

Supporting Information

Asymmetric Total Synthesis of Fawcettimine-Type *Lycopodium* Alkaloid, Lycopoclavamine-A

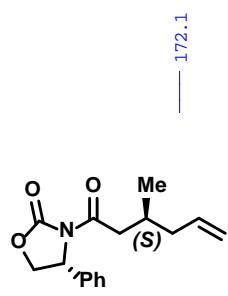
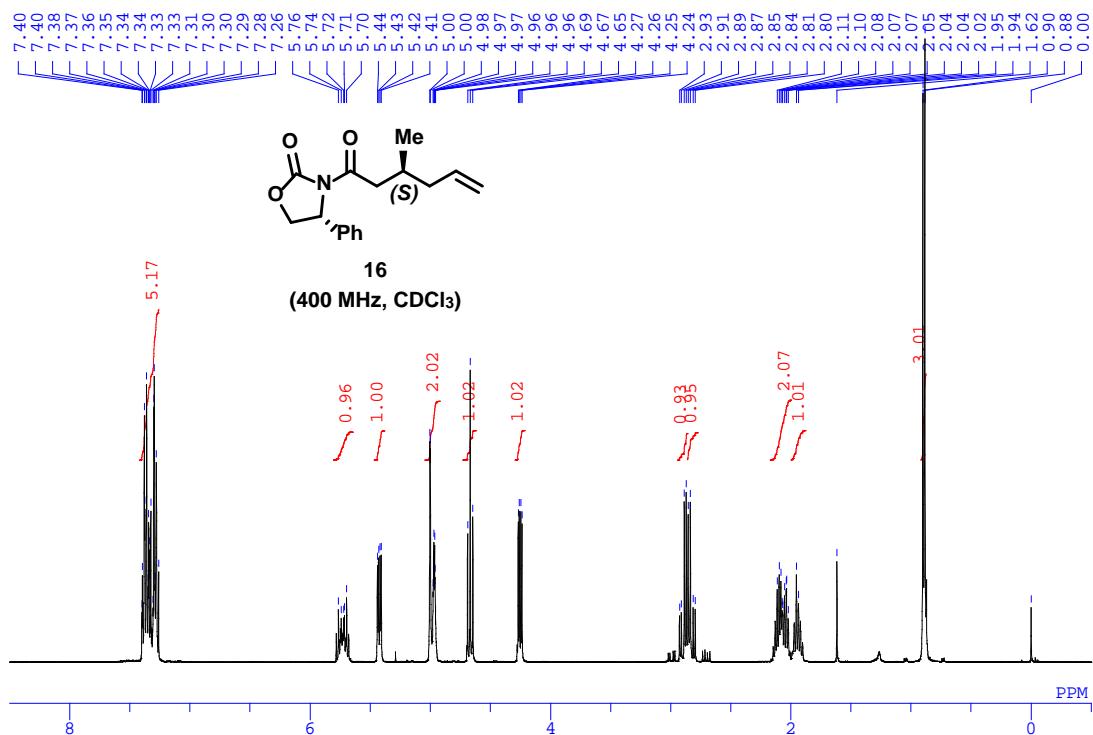
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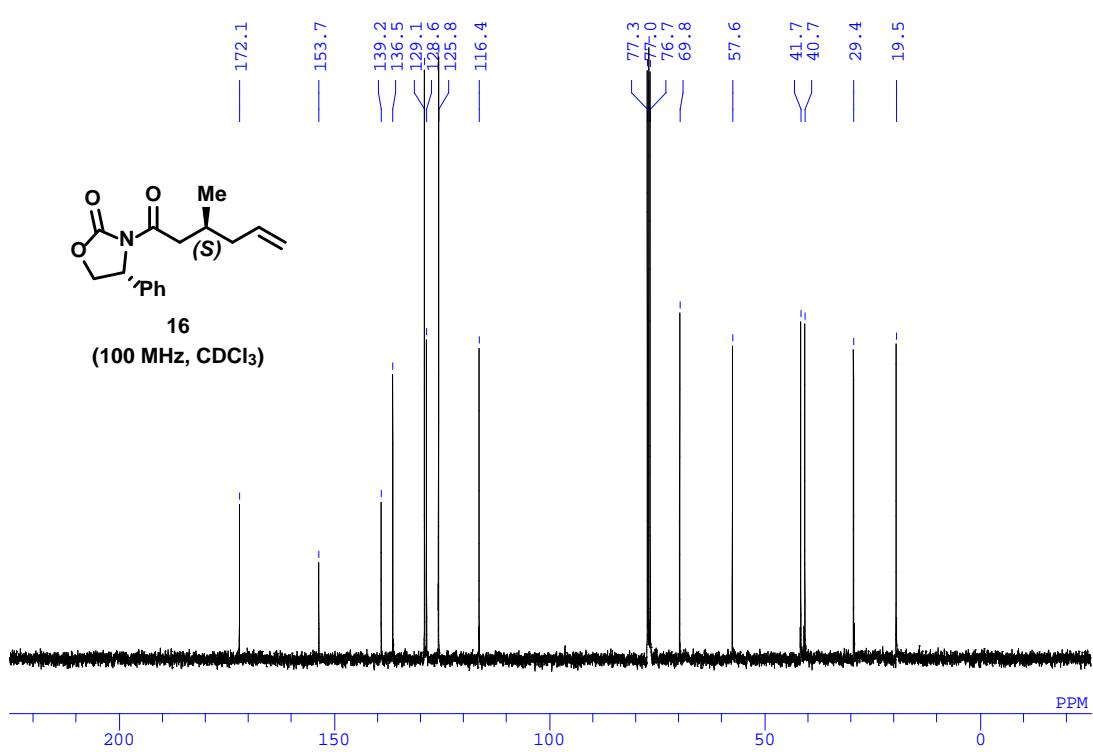
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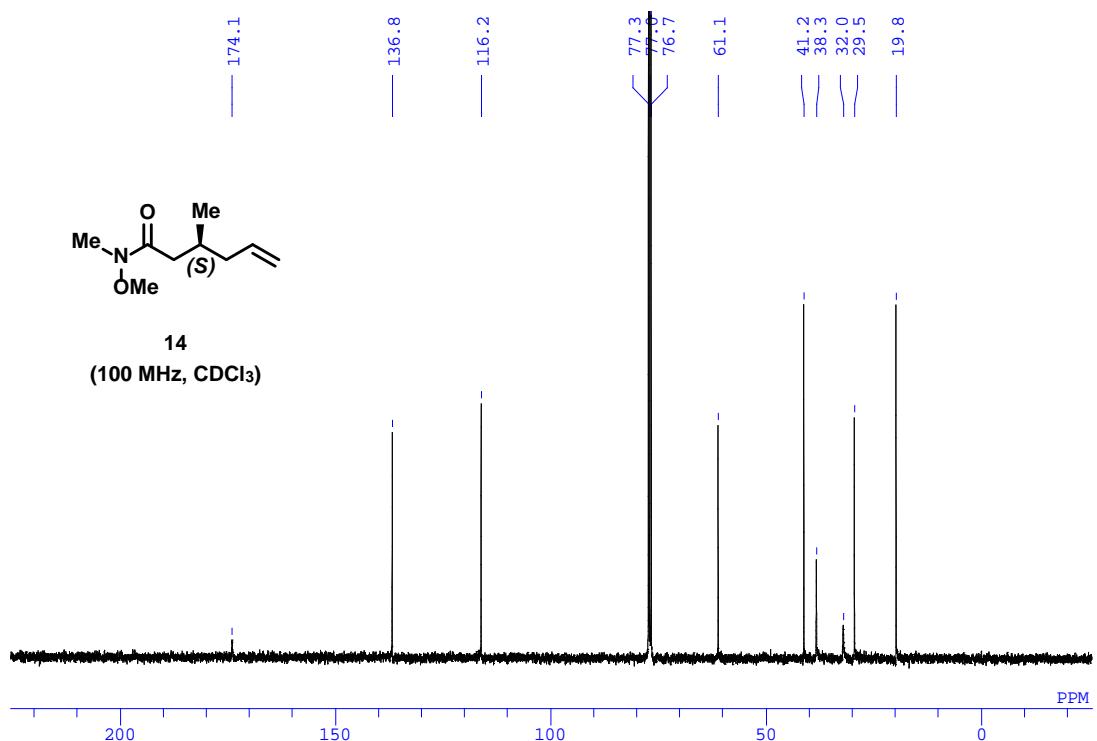
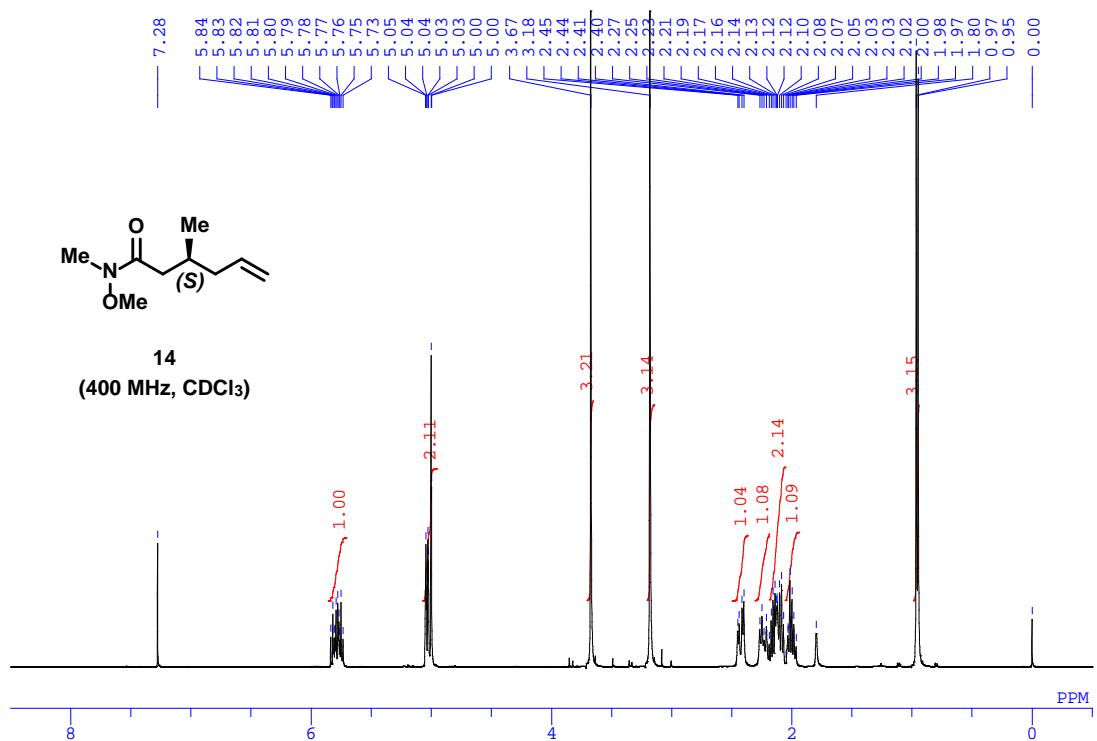
1. ^1H and ^{13}C { ^1H } NMR spectra	S1
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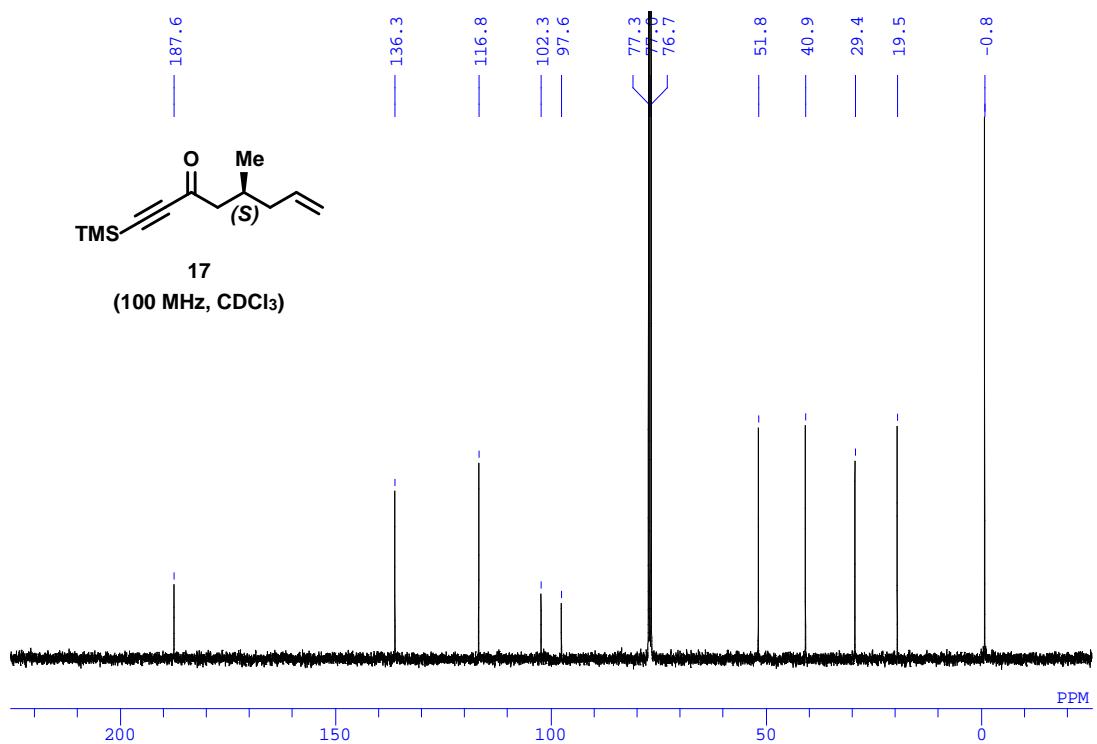
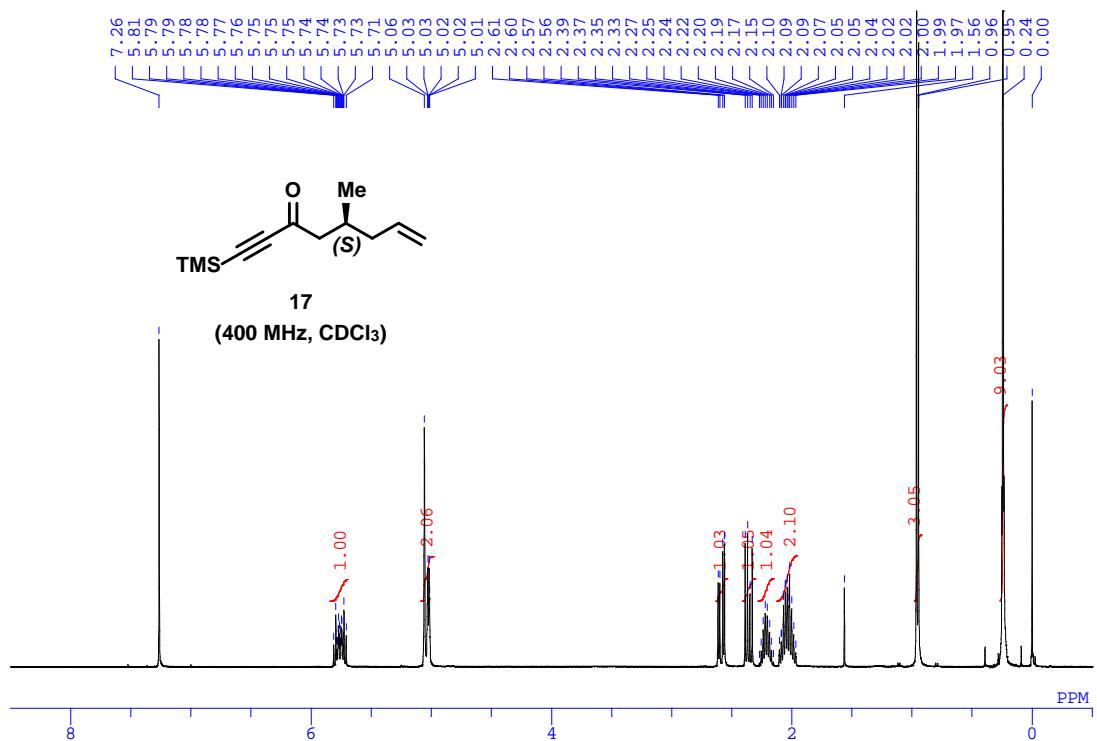
¹H and ¹³C {¹H} NMR spectra

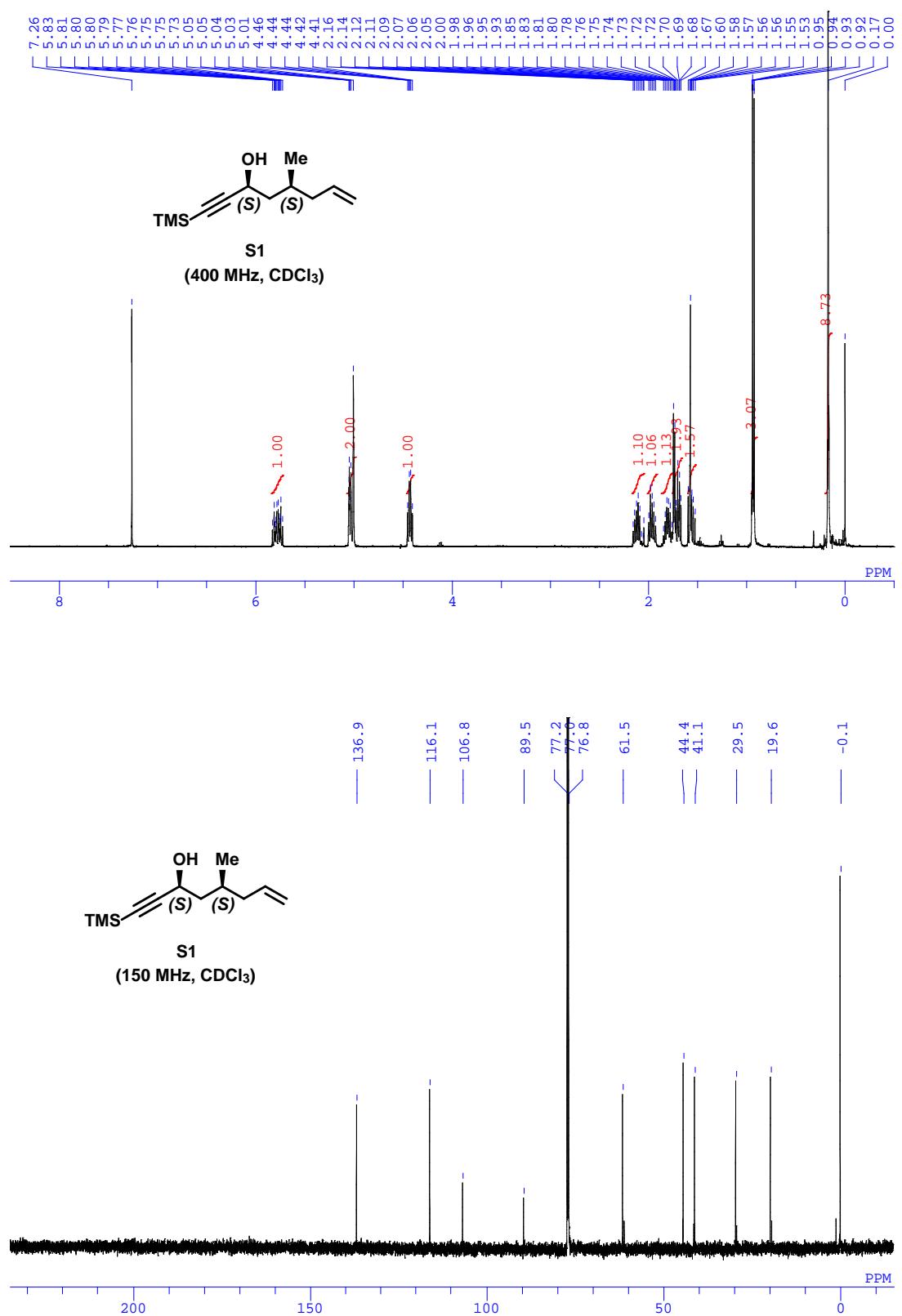


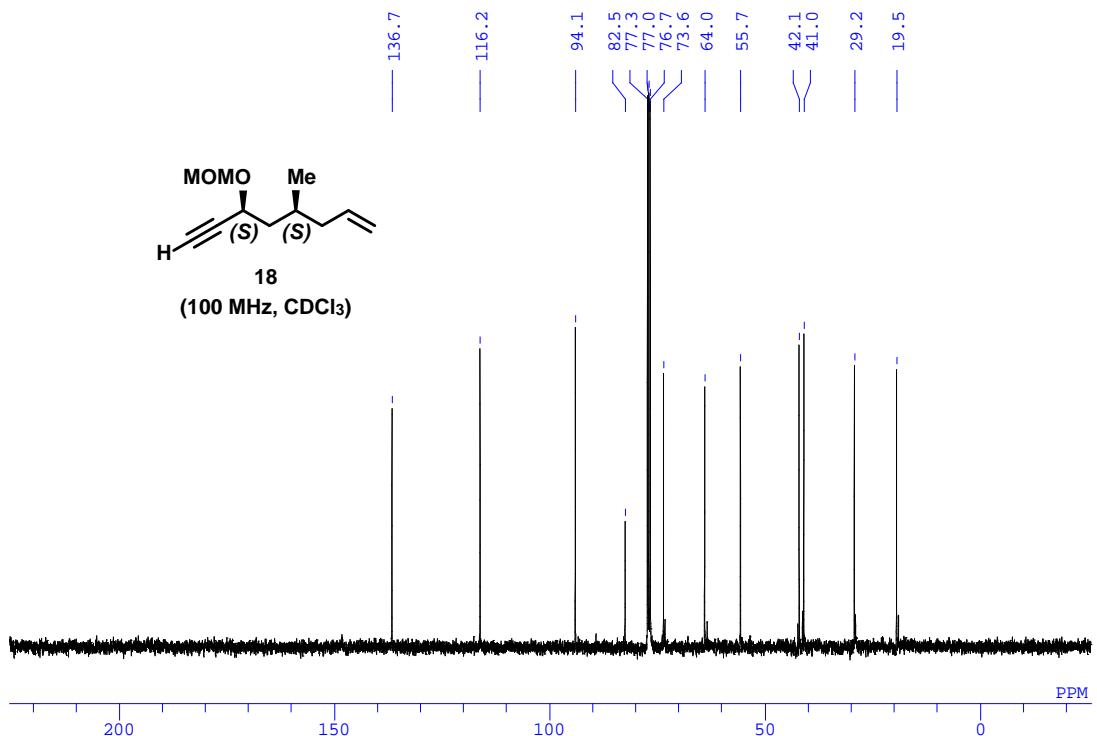
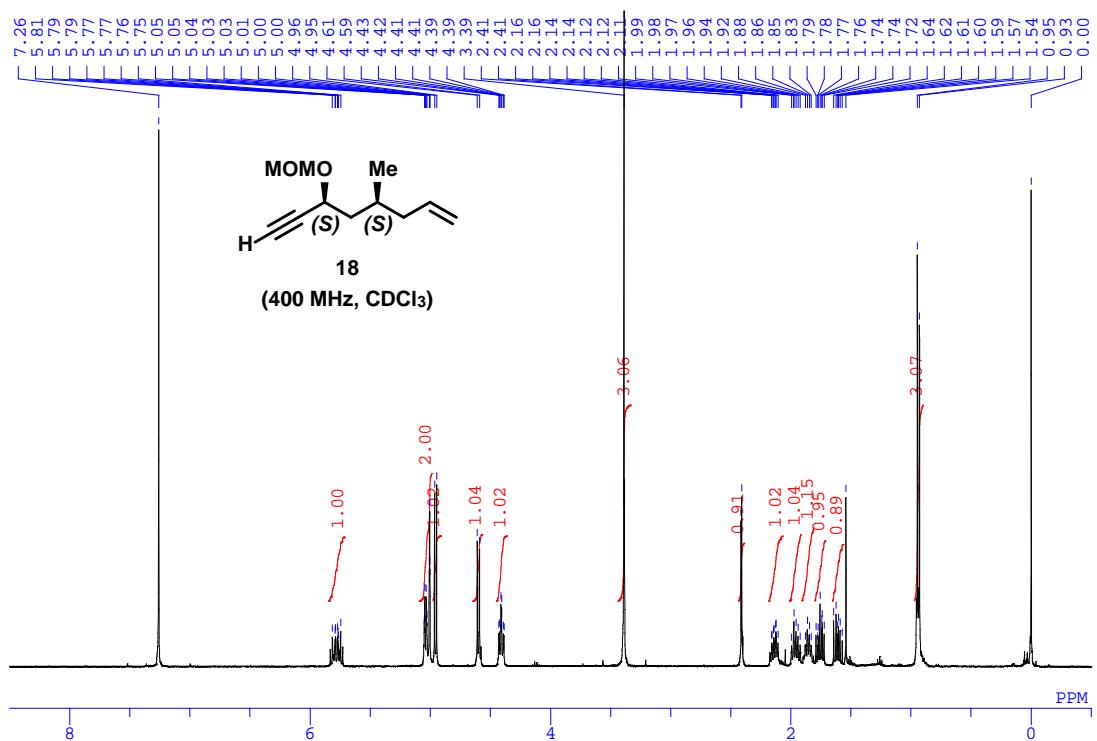
16
(100 MHz, CDCl₃)

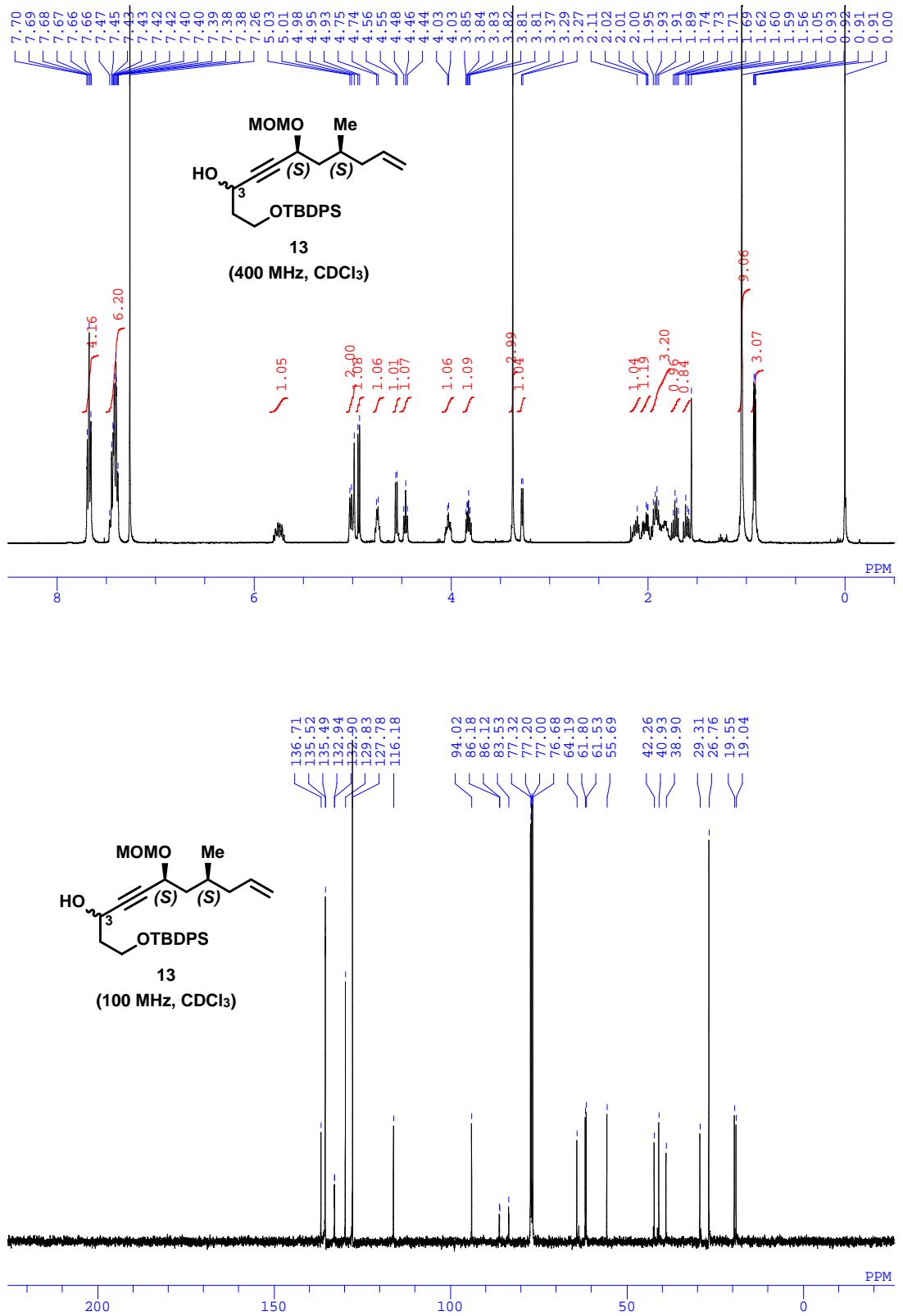


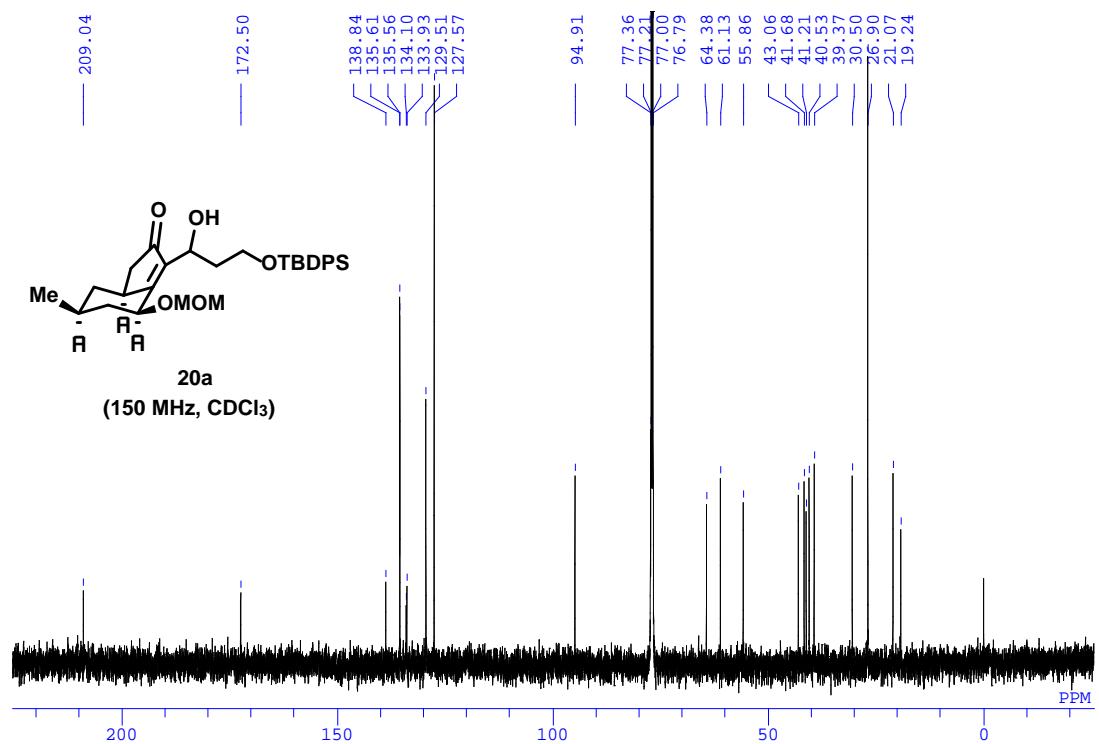
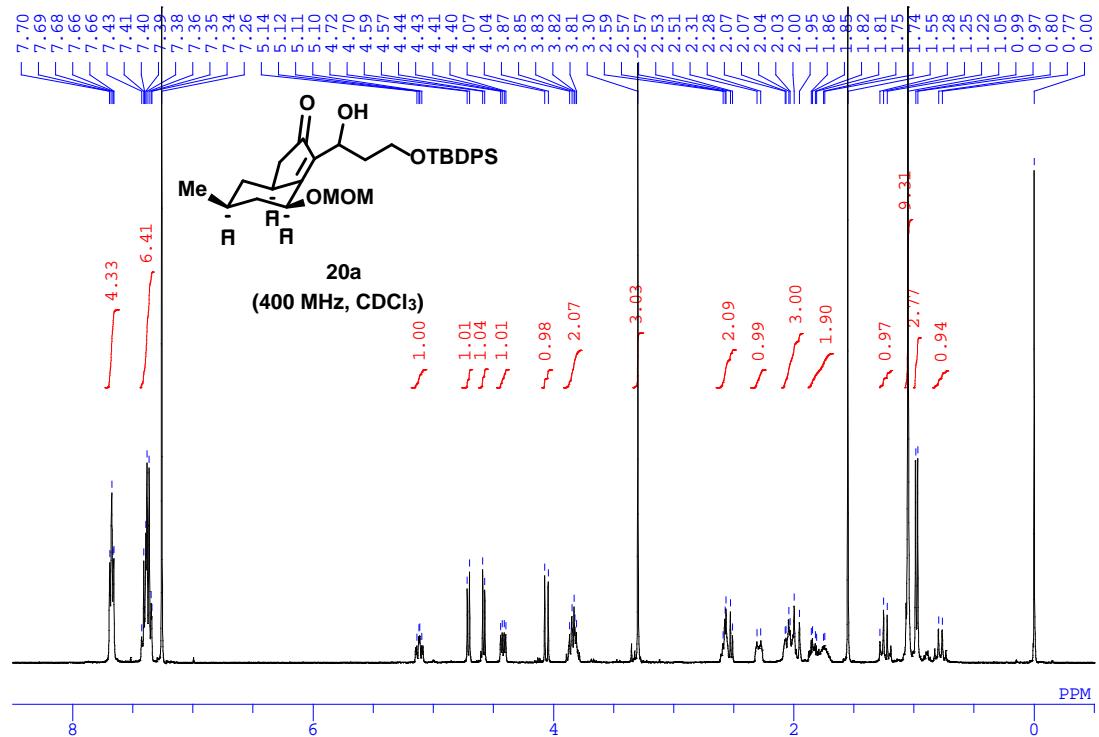


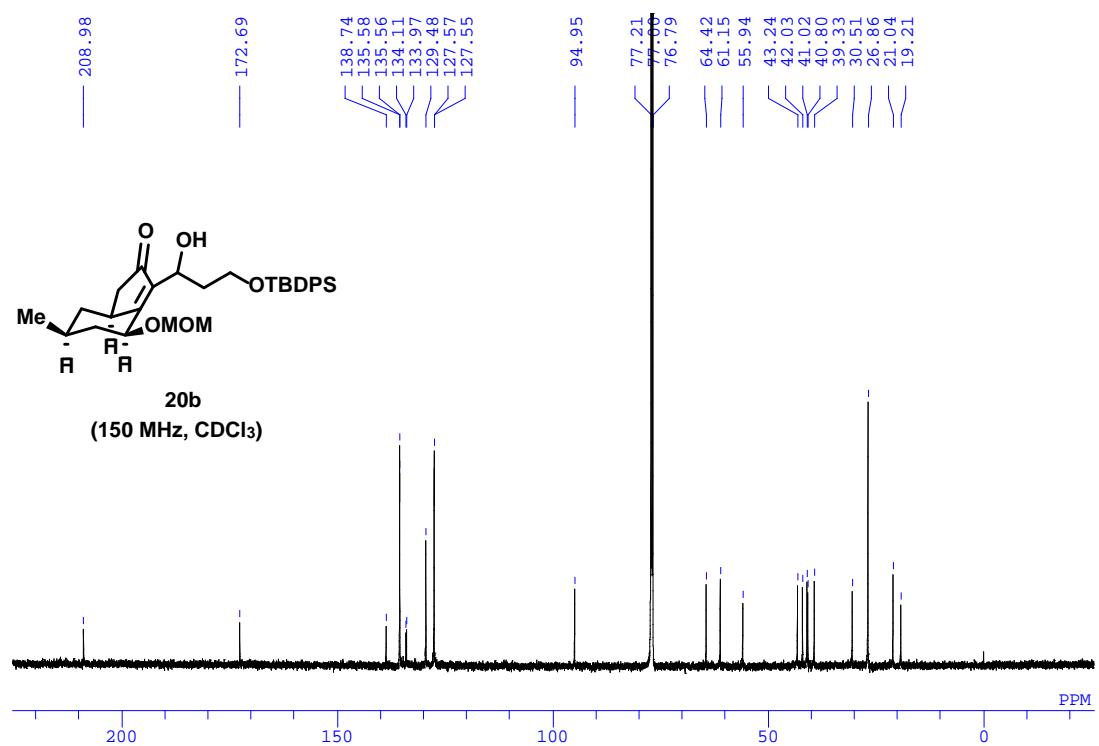
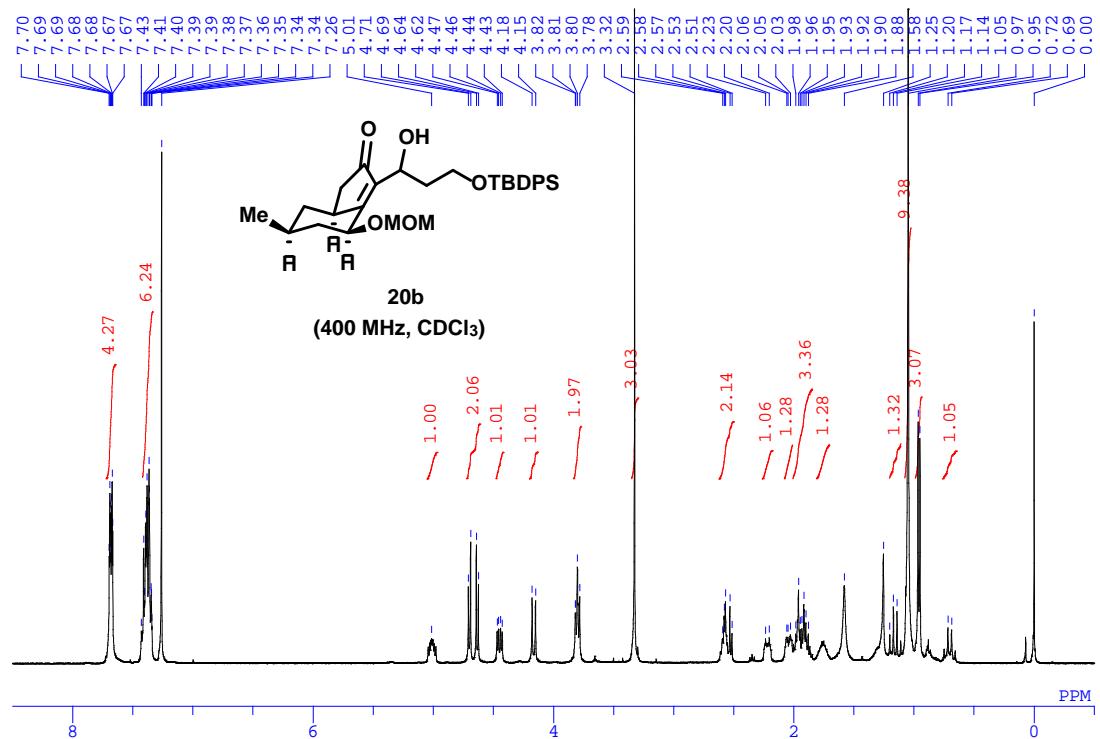


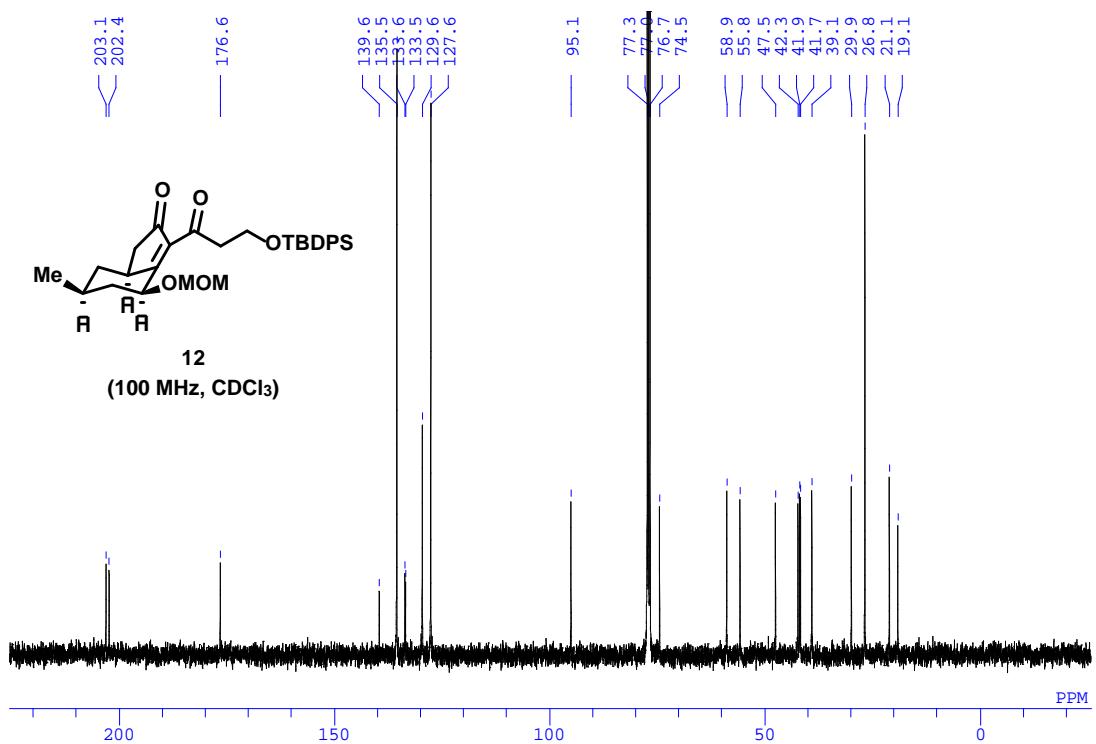
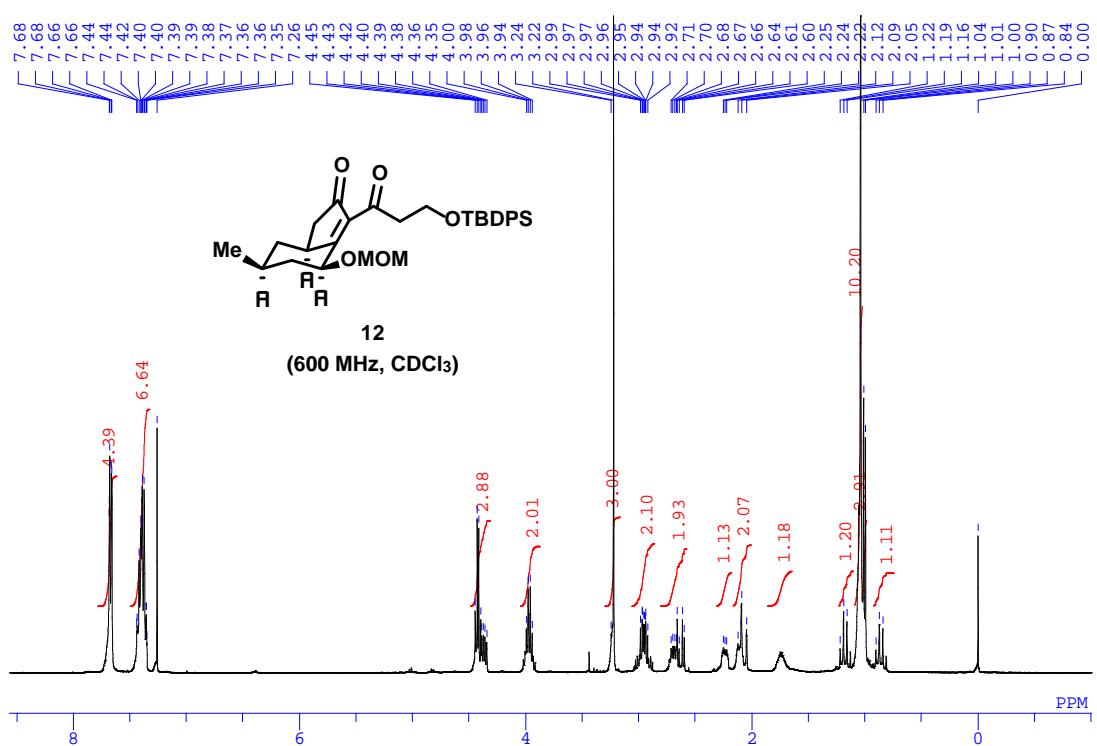


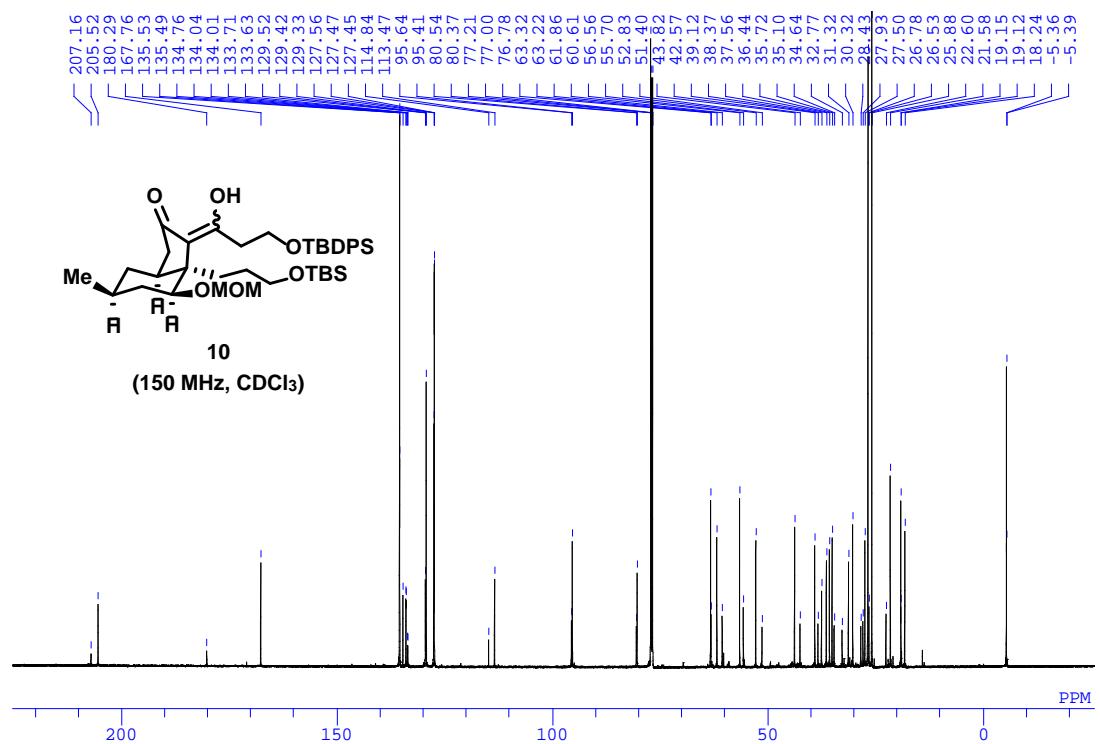
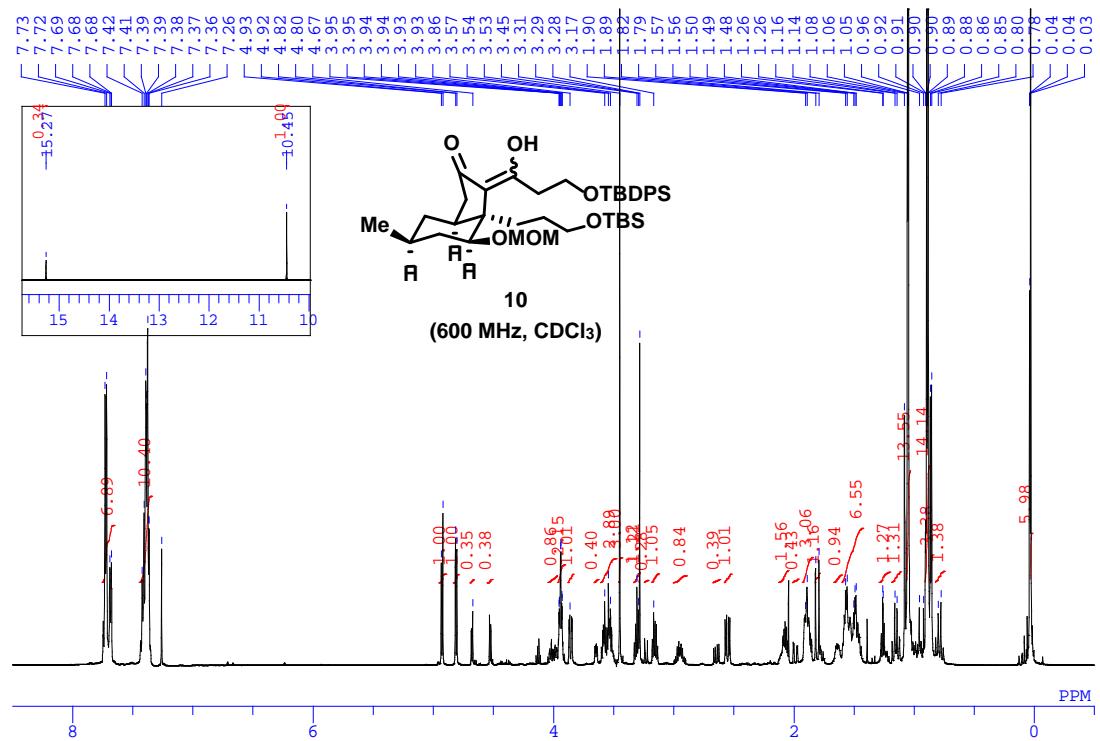


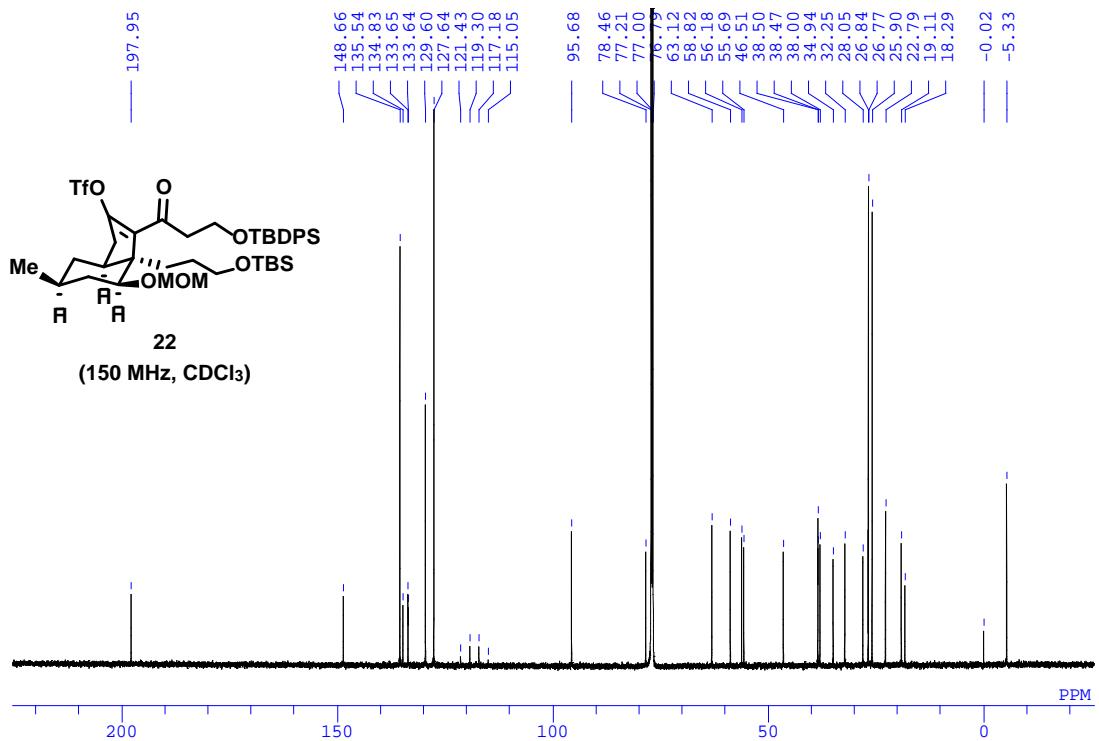
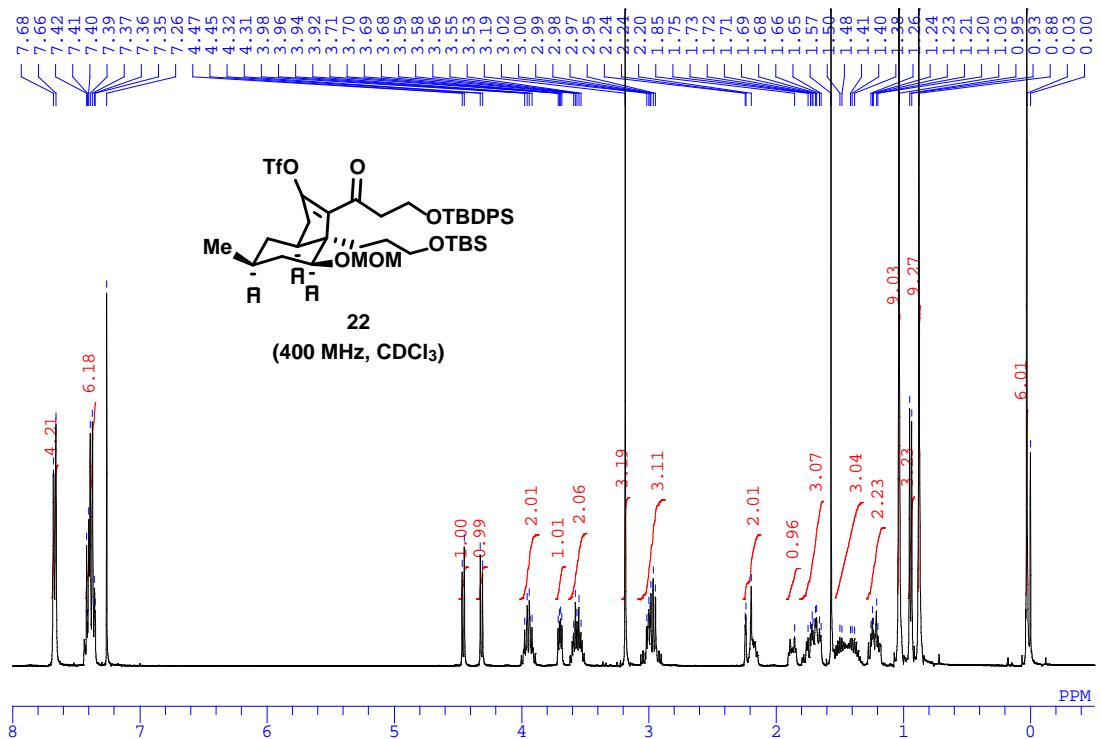


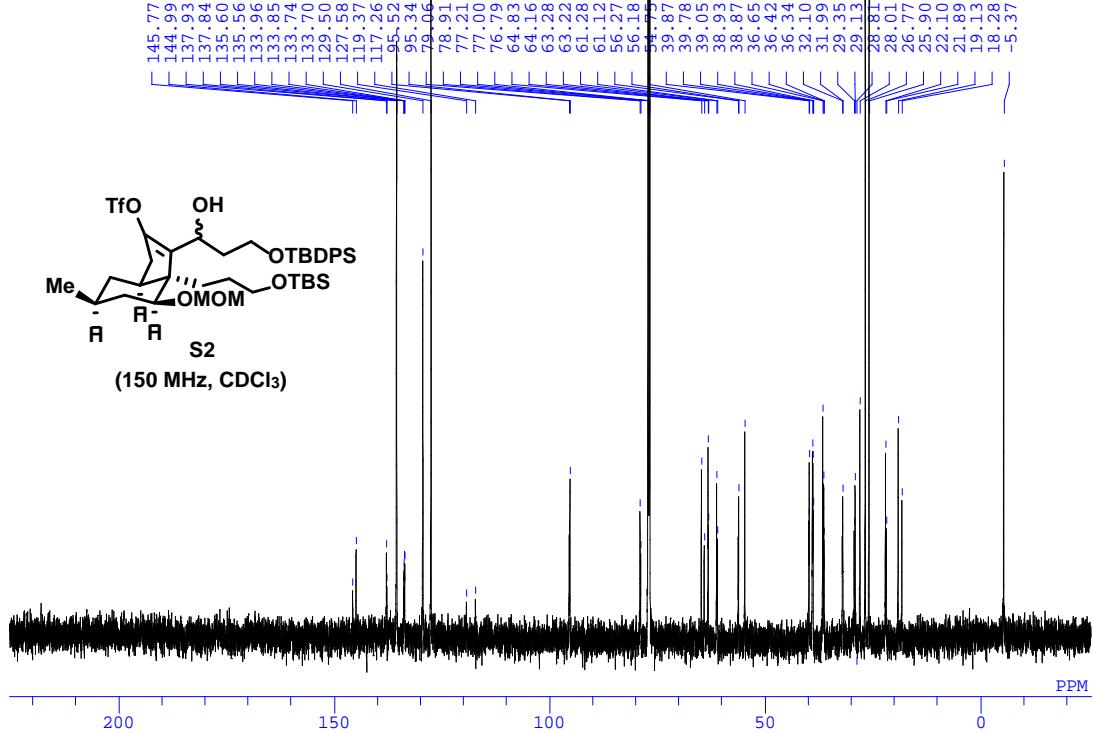
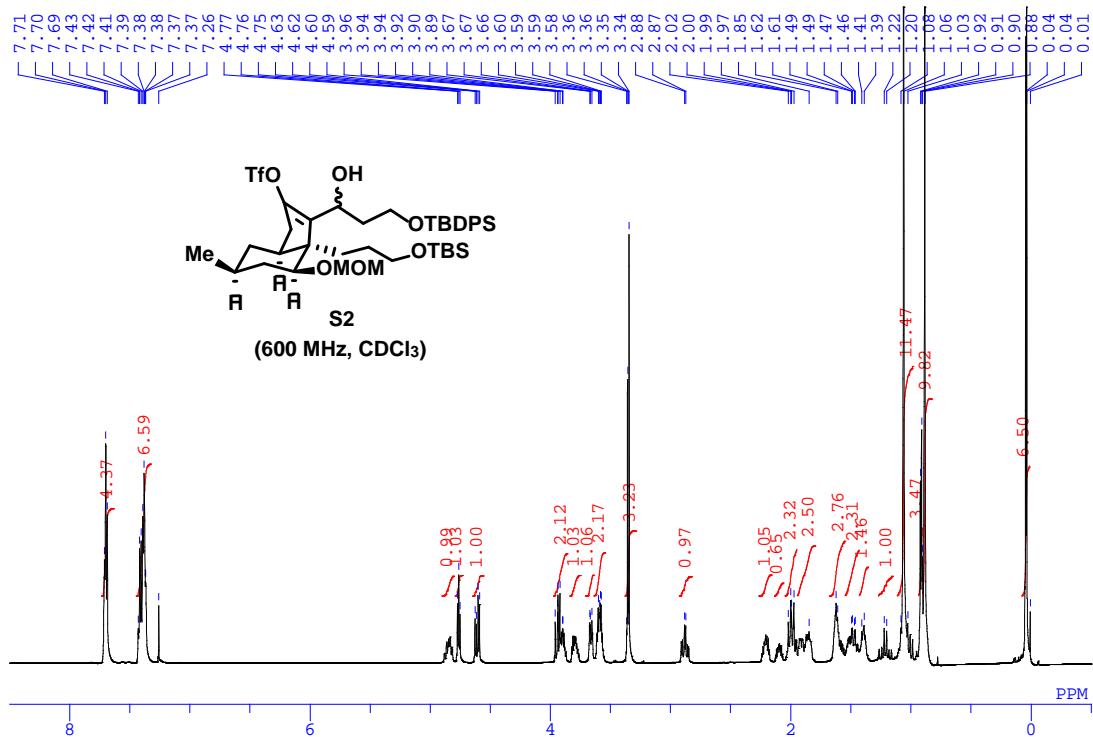


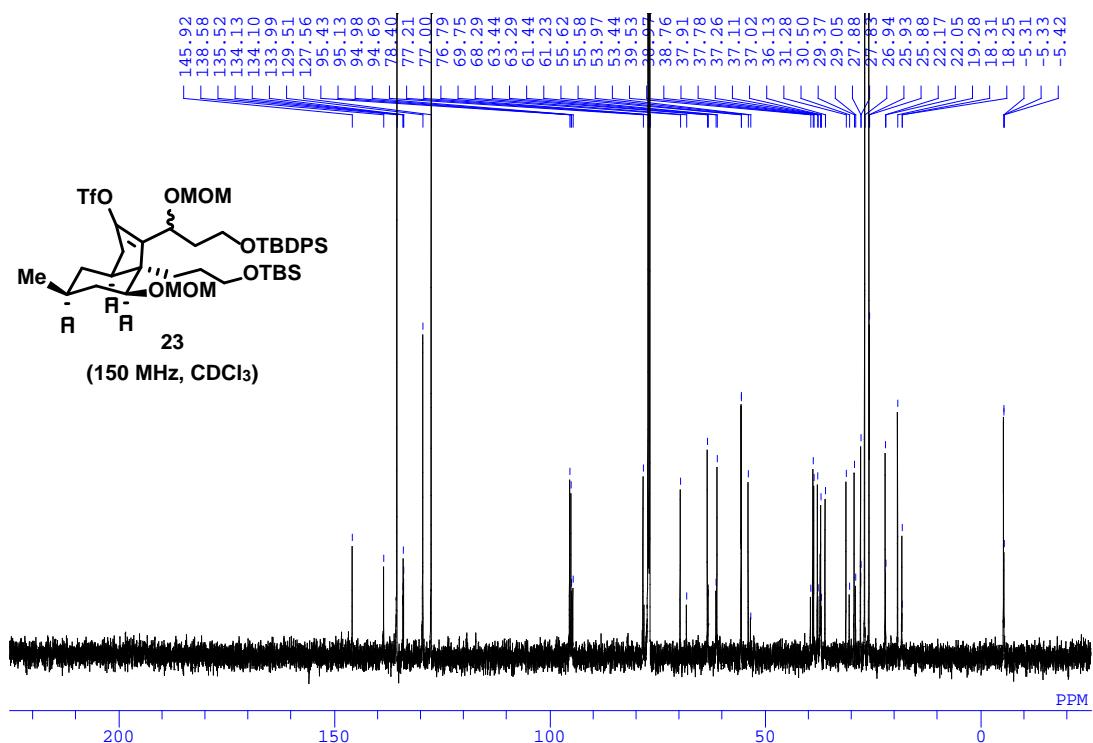
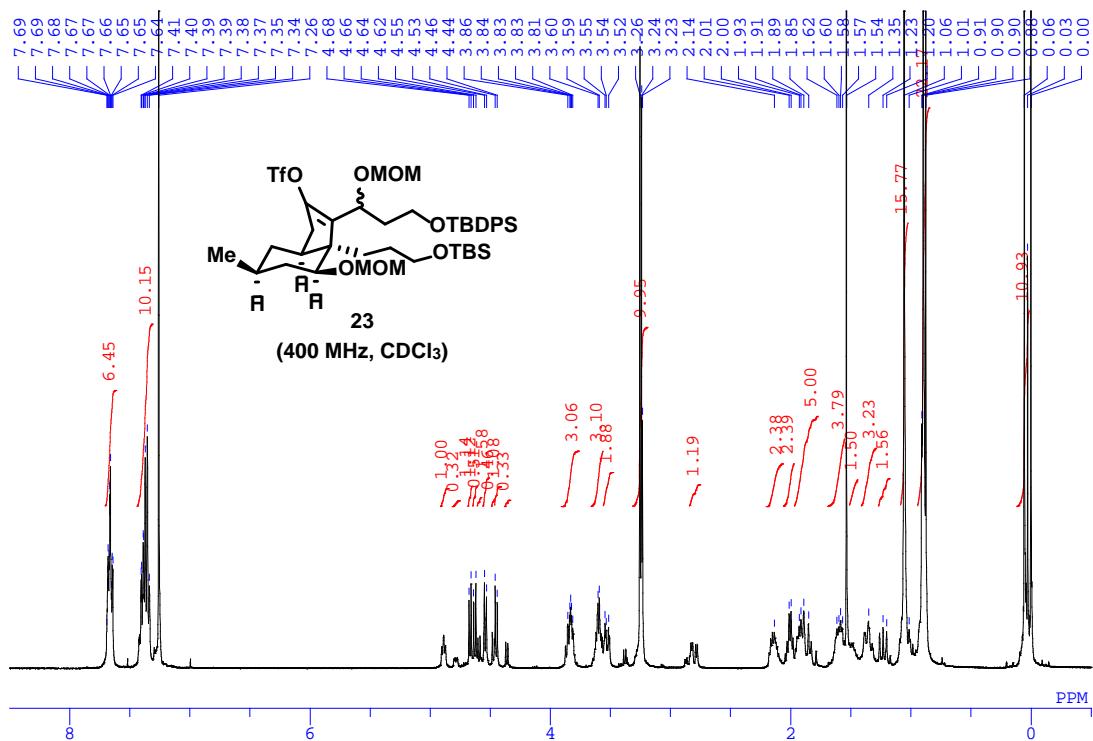


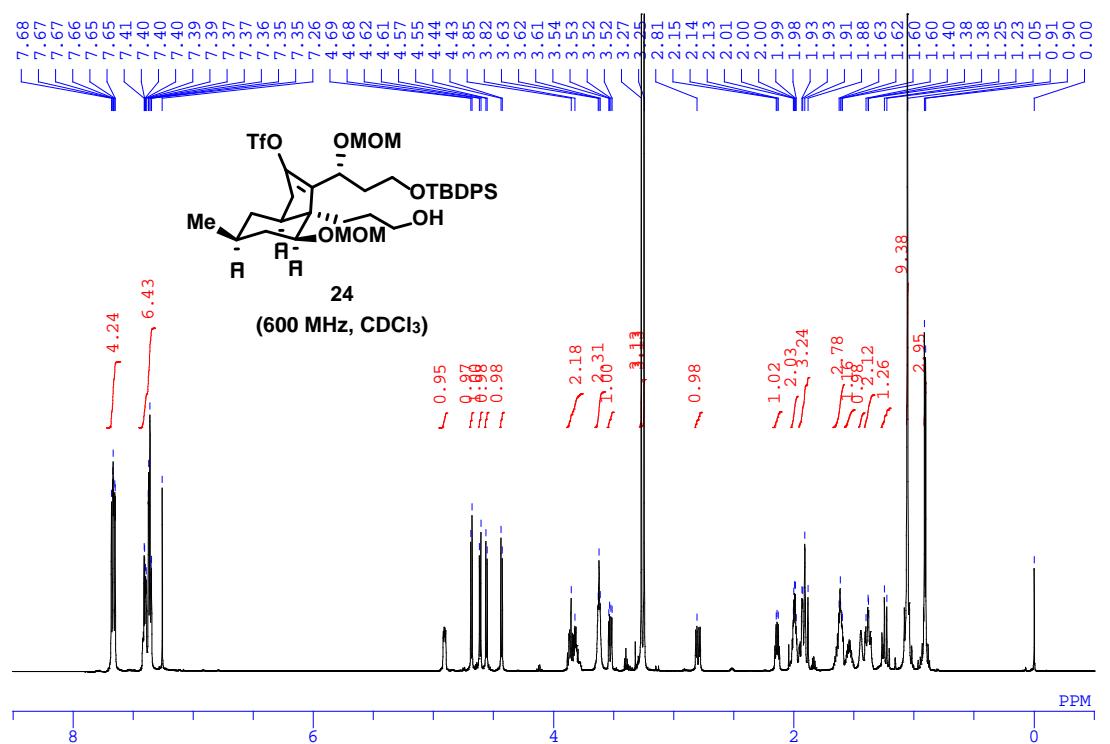


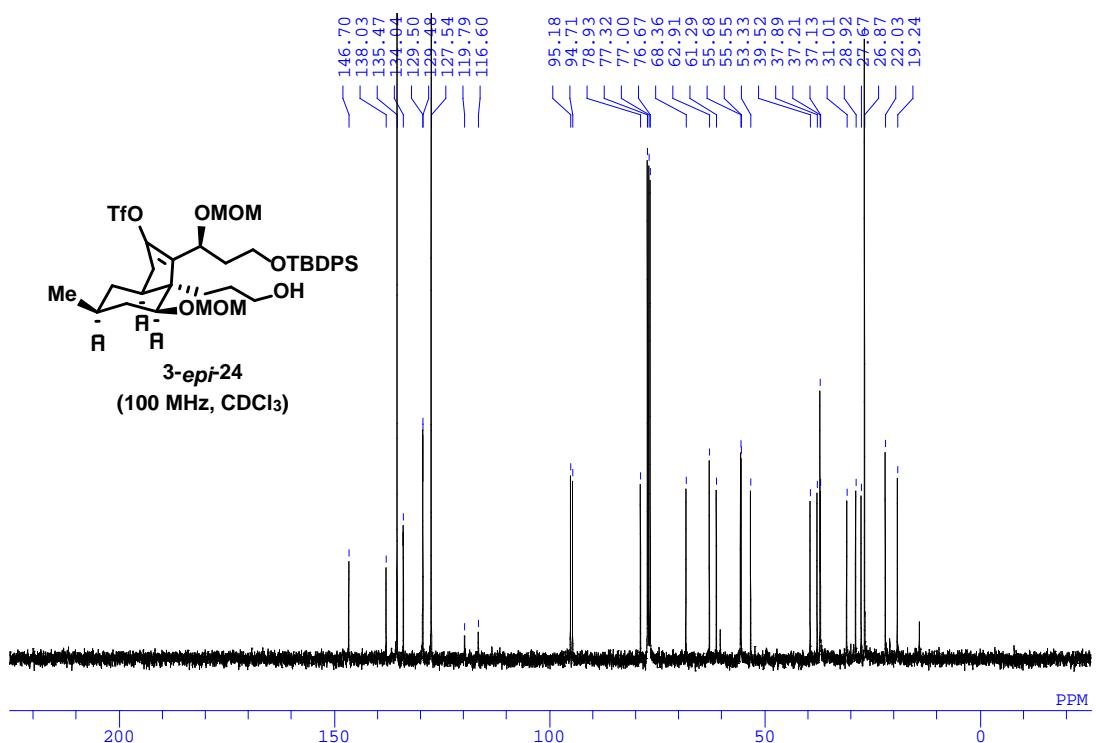
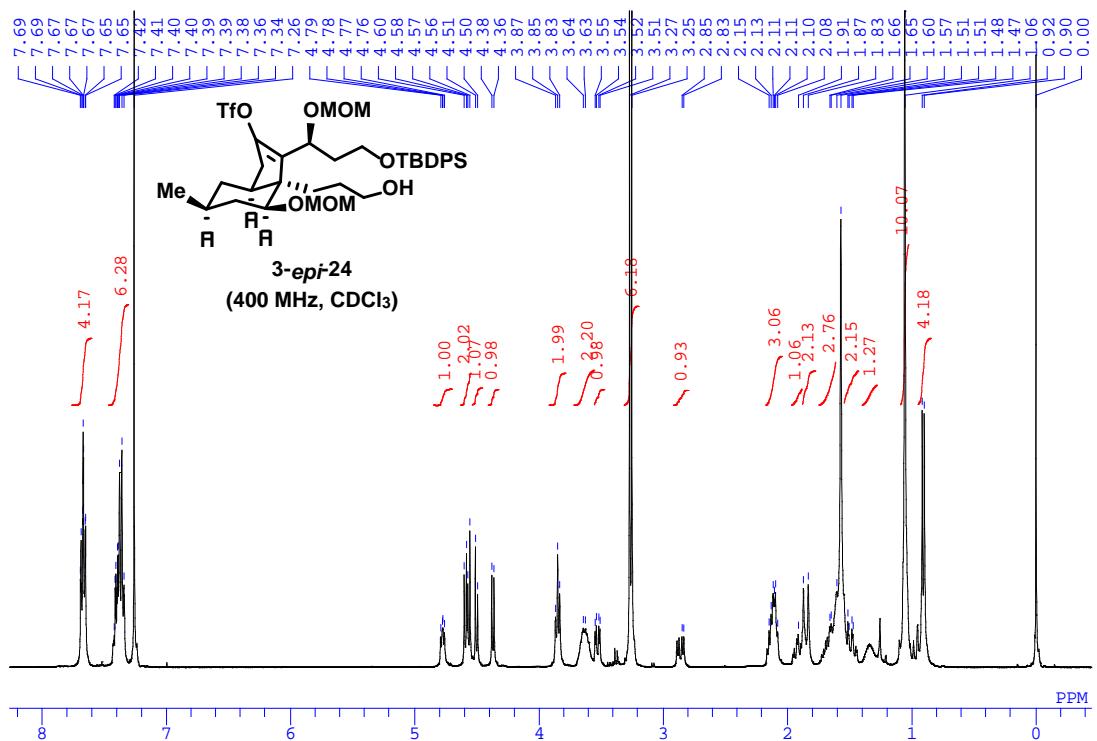


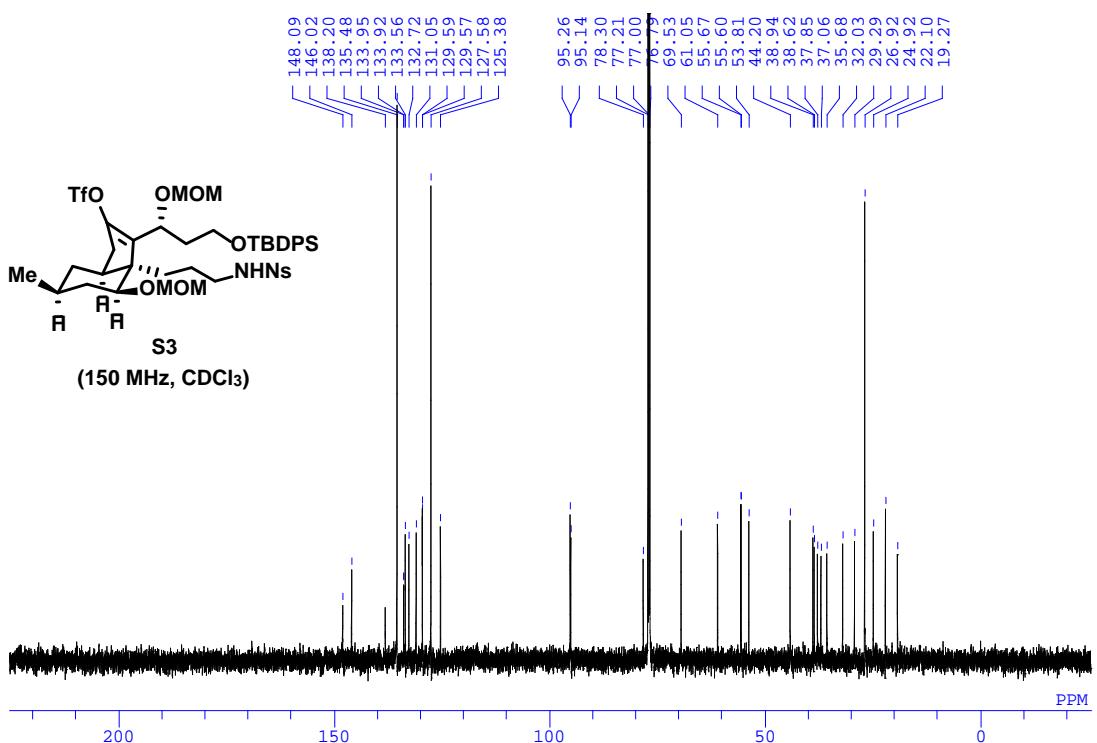
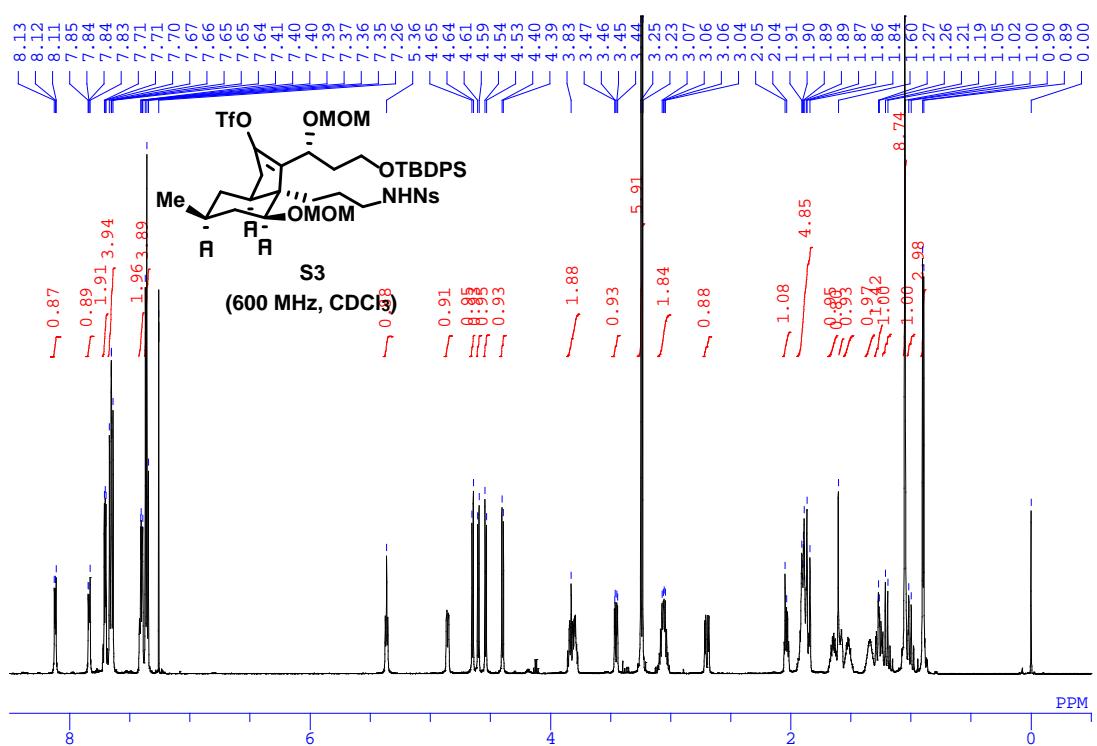


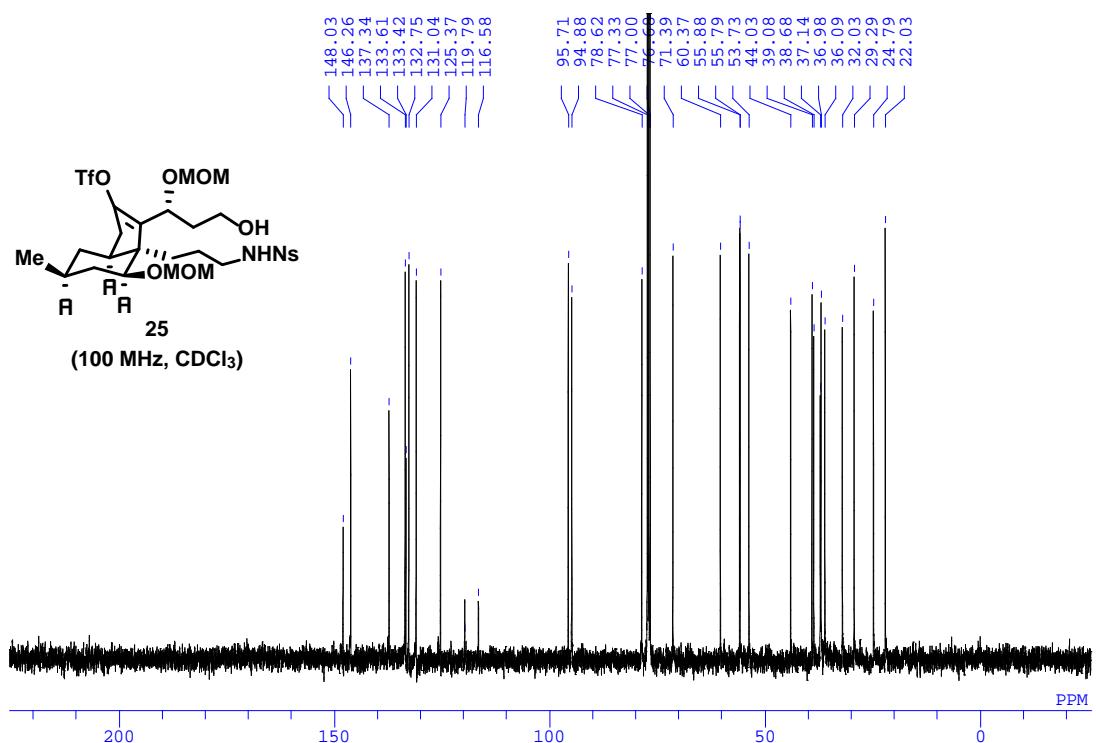
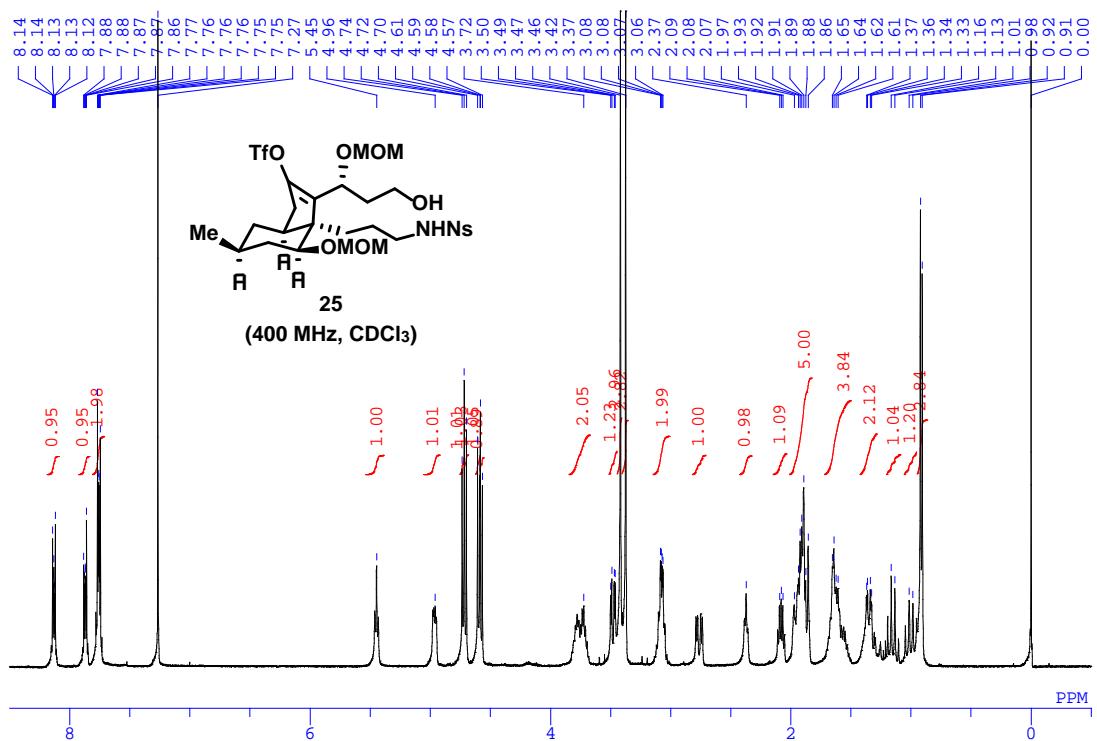


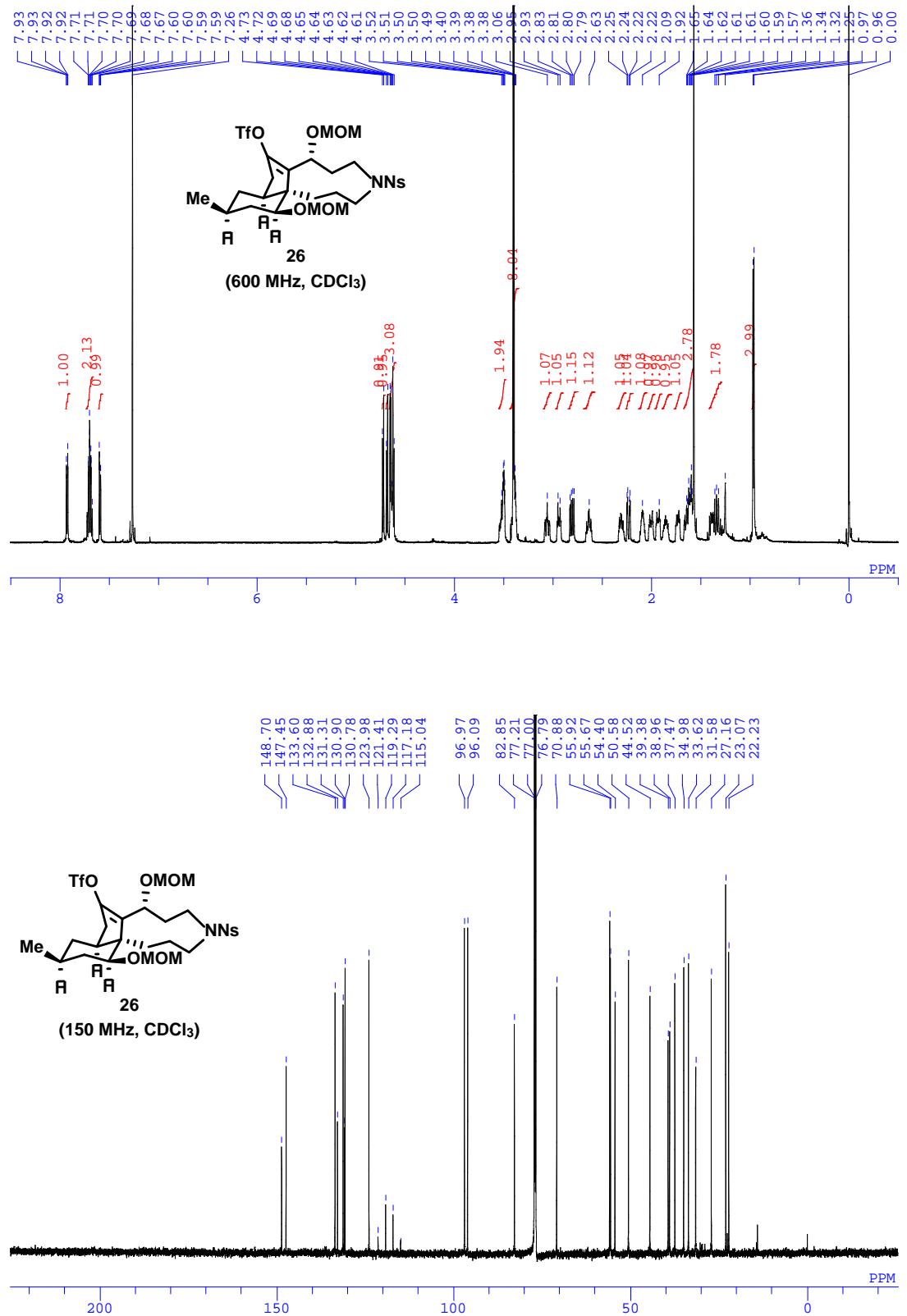


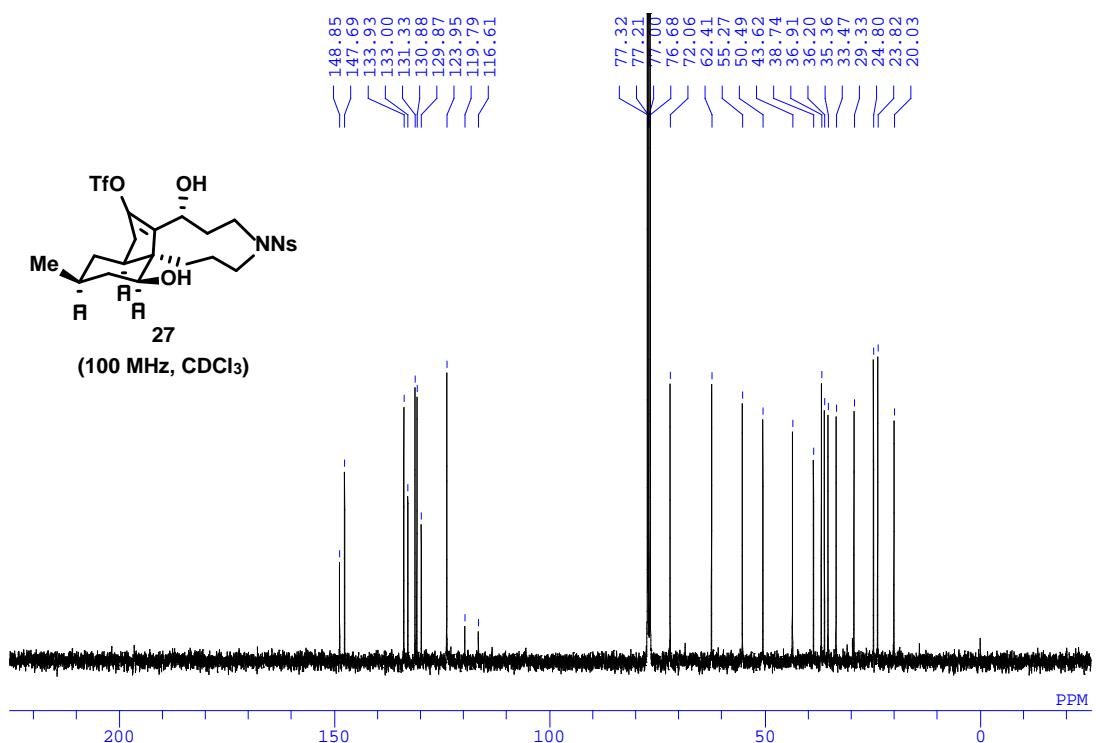
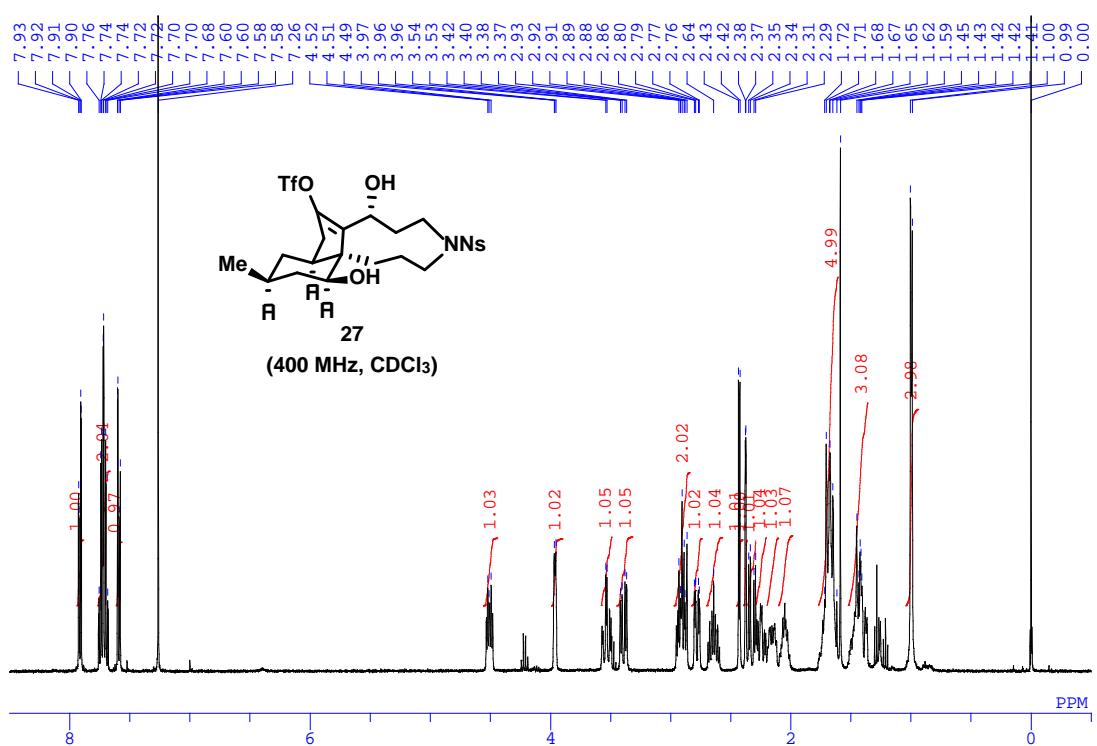


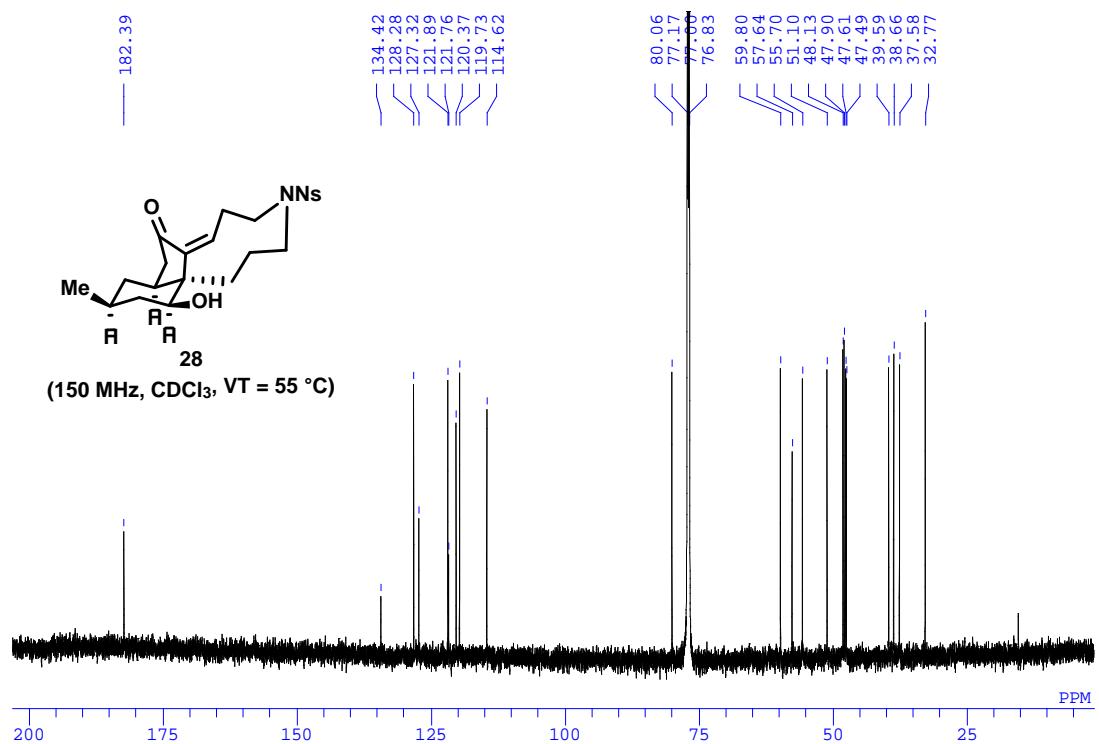
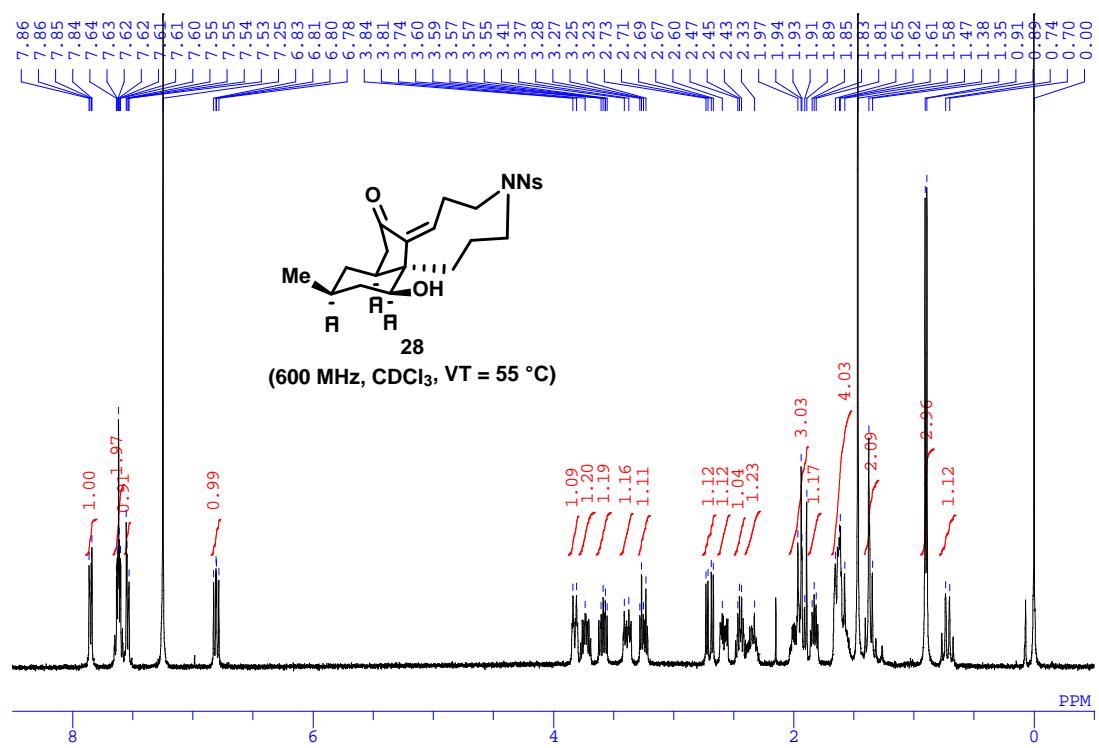


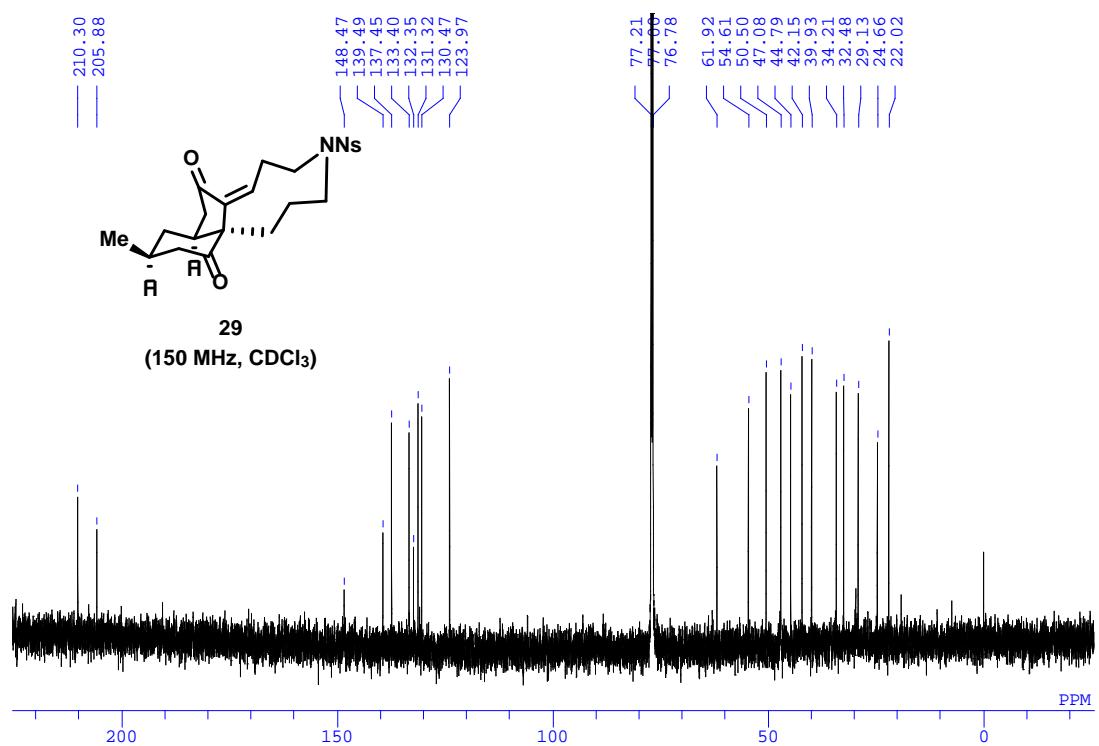
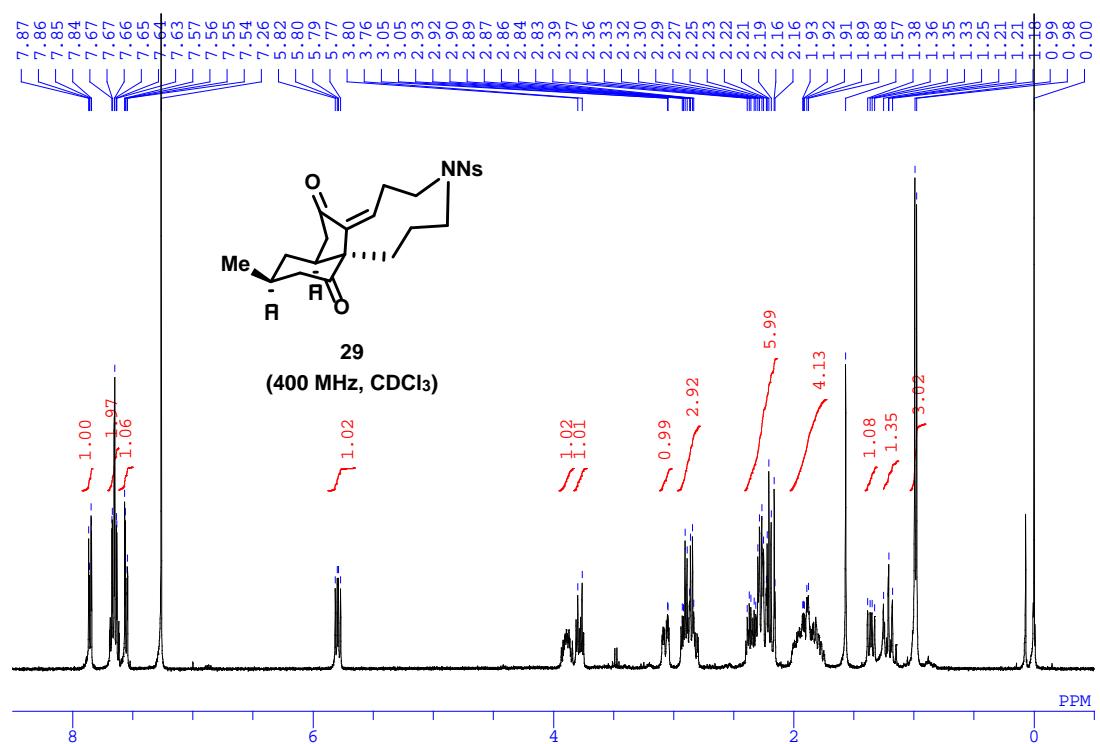


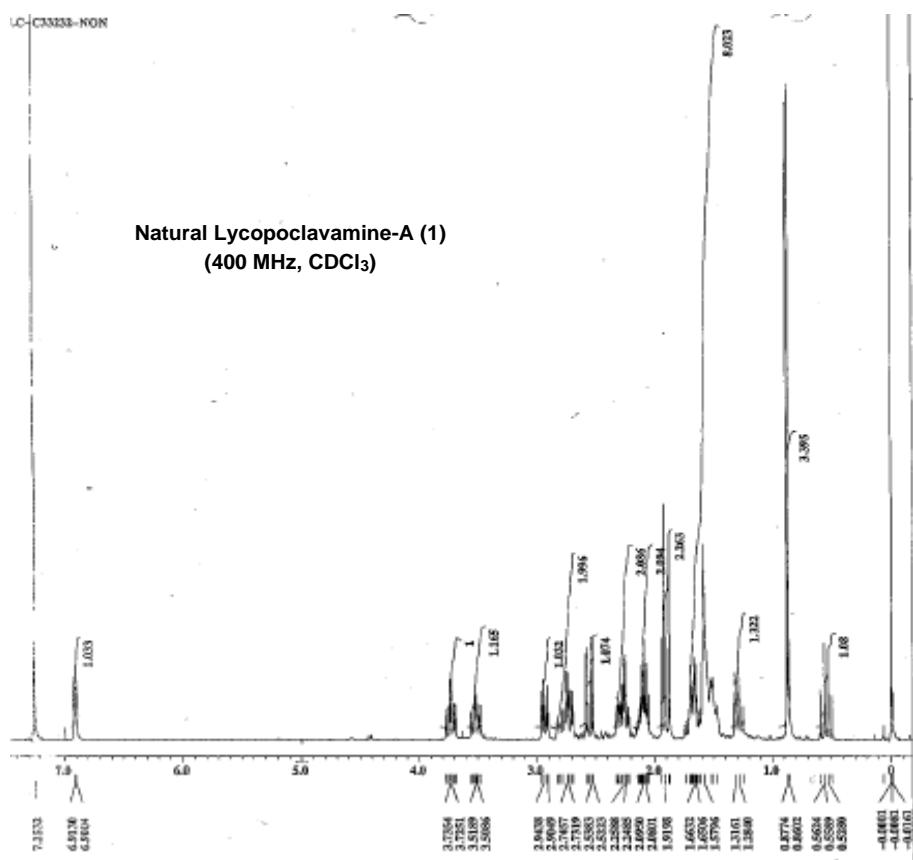
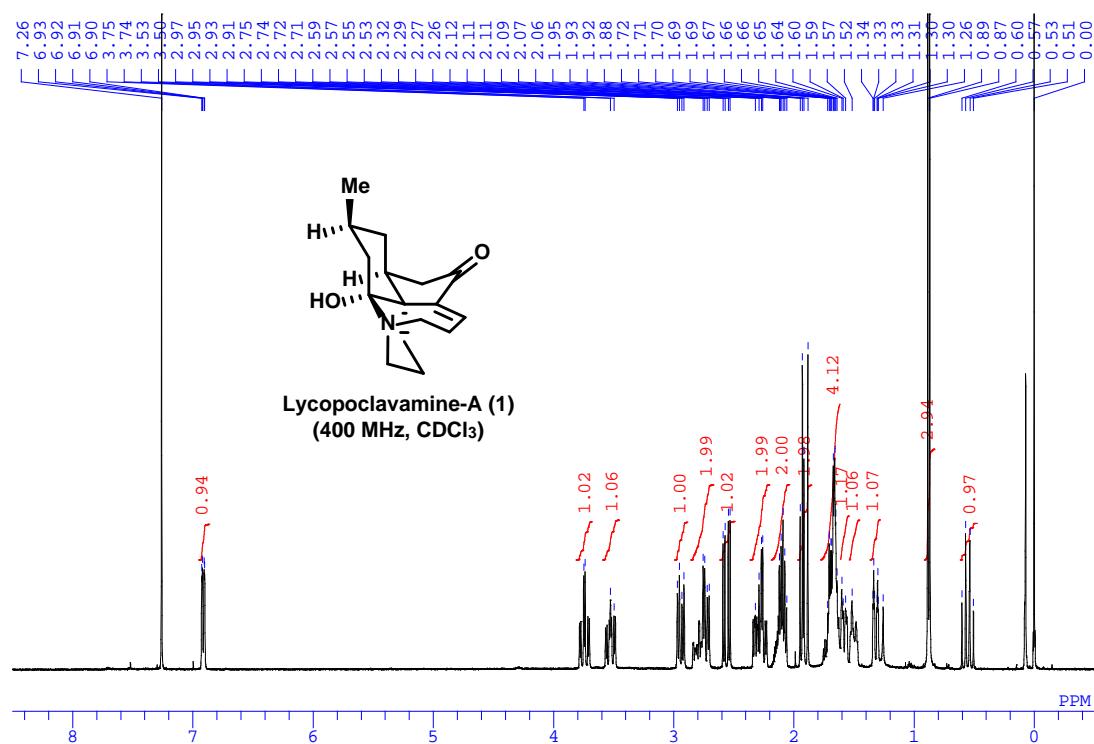


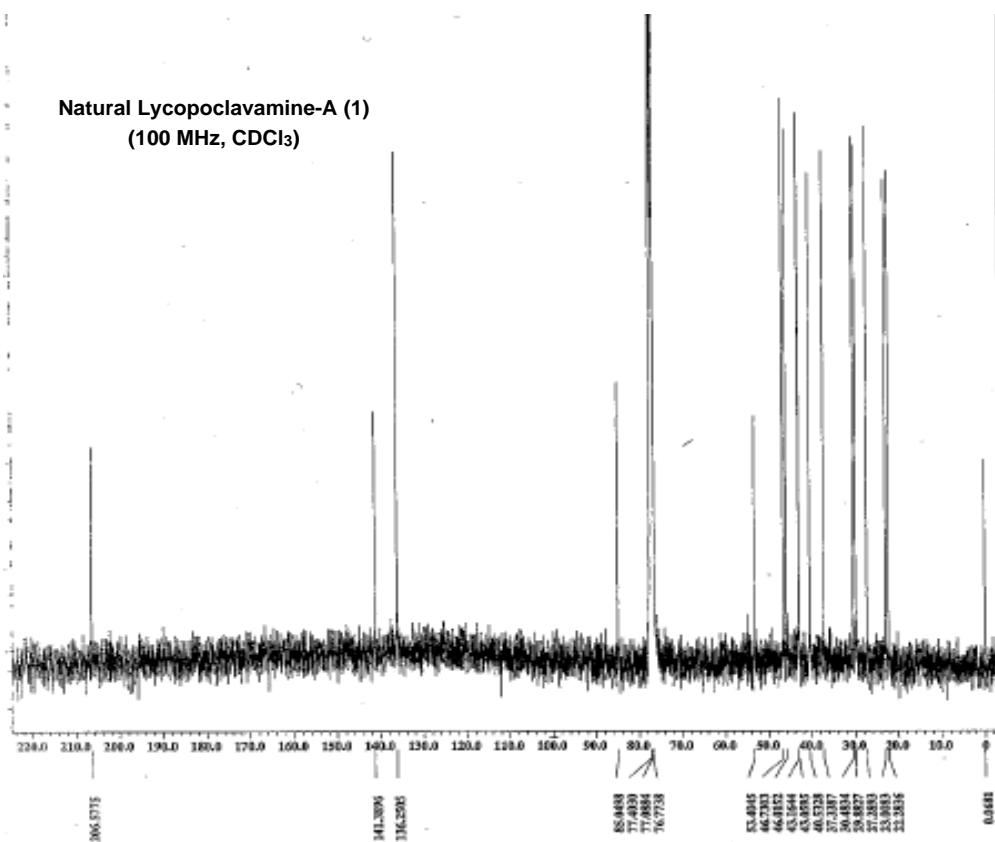
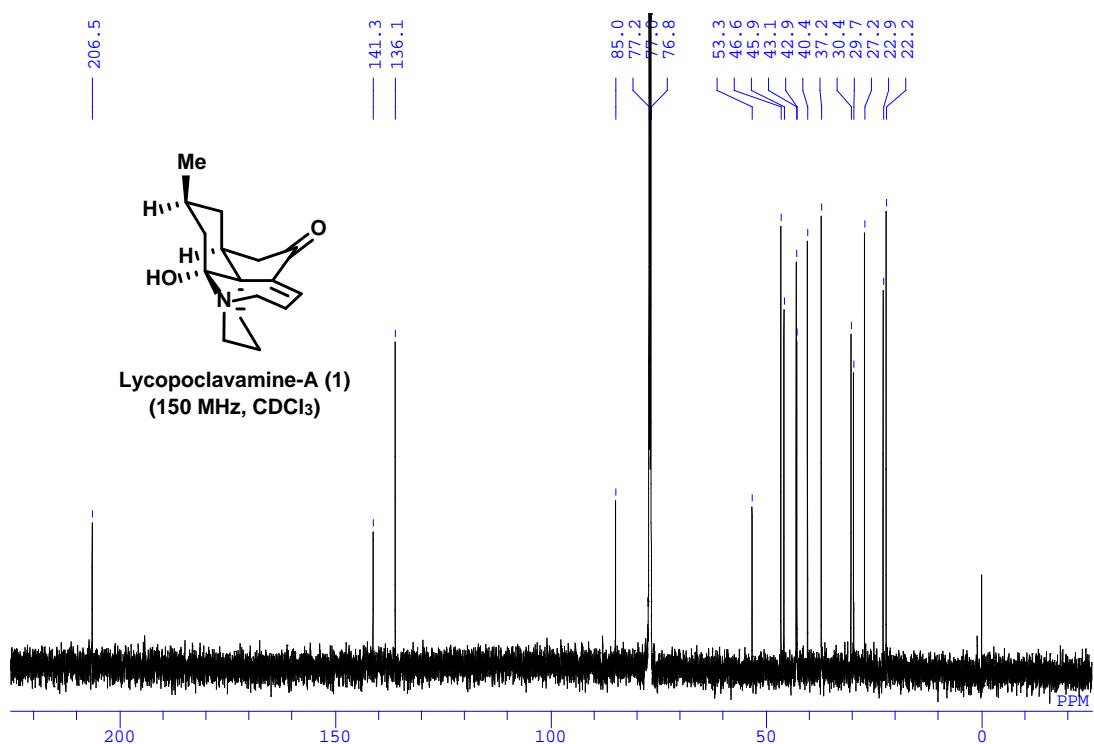


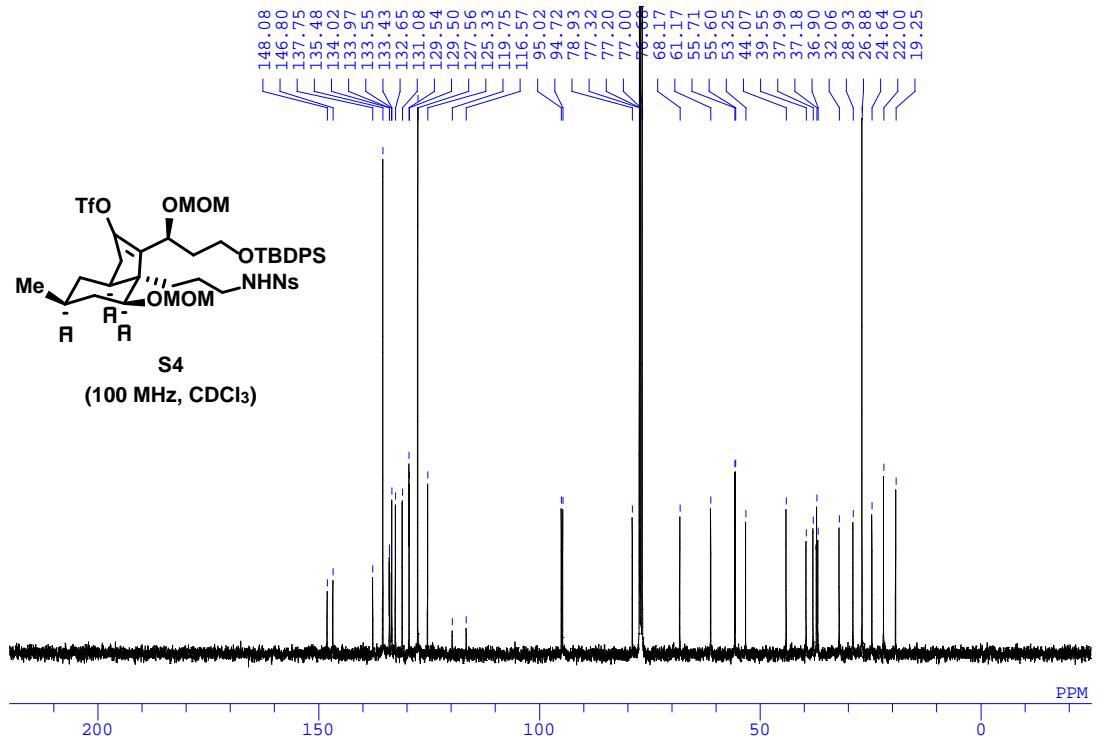
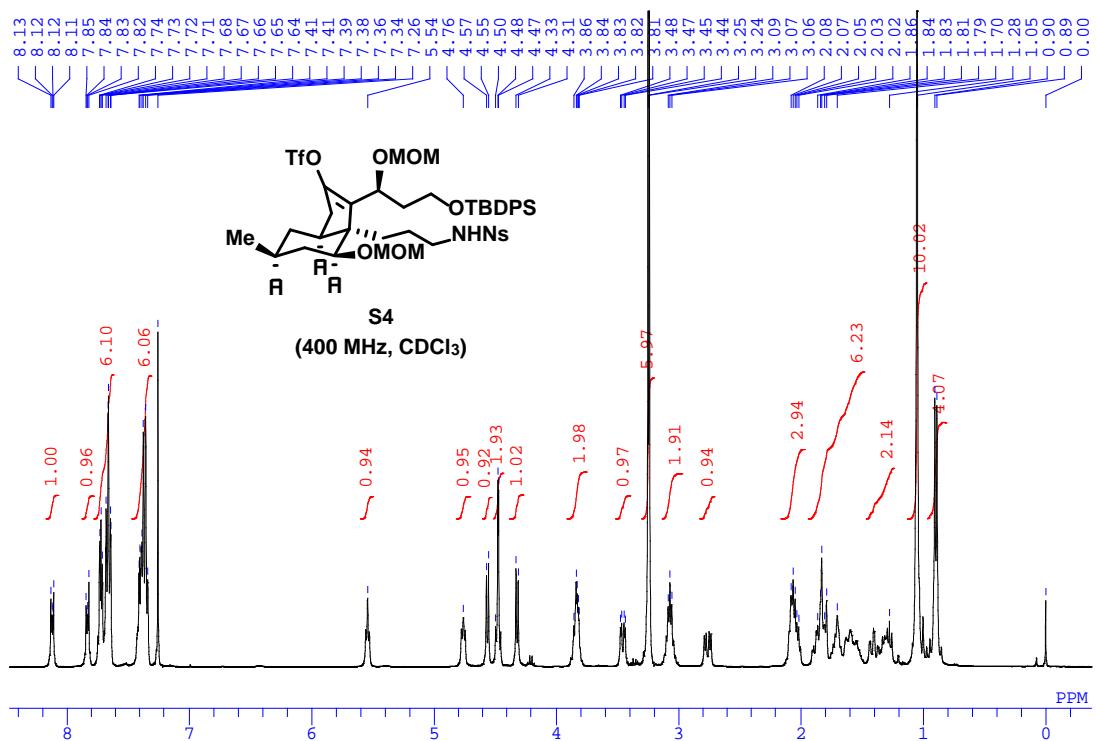


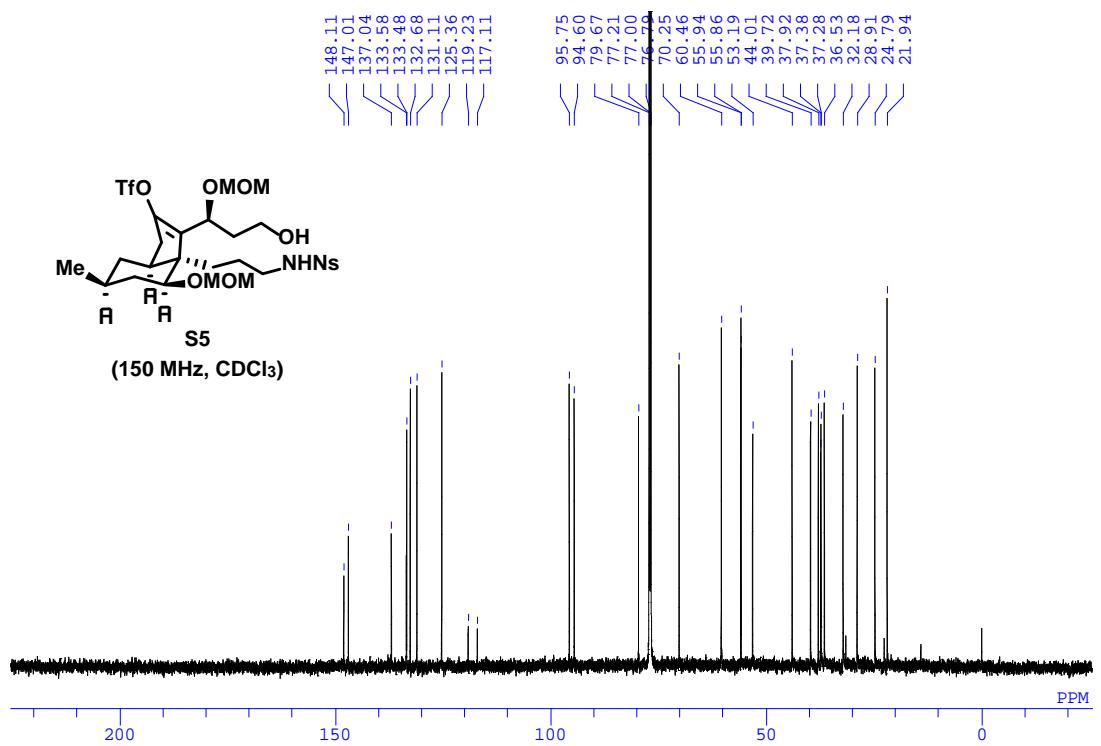
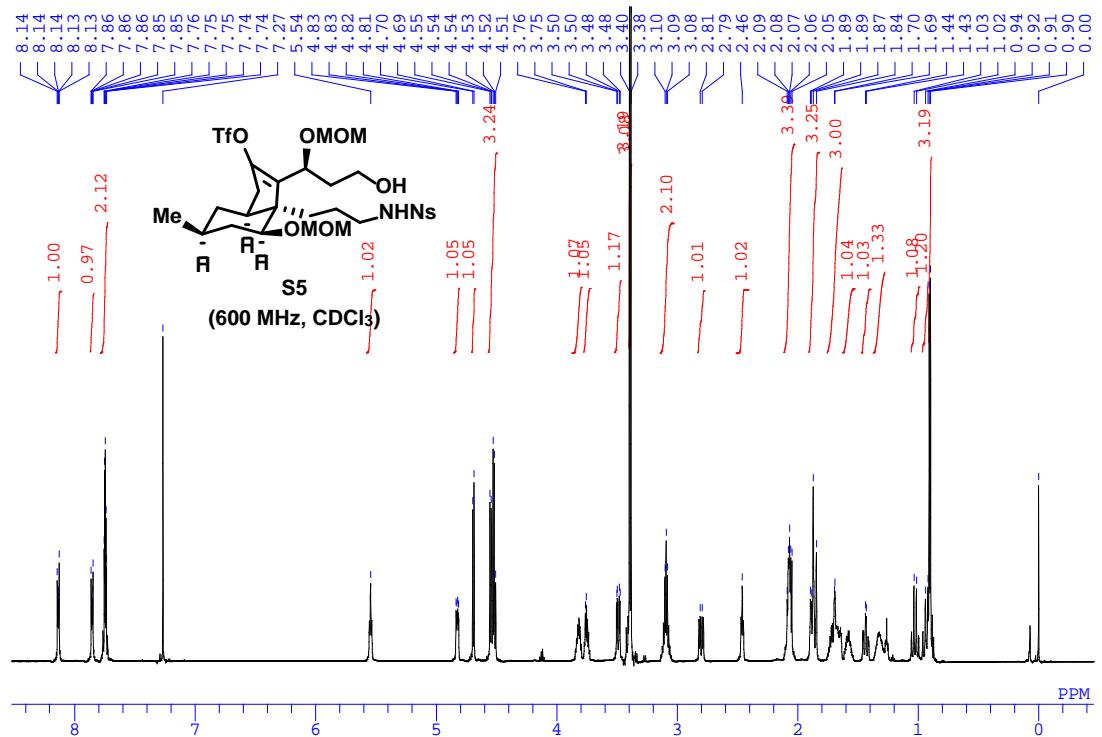


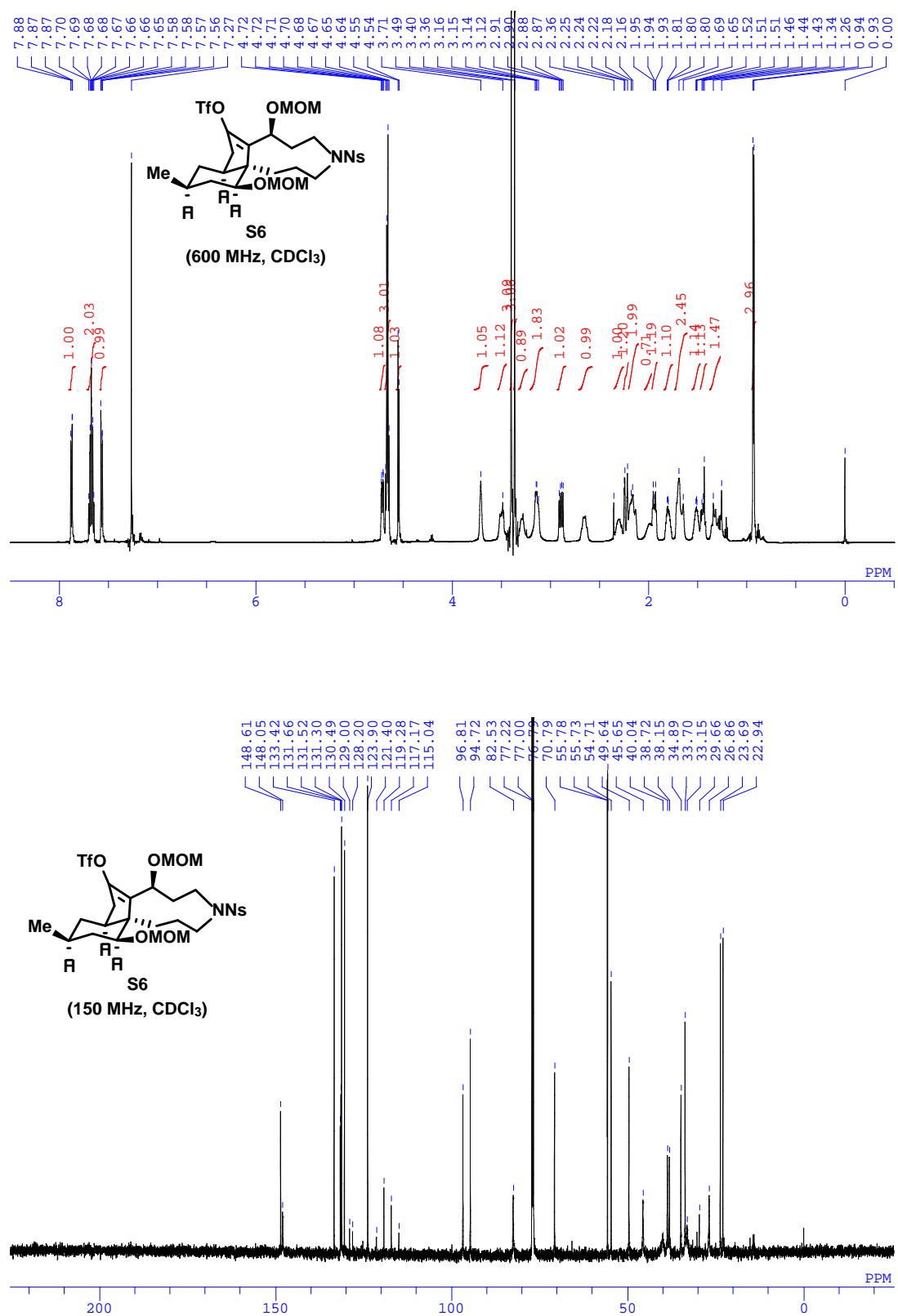


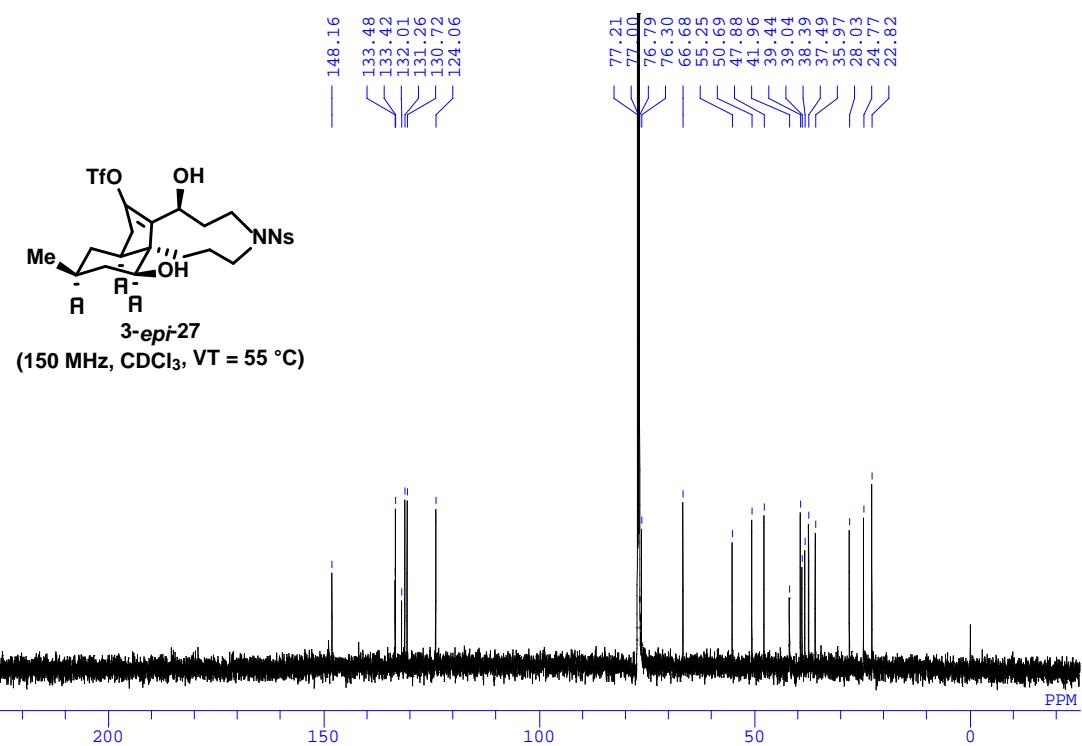
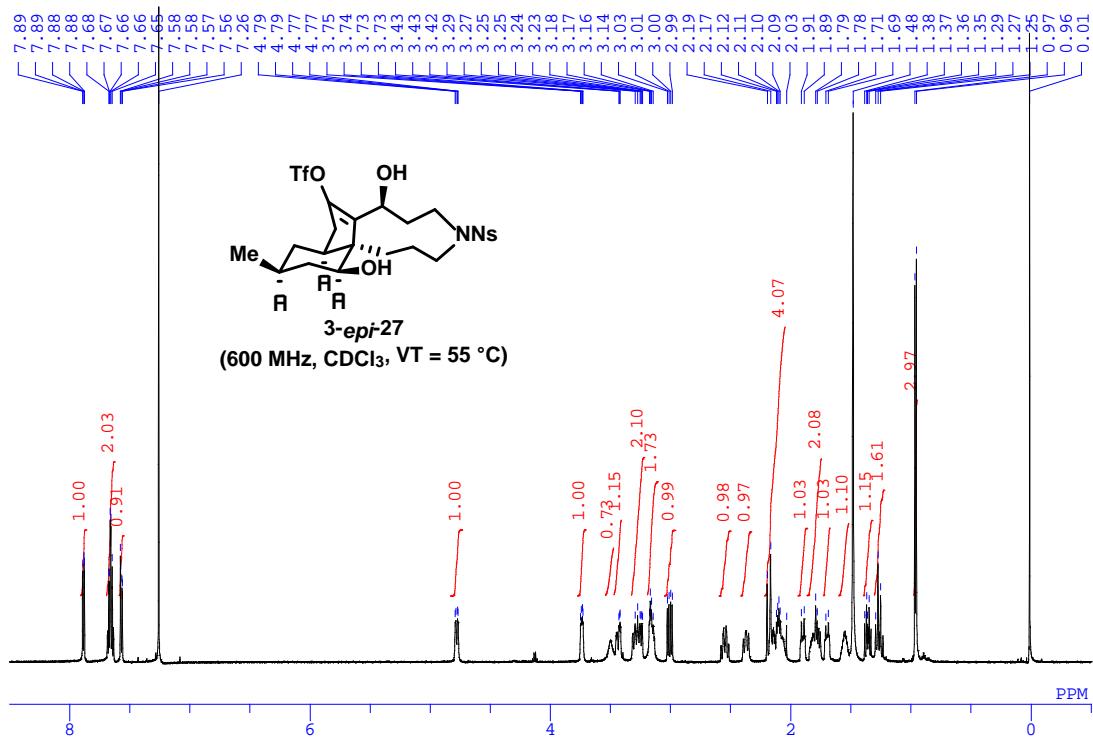


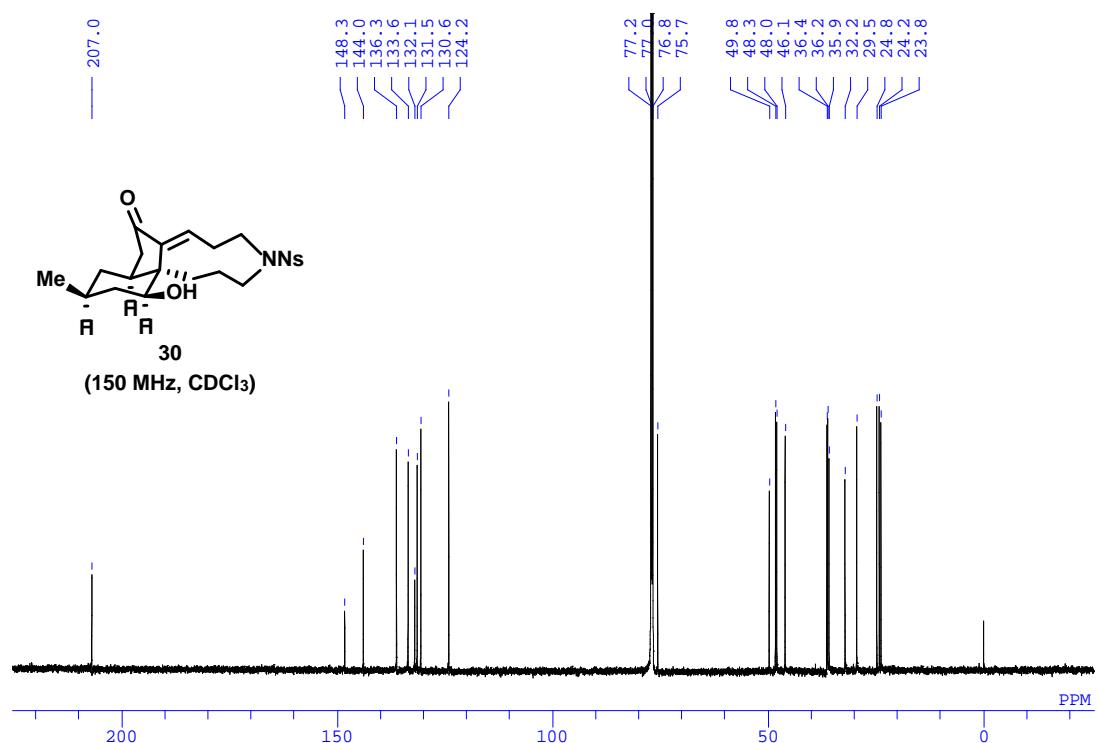
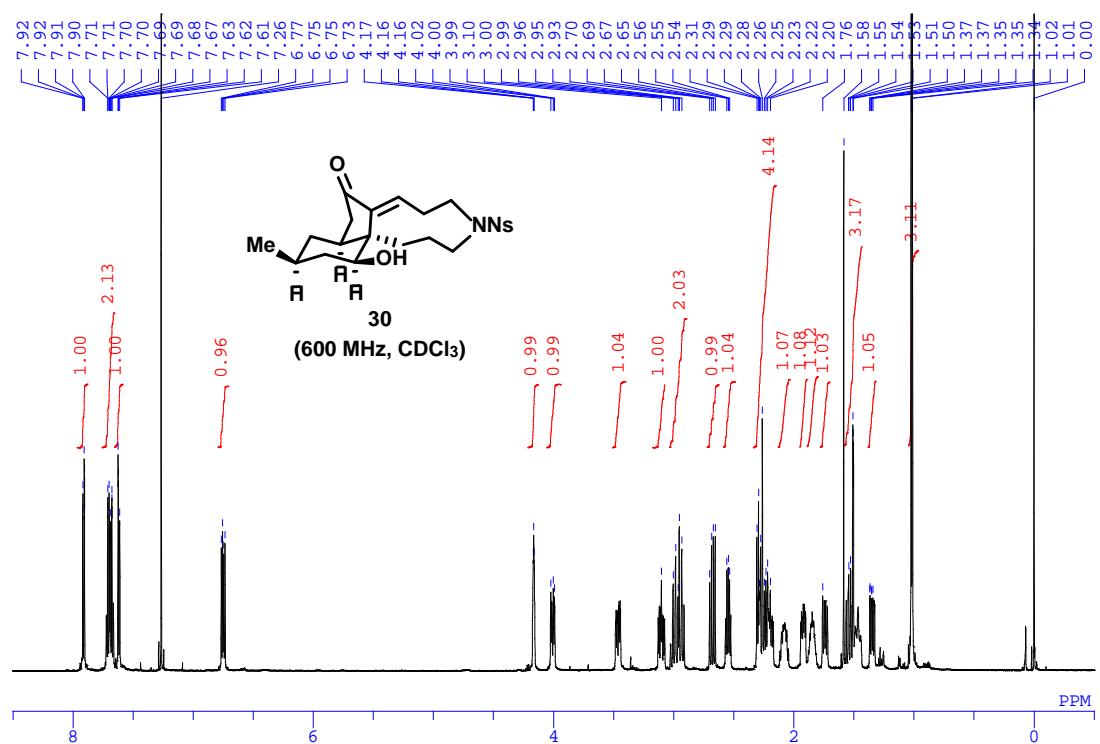


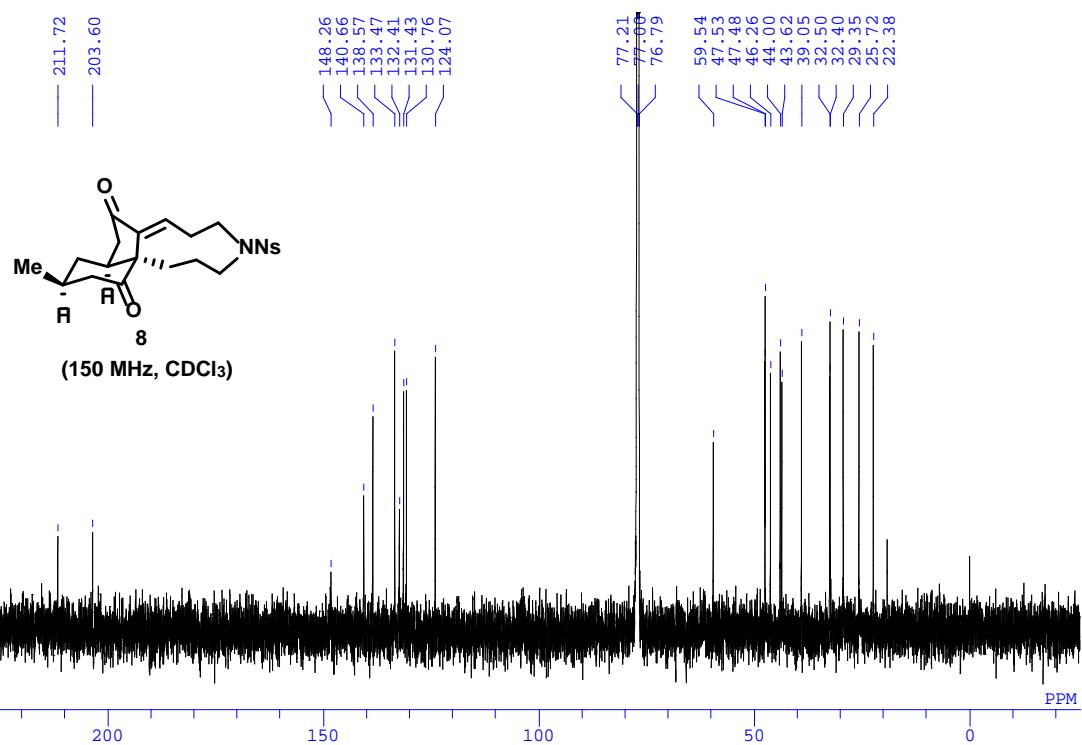
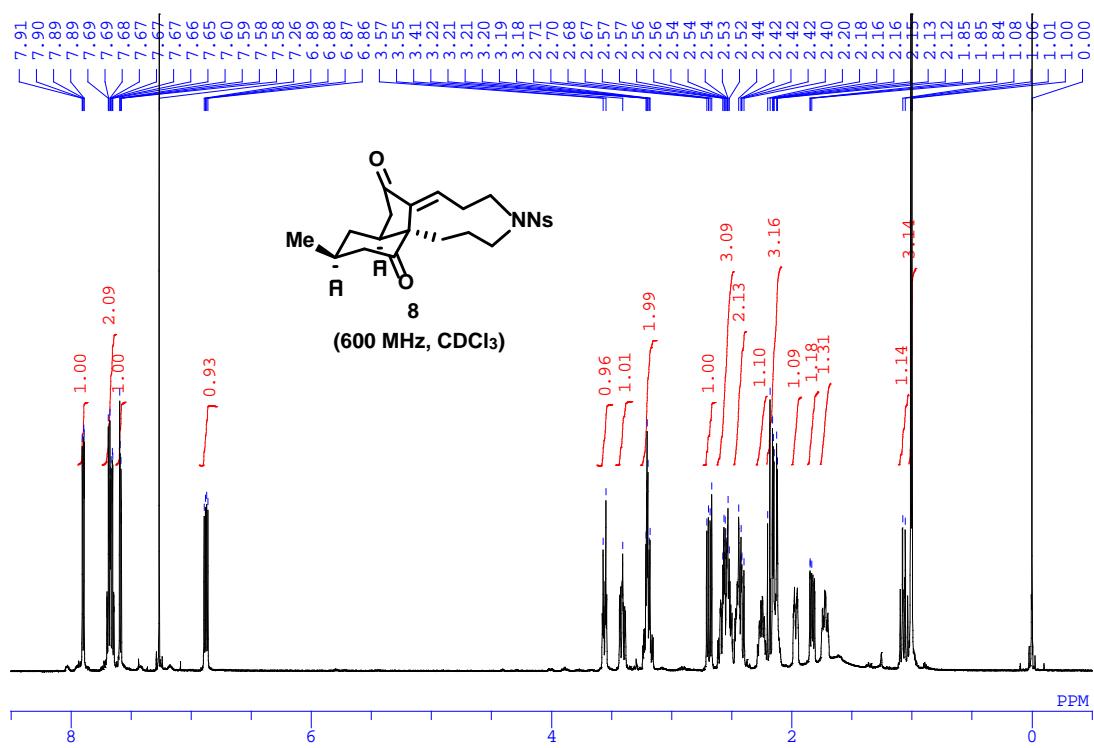












X-ray crystallographic analysis for 3-*epi*27

3-*epi*27 was crystallized from AcOEt/*n*-hexane by the vapor diffusion method. The ORTEP drawing is shown in Figure S1. The crystal structure has been deposited at the Cambridge Crystallographic Centre (deposition number: CCDC 1895333). X-Ray data were collected on a Rigaku R-AXIS RAPID imaging plate diffractometer using a graphite monochromater with Cu-K α radiation ($\lambda = 1.54187 \text{ \AA}$) at 93 K. Crystal Data: C₂₃H₂₉F₃N₂O₉S₂, $M = 598.61$, $a = 14.4545(4) \text{ \AA}$, $b = 14.4545(4) \text{ \AA}$, $c = 11.1066(3) \text{ \AA}$, $\alpha = \beta = 90^\circ$, $\gamma = 120^\circ$, $V = 2009.64(10) \text{ \AA}^3$, space group: P3₁, Z = 3. Refine extinction method: SHELXL.

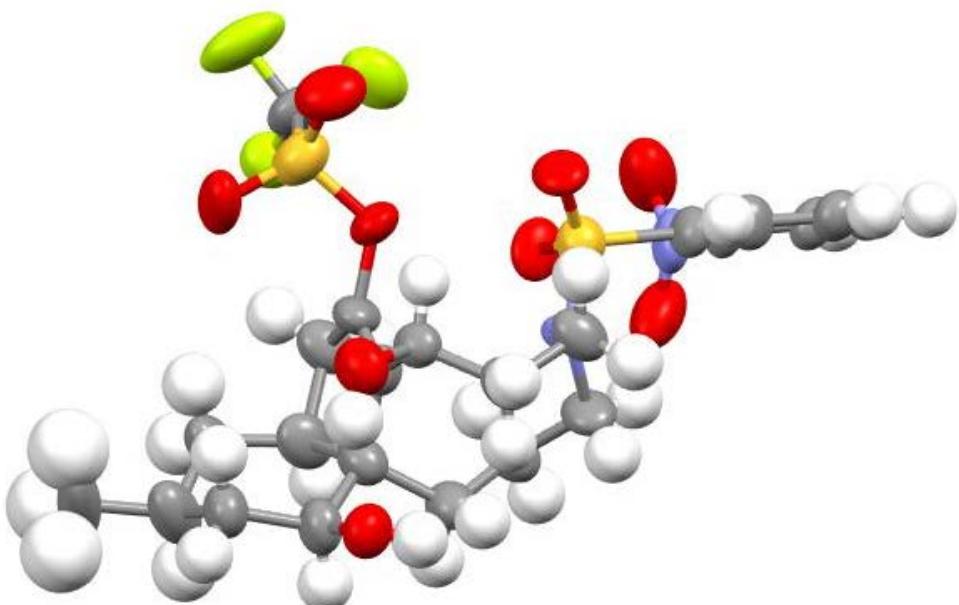


Fig. S1. ORTEP drawing of 3-*epi*27 showing thermal ellipsoids at the 50% probability level.