

Screen mock-ups are approximations and may change. Each optional field is outlined by a red text box.

Chemical Information Submission System

The ‘Chemical Information Submission System’ screen provides a list of submission types from which the user can choose. Based on the selected submission type, the system directs the user to the appropriate application ‘Home’ screen.

The following exhibit shows the screen capture for the ‘Chemical Information Submission System’ screen:

Exhibit 1-1 Chemical Information Submission System – 8(d)

CDX CENTRAL DATA EXCHANGE

Logged in as: John Doe, Primary Authorized Official [Log Out](#)

Chemical Information Submission System

Please Choose a Submission Type

8(d) Health and Safety Data Reporting Form

Under TSCA Section 8(d), EPA has the authority to promulgate rules to require producers, importers, and processors to submit lists and/or copies of ongoing and completed, unpublished health and safety studies. EPA's TSCA Section 8(d) "Health & Safety Data Reporting Rule" was developed to gather health and safety information on chemical substances and mixtures needed by EPA to carry out its TSCA mandates (e.g., to support OPPT's Existing Chemicals Program and Chemical Testing Program and to set priorities for TSCA risk assessment/management activities). EPA has also used its TSCA Section 8(d) authority to gather information needed by other EPA Program Offices and other Federal Agencies. Chemicals that are designated or recommended for testing by the TSCA Interagency Testing Committee (ITC) may be added to the rule via immediate final rulemaking (up to 50 substances/year). Non-ITC chemicals can be added to the Section 8(d) rule via notice and comment rulemaking.

The software includes embedded help files and a downloadable user manual to guide you through the 8(d) Health & Safety Data Reporting submission process. Submit information for all reportable chemical substances at your site in one form. Note that a separate submission is required for each reporting site.

OK

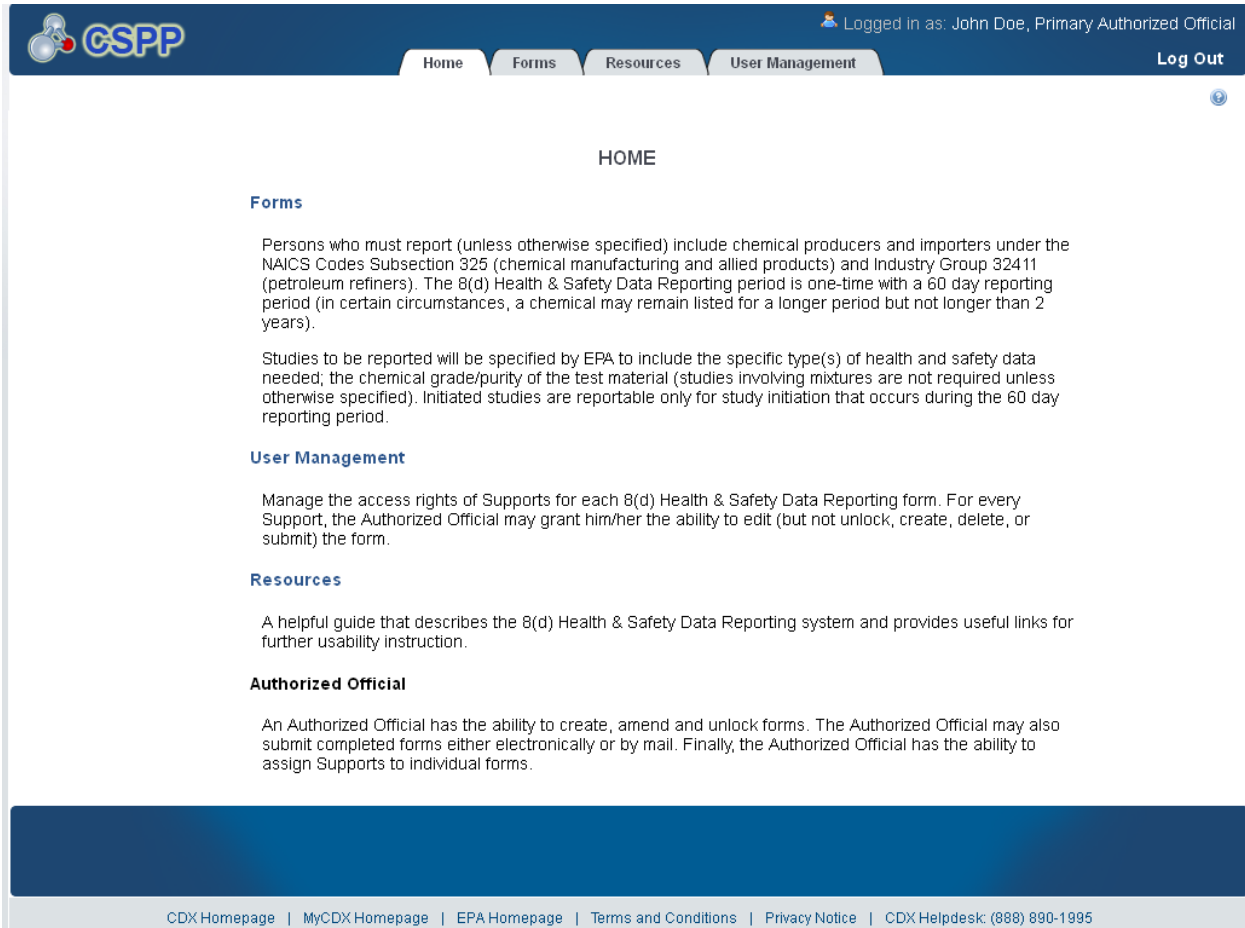
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Home

The 'Home' screen provides the user with links and tabs to access information for the form type, user management, and resources. Additionally, the system provides the user with information relevant to the logged in user's role.

The following exhibit shows the screen capture for the 'Home' screen:

Exhibit 1-2 Home Screen



The screenshot displays the GSPP (Global Support and Reporting Platform) Home Screen. At the top, there is a dark blue header with the GSPP logo on the left and the text "Logged in as: John Doe, Primary Authorized Official" on the right. Below the header is a navigation bar with tabs for "Home", "Forms", "Resources", and "User Management", with "Home" currently selected. A "Log Out" button is located in the top right corner. The main content area is titled "HOME" and contains four sections: "Forms", "User Management", "Resources", and "Authorized Official". Each section provides a brief description of its function. At the bottom of the page, there is a dark blue footer bar with a light gray bar below it containing links: "CDX Homepage", "MyCDX Homepage", "EPA Homepage", "Terms and Conditions", "Privacy Notice", and "CDX Helpdesk: (888) 890-1995".

GSPP Logged in as: John Doe, Primary Authorized Official

Home Forms Resources User Management Log Out

HOME

Forms

Persons who must report (unless otherwise specified) include chemical producers and importers under the NAICS Codes Subsection 325 (chemical manufacturing and allied products) and Industry Group 32411 (petroleum refiners). The 8(d) Health & Safety Data Reporting period is one-time with a 60 day reporting period (in certain circumstances, a chemical may remain listed for a longer period but not longer than 2 years).

Studies to be reported will be specified by EPA to include the specific type(s) of health and safety data needed; the chemical grade/purity of the test material (studies involving mixtures are not required unless otherwise specified). Initiated studies are reportable only for study initiation that occurs during the 60 day reporting period.

User Management

Manage the access rights of Supports for each 8(d) Health & Safety Data Reporting form. For every Support, the Authorized Official may grant him/her the ability to edit (but not unlock, create, delete, or submit) the form.

Resources

A helpful guide that describes the 8(d) Health & Safety Data Reporting system and provides useful links for further usability instruction.

Authorized Official

An Authorized Official has the ability to create, amend and unlock forms. The Authorized Official may also submit completed forms either electronically or by mail. Finally, the Authorized Official has the ability to assign Supports to individual forms.

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Screen mock-ups are approximations and may change. Each optional field is outlined by a red text box.

User Management

The ‘User Management’ screen allows the Authorized Official to manage the access rights of Supports to the appropriate forms within the 8(d) Health & Safety Data Reporting application. For every Support, the Authorized Official may give him/her the ability to edit (but not unlock, create, delete, or submit) the form.

The following exhibit shows the screen capture for the ‘User Management’ screen:

Exhibit 1-3 User Management Screen

CDX CENTRAL DATA EXCHANGE

Logged in as: John Doe, Primary Authorized Official

Home Forms Resources **User Management** Log Out

USER MANAGEMENT

Select a Federal Register Notice from the drop-down menu to assign and unassign Supports to specific 8(d) Health & Safety Data Reporting forms.

8(d) Health & Safety Data Reporting Federal Register Notice: Federal Register Notice ▾

8(d) Health and Safety Data Reporting
CASRN:

Assign Users

Unassigned		Assigned
<input type="text"/>	add >> << remove	<input type="text"/>

Save

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Resources

The 'Resources' screen provides the user with the user guide and helpful web links pertaining to 8(d) Health & Safety Data Reporting requirements. The 8(d) Health & Safety Data Reporting User Guide document provides all necessary information for using the 8(d) Health & Safety Data Reporting web application.

The following exhibit shows the screen capture for the 'Resources' screen:

Exhibit 1-4 Resources Screen

The screenshot displays the 'Resources' screen of the CDX system. At the top, there is a dark blue header with the 'GSPP' logo on the left and 'Logged in as: John Doe, Primary Authorized Official' on the right. Below the header is a navigation bar with tabs for 'Home', 'Forms', 'Resources', and 'User Management', with 'Resources' being the active tab. A 'Log Out' link is also present in the top right corner. The main content area is white and contains the following text:

ADDITIONAL INFORMATION

[eReporting User Guide](#)
[User Guide](#)

This guide describes each screen of the 8(d) Health & Safety Data Reporting software and provides information on how to use the system to complete the form. The guide also contains instructions for registering with CDX, submitting the form, and modifying a previous submission. You can download and print the 8(d) Health & Safety Data Reporting User Guide for quick reference.

Helpful Web Links

[CDX Homepage](#)
[TSCA Chemical Substances Inventory](#)

If you need assistance, please call (202) 564-3011 or e-mail the [TSCA-Hotline](mailto:TSCA-Hotline@epamail.epa.gov) at TSCA-Hotline@epamail.epa.gov.

At the bottom of the page, there is a dark blue footer bar with the following links: [CDX Homepage](#) | [MyCDX Homepage](#) | [EPA Homepage](#) | [Terms and Conditions](#) | [Privacy Notice](#) | [CDX Helpdesk: \(888\) 890-1995](#)

Screen mock-ups are approximations and may change. Each optional field is outlined by a red text box.

8(d) Health & Safety Data Reporting Forms

The '8(d) Health & Safety Data Reporting' form screen presents the user with a list of forms submitted or started by the AO during the current reporting cycle. Additionally, AOs may begin a new submission from this screen.

The following exhibit shows the screen capture for the 'Forms' screen:

Exhibit 1-5 Forms Screen

Logged in as: John Doe, Primary Authorized Official

Home Forms Resources User Management Log Out

8(d) Health & Safety Data Reporting

Federal Register Notice	CASRN	Status	Modify Date	Submission Date	Copy of Record	Action
75 FR 773 January 6, 2010	3425-22-1	Submitted	09/17/2010	09/17/2010	↓	🔒

Select Federal Register Notice:

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Create Passphrase

The 'Create Passphrase' screen allows a user with the AO role to start a new submission and associate a passphrase with the submission. The screen also provides instruction about selecting a valid passphrase.

The following exhibit shows the screen capture for the 'Create Passphrase' screen:

Exhibit 1-6 Create Passphrase Screen

GSPP Logged in as: John Doe, Primary Authorized Official [Log Out](#)

CREATE PASSPHRASE

Please create a passphrase that is at least 10 characters in length. To better protect your account, your passphrase should contain a combination of letters and numbers. Your passphrase may include spaces, but should not contain special characters (for example, +, ?, and *).

New Passphrase:

Confirm New Passphrase:

A passphrase can only be created by an Authorized Official for the reporting company. Your passphrase will be used as an encryption key to protect the contents of your data. As an Authorized Official, you are responsible for remembering your passphrase and distributing it to Supports for your company.

Note: If you lose or forget your passphrase, you will not be able to access your 8(d) Health & Safety Data Reporting form to print, submit, or make changes. You will need to complete a new 8(d) Health & Safety Data Reporting form and create a new passphrase. For security reasons, the system administrator will not have access to your passphrase and will not be able to retrieve it or reset it to a new one.

[Cancel](#) [Next](#)

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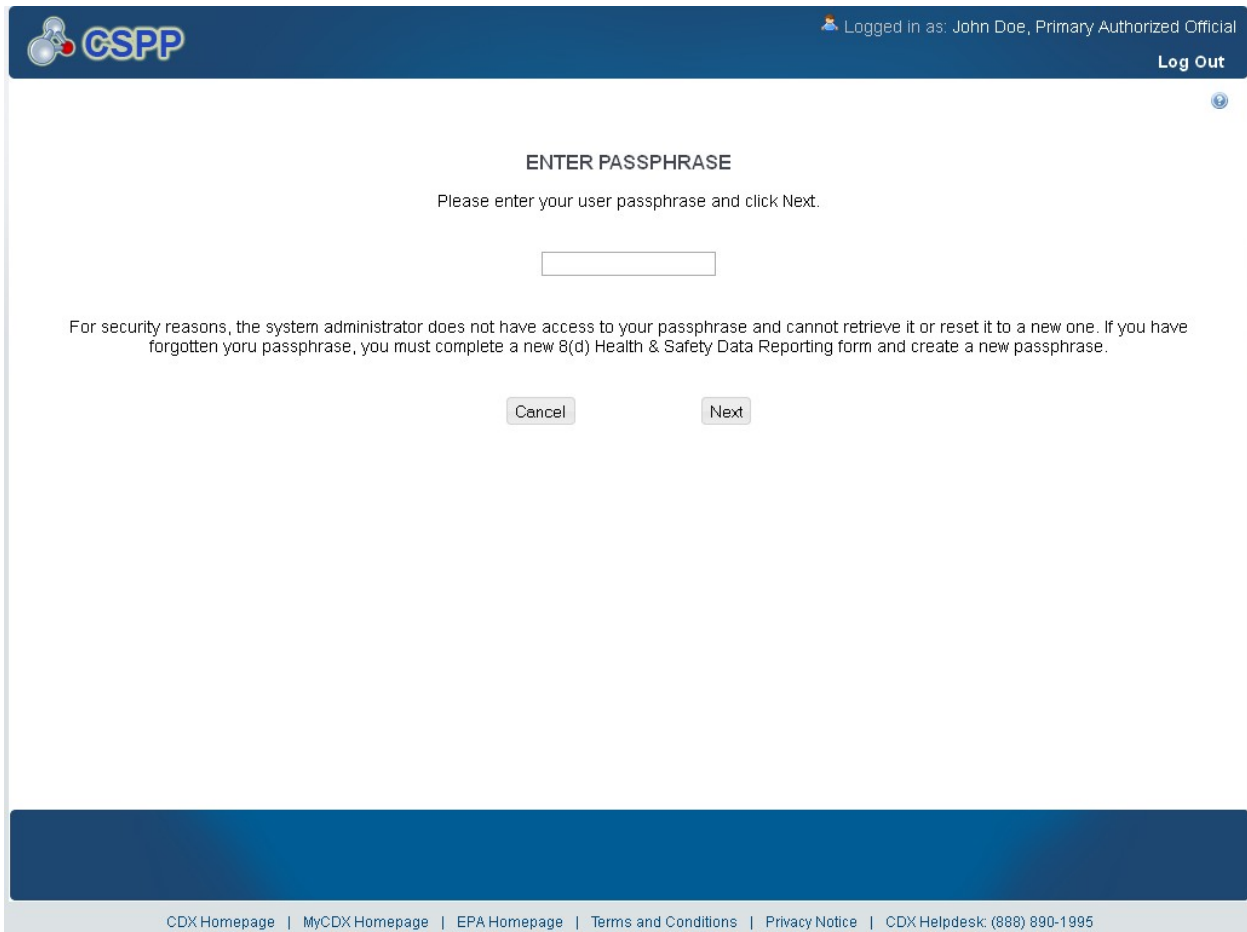
Screen mock-ups are approximations and may change. Each optional field is outlined by a red text box.

Enter Passphrase

The ‘Enter Passphrase’ screen requires the user to enter the passphrase associated with the selected form. The user will have access to the requested form once the entered passphrase has passed validation.

The following exhibit shows the screen capture for the ‘Enter Passphrase’ screen:

Exhibit 1-7 Enter Passphrase Screen



GSPP Logged in as: John Doe, Primary Authorized Official [Log Out](#)

ENTER PASSPHRASE

Please enter your user passphrase and click Next.

For security reasons, the system administrator does not have access to your passphrase and cannot retrieve it or reset it to a new one. If you have forgotten your passphrase, you must complete a new 8(d) Health & Safety Data Reporting form and create a new passphrase.

[Cancel](#) [Next](#)

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Submitting Official Information

The ‘Section A – Submitting Official Information’ allows the user to confirm their contact information. The screen is pre-populated with the information previously entered by the user during CDX registration, but requires the user to enter a response in the ‘Job Title’ field.

The following exhibit shows the screen capture for the ‘Submitting Official Information’ screen:

Exhibit 1-8 Submitting Official Information Screen

8(d) Health and Safety Data Reporting

Primary Authorized Official

Contact Information > Submitting Official Information

SECTION A - SUBMITTING OFFICIAL INFORMATION

This is a submission on behalf of another company:

CBI:

Prefix: Mr.

First Name: John

Middle Initial: D

Last Name: Doe

Suffix:

Company Name: CGI Federal

Job Title:

Telephone: (888) 890-1995

Email Address: cgifederal@cgifederal.com

Mailing Address 1: 12601 Fair Lakes Circle

Mailing Address 2:

City: Fairfax

State: Virginia

Zip: 22033

Next

Add Study

Validate Save Preview Submit

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Screen mock-ups are approximations and may change. Each optional field is outlined by a red text box.

Submitting on Behalf of

The ‘Section A.1 – Submitting on Behalf of’ screen allows users to submit TSCA Section 8(d) Health & Safety Data Reporting forms on behalf of another company covered under the reporting requirement. The ‘Section A.1 – Submitting on Behalf of’ screen presents a list of text fields that allows the user to input contact information of the manufacturing or processing establishment on whose behalf the submission is made.

The following exhibit shows the screen capture for the ‘Submitting on Behalf of’ screen:

Exhibit 1-9 Submitting on Behalf Of

The screenshot displays the GSPP (Global Safety and Performance Portal) interface. At the top, the user is logged in as 'John Doe, Primary Authorized Official'. The navigation menu includes 'Home', 'Forms', 'Resources', 'User Management', and 'Log Out'. The left sidebar shows the '8(d) Health and Safety Data Reporting' section, with 'Submitting on Behalf of' selected under 'Contact Information'. The main content area is titled 'SECTION A.1 - SUBMITTING ON BEHALF OF' and contains the following form fields:

- Prefix:
- First Name:
- Middle Initial:
- Last Name:
- Suffix:
- Company Name:
- Telephone:
- Email Address:
- Mailing Address 1:
- Mailing Address 2:
- City:
- State:
- Zip:

Navigation buttons include 'Previous' and 'Next'. At the bottom, there are buttons for 'Add Study', 'Validate', 'Save', 'Preview', and 'Submit'. The footer contains links to 'CDX Homepage', 'MyCDX Homepage', 'EPA Homepage', 'Terms and Conditions', 'Privacy Notice', and 'CDX Helpdesk: (888) 890-1995'.

Technical Contact

The 'Technical Contact' screen allows the user to add multiple technical contacts, if needed. The system will provide a virtual rolodex address book dropdown menu listing all added technical contacts.

The following exhibit shows the screen capture for the 'Technical Contact' screen:

Exhibit 1-10 Technical Contact Screen

GSPP Logged in as: John Doe, Primary Authorized Official

Home Forms Resources User Management Log Out

8(d) Health and Safety Data Reporting
Primary Authorized Official
75 FR 773 January 6, 2010

Contact Information > Technical Contact

SECTION A.2 - TECHNICAL CONTACT

Select a technical contact from the drop-down menu or enter information for a new contact after clicking Create New Contact. You can select the default button to assign the same technical contact to all your chemical substances or enter a different technical contact for each chemical substance at this site.

CBI:

Prefix: Default Contact:

First Name:

Middle Initial:

Last Name:

Suffix:

Company Name:

Telephone:

Email Address:

Mailing Address 1:

Mailing Address 2:

City:

State:

Zip:

Previous Next

Add Study Validate Save Preview Submit

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Screen mock-ups are approximations and may change. Each optional field is outlined by a red text box.

Chemical Substance Identity of Impurities

The ‘Section B – Chemical Substance Identity of Impurities’ screen allows the user to identify any impurities or additives known to have been present in the substance or listed mixtures as studied. The system provides access to Substance Registry Services Search (SRS) to perform this task.

The following exhibit shows the screen capture for the ‘Chemical Substance Identity of Impurities’ screen:

Exhibit 1-11 Chemical Substance Identity of Impurities Screen

Logged in as: John Doe, Primary Authorized Official

Home Forms Resources User Management Log Out

8(d) Health and Safety Data Reporting

Primary Authorized Official

Chemical Information > Chemical Substance Identity of Impurities

SECTION B - CHEMICAL SUBSTANCE IDENTITY OF IMPURITIES

Identify any impurity or additive known to have been present in the substance or listed mixtures as studied. To search EPA's Substance Registry Services (SRS) for the desired chemical(s), click the magnifying glass below.

SRS	Chemical Identifying Number	Chemical Name	Synonyms	CBI	Actions

Previous Next

Add Study Validate Save Preview Submit

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Substance Registry Services Search

The ‘Substance Registry Services Search’ pop-up helps the user identify chemical impurities in ‘Section B - Chemical Substance Identity of Impurities.’ The user can search for the chemical(s) being reported either by the ‘CASRN’, ‘CAS Index Name or Synonym’, ‘Accession Number’, or ‘Generic Name’.

The following exhibits show the screen captures for the ‘Substance Registry Services Search’ pop-up screen:

Exhibit 1-12 Substance Registry Services Search Pop-Up (Scroll 1)

CSPP

SUBSTANCE REGISTRY SERVICES SEARCH

Enter the specific or partial, currently correct Chemical Abstracts (CA) Index name as listed on the TSCA Inventory and/or the exact corresponding Chemical Abstract Services Registry Number (CASRN) for each reportable chemical substance at your site. Click Search and select the appropriate CA Index name/CASRN combination from EPA's Substance Registry Services (SRS).

Please search by CASRN or CAS Index Name

1. CASRN: Matches exactly

2. CAS Index Name or Other Synonym: Contains

OR

Enter the specific or partial, currently correct Accession Number as listed on the TSCA Inventory and/or the exact or partial corresponding Generic Name for each reportable chemical substance at your site. Click Search and select the appropriate Accession Number/Generic Name combination from EPA's Substance Registry Services (SRS).

Please search by Accession Number and/or Generic Name

1. Accession Number: Contains

2. Generic Name: Contains

Screen mock-ups are approximations and may change. Each optional field is outlined by a red text box.

Exhibit 1-13 Substance Registry Services Search Pop-Up (Scroll 2)

Inventory and/or the exact corresponding Chemical Abstract Services Registry Number (CASRN) for each reportable chemical substance at your site. Click Search and select the appropriate CA Index name/CASRN combination from EPA's Substance Registry Services (SRS).

Please search by CASRN or CAS Index Name

1. CASRN: Matches exactly

2. CAS Index Name or Other Synonym: Contains

OR

Enter the specific or partial, currently correct Accession Number as listed on the TSCA Inventory and/or the exact or partial corresponding Generic Name for each reportable chemical substance at your site. Click Search and select the appropriate Accession Number/Generic Name combination from EPA's Substance Registry Services (SRS).

Please search by Accession Number and/or Generic Name

1. Accession Number: Contains

2. Generic Name: Contains

If you are certain the chemical does not already exist, click the "Chemical Not in SRS" button.

The 'Chemical Not Found in SRS' pop-up allows users to submit information for chemicals not identified in SRS.

Only one of the four fields identified by a blue boundary box is required. Once one of those four fields is populated, the rest become optional.

Exhibit 1-14 Chemical Not Found in SRS Pop-Up

CSPP

CHEMICAL NOT FOUND IN SUBSTANCE REGISTRY SERVICES

Complete all known chemical substance information in the below fields. To add multiple Chemical Synonyms, click the + button to add each synonym. Click the 'OK' button when all known chemical substance information has been fulfilled.

Create New Chemical

Accession Number:	<input type="text"/>
CASRN:	<input type="text"/>
PMN Number:	<input type="text"/>
IUPAC Number:	<input type="text"/>
Chemical Name:	<input type="text"/>
Chemical Synonym:	<input type="text"/> +

OK Cancel

Screen mock-ups are approximations and may change. Each optional field is outlined by a red text box.

Chemical Substance Identity of Impurities

The ‘Section B – Chemical Substance Identity of Impurities’ screen allows the user to identify any impurities or additives known to have been present in the substance or listed mixtures as studied. This screen capture displays how submitted chemical information will display.

The following exhibit shows the screen capture for the ‘Chemical Substance Identity of Impurities’ screen when it is populated:

Exhibit 1-15 Chemical Substance Identity of Impurities Screen (populated)

8(d) Health and Safety Data Reporting

Primary Authorized Official

Home Forms Resources User Management Log Out

Logged in as: John Doe, Primary Authorized Official

Chemical Information > Chemical Substance Identity of Impurities

SECTION B - CHEMICAL SUBSTANCE IDENTITY OF IMPURITIES

Identify any impurity or additive known to have been present in the substance or listed mixtures as studied. To search EPA's Substance Registry Services (SRS) for the desired chemical(s), click the magnifying glass below.

SRS	Chemical Identifying Number	Chemical Name	Synonyms	CBI	Actions
	Accession Number: 123456 ... (+ show more)	Benzene	Synonym1 ... (+ show more)	N	✖

Previous Next

Add Study

Validate Save Preview Submit

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Study Identification

The ‘Study Identificaiton’ screen allows the user to identify and manage documentation related to TSCA Section 8(d) Health & Safety Data Reporting. To identify the submission study type(s) to be reported, the user will check all applicable study types.

Exhibit 1-16 Study Identification Screen

GSPP Logged in as: John Doe, Primary Authorized Official

Home Forms Resources User Management Log Out

8(d) Health and Safety Data Reporting
Primary Authorized Official

Studies > Study Identification

STUDY IDENTIFICATION

Please select which types of studies you will be submitting:

- EPA Request for Further Information
 - Underlying Data
 - Preliminary Reports of Ongoing Studies
 - Copies of Studies
- Full Study Report
- Initiated Studies
- Ongoing Studies
- Robust Summary
- Studies Which are Known but Without Possession of Copies
- Studies Previously Sent to Federal Agencies without Confidentiality Claims

Previous Next

Add Study Validate Save Preview Submit

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Screen mock-ups are approximations and may change. Each optional field is outlined by a red text box.

Submission of Studies – EPA Request for Further Information

The following exhibits show the screen captures for the ‘Submission of Studies – EPA Request for Further Identification’ screens:

Exhibit 1-17 EPA Request for Further Information – Underlying Data Screen

8(d) Health and Safety Data Reporting

Primary Authorized Official

Studies > EPA Request for Further Information > Underlying Data

SECTION C - SUBMISSION OF STUDIES

Underlying Data

3. Browse for the Underlying Data document.

CBI:


Click Add Study to attach a new Underlying Data document.

Underlying Data

File Name	Document Type	CBI	Sanitized	Action
<input type="button" value="Previous"/> <input type="button" value="Next"/>				

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Exhibit 1-18 EPA Request for Further Information – Preliminary Reports of Ongoing Studies Screen (Scroll 1)


Logged in as: John Doe, Primary Authorized Official

[Home](#) | [Forms](#) | [Resources](#) | [User Management](#)

Log Out

8(d) Health and Safety Data Reporting < [Studies > EPA Request for Further Information > Preliminary Reports of Ongoing Studies](#)

SECTION C - SUBMISSION OF STUDIES

Preliminary Reports of Ongoing Studies

3. Browse for the Preliminary Reports of Ongoing Studies document.

CBI:

4. The beginning date of the study, the purpose of the study, types of data to be collected, the anticipated date of completion, and the name and address of the laboratory conducting the study must accompany each entry on the list. Provide the following information for the attached study.

Study Laboratories	
Laboratory Name:	<input type="text"/>
Mailing Address 1:	<input type="text"/>
Mailing Address 2:	<input type="text"/>
City:	<input type="text"/>
State:	<input type="text"/>
Zip:	<input type="text"/>
<input type="button" value="Add Laboratory"/>	

Study Title:

Study Start Date: Study End Date:

Date format: mm/dd/yyyy

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Screen mock-ups are approximations and may change. Each optional field is outlined by a red text box.

Exhibit 1-19 EPA Request for Further Information – Preliminary Reports of Ongoing Studies Screen (Scroll 2)

Logged in as: John Doe, Primary Authorized Official

Log Out
Home Forms Resources User Management

8(d) Health and Safety Data Reporting

Primary Authorized Official

75 FR 773 January 6, 2010

- [-] Contact Information
 - [-] Submitting Official Information
 - [-] Technical Contact
- [-] Chemical Information
 - [-] Chemical Substance
 - [-] Identity of Impurities
- [-] Studies
 - [-] Study Identification
 - [-] EPA Request for Further Information
 - [-] Underlying Data
 - [-] Preliminary Reports of Ongoing Studies**
 - [-] Copies of Studies
 - [-] Full Study Report
 - [-] Initiated Studies
 - [-] Ongoing Studies
 - [-] Robust Summary
 - [-] Studies Which are Known but Without Possession of Copies
 - [-] Studies Previously Sent to Federal Agencies without Confidentiality Claims

State:

Zip:

Study Title:

Study Start Date: Study End Date:

Date format: mm/dd/yyyy

Study Purpose:

Data to be Collected:

Anticipated Completion Date:

Date format: mm/dd/yyyy

Click Add Study to attach a new Preliminary Reports of Ongoing Studies document.

Preliminary Reports of Ongoing Studies

File Name	Document Type	Study Title	Laboratory	CBI	Sanitized	Action
<div style="display: flex; justify-content: center; gap: 10px;"> <input type="button" value="Previous"/> <input type="button" value="Next"/> </div>						

Validate

Save

Preview

Submit

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Exhibit 1-20 EPA Request for Submission of Further Information – Copies of Studies Screen (Scroll 1)

CDX GSPP Logged in as: John Doe, Primary Authorized Official [Home](#) [Forms](#) [Resources](#) [User Management](#) [Log Out](#)

8(d) Health and Safety Data Reporting Studies > EPA Request for Further Information > Copies of Studies

Primary Authorized Official

SECTION C - SUBMISSION OF STUDIES

Copies of Studies

3. Browse for the Copies of Studies document.

CBI: [Browse](#)

4. The name and address of any person known to possess a copy of the unpublished study must accompany each entry on the list. Provide the following information for the attached study:

Study Contacts	
Contact Name	Action
Prefix: <input type="text"/>	
First Name: <input type="text"/>	
Last Name: <input type="text"/>	
Suffix: <input type="text"/>	
Telephone: <input type="text"/>	
Email Address: <input type="text"/>	
Mailing Address 1: <input type="text"/>	
Mailing Address 2: <input type="text"/>	
City: <input type="text"/>	
State: <input type="text"/>	
Zip: <input type="text"/>	
Add Contact	

[Add Study](#) [Validate](#) [Save](#) [Preview](#) [Submit](#)

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Screen mock-ups are approximations and may change. Each optional field is outlined by a red text box.

Exhibit 1-21 EPA Request for Submission of Further Information – Copies of Studies Screen (Scroll 2)

8(d) Health and Safety Data Reporting
Primary Authorized Official

Logged in as: John Doe, Primary Authorized Official

Home Forms Resources User Management Log Out

Prefix:
 First Name:
 Last Name:
 Suffix:
 Telephone:
 Email Address:
 Mailing Address 1:
 Mailing Address 2:
 City:
 State:
 Zip:
 Add Contact

Contact Name Action

Study Title

Click Add Study to attach a new Copies of Studies document.

Add Study

Copies of Studies

File Name	Document Type	Study Title	Contact Name	CBI	Sanitized	Action
		Previous	Next			

Add Study

Validate Save Preview Submit

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Full Study Report

The 'Full Study Report' screen allows the user to submit or make available for review entire, completed studies that have been requested by the EPA. This screen capture displays how submitted 'Full Study Reports' will display.

The following exhibit shows the screen capture for the 'Full Study Report' screen:

Exhibit 1-22 Full Study Report Screen

8(d) Health and Safety Data Reporting **Studies > Full Study Report** Logged in as: John Doe, Primary Authorized Official

Home Forms Resources User Management Log Out

Primary Authorized Official

SECTION C - SUBMISSION OF STUDIES

Full Study Report

2. Browse for the Full Study Report document.

CBI: **Browse**

Click Add Study to attach a new Full Study Report document.

Add Study

Full Study Report

File Name	Document Type	CBI	Sanitized	Action
				Previous Next

Add Study **Validate** **Save** **Preview** **Submit**

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Screen mock-ups are approximations and may change. Each optional field is outlined by a red text box.

Initiated Studies

The ‘Initiated Studies’ screen allows the user to submit a list of ongoing health and safety studies being conducted by or initiated for them to the EPA for each of the listed substances or mixtures called for in the issued Federal Register Notice Citation.

The following exhibits show the screen captures for the ‘Initiated Studies’ screen:

Exhibit 1-23 Initiated Studies (Scroll 1)


The screenshot displays the GSPP (Global Safety and Performance Portal) interface. At the top, the user is logged in as 'John Doe, Primary Authorized Official'. The navigation menu includes 'Home', 'Forms', 'Resources', 'User Management', and 'Log Out'. The main content area is titled 'SECTION C - SUBMISSION OF STUDIES' and 'Initiated Studies'. A sidebar on the left lists various options under '8(d) Health and Safety Data Reporting', with 'Initiated Studies' selected. The main form area contains the following elements:

- Step 2: 'Browse for the Initiated Studies document.' with a 'Browse' button and a 'CBI:
- Step 4: 'Provide the following information for the attached study.' with a 'Study Laboratories' section containing a table with columns 'Laboratory' and 'Action'. Fields include 'Laboratory Name', 'Mailing Address 1', 'Mailing Address 2' (highlighted with a red box), 'City', 'State', and 'Zip'. An 'Add Laboratory' button is also present.
- Fields for 'Study Title', 'Study Start Date', and 'Study End Date' (with a date format of mm/dd/yyyy).
- A 'Study Purpose' text area.

At the bottom of the form, there is an 'Add Study' button and a row of action buttons: 'Validate' (with a green checkmark), 'Save', 'Preview', and 'Submit' (with a database icon).

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Exhibit 1-24 Initiated Studies (Scroll 2)


Logged in as: John Doe, Primary Authorized Official

Home
Forms
Resources
User Management
Log Out

8(d) Health and Safety Data Reporting

Primary Authorized Official

- [-] **Contact Information**
 - Submitting Official Information
 - Submitting on Behalf of
 - Technical Contact
- [-] **Chemical Information**
 - Chemical Substance Identity of Impurities
- [-] **Studies**
 - Study Identification
 - [-] **EPA Request for Further Information**
 - Copies of Studies
 - Initiated Studies**
 - Submitter Requests

City:

State:

Zip:

Study Title:

Study Start Date: Study End Date:

Date format: mm/dd/yyyy

Study Purpose:

Data to be Collected:


Anticipated Completion Date:


Date format: mm/dd/yyyy


Click Add Study to attach a new Initiated Studies document.


Initiated Studies

File Name	Document Type	Study Title	Laboratory	CBI	Sanitized	Action
<div style="display: flex; justify-content: center; gap: 10px;"> <input type="button" value="Previous"/> <input type="button" value="Next"/> </div>						


Validate


Save


Preview


Submit

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Screen mock-ups are approximations and may change. Each optional field is outlined by a red text box.

Ongoing Studies

The ‘Ongoing Studies’ screen allows the user to submit a list of ongoing health and safety studies being conducted by or initiated for them to EPA for each of the listed substances or mixtures called for in the issued Federal Register Notice Citation.

The following exhibits show the screen captures for the ‘Ongoing Studies’ screen:

Exhibit 1-25 Ongoing Studies (Scroll 1)

8(d) Health and Safety Data Reporting Primary Authorized Official

Home Forms Resources User Management Log Out

Logged in as: John Doe, Primary Authorized Official

Studies > Ongoing Studies

SECTION C - SUBMISSION OF STUDIES

Ongoing Studies

2. Browse for the Ongoing Studies document.

CBI: Browse

4. Provide the following information for the attached study.

Study Laboratories

Laboratory	Action
Laboratory Name: <input type="text"/>	
Mailing Address 1: <input type="text"/>	
Mailing Address 2: <input type="text"/>	
City: <input type="text"/>	
State: <input type="text"/>	
Zip: <input type="text"/>	
<input type="button" value="Add Laboratory"/>	

Study Title:


Study Start Date: Study End Date:

Date format: mm/dd/yyyy

Study Purpose:

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Exhibit 1-26 Ongoing Studies (Scroll 2)


Logged in as: John Doe, Primary Authorized Official
Log Out

Home
Forms
Resources
User Management

8(d) Health and Safety Data Reporting

Primary Authorized Official

- [-] Contact Information
 - [-] Submitting Official Information
 - [-] Submitting on Behalf of
 - [-] Technical Contact
- [-] Chemical Information
 - [-] Chemical Substance Identity of Impurities
- [-] Studies
 - [-] Study Identification
 - [-] EPA Request for Further Information
 - [-] Copies of Studies
 - [-] Ongoing Studies
 - [-] Submitter Requests

City:

State:

Zip:

Study Title:

Study Start Date: Study End Date:

Date format: mm/dd/yyyy

Study Purpose:

Data to be Collected:


Anticipated Completion Date:


Date format: mm/dd/yyyy


Click Add Study to attach a new Ongoing Studies document.


Ongoing Studies

File Name	Document Type	Study Title	Laboratory	CBI	Sanitized	Action
<div style="display: flex; justify-content: space-between; width: 100%;"> <input type="button" value="Previous"/> <input type="button" value="Next"/> </div>						


Validate


Save


Preview


Submit

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Screen mock-ups are approximations and may change. Each optional field is outlined by a red text box.

Submission of Studies - Robust Summary

The ‘Robust Summary’ screen allows the user to upload a Robust Summary list for Full Study Reports. The system allows the user to upload a Robust Summary and/or the Full Study Report. This screen capture displays how a submitted ‘Robust Study’ will display.

The following exhibit shows the screen capture for the ‘Submission of Studies – Robust Summary’ screen:

Exhibit 1-27 Robust Summary Screen

The screenshot shows the GSPP (Global Study Processing Platform) interface. At the top, it indicates the user is logged in as John Doe, a Primary Authorized Official. The main navigation bar includes Home, Forms, Resources, User Management, and Log Out. The left sidebar contains a tree view with categories like Contact Information, Chemical Information, Studies, EPA Request for Further Information, and Submitter Requests. The main content area is titled 'SECTION C - SUBMISSION OF STUDIES' and 'Robust Summary'. It contains a 'Browse' button and an 'Add Study' button. Below these is a table with columns for File Name, Document Type, and Action, with 'Previous' and 'Next' buttons. At the bottom, there is a dark blue bar with icons for Add Study, Validate, Save, Preview, and Submit. The footer contains links to CDX Homepage, MyCDX Homepage, EPA Homepage, Terms and Conditions, Privacy Notice, and CDX Helpdesk.

Submission of Studies – Studies which are known but are without Possession of Copies

The ‘Studies Which are Known but are without Possession of Copies’ screen allows the user to submit a list of unpublished health and safety studies known to them , but which they do not have copies.

The following exhibits show the screen captures for the ‘Submission of Studies – Studies which are known but are without Possession of Copies’ screen:

Exhibit 1-28 Studies Which are Known but are without Possession of Copies (Scroll 1)

8(d) Health and Safety Data Reporting

Primary Authorized Official

Home Forms Resources User Management Log Out

Logged in as: John Doe, Primary Authorized Official

Studies > Studies Which are Known but Without Possession of Copies

SECTION C - SUBMISSION OF STUDIES

Studies Which are Known but Without Possession of Copies

2. Browse for the Studies Which are Known but Without Possession of Copies document.

CBI: Browse

4. Provide the following information for the attached study:

Study Contacts

Contact Name	Action
Prefix: <input type="text"/>	
First Name: <input type="text"/>	
Last Name: <input type="text"/>	
Suffix: <input type="text"/>	
Telephone: <input type="text"/>	
Email Address: <input type="text"/>	
Mailing Address 1: <input type="text"/>	
Mailing Address 2: <input type="text"/>	
City: <input type="text"/>	
State: <input type="text"/>	
Zip: <input type="text"/>	
<input type="button" value="Add Contact"/>	

Add Study

Validate Save Preview Submit

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Screen mock-ups are approximations and may change. Each optional field is outlined by a red text box.

Exhibit 1-29 Studies Which are Known but are without Possession of Copies (Scroll 2)

Logged in as: John Doe, Primary Authorized Official
Log Out

Home
Forms
Resources
User Management

8(d) Health and Safety Data Reporting

Primary Authorized Official

- [-] **Contact Information**
 - Submitting Official Information
 - Submitting on Behalf of
 - Technical Contact
- [-] **Chemical Information**
 - Chemical Substance Identity of Impurities
- [-] **Studies**
 - Study Identification
 - [-] **EPA Request for Further Information**
 - Copies of Studies
 - Studies Which are Known but Without Possession of Copies**
 - Submitter Requests

Prefix:
Contact Name
Action

First Name:

Last Name:

Suffix:

Telephone:

Email Address:

Mailing Address 1:

Mailing Address 2:

City:

State:

Zip:

Study Title

Click Add Study to attach a new Studies Which are Known but Without Possession of Copies document.

Studies Which are Known but Without Possession of Copies

File Name	Document Type	Study Title	Contact Name	CBI	Sanitized	Action
<div style="display: flex; justify-content: space-between; width: 100%;"> <input type="button" value="Previous"/> <input type="button" value="Next"/> </div>						

Validate

Save

Preview

Submit

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Submission of Studies – Studies previously sent to Federal Agencies without Confidentiality Claims

The ‘Studies Previously Sent to Federal Agencies without Confidentiality Claims’ screen allows the user to submit a list of unpublished studies which have been sent to a Federal Agency without claims of confidentiality to EPA.

The following exhibits show the screen captures for the ‘Submission of Studies – Studies Previously Sent to Federal Agencies without Confidentiality Claims’ screen:

Exhibit 1-30 Studies Previously Sent to Federal Agencies without Confidentiality Claims (Scroll 1)

GSPP | Logged in as: John Doe, Primary Authorized Official | Home | Forms | Resources | User Management | Log Out

8(d) Health and Safety Data Reporting | **Studies > Studies Previously Sent to Federal Agencies without Confidentiality Claims**

SECTION C - SUBMISSION OF STUDIES

Studies Previously Sent to Federal Agencies without Confidentiality Claims

2. Browse for the Studies Previously Sent to Federal Agencies without Confidentiality Claims document.

CBI:

5. Provide the following information for the attached study:

Study Title:

Study Submission Date: mm/dd/yyyy

Agency:

6. Provide the following information for each Agency Contact.

Contact Name	Action
Prefix: <input type="text"/>	
First Name: <input type="text"/>	
Last Name: <input type="text"/>	
Suffix: <input type="text"/>	
Telephone: <input type="text"/>	
Email Address: <input type="text"/>	
Mailing: <input type="text"/>	

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Screen mock-ups are approximations and may change. Each optional field is outlined by a red text box.

Exhibit 1-31 Studies Previously Sent to Federal Agencies without Confidentiality Claims (Scroll 2)

The screenshot displays the GSPP (Global Study Processing Platform) interface. At the top, the user is logged in as John Doe, a Primary Authorized Official. The navigation menu includes Home, Forms, Resources, and User Management. The main content area is titled "8(d) Health and Safety Data Reporting" and is managed by a Primary Authorized Official. A sidebar on the left contains a tree view with categories: Contact Information (Submitting Official Information, Submitting on Behalf of, Technical Contact), Chemical Information (Chemical Substance, Identity of Impurities), Studies (Study Identification, EPA Request for Further Information, Copies of Studies, Studies Previously Sent to Federal Agencies without Confidentiality Claims, Submitter Requests), and Submitter Requests.

The main form area contains the following fields:

- Prefix:
- First Name:
- Last Name:
- Suffix:
- Telephone:
- Email Address:
- Mailing Address 1:
- Mailing Address 2: (This field is highlighted with a red border in the original image)
- City:
- State:
- Zip:

Below the form is an "Add Contact" button. A message states: "Click Add Study to attach a new Studies Previously Sent to Federal Agencies without Confidentiality Claims document." Below this is an "Add Study" button.

The table below is titled "Studies Previously Sent to Federal Agencies without Confidentiality Claims". It has the following columns: File Name, Document Type, Study Title, Federal Agency, Agency Contact, CBI, Sanitized, and Action. The table currently contains no data rows. Navigation buttons "Previous" and "Next" are located below the table.

At the bottom of the interface, there is a blue bar with an "Add Study" button on the left and four icons with labels: a green checkmark for "Validate", a floppy disk for "Save", a computer monitor for "Preview", and a database icon for "Submit".

The footer contains the following text: "CDX Homepage | MyCDX Homepage | EPA Homepage | Terms and Conditions | Privacy Notice | CDX Helpdesk: (888) 890-1995"

Submitter Requests

The ‘Submitter Requests’ screen allows the user to request an ‘Extension of Time’ or a ‘Withdrawal of Chemical.’ The user can select one or more of the option(s) that apply such as, ‘Request for Extension of Time’, or ‘Request for Withdrawal of Chemical’ and upload supporting documentation.

The following exhibit shows the screen capture for the ‘Submitter Requests’ screen:

Exhibit 1-32 Submitter Requests Screen

The screenshot displays the GSPP (Global Safety and Performance Portal) interface for Submitter Requests. At the top, the user is logged in as John Doe, a Primary Authorized Official. The navigation menu on the left includes sections for Contact Information, Chemical Information, Studies, and Submitter Requests. The main content area is titled "SECTION D - SUBMITTER REQUESTS" and prompts the user to "Select the appropriate request below and upload the the corresponding document." Below this instruction, there is a dropdown menu, a "Browse" button, and an "Upload Document" button. A table with columns for "File Name", "Request Type", and "Actions" is partially visible, with a "Previous" button below it. At the bottom of the screen, a toolbar contains buttons for "Add Study", "Validate", "Save", "Preview", and "Submit". The footer of the page provides links to CDX Homepage, MyCDX Homepage, EPA Homepage, Terms and Conditions, Privacy Notice, and CDX Helpdesk (888) 890-1995.

Screen mock-ups are approximations and may change. Each optional field is outlined by a red text box.

Substantiation Summary

The ‘Substantiation Summary’ screen allows the user to clearly identify the material subject to CBI for each document. The user will manually enter such information into the text box displayed within the ‘Substantiation’ screen.

The following exhibit shows the screen capture for the ‘Substantiation Summary’ screen:

Exhibit 1-33 Substantiation Summary Screen

8(d) Health and Safety Data Reporting | Substantiation Summary

Logged in as: John Doe, Primary Authorized Official | Log Out

Home | Forms | Resources | User Management

Primary Authorized Official

Substantiation Summary

To pass validation, substantiation must be completed for all items claimed CBI before submission. Click **Pending** within the table to complete substantiation for each item below. All items with a Substantiation Status of Complete have been completed. Click **Complete** within the table to view completed substantiation.

Section B.2 Chemical Substance Identity of Impurities

CASRN	Chemical Name	Substantiation Status
51-25-8	2,4-Dinitrophenol	Pending
6379123-37-9	Arsonic acid methyl compd. With 1-octanamine (1:1)	Complete

Section C – Submission of Studies

File Name	Study Classification	Substantiation Status
Upload1.doc	Initiated Studies	Complete
Upload2.doc	Completed Studies	Pending

Previous

Add Study | Validate | Save | Preview | Submit

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Submission of Studies Substantiation

The ‘Substantiation screen allows the user to clearly identify the material subject to CBI for each document. The user will manually enter such information into the text box displayed within the ‘Substantiation’ screen.

The following exhibit shows the screen capture for the ‘Submission of Studies Substantiation’ screen:

Exhibit 1-34 Submission of Studies Substantiation Screen

GSPP Logged In as: John Doe, Primary Authorized Official

Home Forms Resources User Management Log Out

8(d) Health and Safety Data Reporting > Submission of Studies Substantiation

SUBMISSION OF STUDIES SUBSTANTIATION

Any respondent who wishes to assert a claim that part of a study should be withheld from disclosure because disclosure would reveal a confidential process or quantitative mixture composition should briefly state the basis of the claim, e.g. by saying "reveals confidential mixture proportion data," and clearly identify the material subject to the claim

Any respondent may assert a confidentiality claim for company name or address, financial statistics, and product codes used by a company. This information will not be subject to the disclosure requirements of section 14(b) of TSCA

Information other than company name or address, financial statistics and product codes used by a company, which is contained in a study, the disclosure of which would clearly be an unwarranted invasion of personal privacy (such as individual medical records), will be considered confidential by EPA as provided in Title 5, United States Code, section 552(b)(6).

To assert a claim of confidentiality for data contained in a submitted document, the respondent must submit two copies of the document:

One copy must be complete. In that copy, the respondent must indicate what data, if any, are claimed as confidential by bracketing or underlining the specific information. Each page containing data claimed as confidential must also contain a brief statement for the basis of the claim as well as a label such as "confidential," "proprietary" or "trade secret."

The second copy must be complete, except that all information claimed as confidential in the first copy must be deleted. The second copy will be immediately subject to public disclosure.

Failure to furnish a second copy when information is claimed as confidential in the first copy will be considered a presumptive waiver of the claim of confidentiality. EPA will notify the respondent by certified mail that a finding of a presumptive waiver of the claim of confidentiality has been made. The respondent will be given 30 days from the date of his or her receipt of this notification to submit the required second copy. If the respondent fails to submit the second copy within the 30 days, EPA will place the first copy in the public file.

Previous

Add Study Validate Save Preview Submit

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