

CLICK RESEARCH SOFTWARE

An Overview

 University at Buffalo
Research and Economic Development





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WELCOME



Some Background about Click



The Click Portal is a comprehensive software for managing research and streamlining workflow and compliance for research.



The Click Portal was implemented to assist principal investigators, students, compliance and research administration staff with administering sponsored programs



UB is leading the implementation of the Click Portal, working in collaboration with our vendor partner, Huron, and the PACS Leadership Team.

The Click Portal

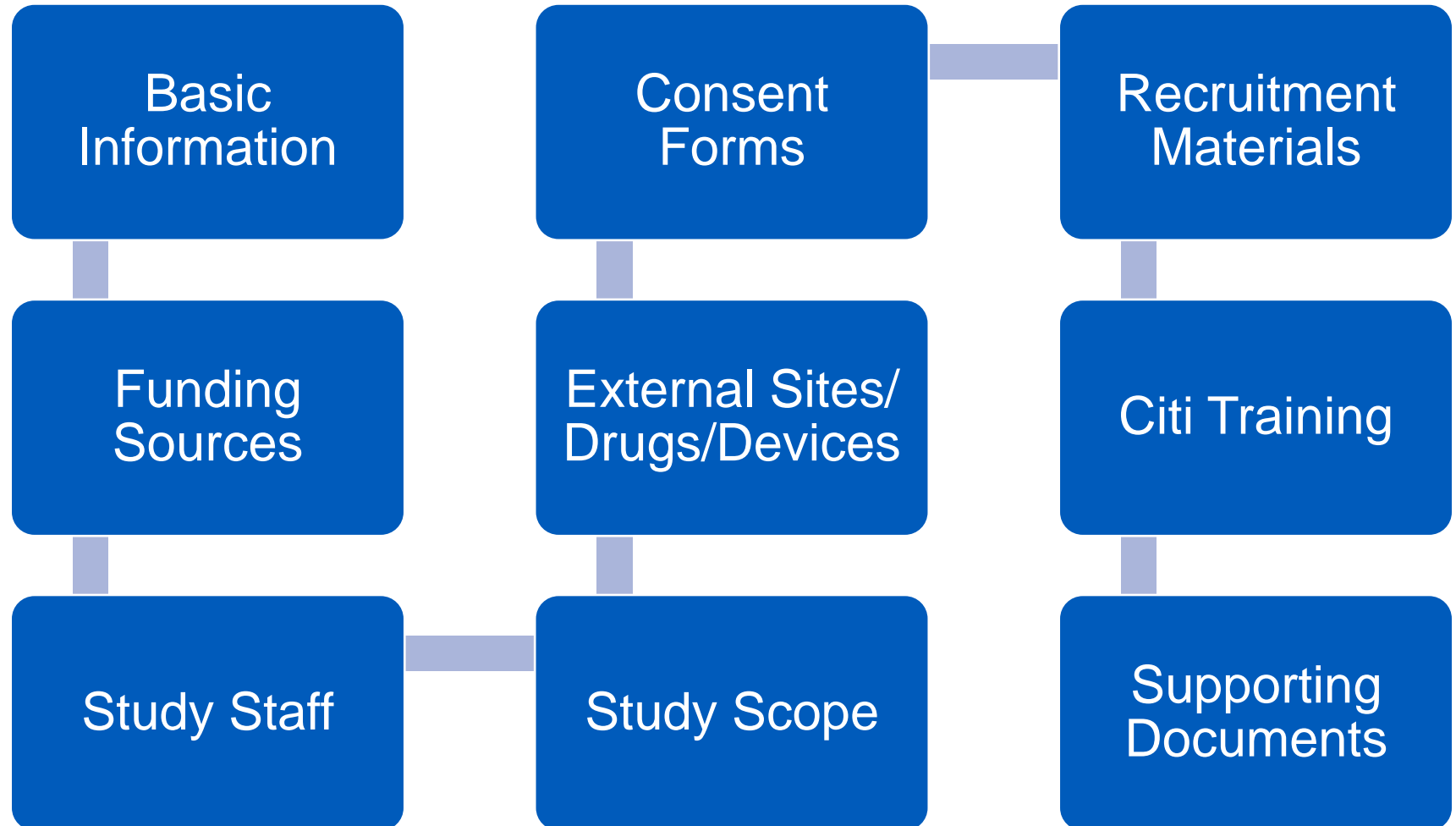
Click automates the submission, review, and approval processes while managing all major administrative aspects of the research and compliance lifecycle:



New Study Entry

SmartForms are a series of forms where you input specific information about a study. Based on your responses, it will determine what form/ questions to complete next

SmartForms



Smartform Basics

On the SmartForm pages, you will see:

1. Required fields (red asterisk)
2. Field help
3. Navigation bar (save or exit)
4. Jump to other pages in protocol
5. Ensure all required fields are completed

The screenshot shows a web form titled "Basic Information" with a navigation bar at the top containing "<< Back", "Save | Exit | Hide/Show Errors | Print...", and "Jump To: - Basic Information". The form contains the following fields and options:

- 1. * Select research team:** A dropdown menu showing "Spine Research Team".
- 2. * Title of protocol:** A text area containing "Spine research in animals".
- 3. * Short title:** A text input field containing "Spine research in animals".
- 4. * Non Scientific Summary of Research:** A text area containing "Spine research in animals".
- 5. * Principal Investigator:** A text input field containing "Rebecca Simms (pi)(030)", with "Select..." and "Clear" buttons.
- 6. * What is the intention of the animal protocol?:** A radio button group with options: "Breeding Only", "Experimental Research" (selected), "Field Research", "Holding Protocol", and "Teaching/Training". A "Clear" button is below the options.

Numbered callouts (1-5) are placed over the form to highlight these features. Callout 1 points to the asterisk on the first field. Callout 2 points to the asterisk on the fourth field. Callout 3 points to the top navigation bar. Callout 4 points to the asterisk on the fourth field. Callout 5 points to the bottom navigation bar.

Filtering and Sorting Data

Many pages contain tables you can filter and sort to help you find the data you want.


- Filtering reduces the list to only the data that meets the criteria. The advanced filter lets you combine multiple filter criteria together.
- Sorting displays the data in ascending or descending order by a particular column.



NOTE: To combine multiple filter criteria, such as *ID number*, *Name*, and *Date Created*, use the Advanced Filters.

Filtering Data

1. Select the column to **Filter by** from the drop-down menu. The menu lists only the columns you can **Filter by**.
2. In the next box, type the beginning characters for the items you want to find. If you do not know the beginning characters, type a % symbol as a wildcard before the characters.



Filter by

3. Click **Go** to apply the filter. The table shows only those rows that are an exact match. If you do not see the expected items in the list, click **Clear** in the **Filter by** area to remove the filter.



Filter by

NOTE: For examples and a list of operators you can use, click the **Help** icon.

The Click Portal Assists 6 Aspects of Research Management

The screenshot shows the Click Portal interface for the University at Buffalo. The navigation bar includes links for Home, Agreements, COI, Facilities, Grants, IACUC, IRB, and Safety. The user is logged in as Rebecca Simms (pi)(030). The main content area is divided into 'Links by Solution' and 'My Inbox'. The 'My Inbox' section is currently displaying 'SPO Tasks' and contains a table of tasks.

6 Modules Make up Click

Links by Solution

- Agreements
- COI
- Grants
- IACUC
- IRB
- Safety

My Inbox

Filter by

ID	Name	Date Created
FP00000013	test	6/30/2016 6:27 PM
DC00000004	Annual Disclosure Certification for Rebecca Simms (pi)(030) 2016	4/13/2016 3:57 PM
030TRA00000004	TRA 10-5-2016	10/5/2016 1:52 PM

Agreements Module

- ❖ Provides electronic system for management of research-related contracts and agreements.
- ❖ Two offices at UB provide oversight for the agreements process:
 - ❖ The Office of Sponsored Projects Services (SPS) assists researchers to prepare and submit proposals for sponsored funding and provides management of awarded funds to ensure compliance with applicable regulations and policies.
 - ❖ The Office of Science, Technology Transfer, and Economic Outreach (STOR) negotiates and executes all Material Transfer Agreements (MTA) and Non-Disclosure Agreements (NDA) in keeping with the university's research guidelines and its obligations to federal research sponsors.

Agreement Upload



1. Principal Investigator/Submitter:

Rebecca Simms (pi)(030)

2. Upload agreement draft (if available): ?

Agreement_for_030MTA00000009.doc(1.07)

3. Title or internal reference number: ?

template MTA form (commercial entity).

4. * Agreement type:

Material Transfer Agreement ([link to agreements definition](#))

If MTA, are the materials being provided/received with the present intention of preparing a joint venture?

Yes No

5. Purpose/Description of Agreement:

6. Supporting documents: ?

Name

There are no items to display

Benefits:

- Customized submission forms for various agreement types
- Eliminates the risk of storing information in individual email accounts
- Interfaces with other modules - simple tracking of relationships to studies and grant awards

COI Module

- Provides electronic system for management of research-related conflict of interest (COI) disclosures.
- Ensures the prevention of bias/ Helps avoid a circumstance in which a researcher's personal interest or relationships might compromise the integrity of research at the University
- The Compliance Team and Conflict of Interest Officer carefully review these disclosures, effectively manage any conflicts, and report to appropriate federal agencies as needed.
- Each Investigators must submit a Disclosure Statement that includes information pertaining to their significant financial interests (SFIs) and any sponsored travel that may be related to their research:
 - when submitting a grant application for research
 - prior to submitting human subject protocols for review
 - annually
 - immediately upon of discovering or acquiring a new SFI.

Rebecca Simms (pi)(030) | My Inbox | Logoff

Home Agreements **COI** Facilities Grants IACUC IRB Safety

COI Submissions > Annual Disclosure Certification for Rebecca Simms (pi)(030) 2016


Review Complete: Preparing Correspondence


Type: Annual Certification | ID# DC00000419 Discloser: **Rebecca Simms (pi)(030)** | Phone:

My Current Actions

View Disclosures

Printer Version

 Manage Ancillary Reviews

 Log Comment

My Disclosures

Meetings

Reports

COI Discloser Quick Reference

COI Advanced Reports Reference

COI QuickStart Guide

Summary History

Notes to Discloser

Your disclosures are currently under review. Please check your email account, listed in the system as , for any communication regarding the status of review. If further action is required, you will be notified at the above listed email address, and will have to log back into this system.

Summary

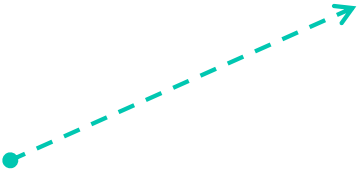
Last COI Training Date: 9/9/2013
Management Plan:
Snapshot:
Date Submitted: 8/3/2016
Review Due Date: 10/2/2016

Routing Information

Discloser has a Significant Financial Interest: Yes
 Discloser is Involved in Research: Yes
 Discloser has Institutional Responsibilities: Yes
 Discloser has Significant Obligations: Yes
Needs Review: Yes

My Disclosures

Export



Benefits:

- Ensures consistent processes
- Eliminates redundant data input by connecting COI input to study protocols and awards
- Fosters a culture of transparency

Click COI Module

The COI Module provides an electronic system for management of research-related conflict of interest disclosures. It ensures the prevention of bias and helps avoid a circumstance in which a researcher's personal interest or relationships might compromise the integrity of research at the University.

If you do not have a Click account, you can request access here:

<https://www.buffalo.edu/research/research-services/click-implementation.html>.

If you do not have access to your annual COI, you can request help here:

<https://www.buffalo.edu/research/about-us/staff-directory/ris/request-it-help.html>

UB's annual filing deadline is Nov. 1.



December 2015

Sunday	Monday	Tuesday	Wednesday	Thursday	Friday	Saturday
1	2	3	4	5	6	7
8	9	10	11 Veterans Day	12	13	14
15	16	17	18	19	20	21
22	23	24	25	26 Thanksgiving Day	27	28
29	30	1	2	3	4	5

Grants Module

- Provides electronic system for the management of grant award applications, budgets, milestones, contract terms, sub-recipients, and compliance status.
- Ensures consistent processes and reminders to researchers and staff for pending work items
- Integrates grant processes with key compliance issues such as IRB and COI
- The Office of Sponsored Projects Services (SPS) helps develop grant proposals, reviews and submits sponsored funding proposals, and assists in managing grant awards.



You Are Here: test

<< Back

Exit | Hide/Show Errors | Print... | Jump To: - Proposal Description & Contacts ▾

Continue >>

Proposal Description & Contacts

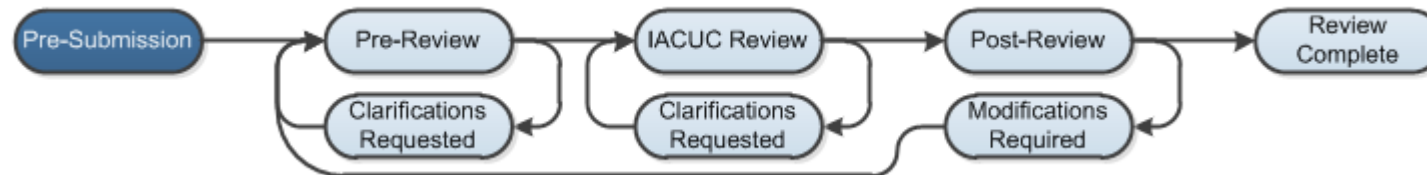
1.0	<p>* Short title of proposal: test</p>	<p><i>For NIH, maximum 200 characters allowed.</i></p>
2.0	<p>* Program Director / Principal Investigator / Project Lead / Fellow: Rebecca Simms (pi)(030)</p> <p>If a fellowship, please identify the Mentor:</p> <p>COI Disclosure: Biosketch: Other Support:</p>	<p>For single proposals: <i>Upload the following PDF documents pertaining to the PD/PI/Fellow:</i> - Biosketch</p> <p>* Note: Other Support: <i>Not usually required. Please refer to the Funding Announcement to see if this is required.</i></p>
3.0	<p>Research Coordinator:</p>	<p><i>Staff chosen for this question will have edit access rights to the proposal.</i></p>
4.0	<p>Administrative Contact:</p>	<p><i>Staff chosen for this question will have edit access rights to the proposal, if they are study staff.</i></p>
5.0	<p>* Select Sponsor: National Institutes of Health</p> <p>If Sponsor does not appear in list:</p> <p>If flow through, select Prime Sponsor:</p>	<p>For single proposals: <i>Sponsor list limited to the following categories:</i> - Industry - Foundation - Government Agency/Sponsor - Division/Department - Institution - State/Local Government</p> <p><i>If you do not find the name of the Sponsor in this list, select "TBD" from the list and enter the name in the next line.</i></p>

Benefits:

- Ensures submissions are accurate and complete
- Standardizes budget preparation and review
- Integrates with existing accounting and HR systems

IACUC Module

- Provides electronic system for management of the use of animal subjects in scientific and educational activities.
- Provides access to species, procedure, substance, training information and inspections
- Assists in the creation of reports
- The Institutional Animal Care and Use Committee (IACUC) reviews and approves all animal research to maintain nationally established standards to ensure that the University's research methods are ethical and compliant.



<< Back Save | Print... Continue >>

Basic Information

1. * **Select admin office:** ?

- IACUC Stony Brook
- IACUC University at Albany
- IACUC University at Buffalo

[Clear](#)

2. * **Title of protocol:**

Hypocholesterolemic and antiatherosclerotic potential of Basella alba leaf extract in hypercholesterolemia-induced rabbits

3. * **Short title:** ?

Basella alba leaf extract in hypercholesterolemia-in

4. * **Summary of research:** ?

Hypercholesterolemia is the major risk factor that leads to atherosclerosis. Nowadays, alternative treatment using medicinal plants gained much attention since the usage of statins leads to adverse health effects, especially liver and muscle toxicity. This study was designed to investigate the hypocholesterolemic and antiatherosclerotic effects of Basella alba (B. alba) using hypercholesterolemia-induced rabbits. Twenty New Zealand white rabbits were divided into 5 groups and fed with varying diets: normal diet, 2% high cholesterol diet (HCD), 2% HCD + 10 mg/kg simvastatin, 2% HCD + 100 mg/kg B. alba extract, and 2% HCD + 200 mg/kg B. alba extract, respectively. The treatment with B. alba extract significantly lowered the levels of total cholesterol, LDL, and triglycerides and increased HDL and antioxidant enzymes (SOD and GPx) levels. The elevated levels of

5. * **Principal Investigator:**

Rebecca Sims (pi)(050) [input] [input]

Benefits:

- Ensures compliance with federal regulations and regulatory reporting requirements
- Provides a library of standard substances and procedures
- Supports Annual/Triennial reviews



- Provides a streamlined way for researchers to create and submit protocols for studies involving human subjects.
- Initiates New Studies, Modifications/Continuing Reviews, and Reportable New Information
- The Institutional Review Board (IRB) approves, modifies, or rejects proposed research based on its perceived risks and benefits to prospective subjects.

IRB



You Are Here: PI1 - Cinnamon and type 2 diab...

<< Back

Exit | Hide/Show Errors | Print... | Jump To: Basic Info

Basic Information

1. * **Title of study:**

PI1 - Cinnamon bark, water-soluble cinnamon extract, and metformin as initial treatment

2. * **Short title:**

PI1 - Cinnamon and type 2 diabetes

3. * **Brief description:**

We are studying whether or not cinnamon bark or water-soluble cinnamon is an effective nutraceutical for the initial treatment of diabetes when compared to standard therapy of metformin 1000 mg (extended-release). This study will enroll 309 patients in randomized, doubled-blind, clinical trial and should be complete in a little over a year. Once consented, all subjects will have hemoglobin A1C, lipid panel, height, weight, and waist circumference measured along with standard demographic info. After randomization, they will receive one of the 3 medications from the pharmacy - thus investigators and subjects will be blinded as to their treatment. Subjects will be strictly counseled to have no further medication adjustments during this time. They will receive standard diabetic teaching. After 90 days of treatment, each subject will again have hemoglobin A1C, lipid panel, height, weight, and waist circumference measured. Subjects will bring in any remaining medication to determine adherence rates to the study protocol.

4. * **Principal investigator:**

Rebecca Simms (pi)(030)

5. * **Does the investigator have a financial interest related to this research?**

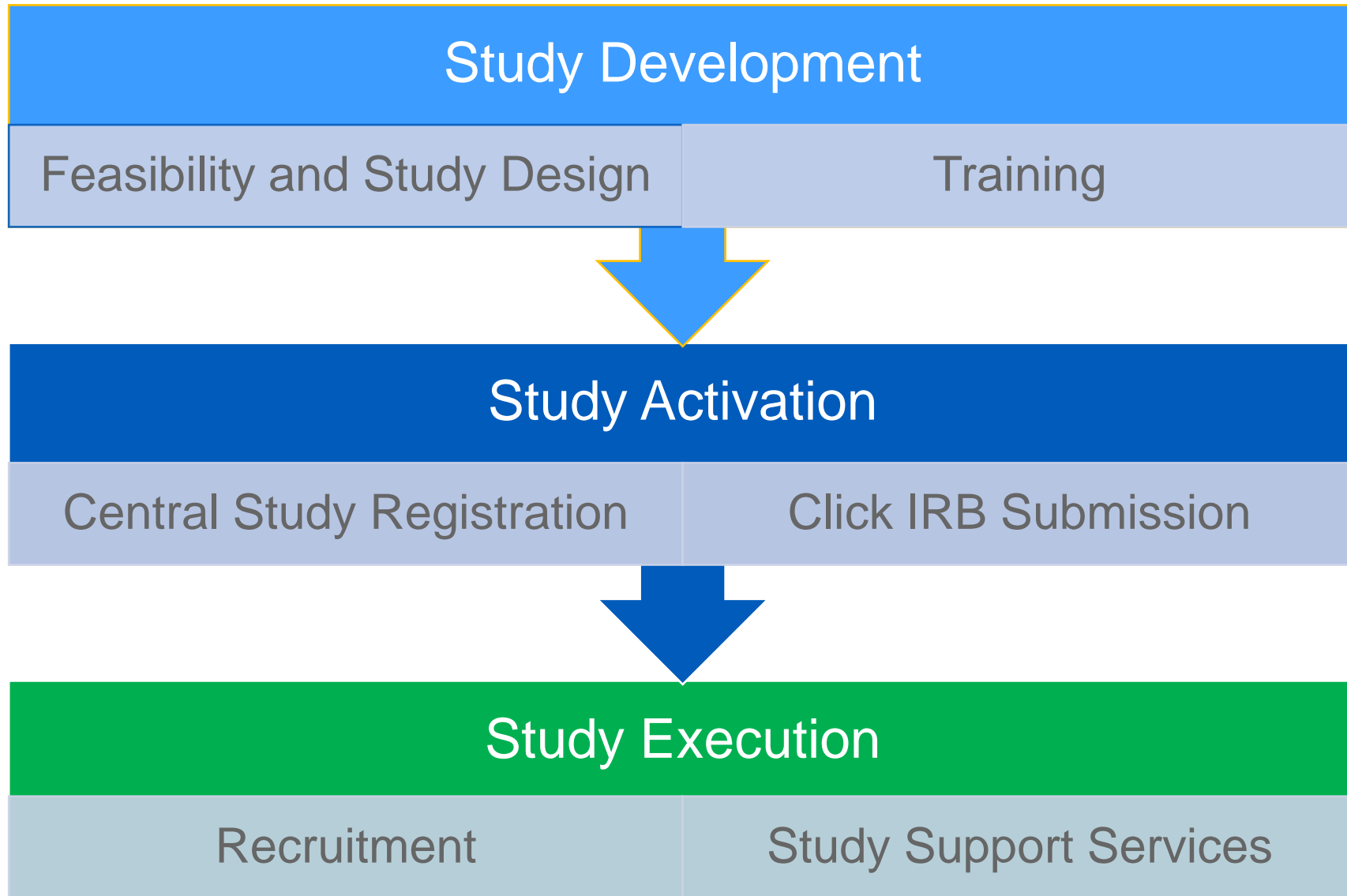
Yes No

6. * **Will an external IRB act as the IRB of record for this study?**

Yes No

Benefits:

- Ensures compliance
- standards
- Reduces approval turnaround times



Central Study Registration CSR:

This is the place to enter your study from any of the Health Sciences Schools, or if your study is deemed a clinical trial.

From the information you provide in CSR, the Clinical Research Office (CRO) will create your IRB submission in CLICK IRB.

It is important to note that CRO will never SUBMIT your study for you, they create it and then turn it over to you to add any outstanding documents that the IRB will need to review (data collection forms, surveys, consent forms and the like...) and submit the study for review when ready.

For help with CSR: contact Kim Brunton
716-888-4840, kbrunton@buffalo.edu

Safety Module

- Provides way for researchers to identify any hazardous materials that will be part of an experimental protocol:
 - Biosafety Hazards
 - Radiation
 - Chemicals
 - Stem Cells
- The Environment, Health and Safety (EHS) department anticipates, recognizes, evaluates, and controls all safety hazards at the University while striving to protect human health and the environment.



View: Click Safety Submission - 030BIO00000017

<< Back Exit | Hide/Show Errors | Print... | Jump To: - Basic Information Continue >>

▼ Reviewer Note

Type	Reviewer	Date Created	Date Modified
There are no items to display			

Basic Information

1. * **Select appropriate safety review:** ⓘ

Biosafety

2. * **Title of protocol:**

GLIOMA INITIATION BY HUMAN OLIGODENDROCYTE PROGENITORS

3. * **Short title:** ⓘ

GLIOMA INITIATION BY HUMAN OLIGODENDROCYTE PROGENITORS

4. * **Summary of research:** ⓘ

Project Overview:

Glioblastoma multiforme (GBM), a lethal brain tumor, likely arises due to oncogenic mutations affecting key signaling pathways that regulate cell growth and proliferation. To determine which pathway mutations are necessary to induce glioma initiation and also the pathways that trigger specific subtypes of GBM. As oligodendrocyte progenitor cells (OPCs) are the largest population of dividing progenitors in the human brain, it is thought that these progenitors represent the most likely source of glioma initiating cell.

The first experiments in this project are designed to establish the timing of tumor initiation of primary human OPCs following the introduction of several oncogenic mutations (e.g., EGFR, KRAS, and PTEN over-expression). In order to model human glioma initiation, human OPCs will be transduced with dominant-negative or shRNAi that target four of the major oncogenes (EGFR, KRAS, PTEN, Rb, and EGFRvIII). The effects of these oncogenic mutations will be first established via a series of in vitro experiments. Then to better model whether these mutations generate bone fide GBM-like tumor, infected OPCs will be injected into immunocompromised mouse brain (i.e. an orthotopic xenograft model similar to Auvergne, Simonsen, et al., 2004). Stereotaxic surgical techniques will be employed to implant 100,000 cells unilaterally into the lateral corpus collosum of 8-10 week old NOD.Cg-Prkdcscid Il2rgtm1Wjl/LtJ mice at various time points following injection to assess tumor formation in vivo.



Benefits:

- Ensures adherence to NIH and other federal, state, and local regulations
- Allows the tracking of Safety protocols in parallel with IRB and IACUC protocols
- Contains full review, committee, and meeting management functionality



There **Click Portal** and **CITI** will automatically add course information for study team members.

In order to ensure that your protocol is not delayed, you should verify that all study personnel have completed the appropriate **CITI Training**.

A screenshot of the Click Portal interface. The top left shows the "Click" logo with "a Huron solution" below it. Below the logo is a breadcrumb trail "You Are Here: [oinj]". A navigation bar contains a "<< Back" button, a menu with "Save | Exit | Hide/Show Errors | Print...", and a "Jump To: CITI Training" dropdown. The main content area is titled "CITI Training" and contains two sections. The first section, "1. PI Completed CITI Coursework:", lists "THOMAS WENDT" with a "View" link. Below this is a table with columns for Curriculum, Group, Stage, and Expire. The second section, "2. Study Team Members Completed CITI Coursework:", has a "Training Details" link and states "There are no items to display". A second navigation bar at the bottom is identical to the one at the top.

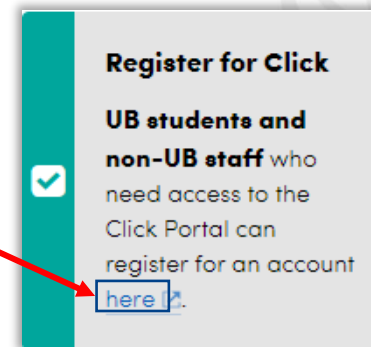
To log in to Citi trainings, navigate to <http://www.citiprogram.org>, enter your **UBIT name** and **Password**.

Getting a Click Account

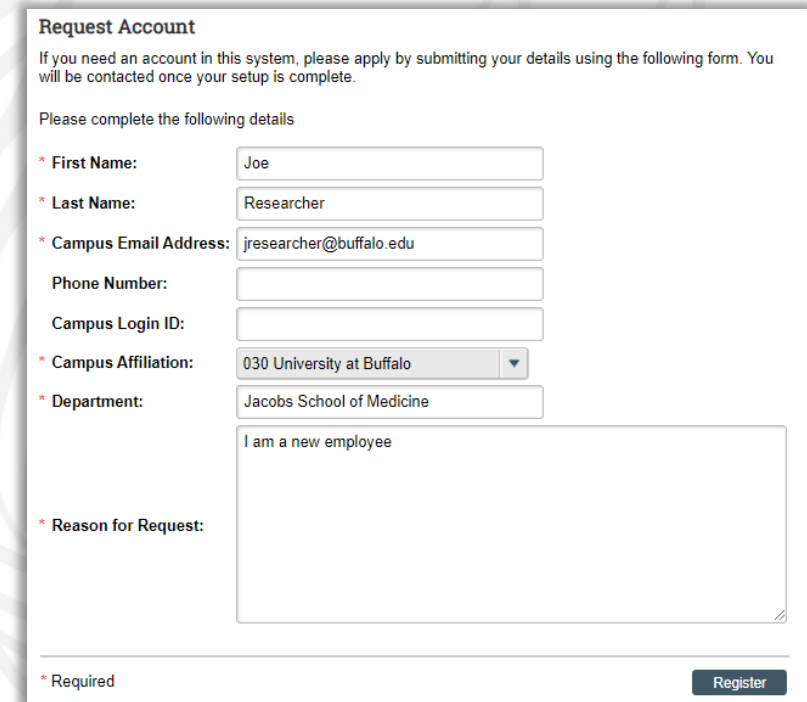
Accounts are created for faculty and staff of the University at Buffalo for access to the Click Portal. Logging into the system will be done using their **UBIT Name** and **Password**.

If you do not have an account, navigate to <https://www.buffalo.edu/research/research-services/training.html>.

1. Locate the **Register for Click** box on the right side of the page
2. Complete the **Request Account** form
3. Click the **Register** button.
4. You will receive an email notification when your account has been activated.



Register for Click
UB students and non-UB staff who need access to the Click Portal can register for an account [here](#).



Request Account
If you need an account in this system, please apply by submitting your details using the following form. You will be contacted once your setup is complete.

Please complete the following details

* First Name:

* Last Name:

* Campus Email Address:

Phone Number:

Campus Login ID:

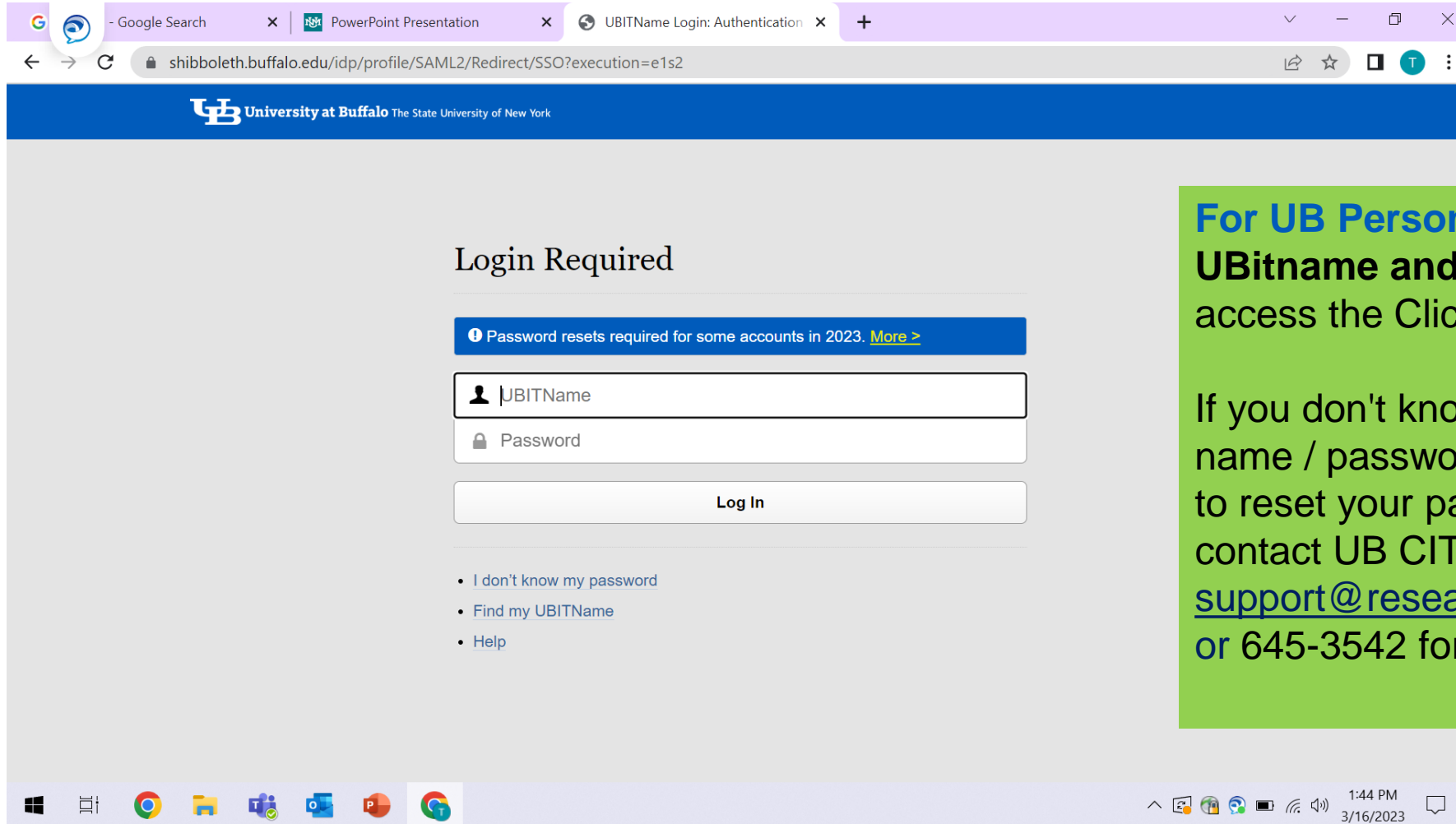
* Campus Affiliation:

* Department:


* Reason for Request:

* Required


Logging In





shibboleth.buffalo.edu/idp/profile/SAML2/Redirect/SSO?execution=e1s2

 University at Buffalo The State University of New York

Login Required

 Password resets required for some accounts in 2023. [More >](#)

 UBITName

 Password

Log In

- [I don't know my password](#)
- [Find my UBITName](#)
- [Help](#)

Windows taskbar: 1:44 PM 3/16/2023

For UB Personnel: Use your **UBITname** and **password** to access the Click Portal.

If you don't know your name / password or you need to reset your password, please contact UB CIT at support@research.buffalo.edu or 645-3542 for help.

During PRE-Submission

1. The research team will complete the SmartForm pages
2. The PI will then submit the study for review

Create New Study

Report New Information

Submissions

Meetings

Reports

Library

Help Center

In-Review	Active	Archived	New Infor
Filter by <input type="text" value="ID"/>			
ID	Name		
STUDY00000102	MLV		
STUDY00000113	MLV		
STUDY00000108	MLV		
STUDY00000104	MLV		
STUDY00000086	NE		
STUDY00000085	;kf		
6 items			

My Current Actions

Edit Study

Printer Version

View Differences

Submit

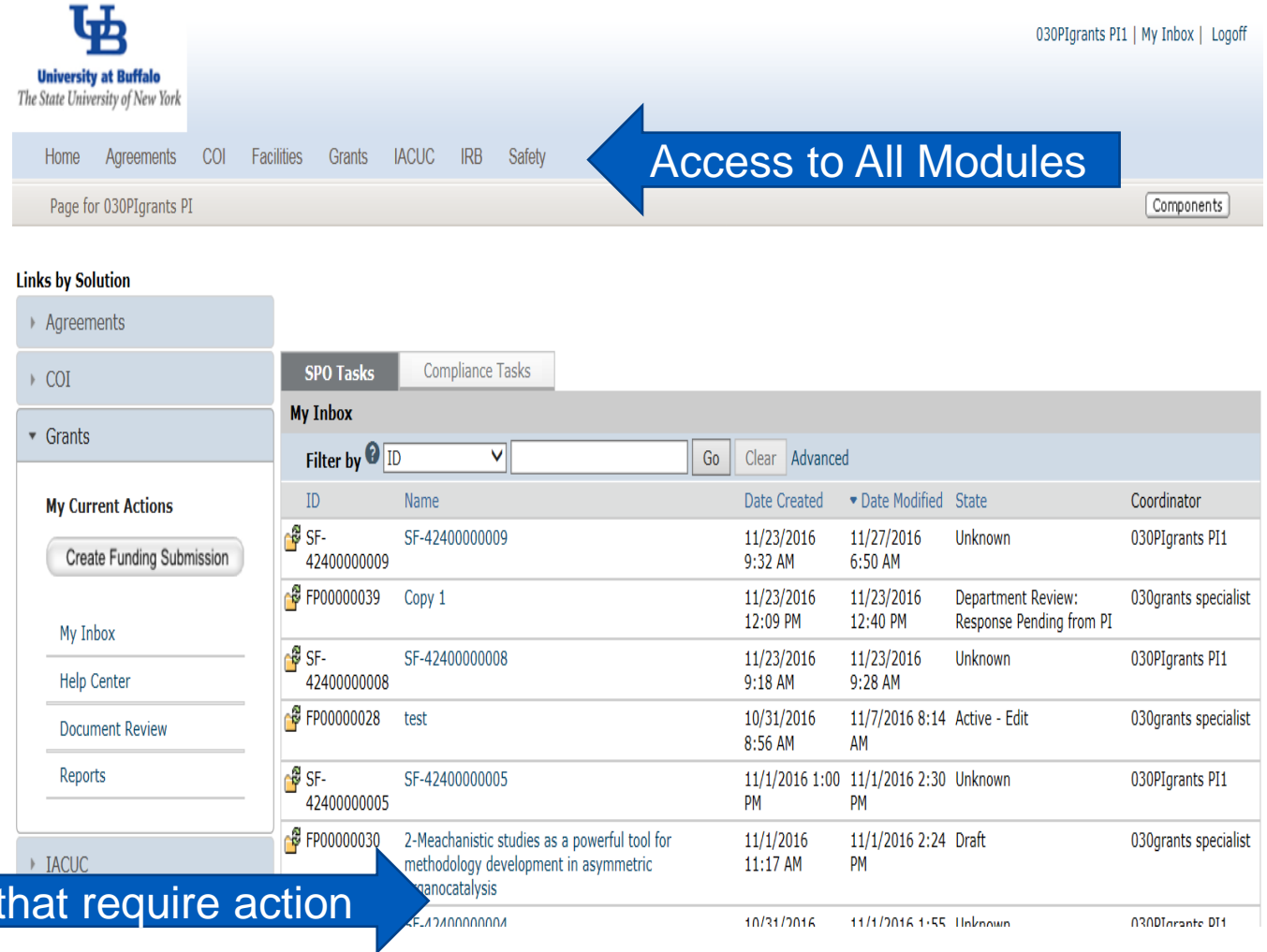
My Inbox

Each time you log into the **Click Portal**, you will be taken to your **Inbox**. This area of the portal contains a list of all submissions that currently require you to take an action.

My Inbox is divided into two tabbed pages:

- Compliance Tasks** -contains a list of all your submissions that currently require you to take an action.

- SPO Tasks** -contains a list of your Agreements and COI Disclosures that currently require you to take an action.



030PIgrants PI1 | My Inbox | Logoff

Home Agreements COI Facilities Grants IACUC IRB Safety

Page for 030PIgrants PI Components

Links by Solution

- Agreements
- COI
- Grants
 - My Current Actions
 - Create Funding Submission
 - My Inbox
 - Help Center
 - Document Review
 - Reports
- IACUC

SPO Tasks Compliance Tasks

My Inbox

Filter by ID Go Clear Advanced

ID	Name	Date Created	Date Modified	State	Coordinator
SF-42400000009	SF-42400000009	11/23/2016 9:32 AM	11/27/2016 6:50 AM	Unknown	030PIgrants PI1
FP00000039	Copy 1	11/23/2016 12:09 PM	11/23/2016 12:40 PM	Department Review: Response Pending from PI	030grants specialist
SF-42400000008	SF-42400000008	11/23/2016 9:18 AM	11/23/2016 9:28 AM	Unknown	030PIgrants PI1
FP00000028	test	10/31/2016 8:56 AM	11/7/2016 8:14 AM	Active - Edit	030grants specialist
SF-42400000005	SF-42400000005	11/1/2016 1:00 PM	11/1/2016 2:30 PM	Unknown	030PIgrants PI1
FP00000030	2-Mechanistic studies as a powerful tool for methodology development in asymmetric organocatalysis	11/1/2016 11:17 AM	11/1/2016 2:24 PM	Draft	030grants specialist
SF-42400000004	SF-42400000004	10/31/2016	11/1/2016 1:55	Unknown	030PIgrants PI1

Submissions that require action

When you first log in to Click, you will be taken to the **Inbox**

- From My Inbox, you will see:
1. Submissions that require action
 2. State in the review process

Click a Huron solution

Rebecca Simms (pi)(demo) | My Inbox | Logoff

Home Agreements COI Facilities Grants IACUC IRB Safety

Page for Rebecca Simms (pi)(demo) Components

Links by Solution

- Agreements
- COI
- Grants
- IACUC
- IRB
- Safety

Compliance Tasks SPO Tasks

My Inbox

Filter by ID Go Clear Advanced

ID	Name	Date Created	Date Modified	State	Coordinator
CR00000001	Continuing Review for Study STUDY00000004	6/5/2015 6:26 PM	6/9/2015 2:55 AM	Clarification Requested (Pre-Review)	Orlando Max (irbc)(demo)
AMEND201500001	Amendment for PROTO201500004	6/5/2015 7:55 PM	6/5/2015 7:55 PM	Pre-Submission	
PROTO201500002	rDNA for Mice	6/2/2015 7:07 PM	6/3/2015 3:23 PM	Pre-Submission	
PROTO201500001	DNA for Mice	6/2/2015 7:07 PM	6/2/2015 7:07 PM	Pre-Submission	
STUDY00000001	ER Re-Admission Study	6/1/2015 9:27 PM	6/2/2015 7:03 PM	Pre-Submission	

5 items page 1 of 1 25 / page

To open a study that you are associated with, click on the study ID.

This will take you to the study workspace.



Home Agreements **IRB** Safety

IRB

IRB

Create New Study

Report New Information

Submissions

Meetings

Reports

Library

Help Center

In-Review				
Active				
Archived				
New Information Reports				
All Submissions				
Filter by [?] ID <input type="text"/> <input type="button" value="Go"/> <input type="button" value="Clear"/> <input type="button" value="Advanced"/>				
ID	Name	Date Modified	State	
STUDY00000102	MLW Test Study	3/30/2016 3:00 AM	Pre-Review	
STUDY00000113	MLW Manual 3.28.16	3/29/2016 1:18 PM	Pre-Submission	
STUDY00000108	MLW Test Study 2.25.2016	2/25/2016 4:41 PM	Pre-Submission	
STUDY00000104	MLW Sample Study 2/18/2016	2/24/2016 4:00 PM	Pre-Review	
STUDY00000086	NEW SUBMISSION	2/2/2016 2:08 PM	Pre-Submission	
STUDY00000085	;kfd;kg;glk;	2/2/2016 2:06 PM	Pre-Review	

6 items page 1 of 1

Study Workspace

This is where anyone with access to the submission can find:

- Details regarding the study
- State in the review process
- Actions that can be taken in the current state
- Tabs with information regarding:
 - **History**
 - **Attached documents**
 - **Important contacts**
 - **Submission information**
 - **Related projects**

The Workspace displays key information about a particular study or submission.



Pre-Review

STUDY0000005 : Study on Improving a condition

IRB office: IRB 1

Entered IRB: 6/9/2015 3:01 AM
 Initial approval:
 Effective:
 Approval end:
 Modified: 6/9/2015 3:05 AM

Principal investigator: Rebecca Simms (pi)(demo) ← 1
Submission type: Initial Study
Primary contact: Rebecca Simms (pi)(demo)
IRB coordinator: Orlando Max (irbc)(demo)

My Current Actions

[View Study](#)
[Printer Version](#)
[View Differences](#)

[Add Related Grant](#)
[Withdraw](#)
[Discard](#)

History | Funding | Project Contacts | Documents | Reviews | Snapshots ← 4

Filter by: Activity [v] [Go] [Clear] [Advanced]

Activity	Author	Activity Date
Response Submitted	Simms (pi)(demo), Rebecca	6/9/2015 3:05 AM
Clarification Requested	Max (irbc)(demo), Orlando	6/9/2015 3:02 AM
IRB Coordinator Assigned	Max (irbc)(demo), Orlando	6/9/2015 3:02 AM

Assigned to Orlando Max (irbc)(demo)

1. Study details
2. The state in the review process
3. Actions you can take
4. Tabs with information about the study and the review process

1. Navigate to <http://www.buffalo.edu/research>
2. Locate the QUICK LINKS section of the page:



A screenshot of a web browser displaying the University at Buffalo research website. The browser's address bar shows "buffalo.edu/research.html". The page features a header with a news article titled "UB launches artificial intelligence institute". Below the header, there are four main sections: "HOW UB SOLVES PROBLEMS" with a photo of Christina Orsi, "NEED TO KNOW" with a list of recent events, "QUICK LINKS" (circled in red) with a list of navigation options, and "HIGHLIGHTED EVENT" with a "Covid-19: Research Guidance" button. The "CLICK Portal Login" link in the QUICK LINKS section is highlighted in yellow. The browser's taskbar at the bottom shows various application icons and the system tray with the date and time (3:46 PM, 2/16/2023).

3. Click on the [Click Portal Login](#) link
4. Enter your UBITName and Password and click the Log In button.

Questions?

- For technical questions regarding Click:
 - Tracie Groves, traciegr@buffalo.edu
- For assistance with UBITNames and Passwords:
 - Electronic Research Administration, support@research.buffalo.edu

