

HMS/HSDM Committee on Human Studies  
**Electronic Training Book**\*

1. Ethical Guidelines
  - Nuremburg Code
  - Declaration of Helsinki
  - Belmont Report
  - CIOMS – International Ethical Guidelines for Research involving Human Subjects
  
2. IRB Regulations
  - DHHS 45 CFR 46
  - FDA 21 CFR 50
  - FDA 21 CFR 56
  - International Compilation of Human Subject Research Protections
  
3. OHRP Guidance
  - Engagement of Institutions in Research
  - Human Subject Regulations Decision Charts
  - IRB Review of (grant) Applications for HHS Support
  - Knowledge of the Local Research Context
  - Informed Consent Checklist
  - Exemption Categories
  - Use of Expedited Review Procedures
  - Expedited Review Categories
  - Research Involving Coded Private Information or Biological Specimens
  - Research Use of Stored Data or Tissues
  - Research Involving Embryonic Stem Cells, Germ Cells and Stem Cell-Derived Test Articles
  - Local IRB Review of Multicenter Clinical Trials
  - Review of Multicenter Clinical Trials Sponsored by DAIDS and NIAID
  - Special Protections for Children as Research Subjects
  - Involvement of Prisoners in Research
  - Continuing Review
  - Reviewing and Reporting Unanticipated Problems and Adverse Events
  
4. Terms of the Federalwide Assurance for the Protection of Human Subjects

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\* *All items listed are hyperlinks to information available on the internet.*