Sponsored Research Compliance Regulations and Requirements

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Learning Objectives

- Before You Begin
- · At the point of proposal
- Compliance Committees
- · At the point of award

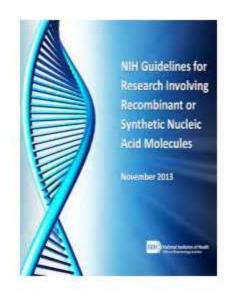


Institutional Biosafety Committee (IBC)

Responsible for the safe use of:

- Recombinant or Synthetic nucleic acids
- Human or NHP blood, cells, tissues, fluids, secretions
- Biological toxins
- Bacteria, virus, fungi, yeast, parasites and prions

In accordance with



Federal
Select Agent
Program

All covered materials must be listed on approved lab IBC Registration prior to use

IBC Areas of Concern

Protecting study participants, research staff, community, and environment

Improve research outcomes

Focus on

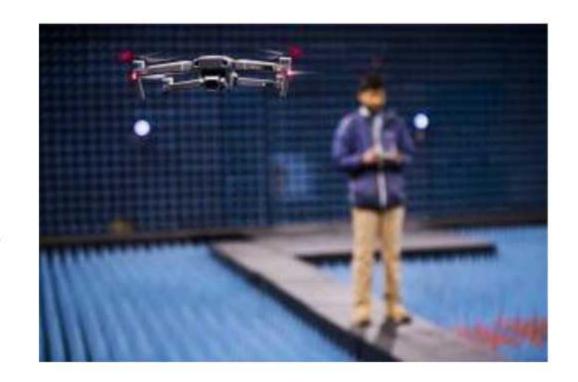
Laboratory Safety & PPE
Biosafety Cabinets, Aerosol Control
Waste Handling & Disinfection



Unmanned Aircraft System (UAS) Review Board

Responsible for:

- To review, approve and monitor the use or proposed use of UAS's, also known as unmanned aerial vehicles and drones
- Adhere to university Policy on Use of Drones
- Further information on federal regulations governing use of drones may be found at the FAA's website



IRB: Is it Human Subjects Research?

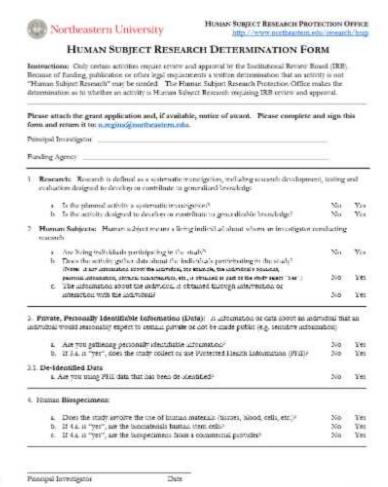
Is it Research?

- Systematic investigation
- Develop or contribute to generalizable knowledge

Does it involve Human Subjects?

- Gather data about living individual participants
- Via intervention or interaction with the individual

Not sure? Use the HSR Determination Form



Andre advance

IRB Regulations

 45 CFR 46: Protection of Human Subjects

 Department of Health & Human Services

Common Rule

Focus On:

- Safety
- Consent
- Privacy





Special Participants (Children, People who are pregnant, incarcerated, or cognitively impaired)

Alterations to Consent (Waiver of Consent, Deception, Documentation of Consent)

Drug or Medical Device Studies

Collaborations or Multi-Site Studies



IRB Timeline

At Proposal

- Determine whether IRB approval is needed
- Complete required CITI training
- Begin developing & submitting IRB protocol

At Award

- IRB Approval should be in place
- Continuing review period indicated on Approval Letter
- For multi-year awards, IRB approval of scope may be acceptable



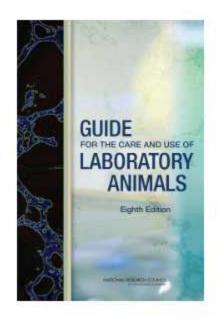
Follow Up with IRB/NU-RES when Human Subjects Research begins

IACUC Regulations

IACUC reviews the use of **live**, **vertebrate animals** in Testing, Research, and Training.

PHS/OLAW

Regulated by PHS Policy and The Guide Covers all live, vertebrate animals

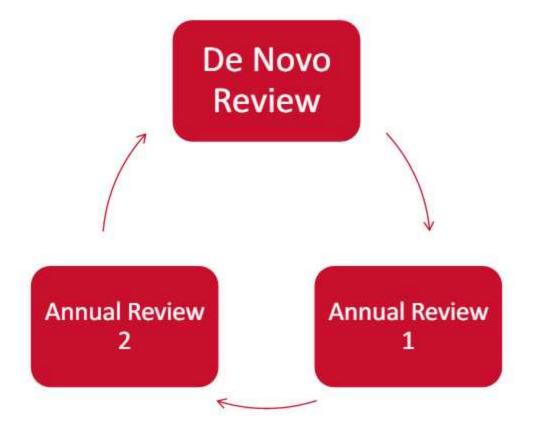


USDA/APHIS

Regulated by the Animal Welfare Act

Covers all warm-blooded animals except for mice, rats, and birds bred for research

IACUC Review Cycle



Submit amendments off-cycle as needed

IACUC Areas of Concern

Use of Modern, Humane Techniques

Justification for animal use procedures & numbers

Management of pain & distress

Use of biohazards, transgenics, and recombinant DNA (in partnership with IBC)

Replace Reduce Refine



IACUC approval must always be secured before work with animals begins.

IACUC Timeline

At Proposal

- Complete required training
- Consult with lab animal vet on procedures and pain/distress management
- Begin developing & submitting IACUC protocol

At Award

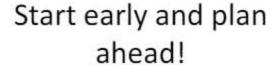
- IACUC Congruency Check
- Submit amendment if necessary



IACUC approval must always be secured before work with animals begins.

Tips from the RRC







If you're unsure about anything, contact Research Compliance



Keep projects and protocols organized

Contact Information

	How to Submit	Who to Contact
IBC	BioRAFT	biosafety@northeastern.edu
IRB	E-Mail Forms on Website	Nan Regina – <u>n.regina@northeastern.edu</u> Kate Skophammer (CPS) – <u>k.Skophammer@northeastern.edu</u>
IACUC	E-Mail Forms on Website	iacuc-office@northeastern.edu
UAS/Drones	E-mail Forms on Website	UAS Review Board at UASRB@northeastern.edu
Research Compliance		researchcompliance@northeastern.edu



Questions?



Thank You!

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