

# Fabry Disease Market Analysis 2025 – Insights For Long-Term Investment & Planning

*The Business Research Company's Fabry Disease Global Market Report 2025 – Market Size, Trends, And Global Forecast 2025-2034*

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Fabry Disease Global Market Report 2025

As per the data, the [fabry disease market](#) size has witnessed strong growth in recent years. It will grow from \$2.10 billion in 2024 to \$2.28 billion in 2025 at a compound annual growth rate CAGR

of 8.2%. Factors such as increased awareness and diagnosis of rare diseases, improvement in specialized healthcare infrastructure, government incentives and orphan drug designations, as well as increased funding for rare disease research and initial launches of therapies are responsible for this growth.

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What Is The Projected Future Growth Of The Fabry Disease Market Size?

Looking ahead, the fabry disease market size is poised for vigorous growth in the coming years. The market is

anticipated to reach \$3.10 billion in 2029, displaying a CAGR of 8.0%. Expansion of newborn screening programs, regulatory support for orphan and rare disease drugs, expansion of healthcare access, growth of telemedicine and remote care, and increasing demand for personalized medicine will dominant drivers in the forecast period. Noteworthy trends include adoption of enzyme replacement therapy, advancement in genetic testing, the use of AI and digital tools for early detection, the development of AI-powered diagnostic algorithms, and advancements in gene therapy.

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## What Are The Key Drivers Propelling The Growth Of The Fabry Disease Market?

A key growth driver propelling the fabry disease market's growth is the rising focus on personalized medicine. This approach uses information about a person's genes, environment, and lifestyle to tailor medical care and therapies specifically for better outcomes. It is anticipated to help improve treatment effectiveness and minimize side effects by tailoring therapies to individual patient needs. For example, in February 2024, the Personalized Medicine Coalition, a US-based non-profit organization, reported that the FDA approved 16 new personalized treatments for rare disease patients, significantly up from the six approvals in 2022.

## What Key Player Strategies Are Driving The Fabry Disease Market?

Major companies in the fabry disease market include Sanofi S.A., GlaxoSmithKline plc, Takeda Pharmaceutical Company Limited, JCR Pharmaceuticals Co. Ltd., Sumitomo Pharma Co. Ltd, Chiesi Farmaceutici S.p.A., Amicus Therapeutics Inc., Idorsia Pharmaceuticals Ltd, Sangamo Therapeutics Inc., uniQure N.V., Protalix BioTherapeutics Inc., Spur Therapeutics Inc., Exegenesis Bio Inc., ISU ABXIS Co. Ltd., Eleva GmbH, AceLink Therapeutics Inc., 4D Molecular Therapeutics Inc., iBio Inc., Freeline Therapeutics Holdings plc, Greenovation Biotech GmbH. These industry players are focusing on developing innovative therapies, such as PEGylated enzyme replacement therapy ERT to enhance treatment efficacy, improve patient compliance, and reduce immunogenic responses associated with conventional ERTs.

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## How Is The [Fabry Disease Market Segmented](#)?

Regarding market segmentation, the fabry disease market report covers the market by Type Type 1, Type 2, Other Types, Treatment Enzyme Replacement Therapy, Oral Therapy, Adjunct Therapy, Other Treatments, Diagnosis Blood Test, Genetic Test, Parenteral Test, Other Diagnosis, Distribution Channel Hospital Pharmacy, Online Pharmacy, Retail Pharmacy, and End User Hospitals, Homecare, Specialty Clinics, Other End Users.

## What Are The Regional Insights In The Fabry Disease Market?

With regards to geography, North America was the largest region in the fabry disease market in 2024. However, Asia-Pacific is anticipated to record the fastest growth in the forecast period.

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