



PHARMA DEVILS

PRODUCTION DEPARTMENT

STANDARD OPERATING PROCEDURE

Title: Transfer of Filled Sterilized Bottle from Unloading Area to Packing 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 transfer of filled sterilized bottle from unloading area to packing area.

2.0 SCOPE:

This SOP is applicable for transfer of filled sterilized bottle from unloading area to packing area.

3.0 RESPONSIBILITY:

Officer / Executive Production

4.0 ACCOUNTABILITY:

Head Production

5.0 ABBREVIATIONS:

BMR Batch Manufacturing Record
IPQA In process Quality Assurance
Ltd. Limited
No. Number
SOP Standard Operating Procedure

6.0 PROCEDURE:

- 6.1 Before transfer of sterilized bottles from unloading area to packing hall, check the cleanliness of packing area & Line clearance of packing hall for the same batch.
- 6.2 Ensures that the BMR is complete till the final operation.
- 6.3 Before transfer of sterilized filled bottles, ensure that the sterilization graph is verified by the production & IPQA with respect to set parameter.
- 6.4 Before transfer to packing area verify that the bottles are at normal temperature and outer surface of the bottles are dried and ready for the packing purpose.
- 6.5 Open the hatch sliding door, connecting to unloading area and packing hall, and transfer the sterilized bottles along with trolley to packing hall, one by one, through the sliding door.
- 6.6 After transfer of sterilized bottles to packing hall close the sliding door of the unloading area using lock & key system to avoid the mix up of the batch.
- 6.7 Verify the 'Status Label' & transfer quantity of sterilized bottles as per BMR and on each trolley containing filled and sealed bottles.
- 6.8 Complete BMR shall be handed over to the Packing Area Officer for the further processing.

7.0 ANNEXURES:

Not applicable

ENCLOSURES: SOP Training Record.



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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