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/26/031
Generic Name	Cefotaxime for Injection IP 1gm	**Sampled qty.**	45 Vials
Batch No.	ARB2628A	**Sampled by**	Shivanand
Batch Size	10,200 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%	-1.83% & +1.72%
4.	Average weight	Informative ±2%.	1096.3 mg
5.	pH	4.5 to 6.5	5.88
6.	Water	NMT 3.0%	1.94%
7.	Related Substances (By HPLC)		
	Area secondary peak	NMT 1.0%	0.9874%
	Sum of all secondary peak	NMT 4.0%	3.0290%
8.	Particulate Matter (IHS)		
	(A.) Light Obscuration Particle Count test (1.) Particles ≥10µm (2.) Particles ≥25µm	NMT 6000/vial NMT 600/vial	1,411/vial 8/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
NAME: NIKHIL SHARMA 02/11/26	NAME: SURJEET KAUR 02/11/26	NAME: MRS. SWATI DHAWAN

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

Format No: PB/QC/045/F05-00

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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	958.9386 mg	95.89%	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. Microbiology
Registration No. : [illegible]
Punjab Technical University, Jalandhar

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