Mark has over 17 years of international experience in a wide range of roles within Pharmaceutical, Medical Device, Sterile and FMCG manufacturing Sites. His experience ranges from Product and Process Transfers, Risk Management, Process Validation, Computerized Systems Compliance, Quality Assurance, Quality Systems, Auditing, and Research & Development.

In his career, he has led and participated in multiple projects involving product and process transfers of new product introductions, process validation, equipment commissioning and qualification, computerized systems validation, continuous improvements, product development and process development.

Information