Clonidine Hydrochloride			
No. : RPB 190	Rev : 08	Eff. date : 18 07 10	Sign :

1 **Quality**

1.1 Farmakope Indonesia V 2014

1.2 USP 39, 2016

2 **Additional Requirements**

2.1 Arrival Date : NMT. 6 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 **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 well - closed containers

5 **Labeling**

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

5.1 Name of manufacturer and address

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

5.3 Quality designation

5.4 Gross/net weight

5.5 Number of packages

6 **Reference Documents**

6.1 Farmakope Indonesia V, 2014

6.2 USP 39, 2016

7 **History**

Purchasing specification is revision of RPB 190 revision 07 with change in the quality of USP 35 to Farmakope Indonesia V and USP 39 and deleting the review point.

8 **Distribution**

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

8.1 Procurement Dept.

8.2 Quality Control Dept.

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9 **Approval**

Preparation steps	Position	Code	Signature	Date
Prepared by	Starting and Packaging Material Testing Assistant Manager	**AM**		180710
Checked by	Analytical Method Assistant Manager	**LB**		180710
Approved by	Quality Control Manager	**AM**	an	180710
	Research & Development Manager	**LB**		180710
	Quality Assurance Manager	**PM**		180710