Investigator Resources

Regulations:

- Common Rule (45 CFR 46)
- eCFR: 21 CFR Part 50-Protection of Human Subjects
- eCFR: 21 CFR Part 56-Institutional Review Boards

National Institutes of Health:

- NIH Home Page
- Office of Recombinant DNA Activities (ORDA)
- Office of Grants and Contracts
- National Human Genome Research Institute
  - Ethical, Legal and Social Implications (ELSI)

UNMC Links:

- General Counsel's Memo on Mandatory Reporting of Child Abuse and Related Statute of Limitations
- Institutional Biosafety Committee (IBC)
- Animal Care and Use Program (IACUC)
- Sponsored Programs Administration (SPA)

Food and Drug Administration:

- FDA Web Site
- FDA - Center for Drug Evaluation and Research (CDER)
- FDA - Center for Devices and Radiological Health

International Standards:

- International Compilation of Human Research Standards

Other Federal Agencies:

- National Archive and Records Administration
  - Federal Register Online
- Office for Civil Rights
  - Medical Privacy - National Standards to Protect the Privacy of Personal Health Information
- Department of Health and Human Services
  - Child Abuse Reporting Memo

Organizations and Other Items of Interest:

- Public Responsibility in Medicine & Research (PRIM&R)
- National Bioethics Advisory Commission (NBAC)
- American Society for Bioethics and Humanities

IRB History and Principles:

- “What Makes Clinical Research Ethical?” Emanuel, et al; JAMA 283(20):2701, 2000
- THE BELMONT REPORT: Ethical Principles and Guidelines for the Protection of Human Subjects of Research(UNMC)
- The Belmont Report(HHS)

Revision #3
Created 12 July 2024 19:24:43 by Robert A Lewis

Updated 8 August 2024 19:17:24 by Robert A Lewis