GOVERNMENT OF INDIA
Directorate General of Health Services
Ministry of Health & Family Welfare
Central Drugs Standard Control Organization

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)[1].

Certificate No: WHO-GMP/Divya-II/Haridwar/02/2019

On the basis of the inspection carried out on 28th&29th Mar.2019 we certify that the site indicated on this certificate complies with Good Manufacturing Practices for the dosage forms, categories *and activities listed in Table 1.*

1. **Name and Address of Site:-** **M/s Divya Pharmacy (Unit-II)**
Khasra No.-210-211, Patanjali Food & Herbal Park,
Padartha, Laksar Road, Haridwar-249404, Uttarakhand

2. **Manufacturer's License Number: -** UK.AY-274/2013 Issued date: 13/12/13

3. **Table: 1**

Dosage form (S)	Category (ies)	Activity (ies)
Oral Powder (Churna)	Gastric Disorder, indigestion, Appetite stimulants, Laxatives, Antacid, Constipation etc	Production Quality Control, Labeling & Packing
Tablets	Hemorrhoids, Appetite stimulants, Gastric Disorder, varna etc.	

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 01 SEP 2022. 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:
Central Drugs Standard Control Organization, FDA Bhawan, ITO
Kotla Road, New Delhi- 110002

Name and Function of responsible Person:
Dr. V. G. Somani
Drugs Controller General (I),
Email: dci@nic.in
Telephone No: - 911123236965
Fax No: - 911123236973

Signature:

Dr. V. G. SOMANI
Drugs Controller General (India)
Dte. General of Health Services
Ministry of Health and Family Welfare
FDA Bhawan, Kotla Road, I.T.O.
New Delhi-110002

Stamp & Date: 02 SEP 2019

[1](This model certificate for GMP is not part of the WHO certification scheme on the Quality of Pharmaceutical Products Moving in International Commerce.)

Explanatory Notes

1. This certificate, which is in the format recommended by WHO, certifies the status of the Site listed in Point 1 of the certificate.
2. The certification 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 case where there is no legal framework for the issuing of a license.
4. Table 1
   List of dosage forms, starting materials, categories and activities.

Example 1

Pharmaceutical Product (s) 2	**Category (ies)**	**Activity (ies)**
Dosage Form (s)[2]		
Tablets	Cytotoxic	Production
	Hormone	Production ,Quality Control & Packing
Injectables	Penicillin	Repacking and Labelling
	Cefalosporin	Aseptic Preparation , Packing, Labelling

Example 2

Pharmaceutical Product (s) [2]	**Category (ies)**	**Activity (ies)**
Starting Material (s): [3]		
Paracetamol	Analgesic	Synthesis ,Purification, Packing , Labelling

[2]Pharmaceutical Products :- Any medicine intended for human use or veterinary product administered to food – Producing animals, presented in its finished dosage for or as a starting material for use in such a dosage form, that is subject to control by pharmaceutical legislation in both the exporting state and the importing state.
[3]Starting Materials: Any substance of a defined quality used in the production of pharmaceutical product, but excluding packaging materials.
Use, whenever available, International nonproprietary Names (INNs) or otherwise national nonproprietary 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 in 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.