Quality, Regulatory Affairs, Medical Device & IT

mediteq FORUM

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Mediteq Forum Digital- Post Market Surveillance

> ″ By failing to prepare, you are preparing to fail″
>
> – Benjamin Franklin

Mediteq Forum is welcoming Paolo Valsecchio and Diego Falletti from BSI Italy who will guide us through the new requirements on post market surveillance under the Medical Device Regulation (MDR). Date of application is later this spring and there is a range of new requirements that will push manufacturers to take a proactive role in the monitoring of their devices. The presentation will cover the PMS requirements in Annex III and will focus on articles 83-86, 88, 32 and the reports that are mentioned in these articles.
The presentation will be held in English.

Date: Thursday 22nd April 2021, 13.00-15.00
Digital: Zoom link will be sent closer to the date
Sign up: here

The presentation will cover the PMS requirements as described above and there will also be a chance for the participants to ask questions.

Please send in your questions ahead of time if you have specific issues you'd like the experts to cover.

Welcome!

Petra Rosén and Jennie Bengtsson
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