

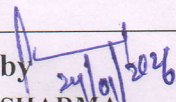
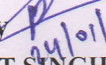
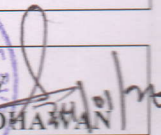
**AMERICAN REMEDIES HEALTHCARE PVT. LTD.**

Vill. Surajpur, Nahan Road, Paonta Sahib, Dist. Sirmour (H.P) 173025

Title : Certificate of Analysis Finished Product

Product Name	FUXORIME – 1.5gm Injection	A.R. No.	ARB/FP/25/022
Generic Name	Cefuroxime Injection IP 1.5gm	Sampled qty.	45 vials
Batch No.	ARB2525A	Sampled by	Anuj Verma
Batch Size	19,930 vials	Sampled on	05/01/2026
Mfg. Date	12/2025	Date of Testing	06/01/2026
Exp. Date	11/2027	Date of Release	24/01/2026

S. No.	Tests	Specifications	Observations
1.	Description	A white or faintly yellow dry powder filled in clear glass vial.	A white dry powder filled in clear glass vial.
2.	Identification (By HPLC)	A. 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. B. It gives the reactions of sodium salts.	Complies Complies
3.	Uniformity of weight	Average weight $\pm 10\%$	-1.13% & +0.60%
4.	Average weight	Informative $\pm 2\%$.	1652.4 mg
5.	Related Substance (By HPLC)		
	Descarbamoyl-cefuroxime	NMT 1.0%	0.76%
	Area of any other secondary peak	NMT 1.0%	0.25%
	Sum of areas of all the secondary peaks	NMT 3.0%	0.39%
6.	pH	6.0 to 8.5	6.58
7.	Water (By K.F.)	NMT 3.5%w/w	1.37%w/w
8.	Particulate Matter (A.) Light Obscuration Particle Count Test (1.) Particles $\geq 10\mu\text{m}$ (2.) Particles $\geq 25\mu\text{m}$ (B.) Visual Particle	NMT 6000/vial NMT 600/vial The reconstituted solution should be essentially free from extraneous, mobile undissolved particles, other than gas bubbles, unintentionally that can be observed on visual inspection.	270/vial 7/vial Complies
9.	Sterility	No microbial growth should be observed.	Complies
10.	Bacterial Endotoxins	NMT 0.1 EU/mg of Cefuroxime.	Less than 0.1 EU/mg of Cefuroxime.

Analysis by 
NAME: NIKHIL SHARMAChecked by 
NAME: RAMANJEET SINGHApproved by 
NAME: MRS. SWATI DHAUMAN

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11.	Assay: (By HPLC) Each glass vial contains:			
Ingredients	Labeled Claim	Found	% of labeled amount	Limits % of labeled amount
Cefuroxime Sodium (Sterile) IP Eq. to Cefuroxime	1500 mg	1524.5809 mg	101.64%	90.0 to 120.0%

Remarks: In the opinion of the undersigned the sample referred to above is/ is ~~not~~ of the standard quality as The Drug & Cosmetic Act 1940 and the rules made there under. Complies/ ~~not~~ complies as per IP/BP/USP/IHS.

Analysis by  NAME: NIKHIL SHARMA	Checked by  NAME: RAMANJEET SINGH	Approved by  NAME: MRS. SWATI DHAWAN
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