





Presented by Management Forum

An Introduction to Pharmaceutical Packaging

30 September-2 October 2025

This course provides an essential overview of pharmaceutical packaging and will be valuable for newcomers to this field. It will also act as a good refresher for those with more experience.



Format:

Live online

(1)

CPD:

12 hours for your records

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Certificate of completion

Course overview

Gain crucial insights into the unique issues and challenges of pharmaceutical packaging to ensure product safety, compliance, and market success.

Effective packaging of pharmaceutical and healthcare products is critical not only for maintaining product integrity and safety but also for regulatory compliance and market acceptance. Packaging serves multiple functions beyond containment, including protection from environmental factors, ensuring product stability, and providing essential information to healthcare providers and patients. Understanding the intricacies of pharmaceutical packaging is vital for professionals involves in drug development, manufacturing, regulatory affairs, quality control, and logistics.

This introductory course is designed to equip delegates with a foundational understanding of pharmaceutical product packaging. It begins with a comprehensive regulatory overview, ensuring participants grasp the essential requirements and compliance standards governing pharmaceutical packaging. The programme also delves into crucial topics such as the selection of packaging materials, considerations for compatibility and stability (including ICH testing and extractables/leachables studies), and the role of packaging in new product development.

Additional topics include:

- Pharmaceutical packaging formats and materials available advantages and applications
- Trade and transit requirements
- Environmental considerations
- Artwork design essentials
- Specialised areas child-resistant closures and tamper-evident packaging

By the end of the course, participants will gain a comprehensive understanding of the key aspects of pharmaceutical packaging, enabling them to contribute effectively to the successful packaging of pharmaceutical packaging, enabling them to contribute effectively to the successful packaging of pharmaceutical products within their organisations.

Benefits of attending

- Gain a useful insight into packaging component and material selection
- Learn key properties of various packaging materials/systems
- **Appreciate** pack testing and evaluation
- Better understand packaging component specifications
- Hear about printing processes and controls
- Acquire knowledge on artwork generation and control
- Ensure that you comply with the regulatory requirements
- Learn about transit packaging
- Consider trade/supply chain requirements

Who should attend?

Whether you are new to the industry, have a basic understanding of pharmaceutical packaging or are more familiar with the area but looking for a refresher, this course will provide you with valuable knowledge and insights from an expert who has worked in the field for more than 25 years.

Those who would benefit from attending include:

- Account managers
- Artwork producers
- Auditors
- Business developers
- Clinical trial suppliers
- Logistics personnel
- Packaging design/labelling personnel
- Project managers
- Purchasers
- Quality assurance and control personnel
- Regulatory personnel
- Suppliers to the industry
- Technical writers



Programme

Day 1

NPD process

The role of pharmaceutical packaging

Regulatory and GMP Requirements

- Agencies, guidelines and legislation
- Dossier requirements
- International Conference on Harmonisation (ICH)
- Common Technical Document (CTD)
- Summary of Product Characteristics (SmPC)
- Differences between submissions in the EU and USA
 - O Bar coding (briefly) EAN, 2D datamatrix, QR
 - Counterfeiting and product security
 - Drug Quality and Security Act
 - Falsified Medicines Directive (FMD)

Product security and emerging pharma legislation

Choice of pharmaceutical packaging

- Compatibility and ICH testing
- Testing and evaluation
 - Extractables and leachables

Pack testing and evaluation - ICH

ICH testing

- Stability testing
- Functionality testing

Special climatic conditions

Secondary and tertiary packaging materials Part 1

- Print processes
- Labels

Special aspects of pharmaceutical packaging

Day 2

ICH testing continued.

- Extraction/migration studies
- Leachables, Extractables & Toxicological Issues
- Case Study
- The EU Perspective
- The FDA Perspective: USP <661>

Key properties of various primary packaging materials Part 1

- Glass
- Plastics (part 1)

Primary packaging materials Part 2

• Plastics (part 2)

Primary packaging materials Part 3

- Plastics (part 3)
- Sachets
- Blisters
- Tubes

Day 3

Primary packaging materials Part 4

- Closures
- Active packaging
- Aerosols

Secondary and tertiary packaging materials Part 2

- Leaflets
- Cartons
- Trade/supply chain requirements
- Specifications
- Environment and sustainability

Trade / transit requirements

Interactive session (Case study: group exercise & feedback)

Takeaway and key messages

Presenter



Chris Penfold

Chris Penfold is an experienced Freelance Packaging Development Specialist and Consultant; a self-motivated, achievement oriented, confident and creative leader with a proven track record in general and technical management. He is a packaging professional with over 25 years packaging development and NPD experience working on £million-brands in the OTC, healthcare and Rx pharma arenas for 'blue-chip' multinational companies such as Glaxo (GSK), CIBA (Novartis), Boots Healthcare and Reckitt Benckiser Healthcare.

He is an MBA graduate and Chartered Marketer with extensive cross-functional experience in a broad spectrum of related disciplines, including marketing, supply chain, QC and production. An 'International Player' with experience working in various European countries, Middle East, North America and extended business trips to the Far East. Underpinned by proven interpersonal skills, the ability to make things happen and experience gained from independent international consultancy projects and running his own business, Design Cognition Ltd.

Course date

30 September-2 October Live online

2025

09:00-16:45 **UK (London)** (UTC+01)

Course code 14945

GBP **1,599** 1,899

EUR **2,239** 2,659

USD 2,571 3,039

Until 26 Aug

How to book



Online:

ipi.academy/2287

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



Email:

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Phone:

+44 (0)20 7749 4749

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

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

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Reviews

Really good. Very thorough. Well constructed in a way that allows the presentation to be used as a reference doc later. The speaker was excellent.



David Bellars

Supply Chain Manager Cycle Pharmaceuticals Ltd May 20 2024



Really enjoyed it. Thought it was well presented and clearly explained by Chris. I would recommend this course to others. Good pace and the information was clearly delivered.



Andy Borg-Myatt

Artwork Specialist Kyowa Kirin International Jan 31 2023



Very informative course with lots of training material



Sanna Sutinen

Specialist, Quality Systems FinVector Oy Jan 31 2023



Webinar was very good and informative. It gave a very good overall regulatory summary and the fact that the group was small enabled to participate and ask direct live questions during the course of the agenda, which was for me very helpful.



Orit Shahaf

Director, Regulatory Affairs Taro Pharmaceutical Industries Jan 31 2023

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