SPARK EUROPE WEBINAR SERIES 2024/25

Wednesdays at 4 pm CET

SPARK Europe Education

Navigating AI Acts in Europe and the USA: Key considerations for MDR and FDA Approvals

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8 January 2025 | 4 – 5 pm (CET) | Online Webinar

In this webinar, we will explore the **AI Acts** in both Europe and the USA, focusing on their broad implications for software and algorithm development in healthcare. The session will provide a high-level overview of the **regulatory frameworks** and highlight important aspects to consider when creating new healthcare solutions. We will compare the **AI regulations** across these two regions, emphasizing potential differences and areas where these Acts intersect or diverge. This session is designed for those involved in the development of AI tools for healthcare.

Key takeaways:

- **Overview of the AI Act** in Europe and its U.S. counterparts
- Key **differences** and **similarities** in regulatory requirements
- Important perspectives to consider when developing healthcare-related **AI algorithms** and **software solutions**

MDS Finland specializes in offering regulatory, quality management, and commercialization consulting services for medical device companies. Jani has extensive expertise in navigating the regulatory landscape for medical devices and assisting companies in bringing their products to market.

Online via Zoom | Please register here!

Registration to the webinar is required in advance in order to receive Zoom meeting link
The event is hosted by SPARK Finland

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