

PHARMA DEVILS

WAREHOUSE DEPARTMENT

STANDARD OPERATING PROCEDURE				
Department: Warehouse	SOP No.:			
Title: Receipt of Raw Materials in Warehouse	Effective Date:			
Supersedes: Nil	Review Date:			
Issue Date:	Page No.:			

1.0 OBJECTIVE:

To lay down a Procedure for Operation and Cleaning of Reverse Laminar Air Flow Unit installed in PM Warehouse.

2.0 SCOPE:

This SOP is applicable for Operation and Cleaning of Reverse Laminar Air Flow Unit installed in PM warehouse.

3.0 RESPONSIBILITY:

Executive / Officer – QC

Executive / Officer – Warehouse

4.0 ACCOUNTABILITY:

Head – Warehouse

5.0 ABBREVIATIONS:

IPA Isopropyl Alcohol
QC Quality Control
PM Packaging Material

RLAF Reverse Laminar Air Flow SOP Standard Operating Procedure

6.0 PROCEDURE:

6.1 OPERATION OF RLAF UNIT:

- **6.1.1** Ensure that the area and RLAF Unit is cleaned.
- **6.1.2** Executive/Officer of user department shall check the due date of cleaning of pre-filter.
- **6.1.3** Executive/officer of user department shall check due date of preventive maintenance
- **6.1.4** Switch "ON" the mains.
- **6.1.5** Switch "ON" the tube light.
- **6.1.6** Switch "ON" the fan & exhaust.
- **6.1.7** Officer/ Executive QC shall sample the primary packaging materials taking Foil / PVC rolls one by one under the RLAF Unit.
- **6.1.8** After completion of the activity switch "OFF" the fan, exhaust, tube light & mains.

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6.1.9 Remark: Switch "ON" the RLAF minimum 15 minutes before starting the activity.

6.2 CLEANING OF RLAF UNIT:

- **6.2.1** Switch "OFF" the mains.
- **6.2.2** Clean all surface with clean lint free cloth.
- **6.2.3** Use vacuum cleaner if required.
- **6.2.4** Pre-filter shall be cleaned as per respective SOP.
- 6.2.5 RLAF Unit shall be mopped with 70% IPA once in a day before starting the activity and enter Details in "Cleaning and Sanitization Record of RLAF".

7.0 ANNEXURES:

Not Applicable.

ENCLOSURES: SOP Training Record

8.0 DISTRIBUTION:

Controlled Copy No.01 Quality Assurance
 Controlled Copy No.02 Warehouse
 Controlled Copy No.03 Quality Control
 Master Copy Quality Assurance

9.0 REFERENCES

• Schedule M of the Drugs & Cosmetics Act 1940

10.0 REVISION HISTORY:

CHANGE HISTORY LOG

Revision	Change Control	Details of Changes	Reason for	Effective	Updated