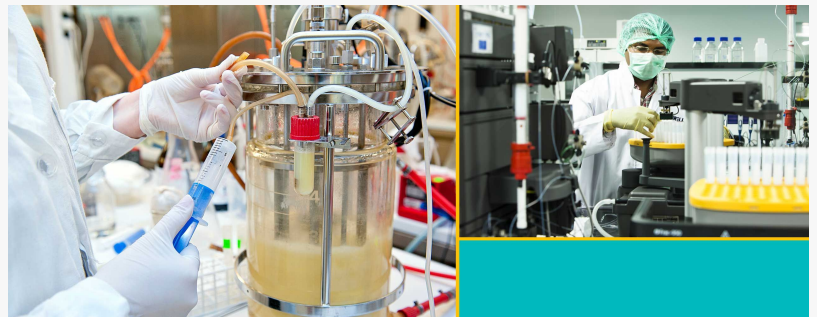


Balancing Progress: The Dynamic Landscape of the Biopharma Buffer Market 2023-2030

| Becton, Lonza Group Ltd, Avantor

biopharma buffer market size was valued at US\$ 3.44 billion in 2022 and is expected to grow at a CAGR of 7.1% to reach US\$ 5.95 billion by end of 2030

BURLINGAME, CALIFORNIA , UNITED STATES, November 24, 2023
/EINPresswire.com/ -- Market Overview:



Biopharma Buffer Market

Biopharma buffers are solutions used to maintain optimum pH levels during biopharmaceutical drug manufacturing and purification processes. They help stabilize products and aid cell growth.

Market Dynamics:

Biopharma buffer market growth is driven by rising biologics production and stringent regulatory standards for optimal pH maintenance. According to market estimates, biologics currently account for over 30% of global drug pipelines. Their large-scale manufacturing requires precise pH and ion balancing at every step, driving buffer consumption. Additionally, regulatory mandates have strengthened over time, with the FDA and EMA enforcing tighter pH specifications of 2.0-3.0 for bioprocessing. This has compelled biopharma companies to rely more on robust and traceable buffering solutions to prevent product failures due to pH drift. With the biologics segment continuing strong growth over the forecast period, demand for associated buffers is also expected to increase significantly.

Increasing investment in R&D boosting biopharmaceutical buffer market growth

The biopharma buffer market has seen significant growth in recent years due to increasing investments made by various companies in research and development activities related to drug discovery and development. Biopharmaceutical companies are allocating large portions of their budgets to R&D to develop novel drug formulations and treatment options for a wide range of complex diseases. Biopharma buffers play a vital role during various stages of drug

manufacturing such as purification, stabilization and transportation of biologic drugs. Therefore, the expanding R&D pipeline of biologic drugs targeting cancers, infectious diseases and rare disorders has augmented the demand for high-quality buffers. Additionally, growing trend of outsourcing manufacturing activities to specialized contract manufacturing organizations is also contributing towards market growth as they require a reliable supply of buffers.

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Stringent regulatory frameworks hampering market expansion

Although the biopharma buffer market is witnessing profitable opportunities, regulatory frameworks set by several agencies have posed challenges for manufacturers.

Biopharmaceutical buffers are classified as critical raw materials and their production and quality is tightly regulated by authorities like the FDA and EMA. Buffers need to undergo rigorous validation testing and meet stringent criteria related to purity, sterility and compatibility with drug formulations. Failing to comply with guidelines can delay new product approvals or lead to recalls. This increases compliance costs for companies. Additionally, changes in specifications can invalidate previous validation data requiring re-validation. Meeting evolving regulatory standards is difficult and resource intensive, slowing the introduction of novel buffer solutions. However, widespread acceptance of quality management systems and regional harmonization of regulations can help alleviate these restraints over time.

Shift towards single-use technologies opens new avenues

The biopharma industry has transitioned significantly towards adoption of single-use technologies over the last decade in an effort to improve flexibility, efficiency and reduce costs. Single-use bioreactors, stainless-steel-free trains and disposable chromatography columns are widely replacing conventional large-scale stainless equipment across various stages of bioproduction. This has created high demand for single-use compatible buffers without leachable and extractable issues. Leading buffer developers are increasingly offering specialized buffer formulations that have been optimized and validated for applications involving single-use systems. Their added benefits include eliminating cleaning and contamination risks involved in re-usable systems. Widespread integration of single-use assemblies within continuous manufacturing workflows is projected to provide further impetus to this emerging opportunity area.

Growing preference for multifunctional excipients

The buffer market is also observing a notable shift towards development and uptake of multifunctional excipients having double or triple buffering capabilities. Traditionally separate buffers were used during different unit operations such as fermentation, purification and filling. However, use of a single excipient performing buffering as well as other utility functions can help

simplify downstream processes and lower production costs. Leading vendors are actively focusing on designing buffers with properties like stabilization, osmolality adjustment and metal ion chelation in addition to pH control. Their advantages over conventional buffers include reduced formulation complexity, improved compatibility with single-use technologies and lower environmental footprint. It is expected that ongoing R&D aimed at exploiting synergistic functionalities of excipients will gain more attention going forward.

Moreover, it will also include the opportunities available in micro markets for stakeholders to invest, a detailed analysis of the competitive landscape, and product services of key players. Analysis of Biopharma Buffer companies, key tactics followed by Leading Key Players:

- Merck KGaA
- Thermo Fisher Scientific Inc.
- Avantor Inc.
- Lonza Group Ltd.
- Bio-Rad Laboratories Inc.
- Sartorius AG
- Corning Inc.
- Becton
- Dickinson and Company
- GE Healthcare
- Promega Corporation

Note: Major Players are sorted in no particular order.

By Type:

Pre-formulated buffers
Customized buffers
Concentrated buffers
Others

By Application:

Cell Culture
Purification
Formulation

By End User:

Biopharmaceutical Companies
Contract Research Organizations
Academic & Research Institutes

By Buffer Component:

Amino acids
Acetic acid
Phosphate
Histidine
Others

By Buffer Preparation:

Liquid
Powder

By Material Form:

Dry
Liquid

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The Study Objectives are:

A comprehensive insight into key players operating in the Biopharma Buffer Market and their corresponding data.

It includes product portfolio, annual revenue, expenditure on research and development, geographical presence, key developments in recent years, and growth strategies.

Regional analysis, which includes insight into the dominant market and corresponding market share.

It also includes various socio-economic factors affecting the evolution of the market in the region.

The report offers a comprehensive insight into different individuals from value chains such as raw materials suppliers, distributors, and stockholders.

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Key Questions Answered:

What is the market size and CAGR of the Biopharma Buffer Market during the forecast period?

How is the growing demand impacting the growth of Biopharma Buffer Market shares?

What is the growing demand of the Market during the forecast period?

Who are the leading vendors in the market and what are their market shares?

What is the impact of the COVID-19 pandemic on the APAC Biopharma Buffer Market?

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