2010 Cosponsored Events

1. 1st Annual Patient-Reported Outcomes (PRO) Consortium Workshop, Co-sponsored by: Food and Drug Administration and Critical Path Institute.
2. 5th Annual FDA/Drug Information Association Statistics Forum, Co-sponsored by: Center for Drug Evaluation and Research and Drug Information Association.
3. 13th Annual FDA-OCRA Educational Conference, Co-sponsored by: Office of Regulatory Affairs Los Angeles District and Orange County Regulatory Affairs Discussion Group.
4. 13th International Paul-Ehrlich-Seminar, Co-sponsored by: Center for Biologics Evaluation and Research, Paul-Ehrlich-Institute and Drug Information Association.
5. 14th Symposium on the Interface of Regulatory and Analytical Sciences for Biotechnology Health Products - "WCBP 2010," Co-sponsored by: Center for Drug Evaluation and Research and California Separation Science Society.
6. 2010 AAMI/FDA Infusion Device Summit, Co-sponsored by: Center Devices and Radiological Health and Association for the Advancement of Medical Instrumentation.
7. 2010 PDA/FDA Vaccine Conference, Co-sponsored by: Center for Biologics Evaluation and Research and Parenteral Drug Association.
8. AAPS Workshop on Advances and Opportunities in Drug Product Manufacturing - A Look at Continuous Manufacturing Process, Co-sponsored by: Center for Drug Evaluation and Research and American Association of Pharmaceutical Scientists.
9. Accelerating Anticancer Agent Development and Validation Workshop, Co-sponsored by: Center for Drug Evaluation and Research and Duke Comprehensive Cancer Center, Duke University Medical Center, Duke University.
10. Adaptive Design for Clinical Trials: FDA Draft Guidance Symposium, Co-sponsored by: Center for Drug Evaluation and Research and Drug Information Association.
11. Antisense Oligonucleotide (AON) Therapies in Neuromuscular Diseases Workshop, Co-sponsored by: Center for Drug Evaluation and Research, National Institutes of Health, Parent Project Muscular Dystrophy, Muscular Dystrophy Association and Children's National Medical Center.
12. Biomarkers in Gaucher's Disease Workshop, Co-sponsored by: Center for Drug Evaluation and Research and National Gaucher Foundation.
13. Cardiac Safety Research Consortium (CSRC) Annual Meeting and Pediatrics Initiative Thinktank Workshop, Co-sponsored by: Food and Drug Administration and Duke University.
14. Cardiovascular Safety in Drug Development: QT, Arrhythmias, Thrombosis, and Bleeding, Co-sponsored by: Center for Drug Evaluation and Research and Drug Information Association.
15. Clinical Investigator Training Course, Co-sponsored by: Food and Drug Administration and Duke University.
16. Computer Methods in Cardiovascular Device Design & Evaluation Workshop, Co-sponsored by: Center for Devices and Radiological Health, National Institutes of Health, Stanford University and Massachusetts Institute of Technology.
17. Debate on Clinical Trials for Pulmonary Arterial Hypertension, Co-sponsored by: Center for Drug Evaluation and Research, Drug Information Association and Pulmonary Vascular Research Institute.

18. Development of Type 2 Diabetes Mellitus Drugs: State of the Art Cardiovascular Safety Assessments, Co-sponsored by: Center for Drug Evaluation and Research and Drug Information Association.
19. Drug Information Association/FDA Orphan Drug Designation Workshop, Co-sponsored by: Food and Drug Administration and Drug Information Association.
20. Digestive System Motility Workshop, Co-sponsored by: Center for Drug Evaluation and Research, American Neurogastroenterology and Molility Society.
21. Do A Designation Workshop, Co-sponsored by: Food and Drug Administration, National Organization for Rare Disorders and Genetic Alliance.
22. Do A Designation Workshop, Co-sponsored by: Food and Drug Administration, Keck Graduate Institute of Applied Life Sciences and Center for Orphan Drug Research, College of Pharmacy, University of Minnesota.
23. Drug-Induced Liver Injury: Getting the Medicine and Science Together Workshop, Co-sponsored by: Center for Drug Evaluation and Research, American Association for the Study of Liver Diseases and Pharmaceutical Research and Manufacturers Association.
24. Drugs & Medical Devices Supplier Management Forum, Co-sponsored by: Office of Regulatory Affairs Baltimore District and Association of Food and Drug Officials.
25. FDA and PAT for Pharma Manufacturing, Co-sponsored by: Center for Drug Evaluation and Research and University of Rhode Island.
26. FDA/Continua/CIMIT Workshop on Medical Device Interoperability, Co-sponsored by: Center for Devices and Radiological Health, Continua Health Alliance and Center for Integration of Medicine and Innovative Technology.
27. FDA Device Requirements Seminar, Co-sponsored by: Office of Regulatory Affairs, Cincinnati District and Xavier University.
28. FDA - ISPE Collaboration: Pharmaceutical Quality Systems Seminar, Co-sponsored by: Center for Drug Evaluation and Research and International Society for Pharmaceutical Engineering.
29. FDA / SoCRA Clincial Trials Seminar, Co-sponsored by: Office of Regulatory Affairs Florida District Office and Society of Clinical Research Associates, Inc.
30. FDA / SoCRA Clinical Trials Seminar, Co-sponsored by: Office of Regulatory Affairs Pacific Region and Society of Clinical Research Associates, Inc.
31. FIP Pharmaceutical Sciences World Congress 2010 in Association with the AAPS Annual Meeting & Exposition, Co-sponsored by: Center for Drug Evaluation and Research, American Association of Pharmaceutical Scientists and International Pharmaceutical Federation.
32. Food Labeling Workshop, Co-sponsored by: Office of Regulatory Affairs Southwest Region and The Robert M. Food & Agricultural Products Center.
33. Food Labeling Workshop, Co-sponsored by: Office of Regulatory Affairs Southwest Region and University of Arkansas.
34. Food Protection Workshop, Co-sponsored by: Office of Regulatory Affairs Southwest Region and University of Arkansas.
35. Global Outsourcing Challenges Seminar, Co-sponsored by: Office of Regulatory Affairs Cincinnati District and Xavier University.
36. Improved Clinical Trials in Imaging: Evaluation of Breast Imaging and CADs (Computer Assisted Devices) Workshop, Co-sponsored by: Center for Devices and Radiological Health and Medical Image Perception Society.
37. International SAEC Phenotype Standardization Conference, Co-sponsored by: Center for Drug Evaluation and Research, Wellcome Trust and International SAE Consortium Ltd.
38. IVD Roundtable 501(k) Workshop, Co-sponsored by: Association of Diagnostics Manufacturers and Center for Devices and Radiological Health.

39. Lessons Learned from Potency Testing of Pandemic (H1N1) 2009 Influenza Vaccines and Considerations for Future Potency Tests Workshop, Co-sponsored by: Center for Biologics Evaluation and Research, Health Canada and World Health Organization.
40. Medical Devices Technology Innovation Workshop, Co-sponsored by: Center for Devices and radiological Health and Rector and Visitors of the University of Virginia.
41. Optimizing Clinical Trail Design for the Development of Pediatric Cardiovascular Devices Workshop, Co-sponsored by: Center for Devices and Radiological Health, American Academy of Pediatrics, American College of Cardiology and Society for Cardiovascular Angiography and Interventions.
42. PDA/FDA Adventitious Virus Workshop, Co-sponsored by: Center for Drug Evaluation and Research, Center for Biologics Evaluation and Research, Center for Devices and Radiological Health and Parenteral Drug Association.
43. PDA/FDA Pharmaceutical Ingredient Supply Chain Workshop, Co-sponsored by: Center for Drug Evaluation and Research and Parenteral Drug Association.
44. Pediatric-Inclusive or Pediatric Focused Safety Databases Meeting, Co-sponsored by: FDA and Critical Path Institute.
45. Process Drift: Detection, Measurement, and Control in the Manufacture of Pharmaceuticals Workshop, Co-sponsored by: Center for Drug Evaluation and Research and Product Quality Research Institute.
46. Public Workshop on Antibacterial Resistance and Diagnostic Device and Drug Development Research for Bacterial Diseases, Co-sponsored by: Center for Drug Evaluation and Research, Center for Devices and Radiological Health, National Institute of Allergy and Infectious Diseases and Infectious Diseases Society of America.
47. Quality by Design (QbD) Topical Conference, Co-sponsored by: Center for Drug Evaluation and Research and American Institute of Chemical Engineers.
48. Quality System Education Forum on Risk Management, Co-sponsored by: Office of Regulatory Affairs Dallas District and FDA Medical Device Industry Coalition, Inc.
49. Scientific Workshop: Patient-Reported Outcomes in Chronic Kidney Disease, Co-sponsored by: Food and Drug Administration and National Kidney Foundation.
50. Statistics and Medical Devices Conference, Co-sponsored by: Center for Devices and Radiological Health and Advanced Medical Technology Association's Medical Technology Learning Institute.
51. The Essentials of FDA Device Regulations: A Primer for Manufacturers and Suppliers Seminars, Co-sponsored by: Office of Regulatory Affairs, Center for Device Evaluation and Research and Advanced Medical Technology Association.
52. The Great Lakes cGMP & Regulatory Science Forum, Co-sponsored by: Office of Regulatory Affairs and University of Illinois Chicago.
53. Thermal Aspects of Radio Frequency Exposure Workshop, Co-sponsored by: Center for Devices and Radiological Health, Mobile Phone Manufacturers Forum and Global System for Mobile Communications Association.