2013 Co-Sponsored Events

The following is a list of FDA cosponsored events that took place in 2013—including workshops, symposia, and training courses.

1. 4th Annual Predictive Safety Testing Consortium Workshop, Co-sponsored by: Center for Drug Evaluation and Research and Critical Path Institute.
2. 7th Annual FDA/Drug Information Association Statistics Forum, Co-sponsored by: Center for Drug Evaluation and Research and Drug Information Association.
3. Benefit-Risk Workshop: Bridging Qualitative and Quantitative Assessments, Co-sponsored by: Center for Drug Evaluation and Research and Society for Clinical Trials.
4. Changing Regulatory and Reimbursement Paradigms for Medical Devices Workshop, Co-sponsored by: Center for Devices and Radiological Health and American Gastroenterological Association.
5. Clinical Trial Design and Endpoints for Advancement of Development of Intravenous Fat Emulsion Products Workshop, Co-sponsored by: Food and Drug Administration and American Society for Parenteral and Enteral Nutrition.
6. Color in Medical Imaging Summit: An International Workshop on the Technical Framework for Consistency and Interoperability Approaches for Dealing with Color in Medical Images, Co-sponsored by: Center for Devices and Radiological Health and The International Color Consortium.
7. Current Practices and the Future of In vitro Testing Technologies and Regulation, Co-sponsored by: American Association of Pharmaceutical Scientists, Center for Drug Evaluation and Research, International Pharmaceutical Aerosol Consortium on Regulation and Science and United States Pharmacopeias Convention.
8. Detecting and Evaluating Drug-Induced Liver Injury: What's Normal? What's Not? What Should We Do About It?, Co-sponsored by: Center for Drug Evaluation and Research, Critical Path Institute and Pharmaceutical Research and Manufacturers Association.
9. Developing Microphysiological Systems for Use as Regulatory Tools – Challenges and Opportunities Workshop, Co-sponsored by: Food and Drug Administration (FDA), Environmental Protection Agency (EPA), National Center for Advancing Translational Sciences (NCATS), National Institutes of Health, National Institute of Environmental

Health Sciences (NIEHS), National Institutes of Health, International Consortium for Innovation and Quality in Pharmaceutical Development (IQ), Johns Hopkins University, Center for Alternatives to Animal Testing (CAAT), Johns Hopkins University Bloomberg School of Public Health.

10. FDA and The American Association of Pharmaceutical Scientists Conference Bringing Together Small and Large Molecules, Co-sponsored by: Center for Drug Evaluation and Research, and American Association of Pharmaceutical Scientists (AAPS).
11. FDA Clinical Trial Requirements Workshop, Co-sponsored by: Office of Regulatory Affairs (Atlanta, GA) and Society of Clinical Research Associates.
12. FDA Clinical Trial Requirements Workshop, Co-sponsored by: Office of Regulatory Affairs (Dallas, Texas) and Society of Clinical Research Associates.
13. FDA Clinical Trial Requirements Workshop, Co-sponsored by: Office of Regulatory Affairs (Seattle, WA) and Society of Clinical Research Associates.
14. FDA New Frontiers in Science Lectureship Program, Co-sponsored by: Food and Drug Administration and Health Research Alliance, Inc.
15. FDA/Pharmaceutical Users Software Exchange Computational Science Symposium, Co-sponsored by: Center for Drug Evaluation and Research, Center for Biologics Evaluation and Research and Pharmaceutical Users Software Exchange.
16. FDA – University of Maryland Center of Excellence in Regulatory Science and Innovation Lecture Series, Co-sponsored by: Food and Drug Administration and University of Maryland Center of Excellence in Regulatory Science and Innovation.
17. First Multiple Sclerosis Outcome Assessments Consortium / Food and Drug Administration Annual Workshop, Co-sponsored by: Center for Drug Evaluation and Research and Critical Path Institute.
18. Food Allergen Workshop, Co-sponsored by: Office of Regulatory Affairs and The Robert M. Kerr Food and Agricultural Product Center.
19. Food Labeling Workshop, Co-sponsored by: Office of Regulatory Affairs and The Robert M. Kerr Food and Agricultural Product Center.
20. Fourth Annual Patient-Reported Outcome (PRO) Consortium Workshop, Co-sponsored by: Center for Drug Evaluation and Research and Critical Path Institute.
21. Gastroenterology Regulatory Endpoints and the Advancement of Therapeutics for Inflammatory Bowel Disease Workshop, Co-sponsored by: Food and Drug Administration, American College of Gastroenterology, American Gastroenterological Association, Crohn's and Colitis Foundation of America, Inc., North American Society for Pediatric Gastroenterology, Hepatology, and Nutrition, and Pediatric Inflammatory Bowel Disease Foundation.
22. Global Quality Systems – An Integrated Approach to Improving Medical Product Safety, Co-sponsored by: Office of Regulatory Affairs and Association of Food and Drug Officials.

23. Great Lakes cGMP & Regulatory Science Forum, Co-sponsored by: Office of Regulatory Affairs and The University of Illinois Chicago (UIC).
24. In Vitro Diagnostic Roundtable Submissions Workshop, Co-sponsored by: The Association of Medical Diagnostics Manufacturers and Center for Devices and Radiological Health.
25. Medical Devices 101, Co-sponsored by: Office of Regulatory Affairs (Oklahoma) and Food and Drug Administration Medical Device Industry Coalition, Inc.
26. Medical Devices 101, Co-sponsored by: FDA Medical Device Industry Coalition, Inc. and the Office of Regulatory Affairs (Texas).
27. Medical Device Conference, Co-sponsored by: Office of Regulatory Affairs and University of Georgia Division of Nontraditional Education & Outreach College of Pharmacy.
28. Minimal Residual Disease (MRD) as a Surrogate Endpoint in Acute Myeloid Leukemia (AML) Workshop, Co-sponsored by: Center for Drug Evaluation and Research and American Society of Clinical Oncology.
29. Minimal Residual Disease (MRD) as a Surrogate Endpoint in Chronic Lymphocytic Leukemia (CLL) Workshop, Co-sponsored by: Center for Drug Evaluation and Research and American Society of Clinical Oncology.
30. Modeling and Simulation Workshop, Co-sponsored by: Center for Drug Evaluation and Research, Critical Path Institute, and International Society of Pharmacometrics.
31. Oligonucleotide-based Therapeutics Conference 2013, Co-sponsored by: Center for Drug Evaluation and Research, Drug Information Association, Oligonucleotide Therapeutic Society, Inc., and American Association of Pharmaceutical Scientists.
32. Parenteral Drug Association / Food and Drug Administration Advanced Virus Detection Technologies for Evaluation of Biologicals Conference, Co-sponsored by: Center for Biologics Evaluation and Research and Parenteral Drug Association.
33. PharmaLink Conference, Co-sponsored by: Office of Regulatory Affairs and Xavier University.
34. Proactive Compliance and Process Validation Conference, Co-sponsored by: Center for Drug Evaluation and Research and International Society for Pharmaceutical Engineering.
35. Public Education campaign on Decorative Contact Lenses, Co-sponsored by: Food and Drug Administration and American Optometric Association and the Entertainment Industries Council.
36. Public Workshop: Innovations in Breast Cancer Drug Development-Neoadjuvant Breast Cancer Workshop, Co-sponsored by: Center for Drug Evaluation and Research and American Society of Clinical Oncology.

37. Summit on Healthcare Technology in Non-Clinical Settings, Co-sponsored by: Center for Devices and Radiological Health and Association for the Advancement of Medical Instrumentation.
38. Trial Designs and Endpoints for Liver Disease Secondary to Nonalcoholic fatty liver disease (NAFLD) Workshop, Co-sponsored by: Food and Drug Administration and American Association for the Study of Liver Diseases (AASLD).
39. Workshop on Developing Novel Endpoints for Premium Intraocular Lenses, Co-sponsored by: Center for Devices and Radiological Health.