

Global AAV Vector-Based Gene Therapy Drugs Market Growth (Status and Outlook) 2024-2030

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Abstracts

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AAV vector-based gene therapy drugs are a class of therapeutic agents used in the treatment of genetic disorders and certain diseases. These drugs utilize AAV vectors, which are small, non-enveloped viruses that have a low pathogenic profile and are capable of delivering genetic material into host cells. The genetic material, which can be a functional gene or a therapeutic gene, is inserted into the AAV vector and then introduced into the patient's cells, where it is intended to correct genetic mutations or produce therapeutic proteins. AAV vector-based gene therapy has shown promising results in clinical trials for conditions such as Leber congenital amaurosis, spinal muscular atrophy, and certain retinal diseases.

The global AAV Vector-Based Gene Therapy Drugs market size is projected to grow from US\$ million in 2024 to US\$ million in 2030; it is expected to grow at a CAGR of % from 2024 to 2030.

LPI (LP Information)' newest research report, the "AAV Vector-Based Gene Therapy Drugs Industry Forecast" looks at past sales and reviews total world AAV Vector-Based Gene Therapy Drugs sales in 2022, providing a comprehensive analysis by region and market sector of projected AAV Vector-Based Gene Therapy Drugs sales for 2023 through 2029. With AAV Vector-Based Gene Therapy Drugs sales broken down by region, market sector and sub-sector, this report provides a detailed analysis in US\$ millions of the world AAV Vector-Based Gene Therapy Drugs industry.

This Insight Report provides a comprehensive analysis of the global AAV Vector-Based Gene Therapy Drugs 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 AAV Vector-Based Gene Therapy Drugs portfolios and capabilities, market entry strategies, market positions, and geographic footprints, to better understand these firms' unique position in an accelerating global AAV Vector-Based Gene Therapy Drugs market.

This Insight Report evaluates the key market trends, drivers, and affecting factors shaping the global outlook for AAV Vector-Based Gene Therapy Drugs 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 AAV Vector-Based Gene Therapy Drugs.

United States market for AAV Vector-Based Gene Therapy Drugs is estimated to increase from US\$ million in 2023 to US\$ million by 2030, at a CAGR of % from 2024 through 2030.

China market for AAV Vector-Based Gene Therapy Drugs is estimated to increase from US\$ million in 2023 to US\$ million by 2030, at a CAGR of % from 2024 through 2030.

Europe market for AAV Vector-Based Gene Therapy Drugs is estimated to increase from US\$ million in 2023 to US\$ million by 2030, at a CAGR of % from 2024 through 2030.

Global key AAV Vector-Based Gene Therapy Drugs players cover uniQure, Spark Therapeutics, AveXis (Novartis), PTC Therapeutics, BioMarin, etc. In terms of revenue, the global two largest companies occupied for a share nearly % in 2023.

This report presents a comprehensive overview, market shares, and growth opportunities of AAV Vector-Based Gene Therapy Drugs market by product type, application, key players and key regions and countries.

Segmentation by Type:

AAV2 Vector

AAV5 Vector

Others

Segmentation by Application:

Hospital

Diagnostic and Testing Laboratories

Academic and Research Organizations

Others

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

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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.

uniQure

Spark Therapeutics

AveXis (Novartis)

PTC Therapeutics

BioMarin

Sarepta Therapeutics

Roche

Pfizer

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Figure 77. Turkey AAV Vector-Based Gene Therapy Drugs Market Size 2025-2030 (\$ millions)

Figure 78. GCC Countries AAV Vector-Based Gene Therapy Drugs Market Size 2025-2030 (\$ millions)

Figure 79. Global AAV Vector-Based Gene Therapy Drugs Market Size Market Share Forecast by Type (2025-2030)

Figure 80. Global AAV Vector-Based Gene Therapy Drugs Market Size Market Share Forecast by Application (2025-2030)

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