

# Investigator Resources

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## Regulations:

- [Common Rule \(45 CFR 46\)](#)
  - [eCFR: 21 CFR Part 50-Protection of Human Subjects](#)
  - [eCFR: 21 CFR Part 56-Institutional Review Boards](#)
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## 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 \(ELSI\)](#)
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## 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 \(SPA\)](#)
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## Food and Drug Administration:

- [FDA Web Site](#)
  - [FDA - Center for Drug Evaluation and Research \(CDER\)](#)
  - [FDA - Center for Devices and Radiological Health](#)
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## International Standards:

- [International Compilation of Human Research Standards](#)
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## 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](#)
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## Organizations and Other Items of Interest:

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