HOF pharma
a better tomorrow

HOF Pharmaceuticals Limited
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a better tomorrow

Delivering hope happily

Company Overview

HOF Pharmaceuticals Limited is a part of the HOF group of companies that has a leading presence in Furniture, Infrastructure real estate and allied industries with an experience of more than 35 years in the manufacturing industry.

Quality, Integrity, Transparency and Customer service are the four pillars on which we have established ourselves and are sure of our growth in the coming times. Our board members comprise of the best brains in the pharmaceutical industry who individually have more than three decades of work experience with the top Global and Indian pharmaceutical companies.

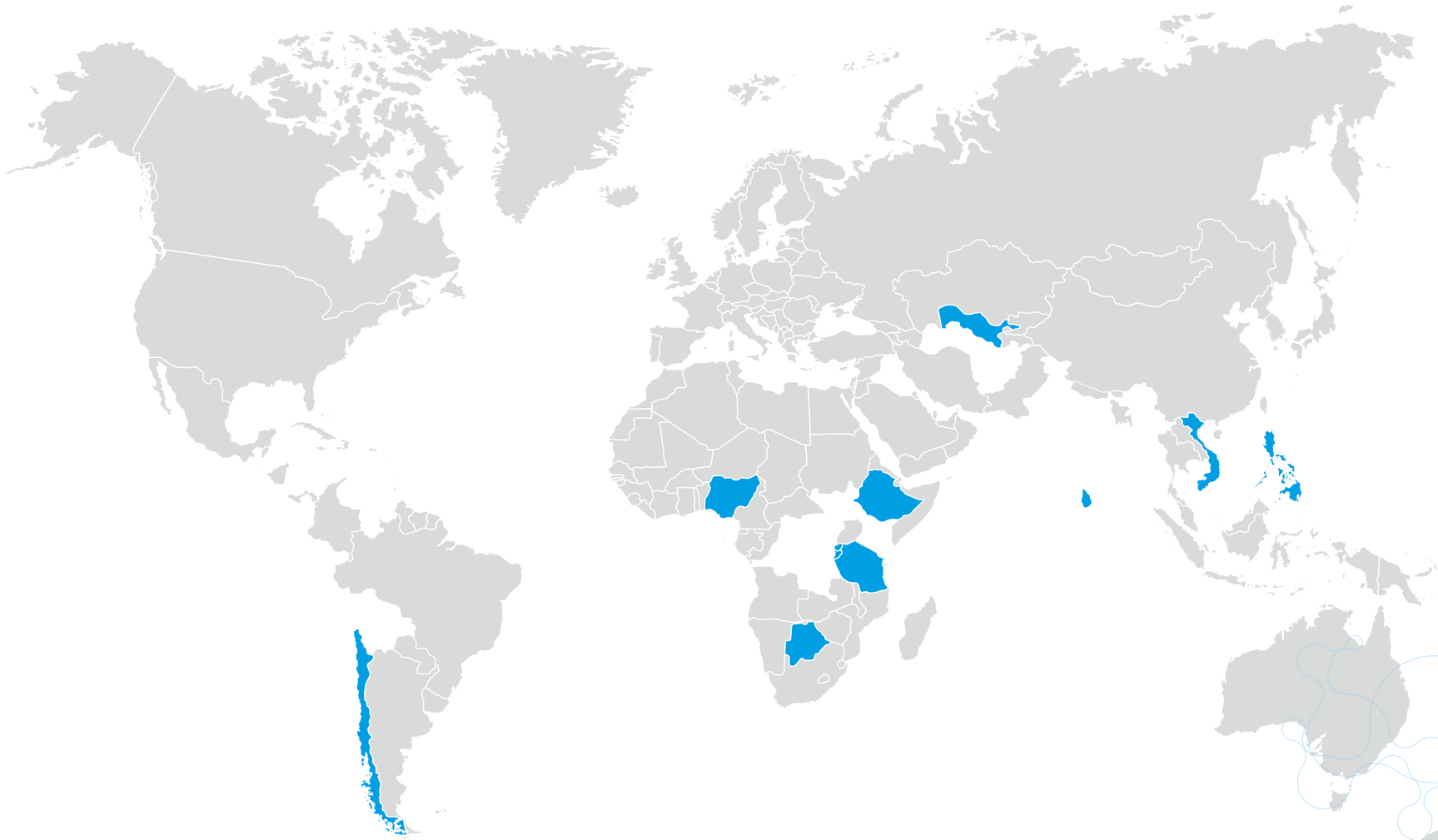
In addition to that, the senior management is aptly supported by young managers who come from a business background and are principally responsible for implementing the strategies and working on the feedback which leads to continuous improvement in our company.

We are a
WHO-GMP certified
manufacturing facility of Tablets,
Capsules and Oral Liquids.
We also offer contract
manufacturing services

Global Presence

India | Kenya | Rwanda | Burundi
Uganda | Tanzania | Ghana | DRC
Botswana | Bolivia | Ecuador
Venezuela | Peru | Chile | Costa Rica
Guatemala | Honduras | Dominican
Republic | Iraq | Myanmar Cambodia |
Vietnam | Philippines Belize |
Afghanistan | Turkmenistan

Our in-house Regulatory support ensures that all our products comply with regulations laid by individual countries and industry standards.



2021 — PLANT COMMISSIONED

2022 — WHO GMP

2023 — KENYA PPB AND CAMBODIA MOH

2024 — UGANDA NDA AND GHANA FDA

2025 — RWANDA FDA, PHILIPPINES FDA

2026 — PERU

2027 — PIC/S

2028 — EU-GMP

Accreditation

We prioritize GMP compliance through robust Quality Management System.

We follow International Standards, ensure quality control and timely execution to satisfy customer requirements

Our focus on Quality, Regulatory Compliance and Innovative Formulation Development allows us to provide our customers with products that are safe, effective and of the highest quality.

Active Presence

ASIA
India | Myanmar | Philippines
Cambodia | Afghanistan

AFRICA
Kenya | DRC | Ghana | Uganda
Rwanda | Botswana | Burundi

LATAM
Guatemala | Costa Rica | Honduras | Venezula | Peru
Ecuador | Bolivia | El Salvador | Dominican Republic

MIDDLE EAST
Iraq

CIS
Turkmenistan

Pipeline

ASIA
Vietnam | Sri Lanka

LATAM
Chile

AFRICA
Tanzania | Ethiopia

CIS
Uzbekistan

Certifications

WHO-GMP | UGANDA NDA | CAMBODIA MOH | RWANDA FDA

KENYA PPB | GHANA FDA | PHILIPPINES FDA | BOTSWANA MRA

PERÚ Ministerio de Salud

Pipeline Certifications

EFDA | TMDA

Mission

Our mission is to serve humanity with quality medicines that spread the joy of good health and thus ensure a better tomorrow.

Vision

We will occupy a prime position in the pharmaceutical industry through stringent quality practices in all our activities, right from manufacturing to customer service. We aim to mark our presence in top global markets around the world

a better tomorrow

Quality Systems

Excellence Through Quality, Compliance & Analytical Precision

At HOF Pharmaceuticals, the Quality Department is structured with integrated QA, QC, and ADL operations to ensure consistent product quality, regulatory compliance, and analytical excellence across all manufacturing processes.

The Analytical Development Laboratory (ADL) supports formulation development through analytical method development, method validation, and technical evaluation aligned with regulatory requirements.

The Quality Control (QC) team performs comprehensive raw material, packing material, in-process, finished product, and stability testing using advanced analytical practices to ensure accuracy, consistency, and product integrity.

Our Quality Assurance (QA) division focuses on GMP compliance, documentation systems, validation activities, and process monitoring to maintain robust quality standards and operational reliability.

Backed by advanced laboratory infrastructure, skilled professionals, and robust quality systems to deliver globally compliant pharmaceutical products.

Core Activities

- Change Control • Deviation Management • CAPA System
- SOP Management • Validation & Qualification • Stability Monitoring
- Documentation Control • GMP Compliance • Annual Product Revision
- Complaints / Recall Procedure • Self Inspiration • Vendor qualification
- Training and development of person

Board of Directors

Our leadership team brings together decades of expertise across pharmaceuticals, manufacturing, operations, technology transfer, and global business development. With a strong foundation in quality, innovation, and strategic execution, the board drives the organization with a shared vision of sustainable growth, operational excellence, and long-term partnerships.

Pravin Patel
Chairman

Leading the group
since last 35+ years

Dr Soumitra Banerjee
Director, Strategy

Pharma Experience:
35+ years
Techno Commercial

Mansukh Dhanani
Director,

Operations & Technology
Pharma Experience:
35+ years
Formulation Development
& Manufacturing

Abhigna Patel
CEO

Business Development

Darshan Dhanani
COO

Operations

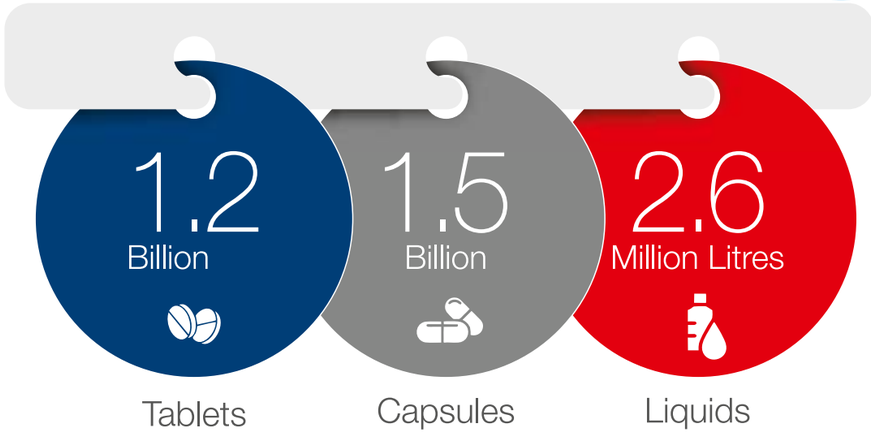
Infrastructure

Plant Area Details & Equipments

World-class manufacturing facility spread over a land area of 1,50,000 sq.ft., our state-of-the-art manufacturing facility which measures 1,00,000 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. Our own state of the art manufacturing facility follows stringent international norms ensuring that every product that we deliver is world-class.

Installed Capacities Annually



Plant Area

01	02	03
Utility	**Stores**	**R&D / Lab**
1400 Sq mtr	3700 Sq mtr	500 Sq mtr
04	**05**	**06**
Admin	**Manufacturing Area**	**Other**
500 Sq mtr	1300 Sq mtr	500 Sq mtr

Total builtup area
7900 Sq mtr

Regulatory Affairs

Ensuring Global Compliance & Market Access

About Department

Our in-house Regulatory Affairs team oversees global regulatory compliance, dossier management, and market authorization processes to support smooth product approvals and international market expansion.

Core Activities

- Product Registration & Dossier Filing
- CTD / e-CTD / ACTD Documentation
- Regulatory Query Management
- Variation & Renewal Support

Key Strengths

- Global Regulatory Expertise
- Timely Submission Management
- Strong Documentation System
- Market-Specific Compliance Support

Delivering compliant pharmaceutical solutions with efficiency, and regulatory excellence.

Formulation & Development (FND)

Innovative Formulations for Global Healthcare

About Department

Our in-house Formulation & Development department operates from a fully dedicated floor spanning over 500 sq. mt., designed to support advanced formulation development, product optimization, and regulatory-focused pharmaceutical innovation.

Infrastructure Highlights

- Modern Development & Trial Facilities
- Efficient Cross-Functional Workflow

Core Activities

- Formulation Development
- Product Optimization
- Technology Transfer