



Curriculum vitae

Frans van den Berg

Qualifications

- **MBCbB** 2000
University of the Free State, South Africa

Registration

- General Medical Council, UK 2002 – present
Registration No. 6062789: full registration & licence to practise

Teaching experience

- Lecturer (*Safety and tolerability*) 2010 – present
MSc course in Clinical Drug Development,
Barts and the London
- Lecturer (*Ethics*) 2009 – present
CPPD course in Clinical Pharmacology Practice,
Thames Valley University

Other training

- Advanced Life Support Course Oct 2009
Kings College, London
- Eudravigilance and electronic submission of case safety reports May 2012
DIA, EMEA, 7 Westferry Circus, London
- Extended Eudravigilance Medicinal Product Dictionary May 2013
DIA, EMEA, 7 Westferry Circus, London

Professional membership

- Affiliate Membership 2007 – present
Faculty of Pharmaceutical Medicine, London
- AHPPI 2009 – present
(Association for Human Pharmacology in the Pharmaceutical Industry)

Current appointment

- Senior Research physician and Principal Investigator Apr 13 - present
Hammersmith Medicines Research
Cumberland Avenue, London, NW10 7EW

Previous appointments

- Research Physician, Hammersmith Medicines Research Jul 05 – Apr 13
Cumberland Avenue, London NW10 7EW



Curriculum vitae

Frans van den Berg

- Senior House Officer, Care of the Elderly
Orthopaedic Rehabilitation Unit
Brighton General Hospital, Brighton Apr 04 – Jul 05
- Senior House Officer, General Medicine/Acute Medicine
Royal Sussex County Hospital, Brighton Mar 04
- Senior House Officer, Care of the Elderly/General Medicine
Brighton General Hospital, Brighton Feb 04
- Senior House Officer, Gastroenterology/General Medicine
Pinderfields General Hospital, Wakefield Jan 04
- Senior House Officer, Cardiology/General Medicine
Belfast City Hospital, Belfast Feb 03 – Sep 03
- Community Medical Officer, Department of Family Medicine
University of the Free State, South Africa Jan 02 – Dec 02
- Internship
Pelonomi Hospital/National Hospital/Universitas Hospital
Training complex, Bloemfontein, South Africa Jan 01 – Dec 01

Experience in pharmaceutical medicine

- Design and execution of human studies in exploratory drug development, since 2005
- More than 50 completed phase 1 and phase 2 studies
- Principal Investigator in 2 studies
- Co-investigator in 50 studies, including phase 2 studies in patients with systemic lupus erythematosus
- Asthma and allergy procedures:
 - allergen dilutions
 - skin prick tests
 - inhaled allergen & methacholine challenge
- FTIH, PET, EEG, bioequivalence, bioavailability, thorough-QT and micro-dosing studies
- Setting up and running clinical pharmacology studies in accordance with Good Clinical Practice
- Main duties:
 - Advising senior management on feasibility of studies
 - Protocol review: design, selection & stopping criteria, rationale
 - IB review
 - Clinical trial risk assessment



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- Calculation and recalculation of starting dose
- Presentation to Ethics Committees
- Consent and screening of subjects
- Day-to-day running of studies
- Medical management of AEs
- Interim and final safety reporting
- Chairing journal club meetings (critical analysis of published papers)
- Presentations and medical lectures to staff
- Medical writing: ICFs, protocols, study procedures and SOPs
- Running training resuscitation scenarios
- Teaching: mentor to new physicians

A handwritten signature in black ink, appearing to read 'FvdB', written in a cursive, stylized script.

19 JUN 2013