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Presented by
Management Forum

Making Financial Sense of GMP: Pharmaceutical Quality System (PQS)

3 October 2025
+ 13 February 2026

In this talk, explore how the Pharmaceutical Quality System (PQS) impacts the financials of an organisation and how it can enhance an organisation's financial strength.



Format:
Live online



CPD:
1.5 hours for your
records



Certificate of
completion

Course overview

Pharmaceutical Quality System (PQS) is outlined in ICH Q10 guideline. The PQS is referred to as ICH Q10 Model and is based on International Standards Organisation (ISO) quality concepts, includes applicable Good Manufacturing Practice (GMP) regulations and complements ICH Q8 'Pharmaceutical Development' and ICH Q9 'Quality Risk Management'.

The implementation of ICH Q10 model aims to achieve three main objectives and complements or enhances regional GMP requirements:

- Achieve product realisation
- Establish and maintain a state of control
- Facilitate continual improvement

There is intrinsic financial impact on each of these objectives and understanding it supports not only improving quality efforts, but aligns with financial and budgetary goals.

ICH Q10 is a model and regulatory expectation. What would happen if the regulatory expectation was removed?

Benefits of attending

Attending this session will help deepen your knowledge and support your understanding on how the objectives of ICH Q10 model benefits an organisation's financial performance.

The benefits of attending this session include:

- **Learn** about the impact of PQS on the gross profit, operational costs and net profit
- **Explore** how PQS can support efforts to reduce costs and improve profits
- **Discuss** PQS elements that contribute to cost reduction
- **Understand** how financial decisions don't have to compromise quality
- **Consider** what would happen if PQS was not a regulatory expectation

Who should attend?

This session is designed for senior life science professionals involved with Pharmaceutical Quality System (PQS) either directly or indirectly.

The stakeholders are typically from executive and management roles responsible for and involved in decision making and striving to balance financial goals with compliance and quality requirements.

Other interested stakeholders not directly involved with compliance or quality include:

- Chief Operating Officers
- Chief Financial Officers
- Finance and costing analysts

Presenter



Jitan Bhudia

Jitan Bhudia is a highly accomplished Pharmaceutical Consultant with extensive expertise in Good Manufacturing Practice (GMP), Good Distribution Practice (GDP), and regulatory compliance within the pharmaceutical industry. With a career spanning over two decades, Jitan has a proven track record of leading business operations, ensuring regulatory compliance, and driving commercial success in pharmaceutical manufacturing and distribution.

As the co-founder of jarmatrixpharma.com, a site dedicated to help organisations “manage knowledge”, Jitan has provided specialized consulting and tailored training solutions for pharmaceutical manufacturing. He has designed and implemented Quality and Document Management Systems, supported regulatory audits, and worked closely with industry stakeholders to ensure compliance with stringent pharmaceutical regulations. His ability to develop cost-effective compliance solutions and deliver expert training on quality management strategies has positioned him as a trusted advisor in the field.

Previously, Jitan served as the Owner, Director, and Head of Production at Percuro Medica Limited, where he played a pivotal role in establishing and expanding the company. Under his leadership, the company gained essential licenses from the MHRA and the Home Office for the manufacture and distribution of controlled drugs. His expertise in facility design, validation, and risk management helped secure major contracts with pharmaceutical companies.

With his deep industry knowledge, regulatory expertise, and business acumen, Jitan continues to support pharmaceutical organizations in navigating complex compliance landscapes, optimizing quality systems, and achieving operational excellence.

Course dates

3 October 2025

Live online

12:00-13:30 **UK (London)** (UTC+01)

Course code 15637

GBP **175** ~~200~~

EUR **245** ~~280~~

USD **280** ~~320~~

Until 26 Sep

13 February 2026

Live online

12:00-13:30 **UK (London)** (UTC+00)

Course code 15638

GBP **175** ~~200~~

EUR **245** ~~280~~

USD **280** ~~320~~

Until 06 Feb

How to book



Online:

ipi.academy/3236

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



Email:

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Phone:

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

Discounts

- Booking more than one delegate on any one date qualifies for a **30% 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

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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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