





Presented by Management Forum

## The Medical Device School - From Concept to CE Marking

This comprehensive course has been designed to map the processes from device concept to marketing and show how regulatory, quality, clinical and other aspects of device development are joined into a continuous process.

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**Format:** Bespoke training

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**CPD:** 30 hours for your records (depending on your requirements)



Certificate of completion

### **Course overview**

Bringing a medical device to the marketplace is a complex and lengthy procedure which requires experience, knowledge and specialist skills. The contribution to successful market placement comes from many different skilled individuals and organisations who should be aware of all the stages involved and be able to relate their responsibilities to the needs of other professionals, scientists, clinicians and regulatory and quality experts.

This comprehensive course has been designed to map the processes from device concept to marketing and show how regulatory, quality, clinical and other aspects of device development are joined into a continuous process. The programme will commence with a general introduction as to what devices are and how they are developed, and continue with lectures, practical workshops and case studies covering each process applicable to device development, marketing and, eventually, postmarket procedures. There will be ample time throughout the five days for informal discussions with our expert faculty and fellow professionals.

#### Benefits of attending:

Understand the procedures for taking a medical device to the marketplace

- Learn what regulations control the manufacture and marketing of devices in the EU
- Ensure compliance with the MDR
- Gain an insight into different aspects of the process for obtaining the CE mark for a device
- Discover a holistic approach to device development and marketing
- Participate in workshops and learn from other people's experiences
- Understand how other professionals affect the process for bringing a device to market
- Network with participants from organisations similar to your own

### Who should attend

This event has been designed primarily for those who wish to understand the processes involved in bringing a medical device to market. General medical devices as well as active implantable, in-vitro diagnostic and drug device products will be covered in the programme.

The course will be of particular interest to those seeking to introduce new medical devices to the market. Previous delegates who have benefited include regulatory, quality, clinical and marketing managers.

## **Reviews**

#### \*\*\*\*

Very good, I achieved what I set out to do at the beginning of the week.

Natasha Smith Marketing Director Bedfont Scientific Ltd Jun 12 2023

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Really engaging and focused on the EU. Enjoyed the fact this was current and mentioned up to date legislation and regulation updates.

> Chelsey Ludlow Regulatory Affairs Coordinator Vision RT Ltd Jun 12 2023

### **★★★☆**☆

Speaker are very good and expert in their area

**Lutfhi Zarkasyi** Plant Head PT. Forsta Kalmedic Global Jun 12 2023

### \*\*\*\*

I really enjoyed the workshops as it gave the opportunity to interact with the other delegates on the course and discuss and develop learnings from the presentations.



Emma Hickson Research Advisor Convatec Jun 12 2023

# Run this programme in-house for your whole team

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