



ICPPS 2014

Proceeding

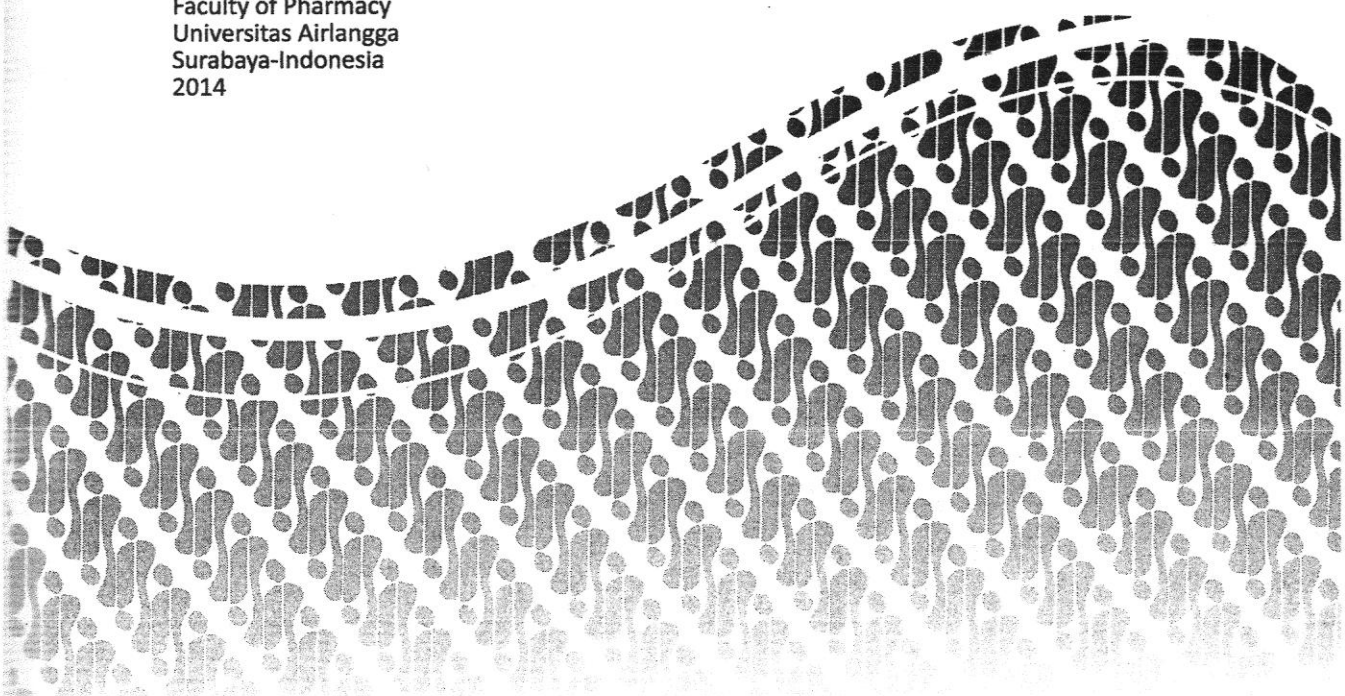
The 1st International Conference on Pharmaceutics & Pharmaceutical Sciences

Drug Delivery Systems:
From Drug-Discovery, Pre-formulation, Formulation and Technological Approaches for
Poorly Soluble Drugs and Protein

Proceeding

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PREFACE From Chairman

It is our pleasure to present you the proceedings of The 1st International Conference on Pharmaceutics and Pharmaceutical Sciences (ICPPS) organized by The Faculty of Pharmacy Universitas Airlangga Surabaya Indonesia.

The proceeding was produced based on papers and posters presented at The 1st International Conference on Pharmaceutics and Pharmaceutical Sciences (ICPPS), held in Surabaya, Indonesia, 14-15 November 2014.

The proceeding clearly reflects broad interest, from the participants that coming from all around the world.

The papers presented were pharmaceutics and biopharmaceutics; requirements on how to evaluate molecules in discovery and their appropriateness for selection as potential candidate; their development in context of challenges and benefits, together with associated time and cost implications and also requirements to progress through pre-clinical and clinical.

In this an opportunity, I would like to express my appreciation to the editorial team of the proceeding who have been working hard to review manuscripts, and making the first edition of this proceeding be possible.

I would like also to thanks to all invited speakers and presenters who participated in The 1st International Conference on Pharmaceutics and Pharmaceutical Sciences (ICPPS) and your contribution to this proceeding.

Finally, I hope this proceeding will give contribution to the Pharmaceutics and Pharmaceutical Sciences research.

Chairman,

Dra. Esti Hendradi, MSI., Ph.D., Apt

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IN VITRO ANTIMALARIAL ACTIVITY OF CHLOROFORM SUBFRACTION OF SALAM BADAK LEAVES (*Acmena acuminatissima*)

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INTRODUCTION

Malaria is still an endemic disease in more than 90 countries, mainly in developing countries (Sanchez et al., 2004). Patients infected with malaria have doubled in the last two decades. This occurred mainly due to the emergence of resistant strains of *Plasmodium falciparum* malaria to the chloroquine and its derivatives (Trape et al., 2002). Perez et al. (1997) stated that the global spread of malaria parasites that are resistant to current antimalarial drug is a major health problem. Therefore, it is need to find new antimalarial substances to replace current drugs that are not sensitive anymore. One attempt to discover new antimalarial substances is through active exploration of the natural sources.

It is known that some species of the family Myrtaceae has an antimalarial activity. Previous studies shown that ethanol extract and chloroform fraction of *A. acuminatissima* leaves (Myrtaceae) were exhibited antimalarial activity with IC₅₀ value of 0.040 µg/ml and 0.006 µg/ml, respectively. Therefore, the separation of the chloroform fraction was done to determine the antimalarial activity substances of *A. acuminatissima*.

MATERIALS AND METHODS

Plant materials

A. acuminatissima leaves was obtained from Purwodadi Botanical Garden on December 2012. Sample was authenticated by the authority of Purwodadi Botanical Garden, Pasuruan, East Java.

Fractionation

Chloroform fractions which obtained from the fractionation of ethanol extract of *A. acuminatissima* leaves was separated by vacuum liquid chromatography (VLC) using hexane, chloroform, and ethanol at gradient condition. This separation was produced six subfractions, then evaporated using a vacuum evaporator and dried. These six subfractions was then analyzed by TLC and tested for antimalarial activity.

In vitro antimalarial activity test

Antimalarial in vitro test was performed based on Budimulya et al. (1997). Sample prepared in serial dilution at concentration of 0.01; 0.1; 1; 10 and 100 µg/ml in microwells. Each microwell was added with 500 µl parasite culture (1% parasitemia, 5% haematocrit) and incubated for 48 hours in 37°C. After incubation, thin blood smears were made and stained using 20% giemsa dye. Percentage of parasitemia was determined by counting infected erythrocytes per 1000 total erythrocytes under microscope.

RESULTS AND DISCUSSION

The separation results of *A. acuminatissima* leaves chloroform fraction was produced six subfractions (A-F). Each subfraction was tested for in vitro antimalarial activity against *P. falciparum* (3D7). According to Rasoanaivo et al. (2004), an extract is very active if IC₅₀ < 5 µg/mL, active 5 µg/mL < IC₅₀ < 50 µg/mL, weakly active 50 µg/mL < IC₅₀ < 100 µg/mL and inactive IC₅₀ > 100 µg/mL. The test re-



sults showed that all chloroform subfractions classified as very active and subfraction E was the most active with IC₅₀ value of 0.007 µg/mL. The result of antimalarial activity test of chloroform subfractions (A-F) can be seen in Table 1.

Identification of chloroform subfraction was performed by TLC method using silica gel GF254 as stationary phase and chloroform: methanol (98:2) as a mobile phase. Then observed under UV light in wavelength of 254 nm, 366 nm and sprayed with H₂SO₄ 10%. TLC chromatogram profile showed that all subfractions contain purple spot (Fig.1). According to Sharifa et al. (2012), terpenoid compounds will form a pink to purple or violet after being sprayed with 10% H₂SO₄. It can be considered that all subfractions were containing terpenoids.

Sub fraction	% Inhibition at a concentration of (µg/ml)					IC ₅₀ (µg/ml)
	100	10	1	0.1	0.01	
A	100	100	75.32	65.18	56.19	0.012
B	100	83.39	63.95	47.69	24.32	0.158
C	86.41	83.94	41.38	35.17	20.40	1.469
D	96.90	92.48	64.83	36.5	26.40	0.172
E	100	85.70	70.42	69.84	56.75	0.007
F	97.19	91.41	68.41	59.81	54.61	0.014

Table 1. IC₅₀ values of Chloroform subfractions against *P. falciparum* 3D7

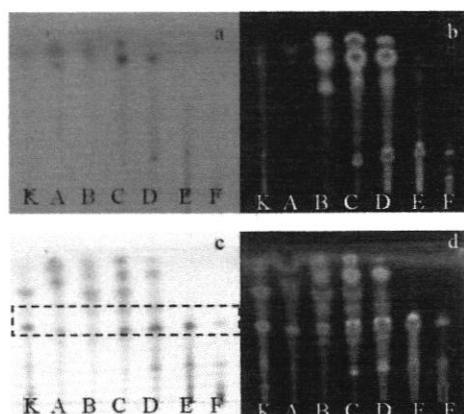


Figure 1. TLC chromatogram of chloroform subfraction using silica gel F254 as stationary phase and chloroform-methanol (98:2) as mobile phase, viewed under UV light : (a) 254 nm; (b) 366 nm; (c) after sprayed with 10% H₂SO₄ and heated at 105°C for 5 minutes. (d) 366 nm after sprayed with 10% H₂SO₄ and heated at 105°C for 5 minutes; K = chloroform fraction, A-F = subfraction.

[---] = considered to contain terpenoids.

Subfraction A-F were exhibited antimalarial activity (IC₅₀ value of 0.007-1.469 µg/ml) and containing terpenoids substances. Based on that result, it is possible to conclude that antimalarial activity of subfractions was derived from terpenoids substances.

CONCLUSION

Subfraction E of *A. acuminatissima* leaves was the most active as antimalarial with IC₅₀ value of 0.007 µg/ml. The antimalarial activity was possible derived from terpenoids substances. Further study needed to isolate and identify the active compounds from *A. acuminatissima*

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