

PROCEEDING

The International Conference on
**Pharmacy and Advanced
Pharmaceutical
Sciences**



Faculty of Pharmacy UGM
Yogyakarta Indonesia
October 2009

9th International Conference on Pharmacy and Advanced Pharmaceutical Sciences

Faculty of Pharmacy UGM



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**The International Conference on Pharmacy
and Advanced Pharmaceutical Sciences
Yogyakarta, Indonesia, 2009**

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Preface from the Editor

The proceeding was produced based on papers and posters presented at the international Conference on Pharmacy and Advanced Pharmaceutical Sciences, held in Yogyakarta, Indonesia, 5 – 6 October 2009.

The proceeding clearly reflects broad interest; from there are participants coming from all around the world. Many contributions on Pharmaceutical Sciences there are quite a substantial number of papers on Pharmacist role in general. The papers presented file into a broad spectrum in Pharmaceutical sciences including Pharmacology, Toxicology, Analytical Chemistry and Drug Design, Drugs Synthesis, Formulation of Drugs, Pharmacy Social, Pharmacoepidemy, Traditional Medicine Natural Product Chemistry and Phytochemistry, etc.

In addition there are substantial numbers of paper deal with professional aspect of Pharmacist in general health care.

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 International Conference on Pharmacy and Advanced Pharmaceutical Sciences and your contribution to this proceeding.

Finally, I hope this proceeding will give contribution to the advanced scientific research in the field of pharmaceutical sciences

Yogyakarta, July 2010

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Welcome Message From the committee

Welcome to Yogyakarta

On behalf of the Scientific and Organizing Committees, it is a great pleasure for me to welcome all participants to Yogyakarta, to the International Conference on Pharmacy and Advanced Pharmaceutical Science 2009.

The international conference is organized by the faculty of Pharmacy UGM to celebrate its 63th anniversary and the Lustrum XII of Gadjah Mada University, as a collaboration work between the Faculty of Pharmacy UGM with the Nara Institute of Science and Technology (Japan) and the Universiti Sains Malaysia (Malaysia). In this conference 15 lectures within the field of Pharmaceutical Care and Advanced Pharmaceutical Science will be given by invited speakers. Besides, 55 posters and 75 paper will be presented in the parallels presentation sessions. Herewith, we express our gratitude to all speakers and presenter, who would like to share their advance knowledge in this scientific event.

The Organizing Committee gratefully acknowledges the Nara Institute of Science and Technology and the Universiti Sains Malysia, for the nice collaboration in bringing forth this conference. A special acknowledgment is addressed to the Rector of Gadjah Mada University and the sponsors, for all supports that make this symposium possible. Furthermore, personally, I want to express my deep appreciation to the members of the Organizing Committee, for the good teamwork and their great effort given in the preparation for this symposium.

Finally, I wish all participants a scientifically rewarding and an enjoyable meeting in Yogyakarta.

Chairman

Dr. Hilda Ismail, M.Si., Apt.

Remark of the Dean Faculty

Assalamu'alaikum wr. wb.

Distinguished ladies & gentlemen.

First of all, on behalf of the Faculty of Pharmacy Universitas Gadjah Mada, I would like to come to all of you in Yogyakarta, thank you very much for your attention to come and to attend the international Symposium on Pharmacy and Advanced Pharmaceutical Sciences. I hope we are all in health condition.

Ladies and gentlemen,

The symposium is organized by the Faculty of Pharmacy UGM in collaboration with the Faculty of Pharmaceutical Sciences Universiti Sains Malaysia and the Nara Institute of Science and Technology Japan, and held as part to celebrate the 63th anniversary of the Faculty of Pharmacy UGM.

In the symposium, I hope we can communicate our recently information concerning social / clinical pharmacy and pharmaceutical sciences. I hope the symposium will be very fruitful, very useful for all of us.

I address special thanks to the plenary speakers both from domestic and abroad, the oral and poster presenters, as well as to those who come just to know the development of clinical or social pharmacy and pharmaceutical science. Your willingness to come, to communicate and to share your experiences is highly appreciated.

Special thanks also I address to my colleague the Dean of Faculty of Pharmacy USM who has been coordinating USM students to attend this symposium. The hope is not to set up networking between the pharmacy students of USM and UGM.

Therefore, during almost whole day discussing scientific matter related to human health and welfare, I hope we can make a wonderful opportunity to make a scientific closer relationship while we enjoy the cultural performances of Yogyakarta presented by our pharmacy student.

Finally, I hope that this meeting will give benefits to all of us, and we may see each other again in a similar event in the near future.

I look forward to thank you all for attending this event.

Wassalamu'alaikum wa rahmatullahi wa barakatuh,
Dean of Faculty of Pharmacy UGM

Prof. Dr. Marchaban, DESS., Apt.

**Speech of the Senior Vice Rector
For Education, Research and Community Services,
Gadjah Mada University**

Assalamu'alaikum wa rahmatulLahi wa barakatuh,

On behalf of the Rector, I would like to welcome all of you to our campus Gadjah Mada University and to our home town Yogyakarta. It is a great honor for me and Gadjah Mada University to host the Two-day International Conference on Pharmacy and Pharmaceutical Sciences that is conducted by the Faculty of Pharmacy, Gadjah Mada University. The increasing problems and new cases of some diseases in the world, both the infectious and the degenerative diseases, have demanded the development of medical and pharmaceutical sciences and technologies for supporting the developments of early detection methods of the diseases, the accurate diagnoses, as well as the appropriate and effective medications or therapy. Pharmaceutical Science and Technology have been developing very fast within recent years. The development trend shows using much more biotechnological approach in both diagnose establishment and medication administrations. For examples the usage of some serums, enzymes, hormones, vaccines, etc., and their recombinant products. The science and technology for finding prevention method against infectious diseases or degenerative diseases now have been developing so amazing, for example the usage of growth hormones, vaccines, and stem cells for it.

Gadjah Mada University has been committed to become World Class University; therefore international networking in education, research and publication is much needed. I really support to this international conference on Pharmaceutical Science and Technology which can keep us in touch with the state of the art of pharmaceutical science. I do believe that by conducting this kind of international meeting, we can get and exchange new information and best practices on pharmaceutical science and technology, and it is very important to inspire our young researchers and enhance our research networking internationally. In this occasion, I would like to express my great gratitude to all the guest speakers and speakers, who have contributed their advanced presentations in this international conference. I also would like to extend my gratitude to the Organizing Committee from the Faculty of Pharmacy, Gadjah Mada University, who has already successfully arranged this international conference. I would also thank to all institutions or companies who have sponsored and supported this conference.

Finally, have a fruitful conference and enjoy Yogyakarta. Thank you
Wassalamu'alaikum wa rahmatulLahi wa barakatuh,

Senior Vice Rector for Education, Research and Community Service
Gadjah Mada University

Prof. Dr. Retno Sunarminingsih, M.Sc., Apt.

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The Influence of Arbutin 3% and Sesame Oil (3,5,7 % w/w) on SPF Values of Oxybenzone and Padimate O (3:7% w/w) in carbomer Gel Base

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Abstract

This research was intended to know the influence of addition 3% Arbutin and (3; 5; 7)% w/w sesame oil on SPF value of sunscreen combination Oxybenzone and Padimate - O (3:7 % w/w) was formulated in carbomer gel base. Characteristic test was done in 2nd day after formulated included organoleptic, pH, and spread-ability. The sunscreen effectiveness determination (SPF value) was performed in vitro by measuring the absorbencies of its 10 ppm solutions in isopropanol between 290 - 400 nm. Organoleptic characteristic all formulas were different with the base, so did the pH and spread ability result. From SPF value determination results conclude that addition of arbutin increased SPF value, so did the increased of sesame oil concentration. Formula with the higher concentration of sesame oil had a higher SPF values. From this research, there was suggested to evaluate in vivo acceptability test and affectivity in vivo trial test on human skin.

Key words: Arbutin, Sesame oil, sunscreens, carbomer gel, Oxybenzone, Padimate - O, SPF values

Introduction

Sunscreens are mostly used to protect the skin from harmful effects of solar radiation. The active ingredients of sunscreens are divided into physical and chemical sunscreen agents. Physical sunscreen agents reflect the UV rays from our skin. It is not transparent and has to be used in a high concentration to be effective, which made them not preferable to some people. The chemical sunscreen agents work by absorbing UV-A radiation like oxybenzone and absorbing UV-B radiation such as Padimate-O (octyldimethyl PABA). To obtain a higher protection and producing a broad spectrum sunscreen, a combination of anti UV-A and anti UV-B sunscreen agents has been used nowadays in many sunscreen product [Widianingsih and Lumintang, 2002].

Normally, our skin has its own protection mechanism against the harmful effect of UV rays such as the thickening of *stratum corneum*, sweat and skin pigmentation. Increasing abnormality of melanin as a result of our skins natural protection can result in a non homogenous skin color which is usually reduce likeliness amongst people. To overcome this problem, whitening agents are used to control the production and metabolism of melanin in the epidermis. One example of a frequently used whitening agent is arbutin, a hydroquinone derivatives which inhibits melanin production. Arbutin has lower toxicity than hydroquinone's and its depigmentation effect is higher than kojic acid and vitamin C [Mashhood, 2006]. This substance is used as whitening agent at a various concentration ranging from 0.5-3.0%. It has a low partition coefficient and penetration rate. Therefore, it needs addition of a penetration enhancer [Zulkarnain, 2003; Galilee, 2008; Mitsui, 1998].

One of enhancers that frequently used in cosmetic formula is sesame oil. Sesame oil is a vegetable oil that obtained from pressing *Sesamum Indicum* seed. Mechanism of sesame enhanced polar pathway in stratum corneum because it contains protein (25%) dan globulin (67,3%) as main substance. Another mechanism is to enhance the oclusiveness (Sarma, K., 1993) to facilitate arbutin penetration. Sesame oil has been widely used in sunscreen and other cosmoceutics preparation for its emollient activity; it is nonirritant, considered as a natural lipid that has the highest compatibility with human skin and has sunburn healing effect [O'Neil, 2006].

Among several variant sunscreen preparations, gel gives us a cool sensation, not sticky, elegant, and smooth and easy to be washed from the skin. A synthetic gelling agent

like *carbomer* 940 usually requires only a small amount of them to produce a gel with good consistency compared to other types of gelling agent.

The aims of this research wants to know whether addition of arbutin 3% w/w and various concentration of sesame oil (3, 5, and 7% w/w) as arbutin's enhancer affect SPF value of sunscreen combination of oxybenzone 3% w/w (as anti UV-A) and Padimate O 7% w/w (as anti UV-B) which is formulated in *carbomer* 940 gel base. SPF values are observed by using a spectrophotometric method.

Methodology

Preparation of sunscreen gel containing arbutin and sesame oil

Oxybenzone, Padimate-O and propylene glycol were mixed and put into the good emulsion system contains arbutin, water, BHT, sesame oil, and tween 80 as emulsifying agent. After that it poured into the *carbomer* gel base and stirred well to form the sunscreen gel. The sunscreen were then kept in a tight container and stored well for further analysis. Preparation of each formula is described as below table.

Table 1: Formula used in research

Composition	Function	Concentration in formula (% w/w)				
		S	S+A	S+A+O 3%	S+A+O 5%	S+A+O 7%
Oxybenzone	Sunscreen agent	3	3	3	3	3
Padimate O	Sunscreen agent	7	7	7	7	7
Arbutin	Whitening agent	0	3	3	3	3
Tween 80	Emulsifying agent	0.5	0.5	0.5	0.5	0.5
Sesame oil	Enhancer	0	0	3	5	7
<i>Carbomer</i>	Gelling agent	1	1	1	1	1
Distilled water up to	Solvent	100	100	100	100	100

* Formula I = S = Sunscreen

Formula II = S+A = Sunscreen + Arbutin

Formula III = S+A+O 3% = Sunscreen + Arbutin + Sesame oil 3%

Formula IV = S+A+O 5% = Sunscreen + Arbutin + Sesame oil 5%

Formula V = S+A+O 7% = Sunscreen + Arbutin + Sesame oil 7%

Characteristics determination of sunscreens gel

The characteristic of preparation were determined include: organoleptic test visually, while the determination of pH and spread ability are done in 2 days after the formula were made by using a digital pH meter Schott CG 842, and a spread-ability apparatus.

Determination of SPF value of sunscreens gel

SPF value was determined by Petro's method.

Firstly, 100.0 mg sunscreens were dissolved in 2.0 ml isopropanol, the solution was then centrifuged for 15 minutes at 50 rpm. 1.0 ml of the filtrate is taken and poured into a 5.0 ml metered flask and shake well until its homogenized (10000 ppm). This solution diluted it and then its shaken well until it reached a concentration of 100 ppm (contains 10 ppm sunscreen's active ingredients). An UV spectrum of this solution was measured at 290-400 nm using Double Beam UV-Vis Spectrophotometer (Perkin Elmer Lambda EZ 201) at an interval of 2 nm which has an absorption that is larger than 0.050.

The AUC of each formula from the shortest and longest wavelength are calculated using the following equation:

$$AUC_{\lambda_{p-\alpha}}^{\lambda_p} = \frac{A_{p-\alpha} + A_p}{2} (\lambda_p - \lambda_{p-\alpha})$$

Whereas:

- AUC = Area under Curve
- A_p = Absorbance on p wavelength
- A_{p-α} = Absorbance on p-α wavelength

The SPF values of a formula were obtained by converting log SPF calculating from the total AUC into the equation below:

$$\text{Log SPF} = \frac{\text{Total area}}{\lambda_n - \lambda_1} \times 2$$

Whereas:

- λ_n = longest wavelength above 290 nm that has an absorption value higher than 0.050
- λ₁ = shortest wavelength 290 nm

Statistical analysis

One-way ANOVA were used to assess the significant of differences only for preparation with the same sunscreen category.

Results and Discussions

The average data of organoleptic observation from all sunscreen formula were shown in table 2. From the data we could see that the addition either of arbutin or sesame oil to the formula affect the color and smell.

Table 2: Organoleptic analysis of the sunscreens formula

Formula	Colour	Smell	Consistency
Gel base	Transparent	Carbomer like	A bit viscous
Sunscreen (S)	Pale Yellow	Padimate-O like	Viscous
S+Arbutin (A)	Pale Yellow	Padimate-O like	Viscous
S+A+Olive oil (O) 3%	Yellowish	Sesame oil like	Viscous
S+A+O 5%	Yellowish	Sesame oil like	Viscous
S+A+O 7%	Yellowish	Sesame oil like	Viscous

Table 3: pH of sunscreen formula

Formula	pH (average)	% var. coefficient
Gel base	6.60 ± 0.07	1.05
Sunscreen (S)	6.31 ± 0.04	0.74
S+Arbutin (A)	6.27 ± 0.05	0.70
S+A+Sesame oil (O) 3%	6.26 ± 0.02	0.28
S+A+O 5%	6.30 ± 0.04	0.63
S+A+O 7%	6.30 ± 0.05	0.83

* The result were obtained from an average of 3 times replication

One of the important factors that influence sunscreen SPF value is pH, besides extinction coefficient and solvent polarity. Therefore it is important to know the mechanism

of SPF changes in the treatment formula. The pH of sunscreen gels also play a major role in the sunscreen gels characteristic as the carbomer consistency were heavily affected by its acidity. Acid condition would lower the gels viscosity. From pH data, value of F calculation = 26.083 > F table = 5.42. Tukey HSD test of the pH data has shown decreasing pH by addition of sunscreen ingredient. However, addition of arbutin and/or sesame oil did not affect the pH even though sesame oil concentration increased. Overall, from the pH data we know that all sunscreen formula were in a range of skin pH.

The spread-ability of the formula was measured and the results were shown in table 4: From the data we could see that the spread-ability of the formula increased with the addition of arbutin but decreased with a rise in sesame oil concentration.

Table 4: Spread-ability of sunscreens formula

Formula	Average slope (mm/g)	% Var. coefficient
Gel base	0.2833 ± -	-
Sunscreen (S)	0.3359 ± 0.004	2.26
S+Arbutin (A)	0.3859 ± 0.004	3.09
S+A+Sesame oil (O) 3%	0.2713 ± 0.002	3.47
S+A+O 5%	0.2474 ± 0.003	2.91
S+A+O 7%	0.2238 ± 0.0007	1.25

* The result were obtained from an average of 3 times replication

The SPF analysis was done by extracting the sunscreen from its base by using isopropanol. In order to assure that the base gel *carbomer* 940 will not give any absorption in the UV spectrum, it is also extracted using isopropanol and observed at the wavelength of 290-400 nm. From the spectra observed it is known that the gel base did not give any absorption at all, Therefore it is assumed that any change within the observation of SPF value were caused by the active ingredient only.

The SPF value that resulted from the calculation (Table 5) was then categorized based on the American Society of Health System Pharmacist standard in order to classify its protection capability.

Table 6: SPF data from the treatment formula

Formula	Average SPF** ± SD	% KV	SPF Category (American Society of Health System Pharmacist Standard)	% increasing SPF value compare with control
I: S (Sunscreen preparation in gel)	6,658 ± 0,15	2,21	Moderate Sun Protection Product	0
II: S + arbutin	9,094 ± 0,24	2,63	High Sun Protection Product	36.58
III: S+A+O 3%	9,615 ± 0,50	5,18	High Sun Protection Product	44.42
IV: S+A+O 5%	11,119 ± 0,34	3,08	High Sun Protection Product	67.01
V: S+A+O 7%	16,435 ± 0,61	3,72	Very High Sun Protection Product	146.84

**The data were obtained from an average of 3 different products in 1 formula

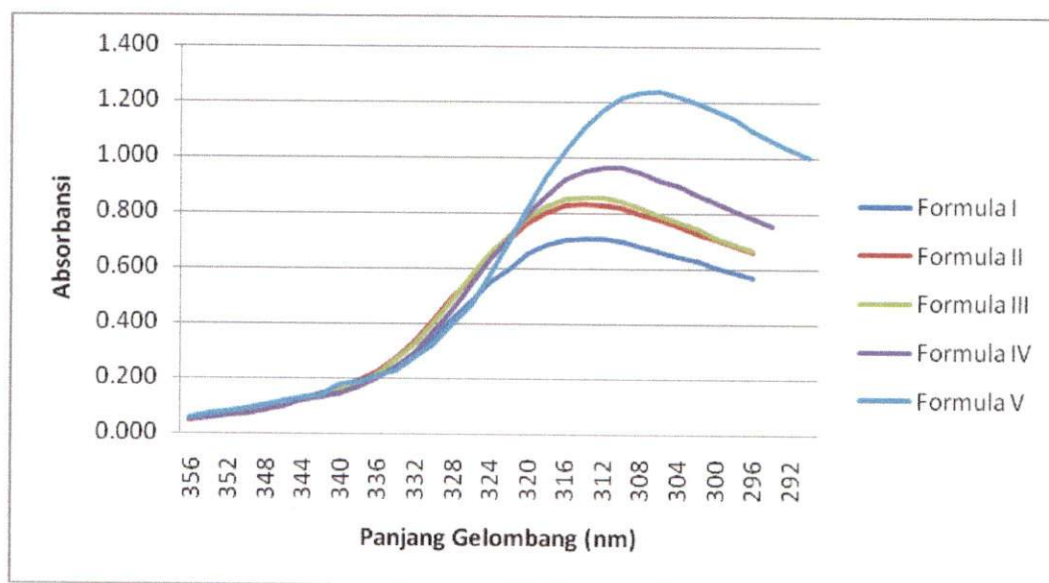


Figure 2: Absorption spectrum of sunscreen

According to category as presented Table 5, we know that arbutin elevated the sunscreen protection category, from "Moderate Sun Protection Product" to "High Sun Protection Product". Increasing sesame oil up to 5% did not alter the category. However addition of 7% sesame provide the maximum protection resulted the sunscreen category enhanced to be "Very High Sun Protection Product"

From the screening spectra profile as shown in Figure 2, it is learned that the addition of arbutin and sesame oil up to 5% did not cause any shiftness on the maximum wavelength. Nevertheless, an increase in the intensity of absorption was observed. Thus it is predicted that an interaction occurred between the molecule of arbutin and sunscreen agent which intensify the effect of aucsochrome group. And it is clear that addition of 7% sesame oil not only increased the absorption intensity but also increased the protection sunscreen, by maximum wavelength shift. This shiftness cause decreasing polarity of sunscreen gels that affect the delocalization of the molecule and resulted in a rise of energy demand needed for excitation to happen and hence, increase the SPF value.

Table 9: HSD test of the sunscreens SPF

		SPF			
Tukey HSD ^a		Subset for alpha = .05			
Kelompok	N	1	2	3	4
Formula I	9	6.6580			
Formula II	9		9.0936		
Formula III	9			9.6153	
Formula IV	9				11.1192
Sig.		1.000	1.000	1.000	1.000

Means for groups in homogeneous subsets are displayed.

a. Uses Harmonic Mean Sample Size = 9.000.

Conclusion

The addition of arbutin and sesame oil affect the physical appearance (organoleptic and consistency) of sunscreen product as well as its spread ability and significant increase in SPF value of the sunscreen formula combination Oxybenzone and Padimate - O (3:7 % w/w) which was formulated in carbomer gel base.

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References

- American Society of Health-System Pharmacists, 2002, AHFS Drug Information, USA: American Pharmaceutical Association.
- Galilee, 2008, Technical Data Sheet: Arbutin (Beta-Arbutin), Israel.
- Mashood, A.A., 2006, Treatment of Hyperpigmentation Disorders, Journal of Pakistan Association of Dermatologists, Vol. 16, p. 65-68.
- Mitsui, T., 1998, New Cosmetic Science, 2nd ed, Amsterdam: Elsevier
- O' Neil, M.J., 2006, The Merck Index, 14th ed, USA: Merck and Co., Inc.
- Petro, A.J., 1981, Correlation of Spectrophotometric Data with Sunscreen Protection Factors, International Journal of Cosmetic Science, Vol. 3, p. 185-196
- Pratiwi, V.K., 2006, Pengaruh Asam Glikolat Terhadap Stabilitas Fisik dan Peningkatan Efektivitas in vitro Sediaan Tabir Matahari Kombinasi Oksibenson-Oktilsimetil PABA (3:7) %b/b, Skripsi, Surabaya: Fakultas Farmasi Universitas Airlangga
- Sarma, K., 1993, Vegetable oil-based skin permeation enhancer compositions and associated method and systems, US Patent, patent no. 5,229,130
- Widianingsih, N.P.T., and Lumintang, H., 2002, Pemakaian Tabir Surya (Sunscreen), Berkala Ilmu Penyakit Kulit dan Kelamin, Vol. 14 No. 2, p. 155-167.