



HOW TO UPLOAD CONTINUING REVIEW APPROVALS IN IREx

IREx is used to document and communicate the sIRB Continuing Review approval for participating sites. IREx does not capture the information the Reviewing IRB needs to conduct the Continuing Review, so this information must be gathered and submitted to the sIRB outside of IREx. Once approved, upload the Continuing Review approvals in IREx by following these three steps:

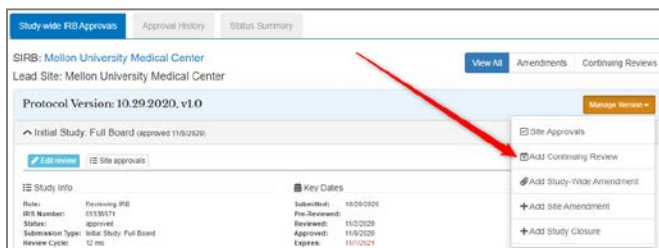


Note: Steps B and C are separate because the newly stamped consents must be uploaded for each site. However, IREx auto-fills the relevant review dates and determination letter for sites, based on those entered for the Lead Site/Overall Study.

STEP A: ADD THE CONTINUING REVIEW IN IREx

The Reviewing IRB Liaison or IREx Study Manager (if permitted by the Reviewing IRB) must first add the Continuing Review for the study and upload the Continuing Review approval for the Lead Site or Overall study. Then, Continuing Review approvals can be uploaded for relying sites (see [Step C](#)).

1. On the Study-wide IRB Approvals tab, click on the orange **Manage Version** button and select **Add Continuing Review**.
2. In the Add Continuing Review dialog, indicate whether the **Continuing review also contained a study-wide amendment approved on the same day** – Yes/No. If **Yes**, provide the required information about whether the protocol was changed and a change summary.



Continuing review also contained a study-wide amendment approved on the same day? Yes No

Which documents were *changed or removed* by this Continuing Review?

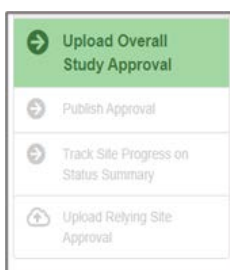
- Investigators Brochure: [INVESTIGATOR'S BROCHURE_1_0.docx](#) changed / removed?
- Package Insert - Package insert: [PackageInsert.docx](#) changed / removed?
- Others - Recruitment: [Recruitment.docx](#) changed / removed?
- Others - Study materials: [Questionnaire.docx](#) changed / removed?
- Others - Survey: [Survey.docx](#) changed / removed?

3. Select the documents that were changed or removed by this Continuing Review. These documents will remain in archived versions. Delete any study documents that are no longer part of the currently approved set of documents. Previously approved consents will be automatically removed. You can upload the newly stamped consents and other updated documents in the steps ahead.

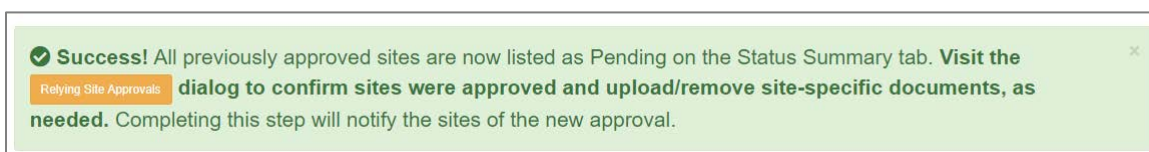
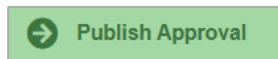
4. Click **Save** to add the Continuing Review. This will add steps to your GETTING STARTED checklist where you finish uploading the approval for the overall study.
Tip: On the Status Summary tab, **sites' approval status will change to "Approval Pending"** until their Continuing Review approval is uploaded in Step C.

STEP B: UPLOAD SIRB APPROVAL FOR THE LEAD SITE/OVERALL STUDY

- Click the **Upload Overall Study Approval** step on the GETTING STARTED checklist to upload the Lead Site/Overall approval.
- In the dialog under Study Information:
 - Set the Status to **Approved**.
 - Confirm **Type of Study** risk level.
 - Ensure the correct **Review Type** (*Expedited or Full Board*) is selected.
 - Enter the **Review Cycle**.
 - Is Site Enrolling?** will default to the most recent enrollment status for the lead site. If the lead site is not enrolling, nor plans to, you will not be required to upload a Consent & Assent.
 - Click Continue.
- Enter the Key Review dates when the Continuing Review was **Submitted, Pre-Review was Completed, Reviewed, and Approved** and click **Continue**.
- Upload the new **IRB Approval Documentation**, the **Continuing Review Application**, the newly stamped **Consent & Assents** and any other new or updated documents for the lead site only. Required documents will be marked in red.
- Review the study information and click **Save**. If required fields are missing, the section of the dialog needing attention will be highlighted.
- To make the documents visible to relying sites, click **Publish Approval** on the GETTING STARTED checklist.
- After the Lead Site/Overall approval is published, a banner will appear at the top of the study page confirming you have added the Continuing Review correctly and the final step is to upload Relying Site approvals.



Type	Document	Date Approved
Protocol [05.02.2021, v3.0]	Protocol Document_v3.0_Amend 5_05.02.2021.docx	5/12/2021

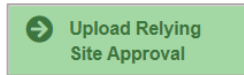
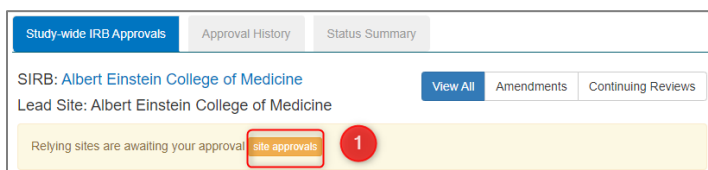


Tip: Relying sites are not notified of the new Continuing Review approval – they are notified of approval when their site-specific approval documents are uploaded in Step C.

STEP C: UPLOAD APPROVAL FOR RELYING SITES

AFTER the Continuing Review Approval has been uploaded for the Lead Site or Overall study, sites' approvals can be added.

- Click on the **site approvals** button on the Study-wide IRB Approvals tab (see **1** in the screenshot). Study Managers can also click the **Uploading Relying Site Approvals** step on their GETTING STARTED checklist.
- In the Relying Site Approvals dialog, select the **name of the site** for which you are uploading approval. (All sites who have ceded review appear, but only sites approved on the previous version will have a review type of Continuing Review: Full/Expedited.)



3. Change the Status to **approved** and
 - o Ensure the correct **Review Type** is selected.
 - o **Is Site Enrolling?** will default to the most recent enrollment status for the site. If the site is not enrolling, nor plans to, you will not be required to upload a Consent & Assent document.
 - o The dates and determination letter from the overall study approval will be auto-populated for all sites that had approval on the previous version.
4. Other **Site Specific Documents** that were approved in the previous version will also be carried forward. Please verify that these documents are still part of the approved set of documents.
5. Upload the new consent forms (or indicate a waiver was approved) and ensure the correct versions of all other approved documents are listed for the site.
6. Click **Save**. IREx will notify the (1) Reviewing IRB Liaisons, (2) site Liaison(s), (3) the IREX Study Manager(s), and (4) site study contacts in IREx of the new approval. The documents will appear on the Site-specific IRB Approvals tab.

ADDITIONAL TIPS:

- You can upload approvals for more than one site at a time by selecting another site name (item 2 above) and completing items 3-5 for each site before saving.
- To save information without notifying a site, change the approval status to *pending* or *leave blank*. You can return later to complete the approval.

The screenshot shows the 'Relying Site Approvals' interface for 'Carnegie Univ'. The form is titled 'Carnegie Univ' and includes a sidebar with site names: Carnegie Univ (selected), Middle-earth College, and Midwest Univ Med Ctr. The main form area contains the following fields and options:

- Status:** approved (marked with red callout 3)
- Review Type:** Continuing Review: Full Board
- Is Site Enrolling?:** Yes (selected)
- Date Submitted:** 02/01/2022
- Date Pre-Reviewed:** mm/dd/yyyy
- Date Reviewed:** 02/18/2022
- Date Approved:** 02/24/2022
- Site Specific Documents:** Consents & Assents (waived), Measures, Recruitment & Advertisements, Additional IRB Approved Documents, Other Documents.

Red callouts 1-6 highlight key elements: 1. Reviewing IRB Liaisons, 2. Site Liaison(s), 3. IREX Study Manager(s), 4. Site study contacts, 5. Upload any newly stamped consents, and 6. Save button. A note states 'Automatically carried over from lead site/overall approval'.