Regional Drugs Testing Laboratory
Directorate General of Health Services, Guwahati (India) - 781022
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Regional Drugs Testing Laboratory Govt. of India Guwahati-22

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** : Mrs. C. Ramdinmawii,Assistant Director (F and D) C.M.O. Office, Champhai-796321 Mizoram
2. **Serial No. and date of Inspector's memorandum** : 020/IPA-CPI/24 , 16-DEC-2024
3. **Number of Sample** : NIL
4. **Date of receipt** : 31-DEC-2024
5. **Names of drugs purporting to be contained in the sample** : Ferrous Sulphate and Folic Acid Tablets I.P.

Lab. Sample No.	Report No.	Batch No.	Date of Mfg.	Date of Exp.	Mfg. By
LSD/GUW/2024-25/1856	GUW/LS/2024-25/1876	CFR-W27	Mar-2024	Feb-2026	Cyano Pharma (P) Ltd., 115-C, Industrial Estate, Pologround, Indore-452 015

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 05-Feb-2025 To 13-Mar-2025

COMPOSITION : Each sugar coated tablet contains:Dried Ferrous Sulphate IP 200 mg. (Eq. to Ferrous Elemental Iron 60 mg.)Folic Acid I.P. 0.5 mg.

Protocol Applied : I.P. 2022

Sr No.	Test Name	Result	Limits
1	Description	Red colored, round, bi-convex, coated tablets, in blister pack.	NA
2	Identification	Gives positive test for Dried Ferrous Sulphate & Folic Acid.	NA
3	Average weight	0.3612 g	NA
4	Disintegration	Complies. (Time taken for disintegration 38 minutes and 19 seconds)	NMT 60 minutes
5	Uniformity of content	Complies for Folic Acid.	NA

Assay

Sr.No	Ingredient Name	Found	Claim	% of claim	Limits	Procedure / Method
1	Folic acid	1.0007 mg/Tablet	0.5 mg/Tablet	200.14	NLT 90 %	I.P. 2022

2	Dried Ferrous Sulphate equivalent to Elemental Iron	59.7 mg/Tablet	60 mg/Tablet	99.5	90 % to 110 %	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: 18-MAR-2025

GOVERNMENT ANALYST

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

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