



Research Administration

Human Research Protection Program

Human Research News

May 20, 2024

1) BruinIRB Transition Update

OHRPP is moving forward with a transition to the BruinIRB system for IRB submissions. A full timeline will be presented in June, but we are on track to accept all new submissions in BruinIRB starting Tuesday, September 3, 2024.

To support the research community during this transition, OHRPP has put together a schedule of trainings. All trainings will be recorded and available on OHRPP's YouTube channel shortly after each session. **All presentations are from noon-1pm.**

Date	Presentation	Registration link
Wednesday May 22, 2024	Learn at Lunch: <i>BruinIRB Phase 2 Roll-out Timeline</i>	Link
Thursday June 6, 2024	Learn at Lunch: <i>BruinIRB Protocol Uploads</i>	Link
Tuesday June 25, 2024	Learn at Lunch: <i>Migration of Existing Studies from webIRB to BruinIRB</i>	Link
Tuesday July 9, 2024	Learn at Lunch: <i>CR/AMs and AMs in BruinIRB</i>	Link
Wednesday July 24, 2024	Learn at Lunch: <i>How to Submit New Studies in BruinIRB</i>	Link
Wednesday August 7, 2024	Learn at Lunch: <i>CRAMs and AMs in BruinIRB (live repeat)</i>	Link
Wednesday August 21, 2024	Learn at Lunch: <i>How to Submit New Studies in BruinIRB (live repeat)</i>	Link

2) OHRPP's Office Hours

[Register once](#) to join any session:

Upcoming Office Hours:

- May 23, 2024, at 2pm

➤ June 6, 2024, at 2pm

3) On-demand Training Opportunities & Resources

- [University of Minnesota's Annual Research Ethics Day Conference - Building Partnerships to Advance Ethical Research: Collaborators, Communities & Companies](#)
 - Interdisciplinary experts discuss the elements of successful collaboration that promote research integrity, inclusivity, mutual respect, and accountability in the conduct of research.
- [OHRP's Participant-Centered Informed Consent Training Program](#)
 - This interactive online program consists of 6 modules and delivers a comprehensive training for creating, designing, and reviewing consent forms or templates for the purposes of ***ensuring that prospective research participants understand the content and appreciate how it relates to them.***
- [The MRCT at Brigham and Women's Hospital new clinical research glossary:](#)
 - This plain-language global standard has been developed at the [MRCT](#) in collaboration with [CDISC](#) for research participants ***and anyone interested in developing easy-to-understand research materials.***

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