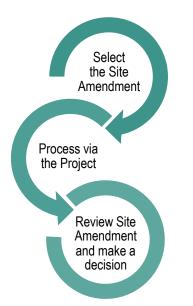
PÊGIS



FOR RESEARCH OFFICES

Processing a Site Amendment

The following diagram is an overview of processing a Site Amendment in REGIS.



Note: This QRG assumes you have already set up your homepage tiles. See <u>QRG – Research Offices – Setting up a</u> <u>Homepage</u> and REGIS User Manual guide – <u>Homepage tiles</u>

Understanding the types of Site Amendments and how they act in the system

When Ethics is in REGIS, the following approved amendments will be **automatically shared** with the Research Governance Office:

 Change to CPI – no RGO action is required. Contact details are automatically updated.

- Request for an Extension of HREC – no RGO action is required. Expiry date is automatically updated
- Change to PI RGO review is required
- Addition of a New Site (Automatically creates a NSW Site Application (STE) for completion)

The following Site Amendments are submitted directly to the RGO through the Site more information form in REGIS. Ethics approval is required before the RGO can finalise;

Site General Amendment

The following Site Amendments are submitted directly to the RGO via REGIS and do not need Ethics approval.

- Changing Site Investigators and Administration Contact
- Contract Changes

Processing a Site Amendment Step 1

Select the site amendment for review and processing from the Decision homepage tile.

Step 2

You will be taken to 'Decisions' in the Site Project (post-approval).

Step 3

Download the form and attachments using the download icon. A temporary .zip file will download.



Review the amendment form and documents, as per your local Research Office policy.

For Site General Amendments

When Ethics is in REGIS, the ethics amendment form ID has been included on the Site General Amendment form.

Ethics Form ID	52500
Note to Research Office: Locate the related Ethics approved documents by typing form ID to sea documents section.	rch filter of the Related

The Research Office can search the Site Project's 'Related Documents' using the Ethics Form ID i.e. 52500, for the ethics approved amendment documents. See below example.

Related documents				
📥 Bulk download				
Filters				
Search				
062454				
Document types				
Please select				
Only modified in latest submission No Q Search X Reset				
Found 4 items				
2023/ETH00078:	062454_2023_ETH0007	24.0 KB		
 Notification of an am 	062454_Notification of	88.3 KB		
Participant Informati	062454_Participant Info	11.7 KB		
 Summary Application 	062454_Summary Appli	11.9 KB		

Step 4

*

Select 'Edit' at the bottom of the screen and click the 'Edit' icon on the decision row.

For Site General Amendments

Use the 'Select Documents' icon to choose the documents related to that amendment. Most recently submitted documents are identifiable by the * next to the System Version number. You can also use the Form ID which appears in the 'Filename' i.e. 030047

Title	🖷 R3447 - Arm-001 - Australasia-1-22-09-2020
Filename	279998-030047_CTRA v1.docx
System Version	1*
Date modified	24/09/2020 10:44 AM
Туре	Clinical Trial Agreement - Medicines Australia (CTRA)

Click the tick at the bottom of the page and click 'Save' on the right-hand side of the screen.

Step 5

In the 'User' section, a list of users will prefill if your Research Office has Research Governance Officers listed in the system. If your name appears progress to step 6.

If there are no users you will need to manually add yourself. Click the three dots at the end of the 'User' cell to search for the person who will or has made the decision.

Search for the user by typing in their first name, last name or email address.

Click on the checkbox next to the user and then on the 'Add user/s' tick icon at the bottom of the panel. This is the person who has been delegated authority to undertake review of Governance business i.e. Research Governance Officer (RGO).

Click 'Save' within the wizard.

Step 6

Select the 'View user decision' icon next to the form you are processing to access the user decision.

Step 7

Click on the available user decision and the management panel will appear.



Step 8

Click on 'Choose' to select the decision:

2

- Accepted
- Rejected
- More information required

When 'More information required' is selected, add a reason in the 'Comments' section and it will pull into the email template. Once comments have been added, click 'Save'.

Step 9

Click on the 'Send email' icon which has generated, triggering the email wizard to appear. The email template will load based off your decision, edit as required.



You can reload the email

template by clicking the 'Reload email template' button. This will revert the email back to the original template, removing any information manually added. If you navigate away from the page, any information added manually will save.

Click 'send'. The rext to the 'Type' of amendment indicates that an email has been sent.

RGO Review of a More Information Required Form

Step 1

Select the More Information Required -RGO Approval form for review and processing from the Decision homepage tile.

Step 2

You will be taken to 'Decisions' in the Site Project (post-approval).

Step 3

Download the form and attachments. A temporary .zip file will download.

Review the More Information Required – RGO Approval form and documents, as per your local Research Office policy.

Follow **steps 4-9** of Processing a Site Amendment.