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Integrated Scientific Services

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BVMA member since 2016
Audit passed in 2016



HISTORY AND MISSION

ISS AG, Integrated Scientific Services provides services to medtech companies for the development of medical devices, their introduction into the market and their maintenance. Typical services include embedded software development, regulatory affairs and clinical evaluation report writing, clinical research, quality management, qualification and validation. The combination of these services with scientific work methods and project management puts ISS AG into a unique position as a service provider for the medtech industry.



CLINICAL SERVICES
EVALUATIONS & CRO

REGULATORY AFFAIRS
WORLDWIDE

QUALITY MANAGEMENT
ISO 13485

SOFTWARE DEVELOPMENT
IEC 62304

REGULATORY AFFAIRS SOFTWARE
REGULA™

HIGHLIGHTS

ISS already has submitted successfully clinical evaluation reports according to MEDDEV 2.7/1 Rev. 4.

With successful product registrations in 99 countries of the world, we are the right partner when it comes to the registration of your product.

Using REGULA™, our web-based RA software, will significantly shorten your time-to-market and offers you instant overview over all your product registrations worldwide incl. deadlines.

Our highly qualified team of software engineers develops embedded and standalone medical software according to IEC 62304.

SERVICES

Clinical Services

- ▷ Clinical Project Management: Study planning (strategy & design), site selection, medical writing and submission of essential documents to Ethics and Comp. Authority, conducting of clinical study (incl. site initiation, routine monitoring, close out, data management and statistics), and completion of the study (data base lock, final statistical analysis, study report, medical writing)
- ▷ Clinical Evaluations according to MEDDEV 2.7/1 Rev. 4

Regulatory Affairs

- ▷ Product registrations worldwide
- ▷ Implementation of REGULA™, our submission management software
- ▷ RA-Intelligence Services, “on demand” and “push”
- ▷ Definition of regulatory strategy within business development processes and projects

Quality Management & Engineering Support

- ▷ Building and maintaining Quality Management Systems for MedTech (ISO 13485)
- ▷ Creation of and support all around the Technical Documentation
- ▷ Interim Quality Manager Services
- ▷ Audit Support, V&V
- ▷ CAPA, Complaint handling

Software Development (IEC 62304)

- ▷ Software Development for medical devices of all risk classes (embedded and stand-alone)
- ▷ Software project management
- ▷ Strong track record in the fields of medical image analysis, numerical simulations, hardware and system related development (firmware), security applications for medical devices (RFID supported)

