

FORM 13  
 (See rule 46)

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

1. Name of Inspector from whom received : MR. R. VANLALRUATA  
 ASSISTANT DIRECTOR (F & D)  
 OFFICE OF CHIEF MEDICAL OFFICER,  
 AIZAWL, MIZORAM, PIN: - 796017
2. Serial No. and date of Inspector's memorandum : MZ/24/A 'E'/13/IPA Date 17.12. 2024
3. Number of Sample : Nil
4. Date of receipt : 02-JAN-2025
5. Names of drugs purporting to be contained in the sample: Pantoprazole Gastro-Resistant Tablets IP 40 mg  
 (Pantojack-40)

Lab. Sample No.	Report No.	Batch No.	Date of Mfg.	Date of Exp.	Mfg. By
LSD/GUW/2024-25/1840	GUW/LS/2024-25/1863	T-1261	APR. 2024	MAR. 2026	Jackson Laboratories Pvt. Ltd. 22-24, Majitha Road, Bye Pass Amritsar-143 001(India)

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 06-Mar-2025 To 10-Mar-2025

COMPOSITION : Each enteric coated tablet contains:  
 Pantoprazole Sodium  
 Sesquihydrate IP  
 eqv. to Pantoprazole 40 mg

**Protocol Applied: I.P. 2022**

Sr No.	Test Name	Result	Limits
1	Description	Yellow colour, round, enteric coated tablet, showing, cross sectional (capping defect) fragment detaches supplied in blister pack.	NA
2	Identification	Gives positive test for Pantoprazole.	NA
3	Average weight	0.1707 gm.	NA
4	Dissolution for Acid stage	<b>Does not comply.</b> (All the six units were found to be dissolved in Acid medium and no residue remained in the vessel). Hence, further test could not be carried out.	NMT 10% of claim

**Assay**

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

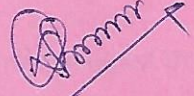


1	Pantoprazole Sodium Sesquihydrate eqv. to Pantoprazole	42.69 mg/Tablet	40 mg/Tablet	106.725	90 % to 110 %	I.P. 2022
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In the opinion of the under signed the sample referred to above **is not of standard quality** as defined in the Drugs and Cosmetics Act, 1940, and Rules thereunder for the reasons given below:-

**The sample does not conform to I.P. with respect to the tests for 'Dissolution' (Acid stage) and is also not of standard quality for the reason stated under 'Description'**

Date: 17-MAR-2025



**GOVERNMENT ANALYST**  
**Dilip Kr. Sarkar**  
Government Analyst  
R.D.T.L., Guwahati-22

Report in red coloured paper indicates the sample is Not of Standard Quality

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

