SPARK EUROPE WEBINAR SERIES 2022

Wednesdays at 4 pm CET

SPARK Europe Education

Medical Devices Regulation

Anne-Mari Håkelien & Ingvild Gudim

4 May 2022 | 4 – 5 pm (CET) | Online Webinar

Medical device compliance is imperative to the success of every medical device company. Failing to achieve compliance will compromise the future of the product.

It is in in everyone's best interest to produce quality medical devices, therefore this industry is heavily regulated.

But, how do you actually achieve compliance for your medical device?

In this **SPARK EUROPE WEBINAR**, **Anne-Mari Håkelien and Ingvild Gudim** will provide some answers and perspectives to these questions, covering both MDR and IVDR.

Anne-Mari is a Senior Adviser at The Norwegian Medicines Agency with an extensive experience from both regulatory authorities and basic cancer research. She has also worked in a cell therapy start-up. Anne-Mari holds a PhD in molecular biology from the University of Oslo.

Ingvild is a Senior Adviser for medical devices and IVD medical devices at The Norwegian Medicines Agency. Prior to that she was working several years as project leader and technical lead for diagnostic immunoassay development in Gentian Diagnostics ASA. Ingvild holds a PhD in biochemistry from the University of Oslo.

Online via Zoom | Please register here!

Registration to the webinar is required in advance. Please register no later than May 3rd 2022.

The webinar is open to all **SPARKees**, **SPARK mentors**, and **students** and **staff** of **SPARK associated organizations**.

In unforeseen cases, the organizers may change and update topics and speakers.

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