

# Tazverik - Drug Insight and Market Forecast - 2030

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## Abstracts

This report can be delivered to the clients within 48 Hours

### OVERVIEW

“Tazverik - Drug Insight and Market Forecast - 2030” report by DelveInsight outlays comprehensive insights of the product indicated for the treatment of its approved condition. A detailed picture of the Tazverik in Seven Major Markets, i.e., United States, EU5 (Germany, France, Italy, Spain, and the United Kingdom), and Japan, for the study period 2017 -2030 is provided in this report along with a detailed description of the product. The product details covers mechanism of action, dosage and administration, route of synthesis, and pharmacological studies, including product marketed details, regulatory milestones, and other development activities. Further, it also consists of market assessments inclusive of the market forecast, SWOT analysis, and detailed analyst views. It further highlights the market competitors, late-stage emerging therapies, and patent details in the global space.

### Drug Summary

Tazemetostat, sold under the brand name Tazverik is a medication for the treatment of Follicular lymphoma, Sarcoma. It is a small molecule which acts as enhancer of zeste homolog 2 protein inhibitors, SMARCA2 protein inhibitors, SMARCA4 protein inhibitors.

On June 18, 2020, the Food and Drug Administration granted accelerated approval to tazemetostat (TAZVERIK, Epizyme, Inc.), an EZH2 inhibitor, for adult patients with relapsed or refractory (R/R) follicular lymphoma (FL) whose tumors are positive for an EZH2 mutation as detected by an FDA-approved test and who have received at least 2 prior systemic therapies, and for adult patients with R/R FL who have no satisfactory alternative treatment options.

## SCOPE OF THE REPORT

The report provides insights into:

A comprehensive product overview including the product description, mechanism of action, dosage and administration, route of synthesis, pharmacological studies (pharmacodynamics and pharmacokinetics) and adverse reactions.

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

The report also highlights the drug marketed details across the United States, Europe and Japan.

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

The report contains historical and forecasted sales for Tazverik till 2030.

Comprehensive coverage of the late-stage emerging therapies (Phase III) in the space with a brief snapshot of the details.

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

## 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.

Tazverik Analytical Perspective by DelveInsight

## In-depth Tazverik Market Assessment

This report provides a detailed market assessment of Tazverik 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 historical and forecasted sales data from 2017 to 2030.

## Tazverik Clinical Assessment

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

## Report highlights

In the coming years, the market scenario for Tazverik is set to change due to the extensive research in the treatment of the indicated condition 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 Tazverik dominance. The therapies under development are focused on novel approaches to treat/improve the disease condition.

Other approved products for the disease are giving market competition to Tazverik 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 market scenario of Tazverik.

Our in-depth analysis of the sales data of Tazverik from 2017 to 2030 will support the clients in the decision-making process regarding their therapeutic portfolio by identifying the overall scenario of the Tazverik in the market.

## Key Questions

What is the prescribed dosage and strengths of Tazverik are available in the market?

What are the common adverse reactions or side effects of Tazverik?

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

What are the chemical specifications of Tazverik?

How are the clinical trials diversified on the basis of the trial status?

What is the history of Tazverik, and what is its future?

What are the marketed details of Tazverik in the seven major countries, including the United States, Europe (Germany, France, Italy, Spain, and the United Kingdom), and Japan?

How many patents have been granted to Tazverik and when these patents will get expire?

What are the pros (benefits) and cons (disadvantages) of Tazverik?

In which countries Tazverik got approval and when it gets launched?

What are the clinical trials are currently ongoing for Tazverik?

How the safety and efficacy results determined the approval of Tazverik?

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

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

What is the historical and forecasted market scenario of Tazverik?

How is the market trend of Tazverik is different in the Seven Major Markets (the United States, EU5 [Germany, France, Italy, Spain, and the United Kingdom], and Japan)?

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