The University of Iowa Human Research Protection Program (HRPP)

Office of the Vice President **Research Integrity Officer** for Research Vice President for Research **UI Institutional Review Boards** Biomedical (IRB-01) & VAMC (IRB-03) Behavioral/Social Science (IRB-02) **Associate VP for Research Preferred IRB Reliance Model:** Administration/ **SMARTIRB** Research Counsel **Institutional Official** Commercial IRBs WIRB. Advarra **Division of Sponsored UI Hospitals & Clinics: Consortiums Programs** Pharmacy & Negotiated on a case by case basis Therapeutics (P&T) Committee P&T IDS (Inv. Drug Service) Research Billing Compliance **Human Subjects** Conflict of Interest **Environmental Health &** Nursing Research in Research & Conflict of **Safety Office** Committee (NRC) Committee **Interest Office** Holden Comprehensive Cancer Center Protocol Review & Medical Radiation Institutional Stem Cell Monitoring Committee Protection Biosafety Committee Additional Entities Committee Committee

- ---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



Last revised 1/2021

OTHER APPROVAL REQUIREMENTS FOR HUMAN SUBJECTS RESEARCH

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:

Approve application <u>before</u> IRB review	Accounting Services, CIRC,PRMC, RBC
Review application <u>concurrent</u> with IRB review. Finalize <u>before</u> IRB approval.	CT.gov, MRPC, NRC, P&T
Concurrent with IRB review. Finalize before IRB releases application.	DSP, P&T IDS, IBC
Review project separately from IRB review and approval process	CRU, Pathology
Review application <u>after</u> IRB approval	VAHCS

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

<u>Learn more</u>

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
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