

Sekhmet Pharmaventures

“Your trusted business partner”

SEKHMET PHARMAVENTURES

SEKHMET GROUP

2020

Sekhmet Pharmaventures was formed.

2020

Anjan Bulk Drugs Pvt. Ltd. was **acquired** - world renowned manufacturer of Intermediates and APIs

2022

Optimus Group was acquired
- a global player in the manufacture of Intermediates, APIs and Formulations



Optimus



Founded in 2004
3 API & 1 FDF
Facilitates



Anjan Drug
Private Limited

Founded in 1990,
2 API Facilities

SEKHMET PHARMAVENTURES AT GLANCE

Revenue: USD 135+ Mn

One of the fastest growing
CDMO/API Manufacturer

Facilities: 6 Mfg. Plants (5
APIs, 1 FDF)

API Plants with Capacities
over 1300 KL+

Products: 70+ APIs, 30+
FDFs

Strong portfolio of API
products with 50+ USDMFs.

Global Accreditations:

USFDA, EU, ANVISA,
WHOGMP, KFDA, TGA

Globally recognized and
accredited mfg. sites to cater
global markets.

Legacy: 20+ Years of
Experience in API
Manufacturing

Optimus & Anjan has rich
experience and Global
recognition.

Key Customers: Sanofi,
Abbott, Pfizer, Sumitomo,
Teva & more.

Trusted supplier to
Innovators and Big Pharma.

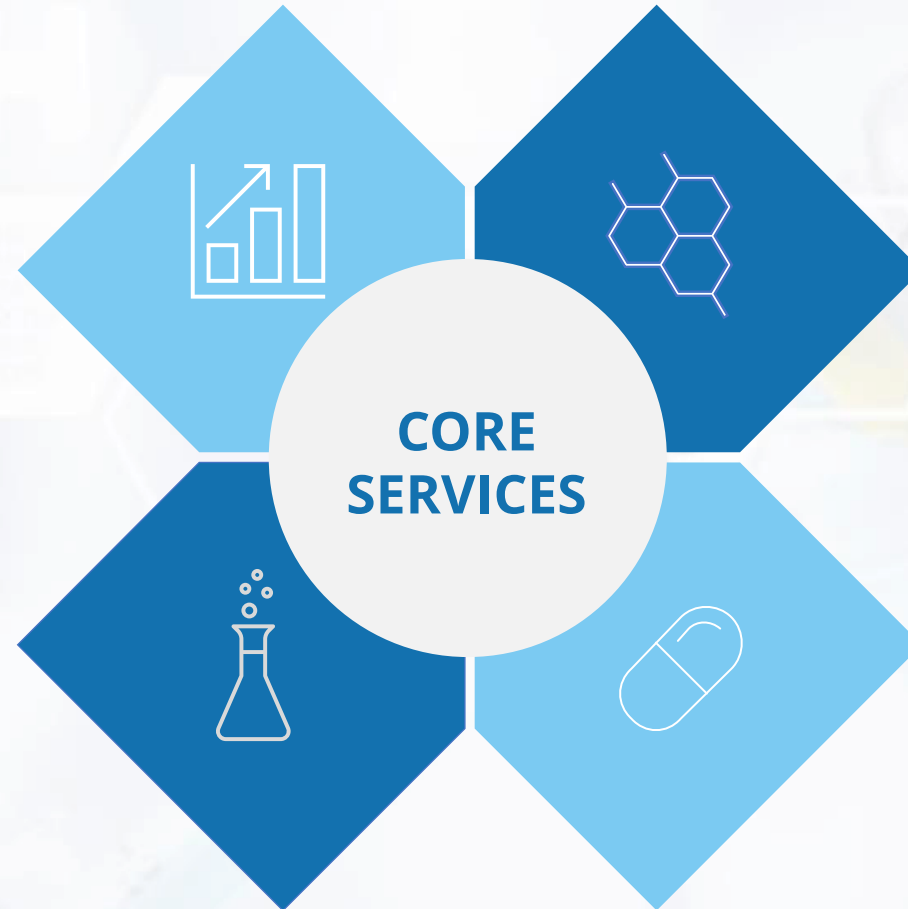
OUR EXPERTISE

CDMO & CMO

Strong R&D to develop and manufacture products for our local and international partners

ACTIVE PHARMACEUTICAL INGREDIENTS

Fastest growing API player with 70+ products across therapeutic segments



ADVANCED INTERMEDIATES

32+ years of experience in producing Advanced Drug Intermediates for the global pharmaceutical industry

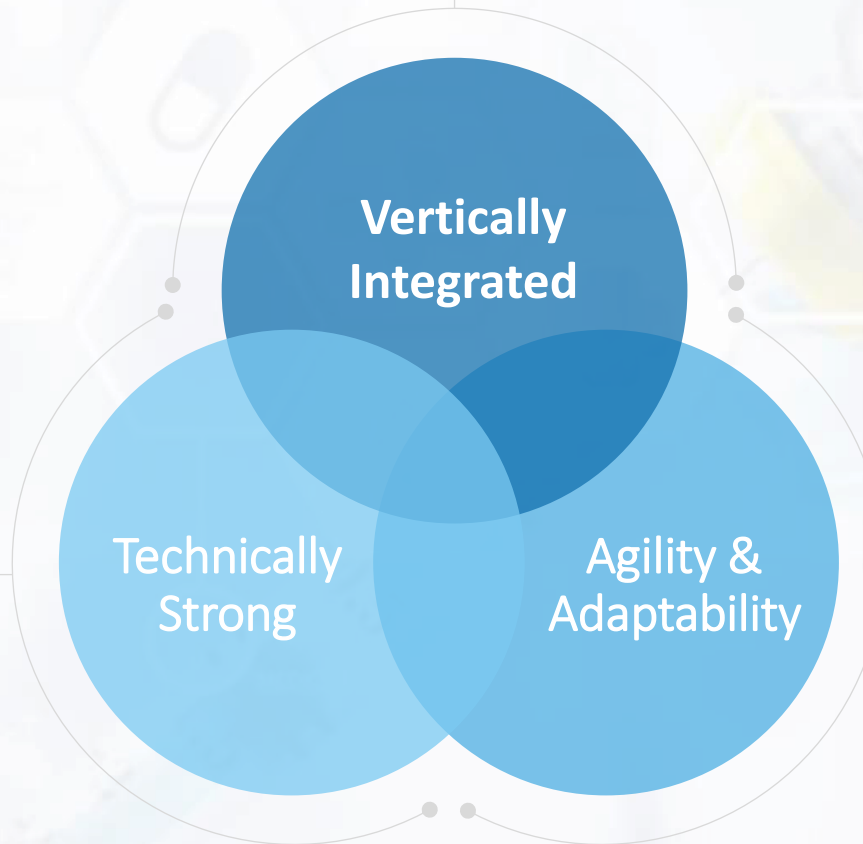
FORMULATIONS

Diverse portfolio & products registered in global markets including the US, Europe, Australia, Russia etc.

CORE COMPETENCIES

Intermediates to APIs to Finished Formulations Best Quality -
20% of our workforce designated to compliance

Focus on Research & Innovation to Develop Complex Molecules.
12% of our annual revenue spent on Research and Development



Technology Transfer from laboratory to Industry scale in **Minimum** lead time

MANAGEMENT TEAM – SEKHMET



Santosh Mahil

Managing Director and Chief Executive Officer
30+ Years



Mukesh Agarwal

Chief Financial Officer
21+ Years



Dr Jayprakash P

Head – R&D
23+ Years



Sumit Kumar

Chief Commercial Officer
24+ Years



Harsha Vardhan

Head – Quality
22+ Years



Durga Prasad

Head – Operations
30+ Years



Jitendra Jalan

Chief Human Resource Officer
25+ Years

KEY OFFERINGS - API & ADVANCED INTERMEDIATES



API

- **Comprehensive API** Solutions from process development to commercial manufacturing, with a dedicated **R&D center** and **5** strategically located **manufacturing** facilities for efficient scale-up and global supply
- Proven track record of developing over **100+** APIs developed involving various types of **complex chemistries and technologies**



RSM & Advanced Intermediates

- Our intermediates are tailored for **CDMO services**, supporting clinical phase development until commercial scale manufacturing
- **Developed over 55+ Advanced intermediates** across various therapeutic categories such as oncology, anti-coagulants, anti-inflammatory among others



Regulatory & Quality Compliance

- **Regulatory Expertise:** Sekhmet Pharma supports clients with over 100+ regulatory filings globally, including **50+ USDMFs** and **14+ CEPs**
- Partnered with Innovators for **NDA** Programs – including Regulatory CMC Support
- **Advanced Risk Assessment:** Expertise in mutagenic impurity classification and **N-nitrosamine** risk assessments ensures safety compliance with regulatory guidelines

KEY OFFERINGS - FORMULATIONS

“Our offerings include tablets, capsules, pellets, creams, and lotions across product categories”



Tablet Dosage Forms

- **Immediate release** ; Extended release by matrix, reservoir, osmotic and bilayer
- **Delayed Release** by enteric coating by MUPS (multi unit particulate System), Micro tablets by multi tip tooling technique
- Tablets **for Oral Suspension**; Dispersible tablets; Chewable tablets ; Effervescent tablets



Powders, Pellets & Capsules

- **Wurster coating**, Extrusion and Spheronization
- Hard gelatin capsules filled with powder, pellets, mini tablets and combination of tablets & Pellets
- Dual drug delivery ; Multi unit particulate system
- Powder For Oral Suspension (**PFOS**)



Semisolids

- **Ointments**, Creams & Lotions
- Controlled/**Extended-Release** Gel

STATE-OF-THE-ART R&D CENTER

“Our R&D collaborates with **Global Pharmaceutical** , Specialty Chemical Companies and **Innovators** to develop innovative, safe, and cost-effective processes for NCEs, RSMs, Advanced Intermediates and APIs”



R&D Center
Hyderabad, India

Salient Features

- 45000 sq. ft. dedicated to R&D infrastructure
- Well-qualified team of over **150+** experienced scientists working on challenging product development
- Adherence to strict regulatory guidelines and compliance with **GLP** and other relevant quality standards
- Advanced Infrastructure such as Kilo Lab, Oncology Facility,

CAPABILITIES – OVERVIEW



Synthesis

- Expertise in **complex** and innovative chemistries
- **Green** Chemistry
- Average time **4 to 5 Stage** Development is ~ **10 to 12 months** (*par with Benchmark)



Formulation Development

- Topicals, Tablets, Capsules, Liquid Orals
- **Extended-release** dosage form available
- Stability Studies , Method Transfers
- **Method validation** support for NDA/ANDA



Analytical Chemistry

- **24*7** Analytical support
- LC-MS, GCMS, XRD capabilities
- ✓ Method Development and validation for **nitrosamine** impurities
- ✓ **21 CFR** Compliance



Value-Added Services

- **100% Regulatory** Support from product development to commercialization
- **Robust IP** Management Support



Technology Transfer (TT) & Scale-Up

- Fast Technology transfer from **Lab** Scale to **Kilo** Scale followed by **Piloting**
- Dedicated **5-member** team for Technology Transfer
- Ensuring Process **Safety** & Quality **Consistency**



End-to-End Project Management

- Risk-Based **Stage Gate Process** implemented for Product Development
- **17** Project Squads for New Product Development and Cost Improvement
- **PMO** – **3** Project Managers + **1** Program Manager

* Time taken from initiation to tech transfer as per benchmarking study

CAPABILITIES – R&D – ANALYTICAL



LCMS



XRD

✓ State-of-the-Art Analytical Facility

- **10,000+ sq. ft.** infrastructure with ultra-modern equipment
- Experienced team of **80+ analysts** providing 24/7 support
- ✓ **GLP-compliant** facility with a dedicated team of scientists for R&D needs

✓ Comprehensive Method Development & Validation

- Method **development** and **validation** aligned with global standards (ICH Q2(R1), Brazil, China, etc.)
- Transfer of analytical methods to QC at respective sites

✓ Chromatography & Specialized Testing

- Systems operating on Empower 3 with 21 CFR compliance.
- Expertise in detecting **genotoxic** impurities, **nitrosamines**, and NDSRIs using **XRD, HPLC, LC-MS, GC-MS**
- Particle size analysis using Malvern PSD analyzers

✓ Stability & Impurities

- Stability studies for development batches
- **Mass** and **NMR** Expertise
- Identification of unknown impurities/solvents via advanced LC-MS and GC-MS

CAPABILITIES - MFG. NETWORK | API

Unit-1 Hyderabad



- ✓ API and Advanced Intermediates
- Total Capacity – **295 KL**
- # of Clean Rooms – **9** (ISO Class 8)
- Area – 18 Acres
- Reactor configuration suitability: **Potent Molecules**
- ✓ Accreditation: USFDA, EUGMP, WHO-GMP, KFDA

Unit-2 Vishakapatnam



- ✓ API and Advanced Intermediates
- Total Capacity – **240 KL**
- # of Clean Rooms – **1** (ISO Class 8)
- Area – 10 Acres
- Reactor configuration suitability: **High Volume Products**

Unit-3 Hyderabad



- ✓ API and Advanced Intermediates
- Total Capacity – **380 KL**
- # of Clean Rooms – **6** (ISO Class 8)
- Area – 20 Acres
- Reactor configuration suitability: **Multi-Product**
- ✓ Accreditation: EUGMP, WHO-GMP, KFDA

Unit-5 Chennai



- ✓ API and Advanced Intermediates
- Total Capacity – **65 KL**
- # of Clean Rooms – **5**
- Area – 2 Acres
- Reactor configuration suitability: **API Purification**
- ✓ Accreditation: USFDA, EDQM, TGA, WHO-GMP, AFM

Unit-6 Chennai



- ✓ Advanced Intermediates
- Total Capacity – **300 KL**
- # of Clean Rooms – **1**
- Area – 12 Acres
- Reactor configuration suitability: **High Volume Products**
- ✓ Accreditation: USFDA, EDQM, WHO-GMP, AFM

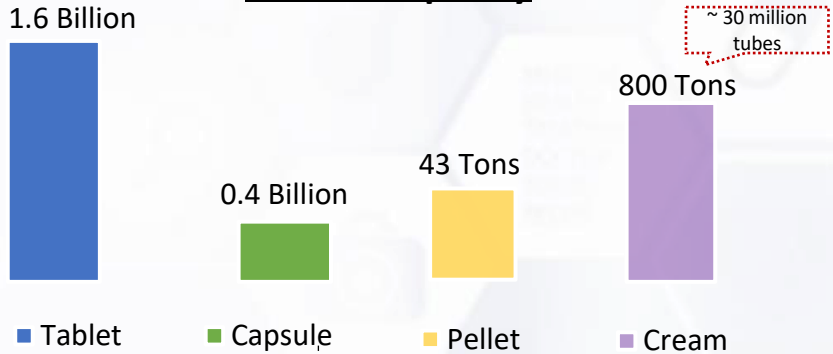
CAPABILITIES - MFG. NETWORK | FDF

Manufacturing Infrastructure Overview

~2 Billion/Year Capacity Tablets & Capsules



Annual Capacity

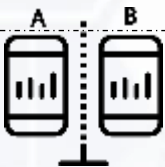


Area	No of line/Suites
Granulation	4 Suites
Compression	2 Suites [Legacy /Fette]
Coating	3 Suites
Encapsulation	1 Suite
Blister packaging	2 Lines (BQS / B-Max)
Bottle packaging	2 Line (Parle / Pharma pack)
Creams manufacturing	2 Suites
Creams/Lotion filling	2 Lines



Dosage Forms

- OSD**
 - Tablets
 - Capsules
 - Pellets
 - PFOS
- SSD**
 - Creams
 - Lotions



Analytical Testing Capabilities

- Equipment: **12 HPLC**, **2 GC** with Empower 3FR4 software
- Server Integration: In-line with **21CFR** requirements
- Microbial Testing**: Dedicated lab
- Chemical Lab: 5 dissolution apparatus with autosampler
- Advanced Tools: **XRD & LC-MS** for R&D investigations
- Expertise: Method transfer, validation, and stability studies



Stability Study Testing Capabilities

- 5 walk-in chambers (**25KL capacity**) with 21 CFR-compliant ICDAS software
- Additional Chambers: Photo stability and **2-8°C cooling chambers**
- Compliance: **GXP SOPs** for OOS, deviations, audit trails, and data backup

ESG - KEY HIGHLIGHTS AT SEKHMET



810 metric tons CO2 emission avoided through introduction of renewable energy



Invested over **Rs. 9+ Crores** in CSR Projects



Zero reportable fatalities and LTIFR during the FY23, FY24, showcases the commitment to safety, health and environment



Reduced Scope 1 & 2 emission from **51,761 tons** of CO2e in FY 23 to **36,489 tons** of CO2e in FY 24



Achieved **30% water recycling** from total water withdrawal



1307 metric tonnes of high calorific hazardous waste was **redirected to co-processing in cement industry** aligning our commitment to responsible waste disposal in FY 23



Achieved Bronze medal in Ecovadis for our commitment to sustainable practices. Optimus 68 (Bronze), Anjan (67 (Bronze). Aspiring Gold Rating by 2027-28.



Emission Intensity reduced from **11.12 KgCO2e in FY 23 to 6 KgCO2e in FY 24**



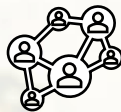
All our sites have embraced the **3R waste hierarchy**



Achieved annual power savings of 66,44.00 kw , **cost savings of Rs. 5,94,656** through installation of BLDC motors in operational sites IN FY 23



73% of total spent is made on significant suppliers in tier 1



Reported **zero cases of bribery and corruption**



Click on report to access Sekhmet's Sustainability Report FY 23

Green & Sustainable Pharmaceutical Manufacturing Solutions

OUR GREEN MANUFACTURING EDGE



Water-Based Processes

Our Chemistry in water solutions to reduce Solvent dependency for safer, cleaner synthesis



Energy-Efficient Operations

Optimized reaction conditions to minimize energy footprint



Solvent-Free Technologies

Neat reactions to eliminate hazardous waste



Waste-Minimization

Closed-Loop recovery and recycling of solvents and reagents



Sustainable Raw Material Sourcing

Suppliers evaluated under Sustainable supply chain framework

Green Chemistry driving CDMO excellence

We integrate advanced green chemistry solutions — including **water-based** and **solvent-free processes**, and waste minimization — into our Contract Manufacturing operations. These innovations **enable effluent reduction**, process intensification, and **cost efficiency** while ensuring high-quality, scalable manufacturing for our clients

Key Water Based Chemistry Capabilities

ESTER HYDROLYSIS

DECARBOXYLATION

ALKYLATION

HOFFMAN
REACTIONS

SALTIFICATION

DE-SALTING

OUR MARQUEE CLIENTS



THANK YOU