



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 : Malsawmtluanga, (Assistant Drugs Inspector (F and D))
Lunglei- 796701
- Serial No. and date of Inspector's memorandum : 1, 31-MAR-2026
- Number of Sample : MS-12
- Date of receipt : 07-APR-2026
- Names of drugs purporting to be contained in the sample : Amoxycillin, Potassium Clavulanate and Lactic Acid
Bacillus Tablets (Pdlin-625 Plus Tablets)

Lab. Sample No.	Report No.	Batch No.	Date of Mfg. & Exp	Marketed By	Mfg. By
LSD/GUW/2026-27/0201	GUW/LS/2026-27/180	TB250682G	Nov-2025 Jul-2027	NA	M/s. Polestar Power Industries, (Pharma Division) Vill.Damuwala, Haripur Road, Barotiwala, Baddi, Distt. Solan (H.P.) - 174 103

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

7. Result of test or analysis with protocols or test or analysis applied : Please see below

Date of Testing : From 16-Jun-2026 To 18-Jun-2026

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

Protocol Applied : Manufacturer Specification

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 Trihydrate and Potassium Clavulanate.	NA
3	Average weight	1.0325 g	NA
4	Uniformity of weight	Complies.	±5% of avg. wt.
5	Disintegration	Complies. (Time taken for disintegration 22 minutes 15 seconds)	NMT 30 minutes

Assay

Sr.No	Ingredient Name	Found	Claim	% of claim	Limits	Procedure / Method
1	Amoxycillin Trihydrate eq. to Amoxycillin	521.29 mg/Tablet	500 mg/Tablet	104.258	90 % to 120 %	Manufacturer Specification
2	Potassium Clavulanate Diluted eq. to Clavulanic Acid	118.84 mg/Tablet	125 mg/Tablet	95.072	90 % to 120 %	Manufacturer Specification

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 Manufacturer Specification in respect of test performed.

Date: 01-JUL-2026

GOVERNMENT ANALYST

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

-----END OF REPORT-----

