



Medical Device Directives **In-Vitro Diagnostic Medical Device Directive**

Course Overview

Companies that sell Medical Devices in Europe are required to comply with the Medical Device Directive or the In-Vitro Diagnostic Medical Device Directive. This course will provide an overview of the two Directives. The course will provide information on which directive to comply with, compliance methods and provide guidance on how to become certified to the directives.

Course Objectives

After attending this training, you will understand:

- What the Medical Device Directive is
- What the In-Vitro Diagnostic Medical Device Directive is
- How to classify a product
- The methods for compliance
- How these directives relate to Quality Management System Standards
- How to become certified