

Administrative Check In

Human Research Protection Program (Last Updated:12/14/2023)



This PowerPoint will guide you through how to submit an Administrative Check In in Huron.

TEXAS A&M UNIVERSITY.

Getting started

- 1. Navigate to the IRB workspace
- 2. Select Submissions tab
- 3. Select All Submissions tab
- 4. Note: **Filter by** allows you to sort through your studies by name, PI first and last name, and submission type.
- 5. Open your study by selecting the **folder symbol** or the **name** of the study.





Reportable New Information

1. Select Reportable New Information

Note: the Administrative Check In for a study is submitted via the Reportable New Information form in Huron.

Approved

Entered IRB: 12/1/2022 11:25 AM Initial approval: 12/1/2022 Initial effective: 12/1/2022 Effective: 12/9/2022 Approval end: 11/30/2023 Last updated: 4/6/2023 10:44 AM

Next Steps

1

View Study	
Printer Version	
Create Modification/CR	
Report New Information	



Reportable New Information

- When completing the **Reportable New** Information page:
 - All questions marked with a red asterisk
 (*) require a response.
- □ Important instructions for completing the Reportable New Information page (*Note:* these instructions are <u>only</u> to be used when submitting an Administrative Check In):
 - **Q1** Type in "Administrative Check In" and the current year
 - **Q2** Select the current date
 - **Q3** Check Administrative Check In

Creating New: IRB Submission

Reportable

Information

Go to forms n

Reportable New Information

- 1. RNI short title: (uniquely identify this new information report) Administrative Check In 20XX
- 2. * Date you became aware of the information:
- 3. Identify the categories that represent the new information: (check all that apply)

Ē

	Name	Description
		Information that indicates a new or increased risk, or a safety issue. For example:
		a. New information (e.g., an interim analysis, safety monitoring report, publication in the literature, sponsor report, or investigator finding) indicates an increase in the frequency or magnitude of a previously known risk, or uncovers a new risk.
		b. An investigator brochure, package insert, or device labeling is revised to indicate an increase in the frequency or magnitude of a previously known risk, or to describe a new risk.
	🗆 Risk	 C. Withdrawal, restriction, or modification of a marketed approval of a drug, device, or biologic used in a research protocol.
,	Administrative	Poriodic roport for studios that do not roquiro
	Check-in Report	renouic report for studies that do not require (



Reportable New Information continued

- □ Important instructions for completing the Reportable New Information page (*Note:* these instructions are <u>only</u> to be used when submitting an Administrative Check In):
 - **Q4** Type in "Administrative Check In" and the current year
 - **Q5** Provide answer
 - **Q6** Provide answer*
 - **Q7** Provide answer*

*If you select **Yes** to Questions 6 or 7, please submit a Modification to the IRB.





Reportable New Information continued

- □ Important instructions for completing the Reportable New Information page (*Note:* these instructions are <u>only</u> to be used when submitting an Administrative Check In):
 - **Q8** The study you submitted the form under automatically populates
 - **Q9** Attach the Administrative Check In form
 - 1. The Administrative Check In template can be found <u>here</u>.
 - 2. Complete the template and attach it to the Huron submission.
 - 3. Instructions on how to attach the Administrative Check In template to the submission can be found on the next slide.

Related studies and	modificatio	ons: 😮				
ID	Short Title	Investigator	State	IRB Office)	
STUDY2023-0012	Test	Denise Puga	Approved	IRB 1	Ø	
+ Add						
There are no items to	display					
			8	Exit	Save	Continue 🔿



How to attach the Administrative Check In

- 1. Click **+Add** in Question 9 of the Reportable New Information page
- 2. Click **Choose File** to locate the desired document from your desktop
- 3. Click OK

The Administrative Check In template can be located on the HRPP website: <u>https://vpr.tamu.edu/human-</u><u>research-protection-program/toolkit/templates/</u>





Submitting the Reportable New Information to the IRB

- 1. Click **Continue** to be directed to the **Final Page**
- 2. From the Final Page, click **Finish**
- 3. From the study workspace, click Submit RNI
- 4. Click **OK**

1	Save Continue
2	Exit Save Finish
	Edit RNI
	Printer Version
8	A Submit RNI
	Manage Ancillary Reviews
	Manage Editors
	Add Related Submission
	Copy Submission
	O Discard