Lanark Laboratories

NEAR ELECTRICITY SUB DIVISION, GONDPUR, PAONTA SAHIB DISTT.SIRMOUR HP INDIA 173025

CERTIFICATE OF ANALYSIS (FINISHED PRODUCT)

(The Drugs & Cosmetics Act 1940 at the under form 39)

Work Order No. : 252611719 **Document No. :** 252613950

Ref. STS No.	8229	**Product Code**	FG01716
Product Name	TAB. AMOLINE-5 (ZLP)		
Generic Name	AMLODIPINE TABLETS IP 5MG		
Batch No.	425-409	**Date Of Sampling**	12/06/2025
Batch Size	1000000.000 TAB **Cumulative Qty.** 985520.000 TAB	**Sample Quantity**	60.000 TAB
Mfg. Date	06/2025 **Exp. Date** 05/2027	**Quantity Release**	985460.000 TAB
A. R. No.	PL25FG01781	**Date Of Release**	12/06/2025

S NO.	Tests	Specification	Results
1	Description	A white colour round shape biconvex uncoated tablets, having central break line on one side and plain on other side.	A white colour round shape biconvex uncoated tablets, having central break line on one side and plain on other side.
2	Identification	Should be positive for Amlodipine Besilate.	Complies
3	Average weight	210 mg ± 7.5% of Average weight	211.26 mg
4	Uniformity of weight	± 7.5 % of Average weight	2.53 % to +3.53%
5	Disintegration Time	Not more than 15:00 minutes	58 Seconds.
6	Friability	Not more than 1.0% w/w	0.38 %w/w
7	Dissolution	Not less than 75.00 %	98.38% to 102.63%
8	Uniformity of content	85.00 to 115.00%	96.25% to 106.29%
9	Composition: Each uncoated tablet contains:	Limit	Assay
10	Amlodipine Besilate I.P. Eq. to Amlodipine 5 mg.	90.0 to 110.0% w/w of the stated amount of Amlodipine Besilate.	104.36 %

Remarks The above sample complies/~~does not comply~~ with prescribed standards of Quality with respect to above tests as per IP/~~IBP/USP/In house~~ specification.

LANARK LABORATORIES QC APPROVED

Prepared By Madhukant **Checked By** QCM **Approved By**

Date 12/06/2025 **Date** 12/06/2025 **Date** 12/06/2025

Appendix No.:- **Confidential**