LEAN MDR WEBINAR

11 DECEMBER
9.00-10.00 AM

The EU regulation for medical devices (MDR) imposes significantly higher demands on manufactures of low-risk devices (class I) than the previous directives. To be compliant requirements must be implemented in the quality management system and the technical documentation.

This webinar will guide you on how to be compliant.

- Go through what applies to class I manufacturers within the EU.
- Examples of what the technical documentation should contain.
- Examples of how to build effective processes and documentation that creates added value.
- Examples of what a management system should contain.

Participate in a short overview of how to bring your com

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