



American Remedies Healthcare Pvt. Ltd.

Surajpur, Nahar Road Paonta sahib Dist. Sirmour (H.P)
QUALITY CONTROL DEPARTMENT

CERTIFICATE OF FINISHED PRODUCT ANALYSIS

Product Name:	Ranican-2ml Injection (Ranitidine Injection IP 25mg/ml)		
Batch No.	26AR013	A.R. No.	AFP260031
Batch Size	100200 Vials	Sample Quantity	60 Nos
Mfg. Date	02/2026	Date of Testing	26/02/2026
Exp. Date	01/2028	Date of Release	14/03/2026
Reference Batch No	26AR013	MFG. Lic. No.	HP-MB-LL-00235

Sr. No.	Test	Specification	Observation
1.	Description	A clear, colourless solution	Clear and colorless solution
2.	Identification	A.) Determine by infrared absorption spectrophotometry (2.4.6) compare the spectrum with that obtained with ranitidine hydrochloride IPRS or with the reference spectrum of ranitidine hydrochloride. (B) In the Assay, the principal peak in the chromatogram obtained with the test solution corresponds to the peak in the chromatogram obtained with the reference solution.	Complies
3.	pH	Between 4.5 to 7.0	5.81
4.	Related Substance		



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	A.) The any secondary impurity B.) The two other secondary impurity C.) Two further secondary impurity D.) The principal impurity	NMT (2.0 per cent) NMT (1.0 per cent) NMT (0.5 per cent) NMT (0.2 per cent)	Complies
5.	Extractable Volume	NLT 2.0 ml	2.0 ml
6.	Bacterial Endotoxins	NMT 7.0 Endotoxin units per mg of ranitidine	Less than 7.0 Endotoxin unit per mg of ranitidine
7.	Sterility	Should be Sterile	Sterile
8.	Assay: Each ml contains: Ranitidine hydrochloride (25 mg)	(90.0% to 110.0%) (22.5mg to 27.5mg)	(101.31 %) (25.33 mg)



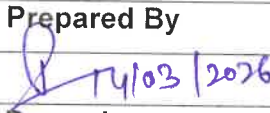
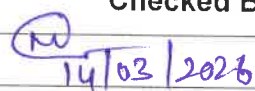
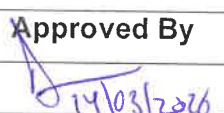
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Remark: The sample is complies as per IP /BP/ USP/IH specification.

	Prepared By	Checked By	Approved By
Sign/Date	 14/03/2026	 14/03/2026	 14/03/2026
Name	Prayash	Manisha Thakur	Bhupinder Navet
Designation	Officer QC	Executive-QC	Head-QC

