# PHYTOCHEMICAL ASSESSMENT ON N-HEXANE EXTRACT AND FRACTIONS OF Marsilea crenata Presl. LEAVES THROUGH GC-MS

# ANALISIS FITOKIMIA EKSTRAK N-HEKSANA DAN FRAKSI DAUN Marsilea crenata Presl. DENGAN GC-MS

Burhan Ma'arif<sup>1\*</sup>, Mangestuti Agil<sup>1</sup> and Hening Laswati<sup>2</sup>

<sup>1</sup>Department of Pharmacognocy and Phytochemistry, Faculty of Pharmacy, Airlangga University, Surabaya, Indonesia

<sup>2</sup>Department of Physical Medicine and Rehabilitation, Faculty of Medicine, Airlangga University, Surabaya, Indonesia

#### ABSTRACT

Estrogen deficiency causes various health problems in postmenopausal women, including osteoporosis. Phytoestrogen emerged as a potential alternative of estrogen with minimum side effects. Green clover (Marsilea crenata Presl.) is a typical plant in East Java which suspected contains estrogen-like substances. The aim of this research was to report the phytochemical properties of M. crenata using GC-MS as a preliminary study. M. crenata leaves were dried and extracted with n-hexane, then separated using vacuum column chromatography to get four fractions, after that the n-hexane extract and four fractions were identified with GC-MS. The results of GC-MS analysis showed some compounds contained in M. crenata leaves like monoterpenoid, diterpenoid, fatty acid compounds, and other unknown compounds. The results obtained in this research indicated a promising potential of M. crenata as medicinal plants, especially as antiosteoporotic agent.

Keywords: Marsilea crenata Presl., phytochemical, antiosteoporosis, GC-MS

#### ABSTRAK

Defisiensi estrogen menyebabkan berbagai masalah kesehatan pada wanita pascamenopause, salah satunya osteoporosis. Fitoestrogen muncul sebagai alternatif yang potensial pengganti estrogen dengan efek samping minimal. Semanggi (Marsilea crenata Presl.) merupakan tanaman khas Jawa Timur yang diduga mengandung senyawa dengan fungsi mirip estrogen (estrogen-like substances). Tujuan dari penelitian ini untuk melaporkan kandungan senyawa kimia dalam M. crenata menggunakan GC-MS sebagai studi pendahuluan. Daun M. crenata diekstraksi dengan n-heksana lalu dipisahkan menggunakan kromatografi kolom vakum, setelah itu ekstrak n-heksana dan empat fraksi yang didapatkan diidentifikasi menggunakan GC-MS. Hasil analisis GC-MS menunjukkan beberapa golongan senyawa yang terkandung dalam daun M. crenata seperti monoterpenoid, diterpenoid, senyawa asam lemak, dan senyawa lain yang belum diketahui. Hasil yang diperoleh dalam penelitian ini menunjukkan potensi menjanjikan M. crenata sebagai tanaman obat, khususnya sebagai antiosteoporosis.

Kata kunci: Marsilea crenata Presl., fitokimia, antiosteoporosis, GC-MS

#### **INTRODUCTION**

*Marsilea. crenata* (Figure 1) is a unique plant that grows in East Java, Indonesia. Its leaves are widely used as an ingredient for traditional food (Afriastini, 2003; Nurjanah *et al.* 2012). *M. crenata* is a kind of ferns which is usually grown in the aquatic environment (Afriastini, 2003; Steenis, 1975). Different from other Marsilea plants such as *Marsilea minuta* Linn., *Marsilea quadrifolia*,

Corresponding Author : Burhan Ma'arif Email : burhan.maarif@gmail.com *Marsilea rajasthansis* Gupta, and *Marsilea drummondii*, the phytochemical properties of *Marsilea crenata* Presl. has not been confirmed yet. Whereas these plants has great potential as a source of food or as medicinal plants and very easy to grow (Yacoeb *et al.* 2010).

Some research shows that *M. crenata* has activity for prevention of osteoporosis (Laswati, 2011). Using radioimmunoassay (RIA), estradiollike compound concentrations in *M. crenata* leaves were detected quite high (Laswati, 2011), and it has potentials in delaying the increment

#### Burhan Ma'arif

of imbalance bone remodeling process in postmenopausal women (Laswati, 2011; Yang, *et al.* 2012; Ososki *et al.* 2003).



Figure 1. M. crenata

This study was aimed to report the phytochemical properties of *M. crenata* using GC-MS as a preliminary study to determine the antiosteoporotic activity of *M. crenata*. GC-MS is a powerful technique used for many applications which has very high sensitivity and specificity and also can save more time (Elezabeth *et al.* 2014; Kumar *et al.* 2014; Bai *et al.* 2014).

#### METHODOLOGY Chemical and Equipment Plant Material

The leaves of *M. crenata* were collected in Surabaya, East Java, Indonesia. The collected plant materials were washed thoroughly with water, then dried carefully under shade, at room temperature so as to retain their fresh green color, and also to prevent decomposition of active compounds (Sarker *et al.* 2006).

## **Extraction and Fractination**

N-hexane and ethyl acetate as solvent and mobile phase, maseration chamber, vacum column chromatography with diameter 6,5 cm and hight 6,5 cm, TLC plate silica gel 60 F254 Merck, silica gel 60 G Merck,  $H_2SO_4$  10% reagent, and 254 nm also 366 nm UV lamp.

#### Method

#### **Extraction and Fractination**

The leaves of *M. crenata* were extracted with *n*-hexane through maseration method. The *n*-hexane extract of *M. crenata* leaves then separated using vacuum column chromatography on 70 g silica gel 60 G Merck as the stationary phase, and *n*-hexane : ethyl acetate as the mobile phase at gradient eluation.

### Analysis with GC-MS

The *n*-hexane extract and fractions were analyzed using an HP gas chromatography model 6890 coupled to an MSD detector. Analysts were separated on an HP-5MS capillary column Agilent 19091S-433 (30.0m X 250µL X 0.25µL) with constant flow of carrier gas 1.0mL/min and initial pressure 11.6 psi, and then applying the following temperature program: 120°C for 2min, 120-200°C at 10°C/min, and 200°C for 10min. Mass detector conditions were acquisition mode scans, source temperature are minimum 230°C and maximum 250°C. Carrier gas was helium at 1.0 mL/min. The tentative identification of volatile components was achieved by comparing the mass spectra with the data system library (NIST02) and (Wiley275), supported by retention index data, which were compared with available literature retention indices.

## **RESULTS AND DISCUSSION**

A total of 1.5 kg green powder was extracted with *n*-hexane to produce 22 g extract. After that, 3.5 g of *n*-hexane extract was separated using vacuum column chromatography on 70 g silica gel 60 G Merck as the stationary phase, and *n*-hexane : ethyl acetate as the mobile phase at gradient eluation. The separation process resulted in four fractions, there are fraction 1 (2,236 mg), fraction 2 (1,100 mg), fraction 3 (403 mg), and fraction 4 (96 mg). The separation process of nhexane extract into several fractions using dry column vacuum chromatography needs to be done before analysis by GC-MS. This separation serves to improve the detection capabilities of GC-MS. Some peak that cannot be detected in the extract can be detected in fractions. This is because the compounds in each fraction, which are separated and identified by GC-MS, is not as much as the compounds in the extract.

The results of GC-MS analysis of the *n*hexane extract and four fractions of M. crenata leaves show some compounds contained as listed in Table I. In Figure 2, we can see the Total Ion Chromatogram (TIC) of *n*-hexane extract and four fractions of *M. crenata* leaves. Mass spectra of each peak of extracts and fractions are further identified and matched with a data system library NIST02 and Wiley275. Each peak is identified with match factor should more than 85%. For examples, see figure 3. Figure 3 is mass spectra of peak from *n*-hexane extract that has a retention time (a) 11,996 and (b) 14,725, its appropriate with mass spectra of (a) neophytadiene with match factor 99%, and (b) n-hexadecanoic acid with match factor 98%.

#### PHYTOCHEMICAL ASSESSMENT ON N-HEXANE EXTRACT



Figure 2. TIC of (a) *n*-hexane extract, (b) fraction 1, (c) fraction 2, (d) fraction 3, and (e) fraction 4 of *M*. *crenata* leaves.



Figure 3. Comparison of spectrums through mass finder. (a) neophytadiene and (b) *n*-hexadecanoic acid

In general, the result of GC-MS show that the *n*-hexane extract of *M. crenata* leaves predicted contains many volatile group compounds such as monoterpenoid and diterpenoid, and fatty acid group compounds. The *n*-hexane extract and fractions of *M. crenata* leaves contain some diterpenoid compounds such as neophytadiene that have activity as antipyretic, analgesic, antiinflammatory, antimicrobial, and antioxidant; dihydroactinidiolide that have function as fragnance or pheromon in animal; phytol that have activity as antiinflammatory, antimicrobial, antioxidant, and diuretic; fatty acid compounds such as palmitic acid that have activity as antioxidant, hypocholesterolemic, antiandrogenic, and 5-Alpha reductase inhibitor; oleic acid that have activity as cancer preventive and antiandrogenic; elaidic acid that have activity as

anticholesterol and lower LDL; and also other unknown compounds. Some of predicted compounds are impossible found in extract and fractions of natural product. For example, methyl ester form of fatty acid is normally detected after saponification process in laboratory work.

Through correlation analysis between TIC GC-MS and literature study, it can be concluded that palmitic acid is estrogen-like substance in *M. crenata* leaves that suspected have a role as antiosteoporotic agent, especially in improvement of osteogenesis (Kim *et al.* 2013). The mechanism of palmitic acid in improving osteogenesis occurs through enhancement of osteoblast's transcription factor in differentiation process such as runt-related transcription factor-2 (runx-2) and osterix (osx) significantly (Kim *et al.* 2013).

Function					[] [] [] [] [] [] [] [] [] [] [] [] [] [	riavout component, itagnance, prietonone ni ammai, muu				Increases plasma cholesterylester transfer protein activit which lowers HDL cholesterol		Antipyretic, analgesic, anti-inflammatory, antimicrobial, antioxidant			Cancer preventive, anemiagenic, insectifuge, antiandroger dermatitigenic	2 Antioxidant, flavor, hypocholesterolemic pesticide, 5-alpha reductase inhibitor		Antioxidant, hypocholesterolemic, nematicide, pesticide, lubr antiandrogenic, flavor, hemolytic, 5-Alpha reductase inhibitor	Antiinflammatory, hypocholesterolemic, cancer preventiv hepatoprotective, nematicide, insectifuge, antihistaminic antiarthritic, anticoronary, antieczemic, antiacne,	5-Alpha reductase inhibitor, antiandrogenic
MM						C <sub>11</sub> H <sub>16</sub> O				C <sub>18</sub> H <sub>34</sub> O		C <sub>20</sub> H <sub>38</sub>			C <sub>18</sub> H <sub>34</sub> O	C17H₃4O		C16H3ZO	C19H3zO	
MF						86				95		66			96	97		68	94	
Compounds	Unknown	Unknown	Unknown	Unknown	2(4H)-Benzofuranone, 5,6,7,7a-tetrahydro-4,4,7a-trimethyl-,	(R)- / dihydroactinidiolide	Unknown	Unknown	Unknown	9-Octadecenoic acid, (E)- / Octadec-9-enoic acid / Elaidic acid	Unknown	Neophytadiene / 7,11,15-trimethyl-3-methylidenehexadec-1- ene	Unknown Thalwarm	OIINIIOWII	Oleic Acid	Hexadecanoic acid, methyl ester	Unknown	n-Hexadecanoic acid / Palmitic acid	9,12,15-Octadecatrienoic acid, methyl ester, (Z,Z,Z)-	Unknown
% Area	0,54	0,91	0,18	0,41		1,60	0,53	4,73	0,77	0,27	1,10	17,56	3,38 5.61	TO'C	1,65	7,64	4,48	33,30	2,09	8,24
Rt	2,46	7,76	7,91	8,29		8,55	8,89	10,35	10,42	10,56	11,28	12,00	12,39	7/171	13,06	13,52	14,35	14,72	18,19	18,75
Peak	H	0	ŝ	4		2	9	7	œ	6	10	11	12	3	14	15	16	17	18	19
Sample										,	pe	e Extr	uexə	H	u					

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ardinec		746	1 39	2 4. Hentadianal (FF).	7	C-H-D	I awen I DI antiovidant
	• 64	5.31	1,66	Unknown	:		
	( m	5,61	1,13	Unknown			
	4	7,76	2,08	3-Buten-2-one, 4-(2,6,6-trimethyl-1-cyclohexen-1-yl)-	93	C13H200	Fragnance, antioxidant, inhibition proliferation of cancer cell
	'n	8,55	0,94	2(4H)-Benzofuranone, 5,6,7,7a-tetrahydro-4,4,7a- trimethyl-, (R)- / dihydroactinidiolide	68	C <sub>11</sub> H <sub>16</sub> O <sub>2</sub>	Flavour component, fragnance, pheromone in animal, inducing gen expression
	9	8,87	0,86	Unknown			
	7	8,99	0,44	Unknown			
	00	9,34	1,13	Unknown			
	6	10,11	3,91	Unknown			
	10	10,27	0,71	9-Octadecenoic acid, (E)- / Octadec-9-enoic acid / Elaidic acid	95	C18H34O2	Increases plasma cholesterylester transfer protein activity which lowers HDL cholesterol
	=	10,47	1,25	Cyclohexane, 1-(1,5-dimethylhexyl)-4-(4-methylpentyl)-	06		
r uc	12	10,61	0,22	Unknown			
ito.	13	11,28	2,51	6-Octadecenoic acid, (Z)-	94	C18H34O2	Cancer preventive, insectifuge
мя	14	12,00	9,60	Neophytadiene / 7,11,15-trimethyl-3-methylidenehexadec- 1-ene	66	C20H38	Antipyretic, analgesic, anti-inflammatory, antimicrobial, antioxidant
	15 16	12,40 12,72	3,36	Unknown Unknown			
	17	13,06	2,63	Oleic Acid	95	C <sub>18</sub> H <sub>34</sub> O <sub>2</sub>	Cancer preventive, anemiagenic, insectifuge, antiandrogenic, dermatitigenic
	18	13,52	20,49	Hexadecanoic acid, methyl ester	67	C <sub>17</sub> H <sub>34</sub> O <sub>2</sub>	Antioxidant, flavor, hypocholesterolemic pesticide, 5-alpha reductase inhibitor
	19	14,74	F 6,08	Unknown			1
	20	14,98	4,67	Cyclopentadecane	86		
	21	17,97	3,86	10,13-Octadecadienoic acid, methyl ester	66		
	22	18,18	12.92	9.12.15-Octadecatrienoic acid. methyl ester. (Z.Z.2)-	8	CosH200	Antiinflammatory, hypocholesterolemic, cancer preventive, hepatoprotective, nematicide, insectifuge, antihistaminic, antiarthritic, anticoronary, antieczemic,
							antiacne, 5. Aluha reductace inhibitore antiandroeenio
	23	18,75	16,10	Unknown			ט־ליוענום ו בעתרנסוכ ווווועיועי ו שווישואי עפרייי

81

# PHYTOCHEMICAL ASSESSMENT ON N-HEXANE EXTRACT

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	1	7,03	4,43	T etradecanoic acid / Myristic acid	86	C14H28O2	Antioxidant, cancer preventive, cosmetic, hypercholesterolemic, nematicide, lubricant
	61	7,32	0,25	Unknown			:
	8	8,55	0,83	2(4H)-Benzofuranone, 5,6,7,7a-tetrahydro-4,4,7a- trimethyl-	93	C11H1602	Flavour component, fragnance, pheromone in animal, inducing gen expression
	4	9,18	0,39	Unknown			
2	s	10,07	1,47	Unknown			
; u	9	11,22	0,68	Unknown			
oitae	2	12,01	24,05	Neophytadiene / 7,11,15-trimethyl-3-methylidenehexadec- 1-me	66	C20H38	Antipyretic, analgesic, and anti-inflammatory, antimized is a surfaction
чu	00	12,41	3,52	Unknown			מוורוווורו ללומון מוורללו למוור
	6	12,73	9,85	Phytol	<u> 6</u>	C20H40O	Antimicrobial, anticancer, cancer preventive, diuretic
	10	14,37	4,92	9,12,15-Octadecatrien-1-ol, (Z,Z,Z)-	68		
	11	14,78	45,18	n-Hexadecanoic acid / Palmitic acid	66	C16H32O2	Antioxidant, hypocholesterolemic, nematicide, pesticide, lubricant, antiandrogenic, flavor, hemolytic,
	;			:			5-Alpha reductase inhibitor
	12	18,78	4,43	Unknown			
		2,29	0,92	Unknown			
	61	7,75	0,66	Unknown			
	m	7,91	0,61	Unknown			
	4	9,72	9,57	Unknown			
	w	10,35	13,00	Unknown			
	9	11,23	5,27	Unknown			
	2	11,74	5,28	Unknown			
E ao	80	12,00	12,69	Neophytadiene / 7,11,15-trimethyl-3-methylidenehexadec- 1-ene	66	C20H38	Antipyretic, analgesic, and anti-inflammatory, antimicrobial, antioxidant
beri							Antiinflammatory, nematicide, jnsectifuge, hypocholesterolemic, cancer preventive,
r -	6	12,40	1,89	9,12-Octadecadienoic acid (Z,Z)- / Linoleic acid	87	C <sub>18</sub> H <sub>32</sub> O <sub>2</sub>	hepatoprotective, antihistaminic, antiacne, antiarthritic, antieczemic, 5-alpha reductase inhibitor,
							antiandrogenic, anticoronary
	10	12,72	2,96	Unknown			
	11	14,34	26,03	9,12,15-Octadecatrien-1-ol, [Z,Z,Z]-	68		
	12	14,75	21,12	n-Hexadecanoic acid / Palmitic acid	95	C16H32O2	Antioxidant, hypocholesterolemic, nematicide, pesticide, lubricant, antiandrogenic, flavor, hemolytic,
							5-Alpha reductase inhibitor

# Burhan Ma'arif

pesticide, lubricant, antiandrogenic, flavor, hemolytic, Increases plasma cholesterylester transfer protein Antioxidant, hypocholesterolemic, nematicide, activity which lowers HDL cholesterol Germination inhibitory, anticancer, immunosuppressive activity 5-Alpha reductase inhibitor Function C18H34O2 C11H16O3 C16H32O2 ΜW MF 32 86 87 86 92 97 9-Octadecenoic acid, (E)- / Octadec-9-enoic acid / Elaidic 11,13-dimethyl-12-tetradecen-1-ol acetate n-Hexadecanoic acid / Palmitic acid 9,12,15-0ctadecatrien-1-ol, (Z,Z,Z)-Tetrahydroxycyclopentadienone Compounds (-)-Loliolide Unknown Unknown Unknown Unknown Unknown Unknown acid % Area 13,13 12,46 39,07 10,01 1,09 0,33 1,99 17,21 0,14 0,85 0,82 2,88 14,78 10,35 10,62 11,76 12,73 14,43 8,55 9,78 12,01 7,76 9,51 2,39 꿃 Peak 9 11 10 12 307 4 2 9 L  $^{\circ\circ}$ Sample Fraction 4

PHYTOCHEMICAL ASSESSMENT ON N-HEXANE EXTRACT

#### CONCLUSIONS

Through GC-MS analysis, it is known that *M. crenata* leaves contain many volatile group compounds such as monoterpenoid, diterpenoid, and fatty acid group compounds that have various activity. Palmitic acid, one of fatty acid contained in *M. crenata* leaves, is suspected to have a role as antiosteoporotic agent, especially in improvement of osteogenesis. Therefore, further research is need to be done to prove it's activity.

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