

UMBRALISIB - Emerging Insight and Market Forecast - 2030

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Abstracts

This report can be delivered to the clients within 48 Hours

“UMBRALISIB - Emerging Insight and Market Forecast - 2030” the report provides comprehensive insights about an investigational product for Follicular lymphoma in 7 Major Markets. A detailed picture of the UMBRALISIB in Seven Major Markets, i.e., United States, EU5 (Germany, France, Italy, Spain, and the United Kingdom), and Japan, for the study period 2020 - 2030 is provided in this report along with a detailed description of the product. The product details cover mechanism of action, dosage and administration, route of synthesis, and Research and development activity including regulatory milestones, and other development activities. Further, it also consists of future market assessments inclusive of the market forecast, SWOT analysis, market competitors, and other emerging therapies.

OVERVIEW

UMBRALISIB is oral, dual inhibitor of PI3K-delta and CK1-epsilon, as a treatment for patients with previously treated marginal zone lymphoma (MZL) who have received at least one prior anti-CD20 based regimen and follicular lymphoma (FL) who have received at least two prior systemic therapies. The NDA for umbralisib was based primarily on data from the umbralisib monotherapy MZL and FL cohorts of the UNITY-NHL Phase 2b trial evaluating patients with relapsed/refractory MZL or FL. It is currently in preregistration phase.

SCOPE OF THE REPORT

The report provides insights into:

A comprehensive product overview including the product description, mechanism of action, dosage and administration, Research and Development activity.

Elaborated details on regulatory milestones and other development activities have been provided in this report.

The report also highlights the drug research and development activity details across the United States, Europe and Japan.

The report also covers the patents information with expiry timeline around UMBRALISIB.

The report contains forecasted sales for UMBRALISIB till 2030.

Comprehensive coverage of the late-stage emerging therapies (Phase III) for Follicular lymphoma.

The report also features the SWOT analysis with analyst insights and key findings of UMBRALISIB.

METHODOLOGY

The report is built using data and information sourced primarily from internal databases, primary and secondary research and in-house analysis by DelveInsight's team of industry experts. Information and data from the secondary sources have been obtained from various printable and nonprintable sources like search engines, news websites, global regulatory authorities websites, trade journals, white papers, magazines, books, trade associations, industry associations, industry portals and access to available databases.

UMBRALISIB Analytical Perspective by DelveInsight

In-depth UMBRALISIB Market Assessment

This report provides a detailed market assessment of UMBRALISIB in Seven Major Markets, i.e., United States, EU5 (Germany, France, Italy, Spain, and the United

Kingdom), and Japan. This segment of the report provides forecasted sales data from 2020 to 2030.

UMBRALISIB Clinical Assessment

The report provides the clinical trials information of UMBRALISIB covering trial interventions, trial conditions, trial status, start and completion dates.

REPORT HIGHLIGHTS

In the coming years, the market scenario for Follicular lymphoma is set to change due to the extensive research and incremental healthcare spending across the world; which would expand the size of the market to enable the drug manufacturers to penetrate more into the market.

The companies and academics are working to assess challenges and seek opportunities that could influence UMBRALISIB dominance. The therapies under development are focused on novel approaches to treat/improve the disease condition.

Other emerging products for Follicular lymphoma are giving market competition to UMBRALISIB and launch of late-stage emerging therapies in the near future will significantly impact the market.

A detailed description of regulatory milestones, development activities, and some key findings provide the current development scenario of UMBRALISIB.

Our in-depth analysis of the forecasted sales data of UMBRALISIB from 2020 to 2030 will support the clients in the decision-making process regarding their therapeutic portfolio by identifying the overall scenario of the UMBRALISIB.

Key Questions

Which company is developing UMBRALISIB along with the phase of the clinical study?

What is the technology utilized in the development of UMBRALISIB?

What is the product type, route of administration and mechanism of action of UMBRALISIB?

What is the clinical trial status of the study and study completion date?

What are the key collaborations, mergers and acquisitions, licensing and other activities related to the UMBRALISIB development?

What are the key designations that have been granted to UMBRALISIB?

What is the forecasted market scenario of UMBRALISIB?

What is the history of UMBRALISIB and what is its future?

What is the forecasted sales of UMBRALISIB in the seven major countries, including the United States, Europe (Germany, France, Italy, Spain, and the United Kingdom), and Japan?

What are the other emerging products available and how these are giving competition to UMBRALISIB?

Which are the late-stage emerging therapies under development for the treatment of the PPD?

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