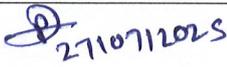
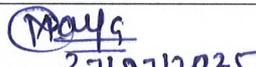
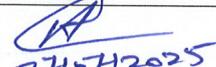


Format No.: KBPL/QC/030/F03-01			
Product Name	SUCONIL INJECTION		
Generic Name	Suxamethonium Chloride Injection BP 50 mg per ml		
Batch No.	K30325003	A.R. No.	FP/07/25/00026
Mfg. Date	06/2025	Sampling Quantity	65 Ampoules
Exp. Date	05/2028	Sampling Date	23/07/2025
Batch Size	110 Litre	Date of Analysis	23/07/2025
Pack Quantity	NA	Date of Release	27/07/2025
Ref. Spec. No.	FP/SPC/0025-01	Page No.	Page 1 of 1
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.	4.2
4.	Hydrolysis products	The volume of 0.1 M sodium hydroxide VS required for the preliminary neutralisation in the Assay is not more than one tenth of the total volume of 0.1 M sodium hydroxide VS required for the preliminary neutralisation and the hydrolysis.	Complies
5.	Extractable Volume	NLT as per label claim	2.2 ml
6.	Particulate Matter		
	≥ 10 µm	NMT 6000 no. of particles per container	#Under analysis
	≥ 25 µm	NMT 600 no. of particles per container	
7.	Sterility	Should be sterile	*Under observation
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 %.	101.3 %

CONCLUSION: The Finish Product Sample ~~Does not complies~~ ^{Complies} as per HH/IP/BP/USP Specification.

Note: 1) #Particulate matter test is under analysis. 2) *Sterility test is under observation.

Function	Prepared By	Reviewed By	Approved By
Name	Pranita Wadje	Maya Sirsat	Avnish Kumar
Designation-Department	Officer - QC	Executive - QC	Manager - QC
Sign & Date	 27/07/2025	 27/07/2025	 27/07/2025

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