Add value.
Inspire trust.

Risk Management ISO 14971:2019

Your training at TÜV SÜD

The ISO standard of 14971:2019 has been published a few months ago. What impact do the new revisions of the standards ISO 14971:2019 and ISO/TR 24971:2020 have on risk management? How does the standard relate to MDR? If you look at the details of the standard, there are numerous details and possibilities to improve important contents in risk management sustainably. In the E-Learning you will get to know the innovation of ISO 14971:2019 as well as an overview of ISO/TR 24971:2020. You will understand how to implement the changes of the new ISO 14971:2019 in your company. The connections between MDR and risk management will be explained to you. At the end of the E-learning you can test your knowledge in a short knowledge test.

- You get to know the essential changes of ISO 14971:2019.
- You can estimate what the ISO 14971:2019 means for your own company.
- The e-learning is available worldwide, 24 hours a day, 7 days a week. You decide for yourself what you would like to learn, when, how often, and for how long.

Course contents
Topics to be covered in this course include:

- New terms and definitions
- Changed structure and contents
- Overview of the changes in ISO 14971:2019 and ISO/TR 24971:2020
- Consideration of the requirements of (EU) 2017/745 MDR
- Overview ISO/TR 24971:2020

Who should attend
This course is specially designed for Product and process developers, medical device manufacturers and their suppliers, employees in quality management and regulatory affairs, QM representatives, safety representatives for medical devices, as well as consultants and service providers

Methodology
The cumulative duration of the programme is 60 min., after which you will be required to pass a quiz to receive your internationally recognised e-certificate with a unique ID.

Sale Modes: Single License, Group License, SCORM Sale

Duration: 60 minutes

Number of Chapters: 5

Learning Mode: Self-Paced (LMS),

Language: English, German