

Medical Device Testing







Toxicology Services

Eurofins Medical Device Testing has more than 30 years of experience in safety testing for biopharmaceuticals, chemicals, and medical devices. This experience will provide you with the technical advice to ensure the success of your product development.

Our toxicologists perform genetic toxicology assessments, alternative toxicology assessments and toxicological risk assessments based on data from extractable and leachable testing to help you understand the safety profile of your medical device. Based on ISO 10993-17, Eurofins Medical Device Testing will identify and evaluate toxicity risks to humans exposed to the final medical device. Evaluations may also be performed on individual chemical compounds, additives, colorants, processing aids, and other potential leachables. Once identified, Eurofins Medical Device Testing can help to determine if additional analytical testing will be needed to help support your product.

Choose Eurofins Medical Device Testing to help you:

- Profit from our network of experts who will help you with advice and action
- Meet international guidelines to achieve the best solution for your product
- Successfully meet testing challenges and prevent costly production problems and delays
- Provide competent support for complex tasks



Biocompatibility Services

- Creation of Certificates and/or Expert Statements
- · Consulting and Customer Support
- Test Strategy Development
- Biological Evaluation Plan and Report Creation
- Gap Analysis
- · Risk Plan Review
- · Biological Risk Plan Review
- Toxicological Risk Assessment of Extractables & Leachables
- Toxicological Support