ISO 13485:2016 Transition Workshop

Attend our half-day workshop and we will provide you the knowledge of the new revisions. With BSI, you can rise above the confusion and know how to prepare your team for the best way to position your organization.

Book your seat now to enjoy 70% off on 2nd participant and be a step ahead of everyone else.

Speaker Profile

Dr Qian Liang
Head of Med Devices, BSI Asia Pacific

Dr Qian Liang is the Head of Medical Device for BSI Asia Pacific Region based in United Kingdom, where she is responsible for the BSI Medical Device business in the Asia Pacific Region including certification, assessment and training.

Dr Qian Liang holds a PhD Degree in Metallurgy and Materials and a BEng Degree in Computer Science and Technology. The device experience of Dr Qian Liang includes the design and development of orthopaedic implants and associated instruments. She also a qualified Lead Auditor for ISO 13485 and technical file reviewer for European Medical Device Directive (MDD).

Agenda & Workshop Details

Agenda

0830 - 0900	Registration
0900 - 0910	Welcome and Introduction
0910 - 1015	Introduction to ISO 13485:2016 Medical Devices Revision
1015 - 1045	Coffee Break
1045 - 1145	Continue session on ISO 13485:2016 Medical Devices Revision
1145 - 1200	Action Manager & VerifEye
1200 - 1230	Question & Answer Session

Workshop Details

Date: 11th April 2016

Time: 0830 - 1230

Venue: BSI Services Malaysia Sdn Bhd
(Click here for map location)

Fees: RM100 per participant
RM30 for 2nd participant
and more

Registration Dateline : 31st March 2016

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