

Scientific Program

3rd World Congress on Bioavailability & Bioequivalence

Pharmaceutical R & D Summit

March 26-28, 2012 Marriott Convention Centre, Hyderabad, India



Conference Secretariat

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Marriott Convention Centre

Day 1 March 26, 2012

08:30-09:30 Registrations 10:00-11:00 Opening Ceremony

Honorable Guests

Shri. R. P. Thakur, IPS

Director General-Drugs Control Administration, Andhra Pradesh, India

Shri. A. Chandra Sekhar Rao

Deputy Drugs Controller-Central Drugs Standard Control Organization, Andhra Pradesh, India

Shri. K. Praveen Kumar

Vice President, Marketing and Investment Promotion, APIIC, Andhra Pradesh, India

Dr. Luigi Silvestro

Co-founder, 3S-Pharmacological Consultation & Research GmbH, Germany

Dr. Gregory Russell-Jones

Director, Mentor Pharmaceutical Consulting, Australia

Dr. Patrick Bennett

Strategic Marketing Director, Thermo Fisher Scientific Pvt. Ltd., USA

Dr. Amit Biswas

Executive Vice President, Dr. Reddy's Laboratories, India

Dr. A.T. Bapuji

Senior Vice President, Aurobindo Pharma, India

Dr. Gedela S. Babu

Director, OMICS Group Inc., USA

		Correebreak: 11:00-11:15
	Keynote Forum	Introduction
11:15-11:45	Title: Research to routine workflows for large n mass spectrometry	nolecules using proteomic tools and high resolution
	Dr. Patrick Bennett, Thermo Fisher Scientific Pvt. Lt	rd., USA
11:45-12:15	Title: Bioavailability and bioequivalence: Do th	ney correlate?
	Dr. Gregory Russell-Jones, Mentor Pharmaceutica	ıl Consulting, Australia
10 15 10 45	Title: BA/BE Challenges to generic pharmaceutical	l industry
12:15-12:45	Dr. A.T. Bapuji, Aurobindo Pharma, India	
		Lunch: 12:45-13:30
13:30-14:00	Title: Anti-angiogenesis therapy: Clinical puzzl	es and prospects
13:30-14:00	Prof. Debabrata Mukhopadhyay, Mayo Clinic Co	llege of Medicine, USA
14:00-14:30	Title: Harmonization in BABE Studies: Regulato	ory aspects
	Dr. T. S. Jaishankar, Quest Life Sciences (P) Ltd, In	dia
14:30-15:00		ted as exploitation- how much is perceptive and how
	much is reality? – Specific to India	
	Dr. S.V. Krishna Prasad, Cito Healthcare Pvt. Ltd,	
		Cofee Break: 15:00-15:15

Marriott Conv	vention Center
	lemiology of Drug Interactions: Biochemical Mechanisms of Drug Toxicity
Track 1.4: Inte	grated Product Development: Structure Based Drug Design
	Session Introduction
15:15-15:40	Title: The role of vitamin D in weight-related metabolic derangements, inflammation and insulin resistance
	Dr. Khalid M Alkharfy, King Saud University, Saudi Arabia
15:40-16:05	Title: In silico design of novel, high-affinity neuraminidase inhibitors for Influenza A/H1N1/2009
	virus
	Dr. Shailendra K. Saxena, Centre for Cellular and Molecular Biology, India
	Title: Salt intake influences the interaction between the renin-angiotensin and renal dopaminergic
16:05-16:30	systems in normotensive humans
	Dr. Aruna R. Natarajan, Georgetown University School of Medicine, USA
16:30-16:55	Title: Photolabile probes to control cellular chemistry and drug discovery
10:30-10:55	Dr. Srinivas Kantevari, Indian Institute of Chemical Technology (IICT), India
	Title: Novel bio-chemical profiling of Indian black teas and development of green tea based radical
16:55-17:20	scavenging conserve as all natural antioxidant for health promotion
	Dr. B.B. Borse, Central Food Technological Research Institute, India
17:20-17:45	Title: Model based drug development: Promises towards success
	Dr. Abhay Sangamwar, National Institute of Pharmaceutical Education and Research, India
Diamond Ha	I
	g Design, Development and Therapy
Track 1.2: Nev	v Strategies for Drug Development
	Session Introduction
15:15-15:40	Title: Developing suitable animal models for drug development
	Prof. Sharma S. Prabhakar, Texas Tech University Health Sciences Center, USA
15:40-16:05	Title: Challenges for trace element analysis in BABE studies
	Dr. Biswajayee Patra, Thermo Fisher Scientific Pvt. Ltd., India
16:05-16:30	Title: Design and synthesis of potent Cyclin C inhibitors
	Dr. P. Sarita Rajender, Osmania University, India
16:30-16:55	Title: Computational technique – A promising pathway for drug discovery: A case study
	Prof. Uma Vuruputuri, Osmania University, India
16:55-17:20	Title: An overview of role of public regulatory authorities in implementing guidelines for drug development and approval globally
10.00 17.20	Dr. Sandeep Arora, Chitkara University, India
	Title: A novel approach of piperine incorporated cubosomal preparations for vitiligo therapy
17:20-17:45	Dr. Vinod K. R, Nalanda College of Pharmacy, India
Ball Room	
15:15-17:00	Poster Presentations (PO-001 to PO-015)
	Day 2 March 27, 2012
Diamond Ha	

Track 2: Identification and Characterization of Drugs

Session Introduction

09:30-09:55 study on few selected plants Prof. Kedam Thyaga Raju, Sri Venkateswara University, India

Title: New compounds that cause inhibition to various reactive oxygen species are from plants: A

09:55-10:20	Title: Role of Zinc on prostate cancer
09:55-10:20	Dr. J. Arunakaran, University of Madras, India
10:20-10:45	Title: Assay of endogenous Drugs
	Dr. D.Vijaya Bharathi, Dr.Reddy's Laboratories Ltd, India
10:45-11:10	Title: Application of NMR spectroscopy for structure elucidation of bioactive natural and synthetic
	compounds
	Dr. T. Narender, Central Drug Research Institute, India
	Coffeebreak 11:10-11:25
11.25 11.50	Title: Recent advances in analytical methodologies for the determination of impurities in drugs
11:25-11:50	Dr. M V Narendra Kumar Talluri, National Institute of Pharmaceutical Education and Research, India
11:50-12:15	Title: Season variation and starvation period influence on the antithrombotic activity of Leech
	Saliva extract from the medicinal Malaysian Leech, Hirudinaria manillensis
	Prof. Abbas Mohammad Ghawi, International Islamic University Malaysia, Malaysia
	Title: Ascorbate defense in response to hyperosmotic stress by adrenal and gonads in albino rat
12:15-12:40	(Rattus norvegicus)
	Dr. Priyanka Mehta, M.J.P. Rohilkhand University, India
12:40-13:05	Title: A fully automated system for LC/MS bioanalysis in regulated laboratories
12.40-13.05	Dr. Doug McIntyre, Agilent Technologies, India
	Lunch 13:05-13:55
Track 5: Bioav	ailability and Bioequivalence of Novel Drug Delivery Systems
	Session Introduction
	Title: Transdermal needle-free delivery of peptides and proteins using water-in-oil microemulsions:
13:55-14:20	High bioavailability, with drug dependent pharmacology
	Dr. Gregory Russell-Jones, Mentor Pharmaceutical Consulting, Australia
14:20-14:45	Title: Oral drug delivery: Pharmaceutical technology archetypes
	Dr. Shivanand P. Puthli, Panacea Biotec Ltd., India
14:45-15:10	Title: Effect of bioenhancers on permeability and In-Vitro release of various anti-tubercular agents
	Dr. Pingale Prashant L, NMIMS Shirpur Campus, India
	Title: Transdermal delivery of lopinavir loaded ultradeformable vesicles (UDVs): An alternate way
15:10-15:35	to circumvent its limited oral bioavailability
	Dr. Hetal Thakkar, M.S.University of Baroda, India
	Coffeebreak 15:35-15:50
	Title: Design, development and evaluation of Ultra-deformable vesicular drug delivery system for topical delivery of itraconazole
15:50-16:15	
	Dr. Shivprasad Majumdar, SVKM's \NMIMS, India
16:15-16:40	Title: Bioavailability assessment of topically applied ufasomal formulation for treatment of Arthritis
	Mr. Arvind Sharma, Chitkara university, India
16:40-17:05	Title: Investigation of DNA integration into reproductive organs following intramuscular injection of DNA in mice
10:40-17:05	Dr. Fatemeh Vahedi, Razi Vaccine & Serum Research Institute, Iran
Marriott Con	vention Center
	nacokinetics, Bioavailability and Bioequivalence
	Session Introduction
09:30-09:55	Title: Metabolism after lung absorption in comparison to oral route. A key to interpret results of
	lung deposition studies based on PK
	Dr. Luigi Silvestro, 3S-Pharmacological Consultation & Research GmbH, Germany

09:55-10:20	Title: A bioequivalence study of two Azithromycin tablet formulations in Indonesian healthy volunteers	
07:55-10:20	Prof. Yahdiana Harahap, University of Indonesia, Indonesia	
10:20-10:45	Title: Does hemolysis affect the pharmacokinetic profile of drugs used in bioavailability studies	
	Dr. Shengjun Zhang, Frontage China Clinical Research Center, China	
	Title: Beyond sensitivity: Improving the performance, productivity and compliance of the	
10:45-11:10	bioanalytical assay process	
	Dr. Gopal Vaidyanathan, Waters India Pvt. Ltd., India	
	Coffeebreak: 11:10-11:25	
11:25-11:50	Title: Enzymatic degradation of chitosan and thiolated chitosan	
	Dr. Flavia Laffleur, University of Innsbruck, Austria	
	Title: Theoretical and in vitro studies of a C-terminal peptide from PGKC of Leishmania mexicana	
11:50-12:15	mexicana	
12:15-12:40	Dr. Vidya Raghunathan, National Institute of Immunology, India	
	Title: Increasing Tamoxifen bioavailibility using Tam:HP β CD inclusion complex	
	Dr. Jaya Shukla, Postgraduate Institute of Medical Education and Research, India	
12:40-13:05	Title: Comparison of biological activities between synthesized ester derivatives and its parent compound Solanesol	
12.40-13.05	Prof. Sridevi Pingala, CM College of Pharmacy, India	
	Lunch: 13:05-13:55	
	Title: Fate of pharmacologically active Isoflavones upon administration in Sprague Dawley rats	
13:55-14:20	Mr. Wahajuddin, CSIR-Central Drug Research Institute, India	
	Title: Diosmetin Pharmacokinetic Following Diosmin Oral Administration in Man; a New Study on	
14:20-14:45	an Old Product with Controversial Pharmacokinetic Findings in the Past	
	Mrs. Adriana Iordachescu, Pharma Serv International SRL, Romania	
Track 4: PK/PI	O Models, Biowaivers and Biosimilars	
	Session Introduction	
14:45-15:10	Title: Free radical scavenging activity of the Malaysian Leech Saliva extract, Hirudinaria manillensis	
	Prof. Abbas Mohammad Ghawi, International Islamic University Malaysia, Malaysia	
15:10-15:35	Title: Bio-Sensor in medication	
	Dr. Awadhesh Kumar Singh, Maharaja Agrasen Institute of Technology, India Coffeebreak: 15:35-15:50	
	Title: Evaluation of the antitrypanosomal potential of the epidermal tissues of Vitex Doniana root	
15:50-16:15	extract using wistar rats	
	Dr. Innocent Omalu, Federal University of Technology, Nigeria	
	Title: Pharmacokinetic driven drug discovery and development strategies	
16:15-16:40	Dr. Rabi Sankar Bhatta, CSIR-Central Drug Research Institute, India	
16:40-17:05	Title: Glucose 6 phosphate dehydrogenase and haematological activities of artemether and	
	lumefantrine in non-malaria infected rats	
	Dr. Abiodun Humphrey Adebayo, Covenant University, Nigeria	
Ball Room		
	Poster Presentation (PO-016 to PO-056)	
15:00-16:30	Poster Presentation (PO-057 to PO-078)	

March 28, 2012 Day 3

Marriott Convention Center

Track 6: Strategies and Challenges in Bioanalysis

	Session Introduction
09:30-09:55	Title: Simplified approach for the development of a bioanalytical DBS assay using a tandem quadrupole with a novel collision cell design
	Dr. Gopal Vaidyanathan, Waters India Pvt. Ltd., India
09:55-10:20	Title: Evaluation of bioequivalence for endogenous molecules; Statistical Impact of different basal levels correction procedures
	Dr. Simona Rizea Savu, 3S-Pharmacological Consultation & Research GmbH, Germany
10:20-10:45	Title: Quality by design (QbD)using design-of-experiments (DoE)approach to develop a LC-MS/MS method
	Dr. Sandeep Sharma, Jubilant Clinsys Ltd., India
10:45-11:10	Title: Gallopamil racemate separation using UV spectrophotometer with preparative chromatographic column
	Dr. Kalaichelvi Ponnusamy, National Institute of Technology, Tiruchirappalli, India
	Coffeebreak: 11:10-11:25
11:25-11:50	Title: A cloud computing system to quickly implement new microarray data normalization methods
	Dr. Dongquan Chen, Univ. of Alabama at Birmingham, USA
11:50-12:15	Title: Harmonizing best practices in bioanalytical methods
	Dr. Maha Tutunji, University of Jordan, Jordan
12:15-12:40	Title: Study of selenium-containing metabolomes in selenium-enriched yeast by ion exchange- inductively coupled plasma- mass spectrometry
	Dr. D. T. Pal, National Institute of Animal Nutrition and Physiology (NIANP), India
	Lunch: 12:40-13:30

Track 10: Oral Bioavailability Enhancement Track 11: Regulatory and Economical Aspects in Bioavailability and Bioequivalence	
	Session Introduction
13:30-13:55	Title: Novel systems for the preparation of oral dosage forms for poorly water soluble curcuminoids co-formulated with water soluble peptides and proteins
	Dr. Gregory Russell-Jones, Mentor Pharmaceutical Consulting, Australia
13:55-14:20	Title: Trend analysis on repeats, a way through
	Ms. Neerja Somyajee, Piramal Healthcare Limited, India
14:20-14:45	Title: Innovation in method development; Drugs solubility and stability during bioanalysis process
	Dr. Naser Rezk, King Abdullah International Medical Research Center, Kingdom of Saudi Arabia
14451510	Title: Federated RHIO solution for HR healthcare data protection
14:45-15:10	Dr. Jayanthi Ranjan, Institute of Management Technology-IMT, India
15:10-15:35	Title: The dysfunction of cGMP-dependent Na/Ca exchange as a gate for age-related dysfunction of cellular activity
	Prof. Sinerik N. Ayrapetyan, UNESCO Chair-Life Sciences International Postgraduate
	Educational Center, Armenia
	Coffeebreak: 15:35-15:50
15:50-16:15	Title: Exemestane solid dispersions: Enhancement of solubility and permeability
	Dr. Suresh Bandari, St Peter's Institute of Pharmaceutical Sciences, India
16:15-16:40	Title: Enhancement of oral bioavailability of ketoprofen by liquisolid compaction technology
	Mr. Vijaykumar Nagabandi, St. Peter's Institute of Pharmaceutical Sciences, India

16:40-17:05	Title: Microwave generated bionanocomposites for solubility and dissolution enhancement of poorly water soluble drug Ibuprofen: In vitro and in vivo evaluation
	Dr. Sachin Kushare, R. G. Sapkal College of Pharmacy, India Title: Simvastatin solid lipid nanoparticles for improving the oral bioavailability
17:05-17:30	Prof. Lankalapalli Srinivas, GITAM University, India
	Title: The novel modified approaches for enhancement of bioavailability of poorly water-soluble
17:30-17:55	drugs
	Dr. Abhishek Kumar Jain, Sagar Institute of Research and Technology, India
17:55-18:20	Title: Regulatory Needs in Bioavailability and Bioequivalence Studies of Pharmaceuticals
	Dr. G. S. Chakraborthy, Noida Institute of Engineering and Technology, India Title: Toxicity and degradation of chlorinated nitroaromatic compounds
18:20-18:45	Dr. Pankaj Kumar Arora, University of Hyderabad, India
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Track 7: Bioan	alytical Methodology Development and Validation Using HPLC and LCMS
Track 9.1: Des	ign, Documentation and Reporting of BA/BE Studies: Protocol Issues
	Session Introduction
09:30-09:55	Title: An Update on Harmonisation of Bioanalytical guidances by GBC (Global Bioanalysis Consortium)
•••••	Dr. S. Ravisankar, GVK Biosciences Pvt Ltd, India
09:55-10:20	Title: Enhancing bioanlytical selectivity beyond conventional LC/MS/MS
07:55-10:20	Dr. Hesham Ghobarah, ABSCIEX, Canada
	Title: Pilot Study informs pivotal trial evaluating bioequivalence of two formulations of Paroxetine
10:20-10:45	40 mg tablet in healthy Chinese subjects Dr. Shengjun Zhang, Frontage China Clinical Research Center, China
	Title: Matrix effects: Termite of analytical methodologies
10:45-11:10	Dr. Chinmoy Ghosh, Cadila Pharmaceuticals Limited, India
	Coffeebreak: 11:10-11:25
	Title: Validated stability-indicating HPLC method for the determination of eberconazole nitrate:
11:25-11:50	Application to hydrolytic, thermal, oxidative and photolytic degradation kinetics Dr. Vamsi Krishna Marothu, Alliance Institute of Advanced Pharmaceutical and Health Sciences, India
	Title: Evaluation of antidiabetic and antioxidant activity of Moringa oleifera in experimental
11:50-12:15	diabetes
	Dr. Rajnish Gupta, University of Rajasthan, India
12:15-12:40	Title: Formulation and evaluation of Terbinafine Hydrochloride Microsponge Gel [THMG]
	Dr. Vedavathi Tavva, CMR College of Pharmacy, India
	Lunch: 12:40-13:30 Title: Challenges for trace element analysis in BABE studies
13:30-13:55	Mr. Sudhakar Kasturi, Thermo Fisher Scientific Pvt. Ltd., India
	Title: Simultaneous determination of amoxicillin and clavulanic acid in human plasma by liquid
13:55-14:20	chromatography coupled with tandem mass spectrometry
	Mr. Chaitanya Krishna Atmakuri, Bombay Bio Research Centre, India
14:20-14:45	Title: Design, documentation and reporting of BA/BE studies on anticancer products: Protocol issues
	Mr. Vikas Kumar, Fresenius-Kabi Oncology Ltd., India Coffeebreak: 15:35-15:50
Ball Room	
	Poster Presentation (PO-079 to PO-119)

Encyclopedia of Bioequivalence and Bioavailability (E-BABE)

Analytical and Bio-Analytical Methodology Database

- World's only database powered by an updated bioanalytical methods of pharmaceuticals for suitable method selection with over thousands of combinations and automatic checks against thousands of methods: all at your finger tips.
- Intelligent processing which understands your analytical terminology and lets you generate, select of your priority and much more in seconds.
- Use the comprehensive analytical and bio-analytical methodology tool to select important method for an insight into regular quality control.
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- All-in-one database with multi-user access.
- As of now it has 5000 methods and is updated regularly.

http://ebabe.gsblifesciences.org

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4th World Congress on Bioavailability & Bioequivalence

Pharmaceutical R & D Summit

April 15-17, 2013 Sheraton Miyako Hotel Osaka, Japan



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