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APPENDICES

APPENDIX A

A. Stock Solution Preparation

Phosphate buffer pH 6.8 stock solution

Preparation of phosphate buffer stock solution 500 mM 1000 mL with K₂HPO₄ (MW 174.18, 42.5521 g) and KH₂PO₄ (MW 136.09, 34.7982 g) in deionizers water. K₂HPO₄ and KH₂PO₄ were dissolved in 900 mL deionizers water and measured pH with pH meter (pH 211 microprocessor pH meters, HANNA Instrument) then adjust pH to 6.8 with 0.1 M HCl and 0.1 M NaOH next adjust volume to 1000 mL.

$$pH = pK_a + \log \frac{[HPO_4^{2-}]}{[H_2PO_4^-]}$$

$$6.8 = 6.82 + \log \frac{[HPO_4^{2-}]}{[H_2PO_4^-]}$$

$$0.2 = \log \frac{[H_2PO_4^-]}{[HPO_4^{2-}]}$$

$$\frac{1.0471}{1} = \frac{[H_2PO_4^-]}{[HPO_4^{2-}]}$$

$$[H_2PO_4^-] = [CA]$$

$$[CA] = (1.0471 / 2.0471) \times 0.5$$

$$[CA] = 0.2557$$

KH₂HPO₄ was used 0.2557 moles, 34.7982 g

$$[HPO_4^{2-}] = [CB]$$

$$[CB] = (1 / 2.0471) \times 0.5$$

$$[CB] = 0.2443$$

K₂HPO₄ was used 0.2443 moles, 42.5521 g

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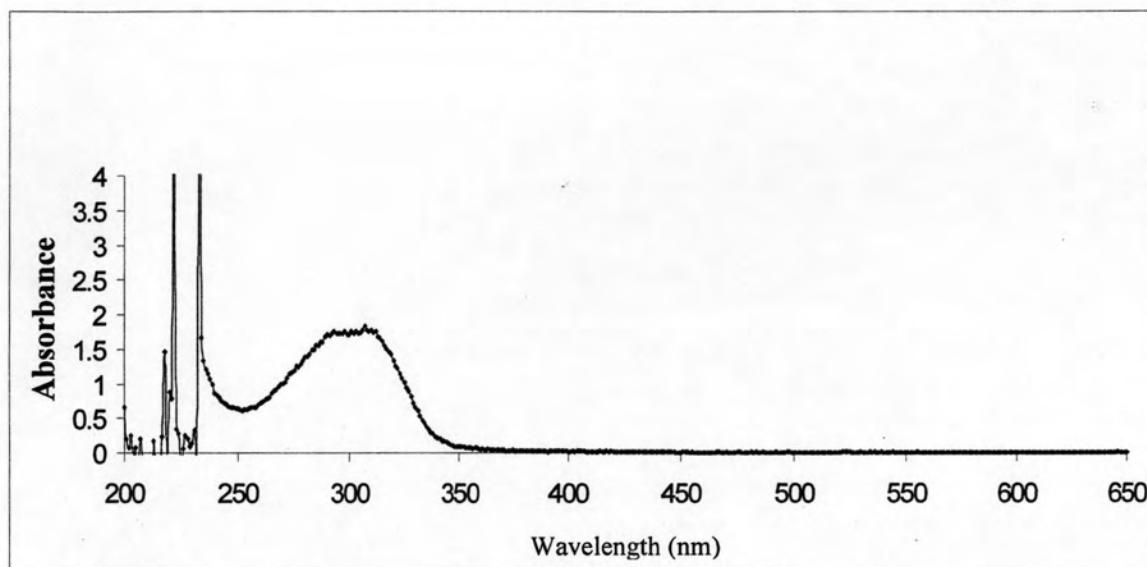
APPENDIX B

Figure B-1 UV spectrum of CH_2Cl_2 crude extract of Chai-nat 1 (CN) strain

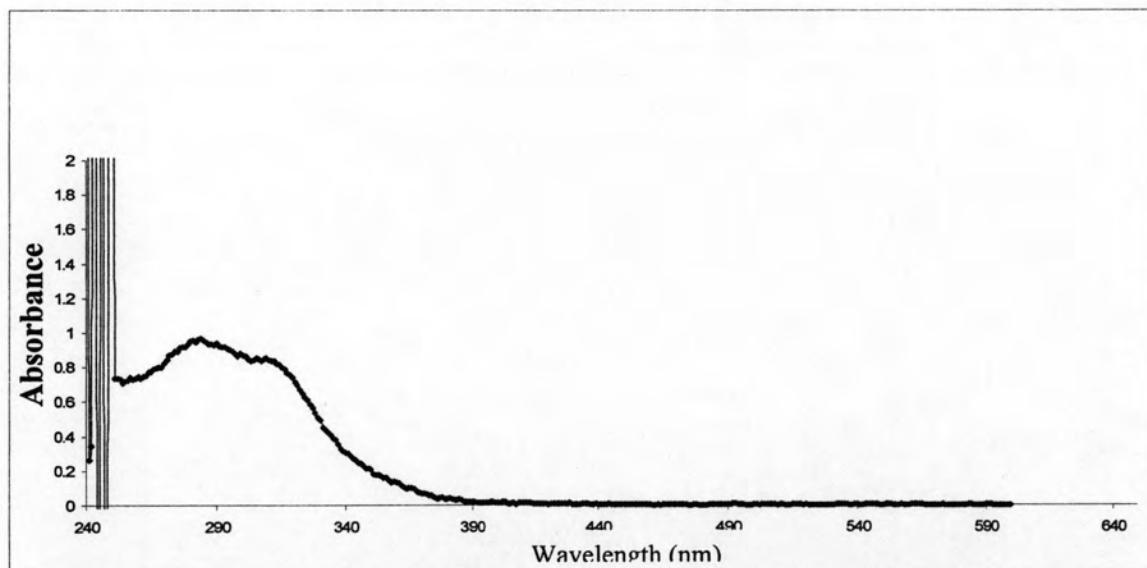


Figure B-2 UV spectrum of EtOAc crude extract of Chai-nat 1 CN strain

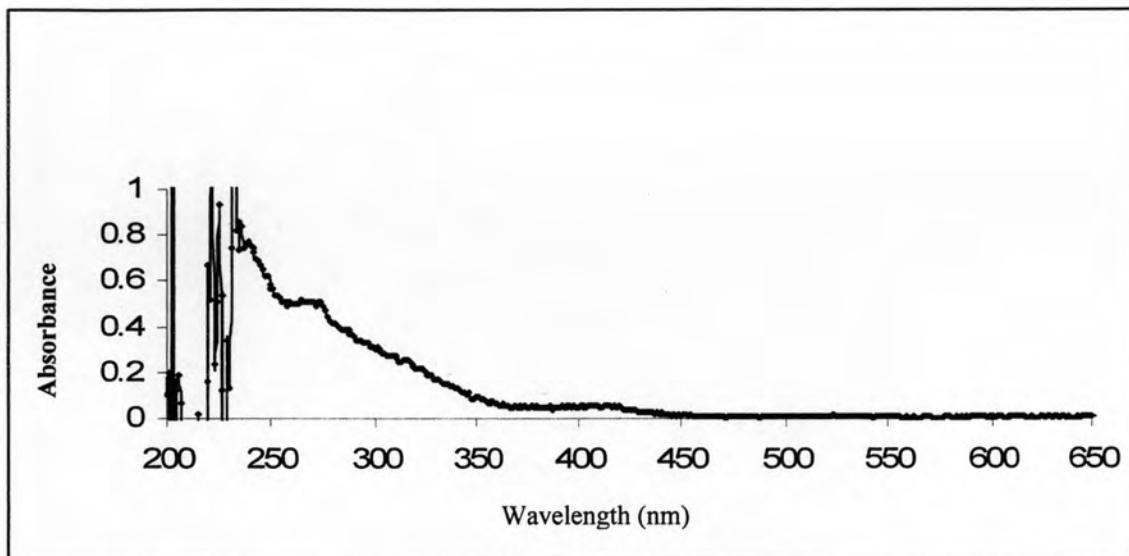


Figure B-3 UV spectrum of CH_2Cl_2 crude extract of Look Daeng Pattani (LD) strain

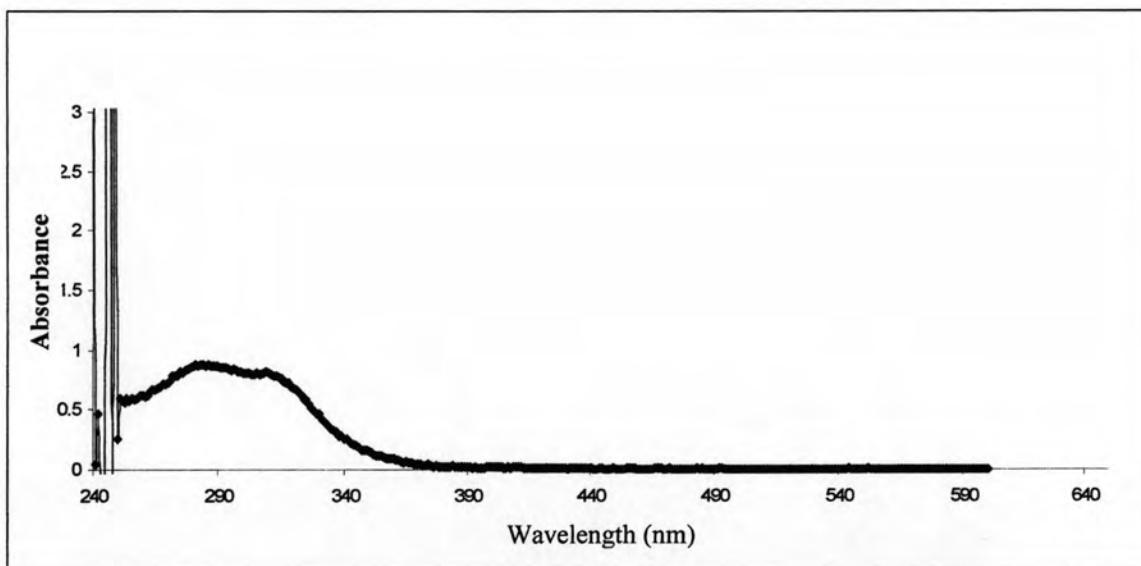


Figure B-4 UV spectrum of EtOAc crude extract of Look Daeng Pattani (LD) strain

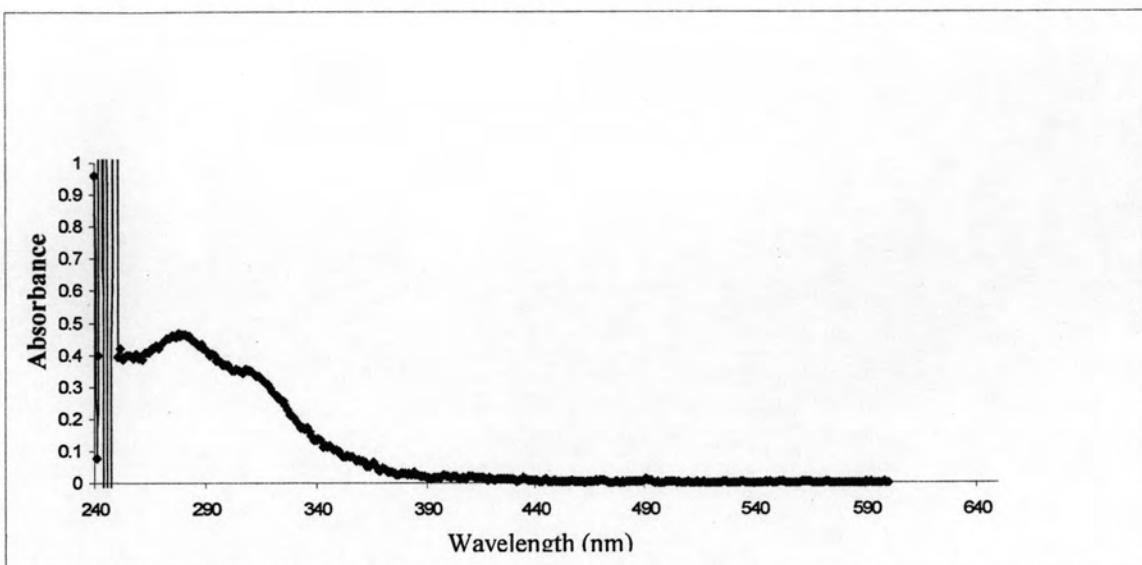


Figure B-5 UV spectrum of EtOAc crude extract of Leb Nok Pattani (LN) strain

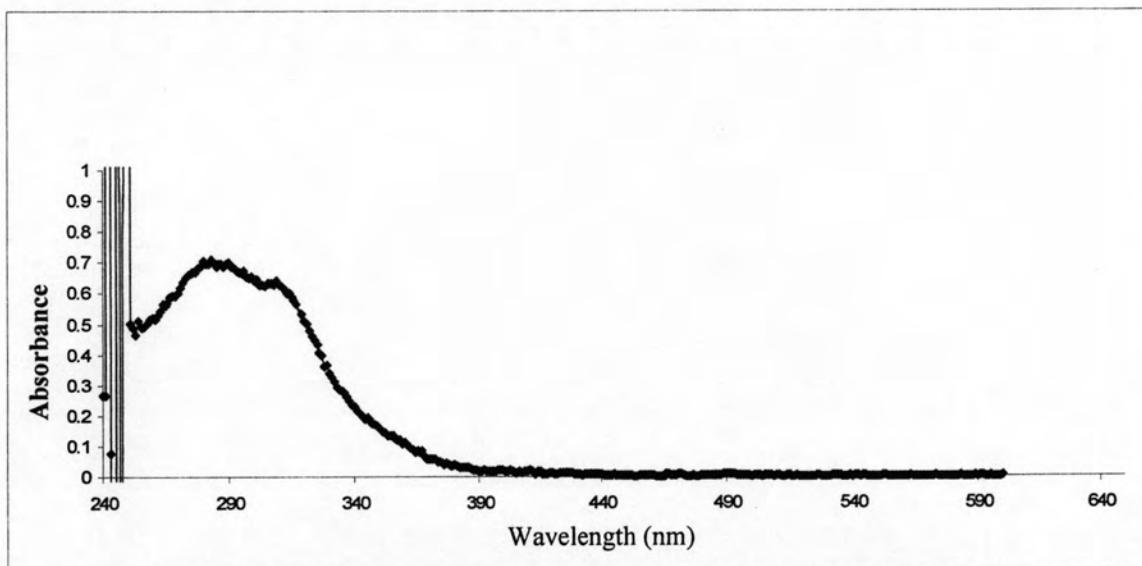


Figure B-6 UV spectrum of EtOAc crude extract of Jasmine (JM) strain

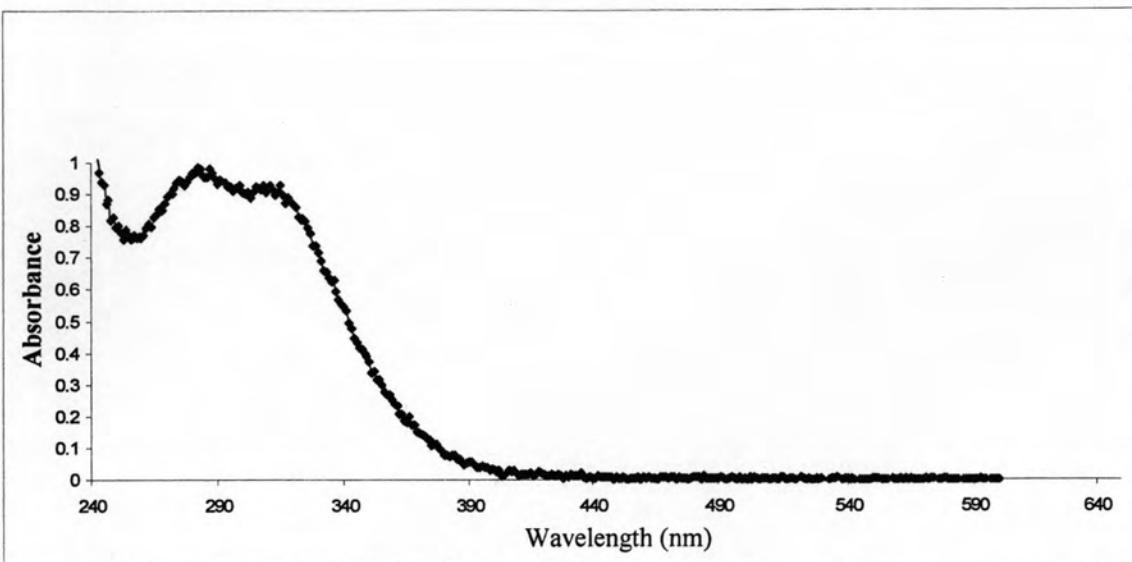


Figure B-7 UV spectrum of MeOH crude extract of Jasmine (JM) strain

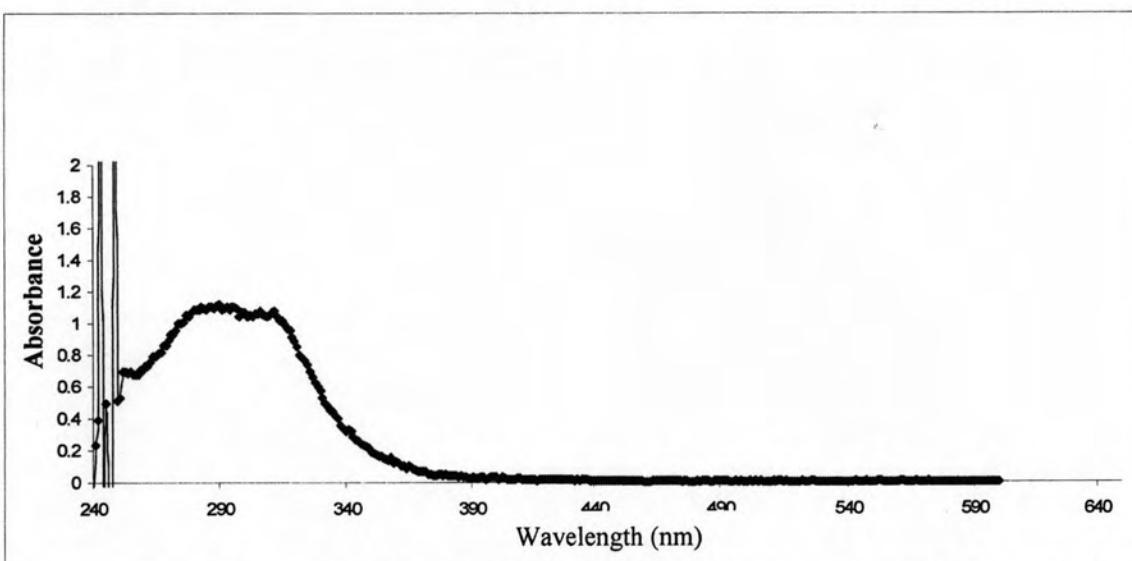


Figure B-8 UV spectrum of EtOAc crude extract of Go Ko 1 (GK) strain

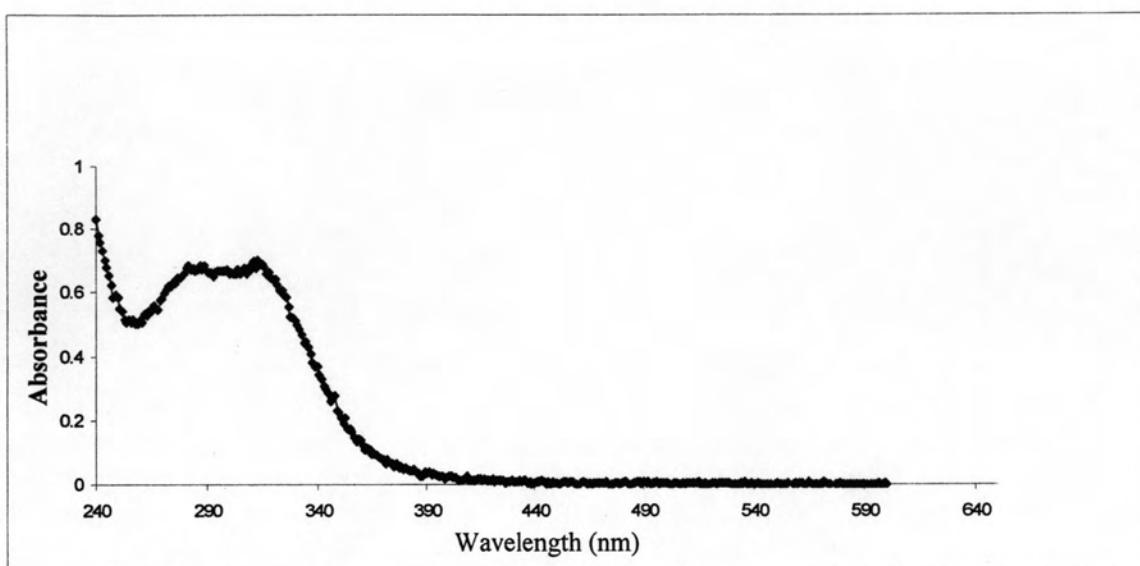


Figure B-9 UV spectrum of MeOH crude extract of Go Ko 1 (GK) strain

APPENDIX C

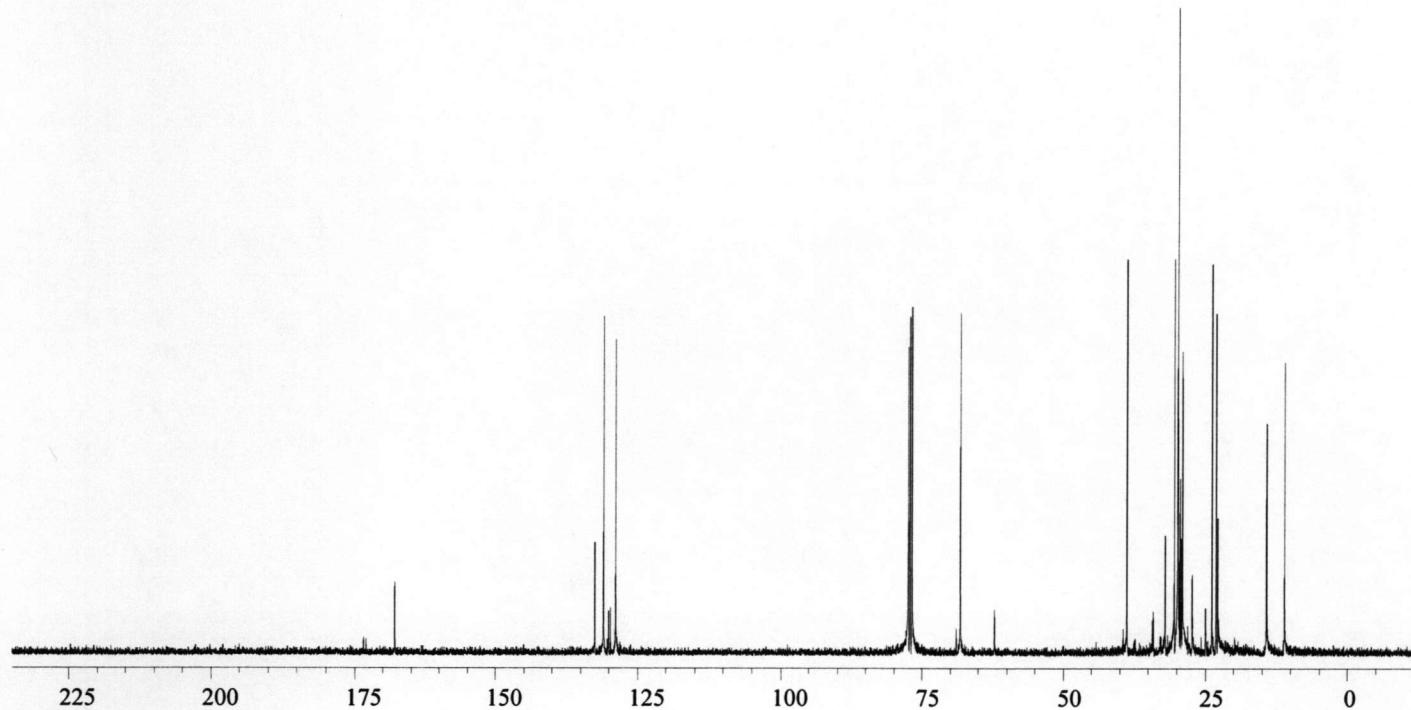


Figure C-1 The ^{13}C NMR spectrum (CDCl_3) of dipentyl phthalate

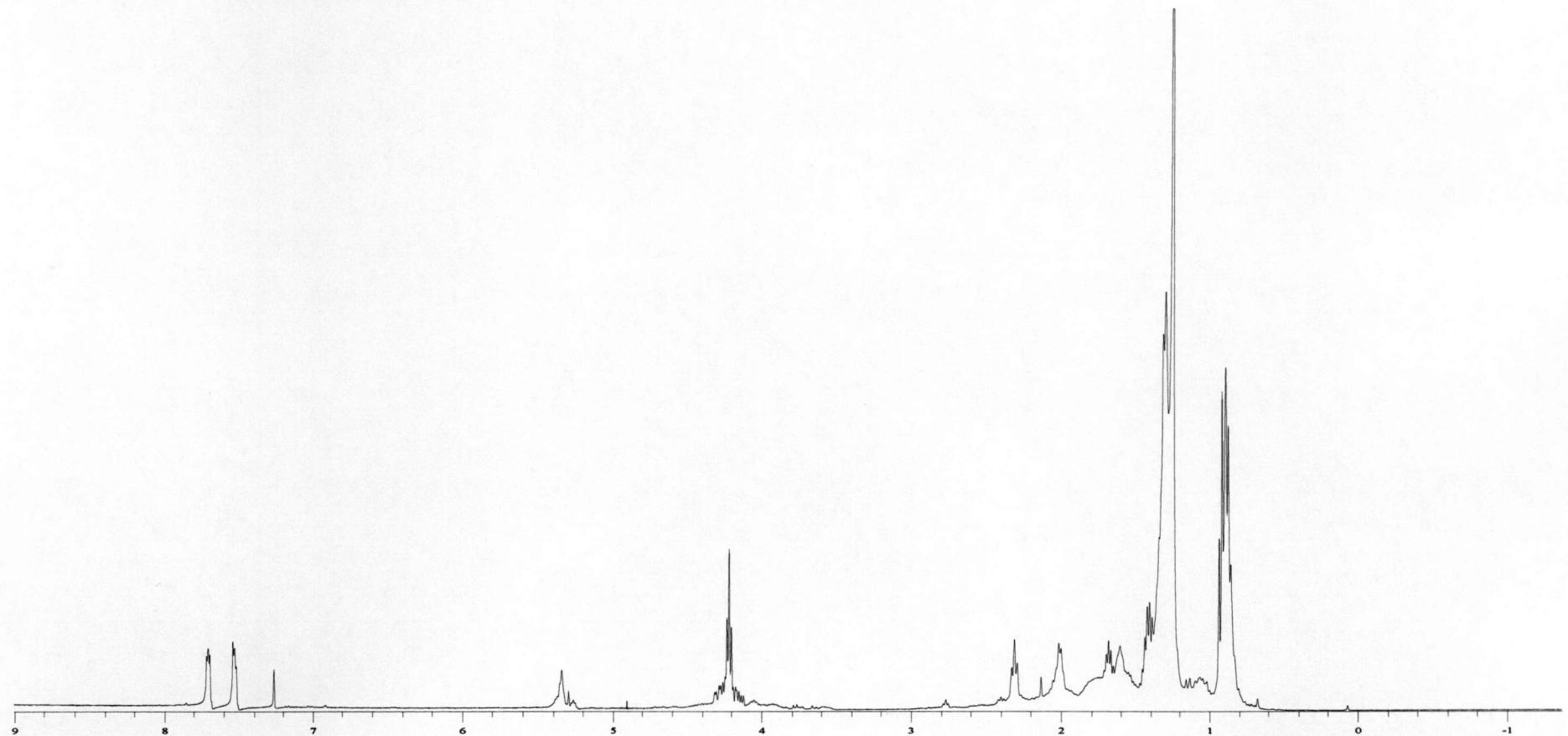


Figure C-2 The ^1H NMR spectrum (CDCl_3) of dipentyl phthalate

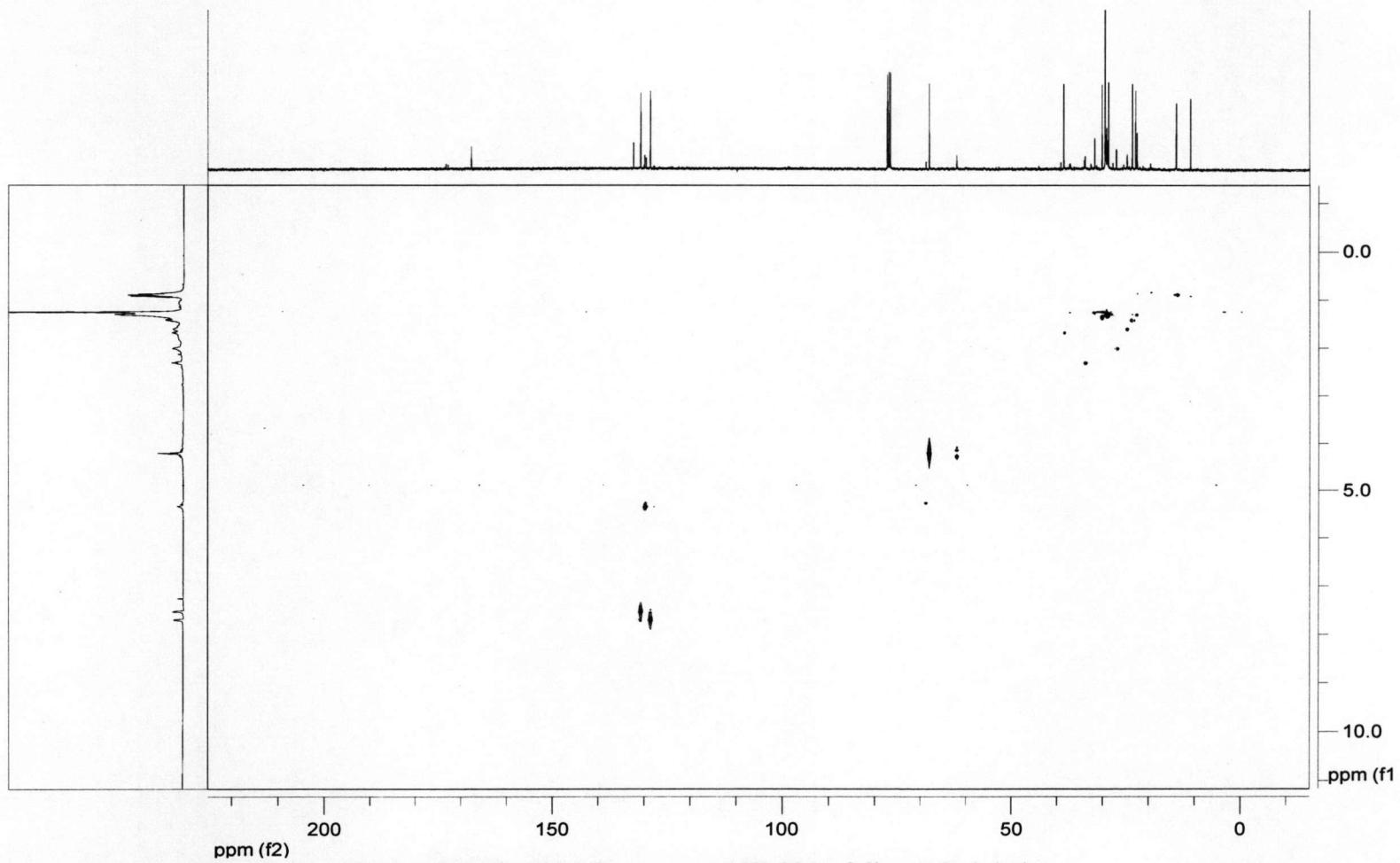


Figure C-3 The HSQC spectrum (CDCl_3) of dipentyl phthalate

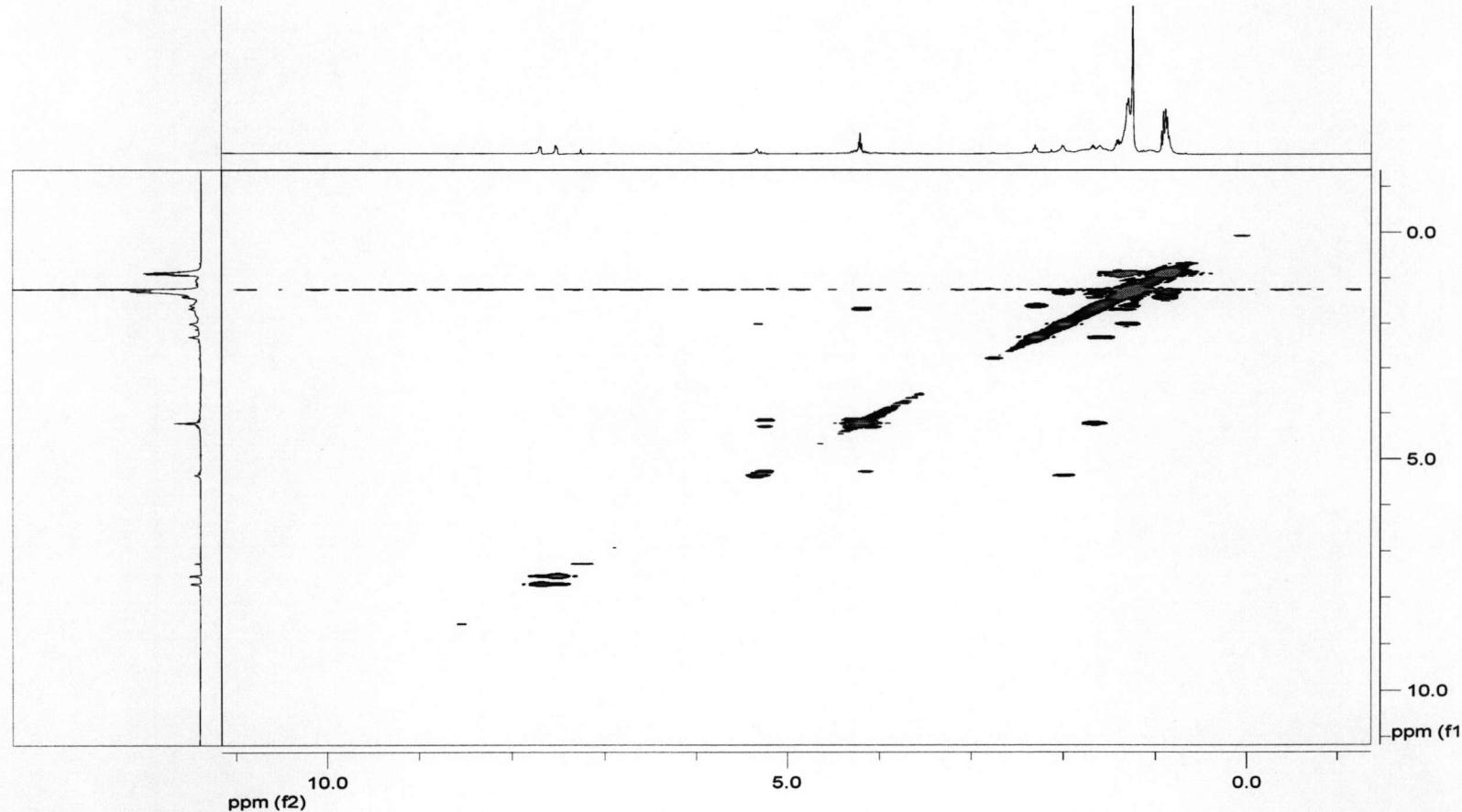


Figure C-4 The COSY spectrum (CDCl_3) of dipentyl phthalate

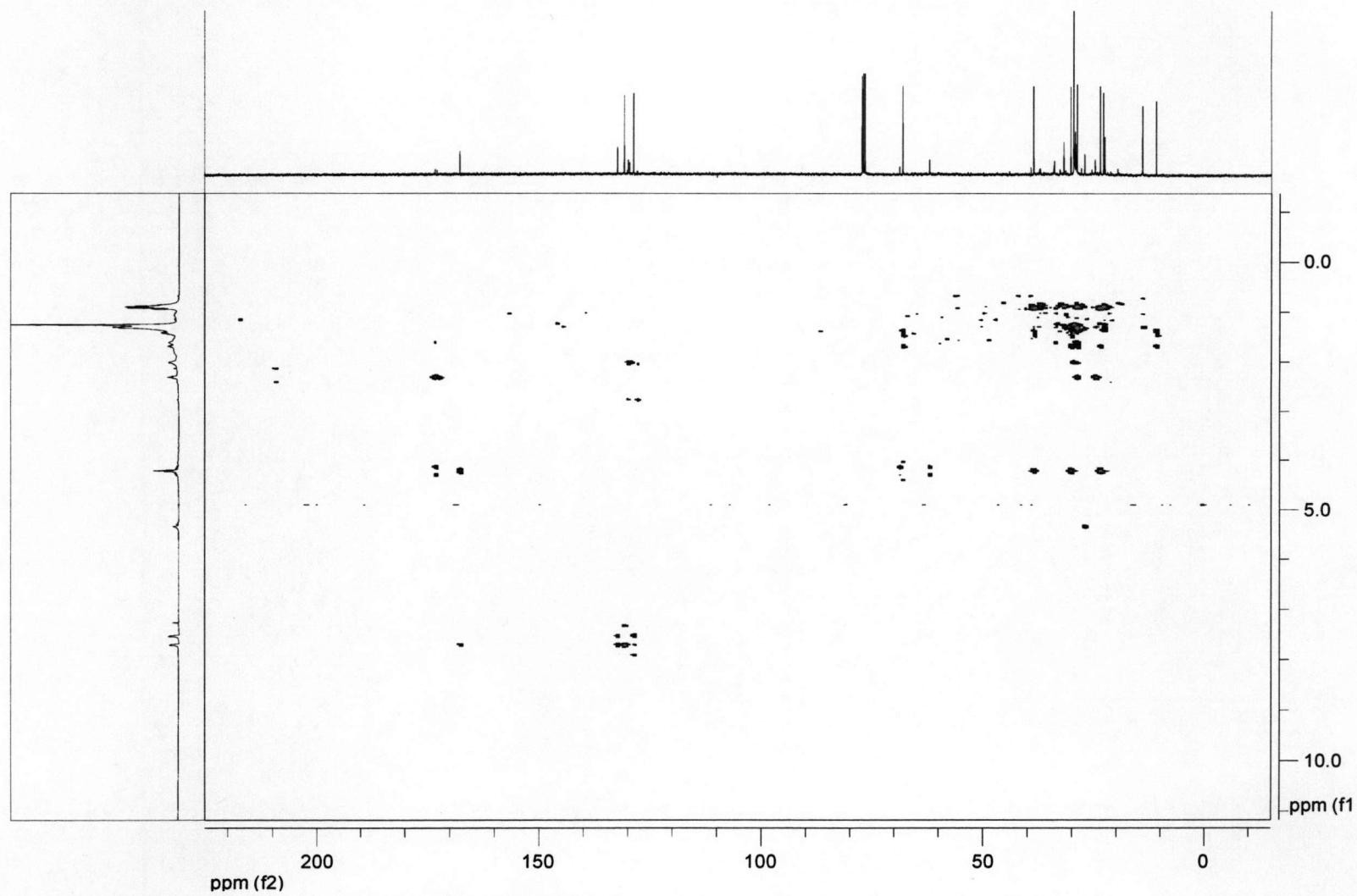


Figure C-5 The HMBC spectrum (CDCl_3) of dipentyl phthalate

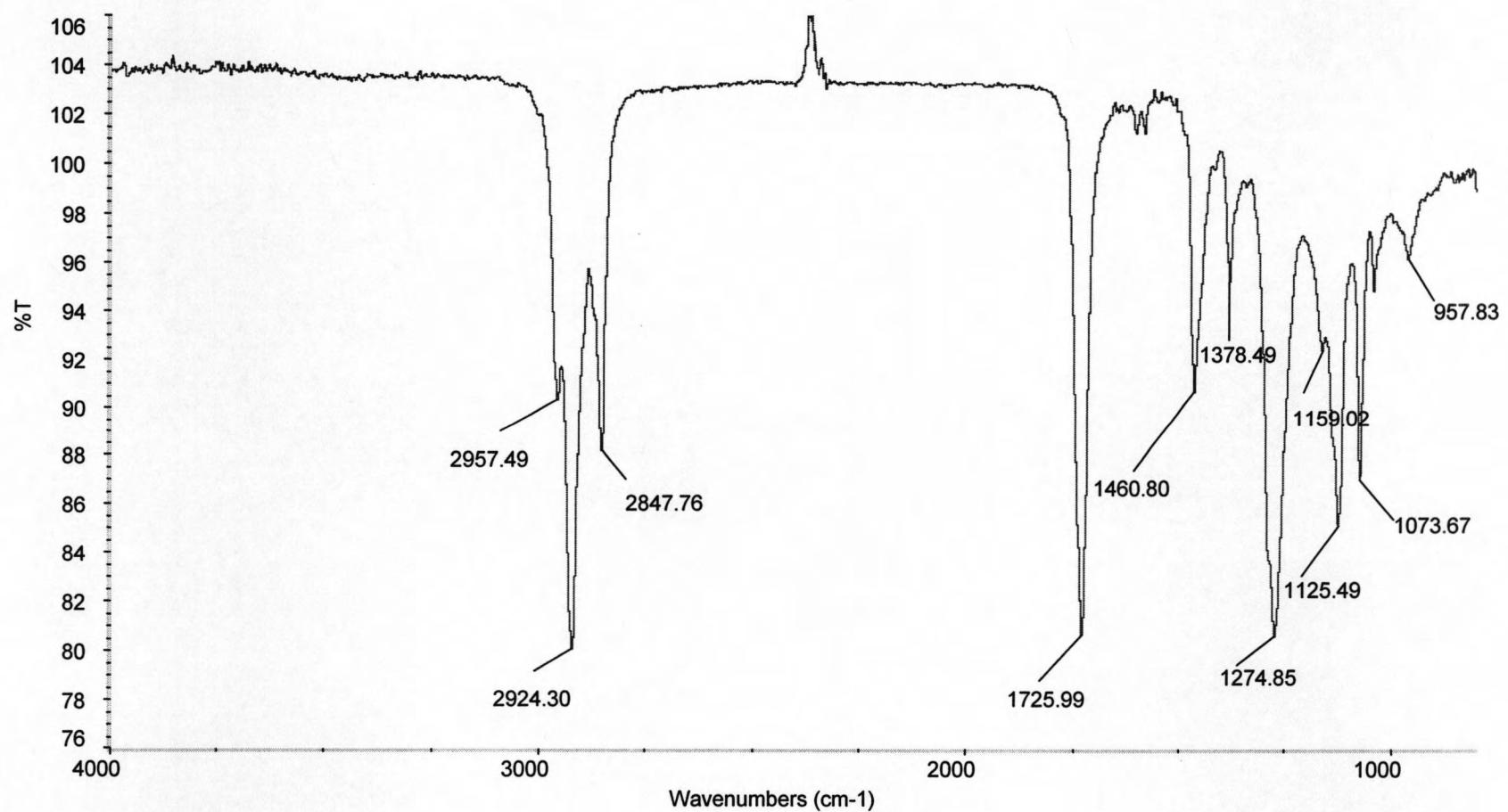


Figure C-6 The IR spectrum of dipentyl phthalate

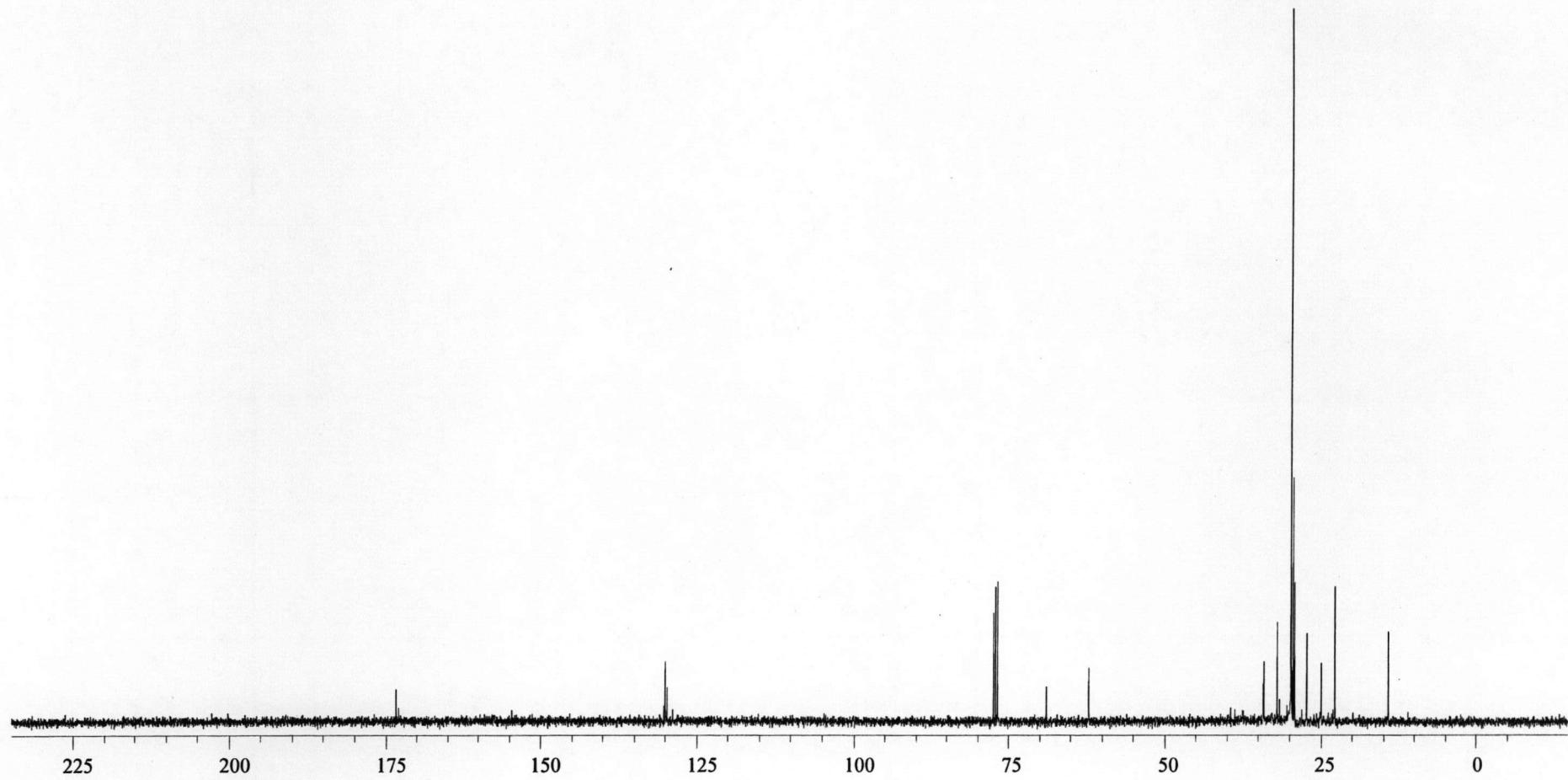


Figure C-7 The ^{13}C NMR spectrum (CDCl_3) of 5-decene

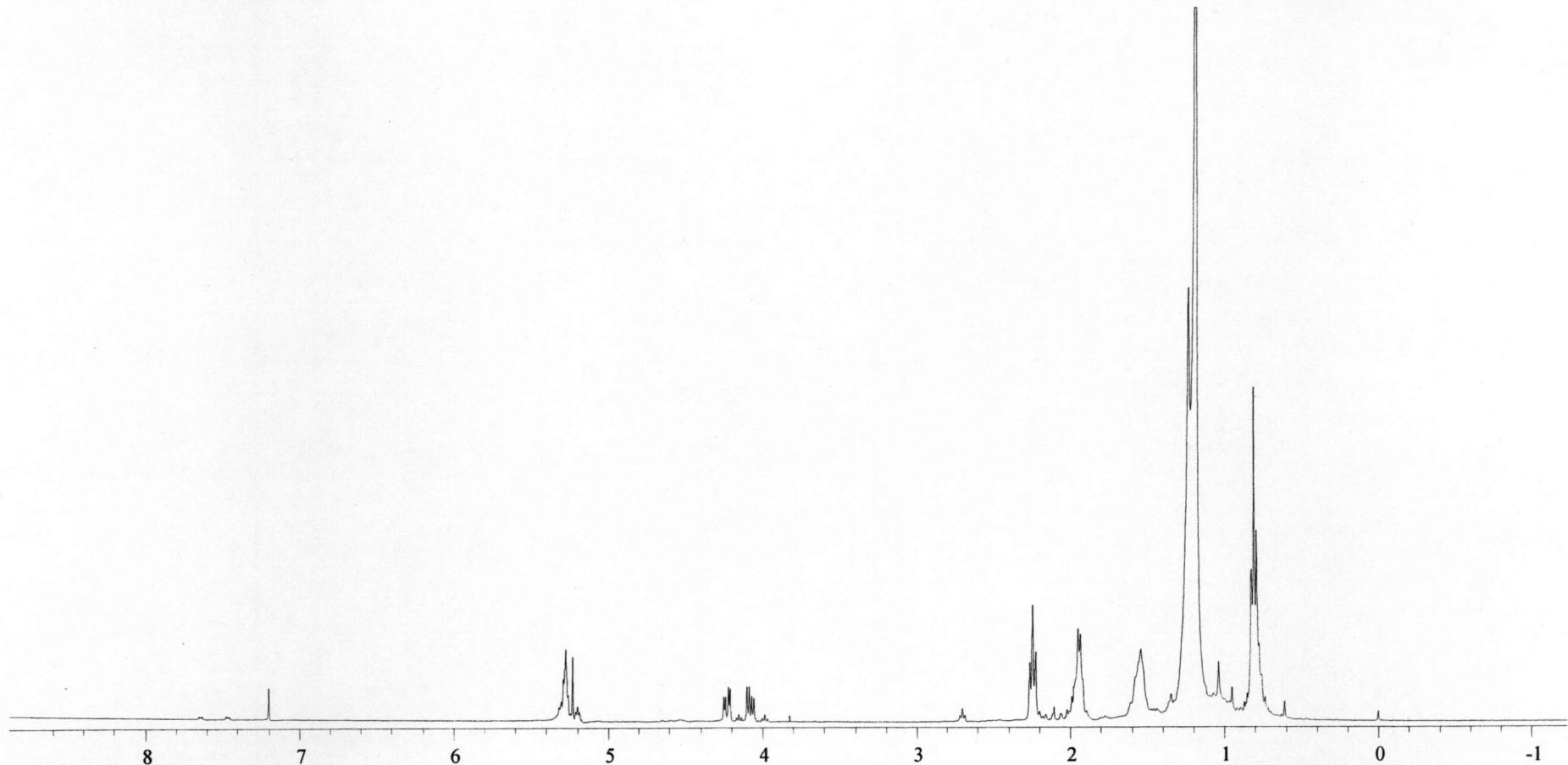


Figure C-8 The ${}^1\text{H}$ NMR spectrum (CDCl_3) of 5-decene

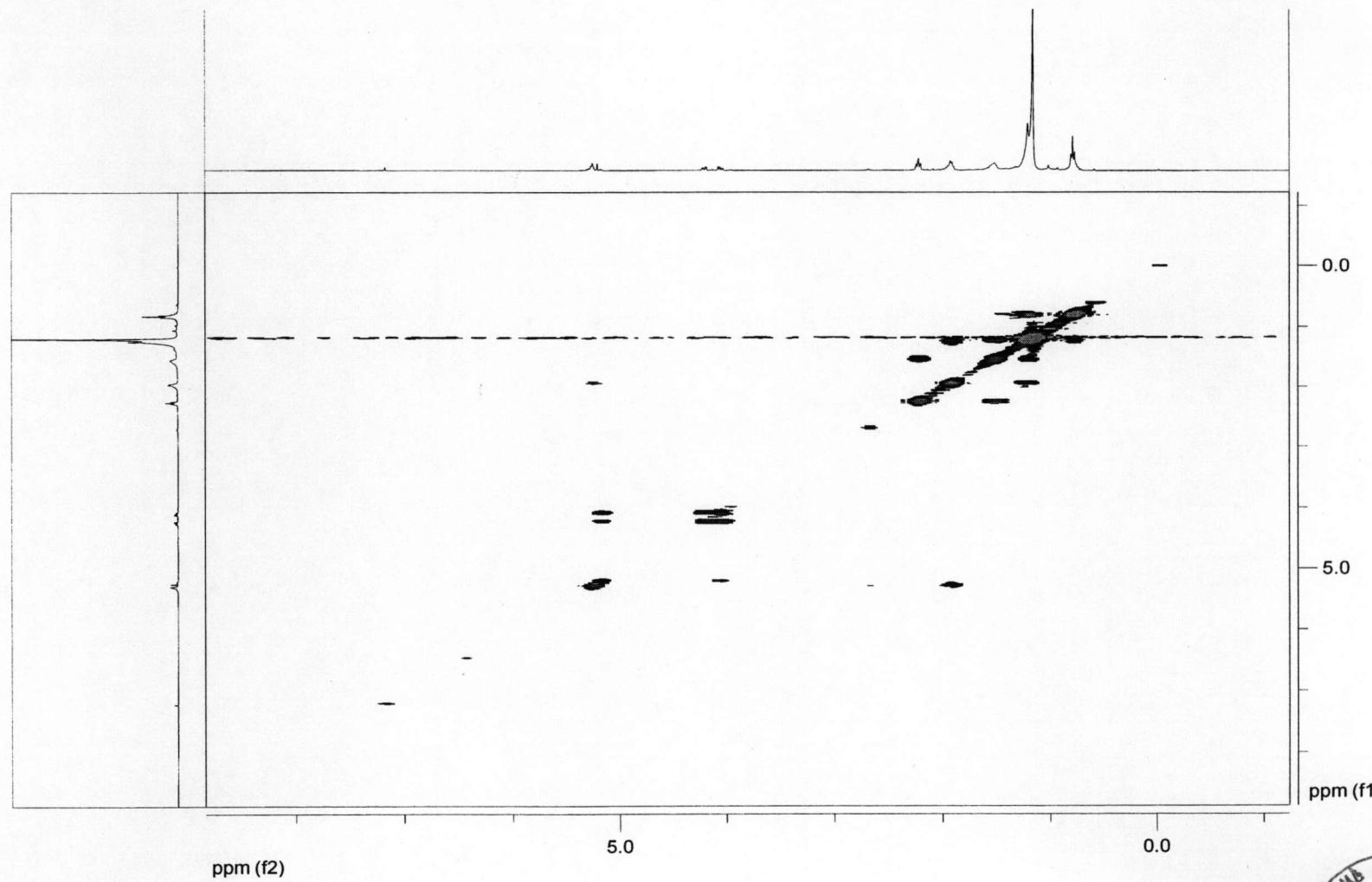


Figure C-9 The COSY spectrum (CDCl_3) of 5-decene



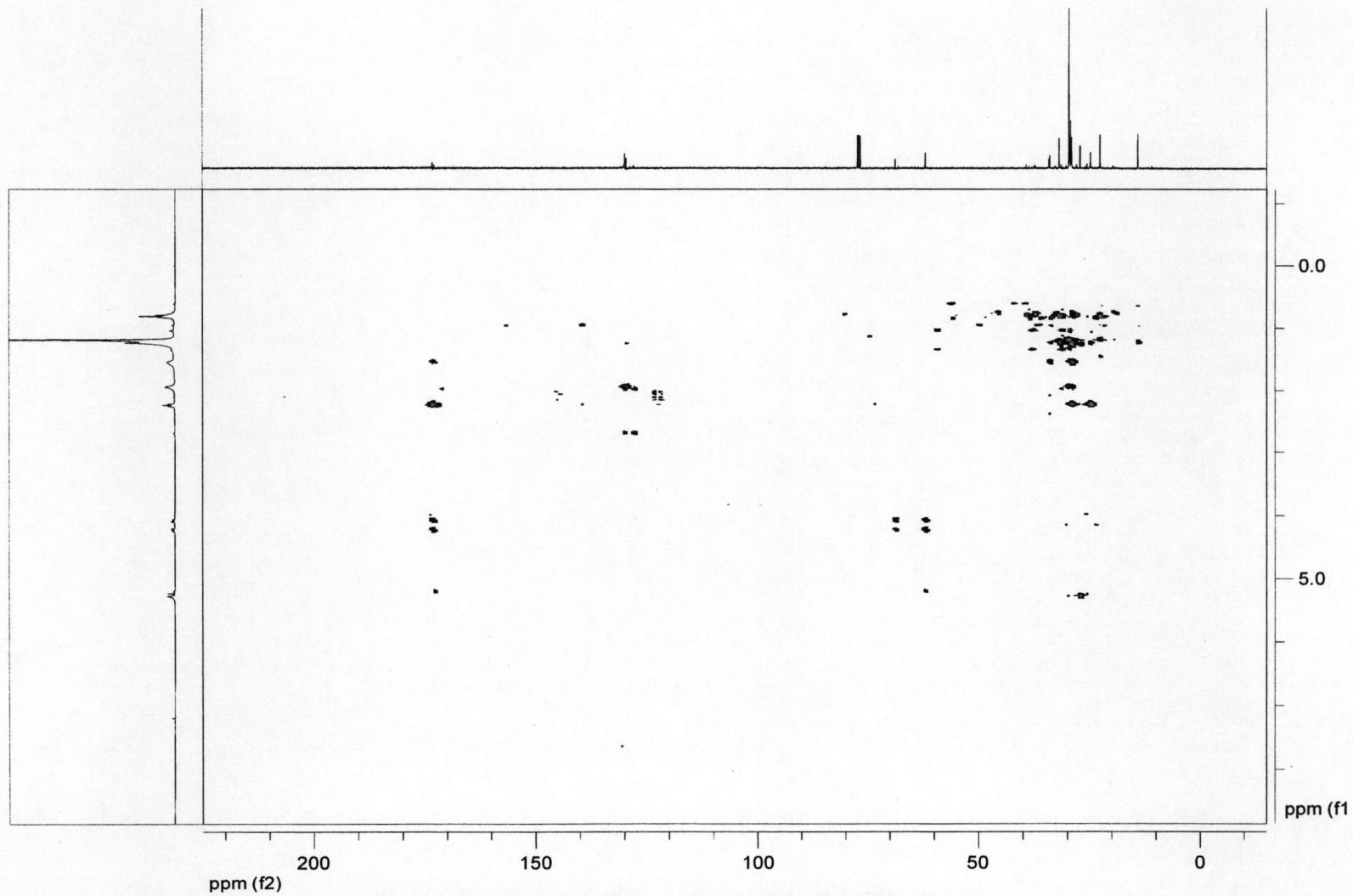


Figure C-10 The HMBC spectrum (CDCl_3) of 5-decene

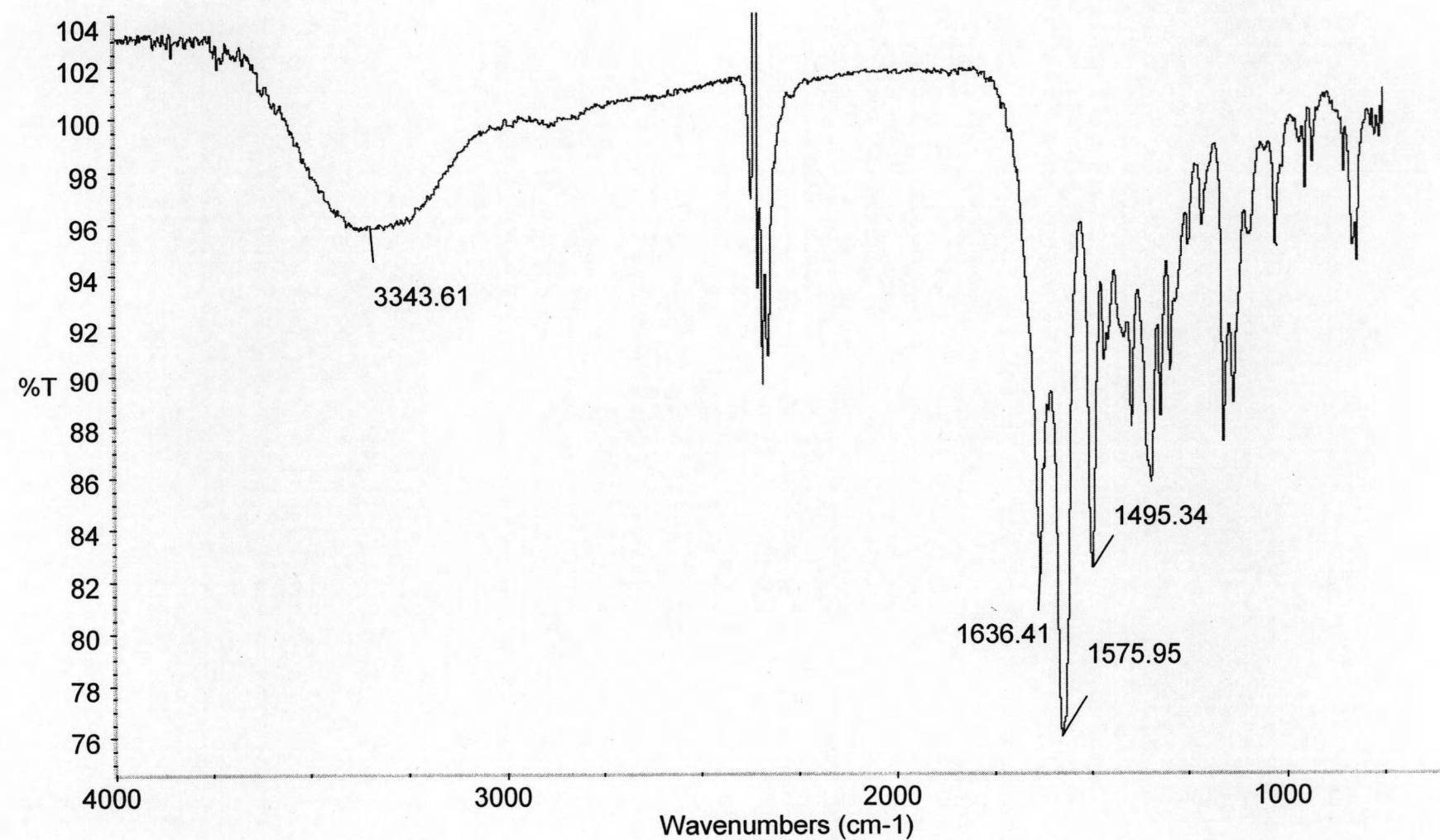


Figure C-11 IR spectrum of 4-hydroxy-3-methoxycinnamaldehyde

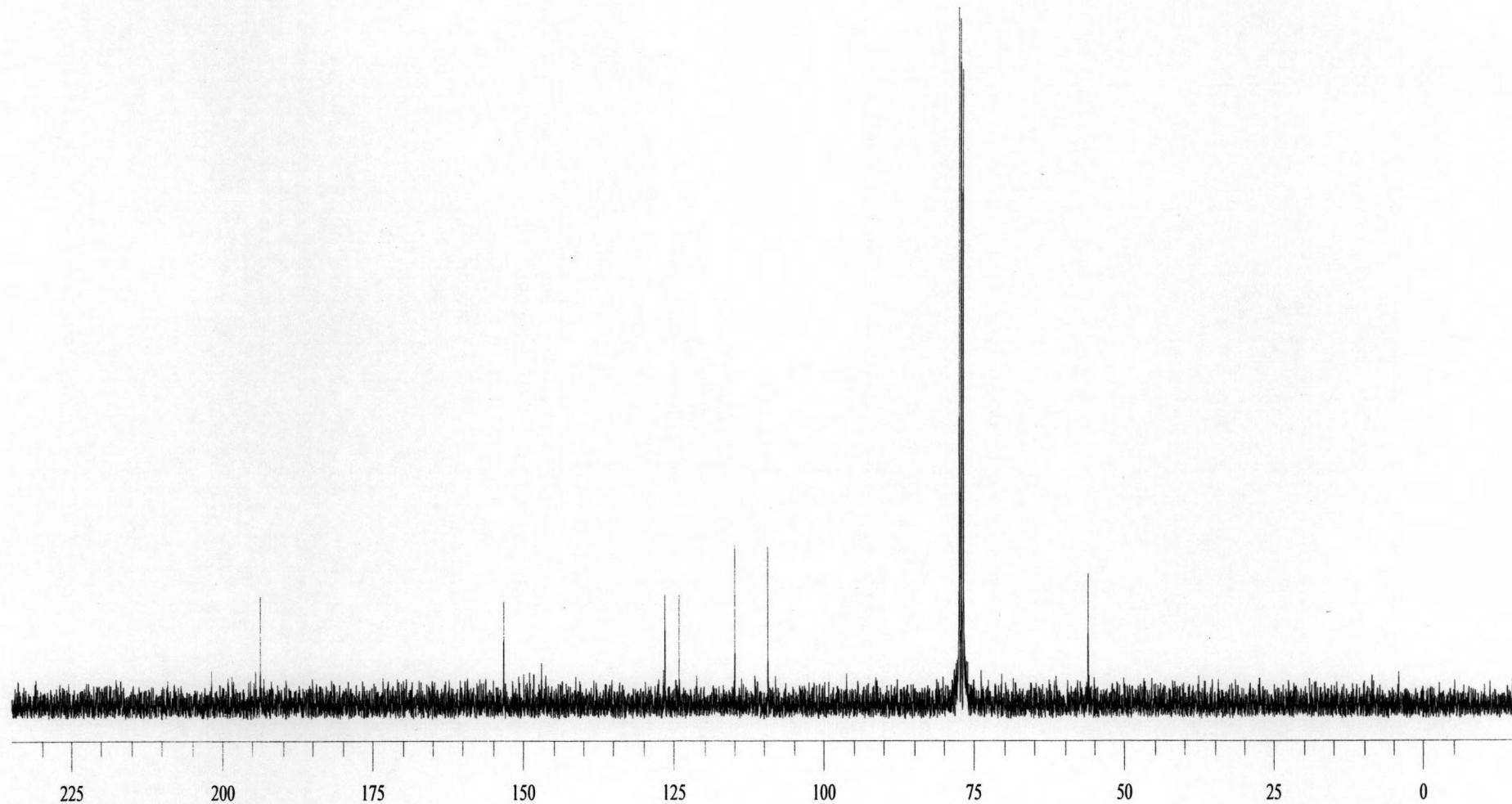


Figure C-12 ^{13}C NMR spectrum (CDCl_3) of 4-hydroxy-3-methoxycinnamaldehyde

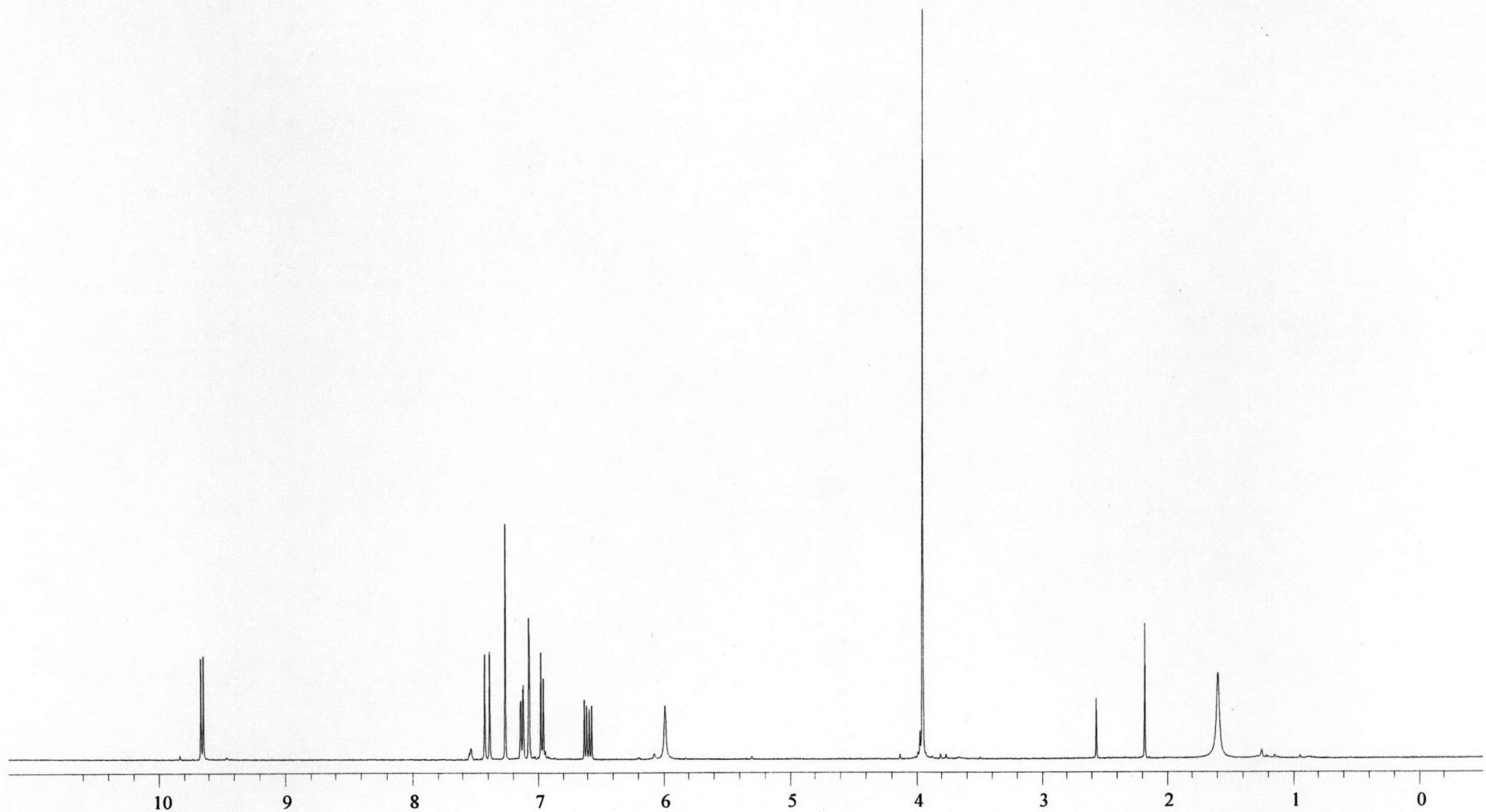


Figure C-13 The ^1H NMR spectrum (CDCl_3) of 4-hydroxy-3-methoxycinnamaldehyde

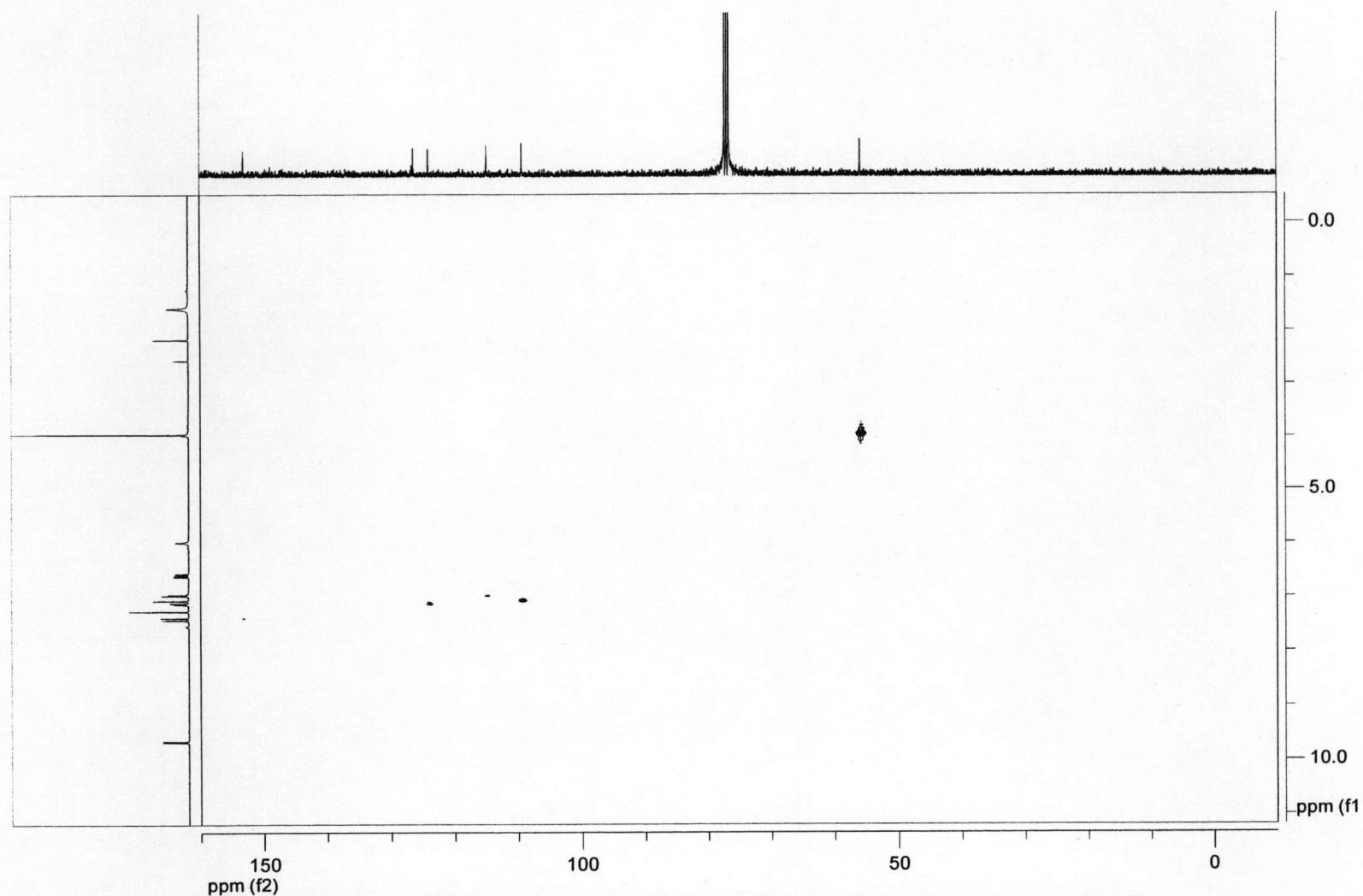


Figure C-14 The HSQC spectrum (CDCl_3) of 4-hydroxy-3-methoxycinnamaldehyde

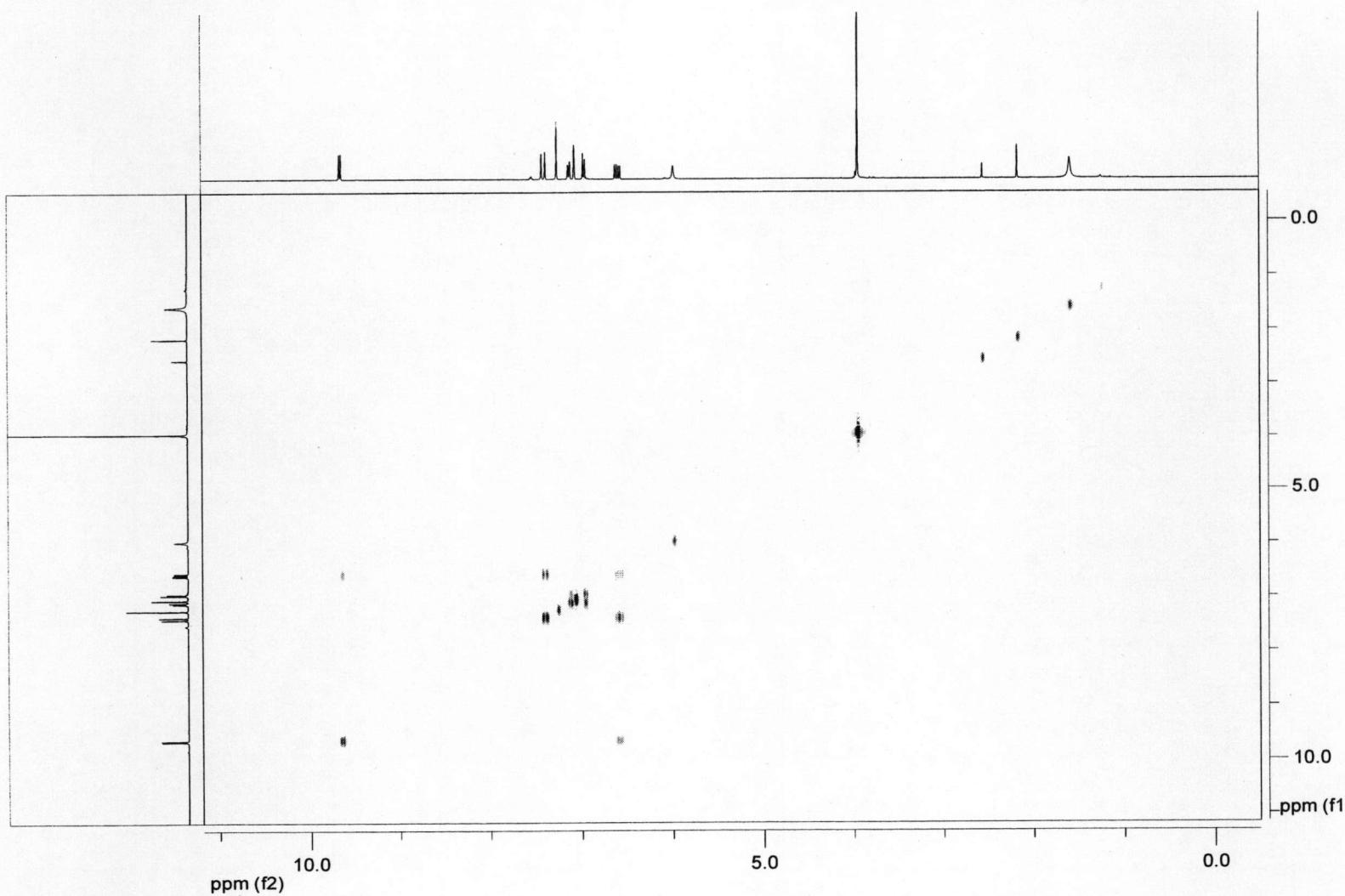


Figure C-15 The COSY spectrum (CDCl_3) of 4-hydroxy-3-methoxycinnamaldehyde

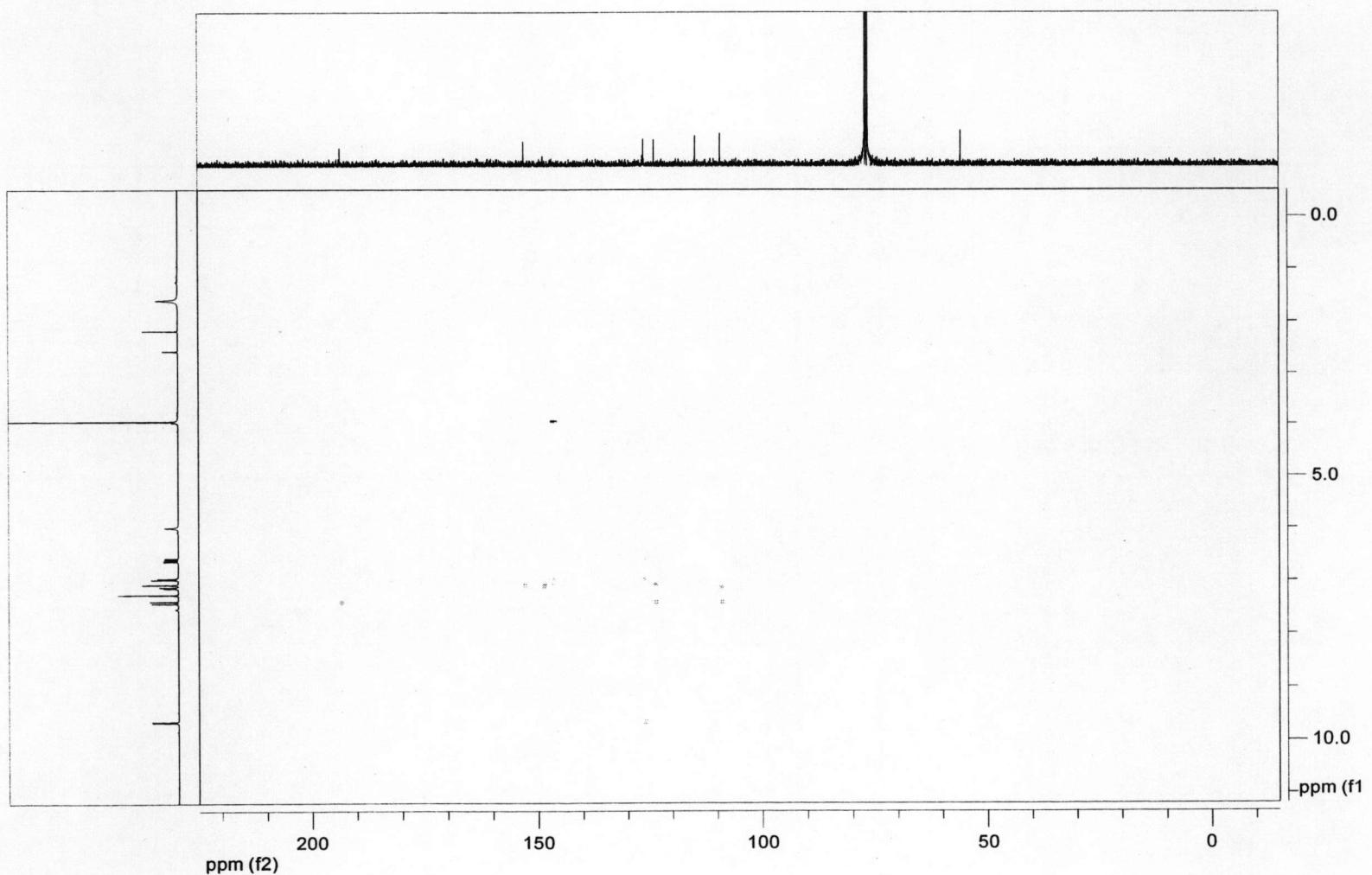


Figure C-16 The HMBC spectrum (CDCl_3) of 4-hydroxy-3-methoxycinnamaldehyde

APPENDIX D

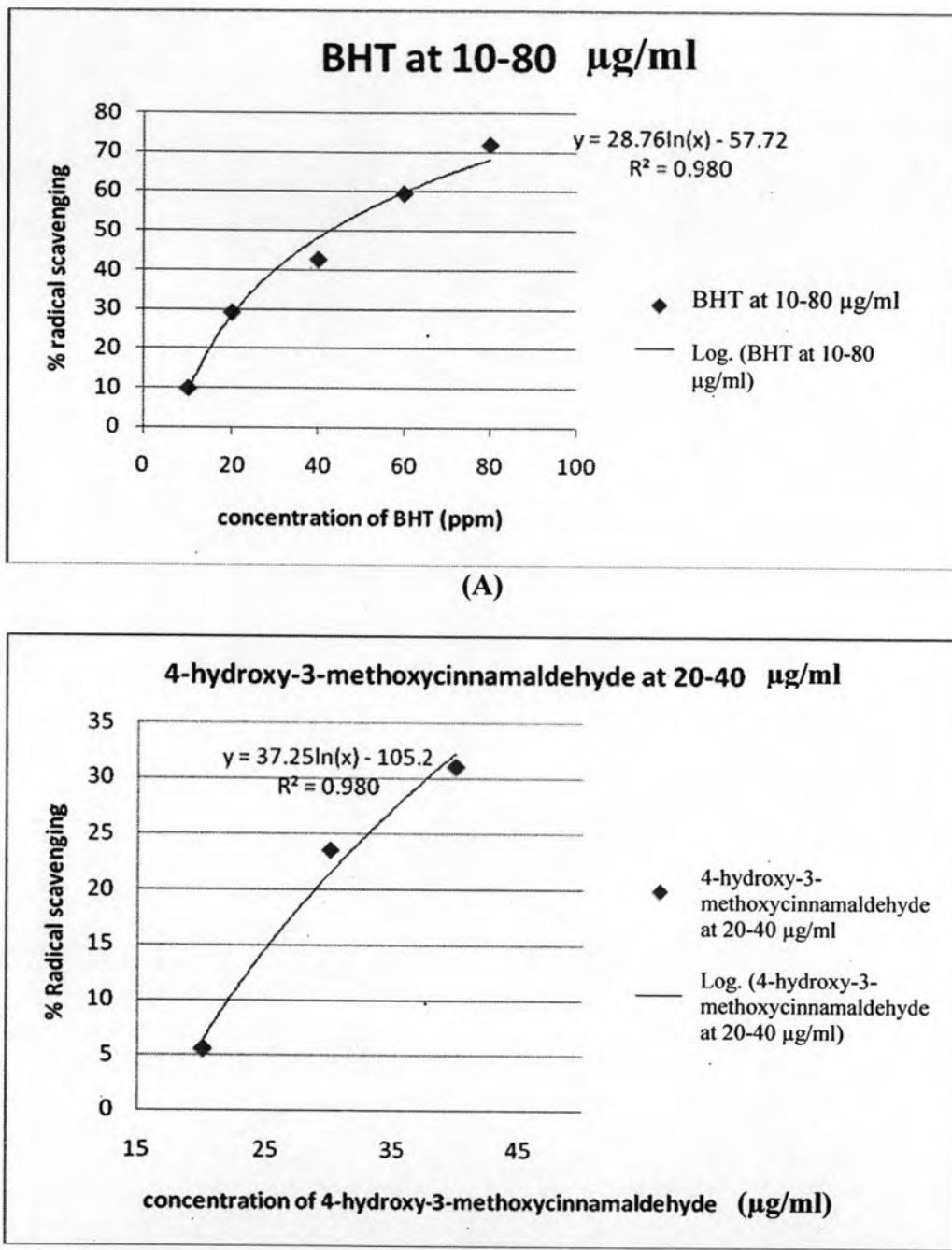


Figure C-1 Free radical scavenging capacity of (A) BHT and (B) 4-hydroxy-3-methoxycinnamaldehyde as determined by the DPPH method. Results are means of three replicates

VITA

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