



Presents

## The 17<sup>th</sup> Annual Fall Conference

### Fresh Perspectives on Recruitment, Regulations and Research

**Dates:** October 21 & 22, 2016

**Time:** 8:00 AM – 5:30 PM (see the detailed agenda at the end of this flyer)

Networking Event October 21, 2016 from 5:30 PM – 7:30 PM sponsored by Covance

**Location:** North Carolina Biotech Center, 15 TW Alexander Drive, Durham, NC, 27709

**Program Description:** The Fresh Perspectives in Recruitment, Regulations, and Research is a two day conference that will include best practices on engaging participants in clinical trials, ethical/compliance considerations associated with regulatory changes, as well as provide insight into topics at the forefront of today's clinical research. The expert presenters will enable clinical research professionals to become knowledgeable about current regulations and the evolving research landscape.

**Objectives:** Upon completion of the conference, attendees should be able to:

1. Demonstrate successful patient recruitment methods using the latest outreach techniques
2. Interpret recent regulatory changes and recognize their implications
3. Distinguish the newest trends in clinical research
4. Gain knowledge of how to be successful in managing clinical research trials.

**Target Audience:** Clinical Research Professionals including: Clinical Research Coordinators, Clinical Research Associates, Investigators, Auditors, Project Managers, Regulatory, Data Managers, Quality Assurance and Clinical Research Administrators

**Registration Cost:** Please use this link to register for the conference. Link to PayPal will be added. You will need to purchase contact hours separately.

RTP Chapter Member	\$125
ACRP Member	\$155
Non-Member	\$175
Student/Volunteer	\$75
One day registration	\$90

**Contact Hours:** 12.25 Contact Hours have been applied for through ACRP. Membership is not required for online registration of contact hours. Contact Hour Cost: \$0 RTP Chapter Member / \$35 ACRP Members / \$50 Non-Members

**To receive contact hours:** Purchase the contact hours separately through ACRP, sign in at the registration desk and attend the program. Log on the ACRP website then “ACRP Learning Portal” to complete the evaluation no later than 30 days following the event and obtain the online certificate. To secure the contact hours the attendee must do all three things – sign in during the conference, purchase the contact hours and complete the evaluations.

**Refund Policy:** There are no refunds unless the program is cancelled by the chapter before the start of the program.

**Level:** Intermediate

### **Program Agenda**

#### **17<sup>th</sup> Annual Fall Conference Fresh Perspectives on Recruitment, Regulations and Research:**

##### ***Friday, October 21, 2016***

8:00am - 8:45am - Registration

8:45am - 9:00am - Welcome

9:00am - 10:00am - **Liz Wool, RN, BSN, CCRA, CMT** Global Head of Training Barnett International. “ICH E6 (R2) Proposed Revisions 2016: A Targeted Overview”

10:00am - 11:00am - **Debbie Rosenbaum, BS, CCRC, CCRA** Sarrison Clinical Research and **Patty Darragh, MS, CCRA** Duke Clinical Research Institute “Exploring Challenges in the Clinical Research Profession”

11:00am - 11:30am - Break

11:30am - 12:30pm - **Juanita Cuffee MPH** Clinical Research Coordinator, Peds-Hematology/Oncology, University of North Carolina – Chapel Hill Chair – UNC Network of Research Professionals “Managing Study Compliance: A Site Perspective”

12:30pm - 1:30pm - Lunch

1:30pm - 2:30pm - **David Resnik J.D., Ph.D.** Bioethicist and NIEHS IRB Chair “Paying Research Participants: Ethical Issues”

2:30pm -3:30pm - **Charles Carter, BS, PharmD, MBA**, Associate Professor, Department of Clinical Research College of Pharmacy & Health Sciences Campbell University and **Melissa A. Holland, PharmD, MSCR**, Assistant Professor, Department of Clinical Research College of Pharmacy & Health Sciences Campbell University “Therapeutic Frontiers of Clinical Research”

3:30pm – 4:00pm - Break

4:00pm – 5:00pm - **Jonathan Zung PhD** Group President, Clinical Development and Commercialization Services for Covance Drug Development “The Clinical Trial of the Future”

5:00pm - 5:30pm - Closing remarks

5:30pm – 7:30pm – Networking event sponsored by Covance

### ***Saturday, October 22, 2016***

8:00am – 8:45am - Registration

8:45am – 9:00am - Welcome

9:00am – 10:00am - **Robert Romanchuk, BSHS, CIP, CCRCP, CCRC**, Vice Chair Schulman IRB “Return of Research Results to Participants: Ethical and Operational Challenges”

10:00am – 11:00am - **Lori Bruhns, BA** “Power-Up your Productivity”

11:00am - 11:30am - Break

11:30am - 12:30pm - **Julie Blasingim MBA** Director of Operations, Clinical Pharmacology and Institutional Research Services, Schulman IRB and **Kirsten Messmer, PhD** Principal Regulatory Affairs Specialist, PPD

12:30pm - 1:30pm - Lunch

1:30pm - 2:15pm - **Jim Kremidas**, Executive Director ACRP “Celebrating Research Professionals”

2:15pm - 3:15pm - **Douglas E. Pierce, Jr.** President Clinical Ink

3:15pm - 3:30pm - Break

3:30pm – 5:00pm – Expert Panelist discussion and question and answer **Social Media Use in Clinical Research**

5:00pm - 5:30pm - Closing remarks