

## Regional Drugs Testing Laboratory

Directorate General of Health Services, Guwahati (India) - 781022

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## **FORM 13** (See rule 46)

CERTIFICATE OF TEST OR ANALYSIS BY GOVERNMENT ANALYST UNDER SECTION 25(1) OF THE DRUGS AND COSMETICS ACT, 1940

Name of Inspector from whom received

: Lalruatadika Varte, Drugs Inspector

Office of the Chief Medical Officer Kolasib, Mizoram-796082

Serial No. and date of Inspector's memorandum

: KLB/03/2025/IPA/DRUGS, 10-JAN-2025

Number of Sample

: NIL

Date of receipt

: 05-FEB-2025

Names of drugs purporting to be contained in the sample : Amoxycillin Trihydrate with Lactic acid bacillus Capsules

(MimoxLB-250 capsules)

Lab. Sample No.	Report No.	Batch No.	Date of Mfg. & Exp	Marketed By	Mfg. By
LSD/GUW/2025-0 26/0014	6UW/LS/2025- 26/39	CF-23012	Jun-2023 May-2025	NA <sub>PO</sub>	Horizon Bioceuticals Pvt. Ltd, Plot No. 3A, Ind. Area, Trilokpur Road, Kala Amb, Distt, Sirmour (H.P.).

6. Condition of seals on

[the packet or on portion of sample or container]

Seals were intact & identical to the specimen impression of the seal received from Drugs Inspector

: Please see below

7. Result of test or analysis with protocols or test or analysis applied

Date of Testing:

From 24-Mar-2025 To 01-Apr-2025

COMPOSITION

Each hard gelatin capsule contains: Amoxycillin Trihydrate I.P. equivalent

Amoxicillin 250 mg.

Protocol Applied: Reference I.P.2022

Sr No.	Test Name	Result	Limits
1	Description	White powder filled in hard gelatin capsules having red colored cap imprinted as "CUREWELL", and white body imprinted "MIMOX-LB 250", in blister pack.	NA
2	Identification	Gives positive test for Amoxycillin Trihydrate.	NA
3	Average filled content	0.3002 g	NA
4	Uniformity of filled content	Complies.	NA

## **Assay**

THE REAL PROPERTY AND PERSONS ASSESSED.	Sr.No	Ingredient Name	Found	Claim	% of claim	Limits	Procedure / Method
	. 1	Amoxycillin Trihydrate eq. to Amoxycillin	244.92 mg/Capsule	250 mg/Capsul e		90 % to 110 %	I.P.2022

In the opinion of the undersigned the sample referred to above **is of standard quality** as defined in the Drugs and Cosmetics Act, 1940, and Rules there under for the reasons given below:-

The sample conform to claim as per I.P.2022 in respect of test performed. Note: Other tests could not be perform due to want of Manufacturer's Specification.

Date: 04-APR-2025

**GOVERNMENT ANALYST** 

Rinku Kalita Government Analyst R.D.T.L., Guwahati-22

---- END OF REPORT ----