2nd International Conference on
Clinical Trials
August 22-24, 2016
Philadelphia, USA

Supporting Sponsors

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47 Churchfield Road, London, UK, W3 6AY
Toll Free: +1-800-014-8923
**Day 1**

**August 22, 2016**

**Independence A**

**Keynote Forum**

<table>
<thead>
<tr>
<th>Time</th>
<th>Title</th>
<th>Speaker</th>
<th>Institution</th>
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<tbody>
<tr>
<td>09:00-09:25</td>
<td><strong>Introduction</strong></td>
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<tr>
<td>09:25-09:50</td>
<td><strong>Title: Patient centricity – A patient’s perspective</strong></td>
<td><strong>Michael Bernstein</strong>, Merck &amp; Co., Inc., USA</td>
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<tr>
<td>09:50-10:15</td>
<td><strong>Title: Why conferences and continued training really matter</strong></td>
<td><strong>Jody Ehrhardt</strong>, Ehrhardt Clinical Research, USA</td>
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<tr>
<td>10:15-10:35</td>
<td><strong>Title: Preclinical and clinical research in wound healing and hyperbaric medicine</strong></td>
<td><strong>Thomas E Serena</strong>, SerenaGroup, USA</td>
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<tr>
<td>10:35-10:55</td>
<td><strong>Title: A mixture model using likelihood-based and Bayesian approaches for identifying responders and non-responders in longitudinal clinical trials</strong></td>
<td><strong>Entsuah Anthony Richard</strong>, Merck &amp; Co., Inc., USA</td>
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<tr>
<td>11:10-11:30</td>
<td><strong>Title: The Burden of Therapy© [BOTH©]. A novel method to evaluate the impact of therapeutic agents on patient safety</strong></td>
<td><strong>Ayad K Abdul-Ahad</strong>, BOTH Analytics GmbH, Germany</td>
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<tr>
<td>11:30-11:50</td>
<td><strong>Title: A framework for investigating scientific and medical research misconduct and fraud</strong></td>
<td><strong>Tamera Norton Smith</strong>, Norton Audits, Inc., USA</td>
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<td>11:50-12:10</td>
<td><strong>Title: Establishing a standard practice to communicate trial results to study volunteer</strong></td>
<td><strong>Jill McNair</strong>, CISCRP, USA</td>
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<td>12:10-12:30</td>
<td><strong>Title: Regulatory and compliance considerations for Direct-To-Patient (DTP) distribution of clinical trial material</strong></td>
<td><strong>Michael Bernstein</strong>, Clinical Supplies Management Inc., USA</td>
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<tr>
<td>12:30-13:00</td>
<td><strong>Title: Wound healing characteristics treated by two newly developed amino acid-based biodegradable polymers in burn and incision porcine wound models</strong></td>
<td><strong>Chih-Chang Chu</strong>, Cornell University, USA</td>
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<td>13:00-13:40</td>
<td><strong>Lunch Break</strong></td>
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<td>13:40-14:00</td>
<td><strong>Title: Bayesian Decision-Optimal Interval (BOIN) designs for Phase I clinical trials</strong></td>
<td><strong>Suyu Liu</strong>, The UT MD Anderson Cancer Center, USA</td>
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<tr>
<td>14:00-14:20</td>
<td><strong>Title: The importance of subject recruitment and retention to clinical trial success</strong></td>
<td><strong>Jody Ehrhardt</strong>, Ehrhardt Clinical Research, USA</td>
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<td>14:20-14:40</td>
<td>Title: Good pharma scorecard-A reform strategy to drive improvement clinical trial transparency</td>
<td>Jennifer Miller</td>
<td>NYU Langone Medical Center, USA</td>
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<td>14:40-15:00</td>
<td>Title: Re-engineering the clinical development process</td>
<td>Anvita Karara</td>
<td>Carnegie Mellon University, USA</td>
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<td>15:00-15:40</td>
<td>Workshop: Clinical research site eSource Readiness Assessment tool (eSRA)- A free tool to assist sites in determining if their systems are ready for regulated clinical research data</td>
<td>Suzanne Bishop</td>
<td>North American Facilitator for the eClinical Forum, USA</td>
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<tr>
<td>15:55-16:15</td>
<td>Title: Low temperature plasmas at atmospheric pressure towards new pharmaceutical products in medicine</td>
<td>N Merbahi</td>
<td>University of Toulouse, France</td>
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<td>16:15-16:35</td>
<td>Title: High patient satisfaction rating for clinical medication counseling services when provided by clinically trained pharmacists</td>
<td>Joe Martinez</td>
<td>Center Point Clinical Services, USA</td>
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<td>16:35-16:55</td>
<td>Title: Functional identification of a CRF-DA microcircuit in mice with relevance to drug abuse</td>
<td>Kyle Gobrogge</td>
<td>Tufts University, USA</td>
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<tr>
<td>16:55-17:15</td>
<td>Title: Big data analytics – Clinical integration and visualization</td>
<td>Ankit Lodha</td>
<td>University of Redlands - School of Business, USA</td>
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<tr>
<td>09:30-09:55</td>
<td>Keynote Forum: Golden Era of Research</td>
<td>Tamera Norton Smith</td>
<td>Norton Audits, Inc., USA</td>
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<td>09:55-10:20</td>
<td>Title: Medical heroes science museum initiative: An innovative approach to patient education</td>
<td>Jill McNair</td>
<td>CISCRP, USA</td>
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<tr>
<td>10:20-10:45</td>
<td>Title: The power of translational research</td>
<td>Thomas E Serena</td>
<td>SerenaGroup, USA</td>
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<td>11:00-11:20</td>
<td>Title: Virucidal capacity of novel ProtecTeaV sanitizer formulations containing lipophilic Epigallocatechin-3-Gallate (EGCG)</td>
<td>Stephen Hsu</td>
<td>Augusta University, USA</td>
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<tr>
<td>11:20-11:40</td>
<td>Title: Response-Adaptive Randomization urn designs in clinical trials</td>
<td>Annalisa Piccorelli</td>
<td>University of Wyoming, USA</td>
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11:40-12:00  
Title: Photodynamic therapy to treat diabetic foot infection  
Joao Paulo Tardivo, Faculdade de Medicina do ABC, Brazil  

Title: Histoscanning™ for detection of prostate cancer: Does it have any role in routine clinical practice?  
Saqib Javed, Leighton Hospital NHS Foundation Trust, UK

Workshop

12:20-13:00  
Title: Combating scientific and medical research misconduct and fraud  
Tamera Norton Smith, Norton Audits, Inc., USA

Lunch Break 13:00-13:40 @ Benjamin’s

13:40-14:00  
Title: Using Artificial Intelligence to Measure and Optimize Adherence in Clinical Trials  
Gordon Kessler, AiCure, USA

14:00-14:20  
Title: Adverse renal effects of novel molecular oncologic targeted therapies: A review of the FDA Adverse Event Reporting System (FAERS)  
Rimda Wanchoo, Hofstra Northwell School of Medicine, USA

14:20-14:40  
Title: Preliminary in vivo screening of experimentally induced gastric ulcer of Ledebouria Ovatifolia  
Eugene Jamot Ndebia, Walter Sisulu University, South Africa

14:40-15:00  
Title: A Modern In-vivo pharmacokinetic (PK) paradigm: Combining snapshot, rapid and full PK approaches to support early drug discovery  
Chun Li, Genomics Institute of the Novartis Research Foundation, USA

15:00-15:20  
Title: Wild leafy vegetable Mormodica foetida, improves metabolic syndrome markers and sperm parameters in diet induced obese male rats  
Constance R Sewani-Rusike, Walter Sisulu University, South Africa

15:20-15:40  
Title: Retrospective analysis of clinical pharmacist medication counseling to improve patient medication compliance and patient retention  
Joe Martinez, Center Point Clinical Services, USA

15:40-16:00  
Title: SharePoint development and implementation in a tertiary care center in Lebanon  
Ulfat Usta Shanouha, American University of Beirut Medical Center, Lebanon

Networking and Refreshments 16:00-16:15 @ Foyer

16:15-16:35  
Title: Reflection of knowledge and concern of ethical principles among biomedical postgraduates: A cross sectional observational study from India  
Krishnan Vengada ragava Chary, Saveetha University, India

Poster Presentations 16:35-17:05 @ Foyer

CLTRS001  
Title: Magnitude and characteristics of clinical trials in Saudi Arabia: A cross-sectional analysis  
Sheraz Ali, King Saud Medical City, KSA

CLTRS002  
Title: Monitoring long term plasma Ochratoxin A levels of female rats: Fluctuating OTA levels and toxicodynamics  
Mehmet Akif Kilic, Akdeniz University, Turkey

CLTRS003  
Title: Assessment of awareness about ethics committee amongst the research scholars/teachers in government medical colleges of punjab, India  
Ramandeep Kaur Brar, Baba Farid University of Health Sciences, India

CLTRS004  
Title: Emerging role of bioinformatics tools and softwares in evolution of clinical research  
Supreet Kaur Gill, Baba Farid University of Health Sciences, India

Panel Discussions

Day 3 August 24, 2016

Independence A

Sessions:

Bioethics and Quality Regulation | Pharmacokinetics and Pharmacodynamics | Toxicogenomics Challenges and Applications | Pharmacovigilance and Drug Safety

Session Chair: Nicola Stagg, Genentech, USA

Session Introduction

09:30-09:50
Title: Gender differences in blood pressure and electrocardiography parameters in response to antihypertensive treatment supplemented with dietary flavonoids
Marina M J Romero-Prado, University of Guadalajara, Mexico

09:50-10:10
Title: Important considerations for successful Direct-to-Patient study implementation
Joe Martinez, Center Point Clinical Services, USA

10:10-10:30
Title: Protective effect of Hypericum triquetrifolium Turra on Cyclophosphamide induced cardiotoxicity in rat
Songul Cetik, Mardin Artuklu University, Turkey

10:30-10:50
Title: Peripheral neuropathy with microtubule inhibitor containing antibody drug conjugates: Challenges and perspectives in translatability from nonclinical toxicology studies to the clinic
Nicola Stagg, Genentech, USA

Networking and Refreshments 10:50-11:05 @ Foyer

11:05-11:25
Title: Proposal of supervised data analysis strategy of plasma miRNAs from hybridisation array data with an application to assess hemolysis-related deregulation
Elena Landoni, University of Milan, Italy

11:25-11:45
Title: Clinical research during the Ebola virus disease outbreak in guinea: Lessons learned and ways forward
Abdoul Habib Beavogui, Rural Health of Maferinyah, Guinea

11:45-12:05
Title: Use of investigative sites in Central & Eastern Europe (CEE) for successful clinical studies
Jeffrey Blum, EastHORN Clinical Services, USA

12:05-12:25
Title: Determination of glyceryl trinitrate and its two main metabolites in human plasma using a new sensitive gas chromatography method
Mohammadreza Sattari, Tabriz University of Medical Sciences, Iran

Awards & Closing Ceremony

Lunch Break 12:30-13:10 @ Benjamin’s

Bookmark your dates

3rd International Conference on Innovations to Advanced Clinical Research and Clinical Trials
June 19-20, 2017 | London, UK

4th International Conference on Clinical Trials
September 11-13, 2017 | San Antonio, USA