

## **BAB V**

### **KESIMPULAN DAN SARAN**

#### **A. Kesimpulan**

Berdasarkan hasil penelitian dapat disimpulkan bahwa :

1. *Co-processed excipient* manitol-Avicel PH 101 mempunyai karakteristik fisik yang baik, meliputi kandungan lembab, viskositas, sifat alir, kompaktibilitas dan titik leleh nya, sehingga dapat digunakan sebagai bahan pengisi-pengikat tablet kempa langsung dengan mutu fisik yang baik.
2. Formula optimum *co-processed excipient* manitol-Avicel PH 101 berdasarkan pendekatan *simplex lattice design* adalah pada perbandingan 38,93 % manitol dan 61,07 % Avicel PH 101.

#### **B. Saran**

Saran dari penelitian ini adalah :

1. *Co-processed excipient* sebagai bahan pengisi-pengikat tablet kempa langsung masih perlu dikembangkan dari material awal yang lain, dengan harga yang lebih murah, tetapi dapat menghasilkan karakteristik fisik *co-processed excipient* yang baik.
2. Optimalisasi pada *co-processing* secara *spray drying* perlu dilakukan agar didapatkan karakteristik fisik *co-processed excipient* yang lebih baik.

**DAFTAR PUSTAKA**

- [Anonim]. 1979. *Farmakope Indonesia*. Edisi III. Jakarta : Departemen Kesehatan Republik Indonesia. hlm 7, 354.
- [Anonim]. 1995. *Farmakope Indonesia*. Edisi IV. Jakarta : Departemen Kesehatan Republik Indonesia. hlm 4 – 6, 31 – 34.
- Ansel HA. 2008. *Pengantar Bentuk Sediaan Farmasi*. Edisi ke-4. Jakarta : Universitas Indonesia Press. hlm 244 – 272.
- Apeji YE, Oyi A, Musa H, Owolosulu AK. 2010. Investigation of the direct compression properties of microcrystallin starch (MCS) as a filler / binder / disintegrant in metronidazole tablet formulation. *International Journal of Pharmaceutical Research and Innovation* 1 : 8 – 14.
- Awasthi *et al.* 2010. Development of directly compressible co-processed excipients for solid dosage form. *Der Pharmacia Lettre* 2 (6) : 151 – 165.
- Banker GS and Anderson NR. 1994. *Tablet*, in Lachman L Lieberman HA Kanig JL (Eds), *The Theory and Practice of Industrial Pharmacy*. Edisi ke-3. Philadelphia : Lea and Febringer. hlm 293 – 345.
- Bolton S. 1997. *Pharmaceutical Statistic, Practical and Clinical Applications*, 3<sup>rd</sup> Ed. New York : Marcel Dekker Inc. hlm 610 – 611.
- Chougule AS, Dikapati A, Trimbake T. 2012. Formulation development techniques of co-processed excipients. *Journal of Advanced Pharmaceutical Sciences* 2 (2) : 231 – 249.
- Gauhar S, Naqfi SBS, Akram M. 2011. Development of co-processed micro granules for direct compression. *International Journal of Pharmacy and Pharmaceutical Sciences* 3 Supl 2 : 64 – 69.
- Gohel MC, Jogani PD. 2005. A review of co-processed directly compressible excipients. *J Pharm Pharmaceut Sci* 8 (1) : 76-93.
- Hauschild K, Picker KM. 2004. Evaluation of a new coprocessed compound based on maize starch for tablet formulation. *AAPS PharmSci* 6 (2) : 1 – 12.
- Khan, KA. 1975. The Concept of Dissolution Efficiency. *J. Pharm. Pharmacol.* 27 : 48-49.

- Kurniawan DW, Sulaiman TNS. 2009. *Teknologi Sediaan Farmasi*. Yogyakarta : Graha Ilmu. hlm 80.
- Lachman L, Lieberman HA, Kanig JL. 1994. *Teori dan Praktek Farmasi Industri*. Edisi III. Suyatmi S, penerjemah. Jakarta : Universitas Indonesia Press. Terjemahan dari : *The Teory and Practice of Industrial Pharmacy*. hlm. 648 – 705.
- Lieberman HA, Lachman L, Schwart JB. 1989. *Pharmaceutical Dosage Forms : Tablet*. Edisi II. Philadelphia : Marcel Dekker. hlm 245 – 246.
- Limwong V, Sutanthavibul N, Kulvanich P. 2004. Spherical composite particles of rice starch and microcrystalin celullosa : A new coprocessed excipients for direct compression. *AAPS PharmSciTech* 5 (2) : 1 – 10.
- Martin A, Swarbrick J, Cammarata A. 2008. *Farmasi Fisik : Dasar-Dasar Kimia Fisik dalam Ilmu Farmasetik*. Edisi III. Yoshita, penerjemah. Jakarta : Universitas Indonesia Press. Terjemahan dari : *Physical Pharmacy*. hlm 845 – 850.
- Marwaha M, Sandhu D, Marwaha RK. 2010. Coprocessing of excipients : A review on excipient development for improved tableting performance. *International Journal of Applied Pharmaceutics* 2 (3) : 41 – 47.
- Nachegari SK, Bansal AK. 2004. Coprocessed excipients for solid dosage forms. *Pharmaceutical Technology* : 52-64.
- Owolosulu AK, Oyi A, Isah AB, Ibrahim MA. 2011. Formulation and evaluation of novel coprocessed excipients of maize starch and acacia gum (StarAc) for direct compression tableting. *International Journal of Pharmaceutical Research and Innovation* 2 : 39 – 45.
- Patel SS, Patel NM. 2009. Development of directly compressible co-processed excipient for dispersible tablets using  $3^2$  full factorial design. *International Journal of Pharmacy and Pharmaceutical Sciences* 1 (1) : 125 – 148.
- Patel RP, Bhavsar M. 2009. Directly compressible materials via co-processing. *International Journal of Pharm Tech Research* 1 (3) : 745-753.
- Patel RP, Patel MP, Suthar AM. 2009. Spray drying technology: an overview. *Indian Journal of Science and Technology* 2 (10) : 44 – 47.
- Patra SR, Kumar GC, Mallick S. 2011. Preparation and physicomechanical characterisation of naproxen tablets by direct compression. *International Journal of Pharma. Research & Development* 3 (2) : 193 – 201.

- Rowe RC, Sheskey PJ, Quinn ME. 2009. *Handbook of Pharmaceutical Excipients*. Ed ke-6. London : Pharmaceutical Press. hlm. 134-135, 206-207, 404-407, 424-428.
- Shahi *et al.* 2008. Formulation and in vitro evaluation of orodispersible tablet of Etoricoxib with emphasis on comparative functionality evaluation of three classes of superdisintegrants. *Rasayan J Chem* 1 (2) : 292 – 300.
- Siregar CJP, Wikarsa S. 2010. *Teknologi Farmasi Sediaan Tablet, Dasar-Dasar Praktis*. Jakarta : EGC. Hlm 33 – 36, 193 – 203, 235 – 265.
- Voigt R. 1995. *Buku Pelajaran Teknologi Farmasi*. Terjemahan: Soendari Noerono. Cet 2. Yogyakarta: Gadjah Mada University Press. hal. 65, 214, 221-222, 563, 566-567.
- Wicaksono *et al.* 2010. Preparasi dan evaluasi eksipien ko-proses pati singkong – kitosan yang dibuat secara spray drying. *Jurnal Farmasi Indonesia* 5 (2) : 78 – 84.
- Yousuf RI, Shoaib MH, Syed AA, Haque N. 2005. Use of Avicel and spray dried lactose in the development of levofloxacin 250 mg tablets by direct compression method. *Pakistan Journal of Pharmacology* 22 (2) : 67 – 74.
- Zhang Y, Law Y, Chakrabarti S. 2003. Physical properties and compact analysis of commonly used direct compression binders. *AAPS PharmSciTech* 4 (4) : 1-11.

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## Lampiran 1. Surat keterangan bantuan bahan baku asetosal



Sukoharjo, 05 Desember 2012

Nomor : 003/KX-RP/XII/12  
 Perihal : Bahan Baku  
 Lamp. : Acetosal 250 g beserta CoA.

Kepada :  
 Yth. Dekan Fakultas Farmasi  
 Universitas Setia Budi  
 Jl. Let. Jend. Sutoyo  
Solo 57127

up. Ibu Prof. Dr. R.A. Oetari SU., MM., Apt.

Dengan hormat,

Sehubungan dengan surat Ibu no. : 560.26/FF.0/A/SPM/XI/2012 tertanggal 26 November 2012 perihal Permohonan permintaan bahan baku Acetosal 250 g untuk penelitian bagi mahasiswa :

No	Nama Mahasiswa	NIM
1	Chaterina Rosalia Muhartoyo	15092662 A
2	Erna Kurniawati	15092681 A

melalui surat ini kami berikan bahan baku beserta CoA-nya sebagai berikut :

1. Acetosal sebanyak 250 g

Demikian, agar diterima dengan baik.

Hormat kami,  
 PT. KONIMEX



Drs. J. Sunarto, Apt.  
 Apoteker Penanggung Jawab

c.c. : file

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## Lampiran 2. Sertifikat analisis asetosal

**Novacyl**  
Peace of Mind

Manufacturing Plant  
NOVACYL établissement Saint-Fons  
RUE PROSPER MONNET  
69190 - SAINT FONTS  
FRANCE

NOVACYL SAS  
29 AVENUE JOANNÈS MASSET  
Le Carré Joannès - CS 10619  
69009 - LYON  
FRANCE

PT KONIMEX  
DESA SANGGRAHAN, KEC. GROGOL KAB SUKOHARJO  
57192 SUKOHARJO / INDONESIA  
INDONESIA

Attn:  
FAX:

## Certificate of analysis

Print date:  
03.04.2012  
Purchase order item/Date  
PO # 120000483 / 14.03.2012  
Delivery item/Date  
82101105 900001 / 02.05.2012  
Order item/Date  
1337033 000010  
Customer  
70456  
Truck number/Seal number  
GRCUPAGE/

Material: Our / Your reference  
11079 RHODINE 3220 / ACETYLSALICYLIC ACID DRUM 50 KG /  
RHODIA PDS : 230 FP RHO3220.2 - Dec 2010  
NOVACYL PDS : ASA-3220-EP-1110-V1

This product complies with the specifications of the current editions of USP and Ph.EUR.  
Our Rhodine is periodically checked in microbiology by CARSO LSEH of Lyon in accordance with the European Pharmacopeia.  
This batch of Rhodine was manufactured in accordance with EU GMP Volume 4 Part II (ICH Q7) and with current files [R2-CEP 1993-007] and [US DMF N°11373].  
\* Compliance guaranteed, skip testing.

Batch FRH1206851 / Manufacturing date 08.03.2012 / Re-Test date 08.03.2015  
Quantity 2,200 KG

Characteristic	Unit	Value	Lower Limit	Upper Limit
Identity	-	Guaranteed conformity	-	-
<i>Ph. Eur. / USP</i> Ethanollic sol.-colour	-	Liquid less coloured than B9	-	-
<i>Ph. Eur.</i> Ethanollic sol.-turbidity	-	Clear	-	-
<i>Ph. Eur.</i> Loss on drying	%	0.01	-	0.50
<i>Ph. Eur.</i> Assay (on a dry basis)	%	100.0	99.5	101.0
<i>Ph. Eur.</i> Sulphated ash	%	< 0.01	-	0.05
<i>Ph. Eur. / USP</i> Heavy Metals *	ppm	< 10	-	10
<i>Ph. Eur. / USP</i>				

R3220-EUR-2  
1 28.12.2011



Material: Our / Your reference  
 11079 RHODINE 3220 / ACETYLSALICYLIC ACID DRUM 50 KG /  
 RHODIA PDS : 230 FP RHO3220.2 - Dec 2010  
 NOVACYL PDS : ASA-3220-EP-1110-V1  
 Batch FRH1206851 /Manufacturing date 08.03.2012/Re-Test date 08.03.2015 / Quantity 2,200 KG

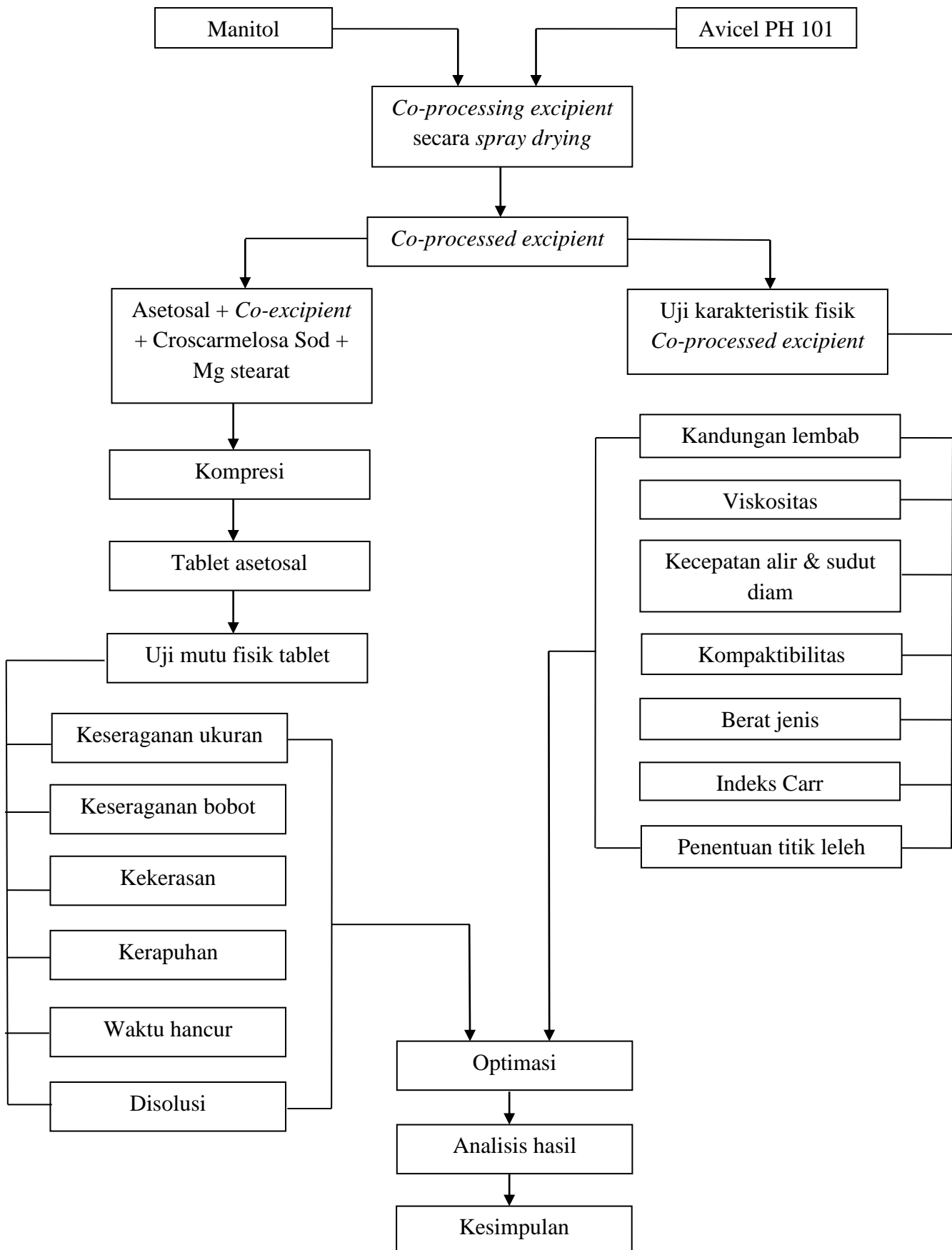
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Anhydride of acetylsalicylic acid	%	< 0.0300	-	0.1500
<i>HPLC</i>				
Acetylsalicylic acid	%	< 0.0300	-	0.1500
<i>HPLC</i>				
Salicylsalicylic acid	%	< 0.0300	-	0.1500
<i>HPLC</i>				
4-hydroxybenzoic acid	%	< 0.0300	-	0.1500
<i>HPLC</i>				
4-hydroxyisophtalic acid	%	< 0.0300	-	0.1500
<i>HPLC</i>				
Free salicylic acid	%	< 0.0300	-	0.1500
<i>HPLC</i>				
Unspecified Impurities>0.05%	-	None	-	-
<i>HPLC</i>				
Sum of related substances	%	< 0.0300	-	0.2500
<i>HPLC</i>				
Retained on 63 mu sieve	%	28.7	0.0	40.0
<i>ALPINE</i>				
Retained on 250 mu sieve	%	0.7	-	1.0

Batch released on :14.03.2012  
 Person responsible for batch release: Bernard Corrias, QA Manager.

R3220-EUR-2  
 1 28.12.2011



**Lampiran 3. Skema jalannya penelitian**

**Lampiran 4. Data kandungan lembab dan viskositas *co-processed excipient***

Kandungan lembab (%)					
Replikasi	Formula I	Formula II	Formula III	Formula IV	Formula V
1.	0,5	2,0	3,5	4,0	11,0
2.	1,0	2,0	3,5	3,5	19,1
3.	0,5	2,0	3,5	3,5	12,0
$\bar{x} \pm SD$	$0,7 \pm 0,29$	$2,0 \pm 0,00$	$3,5 \pm 0,00$	$3,7 \pm 0,29$	$14,0 \pm 4,42$

Viskositas (cP)					
Replikasi	Formula I	Formula II	Formula III	Formula IV	Formula V
1.	300,0	400,0	400,0	500,0	400,0
2.	300,0	400,0	500,0	500,0	400,0
3.	300,0	400,0	400,0	400,0	400,0
$\bar{x} \pm SD$	$300,0 \pm 0,00$	$400,0 \pm 0,00$	$433,3 \pm 57,74$	$466,7 \pm 57,74$	$400,0 \pm 0,00$

### Lampiran 5. Data dan perhitungan kecepatan alir *co-processed excipient*

Berat sampel = 50 gram

$$\text{Kecepatan alir (gram/detik)} = \frac{\text{Berat sampel (gram)}}{\text{waktu alir (detik)}}$$

Waktu alir (detik)					
Replikasi	Formula I	Formula II	Formula III	Formula IV	Formula V
1.	3,44	2,34	2,88	4,37	5,65
2.	3,70	2,47	2,85	4,38	5,90
3.	3,71	2,56	2,69	4,33	5,95
$\bar{x} \pm \text{SD}$	$3,62 \pm 0,15$	$2,46 \pm 0,11$	$2,81 \pm 0,10$	$4,36 \pm 0,03$	$5,83 \pm 0,16$

Kecepatan alir (gram/detik)					
Replikasi	Formula I	Formula II	Formula III	Formula IV	Formula V
1.	14,54	21,37	17,36	11,44	8,85
2.	13,51	20,24	17,54	11,42	8,47
3.	13,48	19,53	17,83	11,55	8,40
$\bar{x} \pm \text{SD}$	$13,84 \pm 0,60$	$20,38 \pm 0,93$	$17,83 \pm 0,66$	$11,47 \pm 0,07$	$8,58 \pm 0,24$

### Lampiran 6. Data dan perhitungan sudut diam *co-processed excipient*

$$\tan \alpha = h/r$$

#### • Formula I

Replikasi	h	r	A
1.	5,876	12,038	26,018
2.	5,862	12,281	25,516
3.	5,562	12,275	24,376
$\bar{x} \pm SD$	$5,767 \pm 0,18$	$12,198 \pm 0,14$	$25,303 \pm 0,84$

#### • Formula II

Replikasi	H	r	A
1.	5,580	11,145	26,595
2.	5,960	11,920	26,565
3.	5,820	11,435	26,974
$\bar{x} \pm SD$	$5,787 \pm 0,18$	$11,500 \pm 0,39$	$26,711 \pm 0,23$

#### • Formula III

Replikasi	H	r	A
1.	6,530	11,200	30,244
2.	6,090	11,465	27,976
3.	6,430	11,860	28,465
$\bar{x} \pm SD$	$6,350 \pm 0,231$	$11,508 \pm 0,332$	$28,895 \pm 1,19$

• **Formula IV**

Replikasi	H	r	A
1.	5,750	12,655	24,445
2.	5,710	12,365	24,787
3.	6,620	12,275	28,338
$\bar{x} \pm SD$	$6,027 \pm 0,51$	$12,432 \pm 0,20$	$25,857 \pm 2,16$

• **Formula V**

Replikasi	H	R	$\alpha$
1.	5,950	11,020	28,366
2.	5,800	10,970	27,866
3.	5,530	10,240	28,371
$\bar{x} \pm SD$	$65,760 \pm 0,21$	$10,743 \pm 0,44$	$28,201 \pm 0,29$

**Lampiran 7. Data kompaktilitas *co-processed excipient***

Kompaktilitas (kg)					
Replikasi	Formula I	Formula II	Formula III	Formula IV	Formula V
1.	6,4	8,0	10,4	10,5	9,3
2.	6,0	8,7	9,4	10,5	9,2
3.	6,2	8,0	9,5	10,4	8,7
$\bar{x} \pm SD$	$6,2 \pm 0,20$	$8,2 \pm 0,40$	$9,8 \pm 0,55$	$10,5 \pm 0,06$	$9,1 \pm 0,32$

### Lampiran 8. Data dan perhitungan keseragaman ukuran tablet

Perhitungan keseragaman ukuran menurut Farmakope Indonesia edisi III :

- **Ketebalan tablet**

- **Formula I**

Replikasi	Tebal tablet	Diameter tablet	Penyimpangan diameter dengan ketebalan tablet (x)
1.	4,00	8,00	2,00
2.	4,00	8,00	2,00
3.	4,00	8,10	2,03
4.	4,05	8,00	1,98
5.	4,10	8,00	1,95
$\bar{x} \pm SD$	$4,03 \pm 0,04$	$8,02 \pm 0,04$	
CV (%)	0,99	0,50	

- **Formula II**

Replikasi	Tebal tablet	Diameter tablet	Penyimpangan diameter dengan ketebalan tablet (x)
1.	4,00	8,00	2,00
2.	4,00	8,10	2,03
3.	3,95	8,10	2,05
4.	4,00	8,00	2,00
5.	4,10	8,00	1,95
$\bar{x} \pm SD$	$4,01 \pm 0,05$	$8,04 \pm 0,05$	
CV (%)	1,25	0,62	

- **Formula III**

Replikasi	Tebal tablet	Diameter tablet	Penyimpangan diameter dengan ketebalan tablet (x)
1.	4,00	8,00	2,00
2.	3,95	8,05	2,04
3.	4,00	8,00	2,00
4.	4,00	8,00	2,00
5.	4,00	8,00	2,00
$\bar{x} \pm SD$	$3,99 \pm 0,02$	$8,01 \pm 0,02$	
CV (%)	0,50	0,25	

- **Formula IV**

Replikasi	Tebal tablet	Diameter tablet	Penyimpangan diameter dengan ketebalan tablet (x)
1.	3,95	8,10	2,05
2.	4,00	8,00	2,00
3.	3,95	8,00	2,03
4.	4,00	8,00	2,00
5.	4,00	8,00	2,00
$\bar{x} \pm SD$	$3,98 \pm 0,03$	$8,02 \pm 0,04$	
CV (%)	0,75	0,50	

- **Formula V**

Replikasi	Tebal tablet	Diameter tablet	Penyimpangan diameter dengan ketebalan tablet (x)
1.	3,95	8,00	2,02
2.	3,95	8,10	2,05
3.	4,00	8,00	2,00
4.	4,00	8,00	2,00
5.	4,00	8,00	2,00
$\bar{x} \pm SD$	$3,98 \pm 0,03$	$8,02 \pm 0,04$	
CV (%)	0,75	0,50	



### Lampiran 9. Data dan perhitungan keseragaman bobot tablet

Perhitungan keseragaman bobot menurut Farmakope Indonesia edisi III

untuk bobot tablet 250 mg :

#### - Formula I

No.	Bobot tablet (mg)	Penyimpangan $ x - \bar{x} $	% penyimpangan $\left  \frac{x - \bar{x}}{\bar{x}} \right  \times 100 \%$
1.	246,8	2,5	1,00
2.	246,8	2,5	1,00
3.	247,8	1,5	0,60
4.	252,8	3,5	1,40
5.	251,8	2,5	1,00
6.	247,5	1,8	0,72
7.	251,5	2,2	0,88
8.	247,0	2,3	0,92
9.	252,5	3,2	1,28
10.	248,4	0,9	0,36
11.	250,1	0,8	0,32
12.	252,8	3,5	1,40
13.	248,0	1,3	0,52
14.	249,8	0,5	0,20
15.	249,9	0,6	0,24
16.	246,6	2,7	1,08
17.	248,1	1,2	0,48
18.	247,2	2,1	0,84
19.	250,3	1,0	0,40
20.	249,6	0,3	0,12
$\bar{x} \pm SD$	$249,3 \pm 2,14$		
CV(%)	0,86		

- **Formula II**

No.	Bobot tablet (mg)	Penyimpangan $[x - \bar{x}]$	% penyimpangan $\left[ \frac{x - \bar{x}}{\bar{x}} \right] \times 100 \%$
1.	247,5	2,3	0,92
2.	246,1	3,7	1,48
3.	253,4	3,6	1,44
4.	252,8	3,0	1,20
5.	251,4	1,6	0,64
6.	249,3	0,5	0,20
7.	250,2	0,4	0,16
8.	251,4	1,6	0,64
9.	248,2	1,6	0,64
10.	252,6	2,8	1,12
11.	250,0	0,2	0,08
12.	250,0	0,2	0,08
13.	249,4	0,4	0,16
14.	252,0	2,2	0,88
15.	248,7	1,1	0,44
16.	247,8	2,0	0,80
17.	250,9	1,1	0,44
18.	248,9	0,9	0,36
19.	247,3	2,5	1,00
20.	248,3	1,5	0,60
$\bar{x} \pm SD$	249,8 $\pm$ 2,02		
CV(%)	0,81		

- **Formula III**

No.	Bobot tablet (mg)	Penyimpangan $[x - \bar{x}]$	% penyimpangan $\left[ \frac{x - \bar{x}}{\bar{x}} \right] \times 100 \%$
1.	248,3	0,4	0,16
2.	250,6	1,9	0,76
3.	249,2	0,5	0,20
4.	250,8	2,1	0,84
5.	246,5	2,2	0,88
6.	248,8	0,1	0,04
7.	245,6	3,1	1,25
8.	247,1	1,6	0,64
9.	247,6	1,1	0,44
10.	249,9	1,2	0,48
11.	246,0	2,7	1,09
12.	247,9	0,8	0,32
13.	251,8	3,1	1,25
14.	247,4	1,3	0,52
15.	253,9	5,2	2,09
16.	248,8	0,1	0,04
17.	247,4	1,3	0,52
18.	251,4	2,7	1,09
19.	247,7	1,0	0,40
20.	247,7	1,0	0,40
$\bar{x} \pm SD$	248,7 $\pm$ 2,12		
CV(%)	0,85		

- **Formula IV**

No.	Bobot tablet (mg)	Penyimpangan $[x - \bar{x}]$	% penyimpangan $\left[ \frac{x - \bar{x}}{\bar{x}} \right] \times 100 \%$
1.	249,9	1,4	0,56
2.	246,2	2,3	0,93
3.	250,9	2,4	0,97
4.	246,2	2,3	0,93
5.	248,3	0,2	0,08
6.	246,3	2,2	0,89
7.	248,6	0,1	0,04
8.	247,6	0,9	0,36
9.	247,2	1,3	0,52
10.	248,3	0,2	0,08
11.	246,9	1,6	0,64
12.	251,0	2,5	1,01
13.	249,8	1,3	0,52
14.	250,4	1,9	0,76
15.	247,3	1,2	0,48
16.	248,1	0,4	0,16
17.	249,0	0,5	0,20
18.	246,7	1,8	0,72
19.	251,6	3,1	1,25
20.	250,2	1,7	0,68
$\bar{x} \pm SD$	248,5 ± 1,74		
CV(%)	0,70		

- **Formula V**

No.	Bobot tablet (mg)	Penyimpangan $[x - \bar{x}]$	% penyimpangan $\left[ \frac{x - \bar{x}}{\bar{x}} \right] \times 100 \%$
1.	242,2	4,1	1,72
2.	242,9	4,8	2,02
3.	234,6	3,5	1,47
4.	234,4	3,7	1,55
5.	235,8	2,3	0,97
6.	238,4	0,3	0,13
7.	239,1	1,0	0,42
8.	236,1	2,0	0,84
9.	239,5	1,4	0,59
10.	240,2	2,1	0,88
11.	236,1	2,0	0,84
12.	238,9	0,8	0,34
13.	243,1	5,0	2,10
14.	240,0	1,9	0,80
15.	231,4	6,7	2,81
16.	236,4	1,7	0,71
17.	242,5	4,4	1,85
18.	237,9	0,2	0,08
19.	236,5	1,6	0,67
20.	235,4	2,7	1,13
$\bar{x} \pm SD$	238,1 $\pm$ 3,19		
CV(%)	1,34		

**Lampiran 10. Data kekerasan tablet**

No.	Kekerasan tablet (kg)				
	Formula I	Formula II	Formula III	Formula IV	Formula V
1.	7,0	7,8	7,6	9,0	7,5
2.	7,0	7,6	8,0	9,2	7,5
3.	7,0	7,8	8,0	9,3	7,6
$\bar{x} \pm SD$	$7,0 \pm 0,00$	$7,7 \pm 0,12$	$7,8 \pm 0,23$	$9,2 \pm 0,15$	$7,5 \pm 0,06$

### Lampiran 11. Data dan perhitungan kerapuhan tablet

#### • Formula I

	Replikasi 1	Replikasi 2	Replikasi 3
Berat sebelum uji (gram)	4,9392	4,9651	4,9491
Berat setelah uji (gram)	4,8956	4,9277	4,9085
% kerapuhan	0,88	0,75	0,82
$\bar{x} \pm SD$			0,82 $\pm$ 0,07

#### • Formula II

	Replikasi 1	Replikasi 2	Replikasi 3
Berat sebelum uji (gram)	4,8696	4,8756	4,9852
Berat setelah uji (gram)	4,8370	4,8439	4,9508
% kerapuhan	0,67	0,65	0,69
$\bar{x} \pm SD$			0,67 $\pm$ 0,02

#### • Formula III

	Replikasi 1	Replikasi 2	Replikasi 3
Berat sebelum uji (gram)	4,9030	4,8897	4,9525
Berat setelah uji (gram)	4,8766	4,8628	4,9217
% kerapuhan	0,54	0,55	0,62
$\bar{x} \pm SD$			0,57 $\pm$ 0,04

• **Formula IV**

	Replikasi 1	Replikasi 2	Replikasi 3
Berat sebelum uji (gram)	5,0488	4,9776	4,9521
Berat setelah uji (gram)	5,0176	4,9438	4,9224
% kerapuhan	0,62	0,68	0,60
$\bar{x} \pm SD$	0,63 $\pm$ 0,04		

• **Formula V**

	Replikasi 1	Replikasi 2	Replikasi 3
Berat sebelum uji (gram)	4,7166	4,7172	4,7221
Berat setelah uji (gram)	4,6876	4,6889	4,6938
% kerapuhan	0,61	0,60	0,60
$\bar{x} \pm SD$	0,60 $\pm$ 0,01		

Contoh perhitungan % kerapuhan adalah sebagai berikut :

$$\begin{aligned} \% \text{ kerapuhan} &= \frac{\text{berat tablet sebelum uji} - \text{bobot tablet setelah uji}}{\text{berat tablet sebelum uji}} \times 100 \% \\ &= \frac{4,9392 - 4,8956}{4,9392} \times 100 \% \\ &= 0,88 \% \end{aligned}$$



**Lampiran 12. Data waktu hancur tablet**

Replikasi	Waktu hancur (menit)				
	F I	F II	F III	F IV	F V
1.	1,32	1,13	0,8	0,9	0,58
2.	1,33	1,17	0,9	1,37	0,67
3.	1,52	1,22	1,12	1,60	1,28
4.	1,62	1,32	1,15	1,87	1,93
5.	1,75	1,40	1,18	2,0	1,95
$\bar{x} \pm SD$	$1,51 \pm 0,19$	$1,25 \pm 0,11$	$1,03 \pm 0,17$	$1,55 \pm 0,44$	$1,28 \pm 0,66$

### Lampiran 13. Data dan perhitungan disolusi tablet asetosal

- Penentuan *operating time*

Waktu (menit)	Absorbansi
1	0,372
2	0,372
3	0,372
4	0,372
5	0,372
6	0,372
7	0,372
8	0,372
9	0,372
10	0,372

- Penentuan panjang gelombang maksimum

Panjang gelombang (nm)	Absorbansi
<b>260</b>	<b>0,372</b>
265	0,362
270	0,355
275	0,312
280	0,237

Panjang gelombang maksimum asetosal adalah 260 nm

Keterangan : larutan yang digunakan adalah larutan asetosal dengan konsentrasi 100 ppm.

- Pembuatan larutan induk asetosal

$$\begin{array}{rcl}
 \text{Kertas timbang + zat} & = & 0,3167 \text{ g} \\
 \text{Kertas + sisa} & = & 0,2667 \text{ g} \\
 \hline
 \text{Bobot zat} & = & 0,0500 \text{ g}
 \end{array}$$

$$\text{Kadar asetosal} = 0,050 \text{ g} / 50 \text{ ml} = 50 \text{ mg} / 50 \text{ ml} = 1000 \text{ mg}/1000 \text{ ml} = 1000 \text{ ppm}$$

- Pengenceran

$$\begin{array}{rcl}
 V_1 \cdot N_1 & = & V_2 \cdot N_2 \\
 2 \cdot 1000 & = & 50 \cdot N_2 \\
 N_2 & = & 80 \text{ ppm}
 \end{array}
 \qquad
 \begin{array}{rcl}
 V_1 \cdot N_1 & = & V_2 \cdot N_2 \\
 2,5 \cdot 1000 & = & 50 \cdot N_2 \\
 N_2 & = & 100 \text{ ppm}
 \end{array}$$

$$\begin{array}{rcl}
 V_1 \cdot N_1 & = & V_2 \cdot N_2 \\
 3 \cdot 1000 & = & 50 \cdot N_2 \\
 N_2 & = & 120 \text{ ppm}
 \end{array}
 \qquad
 \begin{array}{rcl}
 V_1 \cdot N_1 & = & V_2 \cdot N_2 \\
 3,5 \cdot 1000 & = & 50 \cdot N_2 \\
 N_2 & = & 140 \text{ ppm}
 \end{array}$$

$$\begin{array}{rcl}
 V_1 \cdot N_1 & = & V_2 \cdot N_2 \\
 4 \cdot 1000 & = & 50 \cdot N_2 \\
 N_2 & = & 160 \text{ ppm}
 \end{array}$$

- Data kurva baku asetosal

<b>Konsentrasi (ppm)</b>	<b>Absorbansi</b>
80	0,318
100	0,385
120	0,429
140	0,513
160	0,571

Regresi linier :

$$a = 0,0628$$

$$b = 3,17 \times 10^{-3}$$

$$r = 0,9966$$

**Disolusi**

## Formula I Replikasi 1

Waktu (menit)	Absorbansi	F p (kali)	Kadar (mg/1000ml)	Kadar (mg/500 ml)	Faktor koreksi	Kadar terkoreksi	Kadar terdisolusi (%)	Dosis
5	0,441	1	119,306	59,653	0	59,653	74,57	80
10	0,579	1	162,839	81,420	1,193	82,613	103,27	80
15	0,604	1	170,726	85,363	1,628	86,991	108,74	80
30	0,585	1	164,732	82,366	1,707	84,073	<b>105,09</b>	80
45	0,585	1	164,732	82,366	1,647	84,013	105,02	80
60	0,578	1	162,524	81,262	1,647	82,909	103,64	80

## Formula I Replikasi 2

Waktu (menit)	Absorbansi	F p (kali)	Kadar (mg/1000ml)	Kadar (mg/500ml)	Faktor koreksi	Kadar terkoreksi	Kadar Terdisolusi (%)	Dosis
5	0,426	1	114,574	57,287	0	57,287	71,61	80
10	0,545	1	152,114	76,057	1,146	77,203	96,50	80
15	0,554	1	154,953	77,476	1,521	78,997	98,75	80
30	0,566	1	158,738	79,369	1,550	80,919	<b>101,15</b>	80
45	0,568	1	159,369	79,685	1,587	81,272	101,59	80
60	0,574	1	161,262	80,631	1,594	82,225	102,78	80

## Formula I Replikasi 3

Waktu (menit)	Absorbansi	F p (kali)	Kadar (mg/1000ml)	Kadar (mg/500ml)	Faktor koreksi	Kadar terkoreksi	Kadar Terdisolusi (%)	Dosis
5	0,479	1	131,293	65,647	0	65,647	82,06	80
10	0,569	1	159,685	79,842	1,313	81,155	101,44	80
15	0,577	1	162,208	81,104	1,597	82,701	103,38	80
30	0,577	1	162,208	81,104	1,622	82,726	<b>103,41</b>	80
45	0,573	1	160,946	80,473	1,622	82,095	102,62	80
60	0,567	1	159,054	79,527	1,609	81,136	101,42	80

## Formula II Replikasi 1

Waktu (menit)	Absorbansi	F p (kali)	Kadar (mg/1000ml)	Kadar (mg/500ml)	Faktor koreksi	Kadar terkoreksi	Kadar terdisolusi (%)	Dosis
5	0,371	1	97,224	48,612	0	48,612	60,76	80
10	0,513	1	142,019	71,009	0,972	71,982	89,98	80
15	0,551	1	154,006	77,003	1,420	78,423	98,03	80
30	0,577	1	162,208	81,104	1,540	82,644	<b>103,31</b>	80
45	0,547	1	152,744	76,372	1,622	77,994	97,49	80
60	0,547	1	152,744	76,372	1,527	77,900	97,37	80

## Formula II Replikasi 2

Waktu (menit)	Absorbansi	F p (kali)	Kadar (mg/1000ml)	Kadar (mg/500ml)	Faktor koreksi	Kadar terkoreksi	Kadar terdisolusi (%)	Dosis
5	0,347	1	89,653	44,826	0	44,826	56,03	80
10	0,514	1	142,334	71,167	0,897	72,064	90,08	80
15	0,536	1	149,274	74,637	1,423	76,061	95,08	80
30	0,585	1	164,732	82,366	1,493	83,859	<b>104,82</b>	80
45	0,576	1	161,893	80,946	1,647	82,594	103,24	80
60	0,567	1	159,054	79,527	1,619	81,146	101,43	80

## Formula II Replikasi 3

Waktu (menit)	Absorbansi	F p (kali)	Kadar (mg/1000ml)	Kadar (mg/500ml)	Faktor koreksi	Kadar terdisolusi	Kadar terdisolusi (%)	Dosis
5	0,323	1	82,082	41,041	0	41,041	51,30	80
10	0,543	1	151,483	75,741	0,821	76,562	95,70	80
15	0,547	1	152,744	76,372	1,515	77,887	97,36	80
30	0,564	1	158,107	79,054	1,527	80,581	<b>100,73</b>	80
45	0,585	1	164,732	82,366	1,581	83,947	104,93	80
60	0,566	1	158,738	79,369	1,647	81,016	101,27	80

## Formula III Replikasi 1

Waktu (menit)	Absorbansi	F p (kali)	Kadar (mg/1000ml)	Kadar (mg/500ml)	Faktor koreksi	Kadar terkoreksi	Kadar terdisolusi (%)	Dosis
5	0,246	1	57,792	28,896	0	28,896	36,12	80
10	0,372	1	97,539	48,770	0,578	49,348	61,68	80
15	0,400	1	106,372	53,186	0,975	54,162	67,70	80
30	0,468	1	127,823	63,912	1,064	64,975	<b>81,22</b>	80
45	0,511	1	141,388	70,694	1,278	71,972	89,97	80
60	0,526	1	146,120	73,060	1,414	74,474	93,09	80

## Formula III Replikasi 2

Waktu (menit)	Absorbansi	F p (kali)	Kadar (mg/1000ml)	Kadar (mg/500ml)	Faktor koreksi	Kadar terkoreksi	Kadar terdisolusi (%)	Dosis
5	0,295	1	73,249	36,625	0	36,625	45,78	80
10	0,441	1	119,306	59,653	0,732	60,385	75,48	80
15	0,477	1	130,662	65,331	1,193	66,524	83,16	80
30	0,531	1	147,697	73,849	1,307	75,155	<b>93,94</b>	80
45	0,554	1	154,953	77,476	1,477	78,953	98,69	80
60	0,551	1	154,006	77,003	1,550	78,553	98,19	80

## Formula III Replikasi 3

Waktu (menit)	Absorbansi	F p (kali)	Kadar (mg/1000ml)	Kadar (mg/500ml)	Faktor koreksi	Kadar terkoreksi	Kadar terdisolusi (%)	Dosis
5	0,290	1	71,672	35,836	0	35,836	44,79	80
10	0,428	1	115,205	57,603	0,717	58,319	72,90	80
15	0,453	1	123,091	61,546	1,152	62,698	78,37	80
30	0,489	1	134,448	67,224	1,231	68,455	<b>85,57</b>	80
45	0,519	1	143,912	71,956	1,344	73,300	91,63	80
60	0,556	1	155,584	77,792	1,439	79,231	99,04	80

## Formula IV Replikasi 1

Waktu (menit)	Absorbansi	F p (kali)	Kadar (mg/1000ml)	Kadar (mg/500ml)	Faktor koreksi	Kadar terkoreksi	Kadar terdisolusi (%)	Dosis
5	0,304	1	76,088	38,044	0	38,044	47,56	80
10	0,464	1	126,562	63,281	0,761	64,042	80,05	80
15	0,517	1	143,281	71,640	1,266	72,906	91,13	80
30	0,557	1	155,899	77,950	1,433	79,382	<b>99,23</b>	80
45	0,572	1	160,631	80,315	1,559	81,874	102,34	80
60	0,594	1	167,571	83,785	1,606	85,392	106,74	80

## Formula IV Replikasi 2

Waktu (menit)	Absorbansi	F p (kali)	Kadar (mg/1000ml)	Kadar (mg/500ml)	Faktor koreksi	Kadar terkoreksi	Kadar terdisolusi (%)	Dosis
5	0,305	1	76,404	38,202	0	38,202	47,75	80
10	0,482	1	132,240	66,120	0,764	66,884	83,60	80
15	0,528	1	146,751	73,375	1,322	74,698	93,37	80
30	0,531	1	147,697	73,849	1,468	75,316	<b>94,15</b>	80
45	0,562	1	157,476	78,738	1,477	80,215	100,27	80
60	0,572	1	160,631	80,315	1,575	81,890	102,36	80

## Formula IV Replikasi 3

Waktu (menit)	Absorbansi	F p (kali)	Kadar (mg/1000ml)	Kadar (mg/500ml)	Faktor koreksi	Kadar terkoreksi	Kadar terdisolusi (%)	Dosis
5	0,321	1	81,451	40,726	0	40,726	50,91	80
10	0,475	1	130,032	65,016	0,815	65,830	82,29	80
15	0,505	1	139,495	69,748	1,300	71,048	88,81	80
30	0,564	1	158,107	79,054	1,395	80,449	<b>100,56</b>	80
45	0,597	1	168,517	84,259	1,581	85,840	107,30	80
60	0,602	1	170,095	85,047	1,685	86,732	108,42	80



## Formula V Replikasi 1

Waktu (menit)	Absorbansi	F p (kali)	Kadar (mg/1000ml)	Kadar (mg/500)	Faktor koreksi	Kadar terkoreksi	Kadar terdisolusi (%)	Dosis
5	0,218	1	48,959	24,479	0	24,479	30,60	80
10	0,309	1	77,666	38,833	0,490	39,322	49,15	80
15	0,314	1	79,243	39,621	0,777	40,398	50,50	80
30	0,41	1	109,527	54,763	0,792	55,556	<b>69,44</b>	80
45	0,458	1	124,669	62,334	1,095	63,430	79,29	80
60	0,490	1	134,763	67,382	1,247	68,628	85,79	80

## Formula V Replikasi 2

Waktu (menit)	Absorbansi	F p (kali)	Kadar (mg/1000ml)	Kadar (mg/500)	Faktor koreksi	Kadar terkoreksi	Kadar terdisolusi (%)	Dosis
5	0,210	1	46,435	23,218	0	23,218	29,02	80
10	0,317	1	80,189	40,095	0,464	40,559	50,70	80
15	0,352	1	91,230	45,615	0,802	46,417	58,02	80
30	0,42	1	112,681	56,341	0,912	57,253	<b>71,57</b>	80
45	0,457	1	124,353	62,177	1,127	63,303	79,13	80
60	0,495	1	136,341	68,170	1,244	69,414	86,77	80

## Formula V Replikasi 3

Waktu (menit)	Absorbansi	F p (kali)	Kadar (mg/1000ml)	Kadar (mg/500)	Faktor koreksi	Kadar terkoreksi	Kadar terdisolusi (%)	Dosis
5	0,2	1	43,281	21,640	0	21,640	27,05	80
10	0,256	1	60,946	30,473	0,433	30,906	38,63	80
15	0,3	1	74,826	37,413	0,609	38,023	47,53	80
30	0,356	1	92,492	46,246	0,748	46,994	<b>58,74</b>	80
45	0,402	1	107,003	53,502	0,925	54,426	68,03	80
60	0,431	1	116,151	58,076	1,070	59,146	73,93	80

## Kadar rata-rata asetosal yang terdisolusi (%)

Waktu (menit)	Kadar terdisolusi (%)				
	F I	F II	F III	F IV	F V
5	76,08 ±5,39	56,03 ±4,73	42,23±5,32	48,74 ±1,88	28,89±1,78
10	100,40±3,50	91,92 ±3,28	70,02±7,33	81,98 ±1,80	46,16±6,57
15	103,62±5,00	96,82 ±1,55	76,41±7,91	91,10 ±2,28	52,02±5,41
<b>30</b>	<b>103,22±1,98</b>	<b>102,95±2,07</b>	<b>86,91±6,47</b>	<b>97,98 ±3,39</b>	<b>66,58±6,87</b>
45	103,08±1,76	101,89±3,90	93,43±4,63	103,30±3,61	75,48±6,45
60	102,61±1,12	100,03±2,30	96,77±3,22	105,84±3,13	82,16±7,14

## Keterangan :

- Formula I = manitol 100 % : Avicel PH 101 0 %  
 Formula II = manitol 75 % : Avicel PH 101 25 %  
 Formula III = manitol 50 % : Avicel PH 101 50 %  
 Formula IV = manitol 25 % : Avicel PH 101 75 %  
 Formula V = manitol 0 % : Avicel PH 101 100 %

## Contoh perhitungan disolusi pada formula I.

- Kadar mg/1000 ml diperoleh dengan memasukkan absorbansi yang diperoleh pada kurva baku  $y = 0,0628 + 3,17 \times 10^{-3} X$ .  
Absorbansi pada menit ke-5 adalah 0,441 sehingga diperoleh  $x = 119,306$  mg/1000 ml.
- Kadar yang didapat yaitu  $x$  (mg/1000 ml) dikalikan faktor pengenceran yaitu 1x.
- Kadar mg/500 ml adalah kadar zat aktif dalam medium disolusi, dimana volume medium disolusi yang digunakan adalah 500 ml. Kadar yang diperoleh adalah  $500 \text{ ml}/1000 \text{ ml} \times 119,306 \text{ mg} = 59,653 \text{ mg}/500 \text{ ml}$ .
- Pengambilan sampel tiap selang waktu sebanyak 10 ml diganti dengan medium disolusi (dapar asetat pH 4,5) yang baru dengan volume yang sama, karena setiap pengambilan terjadi pengurangan konsentrasi dalam medium disolusi. Agar konsentrasi medium dianggap tetap maka konsentrasi medium

yang diambil tersebut dijadikan faktor koreksi menit ke-10 yang diperoleh dari

$$\frac{10}{500} \times 59,653 \text{ mg} = 1,193$$

5. Kadar terkoreksi pada menit ke-10 diperoleh dari  $1,193 + 81,420 = 82,613$
6. Kadar terdisolusi dalam persen diperoleh dari kadar terkoreksi dibagi dengan dosis asetosal tiap tablet (80 mg) dikalikan 100%.

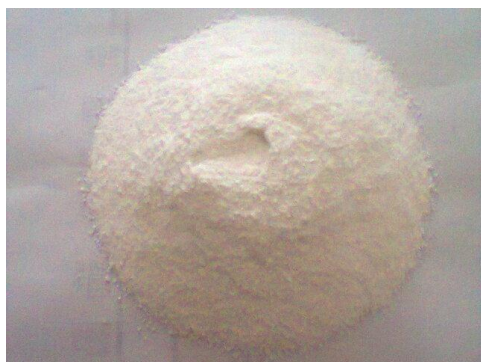
**Lampiran 14. Foto *co-processed excipient* manitol-Avicel PH 101**



*Co-processed excipient* formula I



*Co-processed excipient* formula II



*Co-processed excipient* formula III

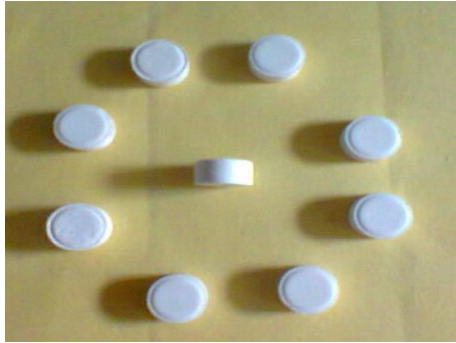


*Co-processed excipient* formula IV

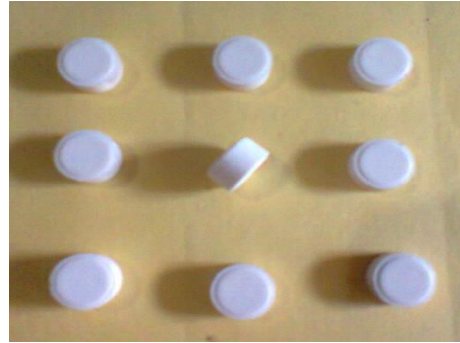


*Co-processed excipient* formula V

**Lampiran 15. Foto tablet kempa langsung asetosal dengan bahan pengisi-pengikat *co-processed excipient* manitol-Avicel PH 101**



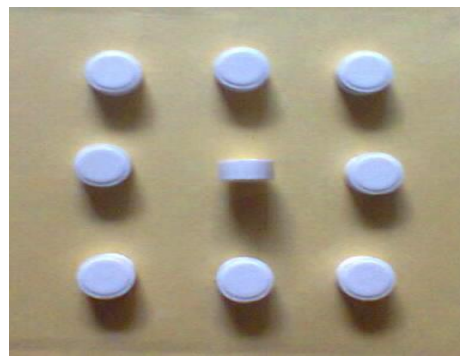
Tablet formula I



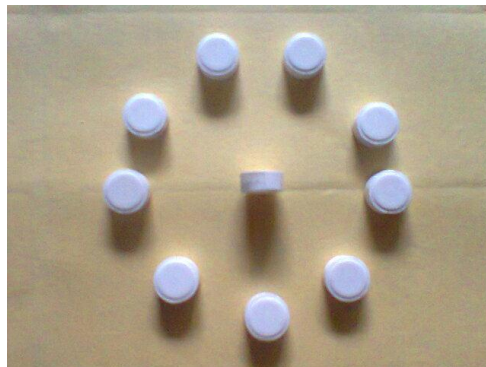
Tablet formula II



Tablet formula III



Tablet formula IV



Tablet formula V

**Lampiran 16. Foto alat yang digunakan dalam penelitian**



*Spray dryer SD basic LabPlant*



*Moisture balance*



*Viskotester*



Alat uji kecepatan alir



Alat uji titik leleh



*Density tester*



Alat pencetak tablet



*Hardness tester*



*Friabilator tester*



*Alat uji waktu hancur*



*Dissolution tester*

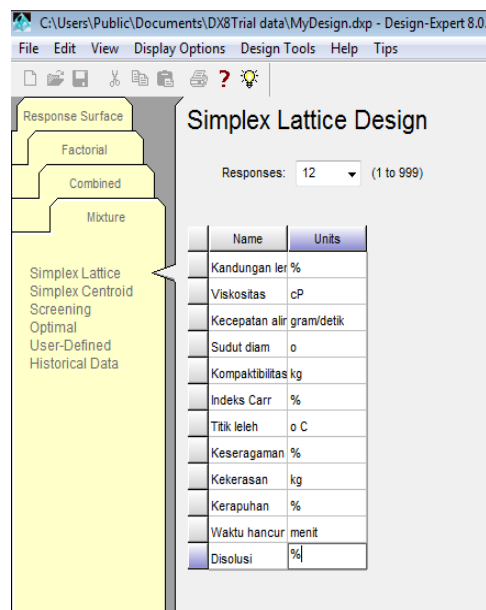
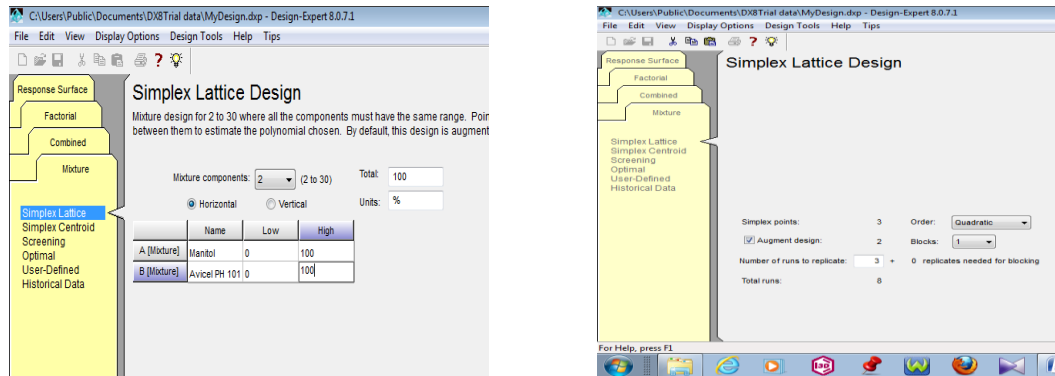


## Spektrofotometer UV-Vis



## Lampiran 17. Optimasi *Simplex Lattice Design* dengan *Software Design*

### Expert versi 8.0.7.1<sup>®</sup>



C:\Users\Public\Documents\DX8Trial data\MyDesign.dxp - Design-Expert 8.0.7.1

File Edit View Display Options Design Tools Help Tips

	Std	Run	Component 1 A:Manitol %	Component 2 B:Avicel PH...	Response 1 Kandungan L... %	Response 2 Viskositas cP	Response 3 Kecepatan alir gram/detik	Response 4 Sudut diam o	Response 5 Kompaktibilitas kg	Response 6 Indeks Carr %	Response 7 Titik leleh o C	Response 8 Kerasagama... %	Response 9 Kerasagama... %
	6	1	100.000	0.000	0.7	300	13.84	25.303	6.2	9.89	122	0.86	
	8	2	50.000	50.000	3.5	433.3	11.47	28.895	9.8	17.38	182	0.85	
	1	3	100.000	0.000	0.7	300	13.84	25.303	6.2	9.89	122	0.86	
	5	4	25.000	75.000	3.5	466.7	11.47	25.857	10.5	14.9	192	0.7	
	3	5	50.000	50.000	3.5	433.3	11.47	28.895	9.8	17.38	182	0.85	
	2	6	0.000	100.000	14	400	8.58	28.201	9.1	10.68	200	1.34	
	4	7	75.000	25.000	2	400	20.38	26.711	8.2	17.01	160	0.81	
	7	8	0.000	100.000	14	400	8.58	28.201	9.1	10.68	200	1.34	

C:\Users\Public\Documents\DX8Trial data\MyDesign.dxp - Design-Expert 8.0.7.1

File Edit View Display Options Design Tools Help Tips

Design (Actual) Transform Fit Summary Model ANOVA Diagnostics Model Graphs

To analyze this response, click on the above icons in succession.

Transformation Equation

None (lambda = 1.0)

$y' = y$

Use with a typical response.

Coding for Analysis: Pseudo

Response ranges from 1.83 to 1.55.  
Ratio of max to min is 1.50485

A ratio greater than 10 usually indicates a transformation is required. For ratios less than 3 the power transforms have little effect.

C:\Users\Public\Documents\DX8Trial data\MyDesign.dxp - Design-Expert 8.0.7.1

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Design (Actual) Transform Fit Summary Model ANOVA Diagnostics Model Graphs

Summary (detailed tables shown below)

Source	p-value	p-value	R-Squared	Predicted R-Squared
R1 Kandungan k	Linear 0.5227	-0.9535	-0.4077	
R2 Viskositas (A)	Quadratic 0.1442	0.1967	-0.1239	
R3 Kecepatan al	Cubic 0.1511	0.4304	-1.2509	
R4 Sudut diam (v)	Quartic <0.0001	1.0000	Suggested	
R5 Kompaktibilitas				
R6 Indeks Carr (				

Source	Sum of Squares	df	Mean Square	F	p-value
Mean vs Total	13.62	1	13.62		
Linear vs Mean	0.021	1	0.021	0.46	0.5227
Quadratic vs L	0.10	1	0.10	2.99	0.1442
Cubic vs Quad	0.077	1	0.077	3.14	0.1511
Quartic vs Cub	0.098	1	0.098	6.3698<007	<0.0001 Suggested
Residual	0.000	3	0.000		
Total	13.92	8	1.74		

C:\Users\Public\Documents\DX8Trial data\MyDesign.dxp - Design-Expert 8.0.7.1

File Edit View Display Options Design Tools Help Tips

Design (Actual) Transform Fit Summary Model ANOVA Diagnostics Model Graphs

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Residual	0.000	3	0.000		
Total	13.92	8	1.74		

"Sequential Model Sum of Squares (Type I)": Select the highest order polynomial where the additional terms are significant and the model is not aliased.

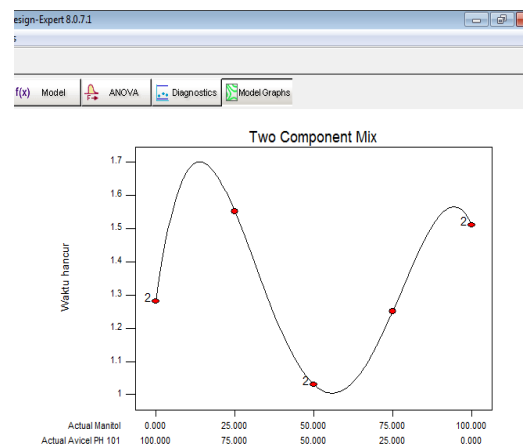
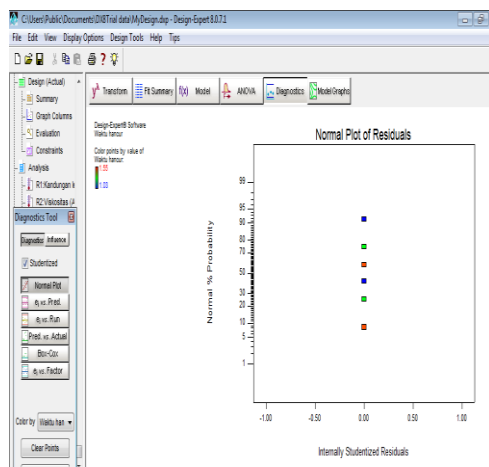
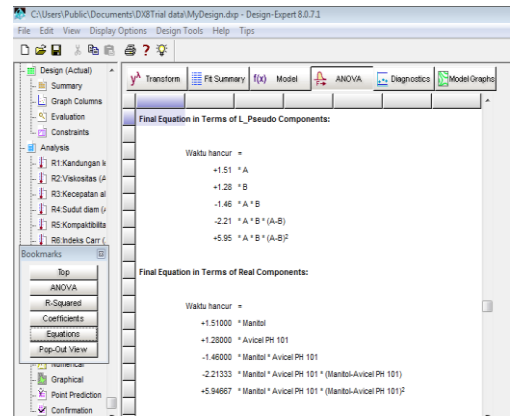
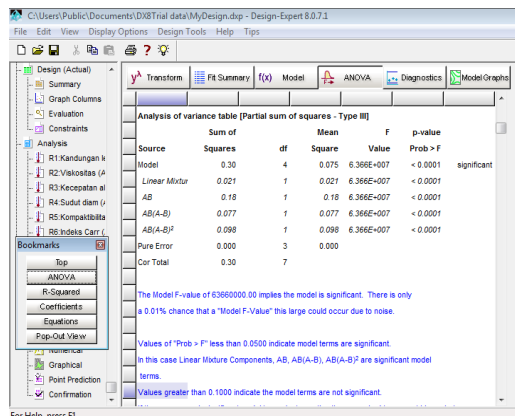
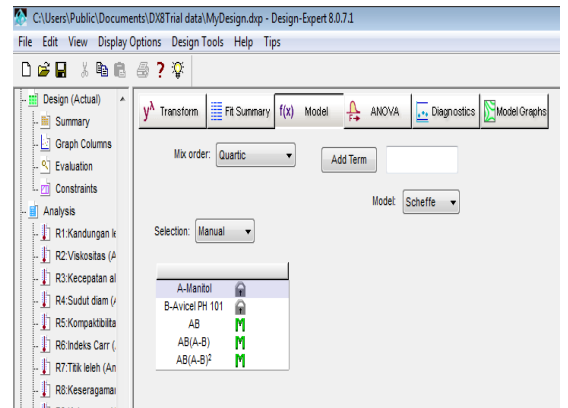
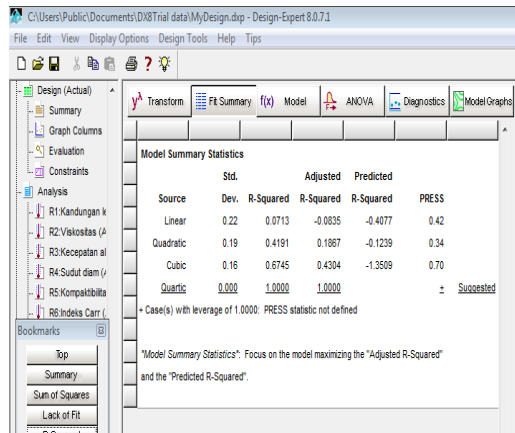
C:\Users\Public\Documents\DX8Trial data\MyDesign.dxp - Design-Expert 8.0.7.1

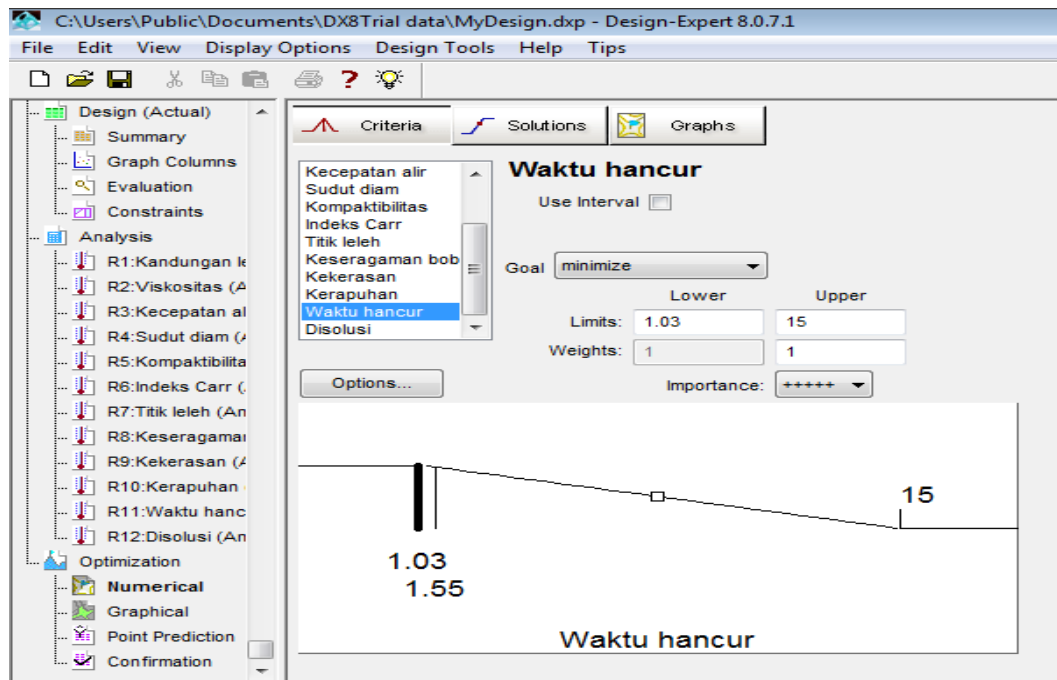
File Edit View Display Options Design Tools Help Tips

Design (Actual) Transform Fit Summary Model ANOVA Diagnostics Model Graphs

Source	Squares	df	Mean Square	F	p-value
Linear	0.28	3	0.093		
Quadratic	0.17	2	0.087		
Cubic	0.098	1	0.098		
Quartic	0.000	0			
Pure Error	0.000	3	0.000		

"Lack of Fit Tests": Want the selected model to have insignificant lack-of-fit.





**Constraints**

Name	Goal	Lower Limit	Upper Limit	Lower Weight	Upper Weight	Importance
A: Mantol	is in range	0	100	1	1	3
B: Avicel PH 10	is in range	0	100	1	1	3
Kandungan ler	is in range	2	4	1	1	3
Viskositas	is in range	300	468.7	1	1	3
Kecepatan alir	maximize	10	20.38	1	1	5
Sudut diam	is in range	25.303	28.895	1	1	3
Kompaktibilitas	maximize	4	10.5	1	1	5
Indeks Carr	is in range	9.89	17.38	1	1	3
Titik leleh	is in range	122	200	1	1	3
Keseragaman	is in range	0.7	1.34	1	1	3
Kekerasan	maximize	7	9.2	1	1	3
Kerapuhan	minimize	0.57	0.82	1	1	3
Waktu hancur	minimize	1.03	15	1	1	5
Disolusi	is target = 90	80	103.22	1	1	5

**Solutions**

Number	Manitol	Avicel PH 10	Kandungan le	Viskositas	Kecepatan ali	Sudut diam	Kompaktibilit	Indeks Carr	Titik leleh	Keseragama	Kekerasan	Kerapuhan	Waktu hancu	Disolu
1	38.927	61.073	3.37048	450.17	15.0872	27.9479	10.2781	16.5028	187.455	0.777952	8.3753	0.583356	1.20831	88.89
2	71.107	28.893	2.40995	405.929	20.4701	27.3458	8.47994	17.38	164.446	0.832116	7.63334	0.644152	1.16935	99.08

2 Solutions found