A Primer on FDA Resources for Clinical Investigators

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Clinical Trials and Human Subject Protection

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See the Office of Good Clinical Practice’s (OGCP’s) mission statement on the OGCP’s Web page.

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Workshops and Meetings

In The News

- FDA and OHRP Announcement to extend the comment period on the draft guidance document entitled, “Minutes of Institutional Review Board (IRB) Meetings: Guidance for Institutions and IRBs.”
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Contact FDA

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10903 New Hampshire Ave.,
WO32-5103
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Workshops and Meetings
Preambles contain useful background information including public comments and FDA’s analysis pertaining to the final rule.
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Guidance Documents (Including Information Sheets) and Notices

Guidance documents accessible from this page represent the Agency's current thinking on good clinical practice (GCP) and the conduct of clinical trials. As with all guidance documents, they do not create or confer any rights for or on any person and do not operate to bind FDA or the public. An alternative approach may be used if such approach satisfies the requirements of the applicable statute and regulations. However, in many places throughout these documents, specific regulations are cited and the requirements of the regulations are reiterated. The regulations are enforceable.

Notices accessible from this page are those that have been published by the Agency that contain important information about good clinical practices and the conduct of clinical trials.

Information Sheets
Final Guidances
ICH Guidance Documents
Most Frequently Referenced Guidances for CIs

• Investigator Responsibilities

• FAQ Statement of Investigator (Form FDA-1572)

• Financial Disclosure by Clinical Investigators

• ICH E6 – Good Clinical Practice Consolidated Guidance
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OGCP’s Listserv for notifications of new guidance documents, regulations, etc. as well as FDA webinars
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**Workshops and Meetings**

- Conferences on FDA clinical trial requirements
Clinical Trials and Human Subject Protection

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Useful References

- **Belmont Report**
  Based on the work of the National Commission for the Protection of Human Subjects of Biomedical and Behavioral Research (1974-1978), the Department of Health and Human Services (HHS) revised and expanded its regulations for the protection of human subjects. 45 CFR part 46, in the late 1970s and early 1980s. In 1978, the Commission’s report "Ethical Principles and Guidelines for the Protection of Human Subjects of Research" was published. It was named the Belmont Report, for the Belmont Conference Center, where the National Commission met when first drafting the report. (Extracted from information posted on the DHHS OHRP web site on the Belmont Report; see [http://www.hhs.gov/ochrp/archive/belmontArchive.html#histReport](http://www.hhs.gov/ochrp/archive/belmontArchive.html#histReport))

- **Comparison of FDA and HHS Regulations**
  A chart comparing FDA's regulations for human subject protection with those of the Department of Health and Human Services.

- **Determination of Mode of Action in Combination Products (PDF - 13KB)**
  This rule defines "mode of action" and "primary mode of action" and sets forth the algorithm FDA will use to assign combination products to an agency component for regulatory oversight.

- **E-Mail Messages**
  Copies of e-mail messages (including the original inquiry and associated reply(ies)) that have been submitted by the public to the Good Clinical Practice Program's [gcp.questions@fda.hhs.gov](mailto:gcp.questions@fda.hhs.gov) e-mail account. These e-mail messages have been redacted to the extent permitted by the Freedom of Information Act.

- **"FDA Issues Advice to Make Earliest Stages Of Clinical Drug Development More Efficient"**
  FDA Press Release (Jan. 12, 2006)

- **Improving Health Through Human Drugs**
  This FDA publication, originally titled From Test Tube to Patient: Improving Health Through Human Drugs, tells the story of new drug development in the United States. Articles discuss various aspects of drug development—from test tube to medicine cabinet. This is an excellent primer for learning about the drug development and approval process.

- **FDA and Clinical Drug Trials: A Short History**

- **GCP Training Information**

- **"Innovation or Stagnation? Challenge and Opportunity on the Critical Path to New Medical Products"**
  FDA issued this major report identifying both the problems and potential solutions to the daunting task of ensuring that the unprecedented breakthroughs in medical science are demonstrated to be safe and effective for patients as quickly and inexpensively as possible. Titled "Innovation or Stagnation? Challenge and Opportunity on the Critical Path to New Medical Products," the report carefully examines the "Critical Path" of medical product development—the crucial steps that determine whether and how quickly a medical discovery becomes a reliable medical treatment for patients.

- **International Compilation of Human Subject Protections (PDF - 877KB)**
  OHRP/HCFA maintains and updates the International Compilation of Human Subject Protections. The
Does FDA Conduct GCP Training?

Yes, the Food and Drug Administration (FDA) conducts GCP training. As described below, the agency conducts some GCP training on site, but also partners with other federal agencies and organizations across the United States to conduct additional training. FDA also has recently made GCP training available online.

In the fall of 2009, FDA's Critical Path Initiative launched a Clinical Investigator Training Course targeted at medical professionals who participate in FDA-regulated clinical trials. This 3-day course includes lectures given by senior FDA experts and guest lecturers from industry and academia. It provides FDA's perspectives on new safety concerns, adverse event monitoring, compliance with legal and ethical obligations of clinical research, and acceptable scientific and analytic standards in clinical study design and conduct. See FDA's Clinical Investigator Training Course for further information.

Throughout the year, FDA district offices co-sponsor two-day workshops with the Society of Clinical Research Associates (SoCRA). These conferences are entitled, "FDA Clinical Trial Requirements, Regulations, Compliance, and GCP Conference." FDA personnel from both the co-sponsoring district office and headquarters participate in these workshops. Locations and dates for future workshops are available at the SoCRA website. FDA personnel, from both headquarters and the district offices, also regularly present at meetings of various professional organizations. In addition to SoCRA, these include the Drug Information Association (DIA), Public Responsibility in Medicine and Research (PRIM&R), the Association of Clinical Research Professionals (ACRP), the Regulatory Affairs Professionals Society (RAPS), and others.

FDA routinely collaborates with the Office for Human Research Protections and the Department of Veterans Affairs on regional programs focused on human subject protection in regulated research. Information about these programs is available at:

- The Educational Materials/Workshops page on FDA's Good Clinical Practice Web site
- The Conferences page on the Office for Human Research Protections Web site

Both the Center for Drug Evaluation (CDER) and the Center for Devices and Radiological Health (CDRH) provide online training relevant to clinical trials on their respective sites. These training programs are accessible at CDERLearn and CDRH Learn respectively.

Other agencies in the Department of Health and Human Services do provide training in this area:

- Online training on human subject protection is provided by the Office for Human Research Protections.
- Clinical Research Training is a course developed by the National Institutes of Health to train its own investigators. It may be accessed by others to enhance their knowledge of clinical research.

Finally, there are numerous references related to good clinical practice (GCP) and human subject protection (HSP), available on FDA's website, including:

- FDA's GCP and HSP regulations found in Title 21 of the Code of Federal Regulations, Parts 50, 56, 312, and 812
- The preambles related to these regulations
- ICH E6 Good Clinical Practice Consolidated Guidance [261KB PDF]
- FDA Information Sheets for IRBs and Clinical Investigators
- Compliance Program Guidance Manuals
- FDA guidance for industry: Finalized documents and draft guidance documents (and proposed regulations).
Compliance/Enforcement

Implementation of CT.gov Databank

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We’re here to help...

Really!!!!!!!!

Thank you!

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