The University of Iowa
Human Research Protection Program (HRPP)

UI Institutional Review Boards
- Biomedical (IRB-01) & VAMC (IRB-03)
- Behavioral/Social Science (IRB-02)

Preferred IRB Reliance Model:
- SMARTIRB
- Commercial IRBs: WIRB, Advarra
- Consortiums: Negotiated on a case by case basis

UI Hospitals & Clinics:
- Pharmacy & Therapeutics (P&T) Committee
- P&T IDS (Inv. Drug Service)
- Research Billing Compliance
- Nursing Research Committee (NRC)

Holden Comprehensive Cancer Center
- Protocol Review & Monitoring Committee

Research Counsel

Research Integrity Officer

Office of the Vice President for Research
- Vice President for Research

Associate VP for Research Administration/Institutional Official

Human Subjects & Conflict of Interest Office

Environmental Health & Safety Office

- Institutional Biosafety Committee
- Stem Cell Committee
- Medical Radiation Protection Committee

Division of Sponsored Programs

Additional Entities:
- Related policies governed by the UI Operations Manual and UI Health Care
- UI HIPAA Privacy Officer and Security Officer
- UI Accounting Services & Grant Accounting
- UI Information Technology Services
- ICTS Clinical Research Unit

Last revised 1/2021
The University of Iowa IRB interacts with other committees in the Human Research Protection Program (HRPP) and other offices or entities that review research projects involving human subjects research. Many of them review information included in the HawkIRB application and attached documents. Others have a separate application form or process. Use the dark boxes in this chart to determine if you need approval from a particular committee, office or entity. Click “Learn More” to go to their web page. For projects overseen by an external IRB, all approvals must be obtained from these other committees/entities prior to submission to the external IRB. Here is a key to the color coding:

<table>
<thead>
<tr>
<th>Approve application before IRB review</th>
<th>Accounting Services, CIRC, PRMC, RBC</th>
</tr>
</thead>
<tbody>
<tr>
<td>Review application concurrent with IRB review. Finalize before IRB approval.</td>
<td>CT.gov, MRPC, NRC, P&amp;T</td>
</tr>
<tr>
<td>Concurrent with IRB review. Finalize before IRB releases application.</td>
<td>DSP, P&amp;T IDS, IBC</td>
</tr>
<tr>
<td>Review project separately from IRB review and approval process</td>
<td>CRU, Pathology</td>
</tr>
<tr>
<td>Review application after IRB approval</td>
<td>VAHCS</td>
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</tbody>
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- **Accounting Services**
  - Compensation for Subjects, Cash or Cash Equivalent
  - Learn more

- **UIHC Research Billing Compliance (RBC)**
  - Research Involves Billable Services from UI Health Care
  - Learn more

- **Pharmacy & Therapeutics Committee (P&T)**
  - Substance Injected, Ingested, or Applied
  - Use of a Contrast Agent
  - Learn more

- **Nursing Research Committee (NRC)**
  - Use of UI Health Care Nursing Resources
  - Learn more

- **Institutional Biosafety Committee (IBC)**
  - Use of Recombinant DNA (rDNA)
  - Learn more

- **Department of Pathology Services**
  - Using Pathology Services for Research Purposes
  - Learn more

- **Clinical Research Unit (CRU)**
  - Using Clinical Research Unit Resources
  - Learn more

- **Conflict of Interest in Research Committee (CIRC)**
  - PI or Team Member Has Financial Conflict of Interest
  - Learn more

- **Protocol Review & Monitoring Committee (PRMC)**
  - Use of Patients or Resources from Holden Comprehensive Cancer Center (HCCC)
  - Learn more

- **ClinicalTrials.gov (CT.gov)**
  - Meets Definition of Applicable Clinical Trial (ACT)
  - Learn more

- **Medical Radiation Protection Committee (MRPC)**
  - Use of Diagnostic or Therapeutic Radiation
  - Learn more

- **P&T Investigational Drug Service (P&T IDS)**
  - Use of Approved or Investigational Drugs
  - Learn more

- **Division of Sponsored Programs (DSP)**
  - External Funding Source
  - Research Related Agreements
  - Learn more

- **Veterans Administration Health Care System (VAHCS)**
  - Use of Patients or Resources from VAHCS
  - Learn more
HRPP Committee Regulatory Mandates

- **CIRC: Conflict of Interest in Research Committee**
  - Public Health Service (PHS), Institutional Policy (Ch.18), & HRPP Accreditation

- **DSP: Division of Sponsored Programs**
  - Federal Acquisition Regulations, Department of Commerce, Federal Uniform Guidance, NIH, Institutional Policy (II–27, V–5, V.6, etc)

- **RBC: Research Billing Compliance**
  - The Centers for Medicare and Medicaid Services’ Clinical Trial Policy

- **RBC/CMS: Research Billing Compliance for device trials only**
  - The Centers for Medicare and Medicaid Services’ Clinical Trial Policy, Social Security Act (Section 1862(m)), HHS Regulations (42 CFR 405, Subpart B)

- **MRPC: Medical Radiation Protection Committee**
  - IDPH, Iowa Administrative Code, National Institute for Occupational Health & Safety, CDC, & UI\UIHC policies

- **PRMC: Holden Comprehensive Cancer Center Protocol Review and Monitoring Committee**
  - NCI (National Cancer Institute)

- **P&T: UIHC Pharmacy & Therapeutics**
  - Joint Commission, Medicare Pharmaceutical standards, & Iowa law on drug dispersing & storage

- **P&T IDS: Pharmacy & Therapeutics Investigational Drug Service**
  - Joint Commission, Medicare Pharmaceutical standards, & Iowa law on drug dispersing & storage

- **NRC: Nursing Research Committee**
  - UIHC Bylaws & American Nurses Magnet Accreditation

- **IBC: Institutional Biosafety Committee**
  - NIH, USDA, CDC, DHHS

- **VA Related Committees**
  - Veterans Health Administration

- **VA Related Committees**
  - Veterans Health Administration