ALVENTA PHARMA LIMITED

VILL. KISHANPURA, TEHSIL BADDI- NALAGARH ROAD, DISTT.- SOLAN (H.P) 174101

Quality Control Department

CERTIFICATE OF ANALYSIS (FINISHED PRODUCT)

Product Name	Diclocan S-AQ	**A.R. No.**	FG/G/25A1932
Generic Name	Diclofenac Sodium Injection IP 25 mg	**Sample Quantity**	65 Ampoules
Batch No.:	AA51007	**Sample Received on**	13/11/2025
Batch Size:	1.0 Lac	**Analysis Date**	13/11/2025
Mfg. Date.	10/2025	**Release Date**	28/10/2025
Exp. Date	09/2027	**Page No.:**	Page 1 of 1

Sr. No.	Test Parameter	Acceptance Criteria	Result
1.	**Description**	A clear colourless liquid filled in 3 ml clear glass ampoule having a blue dot over the neck.	A clear colourless liquid filled in 3 ml clear glass ampoule having a blue dot over the neck.
2.	**Identification**	The principal spot in the chromatogram obtained with the test solution corresponds to that in the chromatogram obtained with the reference solution.	Complies
3.	**pH**	Between 8.1 to 9.0	8.6
4.	**Extractable volume**	Not less than 3.0 ml	3.05 ml
5.	**Particulate Matter**	The Solution should be clear, colourless and free from any visible particle.	The Solution is clear, colourless and free from any visible particle.
6.	**Sterility**	Should be sterile	Sterile
7.	**Bacterial Endotoxins Tests**	NMT 2.33 EU/mg of Diclofenac Sodium.	Less than 2.33 EU/mg Diclofenac Sodium.
8.	**Assay:**		

Each ml contains:	Claim	Limit	mg	%
Diclofenac Sodium IP	**25 mg**	Between 95.0 % to 105.0 % of stated amount of Diclofenac Sodium. (Between 23.75 mg to 26.25 mg)	25.0336 mg	100.13%
Benzyl Alcohol IP (As Preservative)	**4%v/v**	Between 90.0% to 110.0% labeled amount Benzyl Alcohol. (Between 3.6 %v/v to 4.4 %v/v)	3.9658%v/v	99.14%

Remarks: The above test parameters are complies/ ~~not complies~~ as per IP/~~BP~~/USP & In-House Specification.

Particulars	Prepared By	Checked By	Approved By
Name	Bavneen Kumar	Manilal	Kuljeet Kaur
Designation	Executive	Sr. Executive QC	Head QC
Signature			
Date	28/10/2025	28/10/2025	28/10/2025

Format No.: SOP/QC/029/F02-01