

New benzophenone phosphonate derivatives

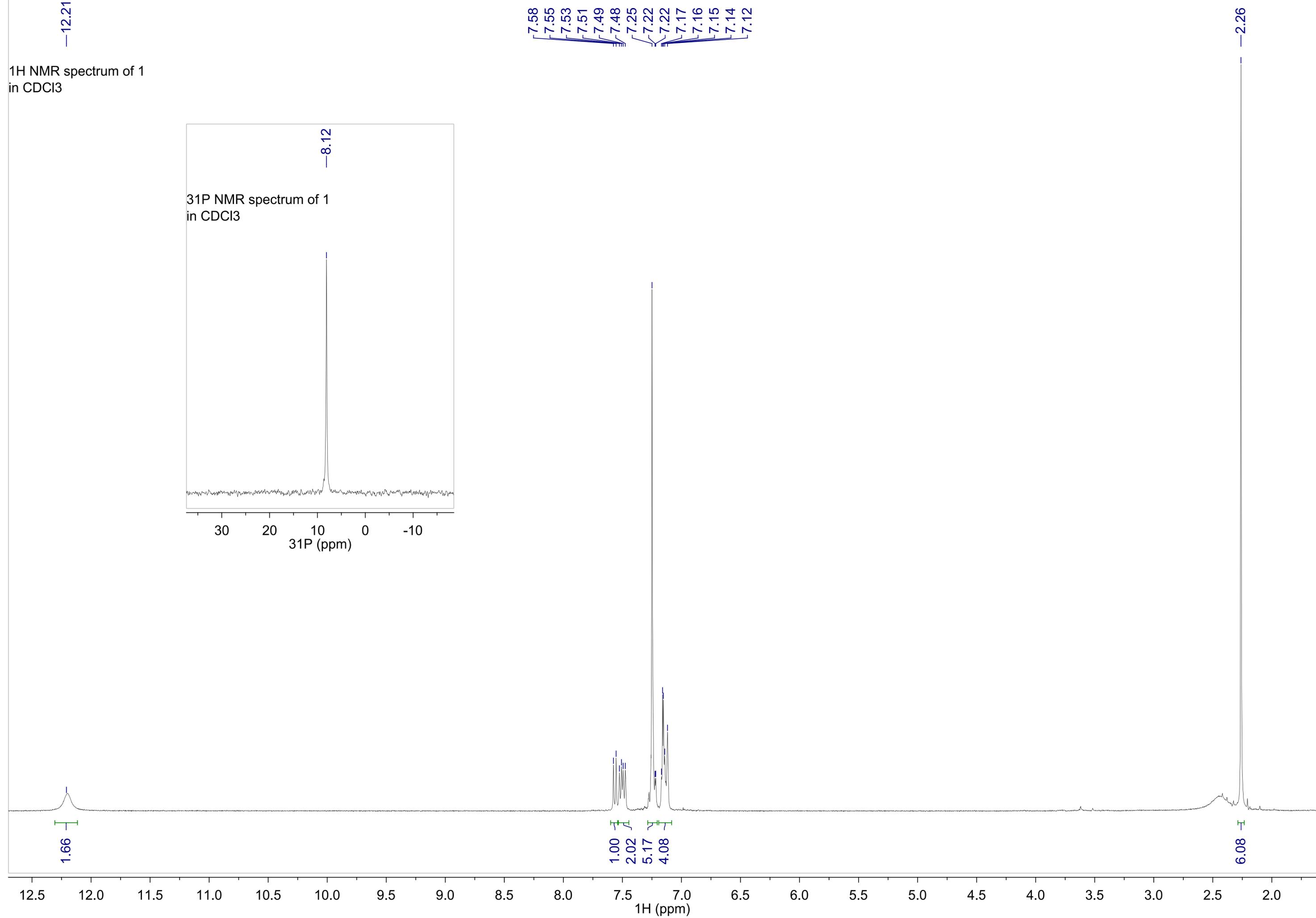
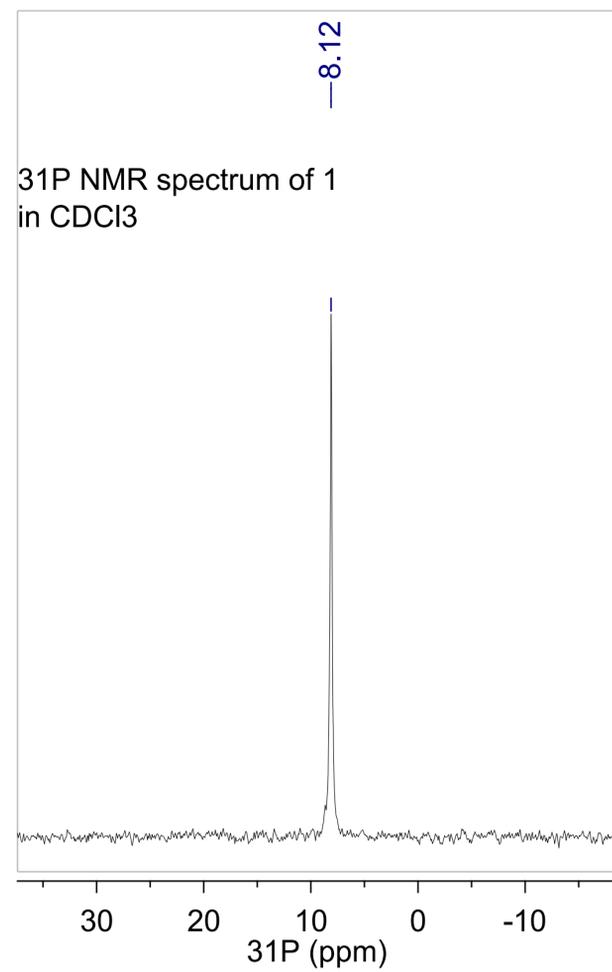
**Maxim S. Gaman, Elena S. Matyugina, Mikhail S. Novikov, Denis A. Babkov,
Pavel N. Solyev, Sergey N. Kochetkov and Anastasia L. Khandazhinskaya**

The reactions were performed with the use of commercial reagents (Acros, Aldrich, and Fluka); anhydrous solvents were purified according to the standard procedures. Column chromatography was performed on Silica Gel 60 0.040–0.063 mm (Merck, Germany) columns, DEAE Toyopearl cellulose and Dowex-50 (H⁺). Preparative liquid chromatography (PLC) was performed on Silica Gel 60 F₂₅₄ with concentrating zone glass plates (Merck, Germany). Thin layer chromatography (TLC) was performed on Silica Gel 60 F₂₅₄ aluminium-backed plates (Merck, Germany).

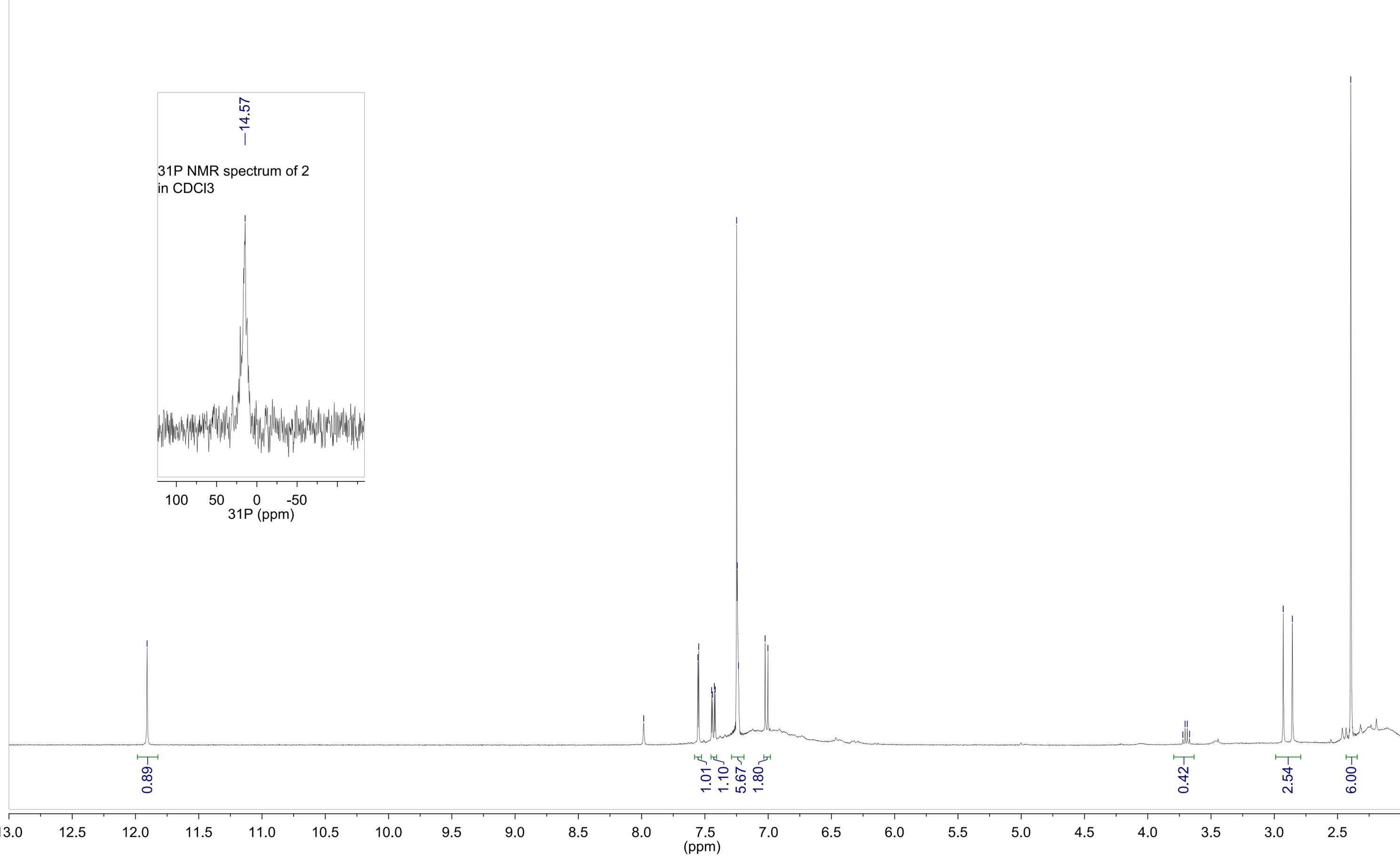
NMR spectra were registered on a Bruker Avance 400 spectrometer («Bruker», Newark, DE) with the working frequency of 400 MHz for ¹H (Me₄Si as internal standard) and 121 MHz for ³¹P (with phosphorus-proton interaction decoupling, 85% phosphoric acid as an external standard) in CDCl₃ or DMSO-d₆.

High resolution mass spectra (HRMS) were registered on a Bruker Daltonics micrOTOF-Q II instrument using electrospray ionization. The measurements were acquired in a negative ion mode with the following parameters: interface capillary voltage – 3700 V; mass range from *m/z* 50 to 3000; external calibration (Electrospray Calibrant Solution, Fluka); nebulizer pressure – 0.3 Bar; flow rate – 3 μL/min; dry gas nitrogen (4.0 L/min); interface temperature was set at 180 or 190°C. A syringe injection was used.

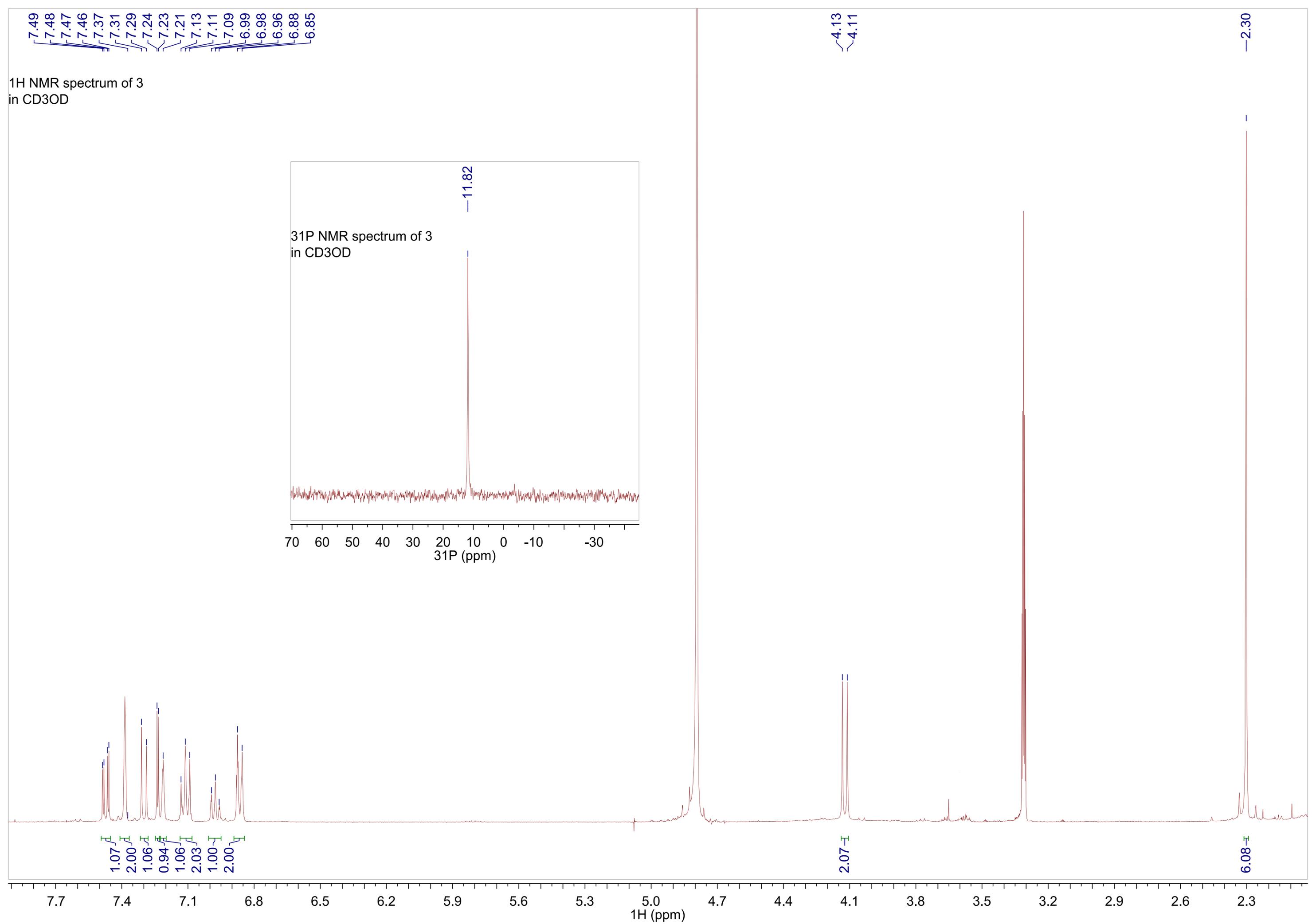
¹H NMR spectrum of 1
in CDCl₃



¹H NMR spectrum of 2
in CDCl₃

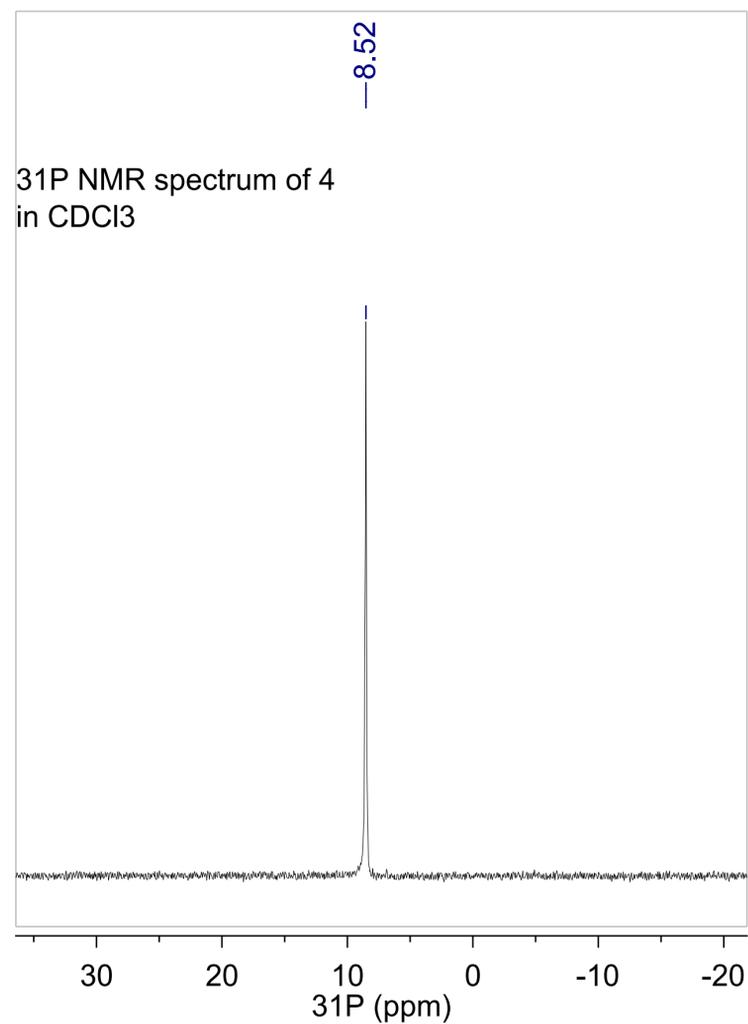


¹H NMR spectrum of 3
in CD₃OD



¹H NMR spectrum of 4
in CDCl₃

7.10
7.08
7.05
7.00
6.99
6.88
6.87



4.69
4.67

4.21
4.19

2.18

4.12
2.10
5.10

2.00

2.25

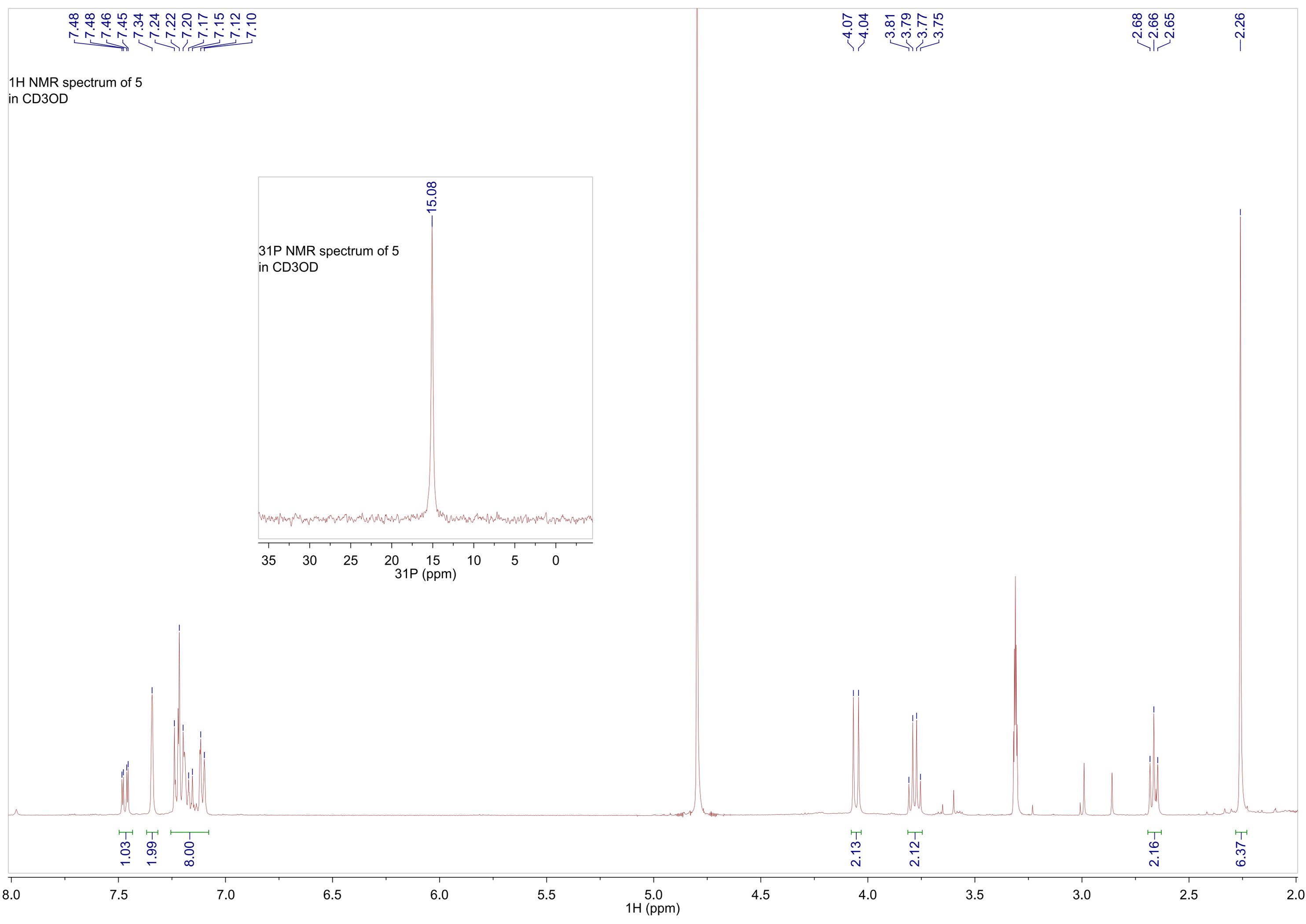
6.44

7.6 7.4 7.2 7.0 6.8 6.6 6.4 6.2 6.0 5.8 5.6 5.4 5.2 5.0 4.8 4.6 4.4 4.2 4.0 3.8 3.6 3.4 3.2 3.0 2.8 2.6 2.4 2.2 2.0 1.8 1.6

¹H (ppm)

Detailed description: The main ¹H NMR spectrum shows several multiplets in the aromatic region (6.8-7.2 ppm) and two multiplets in the aliphatic region (4.1-4.7 ppm). Integration values are shown below the peaks: 4.12, 2.10, 5.10 for the aromatic region; 2.00 and 2.25 for the aliphatic region; and 6.44 for the peak at 2.18 ppm. The x-axis is labeled '¹H (ppm)' and ranges from 7.6 to 1.6 with major ticks every 0.2 units.

¹H NMR spectrum of 5
in CD₃OD



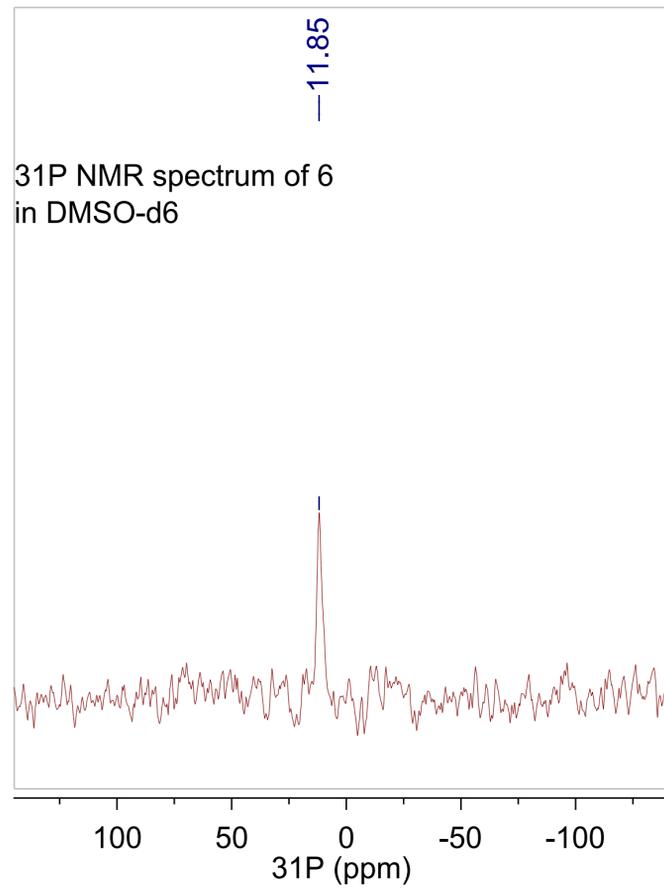
¹H NMR spectrum of 6
in DMSO-d₆

7.50
7.31
7.26
7.25

3.97

3.51

2.27



8.5 8.0 7.5 7.0 6.5 6.0 5.5 5.0 4.5 4.0 3.5 3.0 2.5 2.0

¹H (ppm)

3.01

7.89

1.89

2.12

6.00