Cytotoxicity and antimicrobial activity of isolated compounds from *Monsonia angustifolia* and *Dodonaea angustifolia*

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S1. Spectral data of compound 1

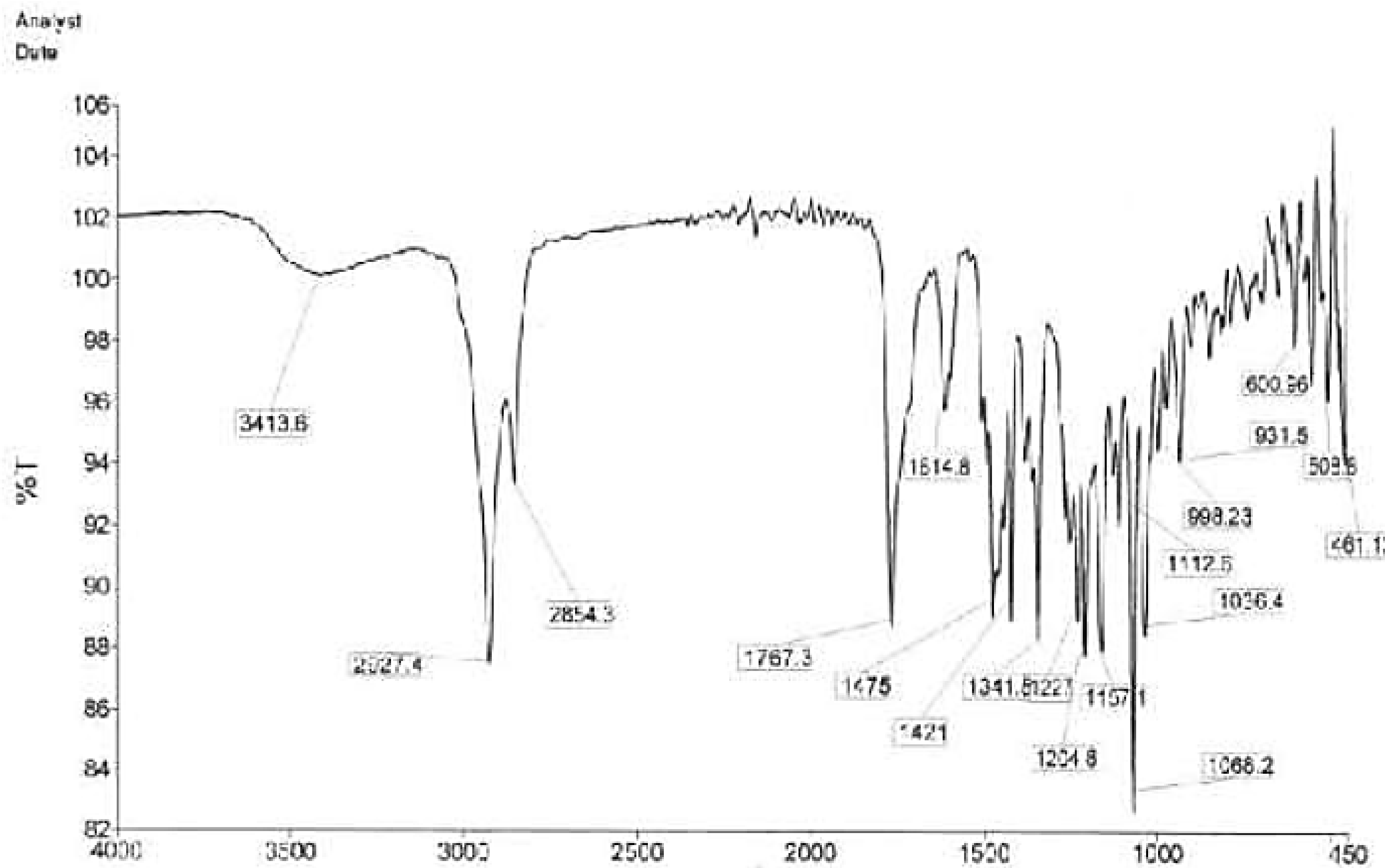


Figure S1.1. Fourier-Transform Infrared Spectroscopy (FTIR) spectrum of 5-methoxyjusticidin A

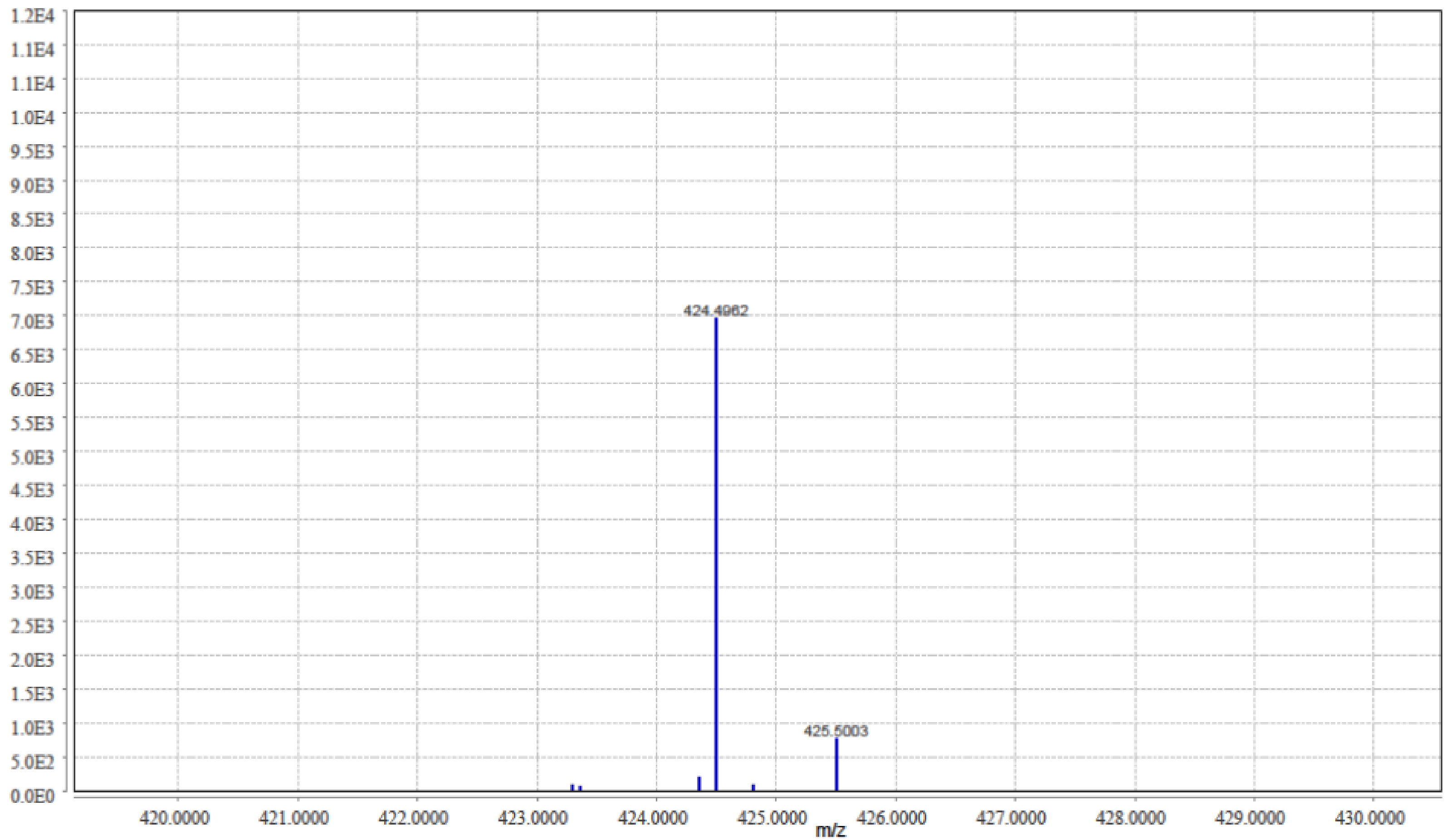
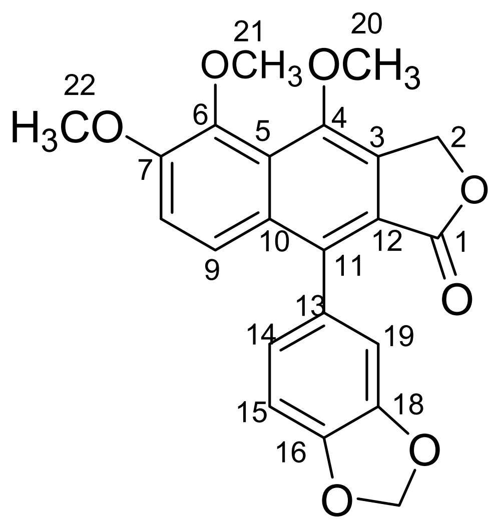


Figure S1.2. High-Resolution Electrospray Ionization Mass spectrum (HR-ESI-MS) of 5-methoxyjusticidin A; $[M+H]^+$ m/z = 425.5003



INDEX	FREQUENCY	PPM	HEIGHT
1	2893.0	7.237	6.5
2	2774.1	6.939	7.8
3	2711.5	6.783	4.8
4	2692.7	6.736	2.4
5	2429.8	6.078	8.0
6	2411.0	6.031	8.7
7	2166.9	5.420	17.1
8	1622.2	4.058	7.2
9	1603.5	4.011	18.5
10	1590.9	3.980	30.4
11	1578.4	3.948	23.5
12	1497.0	3.745	26.3
13	927.4	2.320	2.0
14	808.4	2.022	4.4
15	714.5	1.787	4.9
16	664.5	1.662	5.7
17	626.9	1.568	3.5
18	539.3	1.349	10.9
19	508.0	1.271	11.1
20	489.2	1.224	37.9
21	345.2	0.864	4.4
22	332.7	0.832	4.4

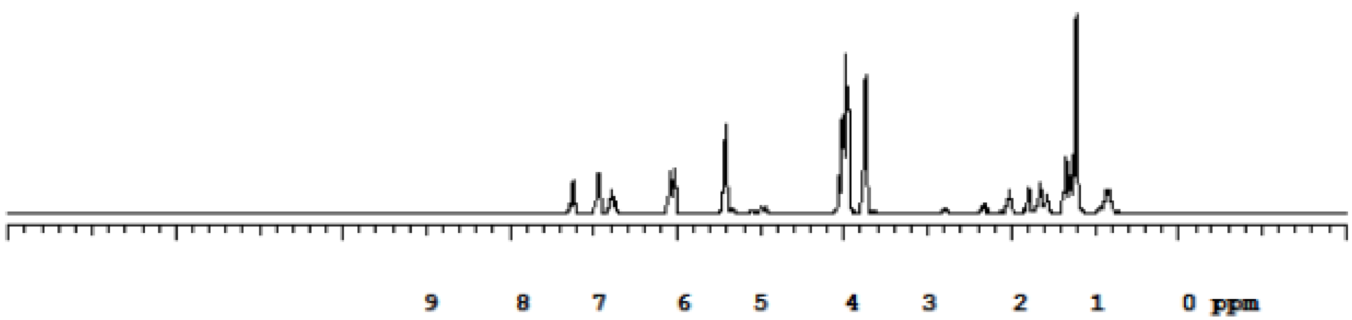


Figure S1.3. Proton Nuclear Magnetic Resonance (^1H NMR) spectrum of 5-methoxyjusticidin A ($CDCl_3$, 400 MHz)

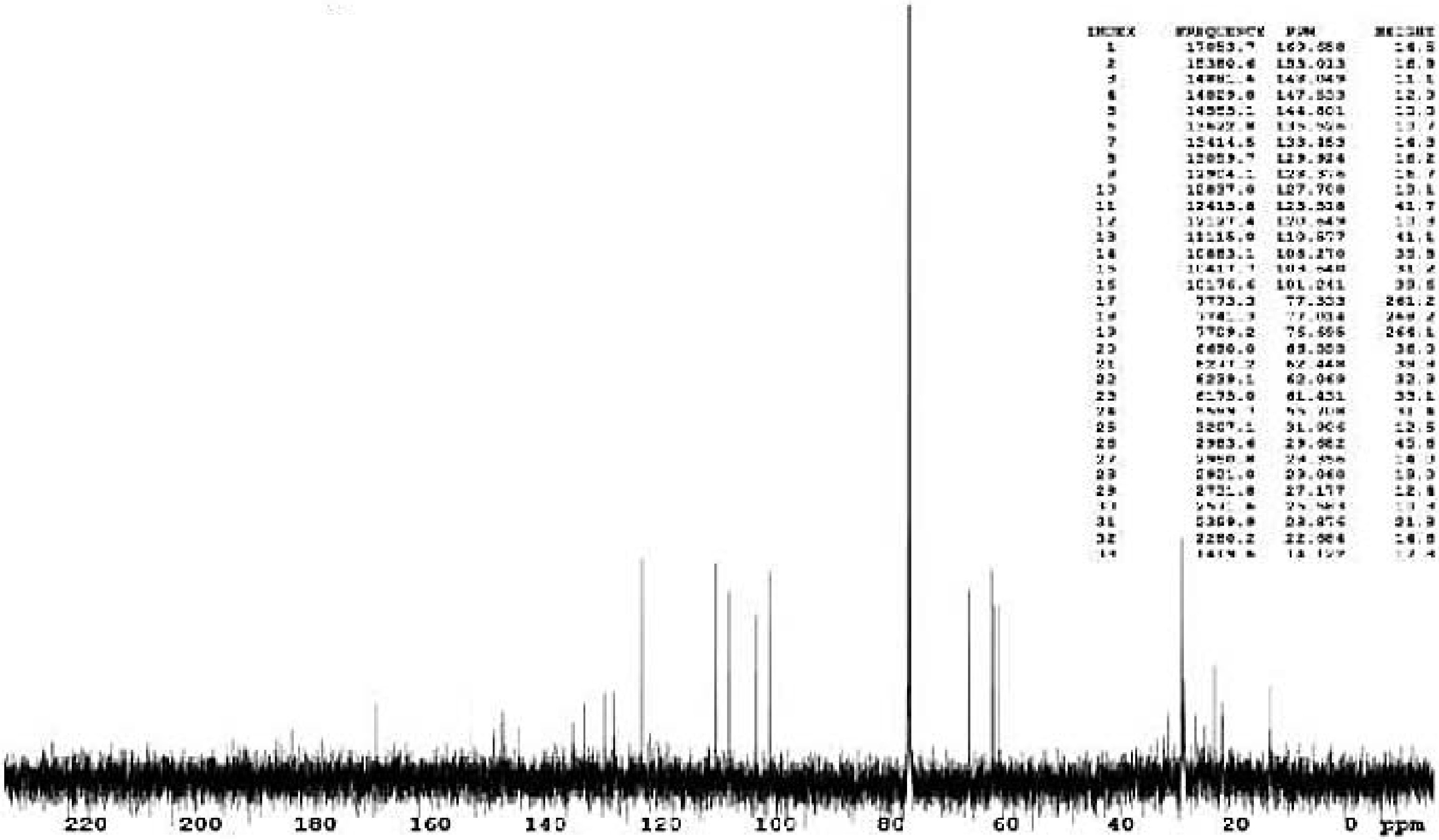


Figure S1.4. Carbon-13 Nuclear Magnetic Resonance (^{13}C NMR) spectrum of 5-methoxyjusticidin A ($CDCl_3$, 100 MHz)

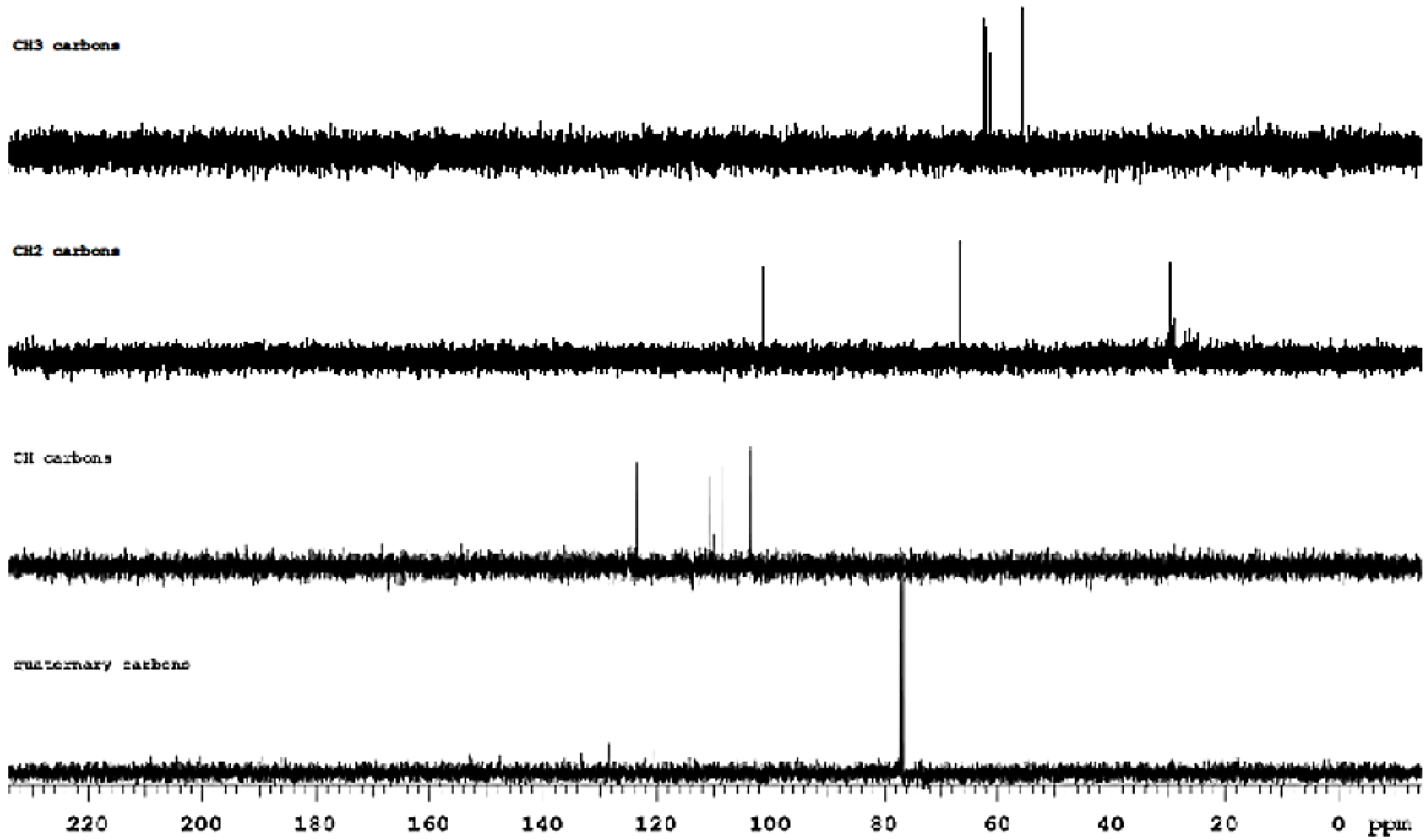


Figure S1.5. Distortionless Enhancement by Polarization Transfer (DEPT) NMR spectra of 5-methoxyjusticidin A ($CDCl_3$, 100 MHz)

S2. Spectral data of compound 2

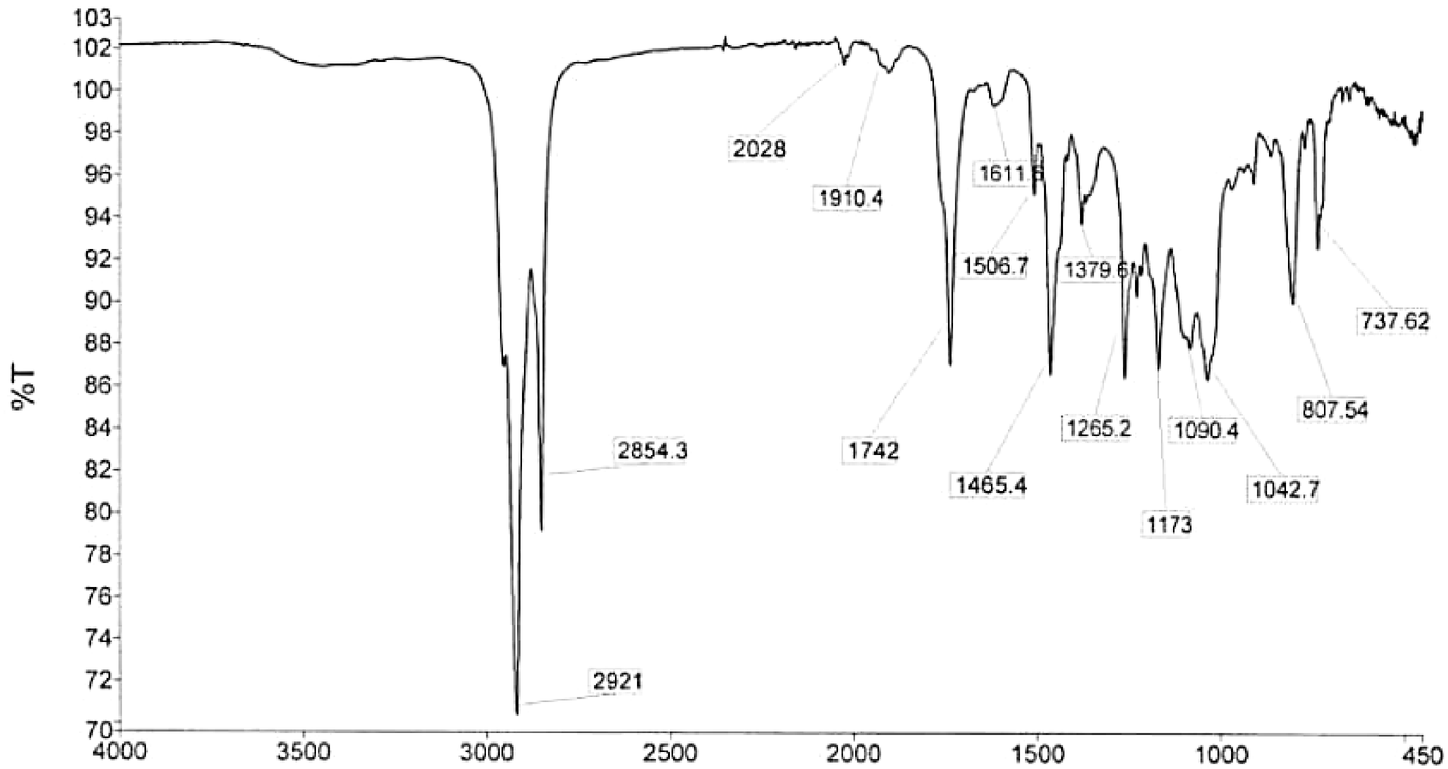


Figure S2.6. Fourier-Transform Infrared Spectroscopy (FTIR) spectrum of *cis*-phytyl diterpenoidal fatty ester

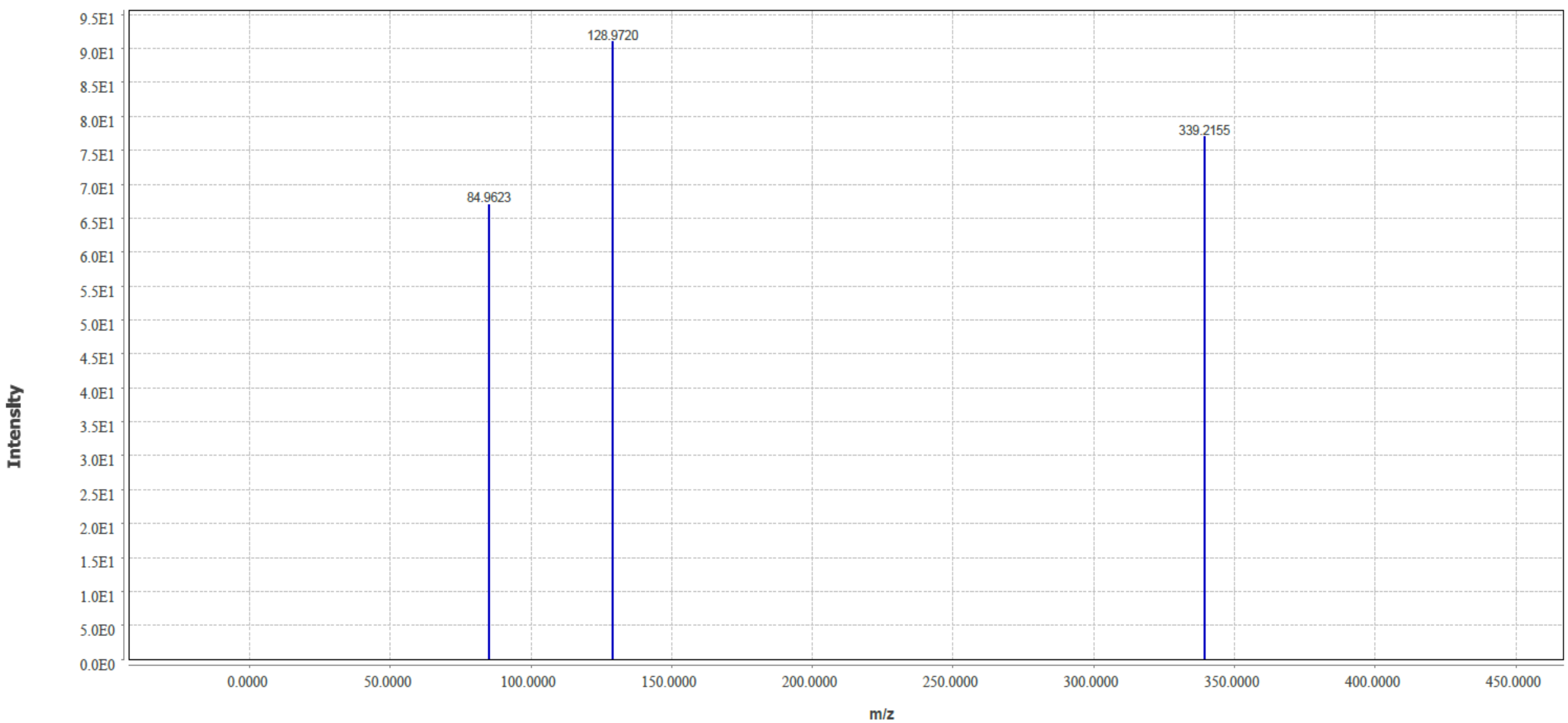
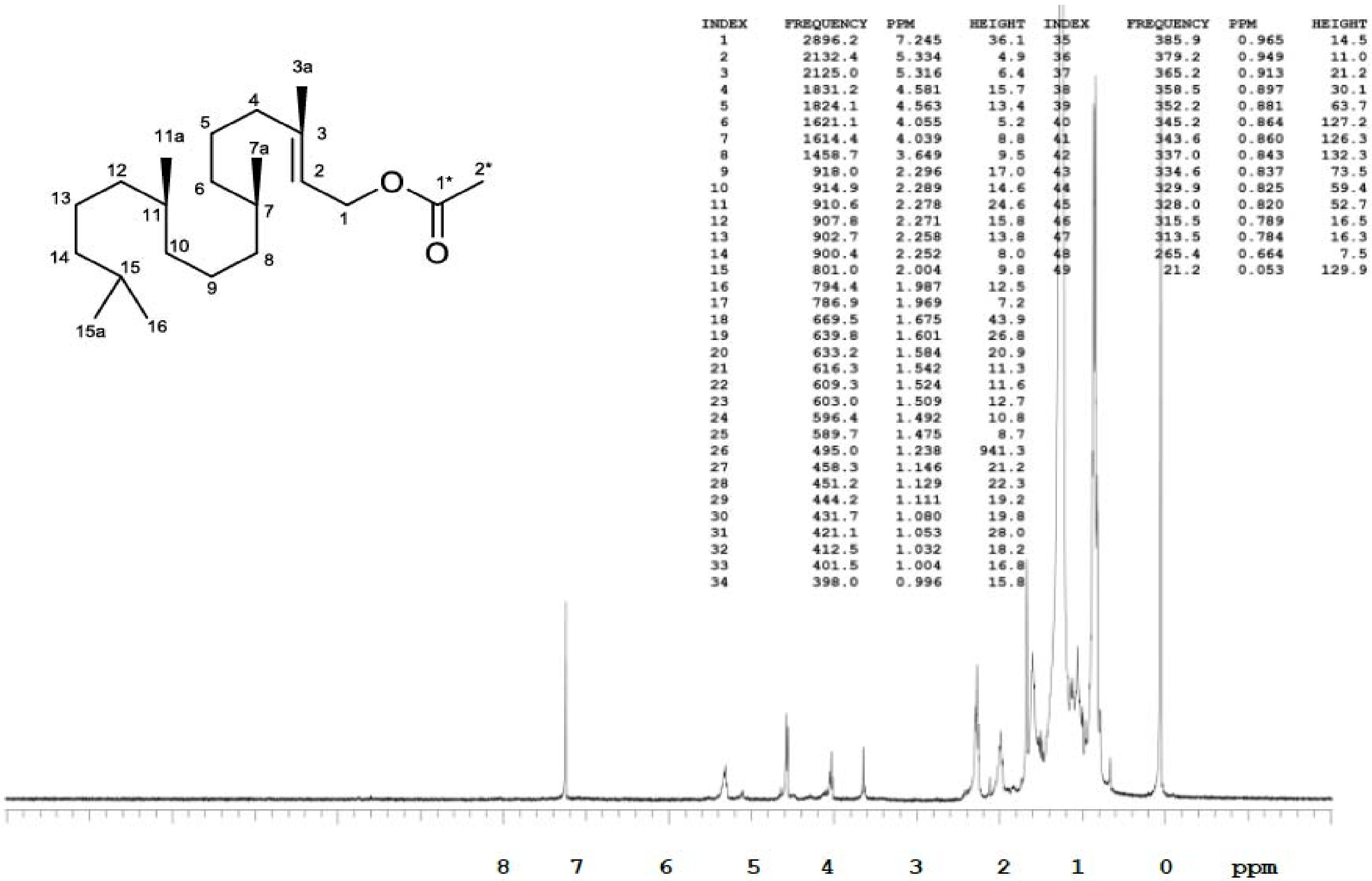
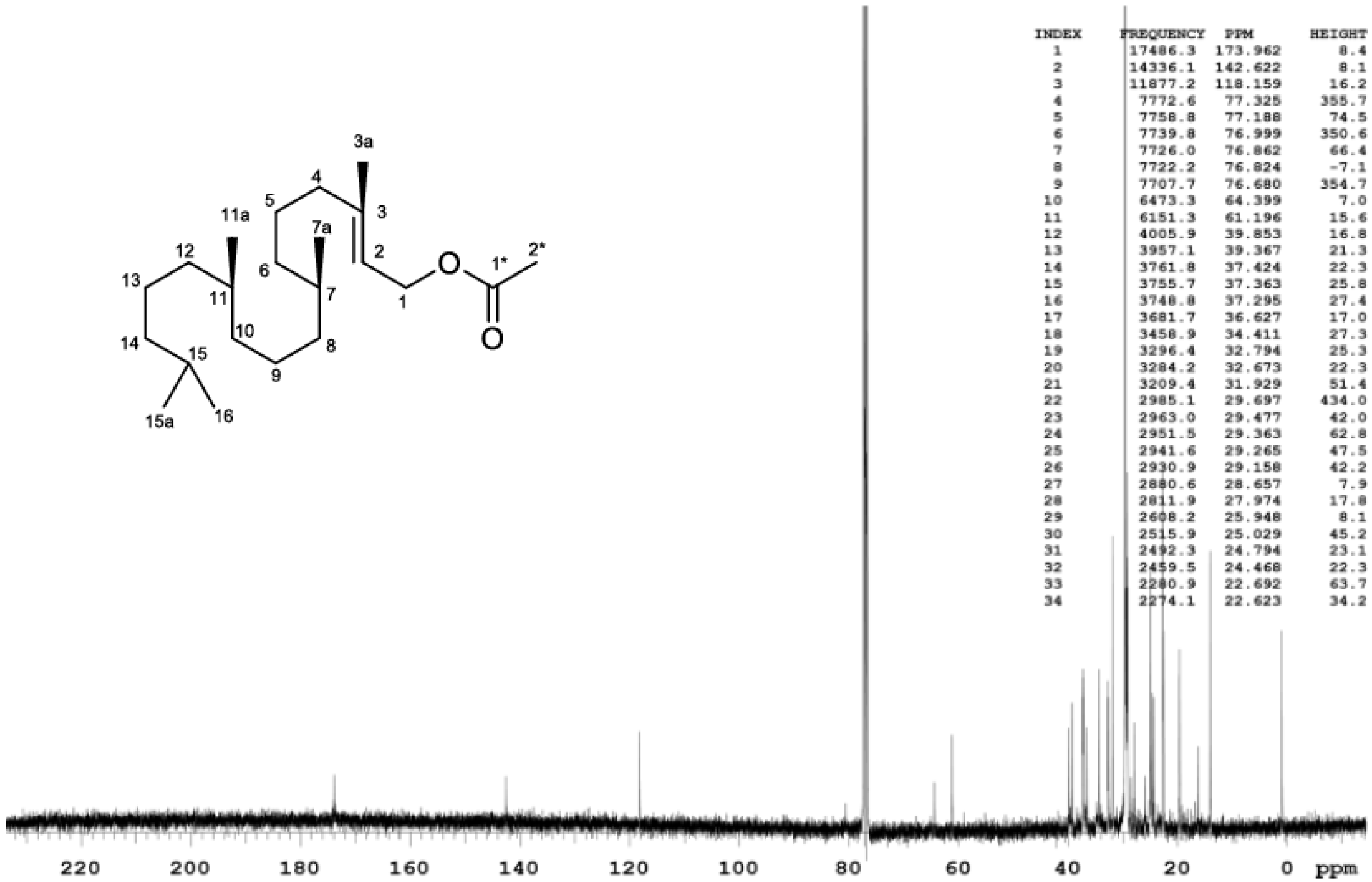


Figure S2.7. High-Resolution Electrospray Ionization Mass spectrum (HR-ESI-MS) of *cis*-phytyl diterpenoidal fatty ester; $[M+H]^+$ m/z = 339.2155



INDEX	FREQUENCY	PPM	HEIGHT	INDEX	FREQUENCY	PPM	HEIGHT
1	2896.2	7.245	36.1	35	385.9	0.965	14.5
2	2132.4	5.334	4.9	36	379.2	0.949	11.0
3	2125.0	5.316	6.4	37	365.2	0.913	21.2
4	1831.2	4.581	15.7	38	358.5	0.897	30.1
5	1824.1	4.563	13.4	39	352.2	0.881	63.7
6	1621.1	4.055	5.2	40	345.2	0.864	127.2
7	1614.4	4.039	8.8	41	343.6	0.860	126.3
8	1458.7	3.649	9.5	42	337.0	0.843	132.3
9	918.0	2.296	17.0	43	334.6	0.837	73.5
10	914.9	2.289	14.6	44	329.9	0.825	59.4
11	910.6	2.278	24.6	45	328.0	0.820	52.7
12	907.8	2.271	15.8	46	315.5	0.789	16.5
13	902.7	2.258	13.8	47	313.5	0.784	16.3
14	900.4	2.252	8.0	48	265.4	0.664	7.5
15	801.0	2.004	9.8	49	21.2	0.053	129.9
16	794.4	1.987	12.5				
17	786.9	1.969	7.2				
18	669.5	1.675	43.9				
19	639.8	1.601	26.8				
20	633.2	1.584	20.9				
21	616.3	1.542	11.3				
22	609.3	1.524	11.6				
23	603.0	1.509	12.7				
24	596.4	1.492	10.8				
25	589.7	1.475	8.7				
26	495.0	1.238	941.3				
27	458.3	1.146	21.2				
28	451.2	1.129	22.3				
29	444.2	1.111	19.2				
30	431.7	1.080	19.8				
31	421.1	1.053	28.0				
32	412.5	1.032	18.2				
33	401.5	1.004	16.8				
34	398.0	0.996	15.8				

Figure S2.8. Proton Nuclear Magnetic Resonance (^1H NMR) spectrum of *cis*-phytyl diterpenoidal fatty ester ($CDCl_3$, 400 MHz)



INDEX	FREQUENCY	PPM	HEIGHT
1	17486.3	173.962	8.4
2	14336.1	142.622	8.1
3	11877.2	118.159	16.2
4	7772.6	77.325	355.7
5	7758.8	77.188	74.5
6	7739.8	76.999	350.6
7	7726.0	76.862	66.4
8	7722.2	76.824	-7.1
9	7707.7	76.680	354.7
10	6473.3	64.399	7.0
11	6151.3	61.196	15.6
12	4005.9	39.853	16.8
13	3957.1	39.367	21.3
14	3761.8	37.424	22.3
15	3755.7	37.363	25.8
16	3748.8	37.295	27.4
17	3681.7	36.627	17.0
18	3458.9	34.411	27.3
19	3296.4	32.794	25.3
20	3284.2	32.673	22.3
21	3209.4	31.929	51.4
22	2985.1	29.697	434.0
23	2963.0	29.477	42.0
24	2951.5	29.363	62.8
25	2941.6	29.265	47.5
26	2930.9	29.158	42.2
27	2880.6	28.657	7.9
28	2811.9	27.974	17.8
29	2608.2	25.948	8.1
30	2515.9	25.029	45.2
31	2492.3	24.794	23.1
32	2459.5	24.468	22.3
33	2280.9	22.692	63.7
34	2274.1	22.623	34.2

Figure S2.9. Carbon-13 Nuclear Magnetic Resonance (^{13}C NMR) spectrum of *cis*-phytyl diterpenoidal fatty ester ($CDCl_3$, 100 MHz)

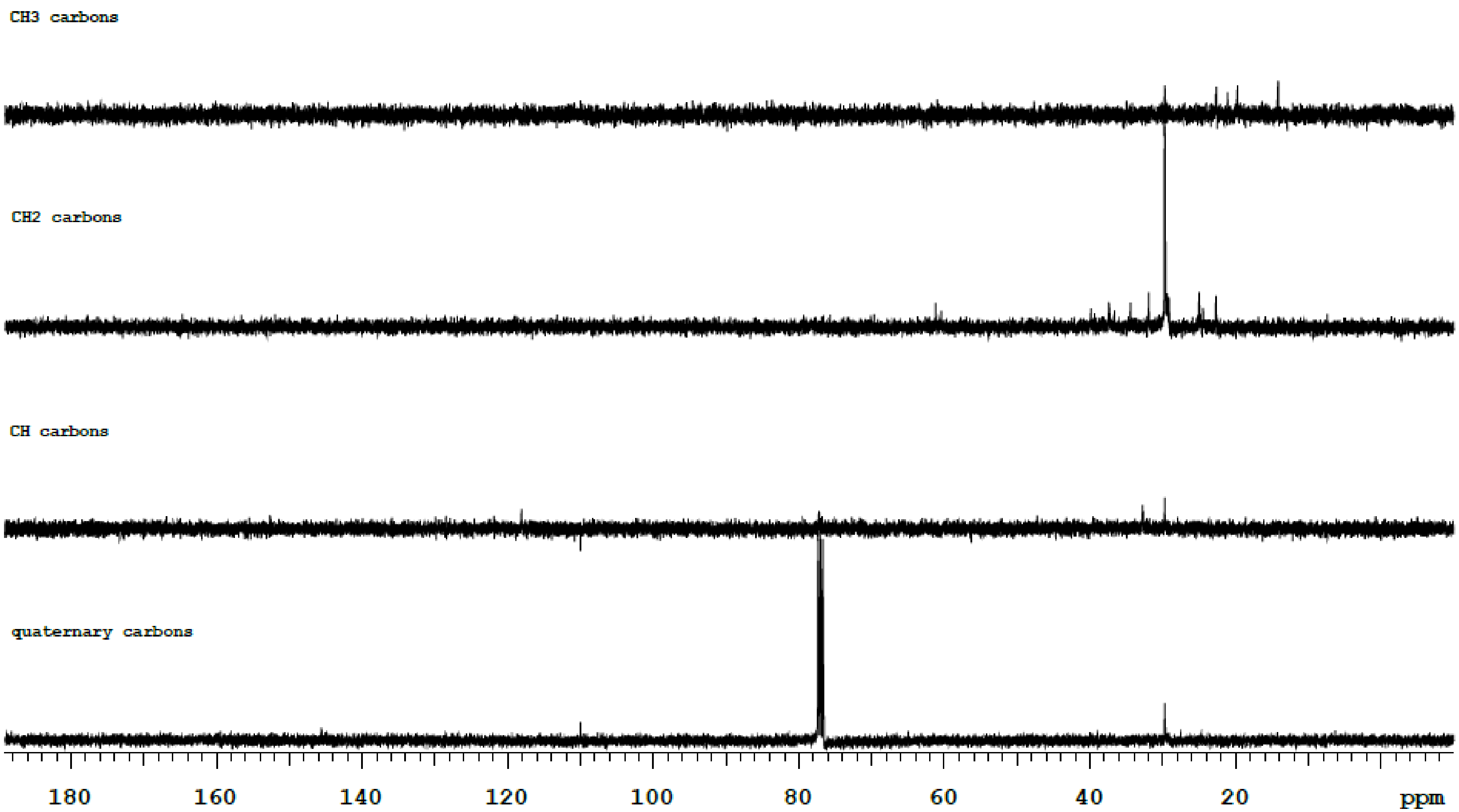


Figure S2.10. Distortionless Enhancement by Polarization Transfer (DEPT) NMR spectra of *cis*-phytyl diterpenoidal fatty ester ($CDCl_3$, 100 MHz)

S3. Spectral data of compound 3

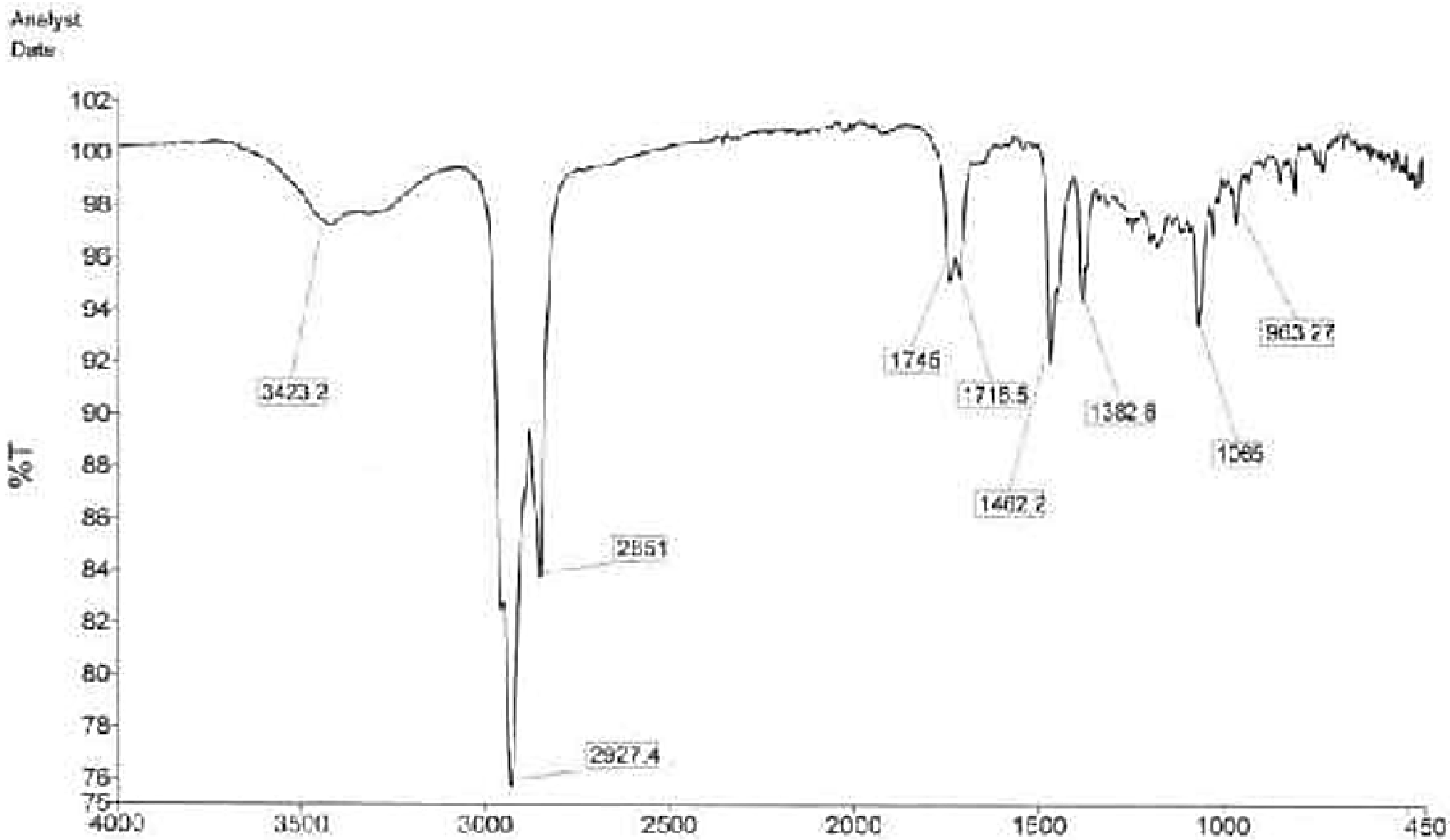


Figure S3.11 Fourier-Transform Infrared Spectroscopy (FTIR) spectrum of stigmasterol

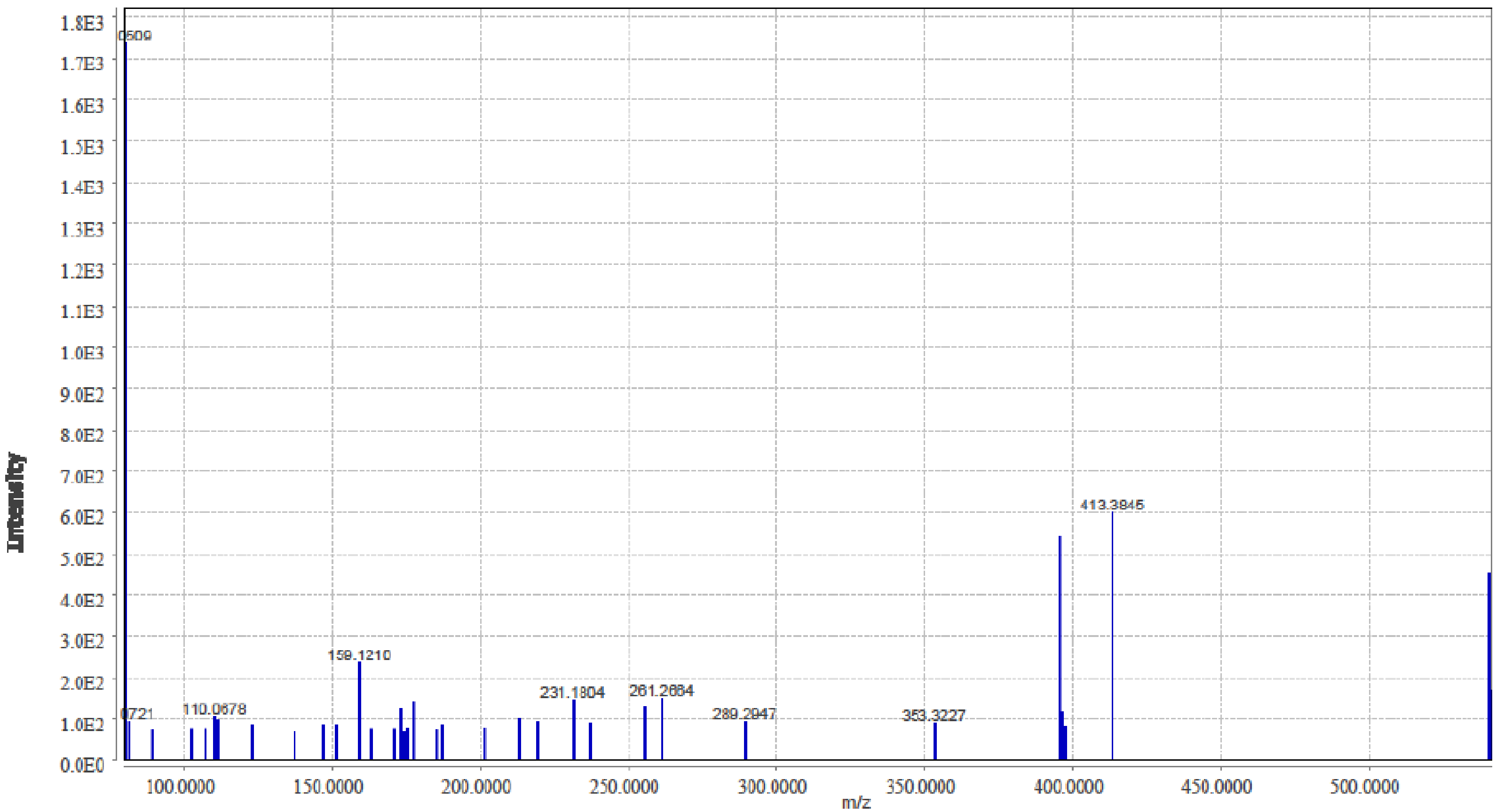
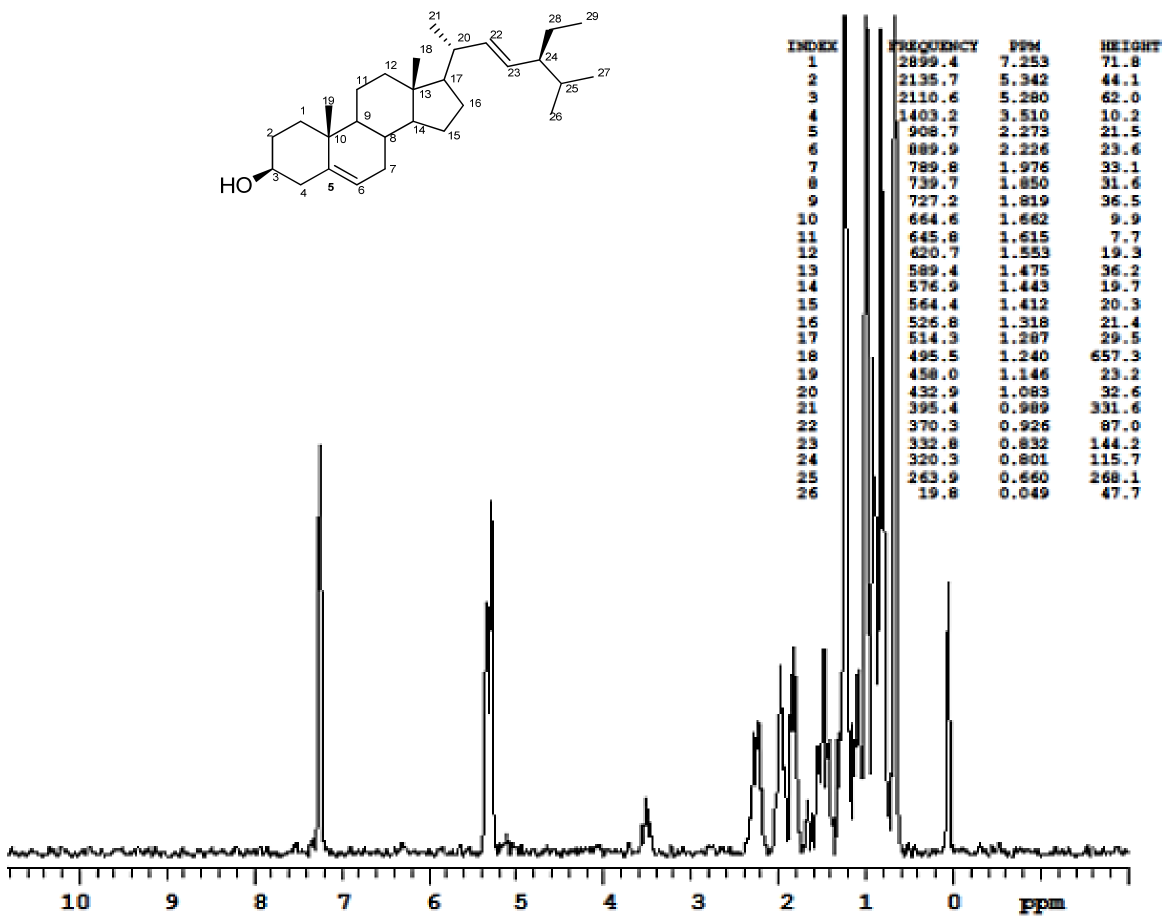
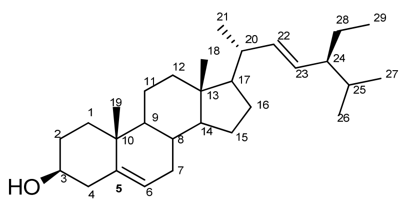


Figure S3.12 High-Resolution Electrospray Ionization Mass spectrum (HR-ESI-MS) of stigmasterol



INDEX	FREQUENCY	PPM	HEIGHT
1	2899.4	7.253	71.8
2	2135.7	5.342	44.1
3	2110.6	5.280	62.0
4	1403.2	3.510	10.2
5	908.7	2.273	21.5
6	889.9	2.226	23.6
7	789.8	1.976	33.1
8	739.7	1.850	31.6
9	727.2	1.819	36.5
10	664.6	1.662	9.9
11	645.8	1.615	7.7
12	620.7	1.553	19.3
13	589.4	1.475	36.2
14	576.9	1.443	19.7
15	564.4	1.412	20.3
16	526.8	1.318	21.4
17	514.3	1.287	29.5
18	495.5	1.240	657.3
19	458.0	1.146	23.2
20	432.9	1.083	32.6
21	395.4	0.989	331.6
22	370.3	0.926	87.0
23	332.8	0.832	144.2
24	320.3	0.801	115.7
25	263.9	0.660	268.1
26	19.8	0.049	47.7

Figure S3.13 Proton Nuclear Magnetic Resonance (^{1}H NMR) spectrum of stigmasterol



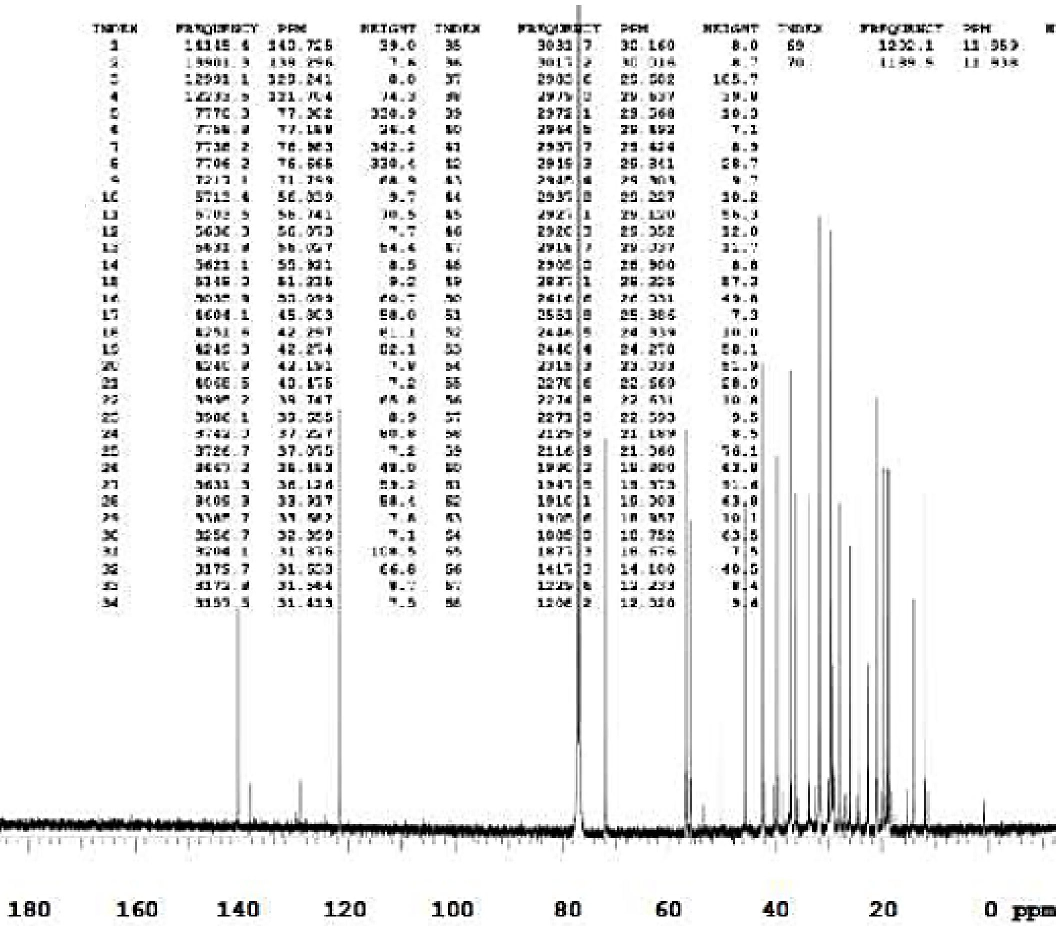


Figure S3.14 Carbon-13 Nuclear Magnetic Resonance (^{13}C NMR) spectrum of stigmasterol

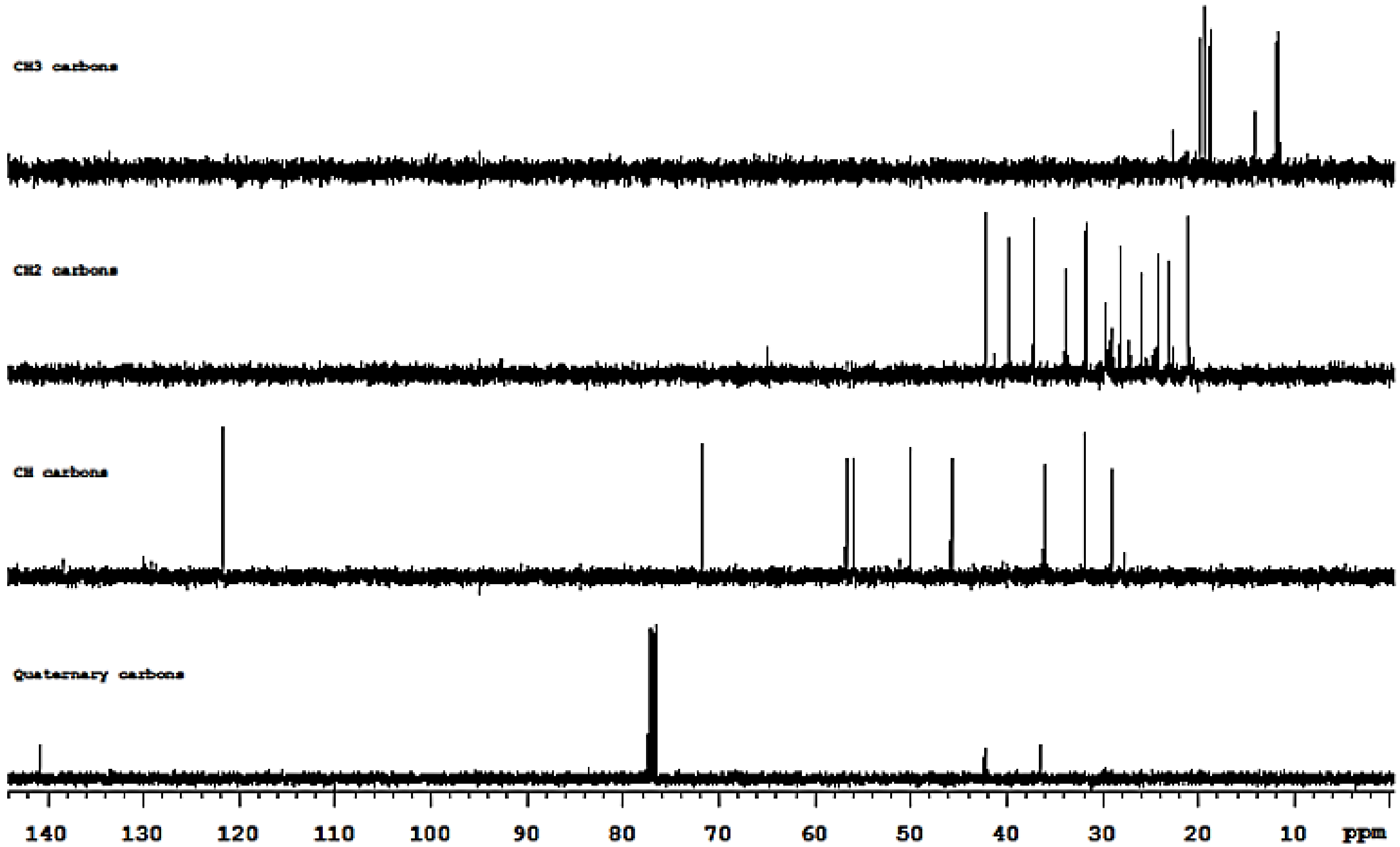


Figure S3.15 Distortionless Enhancement by Polarization Transfer (DEPT) NMR spectra of stigmasterol ($CDCl_3$, 100 MHz)

S4. Spectral data of compound 4

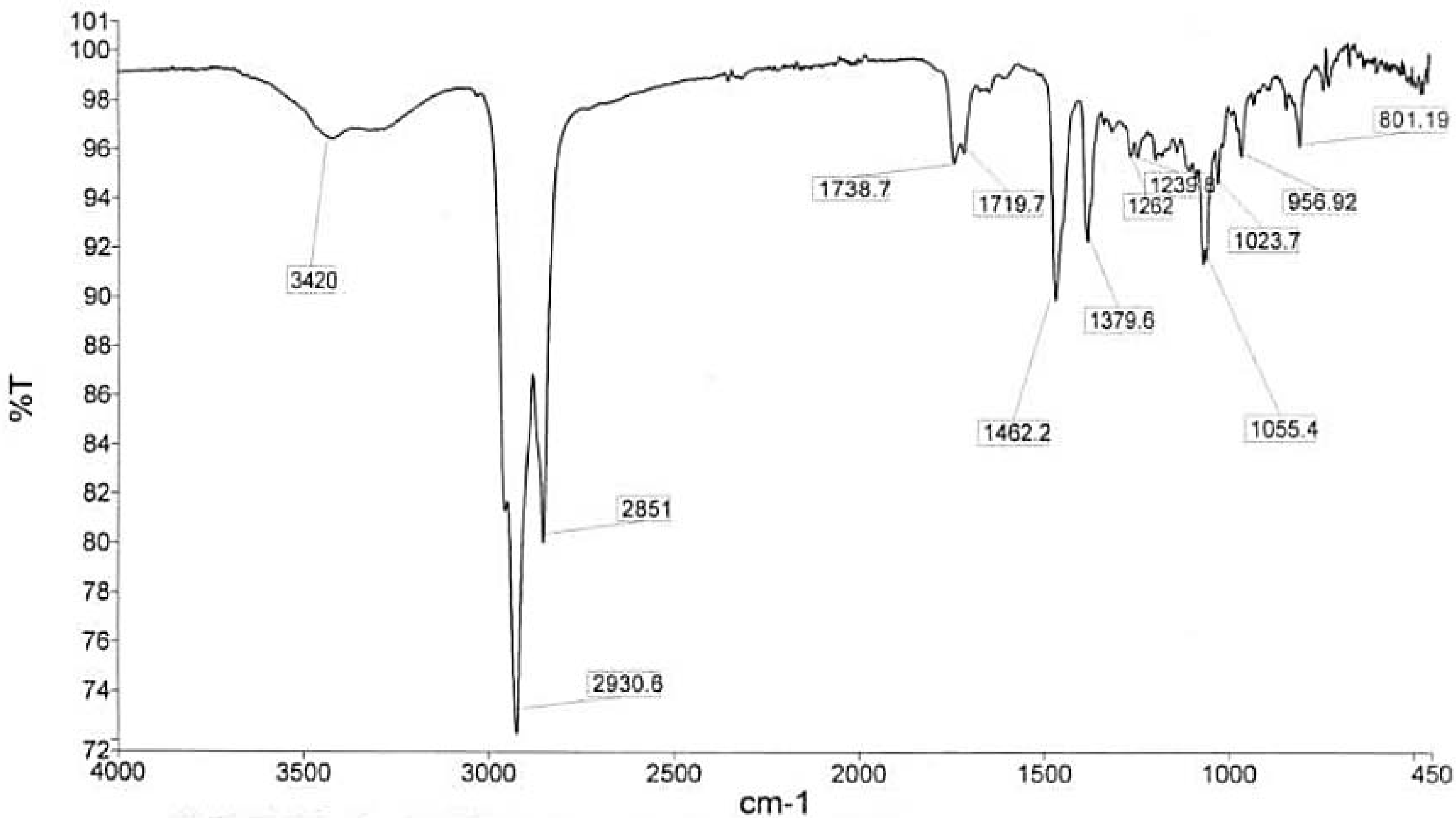


Figure S4.16. Fourier-Transform Infrared Spectroscopy (FTIR) spectrum of β-sitosterol

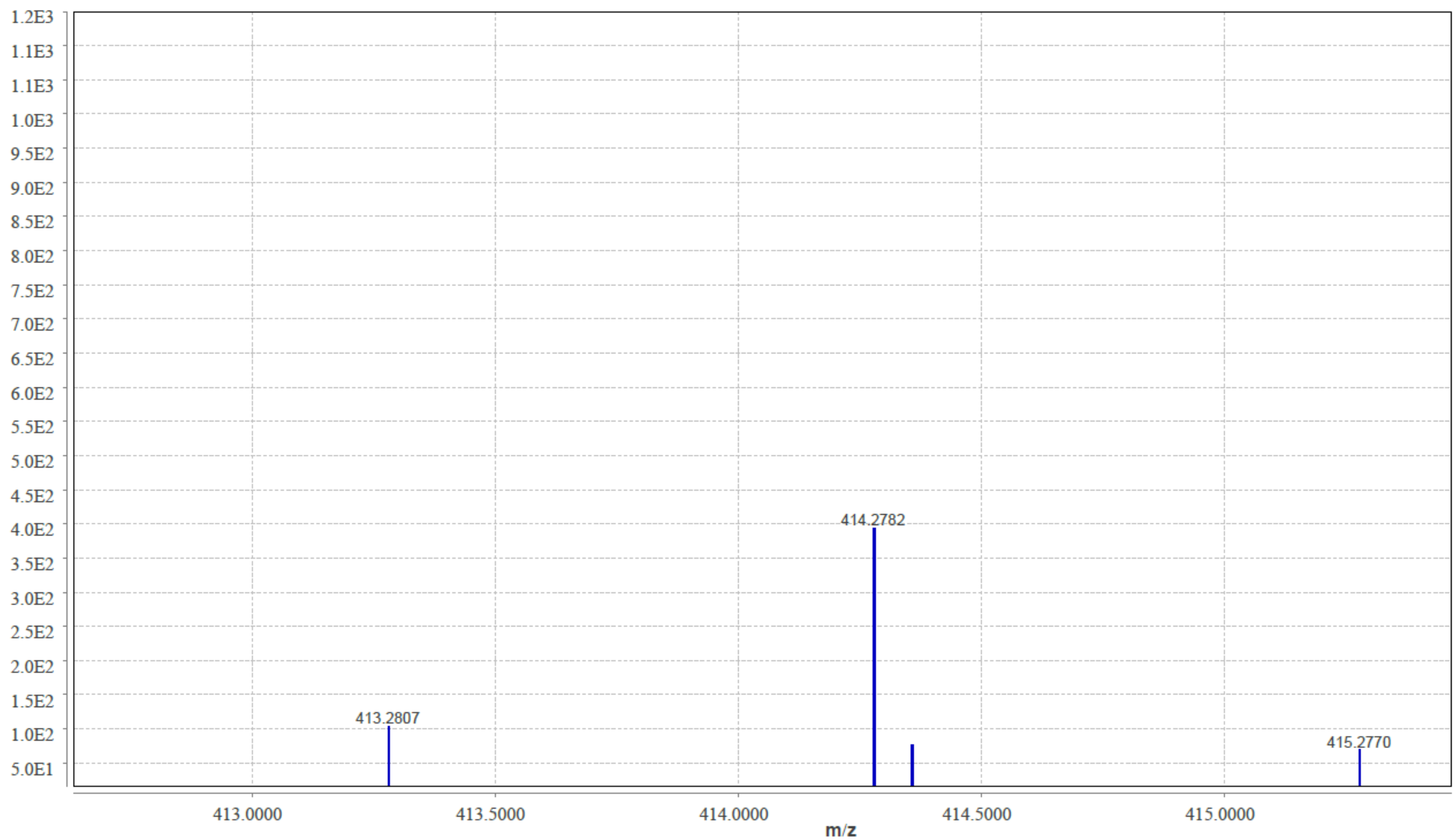


Figure S4.17 High-Resolution Electrospray Ionization Mass spectrum (HR-ESI-MS) of β-sitosterol; [M+H] = 415.27

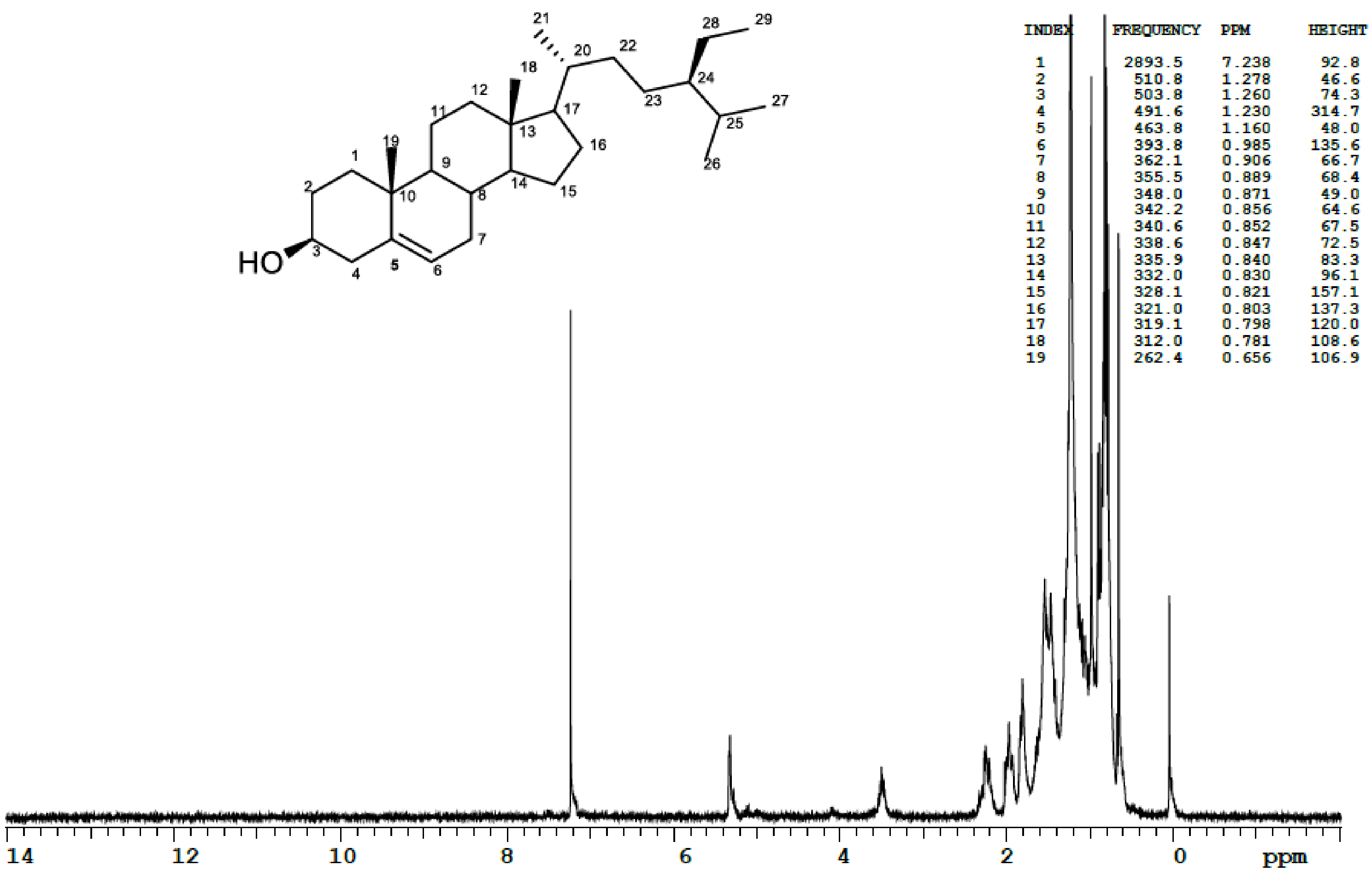
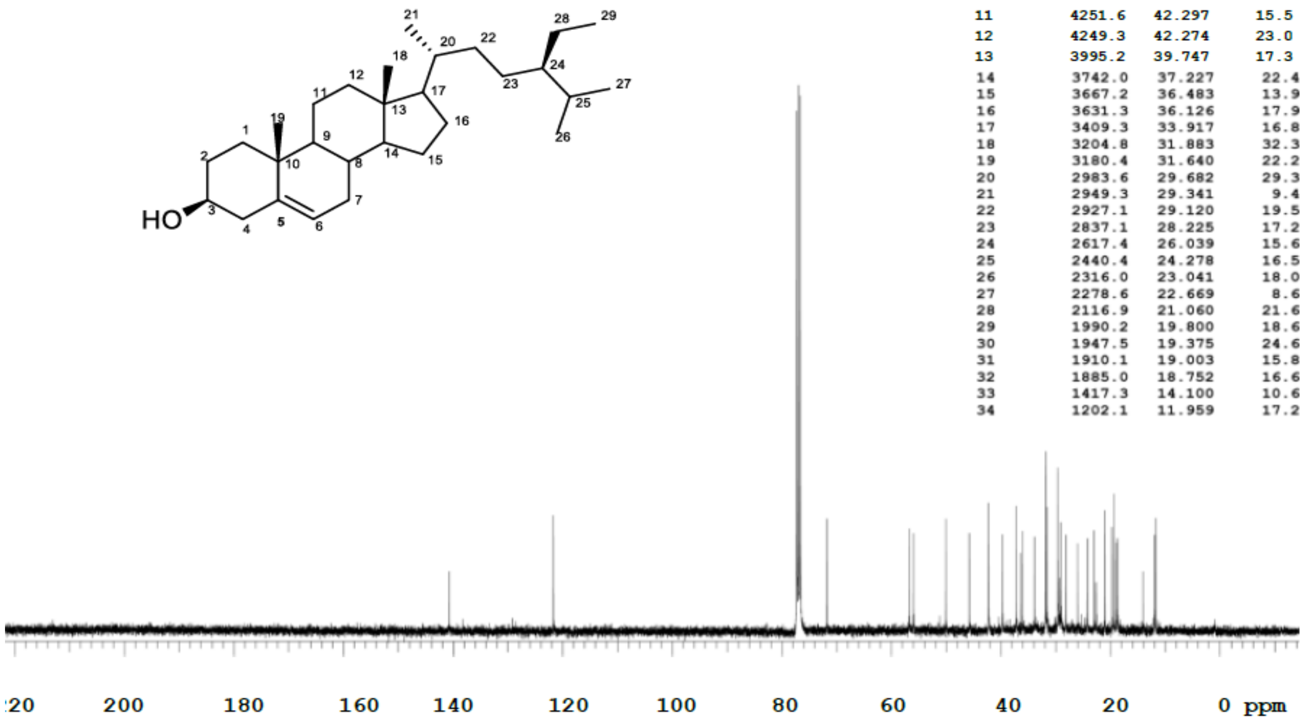


Figure S4.18 Proton Nuclear Magnetic Resonance (^{1}H NMR) spectrum of β-sitosterol



INDEX	FREQUENCY	PPM	HEIGHT	INDEX	FREQ
1	14145.4	140.725	10.6	35	11
2	12233.5	121.704	20.7		
3	7770.3	77.302	93.7		
4	7738.2	76.983	98.4		
5	7706.2	76.665	96.5		
6	7217.1	71.799	20.1		
7	5703.5	56.741	18.3		
8	5631.8	56.027	17.5		
9	5036.7	50.107	20.1		
10	4604.8	45.811	17.5		
11	4251.6	42.297	15.5		
12	4249.3	42.274	23.0		
13	3995.2	39.747	17.3		
14	3742.0	37.227	22.4		
15	3667.2	36.483	13.9		
16	3631.3	36.126	17.9		
17	3409.3	33.917	16.8		
18	3204.8	31.883	32.3		
19	3180.4	31.640	22.2		
20	2983.6	29.682	29.3		
21	2949.3	29.341	9.4		
22	2927.1	29.120	19.5		
23	2837.1	28.225	17.2		
24	2617.4	26.039	15.6		
25	2440.4	24.278	16.5		
26	2316.0	23.041	18.0		
27	2278.6	22.669	8.6		
28	2116.9	21.060	21.6		
29	1990.2	19.800	18.6		
30	1947.5	19.375	24.6		
31	1910.1	19.003	15.8		
32	1885.0	18.752	16.6		
33	1417.3	14.100	10.6		
34	1202.1	11.959	17.2		

Figure S4.19 Carbon-13 Nuclear Magnetic Resonance (^{13}C NMR) spectrum of β-sitosterol

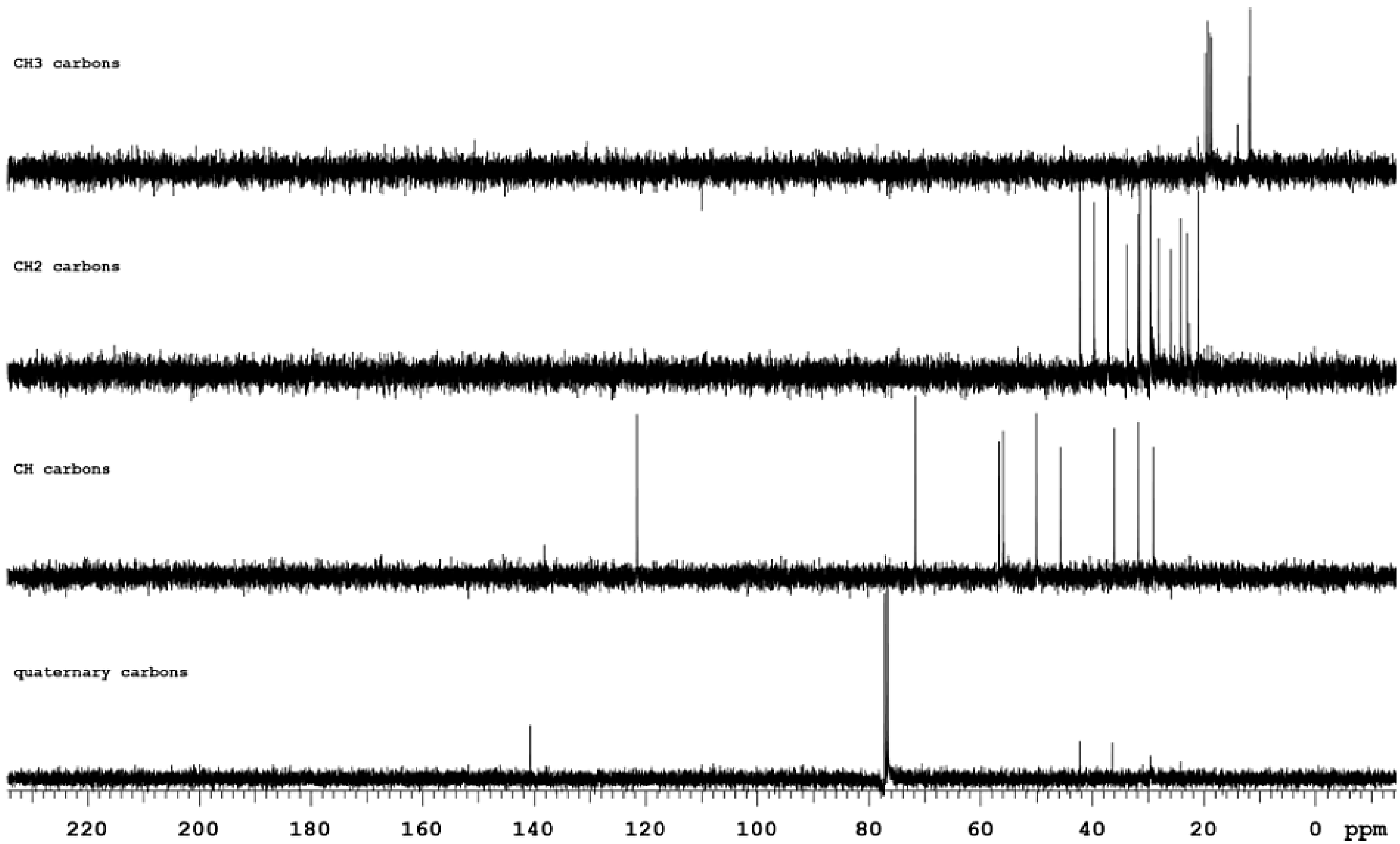


Figure S4.20 Distortionless Enhancement by Polarization Transfer (DEPT) NMR spectra of β-sitosterol ($CDCl_3$, 100 MHz)

S5. Spectral data of compound 5

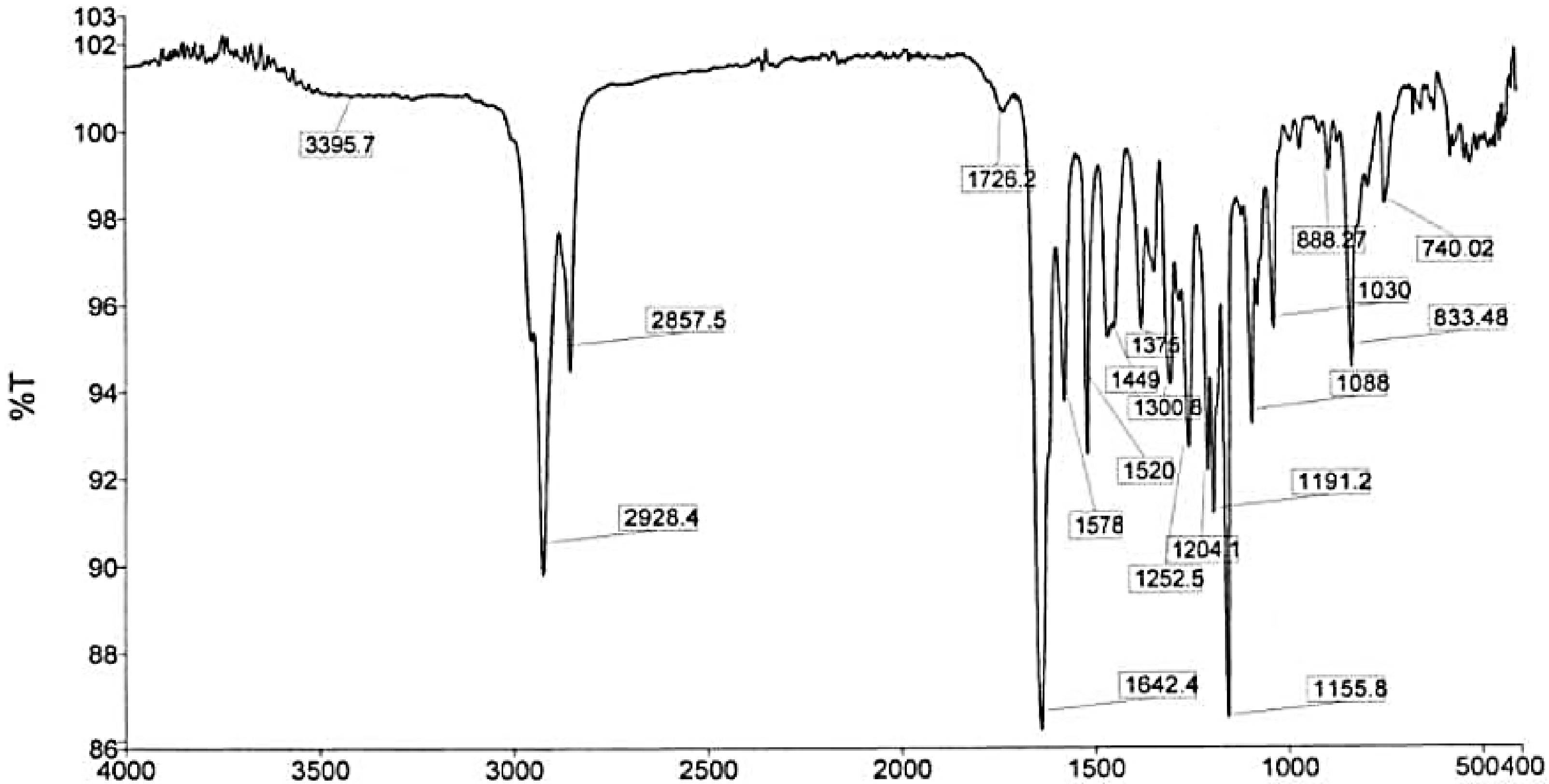


Figure S5.21. Fourier-Transform Infrared Spectroscopy (FTIR) spectrum of 5-hydroxy-7,4`-dimethoxyflavone

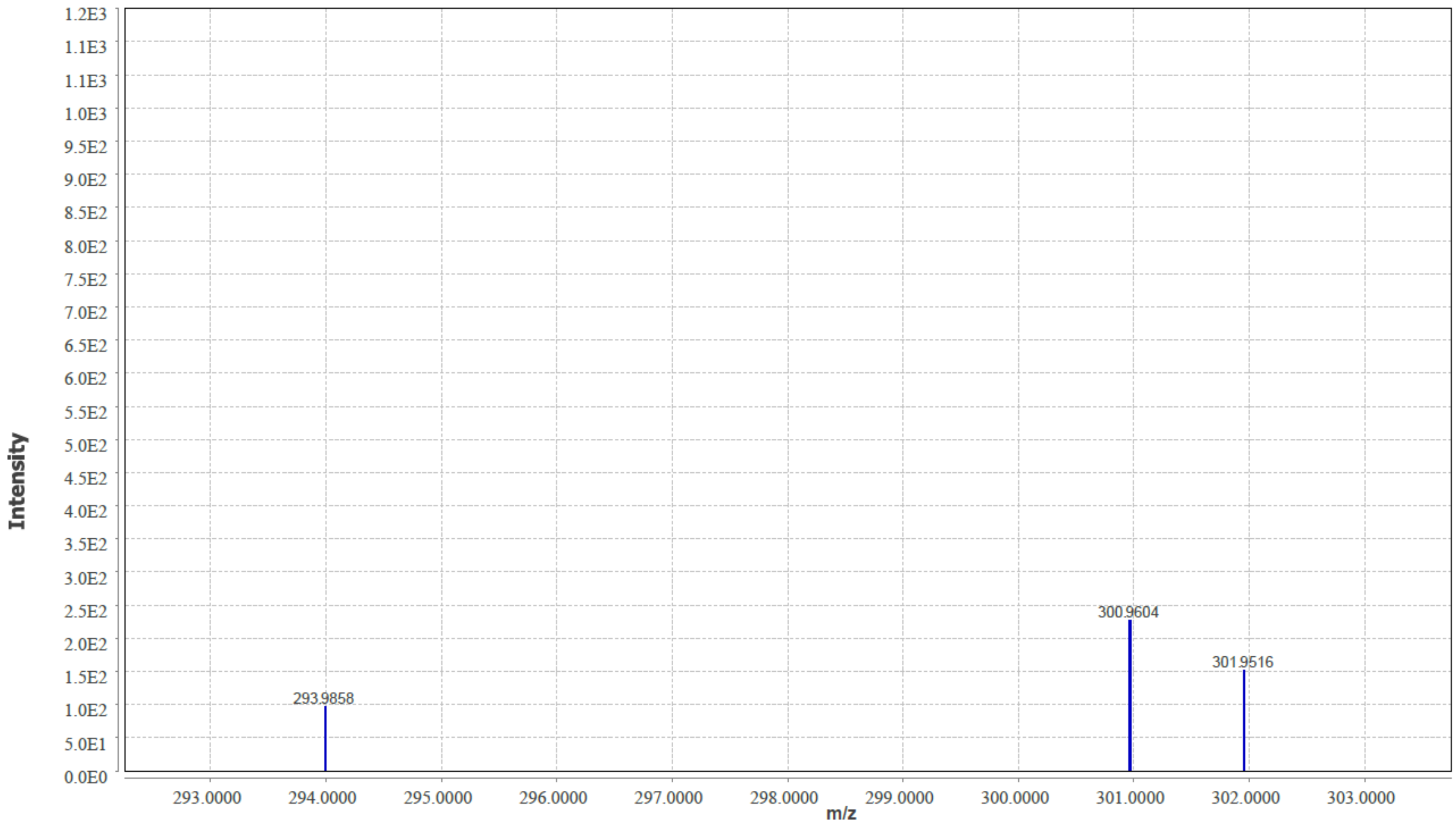
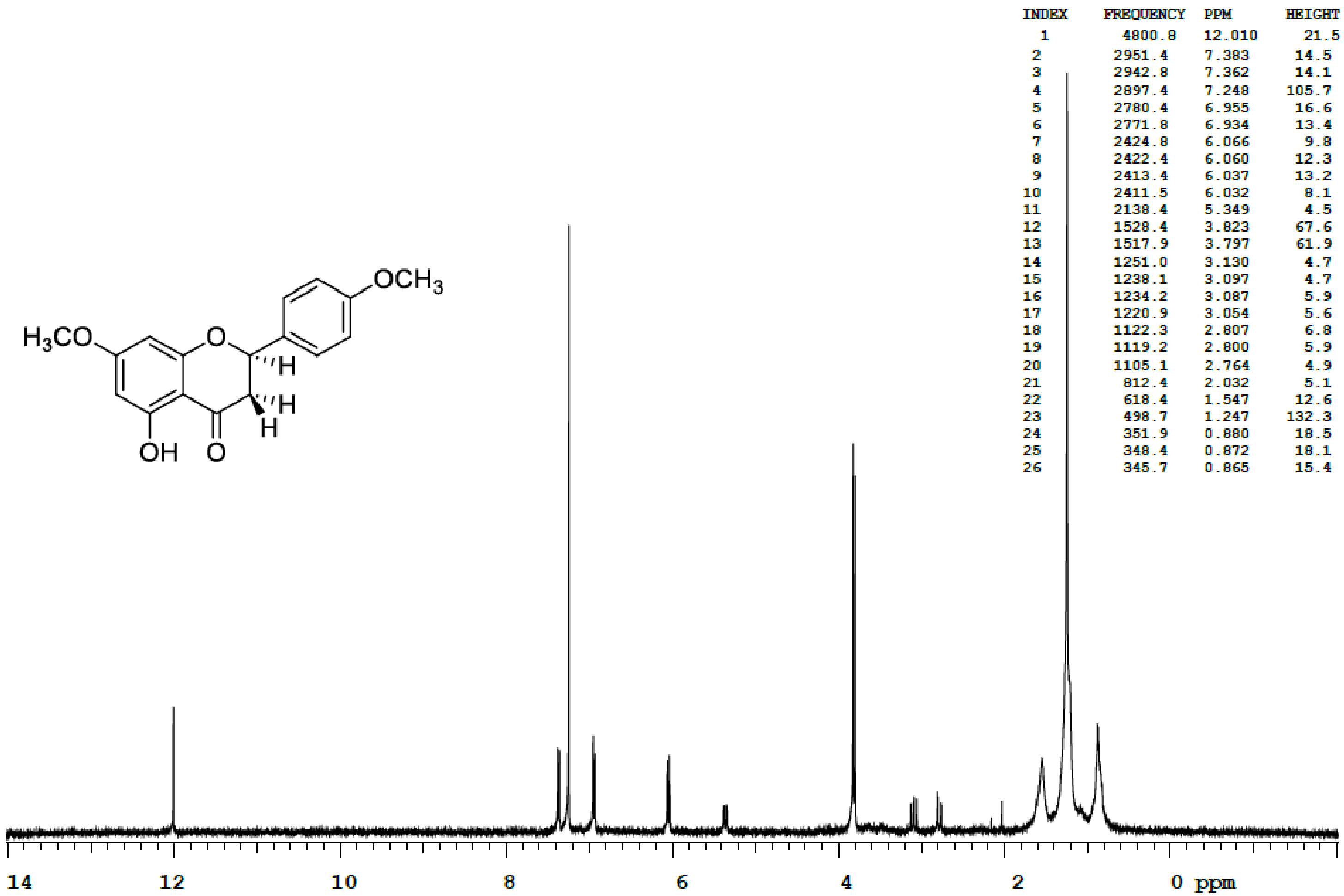


Figure S5.22. High-Resolution Electrospray Ionization Mass spectrum (HR-ESI-MS) of 5-hydroxy-7,4`-dimethoxyflavone; $[M+H]^+$ m/z = 301.9516



INDEX	FREQUENCY	PPM	HEIGHT
1	4800.8	12.010	21.5
2	2951.4	7.383	14.5
3	2942.8	7.362	14.1
4	2897.4	7.248	105.7
5	2780.4	6.955	16.6
6	2771.8	6.934	13.4
7	2424.8	6.066	9.8
8	2422.4	6.060	12.3
9	2413.4	6.037	13.2
10	2411.5	6.032	8.1
11	2138.4	5.349	4.5
12	1528.4	3.823	67.6
13	1517.9	3.797	61.9
14	1251.0	3.130	4.7
15	1238.1	3.097	4.7
16	1234.2	3.087	5.9
17	1220.9	3.054	5.6
18	1122.3	2.807	6.8
19	1119.2	2.800	5.9
20	1105.1	2.764	4.9
21	812.4	2.032	5.1
22	618.4	1.547	12.6
23	498.7	1.247	132.3
24	351.9	0.880	18.5
25	348.4	0.872	18.1
26	345.7	0.865	15.4

Figure S5.23. Proton Nuclear Magnetic Resonance (^{1}H NMR) spectrum of 5-hydroxy-7,4`-dimethoxyflavone ($CDCl_3$, 400 MHz)

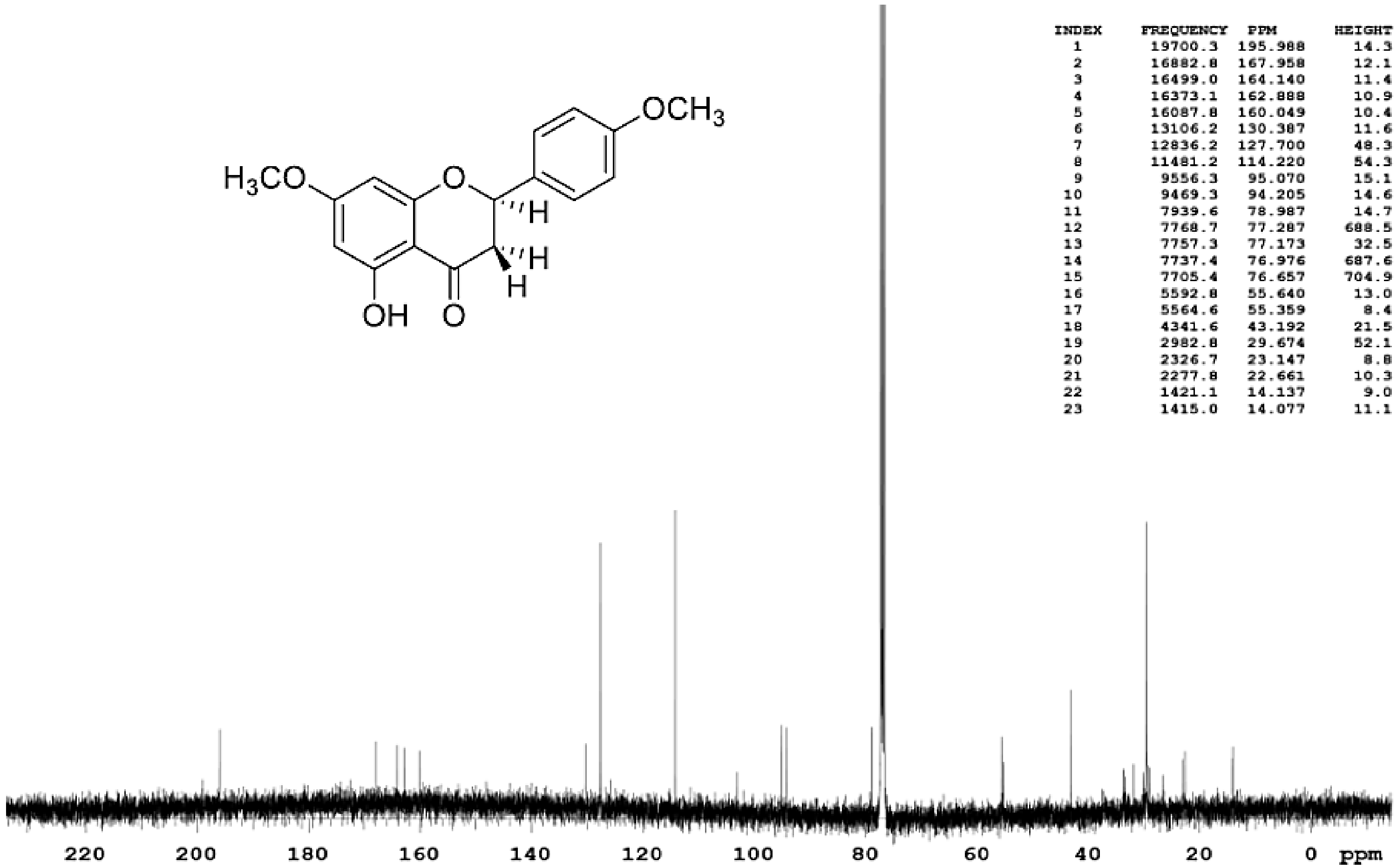


Figure S5.24. Carbon-13 Nuclear Magnetic Resonance (^{13}C NMR) spectrum of 5-hydroxy-7,4`-dimethoxyflavone ($CDCl_3$, 100 MHz)

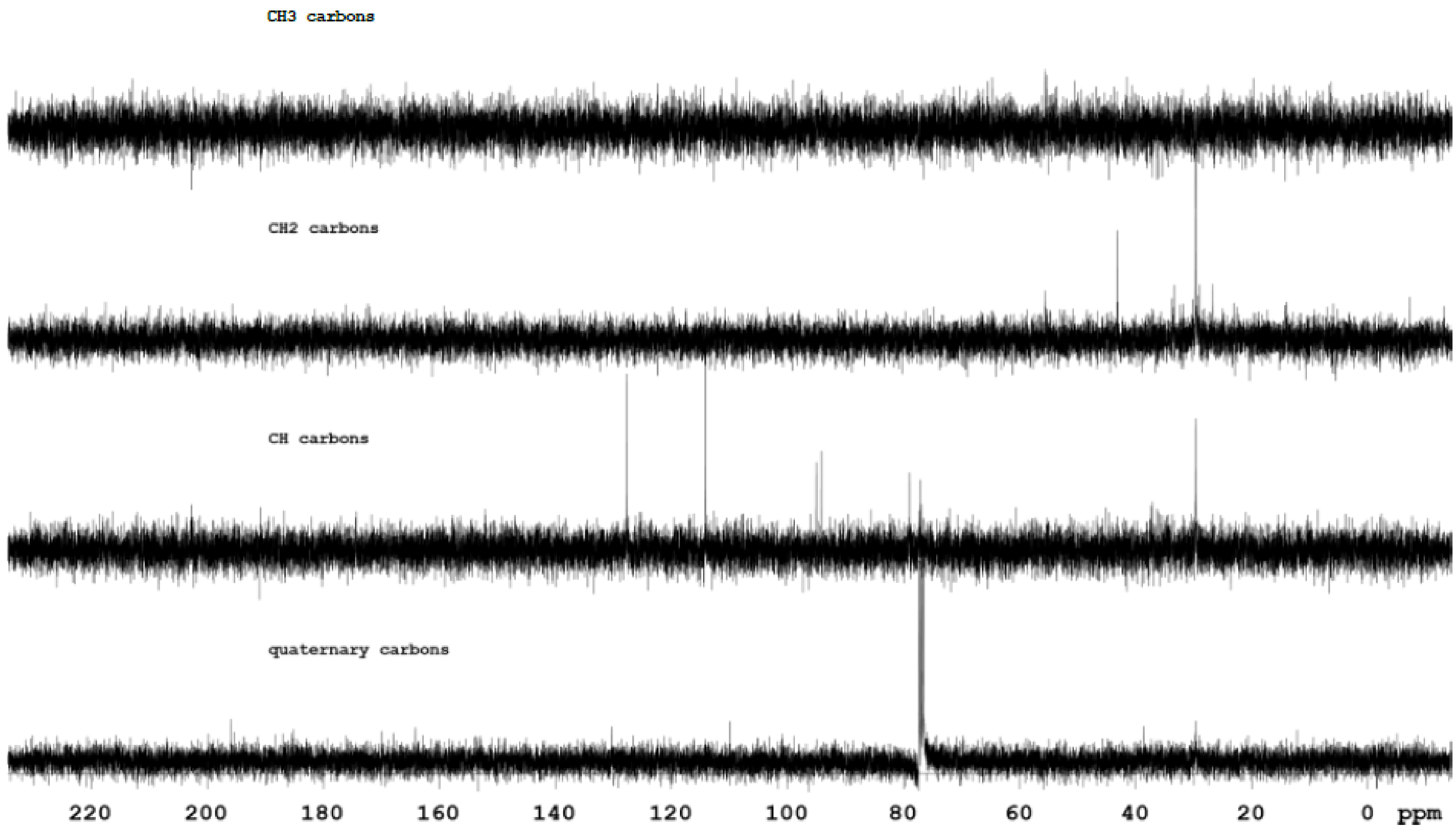


Figure S5.25. Distortionless Enhancement by Polarization Transfer (DEPT) NMR spectra of 5-hydroxy-7,4`-dimethoxyflavone ($CDCl_3$, 100 MHz)