

**KamlaAmrut Pharmaceuticals LLP****QUALITY CONTROL DEPARTMENT****CERTIFICATE OF ANALYSIS**

Generic Name	: LEVOFLOXACIN INFUSION IP		
Product Name	: 0.5% LEVOFLOXACIN INFUSION IP - 100ML(LEVOXICAN IV) - AMERICAN - NH		
Batch No.	: AC2376001	A.R. No.	: KAPL/FP/26/AC2376001
Batch Size	: 1080 Ltr	Quantity sampled	: 25 Nos
Fill Volume	: 100 ml	Received Date	: 09/04/2026
Mfg. Date	: 01/04/2026	Testing Date	: 09/04/2026
Exp. Date	: 31/03/2028	Completion Date	: 30/04/2026

Sr. No.	Tests	Observation	Specifications	Reference to Test Method
1.0	Description	A clear and colourless solution	A clear & yellow solution essentially free from visible particles	IP Monograph
2.1	Identification	confirms	The chromatogram obtained with the test solution corresponds to the chromatogram obtained with standard solution	IP Monograph (2.4.14)
3.0	pH	4.67	Between 3.80 to 5.80	IP Monograph(2.4.24)
4.0	Extractable Volume	102 ml	The volume is not less than the nominal volume.	IP Monograph
5.1	Particulate Matter	15.0 particles per container 0.0 particles per container	Not more than 6000 particles per container Not more than 600 particles per container	IP Monograph(2.5.9)
6.0	Assay of Sodium Chloride	0.9018 %w/v 100.20 %	Between 95.0% to 105.0% (Label Claim: 0.9% w/v)	IP Monograph
7.0	Assay of Levofloxacin	0.5005 %w/v 100.10 %	Between 90.0% to 120.0% (Label Claim: 0.5% w/v)	IP Monograph(2.4.14)
8.0	Bacterial Endotoxins	<0.25 EU/ml	Not more than 0.50 EU/ml	IP Monograph(2.2.3)
9.0	Sterility	sterile	It must be sterile	IP Monograph(2.2.11)

Remarks : The finish product confirms with the defined specifications.

	Analysed by	Checked by	Verified by
Date	30/04/2026	30/04/2026	30/04/2026
Name	Jinal Patel	Baldev Makwana	Kirit Patel
Designation	OFFICER	QC Executive	QA Manager

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Page 1 of 1

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