SMALL ENOUGH TO CARE ... 

... AND BIG ENOUGH TO DELIVER

**BIOCOMPATIBILITY TESTING**
ACC. ISO 10993:
- Cytotoxicity
- Irritation
- Skin Sensitization
- Genotoxicity
- Chemical Characterization
- Nanomaterials

The Medical Device Regulation (EU) 2017/745) requires a certification for Medical Devices to prove that they are safe to use and no adverse effects can be expected after contact with the patient. The thorough evaluation of biological safety (Biocompatibility) as described in the ISO 10993 series is an essential step in this process.

LAUS – The GLP Lab – offers testing according to the principles of Good Laboratory Practice for more than 20 years. But we do more than test your products.

With 30 years of experience in regulatory compliance testing our experts offer their advice regarding the experimental approach and optimized test strategy, customized for your test item. Since we are a small privately owned company, we are able to treat every customer individually.

LAUS can assist you in choosing the appropriate studies and in developing the applicable test strategy customized for your Medical Device.

Testing in accordance with the principles of Good Laboratory Practice (GLP) ensures the acceptance of the studies worldwide.

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