Curriculum Vitae

Personal information

Valeria Zoccano

Work experience

1. Employer: Agenzia Italiana del Farmaco
   - Start date: 042017
   - End date:
   - Position: Quality Assessor and GMP Inspector
   - Activities: Quality assessor and Scientific administrator for centralised procedures (first marketing authorisations and variations to the marketing authorisation) and EMA Scientific Advices; Linguistic revision of centralised products information; Inspector of active pharmaceutical ingredients manufacturing sites (national, in third countries, EMA); draft/edit inspection reports. EMA expert for GMP inspections and Quality Assessment since 2011.
   - Country: Italy
2. Employer: Agenzia Italiana del Farmaco
   - Start date: 092008
   - End date: 042017
   - Position: GMP Inspector and Quality Assessor
   - Activities: Inspector of active pharmaceutical ingredients manufacturing sites in Italy and non_UE countries to ensure GMP and dossier compliance in the framework of Italian law; draft/edit inspection reports; follow_up of inspections.
     Scientific evaluation of dossiers submitted with new applications for active pharmaceutical ingredients manufacturing authorizations in Italy in accordance with Italian law, current technical rules and guidelines.
     Scientific/Technical evaluation of modifications to Site Master Files (i.e. new manufacturing areas, revamping, etc) of active pharmaceutical ingredients and medicinal products manufacturing sites, in accordance with Italian law, current technical rules and guidelines .
   - Country: Italy
3. Employer: Agenzia Italiana del Farmaco
   - Start date: 062007
   - End date: 082008
   - Position: Regulatory Assessor – Mutual Recognition and Decentralized Procedures
   - Activities: Evaluation of CTD Module 1 for new marketing authorization applications.
     Contact_point with applicant and other European licensing authorities in order to facilitate the flow of information concerning items on the work programme.
     Evaluation of marketing authorization variation type 1 according to EU regulation EC/1084/2003.
   - Country: Italy
4. Employer: Catalent Italy SpA
   - Start date: 072000
   - End date: 052007
   - Position: Technical Support Manager – R&D area
   - Activities: Ensuring regulatory compliance between CTD and manufacturing instructions and editing of manufacturing instructions.
     Development and formulation of pharmaceutical products and food supplement, in cooperation with QA, QC, Production and Logistic Departments.
     Technical transfer project leader.
     Technical support for manufacturing process modifications and production troubleshooting .
     Process Validation Protocols check and approval.
     Regulatory evaluation and approval of packaging materials, in compliance to Italian law.
     Preparation of food supplement nutritional labels according to current guidelines and Italian law.
   - Country: Italy

Education and training

1. Subject: università La Sapienza
   - Start date: 091990
   - End date: 121999
   - Qualification: master's degree in chemistry
   - Organisation:
   - Country: Italy

Additional information

Publications

Projects

Memberships

Other Relevant Information