

**DERREN HEALTHCARE PVT. LTD.**

Plot No. #33, 35/p & 36/p XCELON Industrial Park, Lane besides Chak-de-India
Weighbridge, Sarkhej-Bavla Highway Village: Vasna-Chacharvadi,
Ahmedabad-382 213, Gujarat (India)

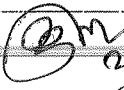

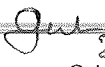
**CERTIFICATE OF ANALYSIS
(FINISHED PRODUCT)**

Name of Product	Milrinone Lactate Injection USP 1 mg/ml		
Batch No.	BM23001	A.R. No.	FP/23/0073
Mfg. Date	11/23	Exp. Date	10/25
Sampled Qty.	34 Nos.	Date of Analysis	10/11/23
Batch Size	60 Ltr.	Date of Release	20/12/23
Mfg. Lic. No.	G/28/1862		

Sr. No.	Test	Result	Specification
1	Description	Clear, Colourless solution free from visible particles and fibers.	Clear, colourless or almost colourless solution free from visible particles and fibers.
2	Extractable volume	10.1 ml	NLT 10.0 ml
3	Identification		
3.1	Identification A by HPLC	The retention time of the milrinone peak of the Sample solution is corresponds to that of the Standard solution, as obtained in the Assay.	The retention time of the milrinone peak of the Sample solution should be corresponds to that of the Standard solution, as obtained in the Assay.
4	pH	3.6	3.2 to 4.0
5	Related Substances		
5.1	Any secondary impurity	Not detected	NMT 0.2%
5.2	Total Impurity	Not detected	NMT 0.5%
6	Bacterial endotoxin	Less than 25.0 EU/mg	NMT 25.0 EU/mg of Milrinone
7	Particulate matter		
7.1	By Visual Inspection	The solution is free from particles that is observed by inspection with the unaided eye.	The solution should be free from particles that can be observed by inspection with the unaided eye.
7.2	By Light Obscuration Method	≥10 microns= 123.0/Container ≥ 25 Microns= 1.67/Container	Sub visible particle by LPC ≥10 microns=NMT 6000/Container ≥ 25 Microns=NMT 600/Container
8	Sterility test	Sterile	Should be sterile
9	Assay		
9.1	Each ml contains: Milrinone Lactate eq. to Milrinone 1.0 mg	99.0%	90.0% to 110.0% of the stated amount of Milrinone

Conclusion: The above sample complies as per USP specification.

In the opinion of the undersigned the sample referred to above is **of standard** / **not-of-standard** quality as defined in the act and the rules made there under for the reasons given above.

Prepared By  20/12/23 QC OFFICER	Checked By  20/12/23 QC MANAGER	Approved By  20/12/23 QA MANAGER
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