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Sample Name	AMLODIPINE BESILATE TABLETS-IP.5.0mg.	**Mfg Lic No**	[redacted]
Submitted By	Quality Control Advisor. (MMDSL) Shillong.	**Report No**	
Address	DHS LaitumkhrahNHM,EastKhasi Hills Meghalaya.	**Receipt Date**	29-Apr-24
Mfg By	N.S.	**Report Date**	16-May-24
Supplied By	N.S.	**Ref No**	N.S.

Batch No	Mfg. Date	Exp Date	Batch Size	Sample Qty
MMDSL/QC-0206	02/2024	01/2026	N.S.	60 TABS.

TEST RESULTS

Date / Period of Performance of test 29/04/2024 to 16/05/2024.

Reference to Protocol :- I.P-2022.

Description :- White round biconvex uncoated tablets.

Identification (by HPLC) :- Complies.

Average weight :- 109.63mg.

Related Substances(by HPLC) :- Complies.

Dissolution (by U.V) :- Complies(84.1% to 89.7%) Limit:- NLT 75%+5%

Uniformity of content (by HPLC) :- Within limit.

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

Contents of	Obtd./Av.wt.	Claim	LIMIT Lower	LIMIT Upper	Method
Amlodipine Besilate cal. as Amlodipine	:- 4.97mg	5.0mg.	4.5mg.	5.5mg.	IP.

NOTE:- SAMPLE CONSUMED IN TESTING.

Report:In Opinion of the undersigned,The sample refer
as defined in the Act and the rules made there under
per IP.

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