HEALTH & FAMILY WELFARE DEPARTMENT
HIMACHAL PRADESH

Certificate of Good Manufacturing Practices

This one page certificate conforms to the format recommended by the World Health Organization [General Instructions and Explanatory Notes attached].

Certificate No. HFW-H(Drugs)47/08

On the basis of the inspection carried out on 25th & 26th March 2021, we certify that the site indicated on this certificate complies with Good Manufacturing Practices for the dosage forms, categories and activities listed in Table I:

1. Names and Address of Site: **M/S Sarv Biolabs Pvt. Ltd., Plot No. 2, Trilokpur Road, Behind IITT College, Kala-Amb, Distt. Sirmaur, Himachal Pradesh, (India) 173030.**

2. Manufacturer's License No: **S-MNB/08/07 on form 25 Valid upto 16.06.2023**

3. Table-I:

Dosage Form[s]	Category[ies]	Activity[ies]
Active Pharmaceutical Ingredients **1.** Thiocolchicoside IP, FP **2.** Thiocolchicoside EP(Crystallized from Ethanol) **3.** Colchicine USP, EP, BP **4.** Thiocolchicoside Hydrate EP	**General**	**Production, Packing & Quality Control**

The responsibility for the quality of the individual batches of the pharmaceutical products manufactured through this process lies with the manufacturer.

This certificate remains valid until **16.06.2023**. It becomes invalid if the activities and/or categories certified herewith are changed or if the site is no longer considered to be in compliance with GMP.

Address of Certifying Authority: **State Drugs Controller**,
Controlling cum Licensing Authority,
2nd floor, HIMUDA Commercial Complex, Phase-I,
Housing Board, Baddi, Distt. Solan [H.P.] 173205, INDIA.

Name & Function of Responsible person: **Navneet Marwaha**
State Drugs Controller
Controlling cum Licensing Authority
01795-244288, sdc4hp@gmail.com

Telephone/Fax No:
Date: 16.06.2021

Signature:
Stamp: Licensing Authority cum Drug Controller * Himachal Pradesh *

(NAVNEET MARWAHA) 16.6.21
State Drugs Controller
Controlling cum Licensing Authority
Baddi Distt.Solan (H.P.)-173205
01795-244288,sdc4hp@gmail.com

Explanatory Notes:

1 This certificate, which is in the format recommended by WHO certifies the status of the site, listed in point I of the certificate.

2 The certificate number should be traceable within the regulatory authority issuing the certificate.

3 Where the Regulatory Authority issues a license for the Site, this number should be specified. Record 'Not Applicable" in cases where there is no legal framework for the issuing of a license.

4 Table I

List the Dosage Forms, starting materials, categories and activities. Examples are given below:

Example 1

Pharmaceutical Product[s]1	Category [ies]	Activity [ies]
Dosage Form [s]:		
Tablets	Cytotoxic	Packaging
	Hormone	Production, Packing, Quality Control
	Penicillin	Repackaging and Labeling
Injectables	Cephalosporin	Aseptic preparation, Packaging, Labeling

Example 2

Pharmaceutical Product[s]1	Category [ies]	Activity [ies]
Starting Material [s]		
Paracetamol	Analgesic	Synthesis, Purification, packing, Labeling

Use, whenever available, International Non proprietary Names [Inns] or otherwise national Non proprietary Names

5 The certificate remains valid until the specified date. The certificate becomes invalid if the activities and/or categories certified are changed or if the site is no longer considered to be in compliance with GMP.

6 The requirements for good practices, the manufacture and quality control of drugs referred to in the certificate are those included in Quality Assurance of Pharmaceuticals: a compendium of guidelines and related materials. Good Manufacturing Practices and Inspection. Volume 2, 1999 World Health Organization. Geneva and subsequent updates.