

Global Hereditary Disease Carrier Screening Market Growth (Status and Outlook) 2026-2032

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Abstracts

The global Hereditary Disease Carrier Screening 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.

Hereditary Disease Carrier Screening is genetic testing used to identify whether an individual carries gene changes associated with inherited disorders that could be passed to a child. It is commonly offered before or during pregnancy to assess reproductive risk, especially for conditions that are inherited in an autosomal recessive or X linked manner. The results help guide next steps such as partner testing and genetic counseling so that individuals and couples can make informed reproductive decisions, while recognizing that a negative result reduces but does not eliminate risk because screening cannot detect every possible disease or genetic variant.

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

Global key Hereditary Disease Carrier Screening 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 “Hereditary Disease Carrier Screening Industry Forecast” looks at past sales and reviews total world Hereditary Disease Carrier Screening sales in 2025, providing a comprehensive analysis by region and market sector of projected Hereditary Disease Carrier Screening sales for 2026 through 2032. With Hereditary Disease Carrier Screening sales broken down by region, market sector and sub-sector, this report provides a detailed analysis in US\$ millions of the world Hereditary Disease Carrier Screening industry.

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

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

This report presents a comprehensive overview, market shares, and growth opportunities of Hereditary Disease Carrier Screening 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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