MEDITRIAL ACADEMY

MDR Fundamentals

REGULATIONS MADE EASY

GET READY TO LEARN ALL YOU NEED TO KNOW ON THE NEW EUROPEAN REGULATION FOR MEDICAL DEVICES.

MDR FUNDAMENTALS

Welcome to the Meditrial MDR Fundamentals. In three easy VIDEO-COURSES, you will learn all the essential information regarding the MDR changes and the new procedures for Clinical Evaluation and clinical trials, including a learning module on the new standard 14155.

Get ready to learn all you need to know on the new European regulation for medical devices. After the courses, you can take a brief test to receive your certification. Enjoy!

WHO SHOULD PARTICIPATE

Members of the Medtech Industry and Clinical Research Community who want to quickly gain a complete skillset to be successful in the new MDR era.

Professionals involved in the clinical, quality assurance and regulatory fields are all welcome to enroll in this course.

A certificate of attendance will be issued to participants who will complete the final exam.

MEDITRIAL EU COMPLIANCE SERIES

- MDR ROADMAP AND ORIENTATION PROGRAM
  This lecture presents the roadmap of the new European regulation for medical devices and provides an orientation to navigate all the key changes.
- CLINICAL EVALUATION 2020
  This video course and tutorial describe all the steps to produce or update your Clinical Evaluation in accordance with the new MDR.
- NEW ISO 14155 2020 KEY CHANGES FOR YOUR TRIAL
  This video course and tutorial describe the key changes in the international ISO Standard 14155 for Clinical Trials.

"The Meditrial team has created a workshop, curriculum and toolset to enable small companies like ours to understand the impact of the new MDR on our business.

In one day, our team recognized the business segments affected, and what updates to make in order to satisfy MDR requirements.

Russ Sampson
CTO at Ancora Heart, Inc.

CERTIFIED FOR CLINICAL TRIALS.
CERTIFIED FOR CLINICAL SOFTWARE.
TRUSTED EUROPEAN REPRESENTATIVE.

CERTIFIED ISO 9001:2015 COMPANY

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