

# Supporting Information File 2

for

## Asymmetric Ugi 3CR on isatin-derived ketimine: synthesis of chiral 3,3-disubstituted 3-aminoxindole derivatives

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### General methods and copies of NMR spectra for all new compounds

#### Table of Contents

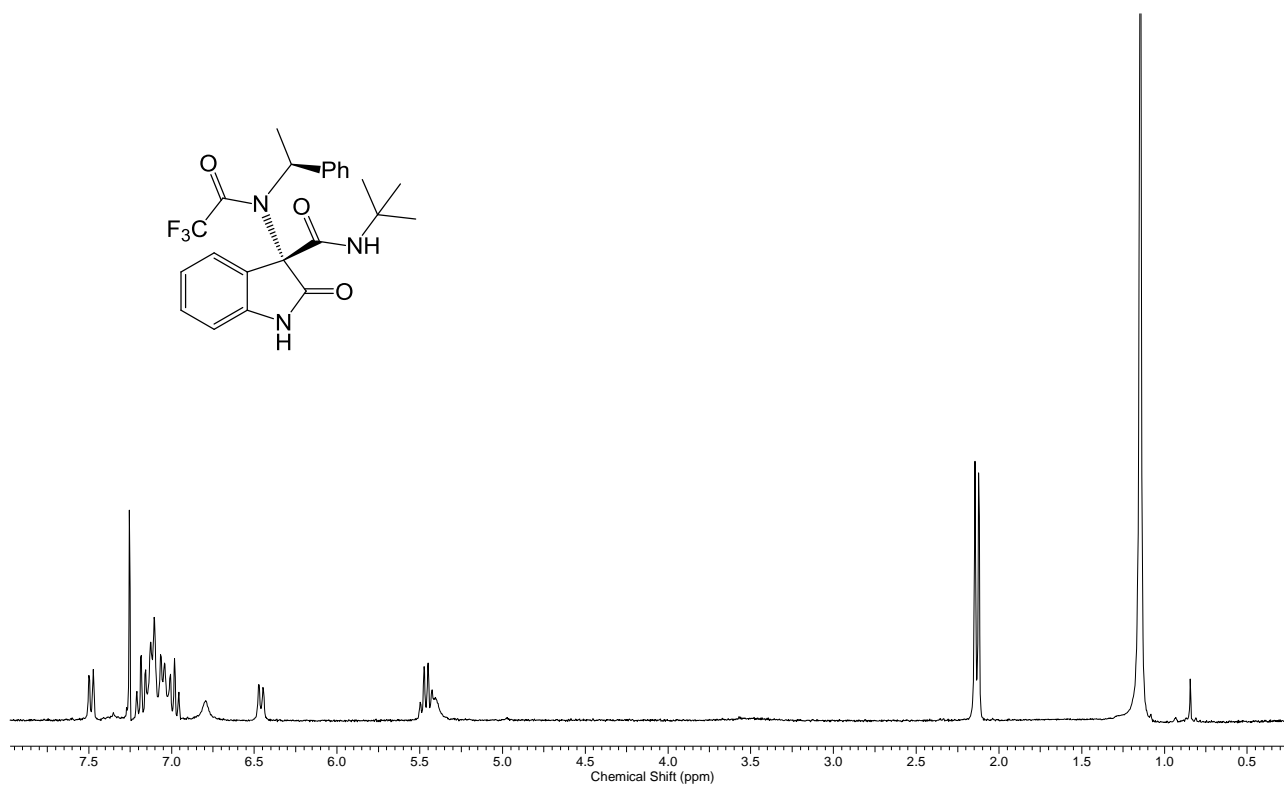
General methods.....	S2
<sup>1</sup> H NMR and <sup>13</sup> C NMR (APT) spectra of compounds <b>4–18</b> .....	S3

## General methods

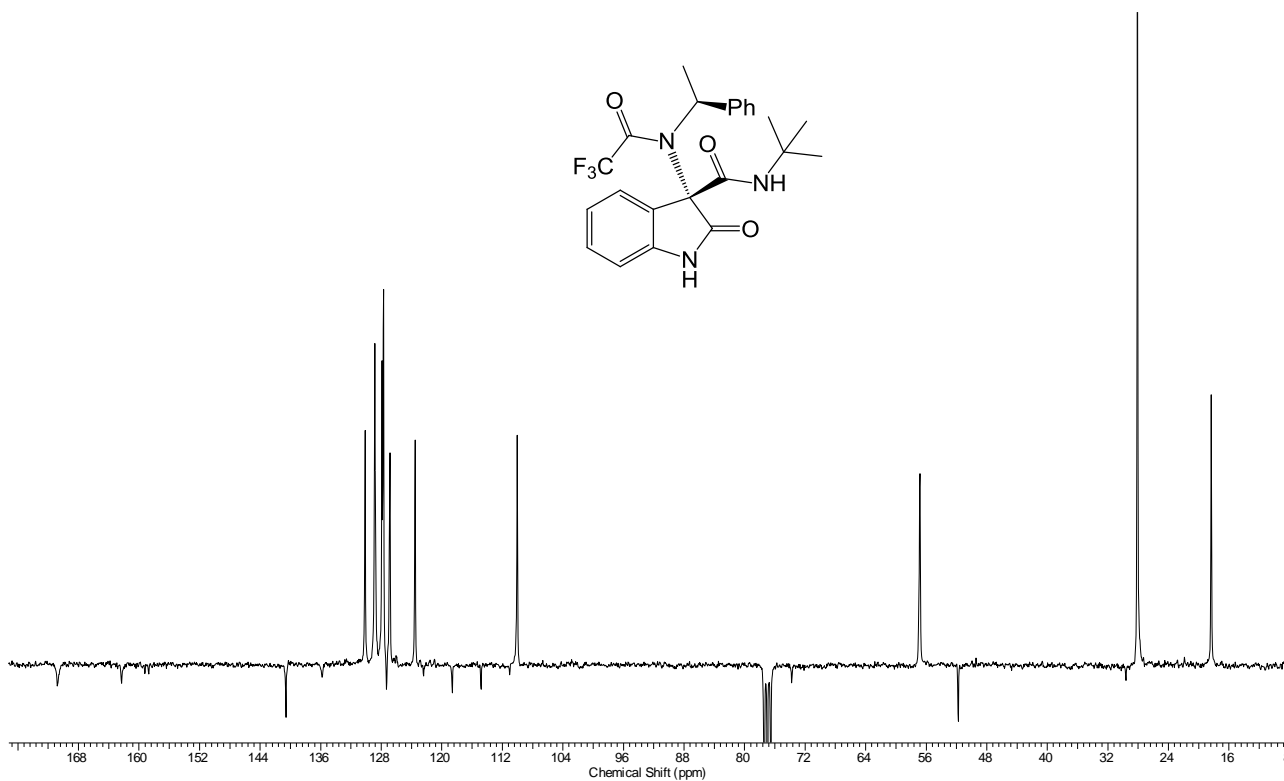
All solvents were distilled and properly dried, if necessary, prior to use. All chemicals were purchased from commercial sources and used directly, unless indicated otherwise. All reactions were run under N<sub>2</sub>, unless otherwise indicated. All reactions were monitored by thin layer chromatography (TLC) on precoated silica gel 60 F254; spots were visualized with UV light or by treatment with 1% aqueous KMnO<sub>4</sub> solution. Products were purified by flash chromatography on silica gel 60 (230–400 mesh).

NMR spectra were recorded with 400 MHz spectrometers. Chemical shifts ( $\delta$ ) are expressed in ppm relative to TMS at  $\delta = 0$  ppm for <sup>1</sup>H NMR and relative to CDCl<sub>3</sub> at  $\delta = 77.16$  ppm for <sup>13</sup>C NMR. <sup>13</sup>C NMR spectra have been recorded using the APT pulse sequence; the signals of CH and CH<sub>3</sub> are positive while CH<sub>2</sub> and quarternary carbons are negative. High-resolution mass spectra were recorded with FT-ICR (Fourier Transform Ion Cyclotron Resonance) instruments, equipped with ESI or EI source.

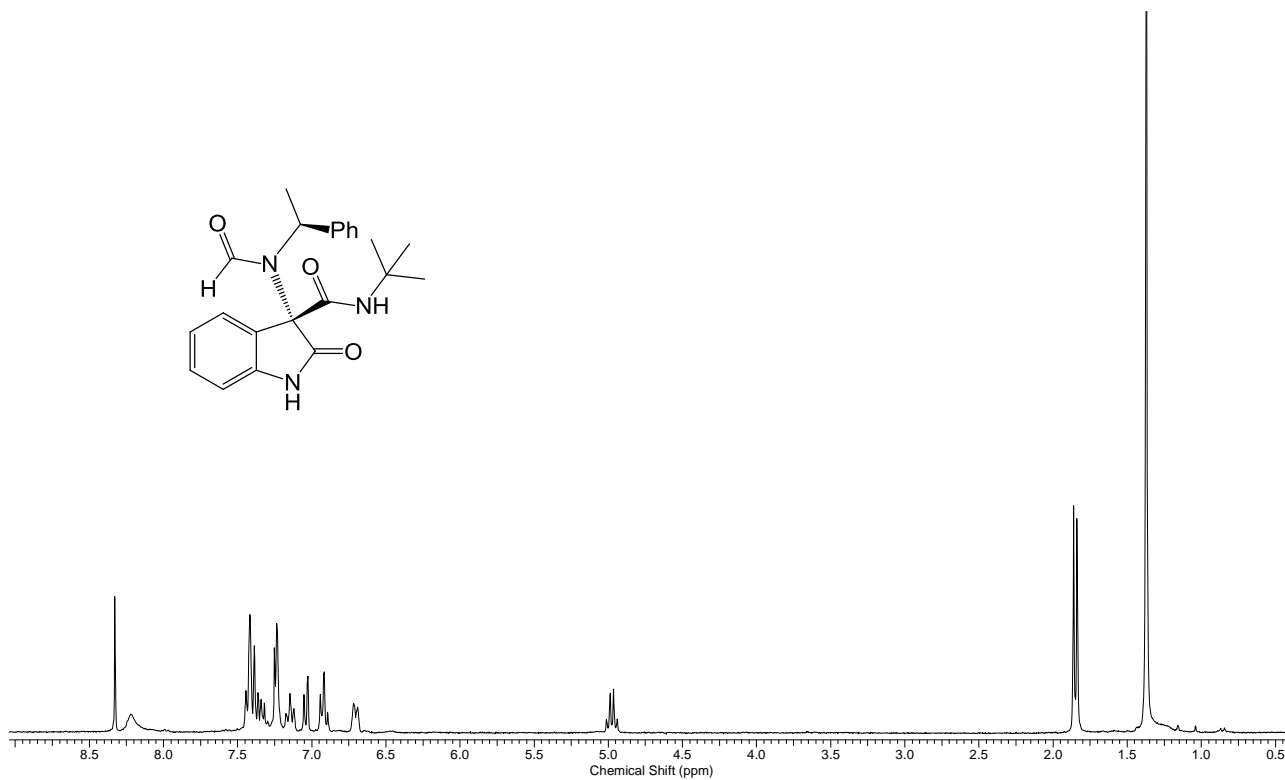
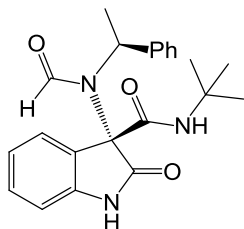
Compound **4a**:  $^1\text{H}$  NMR (300 MHz,  $\text{CDCl}_3$ )



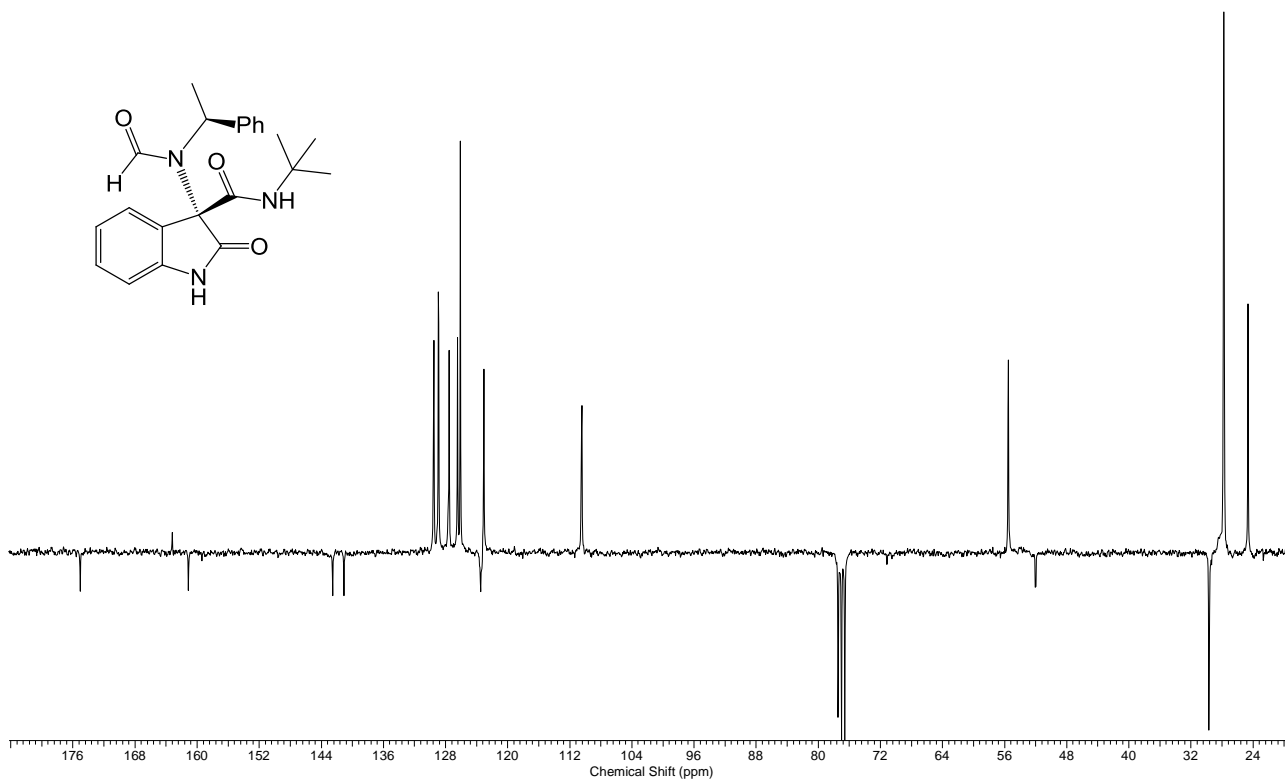
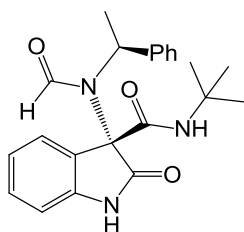
Compound **4a**:  $^{13}\text{C}$  NMR (75 MHz,  $\text{CDCl}_3$ )



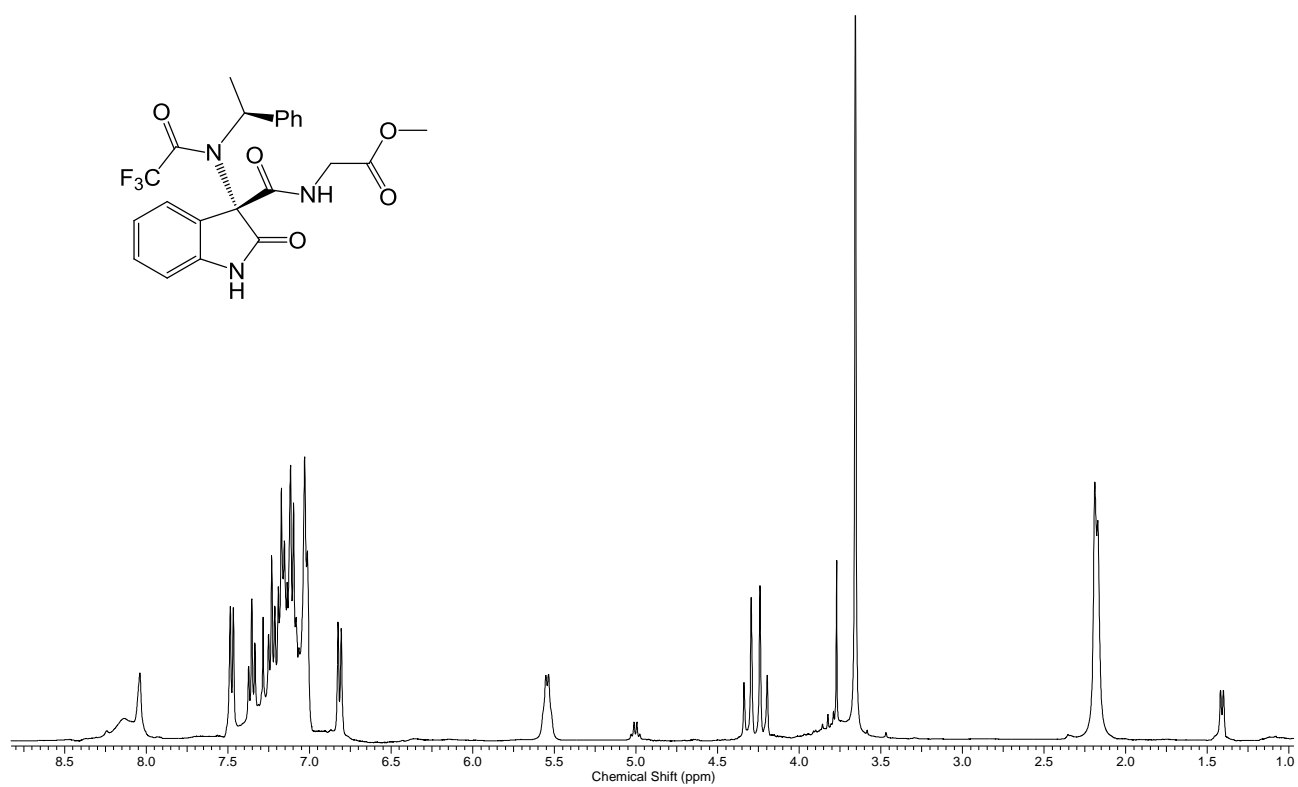
Compound **5a**:  $^1\text{H}$  NMR (300 MHz,  $\text{CDCl}_3$ )



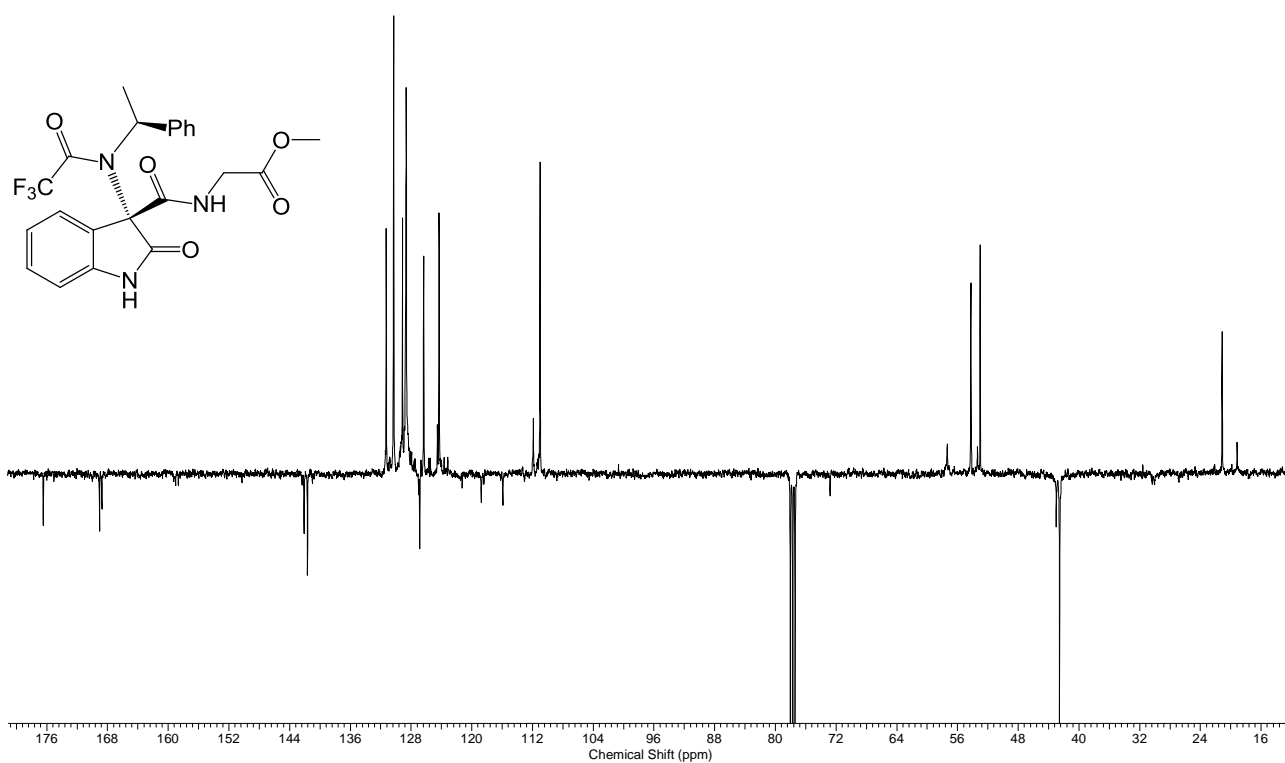
Compound **5a**:  $^{13}\text{C}$  NMR (75 MHz,  $\text{CDCl}_3$ )



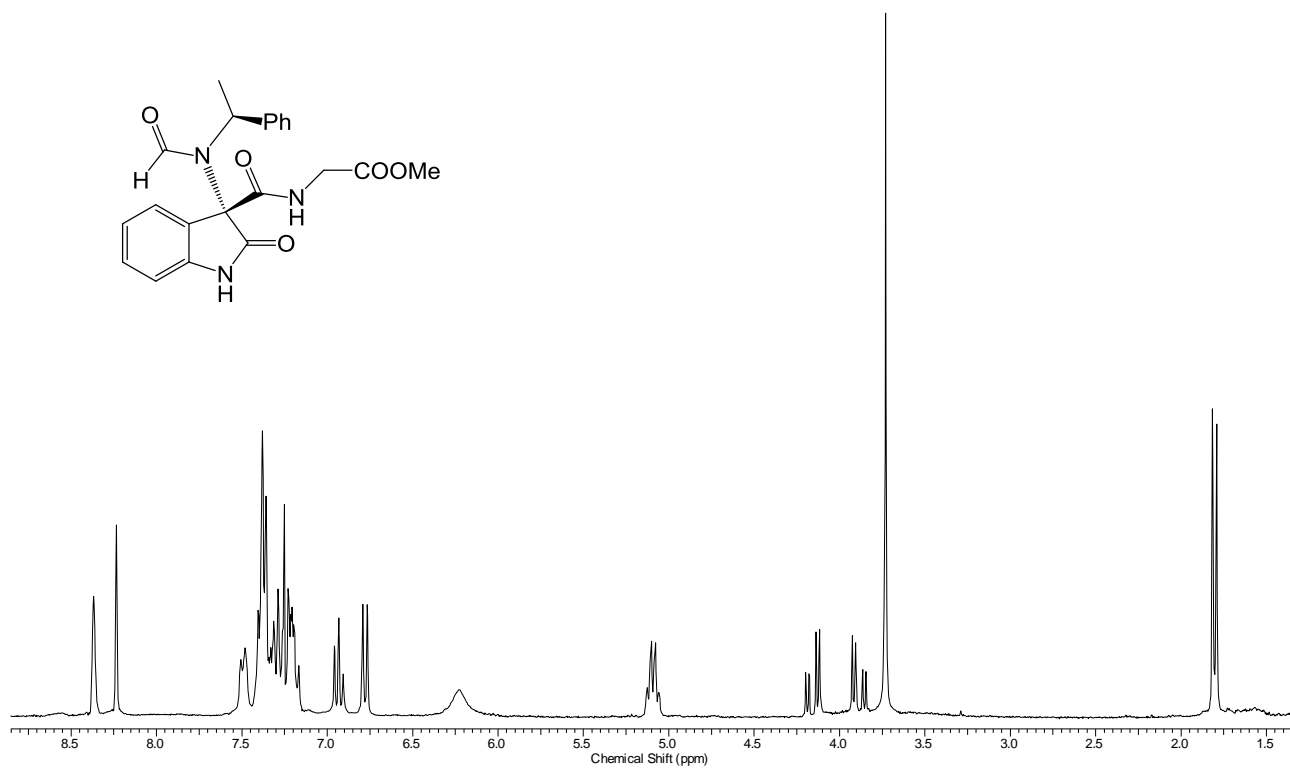
Compound 6:  $^1\text{H}$  NMR (400 MHz,  $\text{CDCl}_3$ , diastereoisomeric mixture)



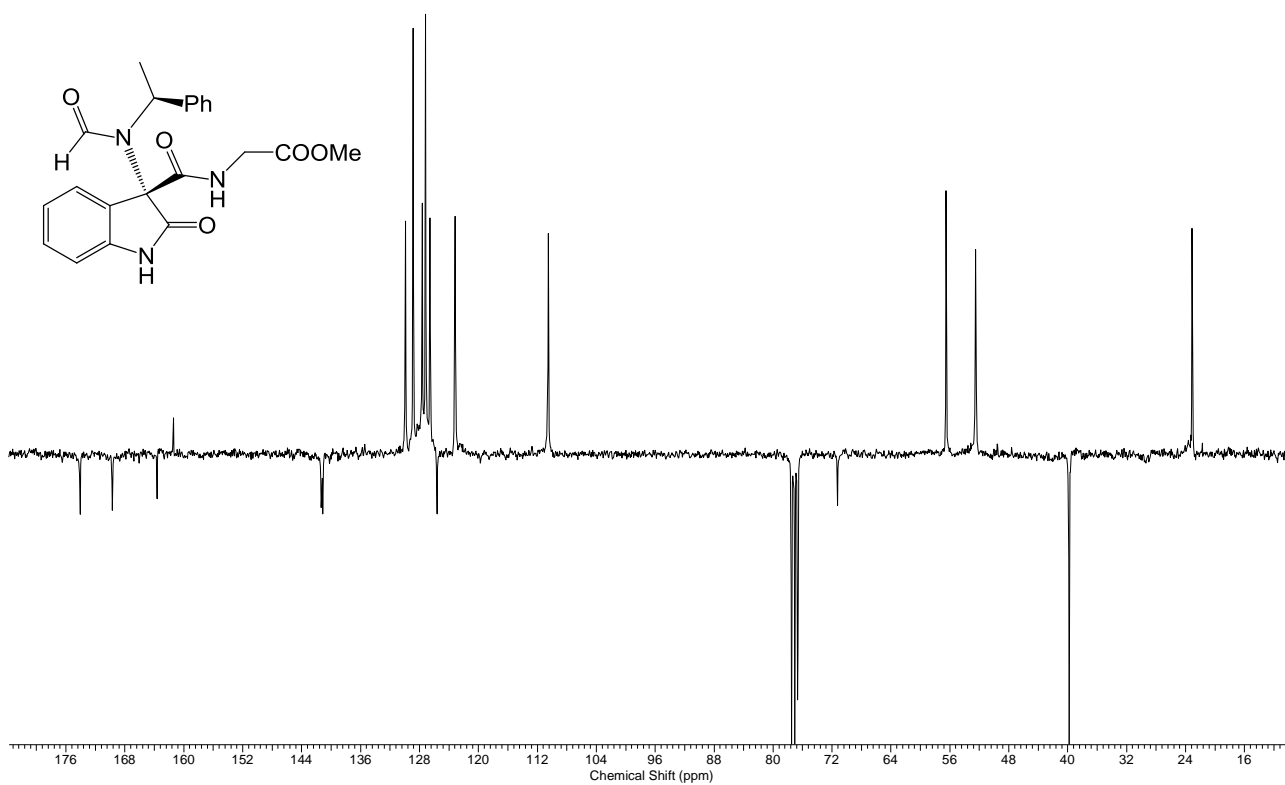
Compound 6:  $^{13}\text{C}$  NMR (100 MHz,  $\text{CDCl}_3$ , diastereoisomeric mixture)



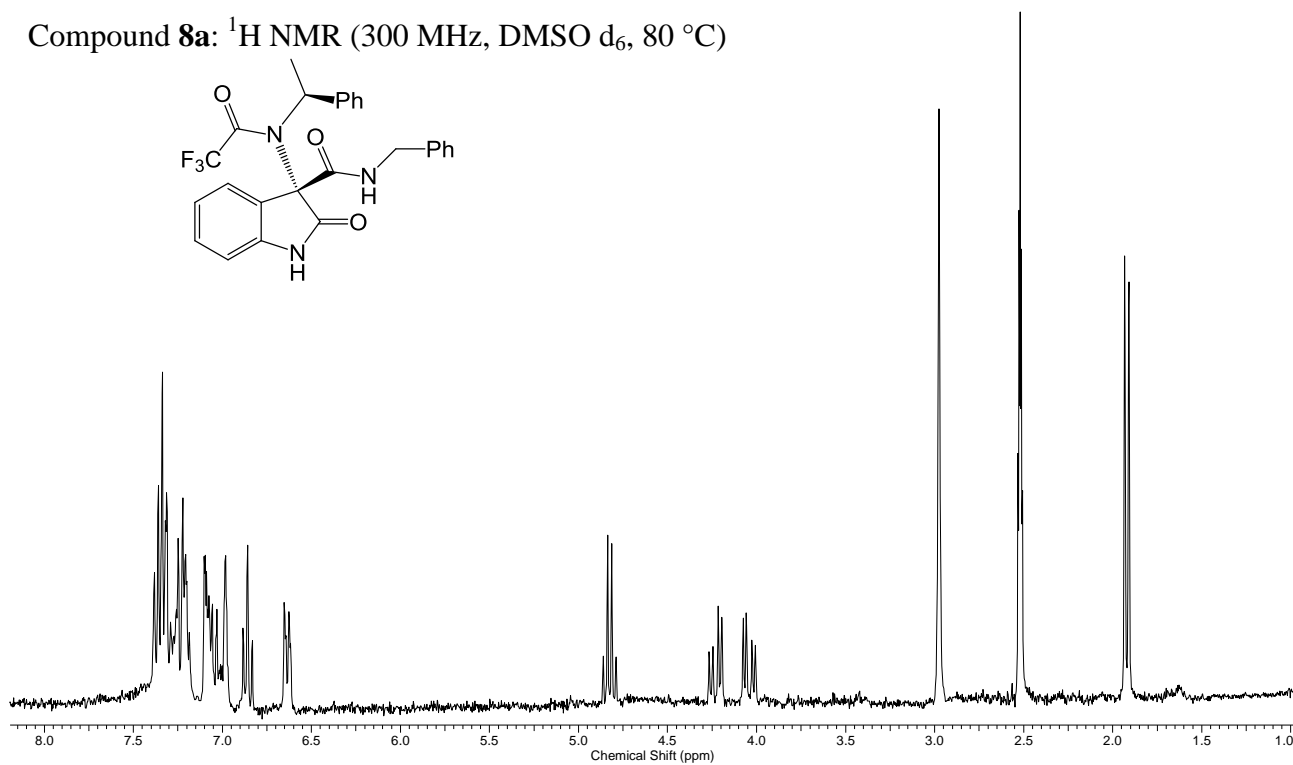
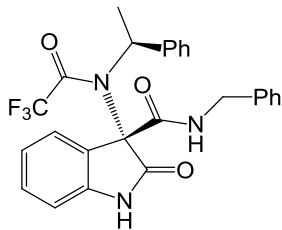
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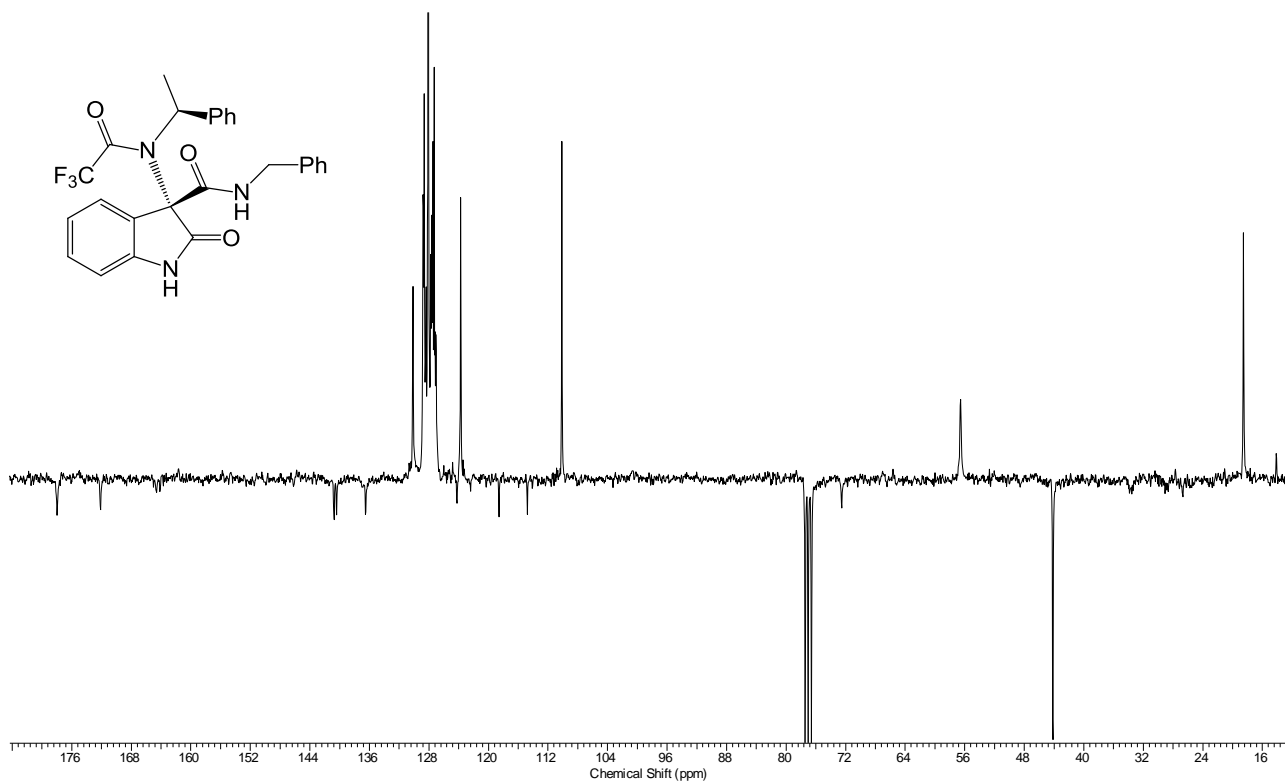
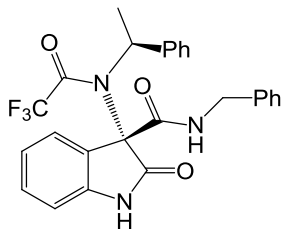
Compound **7a**:  $^{13}\text{C}$  NMR (75 MHz,  $\text{CDCl}_3$ )



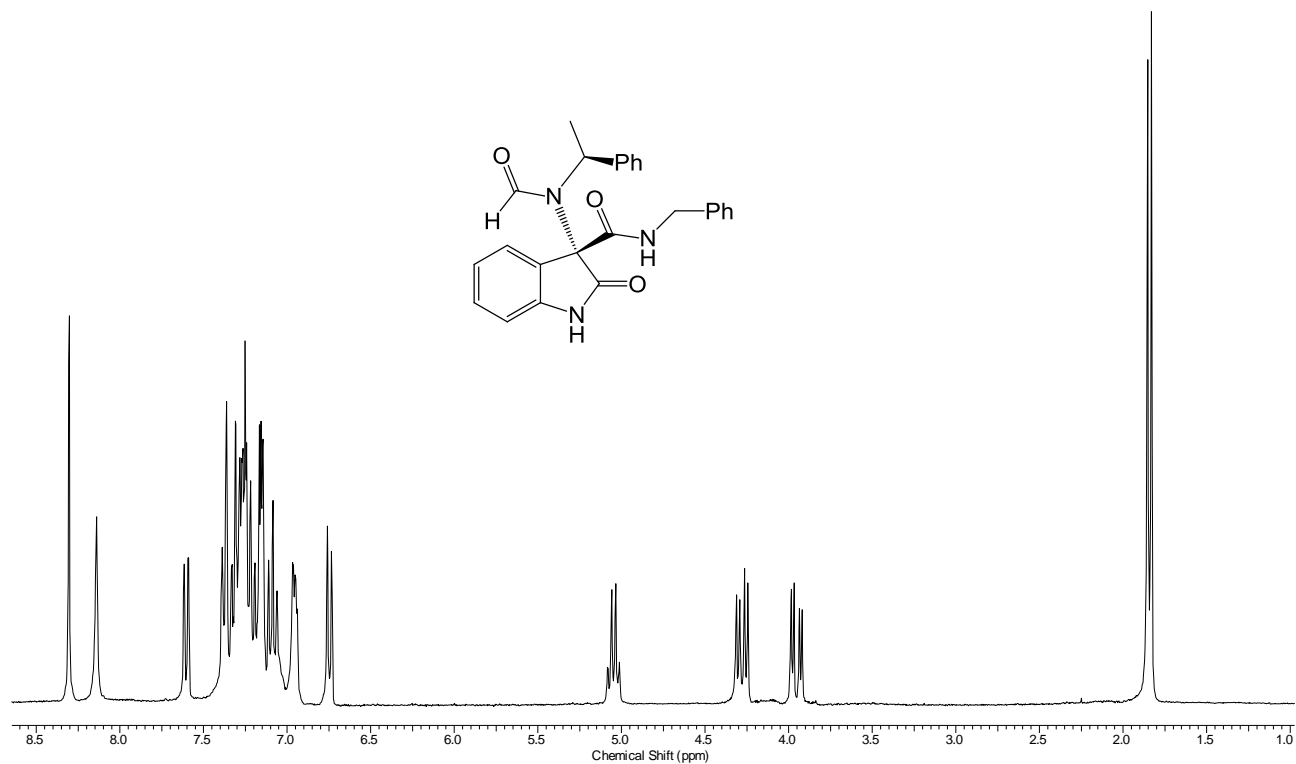
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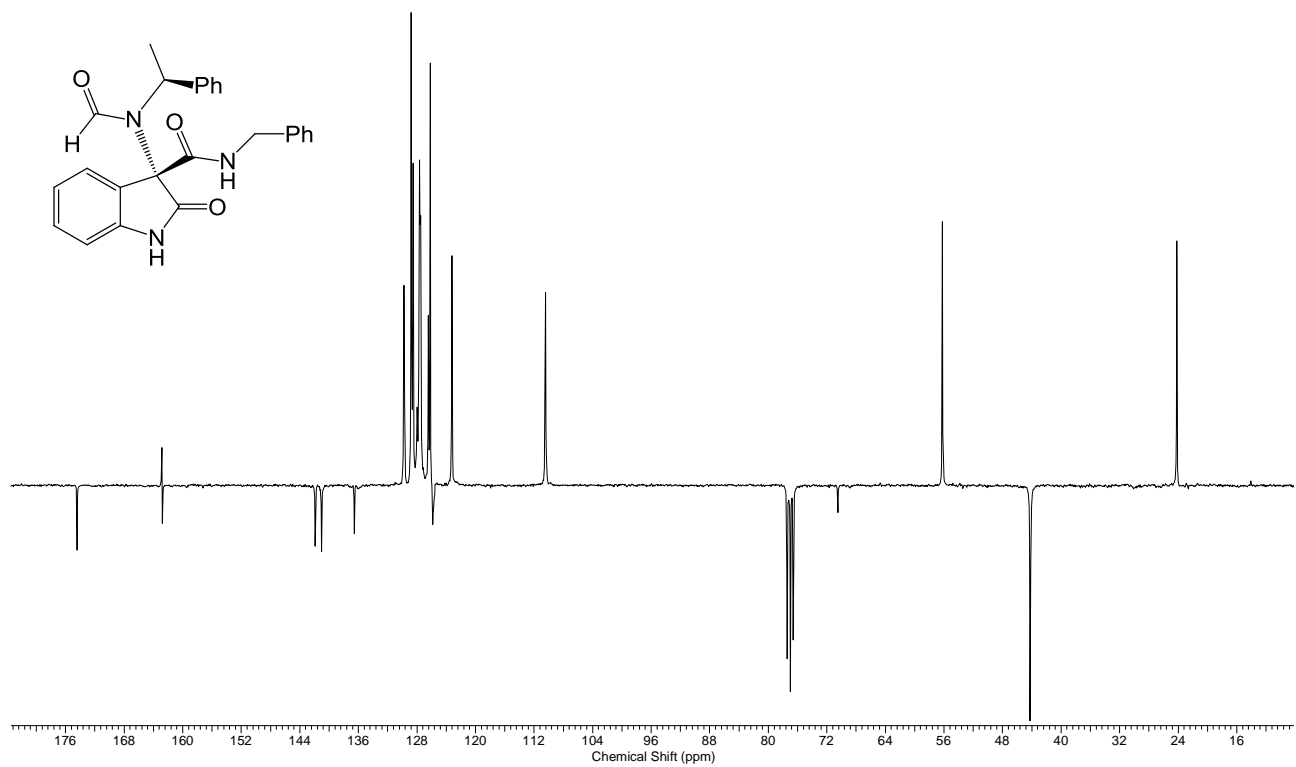
Compound **8a**:  $^{13}\text{C}$  NMR (75 MHz,  $\text{CDCl}_3$ )



Compound **9a**:  $^1\text{H}$  NMR (300 MHz,  $\text{CDCl}_3$ )

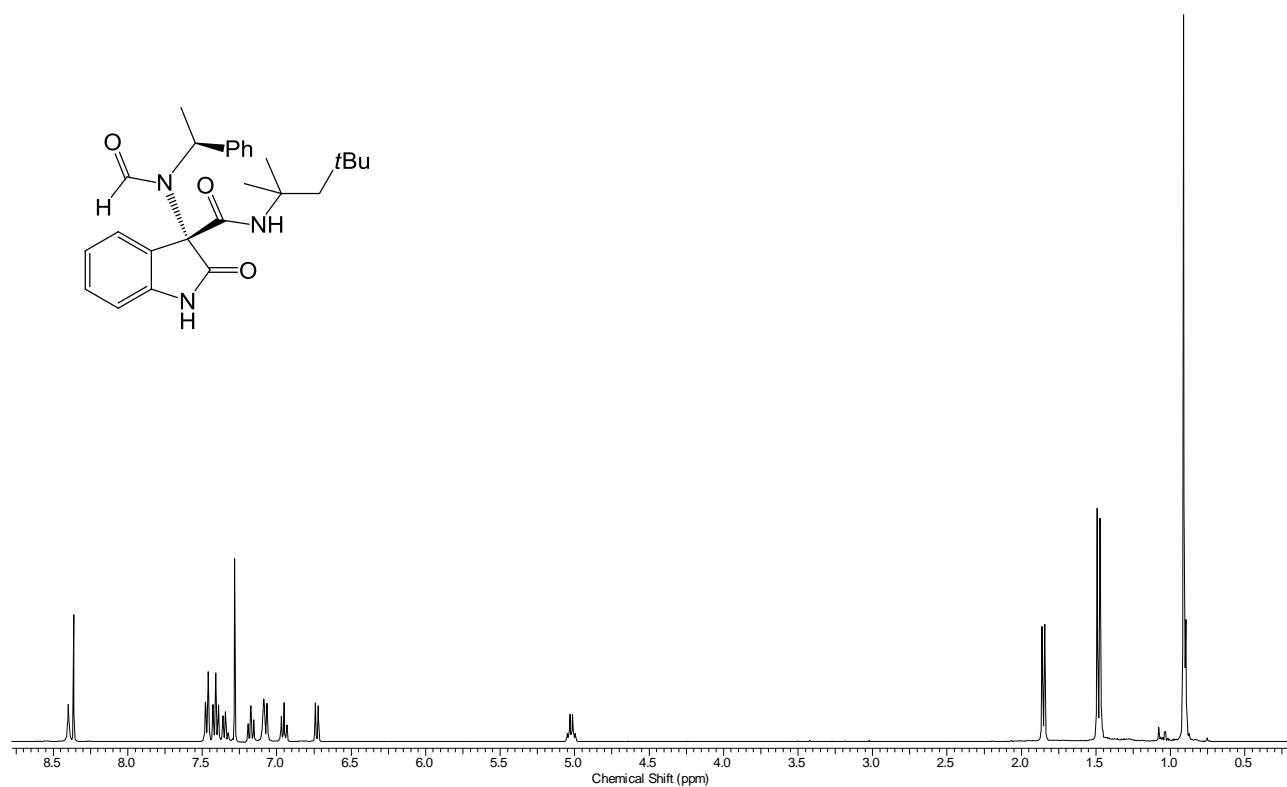


Compound **9a**:  $^{13}\text{C}$  NMR (75 MHz,  $\text{CDCl}_3$ )

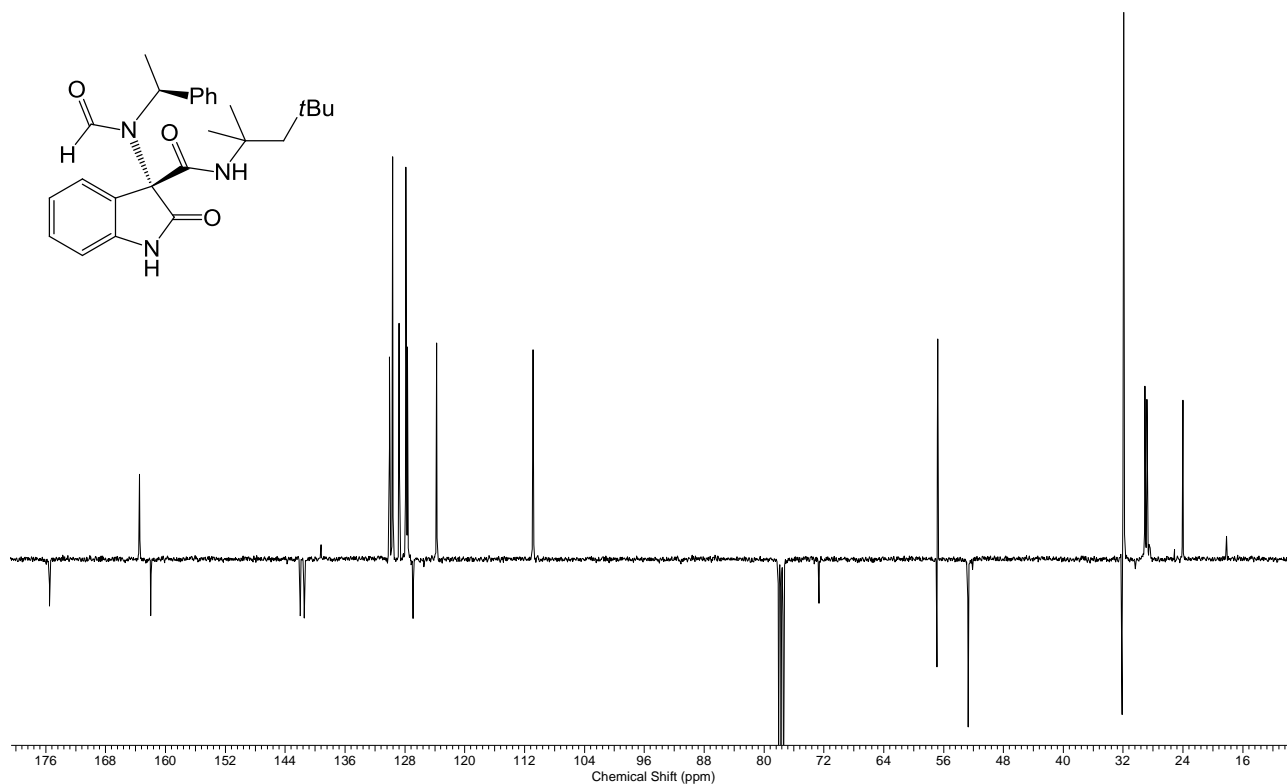




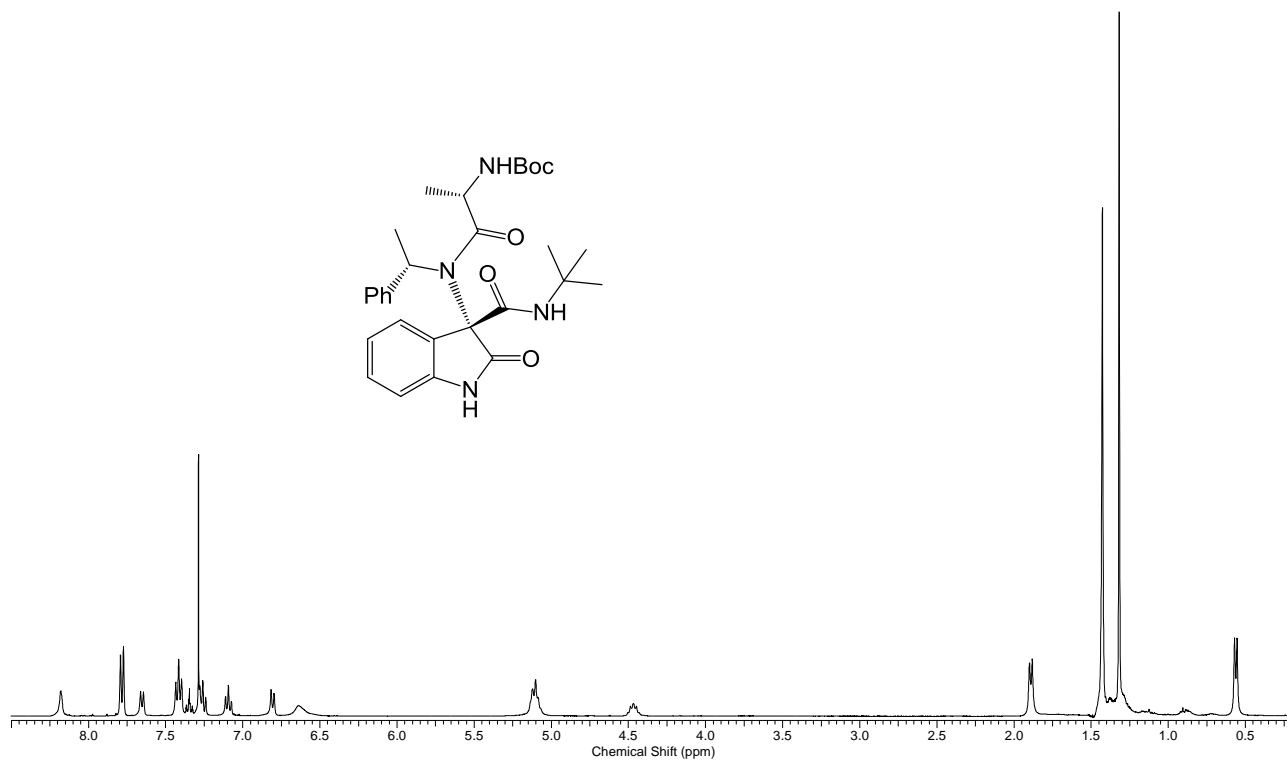
Compound **10a**:  $^1\text{H}$  NMR (400 MHz,  $\text{CDCl}_3$ )



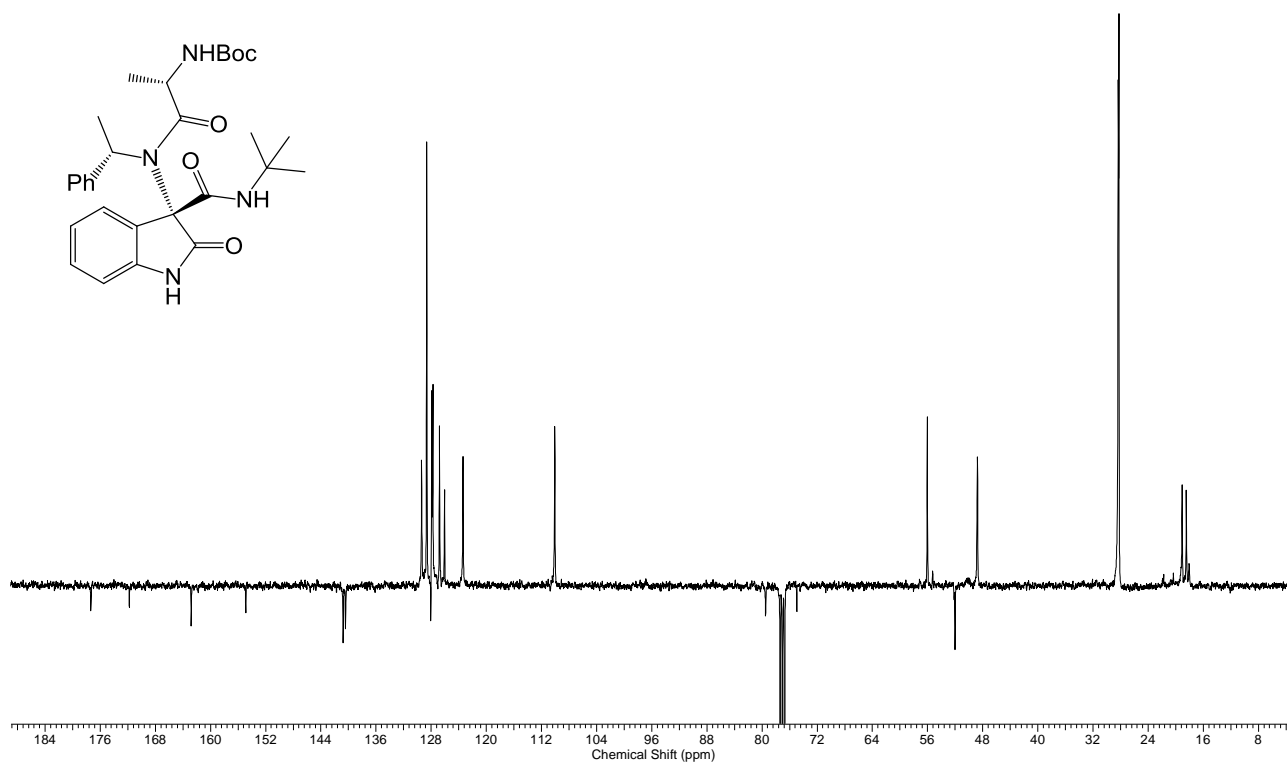
Compound **10a**:  $^{13}\text{C}$  NMR (100 MHz,  $\text{CDCl}_3$ )



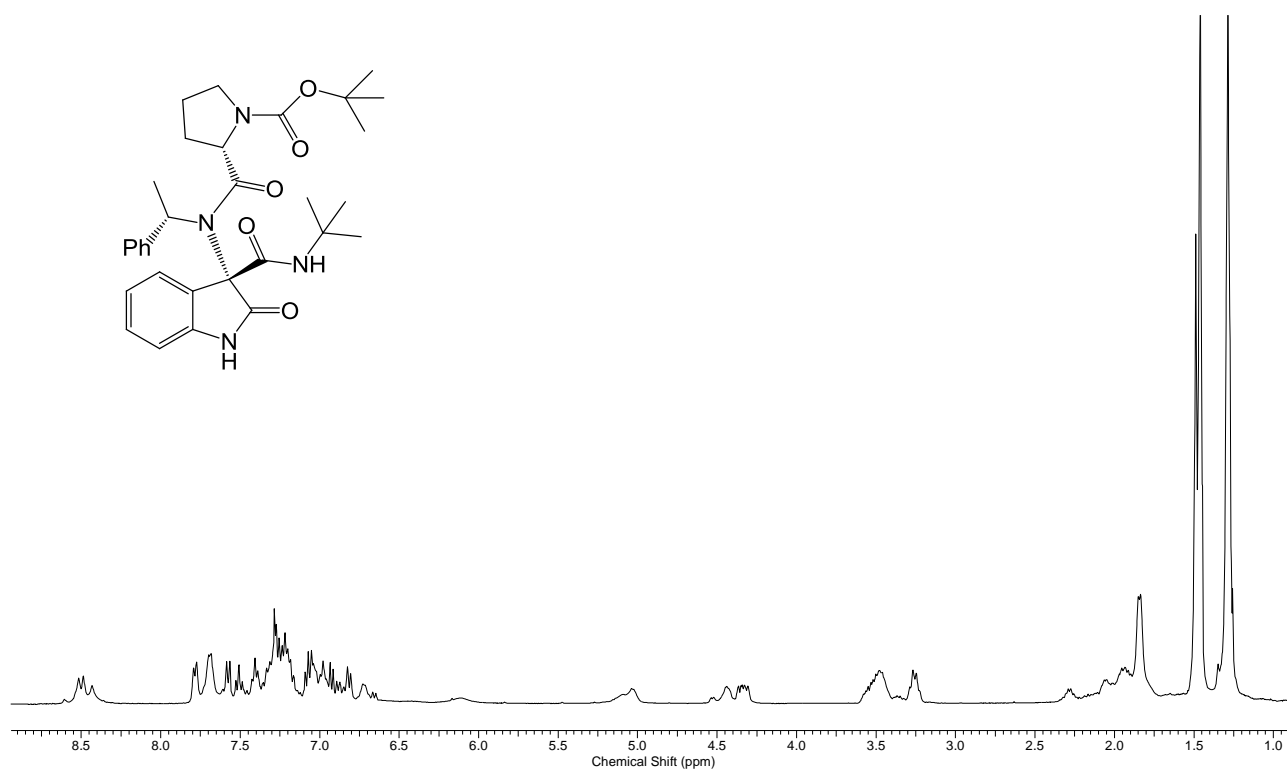
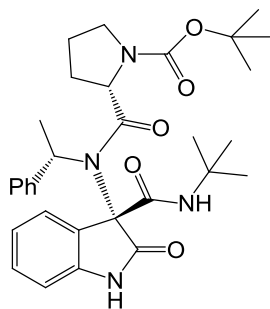
Compound **11a**:  $^1\text{H}$  NMR (400 MHz,  $\text{CDCl}_3$ )



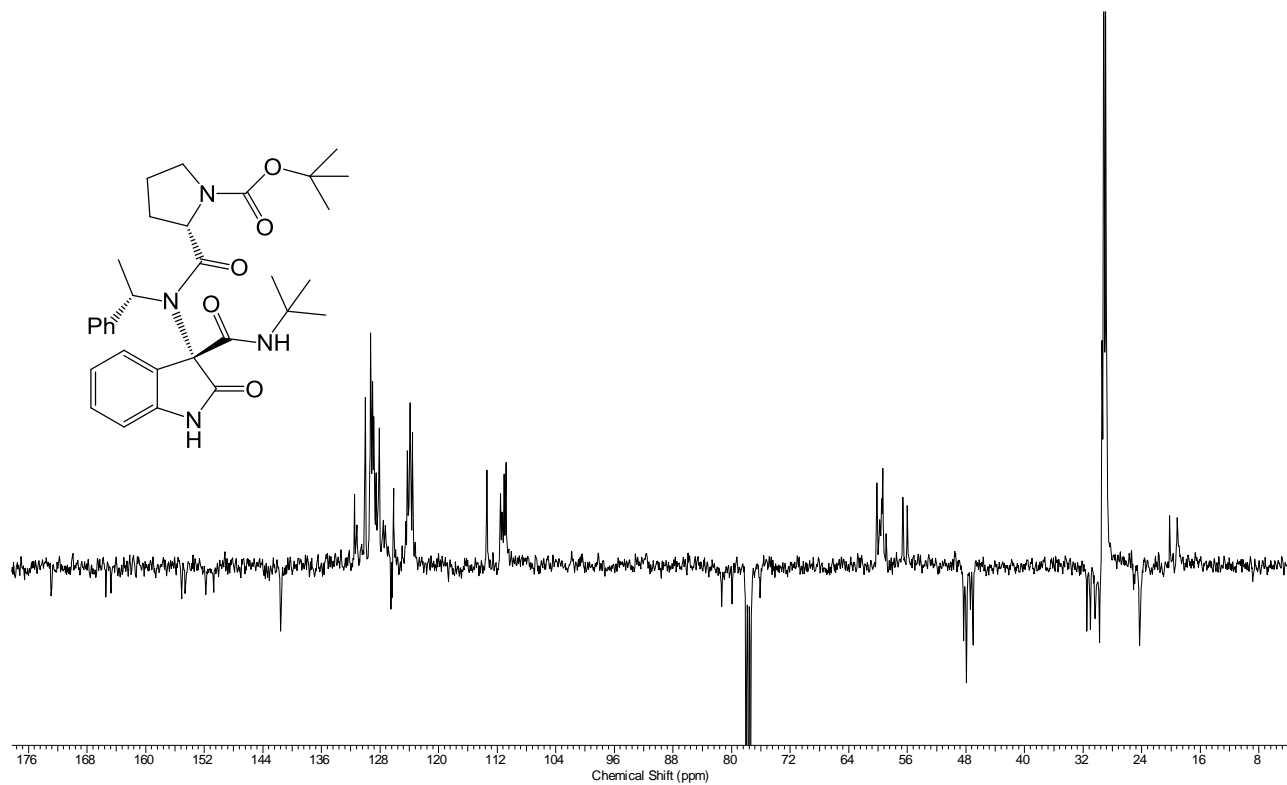
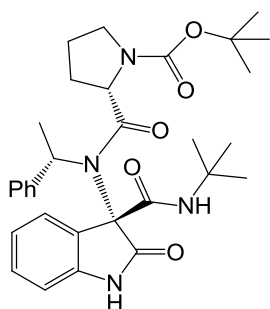
Compound **11a**:  $^{13}\text{C}$  NMR (100 MHz,  $\text{CDCl}_3$ )



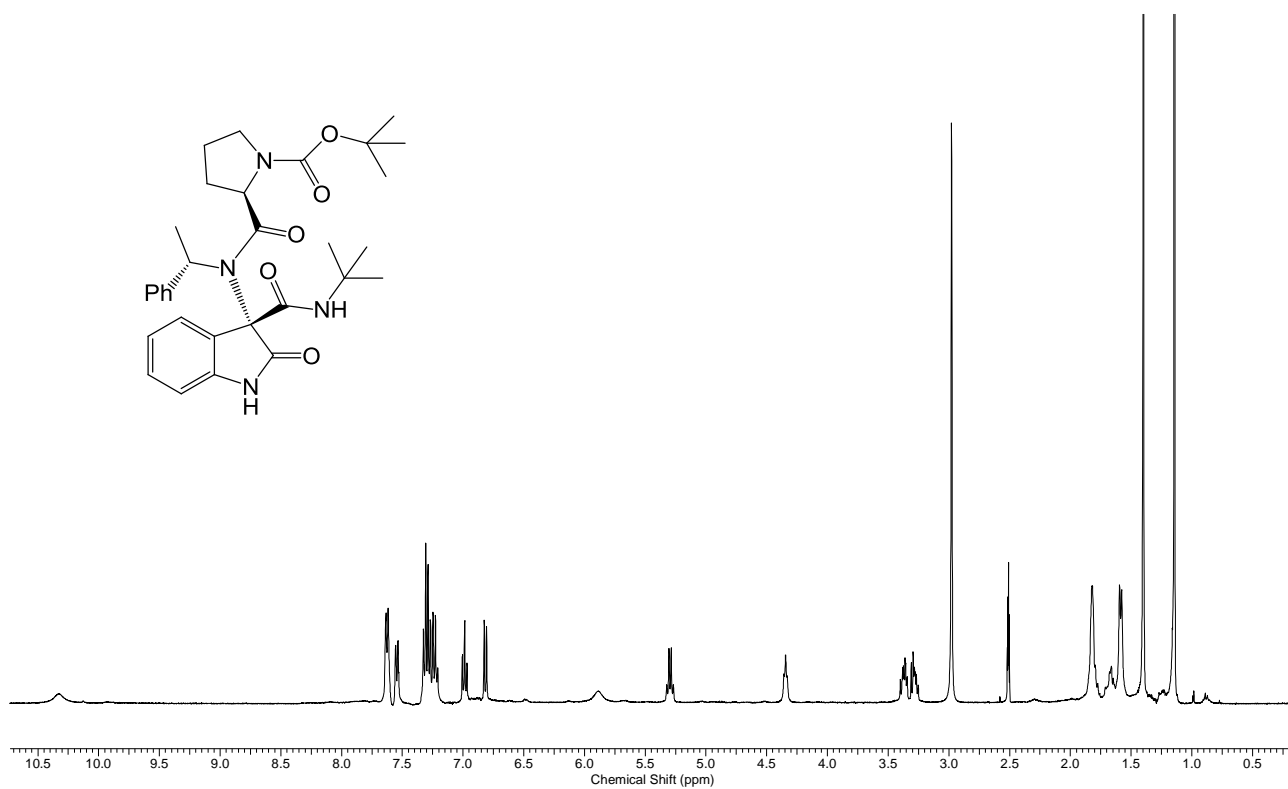
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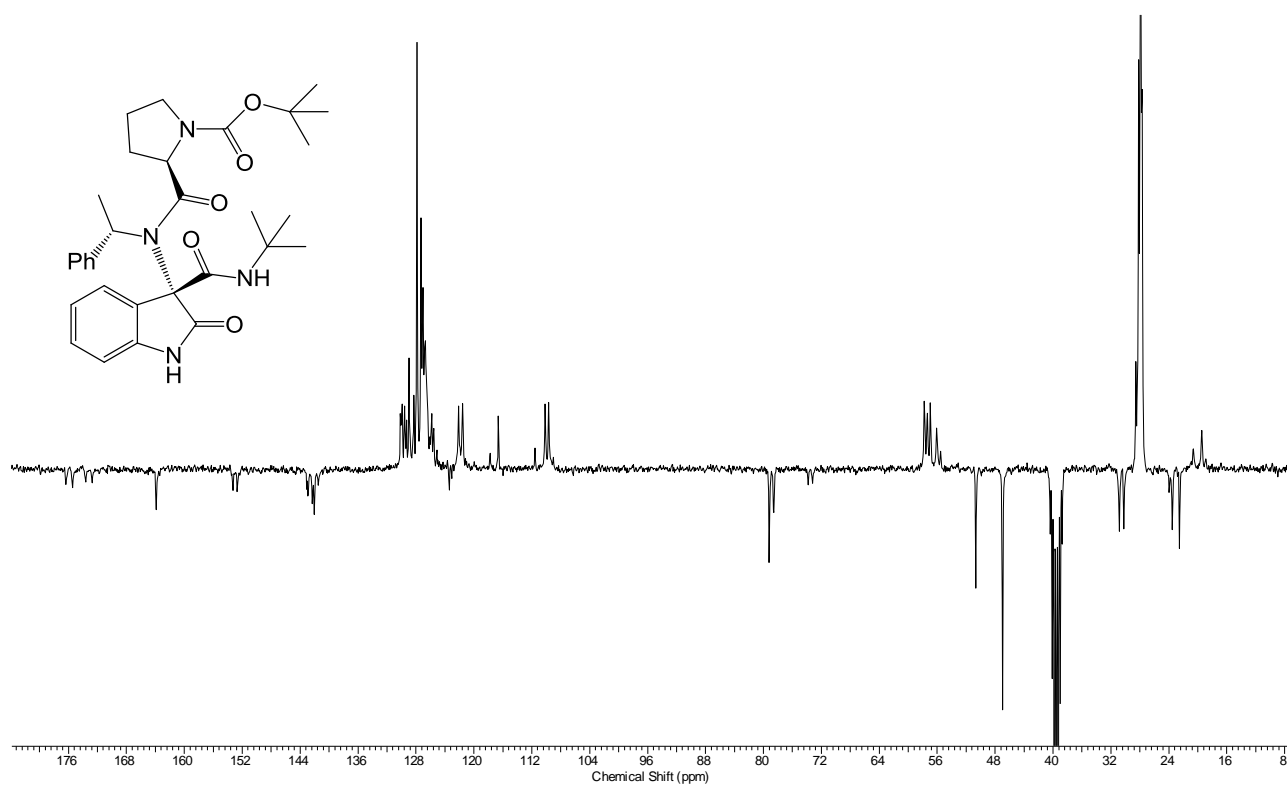
Compound **12a**:  $^{13}\text{C}$  NMR (100 MHz,  $\text{CDCl}_3$ )



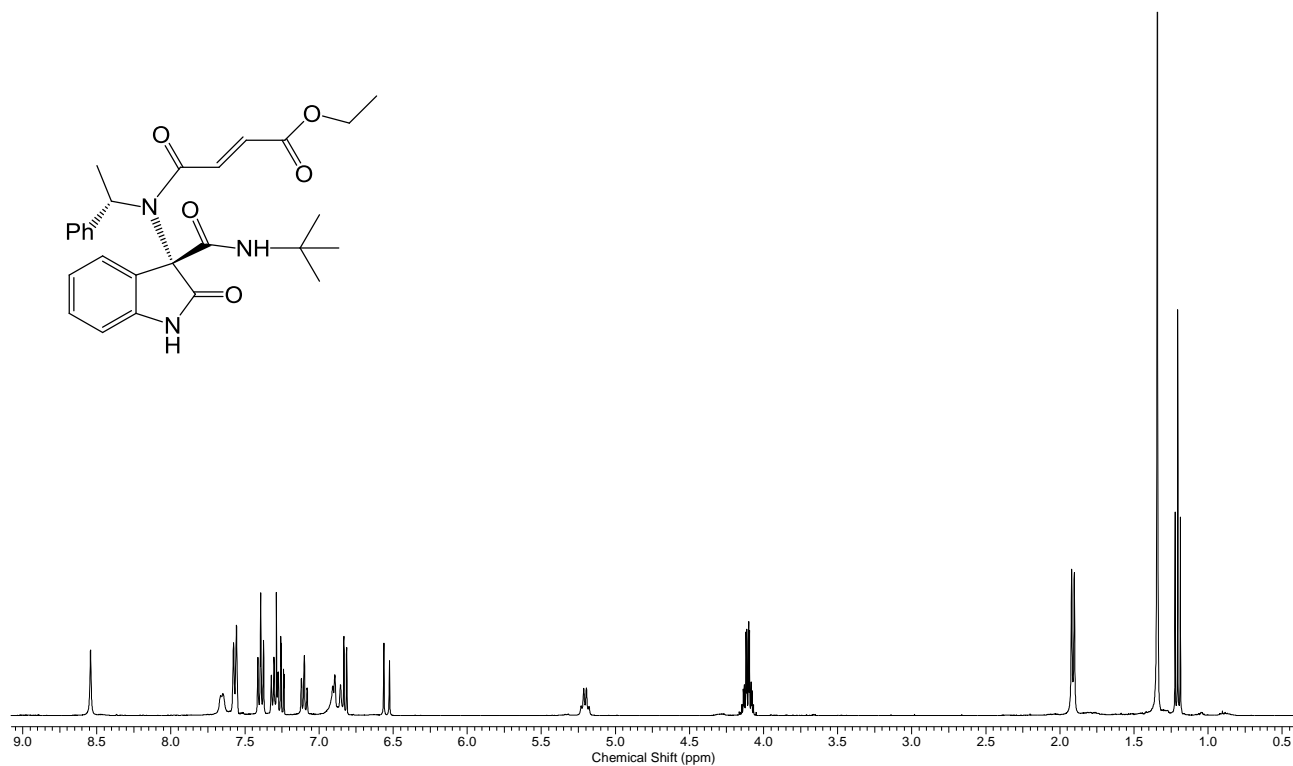
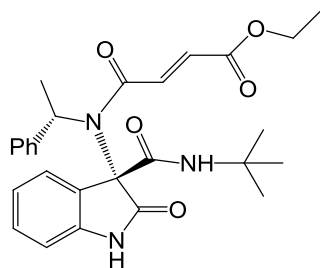
Compound **13a**:  $^1\text{H}$  NMR (400 MHz,  $\text{DMSO-}d_6$ , 100 °C)



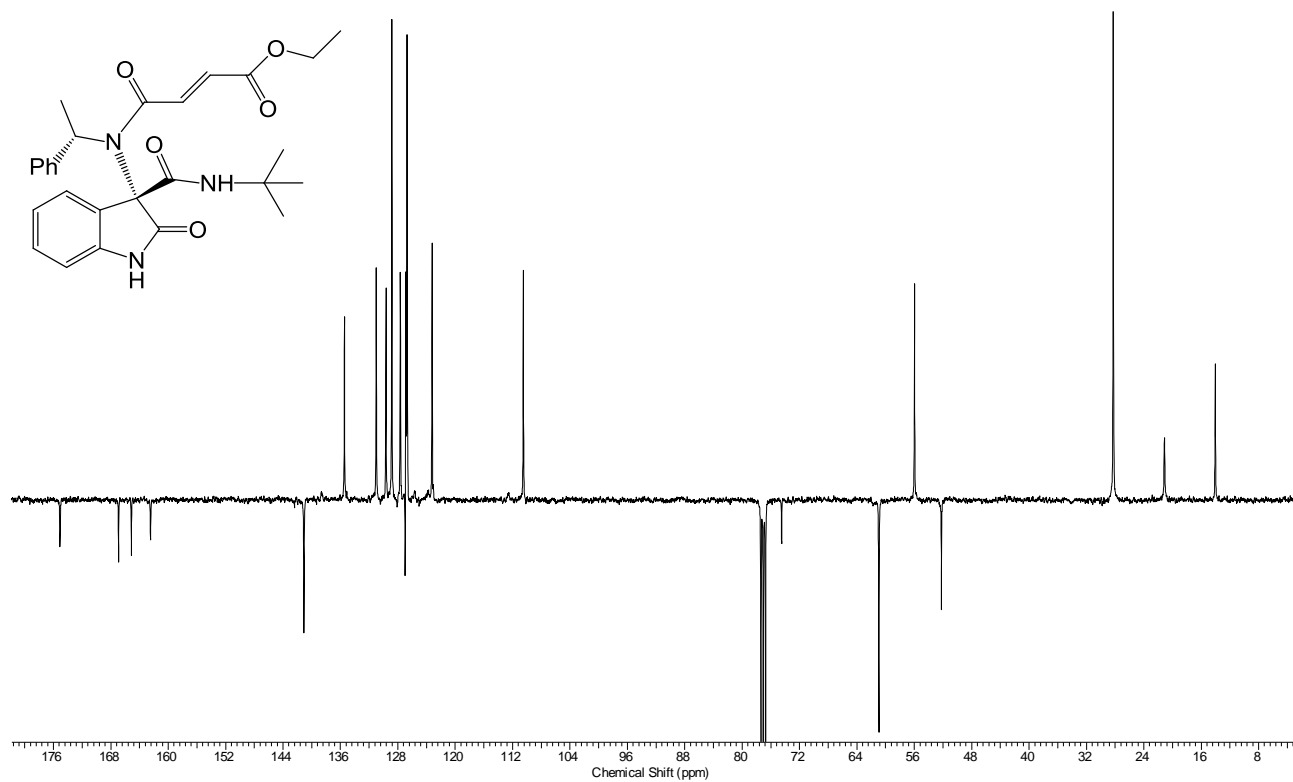
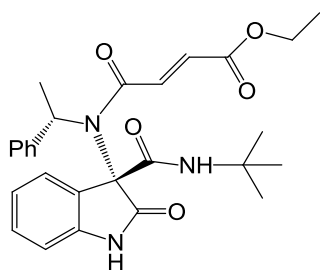
Compound **13a**:  $^{13}\text{C}$  NMR (75 MHz,  $\text{DMSO-}d_6$ , rotameric mixture)



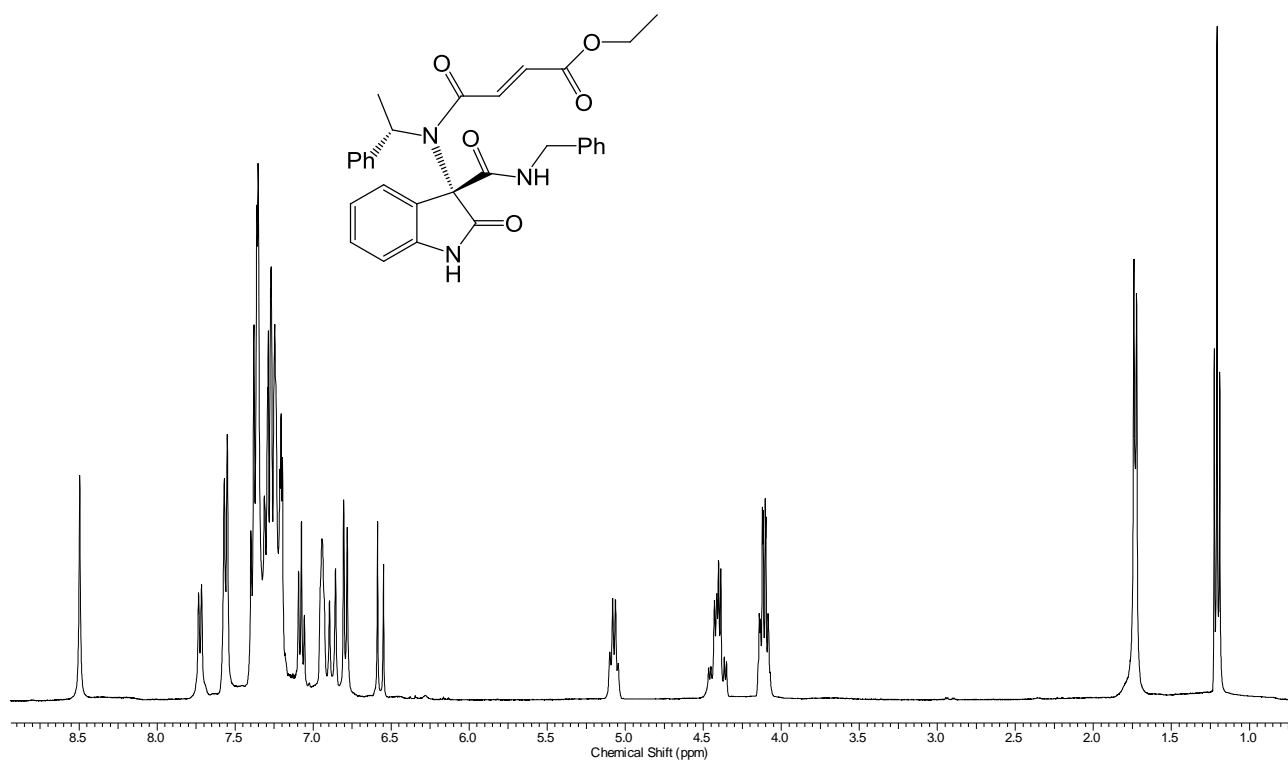
Compound **14a**:  $^1\text{H}$  NMR (400 MHz,  $\text{CDCl}_3$ )



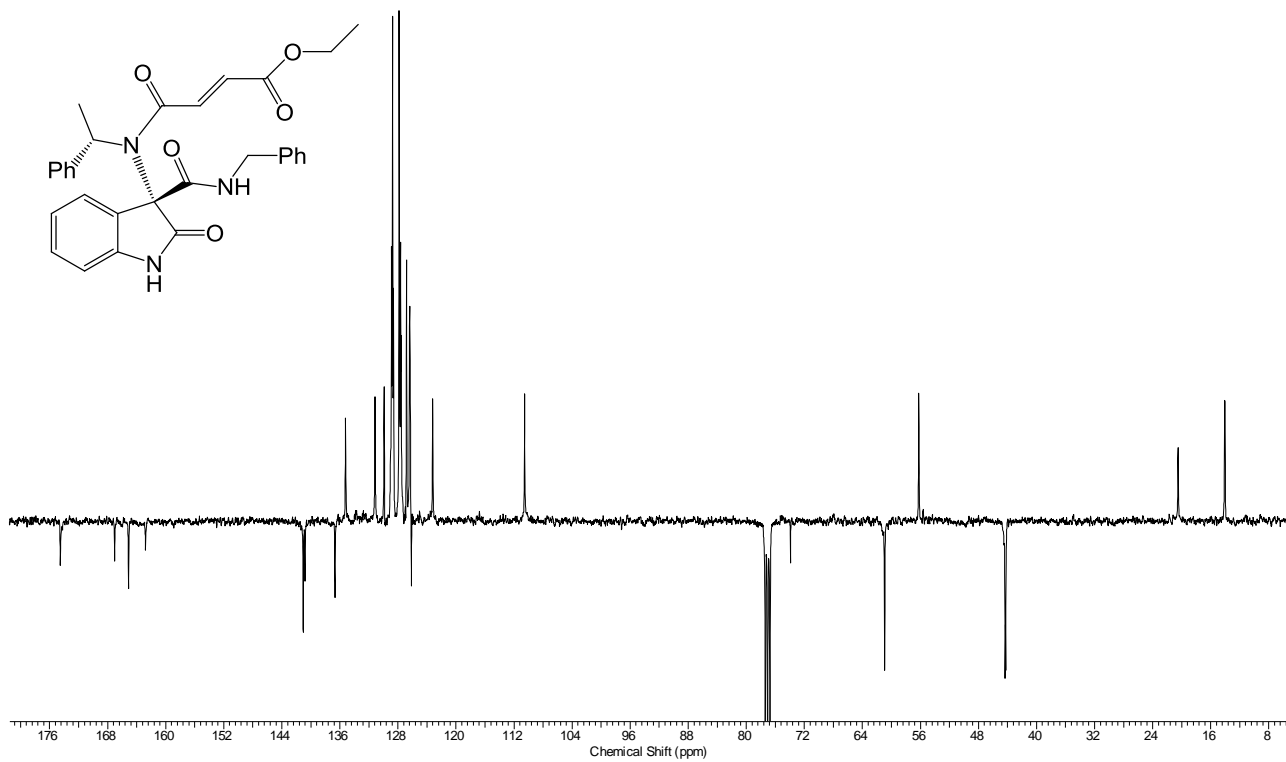
Compound **14a**:  $^{13}\text{C}$  NMR (100 MHz,  $\text{CDCl}_3$ )



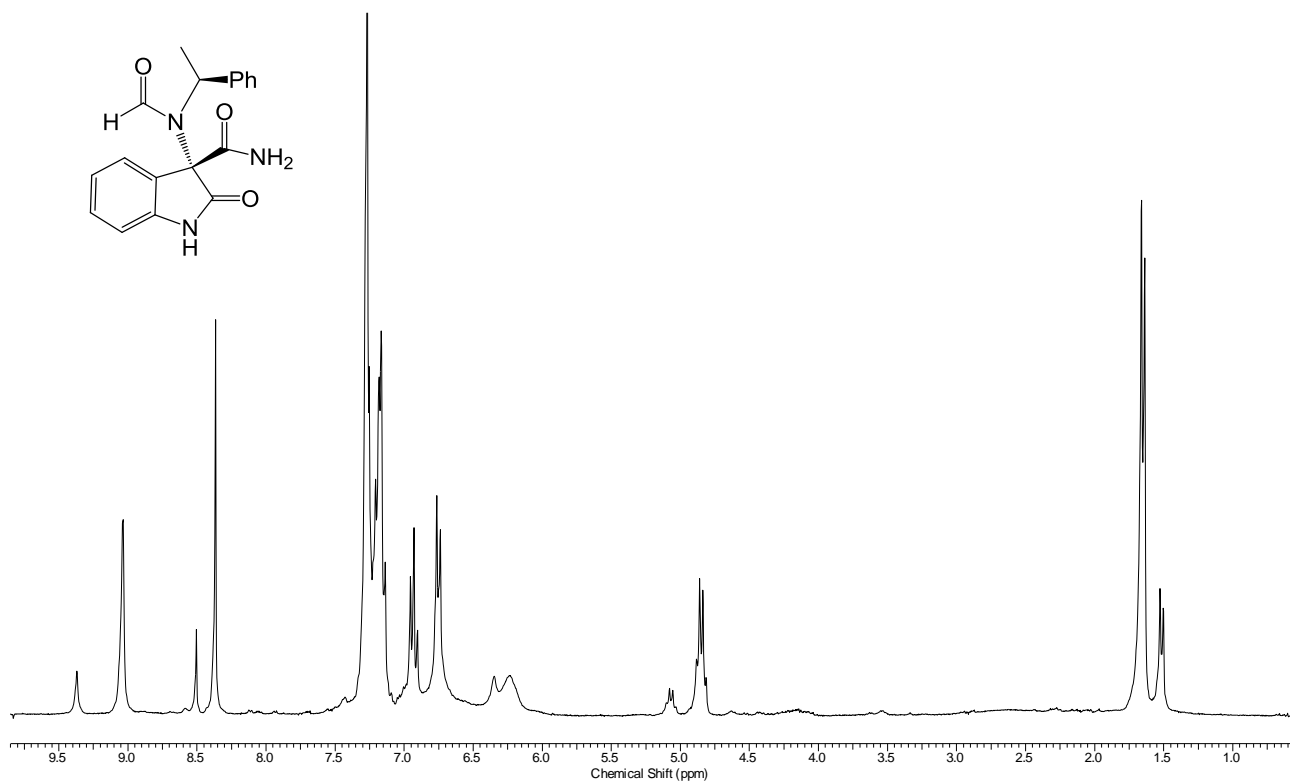
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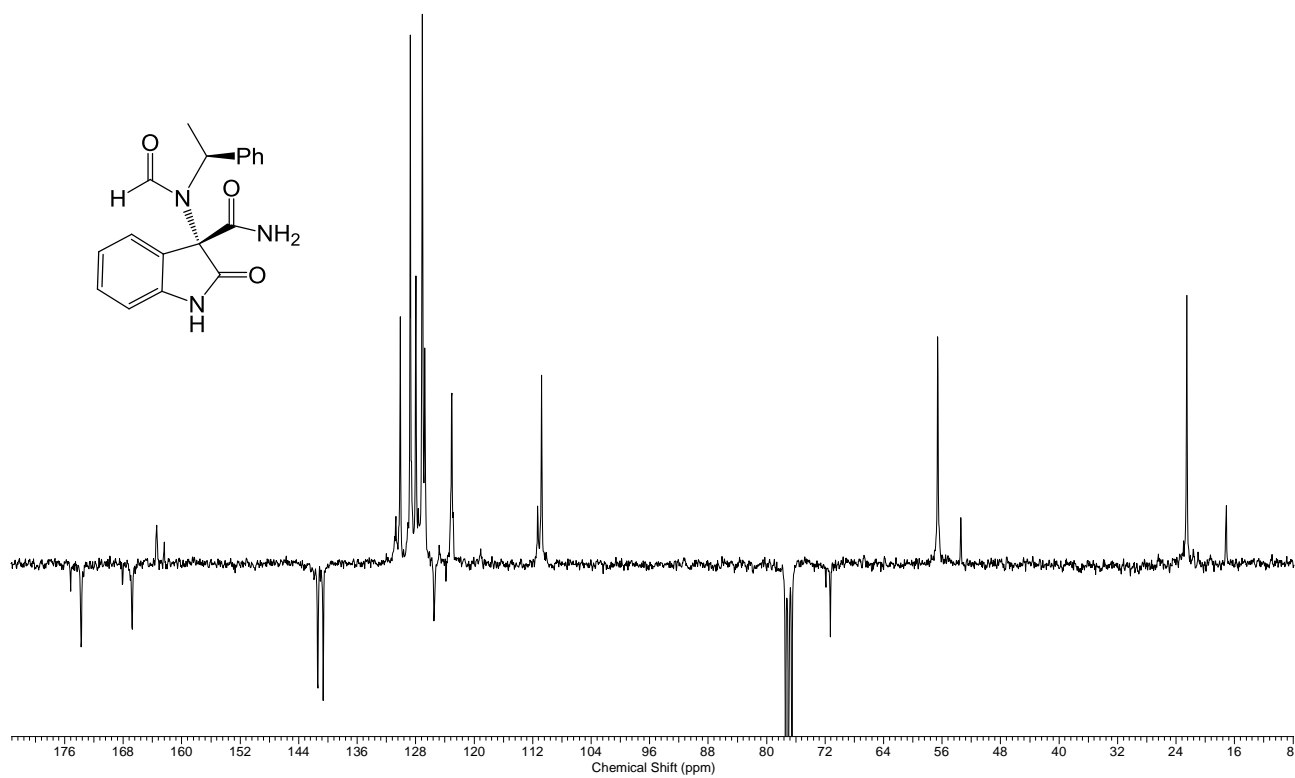
Compound **15a**:  $^{13}\text{C}$  NMR (100 MHz,  $\text{CDCl}_3$ )



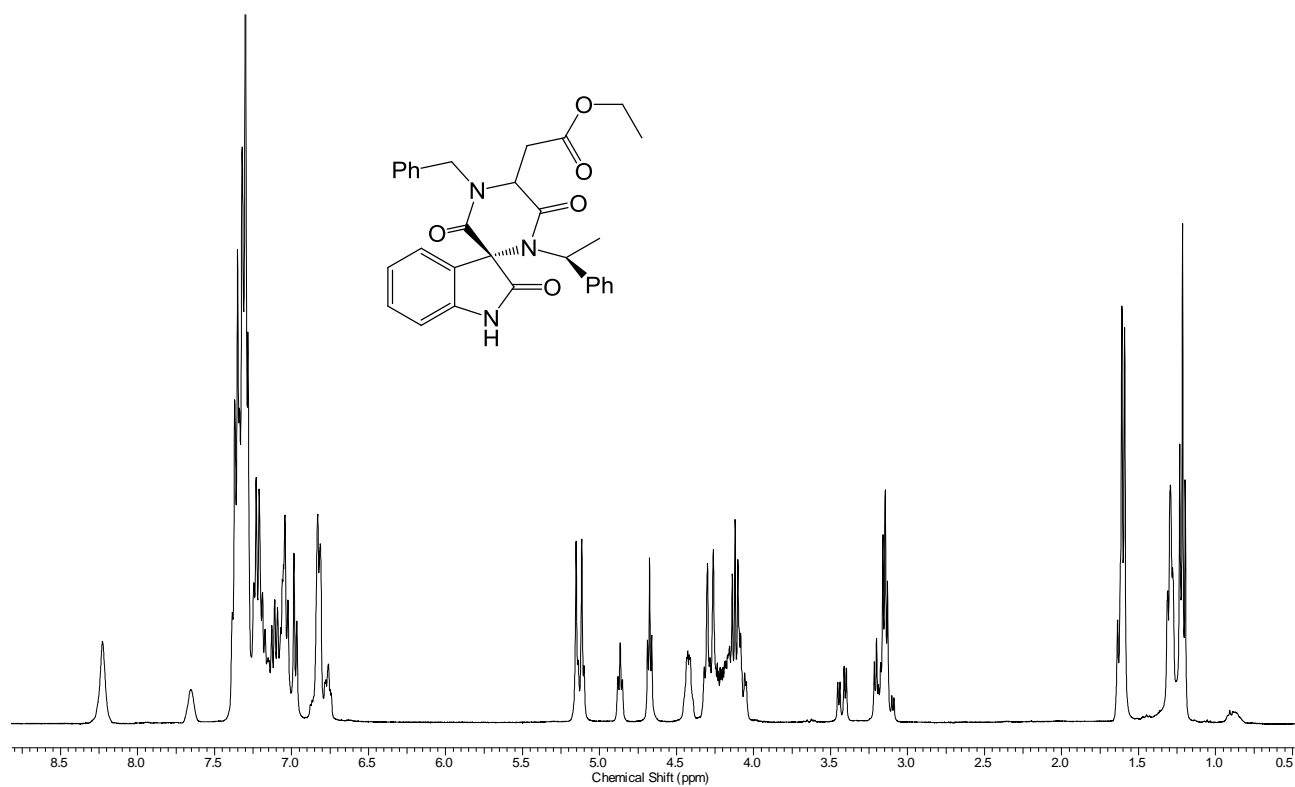
Compound **16**:  $^1\text{H}$  NMR (300 MHz,  $\text{CDCl}_3$ , 0.15:0.85 rotameric mixture)



Compound **16**:  $^{13}\text{C}$  NMR (75 MHz,  $\text{CDCl}_3$ )



Compound **18**:  $^1\text{H}$  NMR (400 MHz,  $\text{CDCl}_3$ )



Compound **18**:  $^{13}\text{C}$  NMR (100 MHz,  $\text{CDCl}_3$ )

