

Phytochemical and Acute Toxicity Studies of Ethanol Extract from Pedada (*Sonneratia caseolaris*) Fruit Flour (PFF)

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Abstract— Studies on the phytochemical and acute toxicity of pedada fruit flour (PFF) were carried out. In acute toxicity test, oral administration of the extract to Swiss albino mice at four levels dose, i.e. 0, 10.50; 15.75 and 21.00 g/kg body weight. Phytochemical analysis of the ethanol extract of PFF showed the presence of saponins, sapogenins, terpenoids, flavonoids, tannins, polyphenols. Phytochemicals such as alkaloids were not detected. The results of acute toxicity (LD₅₀) showed that the ethanol extract of PFF in mice was found more than 21.00 g/kg body weight. It could be concluded that the PFF belongs to relatively less dangerous category 'non-toxic' and 'safe' for food products.

Keywords— pedada fruit flour (PFF), phytochemical, acute toxicity

I. INTRODUCTION

Pedada fruit is a fruit produced from one of mangrove tree species *Sonneratia caseolaris*. The ripened fruits have an appealing flavor and taste [1]. Some research reported that pedada fruit was used as folk medicine, analgesic, anti-inflammatory properties [4], and antibacterial [16],[19]. Bioactive compound found in pedada fruit are steroids, triterpenoids, flavonoids, saponins, and tannins [4],[16],[25]. The fruits have two flavonoid types (luteolin and luteolin 7-O-β-glucoside) which function as an antioxidant [17],[25].

The flour product from pedada was commonly consumed by mangrove farmers in Wonorejo village, Surabaya, and was not known in modern community. In the other hand research about bioactive compounds from pedada fruit flour (PFF) are limited. Previous research had reported that the PFF decreased of blood glucose levels [11] and total plasma cholesterol, LDL-c, triglyceride, but not affecting the HDL-c [10], so it is potential for food ingredients, but research phytochemical on PFF has not been done. The phytochemical screening is qualitative analysis based on color reactions and/ or precipitation [27]. Furthermore the pedada fruit is not toxic [6], but be sure of its safety after processed into the flour so that acute toxicity analysis was required. In most acute toxicity test, a single dose of a test

substance is given to an animal. One measure of the acute toxicity is the lethal dose 50 (LD₅₀) which is the dose of a substance that kills 50 percent of the animals tested [5],[24].

Therefore the PFF will be applied as substitution in food product, because it has pasting properties if mixed with another flour or starch [12], so this study was investigated to determine the phytochemicals and acute toxicity of ethanol PFF extract from *Sonneratia caseolaris*.

II. MATERIALS AND METHODS

A. Chemicals

Dragendorff reagent, Meyer reagent, Wagner reagent, ferric chloride 2%, chloroform, n-hexane, ethyl acetate, anisaldehyde sulfuric acid, hydrochloric acid, sodium chloride, ammonium hydroxide 28%, methanol, butanol.

B. Extract preparation

Fresh Pedada fruits of *Sonneratia caseolaris* (50-55g; ± 2 months) were obtained from Wonorejo village, Surabaya, Indonesia. Pedada fruit flour (PFF) was processed using method by Jariyah *et al.* [10]. The PFF was macerated with ethanol 96% (1:3) in a stoppered container for a defined period with frequent agitation until soluble matter is dissolved. It was filtered through muslin cloth on beaker

glass, the filtrate from ethanol extract of PFF were evaporated at 40°C, 60 rpm, stored in the refrigerator. The ethanol extract of PFF (EPFF) is ready to analyze.

C. Phytochemical analysis

The presence of alkaloids, saponins, saponinins, terpenoids, flavonoids, tannins and polyphenols were determined qualitatively by using methods from YAPI Houphouet *et al.* [27] and Studiawan [22]. The procedure of phytochemicals analysis are as follows :

1) Alkaloids test

5 ml of 2N HCl was added to 0.3 g EPFF, then heated on water bath for 2-3 minutes and stirred. After cooling, 0.3 g NaCl was added and filtered. 5 ml of 2N HCl was added to the filtrate and divided into three part (A, B, C). Meyer reagent was added to part A, Wagner reagent for part B and Dragendorff reagent for part C. The presence of precipitation indicated the presence of alkaloids. Part C was analyzed by thin layer chromatography (TLC) with Kiesel gel GF 254 as stationary phase, ethyl acetate : methanol : water (6:4:2) as mobile phase, Dragendorff reagent as appearance of spot. The appearance of orange color after spraying Dragendorff reagent indicated the presence of alkaloids.

2) Saponins test

10 ml distilled water was added to EPFF 0.3 g, shaken for 30 seconds, and the foam height is measured. The foam height greater than 3 cm and stable for 30 min at a above the liquid surface indicated the presence of saponins

- Saponinins / terpenoids test

5 ml 2N HCl was added to 0.5 g EPFF, boiled and covered with funnel containing a damp cotton for 2 h to hydrolyze saponins. After cooling, neutralized with ammonia, extracted with 3 ml n-hexane for 3 times, evaporated until it reached 0.5 ml. Analysis saponinins by TLC method is the same as alkaloids with n-hexane : ethyl acetate (4:1) as mobile phase, anisaldehyde sulfuric acid as the appearance of spot. The red purple color indicated the presence of saponinins/terpenoids.

- Flavonoids test :

3 ml n-hexane was added to 0.3 g EPFF, extracted and filtered. The residue was diluted in ethanol and divided into 4 parts (A,B,C and D). The procedure of flavonoids analysis are as follows :

- Bate-Smith test

0.5 ml concentrated HCl was added to part B, and heated. The changing color from dark red to bright red indicated the presence of flavonoids.

- Wilstater test

0.5 ml concentrated HCl was added to part C and also added 0.2 g magnesium, diluted with distilled water, stirred. Then 1 ml butanol was added and shaken. The appearance of red orange color layer indicated the presence of flavonones.

- Thin Layer Chromatography (TLC) test for flavonoids

Spotted the filtrate D on stationary phase of Kiesel gel GF 254, with butanol-glacial acetic acid-water (4: 1: 5) as mobile phase and ammonia vapors spray is applied to make

the filtrate D visible. The appearance of clear yellow spot indicated the presence of flavonoids.

3) Polyphenols test :

10 ml hot distilled water was added to 0.3 g EPFF, stirred, and let it cool in room temperature. Then 3-4 drops of 10% NaCl solution was added and filtered. The filtrate was divided into 3 parts (A,B, and C). A drop of aqueous solution 2% FeCl₃ was added to 4 ml filtrate A, the appearance of a bluish color indicated the presence of phenolic compounds. The procedure of polyphenols analysis are as follows :

- Thin Layer Chromatography (TLC) test for polyphenols

Spotted filtrate B on stationary phase of Kiesel gel GF 254, and chloroform-ethyl acetate-formic acid (0.5: 9: 0.5) as mobile phase and aqueous solution of 2% FeCl₃ as appearance of spot. The appearance of black color indicated the presence of polyphenols.

- Tannins test

A drops of aqueous 2% FeCl₃ was added to 4 ml filtrate C, the appearance of blackish green color indicated the presence of tannins.

D. Acute toxicity study

The experimental animals used for this study are white albino mice (20-32 g), 2-3 months old and were obtained from the animal unit at the Department of Pharmacognosy and Phytochemical, University of Airlangga, Surabaya, Indonesia. They were acclimatized at the animal house for seven days before the experiments. The mice were maintained *ad libitum* on water and growers with Comfeed Pars. The acute toxicity test of PFF extracts were carried out using a modified method of acute toxicity by Ghosh [9], and the conversion dose extract of treatment was based on Loomis [15]. The mice were divided into four group with six mice per group, and they were treated orally with 0; 10.50; 15.75 and 21.00 g/kg body weight respectively of the PFF extract. The animals were then observed every 24 h until 168 h for he acute toxicity level.

III. RESULTS AND DISCUSSION

A. Phytochemical ethanol extract of PFF

The results of the qualitative phytochemical composition of EPFF were presented in Table 1. It showed the presence of saponins, saponinins, terpenoids, flavonoids, tannins, polyphenols, and alkaloids were not detected both in the precipitation reaction and thin-layer chromatography test. Although Santoso *et al.* [21] reported that the chloroform and methanol extract of pedada fruit showed the presence of alkaloids, Mingqing *et al.* [16] found the presence of steroids, flavonoids, triterpenoids, and maslinic acid [26], and this acid has antihyperglycemic activity [18], this is also supported by Tiwari *et al.* [23] who reported that the methanol extract of the fruit pedada contains a oleanolic acid, luteolin, and β -sitosterol-3-O- β -D-glucopyranoside.

TABLE I
RESULT OF QUALITATIVE PHYTOCHEMICAL ETHANOL EXTRACT OF PFF

Constituents	Solvent	Reagent	Results
Alkaloids	-----	Mayer	ND
	-----	Wagner	ND
	-----	Dragendorff	ND
Saponins	Distilled water	-----	++
Sapogenins	n-hexane-ethyl acetic	anisaldehyde sulfuric acid	+
Terpenoids	n-hexane-ethyl acetic	anisaldehyde sulfuric acid	+
Flavonoids	-----	Bate Smith and Metcalf	+++
	-----	Wilstater	+++
	Botanol-acetic acid – water	ammonia vapor	+++
Tannins	-----	2% FeCl ₃	+++
Polyphenols	Chloroform-ethyl acetic-formic acid	2% FeCl ₃	+++

Note: +++ = relative abundance of compound,
++ = moderate abundance of compound;
+ = relative low presence of compound; ND = not detected

The phytochemical analysis of flavonoids, tannins and polyphenols in the EPFF extract showed that this compounds are greater in quantity than the other phytochemical test. Result to the research by Sadhu *et al.* [19], this type of flavonoids found in pedada fruit i.e, luteolin and luteolin 7-O-β-glucoside, has a hypoglycemic effect [14], an antioxidant and antidiabetic [20],[21]. The presence of tannins, flavonoids, saponins and steroids have moderate effect on the lipid profile [7], saponins as an active role in treating diabetes [3].

B. Acute toxicity ethanol extract of PFF

The results of acute toxicity ethanol extract of PFF in mice were presented in Table 2 showed that in each treatment group was not found dead mice. Treatment dosing did not influence the behaviour of mice compared to controls during the observation. A dose of 21.00 g/kg in a single oral dose ethanol extract of PFF did not reveal any death until 168 h after administering test the material, so the value of a single dose oral LD₅₀ of the ethanol extract of PFF could not be calculated, due to the absence of an animal dead. So it could be concluded that a dose of 21.00 g/ kg as the LD₅₀ value for the ethanol extract of PFF, with category ‘not toxic’ and ‘safe’ for food product.

TABLE III
THE NUMBER OF DEAD MICE WITH ETHANOL EXTRACT OF PFF FOR 168 H

Group (n=6)	Ethanol extract of PFF (g/kg body weight)	Dead of mice for 0 - 168 h
1	0	0
2	10.50	0
3	15.75	0
4	21.00	0

Ukechukwu *et al.* [24] stated that there were not dead animals in each dose group ratings, the highest dose given to test animal, is considered as the value of LD₅₀. In this experiment the ethanol extract of PFF contained saponins, tannins and flavonoids on acute toxicity test and did not case the death of the animals. This results reported by Kalu *et al.*

[13] who stated that leaf extract of *Combretum dolichopentalum* contain saponins, alkaloids, flavonoids also did not find any dead mice. The same result was also reported by Gadanya *et al.* [8].

IV. CONCLUSIONS

This research has revealed that PFF is rich in phytochemical and the LD50 more than 21.00g/kg body weight, therefore it is ‘non toxic’ and ‘safe’ for consumption as mixture of food product.

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