

Global Newborn Genetic Screening Service Supply, Demand and Key Producers, 2026-2032

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Abstracts

The global Newborn Genetic Screening Service market size is expected to reach \$ 1265 million by 2032, rising at a market growth of 5.0% CAGR during the forecast period (2026-2032).

Newborn Genetic Screening Service is a public health and clinical service that tests babies shortly after birth to identify inherited or congenital conditions that may not be obvious at first but can cause serious harm if untreated. It typically involves collecting a small blood sample and, in some programs, additional hearing or heart checks, then analyzing the sample using laboratory methods that detect metabolic, endocrine, hematologic, and genetic disorders. When a screen suggests risk, the service coordinates confirmatory testing, specialist referral, and rapid initiation of treatment or monitoring, while also providing counseling and follow-up so families understand results and next steps.

The newborn genetic screening service market is characterized by strong policy anchoring, high clinical value, and structurally stable demand driven by universal birth cohorts rather than discretionary utilization. Market growth is shaped less by volume expansion and more by qualitative shifts, including the gradual expansion of screened conditions, the adoption of more advanced molecular methods, and the integration of screening with downstream care coordination and long term follow up. Competitive dynamics favor operators with scale, regulatory credibility, and deep integration into public health systems, creating high entry barriers for new providers while limiting price based competition. At the same time, emerging models such as expanded genomic screening and digital result management introduce opportunities for specialized laboratories and technology partners, suggesting that future differentiation will center on clinical utility, operational reliability, and alignment with public health governance rather

than purely on testing throughput.

This report studies the global Newborn Genetic Screening Service demand, key companies, and key regions.

This report is a detailed and comprehensive analysis of the world market for Newborn Genetic Screening Service, 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 Newborn Genetic Screening Service that contribute to its increasing demand across many markets.

Highlights and key features of the study

Global Newborn Genetic Screening Service total market, 2021-2032, (USD Million)

Global Newborn Genetic Screening Service total market by region & country, CAGR, 2021-2032, (USD Million)

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

Global Newborn Genetic Screening Service revenue by player, revenue and market share 2021-2026, (USD Million)

Global Newborn Genetic Screening Service total market by Type, CAGR, 2021-2032, (USD Million)

Global Newborn Genetic Screening Service total market by Application, CAGR, 2021-2032, (USD Million)

This report profiles major players in the global Newborn Genetic Screening Service 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, Revvity, Quest Diagnostics, MedGenome, Mayo Clinic Laboratories, GeneDx, PreventionGenetics, SYNLAB, Veritas Genetics, ARCHIMEDlife, 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 Newborn Genetic Screening Service 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 Newborn Genetic Screening Service Market, By Region:

United States

China

Europe

Japan

South Korea

ASEAN

India

Rest of World

Global Newborn Genetic Screening Service Market, Segmentation by Type:

NGS

Whole Exome Sequencing

Others

Global Newborn Genetic Screening Service Market, Segmentation by Screening Purpose:

Core Condition Screening

Expanded Condition Screening

Global Newborn Genetic Screening Service Market, Segmentation by Service Model:

Public Health Program

Consumer

Others

Global Newborn Genetic Screening Service Market, Segmentation by Application:

Public Hospitals

Private Hospitals

Others

Companies Profiled:

Thermo Fisher Scientific

Revvity

Quest Diagnostics

MedGenome

Mayo Clinic Laboratories

GeneDx

PreventionGenetics

SYNLAB

Veritas Genetics

ARCHIMEDlife

GGA

Cordlife

Rainbow Genomics

BGI Genomics

Berry Genomics

Annaroad

Jiajian Medical Testing

Genesky

Weihansi Biomedical Technology

Key Questions Answered

1. How big is the global Newborn Genetic Screening Service market?
2. What is the demand of the global Newborn Genetic Screening Service market?
3. What is the year over year growth of the global Newborn Genetic Screening Service market?
4. What is the total value of the global Newborn Genetic Screening Service market?
5. Who are the Major Players in the global Newborn Genetic Screening Service market?
6. What are the growth factors driving the market demand?

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