



Presented by  
Management Forum

# An Introduction to the Design and Development of Medical Devices

23-24 July 2025  
+ 10-11 December 2025

A comprehensive introduction to the design and development of medical devices, including device constituent parts of combination products



**Format:**  
Live online



**CPD:**  
12 hours for your records



Certificate of completion

# Course overview

**This intensive two-day course will introduce those who are new to medical device design and development to the critical elements of the process.** It has been designed to provide delegates with an insight into the tools and techniques required to design and develop a medical device and offer an essential overview of the current regulatory landscape. The importance of safety and efficacy will be covered, as will risk management and documentation. As combination products are a huge market, the programme will also address device constituent parts of combination products.

Attending this event will provide delegates with a comprehensive appraisal of key aspects of this process and an opportunity to discuss the complexities involved with an experienced industry expert.

## Benefits of attending:

- **Gain** a comprehensive overview of the design and development process
- **Comply** with the regulatory requirements and standards
- **Learn** about design controls
- **Review** materials and biocompatibility
- **Access** key information on documentation management and systems
- **Understand** how risk should be managed
- **Consider** human factors and usability studies

## Who should attend?

- Design and development personnel
- Development engineers
- Quality personnel
- Regulatory personnel
- Design control professionals
- Documentation managers
- Programme managers
- Anyone who needs an overview of the medical device design and development process

# Programme

## Day 1

### Overview of the regulations and market routes

- Regulatory pathways
- Medical Device Directive (MDD) vs Medical Device Regulation (MDR) – key differences
- EU vs US (FDA) – markets to consider
- Medical device vs combination product (drug/device and device/drug) – which regulation applies?
- Device classification and the implications for your product
- Resources and sources

### The design and development process

- The stages of design and development
- Key considerations
- Terminology
- Intended use
- Project complexity
- Mandatory requirements
- Design and development tools
- Inspiration, innovation and determination
- Materials and biocompatibility
- DFX, Design for.....?
- Manufacturing – key considerations

## Day 2

### Design control

- Appropriate design and development planning
- Translation of marketing requirements
- SMART design inputs
- Is a trace matrix appropriate?
- Meaningful design outputs
- Verification and validation
- Design reviews
- Design transfer
- Design history file vs technical file
- Change control
- Notified Bodies (NB)

### Risk management – what is required?

- What is risk management and when should it be applied?
- What does the guidance say?
- Help or hindrance?
- How to implement a practical risk management plan
- Tools and techniques to help you succeed

### Clinical evaluation, human factors and usability – how to comply

- Planning your clinical evaluation
- How to incorporate human factors and usability studies into your design and development process – MDR and FDA requirements
- User instructions
- Training considerations – when and who do you need to train?
- Formative studies
- Validation/summative studies

# Presenter



## David Howlett

In 2003 David established PharmaDelivery Solutions Ltd as a specialised consultancy service in the field of drug delivery combination products (especially respiratory) device technology. This has led to involvement in projects with focus in pulmonary, nasal and other delivery routes, with an international client base. Much of the activity of PharmaDelivery Solutions Ltd is focused in the area of development programme support, regulatory GAP analysis and generation of documentation supporting development and test programmes, together with data review and contingency evaluation.

David has over 35 years experience in the development, industrialization and approval of inhalation drug delivery systems, combination products and medical devices.

PharmaDelivery Solutions Ltd has provided input ranging from GAP analysis and comment to complete remedial implementation in areas including

- Design Control
- Risk Management
- Materials strategies
- Regulatory documentation
- Technical reviews and opinion

In addition to activities supporting commercial organizations, David has been involved in the following roles;

Honorary Teaching Fellow in the School of Pharmacy and Pharmaceutical Sciences at the University of Manchester and is author/ tutor for the Pharmaceutical Industry Advanced Training (PIAT) MSc module on Inhalation dosage forms.

A UK national expert representing the British Standards Institute on ISO TC84 developing new international standards for pulmonary and nasal delivery devices syringes and catheters.

David has also worked with the United Nations and various national governments to develop and establish transition strategies from the use of CFC in Metered Dose Inhalers and to secure appropriate budgets from the Multi-lateral fund for the implementation of the Montreal Protocol in emerging markets around the world.

# Course dates

**23-24 July 2025**

**Live online**

09:30-17:00 **UK (London)** (UTC+01)

Course code 14869

GBP **1,299** ~~1,499~~

EUR **1,819** ~~2,099~~

USD **2,087** ~~2,399~~

**Until 18 Jun**

**10-11 December 2025**

**Live online**

09:30-17:00 **UK (London)** (UTC+00)

Course code 15140

GBP **1,299** ~~1,499~~

EUR **1,819** ~~2,099~~

USD **2,087** ~~2,399~~

**Until 05 Nov**

## How to book



**Online:**

[ipi.academy/2047](https://ipi.academy/2047)

Alternatively contact us to book, or if you have any queries:



**Email:**

[info@ipiacademy.com](mailto:info@ipiacademy.com)



**Phone:**

[+44 \(0\)20 7749 4749](tel:+442077494749)

## Discounts

- Booking more than one delegate on any one date qualifies for a **15% discount** on the second and subsequent places.
- Most events qualify for an **early booking discount** prior to 6 weeks before the course date. Be sure to check on our website, where the latest discounts will be shown.

## Further information

### Fee

The fee includes all meals and refreshments for the duration of the course (for venue-based courses) and a complete set of course materials (provided electronically). If you have any particular requirements, please advise customer services when booking.

### Please note

IPI Academy (and our training partners) reserve the right to change the content and timing of the programme, the speakers, the date and venue due to reasons beyond their control. In the unlikely event that the course is cancelled, we will refund the registration fee and disclaim any further liability.

### Terms and conditions

The rest of our terms, the event cancellation policy and the terms and conditions are on our website, please visit [ipi.academy/content/terms-and-conditions](https://ipi.academy/content/terms-and-conditions)

# Reviews



**Good speaker, clearly very knowledgeable in the area. He was happy to have conversations about any specific scenarios/examples/questions we had. Very friendly and i applaud his speaking almost continuously on this topic for two days**



**Michaela Hume**  
Engineer  
42 Technology  
Jul 24 2024



**Very good, lots of relevant material and clearly outlined.**



**Kurt Blinston**  
R&D Manager  
Medtrade Products Ltd  
Jul 24 2024



**Good content, covered all aspects that I required.**



**Anna Short**  
Design Manager  
DM Orthotics  
Jul 24 2024



**David was a very good presenter in going through the content in detail and made it easy to follow. Great with regards to answering questions people had. Very good webinar for all aspects, would recommend to others**



**Simran Sandhu**  
R&D Scientist  
Medtrade Products Limited  
Jul 24 2024

## Run this programme in-house for your whole team

Coming to IPI Academy for your in-house training provides an all-inclusive service which gives you access to a wide variety of content, learning platforms and delivery mechanisms as well as your own personal training adviser who will work with you from the initial enquiry through to feedback and follow-up after the programme.

With over 600 trainers, all practitioners and experts across a huge range of fields, we can provide the training you need, where you need it, when you need it, and at a price which suits your budget. Our approach to tailored learning and development consists of designing and delivering the appropriate solution for each client.

For your FREE consultation and to find out more about how we can work with you to solve your training needs, please contact our training advisers:



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**IPI**  
Academy

IPI Academy is a training initiative of Falconbury and Management Forum; leading providers of industry training for over 30 years, based in the UK.

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