

## **Protein Labeling - Market Share Analysis, Industry Trends & Statistics, Growth Forecasts (2026 - 2031)**

Market Report | 2026-01-16 | 105 pages | Mordor Intelligence

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### **Report description:**

Protein Labeling Market Analysis

The protein labelling market was valued at USD 2.92 billion in 2025 and estimated to grow from USD 3.19 billion in 2026 to reach USD 4.98 billion by 2031, at a CAGR of 9.29% during the forecast period (2026-2031). Demand is propelled by breakthroughs that allow site-specific tags with minimal impact on protein conformation, an essential requirement for today's advanced proteomics, live-cell imaging and biotherapeutic workflows. Widening use of artificial-intelligence tools for structure prediction, a growing pipeline of antibody-drug conjugates and the migration of complex bioconjugation tasks to specialist CDMOs jointly broaden commercial opportunities. At the same time, capital-intensive analytical platforms and evolving regulations around radio-labelled conjugates remain practical hurdles. Competitive intensity is rising as incumbents look to differentiate via proprietary chemistries, integrated service models and strategic alliances with drug developers.

Global Protein Labeling Market Trends and Insights

Expanding R&D in proteomics and genomics

Advances in high-resolution mass spectrometry and AI-driven modelling have sharply raised the volume and depth of protein characterisation studies. The release of ESM-3 and AlphaFold 3 in 2024 improved structure-prediction accuracy, allowing researchers to pinpoint accessible residues for selective tagging and production. Funding programs across the United States and Europe now prioritise multi-omics pipelines that depend on robust labelling chemistries for quantitation and spatial mapping.

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These dynamics sustain premium pricing for next-generation probes and herald broader uptake of high-throughput workflows in academic cores and translational-medicine centres.

#### Growing application in drug discovery and development

Roughly 40% of investigational drugs now focus on historically undruggable proteins, elevating the role of homogeneous, site-specific labelling. Antibody-drug conjugate pipelines, currently exceeding 360 clinical studies, require precise linker-payload ratios for efficacy. Targeted protein degradation platforms similarly depend on live-cell tags that monitor real-time proteolysis. These use cases expand addressable demand across reagents, instrumentation and custom-conjugation services.

#### Limitations of protein labelling assays and methods

Large fluorescent tags above 25 kDa altered sub-cellular localisation in 42% of tested proteins, and nanobody conjugates showed 38% variability in binding efficiency across labs. Such inconsistencies raise validation requirements, extend project timelines and increase consumable usage, particularly in regulated environments.

Other drivers and restraints analyzed in the detailed report include:

Rising outsourcing to CDMOs/CROs  
Increasing use of fluorescence-based techniques  
Regulatory hurdles for protein-radioisotope conjugates

For complete list of drivers and restraints, kindly check the Table Of Contents.

#### Segment Analysis

Reagents and kits accounted for 70.62% of protein labelling market revenues in 2025, underscoring their essential role in day-to-day workflows across discovery, diagnostics, and manufacturing. Ready-to-use chemistries simplify protocol standardisation and support reproducibility, keeping demand resilient even as new techniques emerge. Continuous improvements in dye brightness and click-ready handles sustain incremental pricing power and entice repeat purchases.

The services segment, however, is outpacing headline growth at a 10.31% CAGR. This surge reflects the escalating technical complexity of antibody-drug conjugation, targeted degradation probes and live-animal imaging agents that exceed many in-house capabilities. CDMOs are adding dedicated 1,000-2,000 L suites, such as Lonza's 2024 expansion in Visp, to accommodate large-scale payload conjugation and fill-finish tasks. The protein labelling market size for outsourced projects is projected to expand steadily as pipeline sponsors seek rapid, GMP-compliant solutions.

Immunological techniques remained the largest slice of the protein labelling market with a 34.05% share in 2025, benefiting from decades of optimisation in ELISA, western blot and immunohistochemistry. High throughput and clinically validated protocols ensure enduring relevance in biomarker verification, lot-release testing and routine pathology.

Cell-based assays, however, record the most rapid advance at 10.96% CAGR as researchers prioritise physiologically relevant readouts in oncology, neurology and regenerative medicine. Fluorescent-probe portfolios from Thermo Fisher enable real-time monitoring of receptor activation, trafficking and complex assembly. The protein labelling market share captured by dynamic live-cell platforms is set to widen as automated imaging and high-content analytics permeate screening laboratories.

The Protein Labelling Market is Segmented by Product (Reagents and Kits, Services and Other Products), Application (Immunological Techniques, Cell-Based Assays and More), Labeling Method (In-Vitro Labelling and In-Vivo Labelling), End User

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(Pharmaceutical and Biotechnology Companies, Contract Research and Manufacturing Organisations and More) and Geography. The Market and Forecasts are Provided in Terms of Value (USD).

## Geography Analysis

North America generated 37.35% of 2025 revenues, supported by robust NIH funding, a dense biopharma cluster and early adoption of next-generation labelling technologies. MIT's CuRVE breakthrough, capable of labelling tens of millions of cells in a single day, exemplifies regional leadership in disruptive methodology. A high cadence of FDA approvals for ADCs and radio-immunoconjugates further anchors commercial momentum.

Europe ranks second, buoyed by Horizon Europe grants and strong biomanufacturing corridors in Germany, the United Kingdom and France. Sartorius' acquisition of Polyplus expanded the continent's vector-production capabilities, necessitating advanced tagging for viral envelope proteins. Regulatory harmonisation is progressing, yet divergent approaches to radio-isotope handling keep market entry costs elevated.

Asia-Pacific is the fastest-growing arena, advancing at an 11.40% CAGR. Chinese CDMOs such as WuXi Biologics report vibrant demand for conjugation formats branded under WuXiBody and associated linker chemistries. National industrial policies, including China's 14th Five-Year Plan and India's BIRAC initiatives, funnel capital into proteomic infrastructure, although uneven access to high-end MS platforms persists. South Korea's rapid scale-up of antibody-drug conjugate facilities, backed by Samsung Biologics, further cements regional ascent.

## List of Companies Covered in this Report:

Thermo Fisher Scientific Merck Danaher Corp. Revvity Roche Agilent Technologies Bio-Rad Laboratories KMD Bioscience Co., Ltd. Promega LI-COR Biotech, LLC New England Biolabs Takara Bio Vector Laboratories Jena Bioscience Lumiprobe Corp. Biotium Inc. Abcam Creative Diagnostics Cyanagen Srl Genscript

## Additional Benefits:

The market estimate (ME) sheet in Excel format  
3 months of analyst support

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#### 6.3.5 F. Hoffmann-La Roche AG

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#### 6.3.7 Bio-Rad Laboratories Inc.

#### 6.3.8 KMD Bioscience Co., Ltd.

#### 6.3.9 Promega Corporation

#### 6.3.10 LI-COR Biotech, LLC

#### 6.3.11 New England Biolabs Inc.

#### 6.3.12 Takara Bio Inc.

#### 6.3.13 Vector Laboratories Inc.

#### 6.3.14 Jena Bioscience GmbH

#### 6.3.15 Lumiprobe Corp.

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