

REFERENCES

- เต็ม สมิตินันท์. พันธุ์ไม้ป่าเมืองไทย. กรุงเทพมหานคร: โรงพิมพ์อักษรนตรี, 2518.
- _____ . ชื่อพรรณไม้แห่งประเทศไทย. กรุงเทพมหานคร: โรงพิมพ์ฟันนี่พับลิชชิ่ง, 2523.
- Abeygunawardena, C., Kumar, V., Marshall, D.S., Thomson, R.H., and Wickramaratne, D.B.M. Furanonaphthoquinones from two *Latana* species. **Phytochemistry** 30 (1991): 914-945.
- Adirukmi, N.S., Noor Asimah AB, D., and Noor Saleh, Md. **Essential oil content of three species of *Nepenthes* found in Sarawak, Cameron Highlands and Penang Hill, Malaysia.** ASOMP VIII, Malaysia, 1994.
- Akunyili, D.N., and Houghton, P.J. Meroterpenoids and naphthoquinones from *Kigelia pinnata*. **Phytochemistry** 32 (1993): 1015-1018.
- Ali, S., Read, R.W., and Sotheeswaran, S. Benzisochromanquinones and an isofuranonaphthoquinone from *Ventilago vitensis* (Rhamnaceae). **Phytochmistry** 35 (1994): 1029-1032.
- Alves, A.C., Costa, M.A.C., and Paul, M.I. Naphthaquinones from *Diospyros batocana*. **Planta Medica** 47 (1983): 121-124.
- Amatayakul, T., Cannon, J.R., Dampawan, P., Dechatiwongse, T., Giles, R.G.F., Huntrakul, C., Kusamran, K., Mokkasamit, M., Colin, R.L., Reutrakul, V., and White, A.H. Chemistry and crystal structures of some constituents of *Zingiber cassumunar*. **Australian Journal of Chemistry** 32 (1979): 71-88.

- Bhattacharyya, J., and De Carvalho, V.R. Epi-isoshinanolone from *Plumbago scandens*. **Phytochemistry** 25 (1986): 764-765.
- Bieber, L.W., Messana, I., Lins, S.C.N., Da Silva Filho, A.A., Chiappeta, A.A., and De Mello, J. F. Meroterpenoid naphthoquinones from *Cordia corymbosa*. **Phytochemistry** 29 (1990): 1955-1959.
- Binder, R.G., Benson, M.E., and Flath, R.A. Eight 1,4 naphthoquinones from *Juglans*. **Phytochemistry** 28 (1989): 2799-2801.
- Britton, G. **The biochemistry of natural pigments**. Great Britain: Cambridge University Press, 1983.
- Cannon, J.R., Lojanapiwatna, V., Raston, C.L., Sinchai, W., and White, A.H. The quinones of *Nepenthes rafflesiana*. The crystal structure of 2,5-dihydroxy-3,8-dimethoxy-7-methylnaphtho-1,4-quinone (Nepenthone E) and a synthesis of 2,5-dihydroxy-3-methoxy-7-methylnaphtho-1,4-quinone (Nepenthone C). **Australian Journal of Chemistry** 33 (1980): 1073-1093.
- Carey, F.A., and Sunberg, R.J. **Advanced organic chemistry part A: Structure and mechanisms**. 3 rd ed. New York: Plenum Press, 1993.
- _____. **Advanced organic chemistry part B: Reaction and synthesis**. 3 rd ed. New York: Plenum Press, 1993.
- Colegate, S.M., Dorling, P.R., and Huxtable, C.R. Stypandrone: A toxic naphthalene 1,4-quinone from *Stypandra imbricata* and *Dianella revoluta*. **Phytochemistry** 26 (1987): 979-981.
- Comber, M.F., and Sargent, M.V. Synthesis of Larreantin, a cytotoxic naphthoquinonoid sesquilignan from *Larrea tridentata*. **Journal of Chemical Society, Perkin Transactions I** 11 (1991): 2783-2787.

- Cooke, R.G., Dowd, H., and Segal, W. The chemistry of naphthoquinone I. Directive effects in the substitution of naphthoquinone. **Australian Journal of Chemistry** 1953: 38-43.
- Costa, M.A.C., Lopes, M.H., Paul, M.I., Ferreira, M.A., and Alves, A.C. Naphthoquinones and triterpenoids of *Eucheia divinorum*. **Phytochemistry** 15 (1976): 829.
- Dai, J.R., Decosterd, L. A., Gustafson, K.R., Cardellina II, J. H., Gray, G. N., and Boyd, M. R. Novel naphthoquinones from *Conospermum incurvum*. **Journal of Natural Products** 57 (1994): 1151-1156.
- De L. Duarte Weinberg, M., Gottlieb, O.R., and De Oliveira G.G. Naphthoquinones from *Zeyhera tuberculosa*. **Phytochemistry** 15 (1976): 570.
- De Oliveira, A.B., Raslan, D.S., De Oliveira G.G., and Maia, J.G.S. Lignans and naphthoquinones from *Tabebuia incana*. **Phytochemistry** 34 (1993): 1409-1412.
- Ferreira, M.A., Costa, M.A.C., and Alves, A.C. Identification of methylnaphthazarin in *Diospyros* species. **Phytochemistry** 11 (1972): 2352-2353.
- Ferreira, M.A., King, T.J., Ali, S., and Thomson, R. H. Naturally occurring quinones. Part 27. Sesquiterpenoid quinones and related compounds from *Hibiscus elatus*. Crystal structure of hibicone C (Gmelofuran). **Journal of Chemical Society, Perkin Transaction I** 1 (1980): 249-259.
- Ghera, E., and Ben-David, Y. Annulation reactions leading to naphthalene derivatives. New synthesis of natural 1,2- and 1,4-naphthoquinone. **Journal of Organic Chemistry** 50 (1985): 3855-3859.

Gibb, R.D. **Chemotaxonomy of flowering plants.** Great Britain: McGill Queen's University Press, 1979.

Giles, R.G.F., and Roos, G.H.P. Synthesis of substituted 1,4-naphthoquinone by Diels-Alder addition of methoxy cyclohexadiene to substituted 1,4-benzoquinones. **Journal of Chemical Society, Perkin Transactions I** 19 (1976): 2057-2060.

Gunaherath, G.M.K.B., Gunatilaka, A.A.L., Sultanbawa, M. U., and Balasubramanian, S. 1,2(3)-Tetrahydro-3,3'-biplumbagin: A naphthalenone and other constituents from *Plumbago zeylanica*. **Phytochemistry** 22 (1983): 1245-1247.

Gupta, S., Ali, M., and Alam, M.S. Naphthoquinone from *Lawsonia inermis* stem bark. **Phytochemistry** 33 (1993): 723-724.

Heltzel, C.E., Gunalatilaka, A.A.L., Glass, T.E., and Kingston D.G.I. Bioactive furanonaphthoquinones from *Crescentia cujete*. **Journal of Natural Products** 56 (1993): 1500-1505.

Higa, M., Himeno, K., Yogi, S., and Hokoma, K. A new brominated naphthoquinone from *Diospyros maritima* Blume. **Chemical and Pharmaceutical Bulletin** 35 (1987): 4366-4367.

Hirakawa, K., Ogiue, E., Motoyoshiya, J., and Yajima, M. Naphthoquinones from Juglandaceae. **Phytochemistry** 25 (1986): 1494-1495.

Hooker, J.D. **The Flora of British India vol. V.** India: Jayyed Press, 1975.

Iinuma, M., Ohyama, M., and Tanaka, T. Flavonoids in roots of *Sophora prostrata*. **Phytochemistry** 38 (1995): 539-543.

Inoue, K., Ueda, S., Nayeshiro, H., and Inouye, H. Quinones of *Streptocarpus dunnii*. **Phytochemistry** 22 (1983): 737-741.

- Inouye, H., Okuda, T., and Hayashi, T. Quinones and related compounds in higher plant II¹⁾. On the naphthoquinones and related compounds from *Catalpa* wood. **Chemical and Pharmaceutical Bulletin** 23 (1975): 384-391.
- Isaksen, M. Natural pigments. In Sherma, J., and Bernard, F (eds.), **Handbook of Thin-layer chromatography**, pp. 625-662. New York: Marcel Dekker Inc., 1991.
- Itokawa, H., Matsumoto, K., Morita, H., and Takeya, K. Cytotoxic naphthoquinones from *Mansoa alliacea*. **Phytochemistry** 31 (1992): 1061-1062.
- Joshi, K.C., Singh, P., and Pardasani, R.T. Chemical components of the roots of *Tectona grandis* and *Gmelina arborea*. **Planta Medica** 32 (1977): 71-75.
- Kimura, Y., Kozawa, M., Baba, K., and Hata, K. New constituents of roots of *Polygonum cuspidatum*. **Planta Medica** 48 (1983): 164-168.
- Kodama, O., Ishikawa, H., Akatsuka, T., Santisupasri, V., Kato, A., and Hayashi, Y. Isolation and identification of an antifungal naphthopyran derivative from *Rhinacanthus nasutus*. **Journal of Natural Products** 56 (1993): 292-294.
- Komatsu, M., Yokoe, I., Shirataki, Y. Studies on the constituents of *Sophora* species. XIII¹⁾. Constituents of the aerial parts of *Sophora tomentosa* L. (2). **Chemical and Pharmaceutical Bulletin** 26 (1978): 3863-3870.
- Koyama, I., Ogura, T., Tagahara, K., Konoshima, T., and Kozuka, M. Two naphthoquinones from *Rubia cordifolia*. **Phytochemistry** 31 (1992): 2907-2908.
- Kosuge, T., Yokota, M., Sugiyama, K., Mure, T., Yamazawa, H., and Yamamoto, T. Studies on bioactive substances in crude drugs used for arthritic

- diseases in traditional chinese medicine III¹⁾. Isolation and identification of anti-inflammatory and analgesic principles from the whole herb of *Pyrola rotundifolia* L. **Chemical and Pharmaceutical Bulletin** 33 (1985): 5355-5357.
- Kreher, B., Neszmelyi, A., and Wagner, H. Naphthoquinones from *Dionaea muscipula*. **Phytochemistry** 29 (1990): 605-606.
- Kuwahara, S., Awai, N., and Kodama, O. A revised structure for Rhinacanthone. **Journal of Natural Products** 58 (1995): 1455-1458.
- Letcher, R.M., and Shirley, I.M. o-Naphthoquinones from the heartwood of *Azanza garckeana*. **Phytochemistry** 31 (1992): 4171-4172.
- Liebeskind, L.S., Granberg, K.L., and Jing, Z. A strategy for generalization of the regiospecific synthesis of substituted quinones from cyclobutenediones. **Journal of Organic Chemistry** 57 (1992): 4345-4352.
- Mabberley, D.J. **The plant-book: A portable dictionary of the higher plants**. Great Britain: the Bath Press, 1993.
- Mann, J. **Secondary metabolism**. Oxford: Clarendon Press, 1978.
- March, J. **Advanced organic chemistry**. United States of America: McGraw-Hill, Inc., 1977.
- Mc Makin, P.D. **A field guide to the flowering plants of Thailand**. Bangkok: White Lotus Press, 1988.
- Mock, J., Murphy, S.T., Ritchie, E., and Taylor, W.C. Chemical studies of the proteaceae VI. Two naphthoquinones from *Stenocarpus salignus*. **Australian Journal of Chemistry** 26 (1973): 1121-1130.
- Moir, M., and Thomson, R.H. Naphthoquinones in *Lomatia* species. **Phytochemistry** 12 (1973): 1351-1353.

- Morrison, R.T., and Boyd, R.N. **Organic chemistry.** 4 th ed, Singapore: Allyl and Bacon Inc., 1983.
- Nakanishi K., Sasaki, S., Kiang, A.K., Goh, J., Kakisawa, H., Ohashi, M., Goto, M., Watanabe, J., Yokotani, H., Matsumura, C., and Tegashi, M. Phytochemical survey of Malaysian plants: Preliminary chemical and pharmacological screening. **Chemical and Pharmaceutical Bulletin** 13 (1965): 882-890.
- Papageorgiou, V.P. Naphthoquinones from roots of *Macrotomia cephalotes* DC. **Planta Medica** 37 (1979) 259-263.
- _____. Naturally occurring isohexenylnaphthazarin pigments. A new class of drugs. **Planta Medica** 38 (1980): 194-203.
- Perry, L.M. **Medicinal plants of East and Southeast Asia.** United States of America: The Massachusetts Institute of Technology press, 1980.
- Pluim, H., and Wynberg, H. Catalytic asymmetric induction in oxidation reaction. Synthesis of optically active epoxynaphthoquinone. **Journal of Organic Chemistry** 45 (1980): 2498-2502.
- Sankaram, A.V.B., Narayana Reddy, V.V., and Marthandamurthi, M. ¹³C NMR spectra of some naturally occurring binaphthoquinones and related compounds. **Phytochemistry** 25 (1986); 2867-2871.
- Silverstein, R.M., Bassler, G.C., and Morril, T.C. **Spectrometric identification of organic compounds.** 5th ed. Singapore: John Wiley and son, Inc., 1991.
- Smith, M.B. **Organic synthesis.** Singapore: McGraw-Hill, Inc., 1994.
- Tatum, J.H., Baker, R.A., and Berry, R.E. Naphthoquinones and derivatives from *Fusarium*. **Phytochemistry** 26 (1987) 795-798.

- Tezuka, M., Takahashi, C., Kuroyanagi, M., Satake, M., Yoshihira., K., and Natori, S. New naphthoquinones from *Diospyros*. **Phytochemistry** 12 (1973): 175-183.
- Thomson, R.H. **Naturally occurring quinones**. 2 nd ed. London: Academic Press, 1971.
- Van der Vijver, L.M., and Gerritsma, K.W. Naphthoquinones of *Euclea* and *Diospyros* species. **Phytochemistry** 13 (1974): 2322-2323.
- Wan, A.S., Axel, R.T., Ramsey, R.B., and Nicholas, H.J. Sterols and triterpenes of the pitcher plants. **Phytochemistry** 11 (1972): 456-461.
- Wu, T.S., Tien, H.J., Yeh, M.Y., and Lee, K.H. Isolation and cytotoxicity of rhinacanthin-A and -B, two naphthopuinones from *Rhinacanthus nasutus*. **Phytochemistry** 27 (1988): 3787-3788.
- Yue. J.M., Lin, Z.W., Wang D.Z., Feng, Y.Z., and Sun H.D. Plumbasides A-C three naphthoquinone derivatives from *Ceratostigma minus*. **Phytochemistry** 35 (1994): 1023-1025.
- Zakaria, M. B., Jeffreys, J. A. D., Waterman, P. G., and Zhong, S. M. Naphthoquinones and triterpenes from some Asian *Diospyros* species. **Phytochemistry** 23 (1984): 1481-1484.
- Zhong, S. M., Waterman, P. G., and Jeffreys, J. A. D. Naphthoquinones and Triterpenes from African *Diospyros* species. **Phytochemistry** 23 (1984): 1067-1072.

APPENDIX

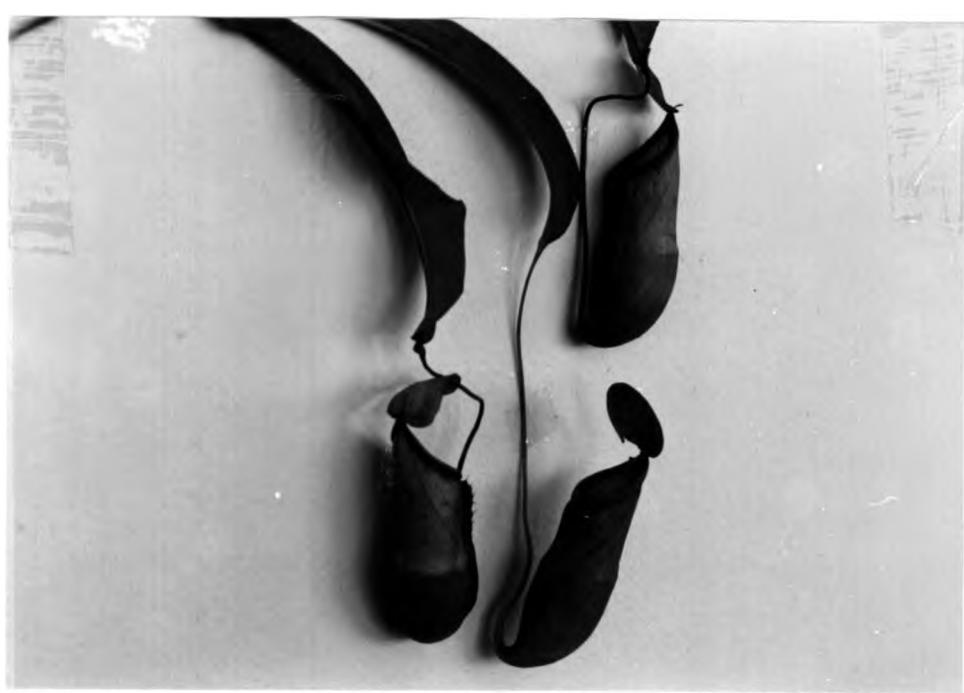


Figure 11 *Nepenthes thorelii* Lec.

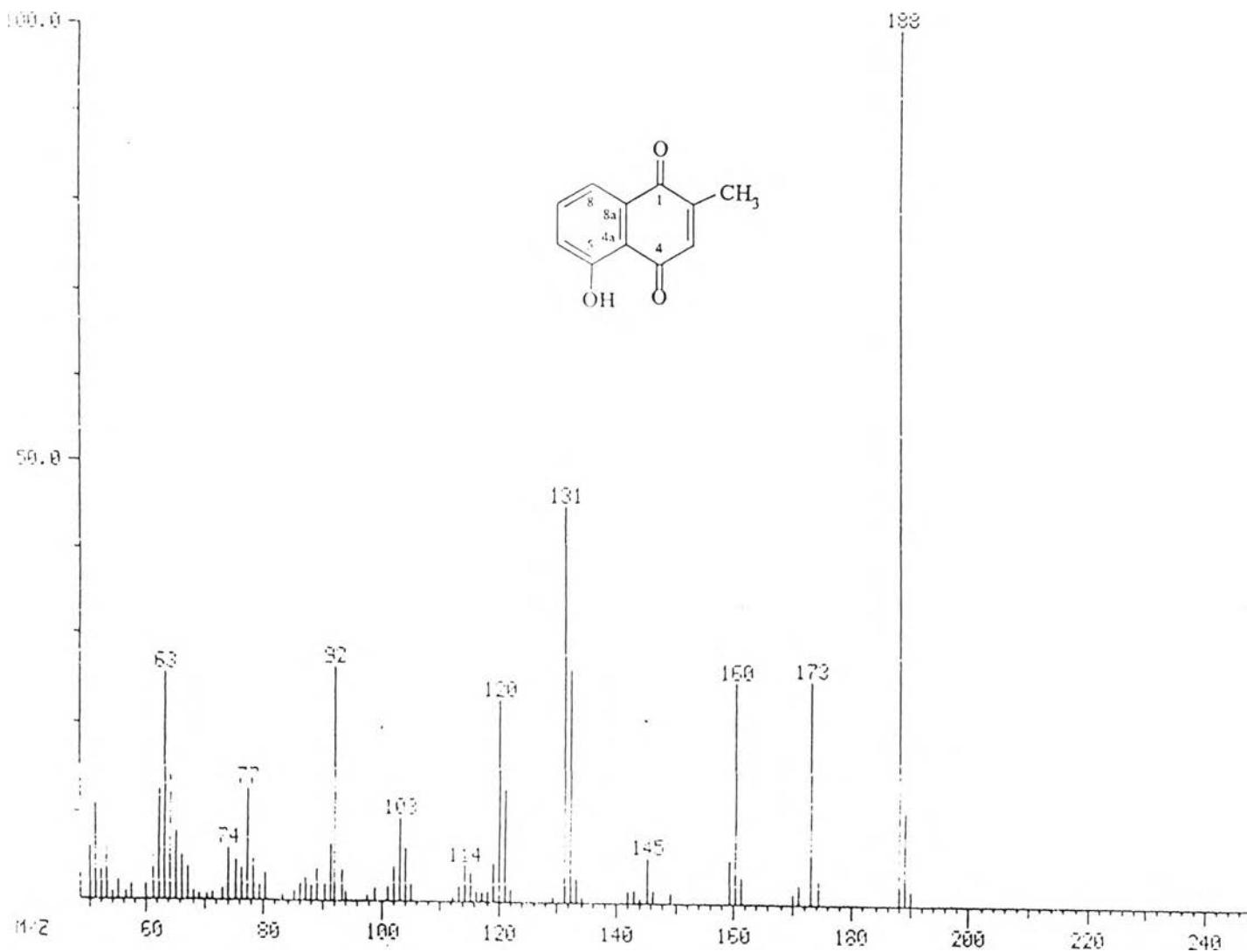


Figure 12 The EI mass spectrum of compound 18

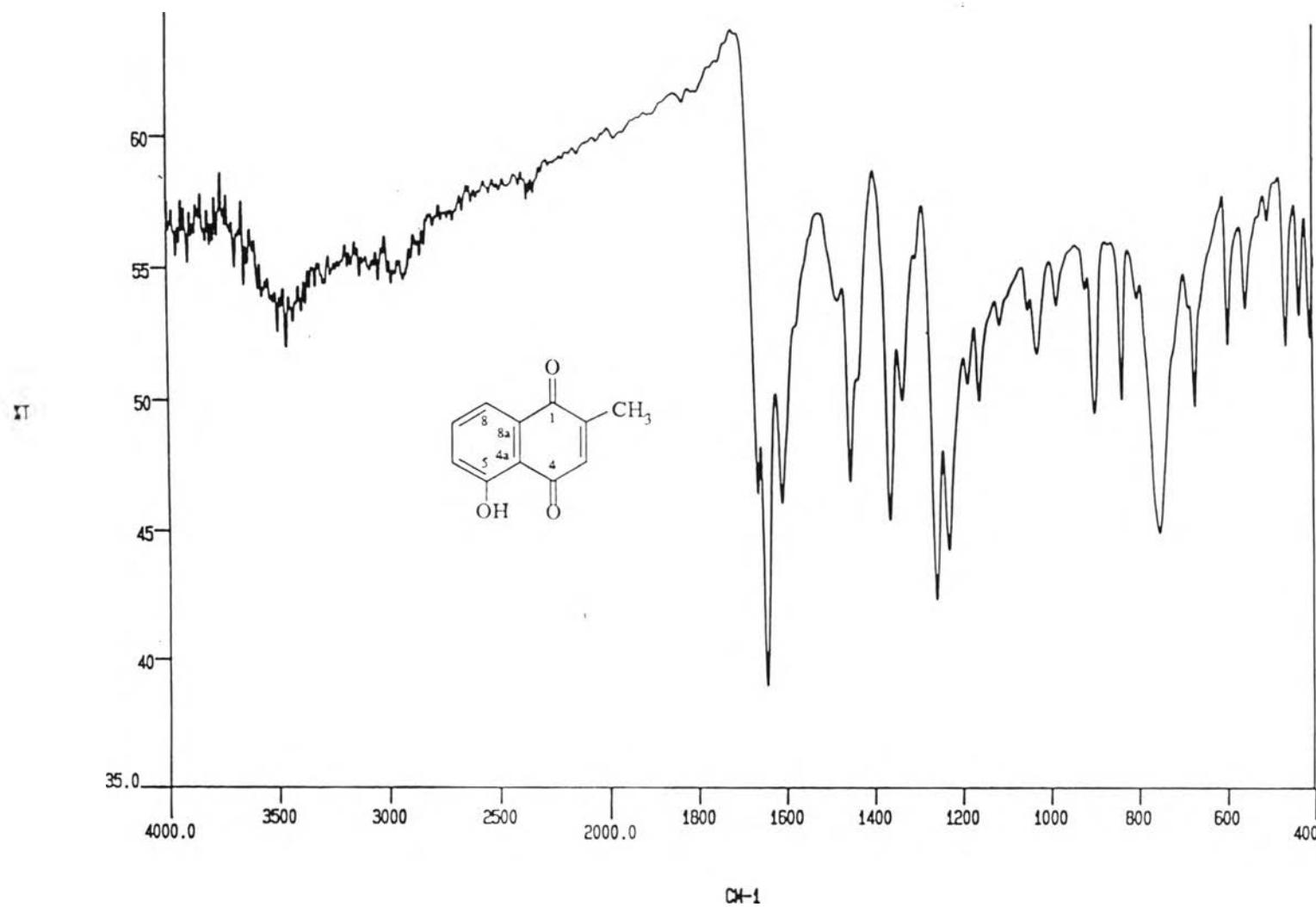


Figure 13 The IR spectrum of compound 18 (in KBr disc)

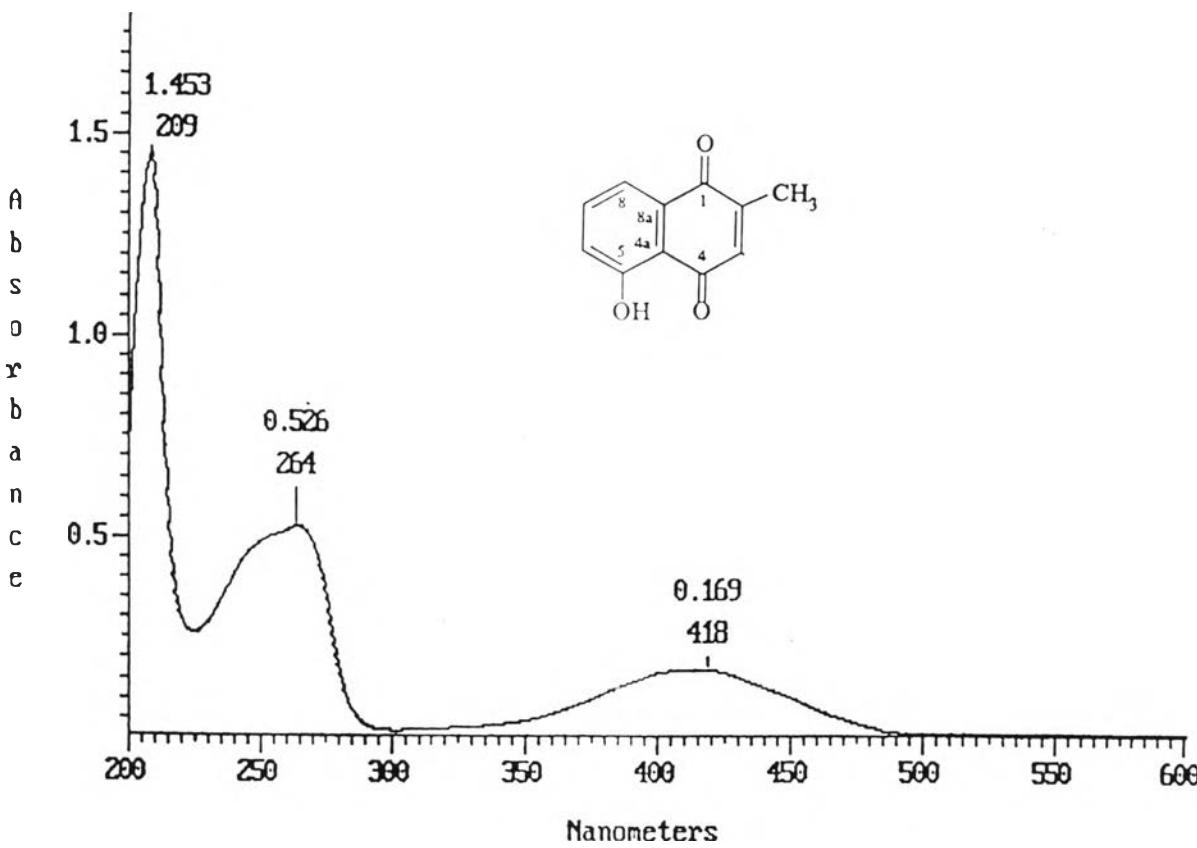


Figure 14 The UV spectrum of compound 18 (in MeOH)

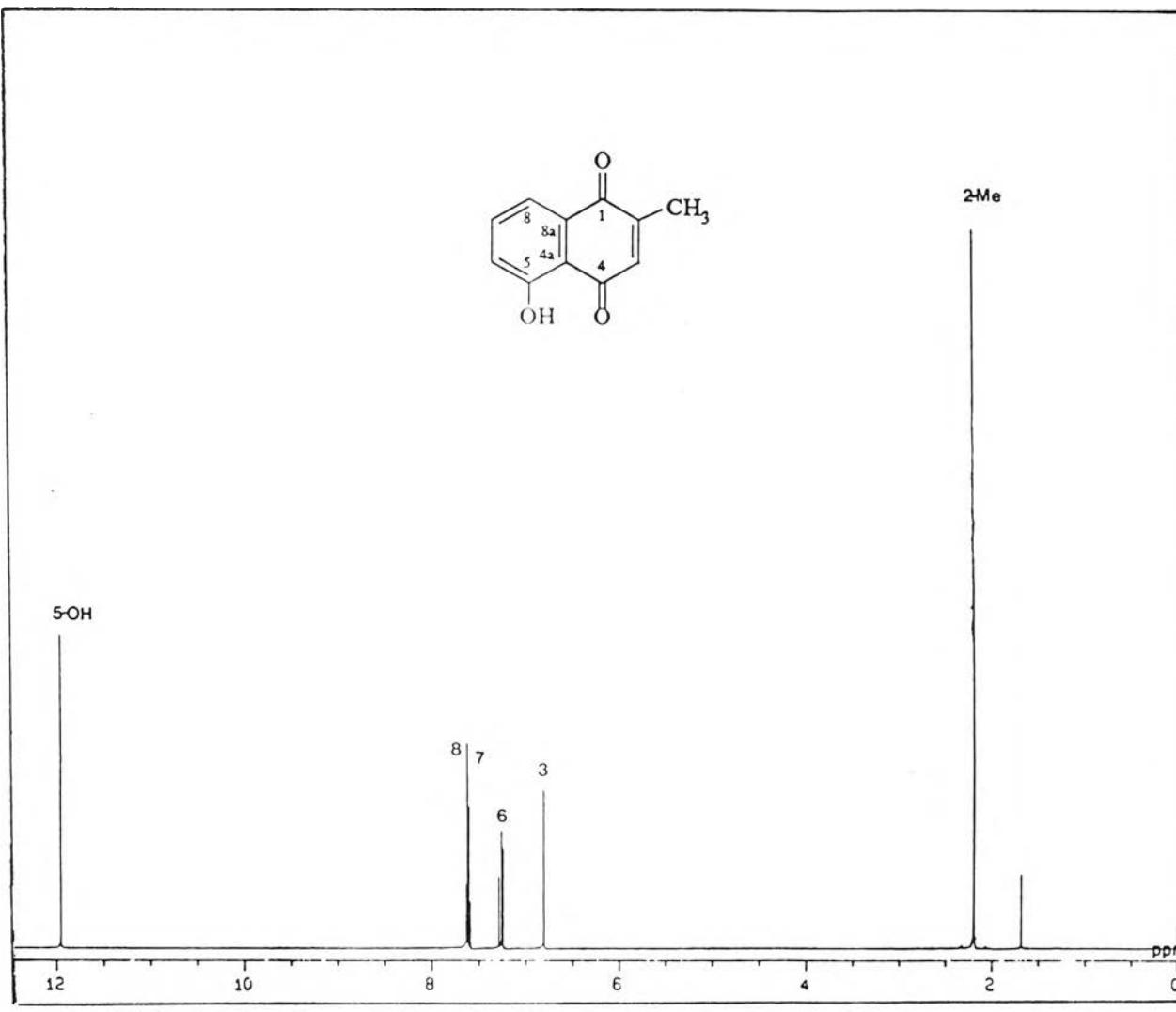


Figure 15 The ¹H NMR (500 MHz) spectrum of compound 18 (in CDCl₃)

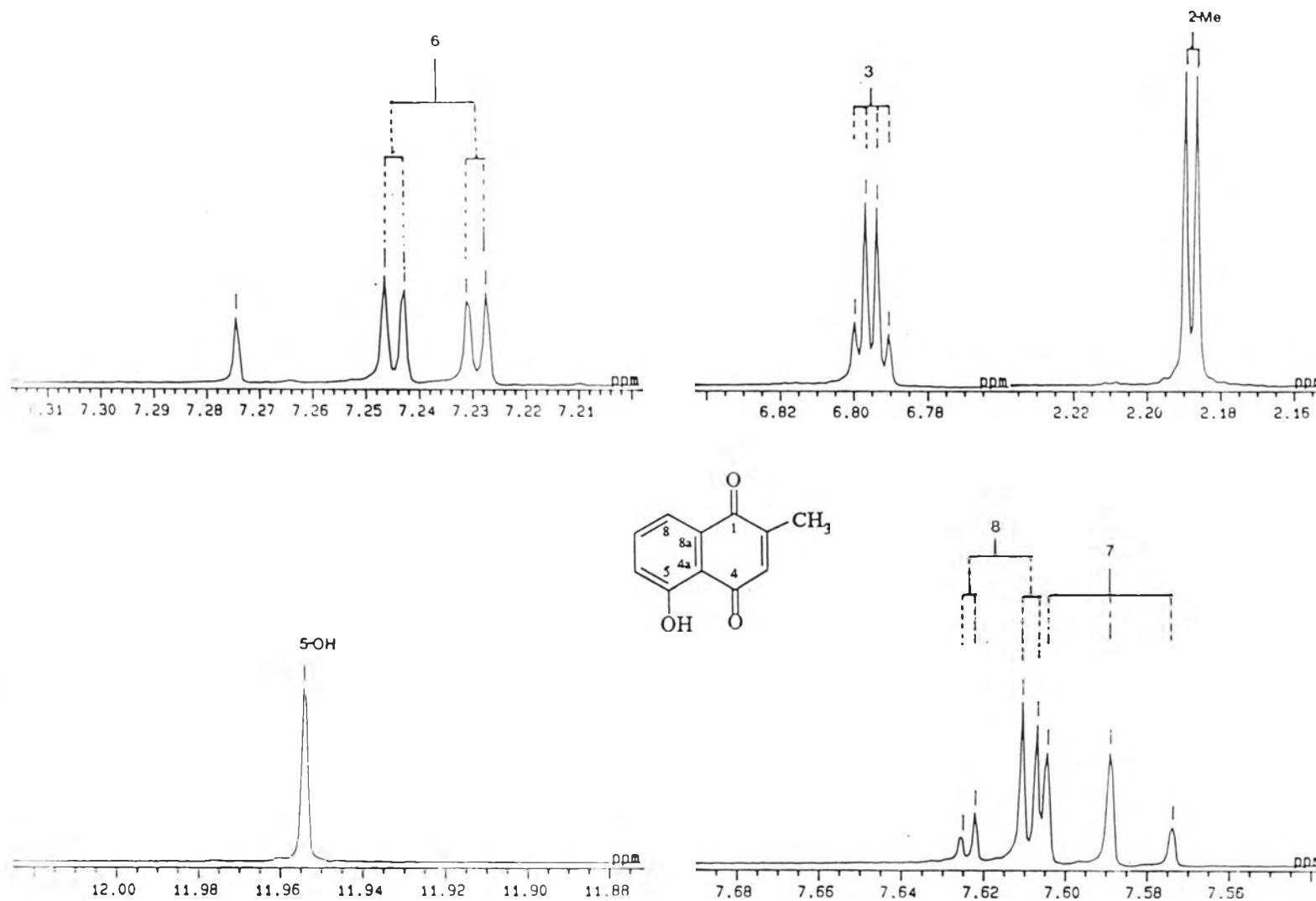


Figure 16 Expansion of the ^1H NMR (500 MHz) spectrum of compound 18

(in CDCl_3) : δ_{H} 2.15-11.96

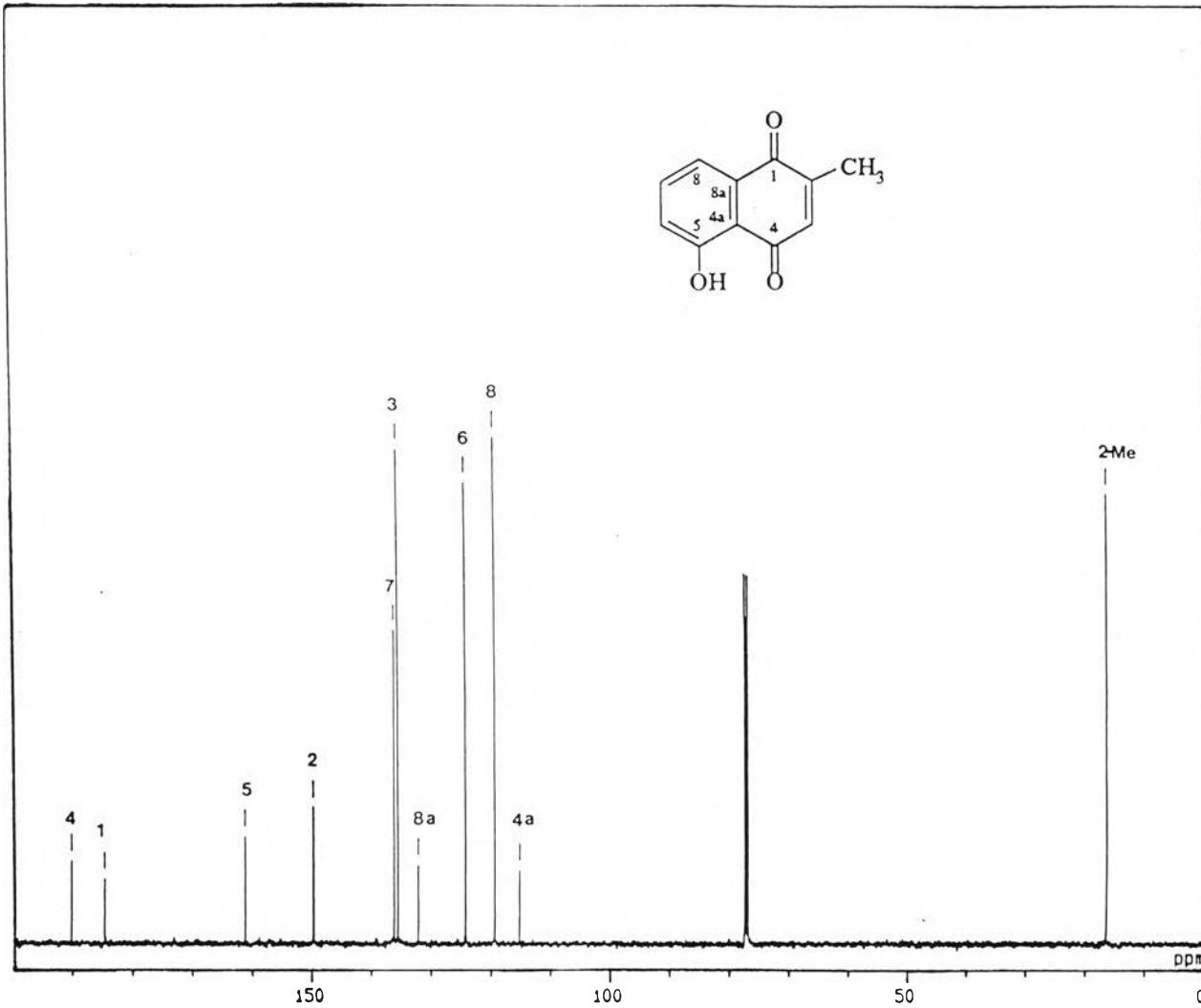


Figure 17 The ^{13}C NMR (125 MHz) spectrum of compound 18 (in CDCl_3)

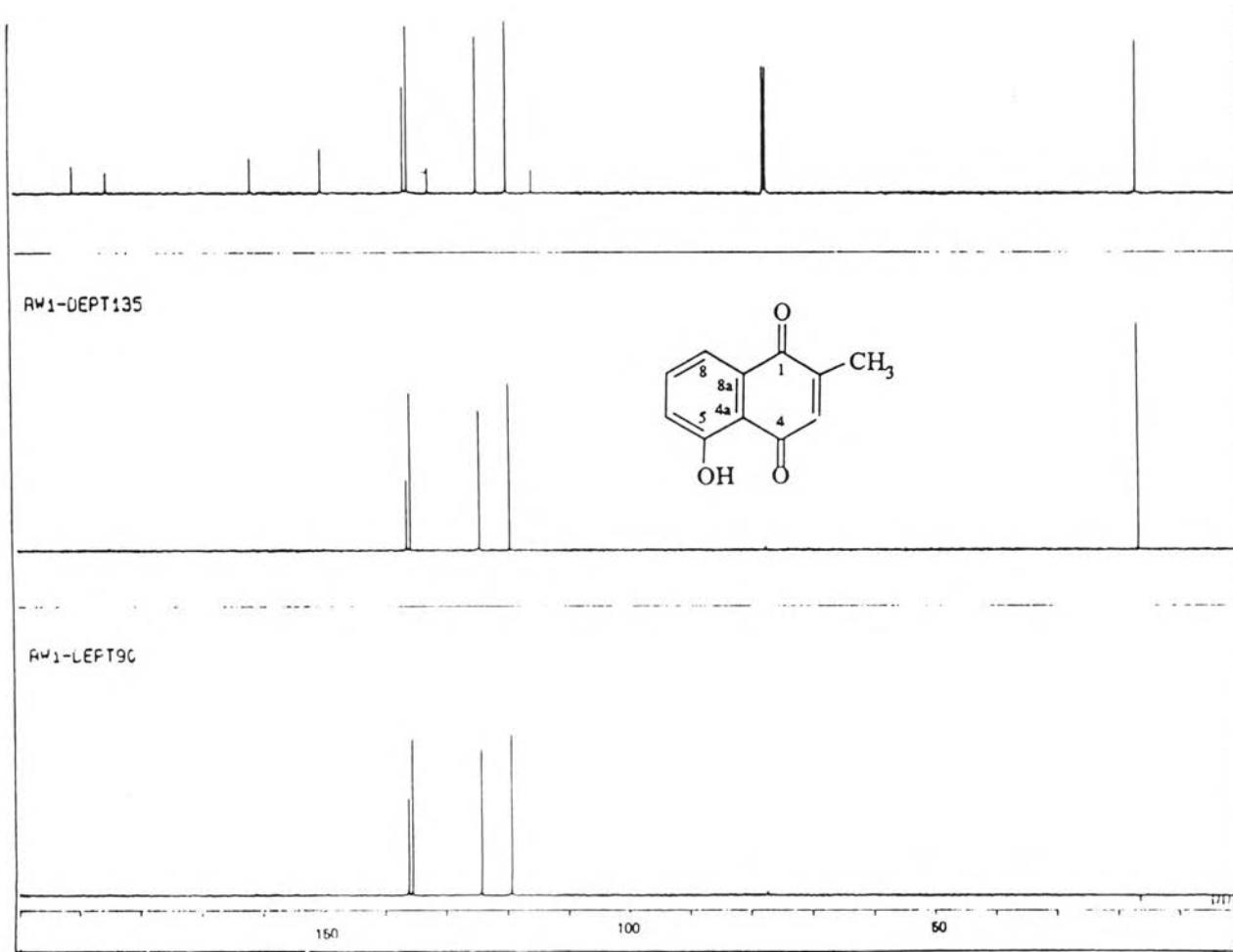


Figure 18 The DEPT (125 MHz) spectrum of compound 18 (in CDCl₃)

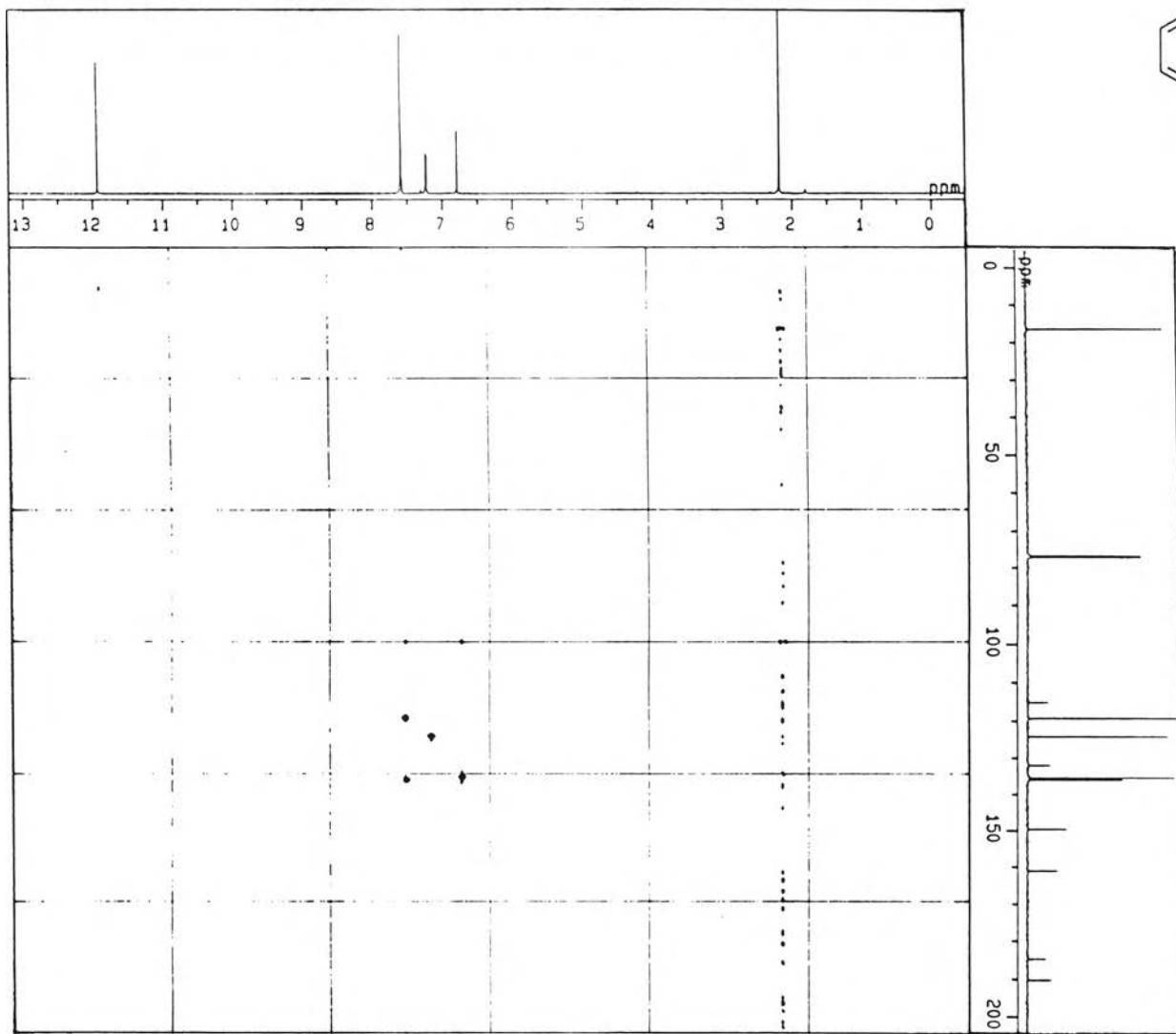
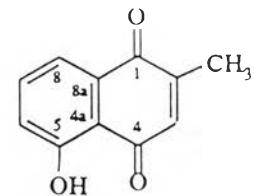


Figure 19 The HMQC spectrum of compound 18 (in CDCl₃)

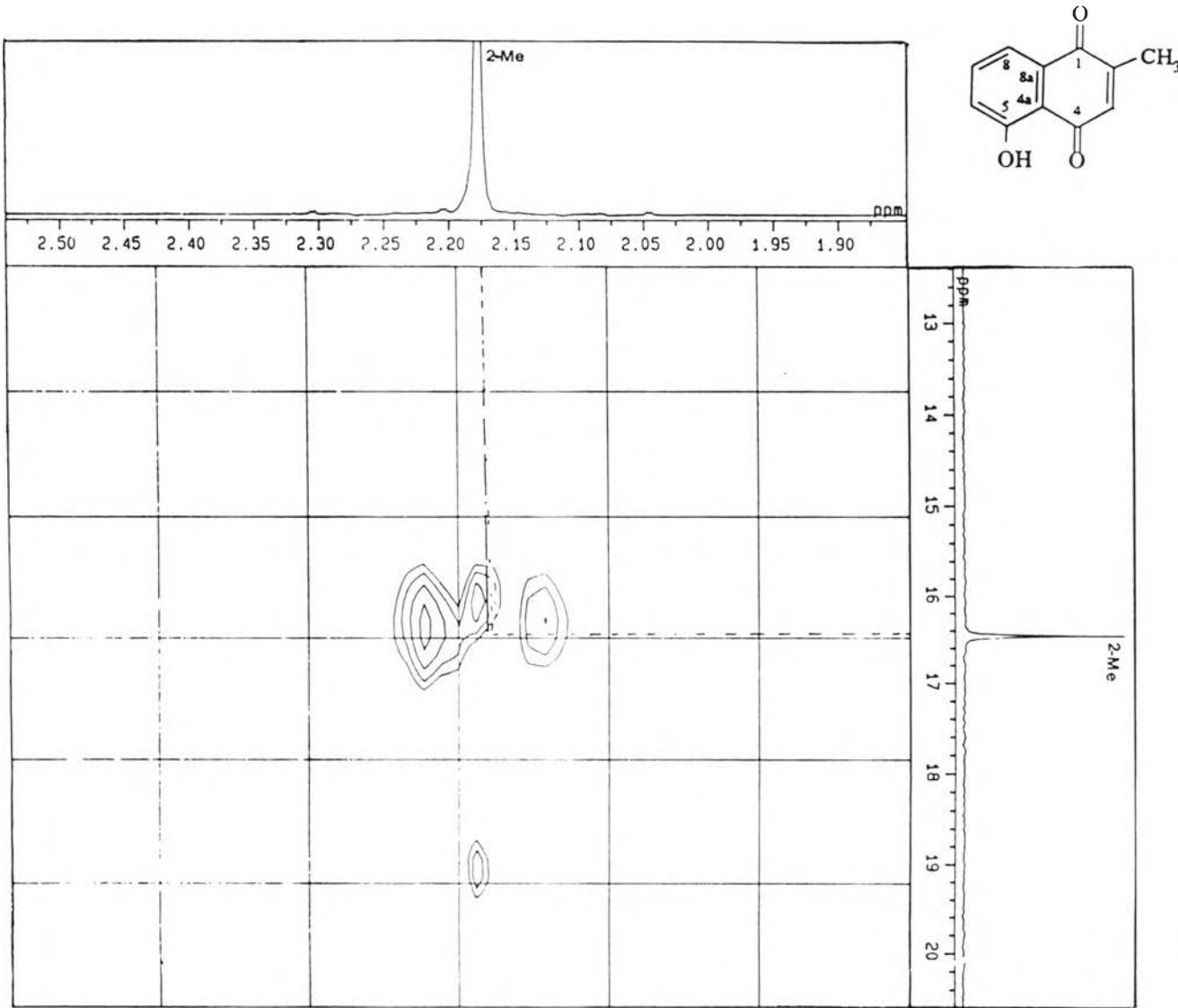


Figure 20 Expansion of the HMQC spectrum of compound 18 (in CDCl_3) :

δ_{H} 1.90-2.50 ; δ_{C} 13.00-20.00 ppm

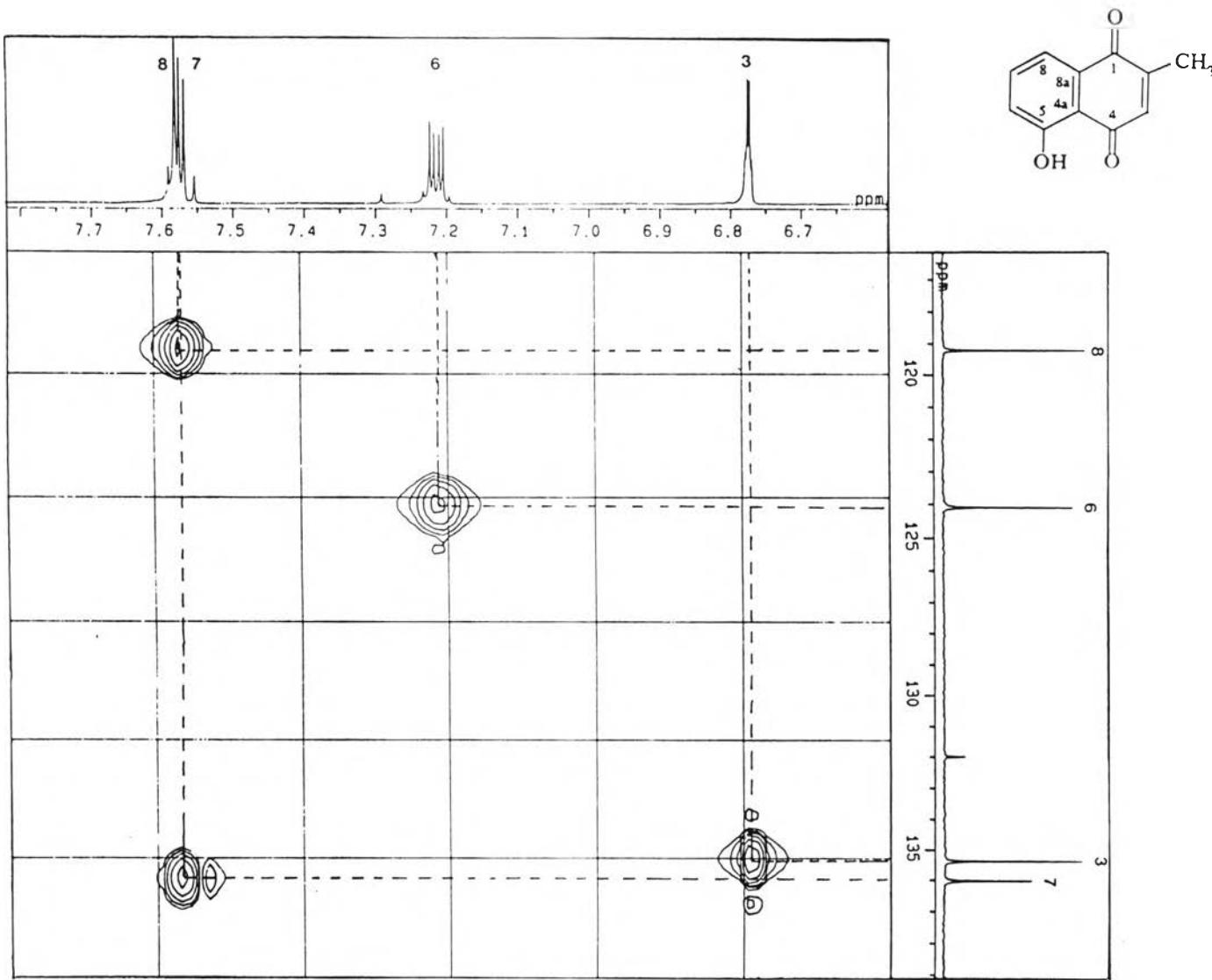


Figure 21 Expansion of the HMQC spectrum of compound 18 (in CDCl_3) :

δ_{H} 6.70-7.70 ; δ_{C} 123.00-139.00 ppm

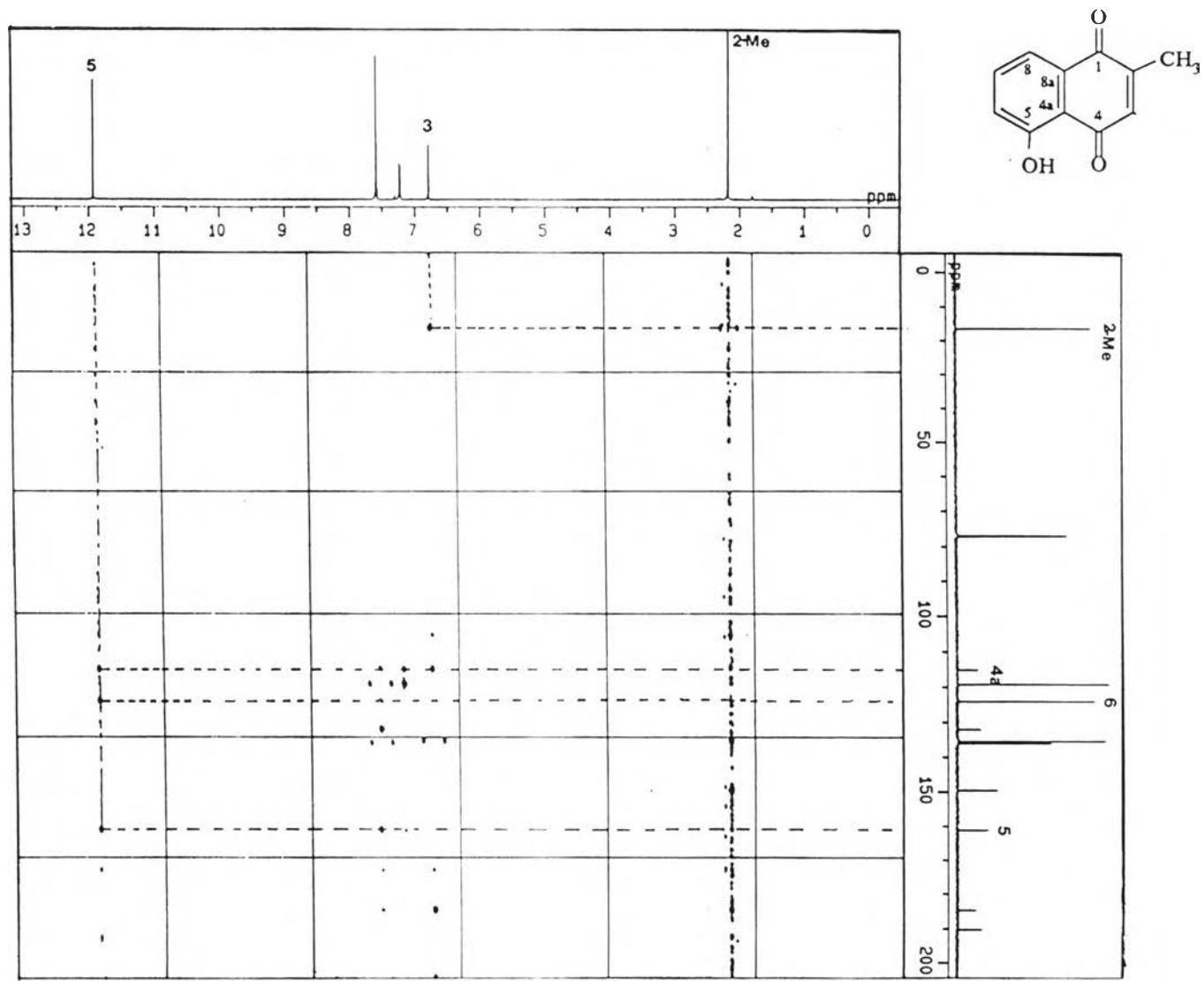


Figure 22 The HMBC spectrum of compound 18 (in CDCl_3)

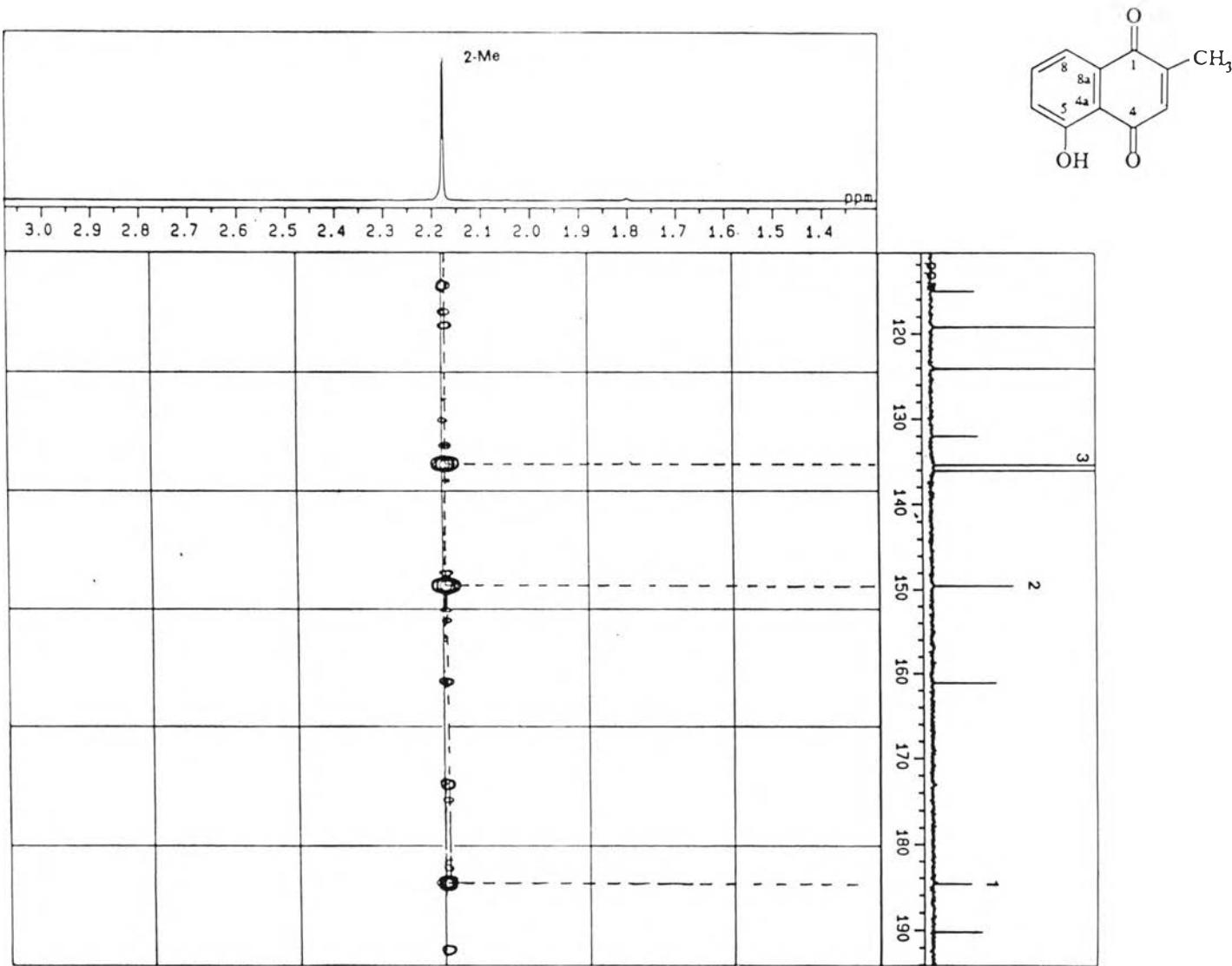


Figure 23 Expansion of the HMBC spectrum of compound 18 (in CDCl_3) :
 δ_{H} 1.40-3.00 ; δ_{C} 116.00-191.00 ppm

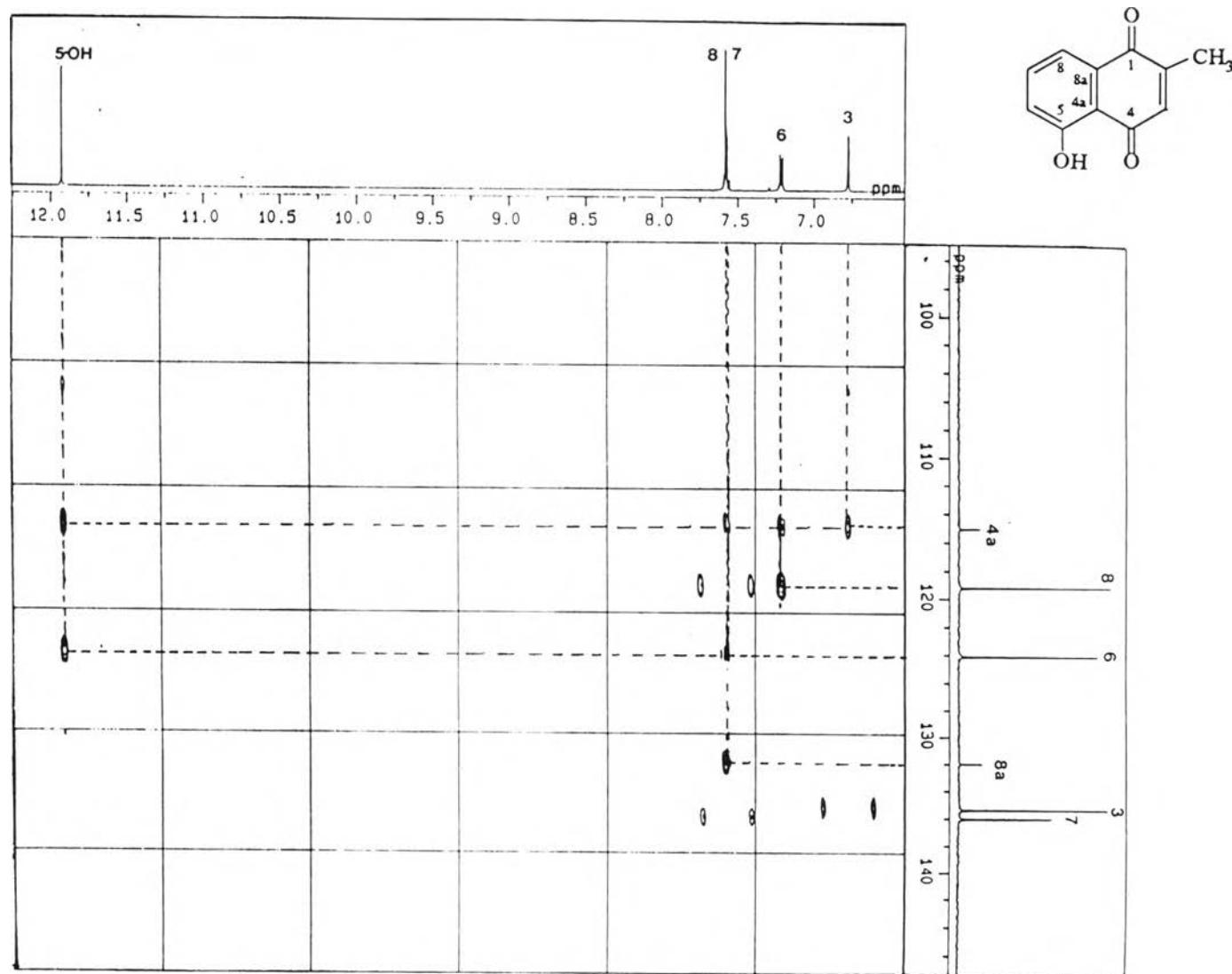


Figure 24 Expansion of the HMBC spectrum of compound 18 (in CDCl₃) :

δ_{H} 6.50-12.00 ; δ_{C} 98.00-143.00 ppm

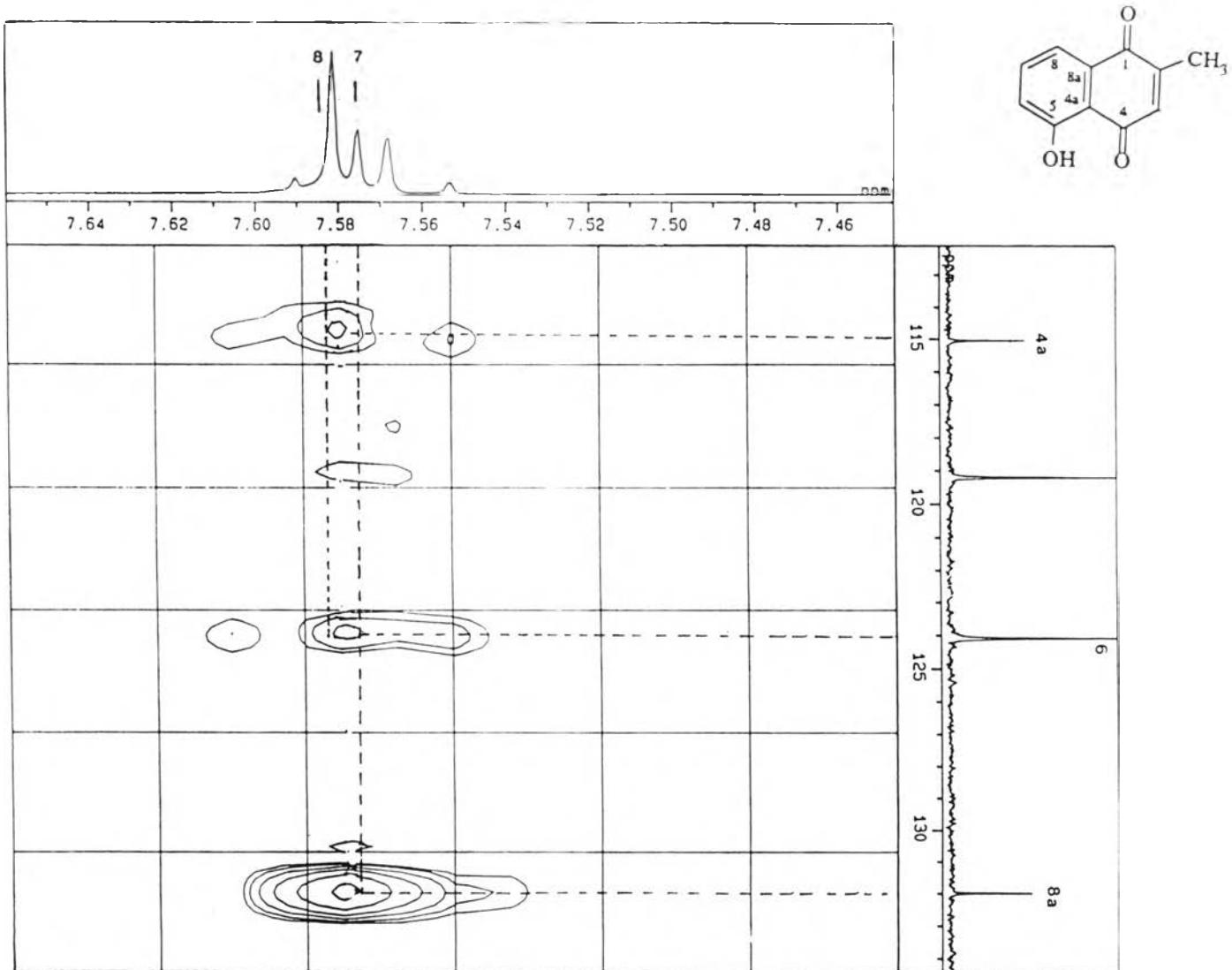


Figure 25 Expansion of the HMBC spectrum of compound 18 (in CDCl_3) :

δ_{H} 7.46-7.64 ; δ_{C} 113.00-134.00 ppm

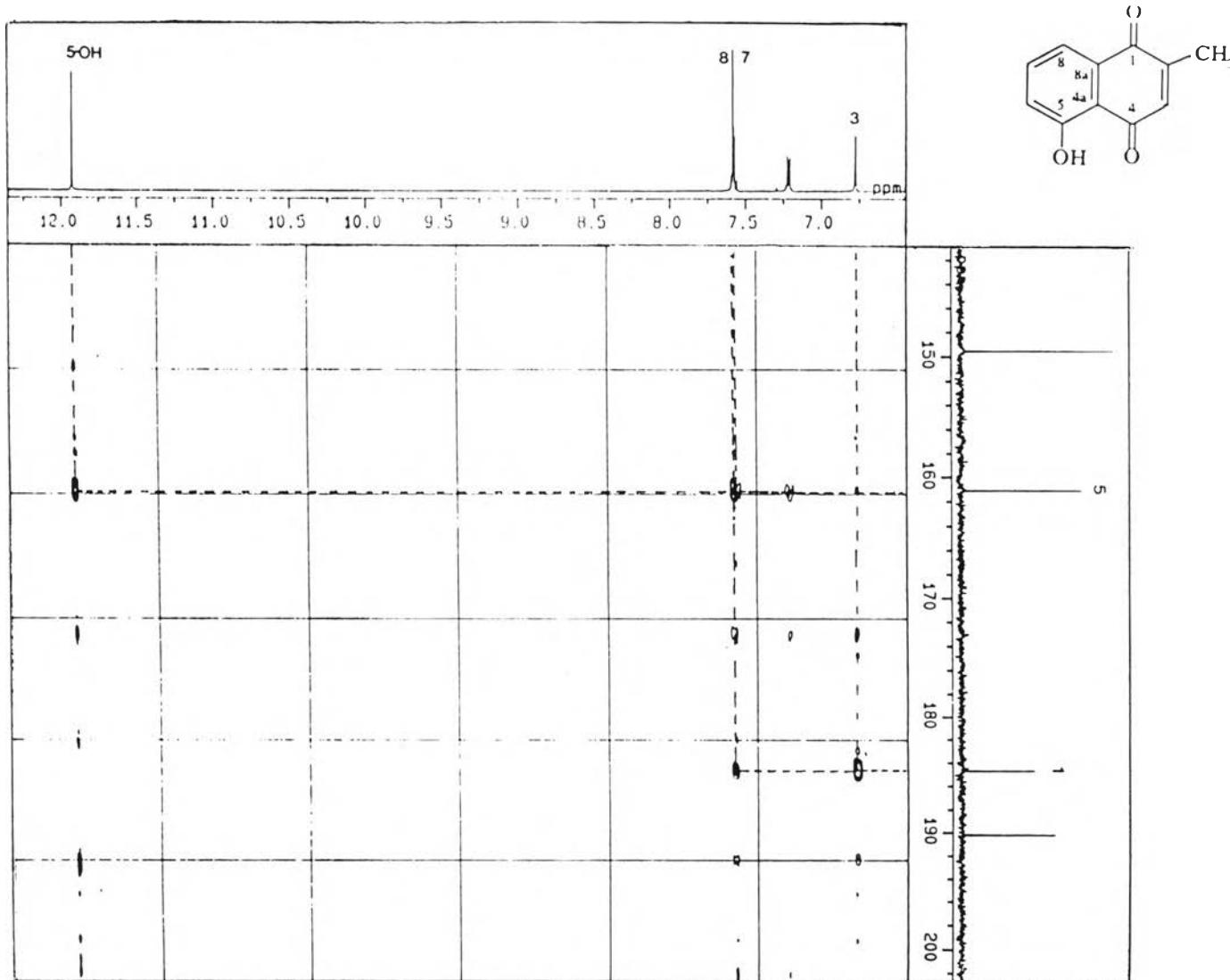


Figure 26 Expansion of the HMBC spectrum of compound 18 (in CDCl_3) :

δ_{H} 6.90-12.10 ; δ_{C} 146.00-201.00 ppm

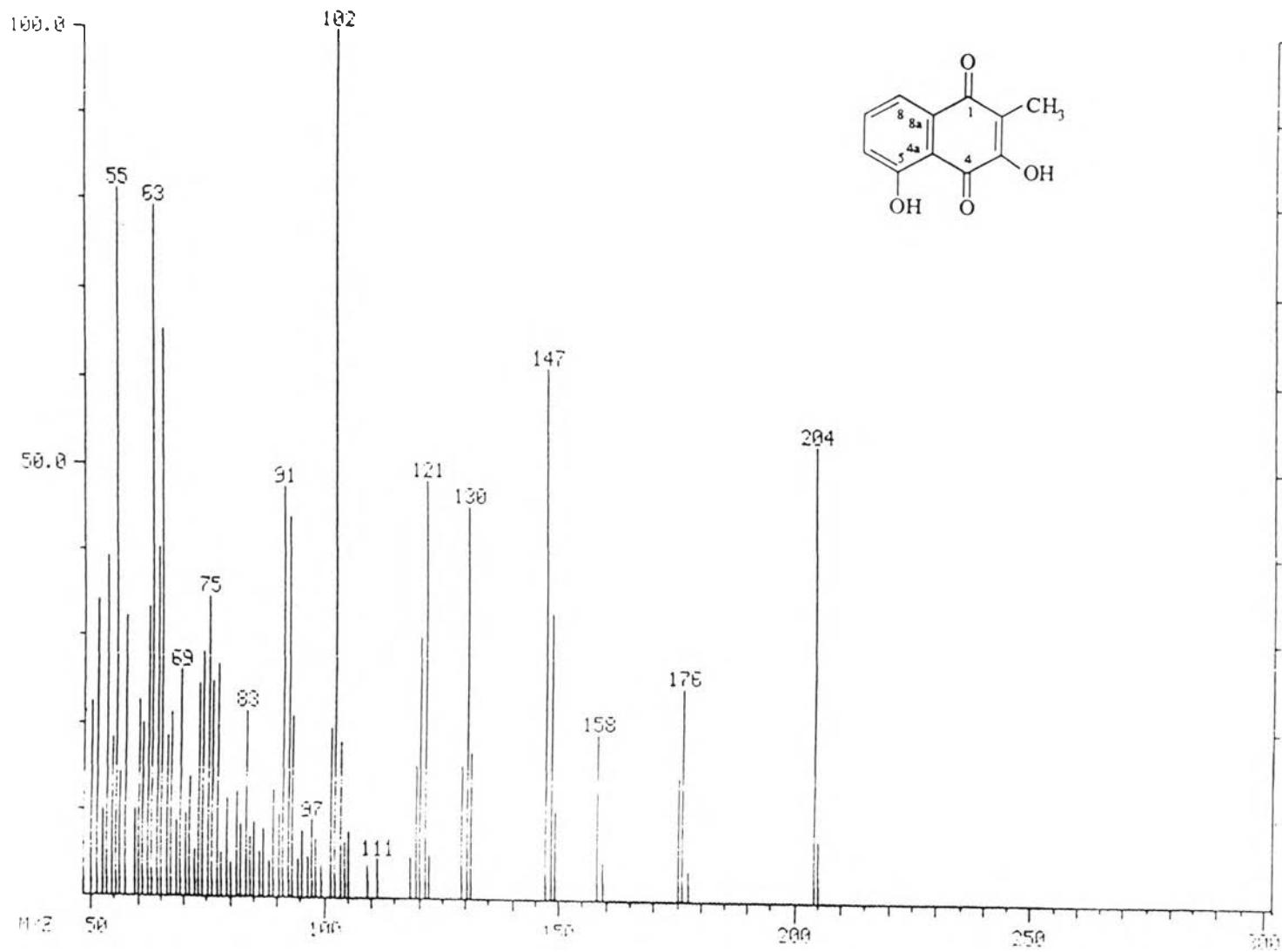


Figure 27 The EI mass spectrum of compound 19

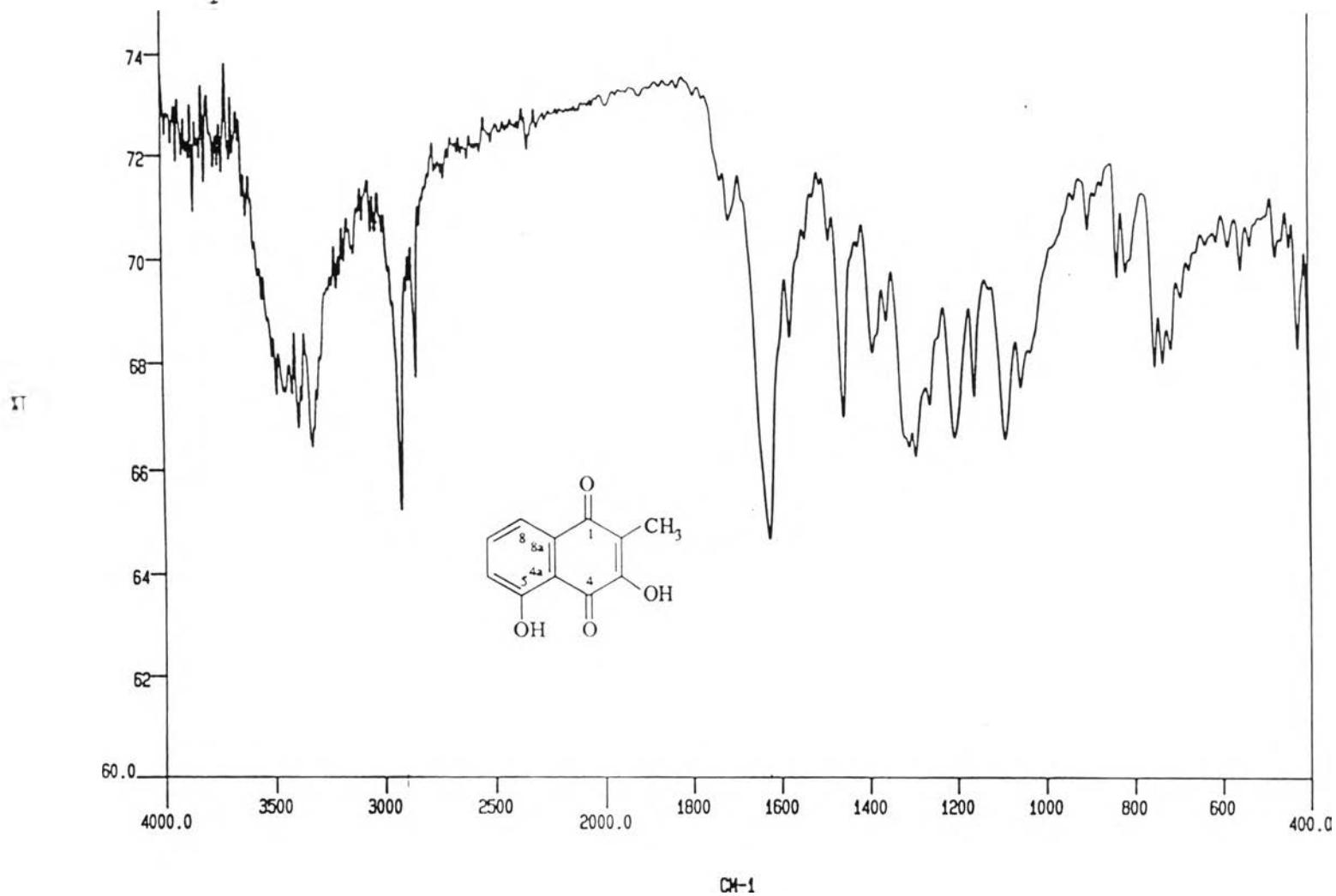


Figure 28 The IR spectrum of compound 19 (in KBr disc)

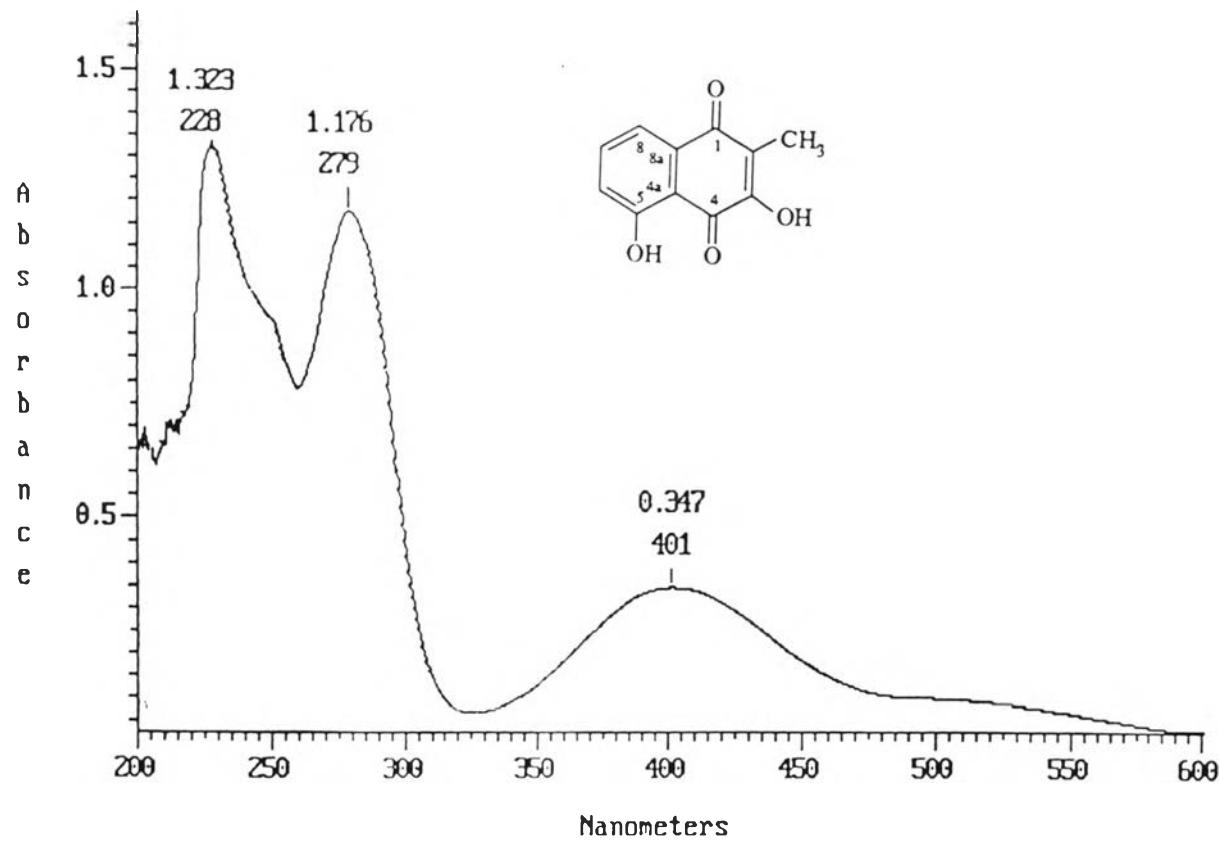


Figure 29 The UV spectrum of compound 19 (in MeOH)

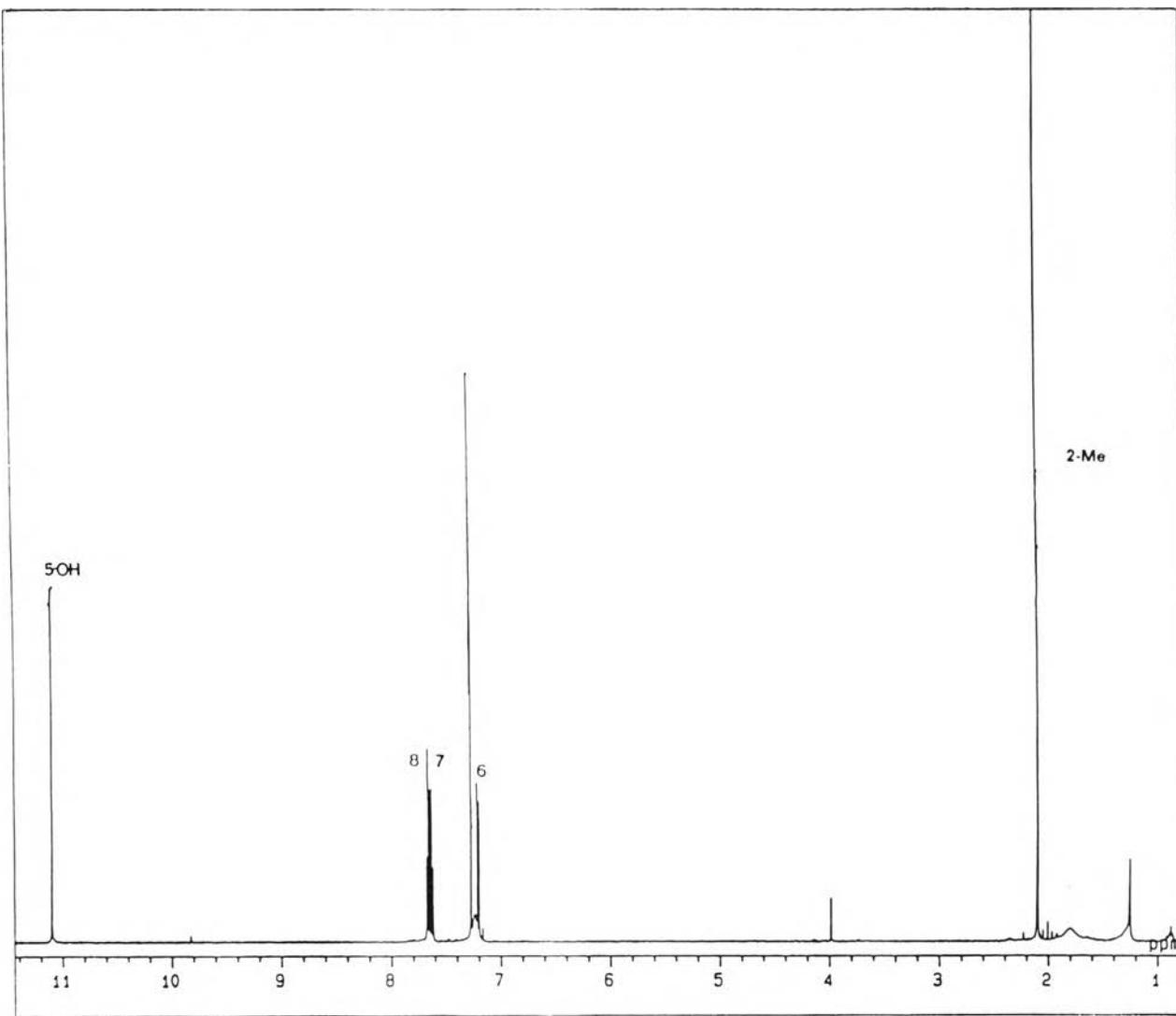
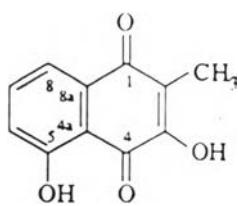


Figure 30 The ¹H NMR (500 MHz) spectrum of compound 19 (in CDCl₃)

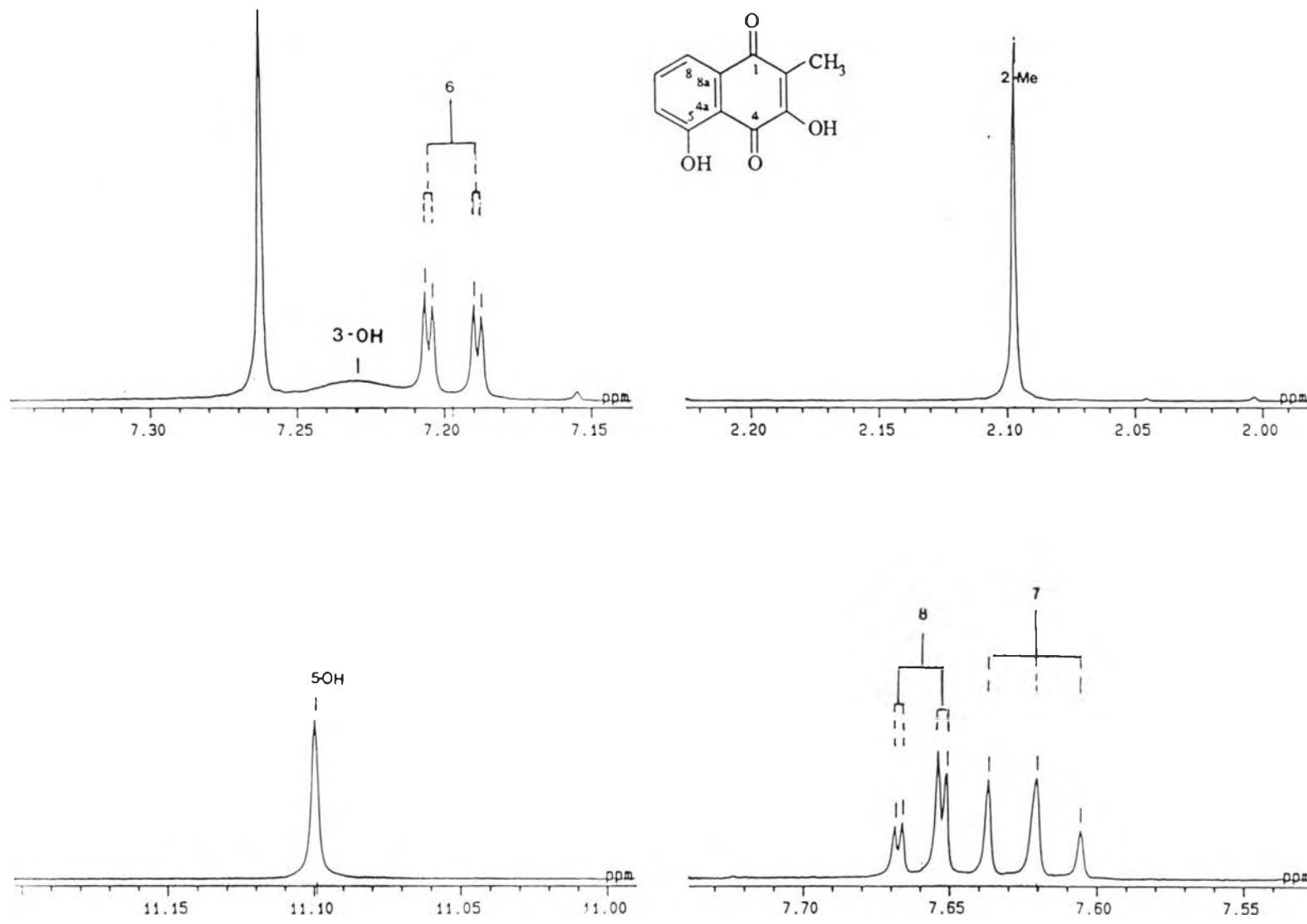


Figure 31 Expansion of the ^1H NMR (500 MHz) spectrum of compound 19
 (in CDCl_3) : δ_{H} 2.00-11.15 ppm

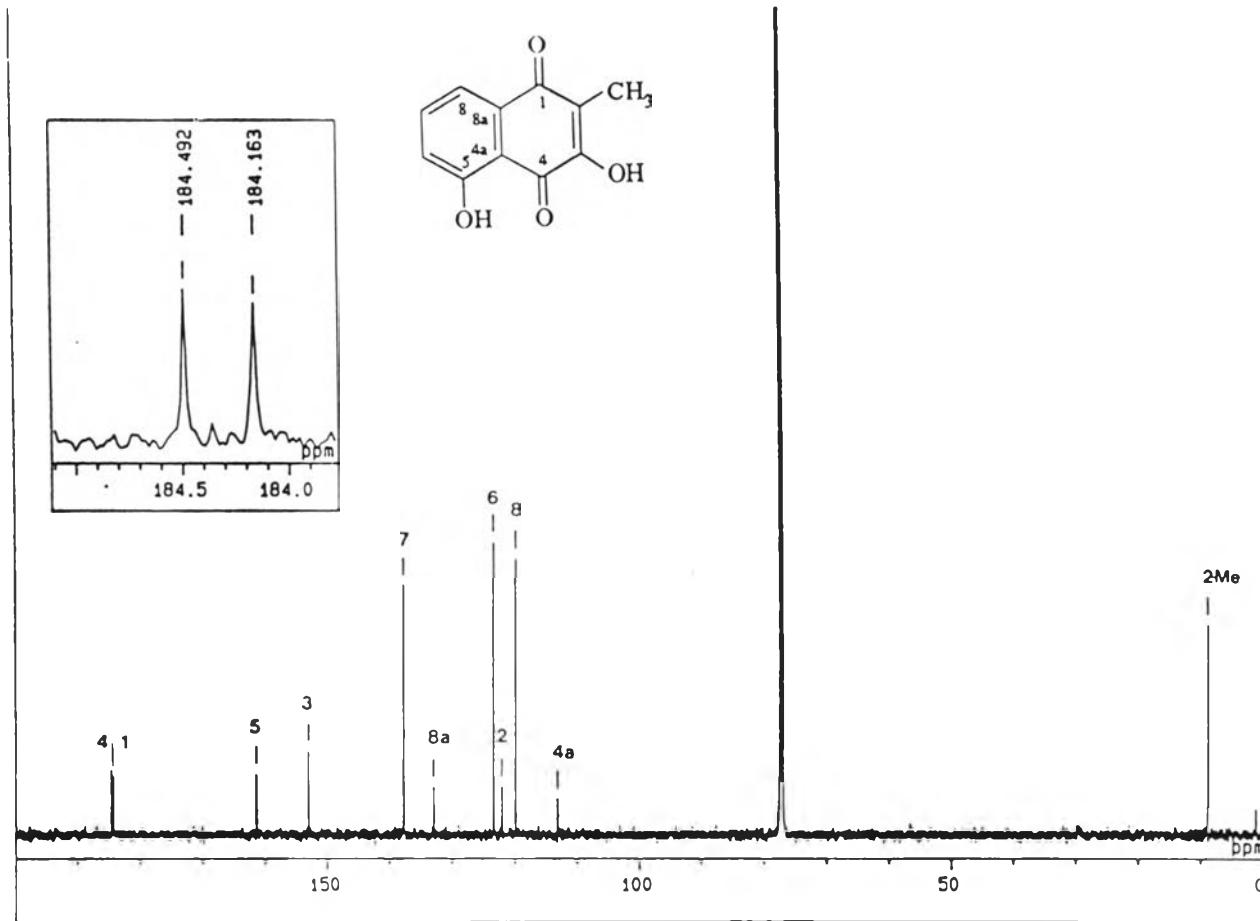


Figure 32 The ^{13}C NMR (125 MHz) of compound 19 (in CDCl₃)

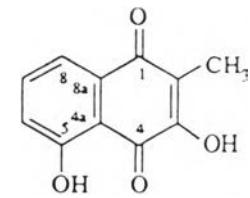
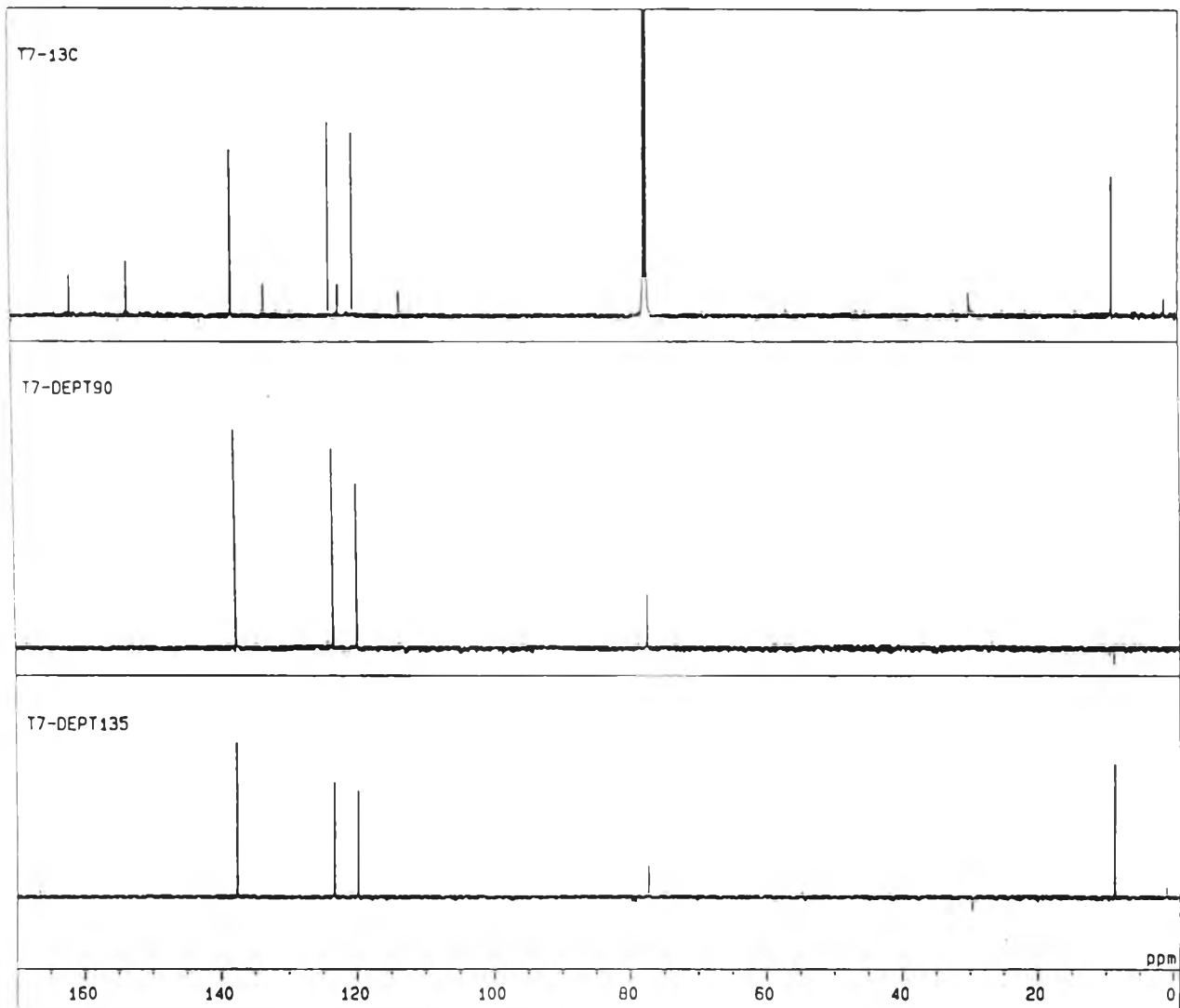


Figure 33 The DEPT (125 MHz) spectrum of compound 19 (in CDCl₃)

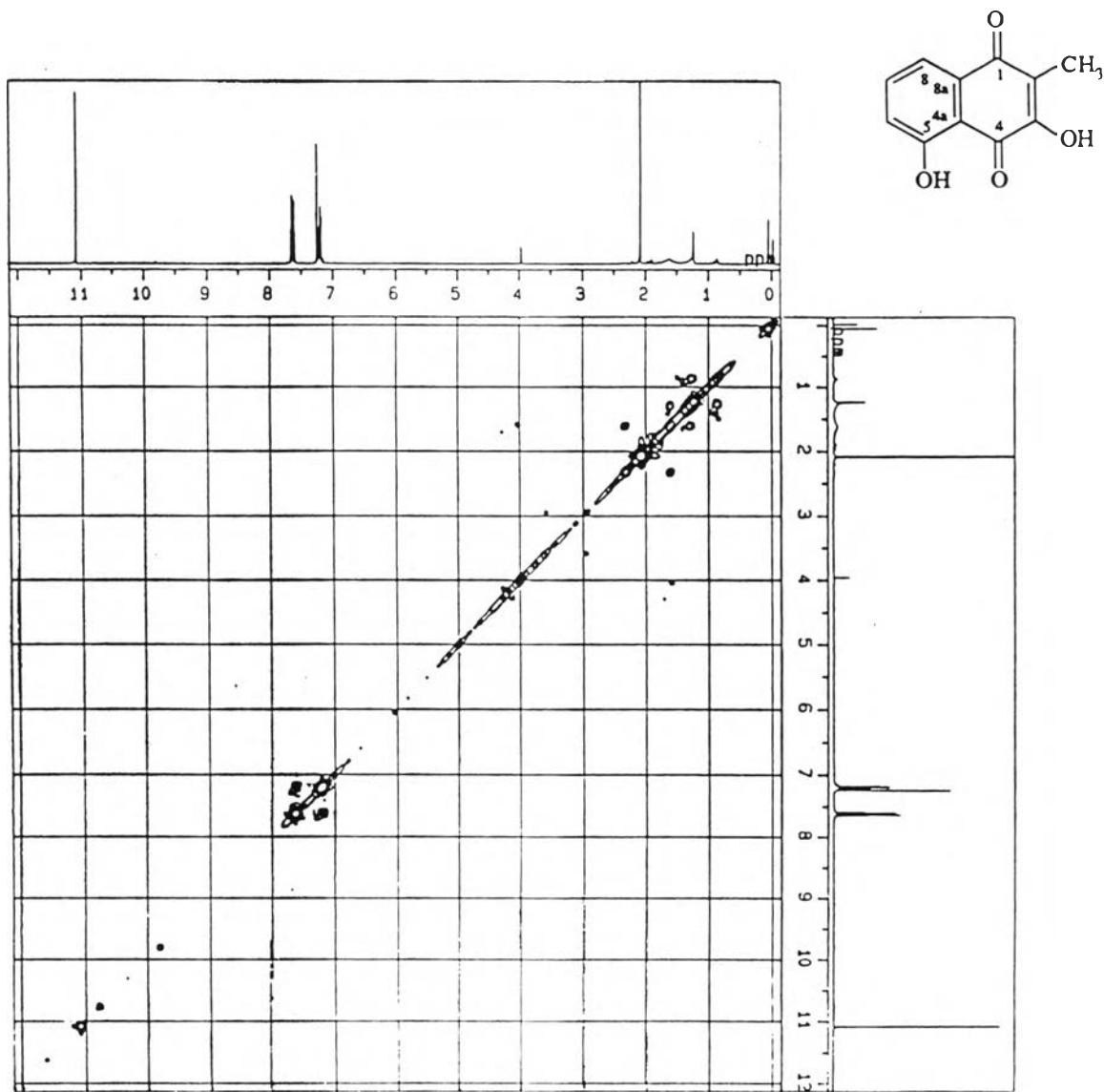


Figure 34 The ^1H - ^1H COSY (500 MHz) spectrum of compound 19 (in CDCl_3)

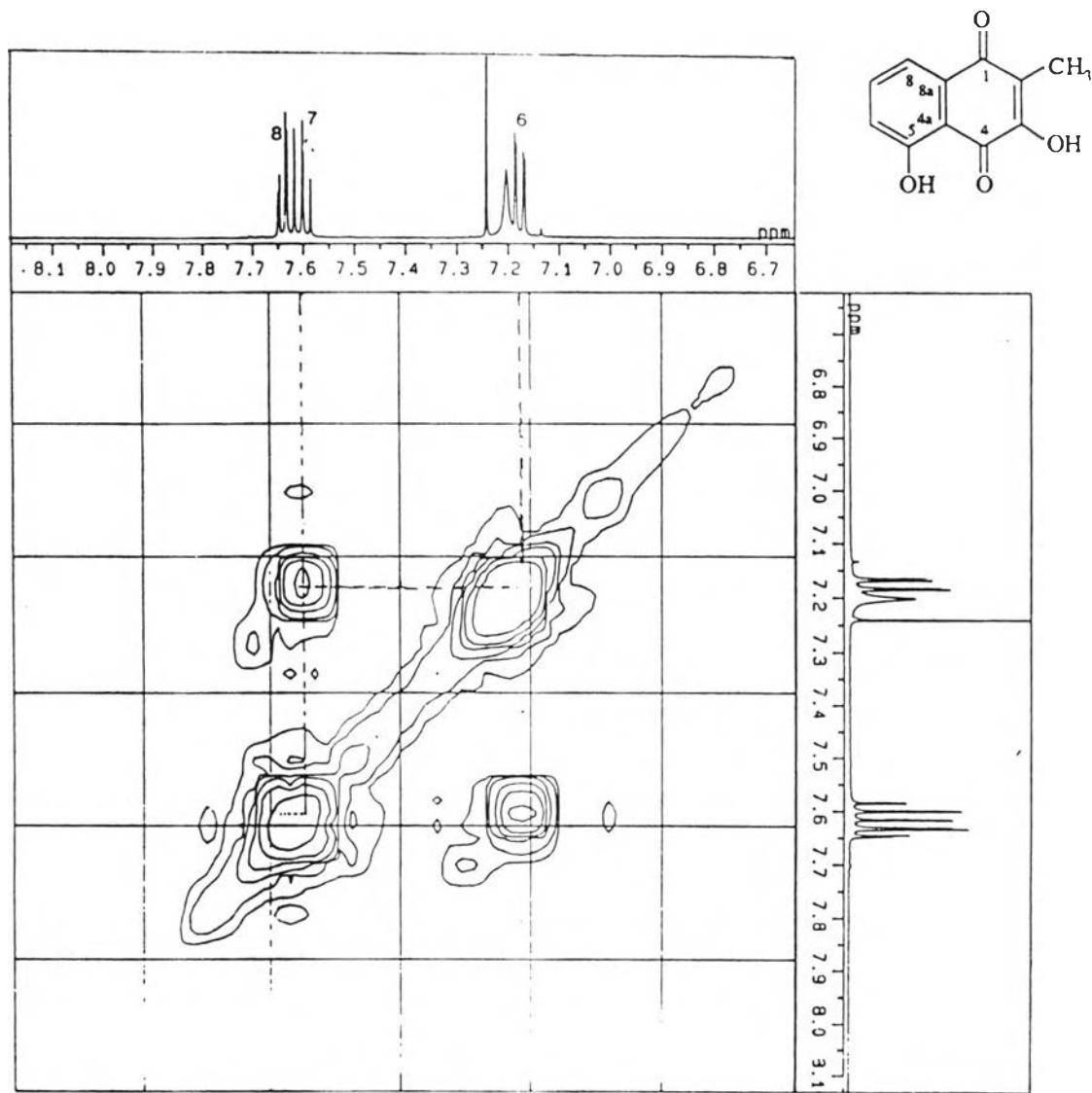


Figure 35 Expansion of the ^1H - ^1H COSY (500 MHz) spectrum of compound 19
(in CDCl_3) : δ_{H} 6.70-8.10 ppm

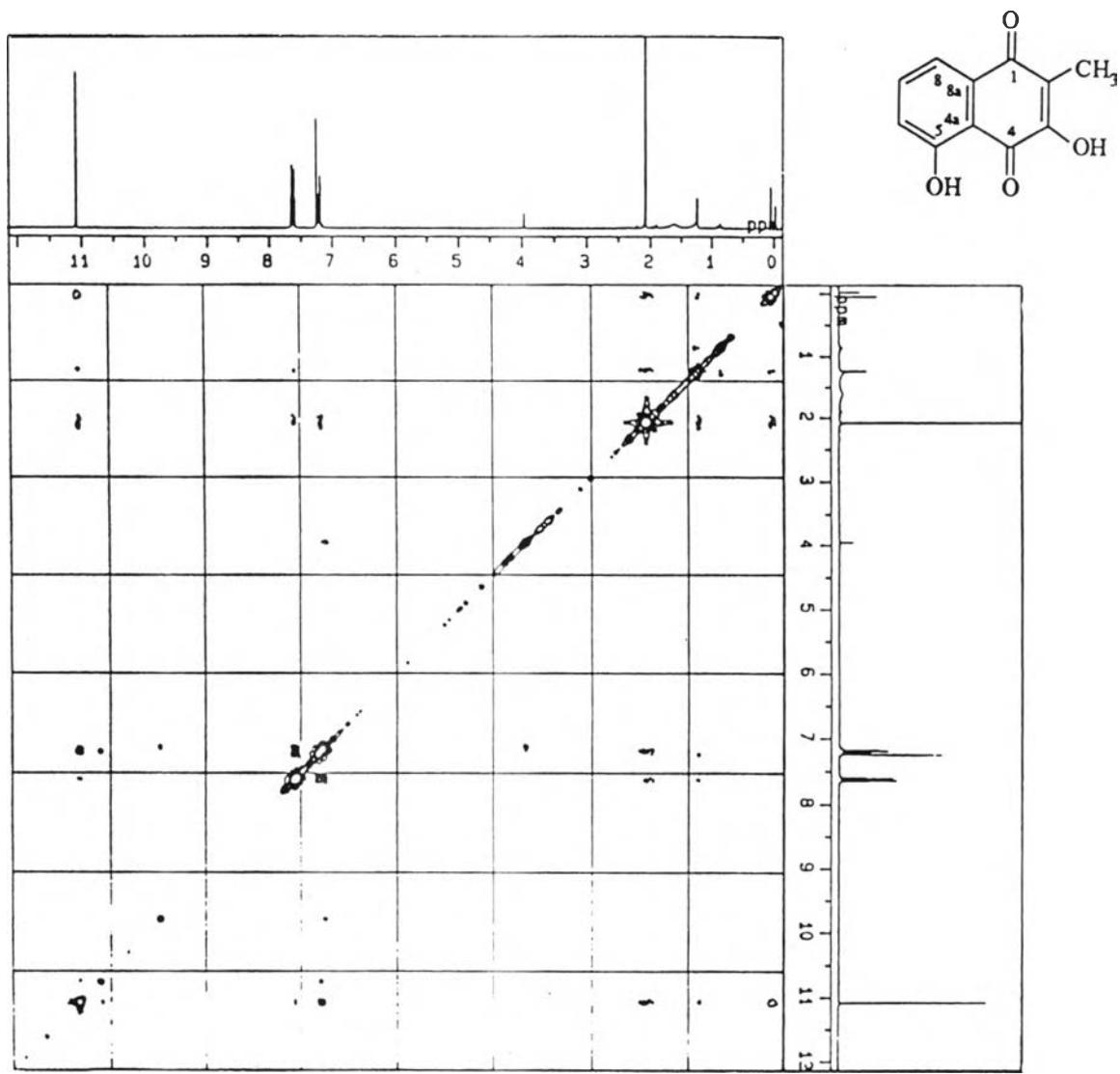


Figure 36 The NOESY (500 MHz) spectrum of compound 19 (in CDCl₃)

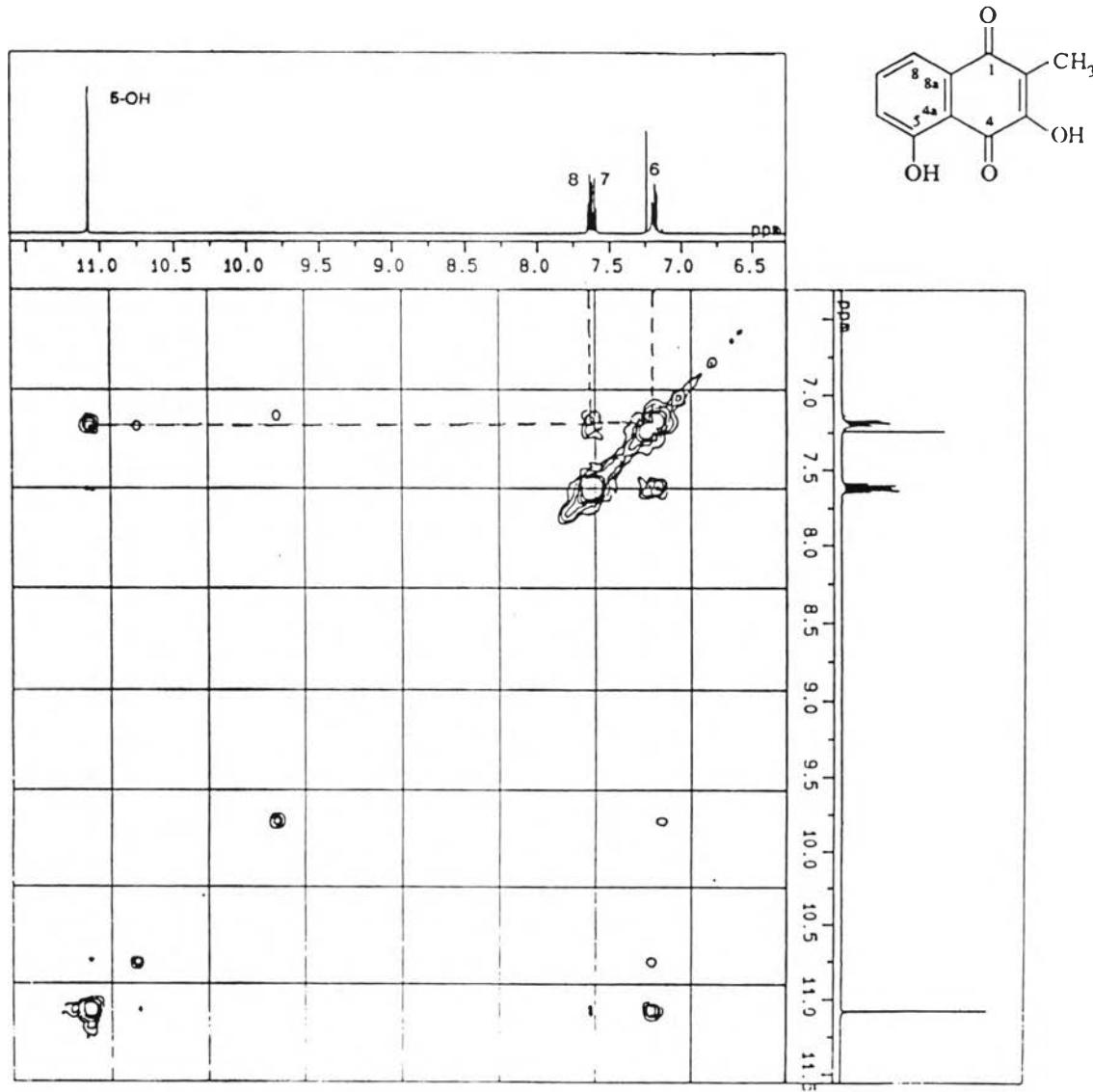


Figure 37 Expansion of the NOESY (500 MHz) spectrum of compound 19
(in CDCl_3) : δ_{H} 6.50-11.50 ppm

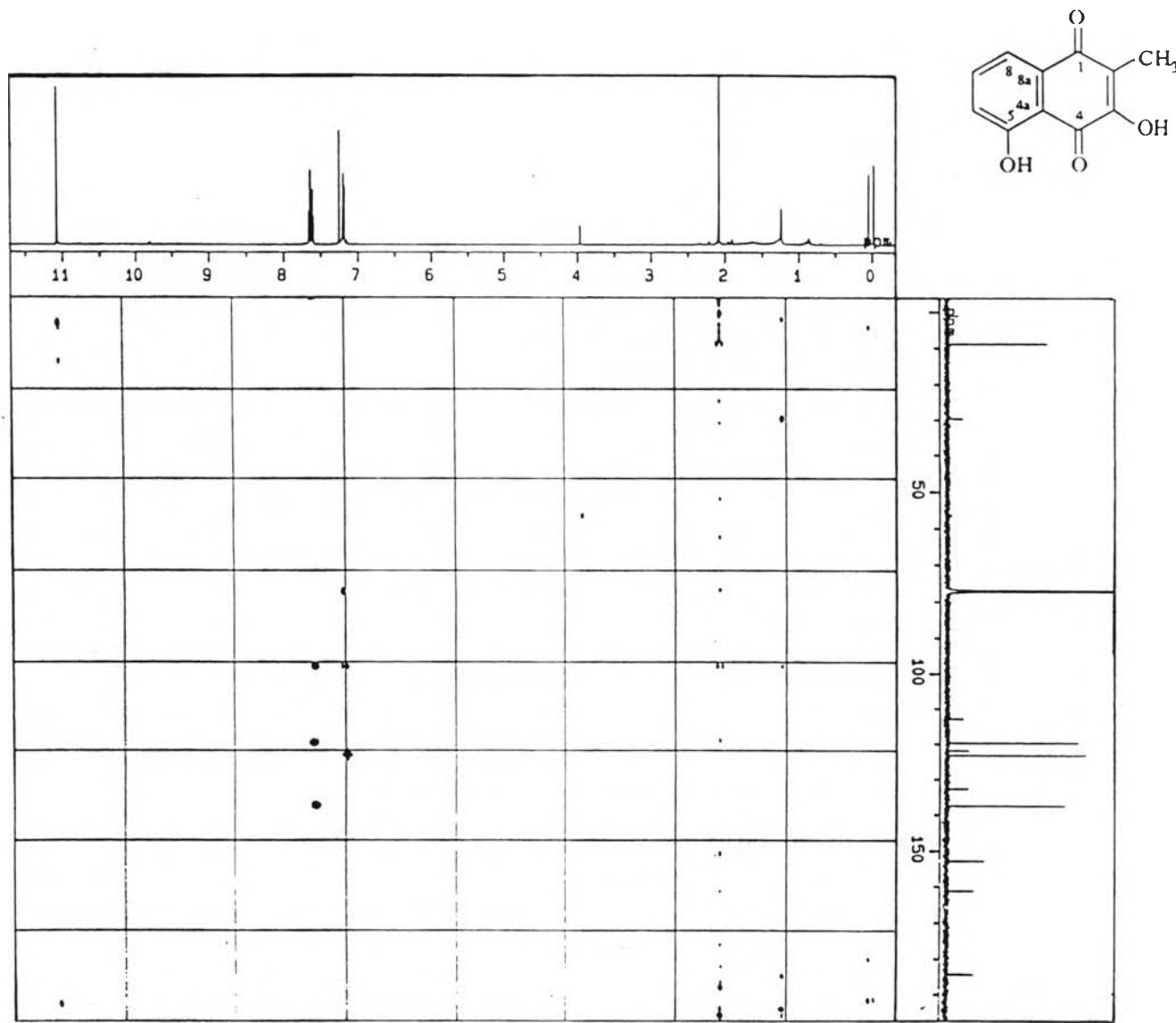


Figure 38 The HMQC spectrum of compound 19 (in CDCl_3)

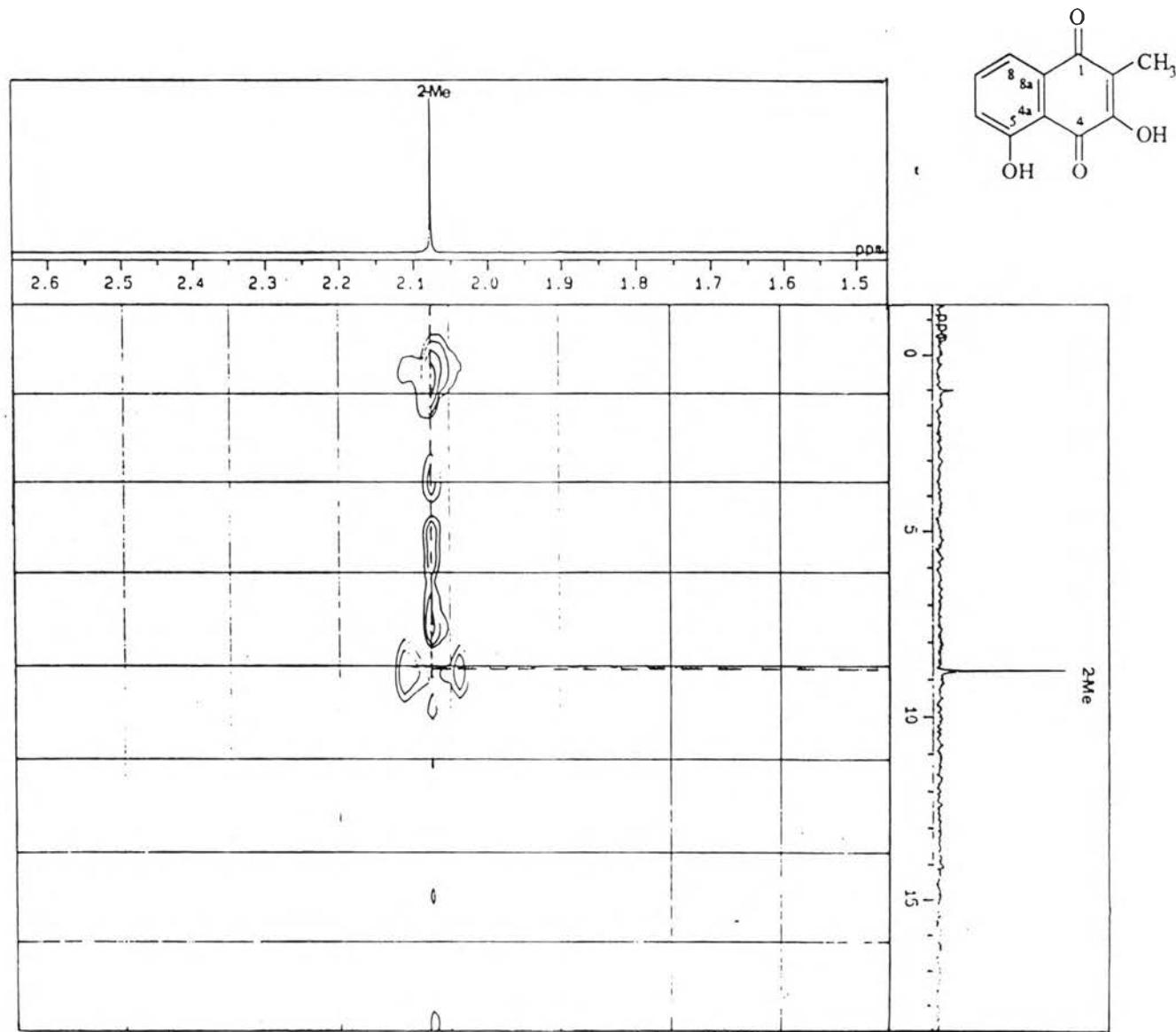
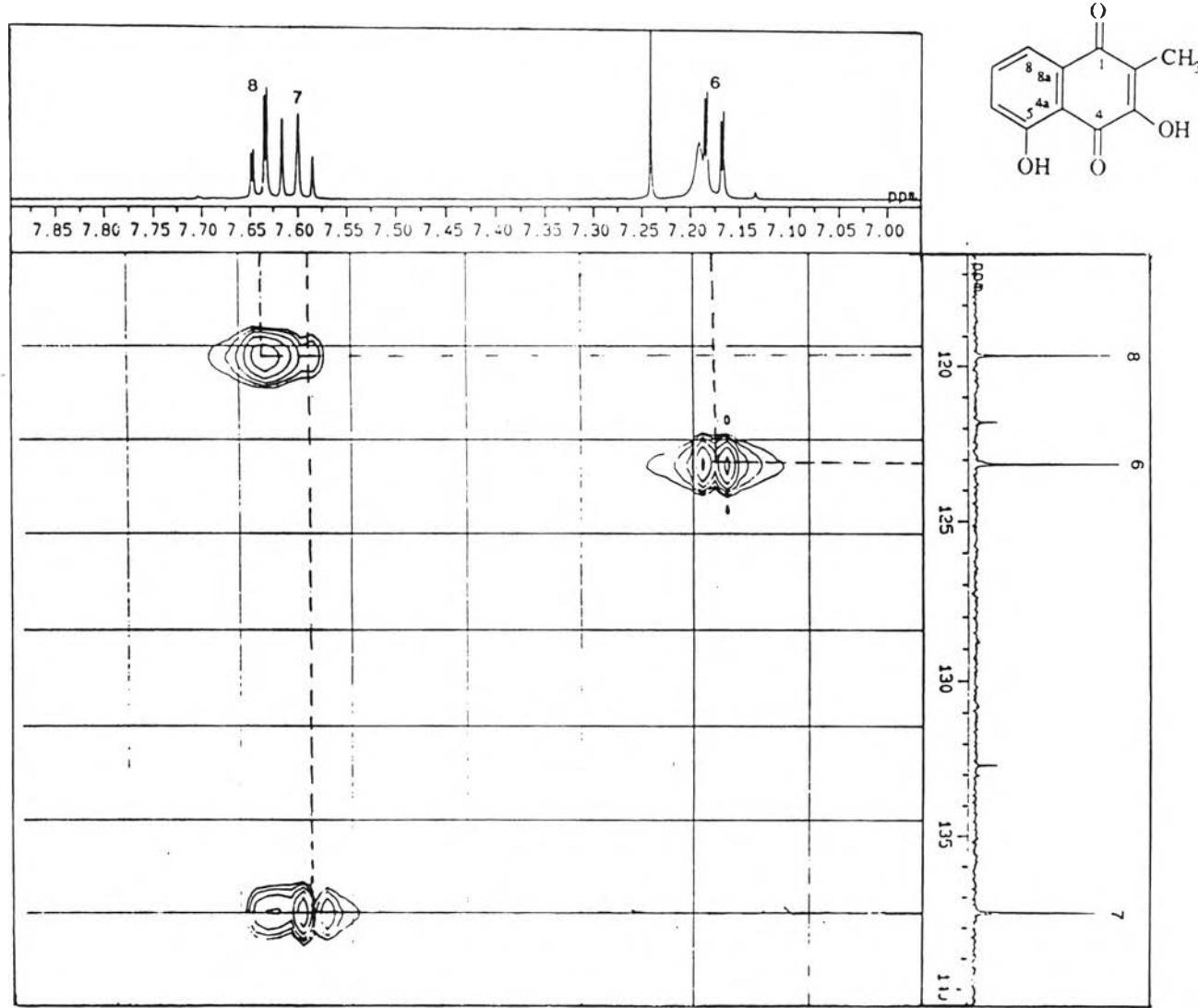


Figure 39 Expansion of the HMQC spectrum of compound 19 (in CDCl₃) :

δ_H 1.50-2.60 ; δ_C 0.00-18.00 ppm



δ_{H} 7.00-7.85 ; δ_{C} 117.00-140.00 ppm

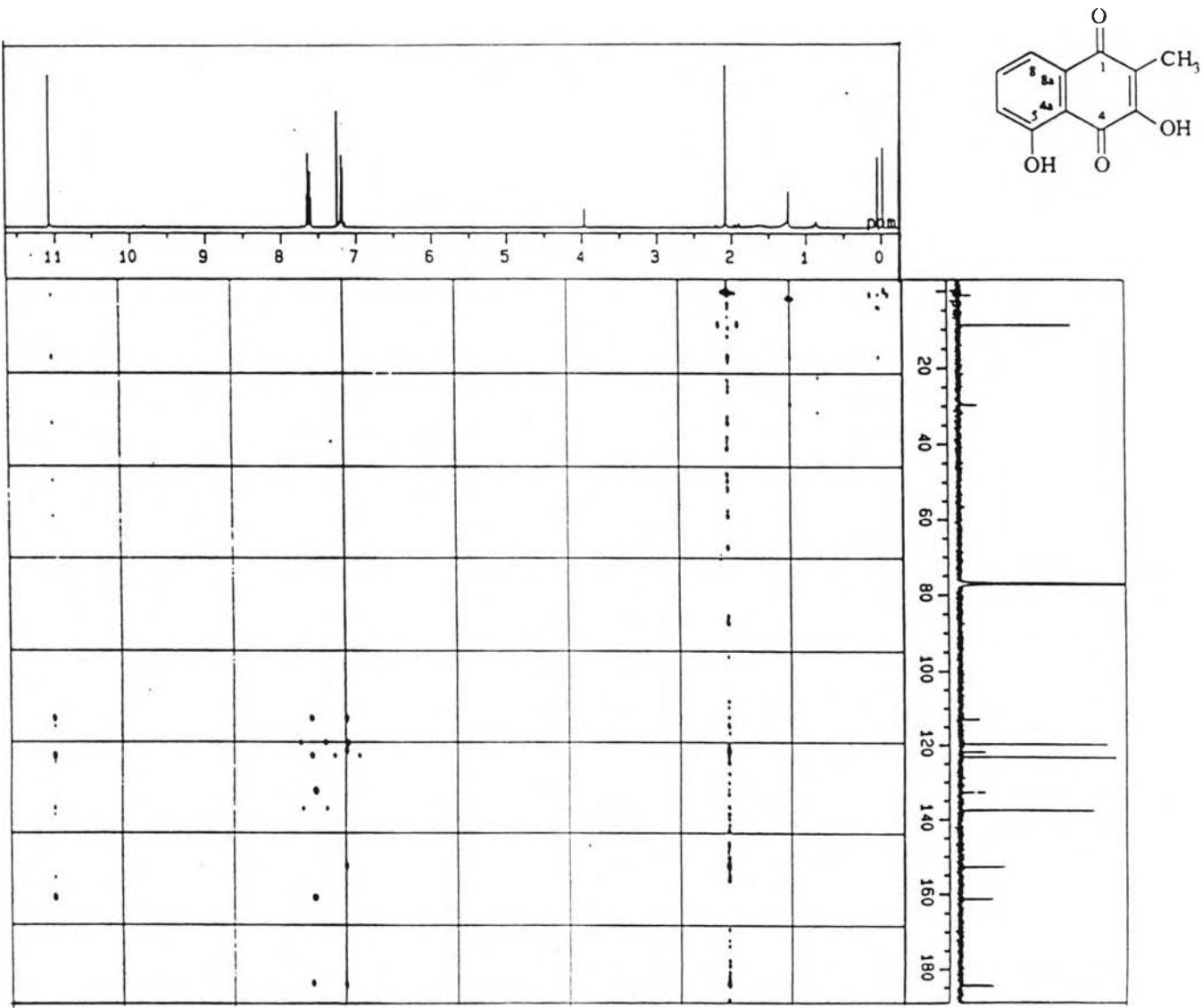


Figure 41 The HMBC spectrum of compound 19 (in CDCl_3)

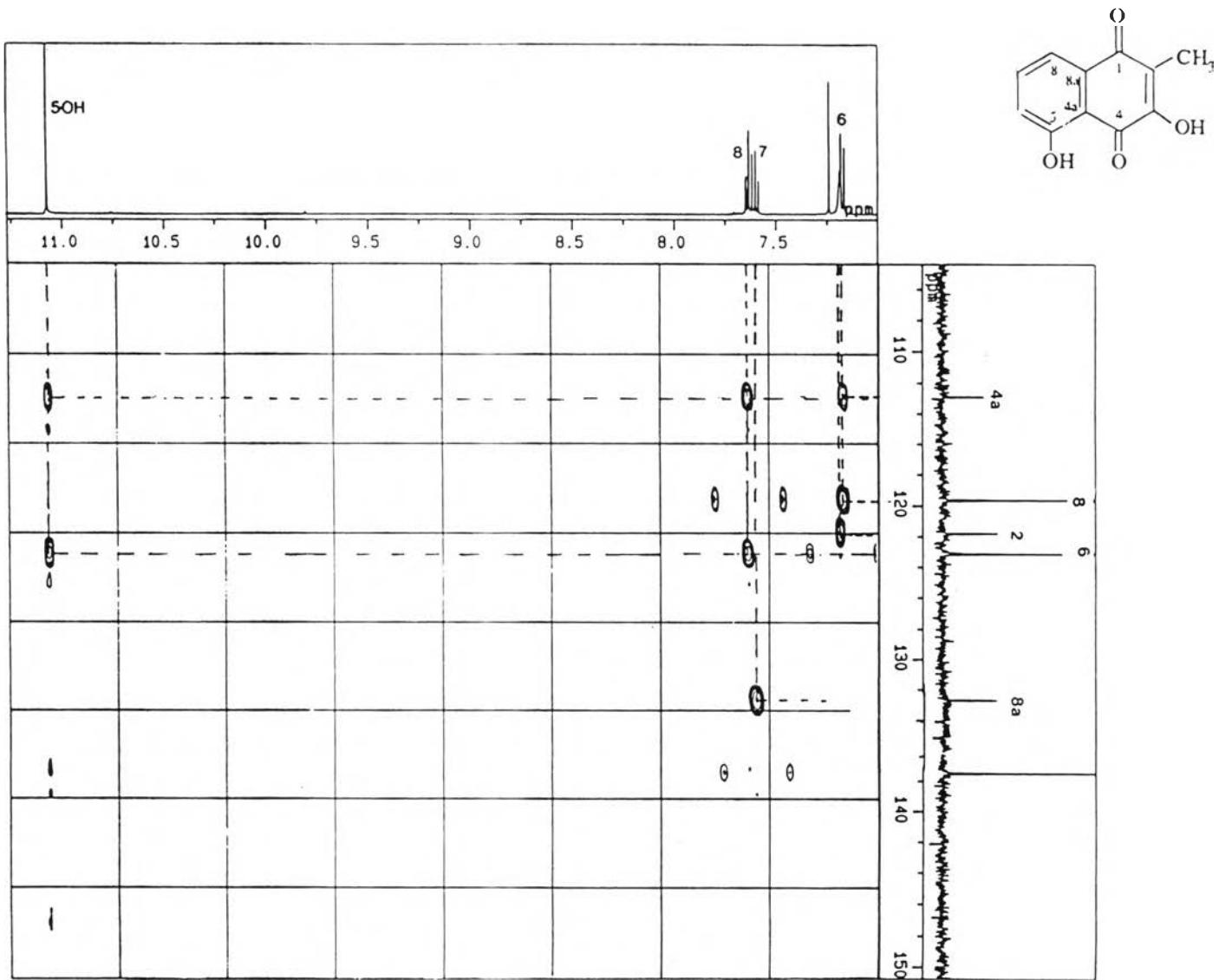


Figure 42 Expansion of the HMBC spectrum of compound 19 (in CDCl_3) :

δ_{H} 7.00-11.50 ; δ_{C} 108.00-150.00 ppm

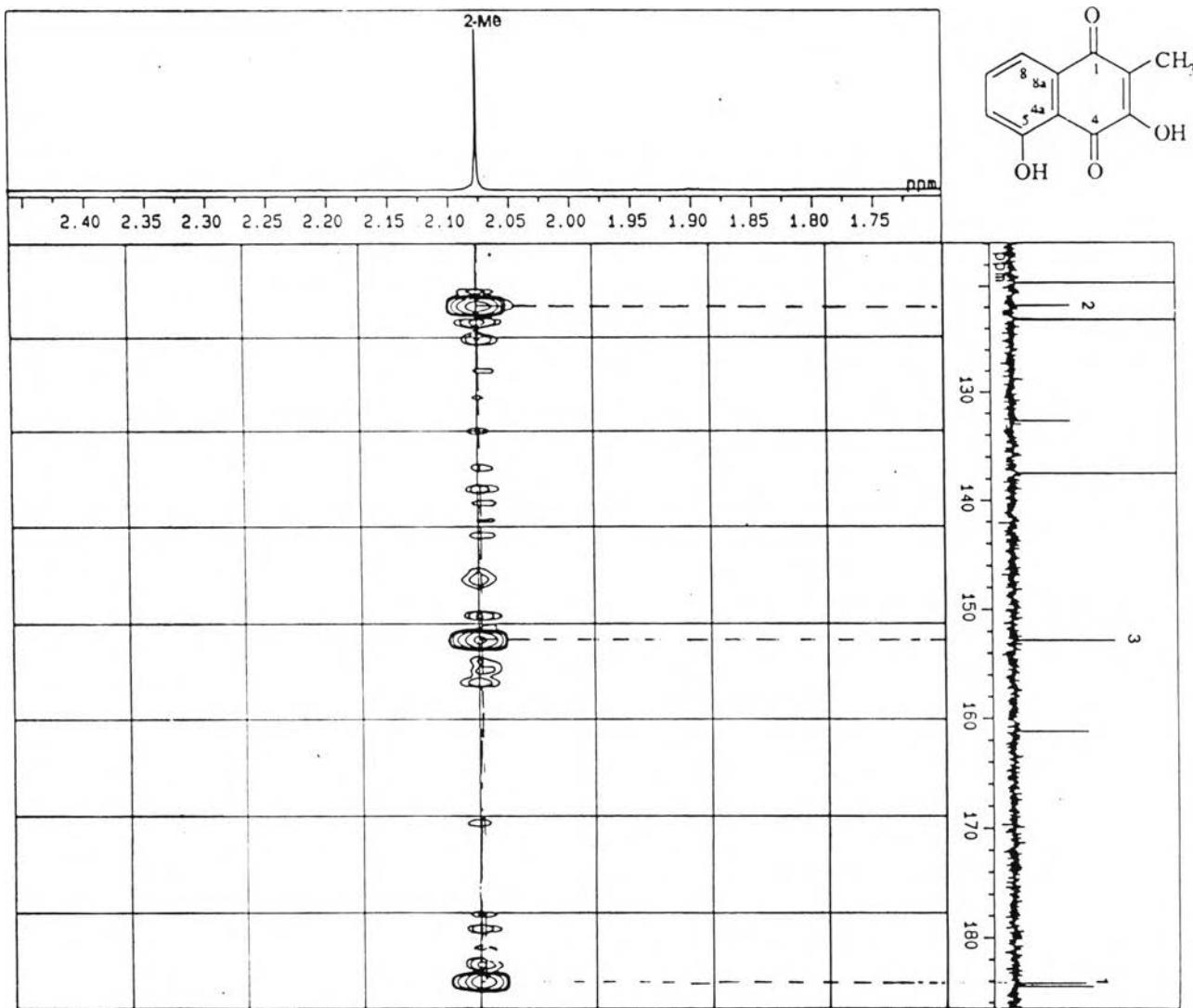


Figure 43 Expansion of the HMBC spectrum of compound 19 (in CDCl_3) :
 δ_{H} 1.75-2.41 ; δ_{C} 124.00-183.00 ppm

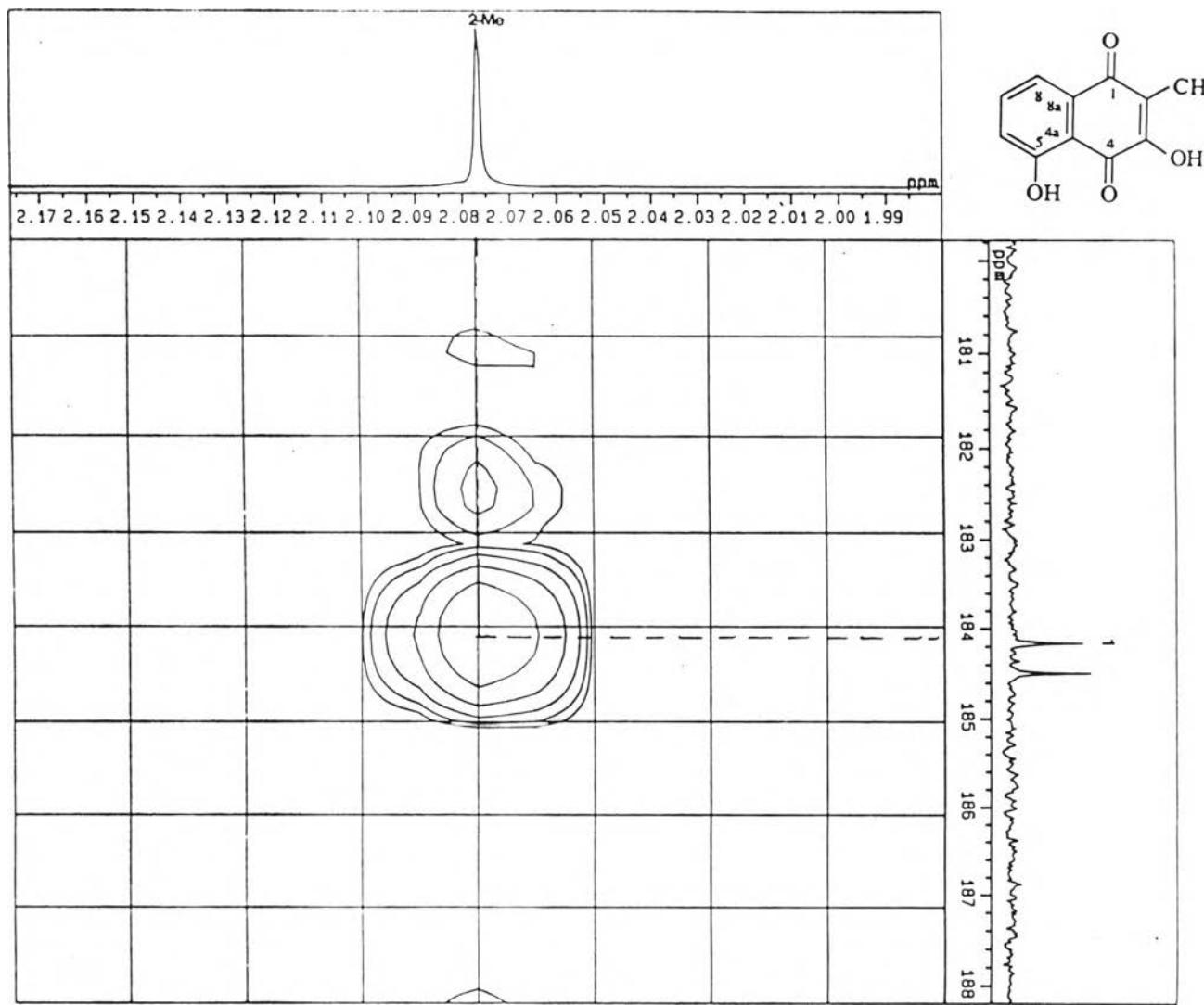


Figure 44 Expansion of the HMBC spectrum of compound 19 (in CDCl_3) :

δ_{H} 1.99-2.17 ; δ_{C} 180.00-188.00 ppm

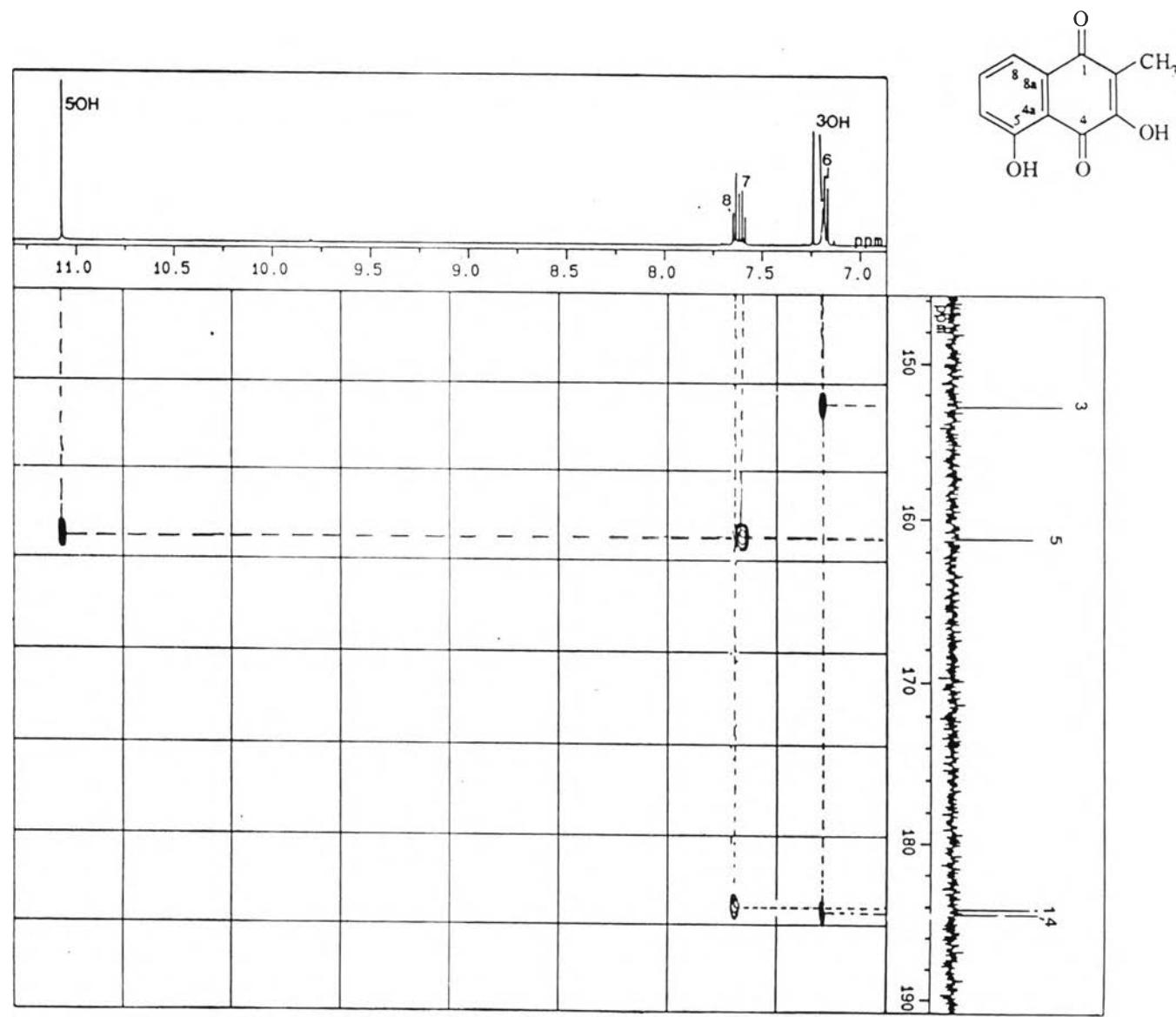


Figure 45 Expansion of the HMBC spectrum of compound 19 :

δ_H 7.00-11.50 ; δ_C 146.00-190.00 ppm

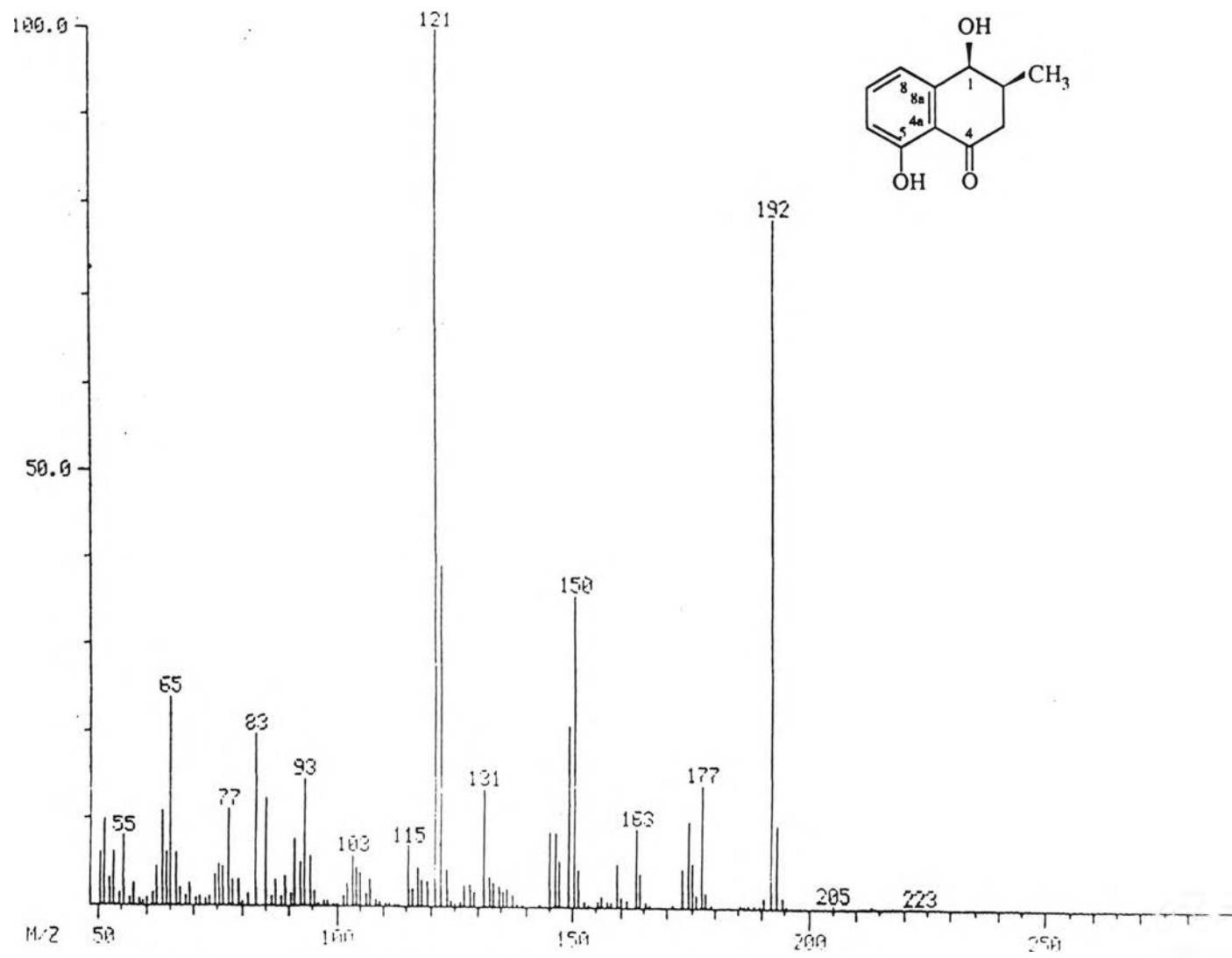


Figure 46 The EI mass spectrum of compound 102

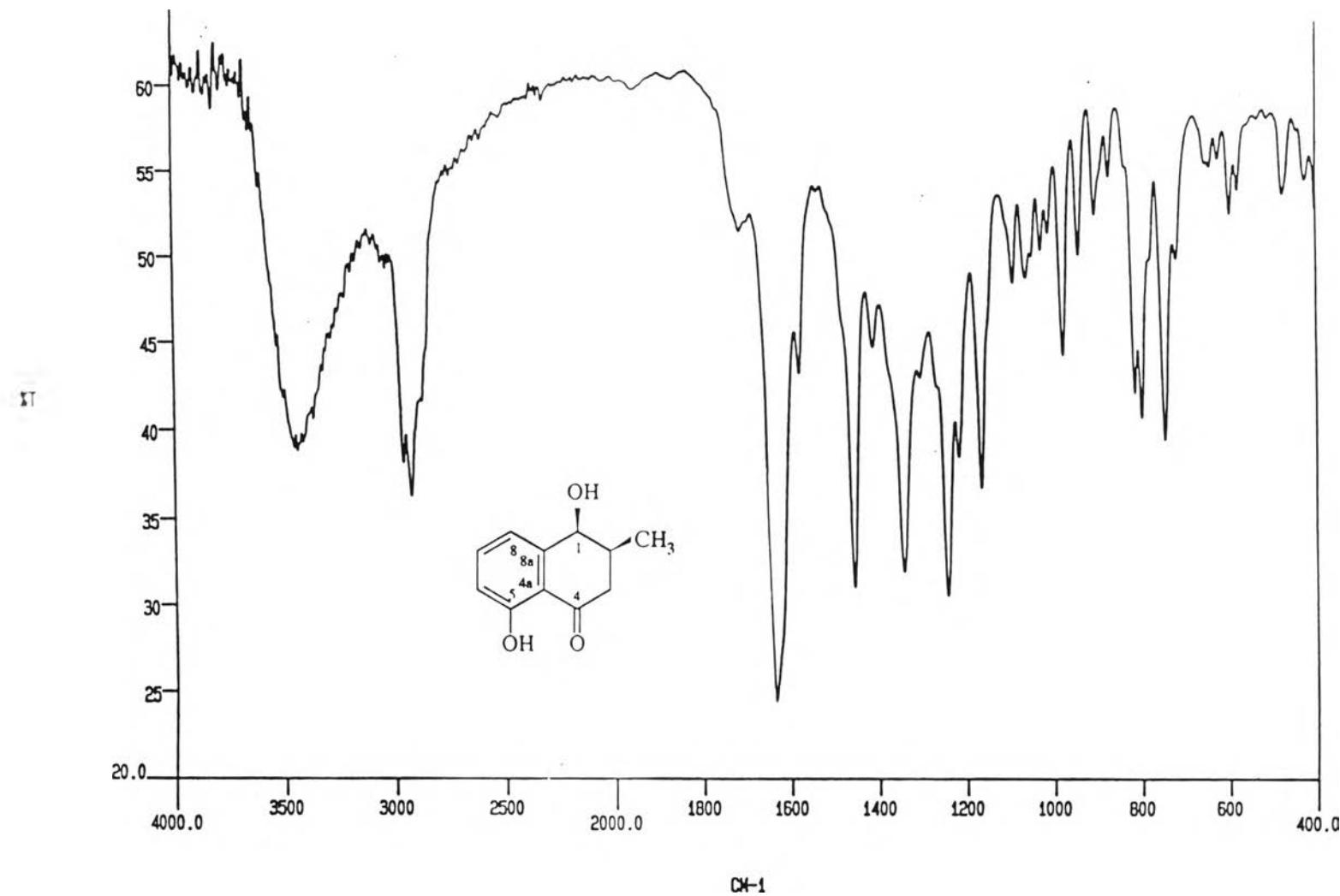


Figure 47 The IR spectrum of compound 102 (in KBr disc)

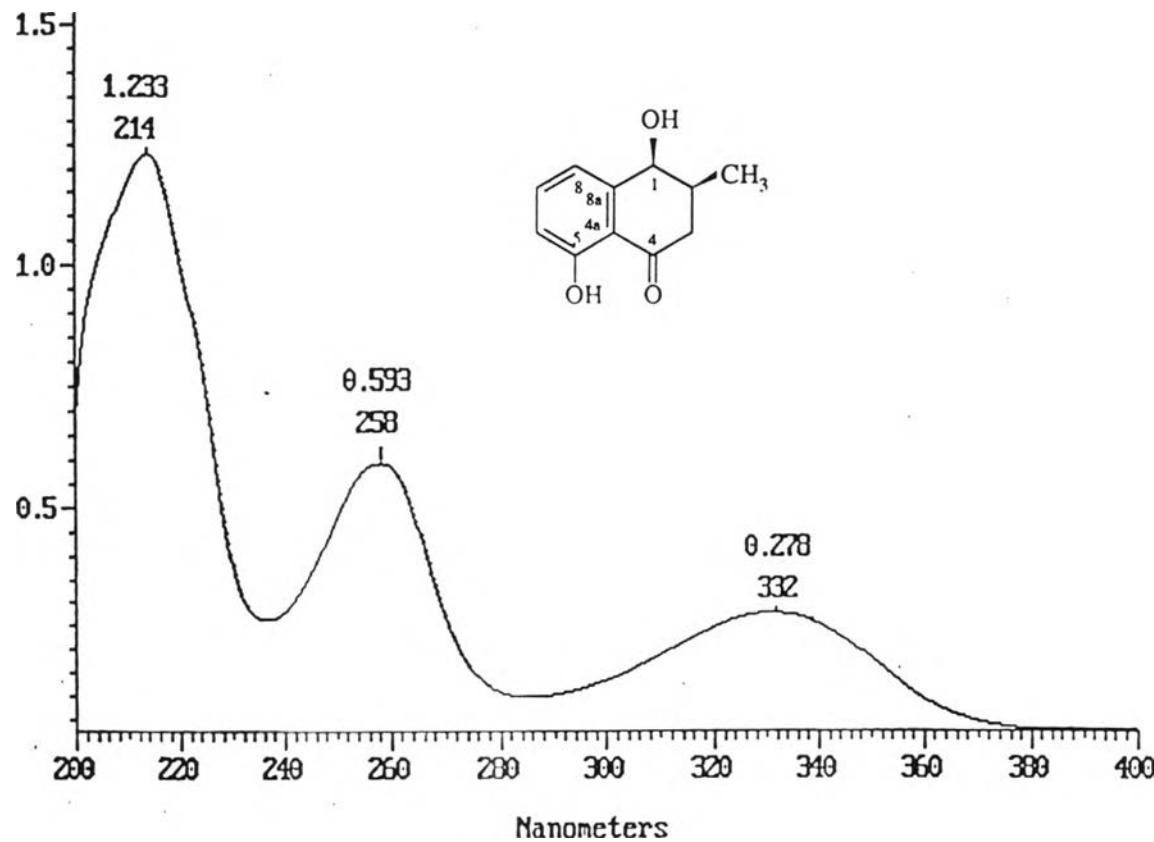


Figure 48 The UV spectrum of compound 102 (in MeOH)

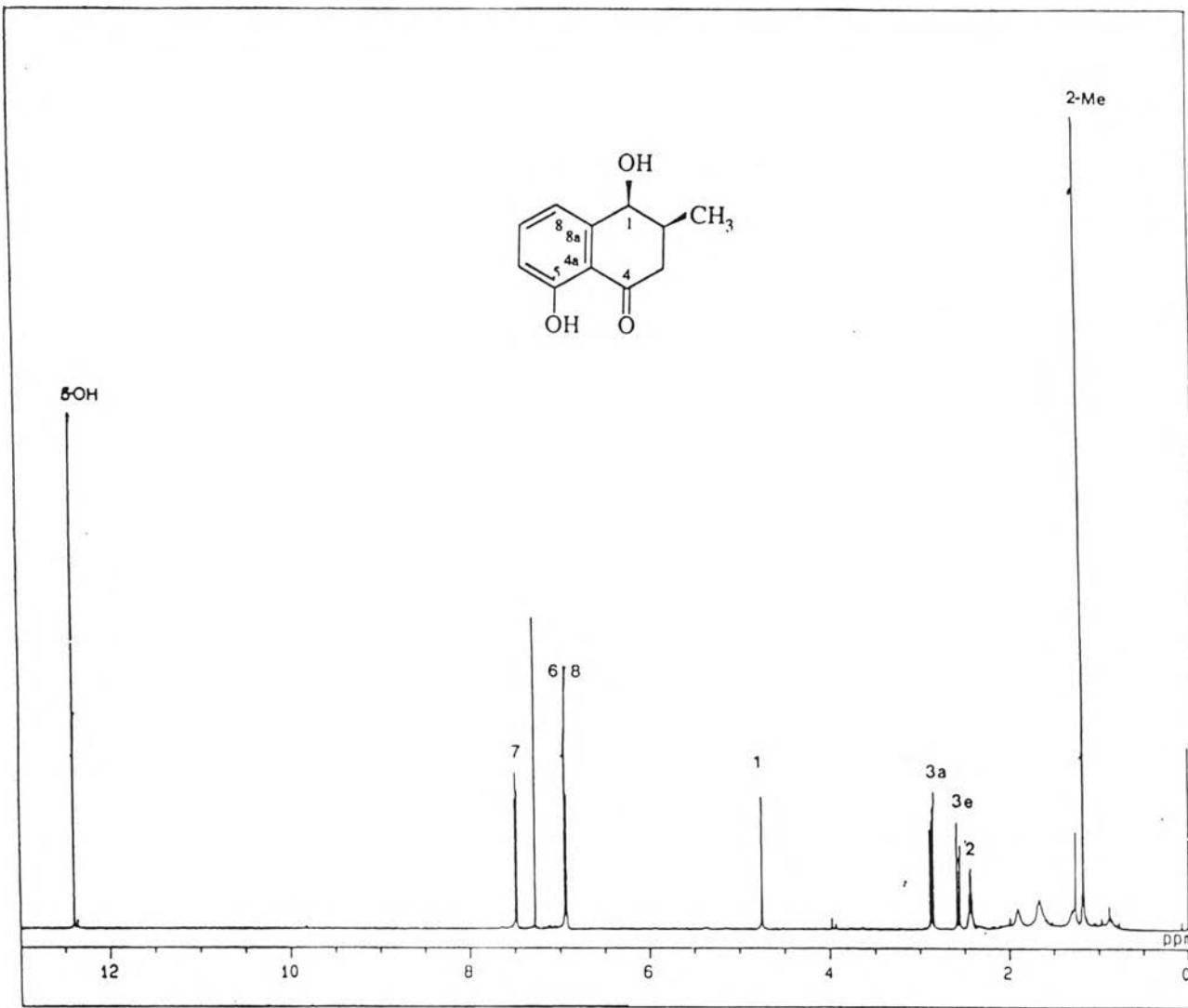


Figure 49 The ^1H NMR (500 MHz) spectrum of compound 102 (in CDCl_3)

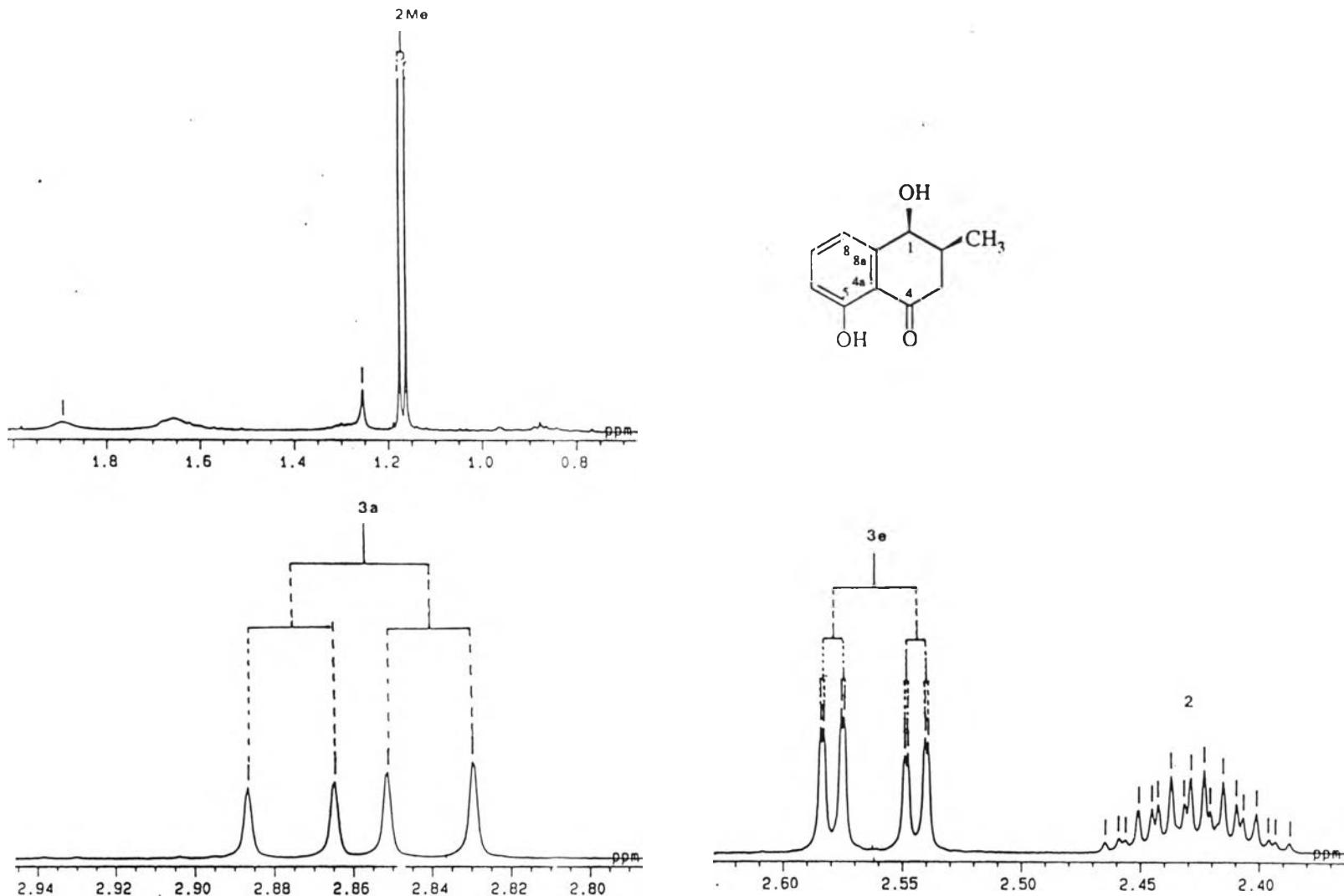


Figure 50 Expansion of the ^1H NMR (500 MHz) spectrum of compound 102

(in CDCl_3) : δ_{H} 0.80-2.94 ppm

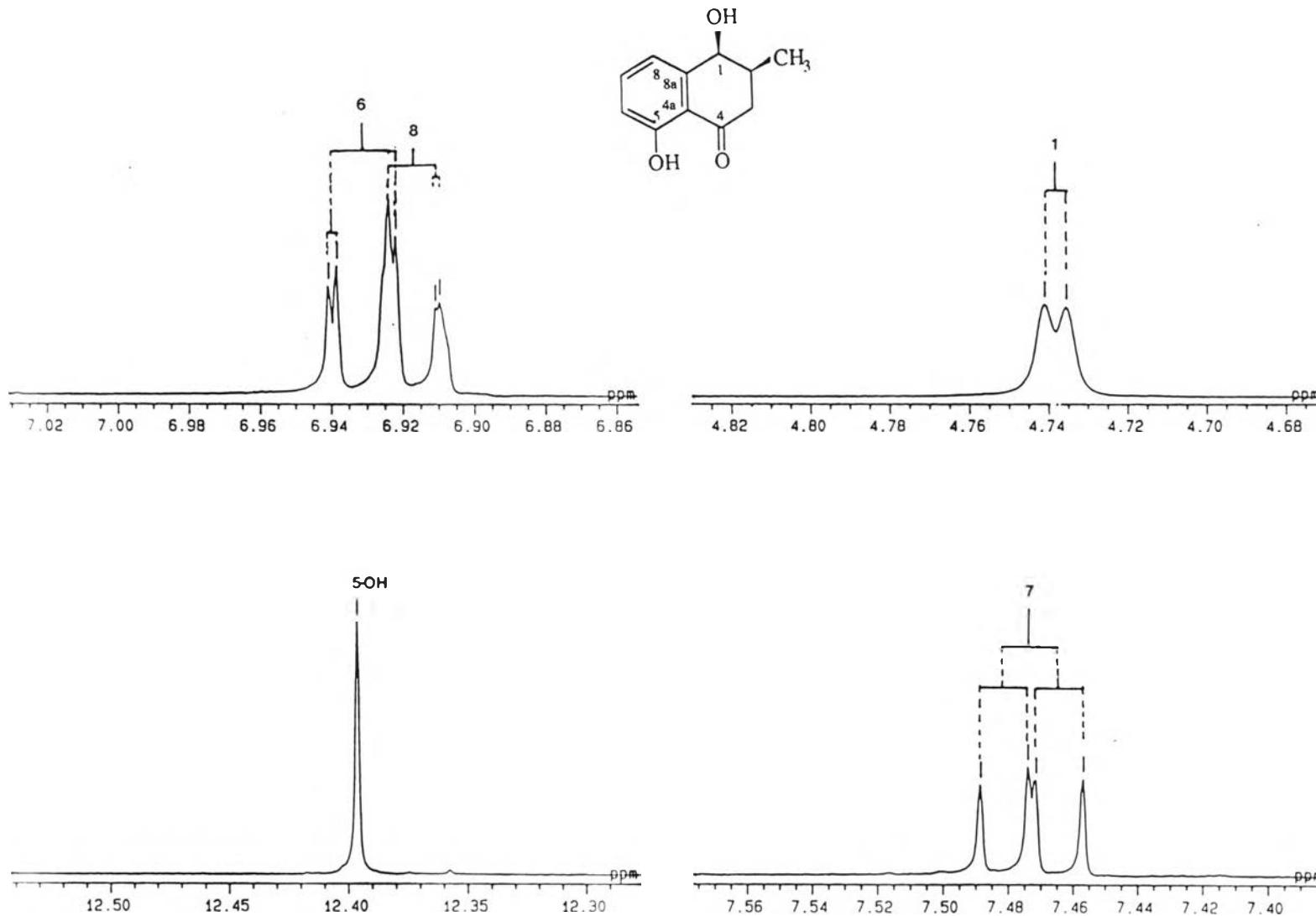


Figure 51 Expansion of the ^1H NMR (500 MHz) spectrum of compound 102

(in CDCl_3) : δ_{H} 4.68-12.50 ppm

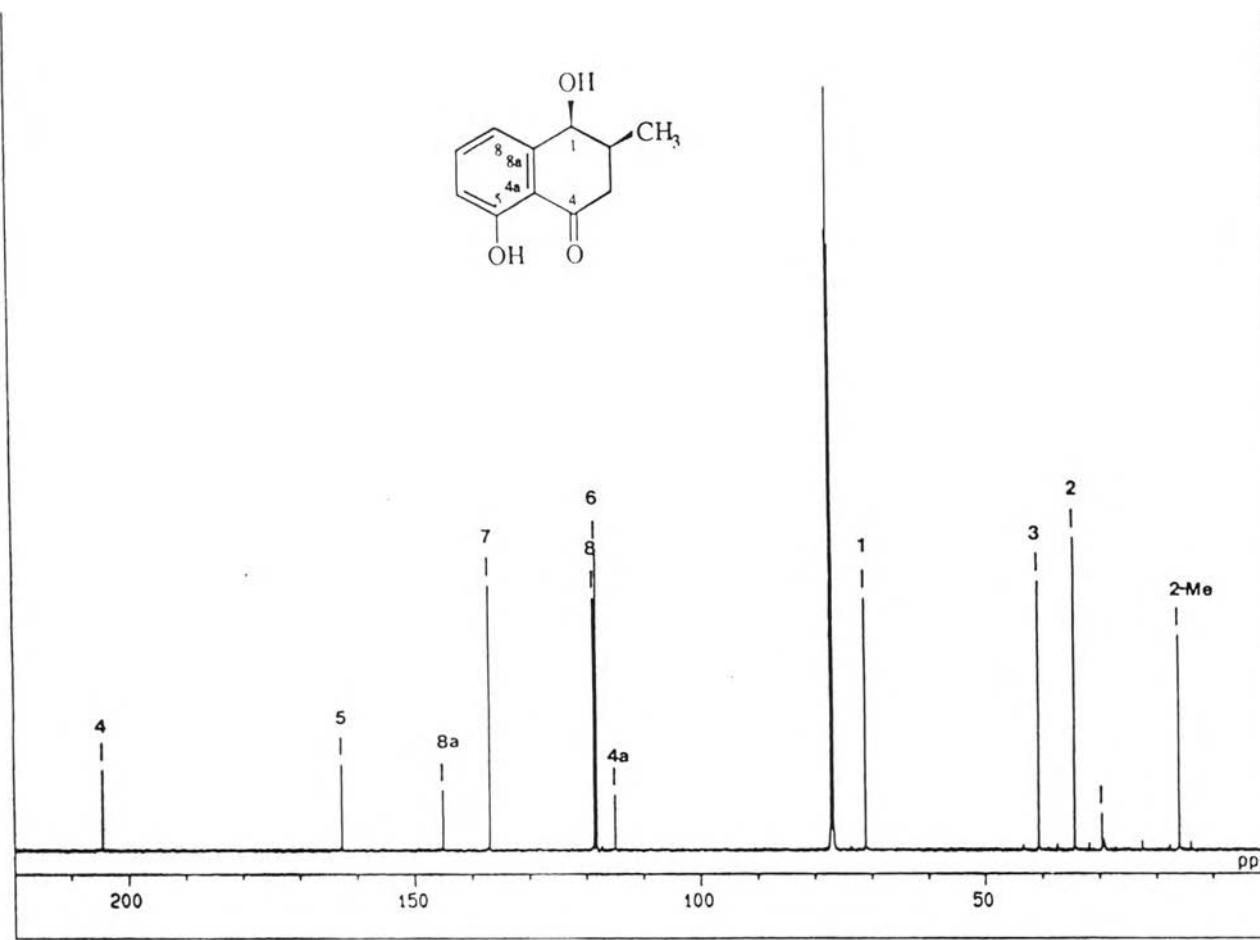


Figure 52 The ^{13}C NMR (125 MHz) spectrum of compound 102 (in CDCl_3)

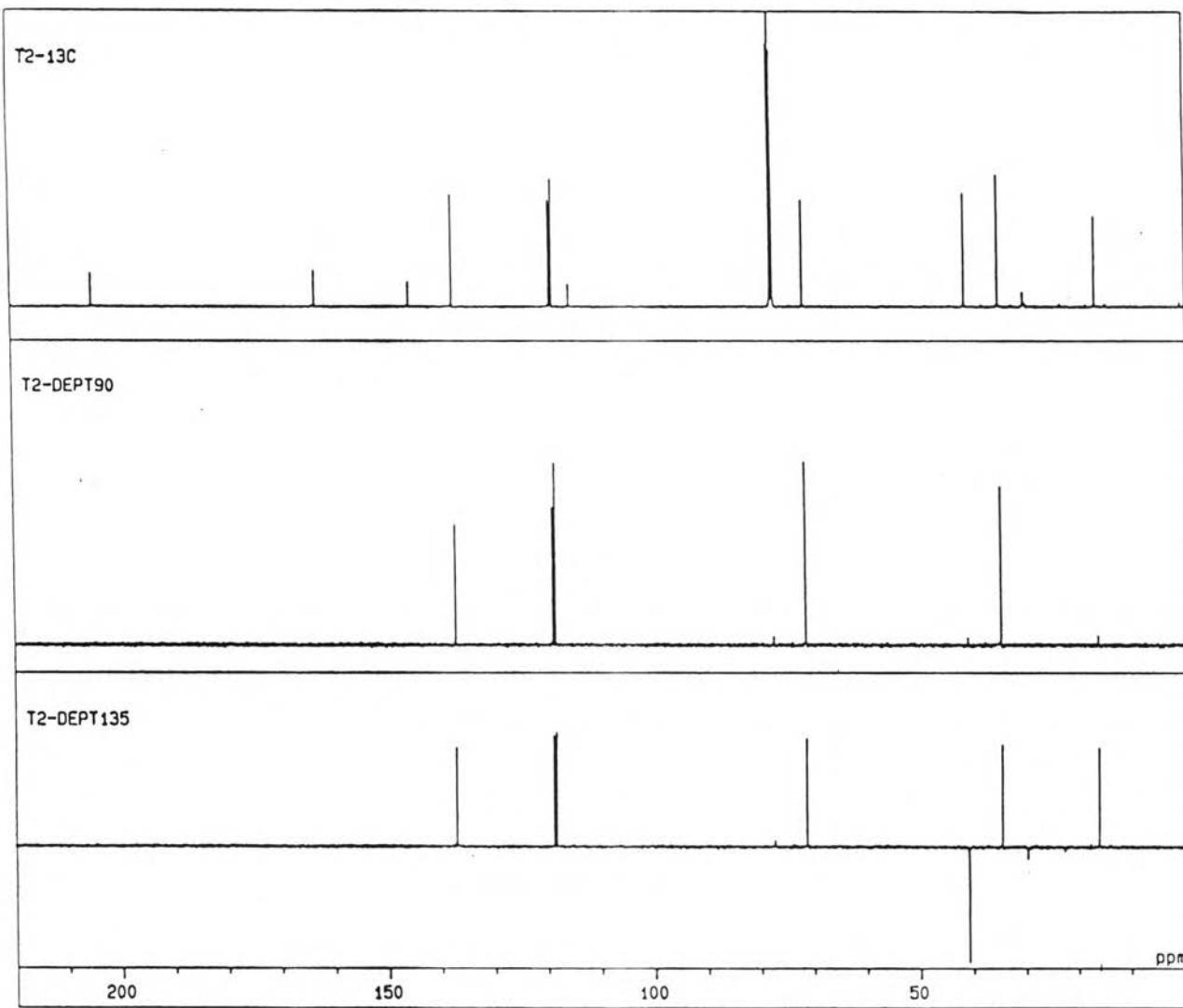
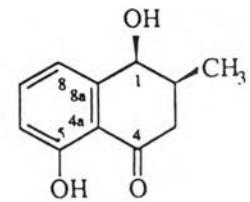


Figure 53 The DEPT (125 MHz) spectrum of compound 102 (in CDCl₃)

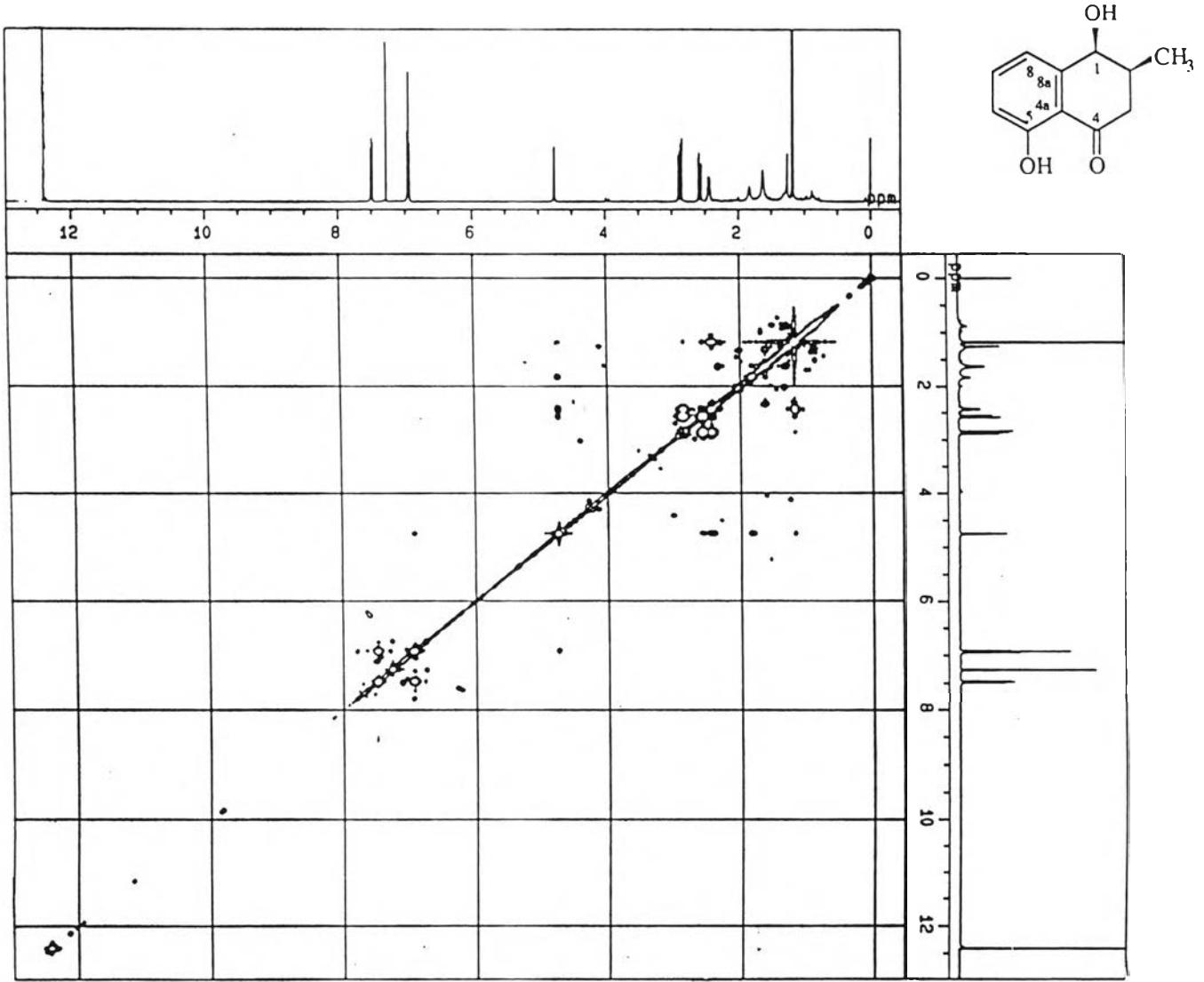


Figure 54 The ^1H - ^1H COSY (500 MHz) spectrum of compound 102 (in CDCl_3)

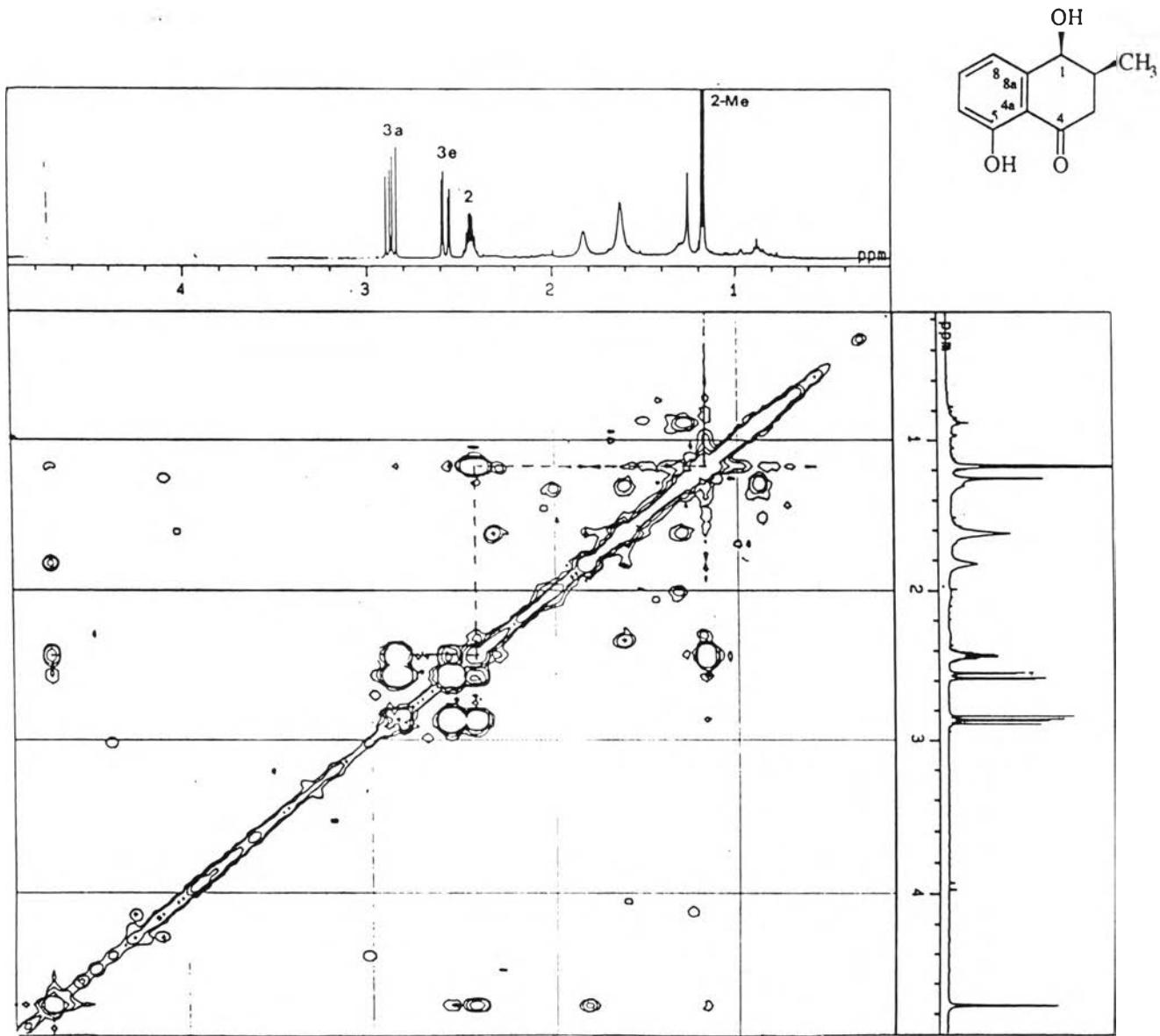


Figure 55 Expansion of the ¹H-¹H COSY (500 MHz) spectrum of compound 102
 (in CDCl₃) : δ_H 0.40-4.80 ppm

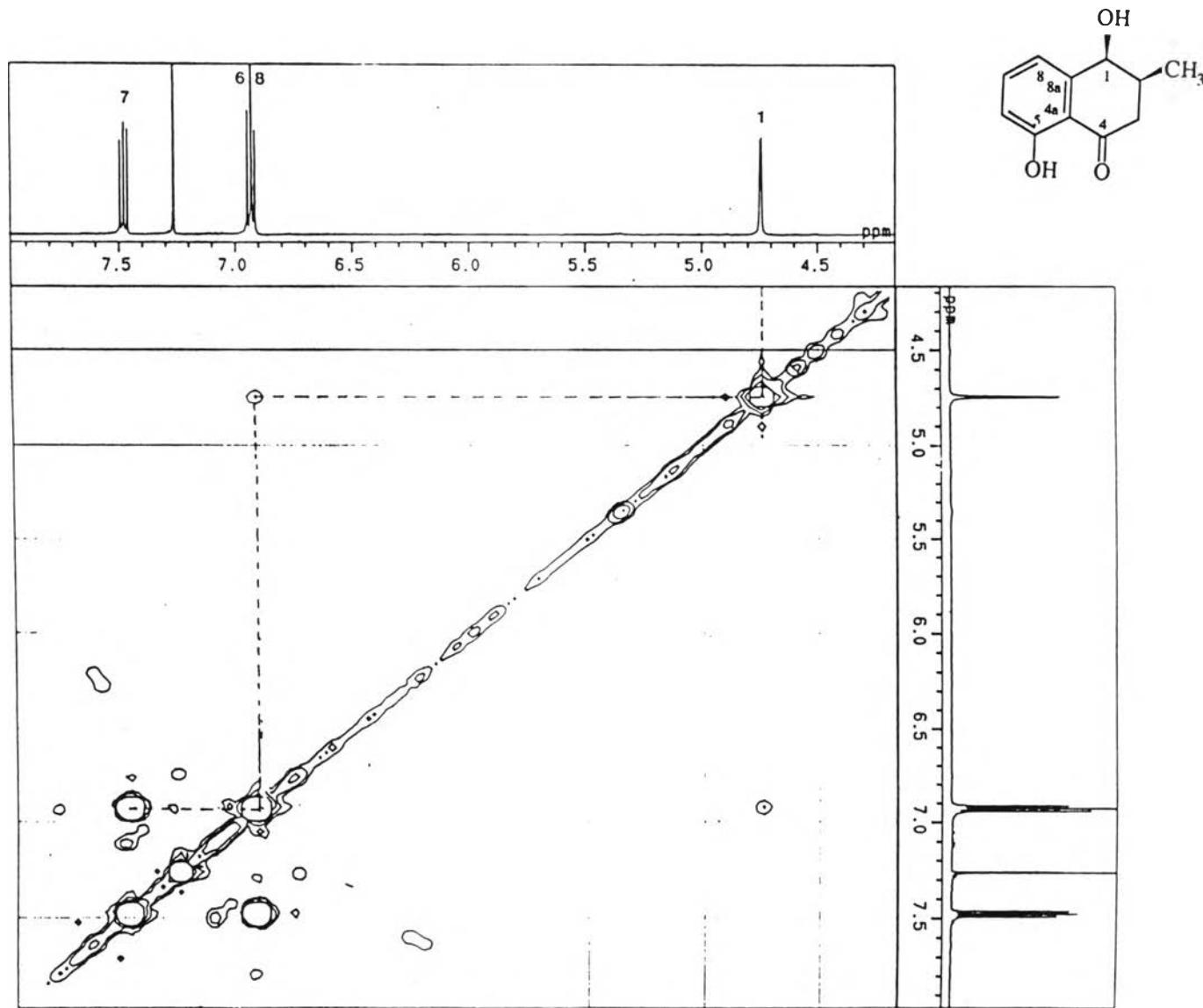


Figure 56 Expansion of the ^1H - ^1H COSY (500 MHz) spectrum of compound 102
(in CDCl_3) : δ_{H} 4.30-7.90 ppm

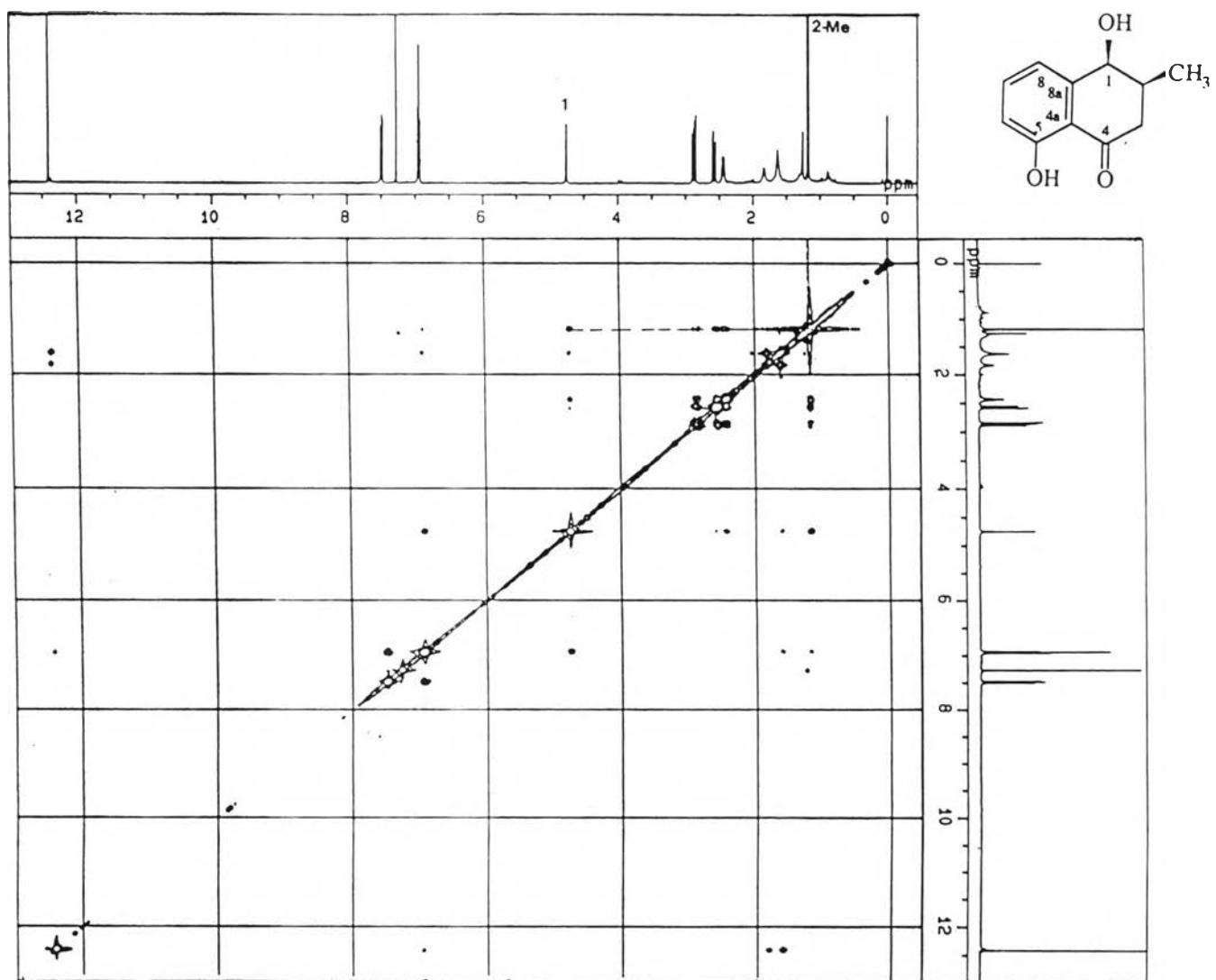


Figure 57 The NOESY (500 MHz) spectrum of compound 102 (in CDCl₃)

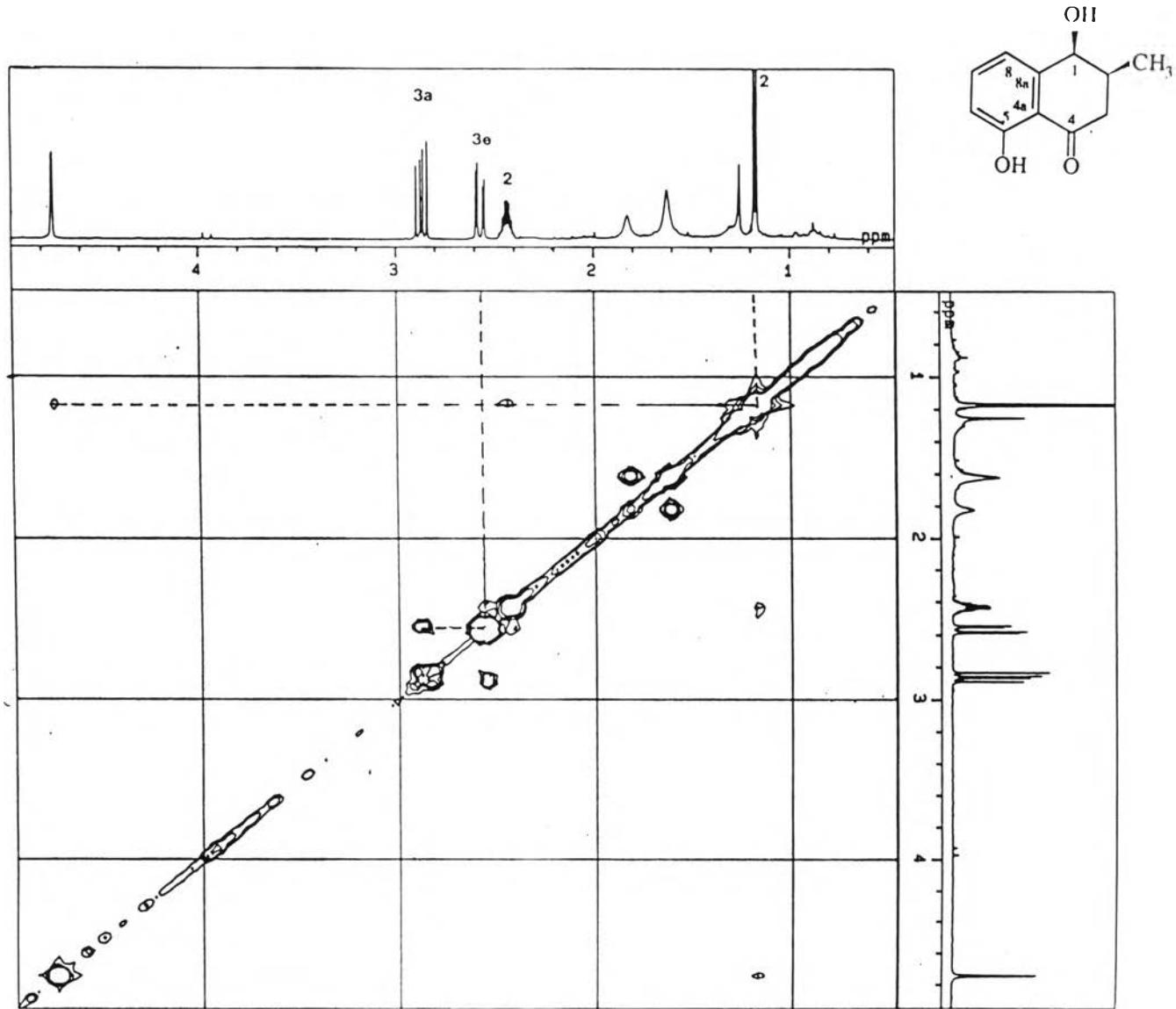


Figure 58 Expansion of the NOESY (500 MHz) spectrum of compound 102
 (in CDCl_3) : δ_{H} 0.60-4.80 ppm

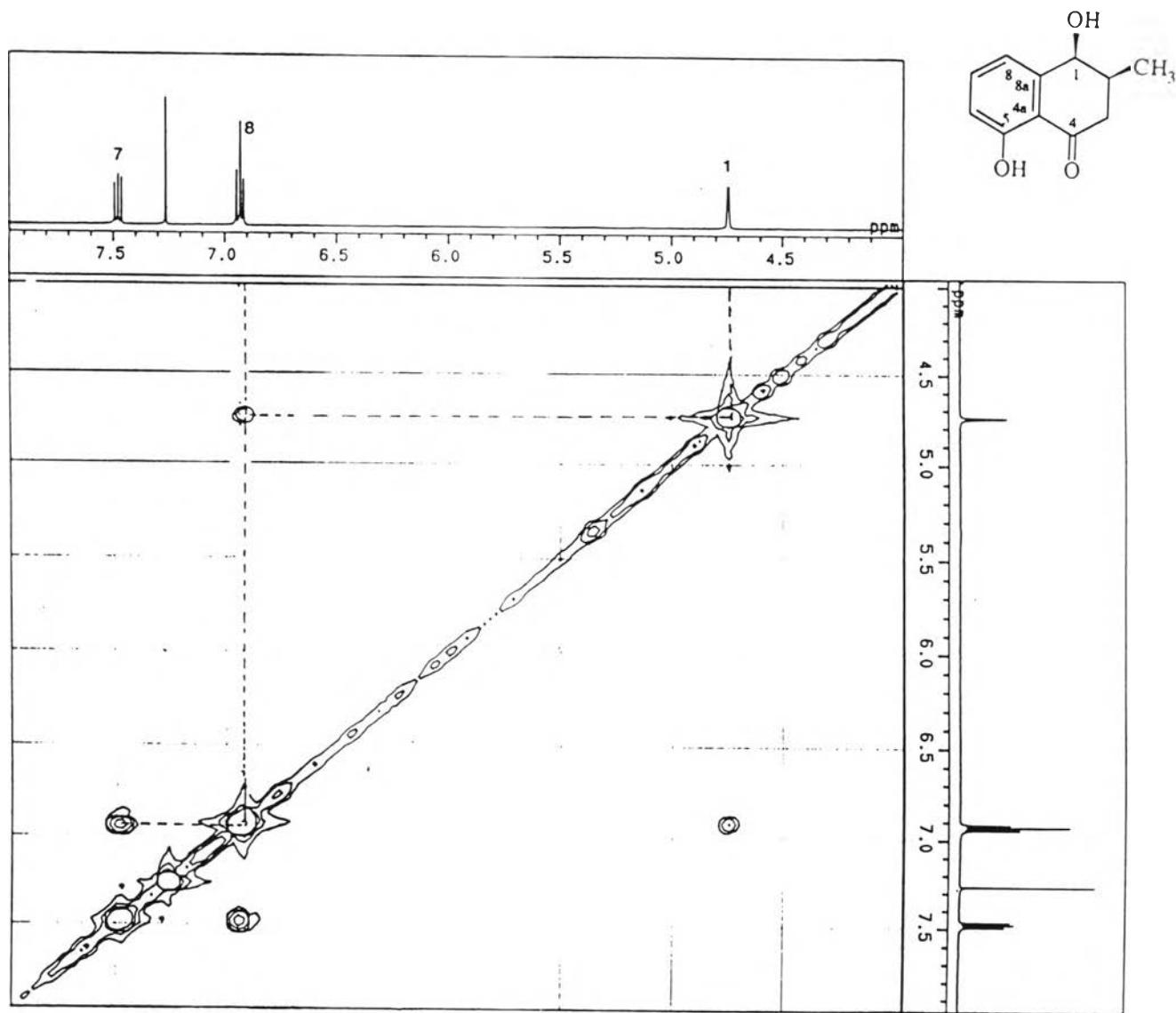


Figure 59 Expansion of the NOESY (500 MHz) spectrum of compound 102
 (in CDCl_3) : δ_{H} 4.10-7.80 ppm

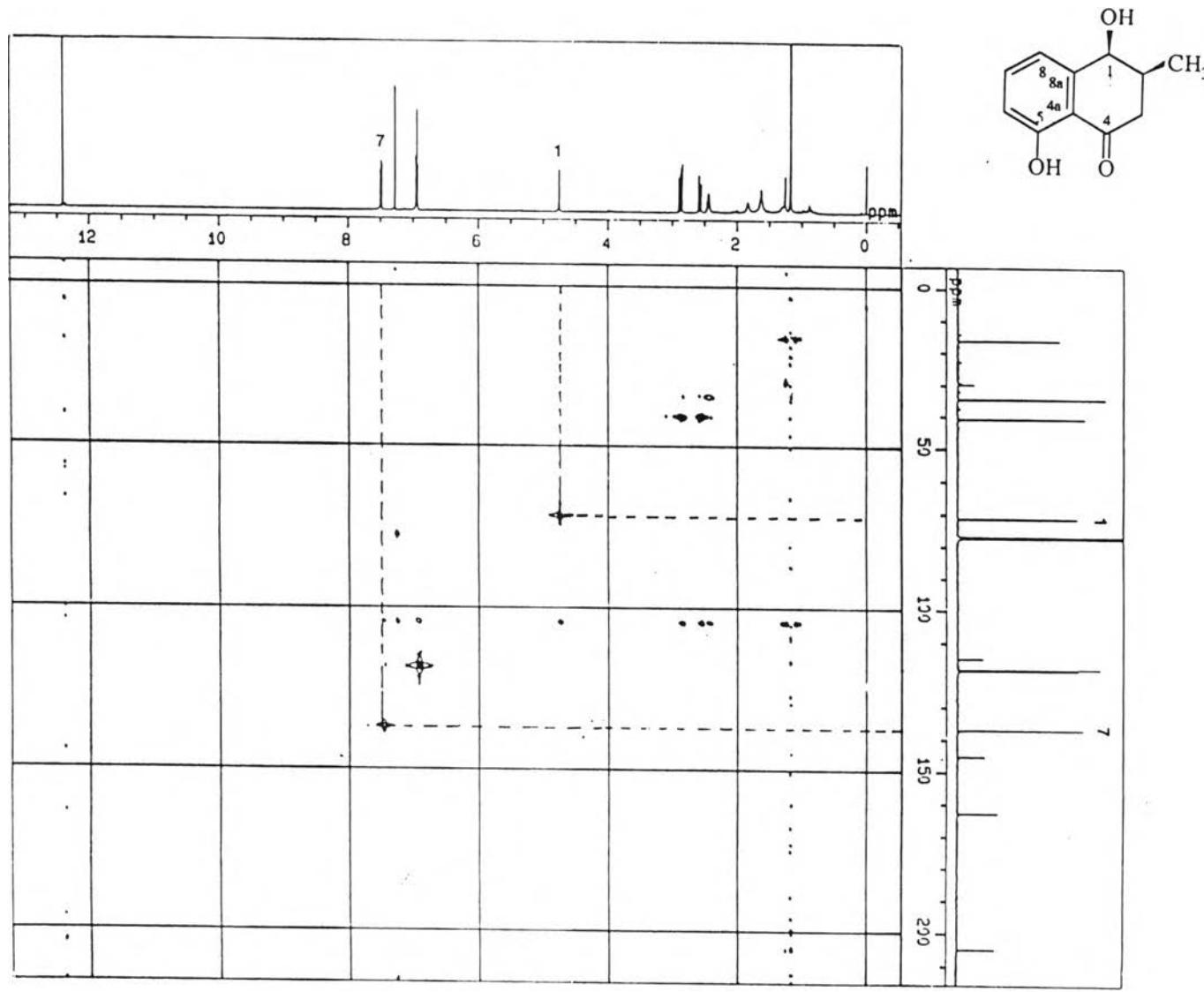


Figure 60 The HMQC spectrum of compound 102 (in CDCl_3)

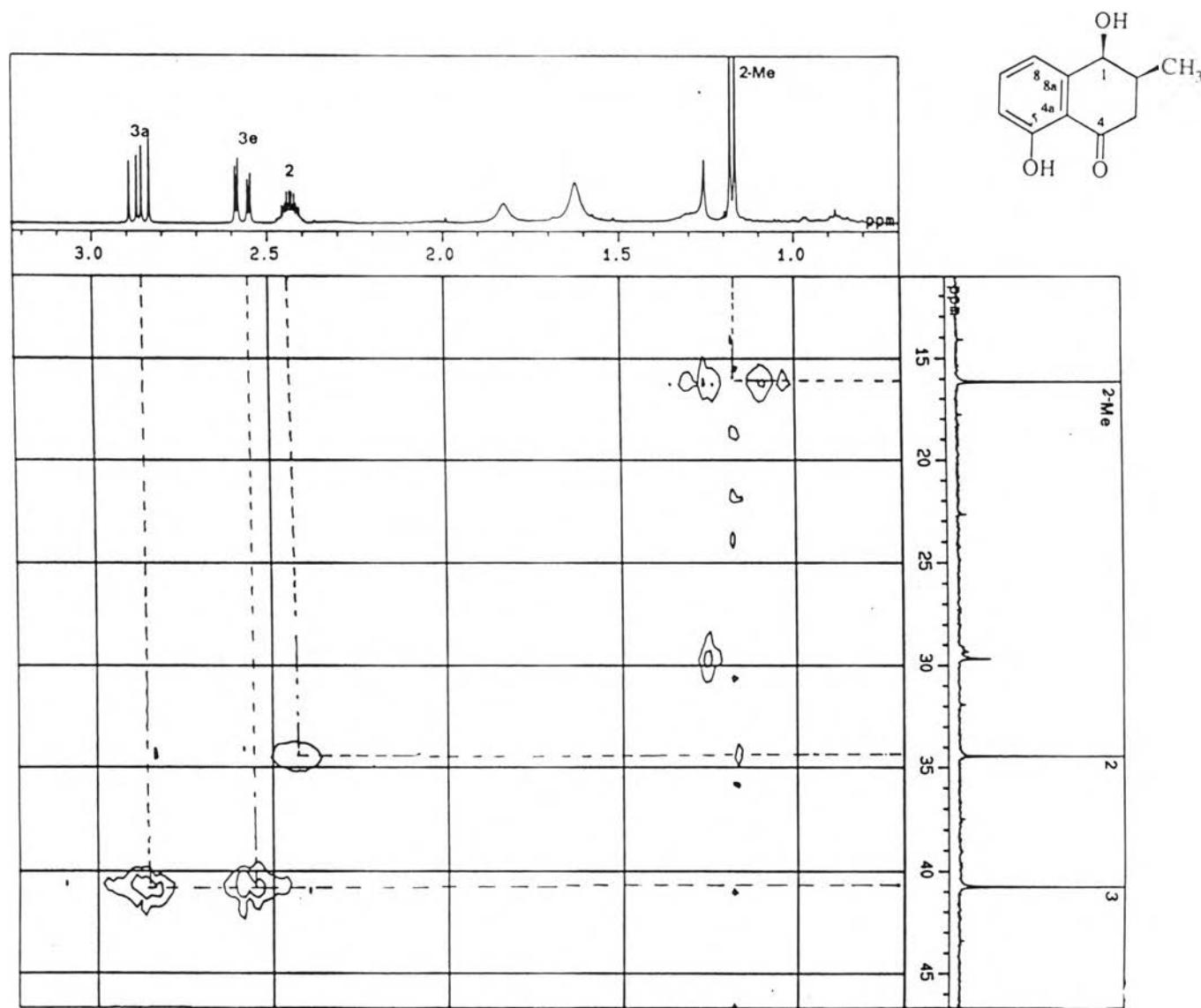


Figure 61 Expansion of the HMQC spectrum of compound 102 (in CDCl_3) :
 δ_{H} 0.60-3.20 ; δ_{C} 12.00-46.00 ppm

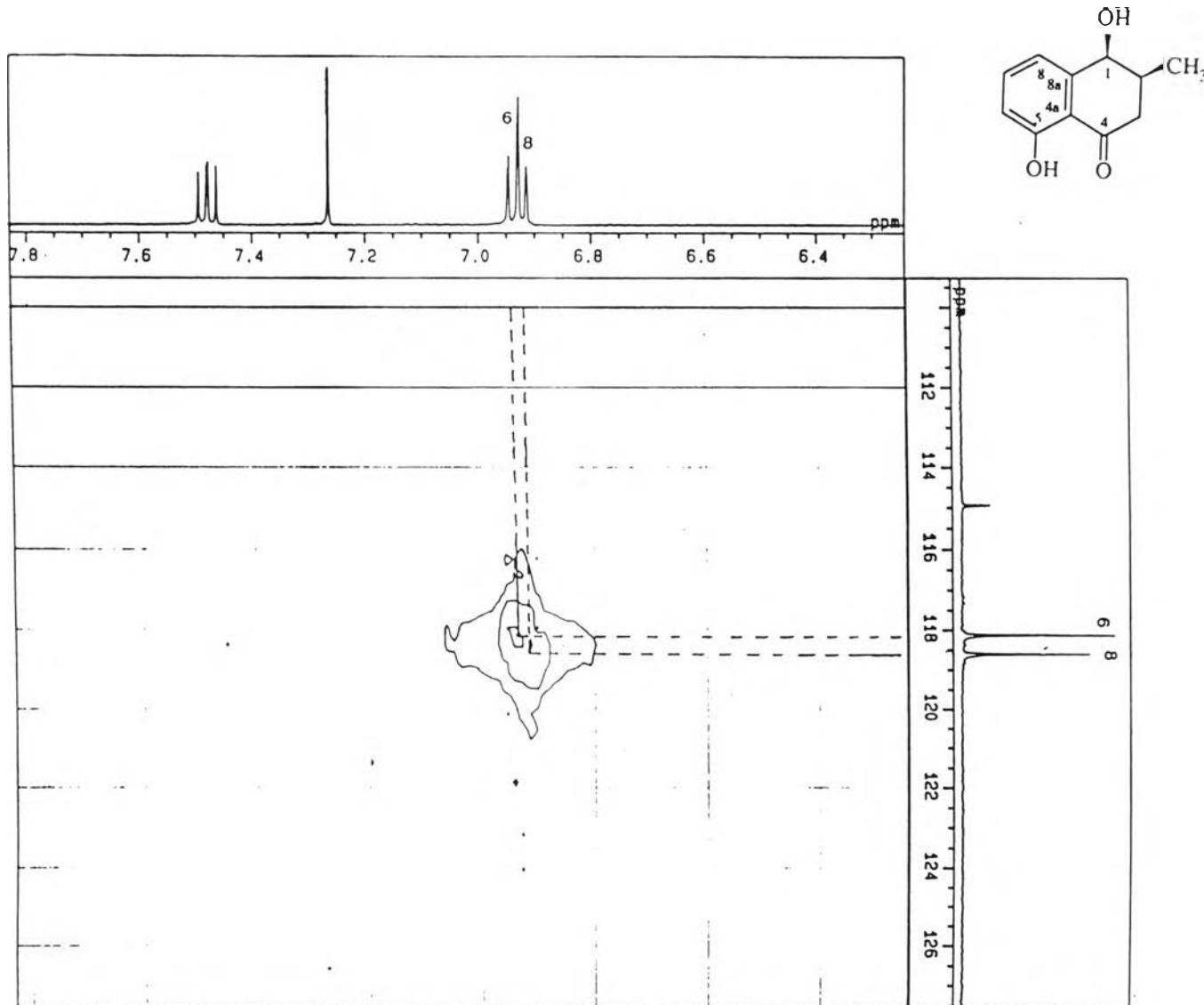


Figure 62 Expansion of the HMQC spectrum of compound 102 (in CDCl_3) :

δ_{H} 6.40-7.80 ; δ_{C} 112.00-126.00 ppm

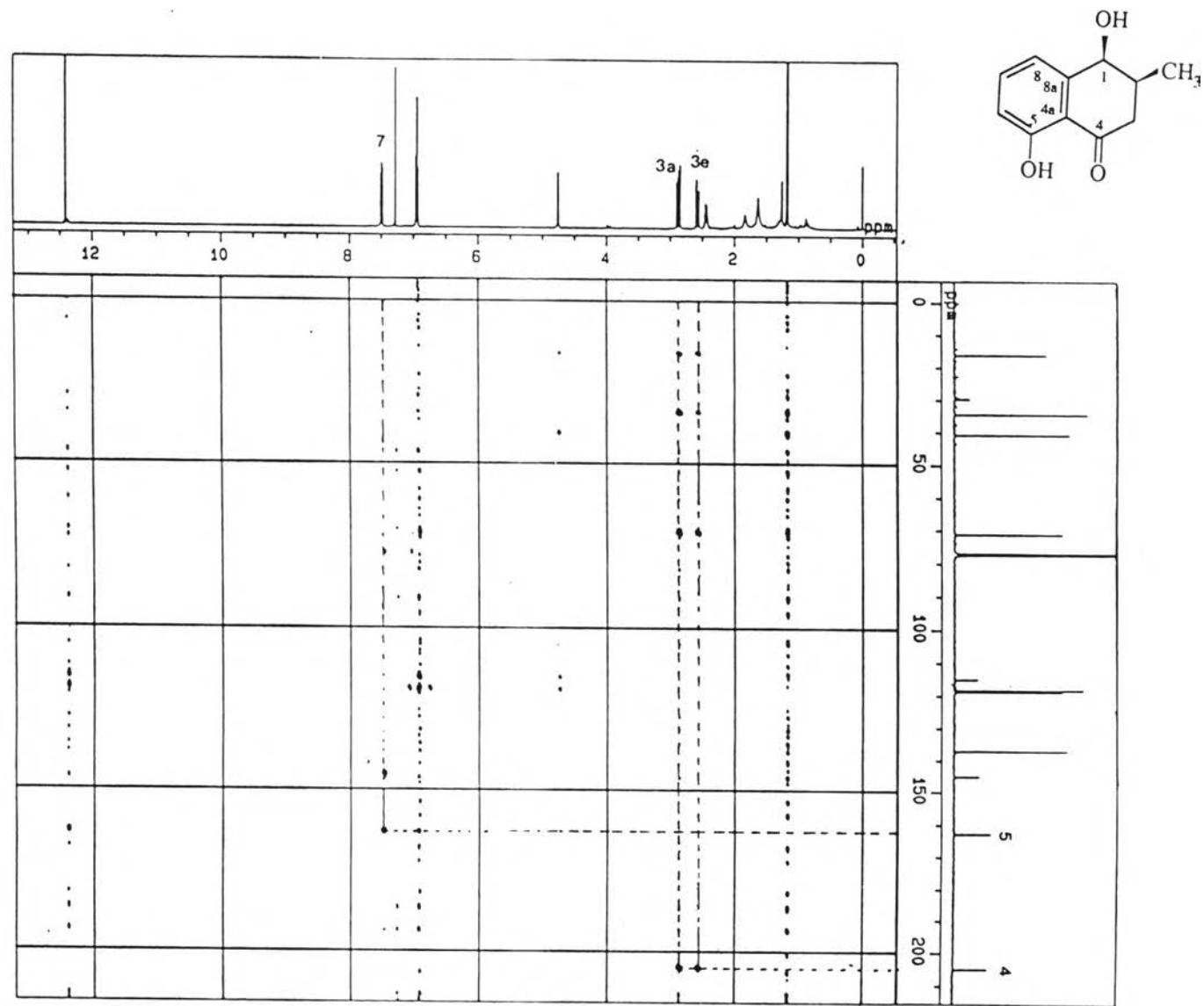


Figure 63 The HMBC spectrum of compound 102 (in CDCl₃)

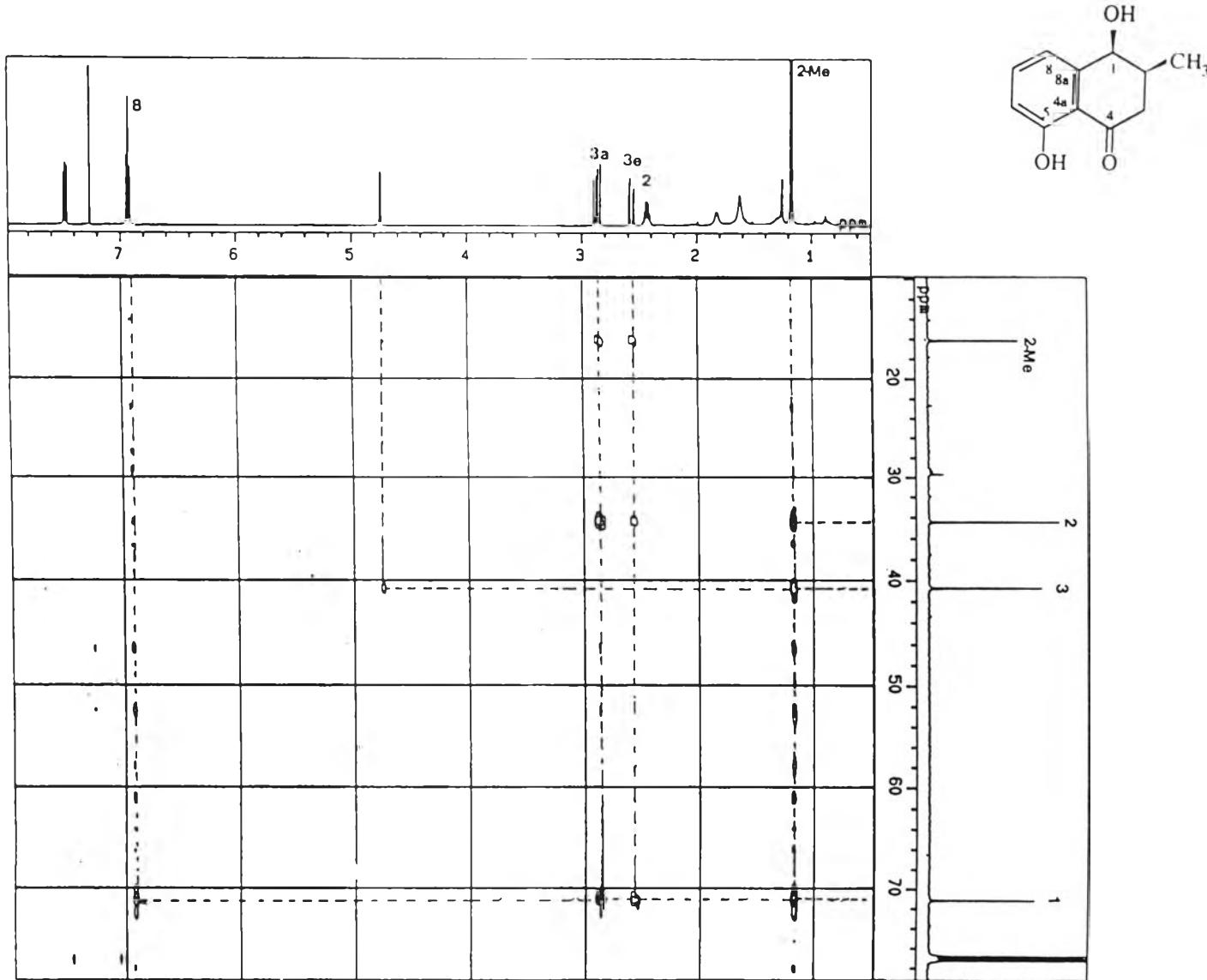


Figure 64 Expansion of the HMBC spectrum of compound **102** (in CDCl_3) :
 δ_{H} 0.60-7.80 ; δ_{C} 10.00-78.00 ppm

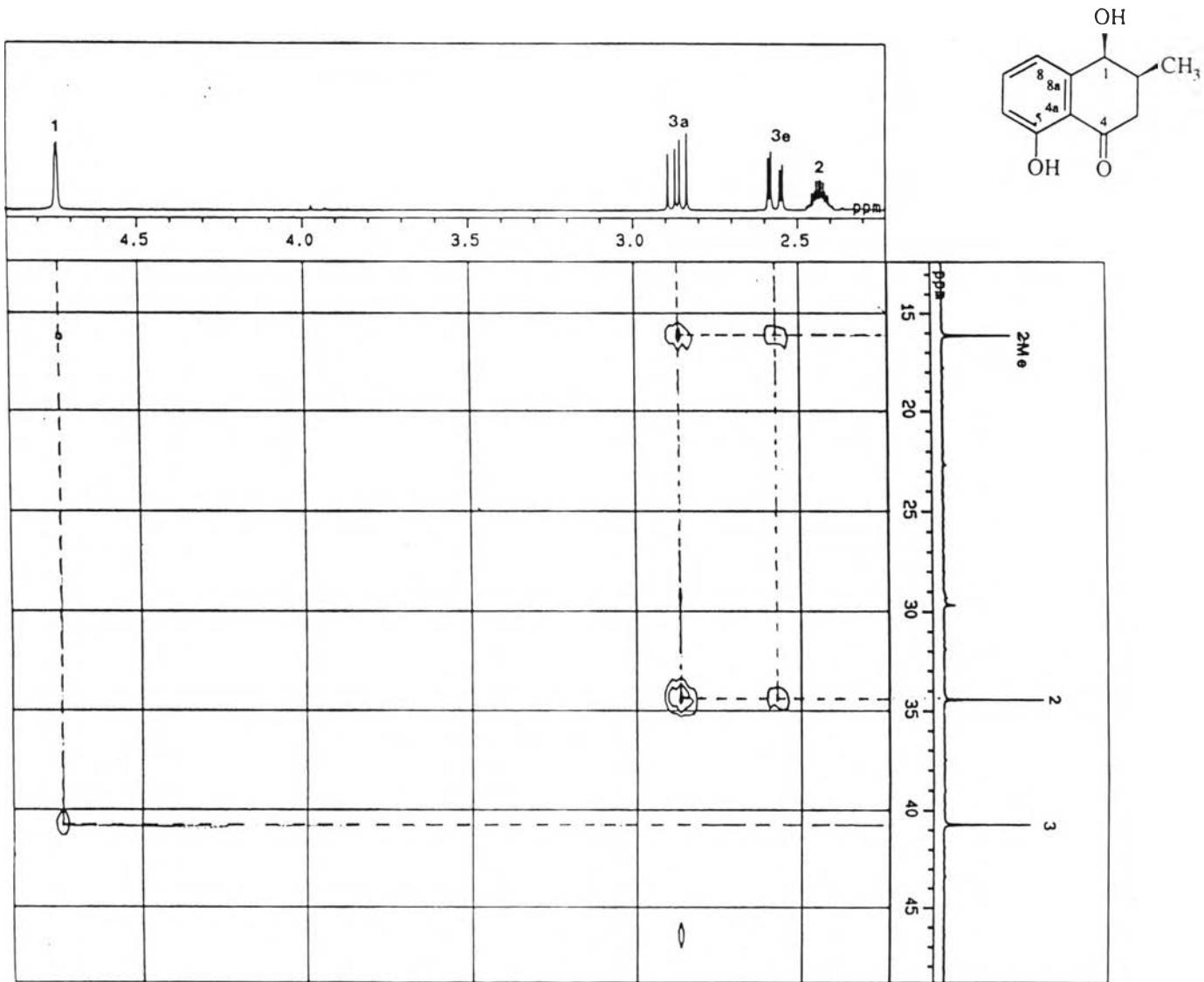


Figure 65 Expansion of the HMBC spectrum of compound 102 (in CDCl_3) :

δ_{H} 2.30-4.80 ; δ_{C} 13.00-48.00 ppm

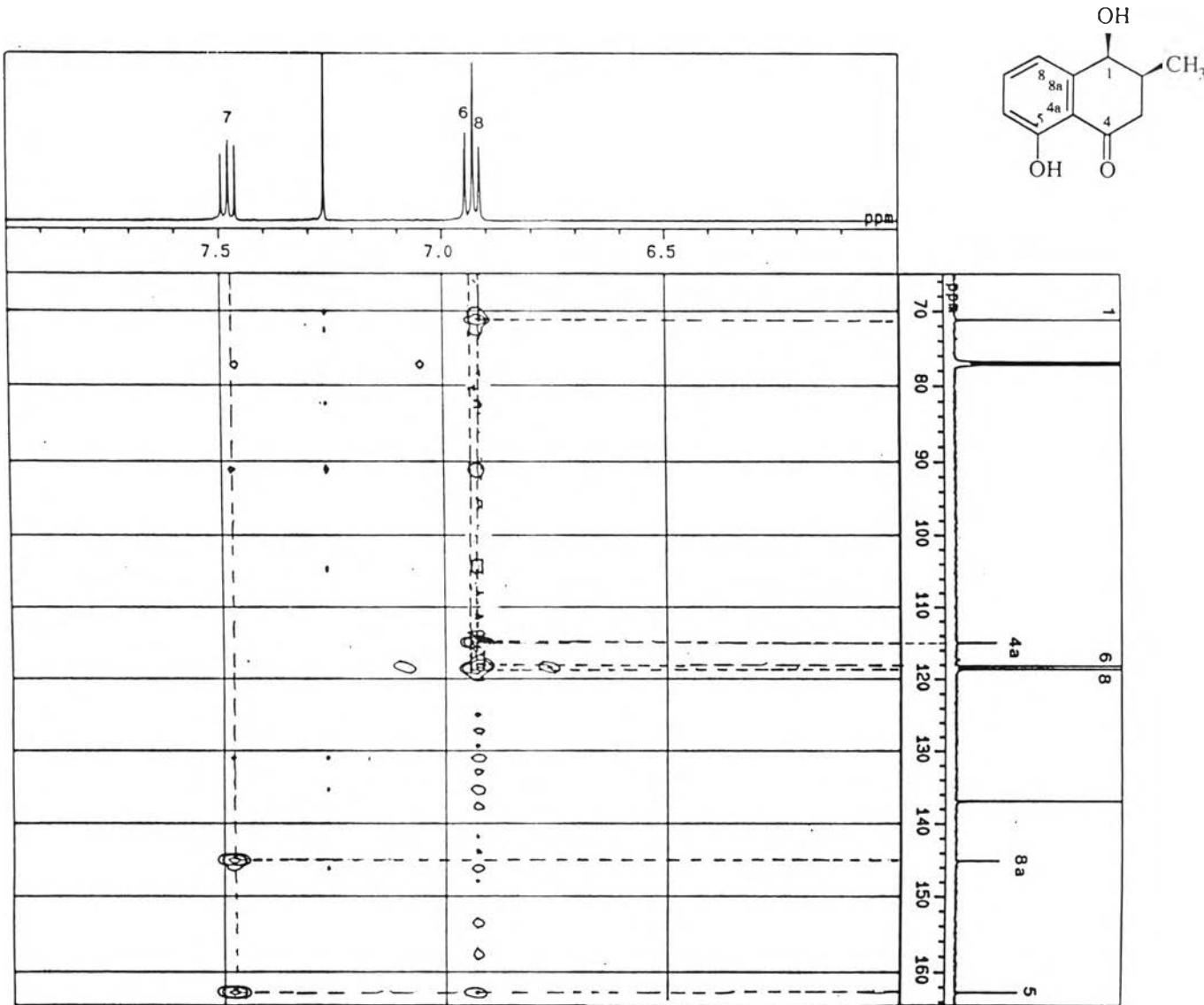


Figure 66 Expansion of the HMBC spectrum of compound 102 (in CDCl_3) :

δ_{H} 6.20-7.90 ; δ_{C} 66.00-164.00 ppm

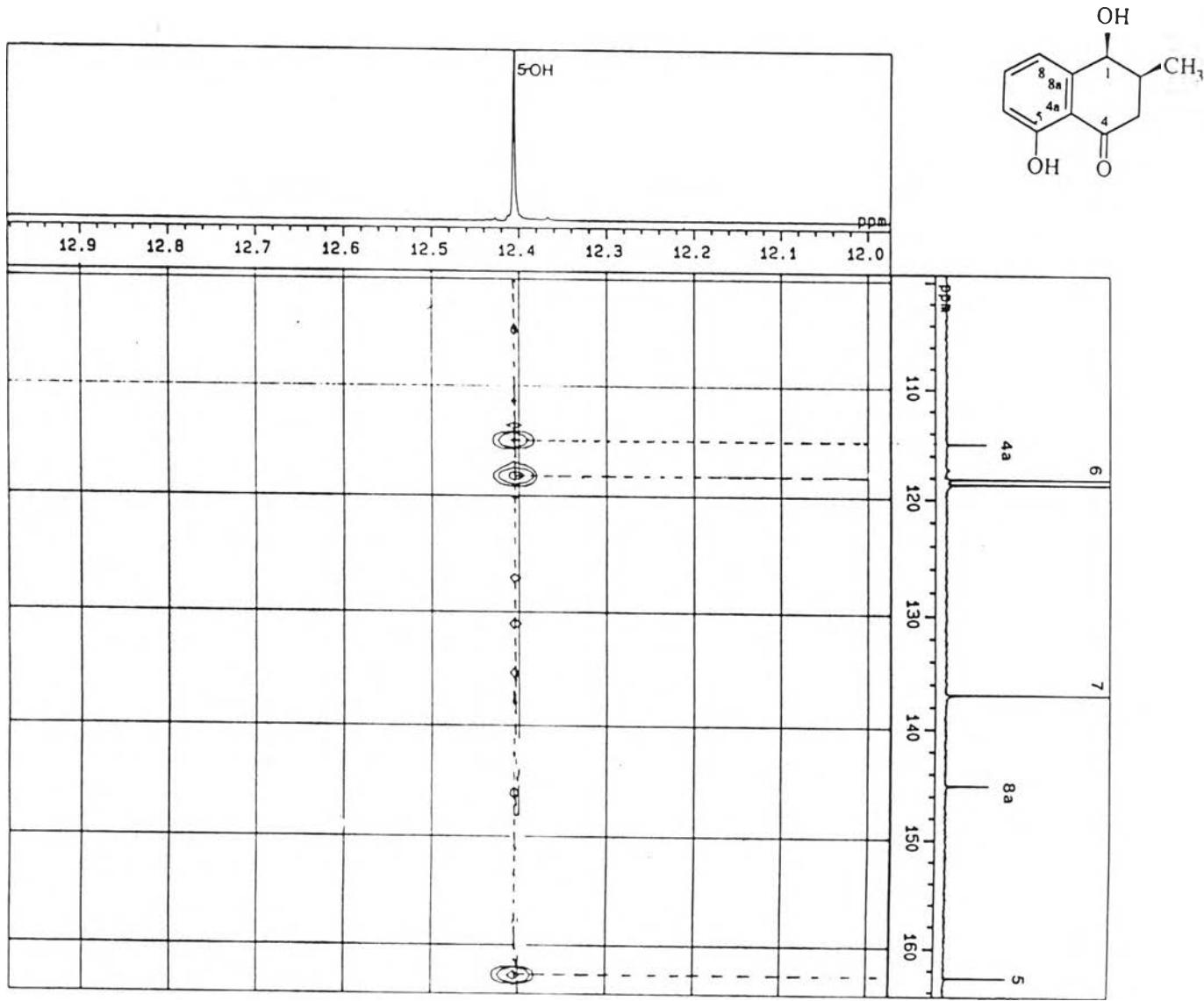


Figure 67 Expansion of the HMBC spectrum of compound 102 (in CDCl₃) :

δ_H 12.00-12.98 ; δ_C 110.00-164.00 ppm

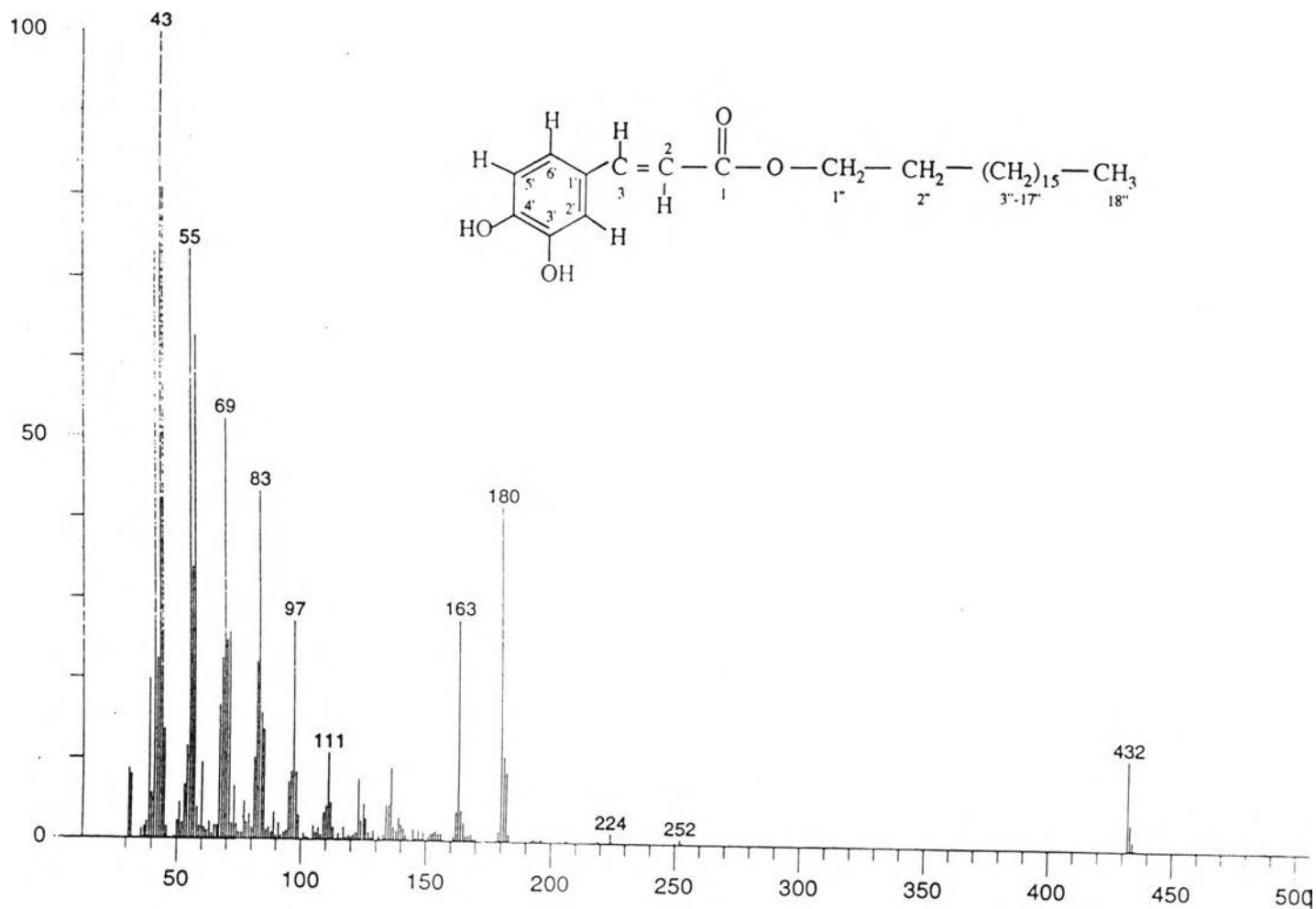


Figure 68 The EI mass spectrum of compound 108

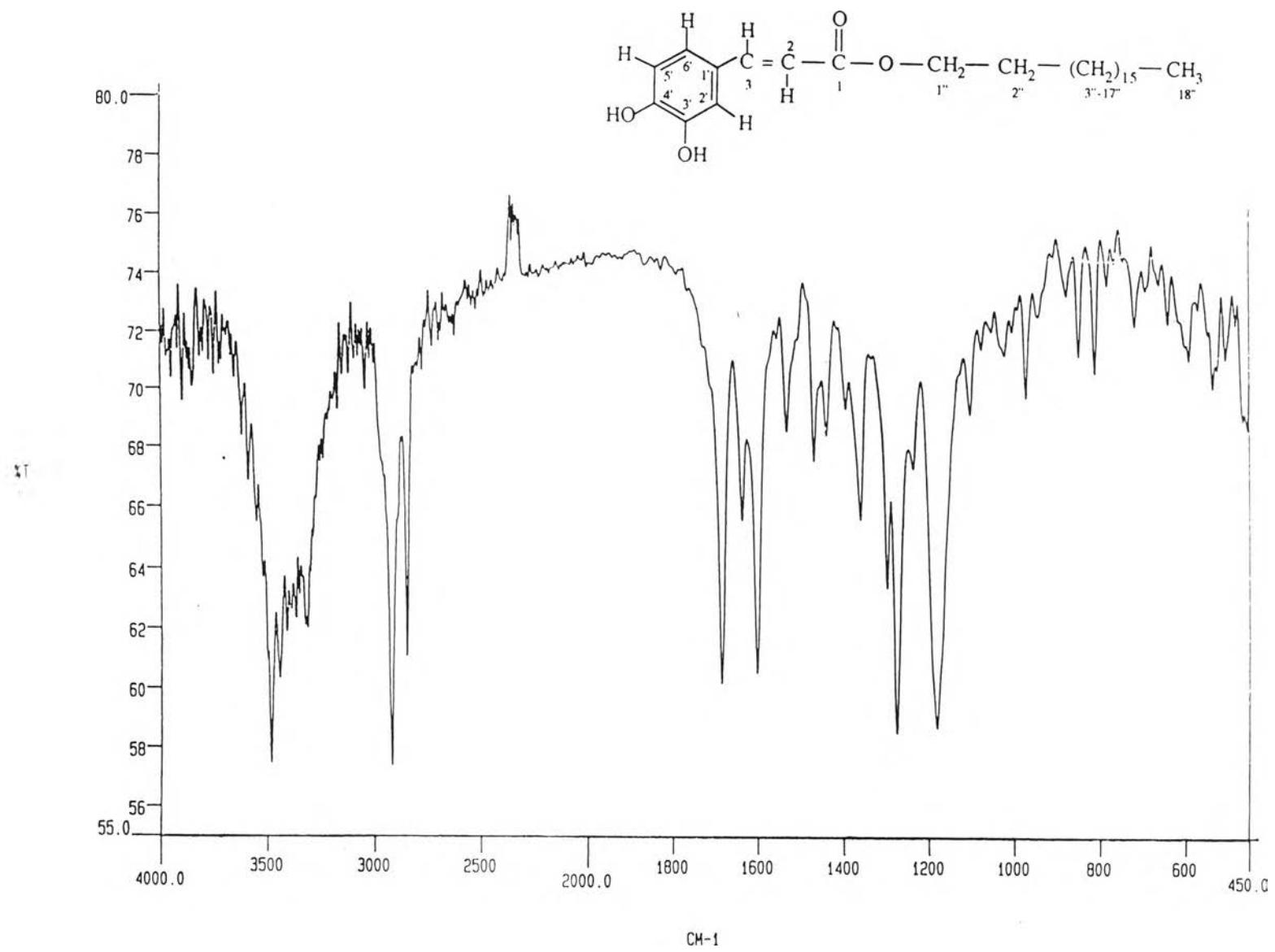


Figure 69 The IR spectrum of compound 103 (in KBr disc)

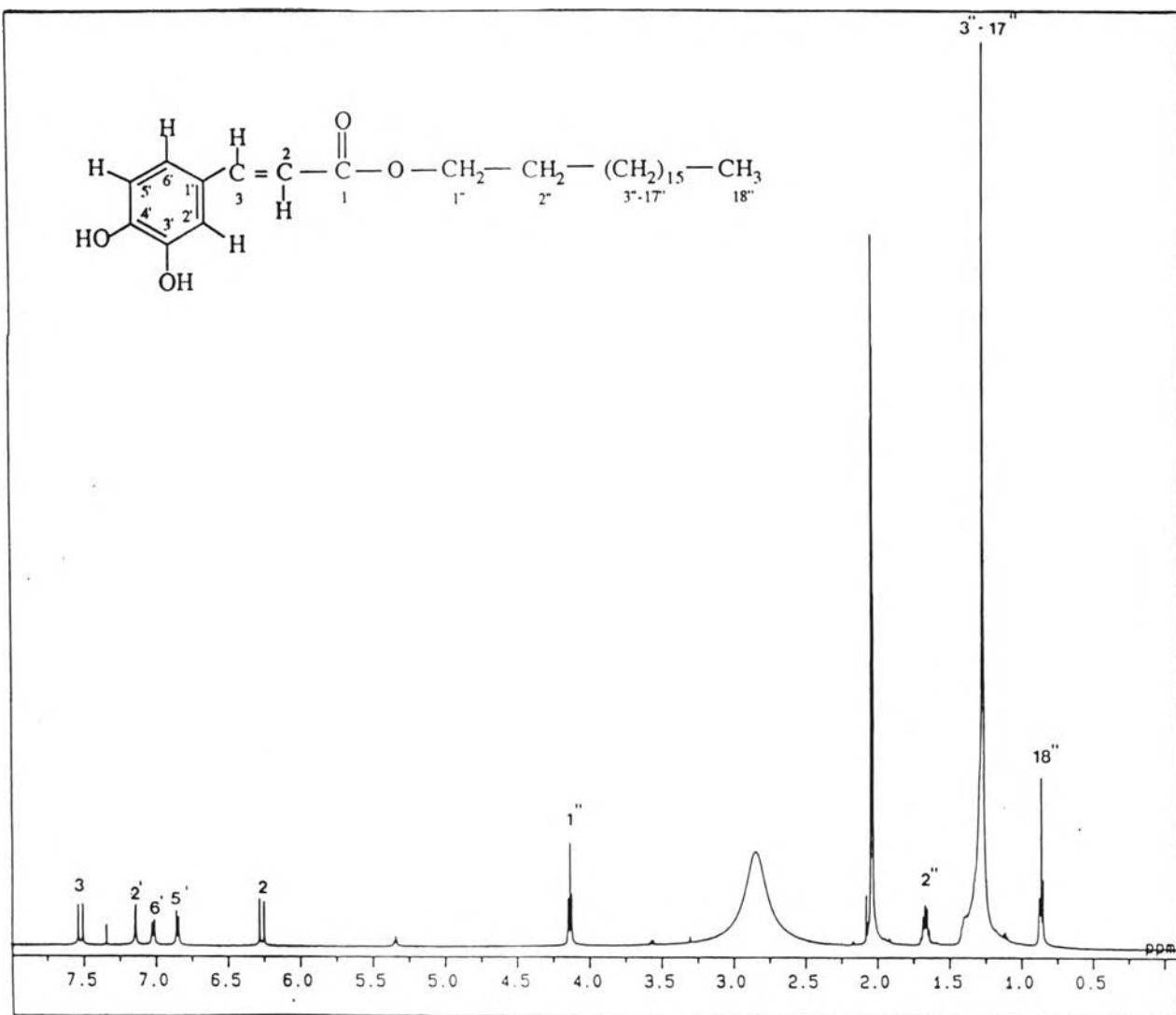


Figure 70 The ^1H NMR(500 MHz) spectrum of compound 103 (in acetone- d_6)

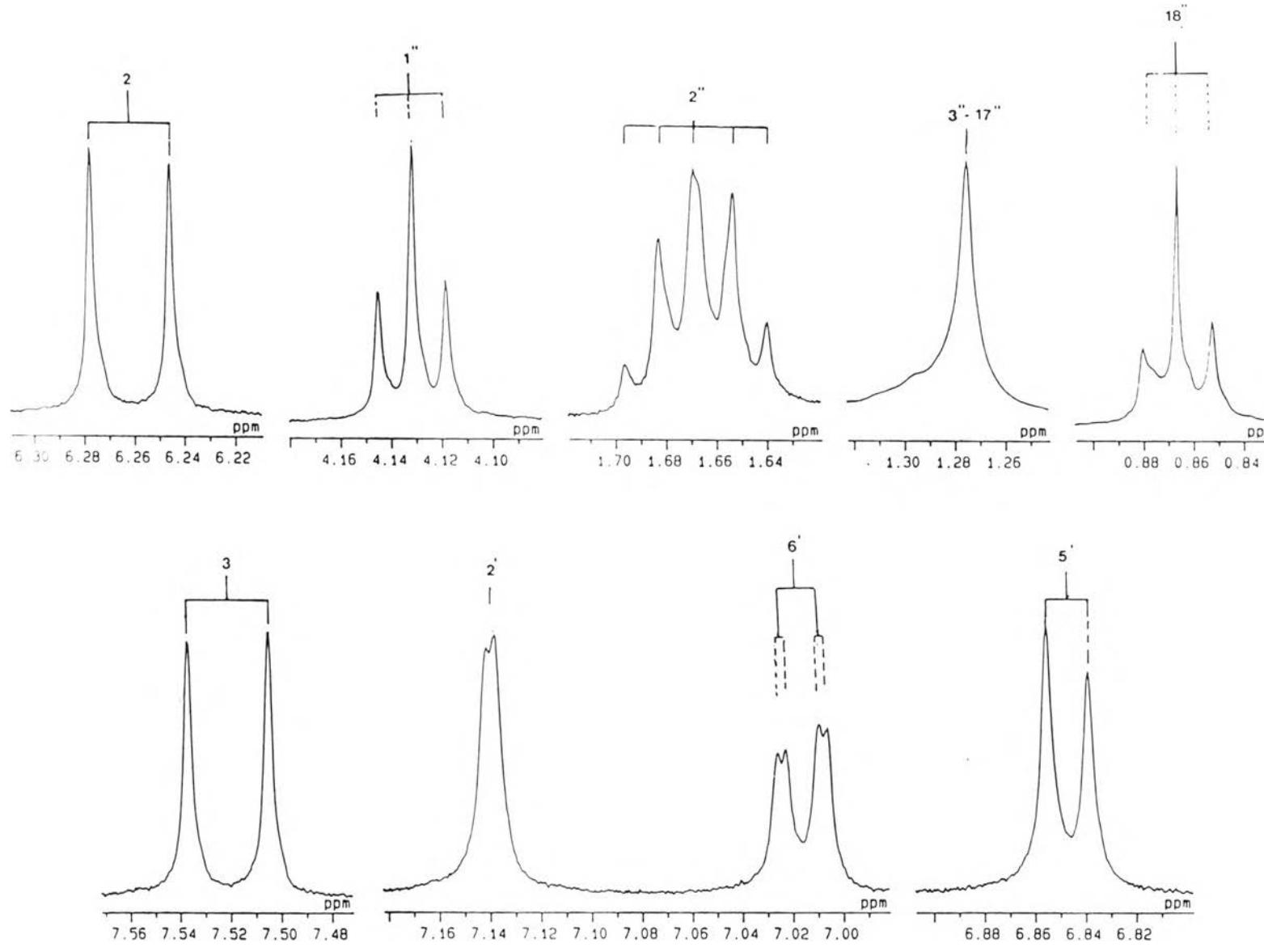


Figure 71 Expansion of the ^1H NMR (500 MHz) spectrum of compound 102

(in acetone- d_6) : δ_{H} 0.84-7.56 ppm

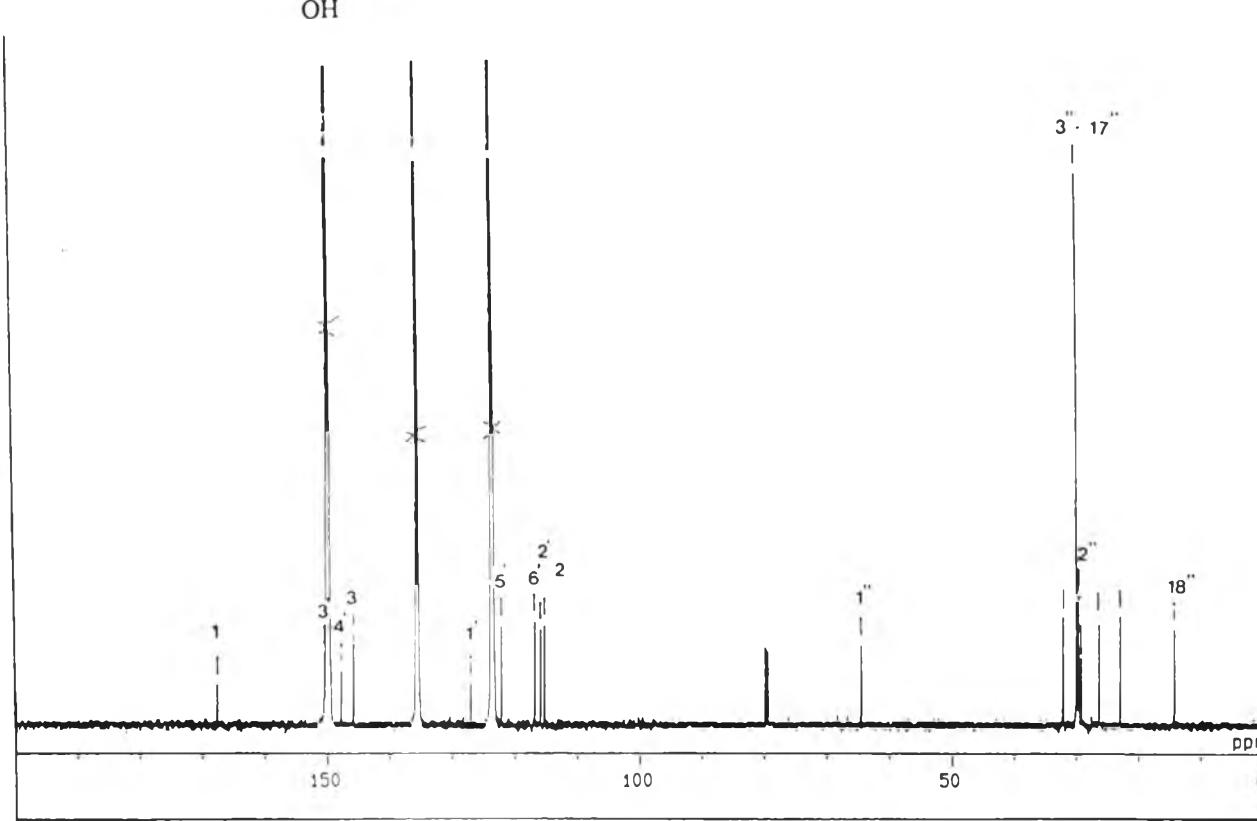
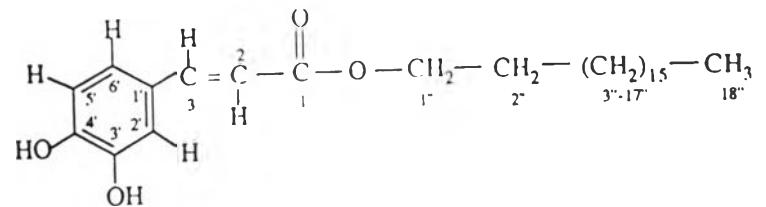


Figure 72 The ¹³C NMR (125 MHz) spectrum of compound 108 (in pyridine-*d*₅)

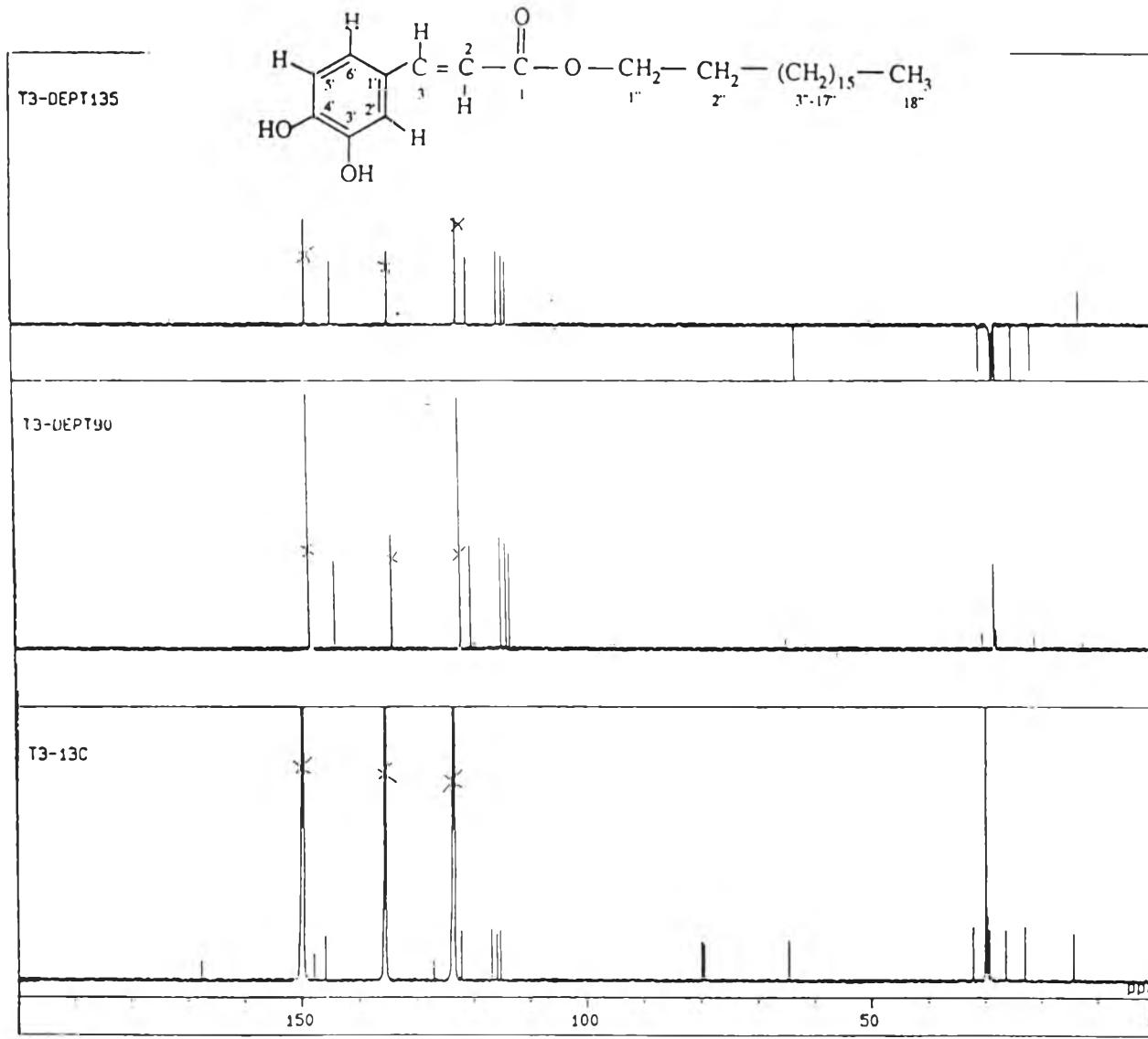


Figure 73 The DEPT (125 MHz) spectrum of compound 103 (in pyridine-*d*₅) :

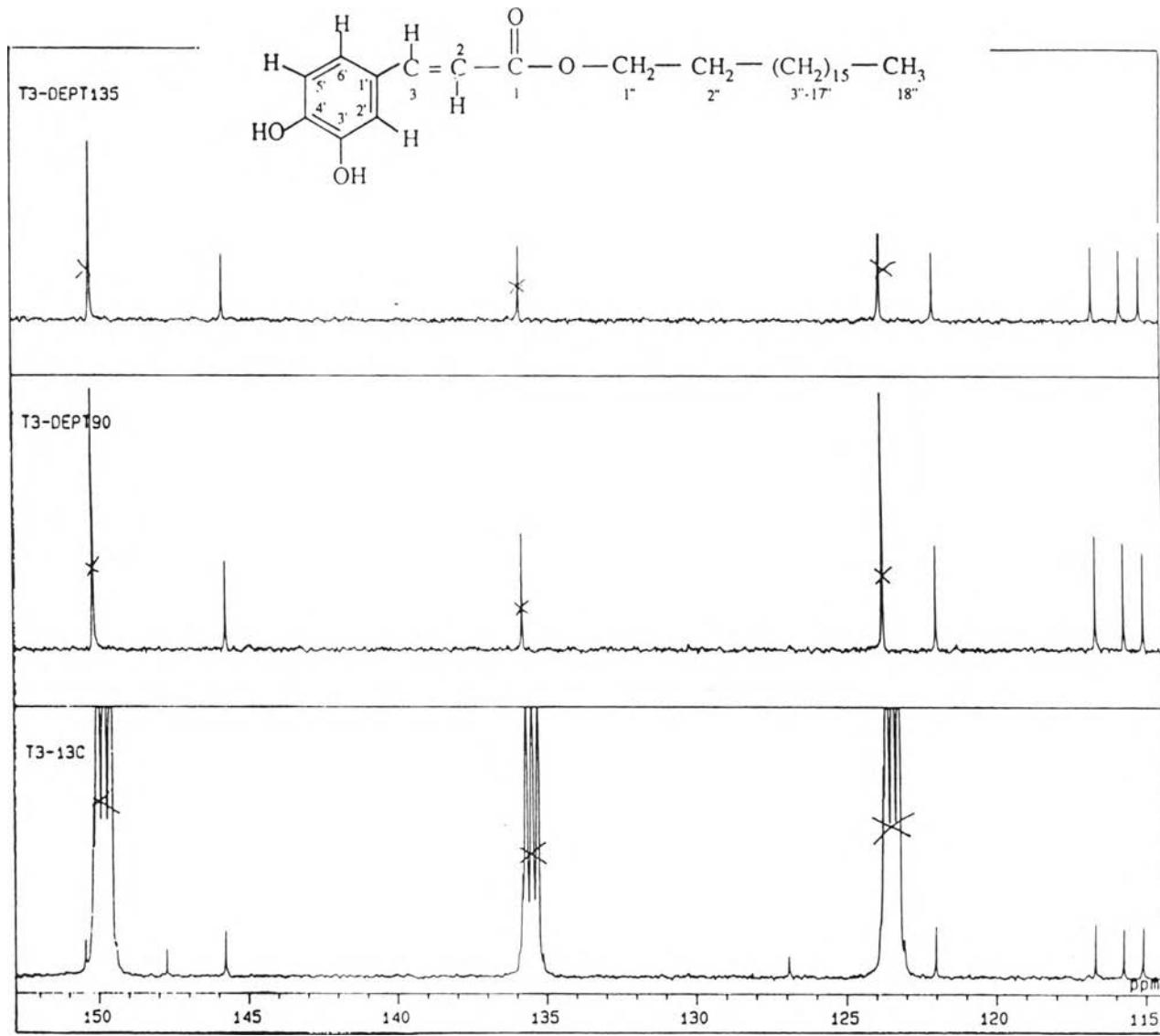


Figure 74 The DEPT (125 MHz) spectrum of compound 108 (in pyridine-*d*₅) :

δ_{C} 115.00-152.00 ppm

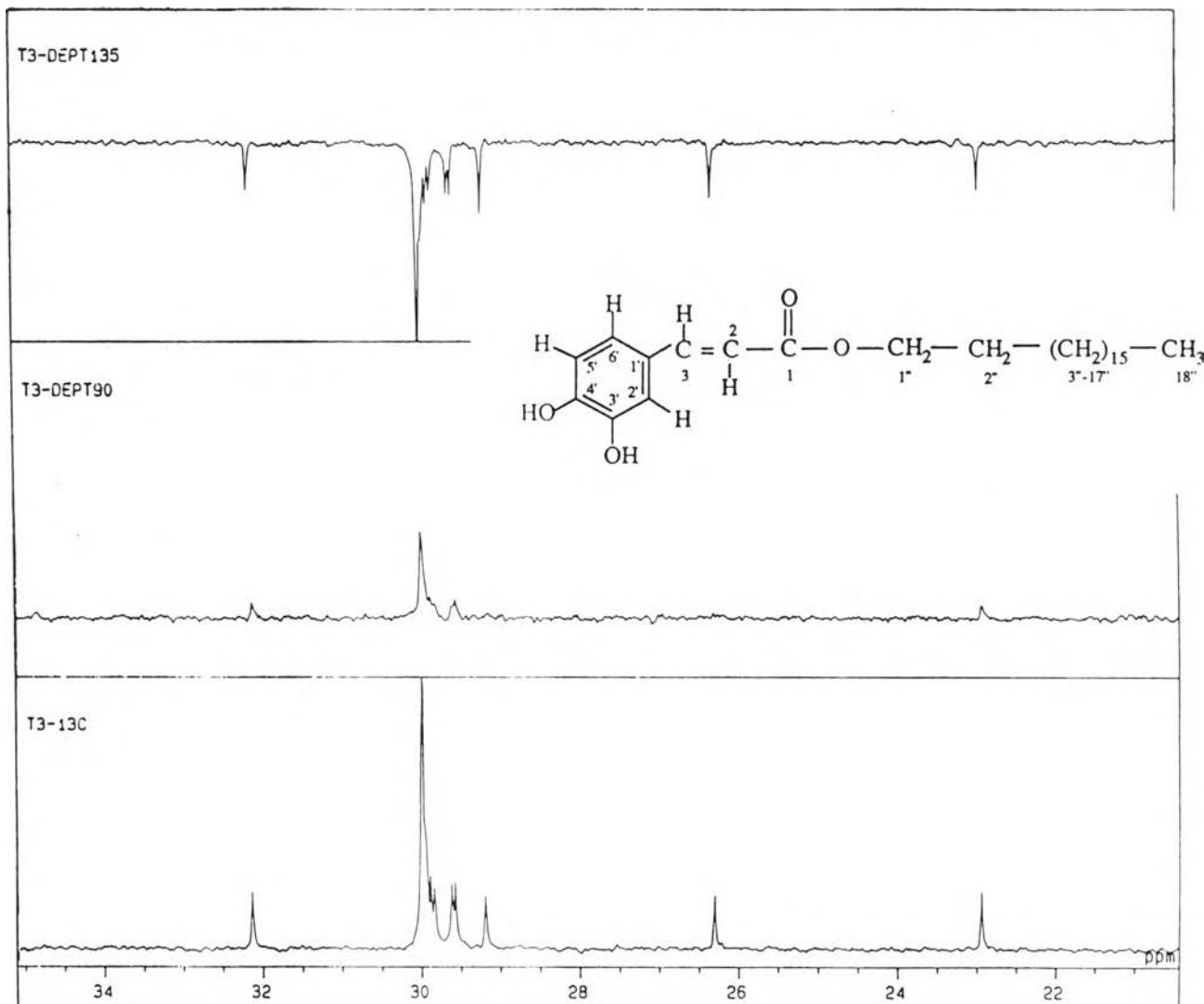


Figure 75 The DEPT (125 MHz) spectrum of compound **103** (in pyridine-*d*₅)
 δ_{C} 23.00-35.00 ppm

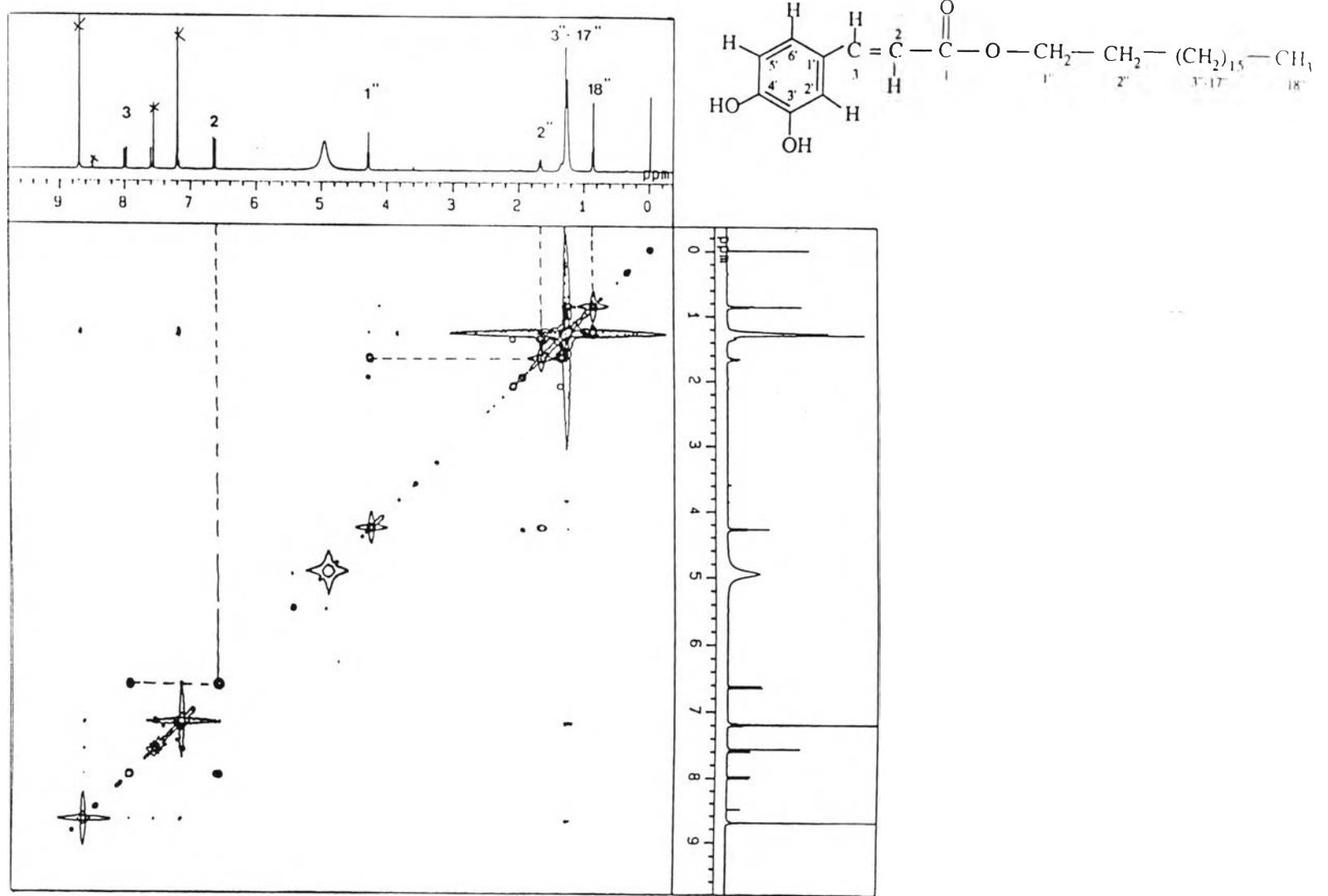


Figure 76 The ^1H - ^1H COSY (500 MHz) spectrum of compound 103
(in $\text{pyridine}-d_5$)

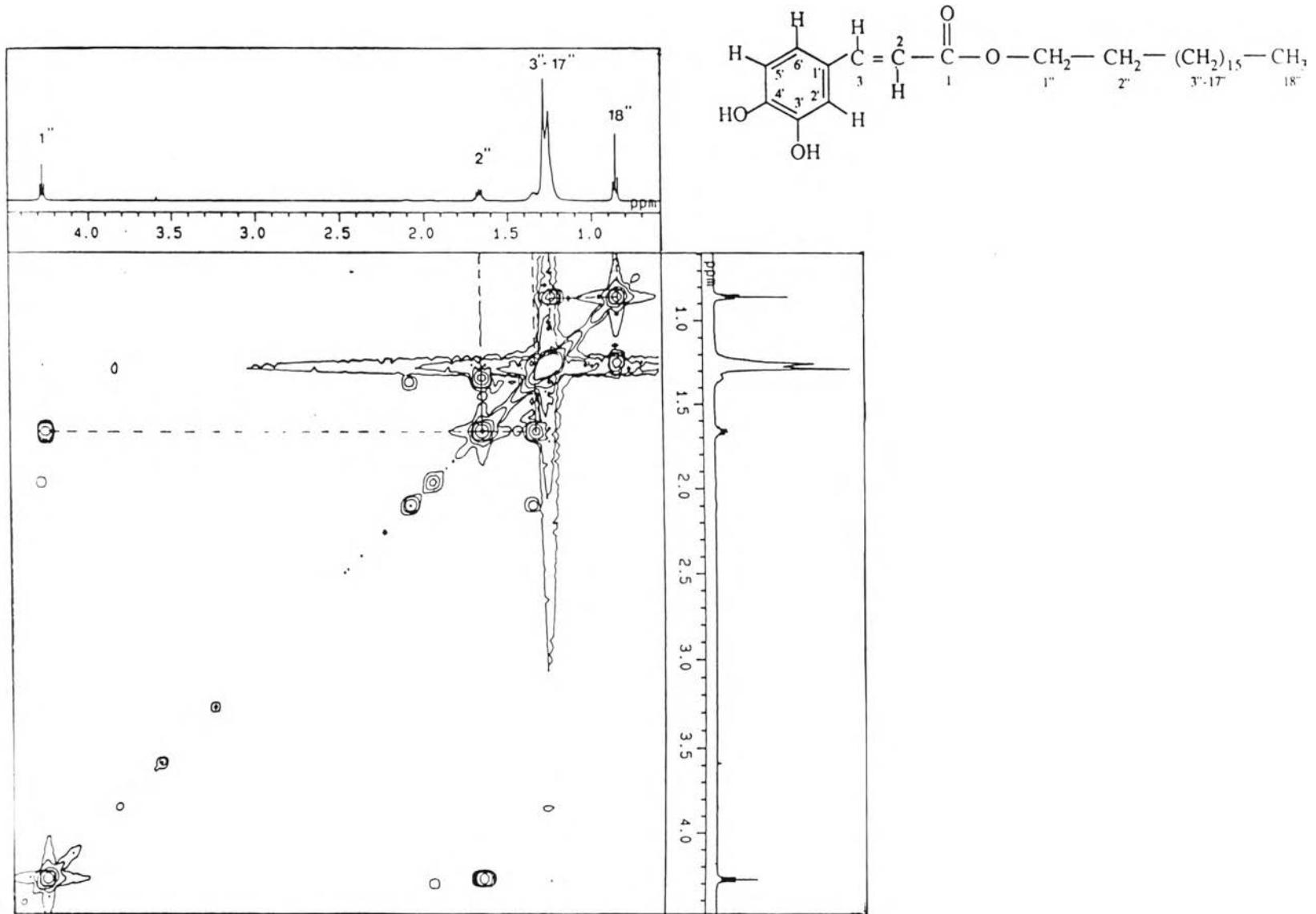


Figure 77 Expansion of the ^1H - ^1H COSY (500 MHz) spectrum of compound 103
(in pyridine- d_5) : δ_{H} 0.70-4.40 ppm

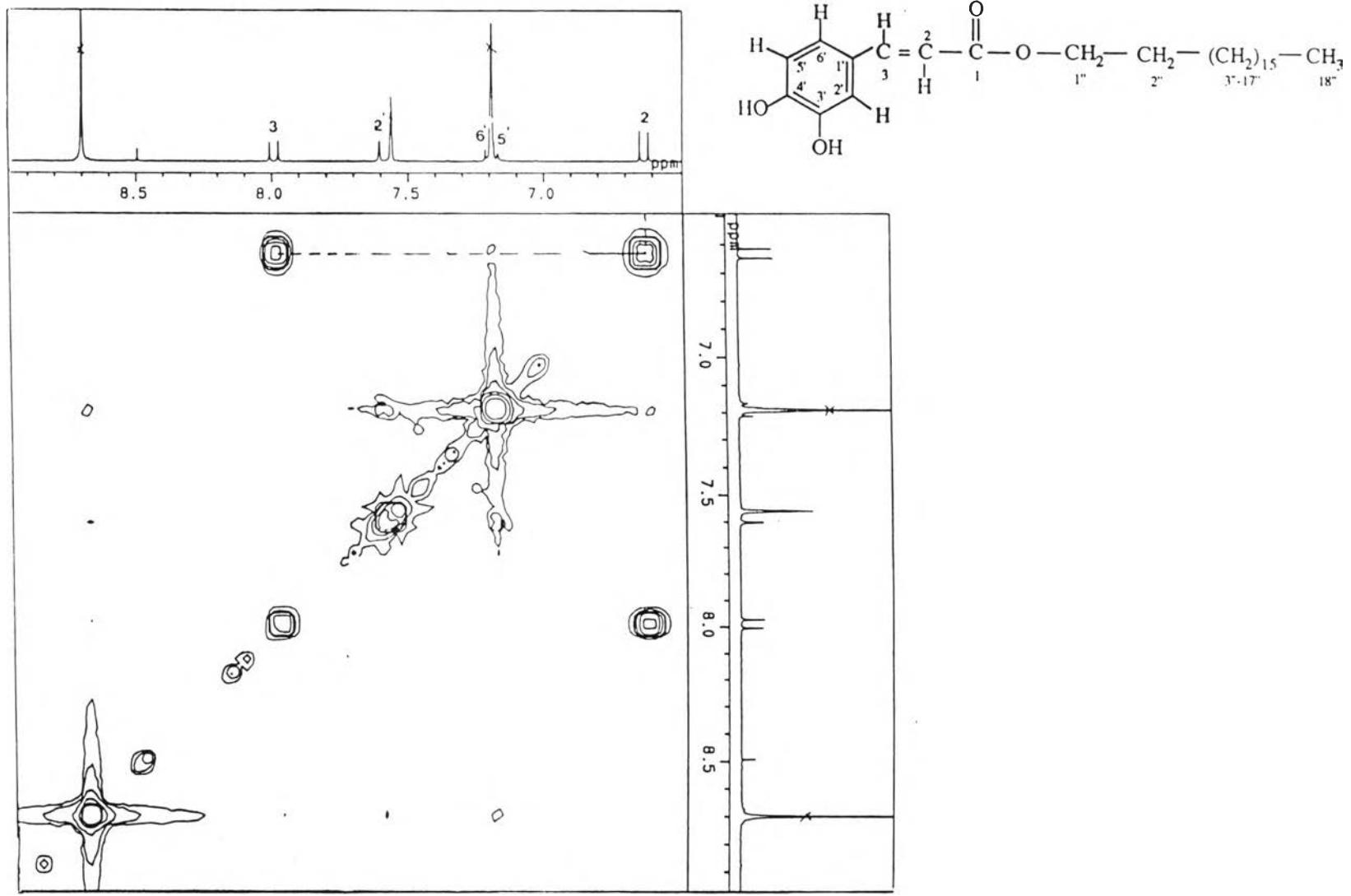


Figure 78 Expansion of the $^1\text{H}-^1\text{H}$ COSY (500 MHz) spectrum of compound 108
 (in pyridine- d_5) : δ_{H} 6.60-8.90 ppm

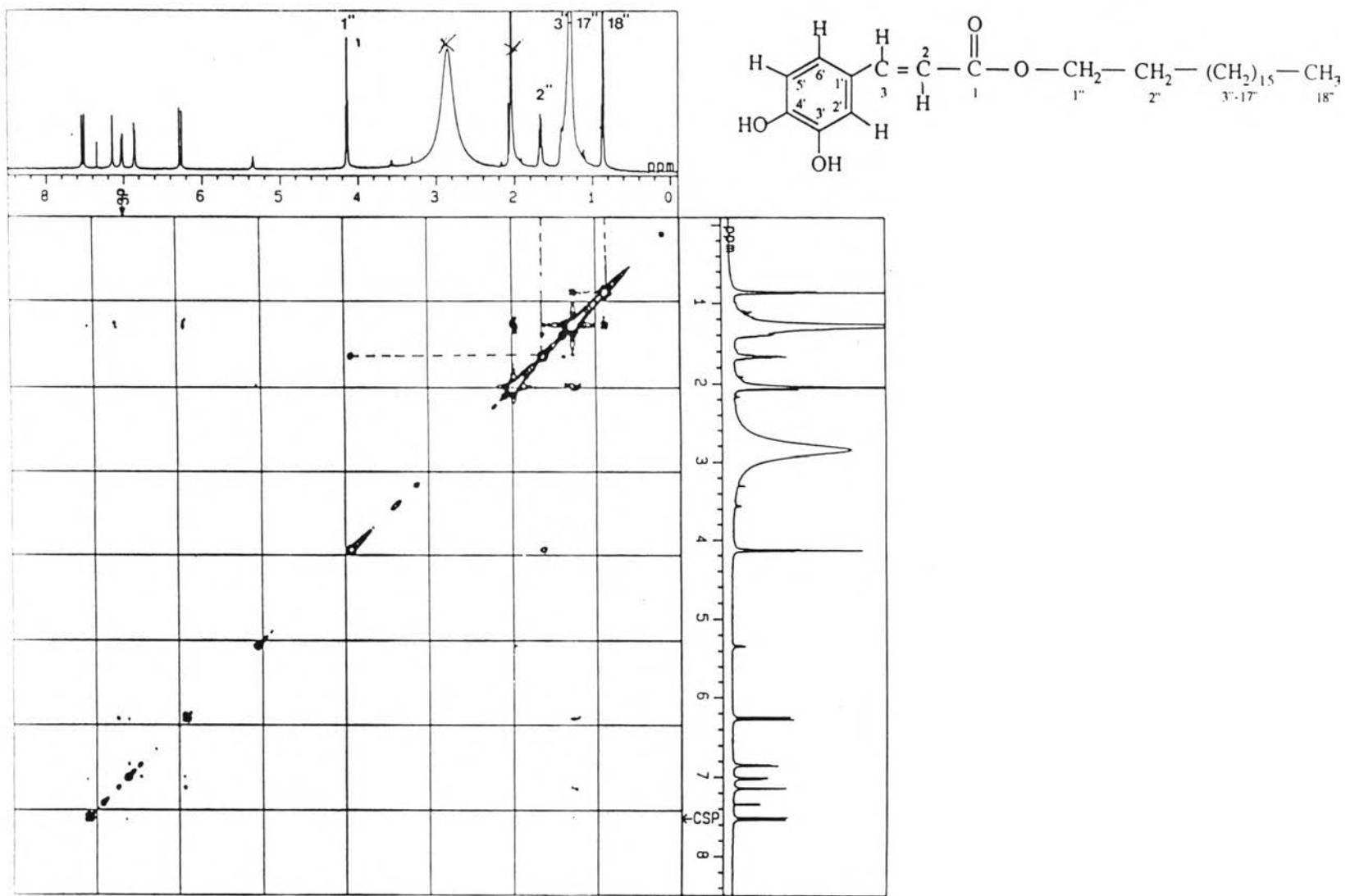


Figure 79 The NOESY (500 MHz) spectrum of compound 108 (in acetone- d_6)

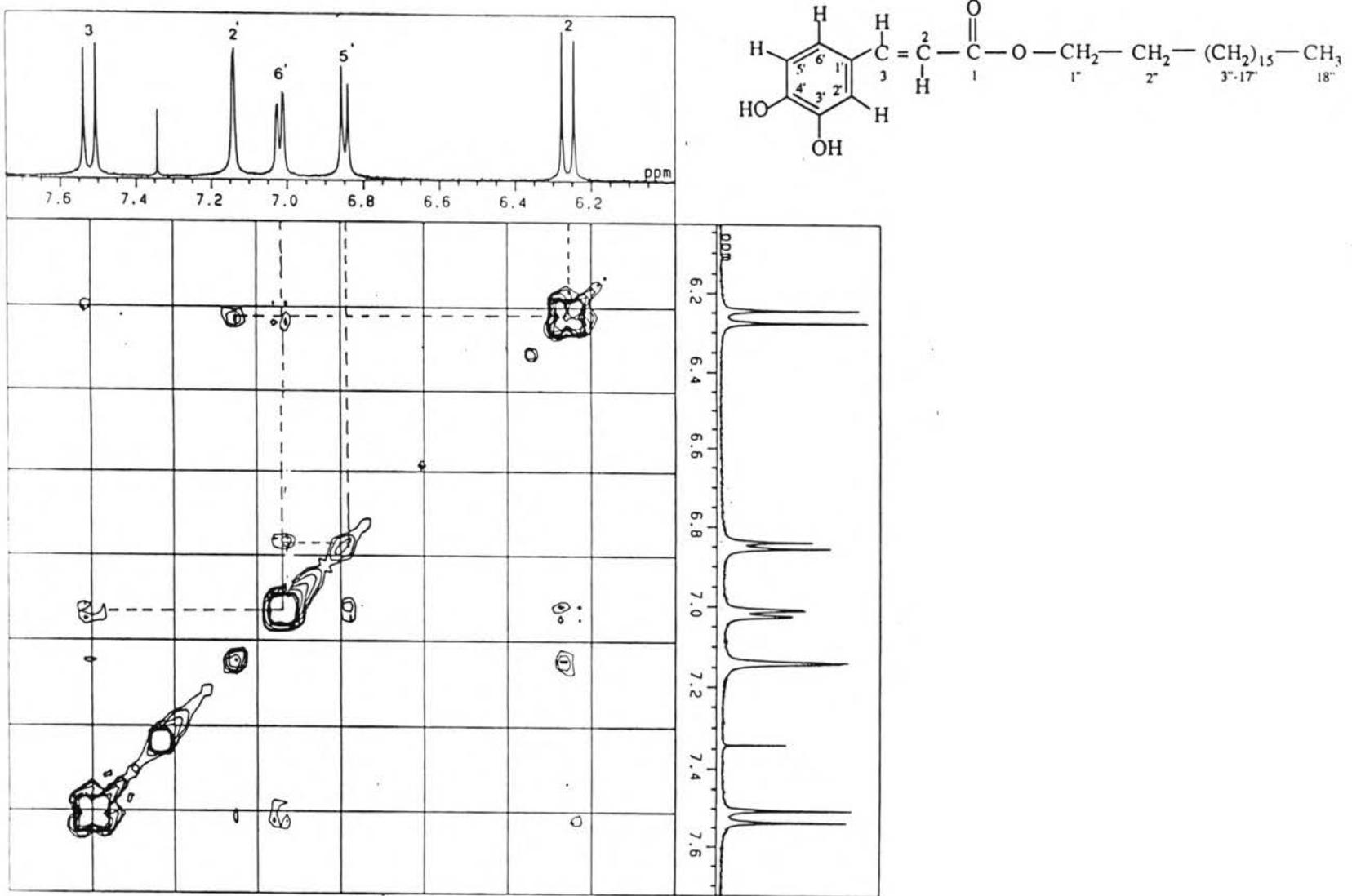


Figure 80 Expansion of the NOESY (500 MHz) spectrum of compound 103

(in acetone- d_6) : δ_H 6.20-7.60 ppm

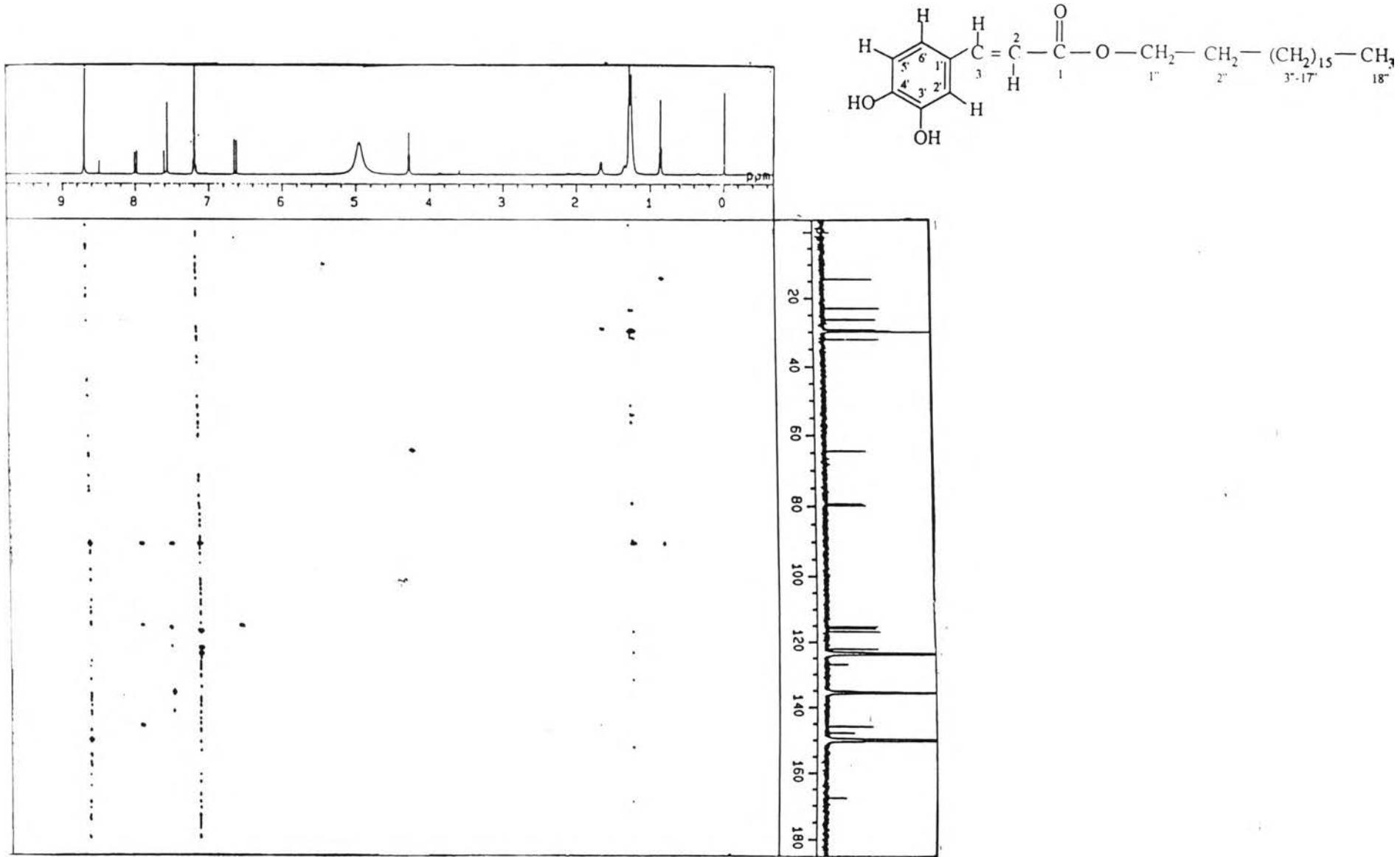


Figure 81 The HMQC spectrum of compound 103 (in pyridine- d_5)

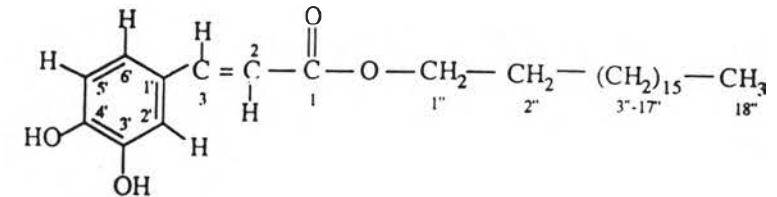
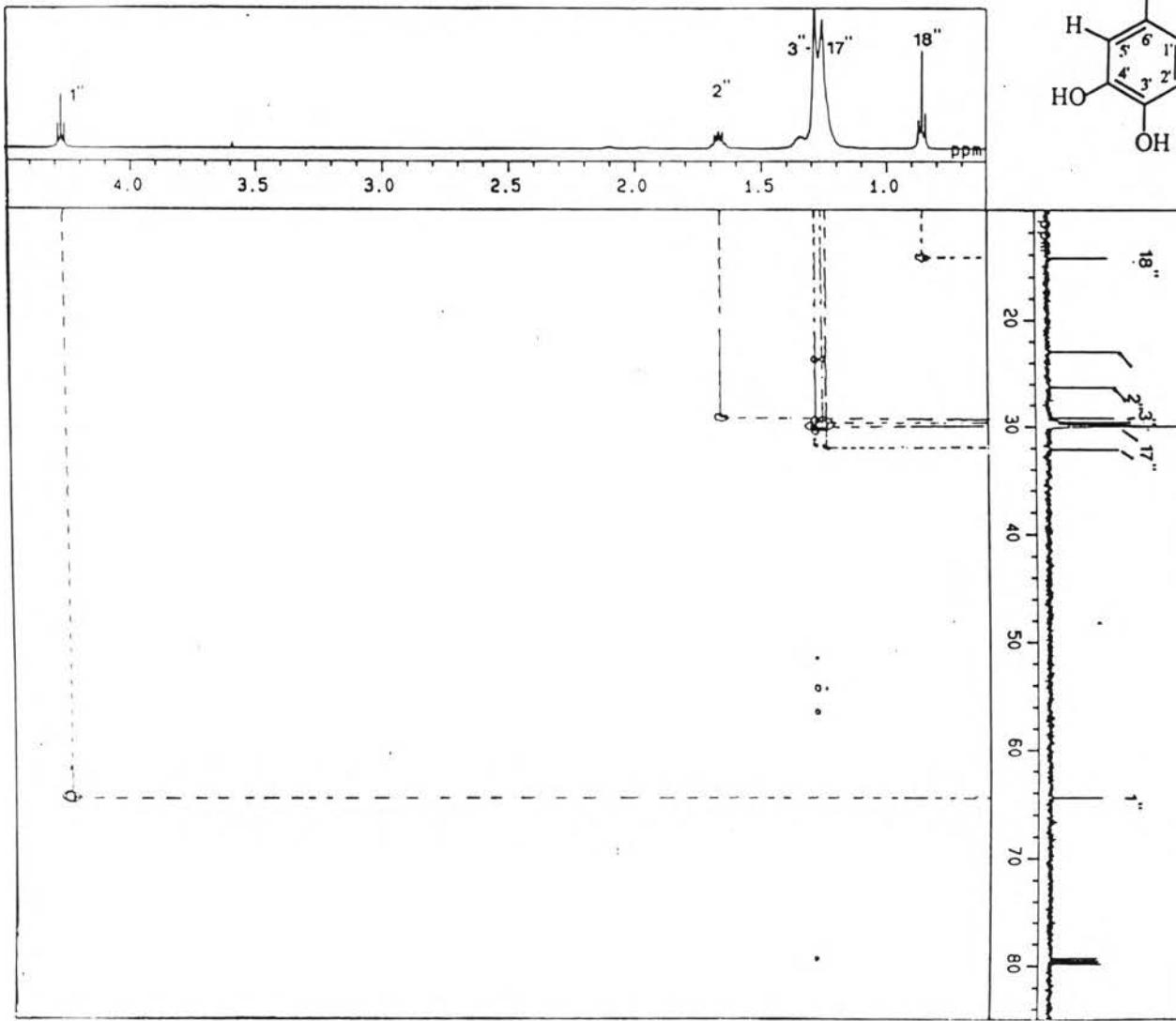


Figure 82 Expansion of the HMQC spectrum of compound 103 (in pyridine- d_5) :

δ_H 0.70-4.40 ; δ_C 10.00-80.00 ppm

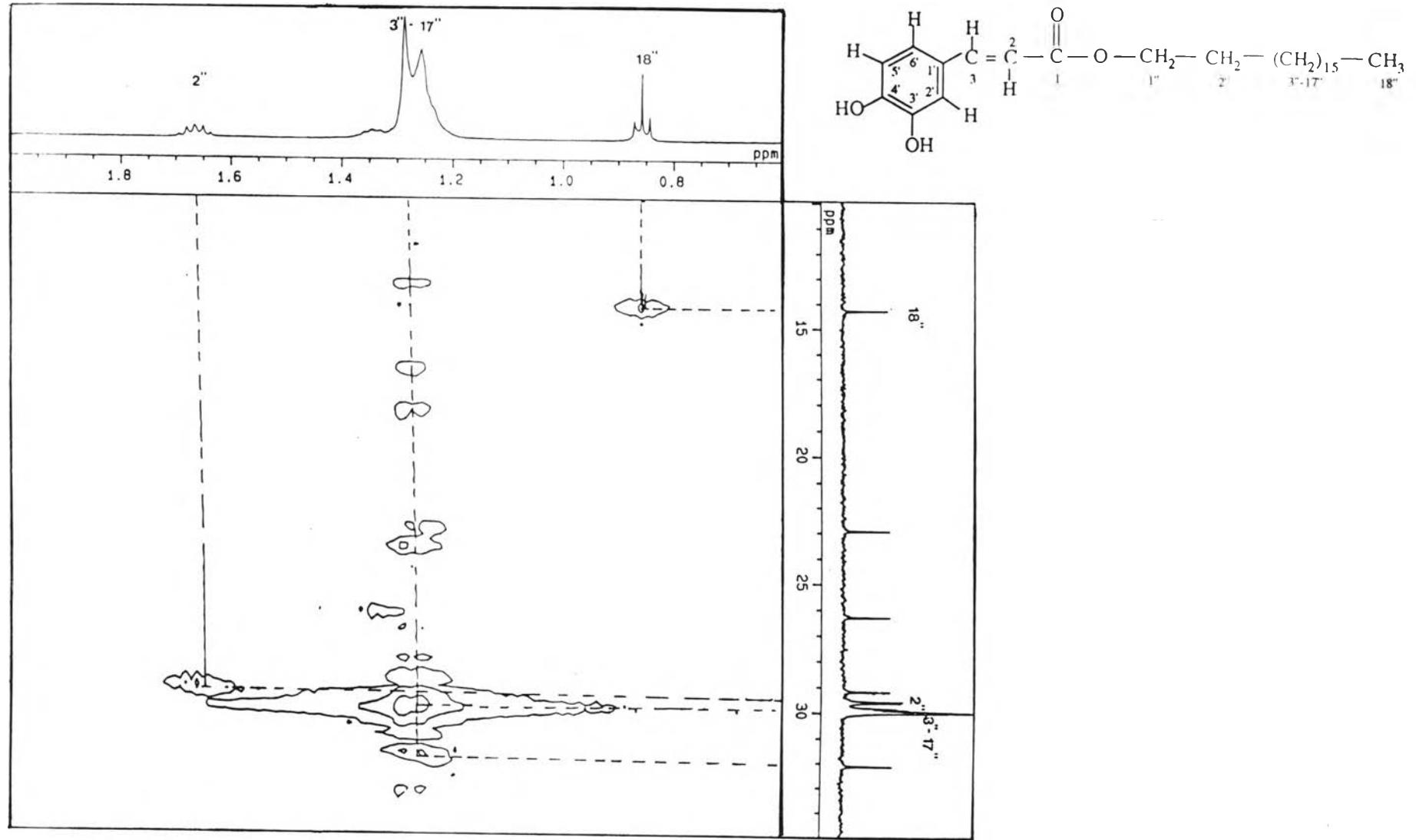


Figure 83 Expansion of the HMQC spectrum of compound 103 (in pyridine- d_5) :

δ_H 0.80-1.80 ; δ_C 10.00-34.00 ppm

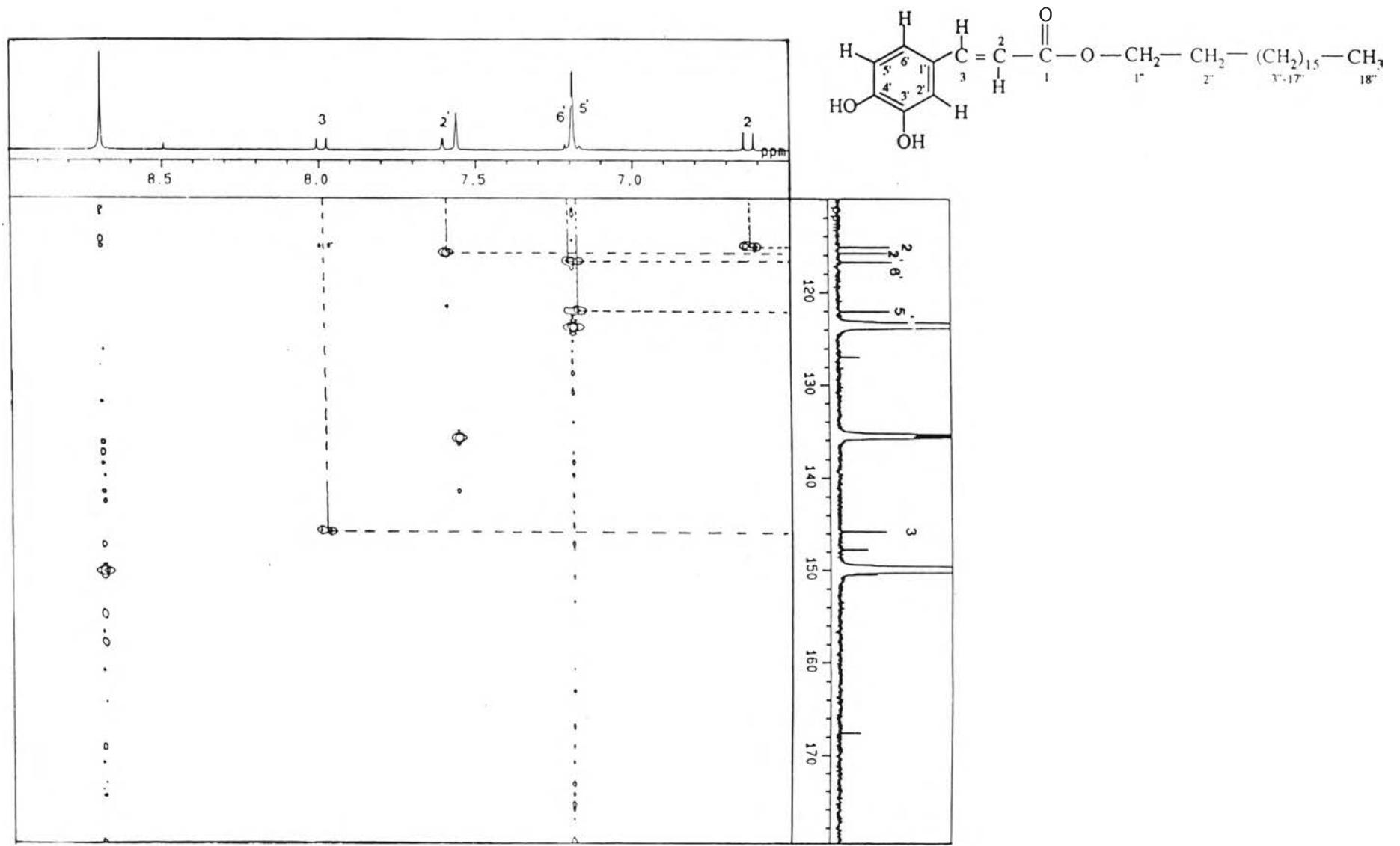


Figure 84 Expansion of the HMQC spectrum of compound 103 (in pyridine- d_5) :

δ_H 6.60-8.90 ; δ_C 110.00-178.00 ppm

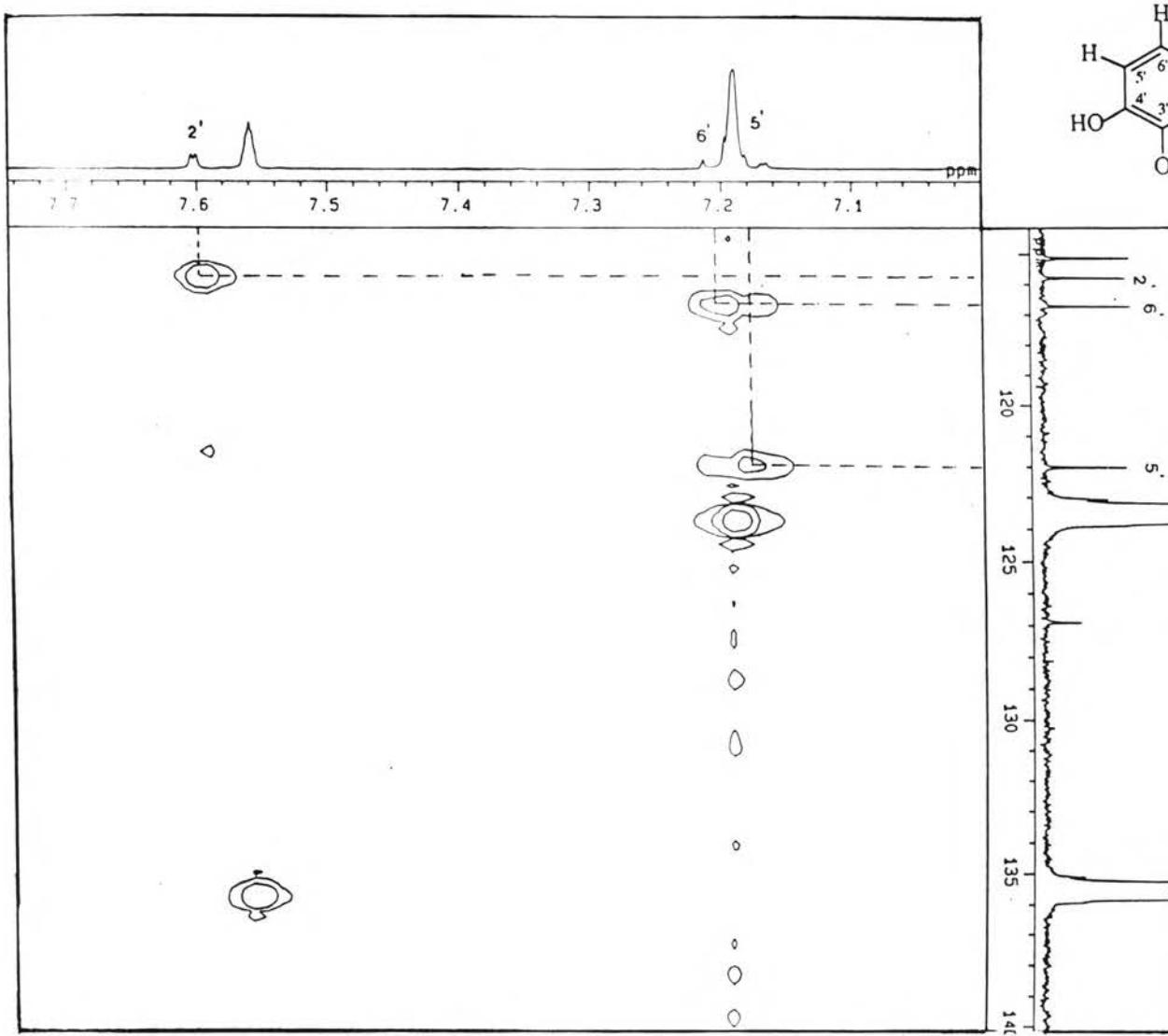


Figure 85 Expansion of the HMQC spectrum of compound 108 (in pyridine- d_5) :

δ_H 7.02-7.74 ; δ_C 115.00-140.00 ppm

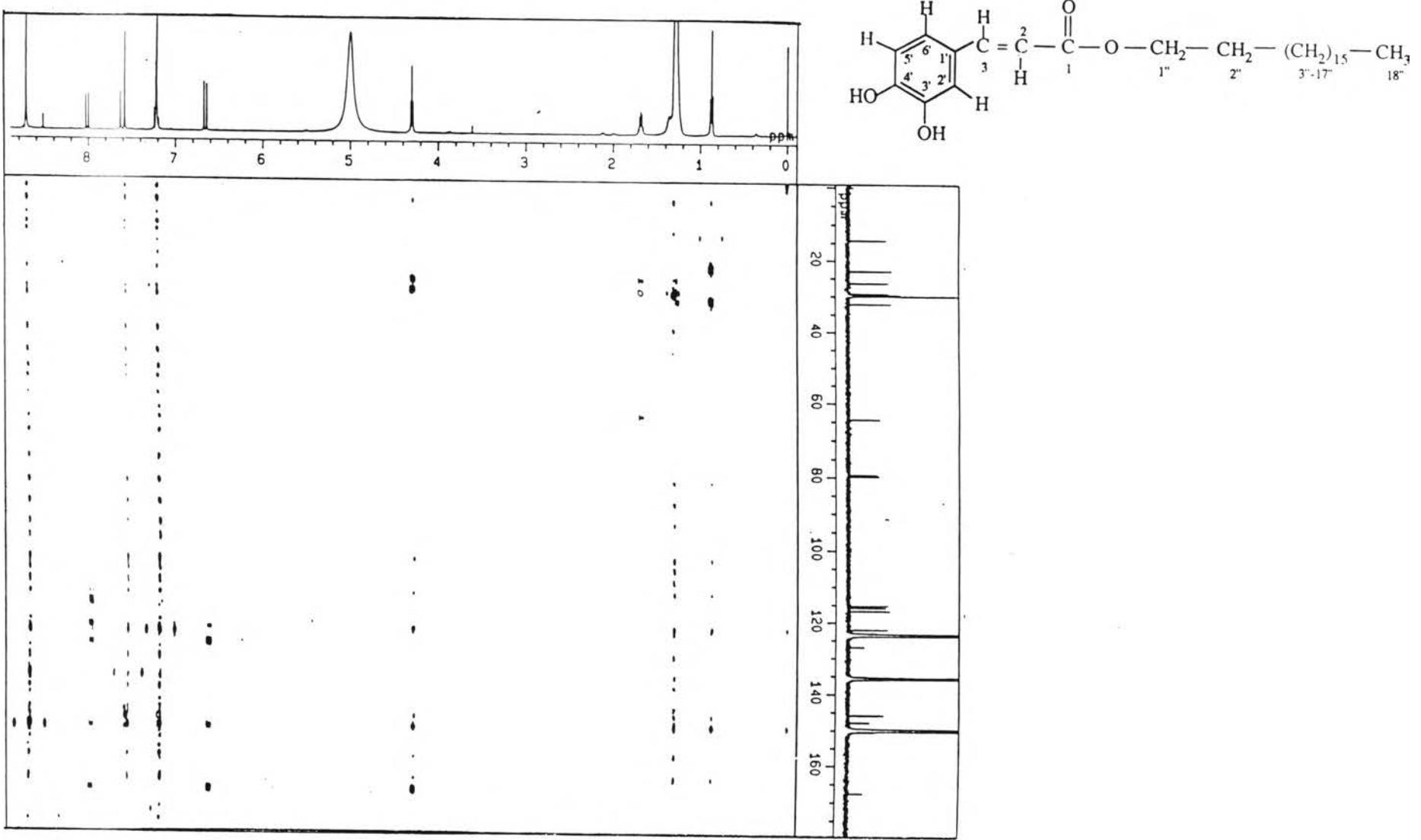


Figure 86 The HMBC spectrum of compound 108 (in pyridine-*d*₅)

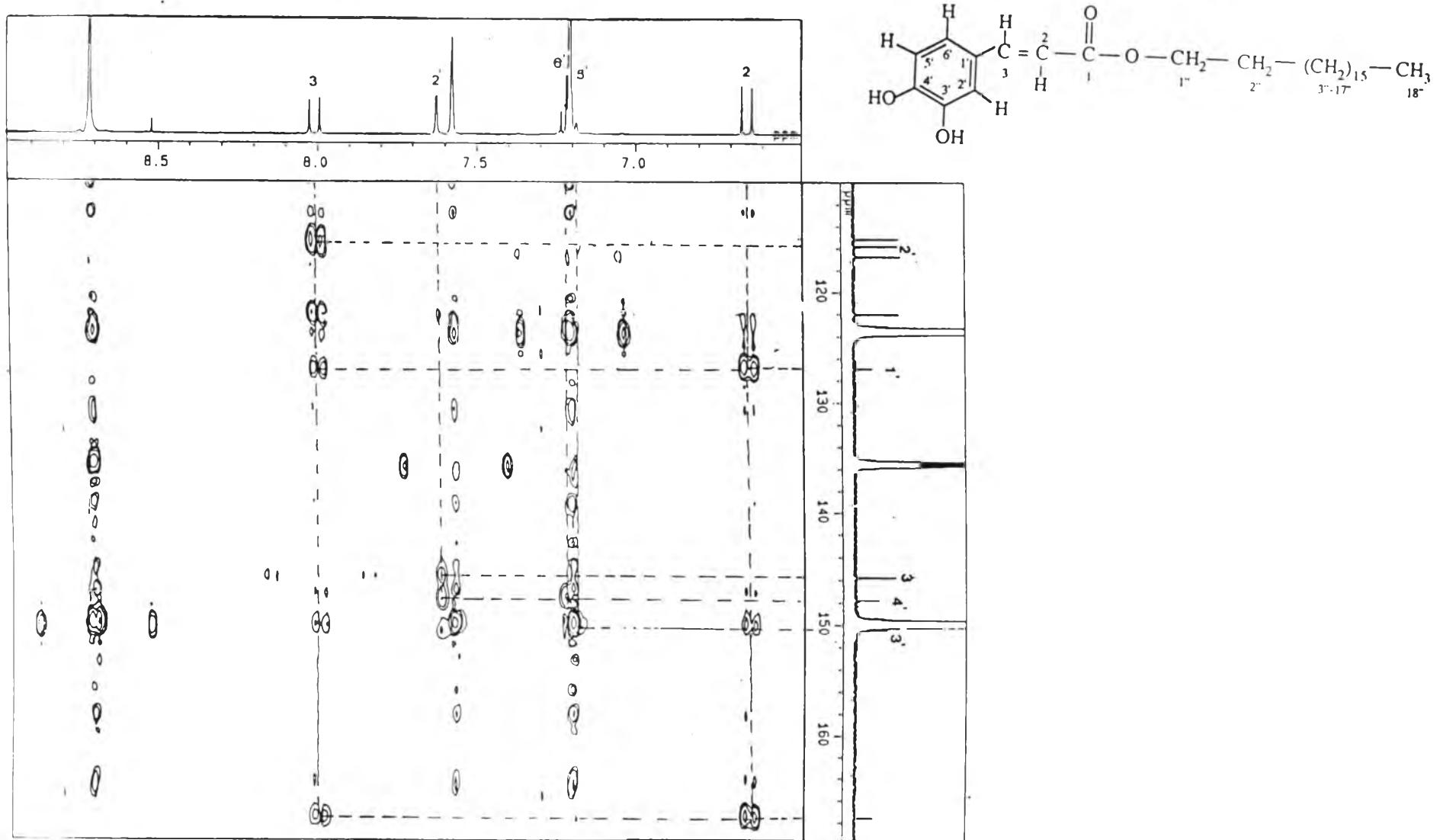


Figure 87 Expansion of the HMBC spectrum of compound 103 (in pyridine- d_5) :

δ_H 6.60-8.90 ; δ_C 110.00-168.00 ppm

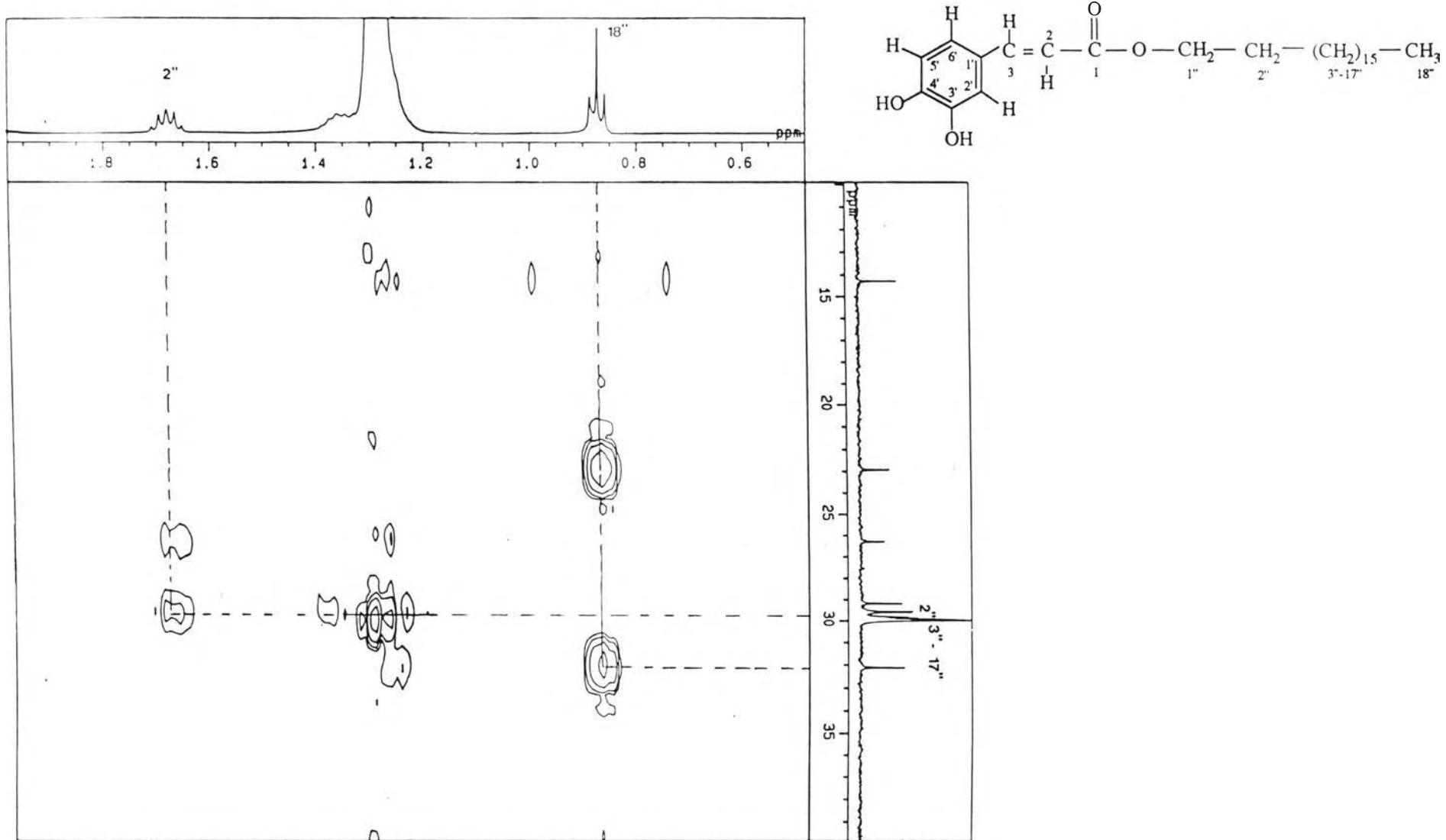


Figure 88 Expansion of the HMBC spectrum of compound 103 (in pyridine- d_5) :

δ_H 0.60-1.80 ; δ_C 10.00-35.00 ppm

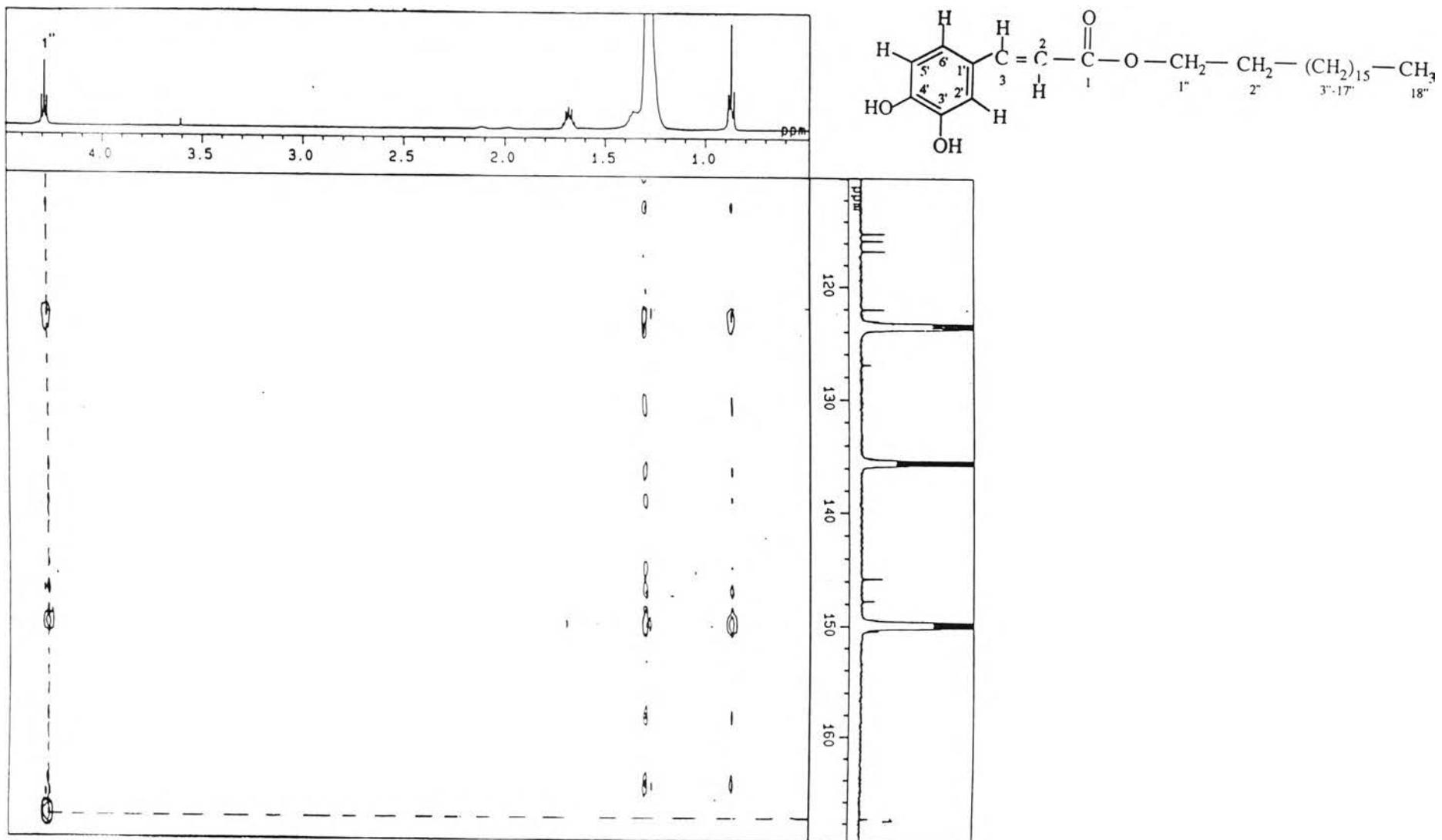


Figure 89 Expansion of the HMBC spectrum of compound 103 (in pyridine- d_5) :

δ_H 0.60-4.40 ; δ_C 110.00-164.00 ppm

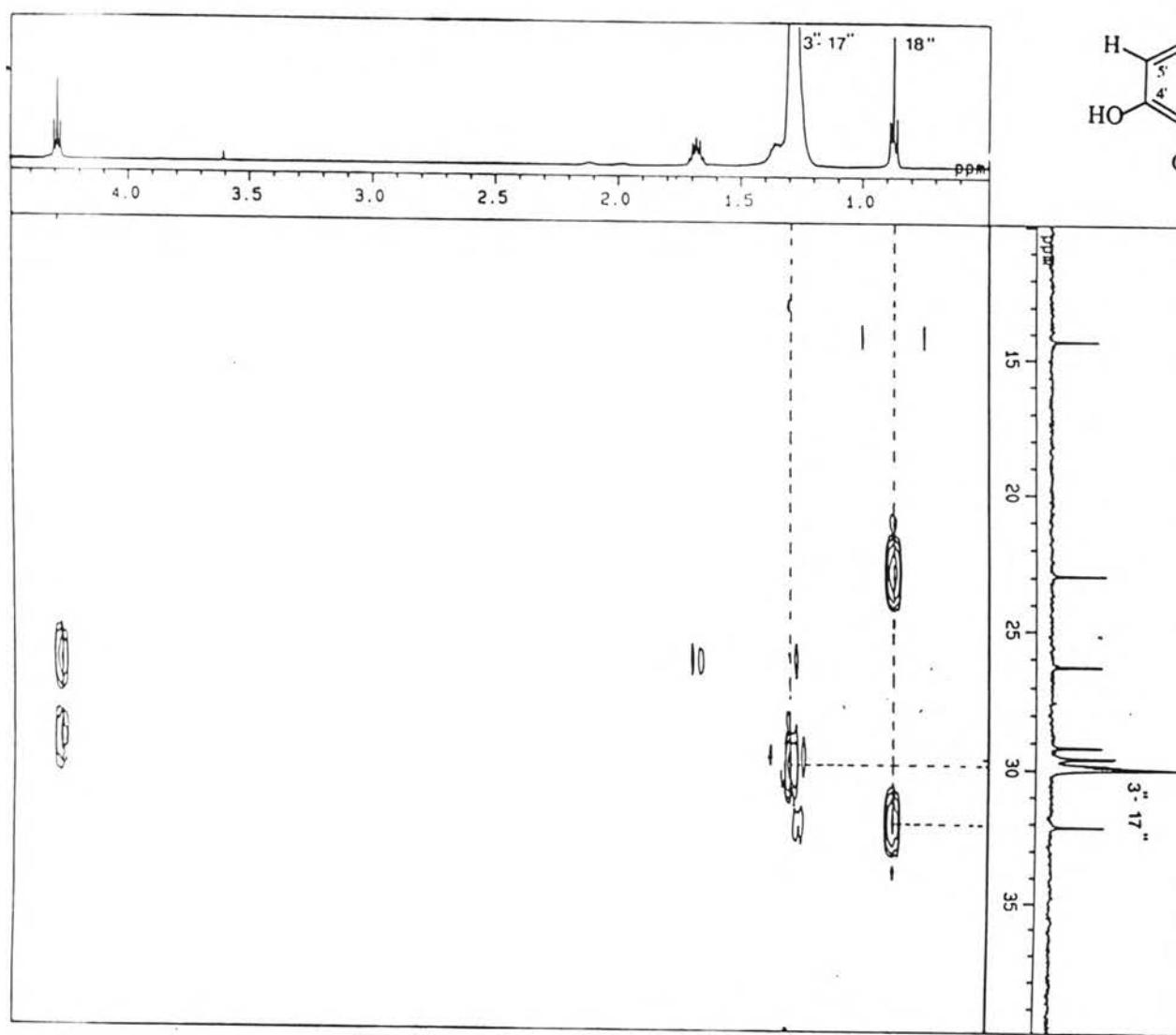


Figure 90 Expansion of the HMBC spectrum of compound 108 (in pyridine-*d*₅) :

δ_{H} 0.60-4.40 ; δ_{C} 10.00-39.00 ppm

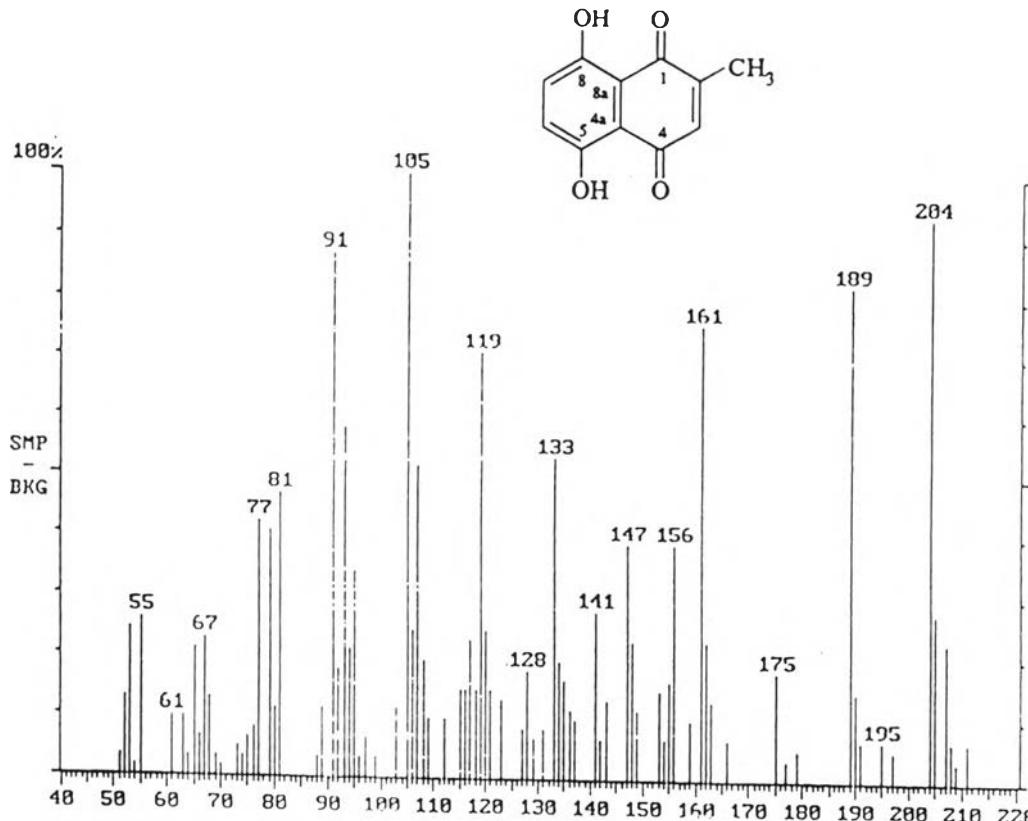


Figure 91 The EI mass spectrum of compound 104

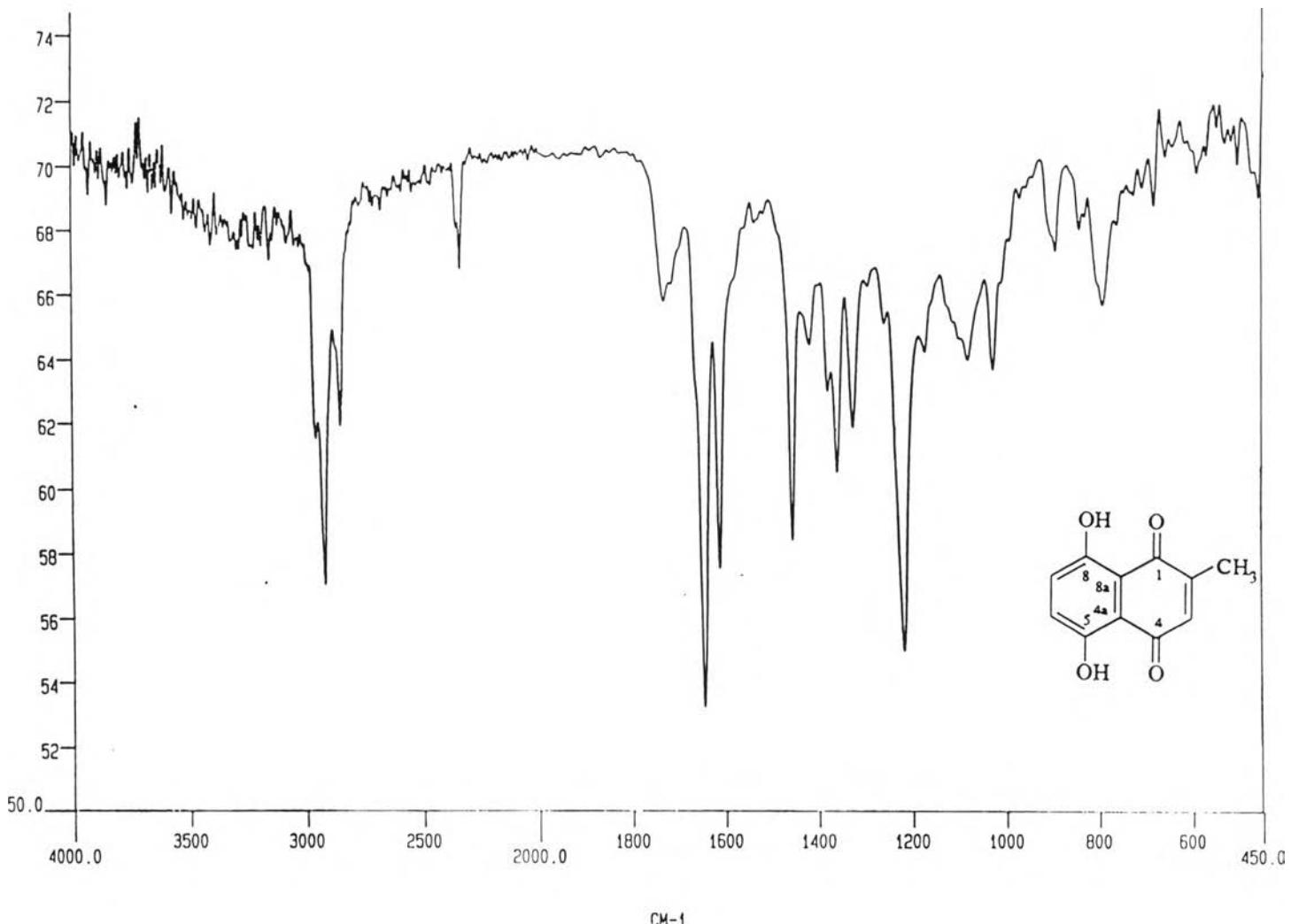


Figure 92 The IR spectrum of compound 104 (in KBr disc)

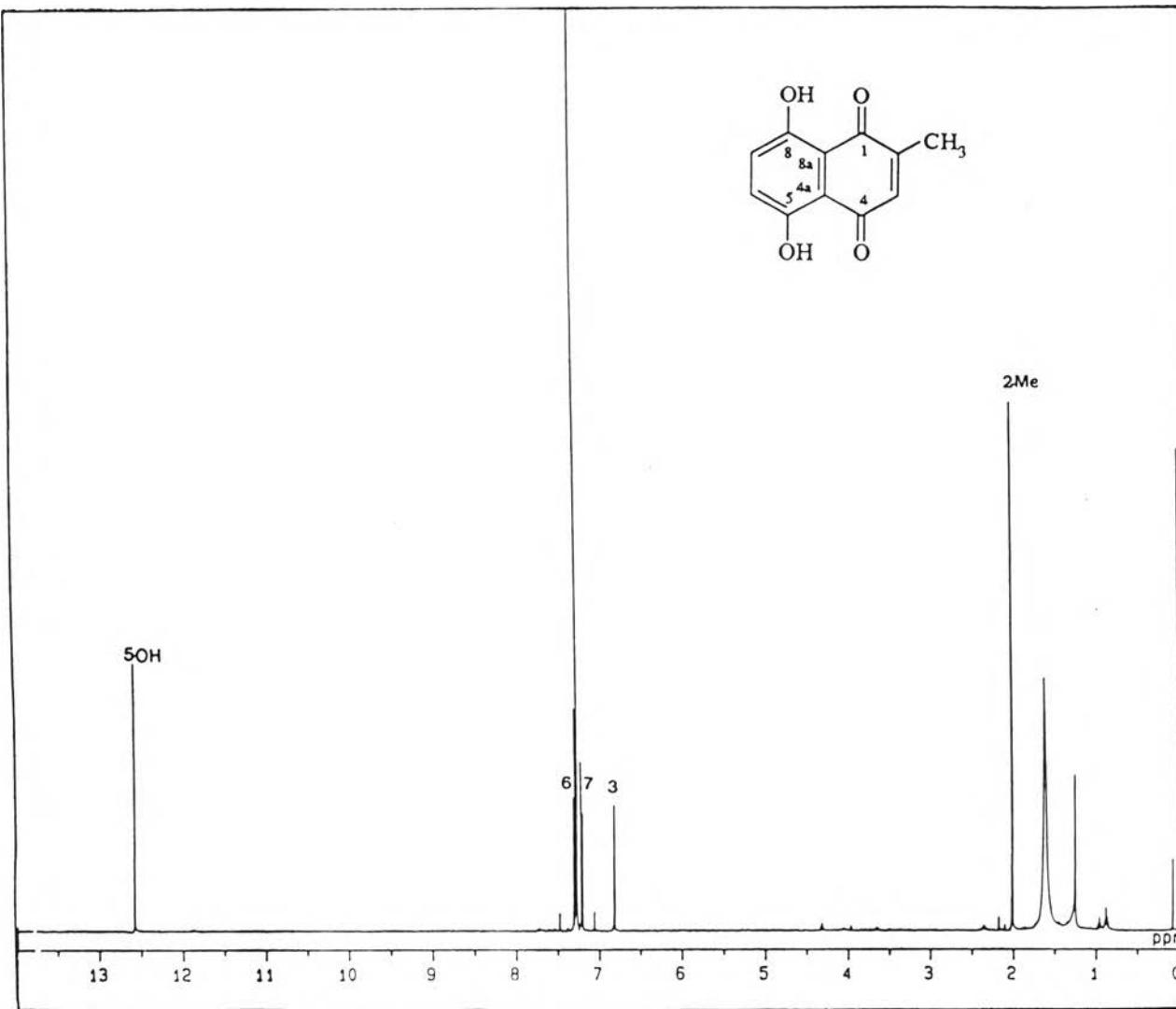


Figure 93 The ^1H NMR (500 MHz) spectrum of compound 104 (in CDCl_3)

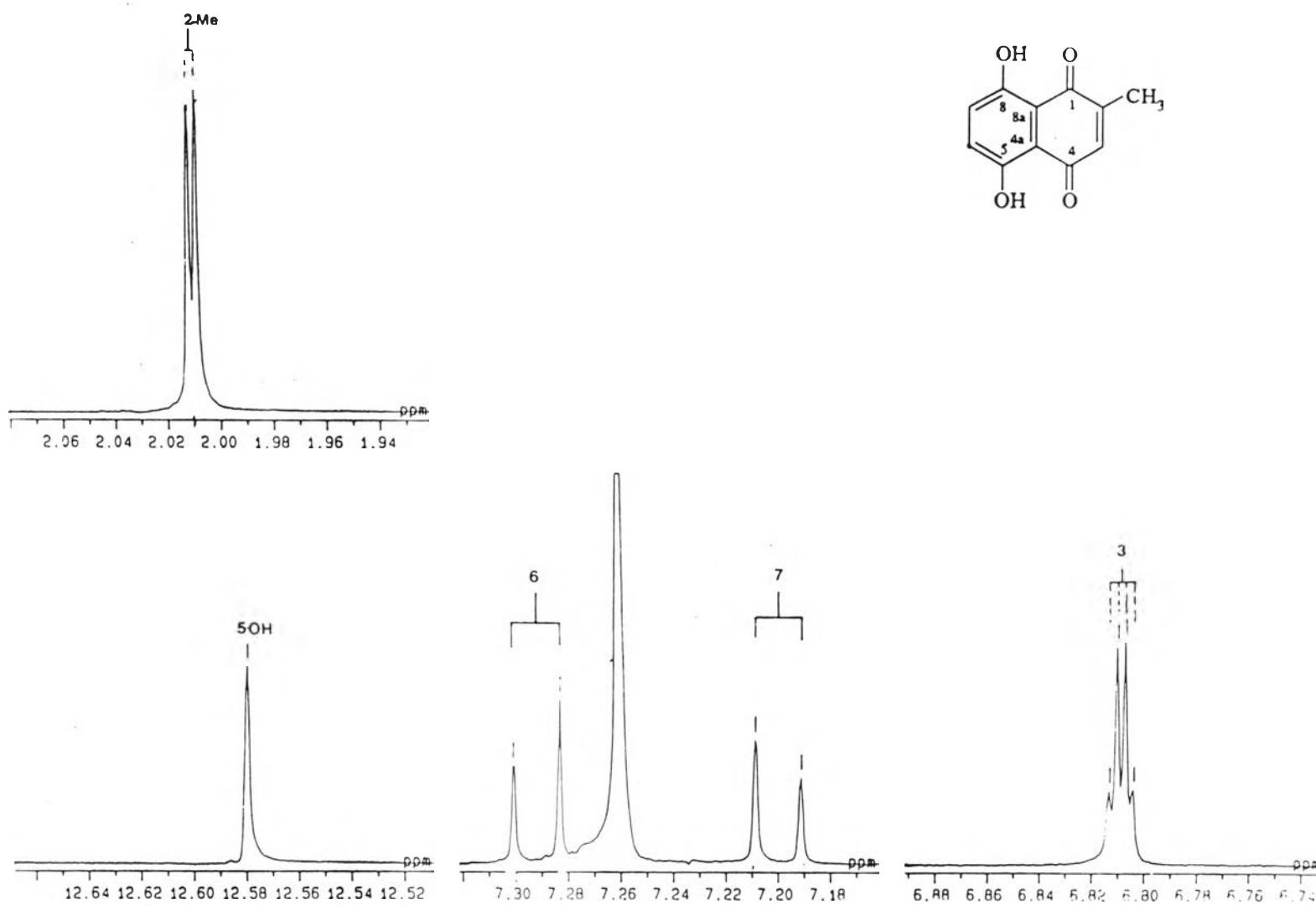
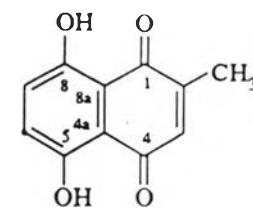


Figure 94 Expansion of the ¹H NMR (500 MHz) spectrum of compound 104

(in CDCl₃) : δ_H 1.94-12.64 ppm

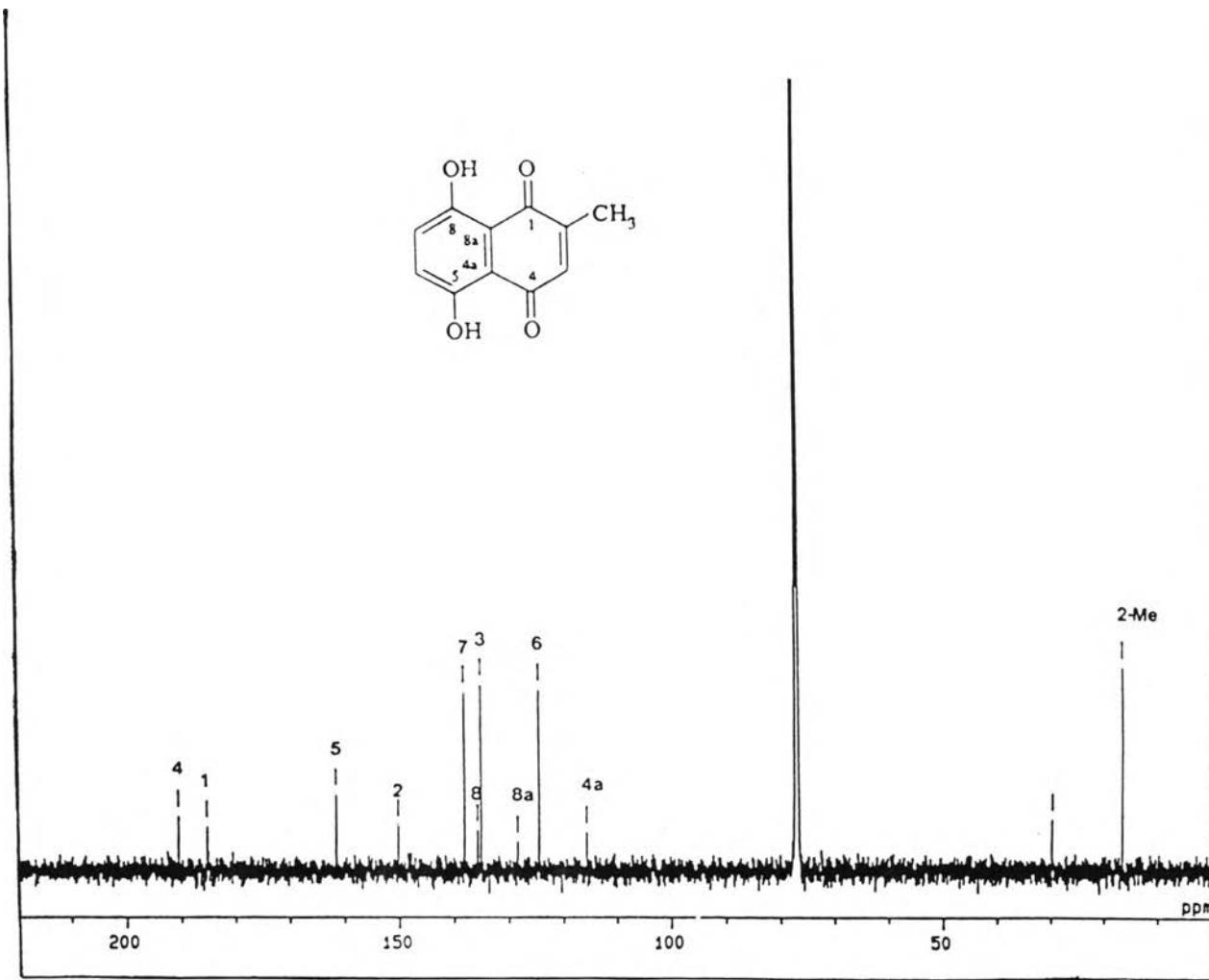


Figure 95 The ^{13}C NMR (125 MHz) spectrum of compound 104 (in CDCl_3)

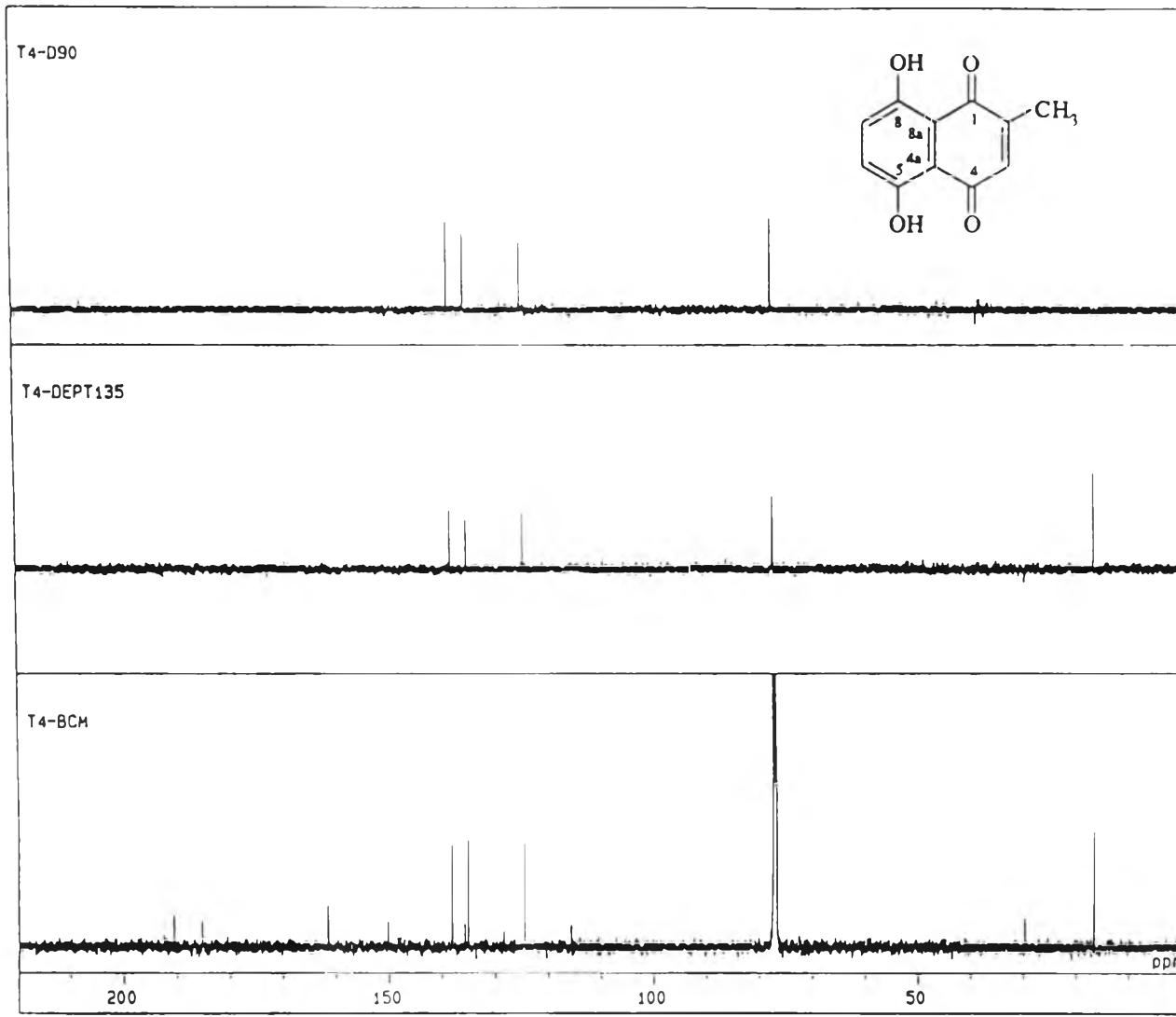


Figure 96 The DEPT (125 MHz) spectrum of compound 104 (in CDCl_3)

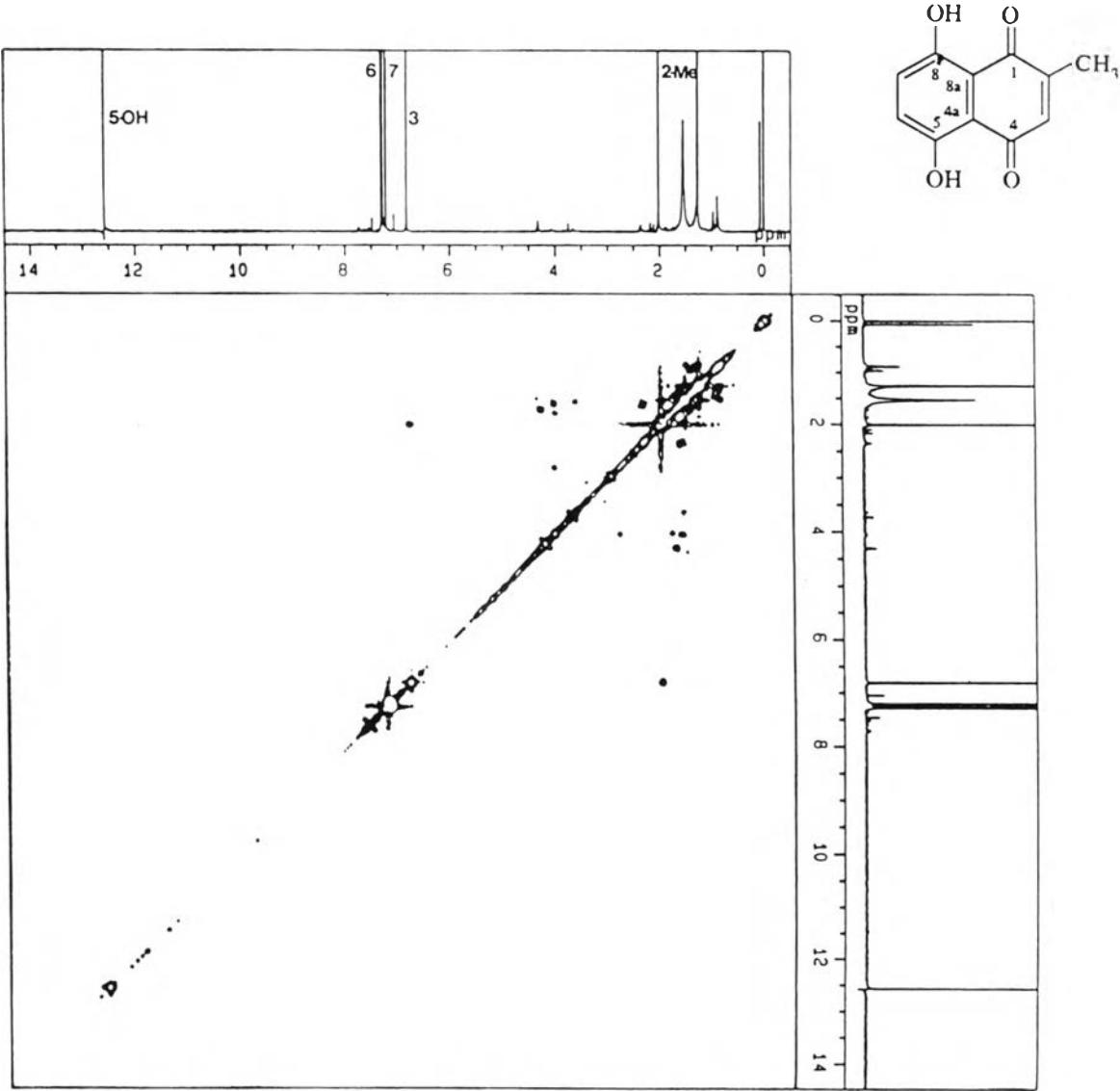


Figure 97 The ^1H - ^1H COSY (500 MHz) spectrum of compound 104 (in CDCl_3)

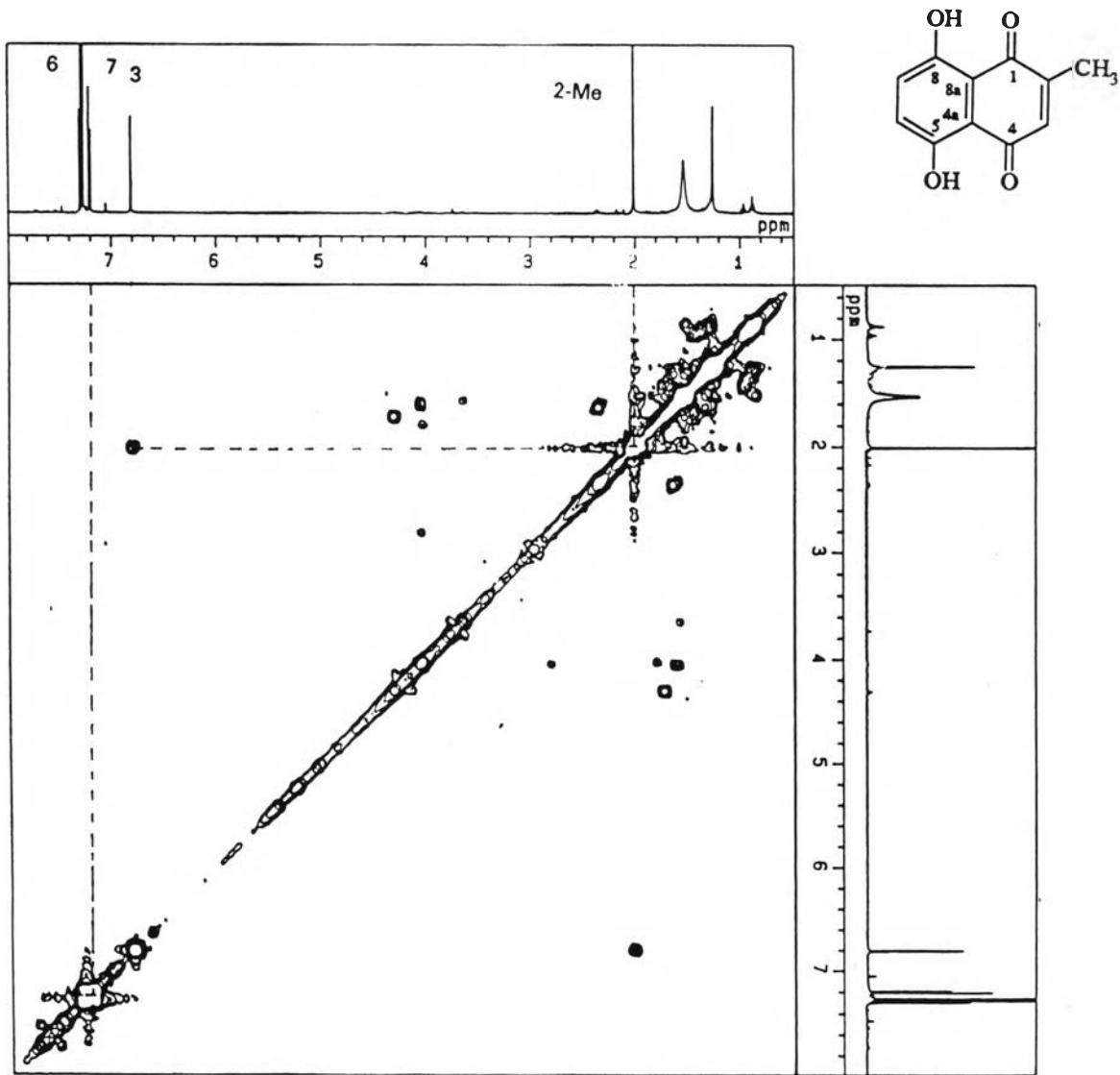


Figure 98 Expansion of the $^1\text{H}-^1\text{H}$ COSY (500 MHz) spectrum of compound 104

(in CDCl_3) : δ_{H} 0.60-7.80 ppm

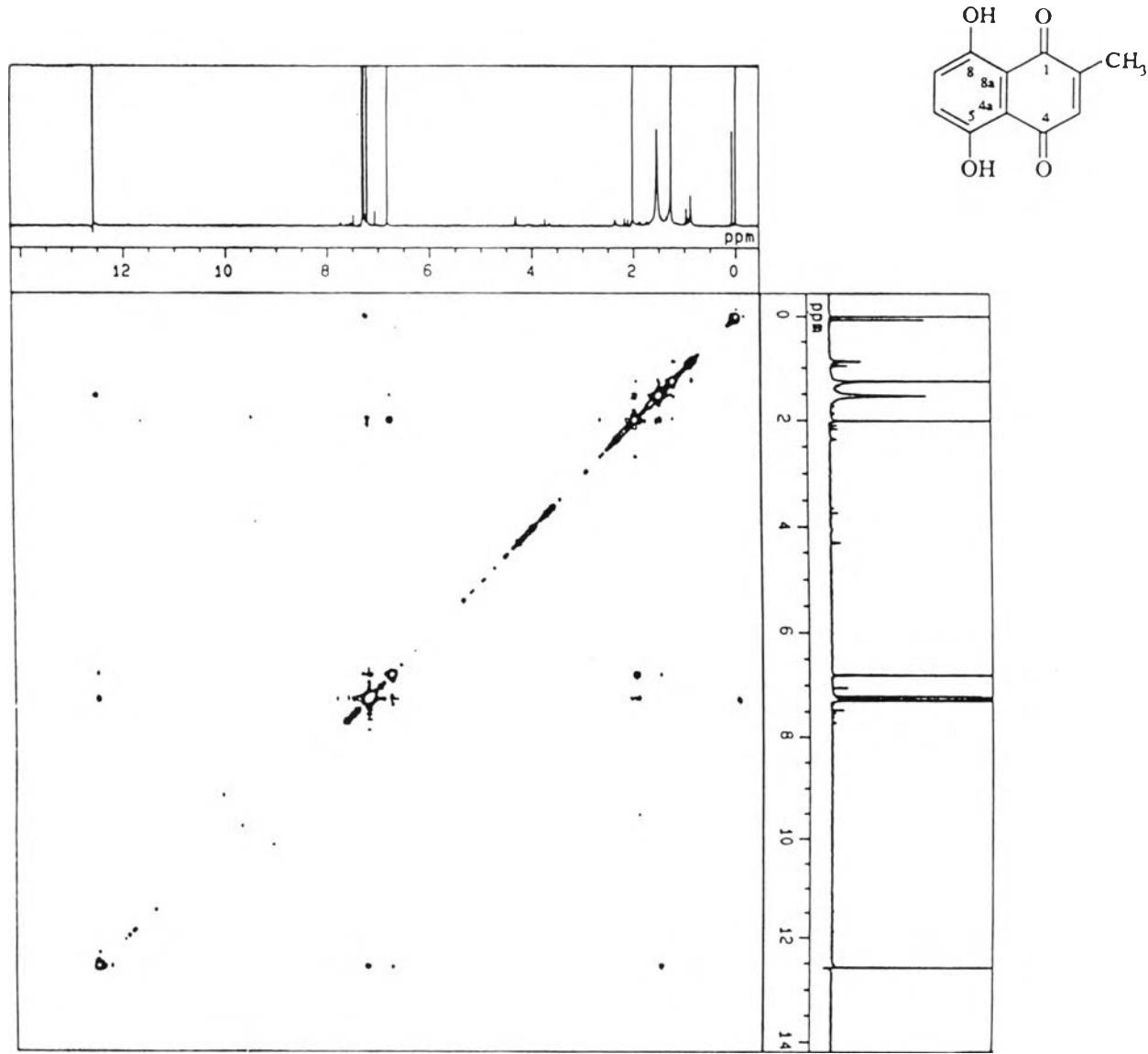


Figure 99 The NOESY (500 MHz) spectrum of compound 104 (in CDCl₃)

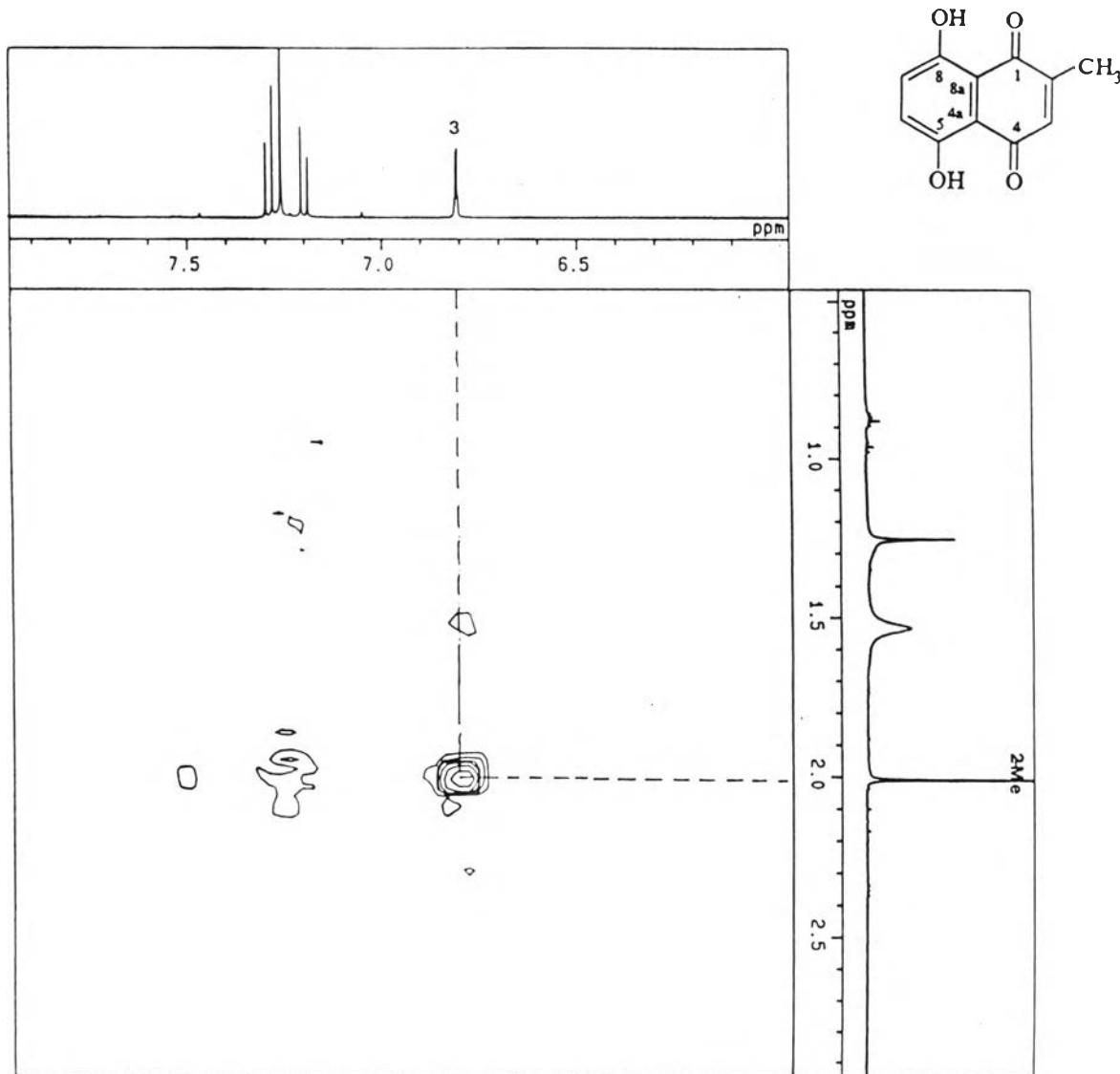


Figure 100 Expansion of the NOESY (500 MHz) spectrum of compound 104

(in CDCl₃) : δ_H 5.60-7.90 ppm

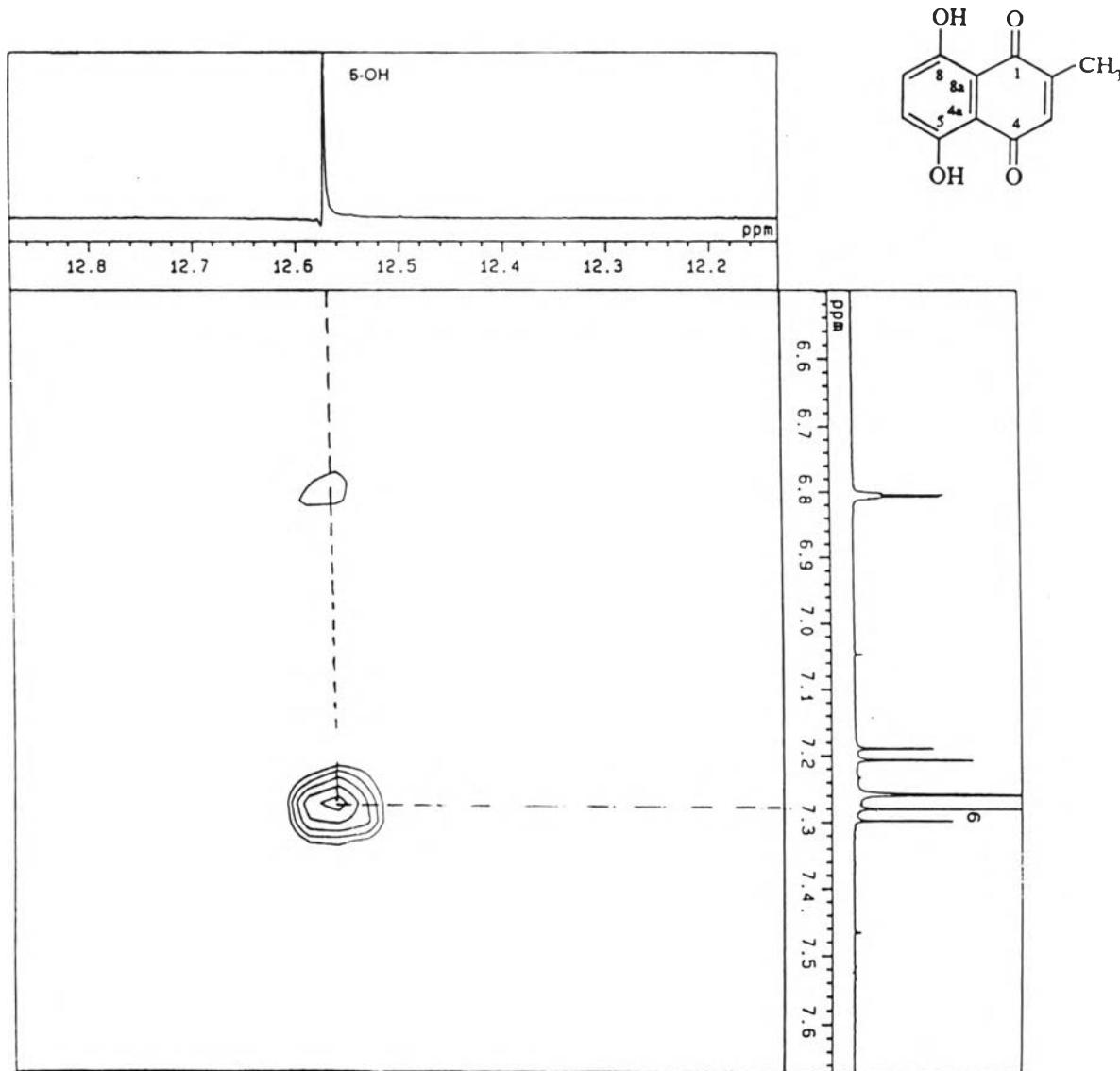
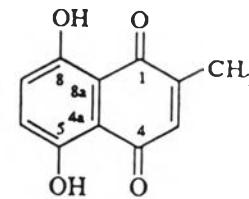


Figure 101 Expansion of the NOESY (500 MHz) spectrum of compound 104
(in CDCl_3) : δ_{H} 12.20-12.80 ppm



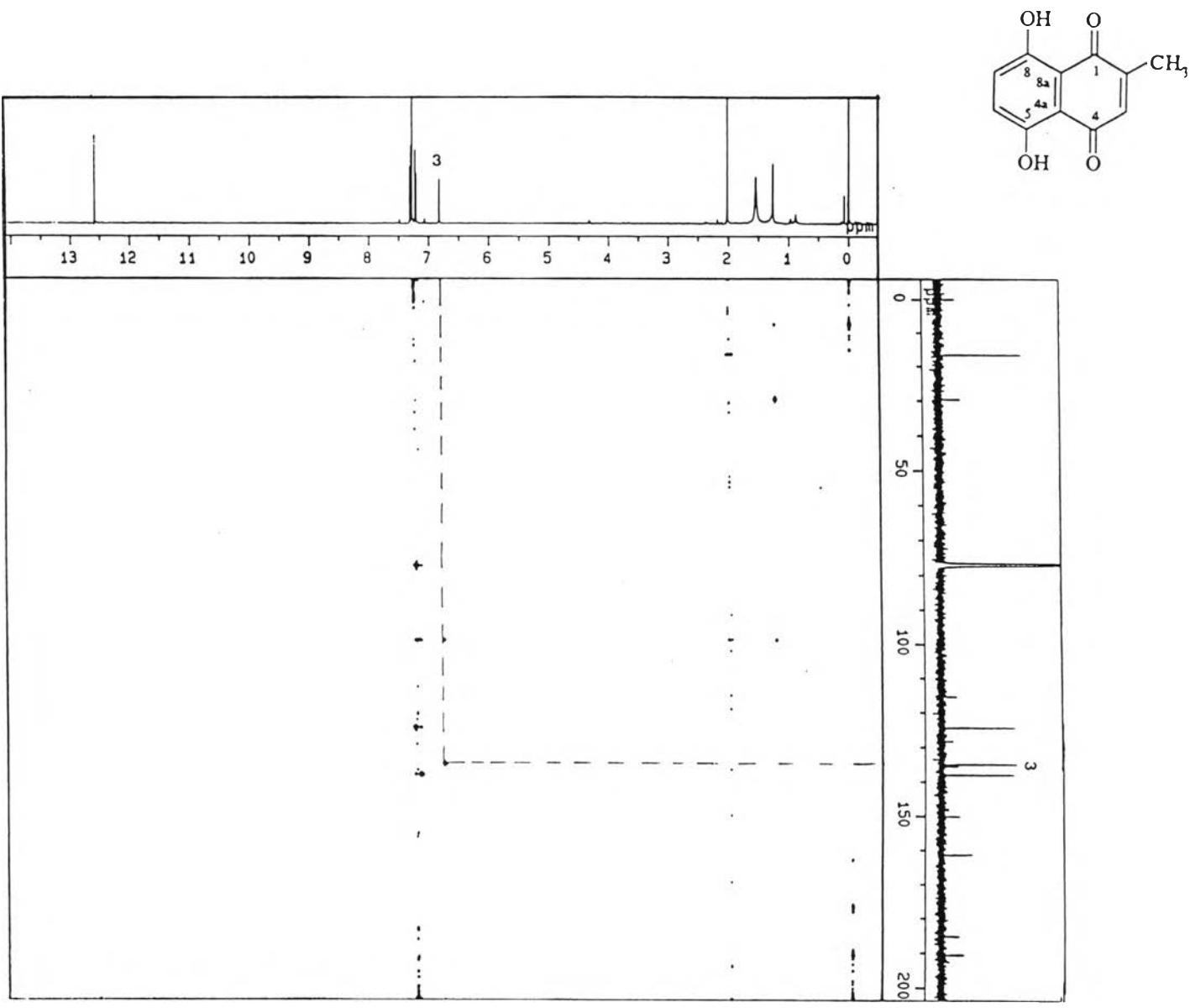


Figure 102 The HMQC spectrum of compound 104 (in CDCl₃)

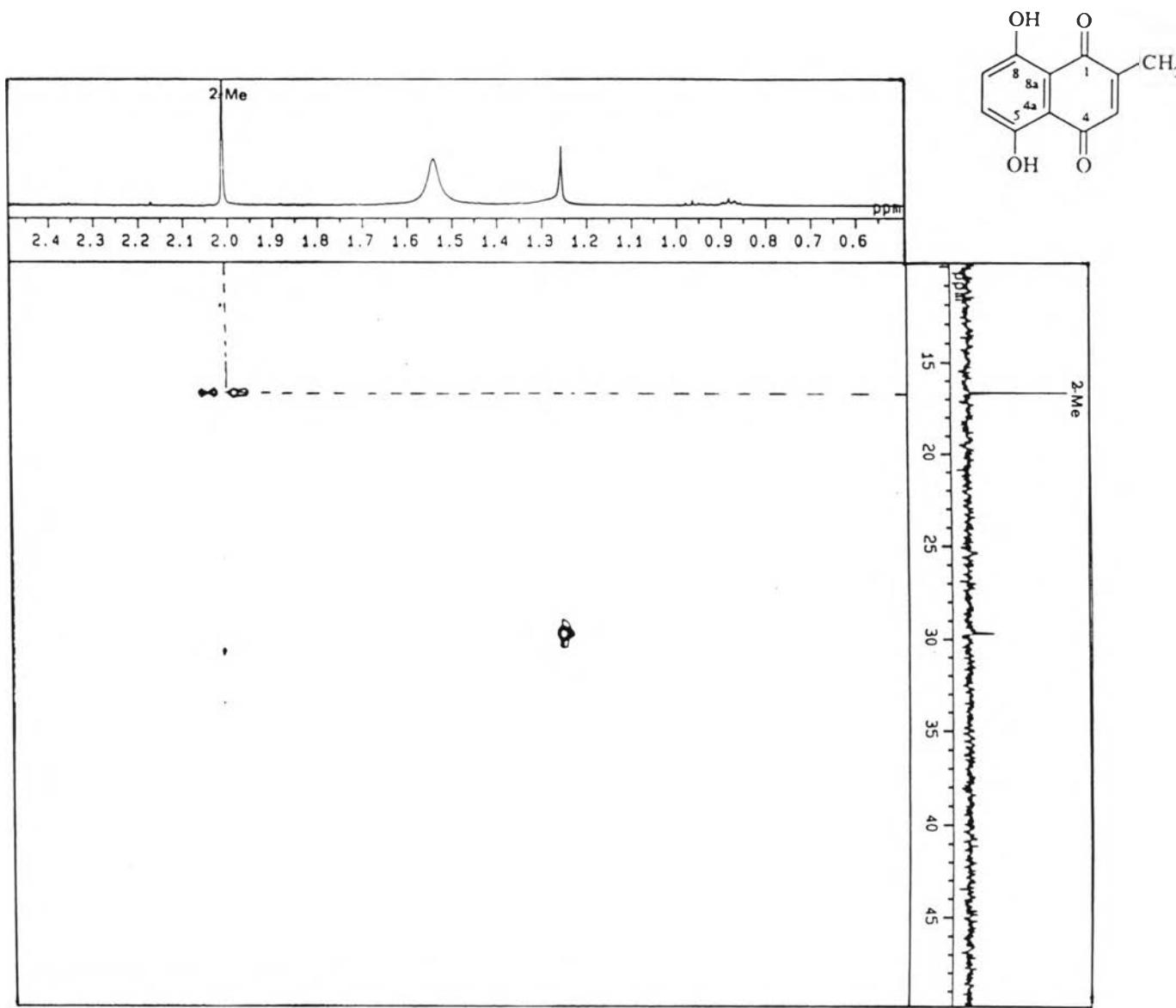


Figure 103 Expansion of the HMQC spectrum of compound **104** (in CDCl_3) :

δ_{H} 0.60-2.40 ; δ_{C} 10.00-45.00 ppm

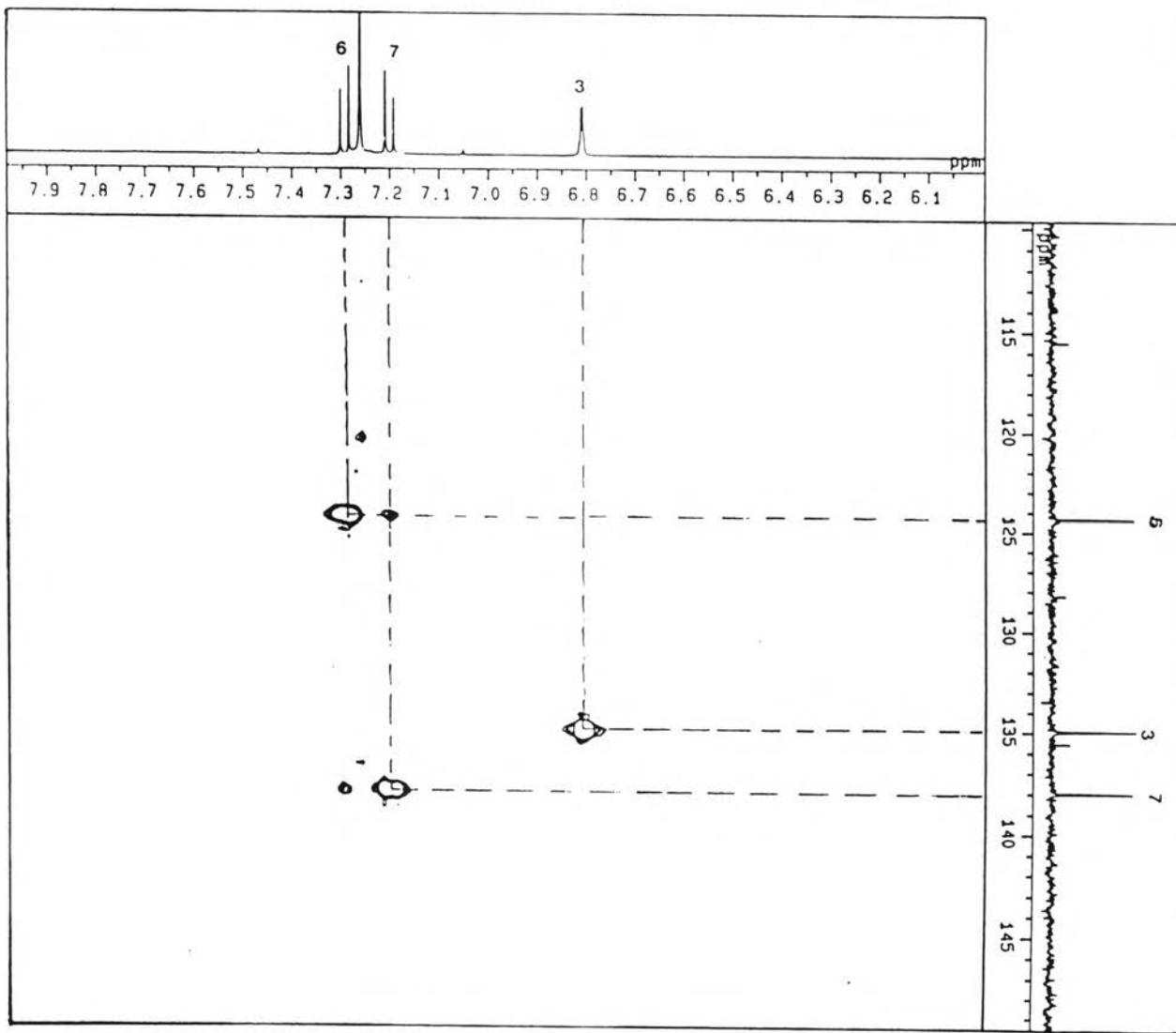
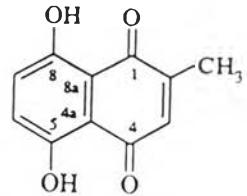


Figure 104 Expansion of the HMQC spectrum of compound 104 (in CDCl₃) :
 δ_{H} 6.10-7.90 ; δ_{C} 110.00-145.00 ppm

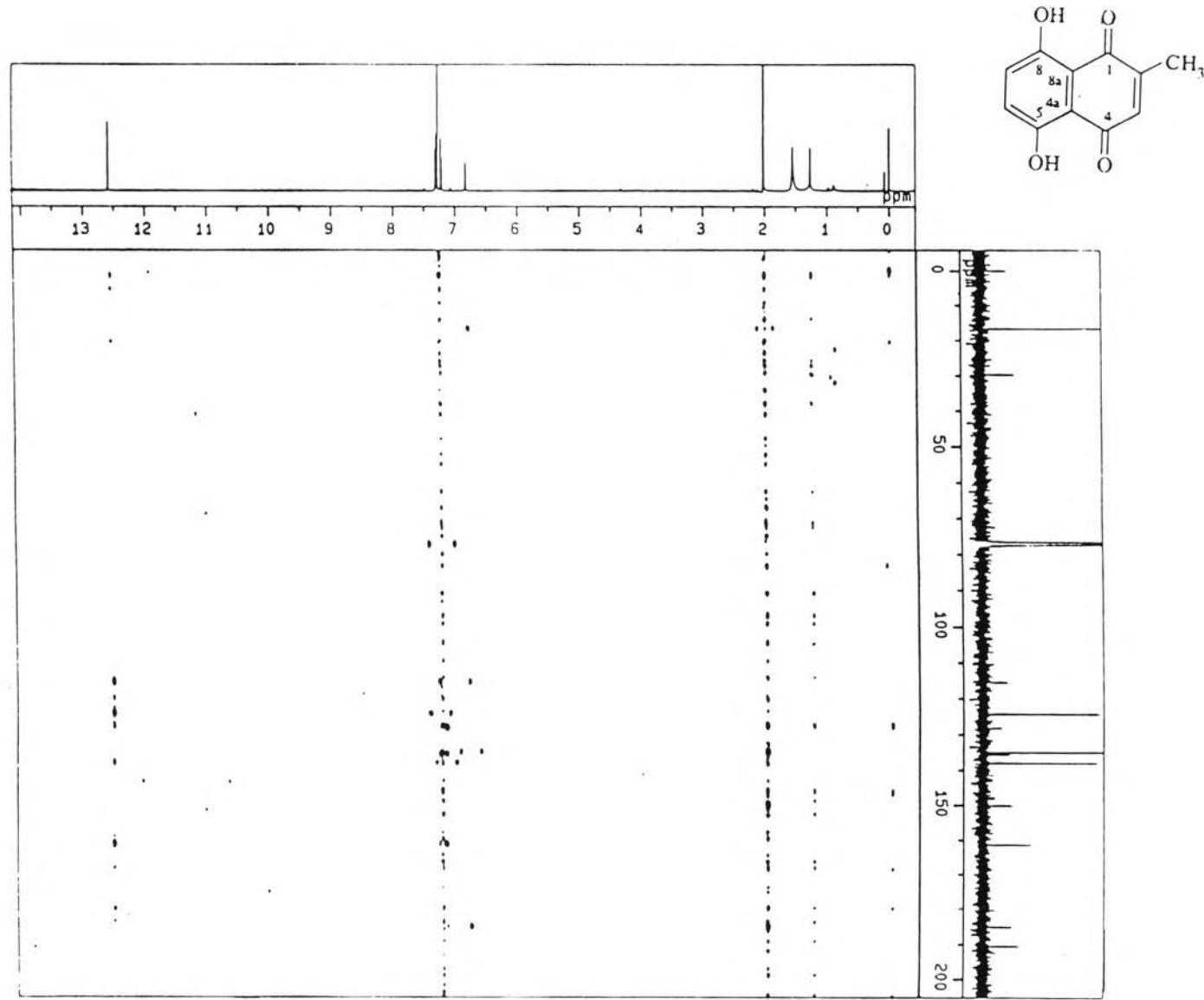


Figure 105 The HMBC spectrum of compound 104 (in CDCl₃)

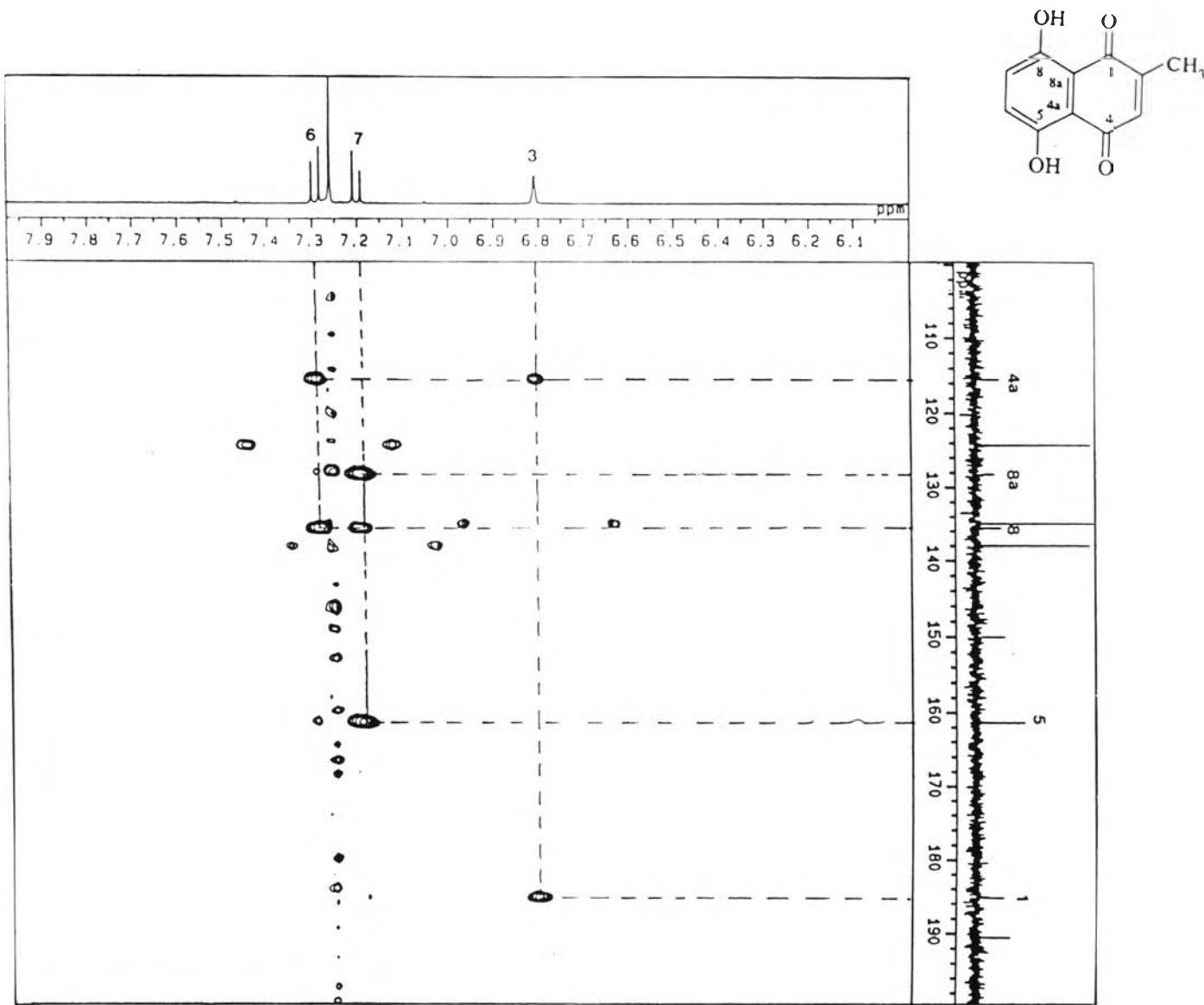


Figure 106 Expansion of the HMBC spectrum of compound 104 (in CDCl_3) :

δ_{H} 6.10-7.90 ; δ_{C} 105.00-194.00 ppm

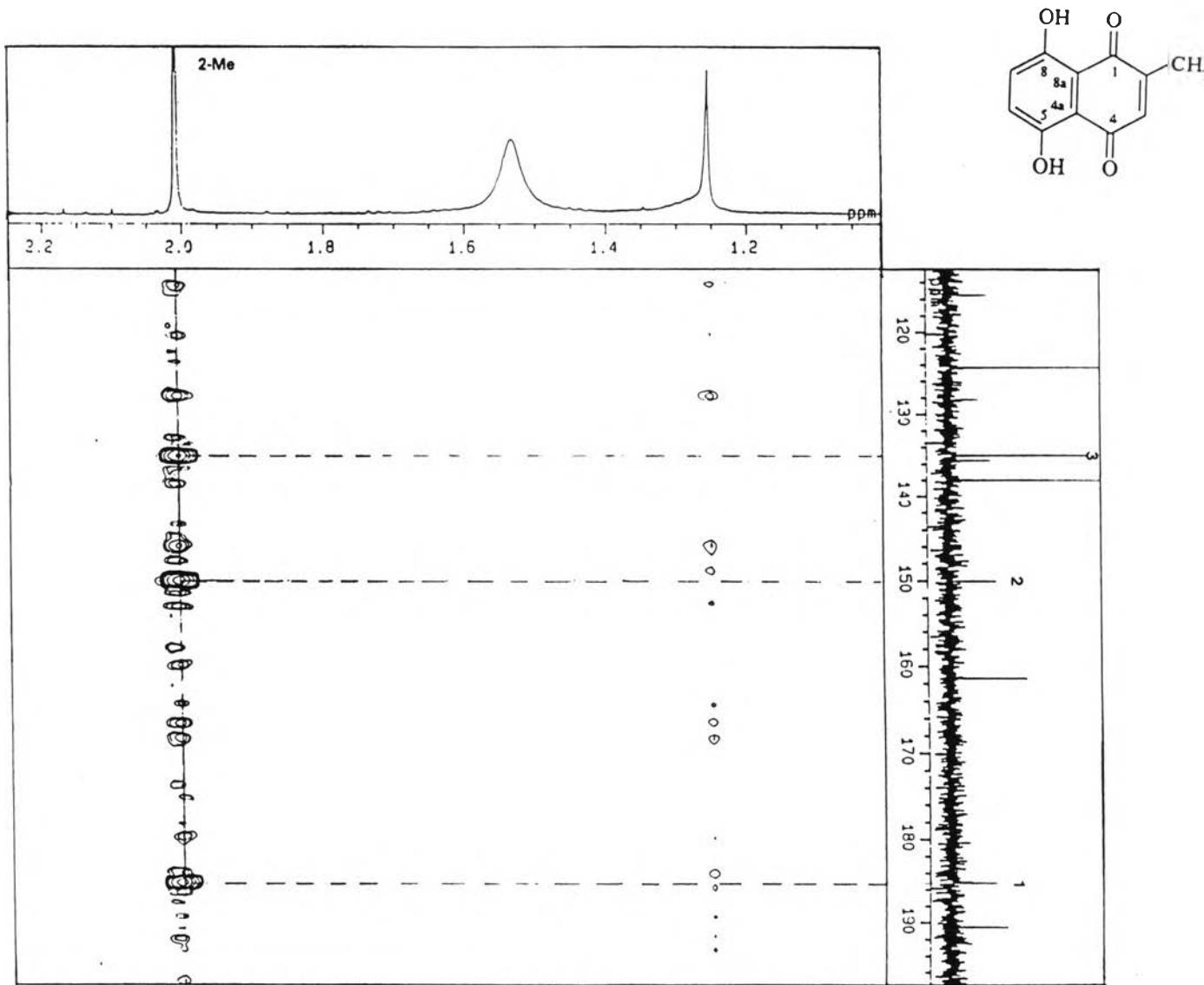


Figure 107 Expansion of the HMBC spectrum of compound 104(in CDCl_3) :

δ_{H} 1.30-2.20 ; δ_{C} 117.00-193.00 ppm

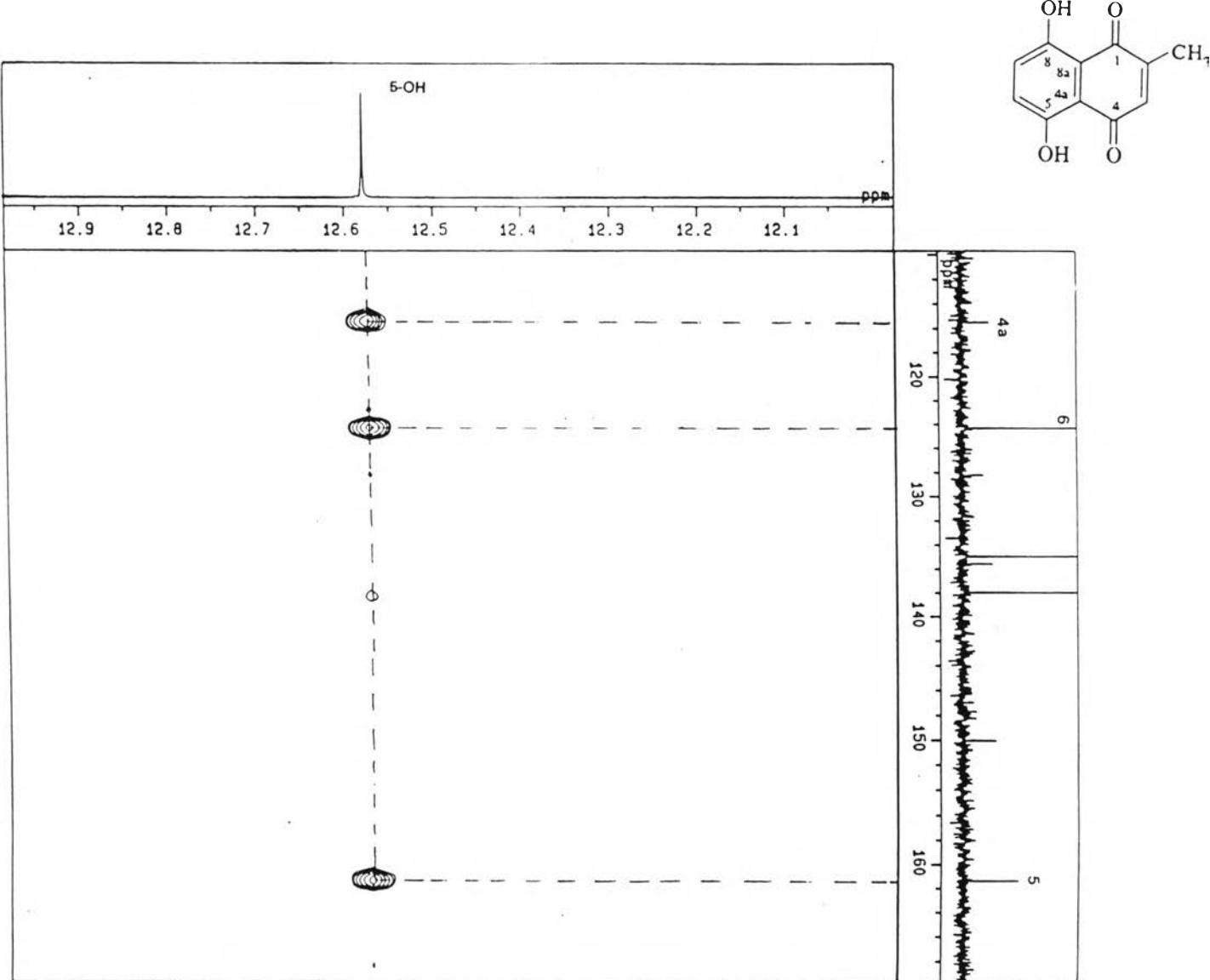


Figure 108 Expansion of the HMBC spectrum of compound 104 (in CDCl_3) :

δ_{H} 1.21-12.90 ; δ_{C} 110.00-168.00 ppm

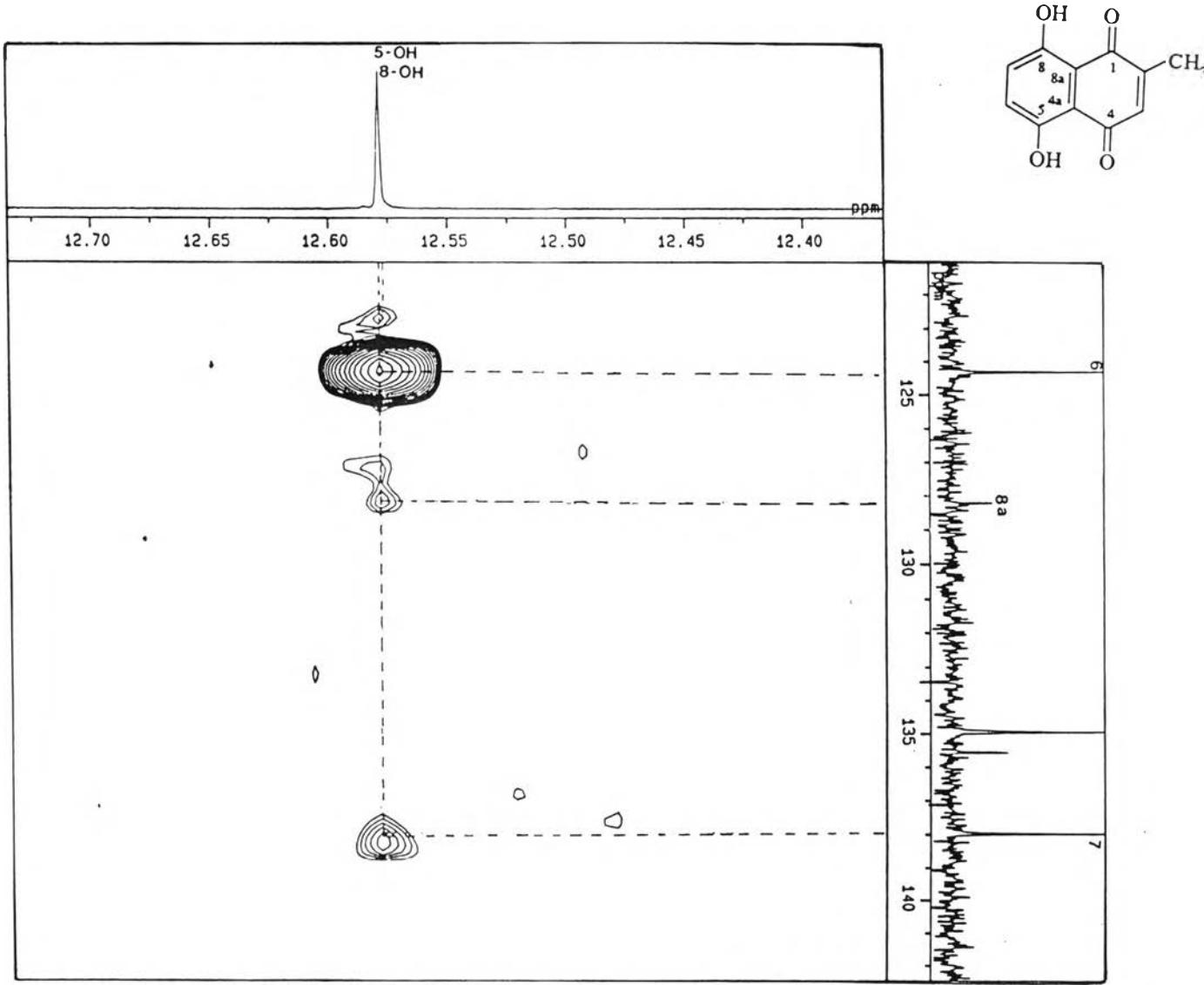


Figure 109 Expansion of the HMBC spectrum of compound 104 (in CDCl_3) :
 δ_{H} 12.40-12.70 ; δ_{C} 122.00-142.00 ppm

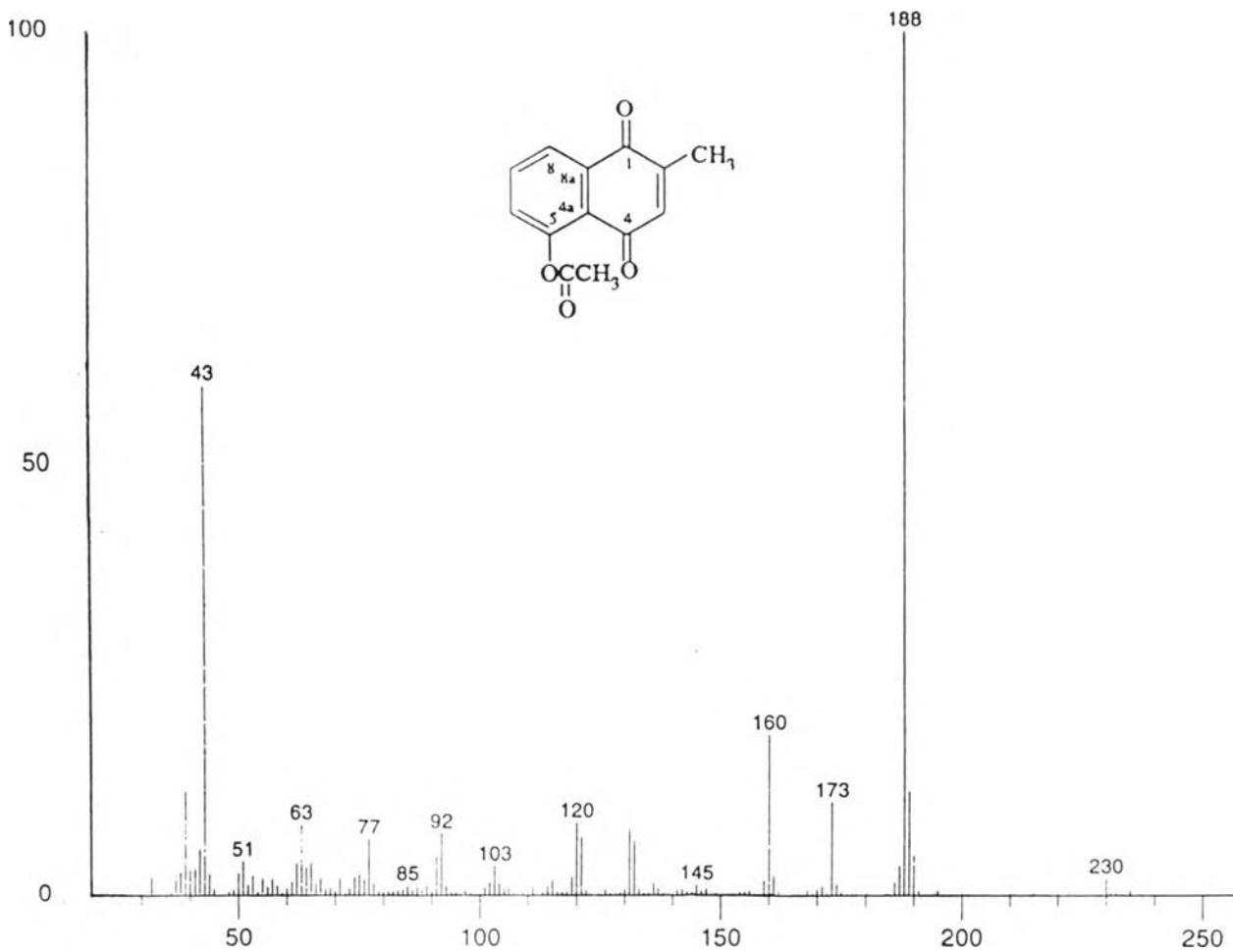


Figure 110 The EI mass spectrum of compound 105

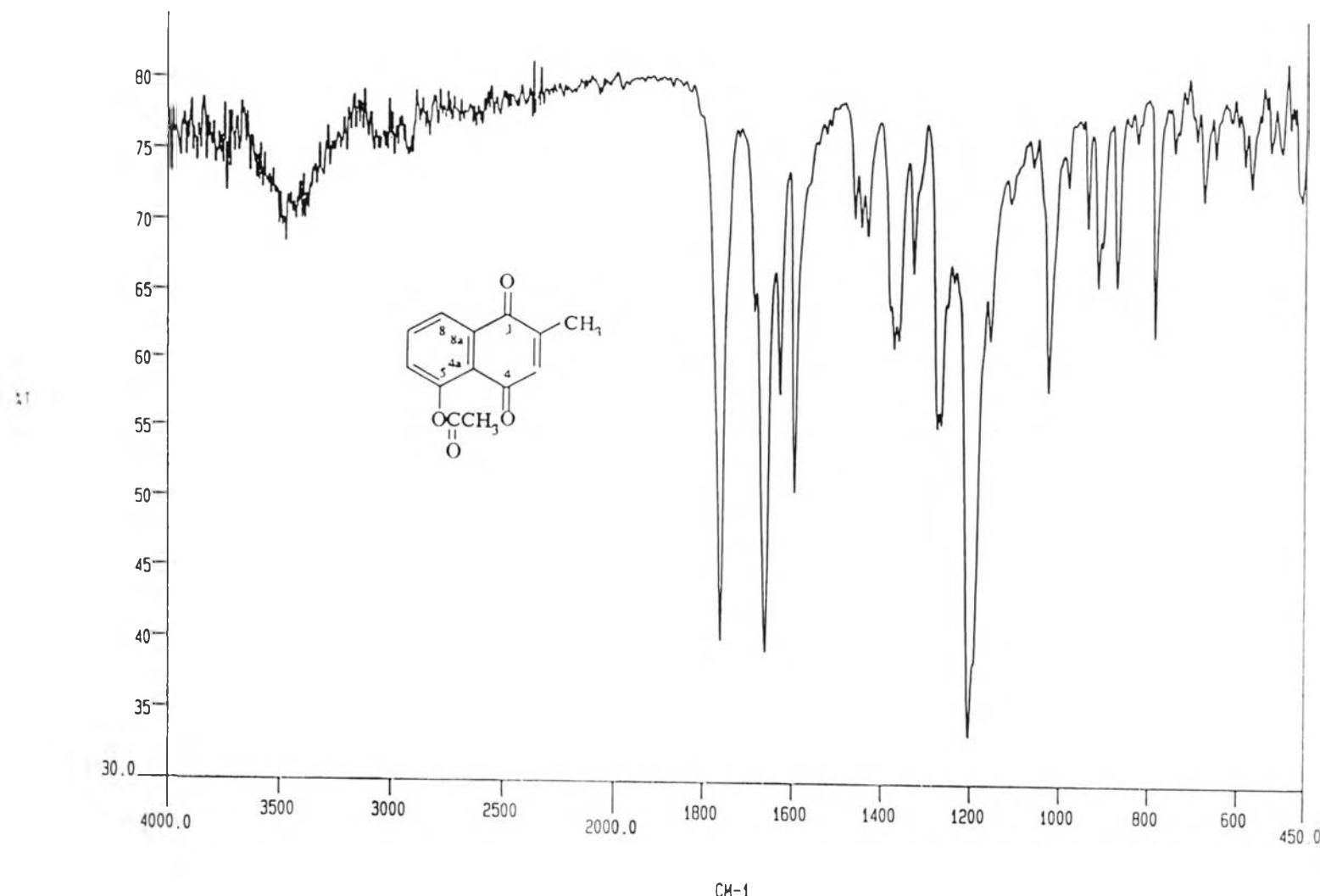


Figure 111 The IR spectrum of compound 105 (in KBr disc)

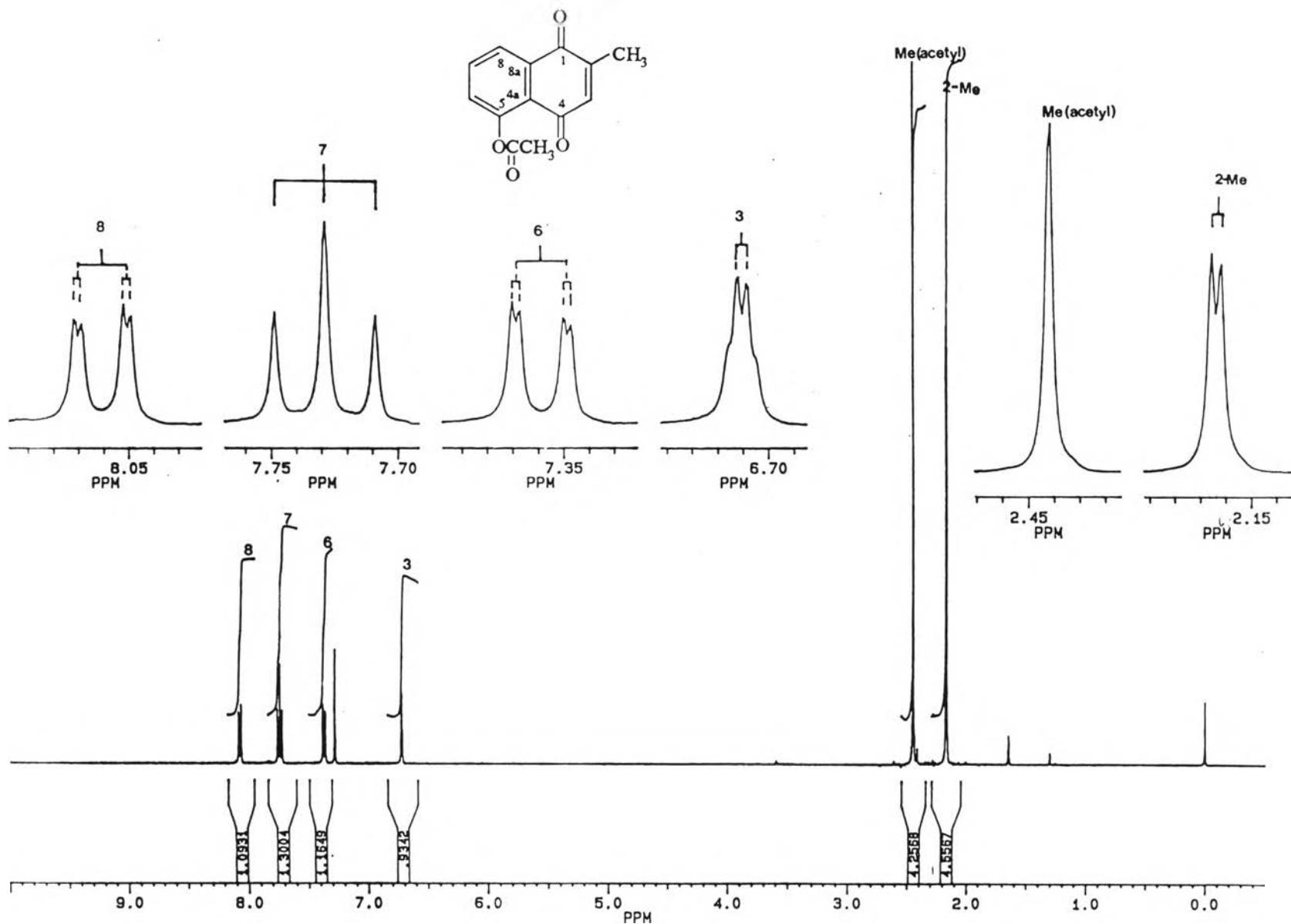


Figure 112 The ^1H NMR (400 MHz) spectrum of compound 105 (in CDCl_3)

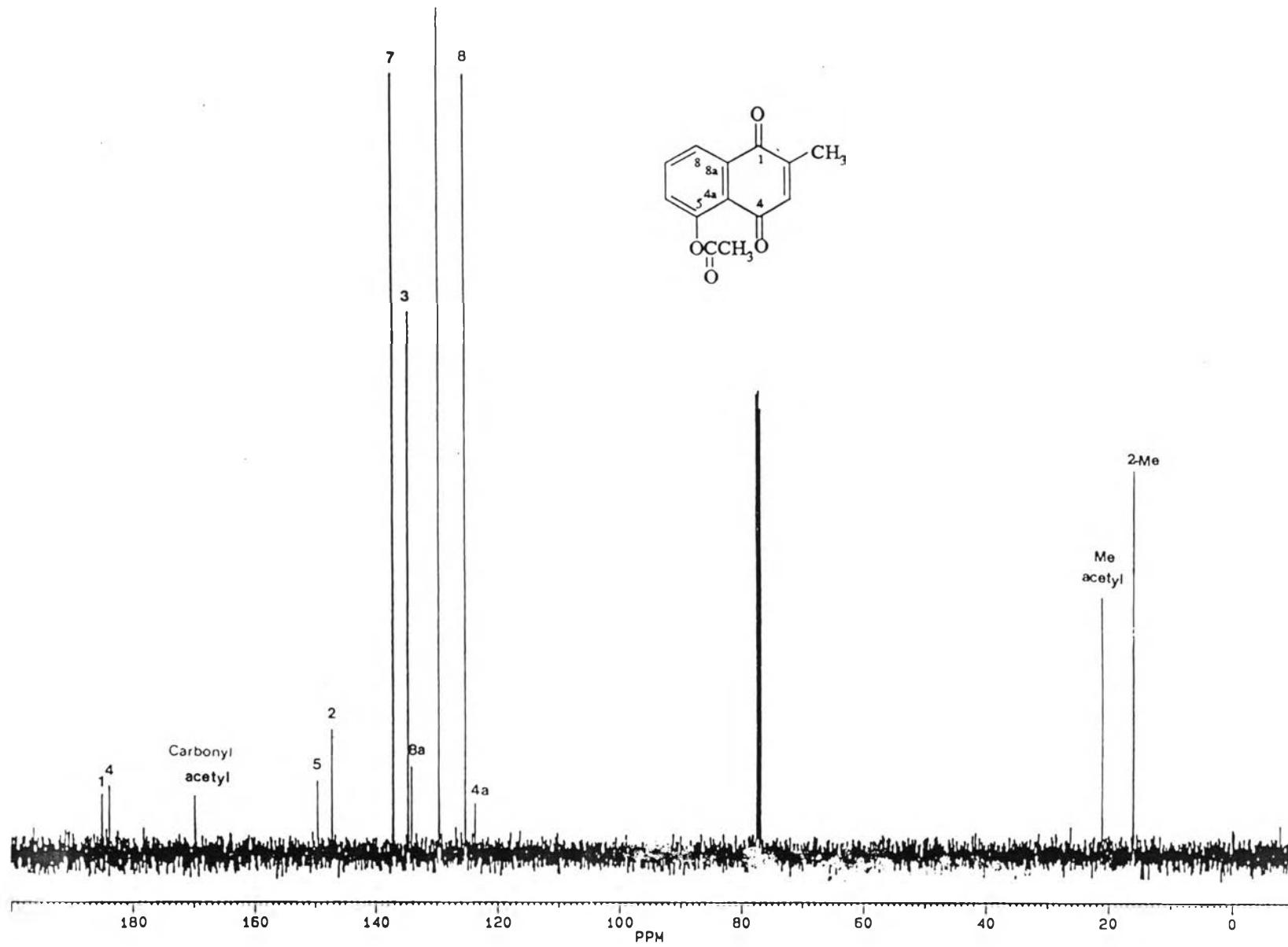


Figure 113 The ^{13}C NMR (100 MHz) spectrum of compound 105 (in CDCl_3)

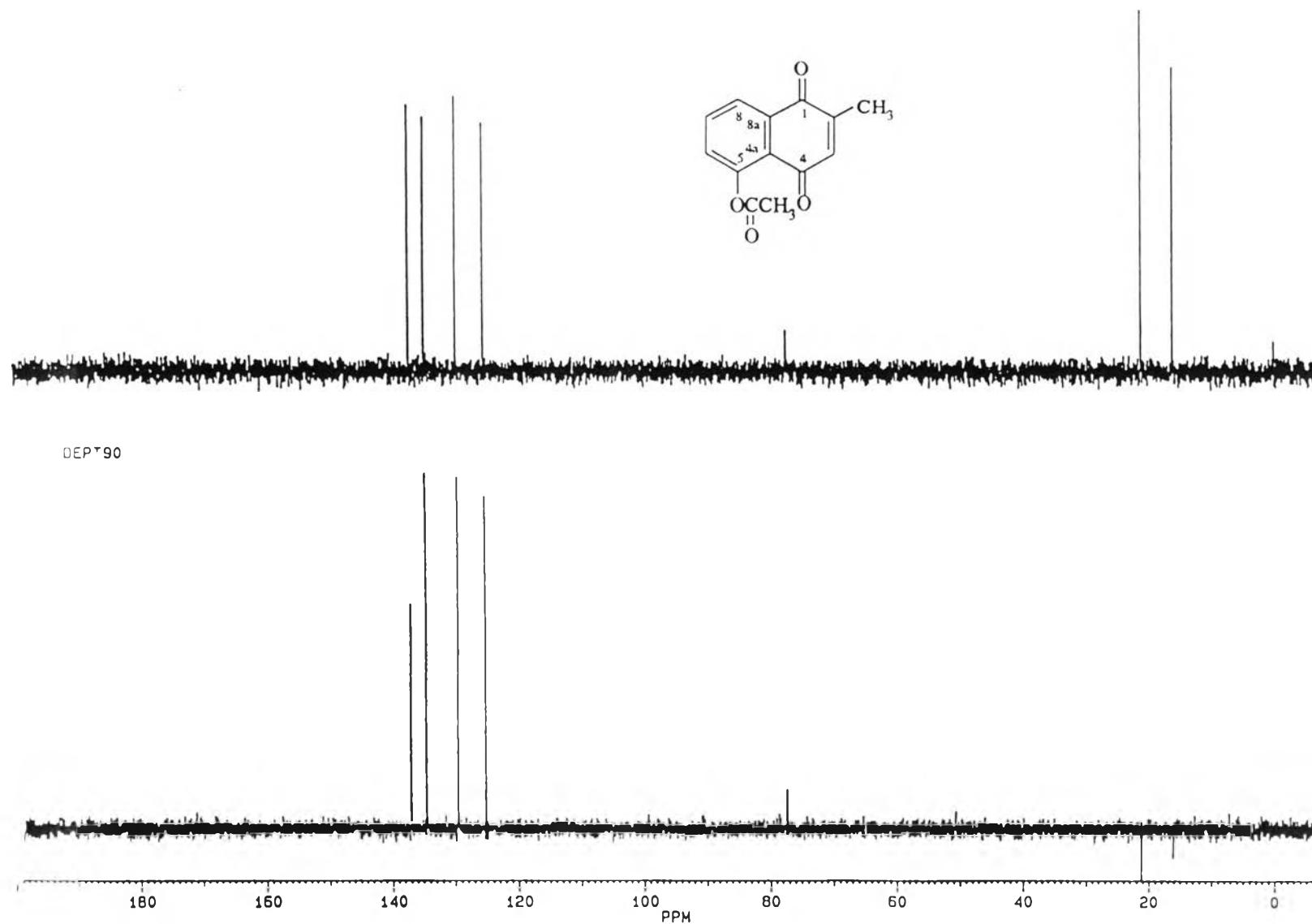


Figure 114 The DEPT (100 MHz) spectrum of compound 105 (in CDCl_3)

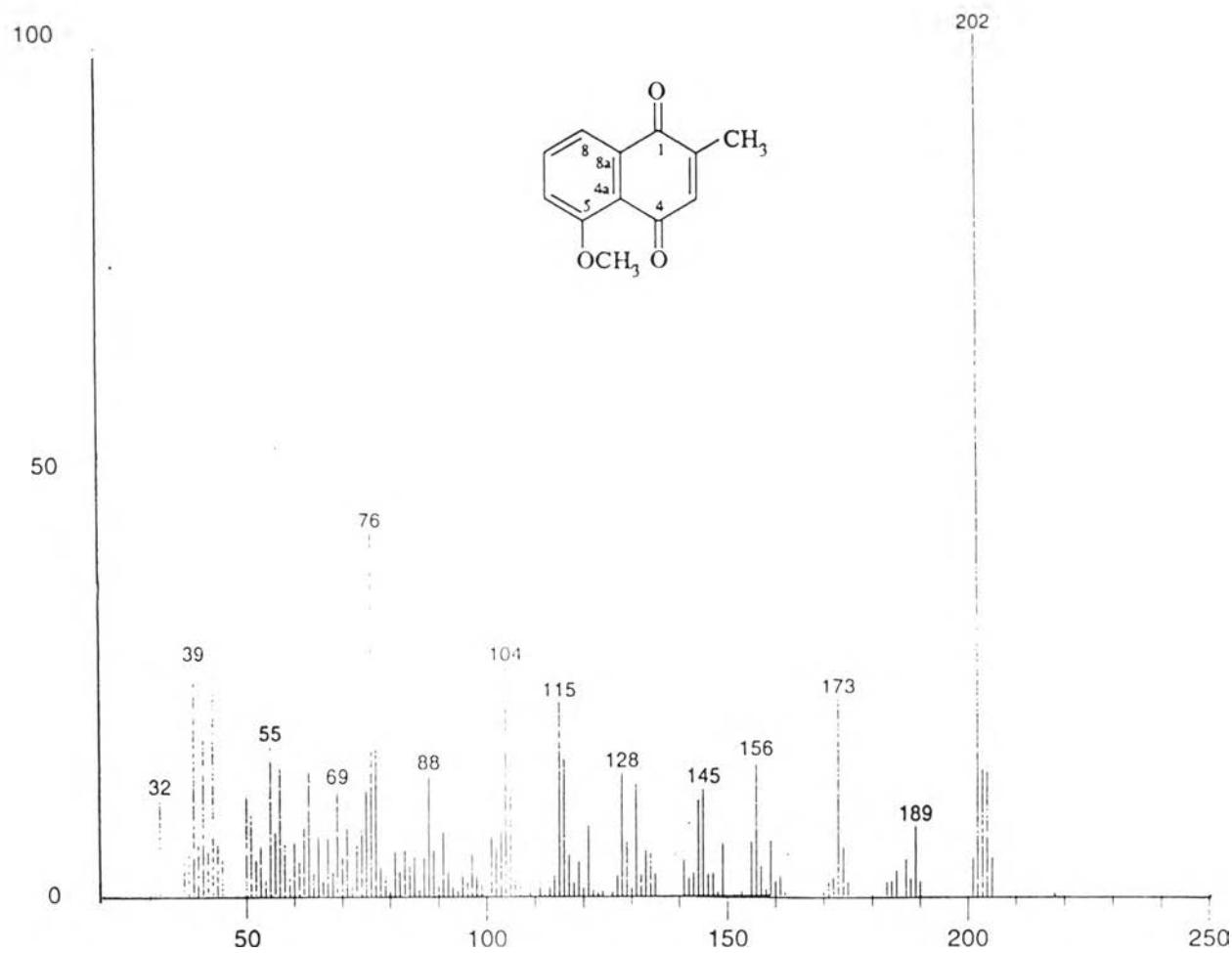


Figure 115 The EI mass spectrum of compound 106

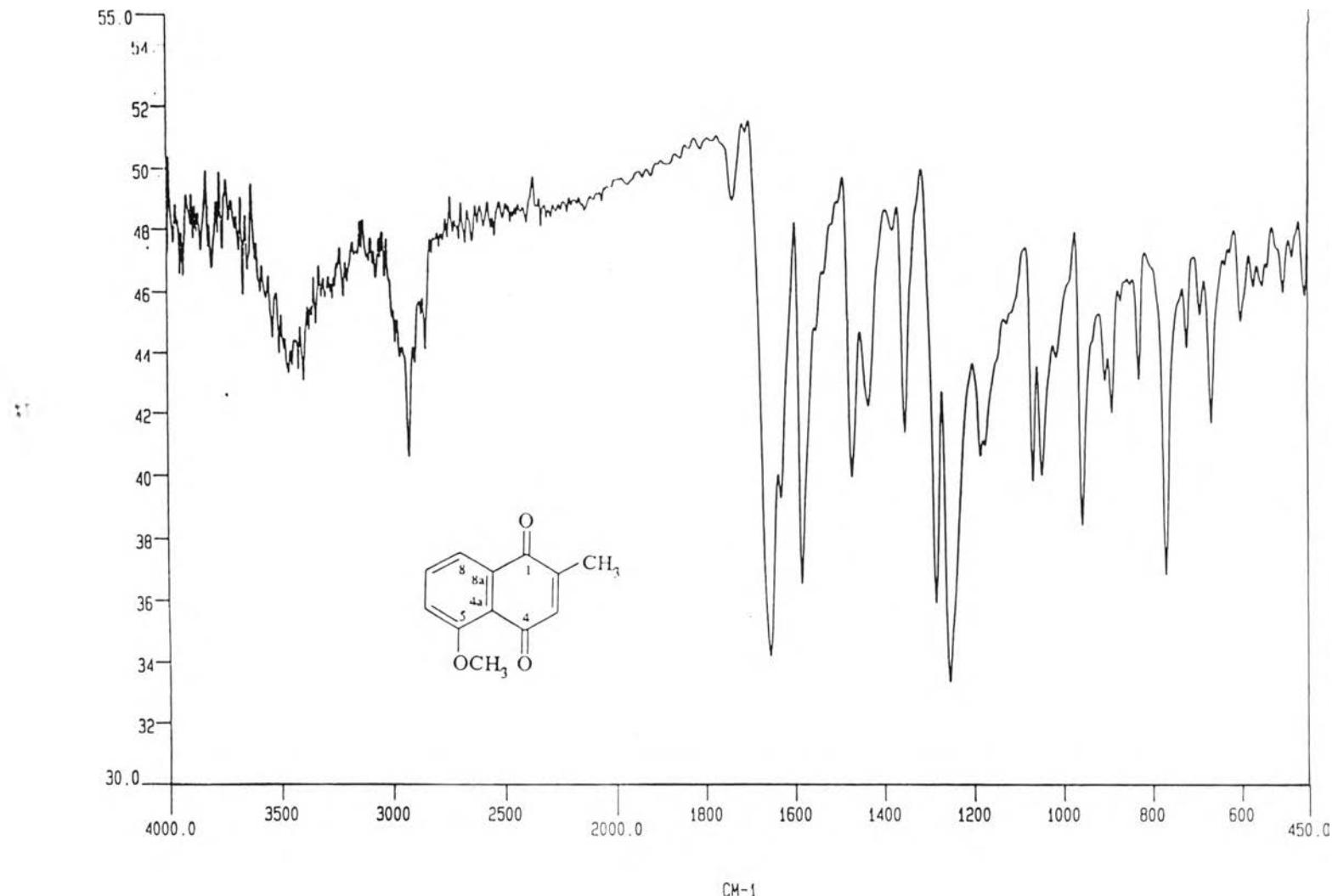


Figure 116 The IR spectrum of compound 106 (in KBr disc)

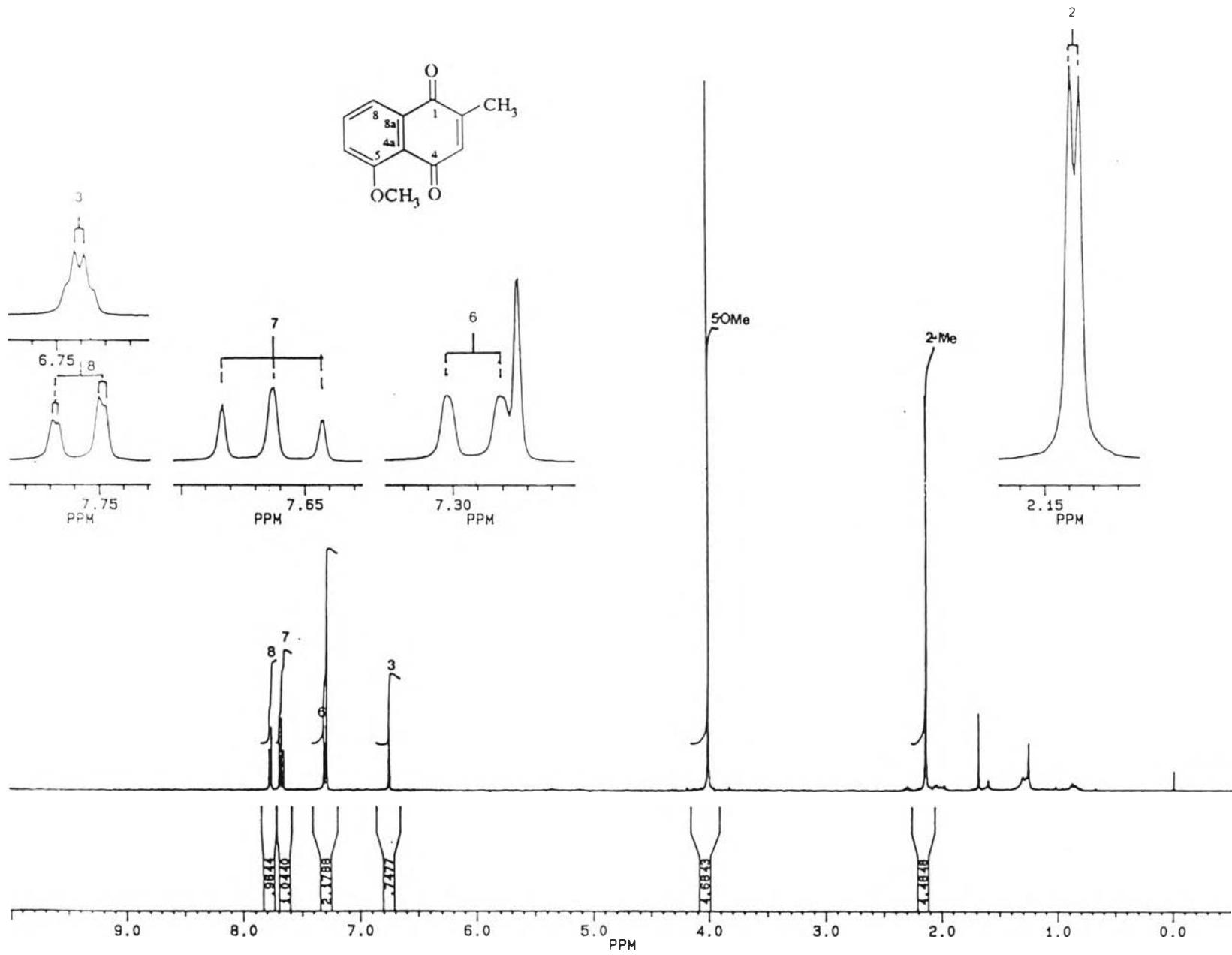


Figure 117 The ^1H NMR (400 MHz) spectrum of compound 106 (in CDCl_3)

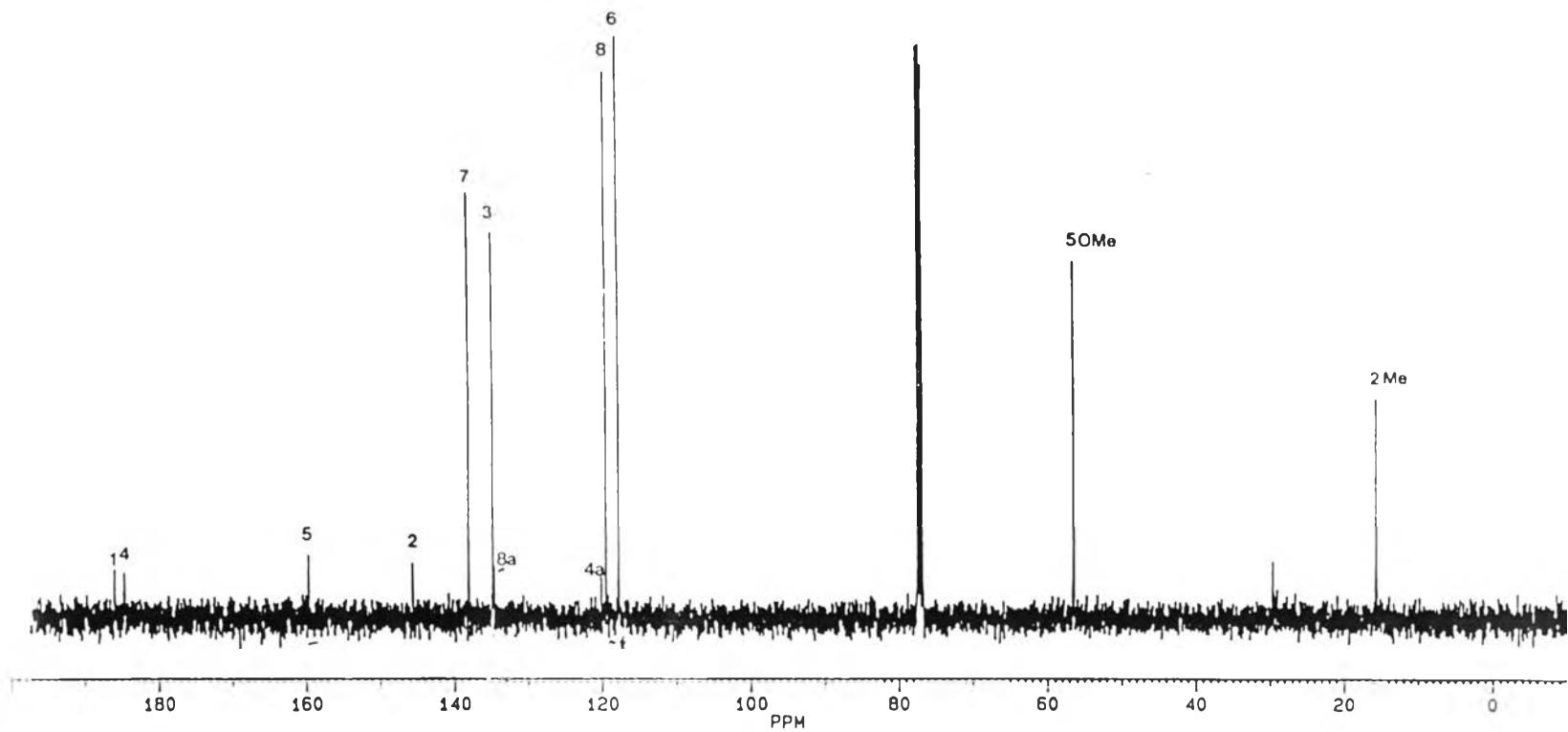
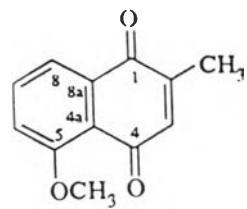
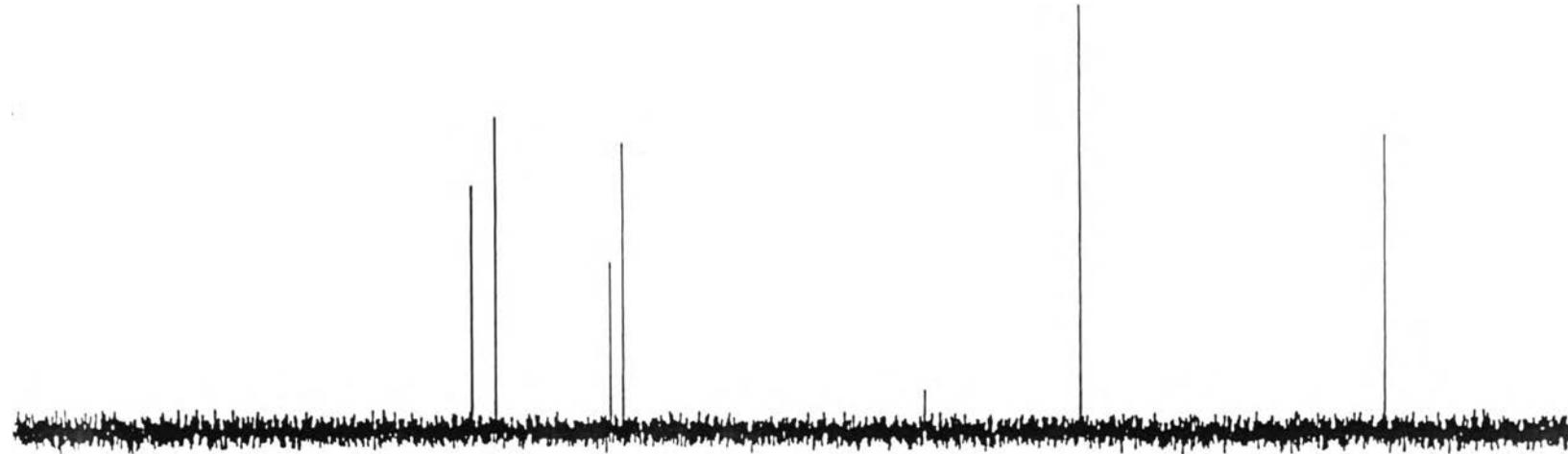


Figure 118 The ¹³C NMR (100 MHz) spectrum of compound 106 (in CDCl₃)

PH DEPT 135



PH DEPT 90

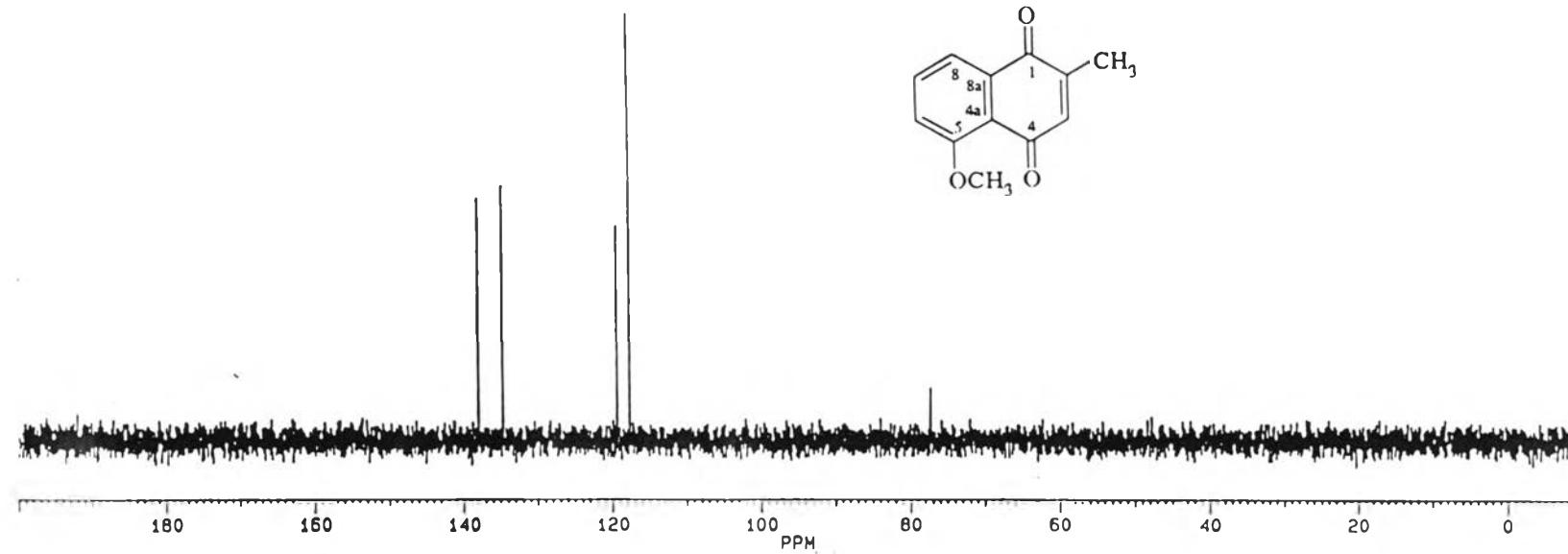


Figure 119 The DEPT (100 MHz) spectrum of compound 106 (in CDCl₃)

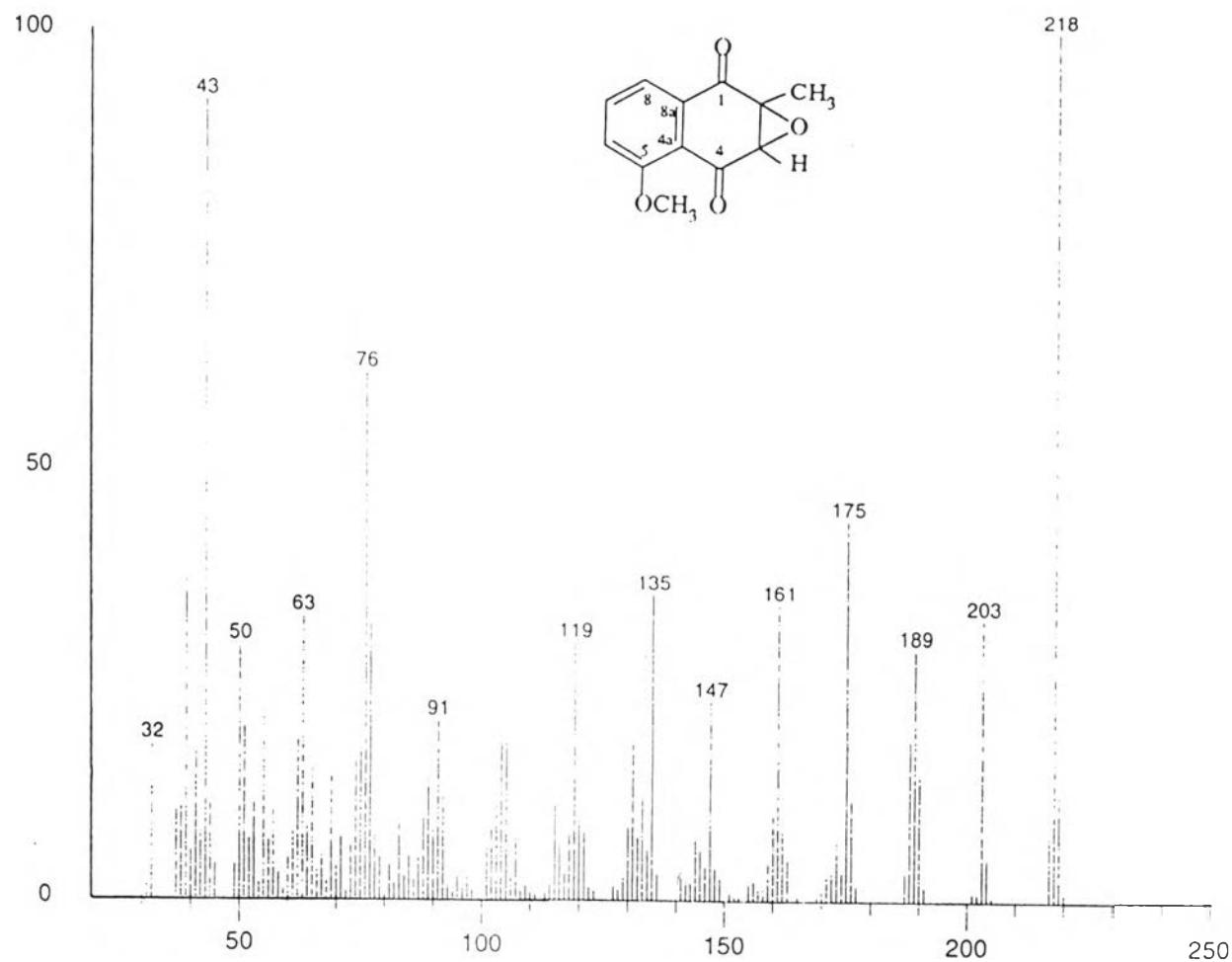


Figure 120 The EIMS spectrum of compound 107

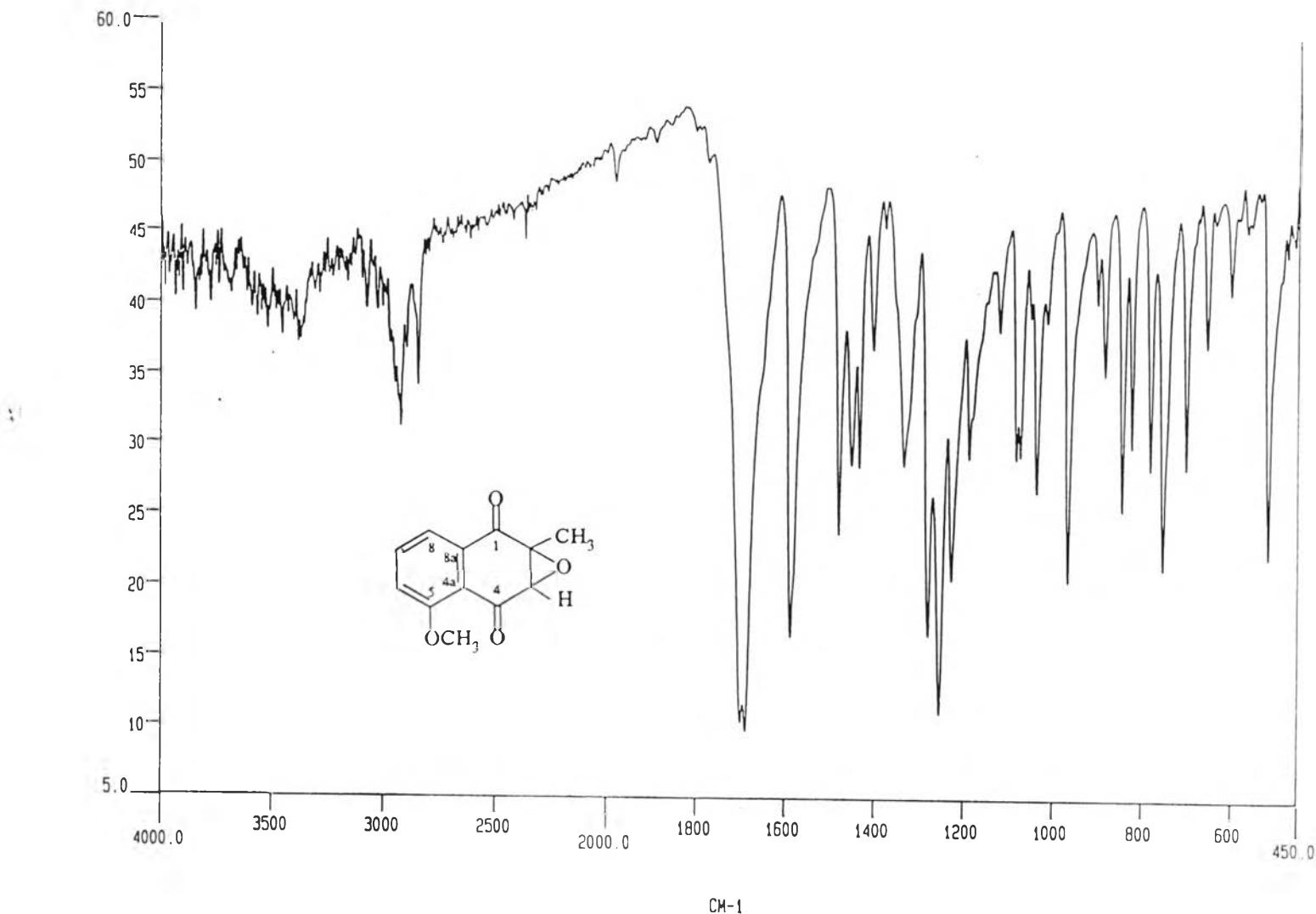


Figure 121 The IR spectrum of compound 107 (in KBr disc)

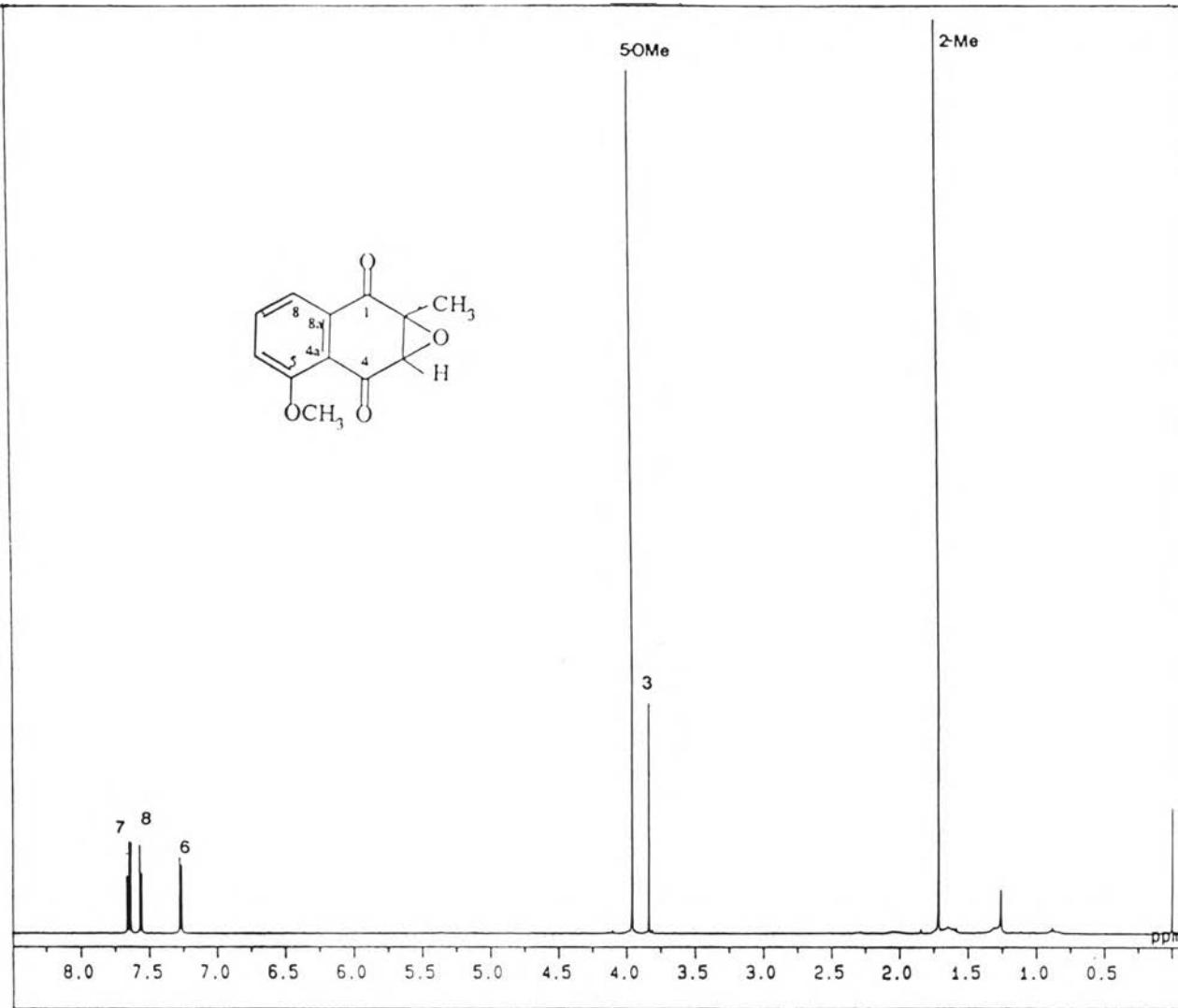


Figure 122 The ^1H NMR (500 MHz) spectrum of compound 107 (in CDCl_3)

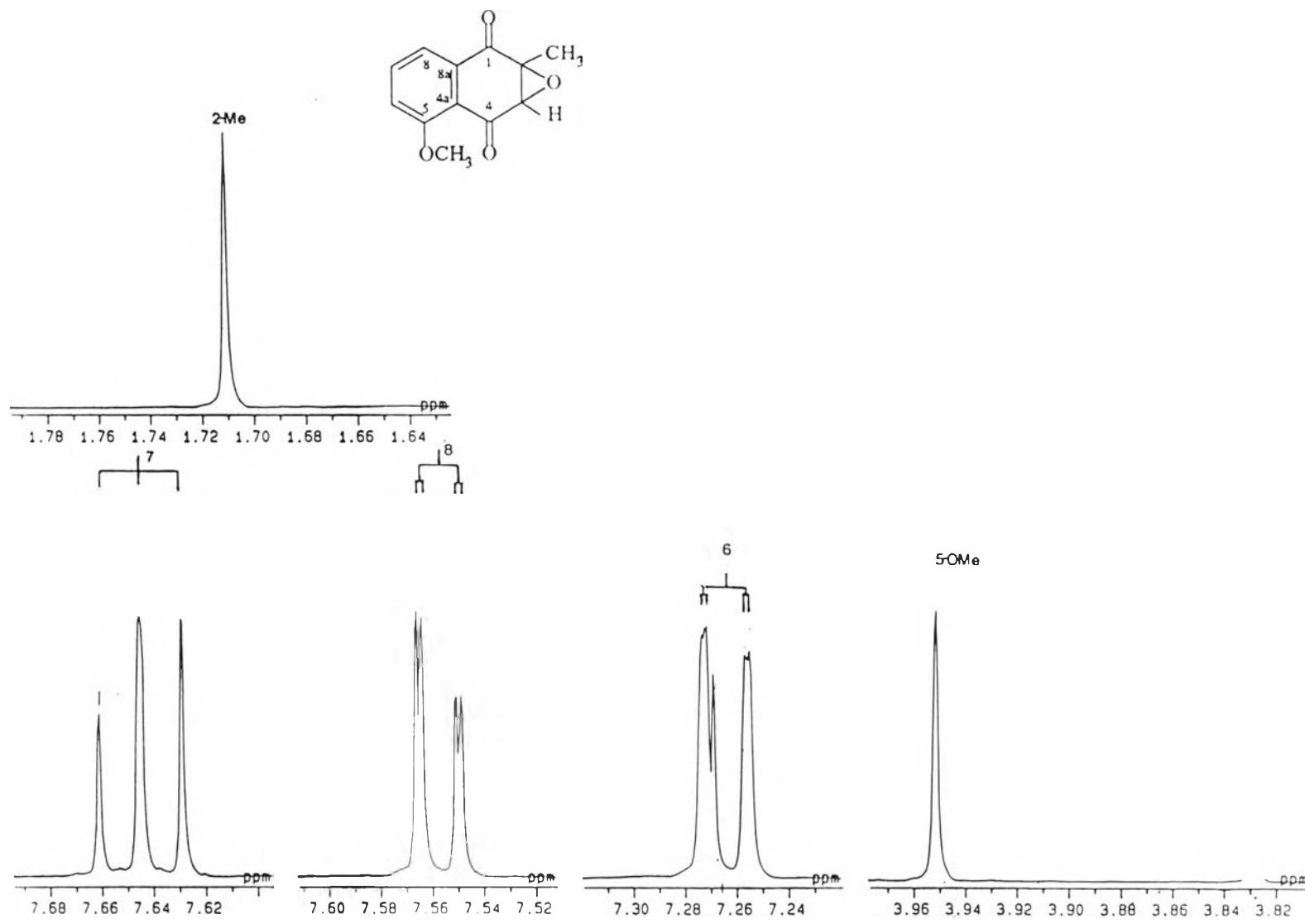


Figure 123 Expansion of the ^1H NMR (500 MHz) of compound 107 (in CDCl_3) :

δ_H 1.64-7.68 ppm

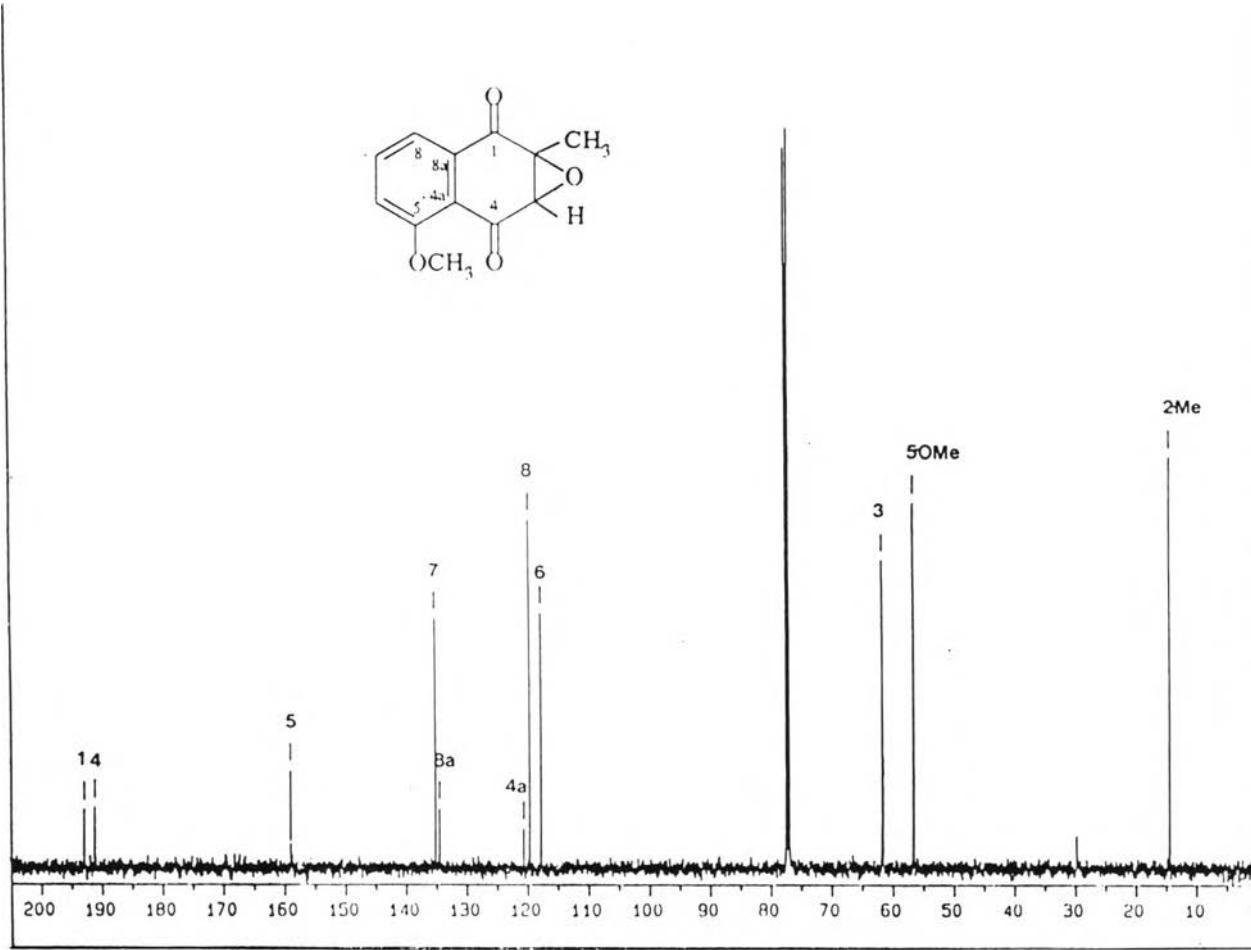


Figure 124 The ^{13}C NMR (125 MHz) of compound 107 (in CDCl_3)

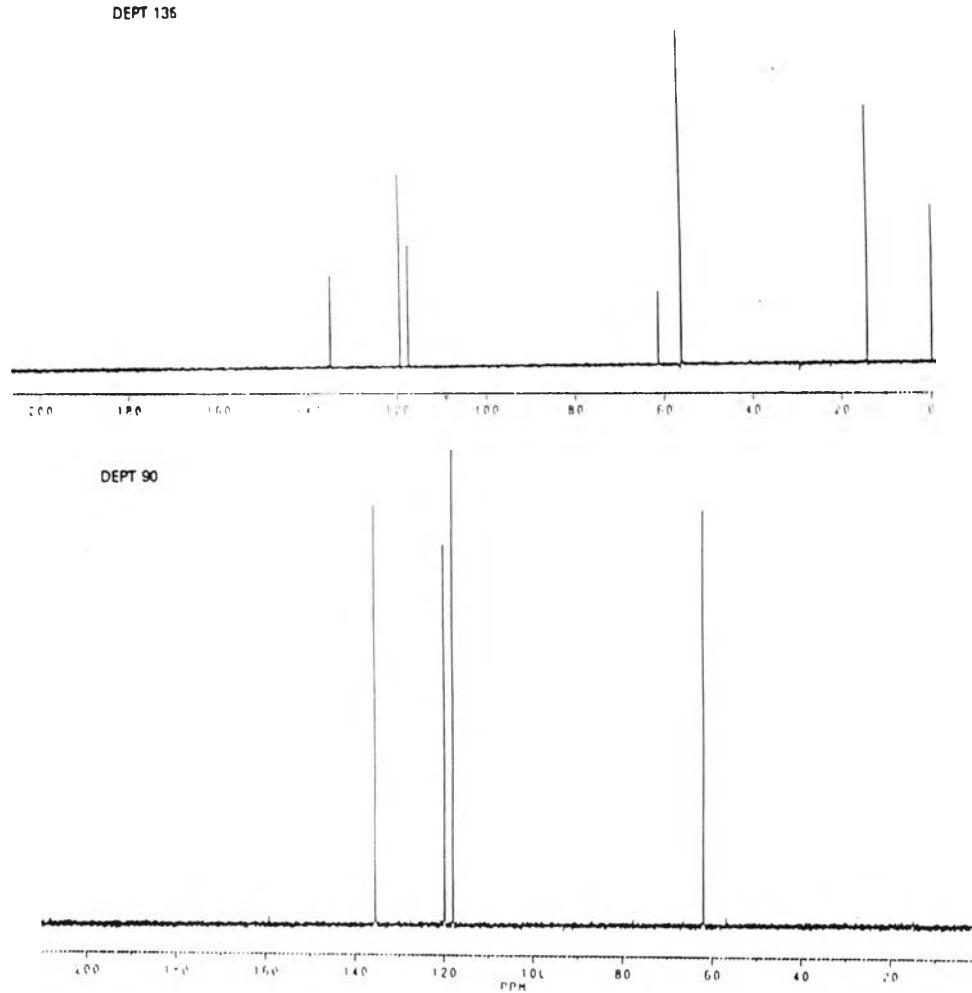


Figure 125 The DEPT (50 MHz) spectrum of compound 107 (in CDCl_3)

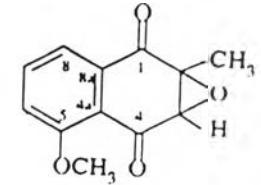
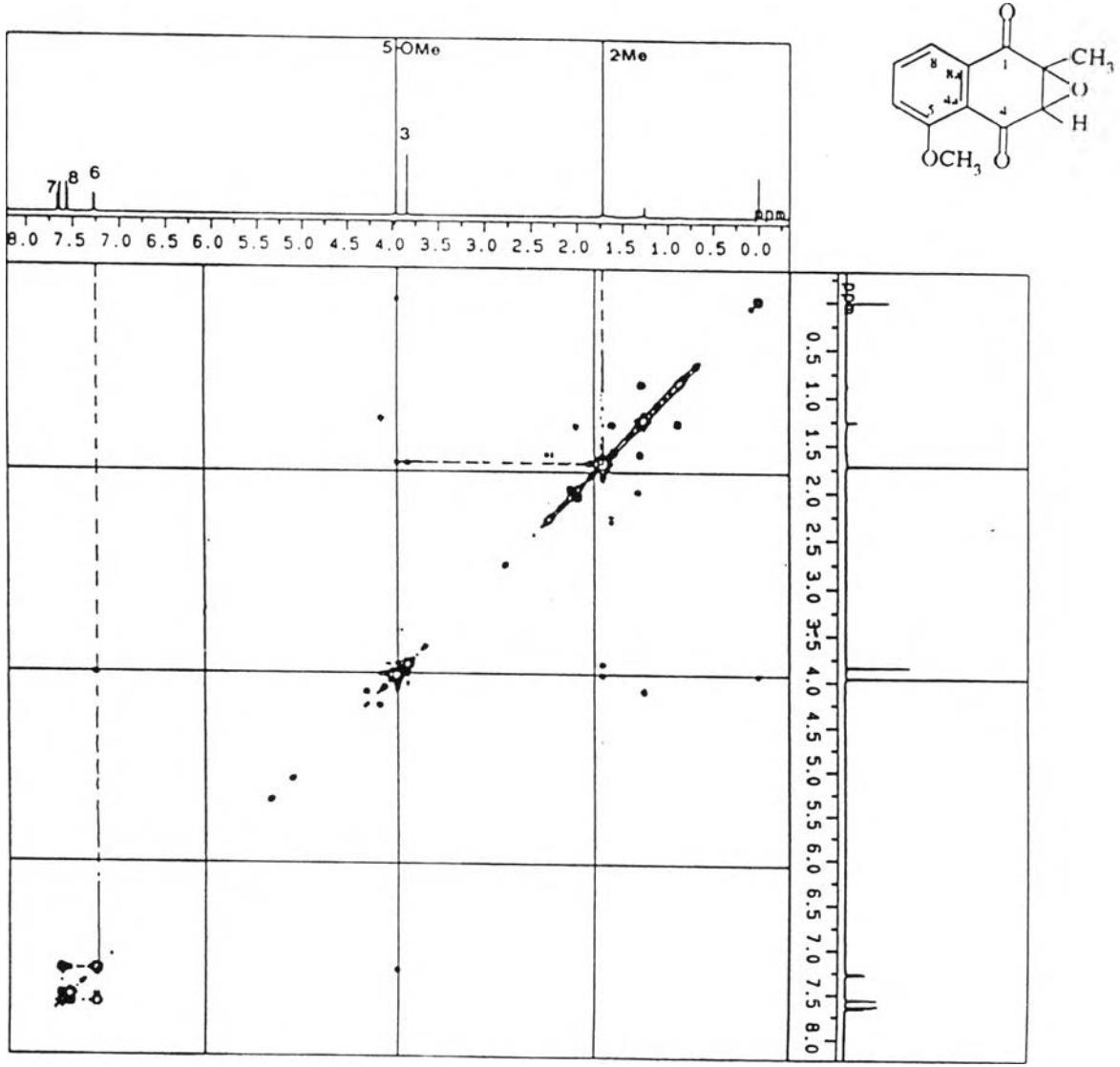


Figure 126 The ^1H - ^1H COSY (500 MHz) spectrum of compound 107 (in CDCl_3)

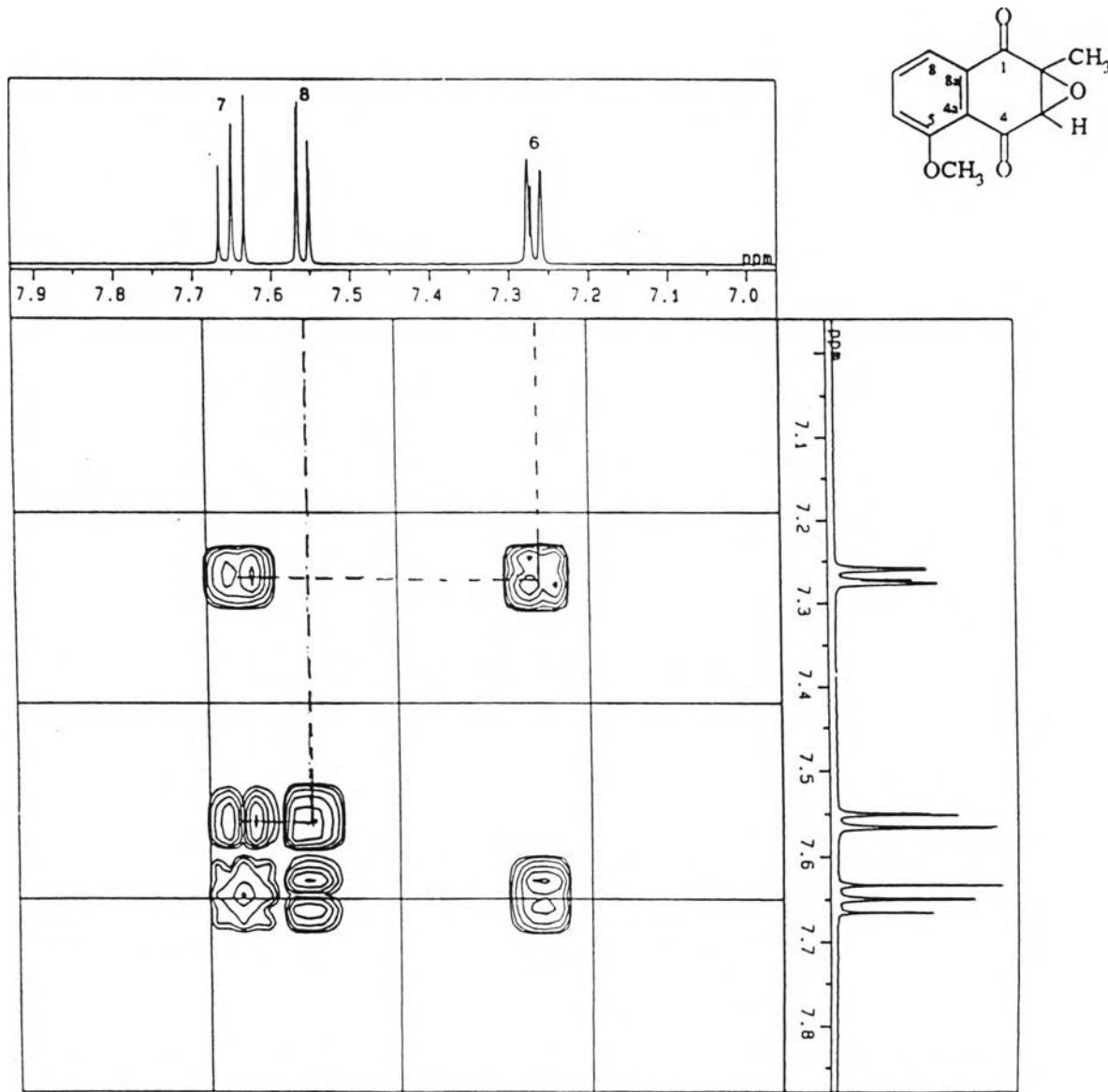


Figure 127 Expansion of the ^1H - ^1H COSY (500 MHz) spectrum of compound 107
(in CDCl_3) : δ_{H} 7.00-7.90 ppm

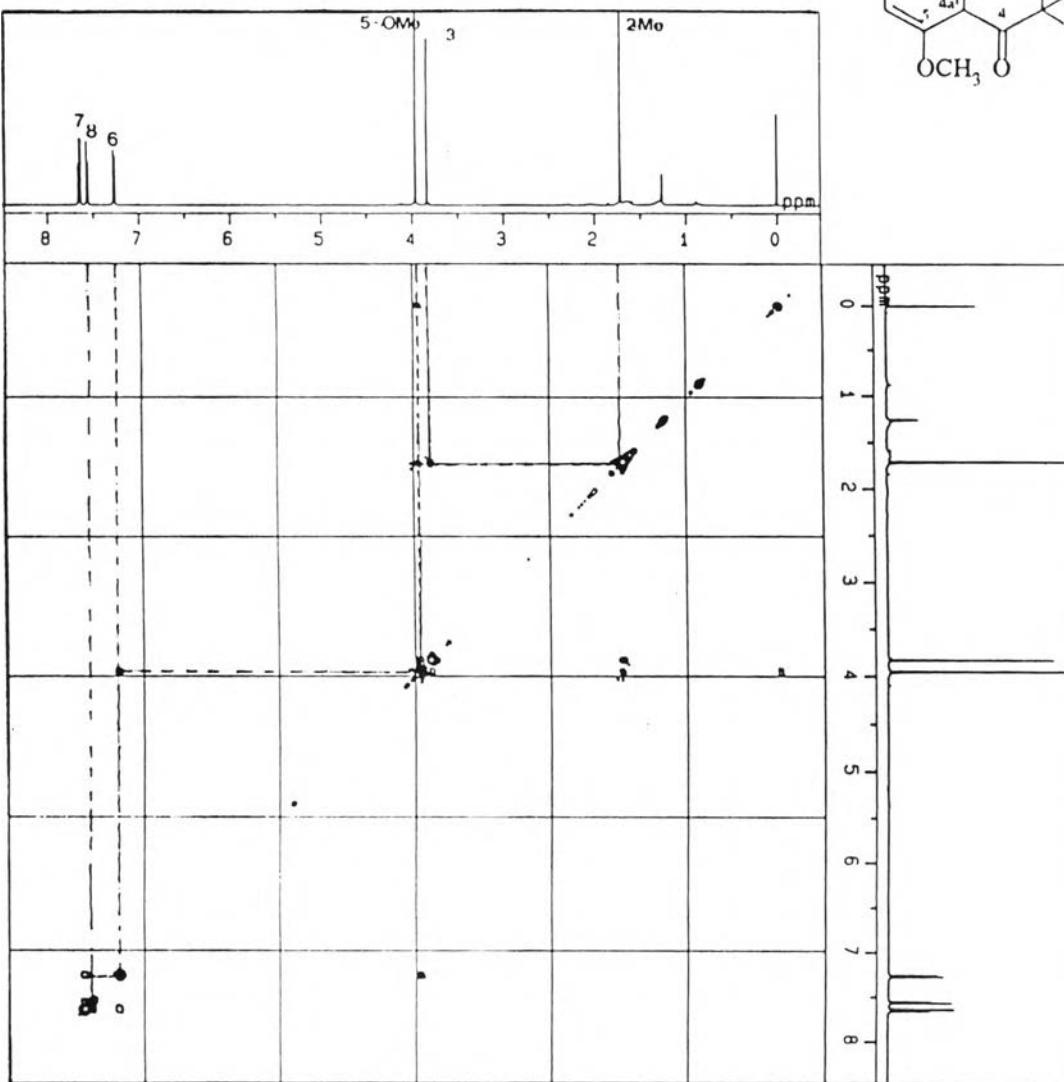
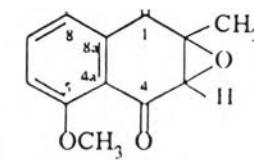


Figure 128 The NOESY (500 MHz) spectrum of compound 107 (in CDCl_3)

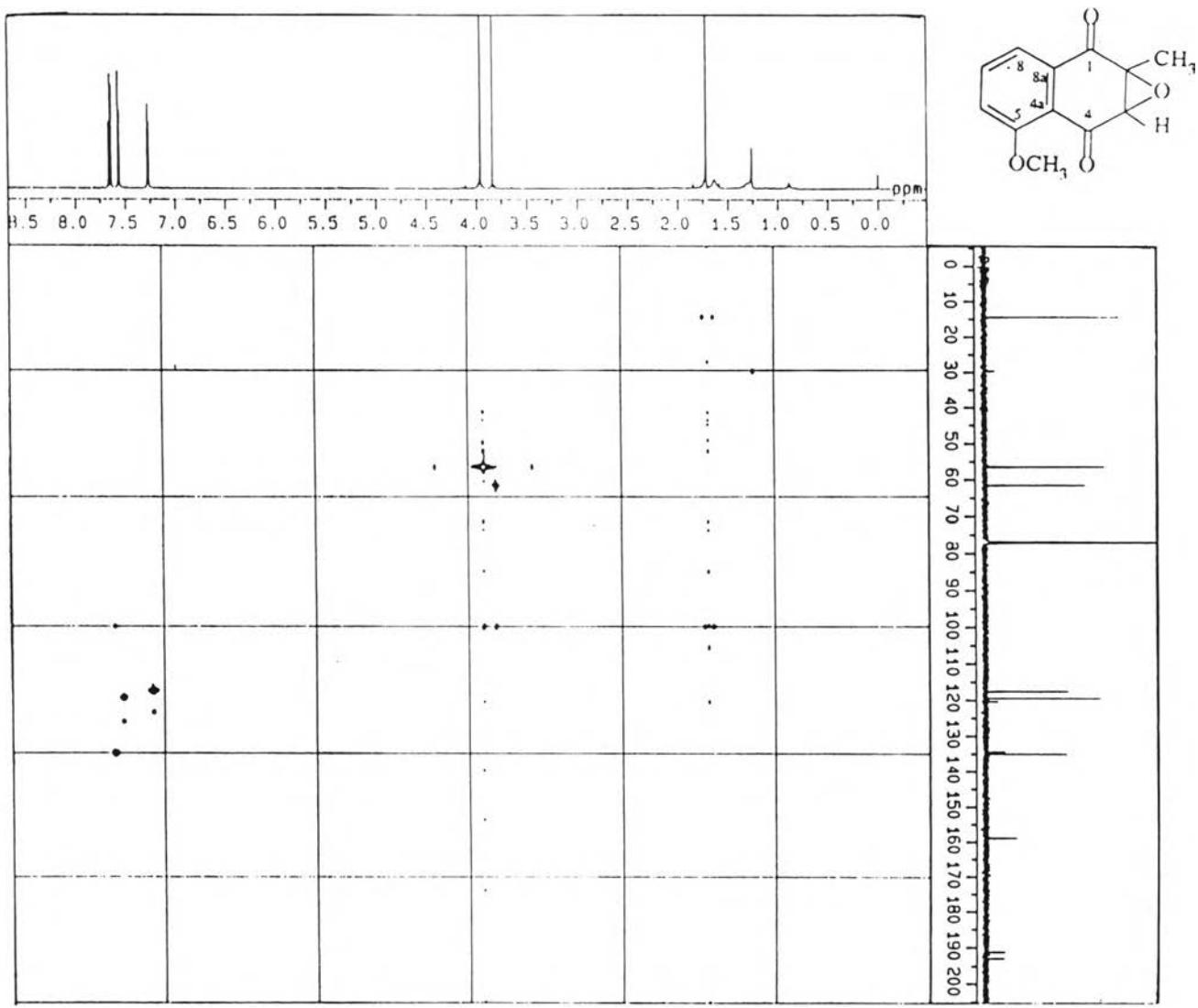


Figure 129 The HMQC spectrum of compound 107 (in CDCl_3)

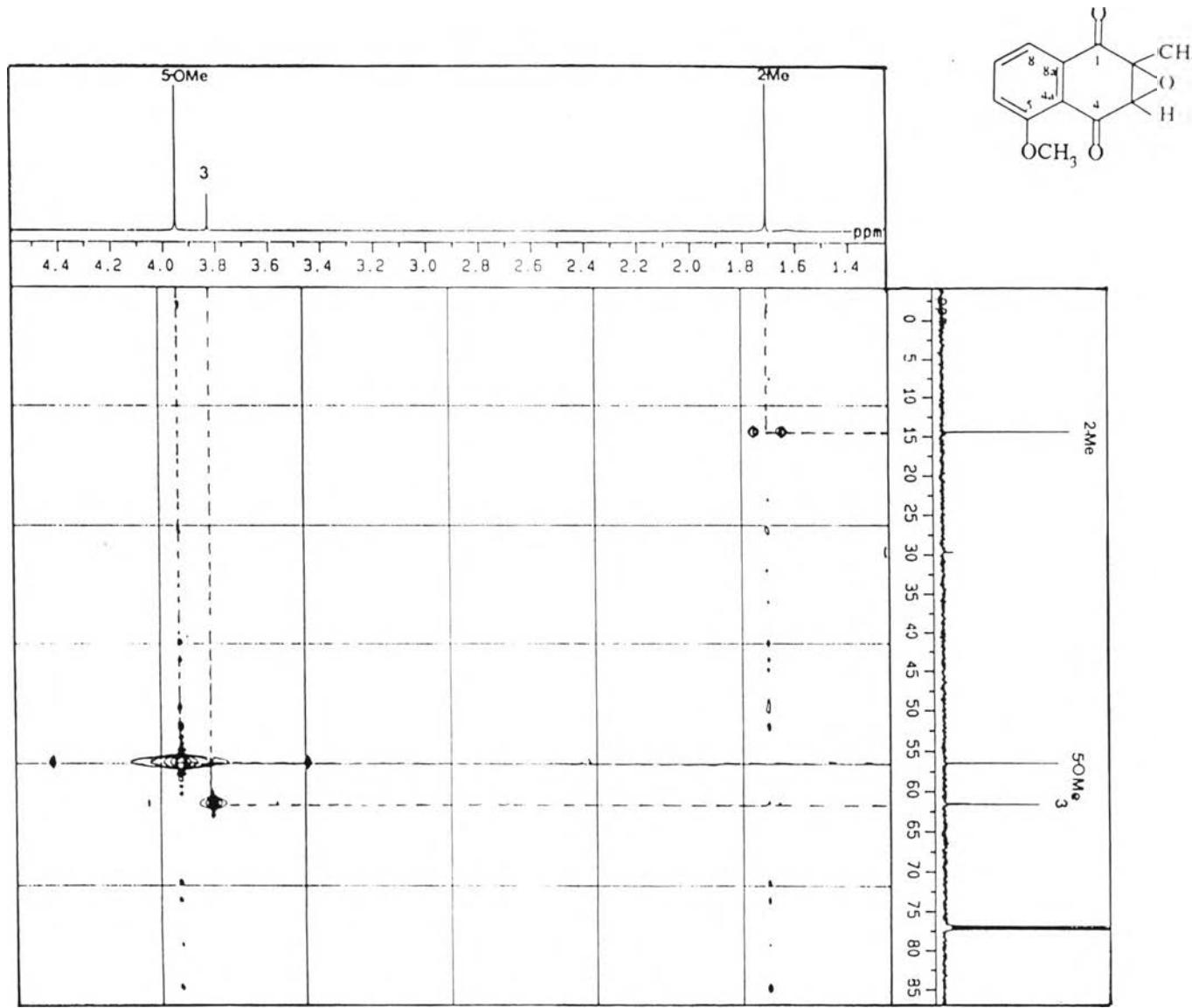


Figure 130 Expansion of the HMQC spectrum of compound 107 (in CDCl_3) :

δ_{H} 1.40-4.40 ; δ_{C} 0.00-0.85 ppm

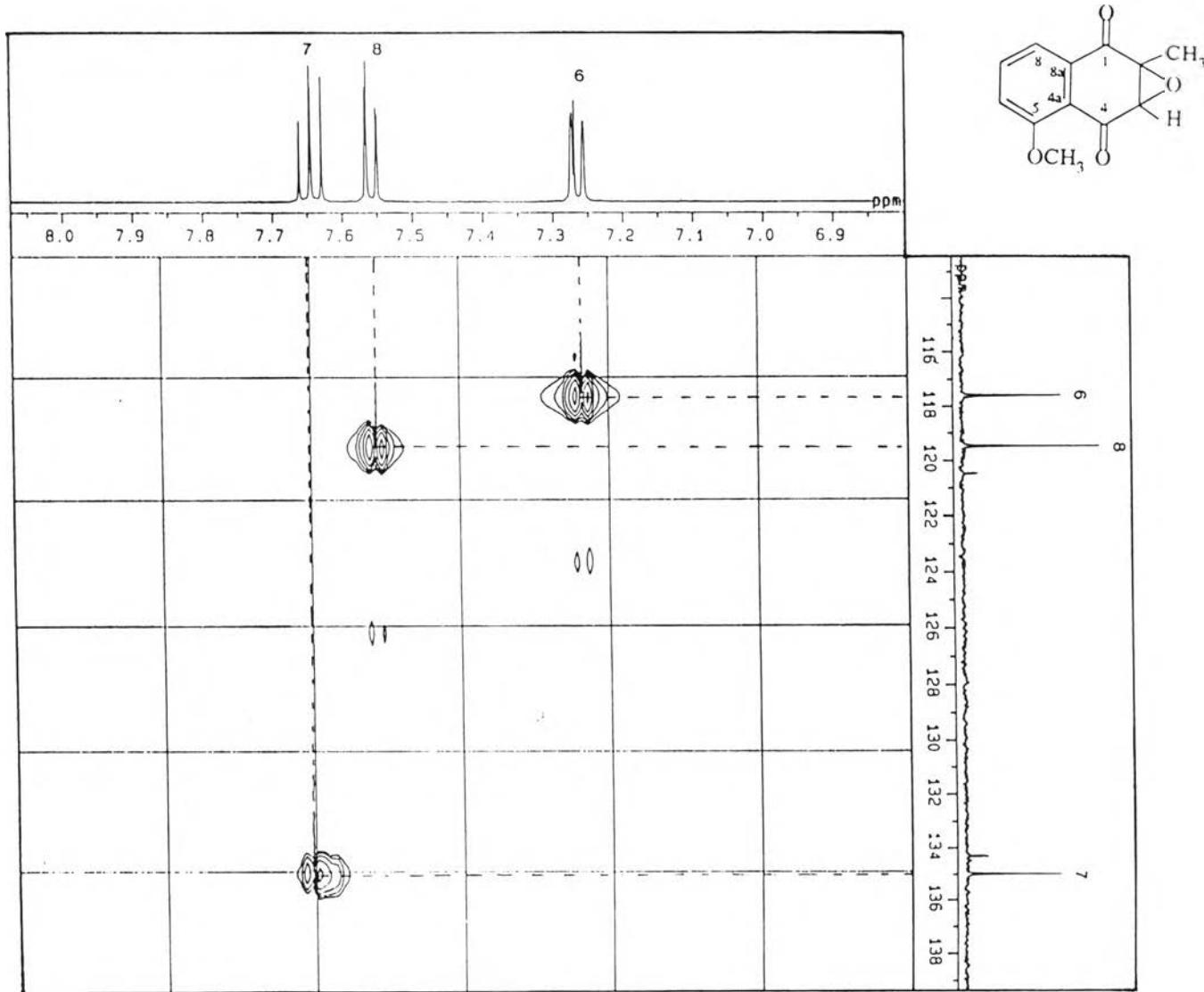


Figure 131 Expansion of the HMQC spectrum of compound 107 (in CDCl_3) :

δ_{H} 6.90-8.00 ; δ_{C} 116.00-138.00 ppm

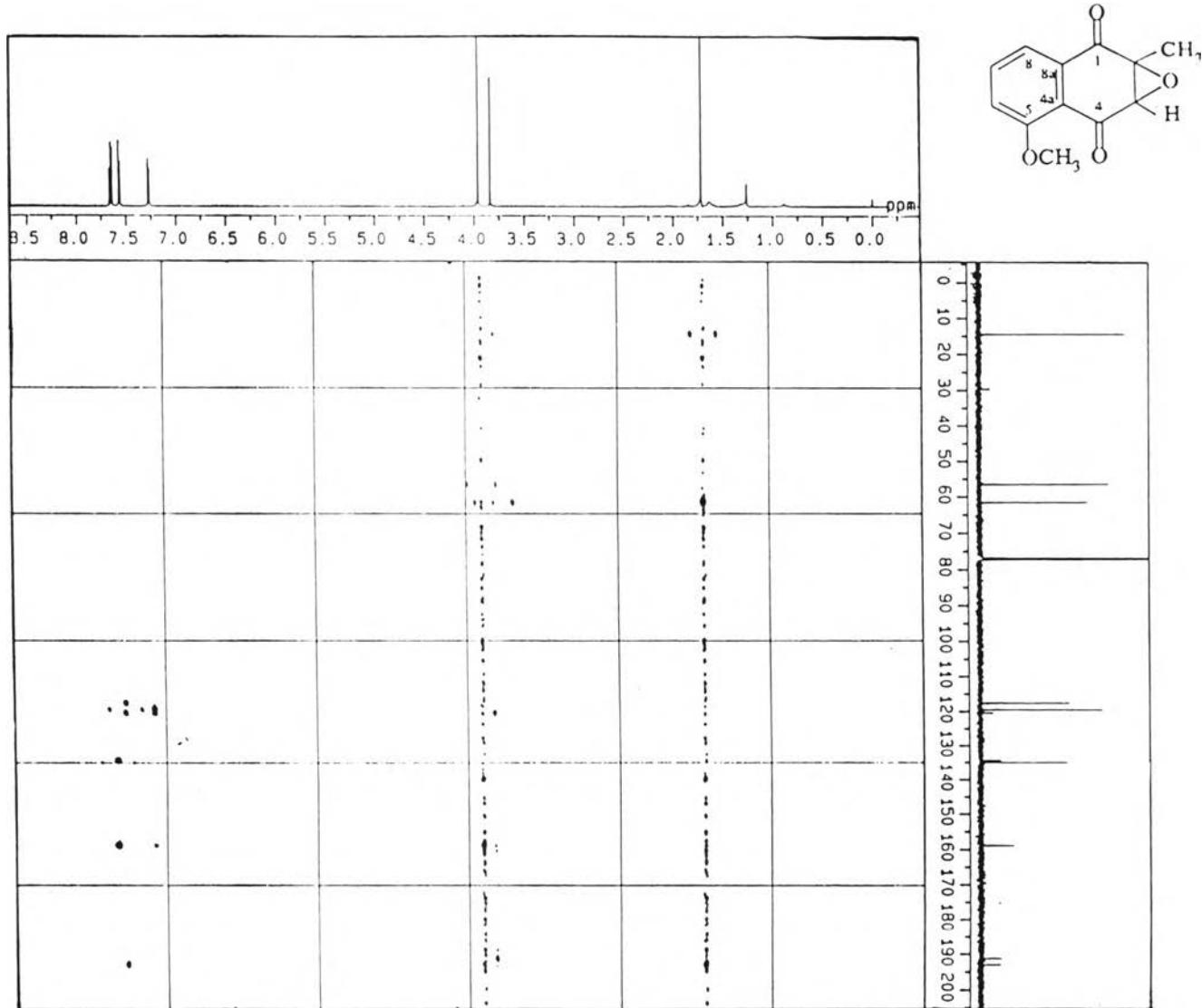


Figure 132 The HMBC spectrum of compound 107 (in CDCl₃)

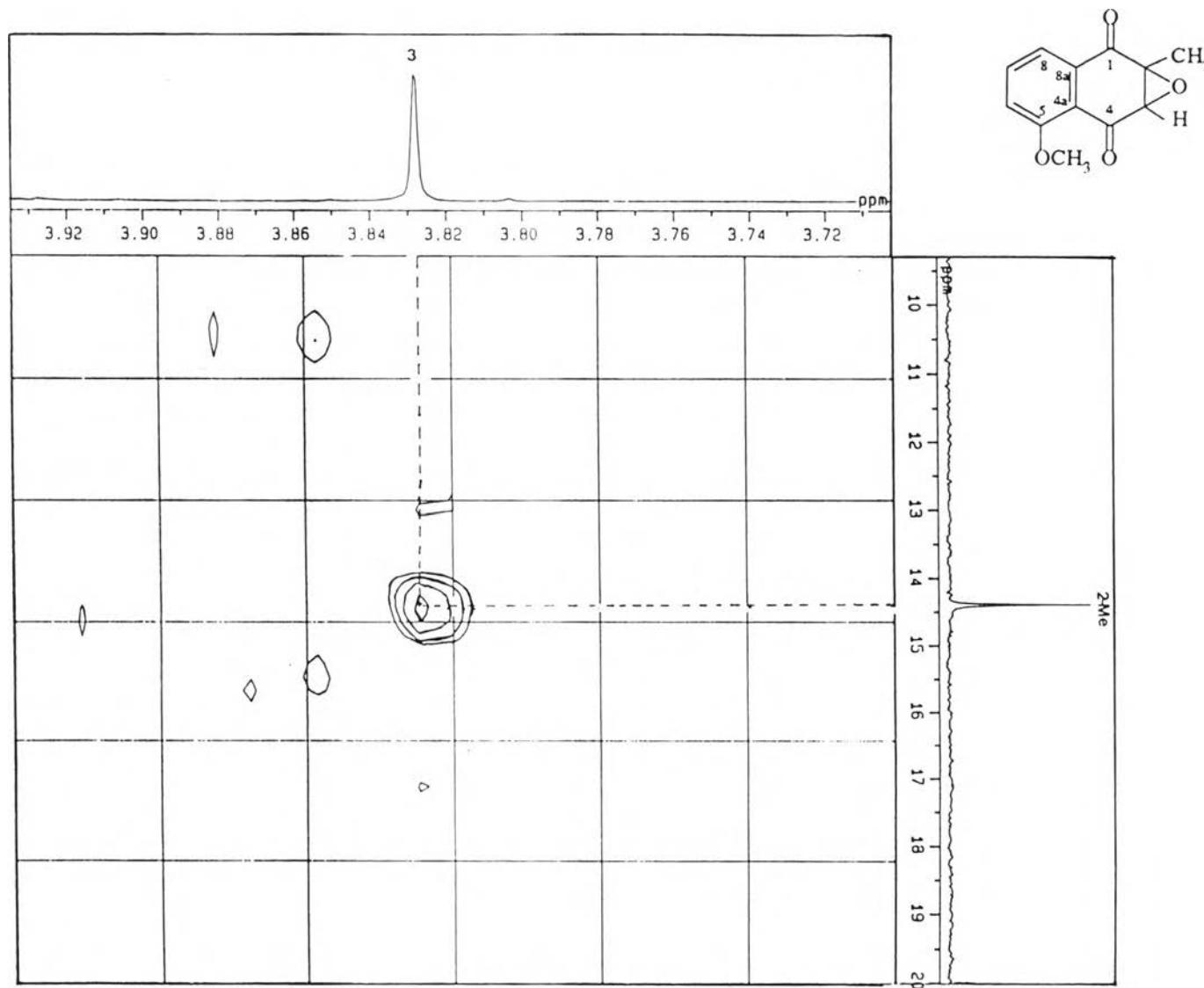


Figure 133 Expansion of the HMBC spectrum of compound 107 (in CDCl_3) :

δ_{H} 3.72-3.92 ; δ_{C} 10.00-20.00 ppm

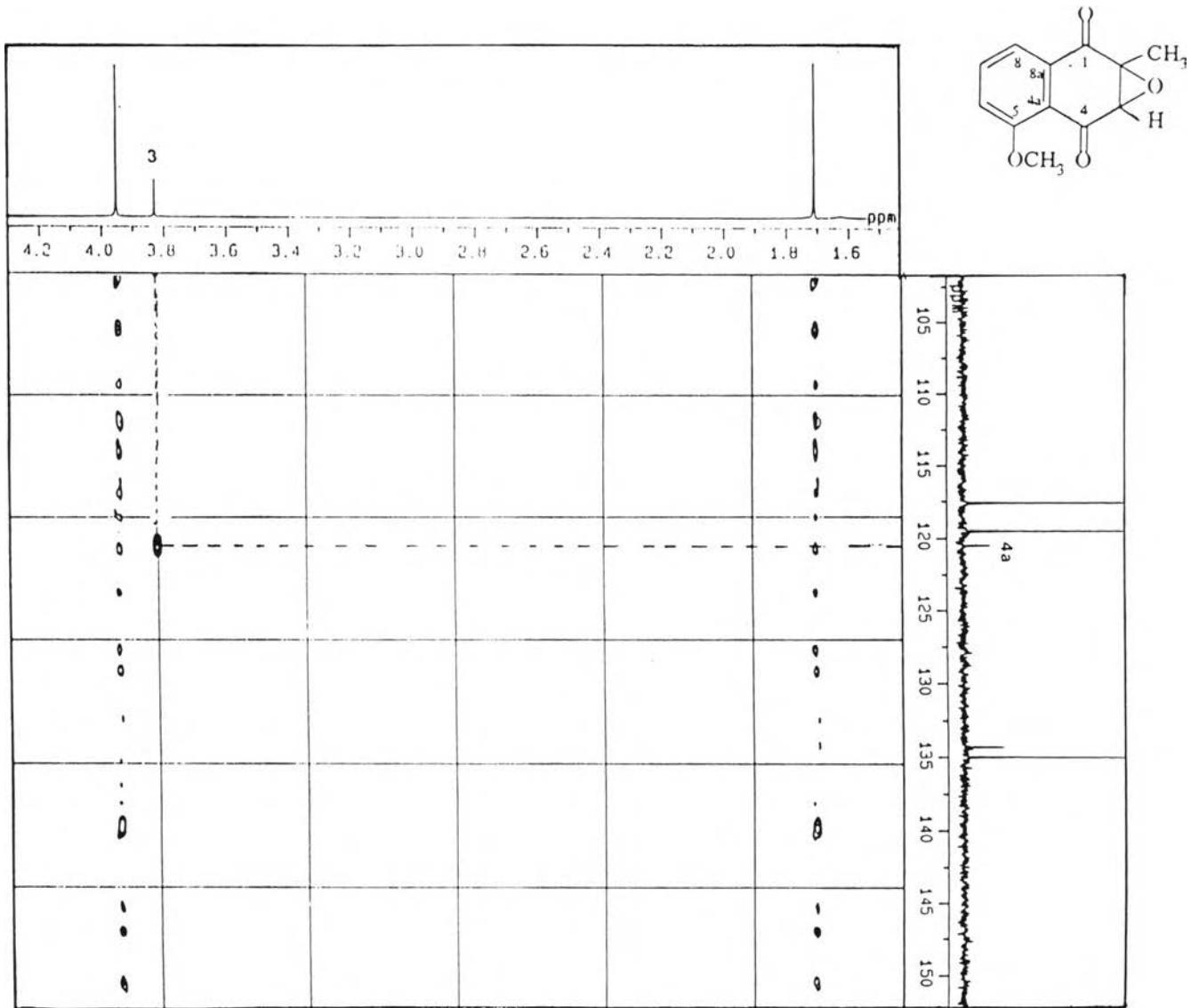


Figure 134 Expansion of the HMBC spectrum of compound 107 (in CDCl_3) :

δ_{H} 1.60-4.20 ; δ_{C} 105.00-150.00 ppm

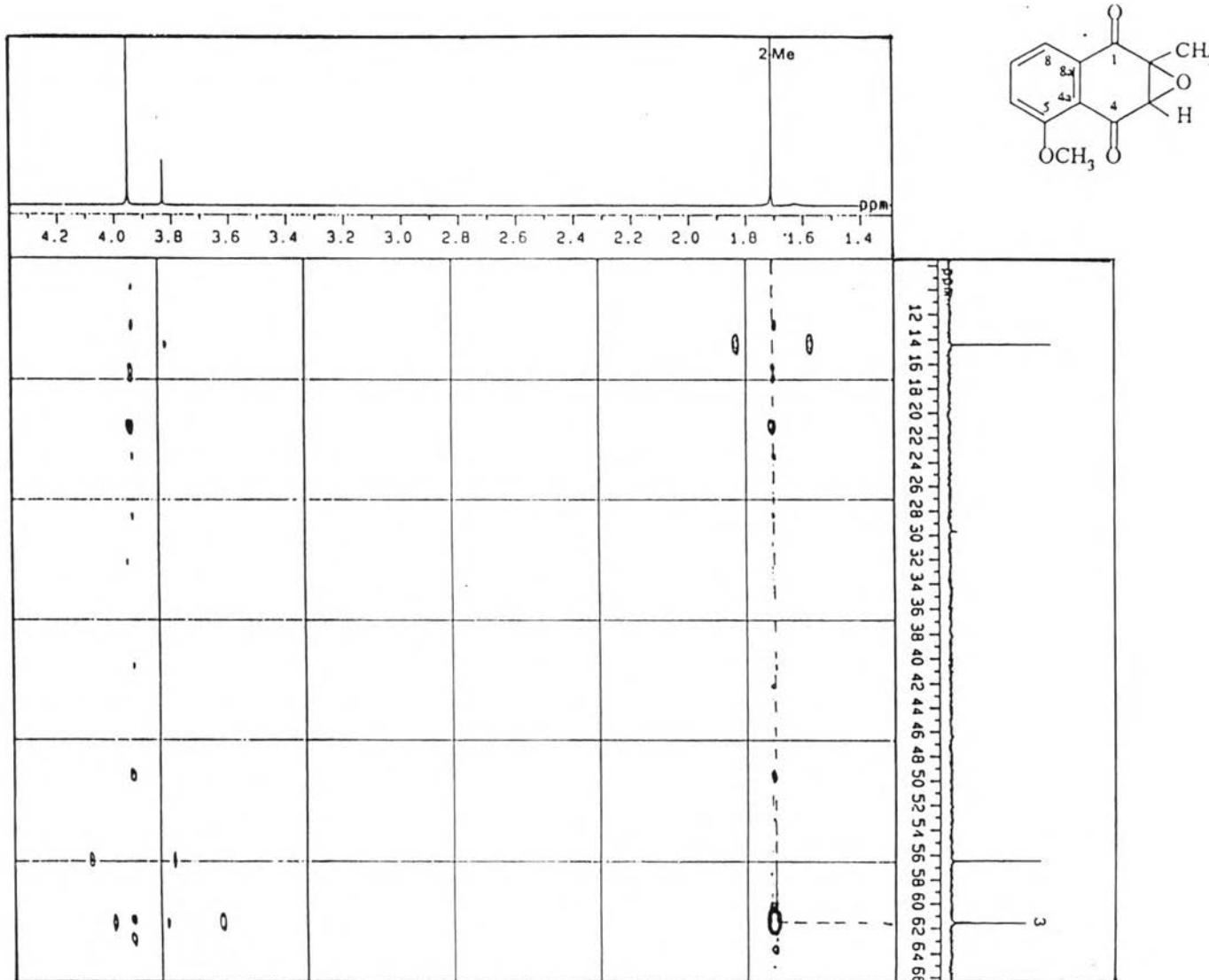


Figure 135 Expansion of the HMBC spectrum of compound 107 (in CDCl_3) :
 δ_{H} 1.40-4.20 ; δ_{C} 12.00-66.00 ppm

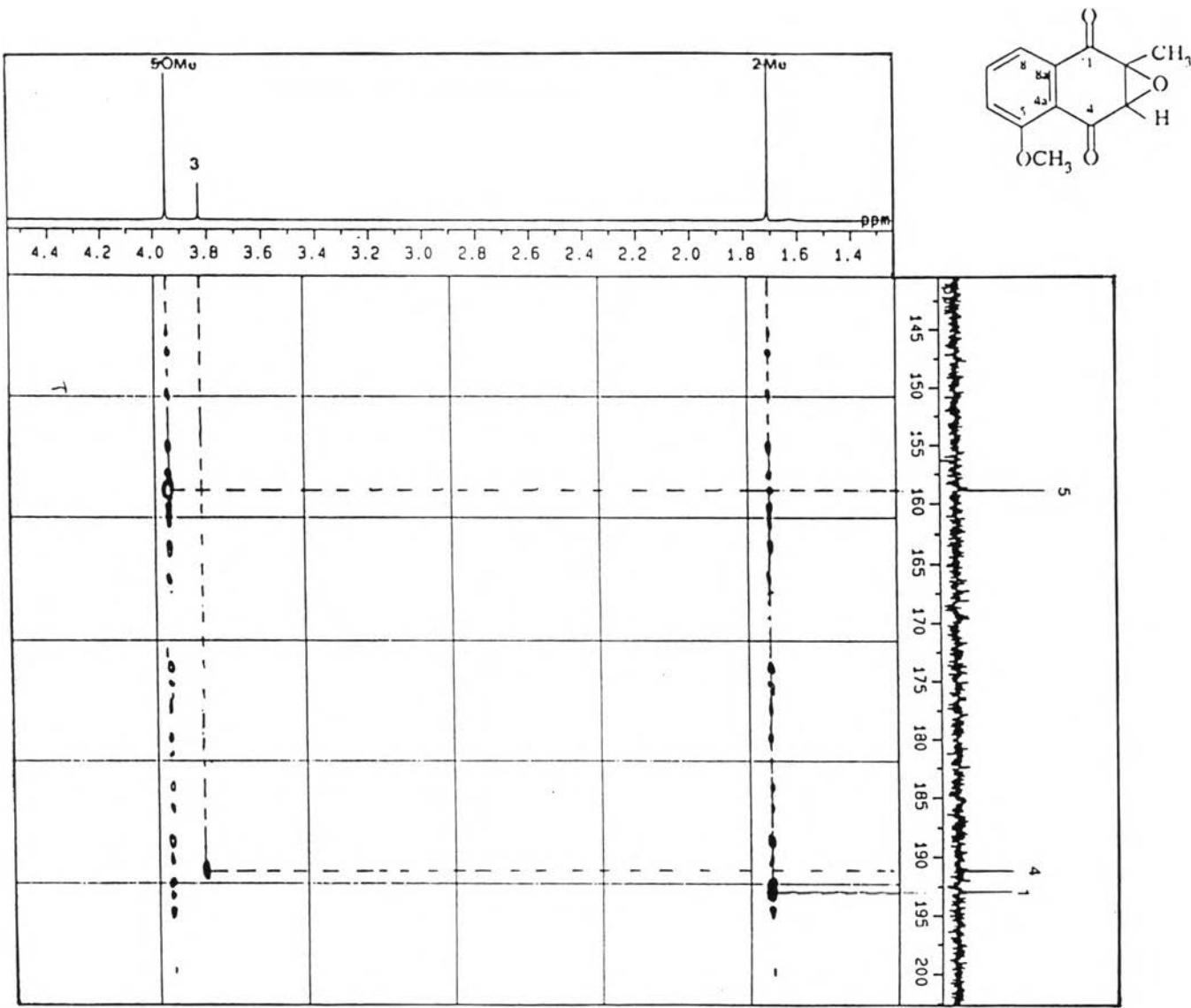


Figure 136 Expansion of the HMBC spectrum of compound 107 (in CDCl_3) :

δ_{H} 1.40-4.40 ; δ_{C} 145.00-200.00 ppm

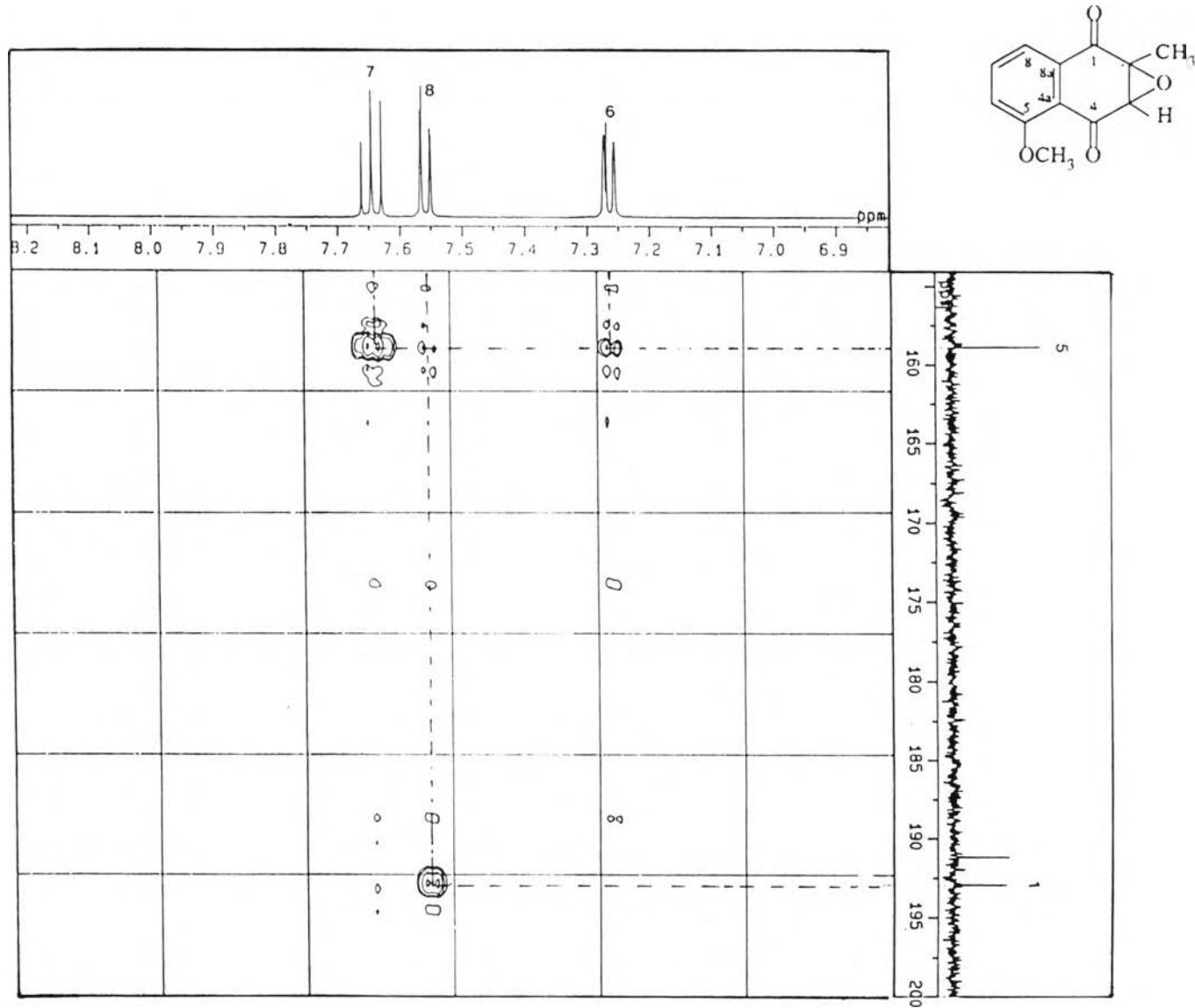


Figure 137 Expansion of the HMBC spectrum of compound 107 (in CDCl₃) :

δ_H 6.90-8.20 ; δ_C 155.00-200.00 ppm

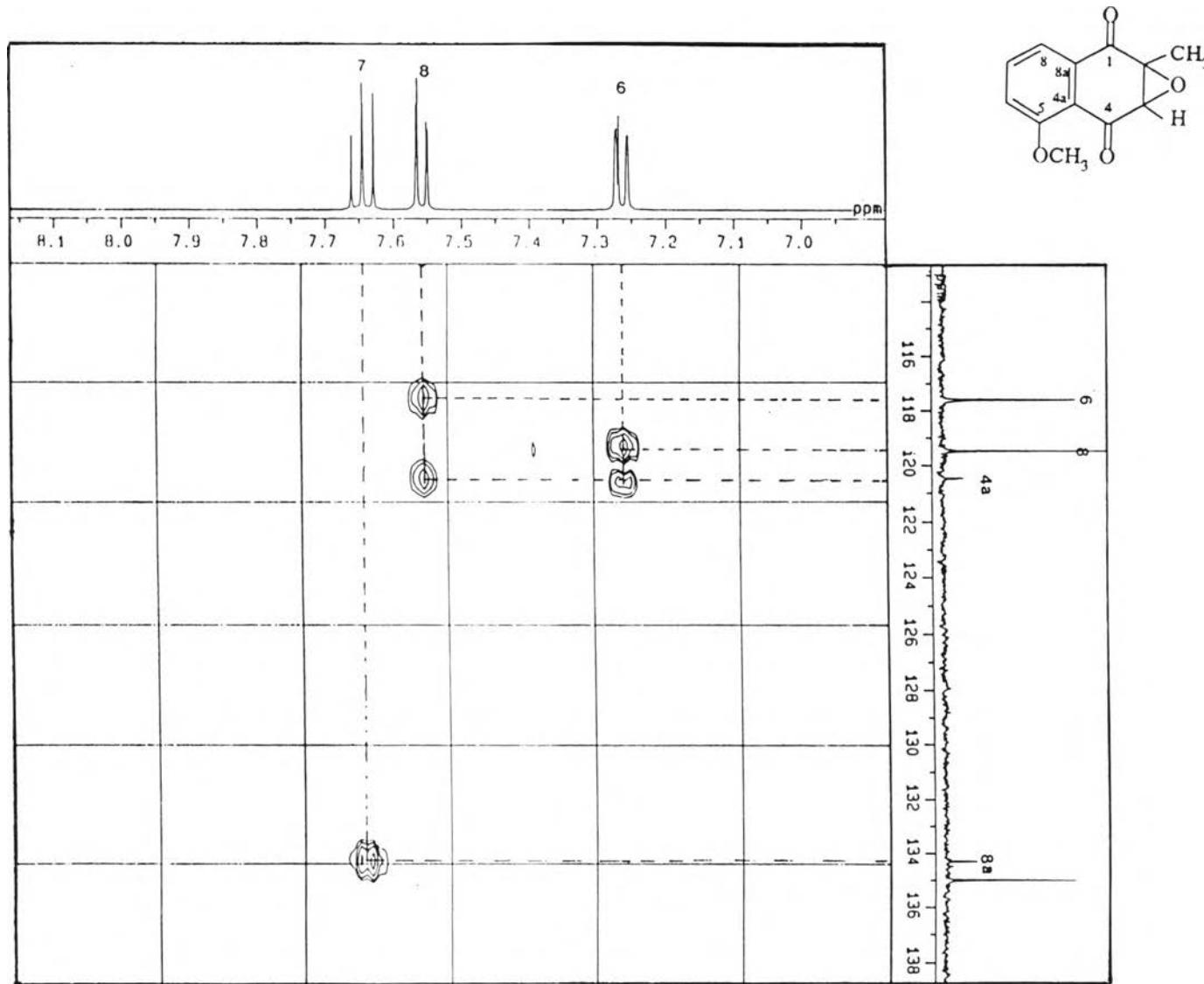
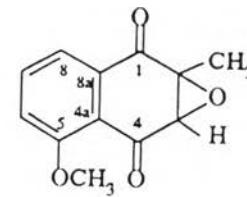


Figure 138 Expansion of the HMBC spectrum of compound 107 (in CDCl_3) :

δ_{H} 7.00-8.10 ; δ_{C} 116.00-138.00 ppm



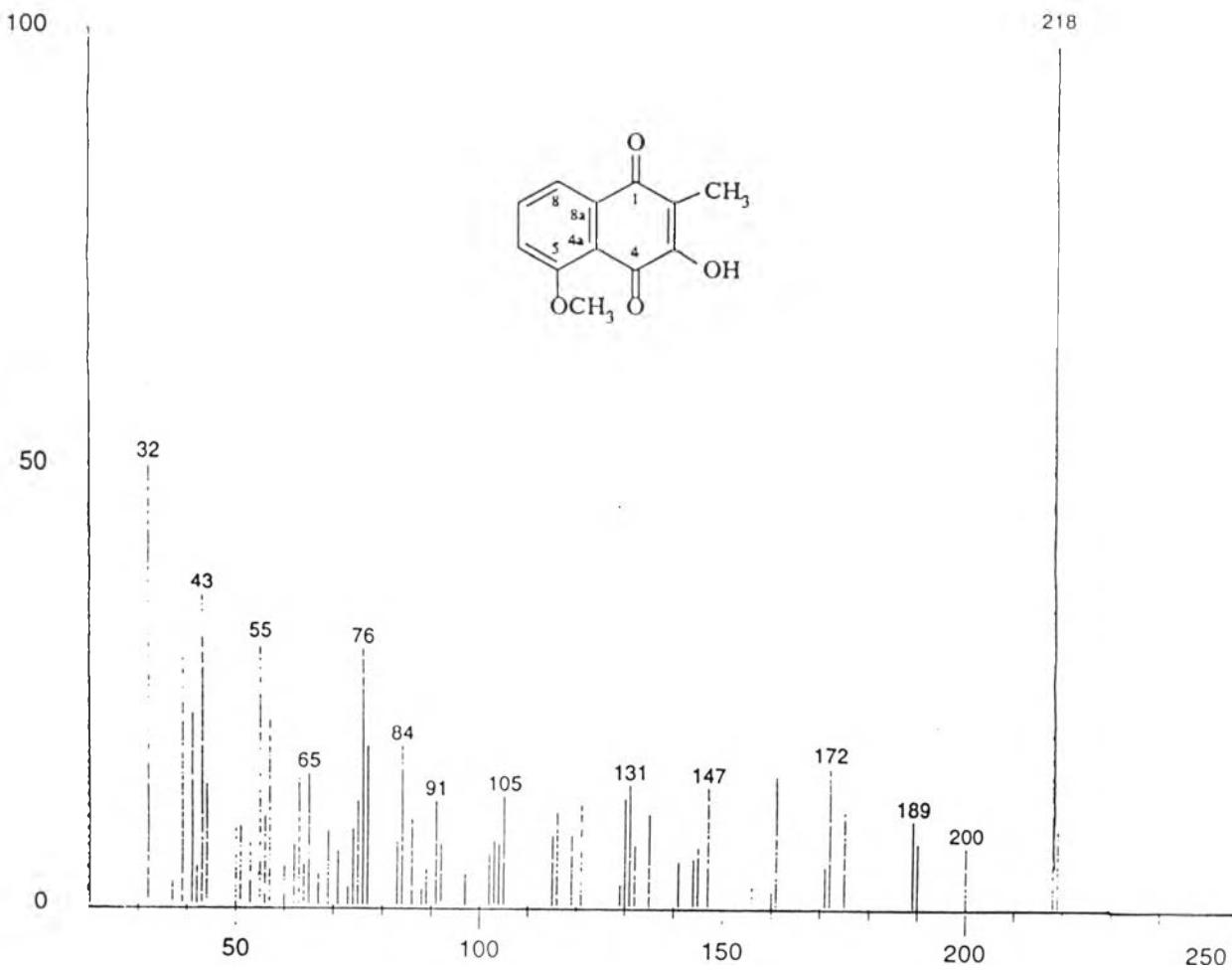


Figure 139 The EI mass spectrum of compound 108

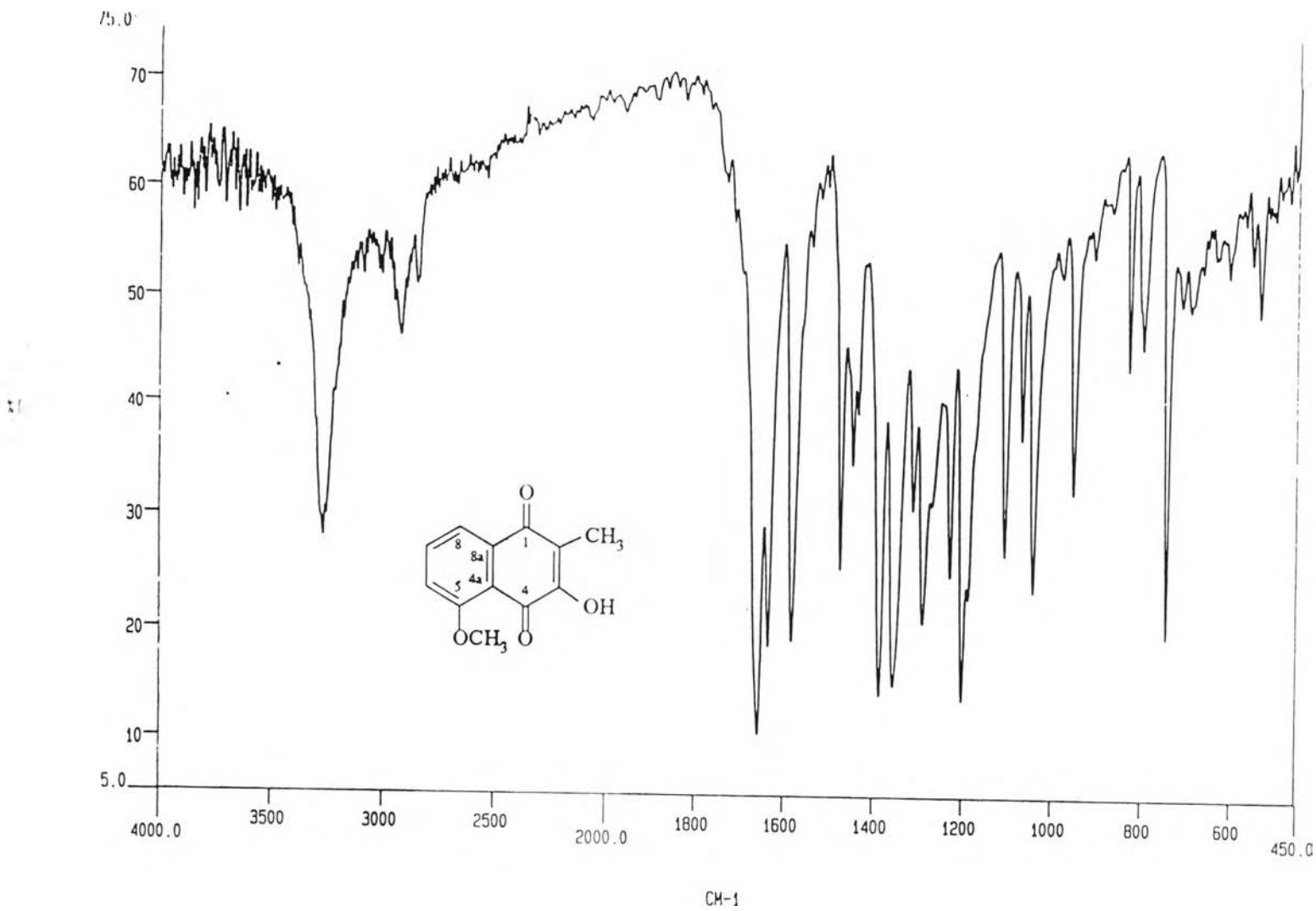


Figure 140 The IR spectrum of compound 108 (in KBr disc)

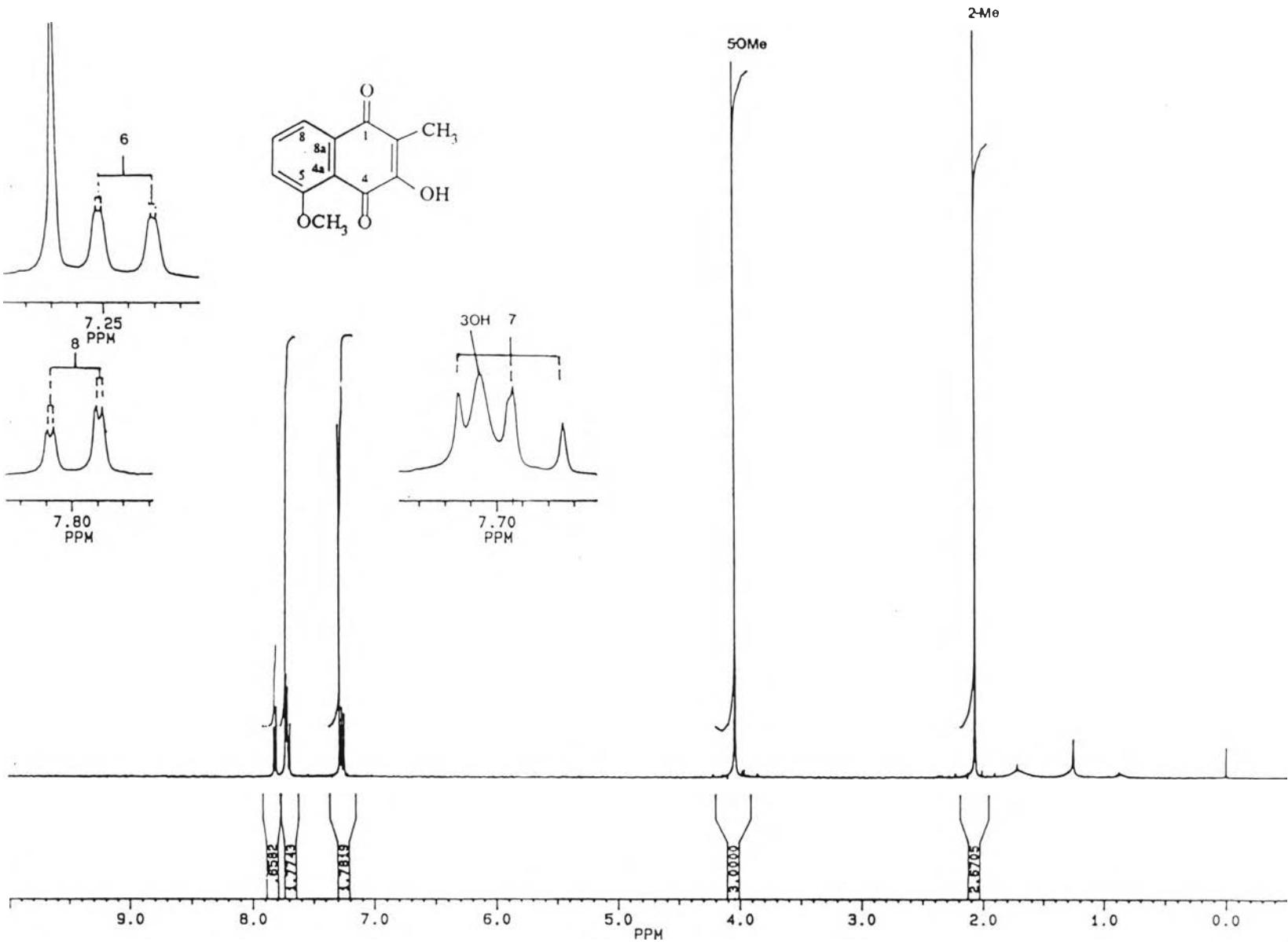


Figure 141 The ^1H NMR (400 MHz) spectrum of compound 108 (in CDCl_3)

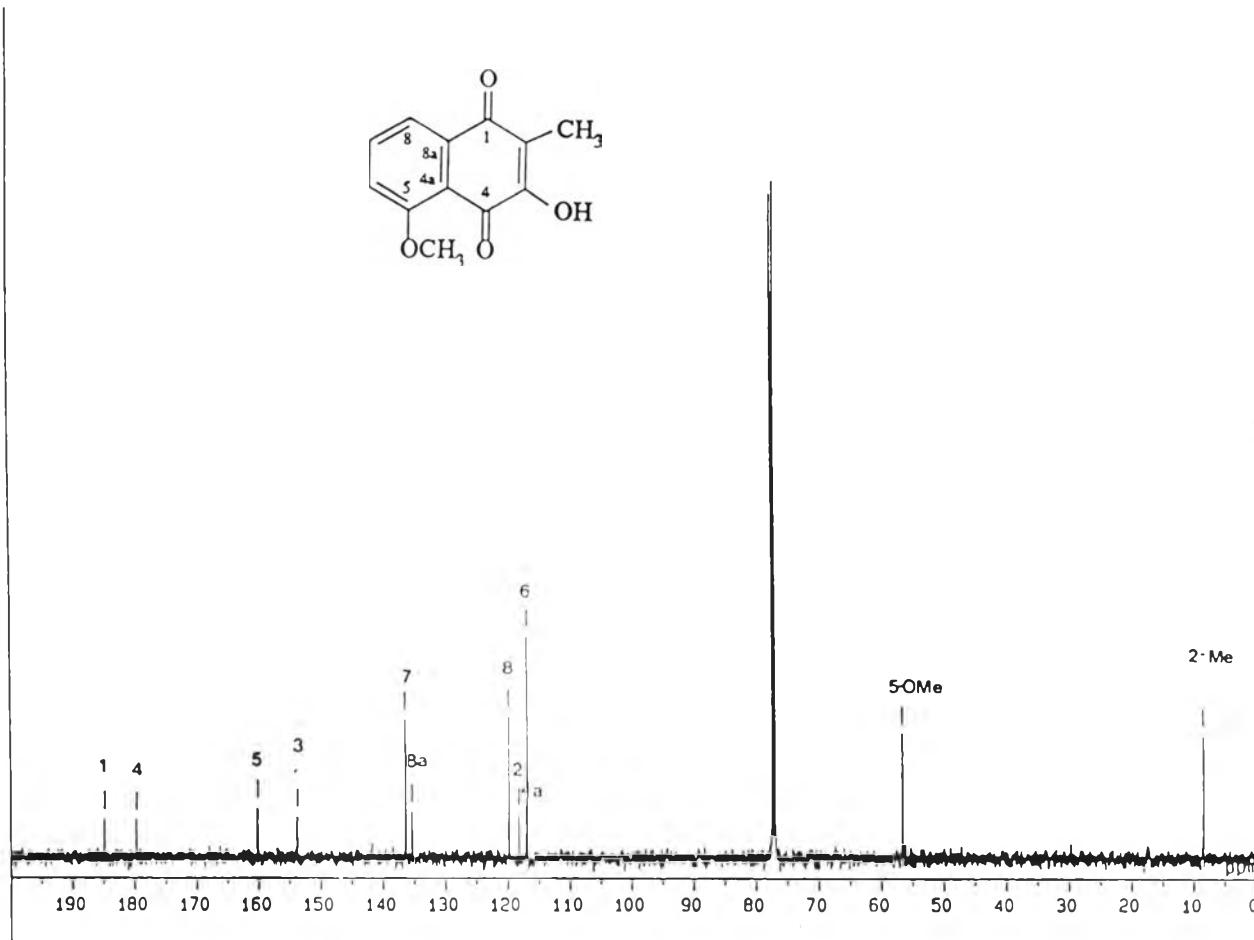
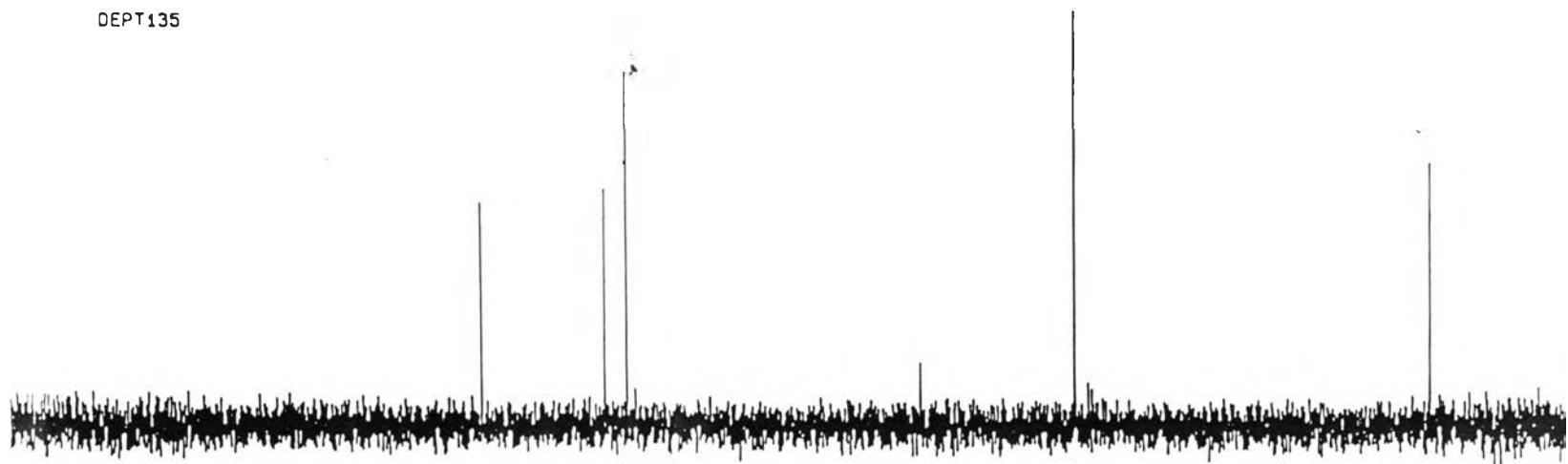


Figure 142 The ^{13}C NMR (125 MHz) spectrum of compound 108 (in CDCl_3)

DEPT135



DEPT90

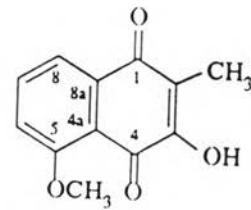
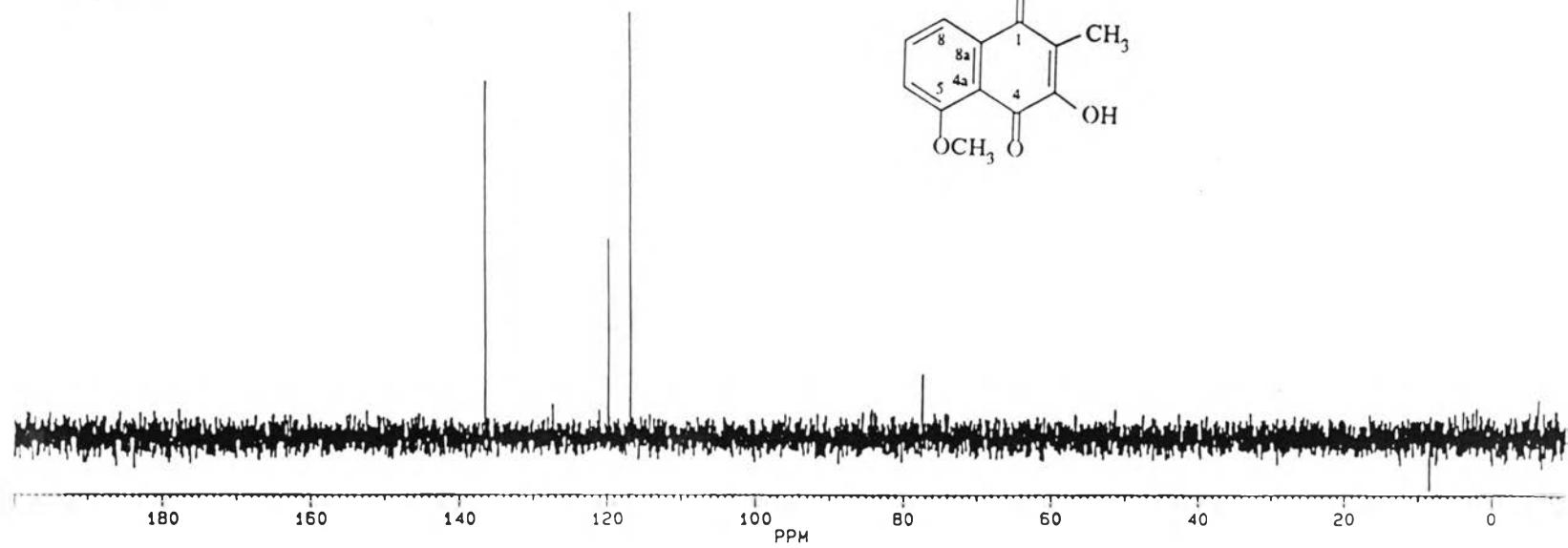


Figure 143. The DEPT (100 MHz) spectrum of compound 108 (in CDCl₃).

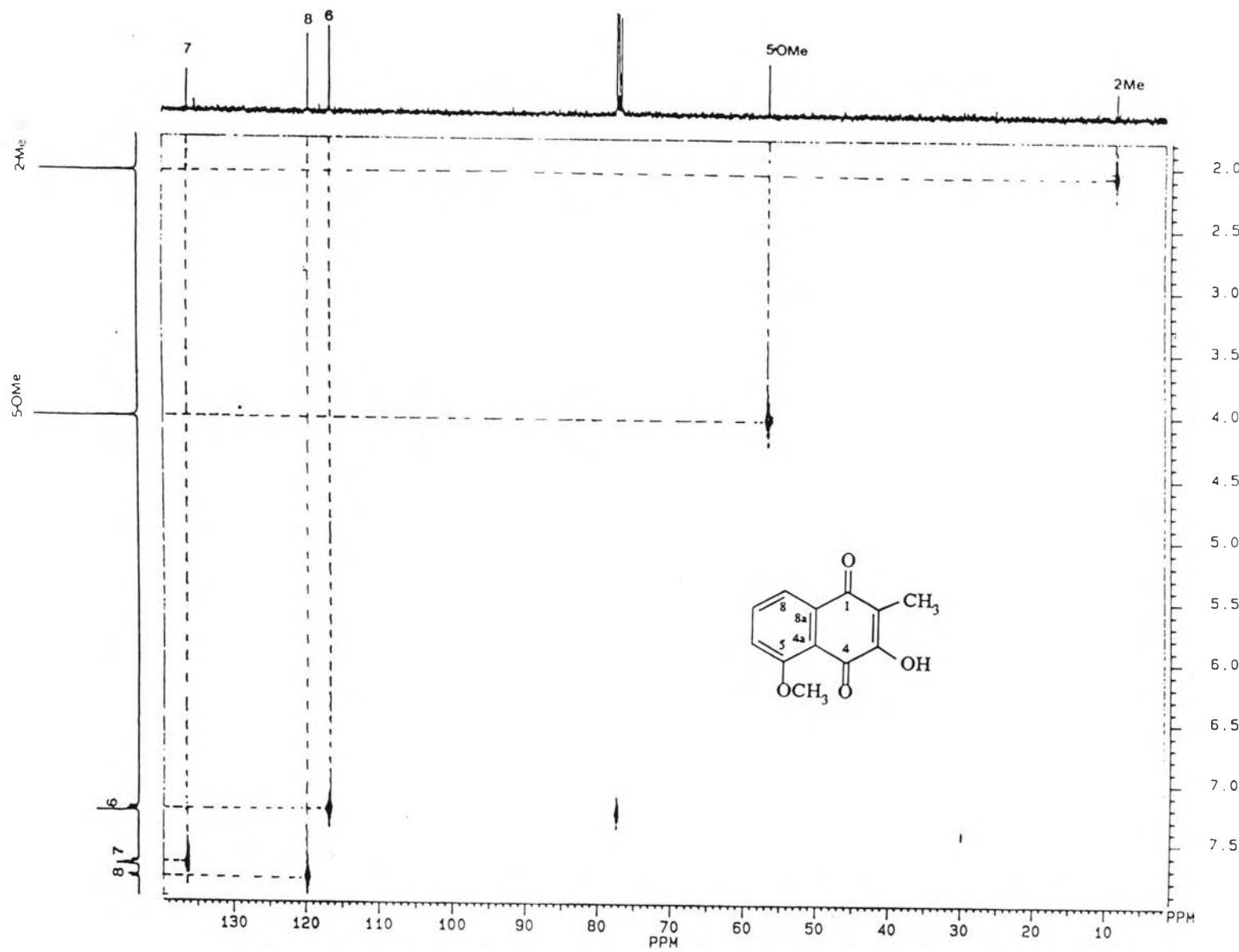


Figure 144 The HETCOR spectrum of compound 108 (in CDCl₃)

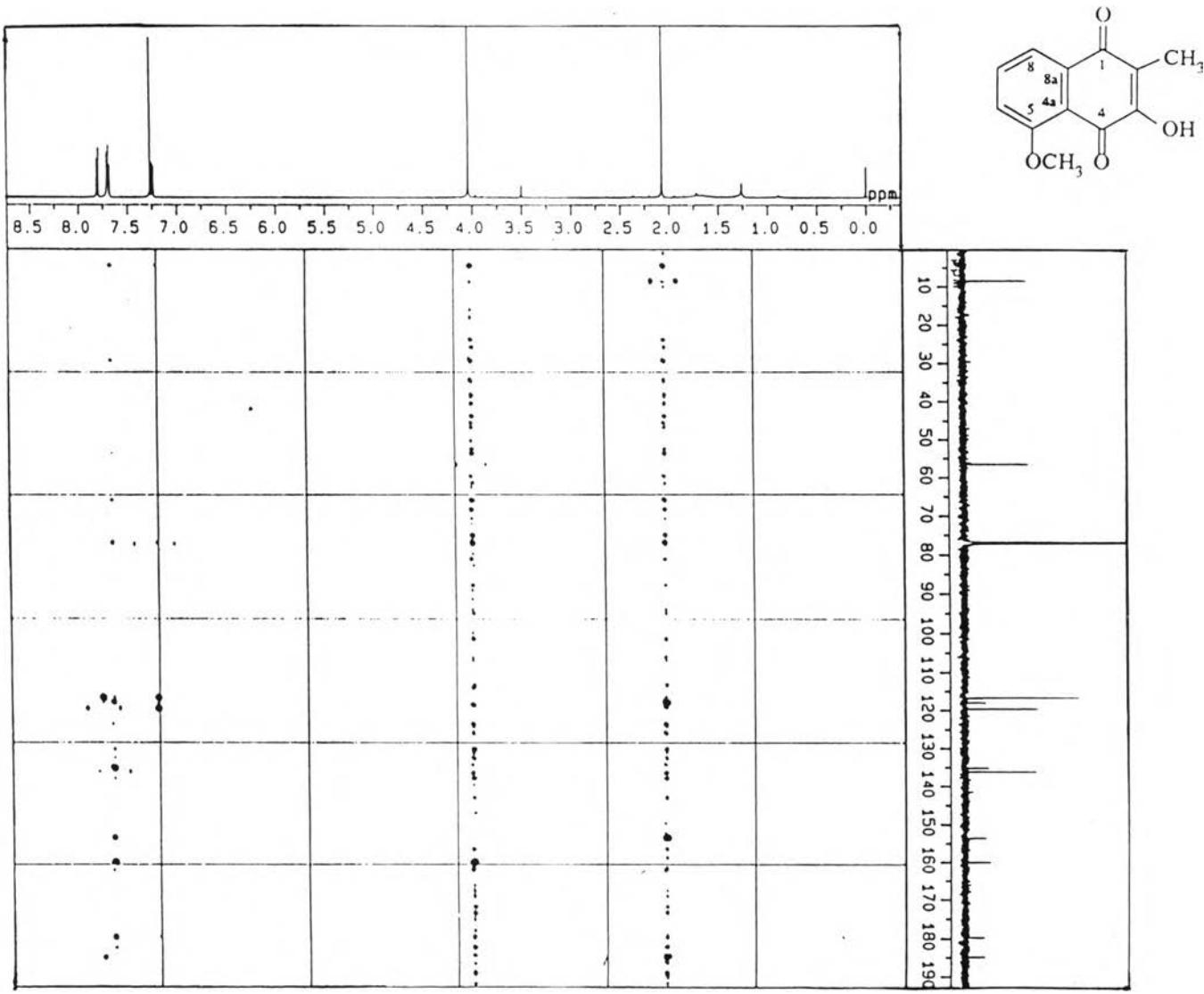


Figure 145 The HMBC spectrum of compound 108 (in CDCl₃)

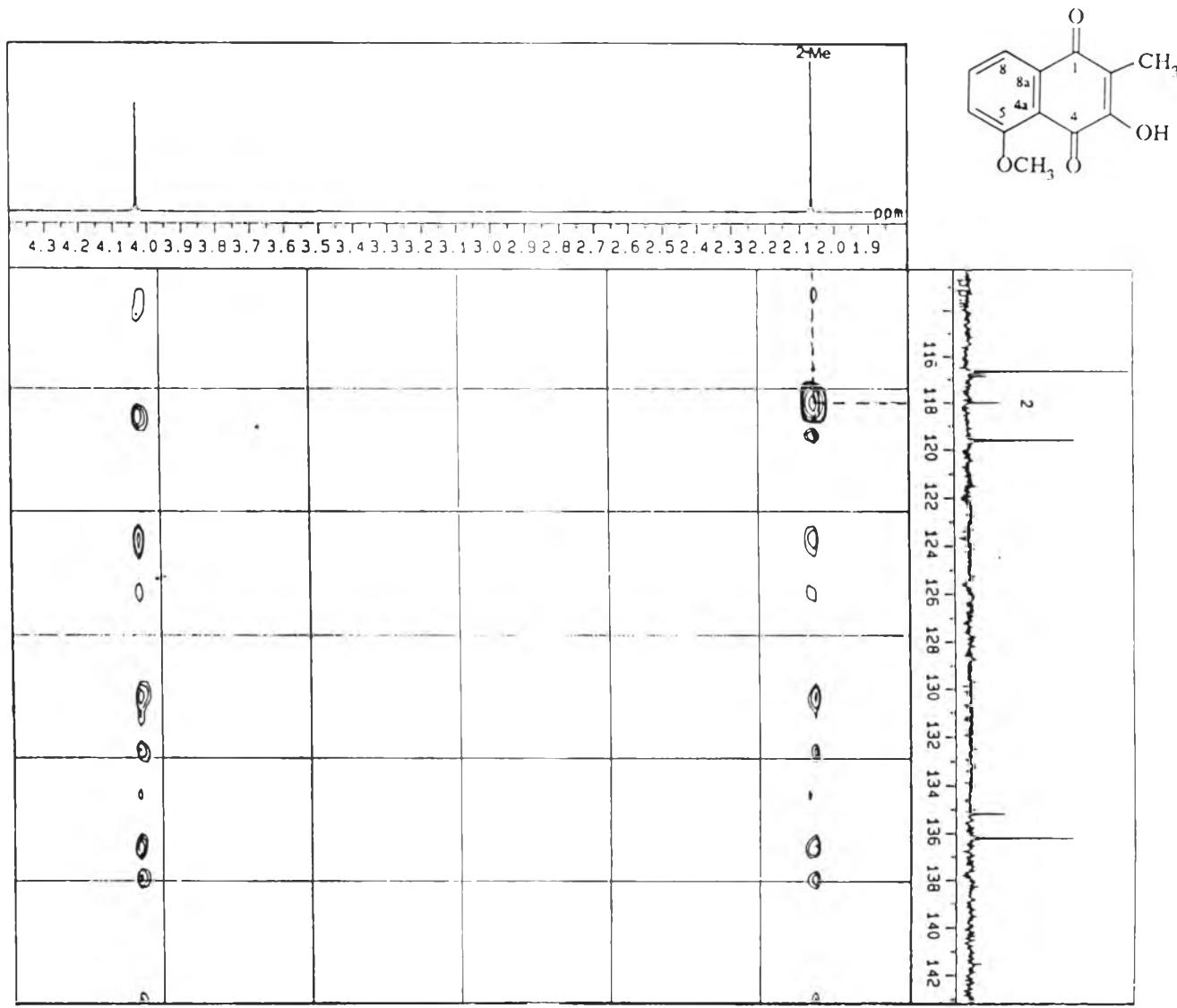


Figure 146 Expansion of the HMBC spectrum of compound 108 (in CDCl_3):

δ_{H} 1.90-4.30 ; δ_{C} 116.00-142.00 ppm

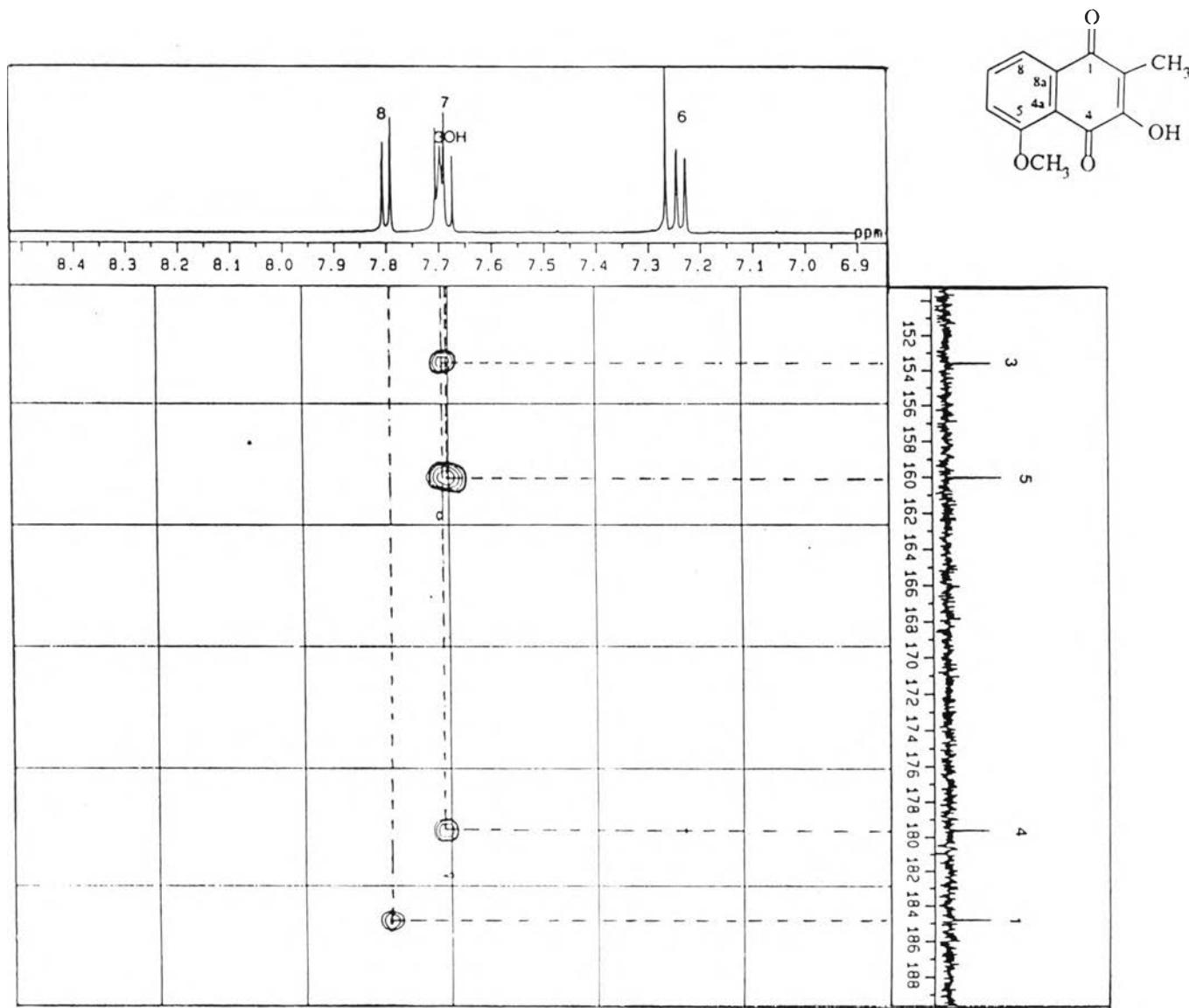


Figure 147 Expansion of the HMBC spectrum of compound 108 (in CDCl_3) :

δ_{H} 6.90-8.40 ; δ_{C} 152.00-188.00 ppm

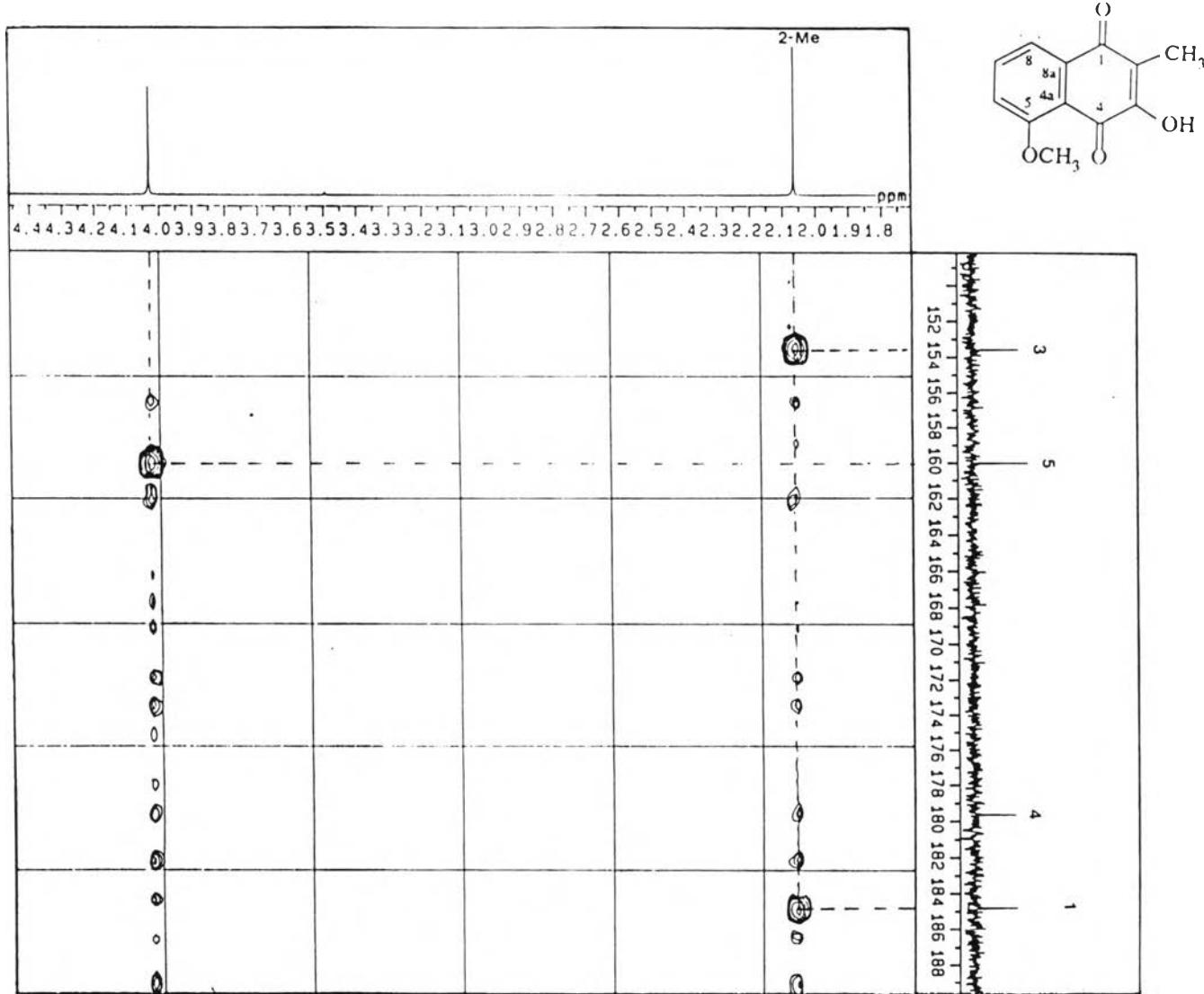


Figure 148 Expansion of the HMBC spectrum of compound 108 (in CDCl_3) :

δ_{H} 1.80-4.40 ; δ_{C} 152.00-188.00 ppm

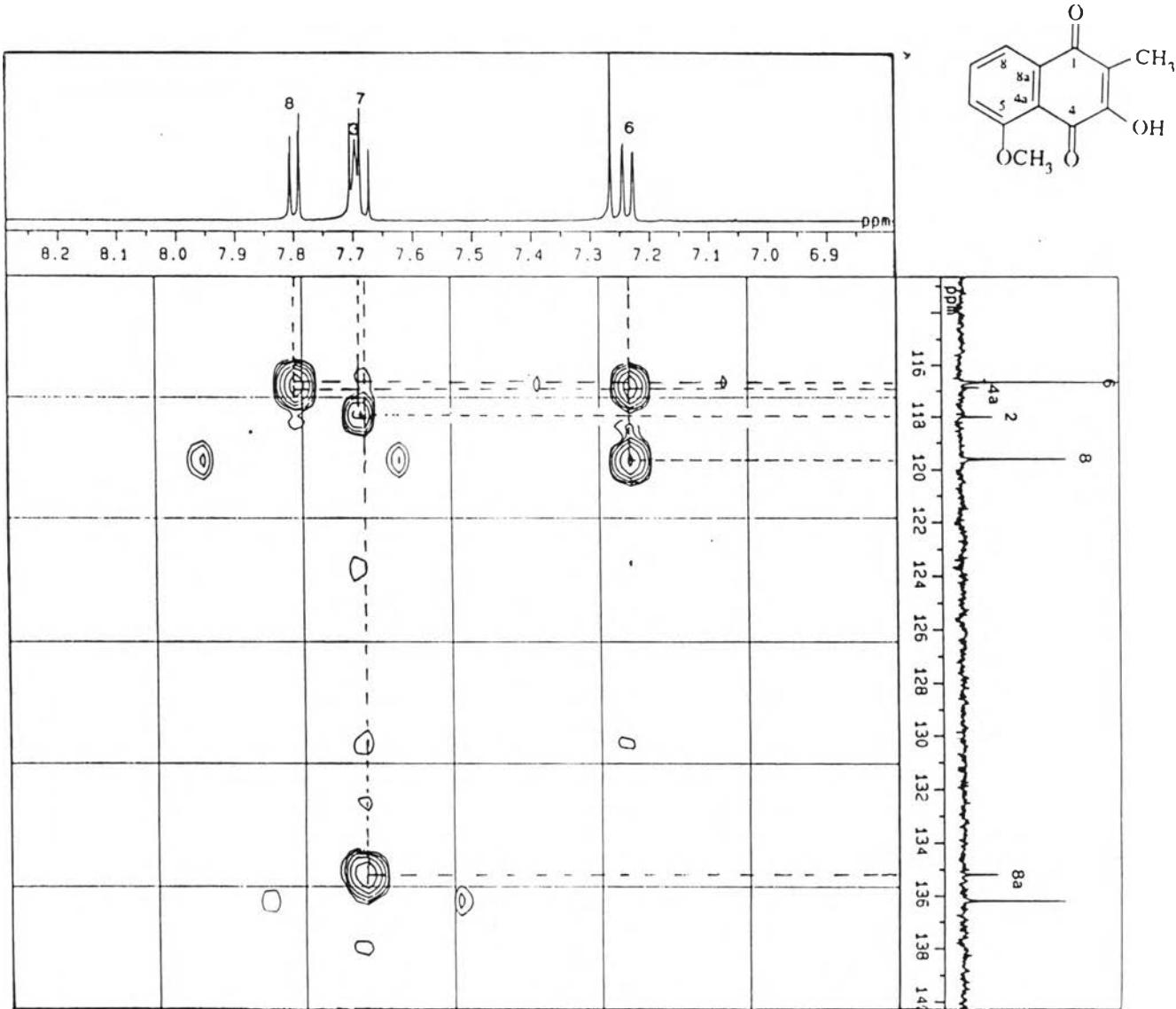


Figure 149 Expansion of the HMBC spectrum of compound 108 (in CDCl_3) :

δ_{H} 6.90-8.20 ; δ_{C} 116.00-140.00 ppm

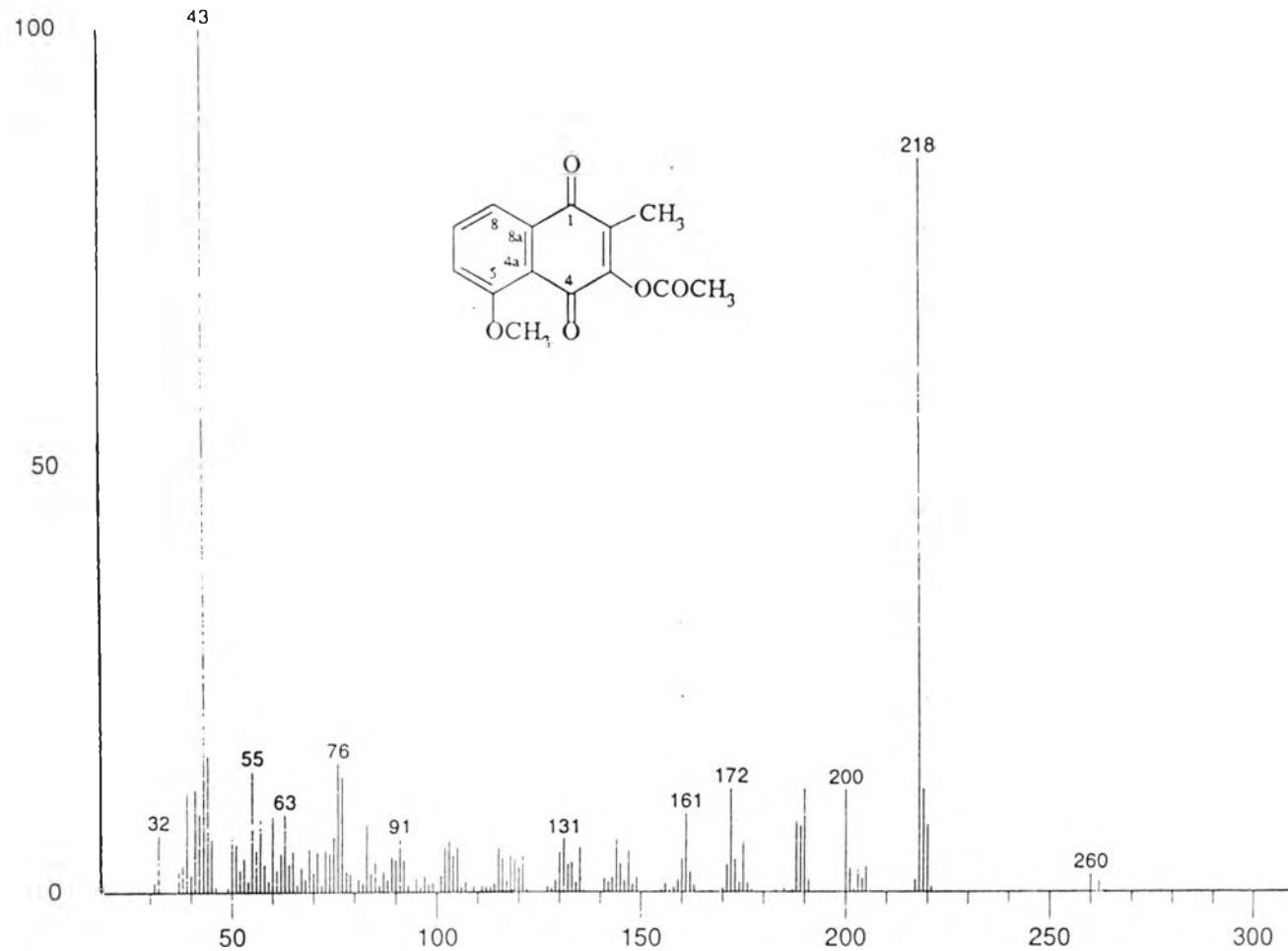


Figure 150 The EI mass spectrum of compound 109

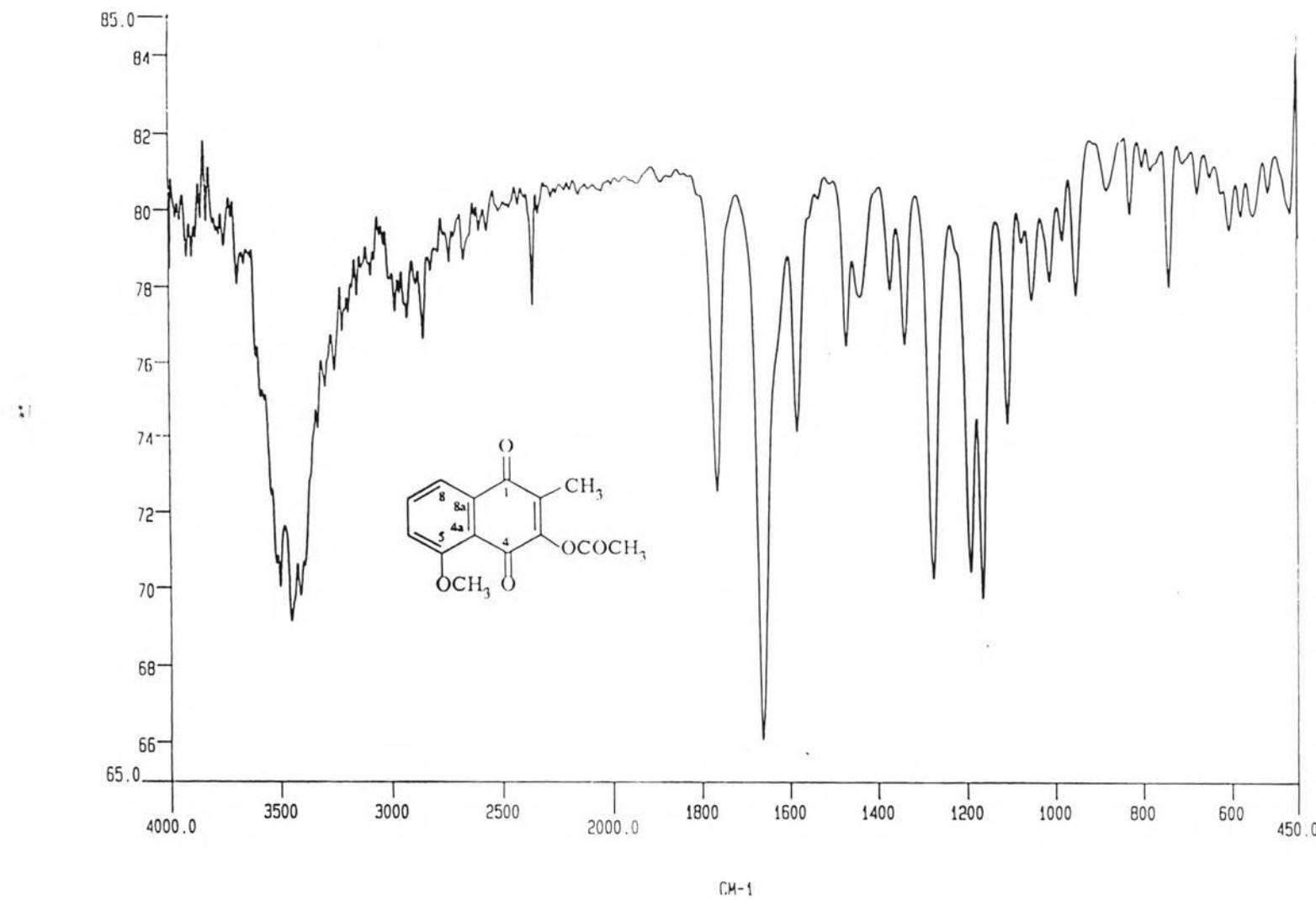


Figure 151 The IR spectrum of compound 109 (in CDCl₃)

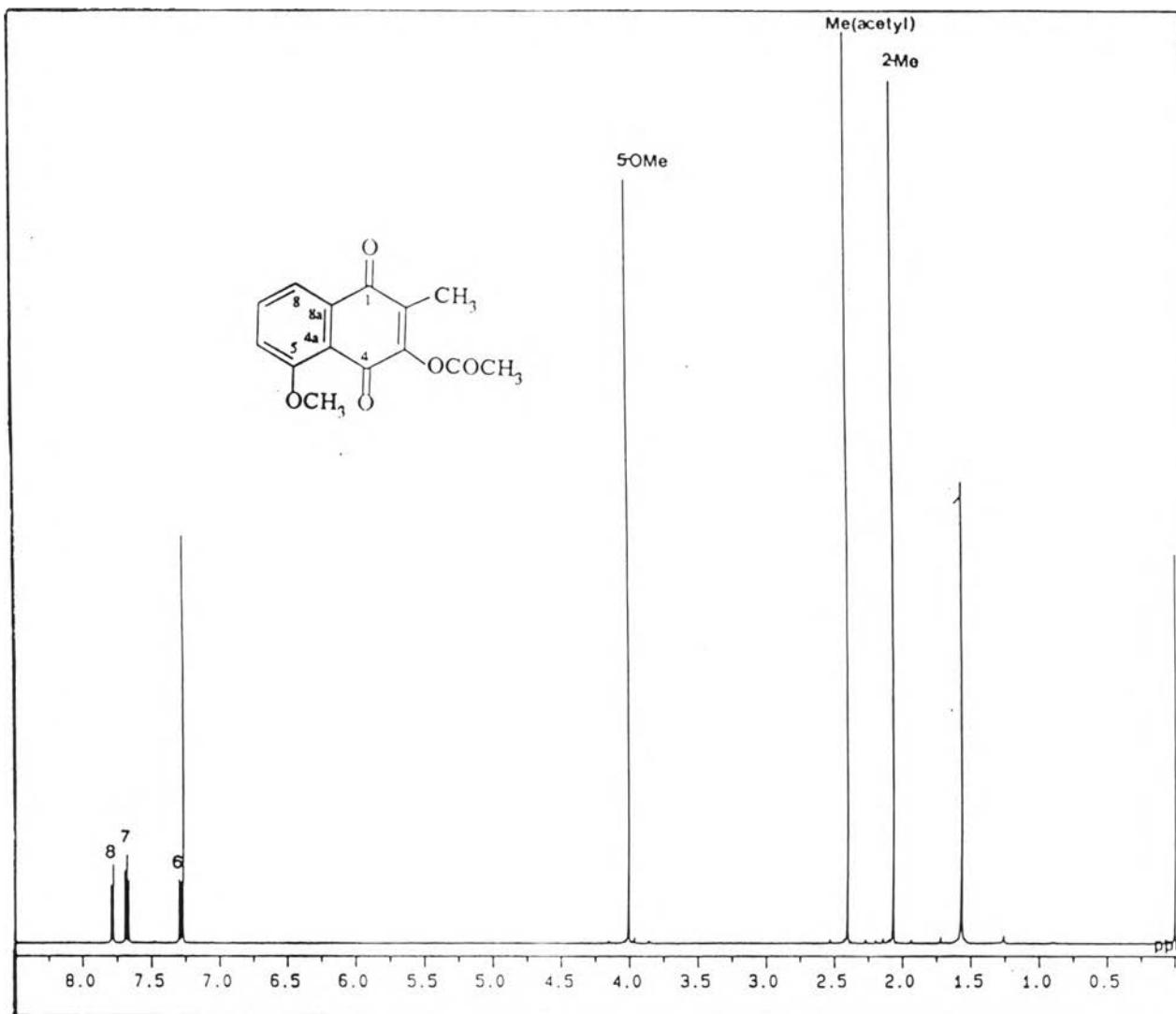


Figure 152 The ^1H NMR (500 MHz) spectrum of compound 109 (in CDCl_3)

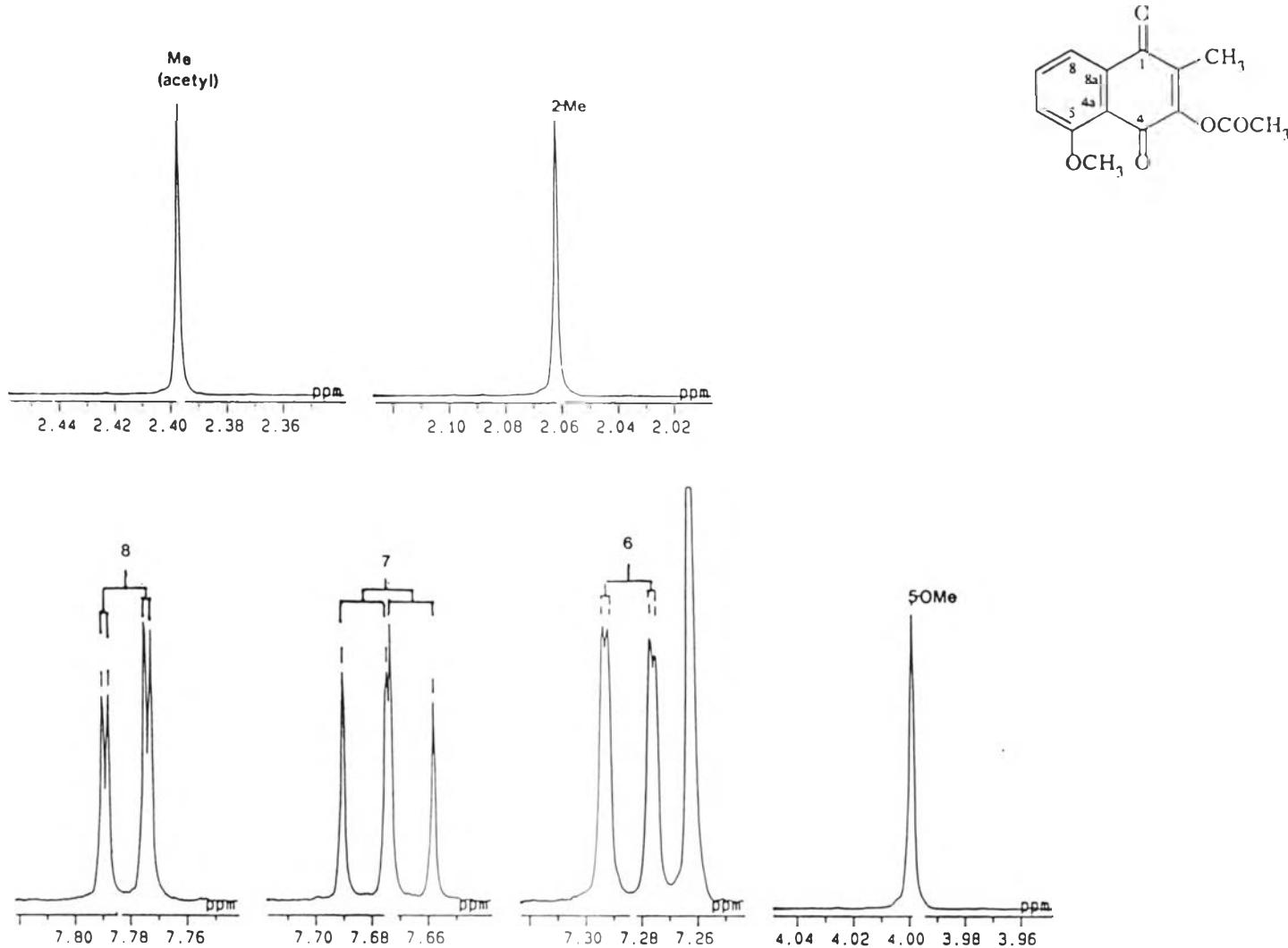


Figure 153 Expansion of the ^1H NMR (500 MHz) spectrum of compound 109
(in CDCl_3) : δ_{H} 2.02-7.80 ppm

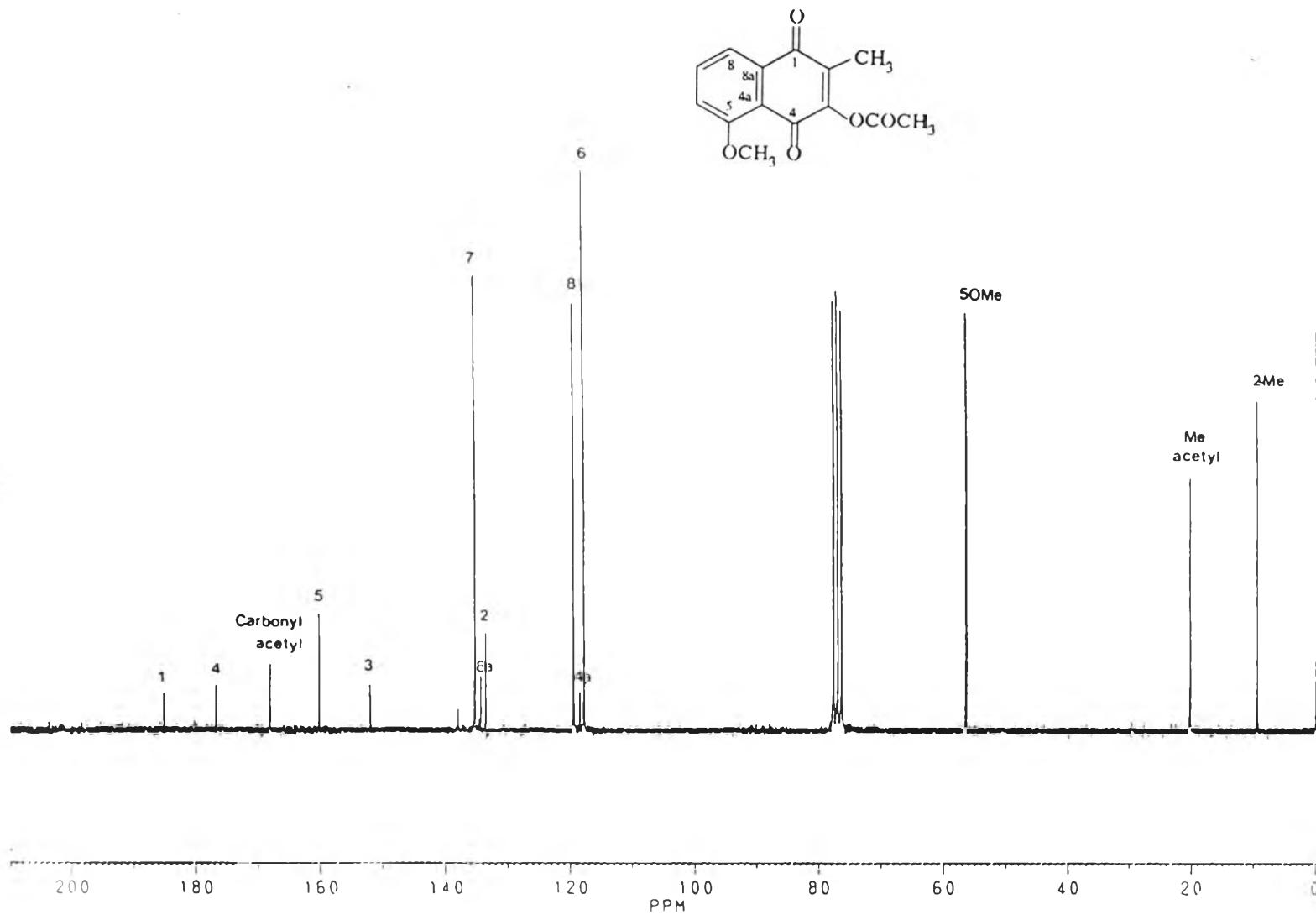


Figure 154 The ^{13}C NMR (50 MHz) spectrum of compound 109 (in CDCl_3)

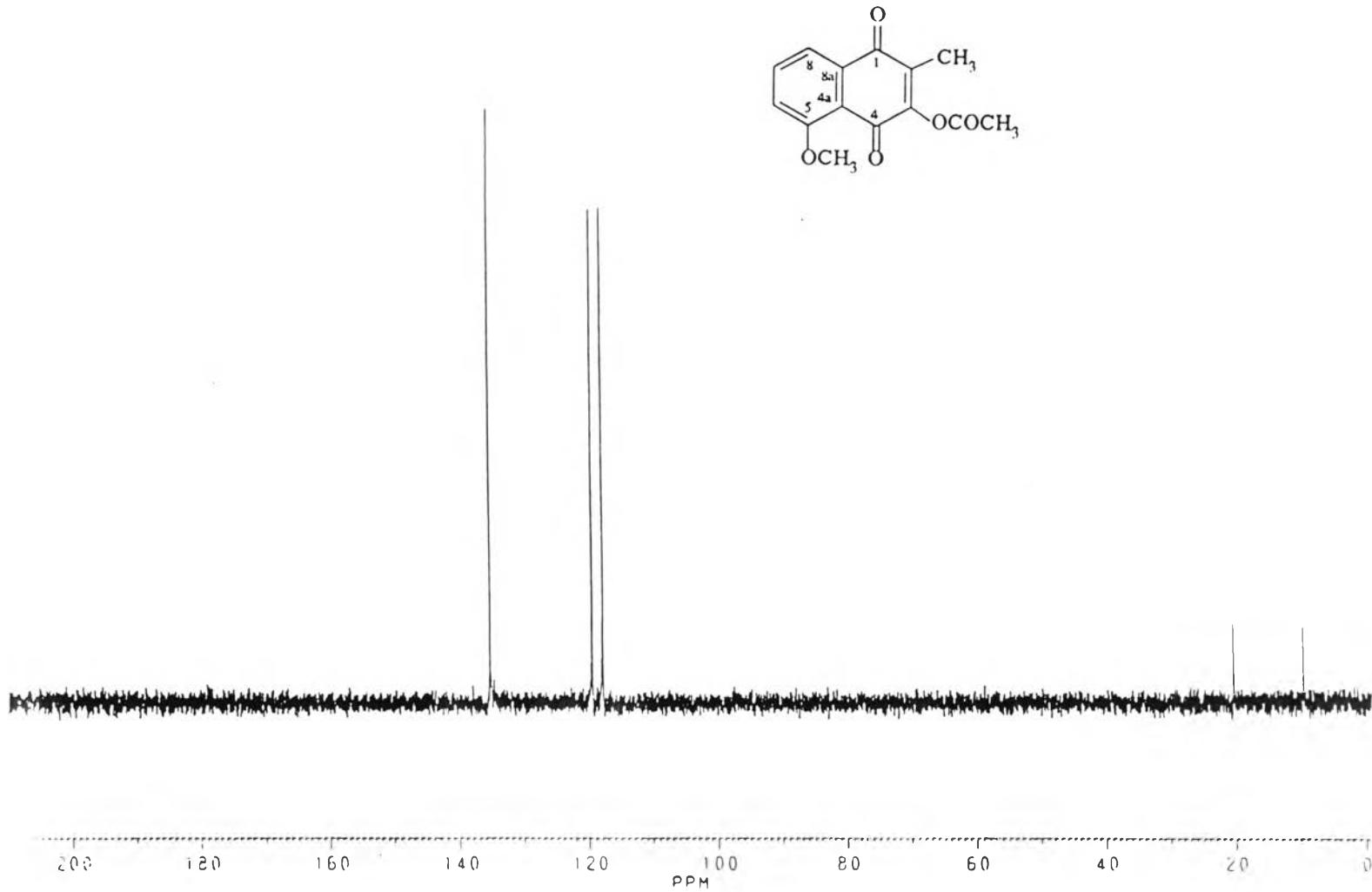


Figure 155 The DEPT 90 (50 MHz) spectrum of compound 109 (in CDCl_3)

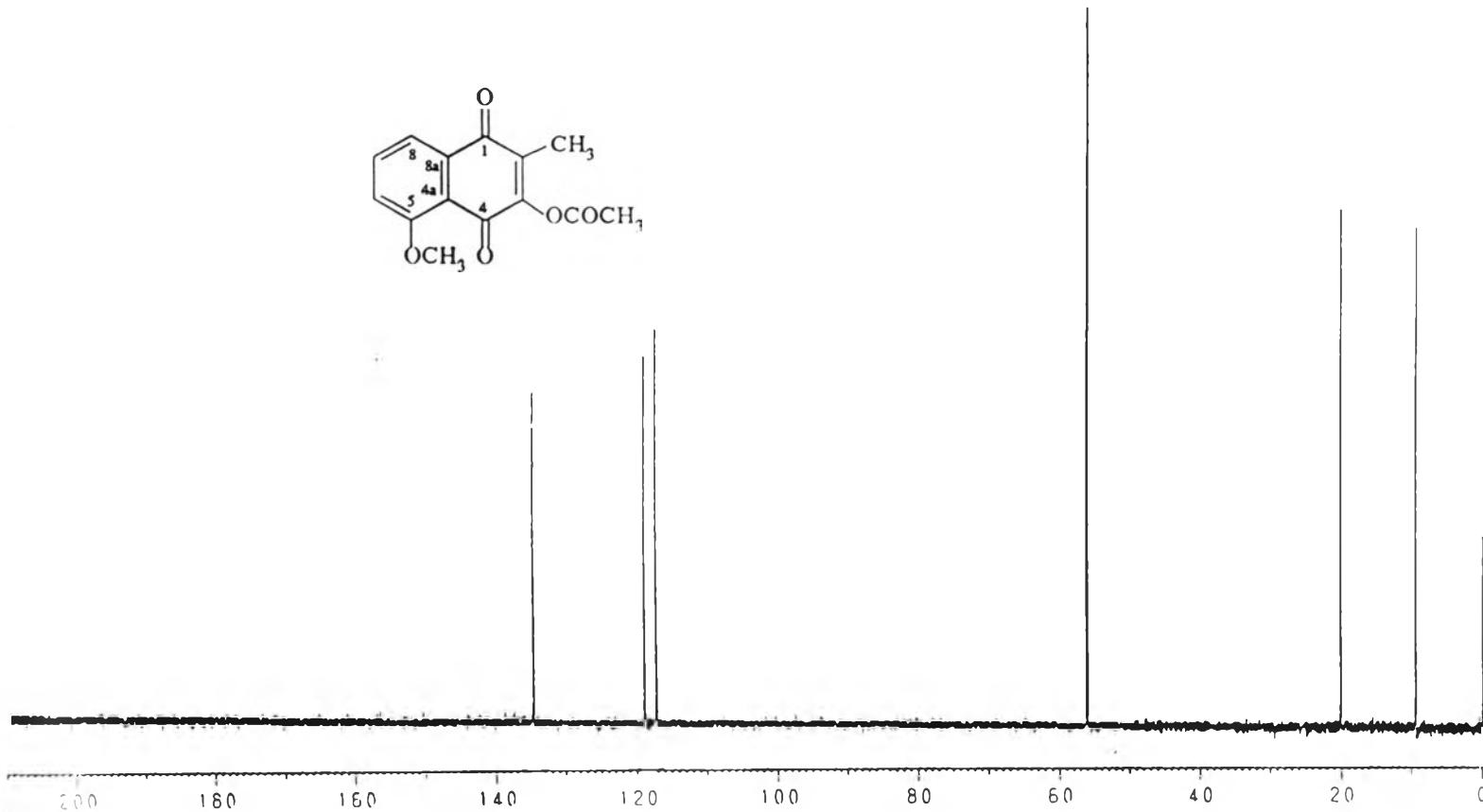


Figure 156 The DEPT 135 (50 MHz) spectrum of compound 109 (in CDCl_3)

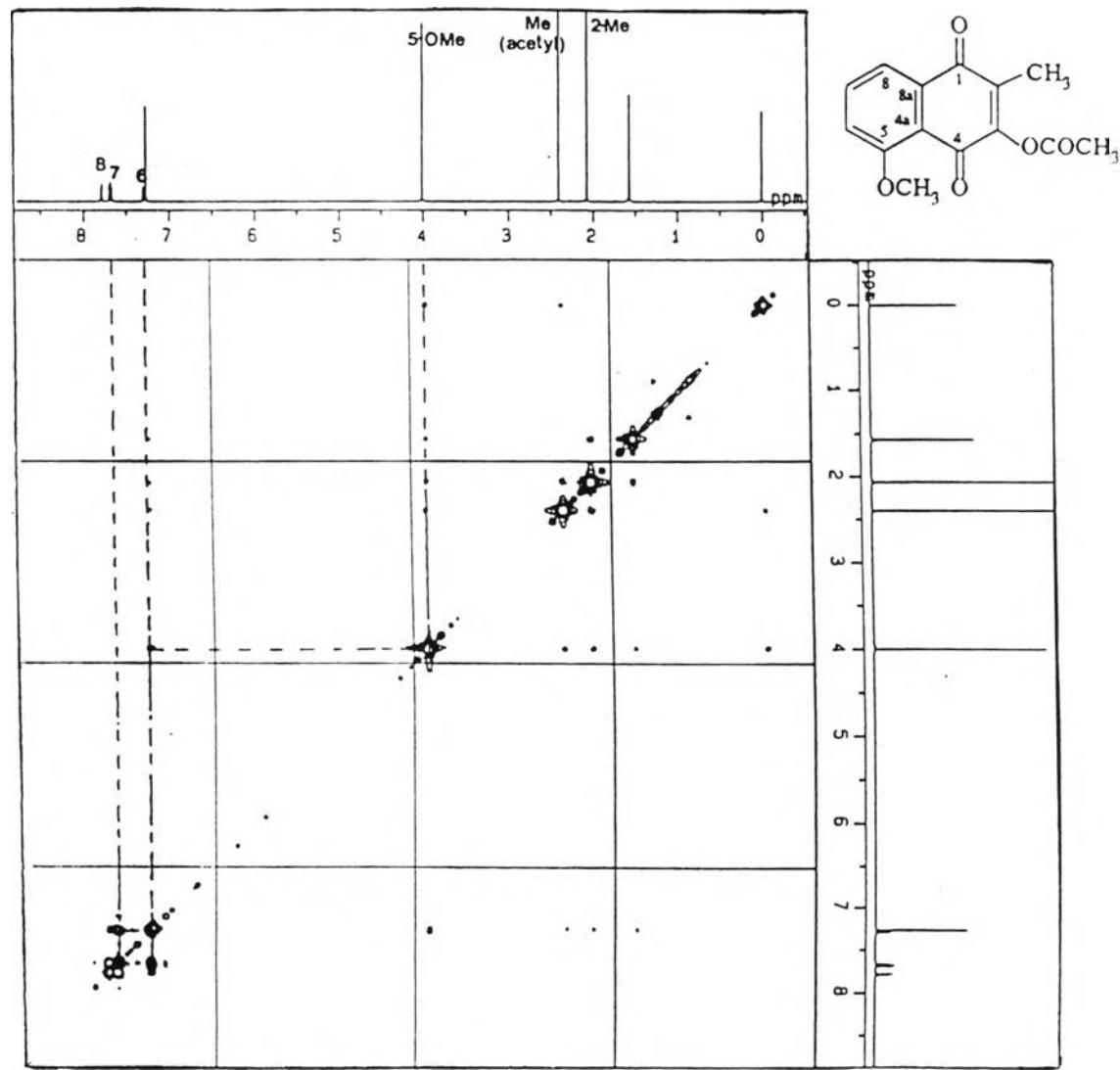


Figure 157 The ^1H - ^1H COSY (500 MHz) spectrum of compound 109 (in CDCl_3)

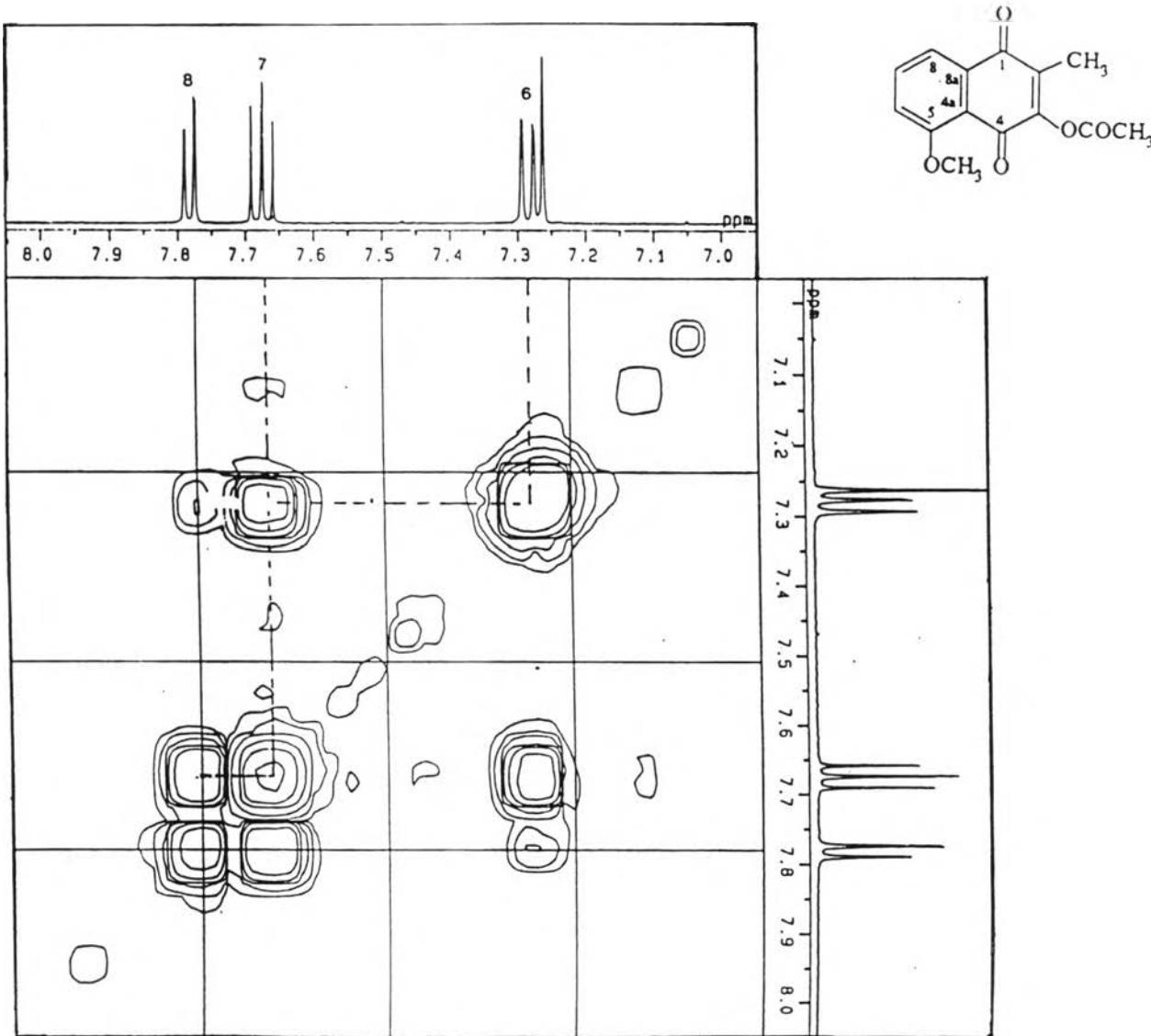


Figure 158 Expansion of the ^1H - ^1H COSY (500 MHz) spectrum of compound 109
(in CDCl_3) : δ_{H} 7.00-8.00 ppm

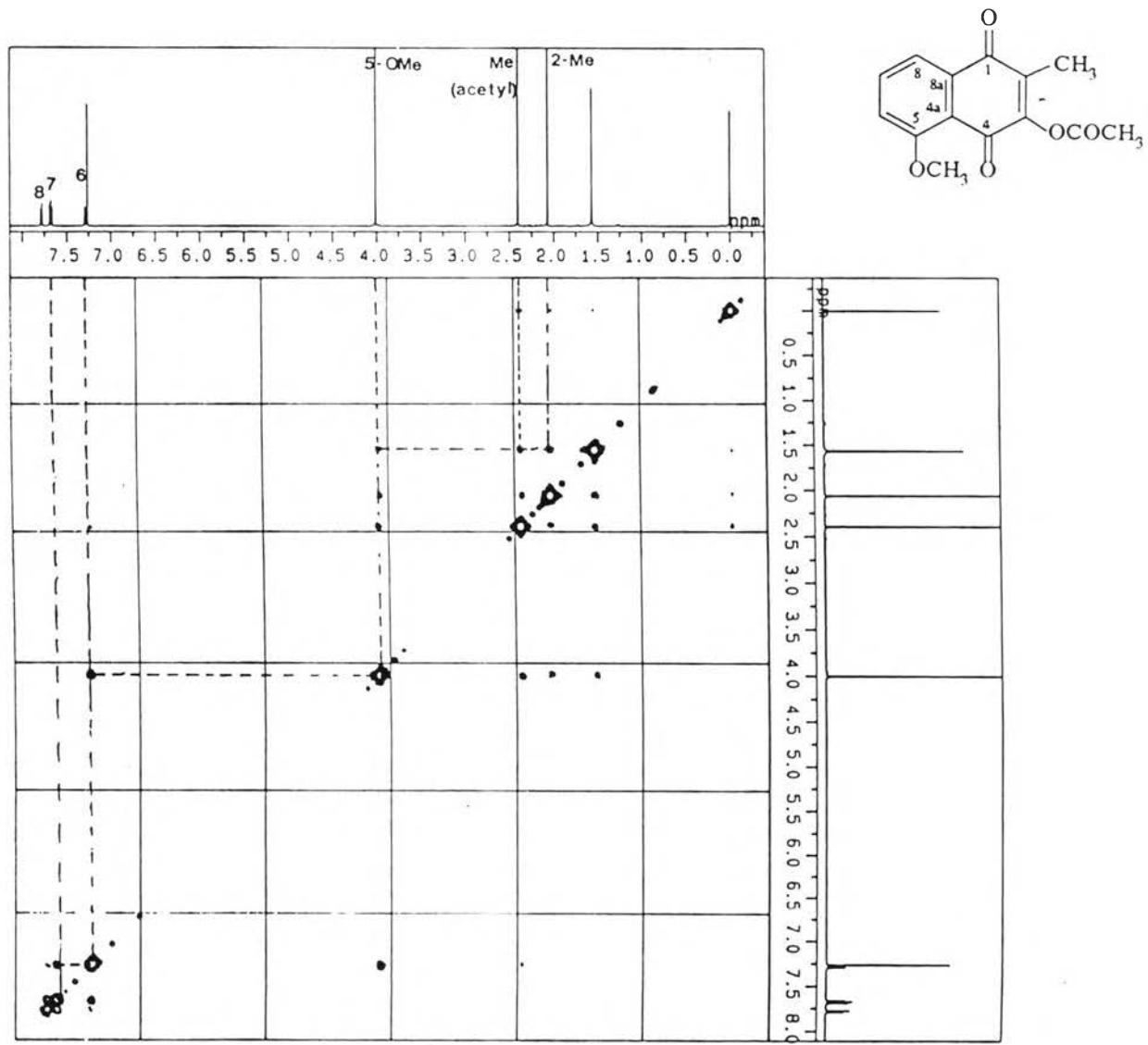


Figure 159 The NOESY (500 MHz) spectrum of compound 109 (in CDCl₃)

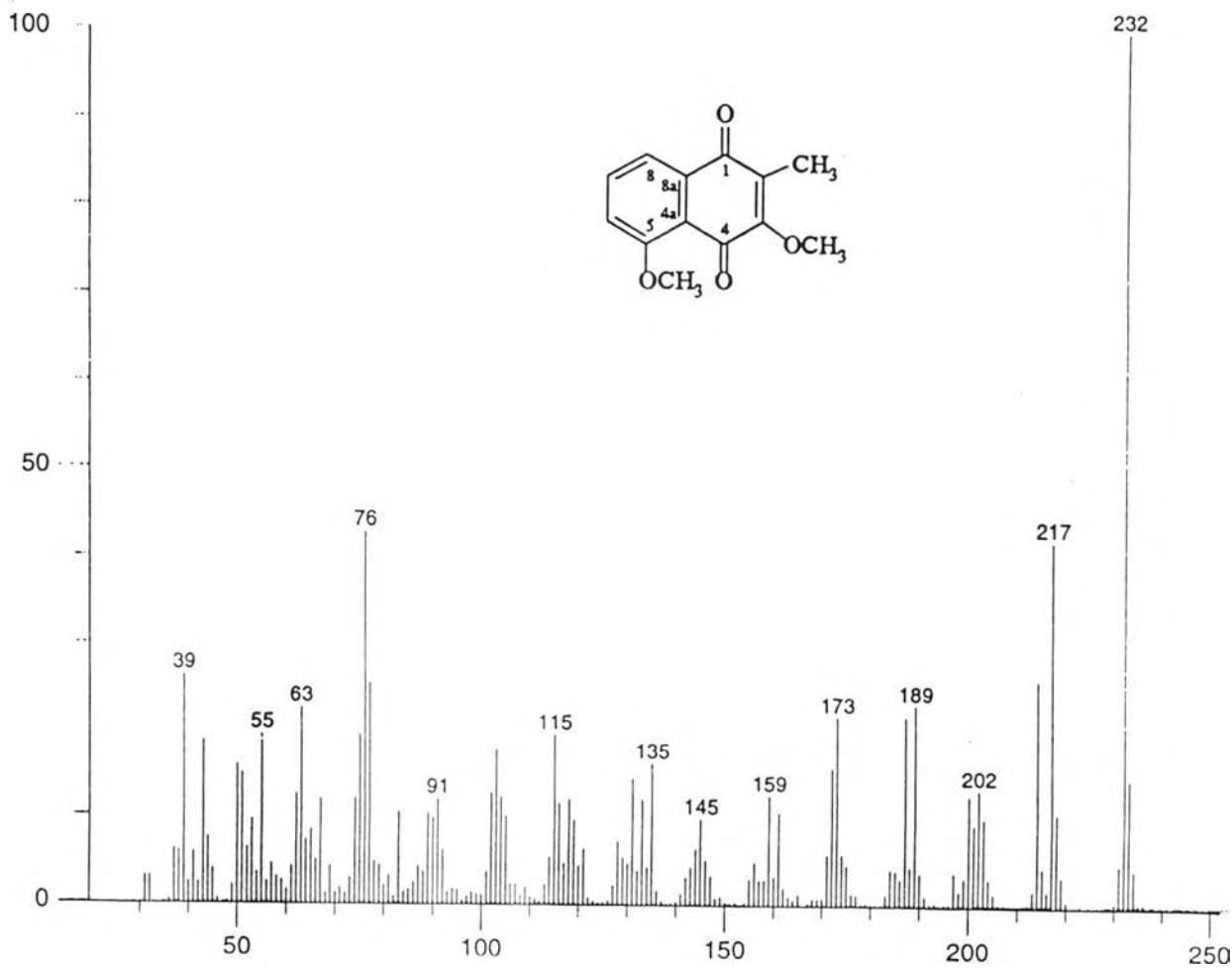


Figure 160 The EI mass spectrum of compound 110

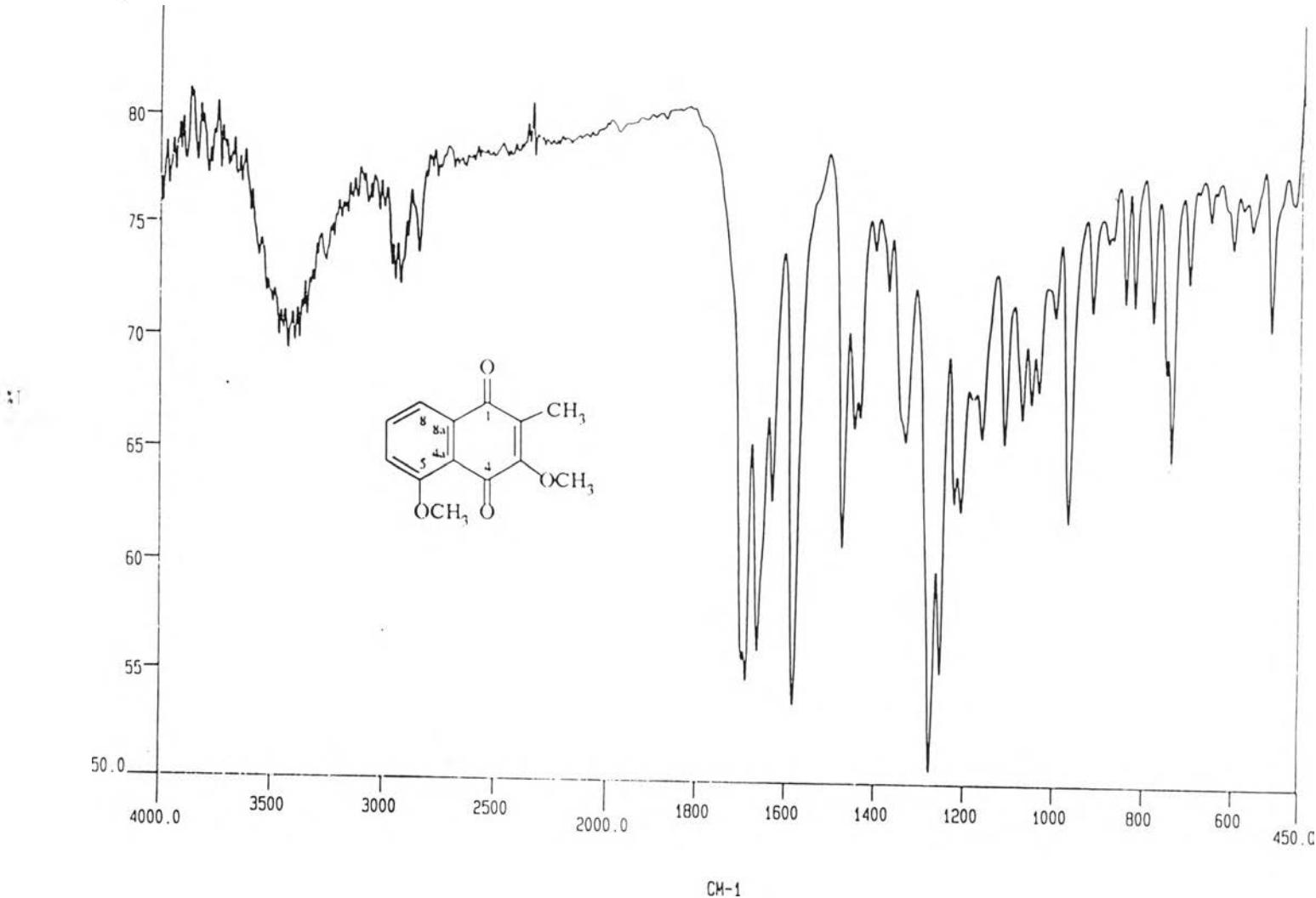


Figure 161 The IR spectrum of compound 110 (in KBr disc)

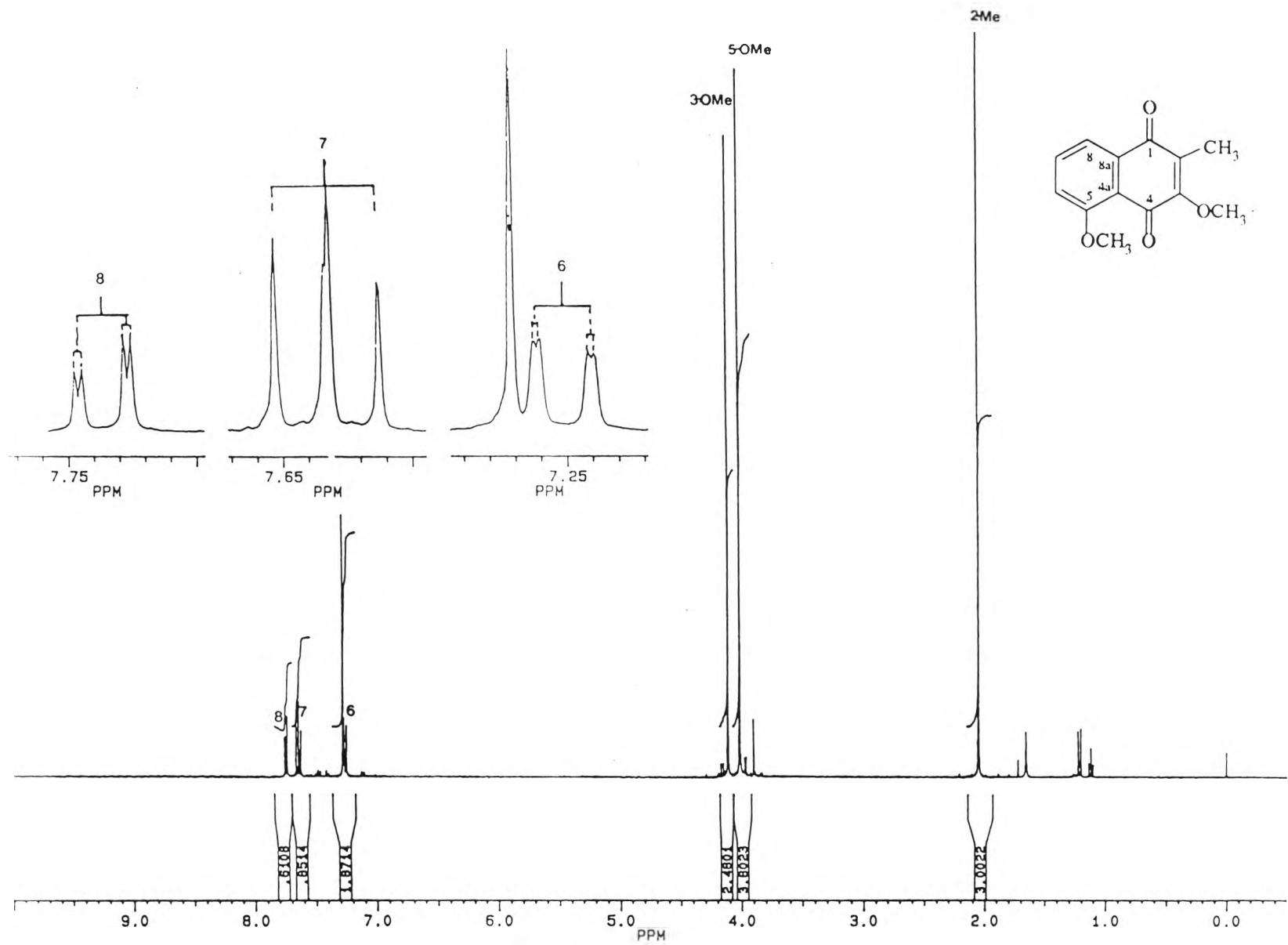


Figure 162 The ^1H NMR (400 MHz) spectrum of compound 110 (in CDCl₃)

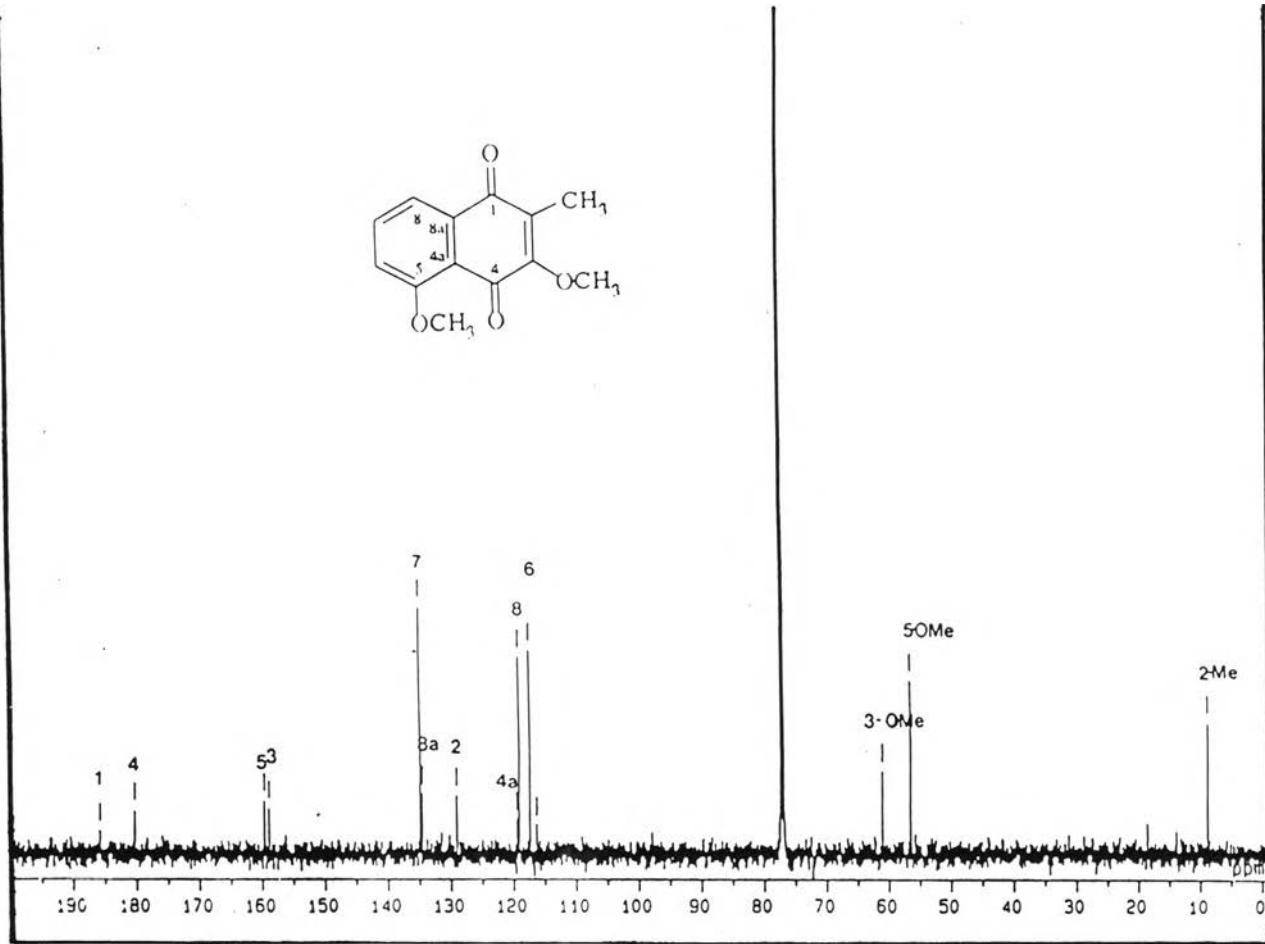


Figure 163 The ^{13}C NMR (125 MHz) of compound 110 (in CDCl_3)

P0M-DEPT135

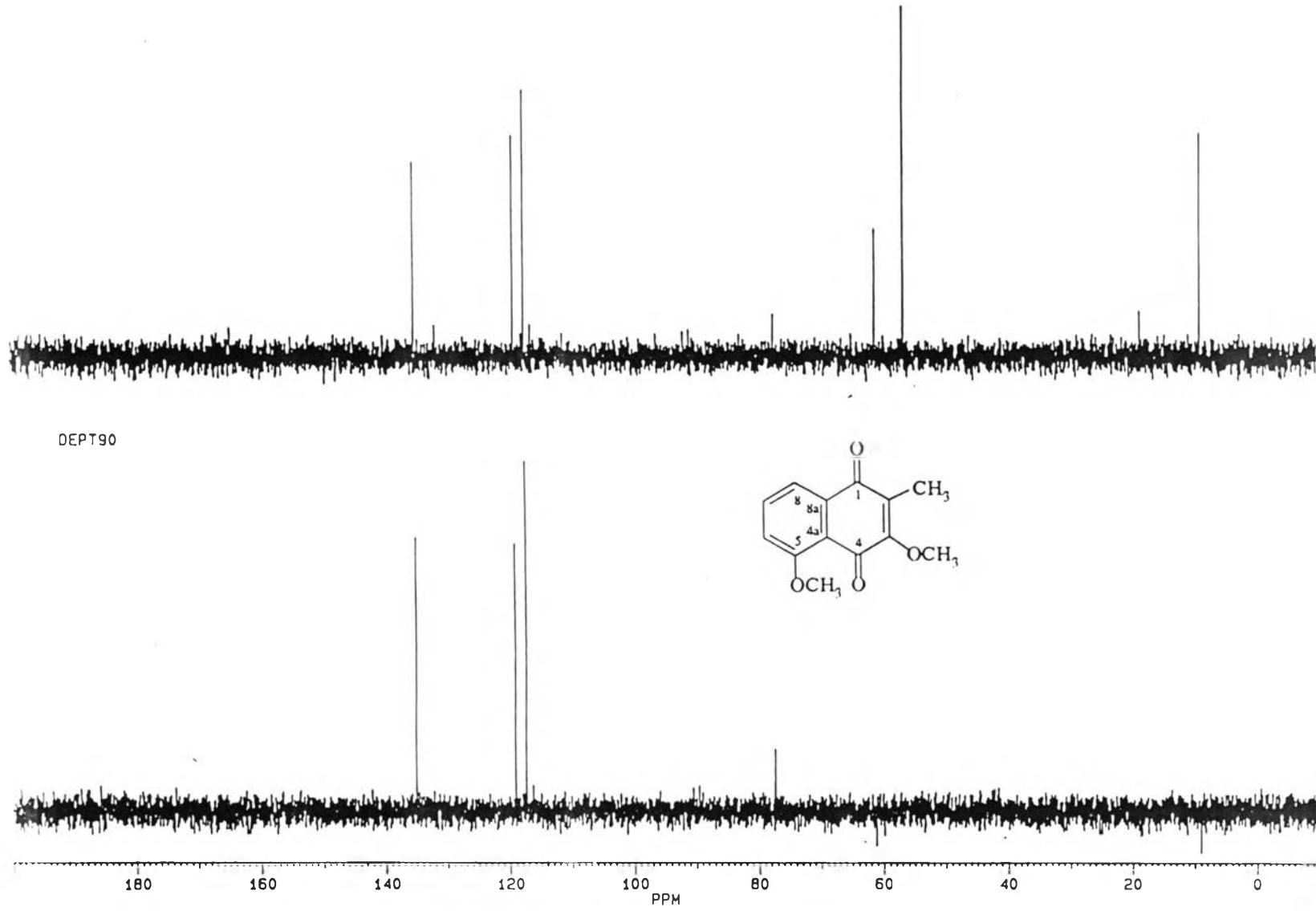


Figure 164 The DEPT (100 MHz) spectrum of compound 110 (in CDCl₃)

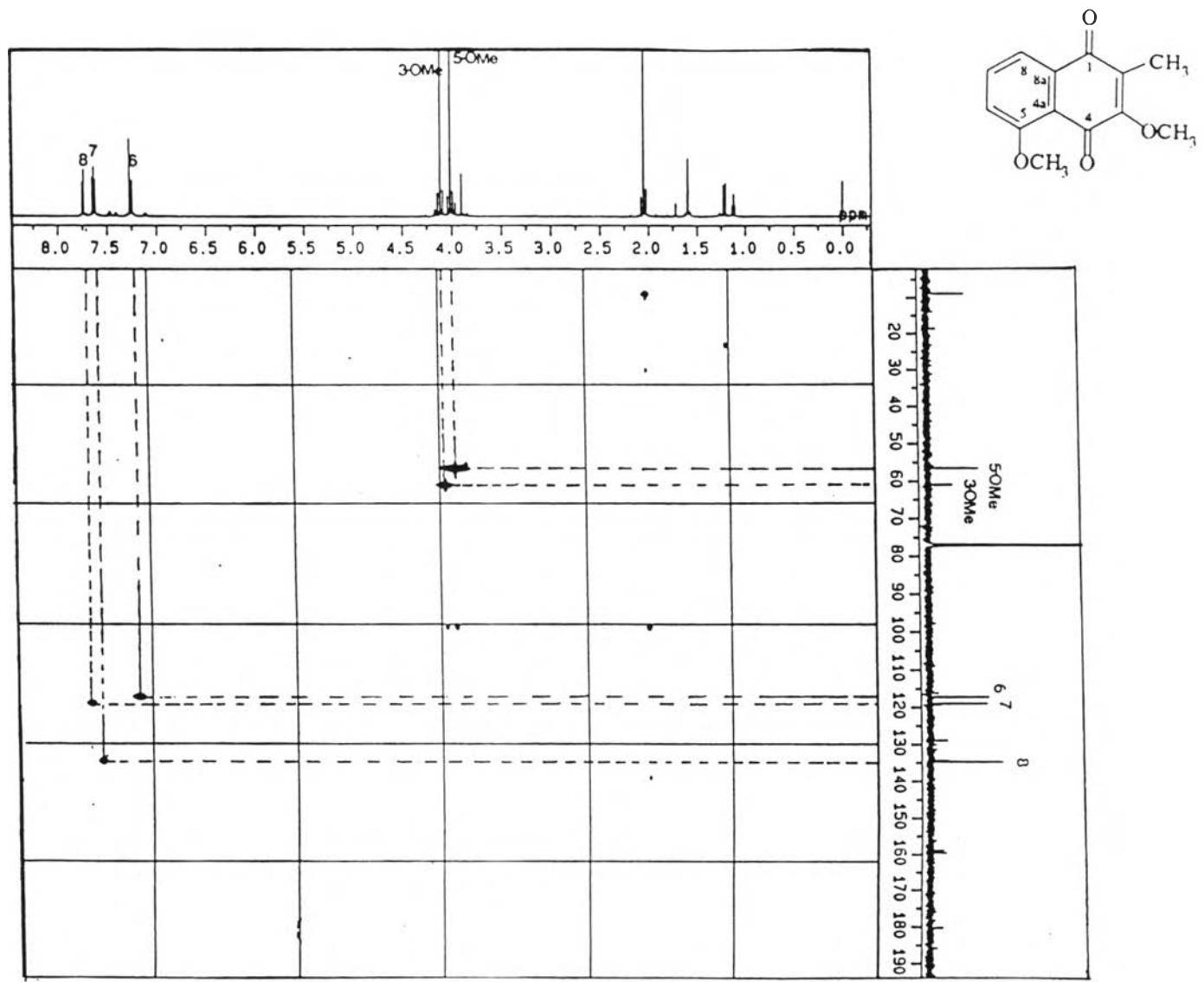


Figure 165 The HMQC spectrum of compound 110 (in CDCl_3)

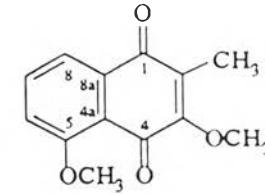
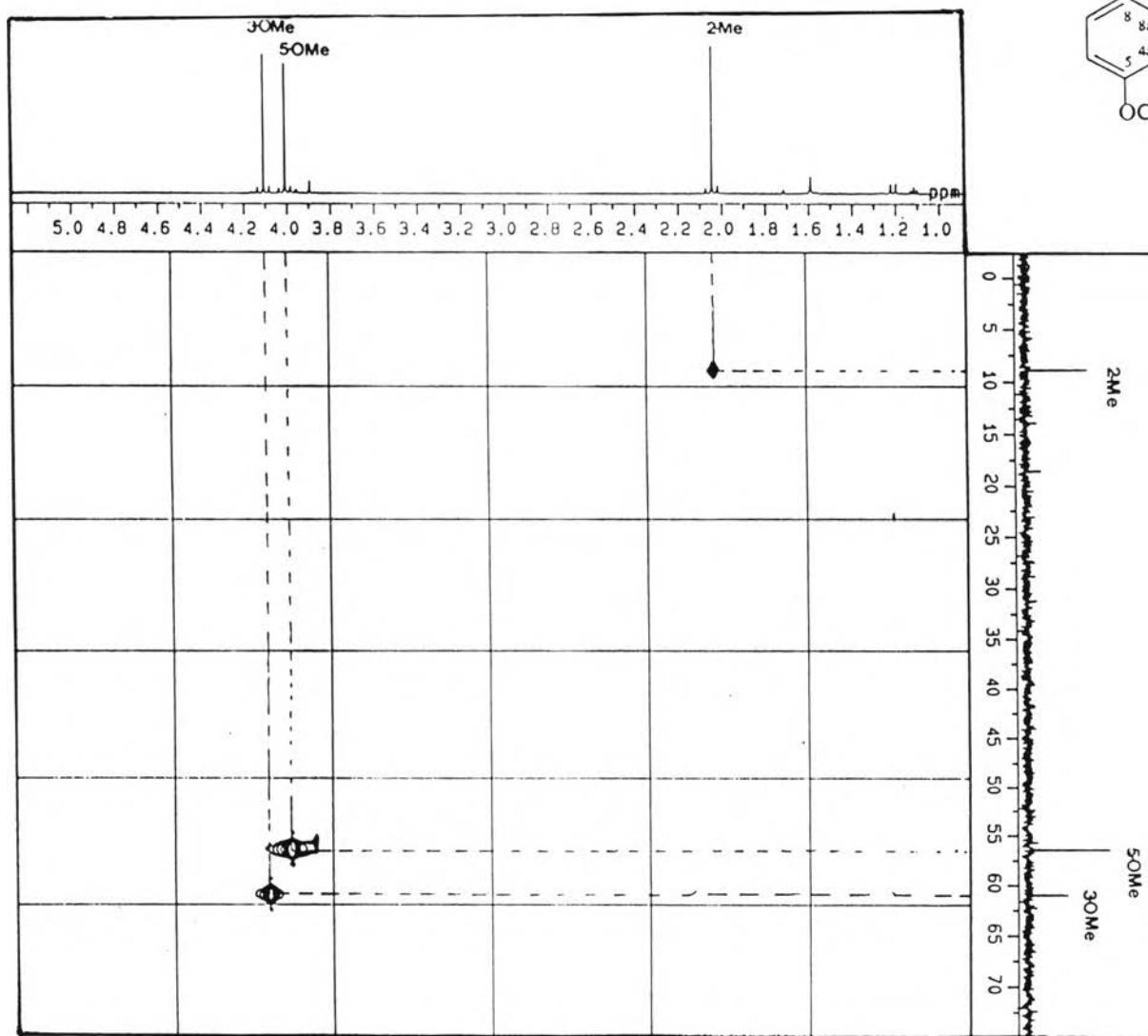


Figure 166 Expansion of the HMQC spectrum of compound 110 (in CDCl_3) :

δ_{H} 1.00-5.00 ; δ_{C} 0.00-70.00 ppm

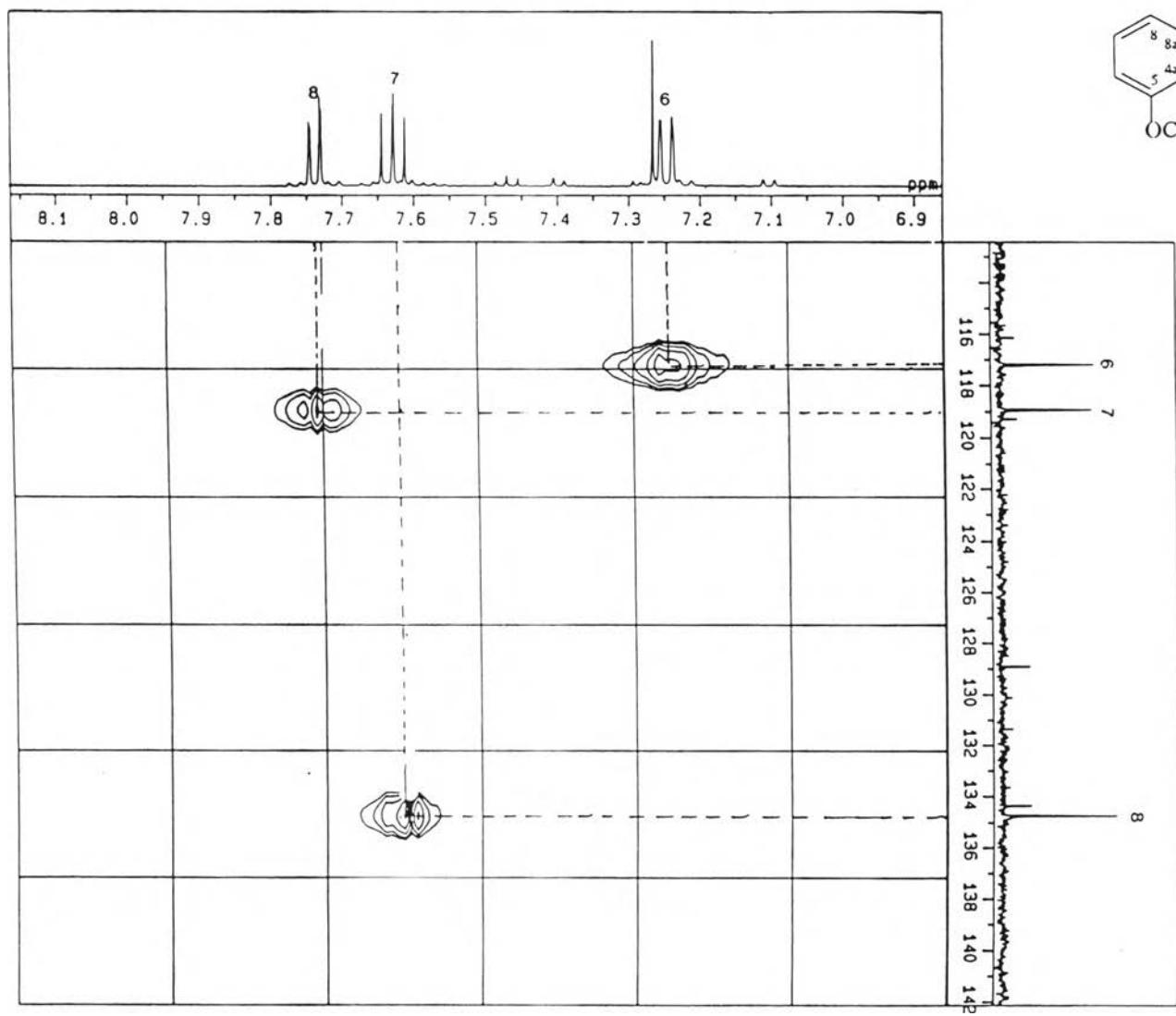
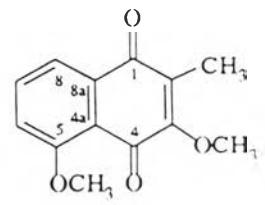


Figure 167 Expansion of the HMQC spectrum of compound 110 (in CDCl₃) :

δ_{H} 6.90-8.10 ; δ_{C} 114.00-142.00 ppm

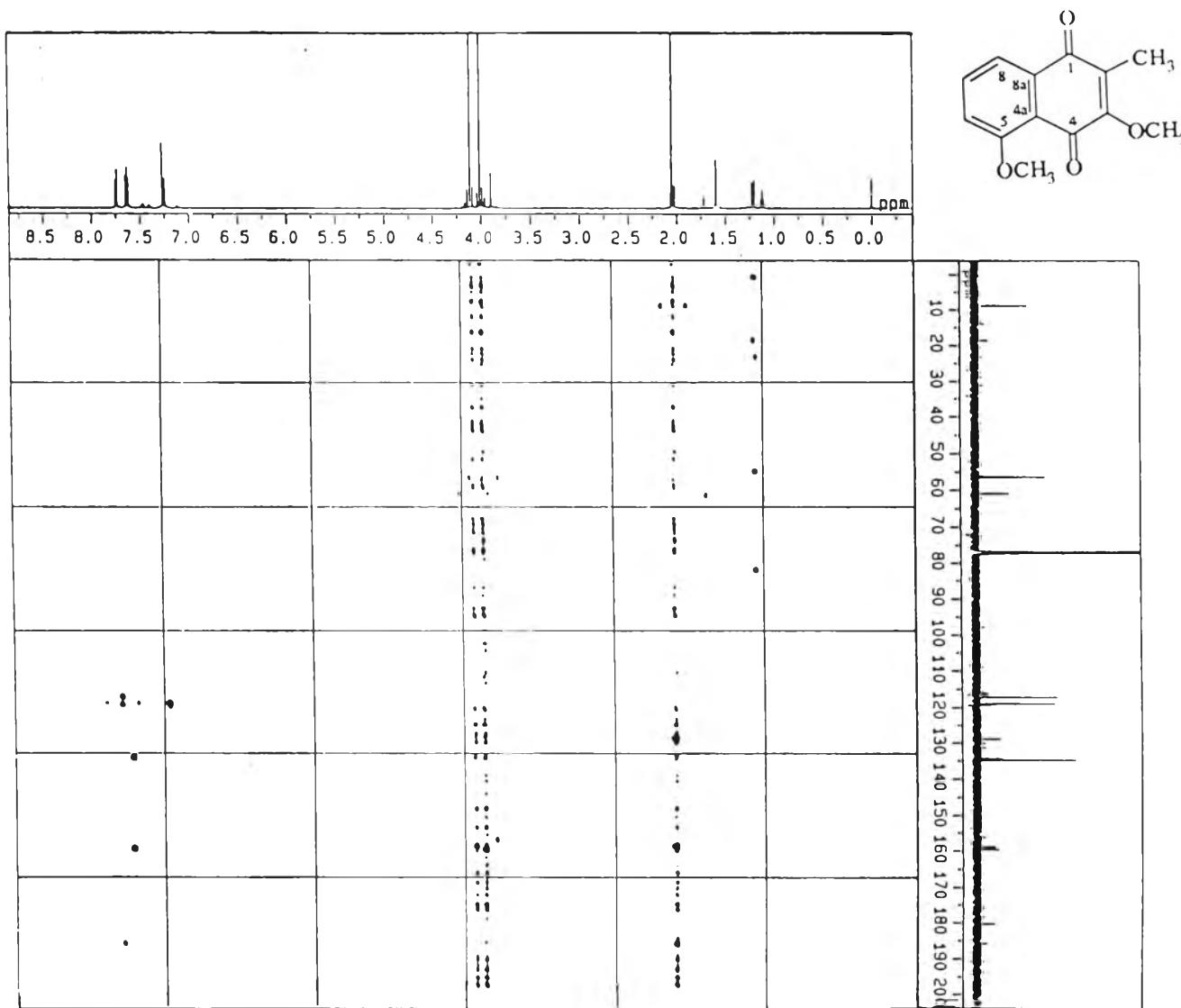


Figure 168 The HMBC spectrum of compound 110 (in CDCl₃)

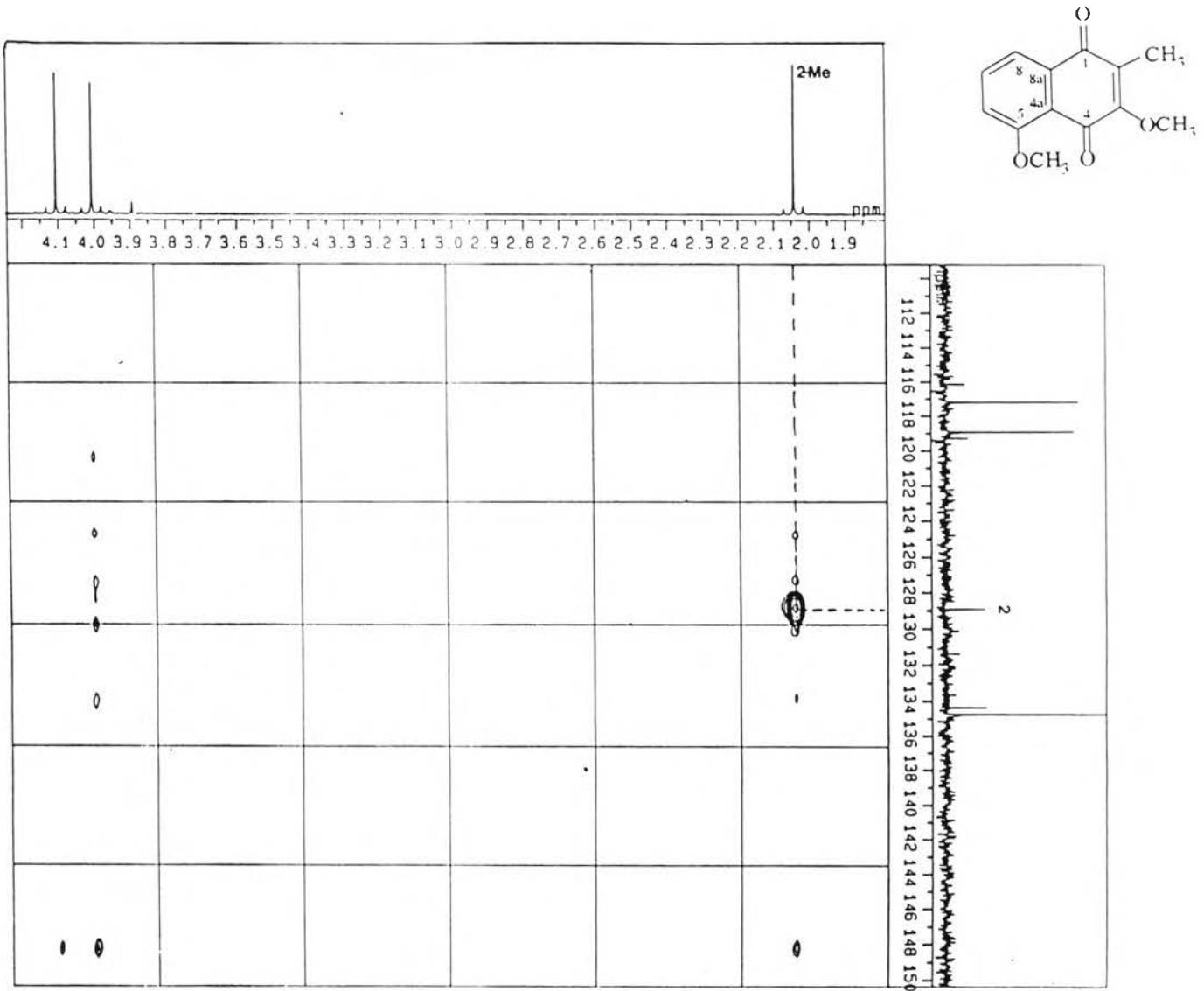


Figure 169 Expansion of the HMBC spectrum of compound 110 (in CDCl₃) :

δ_{H} 1.90-4.10 ; δ_{C} 112.00-150.00 ppm

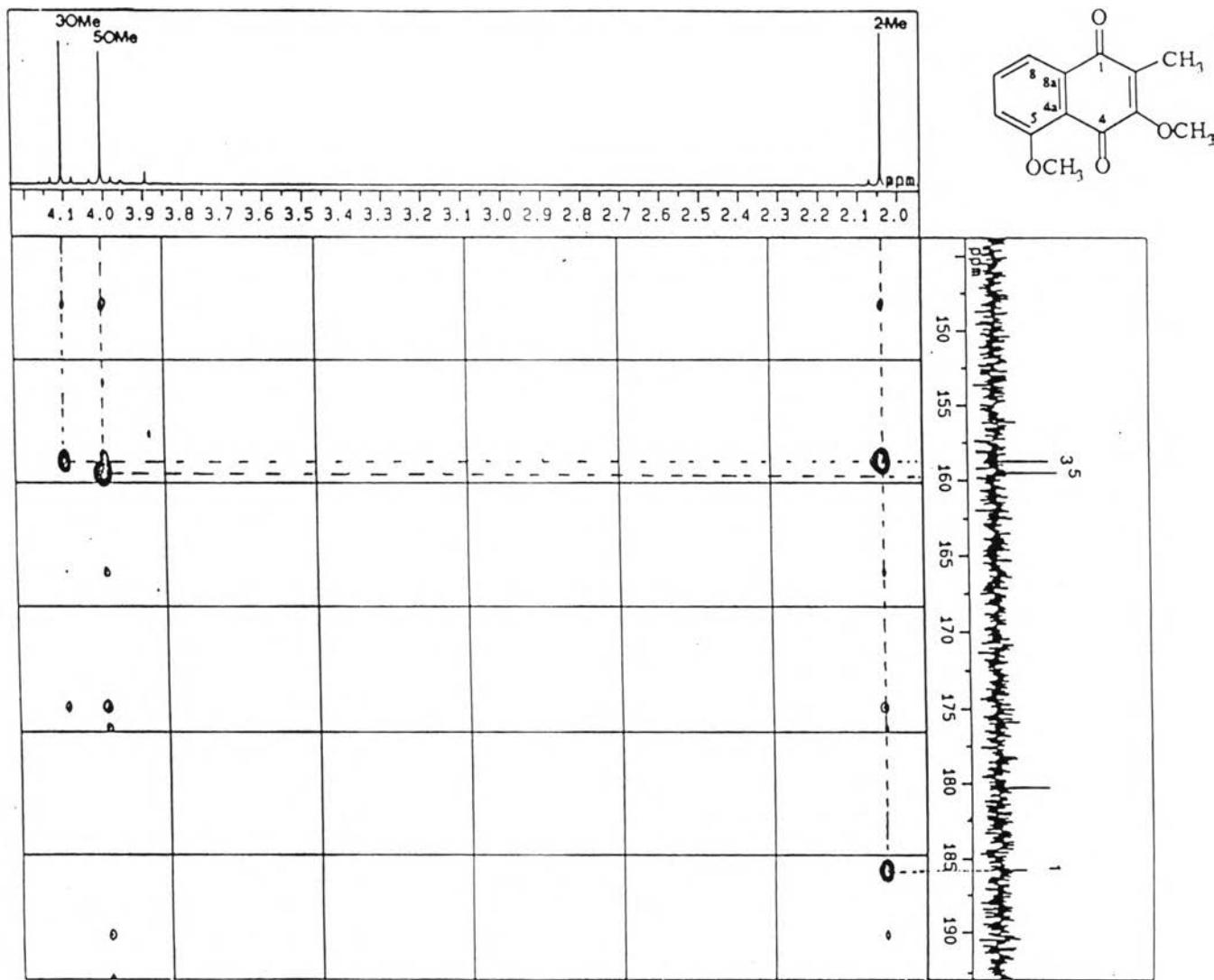


Figure 170 Expansion of the HMBC spectrum of compound 110 (in CDCl_3) :

δ_{H} 2.00-4.10 ; δ_{C} 145.00-190.00 ppm

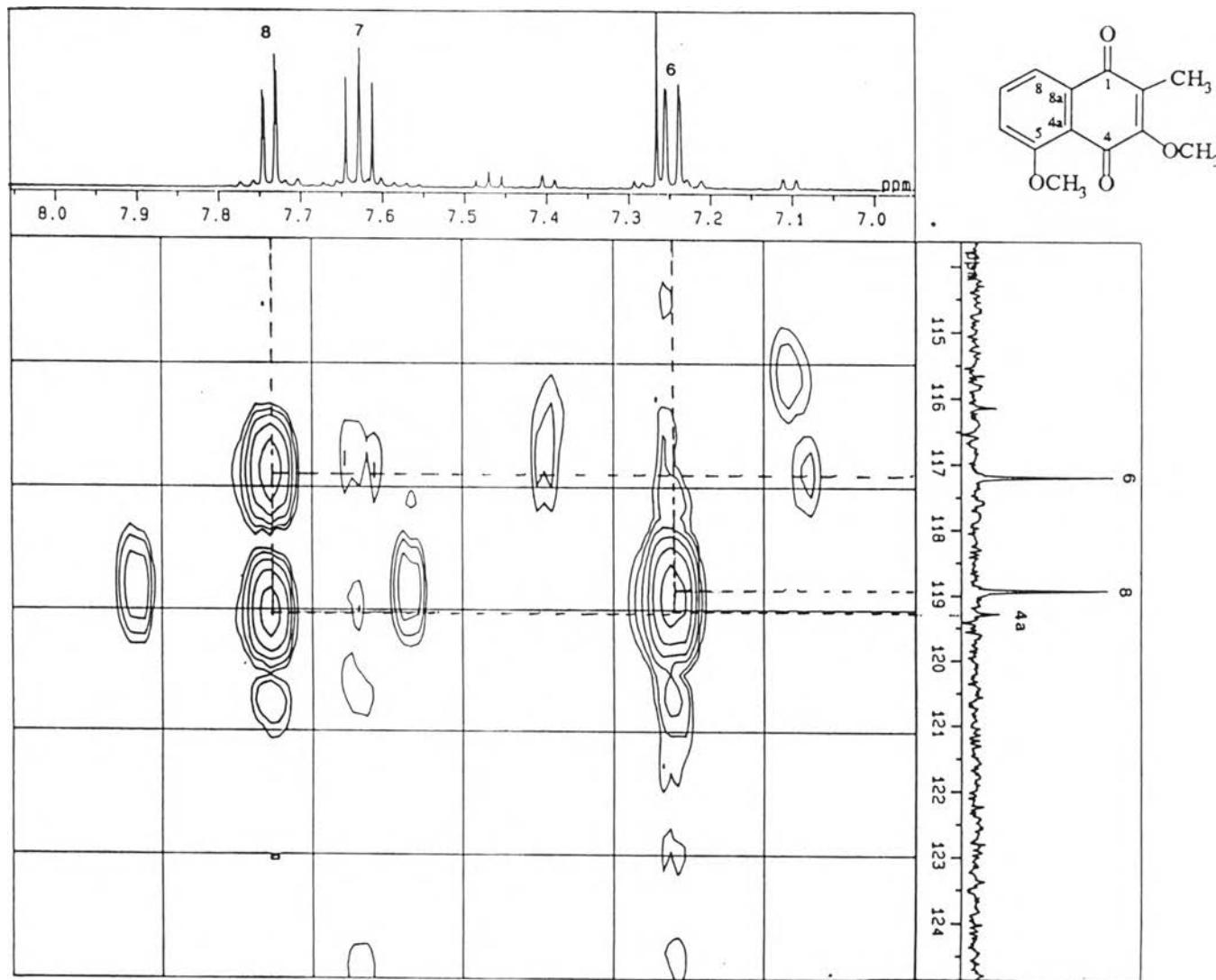


Figure 171 Expansion of the HMBC spectrum of compound 110 (in CDCl_3) :

δ_{H} 7.00-8.00 ; δ_{C} 115.00-124.00 ppm

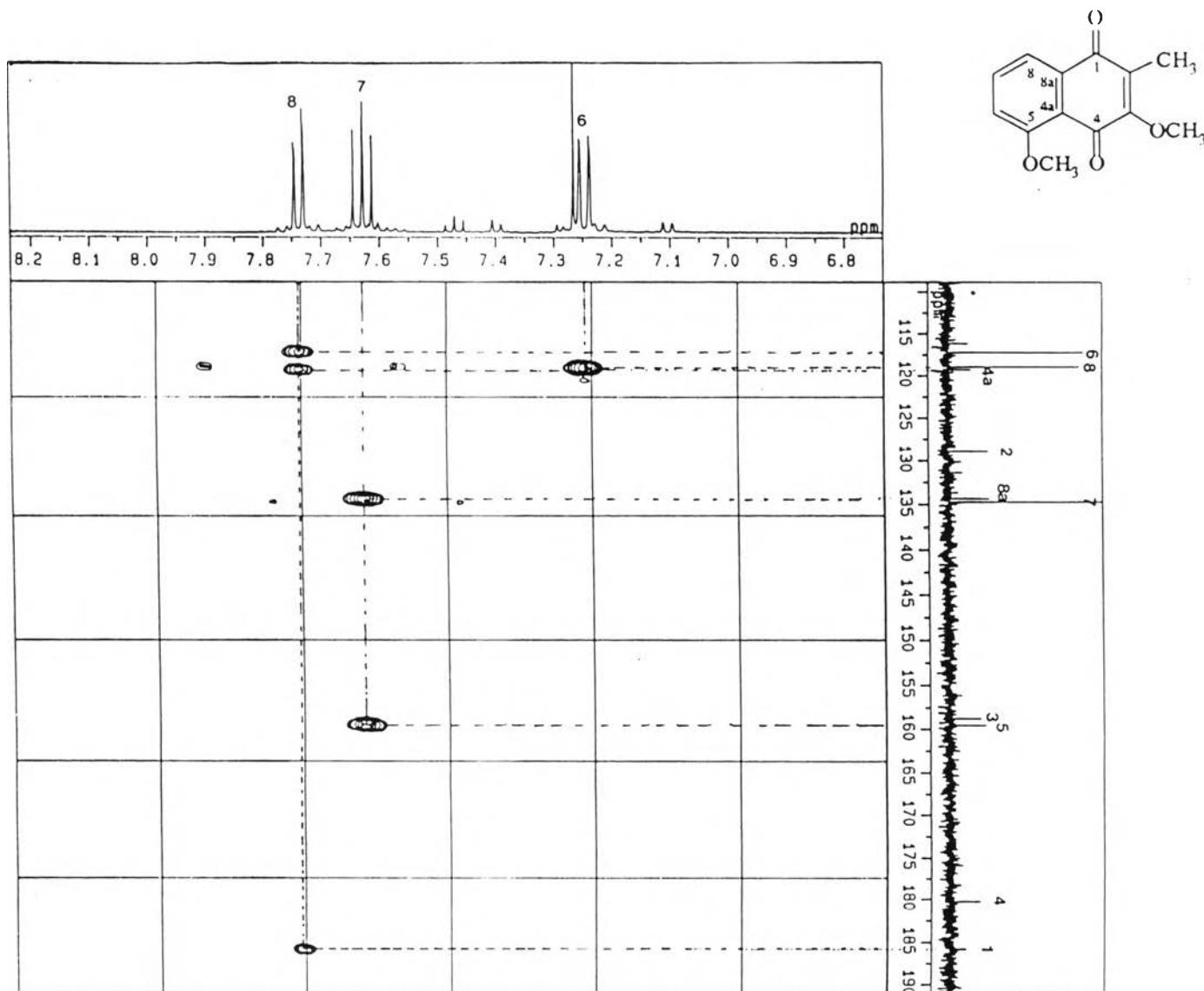


Figure 172 Expansion of the HMBC spectrum of compound 111 (in CDCl_3) :

δ_{H} 6.80-8.20 ; δ_{C} 115.00-190.00 ppm

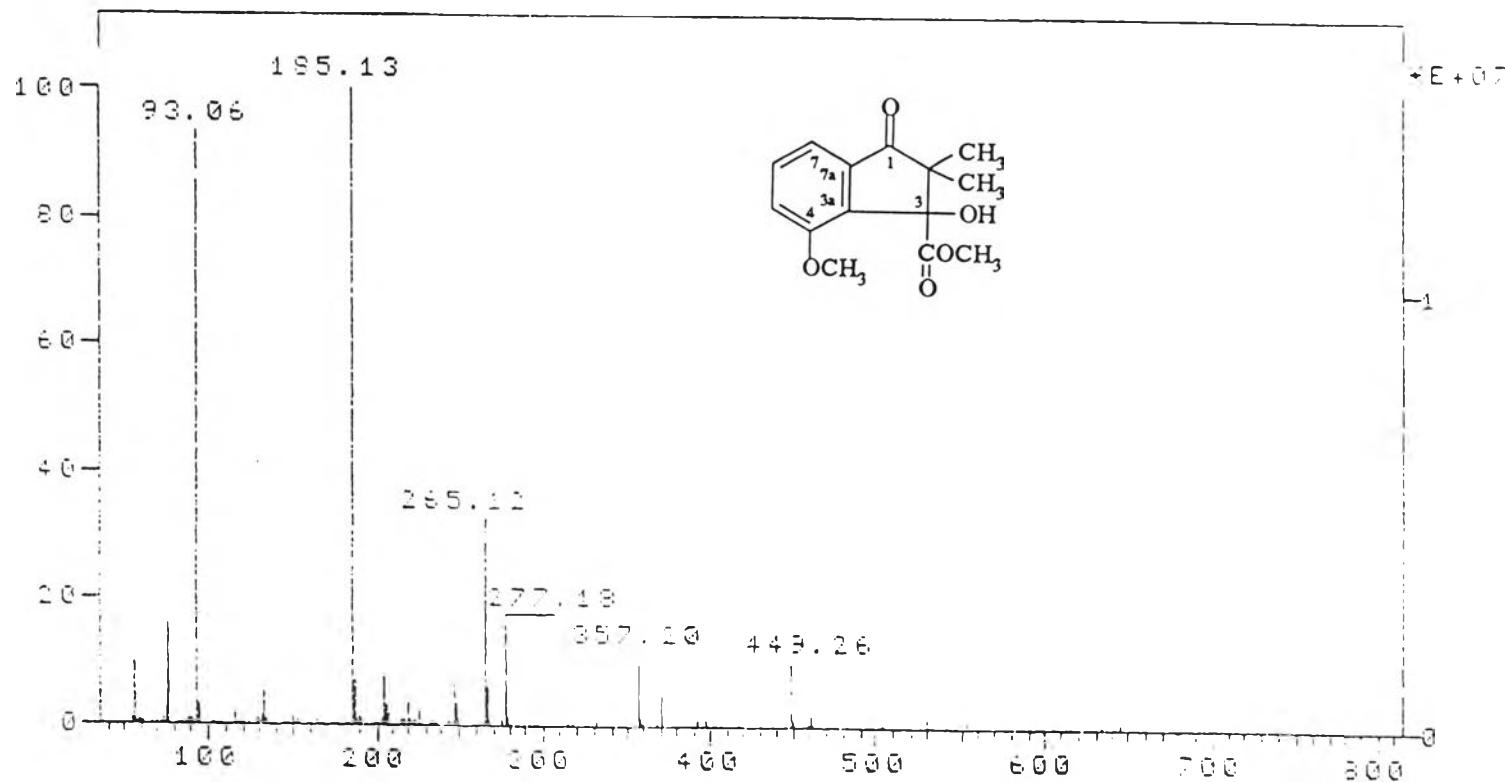


Figure 173 The HRFAB-MS spectrum of compound 111

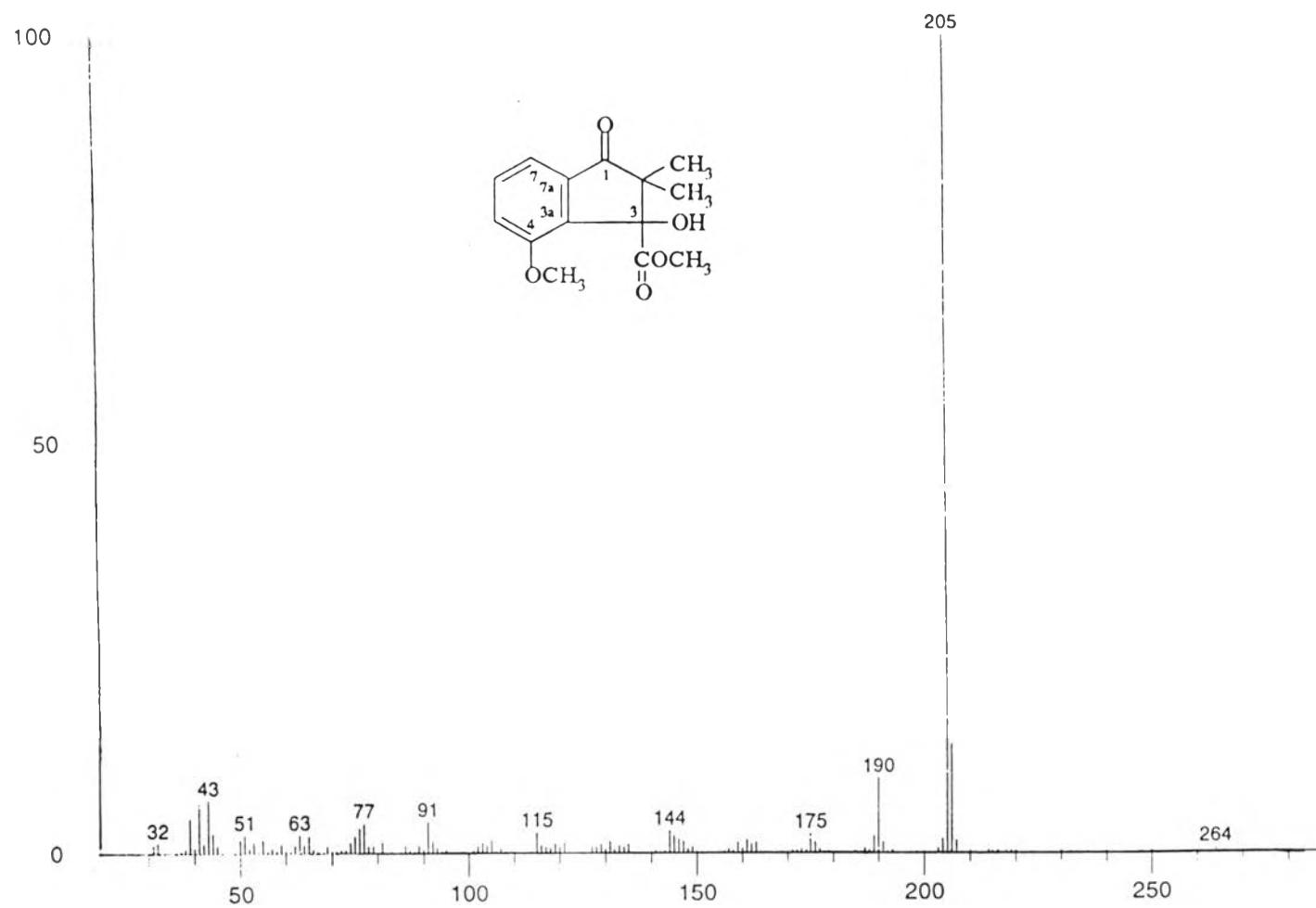


Figure 174 The EIMS spectrum of compound 111

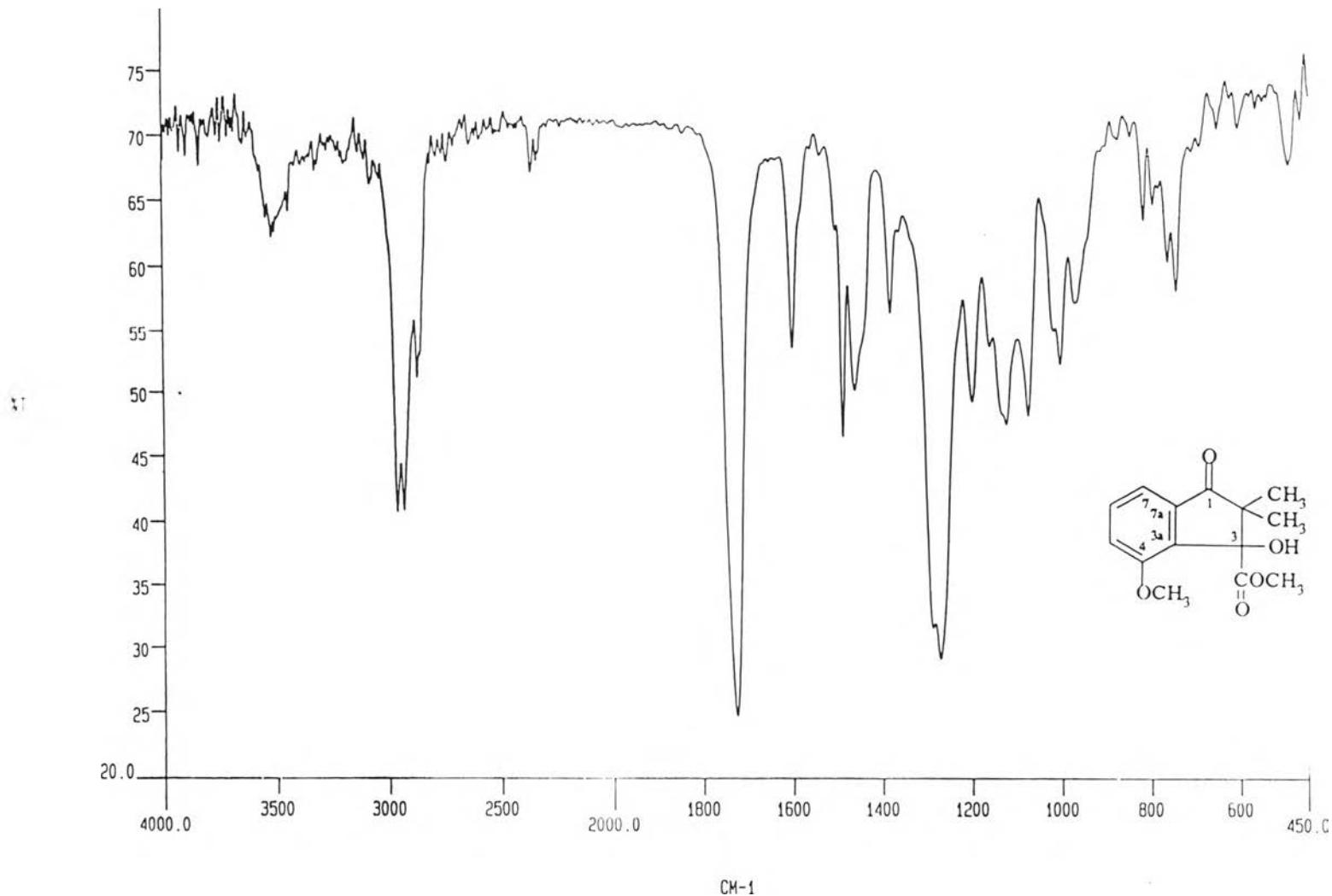


Figure 175 The IR spectrum of compound 111 (in CDCl₃)

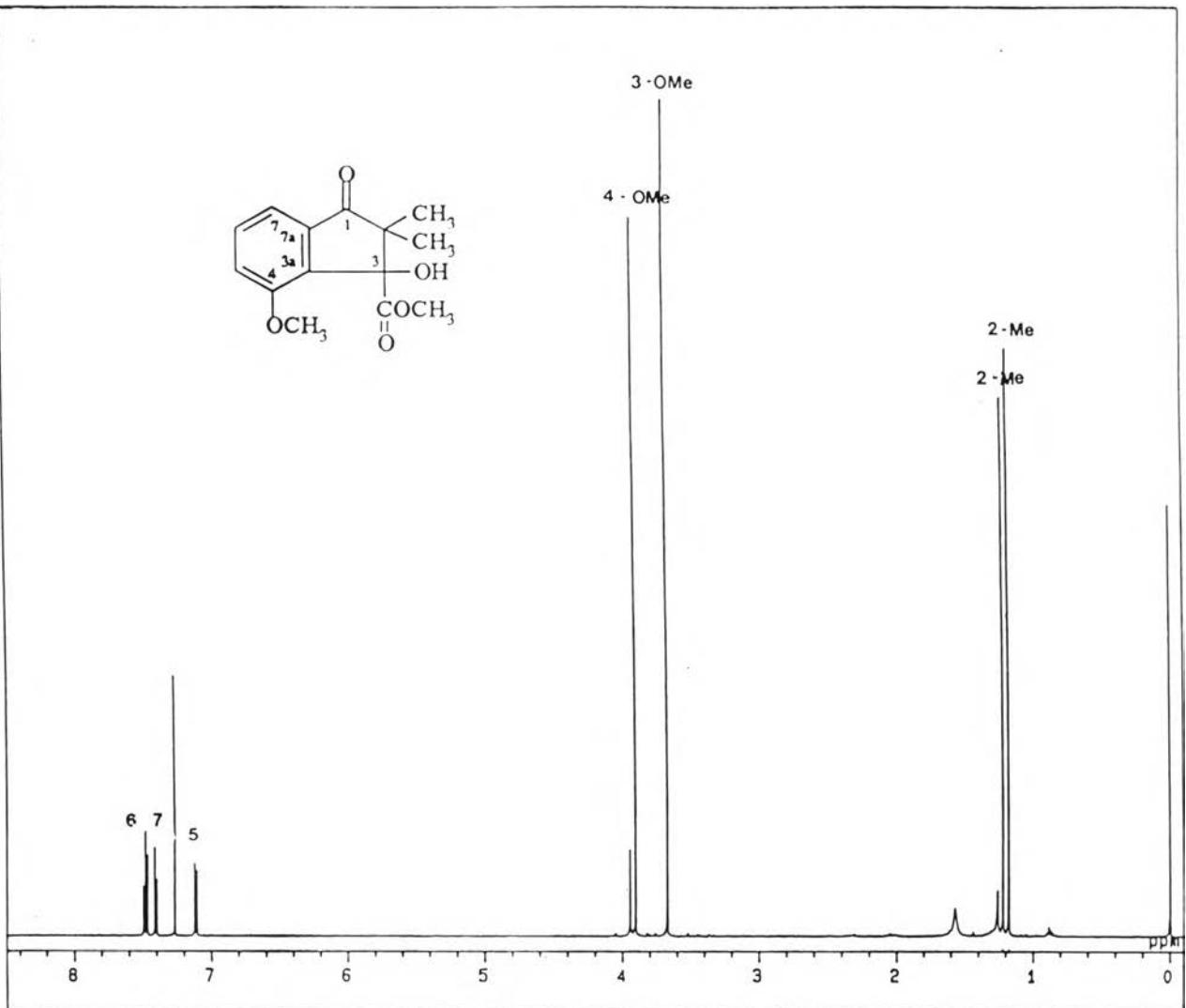


Figure 176 The ^1H NMR (500 MHz) spectrum of compound 111 (in CDCl_3)

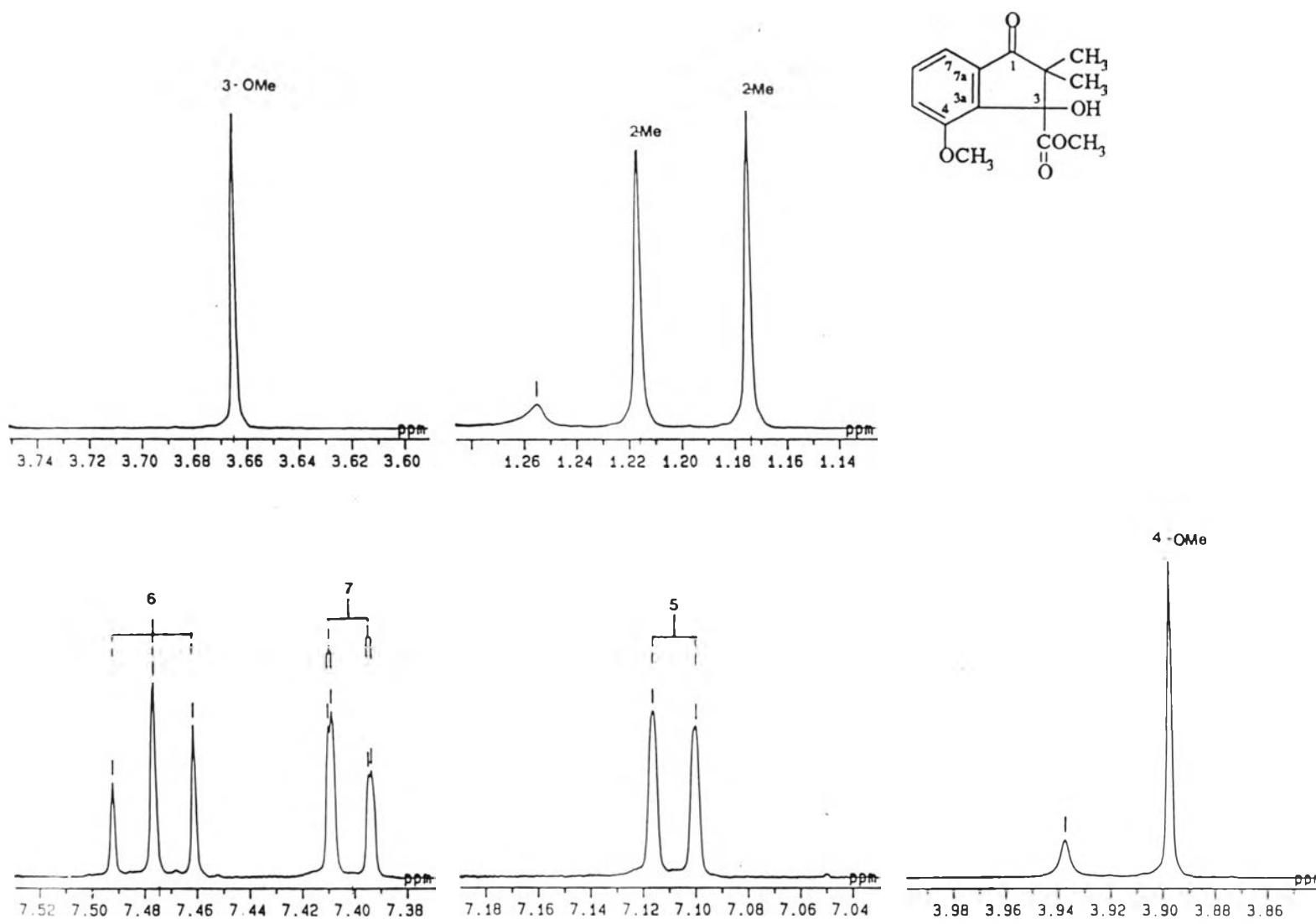


Figure 177 Expansion of the ^1H NMR (500 MHz) spectrum of compound 111
 (in CDCl_3) : δ_{H} 1.14-7.52 ppm

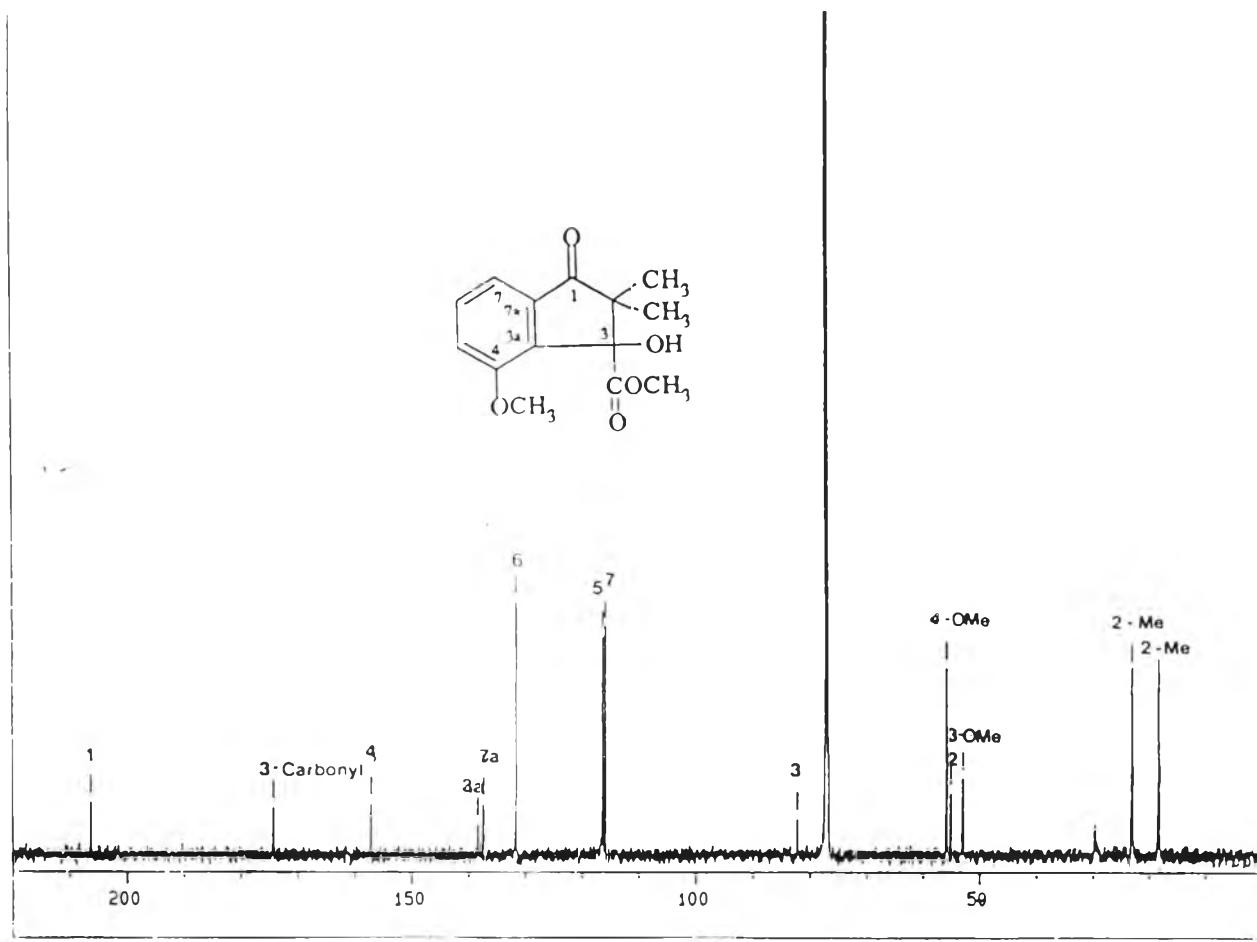


Figure 178 The ^{13}C NMR (125 MHz) spectrum of compound 111 (in CDCl_3)

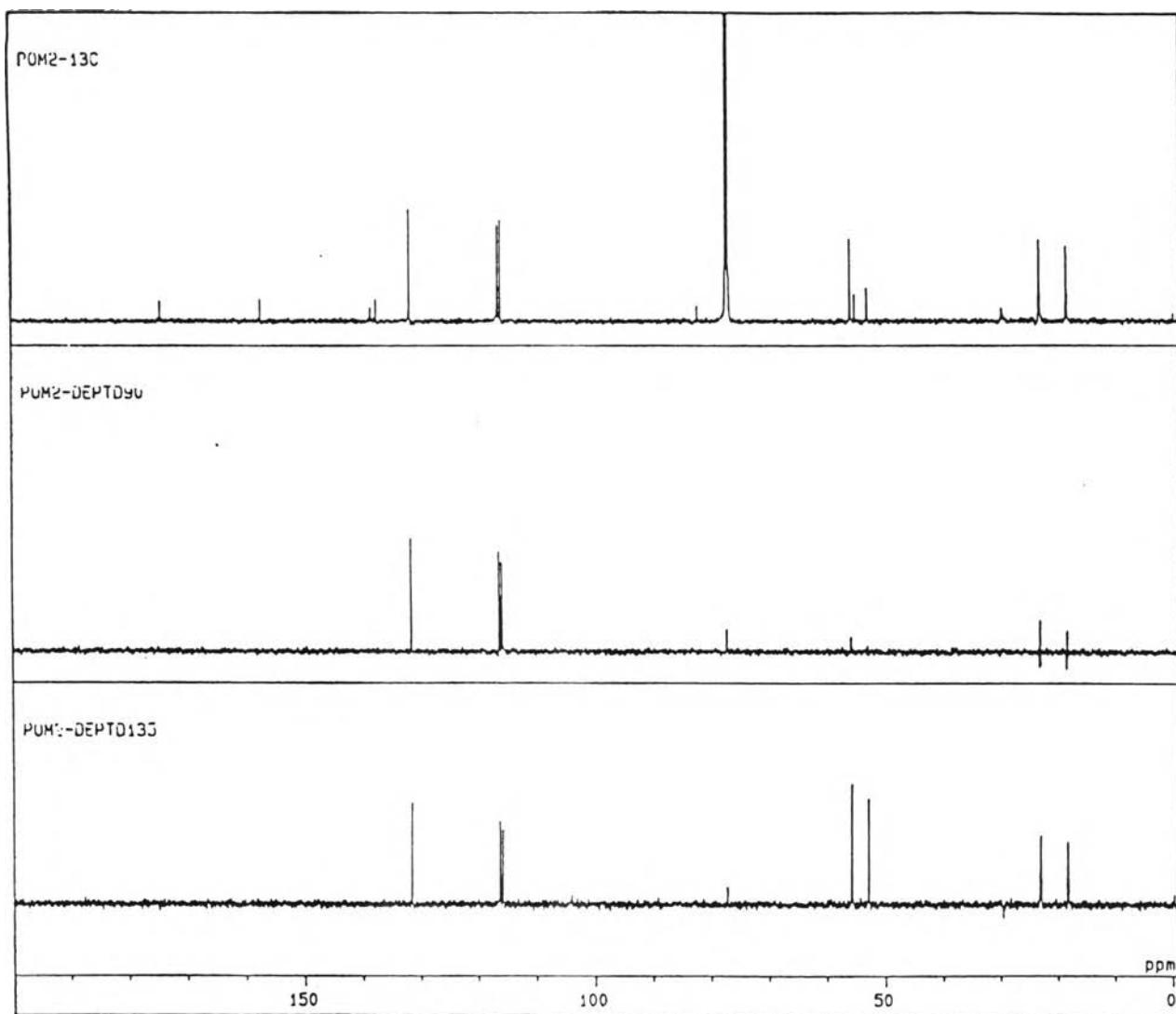
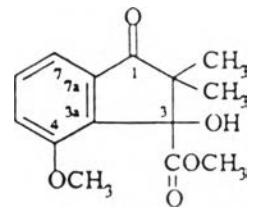


Figure 179 The DEPT (125 MHz) spectrum of compound 111 (in CDCl₃)

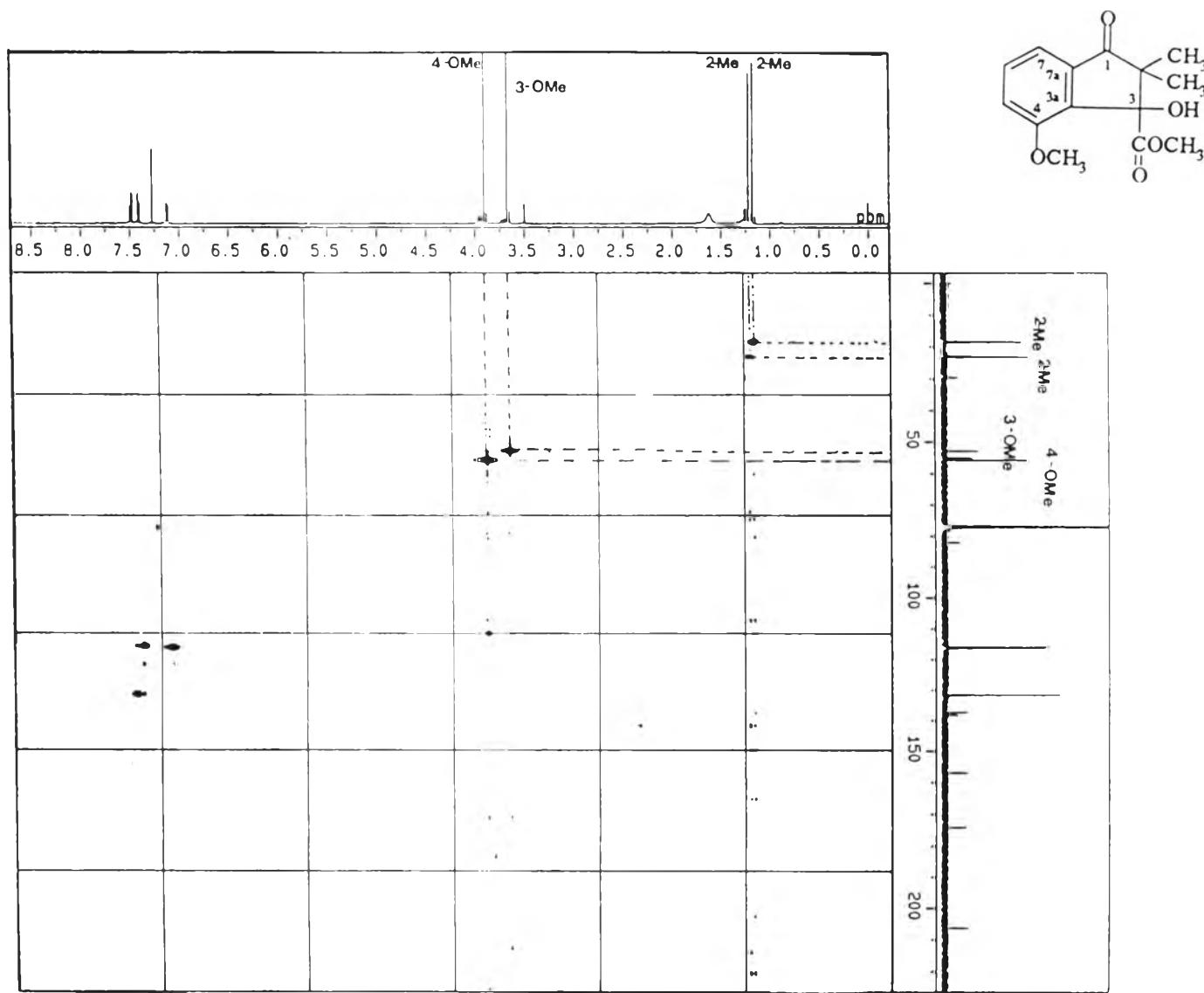


Figure 180 The HMQC spectrum of compound 111 (in CDCl_3)

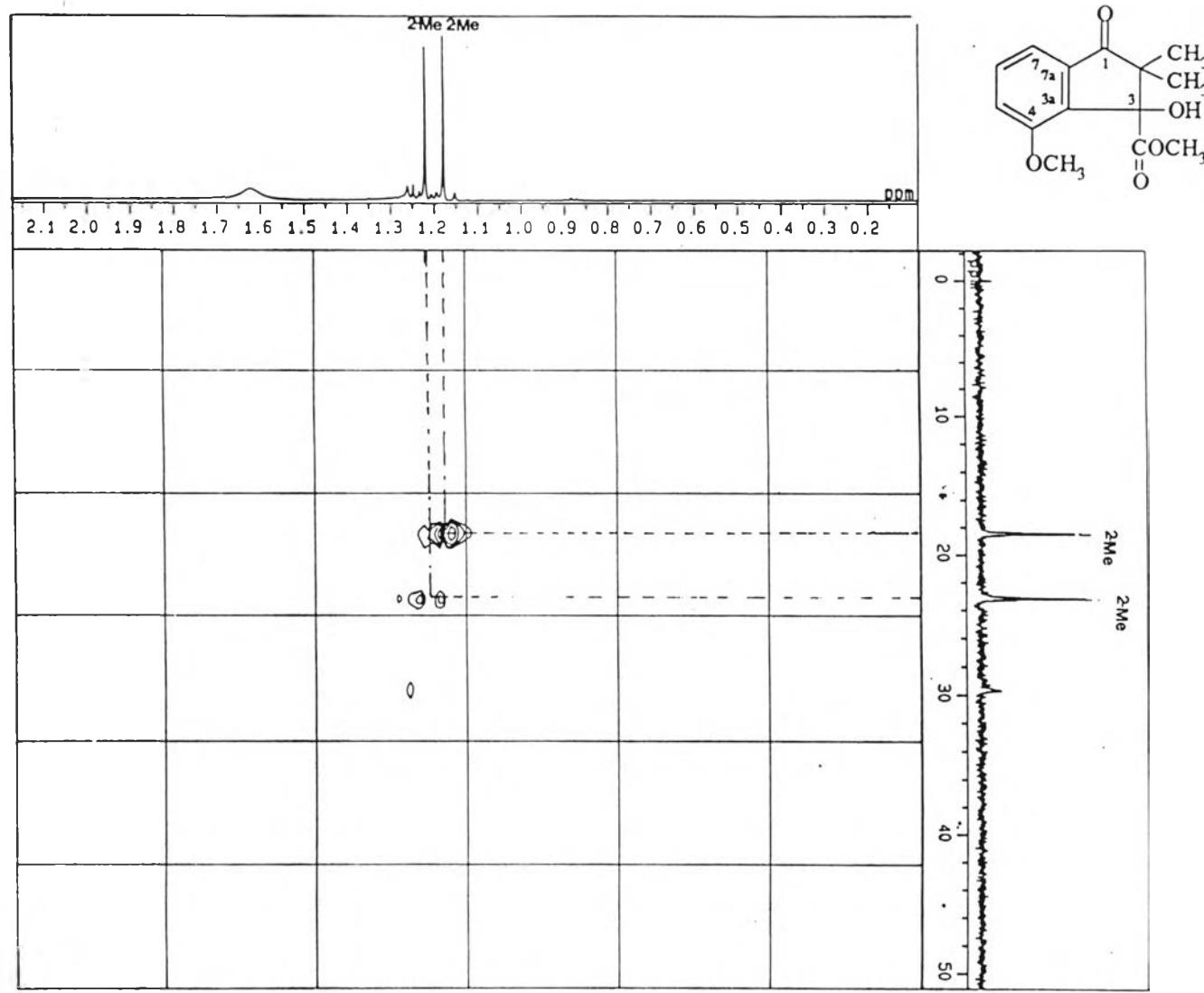


Figure 181 Expansion of the HMQC spectrum of compound 111 (in CDCl₃) :
 δ_{H} 0.20-2.10 ; δ_{C} 0.00-50.00 ppm

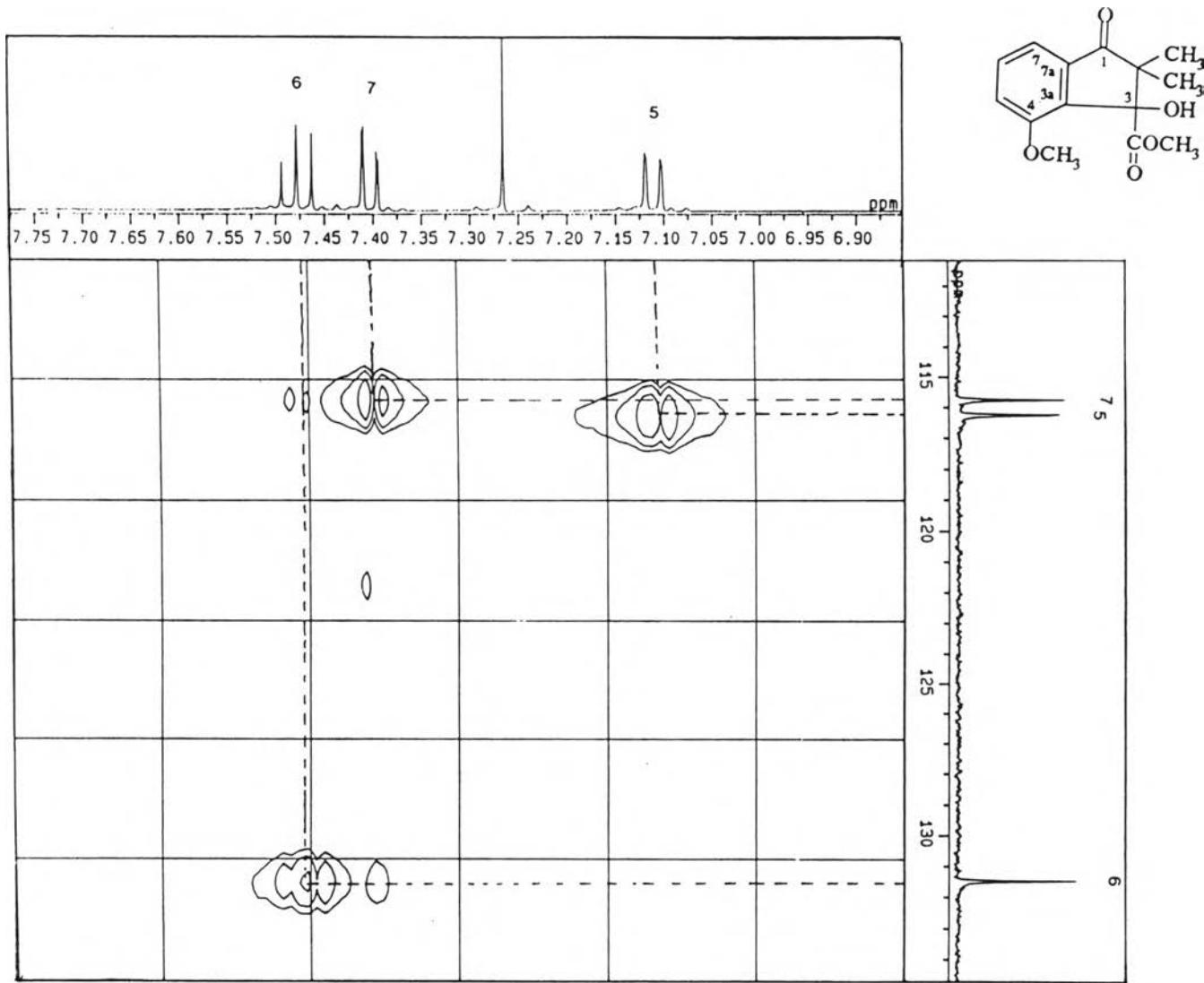


Figure 182 Expansion of the HMQC spectrum of compound 111 (in CDCl₃) :

δ_{H} 6.90-7.75 ; δ_{C} 112.00-134.00 ppm

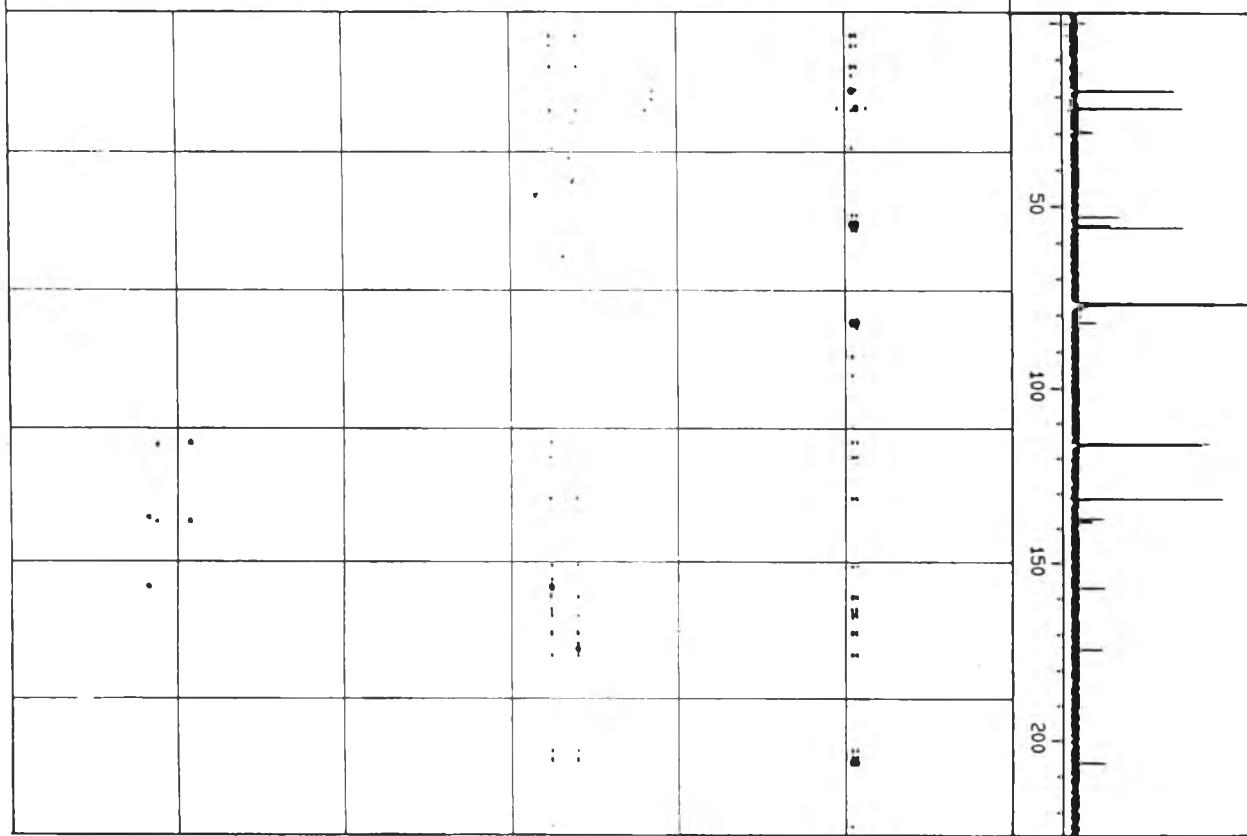
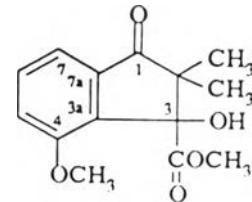
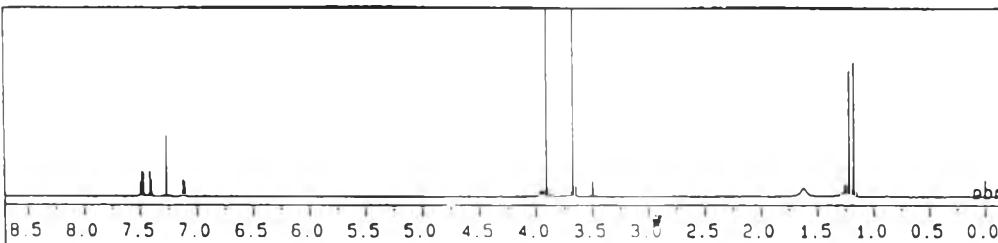


Figure 183 The HMBC spectrum of compound 111 (in $CDCl_3$)

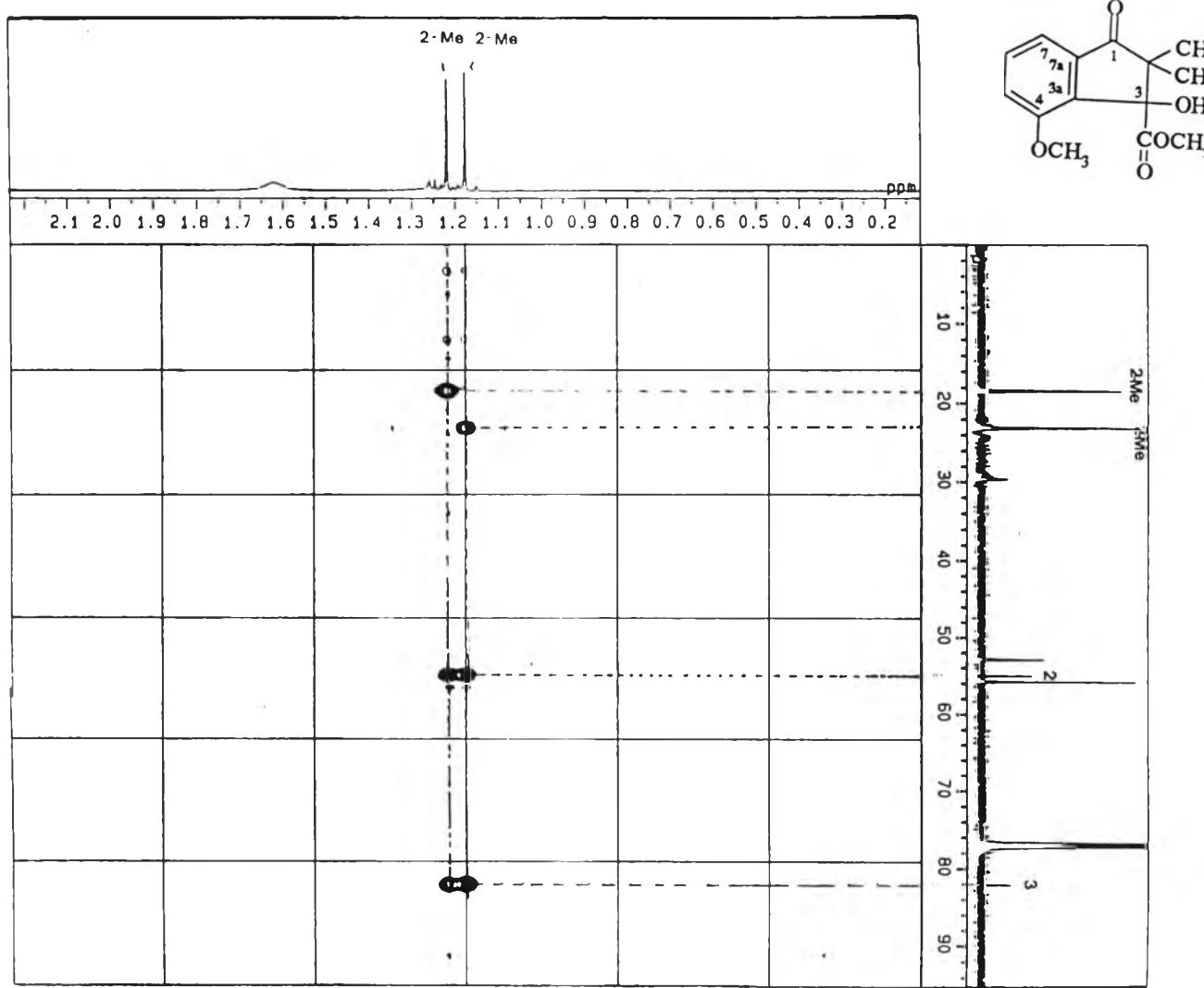


Figure 184 Expansion of the HMBC spectrum of compound 111 (in CDCl_3) :

δ_{H} 0.20-2.10 ; δ_{C} 10.00-90.00 ppm

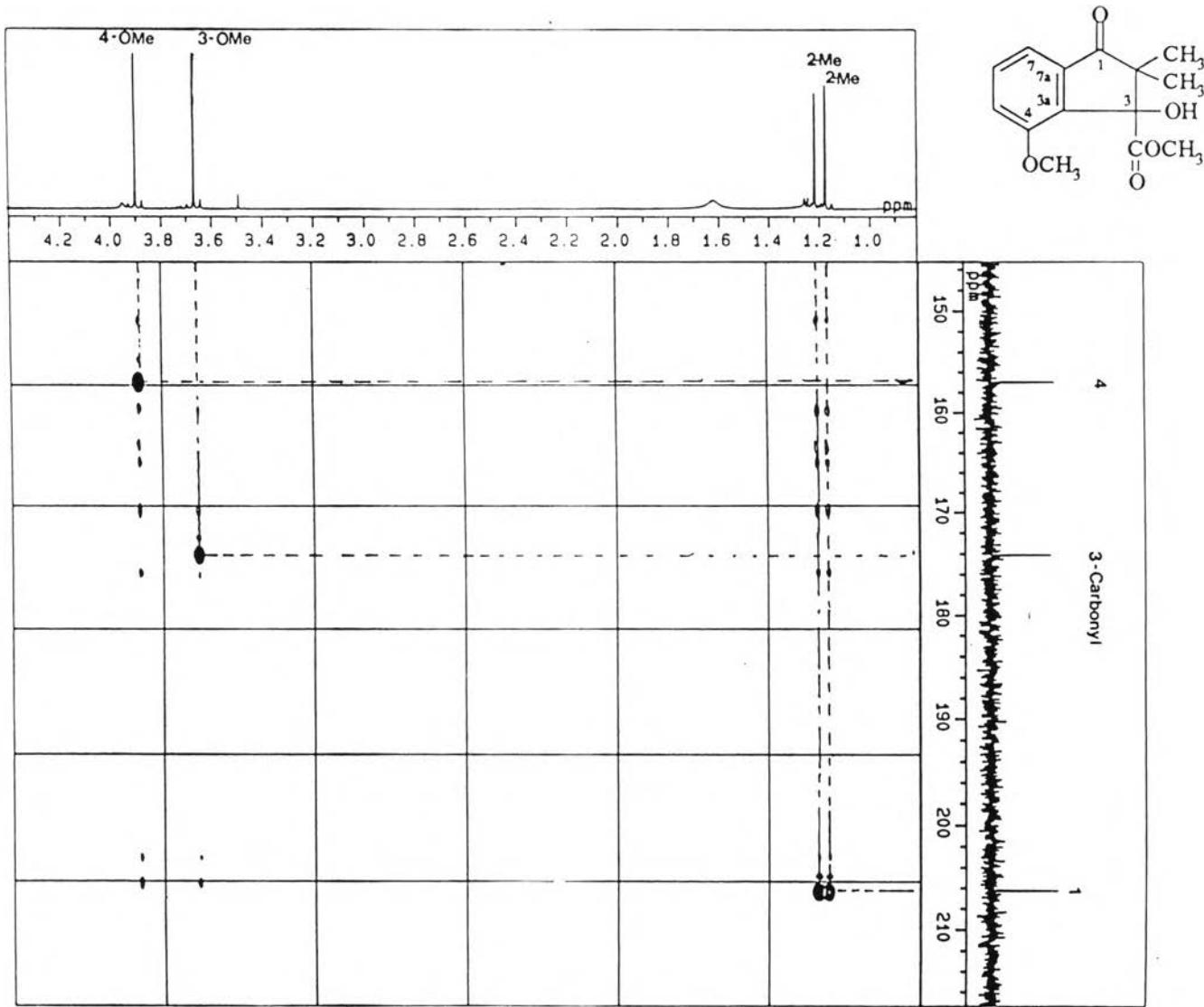


Figure 185 Expansion of the HMBC spectrum of compound 111 (in CDCl_3) :

δ_{H} 1.00-4.20 ; δ_{C} 150.00-210.00 ppm

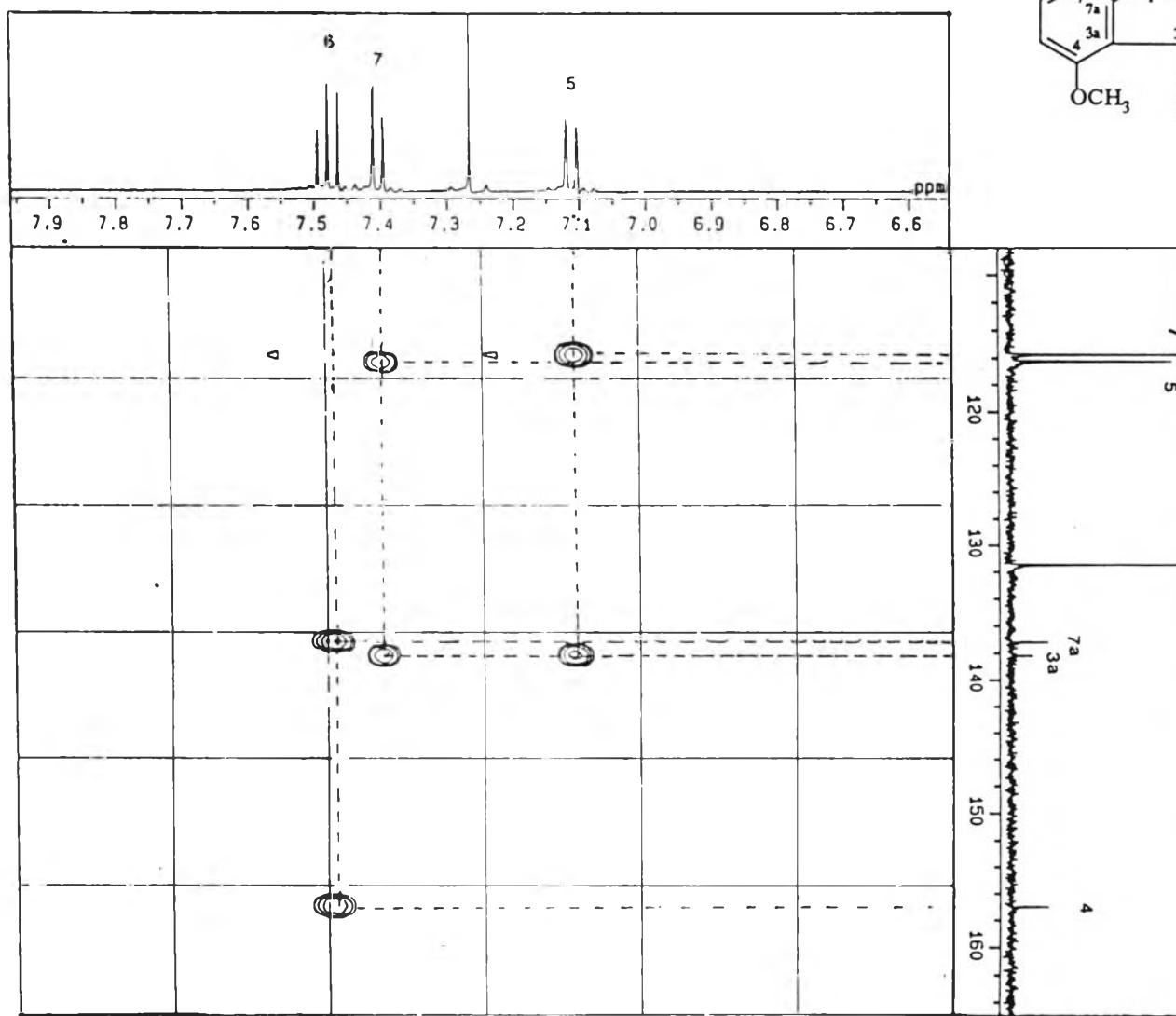
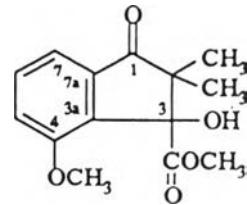


Figure 186 Expansion of the HMBC spectrum of compound 111 (in CDCl_3) :
 δ_{H} 6.60-7.90 ; δ_{C} 110.00-160.00 ppm

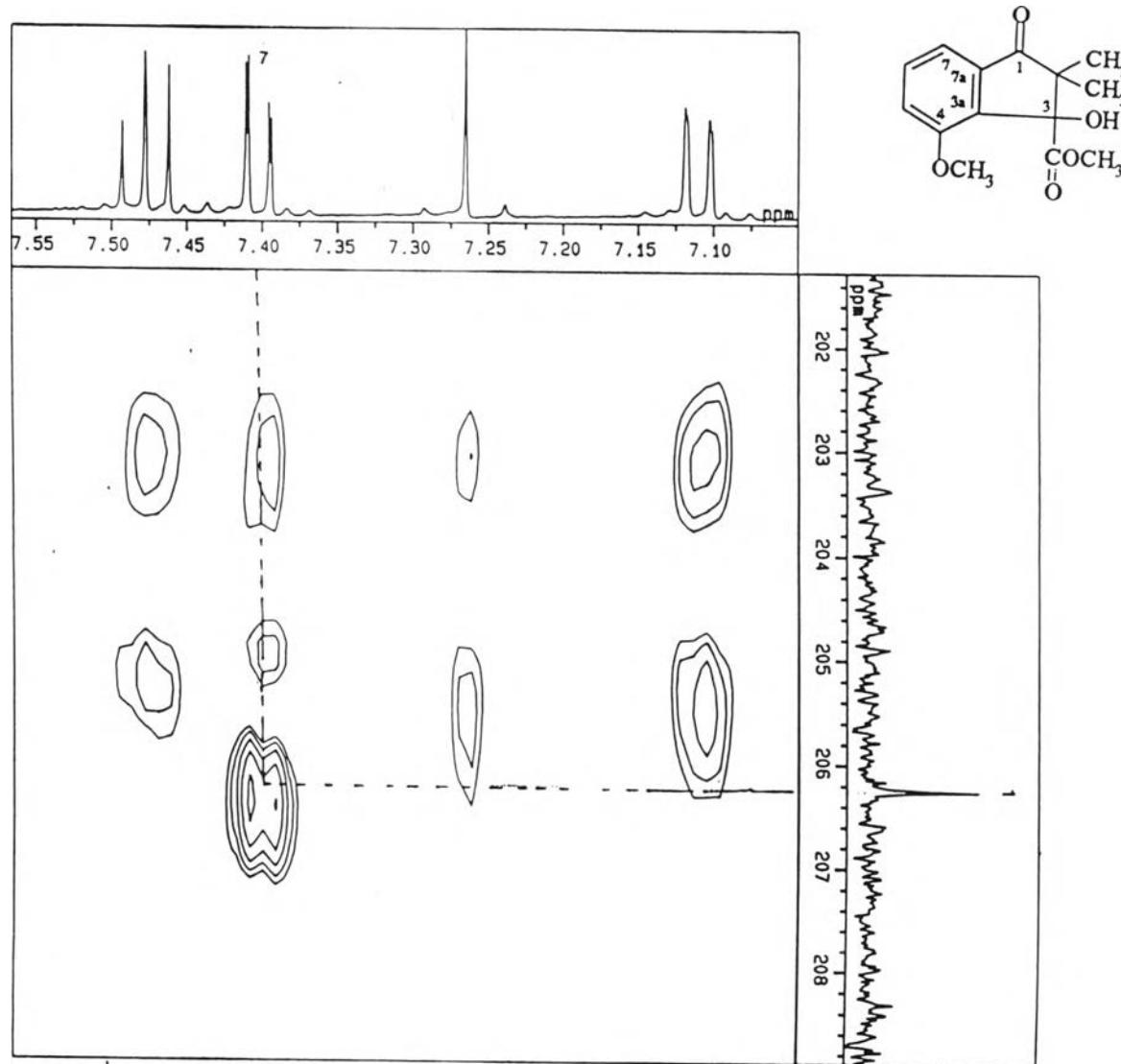


Figure 187 Expansion of the HMBC spectrum of compound 111 (in CDCl_3) :

δ_{H} 7.10-7.55 ; δ_{C} 202.00-208.00 ppm

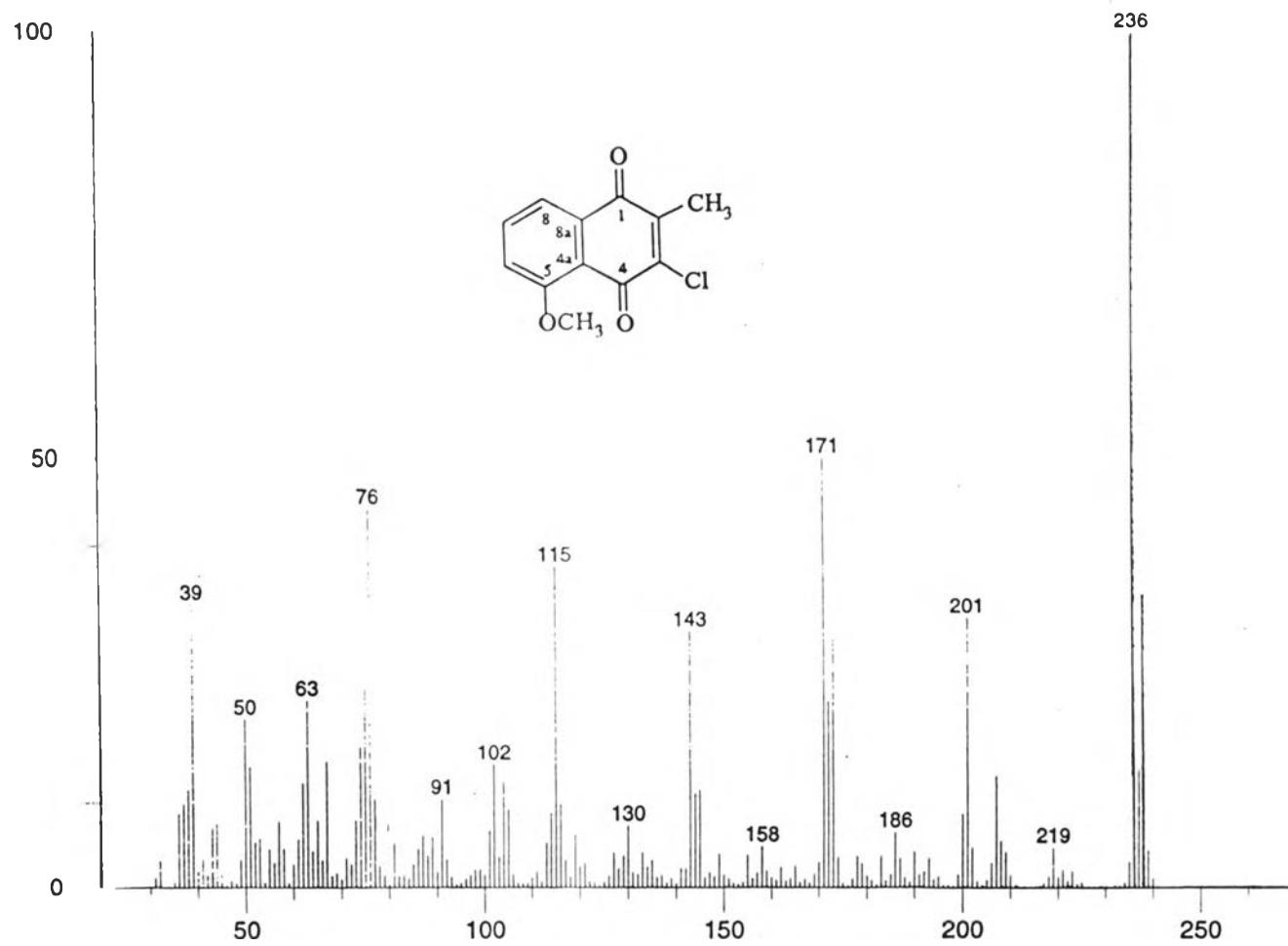


Figure 189 The EI mass spectrum of compound 112

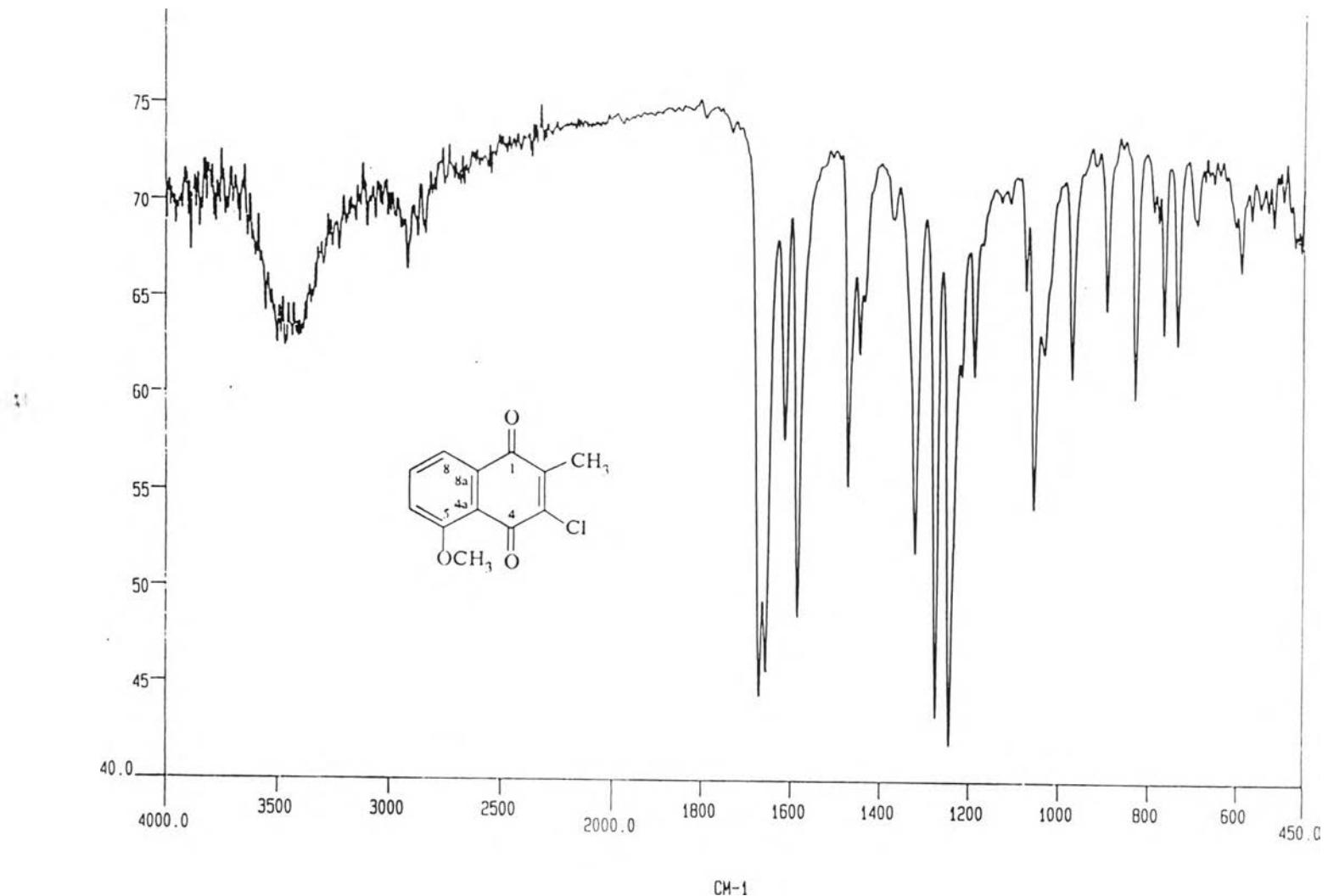


Figure 190 The IR spectrum of compound 112 (in KBr disc)

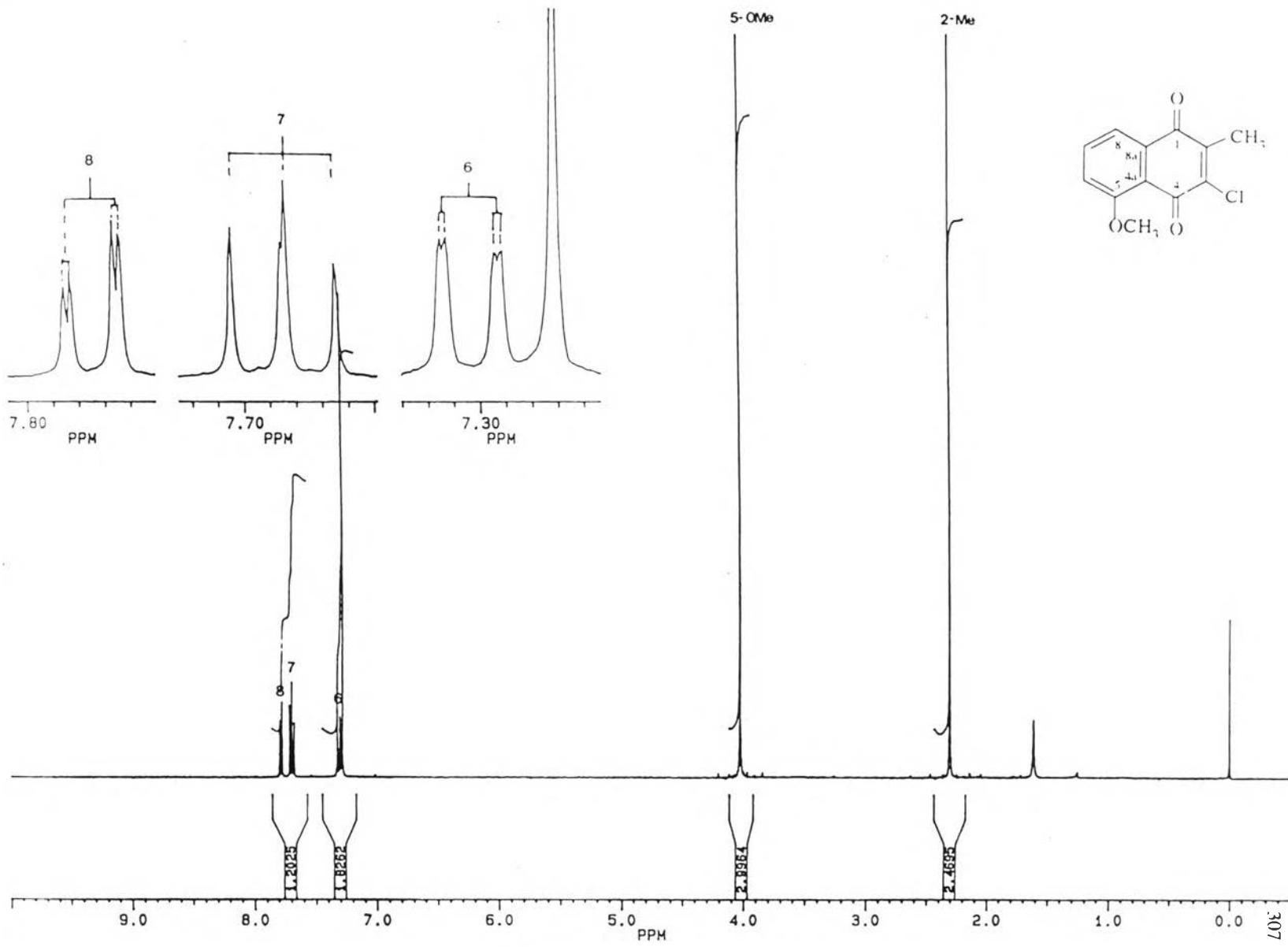


Figure 191 The ^1H NMR (400 MHz) of compound 112 (in CDCl_3)

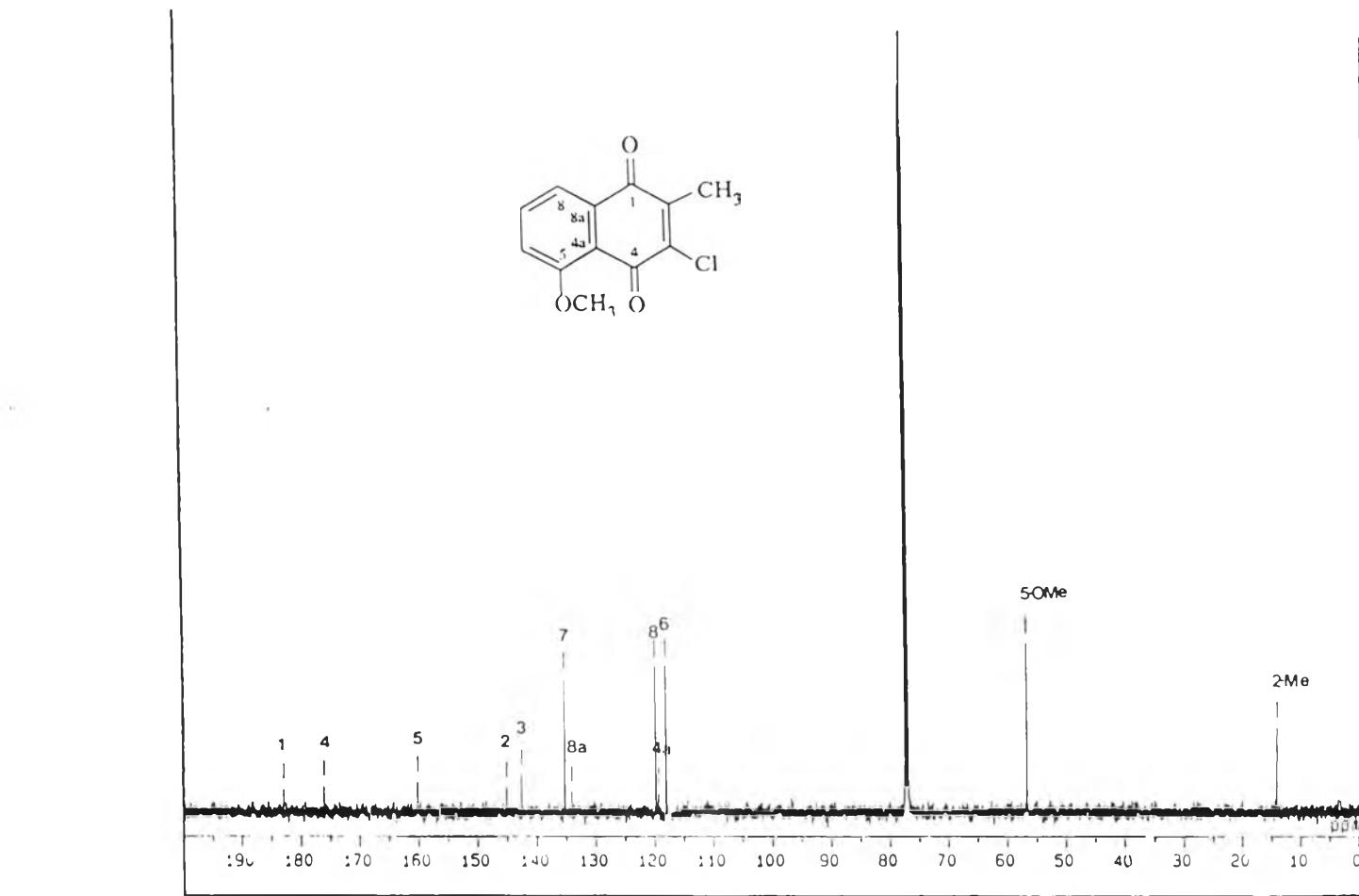
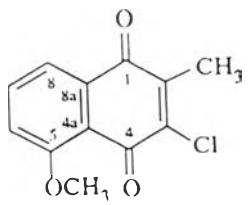
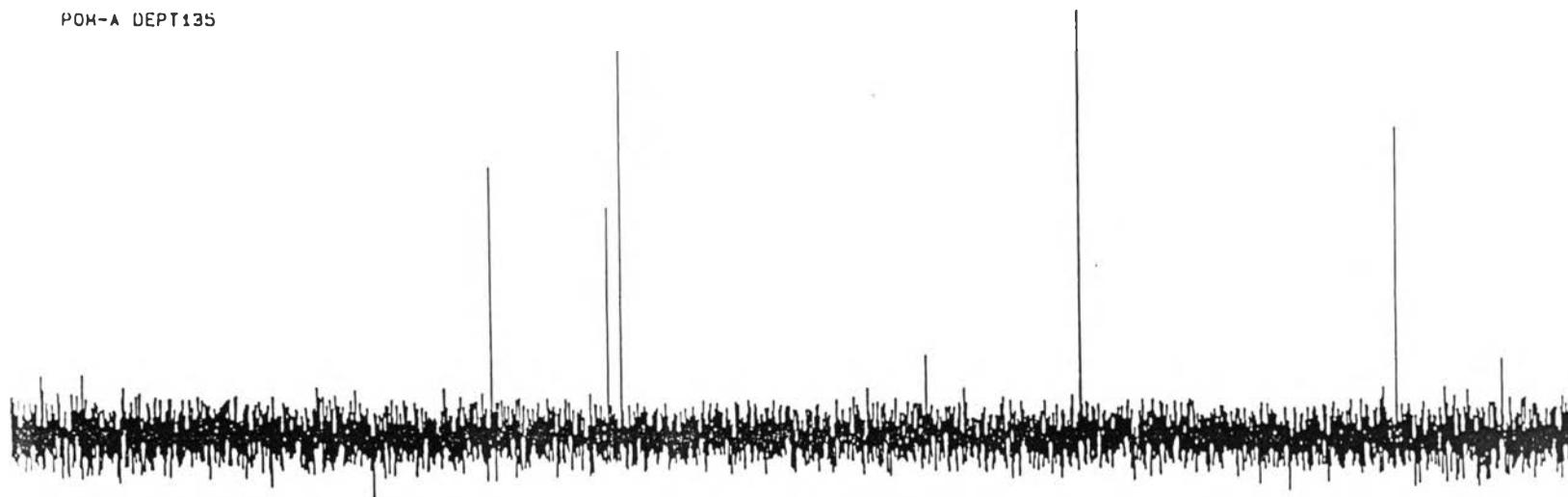


Figure 192 The ^{13}C NMR (125 MHz) of compound **112** (in CDCl_3)

POH-A DEPT135



DEPT90

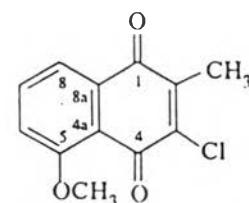
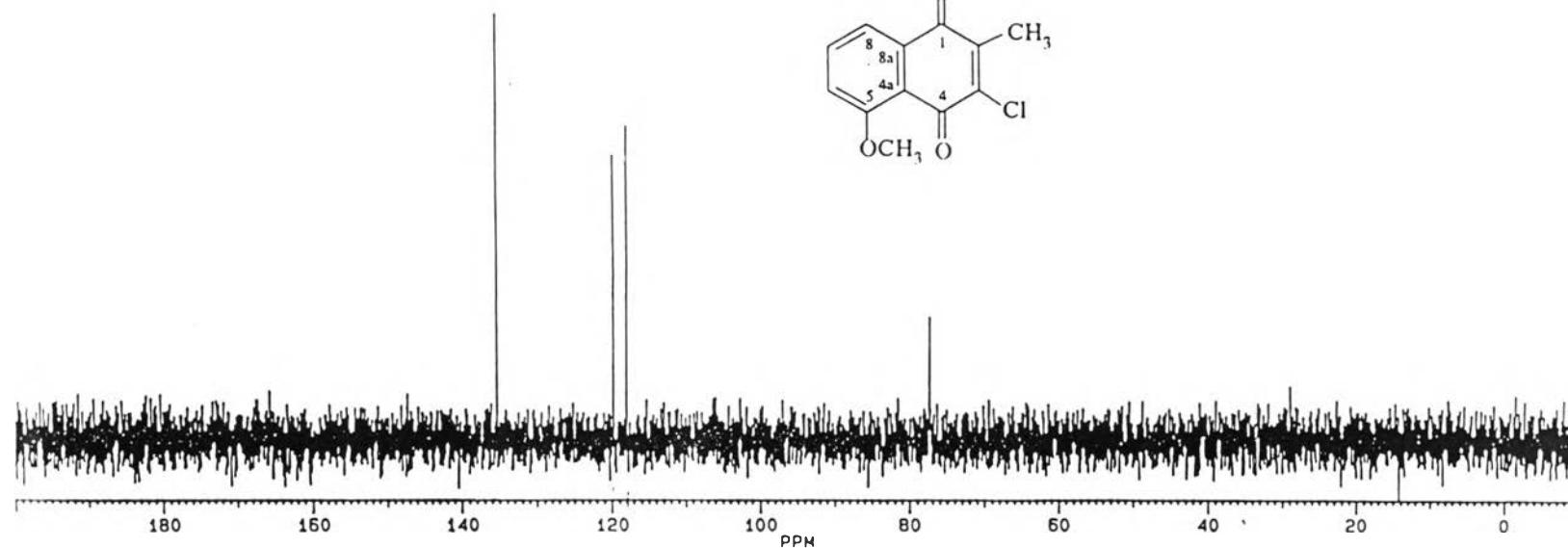


Figure 193 The DEPT (100 MHz) of compound 112 (in CDCl₃)

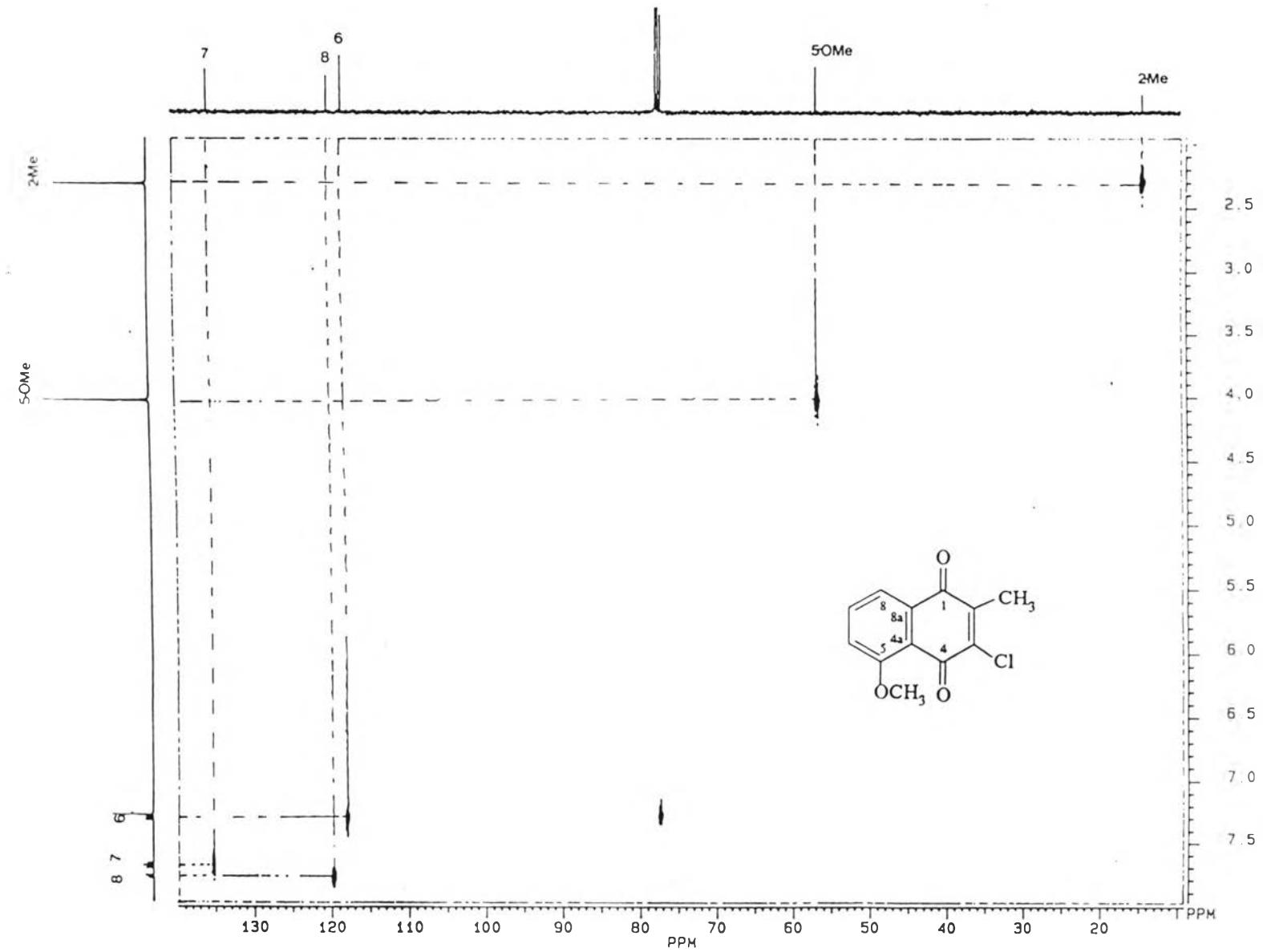


Figure 194 The HETCOR spectrum of compound 112 (in CDCl_3)

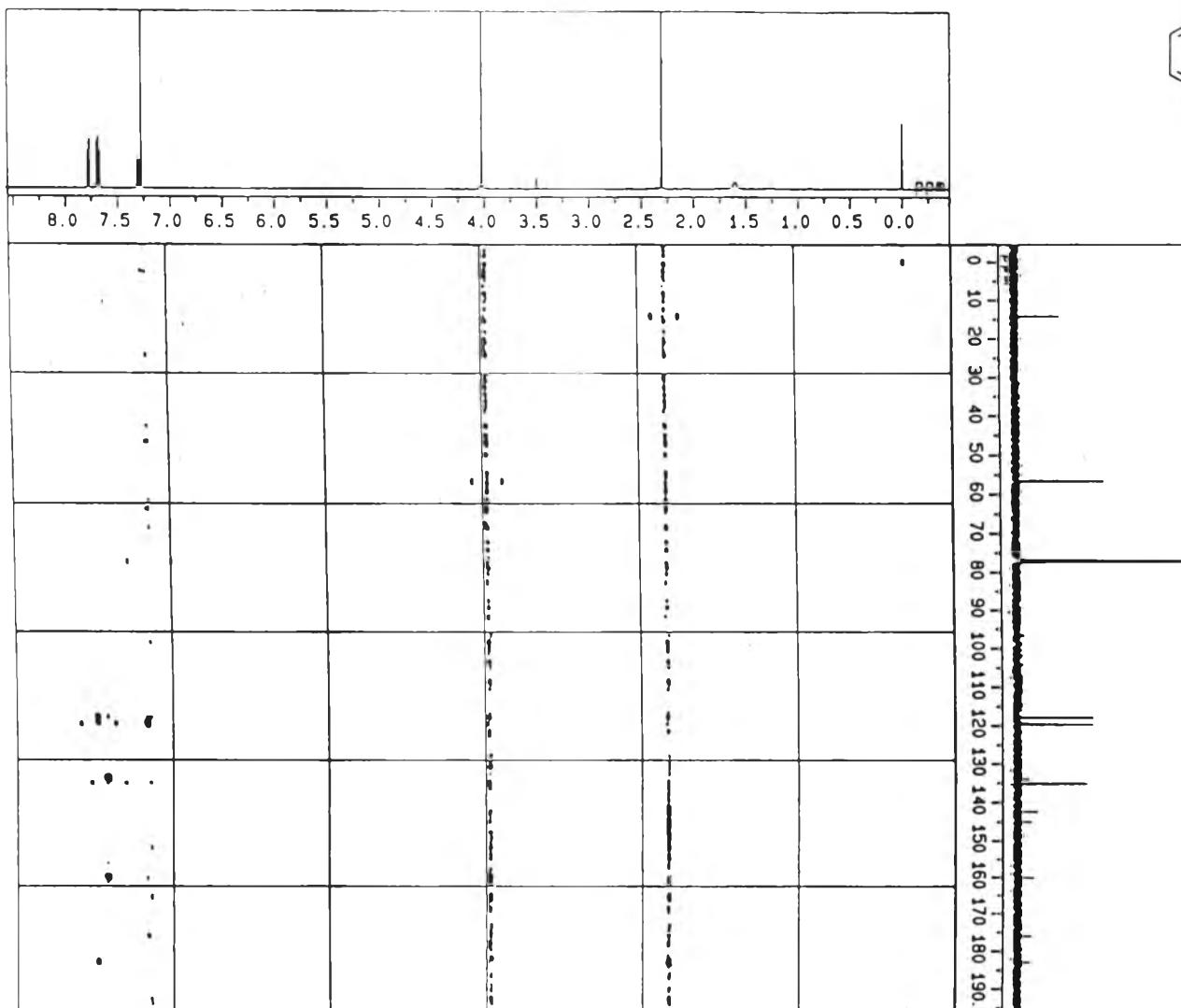
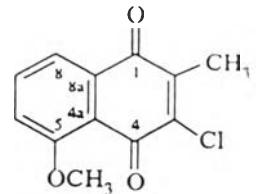


Figure 195 The HMBC spectrum of compound 112 (in CDCl₃)

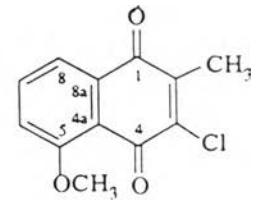
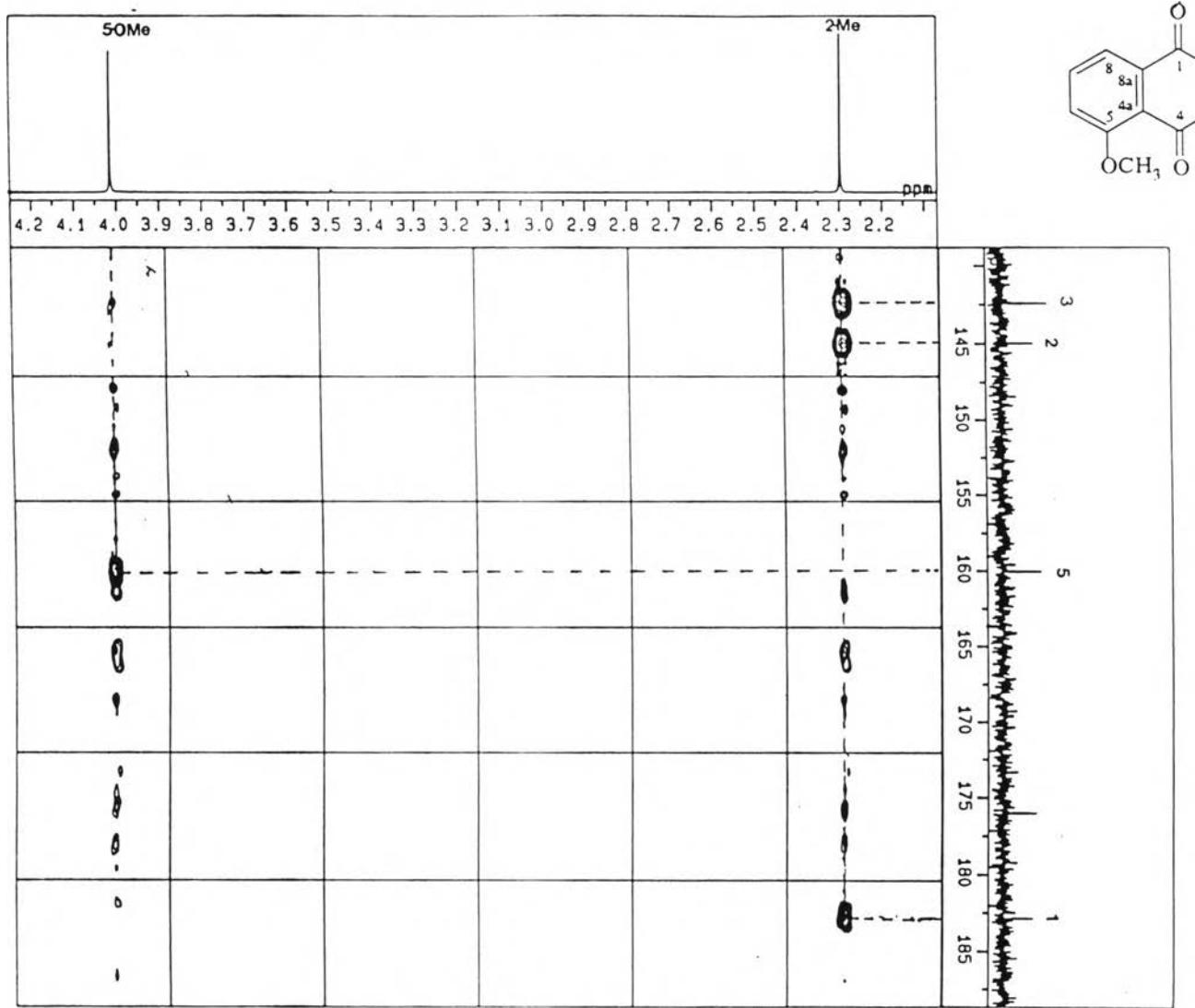


Figure 196 Expansion of the HMBC spectrum of compound 112 (in CDCl_3) :

δ_{H} 2.20-4.20 ; δ_{C} 140.00-185.00 ppm

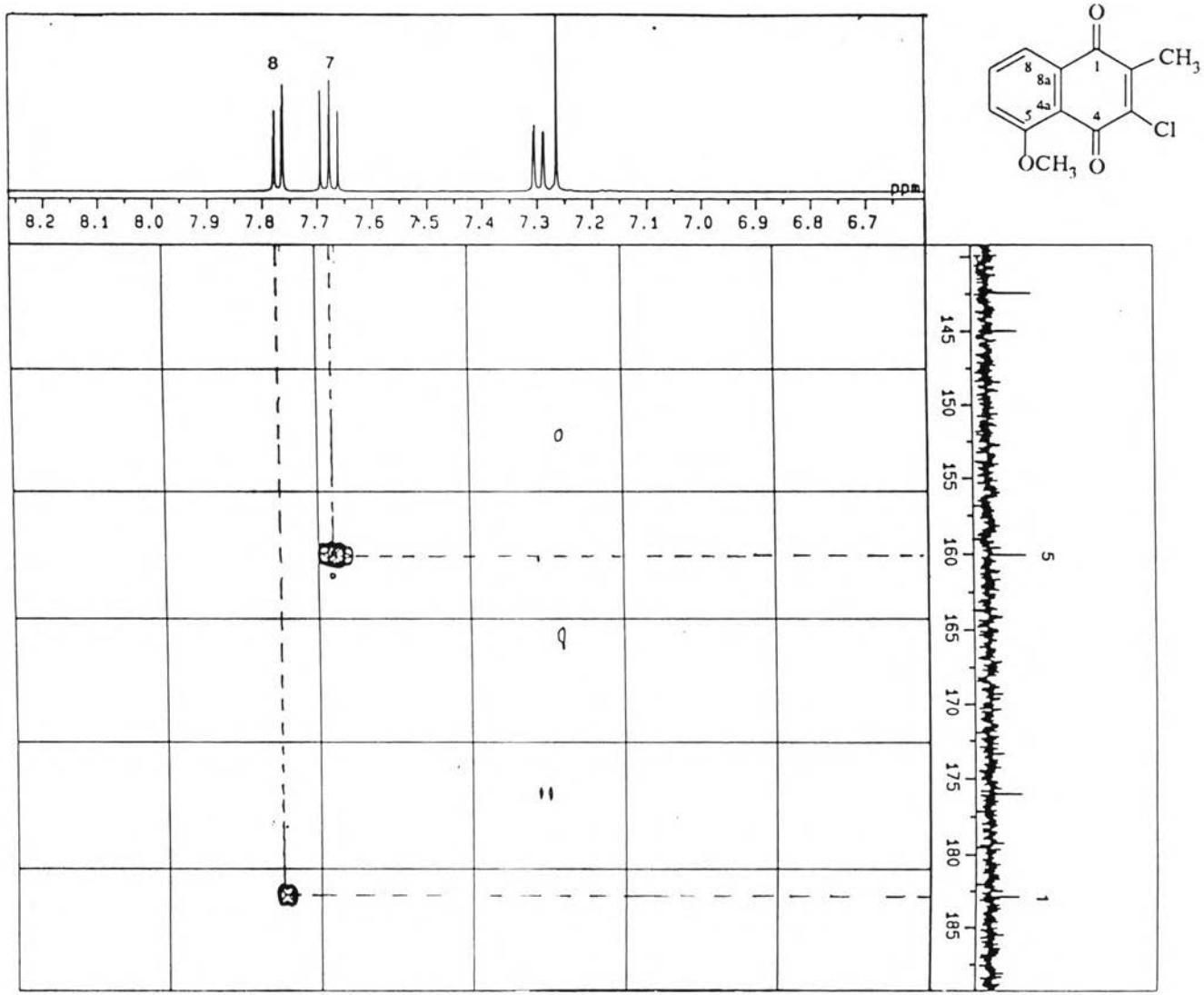


Figure 197 Expansion of the HMBC spectrum of compound 112 (in CDCl₃) :

δ_{H} 6.70-8.20 ; δ_{C} 140.00-185.00 ppm

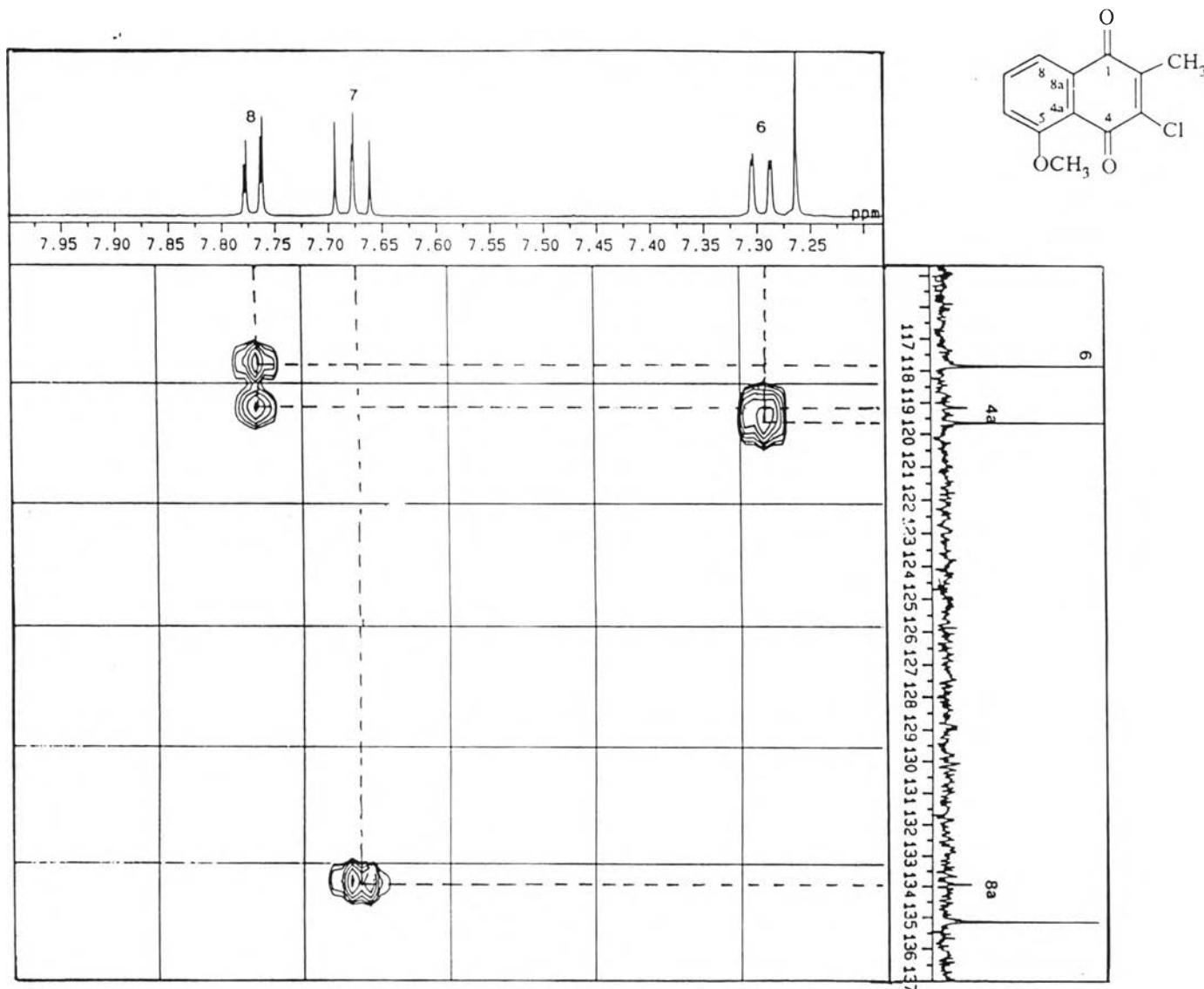


Figure 198 Expansion of the HMBC spectrum of compound 112 (in CDCl_3) :

δ_{H} 7.25-7.95 ; δ_{C} 117.00-135.00 ppm

Vita

Miss Rawiwun Kaewamatawong was born on May, 20, 1970 in Ubon Ratchathani, Thailand. She received her Bachelor of Science in Pharmacy in 1993 from the Faculty of Pharmacy, Rangsit University. Throughout her M.S. study, she received a scholarship from the University Development Commission (UDC) with an obligation to serve, after graduation, as a faculty member at the Faculty of Pharmacy, Ubon Ratchathani University.

