EUROPEAN MEDICINES AGENCY
SCIENCE MEDICINES HEALTH

Curriculum Vitae

Personal information

Alice Truong

Work experience

1. Employer: FRS_F.N.R.S, University of Liège
   - Start date: 102011
   - End date: 032016
   - Position: PhD Student
   - Activities: _ Writing articles and scientific projects (English / French) _ PPT oral presentation of scientific data (English / French) _ Bibliographic search _ Development of a thesis project + defense of thesis _ Fundamental research in an oncology laboratory (experimental manipulation + results analysis)
   - Country: Belgium
2. Employer: GSK, GlaxoSmithKline (via Altran Consulting))
   - Start date: 102017
   - End date: 122018
   - Position: Technical Regulatory Writer, US Facilities Submission Team
   - Activities: _ Project management and planning for the Facilities US section of new vaccine production buildings (HAV) _ Writing of "Performance Qualification" (cleaning, autoclave ...) for the FDA _ Reading, summary in English of deviations and "change controls" _ Training at many SOPs (Validation, Qualification, Cleaning, Autoclaves)
   - Country: Belgium

Education and training

1. Subject: University of Liège
   - Start date: 012012
   - End date: 062012
   - Qualification: Laboratory Animal Science certification for Scientists responsible for directing animal experiments (FELASA Cat. C)
   - Organisation:
   - Country: Belgium
2. Subject: University of Liège, University of Namur
   - Start date: 102016
   - End date: 032017
   - Qualification: Certificate in Regulatory Affairs
   - Organisation: _ Regulatory procedures for medicinal products: regulatory life cycle of medicinal products, PIP & scientific advice, CTD marketing authorization, variations, orphan medicines, pre_clinical development, clinical trials, Biotech, ATMPs, hospital exemption, Health Technology Assessment _ Legislation: legal aspects of pre_marketing, marketing and post_marketing of medicines, Belgian/European/American legislation, legal document research tools, regulatory intelligence _ Regulatory of other health products: cosmetic legislation, food/dietary supplements, CE labelling and notified bodies, REACH, organs, blood, tissue and cells, biocompatibility, diagnostic test _ Introduction to GLP, GMP, QA, QC, validation, biosafety and GMO _ GCP and medical devices
   - Country: Belgium
3. Subject: University of Liège
   - Start date: 092006
   - End date: 092011
   - Qualification: Master in Biomedical Sciences
   - Organisation:
   - Country: Belgium
4. Subject: University of Liège
   - Start date: 102011
   - End date: 032016
   - Qualification: PhD in Biomedical Sciences
   - Organisation:
   - Country: Belgium

Additional information

Publications

Projects

Memberships

Other Relevant Information