MyPeST Application

USER GUIDE

Version 3.1
8/6/2026

MyPeST Support

OPPSupport@epa.gov

When submitting a support request, please include: your name and organization, a brief description of the issue, the action you were trying to complete, any error message received, and a screenshot if available.

DO NOT DOWNLOAD THIS USER GUIDE

This User Guide is regularly updated. Returning to the MyPeST application and viewing the posted user guide ensures you always have the most recent version.

Creating A MyPeST Account

In previous versions of the MyPeST application, your MyPeST account was created and managed through Salesforce. With this update, your MyPeST account is now created and managed through CDX and Login.gov. This change allows MyPeST users to submit new product registrations or update existing product applications directly through MyPeST. You will no longer need to log in to both MyPeST and CDX to view the status of your submissions and create new or update existing applications. MyPeST can do it all. The documents you upload are securely stored and CROMERR compliant. If you already have CDX and Login.gov accounts, you can use those existing accounts to log in to MyPeST. If you have not registered with CDX and Login.gov, please refer to these websites if you need help creating your account:

For help with CDX

888-890-1995

cdx.epa.gov/FAQ#CDX

For help with Login.gov

login.gov/help

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Version History

VERSION	AUTHOR	DATE	CHANGES MADE
1.0	Datawiz Team	12/06/24	Created draft
1.1	Datawiz Team	2/13/25	Updated Screenshots for images 1, 2, 5, 6, 8
1.2	Datawiz Team	3/31/2025	Removed information about Creating an Account, Logging In, and Password Recovery. Added information about the Dashboard (p.11), Dashboard Tiles(p.12-13), the Applications Tab (p. 14), and the Case Details page (p. 15-18).
1.3	Datawiz Team	4/23/2025	Added Appendix B—Subject Codes
2.0	Datawiz Team	8/11/2025	Added information about the Dashboard (p.11) Registration Review Cases Tile (p.12) My DCIs tile (p.13) Products Tab (p.16) Updated information about using Correspondence (p.20) Updated Role Management Deny and Revoke instructions (p.22).
2.1	Datawiz Team	10/2/2025	Added information on the Products Tab (p. 16) How to Certify Bilingual Labeling (p. 17-20)
2.2	EPA	10/17/2025	Updates to the Products Tab (p.16) Changed section title from How to Certify Bilingual Labeling to How Do I Certify That Bilingual Labeling Was Added Or That A Product Is Not To Be Released For Shipment? and updated the instructions (p.18) Added How Do I Provide EPA with Links to Bilingual Labeling? section (p.21) Added a new Appendix B—How to Certify PRIA 5 Bilingual Labeling Compliance (p.29) Changed Subject Codes to Appendix C (p.32)
2.2.1	EPA	10/21/2025	Moved the information from Appendix B into the Bilingual Labeling Added section. Changed Appendix C back to Appendix B.
3.0	BIAC	6/29/2026	Updated log in procedures (p.2) Updated the Purpose (p.6) Updated the Home Page section (p.7) Updated the role descriptions for Company Administrator, Company Representative, and Consultant (p.8) Added Make a Submission (p.32) Added Start New Submission (p.33) Added Update Existing Application (p.36) Added Submit Multiple Applications (p.38) Added Appendix C—Troubleshooting Company Numbers in MyPeST (p.42) Added Appendix D—Troubleshooting Role Requests in CDX (p.46) Added Appendix E—Action Codes (p.49)
3.0.1	BIAC	7/8/2026	Updated compliance deadlines for bilingual labeling requirements (p.21)
3.0.2	BIAC	7/20/2026	Updated Appenix E—Action Codes list (p.54-72)
3.1	BIAC	8/4/2026	Added Respond to a Data Call-In (DCI) (p.40—43)

Purpose

This user guide provides clear instructions for using the MyPeST application for specific functions based on roles. The MyPeST app is a web-based system that tracks your organization's pesticide applications and products. This user guide will cover the following:

- Selecting an organization and requesting a role
- Viewing features based on roles
- The Dashboard and Dashboard Tiles
- The Case Details page
- Correspondence
- Company Administrator responsibilities in the Role Management Tab (Accepting and denying role requests, revoking user access, inviting users to a role.)
- Creating a New Submission and Updating an Existing Application functionality

Scope

This document is meant for all users but is primarily used by Company Admins (CA) as they have full rights to edit and invite users/create user accounts.

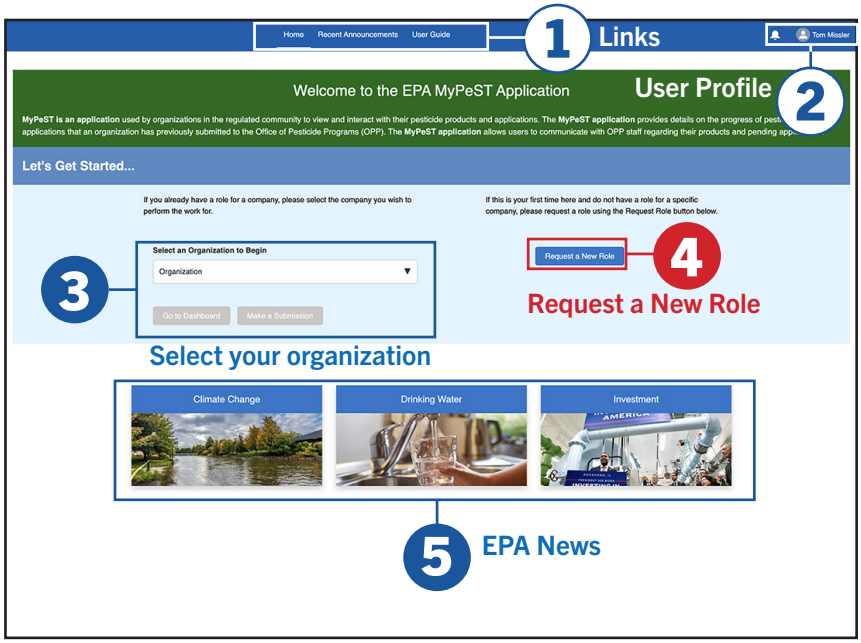
The Home Page

Once you have successfully logged in to My Pesticide e-Submission Tracker (MyPeST), you will see the Home Page.

1. At the top of every page you will see three links: **Home, Recent Announcements,** and **User Guide.**
   - **Home** will take you back to this page
   - **Recent Announcements** will take you to the Pesticides page on EPA.gov
   - **User Guide** will open the latest version of the user guide you are currently reading
2. **User Profile** has a drop-down with the option to view your user profile or log out of the MyPeST application.
3. **Select an Organization to Begin** is a drop-down list of all the organizations with whom you have a role. When you log in for the first time, this is grayed out because you have not requested a role with an organization. Once you have a role, that organization will appear in this drop-down menu.

   The **Go to Dashboard** and **Make a Submission** buttons are grayed out until you select your organization. Clicking the **Go to Dashboard** button will take you to the Dashboard that will give you an overview of all your organization's applications with the EPA. Clicking the **Make a Submission** button will allow you to create a new submission to register a new product or submit documents for an exisiting registration.
4. Click the **Request a New Role** button to start the process of selecting a role with an organization. A role with an organization is required to use the MyPeST app. If this is your first time using the MyPeST app you will need to start here.
5. EPA News provides links to recent articles published by the EPA.

The Home Page screenshot



Roles and Organizations

Creating an account in MyPeST is the first step that gives you access to the application. To see any data, you need to request a role with an organization. There are four different roles: Company Administrator (CA), Company Representative (CR), Consultant, and Contributor (Future). Each role has different responsibilities and levels of access to data which are explained below.

You can have one role at multiple organizations, but you cannot have multiple roles for a single organization. You can have different role types at different organizations—just because you are a Consultant with one organization doesn't mean you can only request Consultant roles. It is up to the organization to approve or deny your role request. So you could be a CA for five organizations, a CR for three others, a Consultant for six others, and never be a Contributor (Future). However, if one of the six organizations where you are a Consultant wants you to become a Company Admin, you can not just request a new role. The one role per organization rule will not let you request a second role. The workaround is explained in Appendix A.

Role Types

There are four roles that a user can request within the MyPeST App:

COMPANY ADMINISTRATOR (CA)

The CA is the highest level in the hierarchy of user roles within the MyPeST App and gives the user the highest level of access to data. CAs can view the status of all current projects and can access product history to view information about previously submitted projects. They can start new submissions and update any existing application. CAs are the only users allowed to invite new users, approve, or deny user requests, and revoke the access of a user. Every organization must have at least one user designated as the CA.

An organization can have multiple CAs. The first CA for an organization is approved directly by the EPA. This process is explained in the next section. No other role can be assigned within an organization until the first CA is approved. After the first CA is approved, that person will approve all other roles including other CA roles.

COMPANY REPRESENTATIVE (CR)

The CR has the same level of data access as a CA. They can start new submissions and update any existing application. However, they are not able to invite users, approve or deny role requests, or revoke user access. An organization can have multiple CRs.

CONSULTANT

Consultants can start a new submission and update an existing applications that they submitted. An organization can have multiple Consultants.

CONTRIBUTOR (FUTURE)

Potential future role.

Who Can Apply For These Roles?

The roles are open to anyone the organization is willing to approve at that level of access. An organization can grant Company Admin access to a contractor who works for a third-party organization. Likewise, a user can be given the Consultant role, even though they are a direct employee of the organization making the submission.

The EPA only approves the first Company Admin for any organization to allow more flexibility in who they appoint to work for them. If an organization wants to make everyone a Company Admin, they are able to do that. If the organization decides that only one person should have CA access, they can do that as well.

If an organization ever finds itself without a Company Admin, the CRs, Consultants, and Contributors (Future) can continue working at their current level of access with no disruption. However, until a new CA is approved, no roles can be approved, denied, or revoked. If there is no acting CA to approve them, the new person requesting the Company Admin role will need to submit a letter and follow the same process the first CA did in order to be approved. It is recommended to have at least two CAs appointed for each organization.

Requirements to be the First Company Administrator for Your Organization

In order to be the first Company Admin for your organization, you need to be approved by EPA. To get get that approval, you need to collect some information outside the app and submit it.

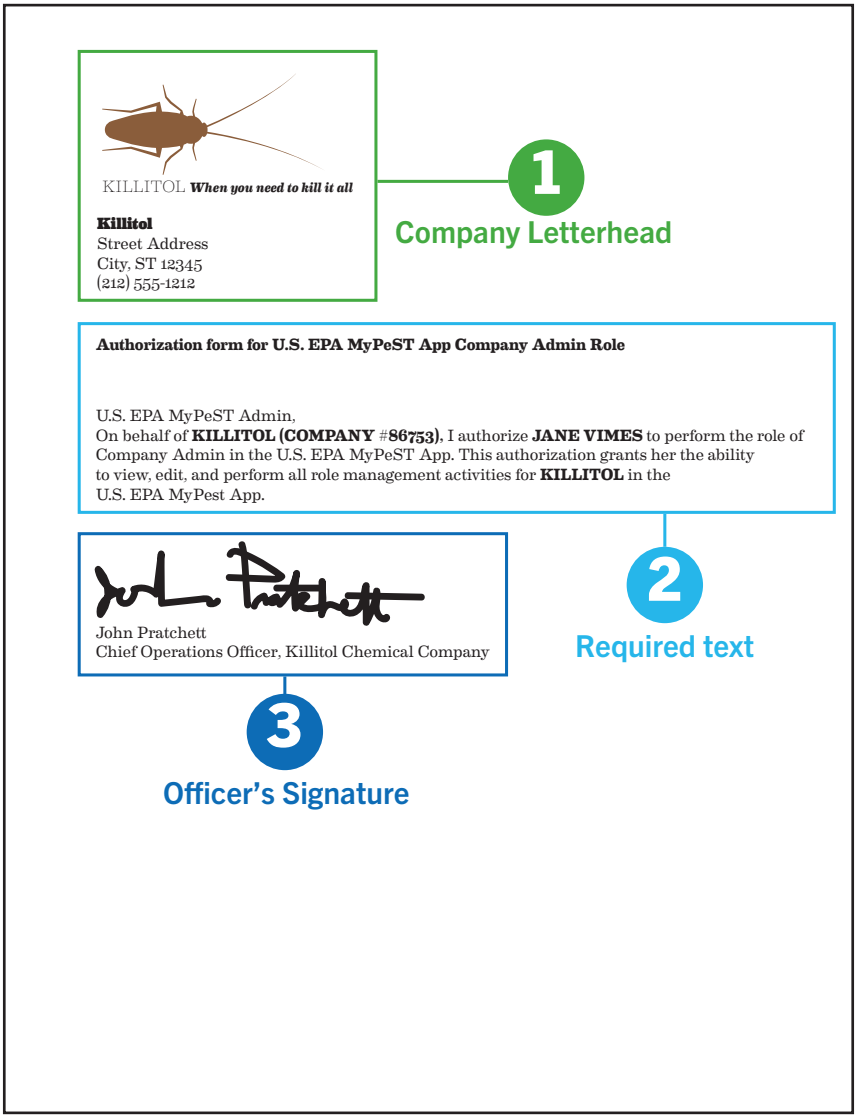
The Authorization Letter

The letter requirements are:

1. The letter must be on the Company's letterhead
2. Use the language from the Authorization Letter Template
3. It must be signed by an officer of the Company
4. Save the letter using the Authorization Filename Conventions: **[Company Number] [Company Name] Mypest Company Admin.pdf**

Every organization with a company number needs a signed authorization letter for each CA. If a organization has 10 different companies with the same name but different company numbers and wants to assign the same person CA for all of them, that organization will need to submit 10 letters to the EPA, one for each company number.

Authorization Letter example



IMPORTANT

You must follow the Authorization Filename Conventions

You **must** name your file as specified below:

[Company Number] [Company Name] **Mypest Company Admin.pdf**

for example:

86753 Killitol Mypest Company Admin.pdf

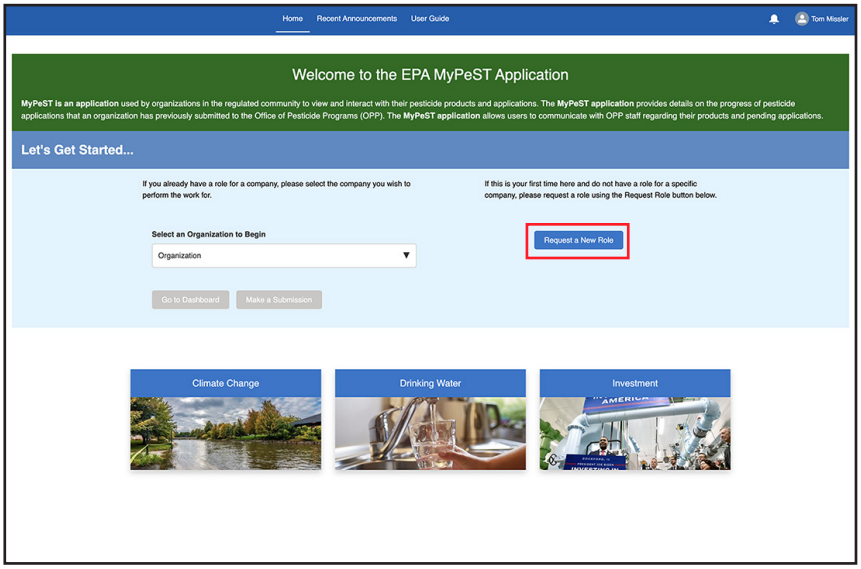
Request To Be The First Company Administrator For Your Organization

The first role your organization needs to establish is the Company Administrator (CA). If you do not have a pdf of your Authorization Letter, get that now. You cannot complete this process without it.

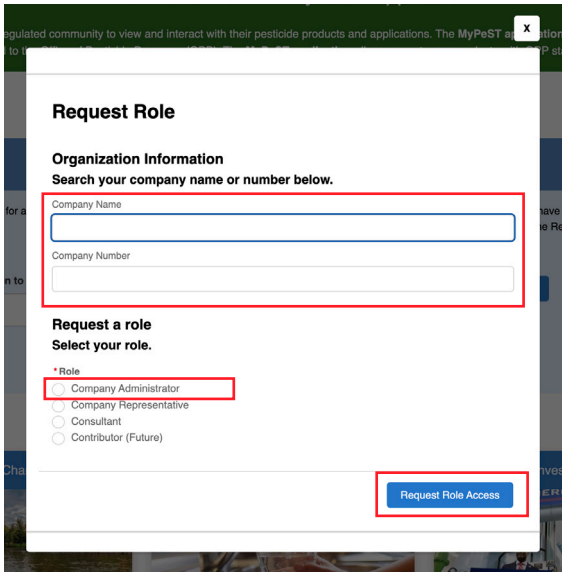
1. On the Home page, click the **Request a New Role** button.
2. The *Request Role* pop-up window appears. You can enter a few letters of the Company Name or you can enter the Company Number (or both). Then click the **radio button next to the role you want.** In this example we are applying for the Company Administrator role for the Big Bear company. Click the **Request Role Access** button.

   If your company does not appear in the Company Search results, check out ***Appendix C—Troubleshooting Company Numbers*** for instructions on how to add your company.
3. Clicking the button brings up the *Company Search* pop-up. This is where knowing the Company Number becomes very important. If the organization you are requesting a role with has multiple divisions or has acquired multiple companies, you might see a list of identical Company Names but different Company Numbers. Each company number represents a different company and they do not share information. So you need to ensure you are selecting the correct company. Once you have selected the organization, click the **Next** button.

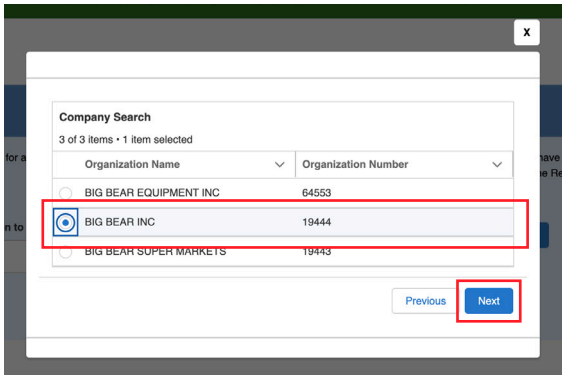
Step 1



Step 2



Step 3



4. The *Confirm Details* popup window will appear. If all the information is correct, click **Confirm Details.** If you see a mistake, you can click **Previous** to return to the *Company Search* page and select a different organization. If at any point you want to cancel your request, click the **X** in the upper right corner of the popup window.
5. Submit your authorization letter.

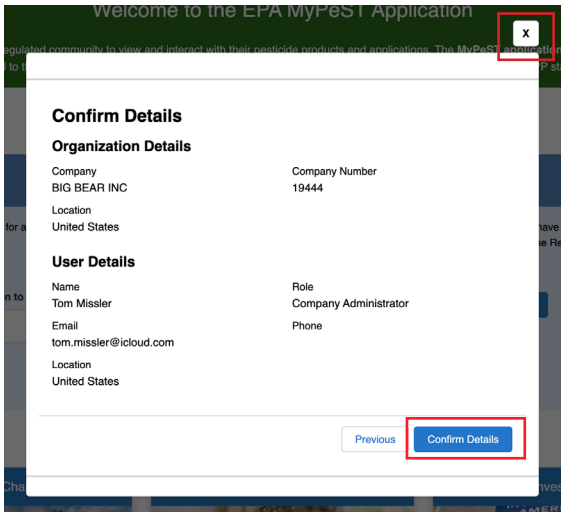
   In the *Upload authorization document* popup you can navigate to a file on your hard drive by clicking **Upload Files** or drag and drop it inside the dotted outline.
6. If you click **Upload Files,** navigate to the file on your hard drive and click **Upload.** If you used drag and drop, click **Upload.**
7. When the progress bar fills in a green check will appear. Click **Done** which will return you to the *Upload authorization document.* Click **Finish.**
8. You will now see the *Role Request Submitted* popup window. Click the **X** in the upper right corner of the popup window to close this window and return to the Home page.
9. You will receive another email from EPA once your request has been approved. The next time you log into MyPeST, the Organization drop-down will be active. Select your organization and click **Go.**

Request A Role With An Organization After the First Company Admin Has Been Approved

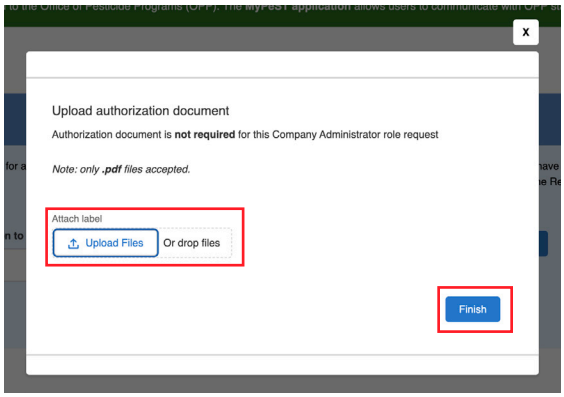
What sets the Company Admin apart from the other roles is the ability to approve all the other role requests. Now that the first Company Admin has been approved, that person will now approve all other role requests.

To request the role of Company Representative, Consultant, Contributor, or to be an additional Company Admin, follow steps 1–4 above. You do not need to submit a letter. If the *Upload authorization document* popup appears, just click **Finish.** Do not upload anything. You will receive an email from EPA after your request has been approved. The next time you log into MyPeST, the Organization drop-down will be active. Select your organization and click **Go.**

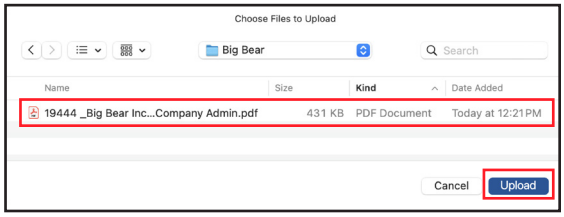
Step 4



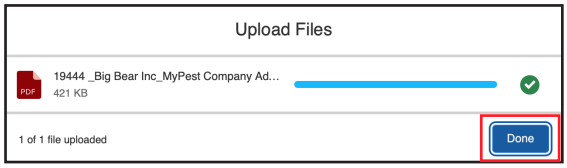
Step 5



Step 6



Step 7



Accepting or Rejecting a Role Invitation

You can be invited by an organization to accept a role they have chosen for you. If you have an invitation waiting for you, the next time you log in to MyPeST you will see the *Accept/Reject Role* popup screen before you see the Home Screen.

Accept

1. If your data is correct and you recognize the organization sending the request, select **Accept Role** from the drop-down.
2. Click **Next.**
3. You will see a popup congratulating you on your new role. Click **OK.**

When you get to the Home page, the organization will appear in your Organizations list.

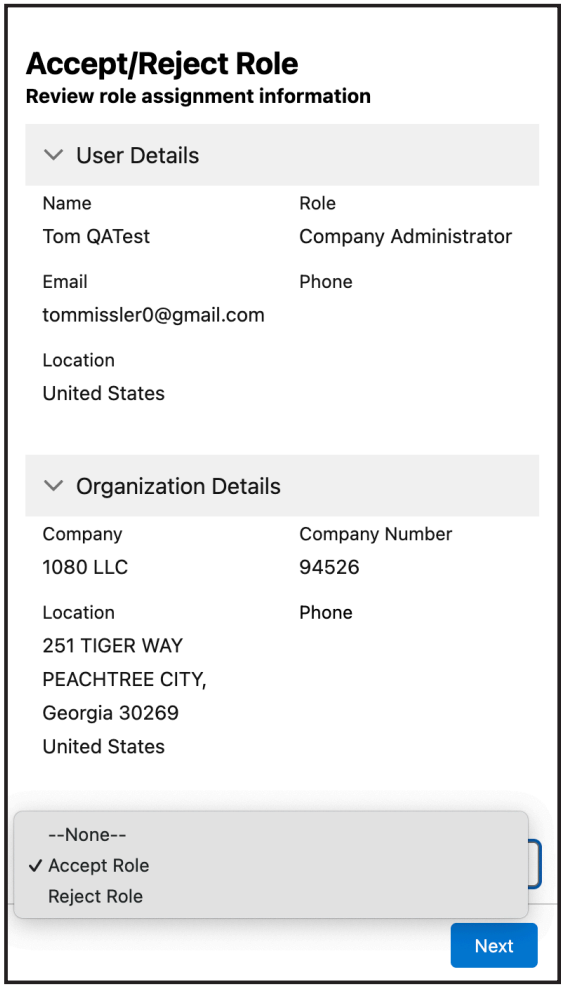
Reject

4. If your data is incorrect or you do not recognize the organization sending the request, select **Reject Role** from the drop-down.
5. Click **Next.**
6. You will see a popup asking *Are you sure you want to reject the role?* Click **Reject Role.**
7. You will see a popup indicating you rejected the role. Click **OK.**

When you get to the Home page, the organization will not appear in your Organizations list.

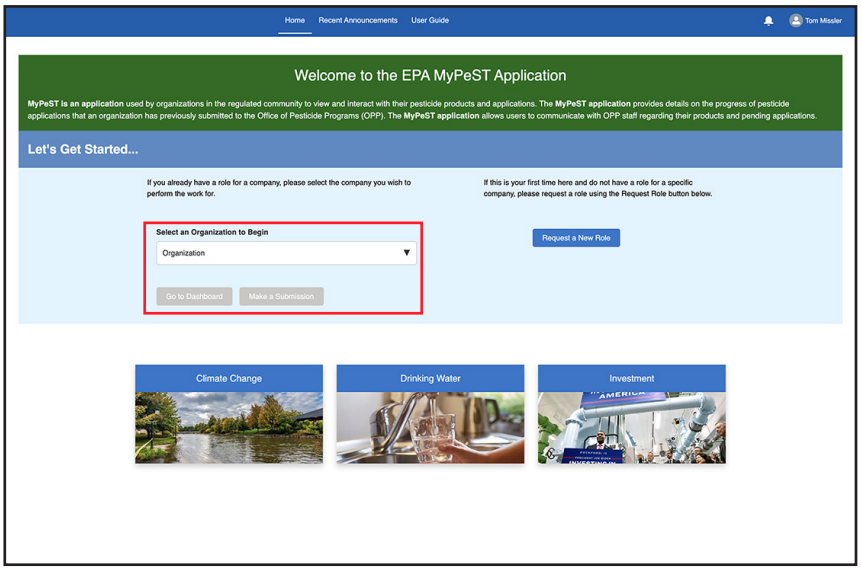
If it turns out you should have accepted the role, you can always request it or the organization can send another invitation.

Accept/Reject Role popup

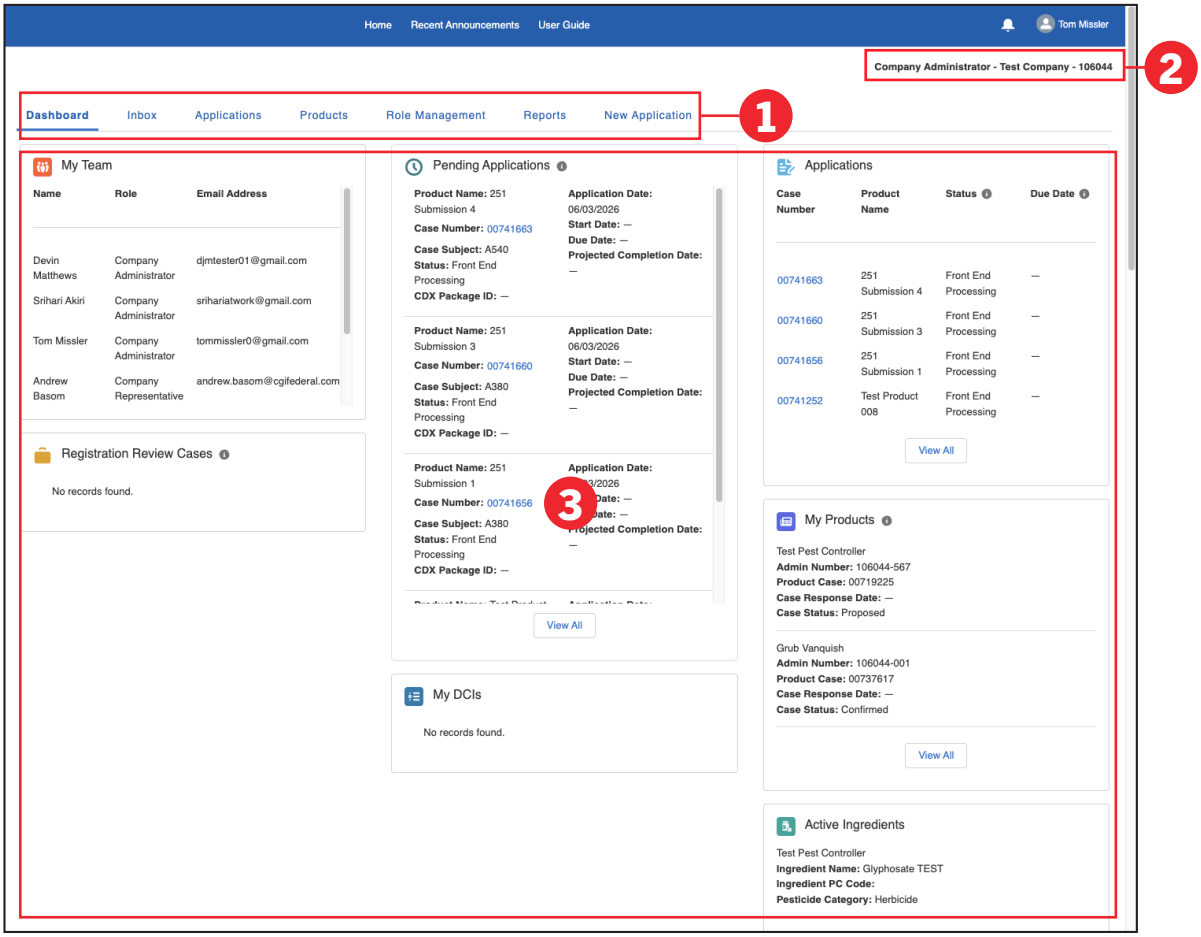


Select An Organization

After you have been accepted for a role, the next time you log in you will see the **Select an Organization to Begin** drop-down is active on the Home screen. Click the drop-down and select your organization from the list. You can click **Go to Dashboard** to see the status of your EPA applications or you can click **Make a Submission** to start a new product registration or update an existing application. We will start with **Go to Dashboard.**



The Dashboard



You will be directed to the Dashboard for your organization after you click the **Go to Dashboard** button on the Home screen.

1. There are seven Tabs on the Dashboard:

   - **Dashboard**
   - **Inbox** (currently inactive)
   - **Applications**
   - **Products**
   - **Role Management**
   - **Reports** (currently inactive)
   - **New Application**

   The Company Administrator can see all seven tabs.

   The Company Representative can see all the tabs except Role Management.

   The Consultant can see Dashboard, Inbox, and Reports

   The Contributor will only see Dashboard and Inbox.

2. This lets you know your role, the company name and number you are currently viewing. If you would like to switch to a different company, click Home in the header and select a different company in the Organizations drop-down.

3. There are seven Tiles on the Dashboard:

   - **My Team**
   - **Registration Review Cases**
   - **Pending Applications**
   - **My DCIs**
   - **Applications**
   - **My Products**
   - **Active Ingredients**

   The Company Administrator and Company Representative can see all seven tiles and all data about all projects regardless if they are complete or in progress.

   The Consultant can see all seven tiles, but will not have any interactive links on any tile.

   Contributors can not see any tiles on the Dashboard.

On the Dashboard and Case Details page, you will see this Tool Tip icon. If you hover over the icon, you will see a popup that gives more information about the field.

Dashboard Tiles

My Team Tile

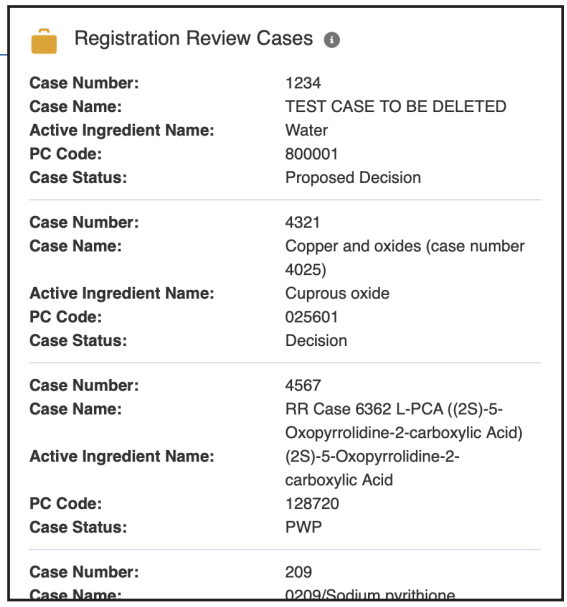
Fields displayed:

- **Name**
- **Role**
- **Email Address**

Registration Review Cases

Fields displayed:

- **Case Number**
- **Case Name**
- **Active Ingredient Name**
- **PC Code**
- **Case Status**—Indicates the stage of EPA review process the application is currently in.

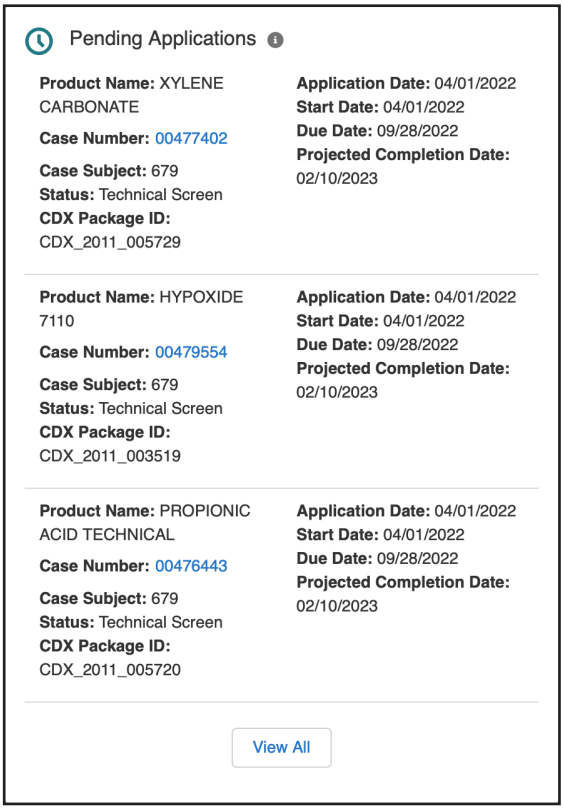


Pending Applications

Fields displayed:

- **Product Name**
- **Case Number**
- **Case Subject**
- **Status**—Indicates the stage of EPA review process the application is currently in.
- **CDX Package ID**—The CDX tracking number
- **Application Date**
- **Start Date**—The date when the action work was started on the application.
- **Due Date**—The date when the action work is to be completed before any negotiation or changes.
- **Projected Completion Date**—The date used to indicate the expected completion date and entered by EPA.

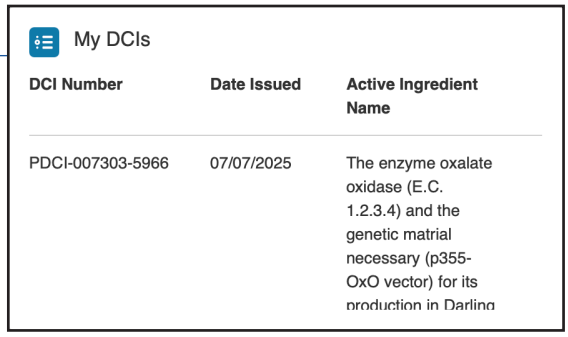
Clicking a **Case Number** link will take you to the Case Details page for that product.
Clicking **View All** will take you to the Applications Tab page.



My DCIs

Fields displayed:

- **DCI Number**
- **Date Issued**
- **Active Ingredient Name**

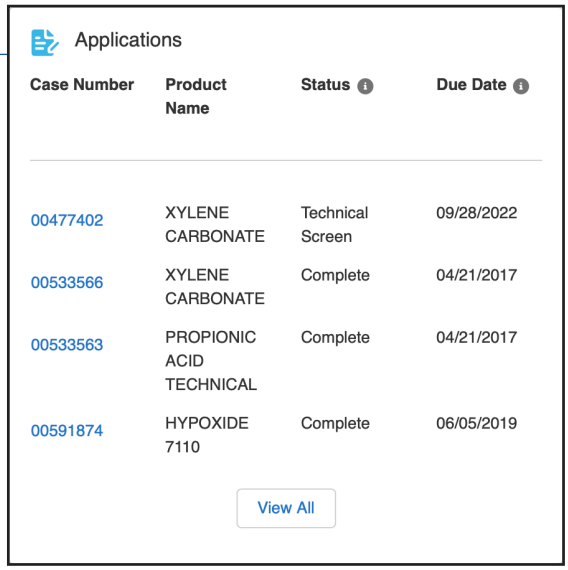


Applications

Fields displayed:

- **Case Number**
- **Product Name**
- **Status**—Indicates the stage of EPA review process the application is currently in.
- **Due Date**—The date when the action work is to be completed before any negotiation or changes.

Clicking a **Case Number** link will take you to the Case Details page for that product.
Clicking **View All** will take you to the Applications Tab page.

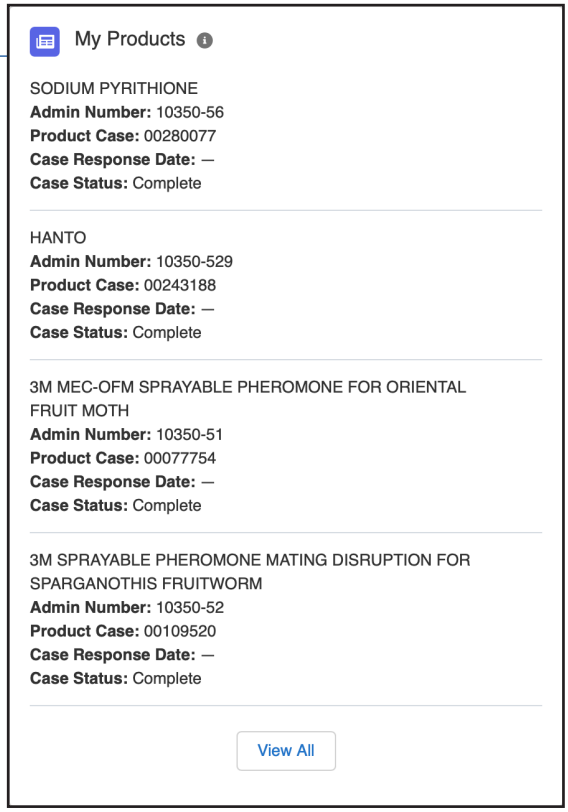


My Products

Fields displayed:

- **Product Name**
- **Admin Number**
- **Product Case**
- **Case Response Date**—the date a response code was added/updated.
- **Case Status**—Indicates the stage of EPA review process the application is currently in.

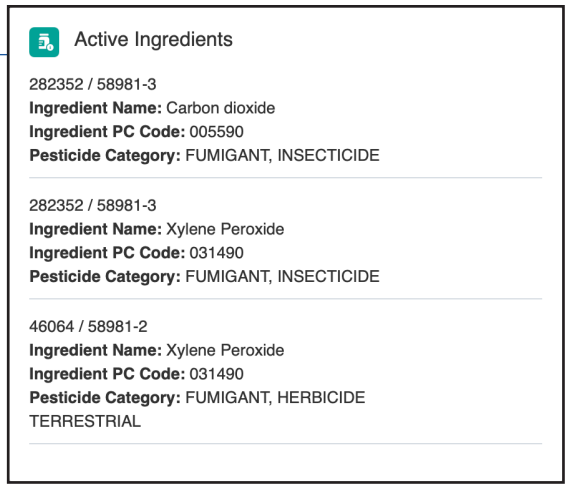
Clicking **View All** will take you to the Products Tab page.



Active Ingredients

Fields displayed:

- **PRISM Ingredient**
- **Ingredient Name**
- **Ingredient PC Code**– or Pesticide Chemical Code is a unique chemical code number assigned by the EPA to a particular pesticide active ingredient, inert ingredient or mixture of active ingredients.
- **Pesticide Category**



Pending Applications & Applications—What's the Difference?

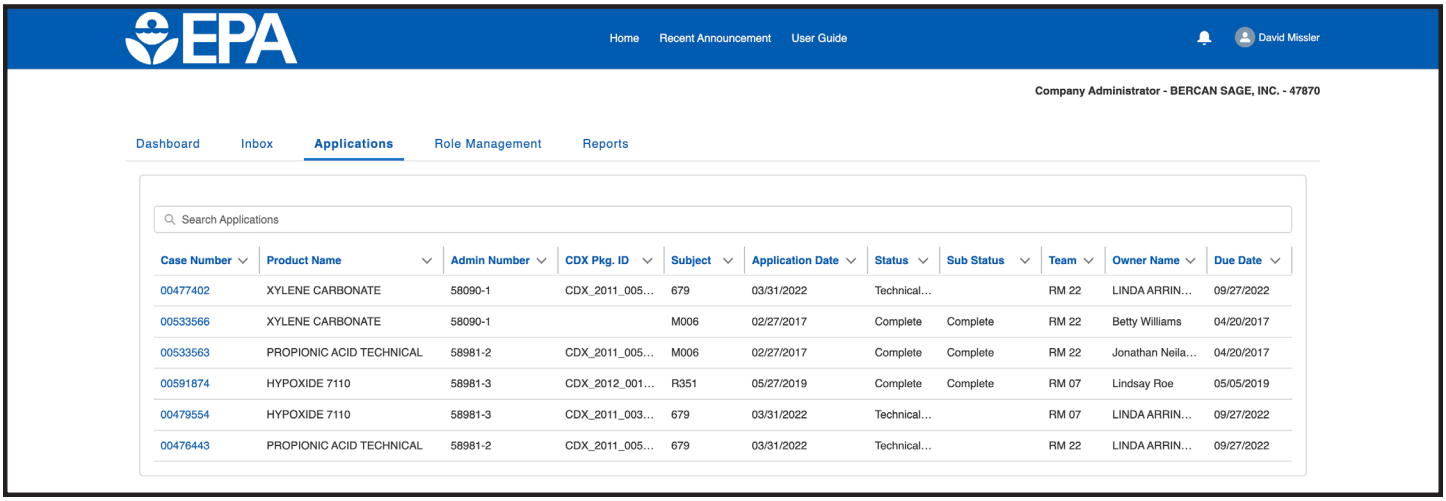
The **Pending Applications** tile shows all currently active applications your organization has with EPA.

The **Applications** tile shows all active and historic applications your organization has with EPA.

Click the **View All** button on the Pending Applications to view the full list of active applications.

Click the **Applications Tab** or the **View All** button on the Applications tile to view the full list of active and historic applications for your organization.

Applications Tab



You can access the Applications Tab by clicking **Applications** in the Tab bar near the top of the screen, or by clicking **View All** in either the Pending Applications or Applications tiles on the Dashboard.

If you click the **Applications Tab** or **View all** in the Applications tile, you will see a list of all current applications as well as all completed applications for your company.

If you click **View All** in the Pending Applications tile, you will only see a list of current applications that have not completed the review process.

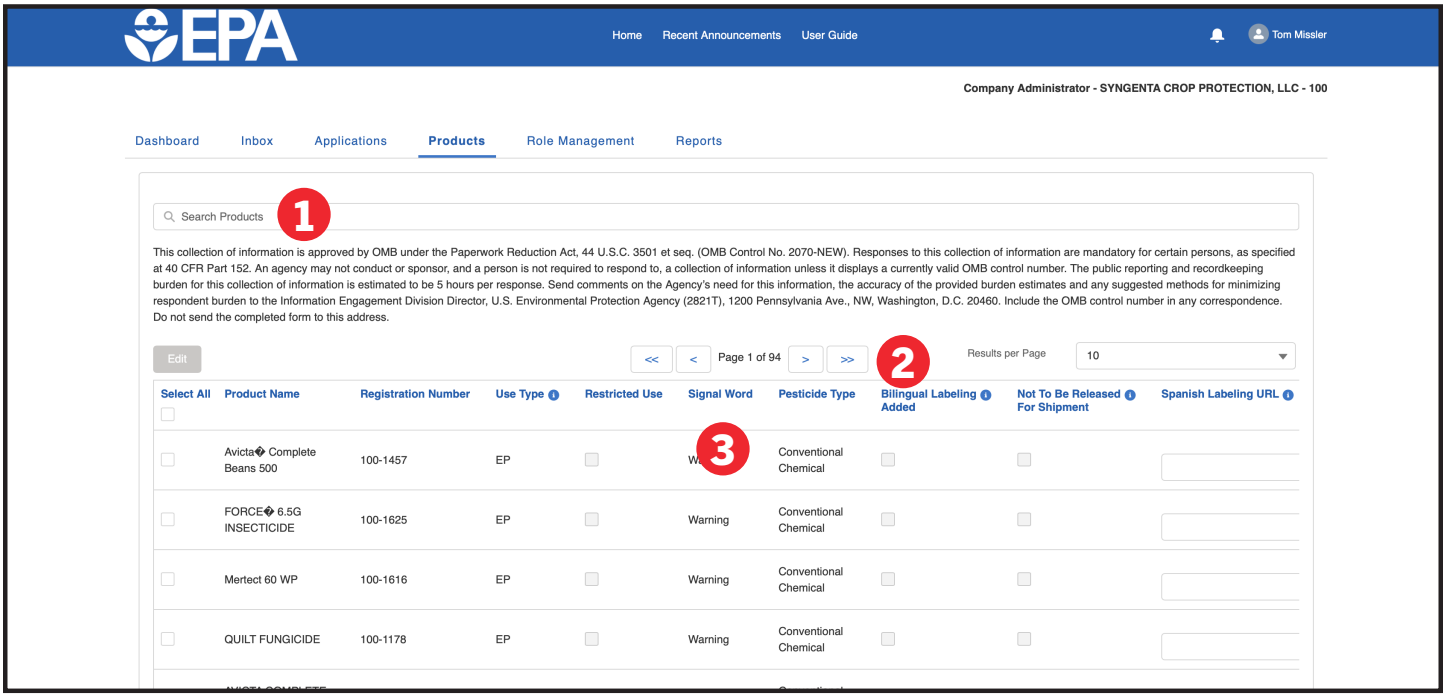
There are 11 fields in the table:

- **Case Number**
- **Product Name**
- **Admin Number**
- **CDX Pkg. ID**—The CDX tracking number
- **Subject**
- **Application Date**
- **Status**—Indicates the stage of EPA review process the application is currently in.
- **Substatus**—This is dependent on what Status the application is in. Not all Statuses have Substatuses.
- **Team**—This is the name of the EPA team working on your application, not the people in the My Team tile.
- **Ower Name**
- **Due Date**—The date when the action work is to be completed before any negotiation or changes.

Clicking a **Case Number** will take you to the Case Details page.

You can search the list of cases by Case Number, Product Name, Admin Number, CDX Pkg. ID, Subject, or Status.

Products Tab



You can access the Products Tab by clicking **Products** in the Tab bar near the top of the screen, or by clicking **View All** in the My Products tile on the Dashboard.

On the Products Tab you will see:

1) The **search** bar. You can search the list by Product Name, Registration/Admin Number, Use Type, Signal Word, or Pesticide Type. As you type, the list will filter the results.

2) The **page navigation** section where you can adjust the number of products that appear on the page and navigate through the pages.

3) The products table. There are 9 fields in the table:

- **Product Name**
- **Registration Number** – Product's EPA registration number.
- **Use Type** – The options are "EP" (end-use product) or "MP" (manufacturing use product)
- **Restricted Use** – A checkmark indicates whether the product is a Restricted Use pesticide. An empty checkbox indicates that it is not an RUP.
- **Signal Word** – The options are Danger, Warning, Caution, or None. These correspond to the product's toxicity category.
- **Pesticide Type** – The options include conventional pesticide, antimicrobial pesticide, and biopesticide. These options can help you determine the PRIA 5 product type, though they do not correspond exactly.
- **Bilingual Labeling Added** – A check indicates the product has bilingual labeling. An empty checkbox indicates a product does not have bilingual labeling.
- **Not To Be Released For Shipment** – A check indicates the product is not going to be released for shipment. An empty checkbox indicates a product may be released for shipment.
- **Spanish Labeling URL** – Users can optionally paste in a URL where the bilingual labeling is stored.

Clicking any column header once will organize the table in ascending order based on that column. Clicking the same column again will sort the table in descending order based on that column.

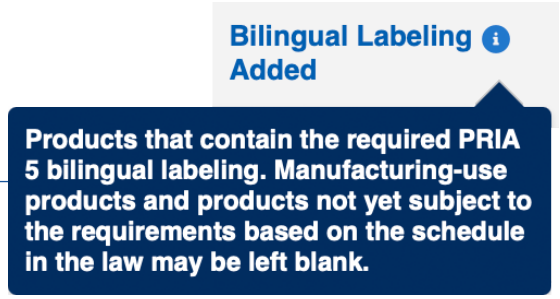
For example, clicking the Product Name column once will sort the table by product name from A to Z. Clicking Product Name again will sort the table by product name from Z to A.

For columns with a checkbox, clicking once will sort the table with empty checkboxes at the top. Clicking a second time will put checked checkboxes at the top.

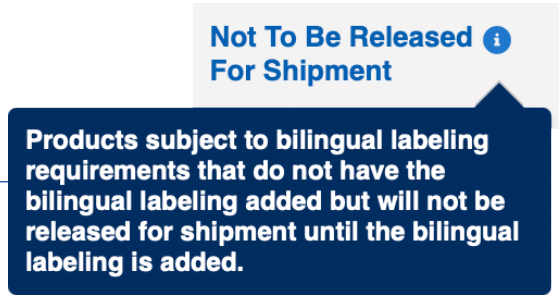
Tool Tips

Column headers also contain “i” icons. If you hover over the icons, tool tips will appear with simplified instructions on what to report.

Bilingual Labeling Added Tool Tip



Not To Be Released For Shipment Tool Tip



Spanish Labeling URL Tool Tip

How to Certify PRIA 5 Bilingual Labeling Compliance

Background

The Pesticide Registration Improvement Act of 2022 (PRIA 5) requires that each registered end-use pesticide product released for shipment include Spanish language translations for parts of the labeling contained in EPA's *Spanish Translation Guide for Pesticide Labeling.*

Translations may be included either on the pesticide product container itself or through a link to such translation via scannable technology or other electronic methods readily accessible on the product label. Antimicrobial pesticide products and non-agricultural/non-restricted use pesticide products may instead provide a link to bilingual safety data sheets (SDS).

There is rolling schedule for the implementation of bilingual labeling requirements, from December 2025 to 2030. That schedule is found in the table below.

Please see https://www.epa.gov/pesticide-labels/bilingual-labeling for more information on PRIA 5 bilingual labeling requirements.

Should I report compliance with PRIA 5 bilingual labeling requirements?

Do report compliance if…

- You have **any** products that are currently required to include bilingual labeling. (***Note:*** this includes both end-use products and products with a special local needs designation.)
- You have **any** products that would currently be required to include bilingual labeling, but you are not releasing them for shipment at this time.

Do not report compliance if…

- **None** of your products are end-use products or have a special local needs designation (i.e., all are manufacturing use products or experimental use permits).
- **None** of your end-use products/special local needs permits are currently required to include bilingual labeling.

If **none** of your products are end-use products/special local needs permits (i.e., all are manufacturing use products or experimental use permits) or **none** are currently required to include bilingual labeling, **DO NOT** check boxes or certify responses in MyPeST.

What are the compliance deadlines for bilingual labeling requirements?

Pesticide Product Type	Bilingual Labeling Due	Report Due in MyPeST
Restricted Use Pesticides (RUPs)	December 29, 2025	July 31, 2026*
Agricultural Products (Non-RUPs)		
Acute Toxicity Category I	December 29, 2025	July 31, 2026*
Acute Toxicity Category II	December 29, 2027	January 28, 2028*
Antimicrobial and Non-Agricultural Products		
Acute Toxicity Category I	December 29, 2026	January 28, 2027*
Acute Toxicity Category II	December 29, 2028	January 28, 2029*
All Other Pesticide Products	December 29, 2030	January 28, 2031*

**Provided OMB has approved EPA's ICR.*

Product types are defined as follows:

Product Type	Definition
Restricted use pesticides	Products classified as "restricted use" under 40 CFR 152.160; they have a "restricted use" statement on the front panel.
Agricultural products	Products that include an "Agricultural Use Requirements" statement on the label. Dual use products are subject to agricultural product deadlines. Animal products are not considered agricultural products.
Antimicrobial products	An antimicrobial pesticide is intended to disinfect, sanitize, reduce, or mitigate growth or development of microbiological organisms or protect inanimate objects, industrial processes or systems, surfaces, water, or other chemical substances from contamination, fouling, or deterioration caused by bacteria, viruses, fungi, protozoa, algae, or slime. Wood preservatives and anti-foulants are classified as antimicrobial pesticides if the products have antimicrobial claims.
All other pesticide products	Agricultural, antimicrobial, and non-agricultural pesticide products classified as acute toxicity categories III and IV.

How do I report compliance with bilingual labeling requirements?

For each pesticide product, check the product's type and acute toxicity category to determine its compliance date. (Some information is available in MyPeST to assist you, but you should verify product information using the product label.)

Then, determine whether the product label includes bilingual labeling by any of the methods permissible under PRIA 5 (e.g., translated text on the container; URL or QR code to the translated text; or translated SDS where applicable).

If a product has reached its compliance date but does not contain the required bilingual labeling, check whether the product has been released for shipment.

Note: Please report all products to which you have added bilingual labeling, even if they are not required to include bilingual labeling at this time (e.g., products that have not reached their compliance dates yet). EPA wishes to acknowledge registrants who have gone above and beyond the statutory requirements.

How Do I Certify That Bilingual Labeling Was Added Or That A Product Is Not To Be Released For Shipment?

1) First, you will select all products you want to certify as having bilingual labeling.
2) Go to the "My Products" tab. Find the product or products you want to certify.

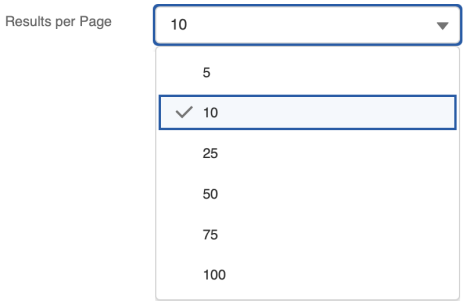
   If you are looking for a specific product, using the **Search** bar is the most efficient method.

   To select products using the **page navigation features,** begin by setting the number of items per page. You can show 5, 10, 25, 50, 75, or 100 products.

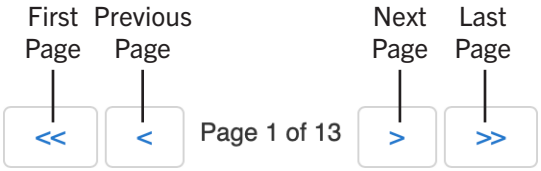
   Clicking the **Select All** checkbox will select all the products displayed on that page, not all products registered to an organization. For example, if a company has 50 products and is showing 10 items per page, clicking **Select All** on page 1 will only select those 10 products; the other 40 products are not selected.

   To choose specific products, you can click the checkbox next to each product to select it and click it again to deselect it. If your company has multiple pages of products, you can select specific products (or click **Select All**) on page one, click the **Next Page** button, and click specific products (or **Select All**) on page two, and repeat the process on each page until you have made all your selections.
3) Click the check boxes next to all products to which you have added bilingual labeling.

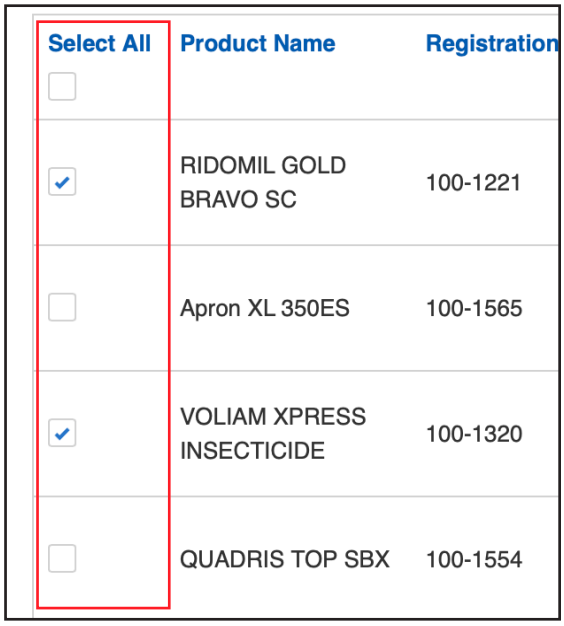
Items per page selector



Items per page navigation



Selecting specific products from the list



Select All	Product Name	Registration
☐		
☑	RIDOMIL GOLD BRAVO SC	100-1221
☐	Apron XL 350ES	100-1565
☑	VOLIAM XPRESS INSECTICIDE	100-1320
☐	QUADRIS TOP SBX	100-1554

4) Clicking a check box will make the **Edit** button active. Once you have finished clicking the check boxes next to all products to which you have added bilingual labeling, click **Edit.** This will bring up the **Certify Bilingual Label Status** popup.

   The **Certify Bilingual Label Status** popup has 4 sections:

   1. An explanation of what you are certifying.
   2. Three bubbles to certify either:
      - Bilingual Labeling Added,
      - Not To Be Released For Shipment, or
      - Remove Product Certification.
   3. The list of products you selected that the certification will be applied to.
   4. Submit and Cancel buttons to confirm your choice.

5) To certify products as having bilingual labeling added, select the **Bilingual Labeling Added** bubble and click **Submit.** This will bring up the **Certify Bilingual Labeling Compliance** popup.

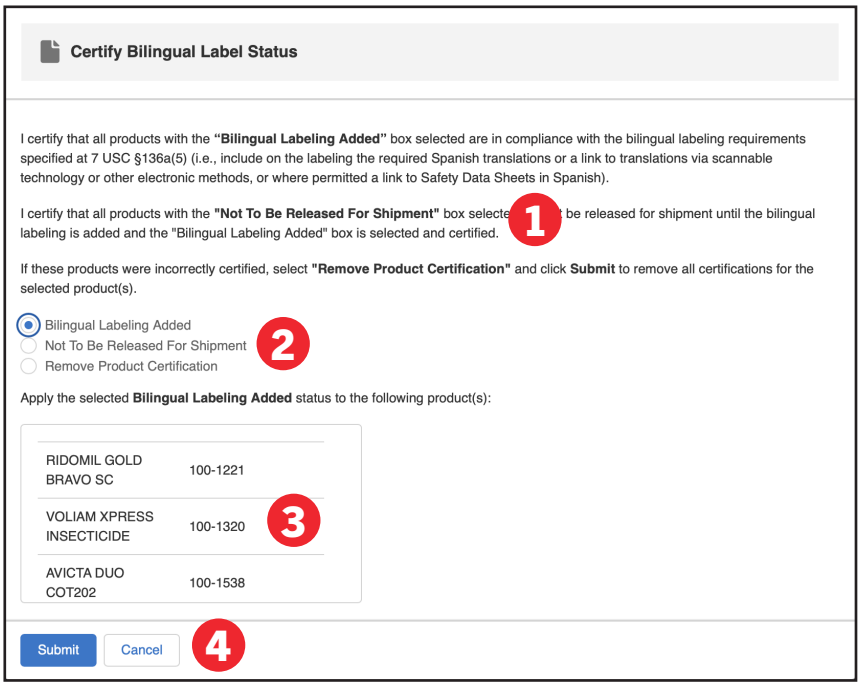
6) Click Yes to confirm that you have added bilingual labeling, or click the X to return to the Products Table to make changes.

7) If you click Yes, you will be returned to the Products Tab. You will now see checkmarks for your selected products in the Bilingual Labeling Added column.

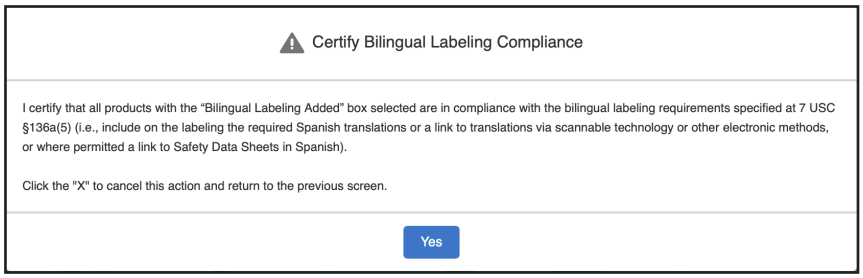
   A green toast message will be displayed at the top of the page for a few seconds, indicating that your certification has been recorded by the EPA.

Note: There is no "Save" button in MyPeST. However, the checkmarks will save when you accept the certification statements, so you do not have to complete the report in one sitting.

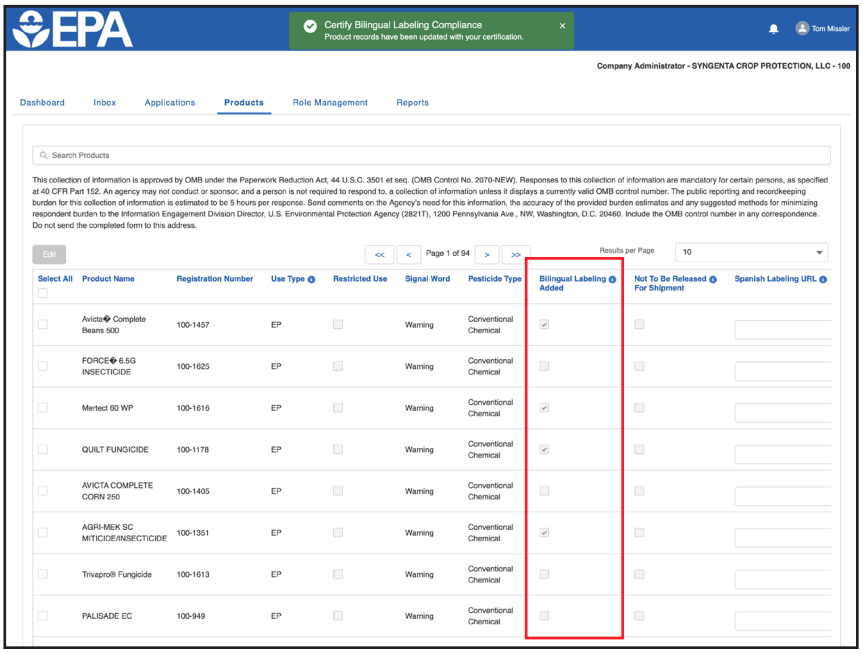
Certify Bilingual Label Status popup with Bilingual Labeling Added selected.



Certify Bilingual Labeling Compliance popup.



Products Tab showing your selected products are now certified Bilingual Labeling Added and the toast message confirming that the product records have been updated.



8) Next, certify any products that would be required to include bilingual labeling if shipped, but which you are not releasing for shipment.

> ***Note:*** **Only** report whether a product as "Not To Be Released for Shipment" if that is the basis for not including bilingual labeling. Otherwise, leave those checkboxes blank.

9) To certify products as "Not To Be Released For Shipment," repeat steps 2 – 7, but this time select the **Not To Be Released For Shipment** bubble instead.

10) Click **Yes** on the acknowledgement popup to confirm that selected products will not be released for shipment until bilingual labeling is added. (Please note that you can release a product for shipment at any time; just add any required bilingual labeling to the product and update your bilingual labeling responses in MyPeST.)

11) You will return to the Products Tab. You will now see checkmarks for your selected products in the **Not To Be Released For Shipment** column.

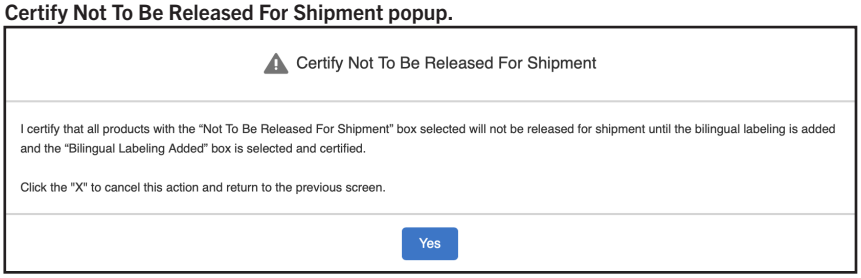
    A green toast message will be displayed at the top of the page for a few seconds indicating that your certification has been recorded by EPA.

12) If you incorrectly certified a product, you can check the box beside it, click **Edit,** click the **Remove Product Certification** bubble, and click **Submit**. There will be no popup. However, you will return to the Products tab and any checkmarks that were in the **Bilingual Labeling Added** or **Not To Be Released For Shipment** columns will be removed.

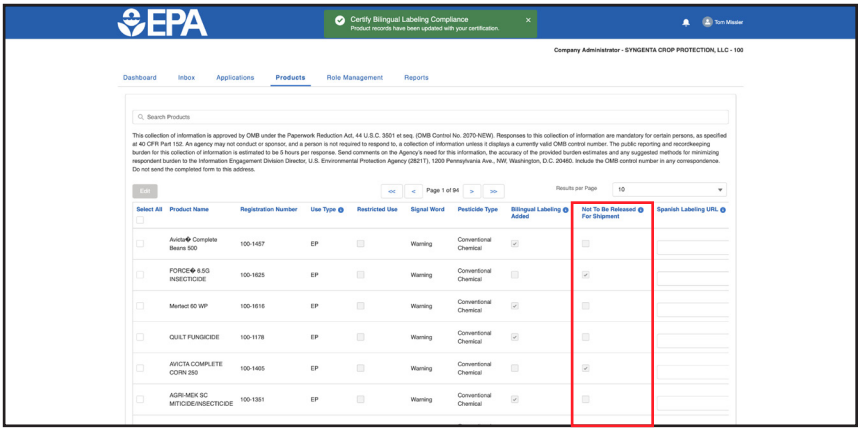
    A green toast message will be displayed at the top of the page for a few seconds indicating that your certification has been updated.

> ***Note:*** You can change your responses in MyPeST **at any time.** If, before the next reporting deadline, you decide to release a selected product for shipment, remove the certification by following the instructions below, and instead certify that you have added the required bilingual labeling.

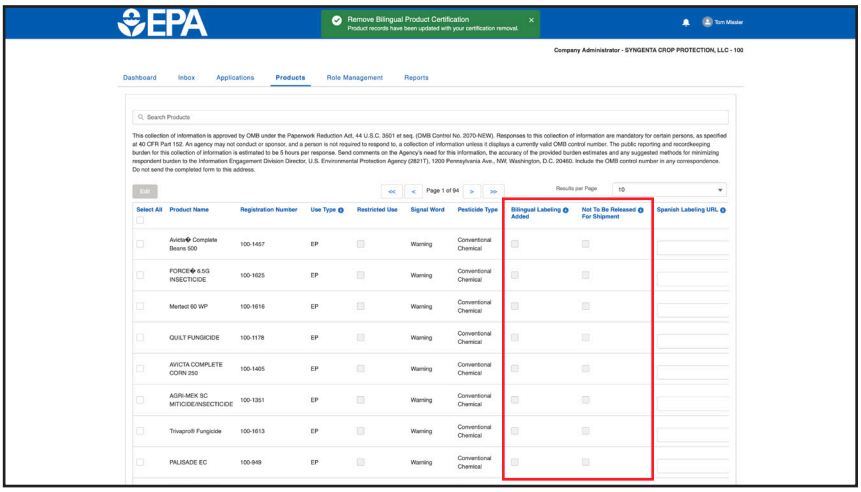
Certify Not To Be Released For Shipment popup.



Products Tab showing your selected products are now certified Not To Be Released For Shipment.



Products Tab showing your selected products have the certifications removed and the toast message confirming that the product records have been updated.



How Do I Provide EPA With Links To Bilingual Labeling?

While reporting on compliance with PRIA 5 bilingual labeling requirements, registrants may also provide a link to their translated labels. Note that providing a URL is not required to report bilingual labeling compliance, nor are registrants who have not put their labeling online expected to create URLs exclusively to share it via MyPeST.

However, EPA hopes that registrants will consider providing URLs to bilingual labeling in MyPeST so that EPA may publish the link on the Pesticide and Product Labeling System (PPLS) to make bilingual labeling accessible to farmworkers and the general public.

1) To add a URL link to a bilingual label, copy the link and paste it into the Spanish Labeling URL field next to the corresponding product.

> ***Note:*** For products that have multiple market labels, note that there is a **255-character** limit on the field. Registrants should include the most representative or complete label, not all possible labels.

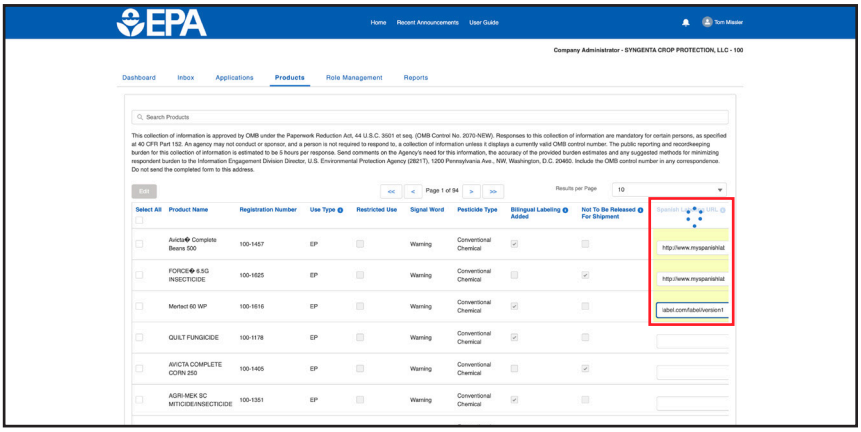
2) When you paste the URL in the field, the background of that table cell will turn yellow. You will see six dots at the top of the column indicating that you have made a change, but that your change has not been saved yet.

   In the example, there are three URLs that are highlighted yellow, which means you entered three URLs, but so far, none of them have been saved. If you close my browser window now, that data will be lost.

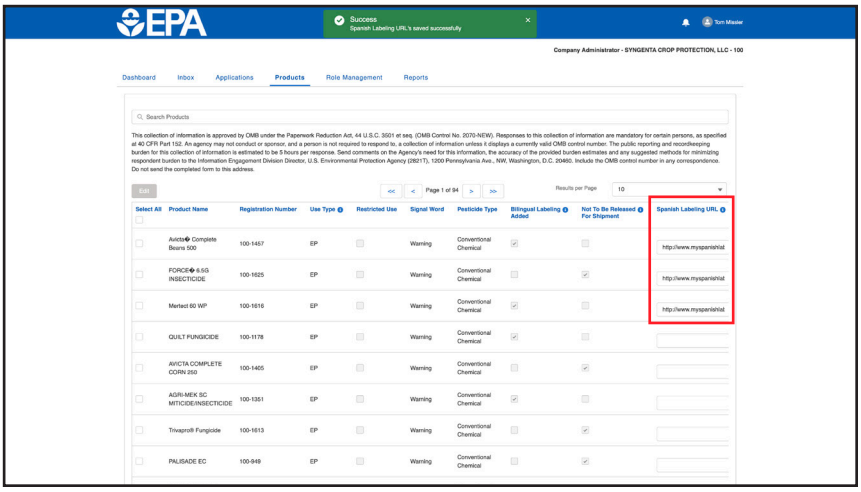
3) After five seconds of inactivity, MyPeST saves all the highlighted data. You will see a green toast message at the top of the screen indicating that your URLs have been saved, and the URLs will have a white background again. Because responses save automatically, you do not have to finish the report in one sitting.

> ***Note:*** the Spanish Labeling URL is separate from the Bilingual Labeling Added checkbox. So, if you select products that have been certified bilingual and click Remove Product Certification, the URL will remain in the Spanish Labeling URL field until you highlight it and delete it.

Products Tab showing three URLS have been entered, but since they have a yellow background and the blue dots are visible, the data has not yet been saved in MyPeST.



Products Tab showing your the URLs have been successfully saved.



I Already Reported Compliance. Should I Update/Recertify My Responses?

Do update/recertify responses if…

- You have any products that have reached their compliance dates since you last reported.
- You have registered any new products that require bilingual labeling per the compliance deadlines, or made label changes (such as adding agricultural uses) that change a product's PRIA 5 compliance dates.
- You released for shipment a product that must include bilingual labeling per the compliance deadlines, but that previously did not have bilingual labeling because it was not released for shipment.

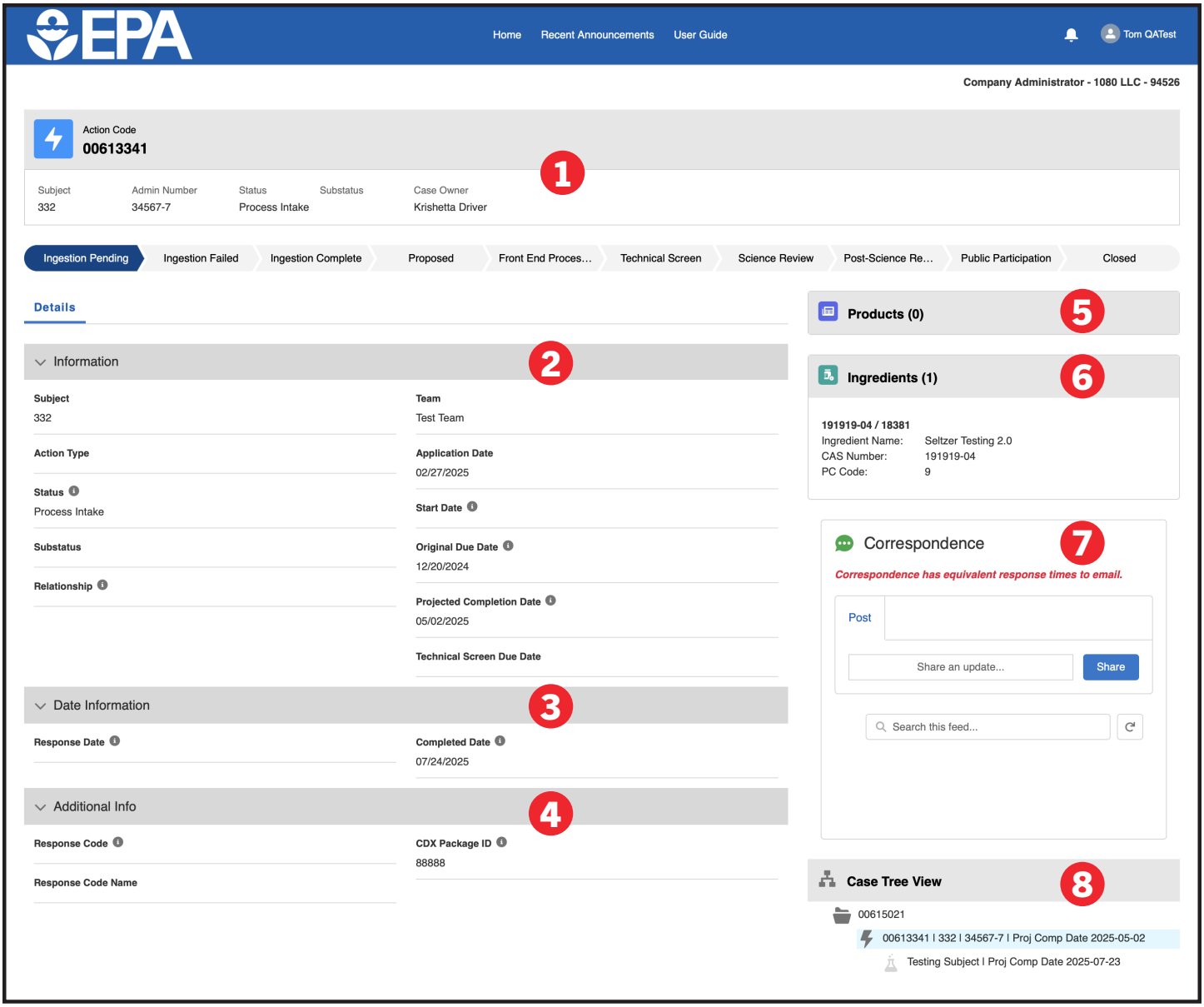
Otherwise, **do not** update or recertify responses.

Assistance

If you have questions about bilingual labeling, you can get assistance by sending an email to **oppbilinguallabels@epa.gov.**

The Case Details Page

When you click on a **Case Number** link on the Dashboard or Applications Tab, the Case Details Page opens in a new tab on your browser. You can only navigate back to the Home page from this page, but you can click on the previous tab in your browser to return to the Dashboard or Applications Tab and navigate from there. This allows you to have multiple cases open at the same time.



There is a lot of information on the Case Details page. We have broken it up into eight sections:

1. **Case Header and Path**
2. **Information**
3. **Date Information**
4. **Additional Information**
5. **Products**
6. **Ingredients**
7. **Correspondence**
8. **Case Tree View**

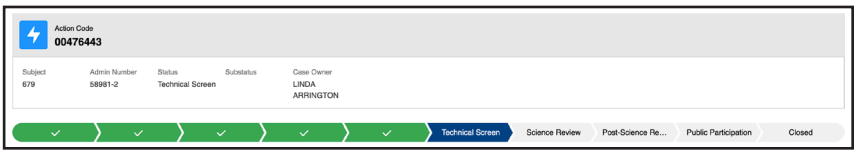
We will cover each section separately.

Case Header and Path

This is a quick reference section that has the most important case information available at a glance. You can see:

- **Subject**
- **Admin number**
- **Status**—indicates the stage of the EPA review process the application is currently in.
- **Substatus**—This is dependent on what Status the application is in. Not all Statuses have Substatuses.
- **Case Owner**

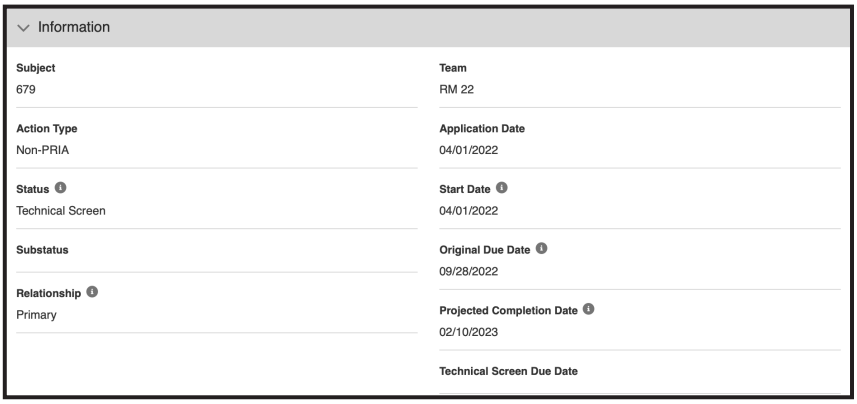
Below those fields you will see the Path. This shows the Status your application is currently in and shows you how many more steps it has before it is closed.



Information

This section has the following fields:

- **Subject**
- **Action Type**—can be Pesticide Registration Improvement Act (PRIA) or Non-PRIA
- **Status**—indicates the stage of the EPA review process the application is currently in.
- **Substatus**—This is dependent on what Status the application is in. Not all Statuses have Substatuses.
- **Relationship**—shows whether the record is Primary or Secondary.
- **Team**—The EPA Team assigned to the case.
- **Application Date**—The date the application was submitted to EPA.
- **Start Date**—the date when the action work was started on the application.
- **Original Due Date**—the date used to indicate the expected completion date and entered by EPA.
- **Technical Screen Due Date**



Date Information

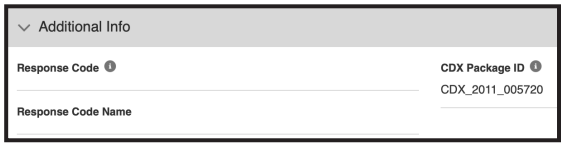
This section has the following fields:

- **Response Date**—the date a Response Code was added/updated.
- **Completed Date**—the date when the Action Work has been completed.

Additional Information

This section has the following fields:

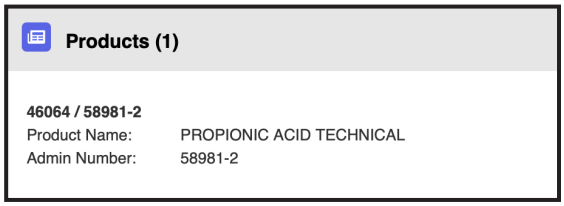
- **Response Code**—the Response Code selected upon Case Closure
- **Response Code Name**
- **CDX Package ID**—CDX tracking number



Products

This is different than the My Products tile on the Dashboard. My Products lists all your company's products. The Products tile on the case details page only shows the products related to that specific case. This section has the following fields:

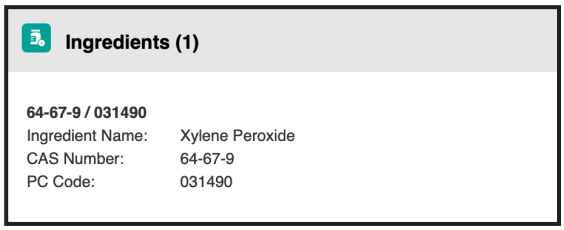
- **Product Name**
- **Admin Number**



Ingredients

This is different than the Active Ingredients tile on the Dashboard. Active Ingredients lists all your company's ingredients. The Ingredients tile on the case details page only shows the ingredients related to that specific case. This section has the following fields:

- **Ingredient Name**
- **CAS Number**—Chemical Abstract Service Numbers are provided for the convenience of the regulated community and the public to aid in the identification of the designated hazardous substances.
- **PC Code**—or Pesticide Chemical Code is a unique chemical code number assigned by the EPA to a particular pesticide active ingredient, inert ingredient or mixture of active ingredients.



Case Tree View

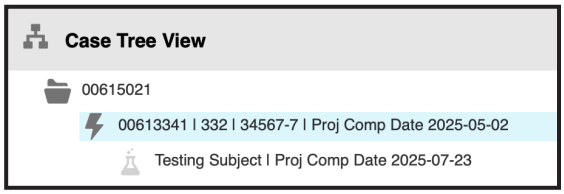
This section shows the following information:

- **Request Case**—Signified by a file folder icon and shows the Case Number
- **Action Code Case**—Signified by a lightning bolt icon and shows the Case Number | Subject | Admin Number | Projected Completion Date
- **Task Group**—Signified by a beaker icon and shows the Subject | Projected Completion Date

The case that is highlighted in the Case Tree View is the case you are currently viewing on the Case Details Page.

If the icon is dark gray, it is an active case. If the icon is light gray, that indicates the case is closed.

In the example, the user is viewing the Action Code Case. The Request Case and Action Code Case are active and the Task Group is closed.



Correspondence

The Correspondence tile provides a place for communication and collaboration about the case you are viewing, allowing you to post updates, share insights, ask questions, and engage in conversations with the EPA. It does not support email integration. Please note, Correspondence has equivalent response times to email.

1. To make a Post, type a message in the **Share an update...** text box. When you are finished click the **Share** button. In the example the Post is "Here is a label with suggested revisions"
2. To Comment on a Post, type a message in the **Write a comment...** text box. When you are finished click the **Comment** button. In the example the Comment is "@Russ Cargill (EPA) Thank you!"
3. Using the "@" followed by a person's name "tags" that person in your comment. If you look at the Comment Riley Schultz made in the example, it starts with "@Russ Cargill (EPA)." That indicates that Russ was tagged in Riley's comment. You can tag people in a Post or a Comment.
4. If another user makes a Comment on your Post or previous Comment, you will receive a Notification—a red circle with a number will appear over the bell icon in the header. The number indicates how many unread notifications you have. Clicking the Notification will take you to that thread where you can see all discussions related to the post.

Emails and Notifications

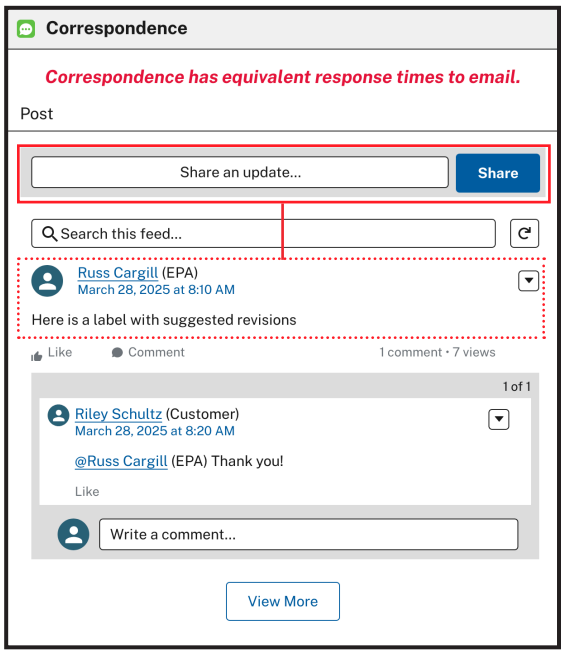
When you make a Post in Correspondence, you are the owner of that thread. You will receive an email and a Notification every time a comment is made. You do not need to be tagged; you will automatically receive the alerts.

If a user makes a Comment in a thread, that user will receive an email every time another comment is made. They will not receive a Notification unless they are tagged. You, as the Post owner will receive an email and Notification.

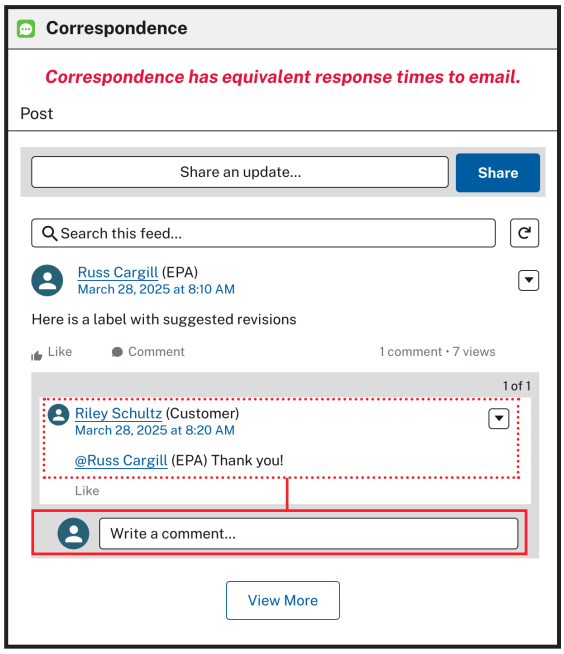
If a user makes a Comment and tags a specific person @User Name, the tagged person will receive an email and a Notification. Everyone else who posted a comment in the thread will receive an email, and the Post owner will receive an email and Notification.

If you reply to the email you received, your response will appear in Correspondence. You cannot tag people using @ from email, but if you type the person's full name in your message, they will receive an email alerting them. If you try to attach a file to your email and send it, the message will go through but the attachment will not.

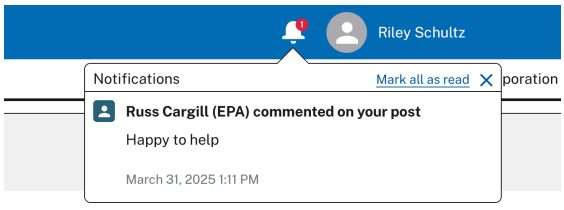
Step 1



Step 2



Step 4



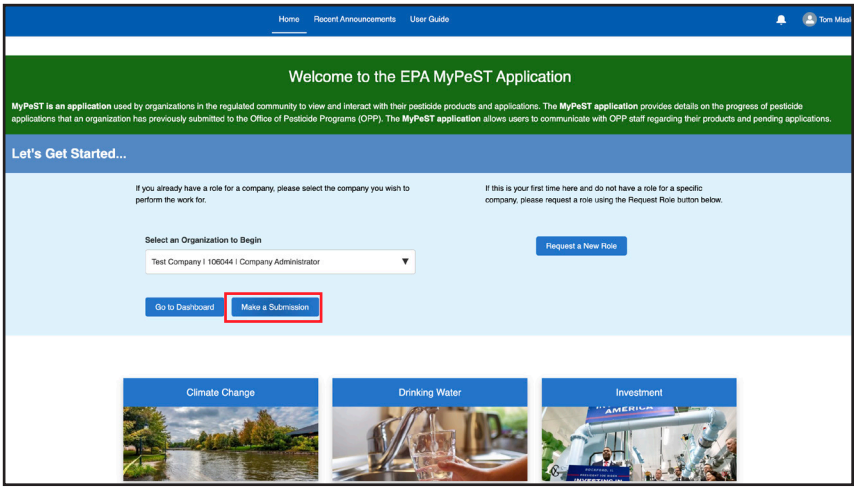
Make a Submission

You are now able to submit a new application or update an existing application for a pesticide through MyPeST.

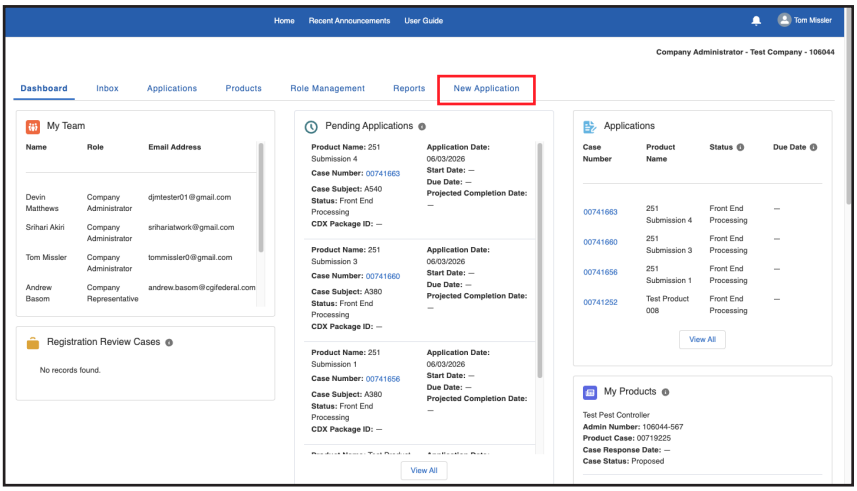
After you select your organization on the Home page, you now have the option to click the **Make a Submission** button. You can also access this functionality by clicking the **New Application** tab from the Dashboard.

To get started, click the **Make a Submission** button on the Home Screen or the **New Application** tab from any other page.

Make a Submission button on the Home page



New Application Tab from any page in MyPeST



Start New Submission

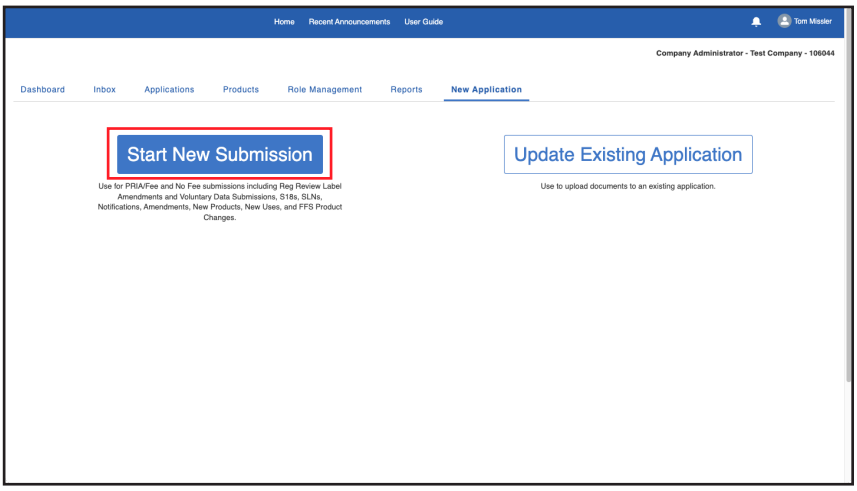
1. You will see two buttons: **Start New Submission** or **Update Existing Application.** Use Start New Submission if you are registering a new product with the EPA. Use Update Existing Application if you need to add documents in support of a product that has begun the registration process. To begin, click **Start New Submission.**

2. You will see the following fields:
   - **Case Contact Name**
   - **Product**
   - **Action Type**
   - **Action Code**
   - **Fee**
   - **Upload File(s)**

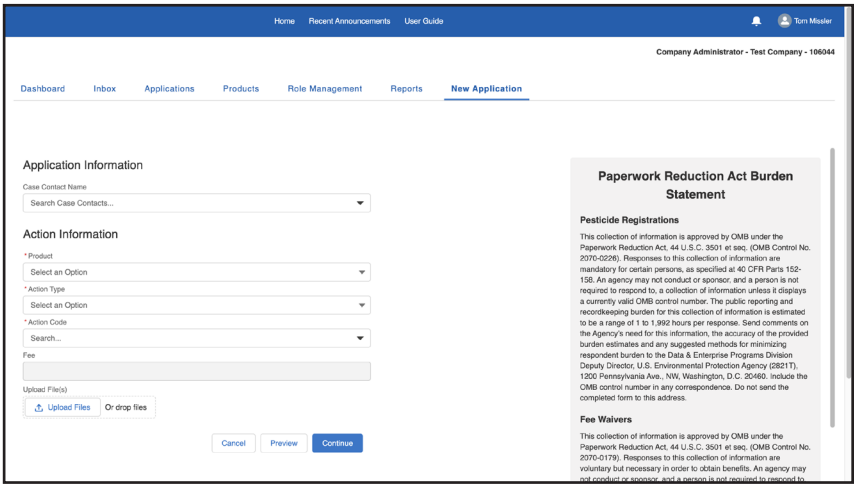
3. Click the **Case Contact Name** dropdown to designate a person in your organization who has registered in CDX and Login.gov. This is the person EPA staff will contact with questions about the product you are registering. If the person you want to designate as contact has not yet registered in MyPeST have them follow the steps in the Account Setup section before making a submission. If a contact is not selected the person making the submission will be designated the contact.

4. Next, click the **Product** drop-down. Your choices are:
   - **No Product**—choose this option for:
     - Tolerance Petition
     - Inert Mixture.
   - **New Product**—choose this option for:
     - PRIA submissions with a Fee
     - Registration Review Label Amendments
     - Section 18 / Emergency Exemptions
     - Notifications
     - New Products
     - Fee For Service (FFS) Product Changes
     - PRIA submission with No Fee
     - Voluntary Data Submissions
     - Special Local Need (SLN)
     - Amendments
     - New Uses
   - A product from the list of your company's registered products. Choose this option if you need to file a new use application for an existing product. Selecting an existing product will create a new case with the EPA for that product.

The screen you see if you click the Make A Submission or the New Application tab.



The fields you first see when you click the Start New Submission button.



5. If you select **New Product**, the **Product Name** field will appear. Enter the name of your product.

6. Click the **Action Type** drop-down. You can select one of two options:
   - PRIA
   - Non-PRIA

7. Click the **Action Code** drop-down. The selection you made for **Action Type** will determine what **Action Codes** are available. For a list of Action Codes for the current fiscal year, please see Appendix E.

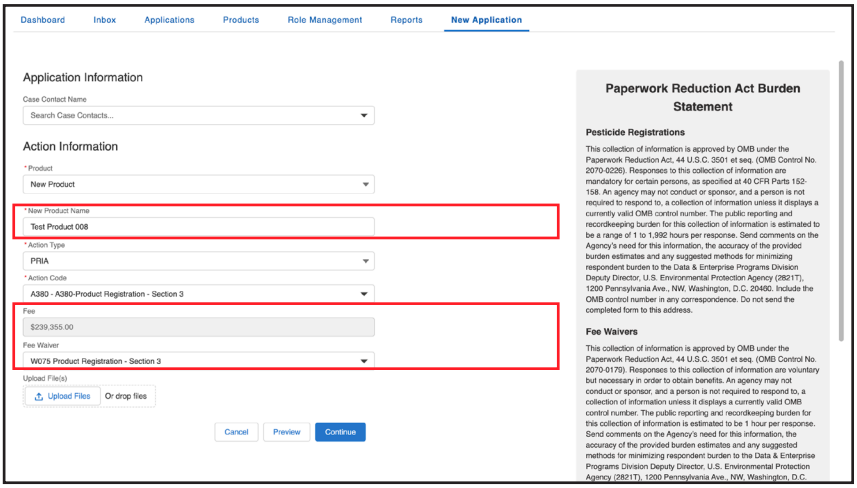
   Some Action Codes require extra information:
   506, 516, 526, 536, 546, 556, or 580—The **Source Product** field will appear. You must select a product from the drop-down list before you can submit.
   676, 678, 679—The **Active Ingredients** field will appear. You must select an ingredient from the drop-down list before you can submit.
   800—The **Source Product** field and **Distributor Company** field will appear. You must select one product and one distributor before you can submit.

8. The **Fee** field will automatically update with amount due based on the **Action Code** you select.
   If a fee is due, the **Fee Waiver** drop down will appear. If you do not wish to claim a fee waiver, you can simply leave this field blank. If you would like to request a fee waiver, select the code corresponding to the percentage of the fee you would like the EPA to waive.

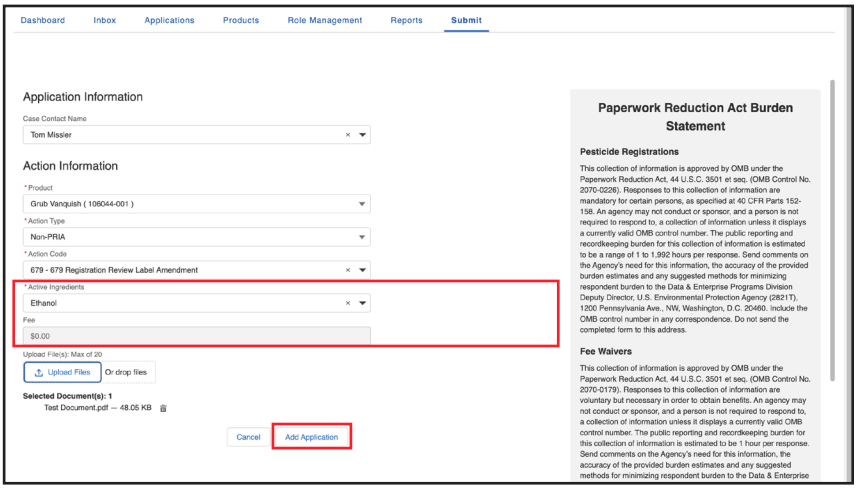
9. Finally, upload your document(s). You must upload at least one and no more than 200 documents. You can drag and drop your files or you can click the Upload Files button and select documents by navigating to the location on your hard drive.
   The accepted file types are:
   - .pdf (Adobe Acrobat)
   - .doc (Microsoft Word 1997–2003)
   - .docx (Microsoft Word 2003–Present)
   - .xls (Microsoft Excel 1997–2003)
   - .xlsx (Microsoft Excel 2003–Present)

10. Click **Add Application.** At the top of the screen a new **Applications Added** section will appear.

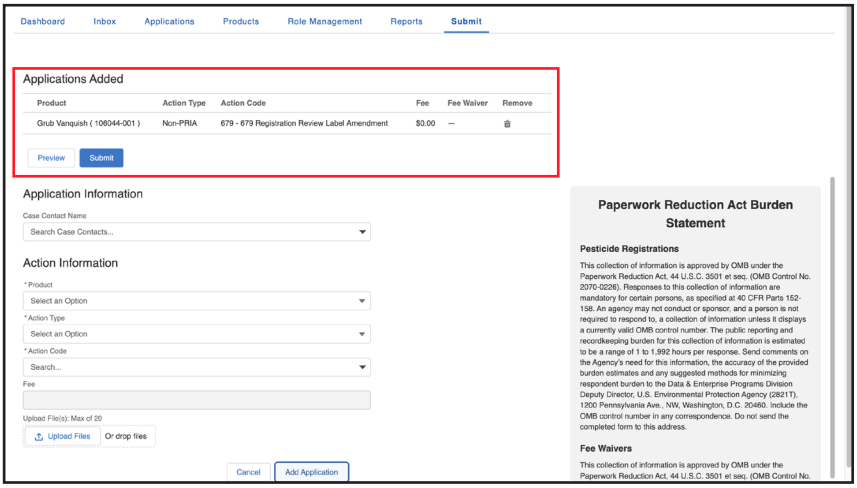
Start New Submission page with the New Product Name and Fee Waiver fields visible and the Fee field populated.



Start New Submission page with the Ingredient field visible and a document uploaded. Since there is no fee associated to this action code the Fee field has $0.00 and the Fee Waiver field is not visible.



After you click Add Application the Applications Added section will appear on the page. Clicking the trashcan icon will delete the application and all files associated to it. You can submit multiple applications at once by filling in the data, uploading at least one additional document and clicking Add Application again.



11. The **Applications Added** section displays the Product, Action Type, Action Code, Fee, and Fee Waiver information you provided.

    If you would like to see a summary of what you are about to send the EPA, click **Preview.** An html file will automatically download. You can open that file with your web browser.

12. If you are ready to send your application to the EPA, click **Submit.**

    Clicking the **trashcan icon** will delete your application and all documents you uploaded and nothing will be sent to the EPA.

13. The **Create Passphrase & Review** popup will appear. Your passphrase must meet the following criteria:
    - At least 8 characters long
    - No more than 20 characters
    - Spaces are allowed
    - No special characters (!,#, ?, etc.)
    - Case sensitive

    When your **Passphrase** and **Confirm Passphrase** field match the Continue button will become active. If you need to see what you typed, click the **eyeball icon** in the field.

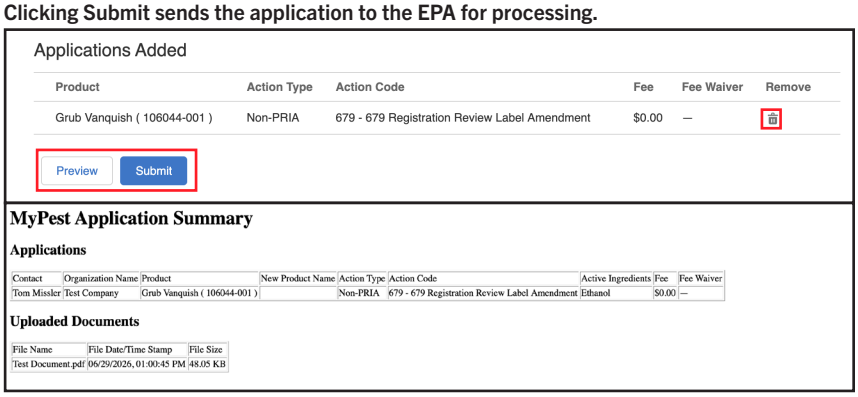
14. Click **Continue.**

15. The **Digital Signature Screen** popup will appear. Click **Accept** if you certify, under penalty of law that the information provided in this document is, to the best of your knowledge and belief, true, accurate, and complete. You are aware that there are significant penalties for submitting false information, including the possibility of fines and imprisonment for knowing violations. If not click **Decline** and cancel your submission.

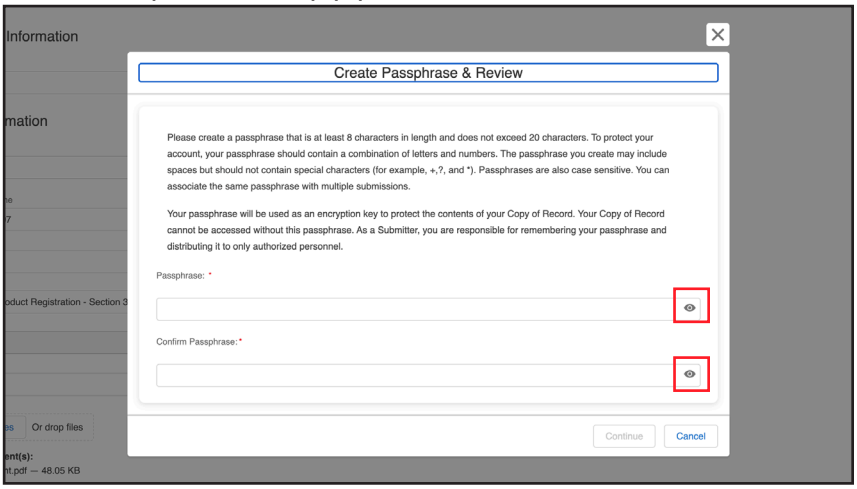
16. If you clicked **Accept** you will return to the submission page that is grayed out. You will see a green Success message appear to indicate that your submission was sent to the EPA.

Congratulations! You have successfully completed a product submission to the EPA.

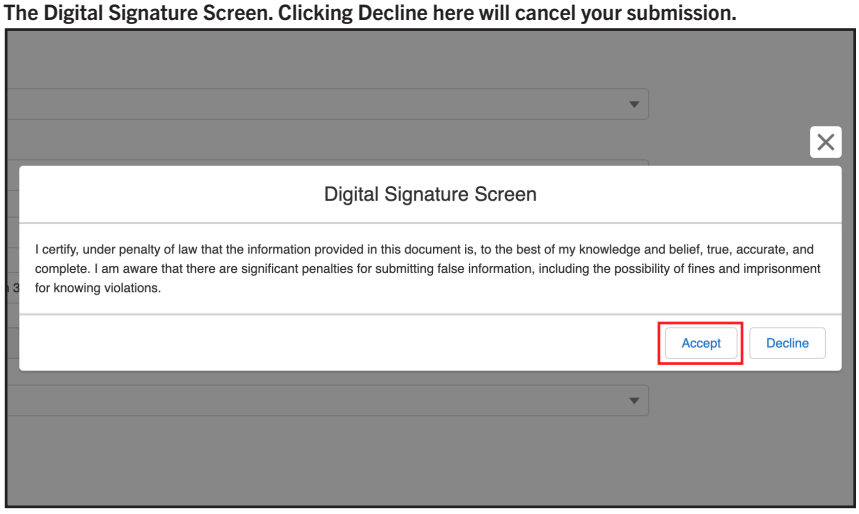
After you click Add Application the Applications Added section will appear on the page. Clicking the Preview button will download an html file of the MyPeST Application Summary. The trashcan icon will delete the application and all files associated to it. Clicking Submit sends the application to the EPA for processing.



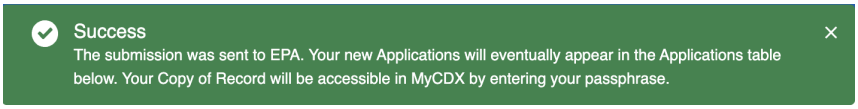
The Create Passphrase & Review popup.



The Digital Signature Screen. Clicking Decline here will cancel your submission.



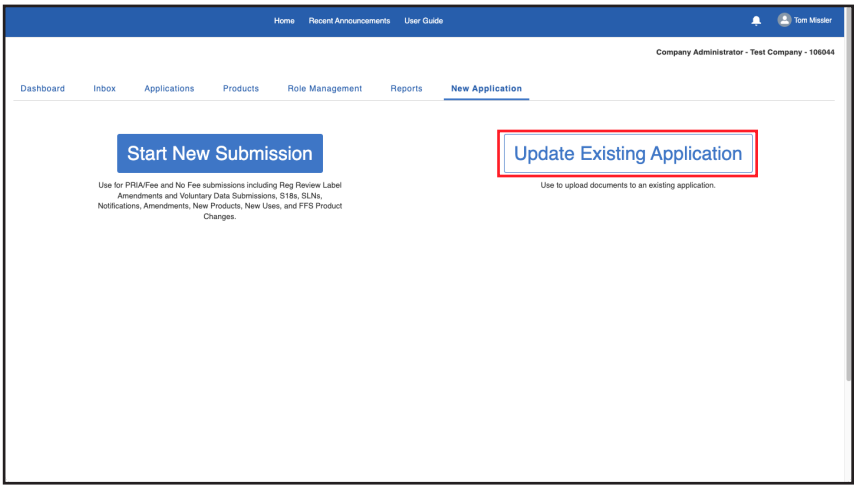
The message you will see when your submission is successfully submitted.



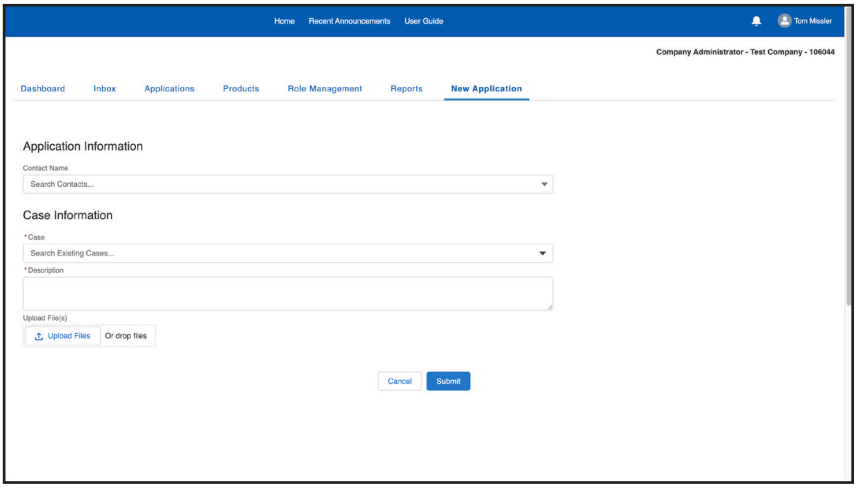
Update Existing Application

1. From the New Application screen, click **Update Existing Application.**
2. You will see the following fields:
   - **Contact Name**
   - **Case**
   - **Description**
   - **Upload File(s)**
3. Click the **Contact** drop-down to see a list of people in your organization who are registered in CDX and Login.gov. If the person you want to designate as contact has not yet registered have them follow the steps in the Account Setup section before making a submission.
4. Next, click the **Case** drop-down. You will see a list of cases your company has created with the EPA. Your case number is an 8 digit number created by Salesforce. You can find the case number by clicking the Applications tab and searching for the product you wish to update.
5. In the **Description** field you can explain what you are uploading and why you are uploading it. If you submitted a form with an error and are resubmitting a corrected version of that form or if the EPA requested more information and the document(s) you are uploading fulfill that request, you can explain that in the Description field.
6. Upload your document(s). You must upload at least one and no more than 200 documents. You can drag and drop your files or you can click the Upload Files button and select documents by navigating to the location on your hard drive.
   The accepted file types are:
   - .pdf (Adobe Acrobat)
   - .doc (Microsoft Word 1997–2003)
   - .docx (Microsoft Word 2003–Present)
   - .xls (Microsoft Excel 1997–2003)
   - .xlsx (Microsoft Excel 2003–Present)
7. Click **Submit.**

The screen you see if you click the Make A Submission or the New Application tab.



The fields you first see when you click the Update Existing Application button.



8. The **Create Passphrase & Review** popup will appear. Your passphrase must meet the following criteria:
   - At least 8 characters long
   - No more than 20 characters
   - Spaces are allowed
   - No special characters (!,#, ?, etc.)
   - Case sensitive

   When your **Passphrase** and **Confirm Passphrase** field match the Continue button will become active. If you need to see what you typed, click the **eyeball icon** in the field.

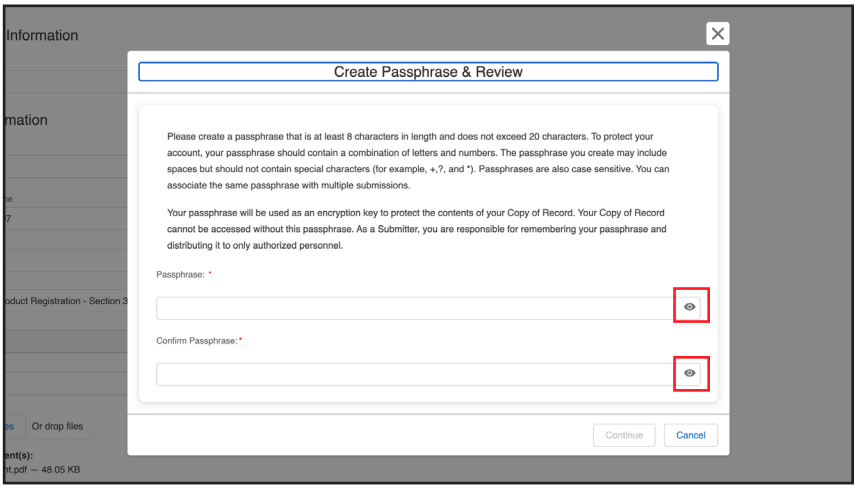
9. Click **Continue.**

10. The **Digital Signature Screen** popup will appear. Click **Accept** if you certify, under penalty of law that the information provided in this document is, to the best of your knowledge and belief, true, accurate, and complete. You are aware that there are significant penalties for submitting false information, including the possibility of fines and imprisonment for knowing violations. If not click **Decline** and cancel your submission.

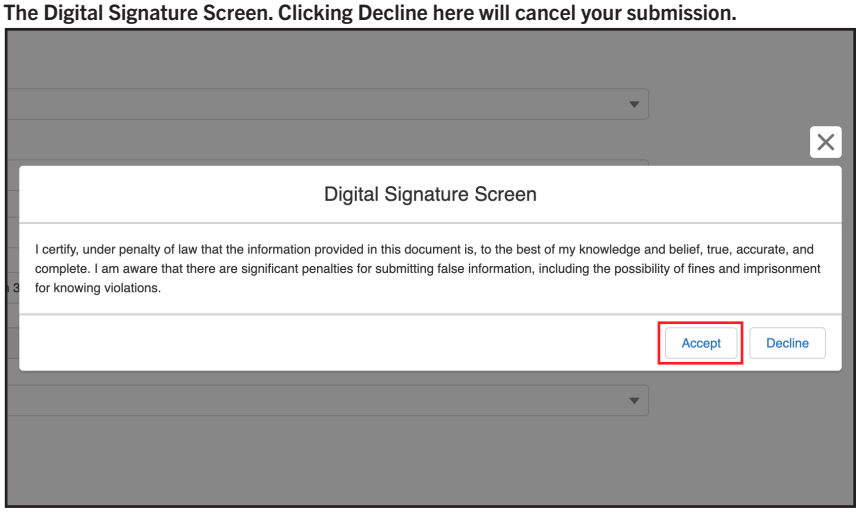
11. If you clicked Accept you will return to the submission page that is grayed out. You will see a green Success message appear to indicate that your submission was sent to the EPA.

Congratulations! You have successfully updated an application with the EPA.

The Create Passphrase & Review popup.



The Digital Signature Screen. Clicking Decline here will cancel your submission.



The notification you will see when your submission has been successfully sent to EPA.

Submitting Multiple Applications

It is possible to include multiple applications in a single submission. To determine if you should submit applications separately or if you can combine them into a single submission, follow these guidelines:

- All applications must be for the same Product.
- Non-PRIA actions cannot be part of a multi-application submission.
- You can only apply one Fee Waiver per submission.
- At least one document must be uploaded for each application in the submission.

The process for submitting multiple applications in a single submission follows many of the same steps that submitting a single application requires.

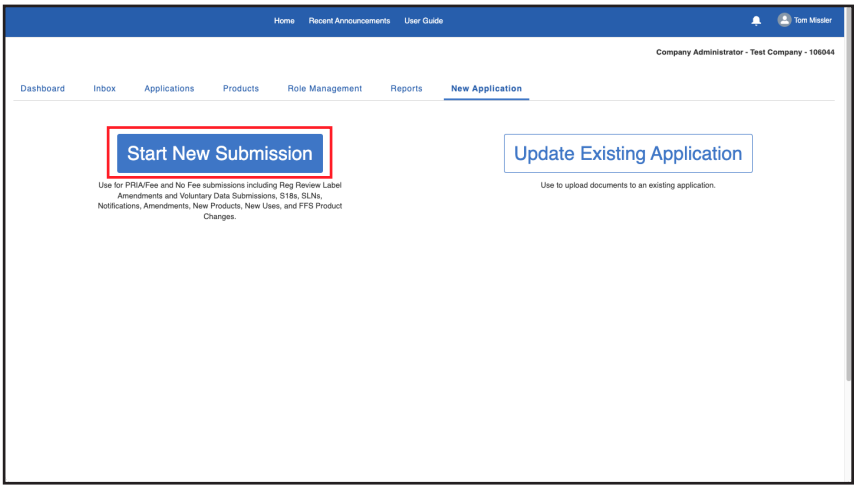
1. From the New Application screen, click **Start New Submission.**
2. Enter data for following fields:
   - **Case Contact Name**
   - **Product**
   - **Action Type—Must be a PRIA action**
   - **Action Code**
   - **Fee**
   - **Upload File(s)**

If any of the Action Codes trigger any additional fields, enter data for those as well.

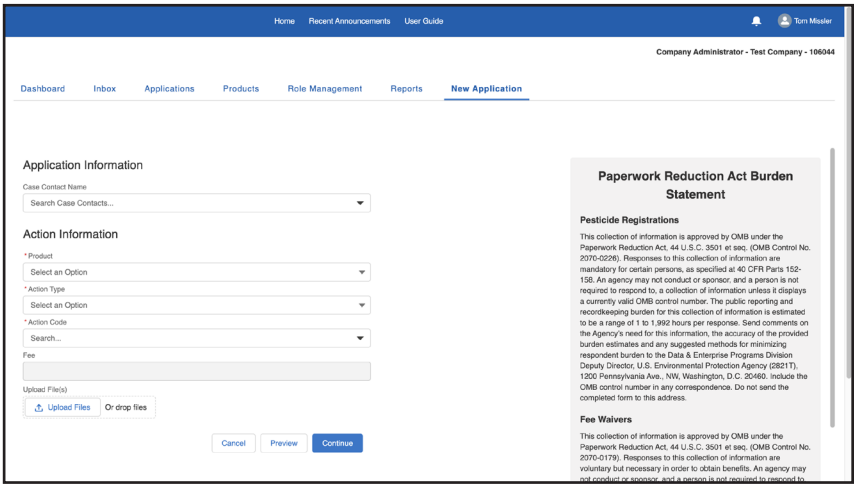
If the **Fee Waiver** field appears, the code you select here will apply to all the other applications included in this submission. Only add one Fee Waiver per submission.

3. Click **Add Application.**

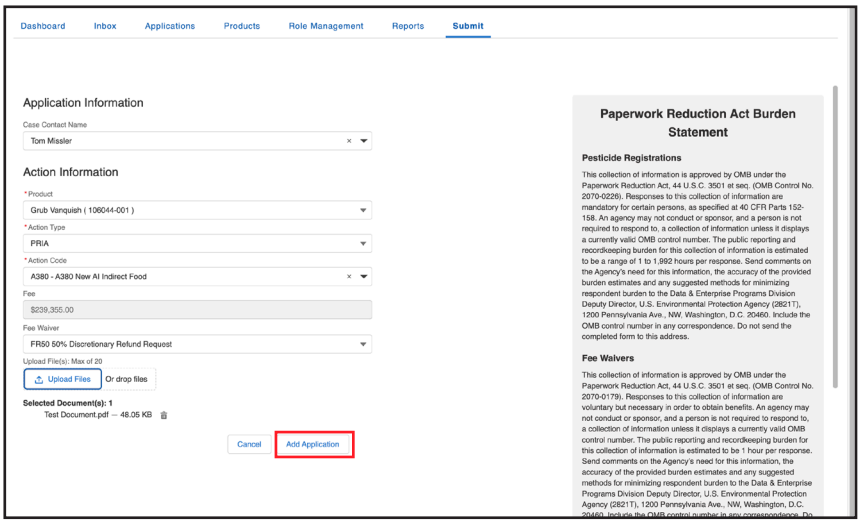
The screen you see if you click the Make A Submission or the New Application tab.



The fields you first see when you click the Start New Submission button.



In this example the product is Grub Vanquish, the Action Type is PRIA and a 50% Fee Waiver is requested and a document has been uploaded for this application.

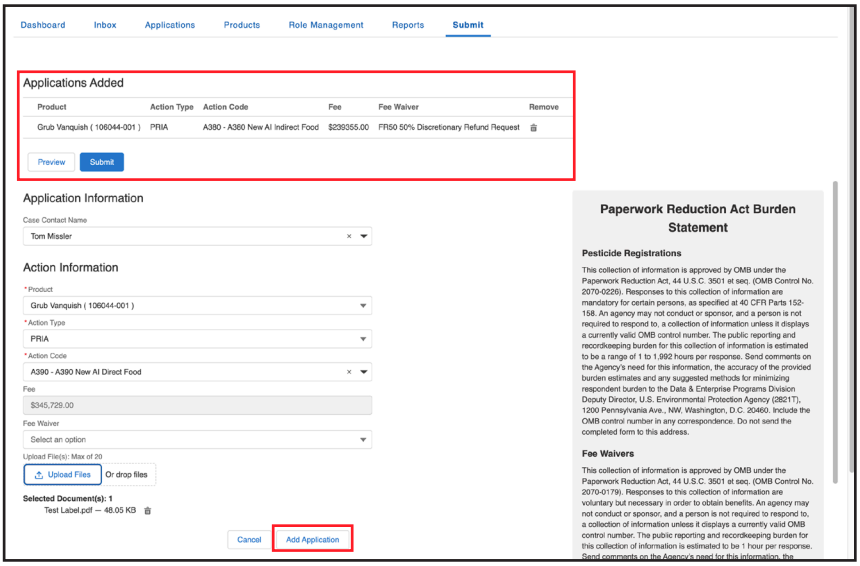


4. The **Application Added** section will appear above the Application Information section showing the product, action type, action code, fee, and fee waiver you just added.
5. To add an additional application to this submission, enter data for:
   - **Case Contact Name**
   - **Product**—Must be the same product you used in step 2.
   - **Action Type**—Must be a PRIA action
   - **Action Code**
   - **Fee**
   - Upload File(s)

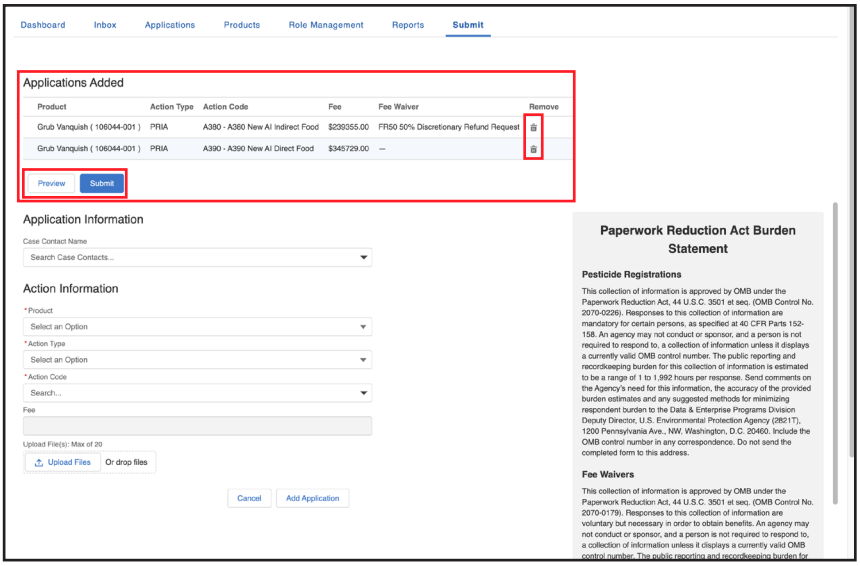
   If the Fee Waiver field appears and you have already entered a Fee Waiver, do not select anything for this application.
6. Click **Add Application.**
7. The application you just created will appear in the Applications Added section.
8. If you would like to remove an Application before submitting, click the **trashcan icon** to remove it.
9. If you would like to view a summary of your application, click **Preview** which will download an html file with the details of what you are submitting. You can open the html file in your web browser.
10. Click **Submit.**

Congratulations! You have successfully submitted both applications to the EPA.

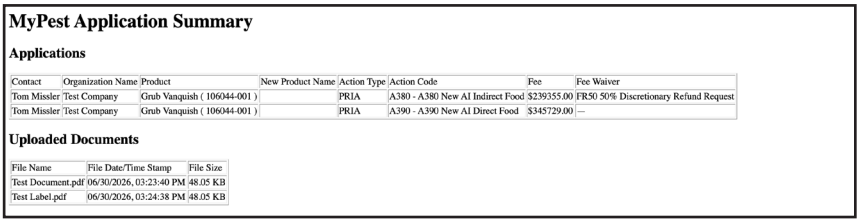
The Applications Added section is now visible. For the second application you must use the same product and the Action Type must be set to PRIA. The Fee Waiver is left blank because you can only submit one fee waiver per submission and one was included with the first application in this submission.



The Applications Added section shows both applications that will be submitted. If you would like to remove an application, click the trashcan icon. If you would like to see a summary of your submission, click Preview. If everything looks good, click Submit to send both applications to the EPA.



Clicking Preview will generate an html file that will automatically download to your computer. You can open it in your web browser.



MyPest Application Summary

Applications

Contact	Organization Name	Product	New Product Name	Action Type	Action Code	Fee	Fee Waiver
Tom Missler	Test Company	Grub Vanquish (106044-001)		PRIA	A380 - A380 New AI Indirect Food	$239355.00	FR50 50% Discretionary Refund Request
Tom Missler	Test Company	Grub Vanquish (106044-001)		PRIA	A390 - A390 New AI Direct Food	$345729.00	—

Uploaded Documents

File Name	File Date/Time Stamp	File Size
Test Document.pdf	06/30/2026, 03:23:40 PM	48.05 KB
Test Label.pdf	06/30/2026, 03:24:38 PM	48.05 KB

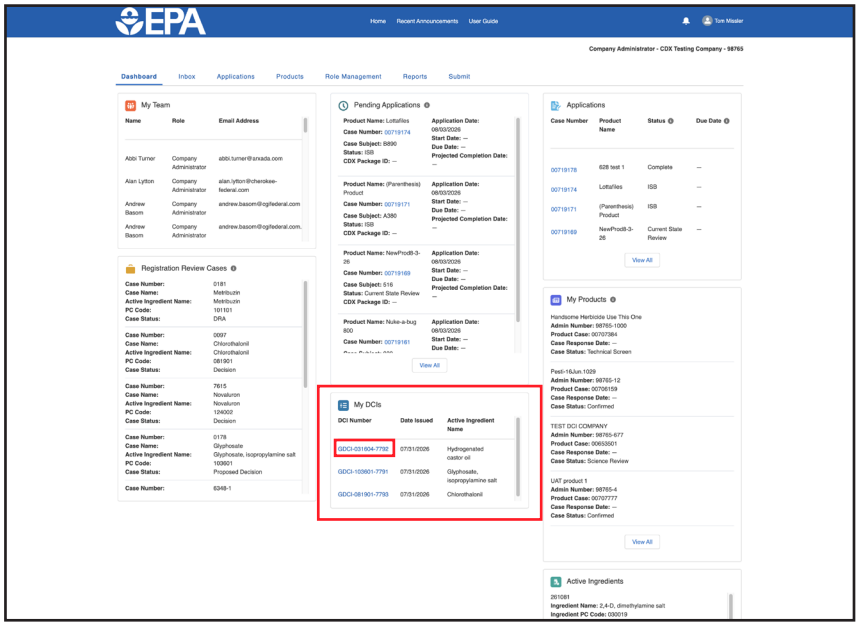
Responding to a Data Call-In (DCI)

It is possible to submit documents (studies, letters, follow-up documentation) and 90-day responses to DCI through MyPeST.

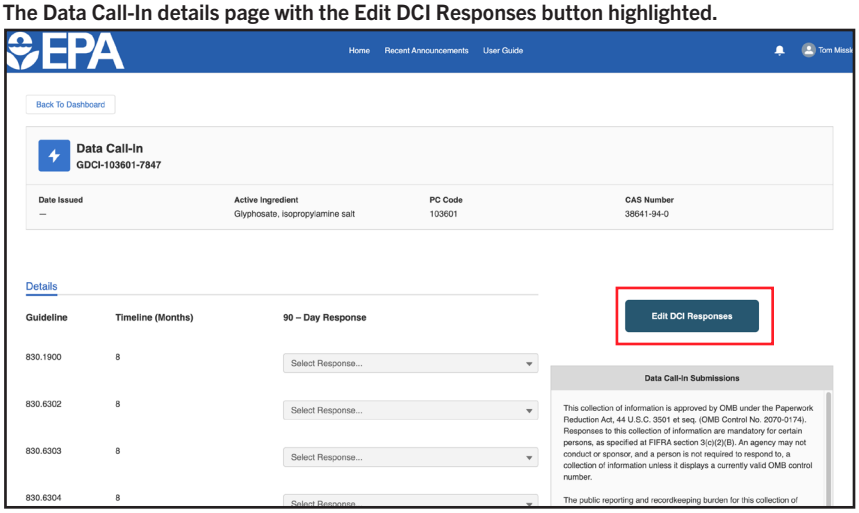
1. On the Dashboard in the center column at the bottom you will find the My DCIs tile. You may need to scroll down to see it. The My DCIs tile displays:
   - **DCI Number**
   - **Date Issued**
   - **Active Ingredient Name**
2. Click the **DCI Number hyperlink** for the DCI for you want to submit responses for.
3. The header on the Data Call-In details page shows:
   - **Date Issued**
   - **Active Ingredient**
   - **PC Code**
   - **CAS Number**

   The Details section shows:
   - **Guideline**—For a complete list of guidelines visit https://www.epa.gov/sites/default/files/2019-10/documents/ocspp-testguidelines_masterlist-2019-09-24.pdf
   - **Timeline (Months)**—Set by EPA and is based on the guideline requirement
   - **90-Day Response**—how your company intends to respond to the DCI
4. Click the **Edit DCI Responses** button

The Dashboard with the My DCIs tile and the DCI Number highlighted.



The Data Call-In details page with the Edit DCI Responses button highlighted.



5. The **Edit DCI Responses** button will dim, **Upload Files or drop files** will appear, and two active buttons (**Cancel** and **Preview**) and one inactive button (**Submit**) will appear, and and the dropdown lists under 90-Day Response will become active.

6. At this point you can do one of three things:

   1) Select a 90-Day response for any or all guidelines and Submit without uploading documents
   2) Select a 90-Day response for any or all guidelines, upload documents and Submit
   3) Upload documents without selecting a 90-Day response

7. There are 11 options in the 90-Day Response dropdown:
   - **Developing Data**
   - **Agreement to Cost Share**
   - **Offer to Cost Share**
   - **Submitting Existing Study**
   - **Upgrading a Study**
   - **Citing a Study**
   - **Deleting Uses**
   - **Low Volume/Minor Use Waiver Request**
   - **Request to Waive Data**
   - **Generic Data Exemption**
   - **Voluntary Product Cancellation**

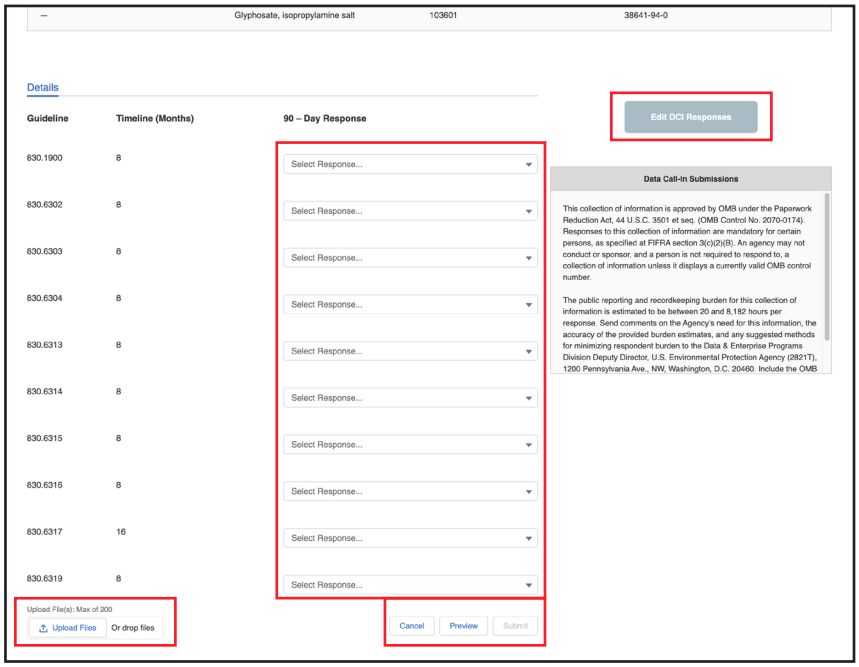
   You will need to make a selection for each guideline separately.

8. If you would like to upload a document, you can drag and drop your files or you can click the Upload Files button and select documents by navigating to the location on your hard drive.

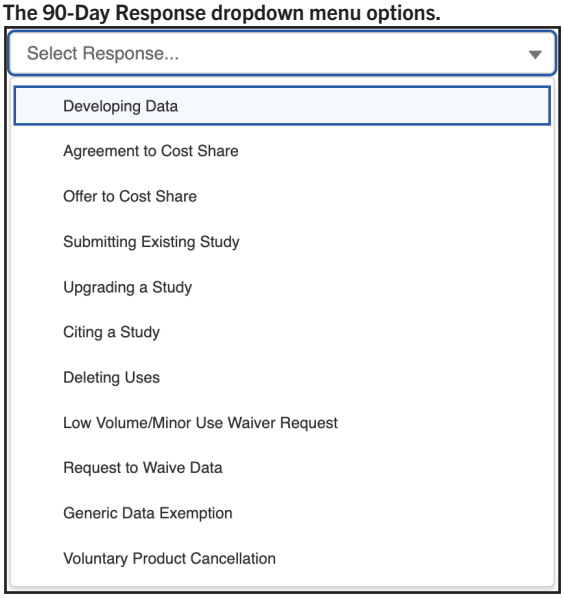
   The accepted file types are:
   - .pdf (Adobe Acrobat)
   - .doc (Microsoft Word 1997–2003)
   - .docx (Microsoft Word 2003–Present)
   - .xls (Microsoft Excel 1997–2003)
   - .xlsx (Microsoft Excel 2003–Present)

   There is a 200 document limit and 200MB per document size limit for uploads. You are not required to upload any documents with your DCI response.

The Data Call-In details page with the Edit DCI Responses button disabled, the 90-Day Response dropdown meus active, Upload Files and the Cancel, Preview, and Submit buttons visible.



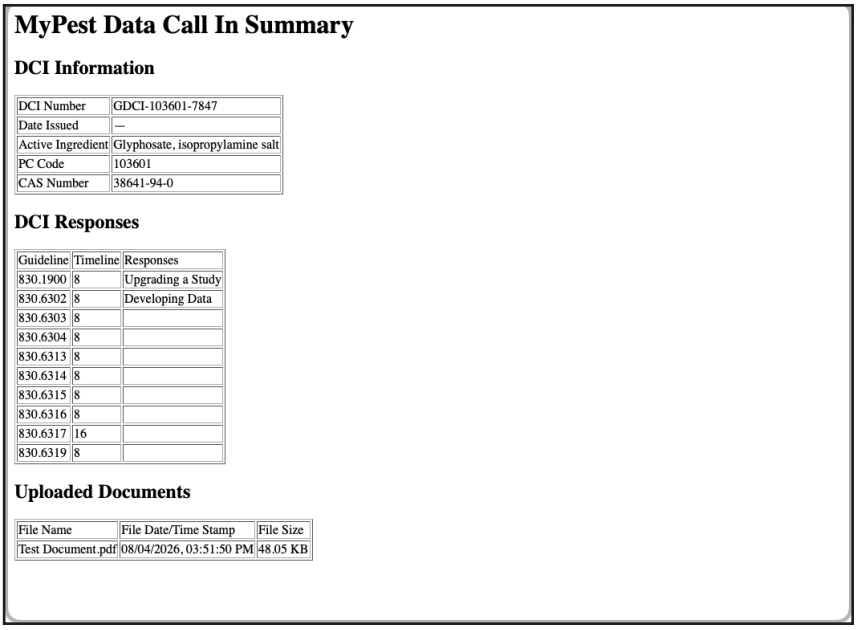
The 90-Day Response dropdown menu options.



9. Once you have selected your 90-Day Responses and uploaded your documents you can click **Preview** to see a receipt summarizing the information you provided. This is an optional step.
10. Click **Submit** to register your responses and submit your documents.
11. The **Create Passphrase & Review** popup will appear. Your passphrase must meet the following criteria:
    - At least 8 characters long
    - No more than 20 characters
    - Spaces are allowed
    - No special characters (!,#, ?, etc.)
    - Case sensitive

    When your **Passphrase** and **Confirm Passphrase** field match the Continue button will become active. If you need to see what you typed, click the **eyeball icon** in the field.
12. Click **Continue.**
13. The **Digital Signature Screen** popup will appear. Click **Accept** if you certify, under penalty of law that the information provided in this document is, to the best of your knowledge and belief, true, accurate, and complete. You are aware that there are significant penalties for submitting false information, including the possibility of fines and imprisonment for knowing violations. If not click **Decline** and cancel your submission.

An example of the receipt that is downloaded when you click the Preview button.



MyPest Data Call In Summary

DCI Information

DCI Number	GDCI-103601-7847
Date Issued	—
Active Ingredient	Glyphosate, isopropylamine salt
PC Code	103601
CAS Number	38641-94-0

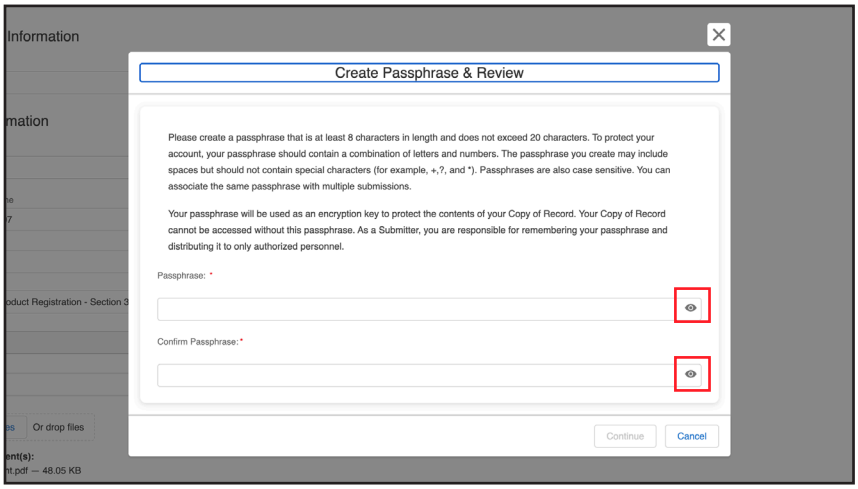
DCI Responses

Guideline	Timeline	Responses
830.1900	8	Upgrading a Study
830.6302	8	Developing Data
830.6303	8	
830.6304	8	
830.6313	8	
830.6314	8	
830.6315	8	
830.6316	8	
830.6317	16	
830.6319	8	

Uploaded Documents

File Name	File Date/Time Stamp	File Size
Test Document.pdf	08/04/2026, 03:51:50 PM	48.05 KB

The Create Passphrase & Review popup.

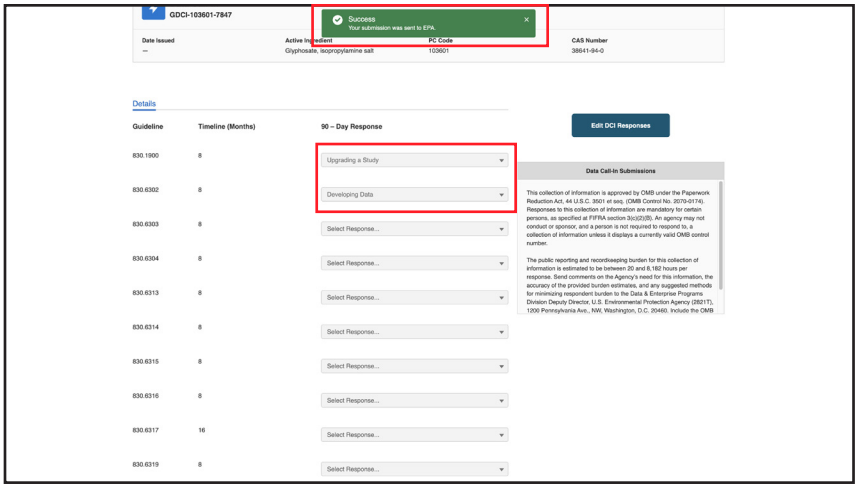


The Digital Signature Screen. Clicking Decline here will cancel your submission.

14. If you clicked Accept you will return to the DCI details page. You will see a green Success message appear to indicate that your submission was sent to the EPA and the 90-Day responses you selected will appear next to the guidelines they apply to.

Congratulations! You have successfully completed a DCI submission to the EPA.

The DCI details page showing the success message indicating your response was received by EPA and the responses appear next to the applicable guidelines.



Appendix A—The Role Management Tab for Company Administrators Only

Pending Requests, Pending Invites, and Decisions

At the top of the Role Management Tab page you will see a search bar. You can search for terms in any of the fields that are displayed in the table. As you type, the table will filter results. In most cases, you will only need to type a few letters before you find the person you are looking for.

On the Role Management Tab you will see a table with three sections: **Pending Requests, Pending Invites,** and **Decisions.** Under each section you will see 6 columns:

- **User:** The requestor's name.
- **Email:** The requestor's email.
- **Role:** The role the requestor is seeking approval for.
- **Status:** Can be Pending, Approved, Denied, or Revoked.
  - **Pending:** No decision has been made about the request.
  - **Approved:** The requestor has been granted the role.
  - **Denied:** The requestor has not been granted the role. They can reapply for a role, but this requested is closed.
  - **Revoked:** The requestor was originally approved for a role but they no longer need access to this company.
- **Down Arrow Buttons:** Click these buttons to Approve, Deny, or Revoke a user's access.

The table is broken down into three sections:

Pending Requests

Users have sent requests to the company for a role. The Company Administrator can **Approve** or **Deny** by clicking the **Down Arrow Button** and selecting from the drop-down options. If you select **Deny**, you will see a popup asking you to confirm your decision. If you click **Approve,** there is no popup.

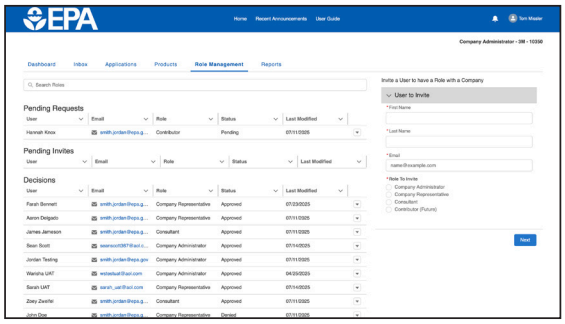
Pending Invites

The company has sent a request to the user for a role. The Company Administrator cannot take any further action on this request.

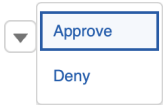
Decisions

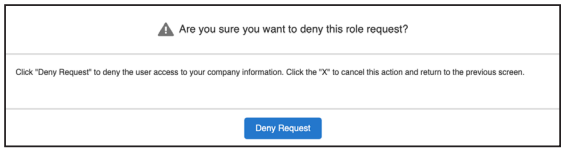
The list of all users who have been approved, denied, or revoked for a role. The Company Administrator can revoke a role for an approved role in this section by clicking the **Down Arrow Button** and selecting **Revoke.** If you select **Revoke,** you will see a popup asking you to confirm your decision.

Role Management Tab

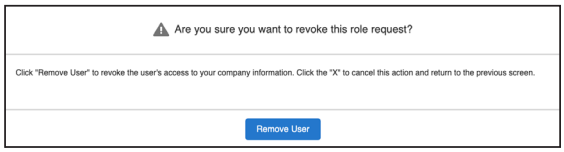


Down Arrow Button with Approve or Deny options in the Pending Requests section





Down Arrow Button with the Revoke option in the Decisions section



Invite A User

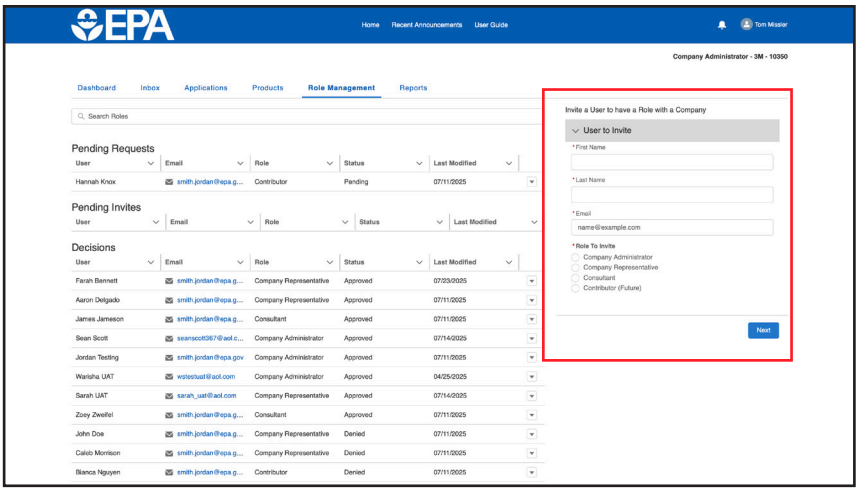
As a Company Administrator you have the ability to invite people to have a role with your company. It works the same way as Requesting a Role, only in reverse. The Company Administrator enters the user's:

- First Name
- Last Name
- Email
- Role

When the Company Administrator clicks **Next,** an invitation is sent to the specified user. They can Accept or Reject the invitation. The Company Administrator will receive an email indicating the user's decision.

If a user rejects a role, they can subsequently request that role or a different role with the company or be invited again for that role or a different one. They are not "locked out" if they refuse an invitation.

If you try to send an invitation to a user who has not yet created an account in MyPeST, you have the option of having an auto-generated message sent to the email address you specified in the invite asking the person to create an account. Once that user has created an account, you can invite them or they can make a request to join your company.



Changing Roles

If a user accepted a role with your organization but now you want them to have a different role, the Company Administrator has the ability to make that change.

1. Find the user's name in the Decisions section of the Role Management tab and select **Revoke** in the drop-down.
2. The CA can now invite the person to take on a new role with the organization since they no longer have a role.
3. When the user accepts the invitation they will be back in the organization with their new permissions and responsibilities based on their new role.

Appendix B—Subject Codes

Code	Description
10	dng 75d
110	dng 75d
117	dng 75d
152	dng 75d
160	dng 75d
163	dng 75d
165	dng 75d
170	dng 75d
172	dng 75d
175	dng 75d
177	dng 75d
184	dng 75d
186	dng 75d
188	dng 75d
192	dng 75d
194	agency-initiated
220	Tolerance
230	Tolerance
268	general correspondence
274	dng 75d
300	amendment label/csf
302	label amend
305	data required
307	data required
312	dng 75d
315	dng 75d
317	dng 75d
320	Amendment (EVP)
322	Amendment
325	dng 75d
330	dng 75d
331	Notification (Covid)
332	notification
335	Use deletion
341	csf amend
345	csf amend
350	general correspondence
352	protocol
354	lab audit
356	label improvement
358	agency-initiated
360	agency-initiated
362	csf amend
385	Special packaging
392	mfa
397	mfa
400	no data required
405	6a2
415	final printed label
455	enforcement case review
457	ATP issue
458	ATP issue
460	Cancellation
462	Use deletion
500	s18 unredg ai
505	s18 renewal
510	s18
513	s18 quar exempt amend
535	s18 quar exempt
570	registration follow-up
575	registration follow-up
576	pet spot-on
577	pet spot-on
578	pet spot-on
579	pet spot-on
580	24c revision
582	24c amend
585	24c
587	24c amend
627	DCI response
676	reregistration decision
677	reregistration decision
678	reg review label
679	reg review label
681	biological opinion mitigation
709	EUP report
714	dng 75d
736	EUP extension
754	EUP amendment
790	Novel microbial notification
888	agency-initiated
900	Cancellation

Appendix C—Troubleshooting Company Numbers in MyPeST

You work for a company that has been submitting documents through CDX for years. Now that you are using MyPeST you are ready to log in and get started. When trying to request a role you search for your company in MyPeST and you see:

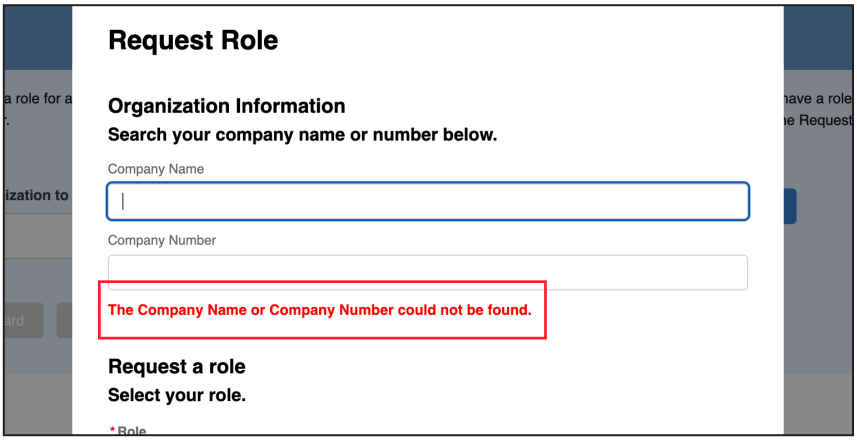
The Company Name or Company Number could not be found.

What is happening here?

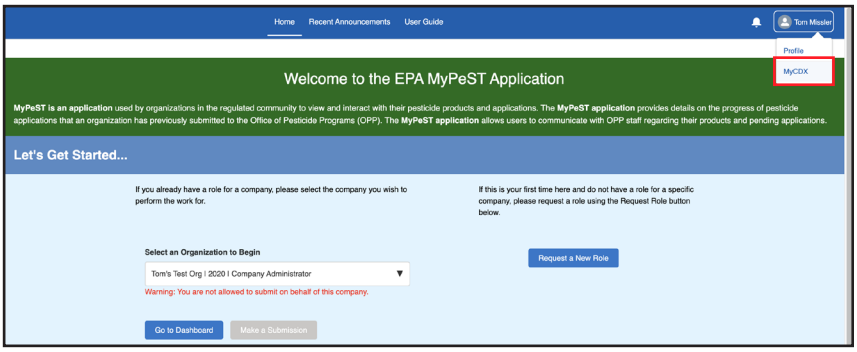
CDX does not require a company number in order to submit documents. MyPeST requires every company to have a company number. You will need to request a company number in CDX, and once it is established, you will be able to request a role in MyPeST.

How do I fix it?

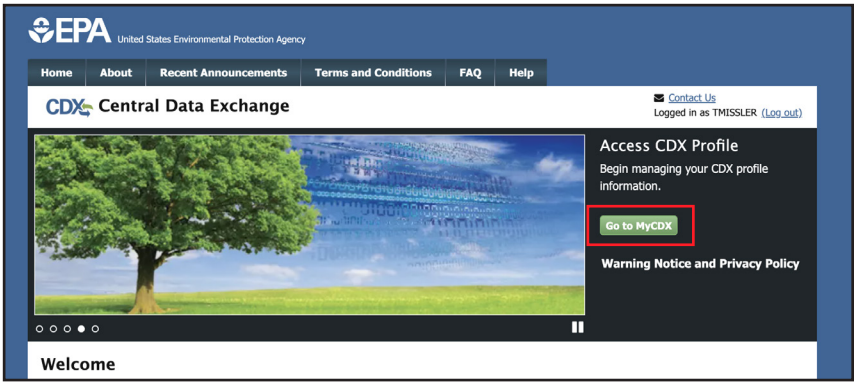
1. Under your profile select **MyCDX.** This will take you out of MyPeST and into CDX.
2. On the Central Data Exchange page click the **Go to MyCDX** button.
3. On the Central Data Exchange page click **Add Program Service.**



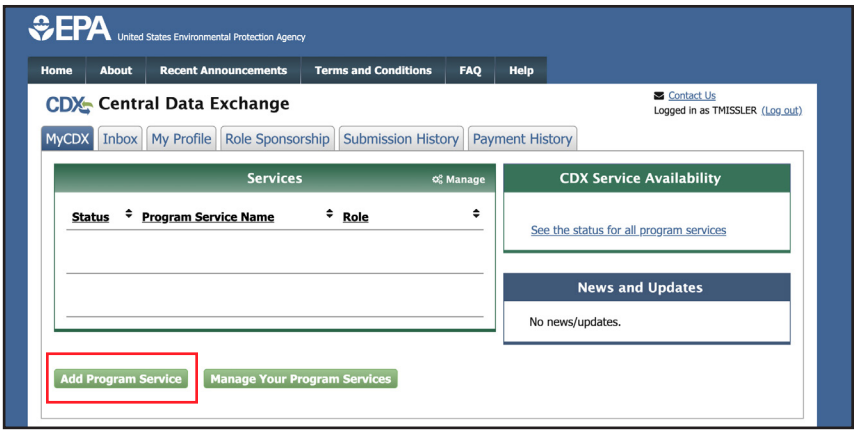
Step 1



Step 2

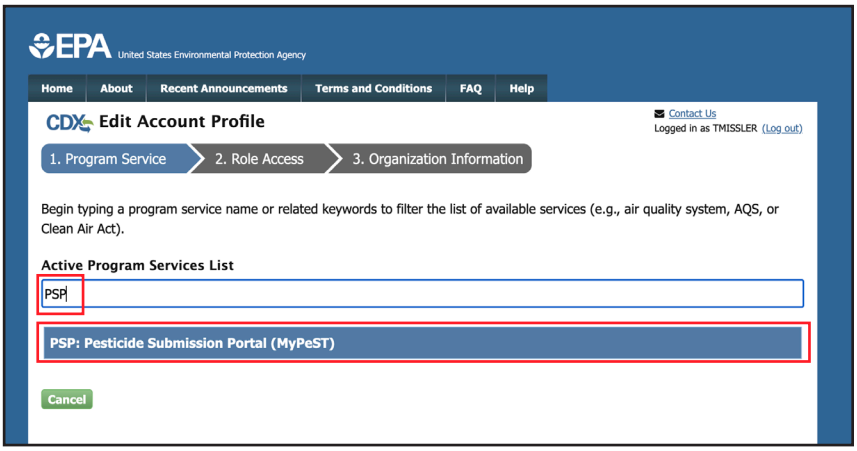


Step 3

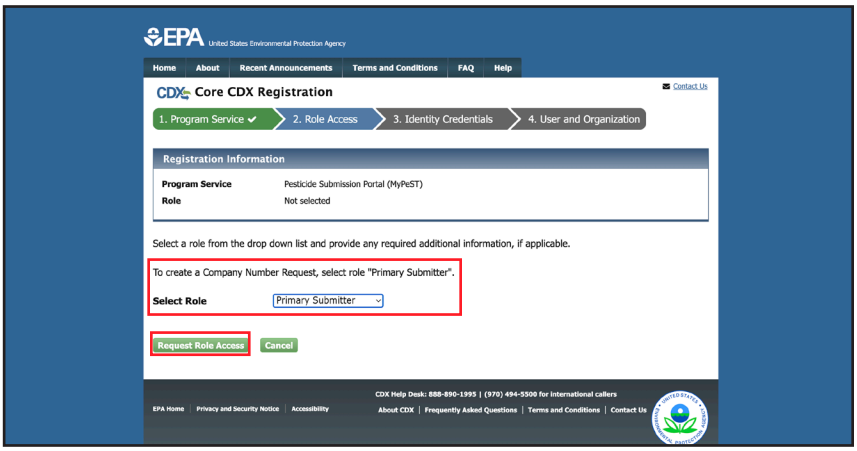


4. In the Active Program Services List type “PSP” then select **PSP: Pesticide Submission Portal (MyPeST).**
5. In the Select Role drop-down, select **Primary Submitter** and click the **Request Role Access** button.
6. Click the **I don’t have a company number yet hyperlink.**

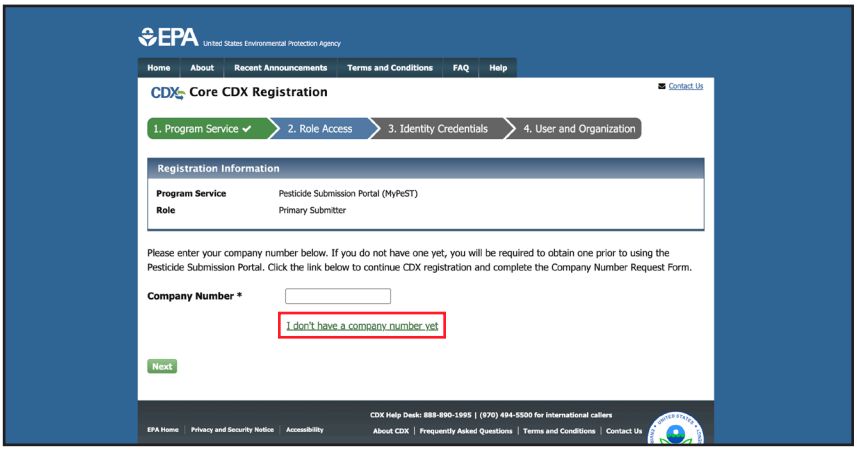
Step 4



Step 5



Step 6

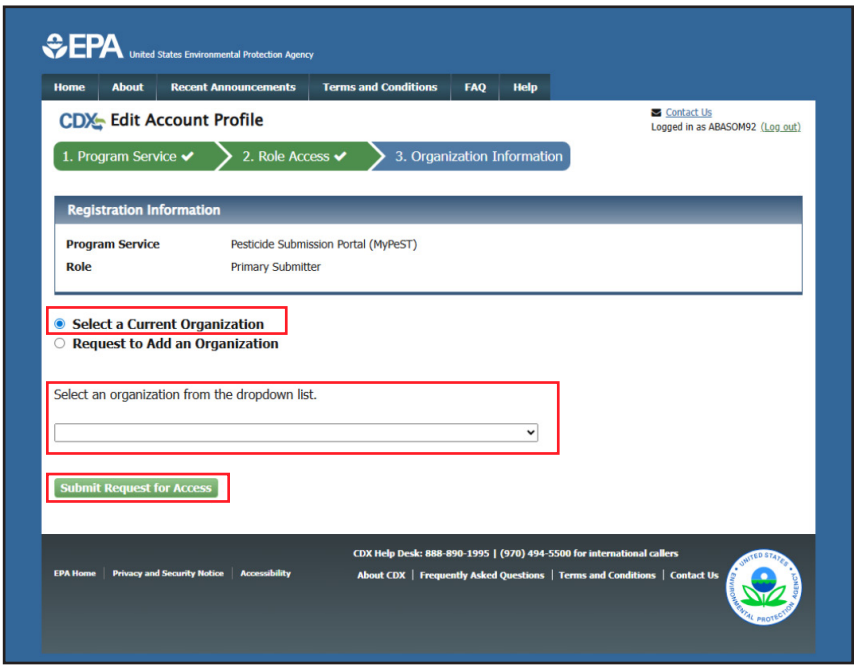


7. Make sure the **Select a Current Organization** radio button is selected. Click the **Select an organization from the drop-down list** and select your company. Then click **Submit Request for Access.**

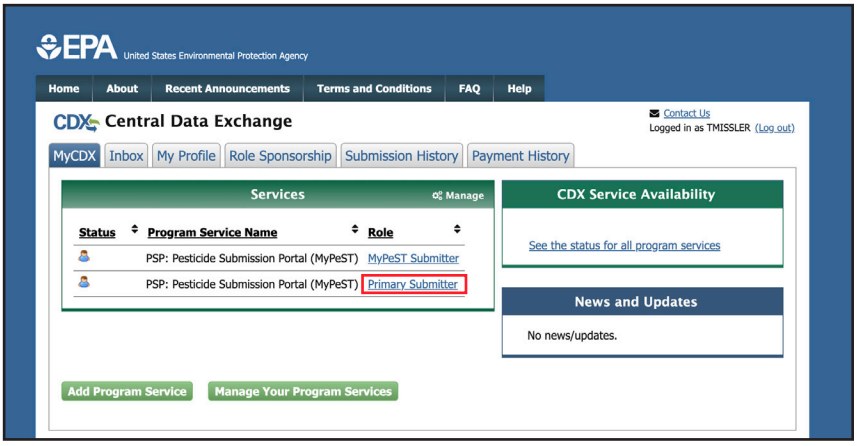
8. You will be returned to the MyCDX page. Click the **Primary Submitter hyperlink.**

8a. If you are the Primary Submitter for more than one company, the Application Profile Settings popup will appear. Select the same organization you selected in step 7 then click **Proceed.** If you are the Primary submitter for only one company, you will not see this popup and will go to Step 9.

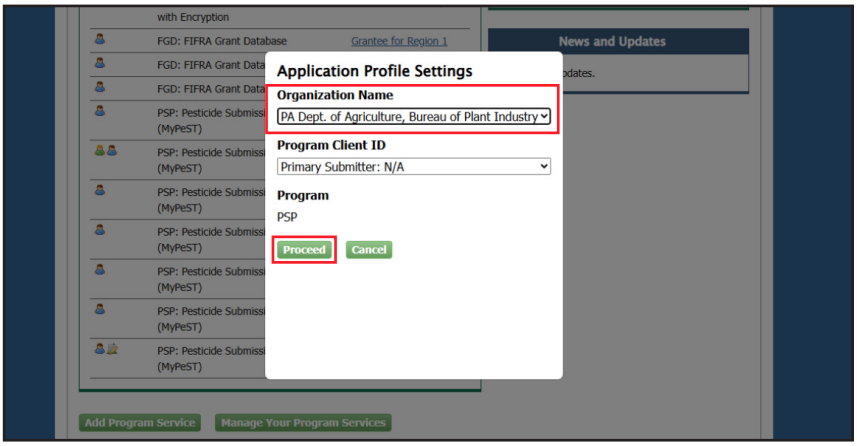
Step 7



Step 8

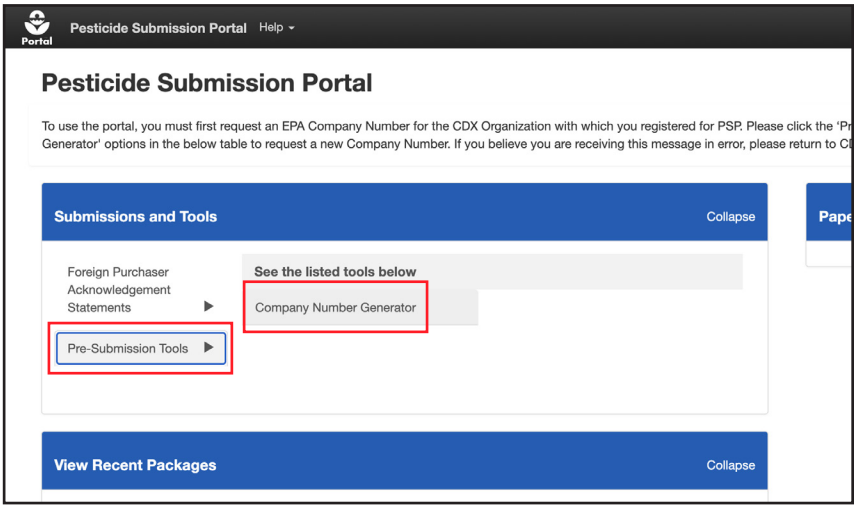


Step 8a—You will only see this if you are Primary Submitter for more than one company.

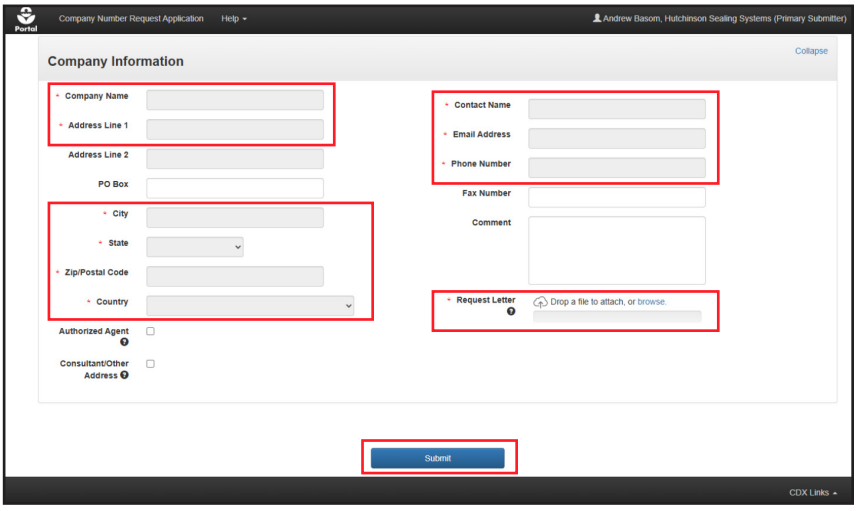


9. You will see the Pesticide Submission Portal page. Click Pre-Submission Tools then click Company Number Generator.
10. The Company Information screen will appear. Enter data for all fields with an asterisk. You must include a **Request Letter.** The letter must have the company letterhead with your corporate contact details (contact person's name, email address, phone number, and official mailing address). Click **Submit.**
11. You will be notified when your company number has been approved. Once you have received notification, log in to MyPeST and click **Request a New Role.** From here, the process follows the same steps outlined begining on page 8.

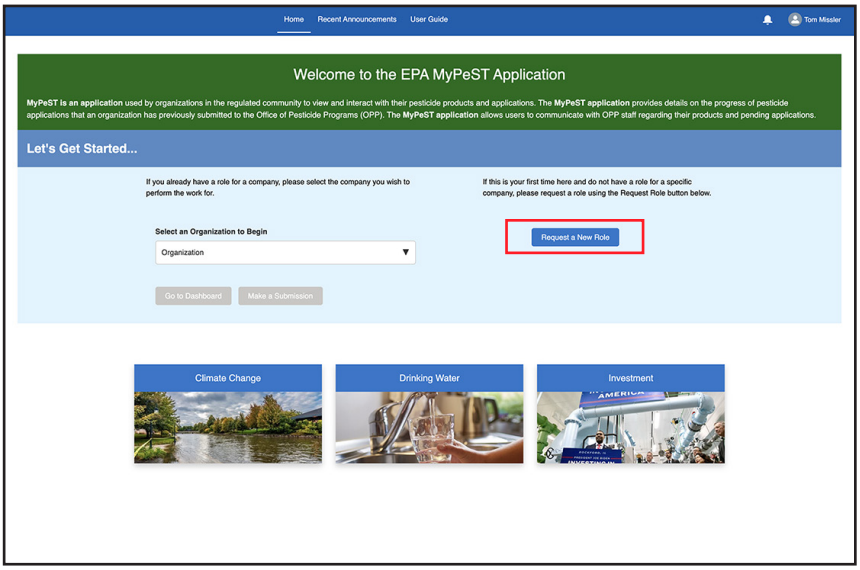
Step 9



Step 10



Step 11

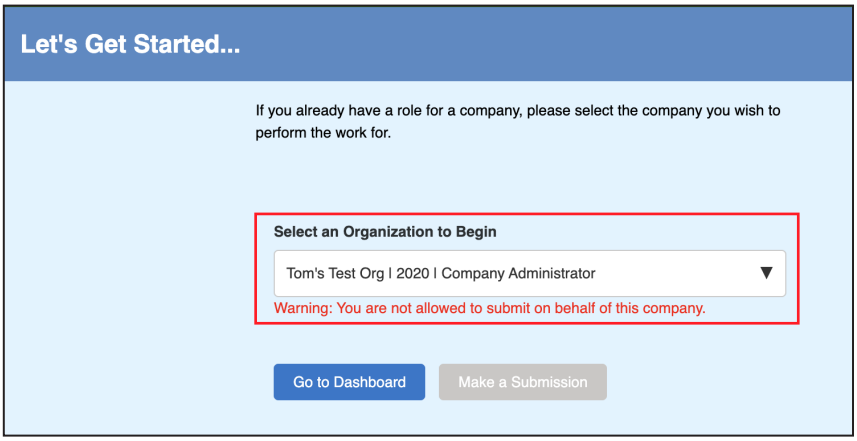


Appendix D–Troubleshooting Role Requests in CDX

You have successfully signed in to MyPeST, requested a role, submitted the required documentation (if necessary), and have been granted the role you requested. If you requested to be a Company Administrator you should be able to see and do everything in MyPeST.

When you select your company to begin working in MyPeST you see this error on the home screen:

Warning: You are not allowed to submit on behalf of this company.



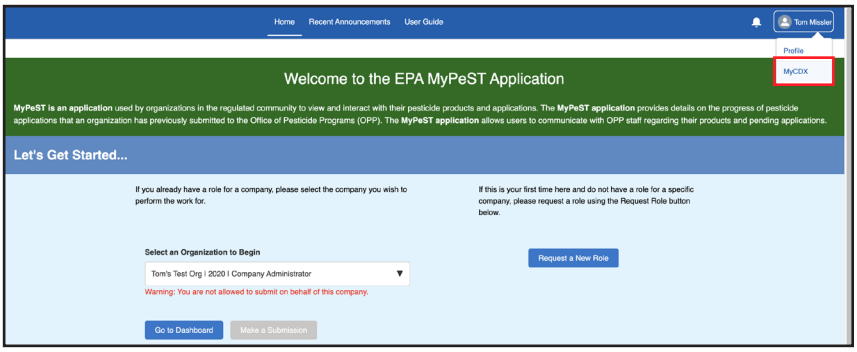
What is happening here?

From MyPeST's perspective you are a Company Admin and have all the permissions to view and submit on behalf of your company. The problem is you don't have the same permissions in CDX.

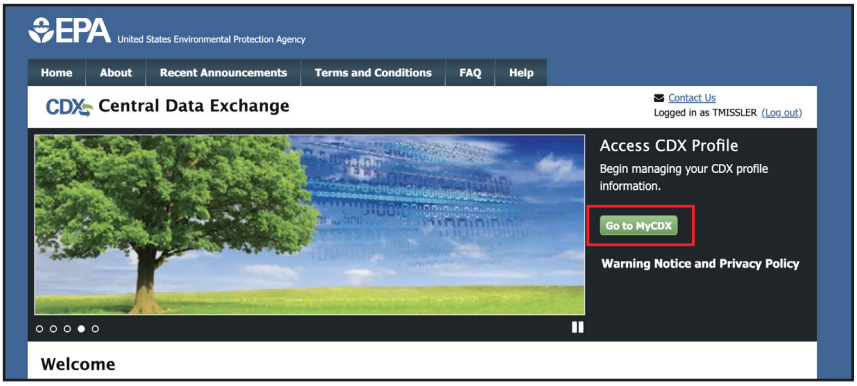
How do I fix it?

1. Under your profile select **MyCDX.** This will take you out of MyPeST and into CDX.
2. On the Central Data Exchange page click the **Go to MyCDX** button.
3. On the Central Data Exchange page click **Add Program Service.**

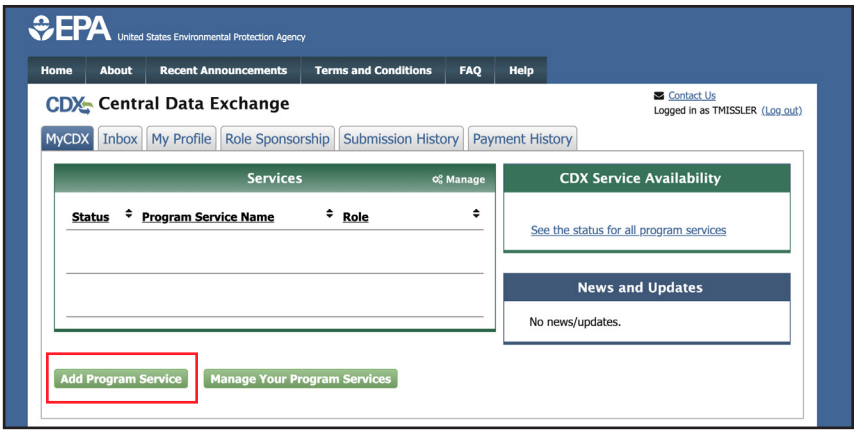
Step 1



Step 2

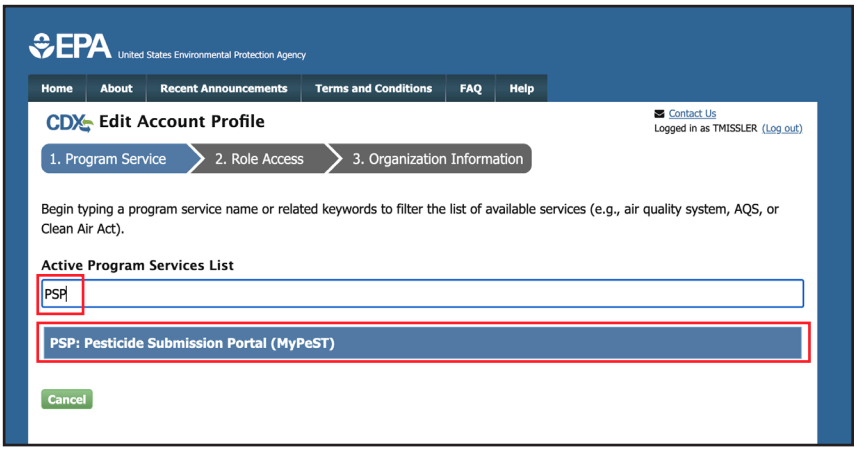


Step 3

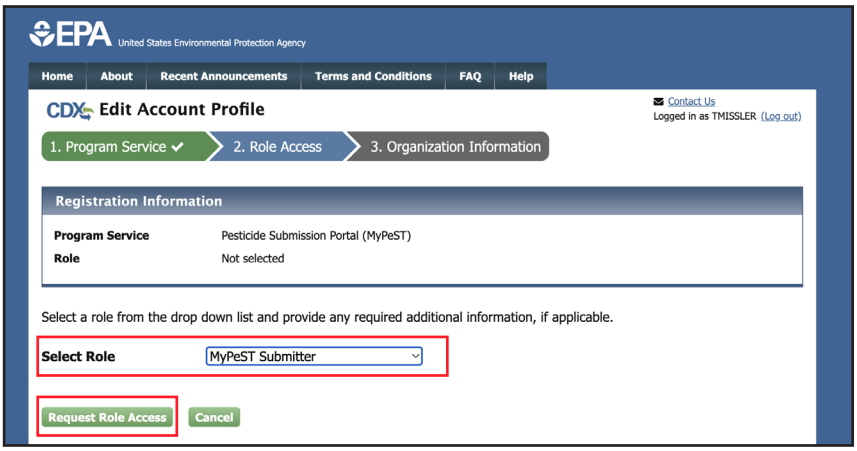


4. In the Active Program Services List type “PSP” then select **PSP: Pesticide Submission Portal (MyPeST).**
5. In the Select Role drop-down, select **MyPeST Submitter** and click the **Request Role Access** button.
6. Search for it by typing the name and clicking **Search.** If you see your company in the search results, click the **Organization ID hyperlink** next to your Organization Name. If your company does not appear in the list, click **request that we add your organization hyperlink** and follow the process to add your company to CDX.

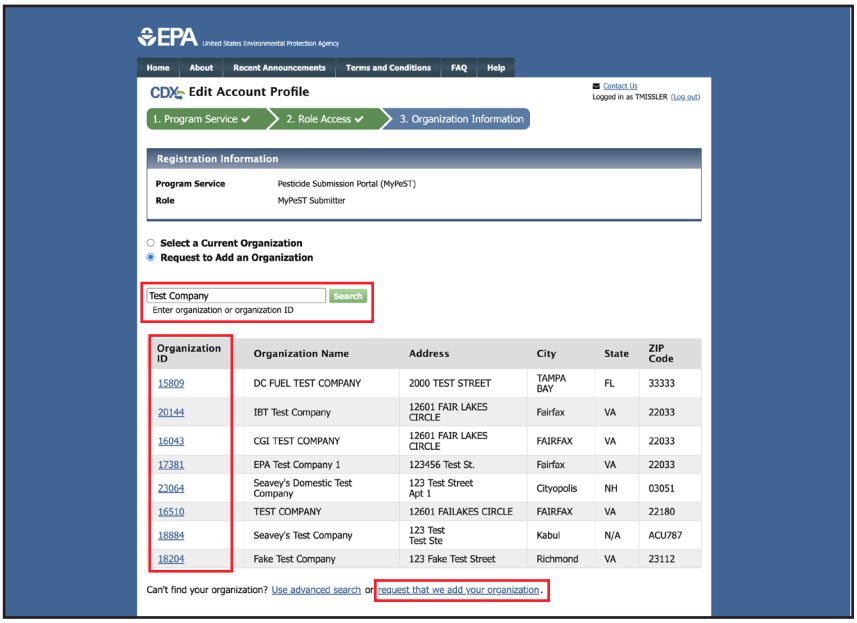
Step 4



Step 5



Step 6

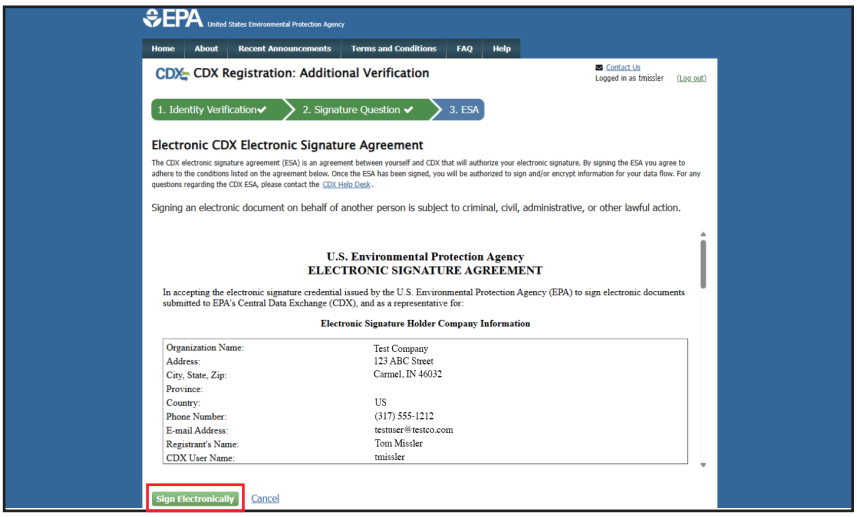


7. You will now see the **Electronic CDX Electronic Signature Agreement** page. Verify the information is correct (some companies have multiple branches registered with CDX. Make sure you have selected the right one. Once you are sure, click **Sign Electronically.**
8. You will return to the Central Data Exchange page. In your list of services you will see **PSP: Pesticide Submission Portal (MyPeST)** under the Program Services Name and **MyPeST Submitter** in the Role. Click the **MyPeST Submitter hyperlink.**
9. This will launch the MyPeST application. Now when you click the organization on the Home Screen, the warning will no longer appear.

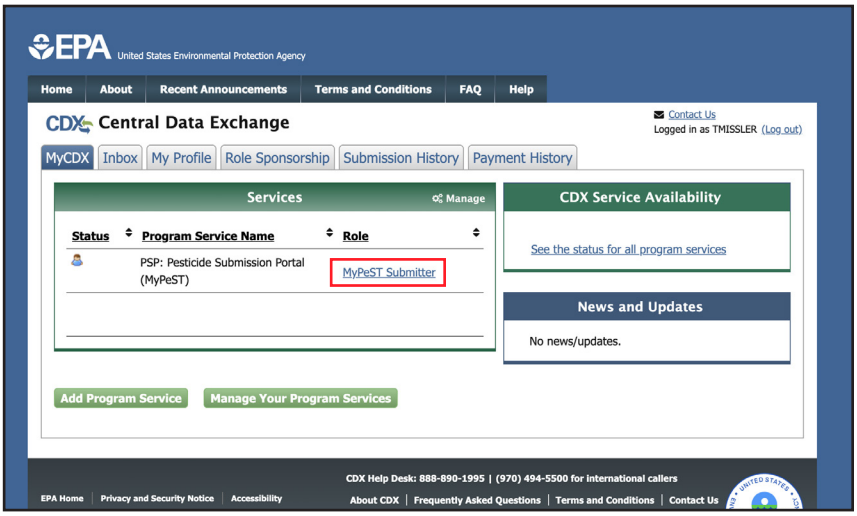
Select An Organization

After you have been accepted for a role, the next time you log in you will see the **Select an Organization to Begin** drop-down is active on the Home screen. Click the drop-down and select
your organization from the list. You can click **Go to Dashboard** to see the status of your EPA applications or you can click **Make a Submission** to start a new product registration or update an existing application. We will start with **Go to Dashboard.**

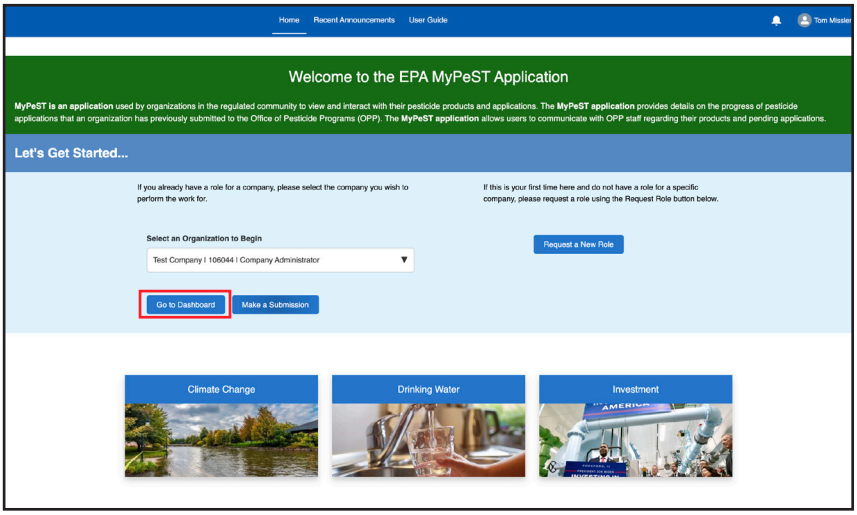
Step 7



Step 8



Step 9 and Select An Organization



Appendix E—Action Codes

Below is a list of all action codes for the 2026 fiscal year.

Code	Category	Type	Fee $	Review Time	Time Type	Description
165	Other	Non-PRIA		190	Days	165 “Not Grant” Follow-up
270	Other	Non-PRIA		90	Days	270 Inert Tradename Clearance Request
300	Amendment	Non-PRIA		90	Days	300 Fast-Track Label Amendment
332	Notification	Non-PRIA		30	Days	332 Notification
335	Cancellation	Non-PRIA		210	Days	335 6(f) Voluntary Use Termination
345	Amendment	Non-PRIA		90	Days	345 Fast-Track Formulation Amendment
350	Other	Non-PRIA		110	Days	350 Miscellaneous (regulatory review only)
360	Other	Non-PRIA		180	Days	360 Agency-Initiated Action
392	Amendment	Non-PRIA		45	Days	392 Minor Formulation Amendment
400	Other	Non-PRIA		110	Days	400 Miscellaneous (science review needed)
405	Incident	Non-PRIA		90	Days	405 6(a)(2) Incident Report with No Study
415	Other	Non-PRIA		190	Days	415 Final Printed Label
506	18/24c	Non-PRIA		45	Days	506 Section 18 New (Specific) Request
516	18/24c	Non-PRIA		45	Days	516 Section 18 New (Quarantine) Request
526	18/24c	Non-PRIA		45	Days	526 Section 18 New (Public Health) Request
536	18/24c	Non-PRIA		45	Days	536 Section 18 New (Crisis) Request
546	18/24c	Non-PRIA		45	Days	546 Section 18 Amendment/Follow-up
556	18/24c	Non-PRIA		45	Days	556 Section 18 Final Report
570	Data	Non-PRIA		110	Days	570 Terms/conditions follow-up product-specific data review only
575	Data	Non-PRIA		110	Days	575 Terms/conditions follow-up generic AI data review needed
576	Data	Non-PRIA		110	Days	576 Pet Spot-On Request for Yearly Reporting
577	Data	Non-PRIA		110	Days	577 Pet Spot-On Quarterly Enhanced Incident Report
578	Data	Non-PRIA		110	Days	578 Pet Spot-On Quarterly Sales Information
579	Data	Non-PRIA		110	Days	579 Pet Spot-On Annual Enhanced Sales and Incident Data
580	18/24c	Non-PRIA		90	Days	580 Section 24(c) New Registration (SLN)
582	18/24c	Non-PRIA		90	Days	582 Section 24(c) Amendment (SLN)
625	Incident	Non-PRIA		180	Days	625 6(a)(2) Incident Report with Study Submission
676	Re-evaluation	Non-PRIA		180	Days	676 Product Reregistration Follow-up
678	Re-evaluation	Non-PRIA		180	Days	678 Registration Review Label Amendment Pre-Decisional Mitigation
679	Re-evaluation	Non-PRIA		180	Days	679 Registration Review Label Amendment

Code	Category	Type	Fee $	Review Time	Time Type	Description
680	Re-evaluation	Non-PRIA		180	Days	680 Registration Review Label Amendment Pre-BiOp Mitigation
681	Re-evaluation	Non-PRIA		180	Days	681 Registration Review Label Amendment BiOp Mitigation
736	Other	Non-PRIA		90	Days	736 Section 5 (Experimental Use Permit) Follow-up
800	Other	Non-PRIA		30	Days	Supplemental Distributor New Registration
801	Other	Non-PRIA		30	Days	Supplemental Distributor Add or Remove Name
802	Other	Non-PRIA		30	Days	Supplemental Distributor Cancel Registration
900	Cancellation	Non-PRIA		210	Days	900 6(f) Voluntary Product Cancellation
950	Other	Non-PRIA		90	Days	950 Biochemical Classification Determination (Pre-PRIA 5)
A380	New Active Ingredients	PRIA	$239,355.00	26	Months	A380 New AI Indirect Food
A380.0	New Active Ingredients	PRIA		26	Months	A380.0 Secondary A380 No fee
A390	New Active Ingredients	PRIA	$345,729.00	26	Months	A390 New AI Direct Food
A390.0	New Active Ingredients	PRIA		24	Months	A390.0 Secondary A390 No Fee
A410	New Active Ingredients	PRIA	$292,592.00	23	Months	A410 New AI Non-Food
A410.0	New Active Ingredients	PRIA		21	Months	A410.0 Secondary A410 No Fee
A431	New Active Ingredients	PRIA	$120,734.00	12	Months	A431 New AI Low Risk Nonfood
A431.0	New Active Ingredients	PRIA		12	Months	A431.0 Secondary A431 No Fee
A440	New Use	PRIA	$48,024.00	21	Months	A440 New Use Indirect Food
A440.0	New Use	PRIA		21	Months	A440.0 Secondary A440 No Fee
A440.1	New Use	PRIA	$12,006.00	21	Months	A440.1 Secondary A440 with Additional Fee
A441	New Use	PRIA	$172,871.00	21	Months	A441 New Use Indirect Food Six or More
A441.0	New Use	PRIA		21	Months	A441.0 Secondary A441 No Fee
A450	New Use	PRIA	$144,058.00	21	Months	A450 New Use Direct Food
A450.0	New Use	PRIA		21	Months	A450.0 Secondary A450 No Fee
A451	New Use	PRIA	$274,400.00	21	Months	A451 New Use Direct Food Six or More
A451.0	New Use	PRIA		21	Months	A451.0 Secondary A451 No Fee
A460	New Product	PRIA	$7,689.00	5	Months	A460 New Product 0 to 10 Public Health Organisms
A460.1	New Product	PRIA	$1,923.00	5	Months	A460.1 Secondary A460 with Additional Fee Product Chemistry Only
A460.2	New Product	PRIA	$7,689.00	5	Months	A460.2 Secondary A460 with Additional Fee Efficacy and/or Acute Tox
A461	New Product	PRIA	$10,666.00	6	Months	A461 New Product 11 to 20 Public Health Organisms
A461.1	New Product	PRIA	$2,667.00	6	Months	A461.1 Secondary A461 with Additional Fee Product Chemistry Only
A461.2	New Product	PRIA	$7,689.00	6	Months	A461.2 Secondary A461 with Additional Fee Efficacy and/or Acute Tox
A462	New Product	PRIA	$13,645.00	7	Months	A462 New Product 21 to 30 Public Health Organisms

Code	Category	Type	Fee $	Review Time	Time Type	Description
A462.1	New Product	PRIA	$3,412.00	7	Months	A462.1 Secondary A462 with Additional Fee Product Chemistry Only
A462.2	New Product	PRIA	$7,689.00	7	Months	A462.2 Secondary A462 with Additional Fee Efficacy and/or Acute Tox
A463	New Product	PRIA	$16,623.00	9	Months	A463 New Product 31 to 40 Public Health Organisms
A463.1	New Product	PRIA	$4,156.00	9	Months	A463.1 Secondary A463 with Additional Fee Product Chemistry Only
A463.2	New Product	PRIA	$7,689.00	9	Months	A463.2 Secondary A463 with Additional Fee Efficacy and/or Acute Tox
A464	New Product	PRIA	$19,602.00	10	Months	A464 New Product 41 to 50 Public Health Organisms
A464.1	New Product	PRIA	$4,901.00	10	Months	A464.1 Secondary A464 with Additional Fee Product Chemistry Only
A464.2	New Product	PRIA	$7,689.00	10	Months	A464.2 Secondary A464 with Additional Fee Efficacy and/or Acute Tox
A465	New Product	PRIA	$22,581.00	11	Months	A465 New Product More Than 50 Public Health Organisms
A465.1	New Product	PRIA	$5,646.00	11	Months	A465.1 Secondary A465 with Additional Fee Product Chemistry Only
A465.2	New Product	PRIA	$7,689.00	11	Months	A465.2 Secondary A465 with Additional Fee Efficacy and/or Acute Tox
A470	Amendment	PRIA	$5,768.00	4	Months	A470 Amendment 0 to 10 Public Health Organisms
A470.0	New Use	PRIA		15	Months	A470.0 Secondary A470 No Fee
A470.1	Pre-Application	PRIA	$1,442.00	4	Months	A470.1 Secondary A470 with Additional Fee
A471	Amendment	PRIA	$8,932.00	5	Months	A471 Amendment 11 to 20 Public Health Organisms
A471.0	Amendment	PRIA		5	Months	A471.0 Secondary A471 No Fee
A472	Amendment	PRIA	$10,730.00	6	Months	A472 Amendment 21 to 30 Public Health Organisms
A472.1	Amendment	PRIA	$2,683.00	6	Months	A472.1 Secondary A472 with Additional Fee
A473	Amendment	PRIA	$12,530.00	7	Months	A473 Amendment 31 to 40 Public Health Organisms
A474	Amendment	PRIA	$14,329.00	8	Months	A474 Amendment 41 to 50 Public Health Organisms
A475	New Product	PRIA	$16,555.00	9	Months	A475 Amendment More Than 50 Public Health Organisms
A500	New Use	PRIA	$48,024.00	15	Months	A500 New Use Nonfood
A500.0	New Use	PRIA		15	Months	A500.0 Secondary A500 No Fee
A500.1	New Use	PRIA	$12,006.00	15	Months	A500.1 Secondary A500 with Additional Fee
A501	New Use	PRIA	$115,253.00	15	Months	A501 New Use Nonfood Six or More
A520	Experimental Use Permits – Other Actions	PRIA	$9,609.00	9	Months	A520 Experimental Use Permit Nonfood
A520.0	Experimental Use Permits – Other Actions	PRIA		9	Months	A520.0 Secondary A520 No Fee

Code	Category	Type	Fee $	Review Time	Time Type	Description
A521	Pre-Application	PRIA	$7,115.00	6	Months	A521 Review of public health efficacy study protocol within AD
A522	Pre-Application	PRIA	$18,296.00	12	Months	A522 Review of public health efficacy study protocol outside AD
A523	Pre-Application	PRIA	$18,296.00	9	Months	A523 Review of protocol other than public health efficacy
A529	Experimental Use Permits – Other Actions	PRIA	$17,203.00	9	Months	A529 Experimental Use Permit Amendment
A530	New Product	PRIA	$1,925.00	4	Months	A530 New Product no data review or only product chemistry data
A531	New Product	PRIA	$2,747.00	4	Months	A531 New Product registered source only product chemistry data
A532	New Product	PRIA	$7,689.00	5	Months	A532 New Product unregistered source only product chemistry data
A533	Experimental Use Permits – Other Actions	PRIA	$3,737.00	4	Months	A533 Exemption from the requirement of an Experimental Use Permit
A534	Pre-Application	PRIA	$7,115.00	4	Months	A534 Rebuttal of agency reviewed protocol
A535	Pre-Application	PRIA	$3,627.00	6	Months	A535 Conditional Ruling on Study Waiver or Data Bridging Argument
A536	Pre-Application	PRIA	$3,737.00	4	Months	A536 Conditional Ruling on Direct Food, Indirect Food, Nonfood
A537	Experimental Use Permits – Other Actions	PRIA	$230,488.00	18	Months	A537 Experimental Use Permit New AI/ Use Direct Food
A537.0	Experimental Use Permits – Other Actions	PRIA		18	Months	A537.0 Secondary A537 No Fee
A538	Experimental Use Permits – Other Actions	PRIA	$144,058.00	18	Months	A538 Experimental Use Permit New AI/Use Indirect Food
A538.0	Experimental Use Permits – Other Actions	PRIA		18	Months	A538.0 Secondary A538 No Fee
A539	Experimental Use Permits – Other Actions	PRIA	$138,699.00	15	Months	A539 Experimental Use Permit New AI/ Use Nonfood
A539.0	Experimental Use Permits – Other Actions	PRIA		18	Months	A539.0 Secondary A539 No Fee
A550	New Product	PRIA	$19,906.00	9	Months	A550 New Product uses other than FIFRA §2(mm)
A550.1	New Product	PRIA	$4,977.00	9	Months	A550.1 Secondary A550 with Additional Fee Product Chemistry Only
A550.2	New Product	PRIA	$7,689.00	9	Months	A550.2 Secondary A550 with Additional Fee Efficacy and/or Acute Tox
A560	New Product	PRIA	$18,957.00	6	Months	A560 New manufacturing use product selective data citation
A560.1	New Product	PRIA	$4,740.00	6	Months	A560.1 Secondary A560 with Additional Fee Product Chemistry Only
A560.2	New Product	PRIA	$7,689.00	6	Months	A560.2 Secondary A560 with Additional Fee Efficacy and/or Acute Tox
A565	New Product	PRIA	$27,442.00	12	Months	A565 New manufacturing-use product unregistered source

Code	Category	Type	Fee $	Review Time	Time Type	Description
A565.1	New Product	PRIA	$6,861.00	12	Months	A565.1 Secondary A565 with Additional Fee Product Chemistry Only
A565.2	New Product	PRIA	$7,689.00	12	Months	A565.2 Secondary A565 with Additional Fee Efficacy and/or Acute Tox
A571	Amendment	PRIA	$144,058.00	18	Months	A571 Science reassessment
A572	New Product	PRIA	$19,906.00	9	Months	A572 New Product or amendment requiring Risk Assessment Branch review
A572.1	New Product	PRIA	$4,977.00	9	Months	A572.1 Secondary A572 with Additional Fee Product Chemistry Only
A572.2	New Product	PRIA	$7,689.00	9	Months	A572.2 Secondary A572 with Additional Fee Efficacy and/or Acute Tox
A575	Pre-Application	PRIA	$3,559.00	4	Months	A575 Efficacy similarity determination
B580	New Active Ingredients	PRIA	$76,832.00	22	Months	B580 New AI with Tolerance Petition
B580.0	New Active Ingredients	PRIA		22	Months	B580.0 Secondary B580 No Fee
B590	New Active Ingredients	PRIA	$48,024.00	20	Months	B590 New AI with Tolerance Exemption Petition
B590.0	New Active Ingredients	PRIA		20	Months	B590.0 Secondary B590 No Fee
B590.1	New Active Ingredients	PRIA	$12,006.00	20	Months	B590.1 Additional Submission Under B590 Action
B590X2	New Active Ingredients	PRIA	$96,048.00	20	Months	B590X2 Two New AIs with Tolerance Exemption Petitions
B590X4	New Active Ingredients	PRIA	$192,096.00	20	Months	B590X4 Four New AIs with Tolerance Exemption Petitions
B600	New Active Ingredients	PRIA	$28,816.00	15	Months	B600 New AI without Petition
B600.0	New Active Ingredients	PRIA		15	Months	B600.0 Secondary B600 No Fee
B600X10	New Active Ingredients	PRIA	$288,160.00	15	Months	B600X10 Ten New AIs without Petitions
B600X2	New Active Ingredients	PRIA	$57,632.00	15	Months	B600X2 Two New AIs without Petitions
B600X3	New Active Ingredients	PRIA	$86,448.00	15	Months	B600X3 Three New AIs without Petitions
B600X4	New Active Ingredients	PRIA	$115,264.00	15	Months	B600X4 Four New AIs without Petitions
B610	Experimental Use Permits – Other Actions	PRIA	$19,211.00	12	Months	B610 Experimental Use Permit New AI with Petition
B610.0	Experimental Use Permits – Other Actions	PRIA		12	Months	B610.0 Secondary B610 No Fee
B614	Pre-Application	PRIA	$3,809.00	3	Months	B614 Conditional Ruling on Waiver Rationale
B614X2	Pre-Application	PRIA	$7,618.00	3	Months	B614X2 Two Conditional Rulings on Waiver Rationales
B614X3	Pre-Application	PRIA	$11,427.00	3	Months	B614X3 Three Conditional Rulings on Waiver Rationales
B614X4	Pre-Application	PRIA	$15,236.00	3	Months	B614X4 Four Conditional Rulings on Waiver Rationales
B614X6	Pre-Application	PRIA	$22,854.00	3	Months	B614X6 Six Conditional Rulings on Waiver Rationales
B614X8	Pre-Application	PRIA	$30,472.00	3	Months	B614X8 Eight Conditional Rulings on Waiver Rationales
B616	Pre-Application	PRIA	$4,951.00	5	Months	B616 Conditional Ruling on Nonfood Determination

Code	Category	Type	Fee $	Review Time	Time Type	Description
B616X2	Pre-Application	PRIA	$9,902.00	5	Months	B616X2 Two Conditional Rulings on Nonfood Determinations
B617	Pre-Application	PRIA	$4,951.00	5	Months	B617 Biochemical Classification Determination
B617X2	Pre-Application	PRIA	$9,902.00	5	Months	B617X2 Two Biochemical Classification Determinations
B620	Experimental Use Permits – Other Actions	PRIA	$9,609.00	9	Months	B620 Experimental Use Permit New AI without Petition
B621	Experimental Use Permits – Other Actions	PRIA	$7,689.00	7	Months	B621 Experimental Use Permit Amendment without Petition
B622	Experimental Use Permits – Other Actions	PRIA	$19,211.00	11	Months	B622 Experimental Use Permit Amendment with Petition
B622.0	Experimental Use Permits – Other Actions	PRIA		11	Months	B622.0 Secondary B622 No Fee
B630	New Use	PRIA	$19,211.00	13	Months	B630 First Food Use with Tolerance Exemption Petition
B630.0	New Use	PRIA		13	Months	B630.0 Secondary B630 No Fee
B630X2	New Use	PRIA	$38,422.00	13	Months	B630X2 Two First Food Uses with Tolerance Exemption Petitions
B640	New Use	PRIA	$28,816.00	19	Months	B640 First Food Use with Tolerance Petition
B640.0	New Use	PRIA		19	Months	B640.0 Secondary B640 No Fee
B641	Amendment	PRIA	$19,211.00	13	Months	B641 Amendment to tolerance or exemption
B641.0	Amendment	PRIA		13	Months	B641.0 Secondary B641 No Fee
B644	New Use	PRIA	$19,211.00	8	Months	B644 New Use without Petition
B644.0	New Use	PRIA		8	Months	B644.0 Secondary B644 No Fee
B645	Experimental Use Permits – Other Actions	PRIA	$19,211.00	12	Months	B645 Experimental Use Permit New Use with Petition
B645.0	Experimental Use Permits – Other Actions	PRIA		12	Months	B645.0 Secondary B645 No Fee
B646	Experimental Use Permits – Other Actions	PRIA	$9,609.00	7	Months	B646 Experimental Use Permit New Use without Petition
B646.0	Experimental Use Permits – Other Actions	PRIA		7	Months	B646.0 Secondary B646 No Fee
B660	New Product	PRIA	$1,925.00	6	Months	B660 New Product no Data Review or Only Product Chemistry
B670	New Product	PRIA	$7,689.00	9	Months	B670 New Product Registered Source
B670.1	New Product	PRIA	$1,923.00	9	Months	B670.1 Secondary B670 with Additional Fee no Data or Product Chemistry Only
B670.2	New Product	PRIA	$7,689.00	9	Months	B670.2 Secondary B670 with Additional Fee Efficacy and/or Acute Tox
B672	New Product	PRIA	$13,723.00	15	Months	B672 New Product Unregistered Source with New Data/Waiver Review
B672.1	New Product	PRIA	$3,431.00	15	Months	B672.1 Secondary B672 with Additional Fee no Data or Product Chemistry Only
B672.2	New Product	PRIA	$7,689.00	15	Months	B672.2 Secondary B672 with Additional Fee Efficacy and/or Acute Tox

Code	Category	Type	Fee $	Review Time	Time Type	Description
B673	New Product	PRIA	$7,689.00	12	Months	B673 New Product Unregistered Source Citing Reviewed Data
B673.1	New Product	PRIA	$1,923.00	12	Months	B673.1 Secondary B673 with Additional Fee no Data or Product Chemistry Only
B674	New Product	PRIA	$1,925.00	4	Months	B674 New Product 100% Repack or Repack of an EP to MP
B677	New Product	PRIA	$13,276.00	12	Months	B677 New Animal Product
B677.1	New Product	PRIA	$3,319.00	12	Months	B677.1 Secondary B677 with Additional Fee no Data or Product Chemistry Only
B680	Amendment	PRIA	$7,689.00	5	Months	B680 Amendment with Data Submission to Product with Registered Source
B680.0	Amendment	PRIA		5	Months	B680.0 Secondary B680 No Fee
B680.1	Amendment	PRIA	$1,923.00	5	Months	B680.1 Secondary B680 with Additional Fee no Data or Product Chemistry Only
B681	Amendment	PRIA	$9,150.00	7	Months	B681 Amendment with Data Submission to Product with Unregistered Source
B681.1	Amendment	PRIA	$2,288.00	7	Months	B681.1 Secondary B681 with Additional Fee no Data or Product Chemistry Only
B682	Pre-Application	PRIA	$3,662.00	3	Months	B682 Protocol Review
B682X2	Pre-Application	PRIA	$7,324.00	3	Months	B682X2 Two Protocol Reviews
B683	Amendment	PRIA	$7,689.00	6	Months	B683 Amendment Updated Risk Assessment without Data Submission
B684	Amendment	PRIA	$13,276.00	8	Months	B684 Amendment with Submission of Animal Safety Data
B685	Amendment	PRIA	$7,689.00	5	Months	B685 Amendment to Add New Unregistered Source of AI
B685.1	Amendment	PRIA	$1,923.00	5	Months	B685.1 Secondary B685 with Additional Fee no Data
B685X2	Amendment	PRIA	$15,378.00	5	Months	B685X2 Amendment to Add Two New Unregistered Sources of AI
B685X3	Amendment	PRIA	$23,067.00	5	Months	B685X3 Amendment to Add Three New Unregistered Sources of AI
B690	New Active Ingredients	PRIA	$3,846.00	7	Months	B690 SCLP New AI
B690.0	New Active Ingredients	PRIA		7	Months	B690.0 Secondary B690 No Fee
B690X2	New Active Ingredients	PRIA	$7,692.00	7	Months	B690X2 SCLP Two New AIs
B700	Experimental Use Permits – Other Actions	PRIA	$1,925.00	7	Months	B700 SCLP Experimental Use Permit
B701	Experimental Use Permits – Other Actions	PRIA	$1,925.00	4	Months	B701 SCLP Experimental Use Permit Amendment
B710	New Product	PRIA	$1,925.00	4	Months	B710 SCLP New Product no Data or Product Chemistry Only
B720	New Product	PRIA	$1,925.00	5	Months	B720 SCLP New Product Registered Source
B721	New Product	PRIA	$4,028.00	7	Months	B721 SCLP New Product Unregistered Source
B721.1	New Product	PRIA	$1,007.00	7	Months	B721.1 Secondary B721 with Additional Fee no Data or Product Chemistry Only

Code	Category	Type	Fee $	Review Time	Time Type	Description
B721.2	New Product	PRIA	$4,028.00	7	Months	B721.2 Secondary B721 with Additional Fee Efficacy and/or Acute Tox
B722	New Use	PRIA	$3,730.00	7	Months	B722 SCLPNew Use/Amendment with Petition
B730	Amendment	PRIA	$1,925.00	5	Months	B730 SCLP Amendment with Data
B740	Experimental Use Permits – Other Actions	PRIA	$144,058.00	9	Months	B740 Experimental Use Permit for New or Existing PIP without Petition
B750	Experimental Use Permits – Other Actions	PRIA	$192,074.00	12	Months	B750 Experimental Use Permit for Existing PIP with Petition
B750.0	Experimental Use Permits – Other Actions	PRIA		12	Months	B750.0 Secondary B750 No Fee
B771	Experimental Use Permits – Other Actions	PRIA	$192,074.00	13	Months	B771 Experimental Use Permit for New PIP with Petition
B771.0	Experimental Use Permits – Other Actions	PRIA		13	Months	B771.0 Secondary B771 No Fee
B772	Experimental Use Permits – Other Actions	PRIA	$19,211.00	3	Months	B772 Amendment to Experimental Use Permit without Petition
B773	Experimental Use Permits – Other Actions	PRIA	$48,024.00	9	Months	B773 Amendment to Experimental Use Permit with Petition
B773.0	Experimental Use Permits – Other Actions	PRIA		9	Months	B773.0 Secondary B773 No Fee
B780	New Active Ingredients	PRIA	$240,090.00	16	Months	B780 New AI PIP without Petition
B800	New Active Ingredients	PRIA	$259,297.00	17	Months	B800 New AI PIP with Petition to Convert Temporary Tolerance or Exemption
B800.0	New Active Ingredients	PRIA		17	Months	B800.0 Secondary B800 No Fee
B820	New Active Ingredients	PRIA	$307,317.00	19	Months	B820 New AI PIP with Petition to Establish or Amend Tolerance or Exemption
B820.0	New Active Ingredients	PRIA		19	Months	B820.0 Secondary B820 No Fee
B820X3	New Active Ingredients	PRIA	$921,951.00	19	Months	B820X3 Three New AI PIPs with Petitions to Establish or Amend Tolerance
B851	New Active Ingredients	PRIA	$192,074.00	9	Months	B851 New Event or Exisiting PIP
B870	New Product	PRIA	$57,626.00	9	Months	B870 New PIP Use without Petition
B870.0	New Product	PRIA		9	Months	B870.0 Secondary B870 No Fee
B880	New Product	PRIA	$48,024.00	9	Months	B880 New PIP Registration without Petition
B880.0	New Product	PRIA		9	Months	B880.0 Secondary B880 No Fee
B880.1	New Product	PRIA	$12,006.00	9	Months	B880.1 Secondary B880 with Additional Fee no Data or Product Chemistry Only
B880.2	New Product	PRIA	$12,006.00	9	Months	B880.2 Secondary B880 with Additional Fee Efficacy and/or Acute Tox
B883	New Active Ingredients	PRIA	$192,074.00	13	Months	B883 New Seed Increase with Petition to Convert Temporary Tolerance
B884	New Active Ingredients	PRIA	$240,090.00	19	Months	B884 New Seed Increase with Petition to Establish Tolerance or Exemption
B884.0	New Active Ingredients	PRIA		19	Months	B884.0 Secondary B884 No Fee
B884X2	New Active Ingredients	PRIA	$480,180.00	19	Months	B884X2 Two New Seed Increase PIPs
B884X3	New Active Ingredients	PRIA	$720,270.00	19	Months	B884X3 Three New Seed Increase PIPs

Code	Category	Type	Fee $	Review Time	Time Type	Description
B885	New Product	PRIA	$48,024.00	6	Months	B885 New Seed Increase Registration
B885.1	New Product	PRIA	$12,006.00	6	Months	B885.1 Secondary B885 with Additional Fee no Data or Product Chemistry Only
B885.2	New Product	PRIA	$12,006.00	6	Months	B885.2 Secondary B885 with Additional Fee Efficacy and/or Acute Tox
B890	Amendment	PRIA	$96,039.00	9	Months	B890 Amendment to Seed Increase Registration
B900	Amendment	PRIA	$19,211.00	6	Months	B900 Amendment to PIP Registration
B900.0	Amendment	PRIA		6	Months	B900.0 Secondary B900 No Fee
B902	Pre-Application	PRIA	$9,609.00	3	Months	B902 PIP Protocol review
B903	New Use	PRIA	$96,039.00	12	Months	B903 Petition to Establish a Tolerance or Exemption for a PIP Inert
B903.0	New Use	PRIA		12	Months	B903.0 Secondary B903 No Fee
B903X3	New Use	PRIA	$288,117.00	6	Months	B903X3 Three Petitions to Establish a Tolerance or Exemption for a PIP Inert
B904	New Use	PRIA	$192,074.00	12	Months	B904 Petition to Establish Import Tolerance or Exemption for PIP Ingredients
B905	Pre-Application	PRIA	$96,039.00	6	Months	B905 FIFRA Scientific Advisory Panel Review
B906	New Use	PRIA	$48,020.00	9	Months	B906 Petition to Establish Emerging Tech Ingredient Tolerance or Exemption
B907	New Use	PRIA	$19,211.00	9	Months	B907 Petition to Convert Emerging Tech Temporary Tolerance or Exemption
B909	Pre-Application	PRIA	$19,211.00	6	Months	B909 PIP Tolerance Exemption Determination
B910	Pre-Application	PRIA	$9,609.00	3	Months	B910 Biotech Notification
B921	Experimental Use Permits – Other Actions	PRIA	$192,074.00	12	Months	B921 Experimental Use Permit Based on GM in Animals without Petition
B922	New Active Ingredients	PRIA	$240,090.00	16	Months	B922 New AI Based on GM in Animals without Petition
B923	Experimental Use Permits – Other Actions	PRIA	$240,091.00	15	Months	B923 Experimental Use Permit Based on GM in Animals with Petition
B924	New Active Ingredients	PRIA	$307,317.00	19	Months	B924 New AI Based on GM in Animals with Petition
B924.0	New Active Ingredients	PRIA		19	Months	B924.0 Secondary B924 No Fee
B925	Experimental Use Permits – Other Actions	PRIA	$28,825.00	11	Months	B925 Experimental Use Permit Based on RNAi Nonfood
B926	New Active Ingredients	PRIA	$86,446.00	17	Months	B926 New AI Based on RNAi without Petition
B927	Experimental Use Permits – Other Actions	PRIA	$57,634.00	14	Months	B927 Experimental Use Permit Based on RNAi with Petition
B928	New Active Ingredients	PRIA	$144,071.00	22	Months	B928 New AI Based on RNAi with Petition
B928.0	New Active Ingredients	PRIA		22	Months	B928.0 Secondary B928 No Fee
B929	New Product	PRIA	$7,689.00	10	Months	B929 New Product Based on RNAi
B929.1	New Product	PRIA	$1,923.00	10	Months	B929.1 Secondary B929 with Additional Fee no Data or Product Chemistry Only

Code	Category	Type	Fee $	Review Time	Time Type	Description
B929.2	New Product	PRIA	$7,689.00	10	Months	B929.2 Secondary B929 with Additional Fee Efficacy and/or Acute Tox
B930	Experimental Use Permits – Other Actions	PRIA	$19,211.00	3	Months	B930 Amendment to Non-PIP Experimental Use Permit without Petititon
B931	Experimental Use Permits – Other Actions	PRIA	$48,024.00	9	Months	B931 Amendment to Non-PIP Experimental Use Permit with Petititon
B932	Amendment	PRIA	$19,211.00	6	Months	B932 Amendment to Non-PIP Registration
B932.0	Amendment	PRIA		6	Months	B932.0 Secondary B932 No Fee
I001	Inert	PRIA	$40,633.00	13	Months	I001 New Approval for Food Use Inert
I001X2	Inert	PRIA	$81,266.00	12	Months	I001X2 Two New Approvals for Food Use Inerts
I002	Amendment	PRIA	$11,288.00	11	Months	I002 Amendment to Inert Tolerance or Exemption with New Data
I003	Amendment	PRIA	$4,980.00	11	Months	I003 Amendment to Inert Tolerance or Exemption with No New Data
I004	Inert	PRIA	$16,594.00	6	Months	I004 New Approval for Nonfood Inert
I005	Amendment	PRIA	$8,299.00	6	Months	I005 New Use for NonFood Inert with New Data
I006	Amendment	PRIA	$4,980.00	3	Months	I006 New Use for NonFood Inert with No New Data
I007	Inert	PRIA	$2,490.00	5	Months	I007 New Approval for Substantially Similar Nonfood Inert
I008	Inert	PRIA	$5,643.00	7	Months	I008 New Use or New Approval for Food Use Polymer
I009	Inert	PRIA	$4,649.00	4	Months	I009 New Use or New Approval for Nonfood Use Polymer
I010	New Use	PRIA	$2,490.00	7	Months	I010 Adding Closely Related CASRNs to Inert Tolerance Exemption
I011	Inert	PRIA	$899,463.00	26	Months	I011 New Approval for Food Use Safener
I012	Inert	PRIA	$624,905.00	21	Months	I012 New Approval for Nonfood Safener
I013	New Use	PRIA	$94,773.00	17	Months	I013 New Food Use for Approved Safener
I014	Inert	PRIA	$37,878.00	15	Months	I014 New Nonfood Use for Approved Safener
I015	Inert	PRIA	$405,919.00	26	Months	I015 New Generic Data for Approved Safener
I016	Inert	PRIA	$83,940.00	15	Months	I016 Amendment to Tolerance for Approved Safener
I017	Amendment	PRIA	$19,906.00	8	Months	I017 New Source of Approved Safener
I018	Inert	PRIA	$2,490.00	3	Months	I018 Petition to Add Approved Inert to Commodity Inert List
M001	New Use	PRIA	$11,947.00	14	Months	M001 HSRB Protocol Review
M002	Pre-Application	PRIA	$11,947.00	14	Months	M002 HSRB Study Review
M003	Pre-Application	PRIA	$96,234.00	12	Months	M003 External Technical Review for PRIA Less Than 12 Months
M004	Pre-Application	PRIA	$96,234.00	18	Months	M004 External Technical Review for PRIA Greater Than or Equal to 12 Months

Code	Category	Type	Fee $	Review Time	Time Type	Description
M005	New Product	PRIA	$33,185.00	9	Months	M005 New Combination Product Multiple Divisions
M006	MISCELLANEOUS	PRIA	$418.00	1	Months	M006 Gold Seal Letter
M007	MISCELLANEOUS	PRIA	$8,299.00	12	Months	M007 Request to Extend Exclusive Use Period
M008	MISCELLANEOUS	PRIA	$2,490.00	15	Months	M008 Request to Grant Exclusive Use Rights
M009	MISCELLANEOUS	PRIA	$3,559.00	6	Months	M009 Non-FIFRA Regulated Determination
M010	Pre-Application	PRIA	$3,559.00	4	Months	M010 Conditional Ruling on Substantial Similarity
M011	Amendment	PRIA	$5,492.00	4	Months	M011 Label Amendment to Add the DfE Logo
M012	MISCELLANEOUS	PRIA	$418.00	1	Months	M012 Certification of Establishment Letter
M013	MISCELLANEOUS	PRIA	$298,352.00	18	Months	M013 Cancer Reassessment
M014	Pre-Application	PRIA	$18,296.00	8	Months	M014 Nanoparticle Determination
R010	New Active Ingredients	PRIA	$1,133,324.00	36	Months	R010 New AI Food Use
R010.0	New Active Ingredients	PRIA		36	Months	R010.0 Secondary R010 No Fee
R010.1	New Active Ingredients	PRIA	$283,331.00	36	Months	R010.1 Secondary R010 with Additional Fee
R020	New Active Ingredients	PRIA	$944,438.00	27	Months	R020 New AI Food Use Reduced Risk
R020.0	New Active Ingredients	PRIA		27	Months	R020.0 Secondary R020 No Fee
R040	Experimental Use Permits – Other Actions	PRIA	$696,028.00	18	Months	R040 New AI Food Use Experimental Use Permit
R040.0	Experimental Use Permits – Other Actions	PRIA		18	Months	R040.0 Secondary R040 No Fee
R060	New Active Ingredients	PRIA	$787,381.00	30	Months	R060 New AI Nonfood Outdoor
R060.0	New Active Ingredients	PRIA		30	Months	R060.0 Secondary R060 No Fee
R060.1	New Active Ingredients	PRIA	$196,846.00	30	Months	R060.1 Secondary R060 with Additional Fee
R070	New Active Ingredients	PRIA	$656,151.00	24	Months	R070 New AI Nonfood Outdoor Reduced Risk
R070.0	New Active Ingredients	PRIA		24	Months	R070.0 Secondary R070 No Fee
R090	Experimental Use Permits – Other Actions	PRIA	$487,127.00	16	Months	R090 New AI Nonfood Outdoor Experimental Use Permit
R110	New Active Ingredients	PRIA	$437,923.00	20	Months	R110 New AI Nonfood Indoor
R110.0	New Active Ingredients	PRIA		20	Months	R110.0 Secondary R110 No Fee
R120	New Active Ingredients	PRIA	$364,934.00	14	Months	R120 New AI Nonfood Indoor Reduced Risk
R121	Experimental Use Permits – Other Actions	PRIA	$274,389.00	18	Months	R121 New AI Nonfood Indoor Experimental Use Permit
R122	New Product	PRIA	$477,253.00	27	Months	R122 New AI Enriched Isomer or Approved AI
R122.0	New Product	PRIA		27	Months	R122.0 Secondary R122 No Fee
R122.1	New Product	PRIA	$119,314.00	27	Months	R122.1 Secondary R122 with Additional Fee

Code	Category	Type	Fee $	Review Time	Time Type	Description
R123	New Active Ingredients	PRIA	$710,111.00	27	Months	R123 New AI Seed Treatment Only no Residues Expected
R123.0	New Active Ingredients	PRIA		27	Months	R123.0 Secondary R123 No Fee
R124	Pre-Application	PRIA	$3,809.00	6	Months	R124 Conditional Ruling on Study Waiver
R124X2	Pre-Application	PRIA	$7,618.00	6	Months	R124X2 Two Conditional Rulings on Study Waiver
R124X3	Pre-Application	PRIA	$11,427.00	6	Months	R124X3 Three Conditional Rulings on Study Waiver
R124X4	Pre-Application	PRIA	$15,236.00	6	Months	R124X4 Four Conditional Rulings on Study Waiver
R124X6	Pre-Application	PRIA	$22,854.00	6	Months	R124X6 Six Conditional Rulings on Study Waiver
R125	Experimental Use Permits – Other Actions	PRIA	$487,127.00	16	Months	R125 New AI Seed Treatment Only Experimental Use Permit
R126	New Active Ingredients	PRIA	$781,122.00	31	Months	R126 New AI Seed Treatment Only with Petition
R130	New Use	PRIA	$288,108.00	23	Months	R130 First Food Use Indoor Only
R130.0	New Use	PRIA		23	Months	R130.0 Secondary R130 No Fee
R140	New Use	PRIA	$67,230.00	17	Months	R140 Additional Food Use Indoor Only
R140.0	New Use	PRIA		17	Months	R140.0 Secondary R140 No Fee
R140X5	New Use	PRIA	$336,150.00	17	Months	R140X5 Five Additional Food Uses Indoor Only
R150	New Use	PRIA	$477,215.00	23	Months	R150 First Food Use
R150.0	New Use	PRIA		23	Months	R150.0 Secondary R150 No Fee
R155	Experimental Use Permits – Other Actions	PRIA	$397,680.00	21	Months	R155 First Food Use Experimental Use Permit
R160	New Use	PRIA	$397,680.00	18	Months	R160 First Food Use Reduced Risk
R160.0	New Use	PRIA		18	Months	R160.0 Secondary R160 No Fee
R170	New Use	PRIA	$119,415.00	17	Months	R170 Additional Food Use
R170.0	New Use	PRIA		17	Months	R170.0 Secondary R170 No Fee
R170X2	New Use	PRIA	$238,830.00	17	Months	R170X2 Two Additional Food Uses
R170X3	New Use	PRIA	$358,245.00	17	Months	R170X3 Three Additional Food Uses
R170X4	New Use	PRIA	$477,660.00	17	Months	R170X4 Four Additional Food Uses
R170X5	New Use	PRIA	$597,075.00	17	Months	R170X5 Five Additional Food Uses
R175	New Use	PRIA	$99,513.00	14	Months	R175 Additional Food Uses Covered by Crop Group Conversion
R175.0	New Use	PRIA		14	Months	R175.0 Secondary R175 No Fee
R180	New Use	PRIA	$99,513.00	12	Months	R180 Additional Food Use Reduced Risk
R180.0	New Use	PRIA		12	Months	R180.0 Secondary R180 No Fee
R180X2	New Use	PRIA	$199,026.00	12	Months	R180X2 Two Additional Food Uses Reduced Risk
R180X3	New Use	PRIA	$298,539.00	12	Months	R180X3 Three Additional Food Uses Reduced Risk
R180X4	New Use	PRIA	$398,052.00	12	Months	R180X4 Four Additional Food Uses Reduced Risk

Code	Category	Type	Fee $	Review Time	Time Type	Description
R180X5	New Use	PRIA	$497,565.00	12	Months	R180X5 Five Additional Food Uses Reduced Risk
R190	New Use	PRIA	$716,475.00	17	Months	R190 Six or More Additional Food Uses
R190.0	New Use	PRIA		17	Months	R190.0 Secondary R190 No Fee
R200	New Use	PRIA	$597,064.00	12	Months	R200 Six or More Additional Food Uses Reduced Risk
R200.0	New Use	PRIA		12	Months	R200.0 Secondary R200 No Fee
R210	Experimental Use Permits – Other Actions	PRIA	$73,721.00	12	Months	R210 Additional Food Use Experimental Use Permit
R210.0	Experimental Use Permits – Other Actions	PRIA		12	Months	R210.0 Secondary R210 No Fee
R220	Experimental Use Permits – Other Actions	PRIA	$29,856.00	6	Months	R220 Additional Food Use Experimental Use Permit Crop Destruct
R230	New Use	PRIA	$47,726.00	16	Months	R230 Additional Nonfood Use Outdoor
R230.0	New Use	PRIA		16	Months	R230.0 Secondary R230 No Fee
R230X2	New Use	PRIA	$95,452.00	16	Months	R230X2 Two Additional Nonfood Uses Outdoor
R230X3	New Use	PRIA	$143,178.00	16	Months	R230X3 Three Additional Nonfood Uses Outdoor
R240	New Use	PRIA	$39,772.00	10	Months	R240 Additional Nonfood Use Outdoor Reduced Risk
R240.0	New Use	PRIA		10	Months	R240.0 Secondary R240 No Fee
R240X2	New Use	PRIA	$79,544.00	10	Months	R240X2 Two Additional Nonfood Uses Outdoor Reduced Risk
R240X3	New Use	PRIA	$119,316.00	10	Months	R240X3 Three Additional Nonfood Uses Outdoor Reduced Risk
R250	Experimental Use Permits – Other Actions	PRIA	$29,856.00	6	Months	R250 Additional Nonfood Use Outdoor Experimental Use Permit
R251	Experimental Use Permits – Other Actions	PRIA	$29,856.00	8	Months	R251 Additional Use Experimental Use Permit Crop Destruct
R260	New Use	PRIA	$23,052.00	12	Months	R260 Additional Nonfood Use Indoor
R260.0	New Use	PRIA		12	Months	R260.0 Secondary R260 No Fee
R260X2	New Use	PRIA	$46,104.00	12	Months	R260X2 Two Additional Nonfood Uses Indoor
R260X3	New Use	PRIA	$69,156.00	12	Months	R260X3 Three Additional Nonfood Uses Indoor
R270	New Use	PRIA	$19,211.00	9	Months	R270 Additional Nonfood Use Indoor Reduced Risk
R271	Experimental Use Permits – Other Actions	PRIA	$14,637.00	6	Months	R271 Additional Nonfood Use Indoor Experimental Use Permit
R272	Pre-Application	PRIA	$3,809.00	3	Months	R272 Review of Protocol
R272.0	Pre-Application	PRIA		3	Months	R272.0 Secondary R272 No Fee
R272X2	Pre-Application	PRIA	$7,618.00	3	Months	R272X2 Two Reviews of Protocol
R272X3	Pre-Application	PRIA	$11,427.00	3	Months	R272X3 Three Reviews of Protocol
R272X4	Pre-Application	PRIA	$15,236.00	3	Months	R272X4 Four Reviews of Protocol
R272X5	Pre-Application	PRIA	$19,045.00	3	Months	R272X5 Five Reviews of Protocol
R272X6	Pre-Application	PRIA	$22,854.00	3	Months	R272X6 Six Reviews of Protocol

Code	Category	Type	Fee $	Review Time	Time Type	Description
R273	New Use	PRIA	$75,918.00	12	Months	R273 Additional Use Seed Treatment Only
R273.0	New Use	PRIA		12	Months	R273.0 Secondary R273 No Fee
R273X2	New Use	PRIA	$151,836.00	12	Months	R273X2 Two Additional Uses Seed Treatment Only
R273X3	New Use	PRIA	$227,754.00	12	Months	R273X3 Three Additional Uses Seed Treatment Only
R274	New Use	PRIA	$455,483.00	12	Months	R274 Six or More Additional Uses Seed Treatment Only
R274.0	New Use	PRIA		12	Months	R274.0 Secondary R274 No Fee
R275	Pre-Application	PRIA	$3,809.00	3	Months	R275 Rebuttal of Agency Review of Protocol
R276	New Use	PRIA	$83,538.00	14	Months	R276 Additional Use Seed Treatment Only with Petition
R276.0	New Use	PRIA		14	Months	R276.0 Secondary R276 No Fee
R276.1	New Use	PRIA		14	Months	R276.1 Secondary R276 with Additional Fee
R276x2	New Use	PRIA	$167,076.00	14	Months	R276x2 Two Additional Uses Seed Treatment Only with Petition
R277	New Use	PRIA	$501,228.00	14	Months	R277 Six or More Additional Uses Seed Treatment Only with Petition
R278	Pre-Application	PRIA	$5,174.00	5	Months	R278 Review of Companion Animal Safety Protocol
R279	Pre-Application	PRIA	$5,460.00	3	Months	R279 Preapplication Determination of Reduced Risk
R280	New Use	PRIA	$480,177.00	22	Months	R280 Import Tolerance New AI or First Food Use
R280.0	New Use	PRIA		22	Months	R280.0 Secondary R280 No Fee
R281	New Use	PRIA	$72,029.00	12	Months	R281 Import Tolerance National Authority Residue Data Review
R281.0	New Use	PRIA		12	Months	R281.0 Secondary R281 No Fee
R281X2	New Use	PRIA	$144,058.00	12	Months	R281X2 Two Import Tolerance National Authority Residue Data Review
R281X3	New Use	PRIA	$216,087.00	12	Months	R281X3 Three Import Tolerance National Authority Residue Data Review
R281X4	New Use	PRIA	$288,116.00	12	Months	R281X4 Four Import Tolerance National Authority Residue Data Review
R281X5	New Use	PRIA	$360,145.00	12	Months	R281X5 Five Import Tolerance National Authority Residue Data Review
R282	New Use	PRIA	$432,159.00	12	Months	R282 Six or More Import Tolerance National Authority Residue Data Review
R290	New Use	PRIA	$96,039.00	16	Months	R290 Import Tolerance Additional Food Use
R290.0	New Use	PRIA		16	Months	R290.0 Secondary R290 No Fee
R290X2	New Use	PRIA	$192,078.00	16	Months	R290X2 Two Import Tolerance Additional Food Uses
R290X3	New Use	PRIA	$288,117.00	16	Months	R290X3 Three Import Tolerance Additional Food Uses

Code	Category	Type	Fee $	Review Time	Time Type	Description
R290X4	New Use	PRIA	$384,156.00	16	Months	R290X4 Four Import Tolerance Additional Food Uses
R291	New Use	PRIA	$576,212.00	16	Months	R291 Six or More Import Tolerance Additional Food Uses
R292	Amendment	PRIA	$68,237.00	12	Months	R292 Amendment to Harmonize Tolerance
R292.0	Amendment	PRIA		12	Months	R292.0 Secondary R292 No Fee
R292X2	Amendment	PRIA	$136,474.00	12	Months	R292X2 Two Amendments to Harmonize Tolerance
R292X3	Amendment	PRIA	$204,711.00	12	Months	R292X3 Three Amendments to Harmonize Tolerance
R292X4	Amendment	PRIA	$272,948.00	12	Months	R292X4 Four Amendments to Harmonize Tolerance
R293	New Use	PRIA	$80,489.00	13	Months	R293 Tolerance for Inadvertant Residues
R293.0	New Use	PRIA		13	Months	R293.0 Secondary R293 No Fee
R294	New Use	PRIA	$482,919.00	13	Months	R294 Six or More Tolerance for Inadvertant Residues
R295	New Use	PRIA	$99,513.00	16	Months	R295 Tolerance for Rotational Crop Application
R295.0	New Use	PRIA		16	Months	R295.0 Secondary R295 No Fee
R295X2	New Use	PRIA	$199,026.00	16	Months	R295X2 Two Tolerances for Rotational Crop Application
R295X3	New Use	PRIA	$298,539.00	16	Months	R295X3 Three Tolerances for Rotational Crop Application
R295X4	New Use	PRIA	$398,052.00	16	Months	R295X4 Four Tolerances for Rotational Crop Application
R296	New Use	PRIA	$597,064.00	16	Months	R296 Six or More Tolerances for Rotational Crop Application
R296.0		PRIA		16	Months	R296.0 Secondary R296 No Fee
R297	Amendment	PRIA	$409,392.00	12	Months	R297 Six or More Amended Tolerances
R298	Amendment	PRIA	$88,137.00	14	Months	R298 Amendment to a Tolerance
R298.0	Amendment	PRIA		14	Months	R298.0 Secondary R298 No Fee
R298X2	Amendment	PRIA	$176,274.00	14	Months	R298X2 Two Amendments to a Tolerance
R298X3	Amendment	PRIA	$264,411.00	14	Months	R298X3 Three Amendments to a Tolerance
R298X4	Amendment	PRIA	$352,548.00	14	Months	R298X4 Four Amendments to a Tolerance
R299	Amendment	PRIA	$429,296.00	14	Months	R299 Six or More Amendments to a Tolerance
R299.0	Amendment	PRIA		14	Months	R299.0 Secondary R299 No Fee
R300	New Product	PRIA	$2,384.00	4	Months	R300 New Product No Data or Product Chemistry Selective Data Owned or LOA
R300X2	New Product	PRIA	$4,768.00	4	Months	R300X2 2 New Products No Data or Product Chemistry Selective Data Owned or LOA
R300X3	New Product	PRIA	$7,152.00	4	Months	R300X3 3 New Products No Data or Product Chemistry Selective Data Owned or LOA

Code	Category	Type	Fee $	Review Time	Time Type	Description
R300X4	New Product	PRIA	$9,536.00	4	Months	R300X4 4 New Product No Data or Product Chemistry Selective Data Owned or LOA
R301	New Product	PRIA	$2,856.00	4	Months	R301 New Product No Data or Product Chemistry Selective Data Not Owned No LOA
R310	New Product	PRIA	$10,990.00	7	Months	R310 New Product with Registered Source Up to 3 Pests
R310.0	New Product	PRIA		7	Months	R310.0 Secondary R310 No Fee
R310.1	New Product	PRIA	$2,748.00	7	Months	R310.1 Secondary R310 with Additional Fee no Data or Product Chemistry Only
R310.2	New Product	PRIA	$10,990.00	7	Months	R310.2 Secondary R310 with Additional Fee Efficacy and/or Acute Tox
R314	New Product	PRIA	$12,983.00	8	Months	R314 New Combination Product Up to 3 AIs with Registered Source Up to 3 Pests
R314.1	New Product	PRIA	$3,246.00	8	Months	R314.1 Secondary R314 with Additional Fee no Data or Product Chemistry Only
R314.2	New Product	PRIA	$10,990.00	8	Months	R314.2 Secondary R314 with Additional Fee Efficacy and/or Acute Tox
R315	New Product	PRIA	$14,779.00	9	Months	R315 New On-Animal Product
R315.1	New Product	PRIA	$3,695.00	9	Months	R315.1 Secondary R315 with Additional Fee no Data or Product Chemistry Only
R315.2	New Product	PRIA	$10,990.00	9	Months	R315.2 Secondary R315 with Additional Fee Efficacy and/or Acute Tox
R316	New Product	PRIA	$17,009.00	9	Months	R316 New Product with Registered Source with 4 to 7 Pests
R316.1	New Product	PRIA	$4,253.00	8	Months	R316.1 Secondary R316 with Additional Fee no Data or Product Chemistry Only
R316.2	New Product	PRIA	$10,990.00	9	Months	R316.2 Secondary R316 with Additional Fee Efficacy and/or Acute Tox
R317	New Product	PRIA	$23,029.00	10	Months	R317 New Product with Registered Source More Than 7 Pests
R317.1	New Product	PRIA	$5,758.00	10	Months	R317.1 Secondary R317 with Additional Fee no Data or Product Chemistry Only
R317.2	New Product	PRIA	$10,990.00	10	Months	R317.2 Secondary R317 with Additional Fee Efficacy and/or Acute Tox
R318	New Product	PRIA	$19,944.00	9	Months	R318 New Combination Product 4 or More AIs with Registered Source Up to 3 Pests
R318.1	New Product	PRIA	$4,986.00	9	Months	R318.1 Secondary R318 with Additional Fee no Data or Product Chemistry Only
R318.2	New Product	PRIA	$10,990.00	9	Months	R318.2 Secondary R318 with Additional Fee Efficacy and/or Acute Tox
R319	New Product	PRIA	$19,002.00	10	Months	R319 New Combination Product Up to 3 AIs with Registered Source 4 to 7 Pests
R319.1	New Product	PRIA	$4,751.00	10	Months	R319.1 Secondary R319 with Additional Fee no Data or Product Chemistry Only
R319.2	New Product	PRIA	$10,990.00	10	Months	R319.2 Secondary R319 with Additional Fee Efficacy and/or Acute Tox

Code	Category	Type	Fee $	Review Time	Time Type	Description
R320	New Product	PRIA	$19,906.00	12	Months	R320 New Product with New Physical Form Review in Science Divisions
R320.0	New Product	PRIA		12	Months	R320.0 Secondary R320 No Fee
R320.1	New Product	PRIA	$4,977.00	12	Months	R320.1 Secondary R320 with Additional Fee no Data or Product Chemistry Only
R320.2	New Product	PRIA	$10,990.00	12	Months	R320.2 Secondary R320 with Additional Fee Efficacy and/or Acute Tox
R321	New Product	PRIA	$25,964.00	11	Months	R321 New Combination Product 4 or More AIs with Registered Source 4 to 7 Pests
R321.1	New Product	PRIA	$6,491.00	11	Months	R321.1 Secondary R321 with Additional Fee no Data or Product Chemistry Only
R321.2	New Product	PRIA	$10,990.00	11	Months	R321.2 Secondary R321 with Additional Fee Efficacy and/or Acute Tox
R331	New Product	PRIA	$3,809.00	3	Months	R331 New Product Repack of EP as MP
R332	New Product	PRIA	$426,215.00	24	Months	R332 New Product Manufacturing Use with New Generic Data Package
R332.1	New Product	PRIA	$106,554.00	24	Months	R332.1 Secondary R332 with Additional Fee no Data or Product Chemistry Only
R332.2	New Product	PRIA	$106,554.00	24	Months	R332.2 Secondary R332 with Additional Fee Efficacy and/or Acute Tox
R333	New Product	PRIA	$29,856.00	11	Months	R333 New Product Unregistered Source Review in Science Divisions Data Owned
R333.1	New Product	PRIA	$7,464.00	11	Months	R333.1 Secondary R333 with Additional Fee no Data or Product Chemistry Only
R333.2	New Product	PRIA	$10,990.00	11	Months	R333.2 Secondary R333 with Additional Fee Efficacy and/or Acute Tox
R333X2	New Product	PRIA	$49,762.00	11	Months	R333X2 R333 with Two Unregistered Sources
R333X3	New Product	PRIA	$69,668.00	11	Months	R333X3 R333 with Three Unregistered Sources
R333X4	New Product	PRIA	$89,574.00	11	Months	R333X4 R333 with Four Unregistered Sources
R334	New Product	PRIA	$34,764.00	12	Months	R334 New Product Unregistered Source Review in Science Divisions Data Not Owned
R334.1	New Product	PRIA	$8,691.00	12	Months	R334.1 Secondary R334 with Additional Fee no Data or Product Chemistry Only
R334.2	New Product	PRIA	$10,990.00	12	Months	R334.2 Secondary R334 with Additional Fee Efficacy and/or Acute Tox
R334X2	New Product	PRIA	$54,670.00	12	Months	R334X2 R333 with Two Unregistered Sources
R334X3	New Product	PRIA	$74,576.00	12	Months	R334X3 R334 with Three Unregistered Sources
R334X4	New Product	PRIA	$94,482.00	12	Months	R334X4 R334 with Four Unregistered Sources
R334X5	New Product	PRIA	$114,388.00	12	Months	R334X5 R334 with Five Unregistered Sources
R340	Amendment	PRIA	$7,508.00	4	Months	R340 Amendment with RD Review Up to 2 Pests

Code	Category	Type	Fee $	Review Time	Time Type	Description
R340.1	Amendment	PRIA	$1,877.00	4	Months	R340.1 Secondary R340 with Additional Fee
R341	Amendment	PRIA	$9,014.00	6	Months	R341 Amendment with RD Review More Than 2 Pests
R341.1	Amendment	PRIA	$2,254.00	4	Months	R341.1 Secondary R341 with Additional Fee
R345	Amendment	PRIA	$13,276.00	7	Months	R345 Amendment to On-Animal Product
R350	Amendment	PRIA	$19,906.00	9	Months	R350 Amendment with Science Division Review
R350.0	Amendment	PRIA		9	Months	R350.0 Secondary R350 No Fee
R350.1	Amendment	PRIA	$4,977.00	9	Months	R350.1 Secondary R350 with Additional Fee
R351	Amendment	PRIA	$19,906.00	8	Months	R351 Amendment to Add Unregistered Source
R351.1	Amendment	PRIA	$4,977.00	8	Months	R351.1 Secondary R351 with Additional Fee
R351X2	Amendment	PRIA	$39,812.00	8	Months	R351X2 Amendment to Add 2 Unregistered Sources
R351X3	Amendment	PRIA	$59,718.00	8	Months	R351X3 Amendment to Add 3 Unregistered Sources
R351X4	Amendment	PRIA	$79,624.00	8	Months	R351X4 Amendment to Add 4 Unregistered Sources
R351X5	Amendment	PRIA	$99,530.00	8	Months	R351X5 Amendment to Add 5 Unregistered Sources
R352	Amendment	PRIA	$19,906.00	8	Months	R352 Amendment to Add Approved Uses With Selective Data Citation
R352.1	Amendment	PRIA	$4,977.00	8	Months	R352.1 Secondary R352 with Additional Fee
R361	New Product	PRIA	$24,570.00	12	Months	R361 New Combination Product Up to 3 AIs with Registered Source >7 Pests
R361.1	New Product	PRIA	$6,143.00	12	Months	R361.1 Secondary R361 with Additional Fee no Data or Product Chemistry Only
R361.2	New Product	PRIA	$10,990.00	12	Months	R361.2 Secondary R361 with Additional Fee Efficacy and/or Acute Tox
R362	New Product	PRIA	$26,618.00	13	Months	R362 New Combination Product 4 or More AIs with Registered Source >7 Pests
R362.1	New Product	PRIA	$6,655.00	13	Months	R362.1 Secondary R362 with Additional Fee no Data or Product Chemistry Only
R362.2	New Product	PRIA	$10,990.00	13	Months	R362.2 Secondary R362 with Additional Fee Efficacy and/or Acute Tox
R363	New Product	PRIA	$8,190.00	6	Months	R363 New Product Repack MP to EP
R371	Experimental Use Permits – Other Actions	PRIA	$15,187.00	6	Months	R371 Amendment to Experimental Use Permit
FR05		Fee Waiver		60	Days	FR05 5% Discretionary Refund Request
FR10		Fee Waiver		60	Days	FR10 10% Discretionary Refund Request
FR15		Fee Waiver		60	Days	FR15 15% Discretionary Refund Request
FR20		Fee Waiver		60	Days	FR20 20% Discretionary Refund Request
FR25		Fee Waiver		60	Days	FR25 25% Discretionary Refund Request

Code	Category	Type	Fee $	Review Time	Time Type	Description
FR30		Fee Waiver		60	Days	FR30 30% Discretionary Refund Request
FR35		Fee Waiver		60	Days	FR35 35% Discretionary Refund Request
FR40		Fee Waiver		60	Days	FR40 40% Discretionary Refund Request
FR45		Fee Waiver		60	Days	FR45 45% Discretionary Refund Request
FR50		Fee Waiver		60	Days	FR50 50% Discretionary Refund Request
FR55		Fee Waiver		60	Days	FR55 55% Discretionary Refund Request
FR60		Fee Waiver		60	Days	FR60 60% Discretionary Refund Request
FR65		Fee Waiver		60	Days	FR65 65% Discretionary Refund Request
FR70		Fee Waiver		60	Days	FR70 70% Discretionary Refund Request
FR75		Fee Waiver		60	Days	FR75 75% Discretionary Refund Request
W02		Fee Waiver		60	Days	W02 50% Small Business Fee Waiver Request
W04		Fee Waiver		60	Days	W04 100% IR-4 Fee Waiver Request
W05		Fee Waiver		60	Days	W05 100% Minor Use Fee Waiver Request
W06		Fee Waiver		60	Days	W06 100% State and Federal Agency Fee Waiver Request
W075		Fee Waiver		60	Days	W075 75% Small Business Fee Waiver Request