

Global Genetic Screening for Carrier Diseases Market Growth (Status and Outlook) 2026-2032

<https://marketpublishers.com/r/G6DB2B22329FEN.html>

Date: May 2026

Pages: 162

Price: US\$ 3,660.00 (Single User License)

ID: G6DB2B22329FEN

Abstracts

The global Genetic Screening for Carrier Diseases market size is predicted to grow from US\$ 2557 million in 2025 to US\$ 4799 million in 2032; it is expected to grow at a CAGR of 9.6% from 2026 to 2032.

Genetic Screening for Carrier Diseases refers to detecting whether an individual carries the causative gene of a certain genetic disease. In human genetics, some diseases are caused by recessively inherited gene mutations. These mutations usually do not cause the disease in the individual themselves, but if two carriers have offspring, they may increase the risk of the offspring suffering from the genetic disease. Genetic screening for genetic disease carriers can help individuals, especially couples planning to have children, by helping to assess the risk of their future offspring developing specific genetic diseases. If both partners are carriers, they are more likely to have offspring with the disease. Through this screening, couples can better understand their genetic status, make more informed reproductive decisions, and seek genetic counseling and medical intervention when needed.

The genetic screening for carrier diseases market is transitioning from a niche, risk-based offering into a more standardized element of reproductive care as clinical practice moves toward earlier identification of inherited risk and simpler pan-ethnic approaches. Growth is supported by declining sequencing costs, expanding awareness among patients and clinicians, and the operational convenience of comprehensive panels that reduce reliance on ancestry assumptions. Competitive differentiation increasingly depends on the quality and consistency of variant interpretation, the clarity of reporting for non-specialist providers, turnaround time, and the ability to embed testing into fertility, obstetrics, and primary care workflows with counseling support and digital ordering. At the same time, the market faces reimbursement and pricing pressure,

which favors scaled laboratory operations, efficient logistics, and evidence generation that links screening to measurable clinical utility and downstream cost avoidance.

LPI (LP Information)' newest research report, the “Genetic Screening for Carrier Diseases Industry Forecast” looks at past sales and reviews total world Genetic Screening for Carrier Diseases sales in 2025, providing a comprehensive analysis by region and market sector of projected Genetic Screening for Carrier Diseases sales for 2026 through 2032. With Genetic Screening for Carrier Diseases sales broken down by region, market sector and sub-sector, this report provides a detailed analysis in US\$ millions of the world Genetic Screening for Carrier Diseases industry.

This Insight Report provides a comprehensive analysis of the global Genetic Screening for Carrier Diseases landscape and highlights key trends related to product segmentation, company formation, revenue, and market share, latest development, and M&A activity. This report also analyses the strategies of leading global companies with a focus on Genetic Screening for Carrier Diseases portfolios and capabilities, market entry strategies, market positions, and geographic footprints, to better understand these firms' unique position in an accelerating global Genetic Screening for Carrier Diseases market.

This Insight Report evaluates the key market trends, drivers, and affecting factors shaping the global outlook for Genetic Screening for Carrier Diseases and breaks down the forecast by Type, by Application, geography, and market size to highlight emerging pockets of opportunity. With a transparent methodology based on hundreds of bottom-up qualitative and quantitative market inputs, this study forecast offers a highly nuanced view of the current state and future trajectory in the global Genetic Screening for Carrier Diseases.

This report presents a comprehensive overview, market shares, and growth opportunities of Genetic Screening for Carrier Diseases market by product type, application, key players and key regions and countries.

Segmentation by Type:

NGS

Whole Exome Sequencing

Others

Segmentation by Screening Timing:

Preconception Screening

Prenatal Screening

Segmentation by Screening Scope:

Single-condition Screening

Targeted multi-condition Screening

Segmentation by Application:

Couple

Individual

This report also splits the market by region:

Americas

United States

Canada

Mexico

Brazil

APAC

China

Japan

Korea

Southeast Asia

India

Australia

Europe

Germany

France

UK

Italy

Russia

Middle East & Africa

Egypt

South Africa

Israel

Turkey

GCC Countries

The below companies that are profiled have been selected based on inputs gathered from primary experts and analyzing the company's coverage, product portfolio, its market penetration.

Thermo Fisher Scientific

Eurofins Scientific

Illumina

Labcorp

Quest Diagnostics

MedGenome

Myriad Genetics

Natera

OPKO Health

23andMe

Mayo Clinic Laboratories

Baylor Genetics

GeneDx

Fulgent Genetics

NxGen MDx

Sonic Genetics

Ambry Genetics

PreventionGenetics

DiaSorin

SYNLAB

BGI Genomics

Berry Genomics

Annaroad

Jiajian Medical Testing

Genesky

Weihansi Biomedical Technology

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