



# PHARMA DEVILS

PRODUCTION DEPARTMENT

## STANDARD OPERATING PROCEDURE

**Title:** Procedure for Decartoning of Bottles

<b>SOP No.:</b>		<b>Department:</b>	Production	
		<b>Effective Date:</b>		
<b>Revision No.:</b>	00	<b>Revision Date:</b>		
<b>Supersede Revision No.:</b>	Nil	<b>Page No.:</b>	1 of 2	

### 1.0 OBJECTIVE:

To lay down a Procedure for Decartoning of bottles.

### 2.0 SCOPE:

This SOP is applicable for Decartoning of bottles in Oral Liquid Section.

### 3.0 RESPONSIBILITY:

Officer / Executive Production

### 4.0 ACCOUNTABILITY:

Head Production

### 5.0 ABBREVIATIONS:

BOM Bill of Material  
Ltd. Limited  
Pvt. Private  
SOP Standard Operating Procedure  
SS Stainless Steel

### 6.0 PROCEDURE:

- 6.1 Ensure the Decartoning area shall be cleaned and free from the previous product material (bottles) and affixed as **CLEANED LABEL**.
- 6.2 Receive the empty bottle of the proposed batch inside the decartoning area through the pass box and keep on pellets.
- 6.3 Open the box and remove the outer packing material (Corrugated box) safely to avoid any damage situation.
- 6.4 Remove the Corrugated boxes scraps from the decartoning area and then clean the area to maintain the area free from the dirt/shredded particle of Corrugated Box.
- 6.5 Finally open the inner packing material (polythene bags) to remove the bottles from the bags on the SS table.
- 6.6 Check the bottle specification against the standards mentioned in BOM for its correct use like shape, size and color etc.
- 6.7 Load the empty bottles in the cleaned trays and then Transfer the bottle through hatch/flush window in bottle washing area for washing the bottles.
- 6.8 Record the bottle decartoning activity in the equipment usage log book.



# PHARMA DEVILS

PRODUCTION DEPARTMENT

## STANDARD OPERATING PROCEDURE

**Title:** Procedure for Decartoning of Bottles

<b>SOP No.:</b>		<b>Department:</b>	Production	
		<b>Effective Date:</b>		
<b>Revision No.:</b>	00	<b>Revision Date:</b>		
<b>Supersede Revision No.:</b>	Nil	<b>Page No.:</b>	2 of 2	

**7.0 ANNEXURES:**  
Not Applicable.

**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