

# **Urokinase API and Finished Dosage Forms Market Size and Forecast (2021 - 2031), Global and Regional Share, Trend, and Growth Opportunity Analysis Report Coverage: By Product Type (Active Pharmaceutical Ingredient and Finished Dosage Forms), Manufacturing Process (Urine-derived, Cell Culture-based, and Recombinant Technology), Indication (Deep Vein Thrombosis (DVT) and Pulmonary Embolism (PE), Catheter Occlusion, Myocardial Infarction (Heart Attack), Peripheral Arterial Occlusive Diseases, and Others), Distribution Channel (Hospital Pharmacies, Retail and Drug Stores, Direct Sales (API and FDF), and Online Pharmacies), and Geography (North America, Europe, Asia Pacific, Middle East and Africa, and South and Central America)**

<https://marketpublishers.com/r/U04B7C155631EN.html>

Date: October 2025

Pages: 372

Price: US\$ 4,450.00 (Single User License)

ID: U04B7C155631EN

## **Abstracts**

The urokinase API and finished dosage forms market size is expected to grow from US\$ 2.37 billion in 2024 to US\$ 3.60 billion by 2031; it is projected to register a CAGR of 6.3% during 2025-2031. The rising occurrence of thromboembolic disorders, surging geriatric population, and advancements in manufacturing techniques to improve yield are noteworthy factors contributing to the expansion of the urokinase API and finished

dosage forms market size. However, the side effects associated with urokinase and high development and production costs hinder the urokinase API and finished dosage forms market.

The urokinase is getting better with the help of biosimilar development and recombinant API technologies. This kind of progress makes the product more affordable, gives higher efficacy, and is more accessible, especially in areas with fewer resources. The pharmaceutical industry sees the rise of one important phenomenon. It is the rapid expansion of biosimilars and recombinant APIs. The main reason for this trend is the request for low-cost, high-quality biologics along with the technical solutions that lead to a biosimilar-like production. Biosimilars provide an inexpensive method of producing biologics already on the market. They do this by providing the same effectiveness but at a lower cost. A biosimilar urokinase product that has the same effectiveness and safety as the reference product but costs less is a good option for healthcare providers and payers everywhere. Thus, with the end of the exclusivity period, biosimilars become more aggressive competitors, leading to lower prices and widening of access.

The APIs source and manufacturing methods are being replaced by new technologies, such as the use of recombinant DNA technology in place of traditional extraction methods. In the past, urokinase would be obtained by complicated surgical procedures on the human body, while producing recombinant urokinase using DNA technology has made it easier, as it is pure and has the same consistency as the original. Low molecular weight urokinase, which is more effective and safer, is made using the recombinant method to ensure the product is of the same quality and quantity. The progress of nanoparticle-delivery platforms and ultrasound-assisted catheter-directed delivery is one of the reasons why the urokinase stability has been greatly enhanced and the chances of bleeding have been reduced, thereby, clinical outcomes have been improved. The proportion of recombinant production has increased (40-45% hybrid models), which has led to improved homogeneity and the resolution of ethical concerns about urine sourcing. The use of recombinant urokinase APIs in the production of different dosage forms, such as injectable thrombolytic agents, has been instrumental in the increase in the safety and the decrease of the immunogenic risks that are always associated with plasma-derived products. As healthcare systems become more and more efficient at getting the most value out of every penny, biosimilars and recombinant APIs will be at the forefront of providing patient access to cost-saving and life-saving thrombolytic therapies. Thus, biosimilars and recombinant API growth are expected to emerge as a significant trend in the urokinase API and finished dosage forms market.

## Distribution Channel -Based Insights

Based on distribution channel, the urokinase API and finished dosage forms market is segmented into hospital pharmacies, retail and drug stores, direct sales (API and FDF), and online pharmacies. The hospital pharmacies segment held the largest urokinase API and finished dosage forms market share in 2024. Urokinase is given intravenously in a hospital setting and is usually a product of a closed system. Crystal urokinase, formed into powder and liquid solutions for emergency administration in cases like cardiovascular surgery or thrombolytic therapy, is a regular supply in hospital pharmacies. The prominence of this segment is also due to the hospital's advanced infrastructures, trained staff, and hospitals' capability to manage the complex storage and handling requirements of urokinase formulations, especially solutions that need to be dosed accurately to avoid errors. Apollo Hospitals in India, one of the leading healthcare providers in the APAC region, which is heavily dependent on hospital pharmacies for managing the supply chain and ensuring delivery on time, has incorporated urokinase as one of the treatment options for pulmonary embolism. Going forward, the hospital pharmacies will continue to play an indispensable role in the distribution and administration of urokinase to patients as the healthcare systems keep on changing, thereby fueling the urokinase API and finished dosage forms market growth.

The World Health Organization and the Centers for Disease Control and Prevention are among the primary and secondary sources referred to while preparing the urokinase API and finished dosage forms market report.

### **Reason to buy**

Save and reduce time carrying out entry-level research by identifying the growth, size, leading players and segments in the global Urokinase API and Finished Dosage Forms market.

Highlights key business priorities in order to assist companies to realign their business strategies

The key findings and recommendations highlight crucial progressive industry trends in the global Urokinase API and Finished Dosage Forms market, thereby allowing players across the value chain to develop effective long-term strategies

Develop/modify business expansion plans by using substantial growth offering developed and emerging markets

Scrutinize in-depth global market trends and outlook coupled with the factors driving the market, as well as those hindering it.

Enhance the decision-making process by understanding the strategies that underpin commercial interest with respect to client products, segmentation, pricing and distribution

## **Companies**

### The List of Companies - Urokinase API and Finished Dosage Forms Market

Taj Pharmaceuticals Ltd

Aetos Pharma Private Limited

Ilex Life Sciences

Jiangxi Haoran Bio-Pharma Co., Ltd.

Microbix Biosystems

Syner-Med (Pharmaceutical Products) Ltd.

Midas Pharma GmbH

Cerbios-Pharma SA

Fresenius Kabi AG

Cadila Pharmaceuticals Ltd.

Chandra Bhagat Pharma

medac GmbH.

Kraeber & Co GmbH

TTK Healthcare Limited

Life Medicare & Biotech Private Limited

Unichem Laboratories Ltd.

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