Oracle® Life Sciences Clinical One Platform

Notifications and Permissions Guide

Release 24.1
F91496-03
May 2024

Oracle Life Sciences Clinical One Platform Notifications and Permissions Guide, Release 24.1

F91496-03

Primary Author: Oracle Life Sciences Documentation Team

Contents

Preface

This preface contains the following sections:

Documentation accessibility

For information about Oracle's commitment to accessibility, visit the Oracle Accessibility Program website at http://www.oracle.com/pls/topic/lookup?ctx=acc&id=docacc.

Diversity and Inclusion

Oracle is fully committed to diversity and inclusion. Oracle respects and values having a diverse workforce that increases thought leadership and innovation. As part of our initiative to build a more inclusive culture that positively impacts our employees, customers, and partners, we are working to remove insensitive terms from our products and documentation. We are also mindful of the necessity to maintain compatibility with our customers' existing technologies and the need to ensure continuity of service as Oracle's offerings and industry standards evolve. Because of these technical constraints, our effort to remove insensitive terms is ongoing and will take time and external cooperation.

Related resources

All documentation and other supporting materials are available on the Oracle Help Center.

Access to Oracle Support

Oracle customers that have purchased support have access to electronic support through Support Cloud.

Contact our Oracle Customer Support Services team by logging requests in one of the following locations:

- English interface Customer Support Portal (https://hsgbu.custhelp.com/)
- Japanese interface Customer Support Portal (https://hsgbu-jp.custhelp.com/)

You can also call our 24x7 help desk. For information, visit https://www.oracle.com/life-sciences/support/ or visit http://www.oracle.com/pls/topic/lookup?ctx=acc&id=trs if you are hearing impaired.

Additional copyright information

This documentation may include references to materials, offerings, or products that were previously offered by Phase Forward Inc. Certain materials, offerings, services, or products may no longer be offered or provided. Oracle and its affiliates cannot be held responsible for any such references should they appear in the text provided.

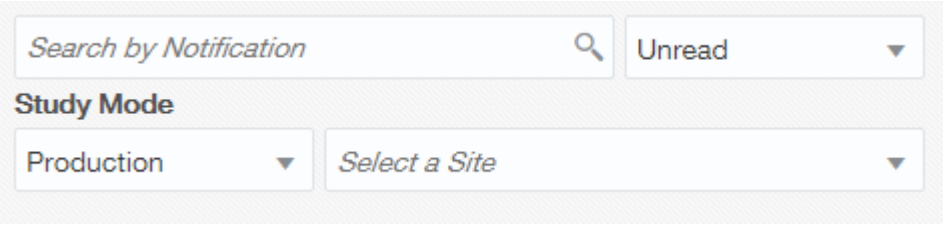
1

View notifications

The notifications that you receive depend on your role during each mode of the study. If you're not getting the notifications that you expect, contact your system administrator. Notifications are also sent to your email.

Show me how!

1. On the Home page, click the notification button ( ) on a study.
2. In the upper right, from the **Study Mode** drop-down, select the mode that you want to view notifications for.
3. Filter your view as needed:
   - To view notifications about specific sites, choose a site from the **Select a Site** drop-down.
     If the Select a Site drop-down is disabled, remove the search terms from the Search by Notification field.
   - To find notifications by name, enter the notification name in the **Search by Notification** field.
     If the Search by Notification field is disabled, remove any filters for sites.
   - To sort notifications, select an option from the drop-down to the right of the Search by Notification field.



4. Click a notification to view its details. For descriptions of the notifications and details about who receives them, see Notifications.

> **Tip:**
>
> A blue circle appears to the left of an unread notification.

2
Permissions

Permissions for report notifications

View the descriptions and required permissions for report notifications.

Notification	Permission required
Report failed to generate notification	None Any role can receive this notification.
Report is ready notification	None Any role can receive this notification.
Scheduled report notifications	*Schedule Reports to Run* This permission is for Sponsor users and should only be granted to one Sponsor level user.

Permissions for rule notifications

View the descriptions and required permissions for rule notifications.

Notification	Permission required
Rule Re-run notification	*Re-run Rules* This permission is for sponsor users only.
Rules Validation Complete notification	*Design Custom Rules* This permission is for sponsor users only.
Rule Lock notification	*Receive Rule Failure Notification for Locked Data* This permission is for sponsor users only.

Permissions for subject notifications

View the descriptions and required permissions for subject notifications.

Notification	Permission required
Code Break notification	*Receive the Code Break Notification* This permission is for both sponsor and site users.
• First subject screened in study notification • First subject screened at site notification • First subject randomized in study notification • First subject randomized at site notification	*Study Limits Notifications* This permission is for both sponsor and site users.
Screening update notification	*Receive the Subject Screening Notification* This permission is for both sponsor and site users.
Signature requests notification	*Receive the Pending Signatures Notification* This permission is for both sponsor and site users.
Subject completion notification	*Receive the Subject Completion Notification* This is a permission for sponsor users.
Subject dispensation notification (with failures)	• *Receive the Unblinded Dispensation Notification* • *Receive the Dispensation Notification* • *Receive the Dispensation with Dosing Instructions Notification* These permissions are for both sponsor and site users. *These permissions can unblind users. Use with caution.*
Subject Number Update notification	*Receive the Subject Number Update Notification* This permission is for both sponsor and site users.
Subject randomization notification (with failures)	• *Receive the Randomization Notification* • *Receive the Unblinded Randomization Notification* These permissions are for both sponsor and site users. *These permissions can unblind users. Use with caution.*
Subject rollover notification	*Receive the Subject Rollover Notification* This is a permission for sponsor users.
Subject screening notification (with error)	*Receive the Subject Screening Notification* This permission is for both sponsor and site users.
Subject transferred notification	*Receive the Subject Transferred Notification* This permission is for both sponsor and site users.
Subject unscheduled visit notification	*Receive the Unscheduled Visit Notification* This is a permission for sponsor users.
Subject visit notification (with failures)	*Receive the Subject Visit Notification* This is a permission for sponsor users.
Subject withdrawal notification	*Receive the Subject Visit Notification* This is a permission for sponsor users.
Training complete notification	None Any role that is assigned training will receive this permission.

Notification	Permission required
Unblinded Pharmacist dispensation notification	*Receive the Unblinded Pharamcist Dispensation Notification* This permission is for both sponsor and site users. *This permission can unblind users. Use with caution.*
User Upload notifications	*Upload Users in Bulk* This permission is for Sponsor users only.

Permissions for supply notifications

View the descriptions and required permissions for supply notifications.

Notification	Permission required
Depot shipment notification	*Receive Notification of Depot Shipments* This permission is for both sponsor and site users. *This permission can unblind users. Use with caution.*
Low kit depot (Unblinded) notification	*Receive Notification of Depot Shipments* This permission is for both sponsor and site users. *This permission can unblind users. Use with caution.*
Kit List Deactivated notification (with failures)	*Upload and Generate Inventory Lists* This permission is for both sponsor and site users.
Kit missing from shipment for destruction notification	*Receive Notification of Shipments* This permission is for both sponsor and site users.
Kit status changed to "Misallocated" notification	*Receive the Unblinded Kit Misallocation Notification* This permission is for both sponsor and site users. *This permission can unblind users. Use with caution.*
Kit status changed to "Not Dispensed to Subject" notification	• *Receive the Not Dispensed to Subject Notification* • *Receive the Not Dispensed to Subject (Unblinded) Notification* • *Receive the Not Dispensed to Subject (Unblinded Pharmacist) Notification* These permissions are for both sponsor and site users. *These permissions can unblind users. Use with caution.*
Kits Quarantined notification	• *Receive the Quarantined Depot Shipment Notification* • *Receive the Quarantined Site Shipment Notification* These permissions are for both sponsor and site users. *These permissions can unblind users. Use with caution.*

Notification	Permission required
Kits Released from Quarantine notification	• *Receive the Released from Quarantine Notification (Depot)* • *Receive the Released from Quarantine Notification (Site)* These permissions are for both sponsor and site users. *These permissions can unblind users. Use with caution.*
Pending shipment notification	*Receive Notification of Shipments* This permission is for both sponsor and site users.
Randomization update notification	*Receive the Randomization Notification* This permission is for both sponsor and site users.
Shipment cancellation notification	*Receive Notification of Shipments* This permission is for both sponsor and site users.
Shipment failure notification	*Receive Notification of Shipments* This permission is for both sponsor and site users.
Shipment for destruction created notification	*Receive Notification of Shipments* This permission is for both sponsor and site users.
Shipment for destruction not received notification	*Receive Notification of Shipments* This permission is for both sponsor and site users.
Shipment Quarantined notification	• *Receive the Quarantined Depot Shipment Notification* • *Receive the Quarantined Site Shipment Notification* These permissions are for both sponsor and site users.
Shipment Released from Quarantine notification	• *Receive the Released from Quarantine Shipment Notification (Depot)* • *Receive the Released from Quarantine Notification (Site)* These permissions are for both sponsor and site users.
Shipment request notification	*Receive Notification of Shipments* This permission is for both sponsor and site users.
State of a site has changed notification	*Receive Site has been Updated Notification* This permission is for both sponsor and site users.
Study inventory notification	*Receive the Quarantined Depot Shipment Notification* This permission is for both sponsor and site users. *This permission can unblind users. Use with caution.*
Unblinded dose hold performed notification	*Receive the Unblinded Dose Hold Notification* This permission is for both sponsor and site users. *This permission can unblind users. Use with caution.*

3

Notifications

Oracle Clinical One Platform notifications

Report notifications

Report failed to generate notification

This notification is sent when a report doesn't generate correctly.

Permissions required to receive the notification

Any role can receive this notification without assigning it a specific permission. You receive the notification if a report you tried to run didn't generate.

Field descriptions

The table below describes all fields and details in the **Report failed to generate notification**.

> **Note:**
>
> This notification also contains an introductory statement indicating that your report has failed to generate in the application.

Field	Description
Title	The title is comprised of two distinct elements: • The study's ID. For example, a study's ID can be P01-123-A3. • A statement about the report failing to generate.
File Name	Indicates the parameters chosen for the report to run.
Study Name	Indicates a study's ID, as specified by the global study creator when they created the study.
Number of Records	Indicates the number of data records in the report.
Server Run Date	Indicates the exact date and time when the report failed to generate. For example: 10-Nov-2022 08:07 AM.
Report Run ID	Indicates the GUID of the report in the application's backend system. For example: 08B6771F54DE45A0BB576E86326BE84D.

Report is ready notification

This notification is sent when a report is ready.

Permissions required to receive this notification

Any role can receive this notification without assigning it a specific permission. You receive the notification after a report that you run is ready.

Field descriptions

The table below describes all fields and details in the **Report is ready notification**.

> **Note:**
>
> This notification also contains an introductory statement indicating that your report is now ready in the application. For example: *Subject Events Report is now ready in Clinical One. You can log on to Clinical One Portal to download the report*.

Field	Description
Title	The title is comprised of two distinct elements: • The study's ID. For example, a study's ID can be P01-123-A3. • A statement about the report being ready. For example: Subject Events Report is ready in Clinical One.
File Name	Indicates the parameters chosen for the report to run.
Site(s) selection	Indicates where the data of the report originated. If all sites of a study have been selected, "All sites" will be displayed. Otherwise, the sites will be listed, separated by commas. Only appears in the Subject Data Extract notification.

Field	Description
Study Name	Indicates a study's ID again, as specified by the global study creator when they created the study.
Number of records	Indicates the number of data records in the generated report.
Server Run Date	Indicates the exact date and time when the report was generated. For example: 10-Nov-2022 08:07 AM.
Report Run ID	Indicates the GUID of the report generation in the application's backend system. For example: 08B6771F54DE45A0BB576E86326BE84D.
File Type	Indicates the format your report has been generated in, such as CSV, CPORT, XPORT, or SAS7BDAT. Only appears in the Subject Data Extract notification.

Scheduled report notifications

These notifications are sent to users whose email address is included in the report schedule configuration.

Permissions required to receive these notifications

Users included in the report schedule configuration, and who have permissions for the report being scheduled can receive the following notifications:

- **Scheduled Report Failure Notification**
- **Scheduled Report sFTP Failure Notification**
- **Scheduled Report sFTP Success Notification**
- **Report Scheduled Created Notification**
- **Report Scheduled Activated Notification**
- **Report Scheduled Deactivated Notification**

Field descriptions

The table below describes all fields and details in the **Report failed to generate notification**.

> **Note:**
>
> sFTP notifications include additional information on sFTP location.
> This notification also contains an introductory statement indicating that your report is now ready in the application. For example: *Scheduled report has run in Clinical One and is successfully uploaded to sFTP.*

Field	Description
Title	The title is comprised of two distinct elements: • The study's ID. For example, a study's ID can be P01-123-A3. • The report name. • The title of the notification, for example, Scheduled Report sFTP Success Notification.
Study Name	Indicates a study's ID again, as specified by the global study creator when they created the study.
Study Mode	Indicates the study mode associated with this notification. Displayed values can be Test, Active, or Training.
Report Name	Indicates the name of the scheduled report.
Upload Date	Indicates the date the report was uploaded.

Rule notifications

- Rule Re-run notification
  This notification is sent when a rule re-run is completed.
- Rules Validation Complete notification
  This notification is sent once the validation of custom rules is complete, after a study has been copied and moved to testing mode, and when a form has been copied.
- Rule Lock notification
  This notification is sent when a rule is unable to update a target due to it being locked.

Rule Re-run notification

This notification is sent when a rule re-run is completed.

Permissions required to receive the notification

Assign the *Re-run rules* permission to anyone who wants to receive this notification.

Field descriptions

The table below describes all fields and details in the **Rule re-run notification**.

> **Note:**
>
> This notification includes details on rules that failed during execution when applicable.

Field	Description
Title	Indicates the title of the notification, for example, Rule re-run is completed.
Jobs group ID	The internal ID name of the job created when a rule is re-run.

Rules Validation Complete notification

This notification is sent once the validation of custom rules is complete, after a study has been copied and moved to testing mode, and when a form has been copied.

Permissions required to receive this notification

Anyone who is assigned the *Design Custom Rules* permission will receive a **Rules Validation Complete notification**. Rules designers will receive this notification by default when the process of validating custom rules is finished.

Field descriptions

The table below describes all fields and details in the **Rules validation complete notification**.

> **Note:**
>
> This notification includes a statement suggesting to run the Rules report for more information on invalid rules below the following fields.

Field	Description
Title	The title is comprised of two distinct elements: • The study's ID. For example, a study's ID can be P01-123-A3. • The title of the notification, for example, Rules Validation Complete.
Total Count of Copied Rules	Indicates the total number of copied rules.
Total Validated Rules	Indicates the total number of validated rules.
Validation Error Count	Indicates the number of rules that encountered an error during validation.
Date/Time Completed	Indicates the exact date and time of copying or rule validation. For example: 10-Nov-2022 08:07 AM.

Rule Lock notification

This notification is sent when a rule is unable to update a target due to it being locked.

Permissions required to receive the notification

Sponsor users assigned the *Receive Rule Failure Notification for Locked Data* will receive this notification.

Field descriptions

The table below describes all fields and details in the **Rule lock notification**.

Field	Description
Title	The title of the notification, for example, Rule Lock Notification.
Study Name	Indicates a study's ID again, as specified by the global study creator when they created the study.

Field	Description
Study Mode	Indicates the study mode associated with this notification. Displayed values can be Test, Active, or Training.
Site Name	Indicates the name of the site as specified by a site manager when they created the site.
Subject Number	Indicates the number of the subject.
Visit Branch	Indicates the name of the visit branch.
Visit	Indicates the name of the visit.
Form	Indicates the name of the form containing the locked target.
Form Sequence Number	Indicates the form sequence number for a repeating form.
Form Section Number	Indicates the form section number for a two-section form.
Question	Indicates the title of the question.
Date & Time of Notification	Indicates the exact date and time of the notification. For example: 10-Nov-2022 08:07 AM.

Subject notifications

- Code Break notification
  This notification is sent when a subject is unblinded due to a code break. The notification does not include unblinded data.
- First subject screened in a study notification
  This notification is sent when the first subject is screened or enrolled in the study.
- First subject screened at a site notification
  This notification is sent when the first subject is screened or enrolled in a site.
- First subject randomized at a site notification
  This notification is sent when the first subject is randomized at a site.
- First subject randomized in a study notification
  This notification is sent when the first subject is randomized in a study.
- Screening update notification
  This notification is sent when the screening limit or a percentage of the screening limit is reached for a study, site, or country.
- Signature requests notification
  This notification is sent when a required signature has a target date that has been reached or become overdue for a site you are assigned to.
- Subject completion notification
  This notification is sent when a subject completes a study. This happens when the site user marks the subject as complete and can only be done once the subject has completed all minimum required visits and all data has been entered.
- Subject dispensation notification (with failures)
  This notification is sent when a dispensation event is successfully completed or when the dispensation failed at a site you're assigned to. Additionally, this notification is sent when a dispensation event with dosing instructions is successfully completed or has failed. Details about partial dispensation are included when they occur.

- Subject randomization notification (with failures)
  This notification is sent when a subject is either successfully randomized or their randomization failed at a site that you're assigned to.
- Subject Number Update notification
  This notification is sent when a site or sponsor user has successfully updated a subject number, and contains the site number, the original subject number, and the new subject number.
- Subject rollover notification
  This notification is sent when a subject is enrolled into a rollover study.
- Subject screening notification (with error)
  This notification is sent when a subject is either successfully screened or their screening failed at a site that you're assigned to.
- Subject transferred notification
  This notification is sent when a subject transfers to another site.
- Subject unscheduled visit notification
  This notification is sent when a subject completes an unscheduled visit.
- Subject visit notification (with failures)
  This notification is sent when a subject completes a non-dispensation or optional visit at a site that you're assigned to. The notification is sent both when the subject successfully completes the visit and when one or more questions have errors.
- Subject withdrawal notification
  This notification is sent when a subject withdraws from a study.
- Training complete notification
  This notification is sent when you complete all training assigned to you.
- Unblinded Pharmacist dispensation notification
  Pharmacists or unblinded site users receive this notification when kits need to be dispensed by them to subjects during a dispensation or randomization visit.
- User Upload notifications
  These notifications are sent after the user upload process completes.

Code Break notification

This notification is sent when a subject is unblinded due to a code break. The notification does not include unblinded data.

Permissions required to receive the notification

Assign the *Receive the Code Break Notification* permission to anyone who wants to receive this notification. Users receive notifications for only the sites that they're assigned to.

Field descriptions

The table below describes all fields and details in the **Code Break notification**.

Field	Descriptions
Title	The title is comprised of three distinct elements: • The study's ID. For example, a study's ID can be P01-123-A3. • A subject's number. For example: MUHC22. • The title of the notification, for example, Code Break.

Field	Descriptions
Study ID	Indicates a study's ID again, as specified by the global study creator when they created the study.
Study Mode	Indicates the study mode associated with this notification. Displayed values can be Test, Active, or Training.
Site Name	Indicates the name of the site as specified by a site manager when they created the site.
Site ID	Indicates a site's ID as specified by the site manager who created the site.
Subject ID	Indicates the number of the subject who was unblinded due to a code break.
Date of Transaction (UTC)	Indicates the exact date and time of the code break. For example: 10-Nov-2022 08:07 AM.
Adverse Event	Indicates whether or not the subject experienced an adverse event.
Performed By	Indicates the username of the user who reported the code break.

First subject screened in a study notification

This notification is sent when the first subject is screened or enrolled in the study.

Permissions that receive the notification
To receive these notifications, assign the *Receive the Study Limits Notifications* permission to anyone who wants to receive this notification. Users receive notifications only for the study and sites that they are assigned to. Both site users and sponsors can receive this notification.

Table 3-1 Field descriptions

Field	Description
Subject line	Indicates the summary of the alert information. The subject would include a short description of the study name and the first subject screened.
Study ID	Indicates a study's ID, as specified by the global study creator when they created the study.
Study Mode	Indicates the study mode associated with this notification. Displayed values can be Test, Active, or Training.
Site Name	Indicates the name of the site where the unscheduled visit took place. The name is displayed as it was specified by a site manager when they created the site.
Site ID	Indicates a site's ID, as specified by the global study creator when they created the site.
Subject ID	Indicates the number of the subject who just completed the scheduled visit.
Date Performed (UTC)	Indicates the exact date and time of the completion of the screening visit. For example: 22-Jan-2023 07:09 AM.
Performed By	Indicates the user name of the user who performed the screening of the subject.

Related Topics

- Specify study, visit, limit, and cohort settings

First subject screened at a site notification

This notification is sent when the first subject is screened or enrolled in a site.

Permissions that receive the notification
To receive these notifications, assign the *Receive the Study Limits Notifications* permission to anyone who wants to receive this notification. Users receive notifications only for the sites that they are assigned to. Both site and sponsor users can receive this notification.

Table 3-2 Field descriptions

Field	Description
Study Name	Indicates the summary of the alert information. The subject would include a short description of the study name and the first subject screened.
Study ID	Indicates a study's ID, as specified by the global study creator when they created the study.
Study Mode	Indicates the study mode associated with this notification. Displayed values can be Test, Active, or Training.
Site Name	Indicates the name of the site where the unscheduled visit took place. The name is displayed as it was specified by a site manager when they created the site.
Site ID	Indicates a site's ID, as specified by the global study creator when they created the site.
Subject ID	Indicates the number of the first subject at a site who just completed the scheduled screening visit.
Date Performed (UTC)	Indicates the exact date and time of the completion of the screening visit. For example: 22-Jan-2023 07:09 AM.
Performed By	Indicates the user name of the user who performed the screening of the subject.

First subject randomized at a site notification

This notification is sent when the first subject is randomized at a site.

Permissions that receive the notification
Assign the *Receive the study limits notifications* permission to anyone who wants to receive this notification. Users receive notifications only for the sites that they are assigned to. Both site and sponsor users can receive this notification.

Only assigning the permissions to the role wouldn't suffice. Site enrollment toggle must be enabled and study managers must activate the **First subject randomized notification** checkbox for the site enrollment row in the study settings. See how to define enrollment settings in Specify study, enrollment, and visits settings.

Table 3-3 Field description

Field	Description
Study Name	Indicates the summary of the alert information. The subject would include a short description of the study name and the first subject screened.
Study ID	Indicates a study's ID, as specified by the global study creator when they created the study.
Study Mode	Indicates the study mode associated with this notification. Displayed values can be **Test**, **Active**, or **Training**.
Site Name	Indicates the name of the site where the unscheduled visit took place. The name is displayed as it was specified by a site manager when they created the site.
Site ID	Indicates a site's ID, as specified by the global study creator when they created the site.
Subject ID	Indicates the number of the first subject at a site who just completed the scheduled randomization visit.
Date Performed (UTC)	Indicates the exact date and time of the completion of the screening visit. For example: 22-Jan-2023 07:09 AM.
Performed By	Indicates the user name of the user who performed the screening of the subject.

First subject randomized in a study notification

This notification is sent when the first subject is randomized in a study.

Permissions that receive the notification
To receive these notifications, assign the *Receive the Study Limits Notifications* permission to anyone who wants to receive this notification. Users receive notifications only for the studies that they are assigned to. Both site and sponsor users can receive this notification.

Only assigning the permissions to the role wouldn't suffice. Site enrollment toggle must be enabled and study managers must activate the **First subject randomized notification** checkbox for the study enrollment row in the study settings. See how to define enrollment settings in Specify study, enrollment, and visits settings.

Table 3-4 Field description

Field	Description
Study Name	Indicates the summary of the alert information. The subject would include a short description of the study name and the first subject screened.
Study ID	Indicates a study's ID, as specified by the global study creator when they created the study.
Study Mode	Indicates the study mode associated with this notification. Displayed values can be **Test**, **Active**, or **Training**.

Table 3-4 (Cont.) Field description

Field	Description
Site Name	Indicates the name of the site where the unscheduled visit took place. The name is displayed as it was specified by a site manager when they created the site.
Site ID	Indicates a site's ID, as specified by the global study creator when they created the site.
Subject ID	Indicates the number of the subject who just completed the scheduled visit.
Date Performed (UTC)	Indicates the exact date and time of the completion of the screening visit. For example: 22-Jan-2023 07:09 AM.
Performed By	Indicates the user name of the user who performed the screening of the subject.

Screening update notification

This notification is sent when the screening limit or a percentage of the screening limit is reached for a study, site, or country.

Permissions that receive the notification

Assign the *Receive the Subject Screening Notification* permission to anyone who wants to receive this notification. Users receive notifications for only the sites that they're assigned to.

Field descriptions

The table below describes all fields and details in the **Screening update notification**.

Field	Description
Title	The title is comprised of three distinct elements: • The study's ID. For example, a study's ID can be P01-123-A3. • A subject's number. For example: MUHC22. • The title of the notification, for example, Screening Notification.
Study ID	Indicates a study's ID again, as specified by the global study creator when they created the study.
Study Mode	Indicates the study mode associated with this notification. Displayed values can be Test, Active, or Training.
Site Name	Indicates the name of the site as specified by a site manager when they created the site.
Subject Number	Indicates the number of the subject who just completed the visit.
Date Performed (UTC)	Indicates the exact date and time of the completion of the screening visit. For example: 10-Nov-2022 08:07 AM.
Performed By	Indicates the username of the user who performed the screening of the subject.
GUID	Indicates the notification's GUID.

Signature requests notification

This notification is sent when a required signature has a target date that has been reached or become overdue for a site you are assigned to.

Permissions required to receive the notification

Assign the *Receive the Pending Signatures Notification* permission to any roles that should receive this notification. Users receive notifications only for the sites that they're assigned to.

Field descriptions

The table below describes all fields and details in the **Signature requests notification**.

Field	Description
Title	The title of the notification, for example, Signature Requests, followed by a message to the recipient stating, "One or more Signature Requests are overdue."
Study Name	Indicates the study name, as specified by the global study creator when they created the study.
Study Mode	Indicates the study mode associated with this notification. Displayed values can be Test, Active, or Training.
Site Name	Indicates the name of the site as specified by a site manager when they created the site.

Subject completion notification

This notification is sent when a subject completes a study. This happens when the site user marks the subject as complete and can only be done once the subject has completed all minimum required visits and all data has been entered.

Permissions required to receive the notification

Assign the *Receive the Subject Completion Notification* permission to anyone who wants to receive this notification. Users receive notifications only for the sites that they're assigned to.

Field descriptions

The table below describes all fields and details in the **Subject completion notification**.

Field	Description
Title	The title is comprised of three distinct elements: • The study's ID. For example, a study's ID can be P01-123-A3. • A subject's number. For example: MUHC22. • The title of the notification, for example, Completion Notification.
Study Name	Indicates a study's ID again, as specified by the global study creator when they created the study.
Study Mode	Indicates the study mode associated with this notification. Displayed values can be Test, Active, or Training.

Field	Description
Site Name	Indicates the name of the site as specified by a site manager when they created the site.
Site ID	Indicates a site's ID as specified by the site manager who created the site.
Subject ID	Indicates the number of the subject who just completed the study.
Date Performed (UTC)	Indicates the exact date and time of the completion. For example: 10-Nov-2022 08:07 AM.
Performed By	Indicates the username of the user who updated the subject's status.

Subject dispensation notification (with failures)

This notification is sent when a dispensation event is successfully completed or when the dispensation failed at a site you're assigned to. Additionally, this notification is sent when a dispensation event with dosing instructions is successfully completed or has failed. Details about partial dispensation are included when they occur.

> **Note:**
>
> If Unblinded Pharmacist kits are assigned to a visit, this notification will also contain a notice for blinded site users to contact their pharmacist or unblinded site user to dispense those Unblinded Pharmacist kits.

Permissions required to receive the notification

Assign all or one of these permissions to anyone who wants to receive this notification:

- *Receive the Unblinded Dispensation Notification*: Includes a kit description for each kit number and the number of dose holds.
- *Receive the Dispensation Notification*: Includes the kit description only if the dispensed kits are of the Blinded distribution type. If not, this information is not included in the notification. The number of dose holds and any titration details are also included in the notification. If a titration does not have a blinded title, then these details are not present in the notification.
- *Receive the Dispensation with Dosing Instructions Notification*: Includes the kit description and dosing instructions, along with the number of dose holds.

Field descriptions

The table below describes all fields and details in the **Subject dispensation notification**.

> **Note:**
>
> Details on treatment arms are included in successful unblinded dispensation notifications.
> Information on kits is displayed in successful dispensation notifications.

Field	Description
Title	The title is comprised of three distinct elements: • The study's ID. For example, a study's ID can be P01-123-A3. • A subject's number. For example: MUHC22. • The title of the notification and whether the dispensation succeeded or not. For example, Dispensation succeeded or Dispensation failed.
Study Name	Indicates a study's ID again, as specified by the global study creator when they created the study.
Study mode	Indicates the study mode associated with this notification. Displayed values can be Test, Active, or Training.
Site Name	Indicates the name of the site where the unscheduled visit took place. The name is displayed as it was specified by a site manager when they created the site.
Site ID	Indicates a site's ID as specified by the site manager who created the site.
Subject ID	Indicates the number of the subject who just completed the dispensation or randomization visit.
Treatment Arm	Indicates the corresponding treatment arm.
Visit Name	Indicates the name of a dispensation or randomization visit.
Number of Kits	Indicates the quantity and type of kits that were dispensed, along with a description. For example, if the kits are locally sourced, and the type, such as a device.
Date of Transaction (UTC)	Indicates the exact date and time of the dispensation. For example: 10-Nov-2022 08:07 AM.
Performed By	Indicates the user who performed the kit dispensation.

Subject randomization notification (with failures)

This notification is sent when a subject is either successfully randomized or their randomization failed at a site that you're assigned to.

Permissions required to receive the notification

Assign either or both of these permissions to anyone who needs to receive this notification:

- *Receive the Unblinded Randomization Notification*: Includes treatment details.
- *Receive the Randomization Notification*: It doesn't include treatment details. However, if the current randomization is unblinded, users who are assigned this blinded permission will also receive information on treatment arms.

Field descriptions

The table below describes all fields and details in both the Unblinded randomization and Randomization notifications.

> **Note:**
>
> Treatment arm details are included only for the **Unblinded randomization notification**.

Field	Description
Title	The title is comprised of three distinct elements: • The study's ID. For example, a study's ID can be P01-123-A3. • A subject's number. For example: MUHC22. • The title of the notification and whether the subject successfully randomizaed or not. For example, Randomization succeeded or Randomization failed.
Study ID	Indicates a study's ID again, as specified by the global study creator when they created the study.
Study Mode	Indicates the study mode associated with this notification. Displayed values can be Test, Active, or Training.
Site Name	Indicates the name of the site as specified by a site manager when they created the site.
Site ID	Indicates a site's ID as specified by the site manager who created the site.
Subject ID	Indicates the number of the subject who just completed the randomization visit.
Visit Name	Indicates the name of a randomization visit.
Date of Transaction (UTC)	Indicates the exact date and time of the randomization. For example: 10-Nov-2022 08:07 AM.
Performed By	Indicates the user who performed the randomization.

Subject Number Update notification

This notification is sent when a site or sponsor user has successfully updated a subject number, and contains the site number, the original subject number, and the new subject number.

Permissions required to receive the notification

Assign the *Receive the Subject Number Update Notification* permission to anyone who wants to receive this notification. Users receive notifications only for the sites that they're assigned to.

Field descriptions

The table below describes all fields and details in the **Subject number update notification**.

Field	Description
Title	The title is comprised of two distinct elements: • A subject's number. For example: MUHC22. • The title of the notification, for example, Subject Number Update.

Field	Description
Study Name	Indicates a study's ID, as specified by the global study creator when they created the study.
Study Mode	Indicates the study mode associated with this notification. Displayed values can be Test, Active, or Training.
Site Name	Indicates the name of the site as specified by a site manager when they created the site.
Original Subject Number	Indicates the original number of the subject that was updated.
New Subject Number	Indicates the new number of the subject.
Date Performed (UTC)	Indicates the exact date and time of the subject number update. For example: 10-Nov-2022 08:07 AM.
Subject Number Changed By	Indicates the username of the user who updated the subject's number.

Subject rollover notification

This notification is sent when a subject is enrolled into a rollover study.

Permissions required to receive the notification

Assign the *Receive the Subject Rollover Notification* permission to anyone who wants to receive this notification. Users receive notifications only for the sites that they're assigned to.

Field descriptions

The table below describes all fields and details in the **Subject rollover notification**.

Field	Description
Title	The title is comprised of three distinct elements: • The study's ID. For example, a study's ID can be P01-123-A3. • A subject's number. For example: MUHC22. • The title of the notification, for example, Rollover visit (Success).
Study ID	Indicates a study's ID again, as specified by the global study creator when they created the study.
Study Mode	Indicates the study mode associated with this notification. Displayed values can be Test, Active, or Training.
Site Name	Indicates the name of the site as specified by a site manager when they created the site.
Site ID	Indicates a site's ID as specified by the site manager who created the site.
Subject ID	Indicates the number of the subject who was enrolled in a rollover study.
Visit Name	Indicates the name of the visit the subject has just completed.
Visit Date / Time	Indicates the date and time the visit took place.

Field	Description
Date Performed (UTC)	Indicates the exact date and time the subject was enrolled in a rollover study. For example: 10-Nov-2022 08:07 AM.
Performed By	Indicates the username of the user who enrolled the subject in a rollover study.

Subject screening notification (with error)

This notification is sent when a subject is either successfully screened or their screening failed at a site that you're assigned to.

Permissions required to receive the notification

Assign the *Receive the Subject Screening Notification* permission to anyone who wants to receive this notification. Users receive notifications for only the sites that they're assigned to.

Field descriptions

The table below describes all fields and details in the **Subject screening notification**.

Field	Description
Title	The title is comprised of three distinct elements: • The study's ID. For example, a study's ID can be P01-123-A3. • A subject's number. For example: MUHC22. • The title of the notification, for example, Screening Notification or Screening Failure Notification.
Study Name	Indicates a study's ID again, as specified by the global study creator when they created the study.
Study Mode	Indicates the study mode associated with this notification. Displayed values can be Test, Active, or Training.
Site Name	Indicates the name of the site as specified by a site manager when they created the site.
Subject Number	Indicates the number of the subject.
Date Performed (UTC)	Indicates the exact date and time of the completion of the screening visit. For example: 10-Nov-2022 08:07 AM.
Performed By	Indicates the username of the user who performed the screening of the subject.

Subject transferred notification

This notification is sent when a subject transfers to another site.

Permissions required to receive the notification

Assign the *Receive the Subject Transferred Notification* permission to anyone who wants to receive this notification. Users receive notifications for only the sites that they're assigned to.

Field descriptions

The table below describes all fields and details in the **Subject transferred notification**.

Field	Description
Title	The title is comprised of three distinct elements: • A subject's number. For example: MUHC22. • The previous site name. • The name of the site the subject was transferred to.
Study Name	Indicates a study's ID again, as specified by the global study creator when they created the study.
Study Mode	Indicates the study mode associated with this notification. Displayed values can be Test, Active, or Training.
Previous Site Name	Indicates the name of the previous site the subject was assigned to, as specified by a site manager when they created the site.
Previous Study Org ID	Indicates the previous study org number.
New Site Name	Indicates the name of the new site the subject was transferred to.
New Study Org ID	Indicates the new study org number.
Previous Subject Number	Indicates the subject's previous number before transferring to another site.
New Subject Number	Indicates the new ID the subject was assigned after transferring.
Date Performed (UTC)	Indicates the exact date and time of the subject transfer. For example: 10-Nov-2022 08:07 AM.
Performed By	Indicates the username of the user who transferred the subject.

Subject unscheduled visit notification

This notification is sent when a subject completes an unscheduled visit.

Permissions required to receive the notification

Assign the *Receive the Unscheduled Visit Notification* permission to anyone who wants to receive this notification. Users receive notifications only for the sites that they're assigned to.

Field descriptions

The table below describes all fields and details in the **Subject unscheduled visit notification**.

Field	Description
Title	The title is comprised of three distinct elements: • The study's ID. For example, a study's ID can be P01-123-A3. • A subject's number. For example: MUHC22. • The title of the notification and whether the visit was successfully completed or not. For example Unscheduled Visit (Success) or Unscheduled Visit (Failure).

Field	Description
Study ID	Indicates a study's ID again, as specified by the global study creator when they created the study.
Study Mode	Indicates the study mode associated with this notification. Displayed values can be Test, Active, or Training.
Site Name	Indicates the name of the site where the unscheduled visit took place. The name is displayed as it was specified by a site manager when they created the site.
Site ID	Indicates a site's ID as specified by the site manager who created the site.
Subject ID	Indicates the number of the subject who just completed the unscheduled visit.
Visit Name	Indicates an unscheduled visit's name, as specified by a study designer when they defined the study's visit schedule.
Visit Date/ Time	Indicates the unscheduled visit's exact date and time of the completion. For example: 10-Nov-2022 08:07 AM.
Date Performed (UTC)	Indicates the exact date and time of the subject completed the unscheduled visit. For example: 10-Nov-2022 08:07 AM.
Performed By	Indicates the username of the user who performed the unscheduled visit completion.

Subject visit notification (with failures)

This notification is sent when a subject completes a non-dispensation or optional visit at a site that you're assigned to. The notification is sent both when the subject successfully completes the visit and when one or more questions have errors.

Permissions required to receive the notification

Assign the *Receive the Subject Visit Notification* permission to anyone who wants to receive this notification. Users receive notifications for only the sites that they're assigned to.

Field descriptions

The table below describes all fields and details in the **Subject visit notification**.

Field	Description
Title	The title is comprised of three distinct elements: • The study's ID. For example, a study's ID can be P01-123-A3. • A subject's number. For example: MUHC22. • The title of the notification and whether the visit was successfully completed or not.
Study ID	Indicates a study's ID again, as specified by the global study creator when they created the study.
Study Mode	Indicates the study mode associated with this notification. Displayed values can be Test, Active, or Training.

Field	Description
Site Name	Indicates the name of the site where the visit took place. The name is displayed as it was specified by a site manager when they created the site.
Site ID	Indicates a site's ID as specified by the site manager who created the site.
Subject ID	Indicates the number of the subject who just completed the visit.
Visit Name	Indicates a visit's name, as specified by a study designer when they defined the study's visit schedule.
Visit Date/ Time	Indicates the visit's exact date and time of the completion. For example: 10-Nov-2022 08:07 AM.

Subject withdrawal notification

This notification is sent when a subject withdraws from a study.

Permissions required to receive the notification

Assign the *Receive the Subject Completion Notification* permission to anyone who wants to receive this notification. Users receive notifications for only the sites that they're assigned to.

Field descriptions

The table below describes all fields and details in the **Subject withdrawal notification**.

Field	Description
Title	The title is comprised of three distinct elements: • The study's ID. For example, a study's ID can be P01-123-A3. • A subject's number. For example: MUHC22. • The title of the notification, for example, Withdrawal Notification.
Study ID	Indicates a study's ID again, as specified by the global study creator when they created the study.
Study Mode	Indicates the study mode associated with this notification. Displayed values can be Test, Active, or Training.
Site Name	Indicates the name of the site as specified by a site manager when they created the site.
Subject Number	Indicates the number of the subject who just withdrew from the study.
Date Performed (UTC)	Indicates the exact date and time of the subject withdrawal. For example: 10-Nov-2022 08:07 AM.
Performed By	Indicates the username of the user who performed the subject withdrawal.

Training complete notification

This notification is sent when you complete all training assigned to you.

Permissions required to receive the notification

Any role that is assigned training will receive this notification after they completed all training assigned to them. Users receive notifications for only the sites that they're assigned to.

Field descriptions

The table below describes all fields and details in the **Training complete notification**.

Field	Descriptions
Title	The title is comprised of two distinct elements: • The study's ID. For example, a study's ID can be P01-123-A3. • The title of the notification, for example, Training complete.
Study ID	Indicates a study's ID again, as specified by the global study creator when they created the study.
Study Mode	Indicates the study mode associated with this notification. Displayed values can be Test, Active, or Training.
Site Name	Indicates the name of the site as specified by a site manager when they created the site.
Site ID	Indicates a site's ID as specified by the site manager who created the site.
Date of Transaction (UTC)	Indicates the exact date and time the training was complete. For example: 10-Nov-2022 08:07 AM.
Performed By	Indicates the username of the user who completed the training.

Unblinded Pharmacist dispensation notification

Pharmacists or unblinded site users receive this notification when kits need to be dispensed by them to subjects during a dispensation or randomization visit.

> **Note:**
>
> Details about partial dispensation are included when they occur.

Permissions required to receive the notification

Assign the *Receive the Unblinded Pharmacist Dispensation Notification* permission to anyone who wants to receive this notification. Users receive notifications for only the sites that they're assigned to.

Field descriptions

The table below describes all fields and details in the **Unblinded Pharmacist dispensation notification**.

> **Note:**
>
> This notification also contains an introductory statement suggesting you see your pharmacist or unblinded site user, as there may be kits for this visit only the pharmacist can dispense.

Field	Description
Title	The title is comprised of three distinct elements: • The study's ID. For example, a study's ID can be P01-123-A3. • A subject's number. For example: MUHC22. • The title of the notification and whether the dispensation succeeded or not. For example Unblinded dispensation succeeded or Unblinded dispensation failed.
Study ID	Indicates a study's ID again, as specified by the global study creator when they created the study.
Study mode	Indicates the study mode associated with this notification. Displayed values can be Test, Active, or Training.
Site Name	Indicates the name of the site where the unscheduled visit took place. The name is displayed as it was specified by a site manager when they created the site.
Site ID	Indicates a site's ID as specified by the site manager who created the site.
Subject ID	Indicates the number of the subject who just completed the dispensation or randomization visit.
Visit Name	Indicates the name of a dispensation or randomization visit.
Number of Kits	Indicates the total number of kits.
Kit Number	Indicates the number of the specific kit(s) that have been dispensed successfully or failed to be dispensed.
Description	A description of the kit, for example, Matching placebo.
Date of Transaction (UTC)	Indicates the exact date and time of the dispensation. For example: 10-Nov-2022 08:07 AM.
Performed By	Indicates the user who performed the kit dispensation.

User Upload notifications

These notifications are sent after the user upload process completes.

Permissions required to receive these notifications

A user associated to a role, that includes the *Upload Users in Bulk* permission, and processes a user import can receive the following notifications:

- **Users uploaded successfully**
- **User upload completed with errors. Error report is ready for your records**

- **User Upload Error Report is ready in Clinical One**

Field descriptions

The table below describes all fields and details in the notifications listed above.

Field	Description
Title	The title is comprised of two distinct elements: • The study's ID. For example, a study's ID can be P01-123-A3. • A statement about users uploading successfully or the report being ready.
Study Name	Indicates a study's ID again, as specified by the global study creator when they created the study.
Number of records	Indicates the number of data records in the generated report.
Server Run Date	Indicates the exact date and time when the report was generated. For example: 10-Nov-2022 08:07 AM.
Report Run ID	Indicates the GUID of the report generation in the application's backend system. For example: 08B6771F54DE45A0BB576E86326BE84D.

Supply notifications

- Depot shipment notification
  This notification is sent when a depot-to-depot shipment is generated. The notification is created for both automatic and manual shipments.
- Kit List Deactivated notification (with failures)
  This notification is sent to a clinical supply manager when a kit list was successfully deactivated or the system failed to deactivate it.
- Kit missing from shipment for destruction notification
  This notification is sent when a depot user marks a kit as missing from a shipment for destruction.
- Kit status changed to "Misallocated" notification
  This notification is sent once a site user changes the status of a kit to Misallocated.
- Kit status changed to "Not Dispensed to Subject" notification
  This notification is sent when a site user updates the status of a kit to Not Dispensed to Subject.
- Kits Quarantined notification
  This notification is sent when individual kits have been marked as Quarantined.
- Low kit depot notification
  This notification is sent when the depot kits run below the limit set up in depot configuration.
- Kits Released from Quarantine notification
  This notification is sent when individual kits have been released from quarantine.
- Pending shipment notification
  This notification is sent when a shipment has not yet been registered at a site within the time span specified in the supply settings.

- Randomization update notification
  This notification is sent when the randomization limit is reached for a study, site, country, or cohort; or a percentage of the randomization limit is reached for a study, site, country, or cohort.
- Replace kits notification
  This notification is sent when a kit has been replaced for a subject.
- Shipment cancellation notification
  This notification is sent every time a shipment is cancelled.
- Shipment failure notification
  This notification is sent when a shipment can't be created, such as because of lack of inventory.
- Shipment for destruction created notification
  This notification is sent when a new shipment for destruction is generated.
- Shipment for destruction not received notification
  This notification is sent when a depot doesn't confirm that they received one or all kits in a shipment.
- Shipment Quarantined notification
  This notification is sent when a shipment has been quarantined.
- Shipment Released from Quarantine notification
  This notification is sent when a shipment has been released from quarantine.
- Shipment request notification
  This notification is sent when a site requests a shipment, and when a new shipment is generated. The notification is created for both automatic and manual shipments.
- State of a site has changed notification
  This notification is sent when a site changes status, such as going from New to Active.
- Study inventory notification
  This notification is sent when available kits at a site or depot are marked as Quarantined.
- Unblinded dose hold performed notification
  This notification is sent once a site user performs a dose hold.

Depot shipment notification

This notification is sent when a depot-to-depot shipment is generated. The notification is created for both automatic and manual shipments.

Permissions required to receive the notification

Assign the *Receive Notification of Depot Shipments* permission to anyone who wants to receive this notification. Users receive notifications only for the depots that they are assigned to.

Field descriptions

The table below describes all fields and details in the **New depot shipment notification**.

Field	Description
Title	The title is comprised of two distinct elements: • The study's ID. For example, a study's ID can be P01-123-A3. • The title of the notification, for example, New Depot Shipment .
Study ID	Indicates a study's ID again, as specified by the global study creator when they created the study.
Study Mode	Indicates the study mode associated with this notification. Displayed values can be Test, Active, or Training.
Destination ID	Indicates a facility's ID as specified by the user who created the facility in the study. For example: • For a site, it indicates a site's ID as specified by a site manager. • For a depot, it indicates a depot's ID as specified by a clinical supply manager.
Destination	Indicates the destination of the shipment, as specified by the user who created the facility in the study. For example: • For a site, it indicates the site's title as specified by a site manager. • For a depot, it indicates a depot's title as specified by a clinical supply manager.
Origin(s)	Indicates the name of the depot facility that is packaging and shipping the depot for its destination. The name appears as specified by a clinical supply manager when they created the depot.
Shipment ID	Indicates a shipment's ID assigned at creation.
Shipment Type	Indicates a shipment's type, as requested by the depot user, the site user, or the clinical supply manager. The following types can be displayed: • Manual • Shipment for Destruction
Date of Transaction (UTC)	Indicates the exact date and time of the shipment creation. For example: 10-Nov-2022 08:07 AM.
Performed By	Indicates the user who created the new depot shipment.

Kit List Deactivated notification (with failures)

This notification is sent to a clinical supply manager when a kit list was successfully deactivated or the system failed to deactivate it.

Permissions required to receive the notification

Assign the *Upload and Generate Inventory Lists* to anyone who wants to receive this notification. Users receive notifications for only the sites that they're assigned to.

Field descriptions

The table below describes all fields and details in the **Kit List Deactivated notification**.

Field	Description
Title	The title is comprised of two distinct elements: • The study's ID. For example, a study's ID can be P01-123-A3. • The title of the notification, for example, Kit List Deactivated.
Study Name	Indicates a study's name, as specified by the global study creator when they created the study.
Study ID	Indicates a study's ID again, as specified by the global study creator when they created the study.
Study Mode	Indicates the study mode associated with this notification. Displayed values can be Test, Active, or Training.
Kit List	Indicates the deactivated kit list's title, as specified by the clinical supply manager or statistician when they created the kit list.
Number of Kits	Indicates the number of kits specified in the deactivated kit list.
Date of Transaction (UTC)	Indicates the exact date and time of the shipment request. For example: 10-Nov-2022 08:07 AM.
Performed By	Indicates the user who deactivated the kit list.

Kit missing from shipment for destruction notification

This notification is sent when a depot user marks a kit as missing from a shipment for destruction.

Permissions required to receive the notification

Assign the *Receive Notification of Shipments* permission to anyone who wants to receive this notification. Users receive notifications for only the sites that they're assigned to.

Field descriptions

The table below describes all fields and details in the **Kit missing from shipment for destruction notification**.

Field	Description
Title	The title is comprised of three distinct elements: • The study's ID. For example, a study's ID can be P01-123-A3. • The shipment ID. • The title of the notification, for example, Kit missing from shipment for destruction.
Study ID	Indicates a study's ID again, as specified by the global study creator when they created the study.
Study Mode	Indicates the study mode associated with this notification. Displayed values can be Test, Active, or Training.
Site Name	Indicates the name of the site where the kit is missing. The name is displayed as it was specified by a site manager when they created the site.

Field	Description
Site ID	Indicates a site's ID as specified by the site manager who created the site.
Depot	Indicates the depot's name as specified by a clinical supply manager when they created the facility in the study.
Depot Shipping Address	Indicates the depot's shipping address as specified by the clinical supply manager when they created the facility in the study.
Shipment	Indicates a shipment's ID.
Missing Kits	Indicates the exact number of missing kits from a shipment, as well as the kit numbers that are missing from that specific shipment.
Date Shipment Received (UTC)	Indicates the exact date and time the kit was marked as missing from the shipment. For example: 10-Nov-2022 08:07 AM.

Kit status changed to "Misallocated" notification

This notification is sent once a site user changes the status of a kit to Misallocated.

Permissions required to receive the notification

Assign the *Receive the Unblinded Kit Misallocation Notification* permission to anyone who wants to receive this notification. Users receive notifications for only the sites that they're assigned to.

Field descriptions

The table below describes all fields and details in the **Kit status changed to "Misallocated" notification**.

> **Note:**
>
> This notification also contains an introductory statement notifying the receiver that a user has changed the status of a kit to *Misallocated*.

Field	Description
Title	The title is comprised of three distinct elements: • The study's ID. For example, a study's ID can be P01-123-A3. • The kit's ID. • The title of the notification, for example, Kit status changed to "Misallocated."
Study ID	Indicates a study's ID again, as specified by the global study creator when they created the study.
Study mode	Indicates the study mode associated with this notification. Displayed values can be Test, Active, or Training.

Field	Description
Site Name	Indicates the name of the site where the kit was marked as Misallocated. The name is displayed as it was specified by a site manager when they created the site.
Site ID	Indicates a site's ID as specified by the site manager who created the site.
Subject ID	Indicates the number of the subject who is associated with the misallocated kit.
Kit Number	Indicates the number of the specific kit(s) whose statuses have been updated.
Date of Transaction (UTC)	Indicates the exact date and time the status of the kit was updated. For example: 10-Nov-2022 08:07 AM.
Performed By	Indicates the user who updated the status of the kit.

Kit status changed to "Not Dispensed to Subject" notification

This notification is sent when a site user updates the status of a kit to Not Dispensed to Subject.

> **Note:**
>
> The unblinded pharmacist user will receive both Blinded and Unblinded Pharmacist kit notifications.

Permissions required to receive the notification

Assign all or one of these permissions to anyone who wants to receive this notification:

- *Receive the Not Dispensed to Subject (Unblinded) notification*: Includes a description for the kit that has been updated with the status Not Dispensed to Subject and notifies the user that the kit has been removed from the subject's history.
- *Receive the Not Dispensed to Subject notification*: Notifies users that the status of a kit has been updated to Not Dispensed to Subject and that the kit has been removed from the subject's history. It includes the kit description only if the updated kit is of Blinded distribution type. If not, this information is not disclosed in the notification.
- *Receive the Not Dispensed to Subject (Unblinded Pharmacist) notification*: Notifies users that the status of a kit has been changed to Not Dispensed to Subject and that the kit has been removed from the subject's history. A description for the updated kit is also included.

Field descriptions

The table below describes all fields and details in the **Kit status changed to "Not Dispensed to Subject" notification**.

> **Note:**
>
> This notification also contains an introductory statement notifying the receiver that a user has changed the status of a kit to *Not Dispensed to Subject*, and that the kit has been removed from the subject's history.

Field	Description
Title	The title is comprised of three distinct elements: • The study's ID. For example, a study's ID can be P01-123-A3. • The kit's ID. • The title of the notification, for example, Kit status changed to "Not Dispensed to Subject."
Study ID	Indicates a study's ID again, as specified by the global study creator when they created the study.
Study mode	Indicates the study mode associated with this notification. Displayed values can be Test, Active, or Training.
Site Name	Indicates the name of the site where the unscheduled visit took place. The name is displayed as it was specified by a site manager when they created the site.
Site ID	Indicates a site's ID as specified by the site manager who created the site.
Subject ID	Indicates the number of the subject who just completed the dispensation or randomization visit.
Kit Number	Indicates the number of the specific kit(s) whose statuses have been updated.
Date of Transaction (UTC)	Indicates the exact date and time of the dispensation. For example: 10-Nov-2022 08:07 AM.
Performed By	Indicates the user who updated the status of the kit.

Kits Quarantined notification

This notification is sent when individual kits have been marked as Quarantined.

Permissions required to receive the notification

Assign all or one of these permissions to anyone who wants to receive this notification:

- *Receive the Quarantined Depot Shipment Notification*: Includes the kit number and quantity of kits quarantined at the associated depot.
- *Receive the Quarantined Site Shipment Notification*: Includes the kit number and quantity of kits quarantined at the associated site.

Field descriptions

The table below describes all fields and details in both the **Quarantined depot shipment** and **Quarantined site shipment notifications**.

Field	Description
Title	The title is comprised of three distinct elements: • The study's ID. For example, a study's ID can be P01-123-A3. • The shipment ID. • The title of the notification, for example, Shipment Quarantined.
Study ID	Indicates a study's ID again, as specified by the global study creator when they created the study.
Study Mode	Indicates the study mode associated with this notification. Displayed values can be Test, Active, or Training.
Shipment ID	Indicates a shipment's ID assigned at creation.
Temperature Excursion ID	Indicates the tracking number associated with the kit(s) that experienced a temperature excursion and have been quarantined.
Kit Number	Indicates the number of the specific kit(s) that have been quarantined.
Origin(s)	Indicates the name of the depot facility that is packaging and shipping the depot for its destination. The name appears as specified by a clinical supply manager when they created the depot.
Destination	Indicates the destination of the shipment, as specified by the user who created the facility in the study. For example: • For a site, it indicates the site's title as specified by a site manager. • For a depot, it indicates a depot's title as specified by a clinical supply manager.
Destination ID	Indicates a facility's ID as specified by the user who created the facility in the study. For example: • For a site, it indicates a site's ID as specified by a site manager. • For a depot, it indicates a depot's ID as specified by a clinical supply manager.
Shipping Address	Indicates the address of the shipment destination.
Date of Transaction (UTC)	Indicates the exact date and time of the kit quarantine. For example: 10-Nov-2022 08:07 AM.
Performed By	Indicates the user who performed the kit quarantine.

Low kit depot notification

This notification is sent when the depot kits run below the limit set up in depot configuration.

The **Low kit depot notification** alerts clinical supplies managers that the depot is running low on supplies. An alert will be sent when the resupply algorithm detects the low inventory limit set in the depot configuration is reached. Users receive it until they restock on supplies.

Permissions that receive the notification
To receive these notifications, assign the *Receive Notification of Depot Shipments* permission to anyone who wants to receive this notification. Users receive notifications only for the depots that they are assigned to.

Table 3-5 Field descriptions

Field	Description
Subject line	Indicates the summary of the alert information.
Study Name	Indicates the name of the study for which the given depot is running low on supplies.
Study ID	Indicates a study's ID, as specified upon study creation.
Study Mode	Indicates the study mode in which the given depot is running low on supplies: *Production*, *Testing* or *Training*.
Depot ID	Indicates a short identifier for the depot.
Depot Name	Indicates the depot's descriptive name..
Depot Address	Indicates the depot's address.
Kit Type ID	Indicates the unique identifier of the given kit that reached the low inventory limit. *Line will repeat for each kit type.*
Kit Type Description	Indicates the description of the given kit that reached the low inventory limit.
Current Inventory Available	Indicates the number of the kits available at the moment of the inquiry. ***In transit*** *and* ***Pre-Quarantined*** *kits will count in the total Current Inventory Available as if they were in Available status.*
Date Alert Raised	Indicates the exact date and time of the alert. For example: 22-Jan-2023 09:15 AM.

Kits Released from Quarantine notification

This notification is sent when individual kits have been released from quarantine.

Permissions required to receive the notification

Assign all or one of these permissions to anyone who wants to receive this notification:

- *Receive the Released from Quarantine Shipment Notification (Depot)*: Includes the kit number and quantity of kits released from quarantine at the associated depot.
- *Receive the Released from Quarantine Notification (Site)*: Includes the kit number and quantity of kits released from quarantine at the associated site.

Field descriptions

The table below describes all fields and details in both the **Shipment released from quarantine (Depot)** and **Shipment released from quarantine (Site) notifications**.

Field	Description
Title	The title is comprised of three distinct elements: • The study's ID. For example, a study's ID can be P01-123-A3. • The shipment ID. • The title of the notification, for example, Shipment released from quarantine (Depot) or Shipment released from quarantine (Site).
Study ID	Indicates a study's ID again, as specified by the global study creator when they created the study.
Study Mode	Indicates the study mode associated with this notification. Displayed values can be Test, Active, or Training.
Shipment ID	Indicates a shipment's ID assigned at creation.
Temperature Excursion ID	Indicates the tracking number associated with the kit(s) that experienced a temperature excursion and have been released from quarantined.
Kit Number	Indicates the number of the specific kit(s) that have been released from quarantine.
Origin(s)	Indicates the name of the depot facility that is packaging and shipping the depot for its destination. The name appears as specified by a clinical supply manager when they created the depot.
Destination	Indicates the destination of the shipment, as specified by the user who created the facility in the study. For example: • For a site, it indicates the site's title as specified by a site manager. • For a depot, it indicates a depot's title as specified by a clinical supply manager.
Destination ID	Indicates a facility's ID as specified by the user who created the facility in the study. For example: • For a site, it indicates a site's ID as specified by a site manager. • For a depot, it indicates a depot's ID as specified by a clinical supply manager.
Shipping Address	Indicates the address of the shipment destination.
Date of Transaction (UTC)	Indicates the exact date and time the kits were released from quarantine. For example: 10-Nov-2022 08:07 AM.
Performed By	Indicates the user who performed the quarantine release.

Pending shipment notification

This notification is sent when a shipment has not yet been registered at a site within the time span specified in the supply settings.

Permissions required to receive the notification

Assign the *Receive Notification of Shipments* permission to anyone who wants to receive this notification. Users receive notifications for only the sites that they're assigned to.

Field descriptions

The table below describes all fields and details in the **Pending shipment notification**.

Field	Description
Title	The title is comprised of two distinct elements: • The study's ID. For example, a study's ID can be P01-123-A3. • The title of the notification, for example, Pending Shipments.
Study Name	Indicates a study's name, as specified by the global study creator when they created the study.
Study ID	Indicates a study's ID again, as specified by the global study creator when they created the study.
Study Mode	Indicates the study mode associated with this notification. Displayed values can be Test, Active, or Training.
Shipment ID	Indicates a shipment's ID assigned at creation.
From Depot ID	Indicates a short identifier for the depot the shipment left from.
To Site ID	Indicates a site's ID, as specified by the site manager who created the site, that the shipment is being sent to.
Date Shipment Left Depot (UTC)	Indicates the exact date and time the shipment left the depot. For example: 10-Nov-2022 08:07 AM.

Randomization update notification

This notification is sent when the randomization limit is reached for a study, site, country, or cohort; or a percentage of the randomization limit is reached for a study, site, country, or cohort.

Permissions required to receive this notification

Assign the *Receive the Randomization Notification* permission to anyone who wants to receive this notification. Users receive notifications for only the sites that they're assigned to.

Field descriptions

The table below describes all fields and details in the **Randomization update notification**.

Field	Description
Title	The title is comprised of two distinct elements: • The study's ID. For example, a study's ID can be P01-123-A3. • The title of the notification.
Study ID	Indicates a study's ID again, as specified by the global study creator when they created the study.
Study Mode	Indicates the study mode associated with this notification. Displayed values can be Test, Active, or Training.

Field	Description
Site Name	Indicates the name of the site where the unscheduled visit took place. The name is displayed as it was specified by a site manager when they created the site.
Site ID	Indicates a site's ID as specified by the site manager who created the site.
Subject ID	Indicates the number of the subject who just completed the dispensation or randomization visit.
Visit Name	Indicates the name of a dispensation or randomization visit.
Date of Transaction (UTC)	Indicates the exact date and time of the randomization. For example: 10-Nov-2022 08:07 AM.
Performed By	Indicates the user who performed the randomization.

Replace kits notification

This notification is sent when a kit has been replaced for a subject.

Permissions required to receive the notification

Assign the *Receive Notification of Shipments* permission to anyone who wants to receive this notification.

Field descriptions

The table below describes all fields and details in the **Replace kits notification**.

Field	Description
Title	The title of the notification, for example, Replace Kits.
Subject number	A system message followed by the subject's number whose kit is being replaced.
Kit number	Indicates the number of the kit the previous kit is being replaced with.

Shipment cancellation notification

This notification is sent every time a shipment is cancelled.

Permissions required to receive the notification

Assign the *Receive Notification of Shipments* permission to anyone who wants to receive this notification. Users receive notifications for only the sites that they're assigned to.

Field descriptions

The table below describes all fields and details in the **Shipment cancellation notification**.

Field	Description
Title	The title is comprised of two distinct elements: • The study's ID. For example, a study's ID can be P01-123-A3. • The title of the notification, for example, Shipment Cancellation Request.
Study ID	Indicates a study's ID again, as specified by the global study creator when they created the study.
Study Mode	Indicates the study mode associated with this notification. Displayed values can be Test, Active, or Training.
Destination ID	Indicates a facility's ID as specified by the user who created the facility in the study. For example: • For a site, it indicates a site's ID as specified by a site manager. • For a depot, it indicates a depot's ID as specified by a clinical supply manager.
Destination	Indicates the destination of the shipment, as specified by the user who created the facility in the study. For example: • For a site, it indicates the site's title as specified by a site manager. • For a depot, it indicates a depot's title as specified by a clinical supply manager.
Origin(s)	Indicates the name of the depot facility that is packaging and shipping the depot for its destination. The name appears as specified by a clinical supply manager when they created the depot.
Shipment ID	Indicates a shipment's ID assigned at creation.
Shipment Type	Indicates a shipment's type, as requested by the depot user, the site user, or the clinical supply manager. The following types can be displayed: • Manual • Shipment for Destruction
Date of Transaction (UTC)	Indicates the exact date and time of the shipment cancellation. For example: 10-Nov-2022 08:07 AM.
Performed By	Indicates the user who performed the cancellation of the shipment.

Shipment failure notification

This notification is sent when a shipment can't be created, such as because of lack of inventory.

Permissions required to receive the notification

Assign the *Receive Notification of Shipments* permission to anyone who wants to receive this notification. Users receive notifications for only the sites that they're assigned to.

Field descriptions

The table below describes all fields and details in the **Shipment failure notification**.

Field	Description
Title	The title is comprised of two distinct elements: • The study's ID. For example, a study's ID can be P01-123-A3. • The reason for a shipment's failure. For example, that reason can be Insufficient kits when creating shipment.
Study ID	Indicates a study's ID again, as specified by the global study creator when they created the study.
Study Mode	Indicates the study mode associated with this notification. Displayed values can be Test, Active, or Training.
Shipment Type	Indicates a shipment's type, as requested by the depot user, the site user, or the clinical supply manager. The following types can be displayed: • Manual • Shipment for Destruction
Destination ID	Indicates a facility's ID as specified by the user who created the facility in the study. For example: • For a site, it indicates a site's ID as specified by a site manager. • For a depot, it indicates a depot's ID as specified by a clinical supply manager.
Destination	Indicates the destination of the shipment, as specified by the user who created the facility in the study. For example: • For a site, it indicates the site's title as specified by a site manager. • For a depot, it indicates a depot's title as specified by a clinical supply manager.
Origin(s)	Indicates the name of the depot facility that is packaging and shipping the depot for its destination. The name appears as specified by a clinical supply manager when they created the depot.
Date of Transaction (UTC)	Indicates the exact date and time of the shipment failure. For example: 10-Nov-2022 08:07 AM.

Shipment for destruction created notification

This notification is sent when a new shipment for destruction is generated.

Permissions required to receive the notification

Assign the *Receive Notification of Shipments* permission to anyone who wants to receive this notification. Users receive notifications for only the sites that they're assigned to.

Field descriptions

The table below describes all fields and details in the **Shipment for destruction created notification**.

Field	Description
Title	The title is comprised of two distinct elements: • The study's ID. For example, a study's ID can be P01-123-A3. • The title of the notification, for example, Shipment for destruction created.
Study ID	Indicates a study's ID again, as specified by the global study creator when they created the study.
Study Mode	Indicates the study mode associated with this notification. Displayed values can be Test, Active, or Training.
Shipment Type	Indicates a shipment's type, as requested by the depot user, the site user, or the clinical supply manager. The following types can be displayed: • Manual • Shipment for Destruction
Destination ID	Indicates a facility's ID as specified by the user who created the facility in the study. For example: • For a site, it indicates a site's ID as specified by a site manager. • For a depot, it indicates a depot's ID as specified by a clinical supply manager.
Destination	Indicates the destination of the shipment, as specified by the user who created the facility in the study. For example: • For a site, it indicates the site's title as specified by a site manager. • For a depot, it indicates a depot's title as specified by a clinical supply manager.
Origin(s)	Indicates the name of the depot facility that is packaging and shipping the depot for its destination. The name appears as specified by a clinical supply manager when they created the depot.
Date of Transaction (UTC)	Indicates the exact date and time the shipment for destruction was created. For example: 10-Nov-2022 08:07 AM.

Shipment for destruction not received notification

This notification is sent when a depot doesn't confirm that they received one or all kits in a shipment.

Permissions required to receive the notification

Assign the *Receive Notification of Shipments* permission to anyone who wants to receive this notification. Users receive notifications for only the sites that they're assigned to.

Field descriptions

The table below describes all fields and details in the **Shipment for destruction not received notification**.

Field	Description
Title	The title is comprised of two distinct elements: • The study's ID. For example, a study's ID can be P01-123-A3. • The title of the notification, for example, Shipment for destruction not received.
Study ID	Indicates a study's ID again, as specified by the global study creator when they created the study.
Study Mode	Indicates the study mode associated with this notification. Displayed values can be Test, Active, or Training.
Shipment Type	Indicates a shipment's type, as requested by the depot user, the site user, or the clinical supply manager. The following types can be displayed: • Manual • Shipment for Destruction
Destination ID	Indicates a facility's ID as specified by the user who created the facility in the study. For example: • For a site, it indicates a site's ID as specified by a site manager. • For a depot, it indicates a depot's ID as specified by a clinical supply manager.
Destination	Indicates the destination of the shipment, as specified by the user who created the facility in the study. For example: • For a site, it indicates the site's title as specified by a site manager. • For a depot, it indicates a depot's title as specified by a clinical supply manager.
Origin(s)	Indicates the name of the depot facility that is packaging and shipping the depot for its destination. The name appears as specified by a clinical supply manager when they created the depot.
Date of Transaction (UTC)	Indicates the exact date and time the shipment for destruction was logged as not received. For example: 10-Nov-2022 08:07 AM.

Shipment Quarantined notification

This notification is sent when a shipment has been quarantined.

Permissions required to receive the notification

Assign all or one of these permissions to anyone who wants to receive this notification:

- *Receive the Quarantined Depot Shipment Notification*: Includes details on the quarantined shipment associated with the depot.
- *Receive the Quarantined Site Shipment Notification* : Includes details on the quarantined shipment associated with the site.

Field descriptions

The table below describes all fields and details in both the **Quarantined depot shipment** and **Quarantined site shipment notifications**.

Field	Description
Title	The title is comprised of three distinct elements: • The study's ID. For example, a study's ID can be P01-123-A3. • The shipment ID. • The title of the notification, for example, Shipment Quarantined.
Study ID	Indicates a study's ID again, as specified by the global study creator when they created the study.
Study Mode	Indicates the study mode associated with this notification. Displayed values can be Test, Active, or Training.
Shipment ID	Indicates a shipment's ID assigned at creation.
Origin(s)	Indicates the name of the depot facility that is packaging and shipping the depot for its destination. The name appears as specified by a clinical supply manager when they created the depot.
Destination	Indicates the destination of the shipment, as specified by the user who created the facility in the study. For example: • For a site, it indicates the site's title as specified by a site manager. • For a depot, it indicates a depot's title as specified by a clinical supply manager.
Destination ID	Indicates a facility's ID as specified by the user who created the facility in the study. For example: • For a site, it indicates a site's ID as specified by a site manager. • For a depot, it indicates a depot's ID as specified by a clinical supply manager.
Shipping Address	Indicates the address of the shipment destination.
Temperature Excursion ID	Indicates the tracking number associated with the kit(s) that experienced a temperature excursion and have been quarantined.
Date of Transaction (UTC)	Indicates the exact date and time of the shipment quarantine. For example: 10-Nov-2022 08:07 AM.
Performed By	Indicates the user who performed the shipment quarantine.

Shipment Released from Quarantine notification

This notification is sent when a shipment has been released from quarantine.

Permissions required to receive the notification

Assign all or one of these permissions to anyone who wants to receive this notification:

- *Receive the Released from Quarantine Shipment Notification (Depot)*: Includes details on the shipment released from quarantine at the associated depot.
- *Receive the Released from Quarantine Notification (Site)*: Includes details on the shipment released from quarantine at the associated site.

Field descriptions

The table below describes all fields and details in both the **Shipment released from quarantine (Depot)** and **Shipment released from quarantine (Site) notifications**.

Field	Description
Title	The title is comprised of three distinct elements: • The study's ID. For example, a study's ID can be P01-123-A3. • A subject's number. For example: MUHC22. • The title of the notification, for example, Shipment released from quarantine (Depot) or Shipment released from quarantine (Site).
Study ID	Indicates a study's ID again, as specified by the global study creator when they created the study.
Study Mode	Indicates the study mode associated with this notification. Displayed values can be Test, Active, or Training.
Shipment ID	Indicates a shipment's ID assigned at creation.
Origin(s)	Indicates the name of the depot facility that is packaging and shipping the depot for its destination. The name appears as specified by a clinical supply manager when they created the depot.
Destination	Indicates the destination of the shipment, as specified by the user who created the facility in the study. For example: • For a site, it indicates the site's title as specified by a site manager. • For a depot, it indicates a depot's title as specified by a clinical supply manager.
Destination ID	Indicates a facility's ID as specified by the user who created the facility in the study. For example: • For a site, it indicates a site's ID as specified by a site manager. • For a depot, it indicates a depot's ID as specified by a clinical supply manager.
Shipping Address	Indicates the address of the shipment destination.
Temperature Excursion ID	Indicates the tracking number associated with the kit(s) that experienced a temperature excursion and have been released from quarantined.
Date of Transaction (UTC)	Indicates the exact date and time the shipment was released from quarantine. For example: 10-Nov-2022 08:07 AM.
Performed By	Indicates the user who performed the quarantine release.

Shipment request notification

This notification is sent when a site requests a shipment, and when a new shipment is generated. The notification is created for both automatic and manual shipments.

Permissions required to receive the notification

Assign the *Receive Notification of Shipments* permission to anyone who wants to receive this notification. Users receive notifications for only the sites that they're assigned to.

Field descriptions

The table below describes all fields and details in the **New shipment request notification**.

Field	Description
Title	The title is comprised of two distinct elements: • The study's ID. For example, a study's ID can be P01-123-A3. • The title of the notification, for example, New Shipment Request.
Study ID	Indicates a study's ID again, as specified by the global study creator when they created the study.
Study Mode	Indicates the study mode associated with this notification. Displayed values can be Test, Active, or Training.
Destination ID	Indicates a facility's ID as specified by the user who created the facility in the study. For example: • For a site, it indicates a site's ID as specified by a site manager. • For a depot, it indicates a depot's ID as specified by a clinical supply manager.
Destination	Indicates the destination of the shipment, as specified by the user who created the facility in the study. For example: • For a site, it indicates the site's title as specified by a site manager. • For a depot, it indicates a depot's title as specified by a clinical supply manager.
Origin(s)	Indicates the name of the depot facility that is packaging and shipping the depot for its destination. The name appears as specified by a clinical supply manager when they created the depot.
Shipment ID	Indicates a shipment's ID assigned at creation.
Shipment Type	Indicates a shipment's type, as requested by the depot user, the site user, or the clinical supply manager. The following types can be displayed: • Manual • Shipment for Destruction
Date of Transaction (UTC)	Indicates the exact date and time of the shipment request. For example: 10-Nov-2022 08:07 AM.
Performed By	Indicates the user who created the new shipment request.

State of a site has changed notification

This notification is sent when a site changes status, such as going from New to Active.

Permissions required to receive the notification

Assign the *Receive Site has been Updated Notification* permission to anyone who wants to receive this notification. Users receive notifications for only the sites that they're assigned to.

Field descriptions

The table below describes all fields and details in the **State of a site has changed notification**.

Field	Description
Title	The title is comprised of three distinct elements: • The site's ID and name. • The kit's ID. • The title of the notification, for example, Site Status Change Alert.
Details	Provides information on the site's status change.

Study inventory notification

This notification is sent when available kits at a site or depot are marked as Quarantined.

Permissions required to receive the notification

Assign all or one of these permissions to anyone who wants to receive this notification:

- *Receive the Quarantined Depot Shipment Notification*: Includes details on Depot inventory kits that have been marked as *Quarantined*.
- *Receive the Quarantined Site Shipment Notification*: Includes details on Site inventory kits that have been marked as *Quarantined*.

Field descriptions

The table below describes all fields and details in **Study Inventory notification**.

Field	Description
Title	The title is comprised of three distinct elements: • The study's ID. For example, a study's ID can be P01-123-A3. • The shipment ID. • The title of the notification, for example, Study Inventory Quarantined.
Study ID	Indicates a study's ID again, as specified by the global study creator when they created the study.
Destination ID	Indicates a facility's ID as specified by the user who created the facility in the study. For example: • For a site, it indicates a site's ID as specified by a site manager. • For a depot, it indicates a depot's ID as specified by a clinical supply manager.

Field	Description
Destination Name	Indicates the destination of the shipment, as specified by the user who created the facility in the study. For example: • For a site, it indicates the site's title as specified by a site manager. • For a depot, it indicates a depot's title as specified by a clinical supply manager.
Shipping Address	Indicates the address of the shipment destination.
Temperature Excursion ID	Indicates the tracking number associated with the kit(s) that experienced a temperature excursion and have been quarantined.
Kit Number(s)	Indicates the number of the specific kit(s) whose statuses have been updated.
Date Quarantined (UTC)	Indicates the exact date and time of the inventory quarantine. For example: 10-Nov-2022 08:07 AM.

Unblinded dose hold performed notification

This notification is sent once a site user performs a dose hold.

Permissions required to receive the notification

Assign the *Receive the Unblinded Dose Hold Notification* to anyone who wants to receive this notification. Users receive notifications for only the sites that they're assigned to.

Field descriptions

The table below describes all fields and details in the **Unblinded dose hold performed notification**.

Field	Description
Title	The title is comprised of three distinct elements: • The study's ID. For example, a study's ID can be P01-123-A3. • A subject's number. For example: MUHC22. • The title of the notification, for example, Unblinded dose hold performed.
Study ID	Indicates a study's ID again, as specified by the global study creator when they created the study.
Study Mode	Indicates the study mode associated with this notification. Displayed values can be Test, Active, or Training.
Site Name	Indicates the name of the site as specified by a site manager when they created the site.
Site ID	Indicates a site's ID as specified by the site manager who created the site.
Subject ID	Indicates the number of the subject.
Treatment Arm	Indicates the corresponding treatment arm.
Visit Name	Indicates the name of the visit the subject has just completed.
Number of Dose Holds	Indicates the total number of dose holds performed.

Field	Description
Kit Type	Indicates the description of the kit.
Date of Transaction (UTC)	Indicates the exact date and time of the dose hold. For example: 10-Nov-2022 08:07 AM.
Performed By	Indicates the user who performed the dose hold.

Oracle CRF Submit notifications

Email and in-application notifications can be customized to notify sponsor and site users about the status of PDF and report requests and inform sponsor users about site-level PDF requests, downloads, and confirmation activities.

The system can even notify you if the Oracle CRF Submit storage limit is reaching capacity. For more information about these notifications and how to configure them, see Email and in-application notifications.

4 Revision history

Date	Part number	Description
21 May 2024	F91496-03	The Notifications section was reformatted so users can easily differentiate between Oracle Clinical One Platform and Oracle CRF Submit notifications.
May 2024	F91496-02	Formatting updates were made throughout the publication.
May 2024	F91496-01	Original version of this document.