



Integrated Quality Management Systems

John Jolley FRPharmS FCQI CQP

A black binder with a silver pen lying on top of it. The binder has a white label that reads "QUALITY MANAGEMENT". The background is a blurred office setting with a computer monitor and papers.

QUALITY MANAGEMENT

**Webinar for members of EIPG
in conjunction with
PIER and University College Cork**

Click here to register

**Monday 15th October 2018
at 16.00 GMT (17.00 CET)**

About the Speaker

John Jolley has held positions in Clinical Research, Product Registration, Manufacturing, Quality Assurance, and General Management. He was Technical Director for Boehringer Ingelheim UK for 15 years before forming an international consultancy PharmaConsult Europe which became PharmaConsult Global. He is a practising Qualified Person (QP) whose experience includes working in sterile product manufacture and clinical trials.

Overview of Webinar

EIPG has invited John Jolley, who has gained a broad and consolidated experience in pharmaceutical quality, to reveal the principles of a quality system and its practical application in the light of the most recent approach to quality management in the pharmaceutical industry.

Supply chains are becoming increasingly complex, and regulations around the quality of products and services are becoming more stringent. Organizations are thus under increasing pressure to demonstrate their commitment to quality based on the implementation of effective quality management systems, good corporate governance practices, sound environmental policies, and a robust safety culture. As a result, companies are seeking to deploy Integrated Quality Management Solutions that can bring business units under one single program, replacing the segmented decision making processes of the past, which can lead to greater controls in supplying.

Learning Outcomes

By the end of this presentation, you will be able to describe:

1. The requirements of an Integrated Quality Management System
2. The principles of quality risk management and their applications
3. The management of product and process deviations , CAPA and Change control
4. Audit Management of Internal and Regulatory Inspections.
5. Management of Enterprise Resource Planning and supplier quality management.

To Join the Webinar

Please register for the event by filling out the form at https://docs.google.com/forms/d/e/1FAIpQLSdB2_nzfddcwwE57GtYqPjalMHiW_gvI22RkoI1ZMHJ9M4wtQ/viewform. Further instructions will then be sent by e-mail.

Continuing Education:

A certificate of attendance will be issued after the webinar if requested upon registration. The session will be an hour of Continuing Education.