GOVERNMENT OF TELANGANA

DEPARTMENT OF AYUSH

CERTIFICATE OF GOOD MANUFACTURING PRACTICES

FORM 26 E-1

Rule 155 - B

Certified that manufacturing unit licensee, namely M/s GREEN CARE BIO SCIENCES

situated at ,QUTHBULLAPUR MEDCHAL - MALKAJGIRI,State Telangana Licence No:**T-2323/Ayurv** Comply with the requirements of Good Manufacturing Practices of AYURVEDA as laid in Schedule 'T' of the Drugs and Cosmetic Rules 1945.

This Certificate is Valid up to 26/04/2025

Date of Issue:27/04/2020
Place**:** Hyderabad

Signature:
K ANASUYA
Additional Director (AYURVEDA)
Drug Licensing Authority
Department of AYUSH
Hyderabad, Telangana State
Digitally Signed by
Date:27-04-2020 18:50:46 PM

CONDITIONS OF LICENCE

- This license and any certificate of renewal in force shall be kept on the approved premises and shall be produced on the request of an Inspector appointed under the Drugs and Cosmetics Act,.
- Any change in the technical staff named in the license shall be forthwith reported to the Licensing Authority.
- This license shall be deemed to extend to such additional items as the Licensee may intimate to the Licensing Authority from time and as may be endorsed by the Licensing Authority.
- The Licensee Shall i nform the Licensing Authority in writing the event of any change in the constitution of the firm operating under the license, where any change in the constitution of the takes palce, the current license shall be deemed to be valid for maximum period of three months from the date on which the change taken place unless in the meantime a fresh license has been taken from the licensing authority in the name of the firm with the changed constitution
- License for preparing eye drops in Ayurvedic System should fulfill the conditions laid in Schedule F.
- The License shall test each batch of the final product containing alcohol for the tests prescrided for each preparation and shall maintain records showing the particulars in respect of such tests, the recors shall be maintained for a period of three years from the date of manufacture.
- The records shall be maintained of Ayurvedic medicindes containing alcohol mentioning the qantities sold, batch numer, names and address of parties to whom sold.
- The licensee shall keep records of the details of each batch of drugs manufactured by him and of raw materials used therein asper particulars Schedule-II (I) and such rcords shall be retained for a period of (3) years.
- The Licensee shall allow in inspector appointed under the act to enter with or without prior notice, any premises where manufacture of drugs, in respect of which licensee is issued is carried on the inspect the premises and to take samples of the raw materials as well as the finished products and the manufactured products unser a recipt and to inspect the records maintained un der these rules.
- The Licensee shall make arrangement for proper storage of drugs manufactured by him.
- The Licensee shall maintain an Inspection Book in form to enable in inspector to record him impressions and the defects noticed.
- The IIcensing authority shall obtain the opinion of the technical experts appointed under the Rules A, regarding the suitability of the drug manufactured by the licensee or the applicant in case of any doubts. The opinion shall be the final if it is not disapproved by the applicant or the licensee within (30) days of the renewed of the opinion.
- This license is granted subject to the conditions that all licensed products will be manufactured and marked strictly following of "Good Manufacturing Practices" as applicable. Any deviations/violations will entail cancellation of License forthwith.