The development of antioxidant peel-off facial masks from cinnamon bark extract (Cinnamomum burmannii)

Sani Ega Priani*, Restianti Mutiara, Dina Mulyanti

Pharmacy Department, Bandung Islamic University (UNISBA)
Taman Sari 1, Bandung, Indonesia

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ABSTRACT

The bark of cinnamon (*Cinnamomum burmannii*) contains cinnamaldehyde and other active substances with potent antioxidant properties. Antioxidants are effective at preventing and reducing UV-induced skin damages and skin aging. This study was intended to formulate and characterize the antioxidant peel-off facial masks containing cinnamon bark extract and the combination of polyvinyl alcohol (PVA) and hydroxypropyl methylcellulose (HPMC) as gelling agents. The ethanol extract of cinnamon bark and the developed peel-off mask were evaluated for their antioxidant activities by the α , α -diphenyl- β -picrylhydrazyl (DPPH) free radical scavenging method and for their physical characteristics. The cinnamon bark extract exhibited a very strong antioxidant activity, as evidenced by IC₅₀= 10.04 \pm 0.08 ppm. As for the formulated peel-off mask, it had excellent physical characteristics, which were identified during organoleptic observations and pH, viscosity, spreadability, and film drying time evaluations. Similar to its constituent extract, this mask produced significantly potent antioxidant effects, with IC₅₀= 47.31 \pm 1.47 ppm. For these reasons, peel-off facial masks containing cinnamon bark extract have not only excellent physical characteristics but also powerful antioxidant properties.

Keywords: peel-off mask, cinnamon bark, ethanol extract, antioxidant

*Corresponding author:

Sani Ega Priani Pharmacy Danart

Pharmacy Department, Bandung Islamic University (UNISBA)

Taman Sari, 1 Bandung , Indonesia Email: egapriani@gmail.com

INTRODUCTION

Cinnamomum burmannii Nees ex. Bl., commonly known as Indonesian cinnamon or Padang cassia, grows in Southeast Asia and is cultivated in parts of Indonesia and the Philippines. C. burmannii stem bark (or cinnamon bark) contains cinnamaldehyde and other active substances that reportedly have antioxidant properties (Al-Dhubiab et al., 2012; Vishnuvardhanaraj et al., 2013; Singh et al., 2007). Previous research confirms that the ethanol extract of cinnamon bark is a highly effective antioxidant with an Inhibitory Concentration (IC₅₀) of <50 ppm (Priani et al., 2014). Antioxidants are effective in preventing and reducing skin damages and aging due to exposure to UV lights. Also, antioxidant compounds can eliminate free radicals and decelerate oxidation and, when applied topically to the skin, can prevent UV-induced oxidative damages (Pandel et al., 2013; Godic et al., 2014).

Previous studies have developed emulsion gel containing black seed oil and proven their antioxidant and UV-protection properties (Priani *et al.*, 2014; Priani *et al.*, 2014). In this study, another dosage form was prepared for non-daily application with high occlusive capacity, namely, peel-off facial masks. Facial masks are one of the topical preparations with many beneficial effects, including skin whitening, skin moisturizing, and freckle removal. Meanwhile, peel-off masks are a kind of facial mask composed of film-forming polymers, such as polyvinyl alcohol or polyvinyl acetate, that create a cohesive plastic layer after entirely dried, allowing manual removal from the skin without leaving any residues. These preparations provide moisturizing action and enhance the effect of active compounds on the epithelium owing to the occlusive reaction caused by plastic polymeric layers (Berigh *et al.*, 2013; Vieria *et al.*, 2009).

The objective of this research was to formulate peel-off facial masks from cinnamon bark extract with excellent physical characteristics and potent antioxidant activities. These facial masks can be used as herbal cosmetic preparations.

MATERIALS AND METHOD

Materials

The plant materials used in this research were collected from Manoko (the Province of Jawa Tengah, Indonesia) and determined by the Herbarium Bandungense, ITB.

Methods

Preparation of cinnamon bark extract

The stem bark was dried, ground into powder, and then extracted by the maceration process using 96% ethanol with a 1:3 ratio. The maceration was carried out for 24 hours, at room temperature, and with three replicates. Afterward, the extract solution was evaporated in a vacuum rotary evaporator at 40°C and 220 mBar (Priani *et al.*, 2014). Preliminary screening of secondary metabolites, such as alkaloids, flavonoids, saponins, quinone, tannin, terpenoids, polyphenol, and steroids, was conducted according to the conventional phytochemical methods (Fransworth, 1966).

Antioxidant activity test of the cinnamon bark extract

The extract was tested for its antioxidant activity by the DPPH method. A DPPH solution (40 ppm, 2 mL) was mixed with 2 mL of the ethanol extract solution at various concentrations (5, 10, 20, 30, and 40 ppm). An equal amount of ethanol was then added to the control DPPH. After 30 minutes of incubation, the absorbance of this mixture was read at 514 nm. The % inhibitions of DPPH, free radicals, were calculated using the following equation:

% inhibition =
$$\frac{Abs\ control - Abs\ Sample}{Abs\ control} \times 100\%$$

where *Abs control* is the absorbance value of the control DPPH solution and *Abs Sample* is the absorbance of the extract and DPPH solution. The extract concentration and % inhibition were plotted to a graph to determine their linear regression equation. The results were expressed in IC₅₀, i.e., the

concentration of sample required to achieve 50% inhibition of the DPPH free radicals (Molyneux *et al.*, 2004).

Preparation of peel-off facial mask from the cinnamon bark extract

The composition of the peel-off mask formulated in this study is shown in Table I. Firstly, the stock gel was prepared by dispersing PVA and HPMC in heated purified water (90°C) separately. Secondly, the cinnamon bark extract, methylparaben, and propylparaben were dissolved in propylene glycol. Finally, the solution and the remaining compounds were added to the gel then stirred continuously using a digital stirrer (Suheri *et al.*, 2016).

Table I. The formulation of facial peel-off masks from the cinnamon bark extract

Ingredients	Percentage in Masks (%)
Cinnamon extract	1
PVA	12
HPMC	1
Propylene glycol	5
Glycerin	5
Methylparaben	0.2
Propylparaben	0.05
Purified water until	100

Physical evaluation of the peel-off facial mask

The peel-off mask gel was observed organoleptically (color, odor, and homogeneity) and evaluated for its pH, viscosity, spreadability, and film drying time.

Homogeneity Test

The gel was applied to the surface of a slide then pressed using another. Its homogeneity was observed under a microscope.

Viscosity Test

The viscosity of the gel was determined using a Brookfield RV viscometer with a spindle number of 15 at 10 rpm.

Spreadability Test

About 1 gram of the mask gel was placed on a glass sized 20×20 cm. Afterward, the sample was closed with another glass, pressed with a predefined mass until the total weight reached 125 g, and allowed to stand for 60 s. Then, the diameter of the gel spread was measured (Hanum and Lia, 2016).

Film Drying Time Measurement

About 1 gram of the sample was spread over a glass plate (5.0 x 2.5 cm) to create a uniformly thin layer, i.e., 1 mm in thickness. The glass plate was heated in an oven at 36.5 ± 2.0 °C, which simulated the temperature of the human body. The formulations were monitored until the drying process was completed and the film could be entirely removed from the glass slide (Vieira *et al.*, 2009).

Antioxidant activity test of the peel-off facial mask

The peel-off facial mask was tested for its antioxidant activity by the same method as the extract, which was until the IC_{50} value of the preparation was obtained.

RESULTS AND DISCUSSION

Cinnamon bark was extracted by maceration using a universal solvent (ethanol), and the yield of this extraction was 27.28%. The phytochemical screening results of the stem bark powder and the extract were presented in Table II. This table shows that both the powder and extract contain various secondary metabolites, such as alkaloids, flavonoids, saponins, tannins, quinones, terpenoids, and steroids, all of which are medicinally important. Flavonoids and polyphenols are the compounds responsible for the antioxidant activity of cinnamon bark (SA *et al.*, 2015).

Table II. The phytochemical screening of the cinnamon bark sample and its extract

Class of compounds	Powder of stem bark	Ethanol extract
Alkaloids	+	+
Flavonoids	+	+
Saponins	+	+
Tannins	+	+
Quinones	+	+
Polyphenol	+	+
Monoterpenes and sesquiterpenes	+	+
Steroids and triterpenoids	+	+

(+) = Presence of constituents

The cinnamon bark extract was developed into peel-off masks. PVA, used as the film-forming agent, can enhance the adhesive capacity of the preparation. Its concentration can influence the viscosity, performance of film formation, film thickness after drying, and applicability of the mask. Meanwhile, HPMC was used to enhance the viscosity of the gel (Berigh *et al.*, 2013; Priani *et al.*, 2015). Based on the physical evaluation results presented in Table III, the peel-off facial mask produced had excellent physical characteristics. Also, the acidity/alkalinity of the gel corresponded to the pH of the face skin (4.5-6.5). As for spreadability, the gel was categorically optimum because the diameter of the spread was in the range of 5-7 cm. Furthermore, the film drying evaluation revealed that the time required by the peel-off mask to become entirely dried was optimum, i.e., <30 min (Wendtner *et al.*, 2006; Garg *et al.*, 2002, Vieira *et al.*, 2009).

Table III. The physical characteristics of the peel-off facial mask

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Parameters	Results	
Organoleptic	Transparent brown color, with the faint odor of	
	cinnamon	
pН	5.92 ± 0.09	
Viscosity	$7233.33 \pm 513.16 \text{ cps}$	
Homogeneity	Homogeneous	
Spreadability	6.10 ± 0.10 cm	
Film Drying Time	19.00 ± 1.73 min	
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The antioxidant activity test was conducted by measuring the inhibition of 2,2-diphenyl-1-picrylhydrazyl (DPPH) free radicals. In the DPPH free radical scavenging assay, when antioxidants react with the DPPH, it changed the color of the mixture from purple to yellow (a-diphenyl-β-picryl hydrazine). The degree of discoloration indicates the radical-scavenging potential of the sample. DPPH assay is one of the most popular antioxidant assays chiefly because it is simple, relatively inexpensive, quick, and efficient (Molyneux *et al.*, 2004).

Table IV. The IC_{50} values of the cinnamon bark extract and peel off facial mask			
Samples	Regression	IC ₅₀ values (ppm)	
	equation		
Cinnamon bark extract	y = 3.69x + 11.48	10.04 ± 0.08	
Cinnamon peel-off facial mask	$y = 10.04 \pm 0.08$	47.31 ± 1.47	

90 80 50 70 inhibition (%) inhibition 50 30 $R^2 = 0.9817$ 40 y = 3,6935x + 11,479 20 30 $R^2 = 0.9957$ 20 10 10 0 0 25 10 15 20 10 25 0 20 30 concentration (ppm) Concentration (ppm) (a) (b)

Figure 1. The calibration curves of the antioxidant activity tests of (a) cinnamon bark extract and (b) cinnamon peel-off facial mask

The cinnamon bark extract and the peel-off mask produced IC₅₀ values of 10.04±0.08 and 47.31 ± 1.47 ppm, respectively (Table IV, Figure 1), indicating powerful antioxidants (IC₅₀ less than 50 ppm) (Shetta et al., 2013). The antioxidant effect of this extract is believed to be the product of its bioactive constituents, including cinnamaldehyde, cinnamic acid, eugenol, polyphenol, and procyanidin (Rao et al., 2014). From the test results, the peel-off facial mask formulated from cinnamon bark extract is concluded to have significant potential as cosmetic preparations. With potent antioxidant properties, it can treat the skin and prevent premature aging. Antioxidants are compounds that can counteract or reduce the negative impact of oxidants on the skin. Propylene glycol at the preparation can act as a penetrant enhancer that improves the infiltration of active antioxidant substances through the skin. It also functions as a chemical penetrant enhancer by improving the partitioning of drugs into the stratum corneum (Haque et al., 2018).

CONCLUSION

The peel-off facial mask formulated from cinnamon bark extract has excellent physical characteristics, as evidenced by organoleptic observation and pH, viscosity, spreadability, and film drying time evaluations. Cinnamon bark extract and the peel-off facial mask prepared from it have highly potent antioxidant properties, with the IC₅₀ of 10.04±0.08 ppm and 47.31 ± 1.47 ppm, respectively.

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