Manufacturers of

- **Pharmaceutical Finished Dosage Forms**
- **Active Pharmaceutical Ingredients**
- **Drug Intermediates**

ULTRATECH INDIA LIMITED

Dedicated GenNext Health Care

Profile

Ultratech India Limited is an integrated pharmaceutical and life sciences company, specialized in the development and manufacturing of **Pharmaceutical Finished Dosage Forms (FDFs), Active Pharmaceutical Ingredients (APIs),** and **Drug Intermediates.**

Established in 1992, in its three decade journey, Ultratech has evolved into a transnational company manufacturing world-class Pharmaceutical products with a diverse product portfolio and a global clientele.

The company's branded generic medicines encompass a **diverse range of therapeutic segments, including cardiology, diabetology, antibiotics, analgesics, respiratory care drugs, anti-malarial treatments, anti-cancer drugs, gynecology, pediatric medications,** as well as **nutraceutical and health supplements**. Ultratech distinguishes itself with expertise in both conventional and advanced dosage forms, with a particular emphasis on pulmonary drug delivery systems. The company offers a comprehensive range of anti-asthmatic Inhalation Drugs in Dry Powder Inhalation **(DPI)** and Pressurized Metered Dose Inhalation **(pMDI)** forms.

In the **Active Pharmaceutical Ingredients** segment, Ultratech specializes in the manufacturing of **synthetic drugs** across a broad spectrum of therapeutic categories. The company supplies its Active Pharmaceutical Ingredients to pharmaceutical companies globally.

Additionally, Ultratech offers comprehensive Custom Research and Manufacturing Services **(CRAMS)**, including the custom synthesis of new molecules, process development for bulk active pharmaceutical ingredients, and the formulation of finished dosage forms.

Ultratech's products are manufactured in compliance with **FDA** guidelines and **cGMP** standards, ensuring the highest quality and safety. The products are supported by **Drug Master Files (DMFs), Dossiers, Free Sale Certificates (FSCs), Certificates of Pharmaceutical Products (COPPs)**, and registration files that adhere to country-specific regulations.

Our Focus

- Developing Quality Healthcare Medicines
- Continued Excellence in R&D and Manufacturing
- Expanding Our Product Portfolio
- Implementing Stringent Quality Control and Assurance

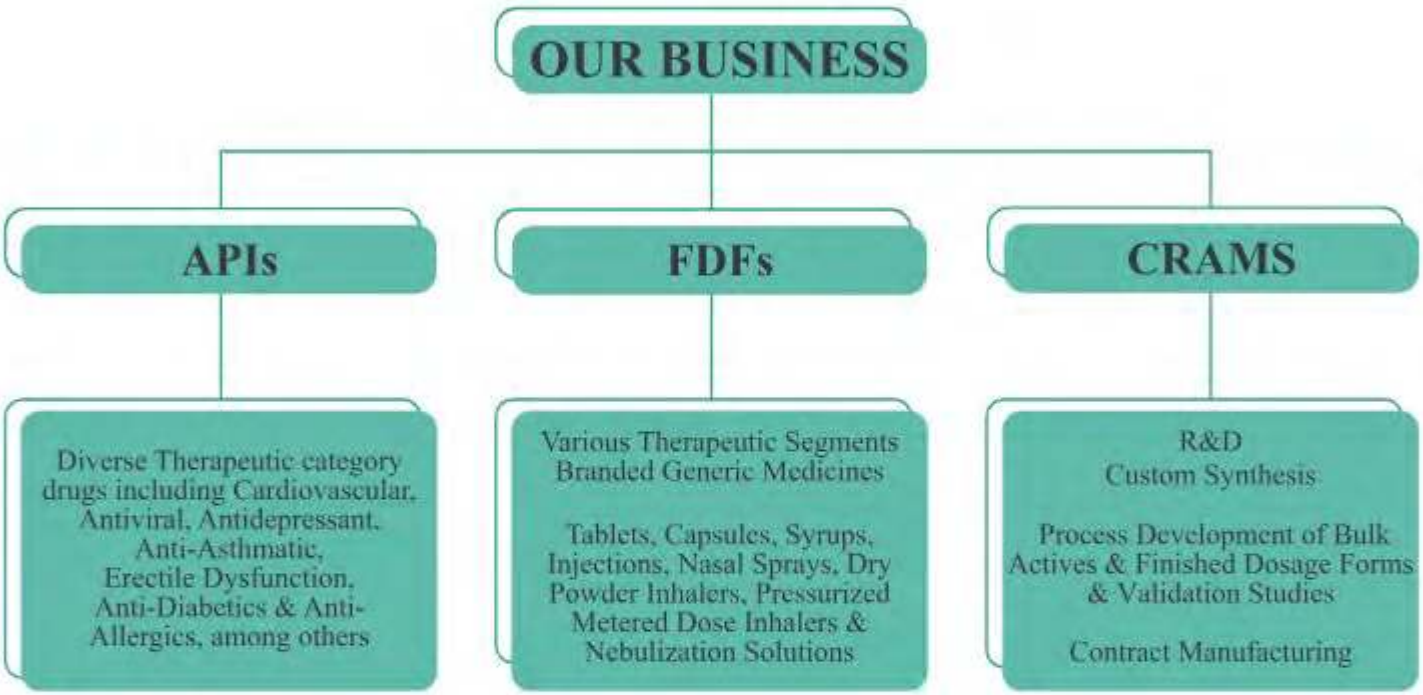
Our Vision

To become a leading global pharmaceutical organization, driven by expertise, pioneering research, and advanced technology, delivering innovative products for a healthier world.

"Quality Healthcare Medicines"

Our Business

Ultratech's core operations are centered around three key segments: large-scale production of Active Pharmaceutical Ingredients, manufacturing Finished Dosage Forms, and offering Custom Research & Manufacturing Services.



"Precision Manufacturing, Ensuring Quality"

Finished Dosage Forms

Ultratech's Finished Dosage Forms are manufactured under stringent quality control standards at our EU-GMP and WHO-GMP approved facilities in India. We provide a comprehensive portfolio of Finished Dosage Forms in conventional dosage forms such as tablets, capsules & injections as well as in advanced dosage forms of Dry Powder Inhalers, Pressurized Metered Dose Inhalers and Nebulization solutions for Pulmonary Drug Delivery.

"We understand our customers' needs and deliver beyond expectations!"

Active Pharmaceutical Ingredients & Intermediates

Ultratech has state-of-the-art manufacturing facilities for the production of Bulk Active Pharmaceutical Ingredients and their Intermediates, addressing a wide array of therapeutic categories. Our advanced infrastructure is designed for flexibility in product mix, enabling increased production volumes, and rapid reconfiguration to align our processes with market demands.

"Backward integration and innovation in complex chemical synthesis - Our obsession!"

Custom Research And Manufacturing Services (CRAMS)

Ultratech provides a comprehensive package that encompasses the entire process - from the development of drug intermediates to the creation of Active Pharmaceutical Ingredients, and the formulation of finished dosage forms, along with commercial manufacturing and packaging. Our dedicated R&D team is committed to optimizing cost efficiency through the adoption of innovative methodologies, process simplification, standardization, and advanced technological enhancements. With a commitment to continuous improvement, our experienced scientists and manufacturing chemists diligently strive to deliver exceptional solutions tailored to meet our customers' needs.

We offer a wide range of services, including:

- Product Development
- Process Development for Pharmaceutical Dosage Forms, APIs, and Drug Intermediates
- Formulation Development Across Various Dosage Forms
- Synthesis Route Development and Optimization
- Impurity Synthesis and Identification
- Development of Environmentally Sustainable Processes
- Analytical Method Development and Validation
- Technology Transfer
- Intellectual Property Rights (IPR) Management
- Stability Studies
- Dossier Development
- Large-Scale Commercial Manufacturing

"Ultratech is a comprehensive result driven partner for clients seeking expertise in Formulations, APIs, and Contract Manufacturing"

Our Niche

- **Pressurized Metered Dose Inhalers (pMDIs)**
- **Nasal Sprays**
- **Dry Powder Inhalers**

Ultratech specializes in the production of a comprehensive range of Nasal Sprays, HFA-based pMDIs, and Dry Powder Inhalers. Our innovative formulations meet a variety of therapeutic needs, encompassing applications such as asthma management, allergic rhinitis, anginal pain relief, sore throat relief, burn treatment, and more.

Ultratech's foray into pMDIs .... A strategic initiative!

Research & Development

Ultratech's R&D unit is accredited by the Department of Scientific and Industrial Research, Government of India. This state-of-the-art facility features dedicated Synthetic Chemistry Laboratories for gram, kilo, and pilot scale development, as well as Formulation Development and Analytical Method Development sections.

The API R&D team specializes in synthesizing key starting materials, optimizing processes, impurity profiling, and in developing non-infringing processes. The Formulation Development section excels in both conventional dosage forms (tablets, capsules, injectables), and advanced forms (pMDIs, Dry Powder Inhalers), demonstrating expertise in Pulmonary Drug Delivery systems. The company fosters synergistic integration across APIs and FDFs.

Intellectual Property Rights

Ultratech has established a robust IPR portfolio and has been awarded a **US PCT patent** for a revolutionary formulation targeting vaginal conditions and designed for optimal vaginal tightening & rejuvenation.

Furthermore, Ultratech has achieved a groundbreaking milestone in overcoming the critical challenge of providing drug delivery of **Remdesivir to patients via the pulmonary route**. This remarkable innovation has earned the company a **prestigious patent from the Controller General of Patents - India,** for **Remdesivir Inhalation Pulmonary Route Delivery Formulations** specifically indicated for **SARS-CoV-2** and other **broad-spectrum antiviral infections**.

These significant achievements reflect Ultratech's steadfast dedication to pioneering innovative healthcare solutions designed to address the evolving needs of next-generation healthcare.

"Pharmaceutical Innovation with Integrated R&D Excellence"

Quality Control & Quality Assurance

Ultratech's Quality Control and Quality Assurance teams operate independently to ensure stringent monitoring and control of all plant activities in alignment with established processes and procedures. We uphold high-quality performance standards through a qualified and experienced workforce, 21-CFR compliant equipment, meticulously documented protocols, and a comprehensive quality assurance system.

We are committed to total quality management and adherence to latest quality and safety standards.

At Ultratech, we recognize that quality is essential for nurturing customer trust, and our operations are guided by stringent manuals and policies to ensure this commitment.

"Ultratech's Third Eye .... Total Quality Management"

Regulatory Compliance

Our products are supported by comprehensive technical packages that include Drug Master Files, Product Dossiers, Free Sale Certificates, and Certificates of Pharmaceutical Products, among others.

Leveraging our capabilities in high-growth and high-value therapeutic areas, we are building a robust pipeline of products and expediting filings to establish leadership across our target markets.

"Compliance Driven, Quality Assured"

Food & Drugs Administration (Maharashtra State)

GMP CERTIFICATE

ULTRA TECH INDIA LTD(70043) PLOT NO. 24, 25 & 26, NEW CHEMICAL ZONE,, M.I.D.C. TALOJA, MAHARASHTRA, INDIA - 410208, Dist - RAIGAD ZONE 1 is holding valid Drugs Manufacturing License in

Form 25, Licence No. KD/310 , Iss Dt: 07/08/1997, Val Dt: 31/12/2027, Ren Dt: 01/01/2023
Form 28, Licence No. KD/327 , Iss Dt: 07/08/1997, Val Dt: 31/12/2027, Ren Dt: 01/01/2023,

issued by this administration under the provision of DRUGS & COSMETICS ACT 1940 & RULES THERE UNDER. Under the said licenses the firm is permitted to manufacture and sell their products covered under the

Categories of : Inhalers /Ointtho, Bulk Drugs / API, Pellets, Aerosols

The firm has employed competent technical persons in manufacturing and quality control departments. The said firm observes GOOD MANUFACTURING PRACTICES (GMP) in the manufacturing and testing of the said categories of products by and large as laid down in revised Schedule 'M' of the Drugs & Cosmetics Rules 1945.

The manufacturing plant is subject to regular inspection by the Competent Authority under The Act.

This Certificate is issued for : purpose of Drug Authority Of Importing County, Any Permissible Destination (Export)

This Certificate is Valid for a period : 26/06/2024 - 25/06/2025

NARENDRA PURUSHOTTAM SUPE
e-Signed on: 24-06-2024 13:46

N. P. SUPE
Licensing Authority
Food & Drugs Administration
KONKAN Division,Maharashtra State

FDA MAHARASHTRA

This Licence/Certificate is ISSUED. Physical Signature is NOT Required

Division	MFG.ID.No	Type/GMP Certificate	CERTIFICATE No	Issue Dt / Validity Dt
KONKAN (RZ1)	70043	GMP/2024-25/002824	412643	26/06/2024 - 25/06/2025

Certificate of Compliance

ULTRATECH INDIA LIMITED

HACCP
Hazard Analysis & Critical Control Points

Environmental Compliance

Ultratech is committed to the effective treatment of effluents to ensure complete compliance with environmental regulations. Our environmental policy prioritizes the production of high-quality products while maintaining ecological balance and minimizing environmental impact. We adhere to all statutory regulations and ISO 14001 standards to uphold our commitment to sustainability.

"Eco-Conscious Manufacturing:
Quality Meets Responsibility"

Certificate of Registration

ULTRATECH INDIA LIMITED

ISO 14001 : 2015
Environmental Management Systems

Certificate of Registration

ULTRATECH INDIA LIMITED

ISO 9001 : 2015
Quality Management Systems

Ultratech: Your strategic partner in Formulations, APIs, & Drug Intermediates

- Comprehensive Development and Manufacturing of Formulations, APIs, & Drug Intermediates
- Expertise in Innovative Dosage Forms
- Strategic Marketing Partnerships
- Proven Expertise from R&D to Large-Scale Production

"Committed to Quality, Dedicated to Health."

Sales Contacts:

- **APIs Business**

  Asia Pacific, Australia, North America, Latin & South America
  Email: sales2@ultratechindia.com

  Europe, Africa, MENA Region, Russia & CIS Countries
  Email: sales3@ultratechindia.com

  India
  Email: sales@ultratechindia.com

- **FDF Business**

  fdf@ultratechindia.com

- **Overall Business Inquiries**

  sales@ultratechindia.com

Websites:

www.ultratechindia.com
www.18again.com
www.femininewonder.com

Company Profile

SCAN ME

ULTRATECH INDIA LIMITED

Regd. Off.: 119, J.K. Chambers, Sector-17, Vashi, Navi Mumbai - 400 703, India.
Plant Address : Plot No's : 24, 25 & 26, M.I.D.C. Taloja, Tal. Panvel,
Dist. Raigad-410 208, Maharashtra, India Tel : +91-22 - 66444000 / 2789 2736 / 2789 1443
Email : info@ultratechindia.com Website : www.ultratechindia.com