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EFFECT OF OPTIMIZATION OF TWEEN 80 AND PROPYLENE GLYCOL AS A SURFACTANT AND COSURFACTANT ON THE PHYSICAL PROPERTIES OF ASPIRIN MICROEMULSION

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ABSTRACT

Background: Aspirin is recommended as a first-line antiplatelet drug for all types of acute diseases that cause thrombosis in the blood vessel, especially in cardiovascular disease. Orally administered aspirin has side effects such as the risk of gastrointestinal bleeding and has presystemic metabolism in the gastrointestinal tract and liver. Transdermal delivery offers an alternative for administering aspirin that passes the gut and is more convenient and safe for long-term use. This study used a form of microemulsion to prevent hydrolysis of aspirin because it contains a high concentration of the surfactant. A microemulsion is a dosage form that can penetrate into the skin for transdermal delivery.

Objectives: The aim of this research was to evaluate the effect of Tween 80 and propylene glycol as the surfactant and cosurfactant on the physical stability of the microemulsion.

Materials and Methods: Various concentrations of Tween 80 and propylene glycol (2:1) were used 54%, 57%, 60%, and 63%, and the physical stability of the different microemulsions was tested for 6 weeks.

Result: The results showed that the formula F3 was the most stable formula. The formula F3 showed the following properties such as pH of 3.74±0.30, viscosity of 1198.76±56.02 cps, BJ of 1.0669±0.005 g/mL, surface tension of 38.77±0.43 dyne/cm, and particle size of 49.46±6.91 nm.

Conclusions: Based on the results concluded that the optimum concentration of Tween 80 as the surfactant and propylene glycol as the cosurfactant with a ratio of 2:1 was 60%.

Keywords: Aspirin, Transdermal, Microemulsions, Tween 80.

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INTRODUCTION

Aspirin is a nonsteroidal anti-inflammatory drug that is efficient as an antiplatelet because it can inhibit the activity of the enzyme cyclooxygenase in platelets. Orally administered aspirin has side effects such as the risk of gastrointestinal bleeding and has presystemic metabolism in the gastrointestinal tract and liver. This situation is exacerbated by the continued use of aspirin that causes serious side effects in the gastrointestinal tract. In view of the poor effectiveness of oral administration, a transdermal drug delivery system is an alternative method that can improve the effectiveness of aspirin use [1,2]. Transdermal delivery by passes the gut and may, therefore, be more convenient and safe for long-term use. A microemulsion is a dosage form used for penetration in transdermal delivery. It contains a surfactant, so it is fine when formulated with aspirin because aspirin is a drug that can be hydrolyzed in the presence of water. The presence of a surfactant can prevent hydrolysis of aspirin by protecting it in micelles that can obstruct the entry of hydroxyl groups that attack the ester groups [3].

The surfactants used in this study were Tween 80. Surfactants help in forming oil-in-water microemulsions with a cosurfactant, which improves the solubilization. Short to medium chain length alcohols (C3-C8) are commonly added as cosurfactants, which further reduce the interfacial tension and fluidity of the interface. This study used propylene glycol as the cosurfactant. The aim of this study was to investigate the effect of Tween 80 and propylene glycol as a surfactant and cosurfactant on the physical properties of aspirin microemulsions.

MATERIALS AND METHODS

Materials

Aspirin was purchased from Yixing City, Xingyu. Virgin coconut oil was purchased from LIPI Cibinong, Bogor. Tween 80 was purchased from PT. KAO Chemical. Propylene glycol was purchased from PT. Dow Chemical Pacific. Methylparaben and Propylparaben were purchased from PT. Clariant.

Pseudoternary phase diagram construction

For preparing the microemulsion first, preliminary experiments were first carried out using pseudoternary phase diagrams. In the preliminary experiment, the microemulsion formula was optimized with virgin coconut oil as the oil phase, a combination of Tween 80 and propylene glycol (2:1) as the surfactant and cosurfactant, and distilled water as the aqueous phase.

Preparation of aspirin microemulsions

Various microemulsions were chosen from the pseudoternary phase diagrams. The concentration of virgin coconut oil was 5%, and combination (54%, 57%, 60%, and 63%) of Tween 80, and propylene glycol (2:1) concentrations were used. Aspirin was dispersed in the virgin coconut oil as the oil phase and then added to a mixture of distilled water and Tween 80. Then, propylene glycol was added to form a clear and homogeneous mixture (Table 1).

Organoleptic

Organoleptic includes observation of the shape, clarity, and smell. Observations were carried out for 6 weeks, on weeks 0, 3, and 6.

Determination of pH

pH measurement was performed using a pH meter, and observations were carried out for 6 weeks, on weeks 0, 3, and 6, at room temperature [4].

Density measurement

Density was measured using a pycnometer. Observations were carried out for 6 weeks, on weeks 0, 3, and 63.

Viscosity measurement

Measurements were carried out using a Brookfield viscometer LVDV-E with spindle number 3 at a speed of 30 rpm for 6 weeks, on weeks 0, 3, and 6.

6

Particle size measurements

The particle size was measured using a nanoparticle size analyzer. The solution was placed in a cuvette that was inserted into the sample holder.

Surface tension measurement

This measurement was carried out for 6 weeks, every week to 0, 3, and 6 at room temperature. The surface tension was measured using a tensiometer with Du Noüy method [5].

Phase separation

The aspirin microemulsions were centrifuged at 3750 rpm for 5 hrs at room temperature, and then, the phase changes were observed. The aspirin microemulsions were also subjected to a freeze-thaw test at 4°C and 45°C to observe the phase separation that occurs.

Data analysis

The data were statistically tested included with two-way analysis of variance (ANOVA) with a 95% confidence level, to determine significant differences between the formula test results.

RESULTS AND DISCUSSION

Preparation of aspirin microemulsions

The results showed that the formula produces a clear microemulsion is virgin coconut oil (5%) as the oil phase, a combination of Tween 80 and propylene glycol as the surfactant and cosurfactant (2:1) (54%, 57%, 60%, and 63%), and distilled water up to 100%. The microemulsion could be held steady during the observation time of 1 week.

Organoleptic

Based on organoleptic observations for 6 weeks, only the F1 formula was physically opaque since the 1st week. In contrast, the formulas F2, F3, and F4 showed no change, indicating that these three formulas have good stability during storage (Table 2).

Density measurement

The 6 weeks observations revealed that a higher surfactant concentration leads to a higher density (Fig. 1). Further, with a higher surfactant concentration, the solubilization process will increase owing to the increase in the micellar concentration [3]. This makes the microemulsions clear, with smaller particles, so that the density increases.

Determination of pH

Results of pH measurement for 6 weeks showed a decrease in pH during storage (Fig. 2). A higher surfactant concentration led to higher pH of the microemulsions. This probably because the formation of micelles with higher surfactant concentration also increased, and thus, more of the aspirin can be protected in the micelles and hydrolysis can be decreased. In contrast, for F1 and F2 with a smaller concentration of surfactant, much of the aspirin was hydrolyzed to salicylic acid and acetic acid, thus decreasing the pH of the microemulsions.

Surface tension measurement

The measurement results showed that with increasing surfactant and cosurfactant concentrations, the surface tension is lower (Fig. 3). Theoretically, increases in the surfactant concentration will cause a

Table 1: Formula of aspirin microemulsions

Materials	Formula (%)				Function
	F1	F2	F3	F4	
Aspirin	3	3	3	3	API
Virgin coconut oil	5	5	5	5	Oil phase
Tween 80	36	38	40	42	Surfactant
Propylene glycol	18	19	20	21	Cosurfactant
Nipagin	0.18	0.18	0.18	0.18	Preservative
Nipasol	0.02	0.02	0.02	0.02	Preservative
Distilled water ad	100	100	100	100	Water phase

API: Active pharmaceutical ingredient

Table 2: Organoleptic observations of aspirin microemulsions

Formula	Organoleptic	Time (week)		
		0	3	6
F1	smell	sp	sp	sp
	clarity	o	o	o
	shape	flu	flu	flu
F2	smell	sp	sp	sp
	clarity	c	c	c
	shape	flu	flu	flu
F3	smell	sp	sp	sp
	clarity	c	c	c
	shape	flu	flu	flu
F4	smell	Sp	sp	sp
	clarity	c	c	c
	shape	flu	flu	flu

flu: Fluidity, sp: Specific; o: Opaque, c: Clear

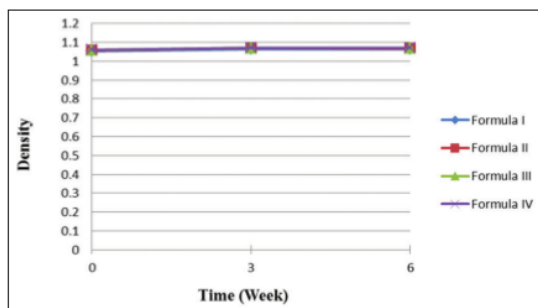


Fig. 1: Graph of density measurement

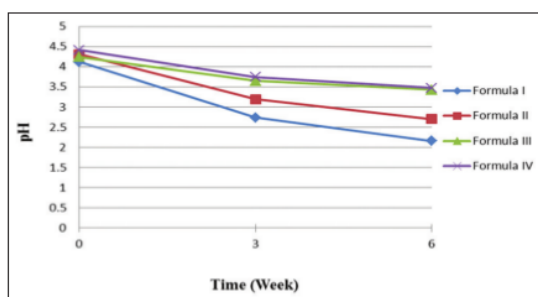


Fig. 2: Differences in pH of four formulas within 6 weeks storage

decrease in the surface tension [6]. From the graph, we can see that the formula F3 tends to be more stable than the other formulas.

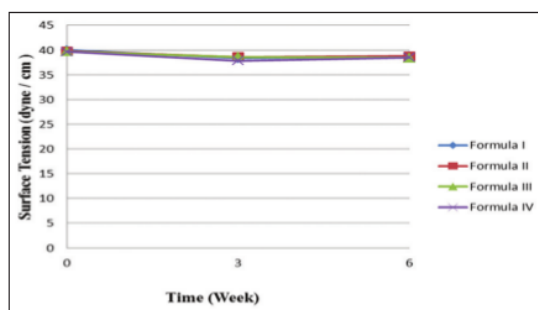


Fig. 3: Graph of surface tension measurement

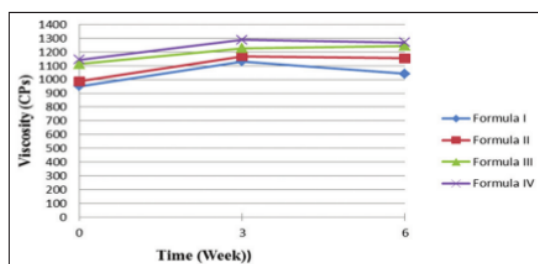


Fig. 4: Graph of viscosity measurements

Viscosity measurements

The viscosity measurement results showed that with increasing surfactant concentration, the viscosity of the microemulsion also increases (Fig. 4). Theoretically, with higher emulsifier concentration, the viscosity also increases [6]. From the graph, it can be seen that the formula F3 tends to be more stable than the others.

Phase separation

Results of the centrifugation test conducted at a speed of 3750 rpm for 5 hrs are listed in Table 3. From these observations, only the F1 formula exhibits separation with the formation of creaming. For F2, F3, and F4, no occurrence of separation was observed, which indicates that these are fairly stable microemulsions. Centrifugation at 3750 rpm for 5 hrs is equivalent to the effect of gravity for about 1 year [3]. Results of the freeze-thaw cycles conducted for 6 cycles are listed in Table 4.

At a temperature of 45°C, the F1 formula underwent phase separation in the 1st cycle, as seen with the formation of creaming at the top of the microemulsion. For formula F2, phase separation occurs in the 3rd cycle, as also seen with the formation of creaming at the top of the microemulsion. For formulas F3 and F4, there is no visible occurrence of phase separation. This suggests that these formulas are stable to changes in extreme temperatures.

6

Particle size measurement

The particle size of the F3 microemulsion was measured, which is considered the most stable as seen from the evaluation results of organoleptic, pH, viscosity, surface tension, freeze-thaw, and centrifugation tests. The results of the particle size measurement are presented in Table 5.

The results show that the F3 formula particles had an average diameter of 49.46±6.91 nm. This is proved that the F3 aspirin microemulsion met

Table 3: Centrifugation of aspirin microemulsions

Formula	Speed (3750 rpm)
F1	+
F2	-
F3	-
F4	-

(+): The separation occurs, (-): The separation does not occur

Table 4: Results of freeze-thaw cycles

Cycles	Temperature	F1			F2			F3			F4		
		A	B	C	A	B	C	A	B	C	A	B	C
1	4°C	-	-	-	-	-	-	-	-	-	-	-	-
	45°C	+	+	+	-	-	-	-	-	-	-	-	-
2	4°C	+	+	+	-	-	-	-	-	-	-	-	-
	45°C	+	+	+	+	+	+	-	-	-	-	-	-
3	4°C	+	+	+	+	+	+	-	-	-	-	-	-
	45°C	+	+	+	+	+	+	-	-	-	-	-	-
4	4°C	+	+	+	+	+	+	-	-	-	-	-	-
	45°C	+	+	+	+	+	+	-	-	-	-	-	-
5	4°C	+	+	+	+	+	+	-	-	-	-	-	-
	45°C	+	+	+	+	+	+	-	-	-	-	-	-
6	4°C	+	+	+	+	+	+	-	-	-	-	-	-
	45°C	+	+	+	+	+	+	-	-	-	-	-	-

(+): The separation occurs, (-): The separation does not occur

Table 5: Particle size measurement of aspirin microemulsion

Formula	Particle size diameter (nm)	PI
F3	42.8	11.967
	56.6	14.995
	49.0	18.309
Average	49.46±6.91	15.09±3.17

PI: Polydispersity index

the particle size requirements. However, more research is needed on the pH of the aspirin microemulsions, to obtain the ideal pH preparations. It is also necessary to test the physical and chemical stability of the aspirin microemulsion.

CONCLUSION

The results of this research indicated that the optimal concentration of Tween 80 as a surfactant and propylene glycol as a cosurfactant (2:1) is 60%. This microemulsion formula exhibited a pH value of 3.74±0.30 pH, viscosity of 1198.76±56.02 cps, BJ of 1.0669±0.005 g/mL, surface tension of 38.77±0.43 dyne/cm, and particle size of 49.46±6.91 nm.

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