

Dissolution, Bioavailability and Bioequivalence

14 – 15 May 2008, Radisson SAS Beke Hotel, Budapest, Hungary

Get up-to-speed with the latest technologies in dissolution testing and ensure regulatory compliance with your bioavailability (BA) and bioequivalence (BE) studies

EXPERT SPEAKERS including...

Prof Ildiko Csoka, *Institute of Drug Regulatory Affairs, University of Szeged*, Hungary

Margherita Zanol, *Merck Serono*, Italy

Professor Jean-Michel Cardot, *Universite d'Auvergne*, France

Dr Paula Muniz Piniella, *Synthon BV*, The Netherlands

Dr Karl Peeters, *Johnson & Johnson Pharm. R&D*, Belgium

Dr Jochen Scher, *Boehringer Ingelheim Pharma*, Germany

REASONS TO ATTEND:

Manage the relationship between BE studies and formulation

Examine pharmacokinetic behaviour of drugs and its implications to BE

Conduct BA/BE studies to manage scale-up issues in drug development

Gain a regulatory update on biosimilars

Hear what regulators are looking for at GCP & GLP inspections

Learn how to avoid the common pitfalls associated with BE studies

Ensure a successful application for a biowaiver

Benchmark strategies to obtain biowaivers from other European countries & Japan

- 2 full-day workshops:
 - Successfully planning and designing a bioequivalence (BE) study
 - Introduction to IVIVC and its practical application to NCE & generic development

Download THE BROCHURE for free:

www.informa-ls.com/beba

REGISTER today:

By phone +44(0)20 7017 7481

By fax +44(0)20 7017 7823

By email registrations@informa-ls.com

Online www.informa-ls.com/beba

Register before 28 March 2008 and SAVE up to £400! Promotion Code: CQ5062BA