

FORM 26 E-1

(Refer Rule 155B)

CERTIFICATE OF GOOD MANUFACTURING PRACTICES (GMP) TO MANUFACTURER OF AYURVEDA, SIDDHA OR UNANI DRUGS

Certified that manufacturing unit Licensee, namely :

M/s. NATIONAL INSTITUTE OF AYURVED

Situated at : **JORAWAR SINGH GATE, AMER ROAD,
JAIPUR (RAJ.) 302002**

State : **RAJASTHAN** License No. **RJ 776-D.**

comply with the requirements of Good Manufacturing Practices (GMP) of Ayurveda-Siddha-Unani drugs as laid down in Schedule T of the Drugs and Cosmetics Rules, 1945.

This certificate is valid for a period of Five years

FROM 17.05.2025 TO 16.05.2030

and the Good Manufacturing Practices (GMP) is valid for the various dosage forms or Rasaushadhis, as follows :-

ANJANA/ PISTI, CHURNA/ NASYA/ MANJAN/ LEPA/ KWATH, PILLS/ VATI/ GUTIKA/ MATRICA/ TABLETS, KUPIPAKVA/ KSARA/ PARPATI/ LAVANA/ BHASMA/ SATVA/ SINDURA/ KARPUR/ UPPU/ PARAM, CAPSULES, PAK/ AVALEH/ KHAND/ MODAK/ LAKAYAM, PANAK/ SYRUP/ PRAVAHI KWATH/ MANAPAKU, ARK/ TINIR, TAIL/ GHRIT/ NEY, SAUNDARYA PRASADAK (SKIN & HAIR CARE).

Place : **AJMER**



(DR. ANAND KUMAR SHARMA)
DIRECTOR & LICENSING AUTHORITY
AYURVED DEPARTMENT RAJASTHAN AJMER

Document certified by ANAND KUMAR SHARMA <sharmavd.anand@yahoo.in>.

Digitally Signed by ANAND KUMAR SHARMA
Designation: Director
Date :20-05-2025 04:22:49



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