OBAVIJESTI

Hrvatsko farmaceutsko društvo

POZIVA

svoje članove da putem Udruženja i Sekcije pošalju

prijedloge kandidata za nagrade HFD-a

(Diploma, nagrada »Mr. ph. Antun Karlovac« i »Medalja prof. dr. Julije Domac«).

Prijedlozi trebaju biti utemeljeni i obrazloženi, a osnovni je uvjet aktivno članstvo i rad predloženog kandidata u HFD-u.

Udruženja i sekcije trebaju poslati svoje prijedloge u HFD, najkasnije do **10. lipnja 2011.**

Povjerenstvo za nagrade HFD-a razmotrit će pristigle prijedloge te mišljenje uputiti Upravnom odboru HFD-a, koji donosi konačnu odluku.

Nagrade zaslužnim članovima bit će dodijeljene na godišnjoj skupštini HFD-a.

Radi poštivanja kriterija pri predlaganju kandidata za pojedine nagrade molimo pogledati Pravilnike o nagradama HFD-a (dodatak Farmaceutskom glasniku, vol. 65, br. 2/2009.)

SEKCIJA FARMACEUTA JUNIORA organizira predavanje

u srijedu, 1. lipnja 2011., u **dvorani Gradske ljekarne Zagreb na Trgu Bana Jelačića 3** (stube lijevo I kat), s početkom u **10:15 sati**.

Tema: BETA BLOKATORI U TERAPIJI HIPERTENZIJE

Predavač: *Jasna Krznar, mr. pharm. spec.*, Opća bolnica Zabok

SEKCIJA ZA ANALITIKU LIJEKOVA organizira predavanje

u srijedu, 16. lipnja 2011., u prostorijama HFD-a (Knjižnica III kat), Masarykova 2, s početkom u **18 sati**.

Tema: MIKROBIOLOŠKA STABILNOST I MIKOTOKSIČKI RIZICI FARMACEUTSKIH SIRUPA

Predavač: *prof. dr. sc. Stjepan Pepeljnjak*

Tvrtka NYCOMED d.o.o. u suradnji s HRVATSKIM FARMACEUTSKIM DRUŠTVOM organizira stručni skup u Zagrebu u Hotelu Esplanade, u **srijedu, 15. lipnja 2011.** s početkom **u 20 sati.**

Teme:

1. VAŽNOST LJEKARNIKA PRI SAMOLIJEČENJU
   Mirna Radošević mag. pharm., Hrvatsko farmaceutsko društvo
2. SMJERNICE ZA SAMOLIJEČENJE SIMPTOMA REFLUKSA – KAKO URAVNOTEŽITI RIZIKE I KORISTI
   prof. dr. sc. Rajko Ostojić, dr. med. spec., Hrvatsko gastroenterološko društvo

Skup će biti vrednovan prema Pravilniku HLJK.

3. PHARMSCIFAIR 2011 'PHARMACEUTSKE ZNANOSTI ZA BUDUĆNOST LIJEKOVA'

'Farmaceutske znanosti za budućnost lijekova' PharmSciFair 2011, održat će se **od 13. do 17. lipnja 2011.** u Pragu (Republika Češka) u organizaciji Europske federacije za farmaceutske znanosti (*European Federation for Pharmaceutical Sciences*, EUFEPS) i brojnih partnerskih organizacija. PharmSciFair je mjesto gdje se susreću znanstvenici u farmaciji, a na tom skupu, trećem po redu, programom su obuhvaćeni raznovrsni znanstveni sadržaji, od otkrića i razvoja do kliničke primjene lijekova. Više informacija može se naći u PharmSciFair 2011 najavi na www.pharmscifair.org.

Putnička agencija TIP TOURS organizira putovanje autobusom na 3. PharmSci Fair od 13. do 17. lipnja 2011. Mogući su i individualni programi (zrakoplovom).
Više informacija: Sanja Ferenčak, e-mail: sanja@tiptours.hr; tiptours@tiptours.hr
tel.: 385 1 461 65 98; 466 37 52; faks: 385 1 466 37 54, www.tiptours.hr

4 BBBB – BLED INTERNATIONAL CONFERENCE ON PHARMACEUTICAL SCIENCES – New Trends in Drug Discovery, Delivery Systems and Laboratory Diagnostics

održat će se **od 29. rujna do 1. listopada 2011., na Bledu** (Slovenija), u organizaciji Slovenskog farmaceutskog društva i Farmaceutskog fakulteta Sveučilišta u Ljubljani.

Uz konferenciju održat će se satelitski simpozij **»Oral Modified Release – From Polymer to Drug Delivery System – Aims and Scopes«**

Cilj skupa je prikazati važna dostignuća u laboratorijskoj dijagnostici, medicinskoj kemiji i farmaceutskoj tehnologiji te omogućiti razmjenu znanstvenih ideja. Cilj je i povezati stručnjake sa sveučilišta, industrije i biomedicinske struke kao i mlade s već uvaženim znanstvenicima i stručnjacima. Glavne teme su:
– Recent strategies in drug delivery formulation: from materials to medicine
– Nanotechnology and nanotoxicity
– Translational medicine: from basic research to new diagnostic and treatment approaches
– All aspects of medicinal chemistry including drug design, discovery and development
– Methods, instrumentation and technologies in pharmacy and laboratory medicine
– Genomics and proteomics in diagnosis and treatment

Više informacija: www.bbbb-eufeps.org

Važni datumi:
Rok za slanje sažetaka usmenih i posterskih prezentacija **do 20. svibnja 2011.**
Autori će biti obavješteni o prihvaćanju sažetka **do 30. lipnja 2011.**
Svi prihvaćeni prošireni sažetci bit će objavljeni u posebnom izdanju *European Journal of Pharmaceutical Sciences.*

Registracija i smještaj:
Kongresni Servis Albatros d.o.o., Ribenska 2, 4260 Bled, Slovenija,
tel: +386 (0)4 5780 350, faks: +386 (0)4 5780 355, e-mail: info@albatros-bled.com, http://www.albatros-bled.com

Međunarodna konferencija

Unapređivanje sigurnosti pacijenata u Europi

(Reinforcing patient safety in Europe)

Zagreb, hotel The Westin Zagreb, 14. i 15. lipnja 2011.

U zajedničkoj organizaciji Europske agencije za lijekove (European Medicines Agency, EMA) i hrvatske Agencije za lijekove i medicinske proizvode (HALMED) u Zagrebu će se, u hotelu The Westin Zagreb, 14. i 15. lipnja 2011. održati međunarodna konferencija »Reinforcing patient safety in Europe« (Unapređivanje sigurnosti pacijenata u Europi). Konferencija je organizirana pod pokroviteljstvom Predsjednika Republike Hrvatske i Ministarstva zdravstva i socijalne skrbi Republike Hrvatske.

Konferencija je organizirana uz potporu EU programa Instrumenti za pomoć u predpristupnom razdoblju (Instruments for Pre-Accesion Assistance programme, IPA) koji, među ostalim, pomaže sudjelovanje država kandidata u aktivnostima europskih institucija.

Cilj konferencije je pružiti pregled europskog zakonodavstva o lijekovima te potaknuti razmjenu znanja, dijalog i suradnju hrvatskih i europskih stručnjaka na području regulative lijekova kao i pripremiti HALMED za buduće sudjelovanje u aktivnostima EMA-e.

Ova konferencija je ujedno nastavak prethodnih konferencija EMA-e održanih u Splitu i Rijeci pod nazivima »Zakonodavstvo za lijekove – hrvatski put ka članstvu u EU« i »EU regulatorna mreža – Izazovi i mogućnosti za Hrvatsku«.

Službeni jezik konferencije je engleski.

Prijava za sudjelovanje

Za sudjelovanje na konferenciji može se prijaviti putem internetskih stranica EMA-e. Registracijski obrazac koji je objavljen pod linkom: http://www.ema.europa.eu/docs/en_GB/document_library/Templates_and_Form/2011/04/WC500105599.doc potrebno je ispuniti i poslati na e-mail: extpax@ema.europa.eu ili fax: +44 (0)20 7418 8501.

Dodatne informacije o konferenciji nalaze se na Internet stranicama EMA-e pod linkom: http://www.ema.europa.eu/ema/index.jsp?curl=pages/news_and_events/events/2011/04/event_ detail_000417.jsp&murl=menus/news_and_events/news_and_events.jsp&mid=WC0b01ac058004d5c3.

Reinforcing patient safety in Europe
14-15 June 2011
Zagreb, Croatia

HALMED
Agency for Medicinal Products and Medical Devices of Croatia

EUROPEAN MEDICINES AGENCY
SCIENCE MEDICINES HEALTH

Zagreb, Croatia, 14-15 June 2011

Reinforcing patient safety in Europe

It is a great pleasure to announce the conference 'Reinforcing patient safety in Europe', to be held in Zagreb, Croatia, on 14-15 June 2011.

This conference is co-organised by the European Medicines Agency and the Croatian Agency for Medicinal Products and Medical Devices (HALMED), under the auspices of the Croatian Ministry of Health and Social Welfare.

The conference is co-funded by HALMED and the European Union's Instrument for Pre-accession Assistance (IPA) Programme, designed to support pre-accession activities of the beneficiaries, i.e. Croatia, the former Yugoslav Republic of Macedonia, Turkey, Albania, Bosnia and Herzegovina, Montenegro, Serbia and Kosovo under UNSC Resolution 1244/99.

Main objectives of the conference

- To exchange views and establish dialogue with interested parties.
- To give an overview of EU legislation on the regulation of medicinal products.
- To provide feedback on experience of applying the *acquis* and related legal aspects.
- To assist the national competent authorities with preparations for their future involvement in the European Medicines Agency's activities.

Conference programme

- Accession preparation.
- Variation legislation.
- Generics.
- Electronic submission in the EU: tools
- Benefit/risk analysis and risk minimisation.
- Product information management.
- Readability testing.
- New pharmacovigilance and risk assessment legislation.
- Crisis management.
- Safety aspects of vaccines.
- Other topics (to be determined).

Target audience

This conference will target scientific audiences and will include specific activities with the aim of informing pharmaceutical stakeholders working in regulatory affairs.

Conference Secretariat

European Medicines Agency
7 Westferry Circus, Canary Wharf
London E14 4HB, United Kingdom
Telephone +44 (0)20 7418 8400
Fax +44 (0)20 7418 8501
E-mail conferences@ema.europa.eu

Further information

European Medicines Agency website:

www.ema.europa.eu
Home > Partners & networks > Europe & the Agency > EU enlargement

HALMED
Agency for Medicinal Products and Medical Devices of Croatia

EUROPEAN MEDICINES AGENCY
SCIENCE MEDICINES HEALTH

20 April 2011
EMA/172829/2011
Administration

Draft conference programme

Reinforcing patient safety in Europe, 14-15 June 2011, Zagreb, Croatia

Tuesday, 14 June 2011

Time	Session
09.30-10.00	Welcome coffee
10.00-10.20	**Welcome address**
10.20-11.10	**Welcome note**
11.10-12.30	**Accession preparation I** ▪ Legal framework from regulatory affairs ▪ Situation in Croatia ▪ Perspectives from a new Member State ▪ Phasing-in (product specific)
12.30-14.00	Lunch
14.00-15.00	**Accession preparation II** ▪ Preparing for dossiers evaluation ▪ Adapting to community procedures (incl. advanced therapies)
15.00-16.00	**Non-clinical assessment requirements** ▪ Non-clinical assessment requirements ▪ Perspectives from a Member State
16.00-16.30	Coffee break
16.30-19.00	**Legislation**
16.30-17.30	Variation legislation
17.30-18.30	Generics

7 Westferry Circus • Canary Wharf • London E14 4HB • United Kingdom
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E-mail info@ema.europa.eu **Website** www.ema.europa.eu
An agency of the European Union

Time	Session
	▪ Tricky issues; ▪ Interchange ability of generics ▪ Non-prescription switching
18.30-19.00	Electronic submission in the EU: tools
21.00-23.00	Welcome drink and conference dinner

Wednesday, 15 June 2011

Time	Session
09.00-10.00	**Benefit risk and risk minimisation** ▪ Assessment and communication ▪ Risk minimisation
10.00-11.45 10.00.10.45	**Product information** Product information management
10.45-11.15	Coffee break
11.15-11.45	Readability testing
11.45-12.45	**Pharmacovigilance** ▪ Summary of the new pharmacovigilance legislation ▪ Perspective from a MS
12.45-14.00	Lunch
14.00-15.30 14.00-14.45	**Crisis management case studies** ▪ Communication ▪ Quality defects
14.45-15.30	Safety aspects of vaccines ▪ Evaluation ▪ Pharmacovigilance
15.30-16.00	Conference closure

AMERICAN ASSOCIATION OF PHARMACEUTICAL SCIENTISTS (AAPS)
and CROATIAN PHARMACEUTICAL SOCIETY (HFD)

organize

Workshop on dissolution testing, biowaivers, and bioequivalence

Location and date:

PLIVA Research Institute Ltd., Prilaz baruna Filipovića 29, Zagreb.

3–4 October 2011

Objectives:

The goal of this workshop is to discuss the following aspects:

- Method development, including Quality by Design concepts
- Disintegration vs Dissolution testing
- GMP in Dissolution laboratory
- *In vitro – in vivo* correlations (IVIVC)
- Biowaivers based on the Biopharmaceutics Classification System (BCS)
- Application for biowaivers of pharmaceutical products following post-approval changes
- Bioequivalence studies (General concepts IR and CR formulations; Current regulatory aspects in Europe and USA)

Academic, industrial and regulatory perspectives will be presented for these aspects.

Target participants: Industry, academia, government, scientists.

Language: The official language of the Workshop is English.

PRELIMINARY PROGRAMME:

REGISTRATION:

Monday, 3 October, 2011 – 08:00–11:00

LECTURE SCHEDULE:

Monday, 3 October 2011 (Day 1)

09:00–9:15 *Opening remarks – Biserka Cetina-Čižmek*

PRINCIPLES AND APPLICATIONS OF DISSOLUTION TESTING

Session Chair: Terry Way

09:15–10:00 *Dragica Raušl, Pliva, Croatia*
Dissolution testing – a powerful tool in drug product development

10:00–10:45 *Vivian Gray, V. A. Gray Consulting, USA*
General concepts – Design of dissolution method development, including Quality by Design (QBD)

10:45–11:15 Break

Session Chair: Johannes Kraemer

11:15–12:00 *Marija Bogataj, University of Ljubljana, Slovenia*
Non-compendial approaches to Dissolution testing

12:00–12:45 *Nikoletta Fotaki, University of Bath, UK*
Predictive Dissolution testing and development of IVIVCs

12:45–13:00	Questions & Answer Session – Panel Discussion Moderator: *Johannes Kraemer*
13:00–14:00	Lunch
	Session Chair: Biserka Cetina-Čižmek
14:00–15:00	*Vivian Gray, V. A. Gray Consulting, USA* Good Manufacturing Practices (GMP's) in the Dissolution laboratory
15:00–15:45	*Terry Way, USP (Europe office)* The role of Dissolution testing: USP perspective
15:45–16:15	Break
16:15–17:00	*Johannes Kraemer, Phast, Germany* Disintegration instead of Dissolution: A case study
17:00–17:30	Question & Answer Session – Panel Discussion Moderator: *Vivian Gray*
20:00	Dinner

Tuesday, 4 October 2011 (Day 2)

	REGULATIONS
	Session Chair: Nikoletta Fotaki
09:15–10:00	*Selvira Zulfikari, HALMED, Croatia* Croatian Regulatory Perspective
10:00–10:45	*Tahseen Mizra, FDA, USA* FDA perspective on IVIVC and BCS
10:45–11:15	Break
11:15–12:00	*Evangelos Kotzagiorgis, EMA* Regulatory perspective (Europe) on Dissolution testing
12:00–12:30	Question & Answer Session and Panel Discussion Moderator: *Nikoletta Fotaki*
12:30–13:30	Lunch
	FROM DISSOLUTION TESTING AND BCS TO BA/BE STUDIES
	Session Chair: Vivian Gray
13:30–14:15	*Johannes Kraemer, Phast, Germany* Examination of Level B Correlations
14:15–15:00	*Nikoletta Fotaki, University of Bath, UK* From dissolution to BA/BE studies
15:00–15:30	Break
15:30–16:15	*Senka Radošević, Pliva, Croatia* Importance of study design and metrics for the proof of bioequivalency
16:15–16:45	Question & Answer Session – Panel Discussion Moderator: *Biserka Cetina-Čižmek*
16:45	**Closing Remarks – *Vivian Gray***

REGISTRATION FEE
Up to 15 July, 2011: 300 EUR
Up to 15 September, 2011: 350 EUR
After 15 September, 2011: 400 EUR

Registration should be made (from 1 June 2011) electronically by using Registration form given at the web site www.tiptours.hr or sending it by fax on +385 1 466 37 54.

Registration fee for participants includes admission to all Workshop scientific events, written materials & badge, certificate of attendance, daily tea/coffee/lunch & workshop dinner.

A complimentary copy of the book titled »Handbook for Dissolution Testing«, 3rd Edition will be provided to attendees.

Program information:
Dragica Raušl
PLIVA Croatia Ltd
Tel.: +385 (0)1 372 30 36
Fax: +385 (0)1 372 15 14
E-mail: dragica.rausl@pliva.com

Exhibition and sponsorship information:
Maja Jakševac-Mikša
Croatian Pharmaceutical Society
Tel.: +385 (0)1 487 28 49
Fax: +385 (0)1 487 28 53
E-mail: hfd-fg-ap@zg.t-com.hr

Travel Agency:
TIP TOURS
Att. Sanja Ferenčak
Vončinina 2/1, 10000 Zagreb, Croatia
Tel. +385 1 461 65 98; 466 37 52
Fax. +385 1 466 37 54
e-mail: sanja@tiptours.hr; tiptours@tiptours.hr
Web page: www.tiptours.hr

Accommodation:
For participants a quotation of rooms was booked at a special conference prices at the Hotel Dubrovnik Zagreb. Reservations will be made on first come first served basis.

Plan tečajeva stručnog usavršavanja magistara farmacije u organizaciji HFD-a, u drugoj polovini 2011. (listopad – studeni)*

Naziv tečaja	Organizatori	Mjesto
Hitna stanja i pomoć u izvanbolničkim uvjetima	HFD i Zavod za hitnu medicinu Grada Zagreba (dvodnevni tečaj)	Zagreb, 17. i 18. listopada
Kozmetika u ljekarni	HFD	Zadar, Rijeka listopad – studeni
Tečaj za voditelje pripravnika	HFD (dvodnevni tečaj)	Zagreb, studeni

*Molimo kolege da se prijave za tečaj kojem žele prisustvovati.
Najavljeni tečajevi će se održati ako se prijavi dovoljan broj zainteresiranih polaznika.

5. MAKEDONSKI KONGRES FARMACIJE s međunarodnim sudjelovanjem održat će se na **Ohridu** (Hotel Metropol), **21.–25. rujna 2011.**, u organizaciji Makedonskog farmaceutskog društva i Farmaceutskog fakulteta - Tema kongresa je **»Znanost i praksa za dobrobit zdravlja«**. Detaljnije informacije: www.mfd.org.mk

2. KONGRES FARMACEUTA BOSNE I HERCEGOVINE s međunarodnim sudjelovanjem održat će se u **Banja Luci, 17.–20. studenoga 2011.**, u organizaciji Farmaceutskog društva Republike Srpske, Farmaceutskog društva Federacije Bosne i Hercegovine i Međukantonalne farmaceutske komore. Tema kongresa je **»Generički lijekovi u farmaciji«**. Detaljnije informacije: www.farmakongres2011.ba

71. MEĐUNARODNI KONGRES FIP-a
SVJETSKI KONGRES FARMACIJE I FARMACEUTSKIH ZNANOSTI 2011
Hyderabad, Indija, 3. – 8. rujna 2011.

Glavna tema kongresa je *Compromising safety and quality: A risky path*.

Sadržaj radnog dijela kongresa:

Pre-satelitski simpoziji (3):
R1-1. Pharmacovigilance and medicines information to enhance patient safety
R1-2. Pharmacovigilance and medicines information to enhance patient safety
R2. Pharmacy in India

Dijelovi glavne teme Compromising safety and quality: A risky path (4):
A1. A primer on quality and safety
A2. Learning from errors and monitoring safety
A3. Building a safer service: tehniques and tools to improve quality and safety
A4. Paying pharmacists for patient outcomes: pay for performance?

Od znanosti do prakse (1):
B1. Environment and pharmaceuticals

Farmaceutske znanosti (7):
C1. WHO guidelines on multisource drugs and interchangeability
C2. Biosimilars
C3. Clinical research
C4. Paradigm shift in drug discovery and development
C5. Pharmaceutical manufacturing
C6. Standardization of herbal products
C7. Dissolution: the pivotal tool for developing quality drugs

Sekcijske skupine izlaganja (23):
D1. Community pharmacy business models: business and financial aspects of implementing and integrating pharmaceutical services - integrating professional services with the business of a pharmacy (Forum for innovators in pharmacy practice) (Part 1)
D2. Community pharmacy business models: business and financial aspects of implementing and integrating pharmaceutical services - integrating professional services with the business of a pharmacy (Forum for innovators in pharmacy practice) (Part 2)
D3. Clinical biology in India health care system: organisation and contribution

D4. Current issues session - vulnerable populations: what are their medicine/health information needs and how can we address these needs?
D5. Solving practical tableting problems
D6. The Basel statements in developing and developed countries - what are the right ingredients?
D7. A glimpse of community pharmacy in 2020 - discussion forum on community pharmacy 2020
D8. Short oral communications of the FIP clinical biology section
D9. Medication safety and risk management
D10. Recent advances and challenges in the safe preparation of cytotoxic agents
D11. The practitioners' day - practical solutions to health problems and service provision (Part I)
D12. The practitioners' day - practical solutions to health problems and service provision (Part II)
D13. Innovations to improve teaching and learning
D14. Quality and safety in pharmacologistics
D15. Pharmacists and mass communication - a job that needs to be done continuously
D16. Aspects of medication and patient safety - social and administrative pharmacy section contributed papers [short oral communications]
D17. Good manufacturing practices - expectations for the coming decade (Part I)
D18. Good manufacturing practices - expectations for the coming decade (Part II)
D19. Ask your pharmacist day (Part 1)
D20. Ask your pharmacist day (Part 2)
D21. Contributed papers - short oral presentations of the FIP academic pharmacy section
D23. Communication and control in an operational setting

Ostalo (11):
F1. Careers and leadership in pharmacy and education
F2. Generics and the patient experience: the pharmacist's role in ensuring safe and effective medicines use
F3. FIP/WHO joint session on public-private mix for tuberculosis care and control
F4. Report of the FIP working group on optimizing the role of pharmacists in improving maternal, newborn, and child health (MNCH)
F5. Outcomes of the pharmacy education taskforce: a report, review and reflection
F6. Mapping a new vision - translating ideas into practice
F7. Symposium on the history of pharmacy (Part 1)
F8. Symposium on the history of pharmacy (Part 2)
F9. FIP member organisations presenting national updates (Part 1)
F10. FIP member organisations presenting national updates (Part 2)
F11. FIP member organisations presenting national updates (Part 3)

Zajednička sekcijska izlaganja: (14)
J1. Building practitioners skills
J2. Your career in industrial pharmacy – from drug development to drug distribution
J3. Regulatory and legislative changes in pharmacy from across the world
J4. Pharmacogenomics in oncology
J5. Careering toward advanced levels of practice
J6. Communicating basic medicines information to patients
J7. Trends in community pharmacy - debating the future of the profession: forum for policy makers
J8. Competition for the best oral industrial presentation (short oral communications)
J9. Pediatric medicines - challenges and opportunities

J10. Building a »toolbox« for practitioner development and support
J12. Pharmaceuticals in the environment
J13. Developing young academics through networking and mentoring
J14. Pharmacovigilance: ensuring serious medication safety concerns are recognized, addressed, reported and monitored

Važni datumi:
Rok za slanje sažetaka: **1. travnja 2011.**
Prihvaćanje sažetaka bit će objavljeno na web stranici: www.fip.org/hyderabad2011 **nakon 1. lipnja 2011.**
Autor koji izlaže rad treba se registrirati i platiti kotizaciju do **15. svibnja 2011.**

Kotizacija	Uplata do 15. svibnja 2011.	Uplata do 1. kolovoza 2011.	Uplata nakon 1. kolovoza 2011. i na licu mjesta
Individualni članovi FIP-a	€ 585	€ 685	€ 995
Nečlanovi	€ 825	€ 925	€ 995
Studenti/nedavno diplomirani farm.	€ 200	€ 250	€ 300
Dnevna kotizacija (na licu mjesta)			€ 300
Pre-satelitski simpozij R1 (prijava moguća uz uplatu kotizacije za kongres)	€ 85	€ 85	€ 85
Osobe u pratnji	€ 150	€ 150	€ 150

Kotizacija uključuje: prisustvovanje radnom dijelu, osim pre-satelitskom simpoziju R1, svečanom otvaranju, prijamu dobrodošlice, ulaz na izložbu, pristup svim apstraktima i biografijama na webu (od 15.8.2011), kongresnu torbu s konačnim programom i popisom sudionika, pristup web stranici s koje se mogu preuzeti prezentacije (od 1.12.2011.)

Informacije:
FIP Congresses & Conferences
P.O. Box 84200
NL-2508 AE The Hague
The Netherlands
Tel.: (+31) (0) 70 302 1982
Fax: (+31) (0) 70 302 1998
E-mail: congress@fip.org
Website: www.fip.org/hyderabad2011

Registracija i sažeci:
New Brooklyn
P.O. Box 73
3620 AB Breukelen
The Netherlands
Tel.: (+31) (0) 346 767 231
Fax: (+31) (0) 346 263 308
E-mail: registration@newbrooklyn.nl
Website: www.newbrooklyn.nl

Mjesto održavanja:
Hyderabad
Convention Centre (HICC)
Novotel &HICC Complex
Hyderabad – 5000 081
India
Tel: +9140 66824422/
66134422

Brošura s programom kongresa može se dobiti u poslovnici HFD-a, Masarykova 2/II, Zagreb.