

Global Medical Device Viral Inactivation Testing Market Growth (Status and Outlook) 2026-2032

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Abstracts

The global Medical Device Viral Inactivation Testing market size is predicted to grow from US\$ 196 million in 2025 to US\$ 536 million in 2032; it is expected to grow at a CAGR of 15.6% from 2026 to 2032.

Medical Device Viral Inactivation Testing is a service project specifically designed for medical devices that come into contact with biological fluids or tissues. This testing ensures that the device effectively eliminates or inactivates viruses, thereby preventing the transmission of infections during medical procedures. The test involves subjecting the device to a controlled environment with a known virus concentration and assessing its ability to reduce the viral load to an acceptable level. The significance of Medical Device Viral Inactivation Testing lies in its role in maintaining patient safety and preventing healthcare-associated infections. The functional benefits of this testing include the validation of the device's effectiveness in inactivating viruses, which is crucial for devices intended for reuse or those that come into direct contact with patients. By confirming the device's viral inactivation capabilities, this testing service contributes to the overall quality and reliability of medical devices.

In the future, the development trends of the Medical Device Viral Inactivation Testing industry will primarily manifest in the innovation of testing technologies and methods, updates to testing standards, the development of personalized testing solutions, the enhancement of automation and intelligence in testing processes, the advancement of internationalization and standardization, improvements in cost-effectiveness, changes in market demand, and shifts in the regulatory environment. With advancements in science and technology, new testing technologies and methods will be developed to enhance the accuracy and efficiency of viral inactivation testing, adapting to the characteristics of different medical devices and viruses. Automated and intelligent

testing processes will utilize advanced automated equipment and analytical software to improve accuracy and efficiency while reducing human errors. The advancement of internationalization and standardization will ensure global recognition of testing results, while improvements in cost-effectiveness will make viral inactivation testing more economically viable. As medical technology progresses and new medical devices emerge, the demand for viral inactivation testing may change, requiring testing institutions to adapt to new market needs. Regulatory agencies may impose higher requirements for the viral inactivation performance of medical devices, necessitating testing institutions to adapt to these changes to ensure compliance of their services. These comprehensive trends will collectively drive the development of the Medical Device Viral Inactivation Testing industry, providing more accurate, efficient, and reliable services to the medical device industry.

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

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

This Insight Report evaluates the key market trends, drivers, and affecting factors shaping the global outlook for Medical Device Viral Inactivation Testing 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 Medical Device Viral Inactivation Testing.

This report presents a comprehensive overview, market shares, and growth

opportunities of Medical Device Viral Inactivation Testing market by product type, application, key players and key regions and countries.

Segmentation by Type:

DNA-based

RNA-based

Segmentation by Biological Characteristics & Structure of Viruse:

Enveloped Viruses

Non-enveloped Viruses

Segmentation by Testing Method:

Biocompatibility Testing

Microbiological & Reusability Validation

Chemical Composition Analysis

Segmentation by Application:

Active Implant Medical Devices

Active Medical Equipment

In Vitro Diagnostic Equipment

Ophthalmic Medical Equipment

Vascular Medical Equipment

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 analyzing the company's coverage, product portfolio, its market penetration.

Charles River

Perfectus Biomed (NAMSA)

Eurofins Scientific

Nelson Labs

Texcell

Intertek Group

LabCorp

Syensqo

SGS

Pacific BioLabs

Creative Biogene

ProPharma Group

Cormica Lab (Wickham Micro)

CTI

WuXi AppTec

The report requires updating with new data and is sent in 48 hours after order is placed.

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