

## Supporting Information

### Stereoselective Formation of Chiral *trans*-4-Hydroxy-5-Substituted 2-Pyrrolidinones: Syntheses of Streptopyrrolidine and 3-*epi*-Epohelmin A

Chang-Mei Si,<sup>[a]</sup> Zhuo-Ya Mao,<sup>[a]</sup> Yi-Wen Liu,<sup>[a]</sup> Zhen-Ting Du,<sup>[c]</sup> Bang-Guo Wei\*,<sup>[a]</sup> and Guo-Qiang Lin<sup>[b]</sup>

<sup>[a]</sup>Department of Natural Products Chemistry, School of Pharmacy Fudan University, 826 Zhangheng Road, Shanghai 201203, China

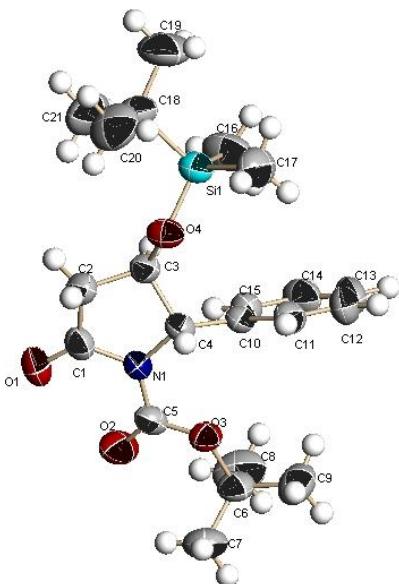
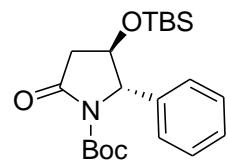
<sup>[b]</sup>Shanghai Institute of Organic Chemistry, Chinese Academy of Sciences, 345 Lingling Road, Shanghai 200032, China

<sup>[c]</sup>College of Science, Northwest Agriculture and Forestry University, Shaanxi, Yangling, 712100, China

## Table of Contents

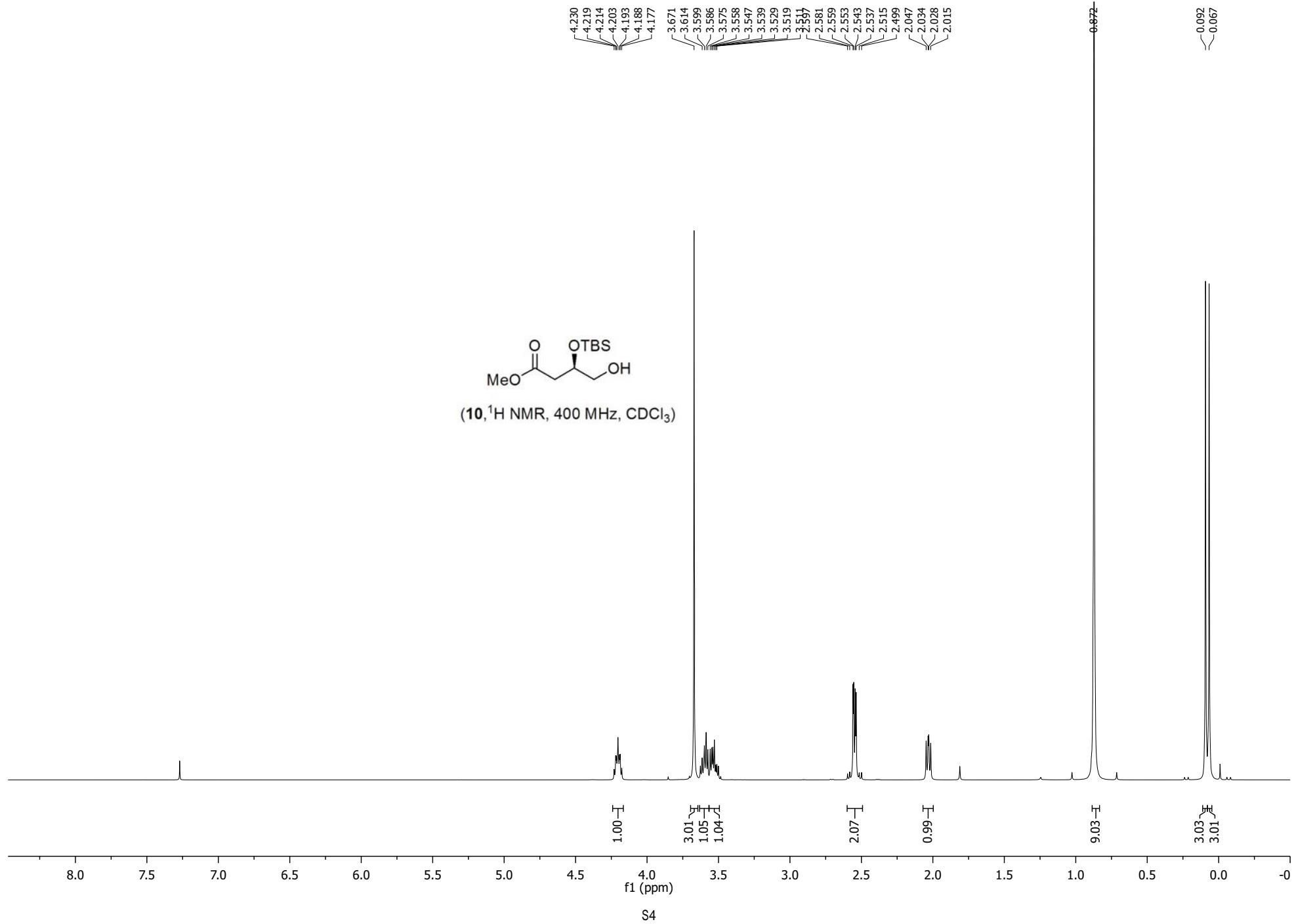
X-Ray Structures of <i>trans</i> - <b>4a</b> .....	S3
NMR Spectra of coupled products.....	S4-S108

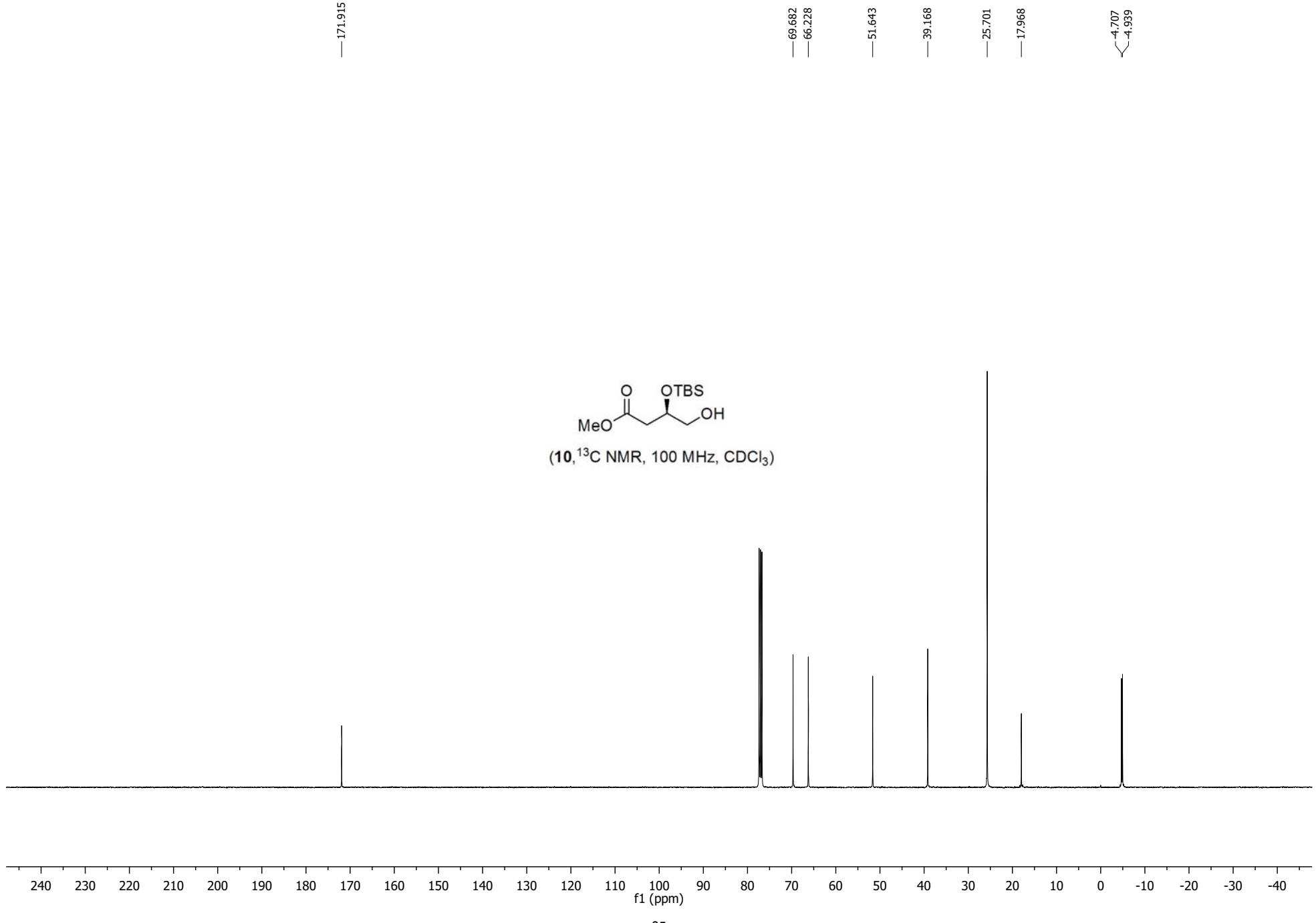
ORTEP drawing of the X-ray crystallographic structure of *trans*-**4a**

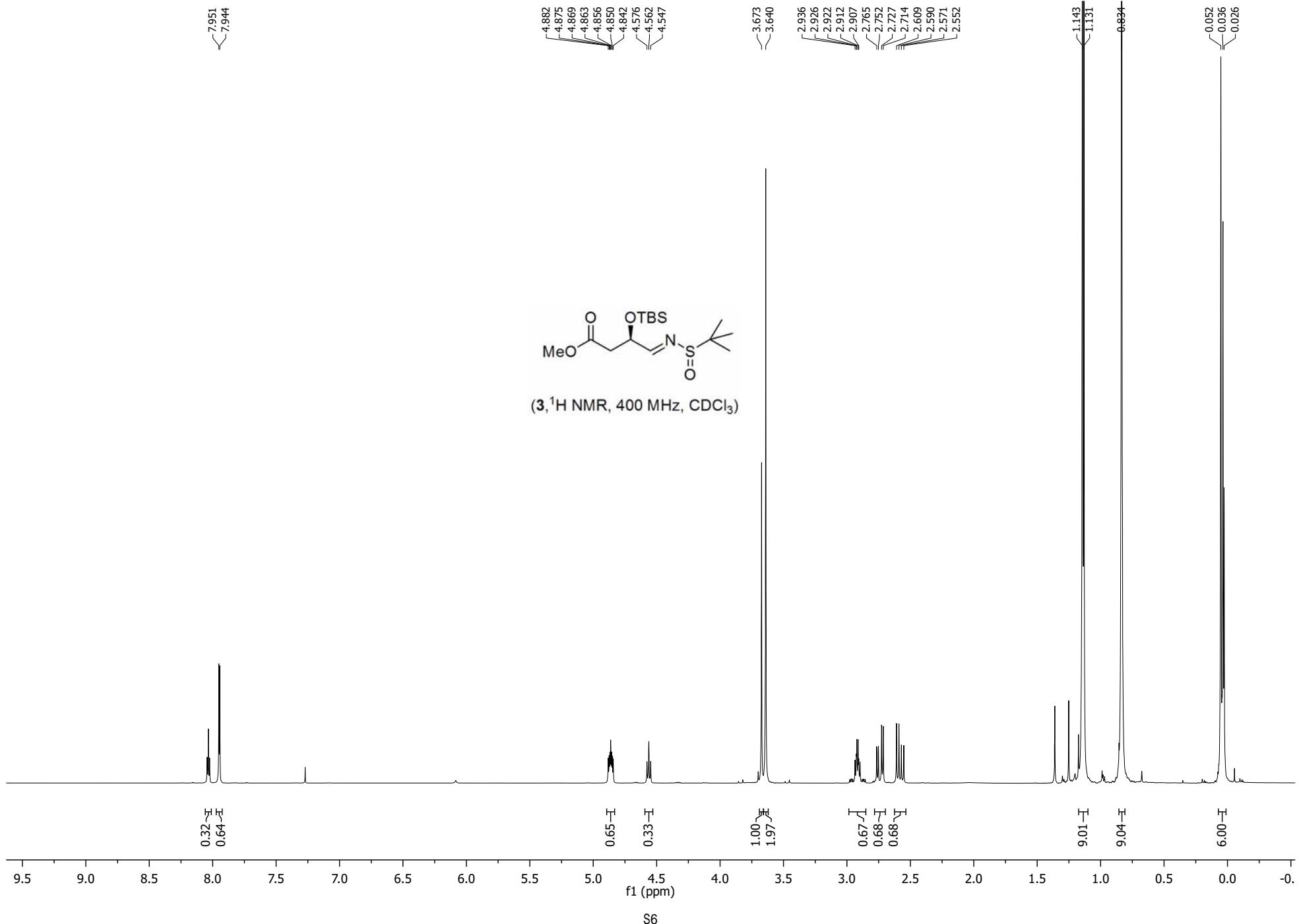


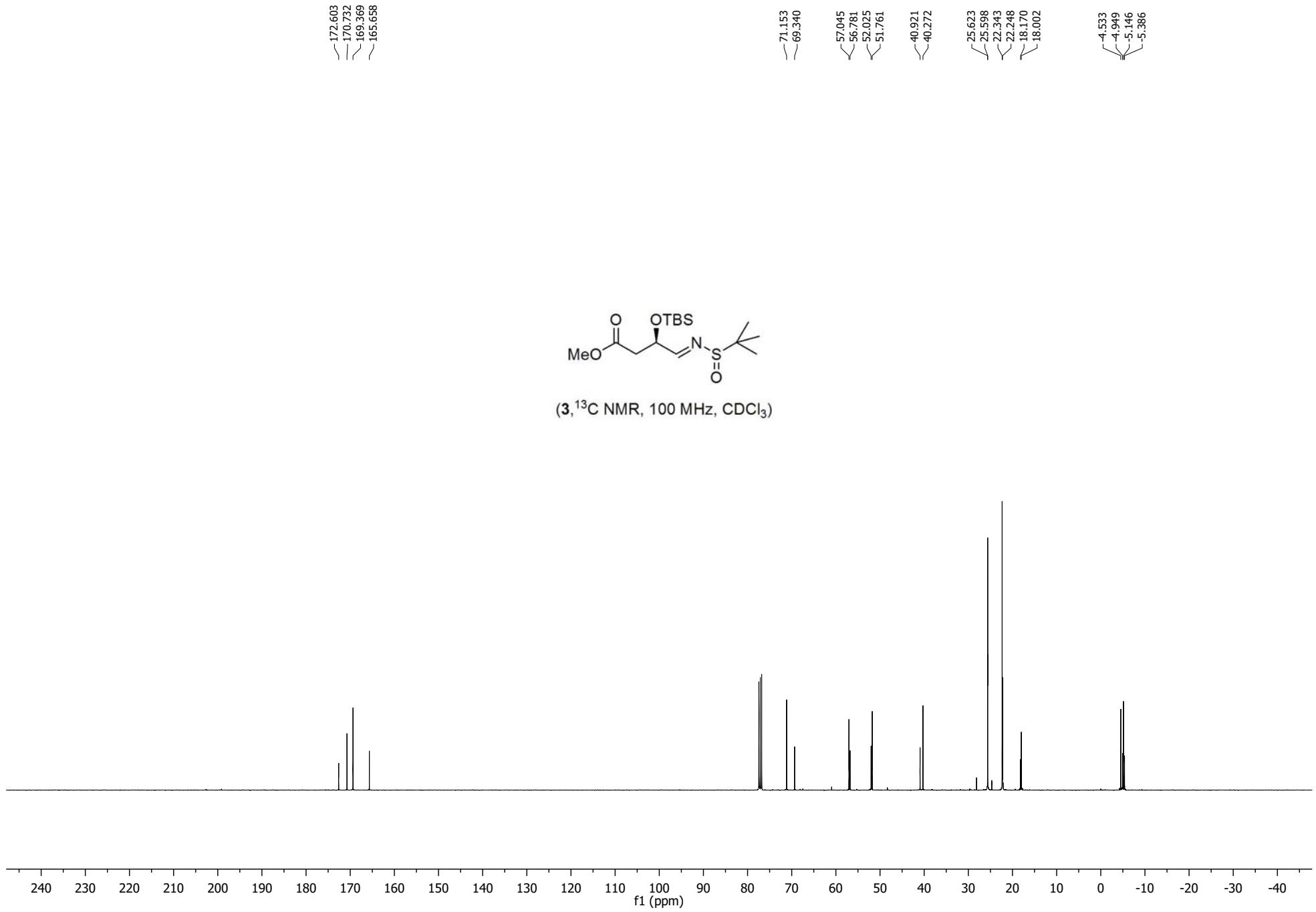
CCDC-1417164. For detailed crystallographic data, please refer to the Cambridge Crystallographic Data Centre at:

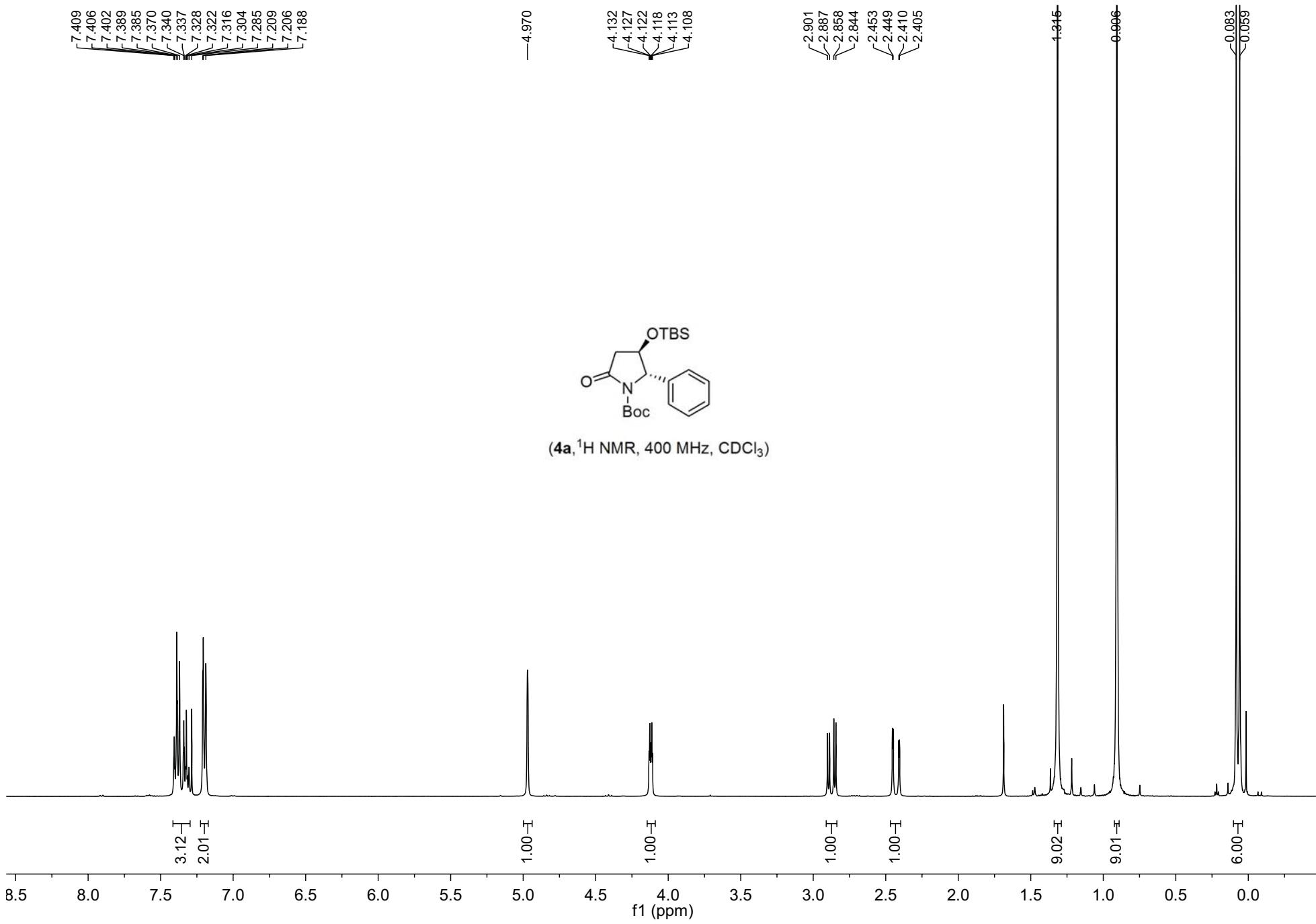
[http://www.ccdc.cam.ac.uk/data\\_request/cif](http://www.ccdc.cam.ac.uk/data_request/cif).

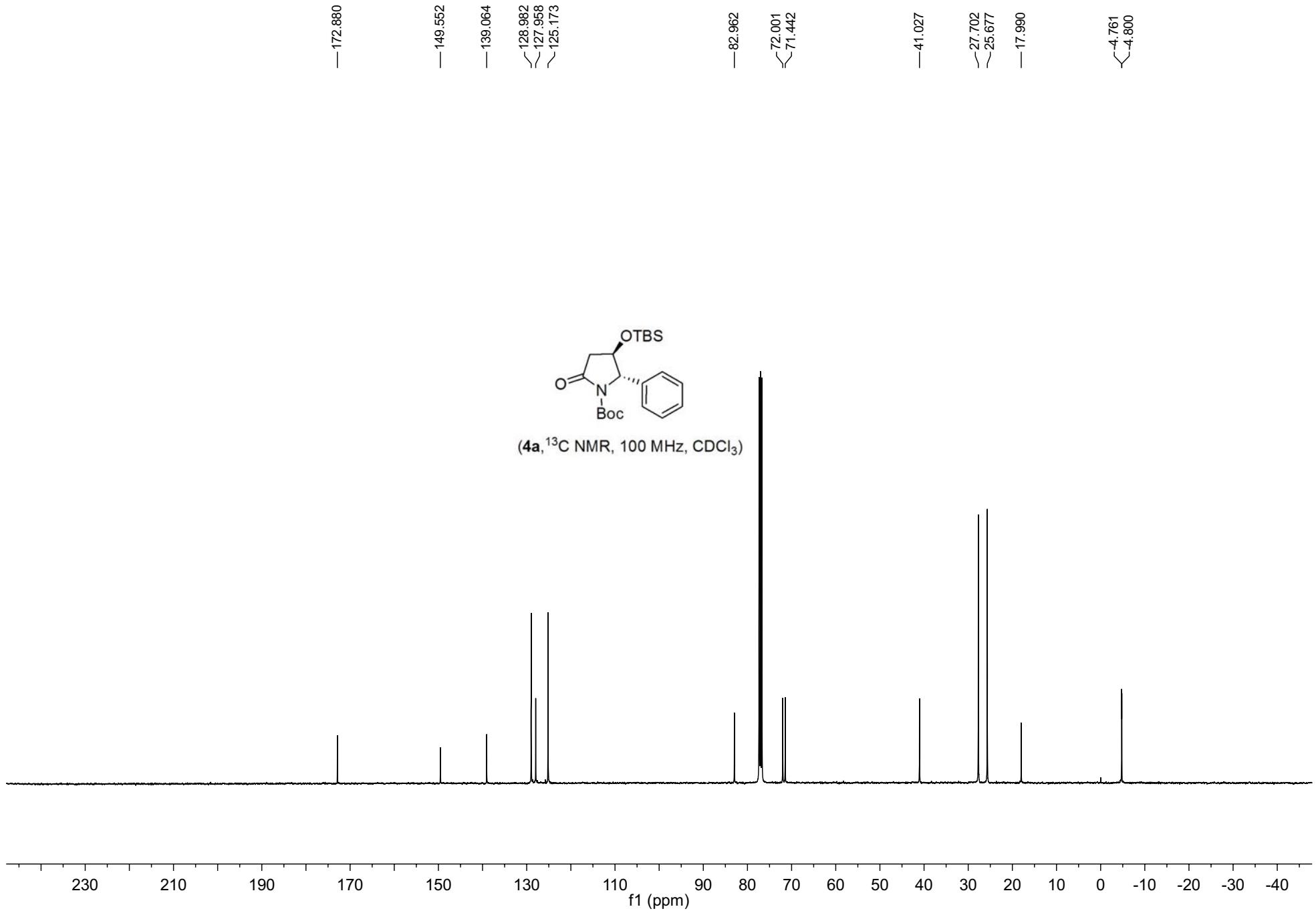


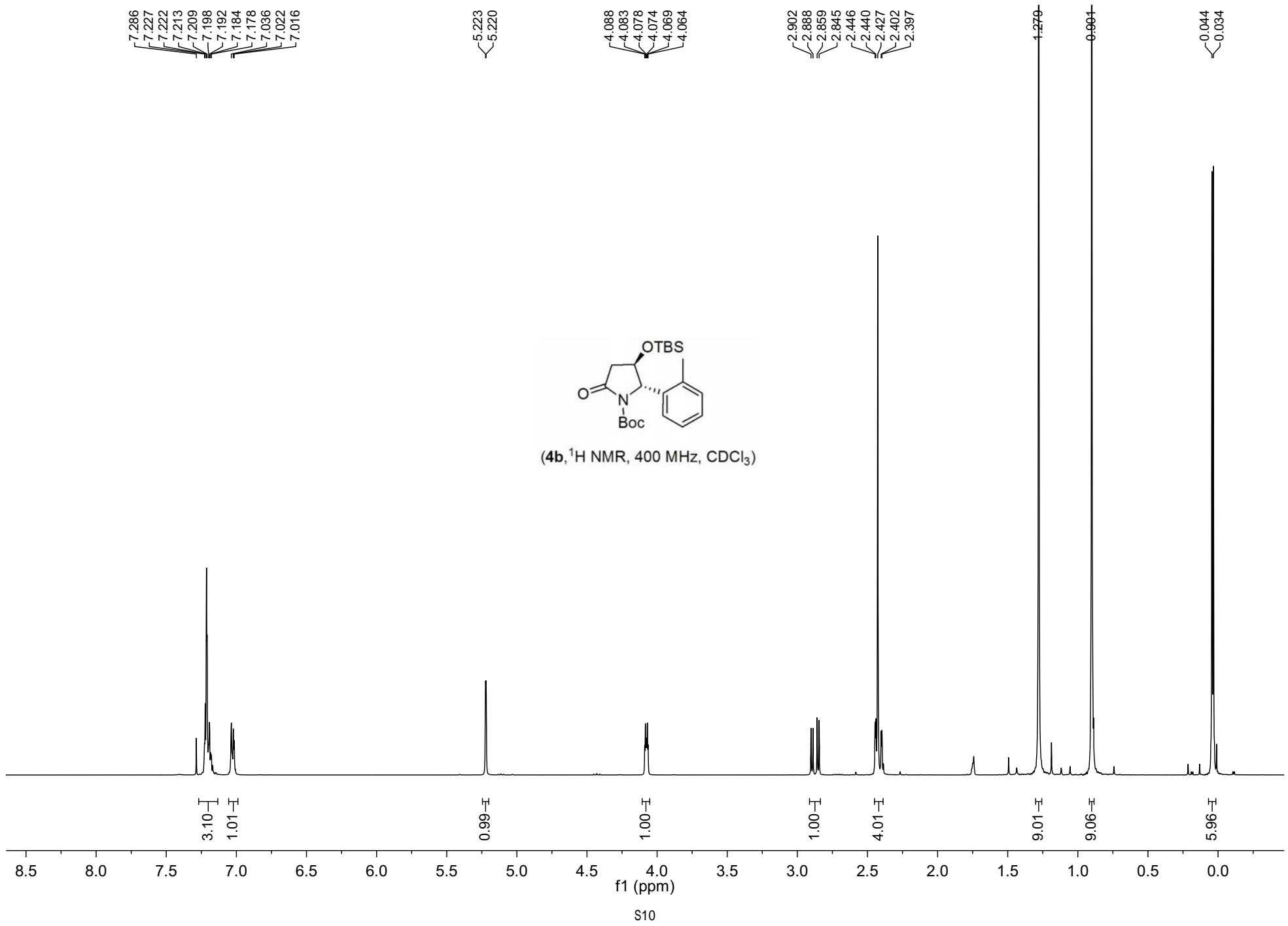


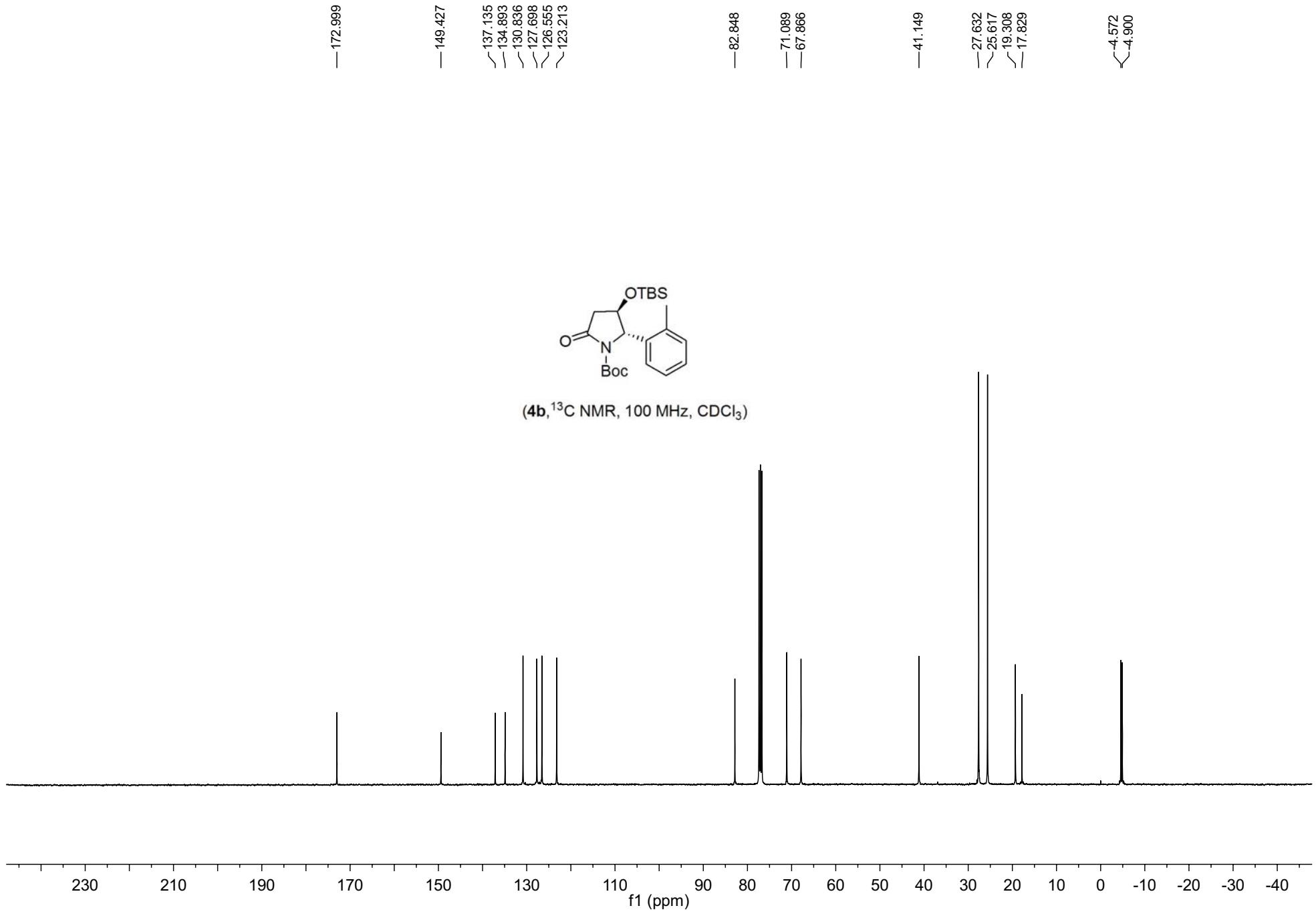


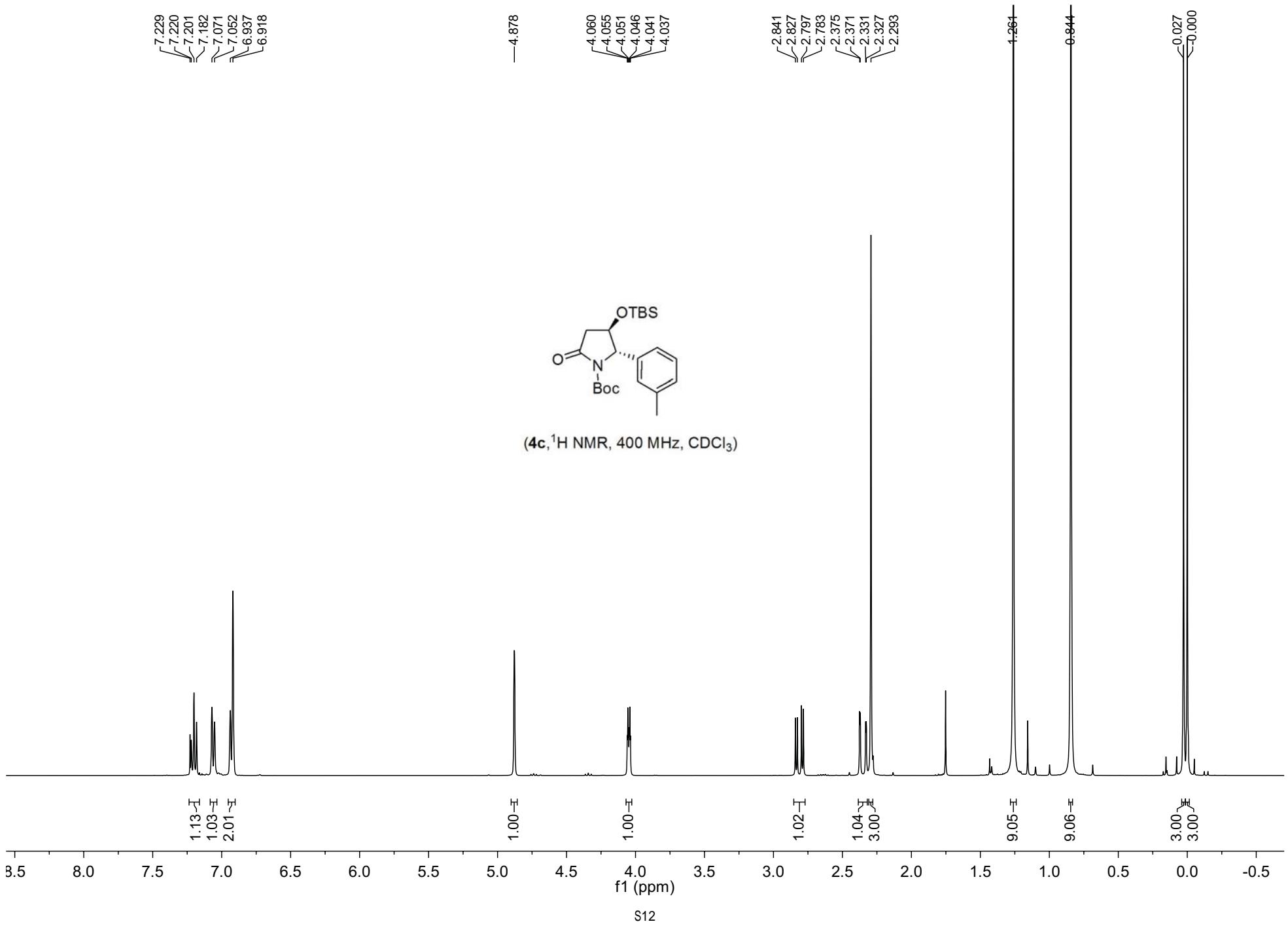


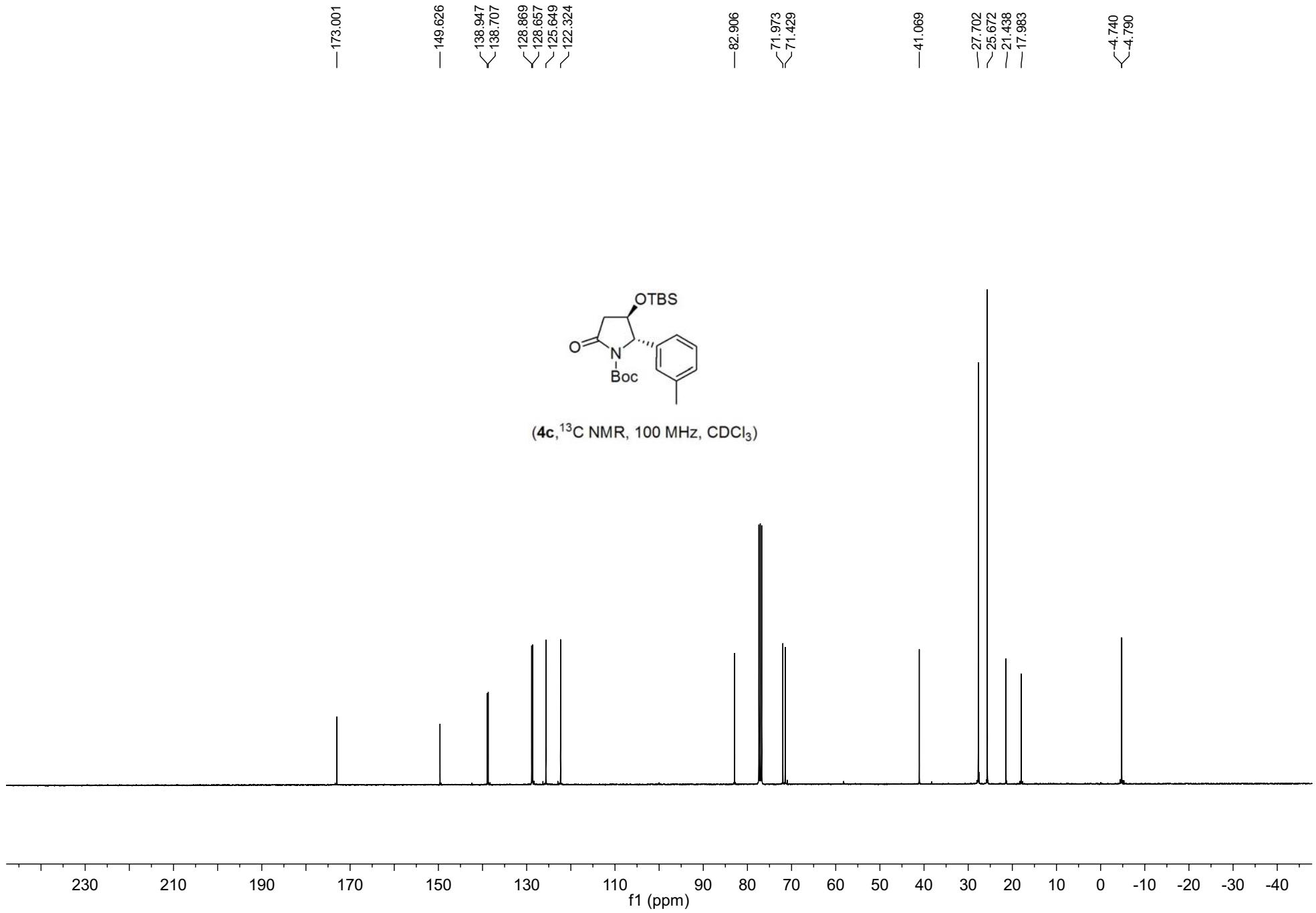


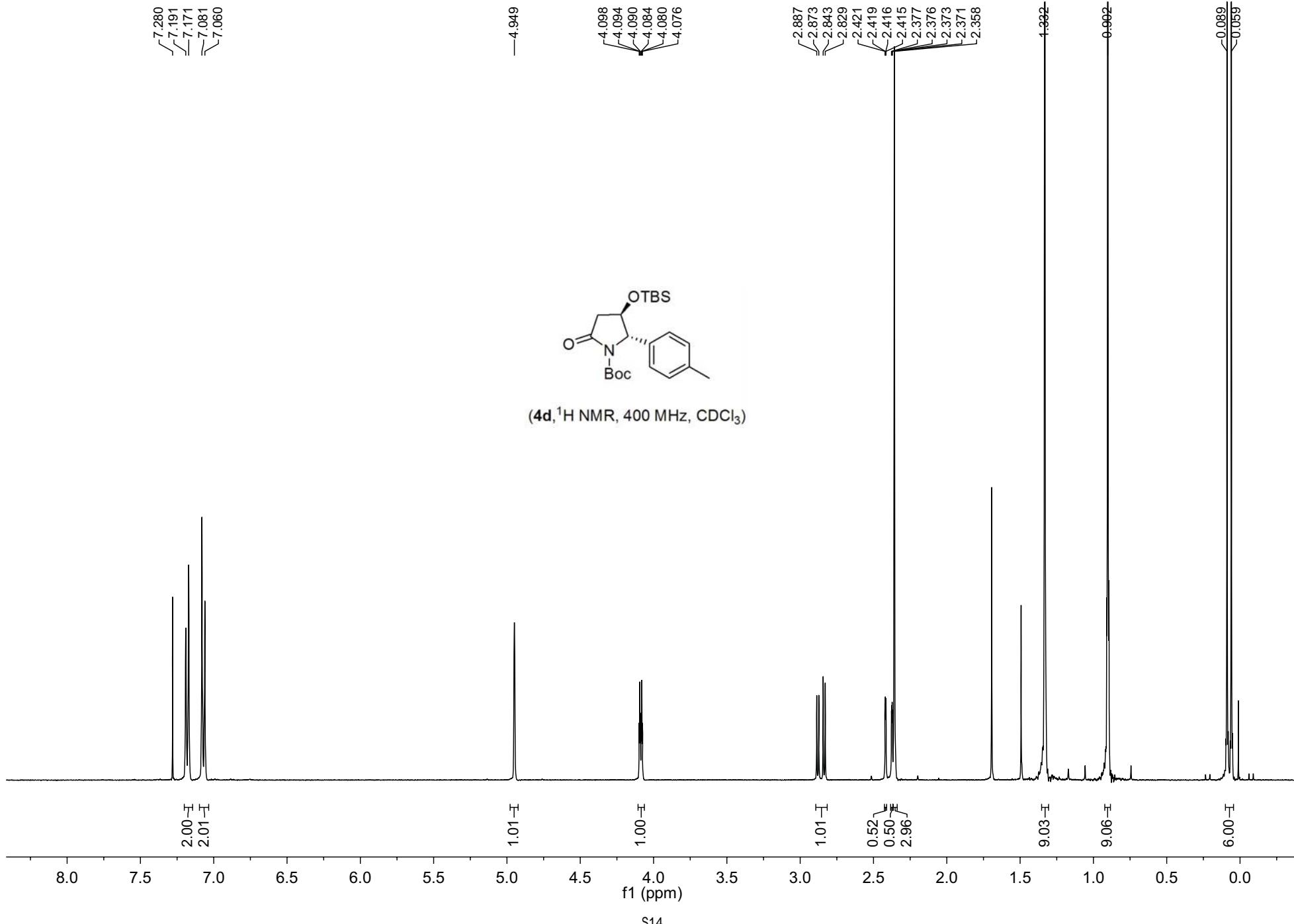


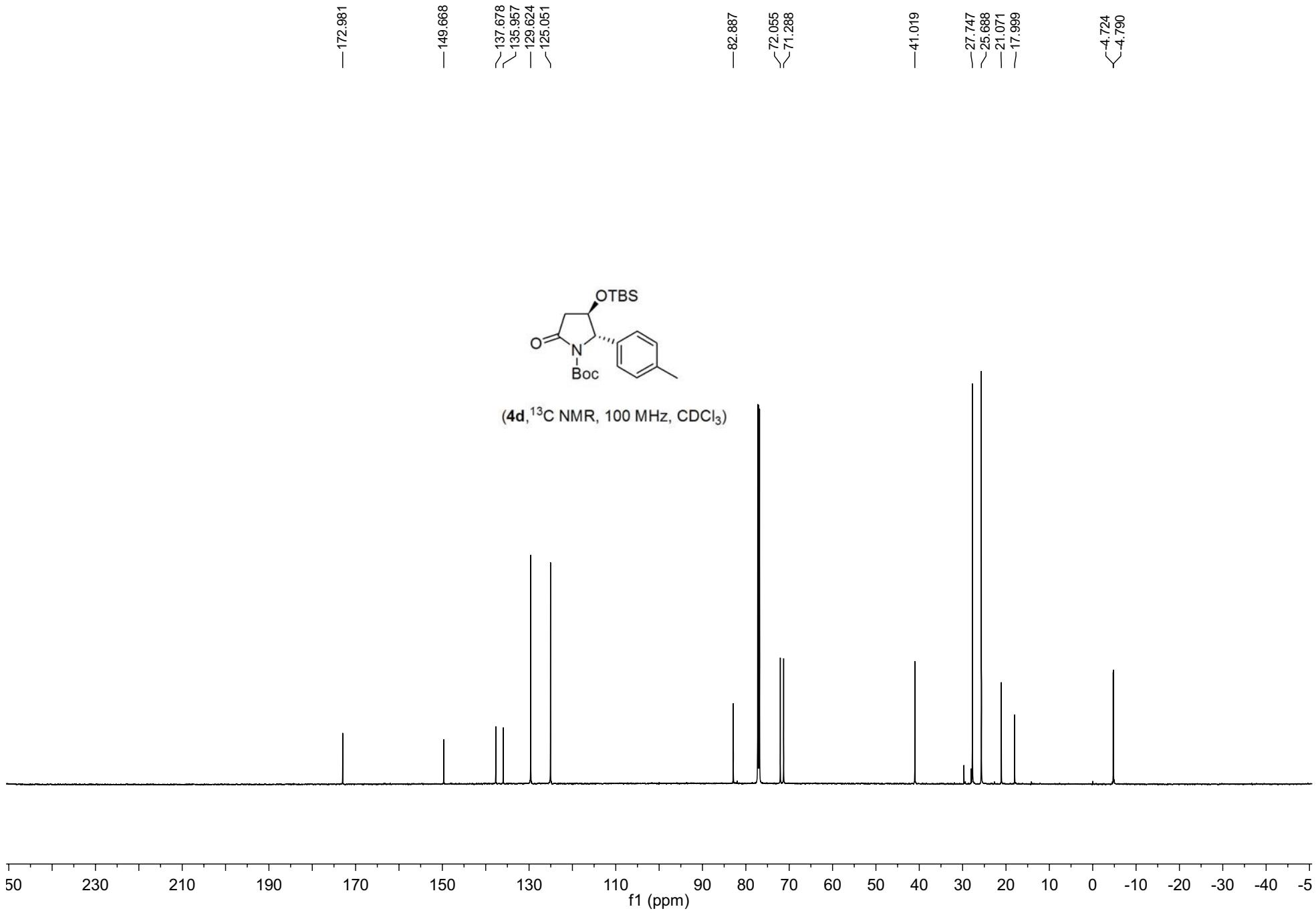


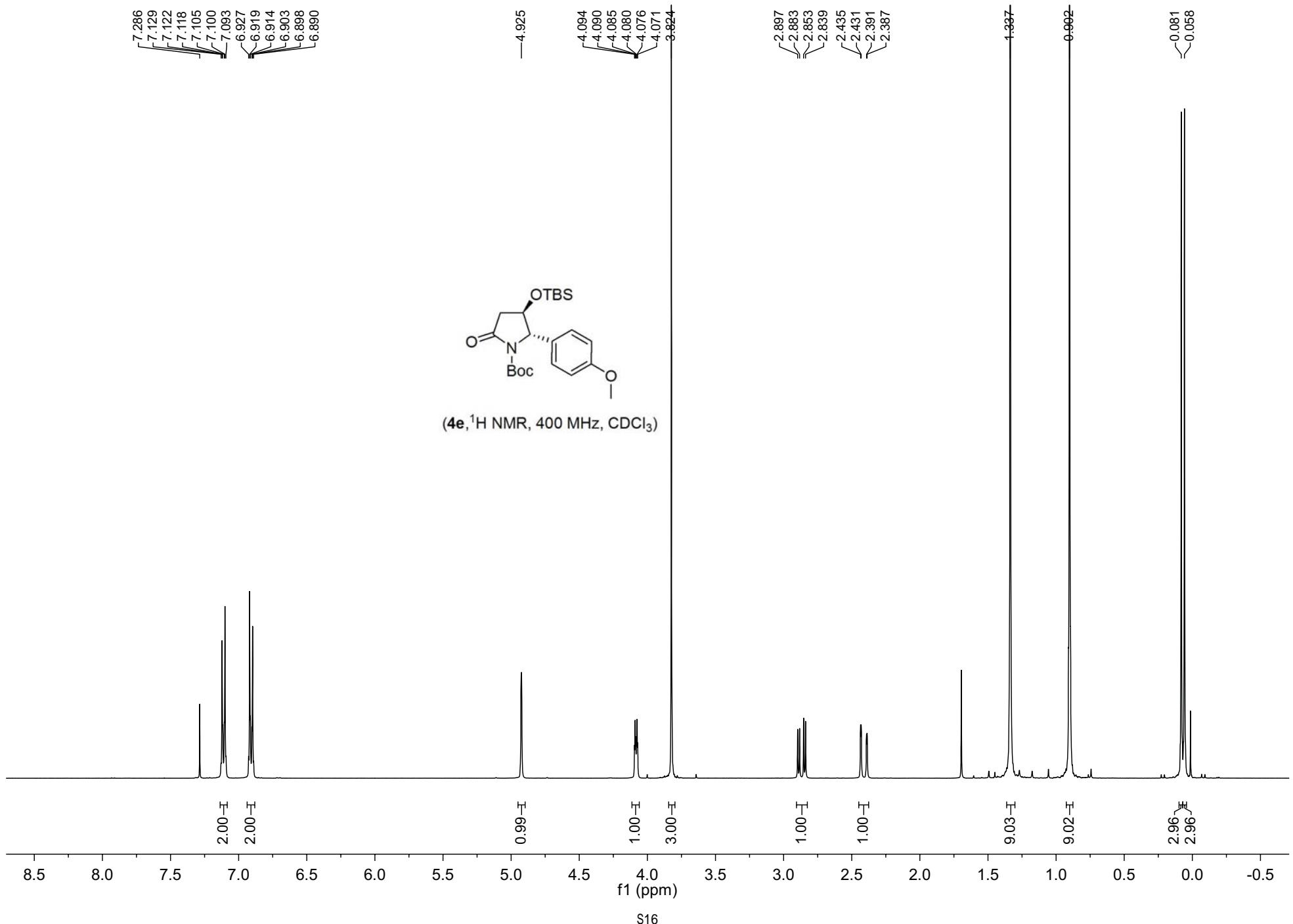


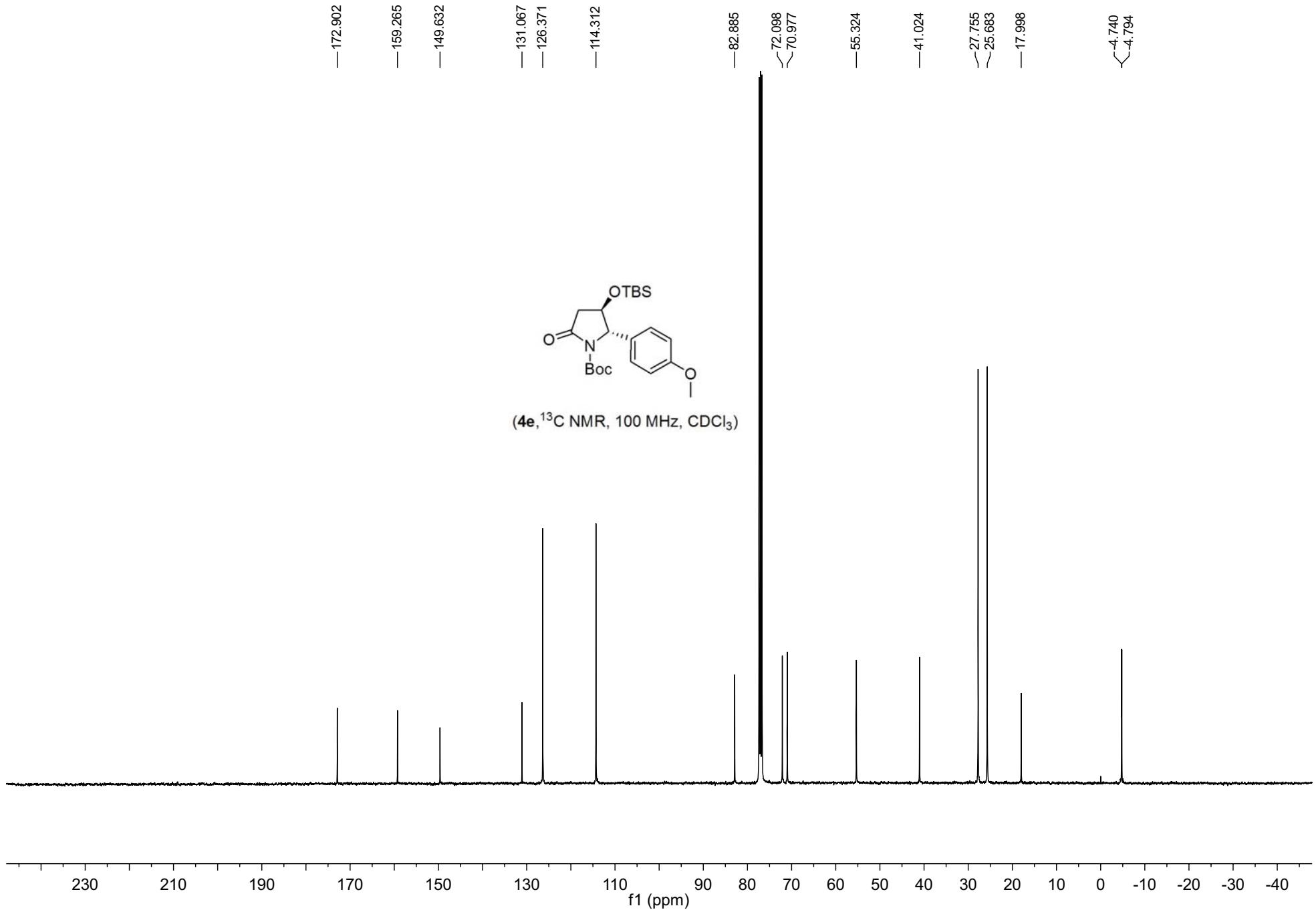


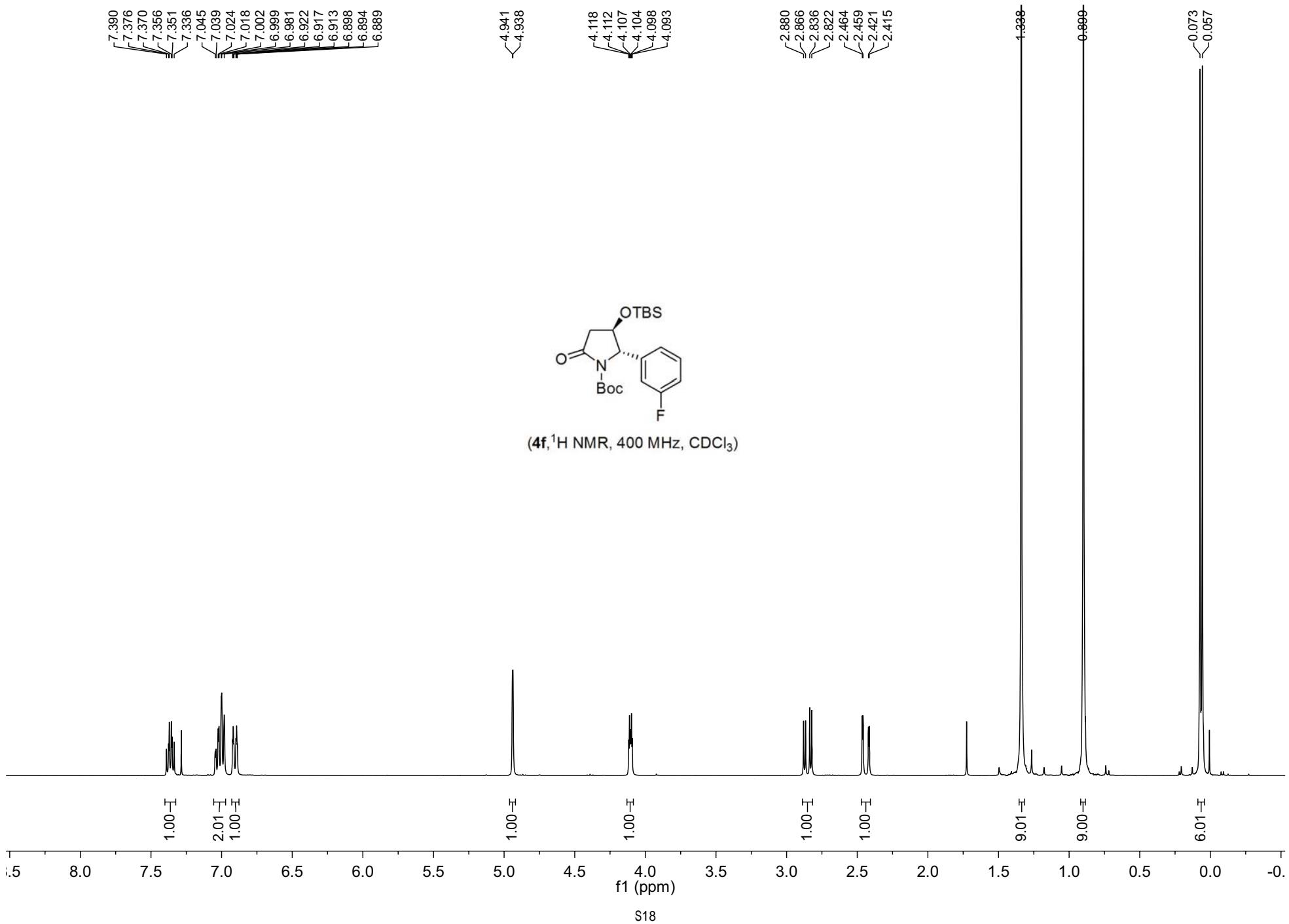


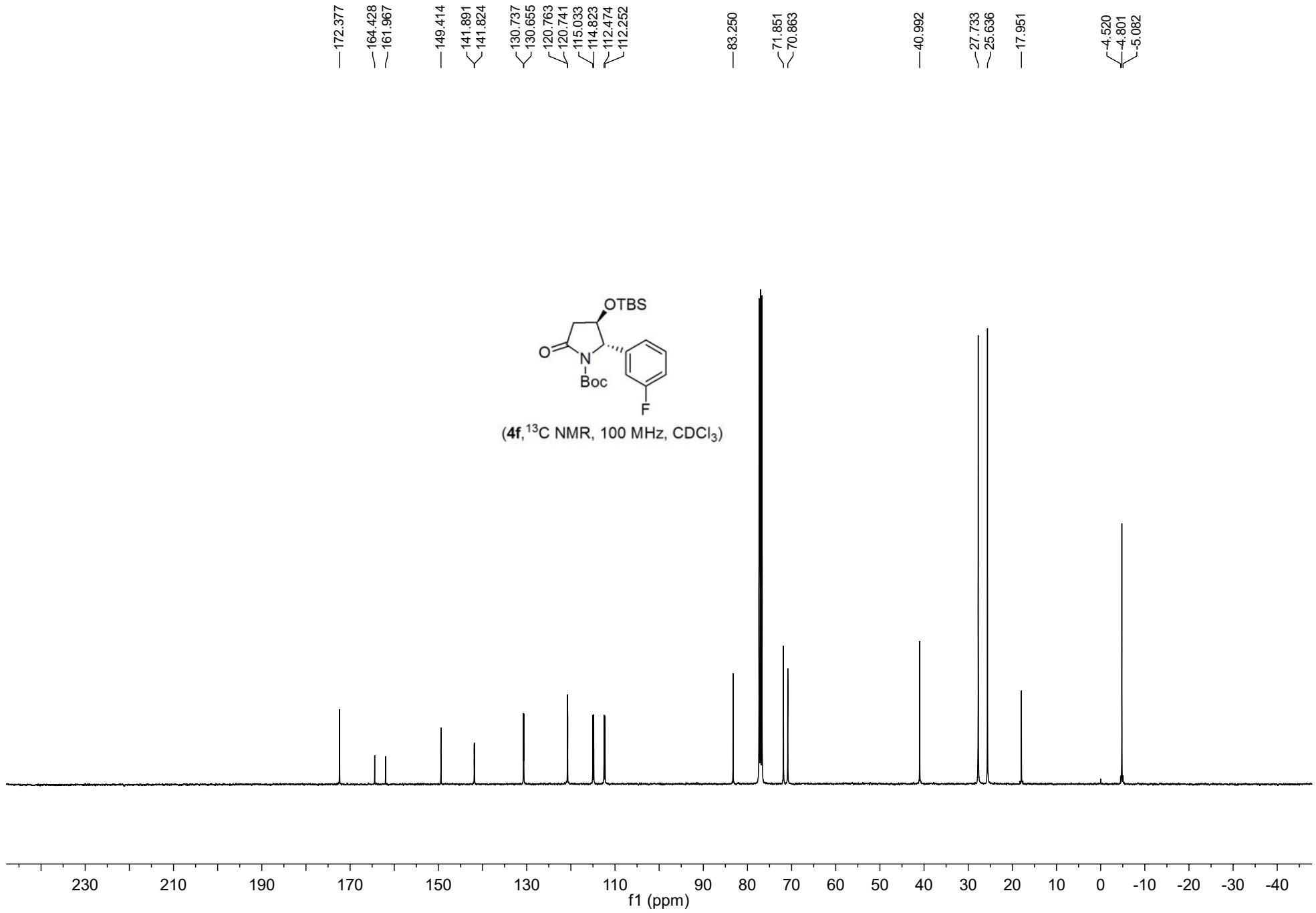




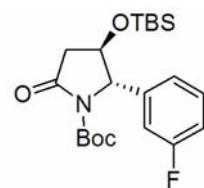




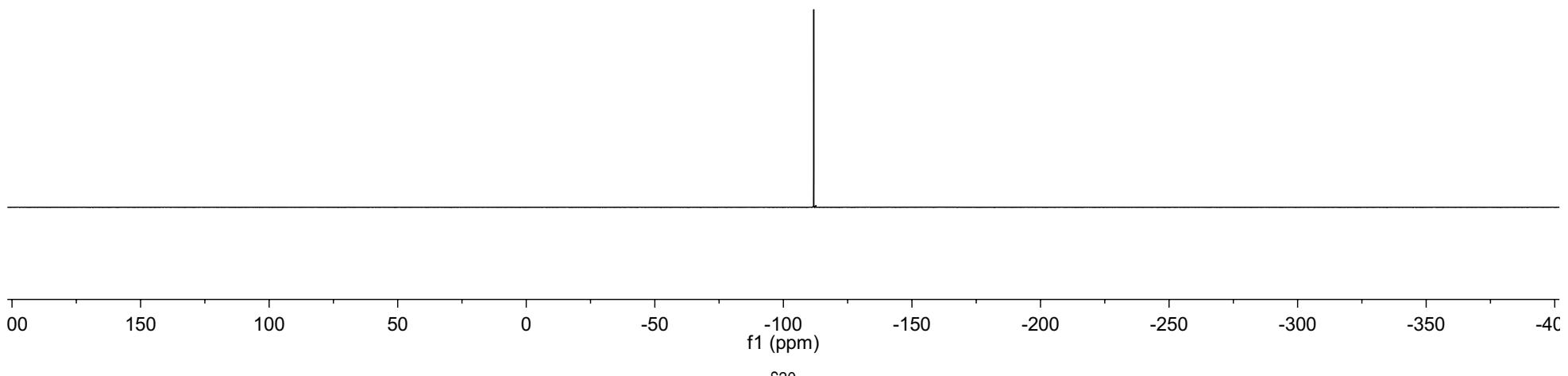


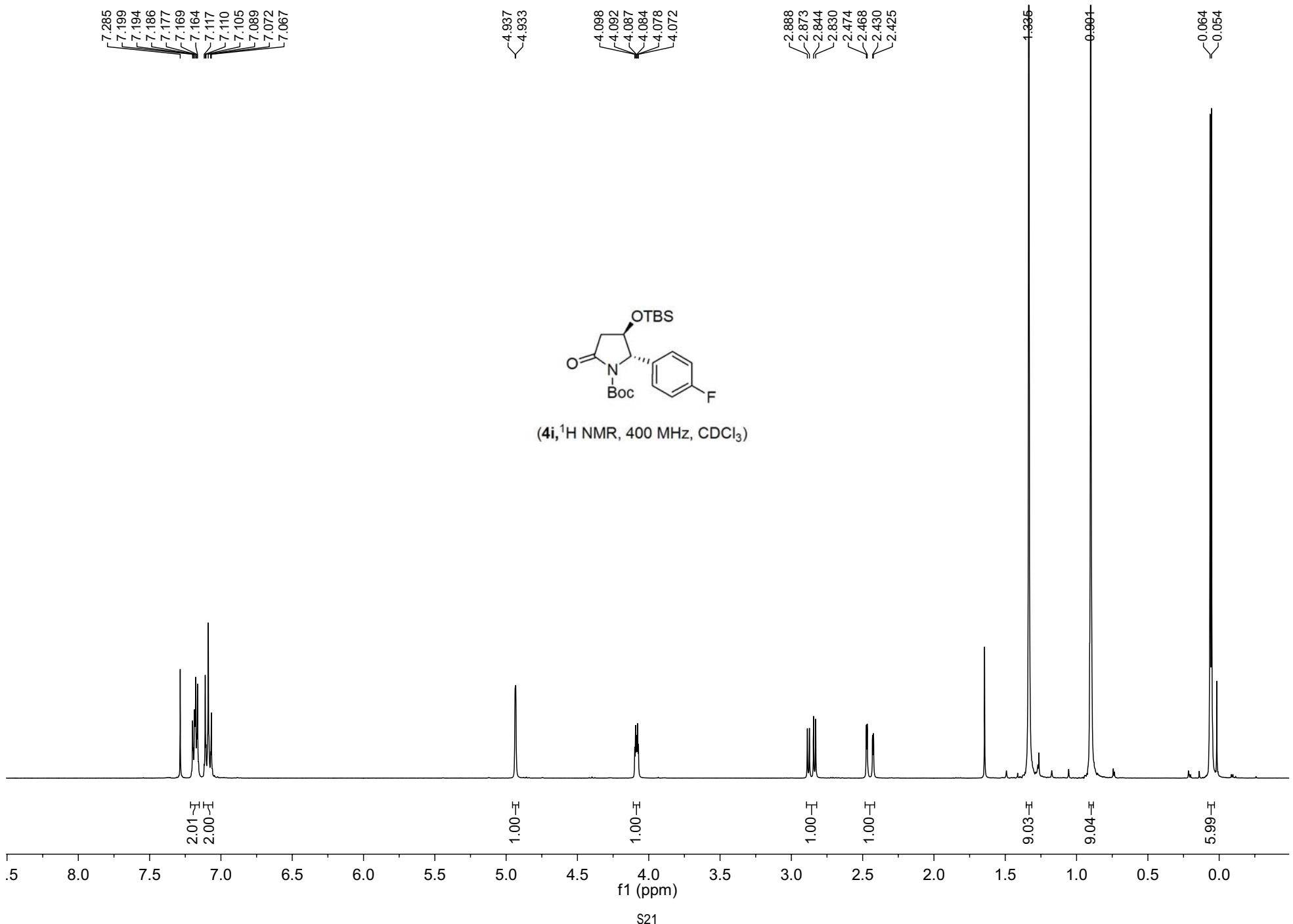


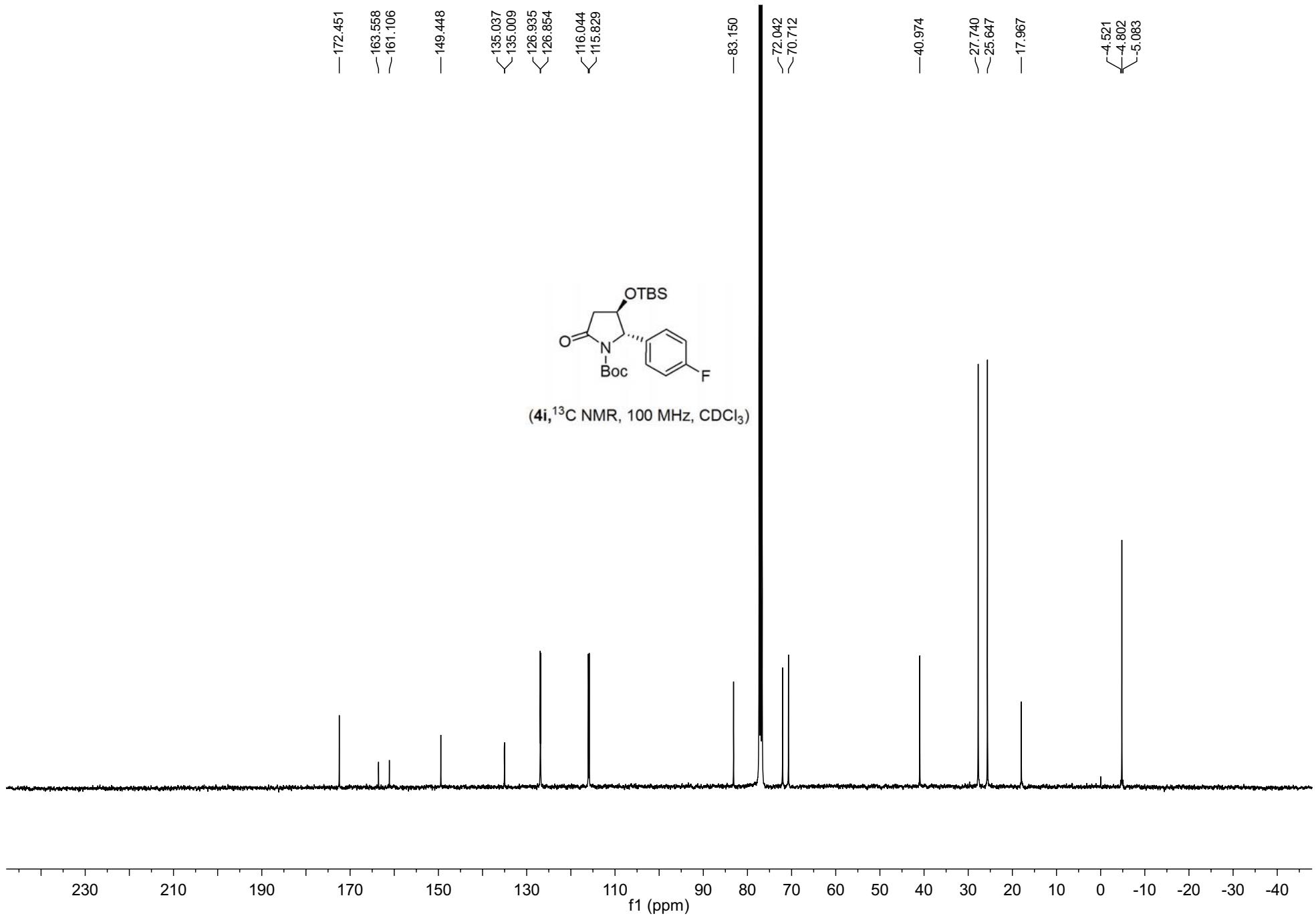
— -111.815



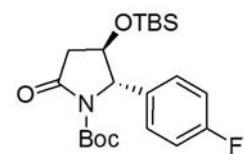
(**4f**,  $^{19}\text{F}$  NMR, 376 MHz,  $\text{CDCl}_3$ )



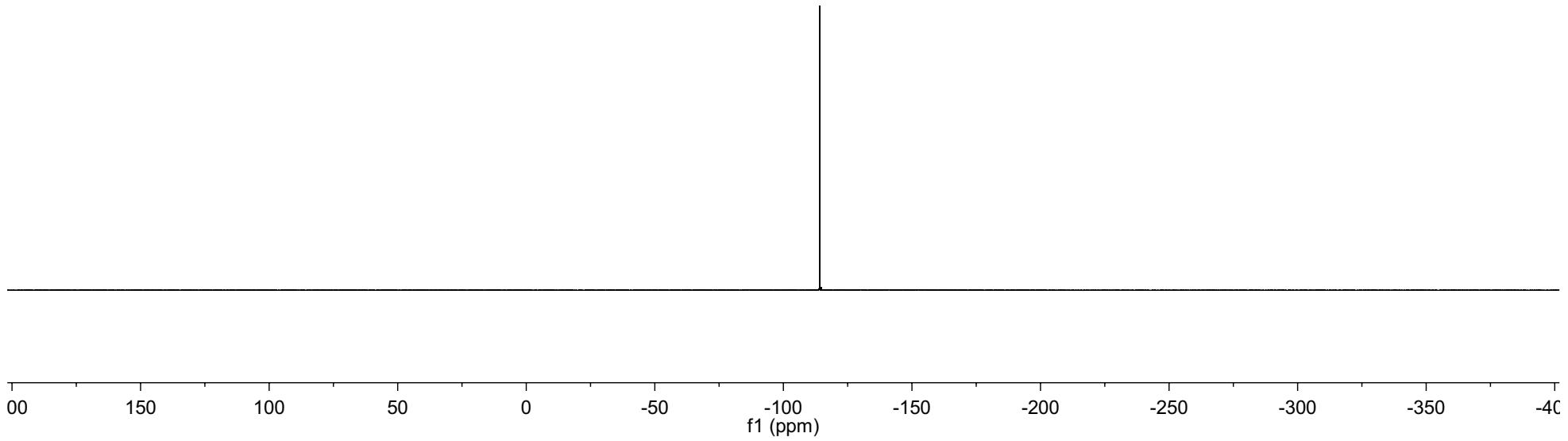


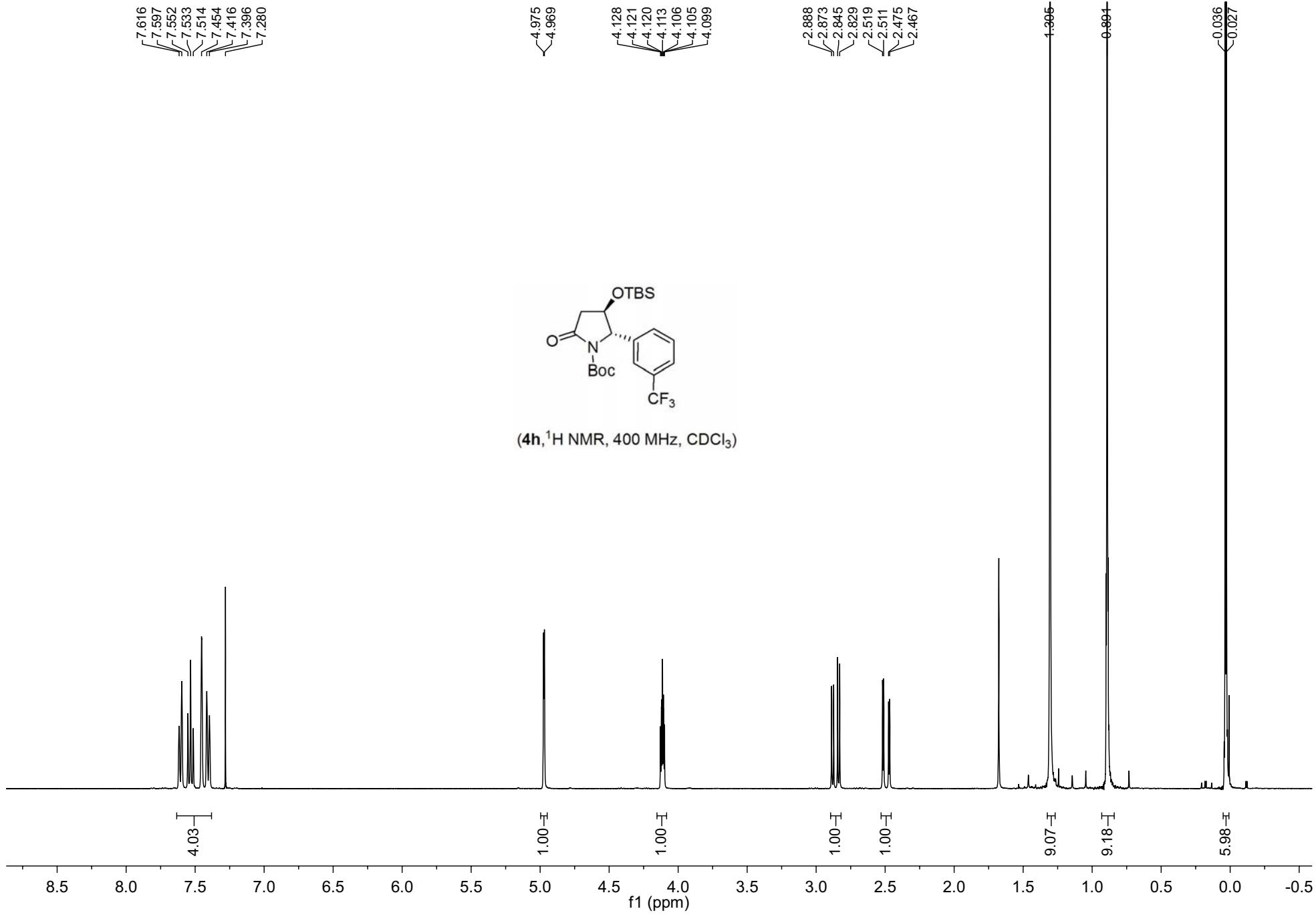


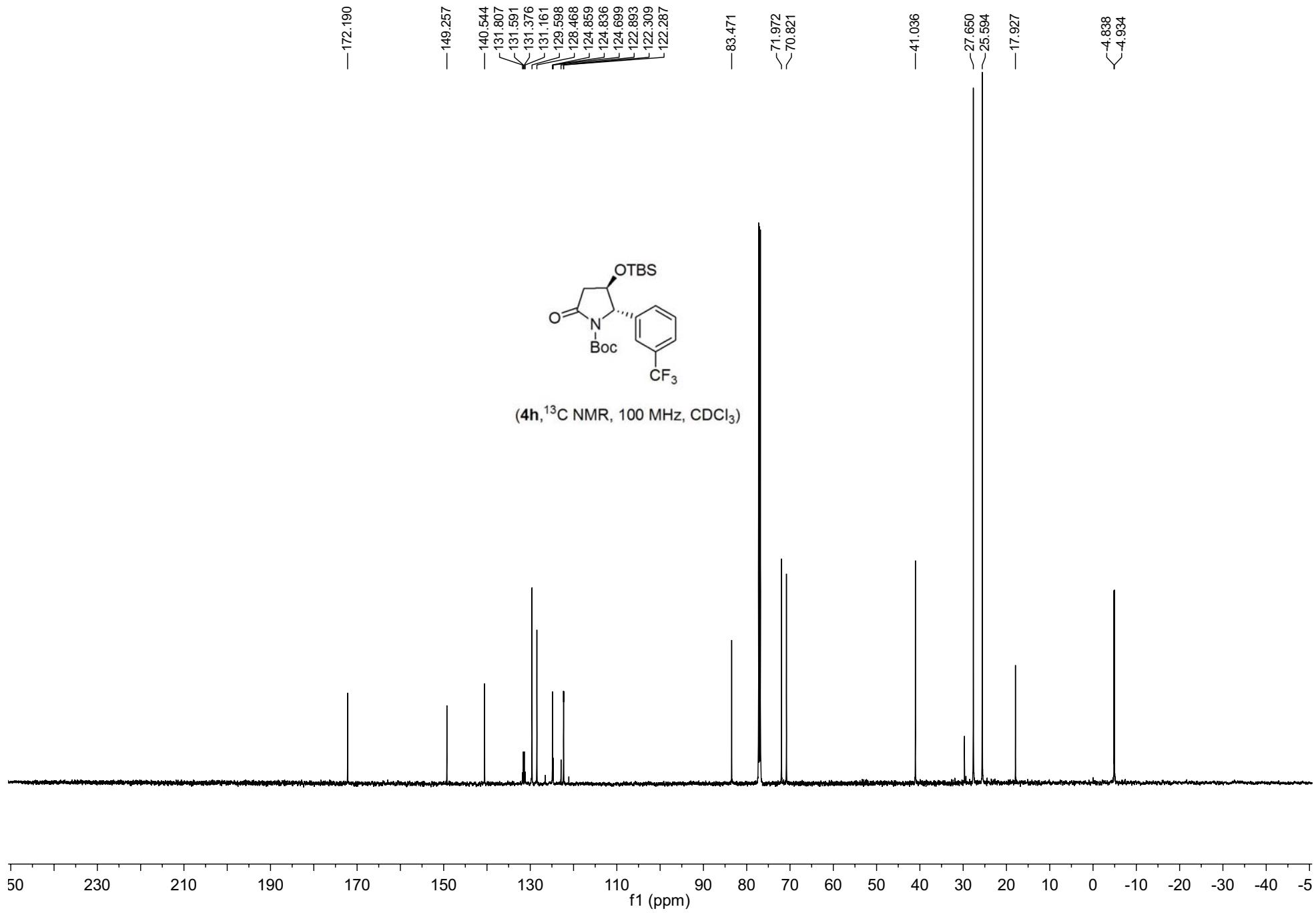
-114.105



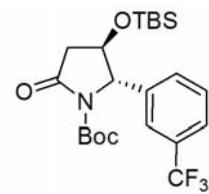
(**4i**,  $^{19}\text{F}$  NMR, 376 MHz,  $\text{CDCl}_3$ )



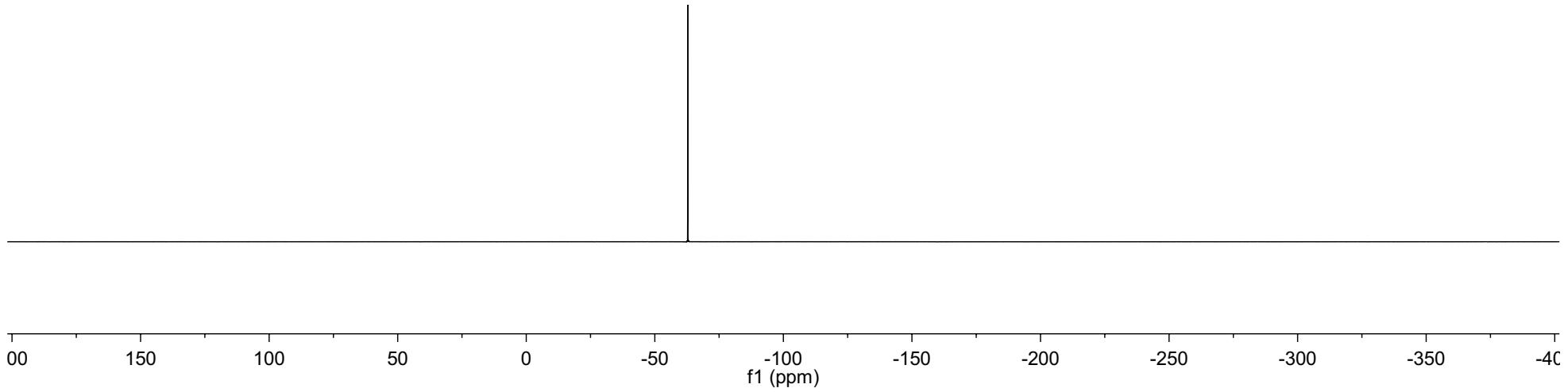


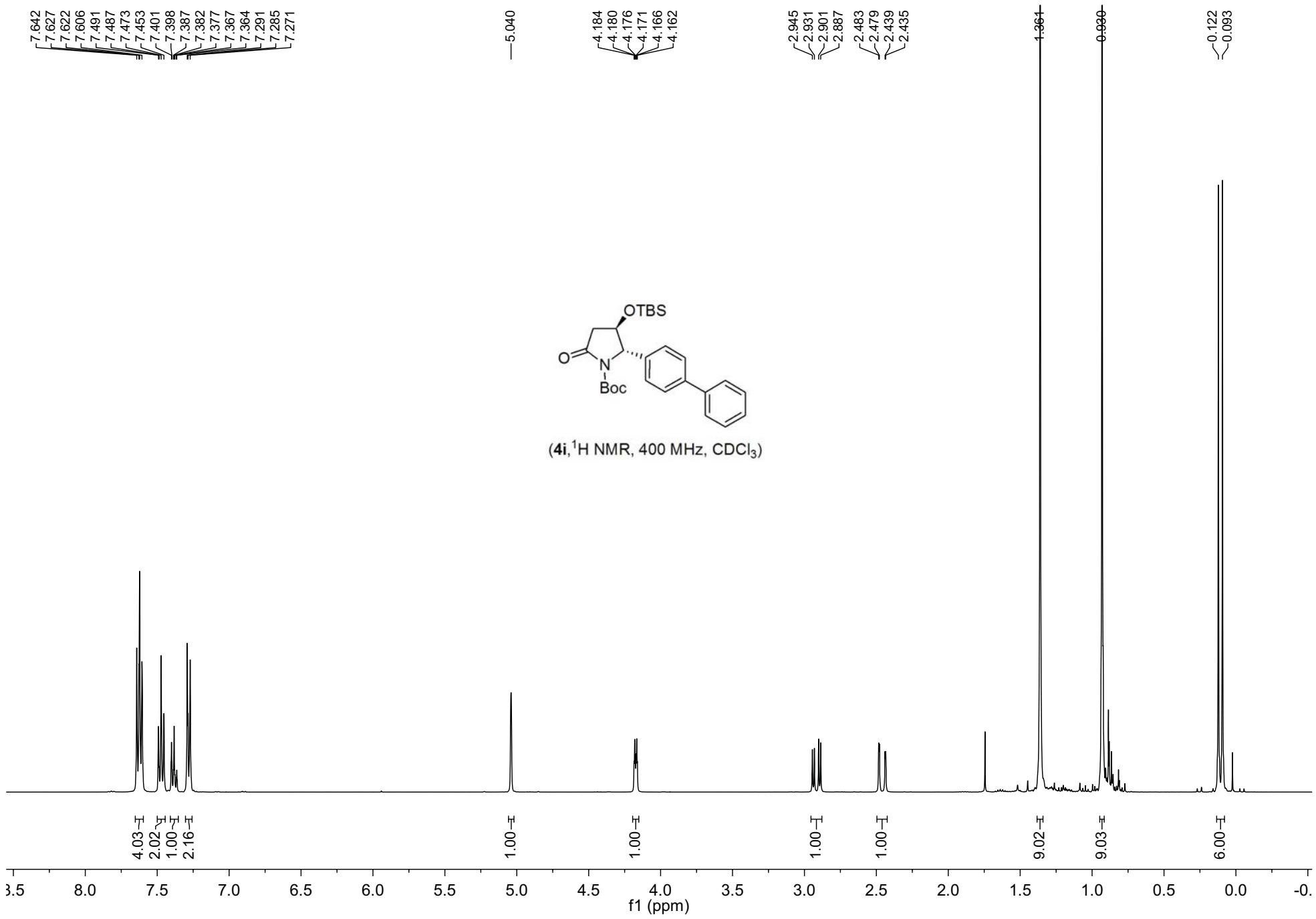


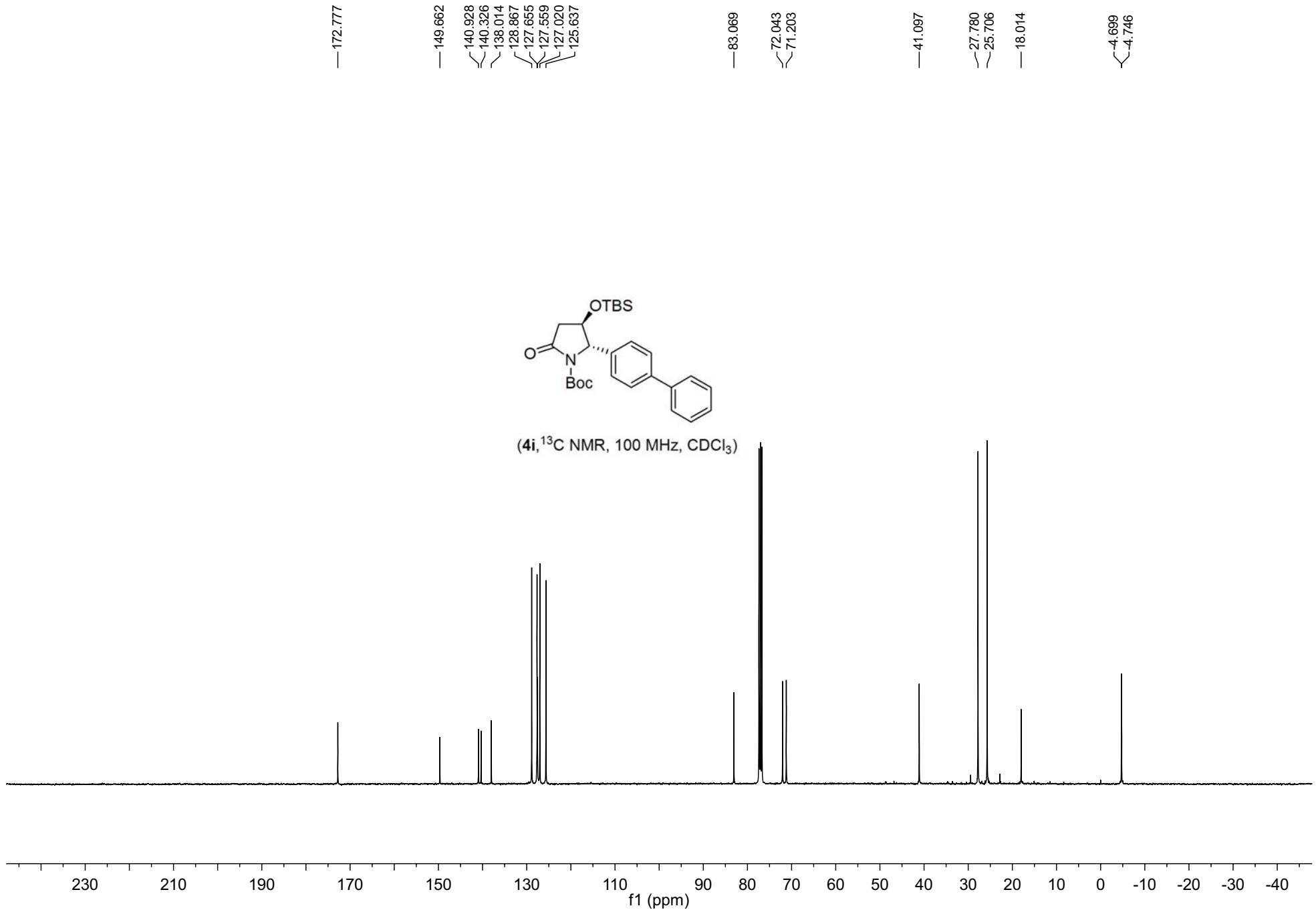
-62.781

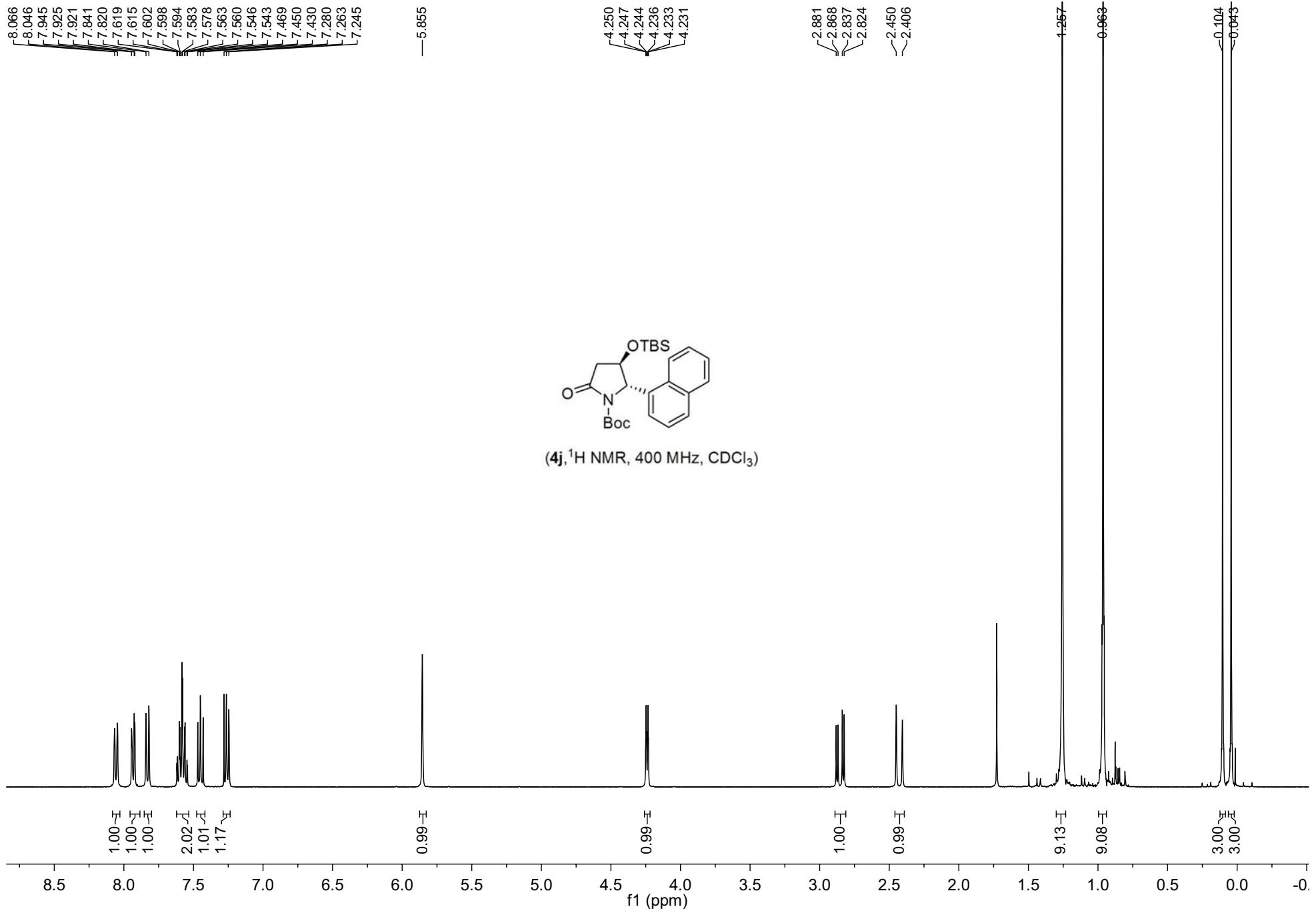


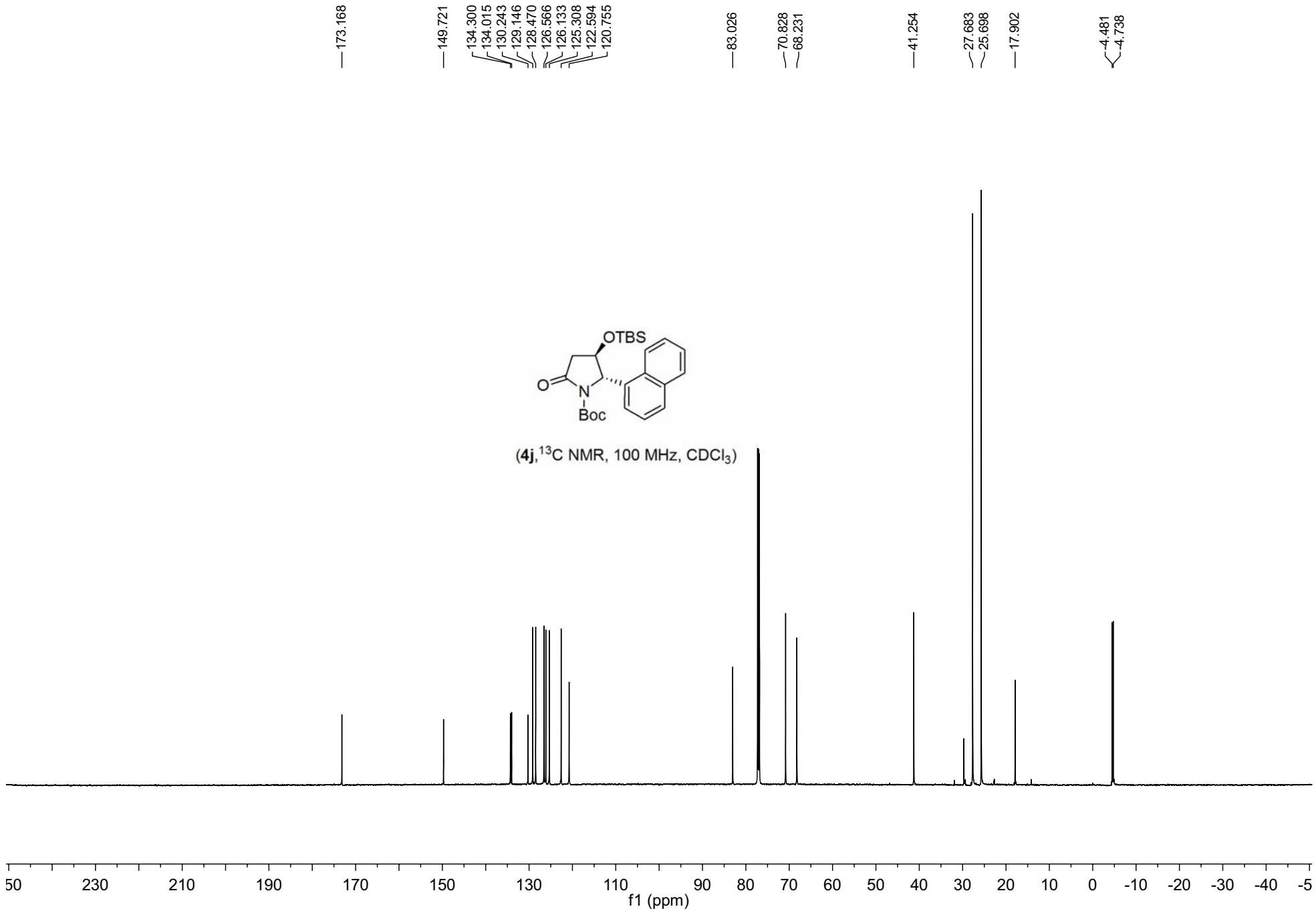
(**4h**,  $^{19}\text{F}$  NMR, 376 MHz,  $\text{CDCl}_3$ )

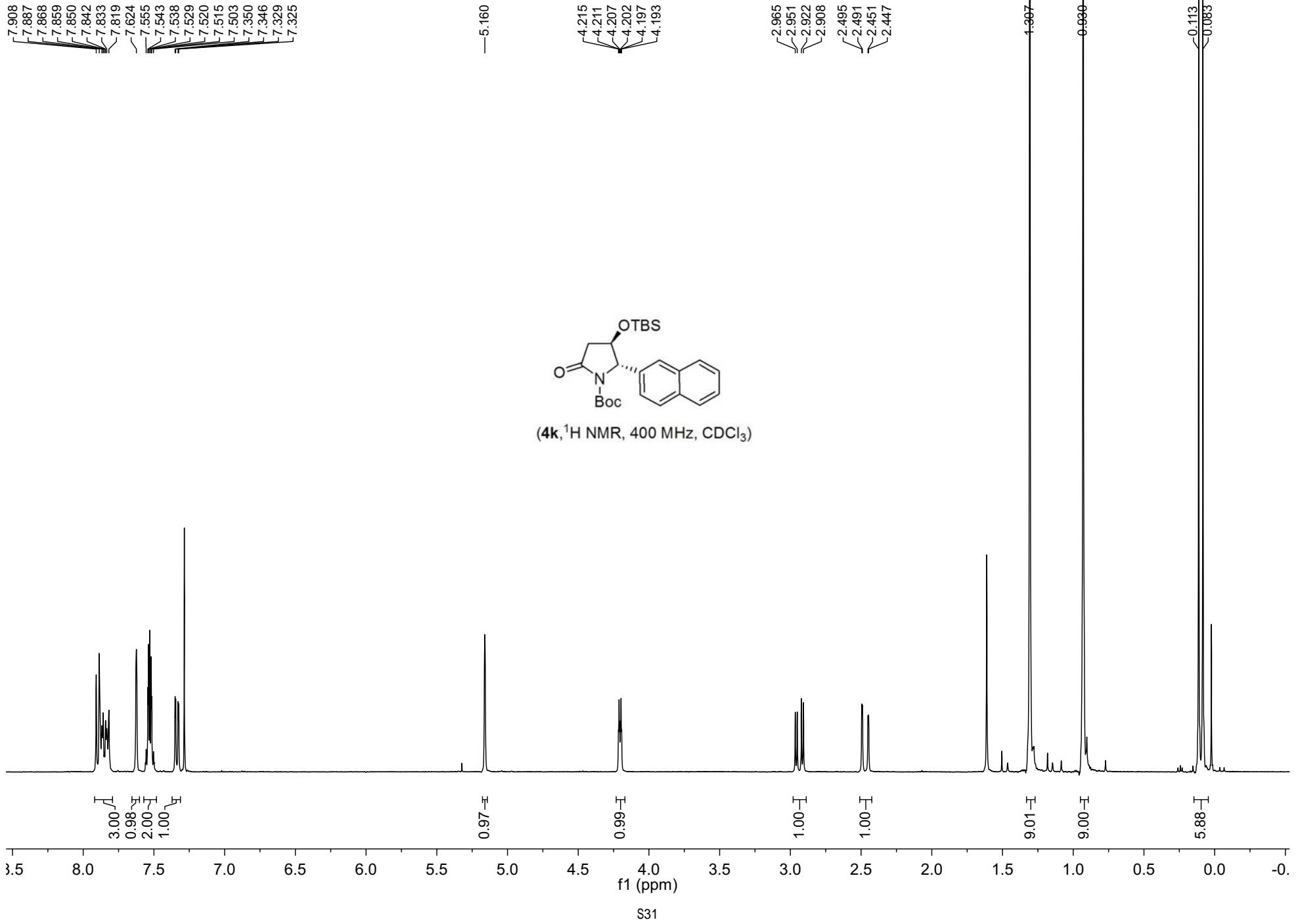


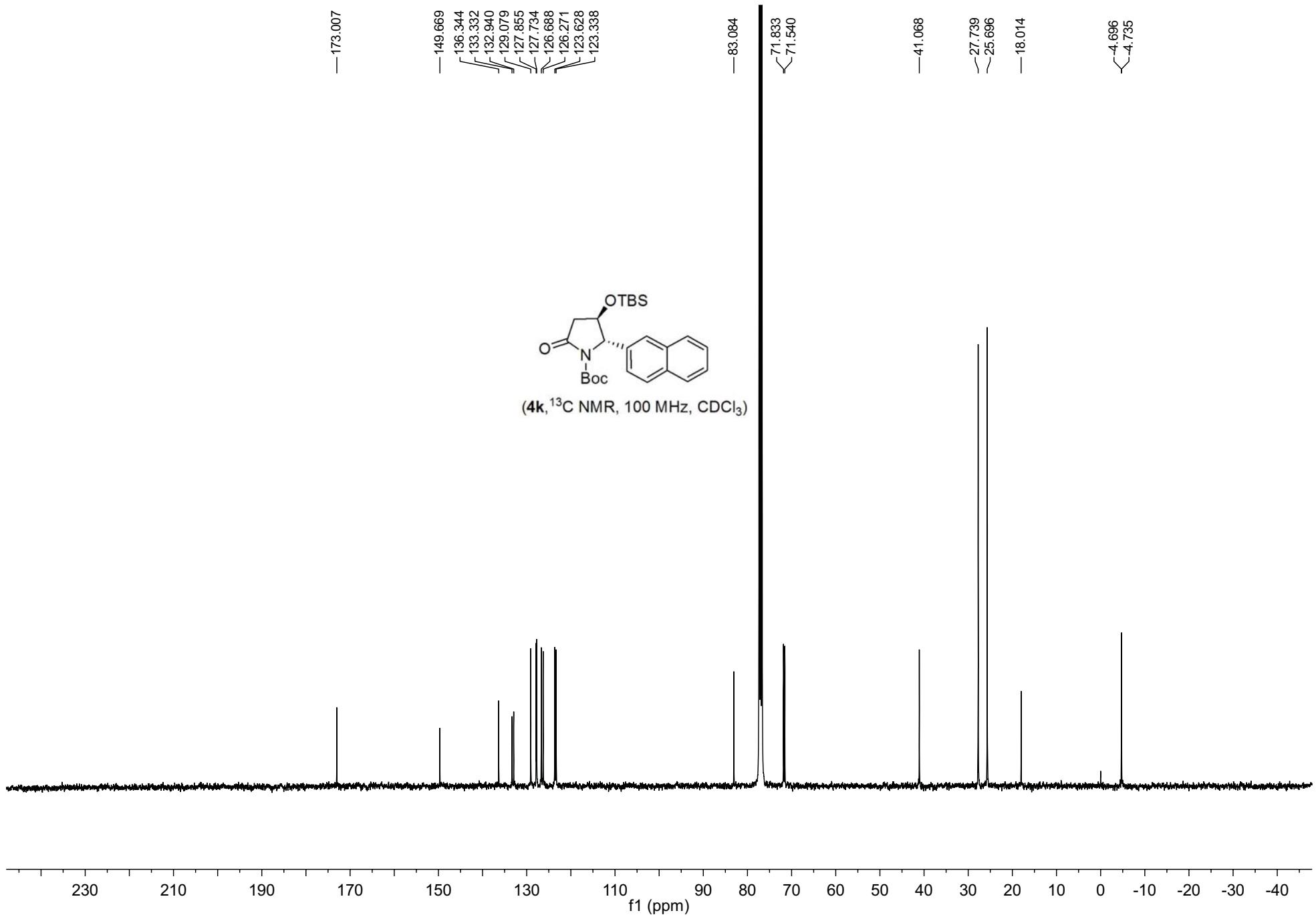


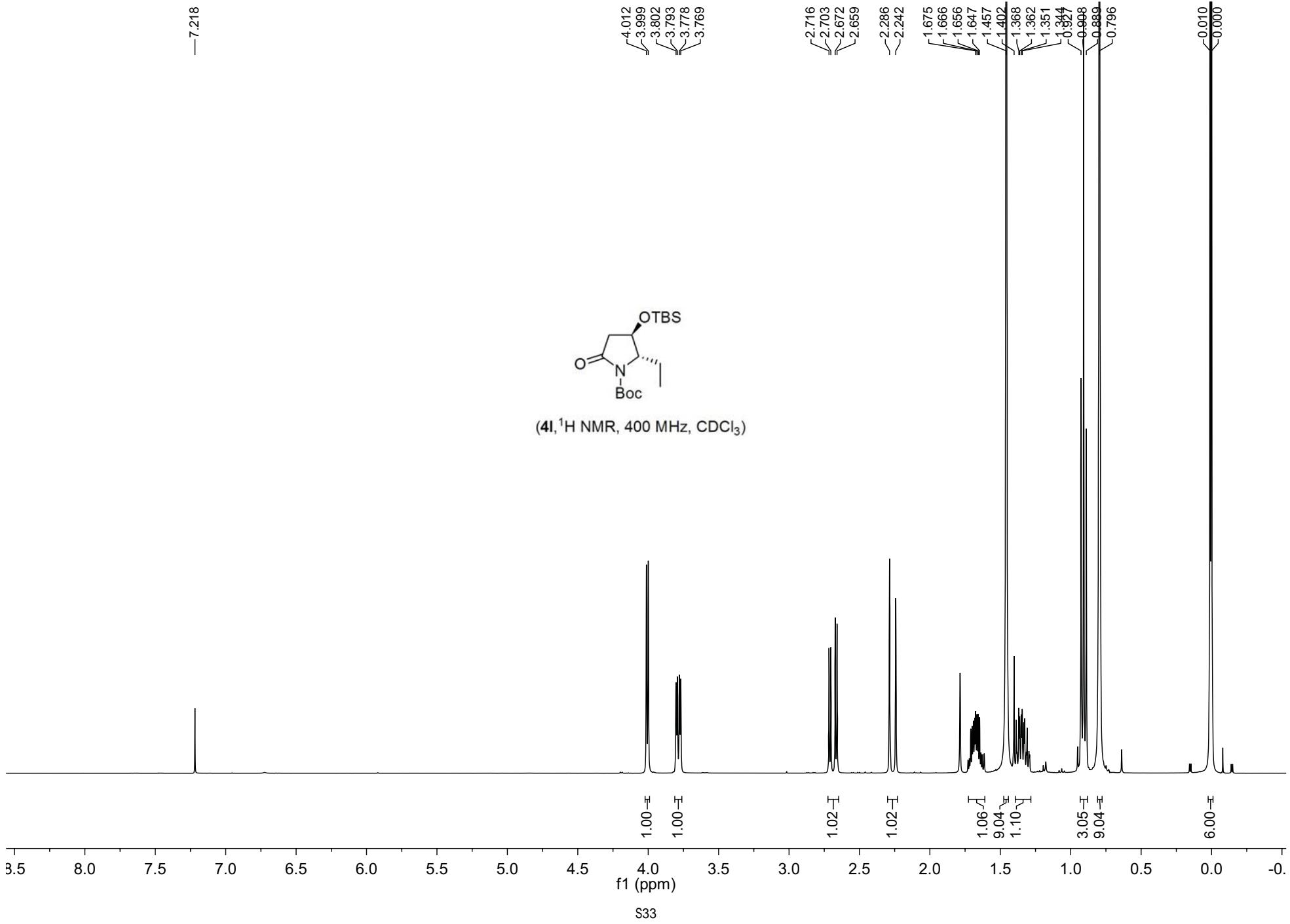


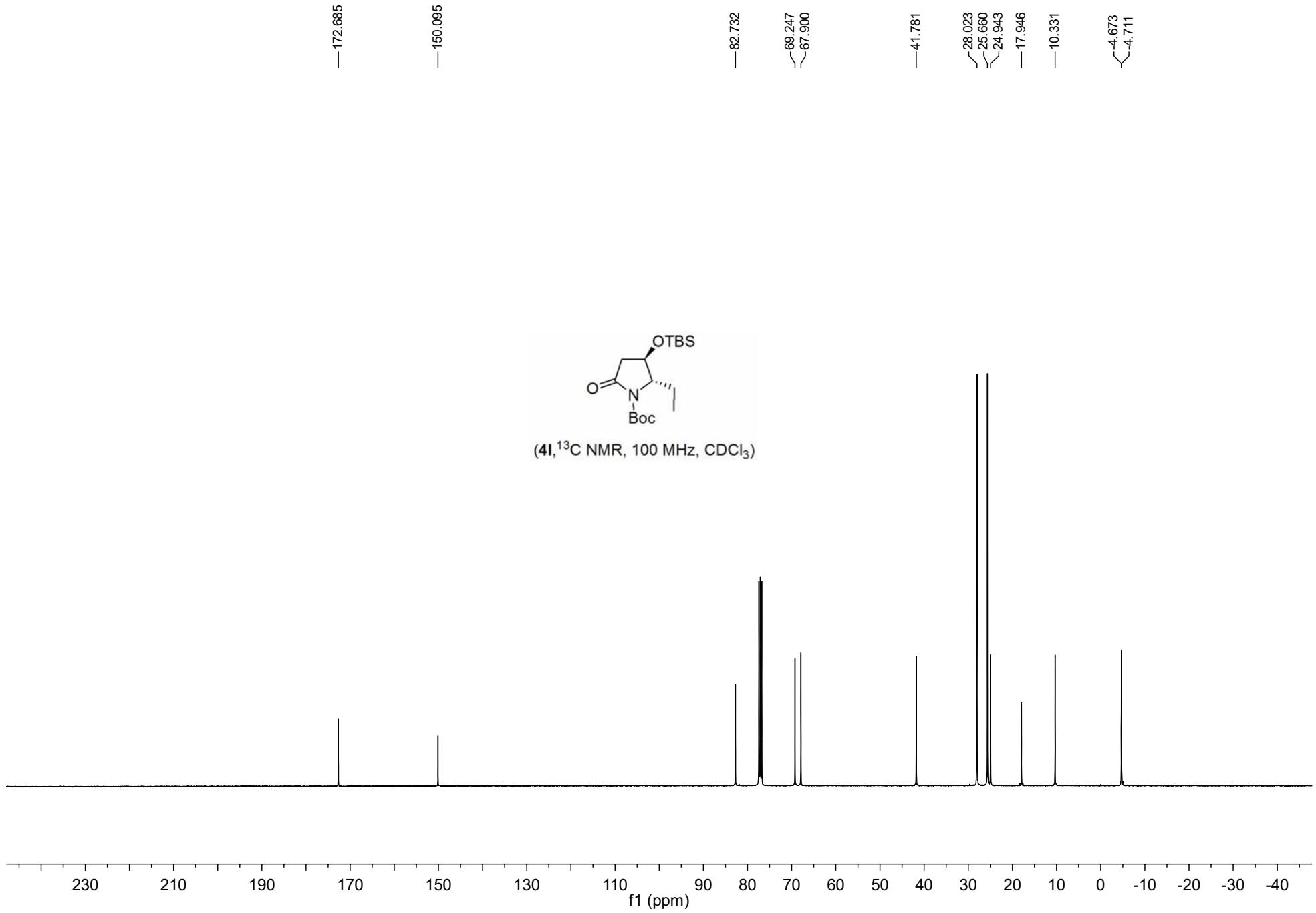


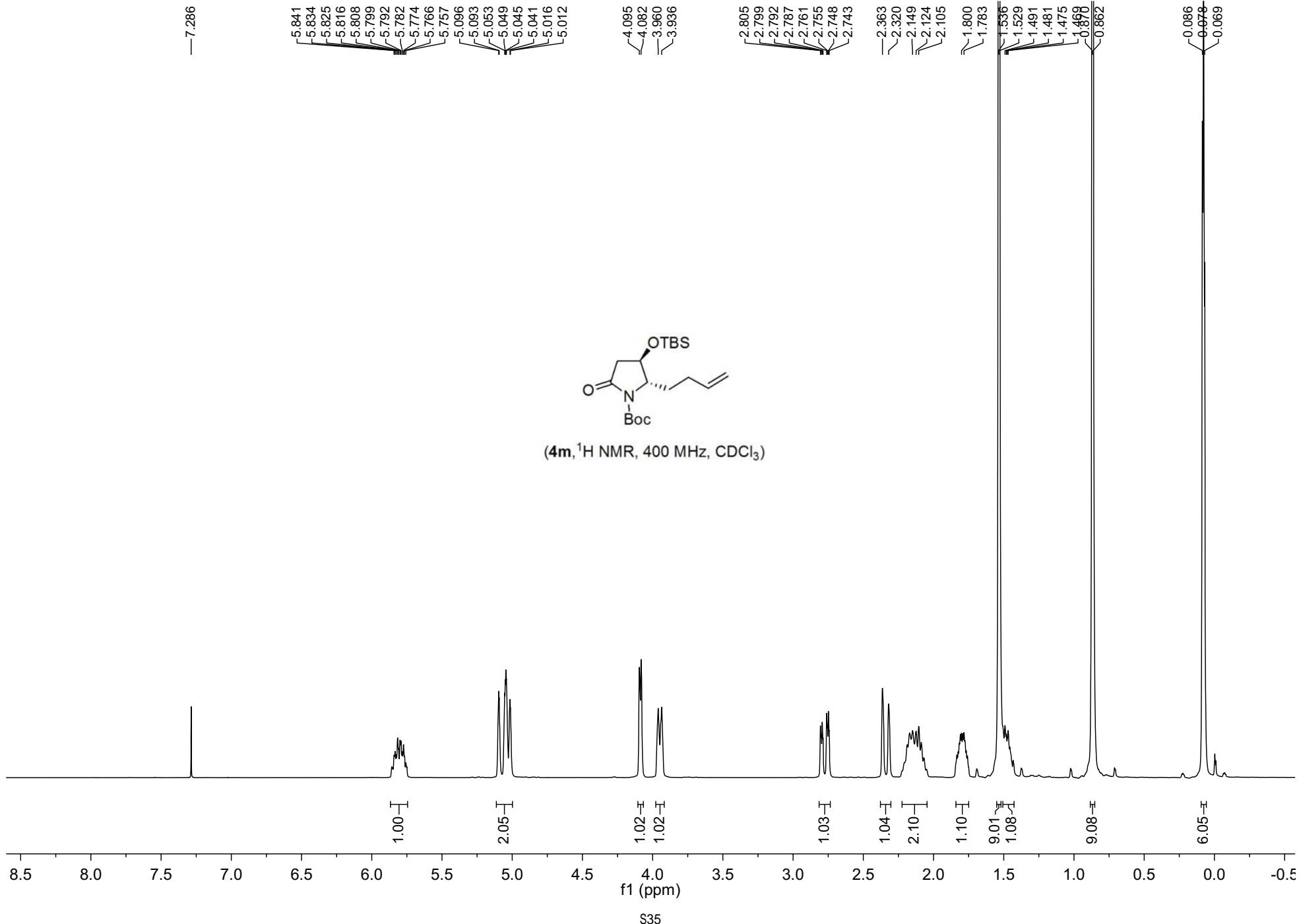


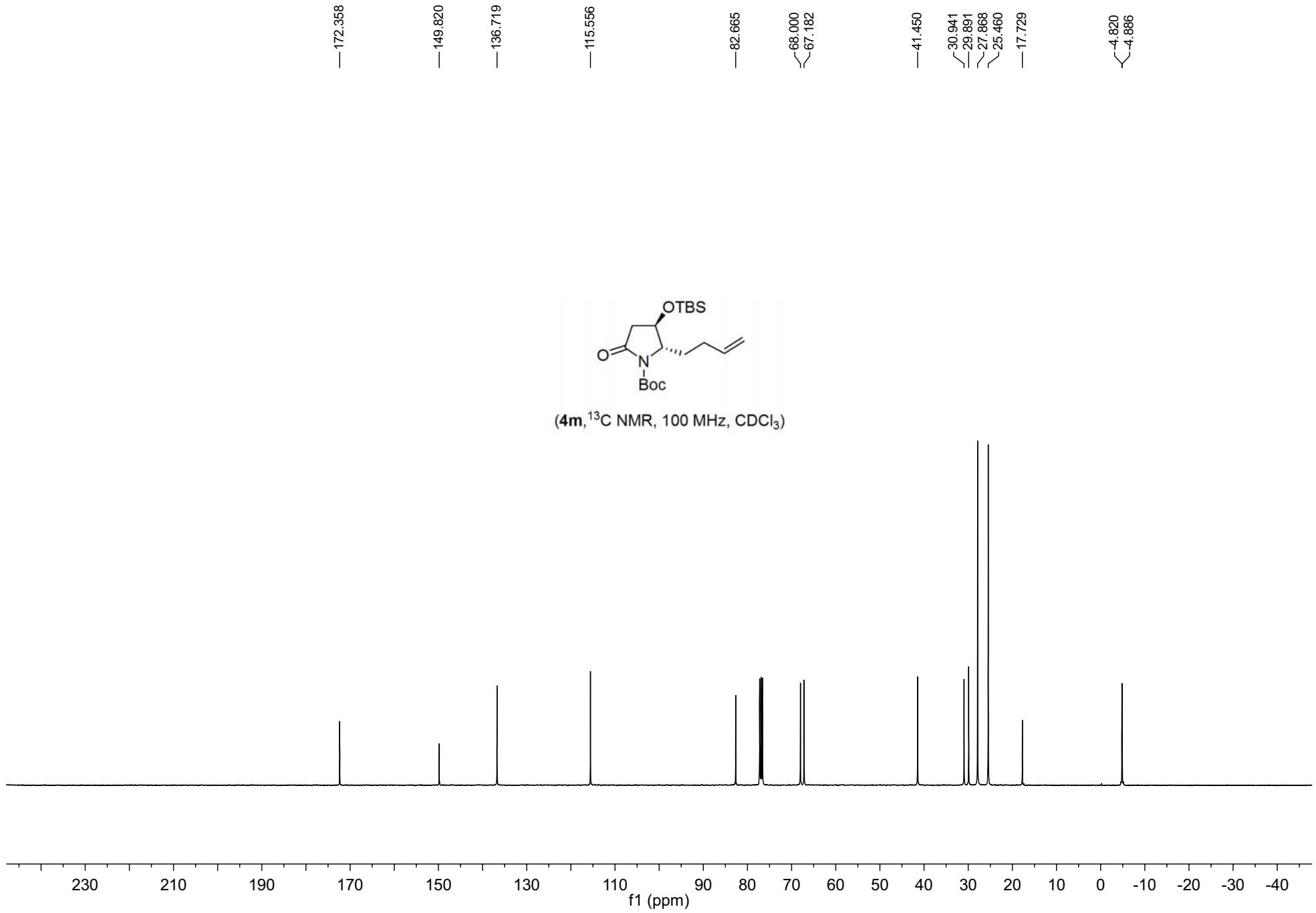


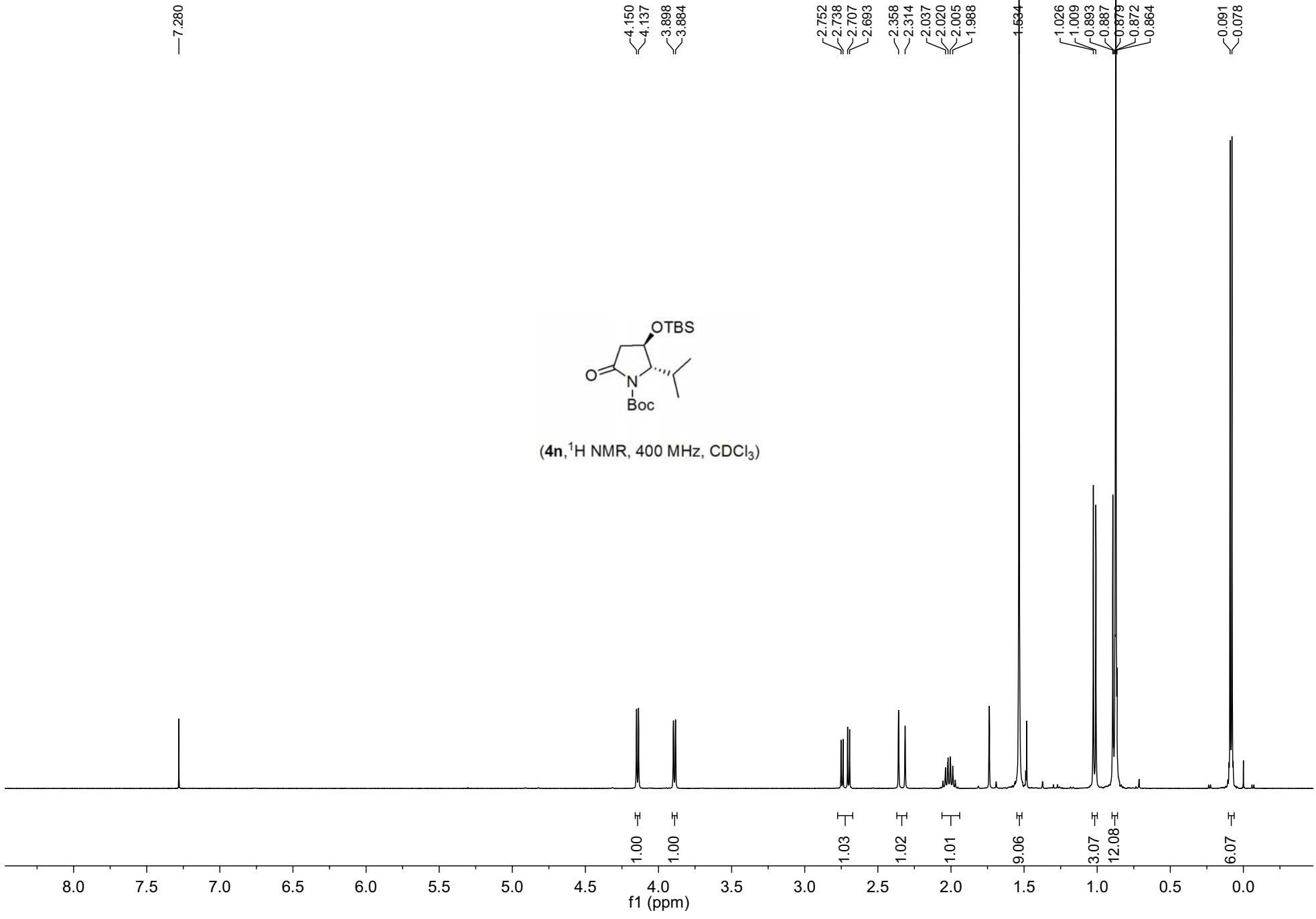


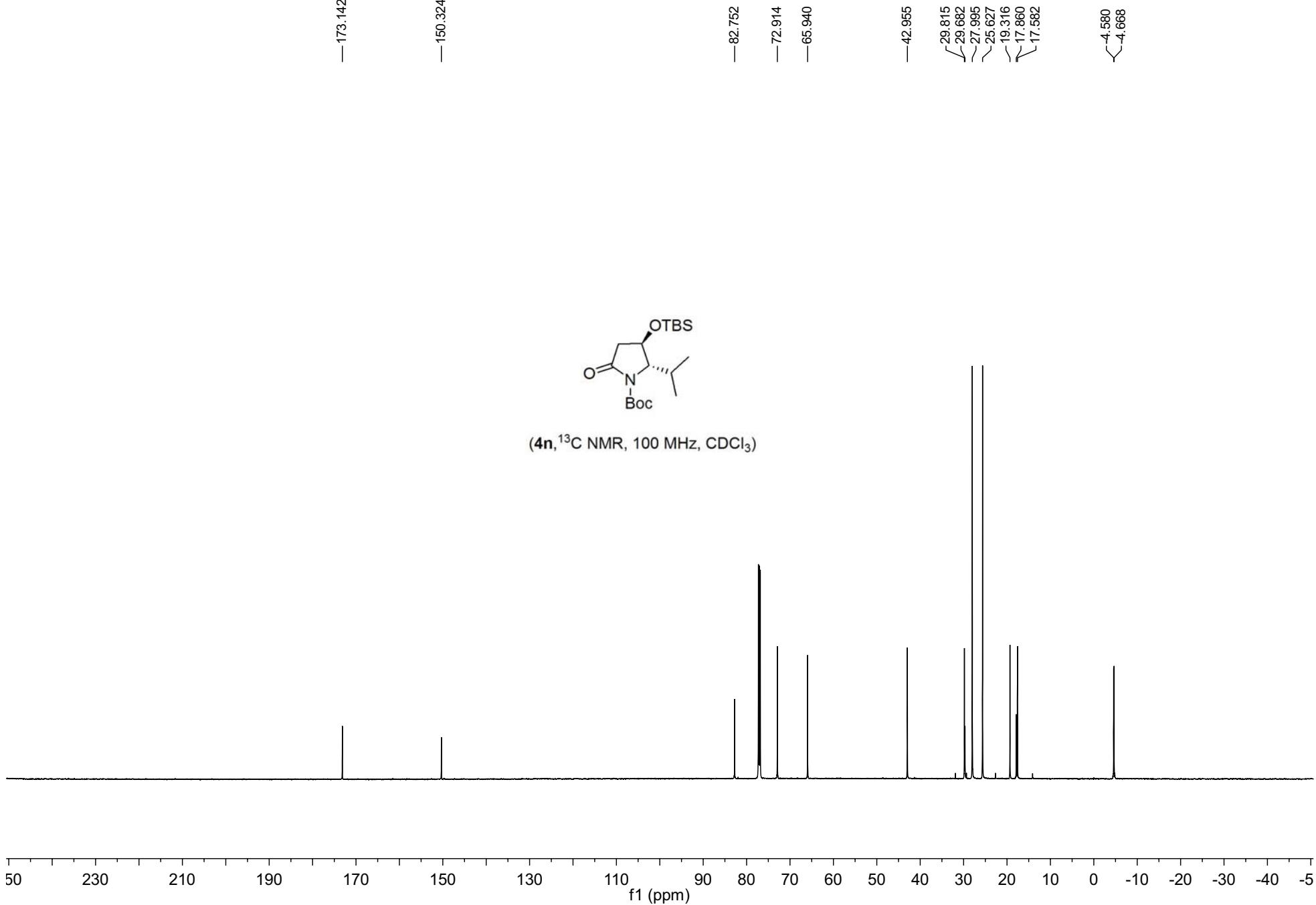


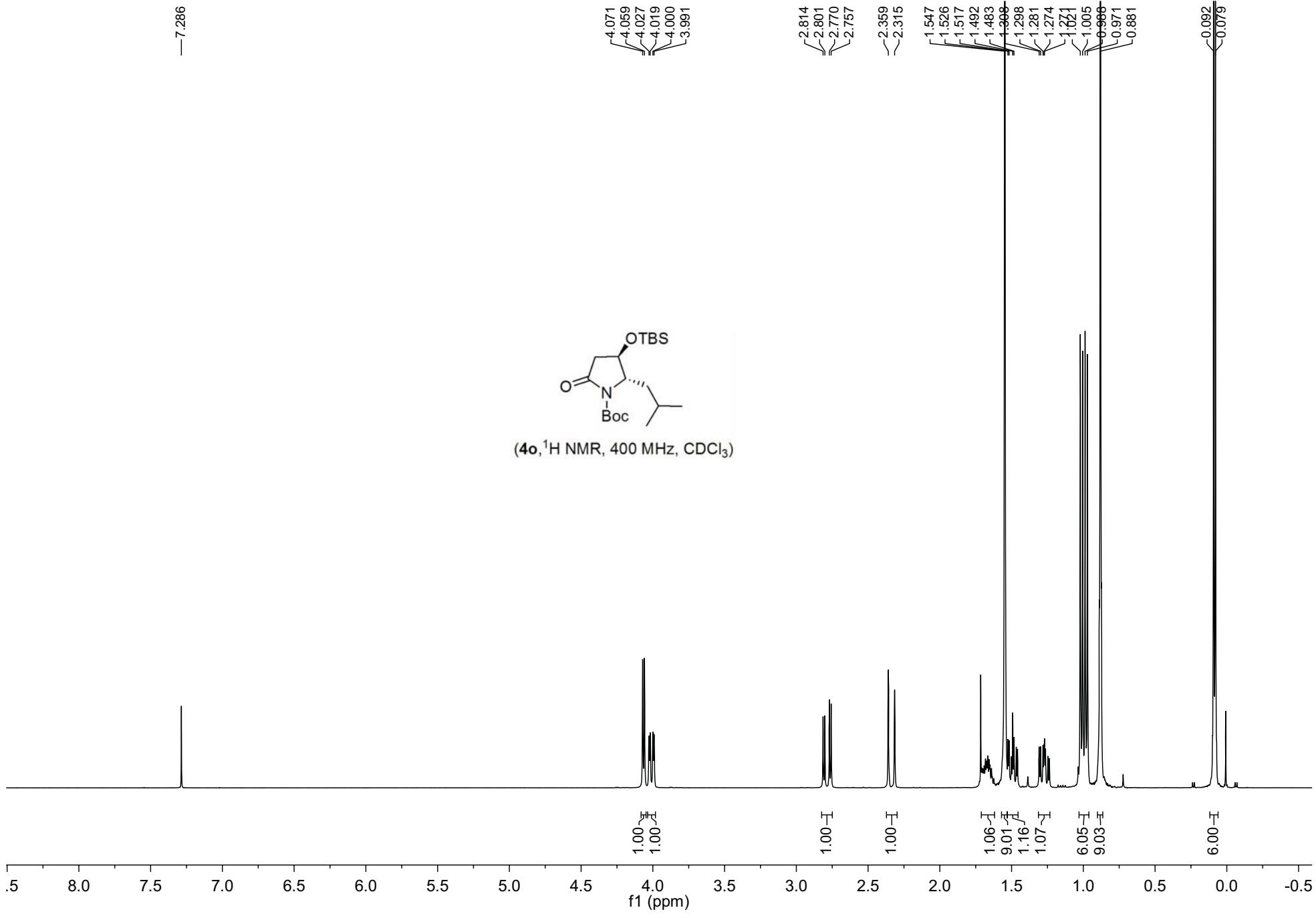


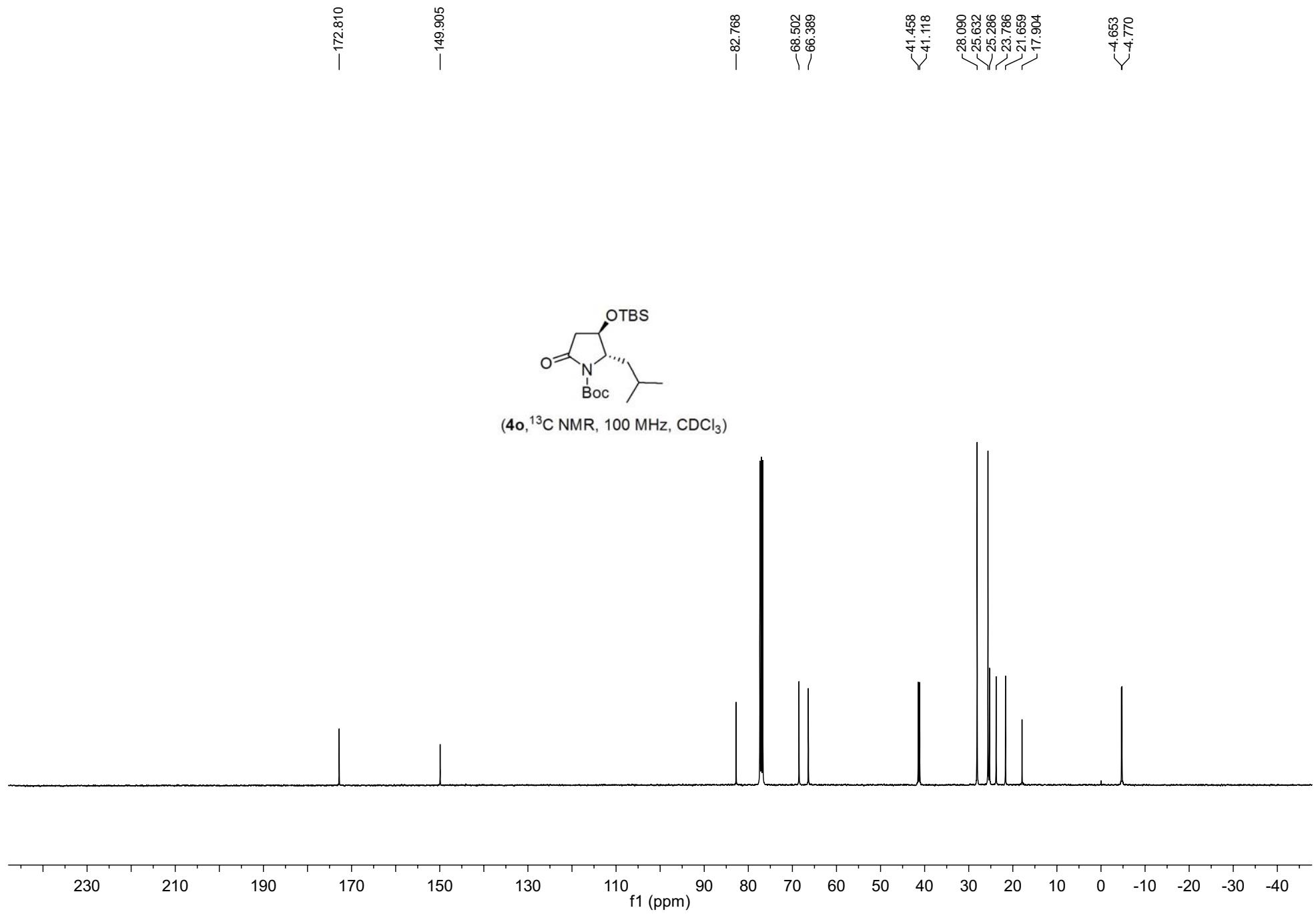


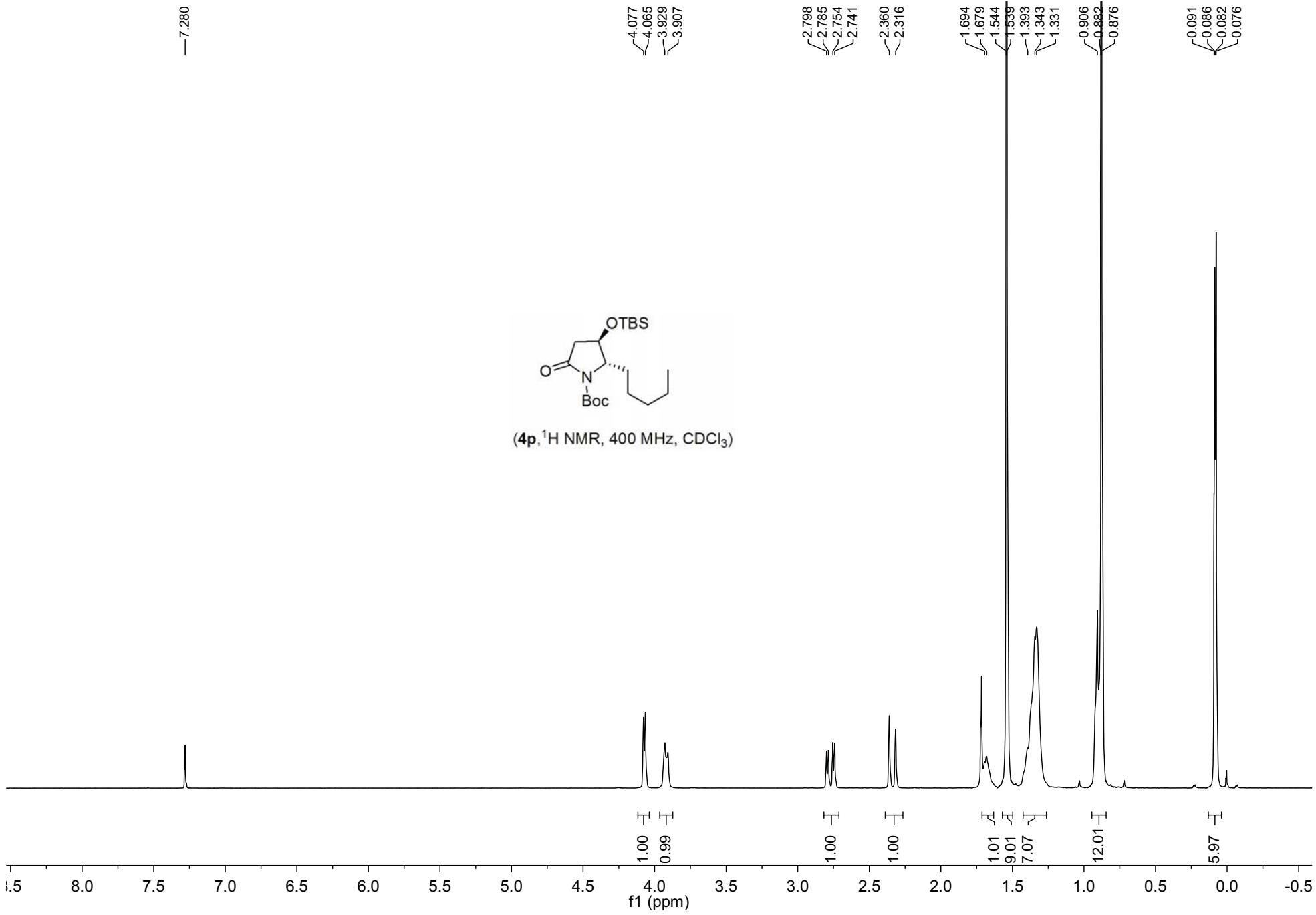


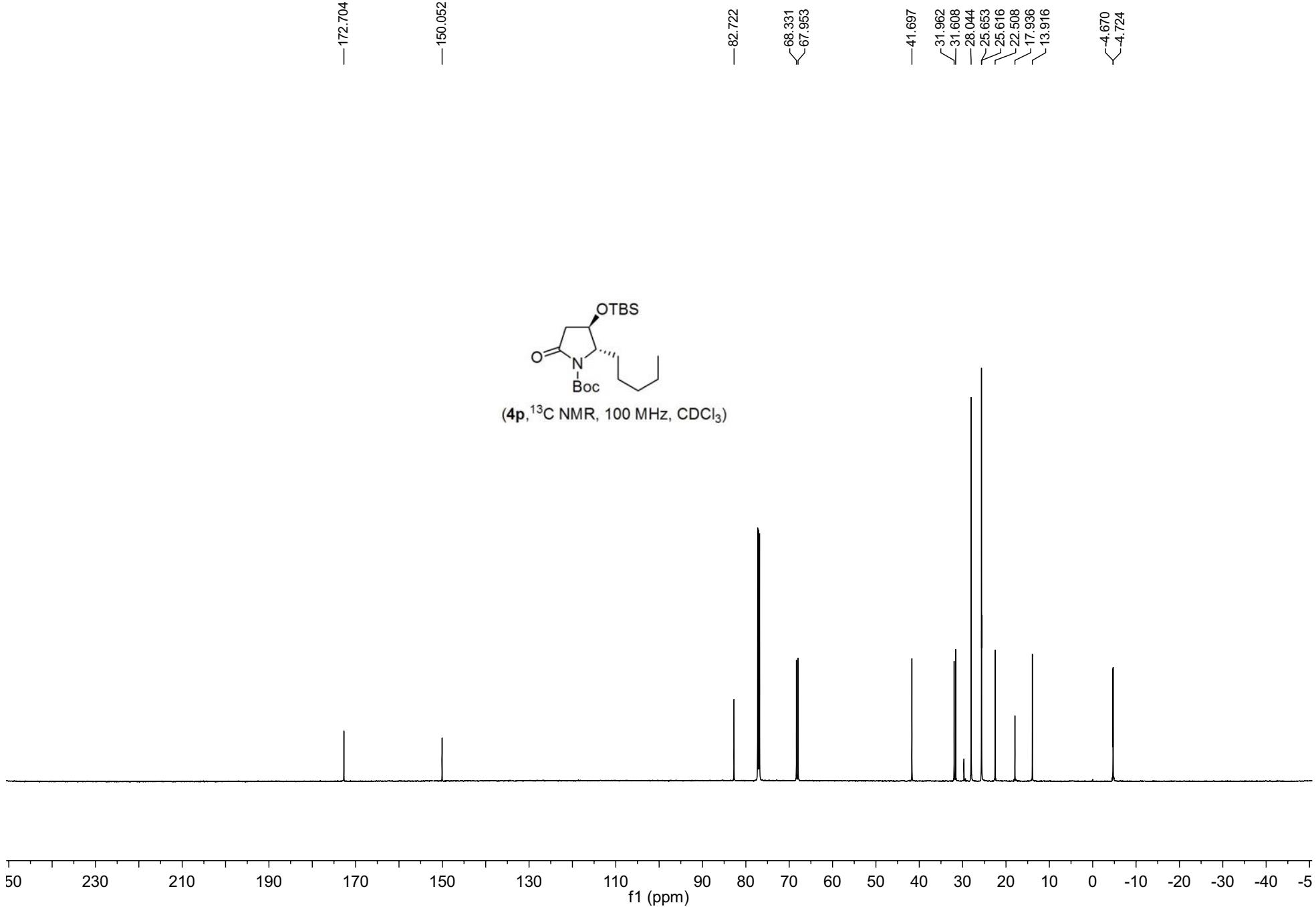


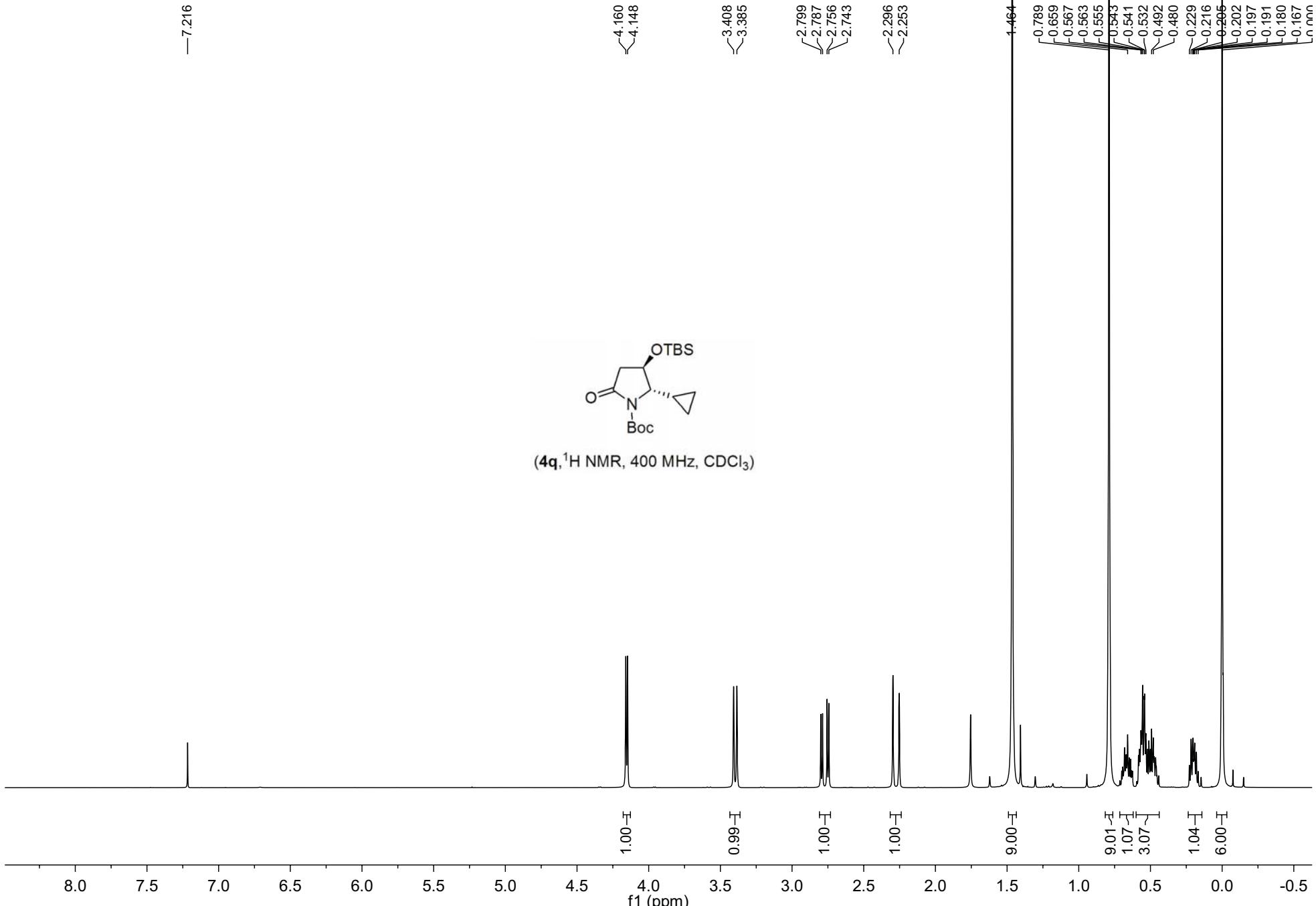




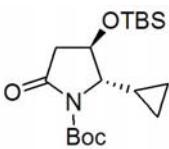


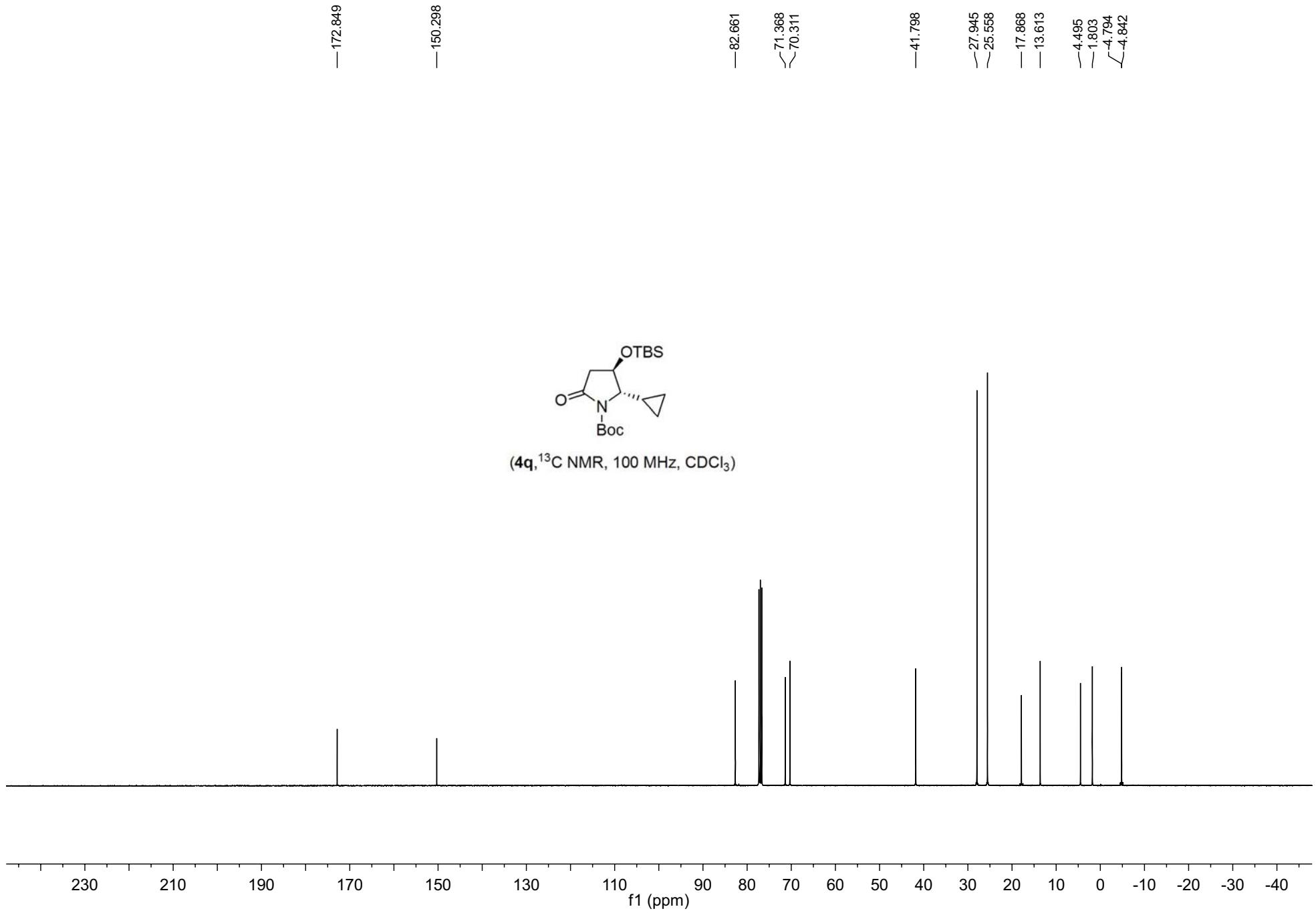


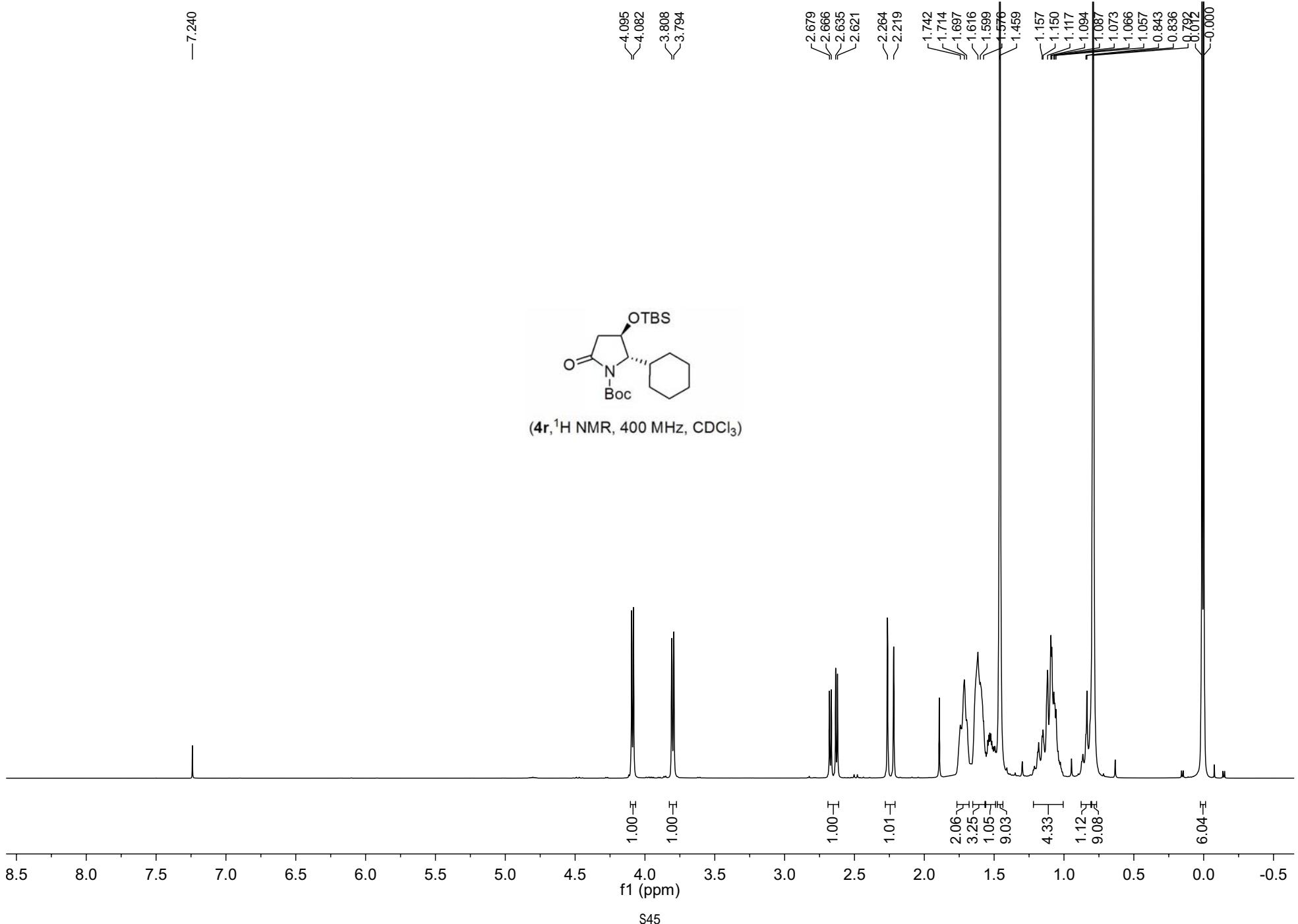


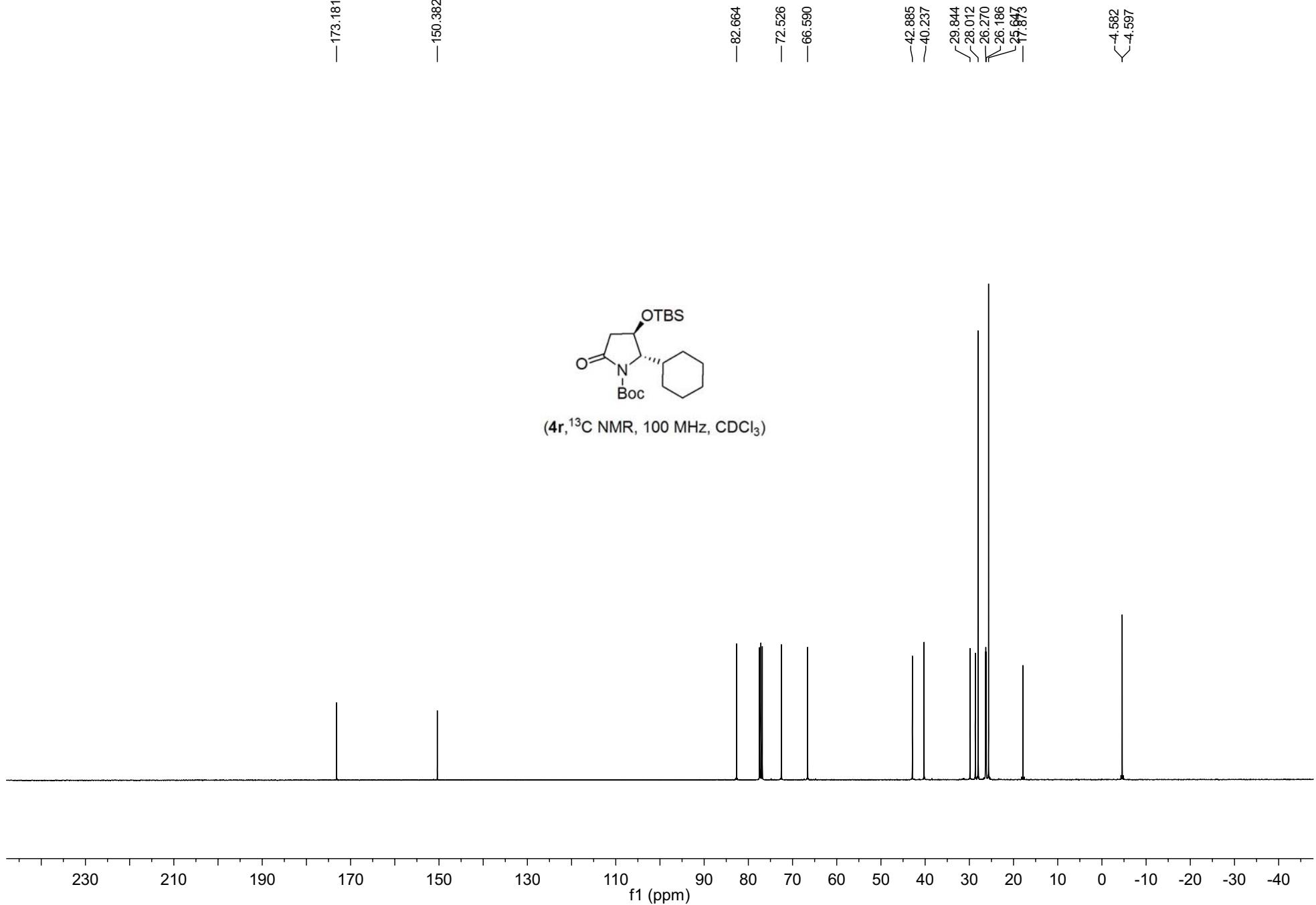


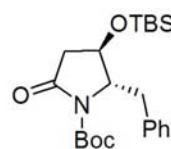
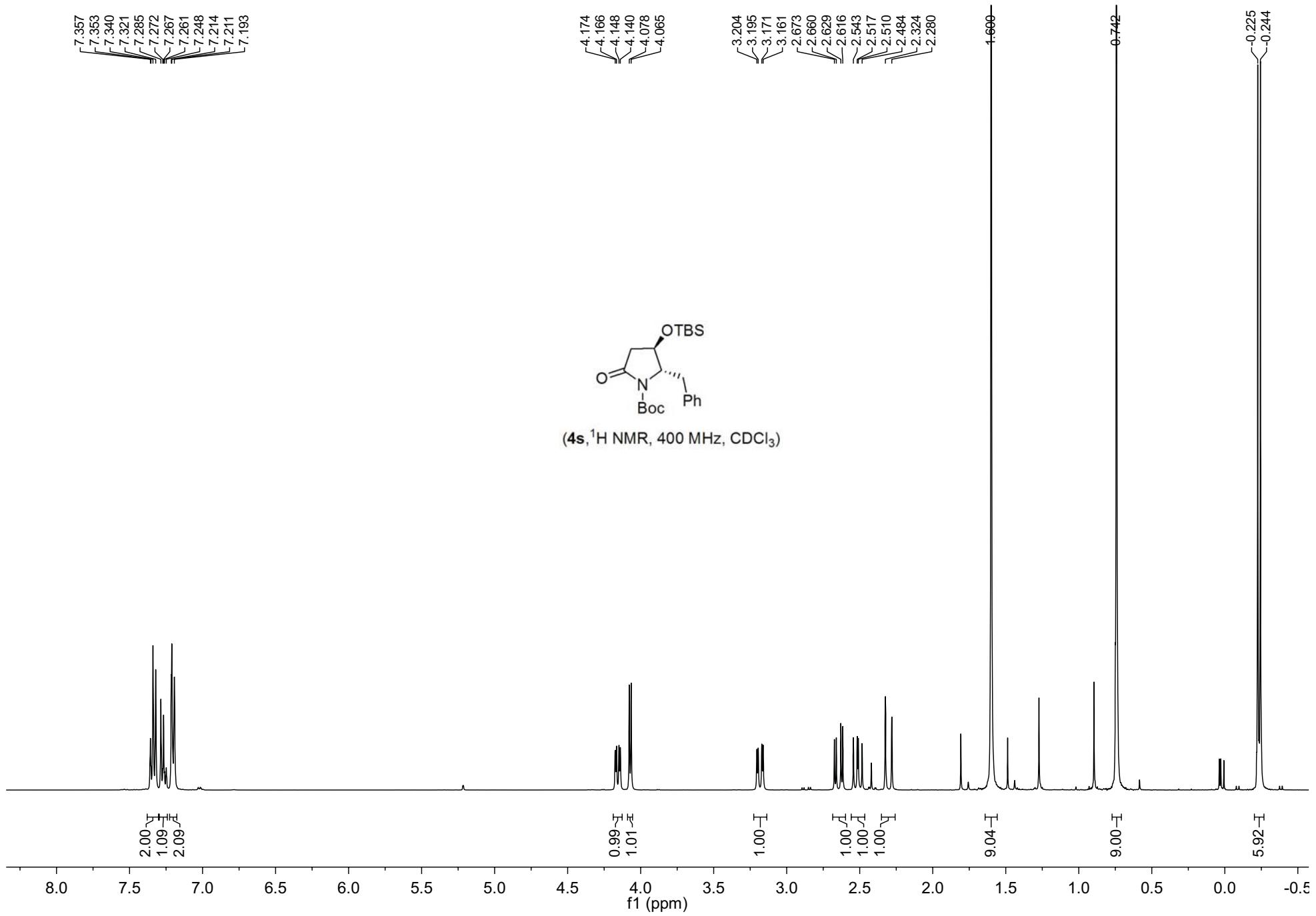
(**4q**,  $^1\text{H}$  NMR, 400 MHz,  $\text{CDCl}_3$ )

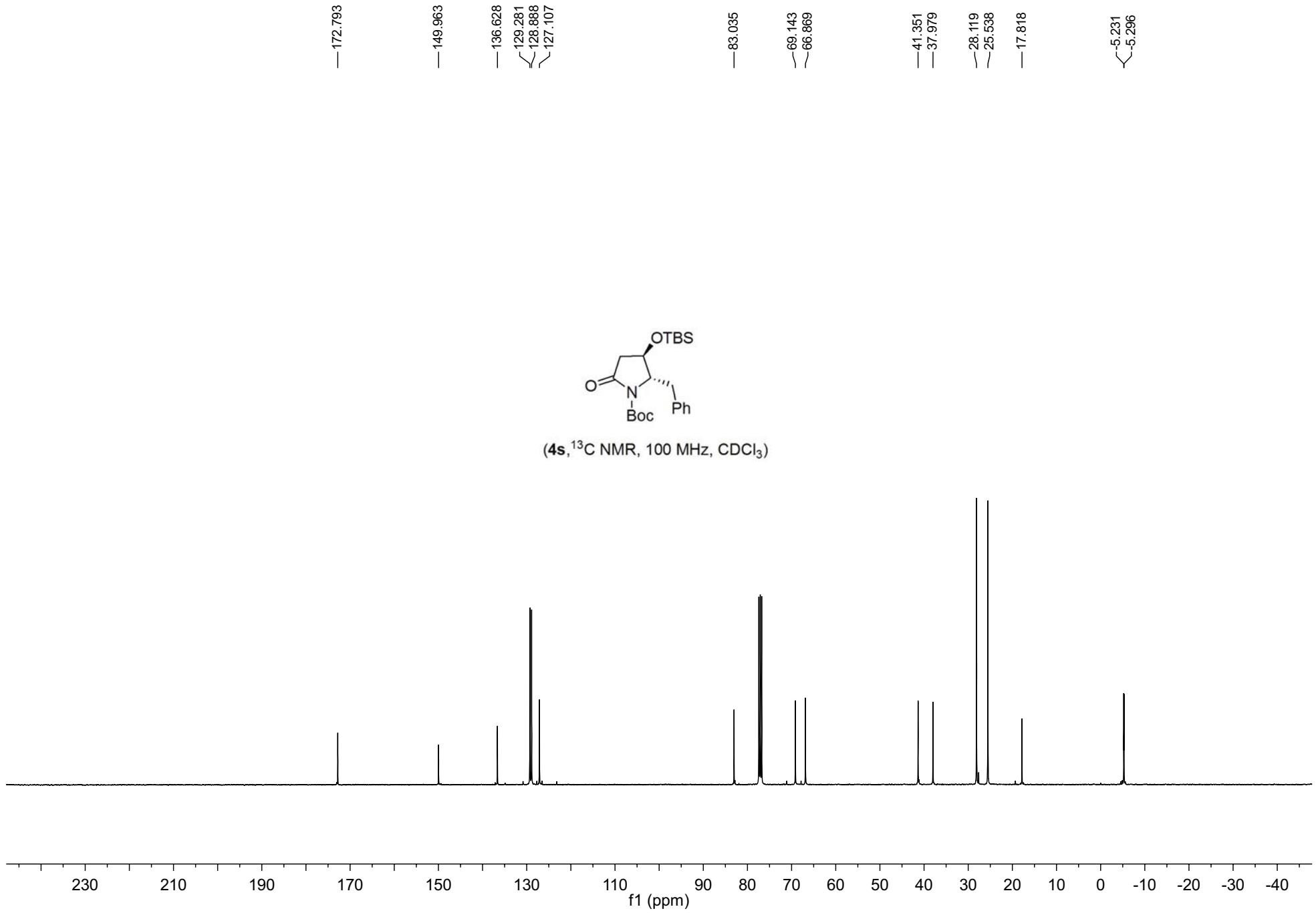


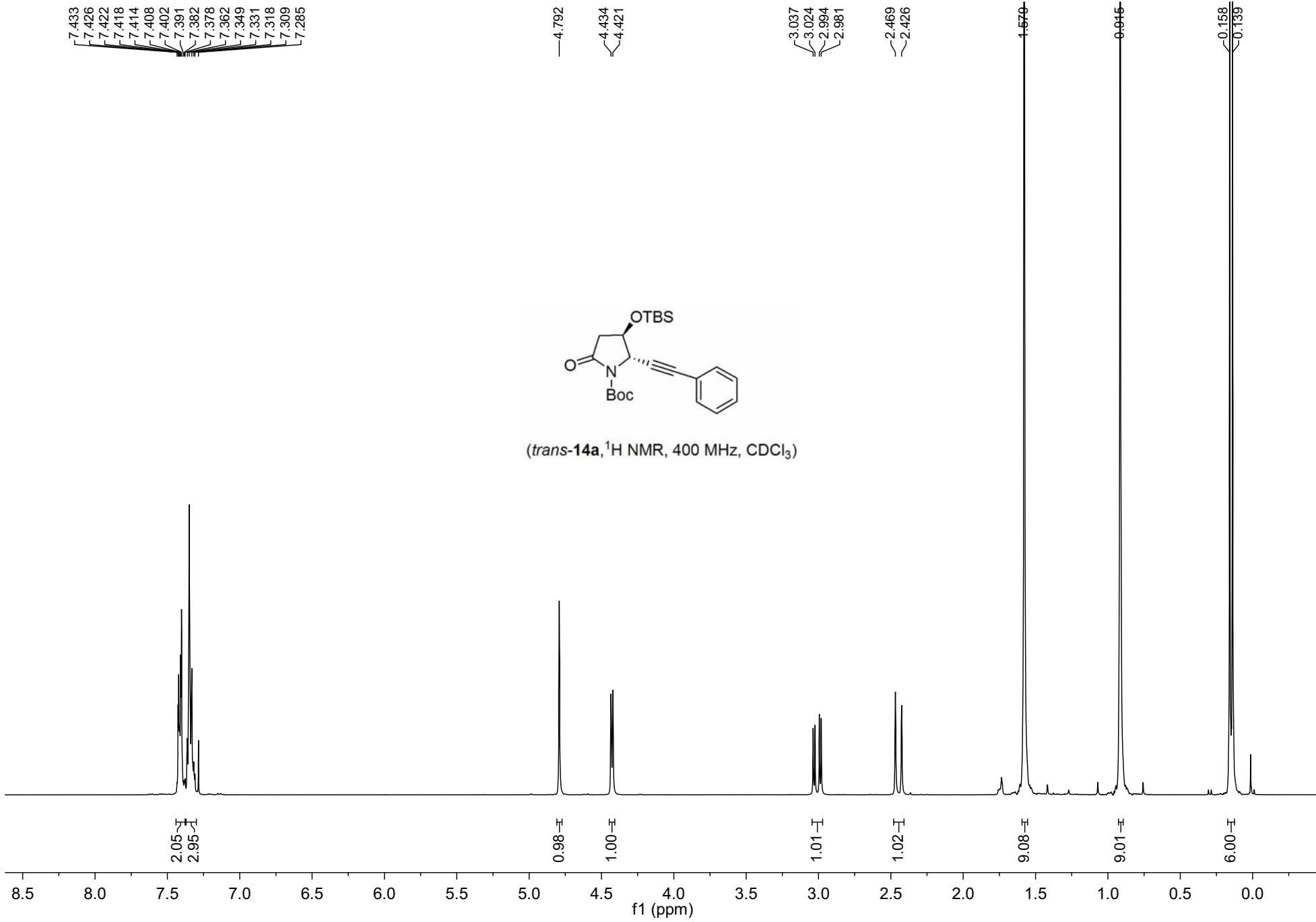


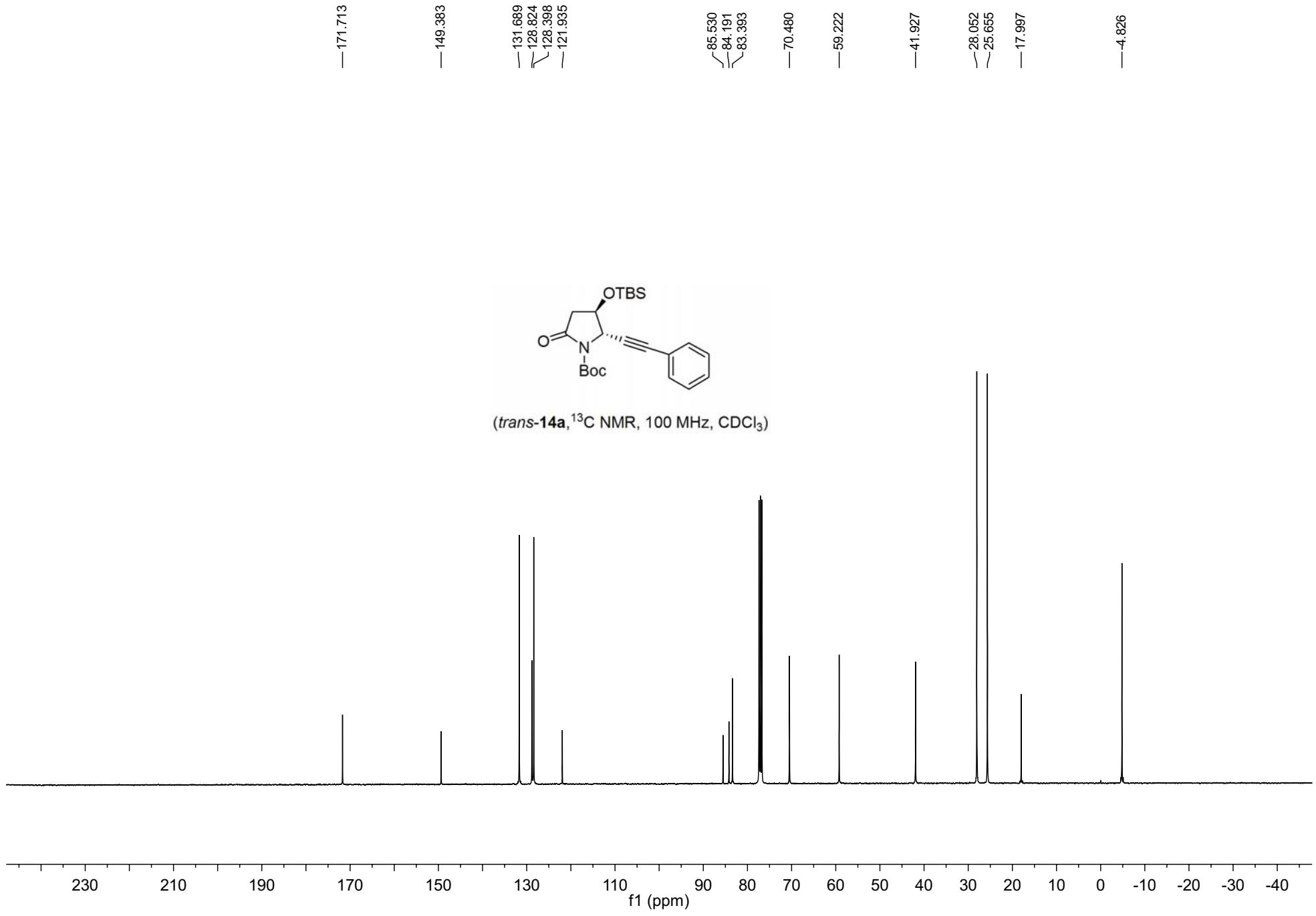


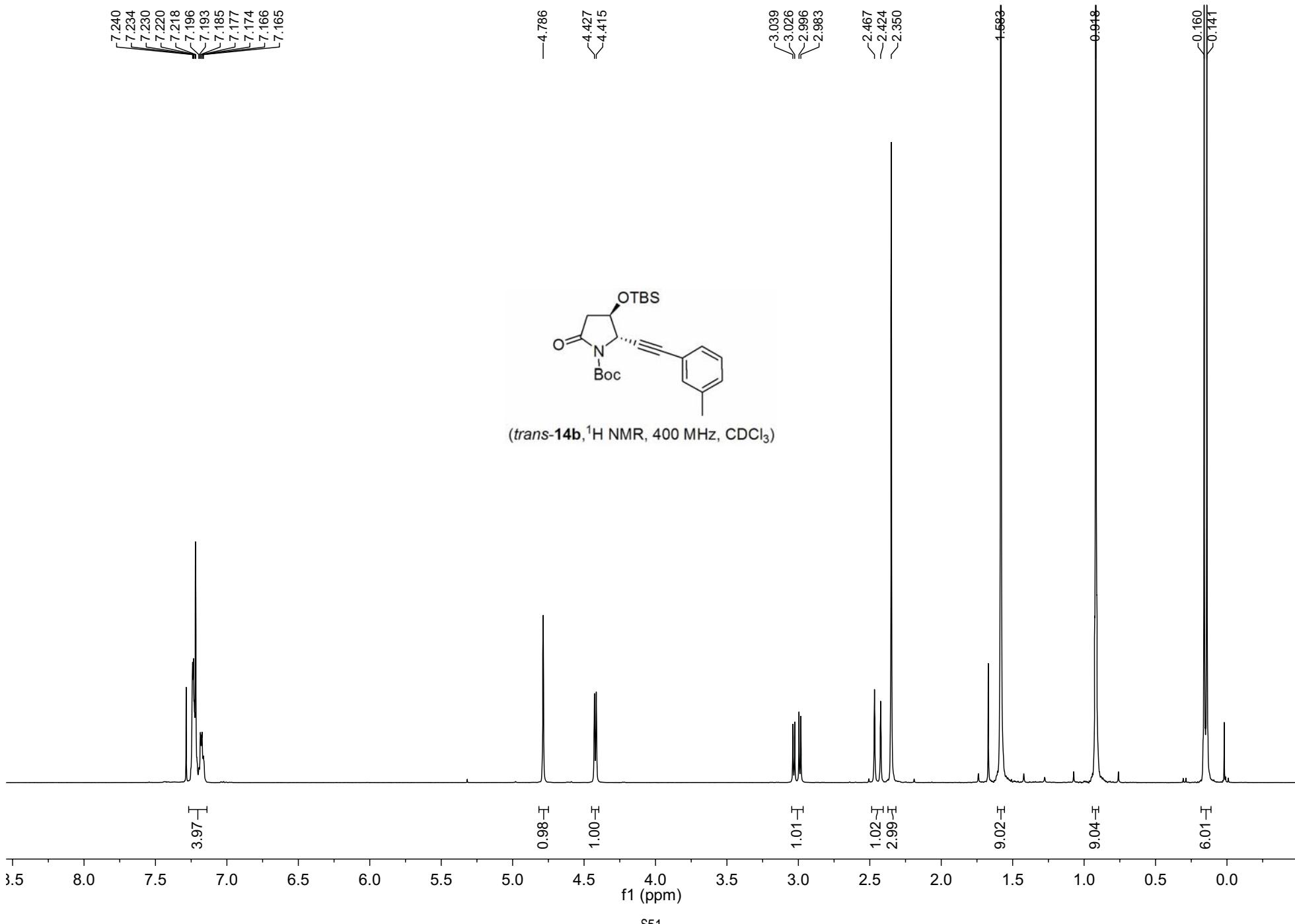


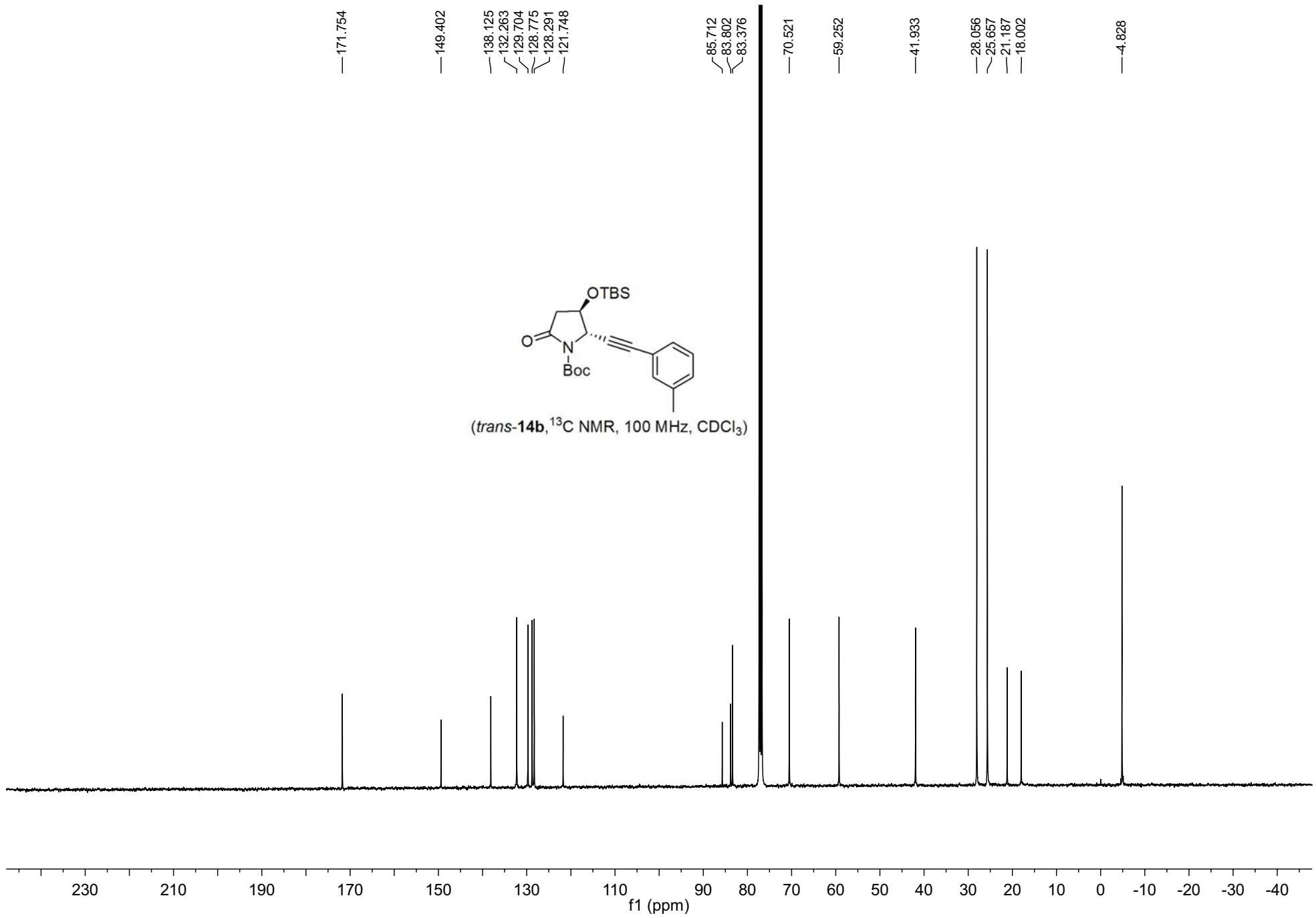


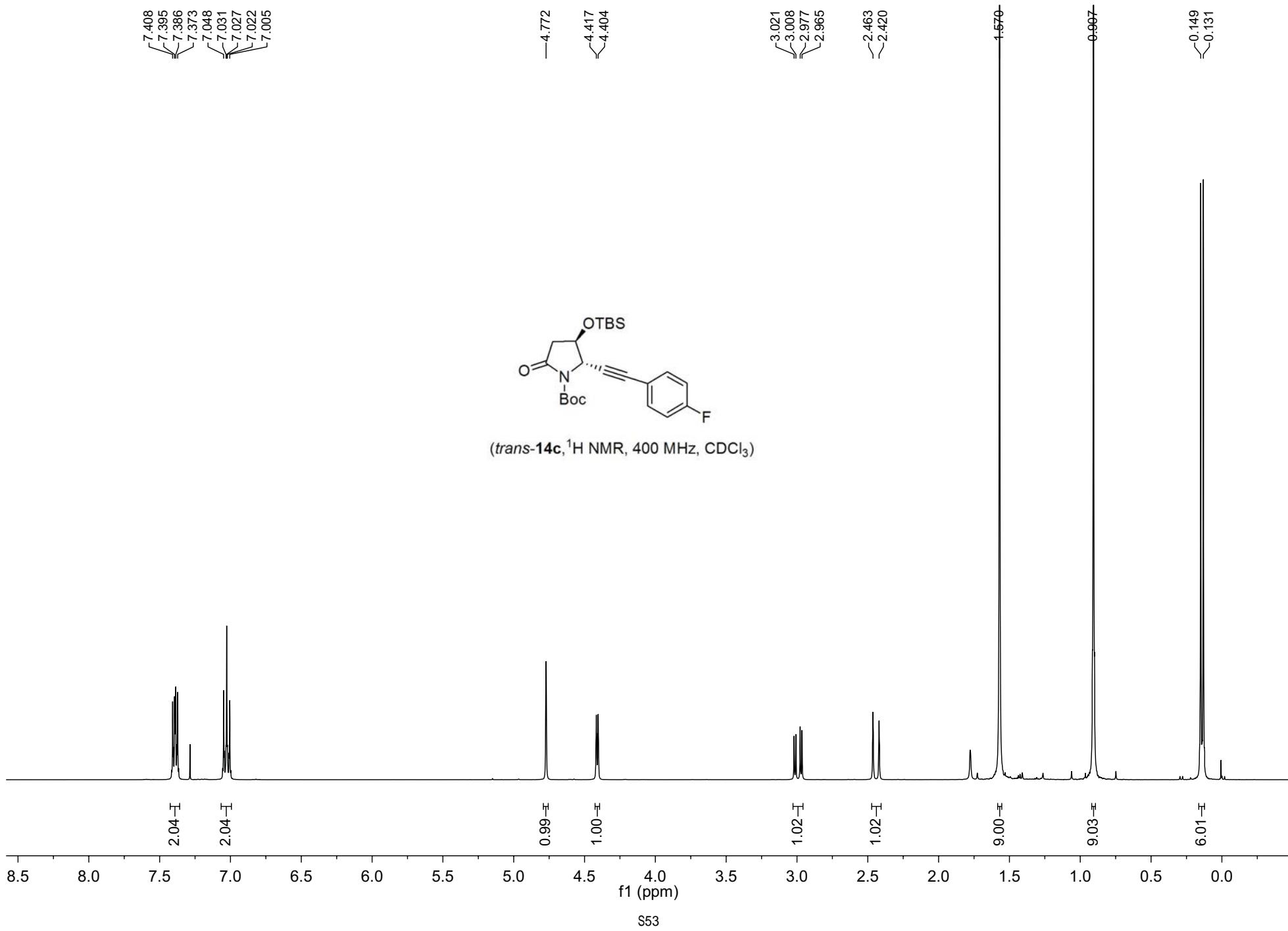


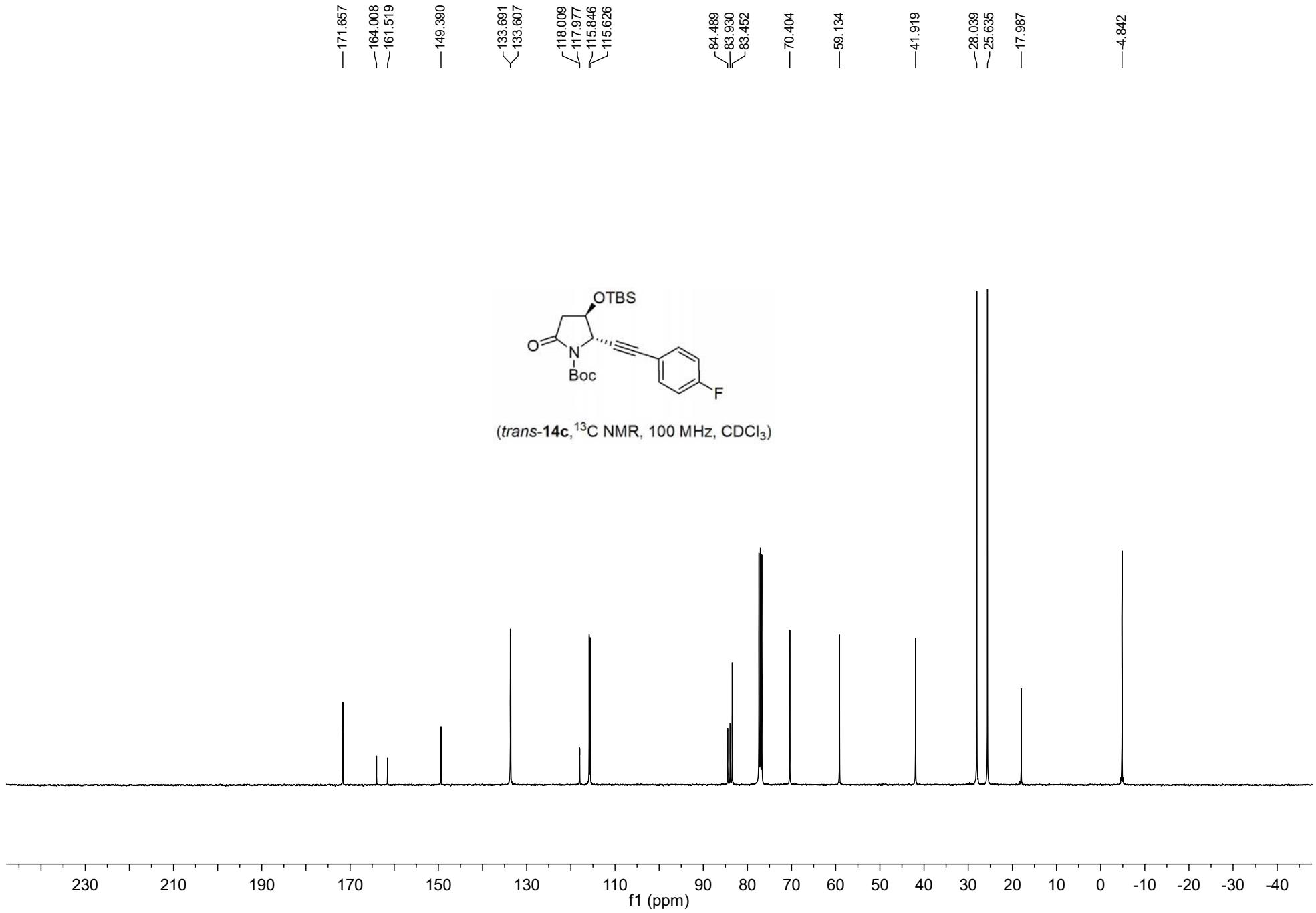




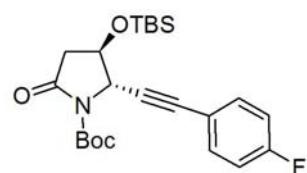




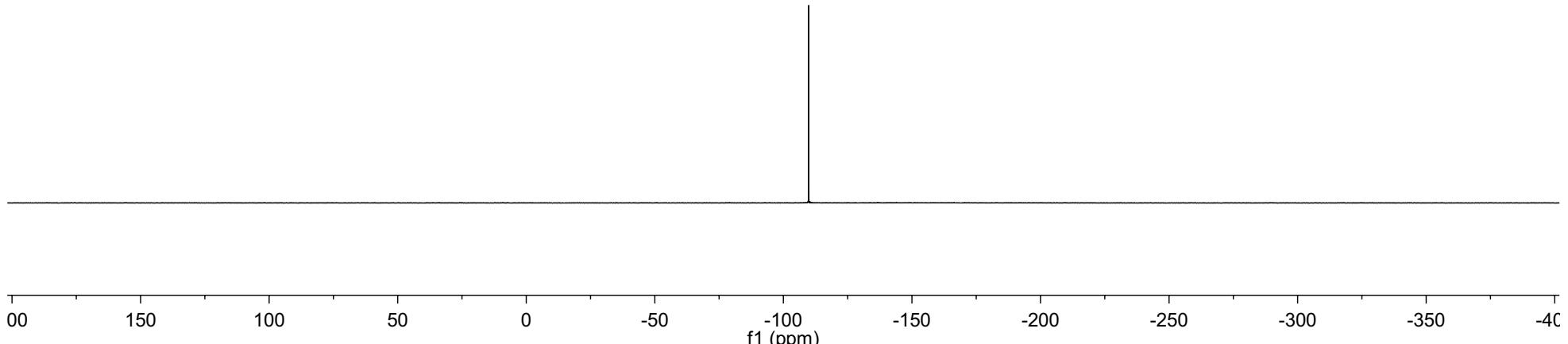


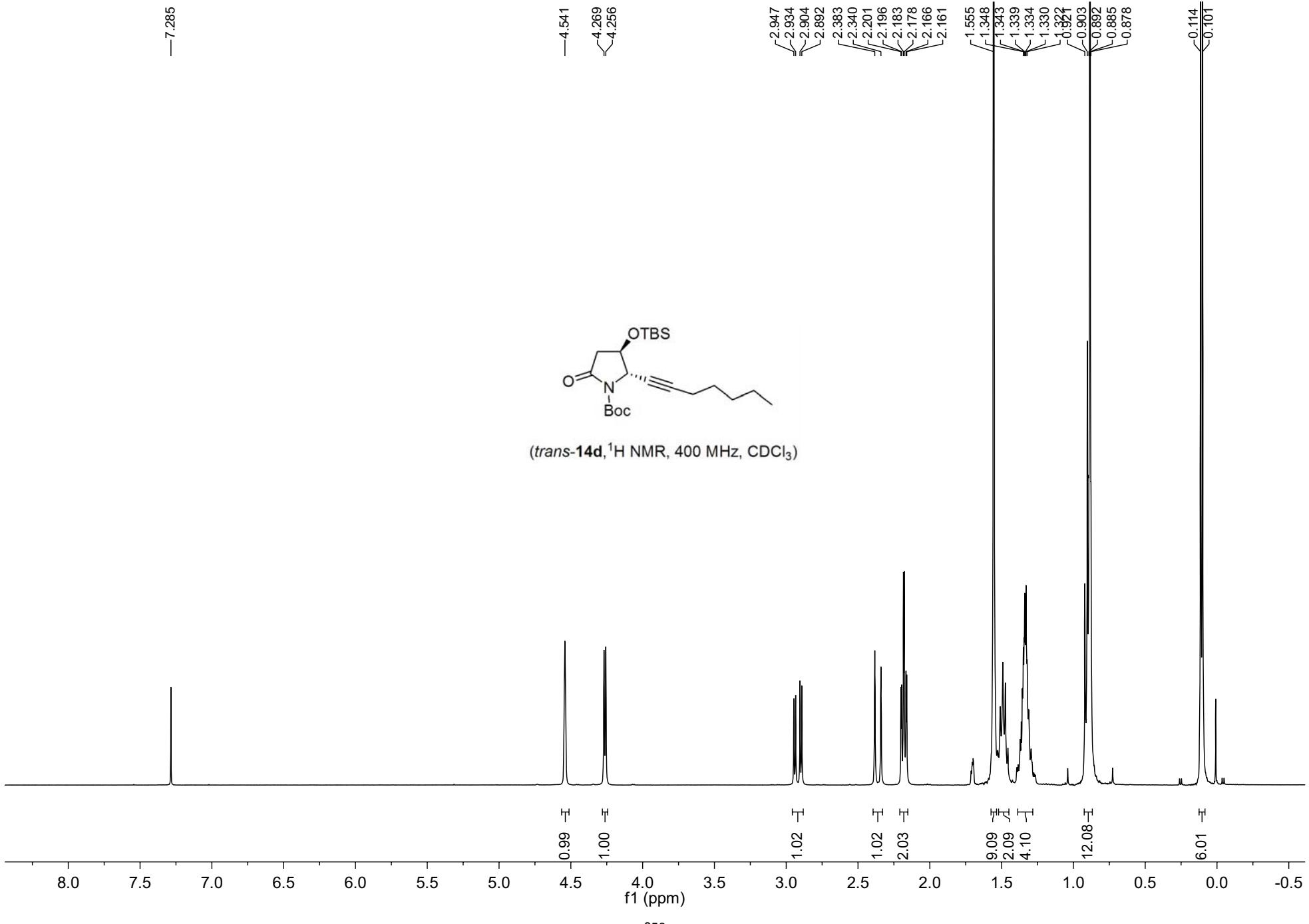


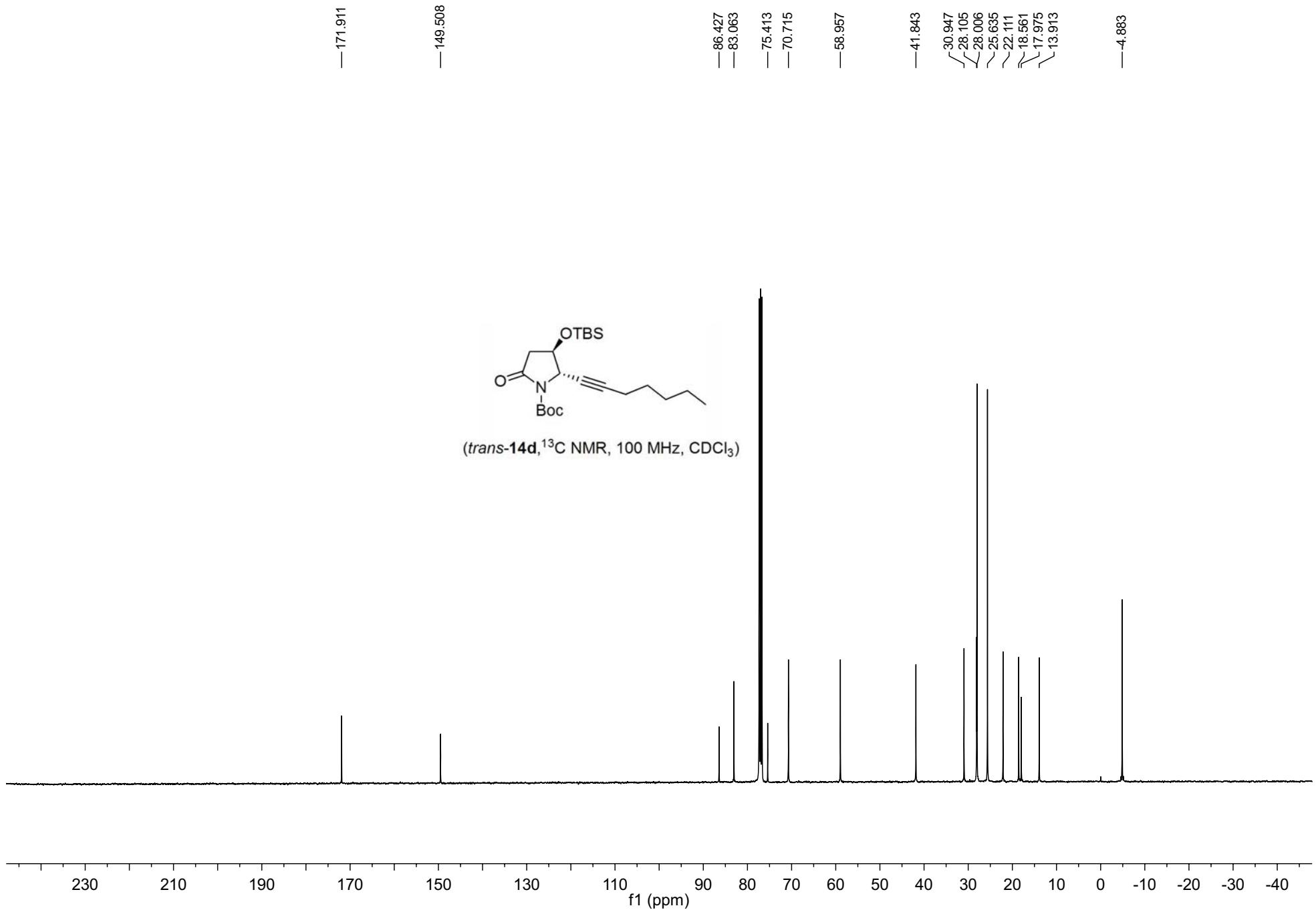
-109.881

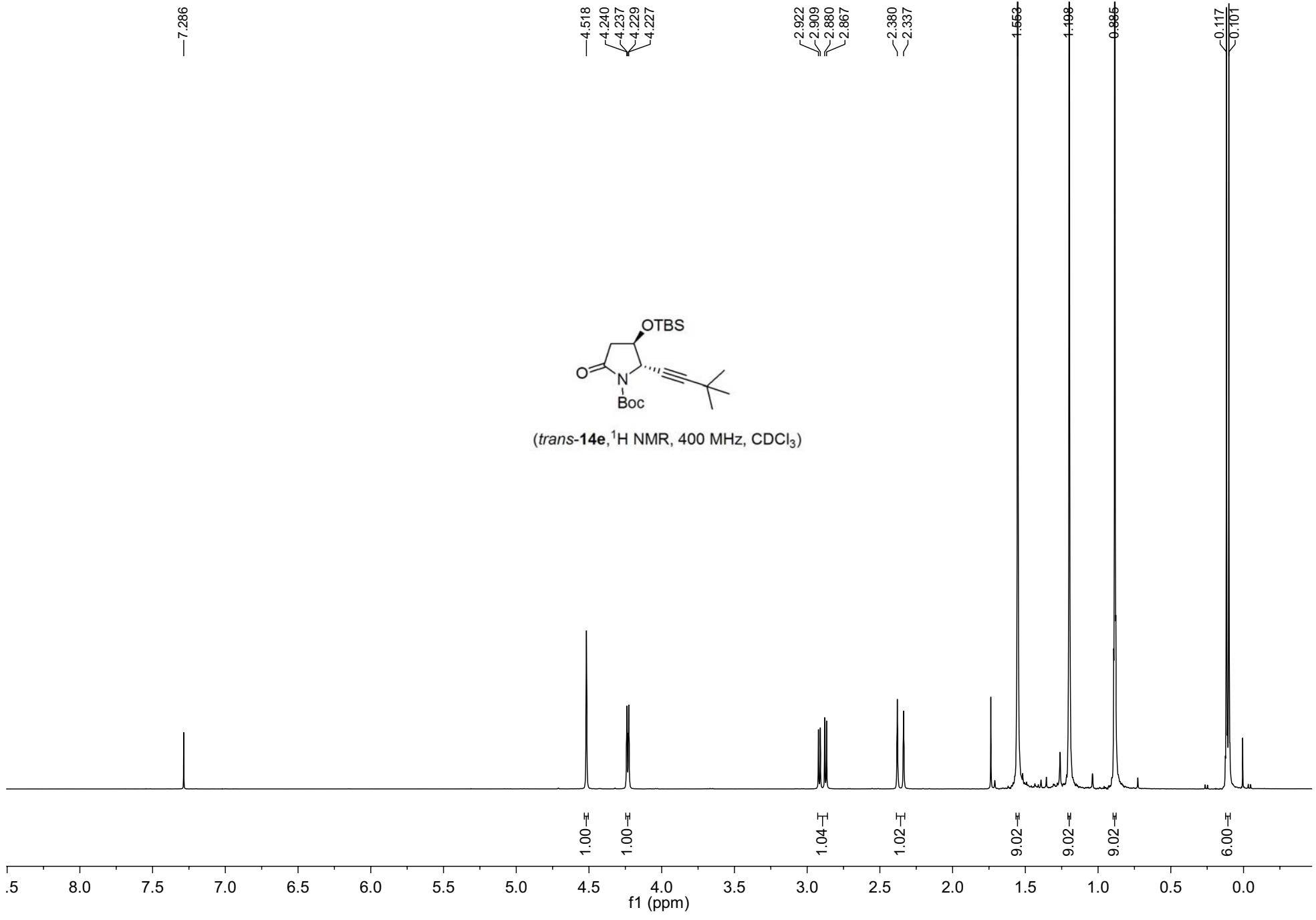


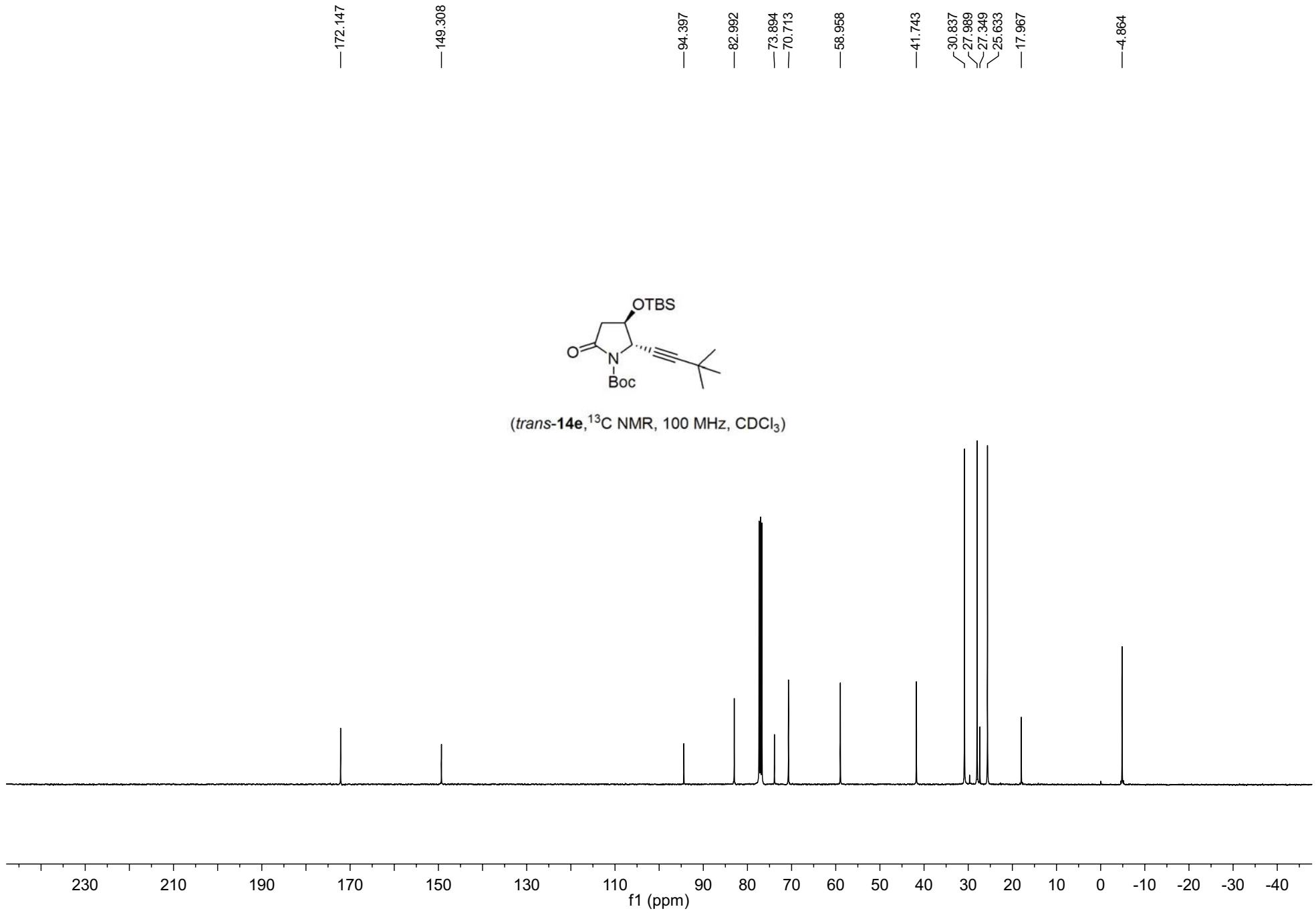
(*trans*-14c,  $^{19}\text{F}$  NMR, 376 MHz,  $\text{CDCl}_3$ )

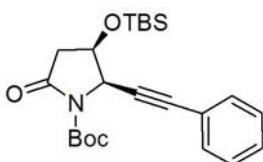
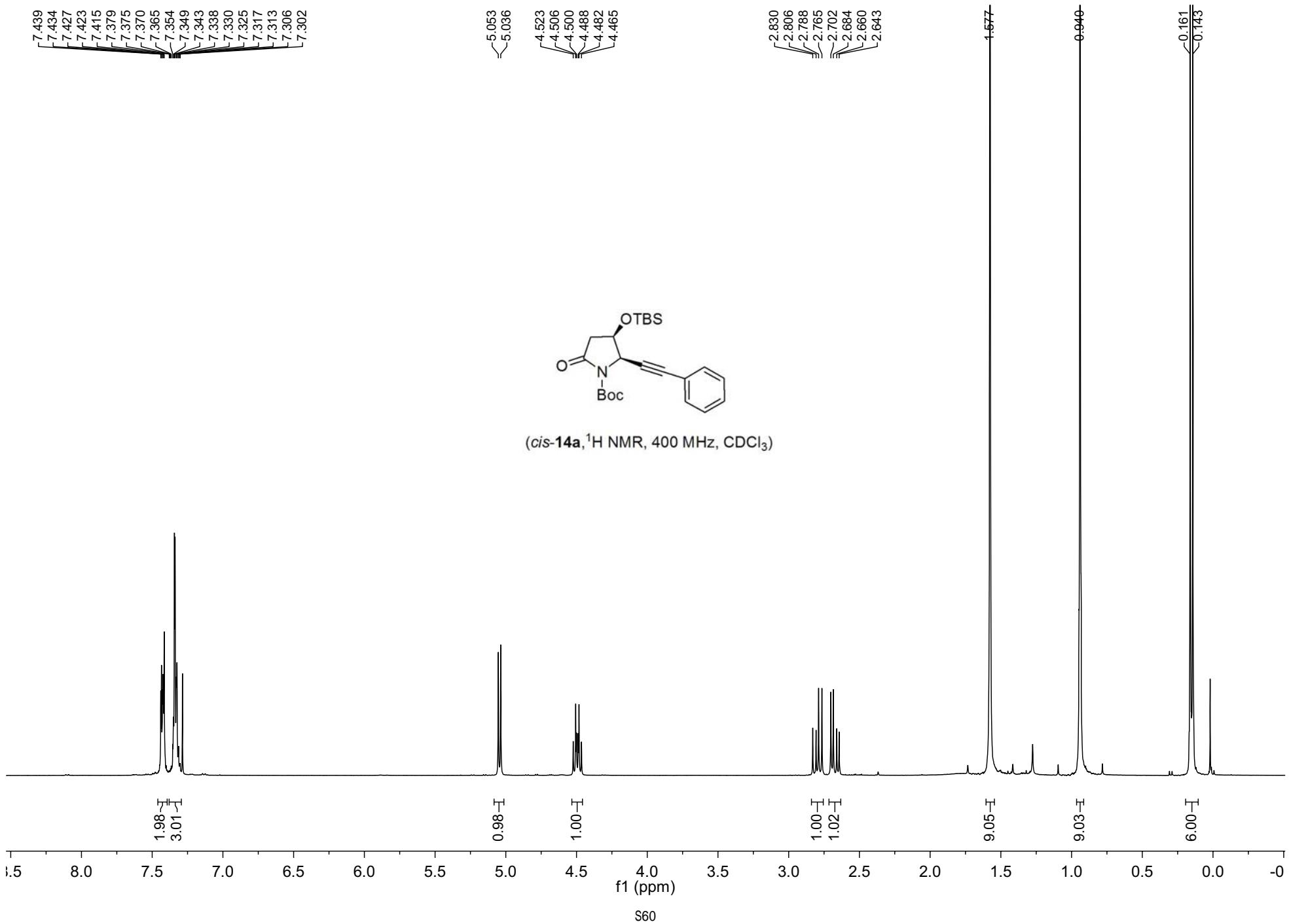


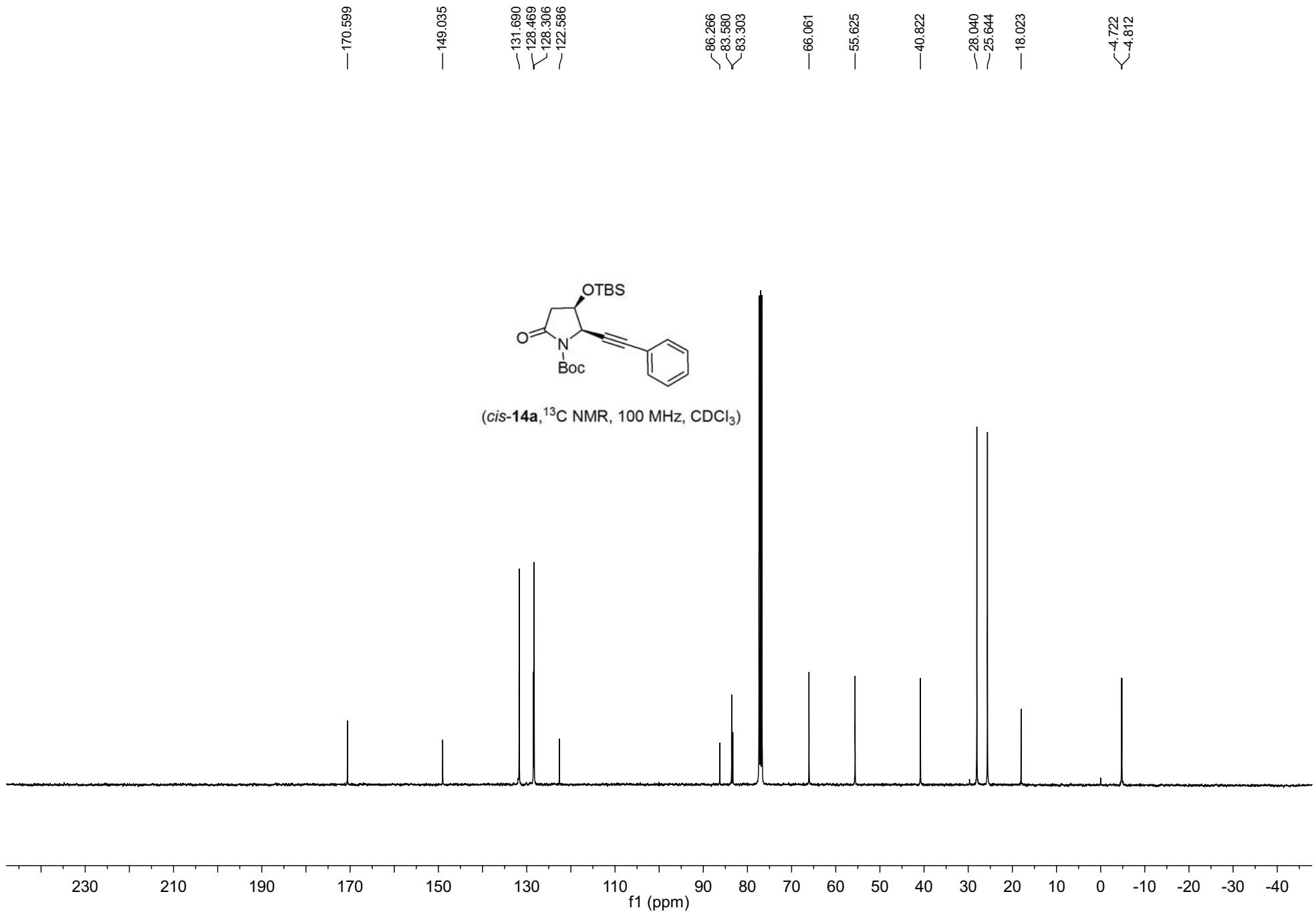


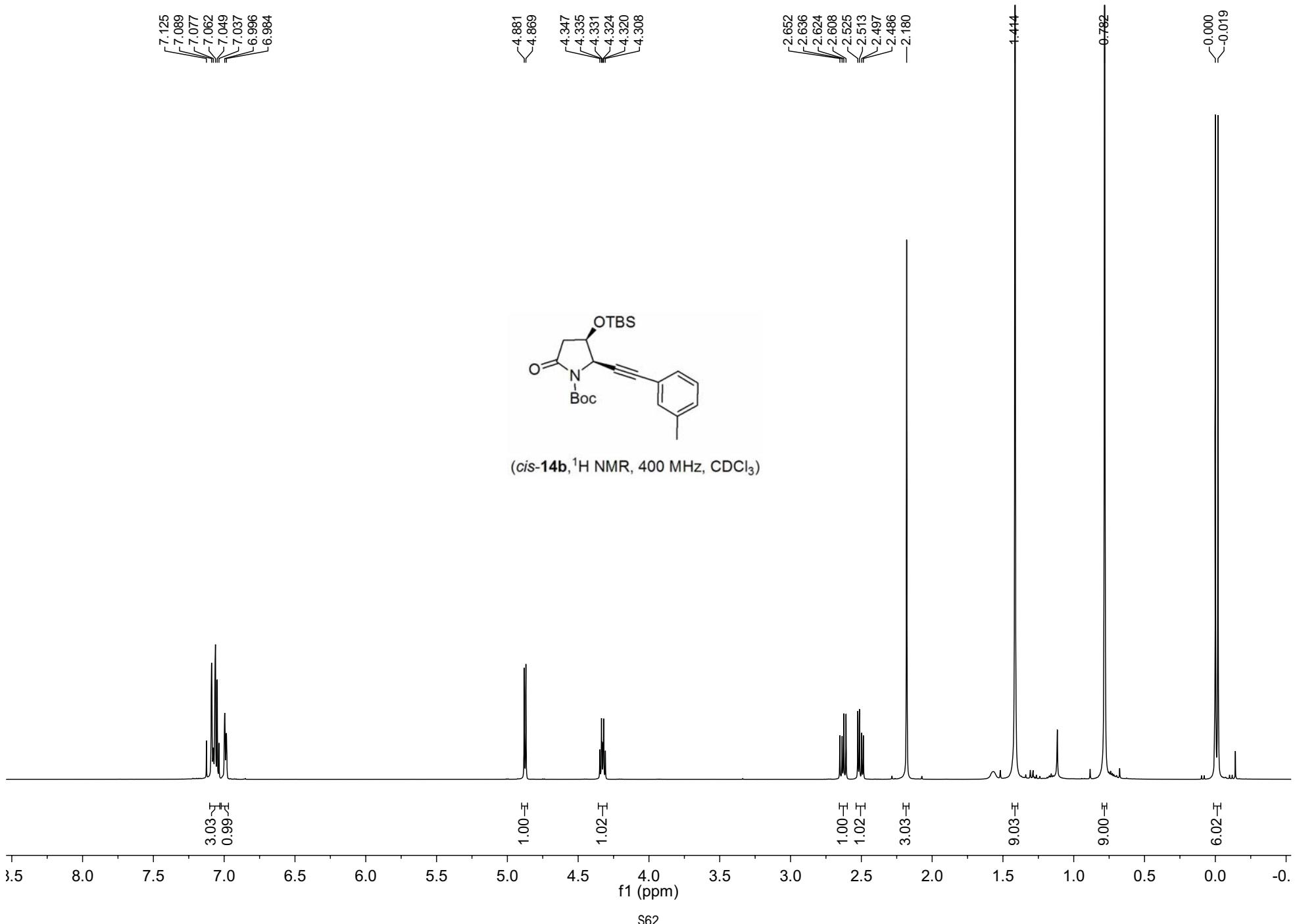


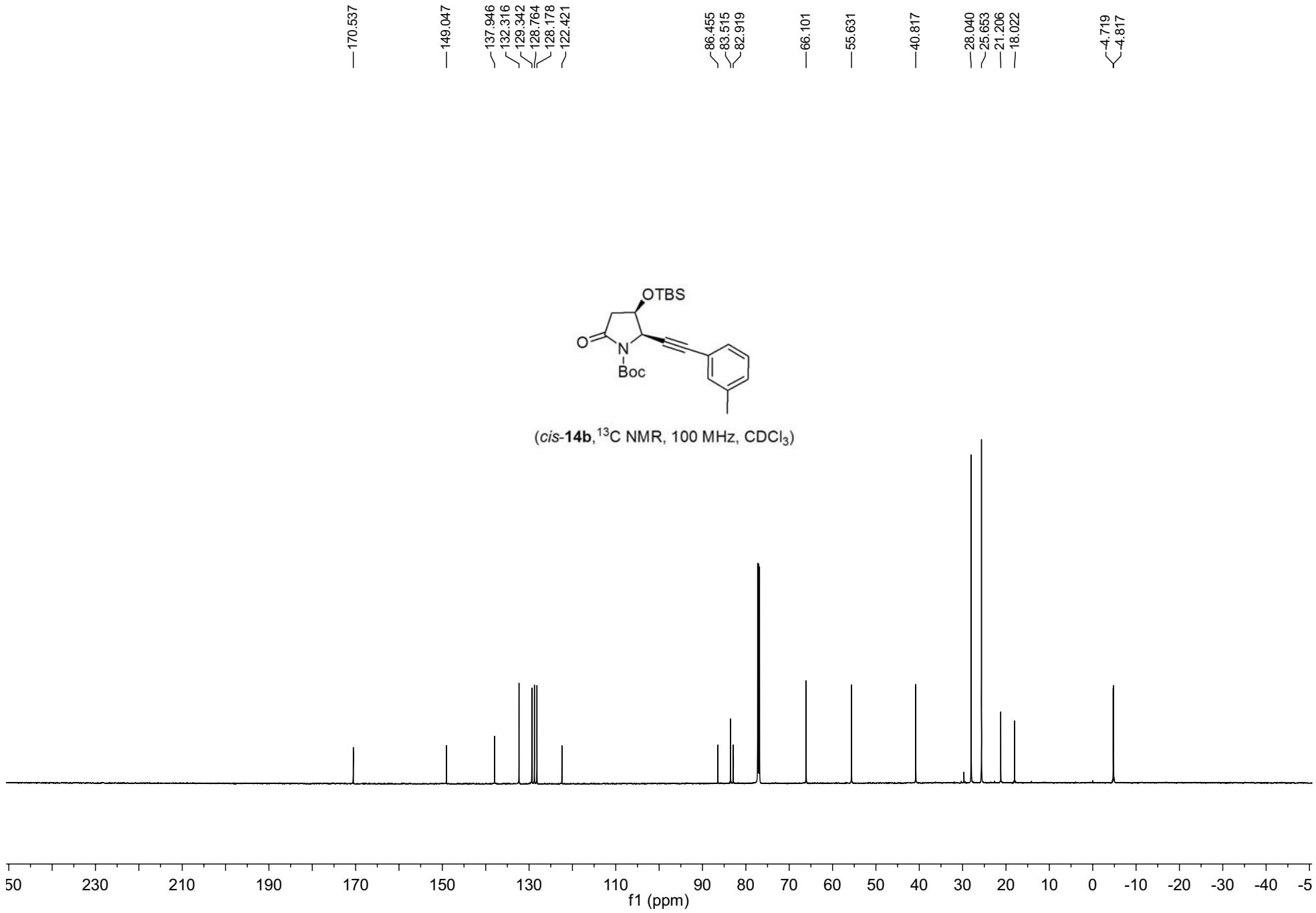


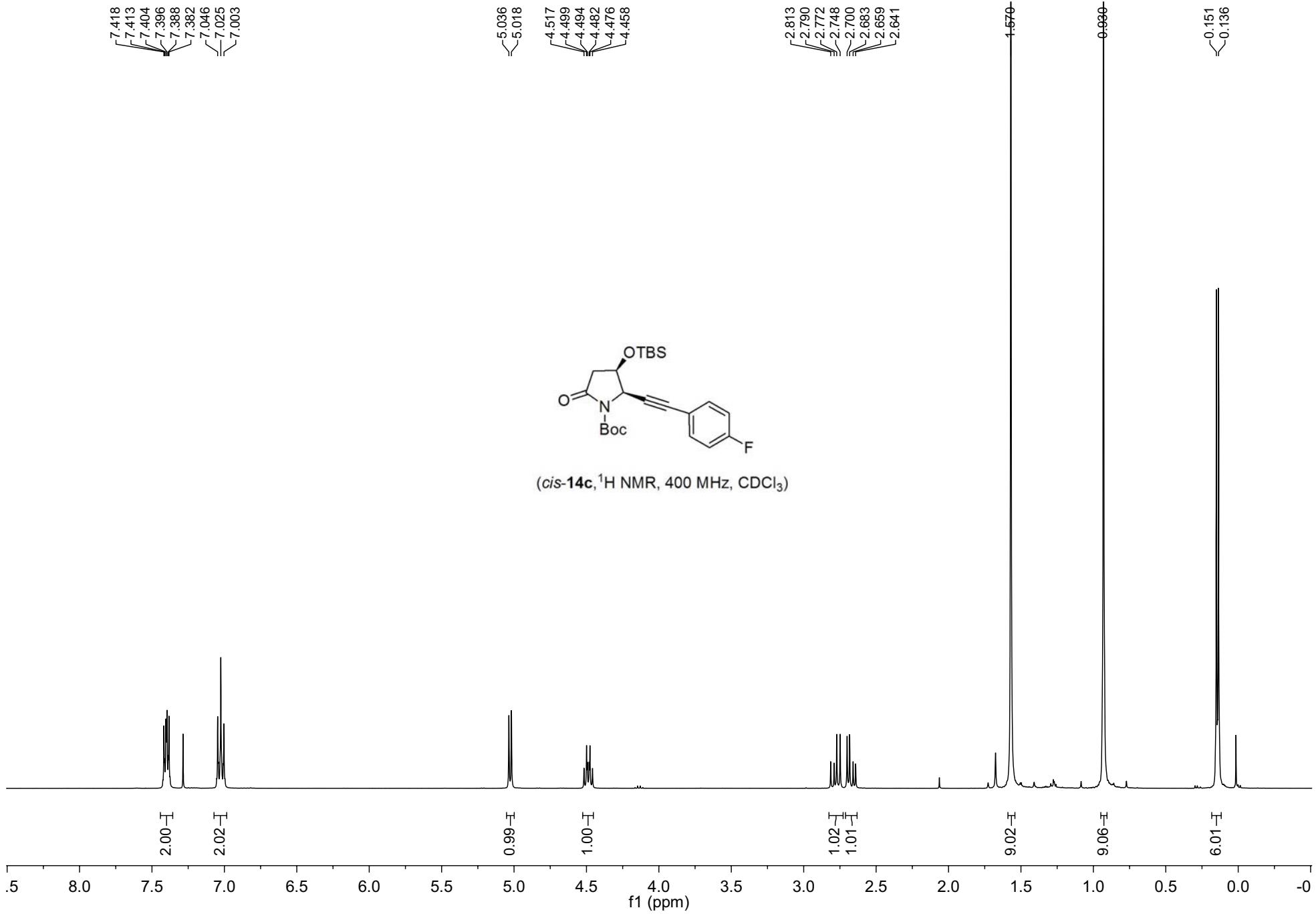


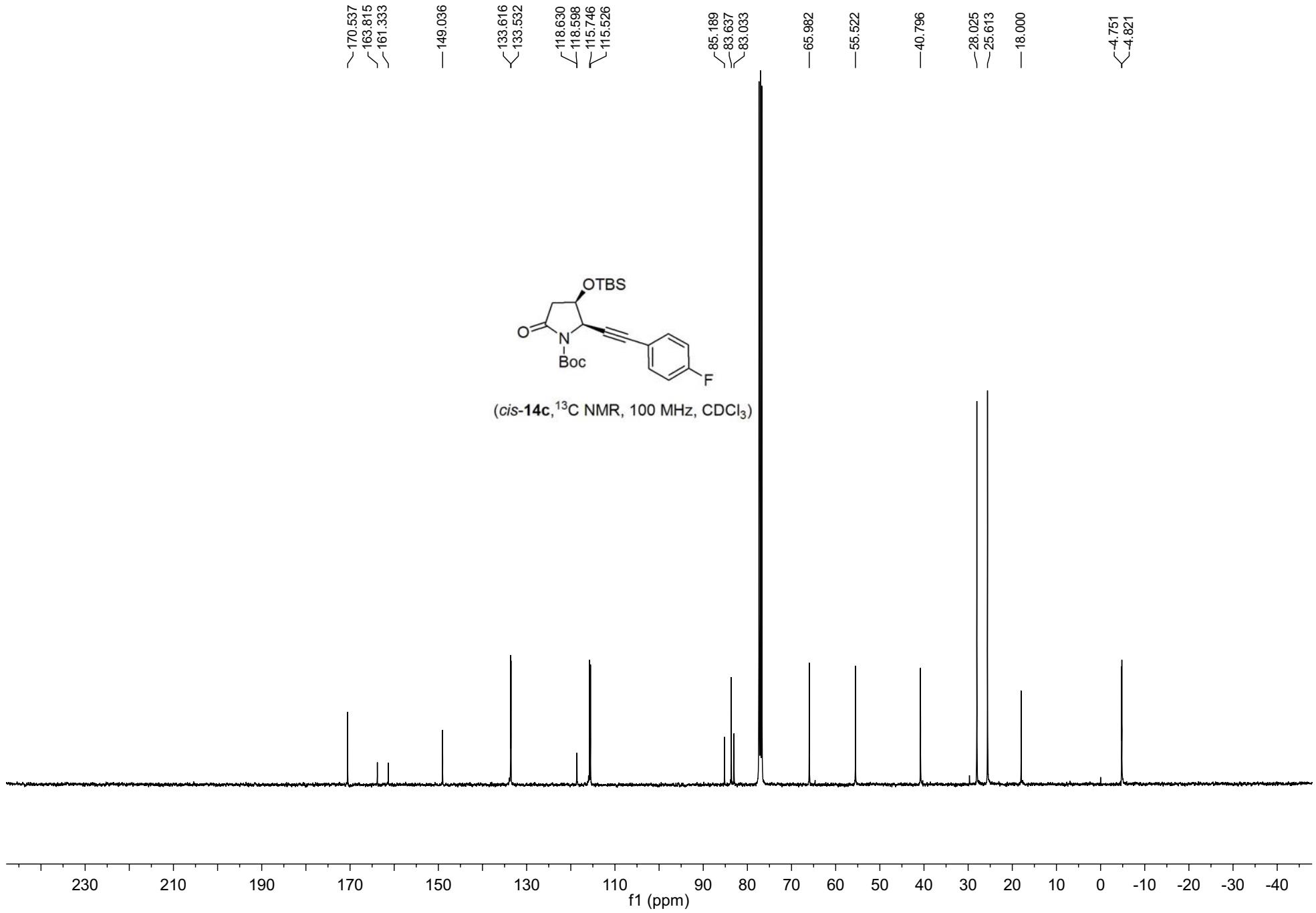


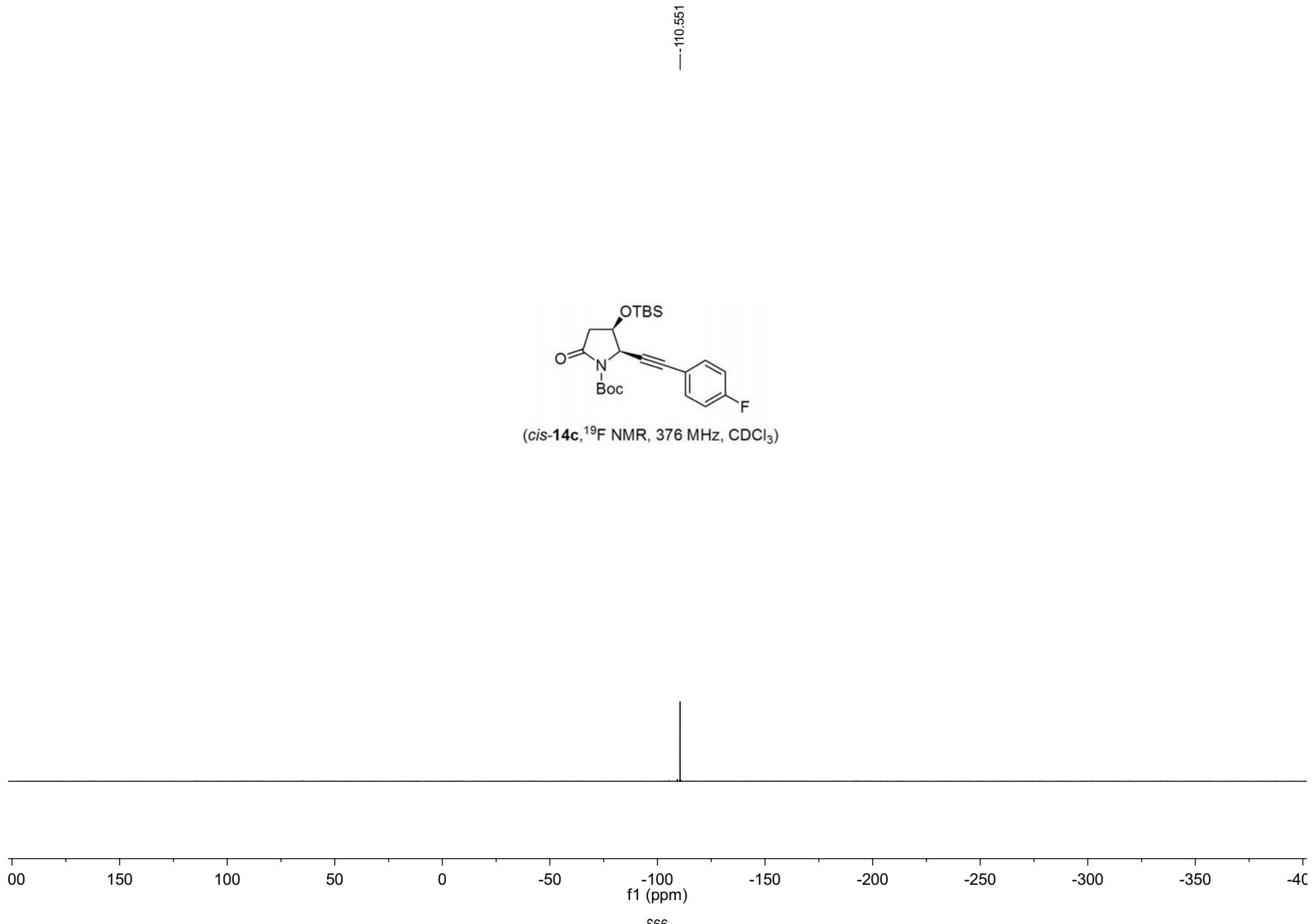


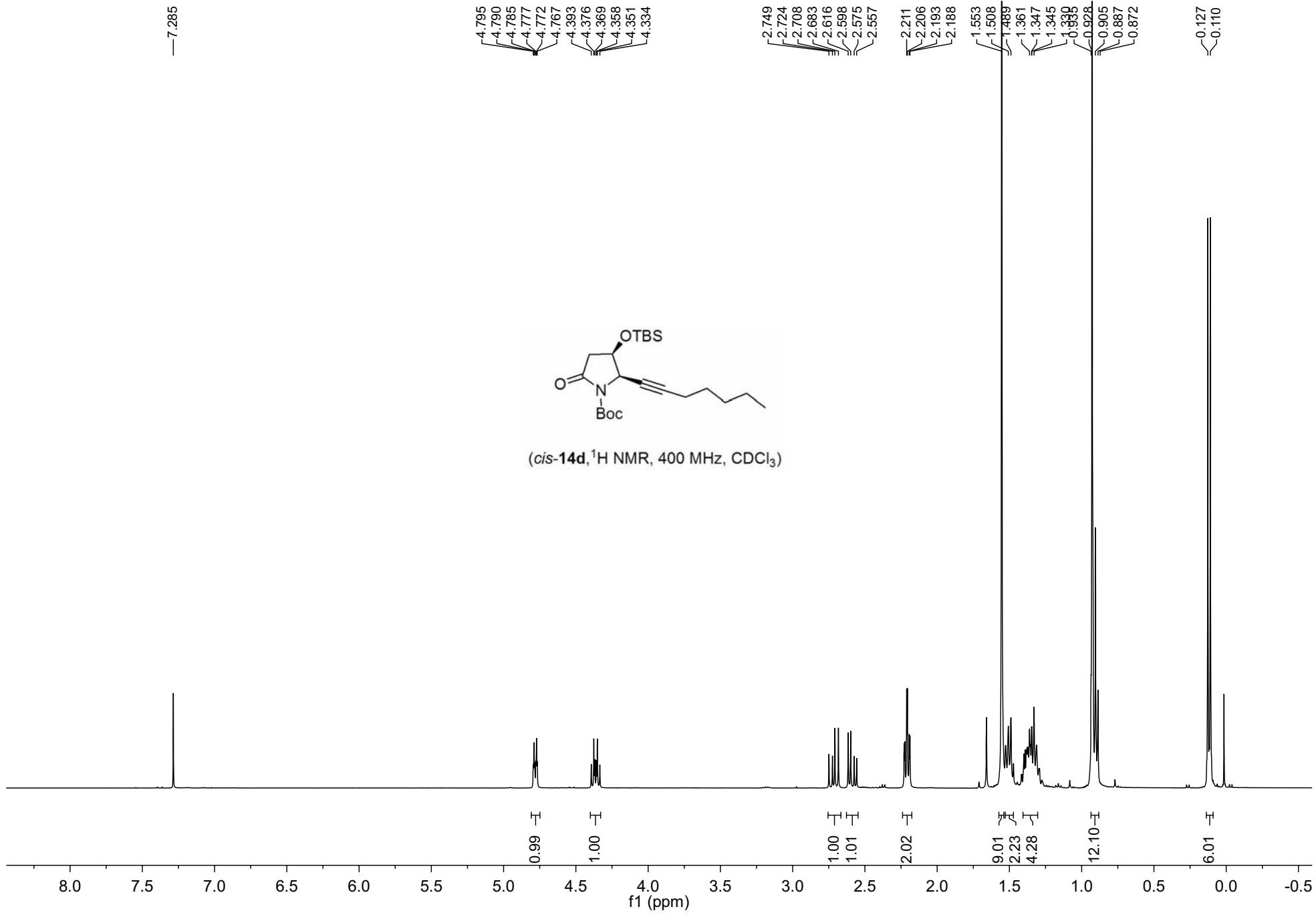


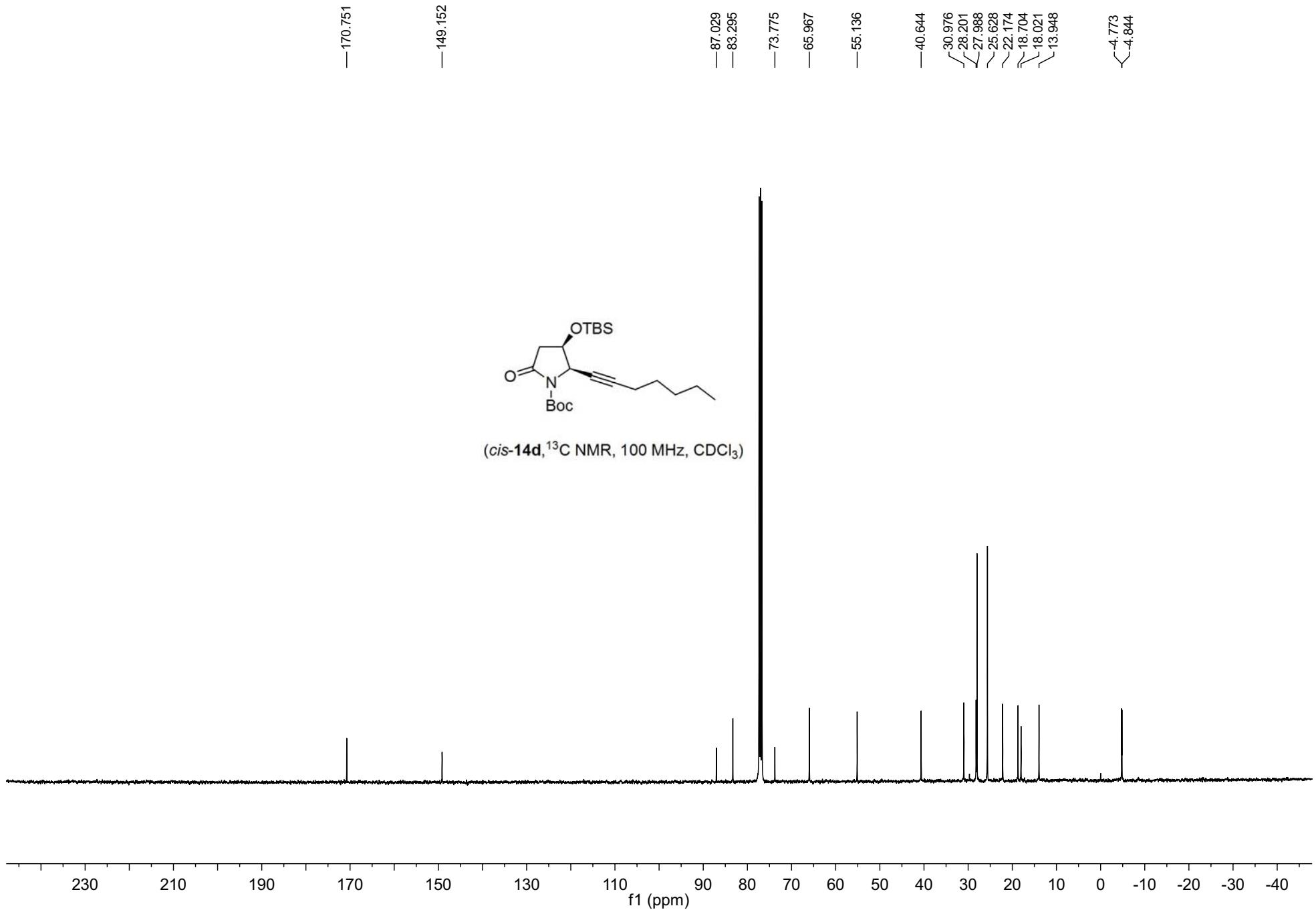


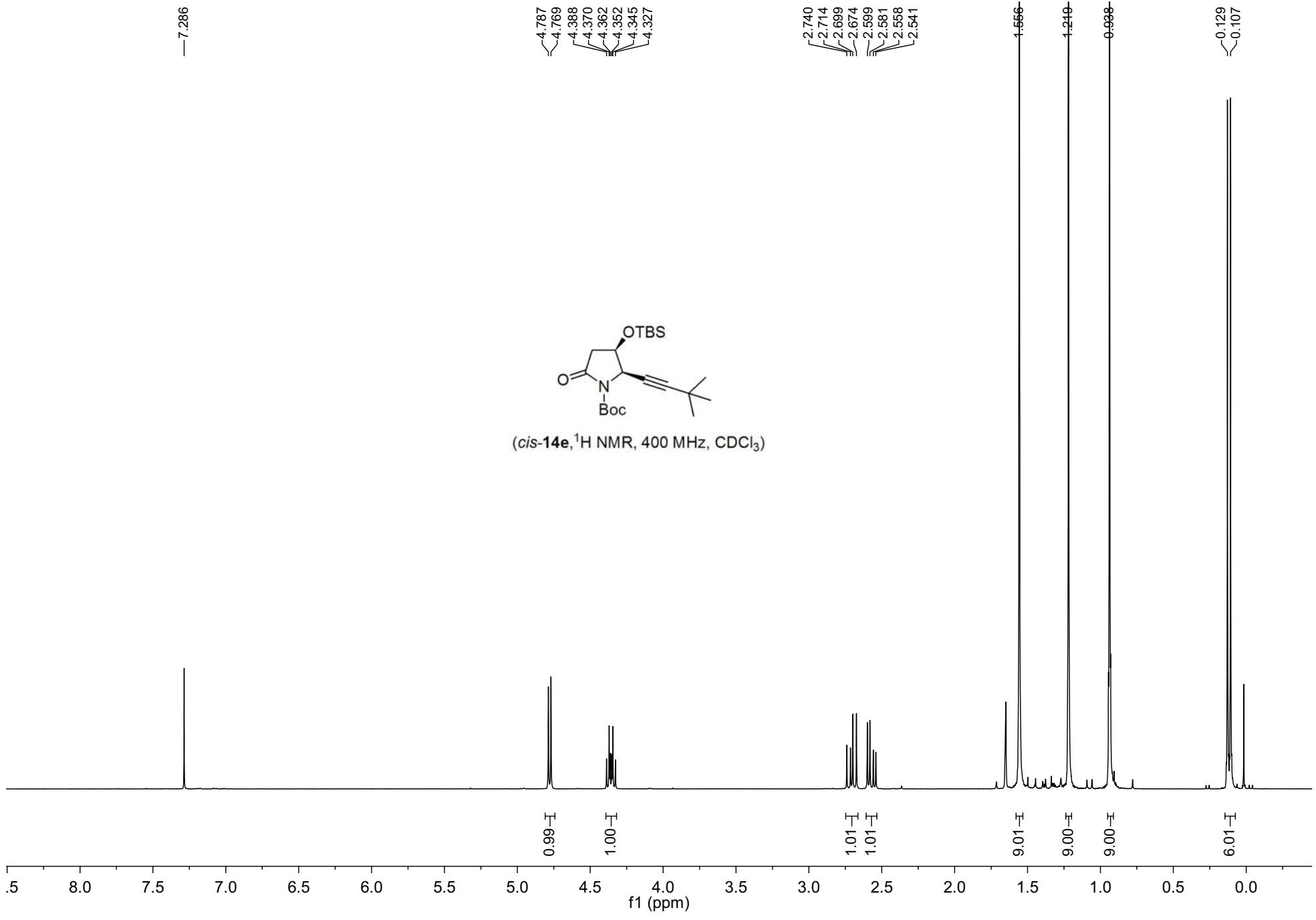


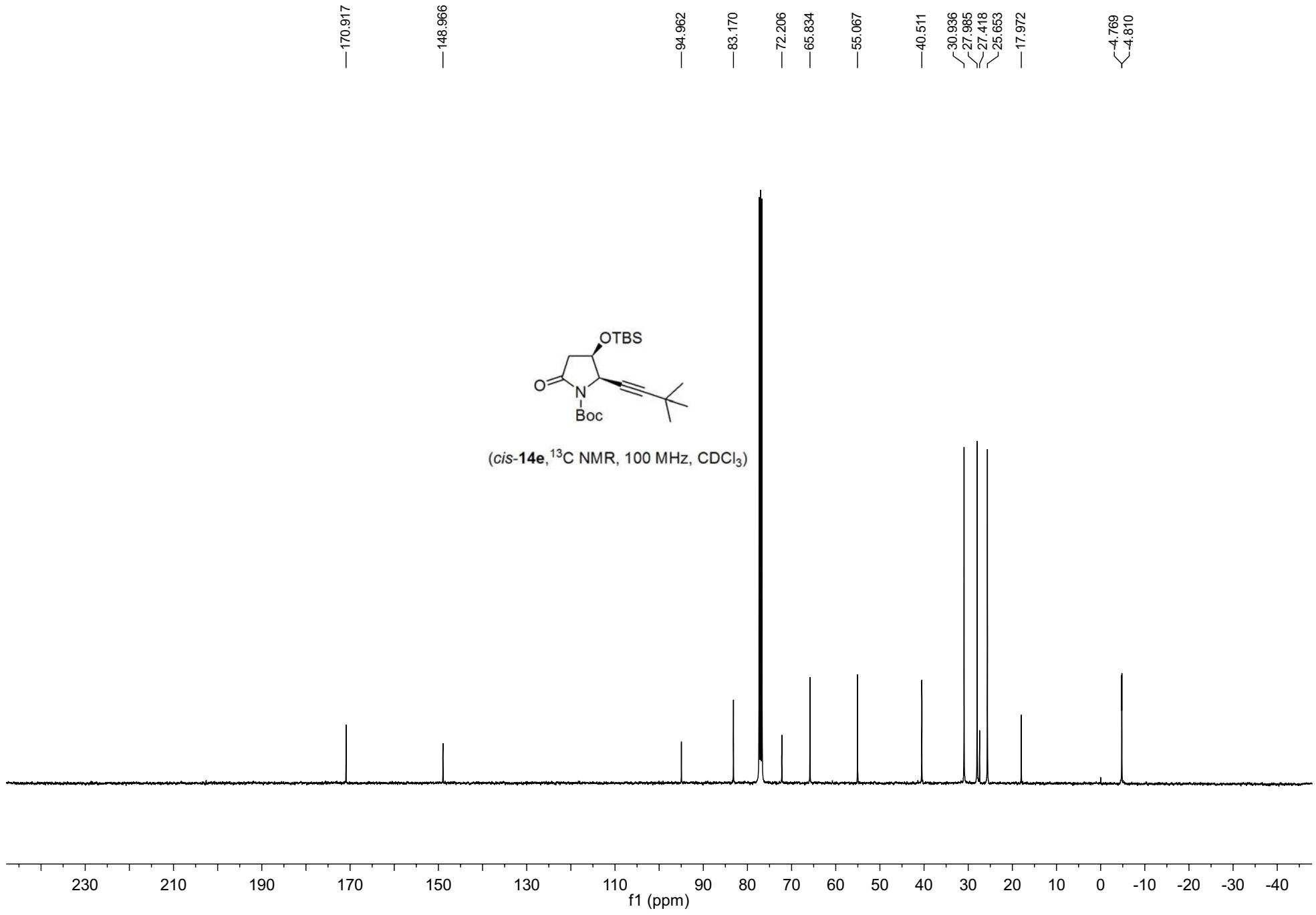


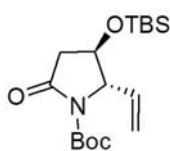
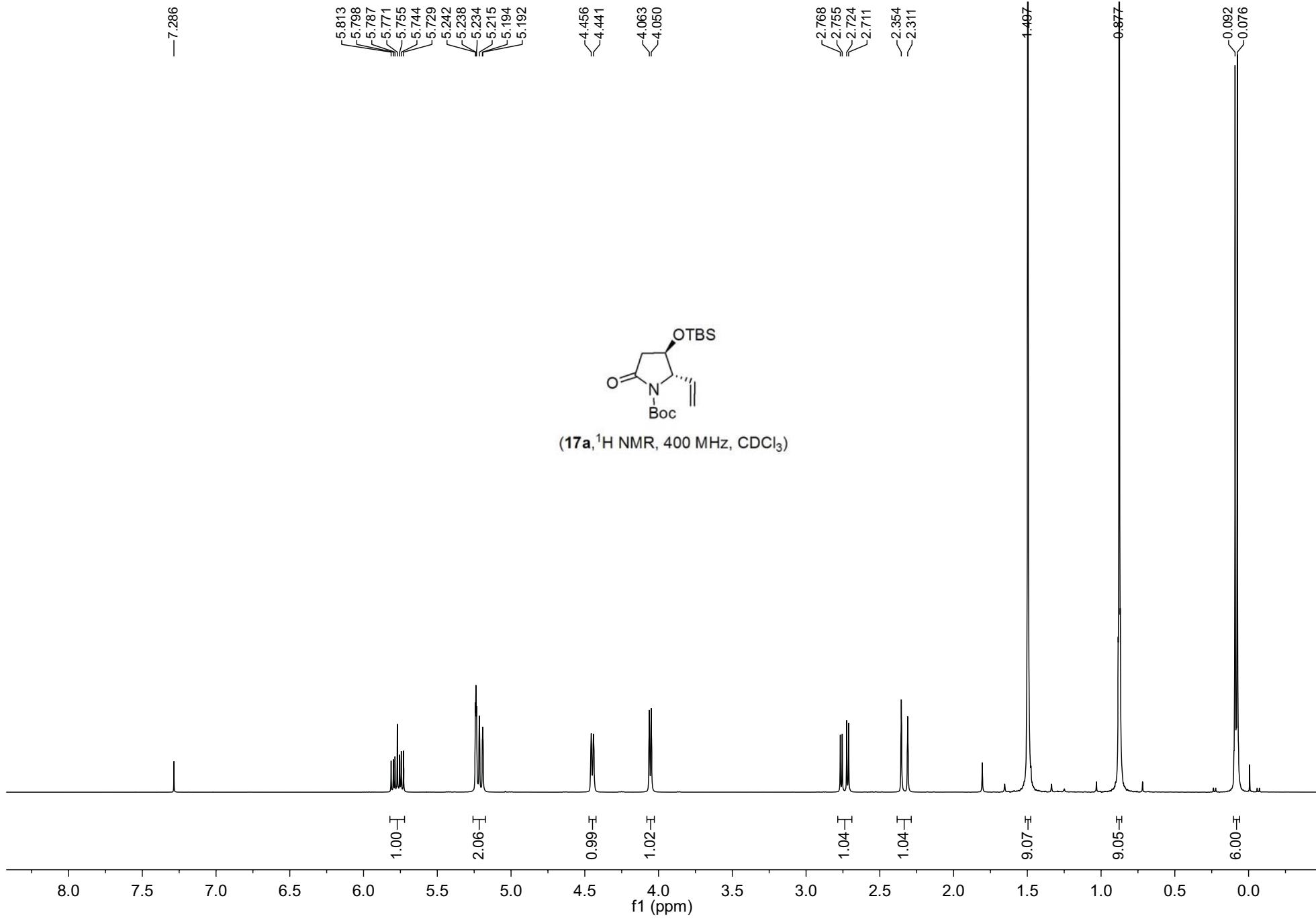


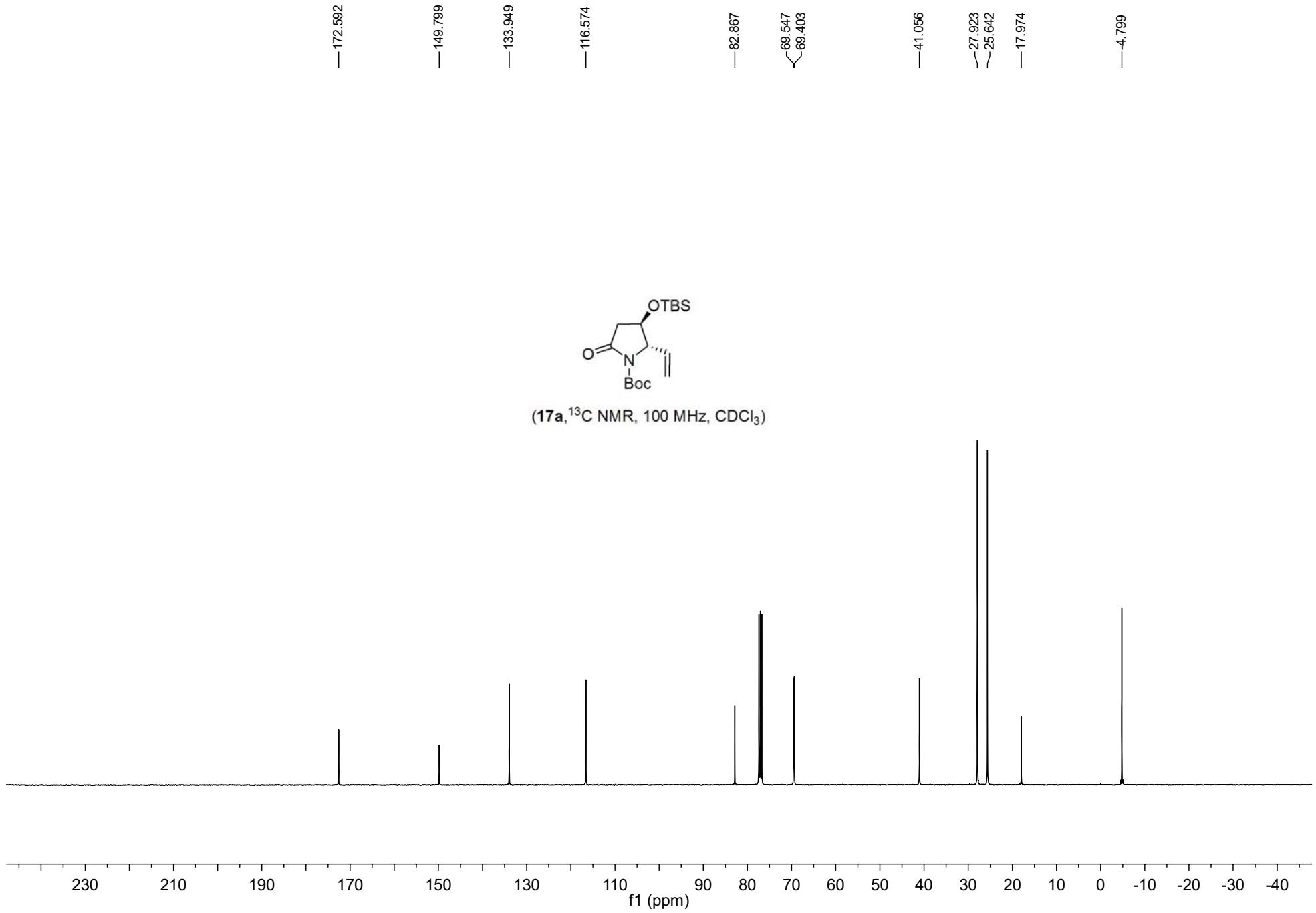


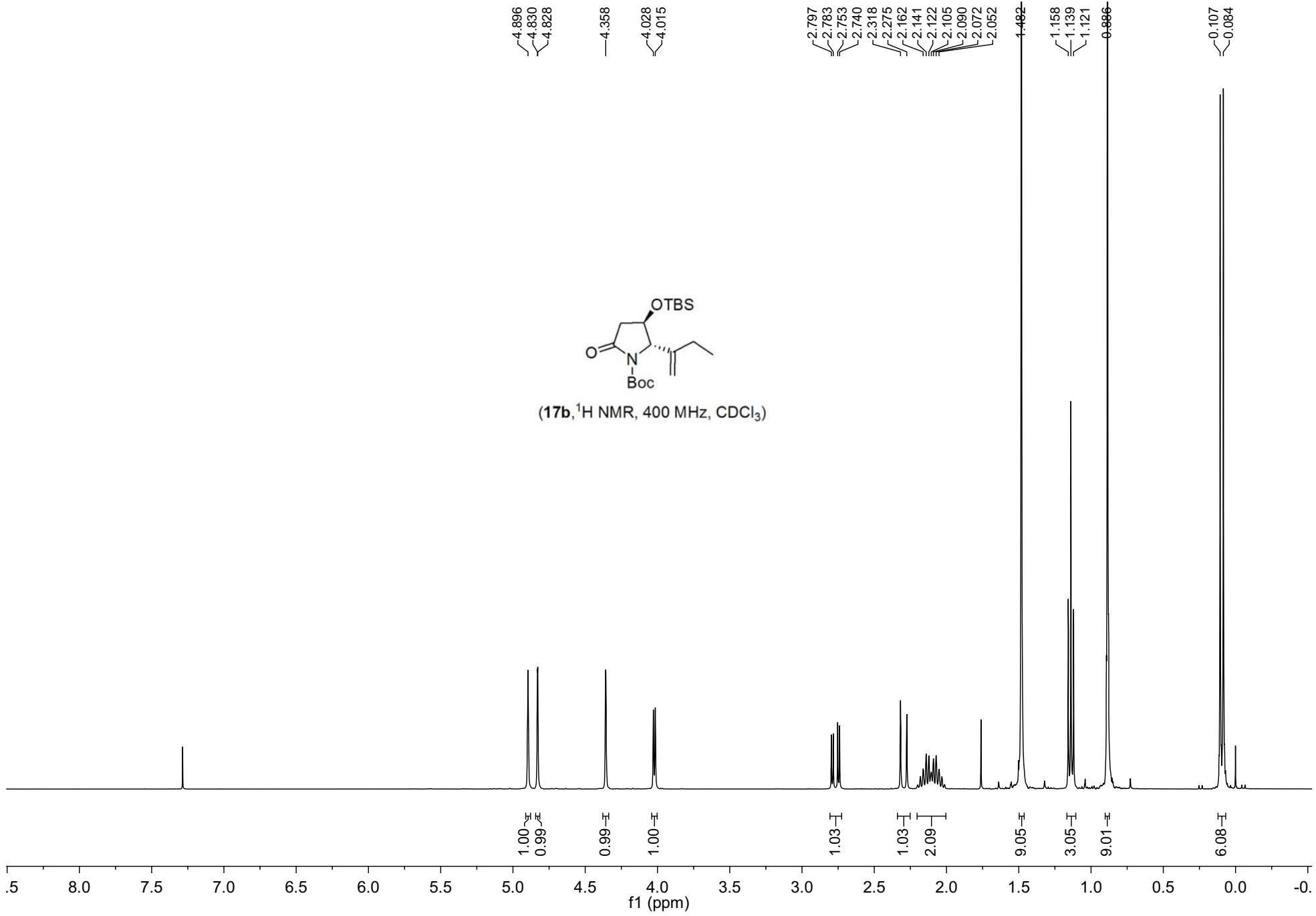


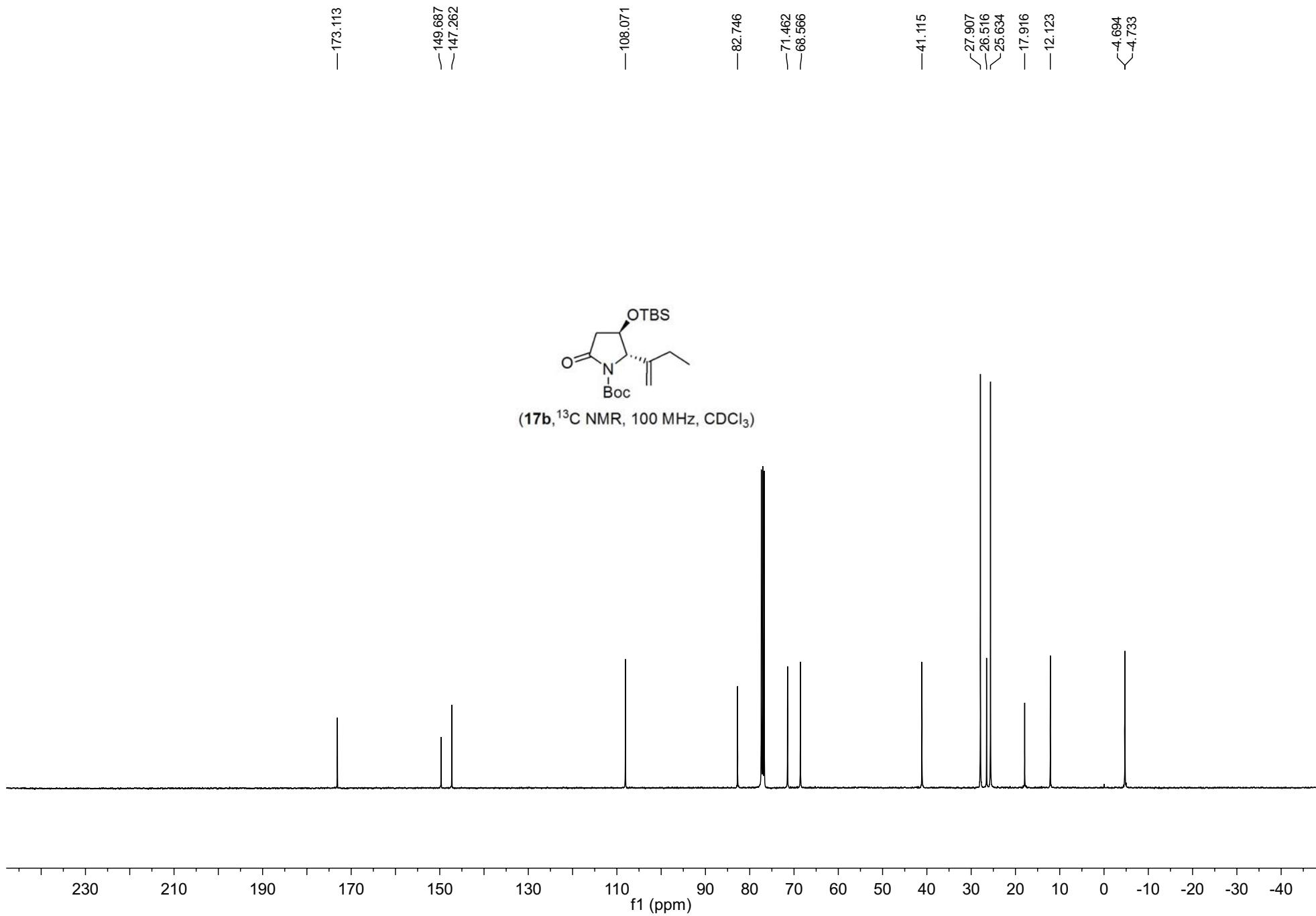


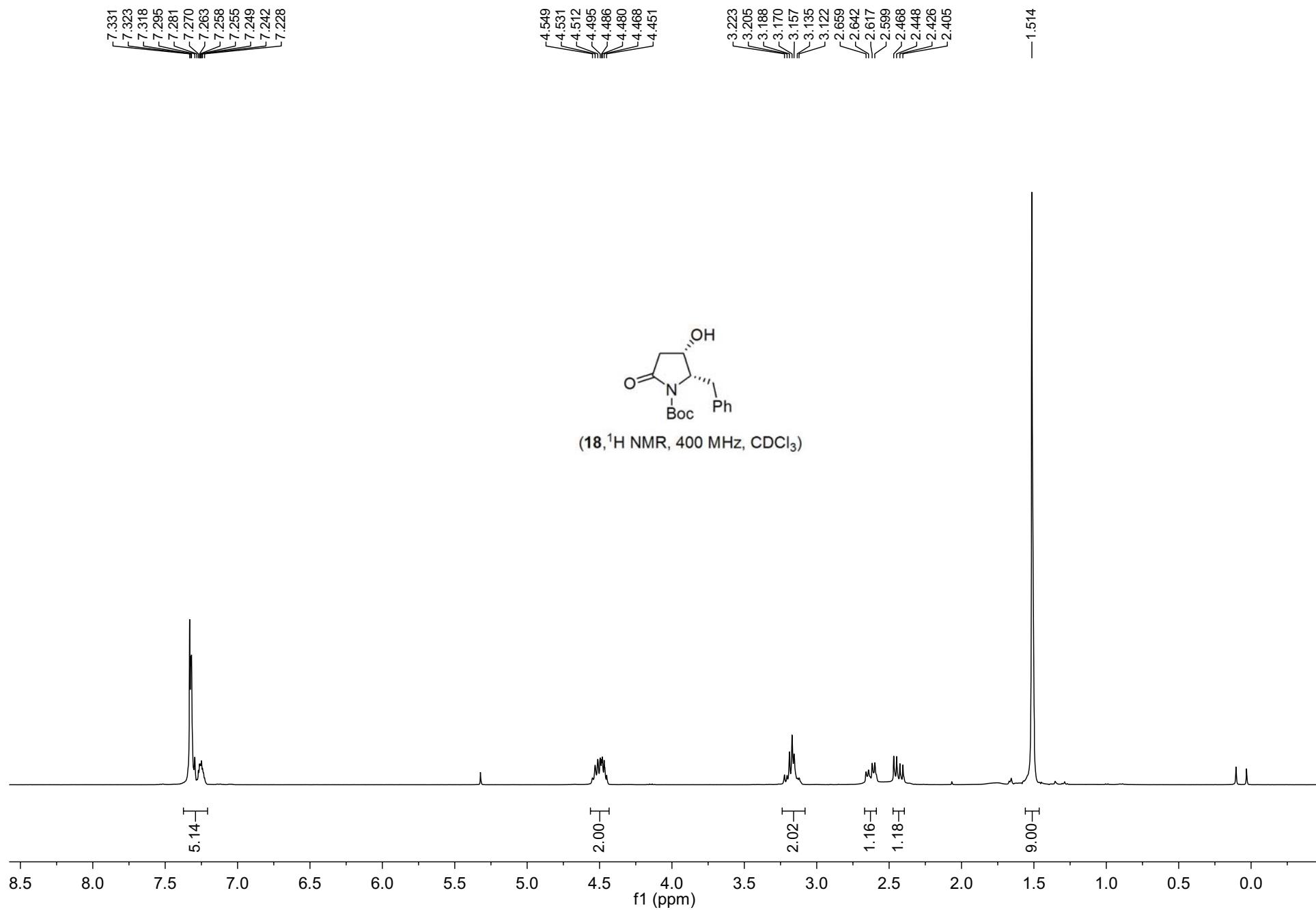


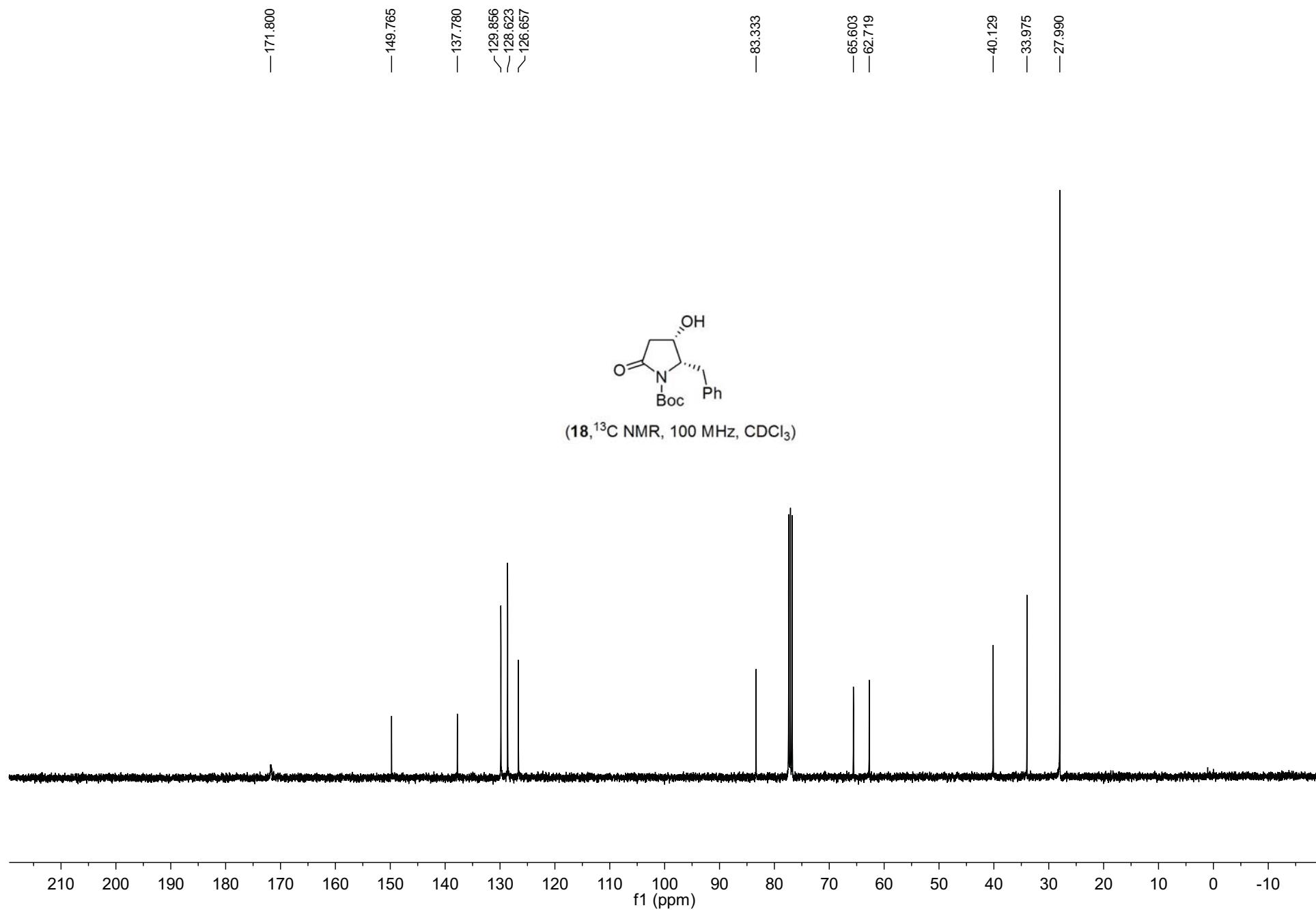


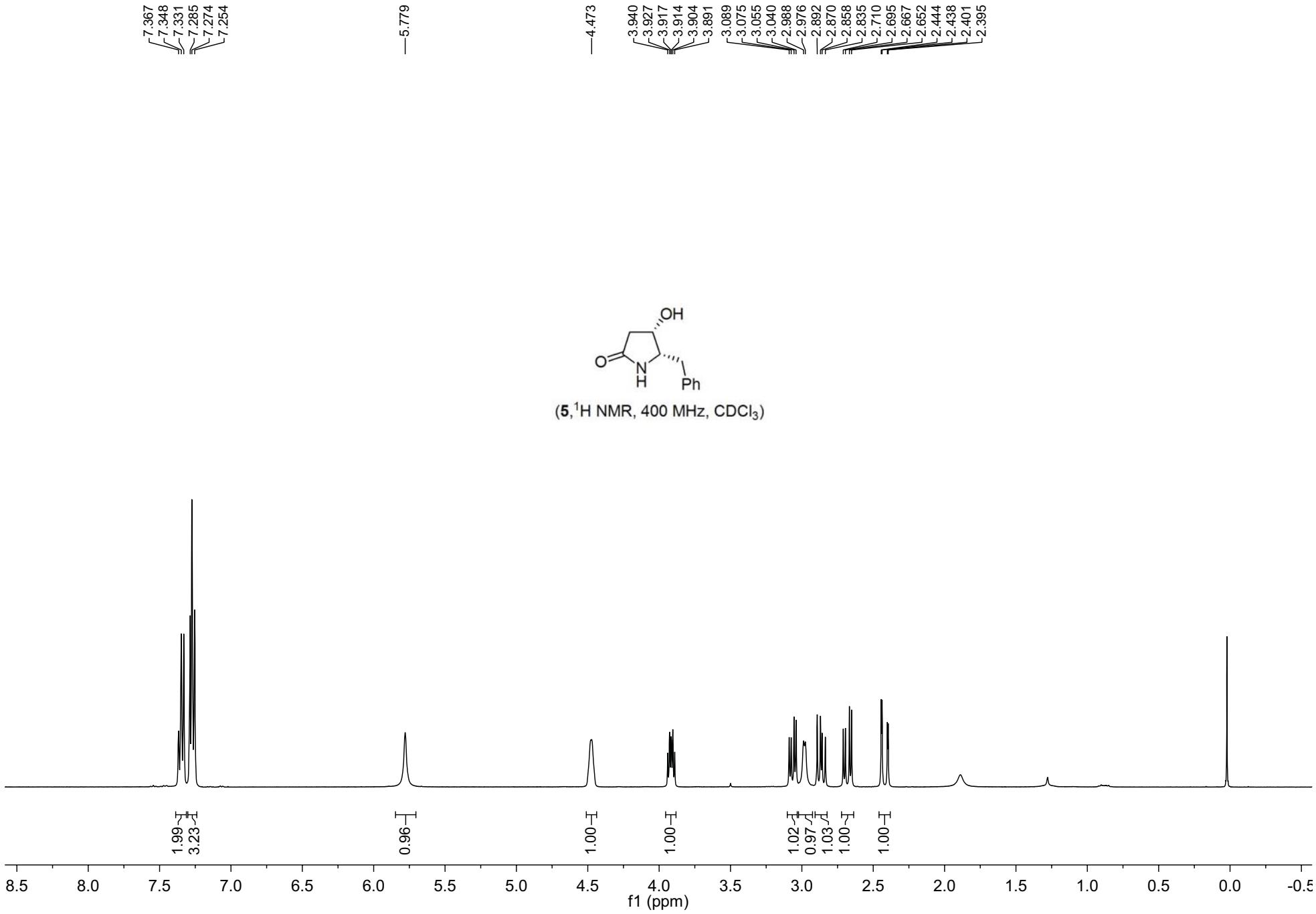


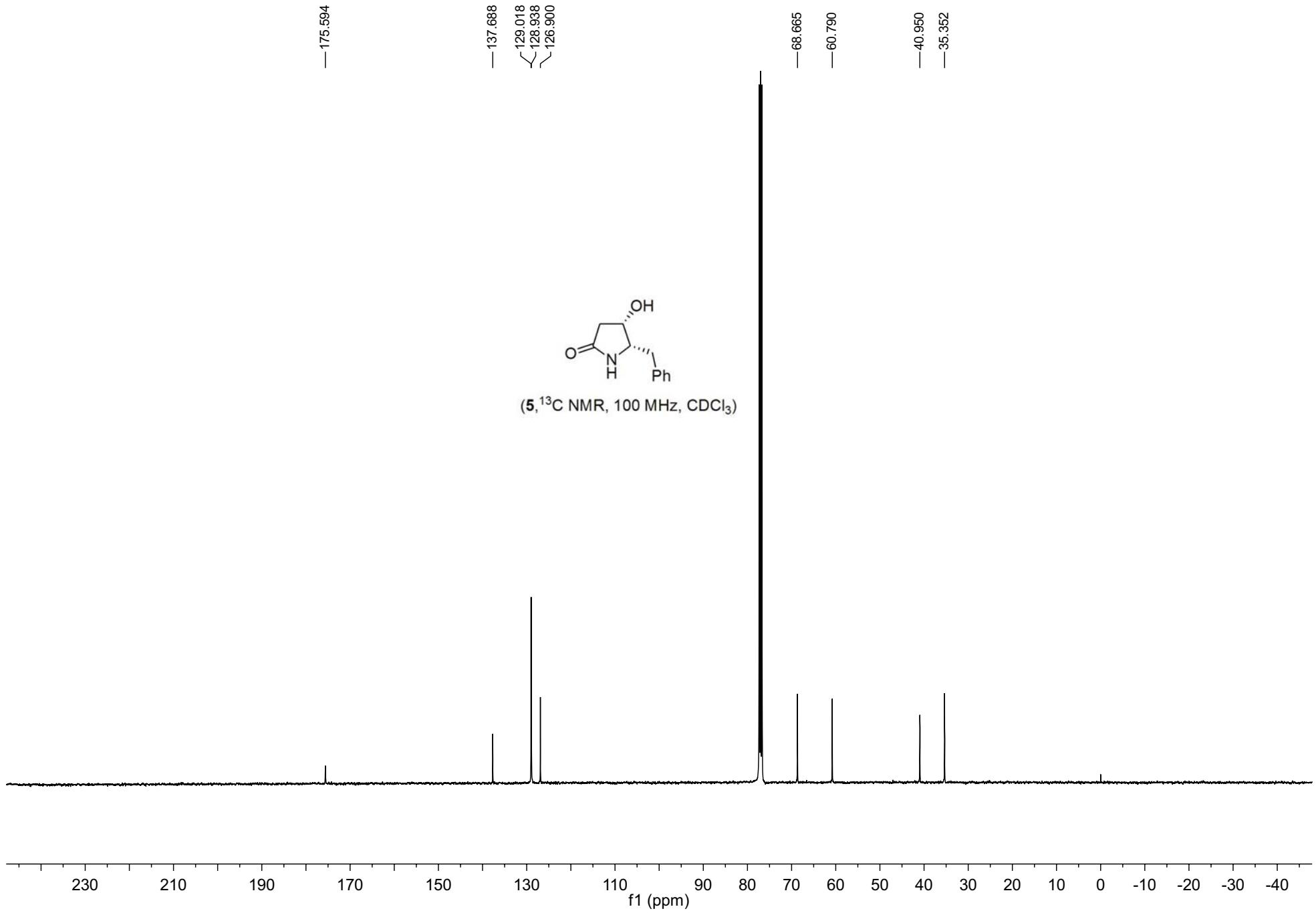


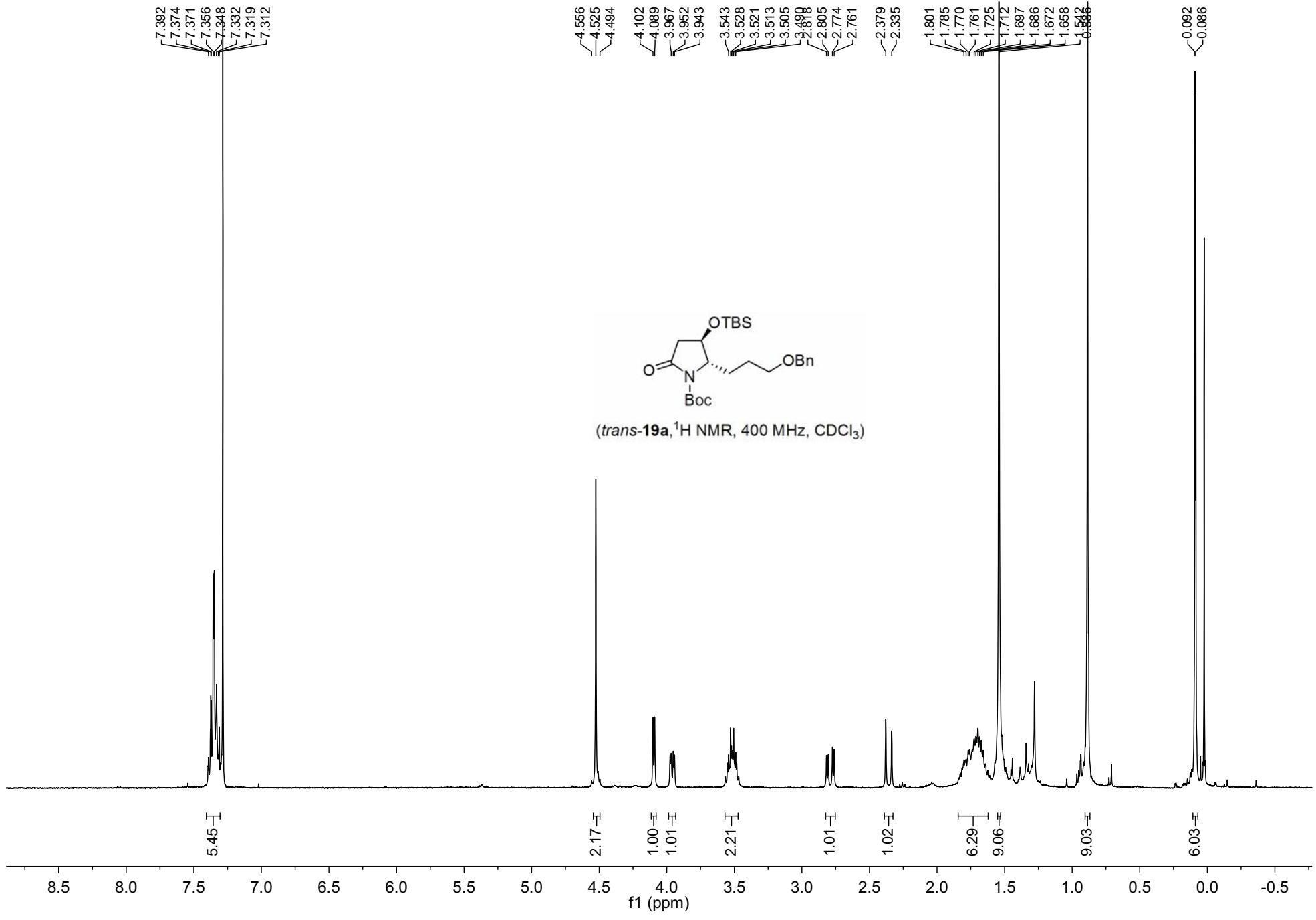




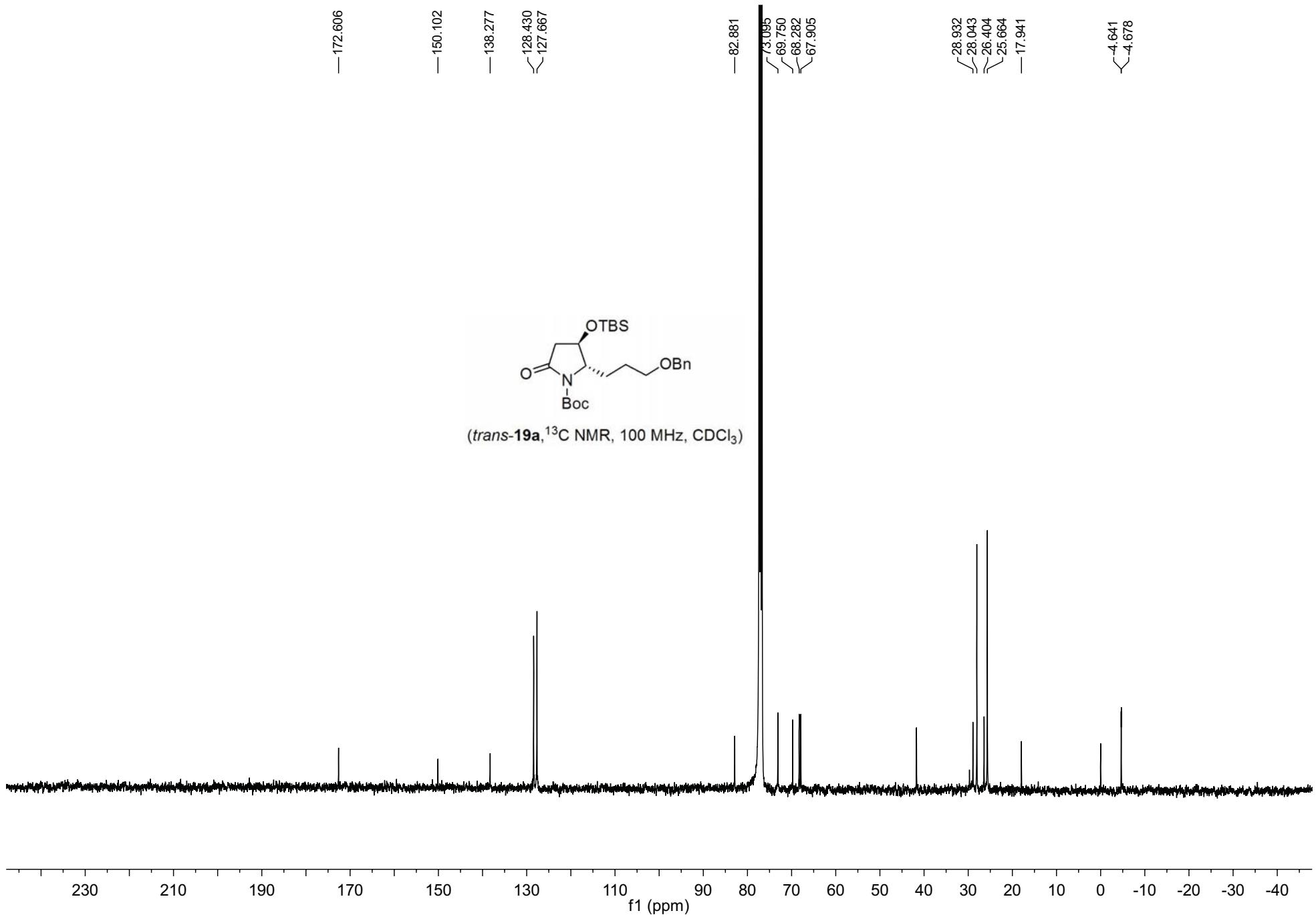


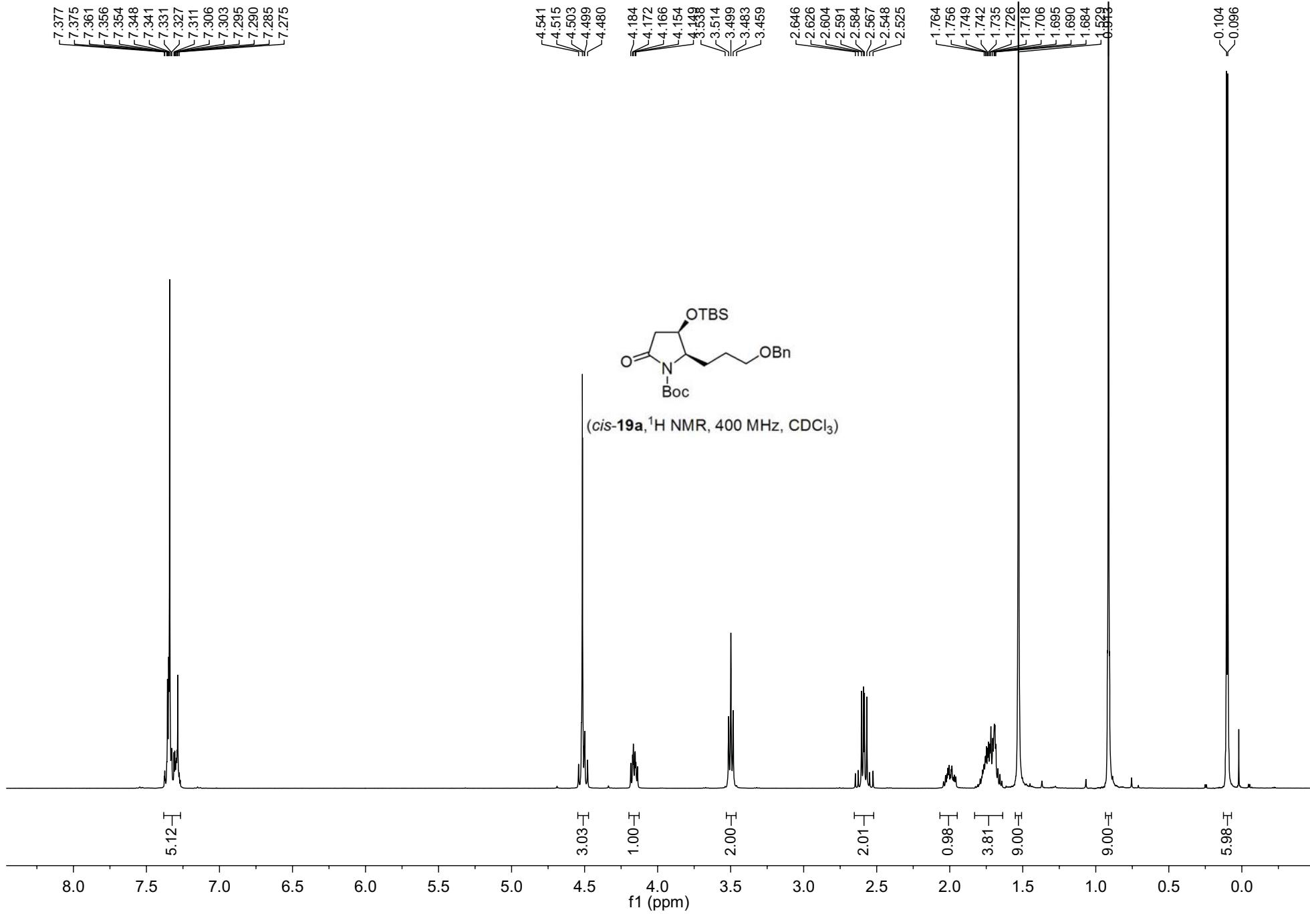


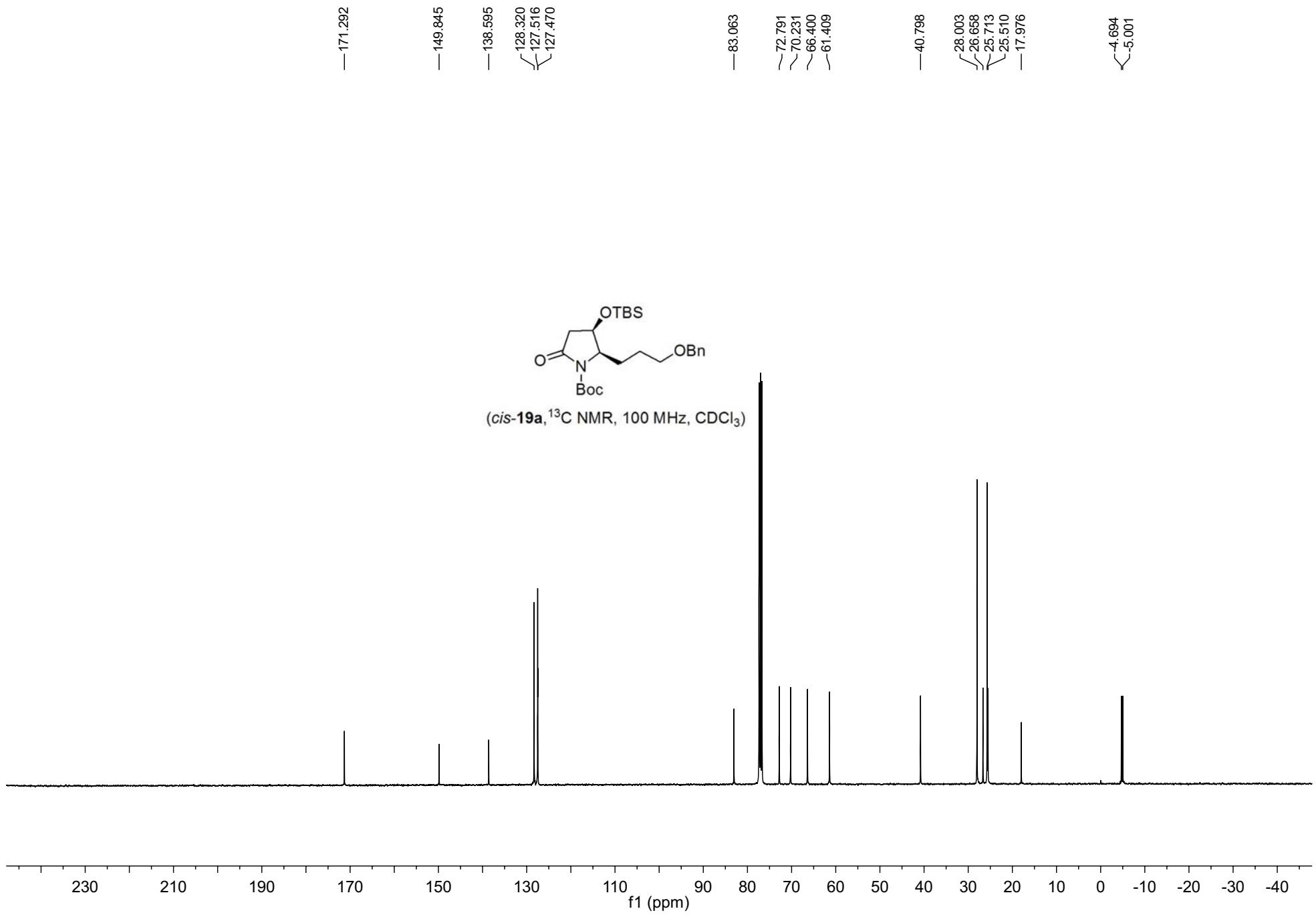


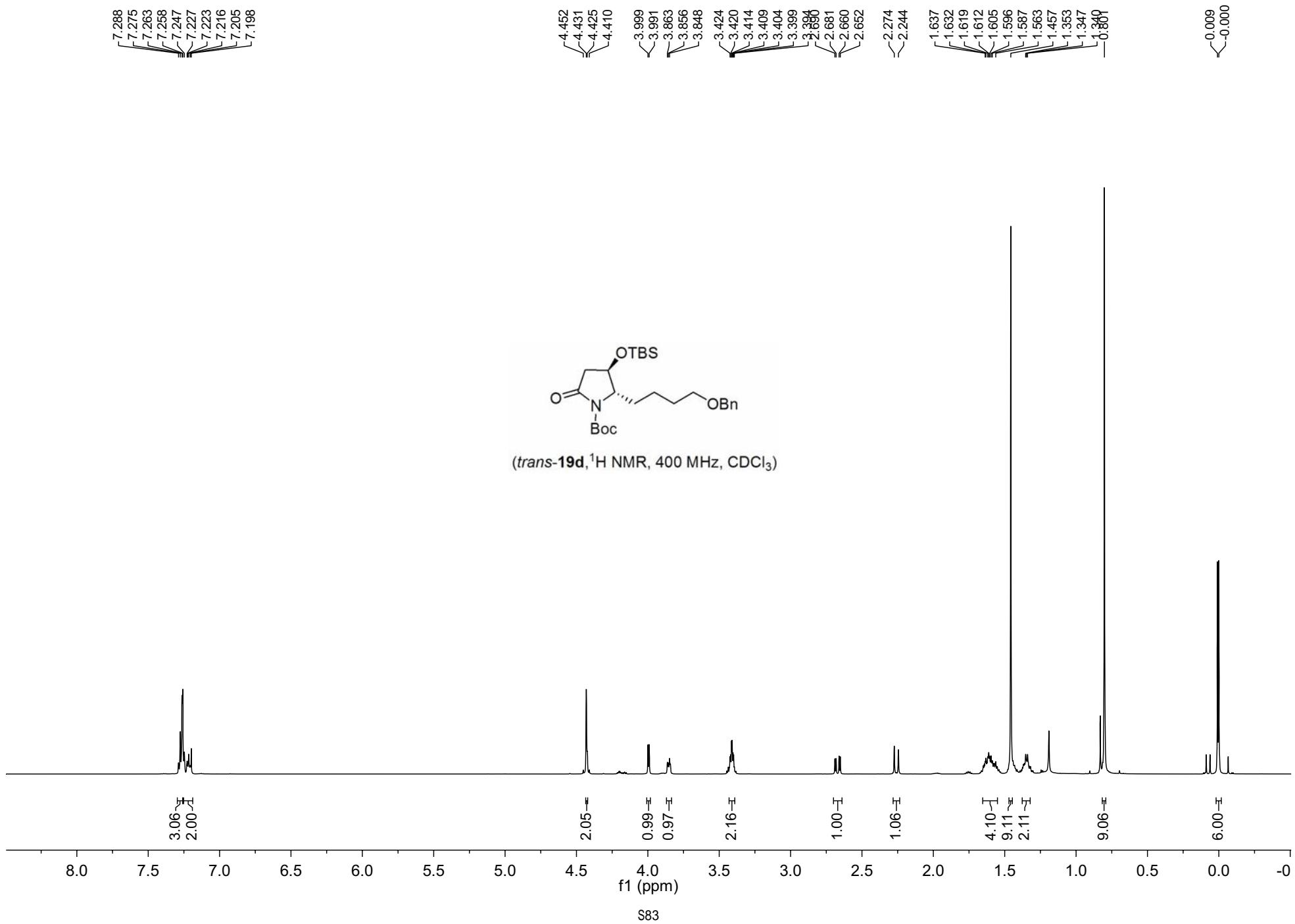


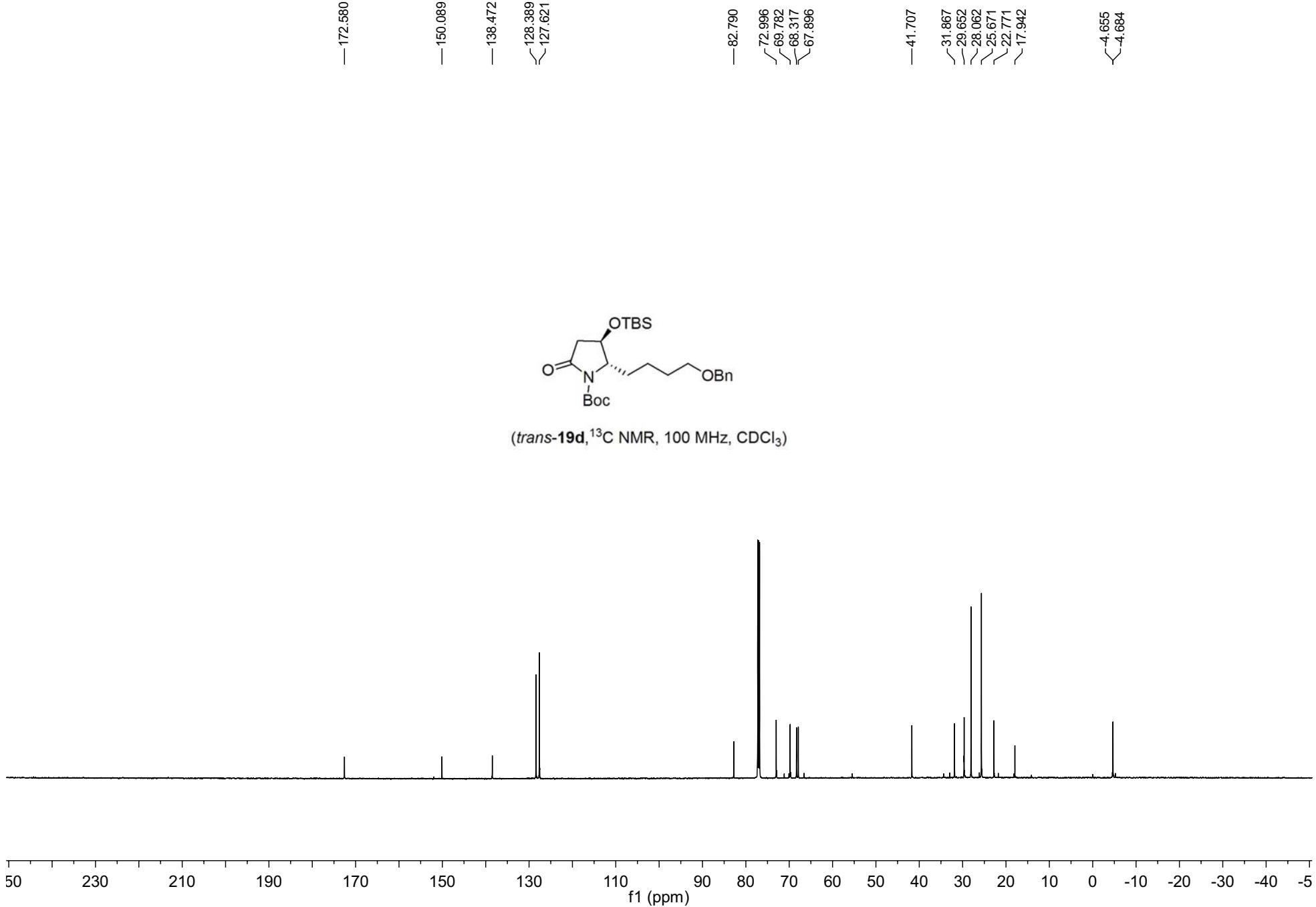
(*trans*-19a,  $^1\text{H}$  NMR, 400 MHz,  $\text{CDCl}_3$ )

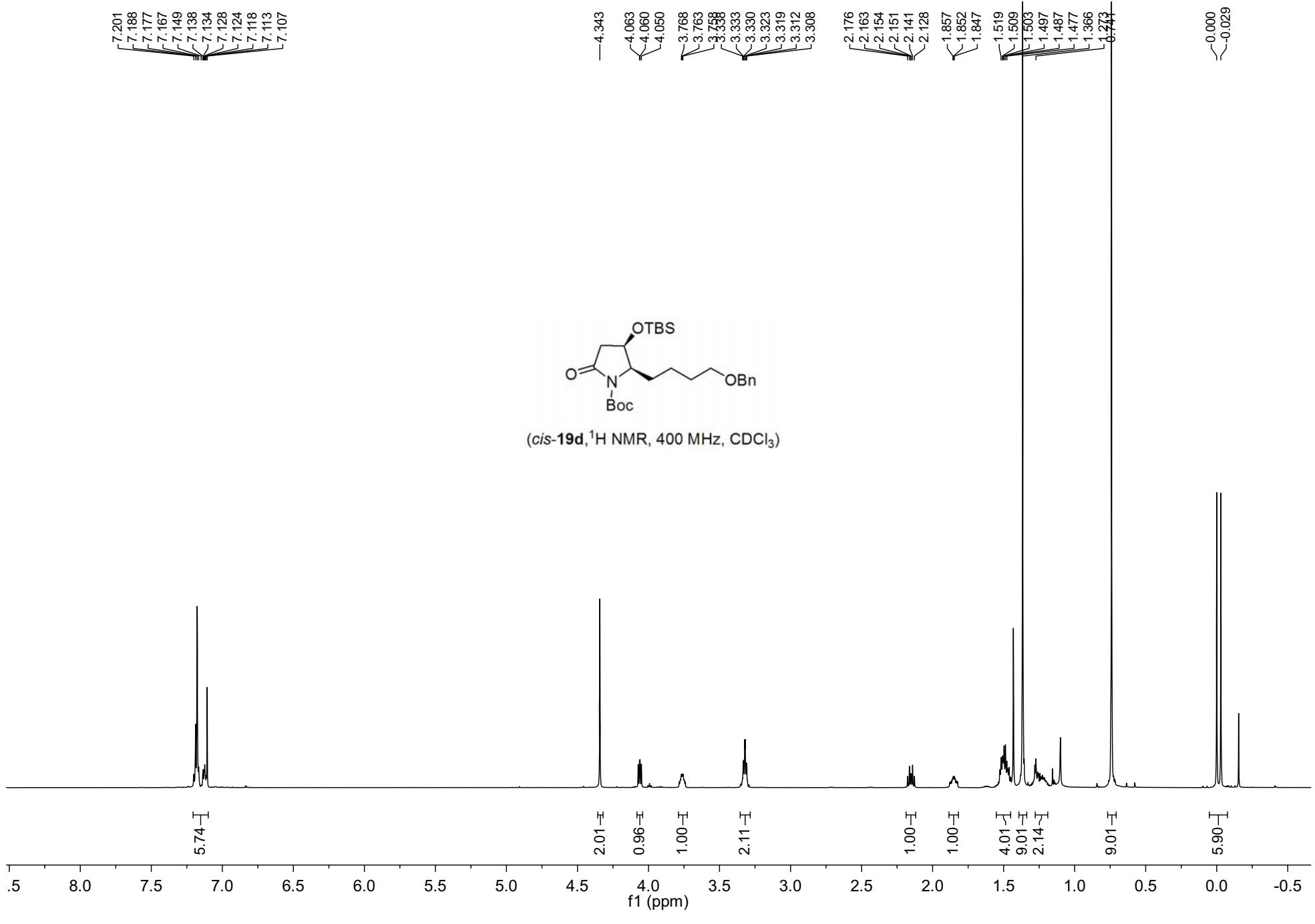


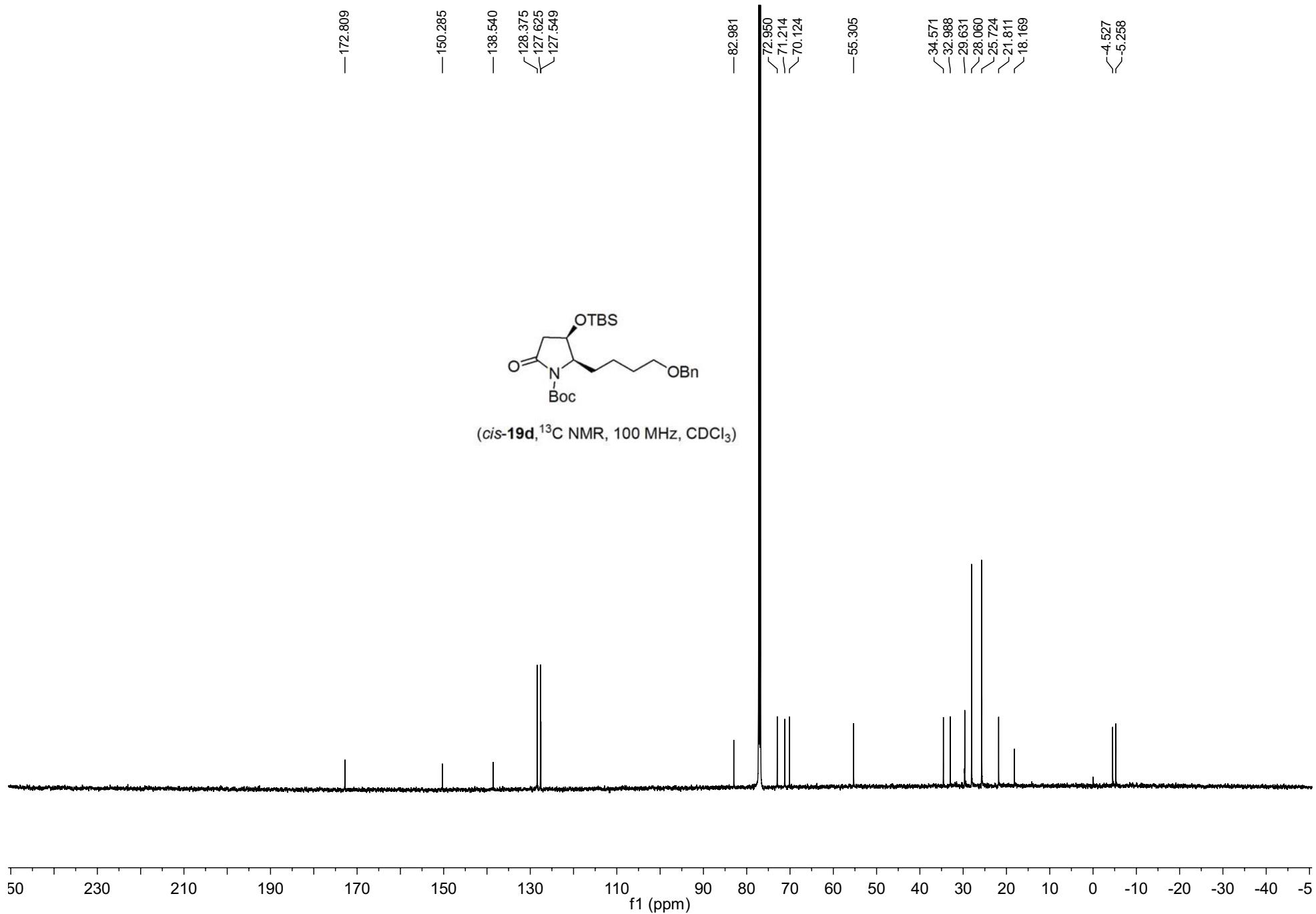


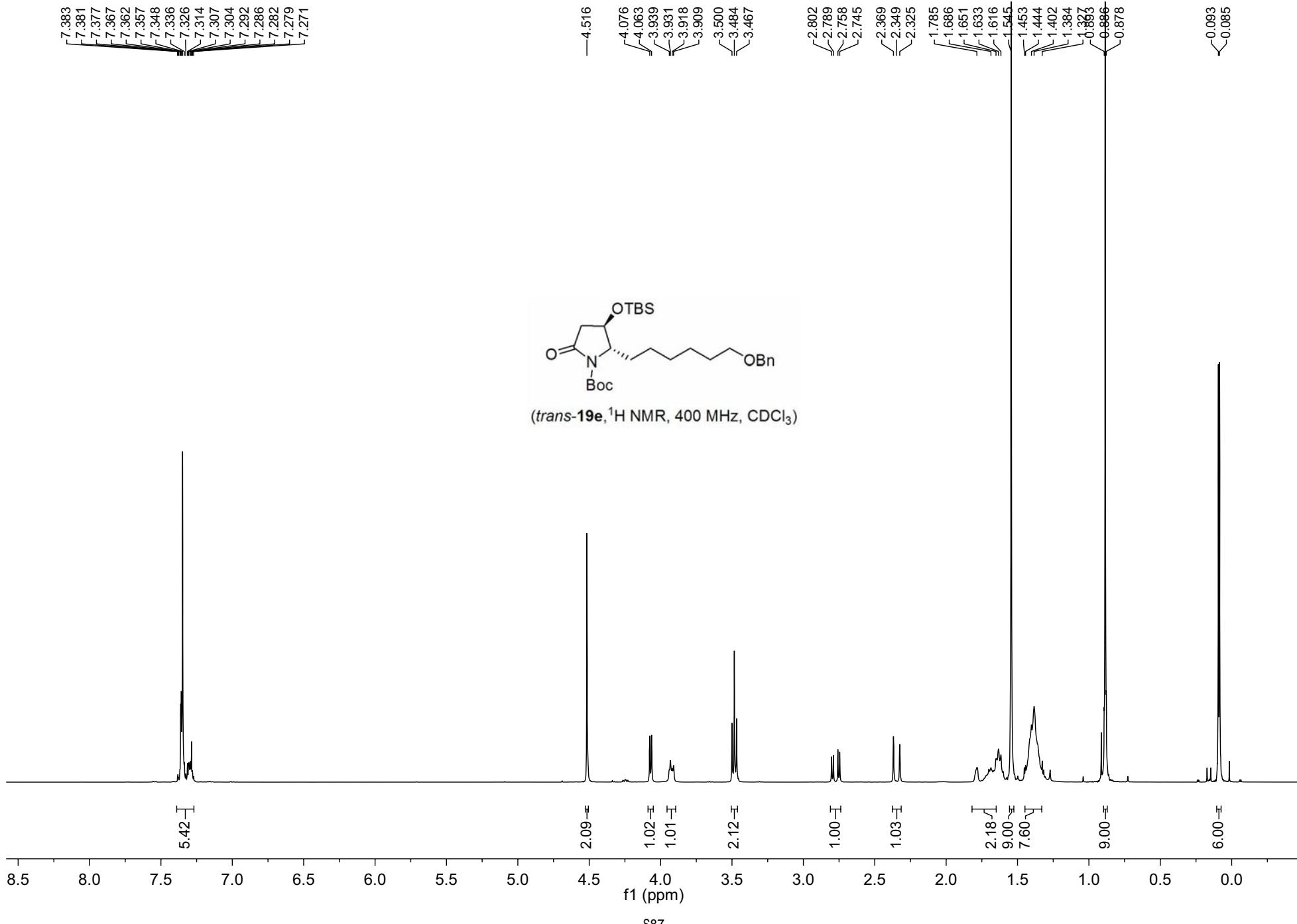


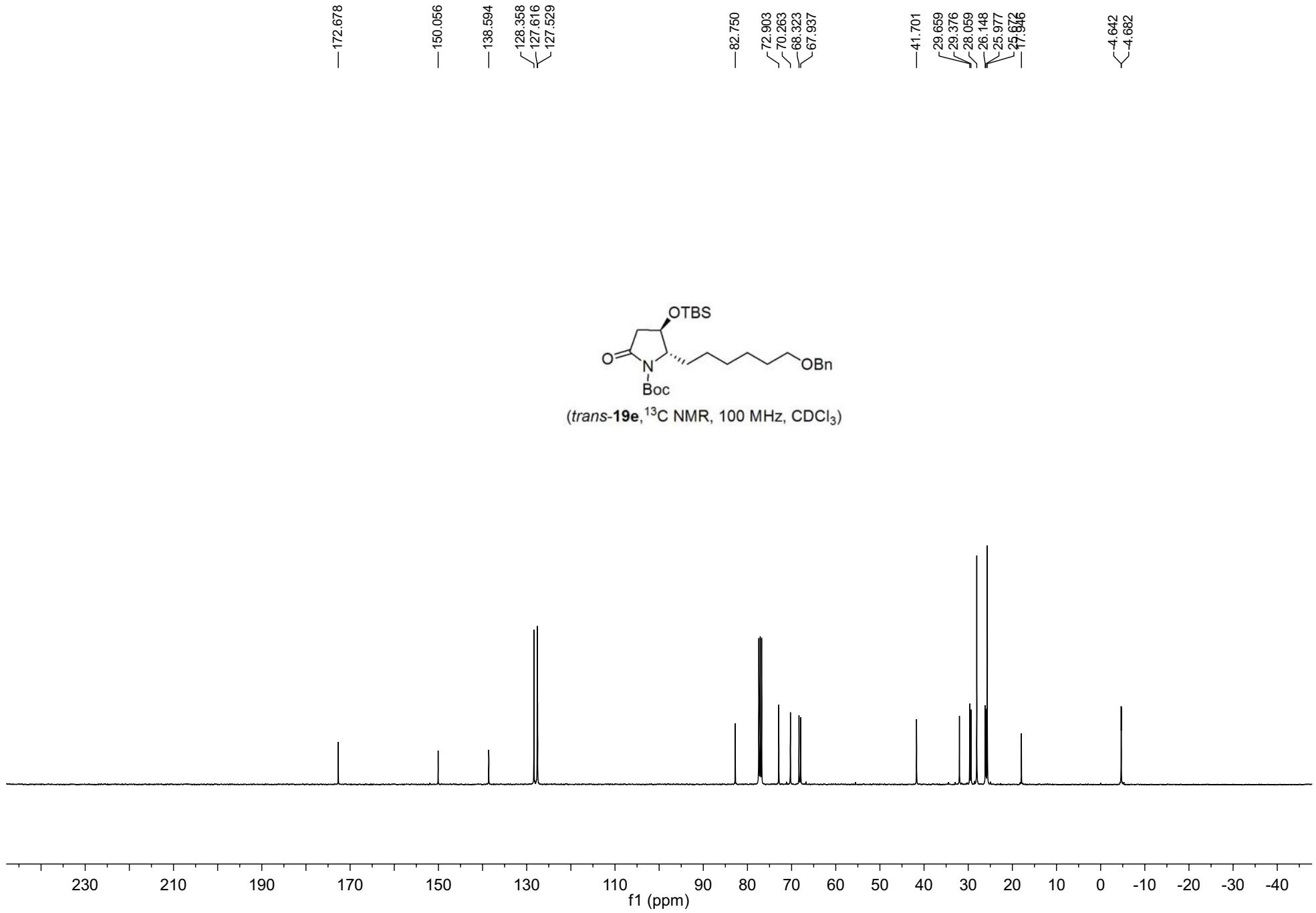


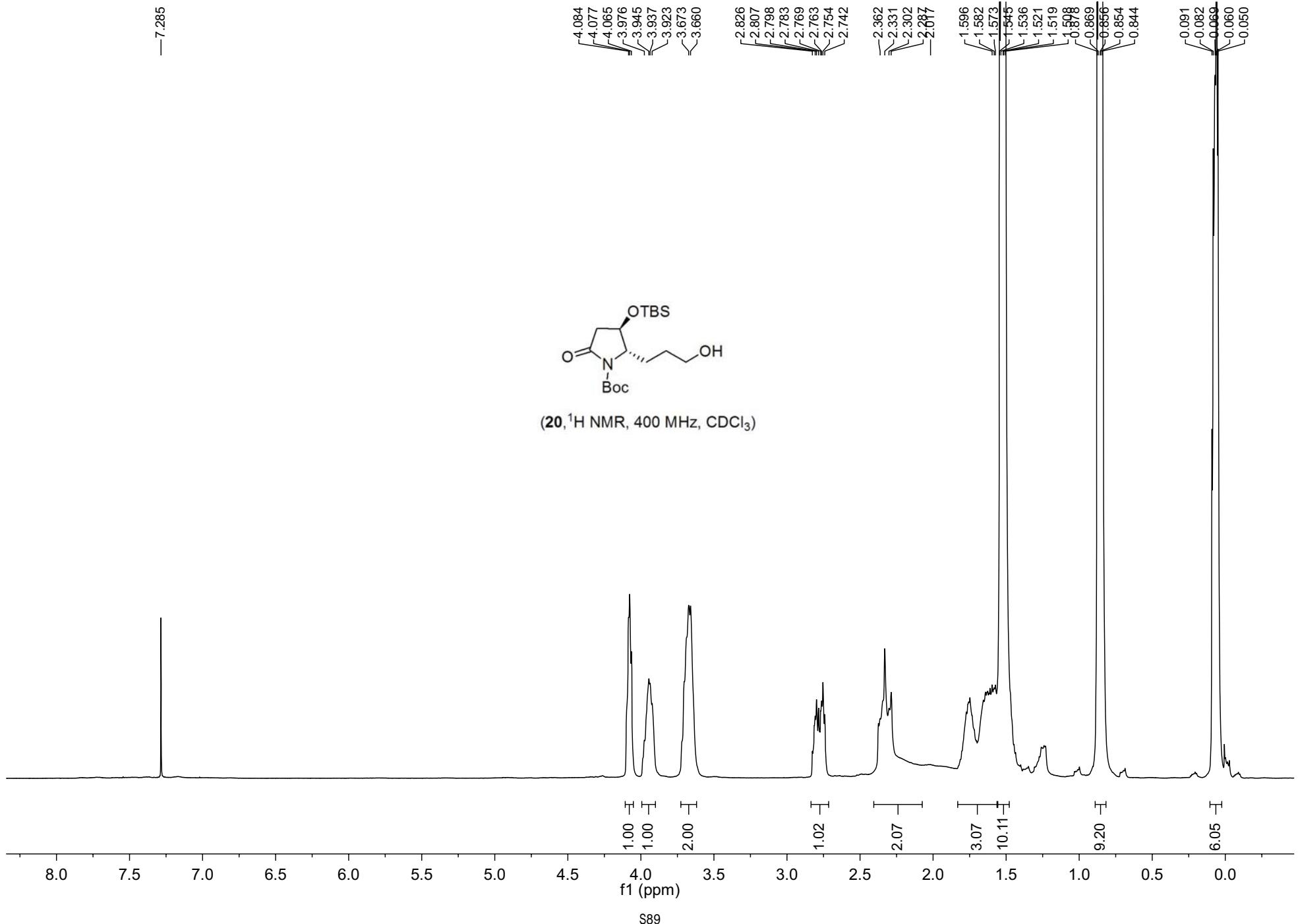


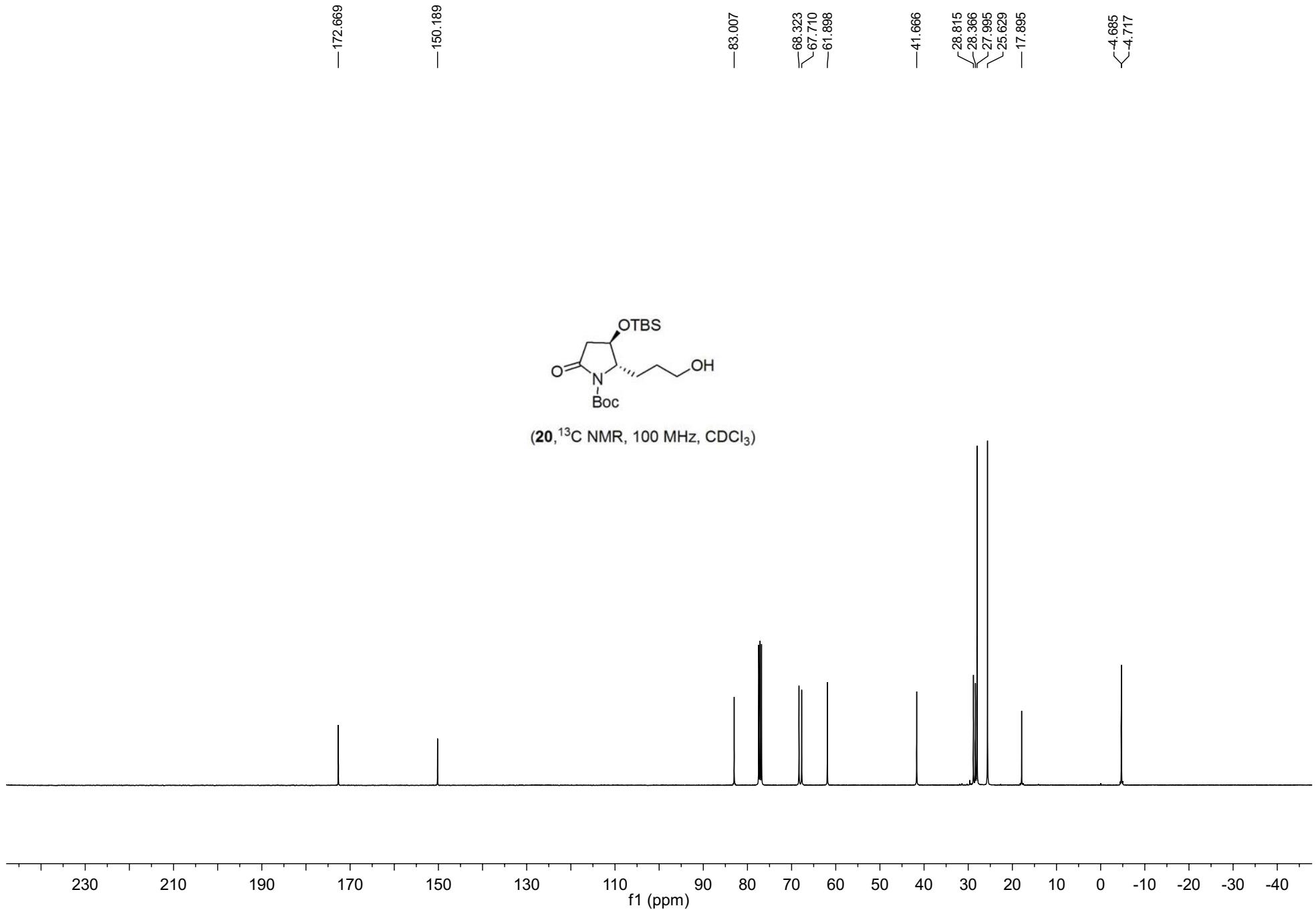


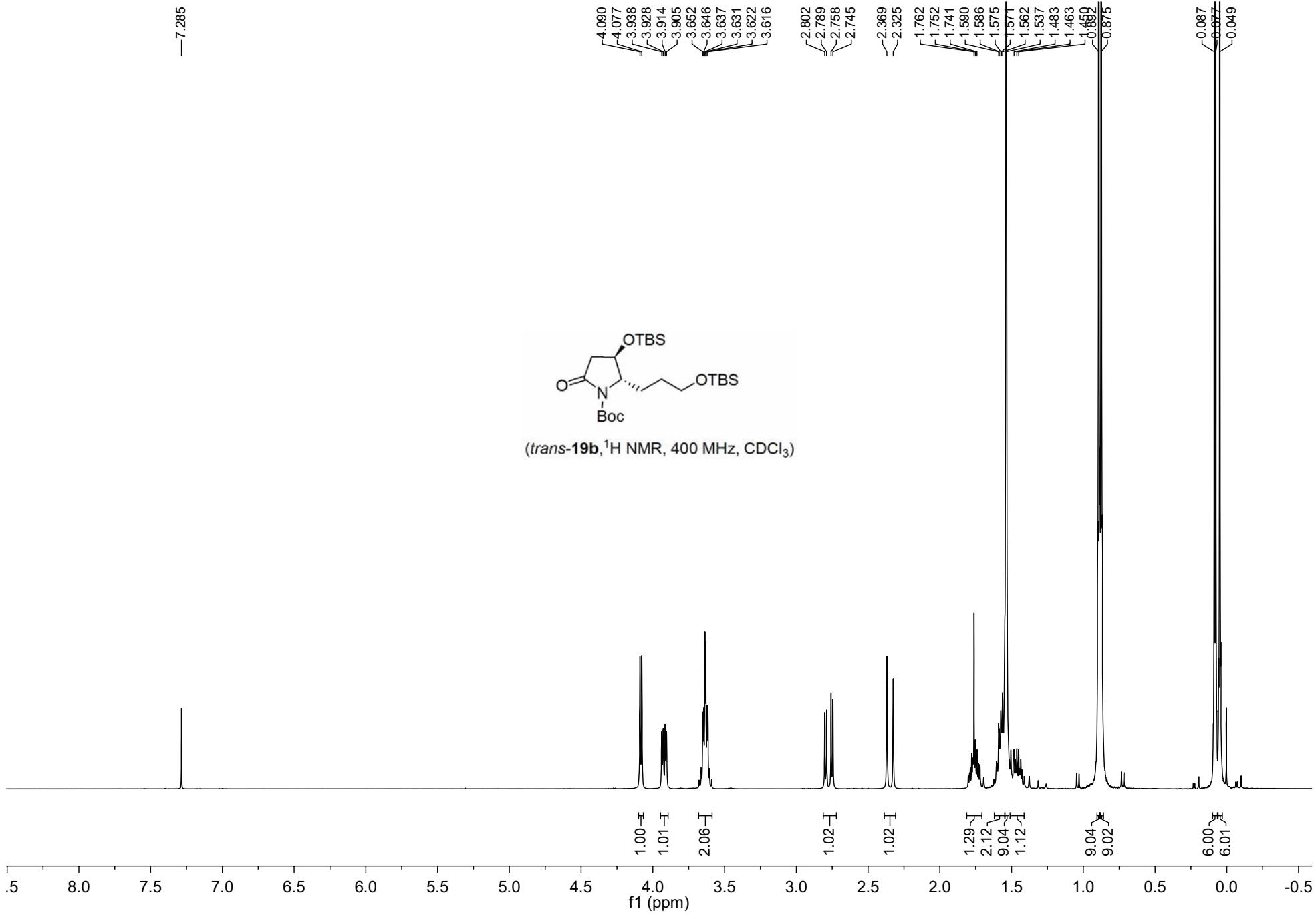


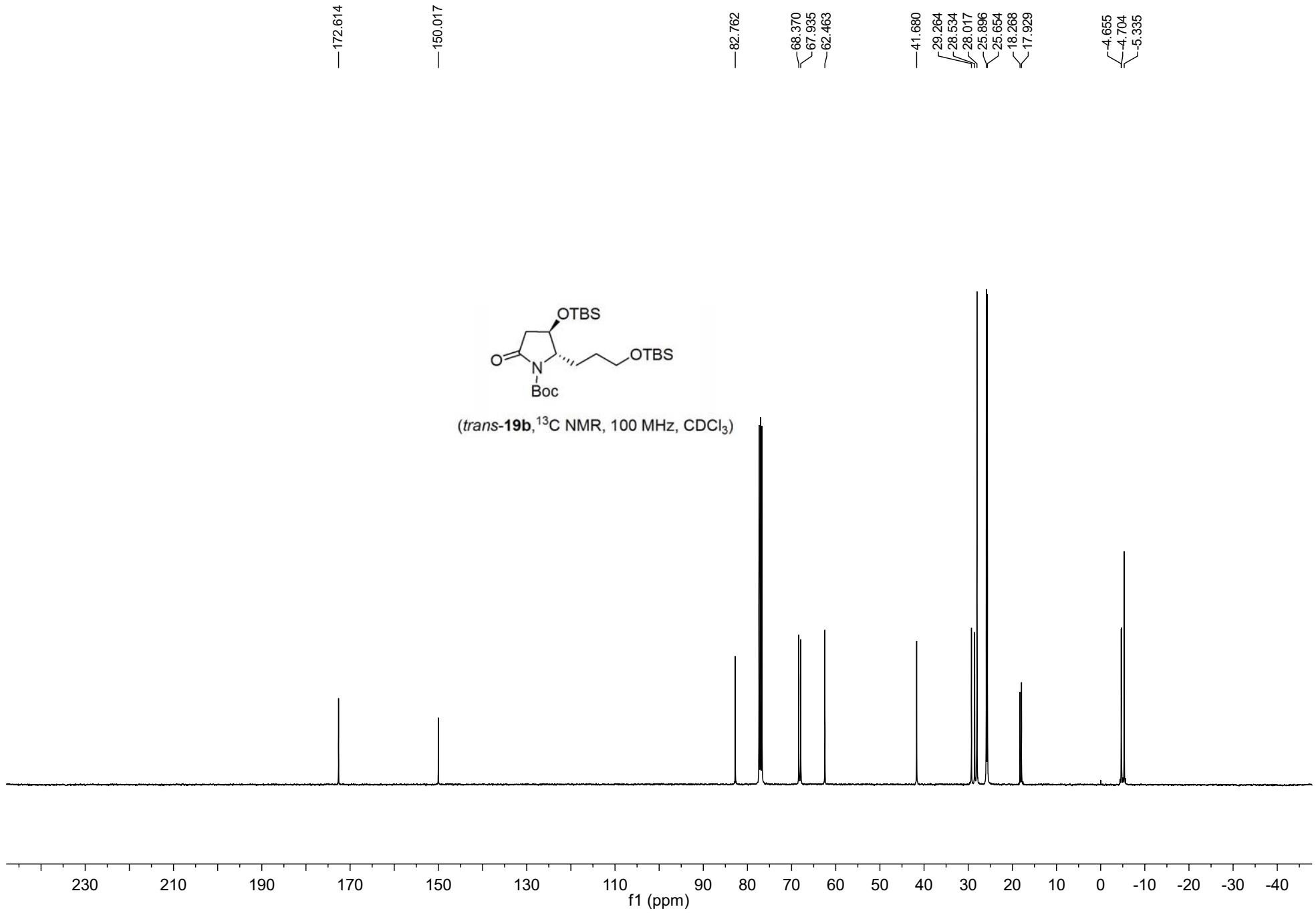


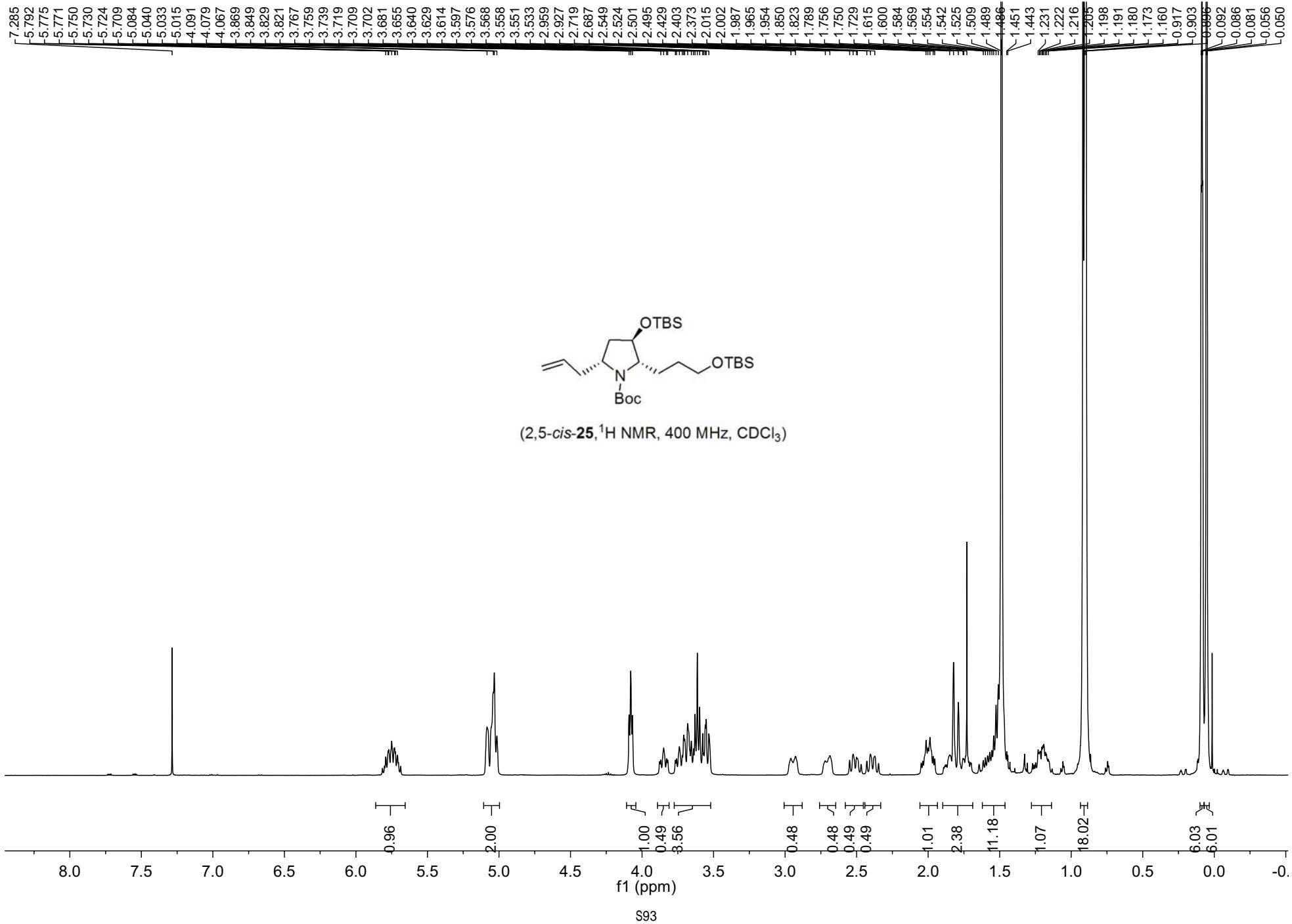


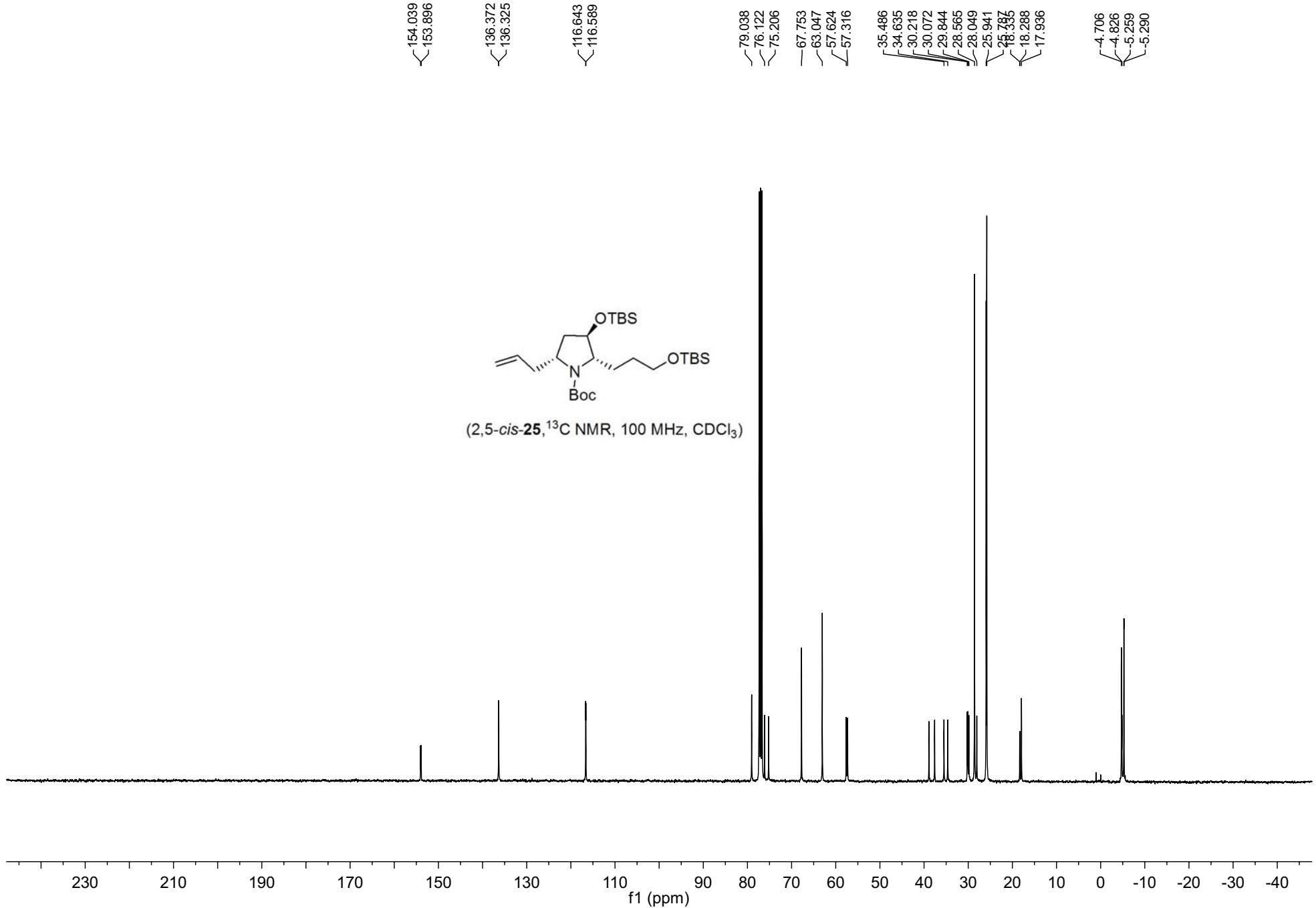


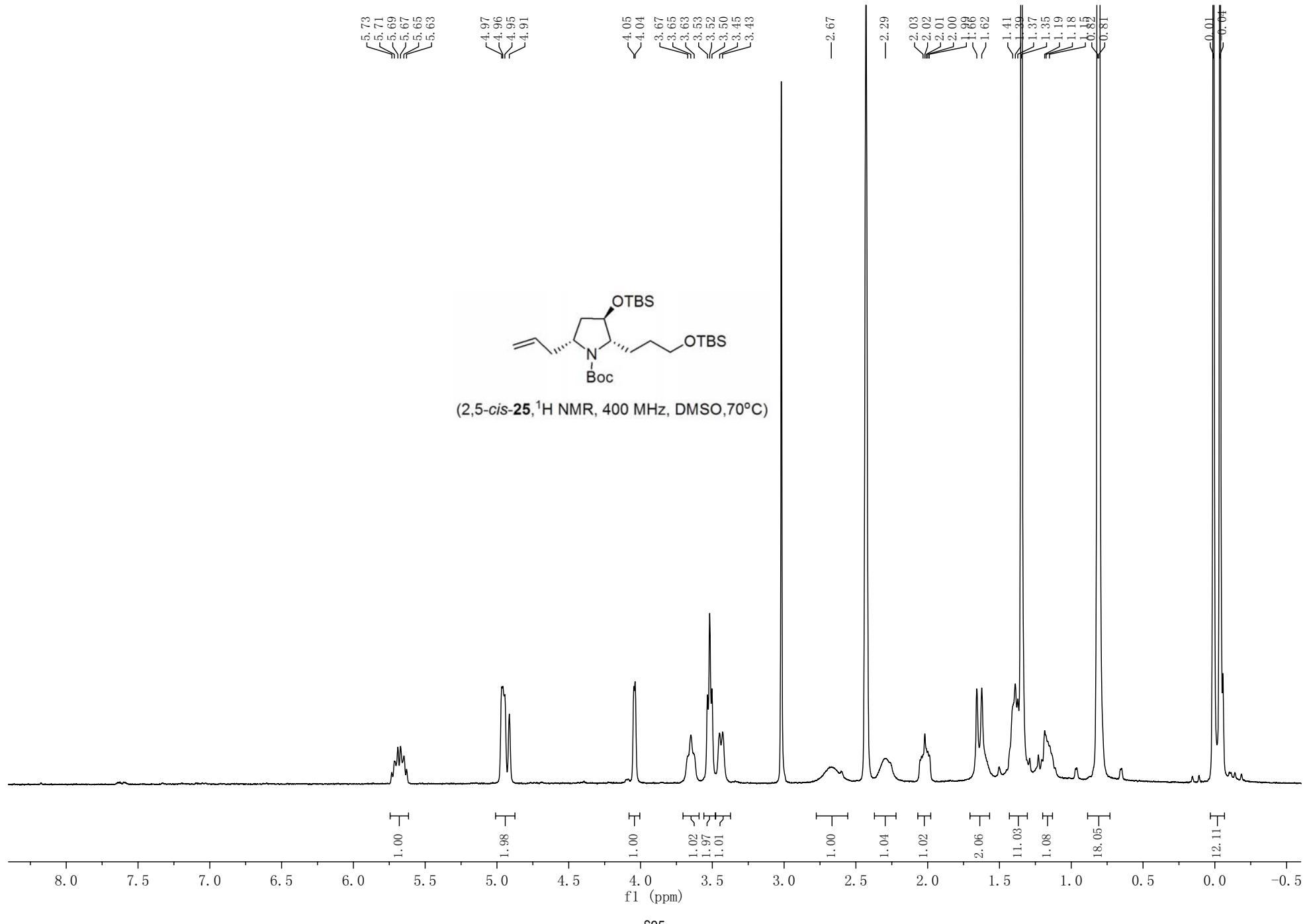


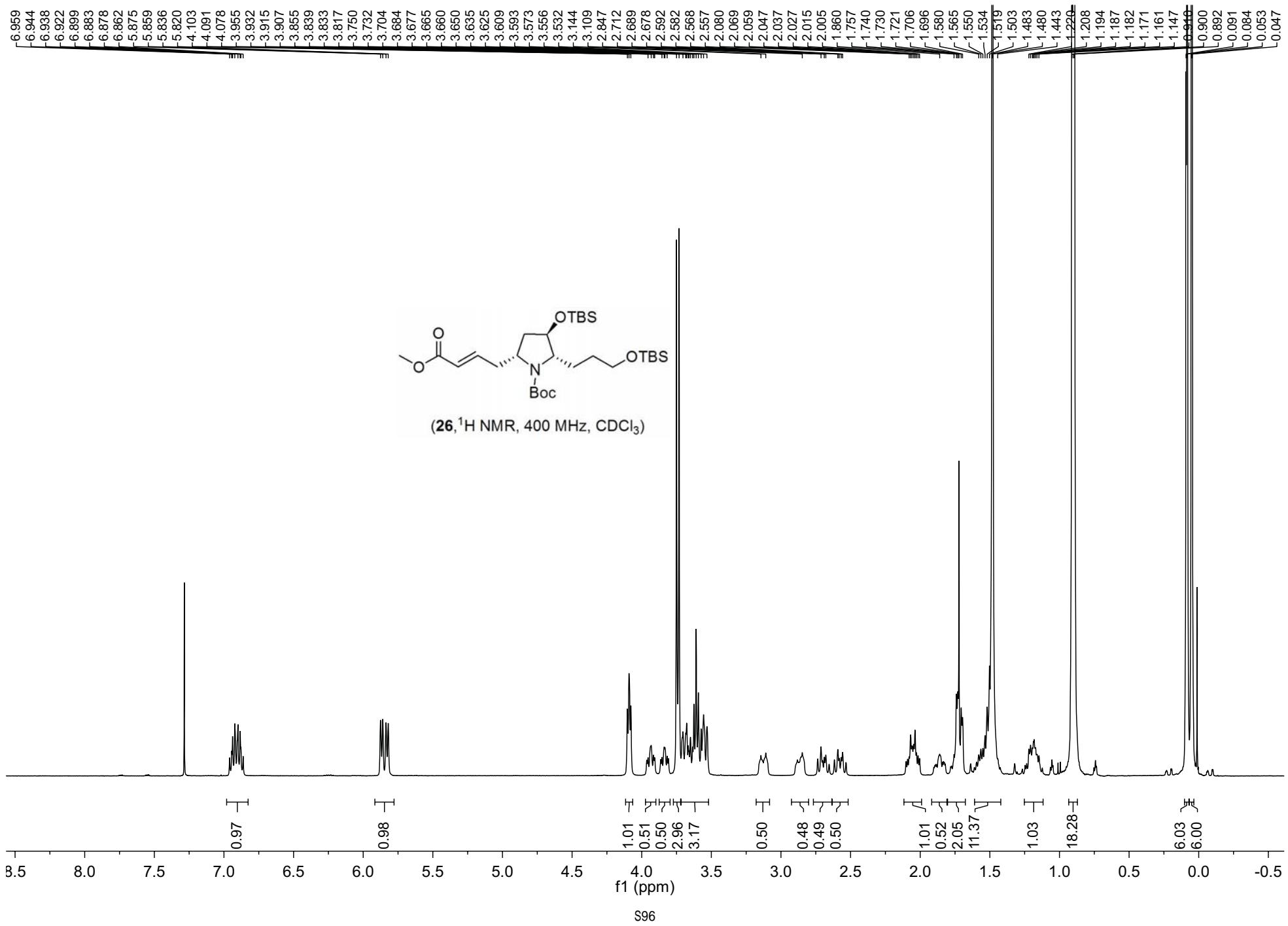


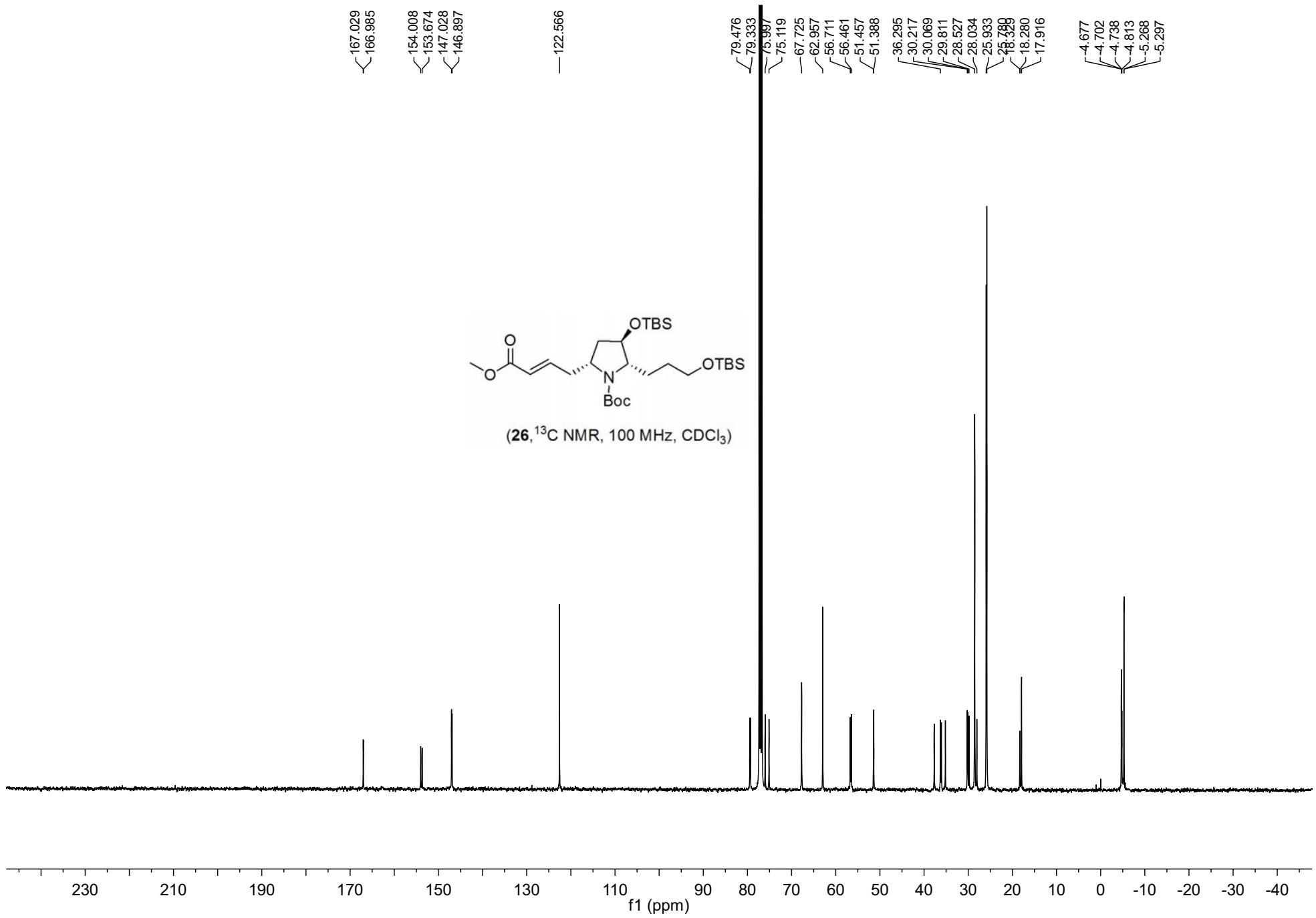


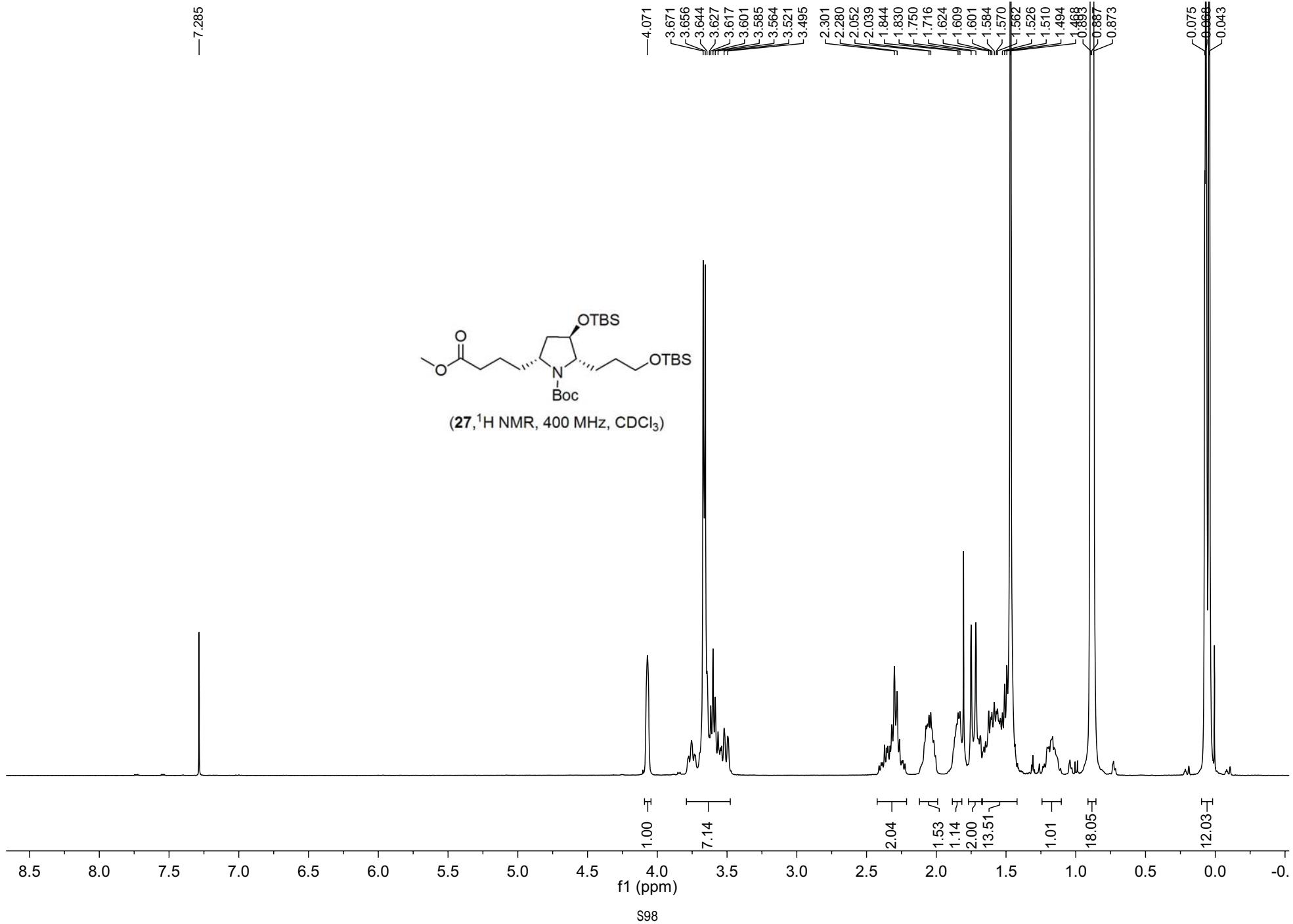


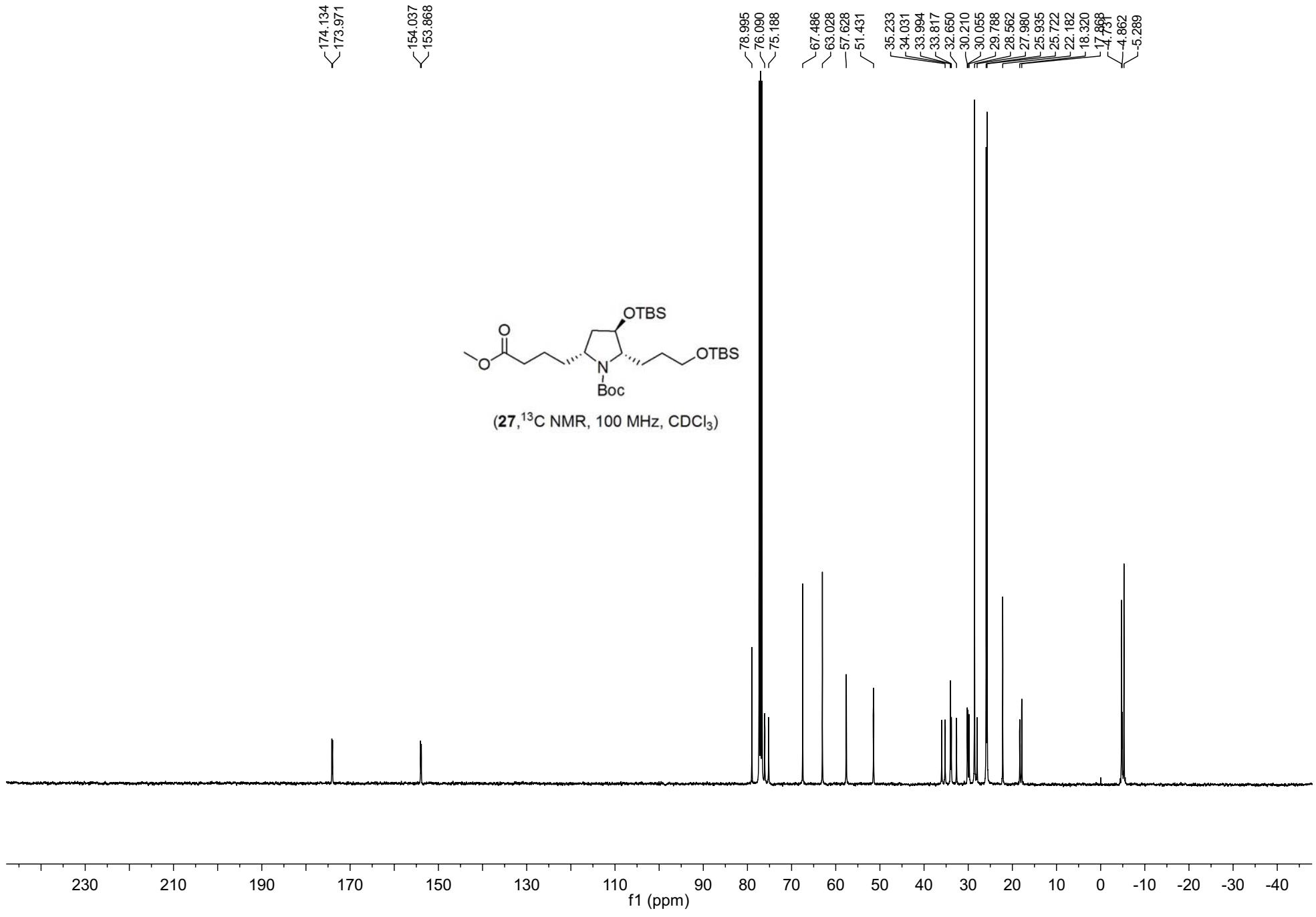


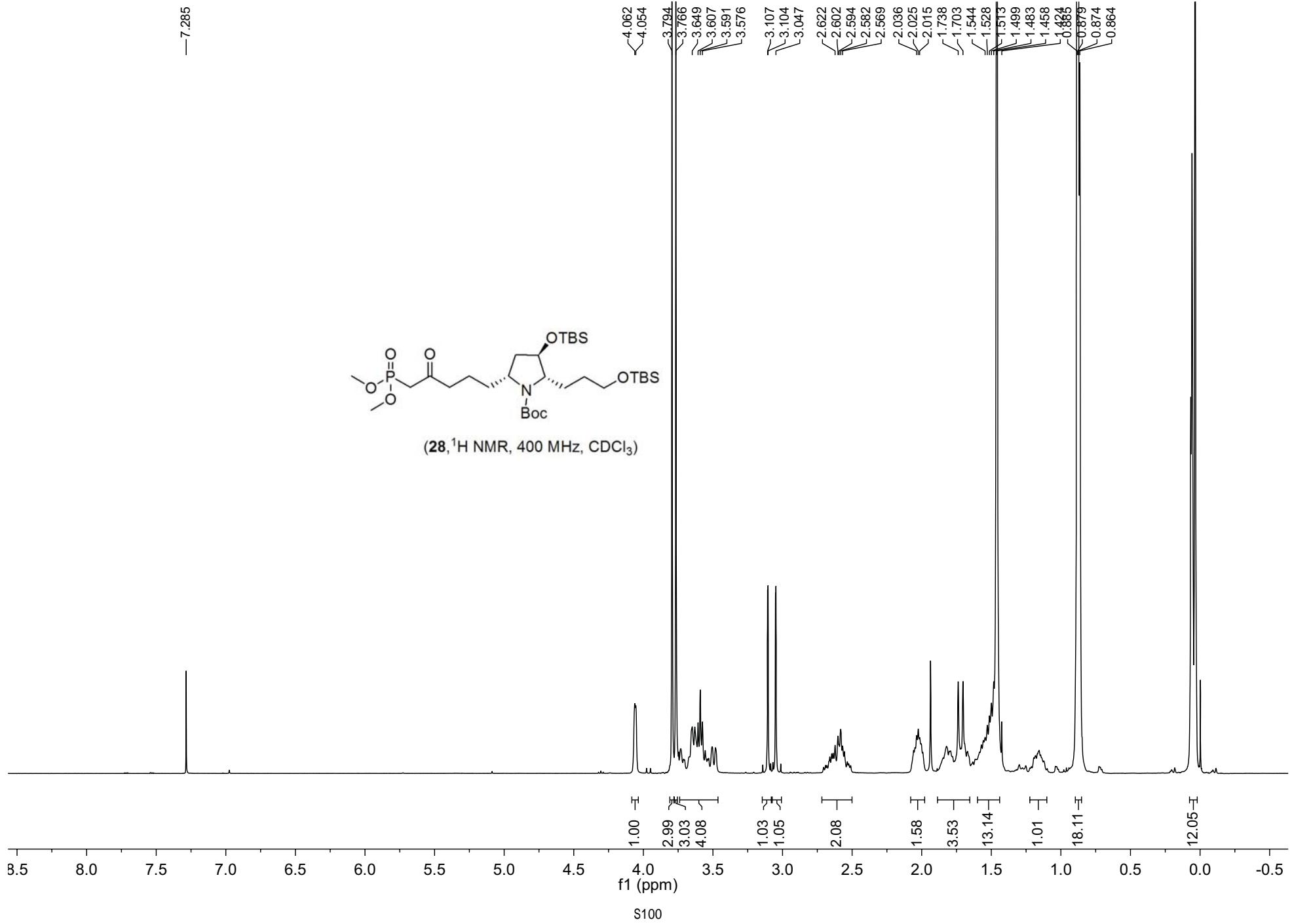


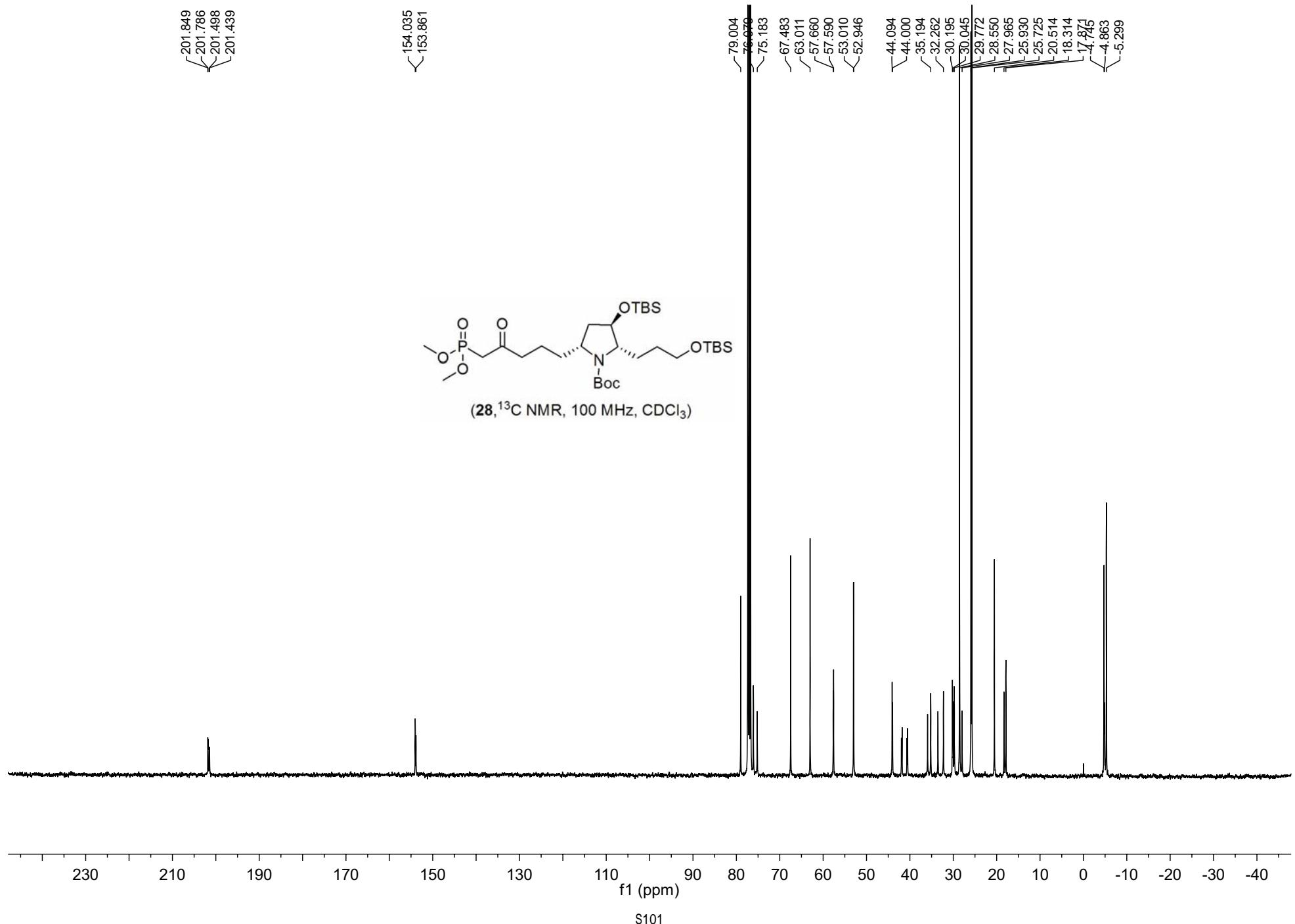




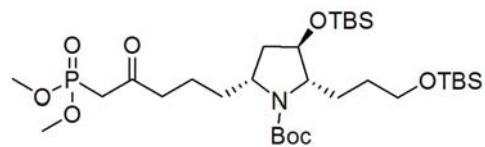




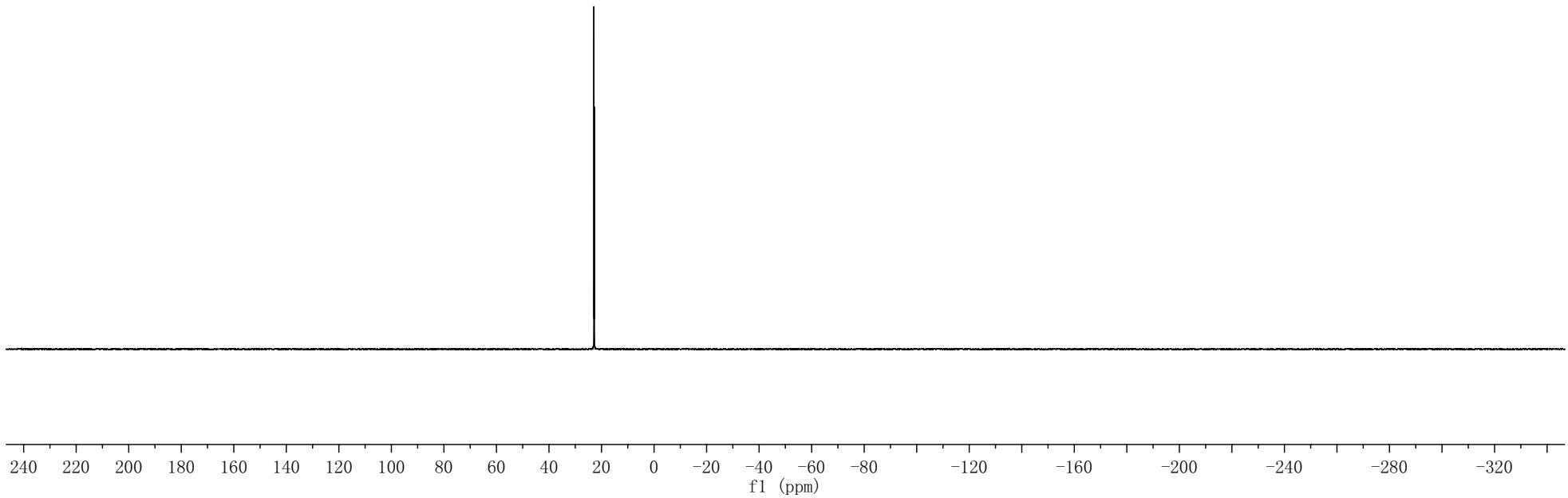


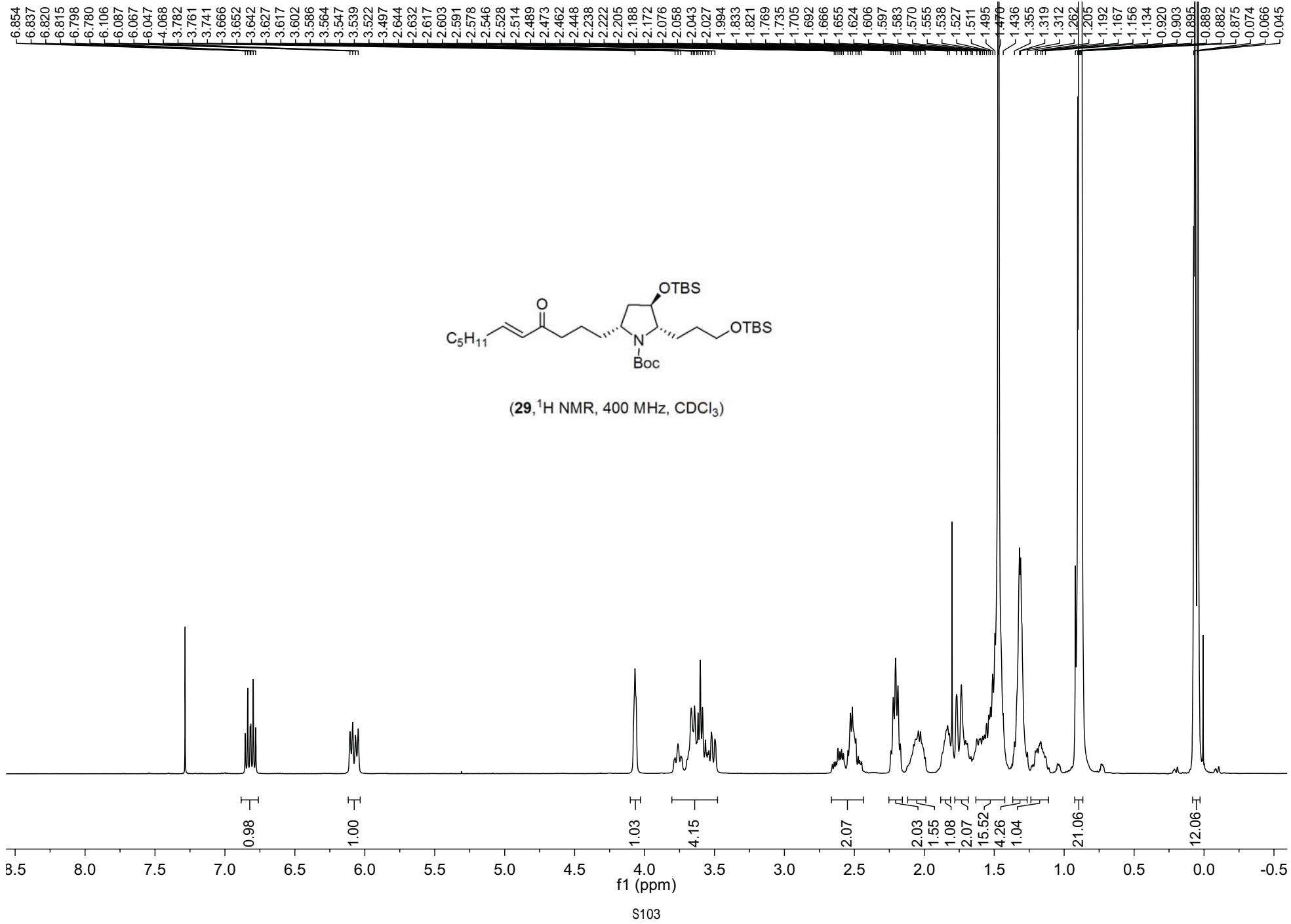


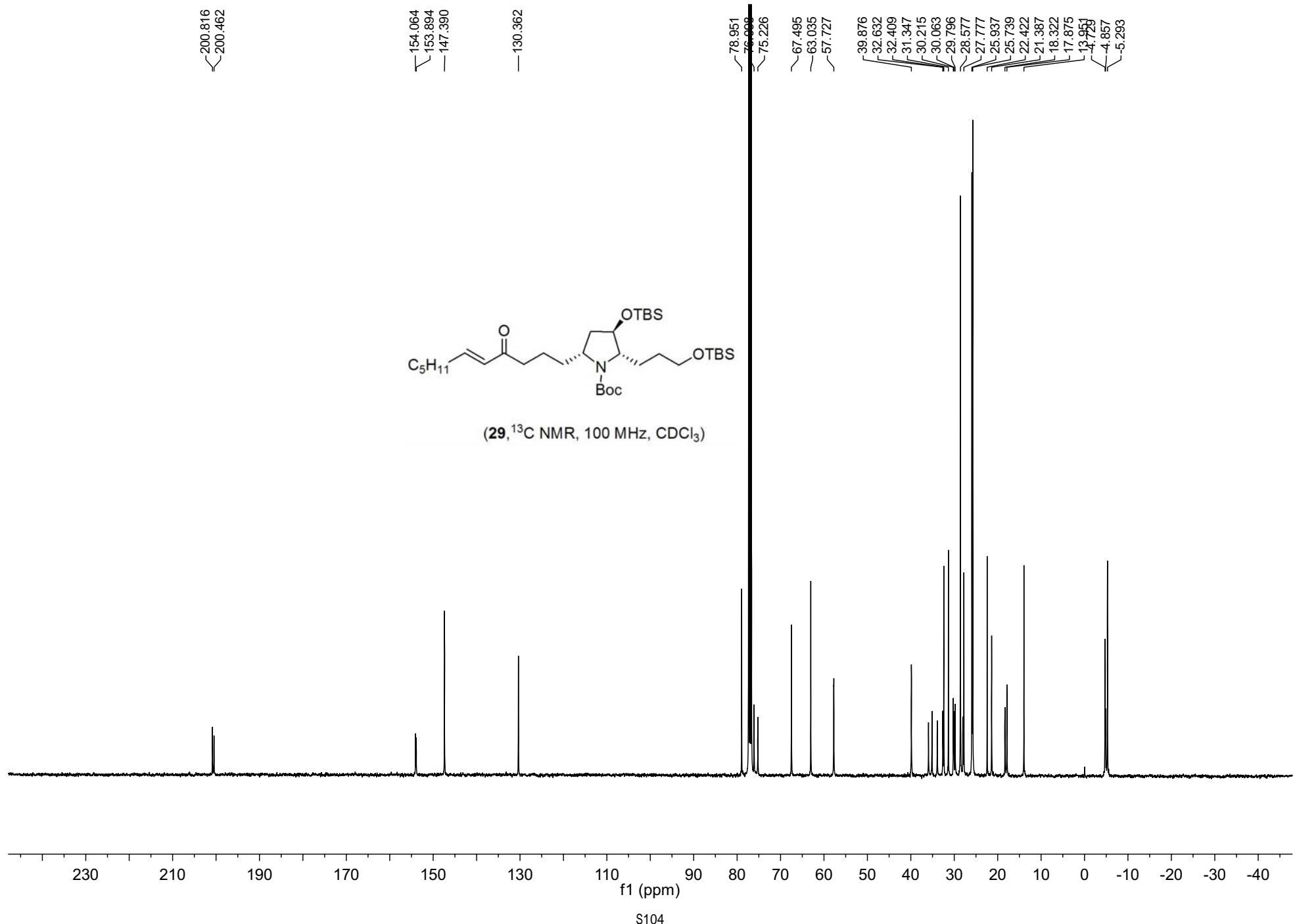
Y<sup>22, 89</sup>  
Y<sup>22, 73</sup>



(28,  $^{31}\text{P}$  NMR, 121.5 MHz,  $\text{CDCl}_3$ )







(**29**,  $^{13}\text{C}$  NMR, 100 MHz,  $\text{CDCl}_3$ )

