CPHS – IRB

- Committee for the Protection of Human Subjects
- Institutional Review Board
- Composition
Overview

- Mission: to protect the rights and welfare of human research participants
- Definition of research: systematic investigation, including research development, testing, and evaluation, designed to develop or contribute to generalizable knowledge
Human Subjects

- An individual about whom an investigator, whether professional or student, conducting research obtains:
  - Data through intervention or interaction with the individual, or
  - Identifiable private information
Types of Review

- **Exempt Research – Short application**
  Processed administratively

- **Expedited Review**
  Reviewed by a committee member

- **Full Committee Review**
  CPHS committees meet four times a month
Exempt Review Examples

- Retrospective chart review without identifiers
- Surveys/interviews/focus groups without sensitive questions
- Evaluation of educational program/course
- Use of already collected biological samples (collected for clinical purposes)
- Secondary analysis of already collected data
Expedited Review Examples

- Retrospective chart review with identifiers
- Surveys/interviews/focus groups with sensitive questions
- Collection of blood samples of limited volume
- Collection of biological samples for research purposes by noninvasive means
Full Board Examples

Clinical trial
- Randomized, double-blind, placebo
- Drug/Device
- When subjects are students, staff or residents
- Vulnerable populations (children, prisoners)
- International research
Submission for Initial Review

Required documents:
- Application
- Informed consent documents
- Protocol/Grant Cover Sheet
- Data Collection/Case Report Forms
- Letters of Support
Areas of IRB Focus

- Autonomy and respect for persons
- Equitable selection of subjects
- Risk versus benefit
- Recruitment methodology
- Consent process
- Privacy and confidentiality
Protocol Components

- Hypothesis/research question
- Background with references
- Subject population
- Recruitment methodology
- Procedures
- Sample size
- Analysis plan
- Security of data
Common Shortfalls

- Readability
- Research vs. Standard of Care
- Sample Size – High/Low
- Explain Tests/Scales/Tools
- Location, Environment
Informed Consent

- Consent is a process
- Document is a guideline
- Contains required elements and language
- Adult/parent/child
- Must be approved by CPHS
- Stamped version
- Signed by subject and research team
After IRB Approval

- Changes/Amendments
- Continuing Review (Annually)
- Protocol Deviations
- Serious Adverse Events
- Unanticipated Problems
- Data Safety Monitoring Reports
Question

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