

**Health & Family Welfare Department  
Himachal Pradesh  
Baddi, Dist. Solan  
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) 56/98

On the basis of the inspection carried out on 19<sup>th</sup> & 20<sup>th</sup> Dec. 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 I:

1. Names and Address of Site: M/s Morepen Laboratories Ltd. Unit-IV,  
(Morepen Village), Malkumajra,  
Baddi-Nalagarh Road,  
Baddi, Distt. Solan (H.P.) INDIA
2. Manufacturer's License No: NB/98/05  
B/98/06 valid upto 31.12.2021

3. Table-I:

Pharmaceutical Product [s]	Category[ies]	Activity[ies]
Starting Material [s]		Synthesis, Purification, Packing, Labeling
Rosuvastatin Calcium IP/EP/IH/BP/USP	Antilipemic	Synthesis, Purification, Packing, Labeling
Atorvastatin Calcium USP/IP/JP/IH	Antihyperlipidemic	Synthesis, Purification, Packing, Labeling
Fexofenadine Hydrochloride USP/EP/BP/JP/IP	Antihistaminic	Synthesis, Purification, Packing, Labeling
Loratadine USP/EP/BP/IP	Antihistaminic	Synthesis, Purification, Packing, Labeling
Olmesartan Medoxomil USP/EP/JP/IH/IP	Antihypertensive	Synthesis, Purification, Packing, Labeling
Sitagliptin Phosphate USP/EP/IP/IH	Antidiabetic	Synthesis, Purification, Packing, Labeling
Montelukast Sodium USP/EP/BP/IP	Antiasthmatic	Synthesis, Purification, Packing, Labeling
Atorvastatin Calcium Amorphous	Antihyperlipidemic	Synthesis, Purification, Packing, Labeling
Atorvastatin Calcium Trihydrate EP	Antihyperlipidemic	Synthesis, Purification, Packing, Labeling
Saxagliptin Hydrochloride IH	Antidiabetic	Synthesis, Purification, Packing, Labeling
Empagliflozin IH	Antidiabetic	Synthesis, Purification, Packing, Labeling
Linagliptin IH	Antidiabetic	Synthesis, Purification, Packing, Labeling
Dapagliflozin Propanediol Monohydrate IH	Antidiabetic	Synthesis, Purification, Packing, Labeling
Candesartan Cilexetil IH/USP/BP/EP	Antihypertensive	Synthesis, Purification, Packing, Labeling

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

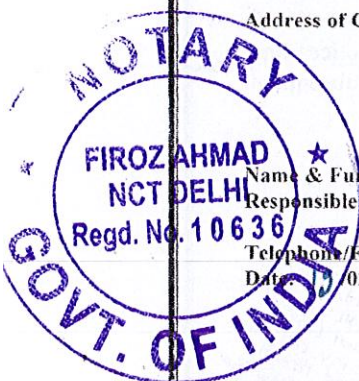
This certificate remains valid until 31-12-2021. It becomes invalid if the activities and/or categories certified herewith are changed or if the site is no longer considered to be in compliance with GMP.

Address of Certifying Authority:

Dy. Drugs Controller,  
Cum Licensing Authority,  
O/o State Drugs Controller,  
Baddi, Distt. Solan, H.P.173205  
01795-244288, sdc4hp@gmail.com

Manish Kapoor  
Dy. Drugs Controller,  
Cum- Licensing Authority,  
01795-244288

Signature:  
Stamp:



Manish Kapoor  
DEPUTY DRUGS CONTROLLER  
-cum- LICENSING AUTHORITY  
O/O STATE DRUGS CONTROLLER  
BADDI, DISTRICT SOLAN, H.P.-173205  
E-mail: sdc4hp@gmail.com  
Phone: 01795-244288

**ATTESTED**

**NOTARY PUBLIC  
DELHI (INDIA)**

**GAURAV RAVISH  
Addl. Director**

**Authorized Signatory  
Millennial India International  
Chamber of Commerce  
Industry & Agriculture  
New Delhi, (INDIA)**

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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 A
4. Applicable" in cases where there is no legal framework for the issuing of a license.
5. Table I

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

Example 1

Pharmaceutical Product[s]	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]	Category [ies]	Activity [ies]
Starting Material [s]		
Paracetamol	Analgesic	S...

Use, whenever available, International Non proprietary Names

6. The certificate the activities considered to

7. The requirements referred to in the a compendium of Inspection. Volume updates.



(सुनील चनाप)  
(SUNIL CHANAP)  
अनुभाग अधिकारी (आ.अ.)  
Section Officer (OI)  
सी. पी. वे. विभाग / C.P.V. Division  
निदेशक, नई दिल्ली  
Ministry of External Affairs, New Delhi