



PHARMA DEVILS

**PERFORMANCE QUALIFICATION REPORT
FOR
JACKETED SUGAR MELTING TANK**

PROTOCOL No.:

**PERFORMANCE QUALIFICATION
REPORT
FOR
JACKETED SUGAR MELTING TANK**

EQUIPMENT ID. No.	
LOCATION	Sugar Melting Room
DATE OF QUALIFICATION	
SUPERSEDES REPORT No.	NIL



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PROTOCOL No.:

1.0 REPORT PRE – APPROVAL:

PREPARED BY:

DESIGNATION	NAME	SIGNATURE	DATE
OFFICER/EXECUTIVE (QUALITY ASSURANCE)			

REVIEWED BY:

OPERATING MANAGER (QUALITY ASSURANCE)			
HEAD (QUALITY CONTROL)			
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 scope of this qualification protocol is limited to qualification of Sugar Melting Tank Installed in Sugar Melting Room Liquid Line.
- This report provides all the relevant information of the performance qualification activity, In-process observations and analytical data of testing of collected samples.



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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 execution of Performance Qualification Report.

DEPARTMENTS	RESPONSIBILITIES
Quality Assurance	<ul style="list-style-type: none">• Preparation, Approval and Compilation of the Performance Qualification Report.• Co-ordination with Quality Control, Production and Engineering to carryout Performance Qualification Activity.• Monitoring of Performance Qualification Activity.• Post Approval of Performance Qualification Report after Execution.
Production	<ul style="list-style-type: none">• Review of Performance Qualification Report.• To co-ordinate and support Performance Qualification Activity.• Post Approval of Performance Qualification Report after Execution.
Quality Control	<ul style="list-style-type: none">• Analytical Support (Microbial Testing/ chemical Analysis).
Engineering	<ul style="list-style-type: none">• Reviewing of qualification report for correctness, completeness and technical excellence• Responsible for trouble shooting (if occurred during execution).• Maintenance & preventive maintenance as per schedule.• Post Approval of Performance Qualification Report after Execution.



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5.0 EQUIPMENT DETAILS:

Equipment Name	SS Jacketed Sugar Melting Tank
ID. Number	
Capacity	1000 Ltr.
Gross Capacity	1200 Ltr.
Manufacturer's Name	Bright Pharma Engineering Pvt. Ltd.
Sr. No.	
Supplier's Name	Bright Pharma Engineering Pvt. Ltd.
Location of Installation	Sugar Melting Area

6.0 PRE – QUALIFICATION REQUIREMENTS:

6.1 Training Record of Validation Team:

All the persons involved in the execution of qualification activity must be trained in all aspects of the qualification activity including the test methodology, acceptance criteria and safety precautions to be followed during working.

6.2 Verification of Documents:

Verify that the DQ/IQ/OQ of Sugar Melting Tank has been executed and approved.

Verify that SOP for Operating, Cleaning and Preventive Maintenance of the Sugar Melting Tank has been prepared.

S.No.	Document Name	Completed (Yes/No)	Checked By Engineering Sign/Date	Verified By (QA) Sign/Date
1.	Executed and approved DQ Protocol Cum Report			
2.	Executed and approved IQ Protocol Cum Report			
3.	Executed and approved OQ Protocol Cum Report			
4.	Approved PQ Protocol			
5.	SOP for Operating, Cleaning of the Sugar Melting Tank			
6.	SOP for Preventive Maintenance of the Sugar Melting Tank			



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7.0 TESTS AND CHECKS :

7.1 Equipment Volumetric Capacity (In Liters) Test:

Name of equipment		Capacity of vessel	
Make		Equipment id no.	

Date of test	Trial no.	Acceptance criteria	Observation
		1000 liter \pm 0.3%	
		200 liter \pm 0.3%	
		1000 liter \pm 0.3%	
		200 liter \pm 0.3%	
		1000 liter \pm 0.3%	
		200 liter \pm 0.3%	

Checked By
Production
Sign/Date:

Verified By
Quality Assurance
Sign/Date:.....

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Reviewed By
Manager QA
Sign/Date:



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PROTOCOL No.:

7.2 Equipment Volumetric Capacity (In Liters) Test by chemical assay method at Minimum Speed of stirrer:

Trial No.:01

Name of equipment		Capacity of vessel	
Make		Equipment Id no.	
Stirrer Speed		Area	Syrup Mfg. room

Date	Volume of tank	Weight of Nacl	Result	Acceptance criteria
			Assay	
	200 Ltr.			0.882% to 0.912%
	400 Ltr.			
	600 Ltr.			
	800 Ltr.			
	1000 Ltr.			

Checked By
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PROTOCOL No.:

Trial No.:02

Name of equipment		Capacity of vessel	
Make		Equipment Id no.	
Stirrer Speed		Area	Syrup Mfg. room

Date	Volume of tank	Weight of Nacl	Result	Acceptance criteria
			Assay	
	200 Ltr.			0.882% to 0.912%
	400 Ltr.			
	600 Ltr.			
	800 Ltr.			
	1000 Ltr.			

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FOR
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PROTOCOL No.:

Trial No.:03

Name of equipment		Capacity of vessel	
Make		Equipment Id no.	
Stirrer Speed		Area	Syrup Mfg. room

Date	Volume of tank	Weight of Nacl	Result	Acceptance criteria
			Assay	
	200 Ltr.			0.882% to 0.912%
	400 Ltr.			
	600 Ltr.			
	800 Ltr.			
	1000 Ltr.			

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PROTOCOL No.:

7.3 Equipment Volumetric Capacity (In Liters) Test by chemical assay method at Maximum Speed of stirrer:

Trial No.:01

Name of equipment		Capacity of vessel	
Make		Equipment Id no.	
Stirrer Speed		Area	Syrup Mfg. room

Date	Volume of tank	Weight of Nacl	Result	Acceptance criteria
			Assay	
	200 Ltr.			0.882% to 0.912%
	400 Ltr.			
	600 Ltr.			
	800 Ltr.			
	1000 Ltr.			

Checked By
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Quality Assurance
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PROTOCOL No.:

Trial No.:02

Name of equipment		Capacity of vessel	
Make		Equipment Id no.	
Stirrer Speed		Area	Syrup Mfg. room

Date	Volume of tank	Weight of Nacl	Result	Acceptance criteria
			Assay	
	200 Ltr.			0.882% to 0.912%
	400 Ltr.			
	600 Ltr.			
	800 Ltr.			
	1000 Ltr.			

Checked By
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PROTOCOL No.:

Trial No.:03

Name of equipment		Capacity of vessel	
Make		Equipment Id no.	
Stirrer Speed		Area	Syrup Mfg. room

Date	Volume of tank	Weight of Nacl	Result	Acceptance criteria
			Assay	
	200 Ltr.			0.882% to 0.912%
	400 Ltr.			
	600 Ltr.			
	800 Ltr.			
	1000 Ltr.			

Checked By
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PROTOCOL No.:

7.4 Test For Verification Of Uniformity Of Mixing at Minimum Speed of stirrer:

Trial No.:01

Date of test		Capacity of vessel	
Name of equipment		Equipment Id no.	
Weight of sodium chloride		Speed	

Sample interval (minute)	Sample location	Critical variables	Acceptance criteria	Observation
After 05	Top	Description	Lump free solution	
	Bottom	Description	Lump free solution	
After 10	Top	Description	Lump free solution	
		pH	5-7	
		Assay	Assay of active content should be within 0.882% W/V – 0.912% W/V	
	Bottom	Description	Lump free solution	
		pH	5-7	
		Assay	Assay of active content should be within 0.882% W/V – 0.912% W/V	
	% RSD of Assay			≤ 2%
After 30	Top	Description	Lump free solution	
		pH	5-7	
		Assay	Assay of active content should be within 0.882% W/V – 0.912% W/V	
	Bottom	Description	Lump free solution	



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PROTOCOL No.:

Sample interval (minute)	Sample location	Critical variables	Acceptance criteria	Observation
		pH	5-7	
		Assay	Assay of active content should be within 0.882% W/V – 0.912% W/V	
	% RSD of Assay		≤ 2%	

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PROTOCOL No.:

Trial No.:02

Date of test		Capacity of vessel	
Name of equipment		Equipment Id no.	
Weight of sodium chloride		Speed	

Sample interval (minute)	Sample location	Critical variables	Acceptance criteria	Observation
After 05	Top	Description	Lump free solution	
	Bottom	Description	Lump free solution	
After 10	Top	Description	Lump free solution	
		pH	5-7	
		Assay	Assay of active content should be within 0.882% W/V – 0.912% W/V	
	Bottom	Description	Lump free solution	
		pH	5-7	
		Assay	Assay of active content should be within 0.882% W/V – 0.912% W/V	
	% RSD of Assay			≤ 2%
After 30	Top	Description	Lump free solution	
		pH	5-7	
		Assay	Assay of active content should be within 0.882% W/V – 0.912% W/V	
	Bottom	Description	Lump free solution	
		pH	5-7	



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Sample interval (minute)	Sample location	Critical variables	Acceptance criteria	Observation
		Assay	Assay of active content should be within 0.882% W/V – 0.912% W/V	
		% RSD of Assay	≤ 2%	

Checked By
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Sign/Date:

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PROTOCOL No.:

Trial No.:03

Date of test		Capacity of vessel	
Name of equipment		Equipment Id no.	
Weight of sodium chloride		Speed	

Sample interval (minute)	Sample location	Critical variables	Acceptance criteria	Observation
After 05	Top	Description	Lump free solution	
	Bottom	Description	Lump free solution	
After 10	Top	Description	Lump free solution	
		pH	5-7	
		Assay	Assay of active content should be within 0.882% W/V – 0.912% W/V	
	Bottom	Description	Lump free solution	
		pH	5-7	
		Assay	Assay of active content should be within 0.882% W/V – 0.912% W/V	
	% RSD of Assay			≤ 2%
After 30	Top	Description	Lump free solution	
		pH	5-7	
		Assay	Assay of active content should be within 0.882% W/V – 0.912% W/V	
	Bottom	Description	Lump free solution	
		pH	5-7	



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PROTOCOL No.:

Sample interval (minute)	Sample location	Critical variables	Acceptance criteria	Observation
		Assay	Assay of active content should be within 0.882% W/V – 0.912% W/V	
		% RSD of Assay		≤ 2%

Checked By
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Quality Assurance
Sign/Date.....

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Reviewed By
Manager QA
Sign/Date:



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JACKETED SUGAR MELTING TANK**

PROTOCOL No.:

7.5 Test For Verification Of Uniformity Of Mixing at Maximum Speed of stirrer:

Trial No.:01

Date of test		Capacity of vessel	
Name of equipment		Equipment Id no.	
Weight of sodium chloride		Speed	

Sample interval (minute)	Sample location	Critical variables	Acceptance criteria	Observation
After 05	Top	Description	Lump free solution	
	Bottom	Description	Lump free solution	
After 10	Top	Description	Lump free solution	
		pH	5-7	
		Assay	Assay of active content should be within 0.882% W/V – 0.912% W/V	
	Bottom	Description	Lump free solution	
		pH	5-7	
		Assay	Assay of active content should be within 0.882% W/V – 0.912% W/V	
	% RSD of Assay			≤ 2%
After 30	Top	Description	Lump free solution	
		pH	5-7	
		Assay	Assay of active content should be within 0.882% W/V – 0.912% W/V	
	Bottom	Description	Lump free solution	



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Sample interval (minute)	Sample location	Critical variables	Acceptance criteria	Observation
		pH	5-7	
		Assay	Assay of active content should be within 0.882% W/V – 0.912% W/V	
	% RSD of Assay		≤ 2%	

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Quality Assurance
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**PERFORMANCE QUALIFICATION REPORT
FOR
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PROTOCOL No.:

Trial No.:02

Date of test		Capacity of vessel	
Name of equipment		Equipment Id no.	
Weight of sodium chloride		Speed	

Sample interval (minute)	Sample location	Critical variables	Acceptance criteria	Observation
After 05	Top	Description	Lump free solution	
	Bottom	Description	Lump free solution	
After 10	Top	Description	Lump free solution	
		pH	5-7	
		Assay	Assay of active content should be within 0.882% W/V – 0.912% W/V	
	Bottom	Description	Lump free solution	
		pH	5-7	
		Assay	Assay of active content should be within 0.882% W/V – 0.912% W/V	
	% RSD of Assay			≤ 2%
After 30	Top	Description	Lump free solution	
		pH	5-7	
		Assay	Assay of active content should be within 0.882% W/V – 0.912% W/V	
	Bottom	Description	Lump free solution	
		pH	5-7	



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Sample interval (minute)	Sample location	Critical variables	Acceptance criteria	Observation
		Assay	Assay of active content should be within 0.882% W/V – 0.912% W/V	
		% RSD of Assay		
			≤ 2%	

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FOR
JACKETED SUGAR MELTING TANK**

PROTOCOL No.:

Trial No.:03

Date of test		Capacity of vessel	
Name of equipment		Equipment Id no.	
Weight of sodium chloride		Speed	

Sample interval (minute)	Sample location	Critical variables	Acceptance criteria	Observation
After 05	Top	Description	Lump free solution	
	Bottom	Description	Lump free solution	
After 10	Top	Description	Lump free solution	
		pH	5-7	
		Assay	Assay of active content should be within 0.882% W/V – 0.912% W/V	
	Bottom	Description	Lump free solution	
		pH	5-7	
		Assay	Assay of active content should be within 0.882% W/V – 0.912% W/V	
		% RSD of Assay		≤ 2%
After 30	Top	Description	Lump free solution	
		pH	5-7	
		Assay	Assay of active content should be within 0.882% W/V – 0.912% W/V	
	Bottom	Description	Lump free solution	
		pH	5-7	



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Sample interval (minute)	Sample location	Critical variables	Acceptance criteria	Observation
		Assay	Assay of active content should be within 0.882% W/V – 0.912% W/V	
	% RSD of Assay		≤ 2%	

Checked By
Production
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Quality Assurance
Sign/Date:.....

Inference:

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Reviewed By
Manager QA
Sign/Date:

8.0 CHECKLIST OF ALL TESTS AND CHECKS:

Tests or checks	Executed [Yes/No]	Remark
Equipment Volumetric Capacity (In Litres) Test		
Verification of Volume of Solution by assay of sodium chloride		
Test For Verification Of Uniformity Of Mixing		

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PROTOCOL No.:

9.0 DOCUMENTS ATTACHED:

- Test Report from QC lab
- Any other Relevant Documents.
- Training Record

10.0 NON COMPLIANCE:

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11.0 DEVIATION FROM PRE-DEFINED SPECIFICATION, IF ANY:

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12.0 CHANGE CONTROL, IF ANY:

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

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14.0 CONCLUSION :

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15.0 RECOMMENDATION :

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16.0 ABBREVIATIONS:

- % : Percentage
- cGMP : Current Good Manufacturing Practices
- CQA : Corporate Quality Assurance
- DQ : Design Qualification
- ID. : Identification
- IQ : Installation Qualification
- LTD. : Limited
- SMT : Sugar Melting Tank
- Nacl : Sodium Chloride
- No. : Number
- OQ : Operational Qualification
- PPQ : Performance Qualification Protocol
- PVT : Private
- RPQ : Report performance qualification
- RSD : Relative standard deviation
- SOP : Standard Operating Procedure



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PROTOCOL No.:

17.0 REPORT POST APPROVAL:

PREPARED BY:

DESIGNATION	NAME	SIGNATURE	DATE
OFFICER/EXECUTIVE (QUALITY ASSURANCE)			

REVIEWED BY:

OPERATING MANAGER (QUALITY ASSURANCE)			
HEAD (QUALITY CONTROL)			
HEAD (ENGINEERING)			
HEAD (PRODUCTION)			

APPROVED BY:

DESIGNATION	NAME	SIGNATURE	DATE
HEAD (QUALITY ASSURANCE)			