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Global Exosome CDMO Market Growth (Status and Outlook) 2026-2032

https://marketpublishers.com/r/G3E4EDE07E4FEN.html
Date: August 2026
Pages: 132
Price: US$ 3,660.00 (Single User License)
ID: G3E4EDE07E4FEN

Abstracts

The global Exosome CDMO market size is predicted to grow from US$ 623 million in 2025 to US$ 1130 million in 2032; it is expected to grow at a CAGR of 9.0% from 2026 to 2032.

Exosome CDMO refers to providing customized R&D and manufacturing services for biopharmaceutical companies across the entire exosome chain, from process development to large-scale production. Its core businesses encompass the construction and culture of exosome production cell lines, development of large-scale exosome extraction and purification processes, rigorous quality control and analysis (including purity, drug loading, and biological function), formulation development, and GMP-compliant manufacturing. This service aims to help clients address the technical, process scale-up, and compliance challenges they face in developing exosomes as drug carriers or therapeutic products, accelerating the translation from laboratory research to preclinical and clinical trials. It is an indispensable key support platform for the industrialization of exosome therapies.

The global Exosome CDMO market exhibits a distinct pattern of "technology leadership, regional concentration, and the rise of the Asia-Pacific region." North America (especially the United States) currently holds an absolute leading position, possessing the most mature GMP production system, the most cutting-edge engineering technology platform, and a dense customer base of early-stage biotechnology companies, dominating the global early-stage clinical trial supply chain. Europe, with its strong foundation in cell and gene therapy, boasts robust capabilities in process development and quality systems. The Asia-Pacific region (represented by China and South Korea) is the fastest-growing market globally. Leveraging its mature biopharmaceutical manufacturing ecosystem, relatively flexible service models, and rapid technological

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advancements, it is actively building a service system from process development to localized production, aiming to become a key node in the regional and even global supply chain. The industry as a whole is still in its early stages, rapidly evolving from fragmented research services to a standardized, large-scale pharmaceutical-grade industry.

LPI (LP Information)' newest research report, the “Exosome CDMO Industry Forecast” looks at past sales and reviews total world Exosome CDMO sales in 2025, providing a comprehensive analysis by region and market sector of projected Exosome CDMO sales for 2026 through 2032. With Exosome CDMO sales broken down by region, market sector and sub-sector, this report provides a detailed analysis in US$ millions of the world Exosome CDMO industry.

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

This Insight Report evaluates the key market trends, drivers, and affecting factors shaping the global outlook for Exosome CDMO 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 Exosome CDMO.

This report presents a comprehensive overview, market shares, and growth opportunities of Exosome CDMO market by product type, application, key players and key regions and countries.

Segmentation by Type:

Preclinical CRO

Clinical CRO

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Segmentation by Service Stage:

- Process Development Type
- GMP Production Type
- Analytical Testing and Quality Control Type

Segmentation by Exosome Source:

- Natural Exosome CDMO
- Engineered Exosome CDMO

Segmentation by Application:

- Laboratory
- Universities
- Bio Industry
- Medical Industry

This report also splits the market by region:

- Americas
  - United States
  - Canada
  - Mexico
  - Brazil

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APAC

China

Japan

Korea

Southeast Asia

India

Australia

Europe

Germany

France

UK

Italy

Russia

Middle East & Africa

Egypt

South Africa

Israel

Turkey

GCC Countries

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

REPROCELL

Histocell

Lonza

MYCENAX

RoosterBio

Altaris LLC

Asahi Glass Co.

Fortis Life Sciences

Scorpius BioManufacturing

Miltenyi Bioindustry

Exothera

EXO Biologics

Porton Advanced

Elise Biopharma

Evox Therapeutics

FUJIFILM Biotechnologies

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