BSI Akademie

New Medical Devices Training courses in Switzerland in Zurich

- ISO 13485:2016 Lead Auditor
  This course teaches the principles and practices of effective quality management system
  I 22.06.2018 I 19.10.2018 I in German and English I

- MDD to MDR Transition
  Learn all about the new requirements and changes
  I 05.03.2018 I 11.06.2018 I in German and English I

- ISO 13485:2016 Transition and Auditor Refresher
  Get to know the latest requirements of the standard since ISO 13485: 2003 and EN 13485: 2012
  I 19.03.2018 I 27.11.2018 I in German I

- Suppliers Auditor 13485:2016 Training Course
  By training as a supplier auditor you will learn principles and dealing with
  auditing based on the international standard ISO 19011
  I 23.03.2018 I 24.08.2018 I in English I

- MDSAP Fundamentals and Readiness Training
  This course will prepare you to host a MDSAP audit and allow you to determine if your own internal QMS
  processes are consistent with the requirements of the MDSAP audit mode for the jurisdictions
  where your products are marketed
  I 26.03.2018 I 10.07.2018 I in English I

- IVDD to IVDR Transition
  In this course you will learn all about the changes and requirements of IVDD to IVDR
  I 23.04.2018 I in English I

- ISO 13485 Introduction
  This one day course has been designed to provide an insight in to the use of ISO 13485
  I 07.05.2018 I 16.07.2018 I 26.11.2018 I in German and English I

bsi.

...making excellence a habit.™

Medical Devices – Training courses

We want to offer you the best possible learning experience. With BSI you can learn in a pleasant learning environment internalize practical knowledge and skills.

Is this the right course for you?

- **ISO 13485 Implementing**
  This seminar course has been designed to provide participants with the knowledge and process steps to enable them to effectively implement a quality management system
  I 09.05.2018 I 18.07.2018 I 28.11.2018 I in German and English I

- **ISO 13485 Internal Auditor**
  This course is intended for medical device quality professionals aiming to build on their current knowledge of ISO 13485 and evaluate the effectiveness of the quality management system in their organization
  I 11.05.2018 I 20.07.2018 I 30.11.2018 I in German and English I

- **Technical Documentation**
  This course is designed to support manufactures by confirming current regulatory requirements of technical documentation
  I 14.05.2018 I in English I

- **Post Market Surveillance**
  This course is designed to help you identify the requirements of the European medical device directives, standards and guidance documents
  I 15.05.2018 I in English I

- **Clinical Evaluation**
  Learn all about the evidence of clinical safety and performance of your product accordance with the requirements of the European MDD and the AMIDD
  I 16.05.2018 I in English I

- **ISO 14971 Risk Managament**
  Gain an understanding of the impact that ISO 14971:2012 has on the decision making process when manufacturing medical devices
  I 17.05.2018 I in English I

- **Medizinprodukteberater**
  In this seminar, we convey the essential requirements, which are made to medical device consultants and the requirements for establishment of an observation and reporting system
  I 22.08.2018 I in German I

Please give us a call or contact us per e-mail !

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