



LIFE VISION HEALTHCARE
Plot No. 138-142, EPIP, Phase-1, Jharmajri,
Baddi, Distt.: Solan (H.P.)-174103

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DEPARTMENT: QUALITY CONTROL

CERTIFICATE OF ANALYSIS
(FINISHED PRODUCT)

Product Name	UZACEF - 50	Volume: 15 g/30 ml
Generic Name	Cefpodoxime Proxetil Suspension IP	
Batch No.	LD - 0575	A.R.No. QC/FP/LD - 0575
Mfg. Date	DEC. 2023	Batch Size 90 Ltr.
Exp. Date	MAY 2025	Sample quantity 12 X 15 g/30 ml
Date of Receipt	20-12-2023	Date of release 28-12-2023
Date of analysis	21-12-2023	Mfg. Lic. No. MNB/09/804 & MB/09/805
Marketed by	HINDUZA LIFE SCIENCE PVT. LTD.	

S. No.	TEST	SPECIFICATION	OBSERVATION	
1	Description	White powder fill in white HDPE bottles, reconstituted with water produced a yellow coloured suspension.	White powder filled in white HDPE bottles, reconstituted with water produced a yellow coloured suspension.	
2	Identification Cefpodoxime Proxetil IP	In assay by HPLC, the RT (Retention Time) of the sample solution should match with the RT of the standard solution.	In assay by HPLC, the RT (Retention Time) of the sample solution match with the RT of the standard solution.	
3	pH	4.00 to 5.50	4.63	
4	Weight per ml	1.00 to 1.30 g	1.1202 g	
5	Average net weight	NLT 15 g/30 ml	15.0196 g/30 ml	
6	Water	NMT 1.5 %	1.12 %	
7	Uniformity of weight	± 9.0 %	Within the limit	
8	Assay			
	Each 5 ml of the reconstituted suspension contains	Claim	Result	Limit
	Cefpodoxime Proxetil IP Equivalent to Cefpodoxime	50 mg	49.955 mg (99.91 %)	45.0 mg to 55.0 mg (90.0 % to 110.0 %)
	Within 7 days of reconstituted suspension			
	Cefpodoxime Proxetil IP Equivalent to Cefpodoxime	50 mg	48.41 mg (96.82 %)	NLT 40.0 mg (NLT 80.0 %)

Remark: The above finished product complies to as per IP & In House specification

	Analyzed by Officer-QC	Checked by Executive-QC	Approved by Manager-QC
Signature			for
Date	28-12-2023	28-12-2023	28-12-2023