

**Office of the Drug Controller (AYUSH) Madhya Pradesh**

(Ground Floor "D" Wing Satpura Bhawan, Bhopal - 462004)

Phone - 0755 - 2763044

No. Drug/20/ 1840-41

Bhopal, Dated: 08/09/2020

To,


M/s. Dindayal Industries Ltd  
62 Malanpur Ind. Area Bhind (M.P.)

Sub :- Renewal of licence in form 26D.

Please find enclosed herewith certificate of renewal of licence No:-  
MP25D/15/392 dated 16/09/2015 to manufacture drugs as per approved list  
for the period 16/09/2020 to 15/09/2025

Any time, it is found that you are not complying schedule "T"  
norms then the licence will be suspended

Encl : Certificate of renewal of licence

  
**LICENSING AUTHORITY**  
**Office of the Drug Controller (AYUSH)**  
**MADHYA PRADESH**

Endt. No. Drug/20/

Bhopal, Dated:

Copy to :-

District AYUSH Officer, Distt.- Bhind M.P. for  
information.

  
**LICENSING AUTHORITY**  
**Office of the Drug Controller (AYUSH)**  
**MADHYA PRADESH**

[Form 26-D, under Rule 155]

***Certificate of renewal of licence to manufacture for sale of  
Ayurvedic / Siddha or Unani Drugs.***


1. Certified that licence No. MP25D/15/392 granted on the 16/09/2015 to Managing Director Shri Anand Mohan Chhapparwal of M/s. Dindayal Industries Ltd., for the manufacture of Ayurvedic Drugs at the premises situated at 62 Malanpur Ind. Area Bhind M.P. has been renewed from 16/09/2020 to 15/09/2025
2. Names of technical staff  
Manufacturing Side -  
1. V.R. Tamhankar  
Quality Control Section -  
1. Dr L.K. Goswami  
Other Testing Laboratories :- Quality Control Lab Bhopal (M.P.)
3. Names of Drugs  
As per approved previous list
4. Sufficient informations on the lable of medicine's packing must be printed as per the provisions of the Drugs and Cosmetics Rules 1945.
5. The record of quality of control test of raw drugs manufacturing process tests for standards during manufacturing process and tests of the finished product must be kept in record as per the provisions of schedule 'T' to comply with the provisions of GMP under rule 157 of Drugs and Cosmetic Rules 1945.
6. Provide Testing Protocol and Safety studies data of patent or proprietary Products before sell in market otherwise licence should be cancelled



AMC/11/122

QC 1339

Bhopal  
Date

  
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**Office of the Drug Controller (AYUSH)**  
**MADHYA PRADESH**