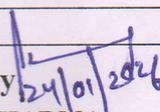
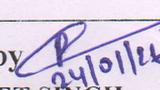
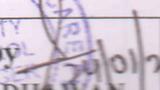
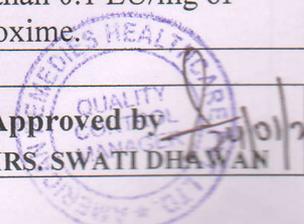


Title : Certificate of Analysis Finished Product

Product Name	FUXORIME Injection	A.R. No.	ARB/FP/25/023
Generic Name	Cefuroxime Injection IP 750mg	Sampled qty.	45 vials
Batch No.	ARB2526A	Sampled by	Aakash
Batch Size	15,000 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 $\pm 10\%$	-1.94% & +1.75%
4.	Average weight	Informative $\pm 2\%$.	820.8 mg
5.	Related Substance (By HPLC)		
	Descarbamoyl-cefuroxime	NMT 1.0%	0.59%
	Area of any other secondary peak	NMT 1.0%	0.20%
	Sum of areas of all the secondary peaks	NMT 3.0%	0.33%
6.	pH	6.0 to 8.5	6.59
7.	Water (By K.F.)	NMT 3.5%w/w	2.97%w/w
8.	Particulate Matter		
	(A.) Light Obscuration Particle Count Test (1.) Particles $\geq 10\mu\text{m}$ (2.) Particles $\geq 25\mu\text{m}$	NMT 6000/vial NMT 600/vial	199/vial 5/vial
	(B.) Visual Particle	The reconstituted solution should be essentially free from extraneous, mobile undissolved particles, other than gas bubbles, unintentionally that can be observed on visual inspection.	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 SHARMA	Checked by  NAME: RAMANJEET SINGH	Approved by  NAME: MRS. SWATI DHAWAN
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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 Injection	A.R. No.	ARB/FP/25/023
Generic Name	Cefuroxime Injection IP 750mg	Sampled qty.	45 vials
Batch No.	ARB2526A	Sampled by	Aakash
Batch Size	15,000 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

11.	Assay: (By HPLC) Each glass vial contains:	Observation		
Ingredients	Labeled Claim	Found	% of labeled amount	Limits % of labeled amount
Cefuroxime Sodium (Sterile) IP Eq. to Cefuroxime	750 mg	755.1538 mg	100.69%	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/HIS.

Analysis by <u>24/01/2026</u> NAME: NIKHIL SHARMA	Checked by <u>24/01/26</u> NAME: RAMANJEET SINGH	Approved by <u>24/01/26</u> NAME: MRS. SWATI DHAWAN
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