AMERICAN REMEDIES HEALTHCARE PVT. LTD.

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

Title : Certificate of Analysis Finished Product

Product Name	**FOXATIM** Injection	**A.R. No.**	ARB/FP/26/030
Generic Name	Cefotaxime for Injection IP 1gm	**Sampled qty.**	45 Vials
Batch No.	ARB2627A	**Sampled by**	Shivanand
Batch Size	30,000 Vials	**Sampled on**	17/10/2026
Mfg. Date	10/2026	**Date of Testing**	18/10/2026
Exp. Date	09/2028	**Date of Release**	02/11/2026

S. No.	Tests	Specifications	Observations
1.	Description	An Off-white to pale yellow, crystalline powder filled in clear glass vial.	A white dry powder filled in clear glass vial.
2.	Identification (By HPLC)	**A.** In, assay, the principal peak in the chromatogram obtained with the test solution corresponds to the peak in the chromatogram obtained with the working reference solution.	Complies
	(By Chemical)	**B.**Gives the reactions of sodium salts.	Complies
3.	Uniformity of weight	Average weight ±10%	-1.82% & +1.56%
4.	Average weight	Informative ±2%.	1095.5 mg
5.	pH	4.5 to 6.5	5.83
6.	Water	NMT 3.0%	1.66%
7.	Related Substances (By HPLC)		
	Area secondary peak	NMT 1.0%	0.9774%
	Sum of all secondary peak	NMT 4.0%	2.8417%
8.	Particulate Matter (IHS)		
	(A.) Light Obscuration Particle Count test		
	(1.) Particles ≥10µm	NMT 6000/vial	1,631/vial
	(2.) Particles ≥25µm	NMT 600/vial	15/vial
	(B.) Visual Particle	The reconstituted solution should be essentially free from extraneous, mobile undissolved particles, other than gas bubbles, unintentionally that can be seen with naked eye.	Complies
9.	Sterility	No microbial growth should be observed.	Complies
10.	Bacterial Endotoxins	NMT 0.20 EU/mg of Cefotaxime.	Less than 0.20 EU/mg of Cefotaxime.
11.	Assay: (By HPLC) Each glass vial contains:		

Analysis by	Checked by	Approved by
02/11/26	02/11/26	02/11/26
NAME: NIKHIL SHARMA	NAME: SURJEET KAUR	NAME: MRS. SWATI DHAWAN

Name : Mrs. Swati Dhawan
Designation : QC HOD
Qualification : M.Sc. Clinical Microbiology
Registration No. : [illegible]

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Care & Cure with Research

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Ingredients	Labeled Claim	Found	% of labeled amount	Limits % of labeled amount
Cefotaxime Sodium (Sterile) IP Eq. to Anhydrous Cefotaxime	1000 mg	972.0925 mg	97.21%	90.0 to 115.0%

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.~~

Name : Mrs. Swati Dhawan
Designation : QC HOD
Qualification : M.Sc. Clinical Microbiology
Registration No. : 580406039598
Name of the Certificate Issuer: Jalandhar

Analysis by	Checked by	Approved by
02/11/26	02/11/26	02/11/26
NAME: NIKHIL SHARMA	NAME: SURJEET KAUR	NAME: MRS. SWATI DHAWAN