

North America RNA Therapeutics Market, By Product (Vaccines, Drugs), By Type (mRNA Therapeutics, RNA Interference (RNAi) Therapeutics, Antisense Oligonucleotide (ASO) Therapeutics, Other Therapeutics), By Indication (Infectious Diseases, Rare Genetic Diseases/Hereditary Diseases, Other Indications), By End User (Hospitals and Clinics, Ambulatory Surgical Centers, Others), By Country, Competition, Forecast and Opportunities, 2019-2029F

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Abstracts

North America RNA Therapeutics Market was valued at USD 12.95 billion in 2023 and is anticipated to project robust growth in the forecast period with a CAGR of 5.48% through 2029.

The North America RNA Therapeutics Market is a dynamic and rapidly growing sector in the field of biopharmaceuticals, characterized by the development and commercialization of RNA-based therapies for a wide range of diseases. RNA therapeutics utilize the inherent biological functions of RNA molecules to modulate gene expression, making them a powerful tool for the treatment of genetic, infectious, and chronic diseases. This market has witnessed significant expansion in recent years, driven by advancements in RNA technology, increasing awareness of precision medicine, and the growing demand for innovative treatment options. The primary factors contributing to the growth of the North America RNA therapeutics market is the development of RNA interference (RNAi) and antisense oligonucleotide (ASO) technologies. These technologies have paved the way for the targeted inhibition or activation of specific genes, allowing for the treatment of various genetic disorders, viral

infections, and rare diseases. This has resulted in a surge of interest from pharmaceutical and biotechnology companies, leading to substantial investments in research and development. The market encompasses a wide spectrum of RNA therapeutics, including small interfering RNA (siRNA), microRNA (miRNA), messenger RNA (mRNA), and aptamers. These RNA molecules have been harnessed to design therapies for conditions such as cancer, cardiovascular diseases, neurodegenerative disorders, and rare genetic diseases. Moreover, the market is also witnessing promising applications in vaccines and gene editing, contributing to its overall growth. The North America RNA therapeutics market is also characterized by collaborations and partnerships between pharmaceutical companies, biotech firms, and academic institutions, which facilitate the sharing of knowledge, resources, and expertise. Additionally, regulatory agencies such as the FDA and EMA are actively engaged in streamlining the approval process for RNA-based therapies, which further supports market growth.

Key Market Drivers

Advancements in RNA Technology

Advancements in RNA technology have played a pivotal role in boosting the North America RNA therapeutics market, revolutionizing the field of biopharmaceuticals and driving innovation in medical treatments. These technological breakthroughs have significantly enhanced the development and efficacy of RNA-based therapies, propelling them to the forefront of the healthcare industry. The most significant advancements in RNA technology is the refinement of RNA interference (RNAi) and antisense oligonucleotide (ASO) technologies. These techniques enable the precise targeting of specific genes, allowing for the modulation or silencing of gene expression, which is critical in treating genetic, infectious, and chronic diseases. Researchers have made substantial progress in optimizing the design and delivery of siRNAs and ASOs, overcoming challenges related to stability, cellular uptake, and off-target effects. This progress has made it possible to develop more effective and targeted RNA-based therapies, expanding their potential applications.

In addition to improved gene modulation, advancements in RNA technology have also led to the development of innovative delivery systems. Effective delivery is crucial for the successful application of RNA therapeutics, as these molecules can be easily degraded or fail to enter target cells. The development of lipid nanoparticles, modified RNA structures, and other delivery methods has greatly enhanced the stability and bioavailability of RNA therapeutics, making them more practical and feasible for clinical

use. The emergence of messenger RNA (mRNA) vaccines, exemplified by the COVID-19 vaccines developed by Pfizer and Moderna, has garnered North America attention and underscored the potential of RNA technology. These mRNA vaccines demonstrated unprecedented efficacy, safety, and rapid development, leading to their widespread adoption. The success of these vaccines has not only accelerated research and investment in mRNA-based therapies but has also encouraged the development of novel delivery systems and the optimization of RNA stability, which will have a positive ripple effect on the broader RNA therapeutics market.

Rising Prevalence of Genetic Disorders

The rising prevalence of genetic disorders has emerged as a significant driver behind the remarkable growth of the North America RNA therapeutics market. Genetic disorders encompass a diverse spectrum of conditions, from rare and orphan diseases to more common disorders with a genetic component. Traditional pharmaceuticals often fall short in treating these conditions effectively, making RNA-based therapies an attractive and promising alternative. Genetic disorders, such as cystic fibrosis, muscular dystrophy, Huntington's disease, and many more, result from mutations in specific genes that lead to debilitating and life-threatening symptoms. The ability of RNA-based therapies to target and correct these genetic mutations or modulate gene expression offers newfound hope to individuals and their families affected by these conditions. RNA therapeutics enable a precise and personalized approach to treatment, addressing the root cause of the disorder rather than merely managing symptoms.

As genetic testing becomes more accessible and affordable, the identification and diagnosis of genetic disorders are on the rise. This increased understanding of an individual's genetic makeup has further driven the demand for tailored treatments, aligning perfectly with the capabilities of RNA-based therapies. The prevalence of genetic disorders is not limited to rare conditions. Even more common diseases like certain forms of breast cancer, Alzheimer's disease, and various cardiovascular disorders have strong genetic components. RNA therapeutics, such as antisense oligonucleotides (ASOs) and small interfering RNA (siRNA), have shown great potential in targeting specific genes or regulating gene expression to mitigate the impact of these diseases. The growth of the North America RNA therapeutics market is closely intertwined with the evolution of personalized medicine, where treatments are custom-tailored to the genetic makeup and specific disease characteristics of each patient. This approach not only enhances treatment efficacy but also reduces adverse effects, making it an appealing option for healthcare providers, patients, and payers alike.

Expanding Applications in Oncology

The expanding applications of RNA therapeutics in oncology have played a pivotal role in boosting the North America RNA therapeutics market. Oncology has remained a focal point of innovation within the biopharmaceutical industry, and the precision and versatility of RNA-based therapies have propelled significant advancements in cancer treatment. One of the most noteworthy developments in this field is the ability to selectively target and modulate genes associated with cancer. RNA therapeutics, such as small interfering RNA (siRNA) and antisense oligonucleotides (ASOs), offer a tailored approach to silencing or regulating cancer-related genes, effectively inhibiting tumor growth and progression. This precision in gene manipulation has led to the development of innovative therapies for various types of cancer. The messenger RNA (mRNA) vaccines have emerged as a breakthrough in cancer immunotherapy. These vaccines encode specific antigens or tumor markers, directing the immune system to recognize and attack cancer cells. The success of mRNA vaccines in clinical trials and real-world applications, such as the Pfizer-BioNTech and Moderna COVID-19 vaccines, has inspired research into their potential use for cancer immunotherapy. mRNA-based immunotherapies show promise in training the immune system to target specific cancer cells, offering a novel approach to oncology treatment.

MicroRNA (miRNA)-based therapies have also come to the forefront in oncology research. MiRNAs are small non-coding RNA molecules that play a significant role in gene regulation. Dysregulation of miRNA expression is a common occurrence in cancer, contributing to the development and progression of the disease. RNA therapeutics targeting miRNAs aim to restore their normal function, controlling the activity of oncogenes and tumor suppressor genes. This approach holds potential in halting tumor growth and preventing metastasis. The expanding applications of RNA therapeutics in oncology are reshaping the landscape of cancer treatment by offering highly targeted and precise therapies. The potential to silence or modulate cancer-associated genes, harness the power of mRNA vaccines, and regulate miRNA expression has ushered in a new era of innovation in the fight against cancer.

Key Market Challenges

Delivery Challenges

The North America RNA therapeutics market holds immense potential for revolutionizing the treatment of various diseases, but it is not without its share of challenges. Among the most critical obstacles facing the field are delivery challenges.

RNA molecules, which include small interfering RNA (siRNA), messenger RNA (mRNA), antisense oligonucleotides (ASOs), and microRNA (miRNA), need to be efficiently and precisely delivered to their intended targets within the body. Overcoming these delivery challenges is crucial for the successful development and commercialization of RNA therapeutics. The primary concerns in RNA therapeutics delivery is the stability and protection of the RNA molecules as they traverse the complex physiological environment of the human body. Naked RNA is susceptible to degradation by enzymes and can be cleared rapidly from the bloodstream, limiting their therapeutic effectiveness. Researchers have been actively working on developing advanced delivery systems, including lipid nanoparticles, viral vectors, and synthetic carriers, to protect RNA molecules and ensure their safe transport to the target cells or tissues.

Selectivity is another significant issue in RNA therapeutics delivery. Ensuring that RNA therapies reach the intended targets while minimizing off-target effects is a complex challenge. Non-specific delivery can lead to adverse reactions and unintended gene modulation, potentially causing harm to healthy cells or tissues. Developing highly selective RNA molecules, optimizing delivery techniques, and employing innovative design strategies are essential for achieving precise and targeted RNA delivery. Overcoming the delivery challenge is not only about protecting and targeting RNA molecules but also about enhancing their cellular uptake. Many cell types have barriers that limit the entry of foreign genetic material, making it essential to develop efficient delivery systems that facilitate RNA uptake. In this regard, nanoparticles and other delivery vehicles have been designed to improve cellular internalization, increasing the chances of RNA molecules reaching their intended destination.

Manufacturing Scalability and Cost

The production of RNA therapeutics is inherently intricate, involving the synthesis, purification, and formulation of RNA molecules. These processes require specialized equipment, skilled personnel, and adherence to stringent quality control standards. Developing RNA molecules in a consistent and reproducible manner is essential for ensuring their safety and efficacy. The complexity of these manufacturing processes can slow down production and limit the capacity to scale up to meet the growing demand. RNA therapeutics are often costly to manufacture due to the expenses associated with research, development, clinical trials, and large-scale production. The use of specialized materials and equipment, as well as the need for highly skilled professionals, contributes to the high production costs. Additionally, regulatory compliance and quality control measures further increase expenses. High production

costs can result in expensive therapies, limiting patient access and payer willingness to cover the cost of treatment.

Moving from small-scale research and development to commercial-scale production is a significant challenge. Achieving the economies of scale required for large-scale manufacturing can be complicated and resource-intensive. Companies must invest in infrastructure, personnel, and equipment to meet the demand for RNA therapeutics. Ensuring that the quality and consistency of production are maintained at a larger scale is vital for regulatory approval and patient safety. The production of RNA therapeutics requires cutting-edge technology and infrastructure. Manufacturers must invest in state-of-the-art equipment and facilities to ensure the quality, purity, and reproducibility of RNA molecules. These technological requirements can be a barrier for smaller companies and research institutions with limited resources.

Key Market Trends

Emergence of mRNA Vaccines

The emergence of mRNA vaccines has proven to be a game-changer in the North America RNA therapeutics market. mRNA (messenger RNA) vaccines, exemplified by the COVID-19 vaccines developed by Pfizer-BioNTech and Moderna, have demonstrated unprecedented success in preventing viral infections and have generated a wave of excitement and optimism within the medical and pharmaceutical communities. The success of mRNA vaccines against COVID-19 has significantly boosted the credibility and potential of RNA-based therapies. These vaccines, which work by introducing a small piece of synthetic mRNA encoding a viral antigen into the body, instruct the immune system to recognize and attack the virus. This innovative approach not only exhibited remarkable efficacy but also demonstrated a rapid development timeline, with vaccines brought from concept to market within record time. This success story has spotlighted the versatility and power of mRNA technology.

The impact of mRNA vaccines extends far beyond infectious diseases. They have catalyzed research and development in other therapeutic areas, propelling the growth of the broader RNA therapeutics market. Applications in oncology, cardiovascular diseases, autoimmune disorders, and rare genetic conditions are now being explored with renewed vigor. The mRNA platform has showcased its adaptability to target various diseases, providing a foundation for precision medicine. The success of mRNA vaccines has accelerated research in novel delivery systems and enhanced RNA stability, which benefits the overall RNA therapeutics market. Researchers and

companies are actively working on improving the safety, efficacy, and delivery of mRNA-based therapies for various diseases. This progress is not only limited to the COVID-19 pandemic but has opened doors to a new era of medical innovation. The emergence of mRNA vaccines has further propelled investments and collaborations in the RNA therapeutics field. The increased attention and support from both governmental and private sectors have stimulated research and development efforts, ultimately advancing the entire RNA therapeutics market. Collaborative initiatives and partnerships among pharmaceutical companies, biotech firms, academic institutions, and research organizations have become more prevalent, fostering innovation and knowledge sharing.

Adoption of Personalized Medicine

The adoption of personalized medicine is proving to be a significant driver of growth in the North America RNA therapeutics market. Personalized medicine, often referred to as precision medicine, is a healthcare approach that tailors medical treatments to the individual characteristics of each patient. This approach includes considering a patient's genetic makeup, molecular profile, and specific disease characteristics when designing and administering treatments. RNA therapeutics, particularly messenger RNA (mRNA) and antisense oligonucleotide (ASO) therapies, are well-suited for personalized medicine, and their utilization in this context is catalyzing market expansion. The key aspects of personalized medicine is the ability to target and modulate genes associated with specific diseases. RNA-based therapies are uniquely positioned to deliver on this promise. For example, mRNA technology allows for the design and production of custom mRNA molecules encoding therapeutic proteins or antigens tailored to an individual's unique genetic profile. This precision enables treatments that are not only highly effective but also remarkably safe, as they minimize off-target effects.

RNA therapeutics also play a crucial role in addressing rare genetic diseases and hereditary conditions, where conventional treatment options are often limited or entirely absent. These therapies can be customized to correct or compensate for specific genetic mutations responsible for rare diseases, offering newfound hope for patients and their families. The personalization of treatment for these conditions contributes significantly to the adoption of RNA therapeutics and bolsters market demand. The rapidly evolving field of personalized cancer therapy has embraced RNA-based treatments. By analyzing the genetic makeup and molecular characteristics of a patient's tumor, researchers can design RNA therapies that precisely target cancer-specific genes. RNA therapeutics, including small interfering RNA (siRNA) and microRNA (miRNA) therapies, can be fine-tuned to interfere with the oncogenes

responsible for tumor growth while sparing healthy cells. This level of precision is revolutionizing cancer treatment, improving outcomes, and enhancing the patient experience.

Segmental Insights

Type Insights

Based on the Type mRNA Therapeutics emerged as the dominant segment in the North America market for North America RNA Therapeutics Market in 2023. The successful development and widespread adoption of mRNA vaccines against COVID-19, such as the Pfizer-BioNTech and Moderna vaccines, have catapulted mRNA technology into the spotlight. These vaccines demonstrated exceptional efficacy and safety, leading to their North America distribution. This success has significantly boosted the recognition and demand for mRNA therapeutics, given that they are based on a similar platform. mRNA therapeutics have diverse applications beyond vaccines. They can be tailored to encode specific proteins, enabling the development of treatments for a wide range of diseases, including cancer, infectious diseases, autoimmune disorders, and rare genetic conditions. The adaptability and versatility of mRNA technology make it highly sought after in the RNA therapeutics market. The development of mRNA-based therapies is known for its speed and efficiency. Compared to traditional drug development, mRNA therapeutics can be designed, synthesized, and produced relatively quickly. This agility in research and development is particularly valuable in responding to emerging health threats and addressing unmet medical needs.

Indication Insights

Based on the Indication, Infectious Diseases emerged as the dominant segment in the North America market for North America RNA Therapeutics Market in 2023. RNA-based therapies, particularly mRNA, are highly versatile and adaptable. They can be designed to encode specific antigens or proteins that target various pathogens, making them suitable for a broad range of infectious diseases. This adaptability is crucial in addressing different infectious agents, from viruses to bacteria, and even future unknown pathogens. RNA therapeutics, especially mRNA vaccines, are known for their rapid development timelines. This agility is invaluable when dealing with infectious disease outbreaks, where quick responses can help curb the spread of the disease and save lives. The ability to design and produce RNA-based treatments in a time-sensitive manner enhances their demand.

Regional Insights

United States emerged as the dominant country in the North America RNA Therapeutics Market in 2023, holding the largest market share. The United States is home to a robust biotechnology and pharmaceutical industry with numerous leading companies, research institutions, and academic centers. These organizations have been at the forefront of RNA therapeutics research and development, contributing to the growth of the market. Significant Investment in Research and Development: The U.S. government, along with private sector investment, has allocated substantial resources for RNA therapeutics research and development. This funding has facilitated innovation and the development of novel RNA-based therapies. The United States has a well-trained and skilled workforce in the field of biotechnology and RNA therapeutics, including scientists, researchers, and clinicians. This expertise has played a crucial role in advancing the field and bringing RNA therapies to market. The regulatory environment in the United States, led by the Food and Drug Administration (FDA), has been supportive of RNA therapeutics development. Regulatory pathways for RNA-based drugs have been well-established, which has encouraged companies to invest in this market.

Key Market Players

Moderna, Inc.

Anylam Pharmaceuticals, Inc.

Novartis Ag

Ionis Pharmaceuticals, Inc.

Sarepta Therapeutics, Inc

Sanofi Ag

Pfizer Inc

Biontech SE

Orna Therapeutics Inc.

CPISPER Therapeutics AG

Report Scope:

In this report, the North America RNA Therapeutics Market has been segmented into the following categories, in addition to the industry trends which have also been detailed below:

North America RNA Therapeutics Market, By Product:

Vaccines

Drugs

North America RNA Therapeutics Market, By Type:

mRNA Therapeutics

RNA Interference (RNAi) Therapeutics

Antisense Oligonucleotide (ASO) Therapeutics

Other Therapeutics

North America RNA Therapeutics Market, By Indication:

Infectious Diseases

Rare Genetic Diseases/Hereditary Diseases

Other Indications

North America RNA Therapeutics Market, By End-use:

Hospitals & Clinics

Ambulatory Surgical Centers

Others

North America RNA Therapeutics Market, By Country:

United States

Canada

Mexico

Competitive Landscape

Company Profiles: Detailed analysis of the major companies present in the North America RNA Therapeutics Market.

Available Customizations:

North America RNA Therapeutics Market report with the given market data, TechSci Research offers customizations according to a company's specific needs. The following customization options are available for the report:

Company Information

Detailed analysis and profiling of additional market players (up to five).

Contents

1. PRODUCT OVERVIEW

- 1.1. Market Definition
- 1.2. Scope of the Market
 - 1.2.1. Markets Covered
 - 1.2.2. Years Considered for Study
 - 1.2.3. Key Market Segmentations

2. RESEARCH METHODOLOGY

- 2.1. Objective of the Study
- 2.2. Baseline Methodology
- 2.3. Key Industry Partners
- 2.4. Major Association and Secondary Sources
- 2.5. Forecasting Methodology
- 2.6. Data Triangulation & Validation
- 2.7. Assumptions and Limitations

3. EXECUTIVE SUMMARY

- 3.1. Overview of the Market
- 3.2. Overview of Key Market Segmentations
- 3.3. Overview of Key Market Players
- 3.4. Overview of Key Regions/Countries
- 3.5. Overview of Market Drivers, Challenges, Trends

4. NORTH AMERICA RNA THERAPEUTICS MARKET OUTLOOK

- 4.1. Market Size & Forecast
 - 4.1.1. By Value
- 4.2. Market Share & Forecast
 - 4.2.1. By Product (Vaccines, Drugs)
 - 4.2.2. By Type (mRNA Therapeutics, RNA Interference (RNAi) Therapeutics, Antisense Oligonucleotide (ASO) Therapeutics, Other Therapeutics)
 - 4.2.3. By Indication (Infectious Diseases, Rare Genetic Diseases/Hereditary Diseases, Other Indications)
 - 4.2.4. By End User (Hospitals and Clinics, Ambulatory Surgical Centers, Others)

4.2.5. By Country

4.2.6. By Company (2023)

4.3. Market Map

4.3.1. By Product

4.3.2. By Type

4.3.3. By Indication

4.3.4. By End User

4.3.5. By Country

5. UNITED STATES RNA THERAPEUTICS MARKET OUTLOOK

5.1. Market Size & Forecast

5.1.1. By Value

5.2. Market Share & Forecast

5.2.1. By Product

5.2.2. By Type

5.2.3. By Indication

5.2.4. By End User

6. MEXICO RNA THERAPEUTICS MARKET OUTLOOK

6.1. Market Size & Forecast

6.1.1. By Value

6.2. Market Share & Forecast

6.2.1. By Product

6.2.2. By Type

6.2.3. By Indication

6.2.4. By End User

7. CANADA RNA THERAPEUTICS MARKET OUTLOOK

7.1. Market Size & Forecast

7.1.1. By Value

7.2. Market Share & Forecast

7.2.1. By Product

7.2.2. By Type

7.2.3. By Indication

7.2.4. By End User

8. MARKET DYNAMICS

- 8.1. Drivers
- 8.2. Challenges

9. MARKET TRENDS & DEVELOPMENTS

- 9.1. Recent Developments
- 9.2. Product Launches
- 9.3. Mergers & Acquisitions

10. NORTH AMERICA RNA THERAPEUTICS MARKET: SWOT ANALYSIS

11. PORTER'S FIVE FORCES ANALYSIS

- 11.1. Competition in the Industry
- 11.2. Potential of New Entrants
- 11.3. Power of Suppliers
- 11.4. Power of Customers
- 11.5. Threat of Substitute Product

12. COMPETITIVE LANDSCAPE

- 12.1. Moderna, Inc.
 - 12.1.1. Business Overview
 - 12.1.2. Company Snapshot
 - 12.1.3. Products & Services
 - 12.1.4. Current Capacity Analysis
 - 12.1.5. Financials (In case of listed)
 - 12.1.6. Recent Developments
 - 12.1.7. SWOT Analysis
- 12.2. Alnylam Pharmaceuticals, Inc.
- 12.3. Novartis Ag
- 12.4. Ionis Pharmaceuticals, Inc.
- 12.5. Sarepta Therapeutics, Inc
- 12.6. Sanofi Ag
- 12.7. Pfizer Inc
- 12.8. Biontech SE
- 12.9. Orna Therapeutics Inc.

12.10. CPISPER Therapeutics AG

13. STRATEGIC RECOMMENDATIONS

14. ABOUT US & DISCLAIMER

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