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Agenda

- New Post-Award Director
- F&A Costs on Sponsored Awards
- IBISResearch Accuracy Human Subjects & Account Setup Delays
- PG Accounts for Closed & Closing Projects in Workday
- Research Administration Curriculum Deadline
- NSF Safe and Inclusive Working Environment for Off-Campus Research
- NIH Data Management Plan
- Form H live
- Clinical Trial Fee Changes
- Effort Certification and Cross Company PAA's
- New Research IT Director
- UM's COI policy, Annual Training and Disclosure Process in the UDisclose System
- NSF Disclosure Requirement Clarification & Compliance
- Designating Investigators for COI Review in the IBISResearch Grants Module
- Upcoming Huron Upgrades and Implementation
- ORCID ID
- AAHRPP Reaccreditation Site Visit

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New Post Award Director

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Laura Kozma

Associate Vice President Research Administration

New Post Award Director



Stephane Malebranche

Director, Post Award

Email: smalebranche@miami.edu

Telephone: 305-284-3223





F&A Costs on Sponsored Awards

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Lionel Vera

Executive Director
Research Administration

F&A Disguised as Interdepartmental Charges





UG Language

Chapter II / Part 200 Previous / Next / 7

(2) Items such as office supplies, postage, local telephone costs, and memberships must normally be treated as indirect (F&A) costs.

Source: https://www.ecfr.gov/current/title-2/subtitle-A/chapter-II/part-200.

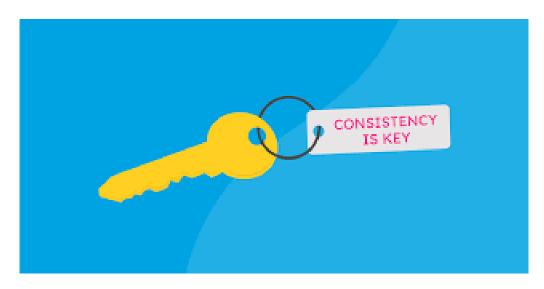




Can we charge the Grant if...

- ☐ we bill the cost as a billing rate (e.g., interdepartmental services)?
- we can allocate the cost to the grant and say, yes, this cost is used exclusively, 100%, for this grant?
- ☐ the cost is in the application budget as a separate line item approved by the sponsor?
- ☐ the cost is all of the above combined?!?

Answers: **No**, **no**, **no**, **and no**, ...not necessarily in that order.







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IBISResearch Accuracy Human Subjects & Account Setup Delays

Laura Kozma

Associate Vice President Research Administration

IBISResearch Accuracy HS & Account Setup

REMINDER!



- Make sure human subject information is consistent and accurate in all areas of IBIS and the application.
- Inconsistent information WILL slow down award setup





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PG Accounts for Closed & Closing Projects in Workday

Laura Kozma

Associate Vice President Research Administration

PG Accounts for Closed & Closing Projects

- A PG account must be provided if there is an anticipated deficit
- Provide on the closeout form or when requested by Research Administration
- Not providing this creates a significant amount of work for Research Administration which impacts service to departments and faculty







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Research Administration Curriculum Deadline

Laura Kozma

Associate Vice President Research Administration

Research Administration at the U

- Deadline extended to August 31, 2023
- Non-compliance will result in loss of roles in IBIS and Workday until the training is complete
- If you have not started, you still have time to complete by the end of the summer!







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NSF Safe and Inclusive Working Environment for Off-Campus Research

Brandon Strickland

Executive Director
Research Administration

NSF Safe Working Environment

NSF Policy (PAPPG 23-1, Chapter II.E.9) requires fostering safe and harassment-free environments wherever science is conducted, including off campus or off site.

Each proposal that proposes to conduct research off campus or off site requires:

- Development of Plan for Safe and Inclusive Working Environment
- Certification of Plan by Institution's Authorized Official (AOR)
- Distribution of Plan to research team prior to departing for off campus or off site research. Pls are responsible for developing plan, maintaining copy of plan, and documentating who received copy of plan. The plan is NOT to be submitted to NSF unless specifically requested by NSF.
- NSF defines "off-campus or off-site research" as "data/information samples being collected off campus or off site, such as fieldwork and research activities on vessels and aircraft".



Developing a Plan

Developing a Plan for Safe and Inclusive Working Environment Required Elements in the Plan:

- 1. Description of how the organization will address the following types of behavior:
 - Abuse of any person, including, but not limited to, harassment, stalking, bullying, or hazing of any kind, whether the behavior is carried out verbally, physically, electronically, or in written form
 - o Conduct that is unwelcome, offensive, indecent, obscene, or disorderly
- 2. Steps the organization will take to nurture an inclusive off-campus or off-site working environment (trainings, processes, codes of conduct, support mechanisms, developmental events, etc)





Developing a Plan

Developing a Plan for Safe and Inclusive Working Environment

Required Elements in the Plan:

- 3. How communications within the team and communications to the organization will be handled, and plans for how any special circumstances will be addressed (such as the involvement of multiple organizations or third parties in the working environment should be taken into account)
- 4. Process or method for handling incident reports How will incident reports be made? How will they be received? How will they be resolved?





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Brandon Strickland

Executive Director
Research Administration

NIH Data Management and Sharing (DMS) Policy - Effective for New and Competing Applications due on/after January 25, 2023.

When is the NIH DMS plan required?

 The data management plan (historically only required for proposals requesting more than \$500K in direct costs annually or projects generating genomic data) is now required in most NIH applications.

DMS Policy Applies*	DMS Policy Does Not Apply
Research Projects	Training (T)
 Some Career Development Awards (Ks) 	 Fellowships (F)
Small Business (SBIR/STTR)	 Construction(CO6)
Research Centers	Resources (Gs)
	 Research Related Infrastructure Programs (SO6)





What is required under this new policy?

- Include Budget Request for Data Management and Sharing costs in NIH grant application
- Submit Data management and Sharing Plan with NIH application
- Adhere to NIH-approved Data Management and Sharing Plan when funded

What resources are available?

Data Plan Resource Document

NIH DMS Plan Template

NIH Data Sharing Summary

Why is there a new policy?

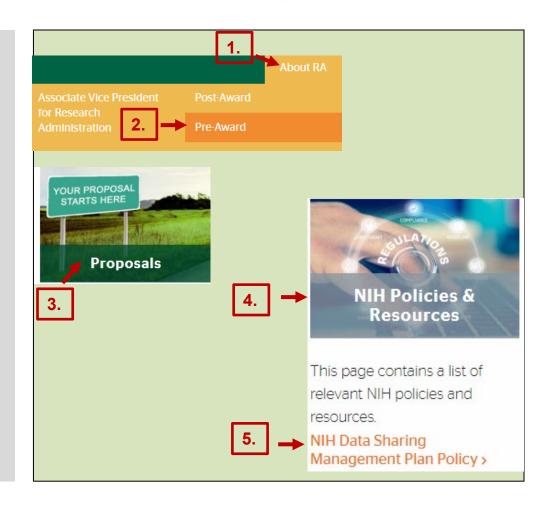
To promote the sharing of scientific data.





For additional information go to the RA website: ora.Miami.edu

- 1. About RA
- 2. Pre-Award
- 3. Proposals
- 4. NIH Policies & Resources
- NIH Data Sharing Management Plan Policy







Form H Live

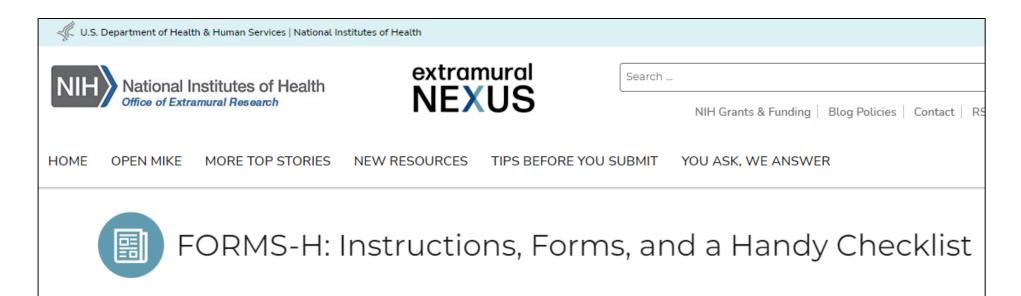
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K. Brandon Strickland, J.D.

Executive Director, Research Administration

Form H live in the system







Clinical Trial Fee Changes

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K. Brandon Strickland, J.D.

Executive Director, Research
Administration

Clinical Trial Fee Changes

Changes to fees on Clinical Trials coming February 15.

- Cris Fee Increase by \$1000
- Amendment \$750
- \$250 Feasibility Fee and F&A to 36%







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Effort Certification & Cross Company PAA's

Kimberly Croft

Executive Director, Research Accounting and Cost Analysis

Effort Certification

Certification Deadline: February 10

Exception requests: Send to kcroft@miami.edu



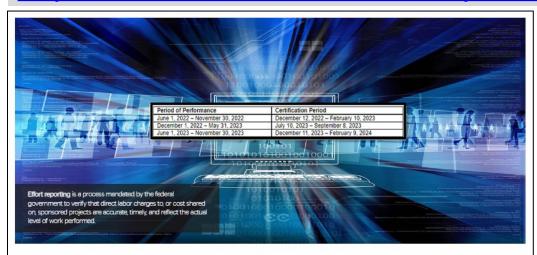




Effort Certification

Effort reporting information:

https://www.ora.miami.edu/compliance/effort-reporting/index.html



Effort reporting is the method of certifying to federal branching agencies that the effort required as a condition of the award has actually been completed. Effort reporting is required by the <u>Uniform Guidance</u>.

The University of Miami Office of Research Administration is deploying a new upgraded version of the Employee Compensation Compliance system known as ECC, (formerly known as ECRT), to serve as the tool to facilitate the effort certification and reporting process. The ECC system is designed to help comply with the provisions of the University of Miami's Effort Reporting-policy, federal policy and other relevant agency policies that govern charging an individual's effort to sponsored projects.

The University of Miami uses Project Confirmation through the Employee Compensation Compliance System (ECC).

Project Confirmation is a streamlined effort certification process, which requires that the Principal Investigator certify his/her effort and the effort of staff on his, or her project if there are other faculty members on the project, each will certify his, or her own effort. This method of certification reduces the number of individuals required to certify effort.





CC System

In the Employee Compensation Compliance (ECC) system Principal Investigators are required to complete Project Confirmation, a process which requires they certify his/her effort and the effort of staff on each project. If there are other faculty members on the project, each will certify their own effort.

ECC: Project Confirmation >



This page contains the University of Miami's Effort Reporting Policy. This policy ensures that effort reporting of externally sponsored activities is managed in a manner consistent with requirements set forth in the Uniform Guidance.

Read More >



Effort reporting is the method of certifying to federal branching agencies that the effort required as a condition of the award has actually been completed. Effort reporting is required by the <u>Uniform Guidance</u>.

Effort Certification Tip Sheet >



Training for the Employee Compensation Compliance (ECC) system is required. CBLs and live offerings are available through

ULearn.

ECC: Project Confirmation for Effort Coordinators >

ECC: Project Confirmation for Principal Investigators >

Read More >



Sponsored Effort, or salary charged to a sponsored project is one of the main components of cost on a sponsored project.

Read More >



This page contains frequently asked questions concerning Effort Reporting. For example, what is effort reporting, what is meant by the term effort, who is subject to effort reporting, etc.

Read More >



Effort Certification Questions

Contact the ECC Administrative Office:

Email: ecrt.admin@miami.edu

Phone: (305) 284-4054



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Effort Reporting Policy

https://www.ora.miami.edu/policies/all-ora-policies/index.html







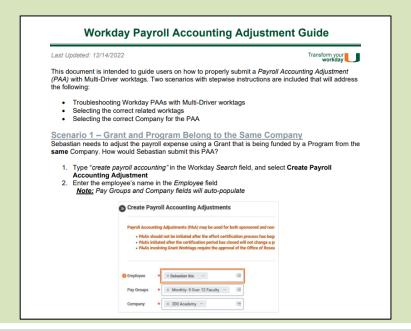
Cross Company PAA's

Step-by-step guide for PAA's with multi-driver worktags:

https://my.it.miami.edu/ assets/pdf/workday-payroll-accounting-adjustment-guide.pdf

Highlights of this guide include:

- Troubleshooting Workday PAAs with Multi-Driver worktags
- Selecting the correct related worktags
- Selecting the correct Company for the PAA



For inquiries contact:

- The EBS team, at workday@miami.edu
- Gloria Gari at ggari@miami.edu





New Research IT Director

UNIVERSITY OF MIAMI



Mikki ONeal

Associate Vice President, Research Operations & Infrastructure (ROI)

Research IT Director



Joshua Rivera
Director, Research Information Technology





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UM's COI policy, Annual Training and Disclosure Process in the UDisclose System

Lory Hayes

Director, Disclosures & Scholarly
Activities Management
(DSAM)

UM's Comprehensive COI Policy

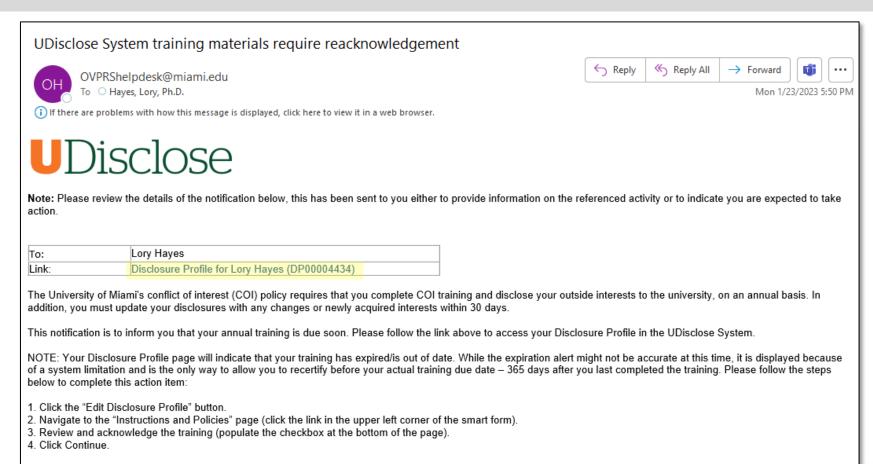
- Conflict of Interest, Conflict of Commitment, Foreign Influence, and Institutional Conflict of Interest Policy
- Promulgated January, 2022
- Policy articulates all <u>Covered Persons</u> must:
 - Complete training on the policy, annually
 - Submit a <u>Disclosure Profile</u>, annually
 - Changes and new interests must be submitted within 30 days





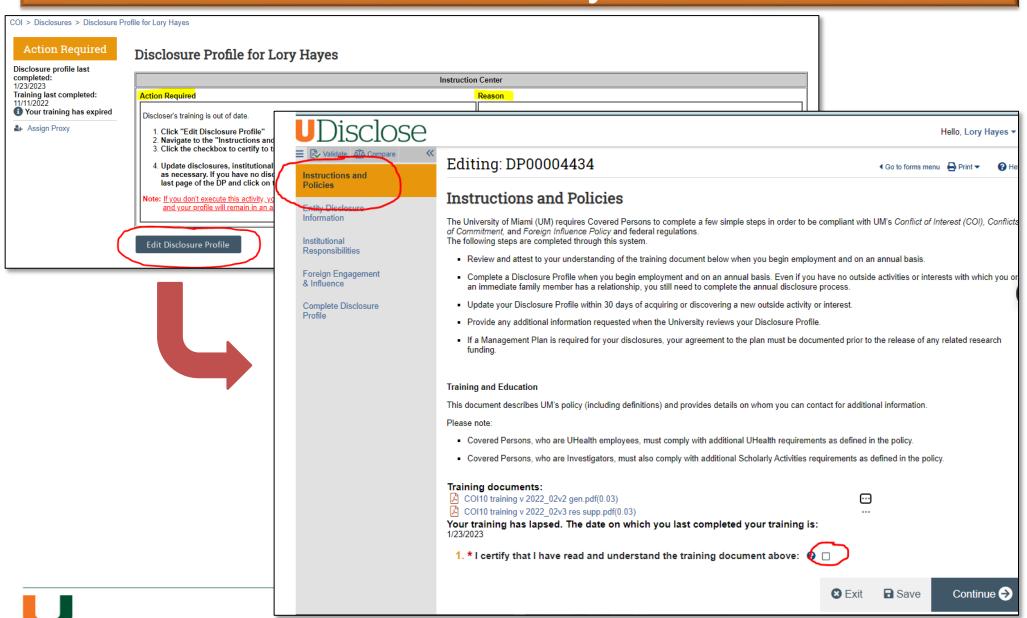
UDisclose System

- Annual training and disclosures are coming due
- Training reminder notifications will be sent ~30 days before expiration





UDisclose System



UDisclose System



Resources (require CaneID):

- UDisclose User Guides & Job Aids (Box folder)
- Discloser Compliance Status Job Aid (PDF)
- Discloser Compliance Status Report (PowerBI)

Questions? Call the UDisclose Help Desk 305-243-0877



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NSF Disclosure Requirement Clarification & Compliance

Lory Hayes

Director, Disclosures & Scholarly
Activities Management
(DSAM)

NSF Disclosure Requirement

- Proposal & Award Policies & Procedures Guide (PAPPG) (NSF 23-1)
- Effective for proposals submitted on or after January 30, 2023
- Must disclose <u>venture or other capital financings</u>
- Qualtrics survey will be completed by all investigators submitting NSF proposals
 - Y/N responses
 - No identifying information will be collected
- RA will have access to a spreadsheet listing names
- If not completed, NSF proposal cannot be submitted





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Designating Investigators for COI Review in the IBISResearch Grants Module

Laura Kozma

Associate Vice President Research Administration

Reminder: Responsible Investigators

- All individuals responsible for the design, conduct or reporting must be identified by clicking "yes" when adding the person in IBIS
- If the FP was created prior to February 2022, review the personnel on the COI Disclosure Status to ensure all responsible people are listed







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Upcoming Huron Upgrades and Implementation

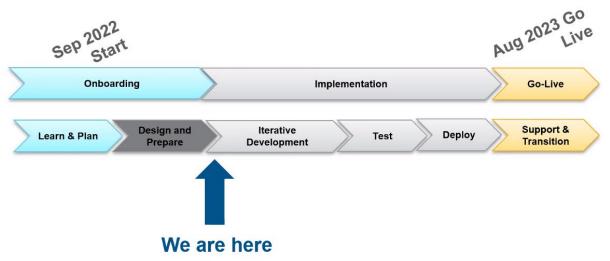
Raquel Zamora

Sr. Manager, IT, Research Intelligence and Data Infrastructure (RIDI)

IACUC Implementation

Extended Timeline





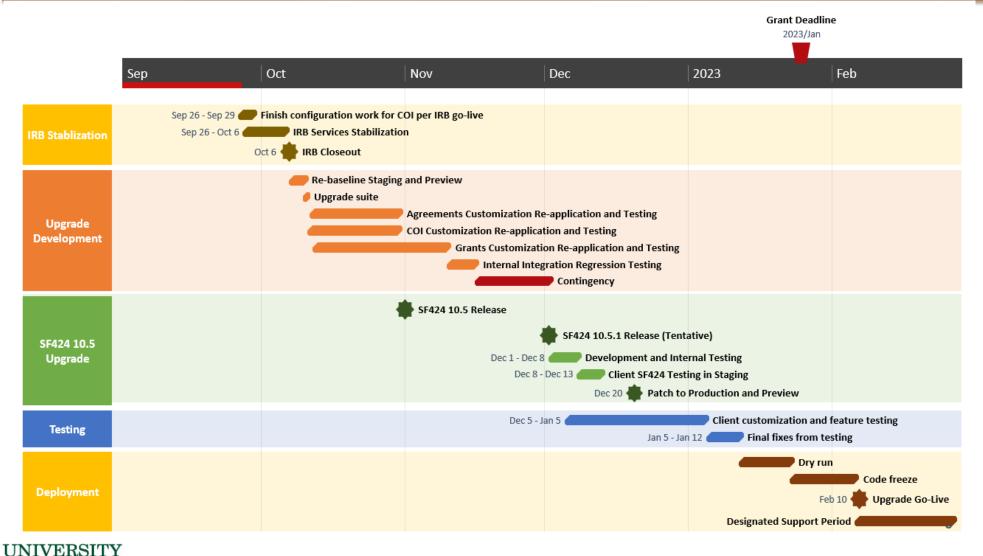
In progress:

- Presentation to Champions January 2023
- Process analysis
- Data transformation





IBISResearch Upgrades





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IBISResearch Upgrades

Date

Upgrade beginning on February 10th, 2023 at 6:00 PM EST.

Maintenance Window Duration

- All sites, including IRB, will be down starting Friday, February 10th at 6:00 PM EST and will be back online by Monday, February 13th at 8:00 AM EST.
- An all clear will be sent to users as soon as the upgrade is completed, and all sites are back online.

<u>Training Resources:</u> Related to the upgrade can be found at the following links:

- 1. Agreements 10.1 Help Links and Videos
- 2. Grants 10.1 Help Links and Videos
- 3. COI 10.3 Help Links and Videos

OVPRS Help Desk (305-243-2314; OVPRShelpdesk@miami.edu)





ORCID ID

OF MIAMI



Patty Atkinson

Research Navigator, Research Development + Education (RDE)

New Initiative to Implement ORCiD UM-Wide

Kick Off: Feb 9th

Sponsored by:













What is an ORCiD?

ORCID: Open Researcher and Contributor ID

- Registry of unique identifiers for researchers and scholars
- Persistent identifier to distinguish each researcher/scholar from all others
- Non-proprietary, not connected to specific database or entity

Transparent, mobile, and community-based







Why ORCiD?

Why Should Researchers Use ORCiD?

- Distinguish you from other researchers
- Get full credit for your work even when name or affiliation has changed
- Increase visibility of your work
- Keep track of all research outputs (publications, professional associations, grants, etc)
- Avoid re-entry of biographical & bibliographical data into multiple systems – ORCiD integrates with UM's <u>Faculty Success</u> and Scholarship@Miami (<u>Esploro</u>) profiles
- Align with funding organizations, publishers, associations, etc that are requiring ORCiD use.





How to Sign Up

Sign Up and Affiliate with UM

Use the ORCiD app developed for UM researchers.
 Available on phone, tablet, or computer.



 Walk through the steps to create a new or to connect an existing ORCiD to the University of Miami.

Connect your ORCID iD Your ORCID Account will be connected with Scholarship@Miami, the University of Miami Libraries' research information hub & institutional repository. Having an ORCID iD will raise the level of discovery for your work and help us create a complete bibliographic record of your research. Please choose one of the two options below to begin the process: I Have an ORCID iD I Don't Have an ORCID iD





ORCID ID

Questions? Please contact:

Angela Clark-Hughes

Director, Rosenstiel School Library Research Impact Strategist UM Libraries a.clark@miami.edu



Research Navigator
Office of the VP for Research & Scholarship
navigator@miami.edu







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AAHRPP Reaccreditation Site Visit

Kenia Viamonte

Director, Human Subject Research (HSRO)

AAHRPP Reaccreditation Update

Association for the Accreditation of Human Research Protection Programs (AAHRPP)

Scheduled for:

- Thursday, March 2nd and Friday, March 3rd
- Fully remote
- 2 site visitors
- 85+ interviewees
- Records review





AAHRPP Reaccreditation Update

General information everyone may be asked

- Who is the Institutional Official?
- Training requirements
- Reporting requirements
- Conflict of Interest
- Workflow
- Communication
- Systems
- Who to contact





Upcoming Site Visit

- **Domain I** (Institution- i.e. Compliance units, HRPP components, ancillaries, resources)- 40 interviewees
- Questions will surround policy & practice for:
 - Conflict of interest (Institutional & Researcher/ study team)
 - Research Pharmacy
 - Monitoring
 - Auditing
 - Scientific design/ feasibility
 - Applicable local laws
 - Contracts





Upcoming Site Visit

- Domain II (Institutional Review board members and staff) -35 interviewees
- Questions will surround policy & practice for:

Authority	Undue Influence
Suspensions/ Terminations	Reporting
Review Criteria/ Intervals	Expedited, Exempt, Full Board, Non Human Subject
Serious/ continuing noncompliance	Waivers/ Alterations of consent
FDA regulated research	Vulnerable Populations
Onboarding/ evaluations	Meeting operations/ preparations
Participant complaints	Adequacy of resources





Upcoming Site Visit

- Domain III (Investigators, Study teams)- 10-20 interviewees
- Questions will surround policy & practice for:
 - Protocol training
 - Recruitment
 - Consenting
 - Equitable selection
 - Reporting requirements
 - FDA regulated research





Questions

AAHRPP contact: Kenia Viamonte

kviamonte@miami.edu

Human Subject Research Office (HSRO) | University of Miami





Open Forum Questions





