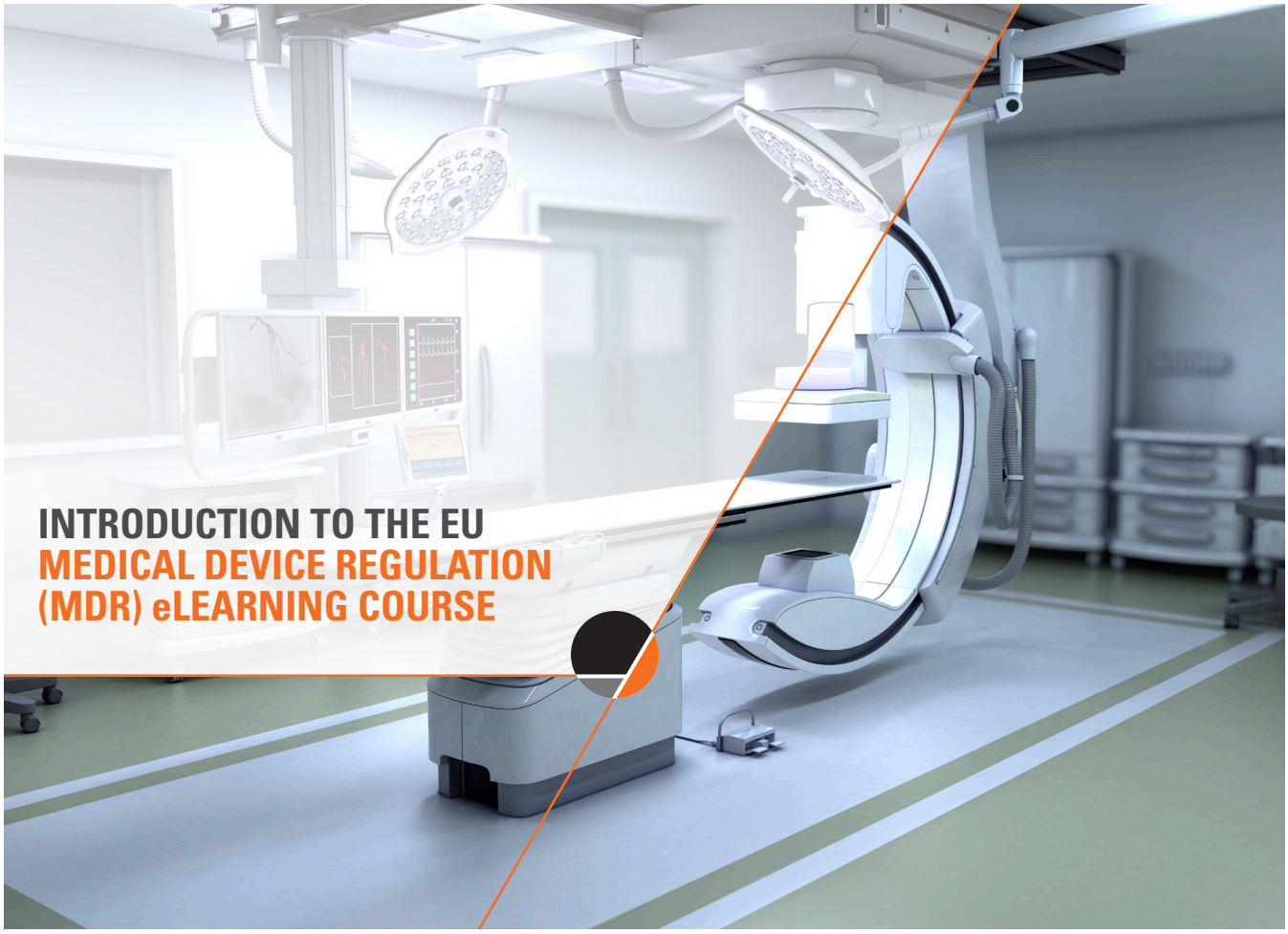

INTRODUCTION TO THE EU MEDICAL DEVICE REGULATION (MDR) eLEARNING COURSE

COURSE OBJECTIVES

This course will provide participants with an overview of the critical changes in relation to EU Medical Device Regulation, along with the implementation timescales.

TARGET AUDIENCE

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

COURSE CONTENT

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

SGS ACADEMY

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COURSE DURATION: 1 HOUR
DELIVERY METHODOLOGY: ELEARNING
COURSE LANGUAGE: ENGLISH
ACCREDITATION: SGS

COMPLETION CRITERIA

Learners will be required to complete the entire course and gain a pass mark of 8/10 or more in the final assessment. Upon successful completion learners will be able to download a certificate of completion. You will be granted 12 months access to this course, effective from the date of purchase.