QUALIFICATION OF SUPPLY CHAINS AND SUPPLIER AUDITS: REMOTE AND ON SITE

CPD/L/___/2026 (TBC)

Why is this course essential?

Manufacturers are required to qualify their supply chains and this means different approaches depending on the type of supplier, such as API, Excipient, 3rd party Manufacturers, Primary Packaging, Laboratories and Printed Matter etc....
The program will provide the attendees with the requisite skills to conduct risk assessment of suppliers, determine levels of documentation and evidence required, develop QTAs and gain experience in remote auditing practices.

Training Modules Include

- PICs rules for qualification of supply chains
- Risk Assessment of suppliers
- Types of suppliers and levels of qualification
- Expected documentation and checklists
- Set up for site and remote audits
- Quality Technical Agreements (QTAs)
- **Simulation of a remote audit**

Mr. Steve Williams
Director & Senior GMP Consultant of CBE Pty. Ltd.

Steve has over 50 years' experience in the Biotechnology, Pharmaceutical and Medical Device industries in quality and manufacturing, including 30+ years consulting in GxP Quality Management and Regulatory compliance. Steve conducts FDA and EU/TGA/PICs compliance audits and gap analyses, assists companies in remediation programs and prepares companies for regulatory inspection. Steve also specializes in GxP improvement programs and re-engineering Pharmaceutical Quality Systems.

Dr. Lyndall Brennan
Senior Consultant of CBE Pty. Ltd.

Lyndall is a former TGA GMP Inspector and senior pharmaceutical quality leader with over 25 years' experience in GMP systems, regulatory compliance, risk management, and data integrity. She has helped shape global standards and specialises in turning complex regulatory expectations into practical, inspection-ready solutionse. Lyndall helps medicine manufacturers strengthen quality systems, embed compliance in operations, and improve patient-safety outcomes.

Mr. Maurice Parlane
Director & Senior GMP Consultant of CBE Pty. Ltd.

A Professional engineer and technical manager with over 30 years' experience in manufacturing management, operational and consulting roles within human & animal health, biotechnology medical device, and medicines sectors. Maurice has been involved with and managed numerous projects of varying scale related to compliance improvement and manufacturing effectiveness for clients for NZ, Australian, US and EU markets.

13 MAY 2026

09:00-17:00

Course Fee (7 CPD Hours)
HKPMA member HKD2,400 Non-Member HKD4,000

Venue
Function Hall, Science Park
1/F, Building 12W, 12 Science Park West Avenue

Important notes:

- Registration deadline: 27th April 2026 (Monday).
- Registration confirmation will be sent on/before 30th April 2026 (Thursday).
- Training material will be sent via email. No hardcopy will be provided.
- Coffee break and lunch are included.
- HKIB reserves all rights to the postponement or cancellation of this training.

For Enquiry
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WEBSITE
https://atp.hkib.org.hk/

Program organized by
Hong Kong Institute of Biotechnology