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 19th & 20th 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]1	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	Antiasthmatics	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 Cilexetin 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

Name & Function of Responsible person:

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

Telephone/Fax No: 01795-244288

Date: 19 /02/2020

DEPUTY DRUGS CONTROLLER-cum-LICENSING AUTHORITY * H.P. *

19 FEB 2020

Signature:

Stamp:

(MANISH KAPOOR)
DEPUTY DRUGS CONTROLLER
-cum-LICENSING AUTHORITY
O/o STATE DRUGS CONTROLLER
BADDI, DISTRICT SOLAN, H.P.-173205
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