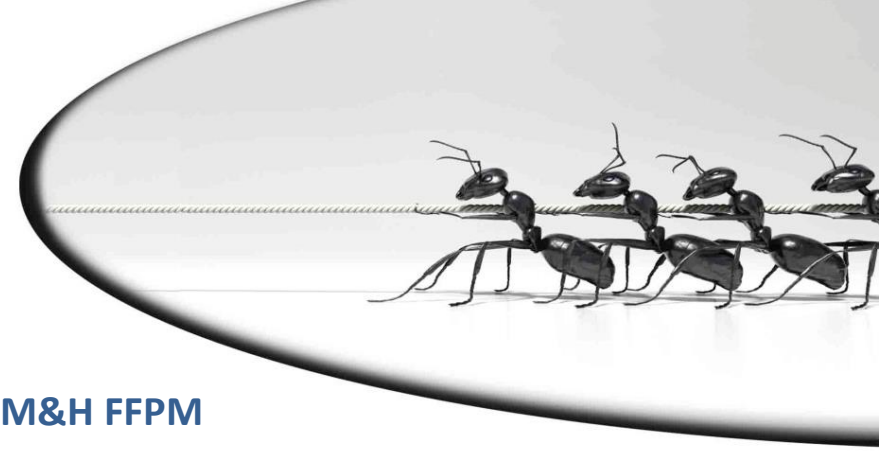
cddi



Grahaem Brown MB FRCP DTM&H FFPM

- Internist
- Clinical dev. strategy
- Clinical operations
- Developed 19 marketed products
- Large Pharma senior roles
- Over 5 years consulting for biotech and pharma

EMPLOYMENT HISTORY

2009 – Current — **Competitive Drug Development International: Managing Partner**

2007 – 2009 — **Grahaem Brown Consulting Ltd: CEO**

2007 – 2010 — **Haemostatix: Non-executive Director**

2004 – 2007 — **UCB: Senior Vice President, Development**

- Responsible for Clinical Pharmacology & Experimental Medicine, Clinical Research, Project Management and Regulatory Affairs
- Led BLA for Cimzia (TNF antibody for Crohn's Disease and Rheumatoid Arthritis) and several sNDAs for Keppra (treatment of epilepsy)

2003 – 2004 — **Celltech: Development Director**

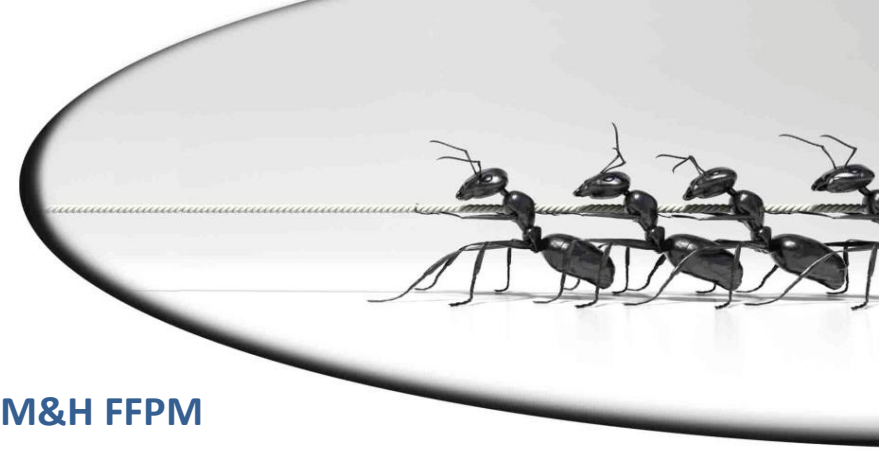
- Responsible for Experimental Medicine, Clinical Development, Pharmacovigilance, Non-Clinical Development, Regulatory Affairs and Project Management

2000 – 2003 — **Pharmacia Corporation: Global Head of Clinical Operations**

- Accountable for global R&D clinical programmes
- Set up global, home-based field monitoring organization, recruiting ~15,000 subjects annually
- Championed successful process re-engineering across Italy, UK and US

1999 – 2000 — **Pharmacia & Upjohn: Vice President Clinical Research, Europe**

- Management of European Clinical Research: Cardiovascular, Ophthalmology and General Medicine Therapeutic areas
- Delivered 5 regulatory submissions: Genotropin (Prader-Willi Syndrome); Detrol (bladder dysfunction); Xalcom (glaucoma); Fragmin (coronary artery disease); Pegvisomant (acromegaly)



Grahaem Brown MB FRCP DTM&H FFPM

1996 – 1999 **Novartis: Head of Clinical Research**

- Global responsibility for Clinical Development, including Therapeutic Areas and Clinical Operations
- Delivered registration dossiers for 6 major NCEs and 11 line extensions

1993 – 1996 **Ciba-Geigy: Head, International Clinical Research**

- Responsible for European Clinical Research, Clinical Pharmacology and Clinical Supplies
- Led the registration programs for 3 major NCEs

1991 – 1993 **Glaxo-Canada: Affiliate Medical Director / SVP R&D**

- Responsible for Regulatory Affairs, Clinical Research, Drug Safety and a corporate Pharmaceutical Development group
- Chaired the International Development Committee for ID & Oncology

1986 – 1991 **Glaxo Group Research: Director of Clinical Research, Infection and Oncology**

- Led a therapeutic area team developing antibiotics, an anti-HIV project, P1/2 Oncology programs and an anti-emetic project
- Led and delivered the filing for Zofran (first 5HT3 antagonist to market)
- Led the early development of Epivir (for HIV)

1982 – 1986 **Glaxo Group Research: Senior Research Physician**

- Managed antibiotic, dermatology and other development projects

1967 – 1986 **Military Medical Service: Consultant in Medicine**

- Internal Medicine, Tropical Medicine, responsibility for Oncology

QUALIFICATIONS

1990	FRCP (London)
1989	FFPM
1982	Specialist Accreditation in Tropical Medicine
1978	Specialist Accreditation in Internal Medicine
1974	DTM&H
1971	MRCP
1967	MB BS