[redacted]

Sample Name	RANITIDINE HCL TABLETS-IP.150mg.	**Mfg Lic No**	[redacted]
Submitted By	Quality Control Advisor. (MMDSL) Shillong.	**Report No**	
Address	DHS LaitumkhrahNHM,EastKhasi Hills Meghalaya.	**Receipt Date**	19-Mar-24
Mfg By	N.S.	**Report Date**	01-Apr-24
Supplied By	N.S.	**Ref No**	N.S.

Batch No	Mfg. Date	Exp Date	Batch Size	Sample Qty
MMDSL/QC-0153	06/2023	05/2025	N.S.	60 TABS.

RESULT OF ANALYSIS

Date / Period of Performance of test 19/03/2024 to 01/04/2024.

Reference to protocol :- I.P-2022.

Description :- Orange coloured round biconvex film coated tablets.

Identification (A) (by IR) :- Complies.
(B)(by HPLC) :- Complies.

Average weight :- 177.1mg

Uniformity of weight :- Within limits.

Dissolution(by UV) :- Complies(88.17% to 94.63%) (Limit NLT-80%+5%)

Related Substances (by HPLC):- Complies.

Assay (by HPLC) :- Each film coated tablet contains:-

	Obtd./Av.wt.	Claim	LIMIT Lower	Upper	Method
Content of Ranitidine Hydrochloride calculated as Ranitidine :-	148.89mg	150.0mg.	135.0mg.	165.0mg.	IP.

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.

[redacted]

NOTE : 1. The result listed refer only to the tested samples and applicable parameter. End
2. Sample not drawn by us. Total liability of this laboratory limited to the invoice amount.
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