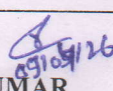
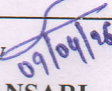


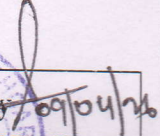
Title	:	Certificate of Analysis Finished Product
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Product Name	AZIDIME Injection	A.R. No.	ARB/FP/25/031
Generic Name	Ceftazidime for Injection IP 1gm	Sampled qty.	45 Vials
Batch No.	ARB2541A	Sampled by	Aakash
Batch Size	10,000 Vials	Sampled on	25/03/2026
Mfg. Date	03/2026	Date of Testing	26/03/2026
Exp. Date	02/2028	Date of Release	09/04/2026

S. No.	Tests	Specifications	Observations
1.	Description	A white or almost white 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. Gives the reactions of sodium salts and reaction A of carbonates.	Complies Complies
3.	Uniformity of weight	Average weight $\pm 10\%$	-1.33% & +1.87%
4.	Average weight	Informative $\pm 2\%$.	1307.2 mg
5.	pH	5.0 to 7.5	6.09
6.	Loss On drying	NMT 13.5%	12.76%
7.	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 solution should be essentially free from extraneous, mobile undissolved particles, other than gas bubbles, unintentionally that can be observed on visual inspection.	10/vial 1/vial Complies
8.	Pyridine	NMT 0.4%	0.20%
9.	Sodium Carbonate (By AAS)	8.0% to 10.0%	9.059%
10.	Sterility	No microbial growth should be observed.	Complies
11.	Bacterial Endotoxins	NMT 0.10 EU/mg of Ceftazidime.	Less than 0.10 EU/mg of Ceftazidime.
12.	Assay: (By HPLC) Each vial contains:		

Analysis by 
NAME: SACHIN KUMAR

Checked by 
NAME: SUFIYAN ANSARI

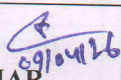
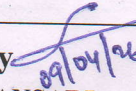
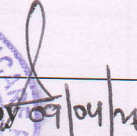
Approved by 
NAME: MRS. SWATI DHAWAN
MANAGER

Title
:
Certificate of Analysis Finished Product

Product Name	AZIDIME Injection	A.R. No.	ARB/FP/25/031
Generic Name	Ceftazidime for Injection IP 1gm	Sampled qty.	45 Vials
Batch No.	ARB2541A	Sampled by	Aakash
Batch Size	10,000 Vials	Sampled on	25/03/2026
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Ingredients	Labeled Claim	Found	% of labeled amount	Limits % of labeled amount
Ceftazidime (Sterile) IP Eq. to anhydrous Ceftazidime (A sterile mixture of sterile Ceftazidime & Sodium carbonate)	1000 mg	1039.0989 mg	103.91%	90.0 to 105.0%

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/HS.

Analysis by 
NAME: SACHIN KUMAR
Checked by 
NAME: SUFIYAN ANSARI
Approved by 
NAME: MRS. SWATI DHAWAN