

Data Submission Guide for Dispensers

North Carolina Controlled Substance Reporting System

December 2024 Version 4.0



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I Document Overview

This document serves as a training guide and support manual for dispensers of Schedule II through Schedule V controlled substances in North Carolina who use Bamboo Health's PMP Clearinghouse repository to report their dispensations. It includes such topics as:

- Reporting requirements for dispensers in or into the State of North Carolina
- Data file submission guidelines and methods
- Creating a PMP Clearinghouse account
- Creating a data file
- Uploading or reporting data
- Understanding and correcting errors

This guide is intended for use by all dispensers in or into the State of North Carolina required to report the dispensing of controlled substances.

2 Data Collection and Tracking

2.1 Data Collection Overview

General Statute 90-113.70 requires that the North Carolina Department of Health and Human Services (NC DHHS) establish and maintain a reporting system of prescriptions for all Schedule II through V controlled substances. This statute requires that all dispensers of licit controlled substances submit information in accordance with the transmission methods and frequency promulgated by the Division of Mental Health, Developmental Disabilities and Substance Abuse Services.

The purpose of this legislation is to improve the ability to identify controlled substance misusers and refer them for treatment, and to identify and stop diversion of prescription drugs in an efficient and cost-effective manner that will not impede the appropriate medical utilization of licit controlled substances.

As per North Carolina General Statute, 90-113.73. Dispensers are required to report to the NC Controlled Substance Reporting System (CSRS) no later than the close of the next business day after the prescription is delivered; however, dispensers are encouraged to report the information no later than 24 hours after the prescription was delivered.

The data shall be reported using one of the electronic formats described in this document. A written report on the Universal Claim Form (UCF) may be accepted, as long as the dispenser has approval from the NC DHHS. All data must be collected in the ASAP 4.2 format defined in <u>Appendix A: ASAP 4.2 Specifications</u>.

Effective January 1, 2019, a person licensed to practice veterinary medicine may submit prescription information by paper form or other means, provided all information required of electronically submitted data is submitted.

Effective March 1, 2024, As of March 1, 2024, Section 11.1, Section 11.2, and Section 11.2A of HB 190 SL2023-65 will become effective.

As of this date, a dispenser's DEA number for prescriptions of controlled substances, and for prescriptions of gabapentin, whether the dispenser has a DEA number, as well as the prescriber's DEA number for prescriptions of controlled substances, and for prescriptions of gabapentin, if the prescriber has a DEA number and the number is known by the dispenser must be reported to the North Controlled Substances Reporting System (CSRS).

Gabapentin is not a scheduled drug and as such may be obtained from prescribers and/or dispensers that do not have a DEA registration. Our vendor, Bamboo Health, has enabled the PMP Clearinghouse system so that the prescribers and dispensers without a DEA registration can submit gabapentin using their NPI.

Effective March 1, 2025, Section 11.3 of HB 190 SL2023-65, which will include a person licensed to practice veterinary medicine as a dispenser required to submit both controlled substances and gabapentin. If a veterinary provider does not have a DEA or NPI number but is prescribing and/or dispensing gabapentin they may utilize the state license failover. To complete this the submitter will leave the PRE01 and PRE02 fields

blank and enter the prescriber's North Carolina registered veterinary license number in PRE04.

Note: See <u>Appendix D: Reporting Gabapentin Prescriptions</u> for guidance on veterinarians.

2.2 Reporting Requirements

All dispensers of controlled substances, as defined by this Act, are required to collect and report the following information to the data repository:

- Dispenser's DEA number
- Name of the patient for whom the controlled substance was dispensed and the patient's:
 - Full address, including city, state, and ZIP code
 - Telephone number (use the customer ID area in the ASAP standards)
 - Date of birth

Note: If you are submitting a veterinary prescription, enter the **owner's** date of birth, not the animal's.

- Date the prescription was written
- Date the prescription was filled
- Prescription number
- Indication of whether the prescription was new or refill
- Metric quantity dispensed
- Estimated days' supply, if provided
- NDC number of the drug dispensed
- Prescriber's DEA number
- National Provider Identifier (NPI) number, for any prescriber that has an NPI number
- Method of payment

Each dispenser must submit the required data to the data repository no later than the close of the next business day after the prescription is delivered; however, dispensers are encouraged to report the information no later than 24 hours after the prescription was delivered.

In the event the dispenser is unable to report the information within the time frame required by this section because the system is not operational or there is some other temporary electrical or technological failure, this inability shall be documented in the dispenser's records. Once the electrical or technological failure has been resolved, the dispenser shall promptly report the information.

The <u>Data Submission</u> chapter provides all the instructions necessary to submit the required information.

Note: If you are a chain pharmacy, your data will likely be submitted from your home office. Please verify this with your home office. If you are an independent pharmacy or other entity, please forward the reporting requirements to your software vendor. They will need to create the data file, and they may be able to submit the data on your behalf. If not, follow the instructions provided in the <u>Data Submission</u> chapter to submit the data.

2.2.1 HIPAA Compliance

Disclosure of prescription monitoring information must comply with the federal Health Insurance Portability and Accountability Act of 1996 (P.L. 104-191) and regulations adopted under it, including 45 C.F.R. parts 160 and 164; federal alcohol and drug treatment confidentiality laws and regulations adopted under those laws, including 42 C.F.R. part 2; and state health information confidentiality laws.

2.2.2 Zero Reports

If a dispenser usually dispenses controlled substances in North Carolina but has no dispenses to report for the preceding reporting period, the pharmacy must report this information to the NC CSRS by filing a zero report, in accordance with IOA NCAC 26E .0602, as described in the Reporting Zero Dispensing topic in this guide.

2.2.3 Reporting Noncompliance

The laws and regulations for reporting to the NC CSRS are continuously subjected to amendments, and it is the dispenser's responsibility to be aware of such updates as they are enacted and promulgated.

As per North Carolina General Statute 90-113.73., any pharmacy that employs dispensers found to have failed to report information in the manner required within a reasonable period of time after being informed that the required information is missing or incomplete may be assessed civil penalties.

2.2.4 Excluded from Reporting

"Dispenser" means a person who delivers a Schedule II through V controlled substance to an ultimate user in North Carolina, but does **not** include any of the following:

- A licensed hospital or long-term care pharmacy that dispenses such substances for inpatient administration
- A wholesale distributor of a Schedule II through V controlled substance
- A physician dispensing a controlled substance medication for no more than a 48-hour supply
- A dispenser is not required to report Schedule V non-narcotic, nonanorectic Schedule V controlled substances provided directly to the ultimate user for the purpose of assessing a therapeutic response when prescribed according to indications approved by the US FDA.

3 Accessing Clearinghouse

This chapter describes how to create your PMP Clearinghouse account and how to log in to the PMP Clearinghouse web portal.

3.1 Creating Your Account

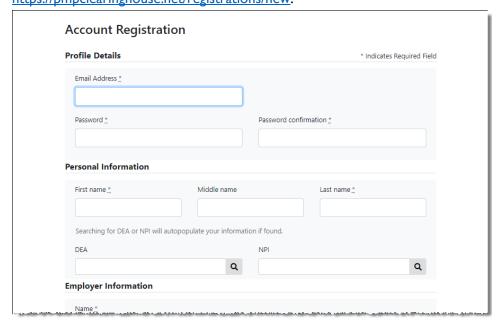
Prior to submitting data, you must create an account. If you are currently registered with the Bamboo Health PMP Clearinghouse system, you do not need to register for a new account—you will be able to add North Carolina to your existing account for data submissions. If you have an existing PMP Clearinghouse account, please refer to Adding PMPs to Your Upload Account to add PMPs to your account.

Notes:

- Data from multiple pharmacies can be uploaded in the same file. For example, chain
 pharmacies may send in one file containing controlled substance dispensing information for
 all their pharmacies licensed in the State of North Carolina. Therefore, chains with multiple
 stores need only to set up one account to upload a file.
- PMP Clearinghouse allows users to submit data through the web portal via manual entry (UCF) or upload of ASAP files. For users who prefer an encrypted transfer method, SFTP access is also available. You may set up your SFTP account during the account creation process.
- If you need to make changes to an existing PMP Clearinghouse upload account, please refer to Managing Your Upload Account.

Perform the following steps to create an account:

 Open an internet browser window and navigate to the PMP Clearinghouse Account Registration page located at https://pmpclearinghouse.net/registrations/new.



2. Complete your Profile Details.

Email Address ** Password ** Password confirmation **	Profile Details		* Indicates Required Fiel
Paraward * Paraward confirmation *	Email Address		
	Password *	Password confirm	ation *

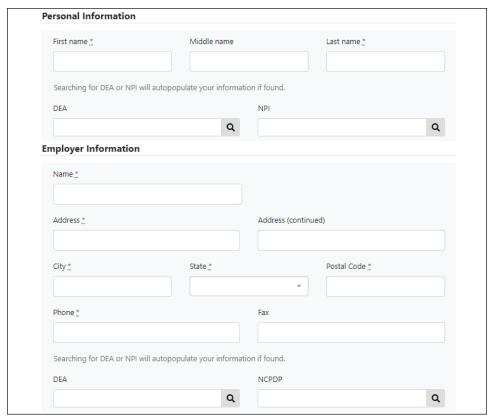
a. Enter your current, valid email address in the **Email Address** field.

Note: The email address you provide here will act as your username when logging into the PMP Clearinghouse system.

b. Enter a password for your account in the **Password** field, then re-enter it in the **Password Confirmation** field. The password requirements are provided below.

Passwords must contain:

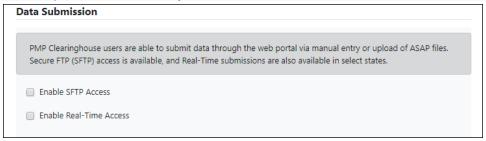
- At least eight (8) characters
- One (1) uppercase letter
- One (1) lowercase letter
- One (1) number
- One (1) special character, such as !, @, #, \$, etc.
- 3. Complete your Personal and Employer information, noting the following:
 - Required fields are marked with a red asterisk (*).
 - You may be able to auto-populate your Personal and/or Employer information by entering your (or your employer's) **DEA**, **NPI**, and/or **NCPDP** number, then clicking the search icon (^Q). If the number you entered is found, your information will automatically be populated.



4. If secure file transfer protocol (SFTP) is required, complete the **Data Submission** section of the page.

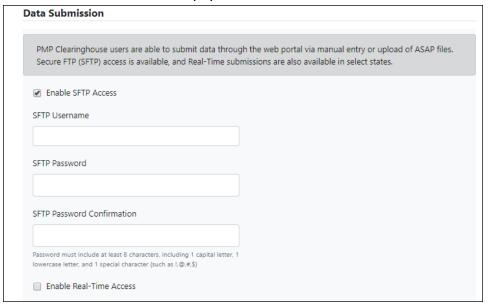
Notes:

- If SFTP access is not required, you do not need to complete the Data Submission section and you may continue to step 5.
- You may add SFTP access to an existing account. Please refer to <u>Adding SFTP Access to an Upload Account</u> for complete instructions.



a. Click to select the **Enable SFTP Access** checkbox.

The **SFTP** access fields are displayed.



- b. Your **SFTP Username** is automatically generated using the first five characters of your employer's name + your employer's phone number. For example, if you entered "Test" as your employer's name and "555-555-5555" as your employer's phone number, your SFTP username would be *test5555555555*.
- c. Enter a password for your SFTP account in the SFTP Password field, then reenter it in the SFTP Password Confirmation field. The password requirements are provided below.

Passwords must contain:

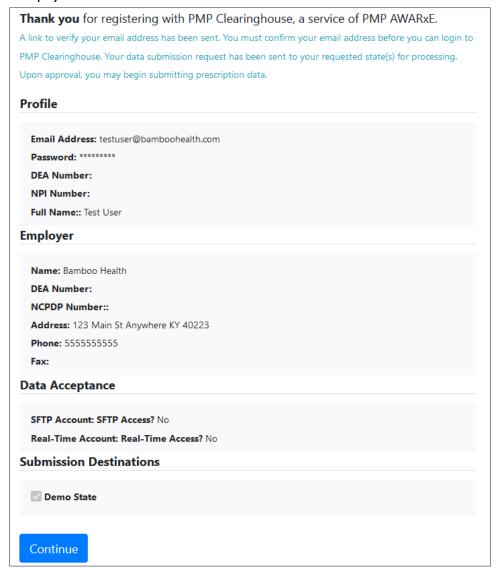
- At least eight (8) characters
- One (1) uppercase letter
- One (1) lowercase letter
- One (1) number
- One (1) special character, such as !, @, #, \$, etc.

This password will be input into the pharmacy software so that submissions can be automated.

Notes:

- This password can be the same as the one previously entered under Profile.
- Unlike the Profile password (i.e., your user account password), the SFTP password does not expire.
- The URL to connect via SFTP is <u>submissions.healthcarecoordination.net/</u>.
- Additional details on SFTP configuration can be found in <u>Appendix C: SFTP</u> Configuration.
- 5. In the **Submission Destinations** section of the page, select the PMP(s) for which you will be submitting data.
- 6. Click Submit.

The request is submitted to the PMP administrator for each of the PMPs you selected for data submission, and the **Registration Information Overview** page is displayed.

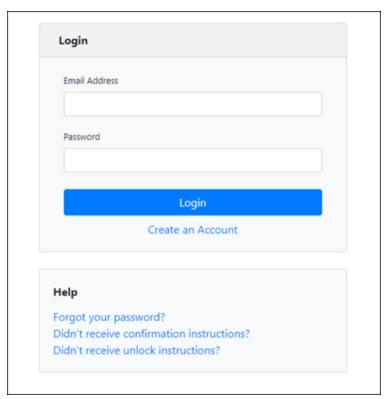


7. Click Continue.

The **PMP Clearinghouse Login** page is displayed. However, you will not be able to log in until your account has been approved. Once the PMP administrator has approved your request, you will receive a welcome email instructing you to confirm your account. Follow the instructions in the email to confirm your account and begin submitting data to PMP AWARxE.

3.2 Logging In to PMP Clearinghouse

I. Open an internet browser window and navigate to the **PMP Clearinghouse Login** page located at https://pmpclearinghouse.net/users/sign in.



- 2. Enter the email address you used to create your account in the **Email Address** field.
- 3. Enter your password in the **Password** field.

Note: If you have forgotten your password, have completed your registration but did not receive the account confirmation email, or your account has been locked and you did not receive the email with instructions for unlocking your account, please refer to the links in the Help section of the page. For detailed instructions on resetting your password, refer to Resetting Your Password.

4. Click Login.

The **PMP Clearinghouse** home page is displayed.



4 Data Submission

This chapter provides information and instructions for submitting data to the PMP Clearinghouse repository.

4.1 Timeline and Requirements

- Pharmacies and software vendors can begin creating their PMP Clearinghouse accounts on or after August 22, 2018. See <u>Creating Your Account</u> for more information.
- You can begin submitting data to PMP Clearinghouse as soon as your account has been created.
- Beginning September 19, 2018, dispensers are required to transmit their data using PMP Clearinghouse in accordance with the guidelines outlined under <u>Reporting</u> <u>Requirements</u>.

Note: If you have any questions about the reporting timeline, please contact the NC CSRS administrator using the information provided in <u>Administrative Assistance</u>.

4.2 Upload Specifications

Files should 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 "20220919.dat". All of your upload files will be kept separate from the files of others.

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

Prescription information must be reported no later than the close of the next business day after the prescription is dispensed; however, dispensers are encouraged to report the information no later than 24 hours after the prescription was dispensed.

4.3 Zero Reports

If you have no dispensations to report for the preceding reporting period, you must report this information to the NC CSRS.

You may submit your zero report through the PMP Clearinghouse web portal by following the steps below or via SFTP using the ASAP Standard for Zero Reports. For additional details on submitting via SFTP, please refer to Appendix B: ASAP Zero Report Specifications.

You may submit zero reports through the PMP Clearinghouse web portal using one of the following methods:

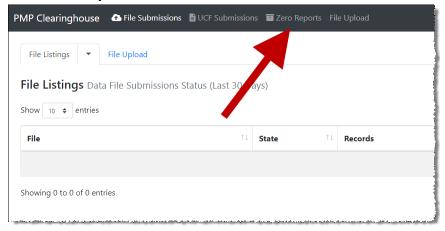
- Submit a single-click zero report
- Create a new zero report

4.3.1 Submit a Single-Click Zero Report

Single-click zero reporting allows you to create a profile for the pharmacy that includes its identifiers (e.g., DEA, NPI, NCPDP), so you do not have to enter it each time you submit a zero report.

To create a pharmacy profile and begin submitting single-click zero reports:

- 1. If you do not have an account, perform the steps in Creating Your Account.
- 2. Log in to PMP Clearinghouse.
- 3. Click Zero Reports.



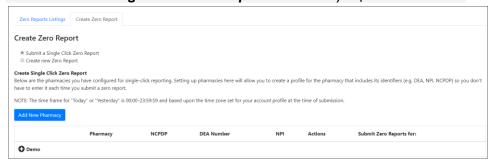
The **Zero Report Listings** page is displayed.



4. Click the **Create Zero Report** tab.

The Create Zero Report page is displayed.

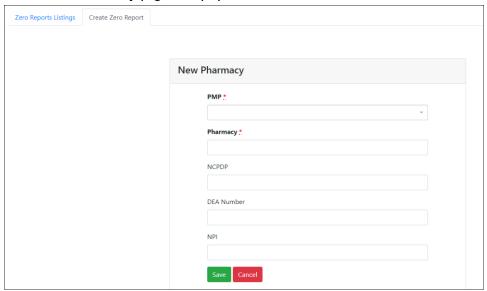
Note: Submit a Single Click Zero Report is selected by default.



- Any pharmacies you have already configured for single-click zero reporting are displayed at the bottom of the page. Continue to <u>step 10</u> to submit a zero report for those pharmacies.
- If you have not configured your pharmacy for single-click zero reporting, continue to step 5.

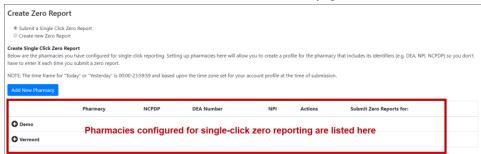
5. Click Add New Pharmacy.

The **New Pharmacy** page is displayed.



- 6. Select the PMP for which you are submitting a zero report from the drop-down list in the **PMP** field.
- 7. Enter the pharmacy's name in the **Pharmacy** field.
- 8. Populate the **NCPDP**, **DEA Number**, and/or **NPI** fields as required by the PMP you selected in step 6. If any of these fields are required, a red asterisk (*) will be displayed next to that field once you have selected a PMP.
- 9. Click Save.

The pharmacy is saved and will be listed under the drop-down for the selected PMP, which is located at the bottom of the page.



10. Click the plus sign ("+") next to the PMP for which you wish to submit a zero report.

The list of pharmacies you have configured for single-click zero reporting for that PMP is displayed.

Note: This page allows you to submit a zero report for the current date (**Today**) or the previous day (**Yesterday**).



- Click **Today** to submit a zero report for the current date;
 Or
- 12. Click **Yesterday** to submit a zero report for the previous date.

Once the report is submitted, the submission is indicated on the screen, and the zero report is displayed on the **Zero Report Listings** tab.

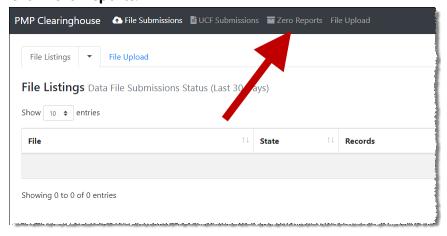


Note: You may edit or delete a pharmacy from this page.

- To edit a pharmacy, click Edit to display the Edit Pharmacy page and make any necessary changes. Refer to steps 6–9 for guidance on entering pharmacy information.
- To delete a pharmacy, click **Delete**. You will be prompted to confirm the deletion. Once you confirm the deletion, the pharmacy configuration will be removed.

4.3.2 Create a New Zero Report

- 1. If you do not have an account, perform the steps in Creating Your Account.
- 2. Log in to PMP Clearinghouse.
- 3. Click **Zero Reports**.



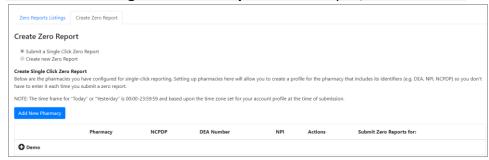
The **Zero Report Listings** page is displayed.



4. Click the Create Zero Report tab.

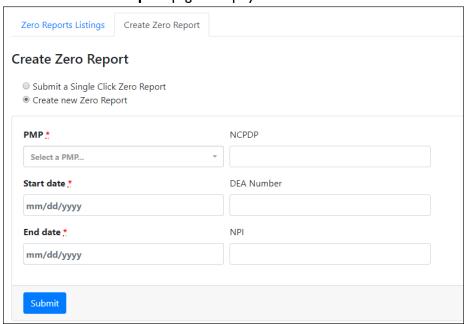
The Create Zero Report page is displayed.

Note: Submit a Single Click Zero Report is selected by default.



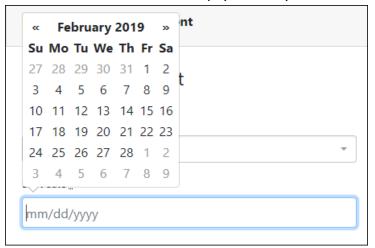
5. Click the button to select **Create new Zero Report**.

The Create Zero Report page is displayed.



6. Select the PMP for which you are submitting a zero report from the drop-down list in the **PMP** field.

7. Enter the start date and end date for the zero report in the **Start date** and **End date** fields using the *MM/DD/YYYY* format. You may also select the dates from the calendar that is displayed when you click in these fields.



8. Enter your NCPDP, DEA, and/or NPI numbers, if required by your PMP.

Note: If any of these fields are required by your PMP, they will be marked with a red asterisk (*).

9. Click Submit.

Your zero report is submitted to PMP Clearinghouse and will be displayed on the **Zero Report Listings** tab.

5 Data Delivery Methods

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

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

Delivery Method	Page
Secure FTP	17
Web Portal Upload	18
Manual Entry (UCF)	20

5.1 Secure FTP

If you are submitting data to PMP Clearinghouse using SFTP, you must configure individual subfolders for the PMP systems to which you are submitting data. These subfolders must be created in the *homedir/directory* folder, which is where you are directed once authenticated, and **should be named using the PMP abbreviation** (e.g., AL, AK, DC, GU, KS, NC, PR, etc.). Data files not submitted to a PMP subfolder will be required to have a manual PMP assignment made on the <u>File Listings</u> page. Please refer to <u>PMP Subfolders</u> for additional details on this process.

I. If you do not have a PMP Clearinghouse account, perform the steps in Creating Your Account.

Or

- 2. If you have a PMP Clearinghouse account but have not enabled SFTP access, perform the steps in Adding SFTP Access to an Upload Account.
- 3. Prepare the data file(s) for submission, using the ASAP specifications described in Appendix A: ASAP 4.2 Specifications.

Notes:

- Files for upload should be named in a unique fashion, with a prefix constructed from the date of submission to PMP Clearinghouse (YYYYMMDD) and a suffix of ".dat". An example file name would be "20220919.dat".
- All of your upload files will be kept separate from the files of others.
- Reports for multiple pharmacies can be in the same upload file in any order.
- 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:
 20220919a.dat, 20220919b.dat, and 20220919c.dat.
- 4. SFTP the file to http://submissions.healthcarecoordination.net/.
- 5. When prompted, enter the username and password you created when setting up the SFTP account.
- 6. Place the file in the appropriate PMP-abbreviated directory.
- 7. You can view the results of the transfer/upload on the Submissions page in PMP Clearinghouse.

Note: If you place the data file in the root directory and not a PDMP sub-folder, a symbol with a mouse over hint of "**Determine PMP**" is displayed on the **File Status** page, and you will be prompted to select a destination PMP to which the data should be sent.

5.2 Web Portal Upload

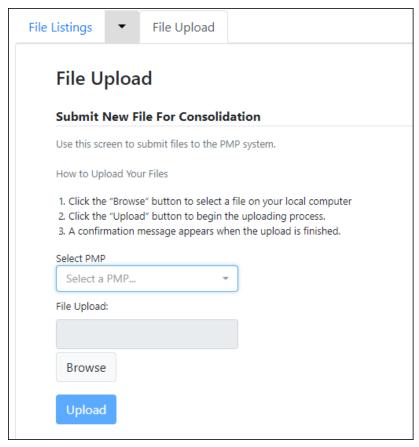
- 1. If you do not have an account, perform the steps in Creating Your Account.
- 2. Prepare the data file(s) for submission, using the ASAP specifications described in Appendix A: ASAP 4.2 Specifications.

Notes:

- Files for upload should be named in a unique fashion, with a prefix constructed from the date of submission to PMP Clearinghouse (YYYYMMDD) and a suffix of ".dat". An example file name would be "20220919.dat".
- All of your upload files will be kept separate from the files of others.
- Reports for multiple pharmacies can be in the same upload file in any order.
- 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: 20220919a.dat, 20220919b.dat, and 20220919c.dat.
- 3. Log in to PMP Clearinghouse.
- 4. From the home page, click the File Upload tab.

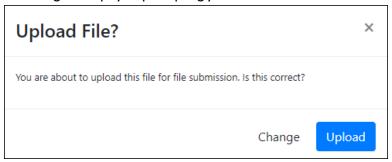


The **File Upload** page is displayed as shown on the following page.



- 5. Select the PMP to which you are submitting the file from the drop-down list in the **Select a PMP** field.
- 6. Click the **Browse** button, located next to the **File Upload** field, and select the file you created in step 2.
- 7. Click **Upload**.

A message is displayed prompting you to confirm the submission.



8. Click **Upload** to continue with the file submission.

Your file is uploaded, and you can view the results of the upload on the File Submissions page.

Note: When uploading a file, the file name must be unique. If the file name is not unique, a message is displayed indicating that the file name has already been taken.

5.3 Manual Entry (UCF)

If you do not have an automated record-keeping system capable of producing an electronic report using the ASAP 4.2 format, prescription information may be submitted on the Universal Claim Form (UCF).

The UCF may be submitted in two ways: online or via paper submission. For more information, see the following topics:

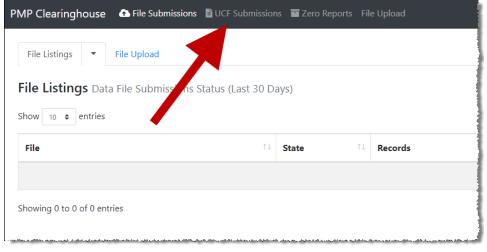
- Online UCF Submission
- Paper UCF Submission

5.3.1 Online UCF Submission

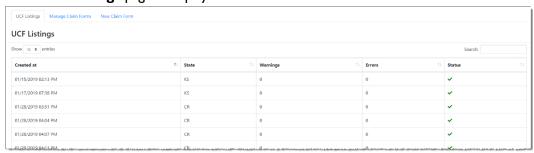
You can manually enter your prescription information into the PMP Clearinghouse system using the Universal Claim Form (UCF) within the PMP Clearinghouse web portal. This form allows you to enter patient, prescriber, dispenser, and prescription information.

Please refer to <u>Reporting Requirements</u> for the complete list of reporting requirements.

- 1. If you do not have an account, perform the steps in Creating Your Account.
- 2. Log in to PMP Clearinghouse.
- 3. Click UCF Submissions.

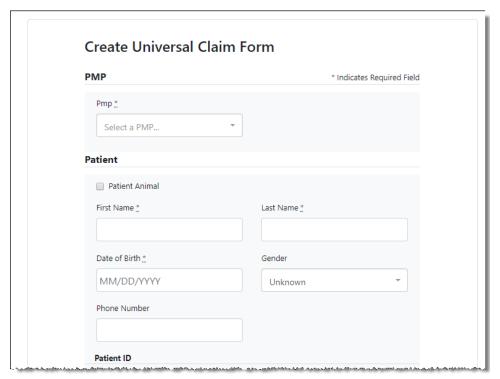


The **UCF Listings** page is displayed.



4. Click the **New Claim Form** tab, located at the top of the page.

The Create Universal Claim Form page is displayed.

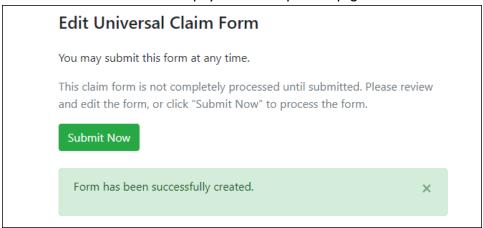


- 5. Select the PMP to which you are submitting data from the drop-down list in the **Select a PMP** field.
- 6. Complete the required fields.

Notes:

- An asterisk (*) indicates a required field.
- If you are entering a compound, click the Compound checkbox in the Drug Information section of the page, complete the required fields for the first drug ingredient, then click **Add New** to add additional drug ingredients.
- 7. Once you have completed all required fields, click Save.

The **Submit Now** button is displayed at the top of the page.



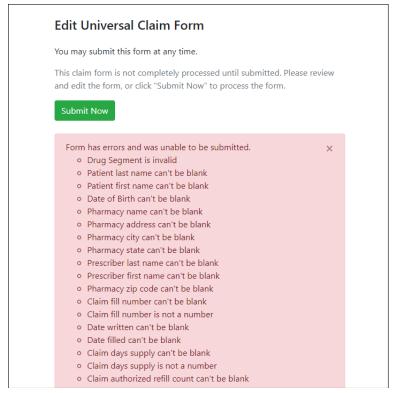
8. Click **Submit Now** to continue with the data submission process.

A message is displayed prompting you to confirm the data submission.



9. Click OK.

Your data will be validated upon submission. If there are any errors on the UCF form, they are displayed at the top of the page.



Note: If there are no errors, you are returned to the Submitted Claim Forms page and your report is listed there.

10. Correct the indicated errors, then repeat steps 7–9.

Once your data has been successfully submitted, your report is listed on the **UCF Listings** page.



5.3.2 Paper UCF Submission

If you do not have an automated record keeping system capable of producing an electronic report following the provided ASAP 4.2 format, and you have been approved by the NC DHHS, you may submit prescription information via paper UCF.

Completed UCFs may be faxed to 1-866-282-7076 or mailed to:

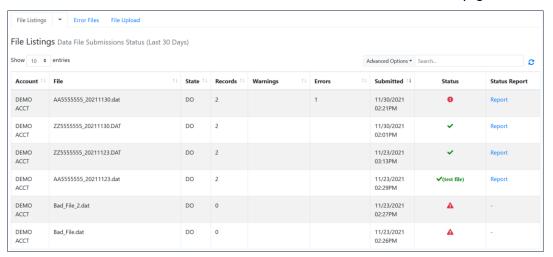
Bamboo Health Attn: Greg Hatcher 9901 Linn Station Rd., Ste. 500 Louisville, KY 40223

6 Data Compliance

This chapter describes how to view the status of your submitted data files and how to correct errors.

6.1 File Listings

The File Listings page displays information extracted from the data files submitted to PMP Clearinghouse, including the file name, number of records identified within the data file, number of records that contain warnings, number of records that contain errors, and the date and time of submission. Click **File Submissions** to access this page.

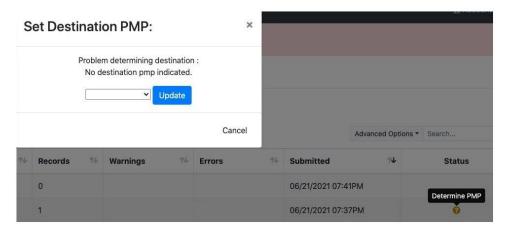


- The **Status** column, located at the end of each row, displays the file status via color-coded icon. Hovering over the icon will display the status message.
- The Status Report column, located next to the Status column, contains a link to
 the status report for that file. Please refer to <u>File Status Report</u> for more
 information on how to read and interpret this report.

If a file contains errors, it will have a • symbol with a mouse over hint of "Pending Dispensation Error" within the status column. You can click the error icon in the Status column to display the Error Correction page, which allows you to view the records containing errors (see View Records for more information). Please refer to Error Correction for instructions on how to correct errors.

If a file is unable to be parsed into the PMP Clearinghouse application, it will have an symbol with a mouse over hint of "ASAP Errors." Clicking the icon will display the detailed error, which indicates what element was missing or malformed. To correct these errors, a new file must be submitted to PMP Clearinghouse. It is not necessary to void a file that failed parsing since it was not successfully submitted to PMP Clearinghouse.

If you submitted a file via SFTP without using a PMP-specific sub-folder, the file will be displayed, and symbol will be displayed in the status column with a mouse over hint of "Determine PMP." Clicking the icon will prompt you to select a destination PMP to which the data file will be transferred.



If you submitted a zero report via file upload or SFTP that is malformed or missing information, the file will be displayed, and an exclamation mark icon inside a red triangle will be displayed in the status column. Hovering over the icon will display the "Invalid Zero Report" error. Clicking on the icon will display the detailed error message. To correct these errors, a new zero report must be submitted. Error example:



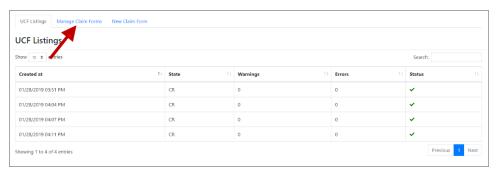
6.2 UCF Listings

The **UCF** Listings page displays information about the UCFs submitted to PMP Clearinghouse, including the number of warnings and errors. Click **UCF** Submissions to access this page.

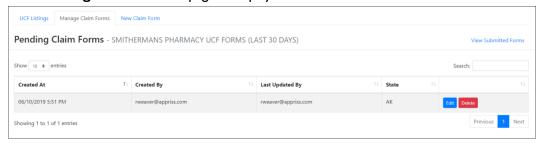


The **Status** column, located at the end of each row, displays the UCF's status. Data entered into the UCF is validated upon submission. Therefore, successfully submitted UCFs should not contain errors. However, if you have attempted to submit a UCF with errors and did not immediately correct those errors and submit the record, you have up to one (1) year to make updates to these records in Clearinghouse.

1. To view pending or incomplete submissions, click the **Manage Claim Forms** tab.



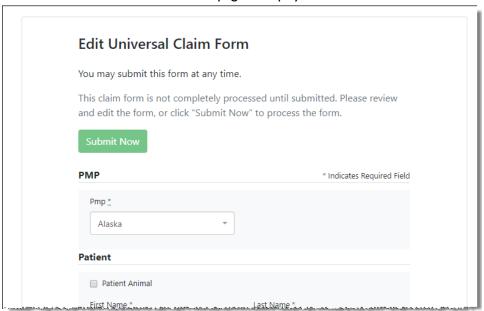
The **Pending Claim Forms** page is displayed.



2. Click **Edit** next to the form you wish to update.

Note: If it has been longer than one (1) year, the **Edit** option will no longer be available. You must click **Delete** to delete the record and start over.

The Edit Universal Claim Form page is displayed.



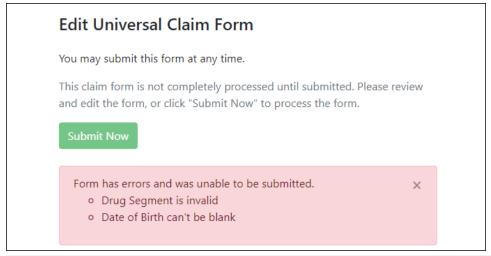
3. Make the necessary corrections or changes, and then click **Submit Now**, located at the top of the page.

A message is displayed prompting you to confirm the data submission.



4. Click OK.

Your data will be validated upon submission. If there are any remaining errors on the UCF form, they are displayed at the top of the page.



Note: If there are no errors, you are returned to the UCF Listings page and your report is listed there.

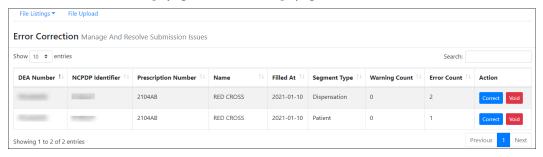
5. Correct the indicated errors, then repeat steps 3-4.

Once your data has been successfully submitted, your report is listed on the **UCF Listings** page.

6.3 Error Correction

6.3.1 View Records

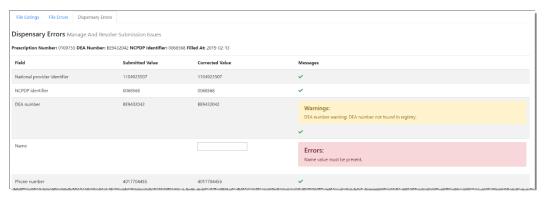
The **Error Correction** page displays more information about the records within a selected data file that need correcting, including **Prescription**Number, Segment Type, Warning Count, and Error Count. To access this page, click the "Pending Dispensation Error" message in the Status column of the File Listings page or UCF Listings page.



The **Correct** button, located at the end of each row, allows you to make corrections to the record.

6.3.2 Error Correction via PMP Clearinghouse

Once you click **Correct** on the **Error Correction** page, the **Errors** page is displayed. This page displays detailed information about the records within a selected data file that need correcting, including all the fields contained within the record and the originally submitted value, and allows you to correct those records.



- The Corrected Value column allows you to enter a new value to correct the error.
- The Message column displays the relevant error message explaining why
 the value entered in that field did not pass the validation rules.

For files that failed to parse, the error identified is "best effort" and any information we could not parse is listed as "unparseable" in the file. In this case, you must submit a corrected file.

To correct records:

- I. Identify the fields that require corrections. Fields containing errors are highlighted in red, as shown in the screenshot above.
- 2. Enter the corrected value in the **Corrected Value** column.
- 3. Click Submit.

The error is processed through the validation rules.

- a. If the changes pass the validation rules, the record is valid, and a
 message is displayed indicating that the errors have been corrected. The
 File Listings and Error Correction pages are also updated.
- b. If the changes fail the validation rules, a message is displayed indicating that there was a problem correcting the errors, and the **Message** column is updated with any new error message. Repeat steps 2–3 until the errors have been corrected and the file can be successfully submitted.

6.3.3 Error Correction via File Submission

The ASAP 4.2 standard requires a pharmacy to select an indicator in the **DSP01** (Reporting Status) field. These indicators allow you to 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 previouslysubmitted record have been revised

To revise a record:

- a. Create a record with the value "01" in the **DSP01** field.
- b. Populate the following fields with the same information originally submitted in the record that is being revised:
 - PHA03 (DEA Number)
 - **DSP02** (Prescription Number)
 - **DSP05** (Date Filled)
- c. 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.
- d. Submit the record.

Important Note: When submitting revisions for the Prescription Number (DSP02), Pharmacy DEA (PHA03), Date Filled (DSP05), Quantity Filled (DSP09), and/or Refill Number (DSP06) fields, a Void submission (02) on the original record should be processed before re-submitting a New Record (00). Submitting Revise (01) for one of these five fields will process as a new prescription and both submissions will appear. All other field revisions may be processed as 01.

- 02 Void indicates that the original record should be removed
 To void a record:
 - a. Create a record with the value "02" in the DSP01 field.
 - b. Fill in all other data identical to the original record.
 - c. Submit the record. This will void the original record.

7 Email Reports

Email status reports are automatically sent to all users associated with a specific data submitter account. These reports are used to identify errors in files that have been submitted and to confirm zero report submissions. This chapter describes the status reports you may receive via email.

7.1 File Failed Report

You will receive the File Failed Report if a submitted file was not able to be parsed and was not processed into PMP Clearinghouse. The report contains a description of the error encountered within the file. In the event of a failed file, a new file should be submitted with the necessary corrections.

Note: Failed files are not parsed into Clearinghouse and do not require a voided ASAP file to remove it from the system.

An example File Failed Report is provided below.

SUBJ: North Carolina ASAP file: fake-test3.txt - Parse Failure

BODY:
Error Message
-----Failed to decode the value '04' for the bean id 'transactionControlType'.

Summary:

- * File Name: fake-test3.txt
- * ASAP Version: 4.2
- * Transaction Control Number: unparseable
- * Transaction Control Type: unparseable
- * Date of Submission: September 19, 2022

NOTE: This file could not be received into the system because the system could not recognize its content as a valid ASAP format. Action is required to resolve the issues and a subsequent file should be submitted. As such the information provided in this report is "best effort" and any information we could not parse is listed as "unparseable" in the fields above.

7.2 File Status Report

The File Status Report serves as notification that a data file is currently being parsed by the PMP system.

This report identifies specific records in the submitted data file and returns identifying information about the record, including specific errors identified during the validation process. It uses fixed-width columns and contains a summary section after the error listings. Each column contains a blank two-digit pad at the end of the data.

The columns are set to the following lengths:

Column	Length
DEA	II (9 + pad)
NCPDP	9 (7 + pad)
NPI	12 (10 + pad)
Prescription	27 (25 + pad)
Filled	10 (8 + pad)
Segment	18 (16 + pad)
Field	18 (16 + pad)
Туре	9 (7 + pad)
Message	Arbitrary

The File Status Report notifies you of the following scenarios:

- Total records: The total number of records contained in the submitted data file.
- Duplicate records: The number of records that were identified as already existing
 within the PMP system. Duplicate records are not imported to prevent improper
 patient information.
- Records in process: The number of records remaining to be processed into the system (usually only displays a number if the file has not finished loading at the time the report is sent out).

Note: Records remaining to be processed will continue to be processed even after the status report is sent.

- Records with errors: The number of records that contain errors. These errors
 must be corrected for the record to be imported into the system. If a zero (0) is
 displayed, there are no errors in the data. Please refer to Error Correction for
 instructions on correcting errors.
- **Records with warnings**: The number of records that contain warnings. These warnings do not need to be corrected for the record to be imported into the system. If a zero (0) is displayed, there are no warnings in the data.
- Records imported with warnings: The number of records with warnings that
 were imported. If a record contains both warnings and errors, the errors must be
 corrected to be submitted to the system. Please refer to Error Correction for
 instructions on correcting errors.
- **Records imported without warnings**: The number of records without warnings that were imported.

Note: The initial File Status Report is sent out two (2) hours after the file has been submitted to the system. Additional reports will be sent out every 24 hours if errors continue to be identified within a submitted data file.

An example File Status Report is provided on the following page.

SUBJ: North	SUBJ: North Carolina ASAP file: fake-test3.txt - Status Report							
BODY: DEA	NCPDP	NPI	Prescription	Filled	Segment	Field	Type	Message
DEA								
BE1234567	1347347	9034618394	123486379596-0	20220918	Dispensation	refill_number	WARNING	message example
DE9841394	3491849	4851947597	357199504833-345		Dispensation	days_supply	ERROR	message example
Summary:								
* File Name:	: fake-test	3.txt						
* ASAP Vers	ion: 4.2							
* Transactio	n Control	Number: 234	89504823					
* Transaction Control Type: send								
* Date of Submission: September 19, 2022								
* Total Reco	* Total Record Count: ###							
* Duplicate	* Duplicate Records: ###							
* In Process	* In Process Count: ###							
* Records w								
* Imported I	Records C	ount: ###						
* Records In	nported w	rith Warning (Count: ###					

7.3 Zero Report Confirmation

You will receive a Zero Report Confirmation after successfully submitting a zero report to PMP Clearinghouse. This report displays the PMP to which the zero report was submitted, date range for the zero report, date the zero report was submitted to PMP Clearinghouse, and date the report was originally created.

An example Zero Report Confirmation is provided below.

SUBJ: ASAP Zero Report: zero_reports_20220306KSMCPS.DAT

BODY:

Summary:

* File Name: zero_reports_20220306KSMCPS.DAT

* PMP Name: North Carolina

* Date Range: 2022-03-06 - 2022-03-06

* Submission Date: 2013-03-07 * ASAP Creation Date: 2022-03-07

8 Managing Your Upload Account

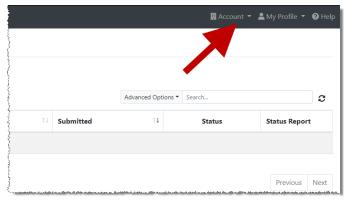
The **Account** menu option allows you to manage the information associated with your organization's upload account, including adding users, PMPs, and SFTP access to your account as well as editing your organization's account information.

Note: This chapter contains information for managing the upload account with which your user account is associated. For information about editing and managing your individual user account, including how to change your password, please refer to <u>Managing Your User Profile</u>.

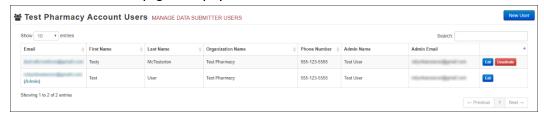
8.1 Adding Users to Your Upload Account

PMP Clearinghouse allows data submitters to add new users to the system who have the same rights and access to submitting data and viewing file status. This practice allows you to create an account to be used for a backup individual.

- I. Log in to PMP Clearinghouse.
- 2. Click Account.

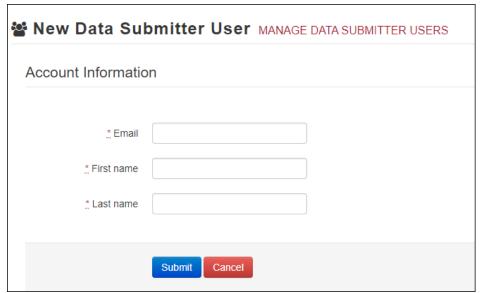


3. Select **Users** from the **Account** drop-down menu. The **Account Users** page is displayed.



4. Click **New User**, located in the top right corner of the page.

The **New Data Submitter User** page is displayed.



5. Enter the new data submitter's email address, first name, and last name in the appropriate fields.

Note: All fields are required.

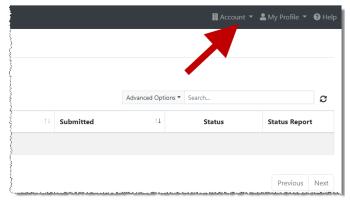
6. Click Submit.

The user is added to the list of data submitters for your organization, and you are returned to the **Account Users** page.

- 7. Please inform the new user of the account creation.
 - a. The user will receive an email with a link for them to confirm their account.
 - b. Once the account has been confirmed, the user will need to navigate to the PMP Clearinghouse Login page and click **Forgot your password?** to create a password for their account and log in.
 - c. Upon logging in, the user will be able to view all files submitted for your organization's upload account.

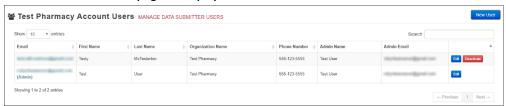
8.1.1 Changing Another User's Password

- I. Log in to PMP Clearinghouse.
- 2. Click Account.



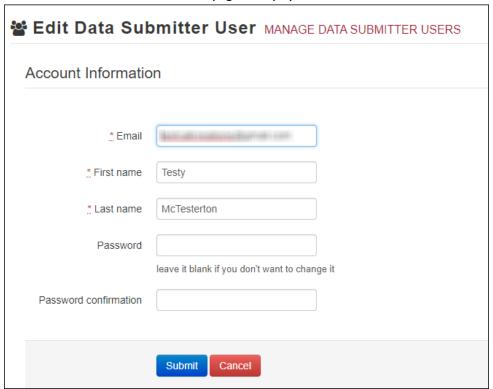
3. Select **Users** from the **Account** drop-down menu.

The **Account Users** page is displayed.



4. Click the **Edit** button, located to the right of the user's information.

The Edit Data Submitter User page is displayed.



5. Enter a new password for the user in the **Password** field, then re-enter it in the **Password confirmation** field. The password requirements are provided below.

Passwords must contain:

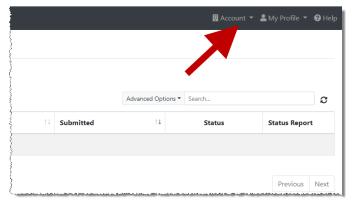
- At least eight (8) characters
- One (1) uppercase letter
- One (1) lowercase letter
- One (1) number
 - One (1) special character, such as !, @, #, \$, etc.
- 6. Click **Submit**.

The password is changed.

8.2 Adding PMPs to Your Upload Account

If your organization needs to submit data files to an additional PMP that uses PMP AWARXE, you can submit the request through PMP Clearinghouse.

- I. Log in to PMP Clearinghouse.
- 2. Click Account.



3. Select Multi State Approval from the Account drop-down menu.

The **Multi State Approval** page is displayed. This page displays all PMPs currently using the PMP AWARxE system as well as your data sharing status with each PMP.



4. To request to submit data to another PMP, click to select the checkbox next to that PMP.

PMP Clearinghouse automatically saves your changes, and your request is submitted to the PMP administrator for review and approval. Once the request has been approved, the status for that PMP will change from "Pending" to "Approved," and you may begin submitting data to that PMP.

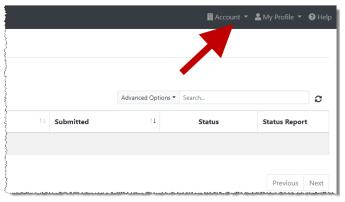
Notes:

- If you are submitting data via SFTP, the file must be located in the proper subfolder to ensure delivery to the desired PMP.
- To cancel data submission to a PMP, uncheck the box for that PMP. Note that if you
 need to submit data to that PMP again in the future, you will have to go through the
 approval process again.

8.3 Adding SFTP Access to an Upload Account

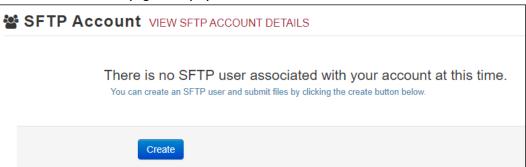
If a registered upload account did not request an SFTP account during the account creation process, you can request one at any time using the **Account** menu option.

- I. Log in to PMP Clearinghouse.
- 2. Click Account.



3. Select SFTP Details.

The SFTP Account page is displayed.



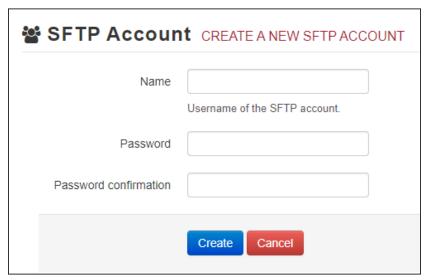
Note: If an SFTP account already exists for the upload account, the username is displayed on the SFTP Account page.



You cannot change the SFTP account username. However, you can update the password by clicking **Edit**.

4. Click Create.

The Create a New SFTP Account page is displayed.



5. Enter a username for the account in the **Name** field.

Notes:

- The username must contain a minimum of eight (8) characters.
- Once the SFTP account has been created, you cannot change the username.
- 6. Enter a password for the account in the **Password** field, then re-enter it in the **Password confirmation** field. The password requirements are provided below.

Passwords must contain:

- At least eight (8) characters
- One (1) uppercase letter
- One (1) lowercase letter
- One (1) number
- One (1) special character, such as !, @, #, \$, etc.

Once the account has been successfully created, this password will be input into the pharmacy software so that submissions can be automated.

Notes:

- This password can be the same as the one used when the upload account was created.
- Unlike your Profile password (i.e., your user account password), the SFTP password does not expire.
- The URL to connect via SFTP is <u>submissions.healthcarecoordination.net/</u>.
- Additional details on SFTP configuration can be found in <u>Appendix C: SFTP</u> <u>Configuration</u>.
- 7. Click Create.

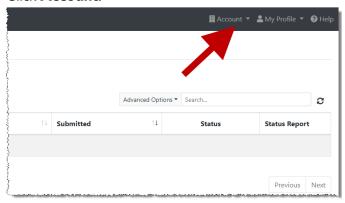
The account is created, and the username is displayed.



8.4 Editing Your Upload Account

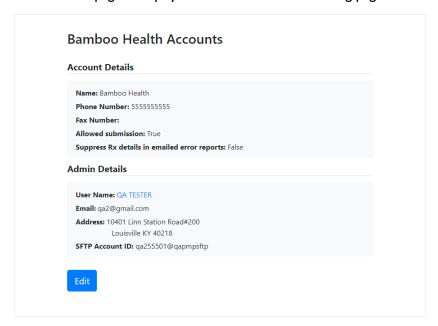
Note: This function only allows you to edit your organization's upload account. If you need to edit your individual profile information, please refer to <u>Editing Your Profile</u>.

- I. Log in to PMP Clearinghouse.
- 2. Click Account.



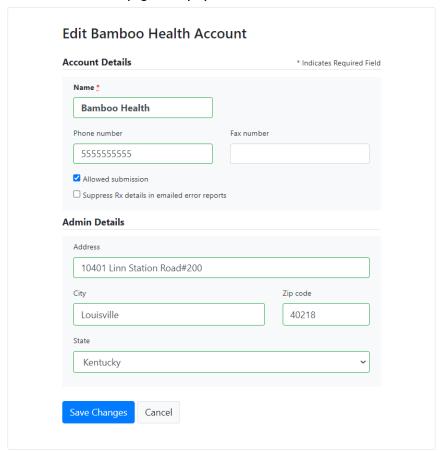
3. Select Account Details.

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



4. Click Edit.

The Edit Account page is displayed.



5. Update the information as necessary, then click **Submit**. The account information is updated.

9 Managing Your User Profile

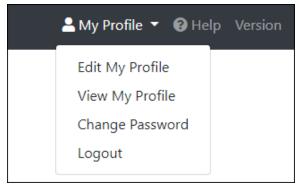
This chapter describes how to manage your individual user profile, including how to edit your profile and manage your password.

Note: This chapter contains information for managing your individual user profile. For information about managing your organization's upload account, including how to add users, please refer to <u>Managing Your Upload Account</u>.

9.1 Editing Your Profile

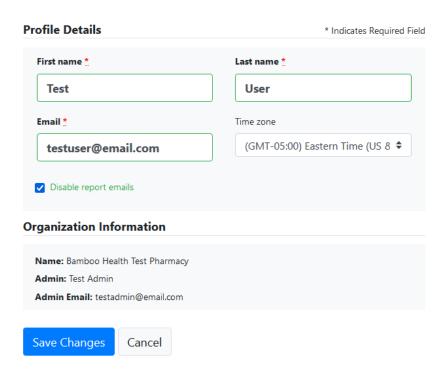
Note: This function only allows you to edit your individual profile information. If you need to edit the Organization Information, please refer to <u>Editing Your Upload Account</u>.

- I. Log in to PMP Clearinghouse.
- 2. Click My Profile.



3. Select Edit My Profile.

Edit Profile

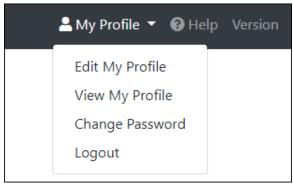


4. Update your information as necessary, then click **Submit**. Your changes are saved, and your updated profile is displayed.

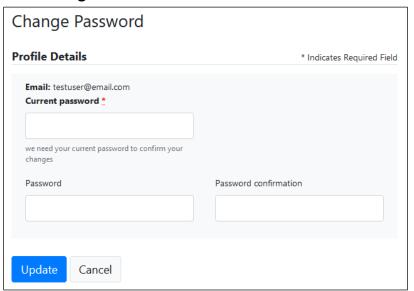
9.2 Changing Your Password

Note: Clearinghouse passwords expire every 90 days. You can use this function to proactively change your password before it expires. If your password has already expired, or you have forgotten your password, navigate to the PMP Clearinghouse Login page and click **Forgot your password?** to reset it. Please refer to <u>Resetting Your Password</u> for more information.

- I. Log in to PMP Clearinghouse.
- 2. Click My Profile.



3. Select Change Password.



- 4. Enter your current password in the **Current Password** field.
- 5. Enter your new password in the **Password** field, then re-enter it in the **Password confirmation** field. The password requirements are provided below.

Passwords must contain:

- At least eight (8) characters
- One (1) uppercase letter
- One (1) lowercase letter

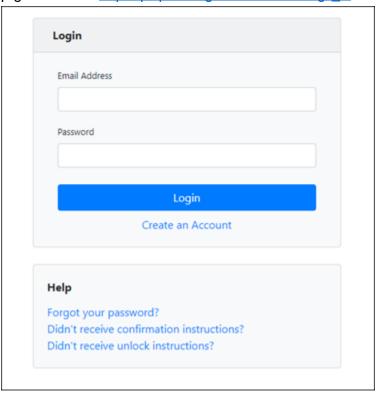
- One (1) number
- One (1) special character, such as !, @, #, \$, etc.
- 6. Click **Update**.

Your password is updated, and you will use it the next time you log in to PMP Clearinghouse.

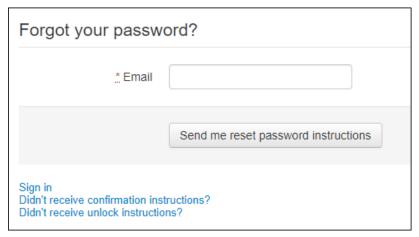
9.3 Resetting Your Password

If you have forgotten your password or your password has expired, perform the following steps to reset it.

I. Open an internet browser window and navigate to the PMP Clearinghouse Login page located at https://pmpclearinghouse.net/users/sign_in.

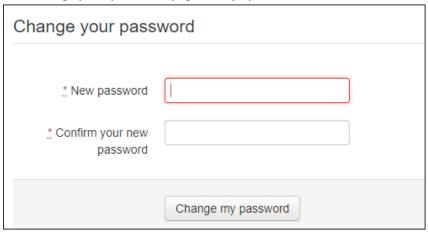


2. Click the **Forgot your password?** link, located in the Help section of the page. The Forgot your password page is displayed.



- 3. Enter the email address associated with your user account, then click **Send me** reset password instructions.
- 4. Once you receive the reset password email, click the **Change my password** link within the email.

The Change your password page is displayed.



5. Enter your new password in the **New password** field, then re-enter it in the **Confirm your new password** field. The password requirements are provided below.

Passwords must contain:

- At least eight (8) characters
- One (1) uppercase letter
- One (1) lowercase letter
- One (1) number
- One (1) special character, such as !, @, #, \$, etc.
- 6. Click Change my password.

Your password is changed, and you can now use it to log in to PMP Clearinghouse.

10 Assistance and Support

10.1 Technical Assistance

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

Contact Bamboo Health at 1-855-962-4767;
 OR

 Create a support request at the following URL: https://pmpclearinghouse.zendesk.com/hc/en-us/

Technical assistance is available 24 hours per day, 7 days per week.

10.2 Administrative Assistance

If you have non-technical questions regarding the NC CSRS, please contact:

North Carolina Controlled Substances Reporting System
Division of Mental Health, Developmental Disabilities, and Substance Abuse Services
3008 Mail Service Center
Raleigh, North Carolina 27699-3008

Phone: 919-733-1765

E-mail: nccsrs@dhhs.nc.gov

II Document Information

II.I Disclaimer

Bamboo Health has made every effort to ensure the accuracy of the information in this document at the time of printing. However, information is subject to change.

11.2 Change Log

Version	Date	Chapter/Section	Change Made
Draft Vers	ions		
DRAFT vI	06/19/2018	N/A	N/A; initial DRAFT publication
DRAFT v2	07/17/2018	Global	Response to State revisions
DRAFT v3	07/26/2018	Global	Response to State revisions
Published \	Versions		
1.0	12/18/2018	Global	Finalized for publication
1.1	06/19/2019	Cover page	Updated address in footer
		10.1/Technical Assistance	Updated phone number
1.2	09/30/2019	Global	Updated screenshots to reflect updated user interface (note that this is only a cosmetic change; no functionality changes are included)
		2.2/Reporting Requirements	Added a note specifying that the owner's date of birth should be entered for veterinary prescriptions
		6.2/UCF Listings	Added clarification on correcting UCF errors
1.3	03/11/2020	4.3/Zero Reports	Separated into two sections (Submit a Single-Click Zero Report and Create a New Zero Report) to reflect the addition of the single-click zero report submission functionality
		4.3.1/Submit a Single- Click Zero Report	Added new section with instructions for submitting a single-click zero report
2.0	03/18/2022	Global	Updated guide to reflect Bamboo Health branding
3.0	04/03/2024	Global	Updated sFTP hostname information
		6.2/UCF Listings	Updated UCF time to edit/delete from 30 days to 1 year
		2/Data Collection and Tracking	Updated guide per state's request to include upcoming legislative changes related to gabapentin submissions

3.1	04/17/2024	Appendix A/ASAP 4.2 Specifications	Added notes to PRE01 and PRE02 fields on Gabapentin reporting
		Appendix D/Reporting Gabapentin Prescriptions	Added new appendix
4.0	12/04/2024	2.1 Data Collection Overview	Updated language to reflect HB 190 SL2023-65 veterinarian effective date
		General	Updated guide to reflect new branding guidelines
		6.3.3/Error Correction via File Submission	Added additional guidance on voiding and correcting files.

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 the NC CSRS 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 (~).
- **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 (~~).

Requirement

- R = Required by NC CSRS
- N = Not required but accepted if submitted
- S = Situational (not required; however, supply if available)
- P = Preferred, please submit

Note: For more information, contact the American Society for Automation in Pharmacy for the full Implementation Guide for the ASAP Standard for Prescription-Monitoring Programs. That guide includes field lengths, acceptable attributes, and examples.

Segment	Element	Element Name	Requirement
	cate the start	er (required) of a transaction. It also assigns the data element separator, segment ter	minator, and
	TH01	Version/Release Number	R
		Code uniquely identifying the transaction.	
		Format = x.xx	
	TH02	Transaction Control Number	R
		Sender assigned code uniquely identifying a transaction.	
	TH03	Transaction Type	S
		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	,	S
	1 1104	Response ID Contains the Transaction Control Number of a transaction that	3
		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 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.	
IC. Informa	<u> </u>		
	tion Source	and identification numbers of the entity supplying the information.	
Osed to con-	T .		
	IS01	Unique Information Source ID Reference number or identification number.	R
		(Example: phone number)	
	1502	+	D
	IS02	Information Source Entity Name	R
	1000	Entity name of the Information Source.	•
	IS03	Message	S
		Free-form text message.	

Segment	Element ID	Element Name	Requirement
PHA: Phari	macy Heade	er (required)	
Used to ider	ntify the pharn	nacy.	
Note: It is r PHA03.	equired that i	nformation be provided in at least one of the following fields: PHA01, I	PHA02, or
	PHA01	National Provider Identifier (NPI)	S
		Identifier assigned to the pharmacy by CMS.	(Required if pharmacy does not have a DEA)
	PHA02	NCPDP/NABP Provider ID	S
		Identifier assigned to pharmacy by the National Council for Prescription Drug Programs.	
	PHA03	DEA Number	R
		Identifier assigned to the pharmacy by the Drug Enforcement Administration.	
	PHA04	Pharmacy Name	S
		Free-form name of the pharmacy or dispensing practitioner's name.	
	PHA05	Address Information - I	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
	D11400	U.S. Postal Service state or other regional jurisdiction code.	
	PHA09	ZIP Code Address U.S. Postal Service ZIP Code.	S
	PHAI0	Phone Number	S
		Complete phone number including area code.	
	PHAII	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.	
		ion (required)	
Used to repo	ort the patien	t's name and basic information as contained in the pharmacy record.	
	PAT01	ID Qualifier of Patient Identifier	S
		Code identifying the jurisdiction that issues the ID in PAT03.	

Segment	Element ID	Element Name	Requirement
	PAT02	ID Qualifier Code to identify the type of ID in PAT03. If PAT02 is used, PAT03 is required. 0 I Military ID 0 2 State Issued ID 0 3 Unique System ID 0 4 Permanent Resident Card (Green Card) 0 5 Passport ID 0 6 Driver's License ID 0 8 Tribal ID	S
	PAT03	 99 Other (agreed upon ID) ID of Patient Identification number for the patient as indicated in PAT02. An example would be the driver's license number. 	S
	PAT04	ID Qualifier of Additional Patient Identifier Code identifying the jurisdiction that issues the ID in PAT06. Used if the PMP requires such identification.	S
	PAT05	Additional Patient ID Qualifier Code to identify the type of ID in PAT06 if the PMP requires a second identifier. If PAT05 is used, PAT06 is required. OI Military ID O2 State Issued ID O3 Unique System ID O4 Permanent Resident Card O5 Passport ID O6 Driver's License ID O8 Tribal ID 99 Other (agreed upon ID)	S
	PAT06	Additional ID 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.	S
	PAT07	Last Name Patient's last name.	R
	PAT08	First Name Patient's first name.	R
	PAT09	Middle Name Patient's middle name or initial if available.	S
	PATI0	Name Prefix Patient's name prefix such as Mr. or Dr.	S
	PATII	Name Suffix Patient's name suffix such as Jr. or the III.	S

Segment	Element ID	Element Name	Requirement
	PAT12	Address Information - I	R
		Free-form text for street address information.	
	PAT13	Address Information – 2	S
		Free-form text for additional address information.	
	PATI4	City Address	R
		Free-form text for city name.	
	PAT15	State Address	R
		U.S. Postal Service state or other regional jurisdiction code	
	PATI6	ZIP Code Address	R
		U.S. Postal Service ZIP code.	
		Note: Populate with zeros if patient address is outside the U.S.	
	PATI7	Phone Number	R
		Complete phone number including area code.	
	PAT18	Date of Birth	R
		Date patient was born.	
		Format: CCYYMMDD	
	PAT19	Gender Code	S
		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	
	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	
		II Hospice	
		98 Unknown	
		• 99 Other	

Segment	Element	Element Name	Requirement
	ID		
	PAT22	Country of Non-U.S. Resident	S
		Used when the patient's address is a foreign country.	
	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.	
-	•	rd (required)	
Used to ider quantity.	ntify the basic	components of a dispensing of a given prescription order including the	date and
	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) 	
		OI 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	Prescription Number	R
		Serial number assigned to the prescription by the pharmacy.	
	DSP03	Date Written	R
		Date the prescription was written (authorized).	
		Format: CCYYMMDD	
	DSP04	Refills Authorized	R
		The number of refills authorized by the prescriber.	
	DSP05	Date Filled	R
		Date prescription was filled.	
		Format: CCYYMMDD	
	DSP06	Refill Number	R
		Number of the fill of the prescription.	
		0 indicates New Rx fill; 01-99 identifies any additional fills.	
	DSP07	Product ID Qualifier	R
		Used to identify the type of product ID contained in DSP08.	
		• 01 NDC	
		 06 Compound (indicates a compound; if used, the CDI segment becomes a required segment) 	
	DSP08	Product ID	R
		Full product identification as indicated in DSP07, including leading zeros without punctuation.	
	DSP09	Quantity Dispensed	R
	30.07	Number of metric units dispensed in metric decimal format.	
		Example: 2.5	
		Note: For compounds show the first quantity in CDI04.	

Segment	Element	Element Name	Requirement
J	ID		
	DSP10	Days' Supply	R
		Estimated number of days the medication will last.	
	DSPII	Drug Dosage Units Code	R
		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 Prescription	
		02 Telephone Prescription	
		03 Telephone Emergency Prescription	
		04 Fax Prescription	
		05 Electronic Prescription	
		• 99 Other	
	DSP13	Partial Fill Indicator	S
		Used when the quantity in DSP 09 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 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.	
	DSP15	Pharmacist State License Number	S
		This data element can be used to identify the pharmacist dispensing the medication.	
		Assigned to the pharmacist by the Licensing Board.	
	DSP16	Classification Code for Payment Type	R
		Code identifying the type of payment (i.e., how it was paid for).	
		OI Private Pay	
		02 Medicaid	
		03 Medicare	
		04 Commercial Insurance	
		05 Military Installations and VA	
		 06 Workers' Compensation 	
		07 Indian Nations	
		• 99 Other	

egment	Element	Element Name	Requiremen
	ID		
	DSP17	Date Sold	S
		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.	_
	DSP18	RxNorm Product Qualifier	S
		01 Semantic Clinical Drug (SCD)	
		02 Semantic Branded Drug (SBD)	
		03 Generic Package (GPCK)	
		04 Branded Package (BPCK)	
	DSP19	RxNorm Code	S
		Used for electronic prescriptions to capture the prescribed drug product identification.	
	DSP20	Electronic Prescription Reference Number	S
		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.	
	DSP21	Electronic Prescription Order Number	S
		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.	
		one, if not both, it decides to capture. nation (required)	
	ntify the preso	one, if not both, it decides to capture. nation (required) criber of the prescription.	
		one, if not both, it decides to capture. nation (required) criber of the prescription. National Provider Identifier (NPI)	S
	ntify the preso	one, if not both, it decides to capture. nation (required) criber of the prescription. National Provider Identifier (NPI) Identifier assigned to the prescriber by CMS.	S
	ntify the preso	one, if not both, it decides to capture. nation (required) criber of the prescription. National Provider Identifier (NPI)	S
	PRE01	one, if not both, it decides to capture. nation (required) criber of the prescription. National Provider Identifier (NPI) Identifier assigned to the prescriber by CMS. Note: See Appendix D: Reporting Gabapentin Prescriptions for guidance on veterinarians.	
	ntify the preso	one, if not both, it decides to capture. nation (required) criber of the prescription. National Provider Identifier (NPI) Identifier assigned to the prescriber by CMS. Note: See Appendix D: Reporting Gabapentin Prescriptions for guidance on veterinarians. DEA Number	S R
	PRE01	one, if not both, it decides to capture. nation (required) criber of the prescription. National Provider Identifier (NPI) Identifier assigned to the prescriber by CMS. Note: See Appendix D: Reporting Gabapentin Prescriptions for guidance on veterinarians.	
	PRE01	one, if not both, it decides to capture. nation (required) criber of the prescription. National Provider Identifier (NPI) Identifier assigned to the prescriber by CMS. Note: See Appendix D: Reporting Gabapentin Prescriptions for guidance on veterinarians. DEA Number Identifying number assigned to a prescriber or an institution by the	
	PRE01	one, if not both, it decides to capture. nation (required) criber of the prescription. National Provider Identifier (NPI) Identifier assigned to the prescriber by CMS. Note: See Appendix D: Reporting Gabapentin Prescriptions for guidance on veterinarians. DEA Number Identifying number assigned to a prescriber or an institution by the Drug Enforcement Administration (DEA). Note: For Gabapentin, see Appendix D: Reporting Gabapentin Prescriptions	
	PRE01	one, if not both, it decides to capture. nation (required) criber of the prescription. National Provider Identifier (NPI) Identifier assigned to the prescriber by CMS. Note: See Appendix D: Reporting Gabapentin Prescriptions for guidance on veterinarians. DEA Number Identifying number assigned to a prescriber or an institution by the Drug Enforcement Administration (DEA). Note: For Gabapentin, see Appendix D: Reporting Gabapentin Prescriptions for guidance.	
	PRE01	one, if not both, it decides to capture. nation (required) criber of the prescription. National Provider Identifier (NPI) Identifier assigned to the prescriber by CMS. Note: See Appendix D: Reporting Gabapentin Prescriptions for guidance on veterinarians. DEA Number Identifying number assigned to a prescriber or an institution by the Drug Enforcement Administration (DEA). Note: For Gabapentin, see Appendix D: Reporting Gabapentin Prescriptions	
	PRE01	one, if not both, it decides to capture. nation (required) criber of the prescription. National Provider Identifier (NPI) Identifier assigned to the prescriber by CMS. Note: See Appendix D: Reporting Gabapentin Prescriptions for guidance on veterinarians. DEA Number Identifying number assigned to a prescriber or an institution by the Drug Enforcement Administration (DEA). Note: For Gabapentin, see Appendix D: Reporting Gabapentin Prescriptions for guidance. Do not enter XDEA in this field. XDEA number should now be	
	PRE01 PRE02	one, if not both, it decides to capture. nation (required) criber of the prescription. National Provider Identifier (NPI) Identifier assigned to the prescriber by CMS. Note: See Appendix D: Reporting Gabapentin Prescriptions for guidance on veterinarians. DEA Number Identifying number assigned to a prescriber or an institution by the Drug Enforcement Administration (DEA). Note: For Gabapentin, see Appendix D: Reporting Gabapentin Prescriptions for guidance. Do not enter XDEA in this field. XDEA number should now be reported in PRE09.	R
	PRE01 PRE02	one, if not both, it decides to capture. nation (required) criber of the prescription. National Provider Identifier (NPI) Identifier assigned to the prescriber by CMS. Note: See Appendix D: Reporting Gabapentin Prescriptions for guidance on veterinarians. DEA Number Identifying number assigned to a prescriber or an institution by the Drug Enforcement Administration (DEA). Note: For Gabapentin, see Appendix D: Reporting Gabapentin Prescriptions for guidance. Do not enter XDEA in this field. XDEA number should now be reported in PRE09. DEA Number Suffix Identifying number assigned to a prescriber by an institution when	R
	PRE02 PRE03	one, if not both, it decides to capture. nation (required) criber of the prescription. National Provider Identifier (NPI) Identifier assigned to the prescriber by CMS. Note: See Appendix D: Reporting Gabapentin Prescriptions for guidance on veterinarians. DEA Number Identifying number assigned to a prescriber or an institution by the Drug Enforcement Administration (DEA). Note: For Gabapentin, see Appendix D: Reporting Gabapentin Prescriptions for guidance. Do not enter XDEA in this field. XDEA number should now be reported in PRE09. DEA Number Suffix Identifying number assigned to a prescriber by an institution when the institution's number is used as the DEA number.	R
	PRE02 PRE03	one, if not both, it decides to capture. nation (required) criber of the prescription. National Provider Identifier (NPI) Identifier assigned to the prescriber by CMS. Note: See Appendix D: Reporting Gabapentin Prescriptions for guidance on veterinarians. DEA Number Identifying number assigned to a prescriber or an institution by the Drug Enforcement Administration (DEA). Note: For Gabapentin, see Appendix D: Reporting Gabapentin Prescriptions for guidance. Do not enter XDEA in this field. XDEA number should now be reported in PRE09. DEA Number Suffix Identifying number assigned to a prescriber by an institution when the institution's number is used as the DEA number. Prescriber State License Number	R
	PRE02 PRE03	one, if not both, it decides to capture. nation (required) criber of the prescription. National Provider Identifier (NPI) Identifier assigned to the prescriber by CMS. Note: See Appendix D: Reporting Gabapentin Prescriptions for guidance on veterinarians. DEA Number Identifying number assigned to a prescriber or an institution by the Drug Enforcement Administration (DEA). Note: For Gabapentin, see Appendix D: Reporting Gabapentin Prescriptions for guidance. Do not enter XDEA in this field. XDEA number should now be reported in PRE09. DEA Number Suffix Identifying number assigned to a prescriber by an institution when the institution's number is used as the DEA number. Prescriber State License Number Identification assigned to the prescriber by the Licensing Board. Note: See Appendix D: Reporting Gabapentin Prescriptions for guidance	R

Segment	Element ID	Element Name	Requirement
	PRE06	First Name	S
		Prescriber's first name.	
	PRE07	Middle Name	S
		Prescriber's middle name or initial.	
	PRE08	Phone Number	S

CDI: Compound Drug Ingredient Detail (situational)

Use of this segment is 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.

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

CDI01	Compound Drug Ingredient Sequence Number	R
	First reportable ingredient is 1; each additional reportable ingredient is incremented by 1.	
CDI02	Product ID Qualifier Code to identify the type of product ID contained in CDI03. 01 NDC 02 UPC 03 HRI	R
CDI03	 04 UPN 05 DIN 06 Compound (this code is not used in this segment) Product ID Full 11-digit NDC number as indicated in CDI02, created by adding 	R
	a leading zero to the appropriate segment to result in a 5-4-2 formatted NDC number and without punctuation.	
CDI04	Compound Ingredient Quantity Metric decimal quantity of the ingredient identified in CDI03. Example: 2.5	R
CD105	 Compound Drug Dosage Units Code 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) 	S

AIR: Additional Information Reporting (situational)

Used when serialized Rx pads are used, the PMP 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.

All	IR0 I	State Issuing Rx Serial Number	S
		U.S.P.S. state or other regional jurisdiction code that issued serialized prescription blank. This is required if AIR02 is used.	

Segment	Element	Element Name	Requirement
	ID		
	AIR02	State Issued Rx Serial Number	S
		Number assigned to 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	
		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.	
	AIR06	Relationship of Person Dropping Off or Picking Up Rx	S
		Code indicating the relationship of the person.	
		OI 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.	
	AIRII	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.	

Segment	Element ID	Element Name	Requirement
TP: Pharma	acy Trailer (ı	required)	
		data for a given pharmacy and provide the count of the total number pharmacy, including the PHA and TP segment.	of detail
	TP01	Number of detail segments included for the pharmacy including the pharmacy header (PHA) and the pharmacy trailer (TP) segments.	R
	ction Trailer cate the end o	r (required) f the transaction and provide the count of the total number of segmen	ts included in the
	ТТОІ	Transaction Control Number Identifying control number that must be unique. Assigned by the originator of the transaction. Must match the number in TH02.	R
	TT02	Segment Count Total number of segments included in the transaction including the header and trailer segments.	R

Appendix B: ASAP Zero Report Specifications

The following table contains the required definitions for submitting zero reports via SFTP or manual upload to the NC CSRS. It lists the **Segment** and **Element ID** with pre-populated data to be used as an example for constructing a zero report. For more details regarding these Segment or Elements IDs, or for details on reporting actual dispensations, please refer to Appendix A: ASAP 4.2 Specifications.

Segment	Element ID	Element Name	Requirement				
TH: Transaction Header (required)							
	TH01	4.2	R				
	TH02	123456	R				
	TH05	20220401	R				
	TH06	223000	R				
	TH07	P	R				
	TH09	11	R				
IS: Information Source (required)							
	IS01	770555555	R				
	IS02	PHARMACY NAME	R				
	IS03	Date Range of Report #YYYYMMDD#-#YYYYMMDD#	R				
PHA: Phar	macy Header (re	equired)					
	PHA03	ZZ1234567	R				
PAT: Patie	PAT: Patient Information (required)						
	PAT07	REPORT	R				
	PAT08	ZERO	R				
DSP: Dispe	DSP: Dispensing Record (required)						
	DSP05	20220401	R				
PRE: Presc	riber Informatio	n (required; can be null as follows: PRE******\)					
CDI: Comp	ound Drug Ingr	edient Detail					
AIR: Additional Information Reporting							
TP: Pharmacy Trailer (required)							
	TP01	7	R				
TT: Transa	TT: Transaction Trailer (required)						
	TT01	123456	R				
	TT02	10	R				

Sample Zero Report

The following example illustrates a zero report using the above values.

```
TH*4.2*123456*01**20220108*223000*P**\\
IS*77055555555*PHARMACY NAME*#20220101#-#20220107#\
PHA*** ZZ1234567\
PAT******REPORT*ZERO********\
DSP*****20220108******\
PRE*\
CDI*\
AIR*\
TP*7\
TT*123456*10\
```

Appendix C: SFTP Configuration

This appendix describes the SFTP configurations required to upload your data to PMP Clearinghouse.

Note: Submitting data via SFTP requires that you have an existing PMP Clearinghouse account with SFTP access.

- If you need to create a PMP Clearinghouse account, please refer to <u>Creating Your Account</u>. You will be able to set up your SFTP account during the account creation process.
- If you have an existing PMP Clearinghouse account but do not have SFTP access, please refer to Adding SFTP Access to an Upload Account.

SFTP Connection Details

Hostname: submissions.healthcarecoordination.net/

Bamboo Health recommends that you use the hostname when configuring the connection rather than the IP address, as the IP address is subject to change.

Port: 22

Note: The port will always be 22.

 Credentials: Your SFTP account credentials (username and password) can be found within the PMP Clearinghouse website. To locate your credentials, <u>log in to PMP Clearinghouse</u>, then click Account > SFTP Details > Edit.

Your username cannot be modified; however, you can update your password.

Note: Your current SFTP password cannot be seen or recovered. If you have forgotten or lost it, you will need to create a new one. For more information on changing the SFTP password, please refer to <u>Adding SFTP Access to an Upload Account</u>.

Once you have established SFTP access, you can test the SFTP connection, but you will not be able to submit data to a PMP until your account has been approved by the PMP administrator.

PMP Subfolders

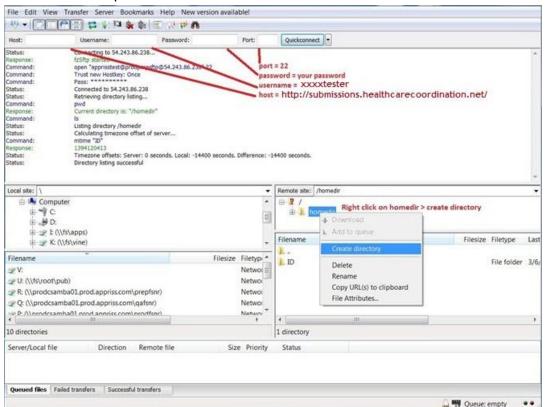
PMP Clearinghouse is the data repository for numerous PMPs. As such, data submitted via SFTP must be placed in the appropriate folder for the PMP for which you are submitting data so that it can be properly imported to that PMP. The creation of subfolders must be done outside of the PMP Clearinghouse website using third-party software, such as an SSH client or a command line utility. Files placed in the root/home directory of the SFTP server will not be imported, as this will cause the dispensing entity to appear as noncompliant/delinquent.

Your pharmacy software will need to be configured to place files in the appropriate PMP folder when submitting. You may need to contact your software vendor for additional assistance with this process.

NOTE: Capitalization of the abbreviated PMP folders' names has no bearing on whether or not Clearinghouse processes the files; however, some pharmacy systems, especially *nix-based systems, will require that the exact case is used when specifying the target folder.

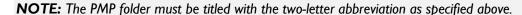
There are two methods by which to create PMP subfolders for SFTP submissions:

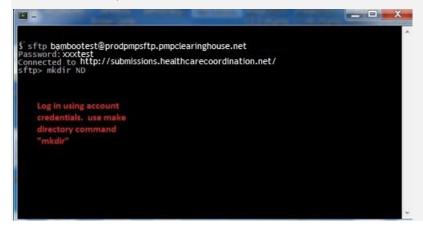
- 1. **Via SSH client** (e.g., WinSCP, FileZilla, etc.)
 - a. Log in to your SFTP account.
 - b. Create the required directories under /homedir.



2. Via command prompt

- a. Log in to your SFTP account using command prompt.
- b. Type "**mkdir**" followed by a space and then the PMP abbreviation you are using (e.g., **mkdir NC**).



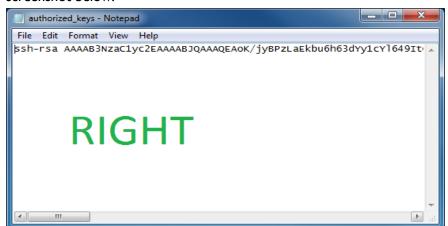


Public (SSH/RSA) Key Authentication

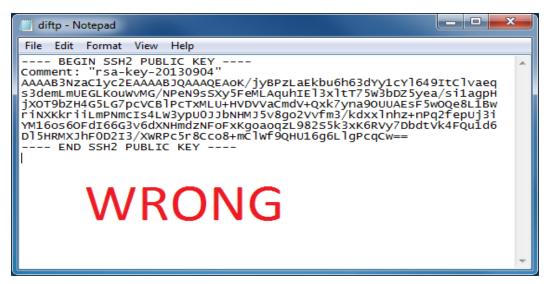
PMP Clearinghouse supports SSH key authentication. The generation of the key is outside the scope of this document; however, general guidelines about the key, along with how to import/load it, are provided below.

Note: PGP Encryption is not supported.

- Supported Key Types:
 - SSH-2 RSA 2048 bit length
- Unsupported Key Types:
 - SSH-I RSA
 - SSH-2 DSA
- Correct Public Key Format: If opened in a text editor, the key should look like the screenshot below.



• Incorrect Public Key Format: If opened in a text editor, the key SHOULD NOT look like the screenshot below.



Once the key has been generated, it should be named "authorized_keys".

Notes:

- There is no file extension.
- There is an underscore between the words **authorized** and **keys**.

A .ssh subfolder needs to be created in the SFTP account's home directory. The "authorized_keys" file must be placed in the .ssh folder. The creation of this folder follows the same process as creating a PMP subfolder. Please refer to PMP Subfolders for steps on creating subfolders.

Appendix D: Reporting Gabapentin Prescriptions

Typically, records submitted to the NC PDMP requires a prescriber's DEA or NPI. Gabapentin is a substance of concern, required to be reported by G.S. 90-113.73, and as such may be obtained from prescribers who do not have a DEA registration and from veterinarians who have neither a DEA registration nor NPI. Refer to the scenarios below for guidance.

Scenario I:

Prescriber does not prescribe Schedule II–V controlled substances but does prescribe gabapentin. Prescriber does not have a DEA number.

Guideline: Leave the prescriber DEA number field blank and input the prescriber NPI number in the appropriate data submission field.

Scenario 2:

Prescriber is a veterinarian who does not prescribe controlled substances in Schedules II–V but does prescribe gabapentin. Veterinarians are not eligible to have NPI numbers.

Guideline: Leave the prescriber DEA number and prescriber NPI number fields blank. Enter the prescriber's professional license number in the PRE04 field.

If there are any other circumstances for which reporting of gabapentin presents issues for your pharmacy, please contact the NC PDMP at nccsrs@dhhs.nc.gov.