



Identification of New Aromatic Compounds in the New Zealand Manuka Honey by Gas Chromatography-Mass Spectrometry

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Abstract: Analysis of aromatic compounds in the New Zealand manuka honey was carried out by solid phase microextraction followed by gas chromatography-mass spectrometry. A total of 38 compounds were detected. Seven of them such as; 1,4-bis(*x*-methoxyphenyl)-but-2-en-1-one, 1,5-bis(*x*-methoxyphenyl)-pent-3-en-1-one, 1,4-bis(*x*-methoxyphenyl)-1-pentanone, 1,6-bis(*x*-methoxyphenyl)-3-heptene, 1,6-bis(*x*-methoxyphenyl)-hex-2(3 or 4)-en-1-one and 2(3, 4 or 5)-hydroxy-1,6-bis(*x*-methoxyphenyl)-1-hexanone, had never before been identified as natural products. Their structures were deduced from the mass spectral data. Seven other compounds; 2,3-dimethoxynaphthalene, 4-(*x*-methoxyphenyl)-1-phenyl-1-butanone, desoxyanisoin, 2,6-dimethoxybenzoic acid benzyl ester, 4,4'-dimethoxystilbene, 3,3,4,5,5,8-hexamethyl-2,3,5,6-tetrahydro-*s*-indacene-1,7-dione and 1,5-bis(4-methoxyphenyl)-pentane-1,5-dione, were found in honey for the first time. Methyl syringate, ortho-methoxyacetophenone and 3-phenyllactic acid were the most abundant components.

Keywords: Manuka honey, Aromatic compounds, Solid phase microextraction, GC-MS.

Introduction

Manuka honey, produced from the New Zealand's manuka tree (*Leptospermum scoparium*), has been shown to have a high antibacterial activity¹⁻⁵. It is used against the *helicobacter pylori*^{3,6} and as a dressing for the treatment of chronic wounds^{5,7} or ulcers⁸. The antibacterial activity of the honey is generally attributed to the presence of hydrogen peroxide. However, after destruction of this, a residual non-peroxide antibacterial activity was also observed

in some honeys, for example certain active manuka honeys derived from specific localities^{2,9}. Several studies have attempted to give evidence of the principal factor responsible for such activity^{1,10,11}, but none have been successful thus far¹². Inoue *et al.*¹³ demonstrated that manuka honey had a radical scavenging activity for superoxide anion radicals due to the presence of methyl syringate.

In a recent work, we demonstrated the usefulness of solid-phase microextraction (SPME) followed by gas chromatography-mass spectrometry (GC-MS) for the analysis of phenolic and other aromatic constituents in honeys¹⁴. In the present paper we report the results from application of this method to the manuka honey that led to the identification of seven new aromatic compounds. The determination of their structures was made by interpretation of their mass spectral fragmentation patterns. In addition, seven compounds were found in the honey for the first time.

Several aromatic compounds identified had one or more methoxy groups. It is known that methoxylation increases the degree of a product's lipophilicity, improves metabolic stability and consequently the biological activity of that product^{15,16}.

Experimental

Two samples of manuka honey from New Zealand were purchased from a local market. The beekeeper associations supplied honey samples to this market asserting their floral and geographical origins.

Standards were purchased from Sigma-Aldrich Chemie, Buchs, Switzerland, benzoic acid, 4-methoxy-benzoic acid, 2-methoxybenzoic acid, 4-hydroxy-3-methoxybenzoic acid methyl ester, methyl syringate, chrysin and *n*-paraffin mix (C10-C36). The polyacrylate (PA) 85 μm SPME fiber was also obtained from Sigma-Aldrich Chemie.

Solid phase microextraction (SPME)

SPME extraction was carried out as already described¹⁴. Briefly, 1 g of honey was placed into a 10 mL vial. The sample was diluted with 4 mL of distilled water and then 1 g of sodium chloride and 0.2 mL of acetic acid were added. The melange was shaken at room temperature until homogenized. The solution was extracted by immersing the PA fiber into the solution for 30 min at room temperature. The fiber was then removed from the vial and immediately inserted into the GC-MS injector for thermal desorption of the analytes. The PA fiber was initially conditioned for 2 h at 300 °C in a GC injector under a helium flow.

GC-MS analysis

Analyses were performed on a Hewlett Packard 5890 gas chromatograph coupled to a VG Masslab Trio-2 mass spectrometer. GC separations were carried out using a J&W DB-5HT column (30 m x 0.32 mm x 0.1 μm of film thickness). A split/splitless injector was used in the splitless mode (2 min) at a temperature of 300 °C. The oven temperature was programmed as follows: 40 °C for 1 min, increased to 300 °C at 3 °C/min, and held at 300 °C for 5 min. Helium was the carrier gas at a constant flow of 1 mL/min. The mass spectrometer was used in the electron impact mode at 70 eV and with a 1.2 s cycle time in the mass range m/z 45-600.

Compounds were identified by comparison of their mass spectra with those in the NIST mass spectral library and identities were confirmed by co-injections of standards when available. Distributions of the studied compounds were calculated from the peak areas in the total ion chromatograms. Kovats indices were determined from the retention times of an *n*-alkane mixture (C10-C36) analyzed under identical conditions.

Results and Discussion

The SPME extracts from the two samples (**A** and **B**) of manuka honey investigated in this work gave GC-MS chromatograms globally similar. Figure 1 shows the chromatogram of the sample **A** as example. This profile is very distinct from those of the European honeys analysed previously in the same conditions¹⁴. A total of 38 compounds were detected and most of them were identified (Table 1).

Table 1. Distribution (% of total) of aromatic compounds extracted from two samples of manuka honey using SPME/GC-MS

Compounds	No	MW	KI	Distribution, %	
				Manuka A	Manuka B
4-Methylacetophenone	1	134	1111	0.1	0.1
4-Hydroxyacetophenone	2	136	1132	0.2	0.1
Benzoic Acid	3	122	1197	0.5	0.9
4-Methoxybenzaldehyde	4	136	1233	0.3	0.6
<i>Ortho</i> -Methoxyacetophenone	5	150	1277	15.0	7.0
1-Methoxy-4-propyl-benzene	6	150	1281	1.2	5.3
Salicylic Acid	7	138	1323	1.6	1.3
1-(2-Hydroxy-6-methoxyphenyl)-ethanone	8	166	1374	0.1	0.1
4-Methoxybenzoic acid	9	152	1444	0.5	1.2
2-Methoxybenzoic acid	10	152	1478	1.3	0.5
4-Hydroxy-3-methoxybenzoic acid methyl ester	11	182	1500	0.1	0.2
(4-Methoxyphenyl)-acetic acid	12	166	1507	0.5	1.4
3-Phenyllactic acid	13	166	1548	10.2	6.5
2,3-Dimethoxynaphthalene	14	188	1553	0.3	0.1
3,5-Dimethoxybenzoic acid methyl ester	15	196	1558	6.6	0.8
3,5-Dimethoxybenzoic acid	16	182	1644	0.9	1.7
Methyl syringate	17	212	1772	32.3	53.5
Unknown	18		1785	5.7	0.1
Isomer of 18	19		1812	3.8	5.1
Unknown	20		1904	1.1	1.0
4-(<i>x</i> -Methoxyphenyl)-1-phenyl-1-butanone	21	254	1965	0.8	0.4
Isomer of 21	22	254	2112	0.1	0.1
Desoxyanisoin	23	256	2155	1.0	0.6
2,6-Dimethoxybenzoic acid benzyl ester	24	272	2164	0.8	0.1
4,4'-Dimethoxystilbene	25	240	2254	0.3	0.4
3,3,4,5,5,8-Hexamethyl-2,3,5,6-tetrahydro- <i>s</i> -indacene-1,7-dione	26	270	2303	0.5	0.1
Pinostrobin chalcone	27	270	2348	0.1	0.3
1,4-Bis(<i>x</i> -methoxyphenyl)-but-2-en-1-one	28	282	2439	0.2	0.1
Pinocembrin	29	256	2450	0.3	2.4
Tectochrysin	30	268	2528	0.0	0.4
1,5-Bis(4-methoxyphenyl)-pentane-1,5-dione	31	312	2631	3.0	0.9
Chrysin	32	254	2641	0.0	<0.1
1,5-Bis(<i>x</i> -methoxyphenyl)-pent-3-en-1-one	33	296	2652	0.5	0.0
1,4-Bis(<i>x</i> -methoxyphenyl)-1-pentanone	34	298	2656	0.4	1.3
1,6-Bis(<i>x</i> -methoxyphenyl)-3-heptene	35	310	2731	0.8	0.6
1,6-Bis(<i>x</i> -methoxyphenyl)-hex-2(3 or 4)-en-1-one	36	310	2773	0.3	0.1
2(3, 4 or 5)-Hydroxy-1,6-bis(<i>x</i> -methoxyphenyl)-1-hexanone	37	328	2796	6.3	3.3
Isomer of 37	38	328	2825	2.4	1.3

No= Compound number in figure 1, *MW*= Molecular weight, *KI*=Kovats indices

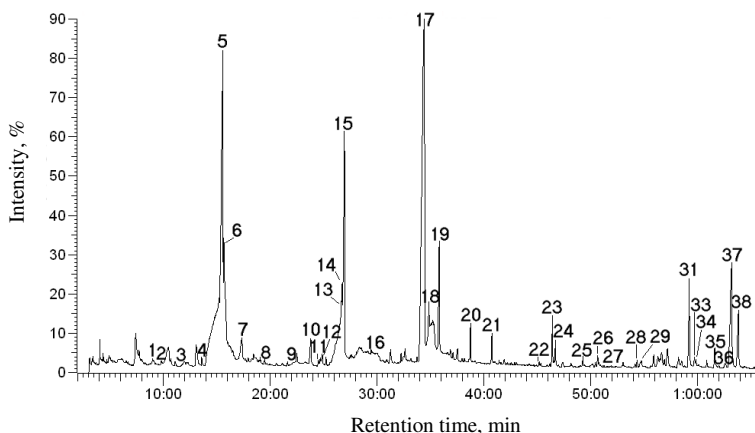


Figure 1. GC-MS chromatogram of aromatic compounds in manuka honey (A), Peak numbers refer to compounds numbered in Table 1

Pinocembrin (**29**) was the only flavonoid detected in sample A. In addition to this compound, tectochrysin (**30**) and chrysin (**32**) were observed in sample B. In comparison with European honeys analysed using the same procedure¹⁴, the manuka honeys are significantly poor in flavonoids; both in variety and relative amount. Several compounds in Table 1 have never been reported to occur in honeys. They are; 2,3-dimethoxynaphthalene (**14**), 4-(*x*-methoxyphenyl)-1-phenyl-1-butanone (**21**), desoxyanisoin (**23**), 2,6-dimethoxybenzoic acid benzyl ester (**24**), 4,4'-dimethoxystilbene (**25**), 3,3,4,5,5,8-hexamethyl-2,3,5,6-tetrahydro-*s*-indacene-1,7-dione (**26**) and 1,5-bis(4-methoxyphenyl)-pentane-1,5-dione (**31**). Their structures are given in Figure 2. Mass spectra of compounds **21** and **24** are not available in the literature and their structures were deduced from interpretation of their mass spectra as shown in Figure 3.

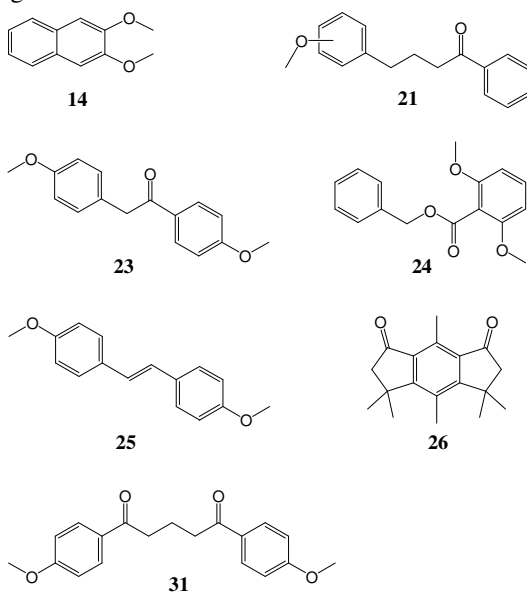


Figure 2. Structures of aromatic compounds reported as honey constituents for the first time

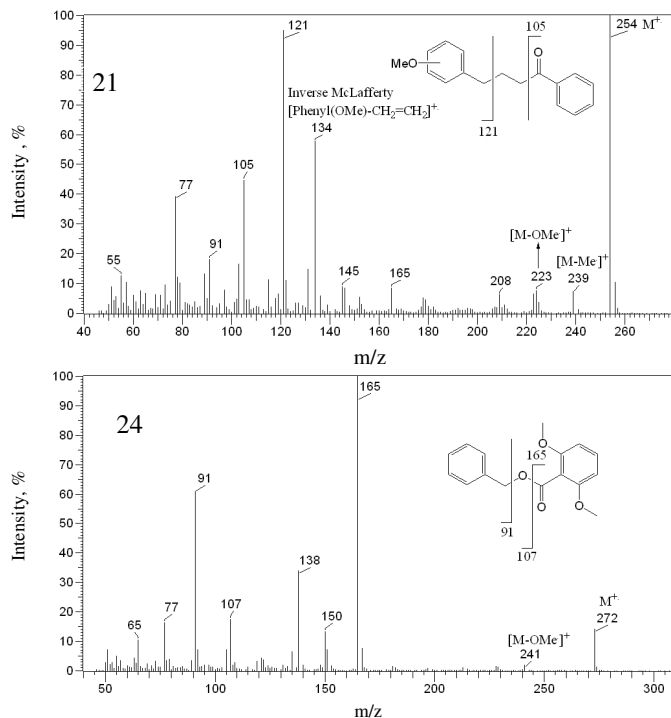
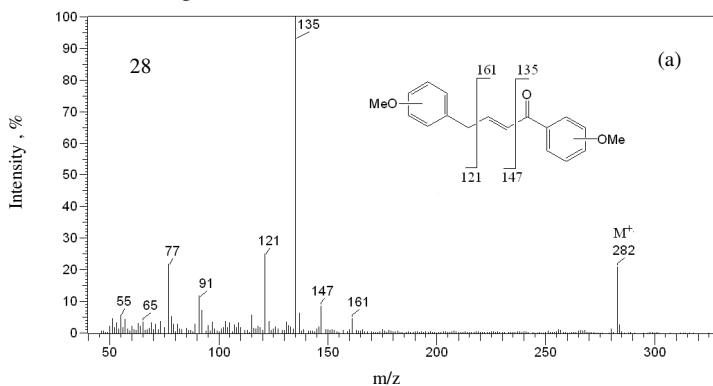
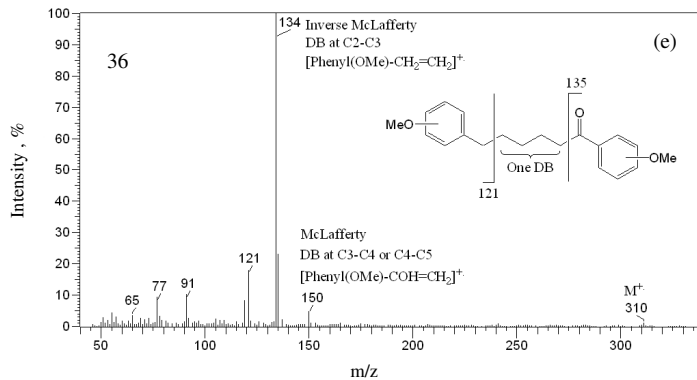
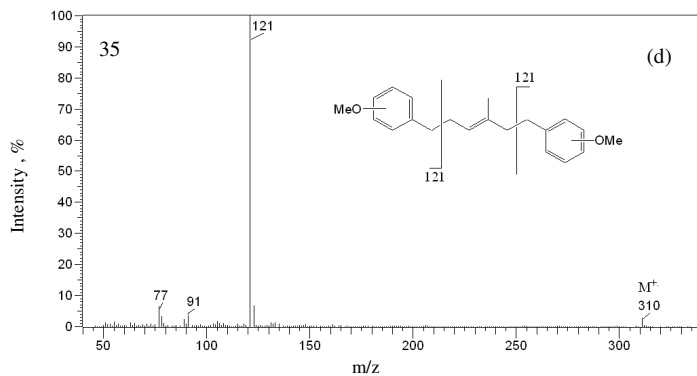
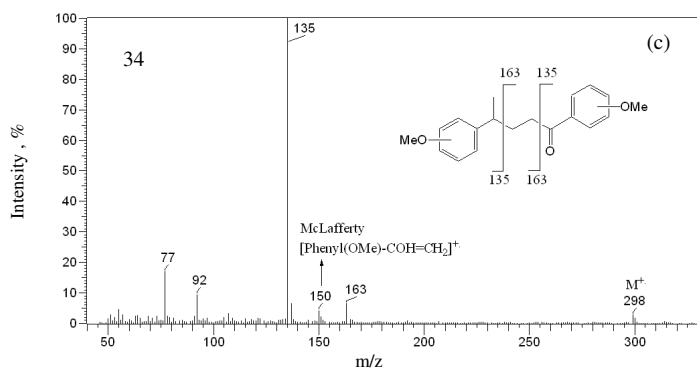
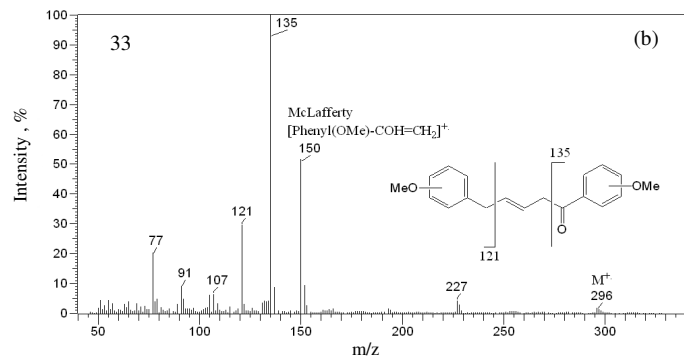


Figure 3. Spectra, structures and important fragments of 14-(*x*-methoxyphenyl)-1-phenyl-1-butanone (**21**) and 2,6-dimethoxybenzoic acid benzyl ester (**24**)

Seven compounds, which are all 1, *n*-bis(*x*-methoxyphenyl)alkane derivatives are tentatively identified on the basis of their fragmentation patterns: 1,4-bis(*x*-methoxyphenyl)-but-2-en-1-one (**28**), 1,5-bis(*x*-methoxyphenyl)-pent-3-en-1-one (**33**), 1,4-bis(*x*-methoxyphenyl)-1-pentanone (**34**), 1,6-bis(*x*-methoxyphenyl)-3-heptene (**35**), 1,6-bis(*x*-methoxyphenyl)-hex-2(3 or 4)-en-1-one (**36**) and 2(3, 4 or 5)-hydroxy-1,6-bis(*x*-methoxyphenyl)-1-hexanone (**37** and **38**). It is probable that the methoxy-substituents of these compounds are in the *para*- position but, in the absence of authentic standards this cannot be ascertained. These are all new compounds hitherto not reported in the chemical literature. Their mass spectra and formation of diagnostic fragments are shown in Figure 4(a-f).





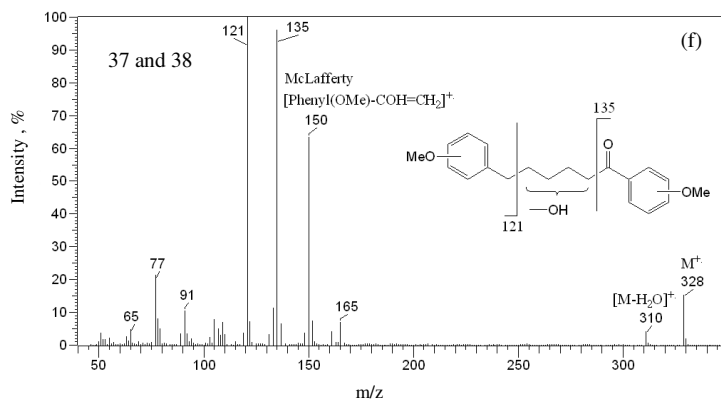


Figure 4. Spectra, structures and important fragments of new aromatic compounds (a) **28**, (b) **33**, (c) **34**, (d) **35**, (e) **36**, (f) **37** and **38** found in manuka honey

Finally, three unidentified components **18**, **19** and **20** were observed in both samples. The major fragments in the mass spectra of these compounds are listed in Table 2. Using liquid-liquid solvent extraction, Wilkins *et al.*¹⁷ found that phenyllactic acid and syringic acid were the most abundant organic compounds in manuka honey. Weston *et al.*⁴ showed moreover, that methyl syringate, which constituted 70% (w/w) of the phenolic fraction, can be considered as a floral marker for this honey. In the present work, methyl syringate (**17**), *ortho*-methoxyacetophenone (**5**) and 3-phenyllactic acid (**13**) were found to be the major components of the SPME extracts: 32.3, 15.0 and 10.2 % of total in sample **A** and 53.5, 7.0 and 6.5% of total in sample **B**, respectively.

Table 2. Mass spectral data of unknown compounds listed in Table 1

Compound No.	Mass (<i>m/z</i>) and Intensity (% , in parentheses) of prominent MS peaks
18	77 (20), 91 (20), 131 (65), 159 (12), 202 (100)
19	55 (26), 77 (14), 91 (12), 102 (22), 115 (22), 131 (45), 158 (13), 202 (100)
20	77 (15), 115 (12), 130 (28), 158 (35), 172 (10), 184 (6), 201 (100)

Rivero-Cruz *et al.*¹⁸ showed the smooth muscle relaxant effect of benzylbenzoates, isolated from *Brickellia veronicifolia*, on guinea-pig ileum. Recently they demonstrated that among a series of spasmolytic benzylbenzoates, some compounds such as 2,4- and 3,4-dimethoxybenzoic acid benzyl ester (isomers of **24**) were the most active inhibitors of the system calmodulin-bovine brain¹⁹.

4,4'-Dimethoxystilbene (**25**) has been reported to be an agent that can decrease cellulite deposits²⁰. This property has been reported also for other substituted *trans*-stilbenes for their ability to inhibit the human tumor necrosis factor alpha-induced (TNF- α) activation of transcription factor nuclear factor kappaB²¹.

Conclusion

The retailers of dietary products sell the manuka honey by claiming that it helps balance hormone levels. It normalizes menstruation and the symptoms of menopause. Isolation of the new compounds identified in this work and study of their biological activity (hormone balance) might provide new insight on the nutritional benefits of manuka honey.

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