# PHARMA DEVILS



PRODUCTION DEPARTMENT

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	S	FANDARD OPERATING PROCE	EDURE				
Departn	ent: Production	SOP No.:					
Title: Cl	eaning and Operation of L	Effective Date:					
Superse	les: Nil	Review Date:					
Issue Da	te:	Page No.:					
Vernao	cular SOP: No						
1.0	OBJECTIVE:						
1.1.	To lay down a procedure for Cleaning and Operation of Lifting and Positioning Device.						
2.0	SCOPE:						
2.1.	The procedure is applicable to the Cleaning and Operation of Lifting and Positioning Device in						
	Production Department.						
3.0	<b>RESPONSIBILITY:</b>						
3.1.	Technical Associate : Operation and cleaning of equipment						
3.2.	Officer/ Executive Production: Supervision						
3.3.	Head Production	: SOP Compliance					
3.4.	IPQA	: SOP Compliance					
4.0	<b>DEFINITION(S):</b>						
4.1.	NA						
5.0	PROCEDURE:						
5.1	CLEANING						
5.1.1	Affix dully filled "UNDER CLEANING" status label on equipment with date and signature of the						
	Production Officer as per SOP ("Status labeling").						
5.1.2	Ensure that power supply is turned "OFF".						
5.1.3	Clean the lifting and positioning device with dry lint free cloth.						
5.1.4	Wipe the surface of lifting and positioning device with 70% v/v IPA solution.						
5.1.5	Replace the "UNDER CLEANING" status label with "CLEANED" status label with date and signature of the Production Officer/QA officer as per SOP ("Status Labeling").						
5.2	<b>OPERATION:</b>						

- 5.2.1 Ensure "CLEANED" label is dully filled and signed is affixed on the equipment.
- 5.2.2 Ensure cleanliness of equipment and area. Remove the "CLEANED" label. Affix "UNDER PROCESS" label dully filled.
- 5.2.3 Switch 'ON' the mains (on main panel).

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ssue Dat	e:	Page No.:						
5.2.4	Bring the IPC or Bin perfectly at the front of lifting and positioning device arms.							
5.2.5	Bring the IPC or Bin to the arm of lifting and positioning de	vice. After insert to the arms fix the						
	locking pin.							
5.2.6	Push the blue UP button to bring the IPC or Bin at the required height, then with holding of holding							
	arms (attached at both sides of lifting and positioning device), bring the IPC or Bin perfectly on the							
	opening of machine hopper, then down the IPC or Bin by pushing the Yellow push button, bring the							
	perfectly in position.							
5.2.7	For charging the material to the compression machine, RMG, sifter, multi-mill, inspection Machine,							
	IPC or Bin (for unloading) and in Packing machine ensure that the equipment charging hopper is							
	properly below the butterfly valve of the lifting and positioning device cone. Lock the L&P with the							
	locking pin.							
5.2.8	Connect the IPC or Bin with hopper by the sleeves.							
5.2.9	Open the butterfly valve of Bin or IPC when lifting and positioning device come over the charging							
	hopper of the equipment perfectly.							
5.2.10	After the operation is over close the butterfly valve, remove sleeves, then slightly UP the lifting and							
	positioning device and then with the help of holding arms move the IPC or Bin.							
5.2.11	By pushing Blue DOWN push button brings the IPC or Bin down to the IPC or Bin stand and perfectly							
	adjust on the stand. Remove the locking pin.							
5.2.12	Remove and destroy "UNDER PROCESS" label and affix "TO BE CLEANED" label dully filled and							
	signed by production officer on the equipment as per SOP ("Status Labeling").							
5.3	Precautions:							
5.3.1	At the time of movement of IPC or Bin, no material or man movement done below the lifting and							
	positioning device.							
5.3.2	Before start operation ensures that lifting and positioning device	arms lock with locking pin.						
6.0	ABBREVIATION (S):							
6.1	IPC : In Process Containers							
6.2	IPA : Iso Propyl Alcohol							
6.3	SOP : Standard Operating Procedure							
6.4	v/v : Volume/ Volume							
6.5	BMR : Batch Manufacturing Record							

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#### 7.0 **RERERENCE** (S):

7.1 SOP No.: Status Labeling

#### 8.0 ANNEXURE(S):

8.1 Nil

#### 9.0 **DISTRIBUTION:**

9.1. Master Copy : Quality Assurance

### 9.2. **Controlled Copy (S) :** Production department (02), Quality Assurance (01)

9.3. **Reference Copy (S) :** Production department (06)

### **10.0 REVISION HISTORY:**

S.	Revision	Change	Reason (S) For	Details Of revision	Revision
No.	No.	Control No.	revision		Date
01	00		New SOP	NA	