Tentative Program

8th International Conference 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

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# Tentative Program

### 8th International Conference on cGMP, GCP & Regulatory Affairs

**June 08-09 2018, Baltimore, Maryland, USA**

## SCIENTIFIC PROGRAM

### DAY 1

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

**09:45-10:15**
- **Plenary Session 1 | Tracks 1, 2 & 3**
  - Good Manufacturing Practices | Current Regulations and Quality Standards | Current GMP Guidelines

**10:15-10:45**
- **Coffee/Tea Break 10:45-11:00**
- **Pre-Lunch Session**
  - 11:00-12:40
  - **Good Manufacturing Practices | Current Regulations and Quality Standards | Current GMP Guidelines**

**11:00-12:40**
- **Networking Lunch 12:40-13:30**
- **Post Lunch Session**
  - 13:30-15:30
  - GxP in Pharmaceuticals | The Role of c in cGMP | Good Clinical Practices

**13:30-15:30**
- **Coffee/Tea Break 15:30-15:45**
- **Evening Session**
  - 15:45-18:00
  - **Good Laboratory Practices | Quality Control | Quality Assurance**

**15:45-18:00**
- **Panel Discussions | Extended Networking**

### DAY 2

**09:15-09:45**
- **Keynote Session**
  - Keynote Talk 5
  - Keynote Talk 6
  - Keynote Talk 7

**09:45-10:15**
- **Plenary Session 2 | Tracks 4, 5 & 6**
  - GMP in Food Industry | Best Industry Practices | GMP in Microbiology and Biotechnology | Supply Chain Controls

**10:15-10:45**
- **Coffee/Tea Break 10:45-11:00**
- **Pre-Lunch Session**
  - 11:00-12:40
  - **Validation | Contract & Sterile Manufacturing | Formulation Development**

**11:00-12:40**
- **Networking Lunch 12:40-13:30**
- **Post Lunch Session**
  - 13:30-15:30
  - GMP in Food Industry | Best Industry Practices | GMP in Microbiology and Biotechnology | Supply Chain Controls

**13:30-15:30**
- **Coffee/Tea Break 15:30-15:45**
- **Evening Session**
  - 15:45-18:00
  - **Softwares in GMP and GCP | Clinical Affairs and Regulatory Strategies Implementation | Equipment Qualification**

**15:45-18:00**
- **Panel Discussions | Extended Networking**

**Closing and Award Ceremony**
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
gmpsummit@pharmaceuticalconferences.org

Conference Venue

Chicago, USA
Best Tourist Destinations in Baltimore

- Washington Monument
- Walters Art Museum of Baltimore
- Johns Hopkins
- Walters Art Gallery
- Harbor place
- Patterson Park
- Oriole Park at
5th International Summit on
GMP, GCP &
Quality Control

August 12-13, 2016
Toronto, Canada

Hosting Organizations: Conference Series LLC
2360 Corporate Circle., Suite 400 Henderson, NV 89074-7722, USA

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Conference Day One | Friday August 12, 2016

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

http://gmp-gcp-quality-control.pharmaceuticalconferences.com
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 Diaz Cinza, Havana University, Cuba

Lunch Break @ Foyer
4th International Summit on GMP, GCP & Quality Control

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October 26-28, 2015  Hyderabad, India

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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

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

Peter D Smith
Parexel International, USA

Networking & Refreshments Break @ Hall-6

**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 Vasierddy, 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

Panel Discussion

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**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 International, 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

Lunch Break @ Hall-6

Session Introduction

**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
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: Overview 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- switch 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 International, 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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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, MMR College of Pharmacy, India

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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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 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

4th International Summit on
GMP, GCP & Quality Control

October 26-28, 2015 - Hyderabad, India
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Proposals are invited for organizing Symposia/Workshops at Conference Series Conferences or Conference Series will sponsor small events at your universities in related areas under the title of your own. These proposals can be sent to respective conference mail ids or to symposia@omicsonline.org

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Email: gmp2013@omicsonline.us
Day 1  
November 12, 2013  

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  

Coffee Break @ Foyer  

**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

Cocktails Sponsored by Journal of Developing Drugs @ Athens

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

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, VaQual 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

Highlights of Process Analytical Technology (PAT) & FDA Directives Workshop
Shib Mookherjea, VaQual 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 Younus, 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

Bookmark your dates

3rd International Summit on
GMP, GCP & Quality Control

September 25-26, 2014   Valencia, Spain