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Bioactive Compounds Profile of Alkaloid on Elaeocarpus sphaericus Schum Seeds by Liquid Chromatography-Mass Spectrometry

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ABSTRACT

Elaeocarpus sphaericus Schum with the local name genitri has medicinal plant properties. Therefore, this study aims to identify the diversity of alkaloid compounds in genitri seeds using the Liquid Chromatography-Mass Spectrometry (LC-MS) method. The Extract Purification through Solid Phase Extraction (SPE) was conducted using C18 Sep-Pak. Based on the LCMS results, several alkaloids were obtained in the genital seeds with different compositions namely, Elaeokanine C 0.90%, (+)-elaeocarpine 1.59%, Elaeocarpenine 1.88%, Isoelaeocarpine 1.36%, Grandisine B 0.86%, Isoelaeocarpiline 1.14%, Grandisine D 1.51%, Elaeocarpidine 1.03%, Isoelaeocarpicine 1.89%, Grandisine F 1.39%, Grandisine A 0.89%, Grandisine C 0.98%, Grandisine E 1.14%, Habbemine A 1.24%, Habbemine B 1.46%, Grandisine G 0.85%. Furthermore, the bioavailability of these compounds was determined through Phytochemical Analysis.

Keywords: alkaloids, compounds profile, Elaeocarpus sphaericus, LCMS

1. INTRODUCTION

Biodiversity of plants can thrive in Indonesia due to geographical conditions, especially with a tropical climate, and they have medicinal properties. Previous studies showed that plants have various active compounds used as medicine. Elaeocarpus sphaericus Schum medicinal plant is one of the 360 species of the Elaeocarpaceae family that can grow in tropical and sub-tropical areas. This species is a local endemic plant that has not been fully utilized, and it is used as shade, firewood, seeds are used for jewelry. Several studies showed that genitri can be used as a medicinal plant [1-3] due to the complex compounds [4-5]. Furthermore, the chemical compounds have pharmacological activities that can be used in the treatment of diseases [3],[6]. The ethanolic extract of E. ganitrus seed of 250 mg/kg and 500 mg/kg (p.o) affected the normal kidney tissue structure of rabbits fed with cholesterol [7]. Meanwhile, E. tectorius seed extract 40.06 mg has antioxidant and antimicrobial properties [8]. E. tonkinesis leaf extract has anti-influenza ability [5]. One of the chemical compounds in plants are alkaloids, and they contain heterocyclic nitrogen atoms [9-10] that are synthesized in various plants. Alkaloids in pure compounds have the characteristics of colorless, nonvolatile, crystalline, tend to taste bitter [11], and they have various physiological effects on the human body [12-13]. Qualitative and quantitative methods are conducted to identify the phytochemical properties of natural materials [14]. Liquid Chromatograph Mass Spectrometry (LC-MS) is one of these analytical methods that combines the separation capabilities of liquid chromatography (LC) against a target compound. Furthermore, this compound is obtained from the mixture of mass spectroscopic capabilities (mass spectrometry/MS) which can read 4 spectrum of the separated target compound [15]. This study aims to analyze the compounds of the alkaloid group of Elaeocarpus sphaericus Schum seeds using the LC-MS method.

2. METHOD

2.1. Materials

Dried genetri seeds were obtained from a forest area in Magetan, East Java, Indonesia.



2.2. Sample Preparation

The seeds were washed and dried using a microwave at 60 °C, then mashed using a blender, and 100-250 g were taken after being \$200th. Furthermore, the seed powder was dissolved in 95% methanol in a ratio of 1:5 (sample: methanol), and 2e solution was stirred until homogeneous. It was left to stand for 24 hours at a cold temperature in a closed bottle, and the filtration process was further conducted to obtain the filtrate and dregs using an Erlenmeyer vacuum filter. The process was repeated three times by dissolving with methanol until the dregs were obtained. The combined filtrate was evaporated from the methanol to obtain a semi-viscous extract using half of the previous solution.

2.3. Extract Dilution

A vortex was used to dissolve the extract with methanol to a concentration below 100 ppm until a homogeneous solution was obtained. Furthermore, the glids were separated by centrifuging at a speed of 8000 rpm for 10 minutes, and the obtained supernatant was used in the next stage.

2.4. Protein Precipitation

In the centrifuge tube, 2 ml of supernatant extract was added to 3 ml of acetonitrile acidified with 0.2% formic acid. The centrifugation was conducted at a speed of 8000 rpm for 30 seconds to obtain another supernatant.

2.4. Purification with SPE (Solid Phase Extraction)

The supernatant was put into a Sep-Pak C18 Cartridge (1 ml, 100 mg) which had been conditioned with 1 ml 80:20 (acetonitrile: water), and about 0.5 ml of the solution was accommodated with 1 ml of the sample into the Sep – Pak column. Furthermore, 0.5 ml of 80:20 acetonitrile/water solution was added into the Sep – Pak column, and 0.5 ml of the solution that comes out was accommodated. Then, 0.25 ml of 200 mM

ammonium formate was added in a 50:50 solution of acetonitrile/methanol to the Sep-Pak column. The obtained 0.5 ml was added with 0.2 ml of 25:75 acetonitrile: buffer (25 mM ammonium formate pH 4.5), and the solution was used for injection in LCMS. Furthermore, the filtration was conducted using a membrane filter of 0.45 m cellulose acetate, and Table 1 showed the LCMS specifications.

Table 1. Specifications of LC-MS

LC-MS apparatus	Shimadzu LC-MS – 8040
model	LC/MS
Column	Shimadzu Shim Pack FC-ODS
	(2 mm x 150 mm, 3 μm)
Injection volume	1 <i>µ</i> I
Capillary voltage	3,0 kV
Column	35 ℃
temperature	
Mobile phase	Isocratic
mode	
Flow rate	0,5 ml/min
Sampling cone	23,0 V
60lvent	Ethanol 95%
MS focused ion	lo type [M]+
mode	
Collison energy	5,0 V
Desolvation gas	60 ml/hr
flow	6
Desolvation	350 °C
temperature	
Fragmentation	Low energy CID
method	
Ionization	ESI
Scanning	0,6 sec/scan (mz: 10-1000)
Source	100 °C
temperature	
Run time	80 minutes
LCMS method	LCMS method

3. RESULTS

Table 2 showed the analysis of genitri seed alkaloids using the LC-MS method, with specifications. In addition, it showed the composition of each compound based on the analysis of the alkaloid group using the LC-MS method of genitri seeds.

Table 2. Results of LC-MS analysis of compounds from the seed alkaloid group Elaeocarpus sphaericus Schum

Peak number	RT (min)	Similarity index (%)	Curve area	Composition (%)	Compound Result	
Hullibei	(11111)	IIIuex (76)		(/0)	Analysis	Structure
15	6,409	92	605,71886	0,90496	elaeokanine C	
16	8,034	92	1068,8228	1,59685	(+)-elaeocarpine	



Peak number	RT (min)	Similarity index (%)	Curve area	Composition (%)	Compound Result	
number	(11111)	muex (76)		(70)	Analysis	Structure
17	8,041	92	1261,48853	1,88470	elaeocarpenine	76
18	8,045	92	911,78578	1,36223	isoelaeocarpine	
19	8,165	92	578,55872	0,86438	grandisine B	
20	8,304	92	760,64328	1,13642	isoelaeocarpiline	
21	8,322	92	1009,42285	1,50810	grandisine D	
22	8,74	92	690,42761	1,03152	elaeocarpidine	
23	9,732	92	1515,8679 7	2,26475	naringenin	NO OH
24	9,805	92	1261,78616	1,88514	isoelaeocarpicine	Di HO Mano H
25	9,847	92	927,65515	1,38594	grandisine F	H ₂ (v) H ₃ (v) H ₄ (v) H ₄ (v) H ₅ (v

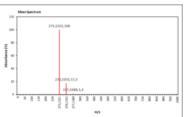


Peak number	RT (mains)	Similarity	Curve area	Composition	Compound Result	
number	(min)	index (%)		(%)	Analysis	Structure
26	9,851	92	596,84565	0,89170	grandisine A	
27	9,855	92	659,15586	0,98480	grandisine C	HO THE THE PART OF
28	9,859	92	760,66456	1,13645	grandisine E	To the state of th
29	9,97	92	831,55537	1,24237	habbemine A	HO NH
30	9,973	92	976,88686	1,45949	habbemine B	HO N H
34	10,52	92	568,08828	0,84874	grandisine G	THE THE PART OF TH

4. DISCUSSION

The analysis results of the alkaloid group using the LC-MS method of genitri seeds showed different compounds, namely Elaeokanine C, (+)-elaeocarpine, Elaeocarpenine, Isoelaeocarpine, Grandisine B, Isoelaeocarpiline, Grandisine D, Elaeocarpidine,

Isoelaeocarpicine, Grandisine F, Grandisine A, Grandisine C, Grandisine E, Habbemine A, Habbemine B, Grandisine G. Alkaloids are one of the active compounds in genitri seeds, and the results of the LCMS analysis showed that the highest content was Isoelaeocarpicine compound, which is about 1.89%. Figure 1 showed the structural analysis of Isoelaeocarpicine, with a chemical formula of $C_{16}H_{21}NO_3$ and a molecular weight of 275.3480.



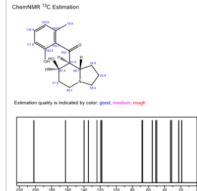


Figure 1. Isoelaeocarpicine, one of the alkaloid compounds in genitri seeds, indicated by the adsorption value and chemical structure

Liquid Chromatography-Mass Spectrometry (LC-MS) is an analytical technique that combines the physical separation capabilities of liquid chromatography with the detection specificity of mass spectrometry [16]. It separates the components of the sample and the charged ions are detected by a mass spectrometer, and the result can be used to provide information about the molecular weight, structure, identity, and quantity of certain sample components [17]. Furthermore, the compounds are separated based on their relative interaction with the chemical layer of the particles (stationary phase) and the elution of the solvent through the column (mobile phase). This method can analyze a wider range of components, such as thermolabile compounds, hon polarity or high molecular mass, and proteins. The font in the row header should be bold and you can use the style available from the style palette.

Alkaloids are one of the secondary metabolites in genitri seeds, and they have medicinal plants properties [18-21]. Furthermore, alkaloids in Uncaria rhynchophylla have been identified as an anti-dengue grus [20]. These compounds can inhibit the growth of Staphyloco gramma sarregions and Pseudomonas aeruginosa bacteria in Callistemon citrinus and Vernonia adoensis leaves [19]. Previous studies showed that alkaloid compounds have immunomodulatory properties [22].

5. CONCLUSION

A phytochemical analysis is an important factor used to determine the components of chemical compounds in plants. Furthermore, Liquid Chromatography-Mass Spectrometry (LC-MS) was used to analyze the alkaloid compounds of genitri seeds, and different compounds were obtained, namely Elaeokanine C 0.90%, (+)-elaeocarpine 1.59%, Elaeocarpenine 1.88%, Isoelaeocarpine 1.36%, Grandisine B 0.86%, Isoelaeocarpiline 1.14%, Grandisine D 1.51%,

Elaeocarpidine 1.03%, Isoelaeocarpicine 1.89%, Grandisine F 1.39%, Grandisine A 0.89%, Grandisine C 0.98%, Grandisine E 1.14%, Habbemine A 1.24%, Habbemine B 1.46%, Grandisine G 0.85%.

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