

Global Genetic Screening for Carrier Diseases Supply, Demand and Key Producers, 2026-2032

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Abstracts

The global Genetic Screening for Carrier Diseases market size is expected to reach \$ 4895 million by 2032, rising at a market growth of 9.0% CAGR during the forecast period (2026-2032).

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.

This report studies the global Genetic Screening for Carrier Diseases demand, key companies, and key regions.

This report is a detailed and comprehensive analysis of the world market for Genetic Screening for Carrier Diseases, and provides market size (US\$ million) and Year-over-Year (YoY) growth, considering 2025 as the base year. This report explores demand trends and competition, as well as details the characteristics of Genetic Screening for Carrier Diseases that contribute to its increasing demand across many markets.

Highlights and key features of the study

Global Genetic Screening for Carrier Diseases total market, 2021-2032, (USD Million)

Global Genetic Screening for Carrier Diseases total market by region & country, CAGR, 2021-2032, (USD Million)

U.S. VS China: Genetic Screening for Carrier Diseases total market, key domestic companies, and share, (USD Million)

Global Genetic Screening for Carrier Diseases revenue by player, revenue and market share 2021-2026, (USD Million)

Global Genetic Screening for Carrier Diseases total market by Type, CAGR, 2021-2032, (USD Million)

Global Genetic Screening for Carrier Diseases total market by Application, CAGR, 2021-2032, (USD Million)

This report profiles major players in the global Genetic Screening for Carrier Diseases market based on the following parameters - company overview, revenue, gross margin, product portfolio, geographical presence, and key developments. Key companies covered as a part of this study include Thermo Fisher Scientific, Eurofins Scientific, Illumina, Labcorp, Quest Diagnostics, MedGenome, Myriad Genetics, Natera, OPKO Health, 23andMe, etc.

This report also provides key insights about market drivers, restraints, opportunities, new product launches or approvals.

Stakeholders would have ease in decision-making through various strategy matrices used in analyzing the world Genetic Screening for Carrier Diseases market

Detailed Segmentation:

Each section contains quantitative market data including market by value (US\$ Millions), by player, by regions, by Type, and by Application. Data is given for the years 2021-2032 by year with 2025 as the base year, 2026 as the estimate year, and 2027-2032 as the forecast year.

Global Genetic Screening for Carrier Diseases Market, By Region:

United States

China

Europe

Japan

South Korea

ASEAN

India

Rest of World

Global Genetic Screening for Carrier Diseases Market, Segmentation by Type:

NGS

Whole Exome Sequencing

Others

Global Genetic Screening for Carrier Diseases Market, Segmentation by Screening Timing:

Preconception Screening

Prenatal Screening

Global Genetic Screening for Carrier Diseases Market, Segmentation by Screening Scope:

Single-condition Screening

Targeted multi-condition Screening

Global Genetic Screening for Carrier Diseases Market, Segmentation by Application:

Couple

Individual

Companies Profiled:

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

Key Questions Answered

1. How big is the global Genetic Screening for Carrier Diseases market?
2. What is the demand of the global Genetic Screening for Carrier Diseases market?
3. What is the year over year growth of the global Genetic Screening for Carrier

Diseases market?

4. What is the total value of the global Genetic Screening for Carrier Diseases market?
5. Who are the Major Players in the global Genetic Screening for Carrier Diseases market?
6. What are the growth factors driving the market demand?

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