

## REFERENCES

1. Jones, J. *Amino Acid and Peptide Synthesis*. Oxford; Oxford University Press: 1992.
2. Williams, R. M.; Hendrix, J. A. Asymmetric Synthesis of Arylglycines. *Chem. Rev.* **1992**, *92*, 889.
3. Červinka, O. *Enantioselective Reactions in Organic Chemistry*, 1<sup>st</sup> ed., Prague: Publishing House of the Academy of Sciences of the Czech Republic, 1995.
4. Baker, D. R.; Fenyes, J. G.; Basarab, G. S. and Hunt, D. A. *ACS Symposium Series: Synthesis and Chemistry of Agrochemicals V*, Washington, D. C.: American Chemical Society, 1998, 95.
5. For reviews, see: (a) wagner, I.; Musso, H. New Naturally Occurring Amino Acids. *Angew. Chem., Int. Ed. Engl.* **1983**, *22*, 816. (b) Greenstein, J. P.; Winitz, M.; Robert E. Krieger *Chemistry of the Amino Acids*; FL 1984; Vols. 1-3. (c) Barrett, G. C.; Ed. *Chemistry and Biochemistry of the Amino Acids*; Chapman and Hall : London, 1985. (d) O'Donnell, M. J., Ed.  $\alpha$  - Amino Acid Synthesis. *Tetrahedron* **1988**, *44*, 5253. (e) Williams, R. M. *Synthesis of optically Active  $\alpha$ -Amino Acids*. Oxford; Pergamon Press: 1989; Vol. 7.
6. For reviews on vancomycin group antibiotics, see : Williams, D. H. Structural Studies on Some Antibiotics of the Vancomycin Group, and on the Antibiotic-Receptor Complexes, by  $^1\text{H}$  NMR. *Acc. Chem. Res.* **1984**, *17*, 364.
7. (a) Harris, C. M.; Kibby, J. J.; Fehlner, J. R.; Raabe, H. B.; Barber, T. A.; Harris, T. M. Amino Acid Constituents of Ristocetin A. *J. Am. Chem. Soc.* **1979**, *101*, 437. (b) Harris, C. M.; Kibby, J. J.; Harris, T. M. The Biphenyl constituent of Ristocetin A. *Tetrahedron Lett.* **1978**, 705. (c) Harris, C. M.; Harris, T. M. Structure of Ristocetin A: Configurational Studies of the Peptide. *J. Am. Chem. Soc.* **1982**, *104*, 363. (d) Harris, C. M.; Fesik, S. W.; Thomas, A. M.; Kannan, R.; Harris, T. M. Iodination of Vancomycin, Ristocetin A, and Ristocetin Pseudoaglycon. *J. Org. Chem.* **1986**, *51*, 1509.

8. (a) Hunt, A. H.; Molloy, R. M.; Occolowitz, J. L.; Marconi, G. G.; Debono, M. Structure of the Major Glycopeptide of the Teicoplanin Complex. *J. Am. Chem. Soc.* **1984**, *106*, 4891. (b) Barna, J. C. J.; Williams, D. H.; Stone, D. J. M.; Leung, T.-W. C.; Doddrell, D. M. Structure Elucidation of the Teicoplanin Antibiotics. *J. Am. Chem. Soc.* **1984**, *106*, 4895.
9. (a) McGahren, W. J.; Martin, J. H.; Morton, G. O.; Hargreaves, R. T.; Leese, R. A.; Lovell, F. M.; Ellestad, G. A. Avoparcin. *J. Am. Chem. Soc.* **1979**, *101*, 2237. (b) McGahren, W. J.; Martin, J. H.; Morton, G. O.; Hargreaves, R. T.; Leese, R. A.; Lovell, F. M.; Ellestad, G. A.; O'Brien, E.; Holker, J. S. E. Structure of Avoparcin Components. *J. Am. Chem. Soc.* **1980**, *102*, 1671. (c) Ellestad, G. A.; Leese, R. A.; Morton, G. O.; Barbatschi, M. F.; Gore, W. E.; McGahren, W. J. Avoparcin and Epiavoparcin. *J. Am. Chem. Soc.* **1981**, *103*, 6522.
10. (a) Hunt, A. H.; Debono, M.; Merkel, K. E.; Barnhart, M. Structure of the Pseudoaglycon of Actaplanin. *J. Org. Chem.* **1984**, *49*, 635. (b) Hunt, A. H.; Elzey, T. K.; Merkel, K. E.; Debono, M. Structure of the Actaplanins. *J. Org. Chem.* **1984**, *49*, 641.
11. Townsend, C. A.; Brown, A. M. Nocardin A: Biosynthetic Experiments with Amino Acid Precursors see also : Total Synthesis of monocyclic  $\beta$ -Lactam Antibiotics, Nocardicin A and D. *J. Am. Chem. Soc.* **1983**, *105*, 913. (See also: Kamiya, T.; Hashimoto, M.; Nakaguchi, O.; Oku, T. *Tetrahedron* **1979**, *35*, 323)
12. (a) Spencer, J. L.; Flynn, E. H.; Roeske, R. W.; Siu, F. Y.; Chauvette, R. R. Chemistry of Cephalosporin Antibiotics. VII. Synthesis of Cephaloglycin and some Homologs. *J. Med. Chem.* **1966**, *9*, 746. (b) Ryan, C. W.; Simon, R. L.; Van Heyningen, E. M. Chemistry of Cephalosporin Antibiotics. XIII. Desacetoxycephalosporins. The Synthesis of Cephalexin and some Analogs. *J. Med. Chem.* **1969**, *12*, 310. (c) Chauvette, R. R.; Pennington, P. A. Chemistry of Cephalosporin Antibiotics. 30. 3-Methoxy- and 3-Halo-3-cephems. *J. Med. Chem.* **1975**, *18*, 403.
13. Meijer, E. M.; Boesten, W. H. J.; Schoemaker, H. E.; van Balken, J. A. M. *Biocatalysts in Organic Synthesis*; Tramper, J., van der Plas, H. C., Linko, P., Eds; Elsevier : Amsterdam, 1985.

14. (a) Knöpfel, T.; Kuhn, R.; Allgeier, H. Metabotropic Glutamate Receptors : Novel Targets for Drug Development. *J. Med. Chem.* **1995**, *38*, 1418. (b) Hatakeyama, S.; Yoshida, M.; Esumi, T.; Iwabuchi, Y.; Irie, H.; Kawamoto, T.; Yamada, H.; Nishizawa, M. A Novel Enantioselective Synthesis of (+)-Myriocin Based on the Chemistry of 1-Trimethylsilylbuta-2, 3-dienes. *Tetrahedron Lett.* **1997**, *38*, 7887.
15. (a) Hayashi, Y.; Sekiyama, N.; Nakanishi, S.; Jane, D. E.; Sunter, D. C.; Bire, E. F.; Udvarhelyi, P. M.; Watkins, J. C. Analysis of Agonist and Antagonist Activities of Phenylglycine Derivatives for Different Cloned Metabotropic Glutamate Receptor Subtypes. *J. Neurosci.* **1994**, *14*, 3370. (b) Ma, D.; Tian, H. Stereoselective Synthesis of (*S*)-(+)– $\alpha$ M4CPG, a Selective Antagonist of Metabotropic Glutamate Receptors. *Tetrahedron : Asymmetry* **1996**, *6*, 1567.
16. Jane, D. E.; Pittaway, K.; Sunter, D. C.; Thomas, N. K.; Watkins, J. C. New Phenylglycine Derivatives with Potent and Selective Antagonist Activity at Presynaptic Glutamate Receptors in Neonatal Rat Spinal Cord. *Neuropharmacology* **1995**, *34*, 851.
17. For reviews, see the following. (a) Wirth, T. New Strategies to  $\alpha$ -Alkylated  $\alpha$ -Amino Acids. *Angew. Chem., Int. Ed. Engl.* **1997**, *36*, 225. (b) Seebach, D.; Sting, A. R.; Hoffmann, M. Self-Regeneration of Stereocenters (SRS)—Applications, Limitations, and Abandonment of a Synthetic Principle. *Angew. Chem., Int. Ed. Engl.* **1996**, *35*, 2708.
18. (a) Baldwin, J. E.; Lee, V.; Schofield, C. J. Synthesis of A bicyclic  $\gamma$ -Lactam Dipeptide Analogue. *Heterocycles* **1992**, *34*, 903. (b) Badorre, R.; Cativiela, C.; Diaz-de-Villegas, M. D.; Galvez, J. A. *Tetrahedron : Asymmetry* **1995**, *6*, 2787. (c) Boyce, R. J.; Mulqueen, G. C.; Pattenden, G. Total Synthesis of Thiangazole, a Novel Naturally Occurring HIV-1 Inhibitor from *Polyangium* sp. *Tetrahedron* **1995**, *51*, 7321. (d) Gershonov, E.; Granoth, R.; Tzeboval, E.; Gaoni, Y.; Fridkin, M. 1-Aminocyclobutanecarboxylic Acid Derivatives as Novel Structural Elements in Bioactive Peptides : Application to Tuftsin Analogs. *J. Med. Chem.* **1996**, *39*, 4833.
19. Strecker, A. *Ann. Chem. Pharm.* **1850**, *75*, 27.
20. March, J. *Advanced Organic Chemistry*, 2<sup>nd</sup> ed., Tokyo, Kogakusha ; McGraw-Hill , 1977, p.874.

21. Calmes, M.; Daunis, J. How to Build Optically Active  $\alpha$ -Amino Acids. *Amino Acids* **1999**, *16*, 215.
22. For recent reviews on this subject, see : (a) Willams, R. M. *Synthesis of Optically Active  $\alpha$ -Amino Acids*. Oxford; Pergamon Press: 1989. (b) Duthaler, R. O. Recent Developments in the Stereoselective Synthesis of  $\alpha$ -amino acids. *Tetrahedron* **1994**, *50*, 1539.
23. Harusawa, S.; Hamada, Y.; Shioiri, T. Diethyl phosphorocyanide (DEPC). A novel Reagent for the classical Strecker's  $\alpha$ -amino nitriles synthesis. *Tetrahedron Lett.* **1979**, *48*, 4663.
24. (a) Harada, K. Asymmetric Synthesis of  $\alpha$ -Amino acids by the Strecker Synthesis. *Nature* **1963**, *200*, 1201. (b) Harada, K.; Okawara, T. Sterically Controlled Syntheses of Optically Active Organic Compounds. XVIII. Asymmetric Syntheses of Amino Acids by Addition of Hydrogen Cyanide to the Schiff Bases. *J. Org. Chem.* **1973**, *38*, 707. (c) Patel, M. S.; Worsely, M. Asymmetric Syntheses of Optically Active  $\alpha$ -Amino Acids by Hydrocyanic Acid Addition to the Optically Active Schiff Bases. *Can. J. Chem.* **1970**, *48*, 1881. (d) Weinges, K.; Gries, K.; Stemmle, B.; Schrank, W. Über die asymmetrische Strecker-Synthese mit (*S*)-(-)-1-Phenylethylamin als chiralem Hilfsreagens. *Chem. Ber.* **1997**, *110*, 2098. (e) Stout, D. M.; Black, L. A.; Matier, W. L. Asymmetric Strecker Synthesis : Isolation of Pure Enantiomers and Mechanistic Implications. *J. Org. Chem.* **1983**, *48*, 5369. (f) Phadtare, S. K.; Kamat, S. K.; Panse, G. T. Asymmetric Synthesis of  $\alpha$ -Amino Acids. *Ind. J. Chem.* **1985**, *24B*, 811. (g) Subramanian, P. K.; Woodard, R. W. An Asymmetric Strecker Synthesis of (*R*)-(+)z-Methyl-3-Phenylalanine: An Efficient Preparation. *Synth. Commun.* **1986**, *16*, 337. (h) Saito, K.; Harada, K. Asymmetric Syntheses of Amino Acids By Addition of Cyanide to the Schiff Bases in the Presence of Cyanide-Modified Hemin-Copolymer. *Tetrahedron Lett.* **1989**, *30*, 4535. (i) Speelman, J. C.; Talma, A. G.; Kellogg, R. M. Molecular Structure of a Chiral 3,5-Bridged Pyridine and the Effect of Structure on Circular Dichroic Spectra. *J. Org. Chem.* **1989**, *54*, 1055. (j) Herranz, R.; Suárez-Gea, M. L.; Vinuesa, S.; García-López, M. T.; Martínez, A. Synthesis of  $\Psi$  [CH(CN)NH] Pseudopeptides. A New Peptide Bond Surrogate. *Tetrahedron Lett.* **1991**, *32*, 7579. (k) Inaba, T.; Fujita, M.;

- Ogura, K. Thermodynamically Controlled 1,3-Asymmetric Induction in an Acyclic System: Equilibration of  $\alpha$ -Amino Nitriles Derived From  $\alpha$ -Alkylbenzylamines and Aldehydes. *J. Org. Chem.* **1991**, *56*, 1274. (l) Chakraborty, T. K.; Reddy, G. V.; Hussain, K. A. Diastereoselective Strecker Synthesis using  $\alpha$ -Phenylglycinol as Chiral Auxiliary. *Tetrahedron Lett.* **1991**, *32*, 7597. (m) Inaba, T.; Kozono, I.; Fujita, M.; Ogura, K. An Efficient and Practical Synthesis of L- $\alpha$ -Amino Acids Using (*R*)-Phenylglycinol as a Chiral Auxiliary. *Bull. Chem. Soc. Jpn.* **1992**, *65*, 2359. (n) Davis, F. A.; Reddy, R. E.; Portonovo, P. S. Asymmetric Strecker Synthesis using Enantiopure Sulfinimines: A Convenient Synthesis of  $\alpha$ -Amino Acids. *Tetrahedron Lett.* **1994**, *35*, 9351. (o) Chakraborty, T. K.; Hussain, K. A.; Reddy, G. V.  $\alpha$ -Phenylglycinol as Chiral Auxiliary in Diastereoselective Strecker Synthesis of  $\alpha$ -Amino Acids. *Tetrahedron* **1995**, *51*, 9179.
25. (a) Kunz, H.; Sager, W. Diastereoselective Strecker Synthesis of  $\alpha$ -Aminonitriles on Carbohydrate Templates. *Angew. Chem., Int. Ed. Engl.* **1987**, *26*, 557. (b) Kunz, H.; Pfrengle, W. Asymmetric Synthesis on Carbohydrate Templates: Stereoselective Ugi Synthesis of  $\alpha$ -Amino Acid Derivatives. *J. Am. Chem. Soc.* **1988**, *110*, 651. (c) Kunz, H.; Sager, W.; Pfrengle, W.; Schanzenbach, D. Reversal of Asymmetric Induction in Stereoselective Strecker Synthesis on Galactosyl Amine as the Chiral Matrix. *Tetrahedron Lett.* **1988**, *29*, 4397. (d) Kunz, H.; Pfrengle, W. Carbohydrates as Chiral Templates: Asymmetric Ugi-Synthesis of Alpha- Amino Acids using Galactosylamines as the Chiral Matrices. *Tetrahedron* **1988**, *44*, 5487.
26. Iyer, M. S.; Gigstad, K. M.; Namdev, N. D.; Lipton, M. A. Asymmetric Catalysis of the Strecker Amino acid Synthesis by a Cyclic Dipeptide. *J. Am. Chem. Soc.* **1996**, *118*, 4910.
27. (a) Oku, J.; Inoue, S. Asymmetric Cyanohydrin Synthesis catalyzed by a Synthetic Cyclic Dipeptide. *J. Chem. Soc., Chem. Commun.* **1981**, 229. (b) Tanaka, K.; Mori, A.; Inoue, S. Cyclic Dipeptide *cyclo[(S)-Phenylalanyl-(S)-histidyl ]* as a Catalyst for Asymmetric Addition of Hydrogen Cyanide to Aldehydes. *J. Org. Chem.* **1990**, *55*, 181. (c) Danda, H.; Nishikawa, H.; Otaka, K. Enantioselective Autoinduction in the Asymmetric Hydrocyanation of 3-

- Phenoxy benzaldehyde Catalyzed by *Cyclo[(R)]-phenylalanyl-(R)-histidyl*. *J. Org. Chem.* **1991**, *56*, 6740.
28. Sigman, M. S.; Jacobsen, E. N. Enantioselective Addition of Hydrogen Cyanide to Imines Catalyzed by a Chiral (Salen) Al (III) Complex. *J. Am. Chem. Soc.* **1998**, *120*, 5315.
29. Sigman, M. S.; Jacobsen, E. N. Schiff Base Catalysts for the Asymmetric Strecker Reaction Identified and Optimized from Parallel Synthetic Libraries. *J. Am. Chem. Soc.* **1998**, *120*, 4901.
30. Sigman, M. S.; Vachal, P.; Jacobsen, E. N. A General Catalyst for the Asymmetric Strecker Reaction. *Angew. Chem., Int. Ed. Engl.* **2000**, *39*, 1279.
31. Krueger, C. A.; Kuntz, K. W.; Dzierba, C. D.; Wirschun, W. G.; Gleason, J. D.; Suapper, M. L; Hoveyda, A. H. Ti-Catalyzed Enantioselective Addition of Cyanide to Imines. A Practice Synthesis of Optically Pure  $\alpha$ -Amino Acids *J. Am. Chem. Soc.* **1999**, *121*, 4284.
32. Takamura, M.; Hamashima, Y.; Usada, H.; Kanai, M.; Shibasaki, M. A Catalytic Asymmetric Strecker-Type Reaction : Interesting Reactivity Difference Between TMSCN and HCN. *Angew. Chem. Int. Ed.* **2000**, *39*, 1650.
33. Mori, M.; Imma, H.; Nakai, T. Asymmetric Catalytic Cyanosilylation of Aldehydes Using a Chiral Binaphthol-Titanium Complex. *Tetrahedron Lett.* **1997**, *38*, 6229.
34. Ishihara, K.; Miyata, M.; Hattori, K.; Tada, T.; Yamamoto, H. A New Chiral BLA Promoter for Asymmetric Aza Diels-Alder and Aldol-Type Reactions of Imines. *J. Am. Chem. Soc.* **1994**, *116*, 10520.
35. Kobayashi, S.; Ishitani, H.; Ueno, M. Catalytic Asymmetric Synthesis of Both Syn-and Anti- $\beta$ -Amino Alcohols. *J. Am. Chem. Soc.* **1998**, *120*, 431.
36. Kobayashi, S.; Komiyama, S.; Ishitani, H. The first Enantioselective Aza-Diels-Alder Reactions of Imino Dienophiles on Use of a Chiral Zirconium Catalyst. *Angew. Chem., Int. Ed. Engl.* **1998**, *37*, 979.
37. Ishitani, H.; Komiyama, S.; Kobayashi, S. Catalytic, Enantioselective Synthesis of  $\alpha$ -Aminonitriles with a Novel Zirconium Catalyst. *Angew. Chem., Int. Ed. Engl.* **1998**, *37*, 3186.

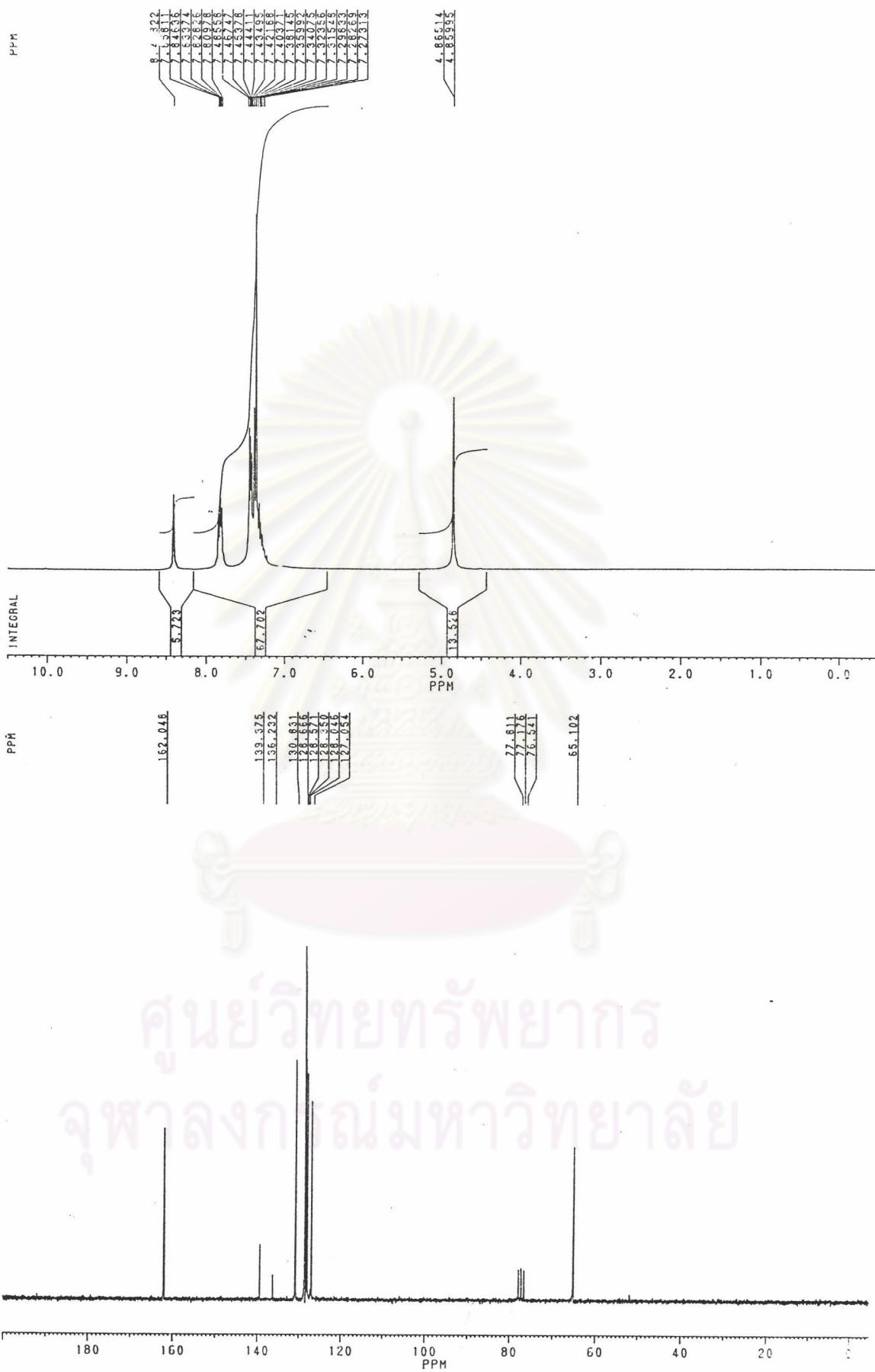
38. Corey, E. J.; Grogan, M. J. Enantioselective Synthesis of  $\alpha$ -Amino Nitriles from *N*-Benzhydryl Imines and HCN with a Chiral Bicyclic Guanidine as Catalyst. *Org. Lett.* **1999**, *1*(1), 157.
39. (a) Zhang, W.; Locbach, J. L.; Wilson, S. R.; Jacobsen, E. N. *J. Am. Chem. Soc.* **1990**, *12*. (b) Zhang, W.; Jacobsen, E. N. *J. Org. Chem.* **1991**, *56*, 2296. (c) Zhang, W.; Muci, A. R.; Eoker, J. R.; Deng, L.; Jacobsen, E. N. *J. Am. Chem. Soc.* **1991**, *113*, 7063.
40. (a) Angelino, M. D.; Laibinis, P. E.; Synthesis and Characterization of a Polymer-Supported Salen Ligand for Enantioselective Epoxidation. *Macromolecules* **1998**, *31*, 7581. (b) Minutolo, F.; Pini, D.; Saluadori, P. Polymer-Bond Chiral (Salen)Mn(III) Complex as Heterogeneous Catalyst in Rapid and Clean Enantioselective Epoxidation of Unfunctionalized Olefins. *Tetrahedron Lett.* **1996**, *37*, 3375. (c) De, B. B.; Lohray, B. B.; Sivaram, S.; Dahl, P. K. Enantioselective Epoxidation of Olefines Catalyzed by Polymer-Bound Optically Active Mn(III)-Salen Complex. *Tetrahedron : Asymmetry* **1995**, *6*, 2105. (d) Peukert, S.; Jacobsen, E. N. Enantioselective Parallel Synthesis Using Polymer-Supported Chiral Co(Salen) Complexes. *Organic Letters* **1999**, *1*, 1245. (e) Angelino, M. D.; Laibinis, P.E. Synthesis and Characterization of a Polymer-Supported Salen Ligand for Enatioselective Epoxidation. *Macromolecules* **1998**, *31*, 7581. (f) Annis, D. A.; Jacobsen, E. N. Polymer-Supported Chiral Co(Salen) Complexes : Synthetic Applications and Mechanistic Investigations in the Hydrolytic Kinetic Resolution of Terminal Epoxides. *J. Am. Chem. Soc.* **1999**, *121*, 4147. (g) Reger, T. S.; Janda, K. D. Polymer-Supported (Salen)Mn Catalysts for Asymmetric Epoxidation : A Comparison between Soluble and Insoluble Matrices. *J. Am. Chem. Soc.* **2000**, *122*, 6929.
41. McEwen, W. E.; Grossi, A. V.; MacDonald, R. J.; Stameyna, A. P. Synthetic Uses of Open-Chain Analogues of Reissert Compounds *J. Org. Chem.* **1980**, *45*, 1301.
42. Aitken, R. A.; Armstrong, D. P.; Galt, R. H. B.; Mesher, S. T. E.; Synthesis and Pyrolytic Behavior of Thiazolidine-2-one 1,1-dioxides. *J. Chem. Soc., Perkin Trans. 1*, **1997**, 2139.

43. (a) Wu, M-J.; Pridgen, L. N. Chiral Oxazolidines as Templates for Stereoselective Synthesis of Chiral Homoallyl Amines. *Syn. lett.* **1990**, 636. (b) Wu, M-J.; Pridgen, L. N. Synthesis of Chiral  $\alpha$ -Alkyl Phenethylamines via Organometallic Addition to Chiral 2-Aryl-1,3-oxazolidines. *J. Org. Chem.* **1991**, *56*, 1340.
44. Takahashi, H.; Suzuki, Y.; Inagaki, H. Asymmetric  $\alpha$ -Substituted Phenethylamines. I. Synthesis of Optically Pure 1-Aryl-N-(2'-hydroxy-1'-isopropylethyl)-2-phenylethylamines *Chem. Pharm. Bull.* **1982**, *30*, 3160.
45. Hayashi, M.; Miyamoto, Y.; Inoue, T.; Oguni, N. Enantioselective Trimethylsilylcyanation of Some Aldehydes Catalyzed by Chiral Schiff Base-Titanium Alkoxide Complexes. *J. Org. Chem.* **1993**, *58*, 1515.
46. Hassan, N. A.; Bayer, E.; Jochims, J. C. Synthese of Optically Active  $\alpha$ -Amino nitriles by Asymmetric Transformation of the Second Kind Using a Principle of O. Dimroth. *J. Chem. Soc., Perkin Trans. I*, **1998**, 3747.
47. Westley, J. W.; Halpern, B. The use of (-)-Menthyl Chloroformate in the Optical Analysis of Asymmetric Amino and Hydroxyl Compounds by Gas Chromatography. *J. Org. Chem.* **1968**, *33*, 3978.
48. Dale, J. A.; Dull, D. L.; Mosher, H. S.  $\alpha$ -Methoxy- $\alpha$ -trifluoromethylphenylacetic Acid, a Versatile Reagent for the Determination of Enantiomeric Composition of Alcohols and Amines. *J. Org. Chem.* **1969**, *34*, 2543.
49. Marvell, C. S.; Noyes, W. A. A Study of the Possible Asymmetry of the Aliphatic Diazo Compounds. *J. Am. Chem. Soc.* **1920**, *42*, 2266.
50. Aitken, R. A. and Kilenyi, S. N. *Asymmetric Synthesis*, Glasgow: Blackie Academic & Professional, 1992.
51. Larrow, J. F. and Jacobson, E. N. A Practical Method for the Large-scale Preparation of [*N*, *N'*-Bis(3,5-*tert*-butylsalicylidene)-1,2-cyclohexane diaminato(2-)]manganese(III)chloride, a Highly Enantioselective Epoxidation Catalyst. *J. Org. Chem.* **1994**, *59*, 1939.



## APPENDICES

ศูนย์วิทยทรัพยากร  
จุฬาลงกรณ์มหาวิทยาลัย



**Fig 1** The  $^1\text{H}$ -NMR and  $^{13}\text{C}$ -NMR spectra of imine 42

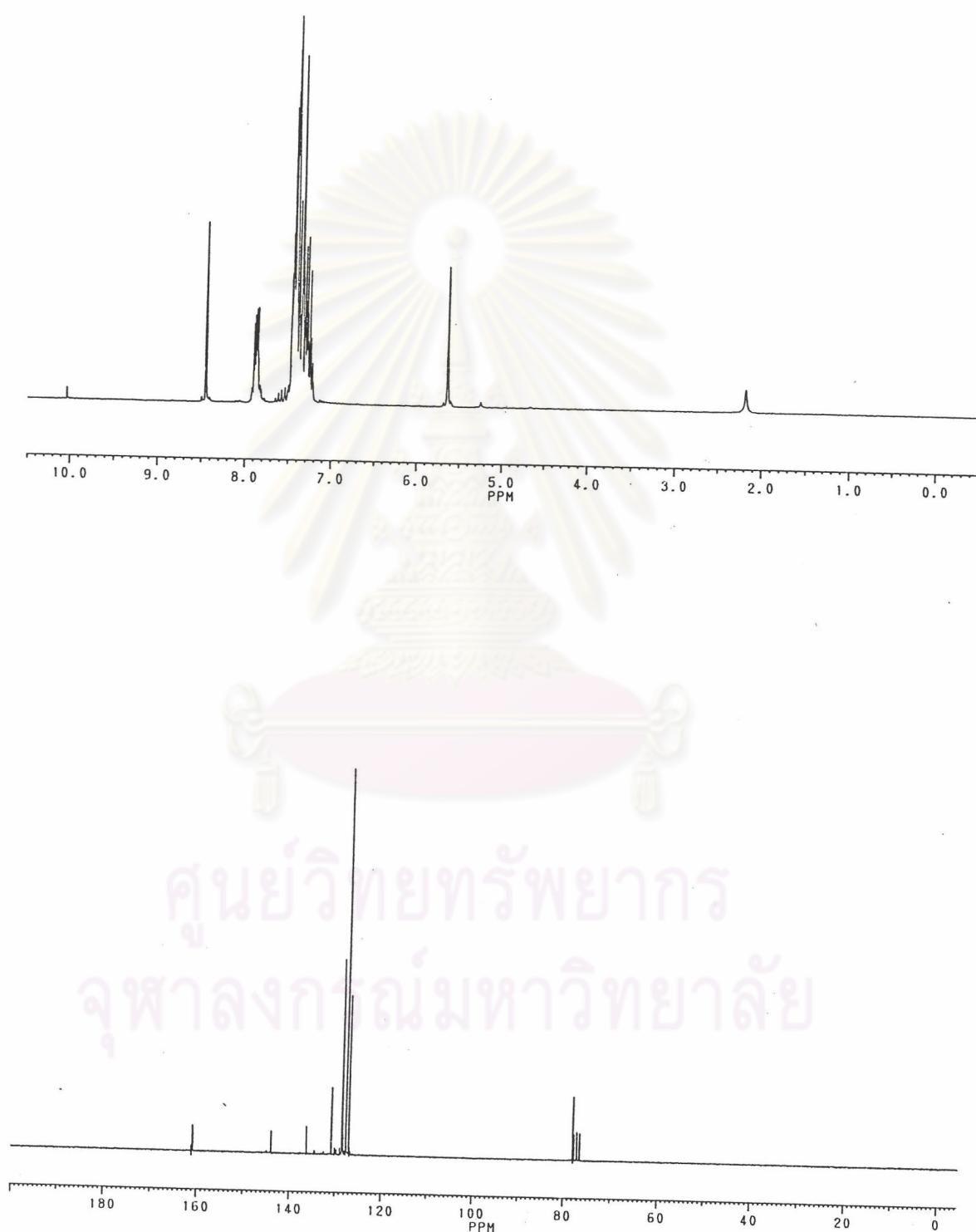
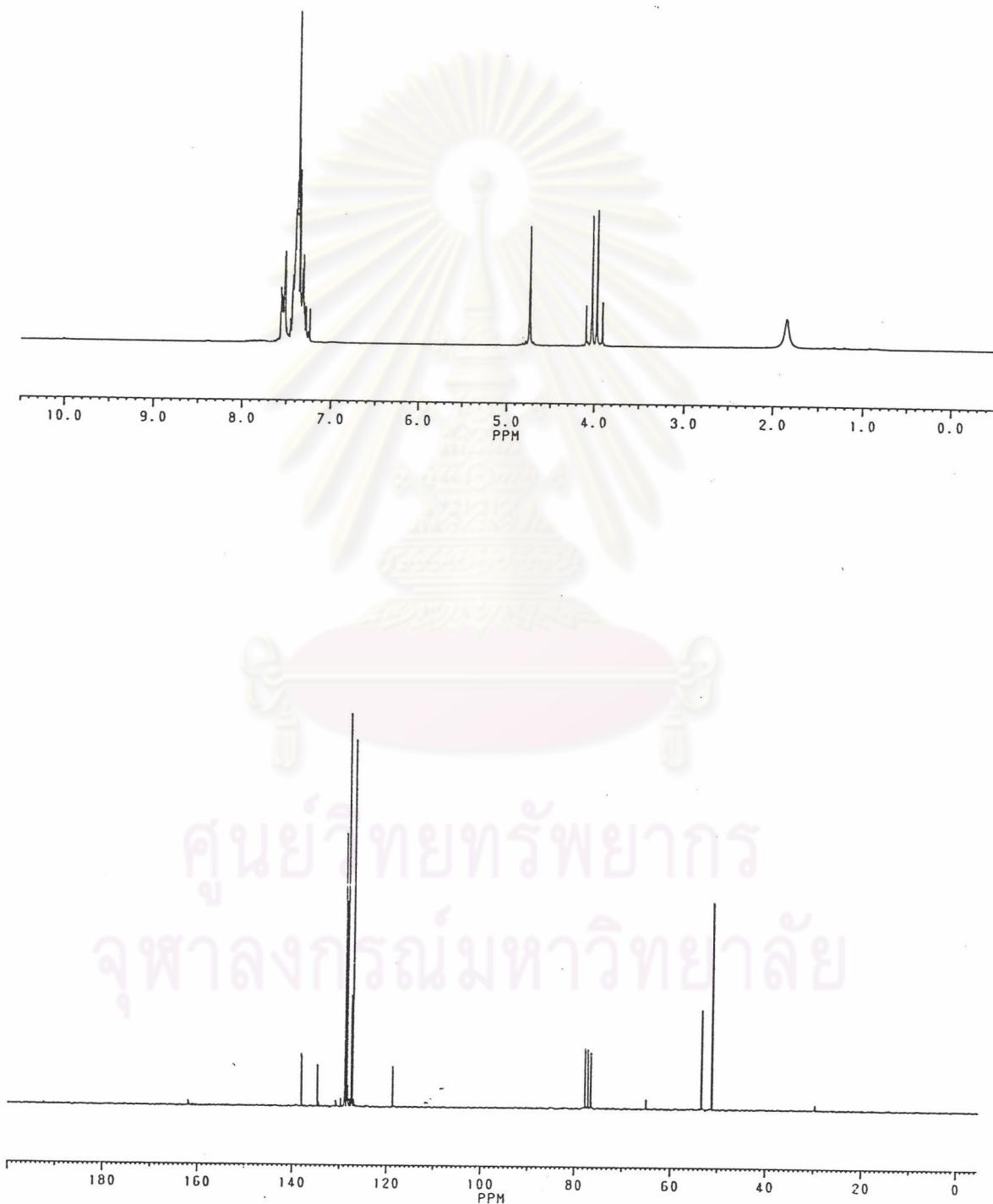
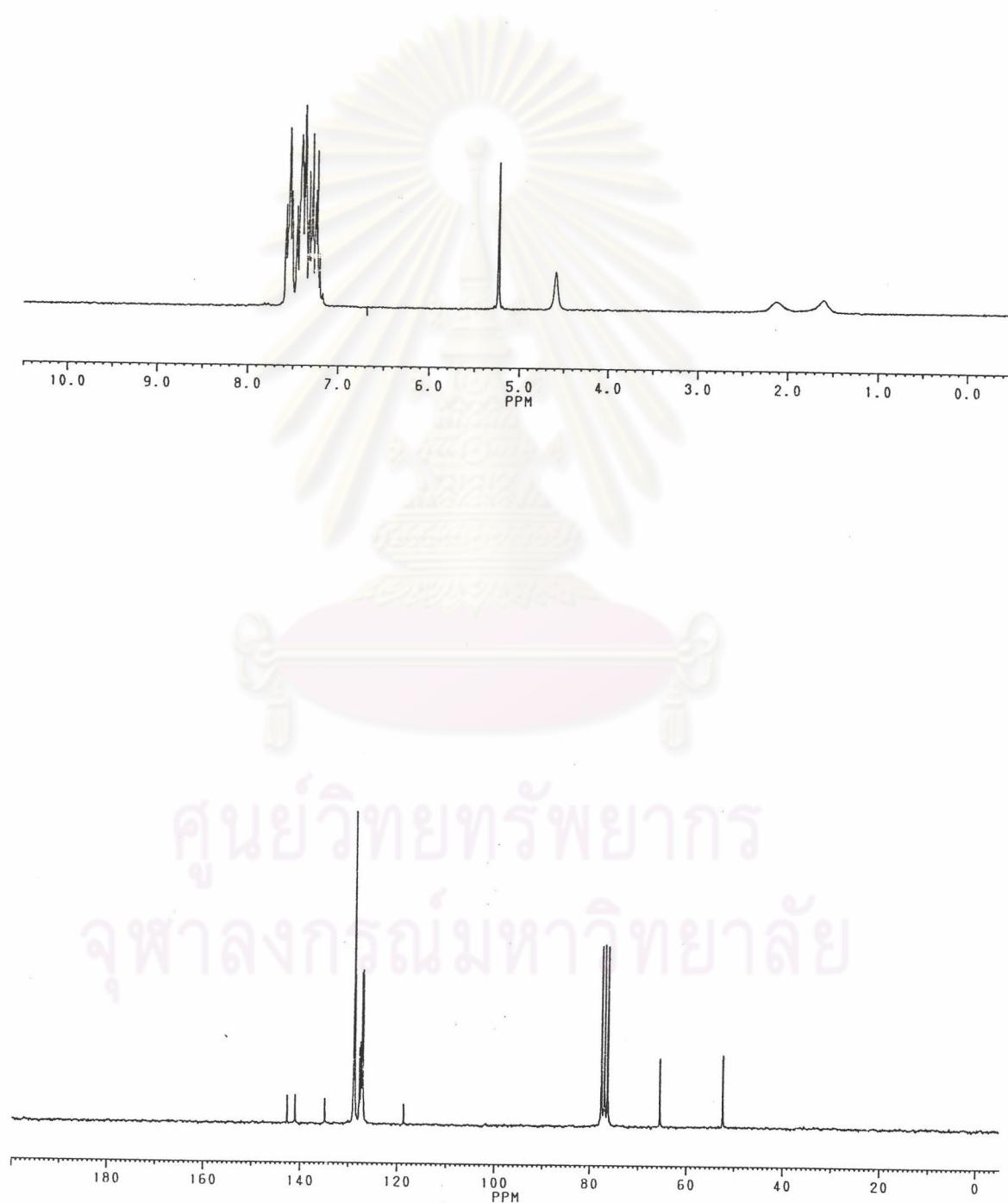


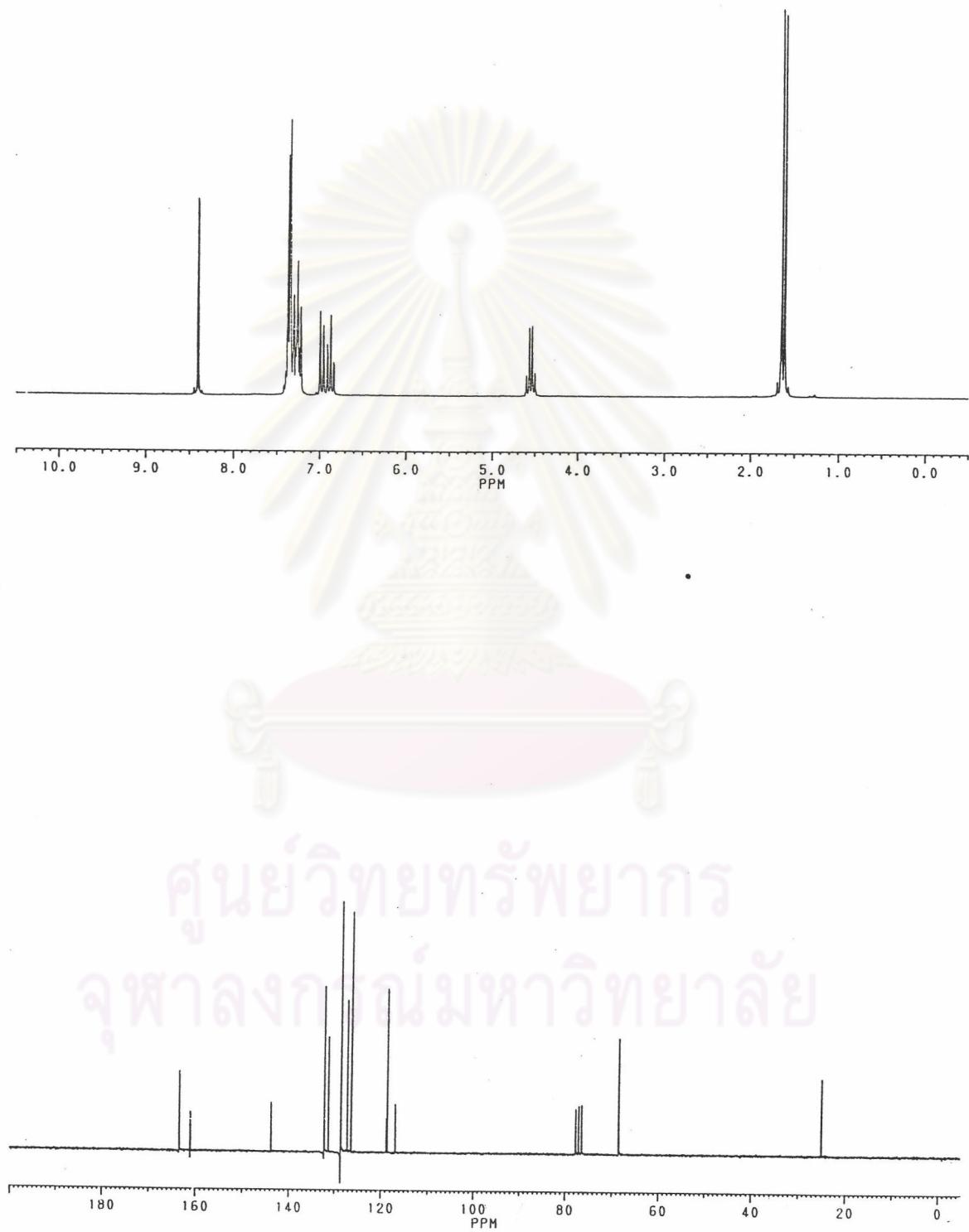
Fig 2 The  $^1\text{H}$ -NMR and  $^{13}\text{C}$ -NMR spectra of imine 24a



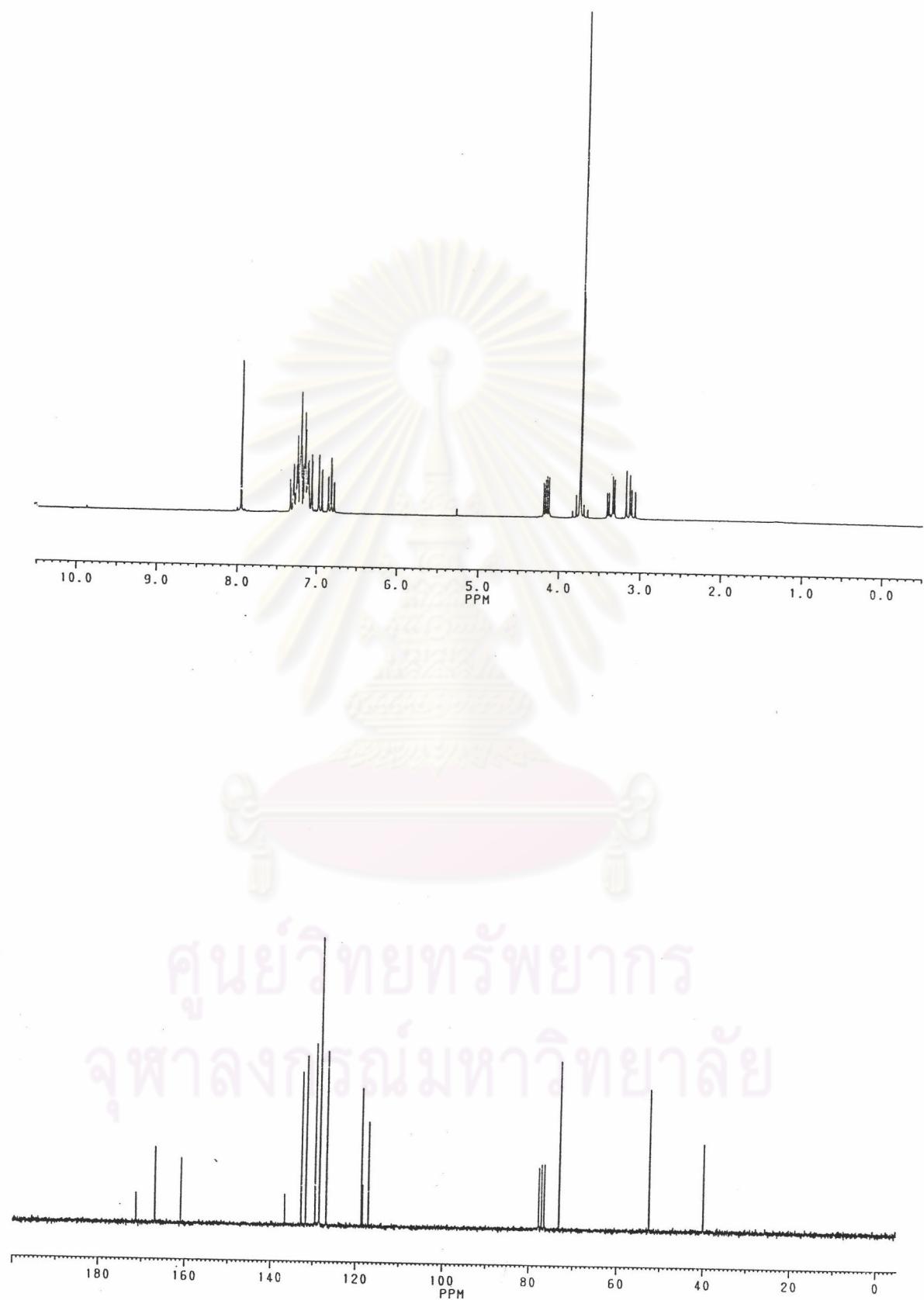
**Fig 3** The  $^1\text{H}$ -NMR and  $^{13}\text{C}$ -NMR spectra of aminonitrile 43



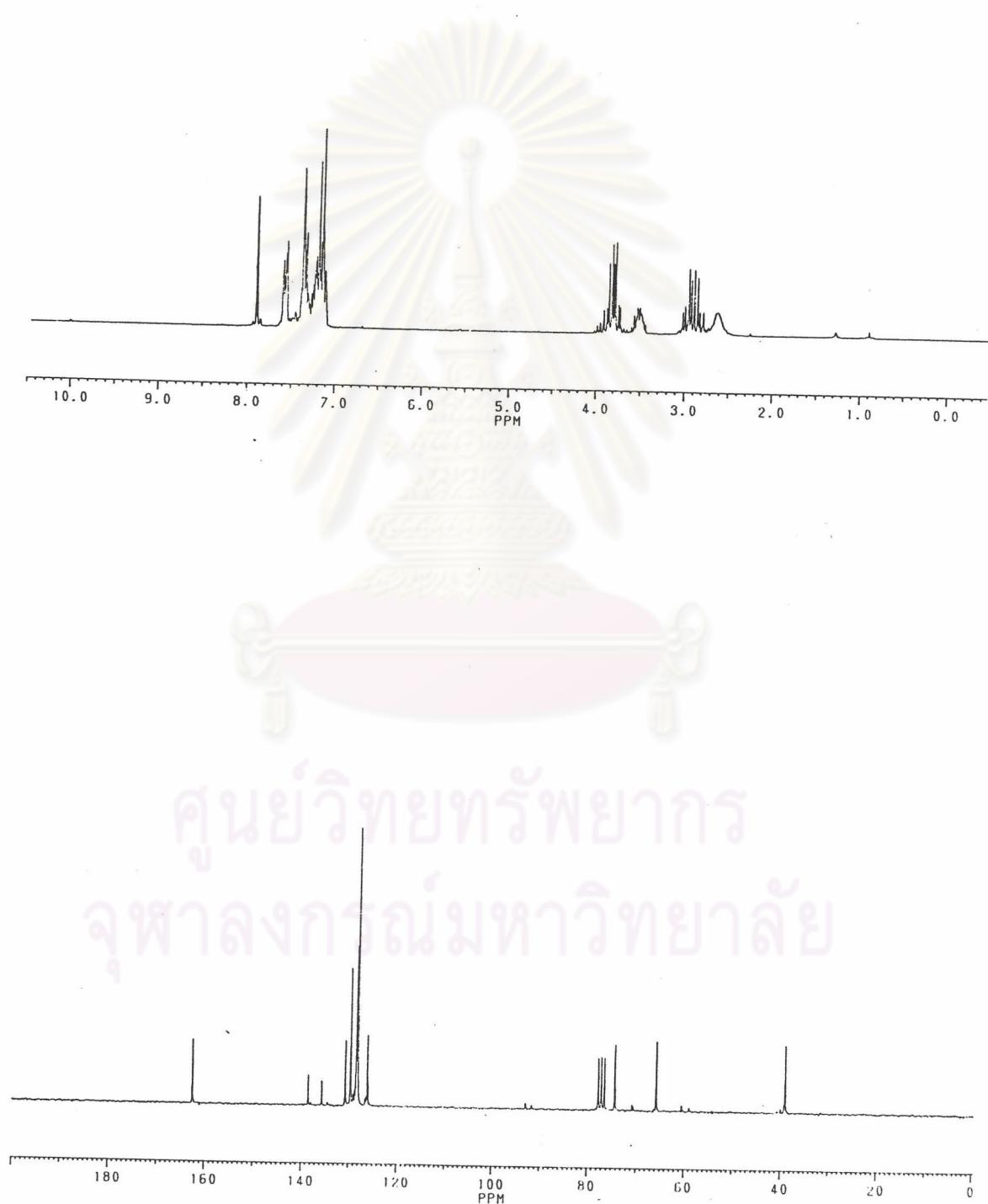
**Fig 4** The  $^1\text{H}$ -NMR and  $^{13}\text{C}$ -NMR spectra of aminonitrile **25a**



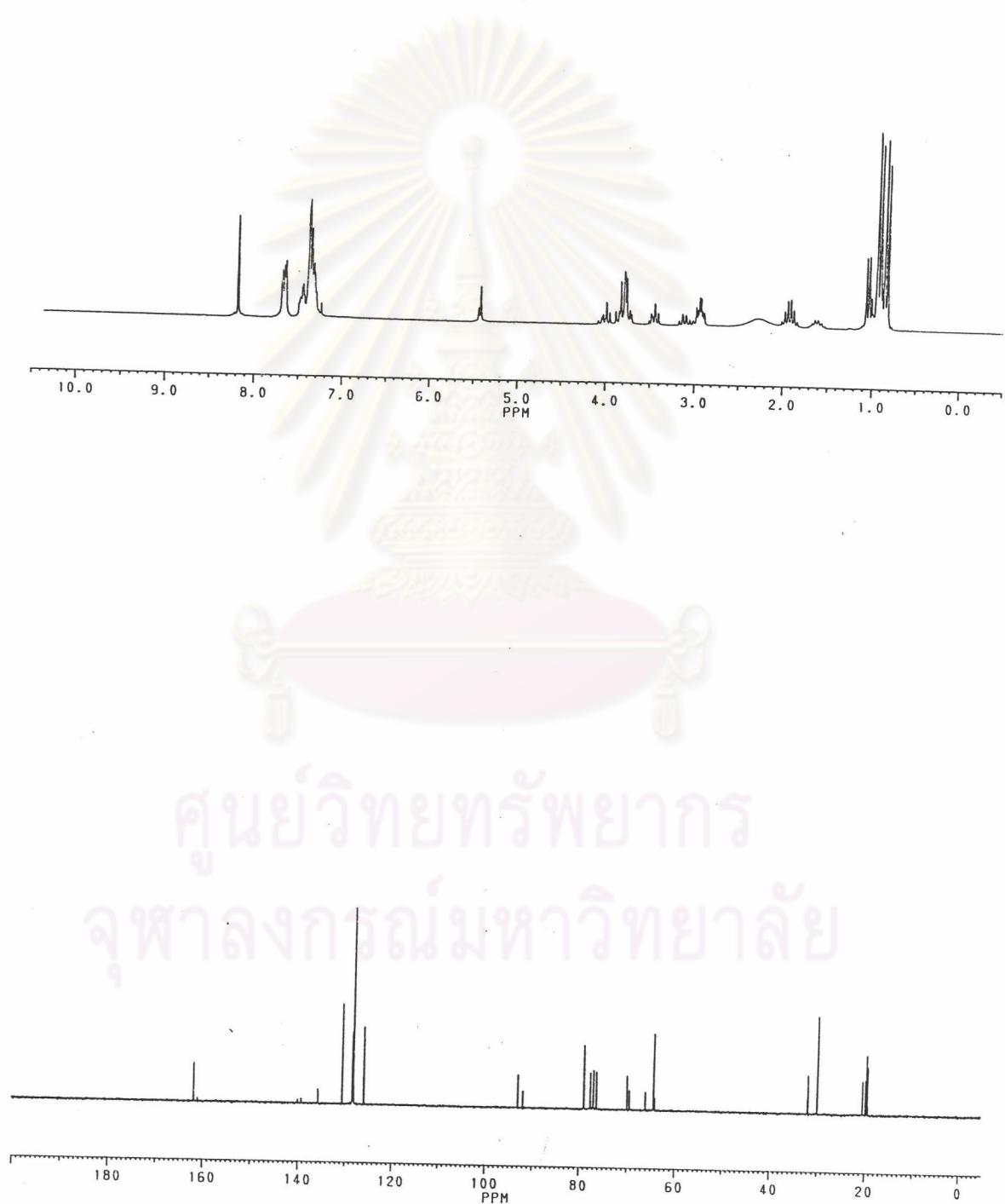
**Fig 5** The  $^1\text{H}$ -NMR and  $^{13}\text{C}$ -NMR spectra of compound 48



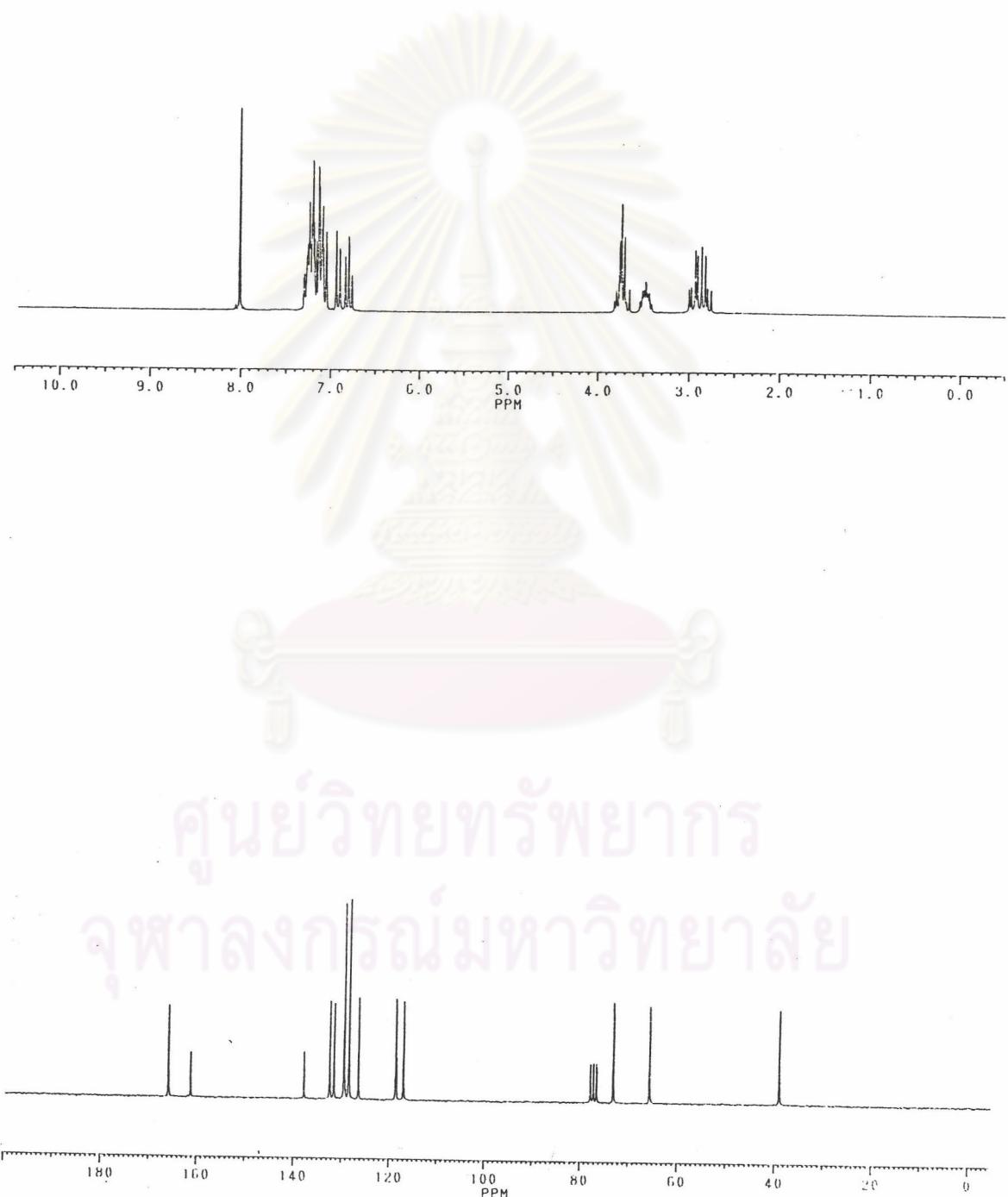
**Fig 6** The  $^1\text{H}$ -NMR and  $^{13}\text{C}$ -NMR spectra of compound 49



**Fig 7** The  $^1\text{H}$ -NMR and  $^{13}\text{C}$ -NMR spectra of compound **50**



**Fig 8** The  $^1\text{H}$ -NMR and  $^{13}\text{C}$ -NMR spectra of compound **51**



**Fig 9** The  $^1\text{H}$ -NMR and  $^{13}\text{C}$ -NMR spectra of compound 52

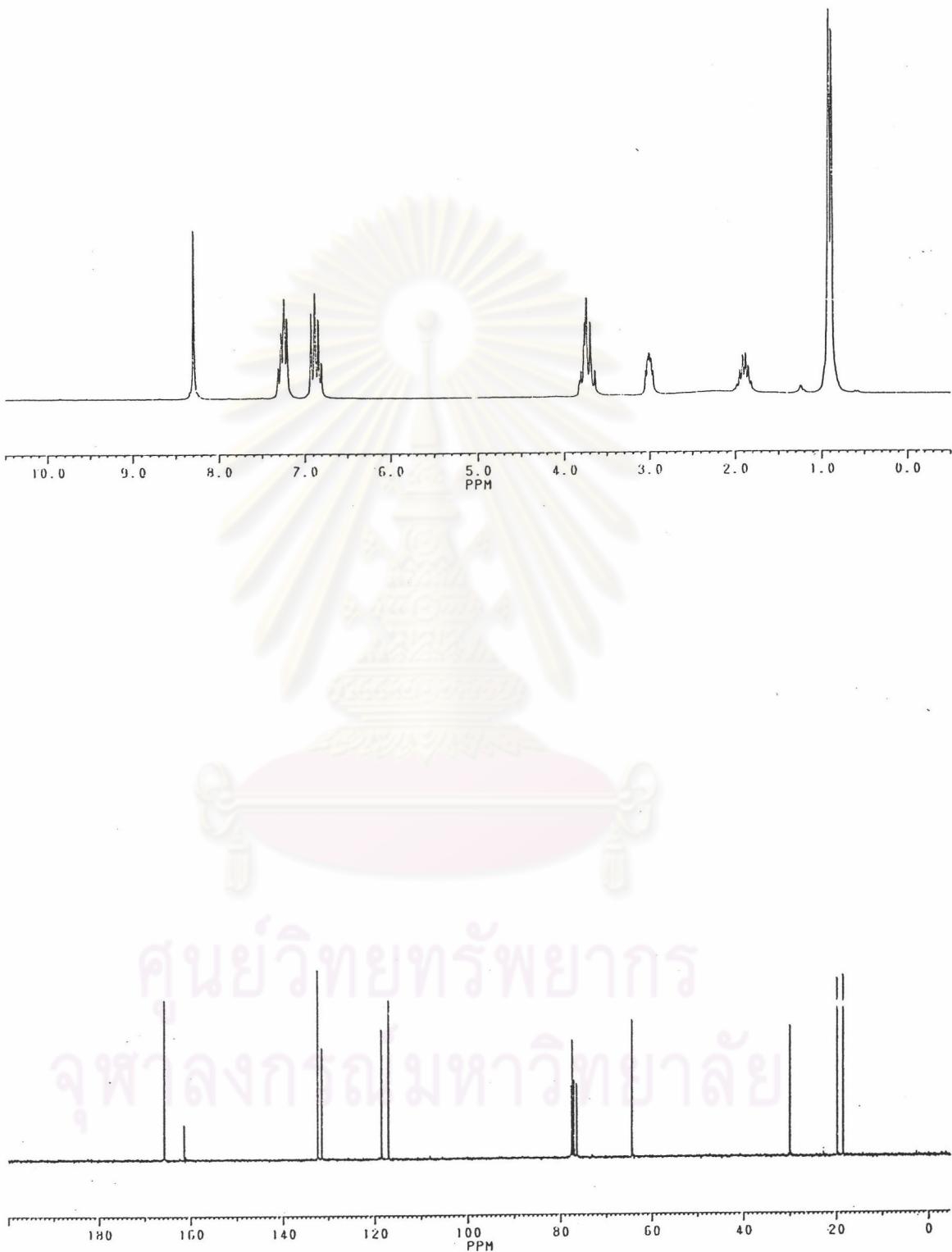


Fig 10 The  $^1\text{H}$ -NMR and  $^{13}\text{C}$ -NMR spectra of compound 53

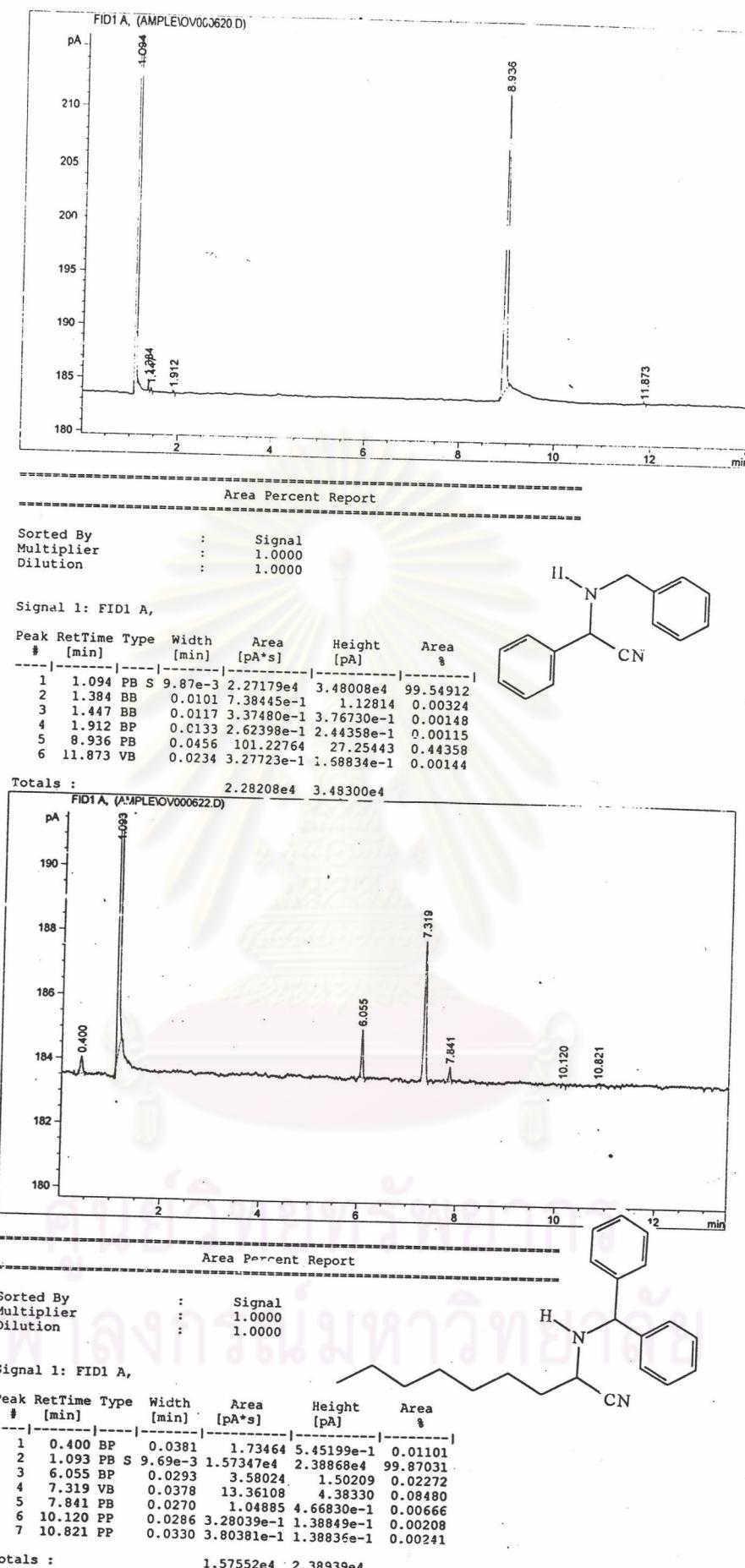
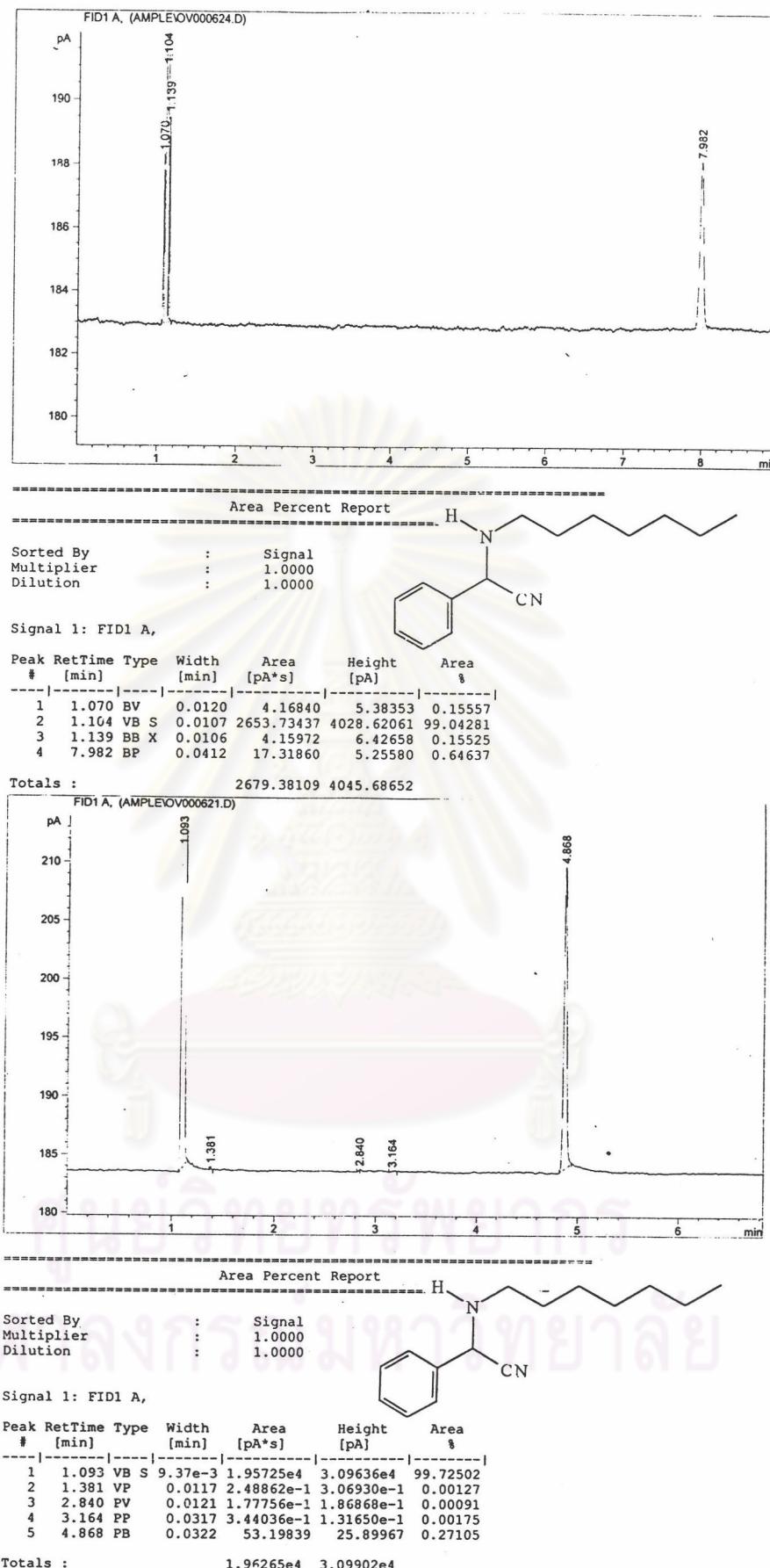
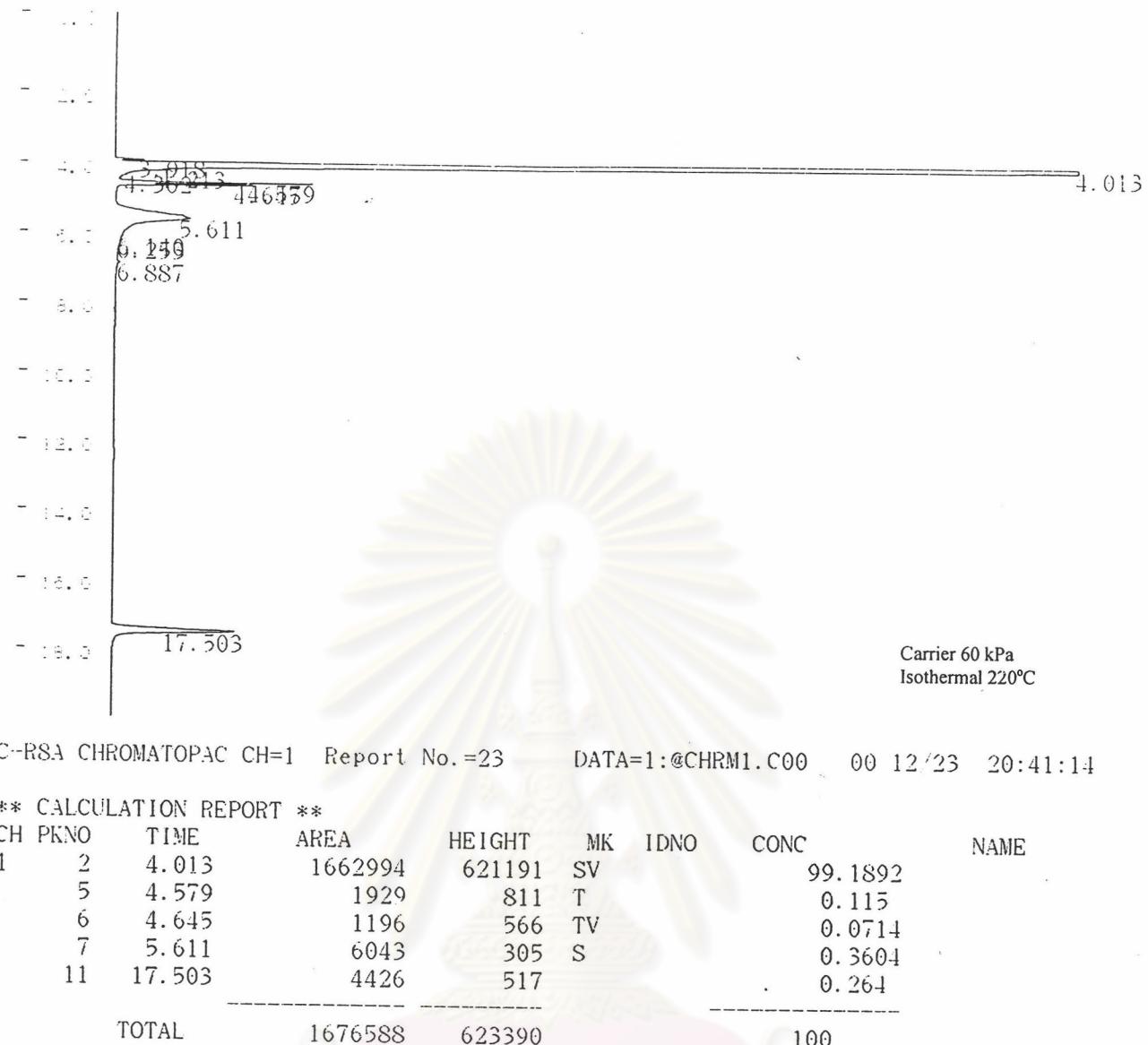


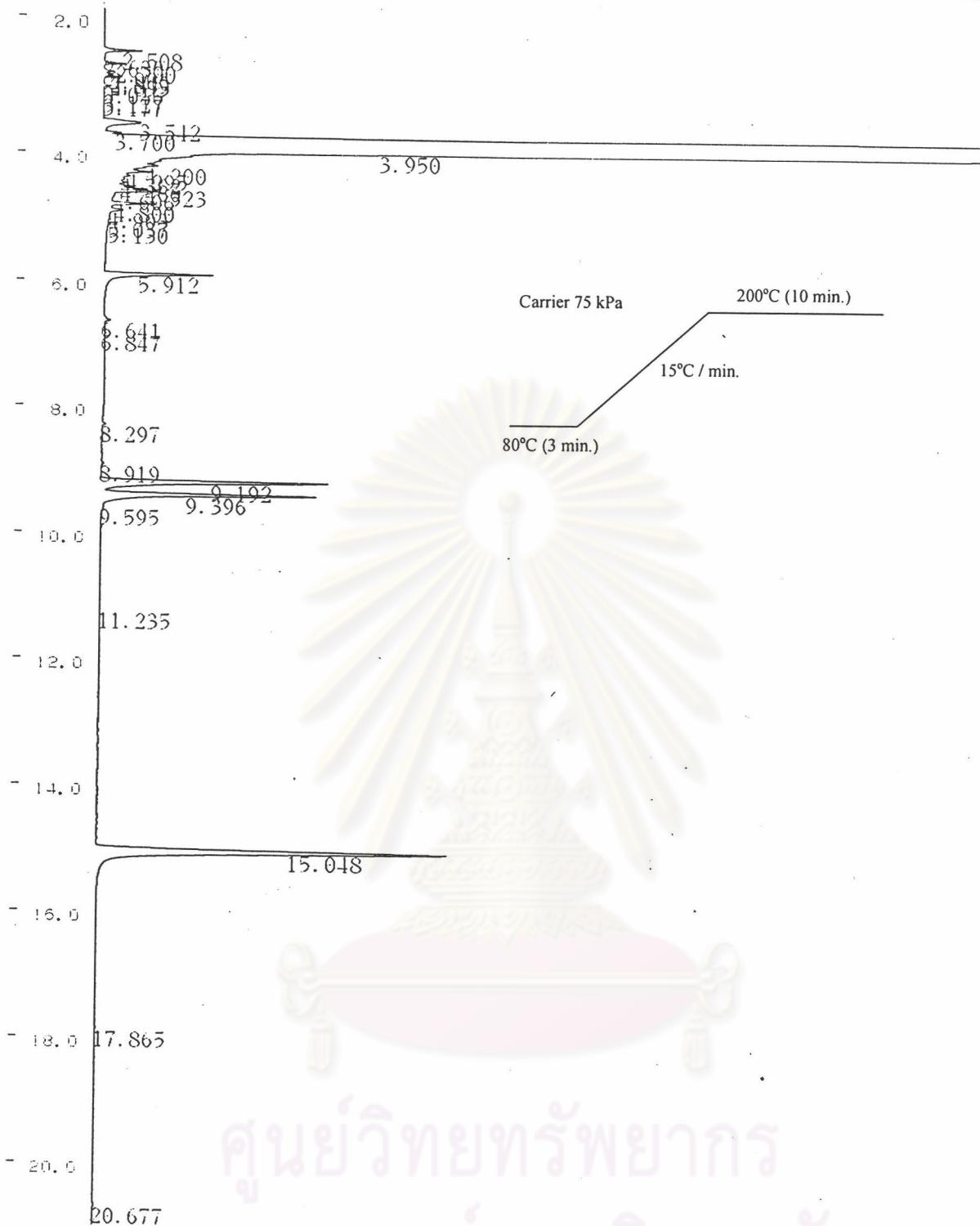
Fig 11 The results of chiral GC analysis; 180 °C isothermal: (a) aminonitrile 43; (b) aminonitrile 56.



**Fig 12** The results of chiral GC analysis; aminonitrile 55: (a) 165 °C isothermal;  
(b) 180 °C isothermal.



**Fig 13** The results of chiral GC analysis; 220 °C isothermal:  
derivatized aminonitrile **25a**.

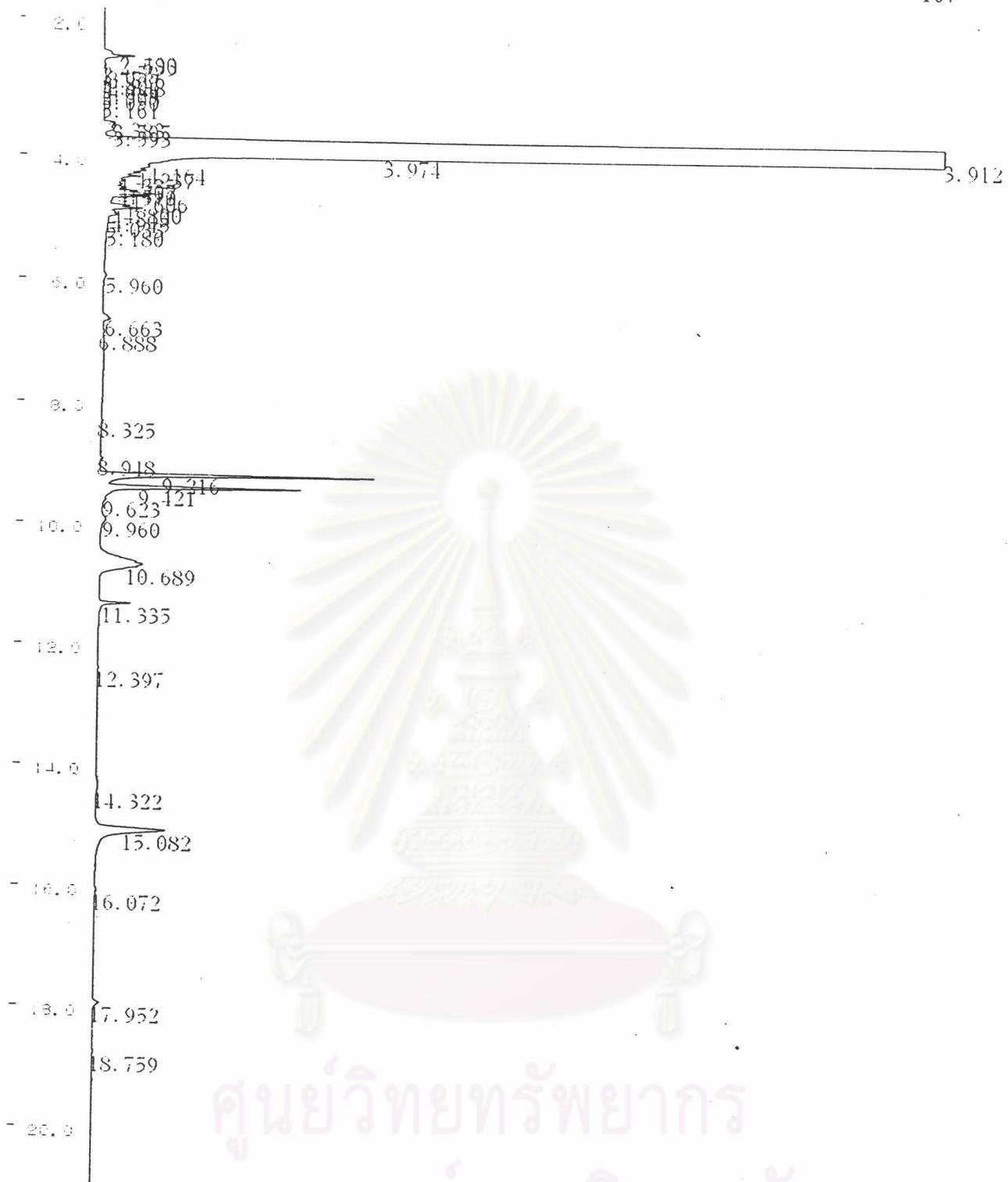


C-RSA CHROMATOPAC CH=1 Report No.=4 DATA=1:@CHRM1.C00 00/12/06 17:54:00

**\*\* CALCULATION REPORT \*\***

CH	PKNO	TIME	AREA	HEIGHT	MK	IDNO	CONC	NAME
1	12	3.95	3307312	477894	SV		99.5073	
	23	5.912		1550		499		0.0466
	28	9.192		3758		1039		0.1131
	29	9.396		3350		970		0.1003
	32	15.048		7717		1599		0.2322
<hr/>		TOTAL	3323688	482001			100	

**Fig 14** The results of chiral GC analysis in temperature program condition:  
derivatized aminonitrile **55**.

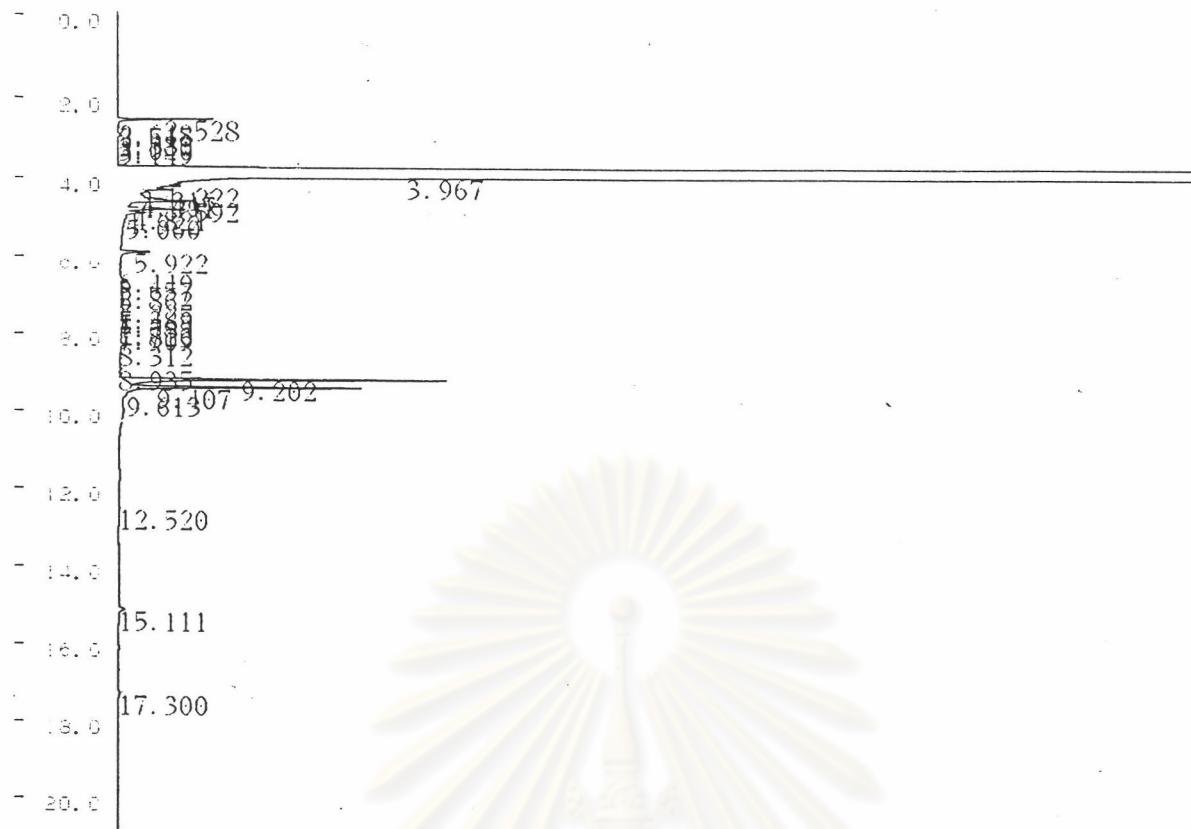


<sup>13</sup>C-RSA CHROMATOPAC CH=1 Report No.=6 DATA=1:@CHRM1.C00 00.12.06 18:48:04

\*\* CALCULATION REPORT \*\*

CH	PKNO	TIME	AREA	HEIGHT	MK	IDNO	CONC	NAME
1	15	3.912	2617404	405924	V		69.1702	
	16	3.974	1154935	390843	SV		30.5215	
	34	9.216	3562	1310			0.0941	
	35	9.421	3111	950	SV		0.0822	
	38	10.689	2525	207			0.0667	
	42	15.082	2471	338			0.0653	
TOTAL			3784007	799572			100	

**Fig 15** The results of chiral GC analysis in temperature program condition:  
derivatized aminonitrile 56.

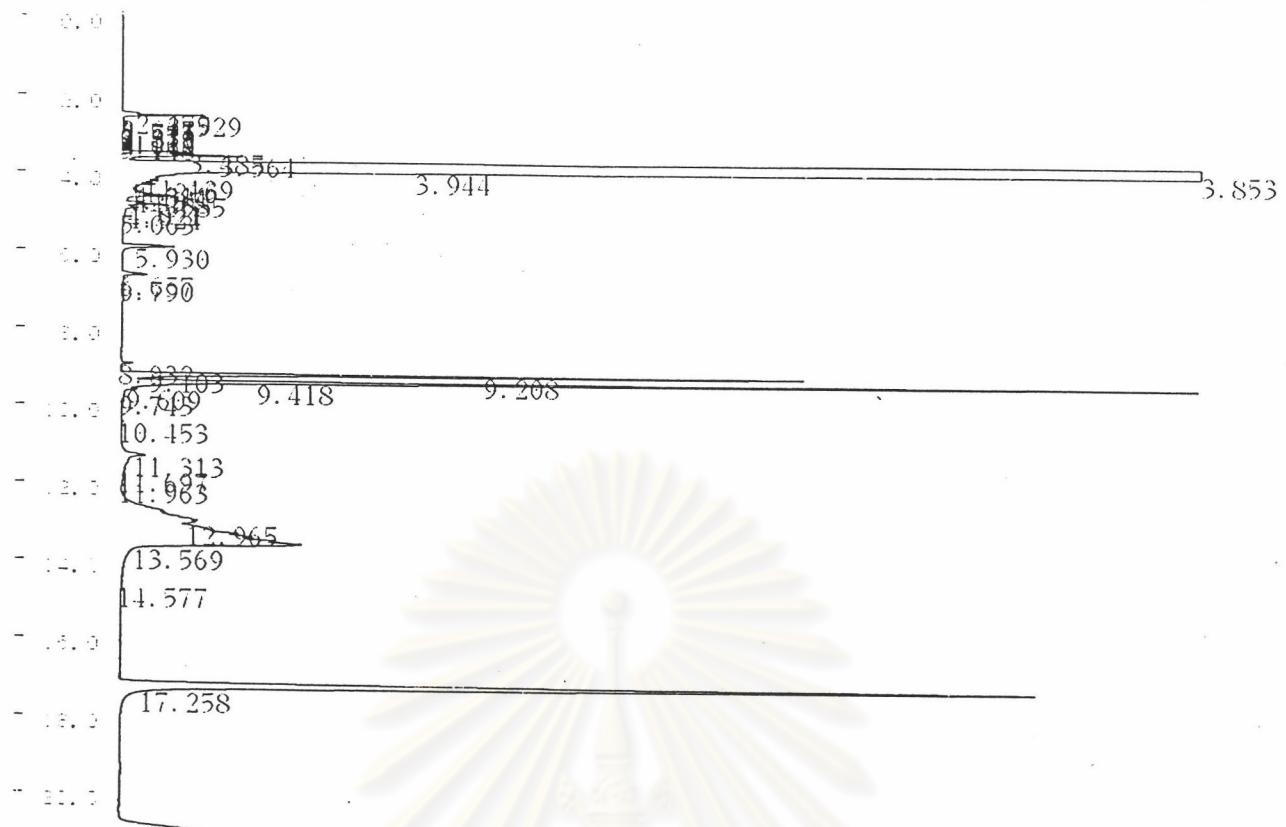


C-R8A CHROMATOPAC CH=1 Report No.=10 DATA=1:\CHRM1.C00 00/12/06 20:30:10

**\*\* CALCULATION REPORT \*\***

CH	PKNO	TIME	AREA	HEIGHT	MK	IDNO	CONC	NAME
1	8	3.967	3682094	424804	S		99.8277	
	30	9.202	3620	1190			0.0981	
	31	9.407	2736	877	SV		0.0742	
TOTAL			3688449	426871			100	

**Fig 16** The results of chiral GC analysis in temperature program condition:  
derivatized aminonitrile **25a**.

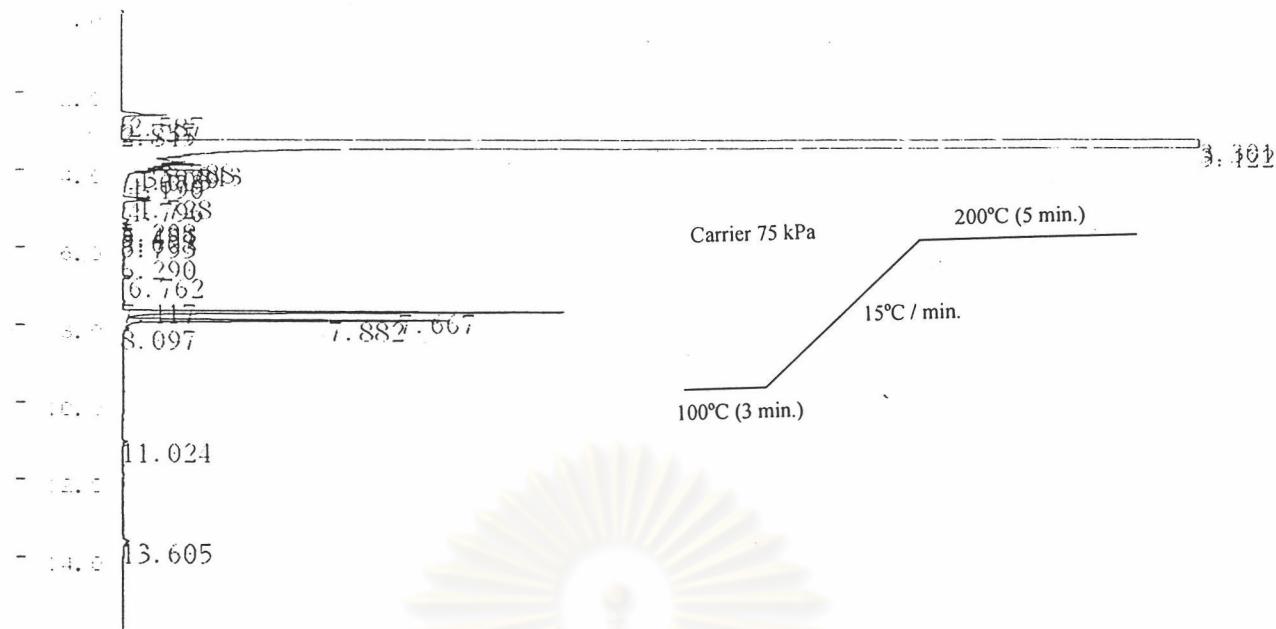


C-RSA CHROMATOPAC CH=1 Report No.=12 DATA=1:@CHRM1.COO 00/12/06 21:22:50

**\*\* CALCULATION REPORT \*\***

CH	PKNO	TIME	AREA	HEIGHT	MK	IDNO	CONC	NAME
1	13	3.564	1618	453	V		0.055	
	14	3.853	1470461	287951	V		49.9344	
	15	3.944	1410010	360207	SV		47.8816	
	31	9.208	7932	2519	V		0.2694	
	32	9.418	12619	4200	SV		0.4285	
	39	12.965	6527	282			0.2216	
	40	13.569	15614	658	SV		0.5302	
	42	17.258	20004	3165			0.6793	
<hr/> TOTAL			2944786	659434			100	

**Fig 17** The results of chiral GC analysis in temperature program condition:  
derivatized aminonitrile **43**.

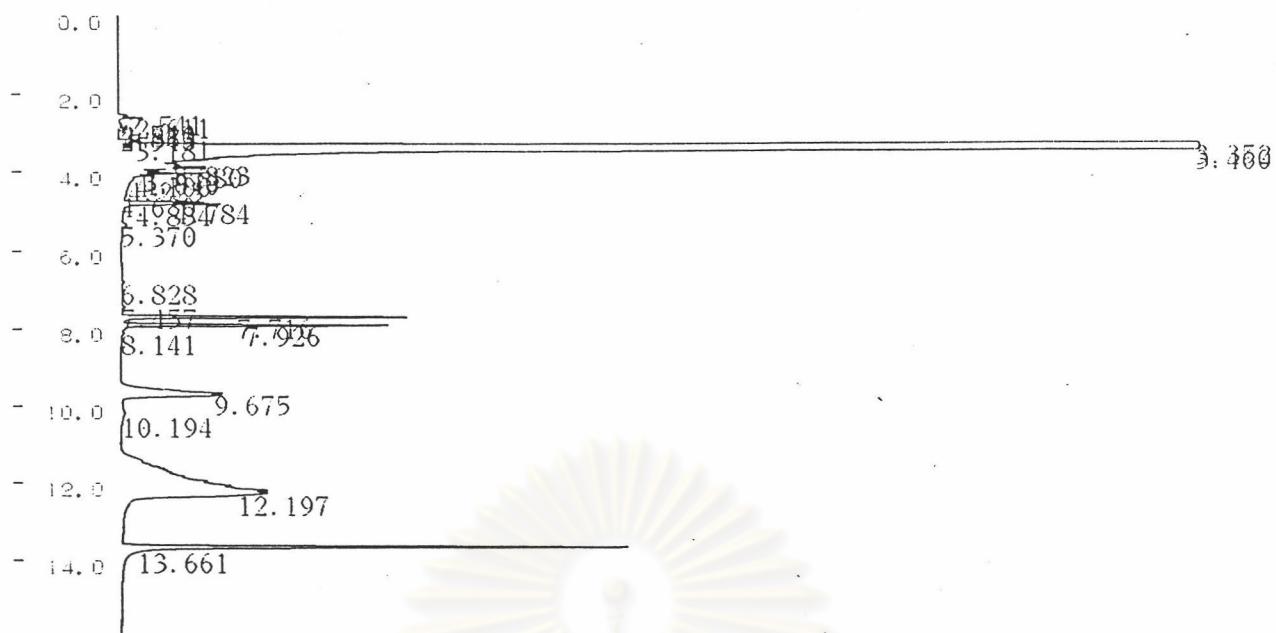


C-RSA CHROMATOPAC CH=1 Report No.=3 DATA=1:@CHRM1.C00 00/12/10 19:23:06

\*\* CALCULATION REPORT \*\*

CH	PKNO	TIME	AREA	HEIGHT	MK	IDNO	CONC	NAME
1	4	3.301	1572539	406329			46.0853	
	5	3.422	1830358	721399	SV		53.6411	
	22	7.667	5448	1633			0.1597	
	23	7.882	3887	1190	SV		0.1139	
		TOTAL	3412232	1130551			100	

**Fig 18** The results of chiral GC analysis in temperature program condition:  
derivatized aminonitrile **25a**.



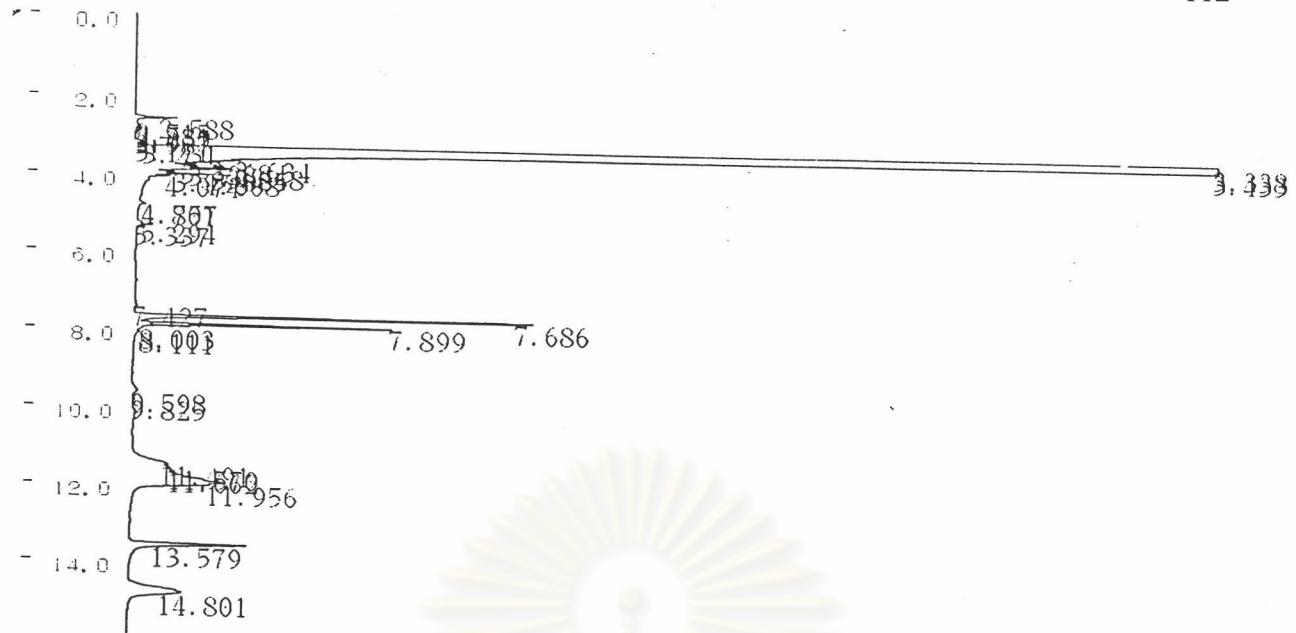
C-R8A CHROMATOPAC CH=1 Report No.=8

DATA=1:@CHRM1.COO 00/12/10 20:57:42

## \*\* CALCULATION REPORT \*\*

CH	PKNO	TIME	AREA	HEIGHT	MK	IDNO	CONC	NAME
1	8	3.352	1161126	360103	V		33.8174	
	9	3.46	2236333	817430	SV		65.1324	
	22	7.716	3470	1044			0.1011	
	23	7.926	3004	983	V		0.0875	
	25	9.675	4227	372			0.1231	
	27	12.197	17883	536			0.5208	
	28	13.661	7477	1786	V		0.2178	
<hr/> TOTAL			3433518	1182254			100	

**Fig 19** The results of chiral GC analysis in temperature program condition:  
derivatized aminonitrile **55**.

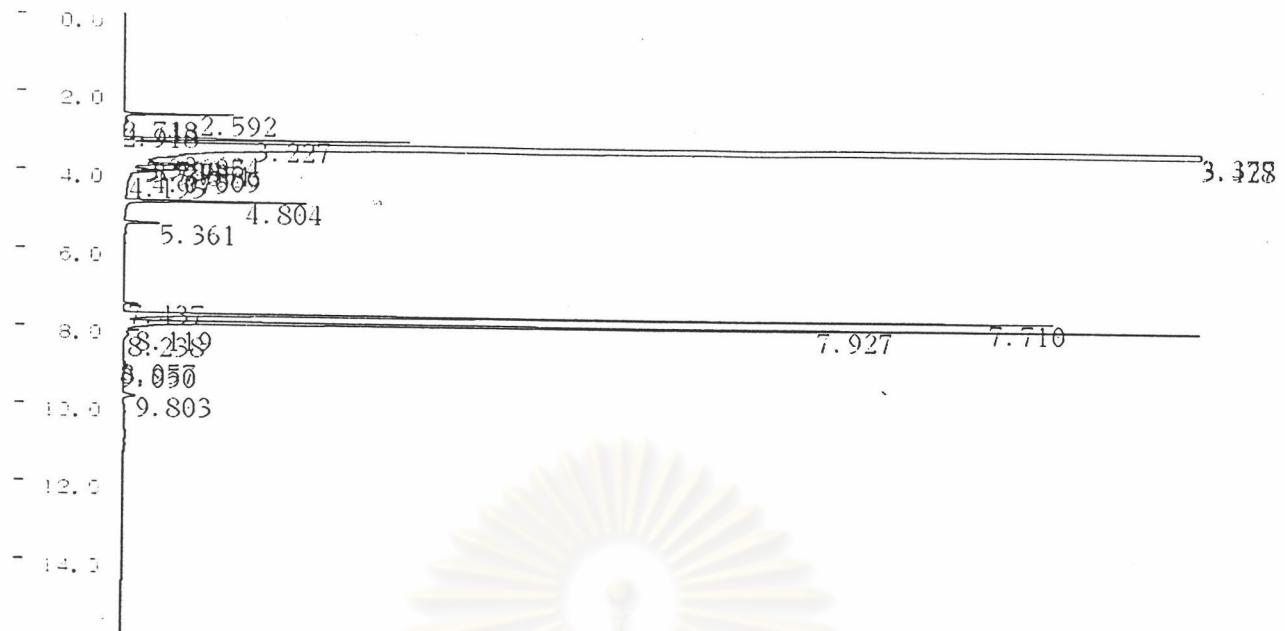


C-R8A CHROMATOPAC CH=1 Report No.=10 DATA=1:@CHRM1.C00 00 12/10 21:45:32

**\*\* CALCULATION REPORT \*\***

CH	PKNO	TIME	AREA	HEIGHT	MK	IDNO	CONC	NAME
1	9	3.338	1253449	412308	V		34.5845	
	10	3.433	2351953	857343	SV		64.8938	
	25	7.686	4264	1462			0.1177	
	26	7.899	2718	946	SV		0.075	
	31	11.491	1205	130			0.0332	
	34	11.956	5605	340	V		0.1546	
	35	13.579	2435	430			0.0672	
	36	14.801	2682	199			0.074	
<hr/> TOTAL			3624310	1273157			100	

**Fig 20** The results of chiral GC analysis in temperature program condition:  
derivatized aminonitrile **56**.

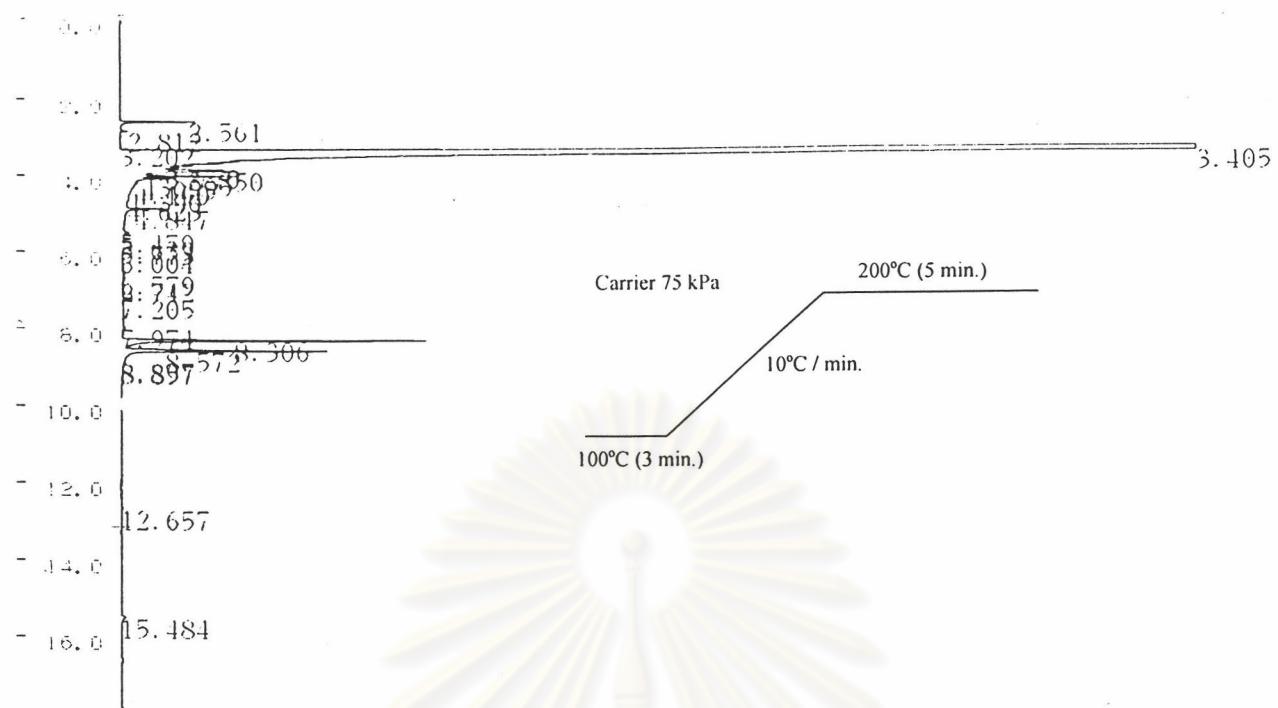


C-R8A CHROMATOPAC CH=1 Report No.=12 DATA=1:@CHRM1.COO 00/12/10 22:23:28

**\*\* CALCULATION REPORT \*\***

CH	PKNO	TIME	AREA	HEIGHT	MK	IDNO	CONC	NAME
1	5	3.227	2506	1052			0.0985	
	6	3.377	814748	370541	V		32.0326	
	7	3.428	1701160	751707	SV		66.8827	
18	4.804		1639	665			0.0644	
21	7.71		9282	3462			0.3649	
22	7.927		14163	5071	SV		0.5568	
<hr/> TOTAL			2543497	1132497			100	

**Fig 21** The results of chiral GC analysis in temperature program condition:  
derivatized aminonitrile **43**.

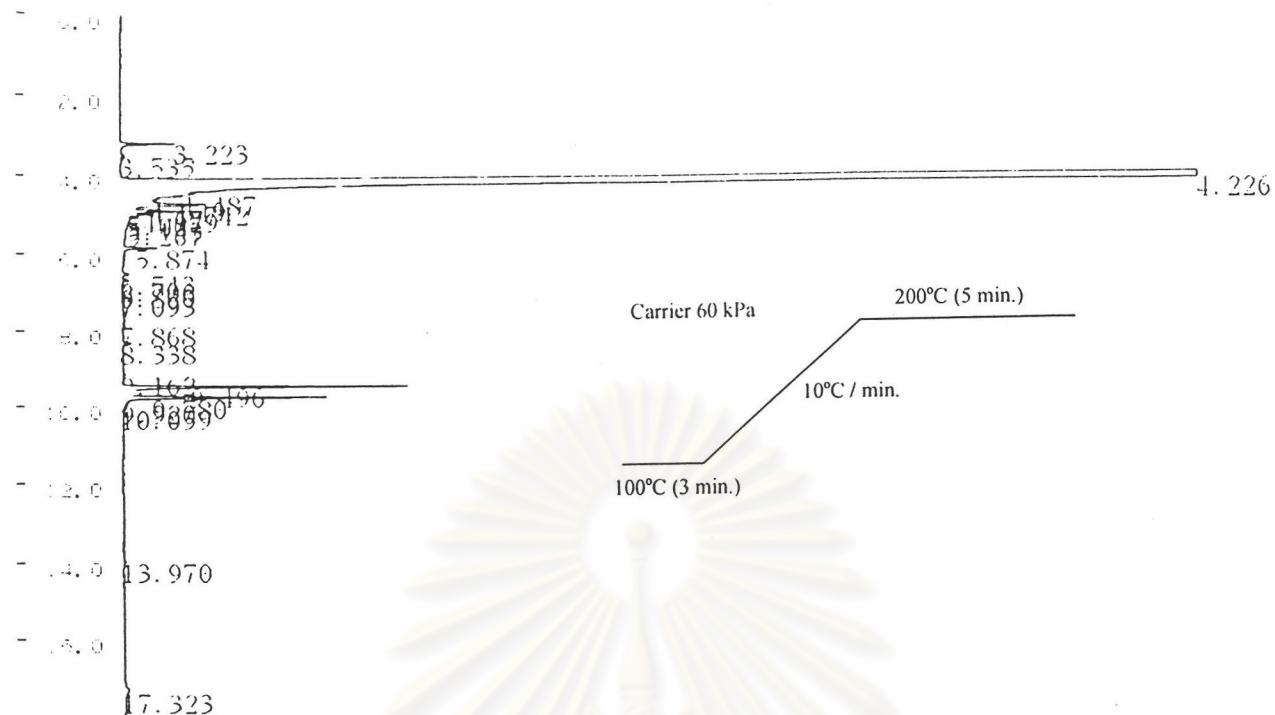


C-R8A CHROMATOPAC CH=1 Report No.=3 DATA=1:@CHRM1.C00 00.12/20 16:47:42

**\*\* CALCULATION REPORT \*\***

CH	PKNO	TIME	AREA	HEIGHT	MK	IDNO	CONC	NAME
1	4	3.405	3141230	936671	SV		99.8189	
	21	8.306	3216	1125			0.1022	
	22	8.572	2482	762	SV		0.0789	
TOTAL			3146929	938558			100	

**Fig 22-Fig 30** The results of GC analysis in various temperature program conditions: derivatized aminonitrile 25a.

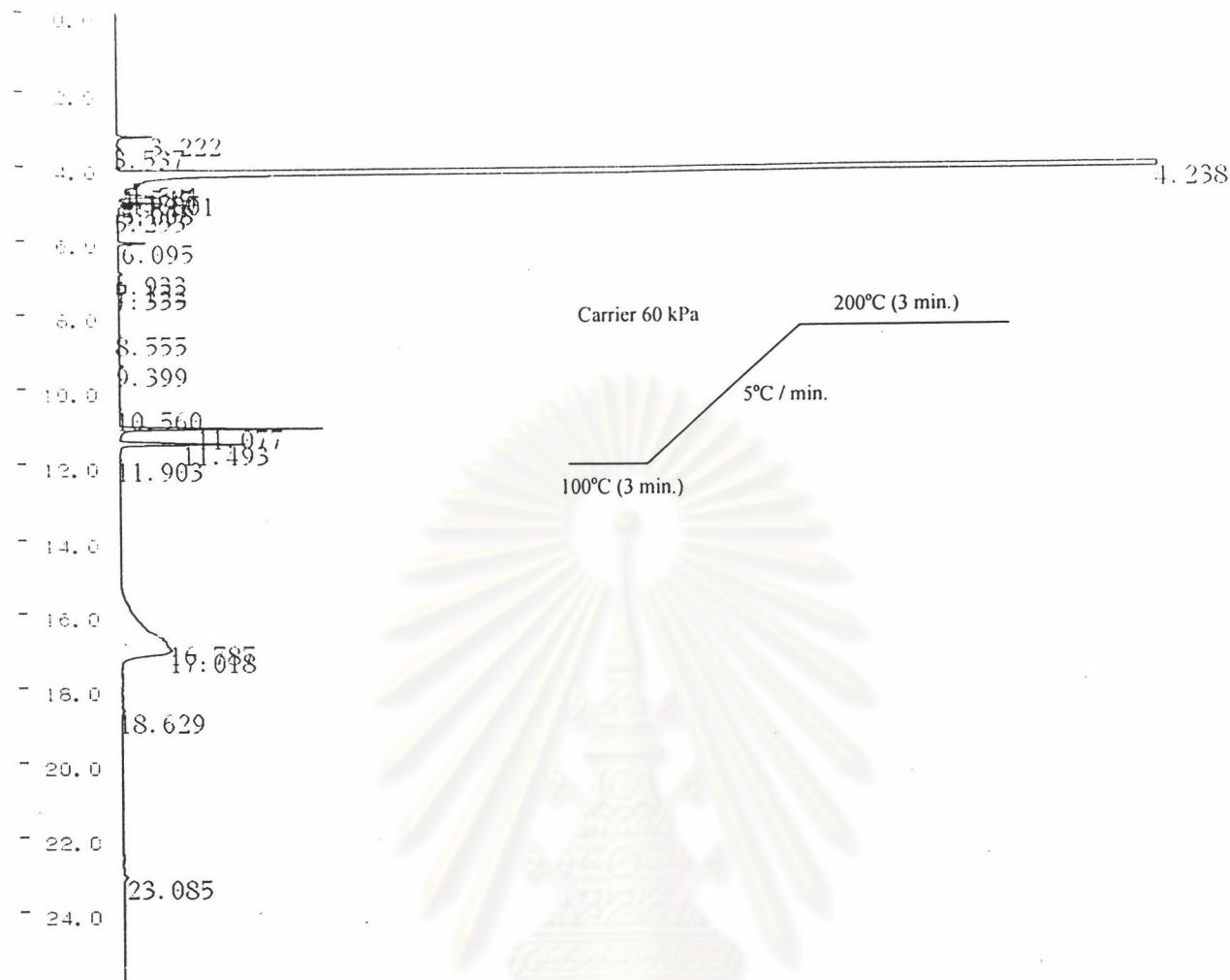


C-RSA CHROMATOPAC CH=1 Report No.=4 DATA=1:@CHRM1.C00 00.12.20 17:10:54

**\*\* CALCULATION REPORT \*\***

CH	PKNO	TIME	AREA	HEIGHT	MK	IDNO	CONC	NAME
1	3	4.226	3234867	725457	S		99.8272	
	19	9.496	3206	1055			0.0989	
	20	9.78	2393	743	V		0.0738	
TOTAL			3240465	727255			100	

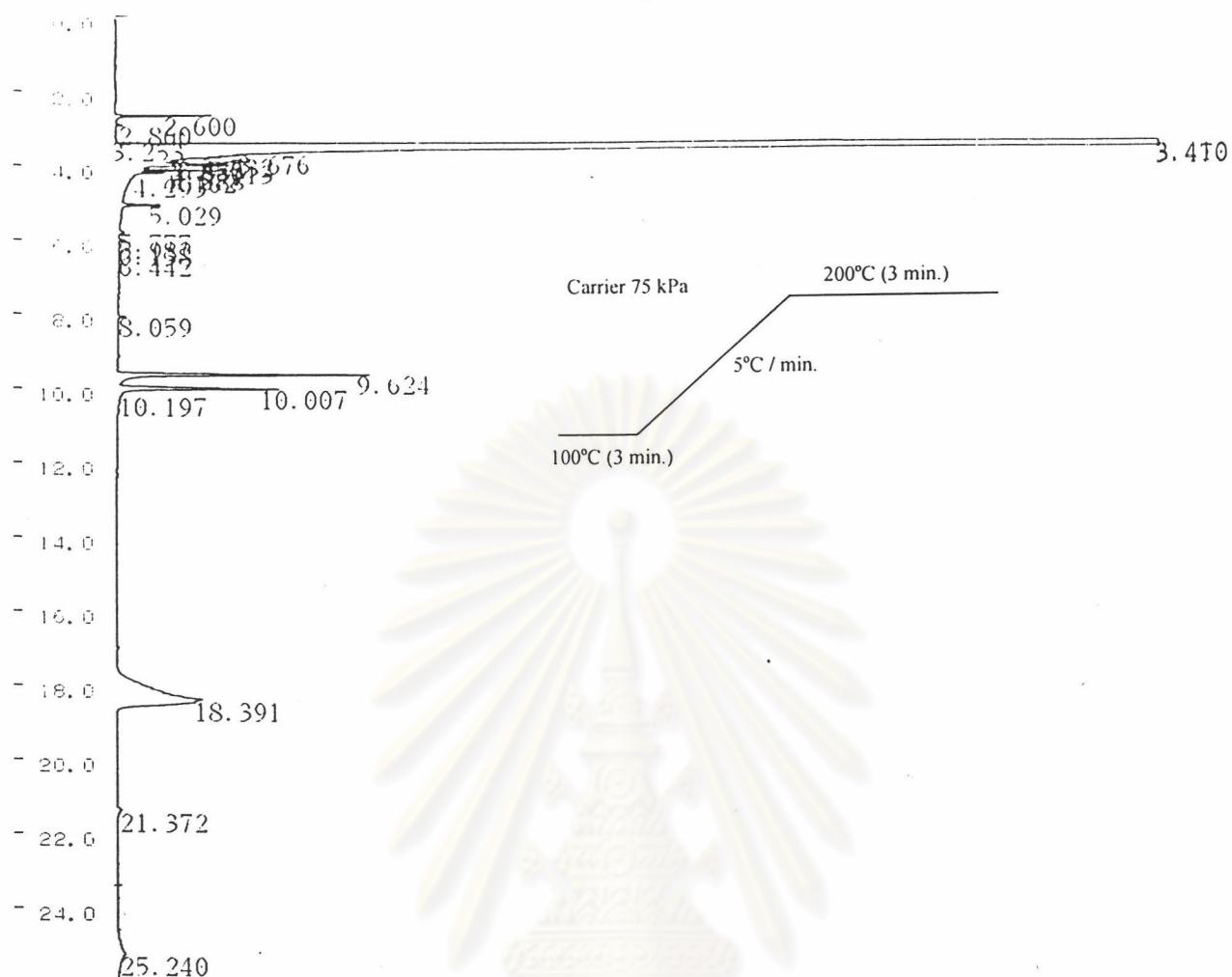
Fig 23



C-RSA CHROMATOPAC CH=1 Report No.=6 DATA=1:@CHRM1.C00 00 12 20 17:35:46

** CALCULATION REPORT **							
CH	PKNO	TIME	AREA	HEIGHT	MK	IDNO	NAME
1	3	4.238	2074014	527847	S		99.3682
	20	11.077	2664	790			0.1277
	21	11.493	1898	480			0.0909
	23	16.787	4828	170			0.2313
	24	17.018	3796	191	V		0.1819
TOTAL			2087200	529477			100

Fig 24

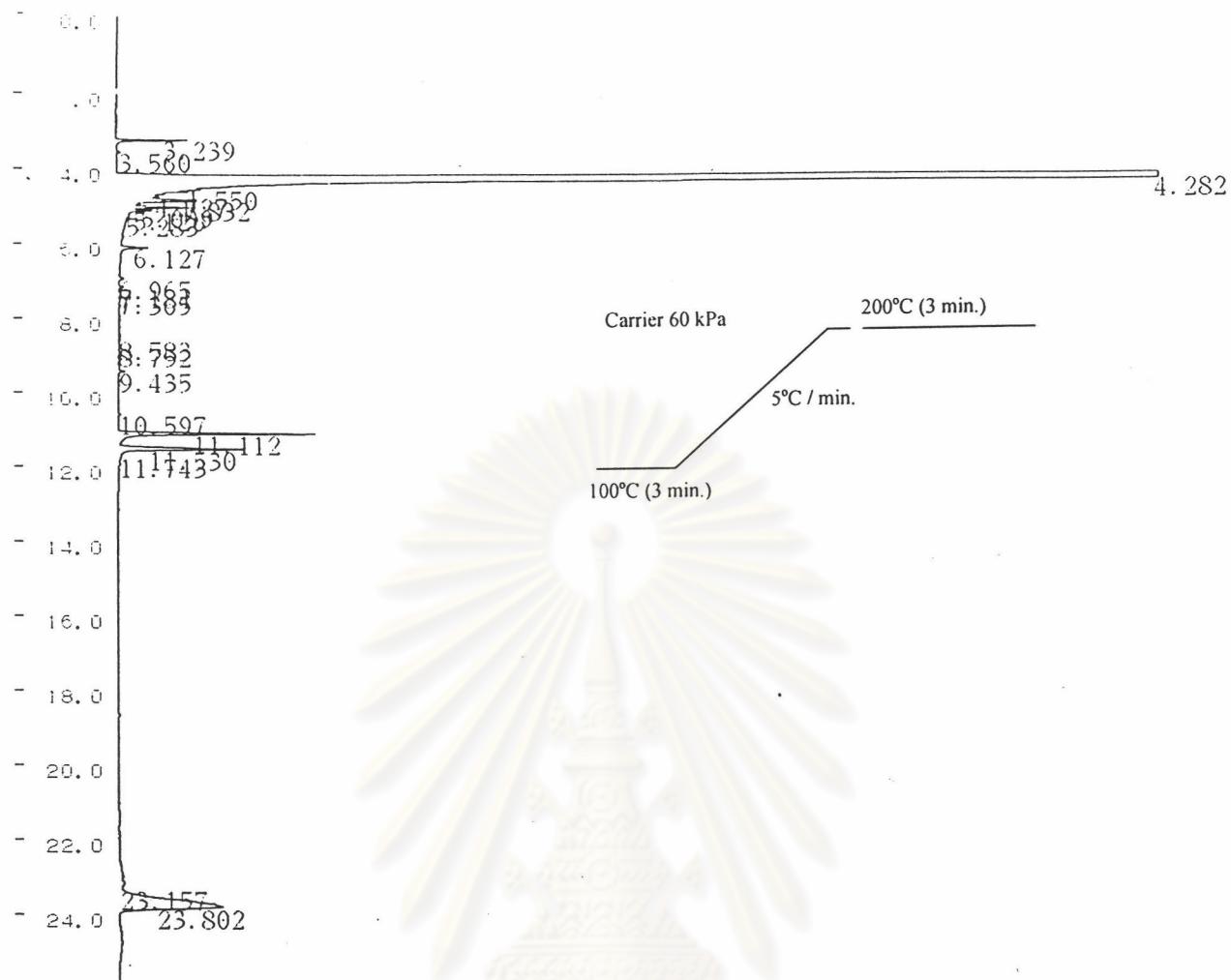


C-R8A CHROMATOPAC CH=1 Report No.=7 DATA=1:@CHRM1.C00 00/12/20 18:07:38

\*\* CALCULATION REPORT \*\*

CH	PKNO	TIME	AREA	HEIGHT	MK	IDNO	CONC	NAME
1	4	3.41	1136864	494161			30.7468	
	5	3.47	2545999	934270	SV		68.8573	
	22	9.624	3656	970			0.0989	
	23	10.007	2631	619	SV		0.0712	
	25	18.391	8349	328			0.2258	
<hr/>								
TOTAL			3697499	1430348			100	

Fig 25

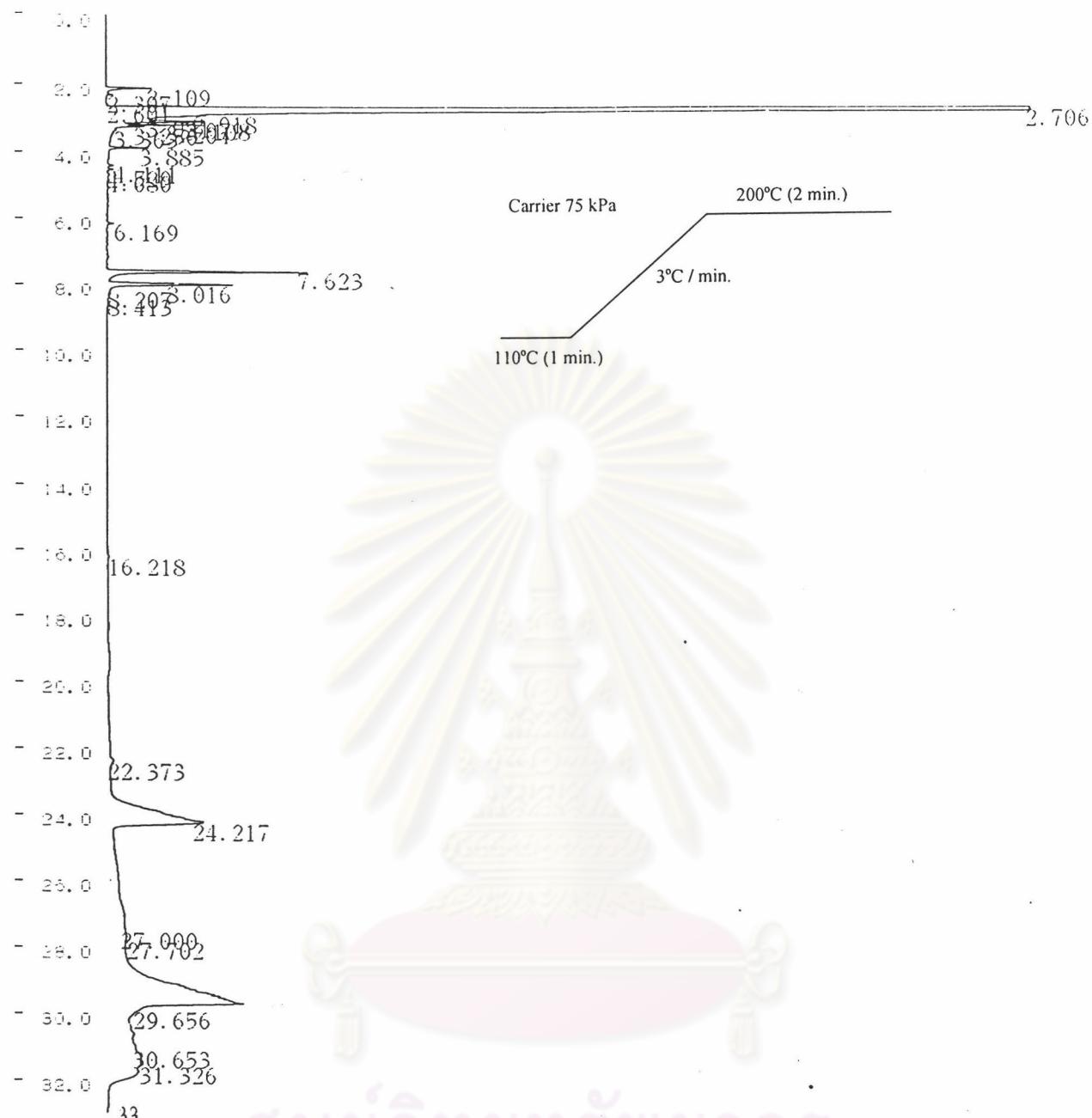


C-R8A CHROMATOPAC CH=1 Report No.=8 DATA=1:@CHRM1:C00 00/12/20 18:42:12

**\*\* CALCULATION REPORT \*\***

CH	PKNO	TIME	AREA	HEIGHT	MK	IDNO	CONC	NAME
1	3	4.282	3180654	735827	S		99.6335	
	18	11.112	3013	759			0.0944	
	19	11.53	2137	478	V		0.0669	
	22	23.802	6549	395	V		0.2051	
TOTAL			3192352	737460			100	

Fig 26

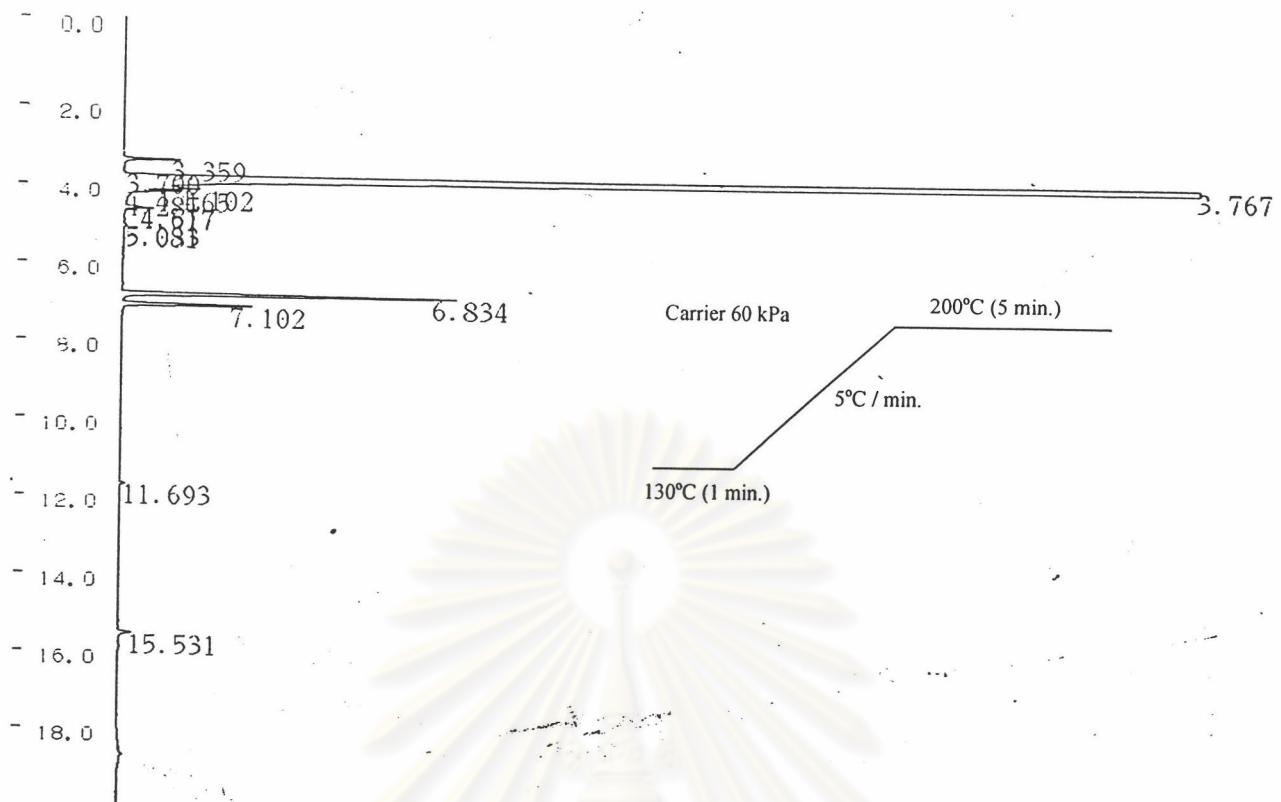


C-R8A CHROMATOPAC CH=1 Report No.=9 DATA=1:@CHRM1.C02 00/12/20 19:22:32

\*\* CALCULATION REPORT \*\*

CH	PKNO	TIME	AREA	HEIGHT	MK	IDNO	CONC	NAME
1	4	2.706	2970941	1120562	SVE		98.2247	
	17	7.623		3451		874	0.1141	
	18	8.016		2528		548	0.0836	
	23	24.217		10068		395	0.3329	
	25	27.702		1259		39	0.0416	
	26	29.656		24834		540	0.821	
	27	30.653		3292		96	0.1088	
	28	31.326		8266		133	0.2733	
<hr/>								
TOTAL			3024639	1123186			100	

Fig 27

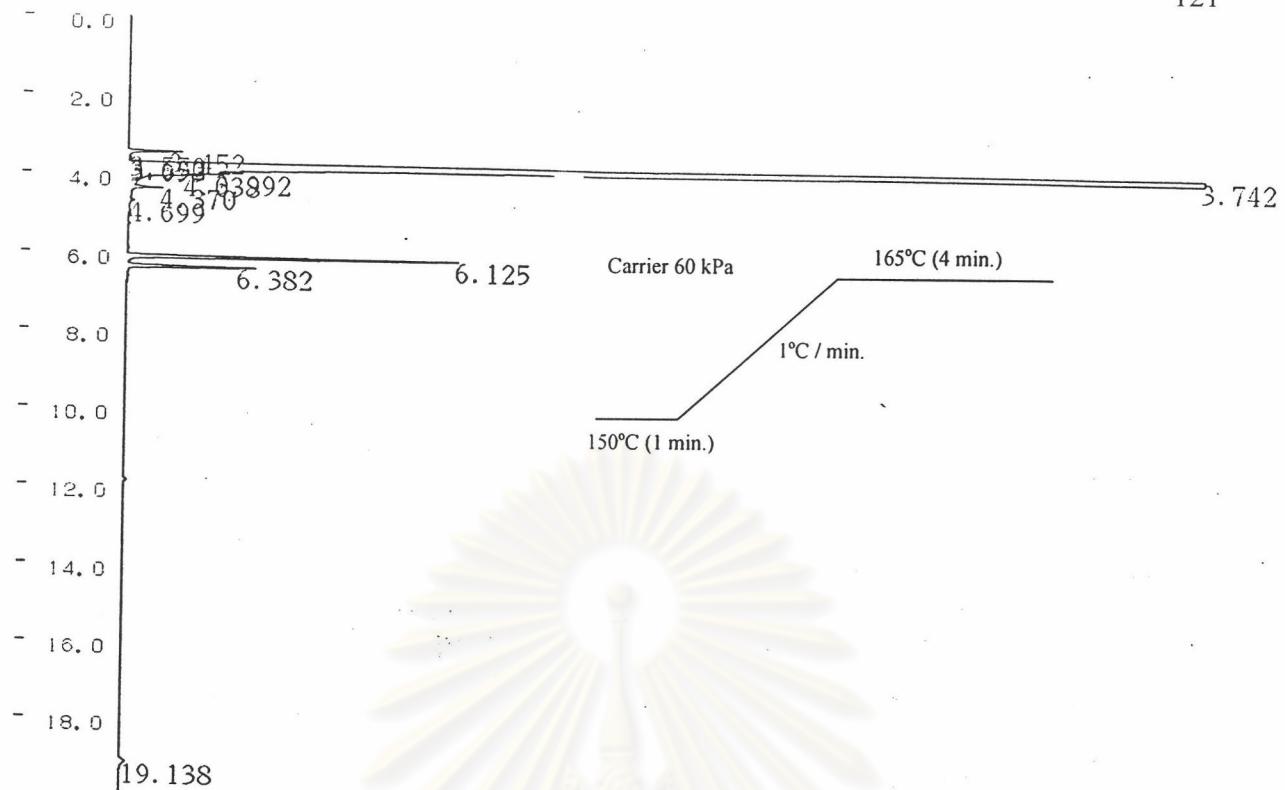


C-R8A CHROMATOPAC CH=1 Report No.=7 DATA=1:@CHRM1.C00 00/12/29 18:55:00

**\*\* CALCULATION REPORT \*\***

CH	PKNO	TIME	AREA	HEIGHT	MK	IDNO	CONC	NAME
1	3	3.767	2469220	735831	SV		99.7532	
	11	6.834	4338	1245			0.1752	
	12	7.102	1772	489	V		0.0716	
<b>TOTAL</b>			<b>2475330</b>	<b>737565</b>			<b>100</b>	

Fig 28

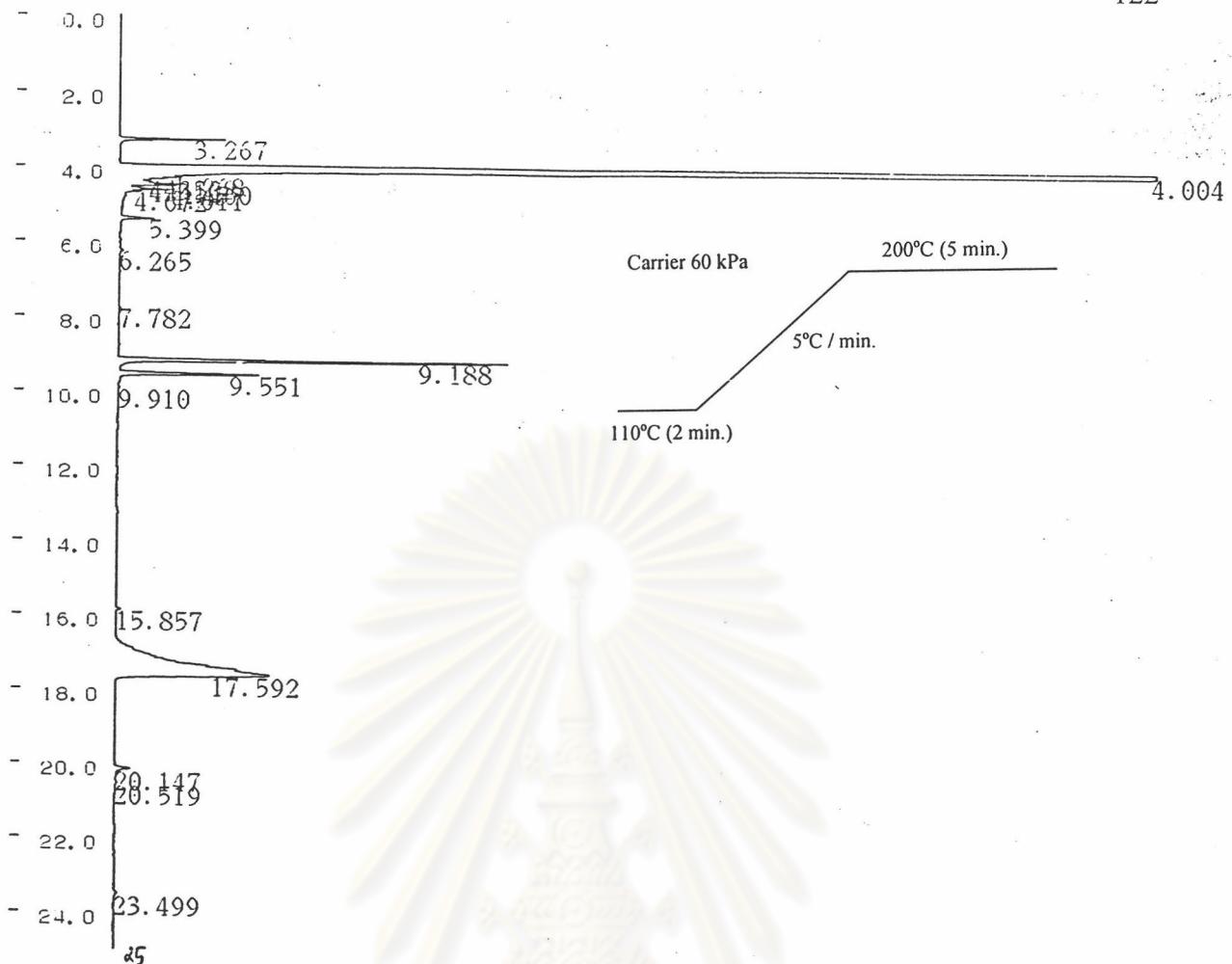


C-R8A CHROMATOPAC CH=1 Report No.=8 DATA=1:@CHRM1.C00 00/12/29 19:19:04

**\*\* CALCULATION REPORT \*\***

CH	PKNO	TIME	AREA	HEIGHT	MK	IDNO	CONC	NAME
1	5	3.742	2021352	644453	SV		99.7426	
	10	6.125	3684	1222			0.1818	
	11	6.382	1533	475	V		0.0756	
TOTAL			2026568	646150			100	

Fig 29



C-R8A CHROMATOPAC CH=1 Report No.=9

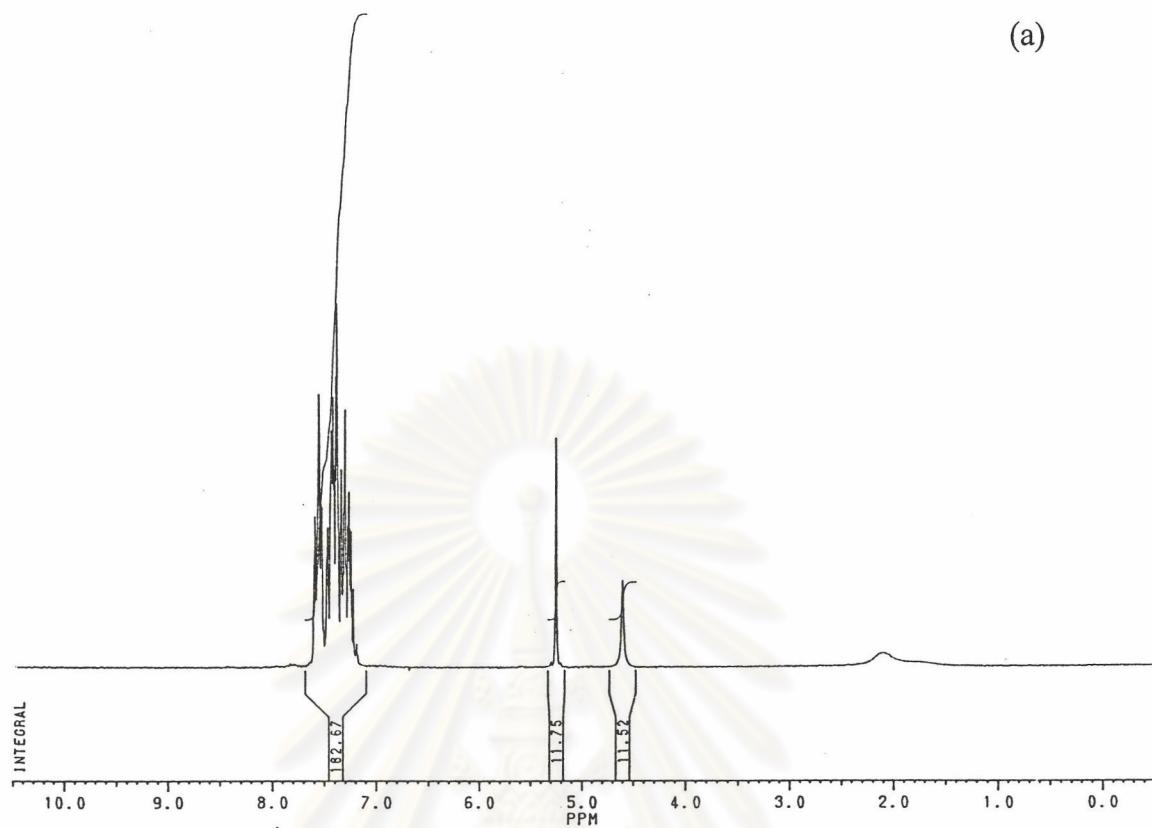
DATA=1:@CHRM1.C00 00/12/29 19:44:22

## \*\* CALCULATION REPORT \*\*

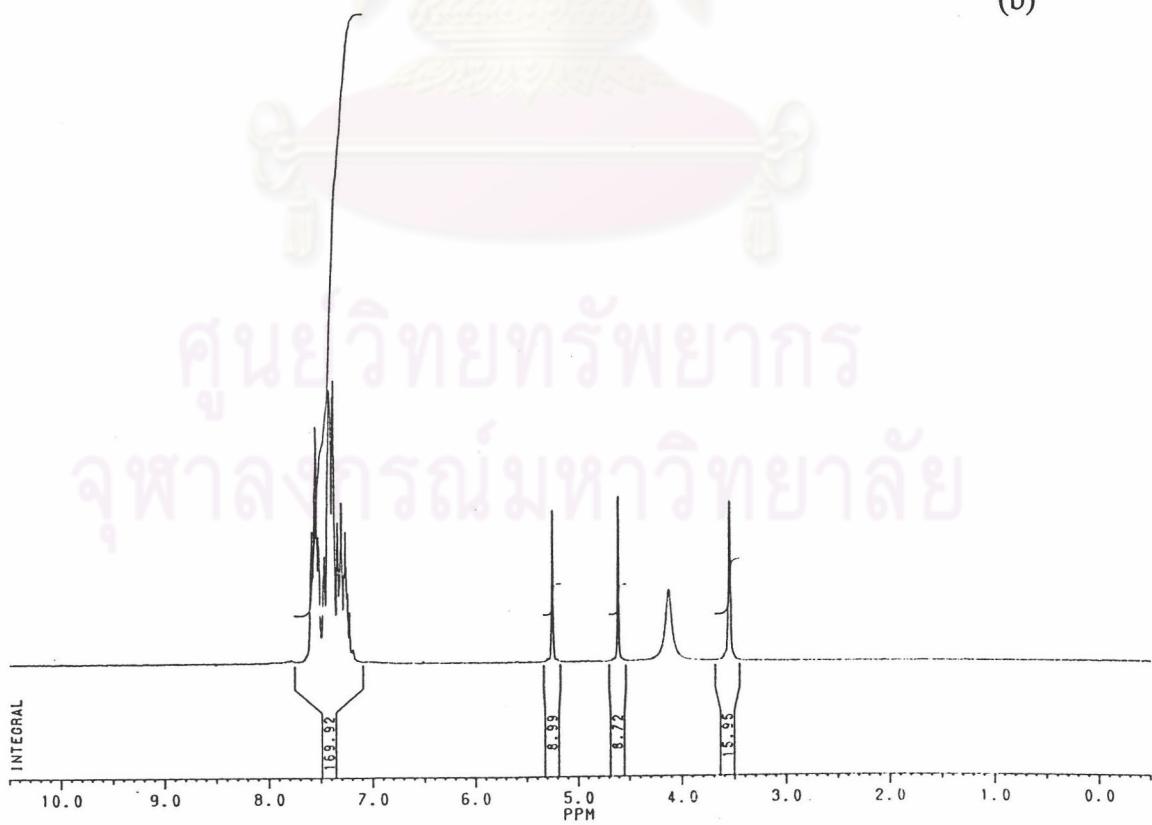
CH	PKNO	TIME	AREA	HEIGHT	MK	IDNO	CONC	NAME
1	2	4.004	2699914	992054	S E		99.2026	
	14	9.188	5240	1489			0.1925	
	15	9.551	2107	538	V		0.0774	
	18	17.592	14355	593			0.5274	
TOTAL			2721616	994675			100	

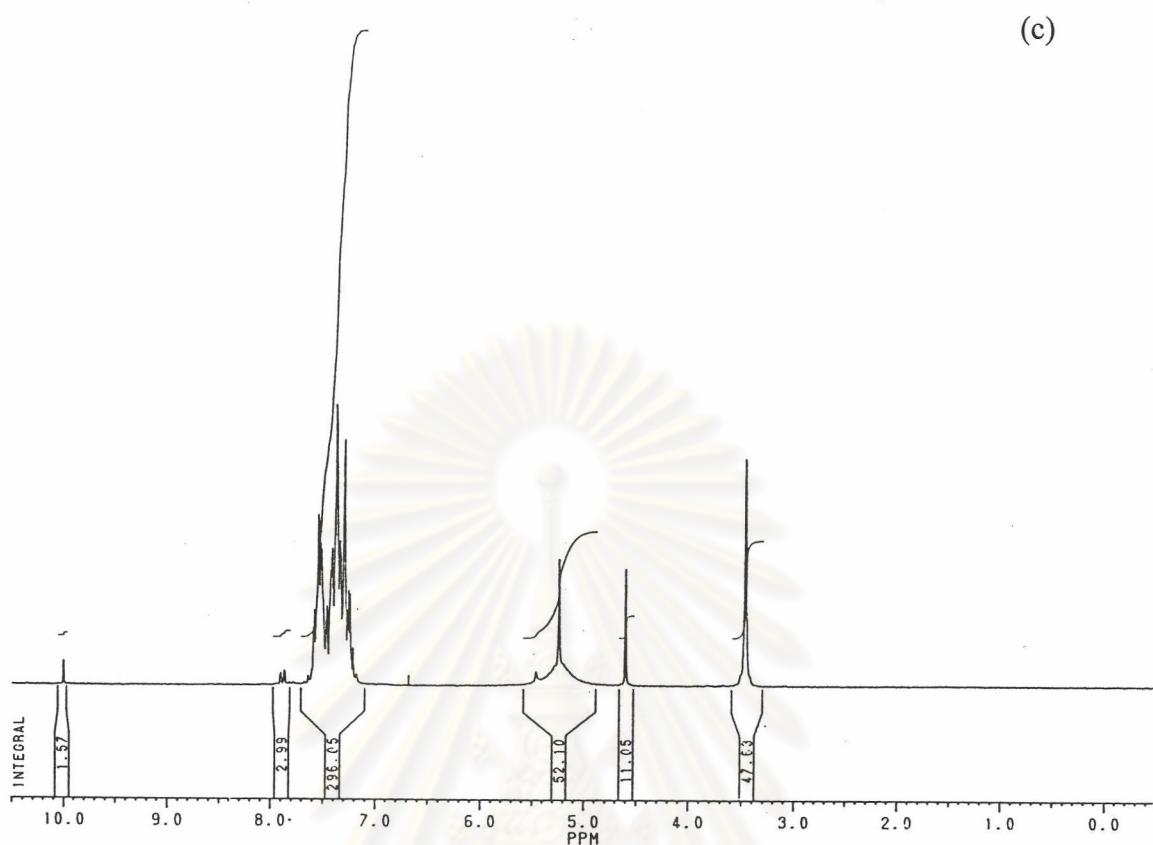
Fig 30

(a)

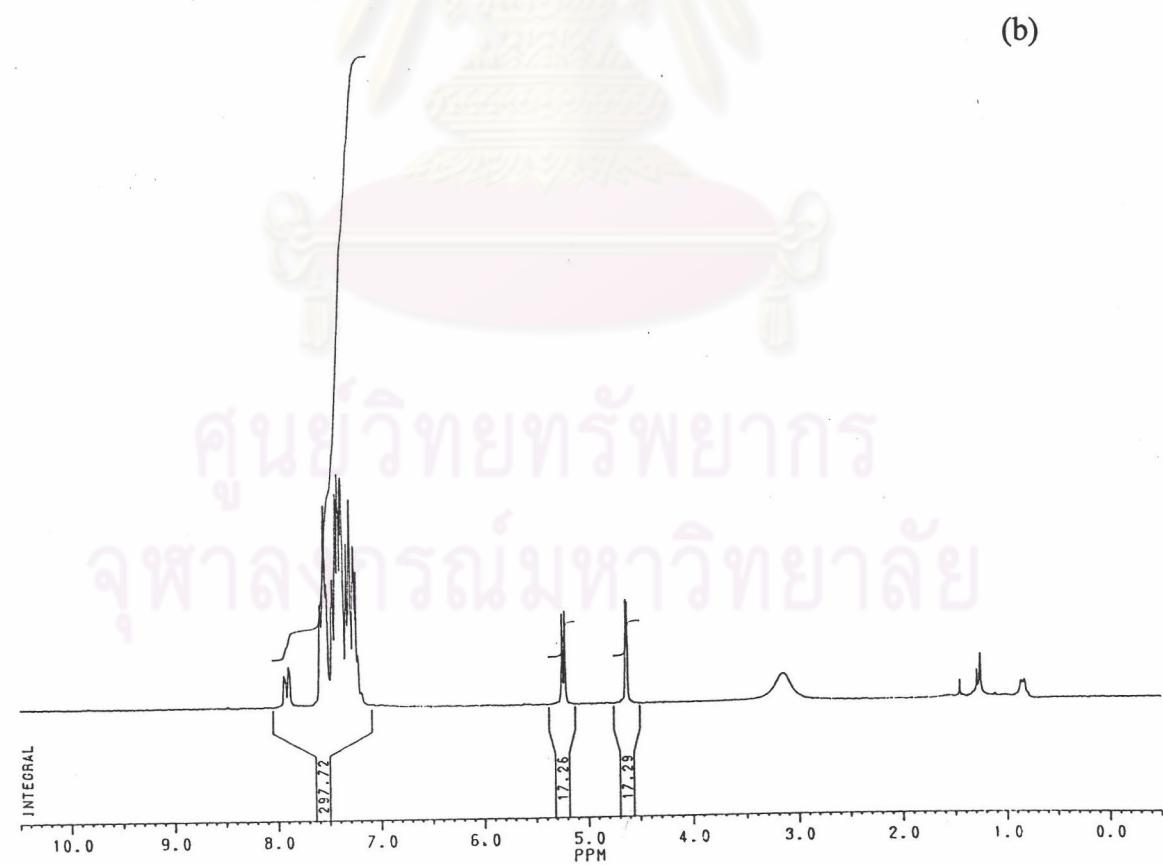
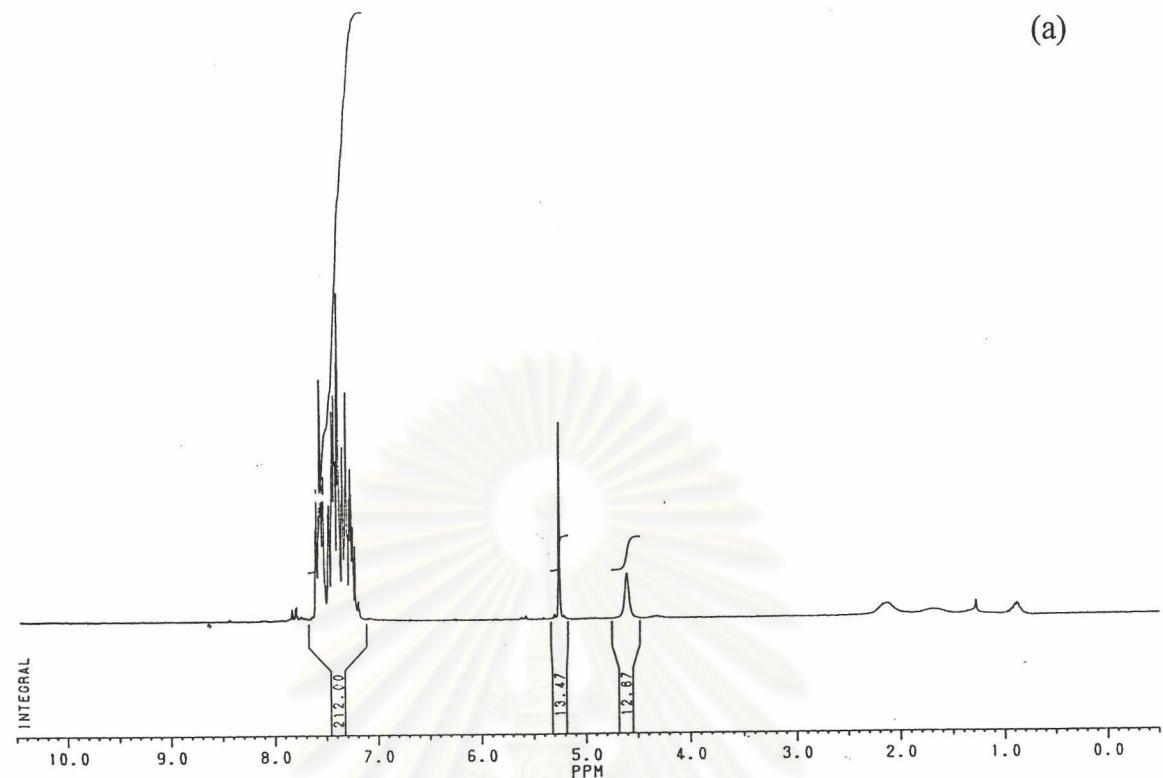


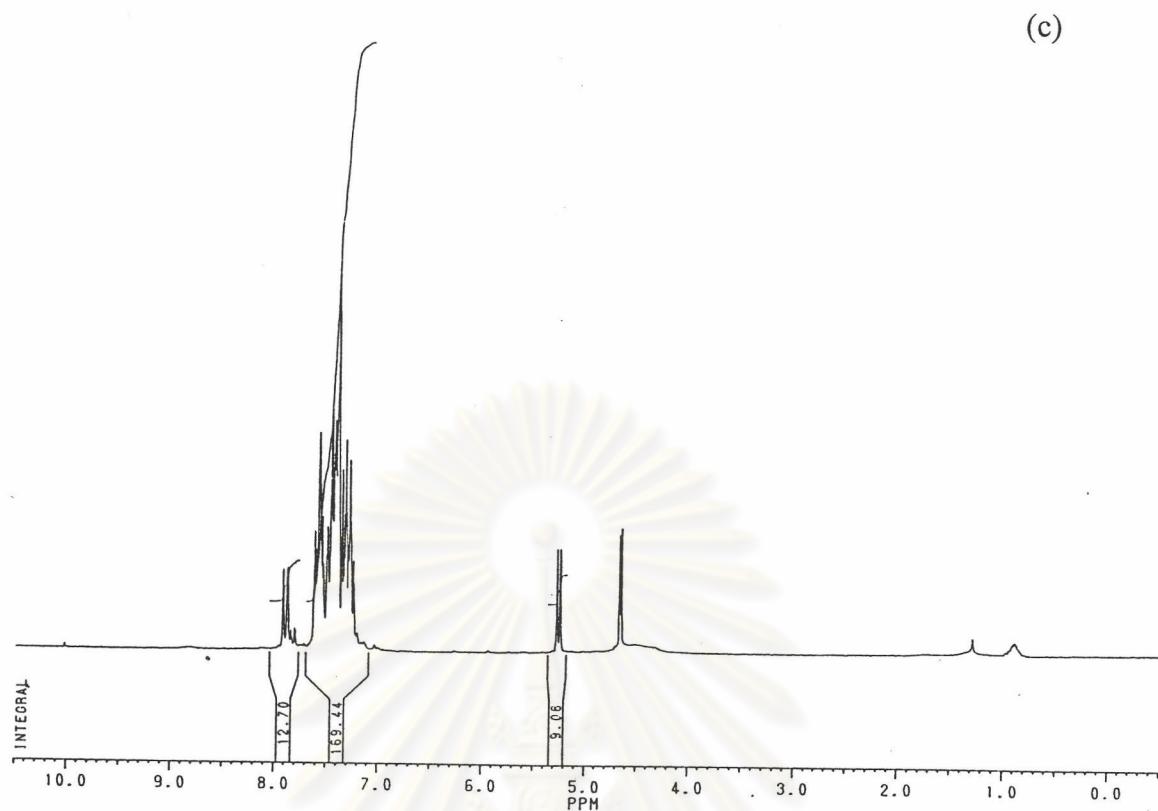
(b)



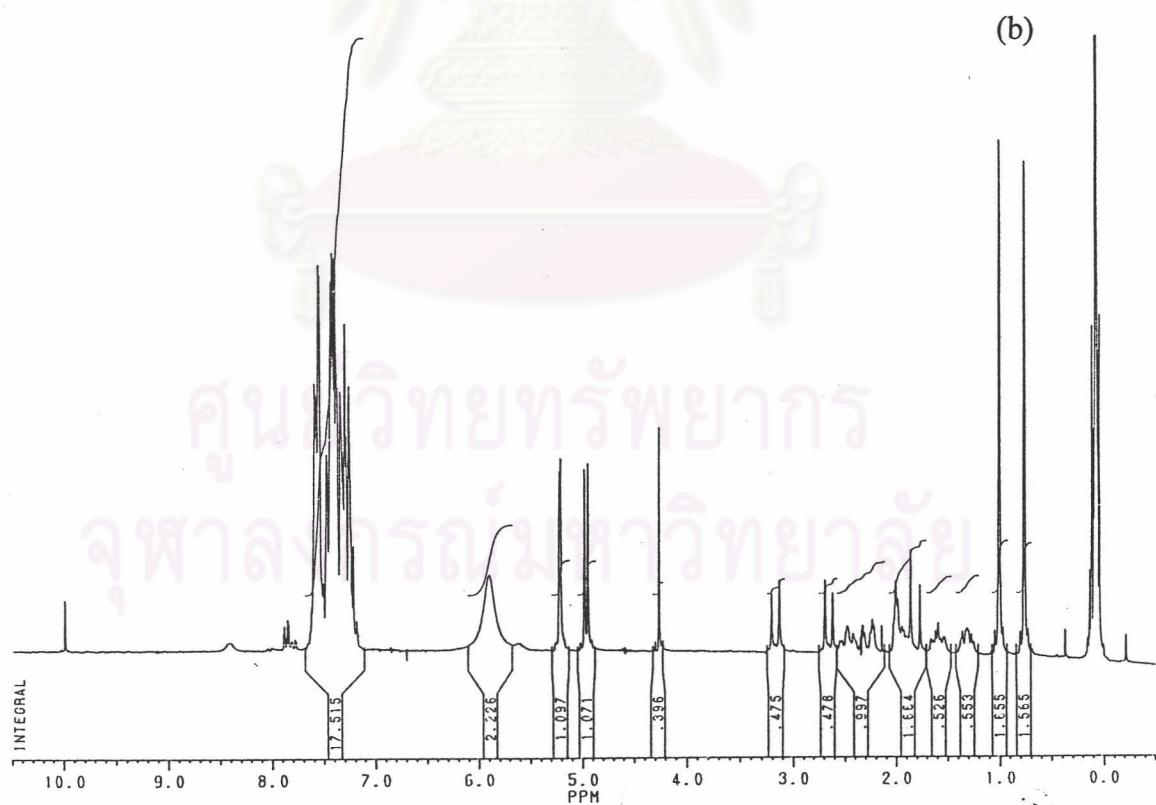
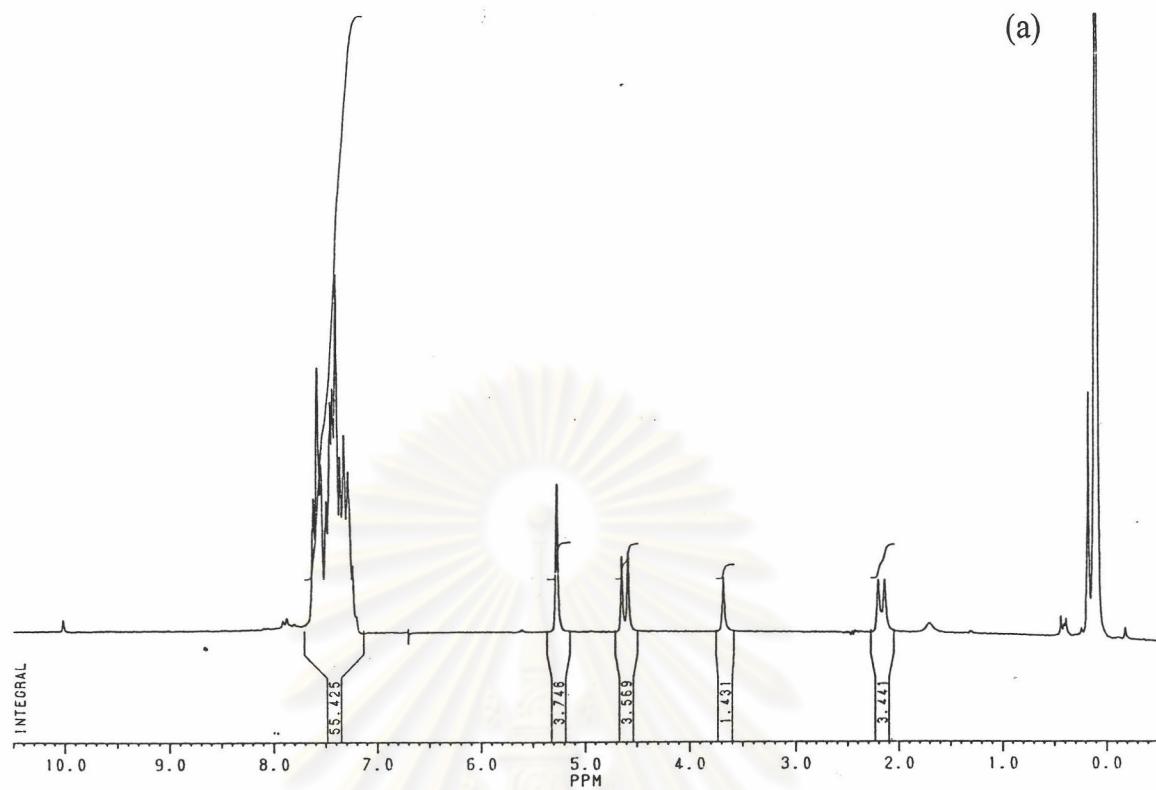


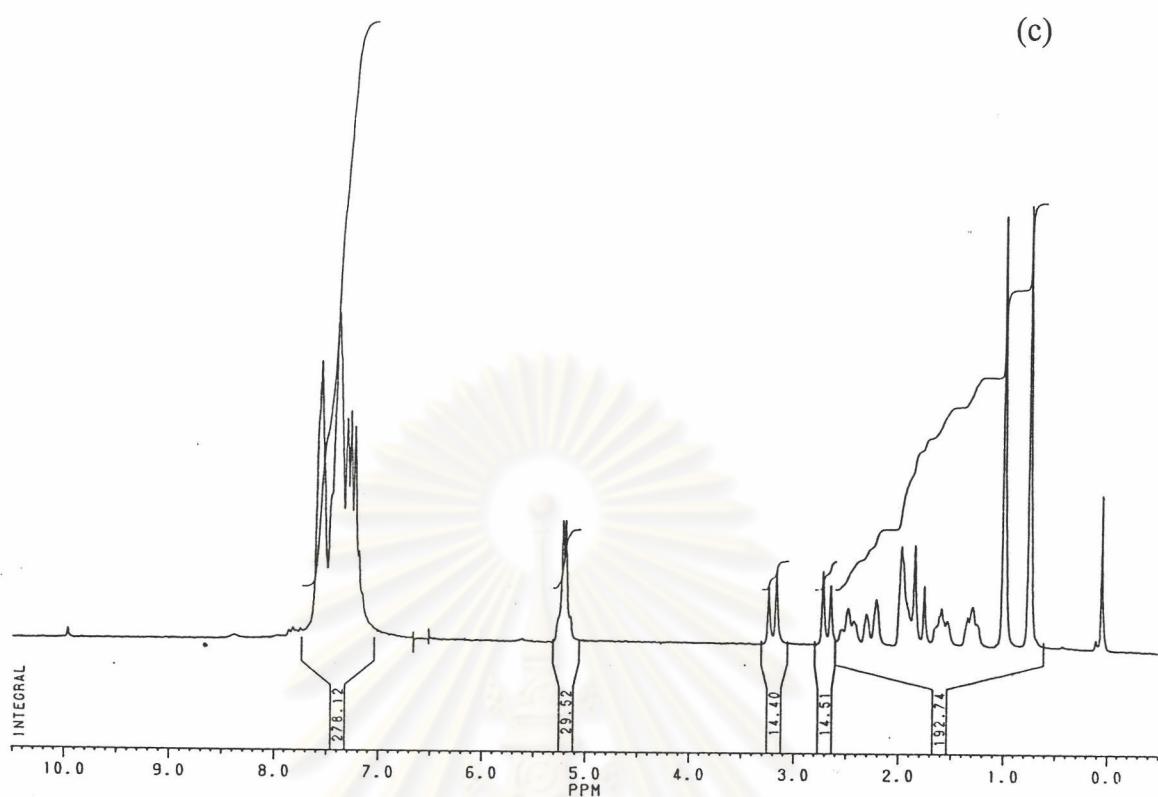
**Fig 31** The NMR spectra of racemic **25a**: (a) Racemic **25a** before adding **58**; (b) After adding 0.5 eq of **58**; (c) After adding 1.0 eq of **58**.





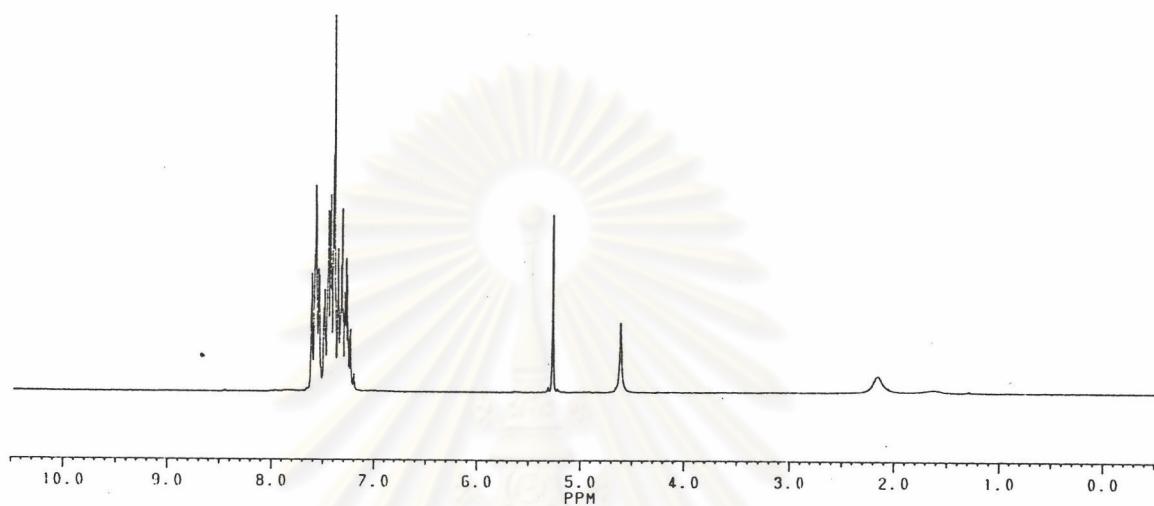
**Fig 32** The NMR spectra of racemic **25a**: (a) Racemic **25a** before adding **59**; (b) After adding 0.25 eq of **59**; (c) After adding 0.5 eq of **59**.



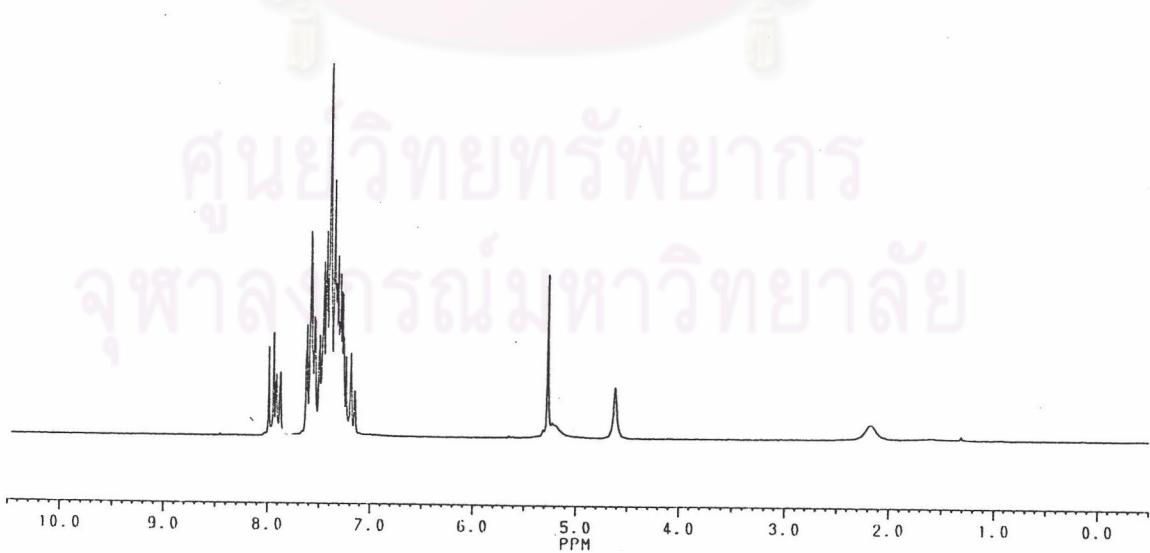


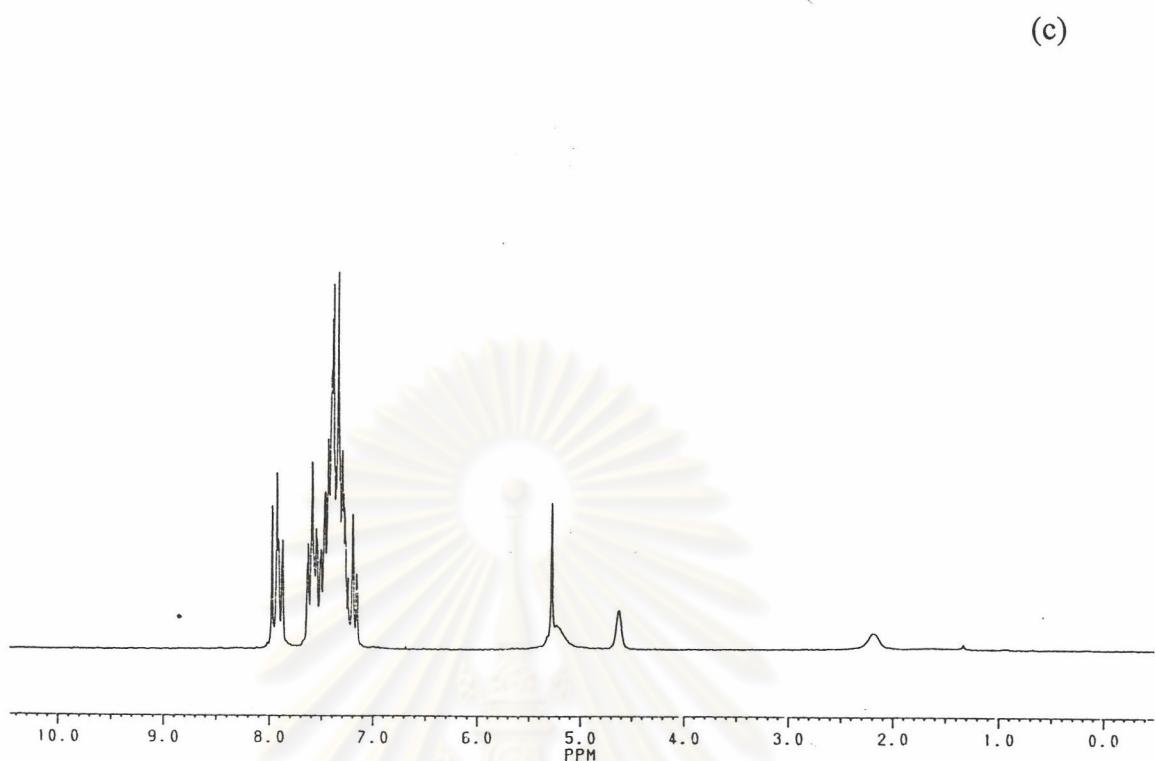
**Fig 33** The NMR spectra of racemic **25a**: (a) Racemic **25a** before adding **60**; (b) After adding 0.5 eq of **60**; (c) After adding 1.0 eq of **60**.

(a)



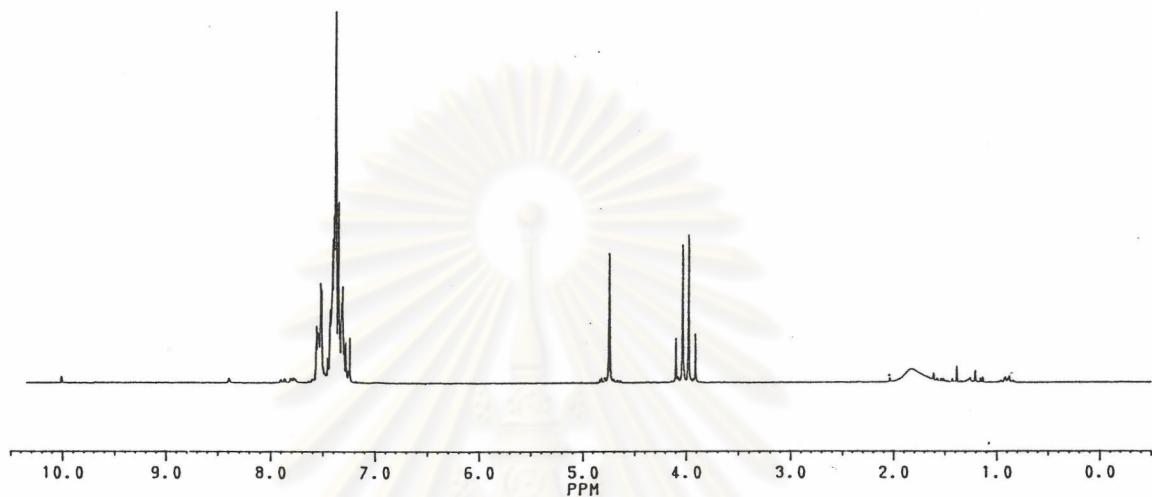
(b)



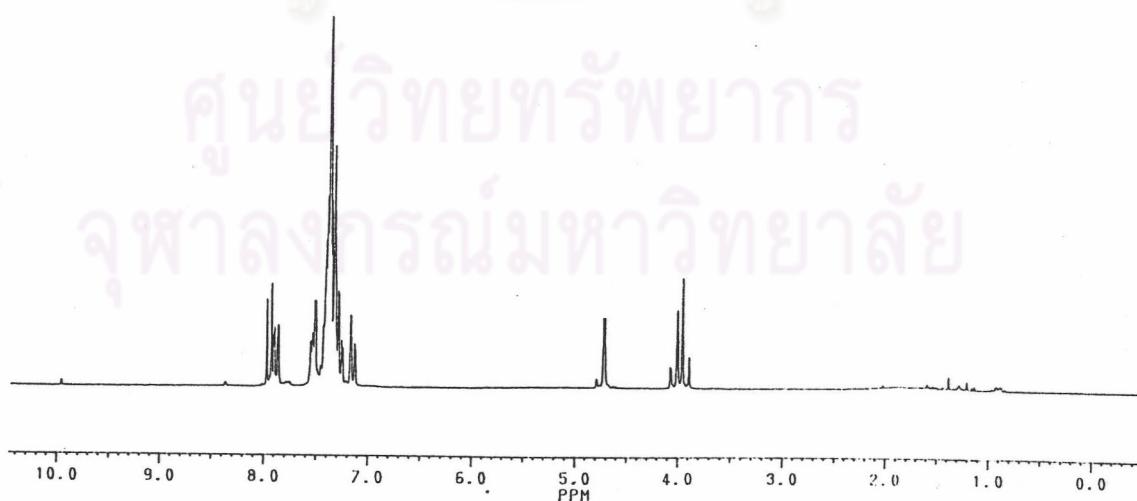


**Fig 34** The NMR spectra of racemic **25a**: (a) Racemic **25a** before adding **61**; (b) After adding 0.5 eq of **61**; (c) After adding 1.0 eq of **61**.

(a)

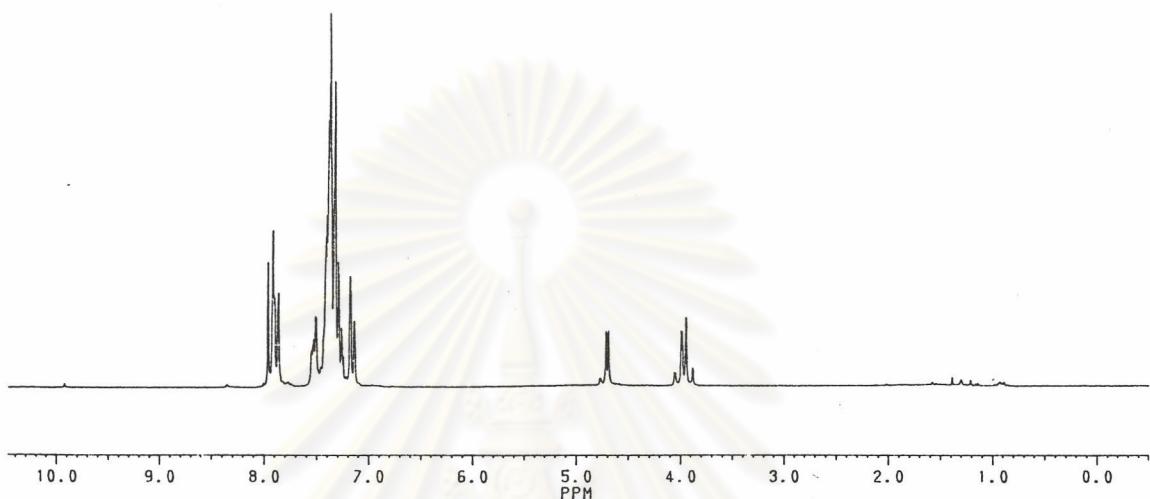


(b)

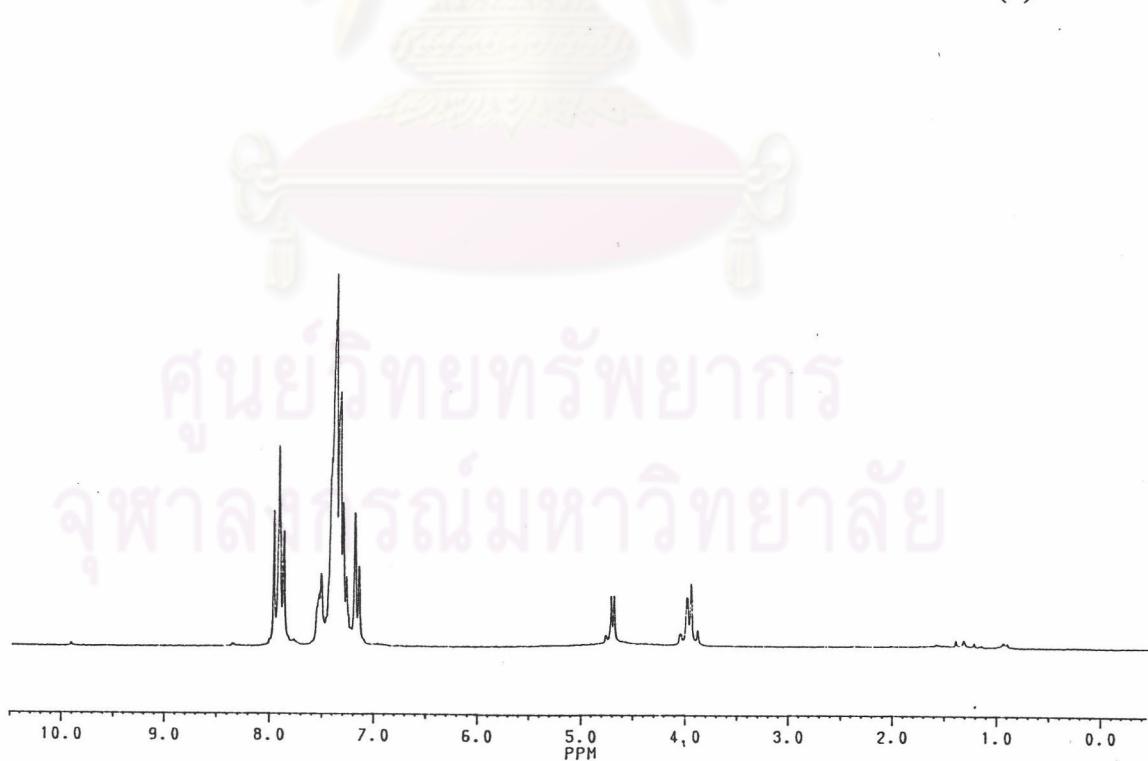


**Fig 35** The NMR spectra of racemic 43: (a) Racemic 43 before adding 61; (b) After adding 0.5 eq of 61.

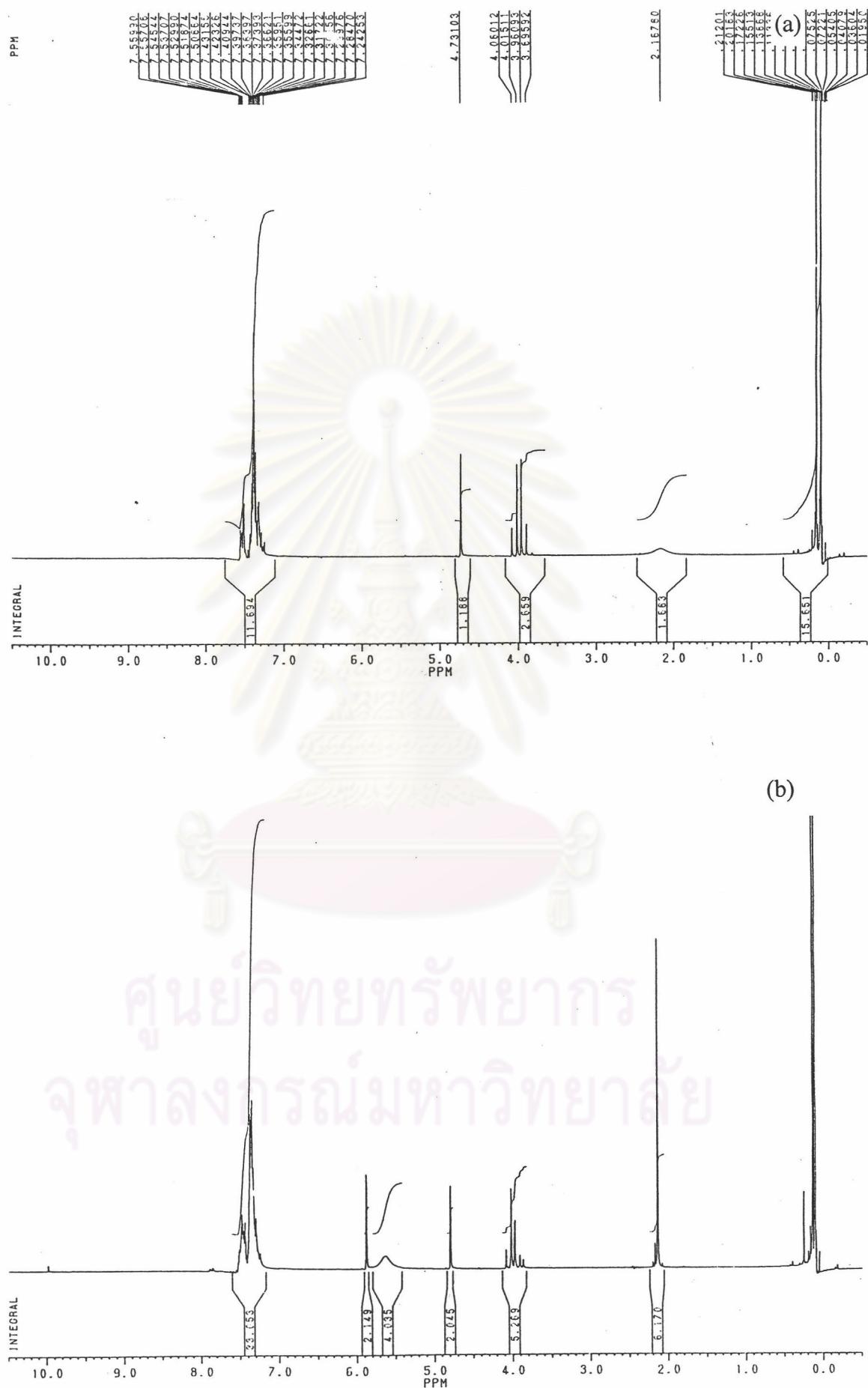
(a)



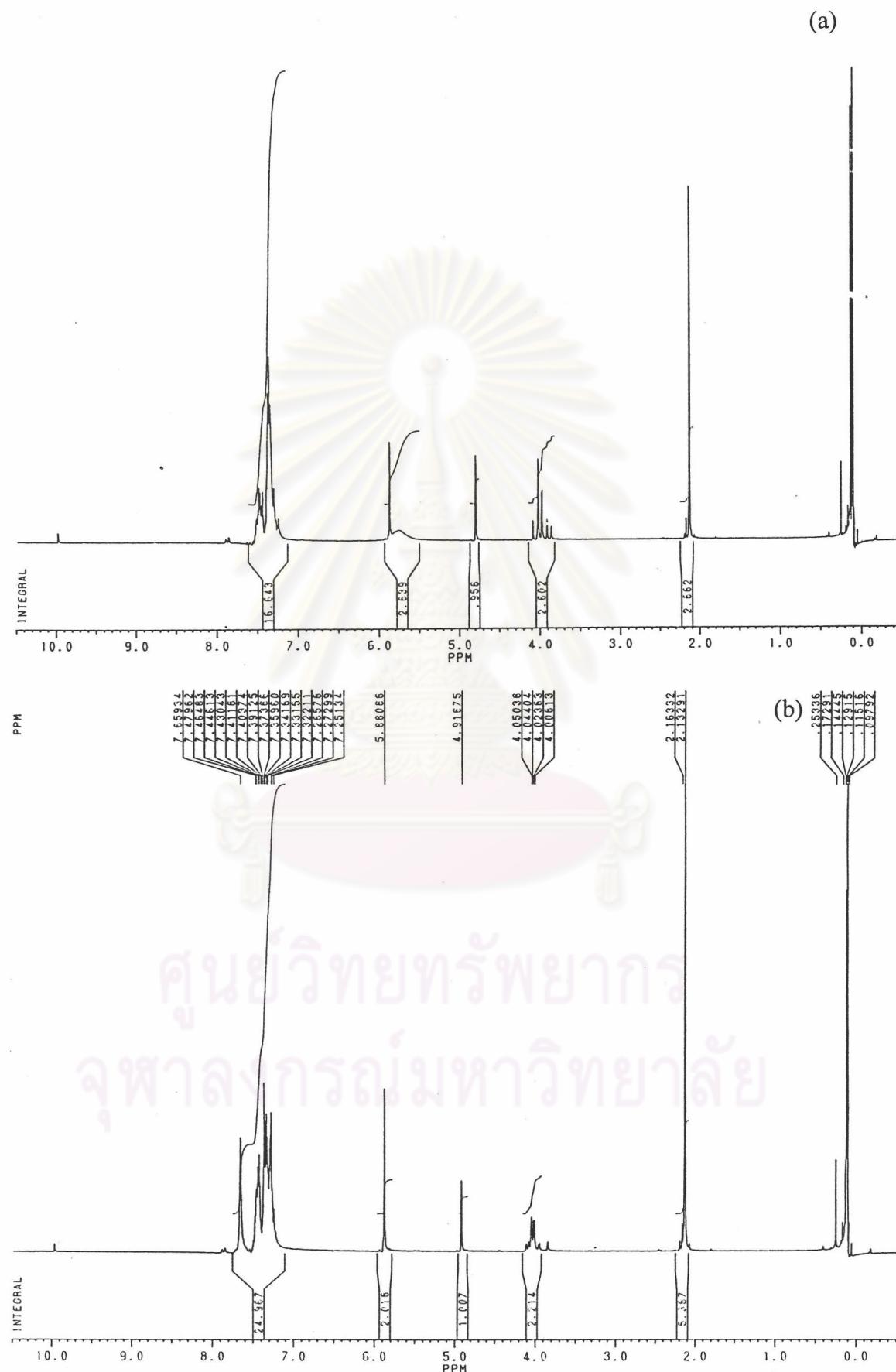
(b)



**Fig 36** The NMR spectra of racemic **43**: (a) After adding 1.0 eq of **61**; (b) After adding 1.5 eq of **61**.



**Fig 37** The NMR spectra of racemic 43: (a) Racemic 43 before adding 62; (b) After adding 1.0 eq of 62.



**Fig 38** The NMR spectra of racemic **43**: (a) After adding 1.0 eq of **62**; (b) After adding 2.0 eq of **62**.

## CURRICULUM VITAE

Miss Woraluk Mansawat was born on June 14<sup>th</sup>, 1976 in Phitsanulok, Thailand. She received a Bachelor Degree of Science, majoring in Chemistry from Chulalongkorn University in 1998. Since 1998, she has been a graduate student studying Organic Chemistry as her major course at Chulalongkorn University. During her studies towards the Master's degree, she was awarded a teaching assistant scholarship by the Faculty of Science during 1998-2001 and was supported by a research grant for her Master degree's thesis from the Graduate School, Chulalongkorn University.

Her present address is 231 Moo 5, Sanamchai District, Muang, Suphanburi, Thailand 72000, Tel. (035)-546335.

ศูนย์วิทยทรัพยากร  
จุฬาลงกรณ์มหาวิทยาลัย