

TSCA Section 8(d) Health Safety Data Reporting

Reporting Tool Screen Shots EPA From 9600-032

OMB Control No. 2027-0224

Expiration 4/30/2029

Chemical Information Submission System Screen

CSPP

Logged in as: John Doe, Primary Support
Log Out

CHEMICAL INFORMATION SUBMISSION SYSTEM

TSCA Section 8(d) Health & Safety Data Reporting

OK

The software includes embedded help files and downloadable user manual to guide you through the 8(d) Health & Safety Data Reporting submission process.

EPA has the authority to publicize rules to require producers, importers, and processors to submit lists and/or copies of ongoing and completed unpublished health and safety studies.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 established to gather health and safety information on chemical substances and mixtures needed by EPA to carry out its TSCA mandates (i.e., to support OPPT's Existing Chemicals Program and Chemical Testing Program and to set priorities for TSCA risk assessment/management activities). OPPT has also used its TSCA Section 8(d) authority to gather information needed by other federal agencies and EPA program offices. 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 per year). Non-ITC chemicals can be added to the Section 8(d) rule via notice and comment rulemaking.

Paperwork Reduction Act Notice

This collection of information is approved by OMB under the Paperwork Reduction Act, 44 U.S.C. 3501 et seq. (OMB Control No. 2070-0224). Responses to this collection of information are mandatory for certain persons, as specified at 40 CFR Part 712. An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. The public reporting and recordkeeping burden for this collection of information is estimated to be 0.4 to 107.5 hours per response. Send comments on the Agency's need for this information, the accuracy of the provided burden estimates and any suggested methods for minimizing respondent burden to the Data & Enterprise Programs Deputy Division Director, U.S. Environmental Protection Agency (2821T), 1200 Pennsylvania Ave., NW, Washington, D.C. 20460. Include the OMB control number in any correspondence. Do not send the completed form to this address.

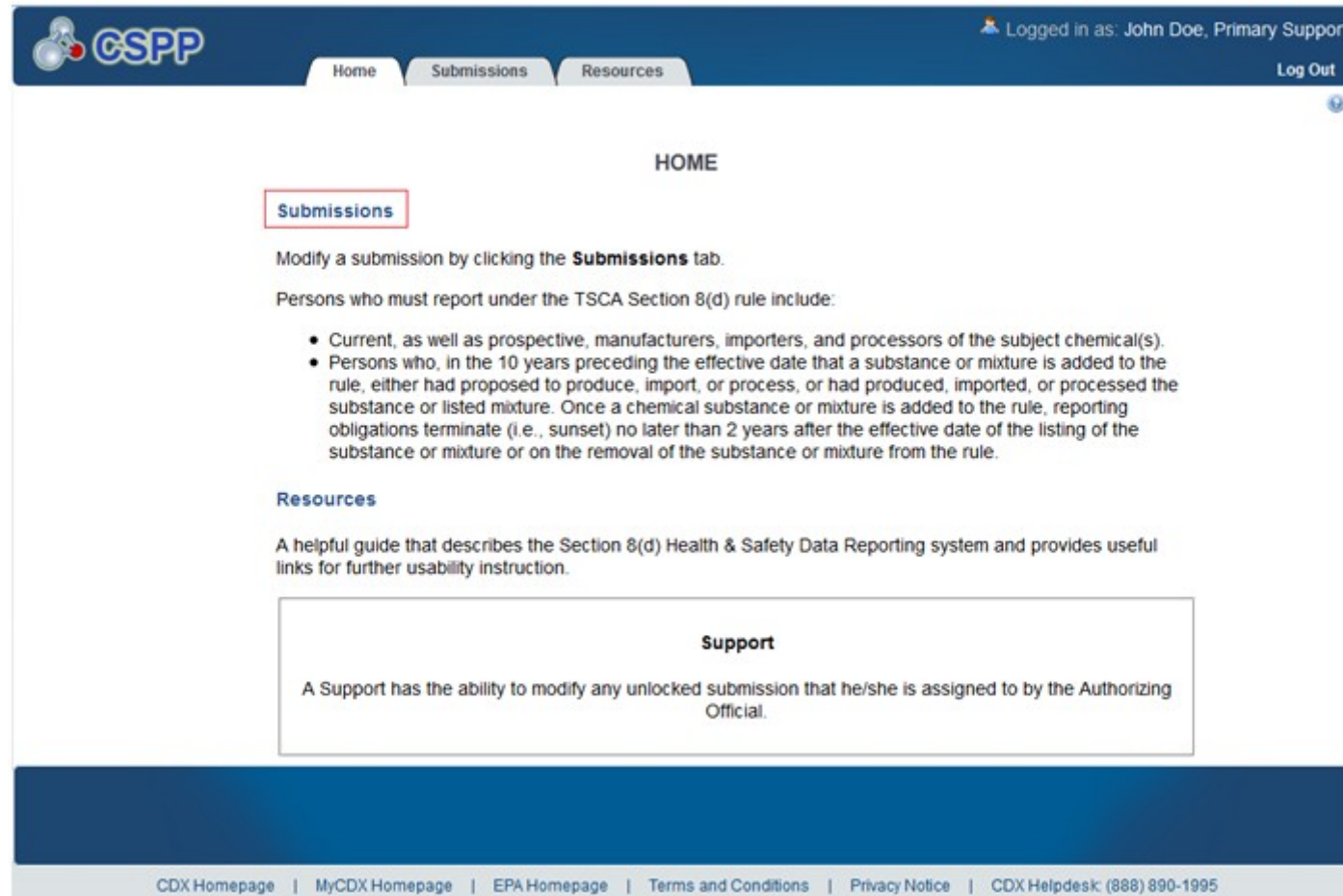
Authority

The Government Paperwork Elimination Act (GPEA) (44 U.S.C. 3504) provides that, when practicable, Federal organizations use electronic forms, electronic filings, and electronic signatures to conduct official business with the public. EPA's Cross-Media Electronic Reporting Regulation (CROMERR) (40 CFR part 3) (Ref. 2), provides that any requirement in title 40 of the CFR to submit a report directly to EPA can be satisfied with an electronic submission that meets certain conditions once the Agency published a document in the **Federal Register** announcing that EPA is prepared to receive certain documents in electronic form. For more information about CROMERR, go to <http://www.epa.gov/cromerr>.

EPA Form 9600-032

Home Screen

OMB No. 2070-0224
Exp 4/30/2029



The screenshot displays the CSPP (Chemical Safety and Hazard Investigation) Home Screen. At the top, a dark blue header bar contains the CSPP logo on the left, a navigation menu with 'Home', 'Submissions', and 'Resources' in the center, and a user status area on the right showing 'Logged in as: John Doe, Primary Support' and a 'Log Out' link. Below the header, the main content area is titled 'HOME'. A red rectangular box highlights the 'Submissions' link in the left sidebar. The main content area includes a paragraph about modifying submissions, a list of persons who must report under the TSCA Section 8(d) rule, a 'Resources' section with a helpful guide link, and a 'Support' section explaining the role of a support user. The footer is a dark blue bar with a light blue background containing links to 'CDX Homepage', 'MyCDX Homepage', 'EPA Homepage', 'Terms and Conditions', 'Privacy Notice', and 'CDX Helpdesk: (888) 890-1995'.

CSPP

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

[Home](#) [Submissions](#) [Resources](#)

HOME

[Submissions](#)

Modify a submission by clicking the **Submissions** tab.

Persons who must report under the TSCA Section 8(d) rule include:

- Current, as well as prospective, manufacturers, importers, and processors of the subject chemical(s).
- Persons who, in the 10 years preceding the effective date that a substance or mixture is added to the rule, either had proposed to produce, import, or process, or had produced, imported, or processed the substance or listed mixture. Once a chemical substance or mixture is added to the rule, reporting obligations terminate (i.e., sunset) no later than 2 years after the effective date of the listing of the substance or mixture or on the removal of the substance or mixture from the rule.

Resources

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

Support

A Support has the ability to modify any unlocked submission that he/she is assigned to by the Authorizing Official.

[CDX Homepage](#) | [MyCDX Homepage](#) | [EPA Homepage](#) | [Terms and Conditions](#) | [Privacy Notice](#) | [CDX Helpdesk: \(888\) 890-1995](#)

Submissions Screen

OMB No. 2070-0224
Exp 4/30/2029

 GSPP

Logged in as: John Doe, Primary Support

Log Out

Home Submissions Resources

SECTION 8(d) HEALTH & SAFETY DATA REPORTING

- To edit an **In Progress** submission, click the submission alias link in the **Submission Alias** column in the table below.

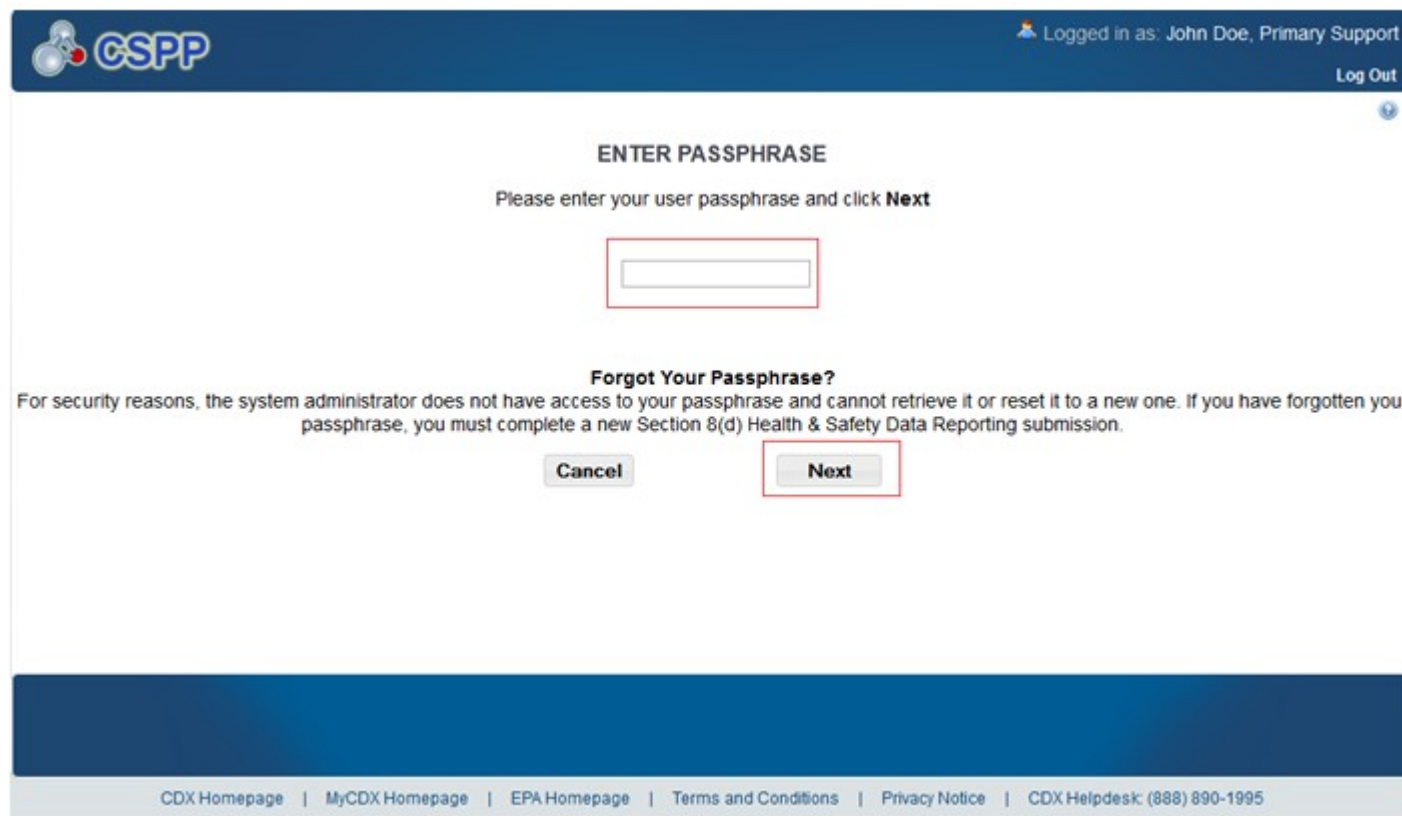
3 items found. Page 1 of 1 Items per page: 25

Submission Alias	FRN/CFR	CASRN	Status	Last Modified	Submission Date
Tue Jun 04 08:55:42 EDT 2013	40 FR 345		In Progress	06/04/2013	
Tue Jun 04 08:59:24 EDT 2013	40 FR 12345		In Progress	06/04/2013	06/04/2013
Tue Jun 04 40 FR 567	40 FR 567		In Progress	06/04/2013	

Export options: [CSV](#) | [Excel](#) | [XML](#) | [PDF](#)

CDX Homepage | MyCDX Homepage | EPA Homepage | Terms and Conditions | Privacy Notice | CDX Helpdesk: (888) 890-1995

Passphrase Screen



The screenshot shows a web interface for the CSPP (Chemical Safety and Hazard Investigation Program). At the top, a dark blue header bar contains the CSPP logo on the left and the text "Logged in as: John Doe, Primary Support" on the right, with a "Log Out" link below it. The main content area is white and features the heading "ENTER PASSPHRASE" in bold. Below this heading is the instruction "Please enter your user passphrase and click **Next**". A text input field is centered below the instruction, highlighted with a red rectangular box. Further down, the section "Forgot Your Passphrase?" is displayed in bold, followed by a paragraph explaining that the system administrator cannot retrieve or reset the passphrase and that a new Section 8(d) Health & Safety Data Reporting submission must be completed if the passphrase is forgotten. At the bottom of this section are two buttons: "Cancel" and "Next", both highlighted with red rectangular boxes. The footer of the page is a dark blue bar with a light gray footer area containing links for "CDX Homepage", "MyCDX Homepage", "EPA Homepage", "Terms and Conditions", "Privacy Notice", and "CDX Helpdesk: (888) 890-1995".

CSPP Logged in as: John Doe, Primary Support
Log Out

ENTER PASSPHRASE
Please enter your user passphrase and click **Next**

Forgot Your Passphrase?
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 Section 8(d) Health & Safety Data Reporting submission.

Cancel **Next**

CDX Homepage | MyCDX Homepage | EPA Homepage | Terms and Conditions | Privacy Notice | CDX Helpdesk: (888) 890-1995

General Submission Information Screen

OMB No. 2070-0224
Exp 4/30/2029

CSPP Logged in as: John Doe, Primary Support Log Out

Home Submissions Resources

Section 8(d) Health & Safety Data Reporting

Section 8(d) Health & Safety Data Reporting > General Submission Information

GENERAL SUBMISSION INFORMATION

You have chosen to report under **75 FR 17645**. Based on this selection, all data entered in this form should pertain to the CFR/FRN selected on the Submissions screen.

The submission alias is an optional field that changes the submission name on the **Submissions Screen**. Its purpose is to make it easier to distinguish between multiple submissions. If an alias is not selected, the field will default to the date and time it was created. The submission alias may be changed at any time.

Please enter a **Submission Alias** in the field below:

Submission Alias:

Next

Validate Save Preview

CDX Homepage | MyCDX Homepage | EPA Homepage | Terms and Conditions | Privacy Notice | CDX Helpdesk: (888) 890-1995

Technical Contact Screen

CSPP

Logged in as: John Doe, Primary Support

Home Submissions Resources Log Out

Section 8(d) Health & Safety Data Reporting

Section 8(d) Health & Safety Data Reporting > Contact Information > Technical Contact Information

TECHNICAL CONTACT INFORMATION

Identify the technical contact who is capable of answering questions related to the chemical(s) submitted to EPA within this submission. Identify if this submission is being submitted on behalf of another company by checking the checkbox. Click the Copy button below to import your CDX registration contact information.

This is a submission on behalf of another company: ☐

Copy CDX Registration

CBI: ☐

Prefix:

First Name:

Middle Initial:

Last Name:

Suffix:

Company Name:

Phone Number: Ext:

(Do not enter any dashes (-) in Phone Number field above.)

Email Address:

Mailing Address 1:

Street address, P.O. box, company name, etc.

Mailing Address 2:

Apartment, suite, etc.

City:

State:

Postal Code:

Country:

CDX Homepage | MyCDX Homepage | EPA Homepage | Terms and Conditions | Privacy Notice | CDX Helpdesk: (888) 890-1995

OMB No. 2070-0224
Exp 4/30/2029

Submitting on Behalf of Company Screen

OMB No. 2070-0224
Exp 4/30/2029

CSPP

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

[Home](#) [Submissions](#) [Resources](#)

Section 8(d) Health & Safety Data Reporting

Primary Support

- General Submission Information
- Contact Information
 - Technical Contact Information
 - Submitting On Behalf Of Company
- 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
- Submitter Requests
- Substantiation Summary

Section 8(d) Health & Safety Data Reporting > Contact Information > Technical Contact Information > Submitting On Behalf Of Company

SUBMITTING ON BEHALF OF COMPANY

Fill out the fields below for the manufacturing or processing establishment on whose behalf this submission is being made.

CBI: ☐

Prefix:

First Name:

Middle Initial:

Last Name:

Suffix:

Company Name:

Phone Number: Ext:
(Do not enter any dashes (-) in Phone Number field above.)

Email Address:

Mailing Address 1:
Street address, P.O. box, company name, etc.

Mailing Address 2:
Apartment, suite, etc.

City:

State:

Postal Code:

Country:

[Previous](#) [Next](#)

[Validate](#) [Save](#) [Preview](#)

[CDX Homepage](#) | [MyCDX Homepage](#) | [EPA Homepage](#) | [Terms and Conditions](#) | [Privacy Notice](#) | [CDX Helpdesk: \(888\) 890-1995](#)

EPA Form 9600-032

Chemical Substance Identity of Impurities Screen

OMB No. 2070-0224
Exp 4/30/2029

CSPP Logged in as: John Doe, Primary Support Log Out

Home Submissions Resources

Section 8(d) Health & Safety Data Reporting

Primary Support

- General Submission Information
- Contact Information**
 - Technical Contact Information
 - Submitting On Behalf Of Company
- Chemical Information**
 - Chemical Substance Identity of Impurities**
- Studies**
 - Study Identification
 - Submitter Requests

Section 8(d) Health & Safety Data Reporting > Chemical Information > Chemical Substance Identity of Impurities

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. If impurities do not apply to the reported chemical substance, check the Not Applicable option below.

☐ Not Applicable

SRS	Chemical Identifying Number	Chemical Name (descriptor)	Synonyms	CBI	Actions

[Previous](#) [Next](#)

Validate Save Preview

[CDX Homepage](#) | [MyCDX Homepage](#) | [EPA Homepage](#) | [Terms and Conditions](#) | [Privacy Notice](#) | [CDX Helpdesk: \(888\) 890-1995](#)

Study Identification Screen

The screenshot displays the GSPP (Global Study Plan Portal) interface. At the top, the GSPP logo is on the left, and the user is logged in as 'John Doe, Primary Support' on the right. Below the logo, there are tabs for 'Home', 'Submissions', and 'Resources'. The left sidebar is titled 'Section 8(d) Health & Safety Data Reporting' and contains a tree view with the following items: 'Primary Support', 'General Submission Information', 'Contact Information' (with a sub-item 'Technical Contact Information'), 'Chemical Information' (with a sub-item 'Chemical Substance Identity of Impurities'), 'Studies' (with sub-items 'Study Identification' and 'Submitter Requests'), and 'Submitter Requests'. The main content area is titled 'STUDY IDENTIFICATION' and contains the text 'Please select which types of studies you will be submitting:'. Below this text is a list of study types with checkboxes: 'Not Applicable' (unchecked), 'Full Study Report' (checked), 'Initiated Studies' (checked), 'Ongoing Studies' (checked), 'Robust Summary' (checked), 'Studies Which are Known but without Possession of Copies' (checked), and 'Studies Previously Sent to Federal Agencies without Confidentiality Claims' (checked). At the bottom of the main content area are two buttons: 'Previous' and 'Next'. The 'Next' button is highlighted with a red box. At the very bottom of the page, there is a footer with links: 'CDX Homepage', 'MyCDX Homepage', 'EPA Homepage', 'Terms and Conditions', 'Privacy Notice', and 'CDX Helpdesk: (888) 890-1995'.

Logged in as: John Doe, Primary Support

Home Submissions Resources Log Out

Section 8(d) Health & Safety Data Reporting

Primary Support

General Submission Information

Contact Information

Technical Contact Information

Chemical Information

Chemical Substance Identity of Impurities

Studies

Study Identification

Submitter Requests

Studies > Study Identification

STUDY IDENTIFICATION

Please select which types of studies you will be submitting:

☐ Not Applicable

☒ 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

Validate Save Preview

CDX Homepage | MyCDX Homepage | EPA Homepage | Terms and Conditions | Privacy Notice | CDX Helpdesk: (888) 890-1995

Study Identification Screen for an Amendment

OMB No. 2070-0224
Exp 4/30/2029

The screenshot shows the CSPP (Chemical Safety and Hazard Investigation) web application. The user is logged in as John Doe, Primary Support. The navigation menu on the left includes sections for General Submission Information, Contact Information, Chemical Information, Studies, EPA Request for Further Information, Submitter Requests, and Substantiation Summary. The main content area is titled 'STUDY IDENTIFICATION' and asks the user to select the types of studies they will be submitting. The options are: Not Applicable, 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, EPA Request for Further Information (which includes Underlying Data, Preliminary Reports of Ongoing Studies, and Copies of Studies). The 'Next' button is highlighted with a red box. At the bottom, there are buttons for Validate, Save, and Preview, and a footer with links to CDX Homepage, MyCDX Homepage, EPA Homepage, Terms and Conditions, Privacy Notice, and CDX Helpdesk.

CSPP

Logged in as: John Doe, Primary Support

Home Submissions Resources Log Out

Section 8(d) Health & Safety Data Reporting

Primary Support

General Submission Information

Contact Information

Technical Contact Information

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

Submitter Requests

Substantiation Summary

Studies > Study Identification

STUDY IDENTIFICATION

Please select which types of studies you will be submitting:

☐ Not Applicable

☒ 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

EPA Request for Further Information

☒ Underlying Data

☒ Preliminary Reports of Ongoing Studies

☒ Copies of Studies

Previous Next

Validate Save Preview

CDX Homepage | MyCDX Homepage | EPA Homepage | Terms and Conditions | Privacy Notice | CDX Helpdesk: (888) 890-1995

Full Study Report Screen

CSPP

Logged in as: John Doe, Primary Support

Log Out

Home Submissions Resources

Section 8(d) Health & Safety Data Reporting

Section 8(d) Health & Safety Data Reporting > Studies > Full Study Report

Primary Support

- General Submission Information
- Contact Information
 - Technical Contact Information
 - Submitting On Behalf Of Company
- Chemical Information
 - Chemical Substance Identity of Impurities
- Studies
 - Study Identification
 - Full Study Report
 - Submitter Requests

FULL STUDY REPORT

Click the **Add Document** button to add a new Full Study Report document.

Add Document

File Name	CBI	Actions
-----------	-----	---------

Previous Next

Validate Save Preview

CDX Homepage | MyCDX Homepage | EPA Homepage | Terms and Conditions | Privacy Notice | CDX Helpdesk: (888) 890-1995

Initiated Studies Screen

Section 8(d) Health & Safety Data Reporting

Primary Support

General Submission Information

Contact Information

Technical Contact Information

Submitting On Behalf Of Company

Chemical Information

Chemical Substance Identity of Impurities

Studies

Study Identification

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

Submitter Requests

Section 8(d) Health & Safety Data Reporting > Studies > Initiated Studies

INITIATED STUDIES

Select the Add Document radio button option to attach a Initiated Studies document, or select the Add Studies radio button option to enter all required Initiated Studies information.

Select: ☒ Add Documents ☐ Add Studies

Click the **Add Document** button to add a new Initiated Studies document.

Add Document

File Name	CBI	Actions
-----------	-----	---------

Previous Next

Validate Save Preview

CDX Homepage | MyCDX Homepage | EPA Homepage | Terms and Conditions | Privacy Notice | CDX Helpdesk: (888) 890-1995

EPA Form 9600-032

Section 8(d) Health & Safety Data Reporting

Primary Support

General Submission Information

Contact Information

Technical Contact Information

Submitting On Behalf Of Company

Chemical Information

Chemical Substance Identity of Impurities

Studies

Study Identification

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

Submitter Requests

Section 8(d) Health & Safety Data Reporting > Studies > Initiated Studies

INITIATED STUDIES

Select the Add Document radio button option to attach a Initiated Studies document, or select the Add Studies radio button option to enter all required Initiated Studies information.

Select: ☐ Add Documents ☒ Add Studies

Click the **Add Study** button to add information pertaining to each listed study. For each listed study, provide the title of the study, beginning date of the study, the purpose of the study, types of data to be collected, and the name and address of the laboratory conducting the study.

Expand All | Collapse All

Cancel

Study Title:

Study Start Date:

Study End Date:

Study Purpose:

Data to be Collected:

Create a new Laboratory or select an existing one from the drop-down. Create New

CBI:

Laboratory Name:

Mailing Address 1:

Mailing Address 2:

City:

State:

Postal Code:

Country:

Save

Add Study

Previous Next

Validate Save Preview

CDX Homepage | MyCDX Homepage | EPA Homepage | Terms and Conditions | Privacy Notice | CDX Helpdesk: (888) 890-1995

Ongoing Studies Screen

This screenshot shows the 'Ongoing Studies' screen in the GSPP system. The user is logged in as 'John Doe, Primary Support'. The left sidebar contains a navigation menu with sections: 'Section 8(d) Health & Safety Data Reporting', 'Primary Support', 'Contact Information', 'Chemical Information', and 'Studies'. The 'Studies' section is expanded, showing options like 'Study Identification', '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', and 'Submitter Requests'. The main content area is titled 'ONGOING STUDIES' and includes instructions: 'Select the Add Document radio button option to attach a Ongoing Studies document, or select the Add Studies radio button option to enter all required Ongoing Studies information.' Below this, there are two radio buttons: 'Add Documents' (selected) and 'Add Studies'. A red box highlights the 'Add Document' button. Below the buttons is a table with columns 'File Name', 'CBI', and 'Actions'. At the bottom of the main content area are 'Previous' and 'Next' buttons. The footer contains links to 'CDX Homepage', 'MyCDX Homepage', 'EPA Homepage', 'Terms and Conditions', 'Privacy Notice', and 'CDX Helpdesk: (888) 890-1995'. A blue bar at the bottom has 'Validate', 'Save', and 'Preview' buttons.

This screenshot shows the 'Add Studies' form in the GSPP system. The user is logged in as 'John Doe, Primary Support'. The left sidebar is the same as the previous screenshot. The main content area is titled 'ONGOING STUDIES' and includes instructions: 'Select the Add Document radio button option to attach a Ongoing Studies document, or select the Add Studies radio button option to enter all required Ongoing Studies information.' Below this, there are two radio buttons: 'Add Documents' and 'Add Studies' (selected). A red box highlights the 'Add Studies' button. Below the buttons is a form with the following fields: 'Study Title', 'Study Start Date', 'Study End Date', 'Study Purpose', 'Data to be Collected', 'Create a new Laboratory or select an existing one from the drop-down.', 'CBI', 'Laboratory Name', 'Mailing Address 1', 'Mailing Address 2', 'City', 'State', 'Postal Code', and 'Country'. A red box highlights the 'Save' button. Below the form are 'Previous' and 'Next' buttons. The footer contains links to 'CDX Homepage', 'MyCDX Homepage', 'EPA Homepage', 'Terms and Conditions', 'Privacy Notice', and 'CDX Helpdesk: (888) 890-1995'. A blue bar at the bottom has 'Validate', 'Save', and 'Preview' buttons.

Robust Summary Add Document Screen

The screenshot shows the CSPP (Chemical Safety and Hazard Investigation) system interface. At the top, the user is logged in as 'John Doe, Primary Support'. The navigation bar includes 'Home', 'Submissions', and 'Resources'. The left sidebar shows the 'Section 8(d) Health & Safety Data Reporting' menu with various options like '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', 'Submitter Requests', and 'Substantiation Summary'. The main content area is titled 'ROBUST SUMMARY' and contains the instruction 'Click the **Add Document** button to add a new Robust Summary document'. The 'Add Document' button is highlighted with a red box. Below this instruction is a table with two columns: 'File Name' and 'Actions'. At the bottom of the main content area are 'Previous' and 'Next' buttons. The footer bar contains 'Validate', 'Save', and 'Preview' buttons, and the bottom-most bar lists links: 'CDX Homepage', 'MyCDX Homepage', 'EPA Homepage', 'Terms and Conditions', 'Privacy Notice', and 'CDX Helpdesk (888) 890-1995'.

CSPP

Logged in as: John Doe, Primary Support

Home Submissions Resources Log Out

Section 8(d) Health & Safety Data Reporting > Studies > Robust Summary

ROBUST SUMMARY

Click the **Add Document** button to add a new Robust Summary document

Add Document

File Name	Actions
-----------	---------

Previous Next

Validate Save Preview

CDX Homepage | MyCDX Homepage | EPA Homepage | Terms and Conditions | Privacy Notice | CDX Helpdesk (888) 890-1995

Studies which are Known but Without Possession of Copies Screens

Section 8(d) Health & Safety Data Reporting

Logged in as: John Doe, Primary Support

Home Submissions Resources Log Out

Section 8(d) Health & Safety Data Reporting > Studies > Studies Which are Known but without Possession of Copies

STUDIES WHICH ARE KNOWN BUT WITHOUT POSSESSION OF COPIES

Select the Add Document radio button option to attach a Studies Which are Known but without Possession of Copies document, or select the Add Studies radio button option to enter all required Studies Which are Known but without Possession of Copies information.

Select: ☒ Add Documents ☐ Add Studies

Click the **Add Document** button to add a new Studies Which are Known but without Possession of Copies document.

Add Document

File Name	CBI	Actions
-----------	-----	---------

Previous Next

Validate Save Preview

CDX Homepage | MyCDX Homepage | EPA Homepage | Terms and Conditions | Privacy Notice | CDX Helpdesk: (888) 890-1995

Section 8(d) Health & Safety Data Reporting

Logged in as: John Doe, Primary Support

Home Submissions Resources Log Out

Section 8(d) Health & Safety Data Reporting > Studies > Studies Which are Known but without Possession of Copies

STUDIES WHICH ARE KNOWN BUT WITHOUT POSSESSION OF COPIES

Select the Add Document radio button option to attach a Studies Which are Known but without Possession of Copies document, or select the Add Studies radio button option to enter all required Studies Which are Known but without Possession of Copies information.

Select: ☐ Add Documents ☒ Add Studies

Click the **Add Study** button to add information pertaining to each listed study. For each listed study, provide the title of the study, and the name and address of the contact conducting the study.

Expand All | Collapse All

Cancel

Study Title:

Create a new Contact or select an existing one from the drop-down.

CBI: ☐

Prefix:

First Name:

Last Name:

Suffix:

Phone Number: Ext:

(Do not enter any dashes (-) in Phone Number field above.)

Email Address:

Mailing Address 1:

Street address, P.O. box, company name, etc.

Mailing Address 2:

Apartment, suite, etc.

City:

State:

Postal Code:

Country:

Save

Add Study

Previous Next

Validate Save Preview

CDX Homepage | MyCDX Homepage | EPA Homepage | Terms and Conditions | Privacy Notice | CDX Helpdesk: (888) 890-1995

Studies Previously Sent to Federal Agencies without Confidentiality Claims Screens

GSPP

Logged in as: John Doe, Primary Support

Home Submissions Resources Log Out

Section 8(d) Health & Safety Data Reporting

Primary Support

- General Submission Information
- Contact Information
 - Technical Contact Information
 - Submitting On Behalf Of Company
- Chemical Information
 - Chemical Substance Identity of Impurities
- Studies
 - Study Identification
 - 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
- Submitter Requests

Section 8(d) Health & Safety Data Reporting > Studies > Studies Previously Sent to Federal Agencies without Confidentiality Claims

STUDIES PREVIOUSLY SENT TO FEDERAL AGENCIES WITHOUT CONFIDENTIALITY CLAIMS

Select the Add Document radio button option to attach a Studies Previously Sent to Federal Agencies without Confidentiality Claims document, or select the Add Studies radio button option to enter all required Studies Previously Sent to Federal Agencies without Confidentiality Claims information.

Select: ☒ Add Documents ☐ Add Studies

Click the **Add Document** button to add a new Studies Previously Sent to Federal Agencies without Confidentiality Claims document.

Add Document

File Name	CBI	Actions
-----------	-----	---------

Previous Next

Validate Save Preview

CDX Homepage | MyCDX Homepage | EPA Homepage | Terms and Conditions | Privacy Notice | CDX Helpdesk: (888) 890-1995

EPA Form 9600-032

GSPP

Logged in as: John Doe, Primary Support

Home Submissions Resources Log Out

Section 8(d) Health & Safety Data Reporting

Primary Support

- General Submission Information
- Contact Information
 - Technical Contact Information
 - Submitting On Behalf Of Company
- Chemical Information
 - Chemical Substance Identity of Impurities
- Studies
 - Study Identification
 - 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
- Submitter Requests

Section 8(d) Health & Safety Data Reporting > Studies > Studies Previously Sent to Federal Agencies without Confidentiality Claims

STUDIES PREVIOUSLY SENT TO FEDERAL AGENCIES WITHOUT CONFIDENTIALITY CLAIMS

Select the Add Document radio button option to attach a Studies Previously Sent to Federal Agencies without Confidentiality Claims document, or select the Add Studies radio button option to enter all required Studies Previously Sent to Federal Agencies without Confidentiality Claims information.

Select: ☐ Add Documents ☒ Add Studies

Click the **Add Study** button to add information pertaining to each listed study. For each listed study, provide the title of the study, submission date of the study, the agency, and the name and address of the agency contact conducting the study.

Expand All | Collapse All

Cancel

Study Title:

Study Submission Date:

Agency:

Create a new Contact or select an existing one from the drop-down.

CBI: ☐

Prefix:

First Name:

Last Name:

Suffix:

Phone Number: Ext:

(Do not enter any dashes (-) in Phone Number field above.)

Email Address:

Mailing Address 1:

Mailing Address 2:

City:

State:

Postal Code:

Country:

Save

Add Study

Previous **Next**

Validate Save Preview

Underlying Data Screen

GSPP

Logged in as: John Doe, Primary Support

Log Out

Home Submissions Resources

Section 8(d) Health & Safety Data Reporting

Primary Support

General Submission Information

Contact Information

Technical Contact Information

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

Section 8(d) Health & Safety Data Reporting > Studies > EPA Request for Further Information > Underlying Data

UNDERLYING DATA

Click the **Add Document** button to add a new Underlying Data document.

Add Document

File Name	CBI	Actions
-----------	-----	---------

Previous Next

Validate Save Preview

CDX Homepage | MyCDX Homepage | EPA Homepage | Terms and Conditions | Privacy Notice | CDX Helpdesk: (888) 890-1995

Preliminary Reports of Ongoing Studies Screens

This screenshot shows the 'Preliminary Reports of Ongoing Studies' screen in the CDX system. The user is logged in as John Doe, Primary Support. The left sidebar contains a navigation menu with sections like 'Section 8(d) Health & Safety Data Reporting', 'Primary Support', 'Contact Information', 'Chemical Information', 'Studies', and 'EPA Request for Further Information'. The main content area has a breadcrumb trail: 'Section 8(d) Health & Safety Data Reporting > Studies > Preliminary Reports of Ongoing Studies'. Below this, the title 'PRELIMINARY REPORTS OF ONGOING STUDIES' is displayed. Instructions state: 'Select the Add Document radio button option to attach a Preliminary Reports of Ongoing Studies document, or select the Add Studies radio button option to enter all required Preliminary Reports of Ongoing Studies information.' The 'Select' section shows two radio buttons: 'Add Documents' (selected) and 'Add Studies'. Below this, a message says 'Click the Add Document button to add a new Preliminary Reports of Ongoing Studies document.' and an 'Add Document' button is highlighted with a red box. A table with columns 'File Name', 'CBI', and 'Actions' is present, but it is empty. At the bottom of the main area are 'Previous' and 'Next' buttons. The footer contains a blue bar with 'Validate', 'Save', and 'Preview' icons, and a grey bar with links to 'CDX Homepage', 'MyCDX Homepage', 'EPA Homepage', 'Terms and Conditions', 'Privacy Notice', and 'CDX Helpdesk: (888) 890-1995'.

EPA Form 9600-032

This screenshot shows the 'Add Studies' form within the 'Preliminary Reports of Ongoing Studies' screen. The user is logged in as John Doe, Primary Support. The left sidebar is the same as the previous screenshot. The main content area has a breadcrumb trail: 'Section 8(d) Health & Safety Data Reporting > Studies > EPA Request for Further Information > Preliminary Reports of Ongoing Studies'. Below this, the title 'PRELIMINARY REPORTS OF ONGOING STUDIES' is displayed. Instructions state: 'Select the Add Document radio button option to attach a Preliminary Reports of Ongoing Studies document, or select the Add Studies radio button option to enter all required Preliminary Reports of Ongoing Studies information.' The 'Select' section shows two radio buttons: 'Add Documents' and 'Add Studies' (selected). Below this, a message says 'Click the Add Study button to add information pertaining to each listed study. For each listed study, provide the title of the study, beginning date of the study, the purpose of the study, types of data to be collected, and the name and address of the laboratory conducting the study.' A modal form is open with a 'Cancel' button and a red 'X' icon. The modal contains the following fields: 'Study Title:', 'Study Start Date:', 'Study End Date:', 'Study Purpose:', 'Data to be Collected:', 'Create a new Laboratory or select an existing one from the drop-down. Create New', 'CBI:', 'Laboratory Name:', 'Mailing Address 1:', 'Mailing Address 2:', 'City:', 'State:', 'Postal Code:', and 'Country:'. A 'Save' button is highlighted with a red box. Below the modal, an 'Add Study' button is visible. At the bottom of the main area are 'Previous' and 'Next' buttons, with the 'Next' button highlighted with a red box. The footer contains a blue bar with 'Validate', 'Save', and 'Preview' icons, and a grey bar with links to 'CDX Homepage', 'MyCDX Homepage', 'EPA Homepage', 'Terms and Conditions', 'Privacy Notice', and 'CDX Helpdesk: (888) 890-1995'.

Copies of Studies Screens

Section 8(d) Health & Safety Data Reporting

Primary Support

General Submission Information

Contact Information

Technical Contact Information

Submitting On Behalf Of Company

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

Completed Studies

Section 8(d) Health & Safety Data Reporting > Studies > Copies of Studies

COPIES OF STUDIES

Select the Add Document radio button option to attach a Copies of Studies document, or select the Add Studies radio button option to enter all required Copies of Studies information.

Select: ☒ Add Documents ☐ Add Studies

Click the **Add Document** button to add a new Copies of Studies document.

Add Document

File Name	CBI	Actions
-----------	-----	---------

Previous Next

Validate Save Preview

CDX Homepage | MyCDX Homepage | EPA Homepage | Terms and Conditions | Privacy Notice | CDX Helpdesk: (888) 890-1995

EPA Form 9600-032

Section 8(d) Health & Safety Data Reporting

Primary Support

General Submission Information

Contact Information

Technical Contact Information

Submitting On Behalf Of Company

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

Submitter Requests

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

COPIES OF STUDIES

Select the Add Document radio button option to attach a Copies of Studies document, or select the Add Studies radio button option to enter all required Copies of Studies information.

Select: ☐ Add Documents ☒ Add Studies

Click the **Add Study** button to add information pertaining to each listed study. For each listed study, provide the title of the study, and the name and address of the contact conducting the study.

Expand All | Collapse All

Cancel

Study Title:

Create a new Contact or select an existing one from the drop-down. Create New

CBI:

Prefix:

First Name:

Last Name:

Suffix:

Phone Number:

Ext:

(Do not enter any dashes (-) in Phone Number field above.)

Email Address:

Mailing Address 1:

Mailing Address 2:

City:

State:

Postal Code:

Country:

Save

Add Study

Previous Next

Validate Save Preview

CDX Homepage | MyCDX Homepage | EPA Homepage | Terms and Conditions | Privacy Notice | CDX Helpdesk: (888) 890-1995

Submitter Requests Screen

CSPP

Logged in as: John Doe, Primary Support

Home Submissions Resources Log Out

Section 8(d) Health & Safety Data Reporting > Submitter Requests

SUBMITTER REQUESTS

Click the **Add Document** button to add a new Submitter Request document.

Add Document

File Name	Request Type	Actions
-----------	--------------	---------

Previous

Validate Save Preview

CDX Homepage | MyCDX Homepage | EPA Homepage | Terms and Conditions | Privacy Notice | CDX Helpdesk: (888) 890-1995

Substantiation Summary Screen

CSPP

Logged in as: John Doe, Primary Support

Home Submissions Resources Log Out

Section 8(d) Health & Safety Data Reporting

Section 8(d) Health & Safety Data Reporting > Substantiation Summary

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.

Chemical Identifying Number	Chemical Name	Substantiation Status
11111	Ricinoleic acid homopolymer	Pending

File Name	Study Classification	Substantiation Status
-----------	----------------------	-----------------------

Previous

Validate Save Preview

CDX Homepage | MyCDX Homepage | EPA Homepage | Terms and Conditions | Privacy Notice | CDX Helpdesk: (888) 890-1995

Submission of Studies Substantiation on Screen



Logged in as: John Doe, Primary Support

Log Out

HomeSubmissionsResources

Section 8(d) Health & Safety Data Reporting

Primary Support

General Submission Information

Contact Information

Technical Contact Information

Submitting On Behalf Of Company

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

Submitter Requests

Substantiation Summary

Section 8(d) Health & Safety Data Reporting > Substantiation Summary > 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

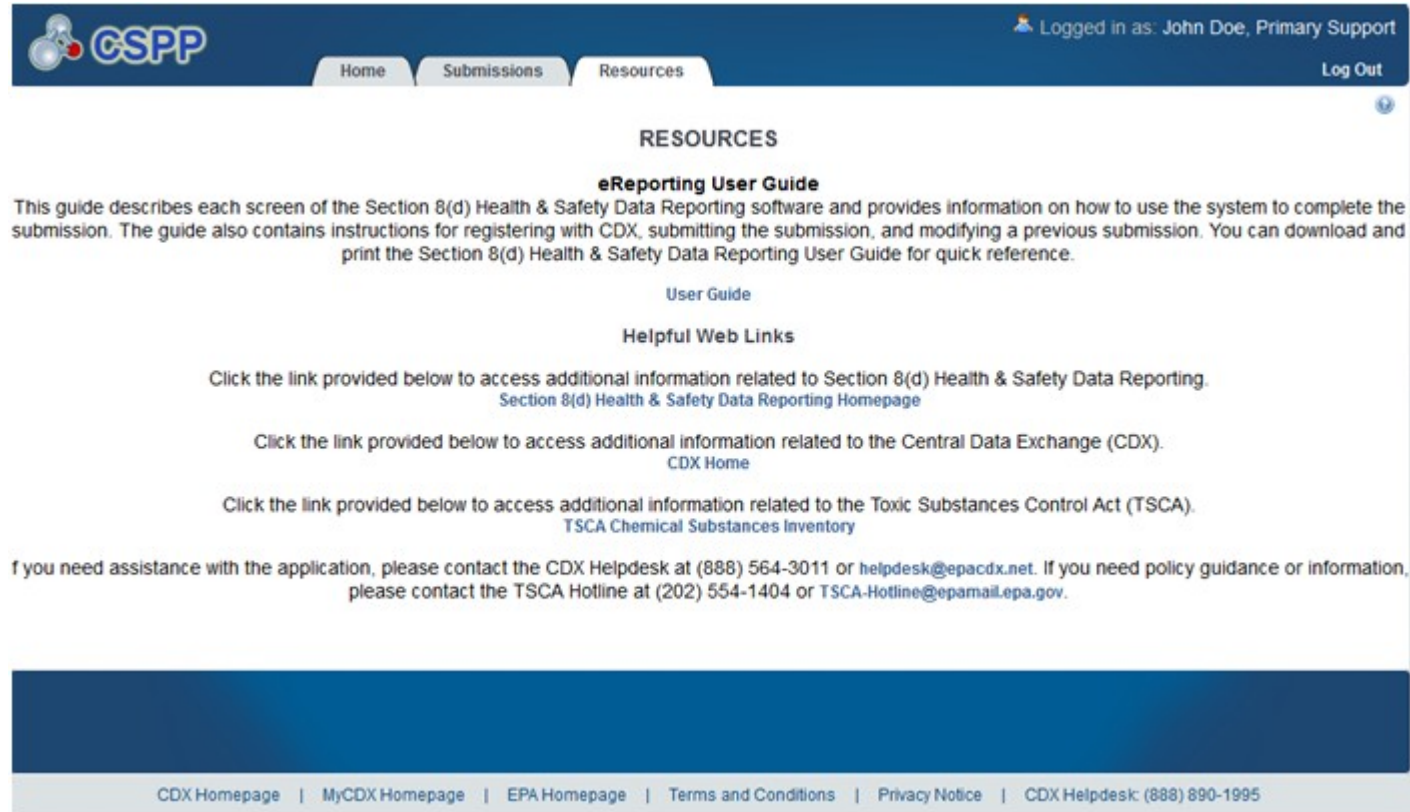
Validate

Save

Preview

CDX Homepage | MyCDX Homepage | EPA Homepage | Terms and Conditions | Privacy Notice | CDX Helpdesk: (888) 890-1995

Resources Screen



The screenshot shows the 'Resources' page of the CSPP (Chemical Safety Portal) application. The header is dark blue with the CSPP logo on the left, navigation tabs for 'Home', 'Submissions', and 'Resources' in the center, and a user login status 'Logged in as: John Doe, Primary Support' with a 'Log Out' link on the right. The main content area is white and titled 'RESOURCES'. It features a section for the 'eReporting User Guide' with a descriptive paragraph and a 'User Guide' link. Below this is a 'Helpful Web Links' section with three links: 'Section 8(d) Health & Safety Data Reporting Homepage', 'CDX Home', and 'TSCA Chemical Substances Inventory', each preceded by a brief instruction. At the bottom, a footer bar contains links to 'CDX Homepage', 'MyCDX Homepage', 'EPA Homepage', 'Terms and Conditions', 'Privacy Notice', and 'CDX Helpdesk: (888) 890-1995'.

CSPP

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

RESOURCES

eReporting User Guide

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

[User Guide](#)

Helpful Web Links

Click the link provided below to access additional information related to Section 8(d) Health & Safety Data Reporting.
[Section 8\(d\) Health & Safety Data Reporting Homepage](#)

Click the link provided below to access additional information related to the Central Data Exchange (CDX).
[CDX Home](#)

Click the link provided below to access additional information related to the Toxic Substances Control Act (TSCA).
[TSCA Chemical Substances Inventory](#)

If you need assistance with the application, please contact the CDX Helpdesk at (888) 564-3011 or helpdesk@epacdx.net. If you need policy guidance or information, please contact the TSCA Hotline at (202) 554-1404 or TSCA-Hotline@epamail.epa.gov.

[CDX Homepage](#) | [MyCDX Homepage](#) | [EPA Homepage](#) | [Terms and Conditions](#) | [Privacy Notice](#) | [CDX Helpdesk: \(888\) 890-1995](#)