Curriculum Vitae

Personal information

Karima Adamo

Work experience

1. Employer: ANSM
   - Start date: 092019
   - End date:
   - Position: Epidemyology Assessor
   - Activities: Evaluation of Post Autorisation Safety Studies Protocols and Reports
   - Country: France
2. Employer: FORTHEIN SASU
   - Start date: 092016
   - End date: 082019
   - Position: President/Manager
   - Activities: find and execute consultancy contracts with CROs.
   - Country: France
3. Employer: Enterome
   - Start date: 102012
   - End date: 122014
   - Position: Director of Clinical development
   - Activities: Clinical studies in the field of inflammatory bowel diseases _ Responsible for the international implementation of projects in accordance with regulations (United States, France, Switzerland, Belgium) _ Responsible for submissions to health authorities and ethics committees _ Responsible for writing clinical operating standards _ Responsible for archiving and maintaining study documents _ Responsible for the selection of service providers _ Responsible for the clinical operations budget _ Responsible for writing study protocols and documents _ Line management
   - Country: France
4. Employer: Guerbet
   - Start date: 112011
   - End date: 092012
   - Position: Clinical Project Manager
   - Activities: Management of international phase III clinical study end activities, pivotal study for obtaining Marketing Authorization in the United States: _ Coordination and management of service providers _ Responsible for the medical review of data, monitoring and resolution of questions relating to study data until the database is frozen _ Review and adaptation of existing documents and development of new information supports intended for the various stakeholders _ Training of investigators, clinical research associates and medical representatives
   - Country: France
5. Employer: Ipsen
   - Start date: 052007
   - End date: 082009
   - Position: Clinical Project manager
   - Activities: International phase III study in endocrinology 13 countries (Europe and Turkey) _ Develop protocol and documents relating to the study _ Feasibility study _ Selection of centers and implementation of investigator contracts _ Selection of service providers, setting up of specifications and negotiation of contracts _ Preparation of submission files to health authorities and ethics committees of the countries involved in the study _ Responses to health authorities and ethics committees _ Establishment of study procedures _ Implementation of the investigators meeting _ Management of therapeutic units _ Management of clinical research associates and validation of on_site data control reports
   - Country: France

Education and training

1. Subject: Central School of Paris
   - Start date: 101993
   - End date: 081998
   - Qualification: PhD Pharmacology/Molecular Biophysics
   - Organisation: Volume regulation in cardiomyocytes
   - Country: France

Additional information

Publications

Projects

Memberships

Other Relevant Information