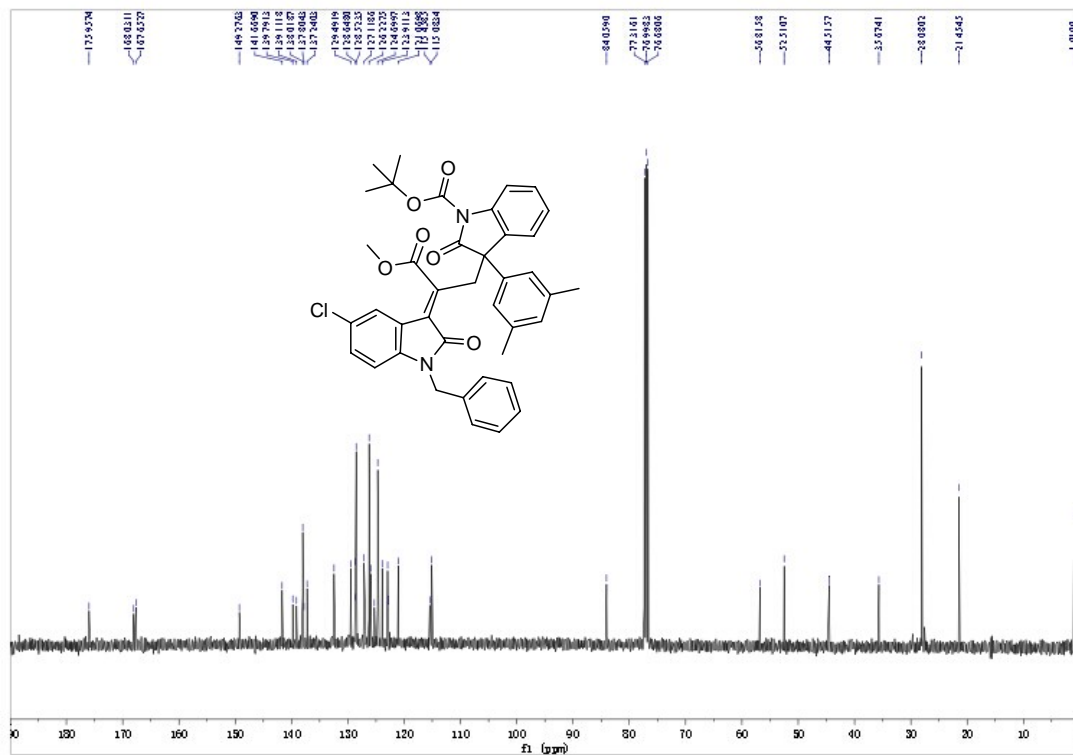
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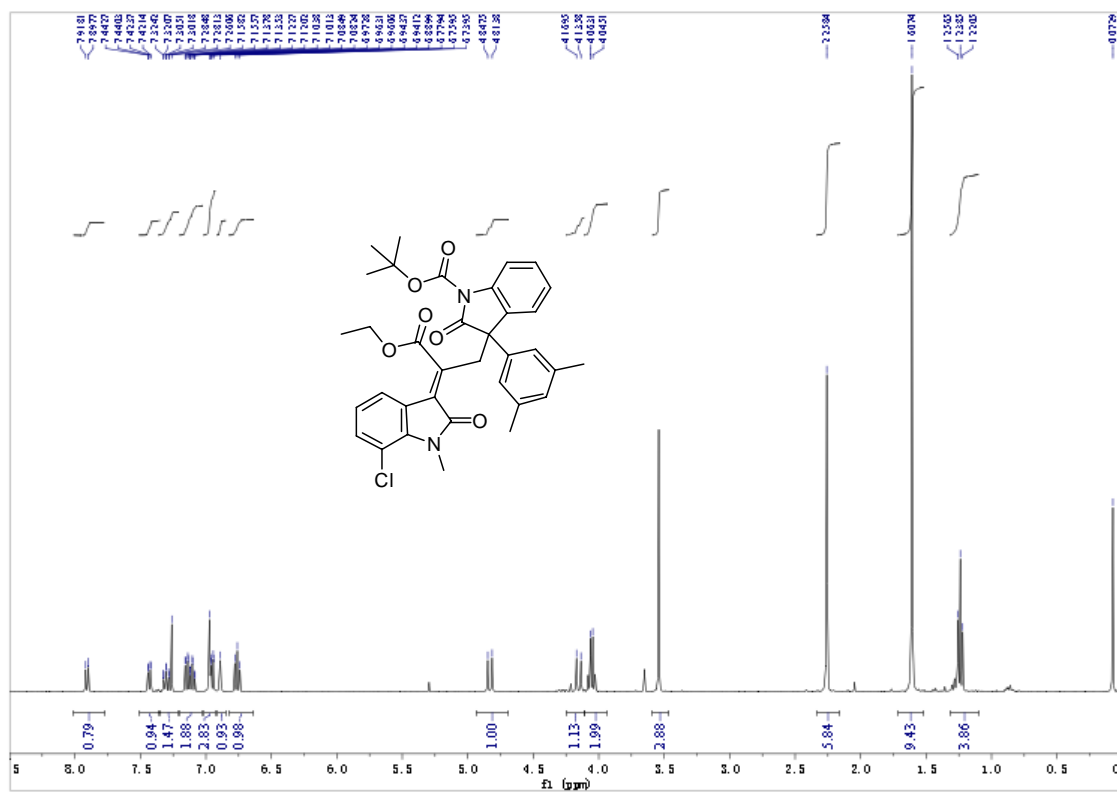


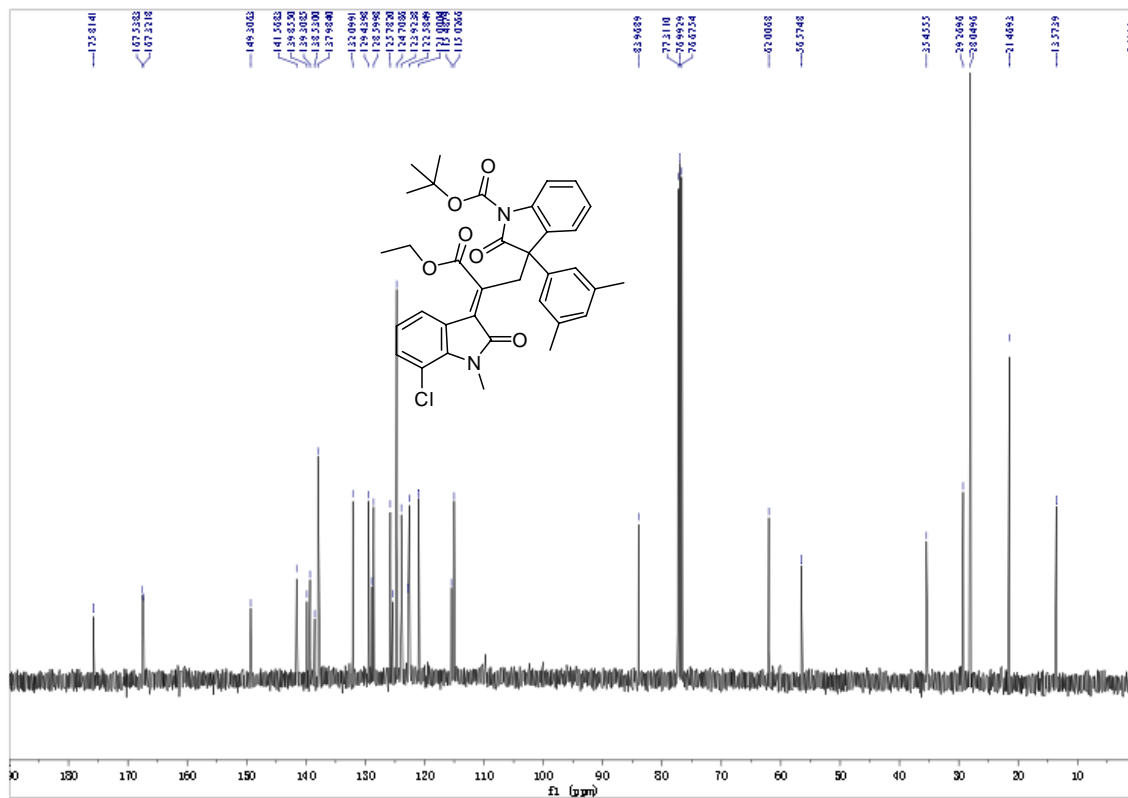
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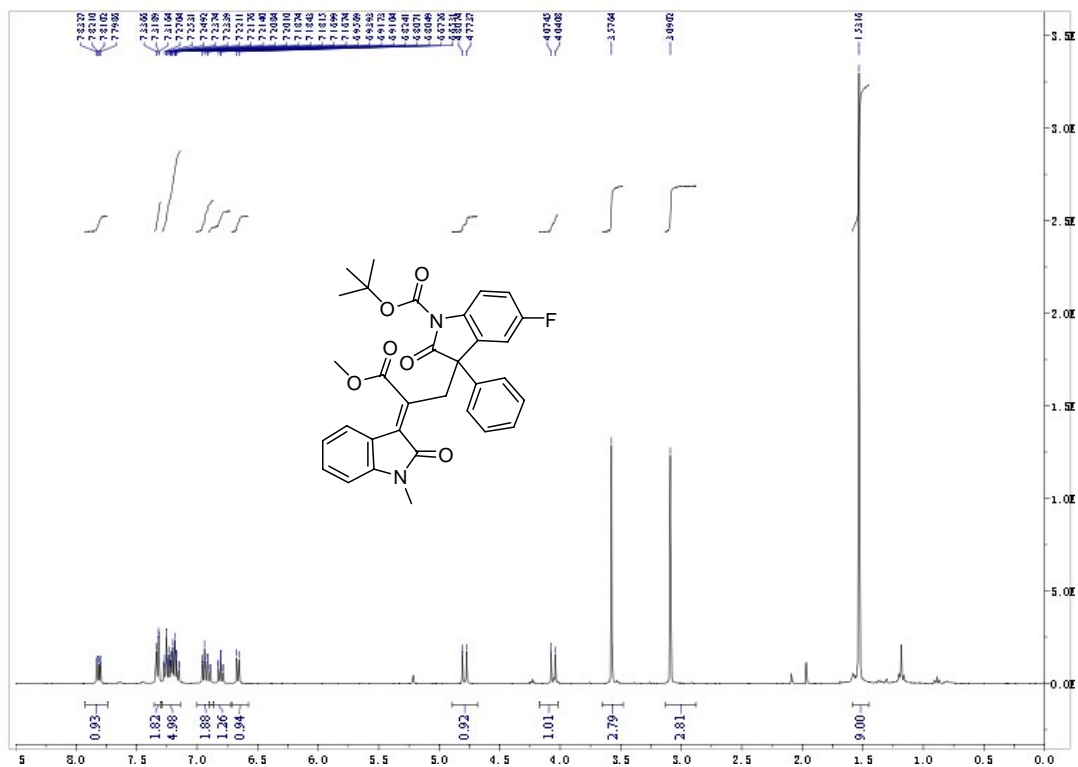


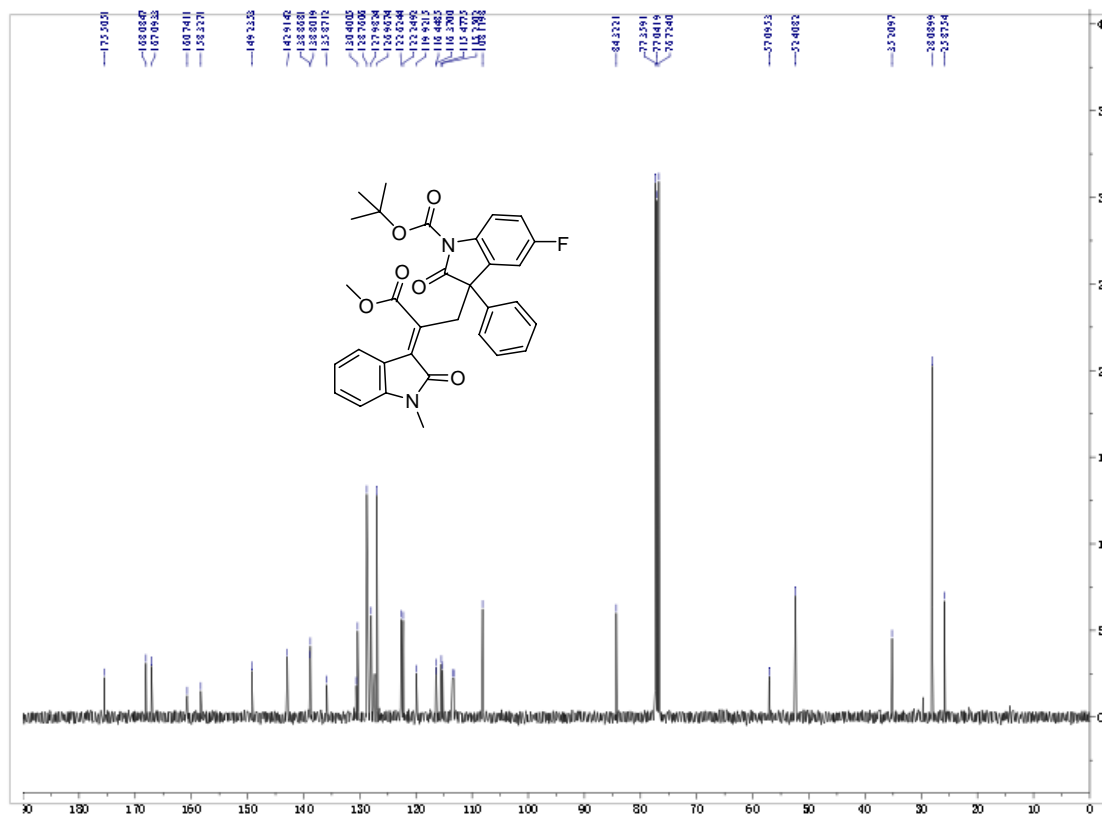
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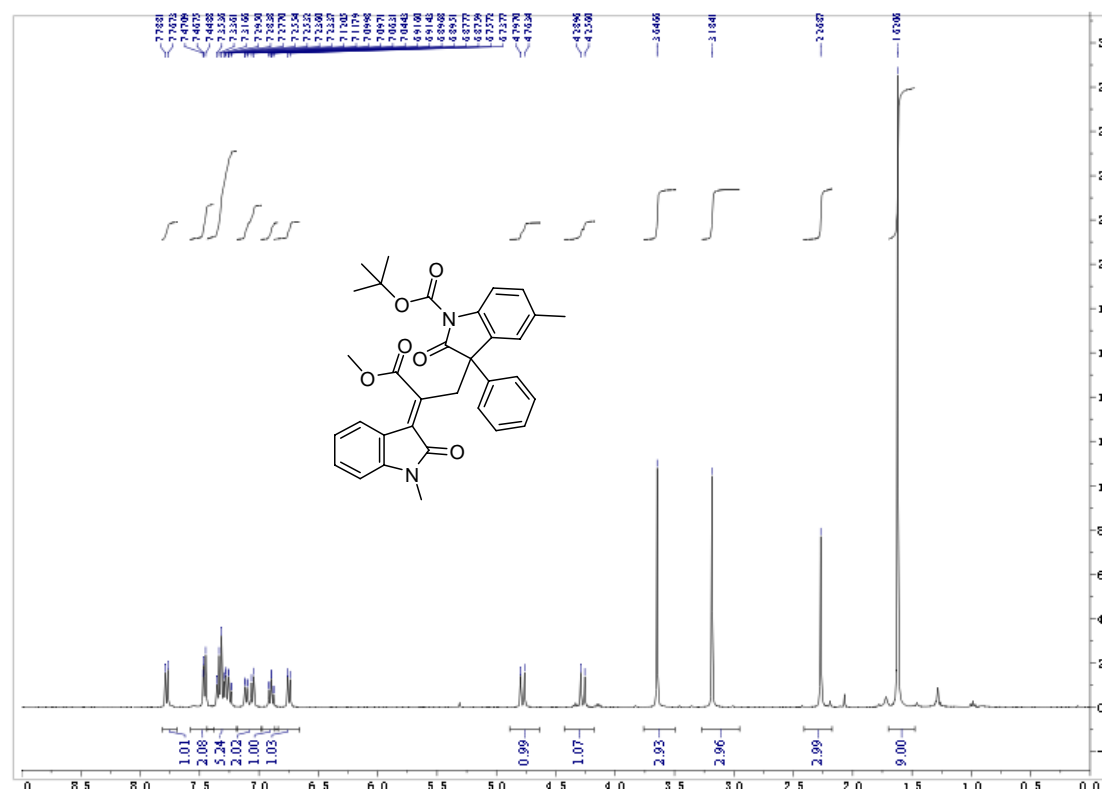


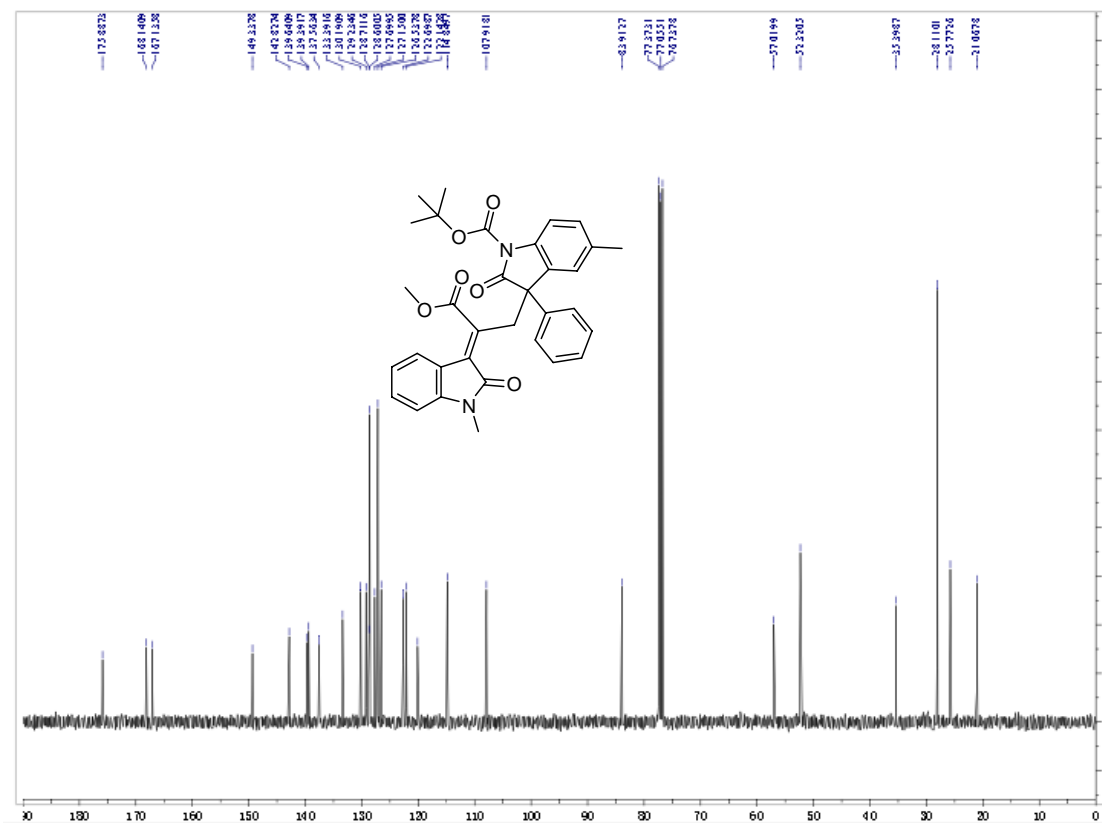
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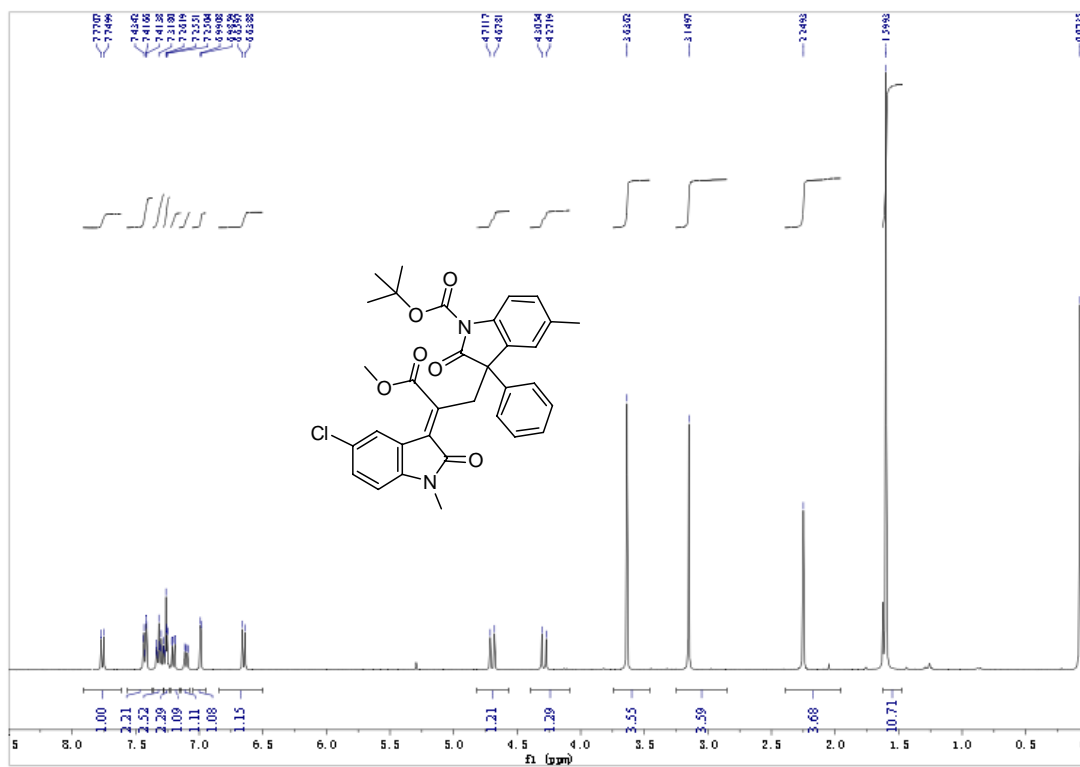


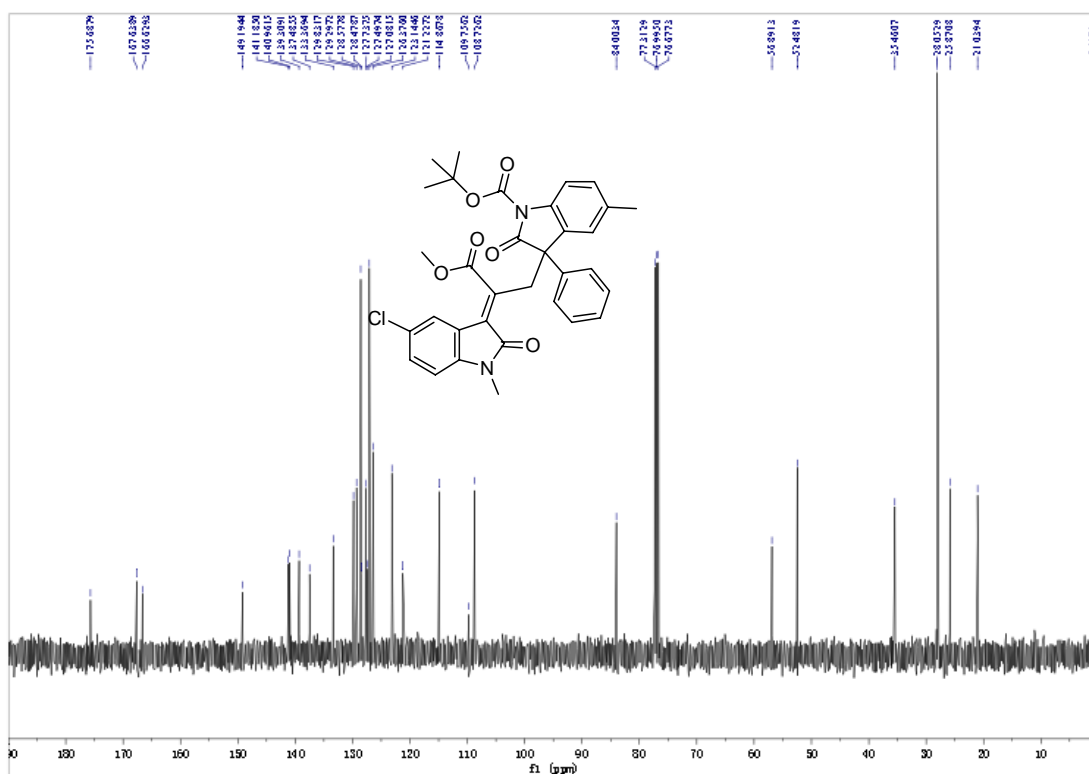
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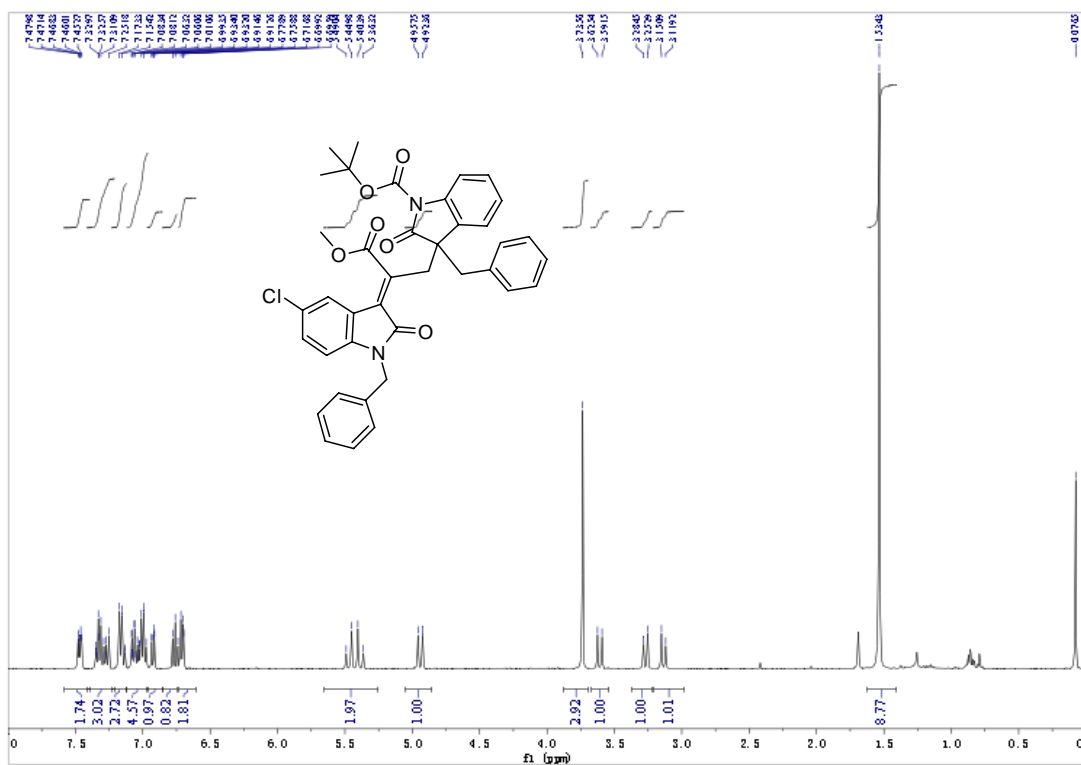


^1H and ^{13}C NMR of 3eb





^1H and ^{13}C NMR of 3fh



C13-HX-030
HX-030

