

PERSONAL INFORMATION

Greet Musch

WORK EXPERIENCE

October 1983–September 1989

Scientific Associate

Free University Brussels (Belgium)

Ph D in Pharmaceutical Sciences : Biomedical HPLC analysis and solid phase extraction

October 1989–April 1997

Director of Quality Development Laboratories JRF

J&J : Janssen Pharmaceutica (Belgium)

Full Responsibilities for all lab activities related to the development of new drugs (clinical trials , stability studies , regulatory applications) for the R&D projects

April 1997–April 2004

senior quality assessor

FPS Health (Belgium)

assessment of scientific data in view of marketing authorisations

Member of CHMP-CVMP QWP at EMA

project management generics , pMDI , HoA Training of Assessors

April 2004–December 2008

Head of R&D department

FPS Health (Belgium)

Responsible for clinical trials , scientific advice and GCP inspections

CTFG member

GCP inspectors group member

January 2009–Present

General Director DG PRE authorisation

Federal Agency for Medicinal and Health products (Belgium)

Scientific advice , clinical trials , first marketing authorisations , veterinary division , homeopathic division , EU Pharmacopea

EDUCATION AND TRAINING

November 1994–November 1995

Middle management

Vlerick (Belgium)

January 2003–November 2003

Public management

KUL (Belgium)

ADDITIONAL INFORMATION

Expertise

Publications

Projects

Memberships

Other Relevant Information