Curriculum Vitae

Personal information

Nanna Aaby Thirstrup

Work experience

1. Employer: Danish Medicines Agency
   - Start date: 012021
   - End date: current
   - Position: Director of Department
   - Activities: Director of Department, Quality evaluation & Clinical Trials
   - Country: Denmark
2. Employer: Danish Medicines Agency
   - Start date: 062001
   - End date: 012020
   - Position: Team Manager, Pharmaceutical Quality, Biologics
   - Activities: Team Manager for the biological pharmaceutical qaulity team (Marketing Authorisations and clinical trials). Scientist with responsibility for evaluation of the Quality (CMC) part of dossiers
   - Country: Denmark
3. Employer: Danish Medicines Agency
   - Start date: 041995
   - End date: 052001
   - Position: Scientist in the Danish Medicines Agency Biological Control Laboratory
   - Activities: Scientist with responsibility for evaluation of the Quality (CMC) part of dossiers and laboratory control of authorised biological products on the Danish / EU marked
   - Country: Denmark
4. Employer: Statens Seruminstitut, Biochemical Sector, Immunological department
   - Start date: 081993
   - End date: 031995
   - Position: Research assistant
   - Activities: Research assistant on a malaria vaccine research project
   - Country: Denmark

Education and training

1. Subject: The Danish University of Pharmaceutical Sciences
   - Start date: 081986
   - End date: 071993
   - Qualification: Master of Science, Pharmacy
   - Organisation:
   - Country: Denmark

Additional information

Publications

Ionic strength_dependent denaturation of Thermomyces lanuginosus lipase induced by SDS. Arch Biochem Biophys. 2011 Feb 1;506(1):92_8 Biosimilars: what clinicians should know. Blood. 2012 120:5111_5117. doi:10.1182/blood_2012_04_425744. Prepublished online October 23, 2012;

Projects

Memberships

Danish representative in CHMP's Biological Working Party (BWP) since 2002. Vice_Chair for BWP April 2014_Sept 2021 Danish member of the Committeee for Adcvanced Therapies (CAT) since 2015, alternate member 2012_2020 Expert member of CHMP's Biosimilar Medicines Working Party (BMWP) since 2010 Expert member of EMA's PAT Team 2013_2020 Danish representative in CHMP's Ad hoc Influenza Working Party 2003_2020 Member of Expert Working Group in ICH Q12. Regulatory Chair since 2017

Other Relevant Information