2012 Cosponsored Events

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

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 Pharmacogenetics; 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.
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.
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.
11. Accelerating Anticancer Agent Development and Validation Workshop, Co-sponsored by: Center for Drug Evaluation and Research and Duke University Medical Center/Duke University.

15. Cardiac Safety Workshops, 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.


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.


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.


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.


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.


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.


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.