



# 11.4 Consent in Study Runner

After the **Consent module** has been activated for your Study and at least one eConsent Form has been configured, you can use **Study Runner** to:

- Monitor Participant consent statuses
- Schedule eConsent Events to make consent Forms available to Participants

You can view eConsent statuses in two places:

- **Participant Matrix** - A separate column is displayed for each eConsent Form in the Study.
- **Participant Details Page** - Status appears in the **General Information** section and on the Form cards in the **Visits** section.

### General Information

Edit

Participant ID	001	Status	Available
Study Name	RGB22	Site Name	a
eConsent Status	Consent Signed		

### Visits

Sort by Date

<b>Consent</b>	30-Mar-2023		<b>eConsent</b>	
				30-Mar-23 by...

The following statuses may appear for eConsent Forms:

**Icon**      **Status**      **Description**



Consent Not Signed

The Participant has not yet signed the eConsent Form.



Consent Signed

The Participant has signed the eConsent Form, but it has not yet been countersigned by a site user (CRC or Investigator).



**Consent**      The Participant has signed the eConsent Form, and it has been Countersigned countersigned by a site user (CRC or Investigator).



**Requires Reconsent**

The Participant's previous consent was removed. This typically occurs when a new version of the eConsent Form is issued and a Data Manager has migrated the record to this new version, or if a site user (CRC or Investigator) has manually unconsented the form due to a process error. The Participant must provide consent again. **Note:** When an eConsent Form is set to *Requires Reconsent*, it is marked as *Data Entry Started* and reappears on the Participant's dashboard so the Participant can re-sign.

## Scheduling and Managing eConsent Forms

### Schedule an eConsent Form

For a Participant to see an eConsent Form on their dashboard, you must schedule the [Event](#) containing the Form.

Schedule the Event the same way you schedule any other Visit Event:

- From the **Participant Matrix**, or
- By clicking **Add New** on the **Participant Details** page.

For more information on scheduling events, refer to [Schedule an Event](#).

### Invite the Participant

To access the eConsent Form, the Participant must also be invited to the Study.

Once the Visit with the eConsent Form is scheduled **and** the Participant has been invited:

- The Participant can log in to their dashboard.
- The Participant can sign the eConsent Form.
- After signing, the Form is marked as **Completed**, and the eConsent status is updated in **Study Runner**.

For more information on inviting participants, refer to [Invite Participants](#) and [Automating Participant Access](#).

### View an eConsent Form

After the Event with an eConsent Form has been scheduled, the Form can be viewed by any user with permission.

- **Participant users** are the only ones who can check the box to sign the Form. It is read only for all other users.
- Other users (e.g., CRCs, Investigators, Data Managers) can open the Form but cannot sign on behalf of the Participant.
- To open the Form:
  - Click the Visit Form card, or
  - Use the actions menu.

When viewing the Form, the top of the page displays signature information:

- When the Participant signed
- The Participant's name (visible only to CRCs and Investigators)
- Who countersigned and when the countersignature was added

□ **Note:** A Form is not visible to the Participant until both conditions are met:

1. The Visit Event containing the eConsent Form is scheduled.
2. The Participant has been invited to the Study.

This form was signed by  on  under the following attestation: "I confirm that this participant has freely and voluntarily provided informed consent. I intend for this electronic signature to be the legally binding equivalent of my written signature."

This form was signed by  on  under the following attestation: "I freely and voluntarily consent to participate in this study. I intend for this electronic signature to be the legally binding equivalent of my written signature."

### 002: Informed Consent

Consent text from the form

I agree

You're in read-only mode.

## Countersign a Signed Consent

After a Participant signs an eConsent Form, its status is updated on both the **Participant Matrix** and the **Participant Details** page.

Clinical Research Coordinators (CRCs) and Investigators can then countersign.

To countersign:

1. On the signed eConsent Form card, open the **Actions** menu.
2. Select **Countersign**.
3. Enter your **Username** and **Password** to confirm.
4. Click **Submit**.

The eConsent status on the Form card updates to **Consent Countersigned**.

## Sign Form eConsent for Participant 003

Enter your user name and password below to signify agreement with the following statement:

I confirm that this participant has freely and voluntarily provided informed consent. I intend for this electronic signature to be the legally binding equivalent of my written signature.

**User Full Name: Riley Bianchi-CRC**

**Date/Time: 03-Apr-2023**

**(The exact date and time will be recorded by the system upon submission of the signature form.)**

**Role: Clinical Research Coordinator**

User Name :

Password :

## Unconsent a Signed Consent

You can also remove consent when a Participant must re-sign (for example, if the participant made a mistake completing the form).

To unconsent:

1. On the signed eConsent Form card, open the **Actions** menu.
2. Select **Unconsent**.
3. In the **Confirm Signature Status** window, review the message.
  - Performing this action will mark the Form as *Requires Reconsent*.
  - The Form will reappear on the Participant's dashboard for re-signing.
4. Click **Confirm**.

The Form's eConsent status updates to **Requires Reconsent** and the Form status changes from **Completed** to **Data Entry Started**.

### Confirm Signature Status

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Performing this action will mark this form as requiring reconsent. This will cause the form to appear on the participant's dashboard again and allow them to resign it. Please confirm this is appropriate for this consent form.

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⚠ **Warning:** Once you unconsent a Form, the action cannot be reversed. The Participant must re-sign to restore consent.

□ **Note:** The eConsent status in the **General Information** section of the **Participant Details** page is updated when the page is refreshed.

## Migrating an eConsent Form to a New Version

When you migrate a Consent Form to a new version, OpenClinica updates the status and clears outdated data to maintain integrity and compliance.

### Behavior during migration

- The Consent signature status is set to **Requires Reconsent**.
- The Form is cleared, removing all data except Participant contact fields.
- The Form status is reset to **Not Started**.
- You must enter a **Reason for Change (RFC)** to complete the migration.

### Audit Log

- Both the *Migration* and *Clear* actions are recorded in the Audit Log.
- The RFC is stored with these records.
- For more details, see [Participant Audit Log](#).

□ **Warning:** Migrating to a new version permanently removes previously signed consent data. Participants must re-sign the updated Form.

## Reassign

Reassigning a Participant transfers them to another site. After reassignment:

- The Participant is associated with the new site.
- Reporting, permissions, and workflow are updated.
- Existing Participant data remains intact.

## Clear

Clearing a Form removes all active data. When you clear a Form:

- The Form status resets to **Not Started**.
- All associated queries are closed.
- Audit history is retained, but active data is permanently removed.

Approved for publication by Kate Lambert. Signed on 2025-09-24 2:24PM

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