



# PHARMA DEVILS

PRODUCTION DEPARTMENT

## STANDARD OPERATING PROCEDURE

<b>Department:</b> Production	<b>SOP No.:</b>
<b>Title:</b> Issuance of Additional Raw / Packing Material	<b>Effective Date:</b>
<b>Supersedes:</b> Nil	<b>Review Date:</b>
<b>Issue Date:</b>	<b>Page No.:</b>

### 1.0 OBJECTIVE:

To lay down the procedure for issuance of additional Raw / Packing Material.

### 2.0 SCOPE:

This procedure is applicable to issuance of additional Raw / Packing Material.

### 3.0 RESPONSIBILITY:

Officer/ Executive- Production Department

Head Production & QA Department- shall ensure compliance of the SOP.

### 4.0 DEFINITION(S):

NA

### 5.0 PROCEDURE:

5.1 In case of any spillage of raw material(s), on line rejection of raw material, extra raw materials and for trial purpose (for both raw material & packing material) may be allowed to issue as per the following procedures:

5.1.1 Requisition for the extra issuance of the raw materials shall be made by the production officer / executive giving proper reason for the extra issuance as per reference annexure-1.

5.1.2 The Head- production shall check and sign the requisition.

5.1.3 The requisition shall be forwarded to the Head- QA for final approval.

5.1.4 After approval, issuance the material from store.

### 6.0 ABBREVIATION(S):

QA : Quality Assurance

SOP : Standard Operating Procedure

### 7.0 REFERENCE(S):

NA

### 8.0 ANNEXURE(S):

Annexure – I : Issuance of additional Raw / Packing Material



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### 9.0 DISTRIBUTION:

- 9.1 **Master copy** : Quality Assurance
- 9.2 **Controlled copy( s)** : Production department, Quality Assurance
- 9.3 **Reference copy (s)** : Production department



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### ANNEXURE I

### ADDITIONAL RAW/PACKING MATERIAL REQUISITION SHEET

<b>From:</b>	<b>To:</b>	<b>Date:</b>
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S.No.	Item code	Item Name	UO M	Required Qty.	Issued Qty	A.R. Number	Issued by.	Checked by	Verified by

**Remarks:**

<b>Prepared By (Production)</b>

<b>Checked By (Head Production)</b>

<b>Authorized By (Head QA)</b>