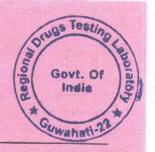


Regional Drugs Testing Laboratory Directorate General of Health Services, Guwahati (India) - 781022 Fax: 0361-2338555/2330555 Phone No.0361 2338555 Email: rdtlguwahati@cdsco.nic.in



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 (Food & Drugs) C.M.O. Office, Champhai-796321

Mizoram.

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

: 366/24

Date: 24/07/2024

3. Number of Sample

: Nil

4. Date of receipt

: 12-AUG-2024

5. Names of drugs purporting to be contained in the sample : Acyclovir Dispersible Tablets IP (Zovibact 800 mg)

Lab. Sample No.	Report No.	Batch No.	Date of Mfg.	Date of Exp.	Mfg. By
LSD/GUW/202 4-25/1290	GUW/LS/2024 -25/1313	SRTH230874B	08/2023	07/2025	SYSTOLE REMEDIES PVT. LTD. Vill. Ogli, Kala Amb, Teh. Nahan, Distt. Sirmour (H.P.) -173 030

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 30-Oct-2024 To 28-Nov-2024

COMPOSITION

Each uncoated dispersible tablet contains:

Acyclovir IP 800 mg

Protocol Applied: I.P. 2022

Sr No.	Test Name	Result	Limits
1	Description	White, elongated, bi-convex, uncoated dispersible tablet, plain on both side packed in blister pack.	NA
2	Identification	Gives positive test for Acyclovir.	NA
3	Average weight	1.241 gm.	NA
4	Uniformity of weight	Complies.	NA
5	Uniformity of dispersion	Does not comply	NA
6	Disintegration	Does not comply. (All six tablets did not disintegrate with & without disc till 3 minutes at 25°C).	NMT 3 minutes

Assay

Sr.No	Ingredient Name	Found	Claim	% of claim	Limits	Procedure / Method
1	Acyclovir	784.54 mg/Tablet	800 mg/Tablet	98.067	95 % to 105 %	I.P. 2022

In the opinion of the undersigned the sample referred to above is not of standard quality as defined in the Drugs and Cosmetics Act, 1940, and Rules there under for the reasons given below:-

The sample does not conform to I.P. with respect to the test for 'Uniformity of dispersion' & 'Disintegration'.

Date: 03-DEC-2024

GOVERNMENT ANALYST

Dilip Kr. Sarkar

Government Analyst

R.D.T.L., Guwahati-22

1

Report in red coloured paper indicates the sample is Not of Standar	d Quality
END OF REPORT	
STANDARD CONTAG	