FORM 39A |(SEE RULE 150 E (f) UNDER THE DRUGS & COSMETICS ACT 1940 & RULES THEREUNDER)

Generic Name: Amoxycillin and Potassium Clavulanate Tablets IP

Sample Name : Amoxycillin and Potassium Clavulanate Tablets

500mg+125mg

Submitted By : M/s Meghalayan Medical Drugs and Services Limited

(MMDSL)

New Colony Shillong, DHS Laitumkhrah, Office of the Managing Director, MMDSL East Khasi Hills District Meghalaya Shillong (Assam) 793003 District Meghalaya

Shillong (Assam) 793003

Mfd. By : NS

Supplied By : NS Ref.No.

: NS

Batch No. MMDSL/QC-0196 Mfg. Date

02/2024

Format No

Mfg. Lic.No.

Report No.

Date of Receipt

Report Date

: 01/05/2024 Period of Analysis : 18/04/2024-01/05/2024

: 18/04/2024

Stability Condition ;

Batch Size

NS

Sample Qty.

60 Tablets

RESULTS OF ANALYSIS

Reference to protocol

IP-2022

White elongated biconvex film coated tablet having scored on one side. Description

Exp. Date

07/2025

Test Parameters

Average weight

Results

Limit

Identification (by HPLC)

Complies

0.00

1089.63mg

Uniformity of weight

-3.51 to +4.49%

±5.0%

To Comply

Uniformity of Content (by HPLC)

For Potassium Clavulanate Eq. to

NMT 15.0

Clavulanic Acid

Disintegration time

10 to 11 Minutes

NMT 30 minutes

Dissolution (by HPLC)

For Amoxycillin Trihydrate Eq. to

Amoxycillin

For Potassium Clavulanate Eq. to

NLT 80.0% (Q)

NLT 85.0% (Q)

Clavulanic Acid

5.12%w/w

NMT 10.0% w/w

Assay (by HPLC)

Each film coated tablet contains.

95.7 to 96.0% (Avg.95.8%)

91.1 to 91.6% (Avg.91.3%)

Particulars

Water

Result

Claim

Lower

Upper

Method

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: 18/04/2024

: 01/05/2024

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Batch Size

NS

Sample Qty.

60 Tablets

RESULTS OF ANALYSIS

		Claim	Lower	<u>Upper</u>	Method
Amoxycillin Trihydrate eq. to Amoxycillin	503.77mg	500.00 mg	450.00 mg	600.00 mg	IP
Potassium Clavulanate eq. to Clavulanic Acid	122.60mg	125.00 mg	112.5 mg	150.00 mg	IP

^{***}End of Report***