

Add Pace To Health

Pace Biotech

Surajpur, Paonta sahib Dist. Sirmour (H.P)

Title	: Certificate of Analysis Finished Product		
Product Name	AZEECAN 500 Injection	A.R. No.	NB/FP/24/145
Generic Name	Azithromycin for Injection USP 500mg	Sampled qty.	45 vials
Batch No.	N24124A	Sampled by	Vineet
Batch Size	7550 Vials	Sampled on	24/08/2024
Mfg. Date	08/2024	Date of Testing	24/08/2024
Exp. Date	07/2026	Date of Release	08/09/2024

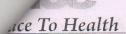
S. No.	Tests	Specifications	Observations
1.	Description	A white dry powder filled in clear glass vial.	A white dry powder filled in clear glass vial.
2.	Identification	In the chromatogram obtained the retention time of major peak in test solution is correspondence with RT of major peak in working standard solution.	Complies
3.	Uniformity of weight	Average weight ±10%	-2.44% & +3.25%
4.	Average weight	Informative.	1089.89 mg
5.	рН	6.4 to 6.8	6.57
6.	Uniformity of Dosage units	Complies as per USP.	Complies
7.	Impurities	Charled Inc.	
	Limit of Azithromycin N-oxide	Complies as per USP	Complies
	Limit of Aminoazithromycin, formamido Analog	Complies as per USP	Complies
	Any other unspecified impurity	NMT- 0.2%	Complies
	Total Impurities	NMT- 3.0%	Complies
8.	Particulate Matter (A.) Light Obscuration Particle Count Test 1.Particles ≥10 μm 2.Particles ≥25 μm	NMT-6000/vial NMT-600/vial	89/vial 1/vial
	(B.) Visual	The solution is free from particles of foreign matter particles that can be observed on visual inspection.	Complies
9.	Water	NMT- 2.0% w/w	1.59%w/w
10.	Sterility	No microbial growth should be observed.	Complies
11.	Bacterial Endotoxins	NMT- 0.7 USP EU/mg of Azithromycin.	Less than- 0.7 USP EU/mg Azithromycin.
12.	Assay: Each glass vial Contains:		PAON

Analysis by

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

Checked by

Approved by m



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Ingredients	Labeled Claim	Found	% of labeled amount	Limits % of labeled amount
Azithromycin Dihydrate (Sterile) IP Lyophilized	ilia cinomi	wyrach chini		
Eq. to anhydrous Azithromycin	500 mg	487.23 mg	97.45%	90.0.0% to 110.0%

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

Checked by W

