eurofins | Viracor BioPharma

Accelerating drug development with regulatory-ready LC-MS/MS solutions

Eurofins Viracor BioPharma delivers **high-quality LC-MS/MS bioanalytical services** that help sponsors move confidently from preclinical research through clinical development. Combining advanced instrumentation, automated workflows, and deep scientific expertise, we generate reliable data that supports critical development and regulatory decisions.

Why Eurofins Viracor BioPharma?



Advanced mass spectrometry and UHPLC platforms



Automated sample preparation for enhanced efficiency and reproducibility



Direct access to experienced LC-MS scientists



Flexible, responsive project management



Regulatory-compliant support from method development through sample analysis

End-to-end bioanalytical support

Method Development/Transfer

Customized LC-MS/MS assay development designed to achieve the sensitivity, robustness, and performance required for your molecule and study objectives.

Method Validation

Validation services performed in accordance with current FDA, EMA, and ICH M10 guidance to ensure confidence in data quality and regulatory readiness.

Sample Analysis

Reliable quantitative analysis supporting:

- Pharmacokinetic (PK) studies
- Toxicokinetic (TK) studies
- Bioavailability/Bioequivalence (BA/BE) studies
- Preclinical studies
- Clinical development programs

Specialized Bioanalysis

Expert solutions for complex analytes and challenging matrices, including:

- Small molecules
- Peptides
- Oligonucleotides
- Metabolites
- Biomarkers
- Antibody-drug conjugates (ADCs)

Accelerating drug development with regulatory-ready LC-MS/MS solutions

Advanced technology. Reliable results.

Industry-leading instrumentation

Our LC-MS/MS laboratory leverages the latest technology to deliver:

- Exceptional sensitivity
- Broad dynamic range
- High-throughput performance
- Robust quantitative accuracy
- High-quality data for regulatory submissions

Automated workflows

Integrated automation platforms include Hamilton, Tecan, KingFisher™, and Biotage Extrahera™ systems, helping to:

- Improve reproducibility
- Minimize manual variability
- Increase throughput
- Support large, complex studies

Quality built into every step

Our quality-focused approach includes:

- GLP-compliant practices where applicable
- FDA, EMA, and ICH M10-aligned processes
- SOP-driven operations
- Comprehensive quality oversight
- Electronic data integrity controls
- Qualified and maintained instrumentation

Scientific partnership beyond testing

Our scientists work as an extension of your team, providing strategic guidance and technical expertise throughout your program. From assay development to sample analysis, we bring a collaborative approach that helps solve complex analytical challenges and keeps studies moving forward.

Experience across therapeutic areas and modalities

Therapeutic expertise

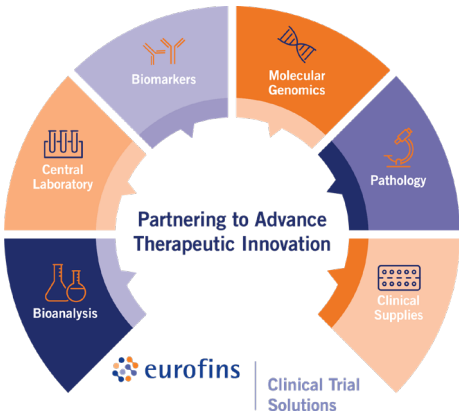
- Oncology
- Rare diseases
- Infectious diseases
- Cardiovascular diseases
- Neuroscience
- Metabolic disorders

Analytical expertise

- Small molecules
- Peptides
- Monoclonal antibodies (mAbs)
- Antibody-drug conjugates (ADCs)
- Oligonucleotides and oligomers
- Metabolites
- Biomarkers

Confident bioanalytical data. Smarter development decisions.

Whether you need a **custom assay**, **validated method**, or **large-scale sample analysis**, Eurofins Viracor BioPharma delivers the technology, expertise, and regulatory rigor required to support your development program from discovery through clinical trials.



Contact: clinicaltrials@vbp.eurofinsus.com

Visit: eurofinsvbp.com

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