

**CERTIFICATE OF FINISH PRODUCT ANALYSIS**

<b>Product Name:</b>	<b>Canaprin (Heparin Sodium Injection IP 1000 IU/ml)</b>		
<b>Batch No.</b>	24VG005H	<b>A.R. No.</b>	PV240665
<b>Batch Size</b>	4000 Nos	<b>Sample Quantity</b>	50 Nos
<b>Mfg. Date</b>	07/2024	<b>Date of Testing</b>	19/07/2024
<b>Exp. Date</b>	06/2026	<b>Date of Release</b>	03/08/2024
<b>Reference Batch No</b>	24VG005	<b>MFG. Lic. No.</b>	N-MB-2023/263

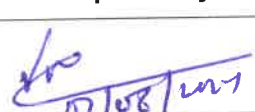
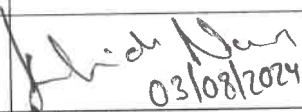
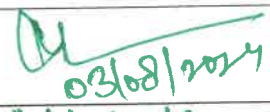
<b>Sr. No.</b>	<b>Test</b>	<b>Specification</b>	<b>Observation</b>
1.	<b>Description</b>	A clear, colourless or straw coloured solution, free from turbidity and matter which deposit on standing.	A clear, colourless, free from turbidity and matter which deposit on standing.
2.	<b>Identification</b>		
	<b>A. By Assay</b>	By Assay	Complies
	<b>B. It gives reaction of Sodium Salt</b>	It gives reaction of Sodium Salt	Complies
3.	<b>Particulate Matter</b>	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
4.	<b>Extractable Volume</b>	NLT 5.0 ml	5.0 ml
5.	<b>pH</b>	Between 5.0 to 7.5.	6.20
6.	<b>Sterility</b>	Should be complies	Sterile
7.	<b>Bacterial Endotoxins</b>	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: Heparin Sodium IP 1000 IU	NLT 900 and NMT 1100 IU/ml (90.0% to 110 %)	998.0 IU/ml 99.8%
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Remark: The sample is complies as per IP /BP/ USP/IH specification.

	Prepared By	Checked By	Approved By
Sign/Date	 03/08/2024	 03/08/2024	 03/08/2024
Name	Vikram Jeth	Bhupinder Navel	Nitish Kumar
Designation	Executive - QC	Head - QC	Head - QA