United States Environmental Protection Agency

Section 4, 8(d), and 8(a) Webinar

December 12, 2013.

Office of Chemical Safety and Pollution Prevention

Agenda

- Speaker: Veronica Shortt, Information Management Division
- Welcome and Overview
- Section 4 highlights
  - Browser-based Application
  - Security
  - Roles
  - Industry Beta Testing
- Demo of CDX Login and Section 4 Web Tool
- Resources
- Questions and Answers

Highlights & Security

- Browser-based Application
- Submission of Section 4 facilitated via EPA's Central Data Exchange (CDX)
  - Enables companies to electronically submit data
  - Provides secure exchange of confidential business information (CBI) data
  - Improves security using digital encryption
    - All data is encrypted in transit to Federal Standards
    - Sensitive data is encrypted at rest to Federal Standards
    - All sensitive data remains encrypted at rest until it is behind at least 2-3 Firewalls

Roles & Responsibilities

- Primary Authorized Official (AO)
  - User who has the ability to authorize and sign a submission
  - Start a form
  - Edit/delete information on a form
  - Submit a form
  - Retrieve Copy of Record
  - Unlock a form for editing
  - Assign Supports
- Primary Support
  - Enter and edit information for in progress and unlocked forms

Demo of CDX Login and Section 4 Tool

Primary Authorized Official

Central Data Exchange

Contact Us

Log in to CDX

User ID

Password

Log In | Register with CDX

Forgot your password?

Warning Notice and Privacy Policy

Welcome

Welcome to the Environmental Protection Agency (EPA) Central Data Exchange (CDX) – the Agency's electronic reporting site. The Central Data Exchange concept has been defined as a central point which supplements EPA reporting systems by performing new and existing functions for receiving legally acceptable data in various formats, including consolidated and integrated data.

Warning Notice and Privacy Policy

Warning Notice

EPA's Central Data Exchange Registration procedure is part of a United States Environmental Protection Agency (EPA) computer system, which is for authorized use only. Unauthorized access or use of this computer system may subject violators to criminal, civil, and/or administrative action. All information on this computer system may be monitored, recorded, read, copied, and disclosed by and to authorized personnel for official purposes, including law enforcement. Access or use of this computer system by any person, whether authorized or unauthorized, constitutes consent to these terms.

Privacy Statement

EPA will use the personal identifying information which you provide for the expressed purpose of registration to the Central Data Exchange site and for updating and correcting information in internal EPA databases as necessary. The Agency will not make this information available for other purposes unless required by law. EPA does not sell or otherwise transfer personal information to an outside third party. [Federal Register: March 18, 2002 (Volume 67, Number 52)][Page 12010-12013].

Central Data Exchange

Contact Us

Last Login: 11/19/2013 1:25:52 PM

Services

Manage Your Program Services

Status	Program Service Name	Role(s)
	CSPP: Submissions for Chemical Safety and Pesticide Programs	Primary Authorized Official

Add Program Service

News and Updates

No news/updates.

CHEMICAL INFORMATION SUBMISSION SYSTEM

TSCA Section 4

OK

The software includes embedded help files and a downloadable user manual to guide you through the Section 4 submission process.

The Toxic Substances Control Act gives EPA authority to issue data development regulations that require manufacturers and processors of existing chemicals to test their chemicals for health and environmental effects. EPA has the broad authority under the law to issue:

Information collection regulations that require the submission of health and safety studies which are known or available to those who manufacture, process, or distribute in commerce specified chemicals; and regulations designed to gather information from manufacturers and processor about production/import volumes, chemical uses and methods of disposal, and the extent to which people and the environment are exposed.

TSCA also requires EPA to develop regulations that establish import/export requirements for chemicals which are subject to certain requirements under TSCA.

Paperwork Reduction Act Notice

The information collection requirements contained in this final rule have been submitted for OMB approval under PRA, 44 U.S.C. 3501 et seq. The ICR document prepared by EPA, identified under EPA ICR No. 2412.01 and OMB control number 2070-0004, is available in the docket for the final rule. The ICR addresses the incremental changes to the currently approved ICR documents that cover the existing reporting and record keeping programs that are approved under OMB control numbers 2070-0004, 2070-0033, and 2070-0054. 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 amended information collection activities contained in this final rule are designed to assist the Agency in meeting its responsibility under TSCA to receive, process, and review reports, data, and other information. As such, responses to the collection of information covered by this ICR would still be mandatory, but with the final rule, respondents would be required to use the CISS reporting tool.

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.

HOME

Submissions

Create, modify, or delete a submission by clicking the Submissions tab.

User Management

Manage the access rights of Supports for each Section 4 submission. For every Support the Authorized Official may grant him/her the ability to edit (but not unlock, create, delete, or submit) the submission.

Resources

A helpful guide that describes the Section 4 system and provides useful links for further usability instruction.

Authorized Official

An Authorized Official has the ability to create, delete, amend, unlock and submit all Section 4 submissions electronically to EPA. The Authorized Official also has the ability to assign Supports to individual submissions.

USER MANAGEMENT

The Authorized Official is responsible for restricting a Support's access to select submissions by assigning or unassigning them to each submission. The Support can access and edit only those submissions for which the Authorized Official has granted access. Select a submission from the drop-down menu, and assign a Support to the submission by highlighting the individual and clicking the **add** link. To unassign a Support, highlight the individual and click the **remove** link. To highlight and assign or unassign multiple Supports, hold down the **Ctrl** or **Shift** keys on the keyboard and click each Support before moving. You must click the **Save** button after each assignment.

Section 4 Submission:

Submission Information

CFR/FRN:
Submission Type:
Last Modified:

Assign Users

Unassigned		Assigned
	add >> << remove	

HOME

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SECTION 4 TEST RULES, ECAS, MOUS, AND 40 CFR 766

- If starting a Section 4 Test Rule, ECA, MOU, or a 40 CFR 766 submission, select the appropriate submission type from the drop-down menu and click the 'Start New Submission' button.
- To edit an **In Progress** submission, click the submission link in the **Submission Alias** column in the table below.
- To access and edit a submission previously **Submitted** through CDX, unlock the submission by clicking the lock icon ( ) and enter your passphrase originally associated with the selected submission. All additional changes made to a submission will be submitted as an amendment.
- Click the green arrow icon ( ) to download a copy of record for a completed submission. It may take up to 15 minutes for the copy of record to become available.
- You may delete any submission that has not yet been submitted by clicking the delete icon ( ).

8 items found. Page 1 of 1 Items Per Page: 25

Submission Alias	CFR/FRN	Status	Modify Date	Submission Date	Copy of Record	Action
CFR766-20131121-12:41:40 EST	40 CFR 766 Dibenzodioxins / Dibenzofurans	In Progress	11/27/2013			
ECA-20131121-12:41:03 EST		In Progress	11/27/2013			
ECA-20131129-15:25:33 EST	70 FR 39630	Submitted	11/29/2013	11/29/2013		
ECA-20131204-15:12:41 EST		In Progress	12/04/2013			
MOU-20131121-12:41:27 EST	MOUForm	In Progress	11/27/2013			
TestRules-20131121-12:41:14 EST	53 FR 22300	In Progress	12/04/2013	11/26/2013		
TestRules-20131127-10:17:23 EST	69 FR 22402	In Progress	11/27/2013			
test amendment	53 FR 22300	Submitted	11/29/2013	11/29/2013		

Export options: CSV | Excel | XML | PDF

Select the submission type and then click **Start New Submission**

Submission Type:

- Test Rules
- ECA
- MOU
- 40 CFR 766 Dibenzodioxins/Dibenzofurans

Section 4: Test Rules/Letter of Intent

CREATE PASSPHRASE

Please create a passphrase that is at least 8 characters in length. To better protect your form, your passphrase should contain a combination of letters numbers. Your passphrase may include spaces, but should not contain special characters (for example, + and *).

As an Authorized Official, you are responsible for remembering your passphrase and distributing it to only authorized individuals. 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 only authorized Supports.

Note: If you lose or forget your passphrase, you will not be able to access your Section 4 Submission to print, submit, or make changes. You will need to complete a new Section 4 Submission and create a new passphrase for the submission. For security reasons, the system administrator will not have access to your passphrase and will not be able to retrieve it or reset it.

New Passphrase:
Confirm New Passphrase:

Cancel Next

Section 4 Test Rules
Primary Authorized Official

- Federal Register Notice
  - Submission Information Type
- Contact Information
  - Technical Contact Information
- Letter of Intent
  - Chemical Identification and Test Rules
  - Sponsoring Firms
  - Additional Information and Submitter Requests
- Optional Substantiation
  - Document Management

Section 4 Test Rules > Federal Register Notice > Submission Information Type

SUBMISSION INFORMATION TYPE

Please select, or start entering, a FRN in the drop-down menu below

Federal Register Notice:

Please enter a submission alias and choose the submission information type(s) here.

Submission Alias: TestRules-20130711-1(

- [ ] Study Plan
- [ ] Results

Next

Section 4 Test Rules
Primary Authorized Official

- 53 FR 22300
  - Submission Information Type
  - Contact Information
    - Technical Contact Information
  - Letter of Intent
    - Chemical Identification and Test Rules
    - Sponsoring Firms
    - Additional Information and Submitter Requests
  - Optional Substantiation
    - Document Management

Section 4 Test Rules > 53 FR 22300 > 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 or consortium by selecting the appropriate radio button. If submitting on behalf of is not applicable, select the N/A radio button. Click the 'Copy CDX Registration' button to copy your information from CDX Registration.

- ( ) N/A
- ( ) This is a submission on behalf of a consortium
- ( ) This is a submission on behalf of another company

Click here to copy your information from CDX Registration: **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:

Previous **Next**

Section 4 Test Rules
Primary Authorized Official

- 53 FR 22300
  - Submission Information Type
  - Contact Information
    - Technical Contact Information
    - Submitting on Behalf of Consortium
  - Letter of Intent
    - Chemical Identification and Test Rules
    - Sponsoring Firms
    - Additional Information and Submitter Requests
  - Optional Substantiation
    - Document Management

Section 4 Test Rules > 53 FR 22300 > Letter of Intent > Chemical Identification And Test Rules

CHEMICAL IDENTIFICATION AND TEST RULES

Click the Assign Chemicals button to identify the chemical substance(s) the sponsor(s) intends to use in each of the tests. To save, delete, or add a chemical test, please use the icons provided.

Assign Chemicals

For the chemical substance(s) listed below, identify the testing requirement(s) to be performed on the specified substance.

Previous Next

CSPP

CHEMICAL MANAGEMENT

Select each chemical substance for which the sponsor(s) intends to use in each of the tests. When all appropriate chemical substances have been assigned, click the Save button.

Assign	Chemical Name	CASRN
☐	Un/check All Chemicals	
☐	2-Fluoroacetamide	640-19-7
☐	4-Bromobenzyl Cyanide	16532-79-9
☐	Bis(2-chloroisopropyl)ether	108-60-1
☐	Endrin	72-20-8
☐	Maleic Hydrazide	123-33-1
☐	Methanethiol	74-93-1
☐	Pentachlorobenzene	608-93-5
☐	Trichloromethanethiol	75-70-7

Save Cancel

Section 4 Test Rules
Primary Authorized Official

- 71 FR 13708
  - Submission Information Type
  - Contact Information
    - Technical Contact Information
  - Letter of Intent
    - Chemical Identification and Test Rules
    - Sponsoring Firms
    - Additional Information and Submitter Requests
  - 1324-76-1
    - Remove
  - 2941-64-2
    - Remove
  - Optional Substantiation
    - Document Management

Section 4 Test Rules > 71 FR 13708 > Letter of Intent > Chemical Identification and Test Rules

CHEMICAL IDENTIFICATION AND TEST RULES

Click the Assign Chemicals button to identify the chemical substance(s) the sponsor(s) intends to use in each of the tests. To save, delete, or add a chemical test, please use the icons provided.

Assign Chemicals

For the chemical substance(s) listed below, identify the testing requirement(s) to be performed on the specified substance.

Benzenesulfonic Acid, [[4-[[4-(phenylamino)phenyl][4-(phenylimino)-2,5-cyclohexadien-1-ylidene]methyl]phenyl]amino]- (CASRN: 1324-76-1)

CBI

Chemical Test	Actions
A	

Carbonochloridothioic Acid, S-ethyl ester (CASRN: 2941-64-2)

CBI

Chemical Test	Actions
B	

Previous Next

Section 4 Test Rules
Primary Authorized Official

- 53 FR 22300
  - Submission Information Type
  - Contact Information
    - Technical Contact Information
    - Submitting on Behalf of Consortium
  - Letter of Intent
    - Chemical Identification and Test Rules
    - Sponsoring Firms
    - Additional Information and Submitter Requests
  - 640-19-7
    - Remove
  - 16532-79-9
    - Remove
  - Optional Substantiation
    - Document Management

Section 4 Test Rules > 53 FR 22300 > Letter of Intent > Sponsoring Firms

SPONSORING FIRMS

Click the Add Sponsoring Firm button to add each Sponsoring Firm and fill out the information below.

Expand All | Collapse All

Firm - Cancel

CBI:
Firm Name:
Phone Number: Ext:
Mailing Address 1:
Mailing Address 2:
City:
State:
Postal Code:
Country:

Save

Click the **Add Sponsoring Firm** button to add a new sponsoring firm. Add Sponsoring Firm

Previous Next

Section 4 Test Rules
Primary Authorized Official

- 53 FR 22300
  - Submission Information Type
  - Contact Information
    - Technical Contact Information
  - Letter of Intent
    - Chemical Identification and Test Rules
    - Sponsoring Firms
    - Additional Information and Submitter Requests
  - 640-19-7
    - Remove
  - 16532-79-9
    - Remove
  - Optional Substantiation
    - Document Management

ADDITIONAL INFORMATION AND SUBMITTER REQUESTS

Select the information or request type from the dropdown and click the Add Information button to complete the required information.

Note: For security purposes, documents are only saved when there are no validation errors. If there are any validation errors, all documents must be reattached.

Information/Request Type: Exemption Application

Add Information

File Name	Document Type	Attachment Date	Action

Nothing found to display.

Previous

Section 4: Submit Letter of Intent

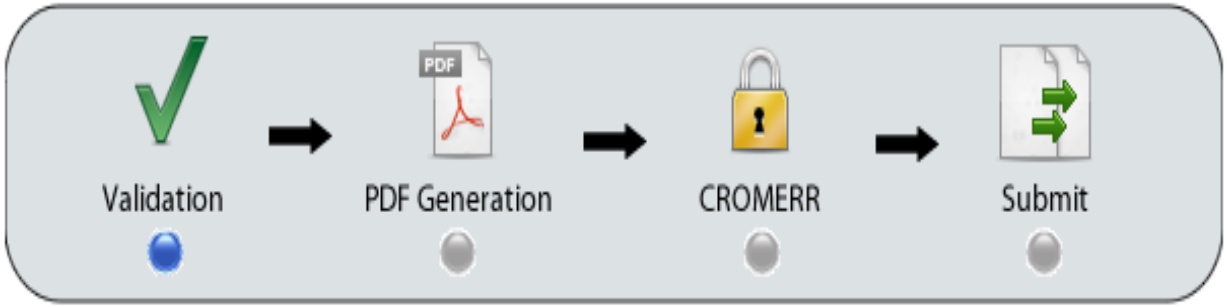
SUBMITTING OFFICIAL INFORMATION

The information below has been prepopulated from CDX registration. If the information listed is incorrect please make the appropriate edits.

CBI:	☐
Prefix:	Mr
First Name:	John
Middle Initial:	
Last Name:	Doe
Job Title:	
Company Name:	TEST Company
Phone Number:	5551231234
Email Address:	johndoe@gmail.com
Mailing Address 1:	123 Test Drive
Mailing Address 2:	534-I
City:	FAIRFAX
State:	VA
Postal Code:	22033

Cancel Next

SUBMISSION PROCESS: VALIDATION



Running validation on your data

Processing. Please wait...

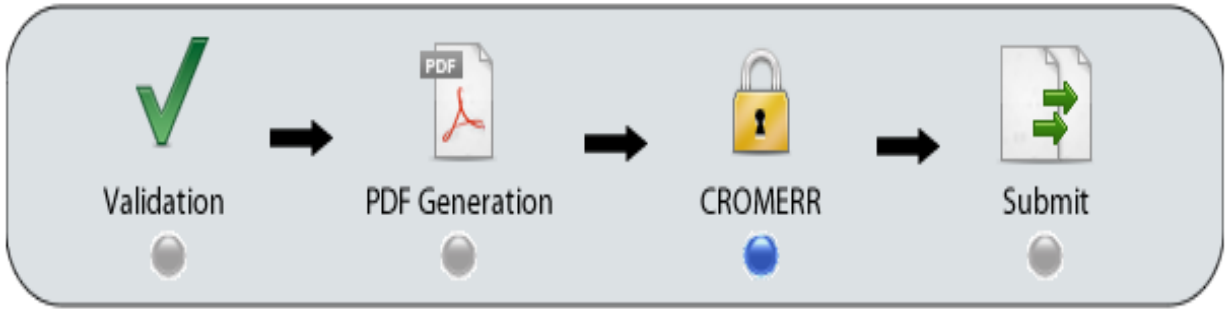
(Please disable any pop-up blockers within your internet browser settings to allow for the validation pop-up to be displayed if validation errors are present.)

SUBMISSION PROCESS: PDF GENERATION

Your PDF preview transaction was successful!

View PDF | Continue

CROSS-MEDIA ELECTRONIC REPORTING REGULATION (CROMERR) CERTIFICATION

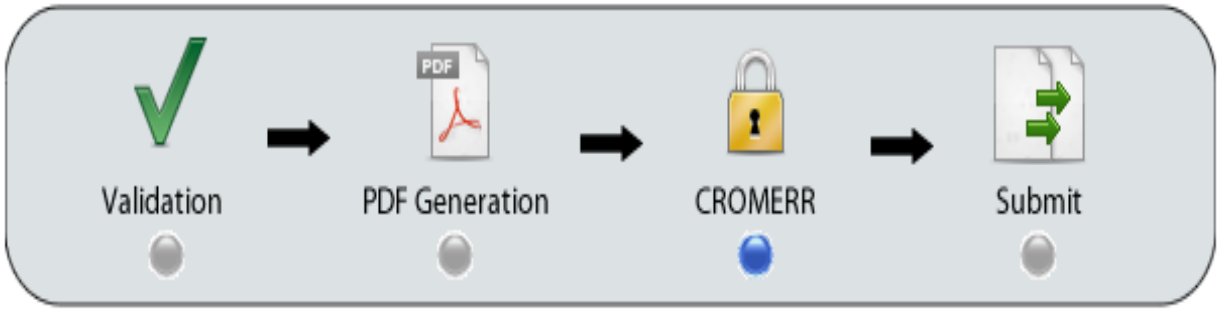


I certify, under penalty of law, that this document and all attachments were prepared under my direction of supervision in accordance with a system designed to assure that qualified personnel properly gather and evaluate the information submitted. Based on my inquiry of the person or persons who manage the system, or those persons directly responsible for gathering the information, the information submitted is, to the best of my knowledge and belief, true, accurate, and complete. I am aware that there are significant penalties for submitting false information, including the possibility of fines and imprisonment for knowing violations.

Cancel

I Certify

CROSS-MEDIA ELECTRONIC REPORTING REGULATION (CROMERR) LOGIN

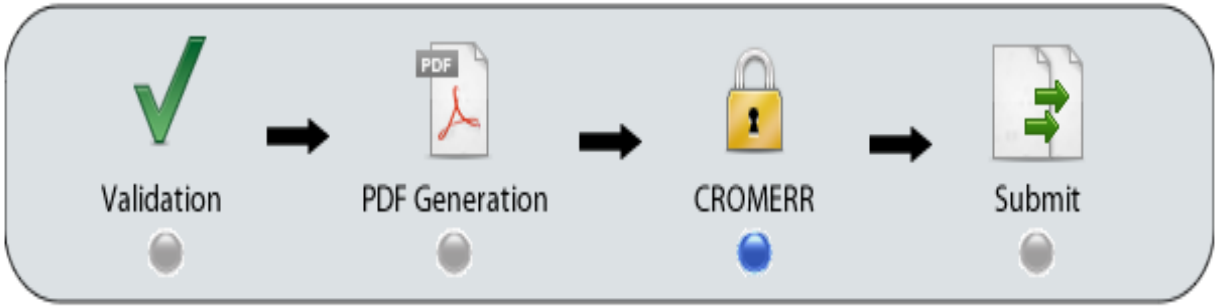


Please login with your CDX password:

Cancel

Next

CROSS-MEDIA ELECTRONIC REPORTING REGULATION (CROMERR) SECURITY QUESTION

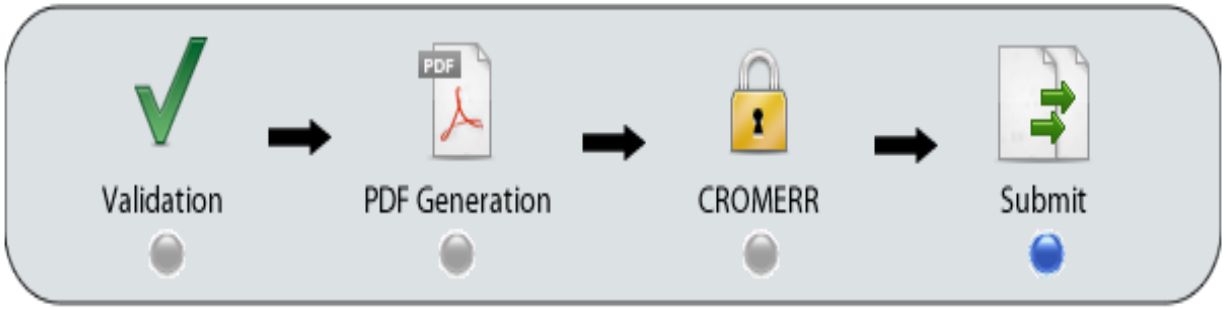


What is your favorite book?

Cancel

Next

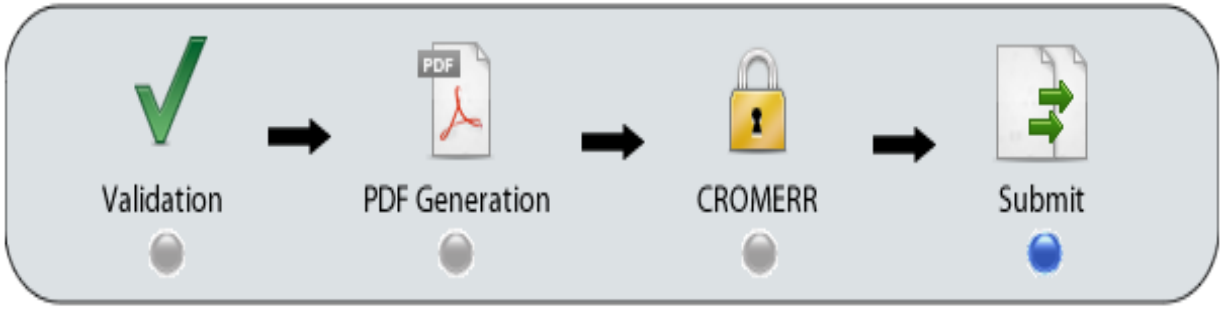
SUBMIT TO CDX



I hereby certify to the best of my knowledge and belief that (1) all information entered on this form is complete and accurate; and (2) any confidentiality claims are true and correct as to that information for which they have been asserted. Any knowing and willful misinterpretation is subject to criminal penalty pursuant to 18 USC 1001

Cancel Confirm

CROSS-MEDIA ELECTRONIC REPORTING REGULATION (CROMERR) SUBMISSION



The submission was sent to the EPA. The Copy of Record link allows you to download of the Copy of Record and signature for this submission. The Copy of Record link will appear in the Submissions list when the EPA receives and processes your submission.

Finish

Section 4: Test Rules/Study Plan

SECTION 4 TEST RULES, ECAS, MOUS, AND 40 CFR 766

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9 items found. Page 1 of 1 Items Per Page: 25

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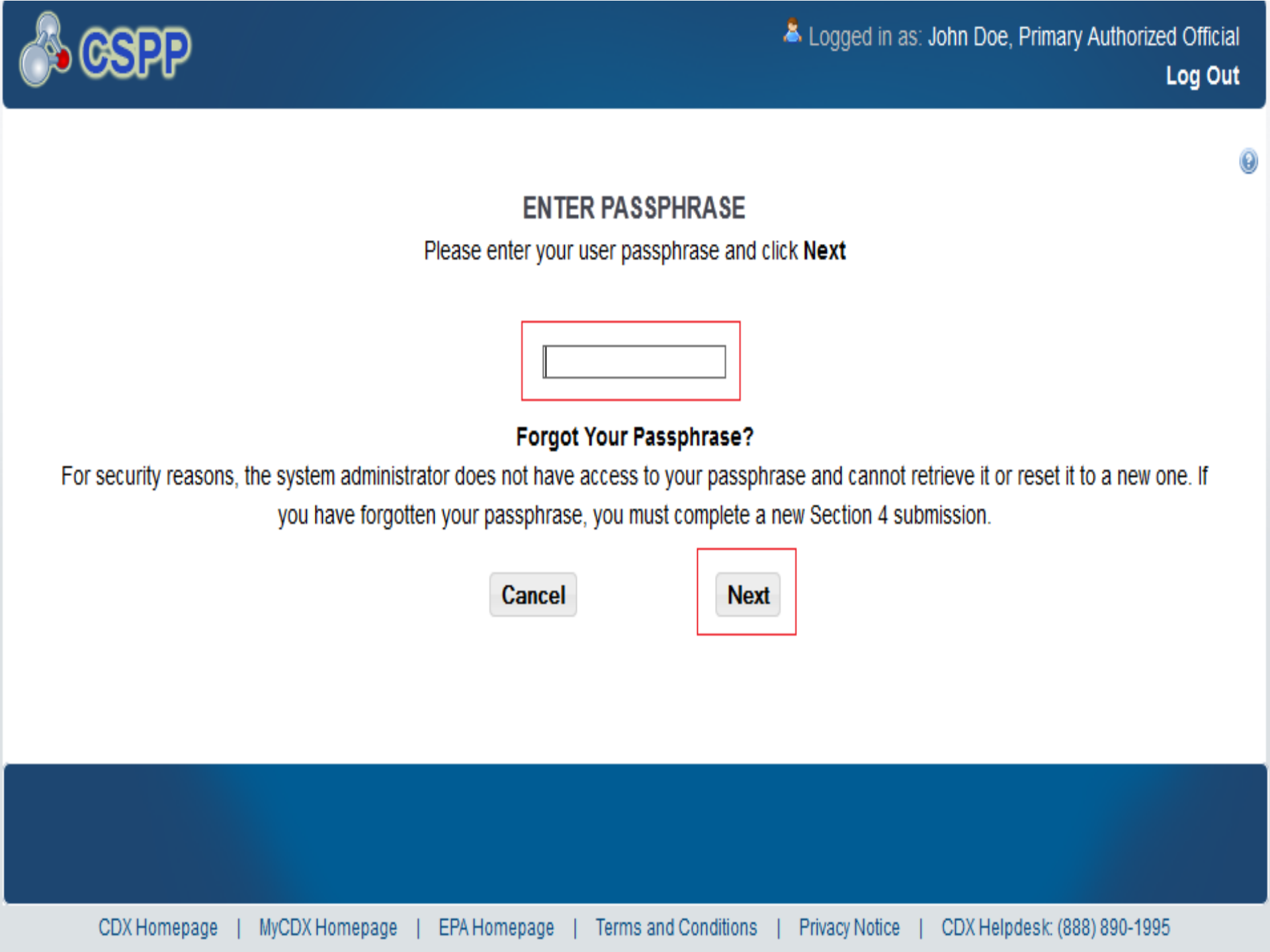
ECA-20131121-12:41:03 EST		In Progress	11/27/2013			
ECA-20131129-15:25:33 EST	70 FR 39630	Submitted	11/29/2013	11/29/2013		
ECA-20131204-15:12:41 EST		In Progress	12/04/2013			
MOU-20131121-12:41:27 EST	MOUForm	In Progress	11/27/2013			
TestRules-20131121-12:41:14 EST	53 FR 22300	In Progress	12/04/2013	11/26/2013		
TestRules-20131127-10:17:23 EST	69 FR 22402	In Progress	11/27/2013			
TestRules-20131206-13:11:56 EST	53 FR 22300	In Progress	12/06/2013			
Test Rule 120613	53 FR 22300	Submitted	11/29/2013	11/29/2013		

Export options: CSV | Excel | XML | PDF

Select the submission type and then click **Start New Submission**
Submission Type:

Start New Submission


CSPP
Logged in as: John Doe, Primary Authorized Official
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 4 submission.
Cancel
Next
CDX Homepage | MyCDX Homepage | EPA Homepage | Terms and Conditions | Privacy Notice | CDX Helpdesk: (888) 890-1995

Section 4 Test Rules
Primary Authorized Official

Section 4 Test Rules > 53 FR 22300 > Study Plan > Document Management

- Submission Information Type
- Contact Information
  - Technical Contact Information
  - Submitting on Behalf of Consortium
- Letter of Intent
  - Chemical Identification and Test Rules
  - Sponsoring Firms
  - Additional Information and Submitter Requests
- 640-19-7
  - Study Plan

DOCUMENT MANAGEMENT

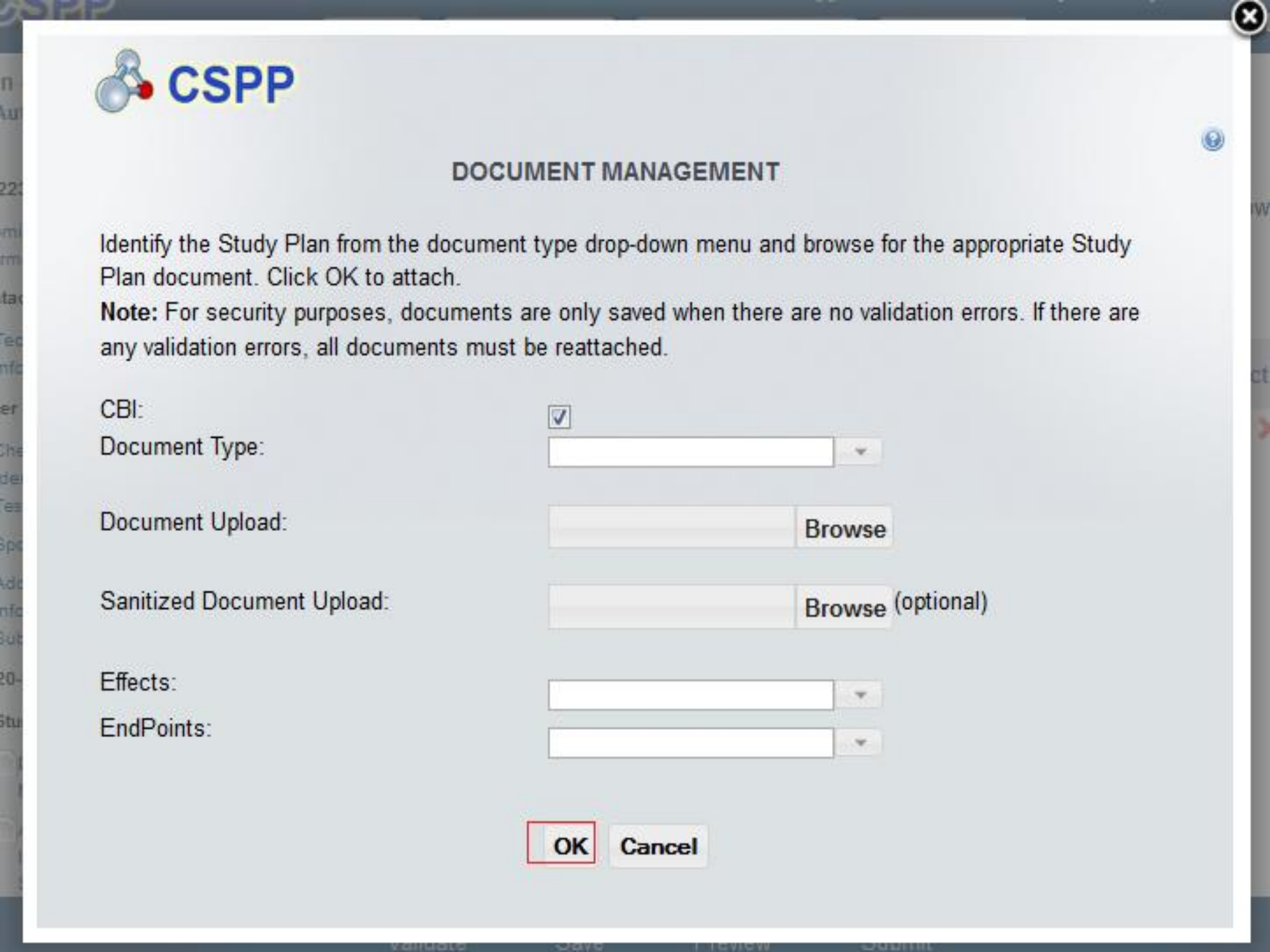
Upload the corresponding Section 4 document(s) by clicking on the Attach Document button below.

Attach Document

File Name	Document Type	Attachment Date	CBI	Action

Nothing found to display.

Previous Next


CSPP
DOCUMENT MANAGEMENT
Identify the Study Plan from the document type drop-down menu and browse for the appropriate Study Plan document. Click OK to attach.
Note: For security purposes, documents are only saved when there are no validation errors. If there are any validation errors, all documents must be reattached.
CBI:
Document Type:
Document Upload:
Browse
Sanitized Document Upload:
Browse
(optional)
Effects:
EndPoints:
OK
Cancel

Section 4 Test Rules
Primary Authorized Official

- 53 FR 22300
  - Submission Information Type
  - Contact Information
    - Technical Contact Information
    - Submitting on Behalf of Consortium
  - Letter of Intent
    - Chemical Identification and Test Rules
    - Sponsoring Firms
    - Additional Information and Submitter Requests
  - 640-19-7
    - Study Plan

Section 4 Test Rules > 53 FR 22300 > Study Plan > Additional Information and Submitter Requests

ADDITIONAL INFORMATION AND SUBMITTER REQUESTS

Select the information or request type from the dropdown and click the Add Information button to complete the required information.

Note: For security purposes, documents are only saved when there are no validation errors. If there are any validation errors, all documents must be reattached.

Information/Request Type:

Add Information

File Name	Document Type	Attachment Date	Action

Nothing found to display.

Previous Next

Section 4 Test Rules
Primary Authorized Official

Section 4 Test Rules > 53 FR 22300 > Study Plan > Test Rules Substantiation Part 1

- 53 FR 22300
  - Submission Information Type
  - Contact Information
    - Technical Contact Information
    - Submitting on Behalf of Consortium
  - Letter of Intent
    - Chemical Identification and Test Rules
    - Sponsoring Firms
    - Additional Information and Submitter Requests
  - 640-19-7
    - Study Plan
      - Document Management
      - Additional Information and Submitter Requests
      - Test Rules Substantiation Part 1
      - Test Rules Substantiation Part 2
    - Results
      - Document Management
  - Remove
  - 16532-79-9
    - Study Plan
      - Document Management
      - Additional Information and Submitter Requests
    - Results

TEST RULES SUBSTANTIATION PART 1

1. Has the information been disclosed in a patent?
   Yes (●) No ( )

2. Would disclosure of the study plan information disclose processes used in the manufacture or processing of a chemical substance or mixture?
   Yes (●) No ( )
   Describe how this would occur.

3. Would disclosure of the study plan information disclose the portion of a mixture comprised by any of the substances in the mixture?
   Yes (●) No ( )
   Describe how this would occur.

4. Has this information been disclosed to the public in any form?
   Yes (●) No ( )
   Describe the circumstances.

Previous Next

Section 4 Test Rules
Primary Authorized Official

Section 4 Test Rules > 53 FR 22300 > Study Plan > Test Rules Substantiation Part 2

- 53 FR 22300
  - Submission Information Type
  - Contact Information
    - Technical Contact Information
    - Submitting on Behalf of Consortium
  - Letter of Intent
    - Chemical Identification and Test Rules
    - Sponsoring Firms
    - Additional Information and Submitter Requests
  - 640-19-7
    - Study Plan
      - Document Management
      - Additional Information and Submitter Requests
      - Test Rules Substantiation Part 1
      - **Test Rules Substantiation Part 2**
    - Results
      - Document Management
  - Remove
  - 16532-79-9
    - Study Plan
      - Document Management

TEST RULES SUBSTANTIATION PART 2

5. For what period of time should confidential treatment be given? Until a specific date, the occurrence of a specific event, or permanently?

- ( ) Event
- ( ) Date
- ( ) Permanently

Why should confidential treatment be given?

6 What harmful effects to your competitive position, if any, do you think would result from disclosure of this information? How would a competitor use such information? How substantial would the harmful effects be? What is the causal relationship between disclosure and the harmful effects?

7. What measures have you taken to guard against disclosure of this information to others?

8. To what extent has this information been disclosed to others? What precautions have been taken in connection with such disclosures?

Section 4 Test Rules
Primary Authorized Official

- 53 FR 22300
  - Submission Information Type
  - Contact Information
    - Technical Contact Information
    - Submitting on Behalf of Consortium
  - Letter of Intent
    - Chemical Identification and Test Rules
    - Sponsoring Firms
    - Additional Information and Submitter Requests
  - 640-19-7
    - Study Plan
      - Document Management
      - Additional Information and Submitter Requests
      - Test Rules Substantiation Part 1
      - Test Rules Substantiation Part 2
    - Results
      - Document Management
  - Remove
  - 16532-79-9
    - Study Plan
      - Document Management

disclosure of this information? How would a competitor use such information? How substantial would the harmful effects be? What is the causal relationship between disclosure and the harmful effects?

7. What measures have you taken to guard against disclosure of this information to others?

8. To what extent has this information been disclosed to others? What precautions have been taken in connection with such disclosures?

9. Has EPA, another Federal Agency, or any Federal court made any pertinent confidentiality determination regarding this information?

Yes (●) No ( )

Click the Browse button and search for the appropriate document(s). Click the Upload button to attach copies of such determinations.

Browse... Upload

File Name	Attachment Date	Action
Nothing found to display.		

Previous Next

Validate Save Preview Submit

Section 4: Test Rules/Results

Section 4 Test Rules
Primary Authorized Official

and Submitter Requests
72-20-8
Study Plan
Document Management
Additional Information and Submitter Requests
Results
Document Management
Remove
Optional Substantiation
Document Management

RESULTS DOCUMENT MANAGEMENT

To add a Results Document, please click the Attach Document button and enter the information in the popup.

Attach Document

File Name	Attachment Date	CBI	Test Type	Action

Previous

CSPP

RESULTS DOCUMENT MANAGEMENT

Identify the Chemical Test from the drop-down menu and browse for the appropriate Results document. Click OK to attach.

Note: For security purposes, documents are only saved when there are no validation errors. If there are any validation errors, all documents must be reattached.

CBI: ☑

Test Type:

Document Upload: Browse

Sanitized Document Upload: Browse (optional)

Effects:

EndPoints:

OK Cancel

Section 4 Test Rules
Primary Authorized Official

- 53 FR 22300
  - Submission Information Type
  - Contact Information
    - Technical Contact Information
  - Letter of Intent
    - Chemical Identification and Test Rules
    - Sponsoring Firms
    - Additional Information and Submitter Requests
  - 640-19-7
    - Remove
  - Optional Substantiation
    - Document Management

OPTIONAL DOCUMENT MANAGEMENT

Upload the Optional substantiation document(s) by clicking on the Attach Document button below.

Attach Document

File Name	Attachment Date	Action

Section 4: Enforceable Consent Agreements (ECAs)

SECTION 4 TEST RULES, ECAS, MOUS, AND 40 CFR 766

- If starting a Section 4 Test Rule, ECA, MOU, or a 40 CFR 766 submission, select the appropriate submission type from the drop-down menu and click the 'Start New Submission' button.
- To edit an **In Progress** submission, click the submission link in the **Submission Alias** column in the table below.
- To access and edit a submission previously **Submitted** through CDX, unlock the submission by clicking the lock icon ( ) and enter your passphrase originally associated with the selected submission. All additional changes made to a submission will be submitted as an amendment.
- Click the green arrow icon ( ) to download a copy of record for a completed submission. It may take up to 15 minutes for the copy of record to become available.
- You may delete any submission that has not yet been submitted by clicking the delete icon ( ).

4 items found. Page 1 of 1 Items Per Page: 25

Submission Alias	CFR/FRN	Status	Modify Date	Submission Date	Copy of Record	Action
CFR766-20131121-12:41:40 EST	40 CFR 766 Dibenzodioxins / Dibenzofurans	In Progress	11/21/2013			
ECA-20131121-12:41:03 EST		In Progress	11/21/2013			
MOU-20131121-12:41:27 EST	MOUForm	In Progress	11/21/2013			
TestRules-20131121-12:41:14 EST		In Progress	11/21/2013			

Export options: CSV | Excel | XML | PDF

Select the submission type and then click **Start New Submission**

Submission Type: ECA

Start New Submission

CREATE PASSPHRASE

Please create a passphrase that is at least 8 characters in length. To better protect your form, your passphrase should contain a combination of letters numbers. Your passphrase may include spaces, but should not contain special characters (for example, + and *).

As an Authorized Official, you are responsible for remembering your passphrase and distributing it to only authorized individuals. 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 only authorized Supports.

Note: If you lose or forget your passphrase, you will not be able to access your Section 4 Submission to print, submit, or make changes. You will need to complete a new Section 4 Submission and create a new passphrase for the submission. For security reasons, the system administrator will not have access to your passphrase and will not be able to retrieve it or reset it.

New Passphrase:
Confirm New Passphrase:

Cancel Next

Section 4 Enforceable Consent Agreement (ECA)
Primary Authorized Official

- Federal Register Notice
  - Submission Information Type
- Contact Information
  - Technical Contact Information
  - ECA Additional Information

Section 4 Enforceable Consent Agreement (ECA) > Federal Register Notice > Submission Information Type

SUBMISSION INFORMATION TYPE

Please select, or start entering, a chemical in the drop-down menu below:

Please enter a submission alias and choose the submission information type(s) here.

Submission Alias: ECA-20130711-13:14:59

- [ ] Study Plans & Conduct of Testing
- [ ] Results

Next

Section 4 Enforceable Consent Agreement (ECA)
Primary Authorized Official

- 68 FR 33125
  - Submission Information Type
  - Contact Information
    - Technical Contact Information
  - Study Plans & Conduct of Testing
    - Principal Test Sponsor
    - Principal Sponsor Organization
    - Testing Facilities
    - Study Professionals
    - Remove
  - Results
    - Document Management
    - Remove
  - ECA Additional Information

Section 4 Enforceable Consent Agreement (ECA) > 68 FR 33125 > 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 selecting the appropriate radio button. If submitting on behalf of is not applicable, select the N/A radio button. Click the 'Copy CDX Registration' button to copy your information from CDX Registration.

- ( ) N/A
- ( ) This is a submission on behalf of another company

Click here to copy your information from CDX Registration: **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:

Previous **Next**

Section 4 Enforceable Consent Agreement (ECA)
Primary Authorized Official

- 68 FR 33125
  - Submission Information Type
  - Contact Information
    - Technical Contact Information
    - Submitting on Behalf of Company
  - Study Plans & Conduct of Testing
    - Principal Test Sponsor
    - Principal Sponsor Organization
    - Testing Facilities
    - Study Professionals
    - Remove
  - Results
    - Document Management
    - Remove
  - ECA Additional Information

Section 4 Enforceable Consent Agreement (ECA) > 68 FR 33125 > Study Plans & Conduct of Testing > Principal Test Sponsor

PRINCIPAL TEST SPONSOR

Fill out the information below for the Principal Test Sponsor.

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

Section 4 Enforceable Consent Agreement (ECA)
Primary Authorized Official

- 68 FR 33125
  - Submission Information Type
  - Contact Information
    - Technical Contact Information
    - Submitting on Behalf of Company
  - Study Plans & Conduct of Testing
    - Principal Test Sponsor
    - Principal Sponsor Organization
    - Testing Facilities
    - Study Professionals
    - Remove
  - Results
    - Document Management
    - Remove
  - ECA Additional Information

Section 4 Enforceable Consent Agreement (ECA) > 68 FR 33125 > Study Plans & Conduct of Testing > Principal Sponsor Organization

PRINCIPAL SPONSOR ORGANIZATION

Fill out the information below for the Administrative Official(s) and Project Manager(s) in the Principal Sponsor's Organization below.

Expand All | Collapse All

Firm - Cancel

CBI:
Prefix:
First Name:
Middle Initial:
Last Name:
Suffix:
Company Name:
Job Title:
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

Click **Add Sponsor Organization** to add a new sponsor organization. **Add Sponsor Organization**

Previous **Next**

Section 4 Enforceable Consent Agreement (ECA)
Primary Authorized Official
68 FR 33125
Submission Information Type
Contact Information
Technical Contact Information
Submitting on Behalf of Company
Study Plans & Conduct of Testing
Principal Test Sponsor
Principal Sponsor Organization
Testing Facilities
Study Professionals
Remove
Results
Document Management
Remove
ECA Additional Information

TESTING FACILITIES

Fill out the information below for the responsible testing facilities.

Expand All | Collapse All

Facility - Cancel

Testing Facility:
Phone Number: Ext:
(Do not enter any dashes (-) in Phone Number field above.)
Mailing Address 1:
Street address, P.O. box, company name, etc.
Mailing Address 2:
Apartment, suite, etc.
City:
State:
Postal Code:
Country
Select the testing facility contact role:
CBI:
Prefix:
First Name:
Middle Initial:
Last Name:
Suffix:
Phone Number: Ext:
(Do not enter any dashes (-) in Phone Number field above)

Validate | Save | Preview | Submit

Section 4 Enforceable Consent Agreement (ECA)
Primary Authorized Official
68 FR 33125
Submission Information Type
Contact Information
Technical Contact Information
Submitting on Behalf of Company
Study Plans & Conduct of Testing
Principal Test Sponsor
Principal Sponsor Organization
Testing Facilities
Study Professionals
Remove
Results
Document Management
Remove
ECA Additional Information

Postal Code:
Country
Select the testing facility contact role:
CBI:
Prefix:
First Name:
Middle Initial:
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

Click the **Add Testing Facility** button to add a new testing facility. Add Testing Facility

Previous Next

Section 4 Enforceable Consent Agreement (ECA)
Primary Authorized Official
- 68 FR 33125
  - Submission Information Type
  - Contact Information
    - Technical Contact Information
    - Submitting on Behalf of Company
  - Study Plans & Conduct of Testing
    - Principal Test Sponsor
    - Principal Sponsor Organization
    - Testing Facilities
    - Study Professionals
    - Remove
  - Results
    - Document Management
    - Remove
  - ECA Additional Information

STUDY PROFESSIONALS

Provide a brief summary of the training and experience of each professional involved in the study below.

Expand All | Collapse All

Study Professional - Cancel

Select the Study Professional Role:

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:

Experience Summary:

Validate | Save | Preview | Submit

Section 4 Enforceable Consent Agreement (ECA)
Primary Authorized Official
68 FR 33125
Submission Information Type
Contact Information
Technical Contact Information
Submitting on Behalf of Company
Study Plans & Conduct of Testing
Principal Test Sponsor
Principal Sponsor Organization
Testing Facilities
Study Professionals
Remove
Results
Document Management
Remove
ECA Additional Information

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:
Experience Summary:

Save

Click the **Add Study Professional** button to add a new study professional. Add Study Professional

Previous Next

Section 4 Enforceable Consent Agreement (ECA)
Primary Authorized Official
Type
Contact Information
Technical Contact Information
Study Plans & Conduct of Testing
Principal Test Sponsor
Principal Sponsor Organization
Testing Facilities
Study Professionals
Remove
Results
Document Management
Remove
ECA Additional

DOCUMENT MANAGEMENT

To add a Results Document, please click the Attach Document button and enter the information in the popup.

Attach Document

File Name	Attachment Date	CBI	Action
Nothing found to display.			

Previous Next

CSPP

DOCUMENT MANAGEMENT

Browse for the appropriate Results document and click the OK button to attach.

Note: For security purposes, documents are only saved when there are no validation errors. If there are any validation errors, all documents must be reattached.

CBI: ☑

Document Upload: Browse

Sanitized Document Upload: Browse (optional)

Effects:

EndPoints:

OK Cancel

Section 4 Enforceable Consent Agreement (ECA)
Primary Authorized Official

- Federal Register Notice
  - Submission Information Type
  - Contact Information
    - Technical Contact Information
  - Study Plans & Conduct of Testing
    - Principal Test Sponsor
    - Principal Sponsor Organization
    - Testing Facilities
    - Study Professionals
    - Remove
  - Results
    - Document Management
    - Remove
  - ECA Additional Information

ECA ADDITIONAL INFORMATION

Select the appropriate option below and upload the corresponding document.

- (•) Amendments to the Study Plan
- ( ) Modification of ECAs

Attach Document

File Name	Document Type	Attachment Date	Action

Nothing found to display.

Previous

Section 4 Enforceable Consent Agreement (ECA)
Primary Authorized Official

- 68 FR 33125
  - Submission Information Type
  - Contact Information
    - Technical Contact Information
    - Submitting on Behalf of Company
  - Study Plans & Conduct of Testing
    - Principal Test Sponsor
    - Principal Sponsor Organization
    - Testing Facilities
    - Study Professionals
    - Remove
  - Results
    - Document Management
    - Remove
  - ECA Additional Information

Section 4 Enforceable Consent Agreement (ECA) > 68 FR 33125 > ECA Additional Information

ECA ADDITIONAL INFORMATION

Select the appropriate option below and upload the corresponding document.

- ( ) Amendments to the Study Plan
- (•) Modification of ECAs

Attach Document

File Name	Document Type	Attachment Date	Action

Nothing found to display.

Previous

Section 4: Memorandum of Understanding (MOUs)

SECTION 4 TEST RULES, ECAS, MOUS, AND 40 CFR 766

- If starting a Section 4 Test Rule, ECA, MOU, or a 40 CFR 766 submission, select the appropriate submission type from the drop-down menu and click the 'Start New Submission' button.
- To edit an **In Progress** submission, click the submission link in the **Submission Alias** column in the table below.
- To access and edit a submission previously **Submitted** through CDX, unlock the submission by clicking the lock icon ( ) and enter your passphrase originally associated with the selected submission. All additional changes made to a submission will be submitted as an amendment.
- Click the green arrow icon ( ) to download a copy of record for a completed submission. It may take up to 15 minutes for the copy of record to become available.
- You may delete any submission that has not yet been submitted by clicking the delete icon ( ).

4 items found. Page 1 of 1 Items Per Page: 25

Submission Alias	CFR/FRN	Status	Modify Date	Submission Date	Copy of Record	Action
CFR766-20131121-12:41:40 EST	40 CFR 766 Dibenzodioxins / Dibenzofurans	In Progress	11/21/2013			
ECA-20131121-12:41:03 EST		In Progress	11/21/2013			
MOU-20131121-12:41:27 EST	MOUForm	In Progress	11/21/2013			
TestRules-20131121-12:41:14 EST		In Progress	11/21/2013			

Export options: CSV | Excel | XML | PDF

Select the submission type and then click **Start New Submission**

Submission Type: MOU

Start New Submission

CREATE PASSPHRASE

Please create a passphrase that is at least 8 characters in length. To better protect your form, your passphrase should contain a combination of letters numbers. Your passphrase may include spaces, but should not contain special characters (for example, + and *).

As an Authorized Official, you are responsible for remembering your passphrase and distributing it to only authorized individuals. 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 only authorized Supports.

Note: If you lose or forget your passphrase, you will not be able to access your Section 4 Submission to print, submit, or make changes. You will need to complete a new Section 4 Submission and create a new passphrase for the submission. For security reasons, the system administrator will not have access to your passphrase and will not be able to retrieve it or reset it.

New Passphrase:
Confirm New Passphrase:

Cancel Next

Section 4 Memorandum of Understanding (MOU)
Primary Authorized Official
- Federal Register Notice
  - Submission Information Type
  - Contact Information
    - Technical Contact Information
  - Study Plans & Conduct of Testing
    - Principal Test Sponsor
    - Principal Sponsor Organization
    - Testing Facilities
    - Study Professionals
    - Remove
  - Results
    - Document Management
    - Remove
  - MOU Additional Information

Section 4 Memorandum of Understanding (MOU) > Federal Register Notice> 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 selecting the appropriate radio button. If submitting on behalf of is not applicable, select the N/A radio button. Click the 'Copy CDX Registration' button to copy your information from CDX Registration.

- ( ) N/A
- ( ) This is a submission on behalf of another company

Click here to copy your information from CDX Registration: **Copy CDX Registration**

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**

Section 4 Memorandum of Understanding (MOU)
Primary Authorized Official

- Federal Register Notice
  - Submission Information Type
  - Contact Information
    - Technical Contact Information
    - Submitting on Behalf of Company
  - Study Plans & Conduct of Testing
    - Principal Test Sponsor
    - Principal Sponsor Organization
    - Testing Facilities
    - Study Professionals
    - Remove
  - Results
    - Document Management
    - Remove
  - MOU Additional Information

Section 4 Memorandum of Understanding (MOU) > Federal Register Notice> Study Plans & Conduct of Testing > Principal Test Sponsor

PRINCIPAL TEST SPONSOR

Fill out the information below for the Principal Test Sponsor.

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

Section 4 Memorandum of Understanding (MOU)
Primary Authorized Official
- Federal Register Notice
  - Submission Information Type
  - Contact Information
    - Technical Contact Information
    - Submitting on Behalf of Company
  - Study Plans & Conduct of Testing
    - Principal Test Sponsor
    - Principal Sponsor Organization
    - Testing Facilities
    - Study Professionals
    - Remove
  - Results
    - Document Management
    - Remove
  - MOU Additional Information

PRINCIPAL SPONSOR ORGANIZATION

Fill out the information below for the Administrative Official(s) and Project Manager(s) in the Principal Sponsor's Organization below.

Expand All | Collapse All

Firm - Cancel

Prefix:
First Name:
Middle Initial:
Last Name:
Suffix:
Company Name:
Job Title:
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

Click **Add Sponsor Organization** to add a new sponsor organization. Add Sponsor Organization

Previous Next

Section 4 Memorandum of Understanding (MOU)
Primary Authorized Official
- Federal Register Notice
  - Submission Information Type
- Contact Information
  - Technical Contact Information
  - Submitting on Behalf of Company
- Study Plans & Conduct of Testing
  - Principal Test Sponsor
  - Principal Sponsor Organization
  - Testing Facilities
  - Study Professionals
  - Remove
- Results
  - Document Management
  - Remove
- MOU Additional Information

Section 4 Memorandum of Understanding (MOU) > Federal Register Notice> Study Plans & Conduct of Testing > Testing Facilities

TESTING FACILITIES

Fill out the information below for the responsible testing facilities.

Expand All | Collapse All

Facility - Cancel

Testing Facility:

Phone Number: Ext:

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

Mailing Address 1:

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

Mailing Address 2:

Apartment, suite, etc.

City:

State:

Postal Code:

Country

Select the testing facility contact role:

Prefix:

First Name:

Middle Initial:

Last Name:

Suffix:

Phone Number: Ext:

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

Email Address:

Section 4 Memorandum of Understanding (MOU)
Primary Authorized Official
- Federal Register Notice
  - Submission Information Type
- Contact Information
  - Technical Contact Information
  - Submitting on Behalf of Company
- Study Plans & Conduct of Testing
  - Principal Test Sponsor
  - Principal Sponsor Organization
  - Testing Facilities
  - Study Professionals
  - Remove
- Results
  - Document Management
  - Remove
- MOU Additional Information

State:
Postal Code:
Country

Select the testing facility contact role:
Prefix:
First Name:
Middle Initial:
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

Click the **Add Testing Facility** button to add a new testing facility. Add Testing Facility

Previous Next

Section 4 Memorandum of Understanding (MOU)
Primary Authorized Official
- Federal Register Notice
  - Submission Information Type
- Contact Information
  - Technical Contact Information
  - Submitting on Behalf of Company
- Study Plans & Conduct of Testing
  - Principal Test Sponsor
  - Principal Sponsor Organization
  - Testing Facilities
  - Study Professionals
  - Remove
- Results
  - Document Management
  - Remove
- MOU Additional Information

STUDY PROFESSIONALS

Provide a brief summary of the training and experience of each professional involved in the study below.

Expand All | Collapse All

Study Professional - Cancel

Select the Study Professional Role:
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:
Experience Summary:

Save

Click the **Add Study Professional** button to add a new study professional. Add Study Professional

Previous Next

Validate | Save | Preview | Submit

Section 4 Memorandum of Understanding (MOU)
Primary Authorized Official

- Federal Register Notice
  - Submission Information Type
  - Contact Information
    - Technical Contact Information
    - Submitting on Behalf of Company
  - Study Plans & Conduct of Testing
    - Principal Test Sponsor
    - Principal Sponsor Organization
    - Testing Facilities
    - Study Professionals
    - Remove
  - Results
    - Document Management
    - Remove
  - MOU Additional Information

DOCUMENT MANAGEMENT

To add a Results Document, please click the Attach Document button and enter the information in the popup.

Attach Document

File Name	Attachment Date	Action

Nothing found to display.

Previous Next

CSPP

DOCUMENT MANAGEMENT

Select the document type and browse for the document. If the document to be attached is not a study or an abstract/summary, it is to be identified as a supplemental document.

Note: For security purposes, documents are only saved when there are no validation errors. If there are any validation errors, all documents must be reattached.

Document Upload: Browse

Effects:

EndPoints:

OK Cancel

Section 4 Memorandum of Understanding (MOU)
Primary Authorized Official

- Federal Register Notice
  - Submission Information Type
  - Contact Information
    - Technical Contact Information
    - Submitting on Behalf of Company
  - Study Plans & Conduct of Testing
    - Principal Test Sponsor
    - Principal Sponsor Organization
    - Testing Facilities
    - Study Professionals
    - Remove
  - Results
    - Document Management
    - Remove
  - MOU Additional Information

Section 4 Memorandum of Understanding (MOU) > Federal Register Notice> MOU Additional Information

MOU ADDITIONAL INFORMATION

Select the appropriate option below and upload the corresponding document.

- ○ Amendments to the Study Plan
- ○ Modification of MOUs

Attach Document

File Name	Document Type	Attachment Date	Action

Nothing found to display.

Previous

Section 4: 40 CFR 766 Dibenzodioxins/Dibenzofurans

SECTION 4 TEST RULES, ECAS, MOUS, AND 40 CFR 766

- If starting a Section 4 Test Rule, ECA, MOU, or a 40 CFR 766 submission, select the appropriate submission type from the drop-down menu and click the 'Start New Submission' button.
- To edit an **In Progress** submission, click the submission link in the **Submission Alias** column in the table below.
- To access and edit a submission previously **Submitted** through CDX, unlock the submission by clicking the lock icon ( ) and enter your passphrase originally associated with the selected submission. All additional changes made to a submission will be submitted as an amendment.
- Click the green arrow icon ( ) to download a copy of record for a completed submission. It may take up to 15 minutes for the copy of record to become available.
- You may delete any submission that has not yet been submitted by clicking the delete icon ( ).

4 items found. Page 1 of 1 Items Per Page: 25

Submission Alias	CFR/FRN	Status	Modify Date	Submission Date	Copy of Record	Action
CFR766-20131121-12:41:40 EST	40 CFR 766 Dibenzodioxins / Dibenzofurans	In Progress	11/21/2013			
ECA-20131121-12:41:03 EST		In Progress	11/21/2013			
MOU-20131121-12:41:27 EST	MOUForm	In Progress	11/21/2013			
TestRules-20131121-12:41:14 EST		In Progress	11/21/2013			

Export options: CSV | Excel | XML | PDF

Select the submission type and then click **Start New Submission**

Submission Type: 40 CFR 766 Dibenzodioxins/Dibenzofurans

Start New Submission

CREATE PASSPHRASE

Please create a passphrase that is at least 8 characters in length. To better protect your form, your passphrase should contain a combination of letters numbers. Your passphrase may include spaces, but should not contain special characters (for example, + and *).

As an Authorized Official, you are responsible for remembering your passphrase and distributing it to only authorized individuals. 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 only authorized Supports.

Note: If you lose or forget your passphrase, you will not be able to access your Section 4 Submission to print, submit, or make changes. You will need to complete a new Section 4 Submission and create a new passphrase for the submission. For security reasons, the system administrator will not have access to your passphrase and will not be able to retrieve it or reset it.

New Passphrase:
Confirm New Passphrase:

Cancel Next

Section 4 § 40 CFR 766 Dibenzodioxins / Dibenzofurans
Primary Authorized Official
- 40 CFR 766 Dibenzodioxins / Dibenzofurans
  - Submission Information Type
  - Contact Information
    - Technical Contact Information
    - Submitting on Behalf of Consortium
  - Chemical Information
    - Chemical Identification

SUBMISSION INFORMATION TYPE

Please enter a submission alias and choose the submission information type(s) here. You can either Select User Requests or any of the other options.

Submission Alias: CFR766-20130826-11:3

- [ ] Letter of Intent
- [ ] Protocol
- [ ] Results
- [ ] User Requests
  - ( ) Waiver
  - ( ) Exclusion
  - ( ) Exemption

Next

Section 4 § 40 CFR 766 Dibenzodioxins / Dibenzofurans
Primary Authorized Official
- 40 CFR 766 Dibenzodioxins / Dibenzofurans
  - Submission Information Type
  - Contact Information
    - Technical Contact Information
    - Submitting on Behalf of Company
  - Chemical Information
    - Chemical Identification
  - Letter of Intent
    - Sponsoring Firms
    - Remove
  - Protocol
    - Document Management
    - Remove
  - Results
    - Document Management
    - Remove

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 or consortium by selecting the appropriate radio button. If submitting on behalf of is not applicable, select the N/A radio button. Click the 'Copy CDX Registration' button to copy your information from CDX Registration.

- ( ) N/A
- ( ) This is a submission on behalf of a consortium
- (•) This is a submission on behalf of another company

Click here to copy your information from CDX Registration: **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:

Previous **Next**

Section 4 § 40 CFR 766 Dibenzodioxins / Dibenzofurans
Primary Authorized Official

- 40 CFR 766 Dibenzodioxins / Dibenzofurans
  - Submission Information Type
  - Contact Information
    - Technical Contact Information
    - Submitting on Behalf of Consortium
  - Chemical Information
    - Chemical Identification
  - Letter of Intent
    - Sponsoring Firms
    - Remove

Section 4 § 40 CFR 766 Dibenzodioxins / Dibenzofurans > Letter of Intent > Chemical Identification

CHEMICAL IDENTIFICATION

Select the chemical that the sponsor is manufacturing or importing to which the submission applies.

☐ CBI

Please select, or begin typing, a **CASRN** in the drop-down menu below:

Previous Next

Section 4 § 40 CFR 766 Dibenzodioxins / Dibenzofurans
Primary Authorized Official

- 40 CFR 766 Dibenzodioxins / Dibenzofurans
  - Submission Information Type
  - Contact Information
    - Technical Contact Information
    - Submitting on Behalf of Company
  - Chemical Information
    - Chemical Identification
  - Letter of Intent
    - Sponsoring Firms
    - Remove
  - Protocol
    - Document Management
    - Remove
  - Results
    - Document Management
    - Remove

SPONSORING FIRMS

Fill out the information below for the Sponsoring Firm.

Expand All | Collapse All

Firm -

Cancel

CBI:

Firm Name:

Phone Number: Ext:

Mailing Address 1:

Mailing Address 2:

City:

State:

Postal Code:

Country:

Save

Click the **Add Sponsoring Firm** button to add a new sponsoring firm. Add Sponsoring Firm

Previous Next

Validate Save Preview Submit

Section 4 § 40 CFR 766 Dibenzodioxins / Dibenzofurans
Primary Authorized Official
Technical Contact Information
Submitting on Behalf of Consortium
Chemical Information
Chemical Identification
Letter of Intent
Sponsoring Firms
Remove
Protocol
Document Management
Remove
Results
Document

PROTOCOL DOCUMENT MANAGEMENT

Upload the corresponding protocol by clicking on the Attach Document button below.

Attach Document

File Name	Document Type	Attachment Date	CBI	Action

Nothing found to display.

Previous Next

CSPP

PROTOCOL DOCUMENT MANAGEMENT

Identify the document type and browse for the appropriate protocol document. Click OK to attach.

Note: For security purposes, documents are only saved when there are no validation errors. If there are any validation errors, all documents must be reattached.

CBI: ☑

Document Type:

Document Upload: Browse

Sanitized Document Upload: Browse (optional)

Effects:

EndPoints:

OK Cancel

Section 4 § 40 CFR 766 Dibenzodioxins / Dibenzofurans
Primary Authorized Official
Information
Submitting on Behalf of Company
Chemical Information
Chemical Identification
Letter of Intent
Sponsoring Firms
Remove
Protocol
Document Management
Remove
Results
Document

DOCUMENT MANAGEMENT

To add a Results Document, please click the Attach Document button and enter the information in the popup.

Attach Document

File Name	Attachment Date	CBI	Action

Nothing found to display.

Previous

CSPP

DOCUMENT MANAGEMENT

Browse for the appropriate document and click OK to attach.

Note: For security purposes, documents are only saved when there are no validation errors. If there are any validation errors, all documents must be reattached.

CBI: ☑

Document Upload: Browse

Sanitized Document Upload: Browse (optional)

Effects:

EndPoints:

OK Cancel

Section 4 § 40 CFR 766 Dibenzodioxins / Dibenzofurans
Primary Authorized Official
- 40 CFR 766 Dibenzodioxins / Dibenzofurans
  - Submission Information Type
  - Contact Information
    - Technical Contact Information
    - Submitting on Behalf of Company
  - Chemical Information
    - Chemical Identification
  - User Requests
    - Waiver
    - Remove

WAIVER

To request a waiver, please identify the appropriate qualifying reason and then click the Attach Document button to attach any supporting documentation.

( ) The chemical substance is produced only in quantities of 100 kilograms or less per year, only for research and development purposes

( ) The cost of testing would drive the chemical substance off the market, or prevent resumption of manufacture or import of the chemical substance, if it is not currently manufactured, and the chemical substance will be produced so that no unreasonable risk will occur due to its manufacture, import, processing, distribution, use, or disposal. (In this case, the manufacturer must submit to EPA all data supporting the determination.)

Attach Document

File Name	Attachment Date	CBI	Action

Nothing found to display.

Previous

Section 4 § 40 CFR 766 Dibenzodioxins / Dibenzofurans
Primary Authorized Official

- 40 CFR 766 Dibenzodioxins / Dibenzofurans
  - Submission Information Type
  - Contact Information
    - Technical Contact Information
    - Submitting on Behalf of Company
  - Chemical Information
    - Chemical Identification
  - User Requests
    - Exclusion
    - Remove

Section 4 § 40 CFR 766 Dibenzodioxins / Dibenzofurans > User Requests > Exclusion

EXCLUSION

To request exclusion, please identify the appropriate qualifying reason and then click the Attach Document button to attach required documentation.

- ( ) Testing of the appropriate grade of the chemical substance has already been carried out, either analytical testing at the lowest LOQ possible, with appropriate QA/QC, or a well-designed bioassay with appropriate QA/QC
- ( ) Process and reaction conditions of the chemical substance such that no HDDs/HDFs could be produced under those conditions.

Attach Document

File Name	Attachment Date	CBI	Action
Nothing found to display.			

Previous

Section 4 § 40 CFR 766 Dibenzodioxins / Dibenzofurans
Primary Authorized Official

- 40 CFR 766 Dibenzodioxins / Dibenzofurans
  - Submission Information Type
  - Contact Information
    - Technical Contact Information
    - Submitting on Behalf of Company
  - Chemical Information
    - Chemical Identification
  - User Requests
    - Exemption
    - Remove

EXEMPTION

To request an exemption, please click the Attach Document button to attach required documentation.

Attach Document

File Name	Attachment Date	CBI	Action

Nothing found to display.

Previous

Section 4: Download a Copy of Record

SECTION 4 TEST RULES, ECAS, MOUS, AND 40 CFR 766

- If starting a Section 4 Test Rule, ECA, MOU, or a 40 CFR 766 submission, select the appropriate submission type from the drop-down menu and click the 'Start New Submission' button.
- To edit an **In Progress** submission, click the submission link in the **Submission Alias** column in the table below.
- To access and edit a submission previously **Submitted** through CDX, unlock the submission by clicking the lock icon ( ) and enter your passphrase originally associated with the selected submission. All additional changes made to a submission will be submitted as an amendment.
- Click the green arrow icon ( ) to download a copy of record for a completed submission. It may take up to 15 minutes for the copy of record to become available.
- You may delete any submission that has not yet been submitted by clicking the delete icon ( ).

5 items found. Page 1 of 1 Items Per Page: 25

Submission Alias	CFR/FRN	Status	Modify Date	Submission Date	Copy of Record	Action
CFR766-20131125-11:08:20 EST	40 CFR 766 Dibenzodioxins / Dibenzofurans	In Progress	11/25/2013			
ECA-20131125-11:05:14 EST	68 FR 33125	Submitted	11/25/2013	11/25/2013		
MOU-20131125-11:08:02 EST	MOUForm	In Progress	11/25/2013			
TestRules 1	53 FR 22300	In Progress	11/25/2013	11/22/2013		
TestRules-20131125-15:31:19 EST	53 FR 22300	In Progress	11/25/2013			

Export options: CSV | Excel | XML | PDF

Select the submission type and then click **Start New Submission**

Submission Type:

Start New Submission

DOWNLOAD COPY OF RECORD

You can now download the Copy of Record for the Section 4 Submission!

Download Copy of Record:

File Name	Actions
Copy of Record	

Download Attachments:

File Name	File Type	File Size	Actions

Home

We will resume in 5 minutes

Section 8(d) Health & Safety Data Reporting Tool

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

The information collection requirements contained in this final rule have been submitted for OMB approval under PRA, 44 U.S.C. 3501 et seq. The ICR document prepared by EPA, identified under EPA ICR No. 2412.01 and OMB control number 2070-0004, is available in the docket for the proposed rule. The ICR addresses the incremental changes to the currently approved ICR documents that cover the existing reporting and record keeping programs that are approved under OMB control numbers 2070-0004, 2070-0033, and 2070-0054. 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 amended information collection activities contained in this final rule are designed to assist the Agency in meeting its responsibility under TSCA to receive, process, and review reports, data, and other information. As such, responses to the collection of information covered by this ICR would still be mandatory, but with the final rule, respondents would be required to use the CISS reporting tool.

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.

HOME

Submissions

Create, modify, or delete 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.

User Management

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

Resources

A helpful guide that describes the Section 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 submissions. The Authorized Official may also submit completed submissions either electronically or by mail. Finally, the Authorized Official has the ability to assign Supports to individual submissions.

USER MANAGEMENT

The Authorized Official is responsible for restricting a Support's access to select submissions by assigning or unassigning them to each submission alias. The Support can access and edit only those submissions for which the Authorized Official has granted access. Select a submission alias from the **Submission Alias** drop-down menu, and assign a Support to the submission by highlighting the individual and clicking the **add** link. To unassign a Support, highlight the individual and click the **remove** link. To highlight and assign or unassign multiple Supports, hold down the **Ctrl** or **Shift** keys on the keyboard and click each Support before moving. You must click the **Save** button after each submission assignment.

Submission Alias:

Submission Alias

CASRN(s):
Federal Register Notice:

Assign Users

Unassigned		Assigned
	add >> << remove	

HOME

Submissions

Create, modify, or delete a submission by clicking the **Submissions** tab.

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- 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.

User Management

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SECTION 8(d) HEALTH & SAFETY DATA REPORTING

- If starting a Section 8(d) Health & Safety Data Reporting submission for the first time in CDX, select a CFR/FRN from the drop-down menu and click the **Start New Submission** button.
- To edit an **In Progress** submission, click the submission alias link in the **Submission Alias** column in the table below.
- To access and edit a form previously Submitted through CDX, unlock the form by clicking the lock icon ( ) and enter your passphrase originally associated with the selected form. All additional changes made to a submission will be submitted as an amendment.
- Click the green arrow icon ( ) to download a copy of record for a submitted form. It may take up to 15 minutes for the copy of record to become available.
- You may delete any submission that has not yet been submitted by clicking the delete icon ( ).

Page 1 of 1

Items per page: 25

2 items found.

Submission Alias	CFR/FRN	CASRN	Status	Last Modified	Submission Date	Copy of Record	Action
Fri Nov 22 10:51:52 EST 2013	75 FR 34076	1336-36-3	In Progress	11/22/2013	11/22/2013		
Thu Nov 21 19:56:06 EST 2013	75 FR 34076	1336-36-3	In Progress	11/21/2013			

Export options: CSV | Excel | XML | PDF

Select Applicable CFR/FRN:

Start New Submission

CREATE PASSPHRASE

Please create a passphrase that is at least 8 characters in length and does not exceed 20 characters. To protect your account, your passphrase should contain a combination of letters and numbers. The passphrase you create may include spaces but should not contain special characters (for example, +,?, and *). You can associate the same passphrase with multiple submissions.

A passphrase can only be created by an Authorized Official for a submission. 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 only authorized Supports.

Note: You will be responsible for remembering the passphrase and distributing it to only authorized Supports. If you forget the passphrase, you will not be able to access the Section 8(d) Health & Safety Data Reporting submission to print, submit, or make changes.'

New Passphrase:

Confirm New Passphrase:

Cancel

Next

Section 8(d) Health & Safety Data Reporting
Primary Authorized Official
- 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

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: Thu Sep 12 14:13:38 EDT 2013

Next

Section 8(d) Health & Safety Data Reporting
Primary Authorized Official
- General Submission Information
- Contact Information
  - Technical Contact Information
- Chemical Information
  - Chemical Substance Identity of Impurities
- Studies
  - Study Identification
- Submitter Requests

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:

Previous **Next**

Section 8(d) Health & Safety Data Reporting

Primary Authorized Official

- General Submission Information
- Contact Information
  - Technical Contact Information
- Chemical Information
  - Chemical Substance Identity of Impurities
- Studies
  - Study Identification
  - Ongoing Studies
- 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.

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

Previous Next

SEARCH SUBSTANCE REGISTRY SERVICES

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 CA Index Name

1. CASRN: Matches exactly
2. CA Index Name or Other Synonym: Matches Exactly

Search

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: Matches Exactly
2. Generic Name: Matches Exactly

Search

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.

Chemical ID

Unknown: ☐

Accession Number:

CASRN:

PMN Number:

IUPAC Name:

Chemical Name (descriptor):

Chemical Synonym: ✚

Cancel OK

Section 8(d) Health & Safety Data Reporting

Primary Authorized Official

- 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

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

Section 8(d) Health & Safety Data Reporting
Primary Authorized Official
- 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

FULL STUDY REPORT

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

File Name	CBI	Actions

Previous **Next**

CSPP

FULL STUDY REPORT

Browse for the appropriate Full Study Report document.
Note: For security purposes, documents are only saved when there are no validation errors. If there are any validation errors, all documents must be reattached.

CBI: ☑

Document Upload: Browse

Sanitized Document Upload: Browse

Effects:

EndPoints:

OK Cancel

Section 8(d) Health & Safety Data Reporting
Primary Authorized Official
- 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

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**

INITIATED STUDIES

Browse for the appropriate Initiated Studies document.
Note: For security purposes, documents are only saved when there are no validation errors. If there are any validation errors, all documents must be reattached.

CBI: ☑

Document Upload: Browse

Sanitized Document Upload: Browse

Effects:

EndPoints:

Section 8(d) Health & Safety Data Reporting
Primary Authorized Official
- 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

Section 8(d) Health & Safety Data Reporting

Primary Authorized Official

- 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

ONGOING STUDIES

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.

Select: (•) Add Documents ( ) Add Studies

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

File Name	CBI	Actions

Previous **Next**

CSPP

ONGOING STUDIES

Browse for the appropriate Ongoing Studies document.
Note: For security purposes, documents are only saved when there are no validation errors. If there are any validation errors, all documents must be reattached.

CBI: ☑

Document Upload: Browse

Sanitized Document Upload: Browse

Effects:

EndPoints:

OK Cancel

Section 8(d) Health & Safety Data Reporting
Primary Authorized Official
- 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 > Ongoing Studies

ONGOING STUDIES

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.

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

Section 8(d) Health & Safety Data Reporting
Primary Authorized Official
- 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

ROBUST SUMMARY

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

File Name	Actions

Previous **Next**

CSPP

ROBUST SUMMARY

Browse for the appropriate Robust Summary document.
Note: For security purposes, documents are only saved when there are no validation errors. If there are any validation errors, all documents must be reattached.

Document Upload: Browse

Effects:

EndPoints:

OK Cancel

Section 8(d) Health & Safety Data Reporting
Primary Authorized Official
- 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

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**

STUDIES WHICH ARE KNOWN BUT WITHOUT POSSESSION OF COPIES

Browse for the appropriate Studies Which are Known but without Possession of Copies document.
Note: For security purposes, documents are only saved when there are no validation errors. If there are any validation errors, all documents must be reattached.

CBI: ☑

Document Upload: Browse

Sanitized Document Upload: Browse

Effects:

EndPoints:

OK Cancel

Section 8(d) Health & Safety Data Reporting
Primary Authorized Official
- 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

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. 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:
Street address, P.O. box, company name, etc.
Mailing Address 2:
Apartment, suite, etc.
City:
State:
Postal Code:
Country:

Save

Add Study

Previous Next

Section 8(d) Health & Safety Data Reporting
Primary Authorized Official
- 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

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**

STUDIES PREVIOUSLY SENT TO FEDERAL AGENCIES WITHOUT CONFIDENTIALITY CLAIMS

Browse for the appropriate Studies Previously Sent to Federal Agencies without Confidentiality Claims document.
Note: For security purposes, documents are only saved when there are no validation errors. If there are any validation errors, all documents must be reattached.

CBI: ☑

Document Upload: Browse

Sanitized Document Upload: Browse

Effects:

EndPoints:

OK Cancel

Section 8(d) Health & Safety Data Reporting
Primary Authorized Official
- 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

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. 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:

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

Mailing Address 2:

Apartment, suite, etc.

City:

State:

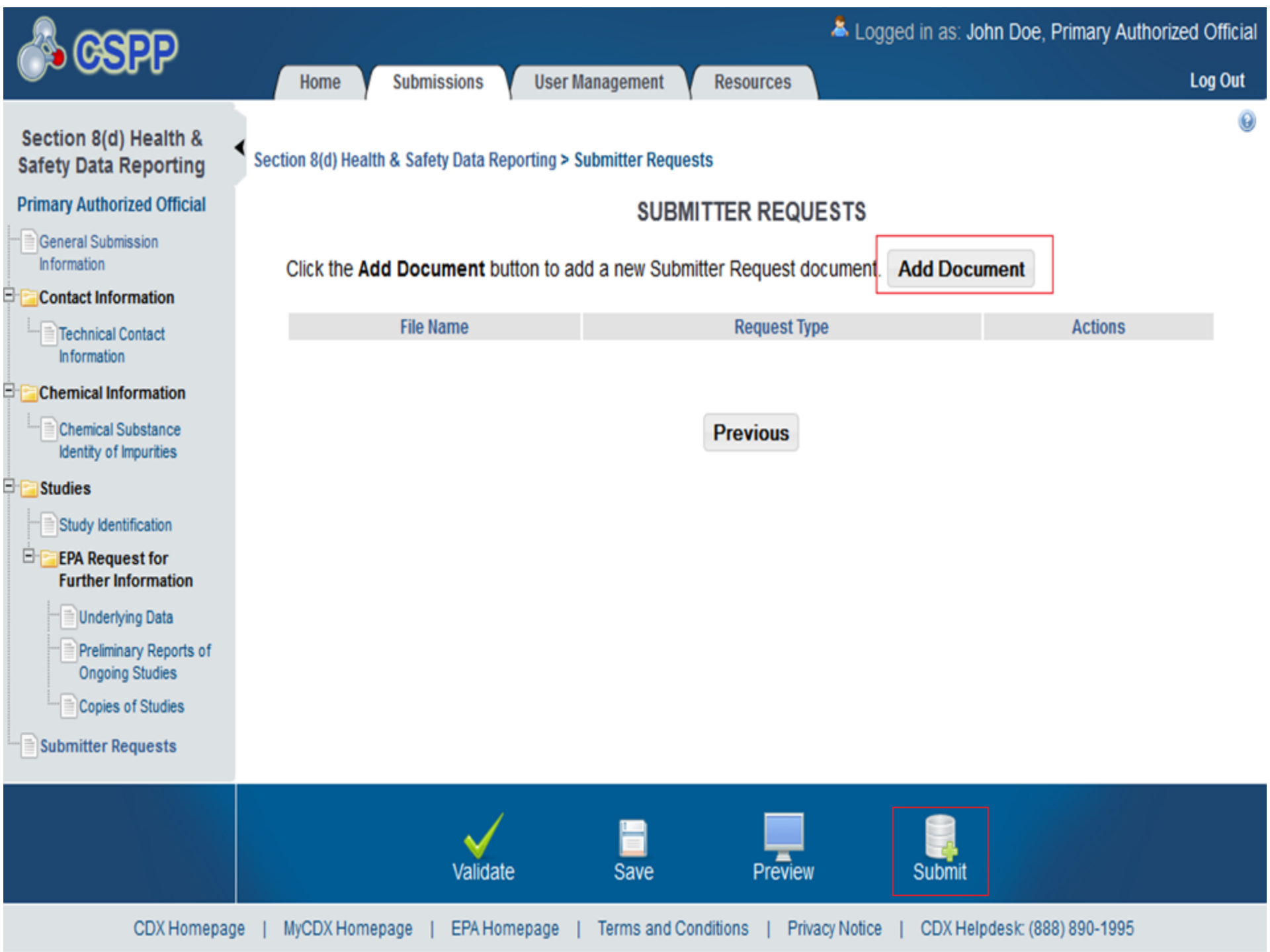
Postal Code:

Country:

Save

Add Study

Previous Next


CSPP
Logged in as: John Doe, Primary Authorized Official
Home
Submissions
User Management
Resources
Log Out
Section 8(d) Health & Safety Data Reporting
Primary Authorized Official
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
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
Submit
CDX Homepage | MyCDX Homepage | EPA Homepage | Terms and Conditions | Privacy Notice | CDX Helpdesk: (888) 890-1995

8(d) User Request

SECTION 8(d) HEALTH & SAFETY DATA REPORTING

- If starting a Section 8(d) Health & Safety Data Reporting submission for the first time in CDX, select a CFR/FRN from the drop-down menu and click the **Start New Submission** button.
- To edit an **In Progress** submission, click the submission alias link in the **Submission Alias** column in the table below.
- To access and edit a form previously Submitted through CDX, unlock the form by clicking the lock icon ( ) and enter your passphrase originally associated with the selected form. All additional changes made to a submission will be submitted as an amendment.
- Click the green arrow icon ( ) to download a copy of record for a submitted form. It may take up to 15 minutes for the copy of record to become available.
- You may delete any submission that has not yet been submitted by clicking the delete icon ( ).

2 items found.

Page 1 of 1

Items per page: 25

Submission Alias	CFR/FRN	CASRN	Status	Last Modified	Submission Date	Copy of Record	Action
Fri Nov 22 10:51:52 EST 2013	75 FR 34076	1336-36-3	Submitted	11/22/2013	11/22/2013		
Thu Nov 21 19:56:06 EST 2013	75 FR 34076	1336-36-3	In Progress	11/21/2013			

Export options: CSV | Excel | XML | PDF

Select Applicable CFR/FRN:

Start New Submission

Logged in as: John Doe, Primary Authorized Official

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

Section 8(d) Health & Safety Data Reporting

Primary Authorized Official

- General Submission Information
- Contact Information
  - Technical Contact Information
- Chemical Information
  - Chemical Substance Identity of Impurities
- Studies
  - Study Identification
- Submitter Requests

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

Section 8(d) Health & Safety Data Reporting

Primary Authorized Official

- 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

UNDERLYING DATA

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

File Name	CBI	Actions

Previous **Next**

UNDERLYING DATA

Browse for the appropriate Underlying Data document.
Note: For security purposes, documents are only saved when there are no validation errors. If there are any validation errors, all documents must be reattached.

CBI: ☑

Document Upload: Browse

Sanitized Document Upload: Browse

Effects:

EndPoints:

OK Cancel

Section 8(d) Health & Safety Data Reporting
Primary Authorized Official
- 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

PRELIMINARY REPORTS OF ONGOING STUDIES

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.

Select: (•) Add Documents ( ) Add Studies

Click the **Add Document** button to add a new Preliminary Reports of Ongoing Studies document. **Add Document**

File Name	CBI	Actions

Previous **Next**

PRELIMINARY REPORTS OF ONGOING STUDIES

Browse for the appropriate Preliminary Reports of Ongoing Studies document.
Note: For security purposes, documents are only saved when there are no validation errors. If there are any validation errors, all documents must be reattached.

CBI: ☑

Document Upload: Browse

Sanitized Document Upload: Browse

Effects:

EndPoints:

OK Cancel

Section 8(d) Health & Safety Data Reporting
Primary Authorized Official
- 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 > Preliminary Reports of Ongoing Studies

PRELIMINARY REPORTS OF ONGOING STUDIES

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.

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

Section 8(d) Health & Safety Data Reporting

Primary Authorized Official

- 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

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**

COPIES OF STUDIES

Browse for the appropriate Copies of Studies document.
Note: For security purposes, documents are only saved when there are no validation errors. If there are any validation errors, all documents must be reattached.

CBI: ☑

Document Upload: Browse

Sanitized Document Upload: Browse

Effects:

EndPoints:

OK Cancel

Section 8(d) Health & Safety Data Reporting
Primary Authorized Official
- 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

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:
Street address, P.O. box, company name, etc.
Mailing Address 2:
Apartment, suite, etc.
City:
State:
Postal Code:
Country:

Save

Add Study

Previous Next

Section 8(a) PAIR Tool

Primary Authorized Official

CHEMICAL INFORMATION SUBMISSION SYSTEM

TSCA Section 8(a) PAIR Reporting

OK

The software includes embedded help files and downloadable user manual to guide you through the 8(a) PAIR submission process.

TSCA 8(a) gives EPA the broad authority to require, by rulemaking, manufacturers (includes importers) and processors of chemical substances to maintain records and/or report such data as EPA may reasonably require to carry out the TSCA mandates.

Section 8(a) regulations can be tailored to meet unique information needs (e.g., via chemical-specific rules) or information can be obtained via use of "model" or standardized reporting rules. One example of a model TSCA Section 8(a) reporting rule is the "Preliminary Assessment Information Rule" (or PAIR).

Paperwork Reduction Act Notice

The information collection requirements contained in this final rule were submitted for OMB approval under PRA, 44 U.S.C. 3501 et seq. The ICR document prepared by EPA, identified under EPA ICR No. 2412.01 and OMB control number 2070-0054, is available in the docket for the proposed rule. The ICR addresses the incremental changes to the five currently approved ICR documents that cover the existing reporting and record keeping programs that are approved under OMB control numbers 2070-0004, 2070-0012, 2070-0033, 2070-0054, and 2070-0156. 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 amended information collection activities contained in this final rule are designed to assist the Agency in meeting its responsibility under TSCA to receive, process, and review reports, data, and other information. As such, responses to the collection of information covered by this ICR would still be mandatory, but with the final rule, respondents would be required to use the CISS reporting tool.

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.

HOME

Forms

Under PAIR, producers and importers of a listed chemical are required to report the following site-specific information:

- Quantity of chemical produced and/or imported
- Amount of chemical lost to the environment during production or importation
- Quantity of enclosed, controlled and open releases of the chemical
- Per release, the number of workers exposed and the number of hours exposed

User Management

Manage the access rights of Supports for each 8(a) PAIR 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(a) PAIR system and provides useful links for further usability instruction.

Authorized Official

An Authorized Official has the ability to create, amend and unlock 8(a) PAIR forms. The Authorized Official may also submit completed forms electronically.

USER MANAGEMENT

The Authorized Official is responsible for restricting a Support's access to select forms by assigning or unassigning them to each Form Alias. The Support can access and edit only those forms for which the Authorized Official has granted access. Select a form alias from the **Form Alias** drop-down menu, and assign a Support to the form by highlighting the individual and clicking the **add** link. To unassign a Support, highlight the individual and click the **remove** link. To highlight and assign or unassign multiple Supports, hold down the **Ctrl** or **Shift** keys on the keyboard and click each Support before moving. You must click the **Save** button after each form alias assignment.

Form Alias:

Form Alias

CASRN:
Code of Federal Regulation:

Assign Users

Unassigned		Assigned
	add >>	
	<< remove	

HOME

Forms

Under 8(a) PAIR, producers, importers, and processors of a listed chemical are required to report the following site-specific information:

- Quantity of chemical produced and/or imported
- Amount of chemical lost to the environment during production or importation
- Quantity of enclosed, controlled and open releases of the chemical
- Per release, the number of workers exposed and the number of hours exposed

User Management

Manage the access rights of Supports for each 8(a) PAIR 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(a) PAIR system and provides useful links for further usability instruction.

Authorized Official

An Authorized Official has the ability to create, amend and unlock 8(a) PAIR forms. The Authorized Official may also submit completed forms electronically.

SECTION 8(a) PAIR

- If starting a Section 8(a) PAIR form for the first time in CDX, select the appropriate CFR from the drop-down menu and click the **Start New Submission** button.
- To edit an **In Progress** form, click the form alias link in the **Form Alias** column in the table below.
- To access and edit a form previously Submitted through CDX, unlock the form by clicking the lock icon ( ) and enter your passphrase originally associated with the selected form. All additional changes made to a submission will be submitted as an amendment.
- Click the green arrow icon ( ) to download a copy of record for a submitted form. It may take up to 15 minutes for the copy of record to become available.
- You may delete any form that has not yet been submitted by clicking the delete icon ( ).

9 items found.

Page 1 of 1

Items Per Page: 25

Form Alias	CASRN	Status	Modify Date	Submission Date	Copy of Record	Action
Mon Sep 09 13:57:34 EDT 2013	79-94-7	Submitted	09/09/2013	09/09/2013		
Mon Sep 16 11:42:12 EDT 2013	37853-61-5	Submitted	09/16/2013	09/16/2013		
Mon Sep 23 11:36:07 EDT 2013		In Progress	09/23/2013			
Mon Sep 23 11:36:56 EDT 2013		In Progress	09/23/2013			
Mon Sep 23 11:41:12 EDT 2013		In Progress	09/23/2013			
Mon Sep 23 11:42:29 EDT 2013		In Progress	09/23/2013			
Mon Sep 23 11:46:32 EDT 2013JD	79-94-7	Submitted	09/23/2013	09/23/2013		
Mon Sep 23 11:50:09 EDT 2013 JD	3389-71-7	Submitted	09/23/2013	09/23/2013		
Thu Sep 19 20:13:49 EDT 2013		In Progress	09/19/2013			

Export options: CSV | Excel | XML | PDF

Select Applicable CFR:

Start New Submission

Section 8(a)

CREATE PASSPHRASE

Please create a passphrase that is at least 8 characters in length and does not exceed 20 characters. To protect your account, your passphrase should contain a combination of letters and numbers. The passphrase you create may include spaces but should not contain special characters (for example, +,?, and *). You can associate the same passphrase with multiple submissions.

New Passphrase:
Confirm New Passphrase:

A passphrase can only be created by an Authorized Official for an individual submission. 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 only authorized Supports.

Note: You will be responsible for remembering the passphrase and distributing it to only authorized Supports for the submission type. If you forget the passphrase, you will not be able to access the Section 8(a) PAIR submission to print, submit, or make changes.

Cancel Next

Section 8(a) PAIR Reporting
Primary Authorized Official
40 CFR 704.43
- General Submission Information
- Contact Information
  - Technical Contact Information
- Chemical Information
  - Plant Site Physical Location
  - Mailing Address
  - Preliminary Assessment Information - Part A
  - Preliminary Assessment Information - Part B

GENERAL SUBMISSION INFORMATION

You have chosen to report under **40 CFR 704.43** . Based on this selection, all data entered in this form should pertain to the following chemical selected from the drop-down menu below.

Please select, or begin typing, a CASRN in the drop-down menu below:

CASRN:

The form alias is an optional field that changes the submission name on the **Forms 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 form alias may be changed at any time.

Form Alias: Mon Sep 23 11:41:12 EDT 2013

Next

Section 8(a) PAIR Reporting
Primary Authorized Official
40 CFR 704.43
- General Submission Information
- Contact Information
  - Technical Contact Information
- Chemical Information
  - Plant Site Physical Location
  - Mailing Address
  - Preliminary Assessment Information - Part A
  - Preliminary Assessment Information - Part B

TECHNICAL CONTACT INFORMATION

Identify the technical contact who is capable of answering questions related to the chemical(s) submitted to EPA within this form. 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:

Previous **Next**

Section 8(a) PAIR Reporting
Primary Authorized Official
40 CFR 704.43
- General Submission Information
- Contact Information
  - Technical Contact Information
  - Submitting On Behalf Of Company
- Chemical Information
  - Plant Site Physical Location
  - Mailing Address
  - Preliminary Assessment Information - Part A
  - Preliminary Assessment Information - Part B

Section 8(a) PAIR Reporting > Contact Information > Submitting On Behalf Of Company

SUBMITTING ON BEHALF OF COMPANY

Please 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

Section 8(a) PAIR Reporting
Primary Authorized Official
40 CFR 704.43
General Submission Information
Contact Information
Technical Contact Information
Submitting On Behalf Of Company
Chemical Information
Plant Site Physical Location
Mailing Address
Preliminary Assessment Information - Part A
Preliminary Assessment Information - Part B

PLANT SITE PHYSICAL LOCATION

Enter the name, physical location address, and Dun & Bradstreet Number of the plant site for which the data are reported.

CBI:

Plant Site:

Company:

Address 1:

Address 2:

City:

State:

Postal Code:

Dun & Bradstreet:

Previous Next

Section 8(a) PAIR Reporting
Primary Authorized Official
40 CFR 704.43
General Submission Information
Contact Information
Technical Contact Information
Submitting On Behalf Of Company
Chemical Information
Plant Site Physical Location
Mailing Address
Preliminary Assessment Information - Part A
Preliminary Assessment Information - Part B

MAILING ADDRESS

Select the appropriate radio button to show whether the plant site or corporate headquarters is submitting this form. Enter the corresponding name and mailing address. Click the Copy button to import information from the Plant Site Physical Location.

- ○ Plant Site
- ○ Corporate Headquarters

CBI: ☐

Company:

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

Address 2:
Apartment, suite, etc.

City:

State:

Postal Code:

Previous Next

Section 8(a) PAIR Reporting
Primary Authorized Official
40 CFR 704.43
- General Submission Information
- Contact Information
  - Technical Contact Information
  - Submitting On Behalf Of Company
- Chemical Information
  - Plant Site Physical Location
  - Mailing Address
  - Preliminary Assessment Information - Part A
  - Preliminary Assessment Information - Part B

Section 8(a) PAIR Reporting > 40 CFR 704.43 > Chemical Information > Preliminary Assessment Information - Part A

PRELIMINARY ASSESSMENT INFORMATION

Part A - Plant Site Activities

Please complete all fields below. Information in part A must be your best estimate from readily obtainable data.

☐ This company is not involved in any manufacturing activity and imports a chemical at one site and processes it at another facility.

					CBI
1.	Total Quantity Imported		kg		☐
2.	Quantity manufactured for sale or use		kg		☐
3a.	Quantity lost to the environment		kg	± %	☐
3b.	Quantity in wastes treated to destroy the chemical		kg	± %	☐
3c.	Quantity in wastes not treated to destroy the chemical		kg	± %	☐
3d.	Quantity lost during manufacture		kg		☐

4. Manufacture of the Chemical CBI: ☐

Process Category	Quantity (kg)	Total Quantity (kg)	Total Workers	Total Worker-Hours
Enclosed				
Controlled Release				
Open				

5. On-Site Use as a Reactant CBI: ☐

Section 8(a) PAIR Reporting
Primary Authorized Official
40 CFR 704.43
- General Submission Information
- Contact Information
  - Technical Contact Information
  - Submitting On Behalf Of Company
- Chemical Information
  - Plant Site Physical Location
  - Mailing Address
  - Preliminary Assessment Information - Part A
  - Preliminary Assessment Information - Part B

5. On-Site Use as a Reactant CBI:

Process Category	Quantity (kg)	Total Quantity (kg)	Total Workers	Total Worker-Hours
Enclosed				
Controlled Release				
Open				

6. On-Site Nonreactant Use of the Chemical Substance CBI:

Process Category	Quantity (kg)	Total Quantity (kg)	Total Workers	Total Worker-Hours
Enclosed				
Controlled Release				
Open				

7. On-Site Preparation of Products CBI:

Process Category	Quantity (kg)	Total Quantity (kg)	Total Workers	Total Worker-Hours
Enclosed				
Controlled Release				
Open				

8. Manufacturer's Products CBI:

a. Products for Export ____ kg

	Domestic Industrial Products	Domestic Consumer Products
Chemical or Mixture	b. ____ kg	e. ____ kg
Article with Some Release	c. ____ kg	f. ____ kg
Article with No Release	d. ____ kg	g. ____ kg

Previous Next

Section 8(a) PAIR Reporting
Primary Authorized Official
40 CFR 704.43
- General Submission Information
- Contact Information
  - Technical Contact Information
  - Submitting On Behalf Of Company
- Chemical Information
  - Plant Site Physical Location
  - Mailing Address
  - Preliminary Assessment Information - Part A
  - Preliminary Assessment Information - Part B

Section 8(a) PAIR Reporting > 40 CFR 704.43 > Chemical Information > Preliminary Assessment Information - Part B

PRELIMINARY ASSESSMENT INFORMATION

Part B - Chemical Substance Processing by Customers

Please complete all fields below. Information in part B must be accurate to within ±50%.

9. Customer Uses and Products CBI: ☐

a. Products for Export _____ kg

b. Quantity of Chemical Consumed as Reactant _____ kg

	Domestic Industrial Products		Domestic Consumer Products	
Chemical or Mixture	c.	kg	f.	kg
Article with Some Release	d.	kg	g.	kg
Article with No Release	e.	kg	h.	kg

Specify Unknown if you do not know within ±50%

i. Unknown Customer Uses _____ kg

10. Customer Process Categories CBI: ☐

Specify Unknown if you do not know within ±50%

a. Enclosed Processes _____ kg

b. Controlled Release Processes _____ kg

c. Open Processes _____ kg

d. Unknown _____ kg

Previous

Section 8(a) PAIR Reporting
Primary Authorized Official
40 CFR 704.43
- General Submission Information
- Contact Information
  - Technical Contact Information
  - Submitting On Behalf Of Company
- Chemical Information
  - Plant Site Physical Location
  - Mailing Address
  - Preliminary Assessment Information - Part A
  - Preliminary Assessment Information - Part B

UPLOAD XML

Click the Upload XML File button below and select the XML file you have populated with information.

Upload XML file

Warning: When importing an XML file into the current form, if XML validation is passed, all form contents will be overwritten with the information stored within the imported XML file.

Export XML

Section 8(a) 40 CFR 766 Dibenzodioxins/Dibenzofurans

- If starting a Section 8(a) PAIR form for the first time in CDX, select the appropriate CFR from the drop-down menu and click the **Start New Submission** button.
- To edit an **In Progress** form, click the form alias link in the **Form Alias** column in the table below.
- To access and edit a form previously Submitted through CDX, unlock the form by clicking the lock icon ( ) and enter your passphrase originally associated with the selected form. All additional changes made to a submission will be submitted as an amendment.
- Click the green arrow icon ( ) to download a copy of record for a submitted form. It may take up to 15 minutes for the copy of record to become available.
- You may delete any form that has not yet been submitted by clicking the delete icon ( ).

Page 1 of 1

3 items found.

Items Per Page: 25

Form Alias	CASRN	Status	Modify Date	Submission Date	Copy of Record	Action
Fri Nov 29 13:39:35 EST 2013		In Progress	11/29/2013			
Fri Nov 29 13:41:24 EST 2013		In Progress	11/29/2013			
Fri Nov 29 13:42:07 EST 2013		In Progress	11/29/2013			

Export options: CSV | Excel | XML | PDF

Select Applicable CFR: 40 CFR 766 Dibenzodioxins/Dibenzofurans

Start New Submission

CREATE PASSPHRASE

Please create a passphrase that is at least 8 characters in length. To better protect your form, your passphrase should contain a combination of letters numbers. Your passphrase may include spaces, but should not contain special characters (for example, + and *).

As an Authorized Official, you are responsible for remembering your passphrase and distributing it to only authorized individuals. 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 only authorized Supports.

Note: If you lose or forget your passphrase, you will not be able to access your Section 4 Submission to print, submit, or make changes. You will need to complete a new Section 4 Submission and create a new passphrase for the submission. For security reasons, the system administrator will not have access to your passphrase and will not be able to retrieve it or reset it.

New Passphrase:
Confirm New Passphrase:

Cancel Next

Section 8(a) PAIR Reporting
Primary Authorized Official
40 CFR 766
General Submission Information
Contact Information
Technical Contact Information
Chemical Information
Document Management

GENERAL SUBMISSION INFORMATION

You have chosen to report under **40 CFR 766** . Based on this selection, all data entered in this form should pertain to the following chemical selected from the drop-down menu below.

Please select, or begin typing, a CASRN in the drop-down menu below:

CASRN: 118-75-2

Chemical Name: **2,3,5,6-Tetrachloro-2,5-cyclohexadiene-1,4-dione**

The form alias is an optional field that changes the submission name on the **Forms 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 form alias may be changed at any time.

Form Alias: Fri Nov 29 13:52:38 EST 2013

Next

Section 8(a) PAIR Reporting
Primary Authorized Official
40 CFR 766
General Submission Information
Contact Information
Technical Contact Information
Chemical Information
Document Management

TECHNICAL CONTACT INFORMATION

Identify the technical contact who is capable of answering questions related to the chemical(s) submitted to EPA within this form. 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:

Previous Next

Section 8(a) PAIR Reporting
Primary Authorized Official

- 40 CFR 766
  - General Submission Information
  - Contact Information
    - Technical Contact Information
    - Submitting On Behalf Of Company
  - Chemical Information
    - Document Management

DOCUMENT MANAGEMENT

Upload the corresponding 40 CFR 766: 8(a) PAIR Reporting document(s) by clicking on the **Add Document** button below.

Add Document

File Name	Attachment Date	CBI	Action

Previous

DOCUMENT MANAGEMENT

Note: For security purposes, documents are only saved when there are no validation errors. If there are any validation errors, all documents must be reattached.

CBI: ☑

Browse for the appropriate document.

Original Document: Browse

Browse for the appropriate sanitized document.

Sanitized Document: Browse (optional)

OK Cancel

Section 8(a) PAIR Reporting
Primary Authorized Official

- 40 CFR 766
  - General Submission Information
  - Contact Information
    - Technical Contact Information
    - Submitting On Behalf Of Company
  - Chemical Information
    - Document Management

DOCUMENT MANAGEMENT

Upload the corresponding 40 CFR 766: 8(a) PAIR Reporting document(s) by clicking on the **Add Document** button below.

Add Document

File Name	Attachment Date	CBI	Action
Submit blank form.docx	12/05/2013	N	✖

Previous

Resources

- Resources
  - Resources screen within Section 8(d) and Section 8(a) web application provides useful links and user guides

- Contacts
  - TSCA Hotline: 202-564-3001, or TSCA-Hotline@epamail.epa.gov
  - CDX Helpdesk: 888-890-1995, or helpdesk@epacdx.net

- Industry Beta Testing (12/16-12/20)
  - Email eTSCAReporting@epa.gov to participate

- The slides and audio will be made available online
  - http://www.epa.gov/oppt/chemtest/ereporting/

Questions

Press *1 to ask a question