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Tentative Program

8th International Confernece on

cGMP, GCP & Regulatory Affairs

June 08-09 2018, Baltimore, Maryland, USA

Theme: "Meeting standards of drug regulations & achieving GxP compliance"



For Available Speaker Slots

gmpsummit@pharmaceuticalconferences.org

19+ Interactive Sessions	15+ Keynote Lectures	75+ Plenary Lectures	5+ Workshops

Conference Secretariat

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http://gmp-gcp-quality-control.pharmaceuticalconferences.com

Tentative Program

8th International Confernece on

cGMP, GCP & Regulatory Affairs

June 08-09 2018, Baltimore, Maryland, USA

SCIENTIFIC PROGRAM

DAY 1

Keynote Session

08:45-09:15	Keynote Talk 1
09:15-09:45	Keynote Talk 2
09:45-10:15	Keynote Talk 3
10:15-10:45	Keynote Talk 4

Coffee/Tea Break 10.45-11.00

Pre-Lunch Session	11.00-12.40	*Plenary Session 1| Tracks 1, 2& 3* Good Manufacturing Practices|Current Regulations and Quality Standards| Current GMP Guidelines
Netwroking Lunch 12.40-13.30		
Post Lunch Sessions	13.30-15.30	*Plenary Session 2 | Tracks 4, 5 & 6* GxP in Pharmaceuticals|The Role of c in cGMP| Good Clinical Practices
Coffee/Tea Break 15.30-15.45		
Evening Sessions	15.45-18.00	*Plenary Session 3 | Track 7, 8 & 9* Good Laboratory Practices| Quality Control| Quality Assurance Panel Discussions | Extended Networking

DAY 2

Keynote Session

09:00 - 09:45	Keynote Talk 5
09:45 - 10:15	Keynote Talk 6
10:15 - 10:45	Keynote Talk 7

Coffee/Tea Break 10.45-11.00

Pre-Lunch Session	11.00-12.40	*Plenary Session 4| Track 10, 11 & 12* Validation| Contract & Sterile Manufacturing| Formulation Development
Netwroking Lunch 12.40-13.30		
Post Lunch Sessions	13.30-15.30	*Plenary Session 5 | Tracks 13 &14* GMP in Food Industry| Best Industry Practices|GMP in Microbiology and Biotechnology| Supply Chain Controls
Coffee/Tea Break 15.30-15.45		
Evening Sessions	15.45-18.00	*Plenary Session 6 | Track 15, 16 & 17* Softwares in GMP and GCP| Clinical Affairs and Regulatory Strategies Implementation| Excipient Qualification Panel Discussions | Extended Networking

Closing and Award Ceremony

Glimpses of GMP Summit Series Conferences



















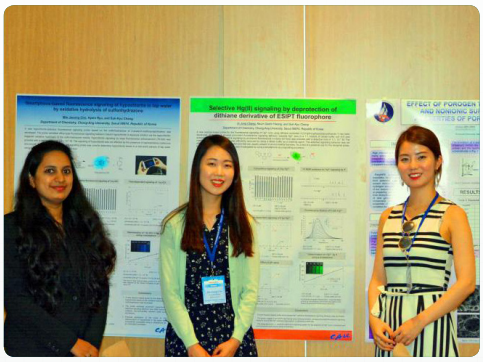






Glimpses of GMP Summit Series Conferences















Major Scientific Sessions

- Good Manufacturing Practices: The Gap within
- Current Regulations and Quality Standards
- Current GMP Guidelines (cGMP) & GxP in Pharmaceuticals
- The Role of c in cGMP
- Good Clinical Practices & Good Laboratory Practices
- Quality Control
- Quality Assurance
- Validation
- Contract & Sterile/Aseptic Manufacturing
- Storage, Distribution, Transportation
- Formulation Development
- GMP in Food Industry
- GMP in Microbiology and Biotechnology
- Softwares in GMP and GCP
- Clinical Affairs and Regulatory Strategies Implementation
- Excipient Qualification and Supply Chain Controls
- Entrepreneurs Investment Meet

Best Poster Award

- You will be given about 5-7 minutes to present your poster including questions and answers. Judges may pose questions during the evaluation of the poster
- Judges will even evaluate the student's enthusiasm towards their study, interest and knowledge in the area of their research
- The winners will be announced at the closing ceremony of the conference. The decision of the winner will be withdrawn if the winner/winners is/are not present at the time of announcement
- Apart from the judging time you may also be present at the poster to share your research with interested delegates





Young Researchers Forum

- Present your research through oral presentations
- Learn about career development and the latest research tools and technologies in your field
- This forum will give pertinent and timely information to those who conduct research and those who use and benefit from research
- Develop a foundation for collaboration among young researchers
- The forum will provide an opportunity for collegial interaction with other young investigators and established senior investigators across the globe
- Interact and share ideas with both peers and mentors

General Queries
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Conference Venue

Chicago, USA

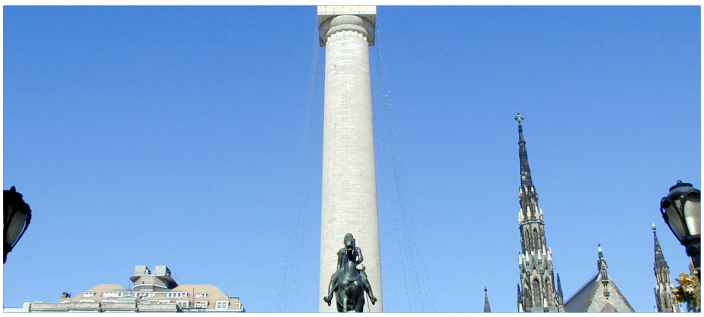
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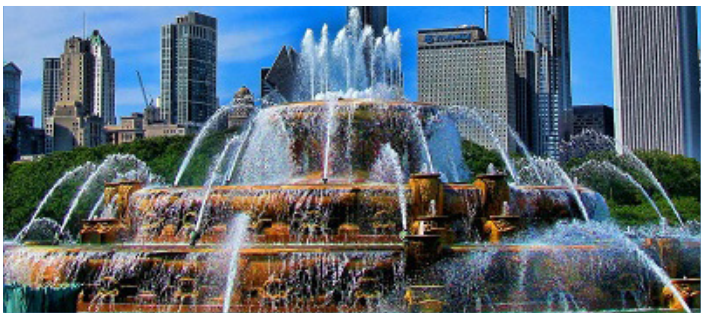
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Harbor place



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Scientifc Program

612th Conference

5th International Summit on

GMP, GCP & Quality Control

August 12-13, 2016
Toronto, Canada





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Conference Day One | Friday August 12, 2016

MacDolald

Registrations

Opening Ceremony

Session Chair: **Boyd L Summers,** BL Summers Consulting. LLC, USA

Keynote Forum

Title: Planning to outsource manufacturing: Have you done your homework?
Mohammed R Khan, Synergex Consulting, Canada

Title: Ensure quality assurance for companies and institutions
Boyd L Summers, BL Summers Consulting LLC., USA

Refreshments and Networking Break 11:00-11:20 @ Foyer

Title: Current FDA audit trends and most common cited drug GMP deficiencies
Kenneth Christie, VTS Consultants, Inc., USA

Title: Importance of characterization of variation in the secondary endpoint measures prior to the trial: a key to a successful outcome of phase 1 trial and progression to a phase
Danuta Radzioch, McGill University and Laurent Pharmaceuticals Inc., Canada

Title: Traceability guide for general food manufacturers
Nadia Narine, Lumar Food Safety Services Ltd., Canada

Lunch Break @ Foyer

Title: Stability considerations from early stage development through phase-IV of pharmaceutical drug products
Dharmi Trivedi, Professional Pharmaceutical Quality and Compliance Specialist, USA

Title: Good clinical practices
Peggy J Berry, Synergy Consulting LLC, USA

B2B Meetings and Networking

Conference Day Two | Saturday August 13, 2016

MacDolald

Session Chair: **Reza Shojaei**, Canadian Plasma Resources, Canada

Keynote Forum

Ramakrishna Pidaparti, Wipro Technologies, USA

Title: Quality risk management
Rashid Mahmood, Surge Laboratories Private Limited, Pakistan

Refreshments and Networking Break 11:00-11:20 @ Foyer

Title: GMP requirements for Canadian blood & blood establishments
Reza Shojaei, Canadian Plasma Resources, Canada

Title: Bioavailability and bioequivalence concerns in pharmaceutical industry
Wael Ebied, SEDICO Pharmaceutical, Egypt

Title: From molecules to market
Luciano Calenti, ACIC Fine Chemicals, Canada

B2B Meetings and Networking

Poster Presentations

Title: Posology in children oral liquid medication studies in Liberia
Jacob Kolawole, University of Liberia, Nigeria

Title: Buccal drug formulation – Pharmacokinetics of Verapamil and its metabolite norverapamil
Wiesław Sawicki, Medical University of Gdansk, Poland

Title: Increase and effectiveness in the activity of audits of quality in the center of genetic engineering and biotechnology
Mariela Díaz Cinza, Havana University, Cuba

Lunch Break @ Foyer

Scientific Program

4th International Summit on

GMP, GCP & Quality Control



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October 26-28, 2015 Hyderabad, India

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Day 1 October 26, 2015

09:00-10:00 Registrations

Hall-1.01 & 1.02

Opening Ceremony

Keynote Forum

Introduction

David Spaulding
SeerPharma, Australia

Shivraj Dasari
SLS Cell Cure Technologies Private Limited, India

Networking & Refreshments Break @ Hall-6

Sunil Kumar Verma
CSIR-Center for Cellular and Molecular Biology, India

Special Session

Peter D Smith
Parexel International, USA

Lunch Break @ Hall-6

Track 1: Good Manufacturing Practices: The Gap within
Track 2: Current Regulations and Quality Standards
Track 3: Current GMP Guidelines (cGMP)
Track 4: The Role of "c" in cGMP

Session Chair: Peter D Smith, Parexel Internationl, USA
Session Chair: Sunil Kumar Verma, CSIR-Center for Cellular and Molecular Biology, India
Session Chair: Shivraj Dasari, SLS Cell Cure Technologies Private Limited, India

Session Introduction

Title: Entropy in Good Manufacturing Practices driven innovative quality tool for pharmaceutical industry: New paradigm approach for manufacturing excellence and quality standards
Pradeep K Jha, IIT Kharagpur, India

Title: Comparison of guidelines of Indian GMP with WHO GMP
Uma Vasireddy, Kakatiya Institute of Pharmaceutical Sciences, India

Title: Role of "c" in cGMP
Abha Doshi, MET Institute of Pharmacy, India

Title: Clinical documentation supporting core labels for generics/OTC products
Aswin Kumar Allupati, Freyr Software Services Pvt Ltd., India

Networking & Refreshments Break

Title: Quality management systems
Vaishali P Nagulwar, Government College of Pharmacy, India

Title: Current GMP guidelines
Abha Doshi, MET Institute of Pharmacy, India

Title: Comparison of regulatory requirements for marketing authorization of biologics in United States and European Union
Shashi Kumar Yadav, Sri Indu Institute of Pharmacy, India

Panel Discussion

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Day 2 October 27, 2015

Hall-1.01 & 1.02

Keynote Forum

Per Nilsson
Profox Company, Sweden

Rama K Pidaparti
Wipro Technologies, USA

Networking & Refreshments Break @ Hall-6

Tanmoy Chakraborty
Manipal University, India

Track 6 & 7: Quality Control & Quality Assurance
Track 8: Validation
Track 9 &11: Contract & Sterile/Aseptic Manufacturing and Formulation Development
Track 12: GMP in Food Industry, Microbiology and Biotechnology

Session Chair: Rama K Pidaparti, Wipro Technologies, USA
Session Chair: Per Nilsson, Profox Company, Sweden

Session Introduction

Title: Food safety management systems-requirements for any organisation in the food chain (ISO 22000:2005)
R Manavalan, Annamalai University, India
Title: HPLC method development and validation as per ICH guidelines
Arunadevi S Birajdar, K T Patil College of Pharmacy, India
Title: Identification of biomarker(s) from polyherbal formulation used in hyperlipidemia for qualitative and quantitative analysis
Charmy S Kothari, Nirma University, India

Lunch Break @ Hall-6

Title: Almighty Astaxanthin: Over view on nutraceutical based approach to aim to combat cancer
Kandra Prameela, GITAM University, India
Title: Effective GMP AUDITS for APIs and Formulation Pharma Companies
G Sundar, PharmQA Compliance Services, India
Title: Pharmaceutical process validation: A tool for pharmaceutical compliance monitor
Sheelpriya Ratnakar Walde, Gurunanak College of Pharmacy, India
Title: Quality Control analytical methods- swith from HPLC to UPLC
Y Padmavathi, G Pullareddy College of Pharmacy, India
Title: Development & characterization of timolol maleate osmotic drug delivery system
B Nagarani , SriKurpa Institute of Pharmaceutical Sciences, India
Title: Effects of lean manufacturing practices to encourage continuous improvement for manufacturing excellence
Dharmvir Uppal, GNA University, India
Title: Quality control methodologies for standardization of herbal medicines: An assessment
Raja Chakaverty, Bengal College of Pharmaceutical Sciences & Research, India

Networking & Refreshments Break @ Hall-6

Poster Presentations @ Hall-6

Poster Judge: Peter D Smith, Parexel Internationl, USA
Poster Judge: R Manavalan, Annamalai University, India

Day 3 October 28, 2015

Hall-1.01 & 1.02

Track 5: Good Clinical Practices & Good Laboratory Practices
Track 10: Storage, Distribution, Transportation

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Session Chair: R Manavalan, Annamalai University, India

Workshop on: How to prevent outbreaks in endemic countries?
Ravi Kumar Tummalacharla, Cleanrooms Containments, India

Networking & Refreshments Break 5 @ Hall-6

Young Researchers Forum

Title: Studies on β-CD complexation of a poorly soluble drug
Mohidurakshan, Sher-I-Kashmir Institute of Medical Sciences, India
Title: Solubility and dissolution rate enhancement of Aceclofenac by solid dispersion technique
Khalid Bashir Mir, The University of Kashmir, India
Title: Good manufacturing practice (GMP): An overview
Firoj A Tamboli, Bharati Vidyapeeth College of Pharmacy, India
Title: A validated UPLC/ESI-MS/MS bioanalytical method for the quantification of Perindopril and Amlodipine in human plasma
Kalaiyarasi Duraisamy, JNTU Hyderabad, India
Title: Adverse drug reaction reporting – A retrospective analysis
Ibel C Fredy, PES College of Pharmacy, India
Title: 3G system in pharmacy practice with vigilance
Khwaja Amtul Raouf Qazi, MRM College of Pharmacy, India

Lunch Break @ Hall-6

Title: Good Clinical Practice (GCP) and declaration of helsinki
Pooja Roy, Vydehi Institute of Medical Sciences and Research Centre, India
Title: Indian clinical trials- The unaddressed challenges of regulatory amendment
N Srinivas, Malla Reddy Institute of Pharmaceutical Sciences, India
Title: Data mining – De-Novo Quality Management tool in Food Sector
S Thiruchenduran, Professor Jayashankar Telangana State Agricultural University, India
Title: Zebrafish as a model system for drug target screening and validation
Bhusnure O G, Channabasweshwar Pharmacy College, India
Title: Estimation of Ramelteon in bulk and tablet dosage form by HPLC
Varaprasad Adepu, JNTU Kakinada, India
Title: QbD approach for the development and optimization of HPLC method for the simultaneous estimation of four component cream formulation: Application to permeability study
Prachi Bhamre, The Maharaja Sayajirao University of Baroda, India
Title: HPLC fingerprinting for quality control of herbal drugs
Nutan Kaushik, The Energy and Resources Institute (TERI), India

Award Ceremony

Bookmark your dates

5th International Summit on

GMP, GCP & Quality Control

August 01-02, 2016 Toronto, Canada

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Scientific Program

3rd International Summit on

GMP, GCP & Quality Control

September 25-26, 2014 Valencia Convention Centre, Spain



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Day 1 September 25, 2014

08:00-09:00 Registrations

Committee Room 5

Opening Ceremony

Keynote Forum

Introduction

Victor Sanchez
Pharma-Bio Serv S.L., Spain

Sundar Chellamani
SysComm Project Management Limited, Ireland

Maria Pellin Amoros
Laboratoires Quinton International S.L., Spain

Coffee Break @ Auditorium 3 Foyer

Rama K Pidaparti
Wipro Technologies, USA

Track 1: Good Manufacturing Practices: The Gap within
Track 2: Current Regulations and Quality Standards
Track 3: Current GMP Guidelines
Track 5: Good Clinical Practices & Good Laboratory Practices

Session Introduction

Session Chair: Magnus Jahnsson, Pharmadule Morimatsu AB, Sweden
Session Co-Chair: Rama K Pidaparti, Wipro Technologies, USA

Workshop on **Natural Health Products site licensing in Canada: How to meet the GMPs regulations**
Jalal Mokhalalati, Quality Medical Regulations Services, Canada

Title: Regulatory requirements and benefits converting to continued process verification
Magnus Jahnsson, Pharmadule Morimatsu AB, Sweden

Title: Greening the pharmaceutical industry to afford Good Laboratory Practice
Salwa Elmeligie, Cairo University, Egypt

Lunch Break @ Multi Purpose Hall 2

Title: Effective methods for software and systems integration for software companies and institutions
Boyd L Summers, BL Summers Consulting, LLC., USA

Title: Reflections about quality control and quality assurance in clinical trials
Fernando Geijo, Telstar, Spain

Title: Role of Good Laboratory Practice in Good Clinical Practice
Salwa Elmeligie, Cairo University, Egypt

Title: Quality excellence through benchmarking quality improvement models
Kamran Atif, Arwan Pharmaceuticals Industries, Lebanon

Track 6: Quality Assurance
Track 7: Quality Control

Session Chairs: Dharmi Trivedi, University of Phoenix, USA
Magnus Jahnsson, Pharmadule Morimatsu AB, Sweden

Title: Ensure quality assurance for software companies and institutions

Boyd L Summers, BL Summers Consulting, LLC., USA

Title: Effective CAPA program, A valuable tool in quality improvement

Dharmi Trivedi, University of Phoenix, USA

Coffee Break @ Auditorium 3 Foyer

Title: Evaluation of bacterial contamination of clean room clothing

Noelle H O Driscoll, Robert Gordon University, Scotland

Title: Optimization solutions for validation procedures in the quality control of enantiomers; Chirality tests for antidepressants Citalopram and Venlafaxine

Ivanka Pencheva, Sofia Medical University, Bulgaria

Title: How to improve quality and consistency of legacy products applying QbD/Six Sigma methodology

Alicia Tebar, Telstar, Spain

Title: Maintain the effectiveness of a QMS by using Lean Six Sigma approach

Peter Jehander, AF Technology AB, Sweden

Title: Analytical method lifecycle management

Gerald de Fontenay, Amatsi Group, France

Panel Discussion

B2B Meetings

Day 2 September 26, 2014

Committee Room 5

Keynote Forum

David L Chesney
Parexel International, USA

Wael Ebied
SEDICO Pharmaceuticals-Merck & Co., Egypt

Aziz Chraibi
Pharma Bio Expert Inc., Canada

Coffee Break @ Auditorium 3 Foyer

Workshop on **Best practices for internal and supplier auditing**

David L Chesney, Parexel International, USA

Track 8: Validation
Track 9: Contract Manufacturing, Sterile/Aseptic Manufacturing
Track 11: Medical Devices

Session Chair: Aziz Chraibi, Pharma Bio Expert Inc., Canada
Session Co-Chair: Jixing Wang, Dalton Pharma Services, Canada

Title: Managing equipment validation using ASTM approach for optimum cost and aggressive schedule

Sundar Chellamani, SysComm Project Management Limited, Ireland

Title: Industrial process validation of metformin tablets to facilitate the scale up of commercial production: A GMP and validation perspective

Gannu Praveen Kumar, Sahasra Institute of Pharmaceutical Sciences, India

Lunch Break @ Multi Purpose Hall 2

Workshop on **ICH Q9 risk management applied to the compliance challenges between cGMP & safety design issues in manufacturing pharmaceutical & biotechnology facilities**

3 Case Studies: High Potent (HP1@5) Plants, Bio-safety containment (BSL1@4) facilities and explosive environment (ATEX 1@3)

Aziz Chraibi, Pharma Bio Expert Inc., Canada

Title: GMP system implementation and certification to manufacture seawater ampoules as a dietary supplement under ISO 5 air quality

Maria Pellin Amoros, Laboratoires Quinton International S.L., Spain

Title: DHF, DMR and DHR - The three Ds of Medical devices

Rama K Pidaparti, Wipro Technologies, USA

Title: Assessing pharmaceutical equipment containment using surrogate monitoring (SMEPAC)

Mootaz El Halawani, Pharmaceutical Quality Expert, Egypt

Coffee Break @ Auditorium 3 Foyer

Title: Challenges of cGMP implementation at different CMO's - role of quality agreements

Rivka Zaibel, Advanced Regulatory Services Ltd. (ADRES), Israel

Title: CMO's challenges and strategies in sterile manufacturing

Jixing Wang, Dalton Pharma Services, Canada

Title: Production of biosimilars in developing countries: Challenges and opportunities: SEDICO case Study

Wael Ebied, SEDICO Pharmaceuticals - Merck & Co., Egypt

Title: The relevance of training in supply chain management of pharmaceutical products

Ibelema Emeh, Setax Training & Consultancy Limited, United Kingdom

Closing Ceremony

B2B

Bookmark your dates



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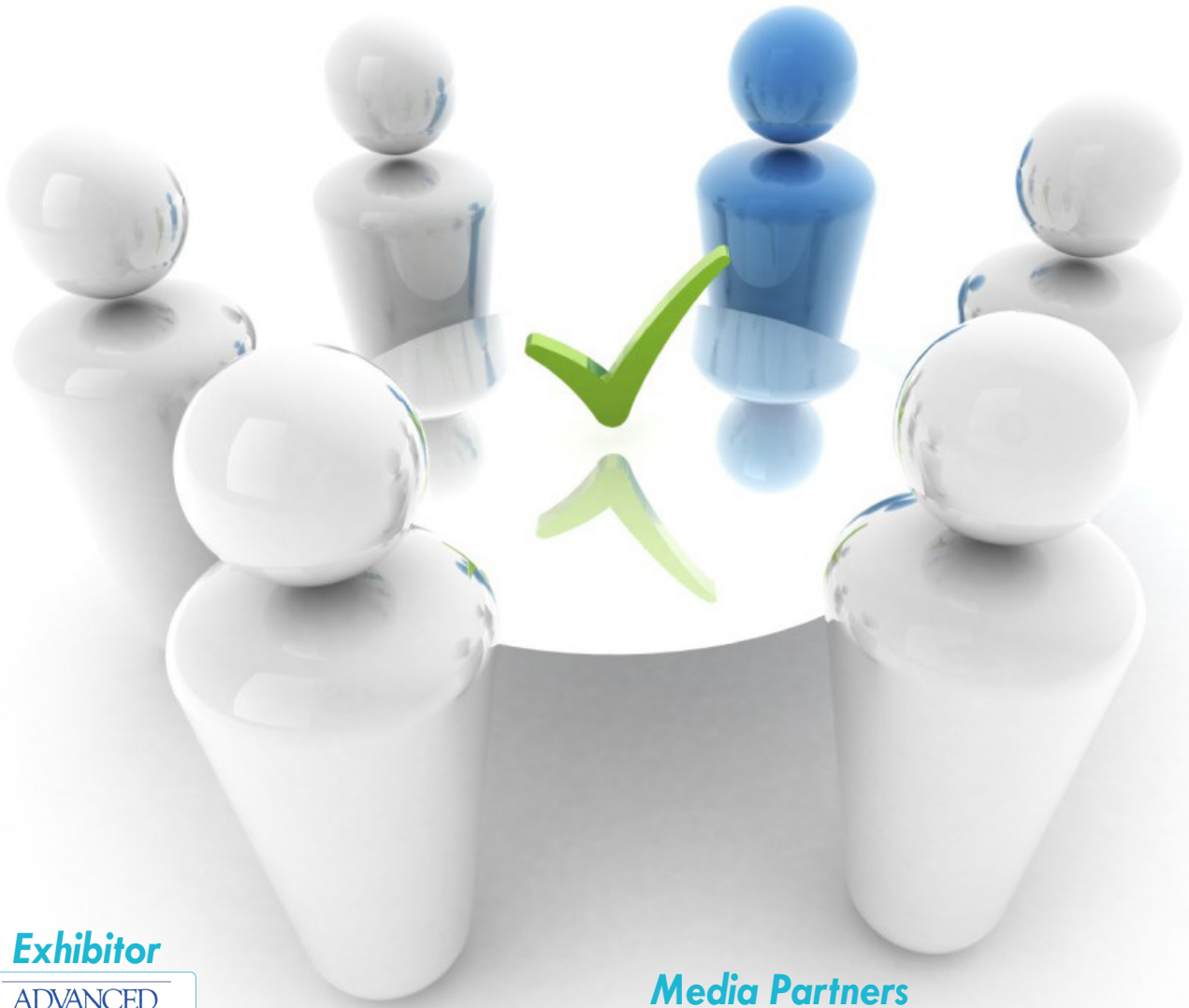
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Scientific Program

2nd International Summit on

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November 12-14, 2013 DoubleTree by Hilton Hotel Chicago-North Shore, IL, USA



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Email: *gmp2013@omicsonline.us*

Day 1 **November 12, 2013**

09:00-10:00 **Registrations**

Berlin

Opening Ceremony

Keynote Forum

Introduction

Gerard Pearce
SQA Services, Inc., USA

Ashraf Youssef
AYPharma Safety Consulting, USA

Coffee Break @ Foyer

Patricia Henley
London School of Hygiene & Tropical Medicine, UK

James Huang
Forest Laboratories, Inc., USA

Track 1: Current Regulations and Quality Standards
Track 2: Current GMP Guidelines for Pharmaceuticals
Track 7: Quality Control, Quality Assurance and Validation

Session Chair: James Huang, Forest Laboratories, Inc., USA

Session Co-Chair: M. S. Lourdes Sanchez, Sintenovo SA de CV, Mexico

Session Introduction

Title: Maintaining your laboratory in a state of Control - The data integrity challenge
Jacqueline McCulloch, Clarkston Consulting, USA

Title: Case study: Quality assurance and quality control partnering to drive compliance, science, and innovation
Jason Ruckert, Advanced Testing Laboratory, USA

Title: Quality assurance on a budget from a non-commercial perspective
Lucy H H Parker, St George's Healthcare NHS Trust, UK

Lunch Break @ Athens

Title: Automation in validated environments
Abitha Sundararajan, Med Manage Systems, USA

Title: Quality accreditation for medical research at Universities?
Brigitte von Rechenberg, University of Zurich, Switzerland

Title: Risk assessment
M.S. Lourdes Sanchez, Sintenovo SA de CV, Mexico

Title: Modern quality systems & Risk management approaches
Naveen Kumar Venkatesham, Laurus Labs Private Limited, India

Title: Diverting from traditional validation approaches - Beefing up and speeding up validation, while minimizing vulnerabilities to new regulatory inspection foci
Constance E. Curts, FDA and EU Regulatory Validation Consultant, USA

Coffee Break @ Foyer

Title: Data integrity issues

Mohammad Iqbal Hossain, Novartis Ltd., Bangladesh

Title: Challenges in benefit-risk evaluations during co-development among partnerships and alliances

Ashraf Youssef, AYPharma Safety Consulting, LLC, USA

Title: Is your company in a state of control?

Brian Hill, Brian Hill & Associates, USA

Title: Prior knowledge assessments: A knowledge compilation process that can be used to bring about more focused process characterization and development studies for platform products

Jim Seely, Bioprocessing Consultant, USA

Title: Mitigating the environmental impact of cleaning processes in GMP regulated facilities

Elizabeth Rivera, STERIS Corporation, USA

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Day 2 November 13, 2013

Berlin

Track 3: The role of "C" in cGMP
Track 8: Legal Requirements for Medical Devices
Track 10: Computational Strategies in GMP/GCP

Session Chair: Pejman Parhami, Hyland's, Inc., USA

Session Co-Chair: Brian Hill, Brian Hill & Associates, USA

Session Introduction

Title: The role of "C" in cGMP

Naveen Kumar Venkatesham, Laurus Labs Private Limited, India

Title: The evolution of the medical device 510 (k) process and impending changes

Jacqueline McCulloch, Clarkston Consulting, USA

Title: Supplier management-Key components of managing suppliers in a cGMP environment

Pejman Parhami, Hyland's, Inc., USA

Title: Bridging the cultural divide: Supplier auditing in a global economy

Jennifer Leny, American Society for Quality, USA

Coffee Break @ Foyer

Title: A risk based scientific approach to analytical method development and validation activities for regulated laboratories

Shib Mookherjea, ValQual International, USA

Title: Moving beyond part 11; Quality assurance considerations for translating cGMP compliance into electronic batch record initiatives

Heather Schwalje, Emerson Life Sciences Industry Solutions Group, USA

Title: Leaderships role in change management

Myriam Ochart, Lean Compliance Partners, USA

Lunch Break @ Athens

Workshops

Zippy Lean Workshop (Production Line Simulation) by

Myriam Ochart, Lean Compliance Partners, USA & **Jennifer Leny,** American Society for Quality, USA

Coffee Break @ Foyer

Highlights of Process Analytical Technology (PAT) & FDA Directives Workshop

Shib Mookherjea, ValQual International, USA

Athens

Poster Presentations

18:30-19:30 Cocktails Sponsored by **Advances in Pharmacoepidemiology & Drug Safety @ Athens**

Day 3 November 14, 2013

Berlin

Track 4: Good Clinical Practice
Track 5: Good Laboratory Practices and Clinical Trials
Track 6: Quality Inspections and Auditing
Track 9: Contract Manufacturing
Track 12: Microbiology, Food and Nutraceuticals

Session Chair: Anthony Grilli, Focus Scientific Solutions, USA

Session Introduction

Title: Performing microbial risk analysis on a standard compounding pharmacy aseptic fill process
Anthony Grilli, Focus Scientific Solutions, USA
Title: The effect of L-arginine on mice placenta
Mohanad AbdulSattar Ali Al-Bayati, University of Baghdad, Iraq
Title: Cleaning challenges in the dietary supplement industry
Elizabeth Rivera, STERIS Corporation, USA

Coffee Break @ Foyer

Title: Myrica rubra fruit drink sub-chronic toxicity and hepatoprotective effect in rats
Mohamed AlAjmi, King Saud University, Saudi Arabia
Title: The informed consent form process: A basic contemporary standpoint for medical research professionals
Peter Odeh, SNBL Clinical Pharmacology Center, USA
Title: DHF, DMR and DHR-The three Ds of medical devices
Rama K. Pidaparti, Wipro Technologies, USA
Title: Immunosuppressive effects of mesenchymal stem cells versus corticosteroid in experimental model of arthritis
Marwa Elhussiny Younnis, Cairo University, Egypt
Title: Comparative pharmacokinetics and compliance issues to optimize art- The Indian scenario
Princy Louis Palatty, Father Muller Medical College, India
Title: A new cGMP bioassay involving electrophysiological recordings of CFTR currents
Antonio Lacerda, ChanTest Corporation, USA

Lunch Break @ Athens

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