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APPENDIX



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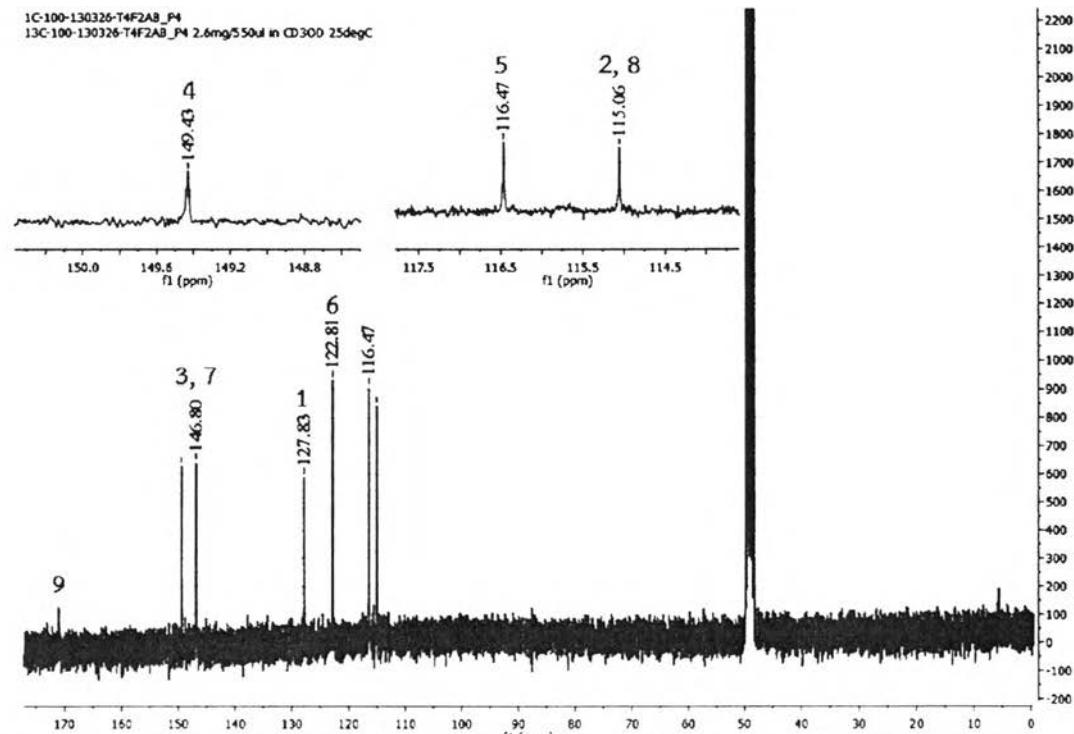
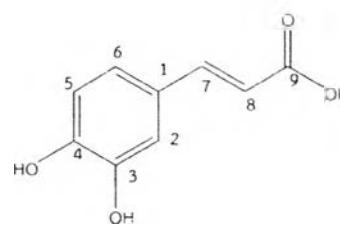


Figure A1 ¹³C NMR (75 MHz) Spectrum of compound T4F2AB (in CD₃OD)

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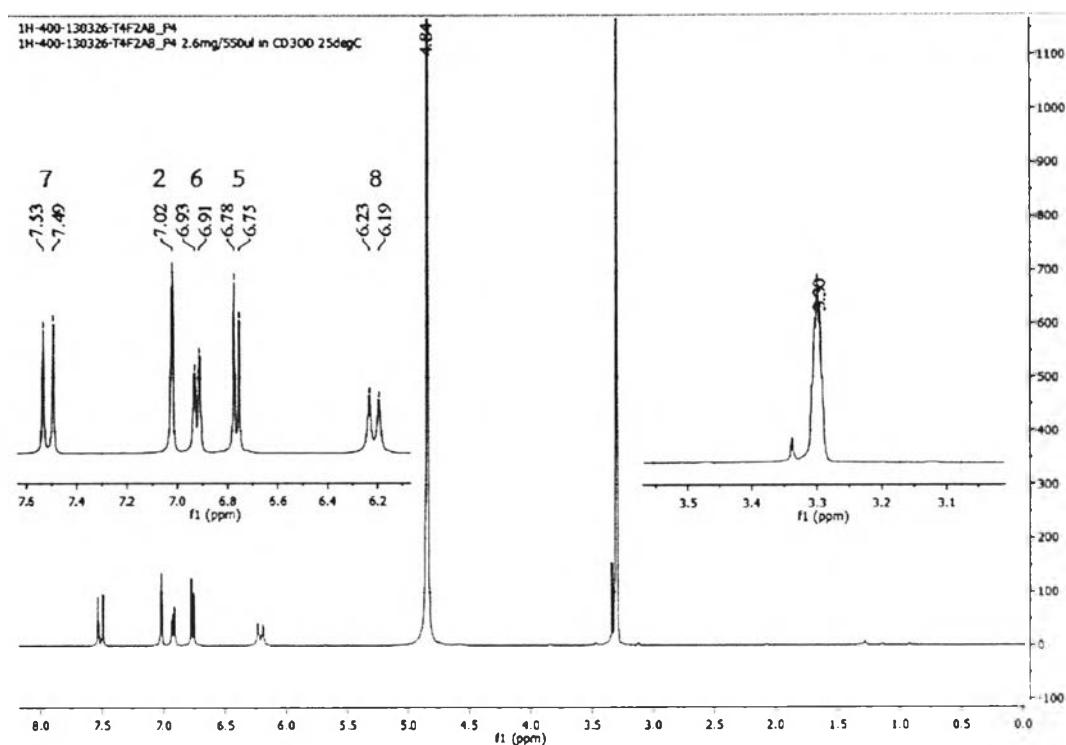
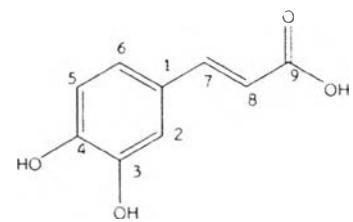


Figure A2 ^1H NMR (300 MHz) Spectrum of compound T4F2AB (in CD_3OD)

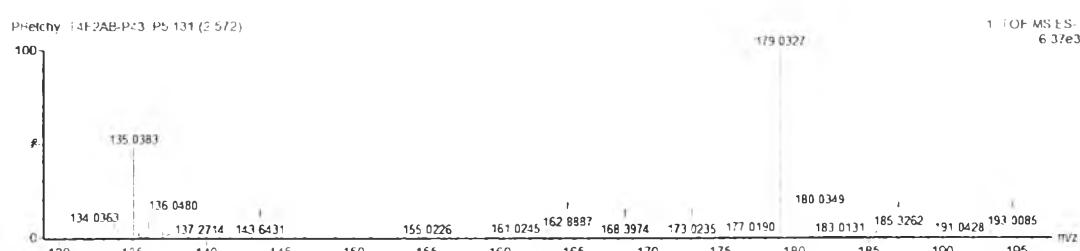
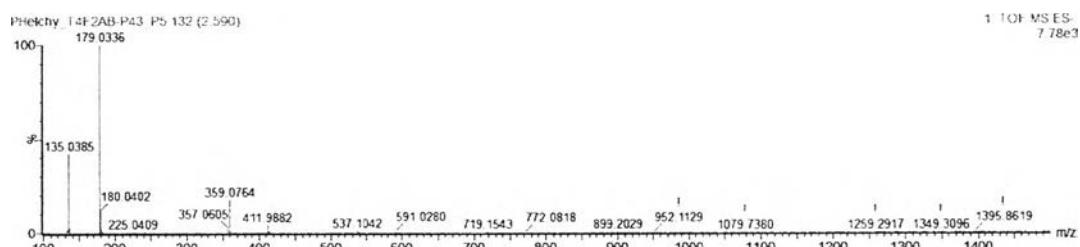
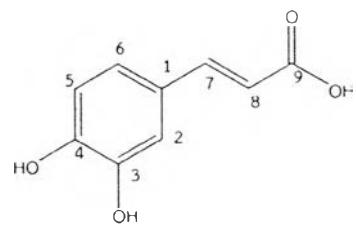
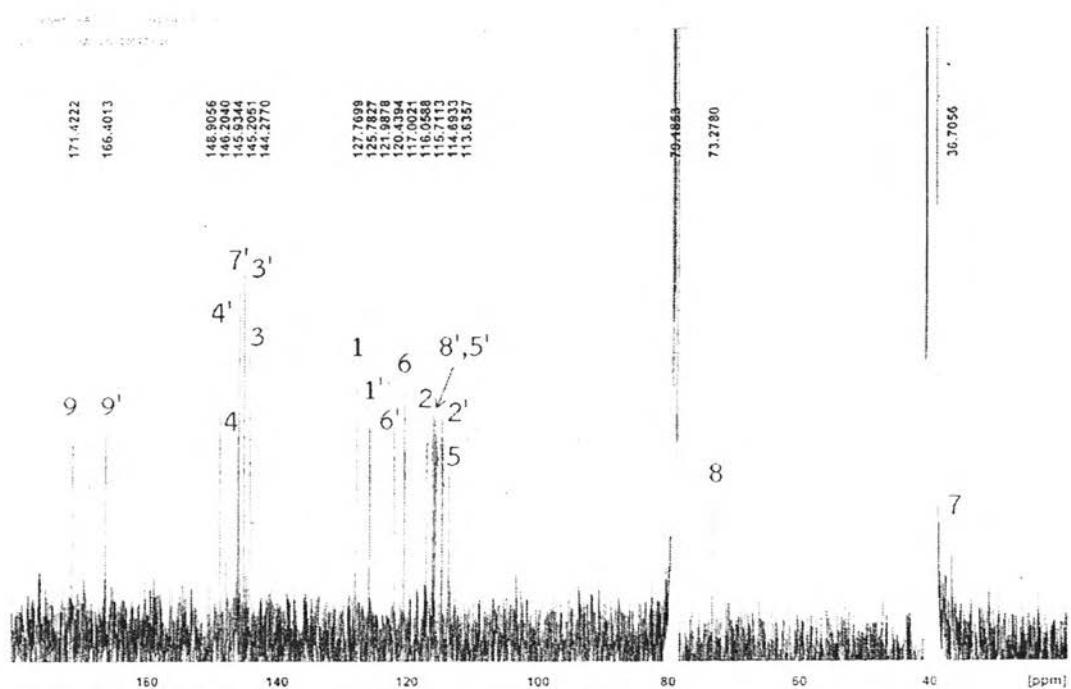
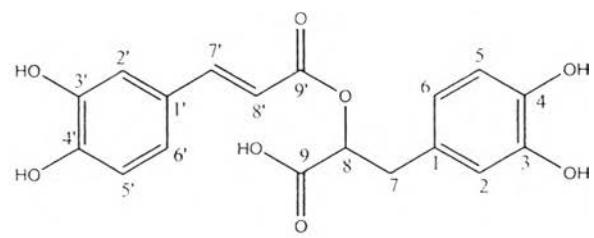


Figure A3 ESI Mass spectrum of compound T4F2AB

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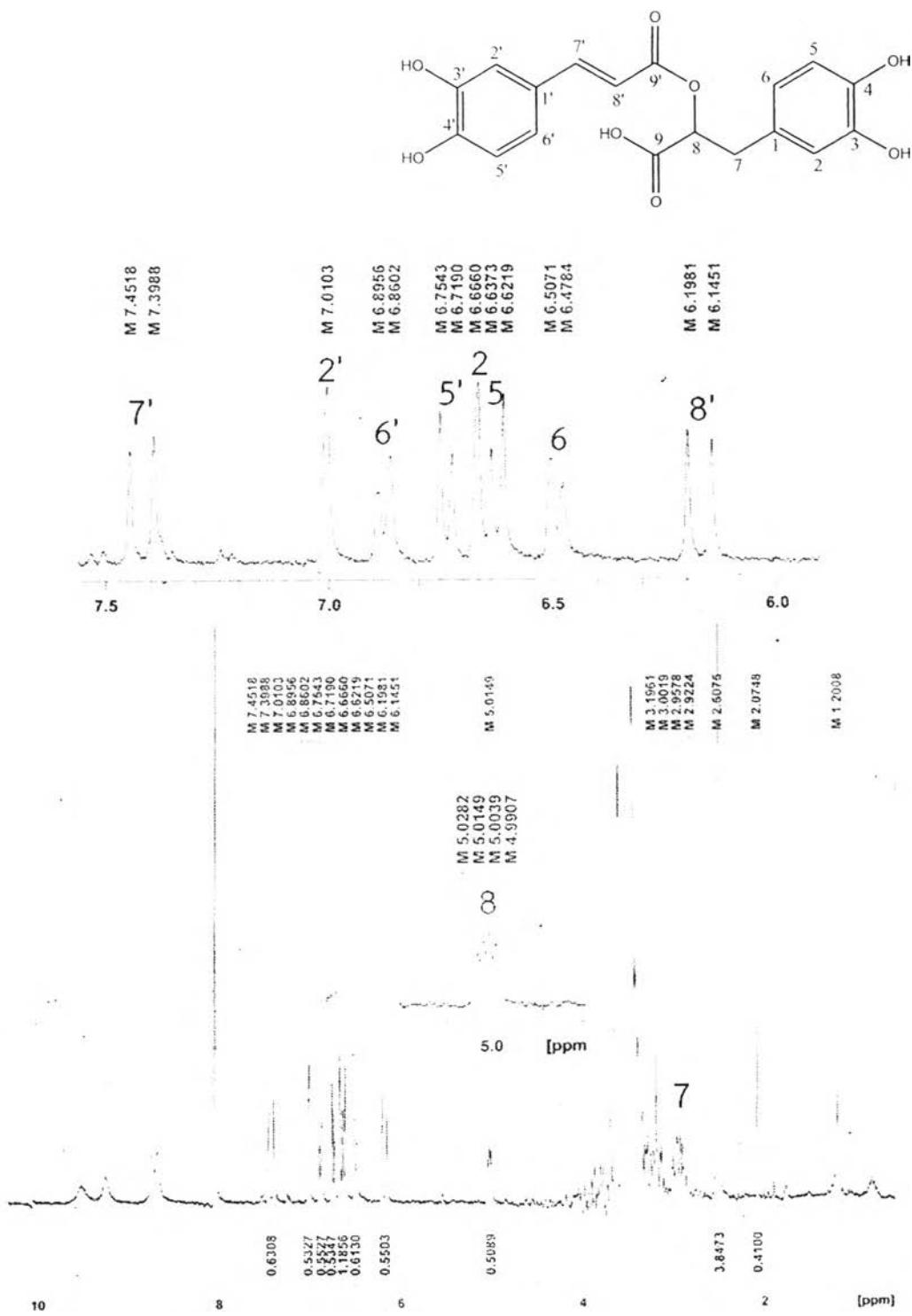
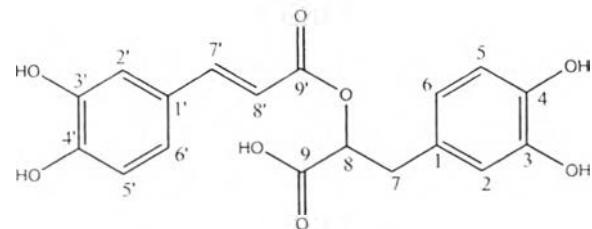


Figure A5 ¹H NMR (300 MHz) Spectrum of compound T5F5 (in DMSO-*d*₆)

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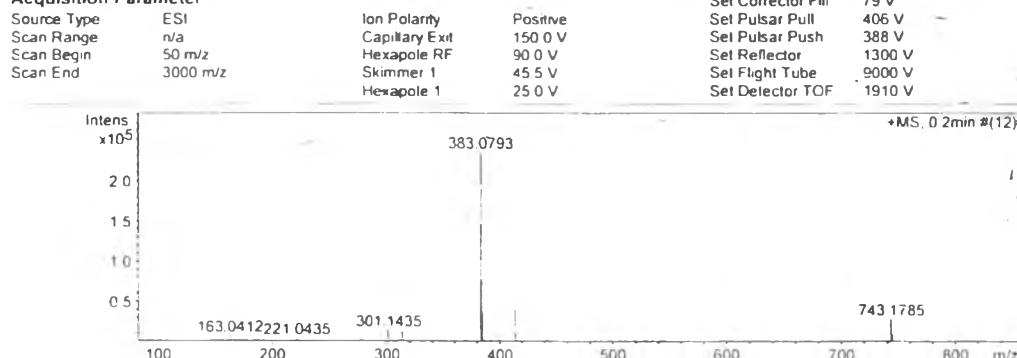
Mass Spectrum List Report

Analysis Info

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Method MKE_tune_low_positive_20130204 m
Sample Name RA
RA

Acquisition Date. 6/5/2014 2:21:58 PM
Operator Administrator
Instrument micrOTOF 72

Acquisition Parameter



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1	163.0412	7550	3.2	902.0	0.0272	6000
2	185.0229	3247	1.4	294.6	0.0293	6308
3	188.0718	1545	0.7	135.6	0.0289	6497
4	221.0435	4795	2.0	312.1	0.0333	5632
5	245.0793	2121	0.9	116.0	0.0341	7177
6	277.1589	2935	1.3	132.5	0.0404	6853
7	302.1467	8427	1.0	96.4	0.0427	7046
8	311.1685	2566	↑ 1	97.7	0.0454	6848
10	313.1802	13029	5.6	492.2	0.0422	7419
11	314.1838	2335	1.0	87.7	0.0442	7112
12	317.1778	2087	0.9	77.3	0.0521	6084
13	333.1540	2234	1.0	77.2	0.0507	6565
14	360.3282	3816	1.6	134.2	0.0470	7668
15	381.3002	1870	0.8	69.3	0.0606	6287
16	383.0793	234309	100.0	8759.7	0.0544	7044
17	384.0821	36335	15.5	1361.8	0.0518	7414
18	385.0831	5738	2.4	215.4	0.0506	7609
19	391.3558	3005	1.3	114.6	0.0586	6676
20	393.3025	1753	0.7	67.1	0.0563	6981
21	399.0525	8848	3.8	345.3	0.0569	7016
22	400.0553	1733	0.7	67.6	0.0560	7140
23	405.0617	2435	1.0	96.5	0.0552	7341
24	413.2717	40662	17.4	1654.1	0.0554	7465
25	414.2743	8915	3.8	363.5	0.0566	7317
26	441.3025	7769	3.3	344.0	0.0593	7444
27	442.3074	1848	0.8	81.8	0.0630	7016
28	743.1785	28418	12.1	2829.2	0.1281	5895
29	744.1809	9501	4.1	948.0	0.1273	5846
30	745.1865	1973	0.8	195.8	0.1378	5408

Figure A6 ESI Mass spectrum of compound T5F5

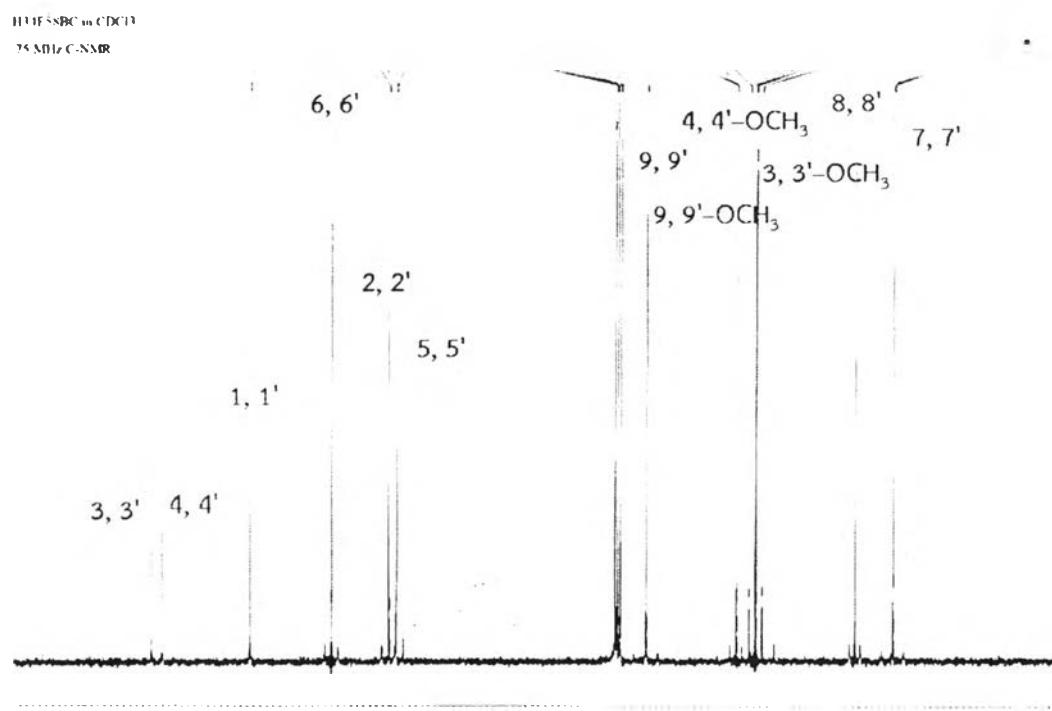
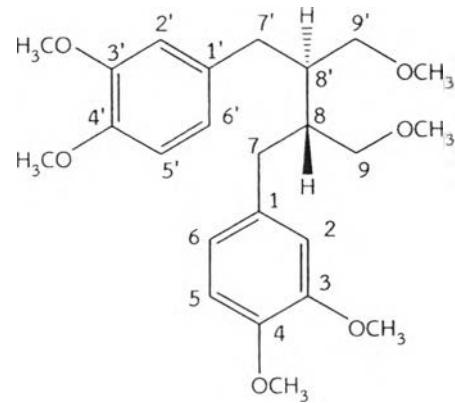


Figure A7 ^{13}C NMR (75 MHz) Spectrum of compound H34F58BC (in CDCl_3)

3697478958

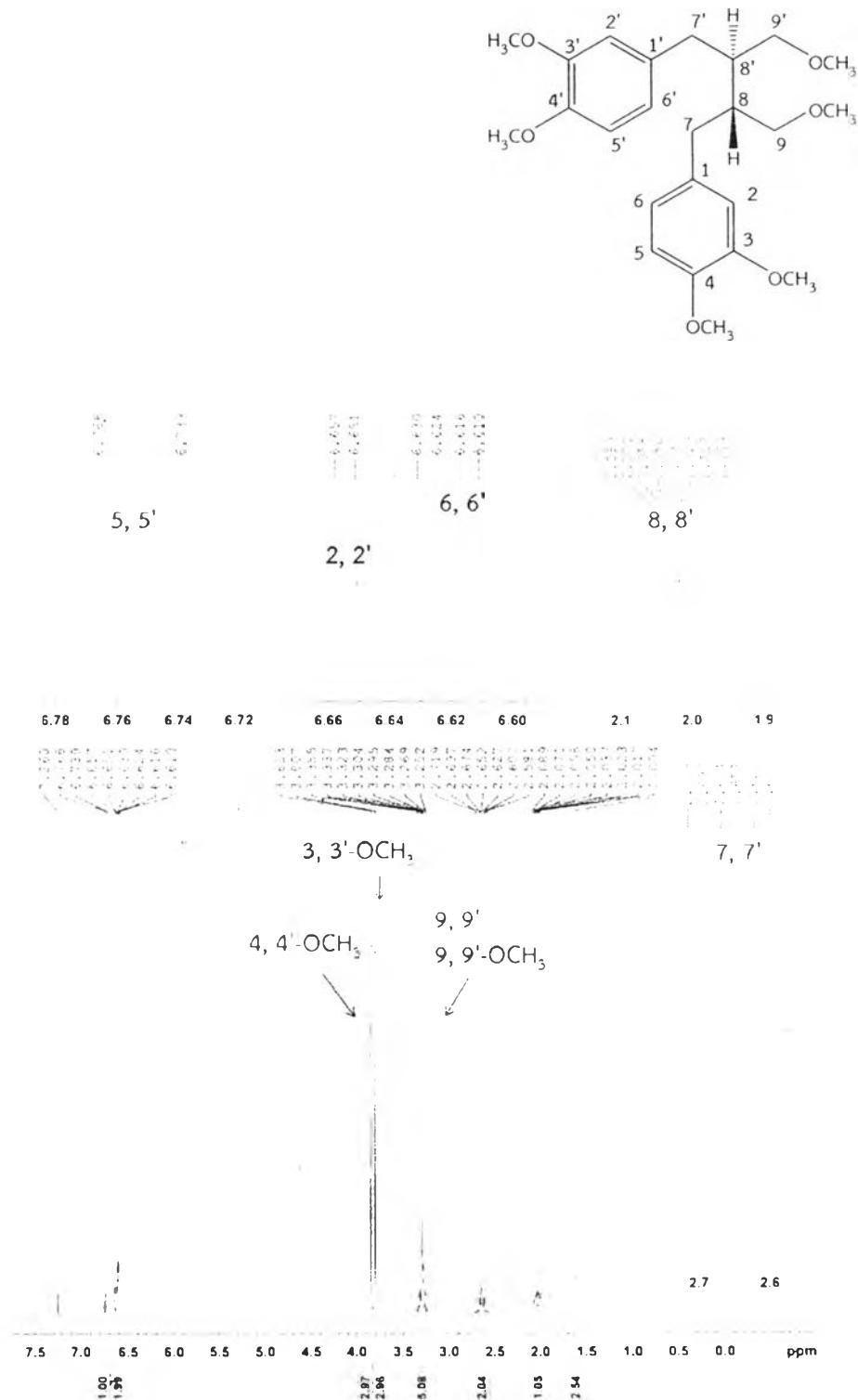
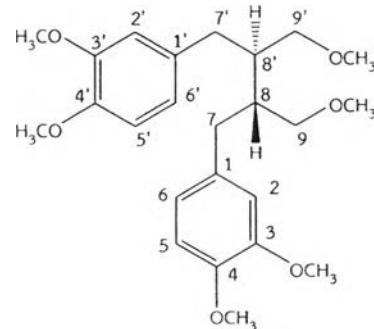


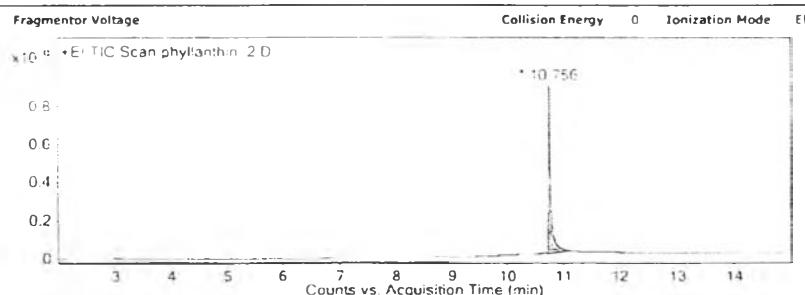
Figure A8 ^1H NMR (300 MHz) Spectrum of compound H34F58BC (in CDCl_3)



Qualitative Analysis Report

Data Filename	phyllanthin_2.D	Sample Name	phyllanthin_2
Sample Type		Position	1
Instrument Name	GCQQQ	User Name	Paweenka Ekkaphan
Acq Method	20140718_Phyllanthin_Phakaporn.CU.M	Acquired Time	7/18/2014 12:53:26 PM
IRM Calibration Status	Not Applicable	DA Method	default.m
Comment			
Expected Barcode		Sample Amount	
Dual Inj Vol	1	TuneName	atunes.eex.tune.xml
TunePath	D:\MassHunter\GCMS\1\17000\tune\	TuneDateStamp	6/17/2014 1:25:15 AM
MSFirmwareVersion	DSP: 7000 2900, q3dServer: G 7090.044 RUN	OperatorName	Paweenka Ekkaphan
RunCompletedFlag	True	Acquisition SW Version	MassHunter GC/MS Acquisition B 07.00 SP2.1654 29 Aug 2013 Copyright © 1989-2013 Agilent Technologies, Inc.

User Chromatograms



Integration Peak List

Peak	Start	RT	End	Height	Area	Area %
1	10.71	10.756	11.144	889494136.2	2643387608	100

User Spectra

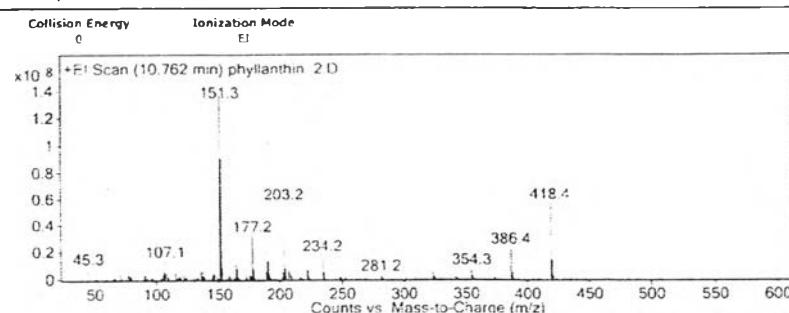


Figure A9 ESI Mass spectrum of compound T5F5

VITA

Miss Pakabhorn Ketmongkhonsit was born on July 31, 1984 in Bangkok, Thailand. She received her B.Sc. in Chemistry in 2006 from the Faculty of Science and Technology, Rajabhat Bansomdejchaopraya University. She was a recipient of THE 90th ANNIVERSARY OF CHULALONGKORN UNIVERSITY FUND (Ratchadaphiseksomphot Endowment Fund) in the year 2010.

Poster presentation

Ketmongkhonsit P. and Chaichantipyuth C. Chemical substances from *Phyllanthus amarus* schum.&thonn. and anti-hepatotoxic activity. Presented at the 9th NRCT-JSPS Joint Seminar “Natural Medicine Research for the Next Decade: New Challenges and Future Collaboration”, December 8-9, 2010, Bangkok, Thailand.

