Mr Darren Jones

Consultant
BSc (Hons), C Biol, MRSB

Darren joins us from the MHRA, where he spent four years as a GMP Inspector. He inspected many sites of varied dosage forms worldwide.

He has a Degree in Applied Biology and joined the industry in 1990, with AstraZeneca, where he spent 18 years in a number of roles, including QC, QA and Production roles. He spent time supporting a large API facility, tablets and packaging operations although much of this career was focussed on complex aseptic and terminally sterilised production, in both the commercial and clinical trials areas.

Darren became a transitional QP under the Clinical Trials Directive 2001/20/EC and went on to qualify as an eligible QP via the permanent provisions on 2001/83/EC.

Areas of Expertise:

- GMP auditing of facilities and processes
- Sterile manufacture
- Clinical trial manufacture
- Supply chain excellence
- Microbiological QC