




COVaxON Inventory Management Job Aid

Target Role: Inventory Manager

<p>Role Description:</p> <p>Responsible for creating shipment and inventory records, transferring inventory, recording wastage, excursion, and recall events in the COVaxON system</p> <p>Permission Level: COVax Inventory Manager</p> <ul style="list-style-type: none">Edit-Access for Shipment, Inventory, & Storage Location tabsRead-Only Client ProfileView DashboardsRead and Export Vaccine Inventory Report & Summary Client and Dose Administration Report	<p>Legend</p> <p> Pencil Icon Click this to edit any data fields</p> <p> Red Asterisks Indicates a required field</p> <p> All COVID public health measures must be followed in alignment with this process.</p>
<p>Your profile has a defined access level and is associated with an Authorized Organization, which means you can perform the above activities within COVaxON on behalf of that Authorized Organization (AO). For further details on setting up your profile, refer to the "Login, Logout, and User Settings" job aid.</p> <p>For any assistance, please email: covaxonsupport@ontario.ca</p>	<p>This document provides training on how to use the COVaxON system for the vaccine management process. Within the clinical package that each site received, there are additional forms and documentation that you will need to use alongside the COVaxON system.</p>

Data Privacy Disclaimer:

Users with access to COVaxON can see the demographic details and health numbers of other clients in the system when searching for a particular person. The information is presented this way to help ensure that users access the correct client record and to reduce the risk of either not locating a client's record or improperly creating duplicate client records.

As required by PHIPA and under the terms of the Acceptable Use Policy, system users are only permitted to access the information of individuals to whom they are providing care or for other purposes that are specifically authorized.

The system records detailed audit transaction logs that inform the Ministry of Health of which client records were accessed by each user, and what actions they took in the system. Any concerns that are identified about improper access to the system will be investigated and appropriate actions taken.

Inventory Management Scenario Overview

Inventory is managed by Authorized Organizations (AOs), which typically includes two scenarios:

- 1. For Hospitals/Clinics Receiving Inventory:** Shipments are received and inspected upon arrival. Inventory Managers will create a shipment and inventory record for each Lot# of inventory received which will be used for vaccinations at the clinic. The Public Health Unit (PHU) may request the hospital to act as a **Vaccine Depot** where they may manage and allocate inventory to other locations (Vaccination Events) such as Long-Term Care Homes.
- 2. For Public Health Units (PHUs) receiving inventory on behalf of LTCHs, RHs and NHs:** Shipments are received by the PHU at their main office storage, acting as the Vaccine Depot where they will be inspected, and shipment and inventory records will be created in the system to reflect what is received. After the initial shipment of inventory arrives at the PHU, they may allocate inventory to the locations (Vaccination Events) such as Long-Term Care Homes.
- 3. For Pharmacies/Primary Care Physicians Receiving Inventory:** Shipments are received and inspected upon arrival. Inventory Managers will create a shipment and inventory record for each Lot# of inventory received which will be used for vaccinations at the pharmacy. Inventory Managers will allocate all the inventory to the pharmacy/primary care location (Vaccination Event)

Inventory Management Activities

The information below reflects the high-level process/activities that Authorized Organizations must follow to manage the inventory within COVaxON. Please refer to the section that is relevant to you.

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MODULE 1 – Inventory Set Up



1. Create Shipment Records

Create a shipment record to document details in the system



2. Create Inventory Record

Create inventory record associated to shipment in the system



3. (Part I) Linking inventory to a Vaccination Event: Allocate the inventory at the Authorized Organization (AO) to a Vaccination Event (VE)
3. (Part II) Modifying a VE Inventory: Staff at the VE to track and update the VE inventory



4. Reconciling the Authorized Organization Inventory based on activities at the Vaccination Event(s)

MODULE 2 – Ad Hoc Inventory Activities



5 & 6 Transferring & Accepting Inventory

Transfer the appropriate amount of inventory to another AO



7. Creating Storage Locations

Create storage locations for inventory if it does not already exist in the system



8. Recording Wastage

Record wastage that occurs through AO/VE operations



9. Excursion Events Record

Record an excursion event that occurred at a specific storage location



10. Adjusting Total Doses

Adjust inventory quantity if required



11. Manage Recalls

When a product is recalled, COVaxON inventory records should be updated to reflect this



12. Rescind Recalls

If the inspection was cleared with no safety issues, Super Users can rescind the recall

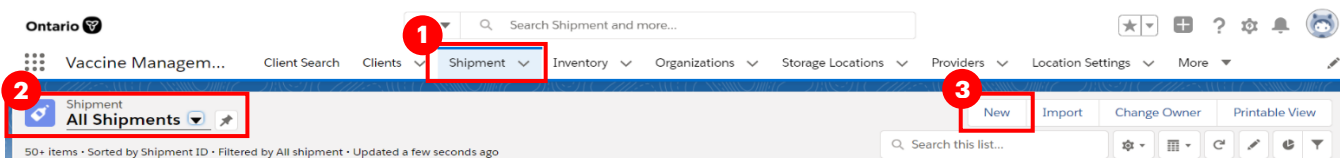
MODULE 1: INVENTORY SET UP

For a new Lot# of inventory that has been received at an Authorized Organization, create a shipment record, then create an inventory record associated to that shipment.



1. Creating Shipment Records

Description: After a shipment is received and inspected, create a shipment record to document the shipment information in the COVaxON system.



1. Go to the **"Shipment"** tab
2. Select **"All Shipments"** from dropdown
3. Select **"New"**

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4. Fill in the details of the new shipment
5. Select “**Save**” or “**Save & New**” for multiple shipment records

Note: all quantities with regards to inventory within COVaxON are measured in **doses, not vials**.

Shipment Details:

- **Shipment ID:** will be pre-populated based on the numerical order of creation
- **Authorized Organization:** Use the search bar to indicate the Authorized Organization (AO) responsible for the inventory. This is a mandatory field and must match the AO as on the User's profile.
- **Product:** This is a lookup field for products that already exist within the system (i.e. “PFIZER-BIONTECH COVID-19 mRNA”, or “MODERNA COVID-19 mRNA-1273” or “ASTRAZENECA COVID-19 Non-rep VV”) *Note: products are pre-loaded and do not need to be created (only selected from shipments and inventory). To view the products available, navigate to the “Products” tab.*
- **Quantity Received:** Indicates number of doses received. *Note: Pfizer previously indicated that there are 5 doses per vial. This has now been changed to 6 doses per vial. Based on the vials received, the Inventory Manager will need to perform the calculation to enter the doses received. This will be the same quantity entered on the inventory record (explained in section below)*
- **Temperature:** Temperature of shipment (UOM: °Celsius)
- **Status:** In most cases the status should be “**Received**”, but if part of the shipment was rejected, the status could be “**Partially Received**”
- **Requisition ID:** Can be entered if known for traceability to the order number
- **Total Shipped Quantity:** Enter quantity of doses that was shipped and planned to be received
- **Damaged Quantity:** Indicate the quantity of damaged doses (if any) that is recognized during shipment inspection
- **Reason:** Indicates the reason for damage (if any)

Further Context

- Pfizer immunizations require both vaccine and diluent inventory, whereas Moderna/AstraZeneca immunizations only require a vaccine (no diluent). Therefore, the following need to be created:
 - Pfizer: shipment and inventory records for vaccine **and** diluent
 - Moderna: shipment and inventory records for vaccine **only**
 - AstraZeneca: shipment and inventory records for vaccine **only**



2. Creating Inventory Records

Description:

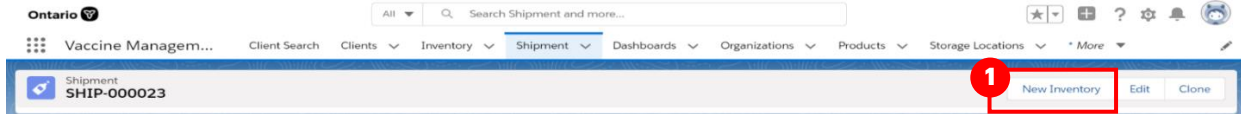
Once shipment record is created, create an inventory record(s) associated to the shipment. Inventory records should be created for each vaccine (Pfizer, Moderna and AstraZeneca) and diluent (Pfizer only) inventory Lot#

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that is received. A specific Lot# of inventory can be managed by one Authorized Organization and “linked” to multiple Vaccine Events. As doses are administered during the Vaccination Event, doses will automatically decrement from the associated Inventory record.

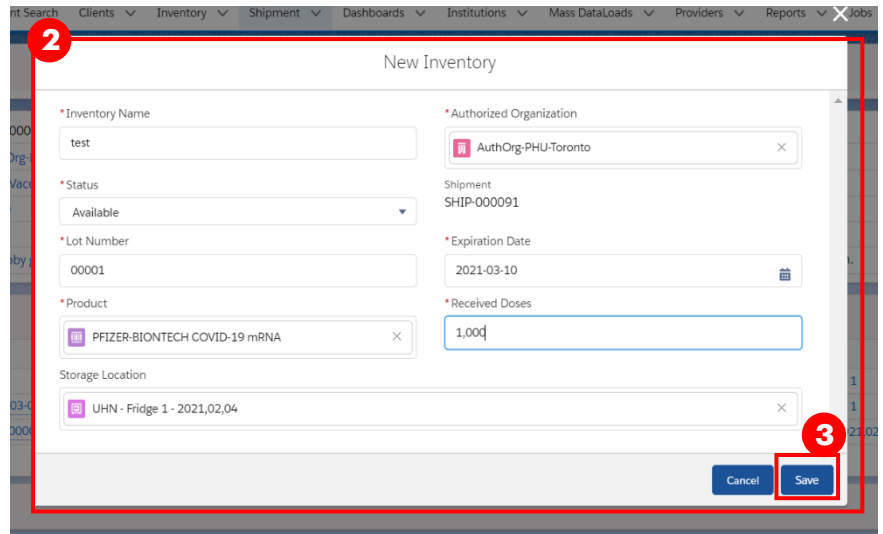
1. After creating a shipment, you will be brought to the shipment’s page. Select **“New Inventory”**



2. Fill in new inventory details:

- **Inventory Name***: Input “test” and continue filling out the other fields. Once saved, this will auto populate with the below naming convention:
[Product Trade Name] [Dosage] [Unit of Measure] - [Lot# Number] [Expiry Date]
Product Trade Name, Dosage and Unit of Measure are pulled from the product record associated with the inventory. Lot# Number and Expiry Date are pulled from the fields on the inventory record.
- **Status***: Select “Available” so the quantity of doses will be reflected as available in inventory.
- **Lot# Number***: Associated with the batch of inventory for traceability
- **Product**: Will be pre-populated based on the shipment (i.e. “PFIZER-BIONTECH COVID-19 mRNA”, or “MODERNA COVID-19 Mrna-1273” or “ASTRAZENECA COVID-19 Non-rep VV”)
- **Storage Location**: Use this search field to identify the storage locations already created in the system (for example: Lakeridge Health Oshawa – Freezer 1). If the accurate storage location does not exist in the system, refer to **Section 7: Creating Storage Locations**.
- **Authorized Organization***: Use the search bar to indicate the Authorized Organization (AO) the inventory is associated to (the party responsible for managing the inventory). This is a mandatory field and must match the AO on the User profile
- **Shipment**: Will be pre-populated with the shipment record associated to the inventory record being created
- **Received Doses***: Input the quantity of doses associated with this Lot# (same quantity as on the shipment record)
- **Expiration Date***: YYYY-MM-DD

3. Select **“Save”**



Key Tips

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- Within the inventory record, there is an option to add files to the record (for example, a temperature logger). This can be done from the right side of the inventory record by selecting "Upload Files"
- The **"Total Doses"** field is not editable (read-only) and is calculated based on **"Received Doses"** plus or minus any form of adjustments. For any future adjustments made to the inventory record (for example, extra doses from vial, no consent doses, etc.), the "Received Doses" will modify the "Total Doses" field automatically.

Received Doses	1,000
Parent Inventory	
Authorized Organization	AuthOrg-PHU-Toronto
> Location	
▼ Dose Information	
Total Doses	1,000
Extra Doses from Vial	



3. (Part I) Linking Inventory Records to Vaccine Events

Description:

Super Users or Inventory Managers will link existing inventory records at the Authorized Organization to a Vaccination Event. This allows Vaccinators to be able to select the inventory when administering doses to clients. The act of linking existing inventory records creates a **"Vaccine Event Inventory"** record associated to a specific Vaccination Event, which is tied to a main Authorized Organization inventory record. As doses are administered, it decrements from the "Doses Available" from the linked Authorized Organization inventory record. One Authorized Organization inventory record can be associated to multiple Vaccination Events.

How:

1. Allocate Inventory to the Vaccination Event by going to the desired AO Inventory Record, and selecting the arrow next to **"Vaccine Event Inventories"**
 2. Select **"New"**
 3. Enter in the details for the inventory being used at the VE:
- **Vaccination Event Inventory Name:** Input **"test"** and continue filling out the other fields. Once saved, this field will auto populate with the same name as the linked AO inventory in the "VE Inventory Name" field on this screen to ensure consistency
 - **Vaccination Event:** Search and select the Vaccine Event to link the inventory to
 - **Inventory:** This field will be auto populated with the inventory associated to that AO (this is the "linked" inventory that will be decremented as doses are administered)
 - **Allocated Doses:** enter the # of doses being allocated to that Vaccination Event

The screenshot shows the 'Vaccine Management' interface. On the right side, there is a dropdown menu labeled 'Vaccine Event Inventories (0)'. A red box highlights this dropdown menu. The main content area shows details for a vaccine inventory record for 'PFIZER-BIONTECH COVID-19 mRNA 0.3 ml - 00001, 2021-03-05'.

The screenshot shows the 'New Vaccine Event Inventory' form. The 'VE Inventory Name' field is highlighted with a red box and contains the text 'test'. The 'Inventory' field is also highlighted with a red box and contains the text 'PFIZER-BIONTECH COVID-19 mRNA 0.3 ml - 00001, 2021-03-05'. The form includes fields for 'Vaccination Event', 'Inventory Type', 'Doses Wasted', 'Allocated Doses', 'Status', 'Inventory Product', 'Inventory Status', 'No Consent for Data Collection Doses', and 'Extra Doses From Vial'.

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Key Tips

- Other fields such as “Doses Wasted”, “Extra Doses from Vial”, “Non-consent Doses”, are not used initially when linking inventory. These fields are part of VE operations and are described below.
- Users may link the same AO inventory record to multiple Vaccine Events by allocating various amounts of that Lot # across various VEs, as long as it remains within the same Authorized Organization.
- Provided that the inventory remains within the same Authorized Organization, Users are no longer required to transfer, accept, or reject inventory, instead they are able to link AO inventory to associated VEs. Transfers may still occur between AOs, refer to **Section 5 “Transferring Inventory”** and **6 “Accepting Inventory”** below for more details on this.



3. (Part II) Modifying Vaccine Events (VE)

Description:

On a regular basis, staff at a Vaccination Event (VE) (Super Users/Inventory Managers) should track and update the VE inventory from the inventory tab based on any wastage, adjustments, etc., that occur at the VE. The updates to the Vaccine Event Inventory do not automatically update the dose quantities on the main AO inventory record linked to the VE. The fields on the VE inventory are used for tracking doses available at the VE based on what was originally allocated to that event.

How:

1. From the **“Inventory”** tab, search and select the AO inventory record that was linked to the VE. The record can be edited by selecting the arrow and selecting **“Edit”**, or by:
2. Under “Vaccine Event Inventories” select **“View All”**
3. This will allow users to see all VE Inventory items associated to the AO inventory record
4. Locate the VE Inventory item and click the arrow on the right-hand side of the screen and click **“Edit”**
5. Edit the relevant quantities for the following fields:
 - **Doses Wasted:** enter quantity of doses wasted through clinical operations
 - **No Consent for Data Collection Doses:** Enter quantity of clients that were administered a dose outside of COVaxON (in the scenario that they did not consent to data collection in COVaxON, but still received a dose, Users must account for one less dose in the system’s inventory)
 - **Extra Doses from Vial:** enter quantity of extra doses achieved from vials
6. Select **“Save”**

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- The “Extra Doses from Vial” & “No Consent for Data Collection” fields are cumulative. Each day as new quantities are identified; Users must add them to the existing quantity in the field.
- Inventory managers who own the main AO Inventory record will be provided with the relevant information so they can reconcile the adjustments from the Vaccine Event Inventory to the main AO Inventory. **See Section 4 “Reconciling the AO Inventory based on Activities at the VE”** for these instructions.



4. Reconciling the Authorized Org. Inventory based on Activities at the Vaccination Event

The Inventory Manager/Super User at the Authorized Organization should reconcile the AO inventory regularly based on any updates made to the VE inventory described above (i.e., wastage, adjustments, etc.).

For the Inventory Manager at the AO to make the required adjustments, they will require information from the staff at the VE outside of COVaxON. This could include a screenshot of the VE inventory (to the right) with the quantities they’ve updated, including additional details related to those adjustments. For example, along with the screenshot, notes such as “5 doses wasted due to broken vials on 2.25.2021”.

The AO Inventory Manager can select “View All” under “Event Inventories” from the Vaccination Event to see a full list of inventory records and adjustment quantities entered.

They can select a specific inventory and under the “Related Tab” of that Vaccine Event Inventory, they can see a history of all changes made to the inventory

The screenshot displays the 'Vaccine Event Inventory' interface. At the top, it shows 'MODERNA COVID-19 mRNA 0.5 ml - EH1234, 2021-06-30'. Below this, a 'Details' tab is active, showing fields for 'VE Inventory Name', 'Vaccination Event', 'Inventory Type', 'Doses Wasted', 'Allocated Doses', 'Inventory', 'Status', 'Inventory Product', 'Inventory Status', 'No Consent for Data Collection Doses', 'Extra Doses From Vial', 'Created By', and 'Last Modified By'. A 'View All' button is visible under the 'Event Inventories' section. Below that, a table lists 'Event Inventories' with columns for 'Event Inventory: Vaccine Event Inventory Name', 'Allocated Doses', 'Extra Doses From Vial', and 'No Consent for Data Collection Doses'. A specific inventory is highlighted: 'PFIZER-BIONTECH COVID-19 mRNA 0.3 ml - EK9999, 2023-02-14'. Below this, an 'Event Inventory History' table shows a list of changes with columns for 'Date', 'Field', 'User', 'Original Value', and 'New Value'. A red box highlights a row where 'Doses Wasted' was updated from 0 to 5 by 'natalie rydell' on '2021-02-24, 7:44 AM'.

Related	Details
VE Inventory Name	MODERNA COVID-19 mRNA 0.5 ml - EH1234, 2021-06-30
Vaccination Event	Lakeridge Health - Alexis Lodge Retirement Residence - Oshawa
Inventory Type	Vaccine
Doses Wasted	
Allocated Doses	5,000
Inventory	MODERNA COVID-19 mRNA 0.5 ml - EH1234, 2021-06-30
Status	Active
Inventory Product	MODERNA COVID-19 mRNA
Inventory Status	Available
No Consent for Data Collection Doses	
Extra Doses From Vial	
Created By	ellen hall, 2021-03-02, 6:18 p.m.
Last Modified By	ellen hall, 2021-03-02, 6:18 p.m.

Event Inventory: Vaccine Event Inventory Name	Allocated Doses	Extra Doses From Vial	No Consent for Data Collection Doses
PFIZER-BIONTECH COVID-19 mRNA 0.3 ml - EK9999, 2023-02-14	1,000		

Event Inventory: Vaccine Event Inventory Name	Allocated Doses	Extra Doses From Vial	No Consent for Data Collection Doses	Doses Wasted
1 PFIZER-BIONTECH COVID-19 mRNA 0.3 ml - EK9999, 2023-02-14	1,000			5

Date	Field	User	Original Value	New Value
1 2021-02-24, 7:44 AM	Doses Wasted	natalie rydell		5

MODULE 2: AD HOC INVENTORY ACTIVITIES

The following activities should be completed on an as-needed basis by Inventory Managers.



5. Transferring Inventory

Description:

Any transfers conducted within COVaxON would be only if inventory is transferred from one Authorized Organization to another Authorized Organization (for example, a transfer of ownership between two PHUs). Each pharmacy is considered to be an individual AO. Doses will not be transferred between Pharmacies.

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Target Role: Inventory Manager

How:

1. Select the **"Inventory"** tab
2. Ensure **"All Inventory"** is selected from the dropdown.
3. If necessary, use the search bar to locate the inventory that is being transferred.
4. Click on the relevant inventory record (initial inventory record from the shipment to the Vaccine Depot; PHU or hospital).
5. From the inventory's page, select **"Transfer Doses"**
6. A pop-up will appear. Populate the **"Transferring to Authorized Organization"** field with the Authorized Organization name.
7. Populate the **"Total Number of Doses"** with the number of doses being transferred
8. Click **"Next"** & **"Next"** again & then **"Finish"**

The screenshot shows the 'Vaccine Management' interface. The 'Inventory' tab is selected. A dropdown menu shows 'All Inventory'. A search bar contains '00001'. A table lists inventory records, with one selected. The 'Transfer Doses' button is highlighted. A pop-up window titled 'Transfer Doses' is shown, with fields for 'Transferring To Authorized Organization' and 'Total Number of Doses'. The 'Next' button is highlighted.

Further Context

- To identify any inventory transfers you have sent from your AO, select the **"Pending Transfer Inventory"** list from the dropdown to view this list
- An error message will return if a user tries to transfer more doses than are available – i.e. if you have 1500 doses to at the primary location, but you try to transfer 1600 doses to another location, you will receive an error
- If Users needed to confirm the correct name of the Authorized Organization they are transferring to, refer to the Authorized Organization tab.
- If a transfer was made from a primary AO to a second AO, and doses need to be returned back to the primary AO, an inventory adjustment should be made instead of a transfer. Also, for any transfers that should not be accepted, it is recommended for the receiving site to accept the transfer, and for both sites to instead create adjustments to reflect the accurate inventory quantities instead of rejecting.
- When a transfer is completed, the "Relocated Doses" field on for the sending AO will be updated with the total amount of doses that has been relocated. The number of doses that have been transferred out of the AO will be decremented from the "Total Doses" field.

Before transferring:

Dose Information	
Total Doses	10
Extra Doses from Vial	
Doses Administered	0
Doses Pending	0
No Consent for Data Collection Doses	
Doses Returned	0
Total Doses Wasted	0
Doses Wasted due to Excursions	0
Doses Relocated	
Doses Impacted by Shipment Incident	0

After transferring 3 doses:

Dose Information	
Total Doses	7
Extra Doses from Vial	
Doses Administered	0
Doses Pending	0
No Consent for Data Collection Doses	
Doses Returned	0
Total Doses Wasted	0
Doses Wasted due to Excursions	0
Doses Relocated	3
Doses Impacted by Shipment Incident	0

COVaxON Inventory Management Job Aid

Target Role: Inventory Manager



6. Accepting Inventory

Description: The receiving Authorized Organization (AO) will need to accept the transfer

How:

1. Return to the “**Inventory**” tab
2. To identify any inventory transfers that are awaiting acceptance within your AO, select the “**Receive Pending Transfer Inventory**” list from the dropdown
3. Select the inventory item that is tagged to the receiving AO
4. From the Inventory’s page, select “**Accept Inventory**”

The screenshot displays the COVaxON Inventory Management interface. The top navigation bar includes the 'Inventory' tab, which is highlighted with a red box and a red circle labeled '1'. Below the navigation bar, the 'Receive Pending Transfer Inventory' dropdown is selected, also highlighted with a red box and a red circle labeled '2'. The main table lists inventory items, with the first item, 'COVID-19 Moderna Vaccine Mod 0.3 ml - OH564, 2022-05-31', highlighted with a red box and a red circle labeled '3'. The 'Status' column for this item shows 'Pending for Transfer'. The bottom section of the interface shows the details for the selected inventory item, 'PFIZER-BIONTECH COVID-19 mRNA 0.3 ml - 00001, 2021-03-05'. The 'Accept Inventory' button is highlighted with a red box and a red circle labeled '4'.

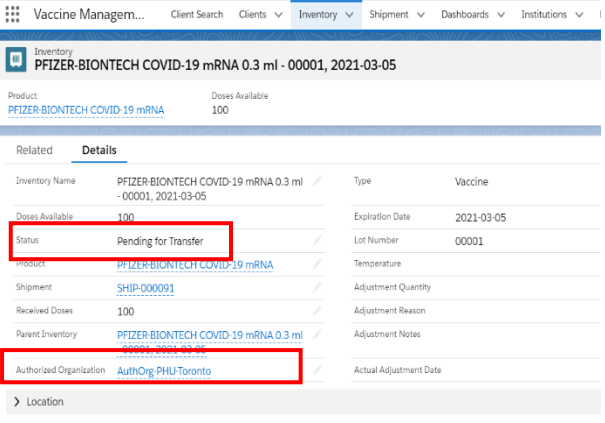
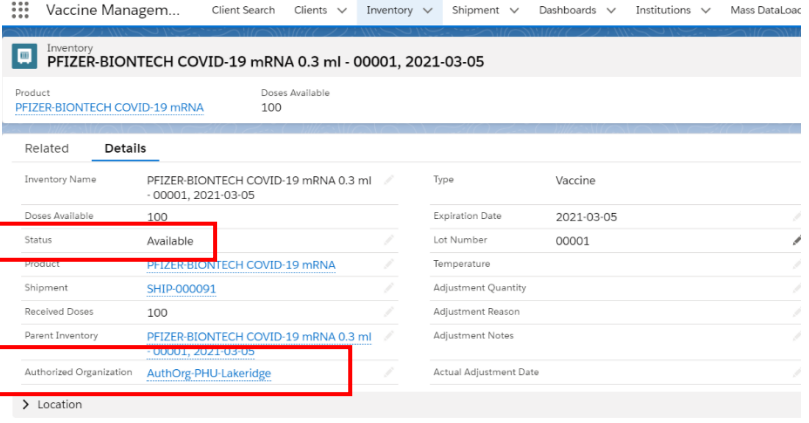
Inventory Name	Transferring To Org	Shipment	Product Trade N...	Lot Nu...	Status	Tot...
COVID-19 Moderna Vaccine Mod 0.3 ml - OH564, 2022-05-31	AuthOrg-PHU-Lakeridge	SHIP-000009	ModernaVaccine	OH564	Pending for Transfer	60
COVID-19 Moderna Vaccine Mod 0.3 ml - OH564, 2022-05-31	AuthOrg-PHU-Lakeridge	SHIP-000009	ModernaVaccine	OH564	Pending for Transfer	1

Related	Details
Inventory Name	PFIZER-BIONTECH COVID-19 mRNA 0.3 ml - 00001, 2021-03-05
Doses Available	100
Status	Pending for Transfer
Product	PFIZER-BIONTECH COVID-19 mRNA
Shipment	SHIP-000091
Received Doses	100
Parent Inventory	PFIZER-BIONTECH COVID-19 mRNA 0.3 ml - 00001, 2021-03-05
Authorized Organization	AuthOrg-PHU-Toronto

5. Click “**Next**”
6. Exit the window. To refresh the status and location of the transferred inventory, please click into the inventory’s shipment, then re-open the inventory record to see the updated status and location.

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Before Acceptance	After Acceptance
	

After Acceptance: (Note: it may take a few moments for the transfer to refresh in the system)

- The inventory's status will move from **"Pending for Transfer"** to **"Available"** for the receiving location
- The inventory's location will change from the primary location to the receiving location

Further Context

- Do not edit any fields on the received inventory record before the inventory has been accepted
 - If a user tries to update the status of the inventory from "Pending for Transfer" to "Available" before it has been properly accepted at that site, the inventory transfer will fail.
 - If a user tries to create an event (i.e. a wastage event) for the inventory before it has been properly accepted at the new site, the event will be unsuccessful
- If a transfer was made from a primary AO to a second AO, and doses need to be returned back to the primary AO, an inventory adjustment should be made instead of a transfer. Also, for any transfers that should not be accepted, it is recommended for the receiving site to accept the transfer, and for both sites to instead create adjustments to reflect the accurate inventory quantities instead of rejecting.



7. Creating Storage Locations

Description:

When populating the Storage Location field in an inventory record, if the accurate location does not exist in the system, use these steps to create a new location.

1. From the **"New Inventory"** window (shown in section 2 above), after searching, if the accurate storage location does not exist in the system, select **"New Storage Location"**. A new window will open called **"New Storage Location"**

Storage Location

Doses Relocated

1 New Storage Location

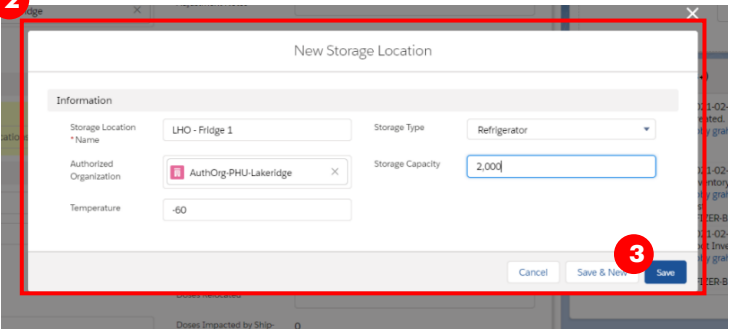
2. Fill in New Storage Location details:

- **Storage Location Name***: the naming convention may differ for each location, for example: *Toronto General Hospital – Floor 1 – Fridge 1*

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- **Authorized Organization:** Use the search bar to indicate the authorized organization the inventory is associated to (the party responsible for managing the inventory). This is a mandatory field and must be the same AO as on your User profile.
- **Storage Type:** Indicates how the inventory is being stored (select from dropdown)
- **Storage Capacity:** Number of doses that the storage location can hold (UOM: doses)
- **Temperature:** Temperature of storage location (UOM: Celsius)



3. Select **"Save"**. The storage location will be saved, and you will return to the New Inventory window. The storage location you created will be auto populated into the **"Storage Location"** field.



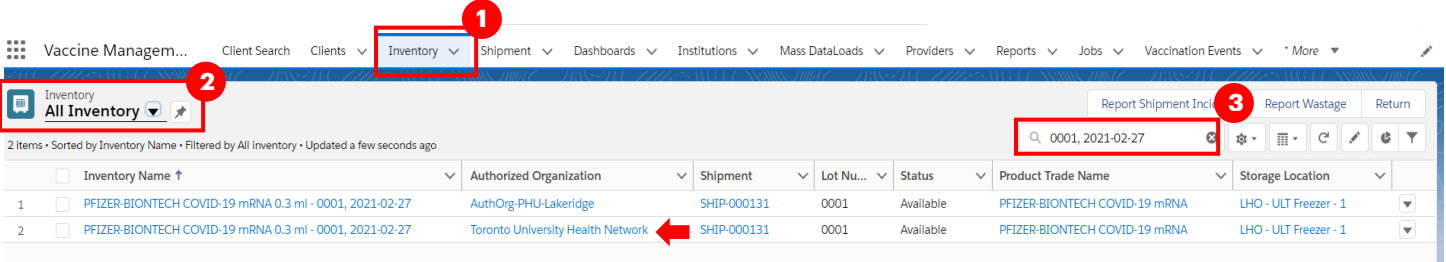
8. Recording Wastage

Description:

When wastage occurs and is identified through clinic operations, use this functionality to record wastage from the Inventory tab.

How:

1. Go to the **"Inventory"** tab
2. Select **"All Inventory"** from dropdown
3. Select the desired inventory you wish to update. You can use the search bar to aid in your search. This will bring you to a specific inventory asset's page



	Inventory Name ↑	Authorized Organization	Shipment	Lot Nu...	Status	Product Trade Name	Storage Location
1	PFIZER-BIONTECH COVID-19 mRNA 0.3 ml - 0001, 2021-02-27	AuthOrg-PHU-Lakeridge	SHIP-000131	0001	Available	PFIZER-BIONTECH COVID-19 mRNA	LHO - ULT Freezer - 1
2	PFIZER-BIONTECH COVID-19 mRNA 0.3 ml - 0001, 2021-02-27	Toronto University Health Network	SHIP-000131	0001	Available	PFIZER-BIONTECH COVID-19 mRNA	LHO - ULT Freezer - 1

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4. From the inventory's page, select the dropdown next to "**Wastage Events**" and select "**New**". 4a) **Note:** If the vaccine inventory is not in the status "Available", the user will not be able to report a wastage event. When the user goes to save the wastage event, they will receive an error message and will be unable to save the event. To proceed, the user will need to return to the inventory record and select the status "available" to log the wastage event.

5. Fill in details of Wastage Event

- **Wastage Event Name***: Follow naming convention: [Location]-[Date]-[#] ("#" would be if there are more than one wastage event on site per day)
- **Vaccine***: Use this search field to identify which vaccine you are recording wastage for
- **Reason for Wastage***: Select reason for wastage from dropdown list (new options are available)
- **Date Wastage Occurred**: YYYY-MM-DD
- **Doses Wasted**: Enter number of doses wasted
- **Entire Vaccine Inventory Wasted**: Checkbox if it was the entire Lot#

6. Select "**Save**" or "**Save & New**" for multiple wastage events

Further Context

- The amount of "Doses Wasted" entered will automatically drawdown the available inventory of that item
- There are 2 methods for recording wastage in the COVaxON system. It can also be recorded from the Inventory tab, by selecting the button on the top right called "Report Wastage". Both methods are acceptable for use.
- If a wastage event is unintentionally created, this can be rectified by editing the wastage event and changing the "Doses Wasted" number to be 0. The doses will then be added back to the "Available" doses. The wastage event will not be deleted.
- From the inventory record, the User can select "View All" under Wastage Events to see the full list of Wastage Events. The columns on this related list have been modified
- Previously Pfizer stated that there were 5 doses per vial. This has been changed to 6 and therefore if upon receiving, the quantity received was calculated at 6 doses per vial, a wastage event can be created if less than 6 doses are extracted from a vial to adjust the inventory, using the "WR - ID - Insufficient Dose(s) From a Single/Multi-Dose Vial" wastage reason

COVaxON Inventory Management Job Aid

Target Role: Inventory Manager



9. Excursion Events

Description: In the case of an Adverse Storage Condition (ASC) that impacts all or a portion of a Lot# of inventory, the "Report Excursion" functionality can be used to report the excursion event.

Note: If the vaccine inventory is not in the status "Available", the user will not be able to report an excursion event. If they try to create an excursion event, they will be unable to select the vaccine inventory and instead, they will receive the error shown. To proceed, they must return to the inventory record and set status to "Available".

How:

1. Go to the "Inventory" tab
2. Select "All Inventory" from dropdown
3. Select the desired inventory you wish to update. You can use the search bar to aid in your search. This will bring you to a specific inventory asset's page.
4. From the inventory's page, select the dropdown next to "Excursion Events" and select "New".

5. Fill in the details of the Excursion Event:

- **Storage Location:** Use search bar to indicate which storage location experienced the excursion event
- **Vaccine:** Select appropriate vaccine from search field
- **Entire Vaccine Inventory Wasted:** Use checkbox to indicate if the entire inventory record was impacted by the excursion. **Do not fill in until after the inspection**
- **Doses Wasted Due to Excursion:** This is a mandatory field. However, the PHU investigates all excursion events upon submission. Therefore, when filling in excursion event information, enter "0" wasted doses until the PHU has approved the event. Once approved, update the excursion field for the correct number of doses wasted.
- **Detected During Annual Inspection:** check this box if the event was found during Annual Inspection
- **Discovery Date/Time:** when the Inventory Manager discovered the excursion

COVaxon Inventory Management Job Aid

Target Role: Inventory Manager

- **Excursion Date:** Select the calendar icon to choose the date of the excursion
- **Excursion Start Date/Time*:** time when the temperature went out of range
- **Excursion End Date/Time*:** time when the temperature was set back in range
- **Time Duration of Excursion (hours):** time elapsed between start time of excursion and end time
- **Excursion Type:** Select "Temperature"
- **Cause*:** Use the drop-down options to indicate the cause of the excursion
- **Recommended Disposition (Resolution):** Select viable or unusable from the drop down. *Do not fill in until after the inspection.*
- **Actions Taken:** Select actions from the available category and use the arrows to move the applicable actions to "Chosen" category

- **Temperature Reading (°C):** Indicate the temperature that the storage location was at during the excursion event
- **Max Temp During Excursion:** Indicate the highest temperature during excursion
- **Min Temp During Excursion:** Indicate the lowest temperature during excursion
- **Last Logged Temperature:** last temperature showing on the data log
- **Temp Last Logged Date/Time:** Enter date/time
- **Number of Times product was exposed:** Indicate # of times vaccine was exposed to excursion event (1 = first time)
- **Duration of Product Exposure:** Indicate the exposure the product has experienced by referencing the "Time duration of Excursion" and "Number of times product exposed"
- **Datalogger Recording Sent for Assessment:** Select the checkbox if you have sent the log to the PHU for inspection
- **Monitor Type (Multi select):** Select the monitor type from the available options and use the right arrow to move the applicable option(s) to "Chosen" category
- **Reported By*:** Insert Inventory Manager who discovered the excursion event
- **Contact #:** Input telephone number for the owner of this record

an
total
was

COVaxON Inventory Management Job Aid

Target Role: Inventory Manager

- Select **“Save”** or **“Save & New”** for multiple events
- Note: If any of the **mandatory fields*** are not filled in when the user attempts to save the record, the system will produce an error and they will be unable to save the excursion event.

7. Return to the inventory record and change the status to **“Suspended Vaccines”**. It should remain in this status *until the investigation of the event is complete*.

8. After/while creating the excursion event in COVaxON, contact the PHU to inform them of the excursion. The PHU will perform an inspection of the incident. While the inspection occurs, the doses remain in “quarantine”

9. The PHU will contact your hub/location with the inspection results. Once the investigation is complete, the record can be updated with the correct number of impacted doses, by editing the excursion event record

- a) Since you are unable to edit an excursion record when the status of the inventory is not in “Available” status, you must update the inventory status to “Available” before editing the excursion record.
- b) Select the drop down from the excursion event record and select **“Edit”**
 - If all inventory was impacted by excursion: check “Entire vaccine inventory wasted” check box and indicate total number of doses wasted (these doses will automatically decrement from the total inventory available) See, *0 Doses Available on inventory record with the status changing to “complete”*. Also note the *9,144 wasted*.
 - If a portion of inventory was wasted, leave check box blank and only fill in the # of doses wasted field. The doses indicated will also decrement from the total available.
 - If the entire inventory Lot# was unaffected from the excursion event, input “0 doses Wasted Due to Excursion”.

COVaxON Inventory Management Job Aid

Target Role: Inventory Manager

10. Indicate the **“Recommended Disposition”** as **viable** or **unusable** depending on if the entire inventory was impacted by the excursion or not

11. Select **“Save”** on the excursion record

12. If the investigation determines that not all the inventory was impacted, you may return to the inventory record and ensure the inventory status is set to **“Available”** for the remaining doses.



10

Recommended Disposition (Resolution)

Unusable

Further Context

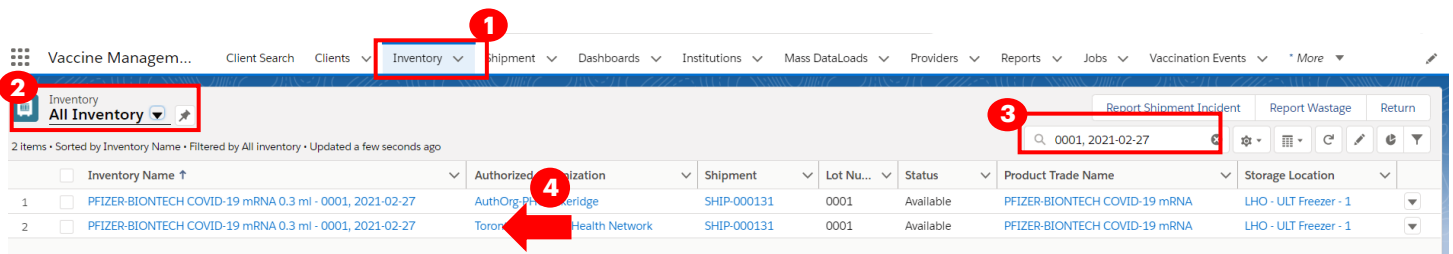
- The Excursion Event differs from the wastage record, as it is describing specifically an ASC (adverse storage condition) malfunction that occurred. Therefore, excursion events are only meant for users who are storing the inventory specifically at their location (meant for storage hubs) where the malfunction was storage related. Therefore, wastage that occurs at a clinic due to on-site error will be recorded as regular wastage events (see Section 8 above).
- If an excursion event is unintentionally created, this can be rectified by editing the excursion event and changing the “Doses Wasted due to Excursion” number to be 0. The doses will then be added back to the “Available” doses. The excursion event will not be deleted in the system and can still be viewed, however the doses wasted will be 0.



10. Adjusting Total Doses

Description:

Steps to adjust shipment and inventory records if the number of available doses requires an adjustment. For example, more doses than expected were extracted from a vial or an immunization was recorded in an offline process (ex. if a client did not consent to digital data capture).



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2

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Vaccine Managem... Client Search Clients Inventory Shipment Dashboards Institutions Mass DataLoads Providers Reports Jobs Vaccination Events More

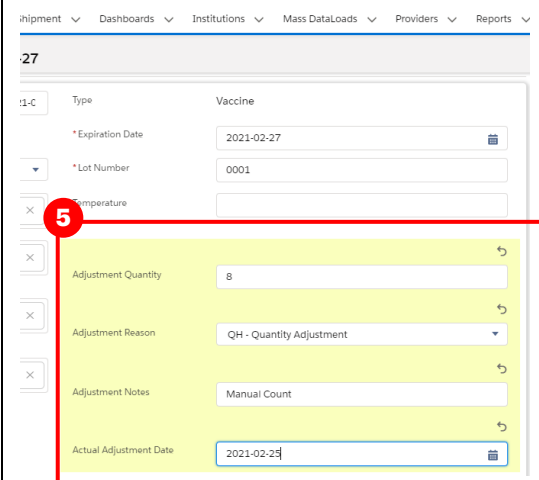
Inventory All Inventory

2 Items • Sorted by Inventory Name • Filtered by All Inventory • Updated a few seconds ago

	Inventory Name ↑	Authorized Organization	Shipment	Lot Nu...	Status	Product Trade Name	Storage Location
1	PFIZER-BIONTECH COVID-19 mRNA 0.3 ml - 0001, 2021-02-27	AuthOrg-Pfizer	SHIP-000131	0001	Available	PFIZER-BIONTECH COVID-19 mRNA	LHO - ULT Freezer - 1
2	PFIZER-BIONTECH COVID-19 mRNA 0.3 ml - 0001, 2021-02-27	Toronto Health Network	SHIP-000131	0001	Available	PFIZER-BIONTECH COVID-19 mRNA	LHO - ULT Freezer - 1

How:

- Go to the **“Inventory”** tab.
- Ensure **“All Inventory”** is selected from the dropdown.
- Use the search bar to identify the inventory you would like to modify.
- Select the relevant inventory and click the pencil icon to edit the following fields:
- “Adjustment Quantity”**: Enter the quantity for the adjustment and add the “Adjustment Reason”, “Adjustment Notes” and “Actual Adjustment Date”. This is used for any other discrepancies or adjustments required outside of “Extra doses from vial”, “Wastage”, and “No Consent for Data



5

Shipment Dashboards Institutions Mass DataLoads Providers Reports

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Type Vaccine

* Expiration Date 2021-02-27

* Lot Number 0001

Adjustment Quantity 8

Adjustment Reason QH - Quantity Adjustment

Adjustment Notes Manual Count

Actual Adjustment Date 2021-02-25

COVaxON Inventory Management Job Aid

Target Role: Inventory Manager

Collection Doses". For example, when performing a physical count.

6. **Extra Doses from Vial**: Enter the quantity of extra doses from vial in this field and **add the "Adjustment Reason", "Adjustment Notes" and "Actual Adjustment Date"**. Note: this will only be for Lot#'s of inventory received at 5 doses per vial. Since Pfizer has updated the vials to be 6 doses per vial, any shipments that had quantity received at 6 doses per vial will not require this adjustment.

7. **"No Consent for Data Collection Doses"**: Enter the quantity of clients that did not consent to data collection and were therefore have no dose administration tracked in COVaxON, and add the "Adjustment Reason", "Adjustment Notes" and "Actual Adjustment Date".

8. Select **"Save"**

The screenshot shows the 'Inventory Adjustment' form for 'PFIZER-BIONTECH COVID-19 mRNA 0.3 ml - 0001, 2021-02-27'. The form is divided into several sections: 'Location', 'Dose Information', 'Recall Information', and 'Adjustment Notes'. The 'Dose Information' section contains fields for 'Doses' (400), 'Extra Doses from Vial' (2), 'Doses Administered' (0), 'Doses Pending' (0), 'Doses Returned' (0), 'Total Doses Wasted' (10), 'Doses Wasted due to Excesses' (0), and 'Doses Relocated' (0). The 'Recall Information' section includes 'Recall Source' (None), 'Recall Date', and 'Recall Date'. The 'Adjustment Notes' section has a 'Manual Count' field and an 'Actual Adjustment Date' field (2021-02-25). Red boxes and numbers 6, 7, and 8 highlight the 'Extra Doses from Vial' field, the 'No Consent for Data Collection Doses' field, and the 'Save' button respectively.

Further Context

- Users cannot modify the "Total Doses" field. Any updates to the "Adjustment Quantity", "Extra Dose from Vial" and "No Consent for Data Collection" will update the "Total Doses" field appropriately
- If a transfer was made from a primary AO to a second AO, and doses need to be returned back to the primary AO, an inventory adjustment should be made instead of a transfer.
- When the inventory hits 0 doses available from being administered, the Inventory status will change to "Completed". If the Inventory Manager has not adjusted the total doses available for that inventory based on additional doses achieved from a vial, there will be a misalignment between actual doses and the doses in the system. To avoid delays to administering doses, the Inventory Manager will need to change the status of the inventory to "Available". Inventory can be administered if the doses available is 0 or negative, as long as the status is "Available"

11. Managing Recalls

Description:

There are offline processes for communicating and handling vaccine recalls:

- Manufacturer determines there is an issue with a specified Lot# and notifies Health Canada
- Health Canada identifies that a specified Lot # had a higher AEFI rate than expected and notifies the provinces/territories
- Public Health Ontario may identify the source of an issue with a specified Lot# and notify the ministry

Health Canada (and often the manufacturer as well) will perform an investigation of the impacted Lot# and communicate to the Users (outside of the system) on next steps. **This is in the form of**

notice/recommendations to the province on further use of the vaccine (i.e. discard or lift the suspension so the vaccines can be distributed for use). Public Health Ontario/the Ministry of Health would inform the public health units who then inform their local providers (Users of COVaxON). Each location has a **designated User (Super User or Inventory Manager)** responsible for reflecting recalls within COVaxON. The User will follow the below process to recall the vaccine within the COVaxON system. For any locations where the impacted Lot# was transferred, the inventory will need to be physically returned to the PHU and the User must recall all impacted inventory records.

COVaxON Inventory Management Job Aid

Target Role: Inventory Manager

How:

1. Identify the impacted Lot # that has been recalled and all the associated inventory records (including any quantity that was transferred as a new inventory record from that Lot# to another location).
2. On the inventory page go to the right-hand corner and select **"Recall Inventory"** button
3. Recall Source: Select "Health Canada"
Recall Date: Select the date that the Lot# was recalled.
4. Select **"Next"** and **"Finish"** the inventory recall record will then save, once saved the following occurs:
 - a. Available doses will now be set to 0 and in turn, the "Number of Doses Recalled" will be automatically populated with the previous number of doses that were available.
 - b. Inventory status will automatically update to "Recalled"

In the example to the right, the inventory record was comprised of 100 doses. Before the recall, 0 doses had been administered. Once the product was recalled, the inventory status changed to "Recalled" and the Doses Available became "0".

Based on the "Recalled" status, vaccinators are now unable to administer additional doses, and under the Dose Information and Recall Information, users can see "100 Total Doses, 0 Doses Administered and 100 Doses recalled".

The screenshots illustrate the Recall Inventory process in the COVaxON system. The first screenshot shows the 'Recall Inventory' modal form with fields for 'Recall Source' (set to 'Manufacturer') and 'Recall Date' (set to 'Jan 26, 2021'). A red box highlights the 'Recall Inventory' button in the top right corner of the inventory page. The second screenshot shows the 'Recall Inventory' modal form with the 'Next' button highlighted. The third screenshot shows the 'Recall Inventory' modal form with the 'Finish' button highlighted. The fourth screenshot shows the inventory record for 'PFIZER-BIONTECH COVID-19 mRNA 0.3 ml - 0001, 2021-02-27'. The 'Doses Available' field is set to 0, and the 'Status' is 'Recalled'. The fifth screenshot shows the 'Dose Information' and 'Recall Information' sections, which display the following data:

Dose Information	
Total Doses	100
Extra Doses from Vial	
Doses Administered	0
Doses Pending	0
No Consent for Data Collection Doses	
Recall Information	
Recall Source	Health Canada
Doses Recalled	100
Recall Date	2021-01-26
Rescind Date	

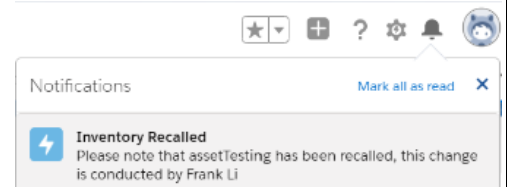
Further Context

- When a recall is saved, the Lot#s that were recalled will be frozen, rendering the inventory unavailable for administration. Functions such as reporting wastage, reporting excursion, returning and transferring doses will also be unavailable. Therefore, the only way to "unfreeze" inventory is for a Super User to rescind a recall event (see section 12 below).

COVaxON Inventory Management Job Aid

Target Role: Inventory Manager

- When inventory status is set to "Recalled" it cannot be changed. Only Super Users will be able to update the status. Inventory Managers will be unable to edit a record once the status is changed from "Available" to "Recalled".
- When the status of an inventory record is changed to "Recalled", a notification will be sent to Super Users, Inventory Managers, Vaccinators, and Site Staff (not Users with the Dashboard profile) in COVaxON so they are aware the recall has been reflected in the system and can no longer select that Lot# or administer doses. The notification is in the format of: "Please note that [inventory name] has been recalled, this change is conducted by [last modified by]"
- When inventory is in the "Recalled" status, and additional doses are added to the "Total Doses" field, those additional doses will be added to the "Recalled Doses" quantity and will not be listed as available inventory. This can only be done by a Super User. The "Doses Available" inventory amount will continue to reflect 0 doses. This is to accommodate a situation whereby the number of total doses requires adjustment after a recall has already been made against that inventory. For example, when reconciling additional doses obtained from a vial, or other forms of inventory adjustments. Please note that the best practice is to conduct adjustments on a regular basis, to avoid any lag in the inventory quantities reflecting the accurate inventory quantities.



12. Rescinding a Recall

Description:

The PHU will advise of the outcome of the investigation and provide recommendations on the use of the vaccine (communication is done outside of COVaxON). If the investigation is determined to have no safety issues, they will notify all Users that the Lot# can be redistributed for administration (and the previous recall can be rescinded). Only Super Users have the ability to rescind an inventory recall.

How:

- Change the status of the Inventory record from "Recalled" to "Available"
- Enter the rescind date in the "Recall Information" section of the inventory page.
- Select Save
- The doses that were initially recalled will now be associated to "Doses Available"

Note: If an inventory is recalled multiple times, users would need to remove the rescind date before recalling the inventory again. The history of the initial recall and rescind are captured in the Inventory History.

At end of shift, log out of COVaxON and clear the browser cache. Refer to the Login, Logout, User Settings job aid for detailed steps. Sanitize shared devices in accordance with location protocols.