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CERTIFICATE OF ANALYSIS
FINISH PRODUCT - QUALITY CONTROL

Product Name	**SUCONIL INJECTION**		
Generic Name	**Suxamethonium Chloride Injection BP 50 mg per ml**		
Batch No.	A04626002	**A.R. No.**	FP/09/26/00002
Mfg. Date	08/2026	**Sampling Quantity**	65 Ampoules
Exp. Date	07/2029	**Date of Sampling**	02/09/2026
Batch Size	200 Liter	**Date of Analysis**	02/09/2026
Pack Quantity	NA	**Date of Release**	16/09/2026
Ref. Spec. No.	FP/SPC/0025-01	**Pack Size/Type**	NA

Sr. No.	Test	Specification	Result
1.	Description	A clear solution.	A clear solution.
2.	Identification		
	By Chemical Method	The chloroform layer is yellow.	Complies
3.	Acidity	3.0 to 5.0.	3.7
4.	Hydrolysis products	The volume of 0.1 M sodium hydroxide VS required for the preliminary neutralization in the Assay is not more than one tenth of the total volume of 0.1 M sodium hydroxide VS required for the preliminary neutralization and the hydrolysis.	Complies
5.	Extractable Volume	NLT as per label claim	2.1 ml
6.	Particulate Matter		
	≥ 10 µm	NMT 6000 no. of particles per container	157 particles per container
	≥ 25 µm	NMT 600 no. of particles per container	10 particles per container
7.	Sterility	Should be sterile	Sterile
8.	Bacterial Endotoxins	NMT 2.0 USP Endotoxin Units/mg of succinylcholine chloride.	Less than 2.0 EU/mg
9.	Assay	90.0 % to 107.5 %.	100.5 %

Remark: The Finish Product test results are Complies / ~~Does not complies~~ as per ~~IH/IP/BP/USP~~ Specification.

Function	Prepared By	Reviewed By	Approved By
Name	**Pranita Wadje**	**Maya Sirsat**	**Kushal Deshmukh**
Designation-Department	**Officer – QC**	**Executive – QC**	**Asst. Manager – QC**
Sign & Date	06/10/2026	06/10/2026	06/10/2026

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