

Global Single-Use Biopsy Device Market Growth 2026-2032

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Abstracts

The global Single-Use Biopsy Device market size is predicted to grow from US\$ 1537 million in 2025 to US\$ 2976 million in 2032; it is expected to grow at a CAGR of 9.9% from 2026 to 2032.

Single-use biopsy devices are disposable biopsy instruments and consumable sets intended for one patient and one procedure, including single-use biopsy needles (FNA/CNB), disposable components for vacuum-assisted biopsy systems, endoscopic/bronchoscopic disposable biopsy forceps and sampling tools, and disposable punch or core biopsy instruments for skin and soft tissue. Their value proposition is to replace the complexity of cleaning, disinfection/sterilization, and reuse governance with a sterile, ready-to-use workflow—reducing cross-contamination risk and reprocessing failures while improving throughput and specimen consistency. As image-guided biopsy, molecular testing, and precision diagnostics raise expectations for specimen integrity and traceability, single-use adoption is increasingly tied to hospital quality systems, compliance, and pathway standardization. In 2025, global Single-use biopsy devices production reached approximately 52.37 Million Units and price is about 30 USD/Unit. The average gross profit margin of this product is 45%.

Adoption is propelled by a dual mandate: patient safety and workflow certainty. Infection prevention and quality governance emphasize that single-use devices are intended for one patient and one procedure, with disposal after use—making disposable biopsy workflows an intuitive way to reduce cross-contamination risk and simplify reprocessing burden and staff training. In parallel, the regulatory landscape around reuse and reprocessing has become more systematized, prompting healthcare facilities to reassess the “true total cost” of reuse under compliance—strengthening the strategic appeal of disposable solutions.

The competitive bar is shifting from “disposable by design” to “diagnostic-grade performance at scale.” Across anatomies and image-guided pathways, clinicians demand consistent tactile control, needle-tip geometry, tissue yield, specimen integrity, and complication management—where engineering reproducibility becomes the decisive differentiator. Meanwhile, reusable instruments and FDA-regulated reprocessing of certain single-use devices remain cost strategies in some facilities, requiring nuanced trade-offs among compliance, quality, and economics; regulators also expect validated reprocessing instructions and clear labeling where applicable, raising the importance of auditable documentation and verification systems.

Demand is moving from “consumable purchasing” toward pathway-level kits. Image-guided biopsy and multidisciplinary care favor standardized sampling workflows, pushing providers to prefer disposable systems that integrate smoothly with imaging platforms and downstream pathology/molecular lab requirements. As outpatient and day-case models expand, throughput, predictability, and supply assurance rise in priority—where single-use solutions gain advantage by removing reprocessing steps. For department leaders, traceability, training simplicity, and quality-control governance are becoming procurement criteria as influential as unit price.

Upstream value is defined by a combined capability in medical-grade materials, precision manufacturing, and sterile barrier systems. Typical inputs include stainless steel or nitinol cannulas and cutting components, engineered plastics/polymers for handles and sheaths, micro-springs and actuation mechanisms, lubricious/surface treatments, and sterile packaging materials (blister packs, medical paper, Tyvek) with validated sterilization services. Because disposable biopsy devices depend on batch-to-batch consistency and sterility assurance, supply chains must deliver rigorous traceability, sterilization validation, and labeling compliance; in the EU, UDI-driven traceability further elevates labeling and supply-chain digitalization as core capabilities.

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

This Insight Report provides a comprehensive analysis of the global Single-Use Biopsy

Device 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 Single-Use Biopsy Device portfolios and capabilities, market entry strategies, market positions, and geographic footprints, to better understand these firms' unique position in an accelerating global Single-Use Biopsy Device market.

This Insight Report evaluates the key market trends, drivers, and affecting factors shaping the global outlook for Single-Use Biopsy Device 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 Single-Use Biopsy Device.

This report presents a comprehensive overview, market shares, and growth opportunities of Single-Use Biopsy Device market by product type, application, key manufacturers and key regions and countries.

Segmentation by Type:

Disposable Biopsy Needle

Disposable Biopsy Forceps

Disposable Puncture Sampler

Other

Segmentation by Target Anatomy:

Breast

Lung / Bronchoscopy

GI / pancreatobiliary

Other

Segmentation by Specimen Type:

Cytology

Histology Core/Tissue

Other

Segmentation by Application:

Hospital

ASCs

Other

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.

BD

Mammutome

Medtronic

Merit Medical Systems

Cook

Boston Scientific

Olympus

ConMed

Micro-tech

KDL

Medax

Sterylab

Hologic

Weigao

Puang Medical

Shanghai Aisiai

Key Questions Addressed in this Report

What is the 10-year outlook for the global Single-Use Biopsy Device market?

What factors are driving Single-Use Biopsy Device market growth, globally and by region?

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

How do Single-Use Biopsy Device market opportunities vary by end market size?

How does Single-Use Biopsy Device break out by Type, by Application?

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