Marian COQUEL

Costermansstede 1
B-1850 Grimbergen
tel. 02/270 95 22
fax 02/306 91 81

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 **1991 - …….**
BRAS, VAPI, CfPA, LuxControl, BIO, DIA, EDQM, Unizo, …

Specialisation Industrial Pharmacist **1989 - 1991**
Universitaire Instelling Antwerpen

Pharmaceutical Sciences **1984 - 1989**
Universitaire Instelling Antwerpen
Rijksuniversitair Centum Antwerpen

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, ...)