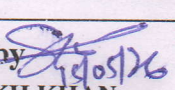
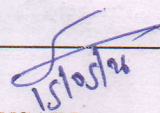


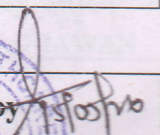
Title : Certificate of Analysis Finished Product

Product Name	CLOCIVIR Infusion	A.R. No.	ARN/FP/26/008
Generic Name	Aciclovir Intravenous Infusion IP 500mg	Sampled qty.	45 vials
Batch No.	ARN2605A	Sampled by	Rakesh
Batch Size	30,000 Vials	Sampled on	30/04/2026
Mfg. Date	04/2026	Date of Testing	01/05/2026
Exp. Date	03/2028	Date of Release	15/05/2026

S. No.	Tests	Specifications	Observations
1.	Description	A white or almost white, crystalline powder filled in clear glass vial.	A white dry powder filled in clear glass vial.
2.	Identification By UV	A. When examined in the range 230 nm to 360 nm the solution prepared in the Assay shows an absorption maximum at about 255 nm and a broad shoulder at about 274 nm.	Complies
	By TLC	B. In the test of Guanine, the principal spot in the chromatogram obtained with test solution (b) corresponds to that in the chromatogram obtained with reference solution (a).	Complies
	By Chemically	C. It gives reaction (A) of sodium salts.	Complies
3.	Uniformity of weight	Average weight $\pm 10\%$	-2.16% & +3.01%
4.	Average weight	Informative $\pm 2\%$.	601.4 mg
5.	Particulate Matter		
	(A.) Light obscuration Particle Count Test (1.) Particles $\geq 10\mu\text{m}$ (2.) Particles $\geq 25\mu\text{m}$	NMT 6000/vial NMT 600/vial	247/vial 2/vial
	(B.) Visual Particle	The solution should be essentially free from extraneous, mobile undissolved particles, other than gas bubbles, unintentionally that can be seen with naked eye.	Complies
6.	Appearance of solution	The solution is not more opalescent than opalescence standard OS2, and not more intensely coloured than reference solution BY55.	Complies
7.	pH	10.7 to 11.7	11.17

Analysis by 
NAME: SHAHRUKH KHAN

Checked by 
NAME: SUFIYAN ANSARI

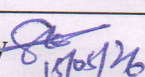
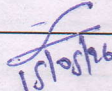
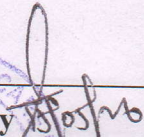
Approved by 
NAME: MRS. SWATI DHAWAN

Title : Certificate of Analysis Finished Product

Product Name	CLOCIVIR Infusion	A.R. No.	ARN/FP/26/008
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8.	Related Substances (By TLC)	Any secondary spot with an R _F value greater than that of the principal spot in the chromatogram obtained with the test solution is not more intense than the spot in the chromatogram obtained with reference solution (0.5 per cent).	Complies		
9.	Guanine (By TLC)	Any secondary spot corresponding to guanine in the chromatogram obtained with test solution (a) is not more intense than the spot in the chromatogram obtained with reference solution (b) (1.0 per cent).	Complies		
10.	Sterility	No microbial growth should be observed.	Complies		
11.	Bacterial Endotoxins	NMT 0.174 EU/mg of Aciclovir.	Less than 0.174 EU/mg of Aciclovir.		
12.	Assay: (By UV) Each glass vial Contains:				
Ingredients		Labeled Claim	Found	% of labeled amount	Limits % of labeled amount
(Sterile) Aciclovir Sodium (Lyophilized) Eq. to Aciclovir IP		500 mg	514.8123 mg	102.96%	95.0% to 105.0%

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.

Analysis by  NAME: SHAHRUKH KHAN	Checked by  NAME: SUFIYAN ANSARI	Approved by  NAME: MRS. SWATI DHAWAN
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