



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 : B.LALZAMLIANA, Drugs Inspector (F and D)
CMO West Office
Kulikawn, Aizawl- 796005
- Serial No. and date of Inspector's memorandum : B.11012/5/2024-CMO (W)/DRUGS/47 , 27-FEB-2025
- Number of Sample : SMP/MZ/TMT/2024-2025/051
- Date of receipt : 04-MAR-2025
- Names of drugs purporting to be contained in the sample : Metformin Hydrochloride & Glimepiride Tablets
(Glimrica G1 Tablets)

Lab. Sample No.	Report No.	Batch No.	Date of Mfg. & Exp	Marketed By	Mfg. By
LSD/GUW/2025-GUW/LS/2025-26/0278	26/294	PT-40622	Feb-2024 Jan-2026	NA	Pharmaroots Healthcare, Khasra No. 411, Village Tipra, P.O. Barotiwala, Tehsil Baddi, Distt. Solan, H.P.

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 02-Apr-2025 To 20-May-2025

COMPOSITION : Each uncoated tablet contains:
Glimepiride IP 1 mg
Metformin Hydrochloride IP 500 mg

Protocol Applied : Manufacturer's Specification

Sr No.	Test Name	Result	Limits
1	Description	White, elongated, bi-convex, having scored on one surface, uncoated tablets, in blister pack.	NA
2	Identification	Gives positive test for Metformin Hydrochloride & Glimepiride.	NA
3	Average weight	1.0278 g	NA
4	Uniformity of weight	Complies.	+5 % of Average weight
5	Disintegration	Complies. (Time taken for disintegration 07 minutes 28 seconds)	NMT 15 minutes

Assay

Sr.No	Ingredient Name	Found	Claim	% of claim	Limits	Procedure / Method

1	Metformin Hydrochloride	479.18 mg/Tablet	500 mg/Tablet	95.836	90 % to 110 %	Manufacturer's Specification
2	Glimepiride	0.98 mg/Tablet	1 mg/Tablet	98	90 % to 110 %	Manufacturer's 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's Specification in respect of test performed.

Date: 27-MAY-2025

Rinku Kalita
GOVERNMENT ANALYST

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

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

