Curriculum Vitae

Personal information

Joelle Warlin

Work experience

1. Employer: FAMPH
   - Start date: 042008
   - End date:
   - Position: Clinical pharmacokinetics assessor
   - Activities: Assessment of the pharmacokinetics documentation in full and abridged applications for the Belgian Agency for Medicines and Health Products (FAMHP)) and for the European Medicines Agency (EMA). Reviewer of the pharmacokinetics assessments. Assessment of the bioequivalence studies in the generic and hybrid applications. Scientific advices and clinical trial applications. Clinical PK team coordination.
   - Country: Belgium
2. Employer: FAMHP
   - Start date: 092001
   - End date: 032008
   - Position: Bioequivalence assessor
   - Activities: Bioequivalence assessment at FAMHP and EMA.
   - Country: Belgium
3. Employer: University of Louvain department of Pk
   - Start date: 092000
   - End date: 092001
   - Position: Assistant at the University
   - Activities: Scientific collaboration between UCL and Belgian Pharmaceutical Inspectorate on generic medicinal products with the objective to assess their bioequivalence and their quality.
   - Country: Belgium
4. Employer: Public pharmacy Forest, Brussels
   - Start date: 071999
   - End date: 071999
   - Position: Pharmacist
   - Activities: Medicinal products delivery
   - Country: Belgium
5. Employer: Pharmacovigilance department Searle Continental Pharma
   - Start date: 2000
   - End date: 2000
   - Position: Student in Pharmaceutical Industry
   - Activities: Pharmacovigilance activities
   - Country: Belgium
6. Employer: Hospital Pharmacy St Luc UCL
   - Start date: 1998
   - End date: 1998
   - Position: Pharmacist
   - Activities: Medicinal products delivery
   - Country: Belgium

Education and training

1. Subject: UCL Catholic University of Louvain (UCL)
   - Start date: 091995
   - End date: 091999
   - Qualification: Graduate of School of Pharmacy, Catholic University of Louvain (UCL).
   - Organisation: Pharmacokinetics: fundamental principles Pharmacokinetics: Clinical aspects
   - Country: Belgium
2. Subject: ULB_UCL_ULg.
   - Start date: 091999
   - End date: 062000
   - Qualification: Specialisation in Industrial Pharmacy, Inter_university program ULB_UCL_ULg.
   - Organisation: Bioequivalence testing Drug metabolism and PK parameters ADME
   - Country: Belgium
3. Subject: ULB, Brussels
   - Start date: 012002
   - End date: 032002
   - Qualification: Post_graduate programme in Pharmacology and Pharmaceutical Medicine: Clinical trials, Pharmacokinetics and Biostatistics. Pharmed, ULB, Brussels.
   - Organisation: Planning the development of new drugs Organising and monitoring clinical trials Good Clinical Practice, standard operating procedures and audits Ethical and legal issues in drug development Biostatistics Pharmacokinetics: basic concepts and applications Drug Metabolism Clinical Pharmacokinetics and linkage between PK and PD data Methods of drug assay Bioequivalence testing Chronopharmacology
   - Country: Belgium

Additional information

Publications

Generic Drug Registration in the European Union: bioequivalence requirements. Roger Verbeeck and Joelle Warlin.

Book Chapter. Isadore kanfer. Informa.

Projects

Collaboration to the INN prescription project.

Collaboration to the development and to the update of a list of narrow therapeutic index (NTI) drugs on the Belgian market.

Collaboration to the development and to the update of a list of biological products on the Belgian market.

Collaboration to the NTI categorization of anti_epileptics to help minimize the risk relating to formulation switching.

Memberships

Additional assessor at the previous Pharmacokinetics Working Party (EMA)

Member of the methodology ESEC (EMA)

Other Relevant Information