



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 : Mr. R. Vanlalruata, (Assistant Director (F and D))
Office of Chief Medical Officer, Aizwal, Mizoram-796017
- Serial No. and date of Inspector's memorandum : 23/AZL E/2026/Local , 25-FEB-2026
- Number of Sample : NIL
- Date of receipt : 11-MAR-2026
- Names of drugs purporting to be contained in the sample : Amoxycillin & Potassium Clavulanate Tablets I.P.
(NIXOX-CL 625)

Lab. Sample No.	Report No.	Batch No.	Date of Mfg. & Exp	Marketed By	Mfg. By
LSD/GUW/2026-27/0115	GUW/LS/2026-27/113	TB250708B	Nov-2025 Jul-2027	NA	M/s. Polestar Power Industries, (Pharma Division) Vill.Damuwala, Haripur Road, Barotiwalla, Baddi, Distt. Solan (H.P.) - 174 103

6. Condition of seals on : Seals were intact & identical to the specimen impression of the seal received from Drugs Inspector
[the packet or on portion of sample or container]

7. Result of test or analysis with protocols or test or analysis applied : Please see below
Date of Testing : From 12-May-2026 To 19-May-2026

COMPOSITION : Each film coated tablet contains:
Amoxycillin Trihydrate I.P. Eq. to Amoxycillin 500 mg.
Potassium Clavulanate Diluted I.P. Eq. to Clavulanic Acid 125 mg.

Protocol Applied : I.P. 2022

Sr No.	Test Name	Result	Limits
1	Description	White, elongated, bi-convex, film coated tablets, in blister pack.	NA
2	Identification	Gives positive test for Amoxycillin & Potassium Clavulanate.	NA
3	Average weight	1.08146 g	NA
4	Uniformity of weight	Complies.	+5 % of Average weight
5	Dissolution (for Amoxycillin)	Complies.	NLT 85 % of claim

6	Dissolution (for Clavulanic Acid))	Complies.	NLT 80 % of claim
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Assay

Sr.No	Ingredient Name	Found	Claim	% of claim	Limits	Procedure / Method
1	Amoxycillin Trihydrate equivalent to Amoxycillin Anhydrous	566.84 mg/Tablet	500 mg/Tablet	113.368	90 % to 120 %	I.P. 2022
2	Potassium Clavulanate Diluted eq. to Clavulanic Acid	137.44 mg/Tablet	125 mg/Tablet	109.952	90 % to 120 %	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.

Date: 29-MAY-2026

GOVERNMENT ANALYST

Rinku Kalita
Government Analyst
R.D.T.L., Guwahati-78

END OF REPORT

