



Method Development and Validation

Whether you need expert testing consultation, method development or validation protocol design, Eurofins BioPharma Product Testing offers you the widest range of laboratory techniques in the industry. We support customers at various stages of the development/validation process and have a proven track record of success on full development and validation studies, optimization of existing research methods and transfer of previously validated methods.

Why Choose Eurofins BioPharma Product Testing?

- We collaborate with you by thoroughly understanding your testing requirements, designing a project approach to achieve your end goals, developing methods in the most efficient manner to maximize your research budget and validating or transferring the method for its intended use.
- We offer phase appropriate method validation. All work is performed under full cGMP compliance.
- We have the most experienced staff, performing more than 400 validation, qualification, verification and transfer protocols for our customers per year.
- We place an emphasis on building relationships and rapport with our clients. Thanks to our extraordinary rate of employee retention, you can be confident that you will have the same management team for the duration of your project, as well as for any future work you may do with us.
- Our Method Development and Validation Team works hand-in-hand with the receiving lab to ensure a seamless transfer of methods.
- We assure timely delivery of services by establishing a project schedule and monitoring progress through weekly conference calls when useful.



Services

- **Method Development** – Prior to starting any project, we will discuss the project requirements with the client to ensure an appropriate scope is defined. We utilize our knowledge of method development on hundreds of products to assist with the development process. We evaluate attributes such as precision, linearity, accuracy and specificity during development to ensure the method is suitable for validation.
- **Method Optimization** – Methods may be optimized for a variety of reasons, including improving separation of a critical peak pair, or supporting a formulation change. In addition, methods not validated to current industry standards may require optimization to allow for successful validation.
- **Method Feasibility** – When methods are submitted for qualification or validation, we recommend performing feasibility prior to proceeding. Feasibility allows us to gain familiarity with the method and evaluate the method for attributes such as precision, linearity, accuracy and specificity with the hopes of avoiding protocol acceptance criteria failures.



- **Method Qualification** – Performed on methods supporting early phase product. Performed to ensure method is appropriate for its intended use. Qualification may or may not be performed under the direction of the protocol.
- **Method Validation** – Performed on methods supporting later phase product. Methods are validated consistent with the EP/USP general chapter and the ICH guidelines. Validation is performed under the direction of a protocol.
- **Method Verification/Suitability** – Performed on compendial methods to ensure that the methods perform acceptably on your API or product matrix. Verification is performed under the direction of a protocol.

Molecular and Cell Biology

- Cell-based Assays
- ELISA
- qPCR

Protocol Writing for

- Method Validation/Qualification/Verification
- Method Transfer
- Stability Studies

Method Development and Validation Services For: Chemistry/Biochemistry

- Assay/Potency, Purity/Impurity, Dissolution
- Cleaning Validation Analysis (Specific and Non-specific analyses)
- Assay, Impurity and Dissolution methods for Comparator products
- Residual/Impurities Testing
- Leachables
- Excipients

Microbiology

- Bacteriostatis/Fungistatis for Sterility
- Inhibition/Enhancement Screening for Endotoxin
- AET and Microbial Limit
- Mycoplasma Testing

Comprehensive GMP Testing Services

Method Development & Validation • Release Testing • Raw Materials Testing
Cell Banking Services • Virology Services • Facility & Process Validation
Chemistry • Biochemistry • Molecular & Cell Biology • Microbiology
Stability Testing & Storage • Primary & Secondary Package Testing

Flexible Service Models

Fee For Service (FFS)
Full-Time-Equivalent (FTE)
Professional Scientific
Services® (PSS)

Global Facilities

Australia	Denmark	India	Japan	Spain	UK
Belgium	France	Ireland	Netherlands	Sweden	US
Canada	Germany	Italy	New Zealand	Switzerland	