

# Global AAV Vector-Based Gene Therapy Drugs Market 2024 by Company, Regions, Type and Application, Forecast to 2030

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## Abstracts

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.

According to our (Global Info Research) latest study, the global AAV Vector-Based Gene Therapy Drugs market size was valued at US\$ million in 2023 and is forecast to a readjusted size of USD million by 2030 with a CAGR of % during review period.

This report is a detailed and comprehensive analysis for global AAV Vector-Based Gene Therapy Drugs market. Both quantitative and qualitative analyses are presented by company, by region & country, by Type and by Application. As the market is constantly changing, this report explores the competition, supply and demand trends, as well as key factors that contribute to its changing demands across many markets. Company profiles and product examples of selected competitors, along with market share estimates of some of the selected leaders for the year 2024, are provided.

Key Features:

Global AAV Vector-Based Gene Therapy Drugs market size and forecasts, in

consumption value (\$ Million), 2019-2030

Global AAV Vector-Based Gene Therapy Drugs market size and forecasts by region and country, in consumption value (\$ Million), 2019-2030

Global AAV Vector-Based Gene Therapy Drugs market size and forecasts, by Type and by Application, in consumption value (\$ Million), 2019-2030

Global AAV Vector-Based Gene Therapy Drugs market shares of main players, in revenue (\$ Million), 2019-2024

The Primary Objectives in This Report Are:

To determine the size of the total market opportunity of global and key countries

To assess the growth potential for AAV Vector-Based Gene Therapy Drugs

To forecast future growth in each product and end-use market

To assess competitive factors affecting the marketplace

This report profiles key players in the global AAV Vector-Based Gene Therapy Drugs 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 uniQure, Spark Therapeutics, AveXis (Novartis), PTC Therapeutics, BioMarin, Sarepta Therapeutics, Roche, Pfizer, etc.

This report also provides key insights about market drivers, restraints, opportunities, new product launches or approvals.

Market segmentation

AAV Vector-Based Gene Therapy Drugs market is split by Type and by Application. For the period 2019-2030, the growth among segments provides accurate calculations and forecasts for Consumption Value by Type and by Application. This analysis can help you expand your business by targeting qualified niche markets.

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#### Market segment by Type

AAV2 Vector

AAV5 Vector

Others

#### Market segment by Application

Hospital

Diagnostic and Testing Laboratories

Academic and Research Organizations

Others

#### Market segment by players, this report covers

uniQure

Spark Therapeutics

AveXis (Novartis)

PTC Therapeutics

BioMarin

Sarepta Therapeutics

Roche

Pfizer

Market segment by regions, regional analysis covers

North America (United States, Canada and Mexico)

Europe (Germany, France, UK, Russia, Italy and Rest of Europe)

Asia-Pacific (China, Japan, South Korea, India, Southeast Asia and Rest of Asia-Pacific)

South America (Brazil, Rest of South America)

Middle East & Africa (Turkey, Saudi Arabia, UAE, Rest of Middle East & Africa)

The content of the study subjects, includes a total of 13 chapters:

Chapter 1, to describe AAV Vector-Based Gene Therapy Drugs product scope, market overview, market estimation caveats and base year.

Chapter 2, to profile the top players of AAV Vector-Based Gene Therapy Drugs, with revenue, gross margin, and global market share of AAV Vector-Based Gene Therapy Drugs from 2019 to 2024.

Chapter 3, the AAV Vector-Based Gene Therapy Drugs competitive situation, revenue, and global market share of top players are analyzed emphatically by landscape contrast.

Chapter 4 and 5, to segment the market size by Type and by Application, with consumption value and growth rate by Type, by Application, from 2019 to 2030.

Chapter 6, 7, 8, 9, and 10, to break the market size data at the country level, with revenue and market share for key countries in the world, from 2019 to 2024. and AAV Vector-Based Gene Therapy Drugs market forecast, by regions, by Type and by Application, with consumption value, from 2024 to 2030.

Chapter 11, market dynamics, drivers, restraints, trends, Porters Five Forces analysis.

Chapter 12, the key raw materials and key suppliers, and industry chain of AAV Vector-Based Gene Therapy Drugs.

Chapter 13, to describe AAV Vector-Based Gene Therapy Drugs research findings and conclusion.

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