



PHARMA DEVILS

PRODUCTION DEPARTMENT

STANDARD OPERATING PROCEDURE

Title: Filtration of Bulk Solution

SOP No.:		Department:	Production
		Effective Date:	
Revision No.:	00	Revision Date:	
Supersede Revision No.:	Nil	Page No.:	1 of 2

1.0 OBJECTIVE:

To lay down a procedure for Filtration of Bulk Solution.

2.0 SCOPE:

This SOP is applicable for Filtration of Bulk Solution in Production Department.

3.0 RESPONSIBILITY:

Officer / Executive Production

4.0 ACCOUNTABILITY:

Head Production

5.0 ABBREVIATIONS:

Ltd.	Limited
No.	Number
Pvt.	Private
QC	Quality Control
CIP	Cleaning In Process
μ	Micron
QA	Quality Assurance
SOP	Standard Operating Procedure

6.0 PROCEDURE:

6.1 Clean the Cartridge Filter Housing containing 1.0 μ and 0.22μ Cartridge Filters respectively.

6.2 Assemble the Cartridge Filter and housing and check for proper fixing of "O-Ring" to ensure complete tightness of the Filtration assembly.

6.3 Perform Bubble Point Test of Cartridge Filter to ensure it's integrity before Sterilization.

6.4 Perform Sterilization of assembled Cartridge filter housing containing Cartridge Filter of 0.22 μ.

6.5 Release the nitrogen from filter prior to start up of the filtration operation.

6.6 Start the Batch Filtration after release of the Batch from the Q.C. and as per instructions from Manufacturing Chemist.

6.7 After completion of filtration, carryout CIP then perform the Bubble Point Test of the Cartridge Filter to confirm the Post Filtration Integrity.

6.8 Record the Filtration related details at specified place in respective BMR.



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7.0 ANNEXURES:

ANNEXURE No.	TITLE OF ANNEXURE	FORMAT No.
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ENCLOSURES: SOP Training Record

8.0 DISTRIBUTION:

- Controlled Copy No.01 Quality Assurance
- Controlled Copy No.02 Production
- Master Copy Quality Assurance

9.0 REFERENCES:

Not Applicable.

10.0 REVISION HISTORY:

CHANGE HISTORY LOG

Revision No.	Change Control No.	Details of Changes	Reason for Change	Effective Date	Updated By