eurofins | BioPharma Product Testing



Product Release Testing for the European Union

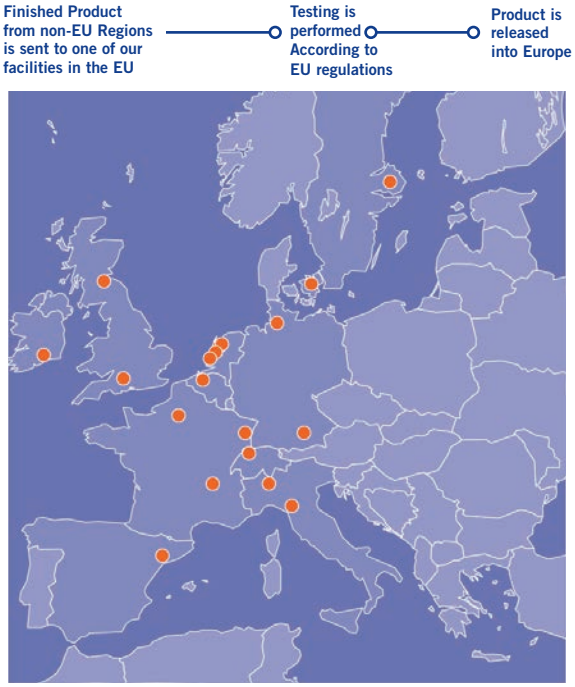
As a global provider of GMP analytical services, Eurofins BioPharma Product Testing is strategically located to provide importation testing for bio/pharmaceutical products on entry into the European Union.

Under EU legislation, all products manufactured outside the EU or states without mutual recognition, must have release testing perfomed upon import into the European market. This testing, combined with appropriate assessments of the related quality systems by a European Qualified Person (QP), enables product to be released to market.

If you need support for effective passage of your product into the EU, Eurofins BioPharma Product Testing can handle and test your product in strict compliance with cGMP. We have been supporting our clients global supply chains and entry into the EU for over 20 years from our facilities in the EU.

Why Choose Eurofins BioPharma Product Testing?

- Our state-of-the-art global facilities offer extensive laboratory capacity and the highest level of instrument technology, enabling our facilities to become an extension of your own operation.
- Our flexible operating systems allow timely testing and reporting of your results to expedite market entry. One of our dedicated client service representatives will monitor and report progress on your release.
- Our 25-year history of cGMP compliance ensures you will receive the highest quality data. We meet your global reporting needs with data that is generated in full compliance with EMA.



- We work extensively with many of the worlds leading global biopharma companies in this area and have also established relationships with logistics organizations, importers and QPs. We also work with many virtual organizations to enable entry into the EU. Many of these organizations use Eurofins BioPharma Product Testing as their primary testing support for entry into the EU.

EU Product Release Services

- Comprehensive Biologic & Pharmaceutical Product Support
- QP Release
- Product Release
- Method Transfer
- Method Validation
- Product Receipt and Storage

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

9122_EU 0817

www.eurofins.com/biopharma