

AIKVIRA PHARMACEUTICALS

www.aikvira.com





Quality Formulations, Quality Lives









WHO WE ARE

- Our Journey started with the aspiration for providing Quality and Affordable Lifecare through Application, Innovation and the Integration of the resources.
- We have evolved from the modest start to emerge as a strong and the established player in the market with more than 10 years standing as an hard core Finished Formulations manufacturing player.
- We have scaled up to developing 173 key Finished Formulations. key Finished Formulations through our consistent efforts which has helped us to achieve a sizable business volume catering our products to all the leading companies in India and overseas, virtual manufactured at various sites while in another couple of years, we plan to develop and commercialize another new molecules.
- Expanding in the manufacturing capacities and capabilities will be the key Growth-Driver for us and the strategic capital investment into our manufacturing facilities at Paithan-Aurangabad is going to help us further in strengthening our journey towards operation excellence and which shall significantly contribute towards increasing the sales turnover, top line and subsequently the bottom line.

VISION

AIK PIRA

PHARMACEUTICALS

—NOVEL • DRUGS • DELIVERY—

"We aspire to be a Global Leader in providing the Quality and Affordable Lifecare through Application, Innovation and the Integration of the resources."

MISSION

"To deliver best Pharmaceutical Products by listening and understanding the customers' requirement and turning ideas into reality, confidentiality, knowledge, and expertise in the fields of the Product Development, Marketing, Industrial Manufacturing, Supply Chain and Regulatory Compliance".



Quality First



Innovation

• OUR IDEOLOGY

A vision to make a difference in healthcare by providing high-quality pharmaceutical products. Our ideology centres around three core principles:



Global Standard



Research Excellence

PURPOSE, STRATEGY AND CULTURE

Committed to Quality and Safety

- Commitment to quality and safety is not just a statement for Aikvira Pharma but it's a guiding principle that shapes the organization's culture, processes, and relationships with stakeholders.
- It fosters trust among customers, employees, and the community, showcasing the organization's dedication to deliver reliable, safe, and high-quality products or services



Innovation

Discover how our innovation

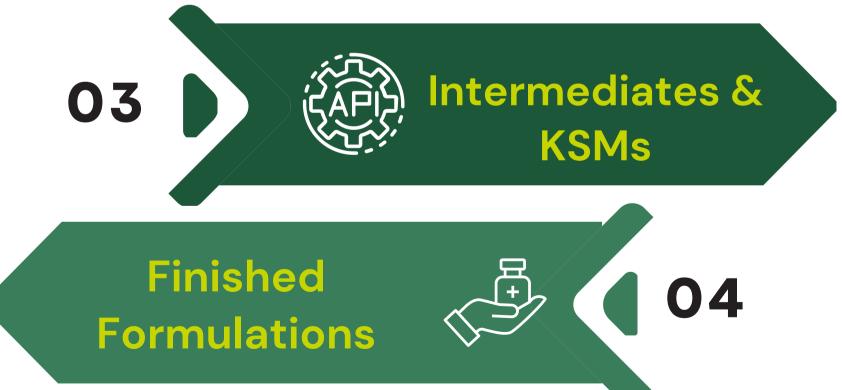
drives excellence in every aspect of pharmaceutical development.





In-House APIs and API Intermediates:









LEADERSHIP Meet our team



Mr. Sanket Patil

Managing Director



Ms. Siya Patil

Director



We are Specialized In:

- DERMATOLOGY
- CARDIOLOGY
- DIABETOLOGY
- GYNECOLOGY
- CNS
- ORTHOPEDICS/ PAIN MANAGEMENT
- IRON REPLACEMENT
- ANTI-ALLERGY/ RESPIRATORY
- GASTRO INTESTINAL TRACK
- HEPATOLOGY
- UROLOGY/NEPHROLOGY
- ANTIBIOTICS
- ANTI PARASITIC

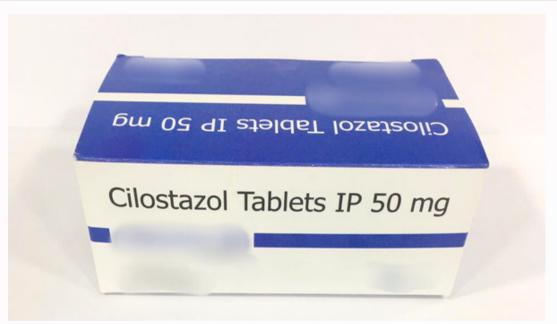
Some of Our Products

Linagliptine Tablets

Cilostazole Tablets

Glimepiride with Metformin Tablets





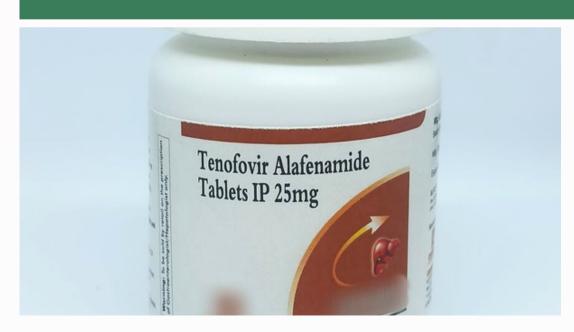


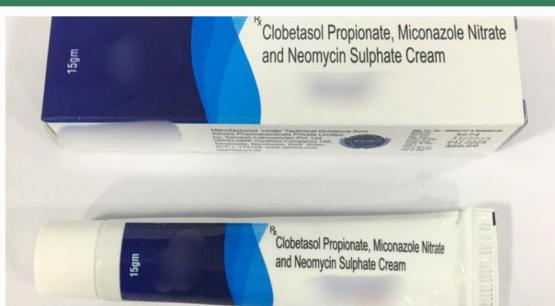


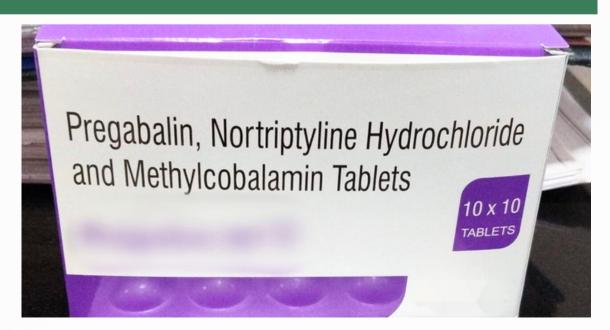
Tenofovir Alafenamide Tablets

Clopidogrel propionate, Miconazole Nitrate, Neomycin sulphate Cream

Pregabalin, Nortriptyline, HCl & Methylcobalamin Tablets









CMO/ CDMO & Third Party Manufacturing (P2P)

Aikvira Pharmaceuticals collaborates closely with clients to bring their pharmaceutical products to market. With high level of Market Intelligence, we identify the opportunities and needs of our customers by providing End to End solutions in Formulation Development, Packaging Development, Commercial manufacturing and Technical services.

We offer our expertise in areas such as drug formulation, analytical testing, regulatory affairs, and quality control along with flexible manufacturing and packaging solutions to save our client's valuable time and resources. These drug development projects can range from small-scale clinical trials to large-scale commercial manufacturing.

We also have the expertise & capacities to develop the product from idea to the commercial scale & getting necessary approvals from CDSCO office after doing Clinical Trials & Bio-equivalency Studies.

Connect with us to stay up to date with the latest technologies and innovations in the constantly evolving Pharmaceutical Industry !!!!





EQUIPMENT & FACILITIES



Manufacturing

Our Drug manufacturing Equipment and Facilities are designed to accommodate changes in production volume according to demand.



Supply Chain Management

Our reliable and robust Supply Chain is always there to help for sourcing raw materials in the required quantity.



Quality Control

We ensure that the Quality
Control measures are in place
at all stages of manufacturing
so that the end product can
meet Safety and Regulations
standards.



Medication Packaging Solutions

Our unique packaging equipments and labelling systems helps us to deliver the right medicine and dose to patients at the right time.



Formulation & Development:

- Aikvira Pharma can help you develop quality finished dosage forms. Our pharmaceutical experts start by understanding your needs and constraints and then work through appropriate pre-formulation assessment to design potential formulations (and processes) that will deliver the high-quality, robust finished drug product you need.
- Our Formulation & Development laboratories are equipped with pilot size rapid mix grinder, fluid bed dryer, compression machine and different instruments such as HPLC, UV Visible spectrophotometers, FTIR, dissolution and disintegration test apparatus etc. for test and analysis of new formulations. A formulation & development laboratories harbor stability chambers for stability test and analysis of finished formulations.
- Our key objective of F &D initiative is to match pace with modern development on solid oral dosages and development of new drug formulations for existing products. We have the expertise, and a proven track record, to formulate a wide range of finished pharmaceutical products, such as:



✓ Topical Ointment , Creams & Lotions





Best Practices followed by us:

- Maintain complete API records.
- Perform appropriate testing to assure conformance to specifications.
- Assure that each chemical step is problem-free before moving to the next step.
- Maintain thorough, complete data from laboratory tests and also include any failed attempts.
- Ensure that all equipment is reliably cleaned.
- Investigate critical deviations adequately.
- Document each step diligently and thoroughly as required by regulators.
- Have all measurement instruments on-site as now required by regulatory bodies.
- Maintain frequent and open communication between the CDMO and sponsor teams – this is critically important.

Aikvira at CPHI INDIA 2023/









THANK YOU



- 8908993399
- Info@aikvira.com
- www.aikvira.com
- Thane(w), Mumbai,
 Maharashtra- 400615, INDIA.

