EUROPEAN BIOMEDICAL INSTITUTE

• WHERE SCIENCE STARTS THE FUTURE •

- **ISO 17025** accredited
- Good Laboratory Practice **(GLP)** certified
- **US FDA 21 CFR 58** compliant
- Laboratories located in Europe: **United Kingdom** and **Poland**

Our studies are recognized by **Competent Authorities** (e.g. FDA, MHLW, MFDS, ANVISA) and **Notified Bodies**

As the European Biomedical Institute, we provide **comprehensive**, in-house **biocompatibility** and **chemistry testing** of medical devices according to the ISO 10993 standards.

>96% studies on time

Customer satisfaction score

4.6 ★ ★ ★ ★ ★

ISO 17025

CERTIFIED GLP GOOD LABORATORY PRACTICE

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List of routinely conducted tests:

Biological Evaluation Plan (**BEP**)	ISO 10993-1
Physical and/or chemical information (**extractables & leachables**)	ISO 10993-18
Toxicological risk assessment	ISO 10993-17
Cytotoxicity	ISO 10993-5
Sensitization	ISO 10993-10
Irritation or Intracutaneous reactivity	ISO 10993-23
Material mediated pyrogenicity	ISO 10993-11
Acute systemic toxicity	ISO 10993-11
Subacute toxicity	ISO 10993-11
Subchronic toxicity	ISO 10993-11
Chronic toxicity	ISO 10993-11
Implantation effects	ISO 10993-6
Hemocompatibility	ISO 10993-4
Genotoxicity	ISO 10993-3
Carcinogenicity	ISO 10993-11
Reproductive/developmental toxicity	ISO 10993-3
Degradation	ISO 10993-13, -14, -15, -16
Nanomaterials	ISO 10993-22
Biological Evaluation Report (**BER**)	ISO 10993-1

BIOLOGICAL SAFETY TESTING - BIOCOMPATIBILITY

CHEMISTRY TESTING

MICROBIOLOGY & STERILITY

PACKAGE & REPROCESSING VALIDATIONS

Please contact us if you would like to receive a **competitive offer** and **professional advice**:

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