



Section 4 User Guide – Primary Authorized Official

Environmental Protection Agency

Office of Pollution Prevention and Toxics

Manage Toxic Substances

Table of Contents

1	Introduction	8
1.1	Labeling Confidential Information	8
2	System Requirements	11
2.1	Supported Browsers	11
2.2	Screen Resolution	11
3	Primary Authorized Official Functions	12
3.1	Log into Section 4 application	12
3.2	Home Screen Navigation	13
3.3	User Management	15
3.4	Create New Submission	17
3.4.1	Section 4 Test Rules, ECAs, MOUs, and 40 CFR 766 Submissions Screen	17
3.4.2	Create Passphrase Screen	20
3.4.3	Enter Passphrase Screen	21
3.4.4	Navigation Tree	21
3.4.5	Action Bar	26
4	Test Rules	29
4.1	Submission Information Type Screen	29
4.2	Technical Contact Information Screen	30
4.3	Submitting on Behalf of Consortium Screen	34
4.4	Submitting on Behalf of Company Screen	36
4.5	Letter of Intent	38
4.5.1	Chemical Identification and Test Rules Screen	39
4.5.2	Sponsoring Firms Screen	42
4.5.3	Additional Information and Submitter Requests Screen	44
4.6	Study Plan	51
4.6.1	Document Management Screen	51
4.6.2	Additional Information and Submitter Requests Screen	54
4.7	Test Rules Substantiation	61
4.7.1	Test Rules Substantiation Part 1 Screen	61
4.7.2	Test Rules Substantiation Part 2 Screen	64
4.8	Results	66
4.8.1	Document Management Screen	67
4.9	Optional Substantiation	70

4.9.1	Optional Document Management Screen	70
5	Enforceable Consent Agreements (ECAs).....	73
5.1	Submission Information Type Screen	73
5.2	Technical Contact Information Screen	74
5.3	Submitting on Behalf of Company Screen	78
5.4	Study Plans	80
5.4.1	Principal Test Sponsor Screen.....	80
5.4.2	Principal Sponsor Organization Screen	81
5.4.3	Testing Facilities Screen.....	83
5.4.4	Study Professionals Screen	87
5.5	Results	90
5.5.1	Document Management Screen	90
5.6	ECA Additional Information Screen.....	94
6	Memorandum of Understanding (MOU).....	98
6.1	Submission Information Type Screen	98
6.2	Technical Contact Information Screen	99
6.3	Submitting on Behalf of Company Screen	103
6.4	Study Plans & Conduct of Testing	104
6.4.1	Principal Test Sponsor Screen.....	104
6.4.2	Principal Sponsor Organization Screen	106
6.4.3	Testing Facilities Screen.....	108
6.4.4	Study Professionals Screen	111
6.5	Results	114
6.5.1	Document Management Screen	114
6.6	MOU Additional Information Screen.....	116
7	40 CFR 766 Dibenzodioxins/Dibenzofurans	119
7.1	Submission Information Type Screen	119
7.2	Technical Contact Information Screen	121
7.3	Submitting on Behalf of Consortium Screen	124
7.4	Submitting on Behalf of Company Screen	126
7.4.1	Chemical Identification Screen.....	127
7.5	Letter of Intent	128
7.5.1	Sponsoring Firms Screen	128
7.6	Protocol	130
7.6.1	Protocol Document Management Screen.....	130

7.7	Results	133
7.7.1	Document Management Screen	133
7.8	User Requests.....	136
7.8.1	Waiver Screen	136
7.8.2	Exclusion Screen	139
7.8.3	Exemption Screen	142
8	Validate	145
9	Submit to EPA via CDX	147
9.1	Submitting Official Information Screen.....	147
9.2	Submission Process: Validation Screen	148
9.3	Cross-Media Electronic Reporting Regulation (CROMERR) Certification Screen	149
9.4	Cross-Media Electronic Reporting Regulation (CROMERR) Login Screen	150
9.5	Cross-Media Electronic Reporting Regulation (CROMERR) Security Question Screen.....	151
9.6	Submit to CDX Screen.....	152
9.7	Cross-Media Electronic Reporting Regulation (CROMERR) Submission Screen.....	153
10	Download Copy of Record.....	155
11	Create an Amendment	158
12	Resources	161
Appendix A. Definitions, Acronyms, and Abbreviations		162
Appendix B. Submission Types, Effects, and Endpoints		163

List of Exhibits

Exhibit 3-1 Chemical Information Submission System Screen	13
Exhibit 3-2: Section 4 Home Screen.....	15
Exhibit 3-3: User Management Screen.....	16
Exhibit 3-4: Section 4 Test Rules, ECAs, MOUs, and 40 CFR 766 Submissions Screen	19
Exhibit 3-5: Create Passphrase Screen.....	20
Exhibit 3-6 Enter Passphrase Screen.....	21
Exhibit 3-7: Navigation Prompt.....	22
Exhibit 3-8: Test Rules Navigation Tree	23
Exhibit 3-9: ECA Navigation Tree.....	24
Exhibit 3-10: MOU Navigation Tree.....	25
Exhibit 3-11: 40 CFR 766 Navigation Tree	26
Exhibit 3-12: Action Bar – Validate	27
Exhibit 3-13: Action Bar – Save.....	27
Exhibit 3-14: Action Bar – Preview	27
Exhibit 3-15: Action Bar – Submit.....	27
Exhibit 3-16: Action Bar – Help Links	28
Exhibit 4-1: Submission Information Type Screen	30
Exhibit 4-2: Technical Contact Information Screen.....	33
Exhibit 4-3: Submitting on Behalf of Consortium Screen	36
Exhibit 4-4: Submitting on Behalf of Company Screen	38
Exhibit 4-5: Chemical Identification and Test Rules Screen.....	40
Exhibit 4-6: Chemical Management Pop-Up Window	41
Exhibit 4-7: Chemical Identification and Test Rules Screen (Populated)	42
Exhibit 4-8: Sponsoring Firms Screen	44
Exhibit 4-9: Additional Information and Submitter Requests Screen.....	47
Exhibit 4-10: Amendment to the Study Plan Pop-Up Window.....	48
Exhibit 4-11: Exemption Application Pop-Up Window.....	48
Exhibit 4-12 Extension of Time to Submit Study Plan Pop-Up Window	49
Exhibit 4-13: Modification of Test Standards or Schedules during Conduct of Test Pop-Up Window.....	49
Exhibit 4-14: Submission of Equivalence Data Pop-Up Window.....	50
Exhibit 4-15: Appeal of Denial of Exemption Application Pop-Up Window.....	50
Exhibit 4-16 Document Management Screen	52
Exhibit 4-17 Document Management Pop-Up Window.....	54
Exhibit 4-18 Additional Information and Submitter Requests Screen.....	57
Exhibit 4-19 Amendment to the Study Plan Pop-Up Window.....	58
Exhibit 4-20 Exemption Application Pop-Up Window.....	58
Exhibit 4-21 Extension of Time to Submit Study Plan Pop-Up Window	59

Exhibit 4-22 Modification of Test Standards or Schedules during Conduct of Test Pop-Up Window.....	59
Exhibit 4-23 Submission of Equivalence Data Pop-Up Window.....	60
Exhibit 4-24 Appeal of Denial of Exemption Application Pop-Up Window.....	61
Exhibit 4-25: Test Rules Substantiation Part 1 Screen	63
Exhibit 4-26: Test Rules Substantiation Part 2 Screen (Scroll 1)	65
Exhibit 4-27: Test Rules Substantiation Part 2 Screen (Scroll 2)	66
Exhibit 4-28 Results Document Management Screen	68
Exhibit 4-29 Results Document Management Pop-Up Window	70
Exhibit 4-30: Optional Document Management Screen.....	71
Exhibit 4-31: Optional Substantiation Document Management Pop-Up Window	72
Exhibit 5-1 Submission Information Type Screen	74
Exhibit 5-2 Technical Contact Information Screen.....	77
Exhibit 5-3 Submitting on Behalf of Company Screen	79
Exhibit 5-4 Principal Test Sponsor Screen	81
Exhibit 5-5 Principal Sponsor Organization Screen.....	83
Exhibit 5-6 Testing Facilities Screen (Scroll 1)	86
Exhibit 5-7: Testing Facilities Screen (Scroll 2)	87
Exhibit 5-8 Study Professionals Screen (Scroll 1)	89
Exhibit 5-9: Study Professionals Screen (Scroll 2)	90
Exhibit 5-10 Document Management Screen	92
Exhibit 5-11 Document Management Pop-Up Window.....	94
Exhibit 5-12 ECA Additional Information Screen	96
Exhibit 5-13 Amendment to the Study Plan Pop-Up Window.....	96
Exhibit 5-14 Modification of ECAs Pop-Up Window.....	97
Exhibit 6-1: Submission Information Type Screen.....	99
Exhibit 6-2 Technical Contact Information Screen.....	102
Exhibit 6-3 Submitting on Behalf of Company Screen	104
Exhibit 6-4 Principal Test Sponsor Screen	106
Exhibit 6-5 Principal Sponsor Organization Screen	108
Exhibit 6-6 Testing Facilities Screen (Scroll 1)	110
Exhibit 6-7: Testing Facility Screen (Scroll 2)	111
Exhibit 6-8 Study Professionals Screen.....	113
Exhibit 6-9 Document Management Screen	115
Exhibit 6-10 Document Management Pop-Up Window.....	116
Exhibit 6-11 MOU Additional Information Screen	117
Exhibit 6-12 Amendment to the Study Plan Pop-Up Window.....	118
Exhibit 6-13 Modification of MOUs Pop-Up Window.....	118
Exhibit 7-1 Submission Information Type Screen	120

Exhibit 7-2 Technical Contact Information Screen.....	123
Exhibit 7-3 Submitting on Behalf of Consortium Screen	125
Exhibit 7-4 Submitting on Behalf of Company Screen	127
Exhibit 7-5 Chemical Identification Screen	128
Exhibit 7-6 Sponsoring Firms Screen	130
Exhibit 7-7 Document Management Screen	132
Exhibit 7-8 Protocol Document Management Pop-Up Window.....	133
Exhibit 7-9 Document Management Screen	135
Exhibit 7-10 Document Management Pop-Up Window.....	136
Exhibit 7-11 Waiver Screen.....	138
Exhibit 7-12 Waiver Document Management Pop-Up Window.....	139
Exhibit 7-13 Exclusion Screen.....	141
Exhibit 7-14 Exclusion Document Management Pop-Up Window.....	142
Exhibit 7-15 Exemption Screen	143
Exhibit 7-16 Exemption Document Management Pop-Up Window	144
Exhibit 8-1 Section 4 Validation Pop-Up Window	146
Exhibit 9-1 Submitting Official Information Screen.....	147
Exhibit 9-2 Submission Process: Validation Screen	148
Exhibit 9-3 Submission Process: PDF Generation Screen	149
Exhibit 9-4 Cross-Media Electronic Reporting Regulation (CROMERR) Certification Screen ..	150
Exhibit 9-5 Cross-Media Electronic Reporting Regulation (CROMERR) Login Screen	151
Exhibit 9-6 Cross-Media Electronic Reporting Regulation (CROMERR) Security Question Screen	152
Exhibit 9-7 Submit to CDX Screen	153
Exhibit 9-8 Cross-Media Electronic Reporting Regulation (CROMERR) Submission Screen ..	154
Exhibit 10-1 Download Copy of Record Screen.....	156
Exhibit 10-2 Download Prompt Pop-Up Window	157
Exhibit 11-1 Section 4 Test Rules, ECAs, MOUs, and 40 CFR 766 Submissions – Amendment	159
Exhibit 11-2 Unlock Prompt Message	160
Exhibit 12-1 Resources Screen.....	161

1 Introduction

This document represents the user guide for the Office of Pollution Prevention and Toxics (OPPT) Section 4 application submission process. This document is the user guide for the Primary Authorized Official (AO) user of the Section 4 tool.

The Primary AO is the person legally responsible for the submission of Section 4 Test Rules, ECA, MOU, and 40 CFR 766 for the reporting company. The AO, typically, is a senior official for the reporting company and may be the supervisor of the person (or persons) completing the form.

As a Primary AO, you can create a new form. You can also assign Supports to edit and complete forms on your behalf. You are responsible for submitting all required information and are legally responsible for the information submitted to EPA within each Section 4 Test Rules, ECA, MOU, and 40 CFR 766 submission.

General information on Section 4 is available at <http://www.epa.gov/oppt/chemtest/ereporting/index.html>. Information on CDX registration, including user roles that apply to completing a Section 4 is available at <http://cdx.epa.gov/About/UserGuide>

Toxic Substances Control Act (TSCA) regulations require industry to provide the Environmental Protection Agency (EPA) with documentation regarding chemical production, manufacture, distribution, use, disposal, import, and export. As a result, the implementation of TSCA has generated a large volume of documents submitted to the Agency by the regulated community.

The Section 4 application is the electronic, web-based tool provided by the Environmental Protection Agency (EPA) for the submission of Section 4 data. As a Primary AO, you can create or modify a Section 4 submission as well as submit it to EPA via the Central Data Exchange (CDX). You can also create Section 4 submission amendments and download the Copy of Record (CoR).

For questions concerning the Toxic Substances Control Act (TSCA) Section 4 software requirements, please contact the CDX Help Desk at helpdesk@epacdx.net or call 1-888-890-1995 between the hours of 8am – 6pm Eastern Standard Time (EST). For TSCA Section 4 requirements, please contact the TSCA Hotline at TSCA-Hotline@epamail.epa.gov or call 202-554-1404

Please note: Any person who submits a positive test result for a specific chemical substance listed under section 766.25 must submit to EPA, no later than 90 days after the date of submission, a completed 7710-51 form. The form can be downloaded from the following link: <http://www.epa.gov/oppt/chemtest/pubs/7710-51.pdf>.

1.1 Labeling Confidential Information

This user guide presents instructions and screenshots for the ‘Section 4 Test Rules, ECA, MOU, and 40 CFR 766 submission’ electronic application as if you were logged in as the Primary AO. As the Primary AO you are able to create, modify, delete, and submit a Section 4 submission. The user may claim certain information as Confidential Business Information (CBI). If the data is claimed as CBI, it will be treated in accordance with Agency confidentiality regulations at 40 CFR part 2, subpart B.

This TSCA Section 4 Test Rules, ECA, MOU, and 40 CFR 766 submission electronic application is optional for the production and submission of these filings. The paper option for filing will continue to be allowed.

Potential users may find that the use of this application saves time, burden and, because it is in an electronic format, provides easy access to reporting forms. Potential users are reminded to carefully read and follow application instructions. Persons who wish to use this application must address all provisions in the application, including those relating to indexing information (i.e., company name, address, chemical name etc.) and those related to CBI submissions, including the requirement for the submission of sanitized or redacted filings.

Please note that certain CBI claims will not be accepted if you choose to use this electronic application for TSCA Section 4 Test Rules, ECA, MOU, and 40 CFR 766 submission. For example, chemical name and/or chemical identity, if the chemical is listed on the public portion of the TSCA Inventory, may not be claimed as CBI in a submission using this application.

If the user believes that these or other data elements should be treated as CBI, then the user may use the paper option to file the Section 4 submission form and assert CBI claims.

If the user claims an uploaded document as CBI, a sanitized document must be provided.

The following fields can be claimed as CBI within the ‘Section 4 Test Rules, ECA, MOU, and 40 CFR 766’ submission application:

- Test Rules:
 - Technical Contact Information
 - Submitting on Behalf of Consortium information
 - Submitting on Behalf of Company information
 - A single chemical substance and associated chemical test(s)
 - Sponsoring Firm information
 - Study Plan document(s)
 - Submission of Equivalence Data document(s)
 - Result document(s)
 - Official Submitter Information
- ECA:
 - Technical Contact Information
 - Submitting on Behalf of Company information
 - Principal Test Sponsor information
 - Principal Sponsor Organization information
 - Testing facility’s contact role information
 - Study Professional information
 - Results document(s)

- Official Submitter Information
- MOU:
 - Technical Contact Information
 - Submitting on Behalf of Company information
 - Principal Test Sponsor information
 - Principal Sponsor Organization information
 - Testing facility's contact role information
 - Study Professional information
 - Results document(s)
 - Official Submitter Information
- 40 CFR 766:
 - Technical Contact Information
 - Submitting on Behalf of Consortium information
 - Submitting on Behalf of Company information
 - A single chemical substance
 - Sponsoring Firm
 - Protocol document(s)
 - Results document(s)
 - Waiver document(s)
 - Exclusion document(s)
 - Exemption document(s)
 - Official Submitter Information

2 System Requirements

To use the Section 4 application the following are required:

- An e-mail account
- A supported web browser (see Section **Error! Reference source not found.** below) with Java Script enabled
- Internet access
- Adobe Acrobat Reader 5.0 or higher
- CDX username and password

2.1 Supported Browsers

One of the following supported browsers is required to access the Section 4 application:

- Internet Explorer 7 or above
 - Go to the following link to download:
<http://windows.microsoft.com/en-US/internet-explorer/downloads/ie>
- Mozilla Firefox 3.5 or above
 - Go to the following link to download:
<http://www.mozilla.com/en-US/firefox/all-older.html>
- Safari 4 or above
 - Go to the following link to download:
<http://support.apple.com/kb/dl877>
- Google Chrome
 - Go to the following link to download:
<http://www.google.com/chrome>

2.2 Screen Resolution

- Screen resolution should be set to 1024 x 768 or greater.

3 Primary Authorized Official Functions

This section describes how to:

- Access the application
- Navigate the Section 4 ‘Home’ screen
- Assign Support users to complete a submission
- Start and complete a Section 4 submission
- Submit to EPA via CDX
- Download the Copy of Record
- Create an amendment

The Primary AO is the person legally responsible for the submission of Section 4 data for the reporting company. The Primary AO is typically a senior official for the reporting company and may be the supervisor of the person (or persons) completing the submission.

As a Primary AO, you can create a new submission and assign Supports to edit and complete the submission on your behalf. You are responsible for submitting all required information and are legally responsible for the information submitted to EPA within each Section 4 submission.

You can save a submission at any point during the data entry process. The save functionality will allow you to return to that same submission at any point in the future. You can print the submission document at any point; however, the ‘Not for Submission’ watermark will be placed on the portable document format (PDF) each time the submission is printed.

3.1 Log into Section 4 application

After you create an account in CDX and click the ‘Primary Authorized Official’ link, you will be taken to the ‘Chemical Information Submission System’ screen.

Note: Information on CDX registration, including user roles that apply to completing a Section 4 is available at <http://cdx.epa.gov/About/UserGuide>.

Exhibit 3-1 shows a screen capture of the ‘Chemical Information Submission System’ screen:

Exhibit 3-1 Chemical Information Submission System Screen



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

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>.

Navigation: Select 'TSCA Section 4' from the drop-down menu and click the 'OK' button to navigate to the Section 4 'Home' screen.

3.2 Home Screen Navigation

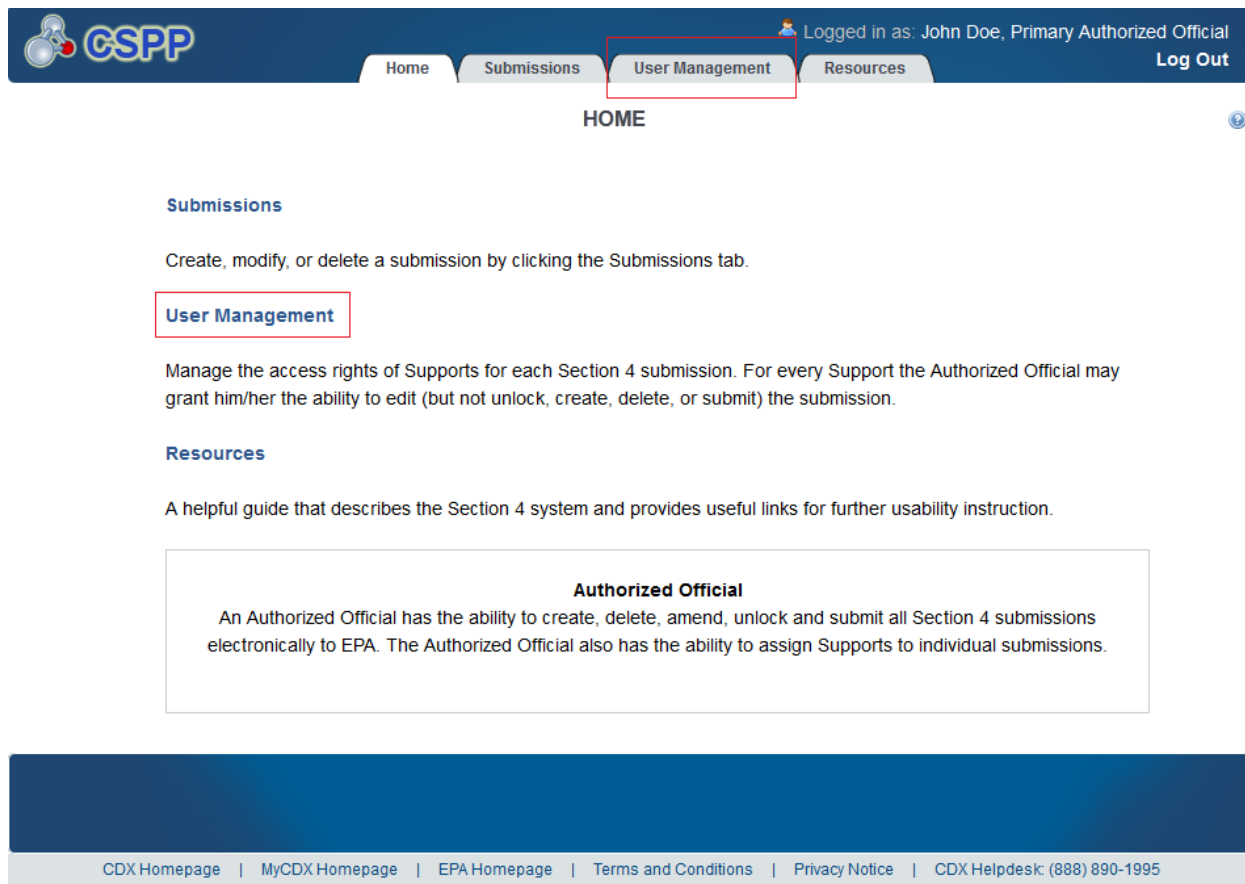
You can access the Section 4 'Home' screen by selecting the 'TSCA Section 4' submission type from the 'Chemical Information Submission System' screen drop-down menu and clicking the 'OK' button.

The Section 4 ‘Home’ screen is the first screen within the Section 4 application. It provides you with links and tabs to access the ‘Section 4 Test Rules, ECAs, MOUs, and 40 CFR 766,’ ‘User Management,’ and ‘Resources’ screens as shown in Exhibit 3-3, Exhibit 3-4, and Exhibit 12-1. To navigate to any of these screens, click the screen link (located in the middle of the Section 4 ‘Home’ screen highlighted in blue), or click the screen tab (located within the application header). The links and tabs provide the same functionality.

- **Submissions:** Click the ‘Submissions’ link or tab to navigate to the ‘Section 4 Test Rules, ECAs, MOUs, and 40 CFR 766’ submissions screen. You can create or modify a Section 4 submission using the ‘Section 4 Test Rules, ECAs, MOUs, and 40 CFR 766’ submissions screen. For additional details about the ‘Section 4 Test Rules, ECAs, MOUs, and 40 CFR 766’ submissions screen, please refer to **Section 3.4.1**.
- **User Management:** Click the ‘User Management’ link or tab to navigate to the ‘User Management’ screen. You can manage the access rights of Supports per submission using the ‘User Management’ screen. For additional details about the ‘User Management’ screen, please refer to **Section 3.3**.
- **Resources:** Click the ‘Resources’ link or tab to navigate to the ‘Resources’ screen. You can view the Section 4 user guide, as well as useful links for further usability instructions on the ‘Resources’ screen. For additional details about the ‘Resources’ screen, please refer to **Section 12**.

Exhibit 3-2 shows a screen capture for the Section 4 ‘Home’ screen:

Exhibit 3-2: Section 4 Home Screen



Navigation: Click the ‘User Management’ link or tab to access the ‘User Management’ screen and assign Supports.

3.3 User Management

You can access the ‘User Management’ screen by clicking the ‘User Management’ link or tab from the Section 4 ‘Home’ screen.

On the ‘User Management’ screen, you can assign your Supports to fill out the Section 4 submission for a given submission type. You have the ability to assign access to one or more Supports to modify any of the Section 4 submissions you have created. To grant a Support access to a Section 4 submission, you must create a new submission and assign a passphrase to that submission. You can also assign different Supports to fill out the Section 4 submission for each submission type.

Section 4 Submission: A submission type will not be available in this menu until it has been created. Select a submission type from the drop-down menu to begin assigning Supports. Once a submission type has been selected, the ‘Assign Users’ group box will update to display only the Supports who have identified you as their Primary AO during CDX registration.

Assign Users: This box shows all the Supports you can assign to complete a Section 4 submission for the submission type selected. To move Supports from the ‘Unassigned’ group

box to the ‘Assigned’ group box, highlight the individual or multiple Supports and click the ‘add >>’ button.

- **Unassigned:** This box contains the registered Supports associated to the Authorized Official who can be assigned to access and edit the selected submission. To highlight multiple Supports hold down the <Ctrl> key on your keyboard while clicking each Support. To highlight multiple consecutive Supports, hold down the <Shift> key on your keyboard while clicking the first and last Support.
- **Assigned:** This box contains the registered Supports who have been given permission, by the Authorized Official, to access and edit the selected submission. To move Supports from the ‘Assigned’ group box to the ‘Unassigned’ group box highlight individual or multiple Supports and click the ‘<< remove’ button. To highlight multiple Supports hold down the <Ctrl> key on your keyboard and click each Support. To highlight multiple consecutive Supports hold down the <Shift> key on your keyboard and click the first and last Support.

When you have finished assigning Supports, click the ‘Save’ button to save the assignments.

Exhibit 3-3 shows a screen capture of the ‘User Management’ screen:

Exhibit 3-3: User Management Screen

Navigation: Click the ‘Submissions’ tab located within the application header to navigate to the ‘Section 4 Test Rules, ECAs, MOUs, and 40 CFR 766’ submissions screen and start or access a submission.

3.4 Create New Submission

As a Primary AO, you can start a new submission and assign your Supports to complete the submission.


3.4.1 Section 4 Test Rules, ECAs, MOUs, and 40 CFR 766 Submissions Screen

You can access the ‘Section 4 Test Rules, ECAs, MOUs, and 40 CFR 766’ submissions screen by clicking the ‘Submissions’ link from the Section 4 ‘Home’ screen, or by clicking the ‘Submissions’ tab located within the application header.

Only one Section 4 submission is created per submission type. Each column within the table of the ‘Section 4 Test Rules, ECAs, MOUs, and 40 CFR 766’ submissions screen can be sorted by clicking the column headers.

Start a New Submission: To create a new Section 4 submission, select the submission type from the ‘Submission Type’ drop-down menu and click the ‘Start New Submission’ button. You will be required to create a passphrase that will be associated with that particular submission. Make sure to create a passphrase that you will remember, as it cannot be retrieved or reset. Refer to **Section 3.4.2** for further instructions on creating a new passphrase.

Edit a Submission: To modify an existing Section 4 submission, click the submission alias link for a submission that has a status of ‘In Progress.’ You will be required to enter the passphrase associated with that particular submission to access and continue editing the submission. Refer to **Section 3.4.3** for further instructions on viewing and modifying a submission that is ‘In Progress.’

Submitted Submission: To edit a submission that has previously been submitted (i.e., to create an amendment), you must first unlock the particular submission by clicking the ‘Lock’ icon () under the ‘Action’ column. You will be required to enter the passphrase associated with that particular submission to gain access to amend the submission. All changes will be submitted as an amendment. Refer to **Section 11** for further instructions on creating an amendment.

The ‘Section 4 Test Rules, ECAs, MOUs, and 40 CFR 766’ submission screen displays the following columns:

- **Submission Alias:** Displays the submission alias that is generated from the ‘Submission Information Type’ screen. The ‘Submission Alias’ column will display as a time stamp unless a unique alias was provided on the ‘Submission Information Type’ screen. The purpose of the submissions alias is to help identify each Section 4 submission when there is more than one.
- **CFR/FRN:** Displays the Code of Federal Regulation/Federal Register Notice (CFR/FRN) that was selected on the ‘Submission Information Type’ screen.
- **Status:** Displays the status of the Section 4 submission. The column will display values as either ‘In Progress’ or ‘Submitted.’
- **Modify Date:** Displays the date the submission was last modified.





- **Submission Date:** Displays the date when you submitted the form to CDX. This date is populated only after you submit.
- **Copy of Record:** Click the ‘Copy of Record’ icon to continue the download copy of record process. The ‘Copy of Record’ icon displays only after you submit to EPA via CDX. The ‘Copy of Record’ icon is confirmation that your submission has been successfully received by EPA. Note that the ‘Copy of Record’ icon may take 15-20 minutes to display on the ‘Section 4 Test Rules, ECAs, MOUs, and 40 CFR 766’ submissions screen. Refer to **Section 10** for further instructions on downloading the copy of record.
- **Action:** Displays a ‘Delete’ icon () for all forms with a status of ‘In Progress.’ Displays a ‘Lock’ icon () for all forms that have been submitted. Only an AO can unlock a form by clicking the ‘Lock’ icon () and unlocking the form. The form will remain unlocked and display the ‘Unlocked’ icon () until the form is resubmitted by the AO.
- **Export Options:** Displays ‘CSV,’ ‘Excel,’ ‘XML,’ and ‘PDF’ links for you to export the ‘Section 4 Test Rules, ECAs, MOUs, and 40 CFR 766’ submissions screen in one of the specified formats as an Excel spreadsheet, Comma-Separated Values (CSV), Extensible Markup Language (XML), or PDF.
- **Submission Type:** Displays the four types of submission you may report on. This drop-down menu contains the following values: ‘Test Rules,’ ‘ECA,’ ‘MOU,’ and ‘40 CFR 766 Dibenzodioxins/Dibenzofurans.’ This field is required.
- **Start New Submission:** Click the ‘Start New Submission’ button once a submission type has been selected from the drop-down menu. You will be taken to the ‘Create New Passphrase’ screen.

Exhibit 3-4 shows a screen capture of the ‘Section 4 Test Rules, ECAs, MOUs, and 40 CFR 766’ submissions screen:

Exhibit 3-4: Section 4 Test Rules, ECAs, MOUs, and 40 CFR 766 Submissions Screen

CSPP

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

Home Submissions User Management Resources

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 (✖).

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

Submission Alias	CFR/FRN	Status	Modify Date	Submission Date	Copy of Record	Action
CFR766-20130717-10:24:00 EDT	40 CFR 766 Dibenzodioxins / Dibenzofurans	In Progress	07/17/2013			✖
ECA-20130718-11:23:20 EDT	70 FR 39630	In Progress	07/18/2013			✖
MOU-20130716-17:23:07 EDT	MOUForm	Submitted	07/17/2013	07/17/2013	↓	🔒
MOU-20130717-13:01:18 EDT	MOUForm	In Progress	07/18/2013			✖
MOU-20130717-16:40:20 EDT	MOUForm	In Progress	07/18/2013			✖
TestRules-20130718-11:04:56 EDT		In Progress	07/18/2013			✖

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

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

Submission Type:

Start New Submission

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

Navigation: Select a submission type from the drop-down menu and click the ‘Start New Submission’ button to navigate to the ‘Create Passphrase’ screen.

3.4.2 Create Passphrase Screen

To start a new Section 4 submission, select a submission type from the ‘Submission Type’ dropdown menu and click the ‘Start New Submission’ button on the ‘Section 4 Test Rules, ECAs, MOUs, and 40 CFR 766’ submissions screen. The ‘Create Passphrase’ screen allows you to create a passphrase and associate that passphrase with the submission.

The application uses the passphrase as an encryption key to protect the contents of the submission. You are responsible for remembering the passphrase and distributing it to the appropriate individuals.

New Passphrase: To associate a passphrase with a submission, enter a passphrase that is at least 8 characters in length. 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.

You will be responsible for remembering the passphrase and distributing it to only authorized Supports for the submission. If you forget the passphrase, you will not be able to access the Section 4 submission to print, submit, or make changes. If you lose or forget the passphrase, you must create a new submission and passphrase. For security reasons, the system administrator does not have access to the passphrase and will not be able to retrieve it or reset it to a new one.

Exhibit 3-5 shows a screen capture of the ‘Create Passphrase’ screen:

Exhibit 3-5: Create Passphrase Screen

The screenshot shows the 'CREATE PASSPHRASE' screen. At the top left is the GSPPP logo. At the top right, it says 'Logged in as: John Doe, Primary Authorized Official' with a 'Log Out' link. The main heading is 'CREATE PASSPHRASE'. Below this, there are instructions: '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 *).' A paragraph follows: '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.' A bolded note states: '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.' Below the text are two input fields: 'New Passphrase:' and 'Confirm New Passphrase:'. A red box highlights both input fields. At the bottom are 'Cancel' and 'Next' buttons. The footer contains links: 'CDX Homepage | MyCDX Homepage | EPA Homepage | Terms and Conditions | Privacy Notice | CDX Helpdesk: (888) 890-1995'.

Navigation: Create a passphrase and click the ‘Next’ button to navigate to the ‘Submission Information Type’ screen.

3.4.3 Enter Passphrase Screen

To edit a submission click the submission alias link for a submission with a status of ‘In Progress’ on the ‘Section 4 Test Rules, ECAs, MOUs, and 40 CFR 766’ submissions screen. The ‘Enter Passphrase’ screen allows you to enter the passphrase associated with the submission.

Enter Passphrase: Enter the passphrase that you originally created and associated with the submission.

Exhibit 3-6 shows a screen capture of the ‘Enter Passphrase’ screen:

Exhibit 3-6 Enter Passphrase Screen

The screenshot shows the 'Enter Passphrase' screen. At the top, the GSPP logo is on the left, and the user is logged in as 'John Doe, Primary Authorized Official' with a 'Log Out' link on the right. The main heading is 'ENTER PASSPHRASE' with the instruction 'Please enter your user passphrase and click Next'. A text input field is provided for the passphrase. Below the input field is a section titled 'Forgot Your Passphrase?' with a message: '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.' At the bottom of the form are 'Cancel' and 'Next' buttons. The 'Next' button is highlighted with a red border. The footer contains links to 'CDX Homepage', 'MyCDX Homepage', 'EPA Homepage', 'Terms and Conditions', 'Privacy Notice', and 'CDX Helpdesk: (888) 890-1995'.

Navigation: Enter the passphrase that you originally created and associated with the submission and click the ‘Next’ button to navigate to the ‘Submission Information Type’ screen.

3.4.4 Navigation Tree

The navigation tree is located on the left hand side of each screen. You can perform the following functions using the navigation tree:

- **Collapse and Expand folders:** Each section of the submission falls under a collapsible folder within the navigation tree, which allows you to save space or easily view items in the navigation tree. When a folder is expanded, you can click the minus sign (-), the folder icon (📁), or the folder title link to collapse that section of the navigation tree. When a folder is collapsed, you can click the plus sign (+), the folder icon (📁), or the folder title link to expand that section of the navigation tree. When you place your cursor over a folder title link, it will highlight in red.
- **Open and Close the Navigation Tree:** To provide more visual form space, a navigation arrow (↔) that opens and closes the navigation tree is located to the right of the navigation tree. Click the arrow (↔) to close the navigation tree and increase the space of the form content. Likewise, click the arrow (↔) on the closed navigation tree to open the navigation tree.

- **Navigate between screens:** You can use the navigation tree to navigate between the various screens within the application. Each screen of the submission is denoted with a form icon (📄). You can click the form icon (📄) or the screen title link to navigate to the selected screen. You are required to save all information entered on a particular screen before navigating to the next screen or all entered information will be lost. A prompt will appear after you click a link in the navigation tree indicating, ‘Are you sure you want to leave the current page? Any unsaved changes will be lost.’ If you click the ‘OK’ button, you will be taken to the requested screen without saving any of the data in the previous screen. If you click the ‘Cancel’ button, the prompt will close and you will not be taken to the requested screen.

Exhibit 3-7 shows a screen capture of the navigation prompt window:

Exhibit 3-7: Navigation Prompt

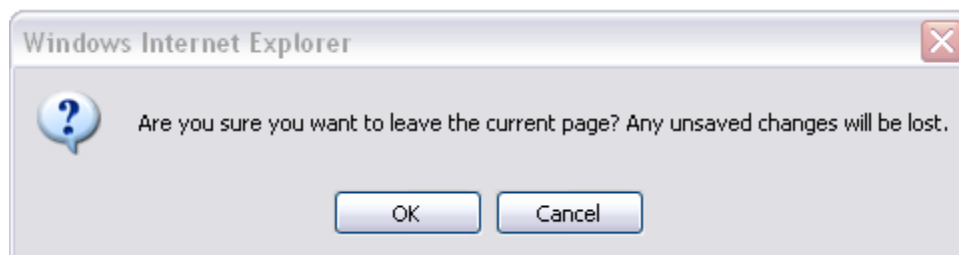


Exhibit 3-8, Exhibit 3-9, Exhibit 3-10, and Exhibit 3-11 show the screen captures of the navigation tree for the Test Rules, ECA, MOU and 40 CFR 766 submissions:

Exhibit 3-8: Test Rules Navigation Tree

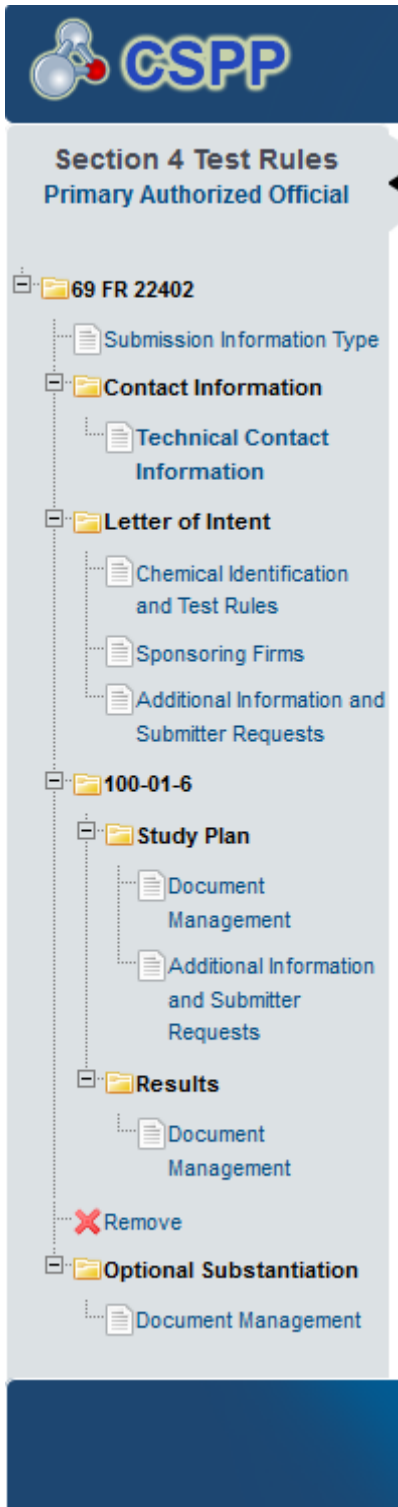


Exhibit 3-9: ECA Navigation Tree

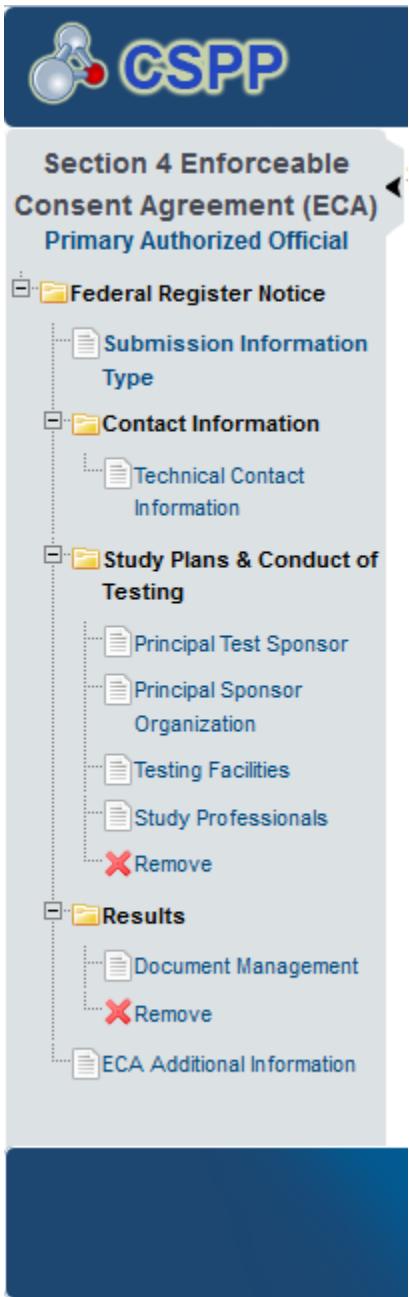


Exhibit 3-10: MOU Navigation Tree

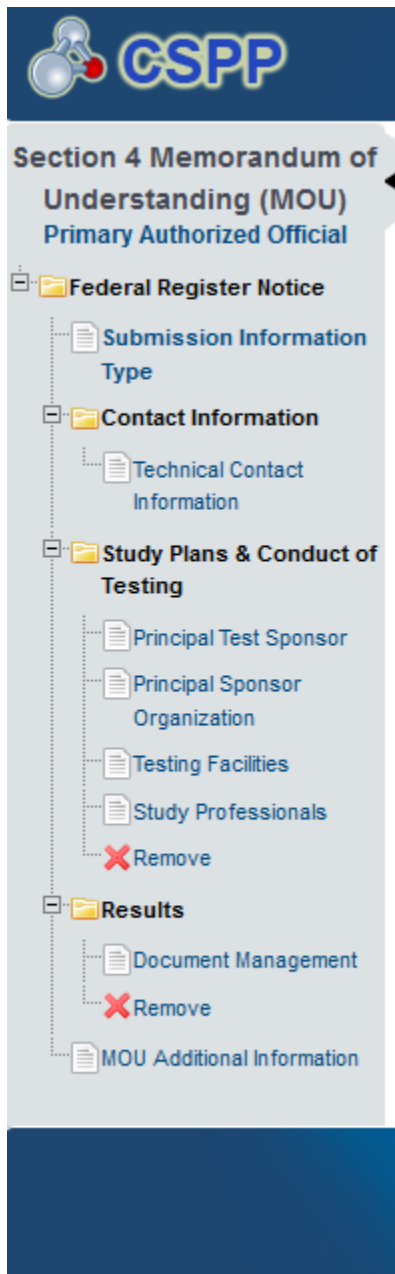
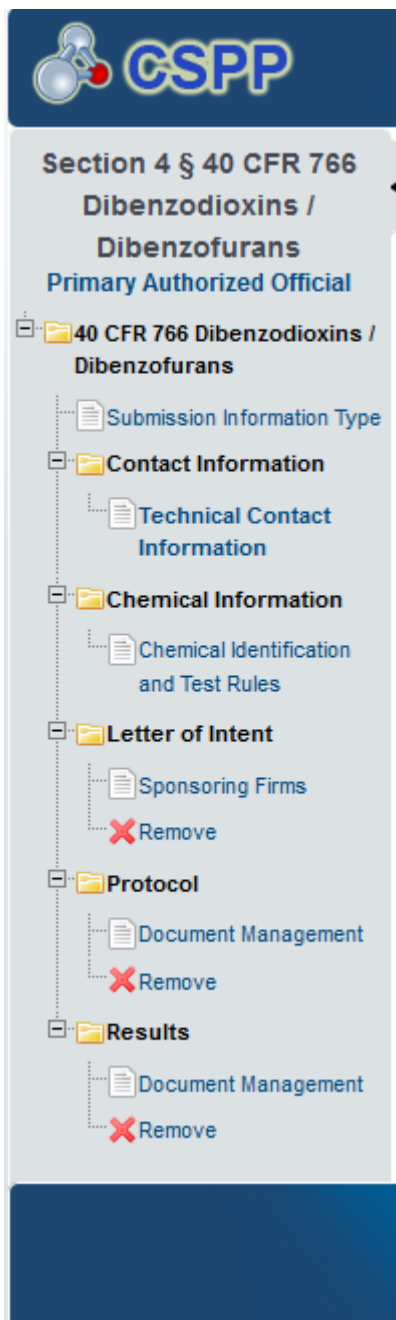


Exhibit 3-11: 40 CFR 766 Navigation Tree



3.4.5 Action Bar

The action bar is located at the bottom of each screen. You can perform the following functions using the action bar:

- **Validate:** You can click the ‘Validate’ icon (✓) at any stage of completing a Section 4 submission to check for certain types of errors in a submission. A ‘Section 4 Validation’ pop-up window generates when you click the ‘Validate’ icon (✓). The ‘Section 4 Validation’ pop-up window displays a report of all validation errors relating to a failed validation. Refer to **Section 8** for more information on validating a submission.

The following exhibits, Exhibit 3-12, Exhibit 3-13, Exhibit 3-14, and Exhibit 3-15 show the different screen captures for the action bar:

Exhibit 3-12: Action Bar – Validate





- **Save:** You can click the ‘Save’ icon () at any stage of completing a Section 4 submission. After you click the ‘Save’ icon (), the data entered on the screen will save. The ‘Save’ function does not validate any data entered.

Exhibit 3-13: Action Bar – Save





- **Preview:** You can click the ‘Preview’ icon () at any stage of completing a Section 4 submission to preview the submission. After you click the ‘Preview’ icon (), you will be taken to the ‘Preview Section 4 Submission’ screen. This screen allows you to view the regular or sanitized PDF. Clicking the ‘Regular’ or ‘Sanitized’ button will generate the ‘Not for Submission’ watermarked PDF version of the submission.

Exhibit 3-14: Action Bar – Preview





- **Submit:** You can click the ‘Submit’ icon () to submit the Section 4 submission after you or the Support have completed all required sections of the Section 4 submission. After you click the ‘Submit’ icon (), the validation process is initiated. Refer to **Section 9** for further instructions on validation errors. You can continue with the submission process only after clearing all validation errors. Refer to **Section 9** for more information on submitting a Section 4 submission.

Exhibit 3-15: Action Bar – Submit

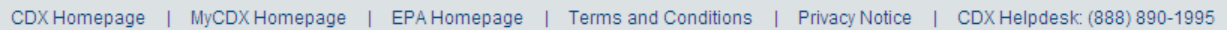


- **Help Links:** You can click any of the Help links, located at the bottom of each screen, at any stage of completing a Section 4 submission.

- If you click the 'CDX Homepage' link, you will be taken to the CDX Homepage.
- If you click the 'MyCDX Homepage' link, you will be taken to the CDX Login.
- If you click the 'EPA Homepage' link, you will be taken to the EPA Homepage.
- If you click the 'Terms and Conditions' link, you will be taken to the CDX Terms and Conditions screen.
- If you click the 'Privacy Notice' link, you will be taken to the CDX Privacy and Security Notice screen.

Exhibit 3-16 shows the screen capture of the action bar 'Help' links:

Exhibit 3-16: Action Bar – Help Links



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

4 Test Rules

4.1 Submission Information Type Screen

You can access the ‘Submission Information Type’ screen by clicking the ‘Next’ button from either the ‘Create New Passphrase’ or ‘Enter Passphrase’ screen. Once you are in the ‘Test Rules’ flow of the Section 4 application, the ‘Submission Information Type’ screen is displayed.

The ‘Submission Information Type’ screen allows you to select the submission information type that you will be submitting to EPA. This screen consists of the following fields:

- **Federal Register Notice:** Allows the user to select a FRN. The selected FRN will dictate which chemicals the user will be able to report on. If you change the FRN at any point while completing the submission, a warning message will display indicating that all data associated with the current FRN will be deleted. This field is required.
- **Submission Alias:** This field is auto-populated with a date and time stamp and is displayed on the ‘Section 4 Test Rules, ECAs, MOUs, and 40 CFR 766’ submissions screen within the ‘Submission Alias’ column. The ‘Submission Alias’ is also used to differentiate between submission types within the ‘User Management’ screen. This field is editable should you choose to provide your own submission alias. This field is required.
- **Study Plan:** The ‘Study Plan’ checkbox will only be available after the Letter of Intent has been submitted. Checking the ‘Study Plan’ checkbox identifies that you are fulfilling the requirements to submit a study plan to EPA. Once you have checked the ‘Study Plan’ checkbox and either saved the information on this screen by clicking the ‘Save’ icon or by clicking the ‘Next’ button, the ‘Study Plan’ folder is generated within the navigation tree allowing you to access the study plans screens required to be completed.
- **Results:** The ‘Results’ checkbox will only be available after the study plan has been submitted. Checking the ‘Results’ checkbox identifies that you are fulfilling the requirements to submit results to EPA. Once you have checked the ‘Results’ checkbox and either saved the information on this screen by clicking the ‘Save’ icon or by clicking the ‘Next’ button, the ‘Results’ folder is generated within the navigation tree allowing you to access the results screens required to be completed.

If you have previously identified a submission type, have accessed the submission type screens, and then unchecked the previously checked submission information type (i.e. Study Plan), you will receive a prompt indicating that all associated information will be lost. Clicking the ‘OK’ button within the prompt will delete all data associated with that submission information type. To close the prompt without losing associated information, click the ‘Cancel’ button.

Once you have identified the type(s) of information you will be submitting to EPA, you are able to access the screen required for completion.

Exhibit 4-1 shows the screen capture of the ‘Submission Information Type’ screen:

Exhibit 4-1: Submission Information Type Screen

Section 4 Test Rules
Primary Authorized Official

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:

Study Plan
 Results

Next

Validate Save Preview Submit

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

Navigation: If this is the first submission, select a federal register notice from the ‘Federal Register Notice’ drop-down menu and click the ‘Next’ button to navigate to the ‘Technical Contact Information’ screen. If you have previously submitted a Letter of Intent, check the ‘Study Plan’ checkbox and click the ‘Save’ or ‘Next’ button to generate the study plan screens and links within the navigation tree. If you have previously submitted a study plan, check the ‘Results’ checkbox and click the ‘Save’ or ‘Next’ button to generate the results screens and links within the navigation tree.

4.2 Technical Contact Information Screen

You can access the ‘Technical Contact Information’ screen by clicking the ‘Next’ button from the ‘Submission Information Type’ screen, or by clicking the ‘Technical Contact Information’ link located within the navigation tree.

The ‘Technical Contact Information’ screen allows you to submit on behalf of another company or submit on behalf of a consortium. To identify if you are submitting on behalf of another entity, the system provides the radio button for ‘This is a submission on behalf of a consortium.’ To identify if you are submitting on behalf of another company, the system provides the radio button ‘This is a submission on behalf of another company.’ The system will not allow you to select more than one radio button per submission.

If you are submitting on behalf of another company, select the ‘This is a submission on behalf of another company’ radio button. This functionality has been implemented to accommodate a company that may use an agent or another individual. If this radio button is selected, you can enter the contact information for the manufacturing or processing establishment on whose behalf

the submission is being made into the ‘Submitting on Behalf Of Company’ screen upon clicking the ‘Next’ button. If you select the radio button for ‘This is a submission on behalf of a consortium,’ you are taken to the ‘Submitting on Behalf of Consortium’ screen upon clicking the ‘Next’ button.

If you are not submitting on behalf of another company or on behalf of a consortium, you must select the ‘N/A’ radio button.

The following fields are displayed on the ‘Technical Contact Information’ screen:

- **CBI:** Checking this checkbox will claim the contact information as Confidential Business Information (CBI).
- **N/A:** Selecting the ‘N/A’ radio button allows you to signify that you are neither making a submission on behalf of a consortium nor making a submission on behalf of another company.
- **This is a submission on behalf of a consortium:** Selecting the ‘This is a submission on behalf of a consortium’ radio button allows you to identify yourself as an individual submitting on behalf of a consortium.
- **This is a submission on behalf of another company:** Selecting the ‘This is a submission on behalf of another company’ radio button allows you to identify yourself as an individual submitting on behalf of another company.
- **Copy CDX Registration:** Clicking the ‘Copy CDX Registration’ button will automatically populate the fields with your contact information provided during CDX registration.
- **Prefix:** Displays all optional prefixes. If applicable, select the appropriate prefix. If you have clicked the ‘Copy CDX Registration’ button, the fields populate with information entered during CDX registration.
- **First Name:** Enter the technical contact’s first name. If you have clicked the ‘Copy CDX Registration’ button, the fields populate with information entered during CDX registration. This field is required.
- **Middle Initial:** Enter the technical contact’s middle initial. If you have clicked the ‘Copy CDX Registration’ button, the fields populate with information entered during CDX registration.
- **Last Name:** Enter the technical contact’s last name. If you have clicked the ‘Copy CDX Registration’ button, the fields populate with information entered during CDX registration. This field is required.
- **Suffix:** Displays all optional suffixes. If applicable, select the appropriate suffix. If you have clicked the ‘Copy CDX Registration’ button, the fields populate with information entered during CDX registration.
- **Company Name:** Enter the technical contact’s company name. If you have clicked the ‘Copy CDX Registration’ button, the fields populate with information entered during CDX registration. This field is required.
- **Phone Number:** Enter the technical contact’s daytime phone number; U.S. or international. This field does not accept dashes. If you have clicked the ‘Copy CDX

Registration' button, the fields populate with information entered during CDX registration. This field is required.

- **Ext:** Enter the extension to the technical contact's daytime phone number.
- **Email Address:** Enter the technical contact's email address. The email address must follow the standard email address format: john.doe@abc.com. If you have clicked the 'Copy CDX Registration' button, the fields populate with information entered during CDX registration. This field is required.
- **Mailing Address 1:** Enter the first line of the technical contact's mailing address. If you have clicked the 'Copy CDX Registration' button, the fields populate with information entered during CDX registration. This field is required.
- **Mailing Address 2:** Enter the second line of the technical contact's mailing address. If you have clicked the 'Copy CDX Registration' button, the fields populate with information entered during CDX registration.
- **City:** Enter the city of the technical contact's mailing address. If you have clicked the 'Copy CDX Registration' button, the fields populate with information entered during CDX registration. This field is required.
- **State:** Enter the state of the technical contact's mailing address. If you have clicked the 'Copy CDX Registration' button, the fields populate with information entered during CDX registration. This is a required field if the 'Country' field is 'United States.'
- **Postal Code:** Enter the postal code of the technical contact's mailing address. If you have clicked the 'Copy CDX Registration' button, the fields populate with information entered during CDX registration. This field is required.
- **Country:** Displays all available countries. Select the country of the technical contact's mailing address. If you have clicked the 'Copy CDX Registration' button, the fields populate with information entered during CDX registration. This is a required field.

Exhibit 4-2 shows the screen capture of the ‘Technical Contact Information’ screen:

Exhibit 4-2: Technical Contact Information Screen

The screenshot displays the 'Technical Contact Information' screen. At the top, the GSPP logo is on the left, and the user is logged in as 'John Doe, Primary Authorized Official' on the right. The navigation menu on the left includes 'Section 4 Test Rules' and 'Primary Authorized Official'. The breadcrumb trail at the top reads 'Section 4 Test Rules > 53 FR 22300 > Contact Information > Technical Contact Information'. The main heading is 'TECHNICAL CONTACT INFORMATION'. Below this, there is a paragraph of instructions: '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.' There are three radio buttons: 'N/A', 'This is a submission on behalf of a consortium', and 'This is a submission on behalf of another company'. Below the radio buttons is a button labeled 'Copy CDX Registration'. The form fields include: CBI (checkbox), Prefix (dropdown), First Name, Middle Initial, Last Name, Suffix (dropdown), Company Name, Phone Number (with an 'Ext.' field), Email Address, Mailing Address 1 (with a note: 'Street address, P.O. box, company name, etc.'), Mailing Address 2 (with a note: 'Apartment, suite, etc.'), City, State (dropdown), Postal Code, and Country (dropdown). At the bottom of the form, there are 'Previous' and 'Next' buttons. The 'Next' button is highlighted with a red box. Below the form, there is a blue bar with icons for 'Validate', 'Save', 'Preview', and 'Submit'. At the very bottom, there is a footer with links: 'CDX Homepage | MyCDX Homepage | EPA Homepage | Terms and Conditions | Privacy Notice | CDX Helpdesk (888) 890-1995'.

Navigation: Click the ‘Next’ button to navigate to the ‘Chemical Identification and Test Rules’ screen if you are not submitting on behalf of a consortium or another company. You can also access this screen by clicking the ‘Chemical Identification and Test Rules’ link within the navigation tree. If you are submitting on behalf of a consortium, you are taken to the ‘Submitting on Behalf of Consortium’ screen. You can also access this screen by clicking the ‘Submitting on Behalf of Consortium’ link within the navigation tree. If you are submitting on behalf of another

company, you are taken to the ‘Submitting on Behalf of Company’ screen. You can also access this screen by clicking the ‘Submitting on Behalf of Company’ link within the navigation tree.

4.3 Submitting on Behalf of Consortium Screen

You can access the ‘Submitting on Behalf of Consortium’ screen by selecting the ‘This is a submission on behalf of a consortium’ radio button and by clicking the ‘Next’ button from the ‘Technical Contact Information’ screen. You can also access this screen by clicking the ‘Submitting on Behalf of Consortium’ link within the navigation tree.

The ‘Submitting on Behalf of Consortium’ screen allows you to enter contact information for each consortium member. The system provides you with the ability to identify the consortium name. The contact information of the logged in user will pre-populate the first consortium field set; this information is editable. To add a consortium member, click the ‘Add’ button to display fields that can be populated and saved. You can click the ‘Save’ button for each consortium member added to capture all information entered on the screen. A minimum of two consortium members must be added to pass validation.

The following fields are displayed on the ‘Submitting on Behalf of Consortium’ screen:

- **Consortium Name:** Enter the name of the consortium.
- **CBI:** Checking this checkbox will claim the consortium information as CBI.
- **Prefix:** The ‘Prefix’ drop-down menu displays all optional prefixes and captures the consortium member contact’s prefix. If applicable, select the appropriate prefix.
- **First Name:** Enter the consortium member contact’s first name. This field is required.
- **Middle Initial:** Enter the consortium member contact’s middle initial.
- **Last Name:** Enter the consortium member contact’s last name. This field is required.
- **Suffix:** The ‘Suffix’ drop-down menu displays all optional suffixes and captures the consortium member contact’s suffix. If applicable, select the appropriate suffix.
- **Company Name:** Enter the consortium member contact’s company name. This field is required.
- **Phone Number:** Enter the consortium member contact’s day time phone number, US or international. This field does not accept dashes. This field is required.
- **Ext:** Enter the extension to the contact’s daytime phone number.
- **Email Address:** Enter the consortium member contact’s email address. The email address must follow the standard email address format: john.doe@abc.com. This field is required.
- **Mailing Address 1:** Enter the first line of the consortium member contact’s mailing address. This field is required.
- **Mailing Address 2:** Enter the second line of the consortium member contact’s mailing address.
- **City:** Enter the city of the consortium member contact’s mailing address. This is a required field.
- **State:** Enter the state of the consortium member contact’s mailing address. This field is required if the ‘Country’ field is set to United States.

-
- **Postal Code:** Enter the postal code of the consortium member contact's mailing address. This field is required.
 - **Country:** Enter the country of the consortium member contact's mailing address. This field is required.

Exhibit 4-3 shows the screen capture for the ‘Submitting on Behalf of Consortium’ screen:

Exhibit 4-3: Submitting on Behalf of Consortium Screen

Section 4 Test Rules > 53 FR 22300 > Contact Information > Submitting on Behalf of Consortium

SUBMITTING ON BEHALF OF CONSORTIUM

Provide the consortium name and click the Add button to enter member contact information.

Consortium Name:

Consortium CBI:

Expand All | Collapse All

CGI FEDERAL (John Doe) Edit ✕

Cancel ✕

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:

Save

Click the **Add** button to add a new consortium member. Add

Previous Next

Validate Save Preview Submit

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

Navigation: Click the ‘Next’ button to navigate to the ‘Chemical Identification and Test Rules’ screen. You can also access this screen by clicking the ‘Chemical Identification and Test Rules’ link within the navigation tree.

4.4 Submitting on Behalf of Company Screen

You can access the ‘Submitting on Behalf of Company’ screen by selecting the ‘This is a submission on behalf of another company’ radio button and clicking the ‘Next’ button from the

‘Technical Contact Information’ screen. You can also access this screen by clicking the ‘Submitting on Behalf of Company’ link within the navigation tree.

The ‘Submitting on Behalf Of Company’ screen presents you with a list of fields that allows you to input the contact information for the manufacturing or processing establishment on whose behalf the submission is being made.

The following fields are displayed on the ‘Submitting on Behalf of Company’ screen:

- **CBI:** Checking this checkbox will claim the manufacturing or processing establishment contact for which the submission is being made as CBI.
- **Prefix:** The ‘Prefix’ drop-down menu displays all optional prefixes and captures the prefix of the manufacturing or processing establishment contact for which the submission is being made. If applicable, select the appropriate prefix.
- **First Name:** Enter the first name of the manufacturing or processing establishment contact for which the submission is being made. This field is required.
- **Middle Initial:** Enter the middle initial of the manufacturing or processing establishment contact for which the submission is being made.
- **Last Name:** Enter the last name of the manufacturing or processing establishment contact for which the submission is being made. This field is required.
- **Suffix:** The ‘Suffix’ drop-down menu displays all optional suffixes and captures the suffix of the manufacturing or processing establishment contact for which the submission is being made. If applicable, select the appropriate suffix.
- **Company Name:** Enter the company name of the manufacturing or processing establishment. This field is required.
- **Phone Number:** Enter the daytime phone number of the manufacturing or processing establishment contact for whom the submission is being made, U.S. or international. This field is required.
- **Ext:** Enter the extension to the contact’s daytime phone number.
- **Email Address:** Enter the email address of the manufacturing or processing establishment contact for which the submission is being made. The email address must follow the standard email address format: john.doe@abc.com. This field is required.
- **Mailing Address 1:** Enter the first line of the mailing address of the manufacturing or processing establishment contact. This field is required.
- **Mailing Address 2:** Enter the second line of the mailing address of the manufacturing or processing establishment contact.
- **City:** Enter the city of the mailing address of the manufacturing or processing establishment contact. This field is required.
- **State:** Enter the state of the mailing address of the manufacturing or processing establishment contact. This is a required field if the ‘Country’ field is defined as ‘United States.’
- **Postal Code:** Enter the postal code of the mailing address of the manufacturing or processing establishment contact. This field is required.

- **Country:** Enter the country of the mailing address of the manufacturing or processing establishment contact. This field is required.

Exhibit 4-4 shows the screen capture of the ‘Submitting on Behalf of Company’ screen:

Exhibit 4-4: Submitting on Behalf of Company Screen

Navigation: Click the ‘Next’ button to navigate to the ‘Chemical Identification and Test Rules’ screen. You can also access this screen by clicking the ‘Chemical Identification and Test Rules’ link within the navigation tree.

4.5 Letter of Intent

The ‘Letter of Intent’ section within the navigation tree must be submitted to EPA before you can complete the ‘Study Plan’ and ‘Results’ screens.

4.5.1 Chemical Identification and Test Rules Screen




You can access the ‘Chemical Identification and Test Rules’ screen by clicking the ‘Next’ button from the ‘Technical Contact Information,’ ‘Submitting on Behalf of Company,’ or the ‘Submitting on Behalf of Consortium’ screen. The ‘Chemical Identification and Test Rules’ screen can also be accessed by clicking the ‘Chemical Identification and Test Rules’ link located within the navigation tree under the ‘Letter of Intent’ folder.

The ‘Chemical Identification and Test Rules’ screen requires you to identify the chemical substance(s) the sponsor(s) intends to use in each of their tests. To identify the chemical substances click the ‘Assign Chemicals’ button. Clicking the ‘Assign Chemicals’ button will generate the ‘Chemical Management’ pop-up window.

The ‘Chemical Management’ pop-up window displays all chemical substances you are able to select. You can select each chemical by checking the checkbox that is located to the left of the chemical. Once the chemical(s) has been checked, click the ‘Save’ button to close the ‘Chemical Management’ pop-up window and display the data on the ‘Chemical Identification and Test Rules’ screen. The chemicals displayed in the ‘Chemical Management’ pop-up window are specific to the FRN selected.

The system allows you to edit previous chemical assignments. To delete a chemical, click the ‘Assign Chemical’ button and uncheck the chemical’s associated checkbox. Click the ‘Save’ button to close the ‘Chemical Management’ pop-up window and save all changes.

Each chemical substance assigned using the ‘Chemical Management’ tool will be displayed on the ‘Chemical Identification and Test Rules’ screen within a field set. Each chemical substance field set requires you to identify a minimum of one test per substance. The system provides a ‘CBI’ checkbox allowing you to claim those tests as CBI.

The system provides a ‘Chemical Test’ drop-down menu for each identified substance. The ‘Actions’ column provides buttons that allow you to save, delete, and add the data entered within the ‘Chemical Test’ text field. To save entered data click the ‘Save’ icon (). To delete entered data click the ‘Delete’ icon (). To add chemical tests click the ‘Add’ icon ().

- **Chemical Identification and Test Rules Screen**

- **CBI:** The ‘CBI’ checkbox allows you to claim information as confidential.
- **Assign Chemicals:** The ‘Assign Chemicals’ button generates the ‘Chemical Management’ window.
- **Chemical Test:** The ‘Chemical Test’ drop-down menu allows you to select the test to be performed using the assign chemical substance. At least one chemical test must be selected.
- **Actions:** The ‘Actions’ column provides you with ‘Save,’ ‘Delete,’ and ‘Add,’ icons that allow you to perform desired functions to the entered chemical test.

- **Chemical Management Pop-Up Window**

- **Save:** Clicking the ‘Save’ button will save all delegated assignments to the ‘Chemical Identification and Test Rules’ screen.
- **Cancel:** Clicking the ‘Cancel’ button will close the ‘Chemical Management’ pop-up window and will not save any changes.

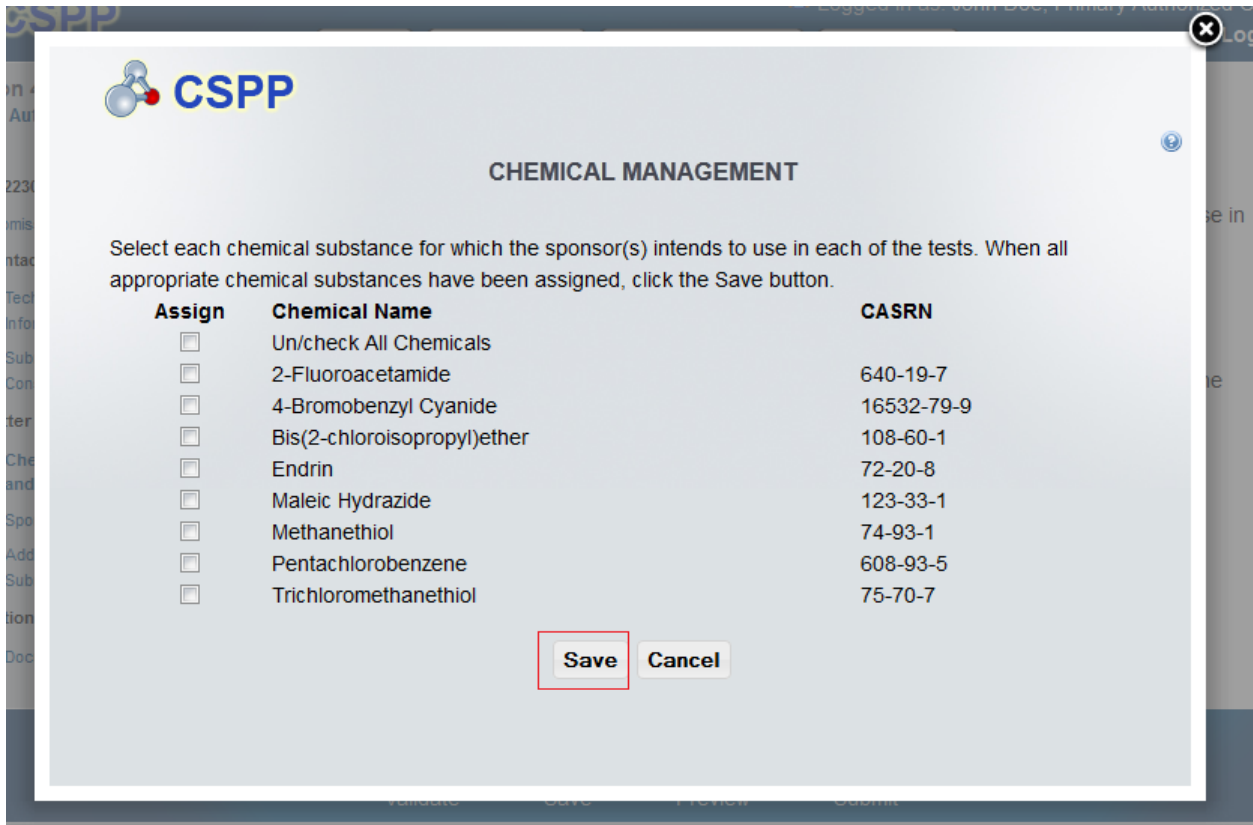
Exhibit 4-5 shows the screen capture of the ‘Chemical Identification and Test Rules’ screen:

Exhibit 4-5: Chemical Identification and Test Rules Screen

The screenshot displays the 'Chemical Identification and Test Rules' screen. At the top left is the GSPP logo. The top right shows the user is logged in as 'John Doe, Primary Authorized Official' with a 'Log Out' link. Below the logo are navigation tabs: 'Home', 'Submissions', 'User Management', and 'Resources'. The left sidebar contains a tree view under 'Section 4 Test Rules Primary Authorized Official', with 'Letter of Intent' expanded to show 'Chemical Identification and Test Rules'. The main content area has a breadcrumb trail: 'Section 4 Test Rules > 53 FR 22300 > Letter of Intent > Chemical Identification And Test Rules'. The title 'CHEMICAL IDENTIFICATION AND TEST RULES' is centered. Below the title, instructions state: '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.' The 'Assign Chemicals' button is highlighted with a red box. Further instructions state: 'For the chemical substance(s) listed below, identify the testing requirement(s) to be performed on the specified substance.' Below this are 'Previous' and 'Next' buttons. At the bottom, a dark blue bar contains icons and labels for 'Validate', 'Save', 'Preview', and 'Submit'. The footer contains links: 'CDX Homepage | MyCDX Homepage | EPA Homepage | Terms and Conditions | Privacy Notice | CDX Helpdesk: (888) 890-1995'.

Exhibit 4-6 shows the screen capture for the 'Chemical Management' pop-up window:

Exhibit 4-6: Chemical Management Pop-Up Window



Navigation: Click the 'Save' button to return to the 'Chemical Identification and Test Rules' screen.

Exhibit 4-7 shows the screen capture of the ‘Chemical Identification and Test Rules’ screen populated:

Exhibit 4-7: Chemical Identification and Test Rules Screen (Populated)

Navigation: Click the ‘Next’ button to navigate to the ‘Sponsoring Firms’ screen. You can also access this screen by clicking the ‘Sponsoring Firms’ link within the navigation tree.

4.5.2 Sponsoring Firms Screen

To access the ‘Sponsoring Firms’ screen click the ‘Next’ button from the ‘Chemical Identification and Test Rules’ screen. You can also access the ‘Sponsoring Firms’ screen by clicking the ‘Sponsoring Firms’ link located within the navigation tree under the ‘Letter of Intent’ folder.

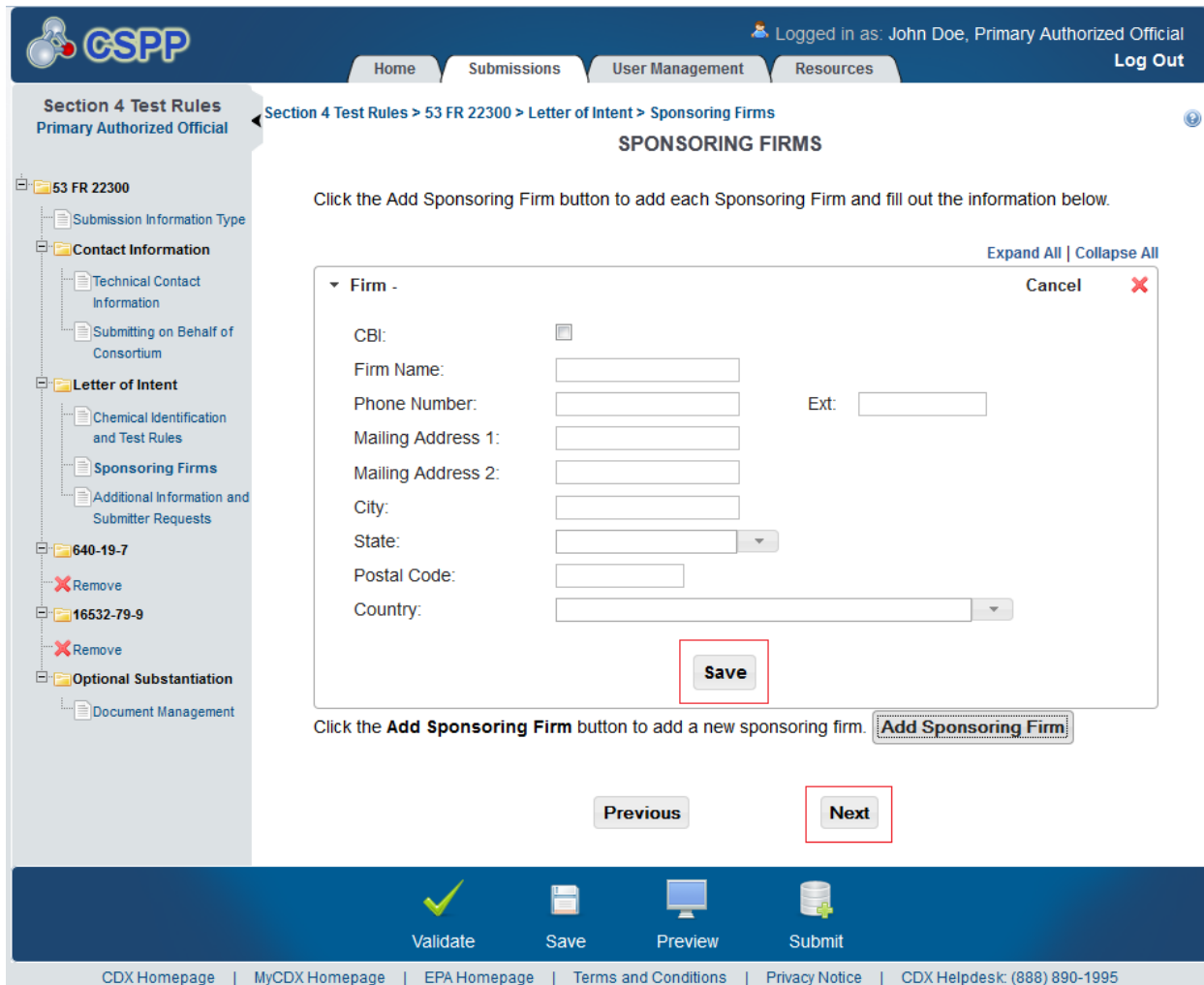
The 'Sponsoring Firms' screen requires you to identify each sponsoring firm involved in the testing of the identified chemical substances. To identify a sponsoring firm click the 'Add Sponsoring Firm' button. Clicking the 'Add Sponsoring Firm' button generates a sponsoring firm field set. Once you have entered data into a sponsoring firm field set, you must click the 'Save' button located within that field set prior to adding another firm. If the data previously entered and saved within a field set needs to be edited, you can click the 'Edit' button located in the upper right corner of that field set. If a sponsoring firm needs to be deleted, you can click the 'Delete' icon that is also located in the upper right corner of each field set.

The system allows you to expand/collapse each field set by clicking the sponsoring firm field set header. If you wish to expand/collapse all field sets simultaneously, click the 'Expand All' or 'Collapse All' links located at the top right of all consortium field sets.

- **CBI:** Check the 'CBI' checkbox to claim the sponsoring firm as CBI. If you claim information as CBI, you are required to complete substantiation.
- **Firm Name:** Enter the sponsoring firm's company name. This field is required.
- **Phone Number:** Enter the sponsoring firm's daytime phone number; U.S. or international. This field does not accept dashes. This field is required.
- **Ext:** Enter the extension to the contact's daytime phone number.
- **Mailing Address 1:** Enter the first line of the sponsoring firm's mailing address. This field is required.
- **Mailing Address 2:** Enter the second line of the sponsoring firm's mailing address.
- **City:** Enter the city of the sponsoring firm's mailing address. This field is required.
- **State:** Enter the state of the sponsoring firm's mailing address. This field is required when 'United States' is identified as the country.
- **Postal Code:** Enter the postal code of the sponsoring firm's mailing address. This field is required.
- **Country:** Enter the country of the sponsoring firm's mailing address. This field is required.

Exhibit 4-8 shows the screen capture of the ‘Sponsoring Firms’ screen:

Exhibit 4-8: Sponsoring Firms Screen



Navigation: Click the ‘Next’ button to navigate to the ‘Additional Information and Submitter Requests’ screen. You can also access this screen by clicking the ‘Additional Information and Submitter Requests’ link within the navigation tree.

4.5.3 Additional Information and Submitter Requests Screen

The ‘Additional Information and Submitter Requests’ screen allows you to submit additional information or special requests. This is not a required screen and does not need to be completed prior to submitting information to EPA.

The system provides you with an ‘Information/Request Type’ drop-down menu that contains the following values:

- Amendment to the Study Plan
- Exemption Application
- Extension of Time to Submit a Study Plan

-
- Modification of Test Standards or Schedules during Conduct of Test
 - Submission of Equivalence Data
 - Appeal of Denial of Exemption Application

To submit additional information or a special request select the desired value from the 'Information/Request Type' drop-down menu and click the 'Add Information' button. Depending on your information or request selection, this choice dictates the pop-up window that generates upon clicking the 'Add Information' button.

If you select 'Amendment to the Study Plan' from the 'Information/Request Type' drop-down menu and click the 'Add Information' button, the 'Amendment to the Study Plan' pop-up window generates. The 'Amendment to the Study Plan' pop-up window allows you to attach documentation. To attach documentation click the 'Browse' button. Clicking the 'Browse' button will generate the 'File Upload' window on your hard drive. Search for your desired document and click the 'Open' button. Clicking the 'Open' button will close the 'File Upload' window and populate the 'Browse' field with the attached document. Click the 'OK' button within the 'Amendment to the Study Plan' pop-up window to return to the 'Additional Information and Submitter Requests' screen. The attached document will be displayed within the table.

If you select 'Exemption Application' from the 'Information/Request Type' drop-down menu and click the 'Add Information' button, the 'Exemption Application' pop-up window generates. The 'Exemption Application' pop-up window allows you to identify the testing requirement(s) from which you seek exemption. The 'Exemption Application' pop-up window also allows you to provide the basis for the exemption request within a text box. Click the 'OK' button within the 'Exemption Application' pop-up window to return to the 'Additional Information and Submitter Requests' screen. The previously entered data will be displayed within the table.

If you select 'Extension of Time to Submit Study Plan' from the 'Information/Request Type' drop-down menu and click the 'Add Information' button, the 'Extension of Time to Submit Study Plan' pop-up window is generated. The 'Extension of Time to Submit Study Plan' pop-up window provides a text box that allows you to identify why EPA should grant the extension. Click the 'OK' button within the 'Extension of Time to Submit Study Plan' pop-up window to return to the 'Additional Information and Submitter Requests' screen. The previously entered data will be displayed within the table.

If you select 'Modification of Test Standards or Schedules During Conduct of Test' from the 'Information/Request Type' drop-down menu and click the 'Add Information' button, the 'Modification of Test Standards or Schedules During Conduct of Test' pop-up window generates. The 'Modification of Test Standards or Schedules During Conduct of Test' pop-up window provides a text box that allows you to enter an explanation and rationale for the modification. Click the 'OK' button within the 'Modification of Test Standards or Schedules During Conduct of Test' pop-up window to return to the 'Additional Information and Submitter Requests' screen. The previously entered data will be displayed within the table.

If you select 'Submission of Equivalence Data' from the 'Information/Request Type' drop-down menu and click the 'Add Information' button, the 'Submission of Equivalence Data' pop-up window generates. The 'Submission of Equivalence Data' pop-up window allows you to attach documentation. To attach documentation click the 'Browse' button. Clicking the 'Browse' button will generate the 'File Upload' window on your hard drive. Search for your desired

document and click the 'Open' button. Clicking the 'Open' button will close the 'File Upload' window and populate the 'Browse' field with the attached document. The system allows you to claim the attached documentation as CBI. When information is claimed as CBI, substantiation of that information is optional. Click the 'OK' button within the 'Amendment to the Study Plan' pop-up window to return to the 'Additional Information and Submitter Requests' screen. The previously attached document will be displayed within the table.

If you select 'Appeal of Denial of Exemption Application' from the 'Information/Request Type' drop-down menu and click the 'Add Information' button, the 'Appeal of Denial of Exemption Application' pop-up window generates. The 'Appeal of Denial of Exemption Application' pop-up window provides a text box that allows you to identify the basis for the reconsideration. The system also allows you to claim a hearing by checking the 'Check here if you wish to request a hearing' checkbox. If you request a hearing, a second text box generates and requires you to explain why a hearing is necessary. Click the 'OK' button within the 'Appeal of Denial of Exemption Application' pop-up window to return to the 'Additional Information and Submitter Requests' screen. The previously entered data will be displayed within the table.

The table on the 'Additional Information and Submitter Requests' screen displays the following columns:

- **File Name:** Displays the file name of the attached document if applicable. If text was entered within the information/request type pop-up window the 'File Name' column will display as 'N/A.' The system allows you to click the 'File Name' link for each information/request type to edit previously attached documentation or text.
- **Information/Request Type:** Displays the information/request type selected from the 'Information/Request Type' drop-down menu.
- **Attachment Date:** Displays the date the information/request type was attached.
- **Action:** Displays the 'Delete' icon (✘), which allows you to delete a desired row within the table and all associated information that pertained to the information/request type.

Exhibit 4-9 shows the screen capture for the ‘Additional Information and Submitter Requests’ screen:

Exhibit 4-9: Additional Information and Submitter Requests Screen

Section 4 Test Rules
Primary Authorized Official

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

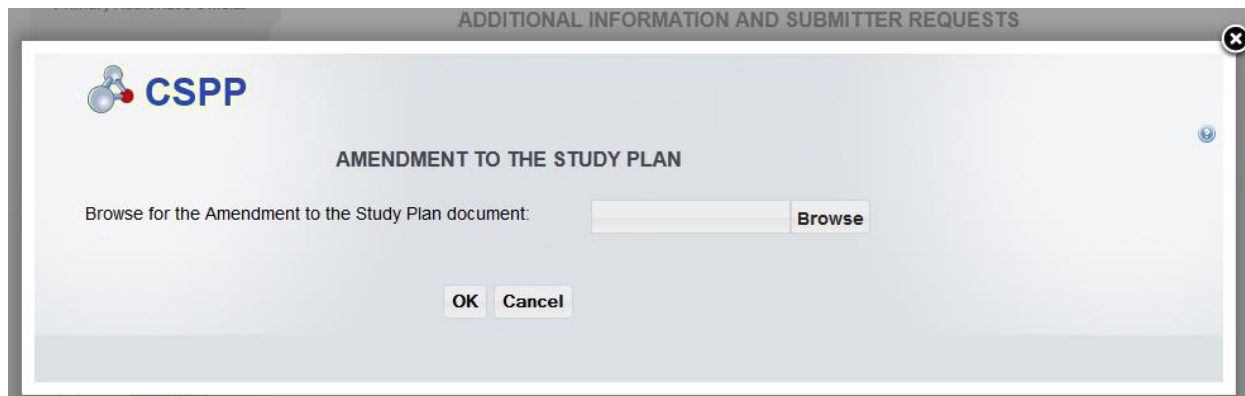
Validate Save Preview Submit

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

Navigation: If you are completing a letter of intent, click the ‘Submit’ icon located in the footer to begin the submission process. If you have previously submitted a letter of intent and checked the ‘Study Plan’ checkbox, click the ‘Next’ button to navigate to the ‘Document Management’ screen located under the ‘Study Plan’ folder within the navigation tree. You can also access this screen by clicking the ‘Document Management’ link under the ‘Study Plan’ folder within the navigation tree.

Exhibit 4-10 shows the screen capture of the ‘Amendment to the Study Plan’ pop-up window:

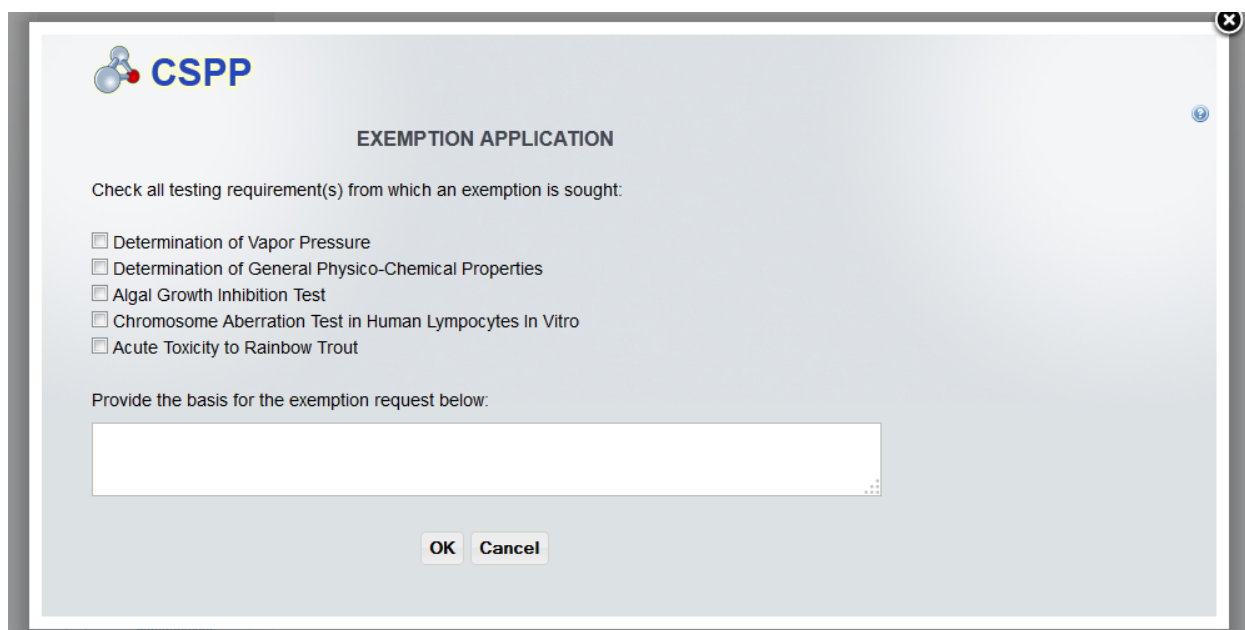
Exhibit 4-10: Amendment to the Study Plan Pop-Up Window



Navigation: Click the ‘OK’ button to return to the ‘Additional Information and Submitter Requests’ screen. Click the ‘Cancel’ or ‘X’ button to close the window and return to the ‘Additional Information and Submitter Requests’ screen without saving the entered information.

Exhibit 4-11 shows the screen capture of the ‘Exemption Application’ pop-up window:

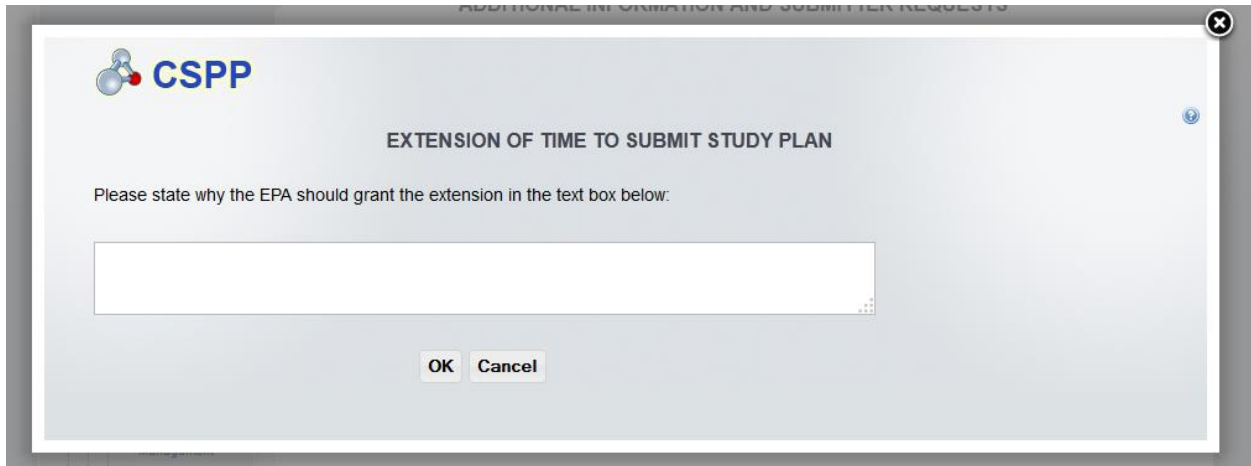
Exhibit 4-11: Exemption Application Pop-Up Window



Navigation: Click the ‘OK’ button to return to the ‘Additional Information and Submitter Requests’ screen. Click the ‘Cancel’ or ‘X’ button to close the window and return to the ‘Additional Information and Submitter Requests’ screen without saving the entered information.

Exhibit 4-12 shows the screen capture of the ‘Extension of Time to Submit Study Plan’ pop-up window:

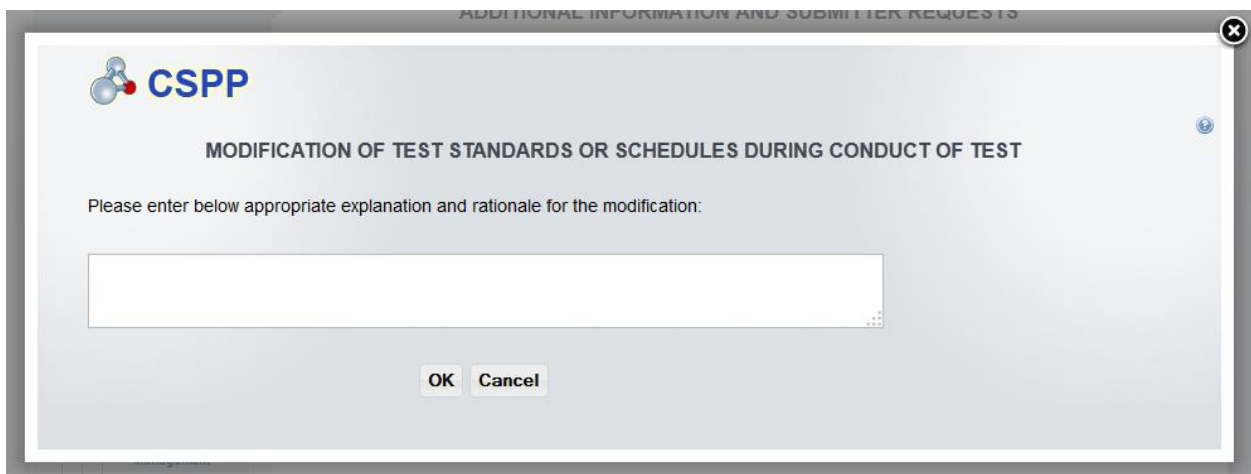
Exhibit 4-12 Extension of Time to Submit Study Plan Pop-Up Window

The screenshot shows a pop-up window titled 'EXTENSION OF TIME TO SUBMIT STUDY PLAN'. The window has a light gray background and a dark gray border. In the top left corner, there is the CSPP logo, which consists of three blue spheres of varying sizes connected by lines, followed by the text 'CSPP' in blue. The title 'EXTENSION OF TIME TO SUBMIT STUDY PLAN' is centered at the top in a dark gray font. Below the title, there is a line of text: 'Please state why the EPA should grant the extension in the text box below:'. Underneath this text is a large, empty white text box with a thin gray border. At the bottom of the window, there are two buttons: 'OK' and 'Cancel', both in a light gray box with dark text. In the top right corner of the window, there is a small blue help icon and a close button (an 'X' in a circle).

Navigation: Click the ‘OK’ button to return to the ‘Additional Information and Submitter Requests’ screen. Click the ‘Cancel’ or ‘X’ button to close the window and return to the ‘Additional Information and Submitter Requests’ screen without saving the entered information.

Exhibit 4-13 shows the screen capture of the ‘Modification of Test Standards or Schedules during Conduct of Test’ pop-up window:

Exhibit 4-13: Modification of Test Standards or Schedules during Conduct of Test Pop-Up Window

The screenshot shows a pop-up window titled 'MODIFICATION OF TEST STANDARDS OR SCHEDULES DURING CONDUCT OF TEST'. The window has a light gray background and a dark gray border. In the top left corner, there is the CSPP logo, which consists of three blue spheres of varying sizes connected by lines, followed by the text 'CSPP' in blue. The title 'MODIFICATION OF TEST STANDARDS OR SCHEDULES DURING CONDUCT OF TEST' is centered at the top in a dark gray font. Below the title, there is a line of text: 'Please enter below appropriate explanation and rationale for the modification:'. Underneath this text is a large, empty white text box with a thin gray border. At the bottom of the window, there are two buttons: 'OK' and 'Cancel', both in a light gray box with dark text. In the top right corner of the window, there is a small blue help icon and a close button (an 'X' in a circle).

Navigation: Click the ‘OK’ button to return to the ‘Additional Information and Submitter Requests’ screen. Click the ‘Cancel’ or ‘X’ button to close the window and return to the ‘Additional Information and Submitter Requests’ screen without saving the entered information.

Exhibit 4-14 shows the screen capture of the ‘Submission of Equivalence Data’ pop-up window:

Exhibit 4-14: Submission of Equivalence Data Pop-Up Window

CSPP

SUBMISSION OF EQUIVALENCE DATA

CBI:

Browse for the appropriate Equivalence Data document: **Browse**

Browse for the appropriate sanitized Equivalent Data document: **Browse (optional)**

OK Cancel

Navigation: Click the ‘OK’ button to return to the ‘Additional Information and Submitter Requests’ screen. Click the ‘Cancel’ or ‘X’ button to close the window and return to the ‘Additional Information and Submitter Requests’ screen without saving the entered information.

Exhibit 4-15 shows the screen capture of the ‘Appeal of Denial of Exemption Application’ pop-up window:

Exhibit 4-15: Appeal of Denial of Exemption Application Pop-Up Window

CSPP

APPEAL OF DENIAL OF EXEMPTION APPLICATION

Please enter below the basis for the request for reconsideration:

Check here if you wish to request a hearing:

Please explain why a hearing is necessary:

OK Cancel

Navigation: Click the ‘OK’ button to return to the ‘Additional Information and Submitter Requests’ screen. Click the ‘Cancel’ or ‘X’ button to close the window and return to the ‘Additional Information and Submitter Requests’ screen without saving the entered information.

4.6 Study Plan

If you have identified you are submitting a study plan, you are required to complete the following screens to satisfy the reporting requirements.

4.6.1 Document Management Screen

You can access the ‘Document Management’ screen by clicking the ‘Next’ button from the ‘Additional Information and Test Rules’ screen or by clicking the ‘Document Management’ link within the navigation tree under the ‘Study Plan’ folder. This screen is only available once a letter of intent has been submitted and you have selected the ‘Study Plan’ checkbox on the ‘Submission Information Type’ screen.

The ‘Document Management’ screen allows you to upload documents related to Section 4 reporting requirements. To add Section 4 study plan documentation, click the ‘Attach Document’ button. Clicking the ‘Attach Document’ button displays the ‘Document Management’ pop-up window.

The ‘Document Management’ pop-up window requires you to identify the document type by selecting ‘Phase I’ or ‘Phase II’ from the ‘Document Type’ drop-down menu. Once you have identified the document type to be attached, click the ‘Browse’ button. Clicking the ‘Browse’ button will generate the ‘File Upload’ window where you can select the desired document from your hard drive. Once you have identified the document to be attached, click the ‘Open’ button. Clicking the ‘Open’ button will close the ‘File Upload’ window and display the attached document within the ‘Browse’ field. To claim attached documentation as CBI, check the ‘CBI’ checkbox. If you claim the attached documentation as CBI you are required to attach the sanitized version of the attached document. To do so, follow the steps outlined above. Once you have attached all desired documentation, click the ‘OK’ button to return to the ‘Document Management’ screen.

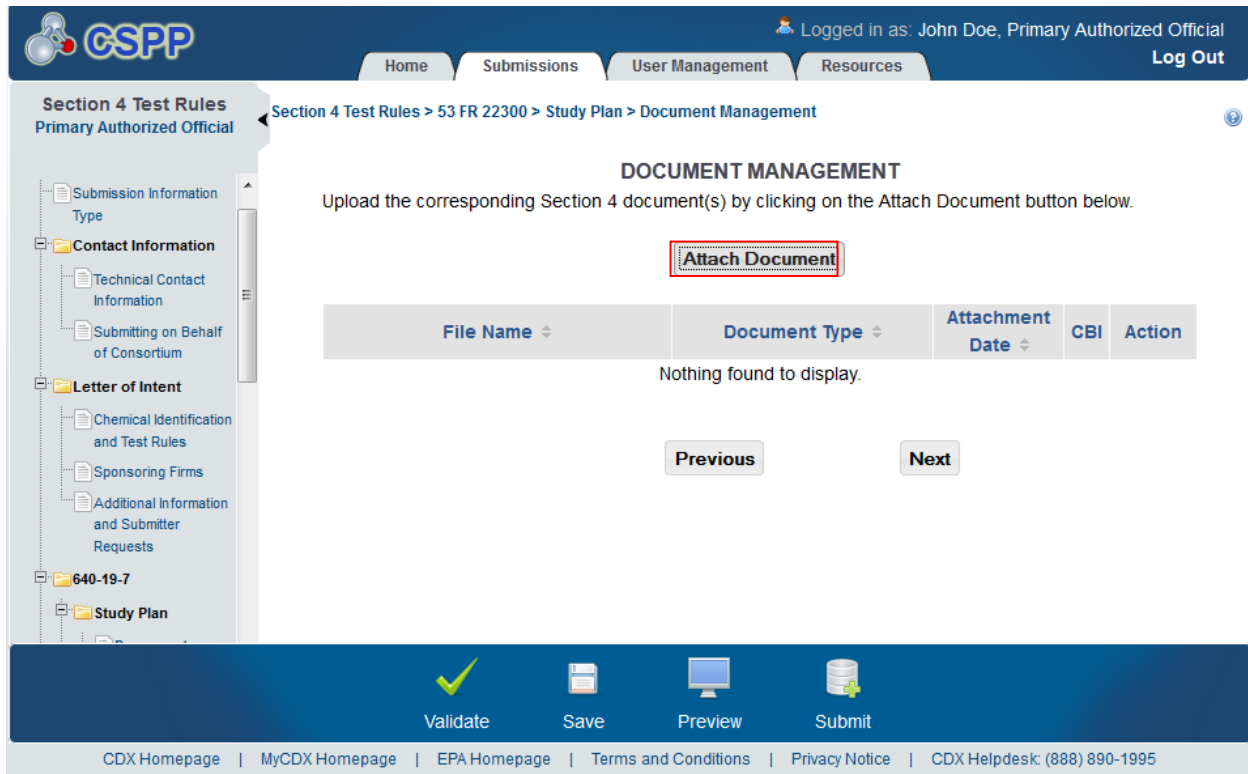
The ‘Document Management’ table displays the file name of the document, the document type, the date the document was attached, and whether the document was claimed as CBI. The system allows you to delete unwanted documents by clicking the ‘Delete’ icon located under the ‘Action’ column for a given row. Clicking the file name links within the ‘File Name’ column generates the ‘Document Management’ pop-up window. This allows you to view and edit previously attached documents.

- **Attach Document:** Clicking the ‘Attach Document’ button displays the ‘Document Management’ pop-up window.
- **File Name:** Displays the name of the attached file. Clicking the file name link generates the ‘Document Management’ pop-up window and allows you to view and edit the attached document.
- **Document Type:** Displays the document type of the attached document.
- **Attachment Date:** Displays the date the document was attached.
- **CBI:** Displays whether the attached document was claimed CBI.

- **Action:** Clicking the ‘Delete’ icon will generate a prompt confirming the command. If you click the ‘OK’ button within the prompt, this will delete the specified row and all contents within.

Exhibit 4-16 shows the screen capture of the ‘Document Management’ screen:

Exhibit 4-16 Document Management Screen



Navigation: Click the ‘Next’ button to navigate to the ‘Additional Information and Submitter Requests’ screen.

The ‘Document Management’ pop-up window allows you to browse for a document from your hard drive and attach one document at a time. You must attach at least one document. Clicking the ‘Browse’ button will generate the ‘File Upload’ window on your hard drive. Select the document you wish to attach and click the ‘Open’ button. If a document is marked as ‘CBI’ then a sanitized copy of the document can be attached. To save the attached document to the ‘Document Management’ screen, click the ‘OK’ button within the ‘Document Management’ pop-up window.

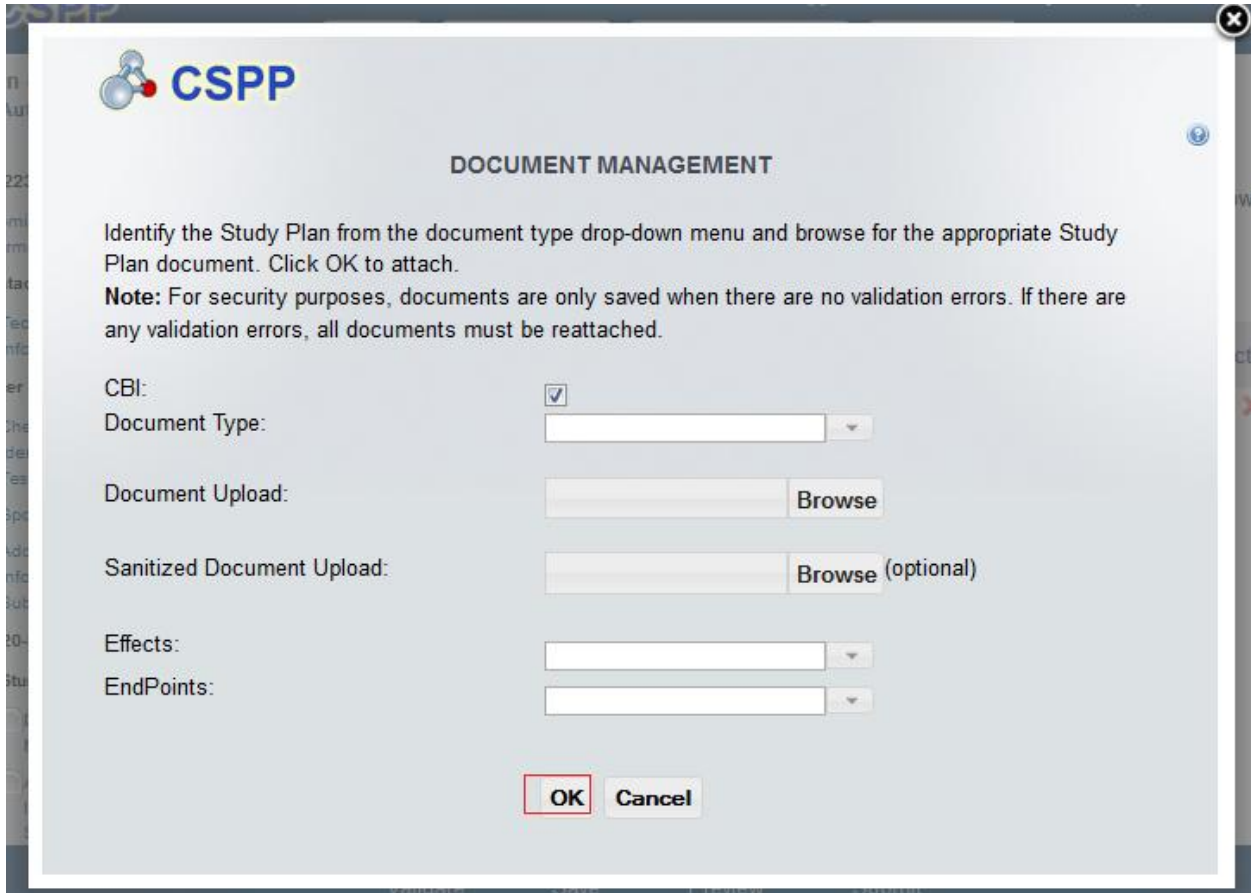
- **Document Type:** Select the document type specific to the uploaded document. Displays the following options: ‘Phase I’ and ‘Phase II.’
- **CBI:** Checking the ‘CBI’ checkbox will claim the uploaded document as CBI and generate a sanitized document upload field.
- **Document Upload:** Clicking the ‘Browse’ button will generate the ‘File Upload’ window for you to search for a particular file on your hard drive.

-
- **Sanitized Document Upload:** This field will only be displayed when the ‘CBI’ checkbox is checked. Clicking the ‘Browse’ button will generate the ‘File Upload’ window for you to search for a particular file on your hard drive.
 - **Effects:** Select the effect specific to the chemical report. Displays the following options: ‘Health Effects,’ ‘Environmental Fate,’ ‘Ecotoxicity,’ and ‘Physical Chemical Properties.’
 - **Endpoints:** Select an endpoint that is specific to the chemical report. The endpoints will be specific to the effect chosen from the ‘Effects’ drop-down menu. If no effect is chosen, no endpoints will display.

Refer to Appendix B. Submission Types, Effects, and Endpoints for a complete list of values that are found within each of the drop-down menus on the ‘Document Management’ pop-up window. If you have multiple documents that share the same effects and/or endpoints, you must upload each individual document separately and select values from the drop-down menus.

Exhibit 4-17 shows the screen capture of the ‘Document Management’ pop-up window:

Exhibit 4-17 Document Management Pop-Up Window



Navigation: Click the ‘OK’ button to return to the ‘Document Management’ screen.

4.6.2 Additional Information and Submitter Requests Screen

You can access the ‘Additional Information and Submitter Requests’ screen by clicking the ‘Next’ button from the ‘Document Management’ screen. The ‘Additional Information and Submitter Requests’ screen can also be accessed by clicking the ‘Additional Information and Submitter Requests’ link within the navigation tree under the ‘Study Plan’ folder.

The ‘Additional Information and Submitter Requests’ screen allows you to submit additional information or special requests. This screen is not a required screen and does not need to be completed prior to submitting information to EPA.

The system provides you with an ‘Information/Request Type’ drop-down menu that contains the following values:

- Amendment to the Study Plan
- Exemption Application
- Extension of Time to Submit a Study Plan
- Modification of Test Standards or Schedules During Conduct of Test

-
- Submission of Equivalence Data
 - Appeal of Denial of Exemption Application

To submit additional information or a special request select the desired value from the ‘Information/Request Type’ drop-down menu and click the ‘Add Information’ button. Depending on your information or request selection, this dictates the pop-up window that will be generated upon clicking the ‘Add Information’ button.

If you select ‘Amendment to the Study Plan’ from the ‘Information/Request Type’ drop-down menu and click the ‘Add Information’ button, the ‘Amendment to the Study Plan’ pop-up window generates. The ‘Amendment to the Study Plan’ pop-up window allows you to attach documentation. To attach documentation click the ‘Browse’ button. Clicking the ‘Browse’ button will generate the ‘File Upload’ window. Search for your desired document on your hard drive and click the ‘Open’ button. Clicking the ‘Open’ button will close the ‘File Upload’ window and populate the ‘Browse’ field with the attached document. Click the ‘OK’ button within the ‘Amendment to the Study Plan’ pop-up window to return to the ‘Additional Information and Submitter Requests’ screen. The previously attached document will be displayed within the table.

If you select ‘Exemption Application’ from the ‘Information/Request Type’ drop-down menu and click the ‘Add Information’ button, the ‘Exemption Application’ pop-up window generates. The ‘Exemption Application’ pop-up window allows you to identify the testing requirement(s) from which you seek exemption. The ‘Exemption Application’ pop-up window also allows you to provide the basis for the exemption request within a text box. Click the ‘OK’ button within the ‘Exemption Application’ pop-up window to return to the ‘Additional Information and Submitter Requests’ screen. The previously entered data will be displayed within the table.

If you select ‘Extension of Time to Submit Study Plan’ from the ‘Information/Request Type’ drop-down menu and click the ‘Add Information’ button, the ‘Extension of Time to Submit a Study Plan’ pop-up window is generated. The ‘Extension of Time to Submit Study Plan’ pop-up window provides a text box that allows you to identify why EPA should grant the extension. Click the ‘OK’ button within the ‘Extension of Time to Submit Study Plan’ pop-up window to return to the ‘Additional Information and Submitter Requests’ screen. The previously entered data will be displayed within the table.

If you select ‘Modification of Test Standards or Schedules During Conduct of Test’ from the ‘Information/Request Type’ drop-down menu and click the ‘Add Information’ button, the ‘Modification of Test Standards or Schedules During Conduct of Test’ pop-up window generates. The ‘Modification of Test Standards or Schedules During Conduct of Test’ pop-up window provides a text box that allows you to enter an explanation and rationale for the modification. Click the ‘OK’ button within the ‘Modification of Test Standards or Schedules During Conduct of Test’ pop-up window to return to the ‘Additional Information and Submitter Requests’ screen. The previously entered data will be displayed within the table.

If you select ‘Submission of Equivalence Data’ from the ‘Information/Request Type’ drop-down menu and click the ‘Add Information’ button, the ‘Submission of Equivalence Data’ pop-up window generates. The ‘Submission of Equivalence Data’ pop-up window allows you to attach documentation. To attach documentation click the ‘Browse’ button. Clicking the ‘Browse’ button will generate the ‘File Upload’ window. Search for your desired document on your hard drive and click the ‘Open’ button. Clicking the ‘Open’ button will close the ‘File Upload’

window and populate the 'Browse' field with the attached document. The system allows you to claim the attached documentation as CBI. When information is claimed as CBI, substantiation of that information is then required. Click the 'OK' button within the 'Amendment to the Study Plan' pop-up window to return to the 'Additional Information and Submitter Requests' screen. The previously attached document will be displayed within the table.

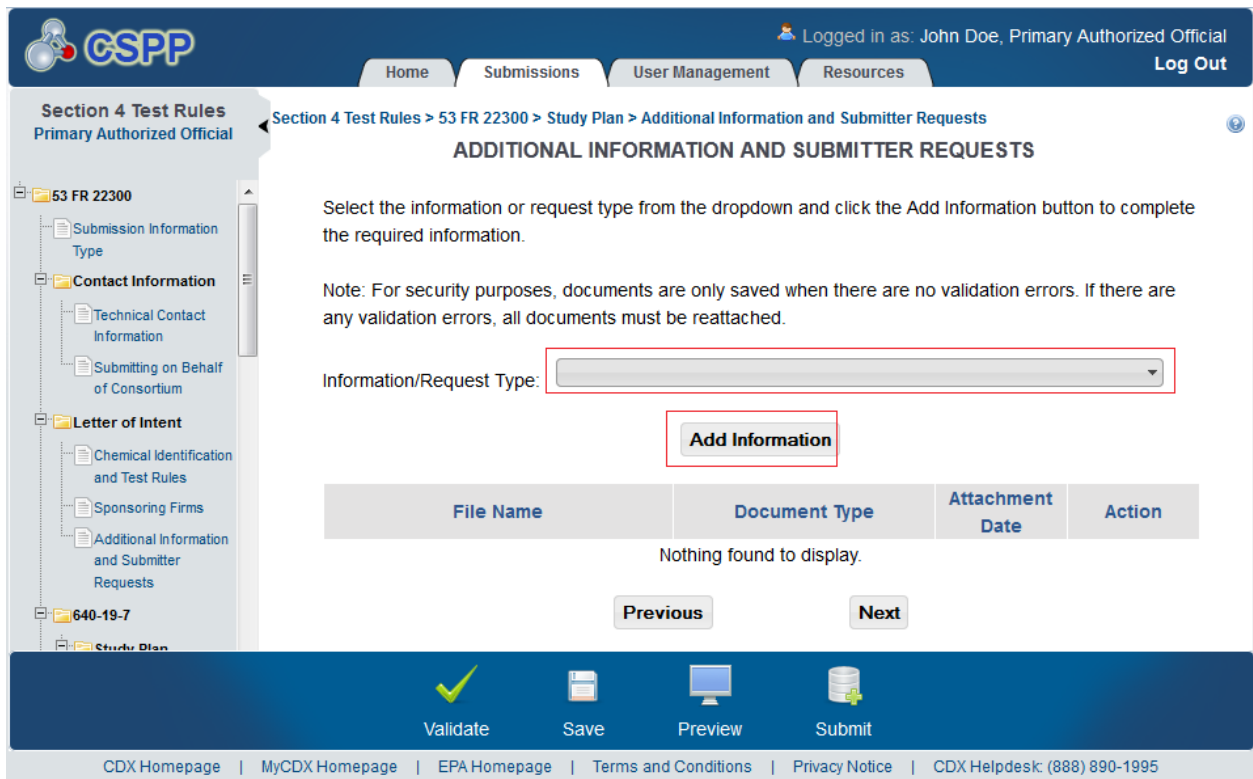
If you select 'Appeal of Denial of Exemption Application' from the 'Information/Request Type' drop-down menu and click the 'Add Information' button, the 'Appeal of Denial of Exemption Application' pop-up window generates. The 'Appeal of Denial of Exemption Application' pop-up window provides a text box that allows you to identify the basis for the reconsideration. The system also allows you to claim a hearing by checking the 'Check here if you wish to request a hearing' checkbox. If you request a hearing, a second text box generates and requires you to explain why a hearing is necessary. Click the 'OK' button within the 'Appeal of Denial of Exemption Application' pop-up window to return to the 'Additional Information and Submitter Requests' screen. The previously entered data will be displayed within the table.

The table on the 'Additional Information and Submitter Requests' screen displays the following columns:

- **File Name:** Displays the file name of the attached document if applicable. If text was entered within the information/request type pop-up window the 'File Name' column will display as 'N/A.' The system allows you to click the file name link for each information/request type to edit previously attached documentation or text.
- **Document Type:** Displays the information/request type selected from the 'Information/Request Type' drop-down menu.
- **Attachment Date:** Displays the date the information/request type was attached.
- **Action:** Displays the 'Delete' icon (✖) which allows you to delete a desired row within the table and all associated information that pertained to the information/request type. Clicking the 'Delete' icon (✖) will generate a prompt confirming the command. Clicking the 'OK' button within the prompt will delete the desired row and all contents within.

Exhibit 4-18 shows the screen capture for the ‘Additional Information and Submitter Requests’ screen:

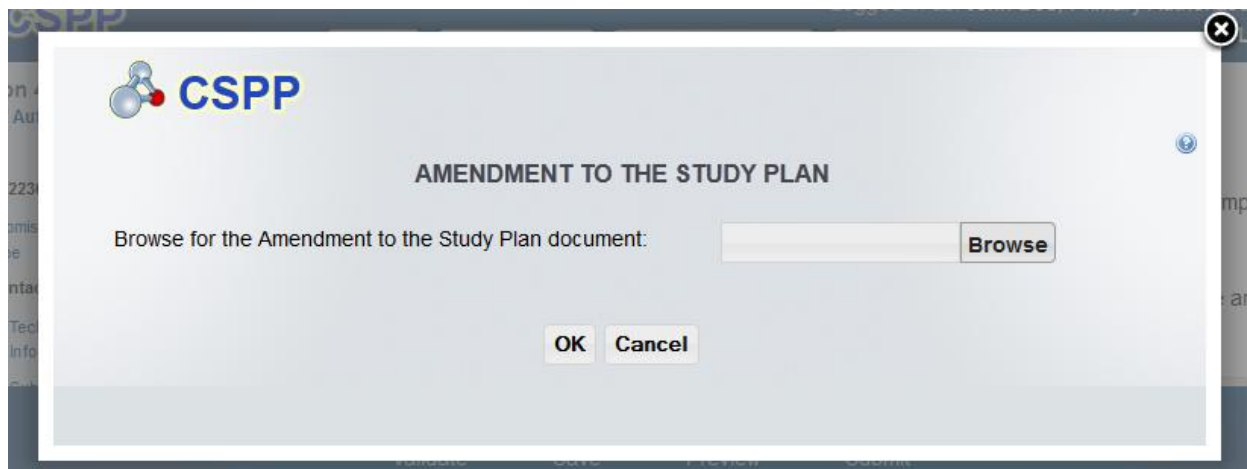
Exhibit 4-18 Additional Information and Submitter Requests Screen



Navigation: Click the ‘Next’ button to navigate to the ‘Test Rules Substantiation Part 1’ screen if information has been claimed CBI. If information has not been claimed CBI and you have not identified other submission information types on the ‘Submission Information Type’ screen, you are ready to submit the Section 4 test rules submission to EPA. If you have identified you are submitting results, you will be taken to the ‘Document Management’ screen under the ‘Results’ folder within the navigation tree.

Exhibit 4-19 shows the screen capture of the ‘Amendment to the Study Plan’ pop-up window:

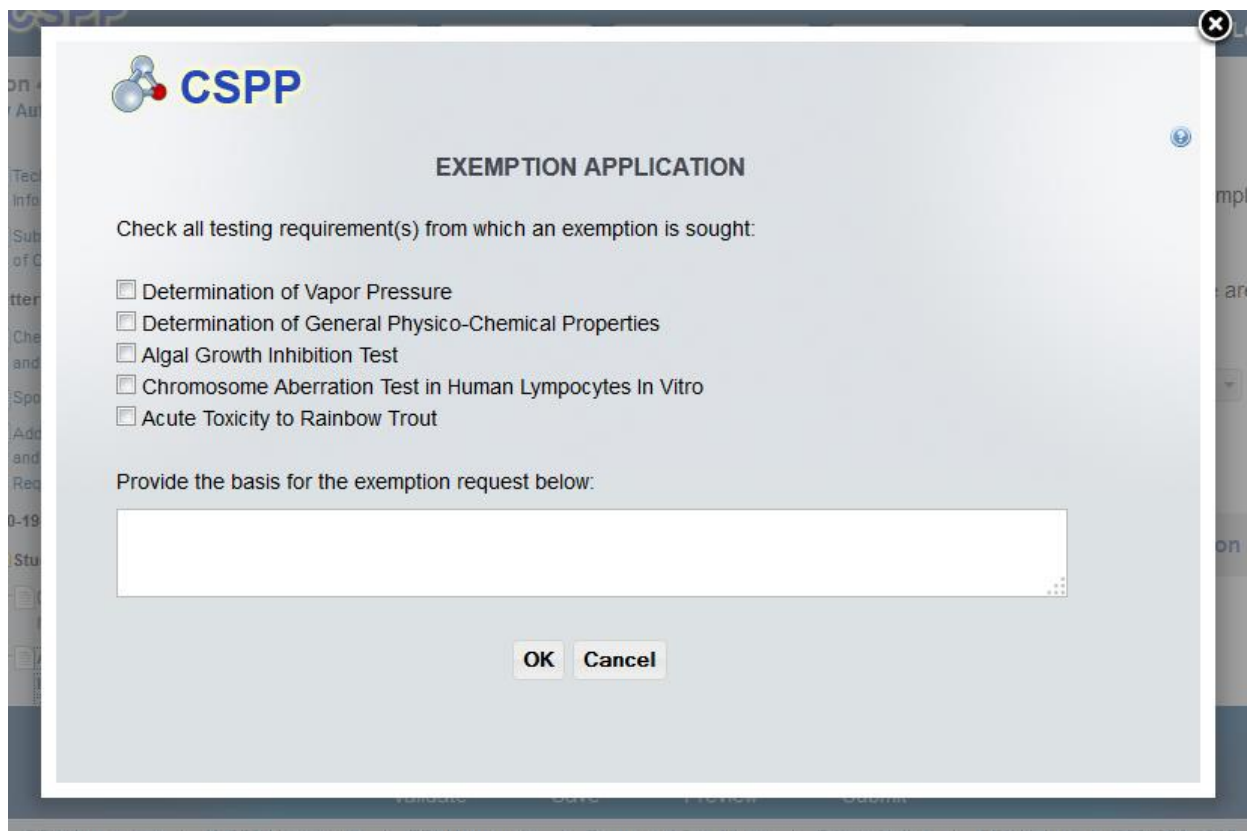
Exhibit 4-19 Amendment to the Study Plan Pop-Up Window



Navigation: Click the ‘OK’ button to return to the ‘Additional Information and Submitter Requests’ screen. Click the ‘Cancel’ or ‘X’ button to close the window and return to the ‘Additional Information and Submitter Requests’ screen without saving the entered information.

Exhibit 4-20 shows the screen capture of the ‘Exemption Application’ pop-up window:

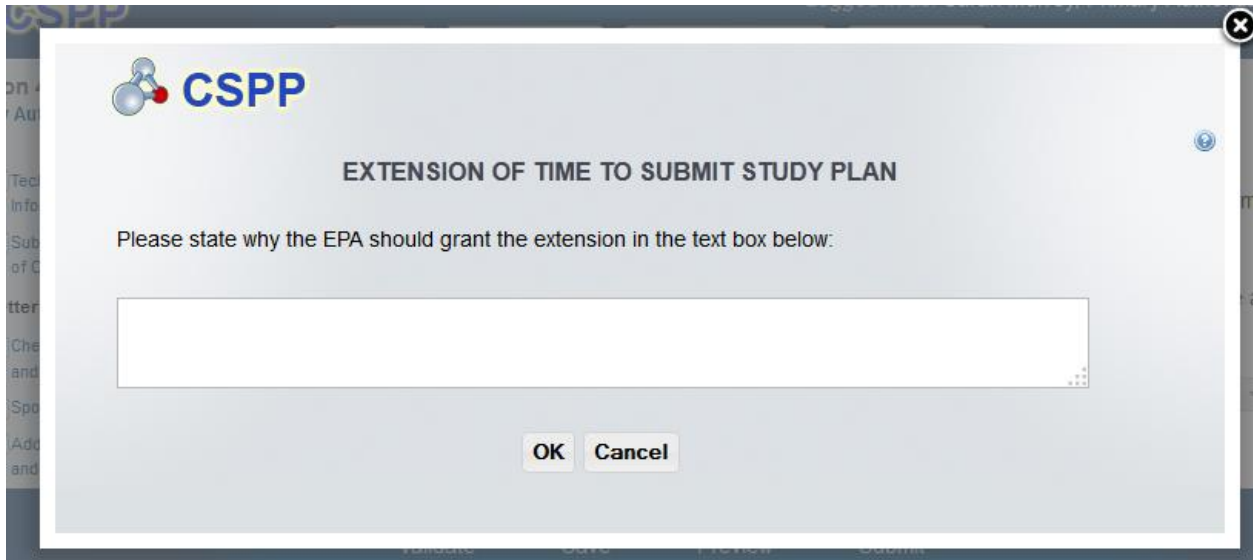
Exhibit 4-20 Exemption Application Pop-Up Window



Navigation: Click the ‘OK’ button to return to the ‘Additional Information and Submitter Requests’ screen. Click the ‘Cancel’ or ‘X’ button to close the window and return to the

‘Additional Information and Submitter Requests’ screen without saving the entered information. Exhibit 4-21 shows the screen capture of the ‘Extension of Time to Submit Study Plan’ pop-up window:

Exhibit 4-21 Extension of Time to Submit Study Plan Pop-Up Window

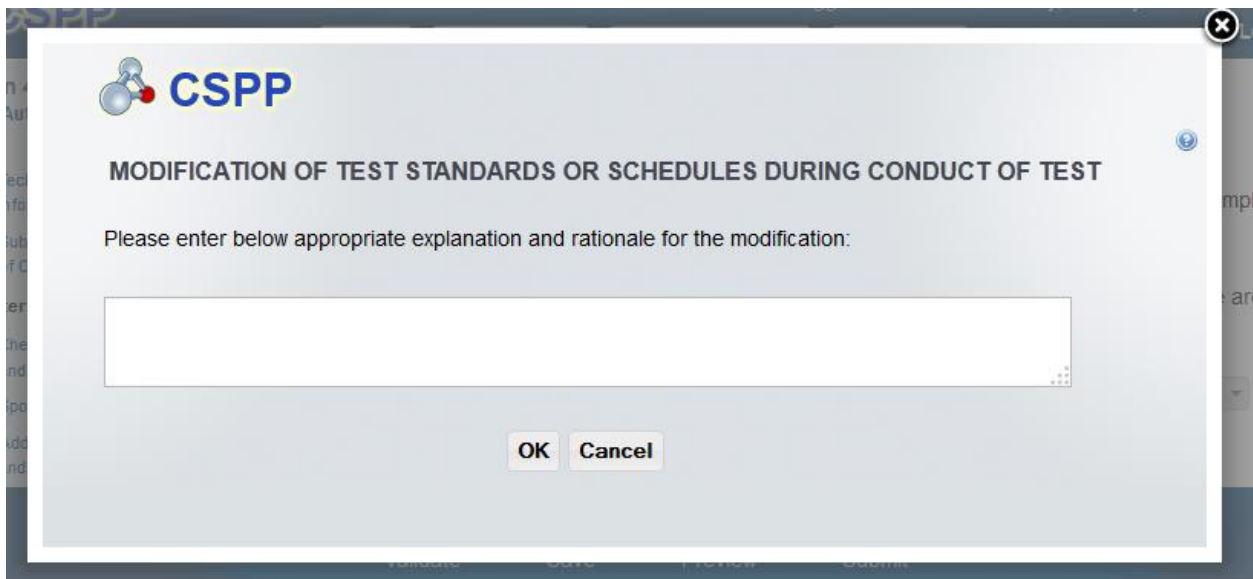


The screenshot shows a pop-up window titled "EXTENSION OF TIME TO SUBMIT STUDY PLAN". The window features the CSPP logo in the top left corner. Below the title, there is a text prompt: "Please state why the EPA should grant the extension in the text box below:". This is followed by a large, empty text input field. At the bottom of the window, there are two buttons: "OK" and "Cancel". The window has a standard close button (an 'X' in a circle) in the top right corner.

Navigation: Click the ‘OK’ button to return to the ‘Additional Information and Submitter Requests’ screen. Click the ‘Cancel’ or ‘X’ button to close the window and return to the ‘Additional Information and Submitter Requests’ screen without saving the entered information.

Exhibit 4-22 shows the screen capture of the ‘Modification of Test Standards or Schedules during Conduct of Test’ pop-up window:

Exhibit 4-22 Modification of Test Standards or Schedules during Conduct of Test Pop-Up Window

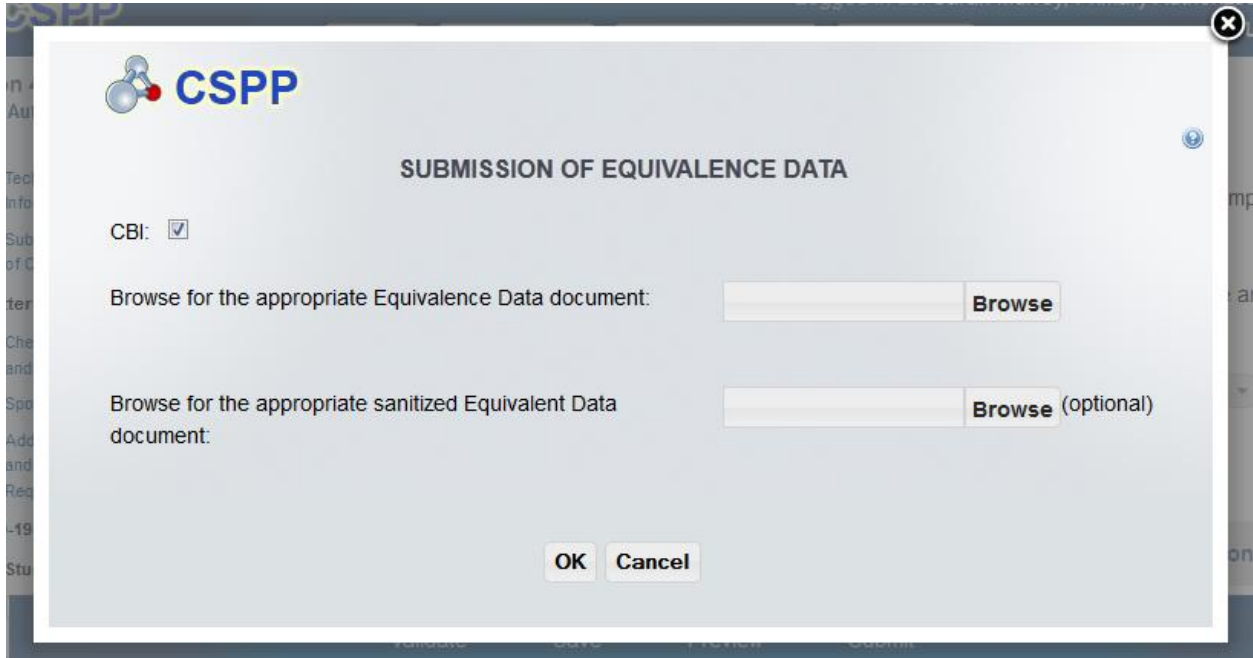


The screenshot shows a pop-up window titled "MODIFICATION OF TEST STANDARDS OR SCHEDULES DURING CONDUCT OF TEST". The window features the CSPP logo in the top left corner. Below the title, there is a text prompt: "Please enter below appropriate explanation and rationale for the modification:". This is followed by a large, empty text input field. At the bottom of the window, there are two buttons: "OK" and "Cancel". The window has a standard close button (an 'X' in a circle) in the top right corner.

Navigation: Click the ‘OK’ button to return to the ‘Additional Information and Submitter Requests’ screen. Click the ‘Cancel’ or ‘X’ button to close the window and return to the ‘Additional Information and Submitter Requests’ screen without saving the entered information.

Exhibit 4-23 shows the screen capture of the ‘Submission of Equivalence Data’ pop-up window:

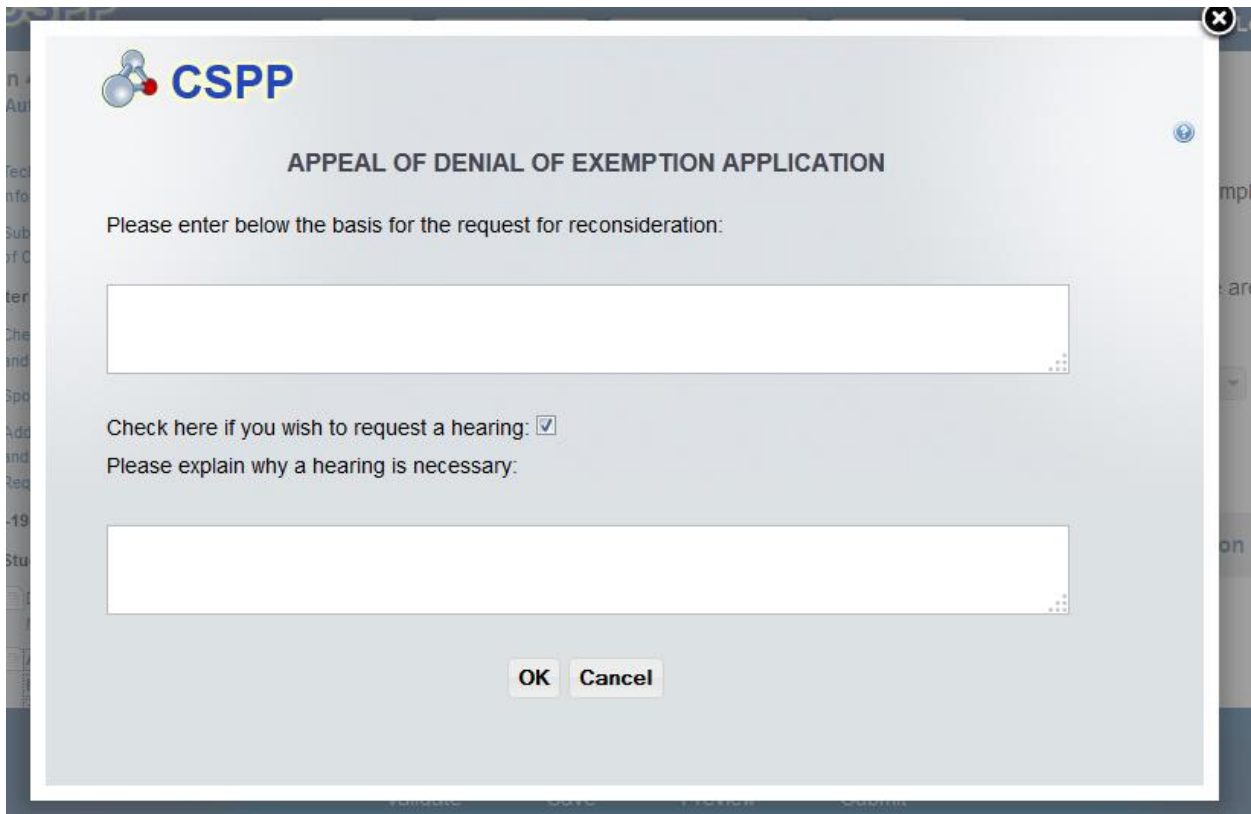
Exhibit 4-23 Submission of Equivalence Data Pop-Up Window



Navigation: Click the ‘OK’ button to return to the ‘Additional Information and Submitter Requests’ screen. Click the ‘Cancel’ or ‘X’ button to close the window and return to the ‘Additional Information and Submitter Requests’ screen without saving the entered information.

Exhibit 4-24 shows the screen capture of the ‘Appeal of Denial of Exemption Application’ pop-up window:

Exhibit 4-24 Appeal of Denial of Exemption Application Pop-Up Window



Navigation: Click the ‘OK’ button to return to the ‘Additional Information and Submitter Requests’ screen. Click the ‘Cancel’ or ‘X’ button to close the window and return to the ‘Additional Information and Submitter Requests’ screen without saving the entered information.

4.7 Test Rules Substantiation

If you claim items within study plan as CBI that require substantiation, you must complete the ‘Test Rules Substantiation Part 1 and Part 2’ screens prior to submission.

4.7.1 Test Rules Substantiation Part 1 Screen

You navigate to the ‘Test Rules Substantiation Part 1’ screen by clicking the ‘Next’ button from the ‘Document Management’ screen or the ‘Additional Information and Submitter Requests’ screen. You can also navigate to the ‘Test Rules Substantiation Part 1’ screen by clicking the ‘Test Rules Substantiation Part 1’ link within the navigation tree.

- **Substantiation Question 1:** Question 1 requires you to identify whether the information has been disclosed in a patent. To complete substantiation for question 1 the system provides ‘Yes’ and ‘No’ radio buttons.
 - Question 1 of the ‘Test Rules Substantiation Part 1’ screen states as follows:
 - Has the information been disclosed in a patent?

- **Substantiation Question 2:** Question 2 requires you to identify whether the disclosure of the study plan information would disclose processes used in the manufacture of a chemical. To complete substantiation for question 2 the system provides ‘Yes’ and ‘No’ radio buttons. If you select the ‘Yes’ radio button, you are required to identify how the process would occur.
 - Question 2 of the ‘Test Rules Substantiation Part 1’ screen states as follows:
 - Would disclosure of the study plan information disclose processes used in the manufacture or processing of a chemical substance or mixture? Describe how this would occur.
- **Substantiation Question 3:** Question 3 requires you to identify whether disclosure of the study plan information would disclose the portion of a mixture. To complete substantiation for question 3 the system provides ‘Yes’ and ‘No’ radio buttons. If you select the ‘Yes’ radio button, you are required to identify how the disclosure would occur.
 - Question 3 of the ‘Test Rules Substantiation Part 1’ screen states as follows:
 - Would disclosure of the study plan information disclose the portion of a mixture comprised by any of the substances in the mixture? Describe how this would occur.
- **Substantiation Question 4:** Question 4 of the ‘Test Rules Substantiation Part 1’ screen requires you to identify if the information claimed as CBI has been disclosed to the public. To complete substantiation for question 4 the system provides ‘Yes’ and ‘No’ radio buttons. If you select the ‘Yes’ radio button, you are required to identify the circumstances in which the information has been disclosed to the public.
 - Question 4 of the ‘Test Rules Substantiation Part 1’ screen states as follows:
 - Has this information been disclosed to the public in any form? Describe the circumstances.

Exhibit 4-25 shows the screen capture of ‘Test Rules Substantiation Part 1’ screen:

Exhibit 4-25: Test Rules Substantiation Part 1 Screen

The screenshot displays the 'TEST RULES SUBSTANTIATION PART 1' screen. The top navigation bar includes 'Home', 'Submissions', 'User Management', 'Resources', and 'Log Out'. The user is logged in as 'John Doe, Primary Authorized Official'. The breadcrumb trail shows 'Section 4 Test Rules > 53 FR 22300 > Study Plan > Test Rules Substantiation Part 1'. The left sidebar contains a tree view with categories like '53 FR 22300', '640-19-7', and '16532-79-9'. The main content area contains four questions:

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.

At the bottom of the form, there are 'Previous' and 'Next' buttons. The 'Next' button is highlighted with a red box. Below the buttons is a dark blue bar with icons for 'Validate', 'Save', 'Preview', and 'Submit'. The footer contains links for 'CDX Homepage', 'MyCDX Homepage', 'EPA Homepage', 'Terms and Conditions', 'Privacy Notice', and 'CDX Helpdesk: (888) 890-1995'.

Navigation: Click the ‘Next’ button to navigate to the ‘Test Rules Substantiation Part 2’ screen. You can also access this screen by clicking the ‘Test Rules Substantiation part 2’ link within the navigation tree.

4.7.2 Test Rules Substantiation Part 2 Screen

You navigate to the ‘Test Rules Substantiation Part 2’ screen by clicking the ‘Next’ button from the ‘Test Rules Substantiation Part 1’ screen. You can also navigate to the ‘Test Rules Substantiation Part 2’ screen by clicking the ‘Test Rules Substantiation Part 2’ link within the navigation tree.

- **Substantiation Question 5:** The system provides a text box to enter information pertaining to question 5. The system also provides a calendar widget to identify the date to which the claim of confidentiality should extend and a text box to provide an explanation as to why confidential treatment should be given. At least one radio button must be selected.
 - Question 5 of the ‘Test Rules Substantiation Part 2’ screen states as follows:
 - For what period of time should confidentiality treatment be given? Until a specific date, the occurrence of a specific event, or permanently? Why should confidential treatment be given?
- **Substantiation Question 6:** The system provides a text box to enter information pertaining to question 6.
 - Question 6 of the ‘Test Rules Substantiation Part 2’ screen states as follows:
 - 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?
- **Substantiation Question 7:** Question 7 requires you to identify what measure you have taken to guard against disclosure of the information and provides a text box to enter information pertaining to the question.
 - Question 7 of the ‘Test Rules Substantiation Part 2’ screen states as follows:
 - What measures have you taken to guard against disclosure of this information to others?
- **Substantiation Question 8:** Question 8 requires you to identify the extent to which the information has been disclosed and the precautions that have been taken to safeguard it. Question 8 provides a text box to enter information pertaining to the question.
 - Question 8 of the ‘Test Rules Substantiation Part 2’ screen states as follows:
 - To what extent has this information been disclosed to others? What precautions have been taken in connection with such disclosures?
- **Substantiation Question 9:** Question 9 requires you to identify if EPA or another Federal Agency or any Federal court has made any confidentiality determinations. If you select the ‘Yes’ radio button, the system requires you to attach such documentation. If you select the ‘No’ radio button, you are not required to provide documentation.
 - Question 9 of the ‘Test Rules Substantiation Part 2’ screen states as follows:

- Has EPA, another Federal Agency, or any Federal court made any pertinent confidentiality determination regarding this information? Click the Browse button and search the appropriate document(s). Click the Upload button to attach copies of such determinations.

Exhibit 4-26 and Exhibit 4-27 show the screen captures of the ‘Test Rules Substantiation Part 2’ screen:

Exhibit 4-26: Test Rules Substantiation Part 2 Screen (Scroll 1)

Section 4 Test Rules
Primary Authorized Official

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

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?

Validate Save Preview Submit

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

Exhibit 4-27: Test Rules Substantiation Part 2 Screen (Scroll 2)

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

Home Submissions User Management Resources

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.

File Name	Attachment Date	Action
Nothing found to display.		

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

Navigation: If you are completing the study plan submission, click the ‘Submit’ icon to begin the submission process. If you have checked the ‘Results’ checkbox on the ‘Submission Information Type’ screen, click the ‘Next’ button to navigate to the ‘Document Management’ screen under the ‘Results’ folder within the navigation tree.

4.8 Results

If you have identified you are submitting results, you are required to complete the following screens to satisfy the reporting requirements.

4.8.1 Document Management Screen

The 'Document Management' screen allows you to upload documents related to Section 4 reporting requirements. To add Section 4 results documentation, click the 'Attach Document' button. Clicking the 'Attach Document' button initiates the 'Document Management' pop-up window.

The 'Document Management' pop-up window allows you to attach documentation by clicking the 'Browse' button. This will generate the 'File Upload' window where you can select the desired document from your hard drive. Once you have identified the document to be attached, click the 'Open' button. Clicking the 'Open' button will close the 'File Upload' window and display the attached document within the 'Browse' field. To claim attached documentation as CBI, check the 'CBI' checkbox. If you claim the attached documentation as CBI you are required to attach the sanitized version of the attached document. To do so, follow the steps outlined above. Once you have attached all desired documentation, click the 'OK' button to return to the 'Document Management' screen.

The 'Document Management' table displays the file name of the document, the date the document was attached, and whether the document was claimed as CBI. The system allows you to delete unwanted documents by clicking the delete icon located under the 'Action' column for a given row. Clicking the file name links within the 'File Name' column generates the 'Document Management' pop-up window. This allows you to view and edit previously attached documents.

- **Attach Document:** Clicking the 'Attach Document' button displays the 'Document Management' pop-up window.
- **File Name:** Displays the name of the attached file. Clicking the file name link generates the 'Document Management' pop-up window and allows you to view the attached document.
- **Attachment Date:** Displays the date the document was attached.
- **CBI:** Displays whether the attached document was claimed as CBI.
- **Test Type:** Displays the test that is associated with the chemical and the result's document.
- **Action:** Clicking the 'Delete' icon will generate a prompt confirming the command. To delete the specified row and all contents within click the 'OK' button within the prompt.

Exhibit 4-28 shows the screen capture of the ‘Results Document Management’ screen:

Exhibit 4-28 Results Document Management Screen

The screenshot shows the 'RESULTS DOCUMENT MANAGEMENT' screen. At the top, it indicates the user is logged in as 'John Doe, Primary Authorized Official'. The navigation menu on the left includes 'Section 4 Test Rules' and 'Primary Authorized Official'. The main content area has a heading 'RESULTS DOCUMENT MANAGEMENT' and a message: 'To add a Results Document, please click the Attach Document button and enter the information in the popup.' Below this is a red-bordered button labeled 'Attach Document'. Underneath is a table with the following columns: 'File Name', 'Attachment Date', 'CBI', 'Test Type', and 'Action'. Below the table is a 'Previous' button. At the bottom of the screen, there is a blue bar with four buttons: 'Validate' (with a green checkmark icon), 'Save' (with a document icon), 'Preview' (with a monitor icon), and 'Submit' (with a document icon and a plus sign). The footer contains links for 'CDX Homepage', 'MyCDX Homepage', 'EPA Homepage', 'Terms and Conditions', 'Privacy Notice', and 'CDX Helpdesk: (888) 890-1995'.

Navigation: Once all documentation has been uploaded, click the ‘Submit’ button to begin the submission process. If multiple chemicals have been selected on the ‘Chemical Identification and Test Rules’ screen, complete all required fields prior to submitting.

The ‘Document Management’ pop-up window allows you to browse for a document from your hard drive and attach one document at a time. You must attach one document. Clicking the ‘Browse’ button will generate the ‘File Upload’ window on your hard drive. Select the document you wish to attach and click the ‘Open’ button. If a document is marked as ‘CBI’ then a sanitized copy of the document can be attached. To save the attached document to the ‘Document Management’ screen, click the ‘OK’ button within the ‘Document Management’ pop-up window.

- **CBI:** Checking the ‘CBI’ checkbox will claim the uploaded document as CBI and generate a sanitized document upload field.
- **Test Type:** Displays the tests that are specific to the chemical that was selected on the ‘Chemical Identification and Test Rules’ screen.
- **Document Upload:** Clicking the ‘Browse’ button will generate the ‘File Upload’ window for you to search for a particular file on your hard drive.
- **Sanitized Document Upload:** This field will only be displayed when the ‘CBI’ checkbox is checked. Clicking the ‘Browse’ button will generate the ‘File Upload’ window for you to search for a particular file on your hard drive.

-
- **Effects:** Select the effect specific to the chemical report. Displays the following options: ‘Health Effects,’ ‘Environmental Fate,’ ‘Ecotoxicity,’ and ‘Physical Chemical Properties.’
 - **Endpoints:** Select an endpoint that is specific to the chemical report. The endpoints will be specific to the effect chosen from the ‘Effects’ drop-down menu. If no effect is chosen, no endpoints will display.

Refer to Appendix B. Submission Types, Effects, and Endpoints for a complete list of values that are found within each of the drop-down menus on the ‘Document Management’ pop-up window. If you have multiple documents that share the same effects and/or endpoints, you must upload each individual document separately and select values from the drop-down menus.

Exhibit 4-29 shows the screen capture of the ‘Results Document Management’ pop-up window:

Exhibit 4-29 Results Document Management Pop-Up Window

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**

Navigation: Click the ‘OK’ button to return to the ‘Results Document Management’ screen.

4.9 Optional Substantiation

If you have claimed any information other than a chemical or a document as CBI, and wish to upload a substantiation version, you can click on the ‘Document Management’ link within the navigation tree.

4.9.1 Optional Document Management Screen

You can access this screen by clicking the ‘Document Management’ link under the ‘Optional Substantiation’ folder.

You can upload substantiation documentation by clicking the ‘Attach Document’ button.

The ‘Document Management’ pop-up window allows you to attach documentation by clicking the ‘Browse’ button. This will generate the ‘File Upload’ window where you can select the desired document from your hard drive. Once you have identified the document to be attached, click the ‘Open’ button. Clicking the ‘Open’ button will close the ‘File Upload’ window and display the attached document within the ‘Browse’ field. Once you have attached all desired documentation, click the ‘OK’ button to return to the ‘Optional Document Management’ screen.

The optional document management table displays the file name of the document and the date the document was attached. The system allows you to delete unwanted documents by clicking

the ‘Delete’ icon located under the ‘Action’ column for a given row. Clicking the file name links within the ‘File Name’ column generates the ‘Document Management’ pop-up window. This allows you to view and edit previously attached documents.

- **Attach Document:** Clicking the ‘Attach Document’ button displays the ‘Optional Substantiation Document Management’ pop-up window.
- **File Name:** Displays the name of the attached file. Clicking the file name link generates the ‘Optional Substantiation Document Management’ pop-up window and allows you to view the attached document.
- **Attachment Date:** Displays the date the document was attached.
- **Action:** Clicking the ‘Delete’ icon (✖) will generate a prompt confirming the command. To delete the specified row and all contents within click the ‘OK’ button within the prompt.

Exhibit 4-30 shows the screen capture of the ‘Optional Document Management’ screen:

Exhibit 4-30: Optional Document Management Screen

The screenshot displays the 'Optional Document Management' screen. At the top, the user is logged in as 'John Doe, Primary Authorized Official'. The navigation menu on the left includes 'Section 4 Test Rules' and 'Optional Substantiation'. The main content area features a red-bordered 'Attach Document' button. Below it is a table with the following structure:

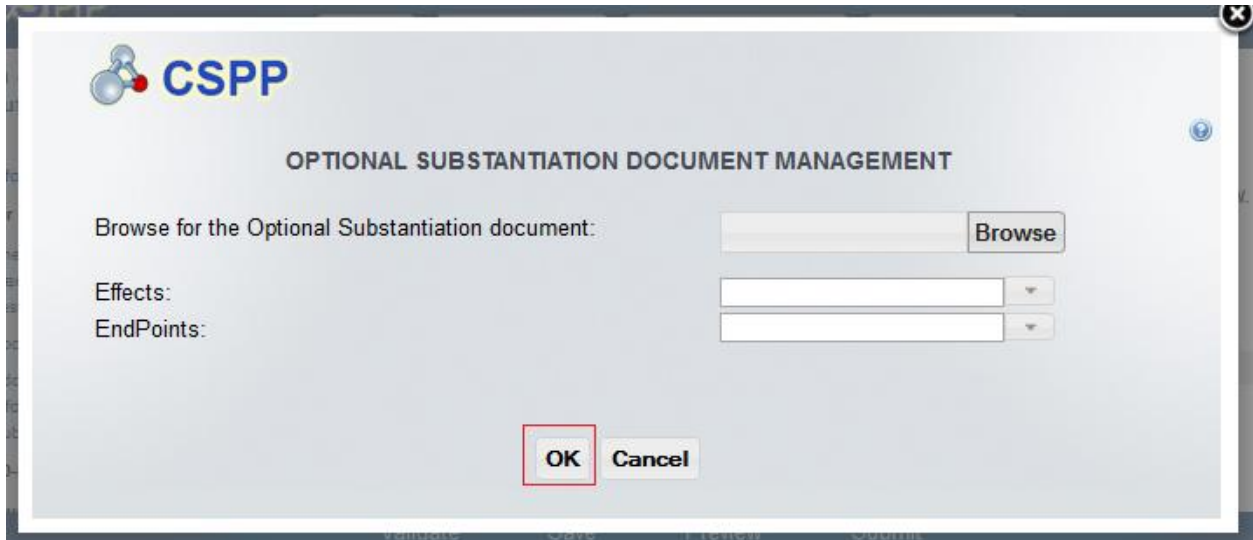
File Name	Attachment Date	Action
-----------	-----------------	--------

At the bottom of the screen, there are four icons: 'Validate' (checkmark), 'Save' (floppy disk), 'Preview' (monitor), and 'Submit' (submit button). The footer contains links to 'CDX Homepage', 'MyCDX Homepage', 'EPA Homepage', 'Terms and Conditions', 'Privacy Notice', and 'CDX Helpdesk: (888) 890-1995'.

Navigation: Click the ‘Attach Document’ button to generate the ‘Optional Substantiation Document Management’ pop-up window. Once all documents have been uploaded and you have completed all required screen you may click the ‘Submit’ icon and begin the submission process.

Exhibit 4-31 shows the screen capture of the ‘Optional Substantiation Document Management’ pop-up window.

Exhibit 4-31: Optional Substantiation Document Management Pop-Up Window



5 Enforceable Consent Agreements (ECAs)

5.1 Submission Information Type Screen

You can access the ‘Submission Information Type’ screen by clicking the ‘Next’ button from either the ‘Create New Passphrase’ or the ‘Enter Passphrase’ screen. Once you are within the ‘Enforceable Consent Agreement’ flow of the Section 4 application, the ‘Submission Information Type’ screen can also be accessed by clicking the ‘Submission Information Type’ link located within the navigation tree.

The ‘Submission Information Type’ screen allows you to select the submission information type that you will be submitting to EPA. This screen consists of the following fields:

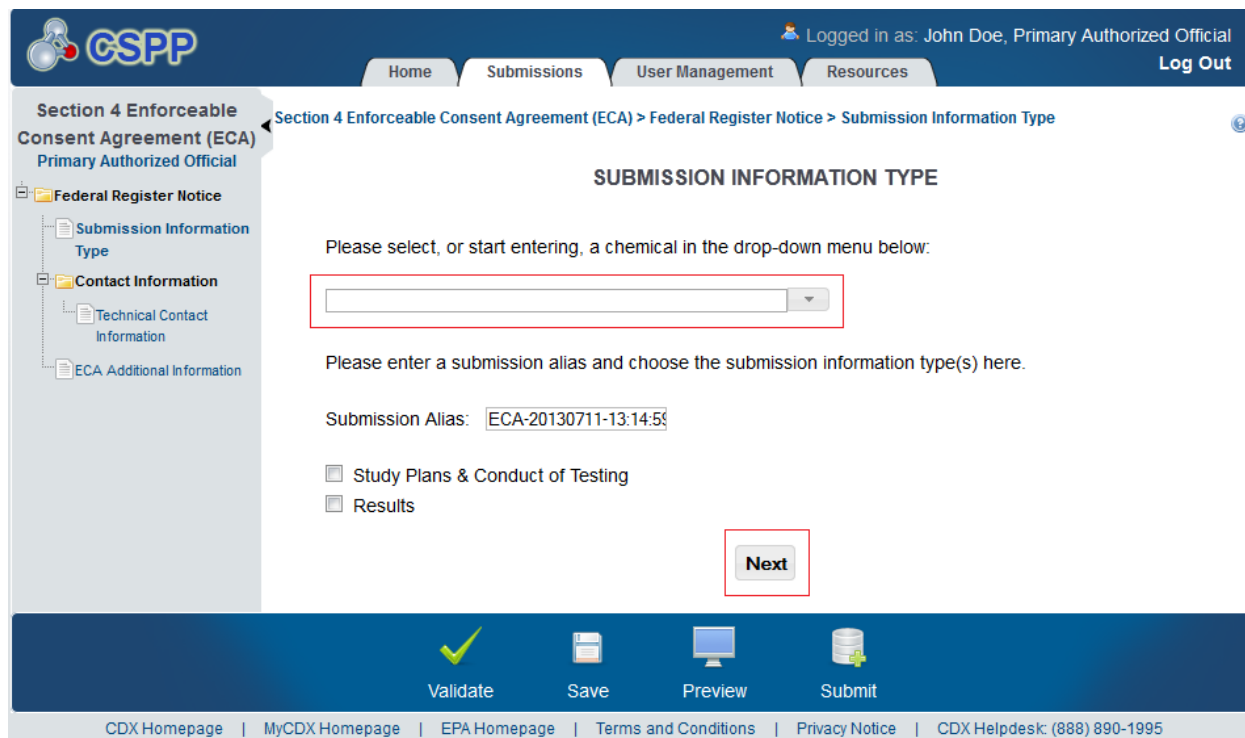
- **Please select, or start entering, a chemical in the drop-down menu below:** Displays all chemicals associated with ECA. This field is required.
- **Submission Alias:** This field is auto-populated with a date and time stamp and is displayed on the ‘Section 4 Test Rules, ECAs, MOUs, and 40 CFR 766’ submissions screen within the ‘Submission Alias’ column. The ‘Submission Alias’ is also used to differentiate between submission types within the ‘User Management’ screen. This field is editable should you choose to provide your own submission alias.
- **Study Plans & Conduct of Testing:** Checking the ‘Study Plans & Conduct of Testing’ checkbox identifies that you are fulfilling the requirements to submit a study plan to EPA. Once you have checked the ‘Study Plans & Conduct of Testing’ checkbox and either saved the information on this screen by clicking the ‘Save’ icon or by clicking the ‘Next’ button, the ‘Study Plans & Conduct of Testing’ folder is generated within the navigation tree allowing you to access the study plan screen required to be completed.
- **Results:** Checking the ‘Results’ checkbox indicates that you are fulfilling the requirements to submit results to EPA. Once you have checked the ‘Results’ checkbox and either saved the information on this screen by clicking the ‘Save’ icon or by clicking the ‘Next’ button, the ‘Results’ folder is generated within the navigation tree allowing you to access the results screens required to be completed.

If you have previously identified a submission type, have accessed the submission type screens, and then unchecked the previously checked submission information type (i.e. Study Plans), you will receive a prompt indicating that all associated information will be lost. Clicking the ‘OK’ button within the prompt will delete all data associated with that submission information type.

Once you have identified the type(s) of information you will be submitting to EPA, you are able to access the screens required to be completed.

Exhibit 5-1 shows the screen capture of the ‘Submission Information Type’ screen:

Exhibit 5-1 Submission Information Type Screen



Navigation: Click the ‘Next’ button to navigate to the ‘Technical Contact Information’ screen. You can also access this screen by clicking the ‘Technical Contact Information’ link within the navigation tree.

5.2 Technical Contact Information Screen

The ‘Technical Contact Information’ screen allows you to submit on behalf of another company. To identify if you are submitting on behalf of another company, the system provides the radio button ‘This is a submission on behalf of another company.’ The system will not allow you to select more than one radio button per submission.

If you are submitting on behalf of another company, select the ‘This is a submission on behalf of another company’ radio button. This functionality has been implemented to accommodate a company that may use an agent or another individual. If this radio button is selected, the information of the contractor/consultant’s company is to be entered into the ‘Submitting on Behalf Of Company’ screen. If you are not submitting on behalf of another company and have selected the ‘N/A’ radio button, you are taken to the ‘Principal Test Sponsor’ screen.

- **N/A:** Selecting the ‘N/A’ radio button allows you to signify that you are not making a submission on behalf of a company.
- **This is a submission on behalf of another company:** Selecting the ‘This is a submission on behalf of another company’ radio button allows you to identify yourself as an individual submitting on behalf of another company.
- **Copy CDX Registration:** Clicking the ‘Copy CDX Registration’ button will automatically populate the fields with your contact information provided during CDX registration.

-
- **Prefix:** The ‘Prefix’ drop-down menu displays all optional prefixes. If applicable, select the appropriate prefix. If you have clicked the ‘Copy CDX Registration’ button, the fields populate with information entered during CDX registration.
 - **First Name:** Enter the technical contact’s first name. If you have clicked the ‘Copy CDX Registration’ button, the fields populate with information entered during CDX registration. This field is required.
 - **Middle Initial:** Enter the technical contact’s middle initial. If you have clicked the ‘Copy CDX Registration’ button, the fields populate with information entered during CDX registration.
 - **Last Name:** Enter the technical contact’s last name. If you have clicked the ‘Copy CDX Registration’ button, the fields populate with information entered during CDX registration. This field is required.
 - **Suffix:** The ‘Suffix’ drop-down menu displays all optional suffixes. If applicable, select the appropriate suffix. If you have clicked the ‘Copy CDX Registration’ button, the fields populate with information entered during CDX registration.
 - **Company Name:** Enter the technical contact’s company name. If you have clicked the ‘Copy CDX Registration’ button, the fields populate with information entered during CDX registration. This field is required.
 - **Phone Number:** Enter the daytime phone number of the manufacturing or processing establishment contact for whom the submission is being made, U.S. or international. This field does not accept dashes. If you have clicked the ‘Copy CDX Registration’ button, the fields populate with information entered during CDX registration. This field is required.
 - **Ext:** Enter the extension to the daytime phone number of the manufacturing or processing establishment contact for whom the submission is being made.
 - **Email Address:** Enter the technical contact’s email address. The email address must follow the standard email address format: john.doe@abc.com. If you have clicked the ‘Copy CDX Registration’ button, the fields populate with information entered during CDX registration. This field is required.
 - **Mailing Address 1:** Enter the first line of the technical contact’s mailing address. If you have clicked the ‘Copy CDX Registration’ button, the fields populate with information entered during CDX registration. This field is required.
 - **Mailing Address 2:** Enter the second line of the technical contact’s mailing address. If you have clicked the ‘Copy CDX Registration’ button, the fields populate with information entered during CDX registration.
 - **City:** Enter the city of the technical contact’s mailing address. If you have clicked the ‘Copy CDX Registration’ button, the fields populate with information entered during CDX registration. This field is required.
 - **State:** Enter the state of the technical contact’s mailing address. If you have clicked the ‘Copy CDX Registration’ button, the fields populate with information entered during CDX registration. This is a required field if the ‘Country’ field is defined as ‘United States.’

- **Postal Code:** Enter the postal code of the technical contact's mailing address. If you have clicked the 'Copy CDX Registration' button, the fields populate with information entered during CDX registration. This field is required.
- **Country:** Displays all available countries. Enter the country of the technical contact's mailing address. If you have clicked the 'Copy CDX Registration' button, the fields populate with information entered during CDX registration. This is a required field.

Exhibit 5-2 shows the screen capture of the ‘Technical Contact Information’ screen:

Exhibit 5-2 Technical Contact Information Screen

The screenshot displays the 'Technical Contact Information' screen. At the top, the user is logged in as 'John Doe, Primary Authorized Official'. The breadcrumb trail indicates the path: 'Section 4 Enforceable Consent Agreement (ECA) > 68 FR 33125 > Contact Information > Technical Contact Information'. The main heading is 'TECHNICAL CONTACT INFORMATION'. Below this, instructions state: '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.' Two radio buttons are present: 'N/A' and 'This is a submission on behalf of another company'. A button labeled 'Copy CDX Registration' is also visible. The form includes fields for CBI, Prefix, First Name, Middle Initial, Last Name, Suffix, Company Name, Phone Number (with an extension field), Email Address, Mailing Address 1, Mailing Address 2, City, State, Postal Code, and Country. At the bottom of the form area, there are 'Previous' and 'Next' buttons, with the 'Next' button highlighted by a red box. The footer contains navigation links: 'CDX Homepage | MyCDX Homepage | EPA Homepage | Terms and Conditions | Privacy Notice | CDX Helpdesk: (888) 890-1995'.

Navigation: If you are submitting on behalf of another company, select the ‘This is a submission on behalf of another company’ radio button and click the ‘Next’ button to navigate to the ‘Submitting on Behalf of Company’ screen. If you are submitting study plan information and are not submitting on behalf of another entity, you are taken to the ‘Principal Test Sponsor’ screen. If you are not submitting either study plan information or results information and are not

submitting on behalf of another entity, the application navigates to the ‘ECA Additional Information’ screen.

5.3 Submitting on Behalf of Company Screen

You can access the ‘Submitting on Behalf of Company’ screen by selecting the ‘This is a submission on behalf of another company’ radio button and by clicking the ‘Next’ button from the ‘Technical Contact Information’ screen. You can also access this screen by clicking the ‘Submitting on Behalf of Company’ link within the navigation tree.

The ‘Submitting on Behalf of Company’ screen presents a list of fields that allow you to input the contact information for the manufacturing or processing establishment on whose behalf the submission is being made.

- **CBI:** Checking this checkbox will claim the information as CBI.
- **Prefix:** The ‘Prefix’ drop-down menu displays all optional prefixes. If applicable, select the appropriate prefix.
- **First Name:** Enter the first name of the manufacturing or processing establishment contact for which the submission is being made. This field is required.
- **Middle Initial:** Enter the middle initial of the manufacturing or processing establishment contact for which the submission is being made.
- **Last Name:** Enter the last name of the manufacturing or processing establishment contact for which the submission is being made. This field is required.
- **Suffix:** The ‘Suffix’ drop-down menu displays all optional suffixes and captures the suffix of the manufacturing or processing establishment contact for which the submission is being made. If applicable, select the appropriate suffix.
- **Company Name:** Enter the company name of the manufacturing or processing establishment. This field is required.
- **Phone Number:** Enter the daytime phone number of the manufacturing or processing establishment contact for whom the submission is being made; U.S. or international. This is a required field.
- **Ext:** Enter the extension to the daytime phone number of the manufacturing or processing establishment contact for whom the submission is being made.
- **Email Address:** Enter the email address of the manufacturing or processing establishment contact for which the submission is being made. The email address must follow the standard email address format: john.doe@abc.com. This field is required.
- **Mailing Address 1:** Enter the first line of the mailing address of the manufacturing or processing establishment contact. This field is required.
- **Mailing Address 2:** Enter the second line of the mailing address of the manufacturing or processing establishment contact.
- **City:** Enter the city of the mailing address of the manufacturing or processing establishment contact. This field is required.

- **State:** Enter the state of the mailing address of the manufacturing or processing establishment contact. This is a required field if the ‘Country’ field is defined as ‘United States.’
- **Postal Code:** Enter the postal code of the mailing address of the manufacturing or processing establishment contact. This field is required.
- **Country:** Enter the country of the mailing address of the manufacturing or processing establishment contact. This field is required.

Exhibit 5-3 shows the screen capture of the ‘Submitting on Behalf of Company’ screen:

Exhibit 5-3 Submitting on Behalf of Company Screen

The screenshot displays the GSPP (Global Study Plan Portal) interface. At the top, the user is logged in as 'John Doe, Primary Authorized Official'. The navigation menu on the left includes 'Section 4 Enforceable Consent Agreement (ECA) Primary Authorized Official' and '68 FR 33125'. The main content area is titled 'SUBMITTING ON BEHALF OF COMPANY' and contains the following form fields:

- CBI:** A checkbox.
- Prefix:** A dropdown menu.
- First Name:** A text input field.
- Middle Initial:** A text input field.
- Last Name:** A text input field.
- Suffix:** A dropdown menu.
- Company Name:** A text input field.
- Phone Number:** Two text input fields for main number and extension (Ext.). A note below states: '(Do not enter any dashes (-) in Phone Number field above)'
- Email Address:** A text input field.
- Mailing Address 1:** A text input field. Below it is the label 'Street address, P.O. box, company name, etc.'
- Mailing Address 2:** A text input field. Below it is the label 'Apartment, suite, etc.'
- City:** A text input field.
- State:** A dropdown menu.
- Postal Code:** A text input field.
- Country:** A dropdown menu.

At the bottom of the form, there are two buttons: 'Previous' and 'Next'. The 'Next' button is highlighted with a red rectangular box. Below the form, there is a blue bar with four icons and labels: 'Validate' (checkmark), 'Save' (floppy disk), 'Preview' (computer monitor), and 'Submit' (database icon with plus sign). At the very bottom, a footer contains links: 'CDX Homepage | MyCDX Homepage | EPA Homepage | Terms and Conditions | Privacy Notice | CDX Helpdesk: (888) 890-1995'.

Navigation: If you are submitting study plans click the ‘Next’ button to navigate to the ‘Principal Test Sponsor’ screen. If you only checked the ‘Results’ checkbox, click the ‘Next’

button to navigate to the ‘Document Management’ screen under the ‘Results’ folder within the navigation tree. If you are only completing the letter of intent, click the ‘Submit’ icon located in the action bar to begin the submission process.

5.4 Study Plans

If you have identified you are submitting information for study plans, you are required to complete the following screens to satisfy the reporting requirements.

5.4.1 Principal Test Sponsor Screen

The ‘Principal Test Sponsor’ screen presents a list of fields that allow you to input the contact information for the principal test sponsor.

- **CBI:** Check the ‘CBI’ checkbox to claim the principal test sponsor contact information as CBI.
- **Prefix:** The ‘Prefix’ drop-down menu displays all optional prefixes. If applicable, select the appropriate prefix.
- **First Name:** Enter the first name of the principal test sponsor contact. This field is required.
- **Middle Initial:** Enter the middle initial of the principal test sponsor contact.
- **Last Name:** Enter the last name of the principal test sponsor contact. This field is required.
- **Suffix:** The ‘Suffix’ drop-down menu displays all optional suffixes and captures the suffix of the principal test sponsor contact. If applicable, select the appropriate suffix.
- **Company Name:** Enter the company name of the principal test sponsor contact. This field is required.
- **Phone Number:** Enter the daytime phone number of the principal test sponsor contact; U.S. or international. This field is required.
- **Ext:** Enter the extension to the daytime phone number of the principal test sponsor contact.
- **Email Address:** Enter the email address of principal test sponsor contact. The email address must follow the standard email address format: john.doe@abc.com. This field is required.
- **Mailing Address 1:** Enter the first line of the mailing address of the principal test sponsor’s company. This field is required.
- **Mailing Address 2:** Enter the second line of the mailing address of the principal test sponsor’s company.
- **City:** Enter the city of the mailing address of the principal test sponsor’s company. This field is required.
- **State:** Enter the state of the mailing address of the principal test sponsor contact’s company. This is a required field if the ‘Country’ field is defined as ‘United States.’
- **Postal Code:** Enter the postal code of the principal test sponsor contact’s company. This field is required.
- **Country:** Enter the country of the mailing address of principal test sponsor contact’s company. This field is required.

Exhibit 5-4 shows the screen capture of the ‘Principal Test Sponsor’ screen:

Exhibit 5-4 Principal Test Sponsor Screen

The screenshot displays the 'Principal Test Sponsor' form within the GSPSP application. The breadcrumb trail indicates the path: Section 4 Enforceable Consent Agreement (ECA) > 68 FR 33125 > Study Plans & Conduct of Testing > Principal Test Sponsor. The form is titled 'PRINCIPAL TEST SPONSOR' and includes the instruction: 'Fill out the information below for the Principal Test Sponsor.' The form fields are as follows:


- 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:

At the bottom of the form, there are two buttons: 'Previous' and 'Next'. The 'Next' button is highlighted with a red border. Below the form, there is a navigation bar with icons and labels for 'Validate', 'Save', 'Preview', and 'Submit'. The footer contains the following links: CDX Homepage | MyCDX Homepage | EPA Homepage | Terms and Conditions | Privacy Notice | CDX Helpdesk: (888) 890-1995.

Navigation: Click the ‘Next’ button to navigate to the ‘Principal Sponsor Organization’ screen. You can also access this screen by clicking the ‘Principal Sponsor Organization’ link within the navigation tree.

5.4.2 Principal Sponsor Organization Screen

The ‘Principal Sponsor Organization’ screen allows you to enter the contact information for each administrative official and project manager within the principal test sponsor’s organization. To add each professional click the ‘Add Sponsor Organization’ button. This will generate a new set of text fields. Once you have entered data into one field set, you must click the ‘Save’ button located within that field set prior to adding another member. If the data previously entered and

saved within a field set needs to be edited, you can click the ‘Edit’ button located in the upper right corner of that field set. If a field set needs to be deleted, you can click the ‘Delete’ icon () that is also located in the upper right corner of each field set.

The system allows you to expand/collapse each field set by clicking the field set header. If you wish to expand/collapse all field sets simultaneously, click the ‘Expand All’ or ‘Collapse All’ links located at the top right of all field sets.

- **CBI:** Check the checkbox to claim the member contact information as CBI.
- **Prefix:** The ‘Prefix’ drop-down menu displays all optional prefixes and captures the member contact’s prefix. If applicable, select the appropriate prefix.
- **First Name:** Enter the member contact’s first name. This field is required.
- **Middle Initial:** Enter the member contact’s middle initial.
- **Last Name:** Enter the member contact’s last name. This field is required.
- **Suffix:** The ‘Suffix’ drop-down menu displays all optional suffixes and captures the member contact’s suffix. If applicable, select the appropriate suffix.
- **Company Name:** Enter the member contact’s company name. This field is required.
- **Job Title:** Enter the member contact’s job title. This field is required.
- **Phone Number:** Enter the member contact’s daytime phone number; U.S. or international. This field is required.
- **Ext:** Enter the extension to the contact’s daytime phone number.
- **Email Address:** Enter the member contact’s email address. The email address must follow the standard email address format: john.doe@abc.com. This field is required.
- **Mailing Address 1:** Enter the first line of the member contact’s mailing address. This field is required.
- **Mailing Address 2:** Enter the second line of the member contact’s mailing address.
- **City:** Enter the city of the member contact’s mailing address. This is a required field.
- **State:** Enter the state of the member contact’s mailing address. This field is required when ‘United States’ is identified as the country.
- **Postal Code:** Enter the postal code of the member contact’s mailing address. This field is required.
- **Country:** Enter the country of the mailing address of the contact. This is a required field.

Exhibit 5-5 shows the screen capture of the ‘Principal Sponsor Organization’ screen:

Exhibit 5-5 Principal Sponsor Organization Screen

Section 4 Enforceable Consent Agreement (ECA) Primary Authorized Official

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

Validate Save Preview Submit

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Navigation: Click the ‘Next’ button to navigate to the ‘Testing Facilities’ screen. You can also access this screen by clicking the ‘Testing Facilities’ link within the navigation tree.

5.4.3 Testing Facilities Screen

The ‘Testing Facilities’ screen allows you to enter the contact information for each facility responsible for testing as well as that facility’s contact. To add each facility and contact, click the ‘Add Testing Facility’ button. This will generate a new set of text fields. Once you have entered

data into one field set, you must click the ‘Save’ button located within that field set prior to adding another facility and contact. If the data previously entered and saved within a field set needs to be edited, you can click the ‘Edit’ button located in the upper right corner of that field set. If a facility needs to be deleted, you can click the ‘Delete’ icon (✖) that is also located in the upper right corner of each field set.

The system allows you to expand/collapse each field set by clicking the field set header. If you wish to expand/collapse all field sets simultaneously, click the ‘Expand All’ or ‘Collapse All’ links located at the top right of all field sets.

- **Testing Facility:** Enter the name of the testing facility. This field is required.
- **Phone Number:** Enter the phone number of the testing facility. This field is required.
- **Ext:** Enter the extension to the phone number of the testing facility.
- **Mailing Address 1:** Enter the first line of the mailing address of the testing facility. This field is required.
- **Mailing Address 2:** Enter the second line of the mailing address of the testing facility.
- **City:** Enter the city of the testing facility’s mailing address. This is a required field.
- **State:** Enter the state of the testing facility’s mailing address. This field is required when ‘United States’ is identified as the country.
- **Postal Code:** Enter the postal code of the testing facility’s mailing address. This field is required.
- **Country:** Enter the country of the mailing address of the testing facility. This is a required field.
- **Select the testing facility contact role:** Select the job title of the contact of the testing facility from the drop-down menu. This drop-down menu contains the following values: ‘Administrative Official,’ and ‘Project Manager.’ This field is required.
- **CBI:** Check the checkbox to claim the contact information of the testing facility as CBI.
- **Prefix:** The ‘Prefix’ drop-down menu displays all optional prefixes and captures the testing facility’s contact’s prefix. If applicable, select the appropriate prefix.
- **First Name:** Enter the testing facility contact’s first name. This field is required.
- **Middle Initial:** Enter the testing facility contact’s middle initial.
- **Last Name:** Enter the testing facility contact’s last name. This field is required.
- **Suffix:** The ‘Suffix’ drop-down menu displays all optional suffixes and captures the testing facility contact’s suffix. If applicable, select the appropriate suffix.
- **Phone Number:** Enter the testing facility contact’s daytime phone number; U.S. or international. This field is required.
- **Ext:** Enter the extension to the testing facility contact’s daytime phone number.
- **Email Address:** Enter the testing facility contact’s email address. The email address must follow the standard email address format: john.doe@abc.com. This field is required.
- **Mailing Address 1:** Enter the first line of the testing facility contact’s mailing address. This field is required.

-
- **Mailing Address 2:** Enter the second line of the testing facility contact's mailing address.
 - **City:** Enter the city of the testing facility contact's mailing address. This is a required field.
 - **State:** Enter the state of the testing facility contact's mailing address. This field is required when 'United States' is identified as the country.
 - **Postal Code:** Enter the postal code of the testing facility contact's mailing address. This field is required.
 - **Country:** Enter the country of the mailing address of the testing facility's contact. This is a required field.

Exhibit 5-6 and Exhibit 5-7 shows the screen capture of the ‘Testing Facilities’ screen:

Exhibit 5-6 Testing Facilities Screen (Scroll 1)

Section 4 Enforceable Consent Agreement (ECA)
Primary Authorized Official

68 FR 33125

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

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**

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Exhibit 5-7: Testing Facilities Screen (Scroll 2)

The screenshot displays the GSPP (Global Site Profile) interface. At the top, the user is logged in as John Doe, a Primary Authorized Official. The navigation menu on the left includes 'Section 4 Enforceable Consent Agreement (ECA) Primary Authorized Official' and a tree view for submission 68 FR 33125, with 'Testing Facilities' highlighted. The main form area contains the following fields:

- 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:

At the bottom of the form is a **Save** button. Below the form, a message states: "Click the **Add Testing Facility** button to add a new testing facility." Below this message are **Previous** and **Next** buttons. The footer contains icons for **Validate**, **Save**, **Preview**, and **Submit**, along with a navigation bar with links: CDX Homepage | MyCDX Homepage | EPA Homepage | Terms and Conditions | Privacy Notice | CDX Helpdesk: (888) 890-1995.

Navigation: Click the 'Next' button to navigate to the 'Study Professionals' screen. You can also access this screen by clicking the 'Study Professionals' link within the navigation tree.

5.4.4 Study Professionals Screen

The 'Study Professionals' screen allows you to provide the contact information and a summary of the training and experience of each professional involved in the study. To add each professional click the 'Add Study Professional' button. This will generate a new set of text fields. Once you have entered data into one field set, you must click the 'Save' button located within that field set prior to adding another professional. If the data previously entered and saved within a field set needs to be edited, you can click the 'Edit' button located in the upper right

corner of that field set. If a professional needs to be deleted, you can click the ‘Delete’ icon (✖) that is also located in the upper right corner of each field set.

The system allows you to expand/collapse each field set by clicking the field set header. If you wish to expand/collapse all field sets simultaneously, click the ‘Expand All’ or ‘Collapse All’ links located at the top right of all field sets.

- **Select the Study Professional Role:** Select the study professional’s role from the drop-down menu. This drop-down menu contains the following values: ‘Chemist,’ ‘Laboratory Assistant,’ ‘Microbiologist,’ ‘Pathologist,’ ‘Study Director,’ ‘Toxicologist,’ ‘Veterinarian,’ and ‘Other.’ This field is required.
- **CBI:** Check the checkbox to claim the professional’s contact information as CBI.
- **Prefix:** The ‘Prefix’ drop-down menu displays all optional prefixes and captures the professional’s prefix. If applicable, select the appropriate prefix.
- **First Name:** Enter the professional’s first name. This field is required.
- **Middle Initial:** Enter the professional’s middle initial.
- **Last Name:** Enter the professional’s last name. This field is required.
- **Suffix:** The ‘Suffix’ drop-down menu displays all optional suffixes and captures the professional’s suffix. If applicable, select the appropriate suffix.
- **Company Name:** Enter the professional’s company name. This field is required.
- **Phone Number:** Enter the professional’s daytime phone number; U.S. or international. This field is required.
- **Ext:** Enter the extension to the professional’s daytime phone number.
- **Email Address:** Enter the professional’s email address. The email address must follow the standard email address format: john.doe@abc.com. This field is required.
- **Mailing Address 1:** Enter the first line of the professional’s mailing address. This field is required.
- **Mailing Address 2:** Enter the second line of the professional’s mailing address.
- **City:** Enter the city of the professional’s mailing address. This is a required field.
- **State:** Enter the state of the professional’s mailing address. This field is required when ‘United States’ is identified as the country.
- **Postal Code:** Enter the postal code of the professional’s mailing address. This field is required.
- **Country:** Enter the country of the professional’s mailing address. This is a required field.
- **Experience Summary:** Enter the summary of the study professional’s experience. This field is required.

Exhibit 5-8 and Exhibit 5-9 show the screen captures of the ‘Study Professionals’ screen:

Exhibit 5-8 Study Professionals Screen (Scroll 1)

The screenshot displays the GSPP (Global Study Plan Portal) interface. At the top, the user is logged in as John Doe, Primary Authorized Official. The navigation menu on the left includes sections for Enforceable Consent Agreement (ECA), Contact Information, Study Plans & Conduct of Testing, and Results. The main content area is titled 'STUDY PROFESSIONALS' and contains a form for adding a new professional. The form fields include:

- Select the Study Professional Role: (dropdown menu)
- CBI: (checkbox)
- Prefix: (dropdown menu)
- First Name: (text input)
- Middle Initial: (text input)
- Last Name: (text input)
- Suffix: (dropdown menu)
- Company Name: (text input)
- Phone Number: (text input) Ext: (text input)
- Email Address: (text input)
- Mailing Address 1: (text input)
- Mailing Address 2: (text input)
- City: (text input)
- State: (dropdown menu)
- Postal Code: (text input)
- Country: (dropdown menu)
- Experience Summary: (text area)

At the bottom of the screen, there are four buttons: Validate (with a green checkmark icon), Save (with a floppy disk icon), Preview (with a monitor icon), and Submit (with a database icon).

Exhibit 5-9: Study Professionals Screen (Scroll 2)

The screenshot displays the 'Study Professionals' screen. At the top, the user is logged in as 'John Doe, Primary Authorized Official'. The navigation menu on the left includes 'Section 4 Enforceable Consent Agreement (ECA) Primary Authorized Official', '68 FR 33125', 'Submission Information Type', 'Contact Information', 'Study Plans & Conduct of Testing', and 'Results'. The main form area contains the following fields:

- 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:

At the bottom of the form, there is a 'Save' button highlighted with a red box. Below the form, there is a message: 'Click the **Add Study Professional** button to add a new study professional.' and an 'Add Study Professional' button. At the bottom of the page, there are 'Previous' and 'Next' buttons, with the 'Next' button highlighted with a red box. The bottom navigation bar includes 'Validate', 'Save', 'Preview', and 'Submit' buttons.

Navigation: If you did not check the 'Results' checkbox on the 'Submission Information Type' screen, click the 'Next' button to navigate to the 'ECA Additional Information' screen. If you checked the 'Results' checkbox on the 'Submission Information Type' screen, click the 'Next' button to navigate to the 'Document Management' screen.

5.5 Results

If you have identified you are submitting results, you are required to complete the following screens to satisfy the reporting requirements.

5.5.1 Document Management Screen

The 'Document Management' screen allows you to upload documents related to Section 4 reporting requirements. To add Section 4 results documentation, click the 'Attach Document'

button. Clicking the ‘Attach Document’ button displays the ‘Document Management’ pop-up window.

The ‘Document Management’ pop-up window allows you to attach documentation by clicking the ‘Browse’ button. This will generate the ‘File Upload’ window where you can select the desired document from your hard drive. Once you have identified the document to be attached, click the ‘Open’ button. Clicking the ‘Open’ button will close the ‘File Upload’ window and display the attached document within the ‘Browse’ field. To claim attached documentation as CBI, check the ‘CBI’ checkbox. If you claim the attached documentation as CBI you are required to attach the sanitized version of the attached document. To do so, follow the steps outlined above. Once you have attached all desired documentation, click the ‘OK’ button to return to the ‘Document Management’ screen.

The document management table displays the file name of the document, the date the document was attached, and whether the document was claimed as CBI. The system allows you to delete unwanted documents by clicking the ‘Delete’ icon located under the ‘Action’ column for a given row. Clicking the file name links within the ‘File Name’ column generates the ‘Document Management’ pop-up window. This allows you to view and edit previously attached documents.


- **Add Document:** Clicking the ‘Add Document’ button displays the ‘Document Management’ pop-up window.
- **File Name:** Displays the name of the attached file. Clicking the file name link generates the ‘Document Management’ pop-up window and allows you to view the attached document.
- **Attachment Date:** Displays the date the document was attached.
- **CBI:** Displays whether the attached document was claimed CBI.
- **Action:** Clicking the ‘Delete’ icon () will generate a prompt confirming deletion. To delete the specified row and all contents within click the ‘OK’ button within the prompt.

Exhibit 5-10 shows the screen capture of the ‘Document Management’ screen:

Exhibit 5-10 Document Management Screen

Section 4 Enforceable Consent Agreement (ECA) > 70 FR 39630 > Results > Document Management

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

Validate Save Preview Submit

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Navigation: Click the ‘Next’ button to navigate to the ‘ECA Additional Information’ screen. You can also access this screen by clicking the ‘ECA Additional Information’ link within the navigation tree.

The ‘Document Management’ pop-up window allows you to browse for a document from your hard drive and attach one document at a time. You must attach at least one document. Clicking the ‘Browse’ button will generate the ‘File Upload’ window on your hard drive. Select the document you wish to attach and click the ‘Open’ button. If a document is marked as ‘CBI’ then a sanitized copy of the document must be attached. To save the attached document to the ‘Document Management’ screen, click the ‘OK’ button within the ‘Document Management’ pop-up window.

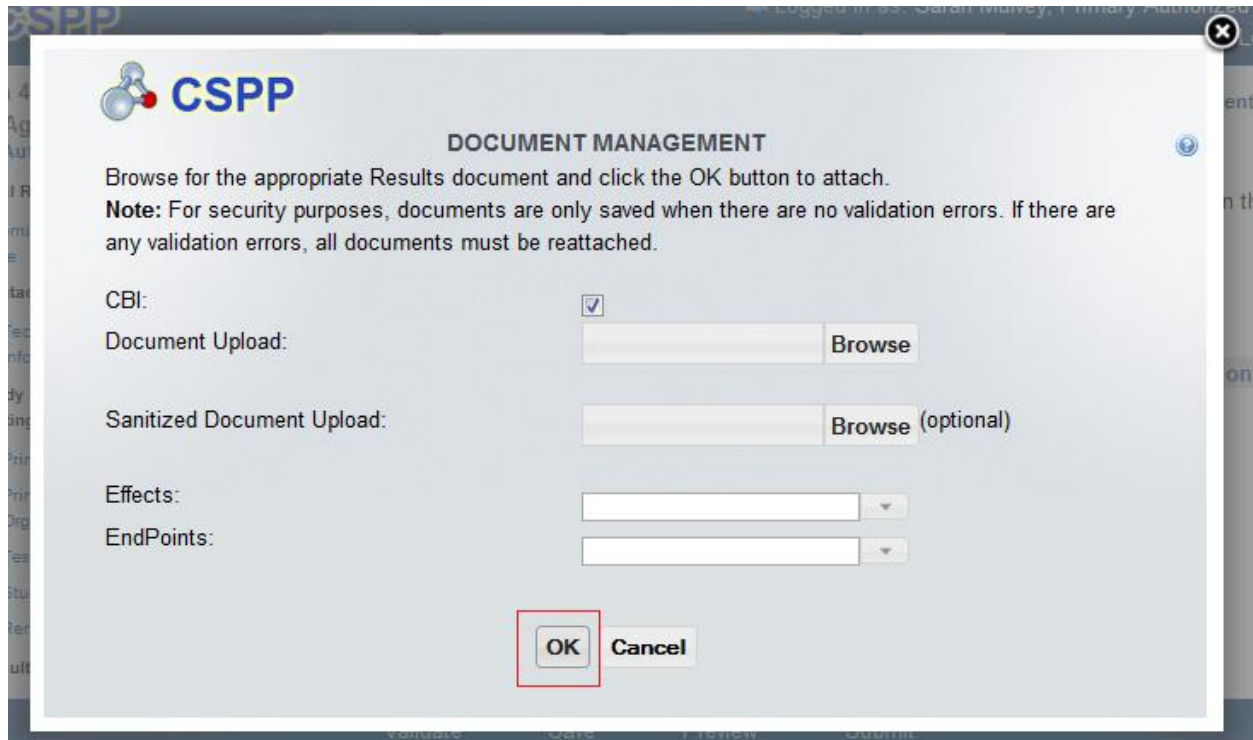
- **CBI:** Checking the ‘CBI’ checkbox will claim the uploaded document as CBI and generate a sanitized document upload field.
- **Document Upload:** Clicking the ‘Browse’ button will generate the ‘File Upload’ window for you to search for a particular file on your hard drive.
- **Sanitized Document Upload:** This field will only be displayed when the ‘CBI’ checkbox is checked. Clicking the ‘Browse’ button will generate the ‘File Upload’ window for you to search for a particular file on your hard drive.
- **Effects:** Select the effect specific to the chemical report. Displays the following options: ‘Health Effects,’ ‘Environmental Fate,’ ‘Ecotoxicity,’ and ‘Physical Chemical Properties.’

-
- **Endpoints:** Select an endpoint that is specific to the chemical report. The endpoints will be specific to the effect chosen from the 'Effects' drop-down menu. If no effect is chosen, no endpoints will display.

Refer to Appendix B. Submission Types, Effects, and Endpoints for a complete list of values that are found within each of the drop-down menus on the 'Document Management' pop-up window. If you have multiple documents that share the same effects and/or endpoints, you must upload each individual document separately and select values from the drop-down menus.

Exhibit 5-11 shows the screen capture of the ‘Document Management’ pop-up window:

Exhibit 5-11 Document Management Pop-Up Window



Navigation: Click the ‘OK’ or ‘Cancel’ buttons to return to the ‘Document Management’ screen.

5.6 ECA Additional Information Screen

You can access the ‘ECA Additional Information’ screen by clicking the ‘Next’ button from the ‘Document Management’ screen within the ‘Results’ folder of the navigation tree. If you did not check the ‘Results’ checkbox on the ‘Submission Information Type’ screen, click the ‘Next’ button from the ‘Study Professionals’ screen. You can also access the ‘ECA Additional Information’ screen by clicking the ‘ECA Additional Information’ link within the navigation tree.

The ‘ECA Additional Information’ screen allows you to upload documents related to Section 4 reporting requirements. To add additional information select the appropriate additional information type and click the ‘Attach Document’ button. Clicking the ‘Attach Document’ button displays the ‘ECA Additional Information’ pop-up window. If you select the ‘Amendments to the Study Plan’ radio button and click the ‘Attach Document’ button, the ‘Amendments to the Study Plan’ pop-up window generates. If you select the ‘Modification of ECAs’ radio button and click the ‘Attach Document’ button, the ‘Modification of ECAs’ pop-up window generates.

The ‘Amendment to the Study Plan’ pop-up window allows you to attach documentation by clicking the ‘Browse’ button. This will generate the ‘File Upload’ window where you can select the desired document from your hard drive. Once you have identified the document to be attached, click the ‘Open’ button. Clicking the ‘Open’ button will close the ‘File Upload’ window and display the attached document within the ‘Browse’ field. Once you have attached all

desired documentation, click the 'OK' button to return to the 'ECA Additional Information' screen.

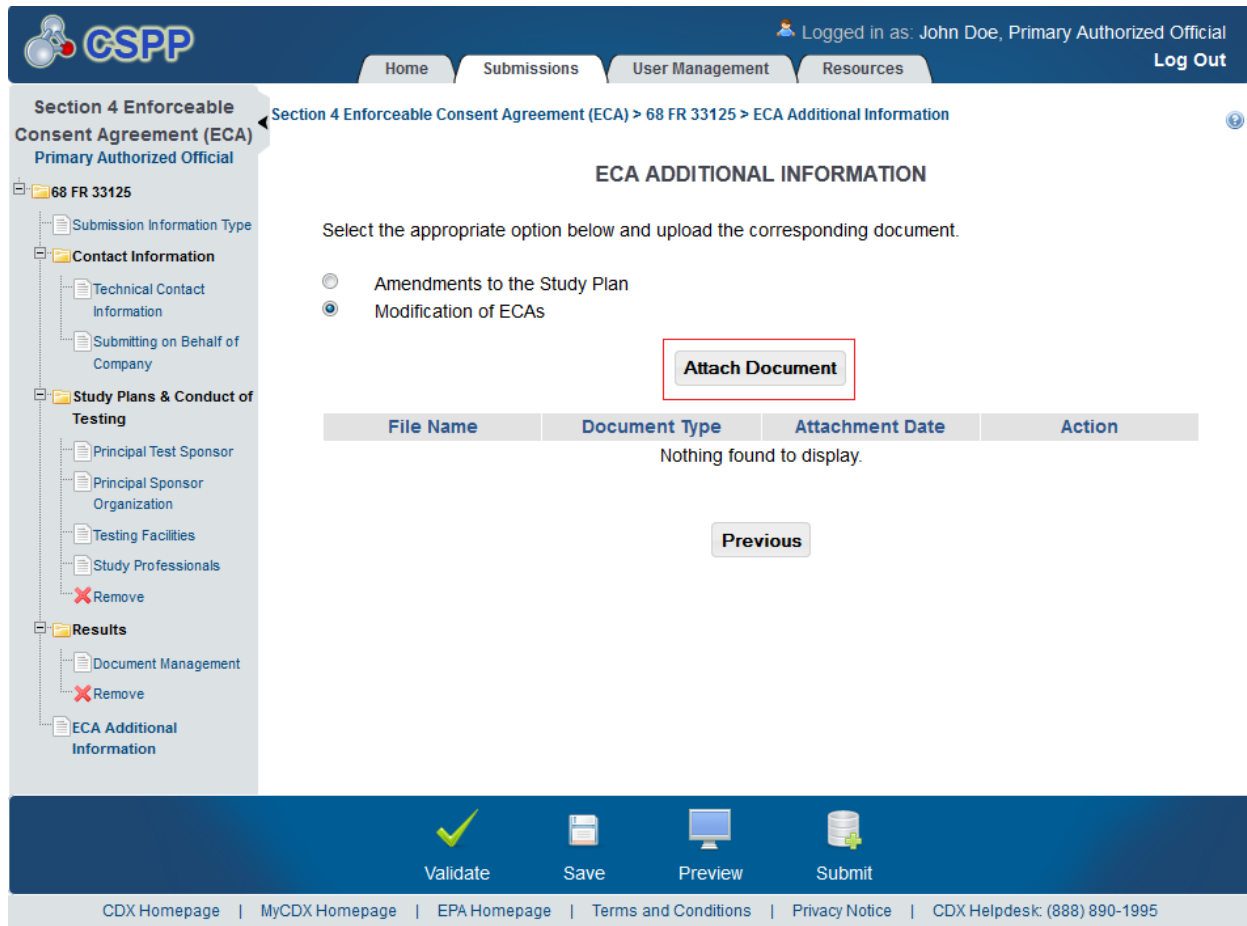
The 'Modification of ECAs' pop-up window allows you to provide an explanation and rationale for the modification of the ECA within a text box. Once you have provided an explanation click the 'OK' button to return to the 'ECA Additional Information' screen.

The 'ECA Additional Information' table displays the file name of the document, the document type, and the date the document was attached. The system allows you to delete unwanted documents by clicking the 'Delete' icon located under the 'Action' column for a given row. Clicking the file name links within the 'File Name' column generates the pop-up window. This allows you to view and edit previously attached documents or information.

- **Amendments to the Study Plan:** Select the radio button to indicate the additional information type.
- **Modification of ECAs:** Select the radio button to indicate the additional information type.
- **Add Document:** Clicking the 'Add Document' button displays the 'Amendments to the Study Plan' or 'Modification of ECAs' pop-up window.
- **File Name:** Displays the name of the attached file if applicable. Clicking the file name link generates the pop-up window and allows you to view the attached document or information.
- **Document Type:** Displays the additional information type.
- **Attachment Date:** Displays the date the document was attached.
- **Action:** Clicking the 'Delete' icon (✖) will generate a prompt confirming deletion. To delete the specified row and all contents within click the 'OK' button within the prompt.

Exhibit 5-12 shows the screen capture of the ‘ECA Additional Information’ screen:

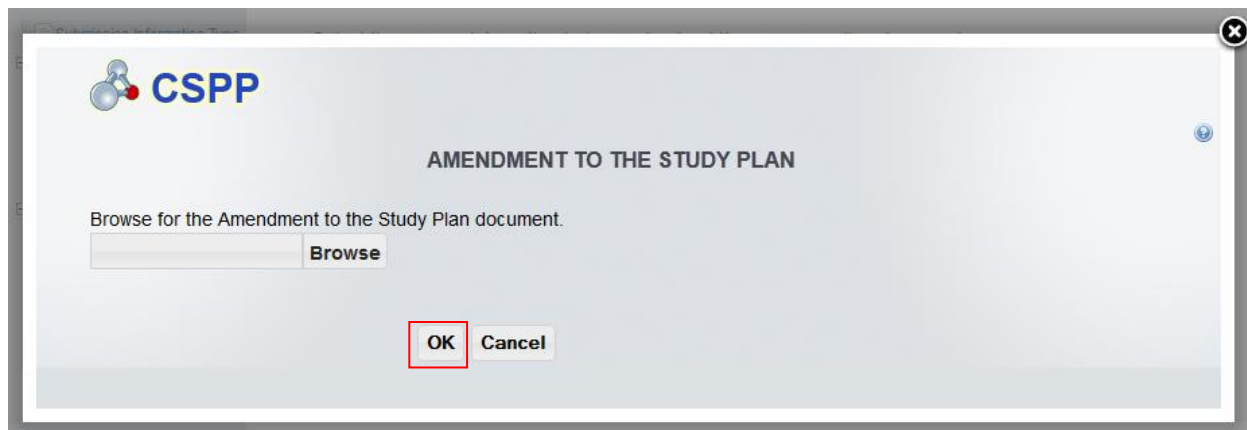
Exhibit 5-12 ECA Additional Information Screen



Navigation: Once all information has been uploaded, click the ‘Submit’ icon located within the action bar to begin the submission process.

Exhibit 5-13 shows the screen capture of the ‘Amendment to the Study Plan’ pop-up window:

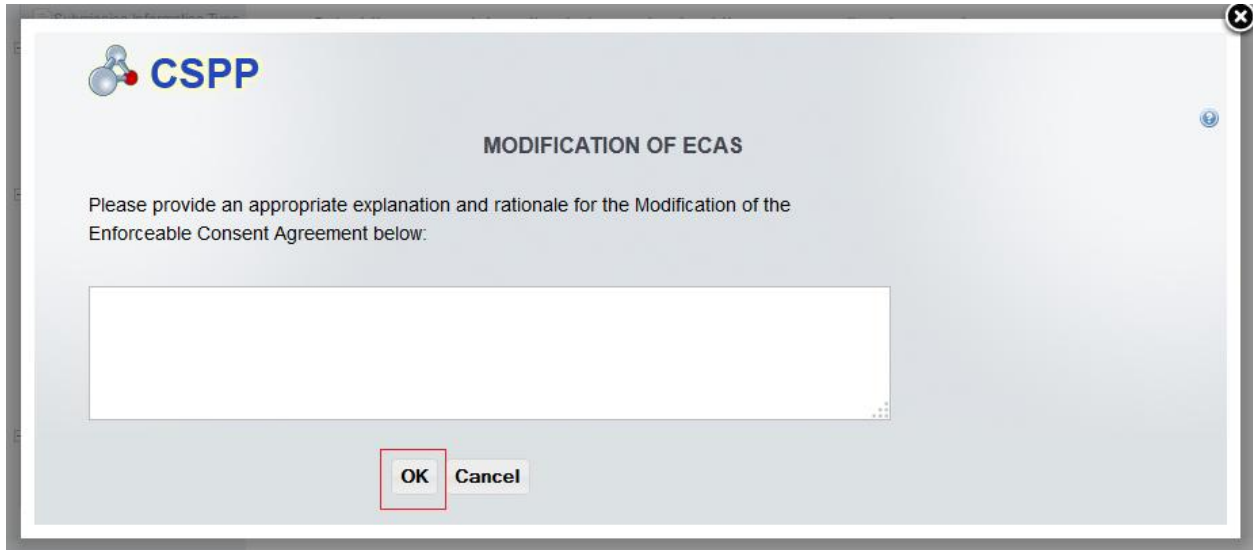
Exhibit 5-13 Amendment to the Study Plan Pop-Up Window



Navigation: Click the ‘OK’ or ‘Cancel’ buttons to return to the ‘ECA Additional Information’ screen.

Exhibit 5-14 shows the screen capture of the ‘Modification of ECAs’ pop-up window:

Exhibit 5-14 Modification of ECAs Pop-Up Window



Navigation: Click the ‘OK’ or ‘Cancel’ buttons to return to the ‘ECA Additional Information’ screen.

6 Memorandum of Understanding (MOU)

6.1 Submission Information Type Screen

You can access the ‘Submission Information Type’ screen by clicking the ‘Next’ button from either the ‘Create New Passphrase’ or the ‘Enter Passphrase’ screen. Once you are within the ‘Memorandum of Understanding’ flow of the Section 4 application, the ‘Submission Information Type’ screen can also be accessed by clicking the ‘Submission Information Type’ link located within the navigation tree.

The ‘Submission Information Type’ screen allows you to select the submission information type that you will be submitting to EPA. This screen consists of the following fields:

- **Please select, or begin typing, a Chemical in the drop-down menu below:** Displays all chemicals associated with MOU. This field is required.
- **Chemical Docket:** This field displays once a chemical is selected. It displays the docket number associated with the selected chemical.
- **Submission Alias:** This field is auto-populated with a date and time stamp and is displayed on the ‘Section 4 Test Rules, ECAs, MOUs, and 40 CFR 766’ submissions screen within the ‘Submission Alias’ column. The ‘Submission Alias’ is also used to differentiate between submission types within the ‘User Management’ screen. This field is editable should you choose to provide your own submission alias.
- **Study Plans & Conduct of Testing:** Checking the ‘Study Plans & Conduct of Testing’ checkbox identifies that you are fulfilling the requirements to submit a study plan to EPA. Once you have checked the ‘Study Plans & Conduct of Testing’ checkbox and either saved the information on this screen by clicking the ‘Save’ icon or by clicking the ‘Next’ button, the ‘Study Plans & Conduct of Testing’ folder is generated within the navigation tree allowing you to access the ‘Study Plans & Conduct of Testing’ screens required to be completed.
- **Results:** Checking the ‘Results’ checkbox indicates that you are fulfilling the requirements to submit results to EPA. Once you have checked the ‘Results’ checkbox and either saved the information on this screen by clicking the ‘Save’ icon or by clicking the ‘Next’ button, the ‘Results’ folder is generated within the navigation tree allowing you to access the ‘Results’ screens required to be completed.

If you have previously identified a submission type, have accessed the submission type screens, and then unchecked the previously checked submission information type (i.e. Study Plans & Conduct of Testing), you will receive a prompt indicating that all associated information will be lost. Clicking the ‘OK’ button within the prompt will delete all data associated with that submission information type.

Once you have identified the type(s) of information you will be submitting to EPA, you are able to access the screens required to be completed.

Exhibit 6-1 shows the screen capture of the ‘Submission Information Type’ screen:

Exhibit 6-1: Submission Information Type Screen

The screenshot displays the 'Submission Information Type' screen within the GSPP system. The top navigation bar includes 'Home', 'Submissions', 'User Management', 'Resources', and 'Log Out'. The user is logged in as 'John Doe, Primary Authorized Official'. The left navigation pane shows the path: Section 4 Memorandum of Understanding (MOU) > Federal Register Notice > Submission Information Type. The main content area is titled 'SUBMISSION INFORMATION TYPE' and contains the following elements:

- A prompt: "Please select, or begin typing, a Chemical in the drop-down menu below:" followed by a text input field with a dropdown arrow.
- A prompt: "Please enter a submission alias and choose the submission information type(s) here."
- A text input field for the submission alias containing "MOU-20130711-14:11:3".
- Two radio buttons: "Study Plans & Conduct of Testing" and "Results".
- A "Next" button.

At the bottom of the screen, there is a blue bar with four icons and labels: "Validate" (checkmark), "Save" (floppy disk), "Preview" (monitor), and "Submit" (submit button). Below this bar is a footer with links: "CDX Homepage", "MyCDX Homepage", "EPA Homepage", "Terms and Conditions", "Privacy Notice", and "CDX Helpdesk: (888) 890-1995".

Navigation: Click the ‘Next’ button to navigate to the ‘Technical Contact Information’ screen. You can also access this screen by clicking the ‘Technical Contact Information’ link within the navigation tree.

6.2 Technical Contact Information Screen

The ‘Technical Contact Information’ screen allows you to submit on behalf of another company or if you are not submitting on behalf of another company. To identify if you are submitting on behalf of another company, the system provides the radio button ‘This is a submission on behalf of another company.’ The system will not allow you to select more than one radio button per submission.

This functionality has been implemented to accommodate a company that may use an agent or another individual. If this radio button is selected, the information of the contractor/consultant’s company is to be entered into the ‘Submitting on Behalf Of Company’ screen. If you are not submitting on behalf of another company and have selected the ‘N/A’ radio button, you are taken to the ‘Principal Test Sponsor’ screen.

- **N/A:** Selecting the ‘N/A’ radio button allows you to signify that you are not making a submission on behalf of another company.
- **This is a submission on behalf of another company:** Selecting the ‘This is a submission on behalf of another company’ radio button allows you to identify yourself as an individual submitting on behalf of another company.

- **Copy CDX Registration:** Clicking the ‘Copy CDX Registration’ button will automatically populate the fields with your contact information provided during CDX registration.
- **Prefix:** The ‘Prefix’ drop-down menu displays all optional prefixes. If applicable, select the appropriate prefix. If you have clicked the ‘Copy CDX Registration’ button, the fields populate with information entered during CDX registration.
- **First Name:** Enter the technical contact’s first name. If you have clicked the ‘Copy CDX Registration’ button, the fields populate with information entered during CDX registration. This field is required.
- **Middle Initial:** Enter the technical contact’s middle initial. If you have clicked the ‘Copy CDX Registration’ button, the fields populate with information entered during CDX registration.
- **Last Name:** Enter the technical contact’s last name. If you have clicked the ‘Copy CDX Registration’ button, the fields populate with information entered during CDX registration. This field is required.
- **Suffix:** The ‘Suffix’ drop-down menu displays all optional suffixes. If applicable, select the appropriate suffix. If you have clicked the ‘Copy CDX Registration’ button, the fields populate with information entered during CDX registration.
- **Company Name:** Enter the technical contact’s company name. If you have clicked the ‘Copy CDX Registration’ button, the fields populate with information entered during CDX registration. This field is required.
- **Phone Number:** Enter the technical contact’s daytime phone number; U.S. or international. If you have clicked the ‘Copy CDX Registration’ button, the fields populate with information entered during CDX registration. This field is required.
- **Ext:** Enter the extension to the technical contact’s daytime phone number.
- **Email Address:** Enter the technical contact’s email address. The email address must follow the standard email address format: john.doe@abc.com. If you have clicked the ‘Copy CDX Registration’ button, the fields populate with information entered during CDX registration. This field is required.
- **Mailing Address 1:** Enter the first line of the technical contact’s mailing address. If you have clicked the ‘Copy CDX Registration’ button, the fields populate with information entered during CDX registration. This field is required.
- **Mailing Address 2:** Enter the second line of the technical contact’s mailing address. If you have clicked the ‘Copy CDX Registration’ button, the fields populate with information entered during CDX registration.
- **City:** Enter the city of the technical contact’s mailing address. If you have clicked the ‘Copy CDX Registration’ button, the fields populate with information entered during CDX registration. This field is required.
- **State:** Enter the state of the technical contact’s mailing address. If you have clicked the ‘Copy CDX Registration’ button, the fields populate with information entered during CDX registration. This is a required field if the ‘Country’ field is defined as ‘United States.’

-
- **Postal Code:** Enter the postal code of the technical contact's mailing address. If you have clicked the 'Copy CDX Registration' button, the fields populate with information entered during CDX registration. This field is required.
 - **Country:** Enter the country of the technical contact's mailing address. If you have clicked the 'Copy CDX Registration' button, the fields populate with information entered during CDX registration. This is a required field.

Exhibit 6-2 shows the screen capture of the ‘Technical Contact Information’ screen:

Exhibit 6-2 Technical Contact Information Screen

Section 4 Memorandum of Understanding (MOU) Primary Authorized Official

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](#)

Validate
 Save
 Preview
 Submit

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Navigation: If you are submitting on behalf of another company and click the ‘Next’ button, you are taken to the ‘Submitting on Behalf of Company’ screen. If you are submitting study plans & conduct of testing information and are not submitting on behalf of another entity, you are taken to the ‘Principal Test Sponsor’ screen. You can also access this screen by clicking the ‘Submitting on Behalf of Company’ link within the navigation tree.

6.3 Submitting on Behalf of Company Screen

You can access the ‘Submitting on Behalf of Company’ screen by selecting the ‘This is a submission on behalf of another company’ radio button and by clicking the ‘Next’ button from the ‘Technical Contact Information’ screen. You can also access this screen by clicking the ‘Submitting on Behalf of Company’ link within the navigation tree.

The ‘Submitting on Behalf Of Company’ screen presents a list of fields that allow you to input the contact information for the manufacturing or processing establishment on whose behalf the submission is being made.

- **Prefix:** The ‘Prefix’ drop-down menu displays all optional prefixes. If applicable, select the appropriate prefix.
- **First Name:** Enter the first name of the manufacturing or processing establishment contact for which the submission is being made. This field is required.
- **Middle Initial:** Enter the middle initial of the manufacturing or processing establishment contact for which the submission is being made.
- **Last Name:** Enter the last name of the manufacturing or processing establishment contact for which the submission is being made. This field is required.
- **Suffix:** The ‘Suffix’ drop-down menu displays all optional suffixes and captures the suffix of the manufacturing or processing establishment contact for which the submission is being made. If applicable, select the appropriate suffix.
- **Company Name:** Enter the company name of the manufacturing or processing establishment. This field is required.
- **Phone Number:** Enter the daytime phone number of the manufacturing or processing establishment contact for whom the submission is being made; U.S. or international. This field is required.
- **Email Address:** Enter the email address of the manufacturing or processing establishment contact for which the submission is being made. The email address must follow the standard email address format: john.doe@abc.com. This field is required.
- **Mailing Address 1:** Enter the first line of the mailing address of the manufacturing or processing establishment contact. This field is required.
- **Mailing Address 2:** Enter the second line of the mailing address of the manufacturing or processing establishment contact.
- **City:** Enter the city of the mailing address of the manufacturing or processing establishment contact. This field is required.
- **State:** Enter the state of the mailing address of the manufacturing or processing establishment contact. This is a required field if the ‘Country’ field is defined as ‘United States.’
- **Postal Code:** Enter the postal code of the mailing address of the manufacturing or processing establishment contact. This field is required.
- **Country:** Enter the country of the manufacturing or processing establishment contact’s mailing address. This field is required.

Exhibit 6-3 shows the screen capture of the ‘Submitting on Behalf of Company’ screen:

Exhibit 6-3 Submitting on Behalf of Company Screen

Section 4 Memorandum of Understanding (MOU) > Federal Register Notice > Contact Information > Submitting on Behalf of Company

SUBMITTING ON BEHALF OF COMPANY

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

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:

Validate Save Preview Submit

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Navigation: If you are submitting study plans & conduct of testing information and click the ‘Next’ button, you are taken to the ‘Principal Test Sponsor’ screen.

6.4 Study Plans & Conduct of Testing

If you have identified you are submitting information for study plans and conduct of testing, you are required to complete the following screens to satisfy the reporting requirements.

6.4.1 Principal Test Sponsor Screen

The ‘Principal Test Sponsor’ screen presents a list of fields that allow you to input the contact information for the principal test sponsor.

-
- **Prefix:** The ‘Prefix’ drop-down menu displays all optional prefixes. If applicable, select the appropriate prefix.
 - **First Name:** Enter the principal test sponsor contact’s first name. This field is required.
 - **Middle Initial:** Enter the principal test sponsor contact’s middle initial.
 - **Last Name:** Enter the principal test sponsor contact’s last name. This field is required.
 - **Suffix:** The ‘Suffix’ drop-down menu displays all optional suffixes and captures the principal test sponsor contact’s suffix. If applicable, select the appropriate suffix.
 - **Company Name:** Enter the principal test sponsor contact’s company name. This field is required.
 - **Phone Number:** Enter the principal test sponsor contact’s daytime phone number; U.S. or international. This field is required.
 - **Email Address:** Enter the principal test sponsor contact’s email address. The email address must follow the standard email address format: john.doe@abc.com. This field is required.
 - **Mailing Address 1:** Enter the first line of the mailing address of the principal test sponsor’s company. This field is required.
 - **Mailing Address 2:** Enter the second line of the mailing address of the principal test sponsor’s company.
 - **City:** Enter the city of the mailing address of the principal test sponsor’s company. This field is required.
 - **State:** Enter the state of the mailing address of the principal test sponsor contact’s company. This is a required field if the ‘Country’ field is defined as ‘United States.’
 - **Postal Code:** Enter the postal code of the principal test sponsor contact’s company. This field is required.
 - **Country:** Enter the country of the mailing address of principal test sponsor contact’s company. This field is required.

Exhibit 6-4 shows the screen capture of the ‘Principal Test Sponsor’ screen:

Exhibit 6-4 Principal Test Sponsor Screen

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:

Validate Save Preview Submit

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Navigation: Click the ‘Next’ button to navigate to the ‘Principal Sponsor Organization’ screen. You can also access this screen by clicking the ‘Principal Sponsor Organization’ link within the navigation tree.

6.4.2 Principal Sponsor Organization Screen

The ‘Principal Sponsor Organization’ screen allows you to enter the contact information for each administrative official and project manager within the principal test sponsor’s organization. To add each professional click the ‘Add Sponsor Organization’ button. This will generate a new set of text fields. Once you have entered data into one field set, you must click the ‘Save’ button located within that field set prior to adding another member. If the data previously entered and saved within a field set needs to be edited, you can click the ‘Edit’ button located in the upper right corner of that field set. If a field set needs to be deleted, you can click the ‘Delete’ icon () that is also located in the upper right corner of each field set.

The system allows you to expand/collapse each field set by clicking the field set header. If you wish to expand/collapse all field sets simultaneously, click the ‘Expand All’ or ‘Collapse All’ links located at the top right of all field sets.

- **Prefix:** The ‘Prefix’ drop-down menu displays all optional prefixes and captures the member contact’s prefix. If applicable, select the appropriate prefix.
- **First Name:** Enter the member contact’s first name. This field is required.
- **Middle Initial:** Enter the member contact’s middle initial.
- **Last Name:** Enter the member contact’s last name. This field is required.
- **Suffix:** The ‘Suffix’ drop-down menu displays all optional suffixes and captures the member contact’s suffix. If applicable, select the appropriate suffix.
- **Company Name:** Enter the member contact’s company name. This field is required.
- **Job Title:** Enter the member contact’s job title. This field is required.
- **Phone Number:** Enter the member contact’s daytime telephone number; U.S. or international. This field is required.
- **Email Address:** Enter the member contact’s email address. The email address must follow the standard email address format: john.doe@abc.com. This field is required.
- **Mailing Address 1:** Enter the first line of the member contact’s mailing address. This field is required.
- **Mailing Address 2:** Enter the second line of the member contact’s mailing address.
- **City:** Enter the city of the member contact’s mailing address. This field is required.
- **State:** Enter the state of the member contact’s mailing address. This field is required when ‘United States’ is identified as the country.
- **Postal Code:** Enter the postal code of the member contact’s mailing address. This field is required.
- **Country:** Enter the country of the member contact’s mailing address. This field is required.


Exhibit 6-5 shows the screen capture of the ‘Principal Sponsor Organization’ screen:

Exhibit 6-5 Principal Sponsor Organization Screen

Navigation: Click the ‘Next’ button to navigate to the ‘Testing Facilities’ screen. You can also access this screen by clicking the ‘Testing Facilities’ link within the navigation tree.

6.4.3 Testing Facilities Screen

The ‘Testing Facilities’ screen allows you to enter the contact information for each facility responsible for testing as well as that facility’s contact. To add each facility and contact, click the ‘Add Testing Facility’ button. This will generate a new set of text fields. Once you have entered

data into one field set, you must click the ‘Save’ button located within that field set prior to adding another facility and contact. If the data previously entered and saved within a field set needs to be edited, you can click the ‘Edit’ button located in the upper right corner of that field set. If a facility needs to be deleted, you can click the ‘Delete’ icon () that is also located in the upper right corner of each field set.

The system allows you to expand/collapse each field set by clicking the field set header. If you wish to expand/collapse all field sets simultaneously, click the ‘Expand All’ or ‘Collapse All’ links located at the top right of all field sets.

- **Testing Facility:** Enter the testing facility’s name. This field is required.
- **Phone Number:** Enter the testing facility’s phone number. This field is required.
- **Ext:** Enter the extension to the testing facility’s phone number.
- **Mailing Address 1:** Enter the first line of the testing facility’s mailing address. This field is required.
- **Mailing Address 2:** Enter the second line of the testing facility’s mailing address.
- **City:** Enter the city of the testing facility’s mailing address. This field is required.
- **State:** Enter the state of the testing facility’s mailing address. This field is required when ‘United States’ is identified as the country.
- **Postal Code:** Enter the postal code of the testing facility’s mailing address. This field is required.
- **Country:** Enter the country of the testing facility’s mailing address. This field is required.
- **Select the testing facility contact role:** Select the job title of the testing facility’s contact from the drop-down menu. This drop-down menu contains the following values: ‘Administrative Official,’ and ‘Project Manager.’ This field is required.
- **Prefix:** The ‘Prefix’ drop-down menu displays all optional prefixes and captures the testing facility contact’s prefix. If applicable, select the appropriate prefix.
- **First Name:** Enter the testing facility contact’s first name. This field is required.
- **Middle Initial:** Enter the testing facility contact’s middle initial.
- **Last Name:** Enter the testing facility contact’s last name. This field is required.
- **Suffix:** The ‘Suffix’ drop-down menu displays all optional suffixes and captures the testing facility contact’s suffix. If applicable, select the appropriate suffix.
- **Phone Number:** Enter the testing facility’s contact’s daytime phone number; U.S. or international. This field is required.
- **Email Address:** Enter the testing facility contact’s email address. The email address must follow the standard email address format: john.doe@abc.com. This field is required.
- **Mailing Address 1:** Enter the first line of the testing facility contact’s mailing address. This field is required.
- **Mailing Address 2:** Enter the second line of the testing facility contact’s mailing address.
- **City:** Enter the city of the testing facility contact’s mailing address. This field is required.

- **State:** Enter the state of the testing facility contact’s mailing address. This field is required when ‘United States’ is identified as the country.
- **Postal Code:** Enter the postal code of the testing facility contact’s mailing address. This field is required.
- **Country:** Enter the country of the mailing address of the testing facility’s contact. This field is required.

Exhibit 6-6 and Exhibit 6-7 show the screen captures of the ‘Testing Facilities’ screen:

Exhibit 6-6 Testing Facilities Screen (Scroll 1)

Section 4 Memorandum of Understanding (MOU) Primary Authorized Official

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:

✓ Validate 📄 Save 🖨 Preview 📤 Submit

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Exhibit 6-7: Testing Facility Screen (Scroll 2)

The screenshot displays the 'Testing Facility Screen' in the CSPP system. The page is titled 'Section 4 Memorandum of Understanding (MOU) Primary Authorized Official'. The user is logged in as 'John Doe, Primary Authorized Official'. The navigation menu on the left includes 'Federal Register Notice', 'Contact Information', 'Study Plans & Conduct of Testing', and 'Results'. The main form area contains the following fields:

- State: [Dropdown]
- Postal Code: [Text]
- Country: [Dropdown]
- Select the testing facility contact role: [Dropdown]
- Prefix: [Text]
- First Name: [Text]
- Middle Initial: [Text]
- Last Name: [Text]
- Suffix: [Text]
- Phone Number: [Text] Ext: [Text]
- Email Address: [Text]
- Mailing Address 1: [Text]
- Mailing Address 2: [Text]
- City: [Text]
- State: [Dropdown]
- Postal Code: [Text]
- Country: [Dropdown]

Buttons: 'Save' (highlighted with a red box), 'Add Testing Facility', 'Previous', and 'Next' (highlighted with a red box).

Bottom navigation bar: 'Validate', 'Save', 'Preview', 'Submit'.

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

Navigation: Click the 'Next' button to navigate to the 'Study Professionals' screen. You can also access this screen by clicking the 'Study Professionals' link within the navigation tree.

6.4.4 Study Professionals Screen

The 'Study Professionals' screen allows you to provide the contact information and a summary of the training and experience of each professional involved in the study. To add each professional click the 'Add Study Professional' button. This will generate a new set of text fields. Once you have entered data into one field set, you must click the 'Save' button located within that field set prior to adding another professional. If the data previously entered and saved within a field set needs to be edited, you can click the 'Edit' button located in the upper right

corner of that field set. If a professional needs to be deleted, you can click the ‘Delete’ icon (✖) that is also located in the upper right corner of each field set.

The system allows you to expand/collapse each field set by clicking the field set header. If you wish to expand/collapse all field sets simultaneously, click the ‘Expand All’ or ‘Collapse All’ links located at the top right of all field sets.

- **Select the Study Professional Role:** Select the study professional’s role from the drop-down menu. This drop-down menu contains the following values: ‘Chemist,’ ‘Laboratory Assistant,’ ‘Microbiologist,’ ‘Pathologist,’ ‘Study Director,’ ‘Toxicologist,’ ‘Veterinarian,’ and ‘Other.’ This field is required.
- **Prefix:** The ‘Prefix’ drop-down menu displays all optional prefixes and captures the study professional’s prefix. If applicable, select the appropriate prefix.
- **First Name:** Enter the study professional’s first name. This field is required.
- **Middle Initial:** Enter the study professional’s middle initial.
- **Last Name:** Enter the study professional’s last name. This field is required.
- **Suffix:** The ‘Suffix’ drop-down menu displays all optional suffixes and captures the study professional’s suffix. If applicable, select the appropriate suffix.
- **Company Name:** Enter the study professional’s company name. This field is required.
- **Phone Number:** Enter the study professional’s daytime phone number; U.S. or international. This field is required.
- **Email Address:** Enter the study professional’s email address. The email address must follow the standard email address format: john.doe@abc.com. This field is required.
- **Mailing Address 1:** Enter the first line of the study professional’s mailing address. This field is required.
- **Mailing Address 2:** Enter the second line of the study professional’s mailing address.
- **City:** Enter the city of the study professional’s mailing address. This is a required field.
- **State:** Enter the state of the study professional’s mailing address. This field is required when ‘United States’ is identified as the country.
- **Postal Code:** Enter the postal code of the study professional’s mailing address. This field is required.
- **Country:** Enter the country of the study professional’s mailing address. This is a required field.
- **Experience Summary:** Enter a summary of the study professional’s experience.

Exhibit 6-8 shows the screen capture of the ‘Study Professionals’ screen:

Exhibit 6-8 Study Professionals Screen

Section 4 Memorandum of Understanding (MOU) Primary Authorized Official

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

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

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Navigation: Click the ‘Next’ button to navigate to the ‘Document Management’ screen. You can also access this screen by clicking the ‘Document Management’ link within the navigation tree if the ‘Results’ checkbox was checked on the ‘Submission Information Type’ screen.

6.5 Results

If you have identified you are submitting results, you are required to complete the following screens to satisfy the reporting requirements.

6.5.1 Document Management Screen

The ‘Document Management’ screen allows you to upload documents related to Section 4 reporting requirements. To add Section 4 results documentation, click the ‘Attach Document’ button. Clicking the ‘Attach Document’ button displays the ‘Document Management’ pop-up window.

The ‘Document Management’ pop-up window allows you to attach documentation by clicking the ‘Browse’ button. This will generate the ‘File Upload’ window where you can select the desired document from your hard drive. Once you have identified the document to be attached, click the ‘Open’ button. Clicking the ‘Open’ button will close the ‘File Upload’ window and display the attached document within the ‘Browse’ field.

The ‘Document Management’ table displays the file name of the document and the date the document was attached. The system allows you to delete unwanted documents by clicking the ‘Delete’ icon located under the ‘Action’ column for a given row. Clicking the file name links within the ‘File Name’ column generates the ‘Document Management’ pop-up window. This allows you to view and edit previously attached documents.


- **Add Document:** Clicking the ‘Add Document’ button displays the ‘Document Management’ pop-up window.
- **File Name:** Displays the name of the attached file. Clicking the file name link generates the ‘Document Management’ pop-up window and allows you to view the attached document.
- **Attachment Date:** Displays the date the document was attached.
- **Action:** Clicking the ‘Delete’ icon () will generate a prompt confirming deletion. To delete the specified row and all contents within click the ‘OK’ button within the prompt.

Exhibit 6-9 shows the screen capture of the ‘Document Management’ screen:

Exhibit 6-9 Document Management Screen

The screenshot displays the 'Document Management' screen within the GSPP system. The top navigation bar includes 'Home', 'Submissions', 'User Management', 'Resources', and 'Log Out'. The user is logged in as 'John Doe, Primary Authorized Official'. The main content area is titled 'DOCUMENT MANAGEMENT' and contains the instruction: 'To add a Results Document, please click the Attach Document button and enter the information in the popup.' The 'Attach Document' button is highlighted with a red box. Below the button is a table with columns for 'File Name', 'Attachment Date', and 'Action'. The table is currently empty, displaying 'Nothing found to display.' Navigation buttons for 'Previous' and 'Next' are visible. The footer includes a status bar with icons for 'Validate', 'Save', 'Preview', and 'Submit', and a navigation bar with links to 'CDX Homepage', 'MyCDX Homepage', 'EPA Homepage', 'Terms and Conditions', 'Privacy Notice', and 'CDX Helpdesk: (888) 890-1995'.

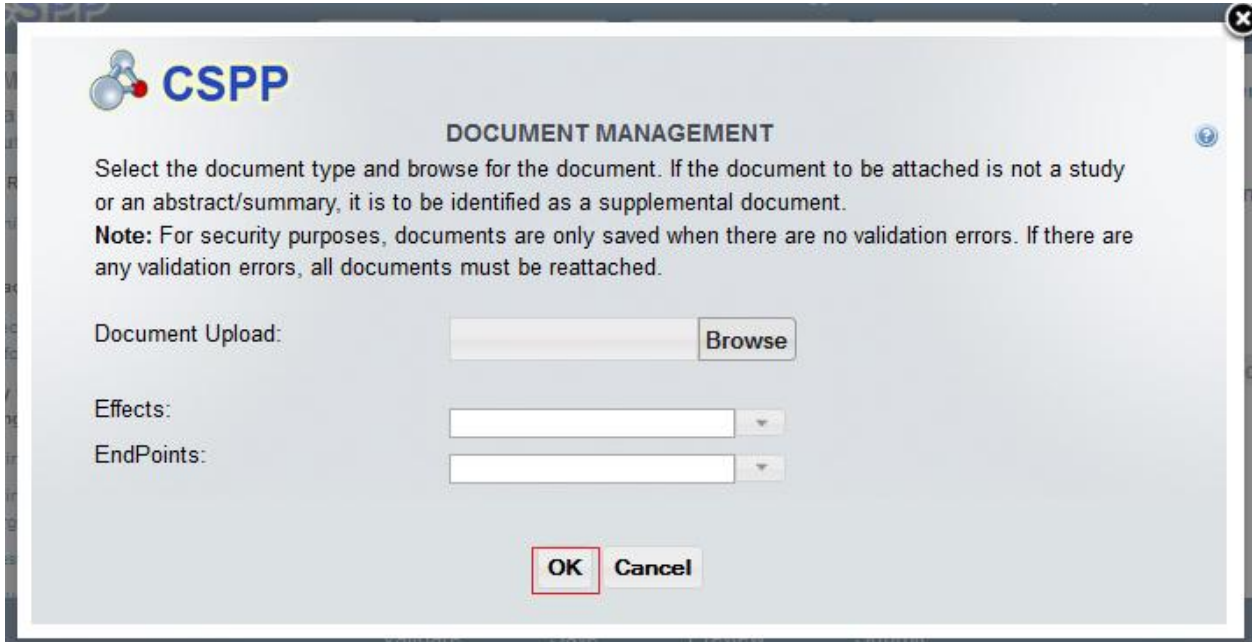
Navigation: Click the ‘Next’ button to navigate to the ‘MOU Additional Information’ screen. You can also access this screen by clicking the ‘MOU Additional Information’ link within the navigation tree.

The ‘Document Management’ pop-up window allows you to browse for a document from your hard drive and attach one document at a time. You must attach at least one document. Clicking the ‘Browse’ button will generate the ‘File Upload’ window on your hard drive. Select the document you wish to attach and click the ‘Open’ button. If a document is marked as ‘CBI’ then a sanitized copy of the document must be attached. To save the attached document to the ‘Document Management’ screen, click the ‘OK’ button within the ‘Document Management’ pop-up window.

- **Original Document:** Clicking the ‘Browse’ button will generate the ‘File Upload’ window for you to search for a particular file on your hard drive.
- **Effects:** Select the effect specific to the chemical report. Displays the following options: ‘Health Effects,’ ‘Environmental Fate,’ ‘Ecotoxicity,’ and ‘Physical Chemical Properties.’
- **Endpoints:** Select an endpoint that is specific to the chemical report. The endpoints will be specific to the effect chosen from the ‘Effects’ drop-down menu. If no effect is chosen, no endpoints will display.

Refer to Appendix B. Submission Types, Effects, and Endpoints for a complete list of values that are found within each of the drop-down menus on the ‘Document Management’ pop-up window. If you have multiple documents that share the same effects and/or endpoints, you must upload each individual document separately and select values from the drop-down menus.

Exhibit 6-10 Document Management Pop-Up Window



Navigation: Click the ‘OK’ or ‘Cancel’ buttons to return to the ‘Document Management’ screen.

6.6 MOU Additional Information Screen

The ‘MOU Additional Information’ screen allows you to upload documents related to Section 4 reporting requirements. To add additional information select the appropriate additional information type and click the ‘Attach Document’ button. Clicking the ‘Attach Document’ button displays the ‘MOU Additional Information’ pop-up window. If you select the ‘Amendments to the Study Plan’ radio button and click the ‘Attach Document’ button, the ‘Amendments to the Study Plan’ pop-up window generates. If you select the ‘Modification of MOUs’ radio button and click the ‘Attach Document’ button, the ‘Modification of MOUs’ pop-up window generates.

The ‘Amendments to the Study Plan’ pop-up window allows you to attach documentation by clicking the ‘Browse’ button. This will generate the ‘File Upload’ window where you can search for the desired document from your hard drive. Once you have identified the document to be attached, click the ‘Open’ button. Clicking the ‘Open’ button will close the ‘File Upload’ window and display the attached document within the ‘Browse’ field. Once you have attached all desired documentation, click the ‘OK’ button to return to the ‘MOU Additional Information’ screen.

The ‘Modification of MOUs’ pop-up window allows you to provide an explanation and rationale for the modification of the MOU within a text box. Once you have provided an explanation click the ‘OK’ button to return to the ‘MOU Additional Information’ screen.

The MOU additional information table displays the file name of the document, the document type, and the date the document was attached. The system allows you to delete unwanted documents by clicking the ‘Delete’ icon located under the ‘Action’ column for a given row. Clicking the file name links within the ‘File Name’ column generates the pop-up window. This allows you to view and edit previously attached documents or information.


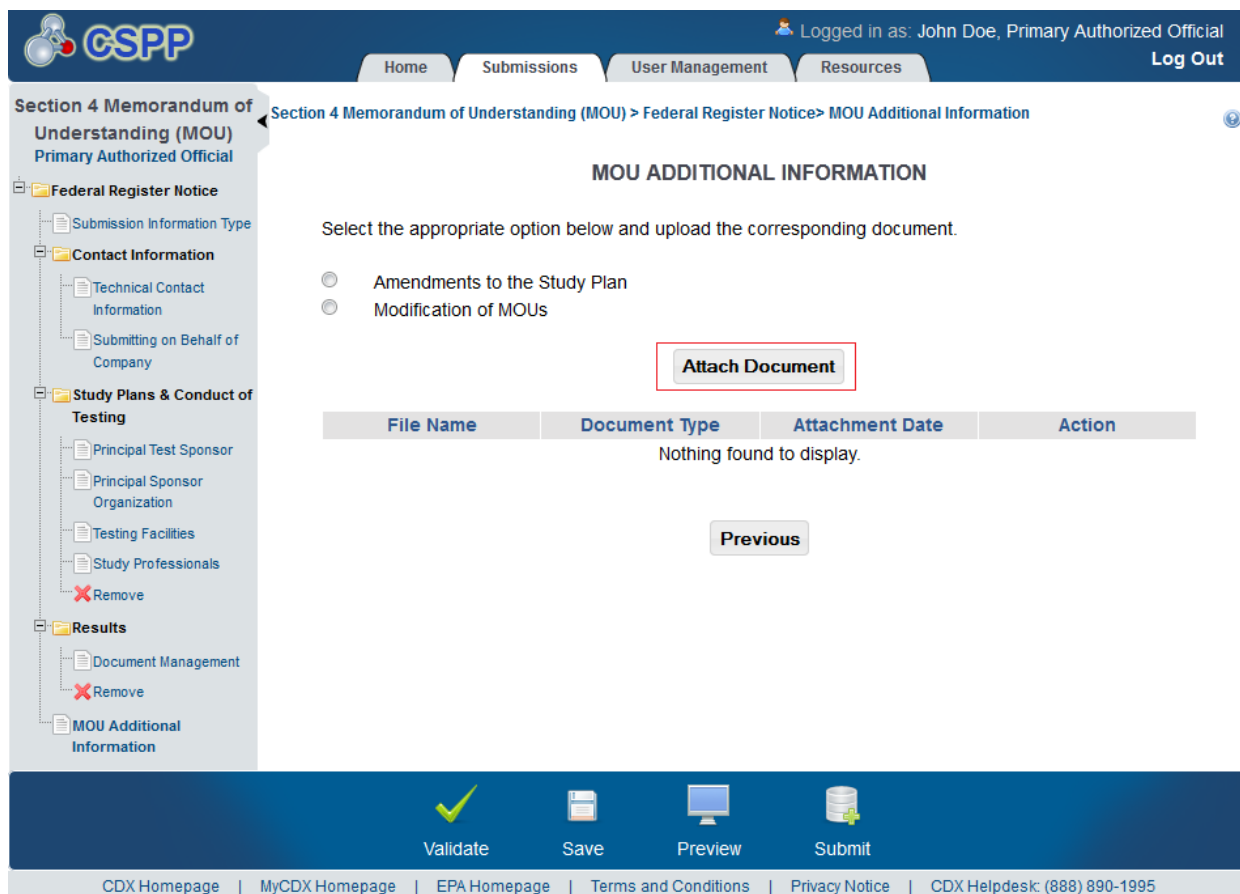
- **Amendments to the Study Plan:** Select the radio button to indicate the additional information type.
- **Modification of MOUs:** Select the radio button to indicate the additional information type.
- **Attach Document:** Clicking the ‘Attach Document’ button displays the ‘Amendments to the Study Plan’ or ‘Modification of MOUs’ pop-up window.
- **File Name:** Displays the name of the attached file if applicable. Clicking the file name link generates the pop-up window and allows you to view the attached document or information.
- **Document Type:** Displays the additional information type indicated on the page.
- **Attachment Date:** Displays the date the document was attached.
- **Action:** Clicking the ‘Delete’ icon () will generate a prompt confirming deletion. To delete the specified row and all contents within click the ‘OK’ button within the prompt.

Exhibit 6-11 shows the screen capture of the ‘MOU Additional Information’ screen:

Exhibit 6-11 MOU Additional Information Screen



The screenshot displays the 'MOU Additional Information' screen. At the top, the user is logged in as 'John Doe, Primary Authorized Official'. The navigation menu on the left includes 'Section 4 Memorandum of Understanding (MOU)', 'Federal Register Notice', 'Contact Information', 'Study Plans & Conduct of Testing', and 'Results'. The main content area is titled 'MOU ADDITIONAL INFORMATION' and contains the instruction: 'Select the appropriate option below and upload the corresponding document.' There are two radio buttons: 'Amendments to the Study Plan' and 'Modification of MOUs'. A red box highlights the 'Attach Document' button. Below this is a table with the following structure:

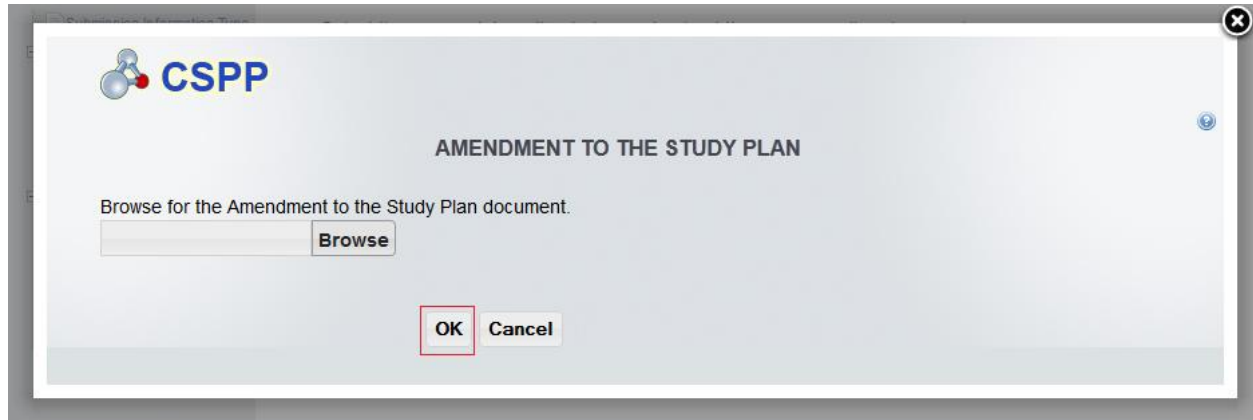
File Name	Document Type	Attachment Date	Action
Nothing found to display.			

At the bottom of the screen, there are four buttons: 'Validate', 'Save', 'Preview', and 'Submit'. The footer contains links to 'CDX Homepage', 'MyCDX Homepage', 'EPA Homepage', 'Terms and Conditions', 'Privacy Notice', and 'CDX Helpdesk: (888) 890-1995'.

Navigation: Click the ‘Previous’ button to navigate to the ‘Document Management’ screen. If you have completed all required fields, click the ‘Submit’ icon located in the action bar to begin the submission process.

Exhibit 6-12 shows the screen capture of the ‘Amendment to the Study Plan’ pop-up window:

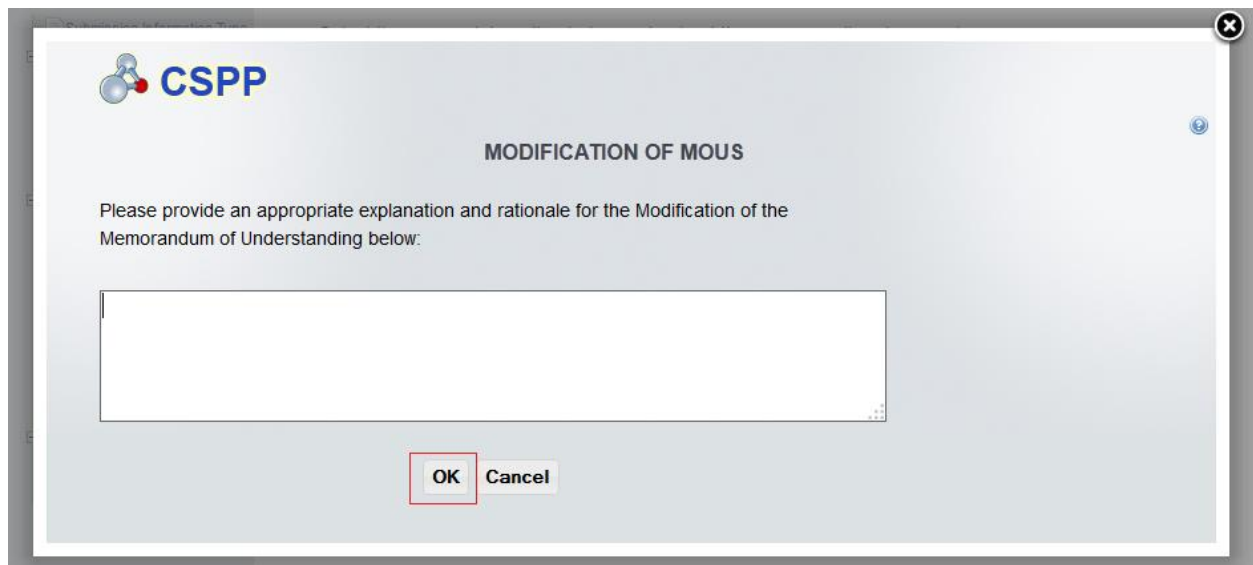
Exhibit 6-12 Amendment to the Study Plan Pop-Up Window



Navigation: Click the ‘OK’ or ‘Cancel’ button to return to the ‘MOU Additional Information’ screen.

Exhibit 6-13 shows the screen capture of the ‘Modification of MOUs’ pop-up window:

Exhibit 6-13 Modification of MOUs Pop-Up Window



Navigation: Click the ‘OK’ or ‘Cancel’ button to return to the ‘MOU Additional Information’ screen.

7 40 CFR 766 Dibenzodioxins/Dibenzofurans

7.1 Submission Information Type Screen

You can access the ‘Submission Information Type’ screen by clicking the ‘Next’ button from either the ‘Create New Passphrase’ or the ‘Enter Passphrase’ screen. Once you are within the ‘40 CFR 766’ flow of the Section 4 application, the ‘Submission Information Type’ screen can also be accessed by clicking the ‘Submission Information Type’ link located within the navigation tree.

The ‘Submission Information Type’ screen allows you to select the submission information type that you will be submitting to EPA. This screen consists of the following fields:

- **Submission Alias:** This field is auto-populated with a date and time stamp and is displayed on the ‘Section 4 Test Rules, ECAs, MOUs, and 40 CFR 766’ submissions screen within the ‘Submission Alias’ column. The ‘Submission Alias’ is also used to differentiate between submission types within the ‘User Management’ screen. This field is editable should you choose to provide your own submission alias. This field is required.
- **Letter of Intent:** Checking the ‘Letter of Intent’ checkbox indicates that you are fulfilling the requirements to submit a letter of intent to EPA. Once you have checked the ‘Letter of Intent’ checkbox and either saved the information on this screen by clicking the ‘Save’ icon or by clicking the ‘Next’ button, the ‘Letter of Intent’ folder is generated within the navigation tree allowing you to access the letter of intent screens required to be completed.
- **Protocol:** Checking the ‘Protocol’ checkbox indicates that you are fulfilling the requirements to submit protocols to EPA. Once you have checked the ‘Protocol’ checkbox and either saved the information on this screen by clicking the ‘Save’ icon or by clicking the ‘Next’ button, the ‘Protocol’ folder is generated within the navigation tree allowing you to access the protocol screens required to be completed.
- **Results:** Checking the ‘Results’ checkbox indicates that you are fulfilling the requirements to submit results to EPA. Once you have checked the ‘Results’ checkbox and either saved the information on this screen by clicking the ‘Save’ icon or by clicking the ‘Next’ button, the ‘Results’ folder is generated within the navigation tree allowing you to access the results screens required to be completed.
- **User Requests:** Checking the ‘User Requests’ checkbox indicates that you are submitting a request to EPA. Checking the ‘User Requests’ checkbox will enable the ‘Waiver,’ ‘Exclusion,’ and ‘Exemption’ radio button options. If you indicate that you are submitting a ‘User Request’ the ‘Letter of Intent,’ ‘Protocol,’ and ‘Results’ checkboxes will become disabled.
 - **Waiver:** Selecting the ‘Waiver’ radio button indicates that you are fulfilling the requirements to submit a waiver to EPA. Once you have checked the ‘User Requests’ checkbox and selected the ‘Waiver’ radio button and either saved the information on this screen by clicking the ‘Save’ icon or by clicking the ‘Next’ button, the ‘User Requests’ folder is generated within the navigation tree allowing you to access the ‘Waiver’ screen.
 - **Exclusion:** Selecting the ‘Exclusion’ radio button indicates that you are fulfilling the requirements to submit an exclusion to EPA. Once you have checked the ‘User Requests’ checkbox and selected the ‘Exclusion’ radio button and either

saved the information on this screen by clicking the ‘Save’ icon or by clicking the ‘Next’ button, the ‘User Requests’ folder is generated within the navigation tree allowing you to access the ‘Exclusion’ screen.

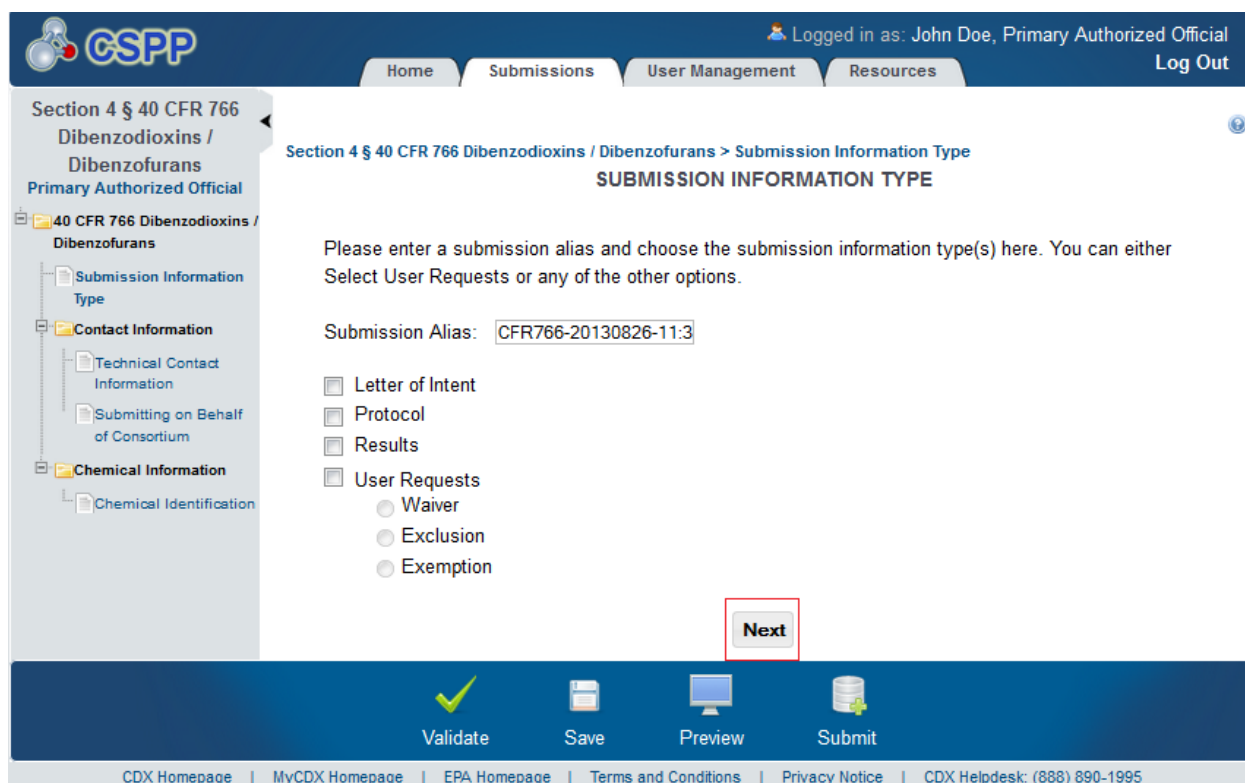
- **Exemption:** Selecting the ‘Exemption’ radio button indicates that you are fulfilling the requirements to submit an exemption to EPA. Once you have checked the ‘User Requests’ checkbox and selected the ‘Exemption’ radio button and either saved the information on this screen by clicking the ‘Save’ icon or by clicking the ‘Next’ button, the ‘User Requests’ folder is generated within the navigation tree allowing you to access the ‘Exemption’ screen.

If you have previously identified a submission type, have accessed the submission type screens, and then uncheck the previously checked submission information type (i.e. Study Plan), you will receive a prompt indicating that all associated information will be lost. Clicking the ‘OK’ button within the prompt will delete all data associated with that submission information type.

Once you have identified the type(s) of information you will be submitting to EPA, you are able to access the screens required to be completed.

Exhibit 7-1 shows the screen capture of the ‘Submission Information Type’ screen:

Exhibit 7-1 Submission Information Type Screen



Navigation: Click the ‘Next’ button to navigate to the ‘Technical Contact Information’ screen. You can also access this screen by clicking the ‘Technical Contact Information’ link within the navigation tree.

7.2 Technical Contact Information Screen

The ‘Technical Contact Information’ screen allows you to submit on behalf of another company or submit on behalf of a consortium. To identify if you are submitting on behalf of another entity, the system provides the radio button for ‘This is a submission on behalf of a consortium.’ To identify if you are submitting on behalf of another company, the system provides the radio button ‘This is a submission on behalf of another company.’ The system will not allow you to select more than one radio button per submission.

If you are not submitting on behalf of another entity and have selected the ‘N/A’ radio button, you are taken to the ‘Chemical Identification’ screen. If you are submitting on behalf of a consortium, select the ‘This is a submission on behalf of a consortium’ radio button. If the radio button for ‘This is a submission on behalf of a consortium’ is selected, you are taken to the ‘Submitting on Behalf of Consortium’ screen. If you are submitting on behalf of another company, select the ‘This is a submission on behalf of another company’ radio button. This functionality has been implemented to accommodate a company that may use an agent or another individual. If this radio button is selected, the information of the contractor/consultant’s company is to be entered into the ‘Submitting on Behalf Of Company’ screen.

- **N/A:** Selecting the ‘N/A’ radio button allows you to signify that you are neither making a submission on behalf of a consortium nor making a submission on behalf of another company.
- **This is a submission on behalf of a consortium:** Selecting the ‘This is a submission on behalf of a consortium’ radio button allows you to identify yourself as an individual submitting on behalf of a consortium.
- **This is a submission on behalf of another company:** Selecting the ‘This is a submission on behalf of another company’ radio button allows you to identify yourself as an individual submitting on behalf of another company.
- **Copy CDX Registration:** Clicking the ‘Copy CDX Registration’ button will automatically populate the fields with your contact information provided during CDX registration.
- **Prefix:** The ‘Prefix’ drop-down menu displays all optional prefixes. If applicable, select the appropriate prefix. If you have clicked the ‘Copy CDX Registration’ button, the fields populate with information entered during CDX registration.
- **First Name:** Enter the technical contact’s first name. If you have clicked the ‘Copy CDX Registration’ button, the fields populate with information entered during CDX registration. This field is required.
- **Middle Initial:** Enter the technical contact’s middle initial. If you have clicked the ‘Copy CDX Registration’ button, the fields populate with information entered during CDX registration.
- **Last Name:** Enter the technical contact’s last name. If you have clicked the ‘Copy CDX Registration’ button, the fields populate with information entered during CDX registration. This field is required.
- **Suffix:** The ‘Suffix’ drop-down menu displays all optional suffixes. If applicable, select the appropriate suffix. If you have clicked the ‘Copy CDX Registration’ button, the fields populate with information entered during CDX registration.

- **Company Name:** Enter the technical contact's company name. If you have clicked the 'Copy CDX Registration' button, the fields populate with information entered during CDX registration. This field is required.
- **Phone Number:** Enter the technical contact's daytime phone number; U.S. or international. If you have clicked the 'Copy CDX Registration' button, the fields populate with information entered during CDX registration. This field is required.
- **Email Address:** Enter the technical contact's email address. The email address must follow the standard email address format: john.doe@abc.com. If you have clicked the 'Copy CDX Registration' button, the fields populate with information entered during CDX registration. This field is required.
- **Mailing Address 1:** Enter the first line of the technical contact's mailing address. If you have clicked the 'Copy CDX Registration' button, the fields populate with information entered during CDX registration. This field is required.
- **Mailing Address 2:** Enter the second line of the technical contact's mailing address. If you have clicked the 'Copy CDX Registration' button, the fields populate with information entered during CDX registration.
- **City:** Enter the city of the technical contact's mailing address. If you have clicked the 'Copy CDX Registration' button, the fields populate with information entered during CDX registration. This field is required.
- **State:** Enter the state of the technical contact's mailing address. If you have clicked the 'Copy CDX Registration' button, the fields populate with information entered during CDX registration. This is a required field if the 'Country' field is defined as 'United States.'
- **Postal Code:** Enter the postal code of the technical contact's mailing address. If you have clicked the 'Copy CDX Registration' button, the fields populate with information entered during CDX registration. This field is required.
- **Country:** Enter the country of the technical contact's mailing address. If you have clicked the 'Copy CDX Registration' button, the fields populate with information entered during CDX registration. This is a required field.

Exhibit 7-2 shows the screen capture of the ‘Technical Contact Information’ screen:

Exhibit 7-2 Technical Contact Information Screen

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

Section 4 § 40 CFR 766 Dibenzenodioxins / Dibenzofurans > 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](#)

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

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

Navigation: If you are submitting on behalf of a consortium, you are taken to the ‘Submitting on Behalf of Consortium’ screen. If you are submitting on behalf of another company and click the ‘Next’ button, you are taken to the ‘Submitting on Behalf of Company’ screen. If you are submitting a Letter of Intent and are not submitting on behalf of another entity, you are taken to the ‘Chemical Identification’ screen. If you are submitting a protocol or results and are not submitting on behalf of another entity, you are taken to the ‘Document Management’ screen.

7.3 Submitting on Behalf of Consortium Screen

You can access the ‘Submitting on Behalf of Consortium’ screen by selecting the ‘This is a submission on behalf of a consortium’ radio button and by clicking the ‘Next’ button from the ‘Technical Contact Information’ screen. You can also access this screen by clicking the ‘Submitting on Behalf of Consortium’ link within the navigation tree.

The ‘Submitting on Behalf of Consortium’ screen allows you to enter the contact information for each consortium member. To add each consortium member click the ‘Add’ button. This will generate a new set of consortium fields. Once you have entered data into one consortium field set, you must click the ‘Save’ button located within that field set prior to adding another member. If the data previously entered and saved within a consortium field set needs to be edited, you can click the ‘Edit’ button located in the upper right corner of that field set. If a consortium member needs to be deleted, you can click the ‘Delete’ icon that is also located in the upper right corner of each field set. There must be at least two consortium members for an entered consortium.

The system allows you to expand/collapse each consortium field set by clicking the consortium field set header. If you wish to expand/collapse all consortium field sets simultaneously, click the ‘Expand All’ or ‘Collapse All’ links located at the top right of all consortium field sets.

- **Consortium Name:** Enter the name of the consortium.
- **Consortium CBI:** Checking the CBI checkbox will claim the consortium as CBI.
- **Prefix:** The ‘Prefix’ drop-down menu displays all optional prefixes and captures the consortium member contact’s prefix. If applicable, select the appropriate prefix.
- **First Name:** Enter the consortium member contact’s first name. This field is required.
- **Middle Initial:** Enter the consortium member contact’s middle initial.
- **Last Name:** Enter the consortium member contact’s last name. This field is required.
- **Suffix:** The ‘Suffix’ drop-down menu displays all optional suffixes and captures the consortium member contact’s suffix. If applicable, select the appropriate suffix.
- **Company Name:** Enter the consortium member contact’s company name. This field is required.
- **Phone Number:** Enter the consortium member contact’s daytime phone number; U.S. or international. This field is required.
- **Ext:** Enter the extension to the consortium member contact’s daytime phone number.
- **Email Address:** Enter the consortium member contact’s email address. The email address must follow the standard email address format: john.doe@abc.com. This field is required.
- **Mailing Address 1:** Enter the first line of the consortium member contact’s mailing address. This field is required.
- **Mailing Address 2:** Enter the second line of the consortium member contact’s mailing address.
- **City:** Enter the city of the consortium member contact’s mailing address. This field is required.
- **State:** Enter the state of the consortium member contact’s mailing address. This field is required when ‘United States’ is identified as the country.

- **Postal Code:** Enter the postal code of the consortium member contact’s mailing address. This field is required.
- **Country:** Enter the country of the consortium contact’s mailing address. This field is required.

Exhibit 7-3 shows the screen capture for the ‘Submitting on Behalf of Consortium’ screen:

Exhibit 7-3 Submitting on Behalf of Consortium Screen

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

Section 4 § 40 CFR 766 Dibenzodioxins / Dibenzofurans > Contact Information > Submitting on Behalf of Consortium

SUBMITTING ON BEHALF OF CONSORTIUM

Provide the consortium name and click the Add button to enter member contact information.

Consortium Name:

Consortium CBI:

Expand All | Collapse All

CGI FEDERAL (John Doe) Edit ✕

Cancel ✕

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:

Save

Click the **Add** button to add a new consortium member. Add

Previous Next

Validate Save Preview Submit

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

Navigation: Click the ‘Next’ button to navigate to the ‘Chemical Identification’ screen if submitting a letter of intent. You can also access this screen by clicking the ‘Chemical Identification’ link within the navigation tree.

7.4 Submitting on Behalf of Company Screen

The ‘Submitting on Behalf of Company’ screen presents a list of fields that allow you to input the contact information for the manufacturing or processing establishment on whose behalf the submission is being made.

- **Prefix:** The ‘Prefix’ drop-down menu displays all optional prefixes. If applicable, select the appropriate prefix.
- **First Name:** Enter the first name of the manufacturing or processing establishment contact for which the submission is being made. This field is required.
- **Middle Initial:** Enter the middle initial of the manufacturing or processing establishment contact for which the submission is being made.
- **Last Name:** Enter the last name of the manufacturing or processing establishment contact for which the submission is being made. This field is required.
- **Suffix:** The ‘Suffix’ drop-down menu displays all optional suffixes and captures the suffix of the manufacturing or processing establishment contact for which the submission is being made. If applicable, select the appropriate suffix.
- **Company Name:** Enter the company name of the manufacturing or processing establishment. This field is required.
- **Phone Number:** Enter the daytime phone number of the manufacturing or processing establishment contact for whom the submission is being made; U.S. or international. This field is required.
- **Email Address:** Enter the email address of the manufacturing or processing establishment contact for which the submission is being made. The email address must follow the standard email address format: john.doe@abc.com. This field is required.
- **Mailing Address 1:** Enter the first line of the mailing address of the manufacturing or processing establishment contact. This field is required.
- **Mailing Address 2:** Enter the second line of the mailing address of the manufacturing or processing establishment contact.
- **City:** Enter the city of the mailing address of the manufacturing or processing establishment contact. This field is required.
- **State:** Enter the state of the mailing address of the manufacturing or processing establishment contact. This is a required field if the ‘Country’ field is defined as ‘United States.’
- **Postal Code:** Enter the postal code of the mailing address of the manufacturing or processing establishment contact. This field is required.
- **Country:** Enter the country of the mailing address of the manufacturing or processing establishment contact. This field is required.

Exhibit 7-4 shows the screen capture of the ‘Submitting on Behalf of Company’ screen:

Exhibit 7-4 Submitting on Behalf of Company Screen

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

Section 4 § 40 CFR 766 Dibenzodioxins / Dibenzofurans > Contact Information > Submitting on Behalf of Company

SUBMITTING ON BEHALF OF COMPANY

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

CBI:

Prefix:

First Name:

Middle Initial:

Last Name:

Suffix:

Company Name:

Phone Number: Ext:

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

Email Address:

Mailing Address 1:

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

Mailing Address 2:

Apartment, suite, etc.

City:

State:

Postal Code:

Country:

Previous Next

Validate Save Preview Submit

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Navigation: If you are submitting a letter of intent and click the ‘Next’ button, you are taken to the ‘Chemical Identification’ screen. If you are submitting a study plan or results, you are taken to the ‘Document Management’ screen.

7.4.1 Chemical Identification Screen

The ‘Chemical Identification’ screen requires you to identify the chemical substance the sponsor(s) intends to use in each of the tests. To identify the chemical substance select a chemical from the ‘Please select, or being typing, a CASRN in the drop-down menu below.’ The system allows you to claim the chemical as CBI, by checking the ‘CBI’ checkbox.

Exhibit 7-5 shows the screen capture of the ‘Chemical Identification’ screen:

Exhibit 7-5 Chemical Identification Screen

Navigation: Click the ‘Next’ button to navigate to the ‘Sponsoring Firms’ screen if you checked the ‘Letter of Intent’ checkbox on the ‘Submission Information Type’ screen. If all information has been completed please click the ‘Submit’ icon in the action bar to begin the submission process.

7.5 Letter of Intent

If you have identified that you are submitting a letter of intent, you are required to complete the following screens to satisfy the reporting requirements.

7.5.1 Sponsoring Firms Screen

To access the ‘Sponsoring Firms’ screen click the ‘Next’ button from the ‘Chemical Identification’ screen. You can also access the ‘Sponsoring Firms’ screen by clicking the ‘Sponsoring Firms’ link located within the navigation tree under the ‘Letter of Intent’ folder.

The ‘Sponsoring Firms’ screen requires you to identify each sponsoring firm involved in the testing of the identified chemical substances. To identify a sponsoring firm click the ‘Add Sponsoring Firm’ button. Clicking the ‘Add Sponsoring Firm’ button generates a sponsoring firm field set. Once you have entered data into a sponsoring firm field set, you must click the ‘Save’ button located within that field set prior to adding another firm. If the data previously entered and saved within a field set needs to be edited, you can click the ‘Edit’ button located in the upper right corner of that field set. If a sponsoring firm needs to be deleted, you can click the ‘Delete’ icon that is also located in the upper right corner of each field set.

The system allows you to expand/collapse each field set by clicking the sponsoring firm field set header. If you wish to expand/collapse all field sets simultaneously, click the 'Expand All' or 'Collapse All' links located at the top right of all consortium field sets.

- **CBI:** Check the checkbox to claim the sponsoring firm as CBI.
- **Firm:** Enter the sponsoring firm's company name. This field is required.
- **Phone Number:** Enter the sponsoring firm's daytime phone number; U.S. or international. This field is required.
- **Ext:** Enter the sponsoring firm's extension to the listed daytime phone number.
- **Mailing Address 1:** Enter the first line of the sponsoring firm's mailing address. This field is required.
- **Mailing Address 2:** Enter the second line of the sponsoring firm's mailing address.
- **City:** Enter the city of the sponsoring firm's mailing address. This field is required.
- **State:** Enter the state of the sponsoring firm's mailing address. This field is required when 'United States' is identified as the country.
- **Postal Code:** Enter the postal code of the sponsoring firm's mailing address. This field is required.
- **Country:** Enter the country of the sponsoring firm's mailing address. This field is required.

Exhibit 7-6 shows the screen capture of the ‘Sponsoring Firms’ screen:

Exhibit 7-6 Sponsoring Firms Screen

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

Section 4 § 40 CFR 766 Dibenzodioxins / Dibenzofurans > Letter of Intent > Sponsoring Firms
SPONSORING FIRMS

Fill out the information below for the Sponsoring Firm.

Expand All | Collapse All

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

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Navigation: Click the ‘Next’ button to navigate to the ‘Document Management’ screen if you have identified that you are submitting a protocol or results. If you are only completing a ‘Letter of Intent,’ click the ‘Submit’ icon located in the action bar to begin the submission process.

7.6 Protocol

If you have identified you are submitting a protocol, you are required to complete the following screens to satisfy the reporting requirements.

7.6.1 Protocol Document Management Screen

The ‘Document Management’ screen allows you to upload documents related to Section 4 reporting requirements. To add protocol documentation, click the ‘Attach Document’ button. Clicking the ‘Attach Document’ button displays the ‘Protocol Document Management’ pop-up window.

The 'Protocol Document Management' pop-up window allows you to upload files from your hard drive. Click the 'Browse' button to generate the 'File Upload' window. The 'File Upload' window allows you to search for the desired document from your hard drive. Once you have identified the document to be attached, click the 'Open' button. Clicking the 'Open' button will close the 'File Upload' window and display the attached document within the 'Browse' field. To claim attached documentation CBI, check the 'CBI' checkbox. If you claim the attached documentation as CBI, you are able to attach the sanitized version of the attached document. To do so, follow the steps outlined above. Once you have attached all desired documentation, click the 'OK' button to return to the 'Document Management' screen.

The document management table displays the file name of the document, the date the document was attached, and whether the document was claimed as CBI. The system allows you to delete unwanted documents by clicking the 'Delete' icon located under the 'Action' column for a given row. Clicking the file name links within the 'File Name' column generates the 'Protocol Document Management' pop-up window. This allows you to view and edit previously attached documents.


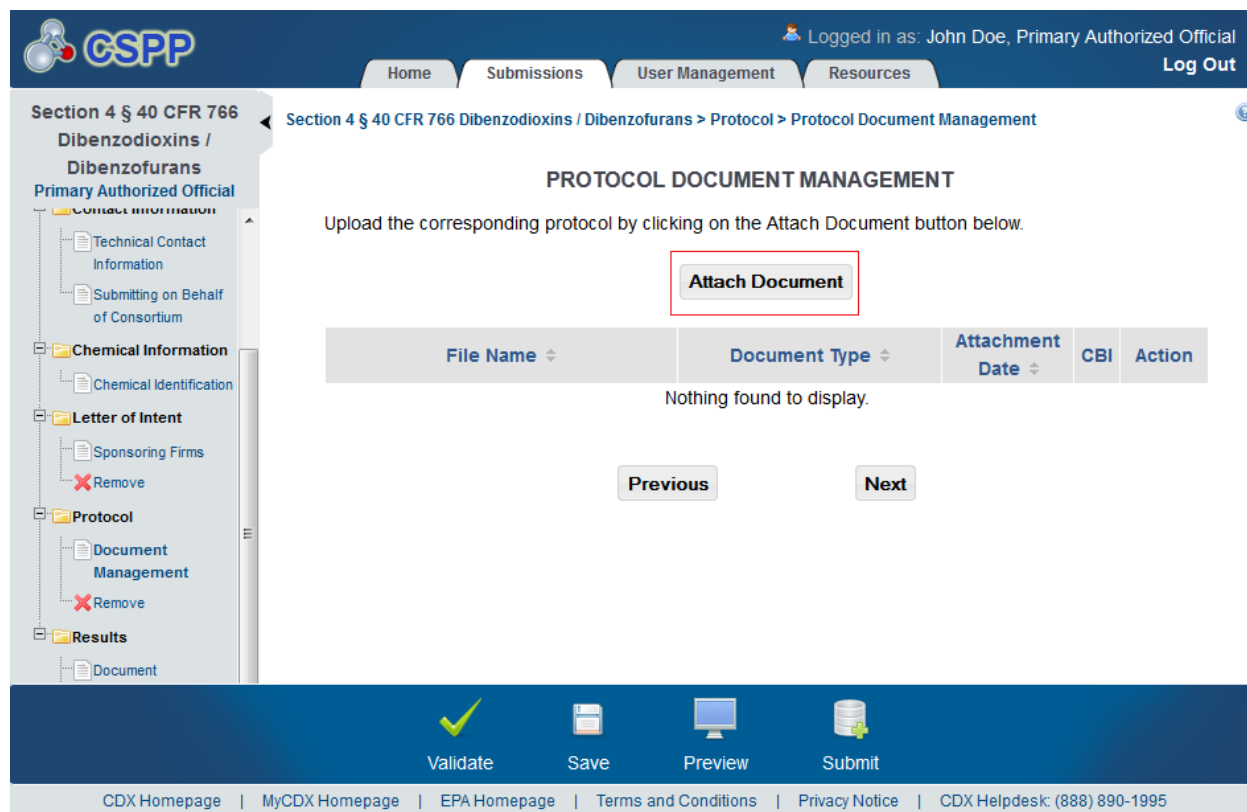
- **Attach Document:** Clicking the 'Attach Document' button displays the 'Protocol Document Management' pop-up window.
- **File Name:** Displays the name of the attached file. Clicking the file name link generates the 'Protocol Document Management' pop-up window and allows you to view the attached document.
- **Document Type:** Displays the type of document attached.
- **Attachment Date:** Displays the date the document was attached.
- **CBI:** Displays whether the attached document was claimed as CBI by displaying a 'Y' or an 'N.'
- **Action:** Clicking the 'Delete' icon () will generate a prompt confirming deletion. To delete the specified row and all contents within click the 'OK' button within the prompt.

Exhibit 7-7 shows the screen capture of the ‘Protocol Document Management’ screen:

Exhibit 7-7 Document Management Screen



Navigation: Click the ‘Next’ button to navigate to the ‘Document Management’ screen if you have identified that you are submitting results to EPA. If you are only submitting a protocol click the ‘Submit’ icon located in the action bar to begin the submission process.

The ‘Protocol Document Management’ pop-up window allows you to browse for a document from your hard drive and attach one document at a time. You must attach at least one document. Clicking the ‘Browse’ button will generate the ‘File Upload’ window on your hard drive. Select the document you wish to attach and click the ‘Open’ button. If a document is marked as ‘CBI’ then a sanitized copy of the document must be attached. To save the attached document to the ‘Document Management’ screen, click the ‘OK’ button within the ‘Protocol Document Management’ pop-up window.

- **CBI:** Checking this ‘CBI’ checkbox will claim the document as CBI and generate a sanitized document upload field.
- **Document Upload:** Clicking the ‘Browse’ button will generate the ‘File Upload’ window for you to search for a particular file on your hard drive.
- **Sanitized Document Upload:** Clicking the ‘Browse’ button will generate the ‘File Upload’ window for you to search for a particular file on your hard drive. This field is only generated when the ‘CBI’ checkbox is checked and is optional.
- **Effects:** Select the effect specific to the chemical report. Displays the following options: ‘Health Effects,’ ‘Environmental Fate,’ ‘Ecotoxicity,’ and ‘Physical Chemical Properties.’

- **Endpoints:** Select an endpoint that is specific to the chemical report. The endpoints will be specific to the effect chosen from the ‘Effects’ drop-down menu. If no effect is chosen, no endpoints will display.

Refer to Appendix B. Submission Types, Effects, and Endpoints for a complete list of values that are found within each of the drop-down menus on the ‘Document Management’ pop-up window. If you have multiple documents that share the same effects and/or endpoints, you must upload each individual document separately and select values from the drop-down menus.

Exhibit 7-8 shows the screen capture of the ‘Protocol Document Management’ pop-up window:

Exhibit 7-8 Protocol Document Management Pop-Up Window

The screenshot shows a web-based form titled "PROTOCOL DOCUMENT MANAGEMENT" from CSPP. The form includes a note about security and validation errors. It has fields for "CBI" (checked), "Document Type" (dropdown), "Document Upload" (text input with "Browse" button), "Sanitized Document Upload" (text input with "Browse" button and "(optional)" text), "Effects" (dropdown), and "EndPoints" (dropdown). "OK" and "Cancel" buttons are at the bottom, with "OK" highlighted by a red box.

Navigation: Click the ‘OK’ button to return to the ‘Document Management’ screen.

7.7 Results

If you have identified you are submitting results, you are required to complete the following screens to satisfy the reporting requirements.

7.7.1 Document Management Screen

The ‘Document Management’ screen allows you to upload documents related to Section 4 reporting requirements. To add Section 4 results documentation, click the ‘Attach Document’ button. Clicking the ‘Attach Document’ button displays the ‘Document Management’ pop-up window.

The ‘Document Management’ pop-up window allows you to attach documentation by clicking the ‘Browse’ button. This will generate the ‘File Upload’ window where you can select the

desired document from your hard drive. Once you have identified the document to be attached, click the 'Open' button. Clicking the 'Open' button will close the 'File Upload' window and display the attached document within the 'Browse' field. To claim attached documentation CBI, check the 'CBI' checkbox. If you claim the attached documentation as CBI you are able to attach the sanitized version of the attached document. To do so, follow the steps outlined above. Once you have attached all desired documentation, click the 'OK' button to return to the 'Document Management' screen.

The document management table displays the file name of the document, the date the document was attached, and whether the document was claimed as CBI. The system allows you to delete unwanted documents by clicking the 'Delete' icon located under the 'Action' column for a given row. Clicking the file name links within the 'File Name' column generates the 'Document Management' pop-up window. This allows you to view and edit previously attached documents.


- **Attach Document:** Clicking the 'Attach Document' button displays the 'Document Management' pop-up window.
- **File Name:** Displays the name of the attached file. Clicking the file name link generates the 'Document Management' pop-up window and allows you to view the attached document.
- **Attachment Date:** Displays the date the document was attached.
- **CBI:** Displays whether the attached document was claimed as CBI by displaying a 'Y' or an 'N.'
- **Action:** Clicking the 'Delete' icon () will generate a prompt confirming deletion. To delete the specified row and all contents within click the 'OK' button within the prompt.

Exhibit 7-9 shows the screen capture of the 'Document Management' screen:

Exhibit 7-9 Document Management Screen

The screenshot displays the 'DOCUMENT MANAGEMENT' interface. At the top, the user is logged in as 'John Doe, Primary Authorized Official'. The navigation menu on the left includes 'Section 4 § 40 CFR 766 Dibenzenodioxins / Dibenzenofurans' and 'Primary Authorized Official'. The main content area contains the text: 'To add a Results Document, please click the Attach Document button and enter the information in the popup.' The 'Attach Document' button is highlighted with a red box. Below this is a table with the following structure:

File Name	Attachment Date	CBI	Action
Nothing found to display.			

Below the table is a 'Previous' button. At the bottom of the page is a blue action bar with icons for 'Validate', 'Save', 'Preview', and 'Submit'. The footer contains links to 'CDX Homepage', 'MyCDX Homepage', 'EPA Homepage', 'Terms and Conditions', 'Privacy Notice', and 'CDX Helpdesk: (888) 890-1995'.

Navigation: Click the 'Previous' button to navigate to the 'Document Management' screen if you have identified you are submitting a protocol to EPA. If you have completed all required fields, click the 'Submit' icon located in the action bar to begin the submission process.

The 'Document Management' pop-up window allows you to browse for a document from your hard drive and attach one document at a time. You must attach at least one document. Clicking the 'Browse' button will generate the 'File Upload' window on your hard drive. Select the document you wish to attach and click the 'Open' button. If a document is marked as 'CBI' then a sanitized copy of the document must be attached. To save the attached document to the 'Document Management' screen, click the 'OK' button within the 'Document Management' pop-up window.


- **CBI:** Checking this 'CBI' checkbox will claim the document as CBI and generate a sanitized document upload field.
- **Document Upload:** Clicking the 'Browse' button will generate the 'File Upload' window for you to search for a particular file on your hard drive.
- **Sanitized Document Upload:** Clicking the 'Browse' button will generate the 'File Upload' window for you to search for a particular file on your hard drive. This field is only generated when the 'CBI' checkbox is checked and is optional.
- **Effects:** Select the effect specific to the chemical report. Displays the following options: 'Health Effects,' 'Environmental Fate,' 'Ecotoxicity,' and 'Physical Chemical Properties.'

- **Endpoints:** Select an endpoint that is specific to the chemical report. The endpoints will be specific to the effect chosen from the ‘Effects’ drop-down menu. If no effect is chosen, no endpoints will display.

Refer to Appendix B. Submission Types, Effects, and Endpoints for a complete list of values that are found within each of the drop-down menus on the ‘Document Management’ pop-up window. If you have multiple documents that share the same effects and/or endpoints, you must upload each individual document separately and select values from the drop-down menus.

Exhibit 7-10 shows the screen capture of the ‘Document Management’ pop-up window:

Exhibit 7-10 Document Management Pop-Up Window



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**

Navigation: Click the ‘OK’ button to return to the ‘Document Management’ screen.

7.8 User Requests

If you have identified you are submitting a user request, you are required to complete the following screens to satisfy the reporting requirements.

7.8.1 Waiver Screen

You can access the ‘Waiver’ screen by clicking the ‘Next’ button from the ‘Chemical Identification’ screen. You can also access this screen by clicking the ‘Waiver’ link within the navigation tree.

The ‘Waiver’ screen requires you to identify the appropriate qualifying waiver reason and allows you to upload documents related to Section 4 reporting requirements. To add Section 4 waiver

documentation, click the 'Attach Document' button. Clicking the 'Attach Document' button displays the 'Waiver Document Management' pop-up window.

The 'Waiver Document Management' pop-up window allows you to attach documentation by clicking the 'Browse' button. This will generate the 'File Upload' window where you can search for the desired document from your hard drive. Once you have identified the document to be attached, click the 'Open' button. Clicking the 'Open' button will close the 'File Upload' window and display the attached document within the 'Browse' field. To claim attached documentation as CBI, check the 'CBI' checkbox. If you claim the attached documentation as CBI you are required to attach the sanitized version of the attached document. To do so, follow the steps outlined above. Once you have attached all desired documentation, click the 'OK' button to return to the 'Waiver' screen.

The waiver table displays the file name of the document, the date the document was attached, and whether the document was claimed as CBI. The system allows you to delete unwanted documents by clicking the 'Delete' icon located under the 'Action' column for a given row. Clicking the file name links within the 'File Name' column generates the 'Waiver Document Management' pop-up window. This allows you to view and edit previously attached documents.


- **The chemical substance is produced only in quantities of 100 kilograms or less per year, only for research and development purposes:** Selecting this radio button option identifies this as the qualifying reason for the Waiver request.
- **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.):** Selecting this radio button option identifies this as the qualifying reason for the waiver request.
- **Attach Document:** Clicking the 'Attach Document' button displays the 'Waiver Document Management' pop-up window.
- **File Name:** Displays the name of the attached file. Clicking the file name link generates the 'Waiver Document Management' pop-up window and allows you to view and edit the attached document.
- **Attachment Date:** Displays the date the document was attached.
- **CBI:** Displays whether the attached document was claimed as CBI by displaying a 'Y' or an 'N.'
- **Action:** Clicking the 'Delete' icon () will generate a prompt confirming deletion. To delete the specified row and all contents within click the 'OK' button within the prompt.

Exhibit 7-11 shows the screen capture of the ‘Waiver’ screen:

Exhibit 7-11 Waiver Screen

Section 4 § 40 CFR 766 Dibenzenodioxins / Dibenzenofurans
Primary Authorized Official

Section 4 § 40 CFR 766 Dibenzenodioxins / Dibenzenofurans > User Requests > Waiver

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

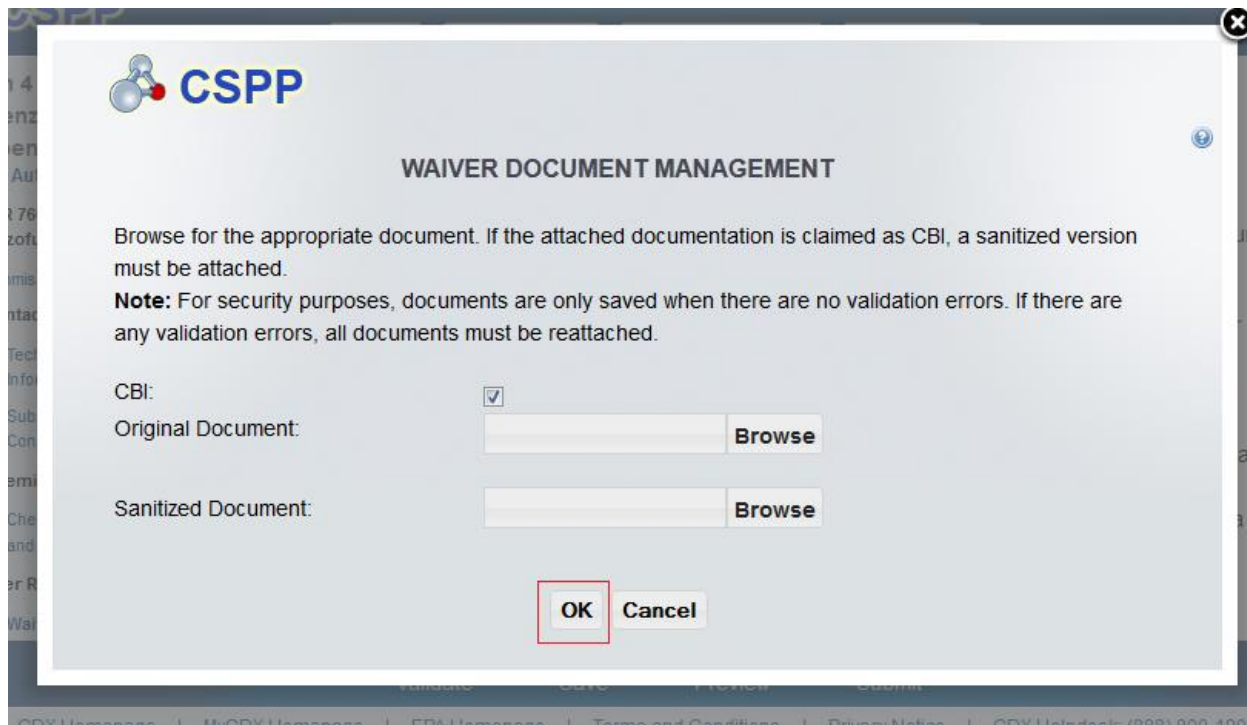
Validate Save Preview Submit

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Navigation: Click the ‘Previous’ button to navigate to the ‘Chemical Identification’ screen.

Exhibit 7-12 shows the screen capture of the ‘Waiver Document Management’ pop-up window:

Exhibit 7-12 Waiver Document Management Pop-Up Window



Navigation: Click the ‘OK’ button to return to the ‘Waiver’ screen.

7.8.2 Exclusion Screen

You can access the ‘Exclusion’ screen by clicking the ‘Next’ button from the ‘Chemical Identification’ screen. You can also access this screen by clicking the ‘Exclusion’ link within the navigation tree.

The ‘Exclusion’ screen requires you to identify the appropriate qualifying exclusion reason and allows you to upload documents related to Section 4 reporting requirements. To add Section 4 exclusion documentation, click the ‘Attach Document’ button. Clicking the ‘Attach Document’ button displays the ‘Exclusion Document Management’ pop-up window.

The ‘Exclusion Document Management’ pop-up window allows you to attach documentation by clicking the ‘Browse’ button. This will generate the ‘File Upload’ window where you can select the desired document from your hard drive. Once you have identified the document to be attached, click the ‘Open’ button. Clicking the ‘Open’ button will close the ‘File Upload’ window and display the attached document within the ‘Browse’ field. To claim attached documentation CBI, check the ‘CBI’ checkbox. If you claim the attached documentation as CBI you are able to attach the sanitized version of the attached document. To do so, follow the steps outlined above. Once you have attached all desired documentation, click the ‘OK’ button to return to the ‘Exclusion’ screen.

The exclusion table displays the file name of the document, the date the document was attached, and whether the document was claimed as CBI. The system allows you to delete unwanted documents by clicking the ‘Delete’ icon located under the ‘Action’ column for a given row.

Clicking the file name links within the 'File Name' column generates the 'Exclusion Document Management' pop-up window. This allows you to view and edit previously attached documents.


- **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:** Selecting this radio button option identifies this as the qualifying reason for the exclusion request.
- **Process and reaction conditions of the chemical substance such that no HDDs/HDFs could be produced under those conditions:** Selecting this radio button option identifies this as the qualifying reason for the exclusion request.
- **Attach Document:** Clicking the 'Attach Document' button displays the 'Exclusion Document Management' pop-up window.
- **File Name:** Displays the name of the attached file. Clicking the file name link generates the 'Exclusion Document Management' pop-up window and allows you to view the attached document.
- **Attachment Date:** Displays the date the document was attached.
- **CBI:** Displays whether the attached document was claimed as CBI by displaying a 'Y' or an 'N.'
- **Action:** Clicking the 'Delete' icon () will generate a prompt confirming deletion. To delete the specified row and all contents within click the 'OK' button within the prompt.

Exhibit 7-13 shows the screen capture of the ‘Exclusion’ screen:

Exhibit 7-13 Exclusion Screen

Section 4 § 40 CFR 766 Dibenzenodioxins / Dibenzenofurans Primary Authorized Official

Section 4 § 40 CFR 766 Dibenzenodioxins / Dibenzenofurans > 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

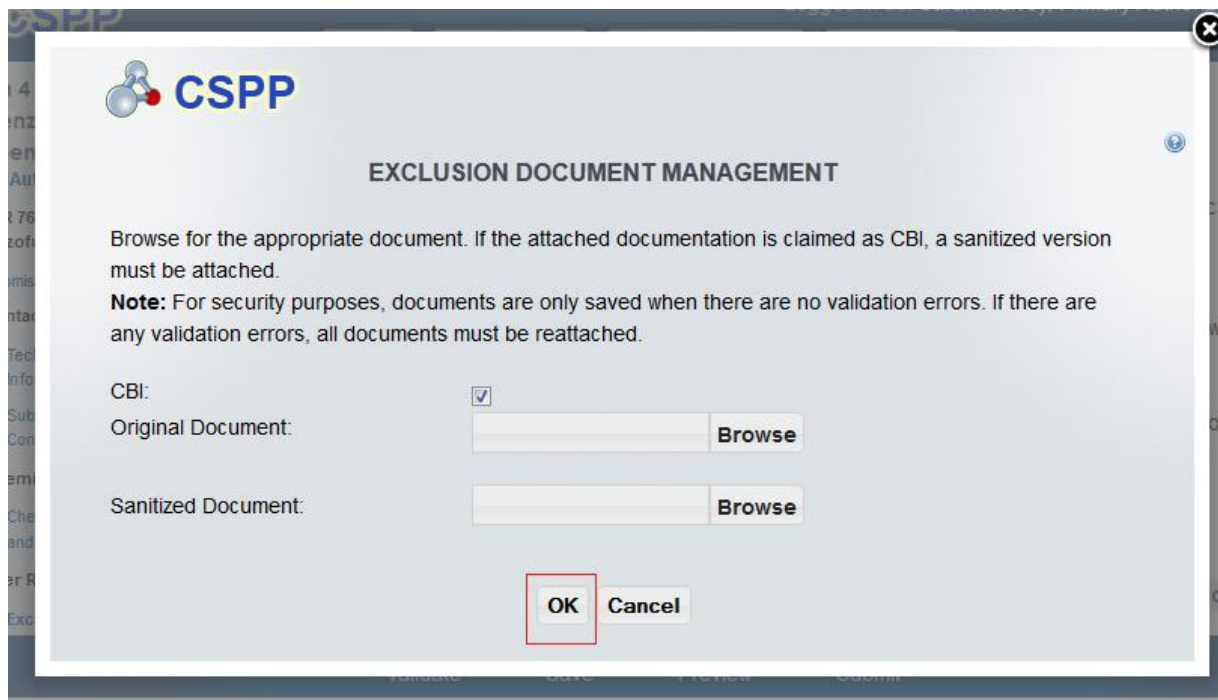
Validate Save Preview Submit

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Navigation: Click the ‘Previous’ button to navigate to the ‘Chemical Identification’ screen.

Exhibit 7-14 shows the screen capture of the ‘Exclusion Document Management’ pop-up window:

Exhibit 7-14 Exclusion Document Management Pop-Up Window



Navigation: Click the ‘OK’ button to return to the ‘Exclusion’ screen.

7.8.3 Exemption Screen

You can access the ‘Exemption’ screen by clicking the ‘Next’ button from the ‘Chemical Identification’ screen. You can also access this screen by clicking the ‘Exemption’ link within the navigation tree.

The ‘Exemption’ screen allows you to upload documents related to Section 4 reporting requirements. To add Section 4 exemption documentation, click the ‘Attach Document’ button. Clicking the ‘Attach Document’ button displays the ‘Exemption Document Management’ pop-up window.

The ‘Exemption Document Management’ pop-up window allows you to attach documentation by clicking the ‘Browse’ button. This will generate the ‘File Upload’ window where you can select the desired document from your hard drive. Once you have identified the document to be attached, click the ‘Open’ button. Clicking the ‘Open’ button will close the ‘File Upload’ window and display the attached document within the ‘Browse’ field. To claim attached documentation as CBI, check the ‘CBI’ checkbox. If you claim the attached documentation as CBI, you are required to attach the sanitized version of the attached document. To do so, follow the steps outlined above. Once you have attached all desired documentation, click the ‘OK’ button to return to the ‘Exemption’ screen.

The exemption table displays the file name of the document, the date the document was attached, and whether the document was claimed as CBI. The system allows you to delete unwanted documents by clicking the ‘Delete’ icon located under the ‘Action’ column for a given row.

Clicking the file name links within the ‘File Name’ column generates the ‘Exemption Document Management’ pop-up window. This allows you to view and edit previously attached documents.

- **Attach Document:** Clicking the ‘Attach Document’ button displays the ‘Exemption Document Management’ pop-up window.
- **File Name:** Displays the name of the attached file. Clicking the file name link generates the ‘Exemption Document Management’ pop-up window and allows you to view the attached document.
- **Attachment Date:** Displays the date the document was attached.
- **CBI:** Displays whether the attached document was claimed as CBI by displaying a ‘Y’ or an ‘N.’
- **Action:** Clicking the ‘Delete’ icon (✖) will generate a prompt confirming deletion. To delete the specified row and all contents within click the ‘OK’ button within the prompt.

Exhibit 7-15 shows the screen capture of the ‘Exemption’ screen:

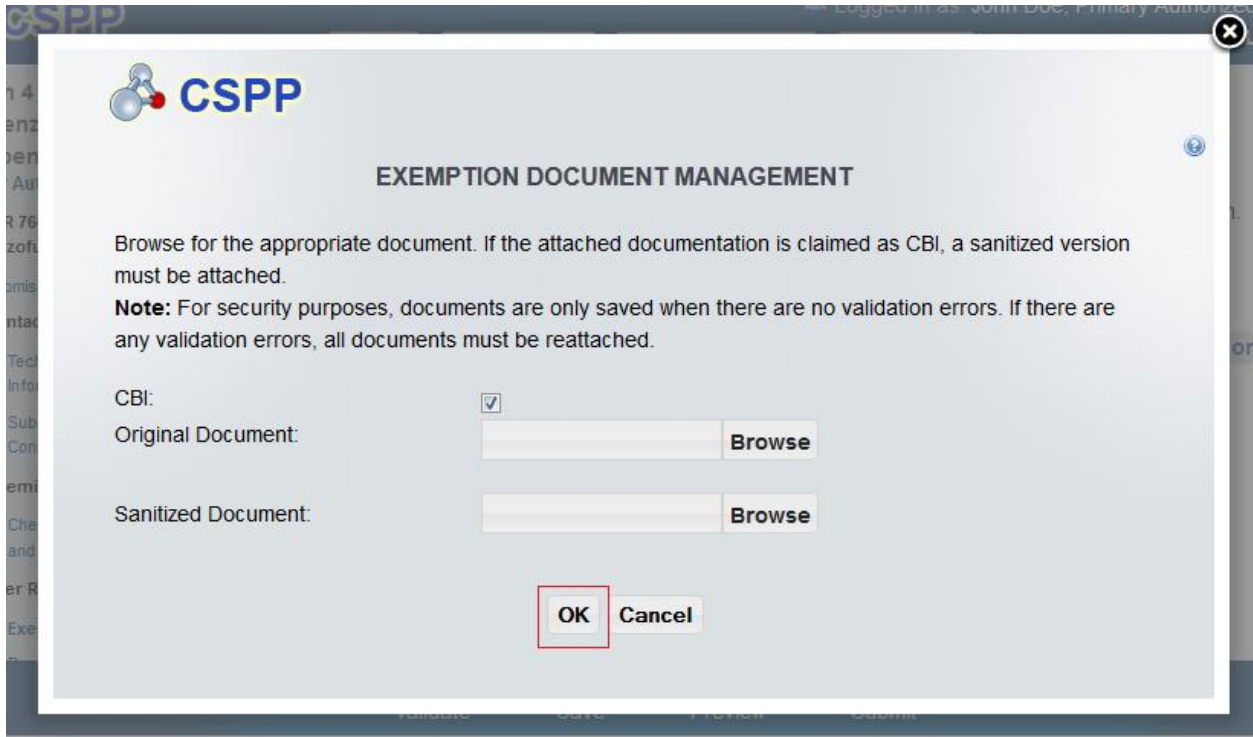
Exhibit 7-15 Exemption Screen

The screenshot shows the GSPP (Global Submission Processing Platform) interface. The user is logged in as John Doe, a Primary Authorized Official. The navigation menu includes Home, Submissions, User Management, and Resources. The current page is titled 'EXEMPTION' and is part of the 'Section 4 § 40 CFR 766 Dibenzodioxins / Dibenzofurans > User Requests > Document Management' path. The main content area contains the text: 'To request an exemption, please click the Attach Document button to attach required documentation.' A red box highlights the 'Attach Document' button. Below this is a table with columns: File Name, Attachment Date, CBI, and Action. The table is currently empty, displaying 'Nothing found to display.' A 'Previous' button is located below the table. The footer contains navigation links: Validate, Save, Preview, and Submit.

Navigation: Click the ‘Previous’ button to navigate to the ‘Chemical Identification’ screen.

Exhibit 7-16 shows the screen capture of the ‘Exemption Document Management’ pop-up window:

Exhibit 7-16 Exemption Document Management Pop-Up Window



Navigation: Click the ‘OK’ button to return to the ‘Exemption’ screen.

8 Validate

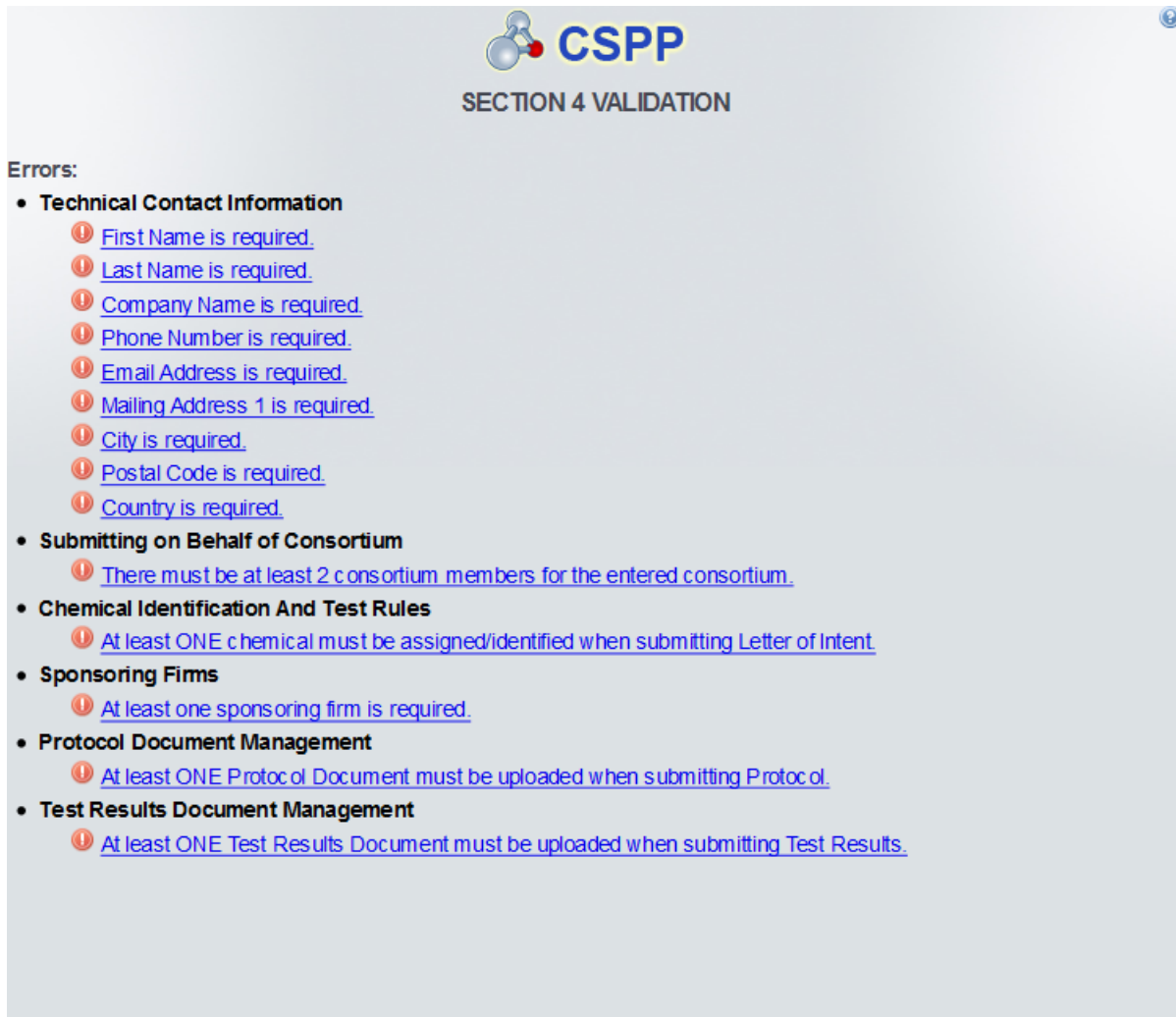
You can click the ‘Validate’ icon (✓) at any stage of completing a Section 4 submission. The ‘Section 4 Validation’ pop-up window is displayed when you click the ‘Validate’ icon (✓). The ‘Section 4 Validation’ pop-up window displays a report of all validation errors. During the validation process, the application validates each screen of the Section 4 submission to find missing and invalid data.

Validation Errors: Errors are denoted by an ‘Error’ icon (ⓘ) and can be fixed by clicking the error link. The links will display the *Screen Title Name* (e.g., Technical Contact Information) and the associated error. After you click a link, the main application screen will display the section where the error occurred so you can easily fix the error. Once you have fixed the error, click the ‘Validate’ icon (✓) again to refresh the ‘Section 4 Validation’ pop-up window. If the information you fixed passes validation, the error will be removed from the ‘Section 4 Validation’ pop-up window. You must fix all validation errors in order to submit the submission to CDX successfully.

You can close the ‘Section 4 Validation’ pop-up window by clicking the ‘X’ button located at the top right of the window.

Exhibit 8-1 shows the screen captures for the 'Section 4 Validation' pop-up window:

Exhibit 8-1 Section 4 Validation Pop-Up Window




9 Submit to EPA via CDX

As the Primary AO, you have the ability to sign and submit a Section 4 submission to EPA. Assigned Supports for your submission cannot submit the Section 4 submission.

Once you complete all required information and correct all items failing validation, the system will allow you to submit successfully.

9.1 Submitting Official Information Screen

Click the 'Submit' icon () that is located in the bottom action bar of the Section 4 application to access the 'Submitting Official Information' screen. This screen only displays to the Primary AO upon submission.


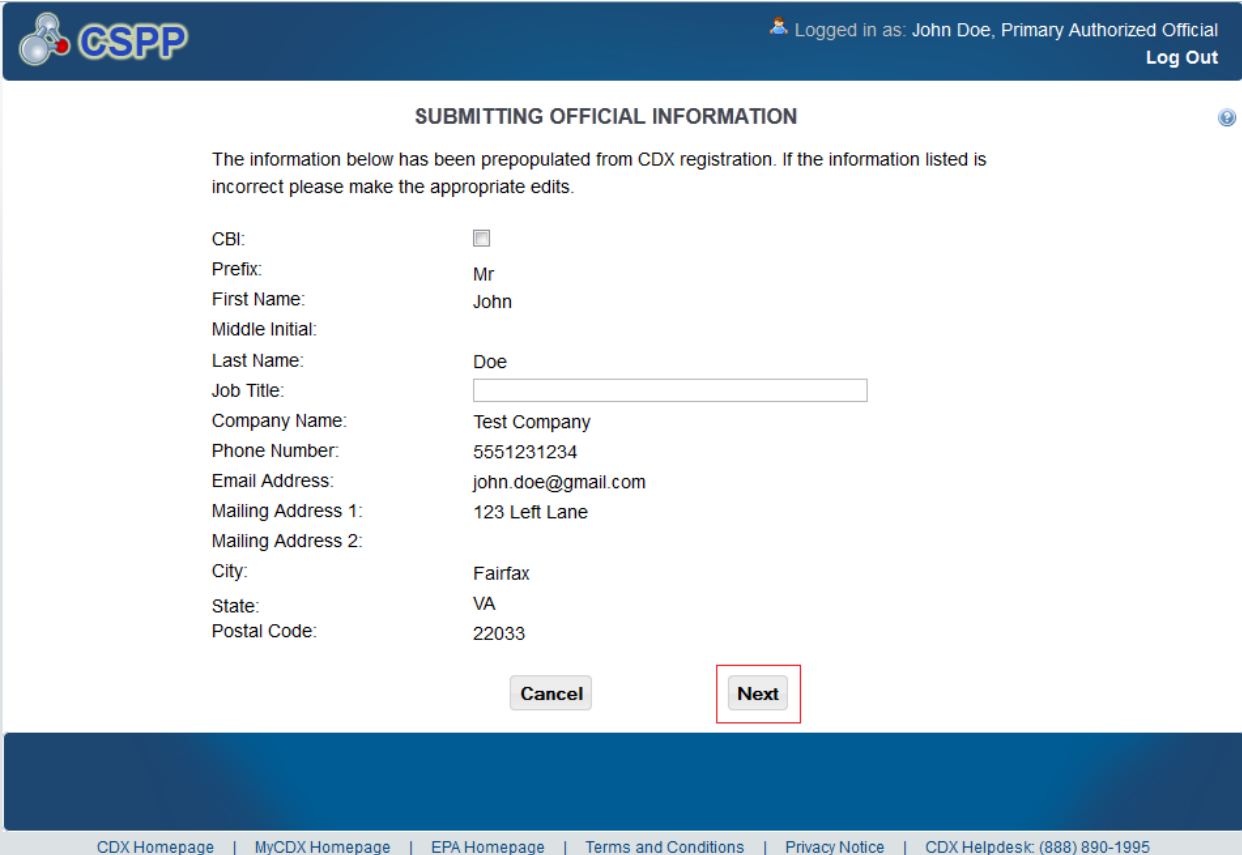
After clicking the 'Submit' icon (), the system requires you to review your contact information provided during CDX registration. The system also requires you to identify your job title.

Exhibit 9-1 shows the screen capture of the 'Submitting Official Information' screen:

Exhibit 9-1 Submitting Official Information Screen



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

SUBMITTING OFFICIAL INFORMATION

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

CBI:	<input type="checkbox"/>
Prefix:	Mr
First Name:	John
Middle Initial:	
Last Name:	Doe
Job Title:	<input type="text"/>
Company Name:	Test Company
Phone Number:	5551231234
Email Address:	john.doe@gmail.com
Mailing Address 1:	123 Left Lane
Mailing Address 2:	
City:	Fairfax
State:	VA
Postal Code:	22033

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

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

Navigation: Click the 'Next' button to continue with the submission process and proceed to the 'Submission Process: Validation' screen.

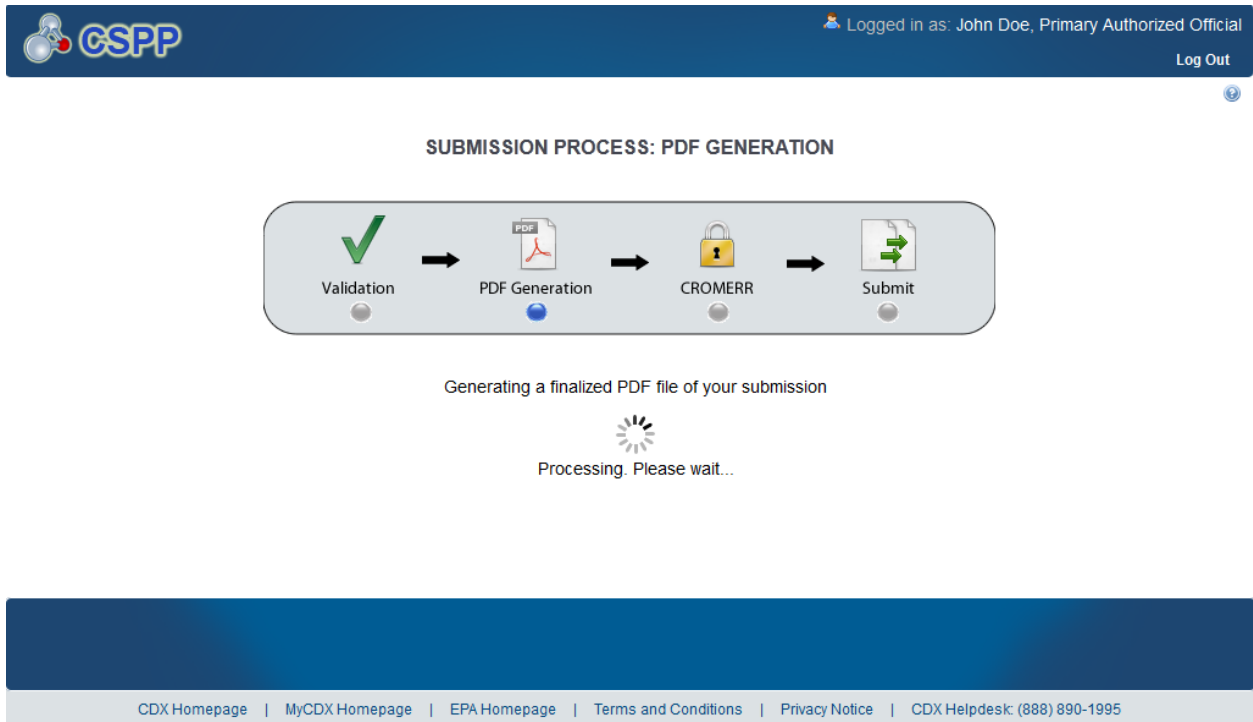
9.2 Submission Process: Validation Screen

Once all information is entered within the ‘Submitting Official Information’ screen, click the ‘Next’ button to proceed to the ‘Submission Process: Validation’ screen.

The ‘Submission Process: Validation’ screen will validate the information entered within the Section 4 submission.

Exhibit 9-2 shows the screen capture of the ‘Submission Process: Validation’ screen:

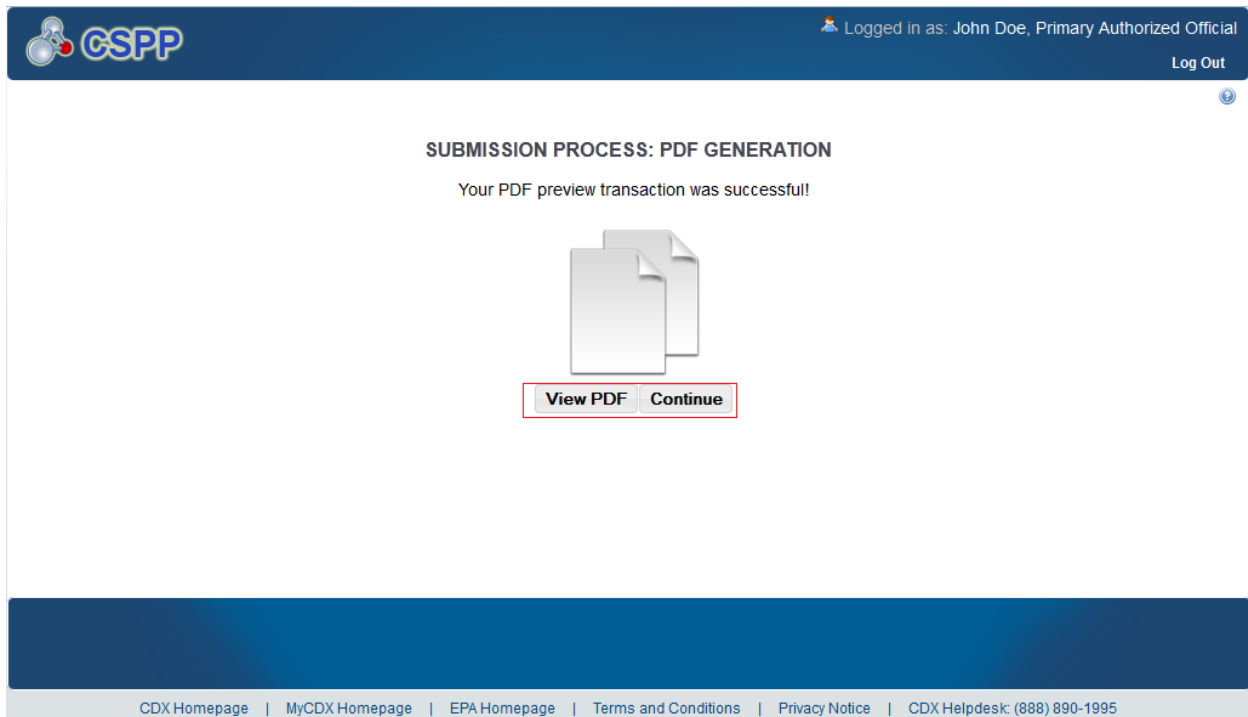
Exhibit 9-2 Submission Process: Validation Screen



After the Section 4 submission being submitted is verified, a PDF is generated for you to view.

Exhibit 9-3 shows the screen capture of the ‘Submission Process: PDF Generation’ screen:

Exhibit 9-3 Submission Process: PDF Generation Screen



Navigation: Click the ‘Continue’ button to continue with your submission and proceed to the ‘Cross-Media Electronic Reporting Regulation (CROMERR) Certification’ screen.

9.3 Cross-Media Electronic Reporting Regulation (CROMERR) Certification Screen

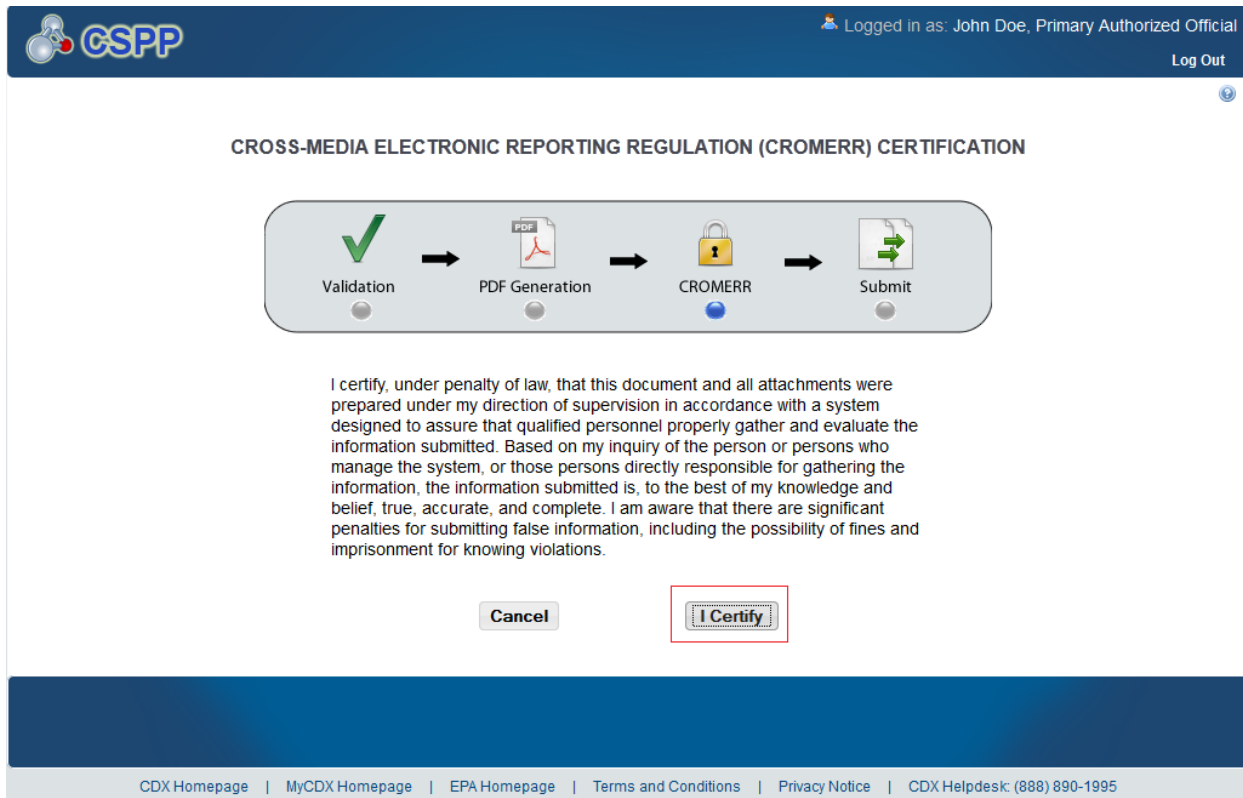
Click the ‘Continue’ button on the ‘Submission Process: PDF Generation’ screen to access the ‘Cross-Media Electronic Reporting Regulation (CROMERR) Certification’ screen.

CROMERR provides the legal framework for electronic reporting under all of EPA’s environmental regulations.

After clicking the ‘Continue’ button, the system requires you to certify that the submission is CROMERR compliant. By clicking the ‘I Certify’ button on the ‘CROMERR Certification’ screen; you authenticate all of the information entered in the submission. Clicking the ‘Cancel’ button will allow you to navigate back to the Section 4 ‘Home’ screen.

Exhibit 9-4 shows the screen capture for the ‘Cross-Media Electronic Reporting Regulation (CROMERR) Certification’ screen:

Exhibit 9-4 Cross-Media Electronic Reporting Regulation (CROMERR) Certification Screen



Navigation: Click the ‘I Certify’ button to continue with your submission and proceed to the ‘Cross-Media Electronic Reporting Regulation (CROMERR) Login’ screen.

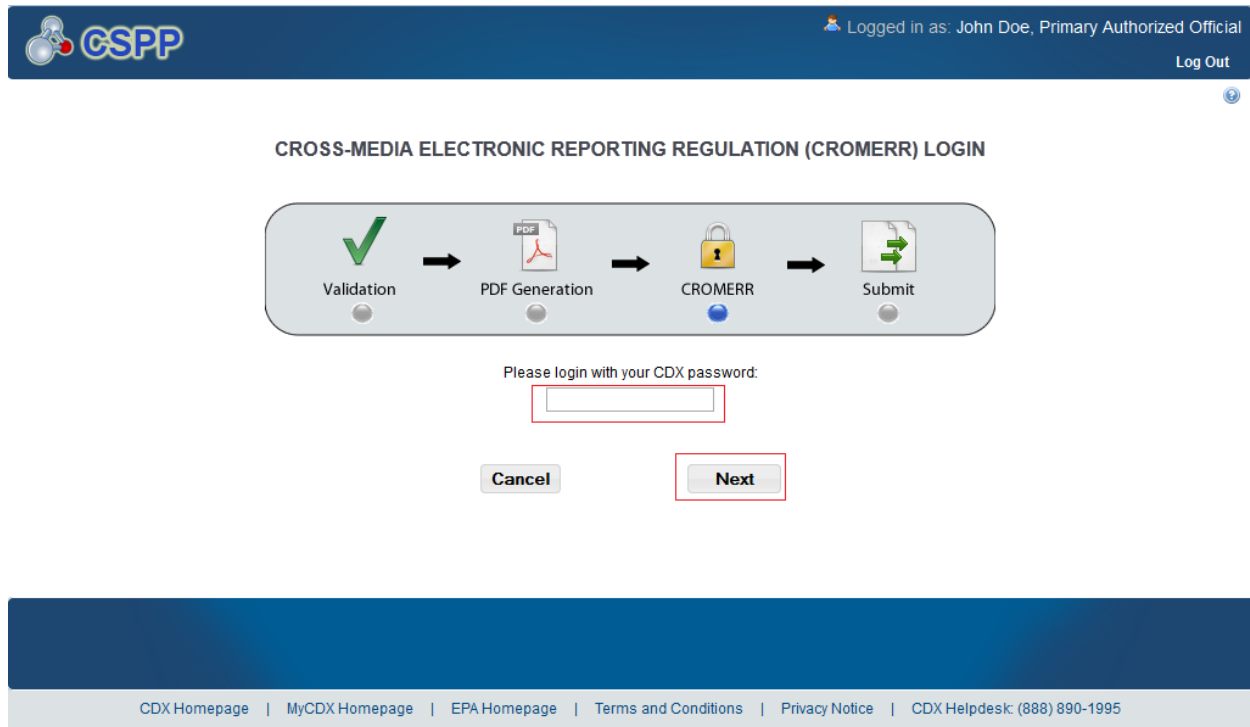
9.4 Cross-Media Electronic Reporting Regulation (CROMERR) Login Screen

You can access the ‘Cross-Media Electronic Reporting Regulation (CROMERR) Login’ screen by clicking the ‘I Certify’ button on the ‘Cross-Media Electronic Reporting Regulation (CROMERR) Certification’ screen.

- **Please login with your CDX password:** Enter your CDX registration password.

Exhibit 9-5 shows the screen capture of the ‘Cross-Media Electronic Reporting Regulation (CROMERR) Login’ screen:

Exhibit 9-5 Cross-Media Electronic Reporting Regulation (CROMERR) Login Screen



Navigation: Enter your CDX registration password and click the ‘Next’ button to navigate to the ‘Cross-Media Electronic Reporting Regulation (CROMERR) Security Question’ screen.

9.5 Cross-Media Electronic Reporting Regulation (CROMERR) Security Question Screen

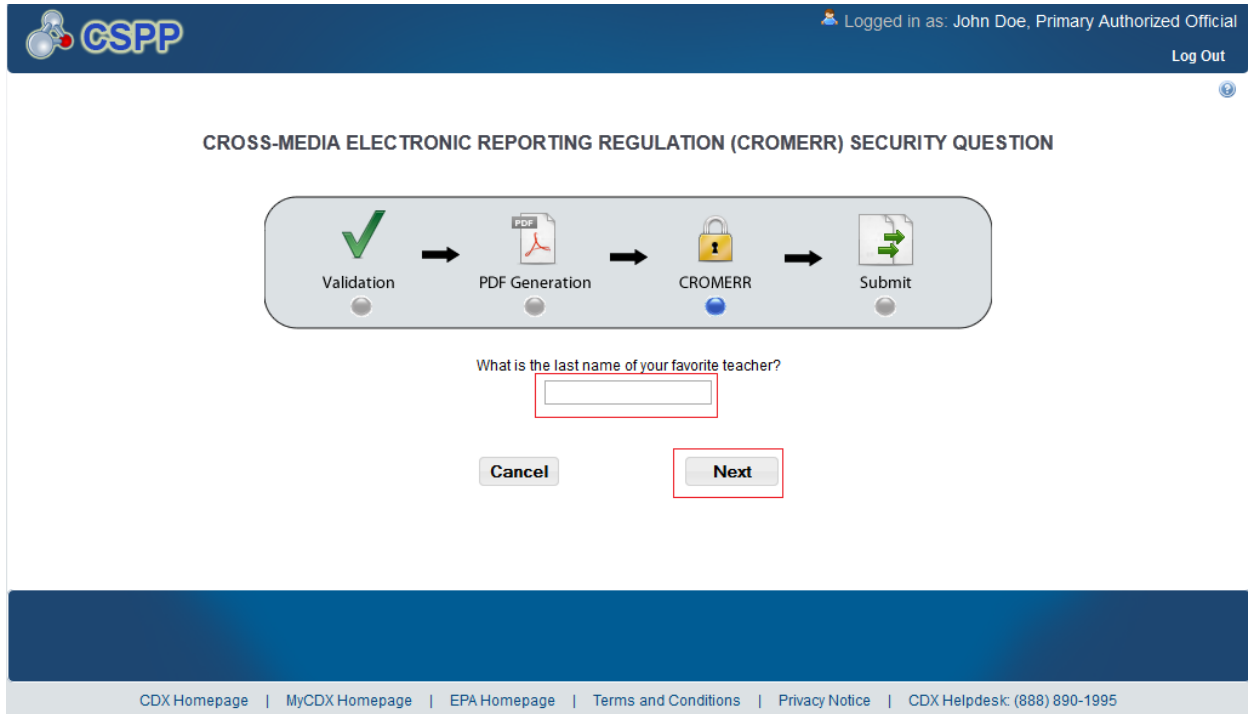
You can access the ‘Cross-Media Electronic Reporting Regulation (CROMERR) Security Question’ screen by entering your CDX registration password on the ‘Cross-Media Electronic Reporting Regulation (CROMERR) Login’ screen and by clicking the ‘Next’ button.

The ‘Cross-Media Electronic Reporting Regulation (CROMERR) Security Question’ screen randomly selects and displays one of the five CROMERR eSIG-PIN questions you answered on the ‘20-5-1 Question’ screen during CDX registration.

eSIG-PIN Authorization Code: Provide one of the five secret answers to the questions that were pulled from the list of 20 CDX security questions.

Exhibit 9-6 shows the screen capture of the ‘Cross-Media Electronic Reporting Regulation (CROMERR) Security Question’ screen:

Exhibit 9-6 Cross-Media Electronic Reporting Regulation (CROMERR) Security Question Screen



Navigation: Answer the security question and click the ‘Next’ button to navigate to the ‘Submit to CDX’ screen.

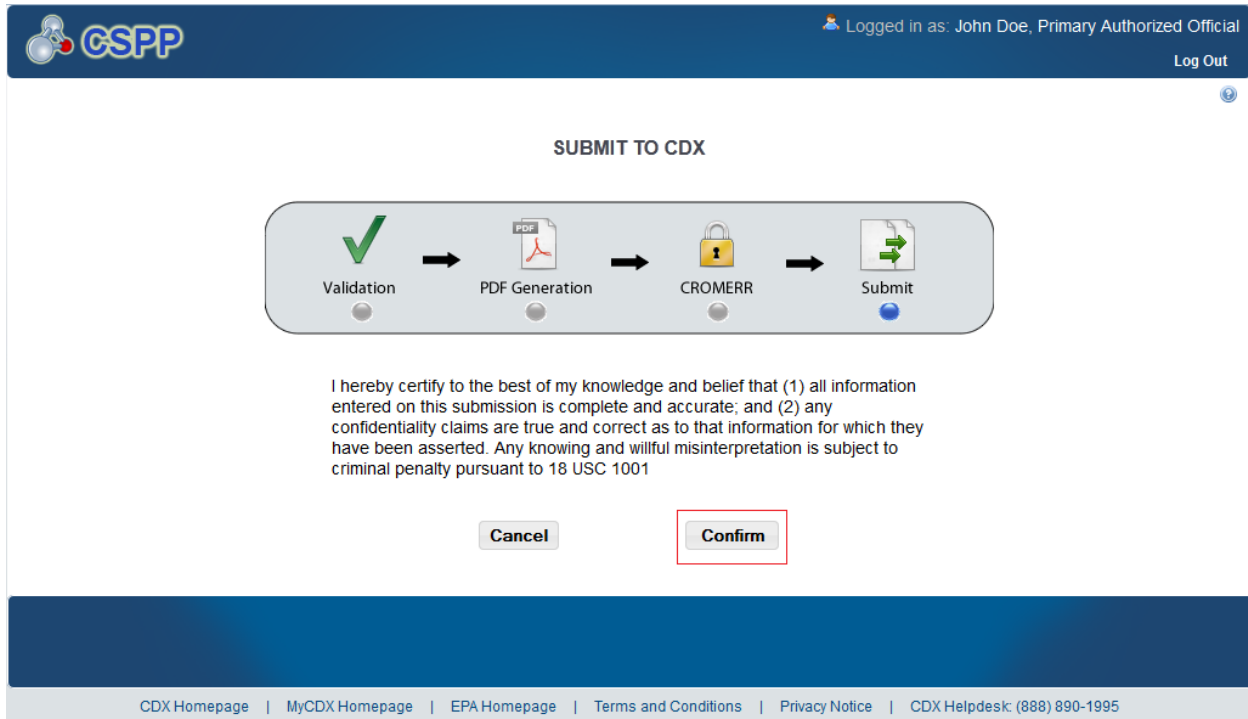
9.6 Submit to CDX Screen

You can access the ‘Submit to CDX’ screen after answering the 20-5-1 security question on the ‘Cross-Media Electronic Reporting Regulation (CROMERR) Security Question’ screen and by clicking the ‘Next’ button.

On the ‘Submit to CDX’ screen, you will see a certification message. Read the message and acknowledge by clicking the ‘Confirm’ button. Once you have clicked the ‘Confirm’ button, your submission will be submitted to CDX.

Exhibit 9-7 shows the screen capture of the ‘Submit to CDX’ screen:

Exhibit 9-7 Submit to CDX Screen



Navigation: Click the ‘Confirm’ button to navigate to the ‘Cross-Media Electronic Reporting Regulation (CROMERR) Submission’ screen.

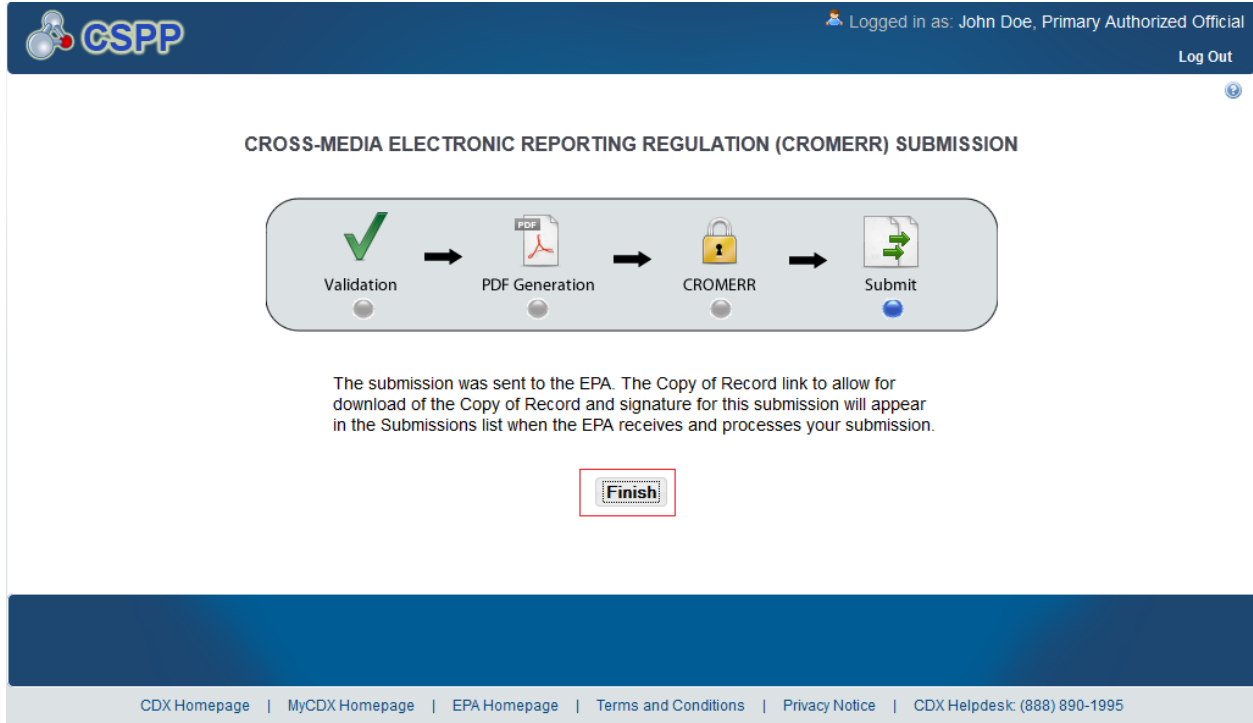
9.7 Cross-Media Electronic Reporting Regulation (CROMERR) Submission Screen

You can access the ‘Cross-Media Electronic Reporting Regulation (CROMERR) Submission’ screen by acknowledging the certification to submit to CDX by clicking the ‘Confirm’ button on the ‘Submit to CDX’ screen.

On the ‘Cross-Media Electronic Reporting Regulation (CROMERR) Submission’ screen, you will see a message indicating the submission has been successfully submitted to EPA.

Exhibit 9-8 shows the screen capture of the ‘Cross-Media Electronic Reporting Regulation (CROMERR) Submission’ screen:

Exhibit 9-8 Cross-Media Electronic Reporting Regulation (CROMERR) Submission Screen



Navigation: Click the ‘Finish’ button to navigate to the Section 4 ‘Home’ screen. Then click the ‘Submissions’ link to find your locked submission. The ‘Section 4 Test Rules, ECAs, MOUs, and 40 CFR 766’ submissions screen will display the submission with a ‘Lock’ icon () under the ‘Action’ column and will display a status of ‘Submitted.’

10 Download Copy of Record

To download a copy of record, navigate to the ‘Section 4 Test Rules, ECAs, MOUs, and 40 CFR 766’ submissions screen and locate the submission type for which you have successfully completed a submission and which has a status of ‘Submitted.’ Note that if you have just submitted to CDX the ‘Copy of record’ icon may take 15-20 minutes to display. Also, you must refresh the ‘Section 4 Test Rules, ECAs, MOUs, and 40 CFR 766’ submissions screen by clicking your browser’s refresh button for the ‘Copy of Record’ icon to display.

Click the ‘Copy of Record’ icon from the row located in the ‘Copy of Record’ column for the submitted submission. You will be taken to the ‘Enter Passphrase’ screen where you will be required to enter the passphrase originally associated with the submission. Once you enter the correct passphrase and click the ‘Next’ button, you will be taken to the ‘Cross-Media Electronic Reporting Regulation (CROMERR) Login’ screen. This screen requires you to enter the password you use to login to CDX. Once the correct CDX password is entered and you click the ‘Next’ button from the ‘Cross-Media Electronic Reporting Regulation (CROMERR) Login’ screen, you will be taken to the ‘Cross-Media Electronic Reporting Regulation (CROMERR) Security Question’ screen. The ‘Cross-Media Electronic Reporting Regulation (CROMERR) Security Question’ screen requires you to provide the answer to a randomly selected 20-5-1 security question. Once you provide the correct answer to the randomly selected 20-5-1 security question and click the ‘Next’ button, you will be taken to the ‘Download Copy of Record’ screen.

The ‘Download Copy of Record’ screen displays the ‘Download Copy of Record’ and ‘Download Attachments’ tables. The ‘Download Copy of Record’ table displays the ‘File Name’ and ‘Actions’ column. The ‘Download Attachments’ table displays the ‘File Name,’ ‘File Type,’ ‘File Size,’ and ‘Actions’ columns in a table format for all documentation that was attached within your submission. You can click the ‘Download’ icon to download a PDF version of the submission or a XML version of the submission. When you click the ‘Copy of Record’ icon, the system will allow you to view or download the individual files that were uploaded during the submission process.

Exhibit 10-1 shows the screen capture of the ‘Download Copy of Record’ screen:

Exhibit 10-1 Download Copy of Record Screen

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
Submit blank form.docx	Results Original Document	12704	↓

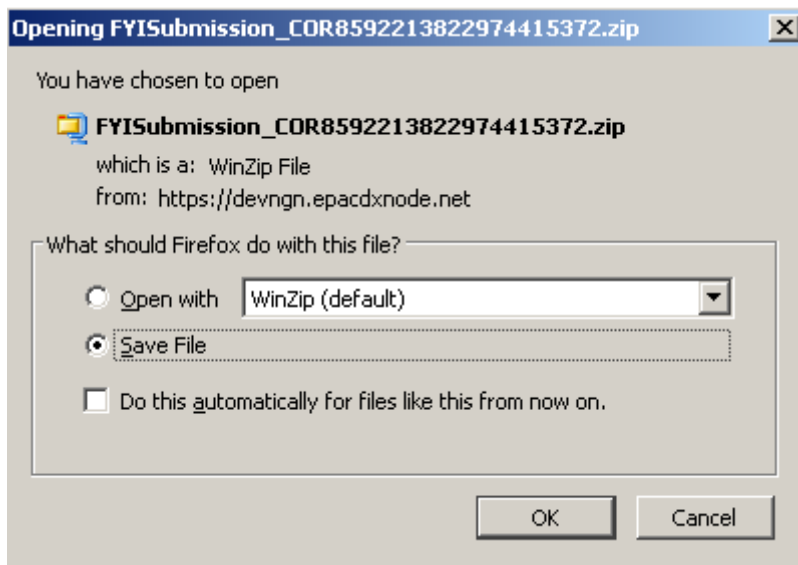
[Home](#)

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Once you click the ‘Copy of Record’ icon a window that allows you to select the appropriate option to open the CoR is generated. You can choose the ‘Open with’ or ‘Save File’ option to either open or save the CoR. If you chose the ‘Open With’ radio button option and open the CoR with WinZip(default), a ‘WinZip’ window is generated where you are able to double click the compressed file or XML file to view the CoR.

Exhibit 10-2 shows a screen capture of the download prompt pop-up window:

Exhibit 10-2 Download Prompt Pop-Up Window



If you click the 'Copy of Record' icon you will download the zip file of the submission that contains:

- A PDF version of the submission
- An XML version of the submission

11 Create an Amendment

To amend a submission, navigate to the ‘Section 4 Test Rules, ECAs, MOUs, and 40 CFR 766’ submissions screen and locate the file type that you previously submitted with a status of ‘Submitted.’ In addition to displaying a status of ‘Submitted’ under the ‘Status’ column, the



‘Lock’ icon () will be displayed under the ‘Actions’ column. Click the ‘Lock’ icon () under the ‘Actions’ column to begin the amendment process.

Exhibit 11-1 shows the screen capture of the ‘Section 4 Test Rules, ECAs, MOUs, and 40 CFR 766’ Submissions - Amendment screen:

Exhibit 11-1 Section 4 Test Rules, ECAs, MOUs, and 40 CFR 766 Submissions – Amendment

CSPP

Logged in as: John Doe, Primary Authorized Official

Home Submissions User Management Resources Log Out

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 (✖).

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

Submission Alias	CFR/FRN	Status	Modify Date	Submission Date	Copy of Record	Action
CFR766-20130717-10:24:00 EDT	40 CFR 766 Dibenzodioxins / Dibenzofurans	In Progress	07/17/2013			✖
ECA-20130718-11:23:20 EDT	70 FR 39630	In Progress	07/18/2013			✖
MOU-20130716-17:23:07 EDT	MOUForm	Submitted	07/17/2013	07/17/2013	↓	🔒
MOU-20130717-13:01:18 EDT	MOUForm	In Progress	07/18/2013			✖
MOU-20130717-16:40:20 EDT	MOUForm	In Progress	07/18/2013			✖
TestRules-20130718-11:04:56 EDT		In Progress	07/18/2013			✖

Export options: CSV | Excel | XML | PDF

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

Submission Type:

Start New Submission

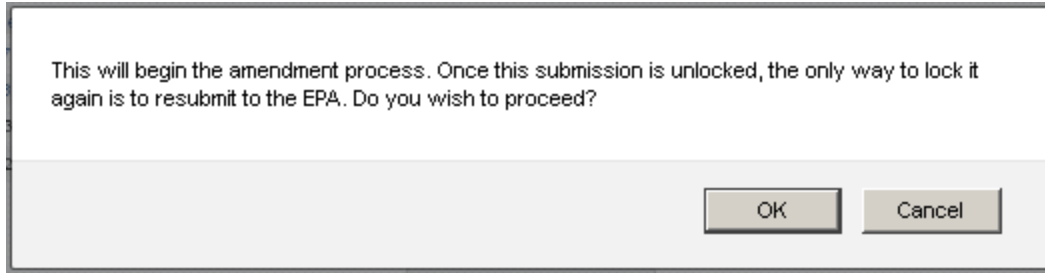
CDX Homepage | MyCDX Homepage | EPA Homepage | Terms and Conditions | Privacy Notice | CDX Helpdes: (888) 890-1995

Navigation: After clicking the ‘Lock’ icon (🔒), you will receive a prompt to confirm the amendment process. Once your submission is unlocked, you will have the ability to update the information within your submission. If you do not want to continue with the amendment, click the ‘Cancel’ button to cancel the amendment process and return to the ‘Section 4 Test Rules,


ECAs, MOUs, and 40 CFR 766' submissions screen. Click the 'OK' button to close the prompt and continue with the amendment process.

Exhibit 11-2 shows the screen capture of the 'Unlock Prompt' message:

Exhibit 11-2 Unlock Prompt Message



After you click the 'OK' button, you will be taken to the 'Enter Passphrase' screen. You must enter the passphrase originally associated with the submission to start the amendment process. After you enter the passphrase and click the 'Next' button you will be taken to the 'Submission Information Type' screen. Refer to **Section 3.4.3** for further instructions on the 'Enter Passphrase' screen.

After a submission has been unlocked, the 'Section 4 Test Rules, ECAs, MOUs, and 40 CFR 766' submissions screen will display a status of 'In Progress' under the 'Status' column, and the 'Unlocked' icon () will display under the 'Action' column; however, you will not see these changes until you navigate back to the 'Section 4 Test Rules, ECAs, MOUs, and 40 CFR 766' submissions screen.

12 Resources

You can access the ‘Resources’ screen by clicking the ‘Resources’ tab located at the top of the screen at any time during the submission process. You can also access the ‘Resources’ screen by clicking the ‘Resources’ link from the Section 4 ‘Home’ screen. The ‘Resources’ screen provides you with links to the Section 4 user guide as well as additional helpful links. Click the ‘User Guide’ link to access the Section 4 user guide document.

Toxic Substance Control Act (TSCA) Section 4 Submission: If you click this link, a new internet browser window will generate displaying the TSCA Section 4 Test Rules page.

CDX Home: If you click this link, a new internet browser window will generate displaying the CDX Homepage.

TSCA Chemical Substances Inventory: If you click this link, a new internet browser window will generate displaying the TSCA Chemical Substances Inventory screen.

Exhibit 12-1 shows the screen capture of the ‘Resources’ screen:

Exhibit 12-1 Resources Screen

RESOURCES
Section 4 Submission User Guide

This guide describes each screen of the Section 4 Submission application software and provides information on how to use the system to complete a Section 4 Submission. The guide also contains instructions for registering with CDX, submitting a Section 4 Submission, and amending a previous submission. You can download and print the guide for quick reference.

[User Guide](#)

Helpful Web Links

Click the link provided below to access additional information related to Section 4 Submission.
[Toxic Substance Control Act \(TSCA\) Section 4 Submissions](#)

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

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

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

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Appendix A. Definitions, Acronyms, and Abbreviations

The following table displays all the definitions, acronyms, and abbreviations used in this document and their full name.

Definitions, Acronyms, and Abbreviations

Acronym	Full Name
AO	Authorized Official
CA	Chemical Abstract
CASRN	Chemical Abstract Service Registry Number
CBI	Confidential Business Information
CDX	Central Data Exchange
CFR	Code of Federal Regulations
CoR	Copy of Record
CROMERR	Cross-Media Electronic Reporting Regulation Security System
CSPP	Submissions for Chemical Safety and Pesticide Programs
ECA	Enforceable Consent Agreements
EPA	Environmental Protection Agency
EST	Eastern Standard Time
FRN	Federal Register Notice
MOU	Memorandum of Understanding
OPPT	Office of Pollution Prevention and Toxics
TSCA	Toxic Substances Control Act

Appendix B. Submission Types, Effects, and Endpoints

The following appendix displays all possible options within each drop-down menu on the 'Document Management' pop-up window.

First Dropdown:

- Title: Submission Type
- Contains the following values
 - Initial Submission
 - Follow-Up Submission
 - Final Submission

Second Dropdown

- Title: Summary/Abstract Submission
- Contains the following values
 - Yes
 - No

Third Dropdown

- Title: Effects
- Contains the following values
 - Health Effect
 - Ecotoxicity
 - Environmental Fate
 - Physical-Chemical Properties

Fourth Dropdown

- Title: Endpoints
- Contains the following values
 - For Health Effect (shows when Health Effect is selected from third dropdown)
 - Basic Toxic kinetics
 - Dermal Absorption
 - Acute Toxicity: oral
 - Acute Toxicity: inhalation
 - Acute Toxicity: dermal
 - Acute Toxicity: other routes
 - Skin irritation/corrosion
 - Eye irritation
 - Skin sensitization
 - Respiratory sensitization
 - Repeated Dose toxicity: oral
 - Repeated Dose toxicity: inhalation

- Repeated Dose Toxicity: dermal
- Repeated Dose Toxicity: other routes
- Genetic Toxicity in vitro
- Genetic Toxicity in vivo
- Carcinogenicity
- Toxicity to Reproduction
- Developmental Toxicity/teratogenicity
- Toxicity to reproduction: other studies
- Neurotoxicity
- Immunotoxicity
- Specific investigations: other studies
- Health surveillance data
- Epidemiological data
- Direct observations: clinical cases, poisoning incidents and other
- Sensitization data (humans)
- Exposure related observations in humans: other data
- Toxic effects on livestock and pets
- Additional toxicological information
- Other
- For Ecotoxicity (shows when Ecotoxicity is selected from third dropdown)
 - Short-term toxicity to fish
 - Long-term toxicity to fish
 - Short-term toxicity to aquatic invertebrates
 - Long-term toxicity to aquatic invertebrates
 - Toxicity to aquatic algae and cyanobacteria
 - Toxicity to aquatic plants other than algae
 - Toxicity to microorganisms
 - Toxicity to other aquatic organisms
 - Sediment toxicity
 - Toxicity to soil macroorganisms except arthropods
 - Toxicity to terrestrial arthropods
 - Toxicity to terrestrial plants
 - Toxicity to soil microorganisms
 - Toxicity to birds
 - Toxicity to other above-ground organisms
 - Biological effects monitoring
 - Biotransformation and kinetics
 - Additional ecotoxicological information
 - Other

-
- For Environmental Fate (shows when Environmental Fate is selected from third dropdown)
 - Photo transformation in air
 - Hydrolysis
 - Photo transformation in water
 - Photo transformation in soil
 - Biodegradation in water: screening test
 - Biodegradation in water and sediment: simulation tests
 - Biodegradation in soil
 - Mode of degradation in actual use
 - Bioaccumulation: aquatic/sediment
 - Bioaccumulation: terrestrial
 - Adsorption/desorption
 - Henry's Law constant
 - Distribution Modeling
 - Other Distribution Data
 - Monitoring Data
 - Field Studies
 - Additional Information on Environmental Fate and Behavior
 - Other
 - For Physical-Chemical Properties (shows when Physical-Chemical Properties is selected from third dropdown)
 - Appearance/Physical
 - Melting Point
 - Boiling Point
 - Density
 - Particle size distribution
 - Vapor Pressure
 - Partition Coefficient
 - Water Solubility
 - Solubility in Organic Solvent
 - Surface Tension
 - Flash Point
 - Auto Flammability
 - Flammability
 - Explosiveness
 - Oxidizing Properties
 - Oxidation Reduction Potential
 - Stability in organic solvents and identity of relevant degradation products
 - Storage stability and reactivity towards container material

- Stability: thermal, sunlight, metals
- pH
- Dissociation constant
- Viscosity
- Additional physic-chemical information
- Other