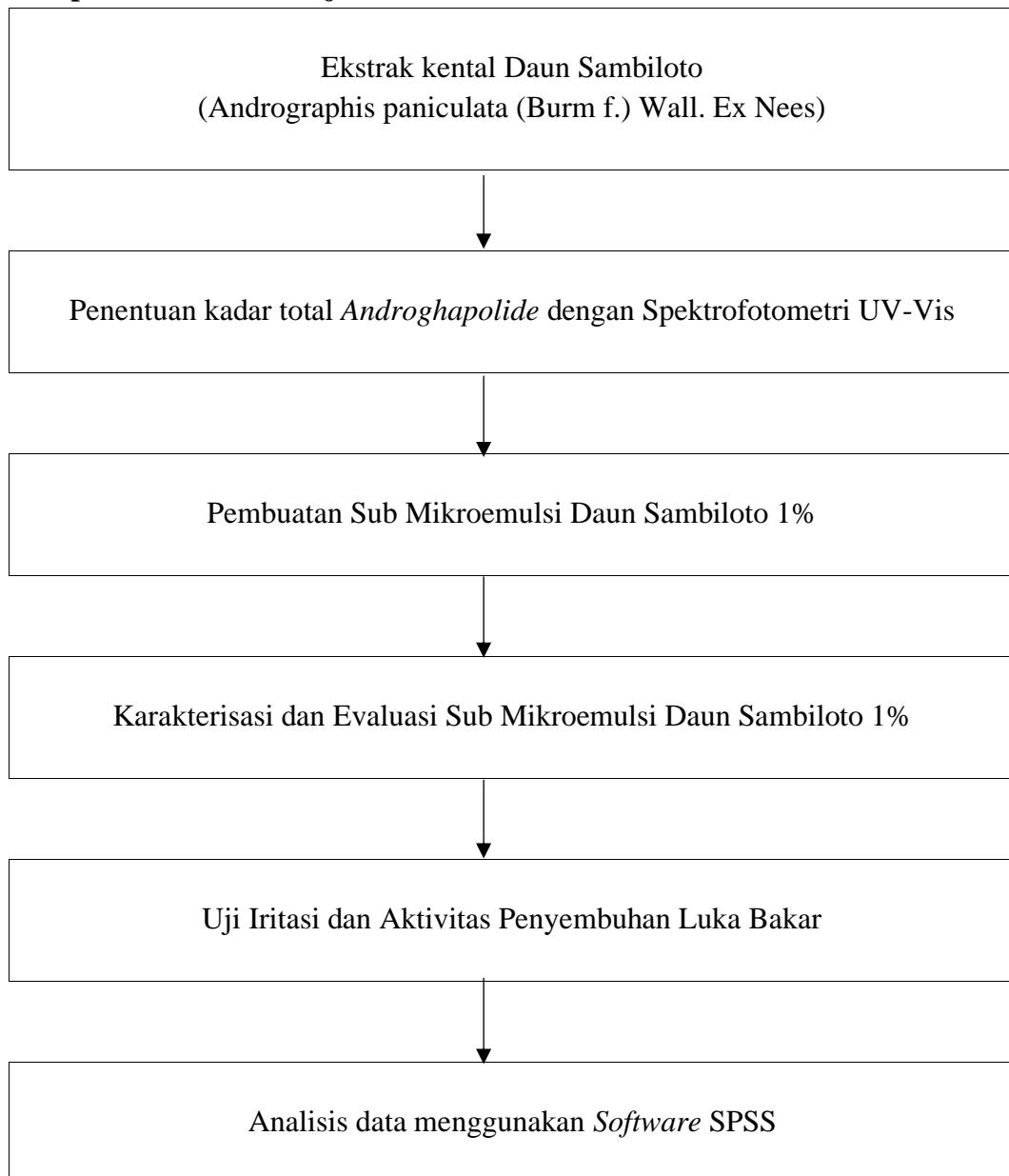


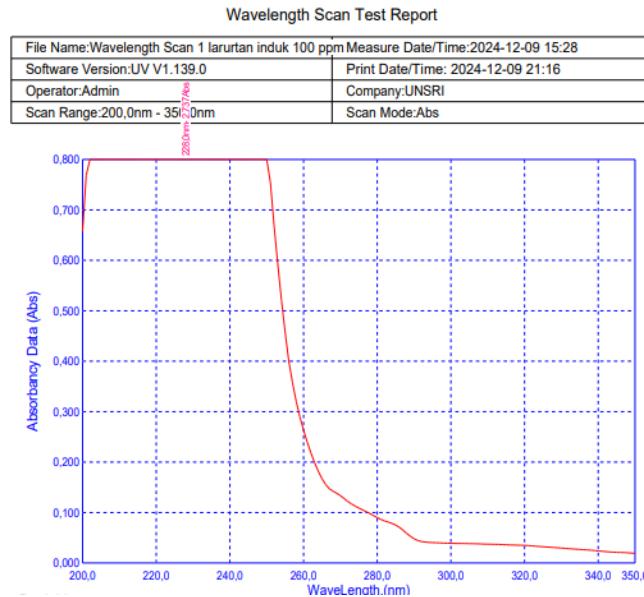
## LAMPIRAN

### Lampiran 1. Skema Kerja Umum



## Lampiran 2. Penentuan Kurva Kalibrasi *Androghapolide*

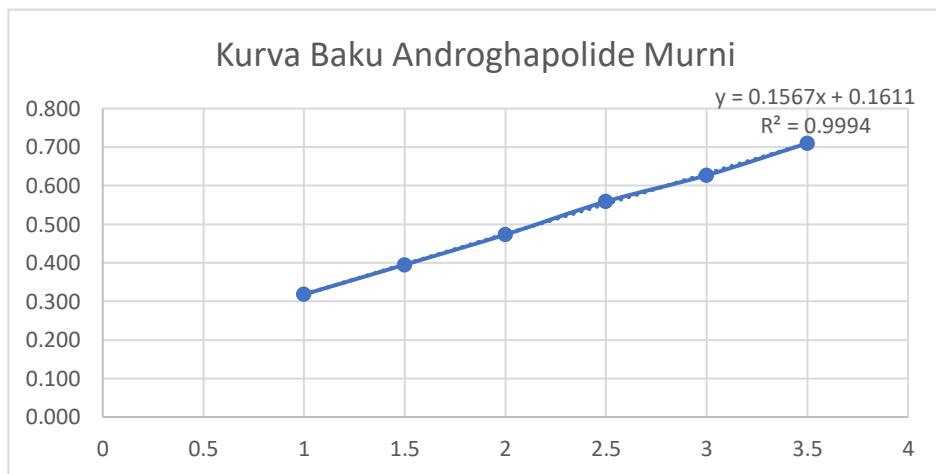
### 1. Panjang Gelombang Maksimum 228 nm



### 2. Tabel Hasil Pembuatan Kurva Baku

Konsentrasi (ppm)	Absorbansi
1	0,318
1,5	0,395
2	0,473
2,5	0,559
3	0,627
3,5	0,710

### 3. Kurva Kalibrasi *Androghapolide*



### Lampiran 3. Perhitungan Kadar Total *Androghapolide Content* Ekstrak

Replikasi	Absorbansi	Faktor Pengenceran	Rata – Rata ± SD	Kadar mg/gr ± SD
1	0.673			
2	0.673	100	0.6733 ± 0,0005	32,688±0,030
3	0.674			

**Persamaan regresi linier pada kurva baku *Androghapolide*:**

$$y = 0,1567x + 0,1611$$

Keterangan:

x = Kadar *Androghapolide*

y = Absorbansi rata – rata

#### Perhitungan kadar *Androghapolide*

$$x = \frac{y - 0,1611}{0,1567}$$

$$x = \frac{0,6733 - 0,1611}{0,1567}$$

$$x = 3.266 \mu\text{g/mL} \text{ konversi mg/mL} = 0,003266 \text{ mg/mL}$$

$$\text{Kadar } Androghapolide \text{ Total} = \frac{x \frac{\text{mg}}{\text{L}} \times \text{volume}}{\text{Gram Ekstrak (g)}} \times \text{FP}$$

$$= \frac{0,003266 \times 10}{0,01} \times 100$$

$$= 326,6751 \text{ mg/gr}$$

1 gram ekstrak etanol daun sambiloto mengandung 326,6751 mg *Androghapolide*

$$\text{Persentase Kadar} = \frac{326,6751}{1000} \times 100\%$$

$$= 32,6675 \%$$

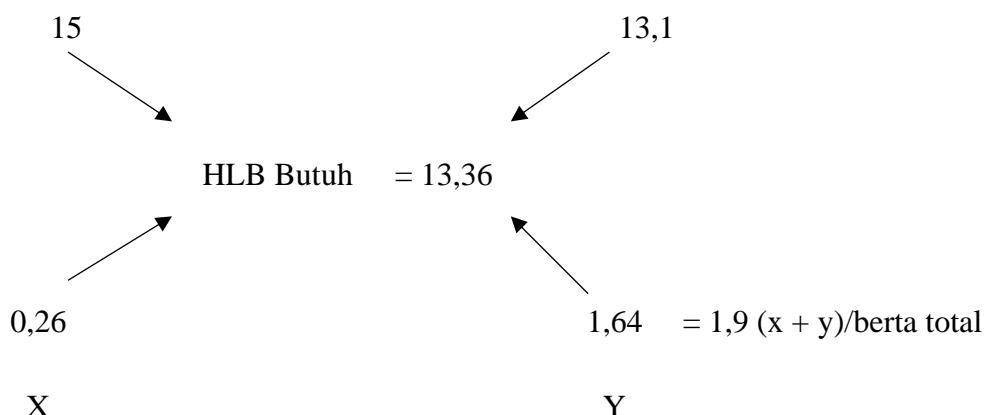
**Lampiran 4. Perhitungan Nilai HLB Sub Mikroemulsi Ekstrak Daun Sambiloto**

HLB Tween 80 = 15

HLB PEG-400 = 13,1

HLB Butuh minyak atsiri daun kemangi = 13,36

Tween 80 PEG-400



HLB Campuran x berat total = (HLB a. x) + (HLB b.y)

HLB Campuran x 1,9 =  $(15 \times 0,26) + (13,1 \times 1,64)$

HLB Campuran x 1,9 =  $3,9 + 21,484$

HLB Campuran x 1,9 = 25,384

HLB Campuran =  $\frac{25,384}{1,9}$

HLB Campuran = 13,36

### Lampiran 5. Perhitungan Kadar *Androghapolide* dalam Sediaan

Replikasi	Absorbansi	X	Kadar	Rerata Kadar	% Kadar	Rerata ± SD
1	0.697	3.420	63,449		19,410	
2	0.706	3.477	64,515	64,199	19,736	19,639±0,163
3	0.707	3.484	64,633		19,772	

Diketahui

Persamaan regresi linier pada kurva baku *Androghapolide* (**Lampiran 2**)

#### Perhitungan kadar *Androghapolide*

$$x = \frac{y - 0,1611}{0,1567}$$

$$x = \frac{0,697 - 0,1611}{0,1567}$$

$$x = 3,420 \text{ ppm}$$

$$\begin{aligned} \text{Kadar } Androghapolide \text{ Total} &= \frac{x \frac{\text{mg}}{\text{L}} \times \text{volume sampel (L)}}{5\text{ml} \times \text{bobot jenis (g)}} \times \text{FP} \\ &= \frac{3,420 \times 0,05 \text{ L}}{5,3925} \times 2000 \\ &= 63,449 \text{ mg/gr} \end{aligned}$$

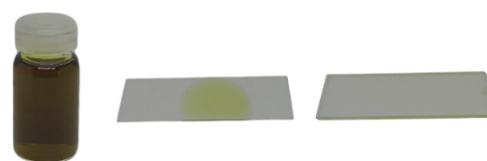
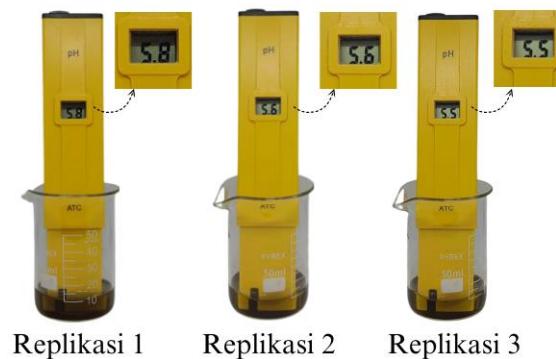
5 mL Sub mikroemulsi mengandung 63,449 mg/gr *Androghapolide*

$$\begin{aligned} \text{Persentase Kadar} &= \frac{\text{Kadar yang didapat}}{\text{Kadar TAC}} \times 100\% \\ &= \frac{63,449}{326,675} \times 100\% \\ &= 19,410\% \end{aligned}$$

$$\text{Persentase Kadar Sebenarnya} = \text{Kadar sediaan} \times \text{Hasil \%EE}$$

$$= 19,410\% \times 0,99\%$$

$$= 0,1909\% \text{ (Kadar 1x pengolesan)}$$

**Lampiran 6. Hasil Evaluasi Karakteristik Sub mikroemulsi****1. Uji Organoleptis****2. Uji pH Submikroemulsi**

### 3. Uji Viskositas



**LABORATORIUM  
KIMIA PENGOLAHAN DAN SENSORIS HASIL PERTANIAN  
JURUSAN TEKNOLOGI PERTANIAN  
FAKULTAS PERTANIAN**

LAPORAN ANALISA  
No. 90/LABKHP/X/2024

Nama Pemesan	: Dian Arifando Rusyadi
Tanggal Terima	: 15 Oktober 2024
Tanggal Selesai	: 24 Oktober 2024
Jumlah Sampel	: 3
Jenis Analisa	: Viskositas (Viskometer NDJ-8S)
Jenis Sampel	: Submikro Emulsi Ekstrak Daun Sambiloto

No	Kode	Viskositas (m.Pa.s)
1	1	11,10
2	2	12,50
3	3	12,86

Palembang, 24 Oktober 2024  
Koordinator Teknis Laboratorium,

Hafsa, ST.,M.T  
NIP.198006202001122001

#### 4. Uji Bobot Jenis



Berat Piknometer + Air

Replikasi	W1	W0	Massa Akhir	Rerata
1	17,3504	12,3354	5,015	
2	17,3503	12,3354	5,0149	5,0148
3	17,3501	12,3354	5,0147	

Keterangan;

W0 : Piknometer Kosong ; W1 : Piknometer Terisi

Berat Piknometer + Sub mikroemulsi 1%

Replikasi	W1	W0	Massa Akhir	Rerata
1	17,8014	12,3354	5,466	
2	17,809	12,3354	5,4736	5,42466
3	17,6698	12,3354	5,3344	

Perhitungan

	Massa				
Massa	Massa	Piknometer +	Densitas	Berat Jenis	
Piknometer	Piknometer +	Sub	Aquademineral	Sub	
Kosong	Aquademineral	Mikroemulsi		Mikroemulsi	
		1%		1%	
12,3354	5,0148	5,42466	0,997	1,078	

$$\begin{aligned}
 \text{Berat Jenis} &= \frac{\text{Berat Piknometer+Sediaan}}{\text{Berat Piknometer+Aquademineral}} \times \text{Densitas Aquademineral} \\
 &= \frac{5,42466}{5,0148} \times 0,997 = 1,078 \text{ (Lebih berat dibandingkan aquademineral)}
 \end{aligned}$$

## 5. Perhitungan Efisiensi Penjerapan

Replikasi	Absorbansi	x	Kadar	%EE	Rerata ± SD
1	0,383	1,416	141,608	99,567	
2	0,381	1,403	190,560	99,571	99,567 ± 0,002
3	0,384	1,422	192,554	99,565	

### Perhitungan

Persamaan regresi linier pada kurva baku *Androghapolide* (**Lampiran 2**)

#### Perhitungan kadar Tak Terjerap

$$x = \frac{y - 0,1611}{0,1567}$$

$$x = \frac{0,383 - 0,1611}{0,1567}$$

$$x = 1,416 \text{ ppm}$$

$$\begin{aligned} \text{Kadar TakTerjerap} &= \frac{x \frac{\text{mg}}{L} \times \text{volume sampel (L)}}{\text{Bobot Sampel (g)}} \times \text{FP} \\ &= \frac{1,416 \frac{\text{mg}}{L} \times 0,00005 L}{0,005} \times 1000 \\ &= 141,608 \text{ mg/gr} \end{aligned}$$

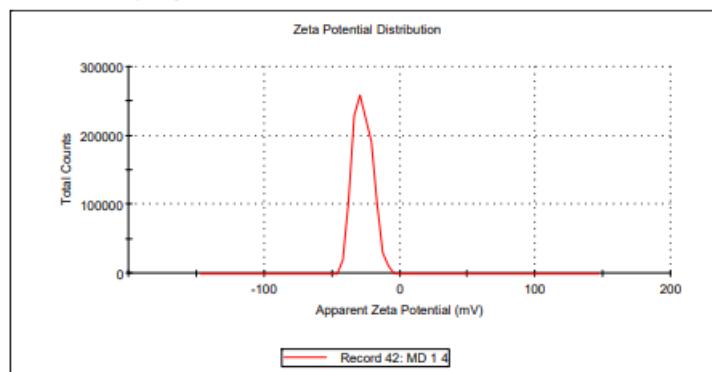
$$\begin{aligned} \% \text{ Efisiensi Penjerapan} &= \frac{\text{Kadar sediaan} - \text{kadar tak terjerap (L)}}{\text{Kadar sediaan}} \times 100\% \\ &= \frac{32688 - 141,608}{32688} \times 100 \\ &= 99,567\% \end{aligned}$$

$$\begin{aligned} \% \text{ EE Tak terjerap dalam sediaan} &= \% \text{ EE} \times \text{Jumlah kadar sediaan formula} \\ &= 99,567 \times 1\% \\ &= 0,995\% \end{aligned}$$

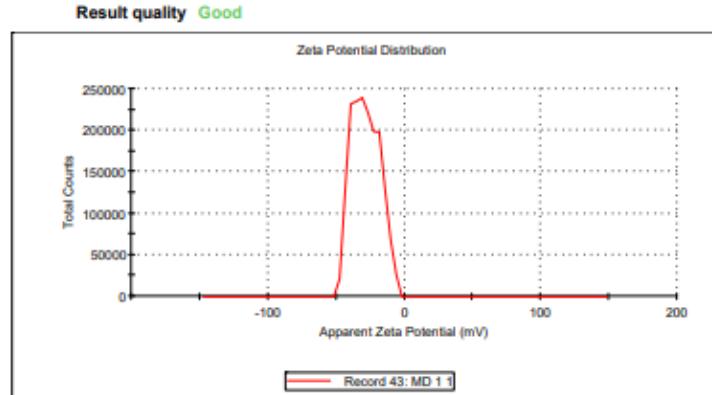
## 6. Uji PSA (Ukuran Partikel, Indeks Polidispersitas dan Zeta Potensial)

### a. Zeta Potensial

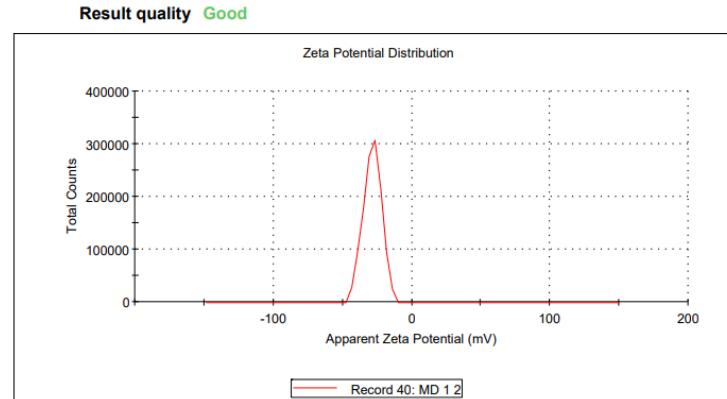
Results			
	Mean (mV)	Area (%)	St Dev (mV)
Zeta Potential (mV): -27.3	Peak 1: -27.3	100.0	6.96
Zeta Deviation (mV): 6.96	Peak 2: 0.00	0.0	0.00
Conductivity (mS/cm): 0.0188	Peak 3: 0.00	0.0	0.00
Result quality	Good		



Results			
	Mean (mV)	Area (%)	St Dev (mV)
Zeta Potential (mV): -28.2	Peak 1: -28.2	100.0	9.86
Zeta Deviation (mV): 9.86	Peak 2: 0.00	0.0	0.00
Conductivity (mS/cm): 0.0184	Peak 3: 0.00	0.0	0.00
Result quality	Good		

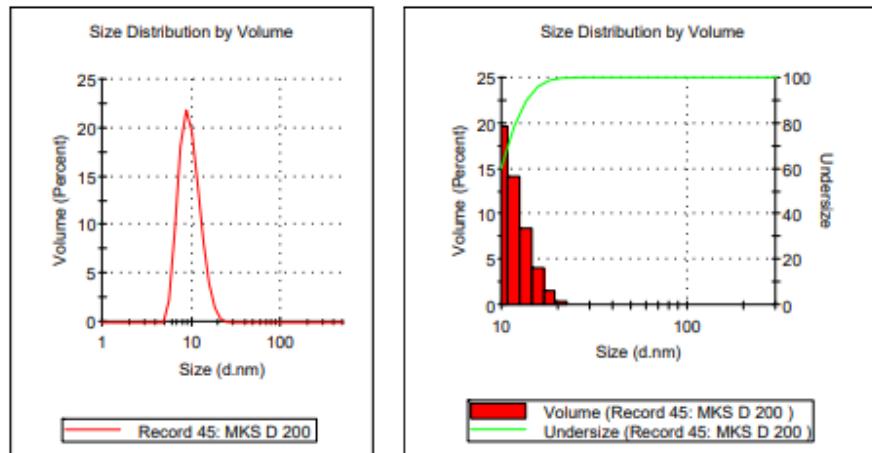


Results			
	Mean (mV)	Area (%)	St Dev (mV)
Zeta Potential (mV): -28.4	Peak 1: -28.4	100.0	6.46
Zeta Deviation (mV): 6.46	Peak 2: 0.00	0.0	0.00
Conductivity (mS/cm): 0.0191	Peak 3: 0.00	0.0	0.00
Result quality	Good		

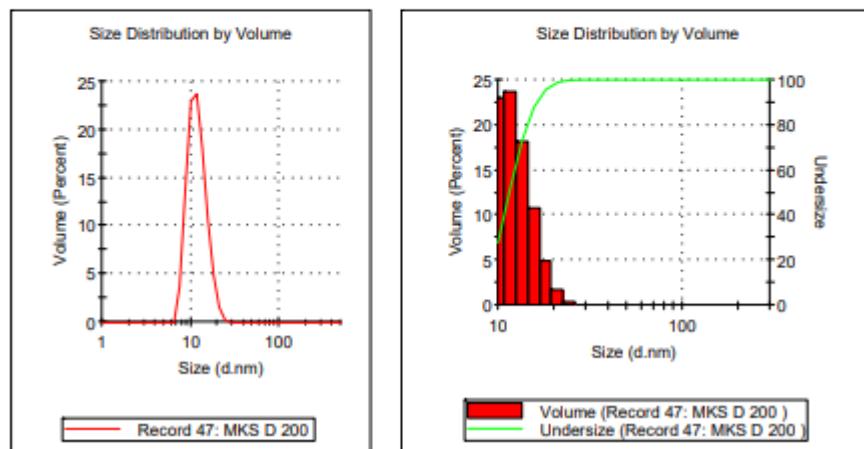


## b. Indeks Polidispersitas dan Ukuran Partikel

Cumulant Results	Distribution Results	Undersize Results
Z-Avg (nm): 61.79	Size (d.nm): % Vol $\sigma$ %Pd	Dv (%) Size (d.nm):
Pd Index: 0.217	Peak 1: 9.810 100.0 2.730 27.8	10 6.72
Pd (nm): 28.8	Peak 2: 288.2 0.0 88.32 30.6	50 9.33
%Pd: 46.6	Peak 3: 0.000 5.4 481.8 9.3	90 13.7
Derived kcps: 2024.7		100 7460



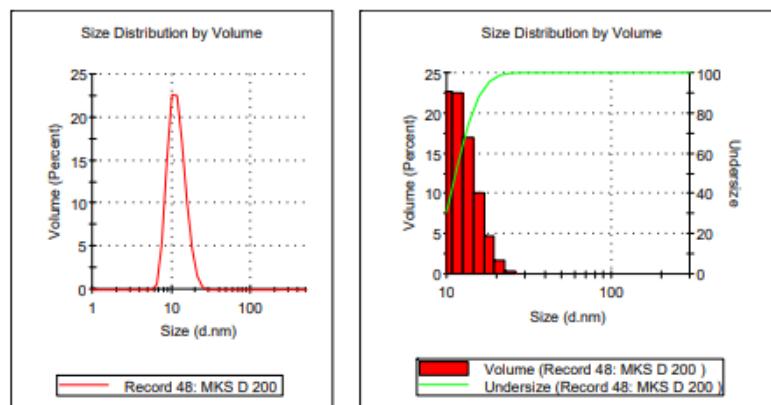
Cumulant Results	Distribution Results	Undersize Results
Z-Avg (nm): 83.50	Size (d.nm): % Vol $\sigma$ %Pd	Dv (%) Size (d.nm):
Pd Index: 0.281	Peak 1: 12.03 99.9 2.943 24.5	10 8.62
Pd (nm): 44.3	Peak 2: 421.4 0.1 124.8 29.6	50 11.5
%Pd: 53.0	Peak 3: 0.000 4.9 460.2 8.8	90 16.4
Derived kcps: 3739.3		100 7460



**Sample Details**

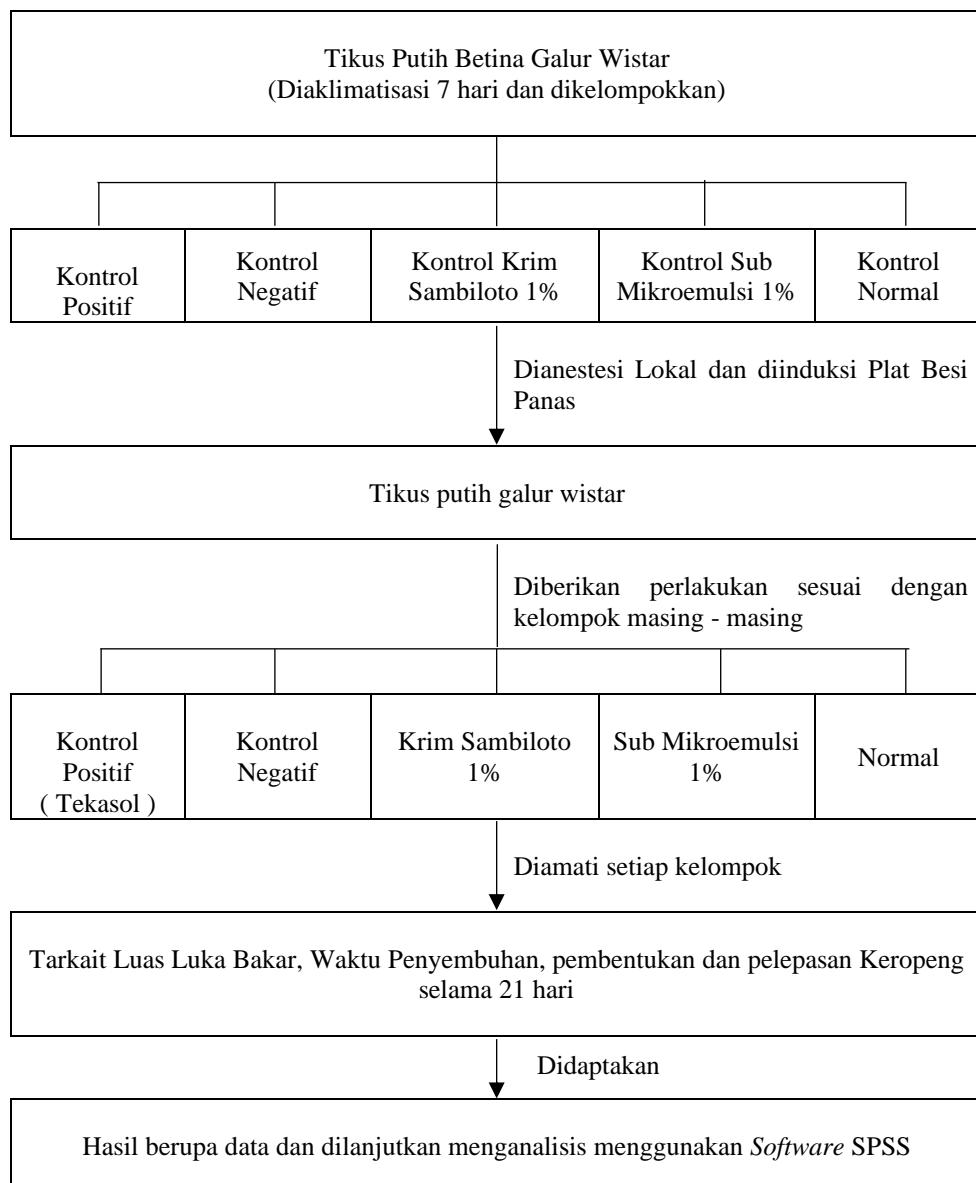
Name: MKS D 200  
 Filename: MKS.dts  
 Operator: User  
 Date and Time: Friday, September 20, 2024 11:08:46  
 SOP: mansettings.nano

<b>Cumulant Results</b>		<b>Distribution Results</b>				<b>Undersize Results</b>	
Z-Avg (nm):	146.9	Size (d.nm):	% Vol	$\sigma$	%Pd	Dv (%)	Size (d.nm):
Pd Index:	0.244	Peak 1:	11.85	99.9	3.033	25.6	10
Pd (nm):	72.6	Peak 2:	316.9	0.1	85.66	27.0	50
%Pd:	49.4	Peak 3:	0.000	2.1	6.104e-5	.0	90
Derived kcps:	4368.5					100	7460



Size d.nm	Mean Volume/Percent	Std Dev Volume/Percent	Size d.nm	Mean Volume/Percent	Std Dev Volume/Percent	Size d.nm	Mean Volume/Percent	Std Dev Volume/Percent
0.4000	0.0		13.54	17.0		468.7	0.0	
0.4632	0.0		15.69	10.1		531.2	0.0	
0.5365	0.0		18.17	4.7		615.1	0.0	
0.6213	0.0		21.04	1.6		712.4	0.0	
0.7195	0.0		24.36	0.5		825.0	0.0	
0.8332	0.0		28.21	0.0		955.4	0.0	
0.9649	0.0		32.67	0.0		1106	0.0	
1.11	0.0		37.84	0.0		1281	0.0	
1.294	0.0		43.62	0.0		1484	0.0	
1.499	0.0		50.75	0.0		1718	0.0	
1.736	0.0		58.77	0.0		1900	0.0	
2.010	0.0		68.06	0.0		2205	0.0	
2.338	0.0		78.82	0.0		2669	0.0	
2.698	0.0		91.28	0.0		3091	0.0	
3.122	0.0		105.7	0.0		3580	0.0	
3.615	0.0		122.4	0.0		4145	0.0	
4.187	0.0		141.8	0.0		4801	0.0	
4.849	0.0		164.2	0.0		5560	0.0	
5.615	0.0		190.1	0.0		6439	0.0	
6.503	0.6		220.2	0.0		7456	0.0	
7.531	5.3		255.0	0.0		8635	0.0	
8.721	15.1		295.3	0.0		1.000e4	0.0	
10.10	22.6		342.0	0.0				
11.70	22.5		396.1	0.0				

### Lampiran 7. Skema Uji Aktivitas Penyembuhan Luka Bakar



### **Lampiran 8. Perhitungan Persiapan Hewan Uji**

Penentuan hewan uji pada masing-masing kelompok dilakukan dengan menggunakan rumus Federer yang dapat dilihat dibawah ini:

$$(n-1)(t-1) \geq 15$$

$$(n-1)(5-1) \geq 15$$

$$4(n-1) \geq 15$$

$$4n-4 \geq 15+4$$

$$n \geq 19/4$$

$$n \geq 4.75$$

Berdasarkan perhitungan yang telah dilakukan menggunakan rumus, hewan uji yang akan digunakan dalam penelitian ini sebanyak 5 ekor tikus pada masing-masing kelompok.

$$\begin{aligned} \textbf{Total Hewan Uji} &= n \times t \\ &= 5 \times 5 \\ &= 25 \text{ Ekor Tikus} \end{aligned}$$

Jadi untuk penelitian ini dibutuhkan 5 ekor tikus Betina galur wistar.

### Lampiran 9. Perhitungan Dosis Anastesi Lidokain 2%

Contoh Perhitungan:

Pemberian dosis lidokain 4 mg/KgBB, berdasarkan perhitungan berikut :

$$\text{Berat badan tikus} = 160 \text{ gram} = 0,16 \text{ kg}$$

$$\text{Dosis Ledokain} = 4 \text{ mg/KgBB}$$

$$\text{Konsentrasi Lidokain 2\%} = 2\% \text{ (2grm/100ml atau } 20\text{mg/mL)}$$

Langkah perhitungan

#### Dosis Total Lidokain (dalam mg)

$$\text{Dosis lidokain (mg)} = 4\text{Mg/kgBB} \times 0,16 = 0,64 \text{ mg}$$

Konversi Dosis kedalam Volume (mL)

Rumus:

$$\begin{aligned} \text{Volume (mL)} &= \frac{\text{Dosis (mg)}}{\text{konsentrasi } (\frac{\text{mg}}{\text{mL}})} \\ &= \frac{0,64 \text{ mL}}{20} \\ &= 0,032 \text{ mL} \end{aligned}$$

Jadi masing-masing tikus yang akan dianastesi mendapatkan lidokain 2% sebanyak 0,032 mL.

## Lampiran 10. Sertifikat Persetujuan Etik



**UNIVERSITAS AHMAD DAHLAN**

**KOMITE ETIK PENELITIAN (KEP UAD)**

Jl. Prof. Dr. Soepomo, S. H, Yogyakarta Telp (0274) 563515, Ekstension 3310.

**Surat Persetujuan Etik (Ethical Approval)  
Untuk Penelitian yang Menggunakan Hewan Coba sebagai  
Subjek Penelitian**

**Persetujuan Etik (Ethical Approval)  
Nomor : 022412171**

Yang bertanda tangan di bawah ini, Ketua Komite Etik Penelitian Universitas Ahmad Dahlan, setelah dilaksanakan pembahasan dan penilaian, dengan ini memutuskan protokol penelitian yang berjudul: **"Uji Aktivitas Penyembuhan Luka Bakar Sub Mikroemulsi Ekstrak Daun Sambiloto dengan Kombinasi Tween-80 dan PEG-400 terhadap Tikus Galur Wistar"** yang menggunakan hewan coba sebagai subjek penelitian, yang diajukan oleh:

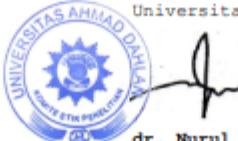
Ketua Peneliti: **apt. Vitri Agustiarini, M. Farm.**

Anggota: **Dian Arifando Rusyadi**

dapat disetujui pelaksanaannya. Persetujuan ini berlaku selama 1(satu) tahun setelah Ethical Approval dikeluarkan.

Pada akhir penelitian, laporan pelaksanaan penelitian harus diserahkan kepada KEP UAD. Jika ada perubahan protokol dan/atau perpanjangan penelitian, harus mengajukan kembali permohonan kajian etik penelitian (amandemen protokol).

Yogyakarta, 27 Desember 2024  
Komite Etik Penelitian  
Universitas Ahmad Dahlan,



dr. Nurul Qomariyah, M.Med., Ed.

**Lampiran 11. Sertifikat Hewan Uji**



**TIKUS PENELITIAN PALEMBANG**

Jalan Seduduk Putih No. 89 RT. 27 RW. 07  
Kec. Ilir Timur II Kel. 8 Ilir Kota Palembang Sumatera Selatan  
Telp. 0813-5068-3378 – 081278811795

**SURAT KETERANGAN**  
Nomor: 004/TPP-010/2024

Yang Bertanda tangan dibawah ini:

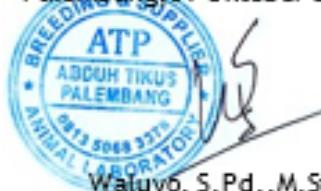
Nama : Dian Arifando Rusyadi  
NIM : 08061182126014  
Jurusan : Farmasi  
Fakultas : Matematika dan Ilmu Pengetahuan Alam  
Universitas : Universitas Sriwijaya  
Judul Skripsi : Uji Aktivitas penyembuhan luka bakar Sub Mikroemulsi Ekstrak Daun Sambiloto dengan kombinasi PEG 400 dan Tween 80 Terhadap Tikus Galur Wistar.

Menerangkan hewan uji dengan spesifikasi:

Galur : Wistar  
Jumlah : 35 ekor  
Jenis Kelamin : Betina  
Berat Badan : 150-250  
Umur : 3 bulan

Demikian surat keterangan ini dibuat untuk dapat digunakan sebaik-baiknya.

Palembang, 01 Oktober 2024



Waluyb, S.Pd., M.Si

### Lampiran 12. Certificate of Analysis Andrographolide

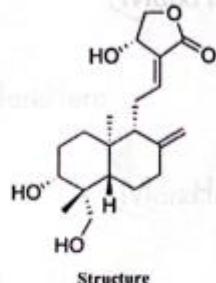


EBM Scientific and Technology, Ltd. (EBM Scitech)  
Jl. Bukit Raya No. 582, Ciumbuleuit, Cidadap, Bandung City 40142,  
West Java, Indonesia  
[www.markherb.com](http://www.markherb.com)

#### CERTIFICATE OF ANALYSIS

*For Laboratory Use Only*

Product name	Andrographolide
Other names	(3E,4S)-3-[2-((1R,4aS,5R,6R,8aS)-6-hydroxy-5-(hydroxymethyl)-5,8a-dimethyl-2-methylidene-3,4,4a,6,7,8-hexahydro-1H-naphthalen-1-yl)methylidene]-4-hydroxyoxolan-2-one ; Andrographolide; Andrographis; Andrographolida (INA)
CAS	5508-58-7
Catalog No.	TER-1-50   TER-1-100
Lot Number	# 190209
Formula	C <sub>21</sub> H <sub>22</sub> O <sub>5</sub>
Molecular mass	350.45 g/mol
Date of sample	13/06/2022
Date of report	16/06/2022



Structure

#### ANALYTICAL RESULTS

Test	Method	Specification	Result
Appearance	NA	White, powder	White, powder
Purity	UPLC-PDA	> 90%	93.22%
UV spectrum	UPLC-PDA	230 nm	230 nm
Mass spectrum	UPLC-TQD MS	Conforms	Conforms
NMR	<sup>1</sup> H-NMR, <sup>13</sup> C-NMR	NA	NA

NA, not available

#### STORAGE CONDITIONS

Storage	Room temperature in a dry place, keep away from direct light
Transport	Room temperature

Digitally signed by  
Andi Rifki Rosandy, Ph.D  
Date: 18.06.2022  
Bandung - Indonesia

### Lampiran 13. Certificate of Analysis Minyak Kemangi

 *Plus que des analyses... des conseils*

Date : February 26, 2018

**CERTIFICATE OF ANALYSIS - GC PROFILING**

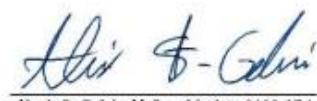
**SAMPLE IDENTIFICATION**

**Customer Identification :** Holy Basil - Tulsi - 3031  
**Type :** Essential oil  
**Source :** *Ocimum sanctum*

**ANALYSIS**

**Method:** PC-PA-014-17J19 - Analysis of the composition of an essential oil, or other volatile liquid, by FAST GC-FID (in French); identifications validated by GC-MS.  
**Analyst :** Sylvain Mercier, M. Sc., Chimiste  
**Analysis date :** February 22, 2018

Checked and approved by :

  
 Alexis St-Gelais, M. Sc., chimiste 2013-174

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**Lampiran 14. Certificate of Analysis PEG-400**



**MATERIAL SAFETY DATA SHEET**

Produced date: 2012-08-10  
Replaces date: 2012-04-16

**1. Identification of the substance/preparation and the company**

<b>Commercial product name</b>	Polyethylene glycol 400, PEG 400
<b>Use of the substance/preparation</b>	Petroleum chemicals, coatings, lubricants, plastics, inks, detergents, cleaning agent
<b>Synonyms</b>	Ethylene oxide polymer, Poly(oxyethylene) glycol
<b>CAS-nr</b>	25322-68-3
<b>EC-nr</b>	500-038-2
<b>REACH registration number</b>	05-2114142277-51-0000
<b>Company</b>	Fred Holmberg & Co AB
<b>Address</b>	Box 60056 S-216 10 Limhamn Sweden
<b>Telephone number</b>	+46 (0)40 15 79 20
<b>Fax</b>	+46 (0)40 16 22 95
<b>e-mail</b>	fred.info@holmberg.se
<b>Contact person</b>	Fred Holmberg
<b>Emergency telephone number</b>	Fred Holmberg 040-15 79 20 (office hours) or. 08-33 12 31 toxicity information central (office hours), 112 for emergency central
<b>Created by</b>	Linus Olofsson, Fred Holmberg & Co AB, Tel. +46 (0)480-42 20 00

**2. Hazards Identification**

This product is not classified as dangerous according to EC criteria.

**3. Composition/Information on ingredients**

<b>EC-nr</b>	<b>CAS-nr</b>	<b>Substance</b>	<b>Conc. % (w/w)</b>	<b>Classification</b>
500-038-2	25322-68-3	Polyethylene glycol	100	Not classified as hazardous material.

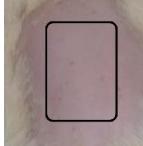
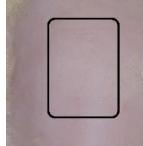
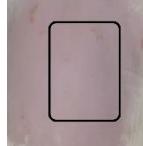
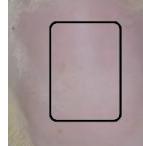
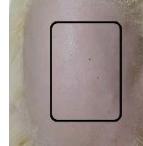
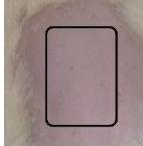
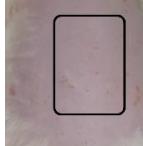
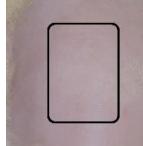
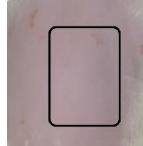
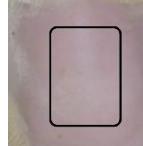
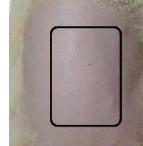
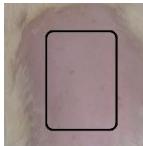
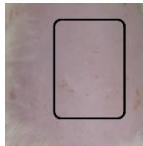
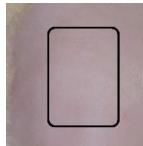
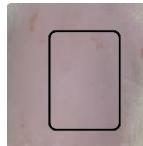
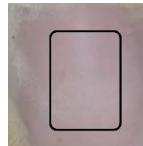
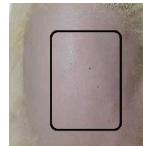
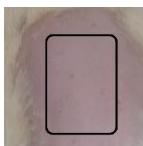
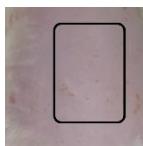
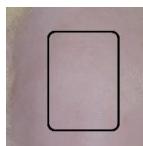
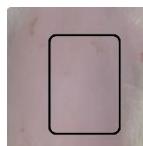
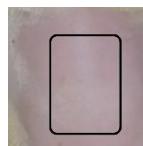
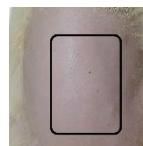
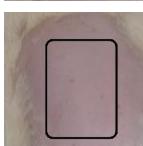
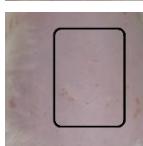
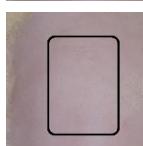
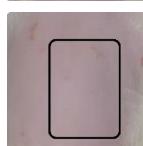
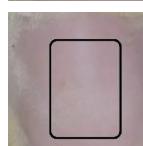
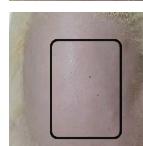
**Lampiran 15. Perhitungan Derajat Indeks Iritasi Primer**

**Indeks Iritasi** =  $\frac{\text{Jumlah eritemia } 0;1;24;48;72 \text{ jam} + \text{jumlah edema } 0;1;24;48;72 \text{ jam}}{\text{jumlah tikus}}$

$$\text{Tikus 1} = \frac{0+0}{3} = 0$$

$$\text{Tikus 2} = \frac{0+0}{3} = 0$$

**Lampiran 16. Gambar Pengamatan Iritasi Primer**

Jam	Kelompok Tikus					
	Tikus Normal			Tikus Sub Mikroemulsi 1%		
	1	2	3	1	2	3
0						
1						
24						
48						
72						



**Lampiran 17. Gambar Pengamatan Perubahan Rerata Luka Bakar**

Kelompok Tikus	Pengamatan Luka Hari Ke										
	0	2	4	6	8	10	12	14	16	18	20
Kontrol Positif											
Kontrol Negatif											
Krim Sambiloto 1%											
Sub Mikroemulsi 1%											

**Lampiran 18. Data Luas Luka Bakar Derajat 2**

Kelompok Tikus	Luas Luka Awal (cm <sup>2</sup> )	Rerata Luas Luka Awal (cm <sup>2</sup> )	Luas Luka Akhir (cm <sup>2</sup> )	Penurunan Luas Luka (cm <sup>2</sup> )	Rerata penurunan luas luka (cm <sup>2</sup> ) ± SD	Persentase Penyembuhan (%)	Rerata Persentase Penyembuhan (%)
Kontrol Positif	6.2		0.44	5.76		92.90	
	6.08	6.193333333	0.8	5.28	5.613 ± 0.236	86.84	90.60
Kontrol Negatif	6.3		0.5	5.8		92.06	
	6.3	6.543333333	2.3	4		63.49	
Kontrol Krim Sambiloto 1%	6.51		2.2	4.31	4.11 ± 0.141	66.20	62.88
	6.82		2.8	4.02		58.94	
Kontrol Sub Mikroemulsi 1%	6.09		1.05	5.04		82.75	
	6.51	6.4	0.78	5.73	5.54 ± 0.356	88.01	86.47
Kontrol Sub Mikroemulsi 1%	6.6		0.75	5.85		88.63	
	6		0.44	5.56		92.66	
Kontrol Sub Mikroemulsi 1%	5.51	6.236666667	0.5	5.01	5.763 ± 0.712	90.92	92.30
	7.2		0.48	6.72		93.33	

### Lampiran 19. Hasil Analisis Statistika Luas Luka Bakar Derajat Dua

#### *Paired Sampel T Test*

Hipotesis :

$H_0$  = Data penurunan luas luka bakar tidak berbeda signifikan

$H_a$  = Data penurunan luas luka bakar berbeda signifikan

Pengambilan Keputusan :

Jika nilai signifikansi  $> 0,05$   $H_0$  diterima

Jika nilai signifikansi  $< 0,05$   $H_0$  ditolak

Kelompok Positif Hari ke-1 dan Hari Ke-20

Paired Samples Test										
Paired Differences										
	Mean	Std. Deviation	Std. Error	95% Confidence Interval of the Difference		t	df	Sig. (2-tailed)		
Pair 1	Positif_Awal	-	5.61333	0.28937	0.16707	4.89451	6.33216	33.599	2	0.001
	Positif_Akhir									

Kesimpulan:

1. Karena Sig. (2-tailed) = 0.001  $< 0.05$ , maka  $H_0$  ditolak.
2. Artinya, terdapat perbedaan yang signifikan antara data awal dan data akhir pada kelompok positif.

Kelompok Negatif Hari ke-1 dan Hari ke-20

<b>Paired Samples Test</b>								
Paired Differences								
	Mean	Std. Deviation	Std. Error Mean	95% Confidence Interval of the Difference		t	df	Sig. (2-tailed)
Pair 1	Negatif_Awal							
	- Negatif_Ahir	4.11000	0.17349	0.10017	3.67902	4.54098	41.032	2 0.001

Kesimpulan:

1. Karena Sig. (2-tailed) = 0.001 < 0.05, maka Ho ditolak.
2. Artinya, terdapat perbedaan yang signifikan antara data awal dan data akhir pada kelompok negatif Kelompok Krim Sambiloto 1% Hari ke-1 dan Hari ke-20

<b>Paired Samples Test</b>								
Paired Differences								
	Mean	Std. Deviation	Std. Error Mean	95% Confidence Interval of the Difference		t	df	Sig. (2-tailed)
Pair 1	Krim_Sambiloto_1%_Awal							
	- Krim_Sambiloto_1%_Akhir	5.54000	0.43715	0.25239	4.45406	6.62594	21.950	2 0.002

Kesimpulan:

1. Karena Sig. (2-tailed) = 0.002 < 0.05, maka Ho ditolak.
2. Artinya, terdapat perbedaan yang signifikan antara data awal dan data akhir pada kelompok Krim Sambiloto 1%

Kelompok Sub mikroemulsi 1% Hari ke-1 dan Hari ke-20

<b>Paired Samples Test</b>											
	Paired Differences					t	df	Sig. (2-tailed)			
	Mean	Std. Deviation	Std. Error Mean	95% Confidence Interval of the Difference							
				Lower	Upper						
Pair 1	Krim_Sub_Mikroemulsi_1%_Awal	-	5.763			11.435	2	0.008			
	Krim_Sub_Mikroemulsi_1%_Akhir	33	0.87295	0.50400	3.59482						
r											

Kesimpulan:

1. Karena **Sig. (2-tailed) = 0.008 < 0.05**, maka  $H_0$  ditolak.
2. Artinya, **terdapat perbedaan yang signifikan antara data awal dan data akhir pada kelompok Sub mikroemulsi 1%**

*One-Way ANOVA*

a. Uji Normalitas

Tujuan : untuk distribusi normal data penurunan luas luka bakar

Hipotesis :

$H_0$  = Data penurunan luas luka bakar terdistribusi normal

$H_a$  = Data penurunan luas luka bakar tidak terdistribusi normal

Pengambilan Keputusan :

Jika nilai signifikan  $> 0,05$   $H_0$  diterima

Jika nilai signifikan  $< 0,05$   $H_0$  ditolak

Tests of Normality						
kelompok_Perlakuan	Kolmogorov-Smirnov <sup>a</sup>			Shapiro-Wilk		
	Statist ic	df	Sig.	Statist ic	df	Sig.
TBSA	Positif	.191	3	.	.997	3
	negatif	.217	3	.	.988	3
	Krim Sambiloto	.324	3	.	.878	3
	1%					.317
Sub mikroemulsi	.276	3	.	.942	3	.537
	1%					

a. Lilliefors Significance Correction

Keputusan: data luas luka bakar atau TBSA seluruh kelompok uji terdistribusi normal.

b. Uji Homogenitas

Tujuan : untuk melihat data penurunan luas luka bakar homogen atau tidak

Hipotesis :

$H_0$  = Data penurunan luas luka bakar terdistribusi homogen

$H_a$  = Data penurunan luas luka bakar tidak terdistribusi homogen

Pengambilan Keputusan :

Jika nilai signifikan  $> 0,05$   $H_0$  diterima

Jika nilai signifikan  $< 0,05$   $H_0$  ditolak

Tests of Homogeneity of Variances					
		Levene Statistic	df1	df2	Sig.
TBSA	Based on Mean	1.865	3	8	.214
	Based on Median	.505	3	8	.689
	Based on Median and with adjusted df	.505	3	5.048	.695
	Based on trimmed mean	1.728	3	8	.238

Keputusan: data luas luka bakar atau TBSA seluruh kelompok uji terdistribusi homogen.

c. Uji *Multipe Comparsion tipe LSD (Least Significant Difference)*

Tujuan : untuk menentukan data penurunan luas luka bakar abnormal kelompok mana yang memberikan nilai berbeda secara signifikan dengan kelompok lainnya.

Hipotesis :

$H_0$  : Data penurunan luas luka bakar tidak berbeda secara signifikan

$H_a$  : Data penurunan luas luka bakar berbeda secara signifikan

Pengambilan Keputusan :

Jika nilai signifikan > 0,05 Ho diterima

Jika nilai signifikan < 0,05 Ho ditolak

<b>Multiple Comparisons</b>						
<b>Dependent Variable: TBSA</b>						
	(I) kelompok_Perlakuan	(J) kelompok_Perlakuan	Mean Difference (I-J)	Std. Error	Sig.	95% Confidence Interval
LSD	Positif	Negative	-.35000	.23870	.181	-.9004 .2004
		Krim Sambiloto 1%	-.20667	.23870	.412	-.7571 .3438
		Sub mikroemulsi 1%	-.28667	.23870	.264	-.8371 .2638
	Negative	Positif	.35000	.23870	.181	-.2004 .9004
		Krim Sambiloto 1%	.14333	.23870	.565	-.4071 .6938
		Sub mikroemulsi 1%	.06333	.23870	.797	-.4871 .6138
	Krim Sambiloto 1%	Positif	.20667	.23870	.412	-.3438 .7571
		negatif	-.14333	.23870	.565	-.6938 .4071
		Sub mikroemulsi 1%	-.08000	.23870	.746	-.6304 .4704
	Sub mikroemulsi 1%	Positif	.28667	.23870	.264	-.2638 .8371
		negatif	-.06333	.23870	.797	-.6138 .4871
		Krim Sambiloto 1%	.08000	.23870	.746	-.4704 .6304

Keputusan: data luas luka bakar atau TBSA seluruh kelompok uji tidak berbeda signifikan.

### Lampiran 20. Hasil Analisis Statistika Persentase Penyembuhan Luka Bakar Derajat Dua

a. Uji Normalitas

Tujuan : untuk distribusi normal data persentase penurunan luas luka bakar

Hipotesis :

$H_0$  = Data persentase penyembuhan luka bakar terdistribusi normal

$H_a$  = Data persentase penyembuhan luka bakar tidak terdistribusi normal

Pengambilan Keputusan :

Jika nilai signifikan  $> 0,05$   $H_0$  diterima

Jika nilai signifikan  $< 0,05$   $H_0$  ditolak

Tests of Normality							
	kelompok_Perla kuan	Kolmogorov-Smirnov <sup>a</sup>			Shapiro-Wilk		
		Statistic	df	Sig.	Statist ic	df	Sig.
Persentase_TBSA	Positif	.338	3	.	.852	3	.245
	Negative	.233	3	.	.979	3	.724
	Krim Sambiloto 1%	.354	3	.	.820	3	.164
	Sub mikroemulsi 1%	.357	3	.	.815	3	.151

a. Lilliefors Significance Correction

Keputusan: data persentase penyembuhan luka bakar seluruh kelompok uji terdistribusi normal

b. Uji Homogenitas

Tujuan : untuk melihat data penurunan luas luka bakar homogen atau tidak

Hipotesis :

$H_0$  = Data penurunan luas luka bakar terdistribusi homogen

$H_a$  = Data penurunan luas luka bakar tidak terdistribusi homogen

Pengambilan Keputusan :

Jika nilai signifikan  $> 0,05$   $H_0$  diterima

Jika nilai signifikan  $< 0,05$   $H_0$  ditolak

Tests of Homogeneity of Variances					
		Levene			
		Statistic	df1	df2	Sig.
Percentase_TBSA	Based on Mean	1.931	3	8	.203
	Based on Median	.493	3	8	.697
	Based on Median and with adjusted df	.493	3	5.789	.701
	Based on trimmed mean	1.770	3	8	.230

Keputusan: data persentase penyembuhan luka bakar seluruh kelompok uji terdistribusi homogen

c. Uji *Multipe Comparsion tipe LSD* (Least Significant Difference)

Tujuan : untuk menentukan data penurunan luas luka bakar abnormal kelompok mana yang memberikan nilai berbeda secara signifikan dengan kelompok lainnya.

Hipotesis :

Ho : Data penurunan luas luka bakar tidak berbeda secara signifikan

Ha : Data penurunan luas luka bakar berbeda secara signifikan

Pengambilan Keputusan :

Jika nilai signifikan > 0,05 Ho diterima

Jika nilai signifikan < 0,05 Ho ditolak

Multiple Comparisons						
Dependent Variable: Persentase_TBSA						
	(I)		Mean Difference	Std. Error	Sig.	95% Confidence Interval
LSD	Positif	Negative	27.72000*	2.20130	.000	22.6438 32.7962
		Krim Sambiloto 1%	5.55333*	2.20130	.036	.4771 10.6295
		Sub mikroemulsi 1%	-1.67096	2.20130	.470	-6.7472 3.4052
	negatif	Positif	-27.72000*	2.20130	.000	-32.7962 -22.6438
		Krim Sambiloto 1%	-22.16667*	2.20130	.000	-27.2429 -17.0905
		Sub mikroemulsi 1%	-29.39096*	2.20130	.000	-34.4672 -24.3148
Krim Sambiloto 1%		Positif	-5.55333*	2.20130	.036	-10.6295 -.4771
		Negative	22.16667*	2.20130	.000	17.0905 27.2429
		Sub mikroemulsi 1%	-7.22429*	2.20130	.011	-12.3005 -2.1481
Sub mikroemulsi 1%		Positif	1.67096	2.20130	.470	-3.4052 6.7472
		Negative	29.39096*	2.20130	.000	24.3148 34.4672
		Krim Sambiloto 1%	7.22429*	2.20130	.011	2.1481 12.3005

\*. The mean difference is significant at the 0.05 level.

Keputusan: data persentase penyembuhan luka bakar seluruh kelompok uji tidak berbeda signifikan.