

Sample Name Amoxycillin & Pot.Clavulanate Tabs-IP. **Mfg Lic No** [REDACTED]
Submitted By Quality Control Advisor. (MMDSL) Shillong. **Report No** [REDACTED]
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 MMDSL/QC-0148 **Mfg. Date** 06/2023 **Exp Date** 09/2025 **Batch Size** N.S. **Sample Qty** 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 :- White oval shaped biconvex film coated tablets.
 Identification (By HPLC) :- Complies.
 Average weight :- 839.2mg
 Uniformity of weight :- Within limits.
 Dissolution for Amoxycillin (by HPLC) :- Complies(90.82% to 95.43%) Limit NLT 85%+5%
 Dissolution for Clavulanic Acid (by HPLC) :- Complies(90.53% to 96.74%) Limit: NLT 80%+5%
 Uniformity of Content by(HPLC) (For Clavulanic Acid) :- Within limits.
 Water(by KF) :- 6.81%w/w Limit:NMT 7.5%w/w

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

			Limit		
Content of	Obtd./Avg.wt.	Claim	Lower	Upper	Method
Amoxycillin Trihydrate eq. to Amoxycillin	:- 249.14mg	250.0mg	225.0mg	300.0mg	IP.
Potassium Clavulanate diluted eq. to Clavulanic Acid	:- 125.06mg	125.0mg	112.5mg	150.0mg	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.

NOTE : 1. The result listed refer only to the tested samples and applicable parameter. Endorsement of products is neither inferred nor implied.
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