EPiC Virtual Compliance Symposium 2024 - From Complaint to Compliant: A Practical Guide to Overcome Current GMDP Challenges

EPiC is proud to present our full-day virtual symposium 2024 event featuring an esteemed lineup of former MHRA Inspector and industry speakers on Thursday 28th November 2024

ABOUT

Join us as we explore the latest regulatory updates and guidance and provide practical guidance to conquer compliance challenges facing the pharmaceutical sector.
We have completed an extensive review of the compliance challenges faced by our clients to identify recurring issues that commonly result in inspection deficiencies and referral to MHRA Inspection Action Group.

We will reflect on the current reality, share common causal factors impacting GMDP compliance, and provide you with practical guidance on how to reframe the situation to avoid complaints and move towards compliance. Our objective is to combine valuable insights using real world examples and experiences from across our consortium of expert pharmaceutical consultants, to provide the knowledge and offer practical solutions to overcome current GMDP compliance challenges.

Expect an interactive and informal experience carefully crafted to include an update on pharmaceutical guidance and regulations, current GMDP compliance challenges, practical guidance on how to avoid Annex 15 equipment qualification and validation common pitfalls, helpful tips to prevent misunderstandings and quality problems associated with outsourced activities, how to 'see the wood for the trees' and maintain perspective critical to GMDP compliance, GDP Master Class exploring current challenges and opportunities, essential factors for effective GMDP refresher training, and a call to action for QPs on how to influence GMDP compliance.

Our panel of experts, drawing on their extensive knowledge and regulatory experience are committed to delivering a thought-provoking and informative event with an emphasis on providing practical guidance. Engage with them throughout the day and participate in a panel session to ensure all your pressing questions are addressed.

Discover how our expertise can add value to your organisations' s compliance improvement strategy by equipping you with the knowledge and practical solutions to overcome current GMDP compliance challenges.

WHO IS IT FOR?

The event is ideal for pharmaceutical industry professionals in quality control, quality assurance, supply chain and production management, as well as Qualified Persons, Responsible Persons, and regulatory and compliance specialists involved in the manufacture and distribution of medicines.

WHAT WILL I LEARN?

The event will provide the latest information and guidance on changing legislation and GMDP compliance challenges to enable you to enhance your knowledge and expand your skillset by broadening your understanding of causal factors and obtain practical solutions to improve GMDP compliance. Through presentations, case studies and Q&A sessions, highlighting regulatory expectations, industry best practice and example deficiencies, you will learn how to avoid common pitfalls to overcome current GMDP compliance challenges.

WHY SHOULD I ATTEND ?

- **LEARN** how to utilise practical guidance to overcome GMDP compliance challenges
- **NETWORK** with other Pharmaceutical Industry delegates
- **CPD** Continuous Professional Development to improve workplace skills and knowledge
- **CONNECT** with EPiC Directors and Senior Managers and expert Ex MHRA Inspectors
- **ENGAGE** in our presentations and panel discussions and hear Industry-focused case studies from our regulatory experts
- **VALUE** for money, presenting a wide range of topics from regulatory experts

NOT AVAILABLE ON THE DAY?

We understand your schedule may not allow you to access all the event sessions. A recording of the event will be made available after the event and will remain open 24/7 for 30 days. Attendees will receive an email once the recordings have been uploaded and can access the presentations whenever it is convenient during this time.

AGENDA

Thursday 28th November 2024 (09.00 to 4.30 pm GMT) to include:

- **Welcome and Introductions** (Richard Andrews, Managing Director & Senior Consultant, EPiC Auditors)

- **Recent Updates to Guidance & Regulations** (Richard Andrews, Managing Director & Senior Consultant, EPiC Auditors)

- **GMDP Compliance - Reflections of Current Reality** (Darren Jones, Director & Senior Consultant, EPiC Auditors)

- **Annex 15: Equipment Qualification and Validation – Avoiding Common Pitfalls** (Mark Thompson, EPiC Auditors Consultant)

- **Coffee Break – 20 Minutes**

- **Chapter 7: Outsourced Activities - Preventing Misunderstandings and Quality Problems** (Michelle Yeomans, Operations Manager & Senior Consultant, EPiC Auditors)

- **Seeing the Wood for the Trees - How to Identify, Prioritise and Maintain Perspective Critical to GMDP Compliance** (Lewis Corbett, Business Manager & Senior Consultant, EPiC Auditors)

- **Lunch Break – 60 Minutes**

- **GDP Master Class - Current Challenges and Opportunities** (Tony Orme, EPiC Auditors Consultant)

- **Effective GMP Refresher Training - Essential Factors for Success** (Expert panel ideas and discussion)

- **Coffee Break – 20 Minutes**

- **How to Influence GMDP Compliance – A Call to Action for QPs** (Phil Rose, EPiC Auditors Consultant)
- **Q&A Panel Session** (Expert panel answer your burning questions)

CONTINUOUS PROFESSIONAL DEVELOPMENT

EPiC is a member of the CPD Certification Service which is recognised as the world's leading and largest CPD accreditation service with over 25 years of experience.

An application will be made to accredit the event and award CPD points. If successful, Certificates will be sent to attendees within approximately 2-3 weeks after the event.

TICKETS PRICE

The ticket price is £300 + VAT per delate

REGISTER

To Register and book your place follow this link: https://academy.epic-auditors.com/register/?utm_source=mpharma

If you have any questions, then email EPiC at enquiries@epic-auditors.com