

# **BLU 782- Emerging Drug Insight and Market Forecast – 2030**

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## **Abstracts**

**This report can be delivered to the clients within 48 Hours**

“BLU 782- Emerging Drug Insight and Market Forecast – 2030” the report provides comprehensive insights about an investigational product for Fibrodysplasia ossificans progressiva (FOP) in 7 Major Markets. A detailed picture of the BLU 782 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.

### **Drug Summary**

BLU-782 was designed by Blueprint Medicines to selectively target mutant ALK2, the underlying cause of FOP, using Blueprint Medicines’ proprietary scientific platform. Blueprint Medicines recently completed dosing in a Phase 1 clinical trial of BLU-782 in healthy volunteers and reported preliminary data at the American Society of Bone and Mineral Research Annual Meeting in September 2019, which showed that BLU-782 was well-tolerated at all doses tested. Previously reported preclinical data in a well-characterized, genetically accurate FOP model showed that BLU-782 prevented injury- and surgery-induced heterotopic ossification, reduced edema and restored healthy tissue response to muscle injury. The FDA has granted a rare pediatric disease designation, orphan drug designation and fast track designation to BLU-782.

## 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 BLU 782.

The report contains forecasted sales for BLU 782 till 2030.

Comprehensive coverage of the late-stage emerging therapies (Phase III) for Fibrodysplasia ossificans progressiva (FOP).

The report also features the SWOT analysis with analyst insights and key findings of BLU 782.

## 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.

## BLU 782 Analytical Perspective by DelveInsight

### In-depth BLU 782 Market Assessment

This report provides a detailed market assessment of BLU 782 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.

### BLU 782 Clinical Assessment

The report provides the clinical trials information of BLU 782 covering trial interventions, trial conditions, trial status, start and completion dates.

### Report highlights

In the coming years, the market scenario for Fibrodysplasia ossificans progressiva (FOP) 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 BLU 782 dominance. The therapies under development are focused on novel approaches to treat/improve the disease condition.

Other emerging products for Fibrodysplasia ossificans progressiva (FOP) are giving market competition to BLU 782 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 BLU 782.

Our in-depth analysis of the forecasted sales data of BLU 782 from 2020 to 2030 will support the clients in the decision-making process regarding their therapeutic portfolio by identifying the overall scenario of the BLU 782.

### Key Questions

Which company is developing BLU 782 along with the phase of the clinical study?

What is the technology utilized in the development of BLU 782?

What is the product type, route of administration and mechanism of action of BLU 782?

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 BLU 782 development?

What are the key designations that have been granted to BLU 782?

What is the forecasted market scenario of BLU 782?

What is the history of BLU 782 and what is its future?

What is the forecasted sales of BLU 782 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 BLU 782?

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