WHO WE ARE:

Eva Grow Medicaps Private Limited is one of the fast-growing Empty Hard Gelatin Capsules Manufacturing Company which is located at one of the Pharmaceutical Manufacturing Hub of India. Eva Grow Medicaps Private Limited is managed under the guidance of experience pharmaceuticals leaders who coach the facility for best compliance of its products as per the requirement of Global Regulatory Market.

The firm was incorporated in the year of 2021 for the supply of Empty Hard Gelatin Capsule Shell and the current yearly capacity is 8.5 Billion Capsules.

Total Buildup Area: 100000 sq-ft, **Green Area:** 40% of buildup area.

evaGROW

EVA GROW MEDICAPS
PRIVATE LIMITED

Eva Grow Medicaps Private Limited

Sai Road, Baddi, HP (India)-173205

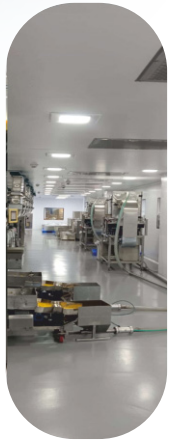
Ph. No.: +91 7903259155, 8091555017, 8091555343

Email: marketing@evagrow.in | gm@evagrow.in

Website: www.evacaps.in

KEY MANUFACTURING EQUIPMENTS:

DBDS HIGH SPEED AUTOMATIC MACHINE



LINE LAYOUT FOR SINGLE PIECE

GMP COMPLIANCE

Manufacturing Block is ISO Class 8 clean room facility which equipped with the online automatic capsule manufacturing and inspection machines. This allowed single piece flow lay-out to control the product-to-product contamination, our HVAC system is fully automatic and External environment is free from bio loads.

INHOUSE WATER PURIFICATION SYSTEM:

WATER PURIFICATION SYSTEM

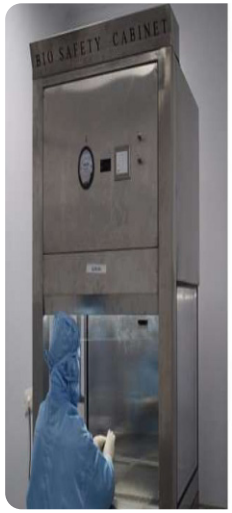
ONLINE OZONE GENERATOR SYSTEM

Water purification is conforming to the cGMP norms for Ozone Generation, RO, DM and Heating mechanism at 70°C which further eliminates the microbial contamination.

KEY LABORATORY

STABILITY STUDY CHAMBERS

We are performing inhouse stability study of our all-Size specific products as well as all key colors. Our product is complying to all zone's conditions required for the stability as per GMP guideline. We ensure the best performance of our Empty Hard Gelatin Capsules Product during filling machine operations.



MLT INCUBATION CHAMBER

CHEMICAL AND INSTRUMENTATION LAB

We have GLP certified Laboratory facility including Chemical, Instrumentation Microbial and Physical capsules parameter measurement.

PRODUCT RANGE: CAPSULE SIZE: #, 0, 1, 2, 3, 4

CUSTOMIZATION ON PRODUCTS:

- Easy fit Capsules.
- Delay Release Capsules.
- Flavored Capsules.
- Printed Capsules.
- Semi-Liquid Fill Capsules.
- Preservative Free Capsules.
- DPI Capsules and Press Fit Capsules.
- Pearl Metallic Capsule.
- SiO2 Free Capsule
- TiO2 Free Capsules.
- SLS Free Capsules.

CERTIFICATIONS:

- QMS - ISO 9001:2015
- Laboratory - GLP
- Manufacturing - GMP & WHO GMP
- Product - EDQM RM, FSSAI and Halal Certified BSE/TSE Free , Bovine Bone 210 Bloom Pharma Grade Gelatin

QAA

Certificate Of Compliance

This is to Certify that the Quality Management System of the

EVA GROW MEDICAPS PRIVATE LIMITED

ISO 9001:2015

"MANUFACTURER OF EMPTY HARD GELATINE CAPSULE SHELL"

ISO:9001-2015

HEALTH & FAMILY WELFARE DEPARTMENT
HIMACHAL PRADESH

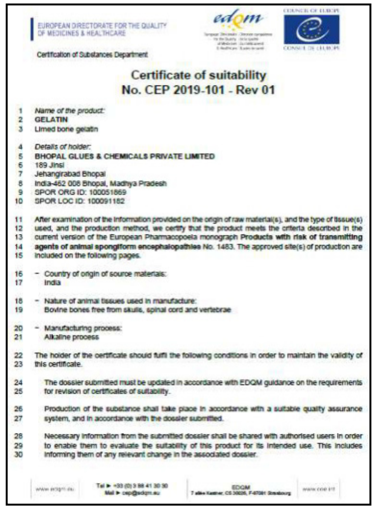
Certificate of Good Manufacturing Practices

WHO-GMP

No.: HFW-H (Drugs) 101/22
HEALTH & FAMILY WELFARE DEPARTMENT
HIMACHAL PRADESH

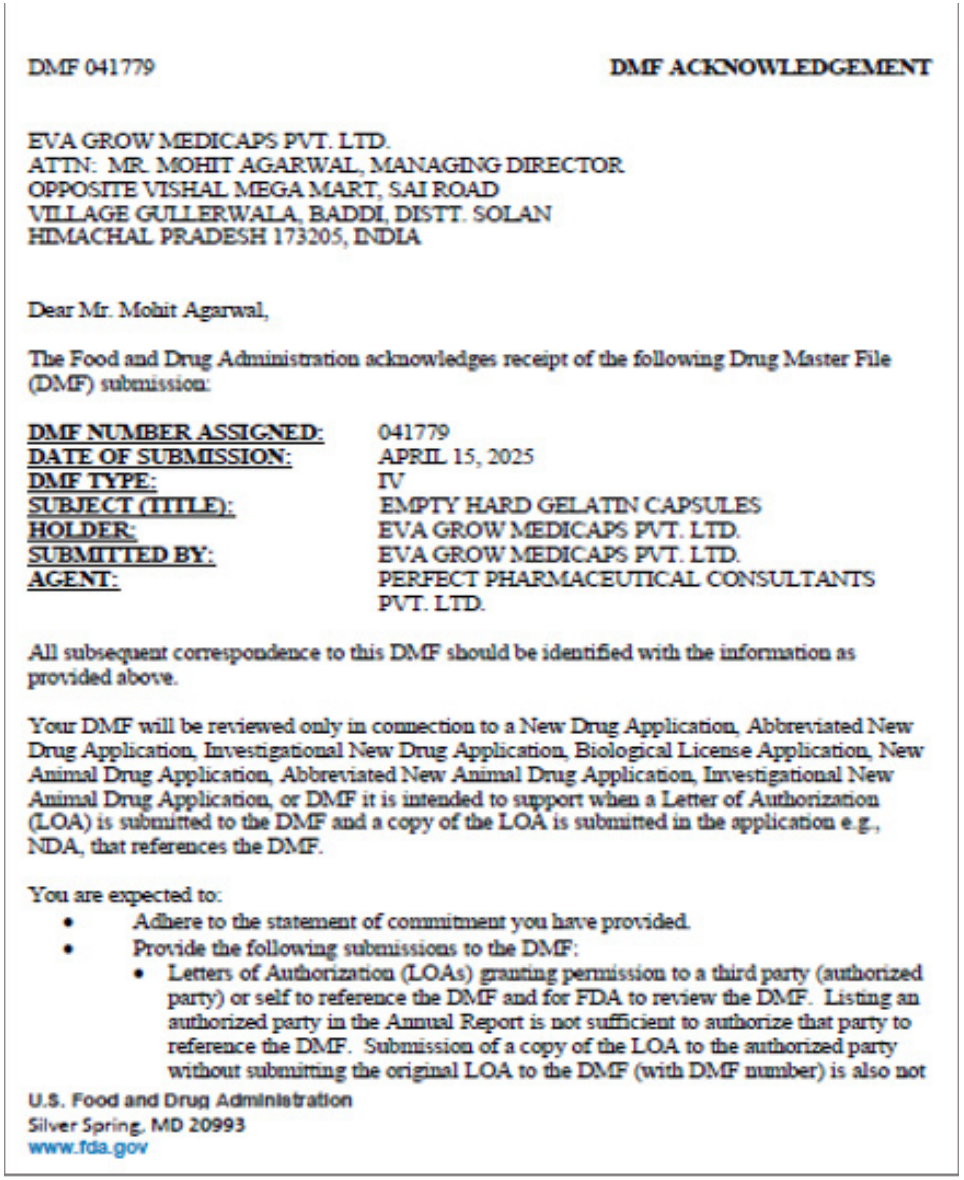
G. L. P. CERTIFICATE

GLP



Certificate of suitability
No. CEP 2019-101 - Rev 01

EDQM OF RAW MATERIAL



DMF 041779 **DMF ACKNOWLEDGEMENT**

EVA GROW MEDICAPS PVT. LTD.
ATTN: MR. MOHIT AGARWAL, MANAGING DIRECTOR
OPPOSITE VISHAL MEGA MART, SAI ROAD
VILLAGE GULLERWALA, BADDI, DISTT. SOLAN
HIMACHAL PRADESH 173205, INDIA

Dear Mr. Mohit Agarwal,

The Food and Drug Administration acknowledges receipt of the following Drug Master File (DMF) submission:

DMF NUMBER ASSIGNED:	041779
DATE OF SUBMISSION:	APRIL 15, 2025
DMF TYPE:	IV
SUBJECT (TITLE):	EMPTY HARD GELATIN CAPSULES
HOLDER:	EVA GROW MEDICAPS PVT. LTD.
SUBMITTED BY:	EVA GROW MEDICAPS PVT. LTD.
AGENT:	PERFECT PHARMACEUTICAL CONSULTANTS PVT. LTD.

All subsequent correspondence to this DMF should be identified with the information as provided above.

Your DMF will be reviewed only in connection to a New Drug Application, Abbreviated New Drug Application, Investigational New Drug Application, Biological License Application, New Animal Drug Application, Abbreviated New Animal Drug Application, Investigational New Animal Drug Application, or DMF it is intended to support when a Letter of Authorization (LOA) is submitted to the DMF and a copy of the LOA is submitted in the application e.g., NDA, that references the DMF.

You are expected to:

- Adhere to the statement of commitment you have provided.
- Provide the following submissions to the DMF:
  - Letters of Authorization (LOAs) granting permission to a third party (authorized party) or self to reference the DMF and for FDA to review the DMF. Listing an authorized party in the Annual Report is not sufficient to authorize that party to reference the DMF. Submission of a copy of the LOA to the authorized party without submitting the original LOA to the DMF (with DMF number) is also not

U.S. Food and Drug Administration
Silver Spring, MD 20993
www.fda.gov

PACKAGING STANDARD:

DOMESTIC PACKAGING MATERIAL:
Primary Packing - Anti-static Poly Bag. 1st
Secondary Packing – Thermocol sheet/box.
2nd Secondary Packing – 5 Ply Corrugated Box.

EXPORT:
Primary Packing – Food Grade Metallic Bag,
Anti-static Poly Bag,
1st Secondary Packing – Thermocol sheet/box.
2nd Secondary Packing – 5 Ply Corrugated box.
3rd Secondary Packing – Shrink film/2PP Straps.

SEALING:
Primary Packing sealing by heat and secondary by BOPP Tape outside on flaps.

OTHERS:
As per customer specific requirement.

SIZE	QUANTITY PER BOX
0	100000
1	125000
2	175000
3	225000
4	300000