Introduction To The EU In Vitro Diagnostic Medical Device Regulation (IVDR) Training Course

COURSE CONTENT

- An overview of the new regulation
- The key changes you need to be aware of
- What EU IVDR means in practice
- Where you can find further information
- A quiz to test your knowledge

DURATION: 60 Minutes	ACCREDITATION: SGS
DELIVERY: eLearning	LANGUAGE: English

> "WHAT EU IVDR MEANS IN PRACTICE"

LEARNING OBJECTIVES

Upon completion of this course, learners will be able to:

- Give an overview of IVD Medical Device Regulation
- Understand implementation timescales

AUDIENCE

This course is intended for, but not limited to, regulatory personnel, managers, CEO, CFO, IVD device industry's employees and IVD device users who want to learn about the new regulation.

COURSE CERTIFICATION

On completion of this course:

- The SGS certificate is available to download once you have achieved a pass mark of 80% or more in the final assessment which completes this course.
- You have 12 months access to this course, effective from the date of purchase.

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