

Our medical devices courses and qualifications allow you to have a detailed look at some of the most important sections of bringing a successful product to market within this highly regulated sector.

Learn more about our courses, upcoming dates and booking options using the course links below:

ISO 13485	Feb	Mar	Apr	May	Jun	Location	Duration	Book
ISO 13485:2016 Introduction	13	20	3 / 4	2		Online*	1 day	LINK
ISO 13485:2016 Requirements	On-demand					eLearning	4 hours	LINK
ISO 13485:2016 Clause by Clause	20-21	20-21	24-25	24-25	26-27	Online*	2 days	LINK
ISO 13485:2016 Implementation		21-22			26-27	Online	2 days	LINK
ISO 13485:2016 Internal Auditor	14-15 27-28	14-15 30-31	26-27	23-24 30-31	14-15 29-30	Online*	2 days	LINK
ISO 13485:2016 Lead Auditor	27-3		11-14		19-23	Online*	5 days	LINK

CE Marking	Feb	Mar	Apr	May	Jun	Location	Duration	Book
Requirements of the In Vitro Diagnostic Regulation (IVDR)					5	Online	1 day	LINK
Implementation of the In Vitro Diagnostic Device Regulation (IVDR)					6-8	Online	3 days	LINK
IVD Directive (IVDD) to IVD Regulation (IVDR) Transition	Contact us					Online	1 day	LINK
Requirements of the Medical Device Regulation (MDR)		27		9		Online*	1 day	LINK
Requirements of the Medical Device Regulation (MDR)	On-demand					eLearning	4 hours	LINK
Implementation of the Medical Device Regulation (MDR)		28-30		9-11		Online*	3 days	LINK
Introduction to Medical Devices Software		10		17		Online	1 day	LINK

Specialist	Feb	Mar	Apr	May	Jun	Location	Duration	Book
Medical Device Single Audit Program (MDSAP): Fundamentals and Readiness			24-25			Online	2 days	LINK
Clinical Evaluation for Medical Devices		15		17		Online	1 day	LINK
ISO 14971:2019 Risk Management for Medical Devices: Requirements	16	6	3	30	12	Online	1 day	LINK
ISO 14971:2019 Risk Management for Medical Devices: Requirements	On-demand					eLearning	4 hours	LINK
Manufacturing Process Validation for Medical Devices: Introduction to Concepts and Methods	20		24			Online	1 day	LINK
Performance evaluation and clinical evidence for In Vitro Diagnostics (IVDs)		23				Online	1 day	LINK
Post Market Surveillance and Vigilance under MDR and IVDR	8		13		14	Online	1 day	LINK
Technical documentation for In Vitro Diagnostic Devices (IVDs)		6				Online	1 day	LINK
Technical documentation for the Medical Device Regulation (MDR)	23	27			8	Online	1 day	LINK

*Some of these course dates are available at classroom venues across the UK. Follow the link to find out more.

Contact our training team today to find out more and book your place.

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