



PHARMA DEVILS

PRODUCTION DEPARTMENT

STANDARD OPERATING PROCEDURE

Title: Filter Transfer from Production Area to Filter Cleaning Area

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 Filter Transfer from production area to Filter cleaning area.

2.0 SCOPE:

This SOP is applicable for Filter transfer from production sterile area to filter cleaning area production department.

3.0 RESPONSIBILITY:

Officer / Executive – Production

4.0 ACCOUNTABILITY:

Head – Production

5.0 ABBREVIATIONS:

Ltd. Limited
No. Number
QA Quality Assurance
SOP Standard Operating Procedure
Pvt. Private

6.0 PROCEDURE:

- 6.1 Filter (Area Return riser/Pre-filter of LAF/ Pre/Return filter of Dynamic pass boxes) cleaning shall be done in Filter cleaning area, service floor.
- 6.2 Only authorize production personals shall be entering into the production core area/ascetic area.
- 6.3 Production person shall intimate to engineering personal for Switch off the AHU's then start dismantling the filters.
- 6.4 In case of LAF, switch off then start dismantling the Pre-filters.
- 6.5 Dirty filter shall be kept in sterile polybag with proper identification.
- 6.6 Production person shall intimate to engineering personal for collecting the filter for cleaning.
- 6.7 Filter shall be transfer through respective area dynamic pass boxes.
- 6.8 Filter cleaning shall be done by Engineering team personals as per defined current version of Filter cleaning SOP.
- 6.9 After completion of cleaning, engineering personals shall kept the filter in cleaned double polybag with proper identification and handover to production personals for affixing in designated areas.



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6.10 Mop the filter with 5% Virosil/ Silvicide and transfer the filter to sterile area through dynamic pass box.

7.0 ANNEXURES:
Not Applicable.

ENCLOSURES: SOP Training Record.

8.0 DISTRIBUTION:

- Controlled Copy No. 01 Quality Assurance
- Controlled Copy No. 02 Production
- Controlled Copy No. 03 Engineering
- 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