



Handbook for Researchers

Link: <https://samc.my.irbmanager.com/Login.aspx>

Contents

Logging into IRBManager	3
Dashboard At-A-Glance	4
Changing Your Password and more	5
Creating a New Study Submission.....	6
Previewing a Form	6
Jumping Between Pages in a Form	6
Adding Collaborators to a Submission Form.....	7
Study Home Page At-A-Glance	8
Submitting Forms on Active Studies	9
Electronic Signatures	10
Checking on Submission Status	11
NEED HELP?.....	11

Logging into IRBManager

To log into IRBManager, enter the following link in your browser:

<https://samc.my.irbmanager.com/Login.aspx>

Enter the *username* and *password* you were provided. Enter the Client name as **samc**, if not already displayed.

Saint Alphonus
A Member of Trinity Health

Login

User Name

Password

Client **samc**

[Forgot Password?](#)

Copyright ©2000-2025 Tech Software. All Rights Reserved.
2025.10.8265.0/Release/3375223 | GCWBWS1 | 2025-12-05 21:25:49Z | 0.009s

Powered By IRBManager

Upon logging in, you will see your dashboard.

Trinity Health

Home Find Study (Ctrl+Q) Help PI's Settings Sign off

My Studies (20 Active)

- You are associated with **20 active** Studies and **37 total** Studies.
- You are the PI for **20 active** and **37 total** Studies.
- You are the Coordinator for **0 active** and **1 total** Studies.
- There are **4 studies** expiring in the next 90 days.
- The next study to expire is **22-1017-Test-GR**.

xForms (0 Active)

- You have **0 unsubmitted** xForms.
- You have **0 xForms** being processed at a later stage.

Events (10 Open)

Only show events where I am:

- You have **1 Amendment** events.
- You have **3 Exempt/NHSR** events.
- You have **4 New Study Submission** events.
- You have **1 Personnel Change** events.
- You have **1 UP / Protocol Deviation** events.
- You have **10 Total Open** events

My Studies (20 Active)

Study	Site	PI	Study Title	Expires	Status
16-1010-test-GR	Trinity Health Grand Rapids	Test, PI	New Test Study for Invoicing Purposes	11/14/2023	Active
17-0721-test-GR	Trinity Health Grand Rapids	Test, PI	HUD Healthy Testing of IRB Application	N/A	Active

Notices

IRB Meeting Dates

IRB Meeting Dates	New Submission Deadlines
August 22, 2023	August 4, 2023
September 26, 2023	September 8, 2023
October 24, 2023	October 6, 2023
November 28, 2023	November 10, 2023
December 19, 2023*	December 1, 2023

*Meeting occurs third week of the month due to a conflicting holiday.

The IRB is no longer accepting QI submissions.
Please complete the **QI vs Research Determination Checklist** and, if all checkmarks are in the **QI** column, retain a copy for your records. **IF ANY checkmarks are in the Research column, you must submit the appropriate IRB Application with the required documents BEFORE any data collection work commences.**

For assistance with IRBManager, contact Cindy Johnston at johnstom@trinity-health.org or 616.685.6198.

Dashboard At-A-Glance



Actions
Add a New User
Exempt or NHSR Application
IRB Application for Initial Review
Start xForm
Show Sponsor Ids

Recent Items
16-0304-Test-GR
21-0910-4-MG
17-0817-Test-SM
17-0825-9-Test-SM
33-3333-33-SM
99-9999-99-SM

Messages
Welcome to IRBManager for Trinity Health West Michigan locations.

My Docs & xForms
0 Attachments
2 xForms

The left sidebar contains 4 separate sections.

Actions is where you can access xForms (electronic submission forms) for *NEW* submissions.

Recent Items allows quick access to anything you've recently viewed in IRBManager.

Messages welcomes you to IRBManager.

My Docs & xForms displays xForms and Attachments.



Home

My Studies

Studies (20 Active)

- You are associated with **20 active** Studies and **37 total** Studies.
- You are the PI for **20 active** and **37 total** Studies.
- You are the Coordinator for **0 active** and **1 total** Studies.
- There are **4 studies** expiring in the next 90 days.
- The next study to expire is **22-1017-Test-GR**.

xForms (0 Active)

- You have **0 unsubmitted** xForms.
- You have **0 xForms** being processed at a later stage.
- There are **1 xForms** awaiting your attention.

Events (10 Open)

Only show events where I am:

- You have **1 Amendment** events.
- You have **3 Exempt/NHSR** events.
- You have **4 New Study Submission** events.
- You have **1 Personnel Change** events.
- You have **1 UP / Protocol Deviation** events.
- You have **10 Total Open** events

My Studies (20 Active)

Study	Site	PI	Study Title
16-1010-test-GR	Trinity Health Grand Rapids	Test, PI	New Test Study for Invoicing Purposes
17-0721-test-GR	Trinity Health Grand Rapids	Test, PI	HUD Healthy Testing of IRB Application
17-0817-Test-GR	Trinity Health Grand Rapids	Test, PI	A funny little study created for testing purposes

The center of your dashboard also contains 4 sections.

Studies displays the number of studies you are associated with, your role, and the next study due to expire.

xForms will display the number of unsubmitted xForms, those being processed at a later stage, and **those awaiting your attention**.

Events specifies the number and type of open study events.

My Studies lists all studies you are associated with, the PI, the expiry date, and the study status. **Click on the study number to open a study.**



Notices

Welcome to IRBManager at SAMC and SAHS

If you are associated with any studies, you will find them listed below. Click on the study link to start an xForm for a particular study.

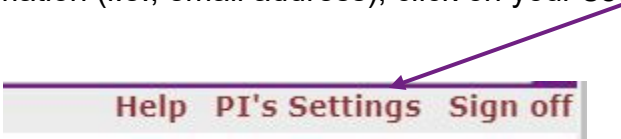
Need to update your Human Subjects Protections training? Go to citiprogram.org and affiliate with **Trinity Health - Michigan Region** to access Trinity standard curriculum. If you have completed CITI training within the last 5 years at another institution, please upload your completion documents to your user profile for review.

Contact THWIRB@saintalphonsus.org or (208) 367-6350 if you have questions or require assistance. Thank you!

Notices is where the IRB Office can post information that may be of help to research staff, including meeting and submission deadline dates, special notices, and contact information.

Changing Your Password and more

To change your password or contact information (i.e., email address), click on your **settings** at the top right side of your landing page.



Help PI's Settings Sign off

It is important to keep your contact information current, especially your email address, as all communication generated in IRBManager is sent via email.

My Settings

Edit Settings

Change My Password

Change My Profile

My Phone Number(s)

My Address(es)

My Expirations

My Attachments

Last 25 Logins

Email Signature

Turn on Dark Mode

Reset Dashboard

Switch Dashboard

Creating a New Study Submission



Actions

- Add a New User
- Exempt or NHSR Application
- IRB Application for Initial Review
- Start xForm
- Show Sponsor Ids

To submit a new application for review, click on the form name, if listed, on the left sidebar of your dashboard or select *Start xForm* and make your selection from the list of the available xForms.

Previewing a Form

To preview a form, click on the printer icon next to the form name. **Please note that all questions may not be applicable to your study.**

Select xForm to start	
Action	Form (Click to start)
	Exempt/NHSR Application
	IRB Application for Initial Review
	IRBManager NEW USER Request

Jumping Between Pages in a Form

The submission forms are built so that you cannot move forward to the next page until all questions on the current page are answered. However, you do have the ability to jump between pages of any form by clicking on the down arrow next to the title of the page you are on and selecting another page to jump to. You can select *Save for Later* at the bottom of any page and return to the form later to resume completion.

The screenshot shows a form page titled "Study Header" with a dropdown menu open. The dropdown menu lists various sections of the form: Funding, Principal Investigator, Additional Study Personnel, Study Sites / Other IRB Review, Study Specific Information, Subject Privacy/Confidentiality, A/V or Photographing, Recruitment Process & Materials, Vulnerable Populations, Access, Use, Disclosure of PHI, Invoicing, Study Documents, Investigator/Study Staff Documents, Confidentiality Agreement, and Application Data Entry Complete. At the bottom of the dropdown is "Check & Submit Form". The main form content includes a yellow warning box about IRB fees, a "Submitter" section for Cindy Johnston, and a "Full Study Title" section. The page number "Page 1 of 16" is visible in the top right corner.

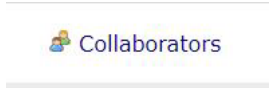
You will find your unsubmitted form on your dashboard.

xForms (1 Active)

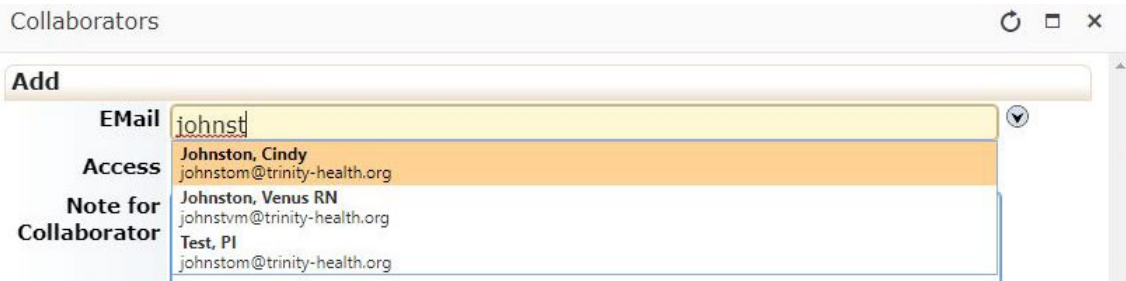
- You have **1 unsubmitted** xForms.

Adding Collaborators to a Submission Form

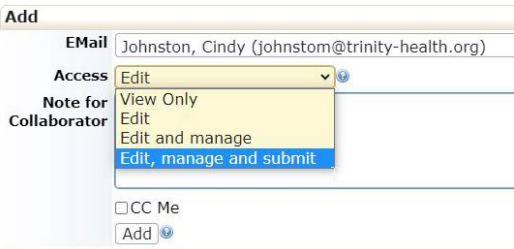
A collaborator can be added to any form. If you initiate a form and want to allow another individual access to review, edit, and/or submit the form, simply [click on the Collaborators link at the top of any page of your form](#).



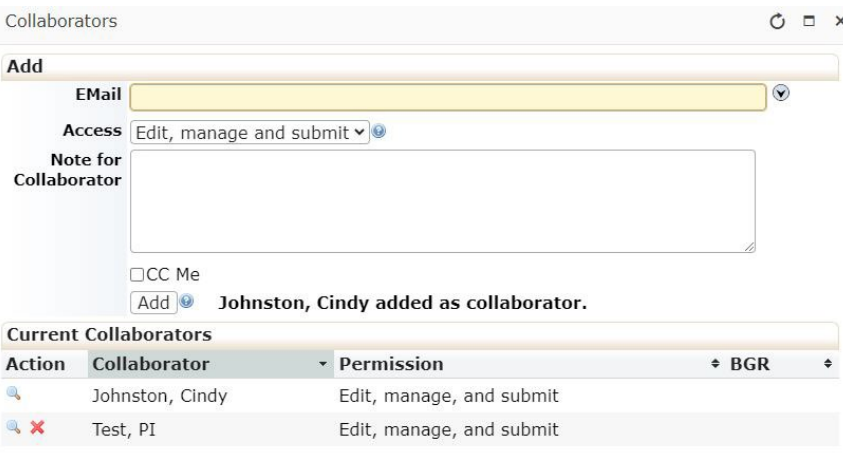
A new window will open. As you begin to enter the email address, the name of the individual should appear. Click on the name of the individual to add as a collaborator.



Next, select the type of access you would like to grant that individual. Add any specific notes for the collaborator if applicable, and click on the **Add** button. An email will be sent to the individual notifying them they have been added as a collaborator to your form.



You will see the individual has been added to the form and their level of access. The list of current collaborators is also updated.



NOTE: The author of the form can also **REMOVE** any collaborator by simply clicking on the red **X** next to the individual's name.

Study Home Page At-A-Glance

To view a specific study, click on the study number under “My Studies” on your dashboard.

Study: 26008 Committee: Test Committee Category: Department: Surgery Last Review: Agent Types: Study Title: Testing expedited approval 2018 Expedited Categories: (2) Risks to subjects are reasonable in relation to anticipated benefits, if any, to subjects, and the importance of the knowledge that may reasonably be expected to result. In evaluating risks and benefits, the IRB should consider only those risks and benefits that may result from the research (as distinguished from risks and benefits of therapies subjects would receive even if not participating in the research). The IRB should not consider possible long-range effects of applying knowledge gained in the research (e.g., the possible effects of the research on public policy) as among those research risks that fall within the purview of its responsibility. Funding Source: IND/IDE/HDE Number(s): Level of Risk: Minimal Risk Significant/Non-Significant Risk: Conditions: Other	Sponsor(s): Sponsor Id: Grants: Next Review: Year: 2026 Children Risk Level: Hospital Departments: IRB of Record: Other External Sites Involved: Study Type: Tissue/Biologic
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Study-Site	Site(s): SAHS - Saint Alphonsus Health System Location(s): Boise Status: Approved Approval: January 27, 2026 for 12 months Initial Approval: January 27, 2026 Approved Number of Participants: 5 Initial Review Level: Expedited Review Vulnerable Populations: Non-English Speaking	PI: Test, Investigator Additional: N Expiration: January 26, 2027 Other Expirations: Closure Date: Latest Review Level: Expedited Review
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Reviews on Open Events (1)						
Action	Event	Type	Reviewer	Review Item	Outcome	Due
	New Research Submission	Expedited Review	Committee Member, Test	New Research Application	Approve	01/3

Events (1)						
Action	Event	Att	Instance/UDF	Start	Complete	Approval Date

The top section is the **Study** section containing basic study information including the study title, assigned study #, name of the reviewing IRB, sponsor, type of review, level of risk and more.

Study Reference Documents will list the current approved protocol.

The **Study-Site** section primarily reflects the study site, study status, approval dates and name of the PI.

The **Study-Site Contacts** section lists all additional research study staff associated with the study.

The **Events** section lists every event submitted and reviewed by the IRB. The most recent submission is listed at the top with all other events listed in reverse chronological order. To the right of the event column is "ATT" where you will find all attachments associated with that specific event.

The **Study-Site Emails** section contains emails generated in IRBManager specific to the study.

Submitting Forms on Active Studies

To submit an amendment, continuing review report, unanticipated problem, item of information, etc. for an active study, **you must be in that specific study**, so the system knows which study to associate your request with. To open an active study, click on the study number under **“My Studies”** on your dashboard.

Study 26008-SAHS (IRB)

Study: 26008 Committee: Test Committee Category: Department: Surgery Last Review: Agent Types: Study Title: Testing expedited approval 2018 Expedited Categories: (2) Risks to subjects are reasonable in relation to anticipated benefits, if any, to subjects, and the importance of the knowledge that may reasonably be expected to result. In evaluating risks and benefits, the IRB should consider only those risks and benefits that may result from the research (as distinguished from risks and benefits of therapies subjects would receive even if not participating in the research). The IRB should not consider possible long-range effects of applying knowledge gained in the research (e.g., the possible effects of the research on public policy) as among those research risks that fall within the purview of its responsibility.	Sponsor(s): Sponsor Id: Grants: Next Review: Year: 2026 Children Risk Level: Hospital Departments: IRB of Record: Other External Sites Involved: No Study Type: Tissue/Biologic
Funding Source: IND/IDE/HDE Number(s): Level of Risk: Minimal Risk Significant/Non-Significant Risk: Conditions: Other	

[Study-Site](#)



Actions

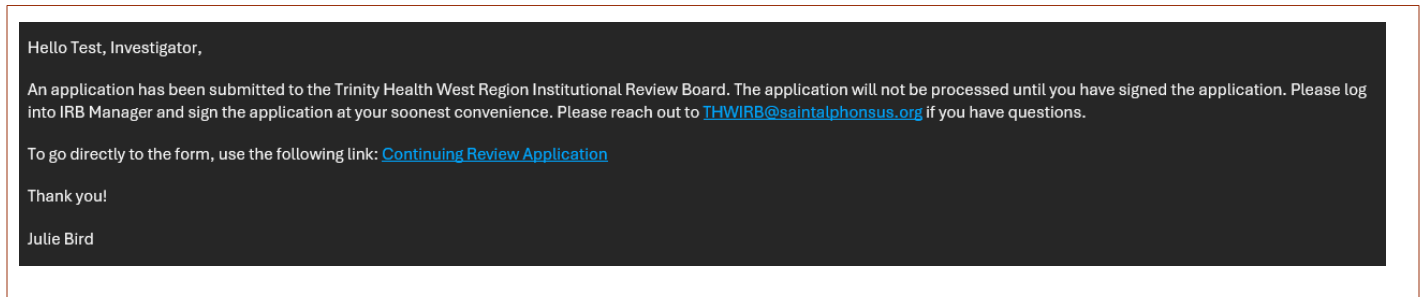
- Study*
- Update
- Add Attachment
- Add Contact
- Add Study-Site

- Study-Site*
- Update
- Add Attachment
- Add Contact
- Add Event
- Add Note
- Add Related Study-Site
- Add Animal
- Expirations
- Generate Doc
- Send EMAIL
- Start xForm
- xForms (0)

Once the study is open, select *Start xForm* under **“Actions”** on the left sidebar, and select the appropriate form from the list of available options.

Electronic Signatures

When a submitter, who is not the PI, submits a form, the PI will receive an email notification that a form was submitted and requires their electronic signature. The PI can go directly to the form by clicking on the link contained in the email (example below). They may also open the form by clicking on "1 xForms awaiting your attention" displayed on their dashboard.



On **new study submissions** and **continuing review reports**, two (or more) signatures are required. The initial signature verifies the information in the form and the signing off on PI Assurance. If a Waiver of HIPAA is requested, the PI will be required to sign off on that as well.

After the PI has provided the required electronic signatures on the submitted form, they may wait 30 seconds or so and refresh their dashboard and the Forms Section should again reflect there is "1 xForms awaiting your attention" or they may wait until they receive an email notification advising their signature is required on financial conflict of interest (FCOI). Click on the link to open the form and provide your electronic signature on FCOI or to complete the FCOI disclosure form. An email is also sent to all research personnel requiring their electronic signature on FCOI as well (see example below).

A new exempt submission was submitted to the Mercy Health Regional Institutional Review Board (IRB). Before this submission can go to the IRB, all research study staff are required to disclose any conflicts of interest and provide their electronic signature. This submission cannot move forward until all signatures are obtained.

If you **do NOT** have any conflicts to disclose, sign off next to your name at the bottom of the Conflicts of Interest page.

If you **DO have a conflict to disclose**, please click on the link below to complete the Conflicts of Interest Disclosure form.

Click [Exempt/NHSR Application](#) to go directly to the form.

If you have questions or require assistance with IRBManager, please contact the IRB Administrator at 616.685.6198.

If any research staff are **not affiliated** with Trinity Health, the submitter should note this where indicated in the application and those individuals will receive an additional email requesting their electronic signature on a Confidentiality Agreement.

NOTE: A study is not fully submitted until ALL research staff have provided the required signatures and the status of the submission is in the "NOTIFY IRB" stage.

Checking on Submission Status

To check on the status of a submission, click on the link of xForms being processed at a later stage found on your dashboard.

My Studies

Studies (9 Active)

- You are associated with **9 active** Studies and **17 total** Studies.
- You are the PI for **3 active** and **7 total** Studies.
- You are the CC Recipient for **0 active** and **1 total** Studies.
- You are the Coordinator for **4 active** and **7 total** Studies.
- You are the Research Assistant for **2 active** and **3 total** Studies.
- There are **1 studies** expiring in the next 90 days.
- The next study to expire is **33-3333-33-GR**.
- You are the reviewer for **0 active** and **5 total** Studies.
- Committee Mercy Health Regional IRB has **83 active** and **955 total** Studies.
- Committee RCD Staff has **0 active** and **0 total** Studies.
- Committee Test IRB Committee has **16 active** and **24 total** Studies.

xForms (2 Active)

- You have **1 unsubmitted** xForms.
- You have **1 xForm** being processed at a later stage. 
- There are **1 xForms** awaiting your attention.

Events (10 Open)

Only show events where I am:

- You have **1 Amendment** events.
- You have **4 Exempt/NHSR** events.
- You have **1 New Study Submission** events.
- You have **1 Personnel Change** events.
- You have **1 Safety Report/Adverse Event** events.
- You have **1 Study Closure** events.
- You have **1 UP / Protocol Deviation** events.

You have **10 Total Open** events

Each submission will display the stage it is in, in the IRB review process.



Action	Form	Identifier	Owner	Stage
	Major Protocol Deviations (Draft)		UP / Protocol Deviation 17-0825-9-Test-GR	Under Review

NEED HELP?

Contact the IRB Office at THWIRB@saintalphonus.org or julie.bird@saintalphonus.org.