

## PHARMA DEVILS

PRODUCTION DEPARTMENT

| STANDARD OPERATING PROCEDURE                           |                        |  |  |
|--|------------------------|--|--|
| <b>Department:</b> Production (External Preparation)   | SOP No.:               |  |  |
| Title: Remove Defective Equipment from Production Area | <b>Effective Date:</b> |  |  |
| Supersedes: Nil  | Review Date:           |  |  |
| Issue Date:  | Page No.:              |  |  |

### 1.0 **OBJECTIVE**:

To lay down a Procedure for Remove Defective Equipment From Production Area.

#### 2.0 SCOPE:

This SOP is applicable for Remove Defective Equipment.

### 3.0 RESPONSIBILITY:

Production Officer/Executive

### 4.0 ACCOUNTABILITY:

**Head Production** 

### **5.0 ABBREVIATIONS:**

I.D. Identification Number QA Quality Assurance

SOP Standard Operating Procedure

### **6.0 PROCEDURE:**

### 6.1 IDENTIFICATION OF DEFECT EQUIPMENT:

- **6.1.1** If any equipment not working properly inform to Engineering department through breakdown.
- **6.1.2** Engineering person received the breakdown & take action immediately.
- **6.1.3** After maintenance of equipment If equipment not working, Ensure the Engineering head Equipment may be preable or not for further use.
- **6.1.4** If any equipment is not reparable for further use in production area then inform to Production & QA head for defect equipment .
- **6.1.5** Affix the out of order lebel on machine.

## **6.2 REMOVE DEFECT EQUIPMENT:**

- **6.2.1** After Ensure equipment can not in useable then QA & plant head give permission to Engineering head for work permit to remove the equipment.
- **6.2.2** QA & Production will remove the I.D. & all type of status label except out of order label.
- **6.2.3** Remove the Defect Equipment from production area to scrap area.
- **6.2.4** QA Personnel remove the equipment I.D. from equipment list.



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### **7.0** ANNEXURES:

| ANNEXURE No. | TITLE OF ANNEXURE    | FORMAT No. |  |  |
|--------------|----------------------|------------|--|--|
| Annexure-I   | Defect Equipment Log |            |  |  |

**ENCLOSURE:** SOPTraining 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 | Change control No. | Details of | Reason for | Effective | Updated |
|----------|--------------------|------------|------------|-----------|---------|
| No.      |                    | Changes    | Change     | Date      | By      |
|          |                    |            |            |           |         |



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## ANNEXURE – I DEFECT EQUIPMENT LOG

| S.No. | Date | Equipment I.D. | <b>Equipment Name</b> | Production<br>Sign/Date | QA<br>Sign/Date | Plant Head<br>Sign/Date |
|-------|------|----------------|-----------------------|-------------------------|-----------------|-------------------------|
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