



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.	26AR031	A.R. No.	AFP260044
Batch Size	104761 Vials	Sample Quantity	60 Nos
Mfg. Date	03/2026	Date of Testing	31/03/2026
Exp. Date	02/2028	Date of Release	16/04/2026
Reference Batch No	26AR031	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	5.83
4.	Related Substance		



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	A.) The any secondary impurity	NMT (2.0 per cent)	Not Detected
	B.) The two other secondary impurity	NMT (0.5 per cent)	Not Detected
	C.) Two further secondary impurity	NMT (0.2 per cent)	Not Detected
	D.) The principal impurity	NMT (1.0 per cent)	Not Detected
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)	(102.56 %) (25.64 mg)



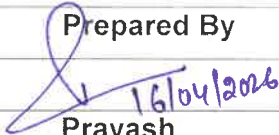
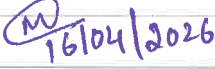
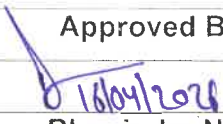
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Remark: The sample is complies as per IP /BP/ USP/IH specification. ✓

	Prepared By	Checked By	Approved By
Sign/Date	 16/04/2026	 16/04/2026	 16/04/2026
Name	Prayash	Manisha Thakur	Bhupinder Navet
Designation	Officer QC	Executive-QC	Head-QC

