Syntheses of (-)-Cryptocaryolone and (-)-

Cryptocaryolone Diacetate via a Diastereoselective Oxy-

Michael Addition and Oxocarbenium Allylation

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General Procedure

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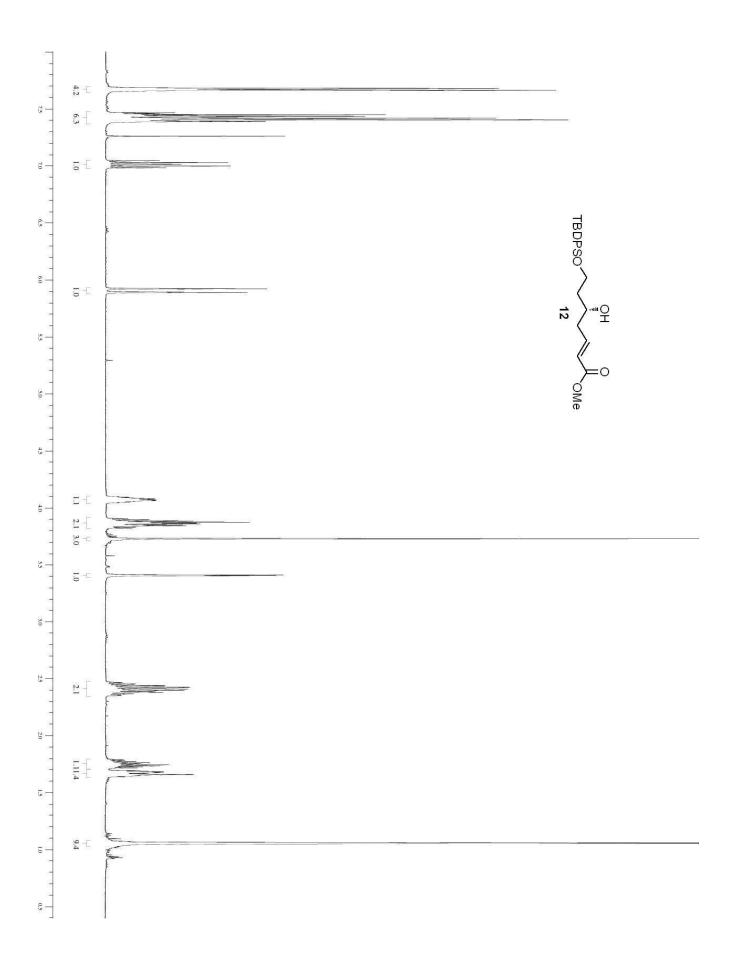
¹H and ¹³C NMR spectra of all compounds

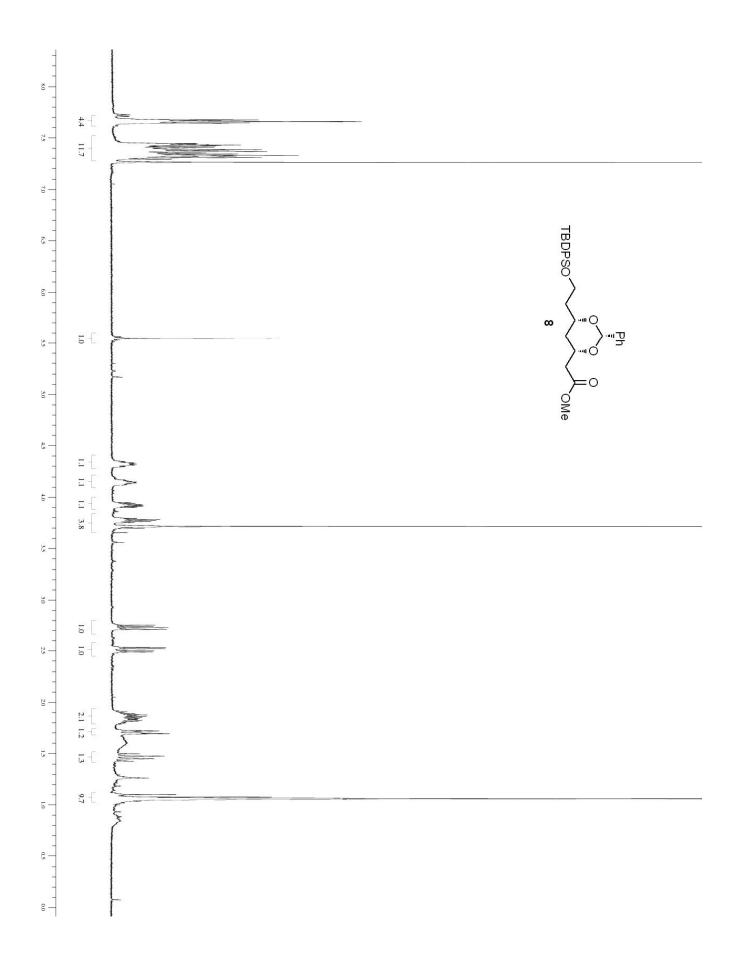
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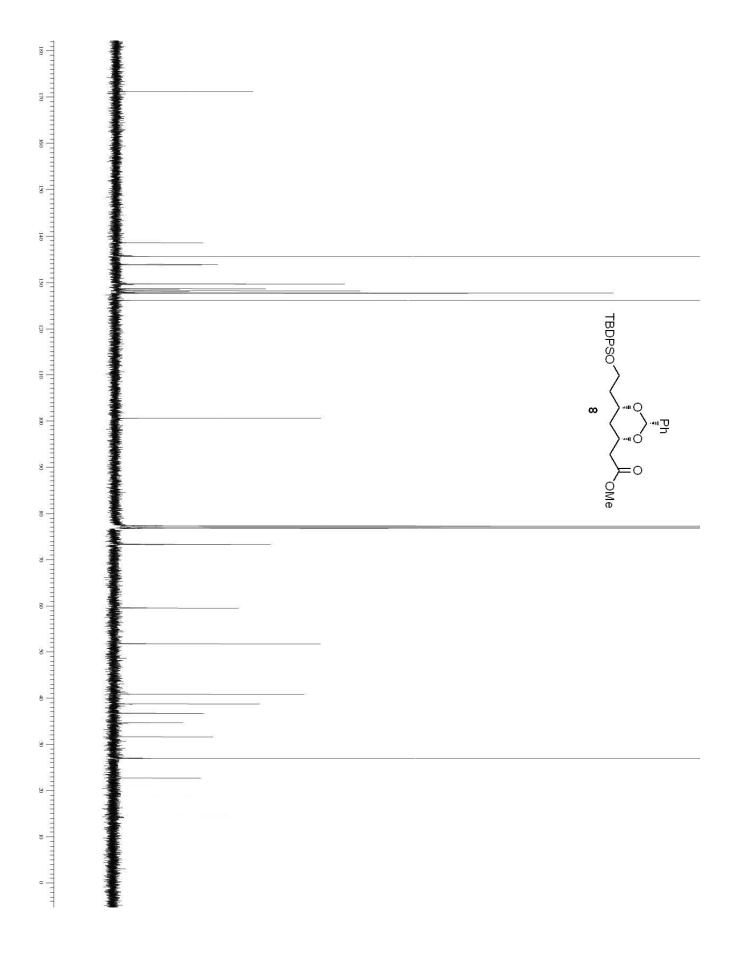
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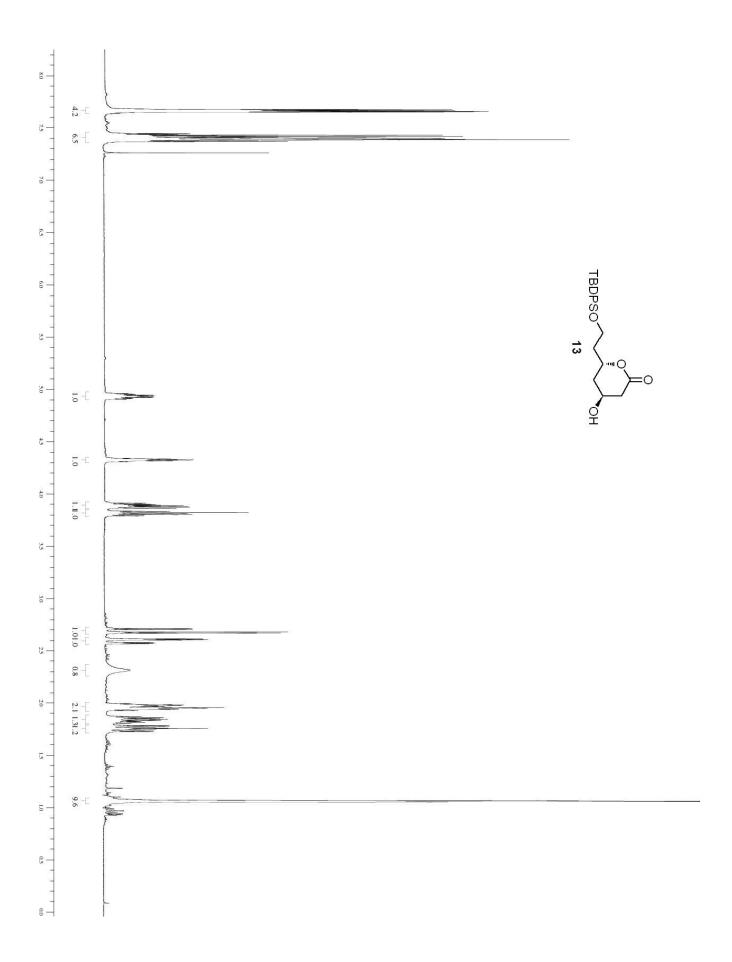
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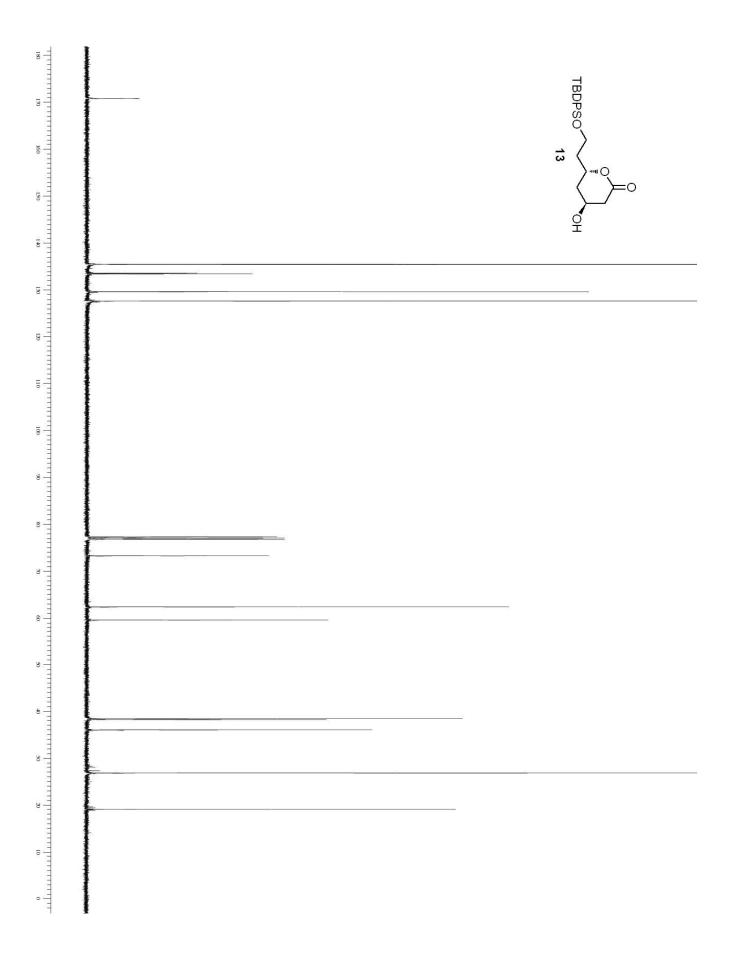
General Procedure. 1 H and 13 C NMR spectra were obtained using either CDCl₃ or C₆D₆ as the solvent with CHCl₃: 7.26 ppm or C₆H₆: 7.15 ppm as internal standards. Chemical shifts are given in δ (ppm) and coupling constants (J) in Hz. All starting materials and solvents were commercially available and were used without further purification. Column chromatography was performed using 60-200 μm silica gel. Analytical thin layer chromatography was performed on silica coated glass plates with F-254 indicator. Visualization was accomplished by UV light (254 nm), KMnO₄, or ceric sulfate-PMA stain. Compound **9** has been previously reported. 1

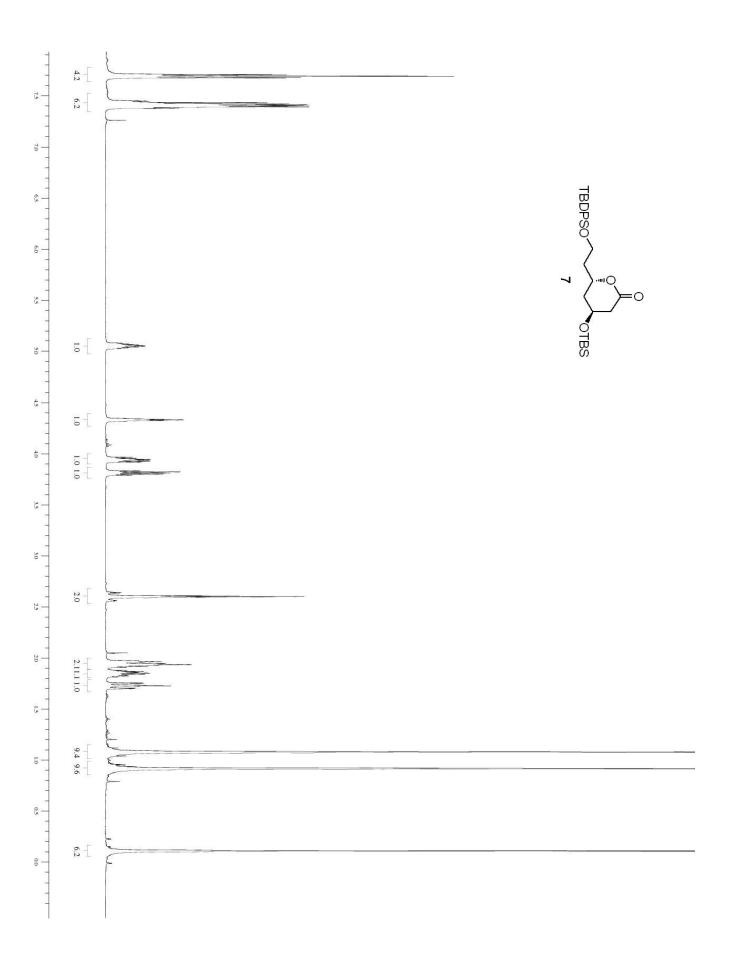


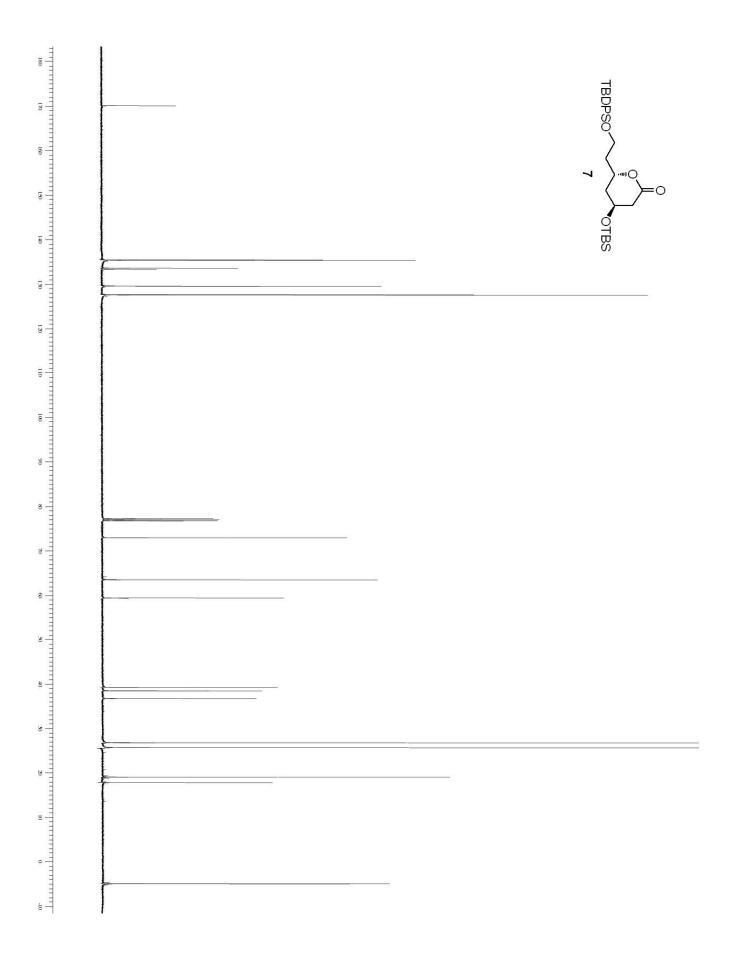


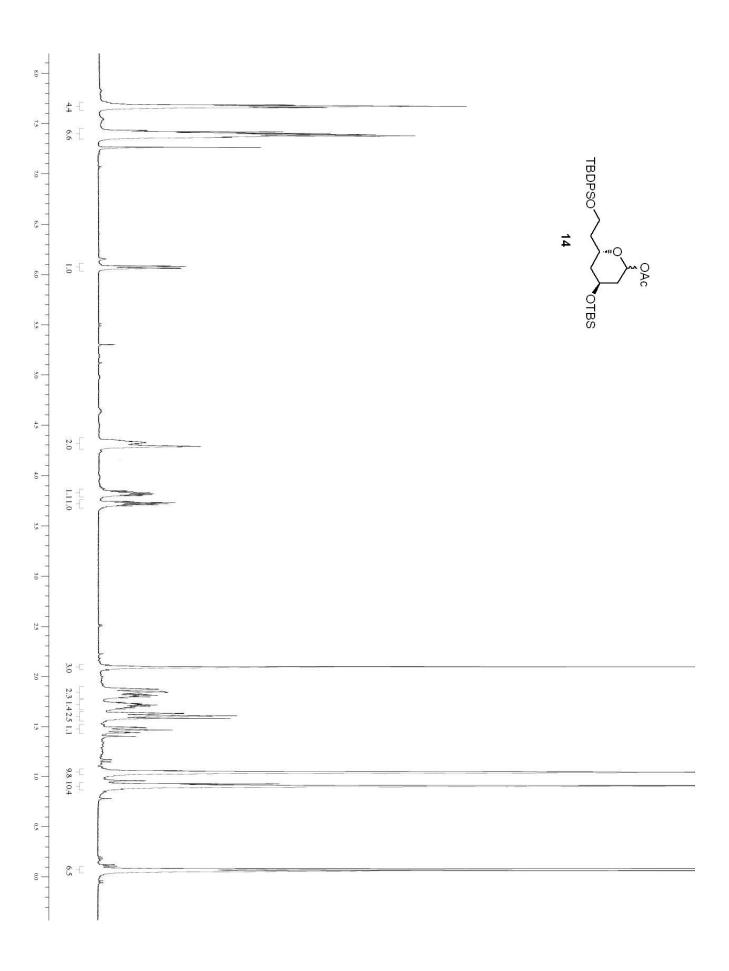


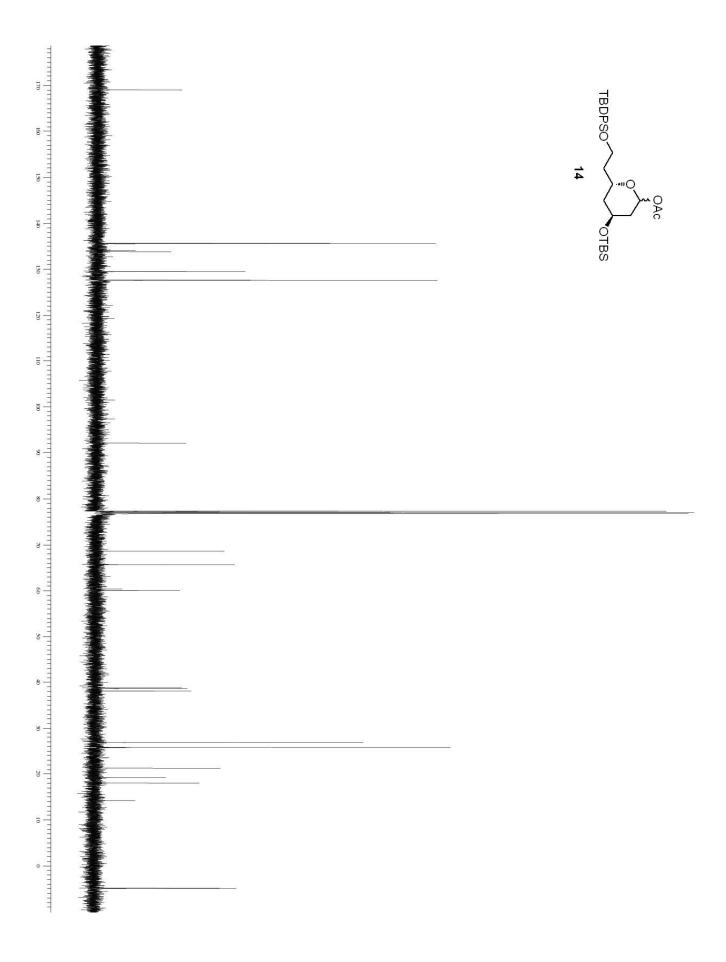


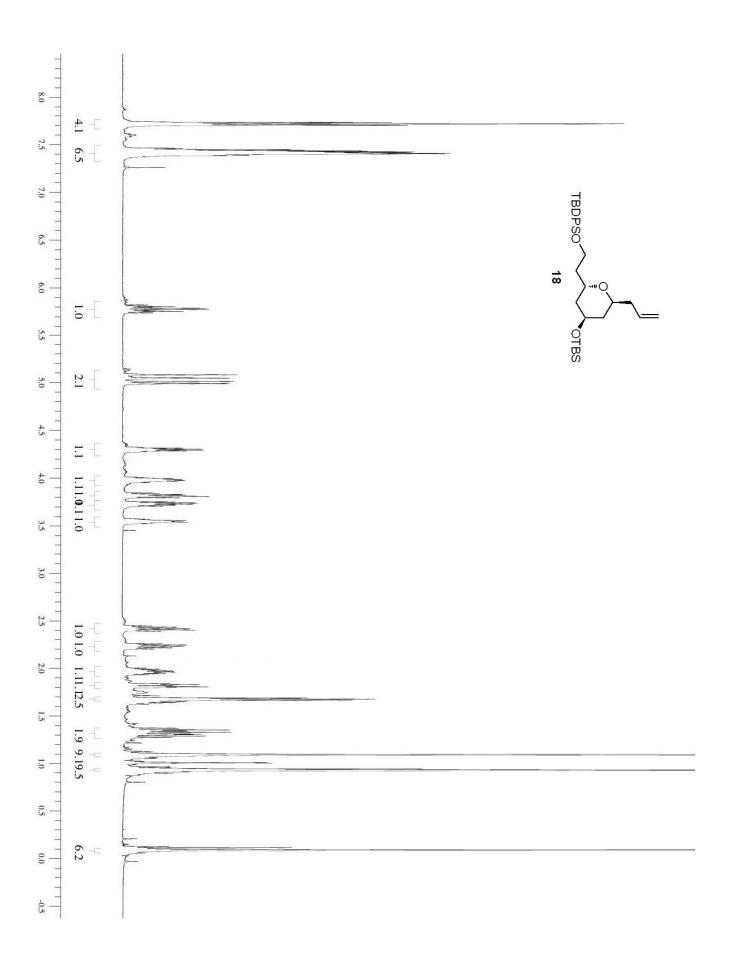


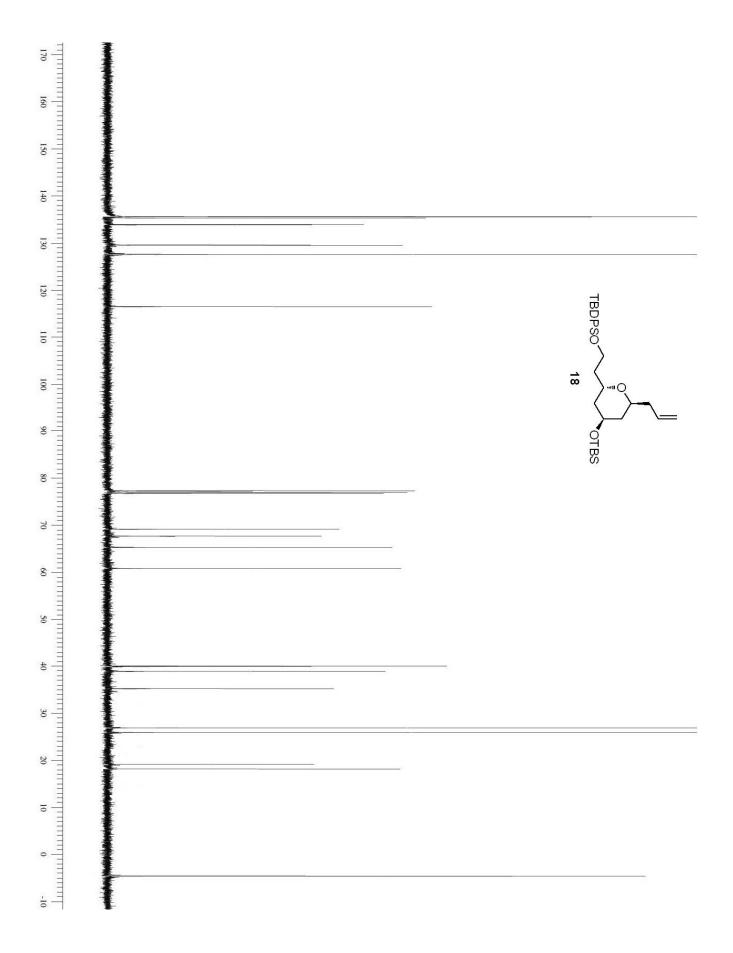


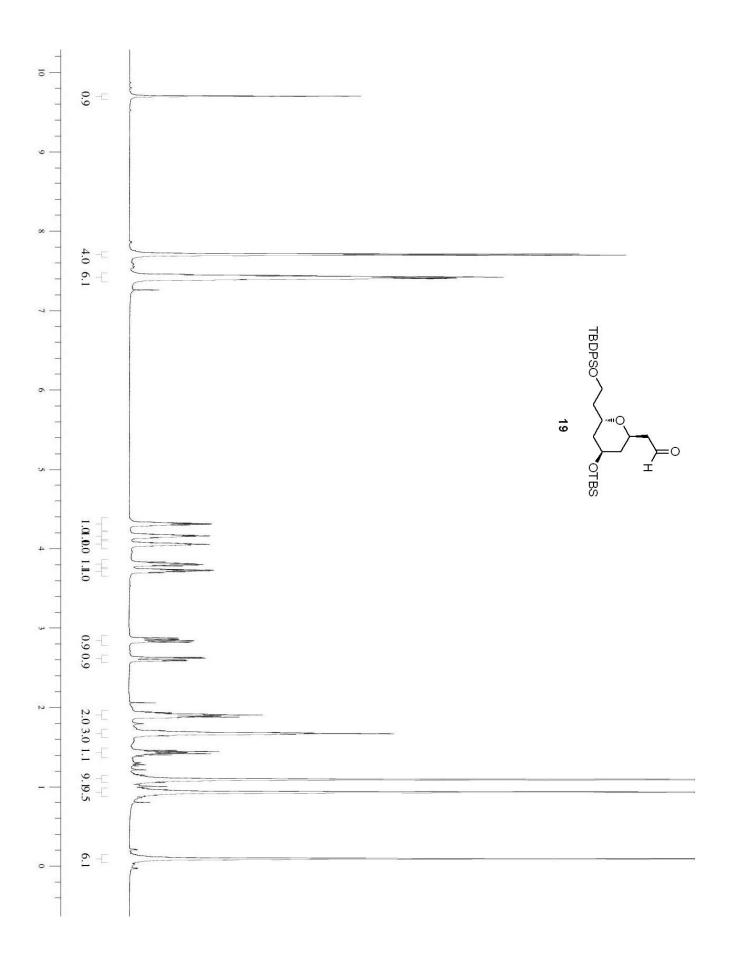


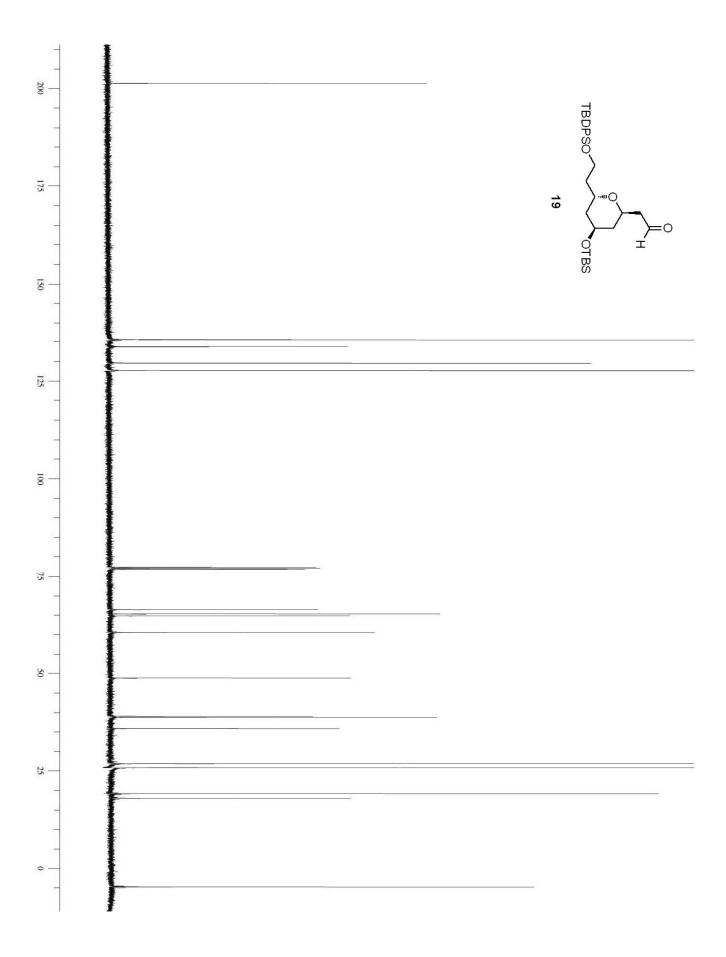


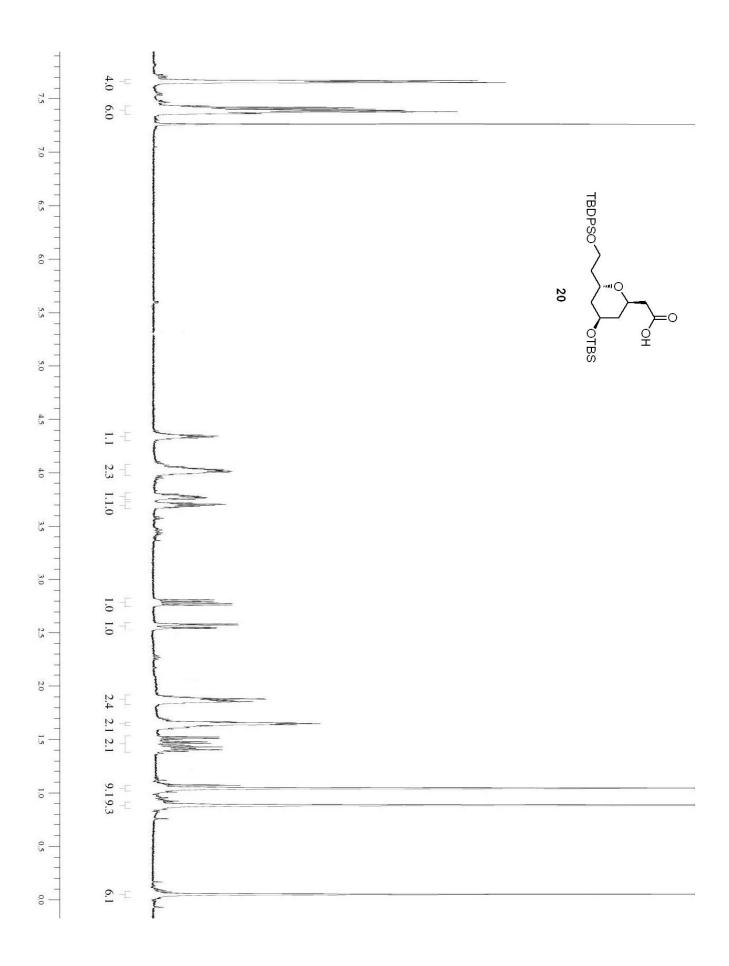


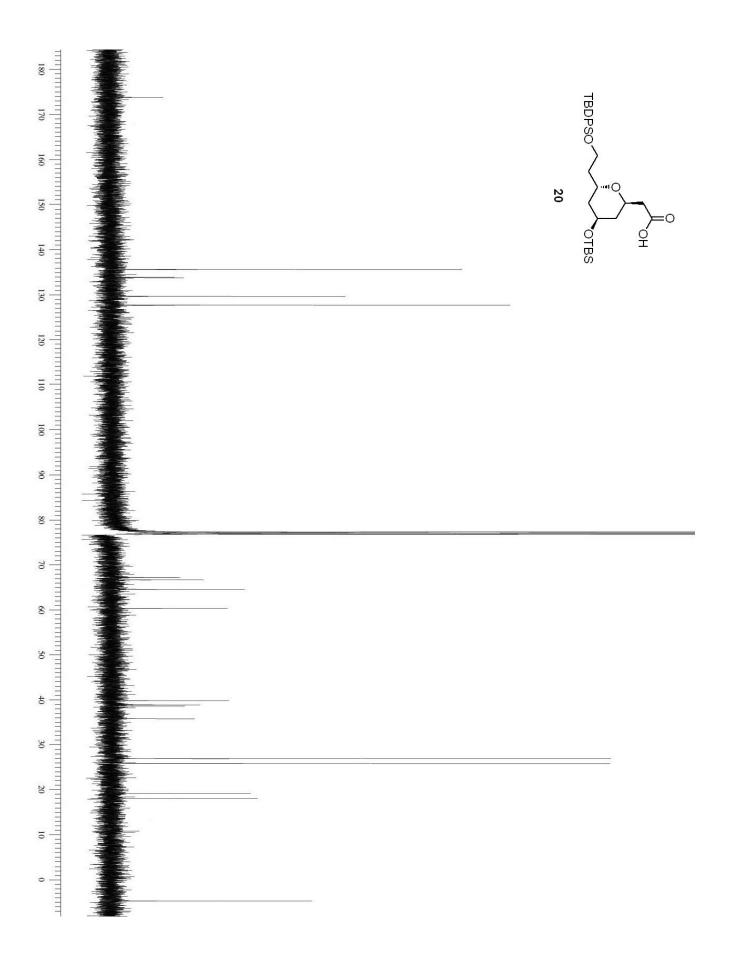


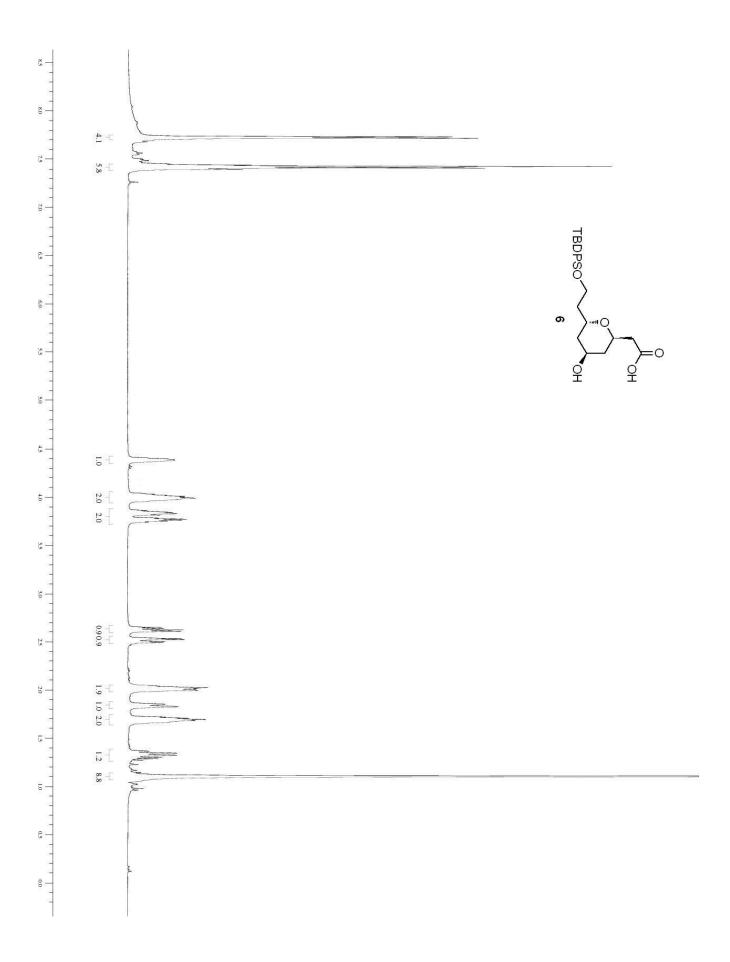


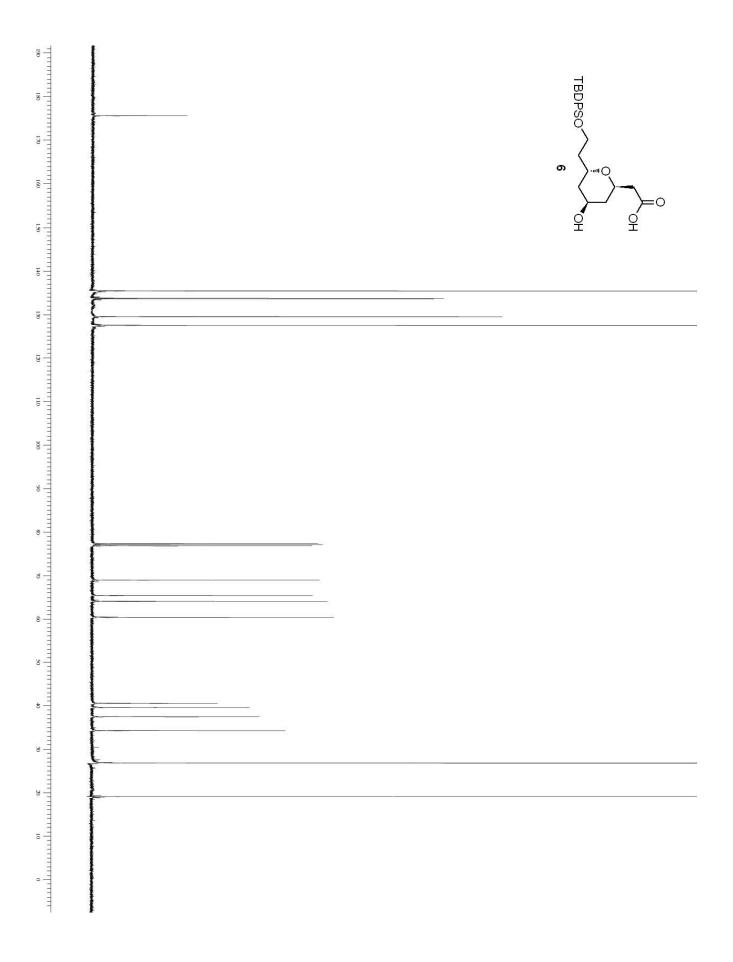


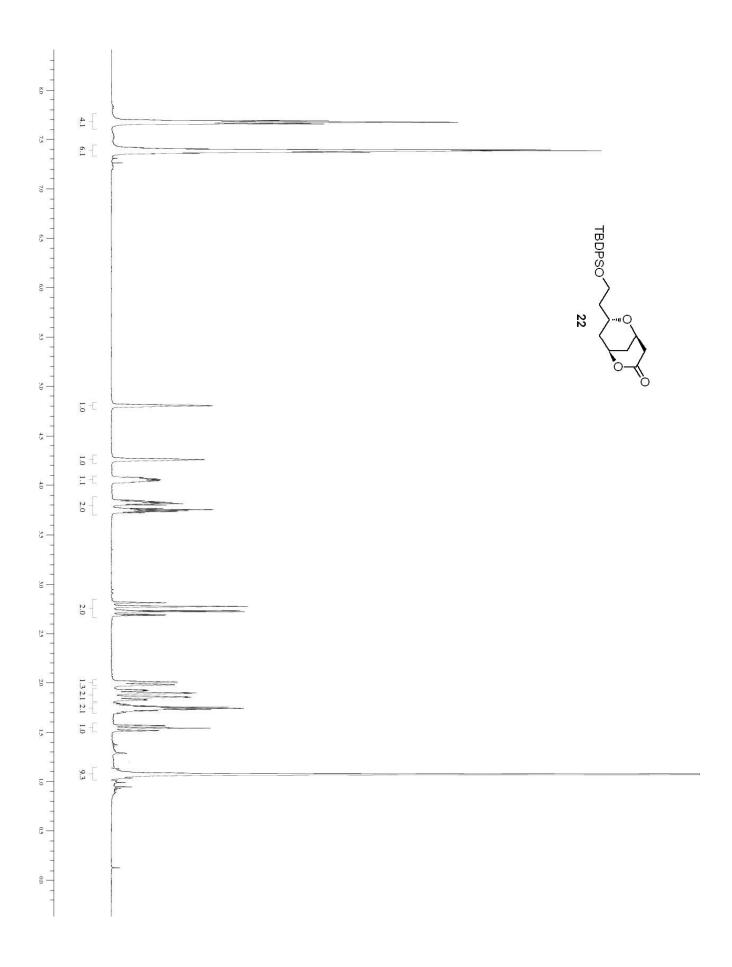


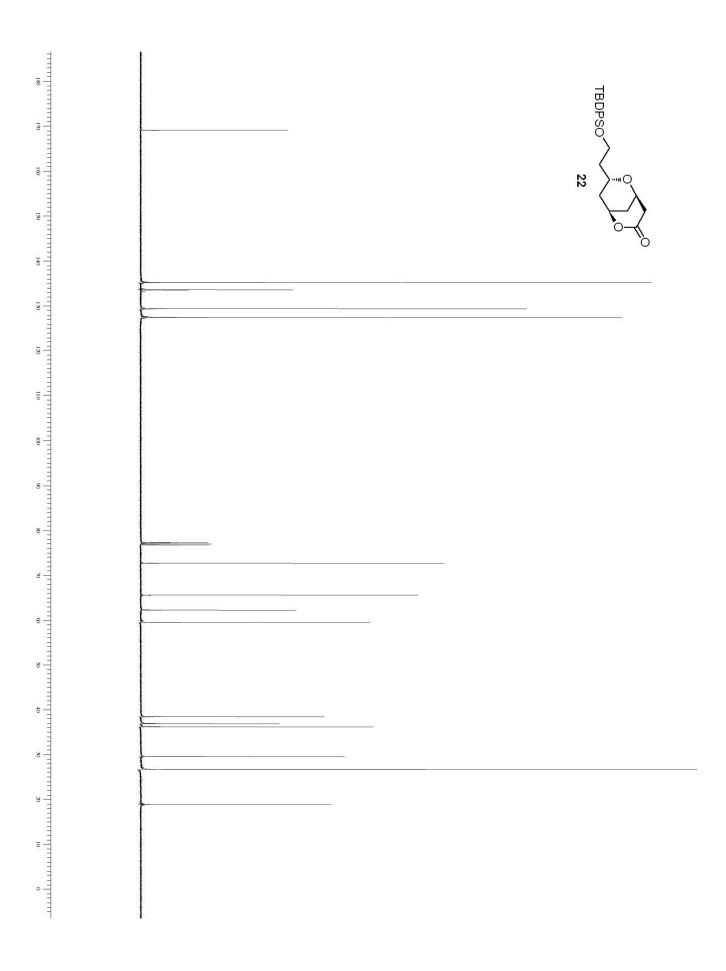


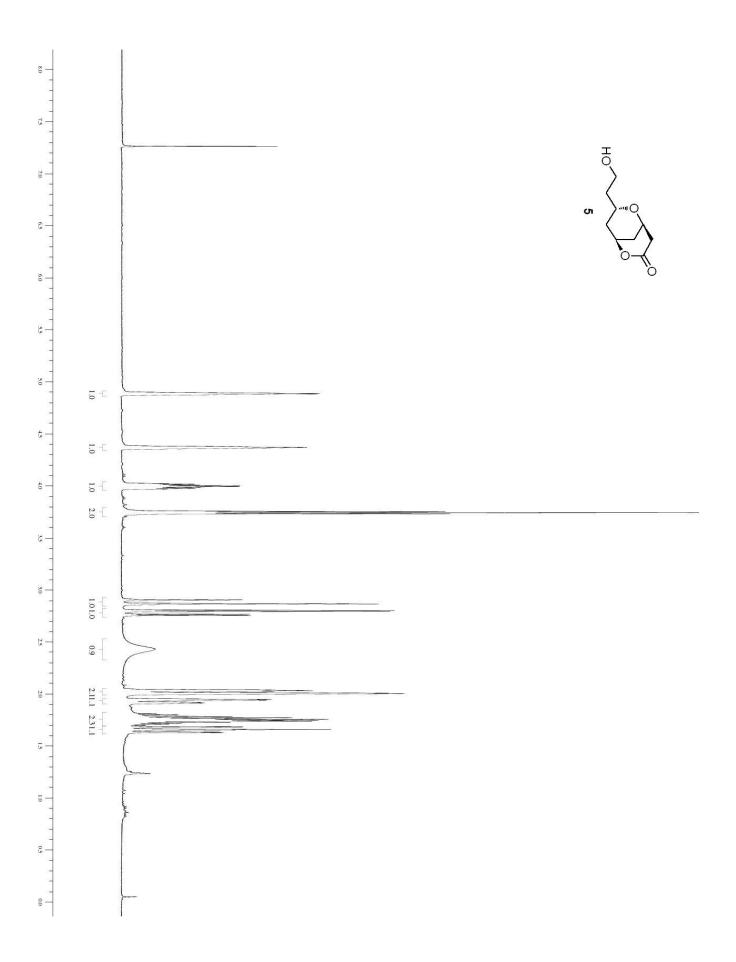


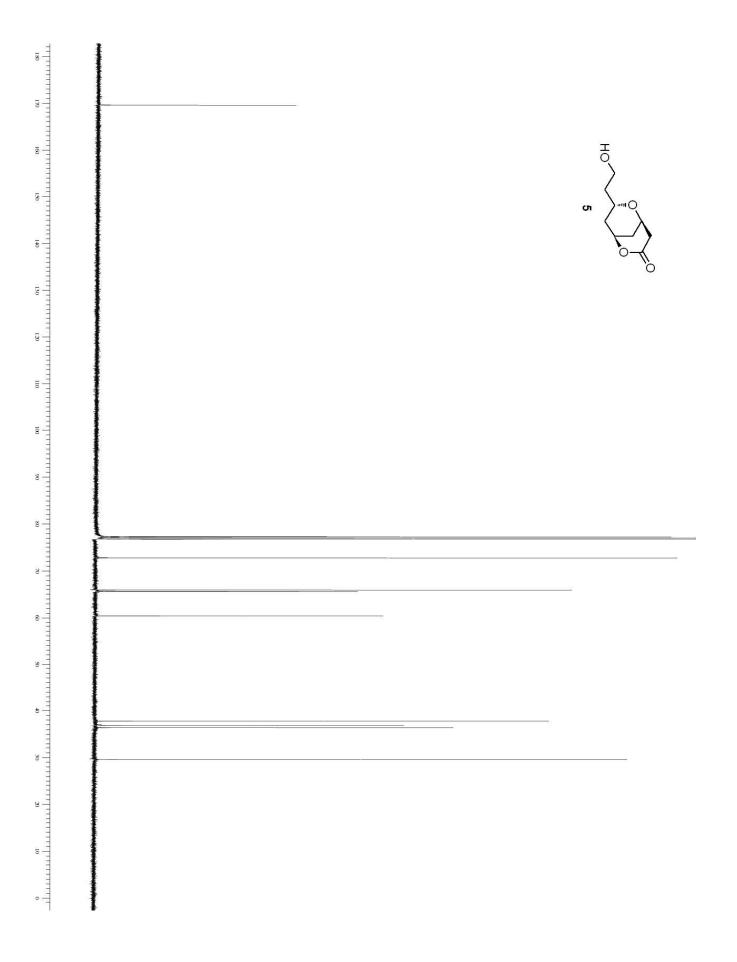


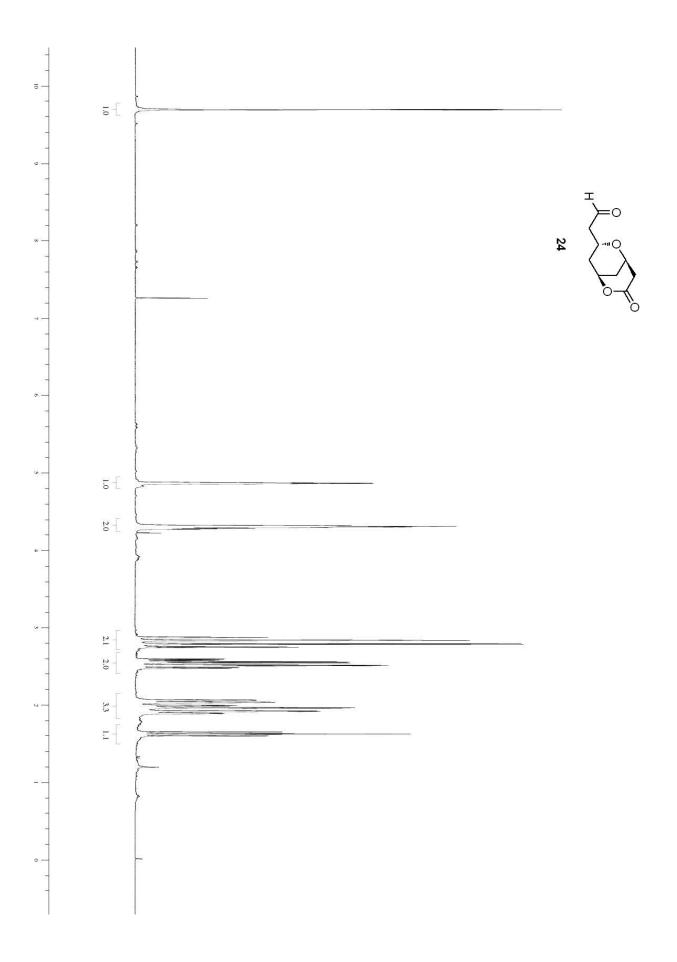


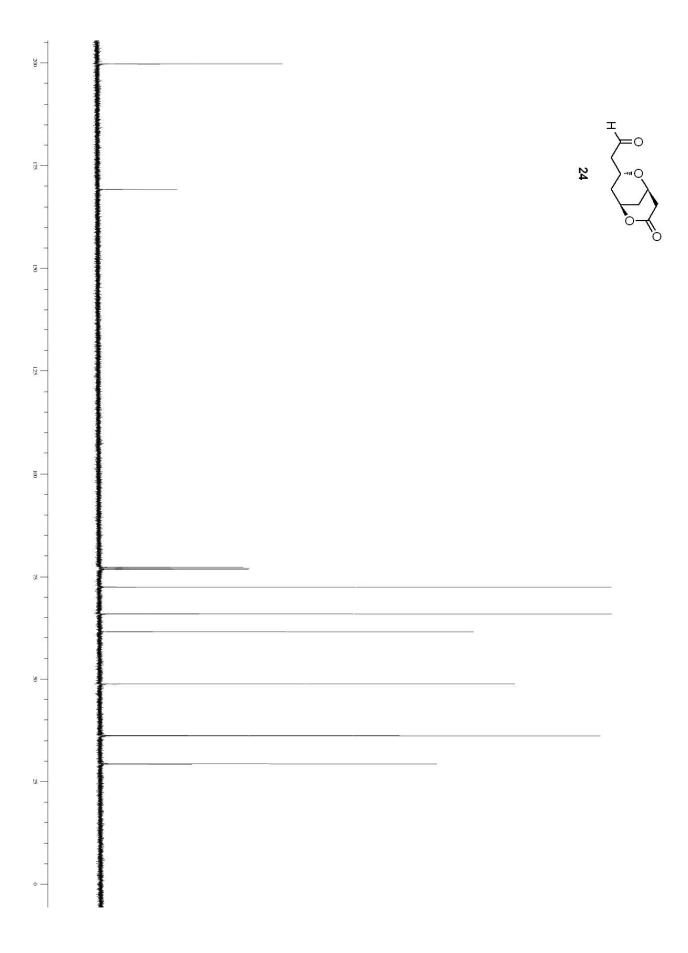


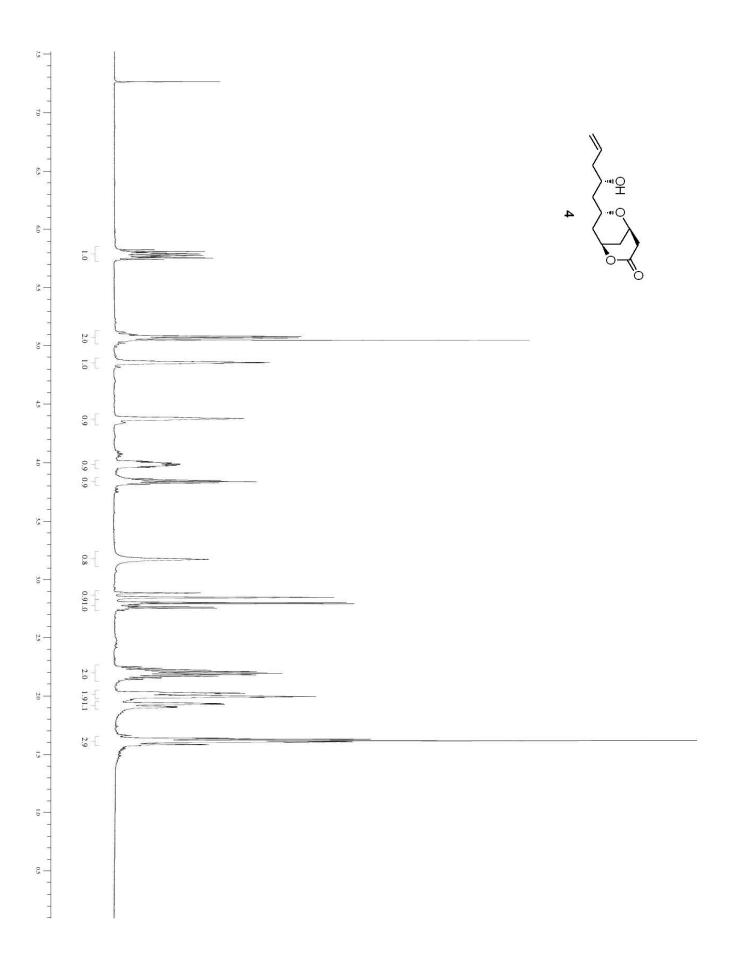


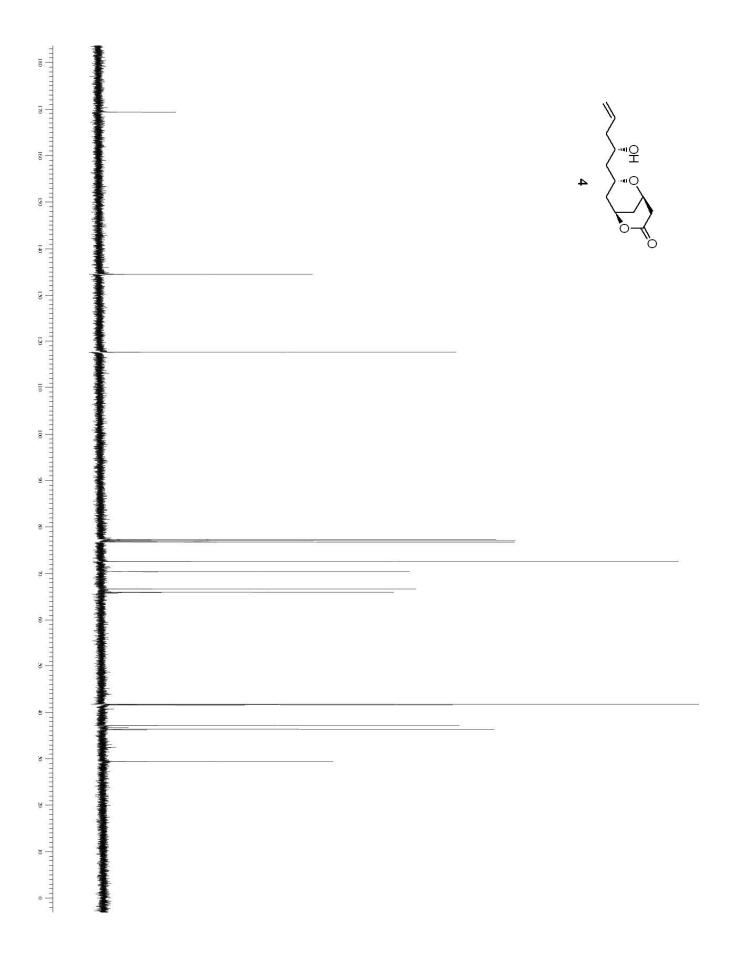


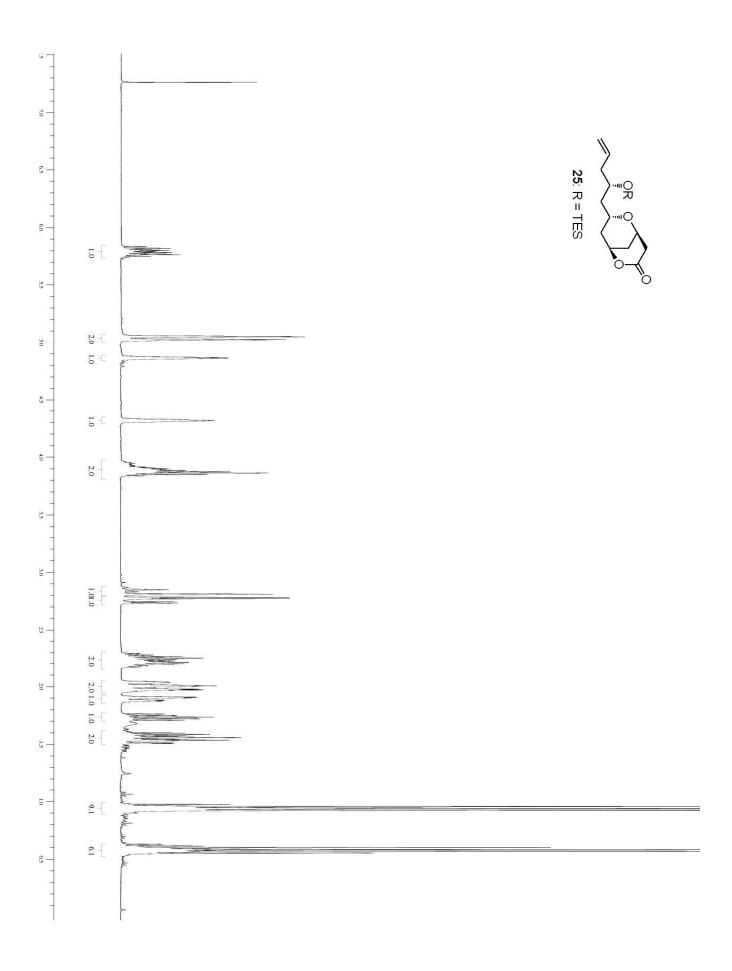


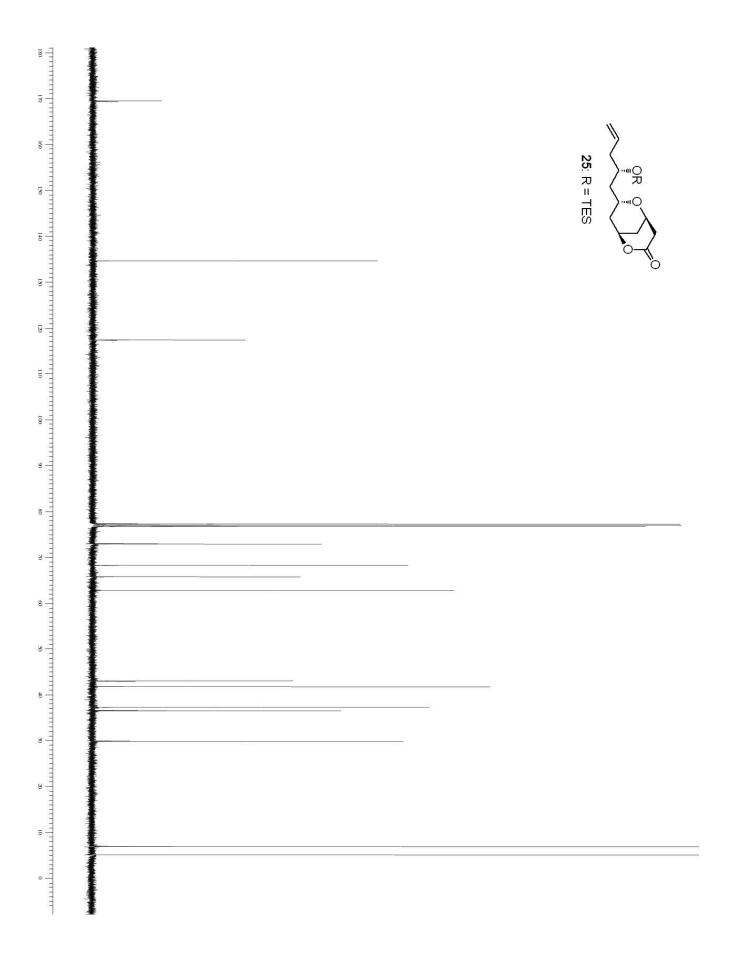


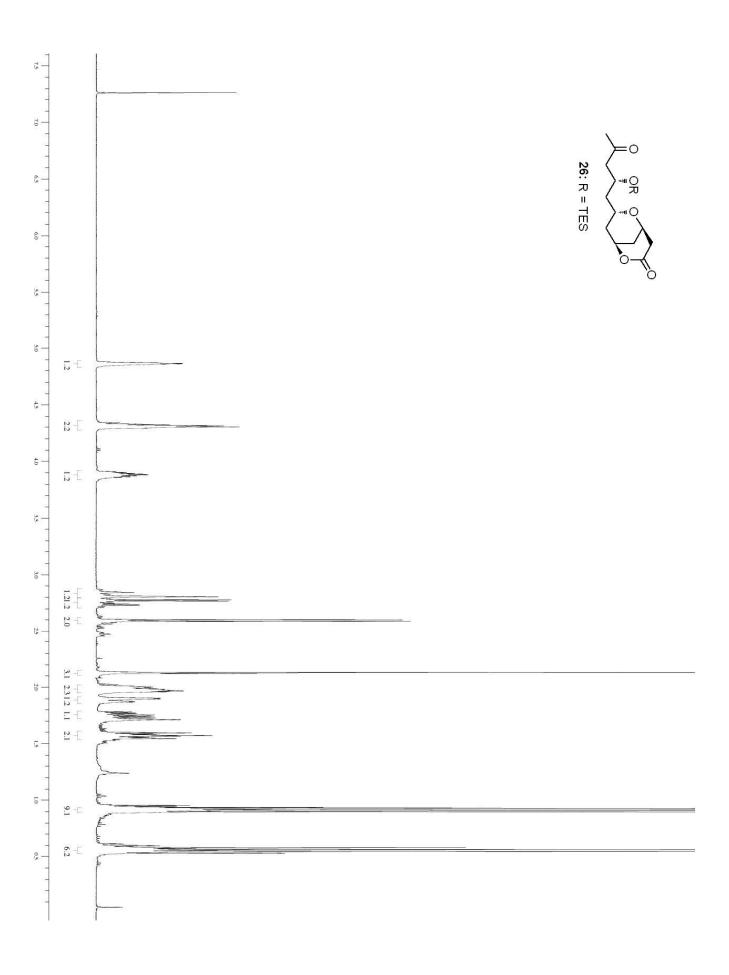


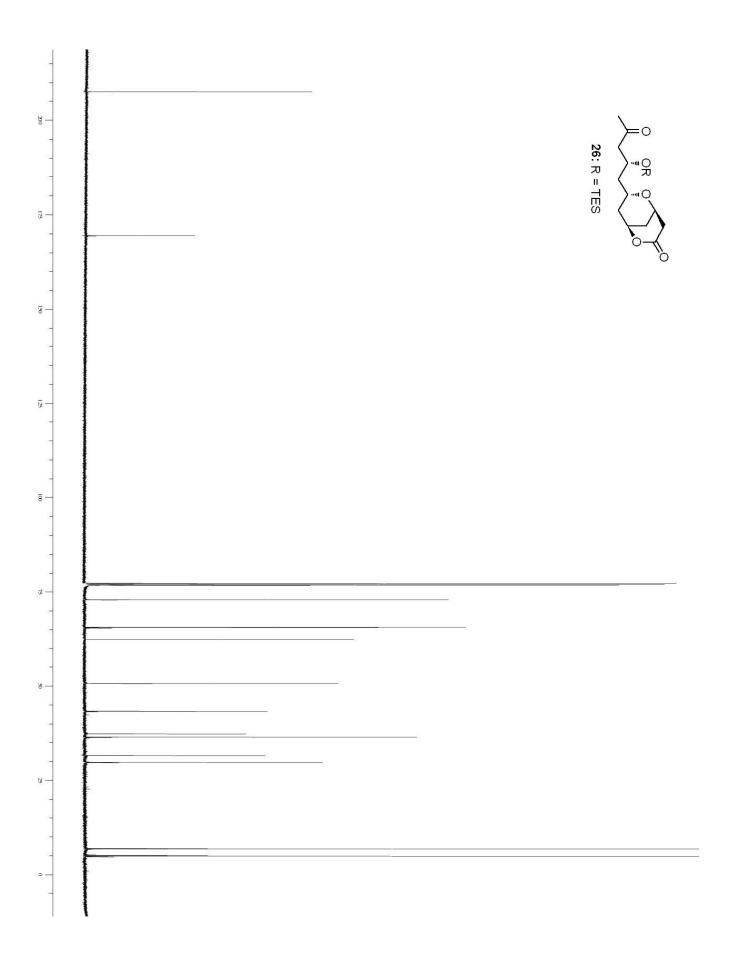


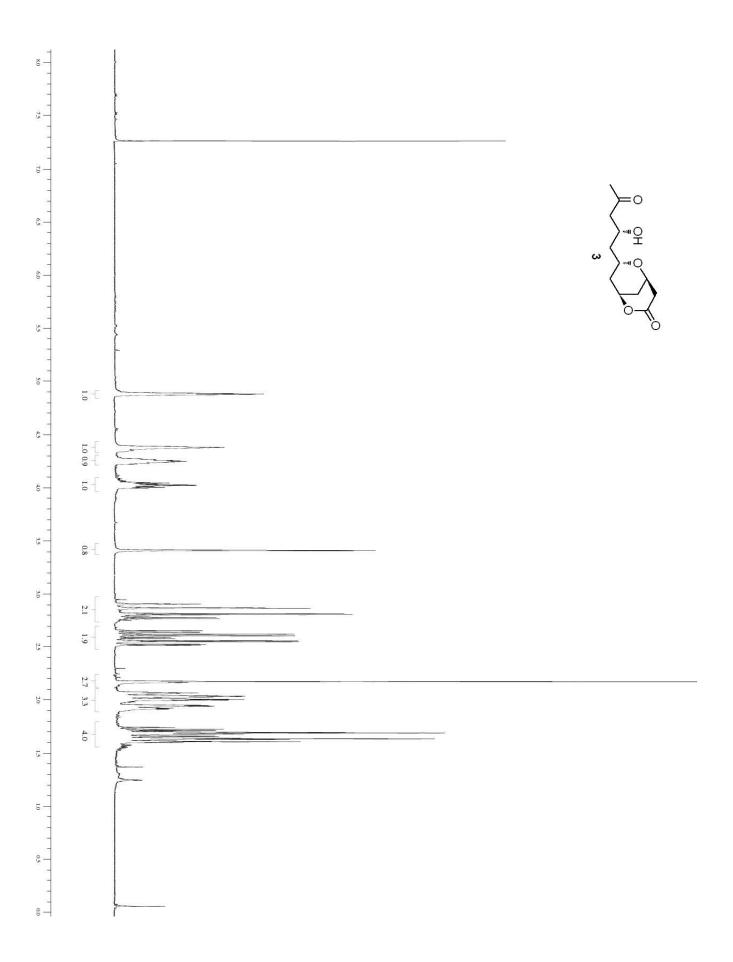


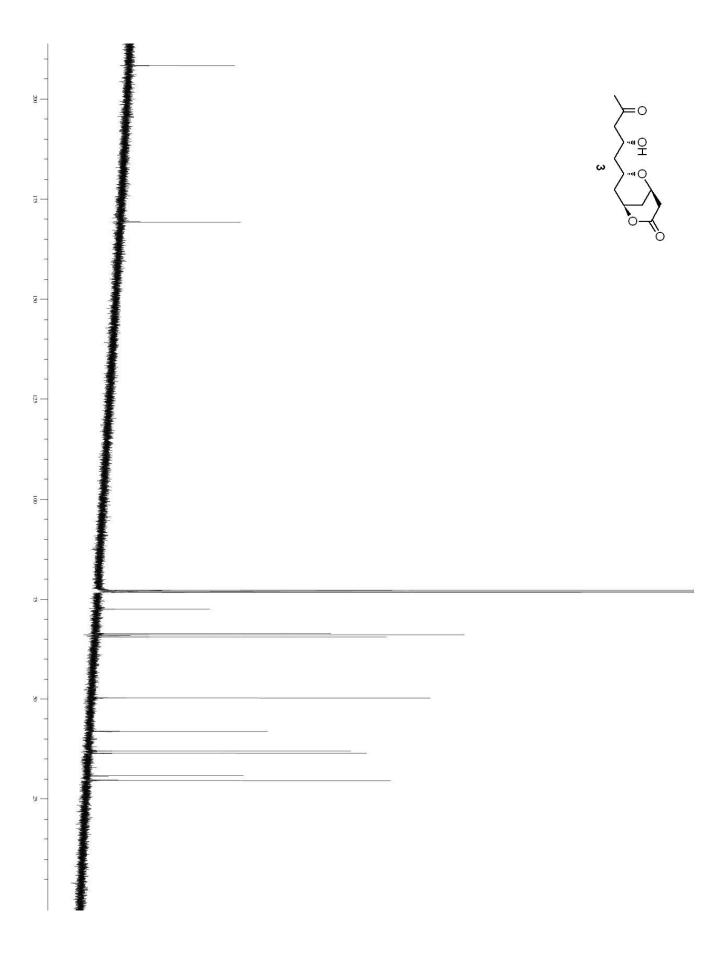


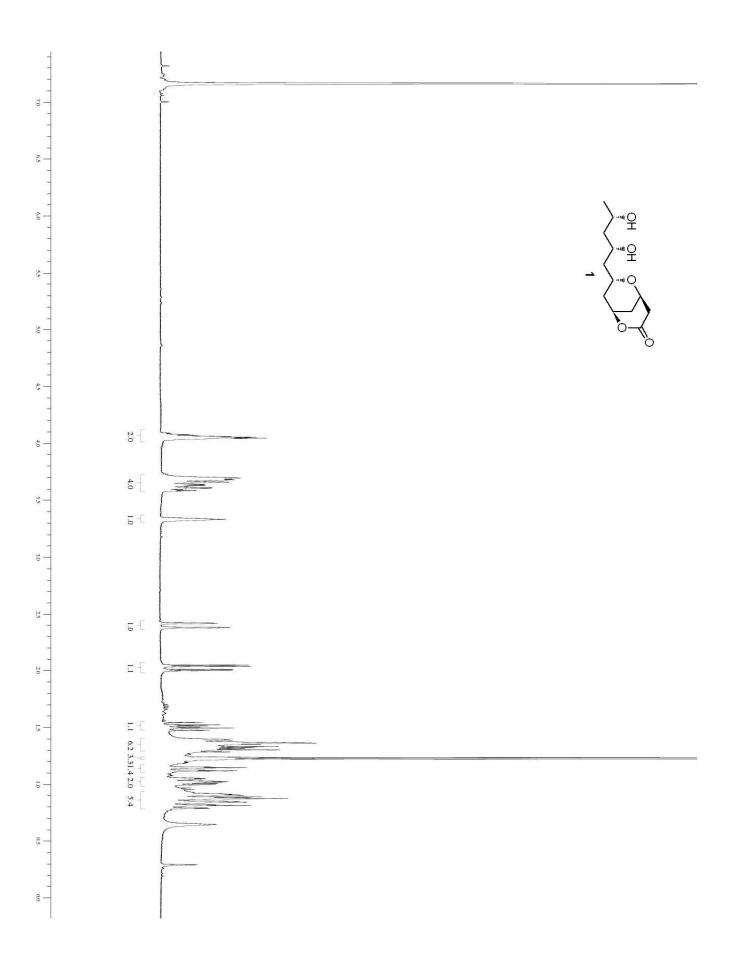


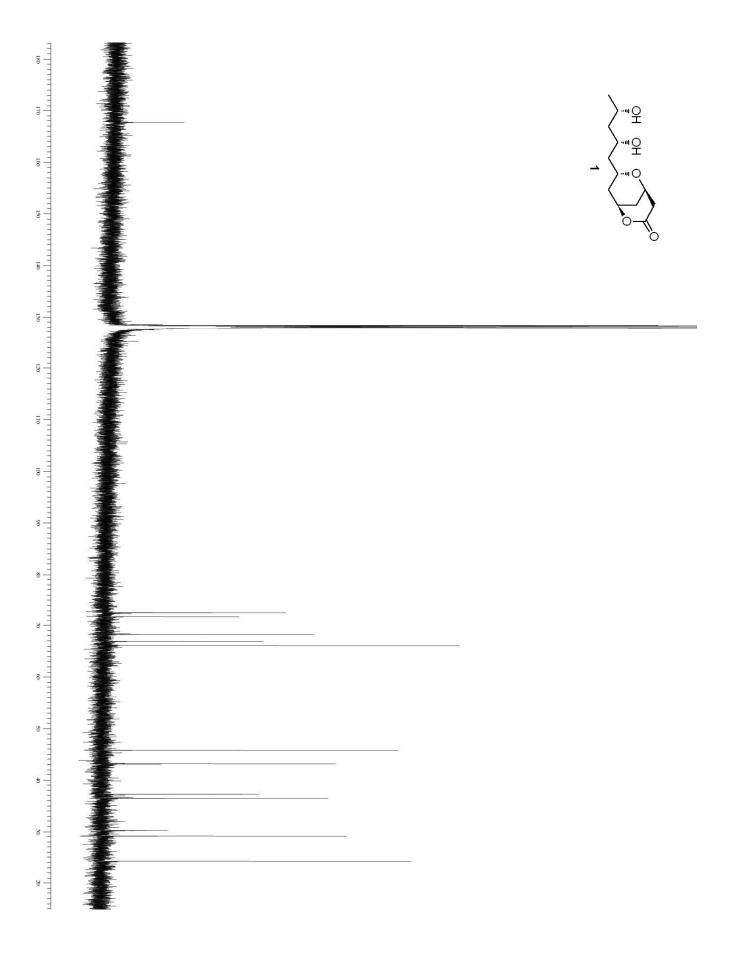


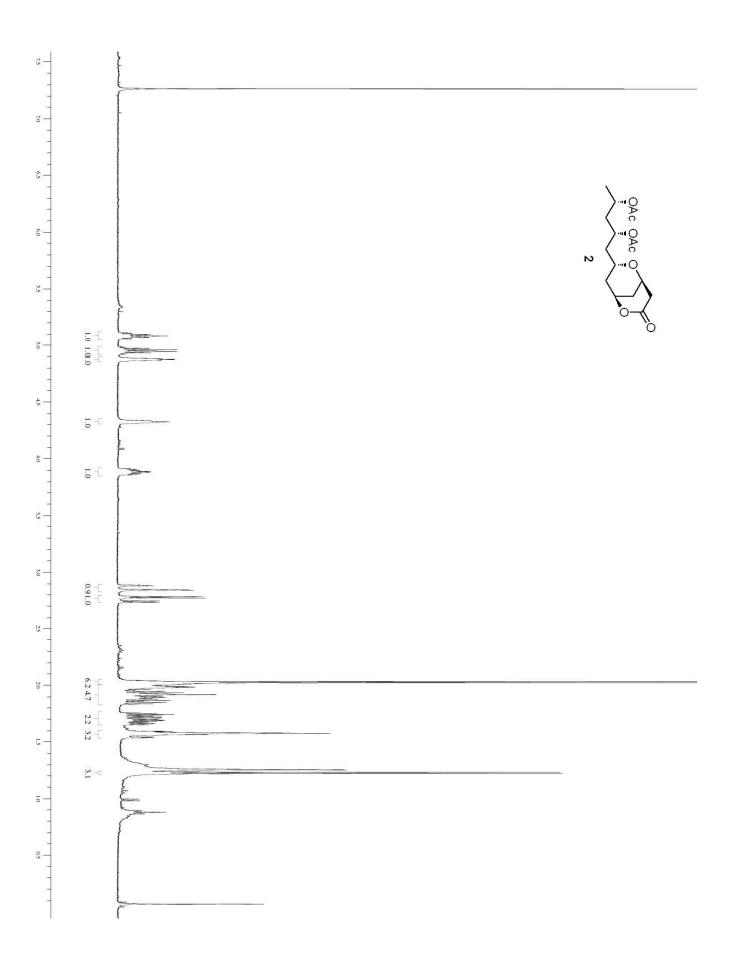


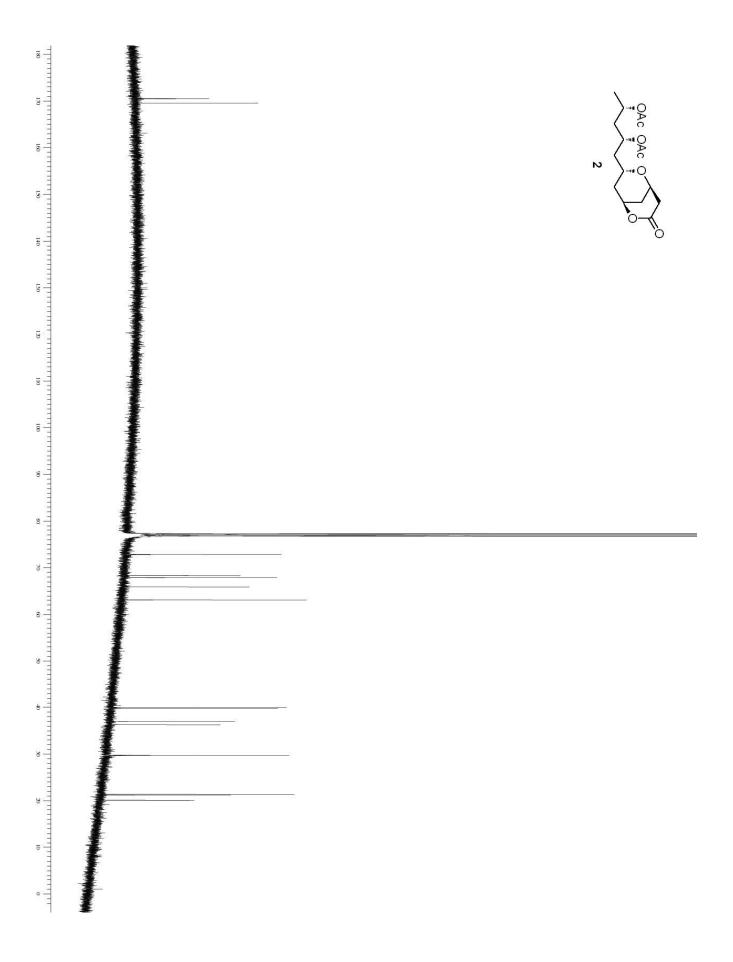












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