

Global Dalteparin Sodium Injection Market Growth 2026-2032

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Abstracts

The global Dalteparin Sodium Injection market size is predicted to grow from US\$ 1834 million in 2025 to US\$ 2816 million in 2032; it is expected to grow at a CAGR of 6.3% from 2026 to 2032.

Dalteparin Sodium Injection is a low molecular weight heparin (LMWH) anticoagulant used primarily for the prevention and treatment of thromboembolic disorders, including deep vein thrombosis (DVT) and pulmonary embolism (PE). Its mechanism of action involves enhancing the activity of antithrombin III, thereby inhibiting coagulation factor Xa and reducing the risk of blood clot formation. This medication was developed to address the limitations of conventional heparin, such as unpredictable dosing, the need for frequent coagulation monitoring, and a higher risk of bleeding.

Dalteparin Sodium was first developed and clinically introduced in the 1990s. Its low molecular weight properties provide more predictable anticoagulant effects and lower adverse risks. The manufacturing process relies on upstream raw materials such as heparin and derivative chemical reagents, while essential components like injectable vials, rubber stoppers, and needles must meet pharmaceutical-grade standards. The supply chain is tightly controlled to ensure the safety and stability of the final product. In 2025, the global production capacity of dalteparin sodium injection was 150 million boxes, with sales volume reaching 118 million boxes, an average unit price of 15.8 USD per box, and corporate gross profit margins ranging between 25% and 35%.

Currently, the market for Dalteparin Sodium Injection is experiencing steady development both globally and within domestic healthcare systems. As a key low molecular weight heparin anticoagulant, it has earned broad clinical acceptance, particularly in perioperative anticoagulation, management of venous thromboembolic

disorders, and prevention of cancer-associated thrombosis. The increasing prevalence of aging populations and cardiovascular-related diseases has led to sustained demand for anticoagulant therapies that are both safe and predictable, underpinning the clinical adoption and market penetration of dalteparin. Moreover, supportive drug reimbursement policies and the push for standardized anticoagulant use in medical institutions have helped normalize its role across multidisciplinary treatment contexts. Nonetheless, the competitive landscape is intensifying, especially with the rapid adoption of novel oral anticoagulants in certain indications, creating a degree of substitution pressure that challenges the dalteparin sodium injection market.

Looking ahead, evolving clinical treatment paradigms and the promotion of individualized anticoagulation strategies are expected to raise expectations for traditional anticoagulants, emphasizing not only efficacy and safety but also patient adherence and ease of use. This environment encourages ongoing innovation in product refinement, dosing regimens, and integrated patient management services for low molecular weight heparins including dalteparin. Advances in production technology and manufacturing processes are also likely to contribute to improved product consistency and cost optimization. Furthermore, deeper exploration in life sciences research may uncover new therapeutic indications and strengthen clinical evidence for combination therapies, offering fresh opportunities for market expansion. Enhanced anticoagulation safety monitoring and improved patient treatment experiences could become significant drivers of industry growth.

Despite a positive outlook, several inhibiting factors remain. Stricter regulatory environments demand higher standards for production quality control, leading to increased R&D, regulatory submission, and compliance costs and risks for manufacturers. At the same time, intense competition from generic products and price pressures compress profit margins, particularly for innovators. Clinical challenges remain in balancing bleeding risk assessments and accounting for patient heterogeneity, necessitating more comprehensive clinical pathways and evidence-based support. Additionally, variations in how healthcare systems select and utilize anticoagulants influence the depth and breadth of dalteparin sodium injection's adoption across different regions.

LP Information, Inc. (LPI) ' newest research report, the "Dalteparin Sodium Injection Industry Forecast" looks at past sales and reviews total world Dalteparin Sodium Injection sales in 2025, providing a comprehensive analysis by region and market sector of projected Dalteparin Sodium Injection sales for 2026 through 2032. With Dalteparin Sodium Injection sales broken down by region, market sector and sub-sector, this report

provides a detailed analysis in US\$ millions of the world Dalteparin Sodium Injection industry.

This Insight Report provides a comprehensive analysis of the global Dalteparin Sodium Injection 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 Dalteparin Sodium Injection portfolios and capabilities, market entry strategies, market positions, and geographic footprints, to better understand these firms' unique position in an accelerating global Dalteparin Sodium Injection market.

This Insight Report evaluates the key market trends, drivers, and affecting factors shaping the global outlook for Dalteparin Sodium Injection 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 Dalteparin Sodium Injection.

This report presents a comprehensive overview, market shares, and growth opportunities of Dalteparin Sodium Injection market by product type, application, key manufacturers and key regions and countries.

Segmentation by Type:

Dalteparin Sodium Injection 0.2ml:2500IU

Dalteparin Sodium Injection 0.2ml:5000IU

Dalteparin Sodium Injection 0.3ml:7500IU

Dalteparin Sodium Injection 0.5ml:12500IU

Dalteparin Sodium Injection 0.6ml:15000IU

Segmentation by Packaging:

Prefilled Syringe

Single-Dose Vial

Segmentation by Manufacturing Process:

Porcine Intestinal Mucosa Origin

Chemical Depolymerization

Nitrous Acid Depolymerization

Segmentation by Application:

Hospital

Nursing Institutions

Home Care

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

The below companies that are profiled have been selected based on inputs gathered from primary experts and analysing the company's coverage, product portfolio, its market penetration.

Pfizer

Hebei Changshan Biochemical Pharmaceutical Co.,Ltd.

Nanjing King-Friend Biochemical Pharmaceutical Co., Ltd.

Changzhou Qianhong Biopharma Co., Ltd.

Qilu Pharmaceutical

Yantai Dongcheng Northern Pharmaceutical

Key Questions Addressed in this Report

What is the 10-year outlook for the global Dalteparin Sodium Injection market?

What factors are driving Dalteparin Sodium Injection market growth, globally and by region?

Which technologies are poised for the fastest growth by market and region?

How do Dalteparin Sodium Injection market opportunities vary by end market size?

How does Dalteparin Sodium Injection break out by Type, by Application?

Contents

1 SCOPE OF THE REPORT

- 1.1 Market Introduction
- 1.2 Years Considered
- 1.3 Research Objectives
- 1.4 Market Research Methodology
- 1.5 Research Process and Data Source
- 1.6 Economic Indicators
- 1.7 Currency Considered
- 1.8 Market Estimation Caveats

2 EXECUTIVE SUMMARY

2.1 World Market Overview

- 2.1.1 Global Dalteparin Sodium Injection Annual Sales 2021-2032
- 2.1.2 World Current & Future Analysis for Dalteparin Sodium Injection by Geographic Region, 2021, 2025 & 2032
- 2.1.3 World Current & Future Analysis for Dalteparin Sodium Injection by Country/Region, 2021, 2025 & 2032

2.2 Dalteparin Sodium Injection Segment by Type

- 2.2.1 Dalteparin Sodium Injection 0.2ml:2500IU
- 2.2.2 Dalteparin Sodium Injection 0.2ml:5000IU
- 2.2.3 Dalteparin Sodium Injection 0.3ml:7500IU
- 2.2.4 Dalteparin Sodium Injection 0.5ml:12500IU
- 2.2.5 Dalteparin Sodium Injection 0.6ml:15000IU
- 2.2.6 Dalteparin Sodium Injection Sales by Type
 - 2.2.6.1 Global Dalteparin Sodium Injection Sales Market Share by Type (2021-2026)
 - 2.2.6.2 Global Dalteparin Sodium Injection Revenue and Market Share by Type (2021-2026)
 - 2.2.6.3 Global Dalteparin Sodium Injection Sale Price by Type (2021-2026)

2.3 Dalteparin Sodium Injection Segment by Packaging

- 2.3.1 Prefilled Syringe
- 2.3.2 Single-Dose Vial
- 2.3.3 Dalteparin Sodium Injection Sales by Packaging
 - 2.3.3.1 Global Dalteparin Sodium Injection Sales Market Share by Packaging (2021-2026)
 - 2.3.3.2 Global Dalteparin Sodium Injection Revenue and Market Share by Packaging

(2021-2026)

2.3.3.3 Global Dalteparin Sodium Injection Sale Price by Packaging (2021-2026)

2.4 Dalteparin Sodium Injection Segment by Manufacturing Process

2.4.1 Porcine Intestinal Mucosa Origin

2.4.2 Chemical Depolymerization

2.4.3 Nitrous Acid Depolymerization

2.4.4 Dalteparin Sodium Injection Sales by Manufacturing Process

2.4.4.1 Global Dalteparin Sodium Injection Sales Market Share by Manufacturing Process (2021-2026)

2.4.4.2 Global Dalteparin Sodium Injection Revenue and Market Share by Manufacturing Process (2021-2026)

2.4.4.3 Global Dalteparin Sodium Injection Sale Price by Manufacturing Process (2021-2026)

2.5 Dalteparin Sodium Injection Segment by Application

2.5.1 Hospital

2.5.2 Nursing Institutions

2.5.3 Home Care

2.5.4 Others

2.5.5 Dalteparin Sodium Injection Sales by Application

2.5.5.1 Global Dalteparin Sodium Injection Sale Market Share by Application (2021-2026)

2.5.5.2 Global Dalteparin Sodium Injection Revenue and Market Share by Application (2021-2026)

2.5.5.3 Global Dalteparin Sodium Injection Sale Price by Application (2021-2026)

3 GLOBAL BY COMPANY

3.1 Global Dalteparin Sodium Injection Breakdown Data by Company

3.1.1 Global Dalteparin Sodium Injection Annual Sales by Company (2021-2026)

3.1.2 Global Dalteparin Sodium Injection Sales Market Share by Company (2021-2026)

3.2 Global Dalteparin Sodium Injection Annual Revenue by Company (2021-2026)

3.2.1 Global Dalteparin Sodium Injection Revenue by Company (2021-2026)

3.2.2 Global Dalteparin Sodium Injection Revenue Market Share by Company (2021-2026)

3.3 Global Dalteparin Sodium Injection Sale Price by Company

3.4 Key Manufacturers Dalteparin Sodium Injection Producing Area Distribution, Sales Area, Product Type

3.4.1 Key Manufacturers Dalteparin Sodium Injection Product Location Distribution

- 3.4.2 Players Dalteparin Sodium Injection Products Offered
- 3.5 Market Concentration Rate Analysis
 - 3.5.1 Competition Landscape Analysis
 - 3.5.2 Concentration Ratio (CR3, CR5 and CR10) & (2024-2026)
- 3.6 New Products and Potential Entrants
- 3.7 Market M&A Activity & Strategy

4 WORLD HISTORIC REVIEW FOR DALTEPARIN SODIUM INJECTION BY GEOGRAPHIC REGION

- 4.1 World Historic Dalteparin Sodium Injection Market Size by Geographic Region (2021-2026)
 - 4.1.1 Global Dalteparin Sodium Injection Annual Sales by Geographic Region (2021-2026)
 - 4.1.2 Global Dalteparin Sodium Injection Annual Revenue by Geographic Region (2021-2026)
- 4.2 World Historic Dalteparin Sodium Injection Market Size by Country/Region (2021-2026)
 - 4.2.1 Global Dalteparin Sodium Injection Annual Sales by Country/Region (2021-2026)
 - 4.2.2 Global Dalteparin Sodium Injection Annual Revenue by Country/Region (2021-2026)
- 4.3 Americas Dalteparin Sodium Injection Sales Growth
- 4.4 APAC Dalteparin Sodium Injection Sales Growth
- 4.5 Europe Dalteparin Sodium Injection Sales Growth
- 4.6 Middle East & Africa Dalteparin Sodium Injection Sales Growth

5 AMERICAS

- 5.1 Americas Dalteparin Sodium Injection Sales by Country
 - 5.1.1 Americas Dalteparin Sodium Injection Sales by Country (2021-2026)
 - 5.1.2 Americas Dalteparin Sodium Injection Revenue by Country (2021-2026)
- 5.2 Americas Dalteparin Sodium Injection Sales by Type (2021-2026)
- 5.3 Americas Dalteparin Sodium Injection Sales by Application (2021-2026)
- 5.4 United States
- 5.5 Canada
- 5.6 Mexico
- 5.7 Brazil

6 APAC

6.1 APAC Dalteparin Sodium Injection Sales by Region

6.1.1 APAC Dalteparin Sodium Injection Sales by Region (2021-2026)

6.1.2 APAC Dalteparin Sodium Injection Revenue by Region (2021-2026)

6.2 APAC Dalteparin Sodium Injection Sales by Type (2021-2026)

6.3 APAC Dalteparin Sodium Injection Sales by Application (2021-2026)

6.4 China

6.5 Japan

6.6 South Korea

6.7 Southeast Asia

6.8 India

6.9 Australia

6.10 China Taiwan

7 EUROPE

7.1 Europe Dalteparin Sodium Injection by Country

7.1.1 Europe Dalteparin Sodium Injection Sales by Country (2021-2026)

7.1.2 Europe Dalteparin Sodium Injection Revenue by Country (2021-2026)

7.2 Europe Dalteparin Sodium Injection Sales by Type (2021-2026)

7.3 Europe Dalteparin Sodium Injection Sales by Application (2021-2026)

7.4 Germany

7.5 France

7.6 UK

7.7 Italy

7.8 Russia

8 MIDDLE EAST & AFRICA

8.1 Middle East & Africa Dalteparin Sodium Injection by Country

8.1.1 Middle East & Africa Dalteparin Sodium Injection Sales by Country (2021-2026)

8.1.2 Middle East & Africa Dalteparin Sodium Injection Revenue by Country
(2021-2026)

8.2 Middle East & Africa Dalteparin Sodium Injection Sales by Type (2021-2026)

8.3 Middle East & Africa Dalteparin Sodium Injection Sales by Application (2021-2026)

8.4 Egypt

8.5 South Africa

8.6 Israel

8.7 Turkey

8.8 GCC Countries

9 MARKET DRIVERS, CHALLENGES AND TRENDS

9.1 Market Drivers & Growth Opportunities

9.2 Market Challenges & Risks

9.3 Industry Trends

10 MANUFACTURING COST STRUCTURE ANALYSIS

10.1 Raw Material and Suppliers

10.2 Manufacturing Cost Structure Analysis of Dalteparin Sodium Injection

10.3 Manufacturing Process Analysis of Dalteparin Sodium Injection

10.4 Industry Chain Structure of Dalteparin Sodium Injection

11 MARKETING, DISTRIBUTORS AND CUSTOMER

11.1 Sales Channel

11.1.1 Direct Channels

11.1.2 Indirect Channels

11.2 Dalteparin Sodium Injection Distributors

11.3 Dalteparin Sodium Injection Customer

12 WORLD FORECAST REVIEW FOR DALTEPARIN SODIUM INJECTION BY GEOGRAPHIC REGION

12.1 Global Dalteparin Sodium Injection Market Size Forecast by Region

12.1.1 Global Dalteparin Sodium Injection Forecast by Region (2027-2032)

12.1.2 Global Dalteparin Sodium Injection Annual Revenue Forecast by Region (2027-2032)

12.2 Americas Forecast by Country (2027-2032)

12.3 APAC Forecast by Region (2027-2032)

12.4 Europe Forecast by Country (2027-2032)

12.5 Middle East & Africa Forecast by Country (2027-2032)

12.6 Global Dalteparin Sodium Injection Forecast by Type (2027-2032)

12.7 Global Dalteparin Sodium Injection Forecast by Application (2027-2032)

13 KEY PLAYERS ANALYSIS

13.1 Pfizer

13.1.1 Pfizer Company Information

13.1.2 Pfizer Dalteparin Sodium Injection Product Portfolios and Specifications

13.1.3 Pfizer Dalteparin Sodium Injection Sales, Revenue, Price and Gross Margin (2021-2026)

13.1.4 Pfizer Main Business Overview

13.1.5 Pfizer Latest Developments

13.2 Hebei Changshan Biochemical Pharmaceutical Co.,Ltd.

13.2.1 Hebei Changshan Biochemical Pharmaceutical Co.,Ltd. Company Information

13.2.2 Hebei Changshan Biochemical Pharmaceutical Co.,Ltd. Dalteparin Sodium Injection Product Portfolios and Specifications

13.2.3 Hebei Changshan Biochemical Pharmaceutical Co.,Ltd. Dalteparin Sodium Injection Sales, Revenue, Price and Gross Margin (2021-2026)

13.2.4 Hebei Changshan Biochemical Pharmaceutical Co.,Ltd. Main Business Overview

13.2.5 Hebei Changshan Biochemical Pharmaceutical Co.,Ltd. Latest Developments

13.3 Nanjing King-Friend Biochemical Pharmaceutical Co., Ltd.

13.3.1 Nanjing King-Friend Biochemical Pharmaceutical Co., Ltd. Company Information

13.3.2 Nanjing King-Friend Biochemical Pharmaceutical Co., Ltd. Dalteparin Sodium Injection Product Portfolios and Specifications

13.3.3 Nanjing King-Friend Biochemical Pharmaceutical Co., Ltd. Dalteparin Sodium Injection Sales, Revenue, Price and Gross Margin (2021-2026)

13.3.4 Nanjing King-Friend Biochemical Pharmaceutical Co., Ltd. Main Business Overview

13.3.5 Nanjing King-Friend Biochemical Pharmaceutical Co., Ltd. Latest Developments

13.4 Changzhou Qianhong Biopharma Co., Ltd.

13.4.1 Changzhou Qianhong Biopharma Co., Ltd. Company Information

13.4.2 Changzhou Qianhong Biopharma Co., Ltd. Dalteparin Sodium Injection Product Portfolios and Specifications

13.4.3 Changzhou Qianhong Biopharma Co., Ltd. Dalteparin Sodium Injection Sales, Revenue, Price and Gross Margin (2021-2026)

13.4.4 Changzhou Qianhong Biopharma Co., Ltd. Main Business Overview

13.4.5 Changzhou Qianhong Biopharma Co., Ltd. Latest Developments

13.5 Qilu Pharmaceutical

13.5.1 Qilu Pharmaceutical Company Information

13.5.2 Qilu Pharmaceutical Dalteparin Sodium Injection Product Portfolios and Specifications

13.5.3 Qilu Pharmaceutical Dalteparin Sodium Injection Sales, Revenue, Price and Gross Margin (2021-2026)

13.5.4 Qilu Pharmaceutical Main Business Overview

13.5.5 Qilu Pharmaceutical Latest Developments

13.6 Yantai Dongcheng Northern Pharmaceutical

13.6.1 Yantai Dongcheng Northern Pharmaceutical Company Information

13.6.2 Yantai Dongcheng Northern Pharmaceutical Dalteparin Sodium Injection Product Portfolios and Specifications

13.6.3 Yantai Dongcheng Northern Pharmaceutical Dalteparin Sodium Injection Sales, Revenue, Price and Gross Margin (2021-2026)

13.6.4 Yantai Dongcheng Northern Pharmaceutical Main Business Overview

13.6.5 Yantai Dongcheng Northern Pharmaceutical Latest Developments

14 RESEARCH FINDINGS AND CONCLUSION

List Of Tables

LIST OF TABLES

Table 1. Dalteparin Sodium Injection Annual Sales CAGR by Geographic Region (2021, 2025 & 2032) & (\$ millions)

Table 2. Dalteparin Sodium Injection Annual Sales CAGR by Country/Region (2021, 2025 & 2032) & (\$ millions)

Table 3. Major Players of Dalteparin Sodium Injection 0.2ml:2500IU

Table 4. Major Players of Dalteparin Sodium Injection 0.2ml:5000IU

Table 5. Major Players of Dalteparin Sodium Injection 0.3ml:7500IU

Table 6. Major Players of Dalteparin Sodium Injection 0.5ml:12500IU

Table 7. Major Players of Dalteparin Sodium Injection 0.6ml:15000IU

Table 8. Global Dalteparin Sodium Injection Sales by Type (2021-2026) & (K Units)

Table 9. Global Dalteparin Sodium Injection Sales Market Share by Type (2021-2026)

Table 10. Global Dalteparin Sodium Injection Revenue by Type (2021-2026) & (\$ million)

Table 11. Global Dalteparin Sodium Injection Revenue Market Share by Type (2021-2026)

Table 12. Global Dalteparin Sodium Injection Sale Price by Type (2021-2026) & (US\$/Unit)

Table 13. Major Players of Prefilled Syringe

Table 14. Major Players of Single-Dose Vial

Table 15. Global Dalteparin Sodium Injection Sales by Packaging (2021-2026) & (K Units)

Table 16. Global Dalteparin Sodium Injection Sales Market Share by Packaging (2021-2026)

Table 17. Global Dalteparin Sodium Injection Revenue by Packaging (2021-2026) & (\$ million)

Table 18. Global Dalteparin Sodium Injection Revenue Market Share by Packaging (2021-2026)

Table 19. Global Dalteparin Sodium Injection Sale Price by Packaging (2021-2026) & (US\$/Unit)

Table 20. Major Players of Porcine Intestinal Mucosa Origin

Table 21. Major Players of Chemical Depolymerization

Table 22. Major Players of Nitrous Acid Depolymerization

Table 23. Global Dalteparin Sodium Injection Sales by Manufacturing Process (2021-2026) & (K Units)

Table 24. Global Dalteparin Sodium Injection Sales Market Share by Manufacturing

Process (2021-2026)

Table 25. Global Dalteparin Sodium Injection Revenue by Manufacturing Process (2021-2026) & (\$ million)

Table 26. Global Dalteparin Sodium Injection Revenue Market Share by Manufacturing Process (2021-2026)

Table 27. Global Dalteparin Sodium Injection Sale Price by Manufacturing Process (2021-2026) & (US\$/Unit)

Table 28. Global Dalteparin Sodium Injection Sale by Application (2021-2026) & (K Units)

Table 29. Global Dalteparin Sodium Injection Sale Market Share by Application (2021-2026)

Table 30. Global Dalteparin Sodium Injection Revenue by Application (2021-2026) & (\$ million)

Table 31. Global Dalteparin Sodium Injection Revenue Market Share by Application (2021-2026)

Table 32. Global Dalteparin Sodium Injection Sale Price by Application (2021-2026) & (US\$/Unit)

Table 33. Global Dalteparin Sodium Injection Sales by Company (2021-2026) & (K Units)

Table 34. Global Dalteparin Sodium Injection Sales Market Share by Company (2021-2026)

Table 35. Global Dalteparin Sodium Injection Revenue by Company (2021-2026) & (\$ millions)

Table 36. Global Dalteparin Sodium Injection Revenue Market Share by Company (2021-2026)

Table 37. Global Dalteparin Sodium Injection Sale Price by Company (2021-2026) & (US\$/Unit)

Table 38. Key Manufacturers Dalteparin Sodium Injection Producing Area Distribution and Sales Area

Table 39. Players Dalteparin Sodium Injection Products Offered

Table 40. Dalteparin Sodium Injection Concentration Ratio (CR3, CR5 and CR10) & (2024-2026)

Table 41. New Products and Potential Entrants

Table 42. Market M&A Activity & Strategy

Table 43. Global Dalteparin Sodium Injection Sales by Geographic Region (2021-2026) & (K Units)

Table 44. Global Dalteparin Sodium Injection Sales Market Share Geographic Region (2021-2026)

Table 45. Global Dalteparin Sodium Injection Revenue by Geographic Region

(2021-2026) & (\$ millions)

Table 46. Global Dalteparin Sodium Injection Revenue Market Share by Geographic Region (2021-2026)

Table 47. Global Dalteparin Sodium Injection Sales by Country/Region (2021-2026) & (K Units)

Table 48. Global Dalteparin Sodium Injection Sales Market Share by Country/Region (2021-2026)

Table 49. Global Dalteparin Sodium Injection Revenue by Country/Region (2021-2026) & (\$ millions)

Table 50. Global Dalteparin Sodium Injection Revenue Market Share by Country/Region (2021-2026)

Table 51. Americas Dalteparin Sodium Injection Sales by Country (2021-2026) & (K Units)

Table 52. Americas Dalteparin Sodium Injection Sales Market Share by Country (2021-2026)

Table 53. Americas Dalteparin Sodium Injection Revenue by Country (2021-2026) & (\$ millions)

Table 54. Americas Dalteparin Sodium Injection Sales by Type (2021-2026) & (K Units)

Table 55. Americas Dalteparin Sodium Injection Sales by Application (2021-2026) & (K Units)

Table 56. APAC Dalteparin Sodium Injection Sales by Region (2021-2026) & (K Units)

Table 57. APAC Dalteparin Sodium Injection Sales Market Share by Region (2021-2026)

Table 58. APAC Dalteparin Sodium Injection Revenue by Region (2021-2026) & (\$ millions)

Table 59. APAC Dalteparin Sodium Injection Sales by Type (2021-2026) & (K Units)

Table 60. APAC Dalteparin Sodium Injection Sales by Application (2021-2026) & (K Units)

Table 61. Europe Dalteparin Sodium Injection Sales by Country (2021-2026) & (K Units)

Table 62. Europe Dalteparin Sodium Injection Revenue by Country (2021-2026) & (\$ millions)

Table 63. Europe Dalteparin Sodium Injection Sales by Type (2021-2026) & (K Units)

Table 64. Europe Dalteparin Sodium Injection Sales by Application (2021-2026) & (K Units)

Table 65. Middle East & Africa Dalteparin Sodium Injection Sales by Country (2021-2026) & (K Units)

Table 66. Middle East & Africa Dalteparin Sodium Injection Revenue Market Share by Country (2021-2026)

Table 67. Middle East & Africa Dalteparin Sodium Injection Sales by Type (2021-2026)

& (K Units)

Table 68. Middle East & Africa Dalteparin Sodium Injection Sales by Application (2021-2026) & (K Units)

Table 69. Key Market Drivers & Growth Opportunities of Dalteparin Sodium Injection

Table 70. Key Market Challenges & Risks of Dalteparin Sodium Injection

Table 71. Key Industry Trends of Dalteparin Sodium Injection

Table 72. Dalteparin Sodium Injection Raw Material

Table 73. Key Suppliers of Raw Materials

Table 74. Dalteparin Sodium Injection Distributors List

Table 75. Dalteparin Sodium Injection Customer List

Table 76. Global Dalteparin Sodium Injection Sales Forecast by Region (2027-2032) & (K Units)

Table 77. Global Dalteparin Sodium Injection Revenue Forecast by Region (2027-2032) & (\$ millions)

Table 78. Americas Dalteparin Sodium Injection Sales Forecast by Country (2027-2032) & (K Units)

Table 79. Americas Dalteparin Sodium Injection Annual Revenue Forecast by Country (2027-2032) & (\$ millions)

Table 80. APAC Dalteparin Sodium Injection Sales Forecast by Region (2027-2032) & (K Units)

Table 81. APAC Dalteparin Sodium Injection Annual Revenue Forecast by Region (2027-2032) & (\$ millions)

Table 82. Europe Dalteparin Sodium Injection Sales Forecast by Country (2027-2032) & (K Units)

Table 83. Europe Dalteparin Sodium Injection Revenue Forecast by Country (2027-2032) & (\$ millions)

Table 84. Middle East & Africa Dalteparin Sodium Injection Sales Forecast by Country (2027-2032) & (K Units)

Table 85. Middle East & Africa Dalteparin Sodium Injection Revenue Forecast by Country (2027-2032) & (\$ millions)

Table 86. Global Dalteparin Sodium Injection Sales Forecast by Type (2027-2032) & (K Units)

Table 87. Global Dalteparin Sodium Injection Revenue Forecast by Type (2027-2032) & (\$ millions)

Table 88. Global Dalteparin Sodium Injection Sales Forecast by Application (2027-2032) & (K Units)

Table 89. Global Dalteparin Sodium Injection Revenue Forecast by Application (2027-2032) & (\$ millions)

Table 90. Pfizer Basic Information, Dalteparin Sodium Injection Manufacturing Base,

Sales Area and Its Competitors

Table 91. Pfizer Dalteparin Sodium Injection Product Portfolios and Specifications

Table 92. Pfizer Dalteparin Sodium Injection Sales (K Units), Revenue (\$ Million), Price (US\$/Unit) and Gross Margin (2021-2026)

Table 93. Pfizer Main Business

Table 94. Pfizer Latest Developments

Table 95. Hebei Changshan Biochemical Pharmaceutical Co.,Ltd. Basic Information, Dalteparin Sodium Injection Manufacturing Base, Sales Area and Its Competitors

Table 96. Hebei Changshan Biochemical Pharmaceutical Co.,Ltd. Dalteparin Sodium Injection Product Portfolios and Specifications

Table 97. Hebei Changshan Biochemical Pharmaceutical Co.,Ltd. Dalteparin Sodium Injection Sales (K Units), Revenue (\$ Million), Price (US\$/Unit) and Gross Margin (2021-2026)

Table 98. Hebei Changshan Biochemical Pharmaceutical Co.,Ltd. Main Business

Table 99. Hebei Changshan Biochemical Pharmaceutical Co.,Ltd. Latest Developments

Table 100. Nanjing King-Friend Biochemical Pharmaceutical Co., Ltd. Basic Information, Dalteparin Sodium Injection Manufacturing Base, Sales Area and Its Competitors

Table 101. Nanjing King-Friend Biochemical Pharmaceutical Co., Ltd. Dalteparin Sodium Injection Product Portfolios and Specifications

Table 102. Nanjing King-Friend Biochemical Pharmaceutical Co., Ltd. Dalteparin Sodium Injection Sales (K Units), Revenue (\$ Million), Price (US\$/Unit) and Gross Margin (2021-2026)

Table 103. Nanjing King-Friend Biochemical Pharmaceutical Co., Ltd. Main Business

Table 104. Nanjing King-Friend Biochemical Pharmaceutical Co., Ltd. Latest Developments

Table 105. Changzhou Qianhong Biopharma Co., Ltd. Basic Information, Dalteparin Sodium Injection Manufacturing Base, Sales Area and Its Competitors

Table 106. Changzhou Qianhong Biopharma Co., Ltd. Dalteparin Sodium Injection Product Portfolios and Specifications

Table 107. Changzhou Qianhong Biopharma Co., Ltd. Dalteparin Sodium Injection Sales (K Units), Revenue (\$ Million), Price (US\$/Unit) and Gross Margin (2021-2026)

Table 108. Changzhou Qianhong Biopharma Co., Ltd. Main Business

Table 109. Changzhou Qianhong Biopharma Co., Ltd. Latest Developments

Table 110. Qilu Pharmaceutical Basic Information, Dalteparin Sodium Injection Manufacturing Base, Sales Area and Its Competitors

Table 111. Qilu Pharmaceutical Dalteparin Sodium Injection Product Portfolios and Specifications

Table 112. Qilu Pharmaceutical Dalteparin Sodium Injection Sales (K Units), Revenue

(\$ Million), Price (US\$/Unit) and Gross Margin (2021-2026)

Table 113. Qilu Pharmaceutical Main Business

Table 114. Qilu Pharmaceutical Latest Developments

Table 115. Yantai Dongcheng Northern Pharmaceutical Basic Information, Dalteparin Sodium Injection Manufacturing Base, Sales Area and Its Competitors

Table 116. Yantai Dongcheng Northern Pharmaceutical Dalteparin Sodium Injection Product Portfolios and Specifications

Table 117. Yantai Dongcheng Northern Pharmaceutical Dalteparin Sodium Injection Sales (K Units), Revenue (\$ Million), Price (US\$/Unit) and Gross Margin (2021-2026)

Table 118. Yantai Dongcheng Northern Pharmaceutical Main Business

Table 119. Yantai Dongcheng Northern Pharmaceutical Latest Developments

List Of Figures

LIST OF FIGURES

- Figure 1. Picture of Dalteparin Sodium Injection
- Figure 2. Dalteparin Sodium Injection Report Years Considered
- Figure 3. Research Objectives
- Figure 4. Research Methodology
- Figure 5. Research Process and Data Source
- Figure 6. Global Dalteparin Sodium Injection Sales Growth Rate 2021-2032 (K Units)
- Figure 7. Global Dalteparin Sodium Injection Revenue Growth Rate 2021-2032 (\$ millions)
- Figure 8. Dalteparin Sodium Injection Sales by Geographic Region (2021, 2025 & 2032) & (\$ millions)
- Figure 9. Dalteparin Sodium Injection Sales Market Share by Country/Region (2025)
- Figure 10. Dalteparin Sodium Injection Sales Market Share by Country/Region (2021, 2025 & 2032)
- Figure 11. Product Picture of Dalteparin Sodium Injection 0.2ml:2500IU
- Figure 12. Product Picture of Dalteparin Sodium Injection 0.2ml:5000IU
- Figure 13. Product Picture of Dalteparin Sodium Injection 0.3ml:7500IU
- Figure 14. Product Picture of Dalteparin Sodium Injection 0.5ml:12500IU
- Figure 15. Product Picture of Dalteparin Sodium Injection 0.6ml:15000IU
- Figure 16. Global Dalteparin Sodium Injection Sales Market Share by Type in 2026
- Figure 17. Global Dalteparin Sodium Injection Revenue Market Share by Type (2021-2026)
- Figure 18. Product Picture of Prefilled Syringe
- Figure 19. Product Picture of Single-Dose Vial
- Figure 20. Global Dalteparin Sodium Injection Sales Market Share by Packaging in 2026
- Figure 21. Global Dalteparin Sodium Injection Revenue Market Share by Packaging (2021-2026)
- Figure 22. Product Picture of Porcine Intestinal Mucosa Origin
- Figure 23. Product Picture of Chemical Depolymerization
- Figure 24. Product Picture of Nitrous Acid Depolymerization
- Figure 25. Global Dalteparin Sodium Injection Sales Market Share by Manufacturing Process in 2026
- Figure 26. Global Dalteparin Sodium Injection Revenue Market Share by Manufacturing Process (2021-2026)
- Figure 27. Dalteparin Sodium Injection Consumed in Hospital

Figure 28. Global Dalteparin Sodium Injection Market: Hospital (2021-2026) & (K Units)

Figure 29. Dalteparin Sodium Injection Consumed in Nursing Institutions

Figure 30. Global Dalteparin Sodium Injection Market: Nursing Institutions (2021-2026) & (K Units)

Figure 31. Dalteparin Sodium Injection Consumed in Home Care

Figure 32. Global Dalteparin Sodium Injection Market: Home Care (2021-2026) & (K Units)

Figure 33. Dalteparin Sodium Injection Consumed in Others

Figure 34. Global Dalteparin Sodium Injection Market: Others (2021-2026) & (K Units)

Figure 35. Global Dalteparin Sodium Injection Sale Market Share by Application (2025)

Figure 36. Global Dalteparin Sodium Injection Revenue Market Share by Application in 2025

Figure 37. Dalteparin Sodium Injection Sales by Company in 2025 (K Units)

Figure 38. Global Dalteparin Sodium Injection Sales Market Share by Company in 2025

Figure 39. Dalteparin Sodium Injection Revenue by Company in 2025 (\$ millions)

Figure 40. Global Dalteparin Sodium Injection Revenue Market Share by Company in 2025

Figure 41. Global Dalteparin Sodium Injection Sales Market Share by Geographic Region (2021-2026)

Figure 42. Global Dalteparin Sodium Injection Revenue Market Share by Geographic Region in 2025

Figure 43. Americas Dalteparin Sodium Injection Sales 2021-2026 (K Units)

Figure 44. Americas Dalteparin Sodium Injection Revenue 2021-2026 (\$ millions)

Figure 45. APAC Dalteparin Sodium Injection Sales 2021-2026 (K Units)

Figure 46. APAC Dalteparin Sodium Injection Revenue 2021-2026 (\$ millions)

Figure 47. Europe Dalteparin Sodium Injection Sales 2021-2026 (K Units)

Figure 48. Europe Dalteparin Sodium Injection Revenue 2021-2026 (\$ millions)

Figure 49. Middle East & Africa Dalteparin Sodium Injection Sales 2021-2026 (K Units)

Figure 50. Middle East & Africa Dalteparin Sodium Injection Revenue 2021-2026 (\$ millions)

Figure 51. Americas Dalteparin Sodium Injection Sales Market Share by Country in 2025

Figure 52. Americas Dalteparin Sodium Injection Revenue Market Share by Country (2021-2026)

Figure 53. Americas Dalteparin Sodium Injection Sales Market Share by Type (2021-2026)

Figure 54. Americas Dalteparin Sodium Injection Sales Market Share by Application (2021-2026)

Figure 55. United States Dalteparin Sodium Injection Revenue Growth 2021-2026 (\$

millions)

Figure 56. Canada Dalteparin Sodium Injection Revenue Growth 2021-2026 (\$ millions)

Figure 57. Mexico Dalteparin Sodium Injection Revenue Growth 2021-2026 (\$ millions)

Figure 58. Brazil Dalteparin Sodium Injection Revenue Growth 2021-2026 (\$ millions)

Figure 59. APAC Dalteparin Sodium Injection Sales Market Share by Region in 2025

Figure 60. APAC Dalteparin Sodium Injection Revenue Market Share by Region
(2021-2026)

Figure 61. APAC Dalteparin Sodium Injection Sales Market Share by Type (2021-2026)

Figure 62. APAC Dalteparin Sodium Injection Sales Market Share by Application
(2021-2026)

Figure 63. China Dalteparin Sodium Injection Revenue Growth 2021-2026 (\$ millions)

Figure 64. Japan Dalteparin Sodium Injection Revenue Growth 2021-2026 (\$ millions)

Figure 65. South Korea Dalteparin Sodium Injection Revenue Growth 2021-2026 (\$
millions)

Figure 66. Southeast Asia Dalteparin Sodium Injection Revenue Growth 2021-2026 (\$
millions)

Figure 67. India Dalteparin Sodium Injection Revenue Growth 2021-2026 (\$ millions)

Figure 68. Australia Dalteparin Sodium Injection Revenue Growth 2021-2026 (\$
millions)

Figure 69. China Taiwan Dalteparin Sodium Injection Revenue Growth 2021-2026 (\$
millions)

Figure 70. Europe Dalteparin Sodium Injection Sales Market Share by Country in 2025

Figure 71. Europe Dalteparin Sodium Injection Revenue Market Share by Country
(2021-2026)

Figure 72. Europe Dalteparin Sodium Injection Sales Market Share by Type
(2021-2026)

Figure 73. Europe Dalteparin Sodium Injection Sales Market Share by Application
(2021-2026)

Figure 74. Germany Dalteparin Sodium Injection Revenue Growth 2021-2026 (\$
millions)

Figure 75. France Dalteparin Sodium Injection Revenue Growth 2021-2026 (\$ millions)

Figure 76. UK Dalteparin Sodium Injection Revenue Growth 2021-2026 (\$ millions)

Figure 77. Italy Dalteparin Sodium Injection Revenue Growth 2021-2026 (\$ millions)

Figure 78. Russia Dalteparin Sodium Injection Revenue Growth 2021-2026 (\$ millions)

Figure 79. Middle East & Africa Dalteparin Sodium Injection Sales Market Share by
Country (2021-2026)

Figure 80. Middle East & Africa Dalteparin Sodium Injection Sales Market Share by
Type (2021-2026)

Figure 81. Middle East & Africa Dalteparin Sodium Injection Sales Market Share by

Application (2021-2026)

Figure 82. Egypt Dalteparin Sodium Injection Revenue Growth 2021-2026 (\$ millions)

Figure 83. South Africa Dalteparin Sodium Injection Revenue Growth 2021-2026 (\$ millions)

Figure 84. Israel Dalteparin Sodium Injection Revenue Growth 2021-2026 (\$ millions)

Figure 85. Turkey Dalteparin Sodium Injection Revenue Growth 2021-2026 (\$ millions)

Figure 86. GCC Countries Dalteparin Sodium Injection Revenue Growth 2021-2026 (\$ millions)

Figure 87. Manufacturing Cost Structure Analysis of Dalteparin Sodium Injection in 2026

Figure 88. Manufacturing Process Analysis of Dalteparin Sodium Injection

Figure 89. Industry Chain Structure of Dalteparin Sodium Injection

Figure 90. Channels of Distribution

Figure 91. Global Dalteparin Sodium Injection Sales Market Forecast by Region (2027-2032)

Figure 92. Global Dalteparin Sodium Injection Revenue Market Share Forecast by Region (2027-2032)

Figure 93. Global Dalteparin Sodium Injection Sales Market Share Forecast by Type (2027-2032)

Figure 94. Global Dalteparin Sodium Injection Revenue Market Share Forecast by Type (2027-2032)

Figure 95. Global Dalteparin Sodium Injection Sales Market Share Forecast by Application (2027-2032)

Figure 96. Global Dalteparin Sodium Injection Revenue Market Share Forecast by Application (2027-2032)

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