

**ZEE LABORATORIES LTD**

Behind 47, Industrial Area, Gondpur Paonta Sahib Sirmour HP INDIA 173025

CERTIFICATE OF ANALYSIS (FINISHED PRODUCT)

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

Work Order No. :

Document No. : 252641687

Ref. STS No.	35	Product Code	FG03022
Product Name	TAB. AMOLINE-LS		
Generic Name	AMLODIPINE & LISINOPRIL TABLETS		
Batch No.	425-2497	Date Of Sampling	26/11/2025
Batch Size	50000.000 Tab	Sample Quantity	60.000 Tab
Mfg. Date	11/2025	Exp. Date	10/2027
A. R. No.	ZL25FG09396	Date Of Release	29/11/2025
S NO.	Tests	Specification	Results
1	Description	A white colour, rounded, uncoated tablets and both sides are smooth.	Complies
2	Identification	Should be positive for Amlodipine Besylate and Lisinopril Dihydrate.	Complies
3	Average Weight of Tablet	120.0 mg $\pm$ 7.5 %	122.4 mg
4	Uniformity of Weight	Average weight of tablet $\pm$ 7.5%.	Complies
5	Disintegration Test	Not more than 15 minutes	06:10 Min
6	Friability Test	NMT 1.0 % w/w	0.22 %
7	Dissolution Test	NLT 75% (Q) of the stated amount of Amlodipine Besylate is dissolved.	88% to 93 %
8	Dissolution Test	NLT 75% (Q) of the stated amount of Lisinopril Dihydrate is dissolved.	90% to 95%
9	Assay Each Uncoated tablet contains Amlodipine Besylate I.P Eq. to Amlodipine 5 mg	NLT 90.0% and NMT 110.0% of the labeled amount of Amlodipine Besylate	102.4 %
10	Lisinopril Dihydrate I.P Eq.toAnhydrous Lisinopril 5 mg	NLT 90.0% and NMT 110.0% of the labeled amount Lisinopril Dihydrate	103.4 %

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

Particulars	Analysis By	Prepared By	Checked By	Approved By
Sign & Date	 29/11/25	 29/11/2025	 29/11/25	 29/11/25
Name:	Nitin	Mr. Raghuveer	Q.C Manager	Q.C Manager
Designation:	officer QC	officer QC	Sr. Operation	Asst. Manager

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