

Title	:	Certificate of Analysis Finished Product
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Product Name	EPTOCAN Injection	A.R. No.	ARN/FP/26/077
Generic Name	Phenytoin Sodium Injection 2ml	Sampled qty.	45 Ampoules
Batch No.	ARN2666A	Sampled by	Shivanand
Batch Size	46,284 Ampoules	Sampled on	04/09/2026
Mfg. Date	08/2026	Date of Testing	04/09/2026
Exp. Date	07/2028	Date of Release	19/09/2026

S. No.	Tests	Specifications	Observations
1.	Description	A clear colourless solution filled in amber glass ampoule.	A clear colourless solution filled in amber glass ampoule.
2.	Identification (By IR) (By HPLC)	A. Residue obtained from the sample meets the requirement, The absorption spectrum of the test preparation should exhibits maxima at same wavelength with that obtained with similar preparation of Phenytoin sodium working standard. B. The retention time of the major peak of the sample solution corresponds to that of the standard solution as obtained in the assay.	Complies Complies
3.	Nominal fill volume	NLT 2ml	2.0 ml
4.	Extractable Volume	NLT Nominal volume or NMT 110% of Nominal volume.	2.1 ml
5.	pH	10.0 to 12.3	11.08
6.	Particulate Matter (A) Light Obscuration Particle Count Test 1. Particles $\geq 10 \mu m$ 2. Particles $\geq 25 \mu m$ (B) Visible Particles	NMT 6000/Ampoule NMT 600/Ampoule The solution should be essentially free from extraneous, mobile undissolved particles, other than gas bubbles unintentionally.	32/Ampoule 0/Ampoule Complies
7.	Sterility	No microbial growth should be observed.	Complies
8.	Bacterial Endotoxins (IHS)	NMT 0.3 EU/mg of Phenytoin Sodium.	Less than 0.3 EU/mg of Phenytoin Sodium.
9.	Assay: Each ml contains:		

Ingredients	Labeled Claim	Found	% of labeled amount	Limits % of labeled amount
Phenytoin Sodium IP	50 mg	50.19772 mg	100.40%	90.0% to 110.0%

Remarks: In the opinion of the undersigned the sample referred to above is/is not of the standard quality as The Drug & Cosmetic Act 1940 and the rules made there under. Complies/not complies as per IP/BP/USP/IHS

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