[redacted]

Sample Name	AMLODIPINE BESILATE TABLETS-IP-10.0mg.	**Mfg Lic No**	[redacted]
Submitted By	Quality Control Advisor. (MMDSL) Shillong.	**Report No**	
Address	DHS LaitumkhrahNHM,EastKhasi Hills Meghalaya.	**Receipt Date**	09-Oct-24
Mfg By	N.S.	**Report Date**	07-Nov-24
Supplied By	N.S.	**Ref No**	N.S.

Batch No	Mfg. Date	Exp Date	Batch Size	Sample Qty
MMDSL/QC-0509	07/2024	06/2026	N.S.	60 TABS.

TEST RESULTS

Date / Period of Performance of test 09/10/2024 to 07/11/2024.

Reference to Protocol :- I.P-2022.

Description :- White round biconvex uncoated tablets.

Identification (by HPLC) :- Complies.

Average weight :- 148.9mg

Related Substances(by HPLC) :- Complies.

Dissolution (by U.V) :- Complies(89.81% to 94.69%) Limit:- NLT 75%+5%

Uniformity of content (by HPLC) :- Within limit

Assay (by HPLC) :- Each uncoated tablet contains:-

Content of	Obtd./Avg.wt.	Claim	Limit Lower	Upper	Method
Amlodipine Besilate eq. to Amlodipine	:- 9.91mg	10.0mg	9.0mg	11.0mg	I.P

END OF TEST REPORT

NOTE:- SAMPLE CONSUMED IN TESTING.

Report:In Opinion of the undersigned,The sample referred to above is of Standard Quality as defined in the Act and the rules made there under for the reason given: Complies as per IP.

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