



PHARMA DEVILS

PRODUCTION DEPARTMENT

STANDARD OPERATING PROCEDURE

Title: Transfer and Receipt of Dispensed Packaging Material from Store

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 and Receipt of Dispensed Packing Material from Store.

2.0 SCOPE:

This SOP is applicable for Transfer and Receipt of Dispensed Packing Material from Store to Production Area.

3.0 RESPONSIBILITY:

Officer / Executive Production

4.0 ACCOUNTABILITY:

Head Production

5.0 ABBREVIATIONS:

A.R No.	Analytical Report Number
BMR	Batch Manufacturing Record
Ltd.	Limited
No.	Number
QA	Quality Assurance
SOP	Standard Operating Procedure

6.0 PROCEDURE:

6.1 PACKAGING MATERIAL – PRIMARY AND SECONDARY PACKAGING MATERIAL:

- 6.1.1 Check the Packing Material Boxes / Bundles (Glass Vials, Rubber Bungs, Aluminium Seals, Vials, Nozzle, Cap, Labels, Cartons, Shippers, and Leaflet etc.) and verify with Packing Material Requisition Slip.
- 6.1.2 Ensure that each box / bundle has identity label and approved label.
- 6.1.3 Verify all primary and secondary material with Material Requisition Slip & Batch Manufacturing Record (BMR) for its quantity and A.R.No.
- 6.1.4 Verify all primary & Secondary Packaging Material packet / box having “**Packaging Material Identification Slip**”.
- 6.1.5 Verify the AR. No. mentioned on issue slip with material requisition cum issue slip.
- 6.1.6 Transfer all the primary & Secondary Packing Material in respective area of production for further processing.
- 6.1.7 Transfer all the over printable packaging material to the Coding Area.
- 6.1.8 Store the coded packing materials separately or under lock and key condition.



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6.1.9 Bring all the over printed packing materials in packing area at the time of use for further packing operation.

7.0 ANNEXURES:

Not Applicable

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