



SAMANTHA CLACK

Independent Consultant

BSc (Hons), PGDip, MSB, CBiol

AREAS OF EXPERTISE

TRAINING

IRCA Lead Auditor
Self-Inspection and Internal Audits
GMP
Cogent GDP/RP/RPi
QP
PQS
Solid Dose, Steriles, Packaging

AUDITING

Parenteral, Ophthalmic, Solid Dose,
Non-sterile Liquid, Cream and
Ointment manufacture and packaging
API, Excipients, packing material
suppliers
Storage and Distribution
PQS

CONSULTANCY

Quality Management Systems
GMP Manufacture and Packaging
Wholesale Operations

KEY KNOWLEDGE AREAS

Qualified Person – Parenteral, Solid
Dose, Biotech
Facility and System Audits
Pharmaceutical Quality Systems
GAP Analysis
Inspection Remediation
Qualified Person Training and
Mentoring

CONTACT

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EXPERIENCE

- > Pharmaceutical Auditor – 20 Years
- > Qualified Person – 15 years
- > Responsible Person – 10 Years
- > Pharmaceutical Quality Systems Implementation and Management
- > Manufacture and Packaging of parenteral, ophthalmic, biotech, solid dose, liquid, cream and ointments products
- > Regulatory Inspection Management and Remediation
- > Wholesale Dealers Quality Management System implementation and Management.
- > QP/RP, GMP/GDP and Auditor Training

KEY JOB ROLES

- > Director/Consultant QP – Quaph Consulting Limited
- > QA Manager/Deputy QA Director – Eisai Manufacturing Limited
- > Group Leader/QP – Eli Lilly and Company Limited
- > Senior Quality Associate – Corporate QA - Eli Lilly and Company Limited
- > Deputy Manager, Sterile and Non-Sterile Topicals manufacture – Hoeschst Marion Roussel

EDUCATION

- > Post Graduate Diploma – Pharmaceutical Quality and GMP – University of Strathclyde
- > Post Graduate Diploma – Business Management – University of Reading
- > Bachelor of Science – Microbiology and Biochemistry – University of Reading

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Training www.nsf.org/info/pharma-training