

SECONDARY METABOLITES AND PHARMACOLOGY OF *FOENICULUM VULGARE* MILL. SUBSP. *PIPERITUM*

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ABSTRACT

From hexane extract of *Foeniculum vulgare* Mill. Subsp. *piperitum* the fatty acids, hydrocarbons and sterols were identified. The furocoumarins imperatorin, psoralen, bergapten, xanthotoxin and isopimpinellin were isolated from the methylene chloride extract. The flavonoids isorhamnetin 3-*O*- α -rhamnoside, quercetin and kaempferol were isolated from the ethyl acetate extract, whereas quercetin 3-*O*-rutinoside, kaempferol 3-*O*-rutinoside and quercetin 3-*O*- β -glucoside were isolated from the methanol extract. The crude hexane, methylene chloride, ethyl acetate and methanol extracts of this plant showed antinociceptive and anti-inflammatory activity.

Keywords: *Foeniculum vulgare* Mill. Subsp. *piperitum*, coumarins, flavonoids, antinociceptive, antiinflammatory.

RESUMEN

Se realizó la investigación de la constitución química y la actividad biológica de *Foeniculum vulgare* Mill. Subsp. *Piperitum*. En el extracto de *n*-hexano de *Foeniculum vulgare* Mill. Subsp. *Piperitum*. se identificaron ácidos grasos, hidrocarburos y esteroides. Las furocumarinas imperatorina, psoraleno, bergapteno, xantotoxina y isopimpinelina se aislaron del extracto de cloruro de metileno. Los flavonoides 3-*O*- α -ramnosido de isoramnetina, quercetina y kaempferol se aislados del extracto de acetato de etilo, en tanto que la 3-*O*-rutinósido de quercetina, el 3-*O*-rutinósido de kaempferol y 3-*O*- β -glucósido de quercetina se aislaron del extracto metanólico. Los extractos crudos de *n*-hexano, cloruro de metileno, acetato de etilo y metanol presentaron actividad antinociceptiva y anti-inflamatoria.

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INTRODUCTION

Foeniculum vulgare Mill. subsp. *piperitum* (Ucria) Coutinho (Fam. Apiaceae) grows wildy in the Mediterranean coastal strip, Egypt (Tackholm, 1974). It has been widely used as a folk remedy by the native people for treatment of various inflammatory ailments. Chemically, *Foeniculum* species are characterized by the presence of essential oils (Ozbek *et al.*, 2003), sterols (Ivanov *et al.*, 1979), coumarins (El-Khrisy *et al.*, 1980; Kwon, *et al.*, 2002) and flavonoids (Kunzemann *et al.*, 1977; Parejo *et al.*, 2004). Certain bioactivities have been attributed to some *Foeniculum* species; *viz*, antioxidant and antimicrobial activities for *F. vulgare* Mill. aerial parts (Ruberto *et al.*, 2000), anti-inflammatory and analgesic activities for the fruits of the same plant (Eun and Jae, 2004). Volatiles reported from the fruits of *F. vulgare* subsp. *piperitum* comprise anethol, methyl chavicol, fenchone and limonene (Muckensturm *et al.*, 1997), as well as piperitenone and piperitenone oxide (Badoc *et al.*, 1994). Chlorogenic acid (Ishikawa *et al.*, 1999), caffeic acid and cynarin (Scarpati, 1957) have also been isolated from the plant. The purpose of this study was the isolation and identification of chemical constituents from *Foeniculum vulgare* Mill. subsp. *piperitum* growing in Egypt, as well as the evaluation of antinociceptive and antiinflammatory activities of crude extracts of the plant.

PLANT MATERIAL

The aerial parts and fruits of *Foeniculum vulgare* Mill. subsp. *piperitum* were collected from North Western Mediterranean coastal strip near El-Salloum, Egypt in October 2005. The plant material was identified by Prof. Dr. S.A. Kawashty, Phytochemistry and Plant Systematics Department, NRC, Cairo, Egypt. Voucher specimen (F121) is kept in Herbarium, Pharmacognosy Department Faculty of Pharmacy, Cairo University.

EXTRACTION AND ISOLATION

Successive extracts of the air-dried powdered aerial parts (1.8 kg) and fruits (150 g) of *Foeniculum vulgare* Mill. subsp. *piperitum* were prepared using *n*-hexane, methylene chloride, ethyl acetate and methanol, in succession. *n*-Hexane extracts (7 g) from each of the aerial parts and fruits were saponified (El-Said and Amer, 1965) to yield the unsaponifiable matter fractions (USM) (3.5 g and 4 g, respectively), and fatty acids fractions (FA) (2.0 g and 2.5 g, respectively). Methylation of FA was carried out by refluxing in 50 ml absolute methanol and 1.5 ml sulphuric acid for 2 h to give fatty acids methyl esters.

GLC of unsaponifiable matter (USM)

Agilent 6890N gas chromatograph (Hewlett Packard) equipped with FID (Flame Ionization Detector) was used. Column: capillary, 30 m length, 0.53 mm internal diameter, film thickness 0.5 μ m, packed with 5% phenyl - 95% dimethyl polysiloxane; temperature program: 80°C, for 1 min, increased at a rate of 8°C/min.; final temperature, 250°C (kept for 20 min). Injector temperature: 280°C; detector temperature: 300°C; carrier gas: nitrogen, flow-rate 30 ml/min; H₂ flow-rate: 30 ml/min.; air flow-rate, 300ml/min.

GLC of fatty acids methyl esters (FAME)

Agilent 6890N gas chromatograph (Hewlett Packard) equipped with FID was used. Column: capillary, 30 m length, HP-INWAX Polyethylene glycol, 320 μ m internal diameter, 0.25 μ m film thickness; temperature program: 70°C for 2 min., increased at a rate of 4°C/min. till 220°C; carrier gas: nitrogen, flow-rate, 30 ml/min.; air flowrate, 230 ml/min.

Isolation of coumarins from methylene chloride extract:

The solvent-free methylene chloride extract (greenish brown, 37g) was subjected to si-

lica gel CC (75 x 3cm, 150 g) and eluted with benzene and step-gradient benzene/ethyl acetate; 50 ml fractions being collected, monitored by TLC using solvent system benzene/ethyl acetate (8:2) and similar fractions were pooled. Fractions eluted with benzene afforded compound **1** (20 mg), while those eluted with benzene/ethyl acetate (9.5:0.5) afforded compound **2** (18 mg). Fractions eluted with benzene/ethyl acetate (9:1) gave a mixture (1.5 g) of compounds **3,4** and **5** which was subjected to silica gel CC (30 x 2 cm, 25g), eluted successively with benzene, benzene/ethyl acetate (9.5: 0.5) and benzene/ethyl acetate (9:1); yielding 60 subfractions (each, 20 ml). Subfractions eluted with benzene/ethyl acetate (9.5:0.5) afforded compounds **3** (20 mg), and **4** (22 mg), while those eluted with benzene/ ethyl acetate (9:1) afforded compound **5** (15 mg).

Isolation of flavonoids from ethyl acetate extract

The ethyl acetate extract (24 g) was subjected to polyamide 6S CC (100 x 4 cm, 200 g). Elution with water, followed by methanol (40 %, 60 %, 80 % and 100 %, in succession) yielded 130 fractions (200 ml, each) which were monitored by PC using BAW and 15 % AcOH as developing systems; similar fractions being pooled. Fractions eluted with 80% methanol yielded the isolate **6** (23 mg) which was further purified by Sephadex LH-20 CC (30 x 2cm, 40g) using methanol as eluent. Fractions eluted with methanol afforded a mixture of two compounds which were separated and purified by Sephadex LH-20 CC (30x2cm, 40g) using methanol as eluent, to give the compounds **7** (30 mg) and **8** (28 mg).

Isolation of flavonoids from the methanolic extract

The methanolic extract (77 g) was treated as the ethyl acetate extract. The fractions eluted with 40 % MeOH yielded the isolate **9** (22 mg), those eluted with 60% MeOH

afforded the isolate **10** (30 mg), while fractions eluted with 80% MeOH yielded the isolate **11** (20 mg). Each isolate was further purified by Sephadex LH-20 CC, using methanol as eluent.

Evaluation of Bioactivities

The powdered air-dried aerial parts of *F. vulgare* Sups. *piperitum* (1.2 kg) was successively extracted with *n*-hexane, methylene chloride, ethyl acetate and methanol to yield 40, 25, 16 and 70 g of extractives, respectively. 1 mg of the individual extracts was equivalent to 30, 48, 75 and 17.1 mg dry plant, respectively. The extracts were suspended in 7% Tween 80 and biologically tested in different doses. All doses were expressed in terms of extract weight/animal body weight (Berhrens and Kerber, 1953).

Animals and Reference drugs

Adult male albino rats weighing (120-150 g) and adult albino mice of both sex (20-25 g) used in this study, were obtained from the animal-breeding unit of National Research Centre, Cairo. The animals were fed a standard laboratory diet composed of vitamin mix (1 %), mineral mix. (4 %), corn oil (10 %), sucrose (20 %), cellulose (0.2 %), casein (10.5 %) and starch (54.3 %) and allowed free access to water. Animal procedures were in accordance with the recommendations for the proper care and use of laboratory animals. This study was performed according to the international rules and to the guidelines of ethical comity of National Research Centre for experimental animal use. Acetylsalicylic acid (El Nasr Co., Egypt), Ibuprofen (Egyptian International Pharma. Industries Co., Egypt), Carrageenan (BDH, England) and Fluconazole (Pfizer INC., USA) were used in bioactivity testing.

Acute toxicity (LD₅₀)

Acute toxicity (LD₅₀) of each of the extracts from the plant was determined in mice (Finney, 1964). They were subcutaneously

(S.C.) administered in doses ranging from 4 to 13 g/kg body weight. Animals were observed and mortality rates were recorded within the first 24 h after administration. Different extracts assayed up to 5.5 g/kg did not show toxicity. LD₅₀ being: 6.75, 11.0, 6.92 and 15 g/kg for *n*-hexane, methylene chloride, ethyl acetate and methanol extracts, respectively.

Antinociceptive activity

This activity was investigated using the writhing method in mice (Margarita *et al.*, 1995). The tested extracts were administered in

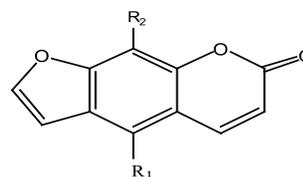
doses ranging 200–2000 mg/kg as well as acetylsalicylic acid (200 mg/kg) as a peripheral antinociceptive reference drug (Margarita *et al.*, 1995) and the control vehicle (7% v/v between 80 in normal saline) were S.C. injected to groups of six animals 15 minutes before the intraperitoneal injection of a freshly prepared acetic acid solution (2% w/v in saline, pH 2.7, 10 ml/kg) and transferred immediately to individual observation cages and the assay was observed over a period of 25 min. The number of abdominal writhes was counted and the percentage of protection was expressed as the following:

$$\frac{C - T}{T} \times 100$$

Where: C = mean writhing of the control group; T = mean writhing of the treated group

Anti-inflammatory activity

Anti-inflammatory activity was studied using Carrageenan-induced rat's paw edema (Winter *et al.*, 1963). Groups of 18 h-fasted male rats (110–130 g, 6 animals each) were orally-dosed with either one of the tested extracts from *Foeniculum vulgare* 1h before induction of Carrageenan foot paw edema by subplanter injection of 0.05 ml of 1% suspension of Carrageenan in saline into the planter tissue of one hind paw. An equal volume of saline was injected into the other hind paw and served as control. Four hours later, the animals were sacrificed and the hind paws were rapidly amputated at the tibiotarsal joint and weighed (Margarita *et al.*, 1995; El-Azzouny *et al.*, 1995). The average weight of edema was estimated for the treated as well as the control group and the percentage inhibition of weight of edema was also evaluated. Ibuprofen (35 mg/kg) (Makhlouf and Maklad, 2004) was employed as a standard.

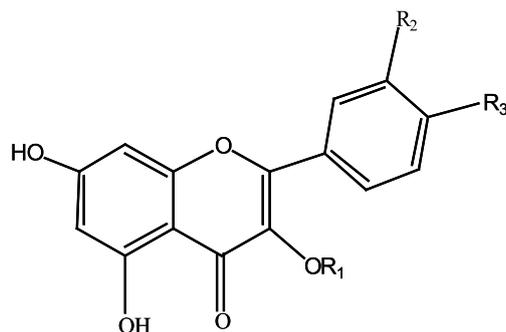


Compounds **1-5**

- 1**, R₁=H, R₂=
- 2**, R₁=H, R₂= H
- 3**, R₁=OCH₃, R₂= H
- 4**, R₁=H, R₂= OCH₃
- 5**, R₁= OCH₃, R₂= OCH₃

IDENTIFICATION: SPECTRAL DATA

GLC analysis of FAME (Table 1) revealed 15 compounds in the aerial parts and 14 compounds in the fruits. In the aerial parts and fruits, saturated fatty acids represent 68.43 and 75.03%, respectively; the major being: palmitic, undecanoic and myristic acids, while the unsaturated acids represent 22.01% and 20.32%, respectively; the major being pentadecadienoic and pentadecenoic acids. GLC of USM revealed 27 compounds in the aerial parts, as well as the fruits (Table 2). Hydrocarbons (56.51,

Compounds **6-11****6**, R₁= rhamnosyl, R₂=OCH₃, R₃ = OH**7**, R₁=H, R₂ = OH, R₃ = OH**8**, R₁=H, R₂=H, R₃ = OH**9**, R₁= rutinoyl, R₂ = OH, R₃ = OH**10**, R₁= rutinoyl, R₂ = H, R₃ = OH**11**, R₁=glucosyl, R₂ =OH, R₃ = OH**Fig. 1:** Structures of isolated compounds**Table 1:** Fatty acids identified as methyl esters (%) in the aerial parts and fruits of *Foeniculum vulgare* Mill. subsp. *piperitum*.

Identified Compounds	Rt(min.)	Aerial Parts	Fruits
Capric acid	9.01	2.83	3.23
Undecanoic acid	10.44	18.21	20.09
Lauric acid	11.76	2.13	2.45
Myristic acid	12.80	10.51	11.20
Pentadecanoic acid	14.27	1.79	2.10
Pentadecenoic acid	15.11	7.33	7.68
Pentadecadienoic acid	16.32	9.29	10.91
Palmitic acid	17.30	31.51	33.47
Stearic acid	18.41	0.69	0.77
Oleic acid	20.85	1.55	0.40
Linoleic acid	22.86	0.43	-
Linolenic acid	23.52	0.65	0.38
Arachidic acid	25.53	0.76	1.31
Behenic acid	26.48	-	0.41
Erucic acid	27.28	0.87	1.33
Tetracosenoic acid	29.68	1.89	-
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Total unsaturated fatty acids (%)	-
Total saturated fatty acids (%)	-	22.01	20.32
Total identified fatty acids (%)	-	68.43	75.03
		90.44	95.35

Table 2: Hydrocarbons and sterols identified in the USM of the aerial parts and fruits of *F. vulgare* Mill. subsp. *piperitum*.

Identified Compounds	R _t (min.)	Aerial Parts	Fruits
<i>n</i> -Decane	3.99	3.75	4.29
<i>n</i> -Dodecane	4.91	2.16	2.84
<i>n</i> -Tridecane	6.72	0.88	0.34
<i>n</i> -Pentadecane	10.98	0.47	1.09
<i>n</i> -Hexadecane	11.61	0.49	0.65
<i>n</i> -Heptadecane	12.40	1.64	1.44
<i>n</i> -Octadecane	13.76	1.71	2.19
<i>n</i> -Nonadecane	14.04	-	1.11
<i>n</i> -Eicosane	15.63	5.89	10.43
<i>n</i> -Monocosane	16.06	2.72	1.88
<i>n</i> -Docosane	16.48	0.96	1.39
<i>n</i> -Tricosane	16.64	-	2.37
<i>n</i> -Tetracosane	17.44	0.45	0.89
<i>n</i> -Pentacosane	18.43	2.25	4.59
<i>n</i> -Hexacosane	18.76	2.86	5.51
<i>n</i> -Heptacosane	19.51	0.41	0.27
<i>n</i> -Octacosane	19.92	0.37	0.44
<i>n</i> -Nonacosane	20.36	0.79	1.76
<i>n</i> -Triacontane	20.60	0.53	1.35
<i>n</i> -Monotriacontane	21.43	0.48	0.89
<i>n</i> -Dotriacontane	23.16	2.09	2.28
<i>n</i> -Tetratriacontane	24.48	4.52	3.34
<i>n</i> -Hexatriacontane	25.81	10.27	-
<i>n</i> -Octatriacontane	26.48	7.91	2.83
<i>n</i> -Tetracontane	27.43	1.71	0.84
Cholesterol	29.68	5.52	5.95
β -Sitosterol	31.80	5.52	5.95
Campesterol	32.53	3.33	4.04
Stigmasterol	34.00	14.86	19.04
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Total identified hydrocarbons (%)	-	56.51	54.17
Total identified sterols (%)	-	25.42	29.87
Total identified compounds (%)	-	81.91	84.04

54.17%), cholesterol (1.71, 0.84%), β-sitosterol (5.52, 5.95%), campesterol (3.33, 4.04%) and stigmasterol (14.86, 19.04%) were identified in the aerial parts as well as the fruits, respectively.

Imperatorin (1): UV λ_{max} (MeOH): 262 and 300 nm. EI-MS: *m/z* 270 [M⁺, base peak, C₁₆H₁₄O₄]; *m/z* 202 [M⁺ -C₅H₉]; *m/z*

174 [M⁺ -C₅H₉ - CO]; *m/z* 146 [M⁺ -C₅H₉ - CO - CO]; ¹H-NMR (CDCl₃, 270 MHz): δ 7.76 (H-4, d, *J* = 9.5 Hz), δ 7.73, (H-5, s), δ 7.34 (H-7, d, *J* = 2.5 Hz), δ 6.79 (H-6, d, *J* = 2.5 Hz), δ 6.35 (H-3, d, *J* = 9.5 Hz), δ 5.58 (H, t, *J* = 6.5 Hz), δ 4.99 (two protons for OCH₂, d, *J* = 7.1 Hz), δ 1.69 and δ 1.71 (singlets for the two methyls).

Psoralen (2): UV λ_{\max} (MeOH): 246 and 293 nm. EI-MS: m/z 186 [M^+ , base peak, $C_{11}H_6O_3$] which loses three carbon monoxide molecules, successively, to give the fragments m/z : 158, 130 and 102. 1H -NMR ($CDCl_3$, 270 MHz): δ 7.48 (H-4, d, $J = 9.5$ Hz), δ 7.36 (H-5, s), δ 7.35 (H-9, s), δ 7.13 (H-7, d, $J = 2.5$ Hz), δ 6.50 (H-6, d, $J = 2.5$), δ 6.06 (H-3, d, $J = 9.5$ Hz).

Bergapten (3): UV λ_{\max} (MeOH): 249, 259, 267 and 310 nm. EI-MS: m/z 216 [M^+ , $C_{12}H_8O_3$], m/z 201 [$M^+ - CH_3$], followed by four successive expulsions of carbon monoxide to give the fragments m/z : 173, 145, 117 and 89. 1H -NMR ($CDCl_3$, 270 MHz): δ 8.13 (H-4, d, $J = 9.7$ Hz), δ 7.57 (H-7, d, $J = 2.5$ Hz), δ 7.08 (H-9, s), δ 7.00 (H-6, d, $J = 2.5$ Hz), δ 6.25 (H-3, d, $J = 9.7$ Hz), δ 4.25 (singlet, 5-OCH₃).

Xanthotoxin (4): UV λ_{\max} (MeOH): 240 and 298 nm. EI-MS: m/z 216 [M^+ , $C_{12}H_8O_3$], m/z 201 [$M^+ - CH_3$], followed by four successive expulsions of carbon monoxide to give the fragments: m/z 173, 145, 117 and 89. 1H -NMR ($CDCl_3$, 270 MHz): δ 7.76 (H-4, d, $J = 9.5$ Hz), δ 7.67 (H-7, d, $J = 2.1$ Hz), δ 7.32 (H-5, s), δ 6.80 (H-6, d, $J = 2.1$ Hz), δ 6.35 (H-3, d, $J = 9.5$ Hz), δ 4.26 (singlet, 9-OCH₃).

Isopimpinellin (5): UV λ_{\max} (MeOH): 248, 267 and 309 nm. EI-MS: m/z 246 [M^+ , $C_{13}H_{10}O_5$], m/z 231 [$M^+ - CH_3$], followed by two successive expulsions of carbon monoxide to give the fragments m/z 203 and 175, and loss of methyl group to give the fragment m/z 160. 1H -NMR ($CDCl_3$, 300 MHz): δ 8.14 (H-4, d, $J = 9.5$ Hz), δ 7.64 (H-7, d, $J = 2.5$ Hz), δ 7.06 (H-6, d, $J = 2.5$ Hz), δ 6.36 (H-3, d, $J = 9.5$ Hz) and two singlets at δ 4.13 and 4.09 for 5- and 9-OCH₃.

Isorhamnetin 3-O- α -L-rhamnoside (6): EI-MS: m/z 316 [Aglycone+H]⁺, +ve ESI-MS: m/z 462 [$M+H^+$, $C_{22}H_{22}O_{11}$]. 1H -NMR ($DMSO-d_6$, 270 MHz): δ 12.63 (5-OH, s), δ 7.27 (H-2', d, *meta*-coupling with H-6', $J = 2.0$ Hz), δ 7.25 (H-6' d, $J = 8.8$ Hz), δ 6.85 (H-5', d, $J = 8.3$ Hz), δ 6.32 (H-8, d, $J = 2.0$ Hz), δ 6.13 (H-6, d, $J = 2.0$ Hz), δ 5.24 (broad singlet for anomeric proton of rhamnose),

δ 0.95 (CH₃ methyl rhamnosyl protons, d, $J = 5.3$ Hz), δ 3.95 (OCH₃ at position 3', sharp singlet) (Chang *et al.*, 1998).

Quercetin (7): UV spectral data suggested a flavonol type with free hydroxyl groups at positions 3, 5, 7, 3' and 4'. **EI-MS:** m/z 303 [$M+1$]⁺, $C_{15}H_{10}O_7$.

Kaempferol (8): UV spectral data suggested a flavonol type with free hydroxyl groups at positions 3, 5, 7 and 4'. **EI-MS:** m/z 285 [$M-1$]⁻, $C_{15}H_{10}O_6$.

Quercetin 3-O-rutinoside (9): 1H -NMR: δ 7.55 (H-2' and H-6' overlapping protons, d, $J = 7.6$ Hz), δ 6.85 (H-5', d, $J = 8.0$ Hz), δ 6.38 (H-8, d, $J = 1.8$ Hz), δ 6.19 (H-6, d, $J = 1.8$ Hz), δ 5.32 (anomeric proton of glucose, $J = 8.1$ Hz), δ 4.37 (anomeric proton of rhamnose, broad singlet), δ 0.99 (CH₃-rhamnosyl, d, $J = 5.4$ Hz).

Kaempferol 3-O-rutinoside (10): UV spectra with shift reagents indicated a flavonol type with free hydroxyl groups at positions 5, 7 and 4', while that at position 3 being substituted (Mabry *et al.*, 1970). Complete acid hydrolysis afforded glucose, rhamnose and kaempferol, identified by CoPC in comparison with authentic samples. 1H -NMR ($DMSO-d_6$, 270 MHz): δ 7.61 (H-2' and H-6', d, $J = 8.5$ Hz) δ 6.77 (H-3' and H-5', d, $J = 8.5$ Hz), δ 6.20 (H-8, d, $J = 1.8$ Hz), δ 6.0 (H-6, d, $J = 1.8$ Hz), δ 5.21 (anomeric proton of glucose, d, $J = 7.6$ Hz), δ 4.40 (anomeric proton of rhamnose, broad singlet) δ 1.04 (methyl rhamnosyl protons, d, $J = 6.1$ Hz). **ESI-MS:** m/z 593 [$M-1$]⁻, $C_{27}H_{30}O_{15}$.

Quercetin 3-O- β -glucoside (11): UV spectral data suggested a flavonol type with free hydroxyl groups at positions 5, 7, 3' and 4', while being substituted at position 3. Complete acid hydrolysis afforded glucose. 1H -NMR: δ 7.66 (H-6', d, $J = 8.6$ Hz), δ 7.55 (H-2', d, $J = 2.0$ Hz), δ 6.84 (H-5', d, $J = 6.6$ Hz), δ 6.38 (H-8, d, $J = 1.8$ Hz), δ 6.17 (H-6, d, $J = 1.8$ Hz), δ 5.34 (anomeric proton of glucose, d, $J = 7.5$ Hz). **ESI-MS:** m/z 464 [$M+1$]⁺, $C_{21}H_{20}O_{12}$.

Table 3: Effect of extracts of the aerial parts *F. vulgare* Mill. subsp. *piperitum* on acetic acid-induced writhing in mice.

Animal groups	Dose (mg/kg)	Number of writhing ^b X ± S.E.	Protection %
Vehicle	-	29.60 ± 1.45 ^a	-
Acetylsalicylic acid	200	6.00 ± 1.18*	79.73
<i>n</i> -Hexane extract	600	21.00 ± 0.82 ^{*a}	29.05
	700	11.17 ± 0.87 ^{*a}	62.26
Methylene chloride extract	400	18.00 ± 0.78 ^{*a}	39.19
	500	11.67 ± 0.76 ^{*a}	62.27
Ethyl acetate extract	700	13.00 ± 0.78 ^{*a}	56.08
	800	7.00 ± 0.87*	76.35
Methanol extract	1500	7.17 ± 0.76*	75.78
	2000	4.50 ± 0.94*	84.79

*Significantly different from control value at $p < 0.05$

^aSignificantly different from acetylsalicylic acid (200 mg / kg) value at $p < 0.05$

^bEach value represents the mean (% increase in weight of paw edema) ± s.e. of the number of animals in each group (n = 6).

Table 4: Anti-inflammatory activity of extracts of the aerial parts of *F. vulgare* Mill. subsp. *piperitum*

Animal groups	Dose (mg/kg)	% Increase in weight of paw edema (g) ^b X ± S.E.	Protetion %
Vehicle	-	65.08 ± 1.69 ^a	-
Ibuprofen	35	31.09 ± 1.84*	52.23
<i>n</i> -Hexane extract	600	58.90 ± 2.32 ^{*a}	9.49
	700	52.51 ± 2.76 ^{*a}	19.31
Methylene chloride extract	400	56.74 ± 2.39 ^{*a}	14.35
	500	43.64 ± 1.92 ^{*a}	32.94
Ethyl acetate extract	700	60.04 ± 2.84 ^a	7.74
	800	58.08 ± 3.00 ^{*a}	10.76
Methanol Extract	1500	46.76 ± 1.99 ^{*a}	28.15
	2000	34.63 ± 2.86*	46.79

*Significantly different from control value at $p < 0.05$.

^aSignificantly different from ibuprofen (35 mg / kg) value at $p < 0.05$.

^bEach value represents the mean (% increase in weight of paw edema) ± s.e. of the number of animals in each group (n = 6).

Bioactivities

The methanolic extract of the aerial parts of *Foeniculum vulgare* subsp. *piperitum* exhibited the highest antinociceptive activity at a dose level of 2000 mg/kg (Table 3), while the activity exhibited by the ethyl acetate extract was at (800 mg/kg). On the other hand, *n*-hexane extract (700 mg/kg) and methylene chloride extract (500 mg/kg)

exhibited similar antinociceptive activities, being less than that of acetylsalicylic acid (200 mg/kg). Data presented in Table 4 revealed that the extracts under investigation exhibited significant anti-inflammatory activity. The methanolic extract possessed the highest activity, where it significantly decreased the weight of edema induced by carrageenan in the rat paw at dose levels of 1500 and 2000 mg/kg, it exerted a pro-

protective effect of 28 and 47%, respectively, compared to the control value, while ibuprofen (35 mg/kg), used as a reference drug, exhibited a protective effect of 52.23%. The present study showed that the four extracts of the aerial parts of the plant exhibited antinociceptive activity. Moreover, the potent antinociceptive activity expressed by the methanolic (2000 mg/kg) and ethyl acetate (800 mg/kg) extracts can be attributed to the content of phenolic constituents (Paulino *et al.*, 2003). Also, the four extracts under investigation exhibited a significant anti-inflammatory activity. The highest anti-inflammatory activity of methanolic extract (2000 mg/kg) of the aerial parts of the plant can possibly be attributed to the presence of phenolic content. In conclusion, traditional application of *F. vulgare* Mill subsp. *piperitum* could be beneficial in the management of different inflammatory disease cases. Our current findings suggested that *F. vulgare* Mill subsp. *piperi-*

tum contains a high amount of flavonoids, which exhibited a strong potency towards suppressing inflammation, as evidenced by our *in vivo* models.

CONCLUSION

Foeniculum vulgare Mill. Subsp. *Piperitum* to contain fatty acids, hydrocarbons and sterols in the *n*-hexane extract. Furocoumarins; imperatorin, psoralen, bergapten, xanthotoxin and isopimpinellin were isolated from the methylene chloride extract. Flavonoids; isorhamnetin 3-*O*- α -rhamnoside, quercetin and kaempferol were isolated from the ethyl acetate extract, whereas quercetin 3-*O*-rutinoside, kaempferol 3-*O*-rutinoside and quercetin 3-*O*- β -glucoside were isolated from the methanol extract. The crude extracts exhibited significant antinociceptive and anti-inflammatory activity.

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