

CERTIFICATE OF FINISH PRODUCT ANALYSIS

Product Name:	Canaprin (Heparin Sodium Injection IP 1000 IU)		
Batch No.	24VG006B	A.R. No.	PV240713
Batch Size	3000 Nos	Sample Quantity	50 Nos
Mfg. Date	07/2024	Date of Testing	20/07/2024
Exp. Date	06/2026	Date of Release	05/08/2024
Reference Batch No	24VG006	MFG. Lic. No.	N-MB-2023/263


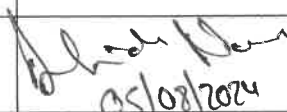
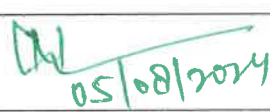
Sr. No.	Test	Specification	Observation
1.	Description	A clear, colourless or straw coloured solution, free from turbidity and matter which deposit on standing.	A clear, colourless, free from turbidity and matter which deposit on standing.
2.	Identification		
	A. By Assay	By Assay	Complies
	B. It gives reaction of Sodium Salt	It gives reaction of Sodium Salt	Complies
3.	Extractable Volume	NLT 5.0 ml	5.0 ml
4.	pH	Between 5.0 to 7.5.	6.30
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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Assay:			
8. Each ml Contains:			
Heparin Sodium IP 1000 IU	NLT 900 and NMT 1100 IU/ml	999.5 IU/ml	

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

	Prepared By	Checked By	Approved By
Sign/Date	 05/08/2024	 05/08/2024	 05/08/2024
Name	Vikram Jeth	Rishinder Navel	Nitish Kumar
Designation	Executive-QC	Head-QC	Head-QA