Achieve EU Market Access for Medical Devices with TÜV SÜD Notified Bodies

01

COMMON CHALLENGES IN MEDICAL DEVICE REGULATION CERTIFICATION

Changing **transition timelines**

Expanded scope of devices that require Notified Body review and approval

Delays in the **review and approval process**

Lack of **notified bodies capacity**

Long **waiting time**

02

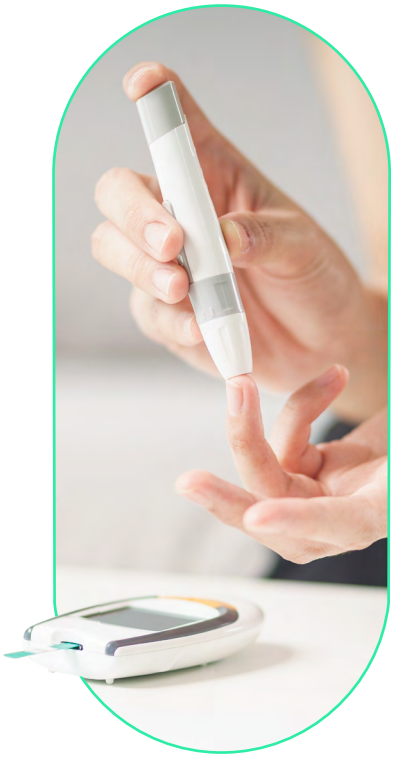
TÜV SÜD NOTIFIED BODIES FOR MEDICAL DEVICE REGULATION

2 **EU notified bodies**
and more than...

750
medical device professionals
in more than...

30
locations worldwide

Among the world's first organisations to receive designation as a Notified Body for the MDR.



03

ACHIEVE MEDICAL DEVICE REGULATION COMPLIANCE WITH TÜV SÜD

1 Our extensive expertise in all areas of medical devices, as well as our niche focus in cardiovascular, software, orthopaedic and neurovascular devices, ensures transparent assessments and support

2 Experts undergo continuous training and development to stay up-to-date with the latest trends and changes

3 Broad designation for most medical devices

4 Team of experienced professionals with extensive industry and notified body knowledge

5 Transparent communication and customer-centric approach

6 Seamless and timely certification experience for each client

REGISTER NOW