When you need
to be sure



Open lecture 13th Nov 13.45 – 16 Arvo, F115

By IVDR Notified Body SGS Fimko Oy

From innovation to the European Medtech market?

This lecture introduces the MDR and IVDR regulations governing medical devices and in vitro diagnostic medical devices, covering their key requirements, CE marking, and the conditions for market access.

Designed for researchers and startups, the lecture offers a practical introduction to regulatory requirements and helps participants incorporate regulatory considerations into the product development process from the earliest stages.

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