



# BIOCOMPATIBILITY TESTING OF MEDICAL DEVICES

The ISO 10993-1:2018 standard requires a structured biological evaluation plan within a risk management process. TÜV SÜD America can assist with the biological evaluation plan generation, toxicological risk assessment, and the conclusive biological evaluation report to help your device meet the requirements of the standard.

## Quality, Safety and Reliability

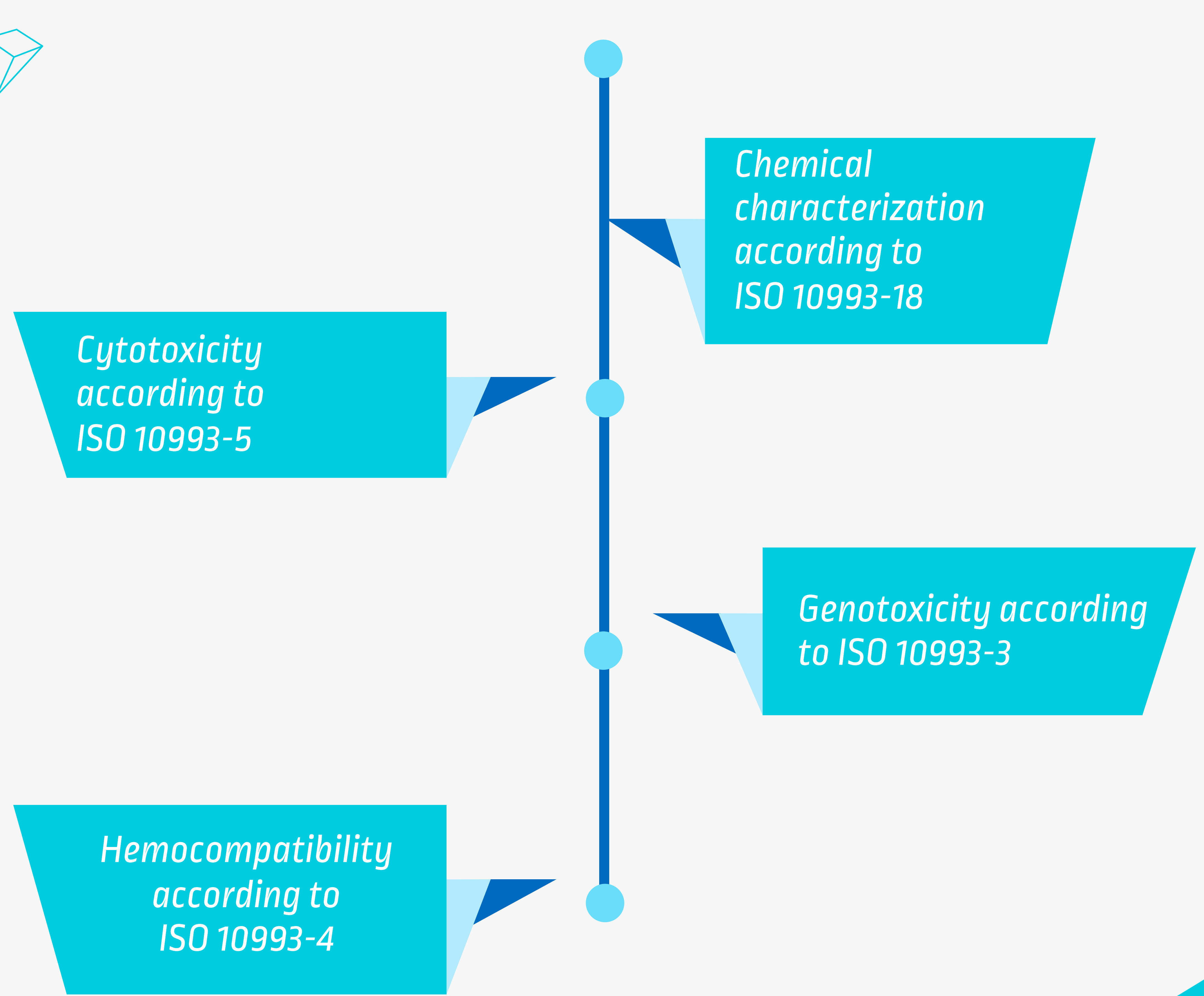
As the most trusted, experienced, and reliable service provider, TÜV SÜD works with 75 percent of the world's largest medical companies. You can be certain of our in-house technical expertise in every field.



A one-stop solution provider for all of your testing and certification requirements.

## Biocompatibility Testing

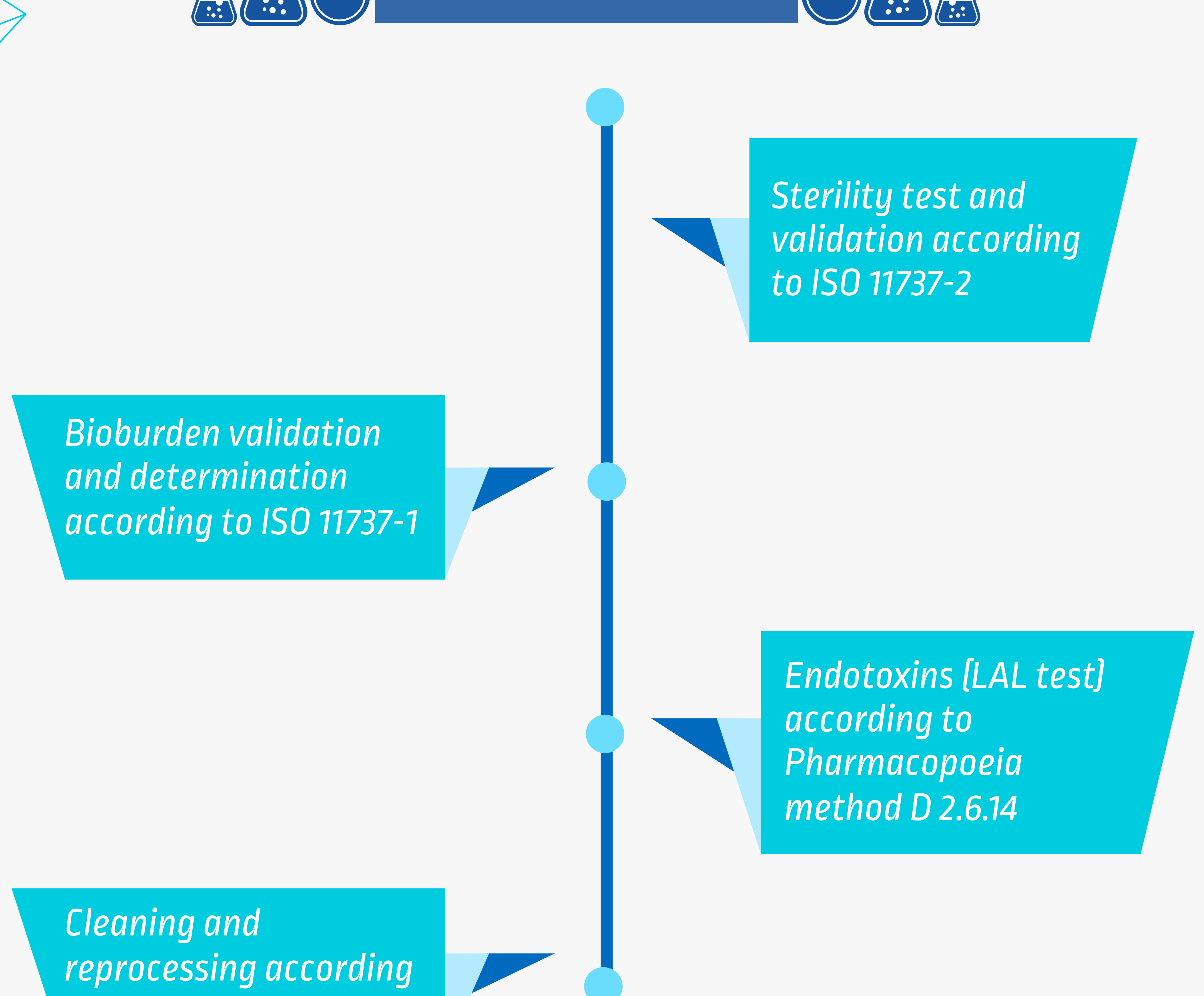
TÜV SÜD provides coverage of assessment services to help manufacturers meet the biocompatibility testing requirements of ISO 10993, FDA, and other regulatory agencies.



## Additional Services

### Microbiology and Sterility Testing

Our state-of-the-art laboratories are fully ISO 17025 accredited. Our facilities are also able to perform testing under Good Laboratory Practices (GLP).



## Choose our unmatched expertise

### One-Stop Shop for Medical Device Testing

By choosing TÜV SÜD, you can be assured that you benefit from the premier standard of clinical assessment reports. With the highest number of medical device experts and the most comprehensive service offering of any Notified Body (NB), our team is ready to assess your innovative design.

### Let's work together!

TÜV SÜD offers biocompatibility testing services for medical devices conducted in our state-of-the-art laboratories and managed by highly skilled experts to ensure the complete availability of high-quality data and reports to meet your testing needs.



TÜV SÜD America

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[www.tuvsud.com/en-us/industries/healthcare-and-medical-devices](http://www.tuvsud.com/en-us/industries/healthcare-and-medical-devices)