IPI Academy

MANAGEMENT forum

Presented by
Management Forum

Making Financial Sense of GMP: Validation Master Plan (VMP)

10 October 2025
+ 27 February 2026

In this talk, explore how the Validation Master Plan (VMP) impacts the financials of an organisation and discover how a well managed VMP enhances an organisation's financial strength.

Format:
Live online

CPD:
1.5 hours for your records

Certificate of completion

Course overview

Validation Master Plans (VMP) govern validation activities for an entire organisation. The VMP serves as the validation roadmap, setting the course, justifying the strategy, outlining the preliminary test and acceptance criteria, and documenting the necessary programs that ensure a continuing state of validation.

The VMP includes the following elements as expected by the regulatory agencies:

- Plan - documenting intent and pathway
- Qualification - confirming design and intent
- Process Validation - assures process consistency
- Reports - Summary of test results and acceptance criteria
- Compliance programs - Ensuring continuous state of validation

The VMP has cost implications from designing and executing to reporting and maintaining. Are these costs justified? What would happen if VMP was not part of regulatory expectations? Would organisations be motivated enough to implement VMP without regulatory expectations?

Benefits of attending

Attending this session will help deepen your knowledge and support your understanding on how the Validation Master Plan (VMP) benefits an organisation's financial performance.

The benefits of attending this session include:

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

Who should attend?

This session is designed for senior life science professionals involved with Validation Master Plan (VMP) 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

10 October 2025	**Live online** 12:00-13:30 **UK (London)** (UTC+01) *Course code 15643*	GBP **175** ~~200~~ EUR **245** ~~280~~ USD **280** ~~320~~ **Until 03 Oct**
27 February 2026	**Live online** 12:00-13:30 **UK (London)** (UTC+00) *Course code 15644*	GBP **175** ~~200~~ EUR **245** ~~280~~ USD **280** ~~320~~ **Until 20 Feb**

How to book

Online:
ipi.academy/3237

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 **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 the 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

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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