

CERTIFICATE OF FINISH PRODUCT ANALYSIS

Product Name:	CANAPARIN 5K (Heparin Sodium Injection IP 25000 IU)		
Batch No.	24VJ044D	A.R. No.	FP240071
Batch Size	5000 Nos	Sample Quantity	60 Nos
Mfg. Date	10/2024	Date of Testing	08/10/2024
Exp. Date	09/2026	Date of Release	24/10/2024
Reference Batch No	24VJ044	MFG. Lic. No.	N-MB-2023/263

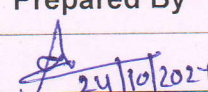
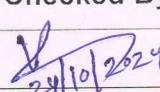
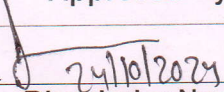
Sr. No.	Test	Specification	Observation
1.	Description	A clear, colourless or straw-coloured solution, free from turbidity and matter which deposit on standing.	A clear, colourless, free from turbidity and matter which deposit on standing.
2.	Identification		
	A. By Assay	By Assay	Complies
	B. It gives reaction of Sodium Salt	It gives reaction of Sodium Salt	Complies
3.	pH	Between 5.0 to 7.5.	6.96
4.	Extractable Volume	NLT 5.0 ml	5.0 ml
5.	Particulate Matter	Injections that are solutions, when examined under suitable conditions of visibility, are clear and practically free from particles that can be observed on visual inspection by the unaided eye.	Complies
6.	Sterility	Should be complies	Sterile
7.	Bacterial Endotoxins	NMT 0.03 Endotoxin unit per unit of heparin.	Less than 0.03 Endotoxin unit per unit of heparin.

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8.	Assay:		
	Each ml Contains:	NLT 4500 and NMT 5500 IU/ml	5060.0 IU/ml
	Heparin Sodium IP 5000 IU	(90.0% to 110 %)	101.20 %

Remark: The sample is complies as per IP /BP/ USP/IH specification.

	Prepared By	Checked By	Approved By
Sign/Date	 24/10/2024	 24/10/2024	 24/10/2024
Name	Abhay Mishra	Vikram Jeet	Bhupinder Navet
Designation	Sr. Officer	Executive QC	Head QC