
Professional Career Narrative Summary

N Peter Maurice, LMSSA, MBBS, DCH, FFPM Independent Clinical Research Quality Assurance and Ethics Consultant

Peter Maurice is a London Medical graduate with 10 years UK clinical experience as a Paediatrician, and Lecturer when in the BMRC Perinatal Research unit at London University.

Subsequently entered the Pharmaceutical Industry in Switzerland, and was involved in the clinical development of many Ciba CVS, CNS and Antibacterial drugs. He then managed an in-house human pharmacology unit and became Global Head of Clinical QA. He developed Ciba's GCP SOPs, training programmes and the global Clinical Audit facility.

**Since 1995 has been a free-lance consultant.
Consulting, lecturing and training on GCP, Clinical QA/Audit, drug safety and clinical research ethics are his areas of expertise.**

Elected chairman of the Swiss (Interpharma) and European Industry (EFPIA) GCP committees he was involved in:

- The first EC Note for Guidance on GCP,
- The ICH GCP harmonisation process(1991-96) and
- Chaired the Experts which developed the ICH GCP Guideline.

He also represented the International Pharma Industry (IFPMA) during the development of:

- The CIOMs International Ethical Guidelines for Biomedical Research
- The WHO GCP Guidelines.

He played a leading role in the revision of the latter and has drafted
The forthcoming WHO Handbook on Health Care Research.

A highly active member of the DIA for >15 years, Peter was on the European Steering Committee and was elected as the 1st European Director on the DIA Board. Received the DIA Award for Outstanding Service in 1993.
Founder member and 1st Executive Committee member SwAPP, >1995

In the academic domain he is External Examiner for higher Medical degrees at Basel and Hong Kong universities and in 2000 initiated Europe's first independent not-for-profit university based clinical research QA service, QuaSwiss, provided by free-lance experts. He has also trained in Industry, Academic and Regulatory GCP programmes in Australia, Canada, China, Denmark, France, Germany, Hong Kong, Japan, Malaysia, New Zealand, Poland, Russia, Slovakia, Switzerland, Thailand, the UK & USA.

1993: 1st MD to become a Certified ISO 9001 auditor. Has performed Clinical QA activities in Europe, N America and Asia for industry, government and at WHO.

2002: Received Institute of Clinical Research, UK, "Outstanding Contribution to Clinical Research" Award.