

Global Gene-based Advanced Therapy Medicinal Market Growth (Status and Outlook) 2023-2029

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Abstracts

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Gene Delivery System or Gene therapy is an experimental technique that uses genes to treat or prevent disease. In the future, this technique may allow doctors to treat a disorder by inserting a gene into a patient's cells instead of using drugs or surgery.

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

This Insight Report provides a comprehensive analysis of the global Gene-based Advanced Therapy Medicinal landscape and highlights key trends related to product segmentation, company formation, revenue, and market share, latest development, and M&A activity. This report also analyzes the strategies of leading global companies with a focus on Gene-based Advanced Therapy Medicinal portfolios and capabilities, market entry strategies, market positions, and geographic footprints, to better understand these firms' unique position in an accelerating global Gene-based Advanced Therapy Medicinal market.

This Insight Report evaluates the key market trends, drivers, and affecting factors shaping the global outlook for Gene-based Advanced Therapy Medicinal 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 Gene-based Advanced Therapy Medicinal.

The global Gene-based Advanced Therapy Medicinal market size is projected to grow from US\$ million in 2022 to US\$ million in 2029; it is expected to grow at a CAGR of % from 2023 to 2029.

United States market for Gene-based Advanced Therapy Medicinal is estimated to increase from US\$ million in 2022 to US\$ million by 2029, at a CAGR of % from 2023 through 2029.

China market for Gene-based Advanced Therapy Medicinal is estimated to increase from US\$ million in 2022 to US\$ million by 2029, at a CAGR of % from 2023 through 2029.

Europe market for Gene-based Advanced Therapy Medicinal is estimated to increase from US\$ million in 2022 to US\$ million by 2029, at a CAGR of % from 2023 through 2029.

Global key Gene-based Advanced Therapy Medicinal players cover Biogen, Novartis, Gilead Sciences, Sarepta Therapeutics, Alnylam Pharmaceuticals, Amgen, Spark Therapeutics, Akcea Therapeutics and Sunway Biotech, etc. In terms of revenue, the global two largest companies occupied for a share nearly % in 2022.

This report presents a comprehensive overview, market shares, and growth opportunities of Gene-based Advanced Therapy Medicinal market by product type, application, key players and key regions and countries.

Market Segmentation:

Segmentation by type

Viral Gene Delivery System

Non-viral Gene Delivery System

Segmentation by application

Neurological Diseases

Cancer

Other

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.

Biogen

Novartis

Gilead Sciences

Sarepta Therapeutics

Alnylam Pharmaceuticals

Amgen

Spark Therapeutics

Akcea Therapeutics

Sunway Biotech

SIBIONO

AnGes

Orchard Therapeutics

Human Stem Cells Institute

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