

ISO 14971: Risk Management of Medical Devices

Audit & Quality Assurance
Sharm El-Sheikh (Egypt)
16 - 20 Mar 2025

UK Training

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ISO 14971: Risk Management of Medical Devices

Ref: 3178_133337 **Date:** 16 - 20 Mar 2025 **Location:** Sharm El-Sheikh (Egypt) **Fees:** 3700 Euro

Course Introduction

On completion of this course, the participant will learn why the risk management process is important, the general requirements of a risk management system and how to perform a risk analysis and control risk in an organization. ISO 14971 specifies a process for a manufacturer to identify the hazards associated with medical devices, estimate and evaluate the associated risks, to control these risks, and to monitor the effectiveness of the controls. The requirements of ISO 14971 are applicable to all stages of the lifecycle of a medical device. This course helps medical device professionals understand how ISO14971 can improve their business and risk management efforts. The student will gain an understanding of the impact that ISO14971 has on the decision-making process when manufacturing medical devices. After finishing the course, the student will gain the ability to work with risk management for medical devices according to the ISO 14971 standard.

Course Objectives of ISO 14971: Risk Management of Medical Devices

- Identifying the links between ISO 13485 QMS and ISO 14971 RM
- Explaining how risk management relates to the product lifecycle
- Defining risk management terminology
- Outlining the stages of the risk management process
- Defining the key deliverables of the risk management process

ISO 14971: Risk Management of Medical Devices Course Outlines

Day 1

Introduction to risk management in medical devices

- Links of risk management to regulatory requirements
- The relationship of ISO14971 to other standards
- ISO13485 and Risk Management
- Understanding ISO13485

Day 2

The development of ISO14971

A graphic of a chessboard with several chess pieces (a king, a queen, a rook, and a pawn) on it. The text 'UK Training PARTNER' is overlaid on the board.

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- Why do we risk manage medical devices using ISO14971?
- The Lifecycle approach
- ISO14971 - Overview of the standard
- Terms and definitions
- General requirements of the standard

Day 3

The risk management process

- The risk management plan
- Risk analysis
- Risk evaluation
- Risk control

Day 4

Residual risk acceptability

- Report/documentation
- The Risk Management File Postproduction
- IEC 606011:2005: Medical Electrical Equipment, General requirements for basic safety and essential performance

Day 5

Tools for Risk management

- Failure modes and effect analysis
- Fault tree analysis
- Hazard analysis and critical control point
- Summary and conclusions

A graphic of a chessboard with several chess pieces (a king, a queen, a rook, and a pawn) on it, set against a background of concentric circles. The text 'UK Training PARTNER' is overlaid on the right side of the board.

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