

Supporting Information

Synthesis of tRNA analogues containing a terminal ribose locked in the South conformation to study tRNA-dependent enzyme

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ETHEVE-QUELQUEJEU^{*a}

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1- NMR Spectra

^1H and ^{13}C spectra of compound **2**

^1H and ^{13}C spectra of compound **3**

^1H and ^{13}C spectra of compound **4**

^1H , ^{13}C and ^{31}P spectra of compound **5**

^1H , ^{13}C and ^{31}P spectra of compound **6**

^1H and ^{31}P spectra of compound **7**

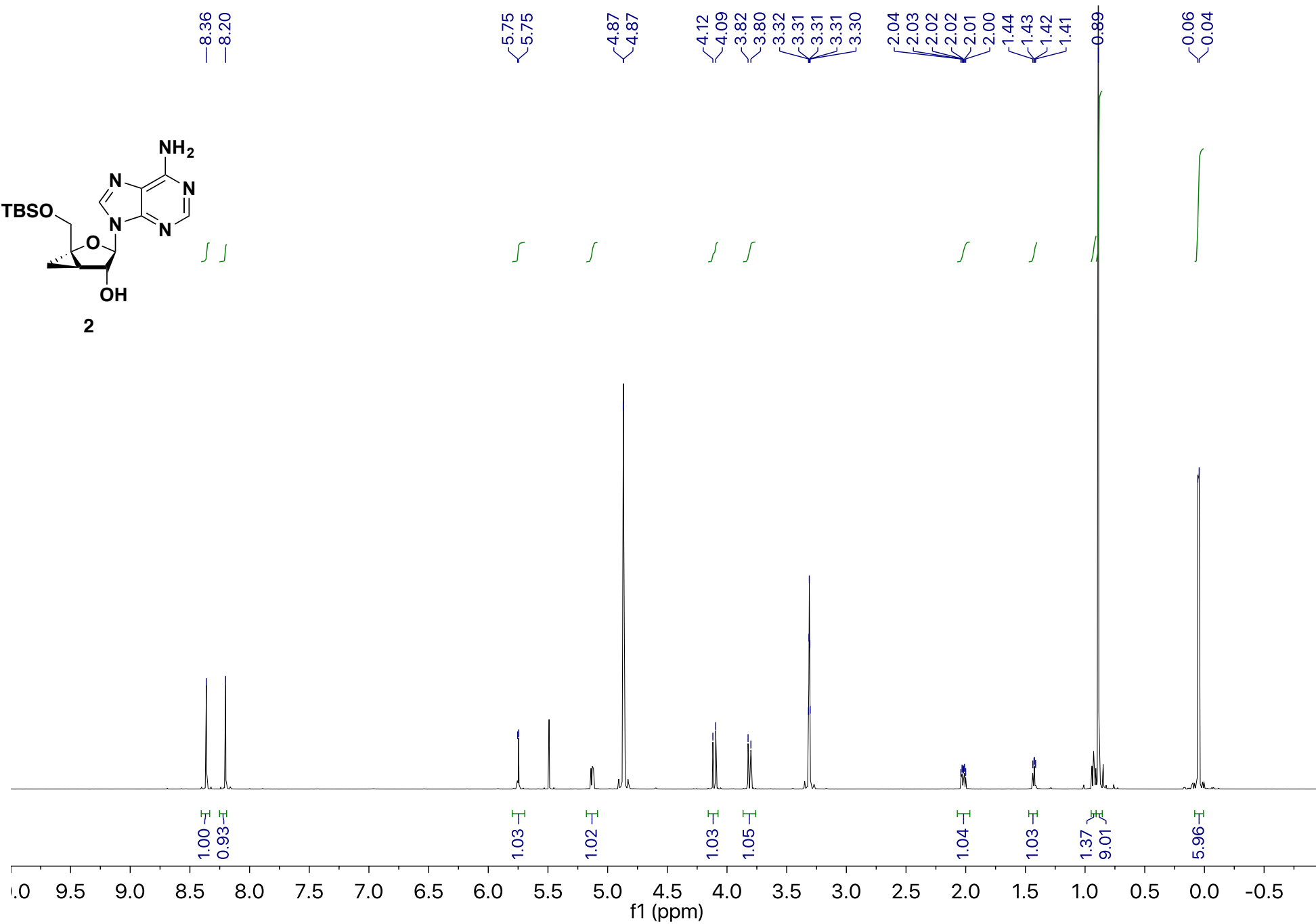
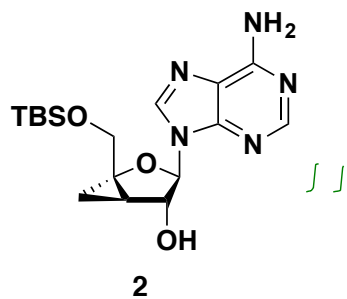
^1H and ^{13}C spectra of compound **8**

^1H and ^{13}C spectra of compound **9**

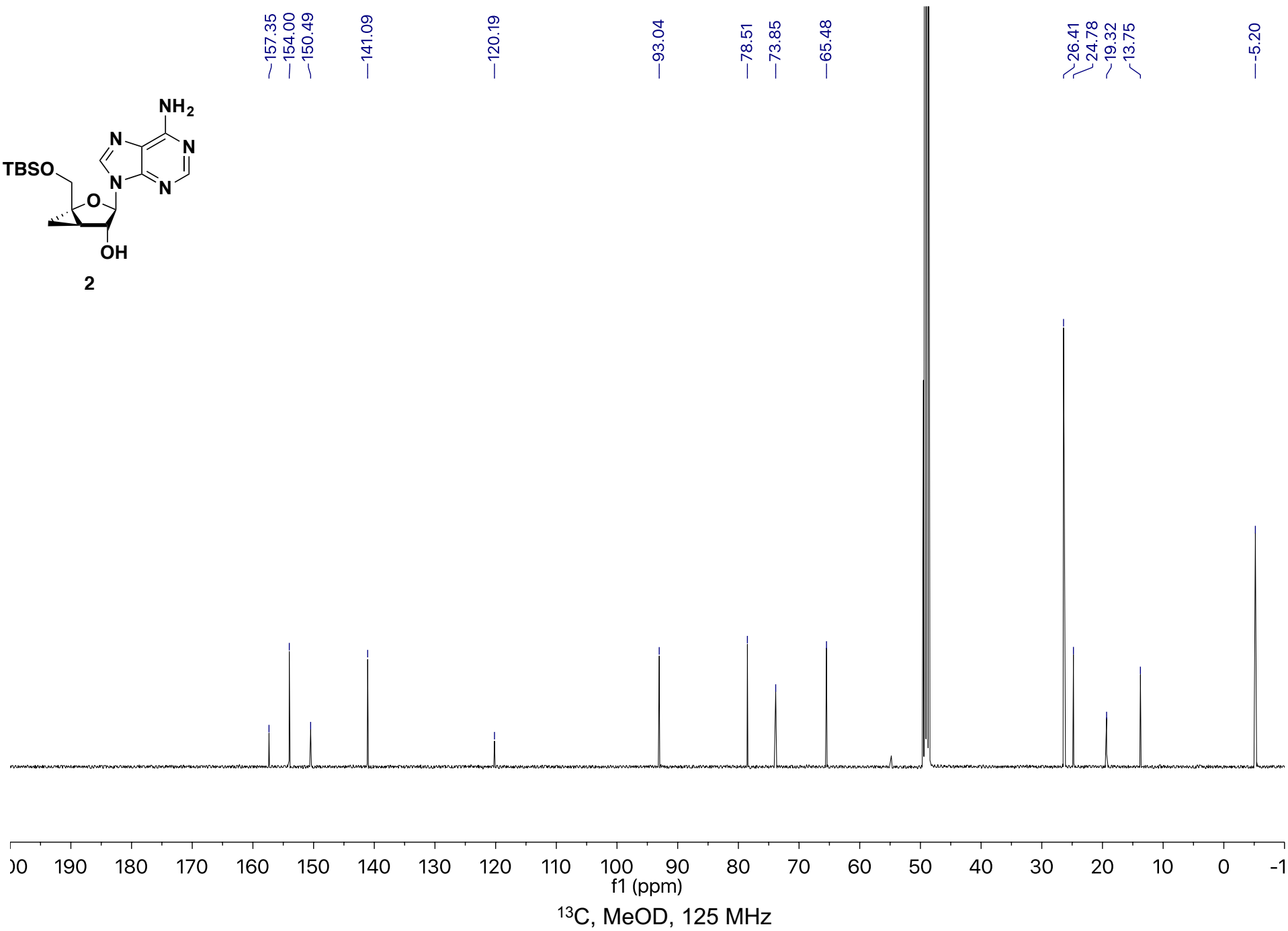
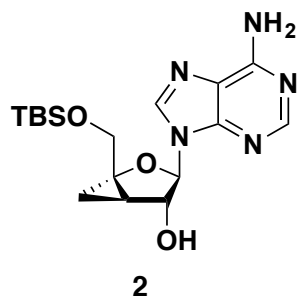
^1H spectrum of compound **10**

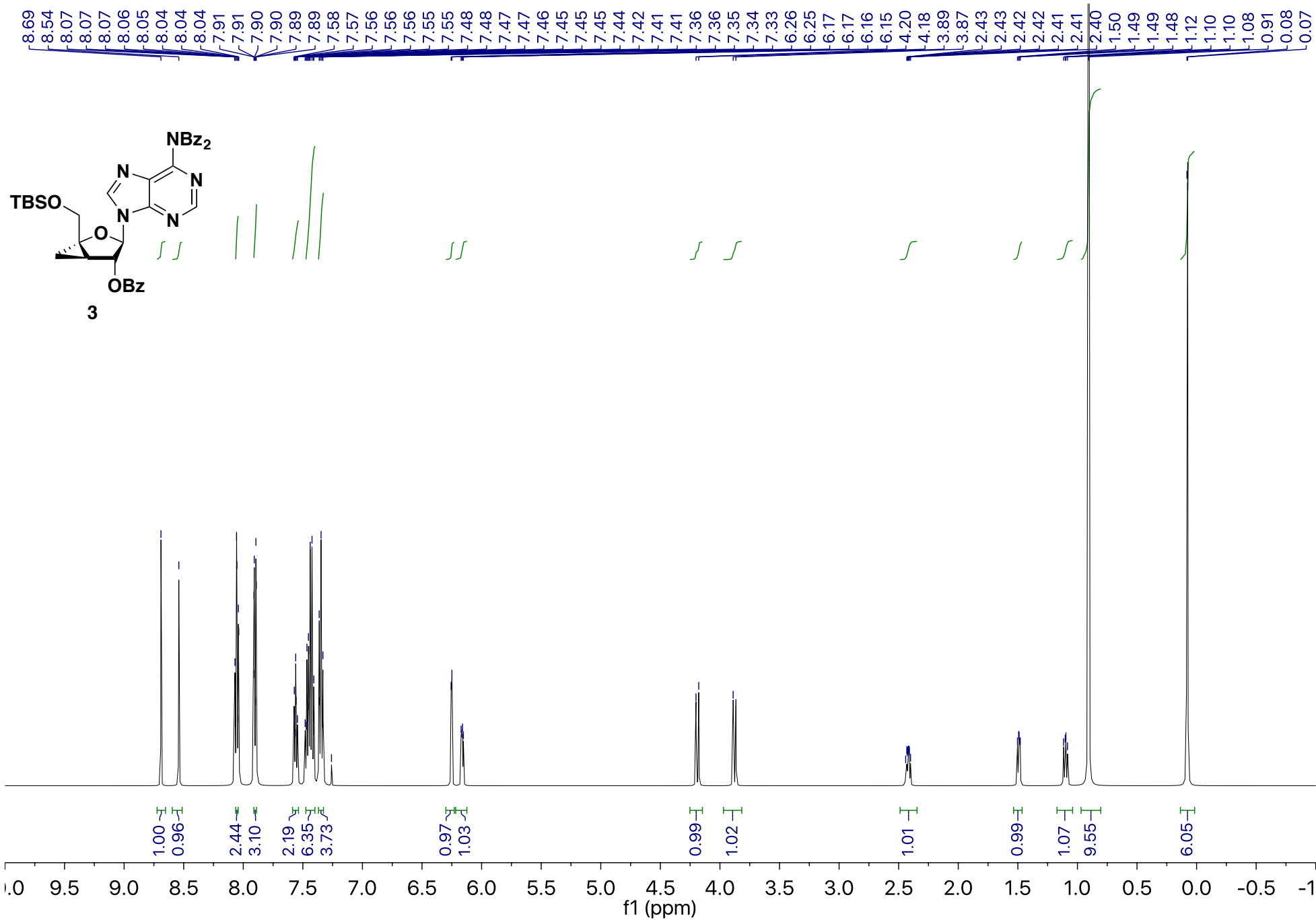
^1H and ^{31}P spectra of compound **11**

^1H spectrum of compound **12**

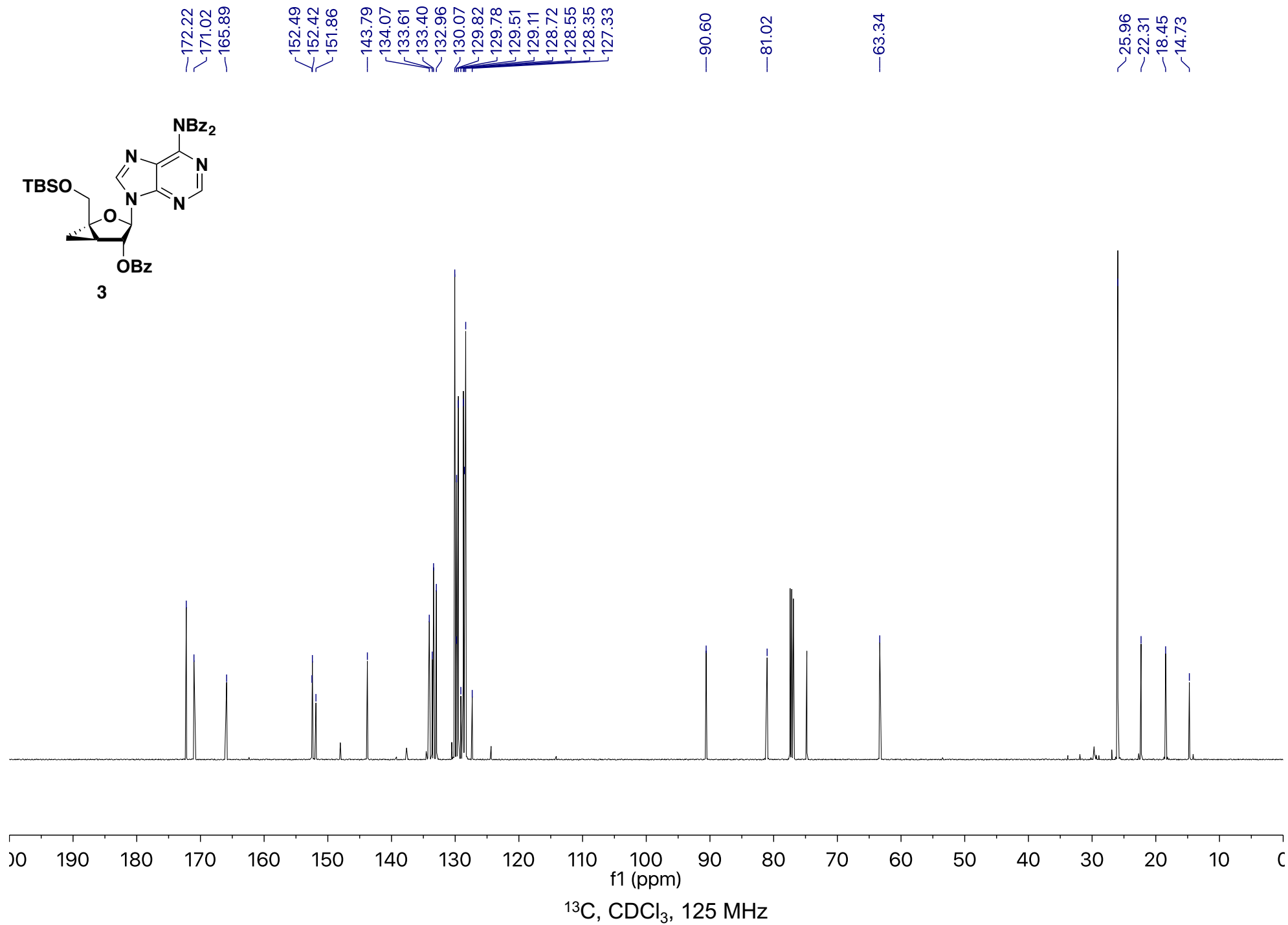
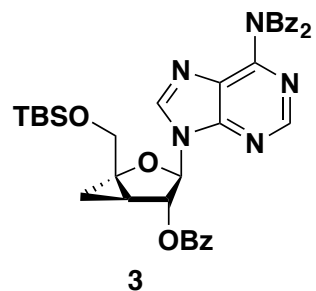


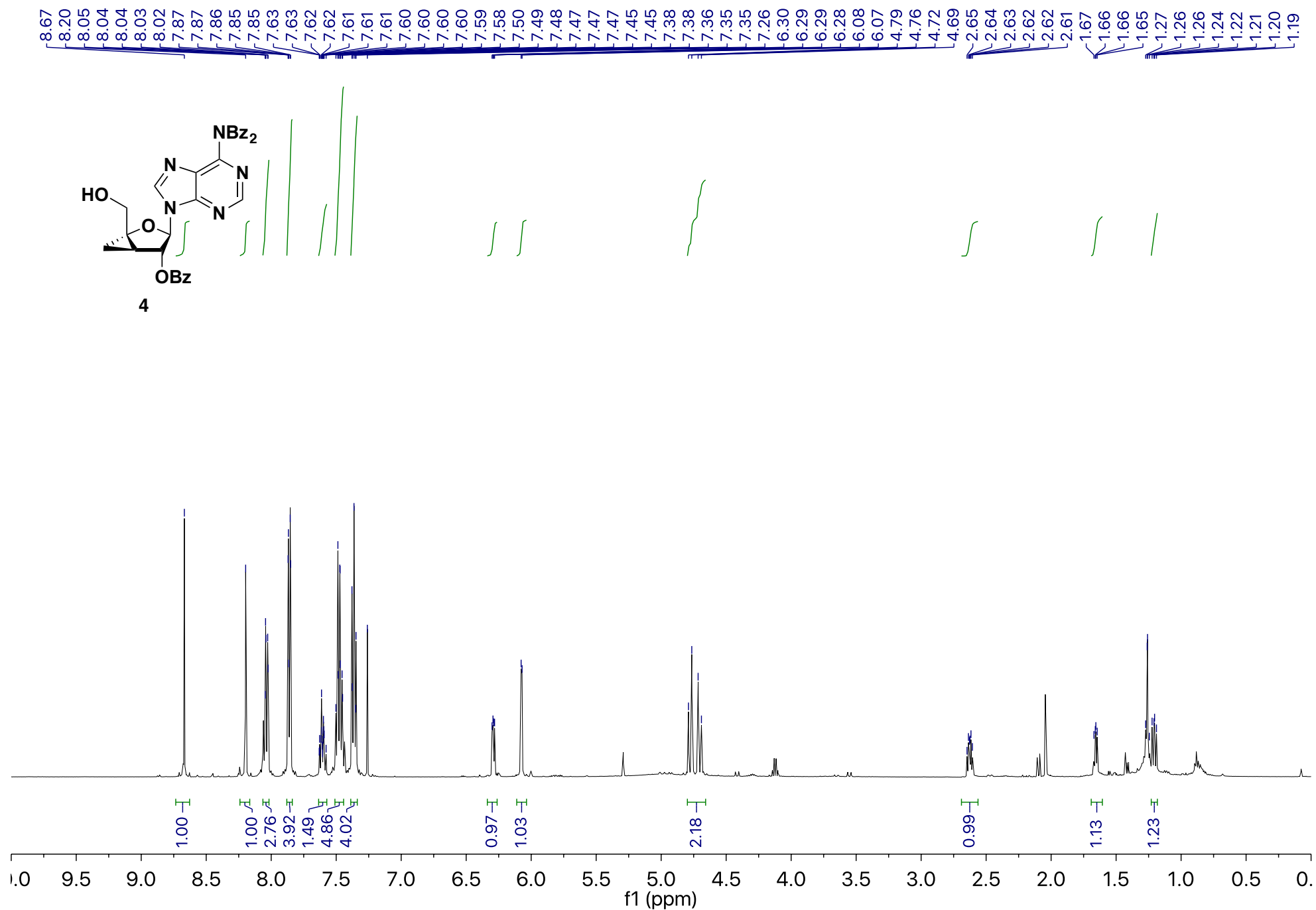
¹H, MeOD, 500 MHz



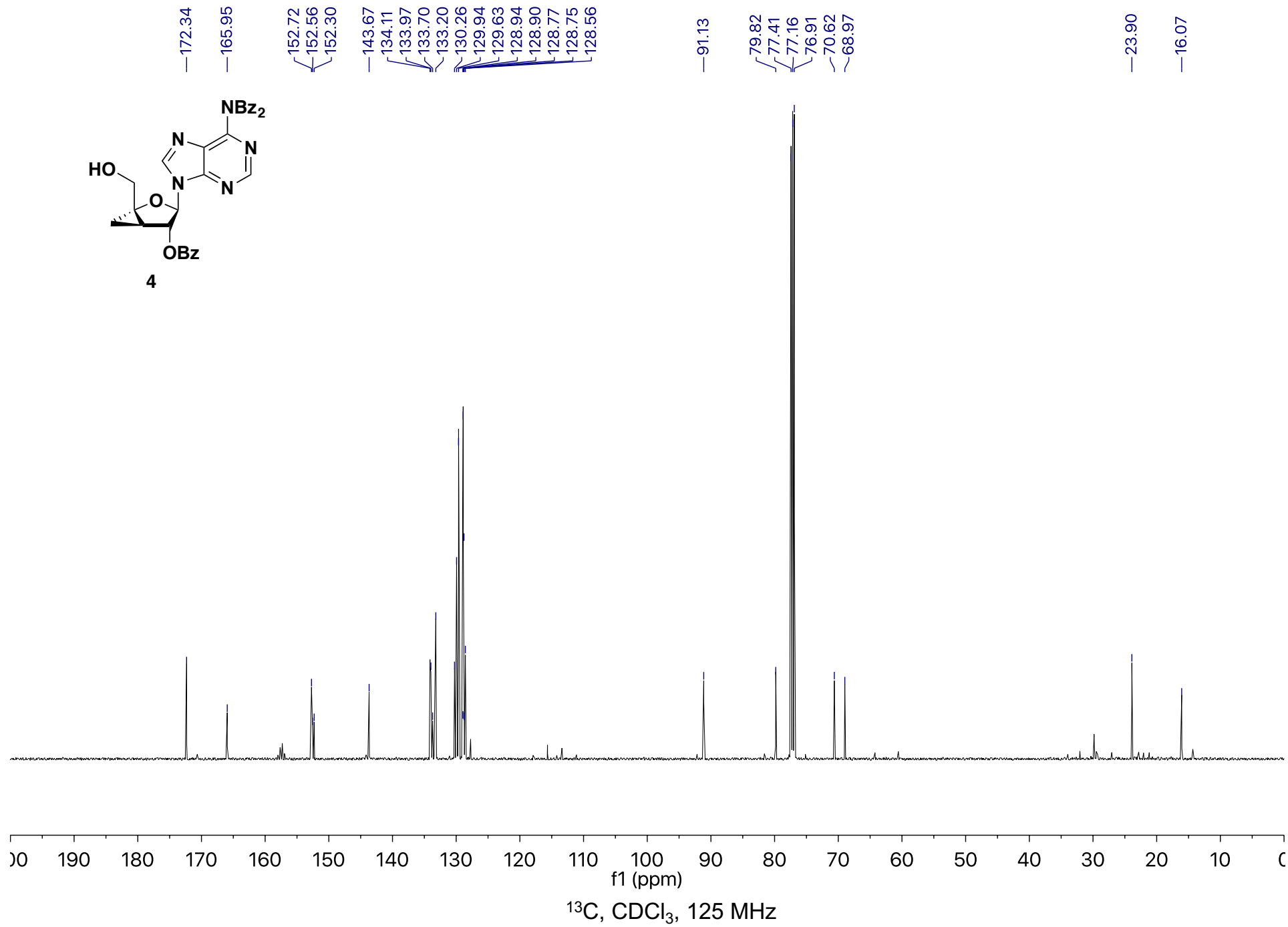
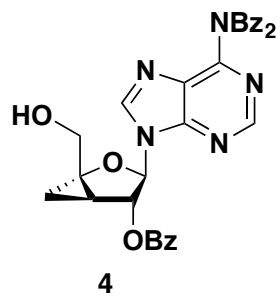


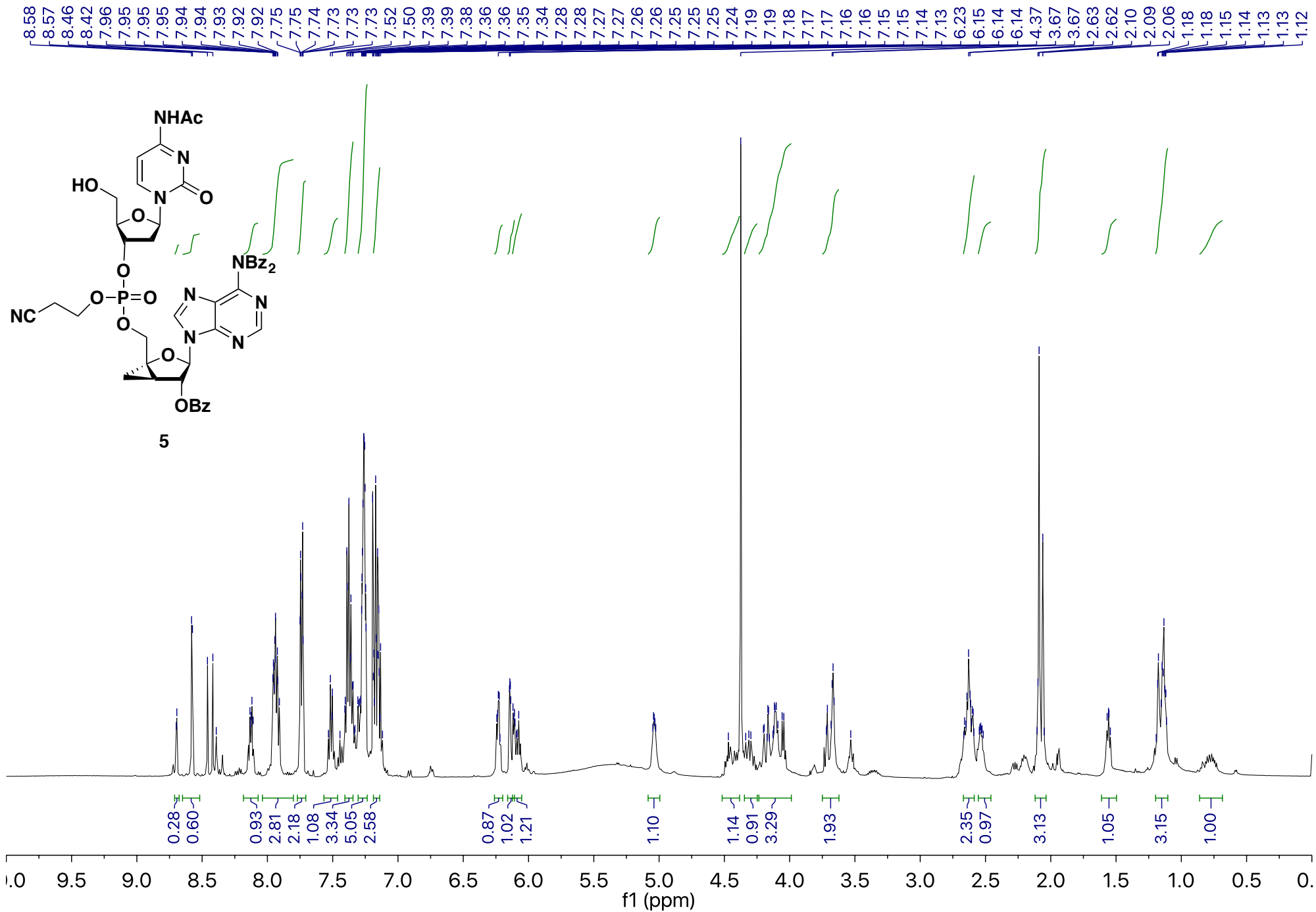
¹H, CDCl₃, 500 MHz



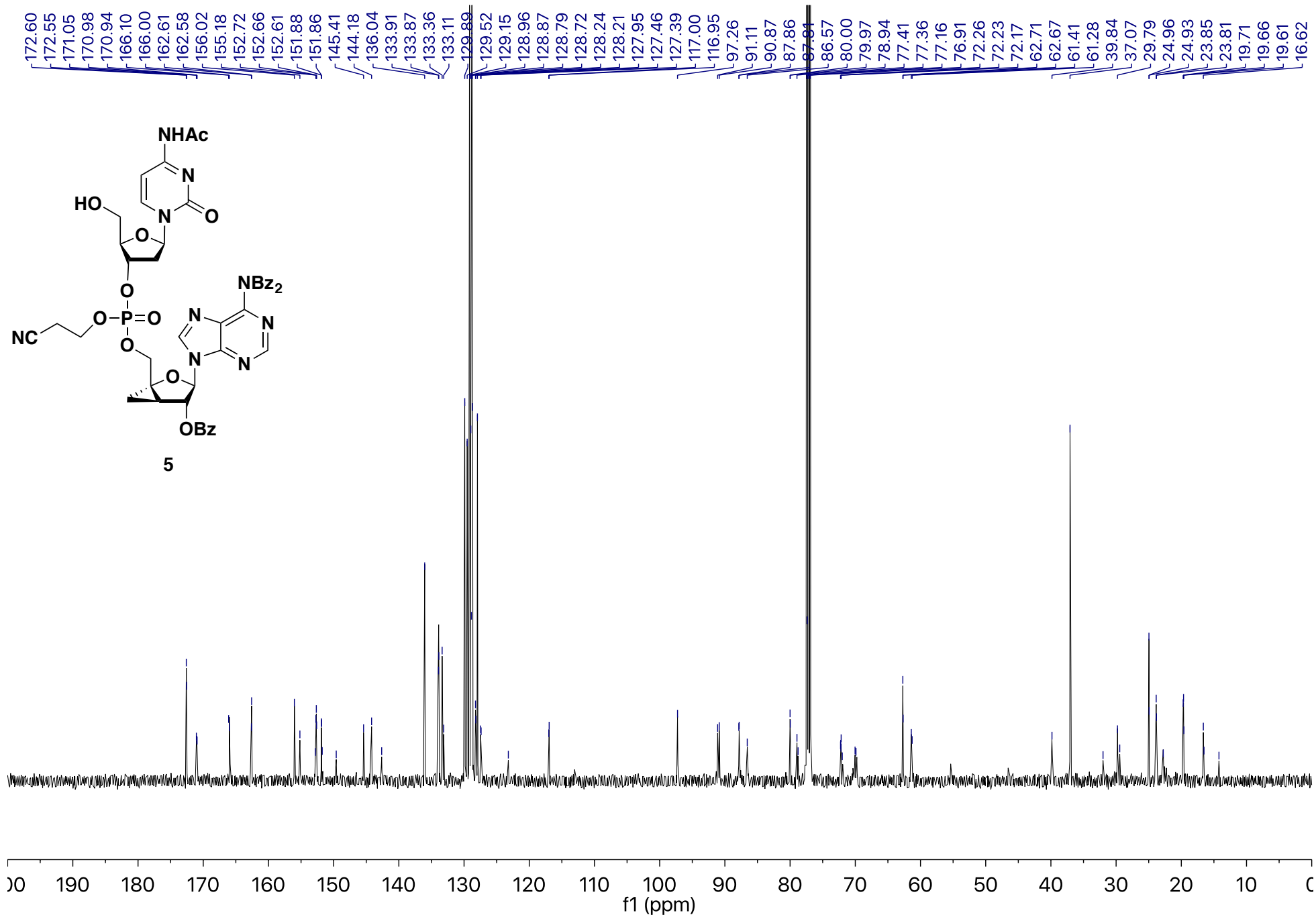


¹H, CDCl₃, 500 MHz

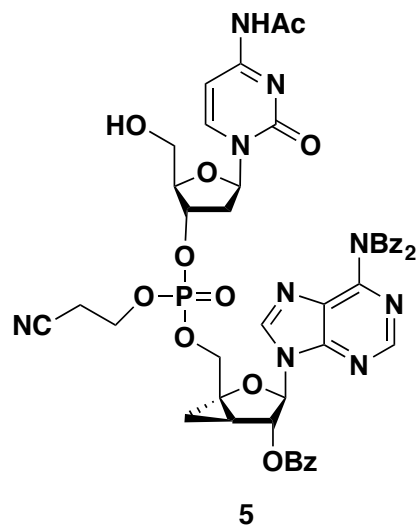




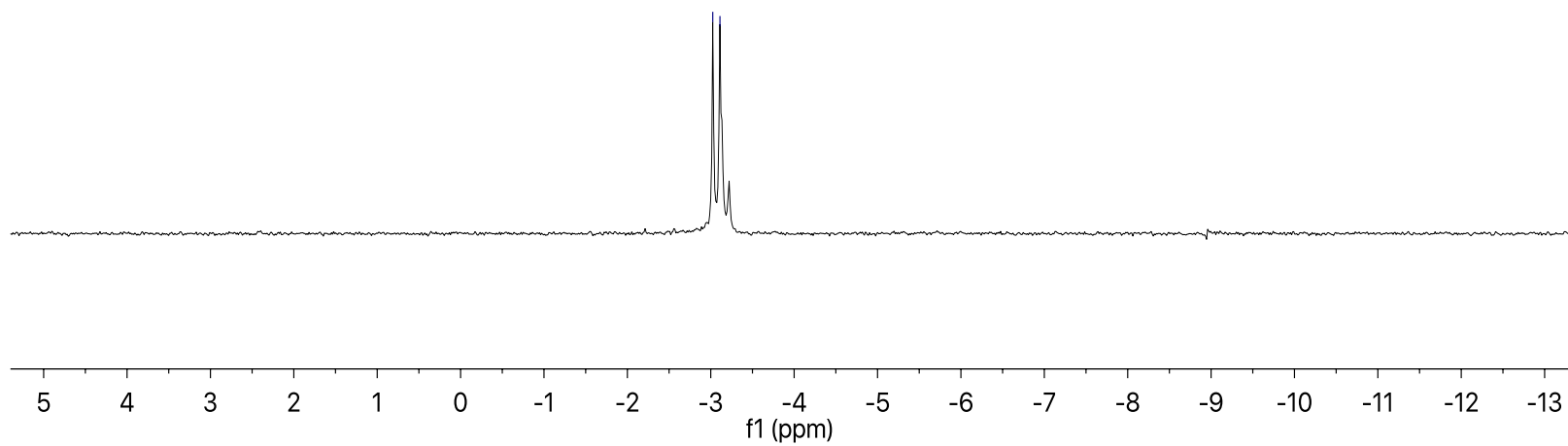
¹H, CDCl₃, 500 MHz



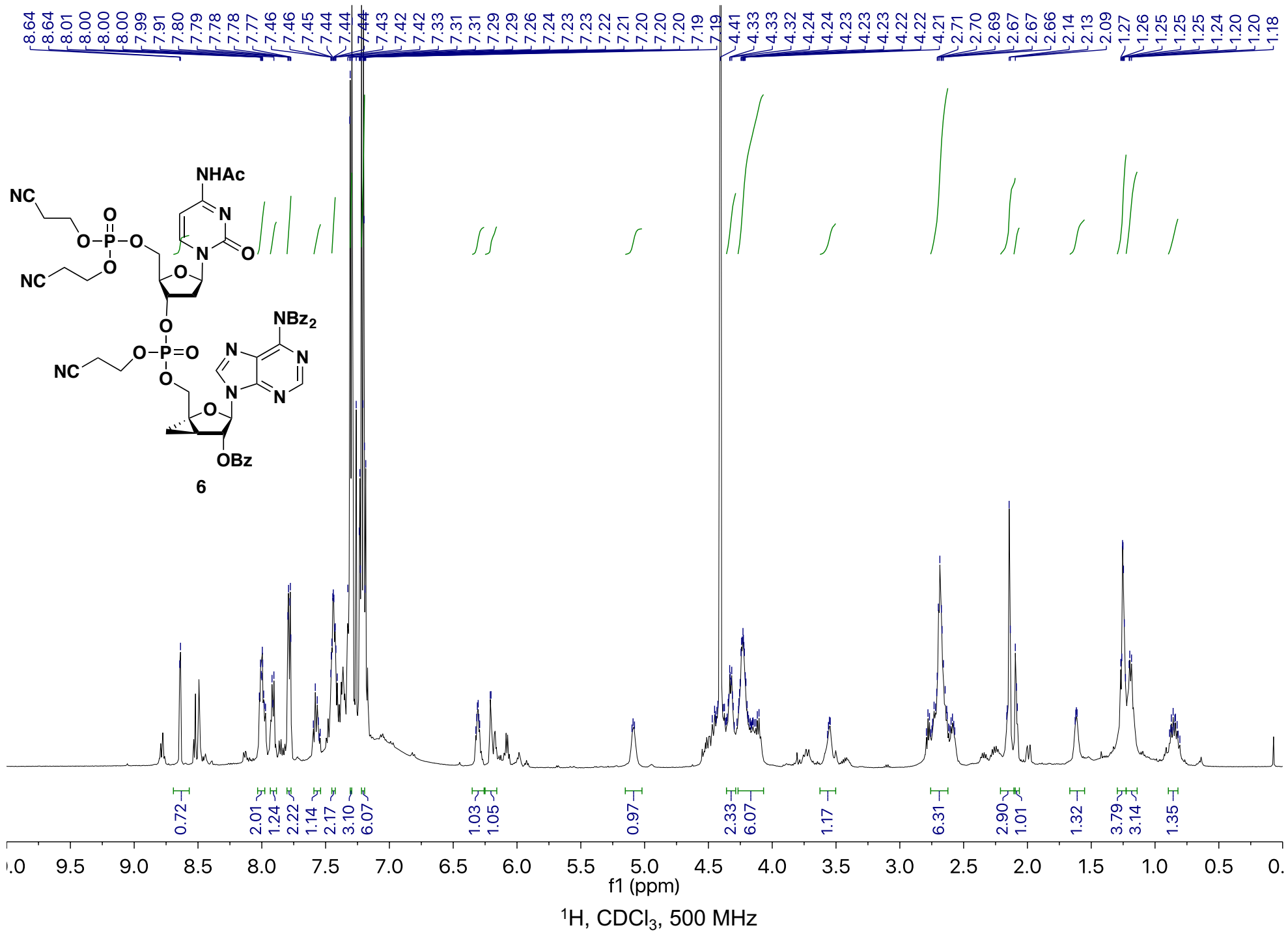
¹³C, CDCl₃, 125 MHz

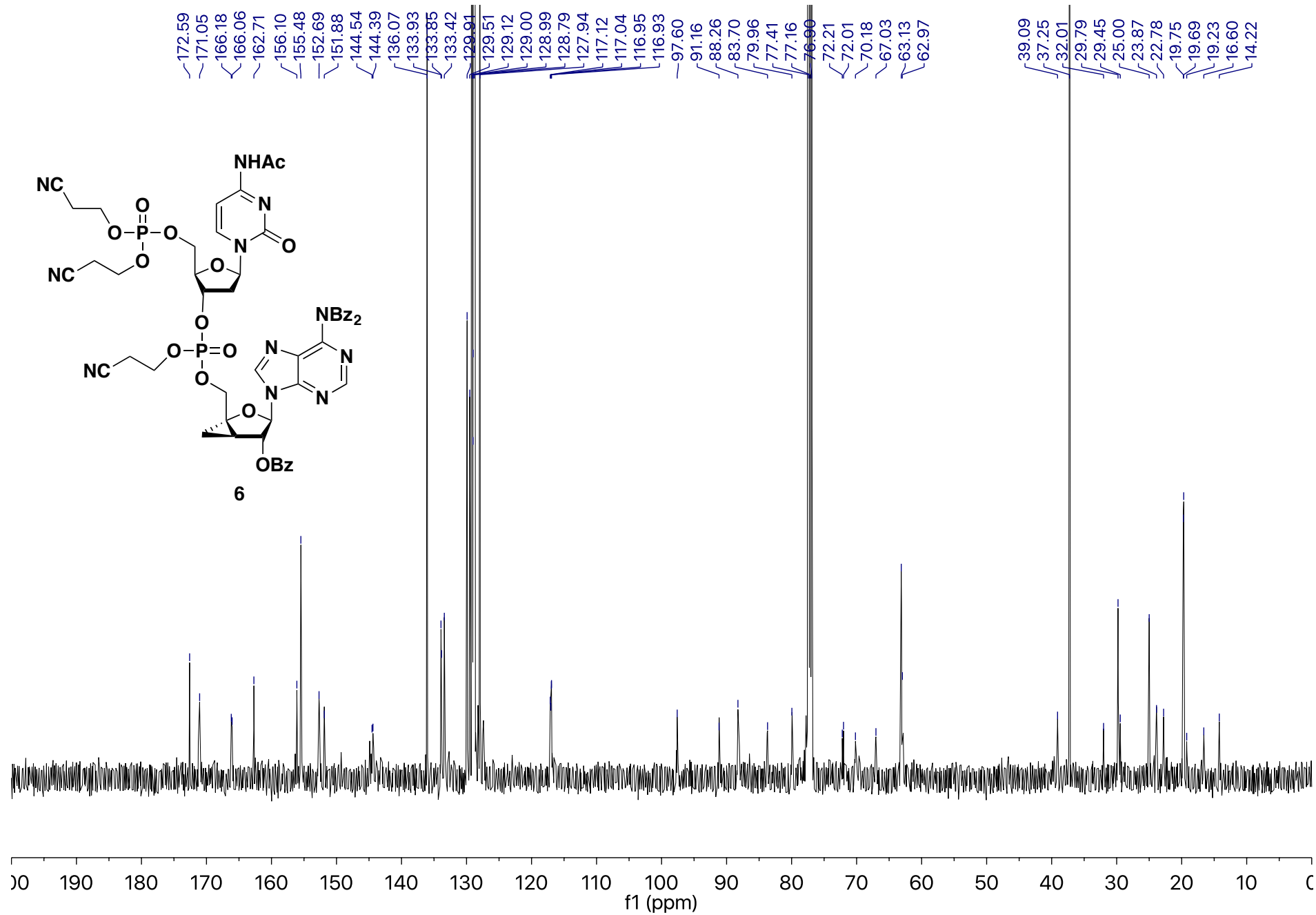


3.02
3.11

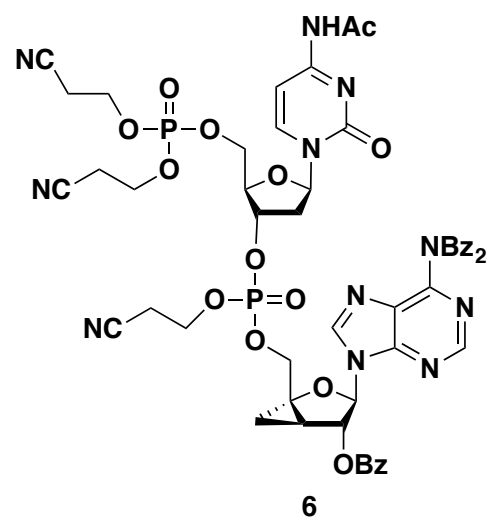


³¹P, CDCl₃, 203 MHz

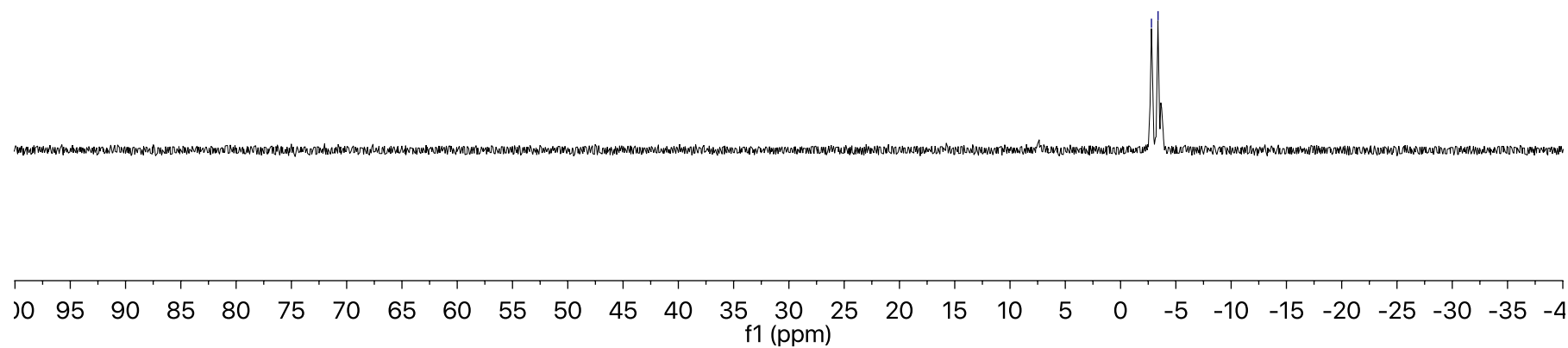




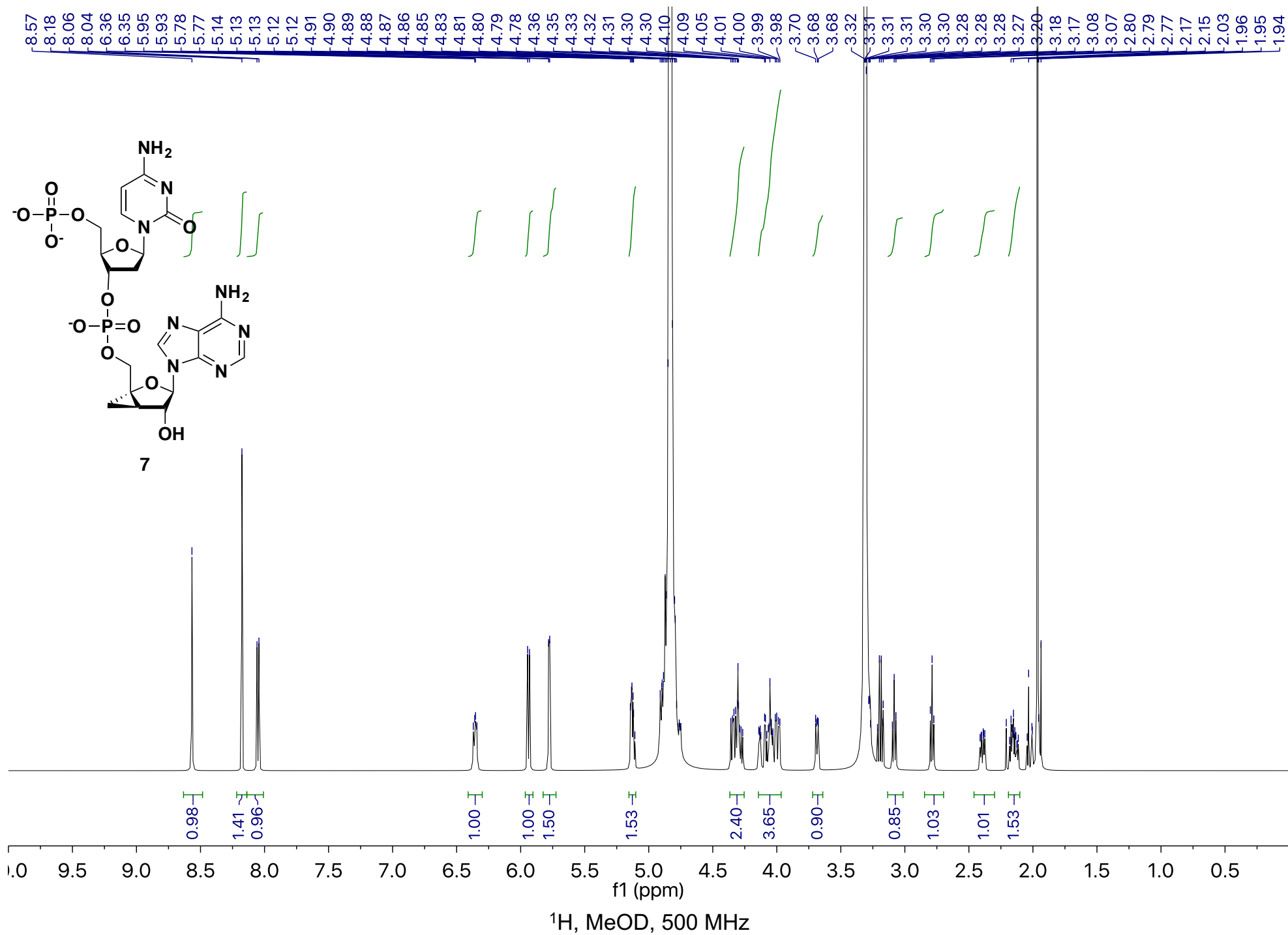
¹³C, CDCl₃, 125 MHz

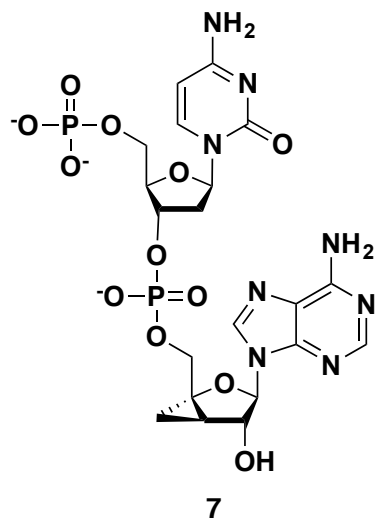


~2.79
~3.39

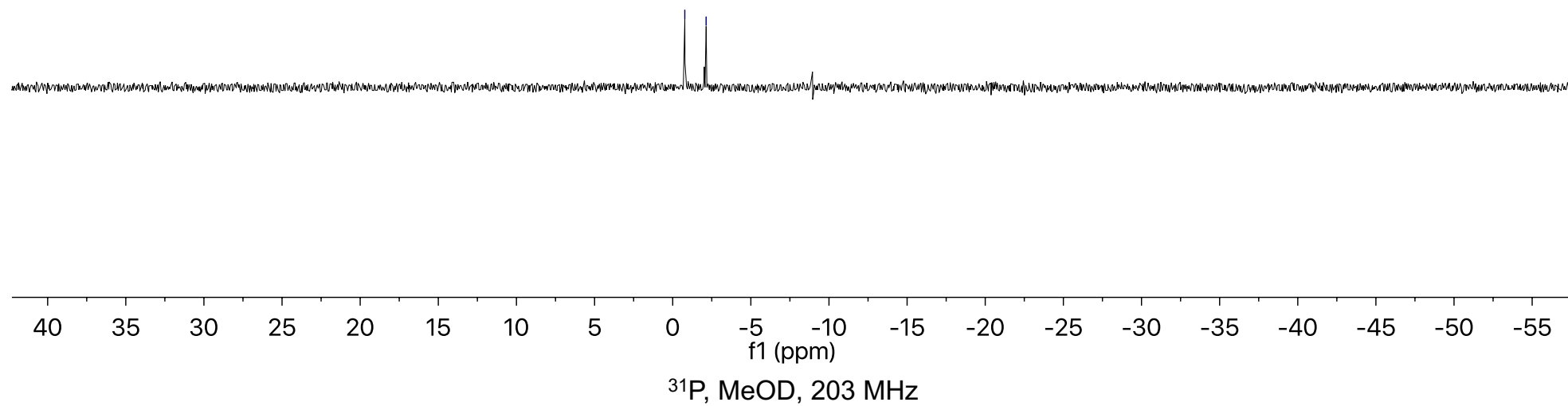


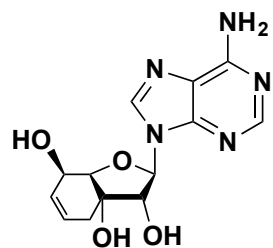
³¹P, CDCl₃, 203 MHz



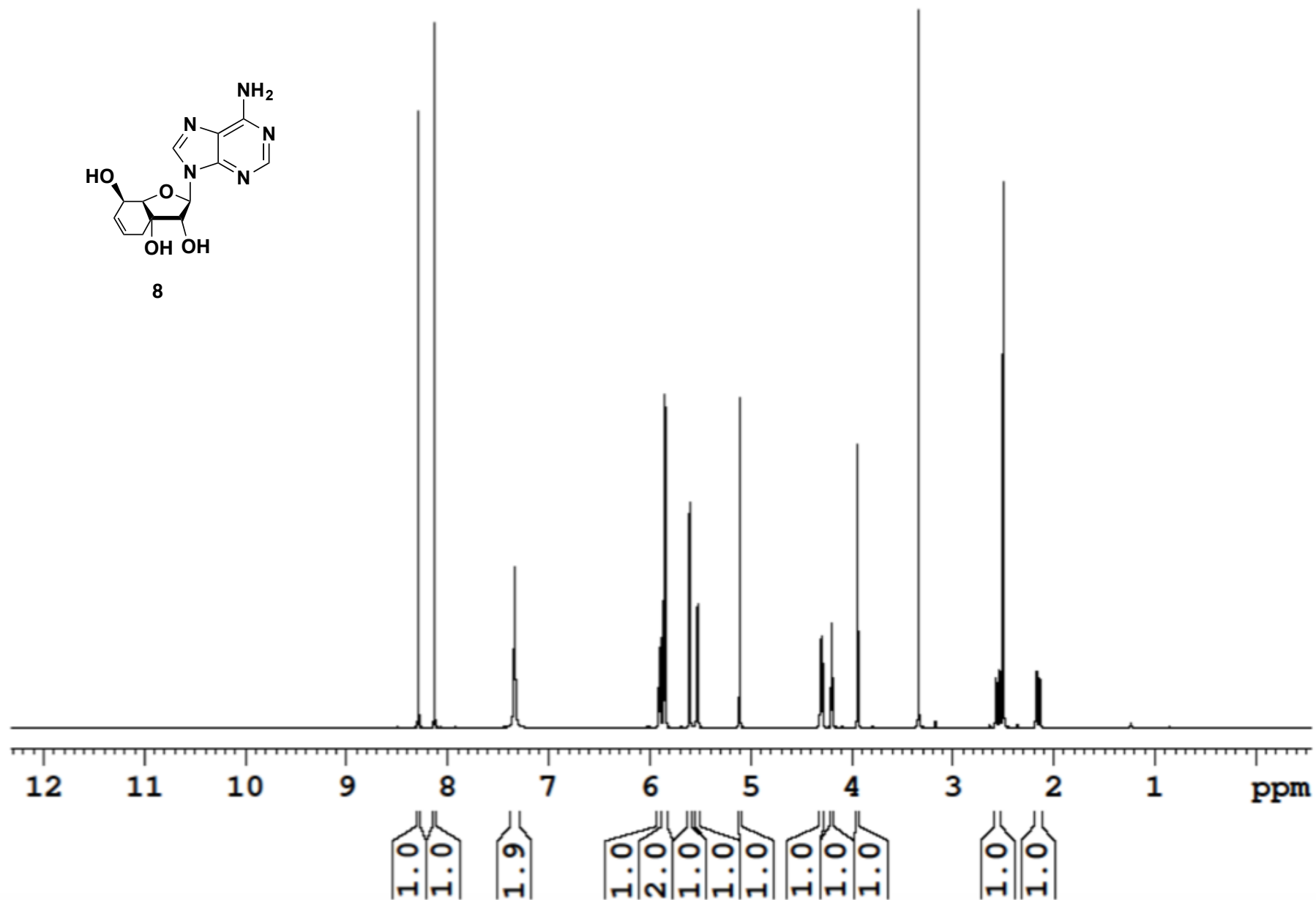


--0.77
--2.14

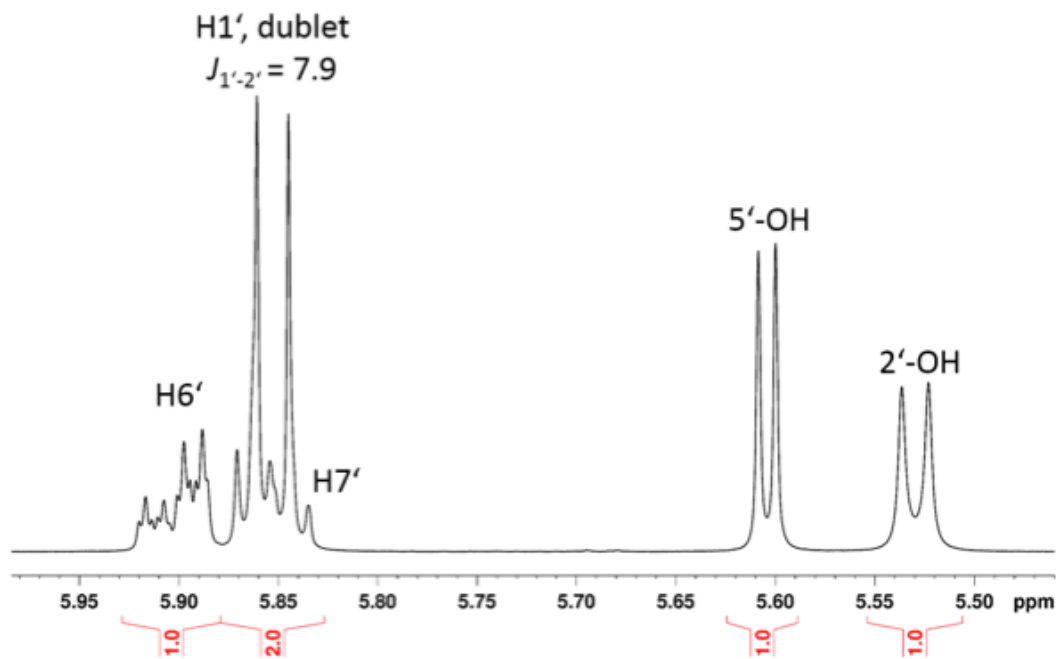
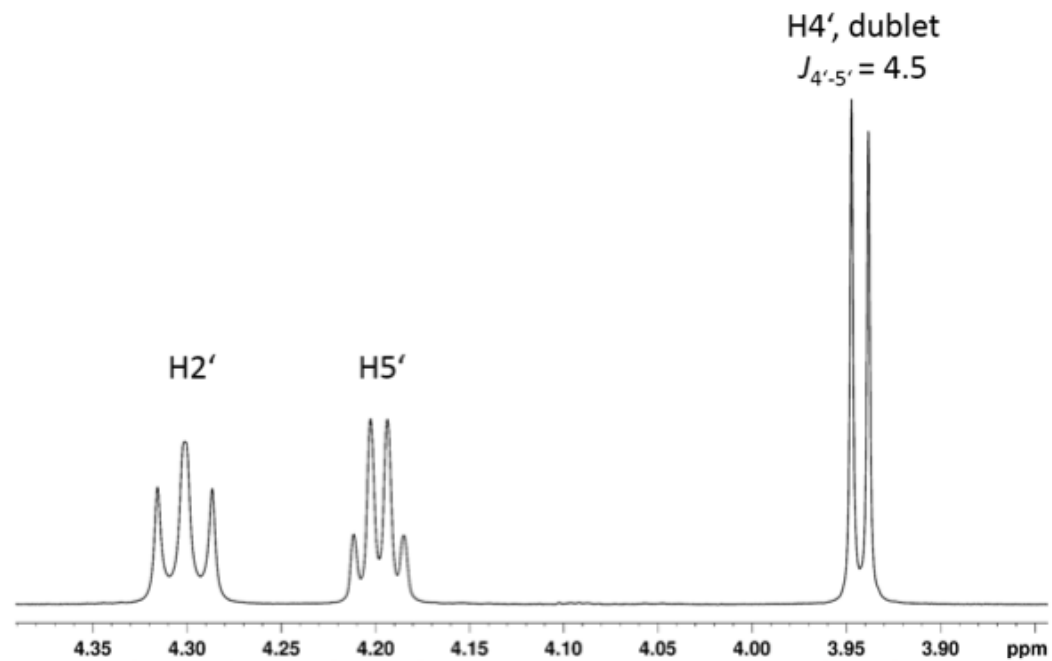
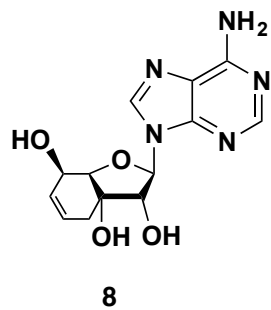




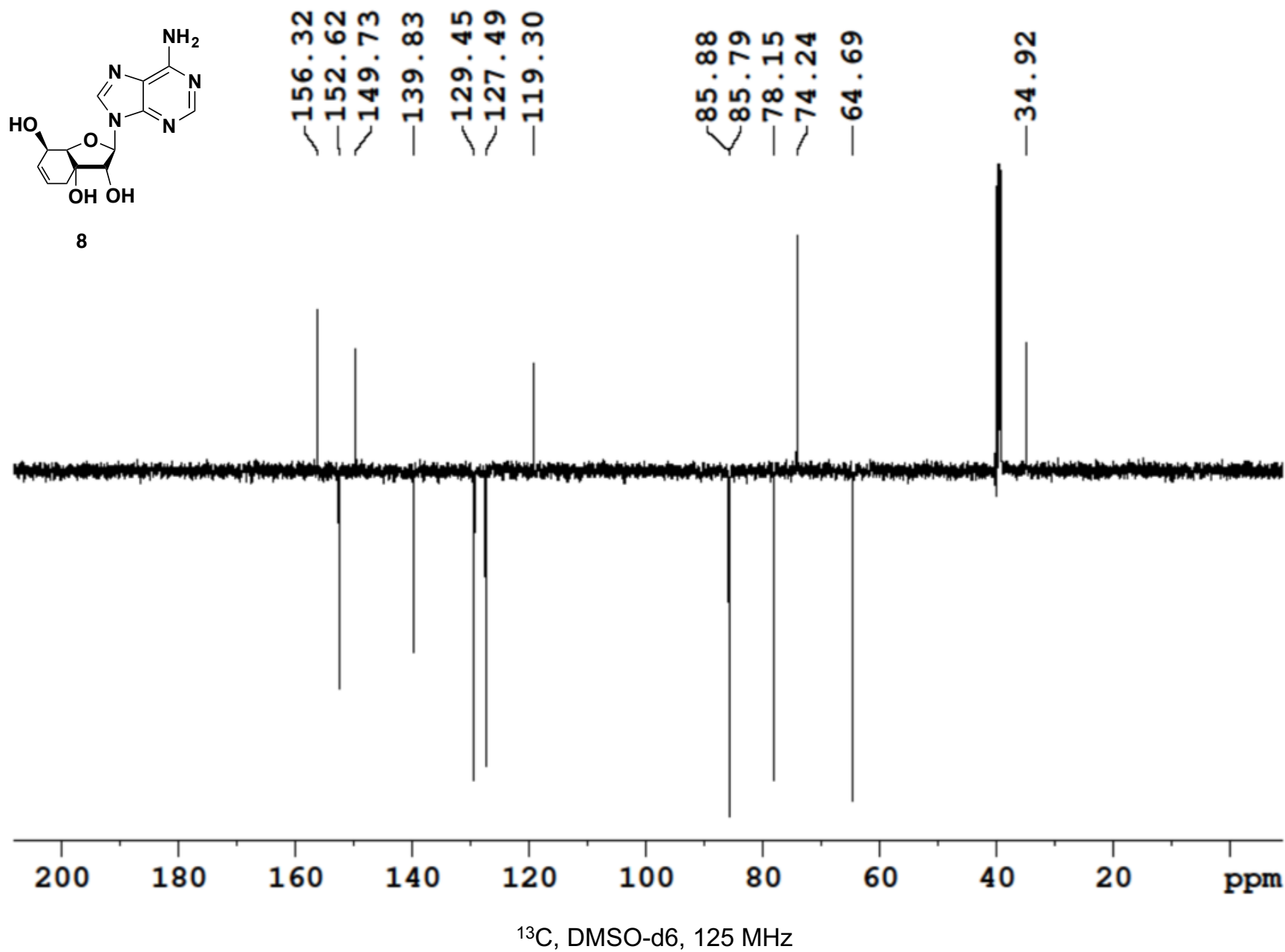
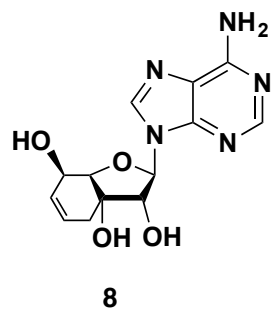
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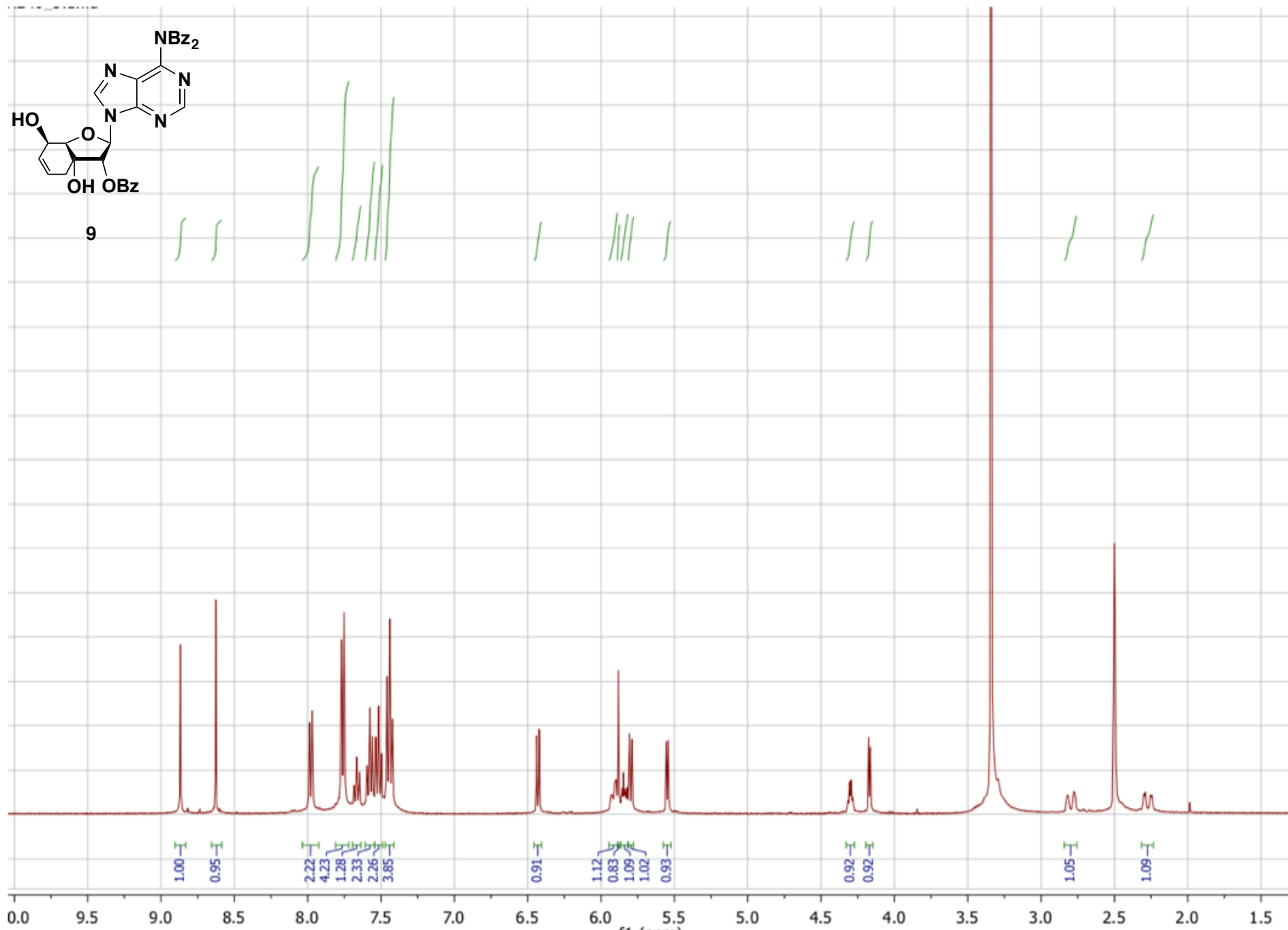


¹H, DMSO-d₆, 500 MHz

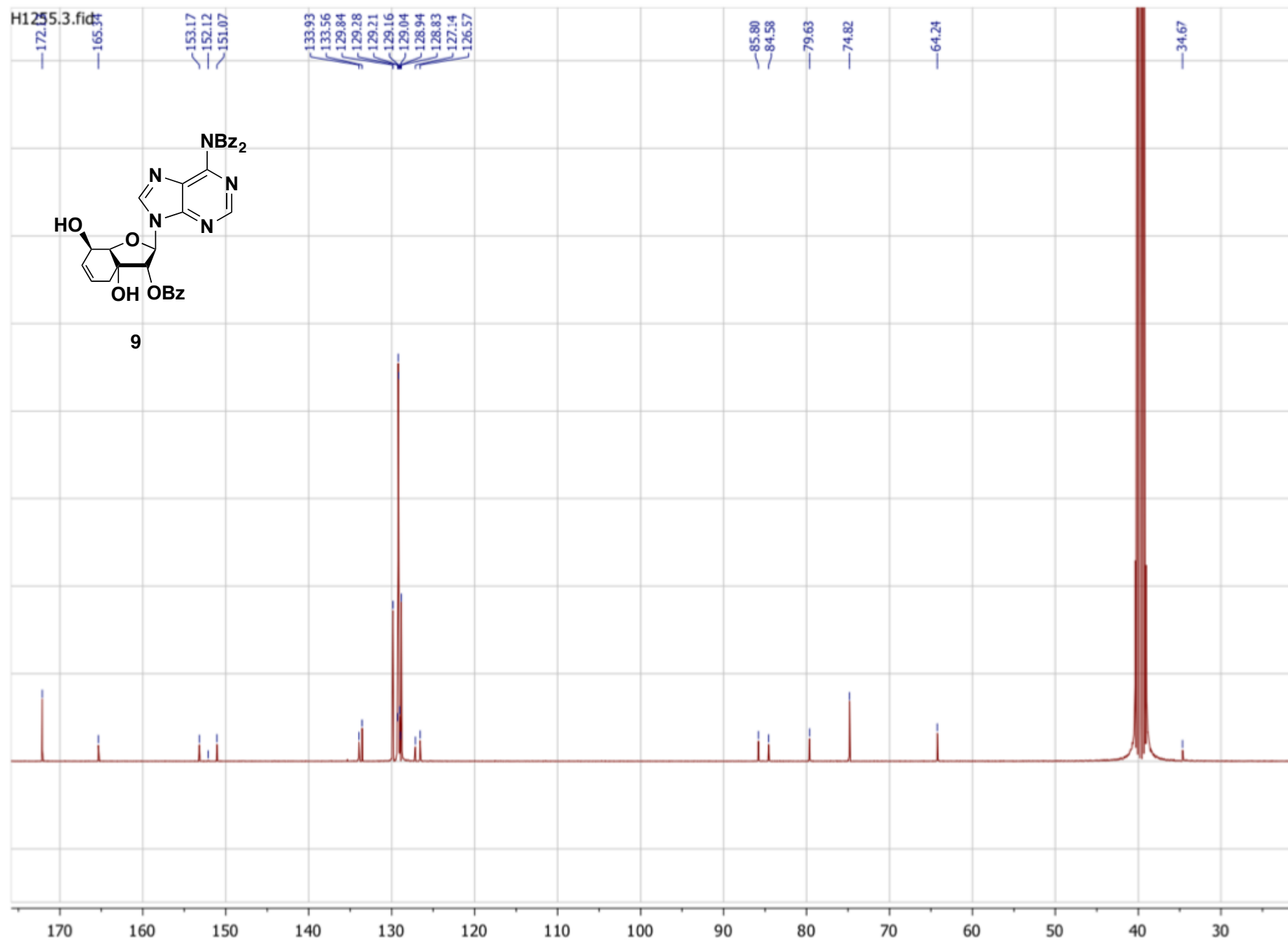


^1H , DMSO- d_6 , 500 MHz

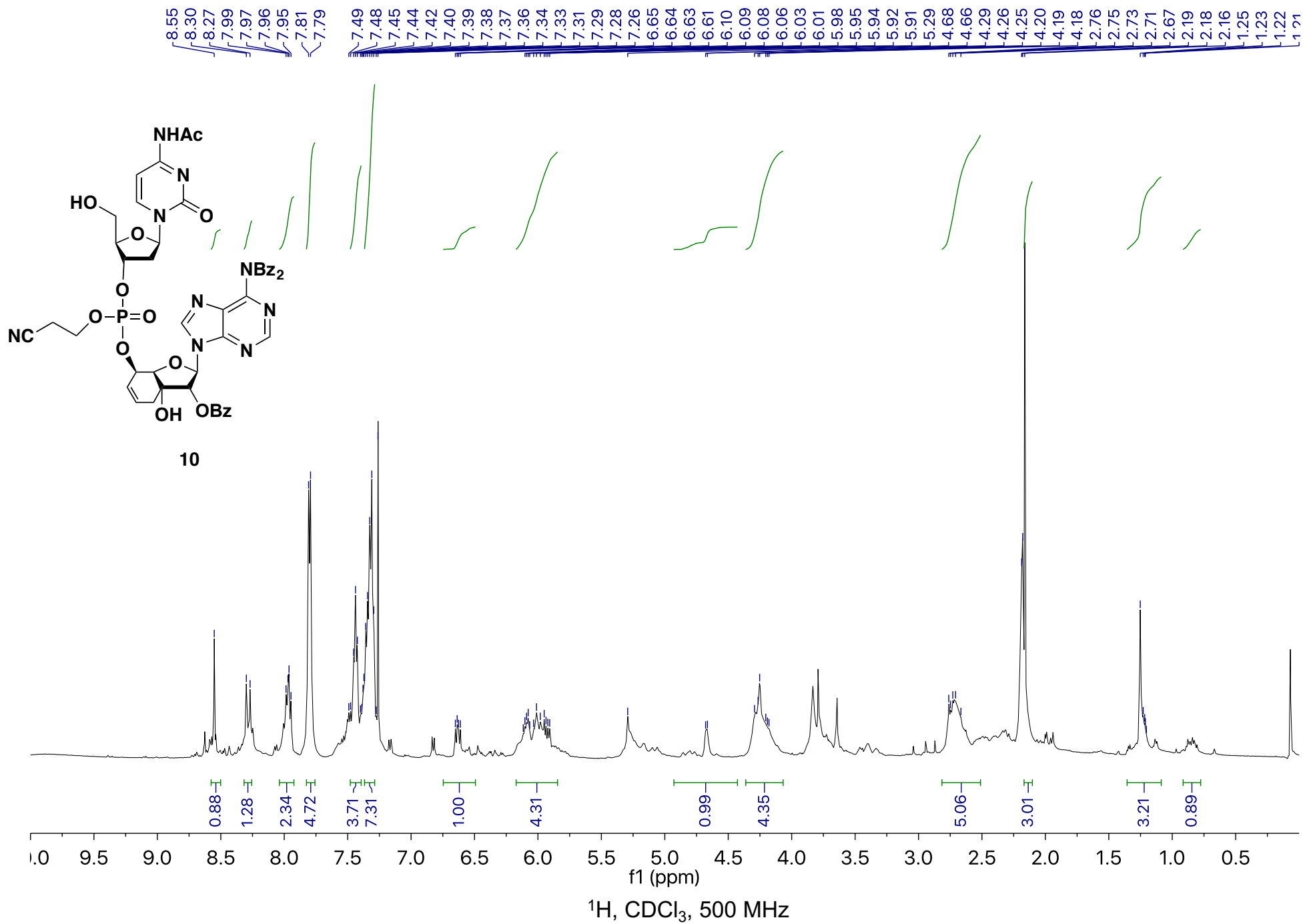


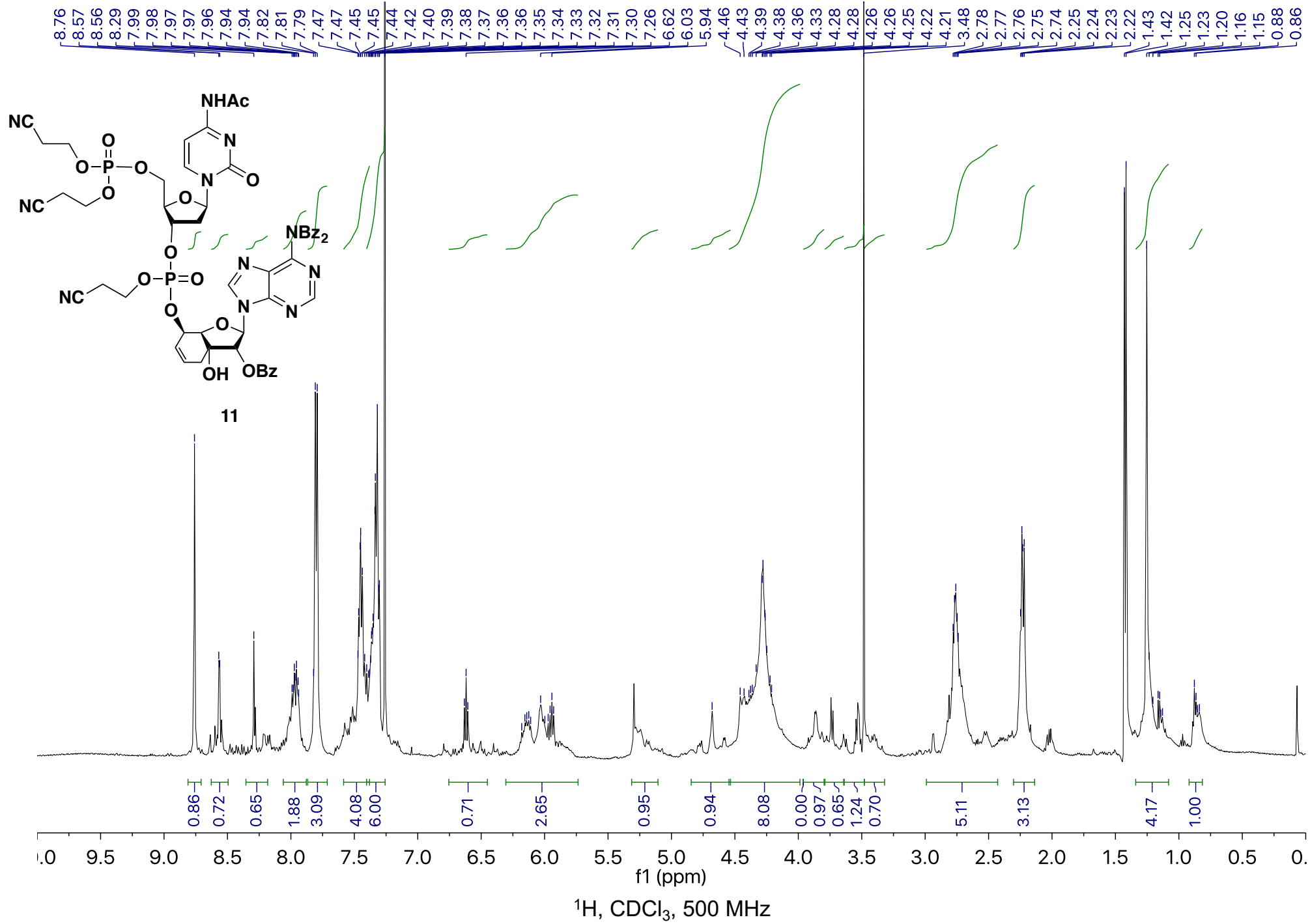


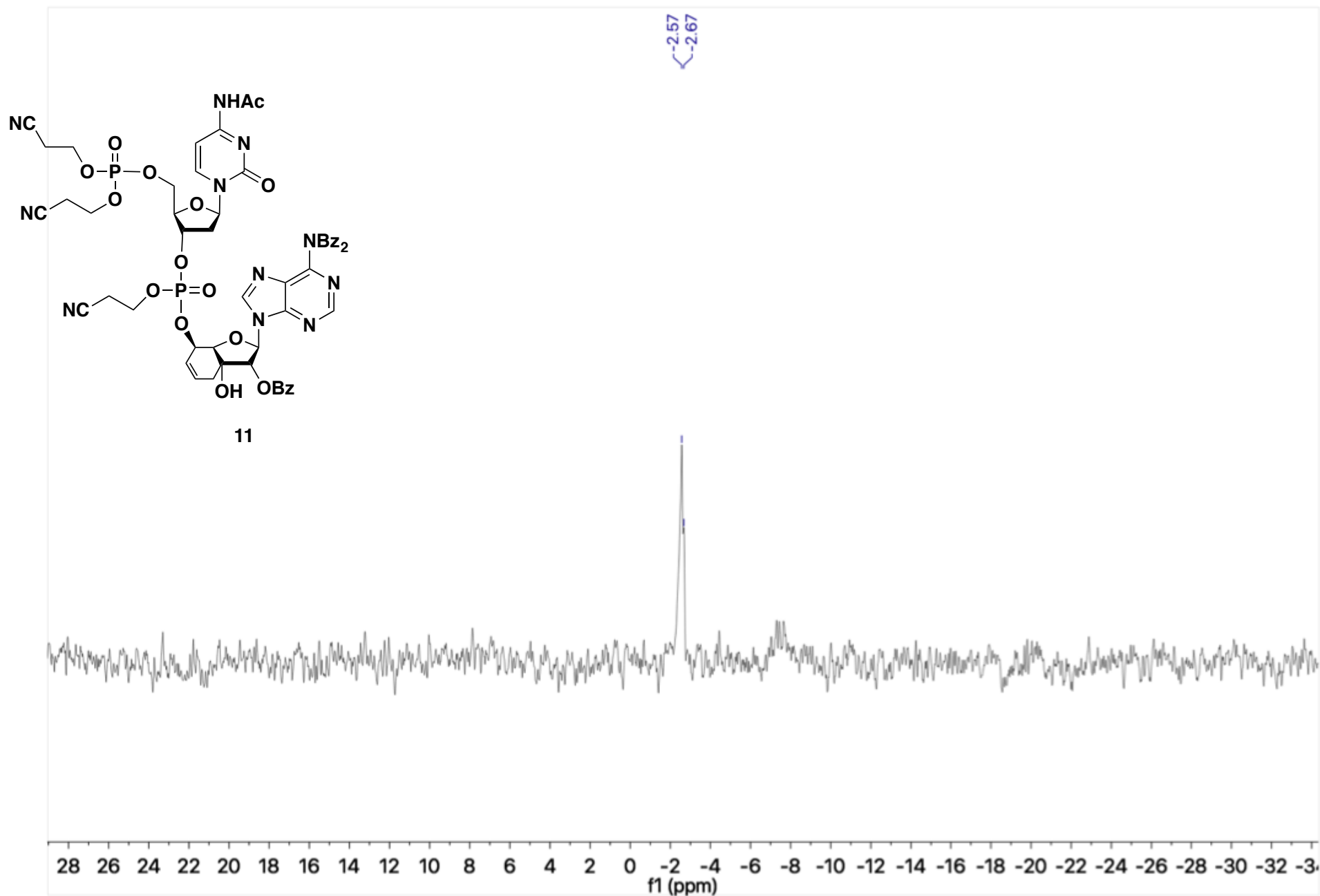
¹H, DMSO, 400 MHz



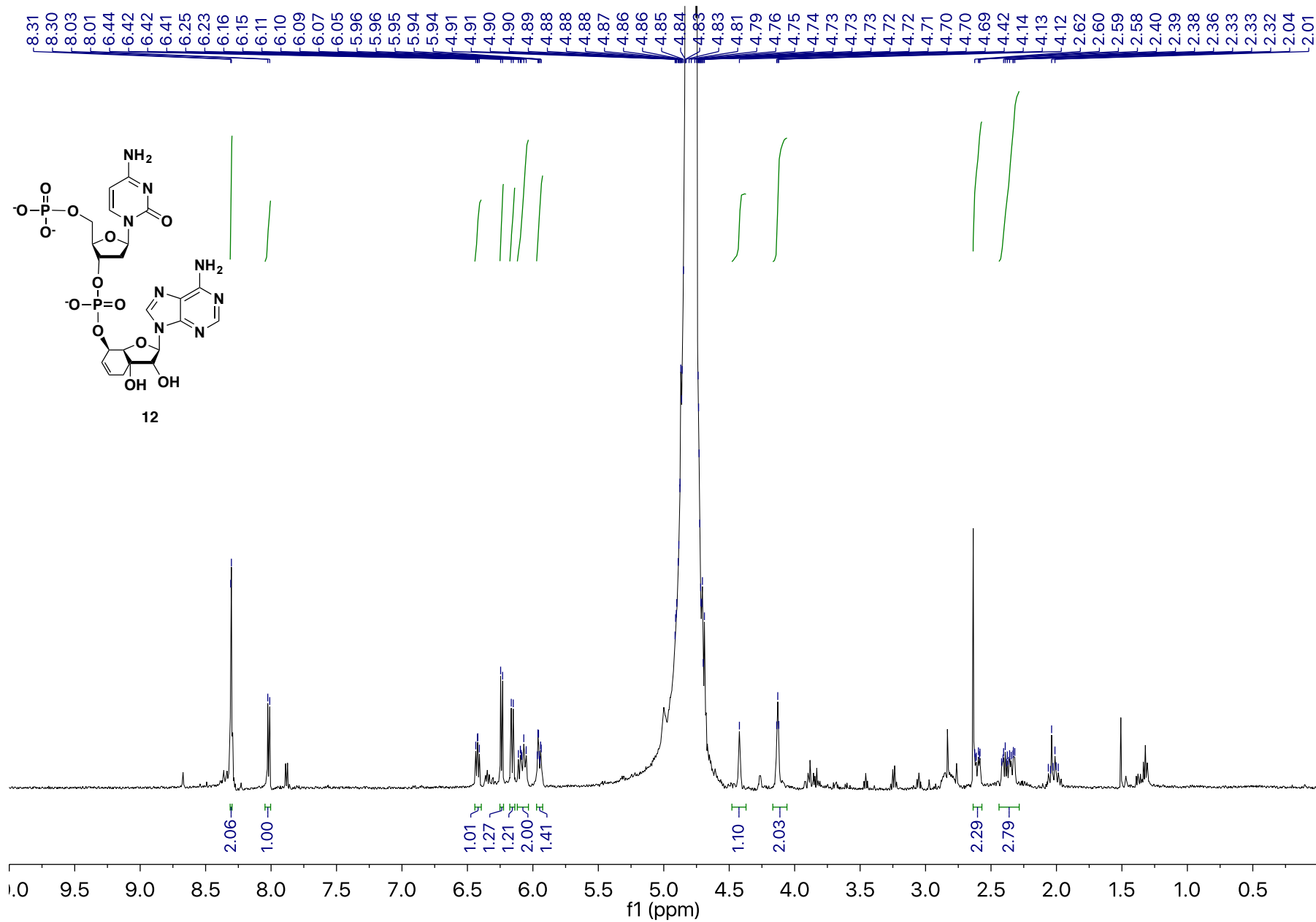
¹³C, DMSO, 100 MHz





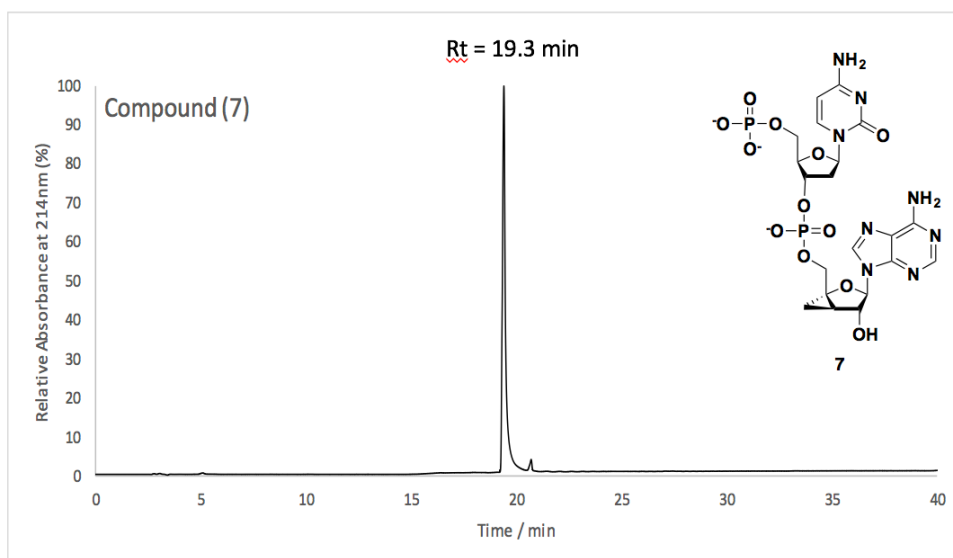


^{31}P , CDCl_3 , 101 MHz



¹H, D₂O, 500 MHz

2- *rp*HPLC analysis of dinucleotides 7 and 12

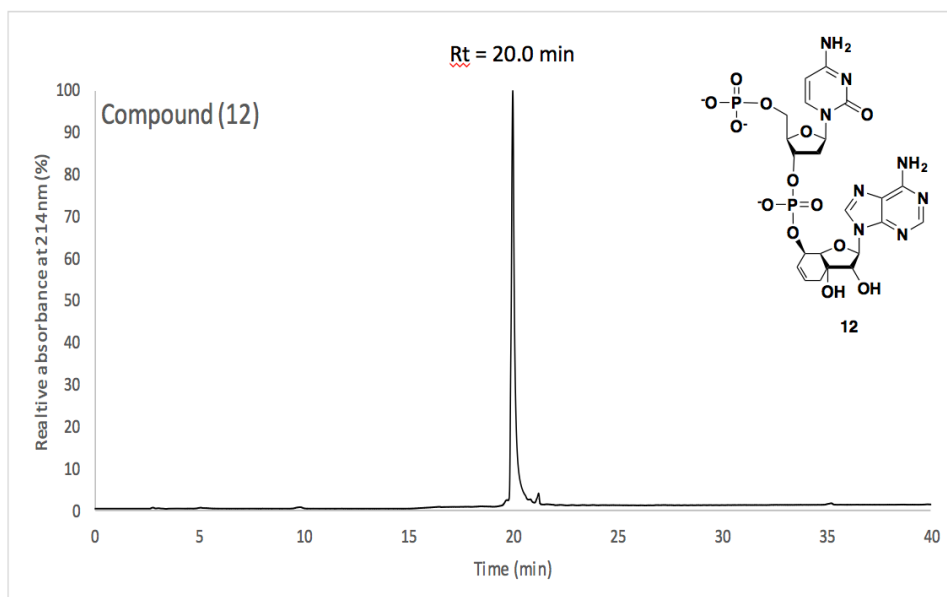


C18 nucleosil 3 μ m; 4.5x250cm (Macherey Nagel)

Gradient: 10.5min A; 0 to 100% B in 30min

A: 0.1%TFA

B:ACN/0.1%TFA



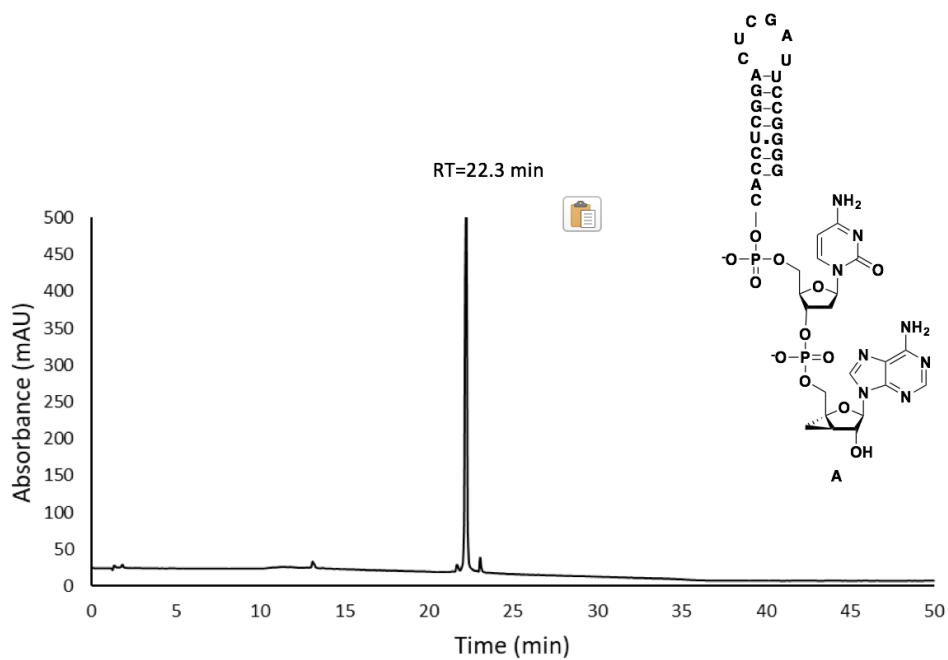
C18 nucleosil 3 μ m; 4.5x250cm (Macherey Nagel)

Gradient: 10.5min A; 0 to 100% B in 30min

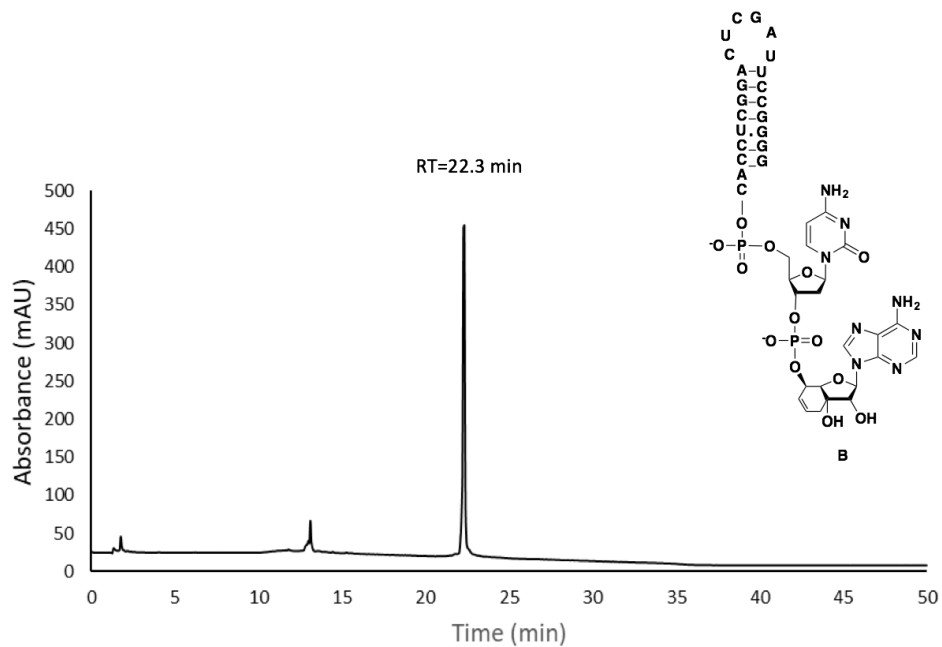
A: 0.1%TFA

B: ACN/0.1%TFA

3- *rp*HPLC analysis of tRNAs A and B



Anionic exchange *rp*HPLC chromatography on a DNAPac PA100 (Thermo Scientific) column. Flow: 1mL.min⁻¹; UV detection at 260nm
 500pmol of each product injected
 Buffer A: Ammonium acetate 25 mM pH 8,0 ; 0,5% ACN
 Buffer B: Ammonium acetate 2500 mM pH 8,0 ; 0,5% ACN
 8,4min at 0%A then a linear 30min gradient from 0 to 100% buffer B.



Anionic exchange *rp*HPLC chromatography on a DNAPac PA100 (Thermo Scientific) column. Flow: 1mL.min⁻¹; UV detection at 260nm
 500pmol of each product injected
 Buffer A: Ammonium acetate 25 mM pH 8,0 ; 0,5% ACN
 Buffer B: Ammonium acetate 2500 mM pH 8,0 ; 0,5% ACN
 8,4min at 0%A then a linear 30min gradient from 0 to 100% buffer B.

4- FemX_{wv} inhibition experiments

