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ISO 14971 Risk Management Requirements for Medical Devices

ISO 14971 Risk Management Requirements for Medical Devices

Semina	r	1 Date available	Certificate of Attendance
P Classro	om training	16 Lessons	

Seminar Number: IDN-I2-PUB

Status: 24.11.2024. All current information can be found at https://academy-id.tuv.com/s/IDN-I2-PUB

Join our comprehensive two-day training program to enhance your proficiency in risk management for medical devices. Gain in-depth knowledge of **ISO 14971**, navigate regulatory requirements, and implement effective control measures to ensure compliance and maximize patient safety.

Ideal for professionals in the medical device industry, this program equips you with the skills to assess, control, and mitigate risks throughout the device lifecycle. Become a skilled risk management professional and drive success in the dynamic field of medical devices!

Benefits

- Understanding of ISO 14971 and equip them with the necessary knowledge and skills to implement risk management practices for medical devices.
- Gain the ability to assess, control, and mitigate risks throughout the device lifecycle, ensuring compliance and enhancing patient safety.

Target group

- Professionals working in the medical device industry
- Regulatory affairs managers and specialists
- Quality assurance and quality control professionals
- Research and development engineers
- Product managers and project leaders



- Compliance officers
- Medical device consultants
- Risk management professionals

Requirements

No Prerequisites required. Anyone can complete this course.

Training outline

- Day 1: Introduction to ISO 14971 and Risk Management Planning
- Day 2: Risk Analysis, Control Measures, Documentation, and Continuous Improvement

Other information

For payment and other inquiries, please contact our Sales Executive:

- 1. Ms. Rezky Citra (Rezky.Citra@tuv.com | +62 896-1257-3270)
- 2. Ms. Novia Yulianti (Novia. Yulianti@tuv.com | +62 814-1350-3255)
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- 4. Ms. Jesseca Cindy (Jesseca.Cindy@tuv.com | +62 812-8183-8854)

ADDITIONAL INFORMATION

- Fees are inclusive of training materials and certificate.
- Training fees exclude 11% VAT.

CANCELLATION POLICY

- TÜV Rheinland Indonesia, reserves the right to postpone and cancel public courses.
- Unless cancelled by TÜV Rheinland Indonesia, training fees are non-refundable.
- Participants with late cancellation (five days prior the training schedule) will not be refunded. Full amount of the training fee will be charged and invoiced.
- Transferability: If you are unable to attend, a substitute delegate may attend in your behalf. Please provide the name and title of the substitute delegate



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Event overview and booking

Book your desired date now directly online at https://academy-id.tuv.com/s/IDN-I2-PUB and benefit from these advantages:

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Alternatively, you can use the order form to order via fax or e-mail.

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I HEREBY BINDING REGISTRATION FOR THE FOLLOWING SEMINAR:

ISO 14971 Risk Management Requirements for Medical Devices

Seminar Number: IDN-I2-PUB					
Please choose an appointment you would like to book:					
28/11/2024 - 29/11/2024, Daerah Khusus Ibukota Jakarta Event number: IDN-I2-PUB- ISO 14971 Risk Management Requirements for Medical Devices Rp 3.500.000,00 (Net price, plus VAT) Rp 3.885.000,00 (Gross price, including VAT)					
All further information about the dates can be found at https://academy-id.tuv.com/s/IDN-I2-PUB.					
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