

### Gene Therapy Platform Market Size, Share & Trends Analysis Report By Platform Type (Viral Vector Platforms, Non-Viral Vector Platforms), By Therapeutic Application, By Delivery Mode, By End Use, By Region, And Segment Forecasts, 2025 - 2030

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#### **Abstracts**

This report can be delivered to the clients within 3 Business Days

Gene Therapy Platform Market Size & Trends

The global gene therapy platform market size was estimated at USD 2.14 billion in 2024 and is projected to grow at a CAGR of 15.2% from 2025 to 2030. The growth is driven by several key drivers, including rapid technological advancements, increasing prevalence of genetic and chronic diseases, growing investments and funding in biopharmaceutical research, and supportive regulatory frameworks facilitating faster approvals.

Innovations in viral vector technologies, gene editing tools including CRISPR, integration of AI and automation in manufacturing are enhancing therapy precision and production efficiency. For instance, in March 2023, Catalent is enhancing its UpTempo AAV platform to speed up the development of gene therapies.

The increasing number of FDA approvals for gene therapies targeting rare and genetic diseases has validated the clinical potential of these platforms, encouraging investment and R&D. For instance, in December 2023[, CRISPR-based therapies, like Exagamglogene autotemcel (Casgevy), received FDA approval for sickle cell disease and beta-thalassemia, marking a significant milestone in gene therapy. These breakthroughs are pushing the field forward, enabling more efficient technology for



various genetic disorders.

The rising prevalence of genetic disorders and chronic diseases worldwide, is fueling demand for gene therapies that target disease at the molecular level. Conditions such as cystic fibrosis, hemophilia, spinal muscular atrophy, and various cancers are increasingly treated with gene therapy approaches that offer potentially curative outcomes. The World Health Organization (WHO) in 2024 projected a sharp increase in cancer cases globally, with over 35 million new cases expected by 2050, up 77% from 2022. This growing patient pool creates urgent demand for advanced treatments, driving investment and innovation in gene therapy technologies to address unmet medical needs.

Moreover, regulatory support for innovative therapies is driving market growth. Regulatory agencies, including the FDA and the European Medicines Agency (EMA), have streamlined approval processes for gene therapies, recognizing their transformative potential. In addition, regulatory incentives such as fast-track designations and orphan drug status further support the development of gene therapies. These measures help reduce development timelines and encourage the introduction of new treatments to address unmet medical needs.

The COVID-19 pandemic significantly impacted the market, disrupting clinical trials, manufacturing, and supply chains. The increased demand for raw materials for COVID-19 vaccine production caused shortages, affecting the availability of essential components for gene therapy manufacturing. Despite these challenges, the pandemic also spurred innovation, with some gene therapy developers adapting their efforts toward COVID-19-related therapeutics and vaccines, highlighting the resilience and potential of the gene therapy sector.

Global Gene Therapy Platform Market Report Segmentation

This report forecasts revenue growth at global, regional, and country levels and provides an analysis of the latest industry trends in each of the sub-segments from 2018 to 2030. For this study, Grand View Research has segmented the global gene therapy platform market report based on platform type, application, delivery method, end use, and region.

Platform Type Outlook (Revenue, USD Million, 2018 - 2030)

Viral Vector Platforms



Adeno-associated Virus (AAV) Lentivirus Retrovirus Adenovirus Herpes Simplex Virus (HSV) Non-Viral Vector Platforms Lipid Nanoparticles (LNPs) Electroporation & Microinjection Platforms Polymer-based Delivery Systems Naked DNA/RNA Delivery Gene Editing Platforms **CRISPR-Cas Systems TALENs ZFNs** Application Outlook (Revenue, USD Million, 2018 - 2030) Oncology Rare Genetic Disorders Cardiovascular Diseases **Neurological Disorders** 



Ophthalmic Diseases

Hematological Disorders (e.g., Hemophilia, Sickle Cell)

Musculoskeletal Disorders

Infectious Diseases (e.g., HIV, COVID-19 adjunct therapies)

Delivery Mode Outlook (Revenue, USD Million, 2018 - 2030)

In Vivo Gene Therapy

Ex Vivo Gene Therapy

Autologous Cell-Based Gene Therapy

Allogeneic Cell-Based Gene Therapy

Others (In-situ Gene therapy)

End Use Outlook (Revenue, USD Million, 2018 - 2030)

Pharmaceutical & Biotechnology Companies

Academic & Research Institutions

Contract Development & Manufacturing Organizations (CDMOs)

Hospitals & Gene Therapy Centers

Regional Outlook (Revenue, USD Million, 2018-2030)

North America

U.S.

Canada

Mexico



	Europe						
		UK					
		Germany					
		France					
		Italy					
		Spain					
		Sweden					
		Denmark					
		Norway					
Asia Pacific		Pacific					
		Japan					
		China					
		India					
		Australia					
		South Korea					
		Thailand					
Latin America		America					
		Brazil					
		Argentina					



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Saudi Arabia

South Africa

UAE

Kuwait

#### **Companies Mentioned**

Lonza

**BioIT** 

Autolomous Ltd

Hypertrust Patient Data Care (Part of Accenture)

**IBM** 

L7 Informatics, Inc.

TrakCel

**IDBS** 

**Novartis** 

SAP SE

**DEEP GENOMICS** 

ElevateBio

Sarepta Therapeutics, Inc.

**CRISPR** Therapeutics

**Precision BioSciences** 

AstraZeneca

Andelyn Biosciences

Renova Therapeutics



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