



**PHARMA DEVILS**  
QUALITY ASSURANCE DEPARTMENT



**DESIGN QUALIFICATION PROTOCOL CUM REPORT FOR VIBRO SIFTER**



**DESIGN QUALIFICATION  
PROTOCOL CUM REPORT  
FOR  
VIBRO SIFTER 30”**

**Pharma Devils**

<b>DATE OF QUALIFICATION</b>	01/01/25
<b>SUPERSEDE PROTOCOL No.</b>	NIL



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### 1.0 PROTOCOL PRE-APPROVAL:

INITIATED BY:



DESIGNATION	NAME	SIGNATURE	DATE
OFFICER/EXECUTIVE (QUALITY ASSURANCE)	QA Executive	<i>Pharmedevils</i>	01/01/25

REVIEWED BY:

DESIGNATION	NAME	SIGNATURE	DATE
HEAD (PRODUCTION)	Head Production	<i>Pharmedevils</i>	01/01/25
HEAD (ENGINEERING)	Head Engineering	<i>Pharmedevils</i>	01/01/25

APPROVED BY:

DESIGNATION	NAME	SIGNATURE	DATE
HEAD (QUALITY ASSURANCE)	Head Quality Assurance	<i>Pharmedevils</i>	01/01/25

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**2.0 OBJECTIVE:**

- To prepare the Design Qualification document for Vibro Sifter on basis of URS and information given by Supplier.
- To ensure that all Critical Aspects of Process/Product Requirement, cGMP and Safety have been considered in designing the equipment and are properly documented.

**3.0 SCOPE:**

- The Scope of this Qualification Document is limited to the Design Qualification of **Vibro Sifter 30” (Make: Elicon Pharma)** for .....
- The Vibro Sifter is a Standalone unit with plug in type electrical connections for operation and is on castor wheel. Hence, may be moved as per requirement to other area of operation which shall not change the performance of equipment.
- The equipment shall be operated under the dust free environment and conditions as per the cGMP requirements.
- The drawings and P & ID's provided by Vendor shall be verified during Design Qualification.







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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 this Protocol cum Report:

DEPARTMENTS	RESPONSIBILITIES
<b>Quality Assurance</b>	<ul style="list-style-type: none"> <li>• Preparation, Review and Approval of the Protocol cum Report.</li> <li>• Assist in the verification of Critical Process Parameters, Drawings as per the Specification.</li> <li>• Co-ordination with Production and Engineering to carryout Design Qualification.</li> <li>• Monitoring of Design Qualification Activity.</li> <li>• Review of Design Qualification Protocol cum Report after Execution.</li> </ul>
<b>Production</b>	<ul style="list-style-type: none"> <li>• Review of Design Qualification Protocol cum Report.</li> <li>• Assist in the verification of Critical Process Parameters, Drawings as per the Specification.</li> <li>• Review of Design Qualification Protocol cum Report after Execution.</li> </ul>
<b>Engineering</b>	<ul style="list-style-type: none"> <li>• Review of Design Qualification Protocol cum Report.</li> <li>• Assist in the Preparation of the Protocol cum Report.</li> <li>• To co-ordinate and support the Activity.</li> <li>• To assist in Verification of Critical Process Parameter, Drawings as per the Specification i.e.               <ul style="list-style-type: none"> <li>➤ GA Drawing</li> <li>➤ Specification of the sub-components/bought out items, their Make, Model, Quantity and backup records/brochures.</li> <li>➤ Details of utilities Required.</li> <li>➤ Identification of components for calibration</li> <li>➤ Material of construction of Product Contact Parts</li> <li>➤ Brief Process Description</li> <li>➤ Safety Features and Alarms</li> </ul> </li> <li>• Review of Design Qualification Protocol cum Report after Execution.</li> </ul>



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### 5.0 PROJECT REQUIREMENTS:

To confirm that safe delivery of the equipment from the supplier site. To ensure that no unauthorized or unrecorded design modification shall take place.

If at any point in time, any change is desired in the mutually agreed design, change control procedure shall be followed and documented.



### 6.0 BRIEF EQUIPMENT DESCRIPTION:

Vibro sifter is an efficient & compact unit self contained & mounted on castor wheels. Vibro sifter have circular unitary vibrating screen used for gradation of material & its proven records over the rotary or longitudinal movement used in the conventional type of sieving machine, both in term of output & uniform grading of materials. Specially designed motor with eccentric weights imparts vibratory motion to the hopper, which have a screen in between them. Material finer than the screen mesh pass through the screen & are collected in the bottom hopper. Coarse material is retained on top of the screen. The amplitude of vibration can be varied from minimum to maximum by adjusting the eccentric weights to suit the process requirement in base minimum time. The machine is generally as per enclosed specs & consists of:

1. **Motor:** It is fitted with top & bottom eccentric weights designed as per required centrifugal force. This whole assembly is covered by SS plate. The motor is flanged mounted & is fixed on the mounting plate by hex. Bolts. The top weights are fixed on the output shaft over the mounting plate.
2. **Spring:** the eight number chrome plated spring are fixed on the base flange at equi- distance. These springs are provided with the ends of the springs. The springs are then screwed on at both the bolts at one end to the base & on the mounting plate at the top. These rugged springs amplify the vibration & restrict them from being transmitted to the floor.
3. **Hopper:** It is a cylindrical, flanged body with an inverted cone at the bottom. This is placed over the mounting plate. The bottom flange is used for clamping to the base plate with a rubber gasket in between the hopper & plate. Hopper is provided with an outlet, tangential to the periphery for discharge of sieved material. The top flange is to provide for holding the charging/ intermediated hopper with a sieve in between them. It is fabricated from stainless steel sheet and works for loading the materials for sifting.
4. **Screen:** based on the product size required a suitable screen is clamped in between the two hopper. Finer mesh sieves can be or with back up cross support to ensure longevity of sieve. This is recommended for sieves finer than 150 meshes.



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5. **Discharge port:** To collect the processed materials.
6. **Conical shape top lid:** It is provided with charging port. Screen is fitted in between the hopper & top lid.

**7.0 EQUIPMENT SPECIFICATION:**

Equipment Specifications are based on User Requirement Specification prepared by ..... The manufacturer of equipment ensures complies with User Requirement Specification.

**8.0 CRITICAL VARIABLES TO BE MET:**

**8.1 PROCESS/PRODUCT PARAMETERS:**

CRITICAL VARIABLES	ACCEPTANCE CRITERIA	REFERENCE
<b>Application:</b> The Vibro Sifter shall be able for sifting of raw material, APIs, Excipients during the manufacturing process.	The Vibro Sifter should be able for sifting of raw material, APIs, Excipients during the manufacturing process.	Process Requirement
<b>Working:</b> Working of Vibro Sifter	Vibro Sifter should capable of sifting the various drugs, raw material, excipients with desired uniformity as per product requirement.	Process Requirement
<b>Electrical Control Panel</b>	The system should have Electrical Control Panel.	Design Requirement

**8.2 UTILITY REQUIREMENTS/LOCATION SUITABILITY:**

CRITICAL VARIABLES	ACCEPTANCE CRITERIA	REFERENCE
<b>Electrical Supply</b>	The electrical system of the equipment shall be housed as per the cGMP and cGEP standards, with adequate safety.	cGMP Requirement
<b>Room Condition</b>	Temperature and RH requirement as per requirement of product	Process Requirement





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### 8.3 TECHNICAL SPECIFICATIONS/KEY DESIGN FEATURES:



S.No.	NAME OF THE COMPONENT	TECHNICAL SPECIFICATION
1.	<b>Model</b>	cGMP
2.	<b>All contact parts</b>	SS316
3.	<b>All non-contact parts</b>	SS304
4.	<b>Capacity</b>	Std.
5.	<b>Dimension</b>	1300 (W) x 800 (D) x 1250 (H) in mm
6.	<b>Charging height</b>	<b>Approx.:</b> 1350 mm,
7.	<b>Discharging height</b>	<b>Approx.:</b> 780 mm, As per your specifications and purchase order.
8.	<b>Electric motor</b>	<b>Make</b> : Vikrant <b>Type</b> : Vibratory <b>H.P</b> : 0.5 HP <b>RPM</b> : 1440 <b>Volt</b> : 415± 10V <b>Amp</b> : 1.2
9.	<b>Screen Diameter</b>	750 mm

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**8.4 MATERIAL OF CONSTRUCTION:**



MACHINE PARTS	ACCEPTANCE CRITERIA	REFERENCE
Top Lid	AISI 316 L	GMP Requirement
Top Deck	AISI 316 L	GMP Requirement
Bottom Deck	AISI 316 L	GMP Requirement
Mesh	AISI 316 L	GMP Requirement
Base	AISI 304	GMP Requirement
'C' - Clamp	AISI 304	GMP Requirement
Gasket	White Food Grade	GMP Requirement
Spring	AISI 304	Design Requirement
Motor Mounting Plate	MS	Design Requirement
Motor	STD	Design Requirement
Castor Wheel	Polyurethane (PU)	GMP Requirement

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**8.5 SAFETY:**

CRITICAL VARIABLES	ACCEPTANCE CRITERIA	REFERENCE
<b>MCB</b>	MCB is provided so that when there is an overload in current or any short circuit then the MCB trips	Safety Requirement
<b>Mechanical Guard</b>	Mechanical guard for all rotating parts.	Safety Requirement
<b>Joints</b>	Welding of joints without any welding burrs	Safety Requirement
<b>Metal Parts</b>	All the metal parts should be Properly grind without any sharp edges.	Safety Requirement
<b>Leveling And Balancing</b>	Equipment should be properly balanced & leveled	Safety Requirement
<b>Electrical Wiring and Earthing</b>	Electrical wiring should be as per approved drawings. Single external Earthing to control machine (panel and motors) and operator should be provided	Safety Requirement
<b>Noise Level</b>	Below 80 db	GMP & Safety Requirement
<b>Emergency Switch</b>	Provided easy access position	GMP & Safety Requirement

**8.6 VENDOR SELECTION:**

CRITICAL VARIABLES	ACCEPTANCE CRITERIA	REFERENCE
<b>Selection of Vendor for supplying the Vibro Sifter</b>	Selection of Vendor is done on the basis of review of vendor. Criteria for review should include vendor background (general/financial), technical knowhow, quality standards, inspection of site, costing, feedback from market (customers already using the equipment)	Process Requirement

**Reference:** (1) User Requirement Specifications (URS).  
(2) Design & Functional Specifications provided by Vendor.

**Verified**  
**(Quality Assurance)** *Maramdevils*  
**Sign/Date:** .....  
*01/01/25*



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**9.0 DOCUMENTS TO BE ATTACHED:**

- Technical details for Equipment Requirement with Engineering Drawings.
- Approved Design and Specifications.
- Any other relevant documents.

**10.0 REVIEW (INCLUSIVE OF FOLLOW UP ACTION, IF ANY):**

All Critical Variables of **Process parameters** (Application/Working/Electrical Control Panel), **Utility requirements or location suitability** (Electrical Supply/Room Condition), **Technical Specification or Key Design features** (Components), **Material of Construction** (Machine parts), **Safety** (MCB/Mechanical Guards/Joints/Metal Parts/Leveling & Balancing/Electrical Wiring & Earthing/Noise Level/Emergency Switch) & **Vendor Selection** (Background/Technical Knowhow/Quality Standards/Inspection of Site/Costing/Feedback from Market) were reviewed and found as per the User Requirement Specification. Hence no any follow up action required.

**11.0 ANY CHANGES MADE AGAINST FORMALLY AGREED PARAMETERS:**

All the Critical Variables of Process Parameters, Utility requirements, Technical Specifications, Material of Construction, Safety & Vendor Selection were observed as per the URS, no any changes were made against the formally agreed parameters.

**12.0 RECOMMENDATION:**

As no any changes were made, Design of the Vibro Sifter was observed as per the provided Design Specification. Hence the Vibro Sifter design is forwarded to vendor for fabrication followed by FAT.

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### 13.0 ABBREVIATIONS:

AISI	:	American Iron Steel Institute
cGMP	:	Current Good Manufacturing Practice
DQ	:	Design Qualification
GA	:	General Arrangement
GB	:	General Block
mm	:	Millimeter
MOC	:	Material of Construction
P & ID	:	Piping and Instrumentation Diagram
QA	:	Quality Assurance
SS	:	Stainless Steel
STD	:	Standard
URS	:	User requirement specification.
VSF	:	Vibro Sifter



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**14.0 REVIEWED BY:**



DESIGNATION	NAME	SIGNATURE	DATE
HEAD (ENGINEERING)	Head Engineering	<i>Pharmadevils</i>	05/01/25

DESIGNATION	NAME	SIGNATURE	DATE
HEAD (PRODUCTION)	Head Production	<i>Pharmadevils</i>	06/01/25

DESIGNATION	NAME	SIGNATURE	DATE
HEAD (QUALITY ASSURANCE)	Head Quality Assurance	<i>Pharmadevils</i>	07/01/25

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## DESIGN QUALIFICATION PROTOCOL CUM REPORT FOR VIBRO SIFTER

### Reference Documents to be attached:

- GA Drawing

