2012 Cosponsored Events

The following is a list of FDA cosponsored events that took place in 2012—including workshops, symposia, and training courses.

1. 2nd International Transporter Consortium Transporter Workshop, Co-sponsored by: Center for Drug Evaluation and Research and American Society for Clinical Pharmacology and Therapeutics.
2. 2nd Latin American Congress of Pharmacogenomics and Personalized Medicine, Co-sponsored by: Food and Drug Administration; Brazilian Ministry of Health, National Cancer Institute, Department of Science Technology; Brazilian Ministry of Education; International Union of Pharmacology; Brazilian Pharmacogenomics Network; Ibero American Network of Phamacogenetics; The Golden Helix Symposia and Pharmacogentics for Every National Initiative.
3. 2nd Scientific Workshop: Analgesic Clinical Trial Translations, Innovations, Opportunities, and Networks Initiative, Co-sponsored by: Center for Drug Evaluation and Research and University of Rochester.
4. 3rd Annual Patient - Reported Outcome (PRO) Consortium Workshop, Co-sponsored by: Center for Drug Evaluation and Research and Critical Path Institute.
5. 3rd Annual Predictive Safety Testing Consortium Workshop, Co-sponsored by: Center for Drug Evaluation and Research and Critical Path Institute.
6. 15th 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.
7. 16th Symposium on the Interface of Regulatory and Analytical Sciences for Biotechnology Health Products - "WCBP 2012," Co-sponsored by: Center for Drug Evaluation and Research and California Separation Science Society.
8. 22nd Annual Association for the Advancement of Medical Instrumentation/ FDA International Conference on Medical Device Standards and Regulation, Co-sponsored by: Center for Devices and Radiological Health and Association for the Advancement of Medical Instrumentation.
9. 2012 Association for the Advancement of Medical Instrumentation /FDA Medical Device Interoperability Summit, Co-sponsored by: Center for Devices and Radiological Health and Association for the Advancement of Medical Instrumentation.
10. 2012 Great Lakes cGMP & Regulatory Science Forum, Co-sponsored by: Office of Regulatory Affairs and University of Illinois Chicago.
11. Accelerating Anticancer Agent Development and Validation Workshop, Co-sponsored by: Center for Drug Evaluation and Research and Duke University Medical Center/Duke University.
12. American Association of Pharmaceutical Scientists/FDA Fifth Bioanalytical Workshop: Quantitative Bioanalytical Methods Validation and Implementation: The 2012 Revised FDA Guidance, Co-sponsored by: Center for Drug Evaluation and Research and American Association of Pharmaceutical Scientists.
13. American Society for Testing and Materials-FDA Workshop on Absorbable Medical Devices: Lessons Learned from Correlations of Bench Testing and Clinical Performance, Co-sponsored by:

Center for Devices and Radiological Health and American Society for Testing and Materials International.

14. Application of IVIVC in Formulation Development, Co-sponsored by: Center for Drug Evaluation and Research and Product Quality Research Institute.
15. Cardiac Safety Workshops, Co-sponsored by: Center for Drug Evaluation and Research, and Duke University.
16. Clinical Investigator Training Course, Co-sponsored by: Center for Drug Evaluation and Research and Duke University.
17. 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.
18. DCD 30-Day Notices and Annual Reports Workshop, Co-sponsored by: Center for Devices and Radiological Health and Advanced Medical Technology Association.
19. Drug Information Association / FDA-Industry PDUFA V Conference, Co-sponsored by: Center for Drug Evaluation and Research and Drug Information Association.
20. Drug Information Association / FDA Revitalizing R&D Productivity in Drug Development, Co-sponsored by: Center for Drug Evaluation and Research and Drug Information Association.
21. FDA/American Glaucoma Society Workshop on the Validity, Reliability, and Usability of Glaucoma Imaging Devices, Co-sponsored by: Center for Devices and Radiological Health and American Glaucoma Society.
22. FDA Clinical Trial Requirements Workshop, Co-sponsored by: Office of Regulatory Affairs Baltimore District and Society of Clinical Research Associates, Inc.
23. FDA Clinical Trial Requirements Workshop, Co-sponsored by: Office of Regulatory Affairs Detroit District and Society of Clinical Research Associates, Inc.
24. FDA Clinical Trial Requirements Workshop, Co-sponsored by: Office of Regulatory Affairs Los Angeles District and Society of Clinical Research Associates, Inc.
25. FDA Coalition Against Major Disease 2012 Annual Meeting, Co-sponsored by: Center for Drug Evaluation and Research, and Critical Path Institute.
26. FDA/ Drug Information Association 2012 Biosimilars Conference: Guidances, Science, and BsUFA, Co-sponsored by: Center for Drug Evaluation and Research and Drug Information Association.
27. FDA / PhUSE Computational Science Symposia, Co-sponsored by: Center for Drug Evaluation and Research, Center for Biologics Evaluation and Research, and Pharmaceutical Users Software Exchange.
28. Food Allergen Workshop, Co-sponsored by: Office of Regulatory Affairs Southwest Region and Robert M. Food & Agricultural Products Center.
29. Food Defense Workshop, Co-sponsored by: Office of Regulatory Affairs Southwest Region and Robert M. Food & Agricultural Products Center.
30. Food Labeling Workshop, Co-sponsored by: Office of Regulatory Affairs Southwest Region and Robert M. Food & Agricultural Products Center.
31. GFR Decline as an Endpoint for Clinical Trials in Chronic Kidney Disease Workshop, Co-sponsored by: Food and Drug Administration and National Kidney Foundation.
32. Global Quality Systems - An Integrated Approach to Improving Medical Product Safety, Co-sponsored by: Office of Regulatory Affairs Northeast Regional Office, Association of Food and Drug Officials and Regulatory Affairs Professionals Society.
33. Global Outsourcing Conference, Co-sponsored by: Office of Regulatory Affairs Cincinnati District and Xavier University.

34. Innovations and Applications of Monitoring Perfusion, Oxygenation and Ventilation, Co-sponsored by: Center for Devices and Radiological Health, Yale University, Society for Technology in Anesthesia and University of Lübeck.
35. IVD Roundtable Submissions Workshop, Co-sponsored by: Center for Devices and Radiological Health and the Association of Medical Diagnostics Manufacturers.
36. MedCon 2012, Co-sponsored by: Office of Regulatory Affairs Cincinnati District and Xavier University.
37. Minimum Clinically Important Difference: An Outcome Metric in Orthopaedic Device Science and Regulation Workshop, Co-sponsored by: Center for Devices and Radiological Health and Georgia Institute of Technology.
38. Oligonucleotides Based Therapeutics Conference 2012: Leveraging Regulators and Industry Knowledge for Future Advancements, Co-sponsored by: Center for Drug Evaluation and Research, Drug Information Association, Oligonucleotide Therapeutic Society, Inc., Health Canada and American Association of Pharmaceutical Scientists.
39. Over-the-Counter Seminars, Co-sponsored by: Center for Drug Evaluation and Research and Consumer Healthcare Products Association.
40. Parenteral Drug Association/FDA Glass Quality Conference, Co-sponsored by: Center for Drug Evaluation and Research and Parenteral Drug Association.
41. 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.
42. Parenteral Drug Association / FDA Pharmaceutical Supply Chain Conference, Co-sponsored by: Center for Drug Evaluation and Research and Parenteral Drug Association.
43. QT Assessment in Early Clinical Development Workshop - Can the Predictive Value be Enhanced to be Similar to that of a TQT Study?, Co-sponsored by: Center for Drug Evaluation and Research and Duke University.
44. Quality System Educational Forum on Medical Device Reporting, Complaints, and Recalls Public Workshop, Co-sponsored by: Office of Regulatory Affairs Dallas District and FDA Medical Device Coalition.
45. Redefining the "C" in CGMP: Creating, Implementing and Sustaining a Culture of Compliance, Co-sponsored by: Center for Drug Evaluation and Research and International Society for Pharmaceutical Engineering.
46. Reproductive and Developmental Toxicology: From In Vivo to In Vitro, Co-sponsored by: Food and Drug Administration, Center for Alternatives to Animal Testing and Middle Atlantic Reproduction and Teratology Association.
47. Statistics and Medical Devices, Co-sponsored by: Center for Devices and Radiological Health and Advanced Medical Technology Association's Medical Technology Learning Institute.