5th Annual Drug Delivery Systems Conference

Assessing Innovations in Dryg Delivery Technologies and Evaluating Business Models

9th-11th February 2010, BSG Conference Centre, London, UK

Key Speakers:

• Dr. Sven Stegemann, Director, Pfizer

- Norbert Rasenack, Fellow Inhalation Development, Novartis
- Dr. Jan Möschwitzer, Head Early Pharmaceutical Development, Solvay
- Dr. Oliver Steinbach, Department Head, Philips Research Laboratories
- Joel Richard, Senior Director, Head of Drug Product Development, Ipsen
- Dr. Duncan F Rogers, Reader, National Heart and Lung Institute, Imperial College London
- Dr. Dejan Djuric, R&D Project Management Excipients, BASF
- Dr. Cheryl Barton, Managing Director, Pharmavision
- Gerry Kamstra, Partner, IP & Life Sciences, Bird and Bird
- Prof. Marc Brown, CSO, MedPharm
- Craig Thomson, Attorney, Murgitroyd & Company
- Dr. Giuseppe Battaglia, Lecturer in Bionanotechnology, University of Sheffield
- Dr. Piet Christiaens, Scientific Director, Toxikon Europe
- Dr Pierandrea Esposito, Associate Director, SiTec PharmaBio

Conference Agenda

Day One, Wednesday 10th February 2010

09:30 Registration and refreshments

10:00 Opening address from the chair

10:10 Analysing key trends in the field of drug delivery

- · Impact of emerging drug delivery technologies on the pharmaceutical industry
- · Evaluating the latest market developments

10:50 Innovations in trans (dermal) drug delivery

- Issues of drug delivery to the skin
- Past successes and failures (trans) dermal drug delivery
- Innovations in active and passive methods of (trans) dermal drug delivery
- · Gene delivery; the final frontier

Dr Marc Brown Chief Technology Officer MedPharm

11:30 Morning refreshments

11:50 Drug delivery systems for oncology products

- Cancer market context and expectations
- Key challenges for delivery of oncology products
- · Oral chemotherapy: opportunities, challenges and delivery solutions
- Drug permeation and bioavailability enhancement
- Delivery strategies based on nanosystems and lipid-based formulations
- I.v. oncology products: challenges and delivery solutions
- Drug solubilisation
- Modification of biodistribution and toxycology profile of drugs
- Delivery strategies based on nanosystems for small molecules and oligonucleotides

· Subcutaneous oncology products: challenges and delivery solutions

- Sustained-release formulations of peptides
- Perspectives for new s.c. formulations of proteins and mAbs

Joel Richard

Senior Director, Head of Drug Product Development Ipsen

12:30 Particle engineering as contributor to a successful pharmaceutical drug product development - challenges, opportunities, trends

- Poorly soluble
- Inhalation delivery systems
- · Technologies: opportunities and drawbacks
- Norbert Rasenack Fellow Inhalation Development Novartis

13:10 Networking lunch

14:30 Stem cell-based therapeutic delivery: challenges & opportunities

• Ethical debates and funding, immune rejection, cancerous potential and issues surrounding the manufacture of stable, pure cell therapeutics

• Delivery of stem cells for CNS disorders

• Autologous products, derived from a patient's own tissue, will be the first to reach the market

Dr Cheryl L Barton Managing Director Pharmavision

Dr Sara Sleigh Senior Associate Pharmavision

15:10 Gatecrashing the cell

- Why it is important to deliver within cells' cytosol
- Biomimetic strategies for cytosolic delivery
- Delivery of DNA, siRNA, ODN, mRNA and their therapeutic implications
- Delivery of contrast agent and how cytosolic delivery can revolutionise diagnostics
- New therapeutic approaches based on cytosolic delivery

Dr Giuseppe Battaglia Biomaterials and Tissue Engineering Group University of Sheffield

15:50 Afternoon refreshments

16:10 Challenges and opportunities in the delivery of small molecules and of biotherapeutics

- Drug delivery role in new drugs development and in lifecycle management
- Trends and changes in delivery of therapeutic biomolecules
- Current issues in small molecules delivery
- New formulation technologies or molecular engineeering?

Dr. Pierandrea Esposito Associate Director SiTec PharmaBio

16:50 Future therapeutic targets for respiratory disease

- Incidence and economic burden of asthma and chronic obstructive pulmonary disease (COPD)
- Overview of pathophysiology of asthma and COPD
- Overview of strategies for target identification and novel drug development

•	Options	for	improving	existing	pharmacotherapy
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- · Hypothesis-driven novel targets
- Natural products, 'phytoceuticals' and non-synthetics
- Assessment of clinical trial progression of new drugs

Dr. Duncan F Rogers Reader, National Heart and Lung Institute Imperial College London

17:40 Closing remarks from the chair

17:30 Networking Drinks Take your discussions further and build new relationships in a relaxed and informal setting

Day two, Thursday 11th February 2010

09:30 Registration and refreshments

10:00 Opening address from the chair

10:10 Design drug delivery for individual patient populations

· Changing patient population and its relevance for therapy

- · Individualisation of drug delivery in development and manufacturing
- Drug delivery for effective medication management
- Drug delivery aligning technology with patient needs

Dr. Sven Stegemann Director Pfizer

10:50 Image-guided ultrasound- triggered drug delivery

Use of drug loaded, temperature (liposomes) and pressure sensitive (microbubbles) systems for ultrasound triggered drug delivery
Tracking, monitoring and quantifying drug release with ultrasound or MRI through the incorporation of imaging lables in drug carriers

· Formulations and applications for different drug formats: small molecules, biologics, nucleic-acid based therapeutics

· Examples from different disease arease.g. oncology, cardiology, metabolic diseases

Dr. Oliver Steinbach Department Head Philips Research Laboratories

11:30 Morning refreshments

11:50 Drug nanocrystals prepared by using different particle size reduction methods

Overview of particles size reduction technologies used in academia and industry, bottom-up, top-down and combinative processes
When can API nanonisation helps to improve the bioavailability of poorly soluble compounds

- Aspects to be considered during development: stabilisation, analysis and solidification
- Aspects to be considered during development, stabilisation, analysis and solid
 The use of nanonised API at different stages of development
- Case study

Dr. Jan Möschwitzer Head Early Pharmaceutical Development, Solvay

12:30 Regulatory Requirements for Biocompatibility Testing of Drug Delivery Devices

• Classification of a Drug Delivery Device: Is the product a Drug or a Device?

- General Considerations regarding Biocompatibility testing
- Typical Biocompatibility Tests to be considered
- Testing/Documentation requirements for the submission of a Medical Device
- Testing/Documentation requirements for the submission of a Medicinal Product/Pharmaceutical Container

Dr. Piet Christiaens Scientific Director Toxikon Europe

13:10 Networking lunch

14:30 Drug synthesis and delivery IP due diligence

• Importance of IP protecting manufacturing process and drug delivery technology in late stage investments

• Typical issues around manufacturing process patents

• Typical issues around drug delivery patents

Subsistence of Supplementary Protection Certificates

Regulatory Data Protection and Orphan Drug

• Protection in late stage investments

Gerry Kamstra Partner, IP & Life Sciences Bird and Bird

15:10 Ensuring collaborations enhance rather than stifle exploitation of new technology

• Problems associated with not addressing ownership of IP early in collaboration

• How the law can help to avoid these problems

• Is there any reason to share IP?

• Examination of collaboration case studies

Craig Thomson Attorney Murgitroyd & Company

15:50 Afternoon refreshments

16:10 Polymeric Solubilizer for Hot Melt Extrusion

Characterization of a bifunctional polymer which is capable of solubilization and of forming solid solutions with poorly soluble drugs
Manufacturing of solid solutions

• Applications for polymeric solubilizers

· Solubility and Bioavailability enhancement

Dr. Dejan Djuric R&D Project Management Excipients BASF

16:50 Presentation to be announced

17:30 Chair's closing remarks

17:40 End of Conference

-Pricing-

Standard Rate for 2 day Conference-£1299 + VAT:£194.85 = Total:£1493.85

Academic & Health Care Organisations Rate for 2 day conference-£899 + VAT:£134.85 = Total:£1033.85

- If you would like to send more than 1 person then take advantage of the 3 for 2 offer. Send 3 people but only pay for 2.

- How to book -

Booking is simple! All you need to register is contact Bijal. You can either call me on TEL:+44(0)2075499946 or email:bijal.patel@ygpharma.com

Please feel free to contact me should you have any questions, I look forward to hearing from you.

Kind Regards, Bijal Patel