



# FINAL PROGRAMME

## *Pharmaceutical Sciences World Congress*

3rd World Congress of the Board  
of Pharmaceutical Sciences of FIP  
PSWC2007/PharmSciFair Exhibition

April 22-25 • 2007 • RAI Congress Centre • Amsterdam • The Netherlands

***Optimising Drug Therapy: An Imperative for World Health***



**abstracts CD  
in your  
congress bag**

**satellites  
pages 32-37  
exhibition  
separate  
brochure**

**career  
centre  
page 21**

### **Co-Sponsored by:**

European Federation for Pharmaceutical Sciences (EUFEPS)  
American Association of Pharmaceutical Scientists (AAPS)  
Association de Pharmacie Galénique Industrielle (APGI)  
Academy of Pharmaceutical Sciences of Great Britain (APSGB)  
Academy of Pharmaceutical Science and Technology, Japan (APSTJ)  
Australasian Pharmaceutical Science Association (APSA)  
Controlled Release Society (CRS)  
Pharmaceutical Society of Japan (PSJ)  
Spanish Society of Pharmaceutics and Pharmaceutical Technology (SSPPT)

### **Supported by:**

American Society of Health-System Pharmacists (ASHP)  
Belgian Society of Pharmaceutical Sciences (BSPS)  
Canadian Society for Pharmaceutical Sciences (CSPS)  
Dutch Federation for Innovative Pharmaceutical Research (FIGON)  
Dutch Royal Pharmaceutical Society (KNMP)  
European Association of Hospital Pharmacists (EAHP)  
European Society of Clinical Pharmacy (ESCP)  
Federación Farmacéutica Sudamericana (FEFAS)  
International Society for Pharmacoepidemiology (ISPE)  
International Society for the Study of Xenobiotics (ISSX)



**[www.fip.org/PSWC](http://www.fip.org/PSWC)**

# Co-sponsoring Organisations



EUROPEAN FEDERATION  
FOR PHARMACEUTICAL  
SCIENCES

**European Federation for  
Pharmaceutical Sciences (EUFPS)**



**aaps**  
American Association of  
Pharmaceutical Scientists

**American Association of  
Pharmaceutical Scientists (AAPS)**



**Association de Pharmacie  
Galénique Industrielle (APGI)**



apsgb.org

**Academy of Pharmaceutical  
Sciences of Great Britain (APSGB)**



The Academy of  
Pharmaceutical Science  
and Technology, Japan

**Academy of Pharmaceutical  
Science and Technology, Japan  
(APSTJ)**



Australasian Pharmaceutical Science Association

**Australasian Pharmaceutical  
Science Association (APSA)**



**Controlled Release Society (CRS)**



**Pharmaceutical Society of Japan  
(PSJ)**



**Spanish Society of Pharmaceutics  
and Pharmaceutical Technology  
(SSPPT)**

# Supporting Organisations



American Society of  
Health-System Pharmacists

**American Society of Health-  
System Pharmacists (ASHP)**



**Belgian Society of Pharmaceutical  
Sciences (BSPS)**



**Canadian Society for  
Pharmaceutical Sciences (CSPS)**



Federatie voor Innovatief Geneesmiddel  
Onderzoek Nederland  
Netherlands Federation for Innovative Drug Research

**Federation  
for Innovative Drug  
Research (FIGON)**



**KNMP**

**Dutch Royal Pharmaceutical  
Society (KNMP)**



**European Association of  
Hospital Pharmacists (EAHP)**



European Society of Clinical Pharmacy

**European Society of  
Clinical Pharmacy (ESCP)**



**Federación Farmacéutica  
Sudamericana (FEFAS)**



**International Society for  
Pharmacoepidemiology (ISPE)**



**International Society for the Study  
of Xenobiotics (ISSX)**

# Dear Colleague

A warm welcome to the Third World Congress of the Board of Pharmaceutical Sciences of FIP (Fédération Internationale Pharmaceutique or International Pharmaceutical Federation).

The new millennium in 2000 saw the staging of the First Pharmaceutical Sciences World Congress (PSWC2000) in San Francisco, bringing together 2,500 pharmaceutical scientists from around the world. The Second Congress, also with a truly global span, was held in the Spring of 2004 in Kyoto, Japan (PSWC2004), and attracted even more participants. The **Third** Congress is now to convene in Amsterdam (PSWC2007) with the theme of:

## Optimising Drug Therapy: An Imperative for World Health

The program committee, chaired by renowned pharmaceutical scientists, has ensured excellence in scientific quality and a high visibility for the conference. Like the previous two conferences, PSWC2007 will cover a broad spectrum of topics from basic to applied and clinical sciences, addressing timely issues of great importance to drug discovery, development, regulation, and medication management. It will also feature interactive round table discussions, poster sessions, a career centre and an exhibition, and will devote significant attention to young pharmaceutical scientists through poster and podium presentations as well as a pre-conference meeting for graduate students and post-doctoral fellows.

PSWC2007 is co-sponsored by many of the world's leading pharmaceutical science and education organisations and members of the FIP, including the European Federation for Pharmaceutical Sciences, Eufeps, the American Association of Pharmaceutical Scientists, AAPS, the Australasian Pharmaceutical Science Association, APSA, the Academy of Pharmaceutical Sciences of Great Britain, APSGB, the Association de Pharmacie Galénique Industrielle, APGI, the Academy of Pharmaceutical Sciences and Technology, Japan, APSTJ, the Controlled Release Society, CRS, the Pharmaceutical Society of Japan, PSJ, and the Spanish Society of Pharmaceutics and Pharmaceutical Technology (SSPPT). In addition, an impressive number of other organisations have pledged their active (financial) support, and they are listed under supporting or sponsoring organisations elsewhere in the programme.

Over 1250 abstracts have been submitted by registrants from 70 countries. In total, PSWC2007 will offer close to 1500 presentations!

Apart from the main 36 symposia, 4 satellite meetings have been organised under the aegis of PSWC2007. These are:

Two pre-satellite meetings:

- As already mentioned, a meeting especially for **Ph.D. students and Postdocs** (co-sponsored by a number of organisations)
- A workshop on **Pharmacy Curriculum Development** (organised by Utrecht University)

Two post-satellite meetings:

- A meeting on **Monoclonal Antibodies** (co-sponsored by Eufeps/EAPB/FIP/AAPS)
- A symposium on **Microdialysis** (Leiden University/FIP/Eufeps)

And, finally,

- **The Mid Year Meeting 2007 of ISPE**, the International Society for Pharmacoeconomics, which partly overlaps with PSWC2007.

Everything is set for a highly exciting event and an important forum for the global community of pharmaceutical scientists!

The co-chairs forming the PSWC2007 Organising Committee, the Scientific Advisory Board and the leadership of FIP's Board of Pharmaceutical Sciences welcome you to this Third Pharmaceutical Sciences World Congress (PSWC2007) in Amsterdam, one of the most colourful cities in the world!

On behalf of the PSWC2007 Co-chairs of the Organising Committee,



Daan J. A. Crommelin, Ph.D.  
Chair of the Organising Committee

## Your Hosts

**International Pharmaceutical  
Federation (FIP)  
Board of Pharmaceutical  
Sciences (BPS)**



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## Organising Committee Leadership

### Chair

Daan J.A. Crommelin (The Netherlands)

### Secretary

Vinod Shah (USA)

### Programme Co-chairs

Geoffrey T. Tucker (United Kingdom)  
Tomohiko Ohwada (Japan)  
Wolfgang Sadee (USA)

### Fund-Raising Co-chairs

Theo Dingermann (Germany)  
Dick Gourley (USA)  
Mitsuru Hashida (Japan)

### Publicity & Hospitality Co-Chairs

Dominique Duchêne (France)  
Jun Haginaka (Japan)  
Mario Rocci (USA)

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Douwe D. Breimer (The Netherlands)  
Kamal Midha (Canada)  
Tsuneji Nagai (Japan)  
Malcolm Rowland (United Kingdom)  
Yuichi Sugiyama (Japan)

## Local Committee

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Bert Leufkens (The Netherlands)  
Ben Westerink (The Netherlands)

## Organisation

Ton Hoek (The Netherlands)  
Gretie R. v.d. Baumen (The Netherlands)  
Hans H. Linden (Sweden)

### Exhibition

Health Links  
Chris Hanney (United Kingdom)  
Email [channey@health-links.co.uk](mailto:channey@health-links.co.uk)

### Congress Registration & Abstract handling

NewBrooklyn  
Barbara de Jong (The Netherlands)  
Email [registration@newbrooklyn.nl](mailto:registration@newbrooklyn.nl)

### Accommodation

RAI Hotel Service (The Netherlands)  
Email [Hotelservice@rai.nl](mailto:Hotelservice@rai.nl)

### Secretariat after the Congress

FIP Congress & Conferences  
Andries Bickerweg 5  
P.O. Box 84200  
2508 AE The Hague (The Netherlands)  
Email [pswc@fip.org](mailto:pswc@fip.org)

### Congress Venue

Amsterdam RAI  
P.O. Box 77777  
1070 MS Amsterdam  
Europaplein  
NL-1078 GZ Amsterdam (The Netherlands)



# Endorsing Organisation

Patron Category



# Sponsoring Organisations

Patron Category

## FPMAJ

The Federation of Pharmaceutical Manufacturers' Associations of Japan

Platinum Category



# Sponsoring Organisations

## Gold Category



## Silver Category



A Global Biopharma Leader



## Bronze Category



## Contributor Category





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Amgen	J. Haginaka
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L. Benet	M. Hashida
V. Shah	M. Rowland

## PSWC2007 Registration Support for PhD Students and Young Postdocs

Belgian Society for Pharmaceutical Sciences (BSPS)  
Dutch Federation of Pharmaceutical Sciences (NVFW)  
German Pharmaceutical Society (DPhG)



# Opening Session

**Pharmaceutical Sciences World Congress (PSWC)**  
3rd World Congress of the Board of Pharmaceutical Sciences of FIP

RAI Auditorium, Sunday, April 22, 2007, from 15:00-17:30

## FIP Fanfare

### Welcome Address on behalf of PSWC2007

*Dr. Daan Crommelin, PSWC 2007 Chair*

### Introduction and Remarks of the Minister of Education, Culture and Science in The Netherlands

*Dr. Ronald Plasterk*

### Welcome Address on behalf of the International Pharmaceutical Federation (FIP)

*Dr. Kamal K. Midha, FIP President*

### Distinguished Lecture The Pharmaceutical Sciences – Getting our Act together

*Dr. Geoffrey T. Tucker, Chair Scientific Programme Committee, Sheffield, United Kingdom*

### Introduction and Presentations of the PSWC Research Achievement Awards

*Dr. Hiroshi Terada – Dr. Elias Fattal – Dr. Kathleen M. Giacomini*

## Entertainment

### Distinguished Lecture The Future of the Global Framework of Regulatory Sciences

*Dr. Thomas Lönngren, EMEA, Executive Director, London, United Kingdom*

### Introduction and Presentations of the PSWC Research Achievement Awards

*Dr. Ole Jannik Bjerrum – Dr. Hans Junginger – Dr. Kevin Shakesheff – Dr. William Charman*

### Distinguished Lecture Life and Times in the Pharmaceutical Industry

*Dr. David Nicholson, Oss, The Netherlands*

### Introduction and Presentations of the PSWC Research Achievement Awards

*Dr. Vladimir Torchilin – Dr. Alfonso Dominguez-Gil Hurlé – Dr. Yuichi Sugiyama*

### Remarks on behalf of the Awardees

*Dr. Yuichi Sugiyama*

### Closure and Reception



Dr. Geoffrey T. Tucker



Dr. Thomas Lönngren



Dr. David Nicholson



For extended curricula vitae of the awardees see pages 28-31.

Day	Time	Activity	Place
<b>Sunday, April 22, 2007</b>	10:00-17:00	Registration	Forum Foyer
	10:00-17:00	Press/Speakers Room	Room M 1st floor Forum Part
	09:00-18:00	Career Centre	Rooms F&G 1st floor Auditorium Part
	09:00-18:00	Career Centre	Commission 1 ground floor Auditorium Part
	15:00-17:30	Opening Session	Main Auditorium
	17:30-19:00	Welcome Reception	Hall 10 st floor Forum Part
<b>Monday, April 23, 2007</b>	07:30-08:30	Hang up Posters	Hall 10 st floor Forum Part
	08:00-17:00	Registration	Forum Foyer
	09:00-17:00	Exhibition & Posters	Hall 10 st floor Forum Part
	09:00-17:00	Press/Speakers Room	Room M 1st floor Forum Part
	09:00-18:00	Career Centre	Rooms F&G 1st floor Auditorium Part
	09:00-18:00	Career Centre	Commission 1 ground floor Auditorium Part
	11:15-12:15	Authors at Poster Board	Hall 10 st floor Forum Part
	17:00-18:00	Take down Posters	Hall 10 st floor Forum Part
<b>Tuesday, April 24, 2007</b>	18:00-19:00	Put up New Poster Numbers	Hall 10 st floor Forum Part
	07:30-08:30	Hang up Posters	Hall 10 st floor Forum Part
	08:00-17:00	Registration	Forum Foyer
	09:00-17:00	Exhibition & Posters	Hall 10 st floor Forum Part
	09:00-17:00	Press/Speakers Room	Room M 1st floor Forum Part
	09:00-18:00	Career Centre	Rooms F&G 1st floor Auditorium Part
	09:00-18:00	Career Centre	Commission 1 ground floor Auditorium Part
	11:15-12:15	Authors at Poster Board	Hall 10 st floor Forum Part
<b>Wednesday, April 25, 2007</b>	17:00-18:00	Take down Posters	Hall 10 st floor Forum Part
	18:00-19:00	Put up New Poster Numbers	Hall 10 st floor Forum Part
	07:30-08:30	Hang up Posters	Hall 10 st floor Forum Part
	08:00-14:30	Registration	Forum Foyer
	09:00-14:30	Exhibition & Posters	Hall 10 st floor Forum Part
	09:00-14:30	Press/Speakers Room	Room M 1st floor Forum Part
	09:00-18:00	Career Centre	Rooms F&G 1st floor Auditorium Part
	09:00-18:00	Career Centre	Commission 1 ground floor Auditorium Part
	11:15-12:15	Authors at Poster Board	Hall 10 st floor Forum Part
	14:00-14:30	Take down Posters	Hall 10 st floor Forum Part

## Programme Time Schedule

### 6 Parallel Symposia & Events

Speaker Presentation	08:30 – 09:05
Speaker Presentation	09:05 – 09:40
Break 09:40 – 10:10	
Speaker Presentation	10:10 – 10:45
Podium Presentations	10:45 – 11:15
Lunch Poster Session	
Keynote Presentations	12:15 – 13:00
Round Table Discussion	13:10 – 14:10
Speaker Presentation	14:15 – 14:50
Speaker Presentation	14:50 – 15:25
Speaker Presentation	15:25 – 16:00
Podium Presentations	16:00 – 16:30
Break 16:30 – 16:45	
EUFEPS Afternoon Sessions	16:45 – 18:15



## Congress Reception


The Welcome Reception is planned on Sunday, April 22, 2007, right after the Opening Session together with the Opening of the PSWC2007/PharmSciFair Exhibition in Hall 10 of the RAI Convention and Exhibition Centre for all participants and accompanying persons.

## Congress Dinner

The Congress Dinner on Tuesday, April 24, 2007, will take place at the famous Wintergarden of the Krasnapolsky Hotel, Dam 9, in the Centre of Amsterdam: 20:00-22:30 hours.


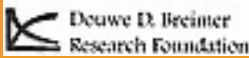
Tickets (Euro 90 including VAT) are limited in number (first-come, first-served). For tickets contact the Registration Desk.

# Programme at a Glance

Scientific Programme Overview • April 22-25 • 2007				
Sunday	Monday	Tuesday	Wednesday	
Registration	<b>Parallel Symposia</b> <ul style="list-style-type: none"><li>■ The era of personalised health care: Impact on drug discovery and development?</li><li>■ Is gene/protein delivery delivering?</li><li>■ Can ADME and PK be predicted from <i>in silico/in vitro</i> data?</li><li>■ Metabolomics: What are the opportunities for biomarker discovery?</li><li>■ What benefits does Process Analytical Technology (PAT) bring to the design and assurance of product quality?</li><li>■ What have we learnt from recent safety cases for new drug development?</li></ul>	<b>Parallel Symposia</b> <ul style="list-style-type: none"><li>■ Druggability: A concept that will fill the pipeline?</li><li>■ Drug targeting: How successful are we?</li><li>■ What is the state-of-the-science in receptor site modelling?</li><li>■ What's new in methods of measuring human drug response?</li><li>■ <i>In silico</i> product development from molecule to man: Dream or reality?</li><li>■ Nutraceuticals: Are new methods of evaluating risk/benefit required?</li></ul>	<b>Parallel Symposia</b> <ul style="list-style-type: none"><li>■ How to manage drug therapy at the extremes of age?</li><li>■ How can nanotechnology and materials science solve drug delivery problems?</li><li>■ How important is genetic and physiological variability in drug transporters?</li><li>■ Systems biology: A driver of drug discovery and development?</li><li>■ How to engineer desired particle properties for drug delivery?</li><li>■ Off-label use of medicines: Abuse or a vehicle for innovation?</li></ul>	
	Posters & Exhibition & Lunch			
Opening Session	<b>Keynote Presentations</b> <ul style="list-style-type: none"><li>■ What are the main challenges for world health? What should be done now and in the future?</li><li>■ Transforming 'art' into 'science' in dosage form design – achievements and challenges</li></ul>	<b>Keynote Presentations</b> <ul style="list-style-type: none"><li>■ Are we meeting the challenges of resistance to anti-infective drugs and of newly emerging infectious diseases?</li><li>■ Will novel approaches to the treatment of cardiovascular disease prove highly effective?</li></ul>	<b>Keynote Presentations</b> <ul style="list-style-type: none"><li>■ Drug resistance in cancer chemotherapy</li><li>■ Recent progress in prion biology</li></ul>	
	<b>Round Table Discussions</b> <ul style="list-style-type: none"><li>■ Science or business as the driver of new drug development?</li><li>■ Does regulation help to 'innovate' or 'stagnate' drug development?</li><li>■ How can the bioavailability of poorly absorbed compounds be enhanced?</li></ul>	<b>Round Table Discussions</b> <ul style="list-style-type: none"><li>■ Translational science: A solution to the productivity gap?</li><li>■ Can microdosing accelerate drug development?</li><li>■ When is a human bioequivalence study not needed?</li></ul>	<b>Round Table Discussions</b> <ul style="list-style-type: none"><li>■ What is the value of observational data post-marketing to assess safety and efficacy?</li><li>■ Life-style drugs: A new burden to the health system?</li><li>■ Is there a consensus on guidelines for the evaluation of biosimilars?</li></ul>	
	<b>Parallel Symposia</b> <ul style="list-style-type: none"><li>■ Pharmacogenetics at the bedside?</li><li>■ What will be the impact of cell-based therapy?</li><li>■ Simulation and modelling in drug development improves decisions, saving time and money?</li><li>■ Dirty vs. selective drugs in the CNS?</li><li>■ How will developments in chemical methods sustain pharmaceutical industrial development?</li><li>■ How effective is the globalisation/harmonisation of pharmacovigilance?</li></ul>	<b>Parallel Symposia</b> <ul style="list-style-type: none"><li>■ Drug-drug interactions: Avoid or understand?</li><li>■ What is the future of smart, feedback, on-demand drug delivery systems?</li><li>■ How are disease and PK-PD connected?</li><li>■ Have omics technologies improved the chance for successful drug development?</li><li>■ Are pharmaceutical manufacturing technologies in stagnation?</li><li>■ Are we using the right outcome measures to ascertain patient benefit from drug therapy?</li></ul>	<b>Parallel Symposia</b> <ul style="list-style-type: none"><li>■ Is the patient taking the tablets?</li><li>■ Developing biotech products: What are the challenges and solutions?</li><li>■ Control of intracellular pharmacokinetics: Advantages for drug therapy?</li><li>■ Molecular targeting in cancer chemotherapy?</li><li>■ Miniaturisation in analytical methods: Is small always beautiful?</li><li>■ Counterfeiting of medicines: Detection and prevention?</li></ul>	
	<b>Round Table Discussion</b> <ul style="list-style-type: none"><li>■ Proactive risk management (plans): Where are we?</li></ul>			
	Short Break			
	<b>EUFEPS Afternoon Sessions</b> <ul style="list-style-type: none"><li>■ Strategic, innovative and critical drug research initiatives: One year later</li><li>■ Pharmaceutical sciences research training and education: Needs and supply</li><li>■ The EU Microdosing AMS Partnership Programme (EUMAPP)</li><li>■ The European Pharma Sciences Leadership Forum (EuPSLF)</li><li>■ Pharmaceutical sciences <i>in silico</i> learning systems: Value and availability</li></ul>	<b>EUFEPS Afternoon Sessions</b> <ul style="list-style-type: none"><li>■ European drug development centres and European growth areas</li><li>■ Reformulation of old drugs: Life cycle management</li><li>■ Vaccine delivery</li><li>■ PharmacoGenetics &amp; Pharmacogenomics Workshop: Outcomes and plans</li><li>■ How to start up a new company?</li><li>■ Drug product quality after new legislation</li></ul>		
Exhibition Opening and Welcome Reception		Congress Dinner		



## SCIENTIFIC PROGR

	<b>Parallel Symposium</b> <b>Room: Auditorium</b> The era of personalised health care: Impact on drug discovery and development?  Sponsoring Organisation	<b>Parallel Symposium</b> <b>Room: Forum</b> Is gene/protein delivery delivering?  Sponsoring Organisation	<b>Parallel Symposium</b> <b>Room: N-O</b> Can ADME and PK be predicted from <i>in silico</i> / <i>in vitro</i> data?  Sponsoring Organisation
	Co-chairs M. E. Brewster, Beerse, Belgium W. Sadee, Columbus, OH USA Y. Sugiyama, Tokyo, Japan	Co-chairs H. Harashima, Sapporo, Japan W. Hennink, Utrecht, The Netherlands	Co-chairs S. Pang, Toronto, Canada H. van de Waaterbeemd, Alderly Park, United Kingdom
08:30	Evaluating molecular genetic factors in drug response: Search for drug targets and biomarkers W. Sadee, Columbus, OH USA	A multi-functional envelope type nano device as a non-viral gene delivery system H. Harashima, Sapporo, Japan	Translation of <i>in vitro</i> data to the whole organ S. Pang, Toronto, Canada
09:05	Predicting drug disposition and response in individual patients: Role of drug transporters for influx and efflux processes Y. Sugiyama, Tokyo, Japan	Cellular mechanisms of non-viral DNA delivery A. Urtti, Helsinki, Finland	<i>In vitro</i> – <i>in vivo</i> extrapolation: Best use of known-knowns to discover unknown-unknowns A. Rostami-Hodjegan, Sheffield, United Kingdom
09:40	<b>Coffee Break</b>		
10:10	Biomarkers in drug development: How do they influence clinical practice? L.J. Lesko, Silver Spring, MD USA	Intravenous siRNA for silencing target genes in solid tumor L. Huang, Chapel Hill, NC USA	A computational systems biology approach to ADME/Tox S. Ekins, New York, NY USA
10:45	Pharmacogenomic analysis reveals determinants of sensitivity and resistance to geldanamycin analogues: Role of membrane transporters Y. Huang, Pomona, USA (MO-S01-1)	Silencing of SOCS genes in cancer cells for effective interferon cancer therapy: Enhancement of antitumor activity of interferons by RNAi-mediated silencing of SOCS gene expression Y. Takahashi, Kyoto, Japan (MO-S02-1)	Assessment of computational and <i>in vitro</i> methods as predictors of oral drug absorption M. Yliperttula, Helsinki, Finland (MO-S03-1)
11:00	Pharmacogenetics: From research results to practical guidelines L. Grandia, The Hague, The Netherlands (MO-S01-2)	Gene expression and silencing for improved islet transplantation R. Mahato, Memphis, USA (MO-S02-2)	Volume of distribution predictions: understanding the processes T. Rodgers, Manchester, United Kingdom (MO-S03-2)
11:15	<b>Posters Lunch Break</b>		
	<b>Keynote Presentation</b> <b>Room: Auditorium</b> Chair H. Leufkens, Utrecht, The Netherlands	<b>Keynote Presentation</b> <b>Room: Forum</b> Chair M. Hashida, Kyoto, Japan	
12:15	What are the main challenges for world health? What should be done now and in the future? R. Laing, Geneva, Switzerland (KLM-1)	Transforming 'art' into 'science' in dosage form design – achievements and challenges P. York, Bradford, United Kingdom (KLM-2)	
	<b>Round Table Discussion</b> <b>Room: L</b>	<b>Round Table Discussion</b> <b>Room: A</b>	<b>Round Table Discussion</b> <b>Room: C-D</b>
13:10	Science or business as the driver of new drug development? Convenor L.Z. Benet, San Francisco, CA USA	Does regulation help to 'innovate' or 'stagnate' drug development? Convenor L.J. Lesko, Silver Spring, MD USA	How can the bioavailability of poorly absorbed compounds be enhanced? Convenor C.-M. Lehr, Saarbrücken, Germany
	<b>Parallel Symposium</b> <b>Room: Auditorium</b> Pharmacogenetics at the bedside?	<b>Parallel Symposium</b> <b>Room: N-O</b> What will be the impact of cell-based therapy?	<b>Parallel Symposium</b> <b>Room: Forum</b> Simulation and modelling in drug development improves decisions, saving time and money?  Sponsoring Organisation
	Co-chairs G.T. Tucker, Sheffield, United Kingdom M. Schwab, Stuttgart, Germany	Co-chairs E. Cattaneo, Milan, Italy S. Nakagawa, Osaka, Japan	Co-chairs D. Stanski, Basel, Switzerland/East Hanover, NJ USA M. Danhof, Leiden, The Netherlands
14:15	Pharmacogenetics – how far is reality from expectation? G.T. Tucker, Sheffield, United Kingdom	Innovative neurogenic neural stem cell lines for neurodegenerative disease E. Cattaneo, Milan, Italy	Applying mechanistic pharmacokinetic-pharmacodynamic (PK/PD) models to drug development M. Danhof, Leiden, The Netherlands
14:50	Multiple gene pharmacogenetics in individualized drug therapy I. Ieri, Yonago, Japan	Cancer immunotherapy using genetically modified dendritic cells S. Nakagawa, Osaka, Japan	Examples of modelling and simulation in the pharmaceutical industry C. Pillai, Basel, Switzerland
15:25	Pharmacogenetics in cancer therapy M. Schwab, Stuttgart, Germany	Cardiac regeneration: Repopulating the heart L.J. Field, Indianapolis, IN USA	The role of innovative model-based trial design to improve drug development S. Duffull, Dunedin, New Zealand
16:00	Pharmacogenetics in paediatric drug development and utilisation: Are we going in the right direction? E. H. J. Kerkels, Leiden, The Netherlands (MO-S07-1)	A study of cell cycle and stem cell markers to identify the factors responsible for cardiac regeneration in mrl mice F. Moseley, Reading, United Kingdom (MO-S08-1)	Using pharmacokinetic-pharmacodynamic analysis in drug discovery. An example on the integration of mechanistic, principle and conceptual effect markers. S. Visser, Seodertaelje, Sweden (MO-S09-1)
16:15	The association of warfarin dosage in clinical use and pharmacogenomics Y.-H. Chen, Taipei, China Taiwan (MO-S07-2)	Alteration of endothelial cell function under high-glucose condition: Association with both disruption of cell-to-cell connection and non-muscle contraction K. Nobe, Tokyo, Japan (MO-S08-2)	Quantification of alpha 1-adrenoceptor concentration, ligand binding kinetics and inotropic response in the perfused rat heart: a PK/PD modeling analysis P. Sermsappasuk, Halle (Saale), Germany (MO-S09-2)
16:45-18:15	<b>EUFEPS Afternoon Sessions</b> See pages 18-20		

Parallel Symposium Room: L Metabolomics: What are the opportunities for biomarker discovery?			Parallel Symposium Room: C-D What benefits does Process Analytical Technology (PAT) bring to the design and assurance of product quality?		Parallel Symposium Room: A What have we learnt from recent safety cases for new drug development? ISPE Midyear Symposium	
Co-chairs <i>T. Hankemeier</i> , Leiden, The Netherlands <i>I. Schuppe-Koistinen</i> , Soedertaelje, Sweden			Co-chairs <i>S. Folestad</i> , Moelndal, Sweden <i>J. Pritchard</i> , Loughborough, United Kingdom		Co-chairs <i>H. Leufkens</i> , Utrecht, The Netherlands <i>M Sturkenboom</i> , Rotterdam, The Netherlands	
08:30	Systems biology & metabolomics: How far are we? <i>T. Hankemeier</i> , Leiden, The Netherlands		Real-time prediction and control of quality – the mechanistic approach to PAT <i>S. Folestad</i> , Moelndal, Sweden		A pharmaco-epidemiological reflection on recent drug safety cases <i>A. Walker</i> , Boston, MA USA	
09:05	The application of metabolic profiling technologies in biomarker discovery during drug R&D <i>I. Schuppe-Koistinen</i> , Soedertaelje, Sweden		The benefits of PAT in ICH and Japanese regulation <i>Y. Hiyama</i> , Tokyo, Japan		Class effects in drug safety and management <i>H. Leufkens</i> , Utrecht, The Netherlands	
09:40	Coffee Break					
10:10	Metabolomics by CE-MS for biomarker discovery <i>T. Soga</i> , Tsuruoka, Japan		Benefits of PAT for bioprocesses: Process design and quality assurance at the example of fermentation processes for recombinant protein production <i>A. Luebbert</i> , Halle, Germany		Industry responding to learning from safety cases <i>S. Perez-Gutthann</i> , Barcelona, Spain	
10:45	A novel immunoassay for monitoring caffeine as an environmental marker for pharmaceuticals input <i>J. J. Carvalho</i> , Berlin, Germany (MO-So4-1)		Evaluation of in-line near infrared spectroscopy for predicting tablet content uniformity during powder mixing <i>H. M. J. Salokangas</i> , Espoo, Finland (MO-So5-1)		Influence of COX-inhibitors on blood-brain barrier properties <i>B. Germann</i> , Vienna, Austria (MO-So6-1)	
11:00	Metabolomic approach for QA/QC on TCM material medica processing procedures-using citrus reticulata as the sample <i>W.-T. Chang</i> , Taichung, China Taiwan (MO-So4-2)		Raman spectroscopy as a PAT in tablet manufacturing <i>A. Sakr</i> , Cincinnati, USA (MO-So5-2)		Monitoring on drug-induced hepatopathy and granulocytopenia using hospital database resources: Prescription and laboratory data linkage <i>J. Kawakami</i> , Hamamatsu, Japan (MO-So6-2)	
11:15	Posters Lunch Break					
Parallel Symposium Room: L Dirty vs selective drugs in the CNS?  Sponsoring Organisation <div>Solvay Pharmaceuticals</div> <div></div>			Parallel Symposium Room: C-D How will developments in chemical methods sustain pharmaceutical industrial development?  Endorsing Organisation <div>c b g</div> <div></div>		Parallel Symposium Room: A How effective is the globalisation/harmonisation of pharmacovigilance?  Endorsing Organisation <div>c b g</div> <div></div>	
Co-chairs <i>H. Meltzer</i> , Nashville, TN USA <i>C. Senneff</i> , Weesp, The Netherlands			Co-chairs <i>T. Ohwada</i> , Tokyo, Japan <i>U. Holzgrabe</i> , Wuerzburg, Germany		Co-chairs <i>F. Lekkerkerker</i> , The Hague, The Netherlands <i>M. Braun</i> , Rockville, MD USA	
14:15	Rational polypharmacy within a single molecule: The basis for current antipsychotic treatment <i>H. Meltzer</i> , Nashville, TN USA		Process chemistry as leverage for drug development and profitability in the pharmaceutical industry <i>T. Konoike</i> , Amagasaki, Japan		The science underlying the practice of pharmacovigilance <i>N. Moore</i> , Bordeaux, France	
14:50	The treatment of major depression: Single or multiple target? <i>F. Artigas</i> , Barcelona, Spain		Active targeting of anticancer agents: Chemical aspects of folate-drug conjugate design <i>I. Vlahov</i> , West Lafayette, IN USA		International variety in interpretation and management of drug safety <i>N. Wathion</i> , London, United Kingdom	
15:25	Muscarinic receptors as a target in the treatment of disorders of the CNS: Antagonism, agonism or both? <i>B. Dean</i> , Melbourne, Australia		The synthetic development of the anti-influenza neuraminidase inhibitor oseltamivir phosphate (Tamiflu®): A challenge for synthesis and process research <i>M. Karpf</i> , Basel, Switzerland		ICH, CIOMS, ISOP, ISPE and other acronymic vehicles to enable harmonisation of pharmacovigilance <i>C-K. Shim</i> , Seoul, South Korea	
16:00	Learning and memory impairments in congenic C57BL/6NTac mice that lack the m2 muscarinic acetylcholine receptor subtype <i>C. Wrenn</i> , Des Moines, USA (MO-S10-1)		Random chemistry as a new tool for the generation of small-compound libraries <i>U. Holzgrabe</i> , Wuerzburg, Germany (MO-S11-1)		Round Table Discussion	
16:15	Neuronal protective effect of recombinant arginine deiminase in a nitric oxide overexpression cell culture system <i>H.-H. Yu</i> , Taipei, China Taiwan (MO-S10-2)		Generation and application of o-benzoquinone methides bearing various substituents on the benzene ring <i>T. Ohwada</i> , Tokyo, Japan (MO-S11-2)		Proactive risk management (plans): Where are we? Convenor <i>F. Lekkerkerker</i> , The Hague, The Netherlands	
16:45-18:15	EUFPS Afternoon Sessions See pages 18-20					

# SCIENTIFIC PROGRAM

	<b>Parallel Symposium</b> <b>Room: Forum</b> <b>Druggability: A concept that will fill the pipeline?</b>  <b>Sponsoring Organisation</b>	<b>Parallel Symposium</b> <b>Room: Auditorium</b> <b>Drug targeting: How successful are we?</b>	<b>Parallel Symposium</b> <b>Room: N-O</b> <b>What is the state-of-the-science in receptor site modelling?</b>
	Co-chairs <i>D. Nicholson</i> , Oss, The Netherlands	Co-chairs <i>M. Hashida</i> , Kyoto, Japan <i>R. Duncan</i> , Cardiff, United Kingdom	Co-chairs <i>S. Dahl</i> , Tromsø, Norway <i>M. Ishiguru</i> , Osaka, Japan
08:30	Druggability and the concept of ADME space <i>D. Smith</i> , Sandwich, United Kingdom	Drug and gene delivery by combination of ultrasound and bubble liposomes <i>K. Maruyama</i> , Kanagawa, Japan	Genomics to drug targets by molecular modelling <i>S. Dahl</i> , Tromsø, Norway
09:05	Drugability and drug-likeness: A medicinal chemist's view <i>B. Testa</i> , Lausanne, Switzerland	Polymeric conjugates as anticancer nano-medicines: Mechanism of action and drug combinations <i>R. Duncan</i> , Cardiff, United Kingdom	Functional structural models of G protein coupled receptors <i>M. Ishiguru</i> , Osaka, Japan
09:40	<b>Coffee Break</b>		
10:10	Concave druggability of protein surfaces for accelerating <i>in silico</i> screening <i>H. Shirai</i> , Tsukuba, Japan	Targeting with molecularly decorated nanoparticles <i>N. Peppas</i> , Austin, TX USA	Structure-based virtual screening <i>J. Irwin</i> , San Francisco, CA USA
10:45	Predicting druggable proteins from amino acid sequence by a machine learning approach <i>C.W. Yap</i> , Singapore, Singapore (TU-S01-1)	Prevention of cytokines responses in cardiac allograft rejection by systemic injection of nf-kappa B decoy/mannosylated cationic liposome complexes <i>Y. Higuchi</i> , Kyoto, Japan (TU-S02-1)	Identification of a conserved hydrophobic asparagine-cage as a constraint for family A GPCR activation <i>A. Jongejan</i> , Amsterdam, The Netherlands (TU-S03-1)
11:00	Medicinal chemistry of hERG optimisations <i>E. Moir</i> , Newhouse, United Kingdom (TU-S01-2)	Squalenoylated-gemcitabine nanomedicine exhibits potential in cancer therapy at preclinic <i>H. R. Lakireddy</i> , Châtenay-Malabry, France (TU-S02-2)	Delineating a powerful virtual screening protocol for G-protein coupled receptors: Application to selective kappa opioid receptor agonist, salvinorin A <i>N. Singh</i> , Mississippi, USA (TU-S03-2)
11:15	<b>Posters</b> <b>Lunch Break</b>		
	<b>Keynote Presentation</b> <b>Room: Auditorium</b> <b>Sponsoring Organisation</b> 	<b>Keynote Presentation</b> <b>Room: Forum</b> <b>Sponsoring Organisation</b> Solvay Pharmaceuticals 	
	Chair <i>W. Sadee</i> , Columbus OH USA	Chair <i>C. Sennef</i> , Weesp, The Netherlands	
12:15	Are we meeting the challenges of resistance to anti-infective drugs and of newly emerging infectious diseases? <i>L. Schlesinger</i> , Columbus OH USA (KLT-1)	Will novel approaches to the treatment of cardiovascular disease prove highly effective? <i>J. Kastelein</i> , Amsterdam, The Netherlands (KLT-2)	
	<b>Round Table Discussion</b> <b>Room: A</b> <b>Sponsoring Organisation</b> 	<b>Round Table Discussion</b> <b>Room: L</b>	<b>Round Table Discussion</b> <b>Room: C-D</b>
13:10	Translational science: a solution to the productivity gap? Convenor <i>D. Nicholson</i> , Oss, The Netherlands	Can microdosing accelerate drug development? Convenor <i>A. Grahnen</i> , Uppsala, Sweden	When is a human bioequivalence study not needed? Convenor <i>L.Z. Benet</i> , San Francisco, CA USA
	<b>Parallel Symposium</b> <b>Room: A</b> <b>Drug-drug interactions: Avoid or understand?</b>	<b>Parallel Symposium</b> <b>Room: N-O</b> <b>What is the future of smart, feed-back, on-demand drug delivery systems?</b>	<b>Parallel Symposium</b> <b>Room: Forum</b> <b>How are disease and PK-PD connected?</b>
	Co-chairs <i>K. Thummel</i> , Seattle, WA USA <i>A. McLachlan</i> , Sydney, Australia	Co-chairs <i>J. Kopecek</i> , Salt Lake City, USA <i>K. Kataoka</i> , Tokyo, Japan	Co-chairs <i>Y. Tanigawara</i> , Tokyo, Japan <i>R. Bruno</i> , Mountain View, CA USA
14:15	Managing herb-drug interactions: Understanding mechanism and educating the public <i>A. McLachlan</i> , Sydney, Australia	Smart drug delivery systems: State-of-the-art and future directions <i>J. Kopecek</i> , Salt Lake City, USA	Bone disease progression and drug action <i>N. Holford</i> , Auckland, New Zealand
14:50	Role of the pharmacist in avoiding drug-drug interactions with patient self-care <i>E. Nakashima</i> , Tokyo, Japan	Light-induced gene and drug delivery by supramolecular nanocarrier <i>K. Kataoka</i> , Tokyo, Japan	Mechanism-based modelling of disease progression – disease system analysis <i>B. Ploeger</i> , Leiden, The Netherlands
15:25	Application of a drug-drug interaction data base in drug development and clinical education <i>K. Thummel</i> , Seattle, WA USA	Smart polymeric carriers for biomolecular drugs <i>P. Stayton</i> , Seattle, WA USA	Modelling of cancer progression and drug effects <i>R. Bruno</i> , Mountain View, CA USA
16:00	Sfinx – construction and implementation of a novel drug drug interaction database <i>B. Eiermann</i> , Stockholm, Sweden (TU-S07-1)	Transdermal iontophoresis of dopamine agonist 5-OH-DPAT: Correlation of in vitro transport to the integrated pk-pd profiles based on non-linear mixed effect modeling <i>A. K. Nugroho</i> , Yogyakarta, Indonesia (TU-S08-1)	Pharmacokinetic-pharmacodynamic model for propofol during long-term sedation in the critically ill patient <i>M. Peeters</i> , Nieuwegein, The Netherlands (TU-S09-1)
16:15	Can we predict the magnitude of drug-drug interaction in a simple way?: Simulation of interaction of rapidly-eliminating drugs with fluvoxamine by dynamo-pk analysis method <i>K. Iga</i> , Kyotanabe, Japan (TU-S07-2)	Biodegradable microparticles containing dexamethasone and spions for intra-articular delivery <i>N. Butoescu</i> , Geneva, Switzerland (TU-S08-2)	Population pharmacokinetic modelling of radioiodine turnover in patients with Graves' disease <i>I. Grabnar</i> , Ljubljana, Slovenia (TU-S09-2)
16:45-18:15	<b>EUFEPS Afternoon Sessions See pages 18-20</b>		



	<b>Parallel Symposium</b> <b>Room: A</b> What's new in methods of measuring human drug response?  Sponsoring Organisation	<b>Parallel Symposium</b> <b>Room: C-D</b> In silico product development from molecule to man: Dream or reality?	<b>Parallel Symposium</b> <b>Room: L</b> Nutraceuticals: Are new methods of evaluating risk/benefit required?
	Co-chairs <i>A. Cohen</i> , Leiden, The Netherlands <i>P. Macheras</i> , Athens, Greece	Co-chairs <i>P. York</i> , Bradford, United Kingdom <i>V. Venkatasubramanian</i> , West Lafayette, IN USA	Co-chairs <i>H. Ohama</i> , Tokyo, Japan <i>R. Oledzka</i> , Warsaw, Poland
08:30	The data intensive first administration to man study – functional outcome replaces tolerability: The impact of new measurement techniques <i>A. Cohen</i> , Leiden, The Netherlands	Structure, thermodynamics and kinetics of pharmaceutical systems from molecular simulation <i>J. Anwar</i> , Bradford, United Kingdom	Risk analysis and evaluation of scientific evidence for nutraceuticals in Japan <i>H. Ohama</i> , Tokyo, Japan
09:05	Pathophysiological concepts as a basis for the measurement of treatment response in inflammatory and obstructive airway disease <i>H. Reddel</i> , Camperdown, Australia	Computer aided design and optimisation for pharmaceutical formulations <i>K. Takayama</i> , Tokyo, Japan	Risk assessment and benefit evaluation for nutraceuticals <i>J. Hathcock</i> , Washington, DC USA
09:40	<b>Coffee Break</b>		
10:10	PET imaging for evaluation of drug effects in neuropsychiatric disease <i>M. Laurelle</i> , Greenford, United Kingdom, and New York, NY USA	Cyberinfrastructure enabled pharmaceutical products design and engineering opportunities and challenges <i>V. Venkatasubramanian</i> , West Lafayette, IN USA	Benefits of antioxidants <i>H.K. Biesalski</i> , Hohenheim, Germany
10:45	Sensitivity of the items of the Montgomery Asberg depression rating scale to treatment response: Impact of different endpoints on clinical study design for antidepressant drugs <i>G. Santen</i> , Leiden, The Netherlands (TU-So4-1)	Expert system software for solid dosage form formulation design. <i>E. Krausbauer</i> , Basel, Switzerland (TU-So5-1)	Organic vs. conventional apple juices: polyphenol profile, anti-oxidant capacity, anti-cancer activity, and inflammatory bowel disease modulator activity <i>E. Mejia-Meza</i> , Pullman, USA (TU-So6-1)
11:00	Application of computation in translational research: A randomized trial of intravesical mitomycin c for superficial bladder cancer with 10-year follow-up <i>J. Au</i> , Columbus, OH USA (TU-So4-2)	Fundamental understanding through simulations? <i>S.-M. Siiriä</i> , Helsinki, Finland (TU-So5-2)	Effects of essential fatty acids on expression level and function of P-glycoprotein in inflammatory bowel diseases <i>A. Nomura</i> , Tokyo, Japan (TU-So6-2)
11:15	<b>Posters Lunch Break</b>		
	<b>Parallel Symposium</b> <b>Room: Auditorium</b> Have omics technologies improved the chance for successful drug development?	<b>Parallel Symposium</b> <b>Room: L</b> Are pharmaceutical manufacturing technologies in stagnation?	<b>Parallel Symposium</b> <b>Room: C-D</b> Are we using the right outcome measures to ascertain patient benefit from drug therapy?
	Co-chairs <i>T. Guentert</i> , Basel, Switzerland <i>M. Bleavins</i> , Ann Arbor, MI USA	Co-chairs <i>J. Fix</i> , Lawrence, KS USA <i>Y. Capan</i> , Ankara, Turkey	Co-chairs <i>G. Skrepnek</i> , Tucson, TX USA <i>A. Hussain</i> , Dubai, United Arab Emirates
14:15	Genomics and drug discovery: Have the promises been fulfilled? <i>K. Lindpaintner</i> , Basel, Switzerland	The changing landscape of pharmaceutical manufacturing: Incremental or breaking new ground? <i>J. Fix</i> , Lawrence, KS USA	Outcomes measurement: Overview of theoretical and applied issues <i>G. Skrepnek</i> , Tucson, TX USA
14:50	Omics and the search for improved biomarkers <i>M. Bleavins</i> , Ann Arbor, MI USA	Innovation needed in pharmaceutical research and technology in the 21st century <i>R. Ibuki</i> , Yaizu, Japan	Outcome measurement: Issues and strategies for pre-market development <i>W.-F. Huang</i> , Taipei, China Taiwan
15:25	Changing the drug development paradigm: Opportunities offered by new technologies <i>J. Kuromitsu</i> , Tsukuba, Japan	Can new production technologies and new excipients meet the demands of future drugs? <i>H. Frijlink</i> , Groningen, The Netherlands	Outcomes measurement: Issues and strategies for post-marketing development <i>J. Cooke</i> , Manchester, United Kingdom
16:00	Microarray analysis of chlamydia pneumoniae infected human epithelial cell line using gene ontology hierarchy <i>J. Alvesalo</i> , Helsinki, Finland (TU-S10-1)	Monitoring the modification of budesonide-lactose interactions within dry powder inhaler formulations using atomic force microscopy <i>F. Buttini</i> , Parma, Italy (TU-S11-1)	Abuse & misuse of lifestyle drugs in Korea <i>K. Kwon</i> , Seoul, South-Korea (TU-S12-1)
16:15	Quantitative proteomic analysis of human renal cell carcinoma using the NBS method <i>J. Matsumoto</i> , Kobe, Japan (TU-S10-2)	Production of beclomethasone and salbutamol loaded poly(lactic acid) nanoparticles by a novel electrospraying technique <i>L. Peltonen</i> , Helsinki, Finland (TU-S11-2)	Combined prescriptions of cardiovascular drugs and Ginkgo biloba in Taiwan: A population-based study <i>L.-C. Chang</i> , Taipei, China Taiwan (TU-S12-2)
16:45-18:15	<b>EUFPS Afternoon Sessions See pages 18-20</b>		


# SCIENTIFIC PROGRAM

	Parallel Symposium Room: N-O How to manage drug therapy at the extremes of age?	Parallel Symposium Room: Forum How can nanotechnology and materials science solve drug delivery problems?	Parallel Symposium Room: Auditorium How important is genetic and physiological variability in drug transporters?
	Co-chairs <i>H. Derendorf</i> , Gainesville, FL USA <i>H. Christensen</i> , Oslo, Norway	Co-chairs <i>C-M. Lehr</i> , Saarbrücken, Germany <i>H. Ghandehari</i> , Baltimore, USA	Co-chairs <i>K. Giacomini</i> , San Francisco, CA USA <i>K. Inui</i> , Kyoto, Japan
08:30	Pharmacotherapy in the elderly <i>H. Derendorf</i> , Gainesville, FL USA	Nanomedicines for overcoming biological barriers <i>C-M. Lehr</i> , Saarbrücken, Germany	Functional genomics of membrane transporters <i>K. Giacomini</i> , San Francisco, CA USA
09:05	Pharmacokinetics and pharmacodynamics in neonates and infants <i>T. Dalla Costa</i> , Porto Alegre, Brazil	Particle design for absorption enhancement using a 4-nozzle spray drier and DNA vaccine by self-organised Tat nanospheres <i>H. Okada</i> , Tokyo, Japan	Pharmacogenomics of MDR1/ABCB1 and CYP3As in tacrolimus therapy after organ transplantation <i>K. Inui</i> , Kyoto, Japan
09:40	<b>Coffee Break</b>		
10:10	Dose optimisation in neonates, infants and children <i>S. Higuchi</i> , Fukuoka, Japan	Can higher definition of the nanoscale result in better drug delivery systems in the 21st century? <i>H. Ghandehari</i> , Baltimore, USA	Assessing the impact of variability in ABC drug transporters using mouse models <i>A. Schinkel</i> , Amsterdam, The Netherlands
10:45	Prediction of the oral bioavailability of midazolam in the first 2 years of life <i>T. N. Johnson</i> , Sheffield, United Kingdom (WE-So1-1)	Development of octaarginine-modified multifunctional envelope-type nano device for gene delivery <i>K. Kogure</i> , Sapporo, Japan (WE-So2-1)	Web-based comprehensive database for all about drug transporters, "TP-Search" <i>K. Maeda</i> , Tokyo, Japan (WE-So3-1)
11:00	Towards the in silico child: midazolam pharmacokinetics using physiologically-based pharmacokinetic vs. non-linear mixed effects modeling <i>S. Willmann</i> , Leverkusen, Germany (WE-So1-2)	In vitro studies into the biological fate of pva nanoparticles for pulmonary delivery <i>M. Orlu</i> , Istanbul, Turkey (WE-So2-2)	Regulatory mechanisms for gene expression of human organic anion transporters <i>K. Ogasawara</i> , Kyoto, Japan (WE-So3-2)
11:15	<b>Posters Lunch Break</b>		
	<b>Keynote Presentation</b> Room: Auditorium Sponsoring Organisation 	<b>Keynote Presentation</b> Room: Forum	
	Chair <i>J. Pritchard</i> , Macclesfield, United Kingdom	Chair <i>K. Midha</i> , Saskatoon, Canada	
12:15	Drug resistance in cancer chemotherapy <i>T. Tsuruo</i> , Tokyo, Japan (KLW-1)	Recent progress in prion biology <i>M. Heikenwaelder</i> , Zurich, Switzerland (KLW-2)	
	<b>Round Table Discussion</b> Room: A	<b>Round Table Discussion</b> Room: A	<b>Round Table Discussion</b> Room: C-D
13:10	What is the value of observational data post-marketing to assess safety and efficacy? Convenor <i>M. Rowland</i> , Manchester, United Kingdom	Life-style drugs: a new burden to the health system? Convenor <i>G. Alvan</i> , Uppsala, Sweden	Is there a consensus on guidelines for the evaluation of biosimilars? Convenor <i>V. Shah</i> , Rockville, MD USA
	<b>Parallel Symposium</b> Room: N-O Is the patient taking the tablets?	<b>Parallel Symposium</b> Room: Auditorium Developing biotech products: What are the challenges and solutions? 	<b>Parallel Symposium</b> Room: Forum Control of intracellular pharmacokinetics: Advantages for drug therapy?
	Co-chairs <i>M. Roberts</i> , Brisbane, Australia <i>J. Kennedy</i> , Cork, Ireland	Co-chairs <i>M. Tsuchiya</i> , Gotenba, Japan <i>S. Frokjaer</i> , Copenhagen, Denmark	Co-chairs <i>H. Kroemer</i> , Greifswald, Germany <i>D. Roden</i> , Memphis, TN USA
14:15	When are dose administration aids of benefit? <i>M. Roberts</i> , Brisbane, Australia	The challenge of the next generation of therapeutic antibodies <i>M. Tsuchiya</i> , Gotenba, Japan	Transporter mediated cellular uptake of drugs as a prerequisite for drug action <i>H. Kroemer</i> , Greifswald, Germany
14:50	Ambulatory patient's variable adherence with prescribed drug dosing regimens: Prevalence, patterns, practicalities for drug trials and patient care <i>J. Urquhart</i> , Palo Alto, CA USA, and Maastricht, The Netherlands	Drug delivery systems for biopharmaceuticals <i>S. Frokjaer</i> , Copenhagen, Denmark	Intracellular pharmacokinetics determines drug action in patients with HIV <i>R. Kim</i> , London, Canada
15:25	Medication errors and human factors in medication use safety <i>P. Schneider</i> , Columbus, OH USA	Efficient transepithelial delivery of biopharmaceuticals <i>J. Mrsny</i> , Cardiff, United Kingdom, and Menlo Park, CA USA	Understanding drug-induced arrhythmias – from intracellular concentrations to candidate genes <i>D. Roden</i> , Nashville, TN USA
16:00	Routine use of dose administration aids (DAAs) in the community - characteristics of Australian consumers making this choice <i>J. Stokes</i> , Brisbane, Australia (WE-So7-1)	Oral delivery of insulin by new polysaccharide nanoparticles <i>B. Sarmento</i> , Porto, Portugal (WE-So8-1)	Methods in drug discovery: measurement of unbound intracellular drug concentrations <i>M. Friden</i> , Uppsala, Sweden (WE-So9-1)
16:15	A cross-national study of persistence of anti-hypertensive medication use in the elderly <i>B. L. Van Wijk</i> , Utrecht, The Netherlands (WE-So7-2)	High pressure treatment for the recovery of active protein from protein aggregates: An enabling technology in comparison to traditional chaotrope-based refolding methods <i>M. Seefeldt</i> , Boulder, USA (WE-So8-2)	Uptake mechanisms of anti-hiv drugs, 2', 3'-dideoxyinosine and 3'-azido-3'-deoxythymidine by a conditionally immortalized syncytiotrophoblast cell line, TR-TBT <i>K. Sato</i> , Tokyo, Japan (WE-So9-2)

# AMME • Wednesday

	<b>Parallel Symposium</b> <b>Room: A</b> Systems biology: A driver of drug discovery and development?  Sponsoring Organisation	<b>Parallel Symposium</b> <b>Room: C-D</b> How to engineer desired particle properties for drug delivery?	<b>Parallel Symposium</b> <b>Room: L</b> Off-label use of medicines: Abuse or a vehicle for innovation?
	Co-chairs C.R. Noe, Vienna, Austria	Co-chairs E. Fattal, Paris, France T. Nagai, Tokyo, Japan	Co-chairs A. Kalis, The Hague, The Netherlands
08:30	Systems Biology: What does it mean for pharmaceutical sciences? A. Aszodi, Vienna, Austria	Particle design for nucleic acids and contrast agents E. Fattal, Châtenay-Malabry, France	Two worlds and why the twain will never meet J. Lisman, Amsterdam, The Netherlands
09:05	The theory of biological robustness and its applications to medicine H. Kitano, Tokyo, Japan	Novel particle design for drug delivery H. Takeuchi, Gifu, Japan	Regulatory and economic aspects of off-label drug use A. Wertheimer, Philadelphia, PA USA
09:40	<b>Coffee Break</b>		
10:10	Reconstruction of the genome-wide human metabolic networks: conceptual and practical uses B. Palsson, La Jolla, CA USA	How to optimize particle properties for pulmonary drug delivery G. Hochhaus, Gainesville, FL USA	The benefits of off-label drug use and its utilisation K. Tsutani, Tokyo, Japan
10:45	From enzymes to cells and back: Integrating biochemical and cellular profiling of small molecule kinase inhibitors J. J. Hornberg, Oss, The Netherlands (WE-So4-1)	The role of particle characterization in the development and dosage form evaluation of a poorly soluble pharmaceutical drug product R. Govoreanu, Beerse, Belgium (WE-So5-1)	On-label and off-label prescribing of erythropoietic agents (epoetin alfa and darbepoetin alfa) in critically ill patients: a multi-center, retrospective study D. Holdford, Richmond, USA (WE-So6-1)
11:00	Pharmbiosim - biosimulation of drug metabolism J. Smolinski, Dresden, Germany (WE-So4-2)	Supercritical fluid particle design for increasing dissolution rate of poorly-soluble active pharmaceutical ingredients F. Deschamps, Champigneulle, France (WE-So5-2)	High rate of off-label use in cardiovascular paediatric pharmacotherapy requires new focus in research L. Hsien, Düsseldorf, Germany (WE-So6-2)
11:15	<b>Posters Lunch Break</b>		
	<b>Parallel Symposium</b> <b>Room: A</b> Molecular targeting in cancer chemotherapy?	<b>Parallel Symposium</b> <b>Room: C-D</b> Miniaturisation in analytical methods: Is small always beautiful?	<b>Parallel Symposium</b> <b>Room: L</b> Counterfeiting of medicines: Detection and prevention?
	Co-chairs J. Au, Columbus, OH USA S. Eck, Ann Arbor, MI USA	Co-chairs S. Lunte, Lawrence, KS USA J. Haginaka, Nishinomiya, Japan	Co-chairs A. Moffat, London, United Kingdom Z. Y. Yang, Guangzhou, China
14:15	The challenges of developing targeted cancer therapies: An industry perspective S. Eck, Ann Arbor, MI USA	Separation based sensors for pharmaceutical analysis using microdialysis and microchip electrophoresis S. Lunte, Lawrence, KS USA	New methods for detection of counterfeit medicines for laboratory and field use A. Moffat, London, United Kingdom
14:50	A systems pharmacology: Targeting p53 networks E. Liu, Singapore, Singapore	Micro and nano chemical systems on chips for analytical and biological sciences T. Kitamori, Tokyo, Japan	Analytical methods to detect and fingerprint counterfeit medicines F. Fricke, Cincinnati, USA
15:25	Translational research on a drug with multiple molecular targets (suramin) J. Au, Columbus, OH USA	Microchip array strategies for biomarker detection using fluorescence and MALDI TOF MS readout T. Laurell, Lund, Sweden	Combating counterfeit drugs in Asia Z. Y. Yang, Guangzhou, China
16:00	Anti-angiogenic actions of liposomal glucocorticoids on tumor growth M. Banciu, Utrecht, The Netherlands (WE-S10-1)	Molecular imaging of redox reaction using OMRI/nitroxyl probe technique K.-I. Yamada, Fukuoka, Japan (WE-S11-1)	Transfer of an NIR method for the authentication of tablets and the detection of counterfeit versions A. J. O'Neil, London, United Kingdom (WE-S12-1)
16:15	Characterization of paclitaxel-loaded immunonanoparticles A. Cirstoiu-Hapca, Geneva, Switzerland (WE-S10-2)		Spurious drugs –epidemic threat to public health and pharma industries V. Mshra, Sagar, India (WE-S12-2)



Monday		
EUFEPS Afternoon Sessions		
<b>Room: O</b> Strategic, innovative and critical drug research initiatives: One year later Chair <i>O.J. Bjerrum</i> , Copenhagen DK  16:45-18:15 Strategic drug research initiatives in Europe: Current and future needs <i>O.J. Bjerrum</i> , Copenhagen DK  The European Innovative Medicines Technology Platform: Current status 2007 <i>B. Rainer</i> , Brussels BE  Precompetitive industry collaboration in Europe: 18 months experience of the InnoMed project on predictive toxicology <i>D. Tweats</i> , Swansea UK  Precompetitive industry collaboration in Europe: 18 months experience of the InnoMed project on biomarkers for Alzheimer's disease <i>P. Francis</i> , London UK  Discussion	<b>Room: L</b> Pharmaceutical sciences research training and education: Needs and supply Chair <i>M. Van der Waart</i> , Oss NL  University perspective <i>S. de Smedt</i> , Ghent BE  Industry perspective <i>J. Dirach</i> , Copenhagen DK  PPP Research Perspective <i>V. Nickolson</i> , Leiden NL  Discussion	<b>Room: N-O</b> The EU Microdosing AMS Partnership Programme (EUMAPP) Chair <i>R.A. de Zeeuw</i> , Assen NL  Microdosing: Pros and cons in translational medicine <i>C. Garner</i> , York UK  EUMAPP: Objectives, approaches and current status <i>B. Oosterhuis</i> , Zuidlaren NL  Microdosing: Servier strategy and expectations from EUMAPP <i>E. Foos-Gilbert</i> , Courbevoie FR  Discussion
Tuesday		
EUFEPS Afternoon Sessions		
<b>Room: E</b> European drug development centres and European growth areas Chair <i>P. Vuorela</i> , Turku FI  16:45-18:15 Center for New Drug Discovery Tools - DDTCC <i>A. Urtti</i> , Helsinki FI  The Pharma game, new rules, new players <i>V. Nickolson</i> , Leiden NL  Drug development in the Medicon Valley: Importance of the binational cluster and its international contacts <i>S. Gestrelus</i> , Copenhagen DK  Discussion	<b>Room: L</b> Reformulation of old drugs: Life cycle management Chair <i>H. Blume</i> , Oberursel DE  Modified drug delivery – development rationale for therapeutic improvement <i>E. Soederlind</i> , Moelndal SE  Product performance in the gastrointestinal tract and perspectives for optimisation <i>W. Weitschies</i> , Greifswald DE  Life Cycle Management: new chances for old drugs <i>H. Blume</i> , Oberursel DE  Discussion Session Sponsor 	<b>Room: C-D</b> Vaccine delivery Co-Chairs <i>J. Bouwstra</i> , Leiden NL <i>W. Jiskoot</i> , Leiden NL  Virosomes as a platform for improved influenza vaccines <i>A. Huckriede</i> , Groningen NL  Vaccines for Hepatitis B using DNA and sub-unit antigens <i>Y. Perrie</i> , Aston UK  Challenges in non-invasive vaccine delivery <i>W. Jiskoot</i> , Leiden NL  Discussion

## Strategic, Innovative and Critical Drug Research Initiatives: One Year Later

Large initiatives to promote drug sciences, supporting the European pharma industry, have recently been launched or are in their late phase of preparation. Major ones include the European 7th Framework Programme for Research and Technological Development, as well as the Innovative Medicines Initiative (IMI) by the European Pharmaceutical Industries and Associations (EFPIA) and the European Commission, including the InnoMed, PredTox and AddNeuromed projects. Substantial experience of working together in new constellations is gained, already, and considered producing a role model for future collaboration in new IMI projects.

Current status of these initiatives and experiences obtained to date will be presented. Presenta-

tions will also be an introduction to a discussion about best practice for joint projects and initiatives.

## Pharmaceutical Sciences Research Training and Education: Needs and Supply

Traditionally, academia is primarily involved in defining and executing training and education programmes, as lectures, workshops and courses. The EUFEPS Committee on Training and Education (CTE; [www.eufeps.org](http://www.eufeps.org)) is also involved in defining and organising training activities, which will be introduced and discussed.

Education and training is an important activity of the Innovative Medicines Initiative (IMI; [www.imi-europe.org](http://www.imi-europe.org)). The IMI is a public-private partnership, including patient organisations, universities, hospitals, and regulatory authorities as well as small and large biopharmaceutical and healthcare companies. Drug effi-

cacy, safety, knowledge management and education and training are all pillars of the IMI.

The newly established Dutch Top Institute Pharma (see other summary) puts strong emphasis on training and education. It is a practical example how three partners, all needing well educated and trained scientists are working closely together.

## The EU Microdosing AMS Partnership Programme (EUMAPP)

Microdosing is a new safe way to obtain essential human drug metabolism and pharmacokinetic (PK) information with minimal animal testing. Both the European and the US regulatory authorities have published microdosing guidance documents and are open for business. The scientific basis of microdosing relies on that there is reasonable predictivity between microdose and pharmacological dose PK.

## Monday

### EUFEPS Afternoon Sessions

#### Room: E

The European Pharma Sciences Leadership Forum (EuPSLF)  
Chair  
C.R. Noe, Vienna AT

16:45-18:15

Background, initiative and progress  
H. H. Linden, Stockholm SE

Aims, ambitions and plans  
R. Pellicciari, Perugia IT

Preparing for a changing world of science  
C.R. Noe, Vienna AT

Discussion

#### Room: C-D

Pharmaceutical sciences *in silico* learning systems: Value and availability  
Chair  
N. Haider, Vienna AT

Computer applications in pharmaceutical education and research  
B. Ernst, Basel CH

Pharmasquare – Blended Learning in Pharmaceutical Sciences  
S. Moss, Bath UK

PharmXplorer, an integrated platform for e-learning in pharmaceutical sciences  
T. Langer, Innsbruck AT

Discussion

Session Sponsor



EUFEPS Afternoon Sessions

Sponsoring Organisation



TI PHARMA

## Tuesday

### EUFEPS Afternoon Sessions

#### Room: O

Pharmacogenetics & Pharmacogenomics Workshop: Outcomes and plans  
Chair  
A-H. Maitland-van der Zee, Utrecht NL

16:45-18:15

Pharmacogenetics of adverse drug reactions  
M. Pirmohamed, Liverpool UK

Pharmacogenetics in paediatrics  
E. Jacqz-Aigran, Paris FR

Methods in pharmacogenetics/genomics  
A-H. Maitland-van der Zee, Utrecht NL

Discussion

#### Room: A

How to start up a new company?  
Co-Chairs  
H. Lennernäs, Uppsala SE  
C. Bogentoft, Stockholm SE

Bridging between entrepreneurs and the pharmaceutical industry  
H. Lennernäs, Uppsala SE

The hands-on experience  
G.T. Tucker, Sheffield UK

The Karolinska Innovation model in starting up companies  
C. Bogentoft, Stockholm SE

Discussion

#### Room: N-O

Drug product quality after new legislation  
Chair  
H. Kőszegi-Szalai, Budapest HU

The impact of new guidance documents on the quality of medicines in Europe  
D. van Riet, Bilthoven NL

The present and the expectable future role of the EP in the standardisation of the quality of medicines in Europe  
H. Kőszegi-Szalai, Budapest HU

Discussion

EUMAPP is a grouping of 10 companies and organisations funded by the EU to a value of over 3 million euros who are undertaking an ambitious programme to (1) examine microdose/pharmacological dose comparisons for seven drugs (2) use microdose PK data to better model human PK in combination with *in silico* and *in vitro* methods (3) compare different analytical approaches and (4) to disseminate the results to both professional and lay people. Will EUMAPP help to put microdosing into the critical path of drug development?

### The European Pharma Sciences Leadership Forum (EuPSLF)

The process of drug discovery, development and utilisation for the improvement of human and animal health and welfare is a complex process that involves many individual scientists and organisations

from a wide variety of scientific backgrounds. They form scientific communities that, directly and indirectly, contribute to better, new, innovative and safe medicines.

Initiatives towards a “European Pharma Sciences Leadership Forum”, comprising the presidents of ten partnering federations and associations, were taken less than one year ago. It is under way and should work for advancing science, engage in European strategic research initiatives, and contribute to relevant training and education of scientists in the wide field of pharmaceutical and related research and development. Speaking up with one voice would, furthermore, demonstrate united commitment to solving problems and setting priorities for both short- and long-range new drug developments and applications in Europe.

### Pharmaceutical Sciences *In-silico* Learning Systems: Value and Availability

Academic education in the pharmaceutical sciences is facing a rapidly increasing amount of available information (new insights, latest research results, emerging new disciplines), which has to be turned into profound knowledge, excellent theoretical and practical skills, and decision-making capabilities of our students and graduates. State-of-the-art information technology is becoming increasingly important in making teaching/learning processes more efficient. This technology has the potential to strengthen the classical link between scientific research and education.

Today, “*in-silico* learning”, “computer-aided learning”, “eLearning”, or “blended learning” is going to grow from scattered pilot projects, initiated by a few pioneers, into larger

initiatives on a regional, national, and increasingly international level, especially in the pharmaceutical sciences. Such larger “*in-silico* learning” initiatives will be our focus, including “live” demonstrations.

#### **European drug development centres and European growth areas**

The Drug Discovery and Development Technology Centre is a new research centre located in the Faculty of Pharmacy, University of Helsinki. It is a multi-disciplinary unit concentrating on improved technologies for preclinical drug discovery and development.

The Duch Top Institute Pharma is a public-private partnership, including universities, academic medical centres, small, medium and large life science companies. Academic and industrial parties contribute know-how as well as other resources. It is their goal to develop novel, cross-disciplinary drug discovery and development processes that will reduce “time and cost to patient”.

The Medicon Valley Academy is a network of universities, hospitals and companies. The cluster itself, and the drug development activities of it are, primarily, in the “Medicon Valley” (Copenhagen DK and in Malmö/Lund SE region). Goals include improving Danish-Swedish collaboration. Collaboration with ScanBalt links to other Nordic/Baltic Sea initiatives and organisations.

#### **Reformulation of Old Drugs: Life Cycle Management**

Could the life cycle of drugs available since long be further expanded? Utilising and combining the wealth of knowledge and experience gained of old drugs with new scientific insights and regulatory requirements should provide such an opportunity for improved drug therapy. Approaches in doing this include, for example, improved and modified drug delivery at the target of drug action. For oral drugs, product performance in the gastrointestinal tract and how to optimise it is crucial. Perhaps, not all old drugs would survive, but for many systematic “Life Cycle Management” will provide new chances.

In this session, the above and related issues will be addressed by experts and further discussed.

#### **Vaccine Delivery**

The proper delivery of antigens is a major hurdle in the search for potent vaccines against several priority diseases, such as tuberculosis, HIV/AIDS, malaria, and pandemic influenza. The choice for a certain delivery vehicle depends on the physicochemical characteristics of the antigen(s), the disease and type of immune response that is desired (e.g., Th1/Th2 ratio), as well as the route of administration.

In this session several hot topics in vaccine delivery will be presented and discussed, including innovative delivery systems for (pandemic) flu and hepatitis B vaccines, delivery issues in the field of genetic (DNA) vaccination, and needle-free vaccine delivery approaches.

#### **Pharmacogenetics & Pharmacogenomics Workshop: Outcomes and Plans**

A workshop was set up, recently, to bring European scientists in pharmacogenetics and pharmacogenomics together to start developing a roadmap for better collaboration and intensified research, including a network among them and their groups, also towards personalised medicines. Development recommendations from the workshop included, for example: Platform for gathering and promoting knowledge about pharmacogenetics in Europe; mechanisms for sharing and extending existing research, databases and bio-banks; extended collaboration between academia and the pharmaceutical industry; better education and training in pharmacogenetics and genomics; and increased European funding.

Presentations of this session will focus on important research areas and tools, all to further encourage and stimulate discussion, new initiatives and collaborations. Input in the discussion on how to proceed as to the network will be welcome.

#### **How to Start up a New Company?**

The first contribution of this session will present a network that supports developments and investments in life science. It links 20 partners with extensive industrial, scientific and entrepreneurial experience, covering the whole value chain, from chemical,

pharmaceutical and clinical development to management, intellectual property IP, financing, and commercial exit of projects.

The second one will contribute hands-on experience of a research-based drug development and information management consultancy, delivering leading-edge science that accelerates drug discovery and development. Development of algorithms, databases and software that incorporate physiological, genetic and epidemiological information are in the main focus.

Promoting results of biomedical research to develop new products and applications are the focus of the third presentation. It supports researchers in developing their research results commercially, providing such as project management, funding for patent protection, legal advice and business development, etc.

#### **Drug Product Quality: Needs and Performance**

An overview will be provided of recent changes in the pharmaceutical legislation („euroreference preparation” concept, tightening timeframes etc.) and of the new guidance documents issued by the ICH and/or the EMEA, which have an impact on the requirements for the content of quality documentation and on the principles and practice of quality assessment. The well detectable shift from analysing the end products to the more thorough understanding, design and control of the API and product manufacture and application of the concept in the daily routine of dossier compilation and assessment are major current issues of the standardisation of quality of medicines.

After this, we will summarise the legal status of EP monographs and general chapters in the current European pharmaceutical regulatory environment. Applicability and limitations of common standards for articles produced by different manufacturers, via different manufacturing techniques, will be illustrated by several examples in the presentation. Advantages and potential risks of inclusion of guidance documents in the pharmacopoea will also be discussed by the speaker.



# Career Centre: Are You Looking for New Career Opportunities?

During the Pharmaceutical Sciences World Congress, FIP is organising a Career Centre. So if you are looking for a new job, want to take your current career in a different direction, or just want to get into contact with representatives from leading pharmaceutical companies, we invite you to participate in this unique event.

Whatever changes or developments you are looking to make within your career, the FIP Career Centre during the PSWC will have something to offer you!

The goal of the Career Centre is to bring Ph.D. students, post doctoral fellows and those individuals already embarked on their career path into contact with representatives from leading pharmaceutical industries and other potential employers. During the FIP Career Centre, opportunities will be available to speak to these representatives, both for informal discussions Centreed around job opportunities, or for a pre-arranged interview for one of the positions posted on the FIP Career Centre website. New job

opportunities can and will also be posted throughout the Congress.

The Career Centre will be on-going throughout the days of the PSWC and will welcome candidates to speak with industry delegates privately in designated recruitment rooms at the Career Centre. Also company presentations will be held. For specific programme details please look on [www.fip.org/careerCenter](http://www.fip.org/careerCenter) or on the message board in the Forum Area.

Thanks are extended to the Founding Partners of this initiative, AstraZeneca and Pfizer.

For more information and free participation by uploading your CV, please visit:  
[www.fip.org/careerCenter](http://www.fip.org/careerCenter)

## Sponsors



## FIP Career Centre at PSWC2007

### What?

A Career Centre where participants are able to have informal discussions with company delegates or be invited for one of the pre-arranged job interviews.

### Who?

The Career Centre is for Ph.D. students, post doctoral fellows but also for individuals working in the pharmaceutical industry.

### How do I participate?

Step 1: Upload your CV on the FIP Career Centre website: [www.fip.org/careercenter](http://www.fip.org/careercenter)  
Step 2: Apply for one of the jobs posted or wait for an invitation for a general interview after posting your CV  
Step 3: Be invited by e-mail for a job interview or a more informal discussion in one of the private recruitment rooms

### When?

April 22 - 25 during the PSWC 2007, every day from 09:00 - 18:00 hours.

### Where?

Amsterdam RAI, at the Career Centre, Rooms F and G on the first floor (Auditorium Area).

### More information

Additional information during the congress can be found on the Message Board (Forum Area) or on the website [www.fip.org/careercenter](http://www.fip.org/careercenter)

# FIRST FLOOR

Symposia  
Short Presentations  
Round Table  
EUFEPS

A

Symposia  
Short Presentations  
Round Table  
EUFEPS

L

Forum Part, 1st floor

M... Press/Speakers  
N... Symposia  
N1... Short Presentations  
O1... Round Table  
O... EUFEPS

...Private Meetings

Auditorium Part, 1st floor

C... Symposia, Short Presentations,  
D... Round Table, EUFEPS  
E... Session Room  
F... Career Centre  
G...



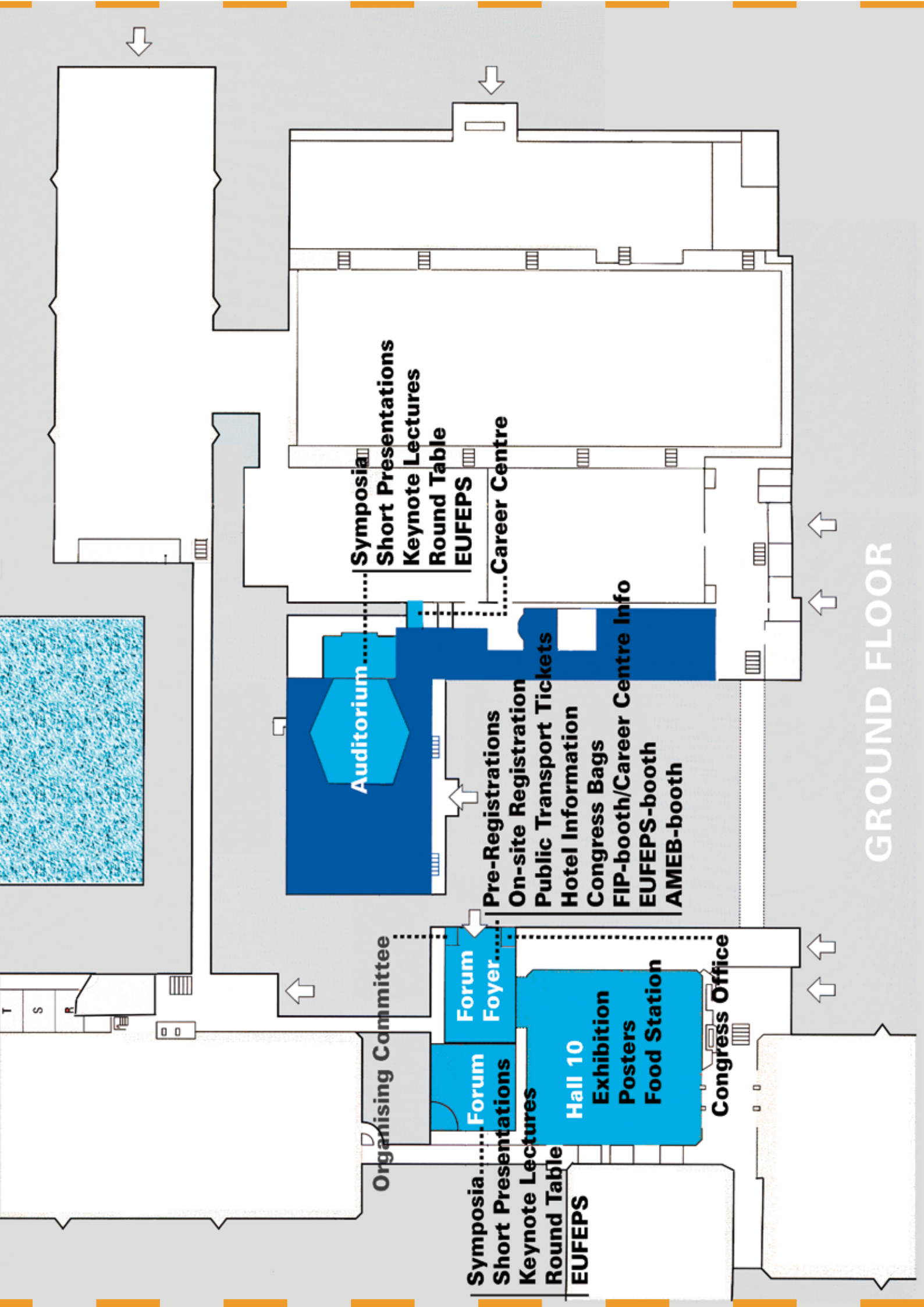
Meeting Rooms



Public Areas

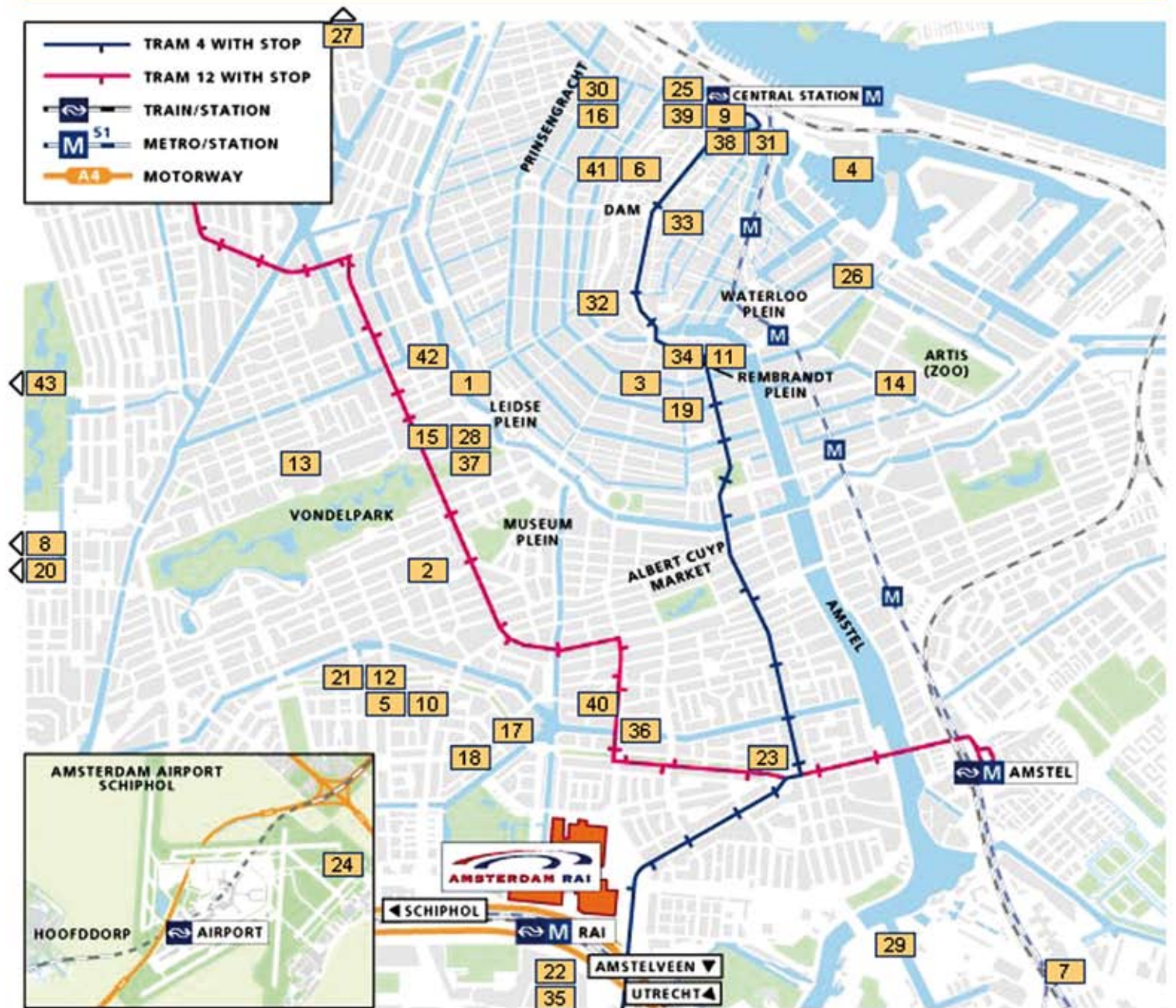


RAI Wireless Hotspot





## MAP OF AMSTERDAM



- |   |   |  |
|---|---|--|
| 1 <a href="#">ACH Leidse Square Hotel</a>                 | 16 <a href="#">Golden Tulip Amsterdam Centre</a>            | 30 <a href="#">New Amsterdam</a>                       |
| 2 <a href="#">ACH Trianon Hotel Amsterdam</a>             | 17 <a href="#">Golden Tulip Apollo Amsterdam</a>            | 31 <a href="#">NH Barbizon Palace</a>                  |
| 3 <a href="#">Albus Grand Hotel</a>                       | 18 <a href="#">Hampshire Hotel Beethoven</a>                | 32 <a href="#">NH City Centre</a>                      |
| 4 <a href="#">Amstel Hotel</a>                            | 19 <a href="#">Hampshire Inn - Prinsengracht</a>            | 33 <a href="#">NH Grand Hotel Krasnapolsky</a>         |
| 5 <a href="#">Apollofirst - A Hampshire Classic Hotel</a> | 20 <a href="#">HEM Hotel Amsterdam</a>                      | 34 <a href="#">NH Schiller</a>                         |
| 6 <a href="#">Avenue Hotel</a>                            | 21 <a href="#">Hilton Amsterdam</a>                         | 35 <a href="#">Novotel Amsterdam</a>                   |
| 7 <a href="#">Bastion Hotel Amsterdam / Amstel</a>        | 22 <a href="#">Holiday Inn Amsterdam</a>                    | 36 <a href="#">Okura Amsterdam</a>                     |
| 8 <a href="#">Bastion Hotel Amsterdam / Centrum-ZVV</a>   | 23 <a href="#">Hotel V</a>                                  | 37 <a href="#">Owl Hotel</a>                           |
| 9 <a href="#">Bellevue</a>                                | 24 <a href="#">Ibis Amsterdam Airport</a>                   | 38 <a href="#">Park Plaza Victoria Hotel Amsterdam</a> |
| 10 <a href="#">Best Western Delphi Hotel</a>              | 25 <a href="#">Ibis Amsterdam Centre hotel - restaurant</a> | 39 <a href="#">Ramada Amsterdam City Centre</a>        |
| 11 <a href="#">Best Western Eden Hotel</a>                | 26 <a href="#">Ibis Amsterdam City Stopera</a>              | 40 <a href="#">Savoy Hotel Amsterdam</a>               |
| 12 <a href="#">Bilderberg Garden Hotel</a>                | 27 <a href="#">Ibis Amsterdam Westcorner</a>                | 41 <a href="#">Sofitel Amsterdam</a>                   |
| 13 <a href="#">De Filosoof</a>                            | 28 <a href="#">Marriott Amsterdam</a>                       | 42 <a href="#">Tulip Inn Amsterdam Centre</a>          |
| 14 <a href="#">Eden Lancaster Hotel</a>                   | 29 <a href="#">Mercure aan de Amstel</a>                    | 43 <a href="#">Tulip Inn Amsterdam City West</a>       |
| 15 <a href="#">Europa 92 Hotel</a>                        |   |  |

The **GVB 72 hours ticket** is by far the easiest way of travelling. The ticket only needs to be stamped once, at the beginning of your first journey. After stamping, you can make unlimited use of GVB trams, buses, metro and night buses for a period of 72 hours (3 days).



# General Information

## Badges

All delegates and accompanying persons will receive a personal badge upon registration in the Forum Foyer of the RAI Exhibition and Congress Center as of Sunday 22 April, 2007. All participants and accompanying persons are kindly requested to wear their badges throughout the congress. Only participants wearing their name badge will be admitted to the sessions, exhibition in Hall 10, Opening Session and Welcome Reception. Accompanying persons wearing their name badge are welcome to the Exhibition Hall 10, Opening Session and Welcome Reception.

## Business Hours

Banks are open from 09:00-16:00 from Monday to Friday. Shops and Department Stores are generally open from 10:00-18:00.

## Congress Reception and Dinner

A Welcome Reception will take place on Sunday, April 22, 2007, after the Opening Ceremony in Hall 10 of the RAI Convention Center.

The Congress Dinner is planned on Tuesday, April 24, 2007 in the famous Wintergarden of the Grand Hotel Krasnapolsky, in the city centre. The event is optional and if there are still tickets available, you can purchase them at the on-site registration desk in the Forum Foyer.

## Credit cards

Hotels, Shops and Restaurants usually accept all credit cards.

## Climate

The congress will take place in April. Temperatures are between 15-20 degrees Celsius.

## Currency

The currency used in The Netherlands is EURO.

## VAT

VAT (value added tax) in the Netherlands is on most items 19%.

## Electrical Appliances

Electrical appliances in The Netherlands operate on 220 volts.

## Food Stations

From 11:15 until 14:00 hours there will be food stations in Hall 10.

## Housing Agent

RAI Hotel Services has a booth in the registration area of the Forum Foyer.

## Insurance

The Organising Committee accepts no liability for personal injuries sustained, sickness or for loss or damage to property belonging to congress participants and/or accompanying persons, incurred either during or as a result of the congress. It is recommended that each participant takes out a personal insurance.

## Internet Hot Spots

There are several Hot Spots in the Lobby of the Auditorium. There are computers, tables and chairs. Prices: Euro 7.50 for 1 hour - Euro 19.00 for 24 hours (can be used over various days and time slots). Credit card payment is Euro 0.20 per minute (minimum of 15 minutes).

## Language

The official congress language is English. No simultaneous interpretation will be provided.

## Public Transport

Train: Amsterdam RAI is easily accessible by train. Air travelers can make use of a direct train connection (4 times per hour). From the RAI station it is a 5 minutes walk to the RAI venue. Trains from/to Roosendaal/Belgium/France call at Schiphol. There are direct connections from Rotterdam, The Hague and Leiden. From Amsterdam Central Station you can take the Amstelveen express tram 51 (travelling time is 12 min.; exit at the RAI station) or tram 4 (travelling time is 30 min.; exit at RAI Europaplein).

## Presentation Equipment and Speakers Preview Room

All Session rooms are equipped with a laptop and LCD projection. All speakers are requested to proceed to the Speakers/Press Room: Room M 1st floor of the Forum



Part to check their presentation. A copy of each presentation will be asked by the technician for use in the Press Room and as general back up. Speakers are requested to hand over their memory stick to the technician in each session room, 30 min prior to the morning and afternoon sessions and collect it from the technician at the end of the session.

In your Registration envelope you will find a form on which speakers will be asked whether their (modified) presentations will be available for electronic dissemination upon request as a pdf file.

### Press Room

All press representatives are requested to register in the Pre-reg-

istration area of the Forum Foyer after which they should proceed to Room M 1st floor Forum Part.

### Tipping

It is not necessary to give tips in taxis or restaurants. Service is included in the bill.

However if you are very satisfied with the performance you could round up the bill.

### Accreditation by KNMP & NVFG

The Royal Dutch Pharmaceutical Society KNMP has accredited PSWC2007 as a continuing education course for community pharmacists (NL) for 0.5 credit point per half day (accreditation number CvD/1829). Participants who wish

to apply have to sign the presence list at the entrance of the session room. The Netherlands Association of Pharmaceutical Physicians NVFG has accredited it as well, with maximum 28 accreditation points ([www.nvfg.nl](http://www.nvfg.nl)).

### Registration

The registration area is open from Sunday, April 22, 2007, until Wednesday, April 25, 2007, in the Forum Foyer of the RAI Convention Centre.

PSWC delegates registration fee includes:

- Access to all scientific sessions
- Access to the exhibition
- Final Congress Programme
- List of Participants
- Congress Bag
- Name Badge
- Invitation to the Opening Session, Sunday, April 22, 2007
- Invitation to the Welcome Reception, Sunday, April 22, 2007

On-site delegates will receive (if applicable)

- Proof of payment on PSWC letterhead
- Poster certificate on PSWC letterhead
- Certificate of attendance on PSWC letterhead

PSWC Accompanying persons registration fee includes:

- Access to the exhibition
- Name Badge
- Invitation to the Opening Session, Sunday, April 22, 2007
- Invitation to the Welcome Reception, Sunday, April 22, 2007

### Payment

Registration fees are to be paid in Euro either in cash or by Credit card (Visa, Eurocard/Mastercard, American Express).

### Student registration

- For on-site registration, please bring with you an official document signed by the Head of Department of your University proving your Ph.D Student/Recent Postdoc graduate status (three years after graduation).



# Useful Addresses

## PSWC2007 Congress Registration & Abstract Handling

NewBrooklyn  
P.O. Box 73  
NL-3620 AB Breukelen  
The Netherlands  
Tel +31 346 266110  
Email  
registration@newbrooklyn.nl

## PSWC2007 Enquiries & Information

International Pharmaceutical Federation  
FIP Congress & Conferences  
Andries Bickerweg 5  
P.O. Box 84200  
NL-2508 AE The Hague  
The Netherlands  
Tel +31 70 3021982  
Fax +31 70 3021998  
Email pswc@fip.org  
www.fip.org/PSWC

## PSWC2007 Promotion & Website

Hans H. Linden  
EUFEPS  
P.O. Box 1136  
SE-11181 Stockholm  
Sweden  
Tel +46 8 7235025  
Fax + 46 8 4113217  
Email hans.linden@eufeps.org  
www.eufeps.org

## PSWC2007 Accommodation & Reservation

RAI Hotel & Travel Service  
P.O. Box 77777  
NL-1070 MS Amsterdam  
The Netherlands  
Tel +31 20 5491927  
Fax +31 20 5491946  
Email hotelservice@rai.nl  
www.rai.nl/hotelservice

## PSWC2007 & PharmSciFair Exhibition

Chris Hanney  
Health Links  
3rd Floor, Windsor House  
11A High Street  
Kings Heath  
Birmingham B14 7BB  
United Kingdom  
Tel +44 121 2483399  
Fax +44 121 2483390  
Email channey@health-links.co.uk  
www.health-links.co.uk



# Public Transport

5 STAR HOTELS	Distance to RAI	Tram to/from RAI - Bus to/from RAI
Bilderberg Garden - Mangerie de Kersentuin	5-10 min by Taxi	tram 5 (direc. Amstelveen Binnenhof) to station Zuid, change to metro 50 (direc. Gein) or metro 51 (direc. Central Station) to RAI
Hilton Amsterdam	5-10 min by Taxi	tram 5 (direc. Amstelveen Binnenhof) to station Zuid, change to metro 50 (direc. Gein) or metro 51 (direc. Central Station) to RAI
Marriott	City Center	tram 7 (direc. Flevopark) or tram 10 (direc. Azartplein) to Frederiksplein, change to tram 4 (direc. RAI) to RAI
NH Barbizon Palace	City Center	tram 4 (direc. RAI) to RAI
NH Grand Hotel Krasnapolsky Okura	City Center	tram 4 (direc. RAI) to RAI
	Walking distance	no tram
4 STAR HOTELS		
Apollofirst	5-10 min by Taxi	tram 5 (direc. Amstelveen Binnenhof) to station Zuid, change to metro 50 (direc. Gein) or metro 51 (direc. Central Station) to RAI
Best Western Delphi	5-10 min by Taxi	tram 5 (direc. Amstelveen Binnenhof) to station Zuid, change to metro 50 (direc. Gein) or metro 51 (direc. Central Station) to RAI
Golden Tulip Inntel Amsterdam Centre	City Center	tram 4 (direc. RAI) to RAI
Holiday Inn	Walking distance	
Le Meridien Apollo	5-10 min by Taxi	bus 15 (direc. Muiderpoort Station) to RAI
Mercure aan de Amstel	5-10 min by Taxi	metro 50 (direc. Isolatorweg) / metro 51 (direc. Amstelveen Westwijk) to RAI
NH Schiller	City Center	tram 4 (direc. RAI) to RAI
Novotel	Walking distance	walk
Savoy Hotel	Walking distance	bus 15 (direc. Muiderpoort) to RAI
Victoria	City Center	tram 4 (direc. RAI) to RAI
Softel	City Center	tram 4 (direc. RAI) to RAI
3 STAR HOTELS		
Albus Grand	City Center	tram 4 (direc. RAI) to RAI
Amstel Botel	City Center	tram 4 (direc. RAI) to RAI

Avenue	City Center	tram 4 (direc. RAI) to RAI
Bellevue	City Center	tram 4 (direc. RAI) to RAI
Best Western Beethoven	5-10 min by Taxi	tram 5 (direc. Amstelveen Binnenhof) to station Zuid, change to metro 50 (direc. Gein) or metro 51 (direc. Central Station) to RAI
Best Western Eden	City Center	tram 4 (direc. RAI) to RAI
Best Western Lancaster	City Center	tram 9 (direc. Centraal Station) or tram 14 (direc. Sloterveer) to Rembrandtplein, change to tram 4 (direc. RAI) to RAI
Bastion Hotel Amsterdam Amstel	5-10 min by Taxi	metro 51 (direc. Amstelveen Westwijk) to RAI
Bastion Hotel Amsterdam Zuidwest	Outskirts of Amsterdam	metro 50 (direc. Gein) to RAI
Europa 92	City Center	tram 12 (direc. Amstel station) to Victorieplein, change to tram 4 (direc. RAI) to RAI
Hampshire Inn Prinsengracht	City Center	tram 4 (direc. RAI) to RAI
Hotel de Filosoof	City Center	tram 12 (direc. Amstel station) to Victorieplein, change to tram 4 (direc. RAI) to RAI
Hotel V	Walking distance	tram 4 (direc. RAI) to RAI
Ibis Airport	Schiphol area	no tram or bus
Ibis Amsterdam Centre	City Center	tram 4 (direc. RAI) to RAI
Ibis Amsterdam Stopera	City Center	metro 51 (direc. Amstelveen Westwijk) to RAI
Ibis Amsterdam Westcorner	Outskirts of Amsterdam	metro 50 (direc. Gein) to RAI
Leidse Square	City Center	tram 7 (direc. Flevopark) or tram 10 (direc. Azartplein) to Frederiksplein, change to tram 4 (direc. RAI) to RAI
NH city centre	City Center	tram 4 (direc. RAI) to RAI
Owl Hotel	City Center	tram 12 (direc. Amstel station) to Victorieplein, change to tram 4 (direc. RAI) to RAI
Ramada city centre	City Center	tram 4 (direc. RAI) to RAI
Tulip Inn City West	Outskirts of Amsterdam	tram 17 (direc. Central Station) to station Lelylaan, from here metro 50 (direc. Gein)
Tulip Inn Amsterdam Centre	Walking distance	tram 4 (direc. RAI) to RAI



# Award Winners

## Ole J. Bjerrum



After finishing his medical internship, Ole J. Bjerrum (MD, DMSc) joined the Protein Laboratory at the Medical Faculty, University of Copenhagen as Assistant Professor (1970-1974), and as Associate Professor (1974-1987). In this period he served five years as director of the laboratory. Further to this, Ole J. Bjerrum has 14 years of industrial experience at Novo Nordisk, first as Director of Biolabs (a diagnostic unit), after that as senior principal scientist at Bioscience (working on various aspects of drug discovery, including HTS), and finally as liaison officer between the company and academia and national and international research organisations. In the period 1989-99 he was Adjunct professor in Immunotechnology at the Technical University of Denmark. From August 1, 2001, he is Professor of Pharmacology at the Danish University of Pharmaceutical Sciences, from 2007 on the University of Copenhagen Scientifically Ole J. Bjerrum has been engaged in the analysis and characterisation of membrane receptors and transporters, employing electrophoresis and immunotechnology tools (thesis 1977). Lately, his interest has been in *in vitro* and *in vivo* pharmacology aspects of chronic pain conditions. His publication list covers 3 books and more than 135 peer reviewed scientific papers.

Ole J. Bjerrum is a Fellow of the Danish Academies of Technical Sciences and Natural Sciences, respectively. He served as member of the Danish Medical Research Council and of the European Science and Technology Assembly, as well as of the EU Commission 4th and 5th Framework Programme Committee on Biotechnology and Quality of Life, respectively. In addition, he was co-founder of the Centre for Proteome Analysis, University of Southern Denmark. He was associated with the Centre as Adjunct

professor from 1999 - 2004.

In 1998, Ole J. Bjerrum joined the EufEps Committee on Industrial Relations (CIR). In 1999 he became member of the Executive Committee and served from 2000 as President-elect, from 2003 as President and from 2005 as Immediate past-president. He has taken an interest to promote the pharmaceutical sciences in Europe, through initiatives such as the formulation of the theme: New Safe Medicines Faster. An initiative which has had significant impact on the EU 6th Framework Programme for 2003-2007 and which paved the way for the Joint Technology Initiative: Innovative Medicines in the 7th Framework Programme.

## William N. Charman



Prof Bill Charman is Dean, Victorian College of Pharmacy, Monash University in Melbourne, Australia. He received his PhD in pharmaceutical chemistry from the University of Kansas in 1985, and from 1986-1989 was a Senior Scientist/Group Leader at the former Sterling-Winthrop Research Institute in Rennselear New York. He returned to Australia in 1989 where his research interests include enhanced absorption and bioavailability of poorly water soluble drugs, lymphatic drug transport, lead candidate optimisation, and the discovery and development of drugs for neglected diseases. He received the GlaxoWellcome International Achievement award in Pharmaceutical Sciences from the Royal Pharmaceutical Society of Great Britain in 1999, the Drug Discovery Project of the Year by the Medicines for Malaria Venture (Geneva, Switzerland) in 2002, the APSA medal in 2005, and the Controlled Release Society international career achievement in oral drug delivery in 2006. He has published over 330 scientific papers and communications, is a member of four international Edi-

torial Boards, and is an Associate Editor of the Journal of Pharmaceutical Sciences. He is an elected Fellow of the American Association of Pharmaceutical Scientists, a previous member of two Corporate Boards, a member of various Scientific Advisory Boards and is Chairman, Seeding Drug Discovery Funding Committee of the Wellcome Trust.

## Alfonso Domínguez-Gil Hurlé



Dr. Alfonso

Domínguez-Gil Hurlé was born in Gijón (Spain) in 1942. He received a fellowship from the Research Per-

sonnel Training Programme of the Spanish Ministry of Education (1968-1971), later receiving a PhD (with Extraordinary Prize). Since 1974 he has been Full professor of Pharmacy and Pharmaceutical Technology at the University of Salamanca (Spain). Currently, he is Professor of Biopharmacy and Clinical Pharmacokinetics. He was the Director of the Department of Pharmacy and Pharmaceutical Technology, Dean of the School of Pharmacy and Vice-Rector responsible for Investigation of the University of Salamanca (1980-1985). He has been a specialist in Hospital Pharmacy since 1975. Currently, he directs the Pharmaceutical Services of the University Teaching Hospital in Salamanca. He has been Director of the specialist courses entitled "Therapeutic Drug Monitoring" run since 1982, of which there have been 24 annual gatherings, attended by 1,800 pharmacists and physicians from Spain, France, Italy, Mexico, Chile, etc. He has delivered many specialist courses dealing with Clinical Pharmacokinetics in Spanish and non-Spanish Universities and at meetings of Spanish Scientific Societies: Hospital Pharmacy, Nephrology, Intensive Care, Chemotherapy, etc.

Dr. Domínguez-Gil Hurlé is the author of more than 300 publications concerning clinical



pharmacokinetics and pharmacoeconomy published in international journals. He has delivered more than 200 scientific contributions at Spanish and international meetings. He has participated in 32 clinical pharmacokinetic studies. He currently directs the Unit of Clinical Pharmacokinetics and Pharmaceutical Care for HIV-Positive Patients of the University Teaching Hospital in Salamanca. He is the co-author of the Spanish-English/English Spanish Terminological Dictionary of the Pharmaceutical Sciences published by the Royal Spanish Academy of Pharmacy (2007). He has authored 10 books addressing Hospital Pharmacy and Clinical Pharmacokinetics published in Spain, the United Kingdom and the United States. He has represented Spain at the European Pharmacopoeia of the European Council in Strasburg and at the International Pharmacopoeia in Geneva; he has been President of the Commission of the Royal Spanish Pharmacopoeia (200-2004) and a member of the Scientific Council the Spanish Drugs Agency. He has been Vice-President of the Spanish Society of Pharmacology and a member of the Spanish Agency for Quality Assessment of Universities and of the Spanish Teaching-staff Accreditation Agency in the area of the Health Sciences. He is a member of the Quality Assessment Committee of Pharmacy Schools in Portugal. Currently he is a member of the Quality Agency of the Universities of the Balearic Islands (Spain). Since 1980, he has been President of the Pharmacy and Therapeutics Commission of the University Teaching Hospital in Salamanca and Vice-President of the Ethical Committee for Clinical Research. Since 1998 he has been the Director of the Institute for Safe Medication Practices (Spain), a delegation of the ISMP in the United States. He has received many awards and distinctions, among them the Laude Award for Pharmaceutical Investigation (Spain) in 1974 and the Award of the American Society of Hospital Pharmacy Research and Education Foundation (1994). He represented Spain in the constitution of the Iberian-Latin-American

Association of Pharmacy Academies held in Valparaiso (Chile) in 2005. Dr. Domínguez-Gil Hurlé is a member of the Scientific Committee of the Spanish Foundation of Drug Sciences, and belongs to the following Academies: the Royal Spanish Academy of Pharmacy; The Royal Academy of Medicine of Salamanca, The Royal Spanish Academy of Pharmacy in Catalonia (Spain), and the Academy of Pharmacy of Galicia (Spain).

### Elias Fattal



Elias Fattal is a Full Professor of Pharmaceutical Technology at the University of Paris-XI in Châtenay-Malabry, France and has been President of APGI since 2003. He received his Pharmacy Degree (1983), and Ph.D. (1990) from the University of Paris-XI. After visiting the Department of Pharmaceutical Chemistry, University of California, San Francisco for a post-doctoral position with Frank Szoka (1990-1991), he became associate Professor (1992) and full Professor at the University of Paris-XI (2000). Elias Fattal is leading the research group "Drug targeting and delivery of poorly stable drugs" in the CNRS research unit UMR CNRS 8612. He is also vice-chair of this department. His research activity deals with the design of nano- and microtechnological approaches for the delivery of peptides/proteins and nucleic acids. His special expertise deals with oral administration of proteins and vaccines and the design of delivery systems for antisense oligonucleotides. Special attention was given in recent years to the ocular delivery of nucleic acids and the use of cyclodextrins as absorption enhancing agents. He is the author and co-author of around 135 publications and book chapters and 10 patents. In 1999, he received the Colloidal Drug Carrier Award (at the 5th Expert meeting on colloidal drug carriers, Berlin, Germany). Elias Fattal is the co-editor of the Journal of Drug Delivery Science and Technology, the European editor of the Journal of Biomedical Nanotech-

nology. He serves on the editorial board of several pharmaceutical journals (Journal of Pharmaceutical Sciences, European Journal of Pharmaceutical Sciences, American Journal of Drug Delivery, and Expert Opinion on Drug Delivery) and nanotechnology dedicated journals (NanoBiotechnology, International Journal of Nanomedicine).

### Kathleen M. Giacomini



Dr. Kathy Giacomini is Professor and Chair of Biopharmaceutical Sciences at the University of California, San Francisco. She received her Ph.D. in Pharmaceutics from the State University of New York at Buffalo and completed a post-doctoral fellowship at Stanford University. Dr. Giacomini is considered a leader in the field of pharmacogenomics of membrane transporters. She led the discovery of coding region variants of about 50 membrane transporters that play a role in drug response in ethnically diverse populations. Dr. Giacomini and her group functionally characterized over 100 transporter variants in cells, discovering both gain of function and loss of function variants that may lead to variation in drug response. She has received numerous awards for her research and teaching including the Dawson Award of the American Association of Colleges of Pharmacy and the Research Achievement Award in Drug Metabolism from the American Association of Pharmaceutical Scientists. In 2006, she was elected to the Institute of Medicine of the National Academies.

### Hans E. Junginger



H.E. Junginger, Ph.D. was Professor of Pharmaceutics and Head of the Division of Pharmaceutical Technology at the Leiden/Amsterdam Center for Drug Research, Leiden University, The Netherlands until 1 February 2004 when he took early retirement

Since 1 January 2004 he is a guest professor at the Faculty of Pharmaceutical Sciences, Naresuan University, Phitsanulok, in Thailand and since 2005 also a Visiting Professor at the National University of Singapore and at the Pharmacy Department of Ljubljana in Slovenia.

He qualified as a pharmacist in 1967, at the University of Munich, Germany. In 1971, he obtained his Ph.D. degree in Pharmaceutical Chemistry at the University of Saarbrücken, Germany. From 1972 to 1980 he worked as a researcher at the Technical University of Braunschweig, Germany to obtain his qualification as professor in Pharmaceutical Technology. From 1980 to 2004 he was the Head of the Department of Pharmaceutical Technology at the Leiden/Amsterdam Center for Drug Research, The Netherlands.

He has published more than 280 articles and 35 book chapters. He is the (co)inventor of 8 patents.

His main research areas included the development of novel controlled drug delivery systems (especially for peptide drugs) for the (trans)dermal and peroral routes, utilizing new (bioadhesive) polymers. Especially, multifunctional polymers as polyacrylates and chitosan derivatives have been identified to be safe and non-toxic penetration enhancers for hydrophilic drugs. Furthermore, they are excellent delivery systems for the nasal, pulmonary and oral route for peptides, protein and (DNA) vaccines. Combining superporous hydrogels or other expanding tablet systems with those multifunctional polymers make the oral absorption of peptides feasible.

Until now 52 Ph.D. students have graduated under his supervision and 25 post-docs from all over the world joined his department in Leiden.

Between 1986 until 1990 he was president of the International Association for Pharmaceutical Technology (APV) and in 1994/5 president of the Controlled Release Society.

He was the Scientific Secretary of the International Pharmaceutical Federation FIP (1995 – 2003)

and as such member of the Executive Committee of FIP.

He has received several major awards and three honorary doctorates (Ghent, Belgium in 1995, Potchefstroom University in South Africa in 2003 and London University, UK in 2004).

He is a frequently invited speaker at international conferences and a consultant to international pharmaceutical industries.

He loves traveling and as result of this his nickname is "Flying Dutchman". When at home he loves to play piano and to read criminal stories.

### Kevin Shakesheff



Kevin Shakesheff is Professor of Tissue Engineering at the University of Nottingham. He is a registered Pharmacist and trained within the NHS as part of his professional qualification in 1992. After obtaining a PhD in polymer science, he moved to the Massachusetts Institute of Technology in the mid 1990s to work on polymers used in tissue engineering. In 1997, he returned to the UK to take up an Engineering and Physical Sciences Research Council Advanced Fellowship. In recent years, he has been named as one of the World's top Young Innovators by MIT's Technology Review, won the 2004 Hanson Award and presented his work at the Royal Institution under the "Scientists for a New Century" series. He was co-founder of the Tissue and Cell Engineering Society and founder of RegenTec Ltd, a biotech company based in Nottingham and developing clinical products based on regenerative medicine.

Scientific highlights of recent work by Professor Shakesheff's team include the demonstration of formation of polymer & cell composites by a novel supercritical fluid processing route (Ginty et al, PNAS, 2006 in press), the use of plasma polymers to control the distribution of cells in 3D polymer scaffolds (Barry et al, Advanced Functional Materials, 15, 1134-1140, 2005), and the selective modification of cell responses

to ECM proteins on scaffolds (De Bank PA et al, Journal of Materials Chemistry 15, 2047-2055, 2005). Our novel scaffold systems are applied in a number of tissue engineering applications including liver (Thomas et al, Cell Tissues Organs 181, 67-79, 2005), skin (Horobin et al, Wound Repair and Regeneration 13, 422-433, 2005), nerve (Teare et al, Neuroreport, 15, 493-498, 2004) and bone (Yang et al, Journal of Bone and Mineral Research, 18, 47-57, 2003).

### Yuichi Sugiyama



Yuichi Sugiyama, Ph.D., born in 1947, is Professor and Chairman, Department of Biopharmaceutics at the University of Tokyo

since 1991. The Department name was recently changed to Molecular Pharmacokinetics. Except for a sabbatical at UCLA in 1979-1981 with Professor Kaplowitz, he has been at the University of Tokyo throughout his career, receiving a B.S. in Pharmacy in 1971, a Ph.D. in Pharmaceutical Sciences with Professor Hanano in 1978. He is a co-author of more than 480 publications in international journals as well as 270 book chapters and review articles (ca.60 written in English). His research focuses on two areas: 1) Physiologically based pharmacokinetics: prediction of drug dispositions from *in vitro* biochemical data; 2) Molecular pharmacokinetics of drug transport in liver, kidney, and brain.

Professor Sugiyama's research on membranes has yielded better understanding of the basic aspects of transport mechanisms. He has discovered several examples in which transporters play a major role in drug disposition by integrating *in vitro* data with *in vivo* pharmacokinetic models. Moreover, his work has highlighted the importance of considering pharmacokinetic properties in drug development, using screening methods to test large numbers of drug candidates. Detailed *in vitro* studies of transporters for the first time appear to predict transporter-mediated drug-drug interaction *in vivo*. Analysis of genetic poly-

morphisms in transporter genes are being identified in his laboratory that can account for inter-individual differences in drug disposition. Overall, he has produced a body of scientific work with an impact on our understanding of how drugs work and how to use them.

His work is internationally recognized receiving several awards, including the Ebert Prize of the American Association of Pharmaceutical Scientists (AAPS) in 1985; Takeru-Aya Higuchi Prize in 1990; Pharmaceutical Scientist of the Year Award of the International Pharmaceutical Federation (FIP) in 1994; the Scientific Achievement Award from the Academy of Pharmaceutical Science and Technology, Japan (APSTJ) in 1995; the Scientific Achievement Award from the "Japanese Society for Xenobiotic Metabolism and Disposition (JSSX)" in 2001.; The Troy C. Daniels Lectureship from UCSF in 2001; the AAPS Distinguished Pharmaceutical Scientist Award in 2003; the Scientific Achievement Award 2004 from the "Pharmaceutical Society of Japan(PSJ)" and the John G. Wagner Pfizer Lectureship Award in Pharmaceutical Sciences from University of Michigan in 2005. According to the information on the website of ISI Essential Science Indicators (ESI), Thomson Scientific<sup>®</sup>USA<sup>®</sup>, Prof. Sugiyama achieved the 2nd top position for the number of citations for the last 10 years (Jan 1, 1995 - Aug 31, 2005) in the field of "Pharmacology & Toxicology". He is currently the president of both the "International Society for the Study of Xenobiotics (ISSX)" and the "Japanese Society for Xenobiotic Metabolism and Disposition (JSSX)".

He is/was an editorial board member of several international journals, especially editor in Japan of "Pharmaceutical Research" (1992-1996) and "AAPS Pharm. Sci." (1999-2001). He served as the chairman of the Board of Pharmaceutical Sciences of FIP (2000-2004) as the successor of Dr. Leslie Benet. As a program co-chair, he was a main contributor to the success of the Millennial World Congress of Pharmaceutical Sciences, San Francisco 2000, and further, he chaired the "Phar-

maceutical Sciences World Conference", Kyoto, Japan, in 2004. (organized by FIP Board of Pharmaceutical Sciences). He was the President of the Academy of Pharmaceutical Science and Technology, Japan (APSTJ) (2002-2003).

### Hiroshi Terada



Dr. Hiroshi Terada, Professor at Tokyo University of Science, is an outstanding scientist in the fields of pharmaceutical sciences and biochemistry. Recently, he took an interest in studying controlled drug delivery formulations, especially those for overcoming infectious diseases.

For a long time, prof. Terada has been a Council Member of FIP, representing the PSJ. He was a co-chair of the Pharmaceutical Sciences World Congress (PSWC2004) in Kyoto. He made every effort to promote PSJ to become a member of FIP when he was a board member of PSJ, and then consistently supported the FIP activity when he was President of PSJ.

### Vladimir P. Torchilin



Vladimir P. Torchilin, Ph.D., D.Sc. is a Distinguished Professor and Chair of the Department of Pharmaceutical Sciences and Director, Center for Pharmaceutical Biotechnology and Nanomedicine, Northeastern University, Boston, Mass. He graduated from the Moscow State University with a MS in Chemistry, and also obtained there his Ph.D. and D.Sc. in Polymer Chemistry, Chemical Kinetics and Catalysis, and Chemistry of Physiologically Active Compounds in 1971 and 1980, respectively. In 1991 Dr. Torchilin joined the Massachusetts General Hospital and Harvard Medical School as the Head of Chemistry Program, Center for Imaging and Pharmaceutical Research, and Associate Professor of Radiology. Since 1998 Dr. Torchilin is with Northeastern

University. His research interests have focused on biomedical polymers, polymeric drugs, immobilized medicinal enzymes, drug delivery and targeting, pharmaceutical nanocarriers for diagnostic and therapeutic agents, and experimental cancer immunology. He has published more than 300 original papers, more than 100 reviews and book chapters, wrote and edited 10 books, made over 250 invited lectures and seminars and holds more than 40 patents. He served on multiple NIH Study Sections and is on the Editorial Boards of many leading journals including Journal of Controlled Release (Review Editor), Bioconjugate Chemistry, Advanced Drug Delivery Reviews, European Journal of Pharmaceutics and Biopharmaceutics, Journal of Drug Targeting, Molecular Pharmaceutics, Journal of Biomedical Nanotechnology, and many others. Among his many awards, Professor Torchilin was the recipient of the 1982 Lenin Prize in Science and Technology (the highest scientific award in the former USSR).

He was elected as a Member of the Russian Academy of Biotechnology and the European Academy of Sciences. He is also a Fellow of the American Institute of Medical and Biological Engineering and of the American Association of Pharmaceutical Sciences and received the 2005 Research Achievements in Pharmaceutics and Drug Delivery Award from the AAPS. He is on the Board of Directors of the International Liposome Society, and in 2005-2006 he served as a President of the Controlled Release Society.



# Monoclonal Antibodies

April 26-27 • 2007 • Amsterdam • The Netherlands

## Scope and Aim

Monoclonal antibodies are one of the most important classes of therapeutic recombinant proteins. They are under clinical evaluation for a broad range of important therapeutic areas including cancer, rheumatoid arthritis and infectious diseases. Classical Biotech companies are no longer the only players and traditional large Pharma companies have now also recognised the potential of recombinantly manufactured monoclonal antibodies.

This Workshop will cover a broad spectrum of topics from the design and engineering of monoclonal antibodies, through process development (e.g. cell line optimisation and down stream processing), analytics, formulation aspects and manufacturing up to clinical applications. An important element of the Workshop will be the attention paid to current regulatory requirements for the introduction of monoclonal antibodies. Trends towards modification of antibodies by conjugation or subsequent glycosylation as well as their reduction in size to antibody fragments such as domain or single chain antibodies will also be discussed.

## Location

Hotel Park Plaza Victoria Amsterdam, Damrak 1-5, 1012 LG Amsterdam.

## Additional Information

Consult the PSWC2007 Website, the EAPB Website or the EUFEPS Online.



## Scientific Planning Committee

Karoline Bechtold-Peters,  
Boehringer Ingelheim, Germany,  
Co-chair

Wim Jiskoot, Leiden/Amsterdam Center for Drug Research (LACDR), Co-chair

Daan J.A. Crommelin,  
Dutch Top Institute Pharma,  
The Netherlands

Barry Moore, XstaltBio,  
United Kingdom

Wayne Gombotz, Omeros Corp.,  
USA

Jan van de Winkel, Genmab,  
The Netherlands

Sven Stegemann, Capsugel,  
Belgium

## Registration

For Registration on line access the Workshop web site of the EUFEPS Online at: [www.eufeps.org](http://www.eufeps.org) or report to the EUFEPS booth at the PSWC2007.

## EAPB [www.eapb.org](http://www.eapb.org)

EAPB is a professional association dedicated to the advancement of biotechnology in pharmaceutical sciences, specifically as applied to industrial materials, processes, products and their associated challenges. Its members constitute scientists employed in industry, government and university laboratories, biotech companies and scientific organisations.

## EUFEPS [www.eufeps.org](http://www.eufeps.org)

Founded in 1991, the mission of EUFEPS is to advance excellence in the pharmaceutical sciences and innovative drug research, and to represent the interests of scientists engaged in drug research and development, drug regulation and drug policymaking. Currently, EUFEPS links 24 Member Societies in 24 European countries.

## AAPS [www.aaps.org](http://www.aaps.org)

AAPS is a professional scientific society of more than 12,000 members employed in academia, industry, government and other research institutes worldwide. Founded in 1986, AAPS provides a dynamic international forum for the exchange of knowledge among scientists to enhance their contributions to public health.

## FIP [www.fip.org](http://www.fip.org)

FIP, founded in The Hague, more than 90 years ago, is the worldwide federation of national pharmaceutical, professional and scientific associations, with a mission to represent and serve pharmacy and pharmaceutical sciences around the globe. Through its member association, FIP connects, represents and serves more than a million pharmacists and pharmaceutical scientists around the world.

## Organisers and Co-sponsors

Organisers of this PSWC 2007 Post-Satellite are the European Association of Pharma Biotechnology (EAPB) and the European Federation for Pharmaceutical Sciences (EUFEPS), and it is co-sponsored by the American Association of Pharmaceutical Scientists (AAPS) and the International Pharmaceutical Federation (FIP).





## Final Programme

### Thursday • April 26, 2007

Welcome, Introduction and Opening Remarks
Keynote Presentation: Past, Present and Future of Antibody Therapeutics <i>Sir Gregory Winter</i> , University of Cambridge, UK
<b>Session I: New technologies for design of antibodies and engineering</b> Chairman: <i>Jan GJ van de Winkel</i> , Genmab, NL
Glycoengineered therapeutic antibodies with increased FcγRIII binding affinity and enhanced biological activity <i>Peter Brunker</i> , F Hoffmann - La Roche, Schlieren - Zürich, CH
Engineering antibody effector function <i>Carl Webster</i> , Cambridge Antibody Technology, UK
Generating novel immuno-therapeutics for small bioactive compounds (haptens) <i>Andy Porter</i> , Haptogen, UK
<b>Session II: Latest advancements in upstream development including expression systems and feed strategies</b> Chairman: <i>Sven Stegemann</i> , Capsugel, BE
Manufacturing therapeutic monoclonal antibodies in chicken eggs <i>Marie-Cecile van de Lavoie</i> , Origen Therapeutics, Burlingame CA USA
Qualitative and quantitative comparison of protein expression systems for the manufacturing of antibodies <i>Rainer Fischer</i> , Fraunhofer-Institut für Molekularbiologie und Angewandte Oekologie, DE
Cell culture points-to-consider: A commercial perspective <i>Ben Bulthuis</i> , Centocor, NL
<b>Session III: Downstream development including purification and recovery optimisation</b> Chairman: <i>Karoline Bechtold-Peters</i> , Boehringer-Ingelheim, DE
Economic considerations for disposable technologies in antibody manufacturing <i>Uwe Gottschalk</i> , Sartorius, DE
Challenges in the development of economic and robust downstream processes for therapeutic antibodies using platform technologies <i>Alexander Jacobi</i> , Boehringer Ingelheim Pharma GmbH & Co. KG, DE
Controlled freeze-thaw technology <i>Gaël Péron</i> , Stedim Biosystems, FR
<b>Session IV: Physical characterisation, formulation and delivery systems</b> Chairmen: <i>Barry Moore</i> , XstalBio, UK, and <i>Wim Jiskoot</i> , Leiden/Amsterdam Center for drug Research (LACDR), NL
Rapid physical characterisation tools for biopharmaceuticals as used in the early development phase <i>Patrick Garidel</i> , Boehringer-Ingelheim, DE
High concentration processing and formulation of Mabs – New and state-of-the-art approaches <i>Wolfgang Friess</i> , University of Munich, DE
Novel dry powder antibody formulations <i>Jan Vos</i> , XstalBio, UK
Analytical challenges addressed <i>Tudor Arvinte</i> , University of Geneva, CH
<b>Session V: Analytics and specs</b> Chairman: <i>Daan JA Crommelin</i> , Top Institute Pharma, NL
Setting specifications for Mabs - Regulatory perspective <i>Bernd Liedert</i> , Paul Erlich Institute, DE
Application of analytical ultracentrifugation as an orthogonal method for protein size distribution analysis <i>James Andya</i> , Genentech Inc., South San Francisco CA USA
Aggregates in biotech products - Regulatory expectations for aggregates <i>Elizabeth Leininger</i> , StemCells Inc, Palo Alto CA USA
End of day I and panel discussion

### Friday • April 27, 2007

<b>Session VI: Scale-up issues and production scale manufacturing</b> Chairman: <i>Wayne Gombotz</i> , Omeros Corporation, Seattle WA USA
How to make successful technology transfers in Mabs production <i>Tim Clayton</i> , Laboratoires Serono SA, CH
Integrated downstream process design in Mabs production – The white knight to slay the cost dragon? <i>Jochen Strube</i> , Clausthal University of Technology, DE
Production of human monoclonal antibodies on the human cell-line PER.C6 <i>Erik Hack</i> , Crucell, NL
<b>Session VII: Clinical update (new antibody therapies), commercial impact and new trends</b> Chairman: <i>Jan GJ van de Winkel</i> , Genmab, NL
Anti-IL12p40 antibody development and clinical data <i>Trudi Veldman</i> , Abbott, USA
Ofatumumab, a novel human CD20 antibody for treatment of lymphoid malignancies and inflammation <i>Jan GJ van de Winkel</i> , Genmab, NL
Commercial impact and potential of Mab products and Mab production <i>Andreas Werner</i> , Boehringer Ingelheim, DE
The next generation of recombinant immunotoxins <i>Stefan Barth</i> , Fraunhofer-Institut IME-MB, DE
<b>Session VIII: Pharmacokinetics and safety</b> Chairman: <i>Wim Jiskoot</i> , Leiden/Amsterdam Center for Drug Research (LACDR), NL
Clinical consequences of development of antibodies to therapeutic monoclonal antibodies <i>Lucien Aarden</i> , Sanquin, NL
Global immunogenicity assessment of antibody therapeutics <i>Philippe Stas</i> , Algonomics, BE
Safety assessment of monoclonal antibody products – Perspective on preclinical requirements to support the determination of safe use conditions <i>James D Green</i> , Biogen Idec Inc, Cambridge MA USA
<b>Session IX: Armed antibodies and new classes of antibodies or antibody derivatives</b> Chairman: <i>Wayne Gombotz</i> , Omeros Corp., USA
Conjugated antibodies – Overview and case study on Mylotarg <i>Boris Gorovits</i> , Wyeth, USA
UniBody: A novel human antibody-based platform for immunotherapy <i>Paul Parren</i> , Genmab, NL
Potent immunoconjugates for cancer therapy <i>Peter Senter</i> , Seattle Genetics, USA
Discussion Forum: Future of antibody therapeutics/what are the next steps? (Prepared questions, participants to forward questions on cards, panel discussion)

### Sponsors



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# Young Pharmaceutical Scientists Meet in Amsterdam



## Scope and Aim

The Pharmaceutical Sciences World Congress 2007 is pleased to offer a pre-satellite symposium dedicated to the next generation of pharmaceutical scientists. Doctorate students and postdoctorate fellows from all over the world convene in Amsterdam to exchange ideas and discuss the latest developments in pharmaceutical sciences.

During the one and a half day event, young scientists present lectures in parallel sessions on such topics as: Target Discovery/Medicinal Chemistry, Pharmacokinetics / Pharmacodynamics, Drug Delivery, Pharmaceutical Analysis / Bioanalysis / Quality Assurance / Regulatory Affairs, Clinical Pharmacy / Pharmacoepidemiology and Toxicology / Safety. In addition, poster presentations are held on the latest research findings in all fields of pharmaceutical sciences. The students/postdoctoral fellows have the opportunity to present their work during "poster walks".

In each scientific section, the most outstanding poster presentation is honored with a "Young Investigator Award". Keynote lecturers in all areas give comprehensive overviews on the current developments in their particular fields. This pre-satellite meeting gives

young pharmaceutical scientists the unique opportunity to network and share professional experiences before the start of the 3rd Pharmaceutical Sciences World Congress on April 22, 2007.

## Location

University of Amsterdam (*Vrije Universiteit Amsterdam*) De Boelelaan 1105, NL-1081 HV Amsterdam, The Netherlands.

## Social Programme

**Saturday afternoon:** Amsterdam tour by foot and boat

**Saturday night:** Pre-satellite party  
Boom Chicago Show, including 3 course dinner and drinks

## Organising Committee

*S.C. De Smedt (Chair); P. Augustijns; J. Commandeur; D.J.A. Crommelin; M. Danhof; G.J. De Jong; S. Deferme; J. Demeester; W. Hennink; U. Holzgrabe; O. Khungel; C.M. Lehr; H.H. Linden; E. Mastrobattista; R. Schiffelers; F. Siepmann; J. Siepmann; N.P.E. Vermeulen; and B.H. Westerink*



**Vrije Universiteit Amsterdam, Amsterdam, The Netherlands**  
**April 20 and 21, 2007**

**PROGRAMME**

<b>Friday, April 20, 2007</b>	
09:00 - 09:15	<b>Introduction</b>
09:15 - 09:40	<b>Key note lecture I</b> Pharmacogenetics and regulation of human cytochromes ( <i>Dr. C. Rodriguez-Antona</i> )
09:40 - 10:05	<b>Key note lecture II</b> Non-viral gene delivery ( <i>Dr. N. Kobayashi</i> )
10:05 - 10:30	<b>Key note lecture III</b> HIV entry inhibitors: ligand-based design of peptidomimetic CXCR4 antagonists ( <i>Dr. J. Vabeno</i> )
10:30 - 11:00	Coffee break
11:00 - 12:30	<b>Oral session I</b> <ul style="list-style-type: none"> <li>■ TD/MC</li> <li>■ DM/DT/Tox-Saf</li> <li>■ DD Techno</li> <li>■ PA/QA/RA</li> <li>■ CP/PK-PD/PE</li> <li>■ DD Bio</li> </ul>
12:30 - 14:00	Lunch and poster viewing
14:00 - 15:30	<b>Oral session II</b> <ul style="list-style-type: none"> <li>■ TD/MC</li> <li>■ DM/DT/Tox-Saf</li> <li>■ DD Techno (1)</li> <li>■ DD Techno (2)</li> <li>■ CP/PK-PD/PE</li> <li>■ DD Bio</li> </ul>
15:30 - 16:00	Coffee break
16:00 - 17:00	<b>Oral session III</b> <ul style="list-style-type: none"> <li>■ TD/MC</li> <li>■ DD Techno</li> <li>■ PA/QA/RA</li> <li>■ CP/PK-PD/PE (1)</li> <li>■ CP/PK-PD/PE (2)</li> <li>■ DD Bio</li> </ul>
17:00 - 19:00	Poster walk - wine & cheese buffet

<b>Saturday, April 21, 2007</b>	
09:15 - 09:40	<b>Key note lecture IV</b> Peptide mapping of biofluids by multidimensional LC/MALDI-TOF MS ( <i>Dr. E. Machtejevas</i> )
09:40 - 10:05	<b>Key note lecture V</b> Gene expression profiling and breast cancer care ( <i>Dr. N. Oestreicher</i> )
10:05 - 10:30	<b>Key note lecture VI</b> Mechanism-based PK-PD modelling of drug efficacy and safety: application to (semi)-synthetic opioids ( <i>Dr. A. Yassan</i> )
10:30 - 10:45	Poster and Galenos award ceremony
10:45 - 11:15	Coffee break
11:15 - 12:45	<b>Oral session IV</b> <ul style="list-style-type: none"> <li>■ TD/MC</li> <li>■ DM/DT/Tox-Saf</li> <li>■ DD Techno</li> <li>■ CP/PK-PD/PE</li> <li>■ DD Bio</li> </ul>
12:45 - 14:00	Closing of the scientific programme / lunch
14:00 - 16:00	<b>Interactive Author Seminar</b> How to write and submit a world-class paper

TD/MC: Target discovery/Medicinal chemistry; DM/DT/Tox-Saf: Drug metabolism/Drug transport/Toxicology-Safety; DD Techno: Drug delivery-Technology; PA/QA/RA: Pharmaceutical analysis/Quality assurance/Regulatory affairs; CP/PK-PD/PE: Clinical pharmacy/Pharmacokinetics-Pharmacodynamics/Pharmacoepidemiology; DD Bio: Drug delivery-Biology.



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Visit our website: <http://pharmacie.univ-lille2.fr/presatellitePSWC>



# Pharmaceutical Curriculum Development: An evolutionary or a revolutionary process?

Saturday, April 21, 2007

Department of Pharmaceutical Sciences, Faculty of Science, Sorbonnelaan 16, Utrecht

## Education Symposium

Over the past several years, pharmacy curricula across the globe have come under the influence of many developments. The explosion of biomedical and pharmaceutical knowledge brought with the unravelling of the human genome, coupled with greater societal needs and expectations placed on pharmacists by an increasingly complex healthcare system, has required changes in the way pharmacists are trained and the skills they bring to the healthcare circle. Competency based education with an emphasis on patient-oriented practice and accountability are the current and future trends in pharmacy education.

Worldwide pharmacy schools face a tremendous challenge in rapidly adapting to these exciting yet sometimes conflicting developments. The intention of the PSWC Education Symposium is to bring together pharmaceutical scientists active in academia to discuss these challenges, possible solutions, and of course the biggest dilemma: How to promote state-of-the-art pharmaceutical sciences in curricula when at the same time there is a great societal demand to shift from product-oriented to patient-oriented pharmacy education and practice.

The programme will consist of plenary lectures and

round table discussions, offering ample opportunity for interaction. In addition, participants are invited to join in a guided tour of the Dept. of Pharmaceutical Sciences in Utrecht, a Department which has recently seen a radical change in its curriculum and organisational structure focussing on problem-oriented and student centred educational methods, in order to optimally prepare future practitioners and scientists.

## For whom?

Academic colleagues who are interested in pharmaceutical curriculum development in particular for 6 years programmes.

## How do delegates get there?

Bus transport to Utrecht will leave at 9:00 hrs sharp! in front of: Hotel Novotel Amsterdam, Europaboulevard 10, Amsterdam.

A hostess/student with the sign "Workshop: Pharmaceutical Curriculum Development: an evolutionary or a revolutionary process?" will be present on the spot to guide the delegates.

After the satellite, there is bus transport back to Hotel Novotel in Amsterdam.

## Programme

Welcome by <i>Bert Leufkens</i> , Dean of the Department of Pharmaceutical Sciences
<i>Anthonius de Boer</i> and <i>Andries Koster</i> Department of Pharmaceutical Sciences, Utrecht, the Netherlands <i>Radical changes of a pharmacy curriculum</i>
<i>Tetsumi Irie</i> Faculty of Medical and Pharmaceutical Sciences, Kumamoto, Japan <i>New six-year pharmacy curricula in Japan</i>
<i>Fe-Lin Lin Wu</i> School of Pharmacy, Taipei, China Taiwan <i>Evolution of pharmacy education in Taiwan</i>
Lunch and guided tour through Education Center of the Department of Pharmaceutical Sciences
<i>Kyenghee Kwon</i> College of Pharmacy, Seoul, Korea <i>The new era of the pharmacy education in Korea: How could we move from subject orientation to professional goals?</i>
<i>Ian Bates</i> School of Pharmacy, London, United Kingdom <i>Is it possible to maintain the science base whilst increasing a patient orientation in pharmacy curricula?</i>
Round Table Discussions





# Microdialysis: The Target Site in Focus

April 26-28, 2007, Leiden, The Netherlands

Prediction of drug target site distribution and effects is of utmost importance for successful drug development. In the last decades a number of useful *in vivo* monitoring techniques have become available, like PET, NMR and microdialysis. The unique characteristic of *in vivo* microdialysis is that it provides specific information on the extracellular tissue space, which represents the target site of many drugs, but also is the space in which the biochemical events may serve as biomarkers of the drug effects or on a pathological process. This makes that *in vivo* microdialysis has gained a special position within Drug Research and Development. Its potential is continuously growing by gain of insight in microdialysis experimentation, together with the improvement of analytical methodologies able to deal with the typical small-volume-low-concentration samples.

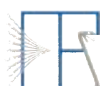
This Symposium will deal with discussions and exchange of knowledge on the latest developments of the role and potential of *in vivo* microdialysis complementary to other techniques and approaches to increase drug candidate selection efficiency, on the basis of the following workshops:

- Methodological Advances and Considerations in Monitoring the Extracellular Space
- Preclinical and Clinical Pharmacokinetics and Target Site Distribution
- Pharmacokinetic – Pharmacodynamic Correlations
- Monitoring Biomarkers and Drug Penetration in Disease Conditions

## Organizing Committee:

- Elizabeth CM de Lange, Chair
- Meindert Danhof
- Martha van der ham
- Erik de Vries

More information  
is available at  
[www.lacdr.nl](http://www.lacdr.nl)



Leiden / Amsterdam  
Center for Drug Research  
Leiden University/Vrije Universiteit Amsterdam



# Poster Overview and Abstract Listing

Poster Overview during PSWC2007 in Hall 10, RAI Amsterdam			
<b>Monday, April 23, 2007</b>	<b>317 posters</b>		<b>Please, note!</b>
CP-M-001	CP-M-013	PSWC: Clinical Pharmacology and Biomarkers	07:30-08:30 hang up poster 11:15-12:15 author at board 17:00- 18:00 take down posters 18:00-19:00 change numbers
DMT-M-001	DMT-M-031	PSWC: Drug Metabolism and Transport	
DD-M-001	DD-M-157	PSWC: Formulation, Delivery, Biopharmaceutics and Pharmaceutical Technology	
MC-M-001	MC-M-043	PSWC: Medicinal Chemistry and Natural Products	
PA-M-001	PA-M-020	PSWC: Pharmaceutical Analysis, Bioanalysis, Quality Assurance/Control and Regulatory Affairs	
PE-M-001	PE-M-015	PSWC: Pharmacoepidemiology and Pharmacovigilance	
KD-M-001	KD-M-025	PSWC: Pharmacokinetics and Pharmacodynamics	
KG-M-001	KG-M-014	PSWC: Pharmacokinetics and Pharmacogenetics	
<b>Tuesday, April 24, 2007</b>	<b>312 posters</b>		<b>Please, note!</b>
CP-T-001	CP-T-012	PSWC: Clinical Pharmacology and Biomarkers	07:30-08:30 hang up poster 11:15-12:15 author at board 17:00- 18:00 take down posters 18:00-19:00 change numbers
DMT-T-001	DMT-T-028	PSWC: Drug Metabolism and Transport	
DD-T-001	DD-T-152	PSWC: Formulation, Delivery, Biopharmaceutics and Pharmaceutical Technology	
MC-T-001	MC-T-035	PSWC: Medicinal Chemistry and Natural Products	
PA-T-001	PA-T-020	PSWC: Pharmaceutical Analysis, Bioanalysis, Quality Assurance/Control and Regulatory Affairs	
PE-T-001	PE-T-015	PSWC: Pharmacoepidemiology and Pharmacovigilance	
KD-T-001	KD-T-025	PSWC: Pharmacokinetics and Pharmacodynamics	
TD-T-001	TD-T-026	PSWC: Target Discovery and Molecular Pharmacology	
<b>Wednesday, April 25, 2007</b>	<b>323 posters</b>		<b>Please, note!</b>
CP-W-001	CP-W-012	PSWC: Clinical Pharmacology and Biomarkers	07:30-08:30 hang up poster 11:15-12:15 author at board 14:00-14:30 take down posters
DMT-W-001	DMT-W-028	PSWC: Drug Metabolism and Transport	
DD-W-001	DD-W-152	PSWC: Formulation, Delivery, Biopharmaceutics and Pharmaceutical Technology	
MC-W-001	MC-W-035	PSWC: Medicinal Chemistry and Natural Products	
PA-W-001	PA-W-021	PSWC: Pharmaceutical Analysis, Bioanalysis, Quality Assurance/Control and Regulatory Affairs	
PE-W-001	PE-W-014	PSWC: Pharmacoepidemiology and Pharmacovigilance	
KD-W-001	KD-W-029	PSWC: Pharmacokinetics and Pharmacodynamics	
TOX-W-001	TOX-W-035	PSWC: Toxicology and Safety Sciences	

**All abstract texts can be found on the CD that has been inserted in your Congress Bag**



Monday  
April 23, 2007

## Clinical Pharmacology and Biomarkers

### CP-M-001

A SIMPLE METHOD FOR PREVENTION OF ADVERSE EFFECTS IN DOUBLE FILTRATION PLASMAPHERESIS (DFPP) Shinji Abe, Mayumi Torii, Yukie Shimooka, Toshihide Kujime, Kazuhiko Teraoka, Kazuyoshi Kawazoe, Kazuo Minakuchi

### CP-M-002

SELECTIVITY OF N-HYDROXYSUCCINIMIDE ESTER-MEDIATED PEPTIDE ACYLATION: APPLICATION FOR THE LABELING OF NITROTROSINE-CONTAINING PEPTIDES

Nicolas Abello, M. Begona Barroso, Huib A.M. Kerstjens, Dirkje S. Postma, Rainer Bischoff

### CP-M-003

BARRIERS TO THE IMPLEMENTATION OF PHARMACEUTICAL CARE IN PSYCHIATRY HOSPITAL IN K.S.A Khalaf Al-Jumaah, Jawza AL-Sabhan, Zeinab Jaglit

### CP-M-004

AN ORAL ADSORBENT, AST-120 PROTECTS AGAINST THE PROGRESSION OF OXIDATIVE STRESS BY REDUCING THE ACCUMULATION OF INDOXYL SULFATE IN THE SYSTEMIC CIRCULATION IN RENAL FAILURE Makoto Anraku, Kazuki Shimoishi, Yuka Tasaki, Toru Maruyama, Masaki Otagiri

### CP-M-005

ACHILLEA SANTOLINA REDUCED OXIDATIVE STRESS IN THE LIVER OF STREPTOZOTOCIN-INDUCED DIABETIC RATS Amin Ardestani, Razieh Yazdanparast

### CP-M-006

IDENTIFYING NONRESPONDERS TO ASTHMA AND COPD THERAPY USING LEUKOCYTE PARAMETERS Madelon Bracke, Karin Velthove, Madelon Bracke, René Schweizer, Patrick Souverein, Maarten Ten Berg, Bert Leufkens, Wouter Van Solinge

### CP-M-007

A MODEL-BASED APPROACH TO BRIDGING PHARMACOKINETICS FROM ADULTS TO CHILDREN. Massimo Cella, Gijs Santen, Meindert Danhof, Oscar Della Pasqua

### CP-M-008

FEMALE HORMONE SECRETION OF TOKI-SHAKUYAKU-SAN (JAPANESE TRADITIONAL MEDICINE) AND RESEARCH ON RELATED GENE EXPRESSION CHANGES: USING OVARIECTOMIZED RATS Mi Hwa Chung, Masao Hattori

### CP-M-009

EFFECT OF SIMVASTATIN MICROEMULSION IN THE TREATMENT OF ABDOMINAL SEPSIS IN WISTAR RATS. Bolívar P. G. L. Damasceno, Amália C. M. Régo, Irami Araújo-Filho, Maria Clara Araújo Silva, Victor Dominici Dominici, Eryvaldo Sócrates Tabosa Egito, Aldo da Cunha Medeiros

### CP-M-010

SIMVASTATIN MICROEMULSION FOR CICATRISATION OF INFECTED WOUNDS ON MALE WISTAR RATS Bolívar P. G. L. Damasceno, Amália C. M. Régo, Irami Araújo-Filho, Maria Clara Araújo Silva, Victor Dominici Dominici, Eryvaldo Sócrates Tabosa Egito, Aldo da Cunha Medeiros

### CP-M-011

SIMULTANEOUS DETERMINATION OF ACE ACTIVITY WITH TWO SUBSTRATES TELLS US ABOUT THE NATIVITY OF SOMATIC ACE AND ALLOWS TO DETECT ACE INHIBITORS IN HUMAN BLOOD. Sergei Danilov, Binevski Petr, Balyasnikova Irina, Albrecht Ronald, Kost Olga

### CP-M-012

ISOSTEVIOL EFFECTS ON THE CONTRACTIVE AND DILATIVE FUNCTIONS OF THE RABBIT HEART FOLLOWING ISCHEMIA-REPERFUSION INJURY. Andrew Keith Davey, Deyi Xu, Yongfang Li, Jiping Wang

### CP-M-013

LC-MS-MS DETECTION OF CORTICOSTERONE IN DIALYSATES SAMPLED FROM THE INTRACEREBRAL LATERAL VENTRICAL OF THE FREELY MOVING RAT Alistair Firth, Nick Andrews, Tamar Idle, Hugh Marston, Paul Scullion, Laura Johnston

## Drug Metabolism and Transport

### DMT-M-001

DRUG DISTRIBUTION INTO SALIVA: THE IMPLICATION OF PROTEIN BINDING Khalid Alkharfy, Tawfeeg Najjar, Sherif Saad

### DMT-M-002

INVESTIGATION OF HUMAN PHASE II ENZYMES INVOLVED IN THE METABOLISM OF ROTIGOTINE Käthe Becker, Susanne Bonsmann, Klaus Hansen, Ute Scharfenecker

### DMT-M-003

GASTROPLUS PBPK SIMULATION OF SIMVASTATIN: UPTAKE TRANSPORT BY HEPATIC SEROSAL OATP, BILIARY EFFLUX TRANSPORT BY MUCOSAL PGP, AND METABOLISM BY CYP3A4 IN GUT AND LIVER. Michael B. Bolger

### DMT-M-004

PROTON-COUPLED BASOLATERAL DIPEPTIDE TRANSPORT IN A HUMAN INTESTINAL CELL LINE, CACO-2. Birger Brodin

### DMT-M-005

PHARMACOKINETIC INTERACTION BETWEEN THE BARK OF MAGNOLIAE OFFICINALIS AND METHOTREXATE IN RATS Ying-Chen Chen, Shang-Yuan Tsai, Yu-Chi Hou, Pei-Dawn Lee Chao

### DMT-M-006

EFFECT OF GER-GEN CHIN-LIEN TANG ON THE PHARMACOKINETICS OF VALPROIC ACID IN RATS Lung-Yen Chen, Chung-Ping Yu, Su-Lan Hsiu, Yu-Chi Hou, Hsiu-Mei Chiang

### DMT-M-007

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**DMT-T-003**  
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**DMT-T-013**  
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Kenji Okada, Toshiimi Kimura, Ken-ichi Sagawa, Kiminori Mohri

**DMT-T-026**  
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