



Curriculum Vitae

Personal information **Karima Adamo**

Work experience

1. Employer: ANSM
 - Start date: 092019
 - End date:
 - Position: Epidemology 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