

MB-102 - Emerging Insight and Market Forecast - 2030

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Abstracts

This report can be delivered to the clients within 48 Hours

“MB-102 - Emerging Insight and Market Forecast - 2030” the report provides comprehensive insights about an investigational product for Blastic Plasmacytoid Dendritic Cell Neoplasm in 7 Major Markets. A detailed picture of the MB-102 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

MB-102 (CD123 CAR T) is a CAR T cell therapy that is produced by engineering T cells to recognize and eliminate CD123-expressing tumors. CD123 is widely expressed on bone marrow cells of patients with myelodysplastic syndromes, as well as in hematologic malignancies including AML, B-cell acute lymphoblastic leukemia, hairy cell leukemia, BPDCN, chronic myeloid leukemia and Hodgkin's lymphoma. The U.S. Food and Drug Administration (FDA) has granted Orphan Drug Designation to MB-102 (CD123 CAR T) for the treatment of blastic plasmacytoid dendritic cell neoplasm (BPDCN), a rare and incurable blood cancer.

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 an expiry timeline around MB-102.

? The report contains forecasted sales for MB-102 till 2030.

? Comprehensive coverage of the Early-stage emerging therapies (Phase I/II) for Blastic Plasmacytoid Dendritic Cell Neoplasm.

? The report also features the SWOT analysis with analyst insights and key findings of MB-102.

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.

MB-102 Analytical Perspective by DelveInsight

? In-depth MB-102 Market Assessment

This report provides a detailed market assessment of MB-102 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.

? MB-102 Clinical Assessment

The report provides the clinical trials information of MB-102 covering trial interventions, trial conditions, trial status, start and completion dates.

REPORT HIGHLIGHTS

? In the coming years, the market scenario for Blastic Plasmacytoid Dendritic Cell Neoplasm 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 MB-102 dominance. The therapies under development are focused on novel approaches to treat/improve the disease condition.

? Other emerging products for Blastic Plasmacytoid Dendritic Cell Neoplasm are giving market competition to MB-102 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 MB-102.

? Our in-depth analysis of the forecasted sales data of MB-102 from 2020 to 2030 will support the clients in the decision-making process regarding their therapeutic portfolio by identifying the overall scenario of the MB-102.

Key Questions

Which company is developing MB-102 along with the phase of the clinical study?

What is the technology utilized in the development of MB-102?

What is the product type, route of administration and mechanism of action of MB-102?

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 MB-102 development?

What are the key designations that have been granted to MB-102?

What is the forecasted market scenario of MB-102?

What is the history of MB-102 and what is its future?

What is the forecasted sales of MB-102 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 MB-102?

Which are the late-stage emerging therapies under development for the treatment of the Blastic Plasmacytoid Cells Neoplasm?

Contents

1. DRUG OVERVIEW

- 1.1. Product Detail
- 1.2. Mechanism of Action
- 1.3. Dosage and Administration
- 1.4. Research and development activity
 - 1.4.1. Clinical Development
 - 1.4.2. Safety and Efficacy
- 1.5. Other Development Activities

2. MARKET ASSESSMENT

- 2.1. 7MM Market Analysis
- 2.2. The United States Market
- 2.3. Germany Market
- 2.4. France Market
- 2.5. Italy Market
- 2.6. Spain Market
- 2.7. United Kingdom Market
- 2.8. Japan Market

3. SWOT ANALYSIS

4. ANALYST VIEWS

5. MARKET COMPETITORS

6. OTHER EMERGING THERAPIES

7. APPENDIX

8. REPORT PURCHASE OPTIONS

List Of Tables

LIST OF TABLES

Table 1 MB-102, Description

Table 2 MB-102, Clinical Trial Description

Table 3 MB-102, 7MM Market Size from 2020 to 2030 (in Million USD)

Table 4 Market Competitors

Table 5 Other Emerging Therapies

List Of Figures

LIST OF FIGURES

Figure 1 The Development Timeline of MB-102

Figure 2 Patent Details, MB-102

Figure 3 MB-102, 7MM Market Size from 2020 to 2030 (in Million USD)

Figure 4 MB-102, US Market Size from 2020 to 2030 (in Millions USD)

Figure 5 MB-102, EU5 Market Size from 2020 to 2030 (in Millions USD)

Figure 6 MB-102, Japan Market Size from 2020 to 2030 (in Millions USD)

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