Curriculum Vitae

Personal information

Pernille Sterling

Work experience

1. Employer: The Danish Medicines Agency
   - Start date: 122004
   - End date:
   - Position: Chief advisor, nonclinical assessor,
   - Activities: _ Clinical trial applications, nonclinical safety evaluation through out the development process. _National scientific advices. _Participant in Clinical Trial Coordination Group (CTCG), incl. ad hoc subgroups in NcWP. Project coordinator. GLP. First in human & early clinical trials. Advanced Therapeutic Medicinal Products and innovative products. Radiopharmaceuticals
   - Country: Denmark
2. Employer: H. Lundbeck A/S
   - Start date: 092002
   - End date: 042004
   - Position: Compliance Manager
   - Activities: Internal audits, maintenance and development of controlled documents, draftet internal guidelines, changes, deviations etc. GMP_related assignments in relation to analysis laboratories.
   - Country: Denmark
3. Employer: National Environmental Laboratories
   - Start date: 011998
   - End date: 052002
   - Position: External consultant
   - Activities: Consultant on the National Monitoring Program for Aquatic Environments (NOVA 2003). Monitoring toxicological responses to pollutants. Responsible for project management, training of staff, data analysis & interpretation
   - Country: Denmark

Education and training

1. Subject: Southern University, Odense
   - Start date: 1992
   - End date: 2000
   - Qualification: Master of Science, toxicology
   - Organisation: Research on the effects of hormone mimicking xenobiotics on animal species and the complex of problems associated with hormone disturbances & regulation.
   - Country: Denmark
2. Subject: Surrey
   - Start date: 2009
   - End date:
   - Qualification: Pathology for Regulators.
   - Organisation: Advanced Modular training programme in applied toxicology for non_clinical assessors. 9_10 juli, Surrey, UK.
   - Country: United Kingdom
3. Subject: MIND, Pharmaceutical Faculty, Copenhagen University
   - Start date: 2005
   - End date:
   - Qualification: Non_clinical Safety; Toxicology
   - Organisation: Non_clinical Safety & toxicology_ studies, signals and regulation
   - Country: Denmark

Additional information

Publications

Projects

Memberships

Other Relevant Information

Expert/paticipant at EMA sub groups

Clinical Trial Coordination Group (CTCG)