2011 Co-sponsored Events

1. 1st Scientific Workshop: Analgesic Clinical Trial Innovations, Opportunities, and Networks Initiative, Co-sponsored by: Center for Drug Evaluation and Research and University of Rochester.
2. 2nd Annual Patient-Reported Outcome (PRO) Consortium Workshop, Co-sponsored by: Center for Drug Evaluation and Research and Critical Path Institute.
3. 6th Annual FDA / Drug Information Association Statistics Forum, Co-sponsored by: Center for Drug Evaluation and Research and Drug Information Association.
4. 14th Annual FDA - Orange County Regulatory Affairs Educational Conference, Co-sponsored by: Office of Regulatory Affairs Los Angeles District and Orange County Regulatory Affairs Discussion Group.
5. 15th Symposium on the Interface of Regulatory and Analytical Sciences for Biotechnology Health Products - "WCBP 2011," Co-sponsored by: Center for Drug and Evaluation Research and California Separation Science Society.
6. 2011 Association for the Advancement of Medical Instrumentation/FDA Medical Device Reprocessing Summit, Co-sponsored by: Center for Devices and Radiological Health and Association for the Advancement of Medical Instrumentation.
7. 2011 Great Lakes cGMP & Regulatory Science Forum, Co-sponsored by: Office of Regulatory Affairs and University of Illinois Chicago.
8. 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.
9. Aiming High for Compliance, Co-sponsored by: Office of Regulatory Affairs and Pharmaceutical Industry Association of Puerto Rico.
10. Cardiac Disease and Safety in Clinical Research: Integrating Cardiology and Oncology Clinical Trials with Practice Workshop, Co-sponsored by: Food Drug Administration and Duke University.
11. Cardiovascular Safety in Drug Development: State-of-the-art Assessments, Co-sponsored by: Center for Drug Evaluation and Research and Drug Information Association.
12. Clinical Investigator Training, Co-sponsored by FDA and Duke University.
13. Comprehensive Quality by Design (QbD) in Pharmaceutical Development and Manufacture Topical Conference, Co-sponsored by: Center for Drug Evaluation and Research and American Institute of Chemical Engineers.
14. Creating Consensus Science: New Tools and Tactics for Next-Gen Drug Development, Co-sponsored by: Center for Drug Evaluation and Research, Critical Path Institute and Clinical Data Interchange Standards Consortium.
15. Device Development in Obesity: Clinical and Outcomes Assessment Workshop, Co-sponsored by: Center for Devices and Radiological Health, Dartmouth Device Development/GI, Dartmouth Medical School and Obesity, Metabolism & Nutrition Institute, Massachusetts General Hospital.
16. Drug Information Association/FDA Best Practices for Regulatory Information Synthesis of Randomized Controlled Trials for Product Safety Evaluation, Co-sponsored by: Center for Drug Evaluation and Research and Drug Information Association.

17. Drug Information Association/FDA CDER/CBER Computational Science Annual Meeting, Co-sponsored by: Center for Drug Evaluation and Research, Center for Biologics Evaluation and Research and Drug Information Association.
18. Drug Information Association/FDA Medical Device Quality System Workshop, Co-sponsored by: Food and Drug Administration and Drug Information Association.
19. Drug Information Association/FDA Orphan Drug Designation Workshops - InDrug Information Association, Co-sponsored by: Food and Drug Administration and Drug Information Association.
20. Drug Information Association/FDA Orphan Drug Designation Workshops, Co-sponsored by: Food and Drug Administration and Drug Information Association European Branch Office.
21. Drug Information Association/FDA Orphan Drug Designation Workshops, Co-sponsored by: Food and Drug Administration and Drug Information Association.
22. Drug Information Association/FDA Quantitative Structure-activity Relationship (Q)SAR Approaches to Assessing Genotoxic Impurities in Pharmaceuticals Workshop, Co-sponsored by: Center for Drug Evaluation and Research and Drug Information Association.
23. Drug Information Association/FDA Tailored Therapeutics: Practical Issues and Methodologies in Selecting the Right Patients, Co-sponsored by: Center for Drug Evaluation and Research and Drug Information Association.
24. Drug Information Association/FDA Training Course on Biostatistical Analytical Methods and Tools for Pharmacovigilance Program of Drug Information Association, Co-sponsored by: Food and Drug Administration and Drug Information Association.
25. Diabetes Drug Development: Challenges and Opportunities Thinktank Meeting, Co-sponsored by: Food Drug Administration and Duke University.
26. Do A Designation Workshop, Co-sponsored by: Food and Drug Administration and Keck Graduate Institute of Applied Life Sciences.
27. Do A Designation Workshop, Co-sponsored by: Food and Drug Administration, National Organization for Rare Disorders and Genetic Alliance.
28. Drug-Induced Liver Injury: Are We Ready to Look? Workshop, Co-sponsored by: Center for Drug Evaluation and Research, American Association for the Study of Liver Diseases and Pharmaceutical Research and Manufacturers Association.
29. Facilitating Oral Product Development and Reducing Regulatory Burden through Novel Approaches to Assess Bioavailability/Bioequivalence Workshop, Co-sponsored by: Center for Drug Evaluation and Research, American Association of Pharmaceutical Scientists and European Federation for Pharmaceutical Sciences.
30. FDA Coalition Against Major Diseases 2011 Coordinating Committee Meeting, Co-sponsored by: Center for Drug Evaluation and Research and Critical Path Institute.
31. FDA New Frontiers in Science Lectureship Program, Co-sponsored by: Food and Drug Administration and Health Research Alliance, Inc.
32. FDA / Society of Clinical Research Associates Clinical Trials Seminar, Co-sponsored by: Office of Regulatory Affairs Denver District and Society of Clinical Research Associates, Inc.
33. FDA / Society of Clinical Research Associates Clinical Trials Seminar, Co-sponsored by: Office of Regulatory Affairs Pacific Region and Society of Clinical Research Associates, Inc.
34. FDA / Society of Clinical Research Associates Clinical Trials Seminar, Co-sponsored by: Office of Regulatory Affairs Philadelphia District and Society of Clinical Research Associates, Inc.
35. Feasibility of Mucosal Healing as a Clinically Significant Endpoint in Inflammatory Bowel Disease Clinical Trials, Co-sponsored by: Center for Drug Evaluation and Research and American College of Gastroenterology.

36. Food Defense Workshop, Co-sponsored by: Office of Regulatory Affairs Southwest Region and Robert M. Kerr Food & Agricultural Products Center.
37. Food Labeling Workshop, Co-sponsored by: Office of Regulatory Affairs Southwest Region and Iowa State University.
38. Food Labeling Workshop, Co-sponsored by: Office of Regulatory Affairs Southwest Region and Robert M. Food & Agricultural Products Center.
39. Food Protection Workshop, Co-sponsored by: Office of Regulatory Affairs Southwest Region, Ozark Food Processors Association and University of Arkansas.
40. Global Outsourcing Conference, Co-sponsored by: Office of Regulatory Affairs Cincinnati District and Xavier University.
41. Hemoglobin Standards and Maintaining Adequate Iron Stores in Blood Donors Workshop, Co-sponsored by: Center for Biologics Evaluation and Research, American Association of Blood Banks, America's Blood Centers and Plasma Protein Therapeutics Association.
42. How to Prepare a HUD Designation Application Workshop, Co-sponsored by: Food and Drug Administration, National Organization for Rare Disorders and Genetic Alliance.
43. How to Prepare a HUD Designation Application Workshop, Co-sponsored by: Food and Drug Administration and Regents of the University of Minnesota
44. Improved Development and Regulation of Transdermal Systems, Co-sponsored by: Center for Drug Evaluation and Research and Drug Information Association.
45. MedCon 2011, Co-sponsored by: Office of Regulatory Affairs Cincinnati District and Xavier University.
46. Over-the-Counter (OTC) Drug Substances and Drug Products Workshop, Co-sponsored by: Center for Drug Evaluation and Research and United States Pharmacopeia.
47. Patient Center Care Conference, Co-sponsored by: Center for Devices and Radiological Health and ECRI Institute.
48. Parenteral Drug Association/FDA Glass Quality Conference, Co-sponsored by: Center for Drug Evaluation and Research and Parenteral Drug Association.
49. Parenteral Drug Association/FDA Joint Regulatory Conference, Co-sponsored by: Center for Biologics Evaluation and Research, Center for Drug Evaluation and Research, Center for Veterinary Medicine, Center for Devices and Radiological Health, Office of Regulatory Affairs and Parenteral Drug Association.
50. Parenteral Drug Association/FDA Pharmaceutical Supply Chain Conference, Co-sponsored by: Center for Drug Evaluation and Research and Parenteral Drug Association.
51. Parenteral Drug Association/FDA Joint Regulatory Conference & Parenteral Drug Association -TRI, Co-sponsored by: Center for Biologics Evaluation and Research, Center for Drug Evaluation and Research, Center for Veterinary Medicine, Center for Devices and Radiological Health, Office of Regulatory Affairs and Parenteral Drug Association.
52. Pharmaceutical Quality System (ICH Q10) Conference: A Practical Approach to Effective Lifecycle Implementation of Systems and Processes for Pharmaceutical Manufacturing, Co-sponsored by: Center for Drug Evaluation and Research and Parenteral Drug Association.
53. Quarantine Release Errors in Blood Establishments, Co-sponsored by: Center for Biologics Evaluation and Research, American Association of Blood Banks and America's Blood Centers.
54. Risk Mitigation Strategies to Address Potential Procoagulant Activity in Immune Globulin Products, Co-sponsored by: Center for Biologics Evaluation and Research and Plasma Protein Therapeutics Association.
55. Safety and Efficacy of Insomnia Drugs, Co-sponsored by: Center for Drug Evaluation and Research and Pharmaceutical Education and Research Institute.

56. Sample Sizes for Decision Making in New Manufacturing Paradigms, Co-sponsored by: Center for Drug Evaluation and Research and Product Quality Research Institute.
57. Science of Abuse Liability, Co-sponsored by: Center for Drug Evaluation and Research, National Institute on Drug Abuse and College on Problems of Drug Dependence.
58. Seizure Detection, Cognitive Function, and TBI/Concussion Devices: Issues in Their Evaluation Workshop, Co-sponsored by: Center for Devices and Radiological Health, American Academy of Neurology, American Epilepsy Society and National Academy of Neuropsychology.
59. Shifting the Balance of Potency and Bleeding Risk for Anti-Coagulant and Anti-Platelet Agents Through Radial Arteriotomy Thinktank Meeting II, Co-sponsored by: Food and Drug Administration and Duke University.
60. Statistics and Medical Devices, Co-sponsored by: Center for Devices and Radiological Health and Advanced Medical Technology Association's Medical Technology Learning Institute.
61. The Future of Medical Products Regulation - Ensuring Safety and Integrity in a Global Market Seminar, Co-sponsored by: Office of Regulatory Affairs and Association of Food and Drug Officials.
62. Women's Health Meeting on Clinical Trial Patient Recruitment and Retention, Co-sponsorship by: Food and Drug Administration and Society for Women's Health Research.