

## Supporting Information

### Indirect Trapping of the Retro-Conjugate Addition Reaction

#### Intermediate Involved in the Epimerization of Lobeline:

#### Application to the Synthesis of (-)-Sedamine

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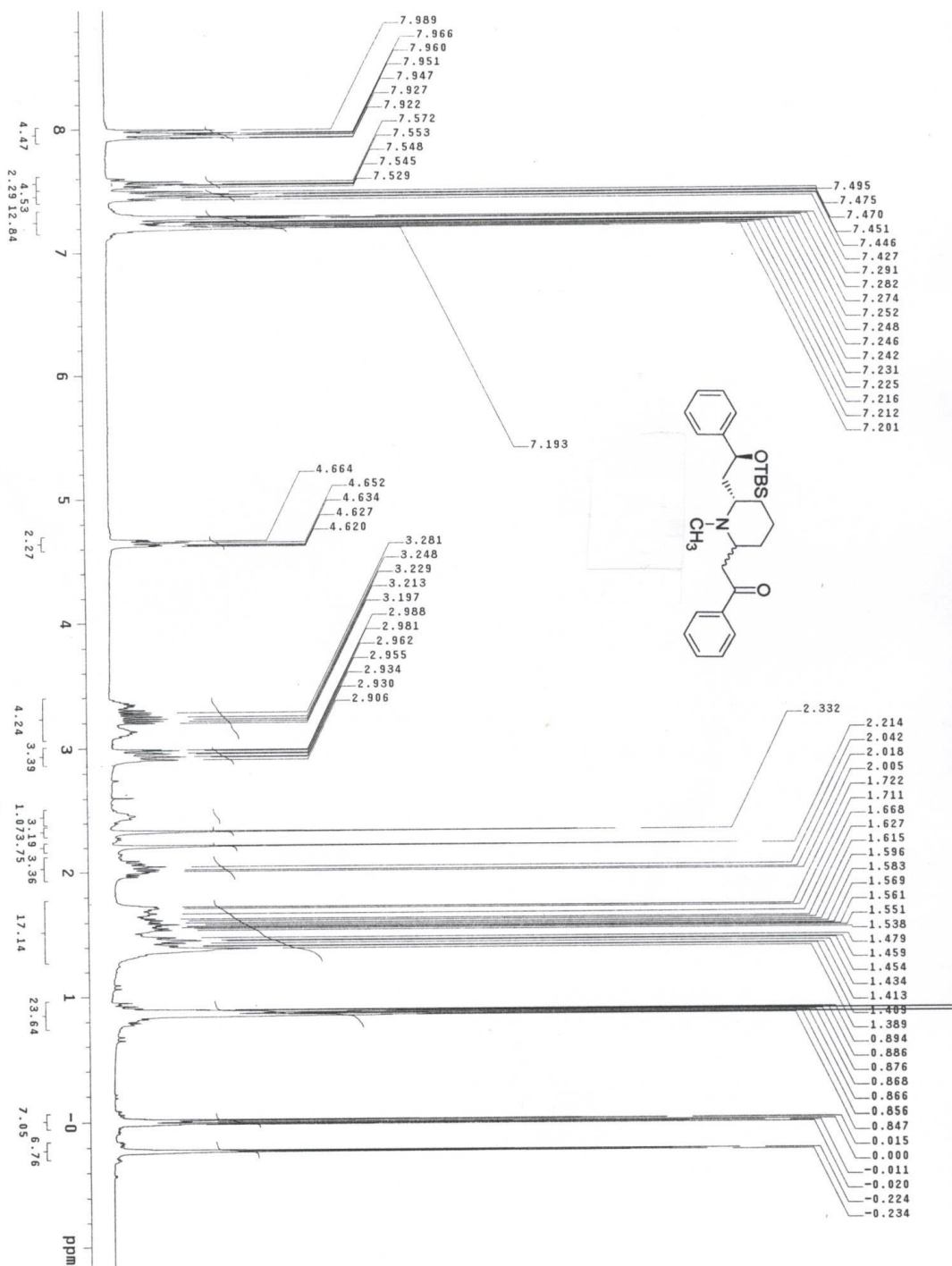
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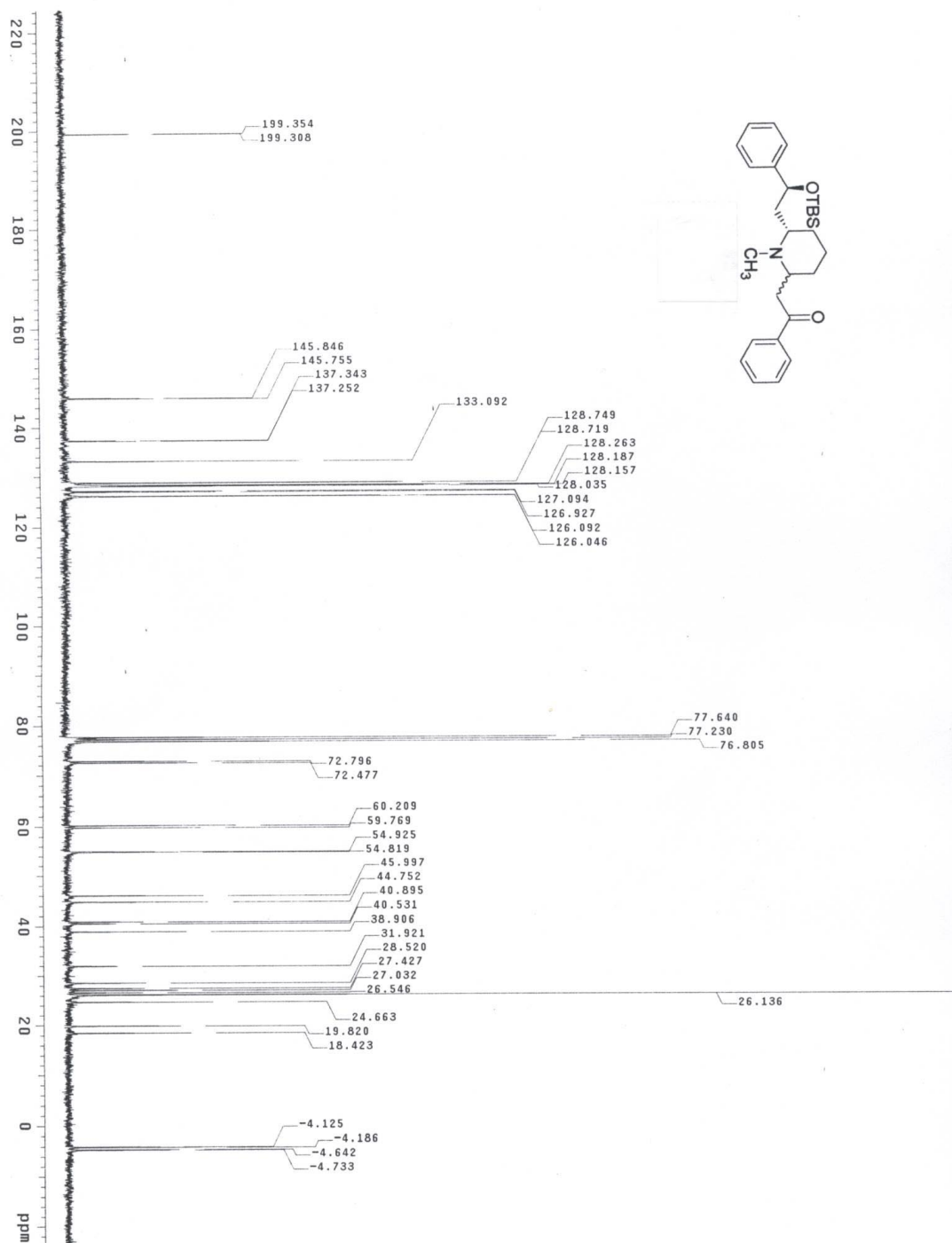
S2	General Methods
S3, S4	<sup>1</sup> H and <sup>13</sup> C NMR spectra of intermediate <b>5</b>
S5, S6	<sup>1</sup> H and <sup>13</sup> C NMR spectra of intermediate <b>6</b>
S7, S8	<sup>1</sup> H and <sup>13</sup> C NMR spectra of intermediate <b>8</b>
S9, S10	<sup>1</sup> H and <sup>13</sup> C NMR spectra of intermediate <b>9</b>
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**General Methods.** Reagents obtained from commercial sources were used without further purification. Flash column chromatography was carried out using ICN SilicTech 32-63, 60 Å silica gel. TLC analysis was carried on glass plates precoated with silica gel 60 F<sub>254</sub>. Melting points were determined on a melting point apparatus and are uncorrected. NMR data were obtained for <sup>1</sup>H at 300MHz and <sup>13</sup>C at 75 MHz using CDCl<sub>3</sub> as solvent and are reported in ppm relative to TMS as internal standard. High resolution electron impact ionization mass spectra were recorded at 70eV at a resolution of greater than 10000.

$^1\text{H}$  NMR spectrum of **5** ( $\text{CDCl}_3$ , 300 MHz)

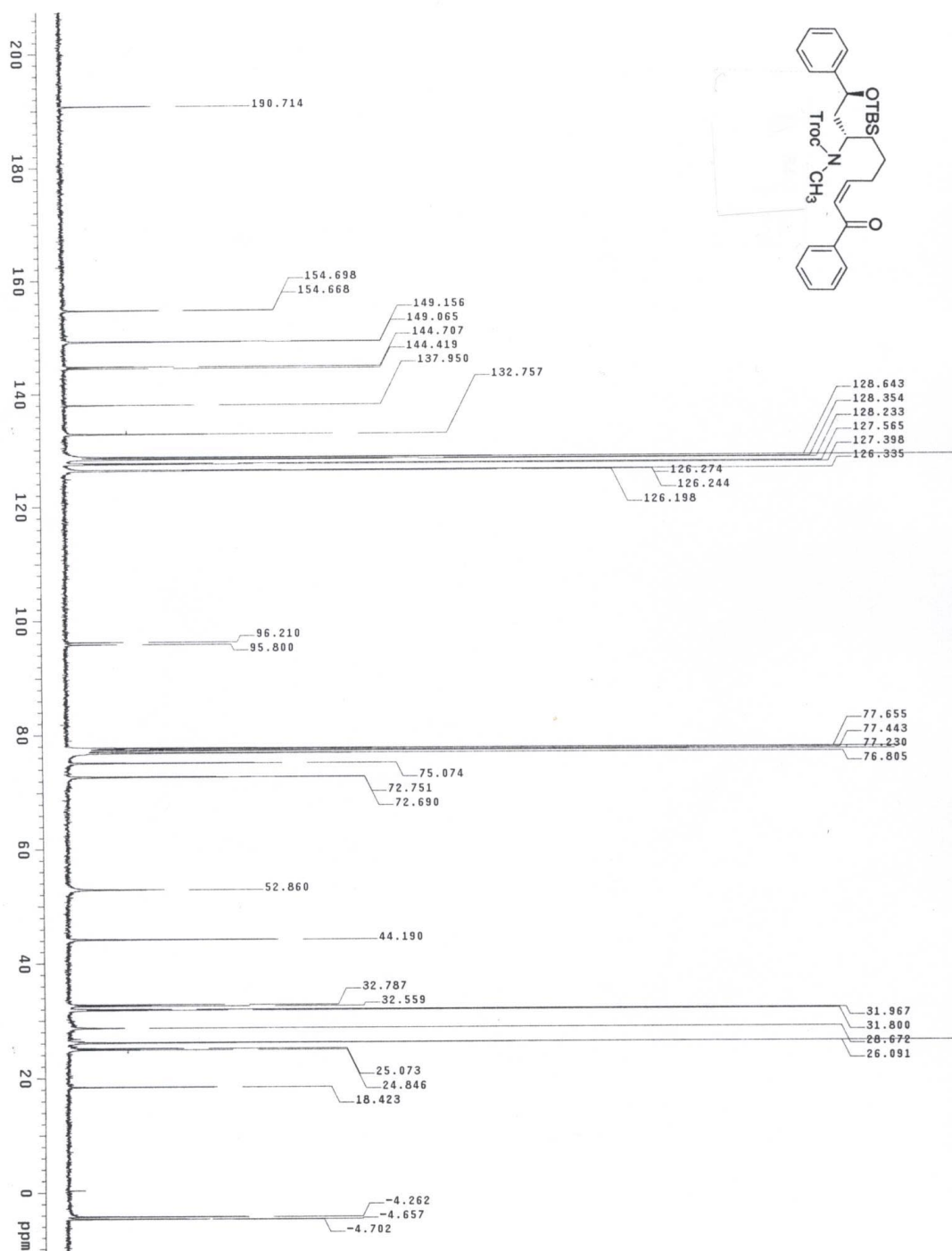


$^{13}\text{C}$  NMR spectrum of **5** ( $\text{CDCl}_3$ , 75 MHz)

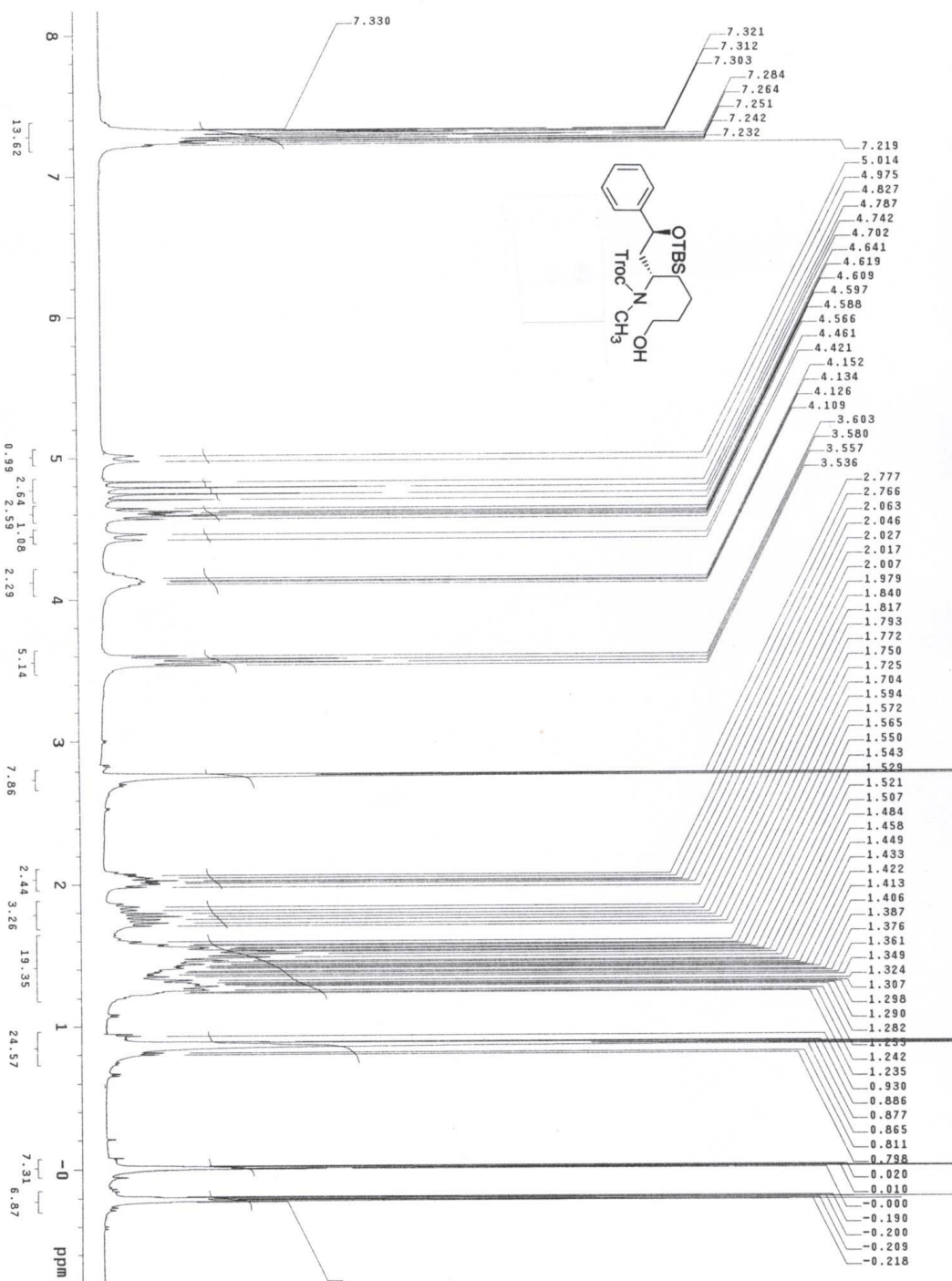




$^{13}\text{C}$  NMR spectrum of **6** ( $\text{CDCl}_3$ , 75 MHz)



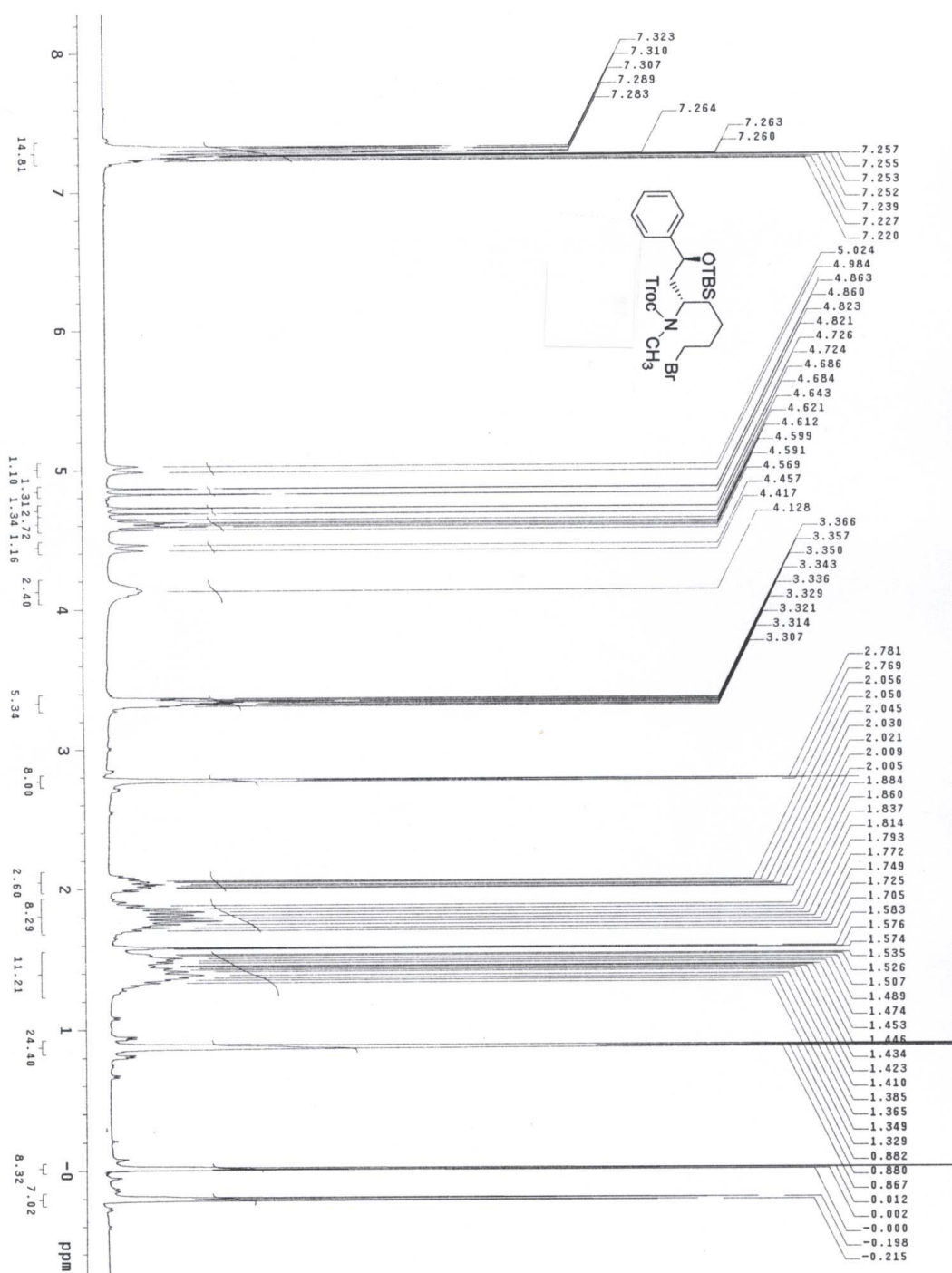
$^1\text{H}$  NMR spectrum of **8** ( $\text{CDCl}_3$ , 300 MHz)





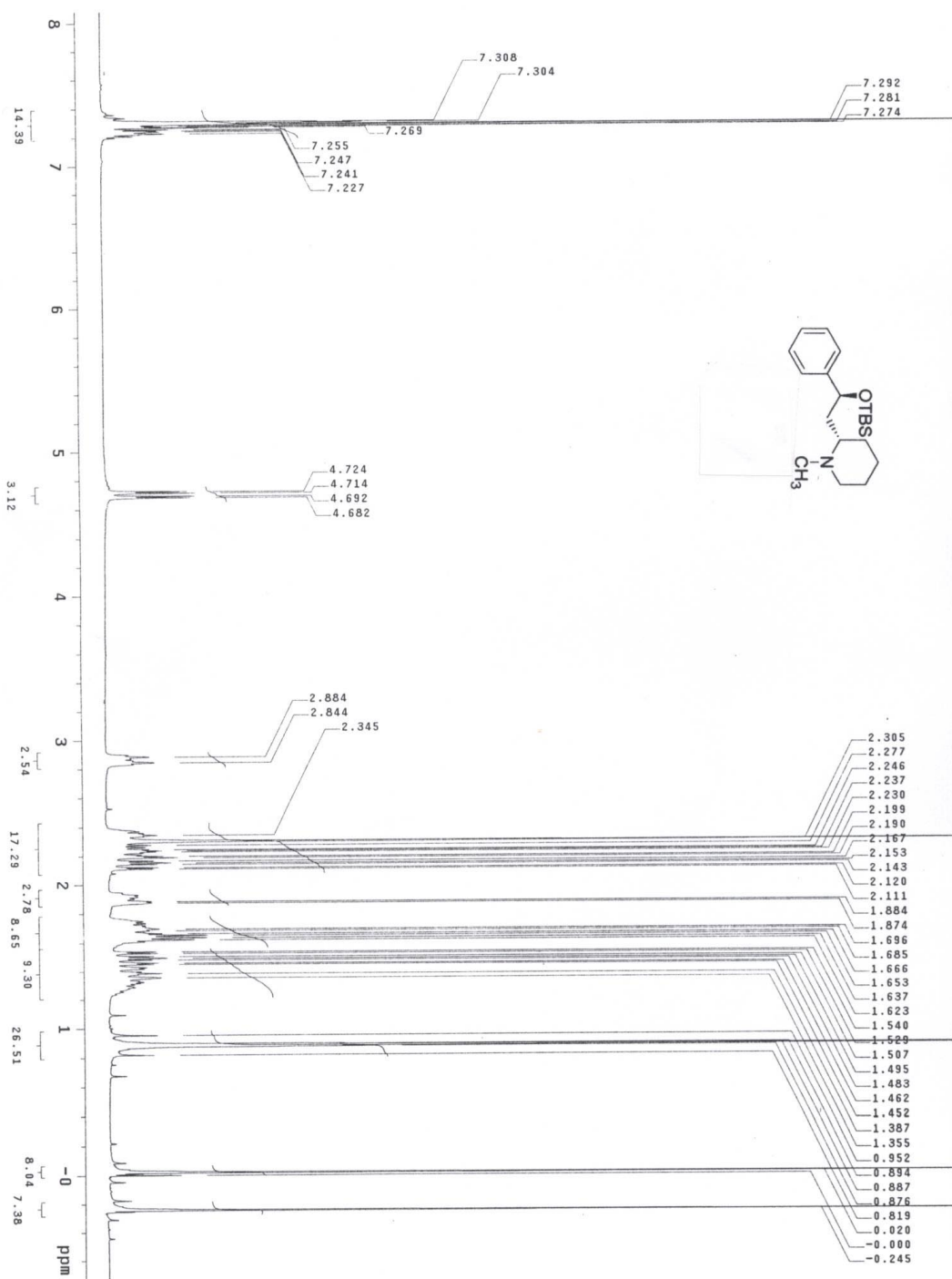


$^1\text{H}$  NMR spectrum of **9** ( $\text{CDCl}_3$ , 300 MHz)





$^1\text{H}$  NMR spectrum of **10** ( $\text{CDCl}_3$ , 300 MHz)



$^{13}\text{C}$  NMR spectrum of **10** ( $\text{CDCl}_3$ , 75 MHz)

