

PHARMA DEVILS QUALITY ASSURANCE DEPARTMENT

PERFORMANCE QUALIFICATION PROTOCOL FOR AUTOMATIC HORIZONTAL STICKER LABELING MACHINE

# **PERFORMANCE QUALIFICATION**

## PROTOCOL

### FOR

# AUTOMATIC HORIZONTAL STICKER LABELING MACHINE

EQUIPMENT ID. No.	
LOCATION	Packing Hall
DATE OF QUALIFICATION	
SUPERSEDES PROTOCOL No.	NIL



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#### **1.0 PROTOCOL APPROVAL:**

#### **PRPARED BY:**

DESIGNATION	NAME	SIGNATURE	DATE
OFFICER/EXECUTIVE (QUALITY ASSURANCE)			

#### **REVIEWED BY:**

DESIGNATION	NAME	SIGNATURE	DATE
OPERATING MANAGER (QUALITY ASSURANCE)			
HEAD (ENGINEERING)			
HEAD (PRODUCTION)			

#### **APPROVED BY:**

DESIGNATION	NAME	SIGNATURE	DATE
HEAD (QUALITY ASSURANCE)			



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#### 2.0 **OBJECTIVE:**

- To provide documented evidence that the Equipment is performing consistently, repeatedly and reproducibly within its established operating range and the results of all the test parameters meet the pre-defined acceptance criteria.
- To confirm the suitability of the Standard Operating Procedures for all routine activities associated with the system.

#### **3.0 SCOPE:**

- The Protocol covers all aspects of Performance Qualification for the Automatic Horizontal Sticker Labeling Machine (**Make:** Harikrushna Machinetech Pvt. Ltd.) installed in packing hall.
- This Protocol will define the methods and documentation used to qualify the Automatic Horizontal Sticker Labeling Machine for PQ.



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#### 4.0 **RESPONSIBILITY:**

The Validation Group, comprising of a representative from each of the following departments, shall be responsible for the overall compliance of Performance Qualification Protocol:

DEPARTMENTS	RESPONSIBILITIES		
	Preparation, Review and, Authorization of the Performance		
	Qualification Protocol.		
Quality Assurance	• Co-ordination with Quality Control, Production and Engineering to		
	carryout Performance Qualification Activity.		
	• Monitoring of Performance Qualification Activity.		
Production	Review & Approval of Performance Qualification Protocol.		
Troduction	• To co-ordinate and support Performance Qualification Activity.		
	Review & Approval of Performance Qualification protocol for		
Engineering	correctness, completeness and technical excellence		
Engineering	• Responsible for trouble shooting (if occurred during execution).		
	• Maintenance & preventive maintenance as per schedule.		



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#### 5.0 **EQUIPMENT DETAILS:**

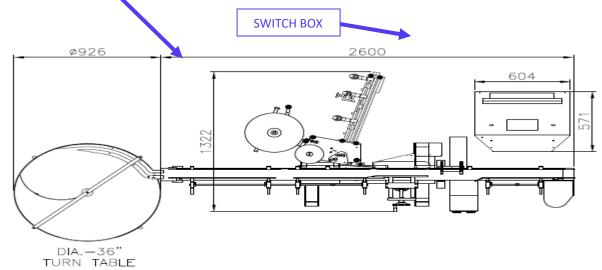
Equipment Name	Automatic Horizontal Sticker Labeling Machine	
Equipment ID.		
Manufacturer's Name	Harikrushna Machinetech Pvt. Ltd.	
Machine No.	<b>.</b> 4486	
Model No.		
Supplier's Name	Harikrushna Machinetech Pvt. Ltd.	
Location of Installation	Packing Hall	

#### 6.0 SYSTEM DESCRIPTION:

This machine is designed to give High Output of Labeling on Bottle. After inspection the Bottle are loaded on In-feed turn table. From in feed turn table, Bottle entered in to infeed conveyor belt. Before the discharge the label device is positioned. Label device having a Printer for printing of batch no/mfg. date/expiry date & then camera system to inspect the printing matter OCR, Pharma code, Barcode etc. & once camera inspect the matter & if found any error then same Bottle with Rejected label is collected into a Box provided for rejected Bottle. The label applicator gets activated as soon as Bottle comes in the position of label; it gets sticks on the Bottles. After this, Bottles move toward the pressing belt meant for proper fixing of label. After pressing of the labels, good Bottles are move forward for further process









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#### 7.0 REASON FOR QUALIFICATION:

• New Machine.

#### 8.0 SITE OF STUDY:

Packing hall.

#### 9.0 FREQUENCY OF QUALIFICATION:

- Once in every 2 Year  $\pm 1$  month.
- After any major breakdown or after major modification.
- After Change of Location.

#### **10.0 PRE – QUALIFICATION REQUIREMENTS:**

- SOP for Operation & Cleaning of Automatic Horizontal Sticker Labeling Machine.
- SOP for Preventive Maintenance of Automatic Horizontal Sticker Labeling Machine.



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#### **11.0 TESTS AND CHECKS:**

#### **11.1 Machine Speed Verification Test:**

**11.1.1 Objective:** To Demonstrate that the machine Perform the smoothly at Different speed claimed by manufacturer and evaluation of the Rejection.

#### 11.1.2 Equipment/ Instruments Used:

Stopwatch.

#### **11.1.3 Method Applied:**

- Switch 'ON' the mains of the machine.
- Start the Machine as per respective SOP of the Equipment Operation.
- Select the Mode of Operation through the PLC i.e. Auto or Manual Mode.
- Then Set the Speed into the Display of Machine.
- Run the machine at Three Different Parameter i.e. Minimum, Optimum and Maximum speed.
- Check the Machine Speed through the Stopwatch after one Minute interval and compare with the Display of Machine.
- Three Trials shall be performed in Both Auto and Manual Mode.

#### 11.1.4 Acceptance Criteria:

- The machine shall give the output of 200-220 labels per minute.
- Machine should run smoothly.



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#### **11.2 Test For Labeling :**

**11.2.1 Objective:** To Demonstrate that the Machine Perform the Smoothly during the Labeling of bottle.

#### **11.2.2 Method Applied:**

- Fix the label roll in to the feeder.
- Run the machine as per Respective SOP.
- Check the Below Mention Parameter at Different Speed.
- Calculate the % age of rejection.
- Three Trials shall be performed.

Parameter	Acceptance Criteria
Labeling Orientation	Should be Uniform
Positioning of Label	Should be proper and should not be tilted
Adhesiveness properties of label	Label should be properly Adhered to bottle
Shrinkage of label	Should be absent
Dent /Rubbing mark on Label	Should be absent
Affixing of labels edges	Label should be intact and properly fixed
Overlapping of Label	Should be absent
Percentage of Rejection	Should be Not More than 2.0%



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#### **11.3 Test for Printing:**

**11.3.1 Objective:** To Demonstrate that the Printing Efficiency of Machine Perform as per manufacturer specification.

#### **11.3.2 Method Applied:**

- Select The Mode of Operation in the Main screen.
- Give the Command to PLC of printing machine as input. Resulting PLC of printing machine gives the Output & Print on Label. Such as

Detail	PLC Input	PLC Out Put
Batch Number:	PC/LB/SLM-001	PC/LB/SLM-001
Mfg Date:	05/18	05/18
Exp Date:	04/20	04/20
MRP:	123.0	123.0

Acceptance Criteria: Coding Imprint Should be Clear & Legible.

#### **11.4 Challenge Test for Camera:**

11.5 Test For Camera: Evaluation of Performance of Camera by Challenge test . Using Defective Printed Label with help of Domino Printer, First of We Set a Coding detail in PLC of Domino Printer and Scan by camera & give Command to PLC.

Batch Number	Mfg. Date	Exp. Date	MRP
PC/LB/SLM-001	6X/Y7	6X/ Y9	123.0



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Test	Set Label Coding Detail		Observation
	Scan by Laptop		Should be Compiles with set Coding Detail
	PC/LB/SLM-001		
<b>Test -01</b>	6X/Y7		
	6X/ Y9		
	123.0		
	Scanning Detail	are	Should be rejected by Camera
	previous		
<b>Test -02</b>	PC/LB/SLM-010		
1050-02	6X/Y7		
	6X/ Y9		
	124.0		
<b>Test -03</b>	Domino Printer not ON		Should be rejected by Camera
Test -04	Spreading of Inc by Domino Printer		Should be rejected by Camera
Test -05	Partial Printing		Should be rejected by Camera

#### 12.0 CHECKLIST OF ALL TESTS AND CHECKS:

S.No.	TESTS AND CHECKS	EXECUTED (Y/N)	REMARKS
1.	Machine Speed Verification		
2.	Test For Labeling		
3.	Test for Printing		
4.	Challenge Test for Camera System.		

#### **13.0 REFERENCES:**

#### The Principle Reference is the following:

- Validation Master Plan
- Schedule-M "Good Manufacturing Practices and Requirements of Premises, Plant and Equipment for Pharmaceutical Products."
- WHO Essential Drugs and Medicines Policy, QA of Pharmaceuticals, Vol-2 Good Manufacturing Practices and Inspection.



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#### 14.0 DOCUMENTS TO BE ATTACHED:

• Any other Relevant Document.

#### **15.0 NON COMPLIANCE:**

• All the Non-compliances of procedure, specifications, and sampling, analysis and documentation activities shall be monitored & recorded.

#### 16.0 DEVIATION FROM PRE-DEFINED SPECIFICATION, IF ANY:

- In case of any deviation observed during PQ, inform to Head QA for necessary action.
- Document the deviation detail in observed deviation section.
- The Head QA will study the impact of deviation. If deviation is acceptable and it does not have an Impact on operation as well as on performance of the machine & prepare final conclusion.

#### 17.0 CHANGE CONTROL, IF ANY:

- If any change control is required during PQ, inform to Head QA for necessary action.
- Document the details observed.
- The Head QA will study the impact of change. If change is acceptable and it does not have an Impact on operation as well as on performance of the machine & prepare final conclusion.

#### **18.0 ABBREVIATIONS:**

- SLM : Automatic Horizontal Sticker Labeling Machine.
- Vol. : Volume
- MRP : Maximum Retail Price
- Mfg. : Manufacturing
- Exp. : Expiry
- GA : General Arrangement
- cGMP : Current Good Manufacturing Practices
- DQ : Design Qualification
- HMI : Human Machine
- IQ : Installation Qualification
- PLC : Programmable Logical Controller
- No. : Number
- WHO : World Health Organization