Curriculum vitae

Personal data

Name	: Fokkens
Given names	: Jasper Gerrit
Date of birth	: 23 December 1950
Nationality	: Dutch
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Employment record as consultant

Fokkens Consultancy B.V.

2001 - : *Managing Director*

Main activities:

- defining and implementation of quality systems mainly in the (bio-)pharmaceutical and food industries
- execution of (mock) audits (GMP)
- guiding (bio-)pharmaceutical companies to pass an FDA inspection
- provide services as Qualified Person (EU clinical trial directive EU2001/20/EC)
- interim management
- project management: coaching of project teams; executing projects
- courses for the pharmaceutical and food industry and their suppliers
- supporting management (e.g. strategic choices; re-organizations)
- coaching managers / management

- *2009 – 2010 Interim Director QA/RA for Sanquin Bloodbank*

- *2008 - Qualified Person for (bio-)pharmaceutical companies*

- *2008 – 2009 Guiding the project Interface Development – Manufacturing, Solvay Pharmaceuticals*

- *2008 – 2009 Interim manager Quality Assurance & Safety, Health and Environment Frieslandfoods*
  - managing the QA&SHE department
  - improvement and implementation of the CAPA system

- *2007 – 2008: DSM Biologics*
  - Manager Quality Assurance and Quality Control
  - Qualified Person

- *2007: Millipore/MicroSafe*
  - support and coaching of persons involved in managing their department, including implementation of relevant management tools

- *2004 – 2007: Givaudan (till 5 March 2007 Quest International); flavours and fragrances*
  - set-up and pilot implementation of a global quality system
  - setting up and managing the flavour global compliance department

- *2005: Diatos S.A.*
  - introduction of the Batch Review system from clinical phase I - consecutive batches

- *2004 - 2005: Purac Biomaterials*
  - increasing compliance to CFR 211 level

- *2003 - 2004:Diatos S.A.*

- executing all quality assurance work
- contract reviews

- *2003 - 2004: DSM Biologics*
  - Preparation, both contents and organisation, for pre-approval inspection by FDA
  - Project Manager; project activities were related to the submission of Biologics License Application

- *2003: KLM Cargo*
  - consultancy for the development of the pharmaceutical services package

- *2003: AVEBE*
  - Managing of the management buy-out of Paragon

- *2002 – 2003: Purac*
  - integration coordinator after takeover of Glucona by Purac

- *2002: Campina*
  - advising on a corporate quality system

- *2001 – 2008: Miscellaneous*
  - audits of various (biopharmaceutical, pharmaceutical and food) companies and suppliers
  - various cGMP training courses
  - guidance of project teams
  - coaching of (QA) managers

Employment record

Avebe Glucona

2001 - 2002 : *Managing Director Avebe Glucona*
Avebe Glucona b.v. was a stand alone Business Unit within Avebe, developing and manufacturing mineral salts and excipients for the pharmaceutical and food / neutraceutical markets. Leading Glucona team selling the business in 2002 to Purac biochem (mineral salts) and DMV International (excipients).

DSM, business group DSM Anti-Infectives

1998 – 2001 : *Director Safety, Health and Compliance*

Gist-brocades, Industrial Pharmaceutical Products Division (IPPD)

1998 : *General Manager site Delft ad interim*

1997 – 1998 : *Director Quality Assurance and Regulatory Affairs*

1996 – 1997 : *Technical Sales Service Manager*

Solvay Human Health

1995 – 1996 : *Licensing Director*

Solvay Duphar

1992 – 1995 : *Manager of the Section Chemistry-Pharmacy*

1990 – 1992 : *International Project Manager*

1990 – 1992 : *Member reorganization team*

1989 : *Portfolio manager Solvay Human Health*

1984 – 1990 : *Manager of the Research Group in pharmaceutical development*

1983 – 1985 : *Manager of the development group for liquids*

State University of Utrecht

1978 – 1983 : *Research scientist (department of biopharmaceutics)*
Research in biopharmacy (Ph.D. study); lecturer at all levels in pharmacy, pharmaceutical technology, physical pharmacy and biopharmaceutics.

1976 – 1978 : *Research scientist (part-time, department of pharmaceutical chemistry)*

Education

1978 : Pharmacist, State University of Utrecht

1983 : Ph.D., State University of Utrecht
(Thesis: 'In vitro drug release from non-aqueous suspensions')

Many courses, technical and managerial.

Languages

Dutch, English, German (moderate) and French (moderate)

Administrative functions

1976 – 1983 : Various technical and managerial functions in the Faculty of Pharmacy, State University of Utrecht.

1988 – 1990 : Founder and lecturer of the course Pharmaceutical Technology in Solvay Duphar, a 6 weeks course, academic level.

1991 – 1994 : Chairman of the Anselmus Stimulation Award Committee

1989 – 1999 : Founder and chairman of the Anselmus Foundation.

Miscellaneous

1983 – 1991 : Member of the editorial board of Pharmacy World & Science

1986 – 1990 : copromotor Dr. E.M.G. van Bommel

1998 : film "Coating Defects"

1990 – 1991 : co-author of the course Pharmaceutical Technician

Referent of tens of articles and books for several scientific journals.
Member of various review and evaluation committees for scientific programs.
Invited speaker on symposia and (post-academic) courses.
Tens of international lectures and poster presentations.
Over 30 scientific publications.
Co-editor 'Innovations in Drug Delivery – impact on pharmacotherapy'.

Hobby

Reading, bridge, tennis, solving cryptograms.

jgf, Jan 2011