

PERSONAL

Belgian, ° 10.12.1966, widow, 2 children (° 19.01.1998 and 15.10.1999).
Officially listed as Industrial Pharmacist (n° 1320), Responsible Pharmacovigilance (n° P154),
Responsible Scientific Information and Publicity (n° 563).
Active member of BRAS Education Group and Management Board.

EDUCATION

Job specific courses/seminars/symposia BRAS, VAPI, CfPA, LuxControl, BIO, DIA, EDQM, Unizo, ...	1991 -
Specialisation Industrial Pharmacist Universitaire Instelling Antwerpen	1989 - 1991
Pharmaceutical Sciences Universitaire Instelling Antwerpen Rijksuniversitair Centrum Antwerpen	1984 - 1989

EXPERIENCE

6. infARAMA BVBA (www.infarama.be) – Partnership of Infarmed and RA-MAnagement

Managing Director Quality and RA - Industrial Pharmacist **founded in Oct 2008**
consultancy services to leading healthcare companies: edition and follow-up of registration
dossiers, audits (Good Distribution Practices / Good Laboratory Practices / Good Clinical
Practices / ISO 9000-compliance), (administrative) batch release, representation for
pharmacovigilance / scientific information and publicity / medical translations

5. RA-MAnagement BVBA (www.ra-management.be)

Managing Director - Industrial Pharmacist **Apr 2006 – Sep 2008**
consultancy services to leading healthcare companies: edition and follow-up of registration
dossiers, audits (Good Distribution Practices / Good Laboratory Practices / Good Clinical
Practices / ISO 9000-compliance), (administrative) batch release, representation for
pharmacovigilance / scientific information and publicity

4. REGIPHARM group

Regulatory Associate - Responsible Pharmacist **Jan 1995 – Mar 2006**
edition of registration dossiers for (inter)national companies, follow-up of bio-equivalence
studies, central pharmacy for clinical studies, administrative controls, audits (Good
Distribution Practices / Good Laboratory Practices / Good Clinical Practices / ISO 9000-
compliance), pharmacovigilance, scientific information and publicity

3. Belgian Red Cross (CDF)

Responsible Pharmacist

Dec 1993 - Dec 1995

release of the raw materials, the intermediate products and the final products, fixing of product specifications, audits of the QC lab and the production department

Responsible production coagulation factors

Oct 1992 - Nov 1993

management of the production-unit for the manufacturing of coagulation factors

2. I.C.C.E. (International Cleanroom Control Engineering)

Consultant - Industrial Pharmacist

Oct 1991 - Sep 1992

advice and support during the installation of clean rooms for pharmaceutical, micro-electronical and biotechnological industries (design, choice of equipment, follow-up installation, validation, control,)

Most important project (client : Belgian Red Cross) : installation of a new production unit for the manufacturing of biotechnological products (coagulation factors)

1. JANSSEN Pharmaceutica

Manager Registration Reports

Apr 1991 - Sep 1991

co-ordination of the editing of pharmaceutical reports as a part of the (inter)national registration dossiers (product specifications, ...)