

Dispenser's Implementation Guide

ASAP 4.2

Florida Department of Health Prescription Drug Monitoring Program



July 2015

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1 Program Overview

Purpose

The Electronic-Florida Online Reporting of Controlled Substances Evaluation program (E-FORCSE) is Florida's Prescription Drug Monitoring Program (PDMP). The PDMP was created by the 2009 Florida Legislature in an initiative to encourage safer prescribing of controlled substances and to reduce drug abuse and diversion within the State of Florida. The purpose of the PDMP is to provide the information that will be collected in the database to healthcare practitioners to guide their decisions in prescribing and dispensing these highly-abused prescription drugs.

E-FORCSE has selected Health Information Designs (HID) to develop a database that will collect and store prescribing and dispensing data for controlled substances in Schedules II, III, and IV. HID's RxSentry® is a web-based program that facilitates the collection, analysis, and reporting of information on the prescribing, dispensing, and use of controlled substance prescription drugs. RxSentry leads the industry in flexibility, functionality, and ease of use.

Section 893.055, Florida Statutes (F.S.) requires healthcare practitioners to report to the PDMP each time a controlled substance is dispensed to an individual. This information is to be reported through the electronic system as soon as possible but not more than seven (7) days after dispensing. This reporting timeframe ensures that health care practitioners have the most up-to-date information available.

E-FORCSE will comply with the Health Insurance Portability and Accountability Act (HIPAA) as it pertains to protected health information (PHI), electronic protected health information (EPHI), and all other relevant state and federal privacy and security laws and regulations. The information collected in the system will be used by the PDMP to encourage safer prescribing of controlled substances and reduce drug abuse and diversion within the State of Florida.

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2 Document Overview

Purpose and Content

The RxSentry Dispenser's Implementation Guide serves as a step-by-step implementation and training guide for dispensers in the State of Florida. A "dispenser" is defined as a pharmacy, dispensing pharmacist, or dispensing healthcare practitioner. It includes such topics as:

- Reporting requirements for dispensers in the State of Florida
- Data file submission guidelines and methods
- Creating your upload account
- Creating a data file
- Uploading or reporting your data
- Understanding upload error codes and definitions

This guide has been customized to target the specific training needs of Florida dispensers. It is intended for use by all dispensers in the State of Florida required to report dispensing of controlled substances.

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3 Data Collection and Tracking

Data Collection Requirements

Each time a controlled substance is dispensed to an individual, the controlled substance shall be reported to the E-FORCSE program using a format approved by the Florida Department of Health (DOH) as soon thereafter as possible, but not more than seven (7) days after the date the controlled substance was dispensed. All dispensers of controlled substances must meet the reporting requirements set forth by section 893.055, F.S., in a secure methodology and format. Such approved formats may include, but are not limited to, submission via the Internet, on a disc, or by use of regular mail.

Reporting Requirements

Any healthcare practitioner who has dispensed a controlled substance, as defined in section 893.03, F.S. (i.e., OxyContin®, Percocet®, Vicodin®, Klonopin®, Xanax®, and Valium®) will be required to report to the database. This includes pharmacies licensed under chapter 465, F.S., including mail order and Internet pharmacies; and healthcare practitioners licensed under chapter 458, 459, 461, 462, 465, or 466, F.S.

For detailed information for each of the fields required by the State of Florida and the fields required by the American Society for Automation in Pharmacy (ASAP), please see Appendix A: ASAP 4.2 Specifications.

Exemptions

A healthcare practitioner will not be required to report to E-FORCSE when he/she:

- Administers a controlled substance directly to a patient if the amount is adequate to treat the patient during that particular treatment session;
- Administers a controlled substance to a patient or resident receiving care as a patient at a
 hospital, nursing home, ambulatory surgical center, hospice, or intermediate care facility for
 the developmentally disabled;
- Administers or dispenses a controlled substance in the healthcare system of the Florida Department of Corrections;
- Administers a controlled substance in the Emergency Room of a licensed hospital;
- Administers or dispenses a controlled substance to a patient under the age of 16; or
- Dispenses a one-time, 72-hour re-supply of controlled substances.

Required Prescription Information

The following information must be reported for each controlled substance dispensed to a patient:

Field Name	Field ID	
Pharmacy Header		
DEA Number	PHA03	
Patient Information		
Last Name	PAT07	
First Name	PAT08	
Address Information – 1	PAT12	
City Address	PAT14	
State Address Note: Populate with "99" if patient address is outside the U.S.	PAT15	
ZIP Code Address	PAT16	
Date of Birth	PAT18	
Gender Code	PAT19	
Dispensing Record		
Prescription Number	DSP02	
Date Written	DSP03	
Date Filled	DSP05	
Refill Number	DSP06	
Product ID (NDC)	DSP08	
Quantity Dispensed	DSP09	
Classification Code for Payment Type	DSP16	
Prescriber Information		
National Provider Identifier-NPI (if available)	PRE01	
DEA Number	PRE02	
Prescriber State License Number (if available)	PRE04	
Last Name	PRE05	
First Name	PRE06	

The Data Submission chapter provides all the instructions necessary to submit the required information.

Reporting Noncompliance

Any dispenser who willfully and knowingly fails to report the dispensing of a controlled substance, as required by section 893.055, F.S., commits a misdemeanor of the first degree, punishable as provided in sections 775.082 or 775.083, F.S.

Zero Reports

If a dispenser usually dispenses controlled substances in Florida but has no dispensing transactions to report for the preceding seven (7) day period, the dispenser must report this information to E-FORCSE by filing a zero report, as described in the Reporting Zero Dispensing topic in this guide.

Reporting Waivers

No Reporting Waiver

If a dispenser is permitted or licensed in the State of Florida, but does not dispense any controlled substances directly to Florida residents, they are not required to report to E-FORCSE. However, the dispenser must notify DOH in writing by completing a waiver form provided by DOH stating that it does not dispense controlled substances in the state.

Electronic Reporting Waiver

DOH may grant a dispenser a waiver of the electronic submission requirement for good cause as determined by the DOH. "Good cause" includes financial hardship and lack of an automated recordkeeping system. The dispenser must notify DOH in writing by completing an electronic reporting waiver form provided by DOH. DOH will work with the dispenser to determine the format, method, and frequency of the alternative non-electronic submissions.

Florida Department of Health Dispenser's Implementation Guide		Data Collection and Tracking
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4 Data Submission

About This Chapter

This chapter provides information and instructions for submitting data to E-FORCSE.

Timeline and Requirements

Registration for dispensers began August 1, 2011, and reporting began September 1, 2011. You may create an account and begin reporting upon receipt of this guide. Instructions for setting up an account are provided in the Creating Your Account topic in this chapter.

- Dispensers are required to report their data within seven (7) days of dispensing a controlled substance. However, dispensers are encouraged to report more frequently if they would like.
- Retroactive data from December 1, 2010 to August 31, 2011 was reported to the E-FORCSE program until November 30, 2011.

Emergency Suspension

In the event that a state of emergency is declared in the State of Florida, and a dispenser is not allowed or is unable to report to E-FORSCE because of the declared state of emergency, a reporting suspension waiver will be granted for the seven (7) day reporting period. Once the state of emergency has been lifted, the dispenser must report the backlog of data as soon as possible in order to bring reporting current.

Upload Specifications

Files must be in the ASAP 4.2 format, as defined in Appendix A: ASAP 4.2 Specifications. Files for upload should be named in a unique fashion, with a prefix constructed from the date (YYYYMMDD) and a suffix of .dat. An example file name would be 20110801.dat. All of your upload files will be kept separate from the files of others.

Reports for multiple dispensers can be in the same upload file in any order.

Controlled substance prescription information must be reported within seven (7) days of dispensing a controlled substance, unless a waiver has been obtained from DOH.

Creating Your Account

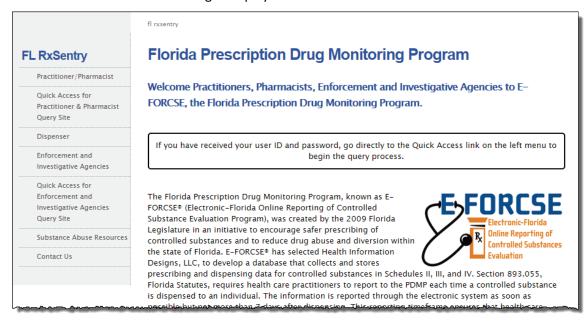
Prior to submitting data, you must create an upload account. If you have already created your account, proceed to the appropriate section of this document that outlines the steps you must follow to upload your data.

Note: Multiple dispensers' reports can be uploaded in the same file. For example, chain pharmacies may send in one file containing controlled substance dispensing information for all of their pharmacies licensed in the State of Florida. Therefore, chains with multiple stores only have to set up one account to upload a file.

Perform the following steps to create an account:

1. Open an Internet browser window and type the following URL in the address bar: http://www.hidesigns.com/flpdmp.

A window similar to the following is displayed:

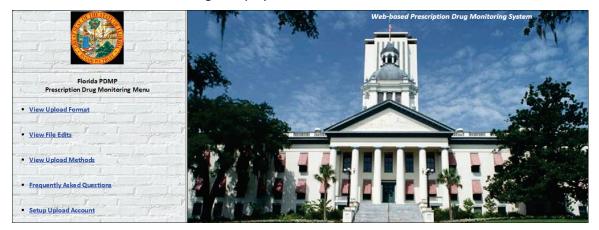


2. Click Dispenser, and then click RxSentry Dispenser's Upload Site.

A login window is displayed.

- 3. Type *newacct* in the **User Name** field.
- 4. Type welcome in the Password field.
- 5. Click OK.

A window similar to the following is displayed:



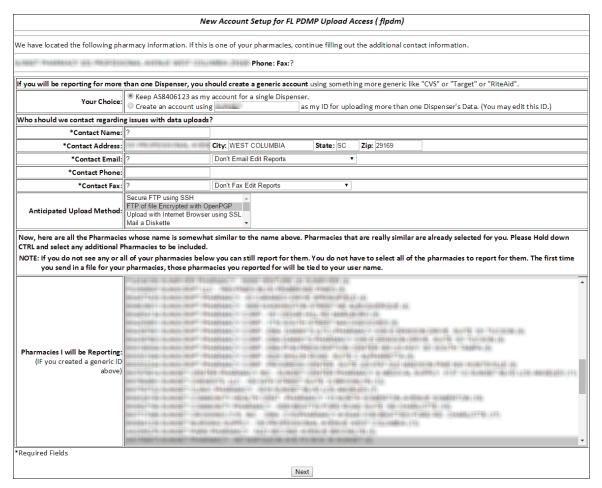
6. Click Setup Upload Account.

A window similar to the following is displayed:



- 7. Enter your DEA number in the **Physician or Pharmacy DEA number** field.
- 8. Type your ZIP code in the **ZIP Code** field, and then click **Next**.

The **New Account Setup for Upload Access** window is displayed as shown on the following page.

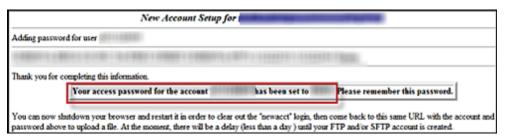


9. Complete all required fields (indicated by an asterisk) on the **New Account Setup for Upload Access** window, using the information in the following table as a guideline:

Field	Description/Usage	
Account selection	 Choose Keep <account number=""> as my account for a single Pharmacy if you wish to use the suggested account name.</account> Choose Create an account using <suggested account="" name=""> as my ID for uploading more than one Pharmacy's Data if you wish to enter an account name of your choosing. If this option is selected, type the desired account name in this field.</suggested> 	
Contact Information Note: Information in this section is used for contact purposes in the event a problem occurs with a data upload.		
Contact Name	(Required) Type the first and last name of the contact person.	
Contact Address	(Required) Type the contact's street address, city, state, and ZIP code in the appropriate fields.	
Contact Email	(Required) Type the contact's e-mail address. Click the down arrow in the field to the right of the Contact Email field to select Email Edit Reports for All Uploads.	

Field	Description/Usage
Contact Phone	(Required) Type the contact's phone number, using the format 999-999-9999.
Contact Fax	(Required) Type the contact's fax number, using the format 999-999-9999. Click the down arrow in the field to the right of the Contact Fax field and select Fax Edit Reports for All Uploads
Anticipated Upload Method	Select the method of data upload you plan to use to report your data: Secure FTP over SSH Encrypted File with OpenPGP Via FTP SSL Website Physical Media (Tape, Diskette, CD, DVD) Universal Claim Form (UCF) Submission Note: You must have received a waiver from DOH to use this method.
Dispensers I will be reporting	A list of all dispensers (pharmacy, dispensing pharmacist, or dispensing health care practitioner) with names similar to the account name you entered above is displayed in this field. To select additional dispensers for which you will be reporting, press the [CTRL] key and then click the name of each dispenser you wish to select. The dispensers you select will be tied to your user name.

10. After completing all required fields, click **Next**. A window similar to the following is displayed:



A randomly-assigned password for the FTP, SFTP, and SSL website upload processes is provided to you.

Software vendors setting up multiple accounts may choose from the following options:

Create each account separately by using the method listed above. After you finish one
dispenser's account, click Setup Upload Account on the home page, and repeat the
process;

Or

• Create multiple accounts using one dispenser's DEA number and ZIP code. If you choose this method, select **Set up user name as a group**.

Note: Data error reports are submitted to the e-mail address(es) supplied for the account(s).

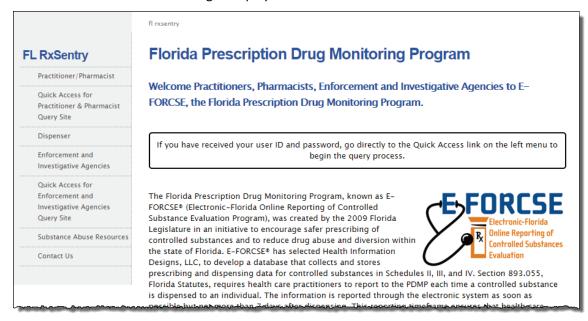
Modifying Your Upload Account

Use this function if you need to modify the information supplied when you originally created your account.

Perform the following steps to update your information:

1. Open an Internet browser window and type the following URL in the address bar: http://www.hidesigns.com/flpdmp.

A window similar to the following is displayed:



2. Click **Dispenser**, and then click **RxSentry Dispenser's Upload Site**.

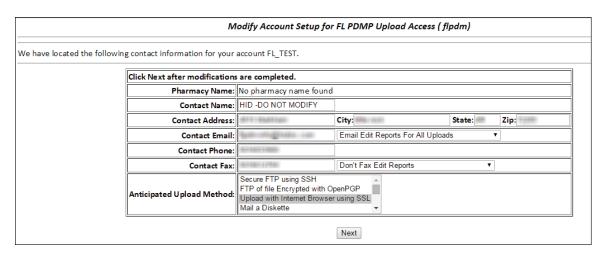
A login window is displayed.

- 3. Type your user name in the **User Name** field.
- 4. Type your password in the **Password** field.
- 5. Click OK.

The RxSentry home page is displayed.

6. Click Modify Upload Account.

The **Modify Account Setup** window is displayed as shown on the following page.



- 7. Update the information as necessary, using the field descriptions provided in the Creating Your Account topic as a guideline.
- 8. Click Next.

A message displays indicating that your account information was successfully updated.

Reporting Zero Dispensing

If you have no dispensing transactions to report for the preceding seven-day period, you must report this information to E-FORCSE.

You may report zero dispensing by using the functionality provided within RxSentry via the Report Zero Activity menu item, or by creating and uploading a zero report data file. The steps you must perform for each method are provided in the following sections.

Report Zero Activity – RxSentry

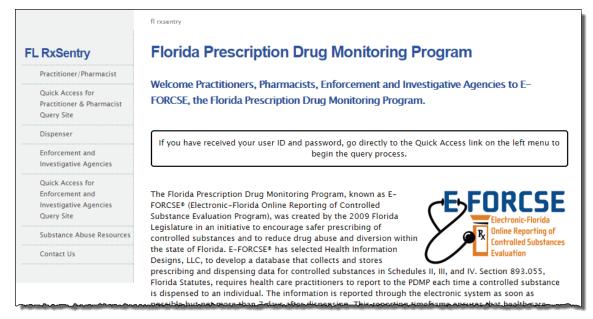
The information in the following topics explains the processes single dispensers and dispensers reporting for a group of pharmacies should use to report zero activity using RxSentry's Report Zero Activity menu item.

Single Dispensers

If you are a single dispenser, perform the following steps to report zero activity using RxSentry's Report Zero Activity menu item:

- 1. If you do not have an account, perform the steps in Creating Your Account.
- 2. Open an Internet browser window and type the following URL in the address bar: http://www.hidesigns.com/flpdmp.

A window similar to the following is displayed:



3. Click **Dispenser**, and then click **RxSentry Dispenser's Upload Site**.

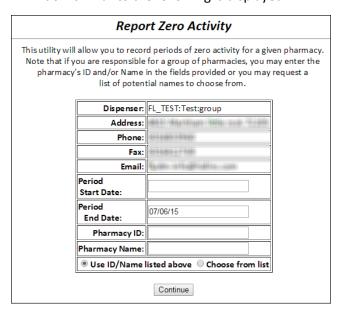
A login window is displayed.

- 4. Type your user name in the **User Name** field.
- 5. Type your password in the **Password** field.
- 6. Click OK.

The RxSentry home page is displayed.

7. Click Report Zero Activity.

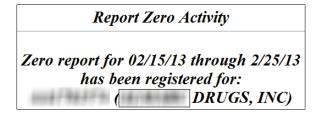
A window similar to the following is displayed:



8. Type the start date for this report in the **Period Start Date** field, using the *mm/dd/yy* format.

Notes:

- The Period End Date field is populated with the current date. You may adjust this date, if necessary.
- All other dispenser information is populated with the information provided when you created your account.
- 9. Click **Continue**. A message similar to the following is displayed, indicating your zero report has been successfully submitted:



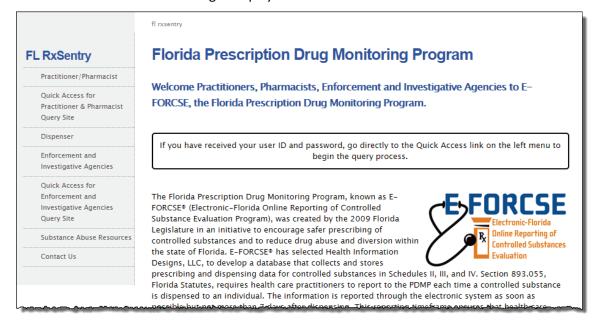
Group Pharmacies

If you are responsible for reporting for a group of pharmacies, perform the following steps to report zero activity using RxSentry.

Note: You are required to repeat this process for every pharmacy for which you are responsible for reporting.

- 1. If you do not have an account, perform the steps in Creating Your Account.
- 2. Open an Internet browser window and type the following URL in the address bar: http://www.hidesigns.com/flpdmp.

A window similar to the following is displayed:



3. Click **Dispenser**, and then click **RxSentry Dispenser's Upload Site**.

A login window is displayed.

- 4. Type your user name in the **User Name** field.
- 5. Type your password in the **Password** field.
- 6. Click OK.

The RxSentry home page is displayed.

7. Click Report Zero Activity.

A window similar to the following is displayed:



8. Type the start date for this report in the **Period Start Date** field, using the *mm/dd/yy* format.

Notes:

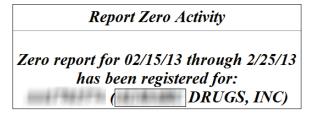
- The Period End Date field is populated with the current date. You may adjust this date, if necessary.
- All other dispenser information is populated with the information provided when you created your account.
- 9. Select the **Use ID/Name listed above** option to manually enter the pharmacy ID whose information you are reporting. If you choose to enter the pharmacy ID manually, type the pharmacy ID in the **Pharmacy ID/Name** field.

Or

Select the **Choose from list** option to select the pharmacy ID whose information you are reporting from a list of pharmacies with a name similar to your pharmacy.

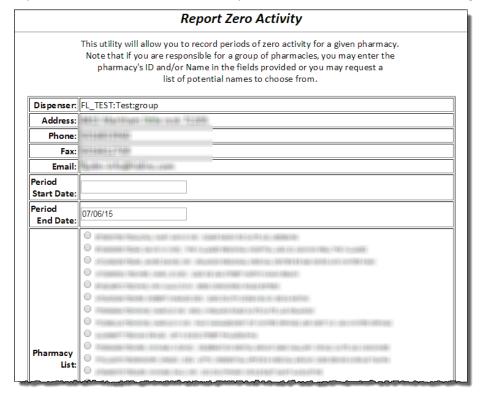
10. Click Continue.

If you selected the **Use ID/Name listed above** option, a message similar to the following is displayed:



Or

If you selected the **Choose from list** option, a window similar to the following is displayed:



- 11. Click the radio button next to the correct pharmacy ID.
- 12. Click Continue.

A window similar to the following is displayed:

Report Zero Activity		
Zero report for 02/15/13 through 2/25/13		
has been registered for:		
(DRUGS, INC)		

Report Zero Activity – File Upload

- 1. If you have not created an account, perform the steps in Creating Your Account.
- 2. Prepare the zero report data file for submission, using the specifications described in Appendix B: Zero Report Specifications.

Important Notes:

- The file name should be constructed using the date of submission to HID as the file name and should have a .dat extension. For example, name the file 20140801.dat if you submit it on August 1, 2014.
- Do not include spaces in the file name.
- If you submit more than one file within the same day, you must uniquely name each file so the system does not overwrite existing uploaded files. For example, if uploading three files within the same day, you could use the following file names: 20140801a.dat, 20140801b.dat, and 20140801c.dat.
- The system will accept zipped files and you should name them using the date of submission to HID. For example, name the file 20140801.zip if you submit it on August 1, 2014.
- Before transmitting your file, rename it to include the suffix .up (e.g., 20140801.dat.up). This will ensure that we do not try to load the file while you are transmitting it. Once transmission is complete, rename the file back to the original name (e.g., 20140801.dat).
- 3. Upload the file using the steps provided in one of the following data delivery topics:
 - Secure FTP over SSH
 - Encrypted File with OpenPGP via FTP
 - SSL Website

HID tracks the use of the web-based tool, date stamps incoming files, and notifies you of a successful file transmission. After the file is reviewed for accuracy, you are notified of the status of the submitted file.

5 Data Delivery Methods

About This Chapter

This chapter provides information about data delivery methods you can use to upload your controlled substance reporting data file(s).

For quick reference, click the desired hyperlink in the following table to view the step-by-step instructions for your chosen data delivery method:

Delivery Method	Page
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Encrypted File with OpenPGP via FTP	22
SSL Website	23
Physical Media (Tape, Diskette, CD, DVD)	24
Universal Claim Form (UCF) Submission	
Online UCF Submission	25
Reporting Requirements for UCF Submissions	25
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Secure FTP over SSH

There are many free software products that support secure FTP. Neither DOH nor HID is in a position to direct or support your installation of operating system software for secure FTP; however, we have information that WinSCP (http://winscp.net) has been used successfully by other pharmacies.

- 1. If an account has not yet been created, perform the steps in Creating Your Account.
- 2. Prepare the data file for submission, using the ASAP specifications described in Appendix A: ASAP 4.2 Specifications.

Important Notes:

- The file name should be constructed using the date of submission to E-FORCSE as the file name, and should have a .dat extension. For example, name the file 20110801.dat if it is submitted on August 1, 2011.
- Do not include spaces in the file name.
- If more than one file is submitted within the same day, each file must be uniquely named so that existing uploaded files are not overwritten. For example, if uploading three files within the same day, the following file names could be used: 20110801a.dat, 20110801b.dat, and 20110801c.dat.
- Zipped files can be accepted and should be named using the date of submission to E-FORCSE. For example, name the file 20110801.zip if it is submitted on August 1, 2011.

- **Before transmitting your file**, rename it to include the suffix .up (e.g., 20110801.dat.up). This will ensure that we do not try to load the file while you are transmitting it. Once transmission is complete, rename the file back to the original name (e.g., 20110801.dat).
- 3. SFTP the file to sftp://flpdmp-reporting.hidinc.com.
- 4. When prompted, type *flpdm* (lower case) in front of your DEA number (or Generic ID) as your user ID, and enter the password supplied when you created your account.
- 5. Place the file in the new directory.
- 6. If desired, view the results of the transfer/upload in your user directory. The file name is **YYYYMMDD.rpt**.
- 7. Log off when the file transfer/upload is complete.

HID tracks the use of the web-based tool, and incoming files are date stamped. You are notified of a successful file transmission. After the file is reviewed for accuracy, you are notified of the status of the submitted file.

Encrypted File with OpenPGP via FTP

There are many free software products that support file encryption using the PGP standard. Neither DOH nor HID is in a position to direct or support your installation of PGP compatible software utilities; however, our usage indicates that software from the GnuPG Project (http://gnupg.org) should be compatible with many operating systems.

- 1. If an account has not yet been created, perform the steps in Creating Your Account.
- 2. Import the PGP public key, supplied during the account creation, into your PGP key ring.
- 3. Prepare the data file for submission, using the ASAP specifications described in Appendix A: ASAP 4.2 Specifications.

Important notes:

- The file name should be constructed using the date of submission to E-FORCSE as the file name, and should have a .pgp extension. For example, name the file 20110801.pgp if it is submitted on August 1, 2011.
- Do not include spaces in the file name.
- If more than one file is submitted within the same day, each file must be uniquely named so that existing uploaded files are not overwritten. For example, if uploading three files within the same day, the following file names could be used: 20110801a.pgp, 20110801b.pgp, and 20110801c.pgp.
- **Before transmitting your file**, rename it to include the suffix .up (e.g., 20110801.pgp.up). This will ensure that we do not try to load the file while you are transmitting it. Once transmission is complete, rename the file back to the original name (e.g., 20110801.pgp).

4. Encrypt the file with the PGP software, using the public key supplied during account creation.

Note: PGP encryption performs a single compression as it encrypts, so there is no need to zip the file.

- 5. FTP the file to ftp://flpdmp-reporting.hidinc.com.
- 6. When prompted, type *flpdm* (lower case) in front of your DEA number (or Generic ID) as your user ID, and enter the password supplied when you created your account.
- 7. Place the file in the new directory.
- 8. Once the transmission is complete, rename the file without the .up extension (e.g., 20110801.pgp).
- 9. If desired, view the results of the transfer/upload in your user directory. The file name is **YYYYMMDD.rpt**.
- 10. Log off when the file transfer/upload is complete.

HID tracks the use of the web-based tool, and incoming files are date stamped. You are notified of a successful file transmission. After the file is reviewed for accuracy, you are notified of the status of the submitted file.

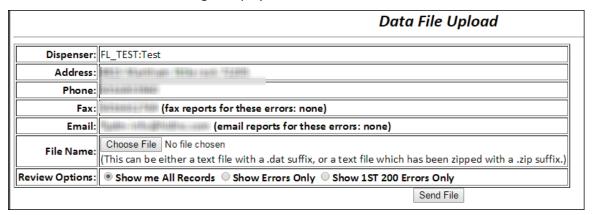
SSL Website

- 1. If an account has not yet been created, perform the steps in Creating Your Account.
- 2. Prepare the data file for submission, using the ASAP specifications described in Appendix A: ASAP 4.2 Specifications.

Important notes:

- The file name should be constructed using the date of submission to E-FORCSE as the file name, and should have a .dat extension. For example, name the file 20110801.dat if it is submitted on August 1, 2011.
- Do not include spaces or parentheses in the file name.
- If more than one file is submitted within the same day, each file must be uniquely named so that existing uploaded files are not overwritten. For example, if uploading three files within the same day, the following file names could be used: 20110801a.dat, 20110801b.dat, and 20110801c.dat.
- Zipped files can be accepted and should be named using the date of submission to E-FORCSE. For example, name the file 20110801.zip if it is submitted on August 1, 2011.
- 3. Open a web browser and enter the following URL: https://flpdmp-reporting.hidinc.com. A login window is displayed.
- 4. Enter the user ID and password supplied when the account was created, and then click **OK**. The RxSentry home page is displayed.
- 5. Click **Upload a File**.

A window similar to the following is displayed:



- 6. Click Choose File, and navigate to the location where you saved the file created in step 2.
- 7. If not previously named according to upload requirements, rename the file using the format *YYYYMMDD.dat*, for example, *20110801.dat*.
- 8. Click to select the file, and then click **Open**.
- 9. Click Send File.

HID tracks the use of the web-based tool, and incoming files are date stamped. You are notified of a successful file transmission. After the file is reviewed for accuracy, you are notified of the status of the submitted file.

Physical Media (Tape, Diskette, CD, DVD)

- 1. If an account has not yet been created, perform the steps in Creating Your Account.
- 2. Prepare the data file for submission, using the ASAP specifications described in Appendix A: ASAP 4.2 Specifications.

Important Notes:

- The file name should be constructed using the date of submission to E-FORCSE as the file name, and should have a .dat extension. For example, name the file 20110801.dat if it is submitted on August 1, 2011.
- Do not include spaces or parentheses in the file name.
- If more than one file is submitted within the same day, each file must be uniquely named so that existing uploaded files are not overwritten. For example, if uploading three files within the same day, the following file names could be used: 20110801a.dat, 20110801b.dat, and 20110801c.dat.
- Zipped files can be accepted and should be named using the date of submission to E-FORCSE. For example, name the file 20110801.zip if it is submitted on August 1, 2011.
- 3. Write the file to the preferred media (tape, diskette, CD, or DVD).
- 4. Add a label to the outside of the media that contains the following information:
 - Dispenser's DEA Number

- Date of Submission
- Contact Person
- 5. Mail the media to:

Health Information Designs, LLC Attn: FL PDMP 391 Industry Drive Auburn, AL 36832

Universal Claim Form Submission

Online UCF Submission

If you have Internet access, but are unable to submit your data in a batch upload, you may submit prescription information using RxSentry's online universal claim form (UCF).

When submitting information using the online UCF, the information provided must be complete and accurate. Only complete and accurate submissions are entered into the Florida PDMP database. Please use the information in the Notes about NDC Numbers topic below as a guideline for providing accurate NDC numbers.

The following new terms are introduced in this topic:

- Record the patient, pharmacy, and prescription information that you enter for one patient on the UCF
- Batch a single record, or group of records, that you upload using the Submit Batch function

Note: Records can be continually added to a batch—a convenient feature that allows you to enter records at your convenience and not all at one time. We recommend that you add as many records as possible to a batch before submitting it; however, you must submit and close batches in accordance with the seven (7) day reporting timeframe.

Reporting Requirements for UCF Submissions

See the Required Prescription Information topic for details regarding reporting requirements.

Notes about NDC Numbers

Use the following information when entering NDC numbers on the UCF:

- NDCs are 11 digits and use the format 99999-999-99.
- When adding an NDC, do not include the dashes, for example, 999999999999.
- NDCs are typically located on the original medication bottle on the top right corner of the label, prefaced with "NDC-" and followed by the number.

• Manufacturers often leave off a zero in the NDC. In these instances, you should add the 0 where appropriate, using the following examples as a guideline:

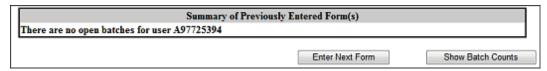
If the NDC appears this way	Enter it this way
1234-5678-90 (missing 0 in first segment)	01234568790
54321-123-98 (missing 0 in 2nd segment)	54321012398

Submitting Information Using the Online UCF

Perform the following steps to use the online UCF to submit prescription information:

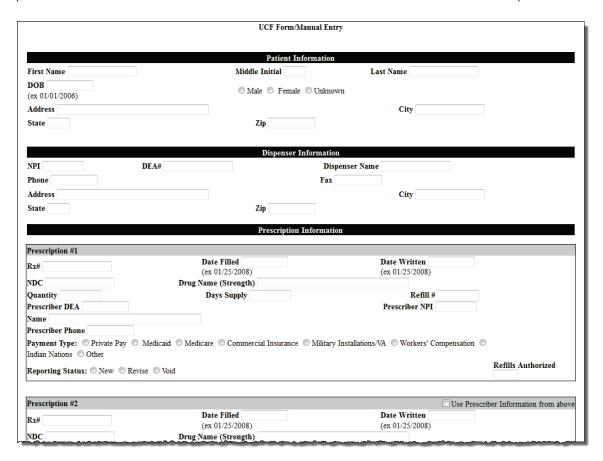
- 1. If you do not have an account, perform the steps in Creating Your Account.
- Open a web browser and enter the following URL: https://flpdmp-reporting.hidinc.com.
 A login window is displayed.
- Enter the user ID and password supplied when the account was created, and then click OK.
 The RxSentry home page is displayed.
- 4. Click **UCF Form Entry**.

A window similar to the following is displayed:



- Enter Next Form allows you to prepare one or more records for submission.
- **Show Batch Counts** displays the number of records in the batch currently being prepared for submission and the number of records that have been previously been submitted.
- 5. Click Enter Next Form.

The online UCF is displayed as shown on the following page.



The UCF contains three sections—Patient Information, Dispenser Information, and Prescription Information. Refer to the following information to complete these sections on the UCF:

- Patient Information Complete all fields in this section.
- Dispenser Information In this section, supply your DEA number in the DEA field. Once
 this information is provided, all associated pharmacy information available within the
 RxSentry database is auto-populated in the appropriate fields.
- Prescription Information Information for up to three prescriptions may be entered in this section, and all fields for each prescription must be completed.

If entering more than one prescription for the same prescriber, you may select the **Use Prescriber Information From Above** check box to auto-populate each prescription with the previously-used prescriber information.

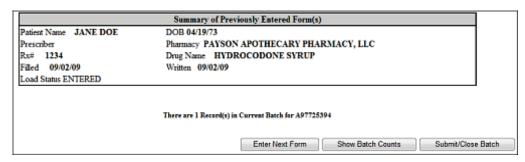
6. Once all information has been entered, click **Submit**.

Notes:

- If information is missing from any required fields on the UCF, the UCF window will display again with the required fields indicated. Click **Modify** to add the missing information, and then click **Submit**.
- If the system indicates that the DEA number or the NDC number you have provided is invalid, and you are certain you have provided the correct number, contact HID using the information supplied in Assistance and Support.

7. The UCF is displayed for your review. If all information is correct, click **Submit**. If you need to modify any information, click **Modify**.

Once you click **Submit**, a window similar to the following is displayed:



- 8. Perform one of the following functions:
 - Click Enter Next Form to add additional records to this batch.
 - Click Show Batch Counts to display the number of records in the current batch.
 - Click Submit/Close Batch to upload this batch of records.

Waiver for Dispensers without Internet Access

DOH may issue a waiver to a dispenser that is unable to submit controlled substance prescription information by electronic means. The waiver may permit the dispenser to submit prescription information by paper form, provided that all required information is submitted.

To request an electronic reporting waiver, please navigate to the Florida PDMP website, http://www.hidesigns.com/flpdmp, and locate the waiver form.

If a waiver is granted, use the paper submission method to submit prescription information. See the Paper UCF Submission topic for instructions.

Important notes:

- As with the online submission method, the information provided must be complete and accurate.
- Use the information in the Notes about NDC Numbers topic as a guideline for providing accurate NDC numbers.

Paper UCF Submission

If you have been granted a waiver to report your controlled substance prescription information on a paper UCF, you may fax the completed forms to 1-888-288-0337 or mail to:

Health Information Designs, LLC

Attn: FL PDMP P.O. Box 3210

Auburn, AL 36832-3210

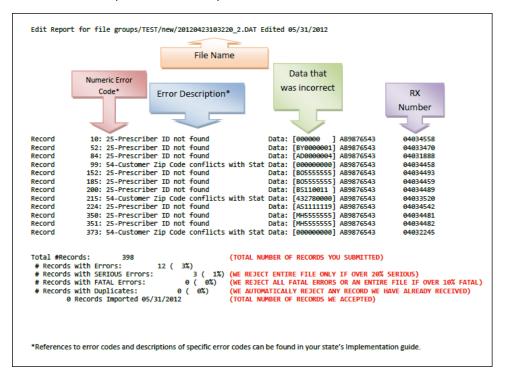
A copy of the UCF is provided in Appendix C: Universal Claim Form.

6 Upload Reports and Edit Definitions

Upload Reports

E-FORCSE provides all submitters of data with an upload report. When creating an account, you are required to submit an e-mail address or a fax number. You must also specify the method by which you wish to receive your upload report. If you FTP/SFTP the data, a report will be placed in your home directory on the FTP server.

Below is an example of an error report:



A single record may be rejected or, if a certain percentage of records are rejected in an individual file, the entire file may be rejected. We track three types of errors:

- Minor Incorrect data in non-vital field
- Serious Record can be loaded with missing or inappropriate data
- Fatal Record cannot be loaded

A single record will be rejected if it contains a fatal error.

An entire batch will be rejected if:

- ALL records have Fatal or Serious errors
- More than 10% of the records have Fatal errors
- More than 20% of the records have Serious errors

Dispensers are required to correct fatal errors and resubmit the records within seven (7) days of the initial record submission.

View Upload Reports

This function provides uploaders access to upload reports that were previously delivered via email or fax following a data submission. By default, the reports that display for reviewing are provided for a 31-day period. However, uploaders can view reports outside of the 31-day default period by entering start and end dates for the desired date range.

Perform the following steps to view upload reports:

1. Open an Internet browser window and type the following URL in the address bar: http://www.hidesigns.com/flpdmp.

A window similar to the following is displayed:



2. Click **Dispenser**, and then click **RxSentry Dispenser's Upload Site**.

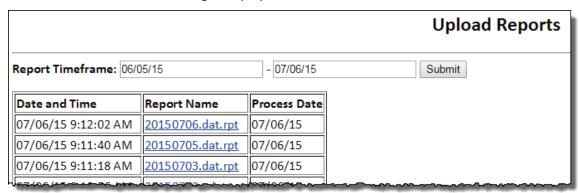
A login window is displayed.

- 3. Type your user name in the **User Name** field.
- 4. Type your password in the **Password** field.
- 5. Click OK.

The RxSentry home page is displayed.

6. Click View Upload Reports.

A window similar to the following is displayed:



- 7. Click a hyperlink in the **Report Name** column to open an upload report for viewing.
- 8. To view reports for a different timeframe, type a start and end date in the **Report Timeframe** fields, and then click **Submit**.

View Zero Reports

This function provides uploaders the ability to view previously submitted zero reports. By default, the reports that display for reviewing are provided for a 31-day period. However, uploaders can view reports outside of the 31-day default period by entering start and end dates for the desired date range.

Perform the following steps to view zero reports:

1. Open an Internet browser window and type the following URL in the address bar: http://www.hidesigns.com/flpdmp.

A window similar to the following is displayed:



2. Click **Dispenser**, and then click **RxSentry Dispenser's Upload Site**.

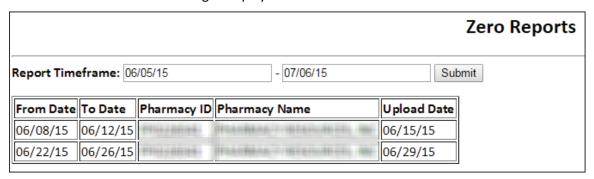
A login window is displayed.

- 3. Type your user name in the **User Name** field.
- 4. Type your password in the **Password** field.
- 5. Click OK.

The RxSentry home page is displayed.

6. Click View Zero Reports.

A window similar to the following is displayed:



7. To view zero report submissions for a different timeframe, type a start and end date in the **Report Timeframe** fields, and then click **Submit**.

Error Correction

Fatal errors will cause a record NOT to be loaded. If this occurs, correct the data that caused the error and resubmit the entire record. Fatal error corrections must be resubmitted within seven (7) days of the initial record submission. If a record with a serious or minor error is loaded and a correction is required, records can be corrected using the DSP01 values as explained below.

Note: Edit Number V1, as shown in the Edit Definitions table, should not be resubmitted. All other records with errors that are not fatal will be loaded unless the batch thresholds are reached. Error thresholds are defined in the Upload Reports section.

The ASAP 4.2 standard requires a dispenser to select an indicator in the DSP01 (Reporting Status) field. Dispensers may submit new records, revise and resubmit records, and void (delete) erroneous records. These actions are indicated by supplying one of the following values in the DSP01 field:

- 00 New Record indicates a new record
- 01 Revise indicates that one or more data elements in a previously-submitted record has been revised
- 02 Void indicates that the original record should be voided

Use the information in the following topics to create, revise/resubmit, or void an erroneous record.

Submit a New Record

Perform the following steps to submit a new record:

- 1. Create a record with the value 00 in the **DSP01** field.
- 2. Populate all other required fields and submit the record.

Note: These steps are used to submit new records *or* to submit records that were previously submitted but received a fatal status on your error report. **Records with fatal errors are not loaded to the system**. The errors in these records must be corrected in your system and resubmitted using the 00 status in the **DSP01** field.

Revise a Record

Perform the following steps to revise a record:

- 1. Create a record with the value 01 in the **DSP01** field.
- 2. Populate the following fields with the same information originally submitted in the erroneous record:
 - PHA03 (DEA Provider ID)
 - DSP02 (Prescription Number)
 - DSP05 (Date Filled)
- 3. Fill in all other data fields with the correct information. This information will override the original data linked to the fields referenced in step 2.
- 4. Submit the record.

Import note: If any of the fields referenced in step 2 are part of the correction, the record must first be voided using the steps provided in the Void a Record section, and then you must re-submit the record using the value 00 in the **DSP01** field.

Void a Record

Perform the following steps to void (delete) a record:

- 1. Send a record with the value 02 in the **DSP01** field.
- 2. Fill in all other data identical to the original record. This will void the original record submission.

Edit Definitions

The following table describes the current list of edits:

Edit Number	Message	Severity
Edit 01	Format of File Error	Fatal
Edit 02	Pharmacy DEA is blank	Fatal
Edit 05	Pharmacy ID not found	Fatal
Edit 09	Invalid DOB	Serious
Edit 10	Gender must be valid	Serious
Edit 14	Reporting Status Invalid	Fatal
Edit 15	Date Dispensed is invalid or irrational	Serious
Edit 17	Refill Code must be a valid number	Minor
Edit 18	Quantity is invalid	Serious
F-J:+ 10	Days Supply is Invalid	Minor
Edit 19	Days Supply is 999	Fatal
5 l'1 20	Days Supply > 150	Minor
Edit 20	Days Supply > 360	Serious
5 lii 24	NDC not found	Serious
Edit 21	NDC not found (used when CDI segment is used)	Fatal
Edit 22	Product ID Qualifier is invalid	Fatal
E-11:4-2E	Prescriber ID not found	Minor
Edit 25	Prescriber ID cannot be blank	Fatal
Edit 28	Date RX Written is invalid	Minor
Edit 29	Number Refill Authorized Invalid	Minor
Edit 50	Customer Last Name blank	Fatal
Edit 51	Customer First Name blank	Fatal
Edit 52	Customer Address blank	Serious
Edit 53	Customer ZIP Code is blank	Serious
Edit 54	Customer ZIP and State Code conflict	Serious
Edit 56	Customer City is blank	Minor
Edit 60	Customer State Code is blank	Serious
Edit 61	Customer State Code is invalid	Serious
Edit 200	Prescription Number is blank	Serious
Edit 360	Date dispensed prior to December 1, 2010	Fatal
Edit V1	Record already exists Note: Duplicate records are not loaded. The number of duplicate records, if any, is displayed on the upload report produced after data file transmission has completed.	Minor

7 Assistance and Support

Technical Assistance

If you need additional help with any of the procedures outlined in this guide, you can:

Contact HID by e-mail at flpdmp-info@hidesigns.com;

OR

Call the HID Help Desk at 877-719-3120.

Administrative Assistance

If you have non-technical questions regarding E-FORCSE, please contact:

E-FORCSE, Florida's Prescription Drug Monitoring Program 4052 Bald Cypress Way, Bin C-16 Tallahassee, Florida 32399

Phone: 850-245-4797

E-mail: e-forcse@doh.state.fl.us Website: www.e-forcse.com

Florida Department of Health Dispenser's Implementation Guide		Assistance and Support
	This page intentionally left blank.	

8 Glossary

ASAP

American Society for Automation in Pharmacy

Batch

Group of files (report or query requests) that are processed in the background while other work is continued

Dispenser

Pharmacy, dispensing pharmacist, or dispensing health care practitioner which dispenses controlled substances

E-FORCSE

Electronic-Florida Online Reporting of Controlled Substances Evaluation (E-FORCSE); the name of Florida's Prescription Drug Monitoring Program

FTP

File Transfer Protocol; commonly-used protocol for exchanging files over any network

Healthcare practitioner

A practitioner who is subject to licensure or regulation by the Florida Department of Health under chapters 458, 459, 461, 462, 464, 465, and 466, F.S.

HID

Health Information Designs, LLC

NDC

National Drug Code; describes specific drugs by drug manufacturer and package size

PMP

Prescription Monitoring Program; term used by ASAP

PDMP

Prescription Drug Monitoring Program

PDMS

Prescription Drug Monitoring System

Prescriber

A practitioner who is authorized by state and federal agencies to prescribe controlled substances

RxSentry

Prescription drug monitoring system developed by Health Information Designs, LLC

SFTP

Secure File Transfer Protocol (also referred to as "SSH File Transfer Protocol"); provides file transfer and manipulation functionality over any reliable data stream

SSL

Secure Sockets Layer; cryptographic protocol that provides secure communications for data transfers

Universal Claim Form (UCF)

Form used by someone who does not have electronic capability to send data; must be approved by E-FORCSE

Uploader

A dispenser that uploads a data file containing controlled substance dispensing information

9 Document Information

Version History

The Version History records the publication history of this document.

Publication Date	Version Number	Comments
06/28/2011	1.0	Initial publication
07/07/2011	1.1	Updated publication
07/20/2011	1.2	Updated publication
07/25/2011	1.3	Updated publication
08/03/2011	1.4	Updated publication
08/10/2011	1.5	Updated publication
08/11/2011	1.6	Updated publication
09/06/2011	1.7	Updated publication
09/12/2011	1.8	Updated publication
10/11/2011	1.9	Updated publication
03/16/2012	1.10	Updated publication
09/10/2012	1.11	Updated publication
07/25/2014	1.12	Updated publication
07/06/2015	2.0	Updated publication

Change Log

The Change Log records the changes and enhancements included in each version.

Version Number	Chapter/Section	Change
1.0	N/A	N/A
1.1	Appendix A: ASAP 4.1 Specifications	Changed the field usage for the following fields from RR to S: PHA05 PHA07 PHA08 PHA09

Version Number	Chapter/Section	Change
1.2	Appendix A: ASAP 4.1 Specifications	Changed the field usage for the following fields from RR to S: PRE01 PRE04 Changed the field usage for the following fields from S to RR: PRE05 PRE06
1.3	Chapter 6/Edit Definitions	Added the following edits and definitions: Edit 14, Edit 17, Edit 19, Edit 20, Edit 29
1.4	Throughout	Replaced screen shots with Florida PDMP system screen shots
1.5	Chapter 5/Universal Claim Form (UCF) Submission	Revised subsections for clarity
1.6	Appendix A: ASAP 4.1 Specifications	Removed information about using a backslash (\) as a segment terminator; only the tilde (~) should be used as a segment terminator
1.7	Appendix A: ASAP 4.1 Specifications	Additional clarification added to Segment Terminator information and in the description for field TH09
1.8	Chapter 5/Online UCF Submission	Updated screenshot of online UCF entry form
1.9	Appendix B: Universal Claim Form	Updated paper UCF to add Gender Code, Prescriber State License Number, and Refill Number
1.10	Chapter 4/Emergency Suspension	Added new topic
1.11	 Chapter 3/ Required Prescription Information Appendix A/ASAP Specifications Table 	Added a note to instruct dispensers to populate field PAT15 with "99" if the patient address is outside the U.S.
1.12	Chapter 4/Report Zero Activity – RxSentry	Separated topic into two sub-topics, Single Dispensers and Group Pharmacies, to clarify the way zero reports must be submitted for each group

Version Number	Chapter/Section	Change
	 Chapter 4/Report Zero Activity – File Upload Chapter 6/View Zero Reports Appendix B: Zero Report Specifications 	Added new topics
2.0	Global	 Updated to new HID document format Updated public site URL Updated screenshots, as necessary Updated to ASAP 4.2
	Chapter 5/Submitting Information Using the Online UCF	Updated the screenshot of the online UCF
	Chapter 7/Technical Assistance	Updated HID Help Desk e-mail address
	Appendix A/ASAP Specifications Table	Updated to ASAP 4.2 specifications
	Appendix C/Paper UCF	Updated paper UCF

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Appendix A: ASAP 4.2 Specifications

The information on the following pages contains the definitions for the specific contents required of uploaded records in the American Society for Automation in Pharmacy (ASAP) format to comply with E-FORCSE requirements.

The following elements are used in each upload file:

- Segment Identifier indicates the beginning of a new segment, for example, PHA.
- Data Delimiter character used to separate segments and the data elements within a segment, for example, an asterisk (*).
 - Each completed field should be followed by an asterisk, and each blank field should contain a single asterisk.
 - If the last field in the segment is blank, it should contain an asterisk and a tilde ($^{\sim}$).
- Segment Terminator character used to mark the end of a segment, for example, the tilde
 (~).

Note: Field TH09 in the Transaction Header segment contains a built-in segment terminator. Since TH09 also signifies the end of the segment, it should contain two tildes (~~).

Field Usage

- R = Required by ASAP
- o RR = Required by E-FORCSE
- S = Situational (not required; however, supply if available)

Both "R" and "RR" fields must be reported.

Note: For more information regarding ASAP 4.2 specifications, please contact the American Society for Automation in Pharmacy at www.asapnet.org for the full *Implementation Guide for the ASAP Standard for Prescription-Monitoring Programs*. This guide includes field lengths, acceptable attributes, and examples.

Segment	Field ID	Field Name	Field Usage			
TH: Transaction Header						
-		indicate the start of a transaction. It also assigns the data element sep	parator, segment			
terminator, a	erminator, and control number.					
	TH01	Version/Release Number	R			
		Code uniquely identifying the transaction.				
		Format = xx.x				
	TH02	Transaction Control Number	R			
		Sender assigned code uniquely identifying a transaction.				
	TH03	Transaction Type	R			
		Identifies the purpose of initiating the transaction.				
		■ 01 Send/Request Transaction				
		02 Acknowledgement (used in Response only)				
		03 Error Receiving (used in Response only)				
		04 Void (used to void a specific Rx in a real-time transmission or				
		an entire batch that has been transmitted)				
	TH04	Response ID	S			
		Contains the Transaction Control Number of a transaction that				
		initiated the transaction. Required in response transaction only.				
	TH05	Creation Date	R			
		Date the transaction was created. Format: CCYYMMDD.				
	TH06	Creation Time	R			
		Time the transaction was created. Format: HHMMSS or HHMM.				
	TH07	File Type	R			
		■ P = Production				
		■ T = Test				
	TH08	Routing Number	S			
		Reserved for real-time transmissions that go through a network				
		switch to indicate, if necessary, the specific state PMP the				
		transaction should be routed to.				
	TH09	Segment Terminator Character	R			
		This terminates the TH segment and sets the actual value of the data				
		segment terminator for the entire transaction.				
IS: Information						
Required segr		o convey the name and identification numbers of the entity supplying t	he information.			
	IS01	Unique Information Source ID	R			
		Reference number or identification number.				
		(Example: phone number)				
	ISO2	Information Source Entity Name	R			
		Entity name of the Information Source.				
	IS03	Message	S			
		Free-form text message.				
l			I .			

Segment	Field ID	Field Name	Field Usage			
PHA: Pharma	acy Header	•				
Required seg	ment; used t	o identify the pharmacy.				
Note: It is re	te: It is required that information be provided in at least one of the following fields: PHA01, PHA02, or PHA03.					
	PHA01	National Provider Identifier (NPI)	S			
		Identifier assigned to the pharmacy by CMS.				
	PHA02	NCPDP/NABP Provider ID	S			
		Identifier assigned to pharmacy by the National Council for Prescription Drug Programs.				
	PHA03	DEA Number	RR			
		Identifier assigned to the pharmacy by the Drug Enforcement Administration.				
	PHA04	Pharmacy Name	S			
		Free-form name of the pharmacy.				
	PHA05	Address Information – 1	S			
		Free-form text for address information.				
	PHA06	Address Information – 2	S			
		Free-form text for address information.				
	PHA07	City Address	S			
		Free-form text for city name.				
	PHA08	State Address	S			
		U.S. Postal Service state code.				
	PHA09	ZIP Code Address	S			
		U.S. Postal Service ZIP Code.				
	PHA10	Phone Number	S			
		Complete phone number including area code.				
	PHA11	Contact Name	S			
		Free-form name.				
	PHA12	Chain Site ID	S			
		Store number assigned by the chain to the pharmacy location. Used when the PMP needs to identify the specific pharmacy from which information is required.				
PAT: Patient Required seg		o report the patient's name and basic information as contained in the p	oharmacy record.			
	PAT01	ID Qualifier of Patient Identifier	S			
		Code identifying the jurisdiction that issues the ID in PAT03.				

Segment	Field ID	Field Name	Field Usage
	PAT02	ID Qualifier	S
		Code to identify the type of ID in PAT03. If PAT02 is used, PAT03 is	
		required.	
		• 01 Military ID	
		02 State Issued ID	
		03 Unique System ID	
		 04 Permanent Resident Card (Green Card) 	
		05 Passport ID	
		• 06 Driver's License ID	
		 07 Social Security Number 	
		08 Tribal ID	
		99 Other (agreed upon ID)	
	PAT03	ID of Patient	S
		Identification number for the patient as indicated in PAT02.	
		An example would be the driver's license number.	
	PAT04	ID Qualifier of Additional Patient Identifier	S
		Code identifying the jurisdiction that issues the ID in PAT06.	
		Used if the PMP requires such identification.	
	PAT05	Additional Patient ID Qualifier	S
		Code to identify the type of ID in PAT06 if the PMP requires a second identifier. If PAT05 is used, PAT06 is required.	
		■ 01 Military ID	
		02 State Issued ID	
		03 Unique System ID	
		 04 Permanent Resident Card 	
		■ 05 Passport ID	
		06 Driver's License ID	
		 07 Social Security Number 	
		08 Tribal ID	
		99 Other (agreed upon ID)	
	PAT06	Additional ID	S
		Identification that might be required by the PMP to further identify the individual. An example might be that in PAT03 driver's license is required and in PAT06 Social Security number is also required.	
	PAT07	Last Name	RR
		Patient's last name.	
	PAT08	First Name	RR
		Patient's first name.	
	РАТО9	Middle Name	S
		Patient's middle name or initial if available.	
	PAT10	Name Prefix	S
		Patient's name prefix such as Mr. or Dr.	

Segment	Field ID	Field Name	Field Usage
	PAT11	Name Suffix	S
		Patient's name suffix such as Jr. or the III.	
	PAT12	Address Information – 1	RR
		Free-form text for street address information.	
	PAT13	Address Information – 2	S
		Free-form text for additional address information.	
	PAT14	City Address	RR
		Free-form text for city name.	
	PAT15	State Address	RR
		U.S. Postal Service state code	
		Note: Field has been sized to handle international patients not	
		residing in the U.S.	
	PAT16	ZIP Code Address	RR
		U.S. Postal Service ZIP code.	
		Populate with zeros if patient address is outside the U.S.	
	PAT17	Phone Number	S
		Complete phone number including area code.	
	PAT18	Date of Birth	RR
		Date patient was born.	
		Format: CCYYMMDD	
	PAT19	Gender Code	RR
		Code indicating the sex of the patient.	
		F Female	
		M Male	
		■ U Unknown	
	PAT20	Species Code	S
		Used if required by the PMP to differentiate a prescription for an	
		individual from one prescribed for an animal.	
		• 01 Human	
		02 Veterinary Patient	

Segment	Field ID	Field Name	Field Usage
	PAT21	Patient Location Code	S
		Code indicating where patient is located when receiving pharmacy	
		services.	
		■ 01 Home	
		02 Intermediary Care	
		03 Nursing Home	
		 04 Long-Term/Extended Care 	
		05 Rest Home	
		 06 Boarding Home 	
		 07 Skilled-Care Facility 	
		 08 Sub-Acute Care Facility 	
		 09 Acute Care Facility 	
		 10 Outpatient 	
		■ 11 Hospice	
		■ 98 Unknown	
		■ 99 Other	
	PAT22	Country of Non-U.S. Resident	S
		Used when the patient's address is a foreign country and PAT12	
		through PAT16 are left blank.	
	PAT23	Name of Animal	S
		Used if required by the PMP for prescriptions written by a	
		veterinarian and the pharmacist has access to this information at	
		the time of dispensing the prescription.	
DSP: Dispens Required Seg the date and	gment; used t	o identify the basic components of a dispensing of a given prescription	order including
	DSP01	Reporting Status	R
		DSP01 requires one of the following codes, and an empty or blank	
		field no longer indicates a new prescription transaction:	
		 00 New Record (indicates a new prescription dispensing transaction) 	
		 01 Revise (indicates that one or more data element values in a previously submitted transaction are being revised) 	
		 02 Void (message to the PMP to remove the original prescription 	
		transaction from its data, or to mark the record as invalid or to be ignored).	
	DSP02		RR
	DSP02	ignored).	RR
	DSP02	ignored). Prescription Number	RR RR
		ignored). Prescription Number Serial number assigned to the prescription by the pharmacy.	
		ignored). Prescription Number Serial number assigned to the prescription by the pharmacy. Date Written	
		ignored). Prescription Number Serial number assigned to the prescription by the pharmacy. Date Written Date the prescription was written (authorized).	

Segment	Field ID	Field Name	Field Usage
	DSP05	Date Filled	RR
		Date prescription was filled. Format: CCYYMMDD	
	DSP06	Refill Number	RR
		Number of the fill of the prescription.	
		0 indicates New Rx; 01-99 is the refill number.	
	DSP07	Product ID Qualifier	R
		Used to identify the type of product ID contained in DSP08.	
		■ 01 NDC	
		06 Compound	
	DSP08	Product ID	RR
		Full product identification as indicated in DSP07, including leading	
		zeros without punctuation.	
	DSP09	Quantity Dispensed	RR
		Number of metric units dispensed in metric decimal format.	
		Example: 2.5	
		Note: For compounds show the first quantity in CDI04.	
	DSP10	Days Supply	R
		Estimated number of days the medication will last.	
	DSP11	Drug Dosage Units Code	S
		Identifies the unit of measure for the quantity dispensed in DSP09.	
		• 01 Each	
		02 Milliliters (ml)	
		• 03 Grams (gm)	_
	DSP12	Transmission Form of Rx Origin Code	S
		Code indicating how the pharmacy received the prescription.	
		01 Written Prescription02 Telephone Prescription	
		03 Telephone Emergency Prescription	
		04 Fax Prescription	
		O5 Electronic Prescription	
		99 Other	
	DSP13	Partial Fill Indicator	S
		Used when the quantity in DSP09 is less than the metric quantity per	_
		dispensing authorized by the prescriber. This dispensing activity is	
		often referred to as a split filling.	
		 00 Not a Partial Fill 	
		01 First Partial Fill	
		Note: For additional fills per prescription, increment by 1. So the	
	D0511	second partial fill would be reported as 02, up to a maximum of 99.	
	DSP14	Pharmacist National Provider Identifier (NPI)	S
		Identifier assigned to the pharmacist by CMS. This number can be used to identify the pharmacist dispensing the medication.	

Segment Field ID Field Name		Field Name	Field Usage
	DSP15	Pharmacist State License Number This data element can be used to identify the pharmacist dispensing the medication. Assigned to the pharmacist by the State Licensing Board.	S
	DSP16	Classification Code for Payment Type Code identifying the type of payment (i.e., how it was paid for). O1 Private Pay O2 Medicaid O3 Medicare O4 Commercial Insurance O5 Military Installations and VA O6 Workers' Compensation O7 Indian Nations 99 Other	RR
	DSP17	Date Sold Usage of this field depends on the pharmacy having a point-of-sale system that is integrated with the pharmacy management system to allow a bidirectional flow of information.	S
	DSP18	RxNorm Product Qualifier 1 01 Semantic Clinical Drug (SCD) 2 02 Semantic Branded Drug (SBD) 3 Generic Package (GPCK) 4 04 Branded Package (BPCK) Note: DSP18 and DSP19 are placeholder fields pending RxNorm becoming an industry standard and should not be required until such time.	S
	DSP19	RxNorm Code Used for electronic prescriptions to capture the prescribed drug product identification. Note: DSP18 and DSP19 are placeholder fields pending RxNorm becoming an industry standard and should not be required until such time.	S
	DSP20	Electronic Prescription Reference Number Used to provide an audit trail for electronic prescriptions. Note: DSP20 and DSP21 should be reported as a pair to the prescription drug monitoring program, and each program decides which one, if not both, it decides to capture.	S
	DSP21	Electronic Prescription Order Number Note: DSP20 and DSP21 should be reported as a pair to the prescription drug monitoring program, and each program decides which one, if not both, it decides to capture.	S

Segment	Field ID	Field Name	Field Usage
PRE: Prescrib	er Informati	ion	
Required segr	ment; used t	to identify the prescriber of the prescription.	
	PRE01	National Provider Identifier (NPI)	S
		Identifier assigned to the prescriber by CMS.	
	PRE02	DEA Number	RR
		Identifying number assigned to a prescriber or an institution by the Drug Enforcement Administration (DEA).	
	PRE03 DEA Number Suffix		S
		Identifying number assigned to a prescriber by an institution when the institution's number is used as the DEA number.	
	PRE04	Prescriber State License Number Identification assigned to the prescriber by the State Licensing Board.	S
	PRE05 Last Name Prescriber's last name.		RR
	PRE06	First Name Prescriber's first name.	RR
	PRE07	Middle Name Prescriber's middle name or initial.	S
	PRE08	Phone Number	S

CDI: Compound Drug Ingredient Detail

Required when medication dispensed is a compound and one of the ingredients is a PMP reporting drug. If more than one ingredient is for a prescription-monitoring program reporting drug, then this would be incremented by one for each compound ingredient being reported.

Used to identify the individual ingredients that make up a compounded drug.

If CDI is filled in, the NDC of DSP08 must be 99999999999.

CDI01	Compound Drug Ingredient Sequence Number First reportable ingredient is 1; each additional reportable ingredient is incremented by 1.	R
CDI02	Product ID Qualifier Code to identify the type of product ID contained in CDI03. 1 01 NDC 1 02 UPC 1 03 HRI 1 04 UPN 1 05 DIN 1 06 Compound (this code is not used in this segment)	R
CDI03	Product ID Full product identification as indicated in CDI02, including leading zeros without punctuation.	R

Segment	Field ID	Field Name	Field Usage
	CDI04	Compound Ingredient Quantity	R
		Metric decimal quantity of the ingredient identified in CDI03.	
		Example: 2.5	
	CDI05	Compound Drug Dosage Units Code	S
		Identifies the unit of measure for the quantity dispensed in CDI04.	
		01 Each (used to report as package)	
		 02 Milliliters (ml) (for liters, adjust to the decimal milliliter equivalent) 	
		 03 Grams (gm) (for milligrams, adjust to the decimal gram equivalent) 	

AIR: Additional Information Reporting

Use of this segment is situational; used when state-issued serialized Rx pads are used, the state requires information on the person dropping off or picking up the prescription, or for data elements not included in other detail segments.

Note: If this segment is used, at least one of the data elements (fields) will be required.

AIR01	State Issuing Rx Serial Number	S
	U.S.P.S. state code of state that issued serialized prescription blank.	
	This is required if AIR02 is used.	
AIR02	State Issued Rx Serial Number	S
	Number assigned to state issued serialized prescription blank.	
AIR03	Issuing Jurisdiction	S
	Code identifying the jurisdiction that issues the ID in AIR05.	
AIR04	ID Qualifier of Person Dropping Off or Picking Up Rx	S
	Used to identify the type of ID contained in AIR05 for person	
	dropping off or picking up the prescription.	
	01 Military ID	
	 02 State Issued ID 	
	 03 Unique System ID 	
	 04 Permanent Resident Card (Green Card) 	
	■ 05 Passport ID	
	 06 Driver's License ID 	
	 07 Social Security Number 	
	■ 08 Tribal ID	
	99 Other (agreed upon ID)	
AIR05	ID of Person Dropping Off or Picking Up Rx	S
	ID number of patient or person picking up or dropping off the	
	prescription.	

Segment	Field ID	Field Name	Field Usage
	AIR06	Relationship of Person Dropping Off or Picking Up Rx	S
		Code indicating the relationship of the person.	
		■ 01 Patient	
		02 Parent/Legal Guardian	
		03 Spouse	
		04 Caregiver	
		• 99 Other	
	AIR07	Last Name of Person Dropping Off or Picking Up Rx	S
		Last name of person picking up the prescription.	
	AIR08	First Name of Person Dropping Off or Picking Up Rx	S
		First name of person picking up the prescription.	
	AIR09	Last Name or Initials of Pharmacist	S
		Last name or initials of pharmacist dispensing the medication.	
	AIR10	First Name of Pharmacist	S
		First name of pharmacist dispensing the medication.	
	AIR11	Dropping Off/Picking Up Identifier Qualifier	S
		Additional qualifier for the ID contained in AIR05	
		■ 01 Person Dropping Off	
		 02 Person Picking Up 	
		 98 Unknown/Not Applicable 	
		Note: Both 01 and 02 cannot be required by a prescription drug	
		monitoring program.	
TP: Pharmacy	Trailer		
		o identify the end of data for a given pharmacy and provide the count of separated for the pharmacy, including the PHA and TP segment.	of the total
	TP01	Detail Segment Count	R
		Number of detail segments included for the pharmacy including the pharmacy header (PHA) and the pharmacy trailer (TP) segments.	
TT: Transaction	n Trailer		
		o indicate the end of the transaction and provide the count of the total	number of
segments inclu	uded in the t	ransaction.	
	TT01	Transaction Control Number	R
		Identifying control number that must be unique.	
		Assigned by the originator of the transaction.	
		Must match the number in TH02.	
	TT02	Segment Count	R
		Total number of segments included in the transaction including the header and trailer segments.	

Florida Department of Health Dispenser's Implementation Guide		Appendix A:ASAP 4.2 Specifications
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Appendix B: Zero Report Specifications

The information on the following pages contains the definitions for the specific contents required by the American Society for Automation in Pharmacy (ASAP) to comply with zero dispense reporting for the FL PDMP.

The zero report specification is a complete transaction that includes the information that would normally be sent with a batch file filled out as it would be for reporting the dispensing of controlled substances. However, for the detail segments, while all the segments and data elements that are required by the FL PDMP are sent, only the Patient First Name, Last Name, and Date Filled fields are populated. The values populating these fields are:

- First Name = Zero
- Last Name = Report
- Date Filled = Date that the report is sent

All other fields in the detail segments should be left blank.

The following elements are used in each upload file:

- Segment Identifier indicates the beginning of a new segment, for example, PHA.
- Data Delimiter character used to separate segments and the data elements within a segment, for example, an asterisk (*).
 - Each completed field should be followed by an asterisk, and each blank field should contain a single asterisk.
 - If the last field in the segment is blank, it should contain an asterisk and a tilde (~).
- Segment Terminator character used to mark the end of a segment, for example, the tilde
 (~).

Note: Field TH09 in the Transaction Header segment contains a built-in segment terminator. Since TH09 also signifies the end of the segment, it should contain two tildes (~~).

For more information, contact the American Society for Automation in Pharmacy for the full Implementation Guide for the ASAP Standard for Prescription Drug Monitoring Programs Zero Reports.

Segment	Field ID	Field Name	Field Usage
TH: Transact	ion Header		
Used to indic		of a transaction. It also assigns the data element separator, segment	terminator, and
	TH01	Version/Release Number	R
	TH02	Transaction Control Number Sender assigned code uniquely identifying a transaction.	R
	TH03	Transaction Type Identifies the purpose of initiating the transaction. 01 Send/Request Transaction	R
	TH04	Response ID	N
	TH05	Creation Date Time the transaction was created. Format: HHMMSS or HHMM.	R
	TH06	Creation Time Time the transaction was created. Format: HHMMSS or HHMM.	R
	TH07	File Type P = Production	R
	TH08	Routing Number	N
	TH09	Segment Terminator Character TH09 also signifies the end of this segment; therefore, it should contain two tildes (~~).	R
IS: Informati		and identification numbers of the entity supplying the information.	
	IS01	Unique Information Source ID Reference number or identification number.	R
	ISO2	Information Source Entity Name Entity name of the Information Source.	R
	IS03	Message Enter the date range in the following format: #yyyymmdd#- #yyyymmdd#~.	R
PHA: Pharma Used to iden	•	macv.	
	PHA01	National Provider Identifier (NPI) Identifier assigned to the pharmacy by CMS.	N
	DUAGO		N.
	PHA02	NCPDP/NABP Provider ID Identifier assigned to pharmacy by the National Council for Prescription Drug Programs.	N
	РНА03	DEA Number Identifier assigned to the pharmacy by the Drug Enforcement Administration.	R

Segment	Field ID	Field Name	Field Usage
PAT: Patient	Information		<u> </u>
Used to repo	ort the patien	t's name and basic information as contained in the pharmacy red	cord.
	PAT01	ID Qualifier of Patient Identifier	N
	PAT02	ID Qualifier	N
	PAT03	ID of Patient	N
	PAT04	ID Qualifier of Additional Patient Identifier	N
	PAT05	Additional Patient ID Qualifier	N
	PAT06	Additional ID	N
	PAT07	Last Name	R
		Required value = Report	
	PAT08	First Name	R
		Required value = Zero	
	PAT09	Middle Name	N
	PAT10	Name Prefix	N
	PAT11	Name Suffix	N
	PAT12	Address Information – 1	N
	PAT13	Address Information – 2	N
	PAT14	City Address	N
	PAT15	State Address	N
	PAT16	ZIP Code Address	N
	PAT17	Phone Number	N
	PAT18	Date of Birth	N
	PAT19	Gender Code	N
DSP: Dispens Used to iden quantity.	_	components of a dispensing of a given prescription order includ	ing the date and
	DSP01	Reporting Status	N
	DSP02	Prescription Number	N
	DSP03	Date Written	N
	DSP04	Refills Authorized	
	D3F04		N
	DSP05	Date Filled Date prescription was filled Format: CCYYMMDD	R
	DSP05	Date prescription was filled. Format: CCYYMMDD	R
	DSP05	Date prescription was filled. Format: CCYYMMDD Refill Number	R N
	DSP05 DSP06 DSP09	Date prescription was filled. Format: CCYYMMDD Refill Number Quantity Dispensed	R N N
DDE: Drosovil	DSP05 DSP06 DSP09 DSP10	Date prescription was filled. Format: CCYYMMDD Refill Number Quantity Dispensed Days Supply	R N
	DSP05 DSP06 DSP09 DSP10 Der Informati	Date prescription was filled. Format: CCYYMMDD Refill Number Quantity Dispensed Days Supply	R N N

Segment	Field ID	Field Name	Field Usage
	PRE02	DEA Number	N
TP: Pharmacy	Trailer		
	=	data for a given pharmacy and provide the count of the total number pharmacy, including the PHA and TP segment.	of detail
	TP01	Detail Segment Count	R
		Number of detail segments included for the pharmacy including the pharmacy header (PHA) including the pharmacy trailer (TP) segments.	
TT: Transaction	n Trailer		
Used to indicat transaction.	e the end of	f the transaction and provide the count of the total number of segmen	ts included in the
	TT01	Transaction Control Number	R
		Identifying control number that must be unique.	
		Assigned by the originator of the transaction.	
		Must match the number in TH02.	
	TT02	Segment Count	R
		Total number of segments included in the transaction including the header and trailer segments.	

Appendix C: Universal Claim Form

The Universal Claim Form is provided on the following page.

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Florida Prescription Drug Monitoring Program Universal Claim Form



The State of Florida now requires that ALL prescriptions for Schedule II-IV Controlled Substances be reported to a data repository managed by the Florida Department of Health. This form may be used with permission from the Florida Department of Health.

Fax: (888) 288-0337 Phone: (877) 719-3120	Fax or Mail to: Health Information Designs, LLC	391 Industry Dr. Auburn, AL 36832
	PATIENT INFORMATION	,
First Name		
First Name	MI Last Name Gender ☐ Female ☐ Male ☐ Unknown	
Address		ZIP
Dispenser Name	DEA NPI	
Phone # (
Address		ZIP
Prescription #1 Reporting Sta	atus New Record Revise Void	
Rx # Date Filled	// Date Written// Refill N	Number
	Drug Name (Strength)	
Quantity Dispensed Days Sup		
	DEA NPI (if available)	
Prescriber State License # (if available)		
Payment Type ☐ Private Pay ☐ Medic ☐ Workers' Compensation	caid Medicare Commercial Insurance Military Indian Nations Other	y Installations/VA
D : : : : : : : : : : : : : : : : : : :		
Prescription #2 Reporting Sta	atus □ New Record □ Revise □ Void	
NDC	Drug Name (Strength)	
Quantity Dispensed Days Sup	pply Refills Authorized	
Prescriber Name		
Prescriber State License # (if available)	Prescriber Phone # (
Payment Type ☐ Private Pay ☐ Medic ☐ Workers' Compensation	eaid Medicare Commercial Insurance Military Indian Nations Other	ry Installations/VA
	PRESCRIPTION INFORMATION	
Prescription #3 Reporting Sta	atus □ New Record □ Revise □ Void	
Rx # Date Filled		Number
NDC	Drug Name (Strength)	
Quantity Dispensed Days Sup	pply Refills Authorized	
Prescriber Name		
Prescriber State License # (if available)		
Payment Type ☐ Private Pay ☐ Medic ☐ Workers' Compensation	caid	ry Installations/VA

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