INDOFARMA

PURCHASING SPECIFICATION

Cimetidine			
No. : RPB 268	**Rev : 08**	**Eff. date : 180518**	**Sign :**

1 **Quality**

1.1 Farmakope Indonesia V, 2014

1.2 USP 37, 2014

2 **Additional Requirements**

2.1 Arrival Date : NMT. 12 months from manufacturing date

3 **Certificate of Analysis from Manufacturer**

To be stated on the certificate

3.1 Manufacturer name and address

3.2 Product name, batch/lot number

3.3 Results of test and assays

3.4 Compliance with Quality

3.5 Quality designation

3.6 Date of manufacturing, expiration (year-month) and date of released (year- month-day)

3.7 Approval/signature of QA/QC responsibility

4 **Attached update Good Manufacture Practice Certificate**

5 **Package**

Cardboard drum @25 kg, double plastic PE, with original seal (inside and outside) from manufacture.
To be stated on goods, label original from manufacture at primary and secondary package each.
Preserve in tight, light-resistant containers.

6 **Labeling**

To be stated on goods **label original from manufacturer**, packing list and invoice :

6.1 Name of manufacturer and address

6.2 Product name, batch/lot number, manufacturing and expiration date

6.3 Quality designation

6.4 Gross/net weight

6.5 Number of packages

7 **Reference Documents**

7.1 Farmakope Indonesia V, 2014

7.2 USP 37, 2014

8 **History**

Purchasing specification is revision of RPB 268 revision 07 with change in the format document and deleting the review point.

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9 **Distribution**

In general, a copy of this specification is distributed to :

9.1 Procurement Dept.

9.2 Quality Control Dept

10 **Approval**

Preparation steps	Position	Code	Signature	Date
Prepared by	Starting and Packaging Material Testing Assistant Manager	**AM**		18 05 18
Checked by	Analytical Method Assistant Manager	**LB**		18 05 18
Approved by	Quality Control Manager	**AM**	an.	18 05 18
	Research & Development Manager	**LB**		18 05 18
	Quality Assurance Manager	**PM**		18 05 18