PRODUCT PORTFOLIO

Quality For Life
Is Our Passion

- Tablets
- Capsules
- Liquid
- Ointment & Cream
- Sachet Powder

DEN MARK®
PHARMACEUTICALS
Quality for Life is our Passion

Delivering Hope Happily.

Where Science and Compassion Converge: Discover the Difference at Den Mark. Driven by a Commitment to Your Health, Den Mark Pioneers Tomorrow's Solutions Today.

"We are the largest independent manufacturing company in the Pharmaceutical space. We are dedicated to providing high quality Tablets, Capsules, Ointment, Sachets, Liquid Orals to our clients with the goal of making people's lives healthier.

MISSION & VISION

The mission of our company is to contribute to the protection and maintenance of the health of as many people as possible by providing quality and affordable pharmaceutical products. Our vision is to take a leading position in the markets where we operate and in the future on a global scale.

GLOBAL PRESENCE

DEN MARK PHARMACEUTICALS PVT. LTD. has significantly expanded its borders, quickly opening new regional operations in economically developing countries. In strict sequence, we have achieved impressive influence in many countries of the world, despite economic indicators, expected profit or complexity of the task. We Work on the significant development of our company in Latin America and Africa in coming future.

BUSINESS MODEL

DEN MARK PHARMACEUTICALS PVT. LTD. specialises in developing new formulations, undertaking bio equivalence studies clinical trials, obtaining the approval from the Drug Controller General of India (DCGI) for manufacturing and marketing new Fixed Dossage Combinations (FDCs) and molecules & Latest Technology driven products.

- New Formulation with DCGI Approvals.
- Contract Research and Mfg.Service
- Technology Transfer
- Loan License
- Technical Collaboration and Joint Venture
- Formulation & Development of Novel Drugs Delivery System (N.D.D.S.)
- Institutional Business
- Global Exports

MANAGEMENT

Mr.Samad H. Patanwala
Managing Director

The young and dynamic senior management of our company brings a fresh perspective and boundless energy to the table. Their innovative thinking and willingness to embrace change propel our organization forward in today's fast-paced business landscape. With their blend of experience and modern insights, they inspire creativity, foster collaboration, and drive results. Their leadership style encourages autonomy, growth, and adaptability, empowering teams to flourish and achieve excellence. Together, they epitomize the spirit of innovation and agility, steering our company toward continued success in an ever-evolving marketplace.

Mr.Shakoor Shaikh
Advisory Board

Mr.Narayan Sangem
Administration Head

Mr.Shahnazar Ali
Manufacturing
Operations Head

Mr. Gulam M Shaikh
International
Business Head

Mr.Aiman Ansari
Domestic Sales Head

Mr.Jimit Gandhi
General
Divisions Head

PRODUCTION CAPACITY

TABLETS
20,00,000/Day

OINTMENTS
20,000/Day

SACHETS
10,000/Day

CAPSULES
10,00,000/Day

LIQUIDS
30,000/Day

01. COATED
02. UNCOATED
03. CONTROLLED RELEASED
04. SUSTAINED RELEASED
05. CHEWABLE
06. DISPERSIBLE
07. BI-LAYERED
08. PVC BLISTER
09. ALU-ALU
10. STRIP

01. HARD GELATIN
02. PET BOTTLES
03. GLASS BOTTLES
04. SYRUP
05. SUSPENSION
06. GEL
07. LOTION
08. CREAM
09. LAMINATE TUBES
10. DISPERSING JARS

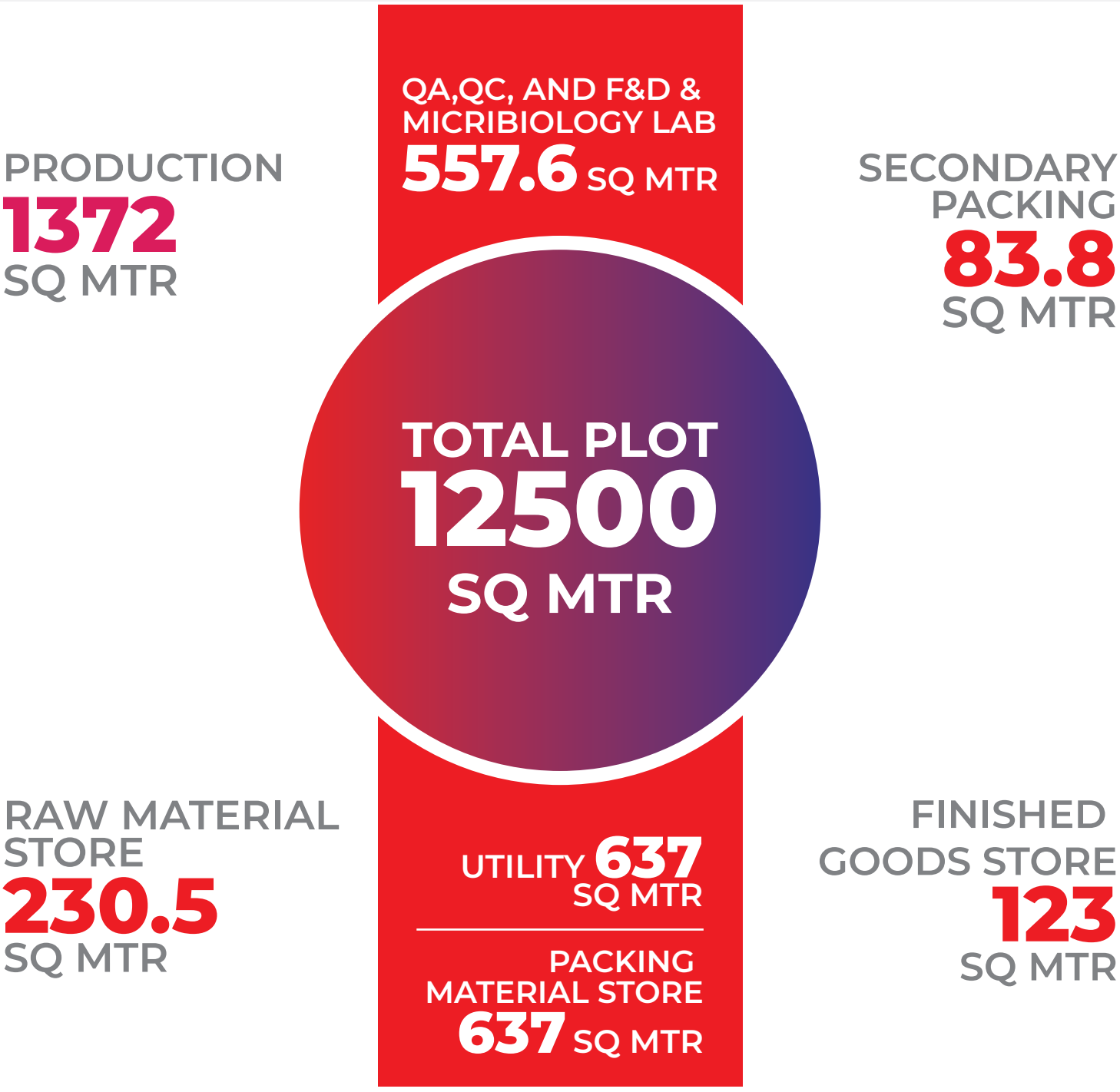


CENTRALISED STABILITY CENTRE

STABILITY CHAMBER *(Temperature/Humidity)*	**CAPACITY** *(Liters)*
Accelerated Stability study (40°C±2/75% RH ±5)	1000 L
Long term Stability study (30°C±2/75% RH ±5)	8000 L
Long term Stability study (25°C±2/60% RH ±5)	8000 L
Long term Stability study (30°C±2/65% RH ±5)	8000 L


Infrastructure Plant Details Area, & Equipment .

World-class manufacturing facilities spread over a land area of 1,20,000 sq.ft., our state-of-the-art manufacturing facility which measures 55,000 sq.ft. & having a green space of 50,000 sq. ft. has been built with a vision to achieve global accreditations like WHO-GENEVA, PIC/S and EU-GMP in foreseeable future.



We have a dedicated section for Formulation Development and Analytical development activity. Having our own manufacturing facility as per stringent international norms ensures that every product that we deliver is world-class.

WE ARE CERTIFIED

FOOD SAFETY & DRUGS ADMINISTRATION, UTTARAKHAND
DIRECTORATE GENERAL OF MEDICAL HEALTH AND FAMILY WELFARE
SAHASTRADHARA ROAD, DEHRADUN (UTTARAKHAND)

. F.No. 26/1/DRUGS/69/2023/ 16194 Dated: 05 -09 -2023

G.M.P. CERTIFICATE

This is to certify that **M/S Den Mark Pharmaceuticals Pvt. Ltd,Situated at Khasra No.764-66,NH-73,Vill-Kisanpur,Jamalpur,Pargana-Bhagwanpur,Tehsil-Roorkee,Disrt- Haridwar Uttarakhand** has been licensed under Drug & Cosmetics Act 1940 & Rules there under. They are holding Valid Drug manufacturing licence bearing no. **09/UA/2023** on **Form 25** and **07/UA/SC/P-2023** on **Form 28 valid up to 07-05-2028** to manufacture for sale of drugs.

In view of report dt. 08-05-2023 of Inspector of Drugs, Roorkee, it is certified that **M/S Den Mark Pharmaceuticals Pvt. Ltd,Situated at Khasra No. 764-66,NH-73,Vill-Kisanpur,Jamalpur,Pargana-Bhagwanpur,Tehsil-Roorkee,Disrt-Haridwar Uttarakhand** conforms to requirements of Rule 71, 74, 76 & 78 and **Good Manufacturing Practices** as laid down under Revised **Schedule "M"** of the Drugs & Cosmetics Rule 1945.

This certificate is issued to the firm on their request for submission to Govt. department / Institutions / Overseas Authority.

This certificate is valid for a period up to three years from the date of issue.

(Tajber Singh)
Drug Licensing & controlling authority
Uttarakhand

FOOD SAFETY & DRUG ADMINISTRATION
SAHASTRADHARA ROAD, DEHRADUN (UTTARAKHAND)

F.no. 26/1/Drugs/69/2023/ 2287 Date: 16 /02/2024

Certificate of Good Manufacturing Practices

Certificate no.: 26/1/Drugs/69/2023/

On the basis of the inspection carried out on 25-01-2024 & 29-01-2024 we certify that the site indicated on this certificate complies with Good Manufacturing Practices for the dosage forms, categories and activities listed in Table 1.

1- Name & Address of site: Mis Den Mark Pharmaceuticals, Pvt.Ltd., Khasra.No.764-66, NH-73, Village-Kishanpur Jamalpur, Pargana-Bhagwanpur Tehsil- Roorkee, Distt-Haridwar, Uttarakhand (India).

2- Manufacturer's license number: **Form 25 - 9/UA/2023**
Form 28 - 7/UA/SC/P-2023

3- Table 1:

Dosage form (s)		Activity(ies)
Tablets	Non Beta Lactam	Manufacturing
Capsules,	Non Beta Lactam	Manufacturing
Oral Liquid		Manufacturing
Sachet	Non Beta Lactam	Manufacturing
External Preparations (Cream, Ointment, Gel, Lotion, Shampoo, Oral rinse & Dusting Powder)		Manufacturing

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 15-02-2027. 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.

The firm is following ***Good Manufacturing Practices as per World Health Organization (WHO) TRS Guidelines TRS No. 908 of 2003*** *in the Manufacturing & testing of the said categories of Products and items in respect of which the* ***Certificate of Pharmaceuticals products*** *have been issued*

Address of certifying Authority:-
Office of Commissioner, Food Safety and Drug Administration Uttarakhand , Dehradun India.

Name & function of responsible person:
Shri Tajber Singh
Drugs Licensing & Controlling Authority
Uttarakhand
Email: *drugscontroluk@gmail.com*
Tel.no. *NA*
Fax. no. *0135-2608874*

(Tajber Singh)
Drugs Controlling & Licensing Authority
(Uttarakhand)

(Tajber Singh)
Drugs Controlling & Licensing Authority
Uttarakhand

FORM-25
[See Rule 70]
License to manufacture for sale [or for distribution] of drugs other than those Specified in Schedules C, C (1) and X

Number of license and date of issue **9/UA/2023** *Date of issue* **08-05-2023**

I, ***M/s Den Mark Pharmaceuticals Pvt. Ltd,*** *is hereby licensed to manufacture the following categories of drugs being drugs other than those specified in Schedules C, C (1), and X to the Drugs and Cosmetics Rules, 1945, on the premises situated at-* ***Khasra No-764-66 NH-73, Village-Kishanpur Jamalpur Pargana-Bhagwanpur, Tehsil-Roorkee, Dist. Haridwar Uttarakhand (India)*** *under the direction and supervision of the following competent technical staff.*

(a) Competent Technical Staff (Names)

For Manufacturing-
1- Mr Sanjay Kumar Patel (Tablets)
2- Mr Shahnazar (Capsules)
3- Mr Prakash Prasad Singh (Liquid Oral &External Preparation)

For Analysis-
1- MOHD Jamshed (Chemical, Instrumental & Microbiological)

(b) Name of Drugs- As per list enclosed

2. The license authorizes the sale by way of wholesale dealing and storage for sale by the licensee of the drugs manufactured under the license, subject to the conditions applicable to license for sale.
3. The license shall be in force from **08-05-2023 to 07-05-2028.**
4. The license is subject to the conditions stated below and to such other conditions as may be specified in the rules for the time being in force under the Drugs and Cosmetics Act, 1940.

Date: 08-05-2023

Signature.....
Designation.....

Conditions of License

1. *This license and any certificate of renewal in force shall be kept on the approved premises and shall be produced at the request of an Inspector appointed under the Drugs and Cosmetics Act 1940.*
2. *Any change in the expert staff named in the license shall be forthwith reported to the Licensing Authority.*
3. *If the licensee wants to manufacture for sale additional items of drugs not included above he should apply to the Licensing Authority for the necessary endorsement as provided in Rule 69 (3). The license will be deemed to extend to the drugs so endorsed.*
4. *The licensee shall inform the Licensing Authority in writing in the event of any change in the constitution of the firm operating under the license. Where any change in the constitution of the firm takes place, the current license shall be deemed to be valid for a maximum period of three months from the date on which the change takes 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.*
5. *The Licensee, Who is Marketing or engaged in manufacture of drug containing psychotropic substance under the Drug License, shall have to register and shall file quarterly return with the Narcotics Commissioner for each of the substance in the form and manner as many be specified by the "Narcotic Commissioner" prior to initiating and activities mentioned therein under the license issued to them. Vide GSR 224(E) dt.* **25-03-2015.**

UQSR.

UQSR.
Certificate

UQSR Global Private Limited has assessed the Quality Management Systems of

DEN MARK PHARMACEUTICALS PRIVATE LIMITED

KHASRA NO.- 764-66, NH-73, KISHANPUR JAMALPUR, PARGANA BHADWANPUR, ROORKEE, HARIDWAR, UTTARAKHAND- 247667, INDIA

And hereby declares that the organization is in conformance with

ISO 9001:2015

For the following scope of activities:

MANUFACTURING OF PHARMACEUTICAL PRODUCTS

Further clarification regarding the scope of this certificate and the applicability of Quality Management Systems standard requirements may be obtained by contacting the organization

Certificate Number : UQSR-3521-DMPPL

Original Issue Date	: 21/02/2024	Current Issue Date	: 21/02/2024	IAF Code: 13
Surveillance Date	: 20/02/2025	Recertification Date	: 20/02/2027	Issue No. 01

* Validity of certificate is subjected to the continued satisfactory performance during surveillance audit

Authorized By

Certification Manager

UQSR Global Private Limited
www.uqsr.org

For information concerning validity of certificate, you can visit the site: www.uqsr.org

This document is property of UQSR and should be returned on request. Any unauthorized alteration, forgery or falsification of the content or appearance of this document is unlawful and offenders may be prosecuted to the fullest extent of the law.

OUR UPCOMING CERTIFICATIONS

Our Manufacturing facility in Roorkee, Uttarakhand, INDIA is spreaded across an area of 12000 Sq .Mtrs (Approx) with Built up area of around 75000 Sq ft.. Our Facility is Compliant of has been marked approved by authorities of WHO-GMP. Our plant pursues to easily clear EU-GMP, ENVIZA, PIC's and many more world-wide country-wise audits by End of this year.

CONTACT INFORMATION

Corporate Office Address:

1st floor, Harish Textiles, Parsi Panchayat Road, Near WESTERN EXPRESS HIGHWAY, Andheri EAST, Mumbai - 400 069, INDIA.

Email: gmshaikh@denmarkpharma.com | Mob.: +91 97691 06504

Manufacturing Facility Address:

Khasra No. 764-66, NH-73, Karondi, Roorkee - 247667, Uttarakhand, INDIA.

Email: shahnazar@denmarkpharma.com | Mob.: +91 96750 04403

Regional Office Address:

205 & 206 Matrix Tower, Corporate Road, Sarkhej -Gandhinagar Hwy, near Vodafone Office, Ahmedabad, Gujarat-380054.

Email: aiman@denmarkpharma.com Mob.: +91 96628 20878

DEN MARK®
PHARMACEUTICALS
Quality for Life is our Passion

WHO-GMP & ISO 9001:2015 CERTIFIED

www.denmarkpharma.com