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Global Trastuzumab Deruxtecan Supply, Demand and Key Producers, 2026-2032

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Abstracts

The global Trastuzumab Deruxtecan market size is expected to reach $ 11233 million by 2032, rising at a market growth of 9.4% CAGR during the forecast period (2026-2032).

Trastuzumab Deruxtecan is a HER2-directed antibody-drug conjugate composed of a humanized anti-HER2 IgG1 monoclonal antibody, a tetrapeptide-based cleavable linker, and the membrane-permeable topoisomerase I inhibitor payload DXd, with approximately eight payload molecules conjugated to each antibody. The commercial product is supplied as a 100 mg lyophilized powder in a single-dose vial for intravenous infusion, providing a concentration of 20 mg/mL after reconstitution. Treatment is generally administered once every three weeks, with 5.4 mg/kg used for breast cancer, HER2-mutant non-small cell lung cancer, and other eligible solid tumors, while 6.4 mg/kg is primarily used for HER2-positive gastric and gastroesophageal junction cancer. This study focuses on global commercial sales of finished Trastuzumab Deruxtecan products and analyzes the market by dosage regimen, treatment regimen, and cancer type, covering monotherapy, approved combination and sequential treatment settings, and major HER2-related cancer indications.

Key Findings

The 5.4 mg/kg every-three-week regimen represents the dominant commercial dose across most approved cancer settings

Breast cancer remains the largest cancer segment, with demand extending from metastatic treatment into first-line and early-stage care

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Current use remains centered on monotherapy, while combination and sequential regimens are becoming increasingly relevant

The US represented approximately 49% of 2025 combined global sales, confirming North America's leading regional position

The market is structured around one product platform and two core partner groups, with manufacturing concentrated under Daiichi Sankyo

Market Trends

The Trastuzumab Deruxtecan market is evolving from a predominantly later-line metastatic treatment market into a broader HER2-targeted therapeutic platform spanning earlier treatment stages, additional biomarker-defined populations, and multiple cancer types. In breast cancer, commercial demand is progressively extending from previously treated HER2-positive and HER2-low disease toward HER2-ultralow patients, first-line metastatic treatment, neoadjuvant therapy, and post-neoadjuvant treatment for patients with residual invasive disease. Treatment formats are also becoming more diversified. Monotherapy remains the central commercial model, but the approval of Trastuzumab Deruxtecan combined with pertuzumab and the use of Trastuzumab Deruxtecan followed by taxane, trastuzumab, and pertuzumab demonstrate a shift toward combination and sequential regimens. At the same time, HER2 testing is changing from a binary positive-or-negative model toward more differentiated identification of HER2-positive, HER2-low, HER2-ultralow, and HER2-mutant populations. This development is increasing the strategic importance of diagnostic accuracy, treatment sequencing, and real-world safety management across oncology centers.

Market Dynamics

Drivers

Market development is primarily driven by continued expansion of approved indications, movement into earlier and first-line treatment settings, and broader identification of patients who may benefit from HER2-directed therapy. Breast cancer contributes the largest demand base, while HER2-mutant non-small cell lung cancer, HER2-positive

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gastric cancer, and other HER2-positive solid tumors provide incremental growth. Improved HER2 testing and growing clinical recognition of HER2-low and HER2-ultralow disease are expanding the addressable population beyond traditional HER2-positive patients. Reimbursement expansion is another important driver, particularly in China, where inclusion in the national reimbursement system has accelerated access and patient adoption.

Restraints

The market remains constrained by the high total cost of treatment, intravenous administration requirements, specialist supervision, and differences in reimbursement and hospital access across countries. Commercial adoption also depends on reliable HER2 testing, appropriate patient selection, and the ability of oncology centers to manage adverse events. The manufacturing process is more complex than that of conventional monoclonal antibodies because it requires controlled antibody production, linker-payload synthesis, conjugation, purification, analytical characterization, and aseptic fill-finish. Concentration of manufacturing and supply under a single controlling group increases the importance of production capacity, quality consistency, and global inventory management.

Opportunities

The largest opportunities are expected to arise from earlier-stage breast cancer, first-line metastatic treatment, broader adoption in HER2-low and HER2-ultralow populations, and expansion into additional HER2-positive solid tumors. Combination treatment with pertuzumab and sequential use before surgery create new revenue pools beyond established monotherapy indications. Emerging markets also provide significant potential as regulatory approvals, reimbursement coverage, HER2 diagnostic capacity, and specialist oncology infrastructure improve. Additional opportunities may develop from greater use in patients with brain metastases, more standardized biomarker testing, improved treatment sequencing, and longer treatment duration in earlier disease settings.

Challenges

Long-term market development depends on balancing strong antitumor activity with safety and treatment tolerability. Interstitial lung disease or pneumonitis remains a major

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clinical risk requiring early identification, dose interruption, and permanent discontinuation in severe cases. Nausea, hematologic toxicity, cardiac monitoring requirements, and treatment-related discontinuation may also affect real-world treatment duration. The market will face increasing therapeutic competition from other HER2-directed ADCs, bispecific antibodies, alternative payload technologies, and new combination regimens. Differences in regulatory labels, diagnostic definitions, reimbursement conditions, and clinical practice among regions may further limit the speed at which new indications translate into commercial sales.

Industry Chain Analysis

The upstream segment of the Trastuzumab Deruxtecan industry chain includes mammalian cell culture media, recombinant antibody production materials, linker and cytotoxic payload intermediates, pharmaceutical excipients, chromatography media, filtration systems, single-use bioprocessing consumables, sterile glass vials, closures, and analytical testing reagents. HER2 immunohistochemistry and molecular diagnostic products also form an important supporting component because patient eligibility depends on accurate assessment of HER2 expression or ERBB2 mutation status. Upstream quality requirements are stringent, particularly for the high-potency DXd payload, conjugation intermediates, and materials that affect drug-to-antibody ratio, impurity profiles, aggregation, and product stability.

The midstream segment covers antibody production, linker-payload synthesis, controlled conjugation, purification, formulation, sterile filling, lyophilization, batch release, pharmacovigilance, and global supply management. Value creation is concentrated in proprietary ADC design, process control, analytical characterization, clinical evidence, regulatory approvals, and reliable commercial-scale manufacturing. The downstream segment includes regional marketing authorization holders, importers and distributors, public and commercial payers, HER2 diagnostic laboratories, general hospitals, cancer centers, and outpatient infusion facilities. Commercial profitability is influenced not only by manufacturing cost, but also by clinical development investment, regulatory expenditure, market-access discounts, profit-sharing arrangements, companion diagnostic availability, and the ability to maintain continuous supply across multiple treatment indications and regions.

Segment Insights

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By Dosage Regimen, the 5.4 mg/kg every-three-week segment represents the dominant market because it covers breast cancer, HER2-mutant non-small cell lung cancer, and most other approved solid tumor settings. The 6.4 mg/kg every-three-week segment is narrower and is primarily associated with HER2-positive gastric and gastroesophageal junction cancer. The commercial performance of the 5.4 mg/kg segment is therefore more closely linked to breast cancer treatment penetration, earlier treatment-line expansion, and the development of tumor-agnostic HER2-positive indications, while the 6.4 mg/kg segment is influenced mainly by gastric cancer diagnosis, prior trastuzumab exposure, and regional treatment practices.

By Treatment Regimen, monotherapy continues to account for the majority of current commercial use because most established indications were initially approved as single-agent treatment. Combination therapy is becoming more important following approval of Trastuzumab Deruxtecan with pertuzumab for first-line HER2-positive metastatic breast cancer. Sequential therapy is an emerging segment supported by neoadjuvant use of Trastuzumab Deruxtecan followed by taxane, trastuzumab, and pertuzumab. By Cancer Type, breast cancer remains the largest segment, while gastric and gastroesophageal junction cancer and non-small cell lung cancer form smaller but clinically established markets. Other HER2-positive solid tumors represent a diversified opportunity with comparatively limited individual cancer populations but broad cumulative potential.

Downstream Market Opportunities

Downstream market opportunities are concentrated in oncology treatment settings with established HER2 testing and intravenous infusion capabilities. Breast cancer centers offer the largest opportunity as treatment expands across HER2-positive, HER2-low, and HER2-ultralow disease and moves from advanced disease toward first-line and early-stage settings. Gastric and gastroesophageal junction cancer provide a differentiated opportunity in patients previously treated with trastuzumab-based regimens, while HER2-mutant non-small cell lung cancer benefits from the increasing adoption of genomic testing. The expansion of tumor-agnostic HER2-positive indications creates additional demand across smaller solid tumor categories. Hospitals and specialized cancer centers that can integrate pathology, molecular testing, multidisciplinary treatment decisions, infusion management, and adverse-event monitoring are expected to capture the greatest downstream value.

Regional Insights

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North America is the largest regional market, led by the US, which accounted for approximately 49% of combined global sales in 2025. Its leading position reflects earlier regulatory approvals, broad specialist adoption, established HER2 testing, and rapid uptake in HER2-low, HER2-ultralow, first-line metastatic, and early-stage breast cancer settings. Europe remains an important market supported by expanding reimbursement and approvals across breast cancer, gastric cancer, non-small cell lung cancer, and other HER2-positive solid tumors, although market access and reimbursement timing vary among countries.

Asia is becoming an increasingly important source of incremental demand. Japan benefits from early adoption and a broadening solid tumor application base, while China is supported by national reimbursement coverage, additional approved cancer indications, and expansion of hospital access. Emerging Markets recorded the fastest disclosed regional growth within AstraZeneca's booked geographies in 2025, supported in part by rapid adoption following reimbursement inclusion in China. Other Asia-Pacific, Latin American, Middle Eastern, and African markets remain at different stages of regulatory approval and reimbursement, creating long-term opportunities but also greater variability in affordability, diagnostic capacity, and market access.

Competitive Landscape Analysis

The Trastuzumab Deruxtecan market has a highly concentrated structure rather than a conventional multi-manufacturer competitive landscape. Daiichi Sankyo controls the proprietary DXd ADC platform and is responsible for manufacturing and global supply, while Daiichi Sankyo and AstraZeneca jointly develop and commercialize the product outside Japan; Daiichi Sankyo retains exclusive rights in Japan. The two companies therefore represent different but complementary positions within the same product ecosystem, and their reported product references should not be treated as separate drug models or duplicated market sales. Competitive differentiation is based on ADC technology, clinical evidence, regulatory coverage, manufacturing reliability, oncology commercialization capability, reimbursement access, and safety management. Direct competition increasingly occurs at the treatment-class level through other HER2-directed ADCs, antibody-based therapies, and combination regimens rather than through additional manufacturers of Trastuzumab Deruxtecan itself. Future competitive intensity will rise as alternative products seek differentiation through safety, activity in brain metastases, broader biomarker coverage, improved convenience, or lower treatment cost.

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Report Scope

This report studies the global Trastuzumab Deruxtecan production, demand, key manufacturers, and key regions.

This report is a detailed and comprehensive analysis of the world market for Trastuzumab Deruxtecan and provides market size (US$ million) and Year-over-Year (YoY) Growth, considering 2025 as the base year. This report explores demand trends and competition, as well as details the characteristics of Trastuzumab Deruxtecan that contribute to its increasing demand across many markets.

Highlights and key features of the study

Global Trastuzumab Deruxtecan total production and demand, 2021-2032, (K Units)

Global Trastuzumab Deruxtecan total production value, 2021-2032, (USD Million)

Global Trastuzumab Deruxtecan production by region & country, production, value, CAGR, 2021-2032, (USD Million) & (K Units), (based on production site)

Global Trastuzumab Deruxtecan consumption by region & country, CAGR, 2021-2032 & (K Units)

U.S. VS China: Trastuzumab Deruxtecan domestic production, consumption, key domestic manufacturers and share

Global Trastuzumab Deruxtecan production by manufacturer, production, price, value and market share 2021-2026, (USD Million) & (K Units)

Global Trastuzumab Deruxtecan production by Type, production, value, CAGR, 2021-2032, (USD Million) & (K Units)

Global Trastuzumab Deruxtecan production by Application, production, value, CAGR, 2021-2032, (USD Million) & (K Units)

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This report profiles key players in the global Trastuzumab Deruxtecan market based on the following parameters - company overview, production, value, price, gross margin, product portfolio, geographical presence, and key developments. Key companies covered as a part of this study include Daiichi Sankyo, AstraZeneca, etc.

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

Stakeholders would have ease in decision-making through various strategy matrices used in analyzing the World Trastuzumab Deruxtecan market

Detailed Segmentation:

Each section contains quantitative market data including market by value (US$ Millions), volume (production, consumption) & (K Units) and average price (US$/Unit) by manufacturer, by Type, and by Application. Data is given for the years 2021-2032 by year with 2025 as the base year, 2026 as the estimate year, and 2027-2032 as the forecast year.

Global Trastuzumab Deruxtecan Market, By Region:

United States

China

Europe

Japan

South Korea

ASEAN

India

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Rest of World

Market Segmentation

Global Trastuzumab Deruxtecan Market, Segmentation by Type:

5.4 mg/kg Every Three Weeks

6.4 mg/kg Every Three Weeks

Global Trastuzumab Deruxtecan Market, Segmentation by Treatment Regimen:

Monotherapy

Combination Therapy

Sequential Therapy

Global Trastuzumab Deruxtecan Market, Segmentation by Cancer:

Breast Cancer

Gastric and Gastroesophageal Junction Cancer

Non-Small Cell Lung Cancer

Other

Global Trastuzumab Deruxtecan Market, Segmentation by Application:

General Hospitals

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Cancer Hospitals

Other

Companies Profiled:

Daiichi Sankyo

AstraZeneca

Key Questions Answered

1. How big is the global Trastuzumab Deruxtecan market?
2. What is the demand of the global Trastuzumab Deruxtecan market?
3. What is the year over year growth of the global Trastuzumab Deruxtecan market?
4. What is the total value of the global Trastuzumab Deruxtecan market?
5. Who are the Major Players in the global Trastuzumab Deruxtecan market?
6. What are the growth factors driving the market demand?

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Contents

1 SUPPLY SUMMARY

1.1 Trastuzumab Deruxtecan Introduction
1.2 World Trastuzumab Deruxtecan Supply & Forecast
  1.2.1 World Trastuzumab Deruxtecan Production Value (2021 & 2025 & 2032)
  1.2.2 World Trastuzumab Deruxtecan Production (2021-2032)
  1.2.3 World Trastuzumab Deruxtecan Pricing Trends (2021-2032)
1.3 World Trastuzumab Deruxtecan Production by Region (Based on Production Site)
  1.3.1 World Trastuzumab Deruxtecan Production Value by Region (2021-2032)
  1.3.2 World Trastuzumab Deruxtecan Production by Region (2021-2032)
  1.3.3 World Trastuzumab Deruxtecan Average Price by Region (2021-2032)
  1.3.4 North America Trastuzumab Deruxtecan Production (2021-2032)
  1.3.5 Europe Trastuzumab Deruxtecan Production (2021-2032)
  1.3.6 China Trastuzumab Deruxtecan Production (2021-2032)
  1.3.7 Japan Trastuzumab Deruxtecan Production (2021-2032)
1.4 Market Drivers, Restraints and Trends
  1.4.1 Trastuzumab Deruxtecan Market Drivers
  1.4.2 Factors Affecting Demand
  1.4.3 Trastuzumab Deruxtecan Major Market Trends

2 DEMAND SUMMARY

2.1 World Trastuzumab Deruxtecan Demand (2021-2032)
2.2 World Trastuzumab Deruxtecan Consumption by Region
  2.2.1 World Trastuzumab Deruxtecan Consumption by Region (2021-2026)
  2.2.2 World Trastuzumab Deruxtecan Consumption Forecast by Region (2027-2032)
2.3 United States Trastuzumab Deruxtecan Consumption (2021-2032)
2.4 China Trastuzumab Deruxtecan Consumption (2021-2032)
2.5 Europe Trastuzumab Deruxtecan Consumption (2021-2032)
2.6 Japan Trastuzumab Deruxtecan Consumption (2021-2032)
2.7 South Korea Trastuzumab Deruxtecan Consumption (2021-2032)
2.8 ASEAN Trastuzumab Deruxtecan Consumption (2021-2032)
2.9 India Trastuzumab Deruxtecan Consumption (2021-2032)

3 WORLD MANUFACTURERS COMPETITIVE ANALYSIS

3.1 World Trastuzumab Deruxtecan Production Value by Manufacturer (2021-2026)

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info@marketpublishers.com

3.2 World Trastuzumab Deruxtecan Production by Manufacturer (2021-2026)
3.3 World Trastuzumab Deruxtecan Average Price by Manufacturer (2021-2026)
3.4 Trastuzumab Deruxtecan Company Evaluation Quadrant
3.5 Industry Rank and Concentration Rate (CR)
3.5.1 Global Trastuzumab Deruxtecan Industry Rank of Major Manufacturers
3.5.2 Global Concentration Ratios (CR4) for Trastuzumab Deruxtecan in 2025
3.5.3 Global Concentration Ratios (CR8) for Trastuzumab Deruxtecan in 2025
3.6 Trastuzumab Deruxtecan Market: Overall Company Footprint Analysis
3.6.1 Trastuzumab Deruxtecan Market: Region Footprint
3.6.2 Trastuzumab Deruxtecan Market: Company Product Type Footprint
3.6.3 Trastuzumab Deruxtecan Market: Company Product Application Footprint
3.7 Competitive Environment
3.7.1 Historical Structure of the Industry
3.7.2 Barriers of Market Entry
3.7.3 Factors of Competition
3.8 New Entrant and Capacity Expansion Plans
3.9 Mergers, Acquisition, Agreements, and Collaborations

4 UNITED STATES VS CHINA VS REST OF THE WORLD

4.1 United States VS China: Trastuzumab Deruxtecan Production Value Comparison
4.1.1 United States VS China: Trastuzumab Deruxtecan Production Value Comparison (2021 & 2025 & 2032)
4.1.2 United States VS China: Trastuzumab Deruxtecan Production Value Market Share Comparison (2021 & 2025 & 2032)
4.2 United States VS China: Trastuzumab Deruxtecan Production Comparison
4.2.1 United States VS China: Trastuzumab Deruxtecan Production Comparison (2021 & 2025 & 2032)
4.2.2 United States VS China: Trastuzumab Deruxtecan Production Market Share Comparison (2021 & 2025 & 2032)
4.3 United States VS China: Trastuzumab Deruxtecan Consumption Comparison
4.3.1 United States VS China: Trastuzumab Deruxtecan Consumption Comparison (2021 & 2025 & 2032)
4.3.2 United States VS China: Trastuzumab Deruxtecan Consumption Market Share Comparison (2021 & 2025 & 2032)
4.4 United States Based Trastuzumab Deruxtecan Manufacturers and Market Share, 2021-2026
4.4.1 United States Based Trastuzumab Deruxtecan Manufacturers, Headquarters and Production Site (States, Country)

+357 96 030922
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4.4.2 United States Based Manufacturers Trastuzumab Deruxtecan Production Value (2021-2026)
4.4.3 United States Based Manufacturers Trastuzumab Deruxtecan Production (2021-2026)
4.5 China Based Trastuzumab Deruxtecan Manufacturers and Market Share
4.5.1 China Based Trastuzumab Deruxtecan Manufacturers, Headquarters and Production Site (Province, Country)
4.5.2 China Based Manufacturers Trastuzumab Deruxtecan Production Value (2021-2026)
4.5.3 China Based Manufacturers Trastuzumab Deruxtecan Production (2021-2026)
4.6 Rest of World Based Trastuzumab Deruxtecan Manufacturers and Market Share, 2021-2026
4.6.1 Rest of World Based Trastuzumab Deruxtecan Manufacturers, Headquarters and Production Site (State, Country)
4.6.2 Rest of World Based Manufacturers Trastuzumab Deruxtecan Production Value (2021-2026)
4.6.3 Rest of World Based Manufacturers Trastuzumab Deruxtecan Production (2021-2026)

5 MARKET ANALYSIS BY TYPE

5.1 World Trastuzumab Deruxtecan Market Size Overview by Type: 2021 VS 2025 VS 2032
5.2 Segment Introduction by Type
5.2.1 5.4 mg/kg Every Three Weeks
5.2.2 6.4 mg/kg Every Three Weeks
5.3 Market Segment by Type
5.3.1 World Trastuzumab Deruxtecan Production by Type (2021-2032)
5.3.2 World Trastuzumab Deruxtecan Production Value by Type (2021-2032)
5.3.3 World Trastuzumab Deruxtecan Average Price by Type (2021-2032)

6 MARKET ANALYSIS BY TREATMENT REGIMEN

6.1 World Trastuzumab Deruxtecan Market Size Overview by Treatment Regimen: 2021 VS 2025 VS 2032
6.2 Segment Introduction by Treatment Regimen
6.2.1 Monotherapy
6.2.2 Combination Therapy
6.2.3 Sequential Therapy

+357 96 030922
info@marketpublishers.com

6.3 Market Segment by Treatment Regimen
6.3.1 World Trastuzumab Deruxtecan Production by Treatment Regimen (2021-2032)
6.3.2 World Trastuzumab Deruxtecan Production Value by Treatment Regimen (2021-2032)
6.3.3 World Trastuzumab Deruxtecan Average Price by Treatment Regimen (2021-2032)

7 MARKET ANALYSIS BY CANCER

7.1 World Trastuzumab Deruxtecan Market Size Overview by Cancer: 2021 VS 2025 VS 2032
7.2 Segment Introduction by Cancer
7.2.1 Breast Cancer
7.2.2 Gastric and Gastroesophageal Junction Cancer
7.2.3 Non-Small Cell Lung Cancer
7.2.4 Other
7.3 Market Segment by Cancer
7.3.1 World Trastuzumab Deruxtecan Production by Cancer (2021-2032)
7.3.2 World Trastuzumab Deruxtecan Production Value by Cancer (2021-2032)
7.3.3 World Trastuzumab Deruxtecan Average Price by Cancer (2021-2032)

8 MARKET ANALYSIS BY APPLICATION

8.1 World Trastuzumab Deruxtecan Market Size Overview by Application: 2021 VS 2025 VS 2032
8.2 Segment Introduction by Application
8.2.1 General Hospitals
8.2.2 Cancer Hospitals
8.2.3 Other
8.3 Market Segment by Application
8.3.1 World Trastuzumab Deruxtecan Production by Application (2021-2032)
8.3.2 World Trastuzumab Deruxtecan Production Value by Application (2021-2032)
8.3.3 World Trastuzumab Deruxtecan Average Price by Application (2021-2032)

9 COMPANY PROFILES

9.1 Daiichi Sankyo
9.1.1 Daiichi Sankyo Details
9.1.2 Daiichi Sankyo Major Business

+357 96 030922
info@marketpublishers.com

9.1.3 Daiichi Sankyo Trastuzumab Deruxtecan Product and Services
9.1.4 Daiichi Sankyo Trastuzumab Deruxtecan Production, Price, Value, Gross Margin and Market Share (2021-2026)
9.1.5 Daiichi Sankyo Recent Developments/Updates
9.1.6 Daiichi Sankyo Competitive Strengths & Weaknesses
9.2 AstraZeneca
9.2.1 AstraZeneca Details
9.2.2 AstraZeneca Major Business
9.2.3 AstraZeneca Trastuzumab Deruxtecan Product and Services
9.2.4 AstraZeneca Trastuzumab Deruxtecan Production, Price, Value, Gross Margin and Market Share (2021-2026)
9.2.5 AstraZeneca Recent Developments/Updates
9.2.6 AstraZeneca Competitive Strengths & Weaknesses

10 INDUSTRY CHAIN ANALYSIS

10.1 Trastuzumab Deruxtecan Industry Chain
10.2 Trastuzumab Deruxtecan Upstream Analysis
10.2.1 Trastuzumab Deruxtecan Core Raw Materials
10.2.2 Main Manufacturers of Trastuzumab Deruxtecan Core Raw Materials
10.3 Midstream Analysis
10.4 Downstream Analysis
10.5 Trastuzumab Deruxtecan Production Mode
10.6 Trastuzumab Deruxtecan Procurement Model
10.7 Trastuzumab Deruxtecan Industry Sales Model and Sales Channels
10.7.1 Trastuzumab Deruxtecan Sales Model
10.7.2 Trastuzumab Deruxtecan Typical Distributors

11 RESEARCH FINDINGS AND CONCLUSION

12 APPENDIX

12.1 Methodology
12.2 Research Process and Data Source
12.3 Disclaimer

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info@marketpublishers.com

List Of Tables

LIST OF TABLES

Table 1. World Trastuzumab Deruxtecan Production Value by Region (2021, 2025 and 2032) & (USD Million)
Table 2. World Trastuzumab Deruxtecan Production Value by Region (2021-2026) & (USD Million)
Table 3. World Trastuzumab Deruxtecan Production Value by Region (2027-2032) & (USD Million)
Table 4. World Trastuzumab Deruxtecan Production Value Market Share by Region (2021-2026)
Table 5. World Trastuzumab Deruxtecan Production Value Market Share by Region (2027-2032)
Table 6. World Trastuzumab Deruxtecan Production by Region (2021-2026) & (K Units)
Table 7. World Trastuzumab Deruxtecan Production by Region (2027-2032) & (K Units)
Table 8. World Trastuzumab Deruxtecan Production Market Share by Region (2021-2026)
Table 9. World Trastuzumab Deruxtecan Production Market Share by Region (2027-2032)
Table 10. World Trastuzumab Deruxtecan Average Price by Region (2021-2026) & (US$/Unit)
Table 11. World Trastuzumab Deruxtecan Average Price by Region (2027-2032) & (US$/Unit)
Table 12. Trastuzumab Deruxtecan Major Market Trends
Table 13. World Trastuzumab Deruxtecan Consumption Growth Rate Forecast by Region (2021 & 2025 & 2032) & (K Units)
Table 14. World Trastuzumab Deruxtecan Consumption by Region (2021-2026) & (K Units)
Table 15. World Trastuzumab Deruxtecan Consumption Forecast by Region (2027-2032) & (K Units)
Table 16. World Trastuzumab Deruxtecan Production Value by Manufacturer (2021-2026) & (USD Million)
Table 17. Production Value Market Share of Key Trastuzumab Deruxtecan Producers in 2025
Table 18. World Trastuzumab Deruxtecan Production by Manufacturer (2021-2026) & (K Units)
Table 19. Production Market Share of Key Trastuzumab Deruxtecan Producers in 2025
Table 20. World Trastuzumab Deruxtecan Average Price by Manufacturer (2021-2026)

+357 96 030922
info@marketpublishers.com

& (US$/Unit)
Table 21. Global Trastuzumab Deruxtecan Company Evaluation Quadrant
Table 22. World Trastuzumab Deruxtecan Industry Rank of Major Manufacturers, Based on Production Value in 2025
Table 23. Head Office and Trastuzumab Deruxtecan Production Site of Key Manufacturer
Table 24. Trastuzumab Deruxtecan Market: Company Product Type Footprint
Table 25. Trastuzumab Deruxtecan Market: Company Product Application Footprint
Table 26. Trastuzumab Deruxtecan Competitive Factors
Table 27. Trastuzumab Deruxtecan New Entrant and Capacity Expansion Plans
Table 28. Trastuzumab Deruxtecan Mergers & Acquisitions Activity
Table 29. United States VS China Trastuzumab Deruxtecan Production Value Comparison, (2021 & 2025 & 2032) & (USD Million)
Table 30. United States VS China Trastuzumab Deruxtecan Production Comparison, (2021 & 2025 & 2032) & (K Units)
Table 31. United States VS China Trastuzumab Deruxtecan Consumption Comparison, (2021 & 2025 & 2032) & (K Units)
Table 32. United States Based Trastuzumab Deruxtecan Manufacturers, Headquarters and Production Site (States, Country)
Table 33. United States Based Manufacturers Trastuzumab Deruxtecan Production Value, (2021-2026) & (USD Million)
Table 34. United States Based Manufacturers Trastuzumab Deruxtecan Production Value Market Share (2021-2026)
Table 35. United States Based Manufacturers Trastuzumab Deruxtecan Production (2021-2026) & (K Units)
Table 36. United States Based Manufacturers Trastuzumab Deruxtecan Production Market Share (2021-2026)
Table 37. China Based Trastuzumab Deruxtecan Manufacturers, Headquarters and Production Site (Province, Country)
Table 38. China Based Manufacturers Trastuzumab Deruxtecan Production Value, (2021-2026) & (USD Million)
Table 39. China Based Manufacturers Trastuzumab Deruxtecan Production Value Market Share (2021-2026)
Table 40. China Based Manufacturers Trastuzumab Deruxtecan Production, (2021-2026) & (K Units)
Table 41. China Based Manufacturers Trastuzumab Deruxtecan Production Market Share (2021-2026)
Table 42. Rest of World Based Trastuzumab Deruxtecan Manufacturers, Headquarters and Production Site (State, Country)

+357 96 030922
info@marketpublishers.com

Table 43. Rest of World Based Manufacturers Trastuzumab Deruxtecan Production Value, (2021-2026) & (USD Million)
Table 44. Rest of World Based Manufacturers Trastuzumab Deruxtecan Production Value Market Share (2021-2026)
Table 45. Rest of World Based Manufacturers Trastuzumab Deruxtecan Production, (2021-2026) & (K Units)
Table 46. Rest of World Based Manufacturers Trastuzumab Deruxtecan Production Market Share (2021-2026)
Table 47. World Trastuzumab Deruxtecan Production Value by Type, (USD Million), 2021 & 2025 & 2032
Table 48. World Trastuzumab Deruxtecan Production by Type (2021-2026) & (K Units)
Table 49. World Trastuzumab Deruxtecan Production by Type (2027-2032) & (K Units)
Table 50. World Trastuzumab Deruxtecan Production Value by Type (2021-2026) & (USD Million)
Table 51. World Trastuzumab Deruxtecan Production Value by Type (2027-2032) & (USD Million)
Table 52. World Trastuzumab Deruxtecan Average Price by Type (2021-2026) & (US$/Unit)
Table 53. World Trastuzumab Deruxtecan Average Price by Type (2027-2032) & (US$/Unit)
Table 54. World Trastuzumab Deruxtecan Production Value by Treatment Regimen, (USD Million), 2021 & 2025 & 2032
Table 55. World Trastuzumab Deruxtecan Production by Treatment Regimen (2021-2026) & (K Units)
Table 56. World Trastuzumab Deruxtecan Production by Treatment Regimen (2027-2032) & (K Units)
Table 57. World Trastuzumab Deruxtecan Production Value by Treatment Regimen (2021-2026) & (USD Million)
Table 58. World Trastuzumab Deruxtecan Production Value by Treatment Regimen (2027-2032) & (USD Million)
Table 59. World Trastuzumab Deruxtecan Average Price by Treatment Regimen (2021-2026) & (US$/Unit)
Table 60. World Trastuzumab Deruxtecan Average Price by Treatment Regimen (2027-2032) & (US$/Unit)
Table 61. World Trastuzumab Deruxtecan Production Value by Cancer, (USD Million), 2021 & 2025 & 2032
Table 62. World Trastuzumab Deruxtecan Production by Cancer (2021-2026) & (K Units)
Table 63. World Trastuzumab Deruxtecan Production by Cancer (2027-2032) & (K

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Units)
Table 64. World Trastuzumab Deruxtecan Production Value by Cancer (2021-2026) & (USD Million)
Table 65. World Trastuzumab Deruxtecan Production Value by Cancer (2027-2032) & (USD Million)
Table 66. World Trastuzumab Deruxtecan Average Price by Cancer (2021-2026) & (US$/Unit)
Table 67. World Trastuzumab Deruxtecan Average Price by Cancer (2027-2032) & (US$/Unit)
Table 68. World Trastuzumab Deruxtecan Production Value by Application, (USD Million), 2021 & 2025 & 2032
Table 69. World Trastuzumab Deruxtecan Production by Application (2021-2026) & (K Units)
Table 70. World Trastuzumab Deruxtecan Production by Application (2027-2032) & (K Units)
Table 71. World Trastuzumab Deruxtecan Production Value by Application (2021-2026) & (USD Million)
Table 72. World Trastuzumab Deruxtecan Production Value by Application (2027-2032) & (USD Million)
Table 73. World Trastuzumab Deruxtecan Average Price by Application (2021-2026) & (US$/Unit)
Table 74. World Trastuzumab Deruxtecan Average Price by Application (2027-2032) & (US$/Unit)
Table 75. Daiichi Sankyo Basic Information, Manufacturing Base and Competitors
Table 76. Daiichi Sankyo Major Business
Table 77. Daiichi Sankyo Trastuzumab Deruxtecan Product and Services
Table 78. Daiichi Sankyo Trastuzumab Deruxtecan Production (K Units), Price (US$/Unit), Production Value (USD Million), Gross Margin and Market Share (2021-2026)
Table 79. Daiichi Sankyo Recent Developments/Updates
Table 80. Daiichi Sankyo Competitive Strengths & Weaknesses
Table 81. AstraZeneca Basic Information, Manufacturing Base and Competitors
Table 82. AstraZeneca Major Business
Table 83. AstraZeneca Trastuzumab Deruxtecan Product and Services
Table 84. AstraZeneca Trastuzumab Deruxtecan Production (K Units), Price (US$/Unit), Production Value (USD Million), Gross Margin and Market Share (2021-2026)
Table 85. AstraZeneca Recent Developments/Updates
Table 86. AstraZeneca Competitive Strengths & Weaknesses
Table 87. Global Key Players of Trastuzumab Deruxtecan Upstream (Raw Materials)

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Table 88. Global Trastuzumab Deruxtecan Typical Customers
Table 89. Trastuzumab Deruxtecan Typical Distributors

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List Of Figures

LIST OF FIGURES

Figure 1. Trastuzumab Deruxtecan Picture
Figure 2. World Trastuzumab Deruxtecan Production Value: 2021 & 2025 & 2032, (USD Million)
Figure 3. World Trastuzumab Deruxtecan Production Value and Forecast (2021-2032) & (USD Million)
Figure 4. World Trastuzumab Deruxtecan Production (2021-2032) & (K Units)
Figure 5. World Trastuzumab Deruxtecan Average Price (2021-2032) & (US$/Unit)
Figure 6. World Trastuzumab Deruxtecan Production Value Market Share by Region (2021-2032)
Figure 7. World Trastuzumab Deruxtecan Production Market Share by Region (2021-2032)
Figure 8. North America Trastuzumab Deruxtecan Production (2021-2032) & (K Units)
Figure 9. Europe Trastuzumab Deruxtecan Production (2021-2032) & (K Units)
Figure 10. China Trastuzumab Deruxtecan Production (2021-2032) & (K Units)
Figure 11. Japan Trastuzumab Deruxtecan Production (2021-2032) & (K Units)
Figure 12. Trastuzumab Deruxtecan Market Drivers
Figure 13. Factors Affecting Demand
Figure 14. World Trastuzumab Deruxtecan Consumption (2021-2032) & (K Units)
Figure 15. World Trastuzumab Deruxtecan Consumption Market Share by Region (2021-2032)
Figure 16. United States Trastuzumab Deruxtecan Consumption (2021-2032) & (K Units)
Figure 17. China Trastuzumab Deruxtecan Consumption (2021-2032) & (K Units)
Figure 18. Europe Trastuzumab Deruxtecan Consumption (2021-2032) & (K Units)
Figure 19. Japan Trastuzumab Deruxtecan Consumption (2021-2032) & (K Units)
Figure 20. South Korea Trastuzumab Deruxtecan Consumption (2021-2032) & (K Units)
Figure 21. ASEAN Trastuzumab Deruxtecan Consumption (2021-2032) & (K Units)
Figure 22. India Trastuzumab Deruxtecan Consumption (2021-2032) & (K Units)
Figure 23. Producer Shipments of Trastuzumab Deruxtecan by Manufacturer Revenue ($MM) and Market Share (%): 2025
Figure 24. Global Four-firm Concentration Ratios (CR4) for Trastuzumab Deruxtecan Markets in 2025
Figure 25. Global Four-firm Concentration Ratios (CR8) for Trastuzumab Deruxtecan Markets in 2025
Figure 26. United States VS China: Trastuzumab Deruxtecan Production Value Market

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Share Comparison (2021 & 2025 & 2032)
Figure 27. United States VS China: Trastuzumab Deruxtecan Production Market Share Comparison (2021 & 2025 & 2032)
Figure 28. United States VS China: Trastuzumab Deruxtecan Consumption Market Share Comparison (2021 & 2025 & 2032)
Figure 29. United States Based Manufacturers Trastuzumab Deruxtecan Production Market Share 2025
Figure 30. China Based Manufacturers Trastuzumab Deruxtecan Production Market Share 2025
Figure 31. Rest of World Based Manufacturers Trastuzumab Deruxtecan Production Market Share 2025
Figure 32. World Trastuzumab Deruxtecan Production Value by Type, (USD Million), 2021 & 2025 & 2032
Figure 33. World Trastuzumab Deruxtecan Production Value Market Share by Type in 2025
Figure 34. 5.4 mg/kg Every Three Weeks
Figure 35. 6.4 mg/kg Every Three Weeks
Figure 36. World Trastuzumab Deruxtecan Production Market Share by Type (2021-2032)
Figure 37. World Trastuzumab Deruxtecan Production Value Market Share by Type (2021-2032)
Figure 38. World Trastuzumab Deruxtecan Average Price by Type (2021-2032) & (US$/Unit)
Figure 39. World Trastuzumab Deruxtecan Production Value by Treatment Regimen, (USD Million), 2021 & 2025 & 2032
Figure 40. World Trastuzumab Deruxtecan Production Value Market Share by Treatment Regimen in 2025
Figure 41. Monotherapy
Figure 42. Combination Therapy
Figure 43. Sequential Therapy
Figure 44. World Trastuzumab Deruxtecan Production Market Share by Treatment Regimen (2021-2032)
Figure 45. World Trastuzumab Deruxtecan Production Value Market Share by Treatment Regimen (2021-2032)
Figure 46. World Trastuzumab Deruxtecan Average Price by Treatment Regimen (2021-2032) & (US$/Unit)
Figure 47. World Trastuzumab Deruxtecan Production Value by Cancer, (USD Million), 2021 & 2025 & 2032
Figure 48. World Trastuzumab Deruxtecan Production Value Market Share by Cancer in

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2025
Figure 49. Breast Cancer
Figure 50. Gastric and Gastroesophageal Junction Cancer
Figure 51. Non-Small Cell Lung Cancer
Figure 52. Other
Figure 53. World Trastuzumab Deruxtecan Production Market Share by Cancer (2021-2032)
Figure 54. World Trastuzumab Deruxtecan Production Value Market Share by Cancer (2021-2032)
Figure 55. World Trastuzumab Deruxtecan Average Price by Cancer (2021-2032) & (US$/Unit)
Figure 56. World Trastuzumab Deruxtecan Production Value by Application, (USD Million), 2021 & 2025 & 2032
Figure 57. World Trastuzumab Deruxtecan Production Value Market Share by Application in 2025
Figure 58. General Hospitals
Figure 59. Cancer Hospitals
Figure 60. Other
Figure 61. World Trastuzumab Deruxtecan Production Market Share by Application (2021-2032)
Figure 62. World Trastuzumab Deruxtecan Production Value Market Share by Application (2021-2032)
Figure 63. World Trastuzumab Deruxtecan Average Price by Application (2021-2032) & (US$/Unit)
Figure 64. Trastuzumab Deruxtecan Industry Chain
Figure 65. Trastuzumab Deruxtecan Procurement Model
Figure 66. Trastuzumab Deruxtecan Sales Model
Figure 67. Trastuzumab Deruxtecan Sales Channels, Direct Sales, and Distribution
Figure 68. Methodology
Figure 69. Research Process and Data Source

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