

Regional Drugs Testing Laboratory Directorate General of Health Services, Guwahati (India)-781022 Fax: 0361-2338555/2330555 Phone No.0361 2338555

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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

1. Name of Inspector from whom received

: SARAH LALDINTLUANGI

ASSISTANT DIRECTOR (FOOD AND DRUGS)

Sr. CMO OFFICE, SERCHHIP-796181,

SERCHHIP

2. Serial No. and date of Inspector's memorandum

:019/IPA/2024

Date 13.12.20124

3. Number of Sample

:Nil

4. Date of receipt

:02-JAN-2025

5. Names of drugs purporting to be contained in the sample: CIPROFLOXACIN TABLETS IP 250 mg (Ciprodac 250)

Lab. Sample No.	Report No.	Batch No.	Date of Mfg.	Date of Exp.	Mfg. By
LSD/GUW/2024- 25/1607	GUW/LS/2024- 25/1642	CT22008	AUG. 2022	JUL. 2025	CADILA PHARMACEUTICALS LIMITED,1389, Dholka- 382 225, Dist.: Ahmedabad

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

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

Date of Testing: From 20-Jan-2025To 27-Jan-2025

COMPOSITION

Each film-coated tablet contains: Ciprofloxacin Hydrochloride IP equivalent to Ciprofloxacin 250 mg

Sr No.	Test Name	Result	Limits	
1	Description	White, round.bi-convex, film coated tablet, having a dosage line on one side packed in blister pack.	NA	
2	Identification	Gives positive test for Ciprofloxacin.	NA	
3	Average weight	0.3324 gm.	NA	
4	Uniformity of weight	Complies.	NA	
5	Dissolution	Complies {Average drugs release (S ₁ +S ₂₎ found: 84.07%}.	Average of 12 units (S_1+S_2) is $\geq 80\%$ ($\geq Q$) and no unit is $< 65\%$ of claim (Q-15%).	

<u>Assay</u>

Sr.No	Ingredient Name	Found	Claim	% of claim	Limits	Procedure / Method
		- 5 ™	a			

	Hydrochloride	250.87 mg/Tablet	250 mg/Tablet	100.348	90 % to 110 %	I.P. 2022
ioni .	equivalent to Ciprofloxacin	Sedantic	reserve to			

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

The sample conforms to I.P. with respect to the above tests only.

Date: 03-FEB-2025

GOVERNMENT ANALYST

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

- END OF REPORT -