

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

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Abstracts

The global Carrier Screening for Genetic Conditions 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.

Carrier Screening for Genetic Conditions is a genetic testing approach used to determine whether an individual carries a gene variant associated with inherited disorders that may be passed on to offspring. Carriers typically do not show symptoms of the condition but can transmit the altered gene to their children, particularly in autosomal recessive or X-linked inheritance patterns. The screening is commonly performed before or during pregnancy to assess reproductive risk, support informed family planning decisions, and guide further diagnostic or counseling steps when both partners are identified as carriers of the same genetic condition.

United States market for Carrier Screening for Genetic Conditions is estimated to increase from US\$ million in 2025 to US\$ million by 2032, at a CAGR of % from 2026 through 2032.

China market for Carrier Screening for Genetic Conditions is estimated to increase from US\$ million in 2025 to US\$ million by 2032, at a CAGR of % from 2026 through 2032.

Europe market for Carrier Screening for Genetic Conditions is estimated to increase from US\$ million in 2025 to US\$ million by 2032, at a CAGR of % from 2026 through 2032.

Global key Carrier Screening for Genetic Conditions players cover Thermo Fisher Scientific, Eurofins Scientific, Illumina, Labcorp, Quest Diagnostics, etc. In terms of

revenue, the global two largest companies occupied for a share nearly % in 2025.

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

This Insight Report provides a comprehensive analysis of the global Carrier Screening for Genetic Conditions 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 Carrier Screening for Genetic Conditions portfolios and capabilities, market entry strategies, market positions, and geographic footprints, to better understand these firms' unique position in an accelerating global Carrier Screening for Genetic Conditions market.

This Insight Report evaluates the key market trends, drivers, and affecting factors shaping the global outlook for Carrier Screening for Genetic Conditions 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 Carrier Screening for Genetic Conditions.

This report presents a comprehensive overview, market shares, and growth opportunities of Carrier Screening for Genetic Conditions 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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