

**Broad Agency Announcement Industry Day**  
**U.S. Food and Drug Administration**  
**Department of Health and Human Services**  
**March 25, 2013**  
**White Oak Campus**  
**10903 New Hampshire Avenue**  
**Silver Spring, MD 202993**  
**Building 2 Room 2047E**  
**10:00am – 12:00pm**

**AGENDA**

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| I.   | Opening Remarks  | Sean Wybenga, Contracting Officer<br>Office of Acquisitions and Grants                                     |
| II.  | Background of Advancing Regulatory Science Plan  | Dr. Frank Weichold, M.D, Ph.D<br>Director<br>Office of Critical Path and Regulatory Science and Innovation |
| III. | Overview of the BAA Review Process   | Robyn Barringer, PMP<br>BAA Program Manager<br>Office of Regulatory Science and Innovation                 |
| IV.  | Research Area Panel Discussion<br>(Each Research Area is limited to 8 min)                     |  |
|      | i. Social and Behavioral Science   | Miriam Campbell, Ph.D, MPH<br>Office of Planning/Risk Communications Staff                                 |
|      | ii. Pharmaceutical Quality Surveillance<br>Harvesting Clinical Study Data<br>Pharmacovigilance | Binh Ta, PMP<br>Center for Drug Evaluation and Research, Office of Business Informatics                    |
|      | iii. Medical Counter Measures  | Tracy MacGill, Ph.D. CDR, USPHS<br>Office of Counterterrorism and Emerging Threats                         |
|      | iv. Harnessing Diverse Data  | Ross Filice, M.D.<br>Office of Regulatory Science and Innovation   |
|      | v. Innovation in product development and evaluation  | Nancy Collazo-Braier, Ph.D.<br>Center for Devices and Radiological Health, Office of the Center Director   |

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| vi.   | Supporting New Approaches to Improve Product Manufacturing and Quality | Monica L. Young, Ph.D.<br>Center for Biologics Evaluation & Research, Office of the Director                 |
| vii.  | Modernizing toxicology and risk assessment to better predict safety    | Dr. Peggy Miller<br>Associate Director for Regulatory Activities, National Center for Toxicological Research |
| viii. | Questions/Comments   | ALL  |

**REGISTRATION INFORMATION:** Pre-registration for this event is required. Please send e-mails indicating attendance to: [FDABAA@fda.hhs.gov](mailto:FDABAA@fda.hhs.gov).

\*\* All registration e-mails are due no later than Thursday, March 21, 2013. \*\*

**CONFERENCE ATTENDANCE:** Please be sure to bring government issued ID card along for building access. Visitor parking is available across from Building 51 and is accessible via Southwest loop road. Additional visitor parking can be accessed the Northwest Parking Garage across from Building 22. A WO campus parking map is included to assist with planning your visit.

**POC Information:**

Robyn Barringer

E-mail: [robyn.barringer@fda.hhs.gov](mailto:robyn.barringer@fda.hhs.gov)

Telephone: 301-796-8485