



## 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

| 18.06.2018 | 15.10.2018 | in German and English |

- **MDD to MDR Transition - Zurich and Basel**

Learn all about the new requirements and changes

| 05.03.2018 | 11.06.2018 | in German and English |

| Also in Basel | 04.06.2018 | 04.06 - 05.06.2018 |

- **ISO 13485:2016 Transition and Auditor Refresher**

Get to know the latest requirements of the standard since ISO 13485: 2003 and EN 13485: 2012

| 19.03.2018 | 27.11.2018 | in German |

- **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

| 23.03.2018 | 24.08.2018 | in English |

- **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

| 26.03.2018 | 09.07.2018 | in English |

- **IVDD to IVDR Transition**

In this course you will learn all about the changes and requirements of IVDD to IVDR

| 23.04.2018 | in English |

- **ISO 13485 Introduction**

This one day course has been designed to provide an insight in to the use of ISO 13485

| 07.05.2018 | 16.07.2018 | 26.11.2018 | in German and English |

# 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

| 08.05.2018 | 18.07.2018 | 28.11.2018 | in German and English |



- **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

| 10.05.2018 | 20.07.2018 | 30.11.2018 | in German and English |

- **Technical Documentation**

This course is designed to support manufactures by confirming current regulatory requirements of technical documentation

| 14.05.2018 | in English |

- **Post Market Surveillance**

This course is designed to help you identify the requirements of the European medical device directives, standards and guidance documents

| 15.05.2018 | in English |

- **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

| 16.05.2018 | in English |

- **ISO 14971 Risk Management**

Gain an understanding of the impact that ISO 14971:2012 has on the decision making process when manufacturing medical devices

| 17.05.2018 | in English |

- **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

| 22.08.2018 | in German |

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



Oliver Schaer  
Phone: +49 (0) 69 2222 89 265  
Email: [Oliver.Schaer@bsigroup.com](mailto:Oliver.Schaer@bsigroup.com)

