



About Us



Mission:

To help our members, and the communities we serve, be healthy.

Vision:

To be the most highly regarded managed care plan in California.





Agenda

- QIP vs. HEDIS Measures
- Purpose
- Introduction to HEDIS
- Coding / File Layout Overview
- ROADMAP
- Audit Process
- Primary Source Verification (PSV)





Purpose



Measure Description

Electronic Clinical Data Systems (ECDS) allows for data exchange from Provider Electronic Health Records (EHR) to PHC in order to capture depression screening and follow-up care. ECDS implementation is a vital component of furthering **Perinatal** and **PCP QIP** technical advancement toward 100% administrative data capture, and is a vital component of furthering the quality of care for covered PHC members.

*Note that NCQA plans to convert most hybrid measures to ECDS measures in the coming years. DHCS continues to make PHC accountable to several ECDS measures, **this process will continue to increase in emphasis.**

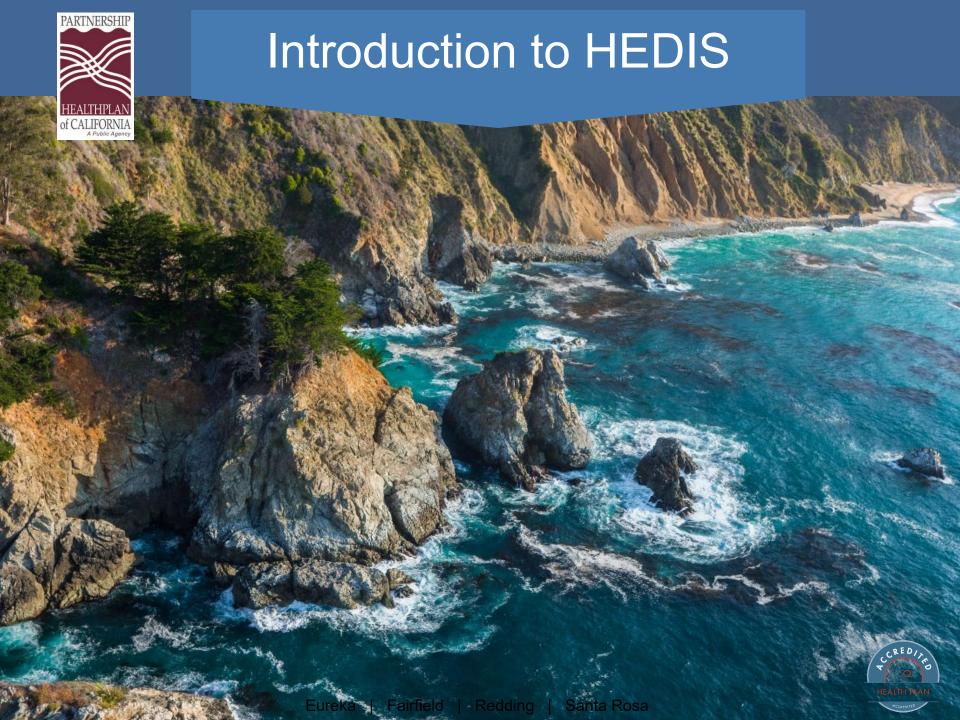
Incentive Amount - New! (Same for PCP and Perinatal QIPs)

- \$5,000 per Parent Organization with EHR vendor support
- \$10,000 per Parent Organization with no EHR vendor support

Documentation Source - New! (Same for PCP and Perinatal QIPs)

- Generate ECDS output and submit test file via Secure File Transfer Protocol (sFTP) by October 15, 2023
- Any corrections to the data must be completed by November 1, 2023.
- Submit final data file (using the exact programming used for the test file) via sFTP between **January 7**, **2024** and **January 14**, **2024**







HEDIS Overview

HEDIS stands for:

Healthcare Effectiveness Data and Information Set

Why does HEDIS exist?

- HEDIS is a measurement tool established and maintained by the National Committee for Quality Assurance (NCQA).
- HEDIS is used to evaluate clinical quality measures in a standardized way.
- The California Department of Healthcare Services (DHCS) selects a subset of the NCQA measures for Medi-Cal plans to report on annually as required for the State.
- DHCS uses HEDIS data to evaluate clinical quality outcomes in a standardized way and is utilized to evaluate the quality of health care across all health plans





Why ECDS?

- Electronic Clinical Data Systems (ECDS) measures provide information about the organization's use of clinical data to document high-quality patient care.
- ECDS encourages exchange of the information needed to provide high-quality services, ensuring it reaches the right people at the right time.
- ECDS may also support other care-related activities directly or indirectly, including evidence-based decision support, quality management and outcome reporting.
- ECDS is a reporting requirement by NCQA.





ECDS Timeline

- October 15, 2023 Submit test file via Secure File Transfer Protocol (sFTP)**
- October 15, 2023 Submit draft of ECDS ROADMAP to PHC**
- November 1, 2023 Deadline for sites to submit the final corrected data if discrepancies were identified. <u>Highly encouraged to submit</u> the corrected data file immediately in case there are additional errors**
- November 1, 2023 and January 31, 2024 Partnership HEDIS team will conduct Primary Source Verification.
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Associated Documents (All 2023 Editions)

- ECDS Measure Requirements
- ASF Data Reporting Template
- DEP Data Reporting Template
- Value Set File
- Sample SQL Code





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ECDS Measure Requirements

- General instructions
- Working with the submission templates
- Description of each measure, including definitions and logic of data pull
- List of report columns and definitions for each measure submission
- Background information on the measure, including technical definitions





Reporting Templates

- One for Alcohol Use Screening and Follow-Up (ASF Data Reporting Template)
- One for the combined Depression measures (DEP Data Reporting Template)
- Both are Excel files





Reporting Templates: Column Definitions

- Columns listed in the specifications document correspond to the columns on the reporting template
- The reporting template columns are in a particular order and have a particular format

Table in specifications document

Field Name	Data Type (Max Length)	Notes
Date of Service	Date	Visit date to the service provider (MM/DD/YYYY)
Site Name	Varchar (255)	Name of PCP or Specialist site where care was provided
Site ID Number	Varchar (50)	NPI number assigned to site where care was provided
Provider Key	Varchar (25)	A unique identifier for a provider. PCP ID (PHC assigned)
Clinician NPI Number	Varchar (179)	Provider NPI number for clinician who saw the patient on the date of service



Columns in submission template

4	А	В	С	D	E
1	Date_of_Service	Site_Name	Site_ID_Number	Provider_Key	Clinician_NPI_Number
2	Date	Varchar (255)	Varchar (50)	Varchar (25)	Varchar (179)
3					
4					
5					





Reporting Templates Are "Locked"

- In order for Partnership HealthPlan to integrate the data in a systematic way, all columns must be present and remain in the original order and format
- Therefore, the templates are "locked" so that the structure is preserved (for example, users cannot add or delete columns, or change the cell format)
- If the initial output needs to be populated in the template, data should be populated in mass (not row by row). Initial output must be saved for auditing.





Reporting Template Submission Names

- Save the template with your data in a specific manner
- The name should follow this convention: MEASUREID_SITENAME_DATE
- MEASUREID: "ASF" or "DEP"
- SITENAME: an abbreviation or shortened name of your organization
- DATE: in format YYYYMMDD with no dashes, periods or slashes
- Examples: ASF_SHASTA_20230930 DEP_PETALUMA_20240115





Value Set File

- For reference only (to possibly help with validation or understanding the measures)
- The file contains six tabs with lists of value set codes:
 - ✓ ASF: Alcohol counseling codes
 - ✓ ASF: Exclusion codes
 - ✓ DEP: Outpatient visit codes (standard)
 - ✓ DEP: Outpatient visit codes (non-standard)
 - ✓ DEP: Depression diagnosis codes
 - ✓ DEP: Exclusion codes





Sample SQL Code File (or "SQL Template")

- Aimed at SQL programmers. Gives ideas on how to pull the data and display it for the templates
- Value sets are incorporated in the code. Output columns are in correct order and format for the template
- Uses POSTGRES SQL language based on eCW and Nextgen base tables
- The SQL code can be modified for the table/field names at your organization and the SQL language used by your analytic software. This modified code must be saved for auditing and submitted as an attachment to the ROADMAP.
- Use your own "best practices" for specific data definitions and extraction (for example, the estimated date of delivery for the depression submission). Non-standard, individual billing codes used in the reporting programming must be explained in the ROADMAP.





Sample SQL Code

- Some sections are highlighted in blue where you will have to customize the code. The specifications document contains definitions for these items
- ASF: alcohol screen date and result (up to three types of screens accepted)
- ASF: alcohol counseling and follow-up (see definition in specifications)
- DEP: depression screen date and result (several possible screens)
- Both ASF and DEP: Join patients to the Partnership enrollment lists





Submit Only PHC Members

- Only patients enrolled in Partnership anytime during the measurement year should be submitted
- These patients have a unique CIN
- The CIN of patients assigned to the organization appears in the monthly enrollment file available in provider online services





Include Only Partnership Members from Enrollment Lists

- Match the Partnership members file with your total population in your analytics software electronically
- This process will be unique to your organization and software
- The default SQL code in the sample file identifies
 Partnership patients by their primary insurance within a Partnership insurance group (assuming that exists in your system)
- Use the best method available in your system to identify Partnership members





Other Notes About Report Design

 If you used and validated the depression report from the 2022 SQL template file (using some or all of the sample code), make a change to the Value Set definition section



SQL Template File, Depression Measures Section

Copy and replace the TEMPORARY TABLE temp_value_sets that contains the updated 2023 Value Set codes

Codes added and removed from the Value Sets since last year are highlighted in the Value Set file





Other Notes About Report Design

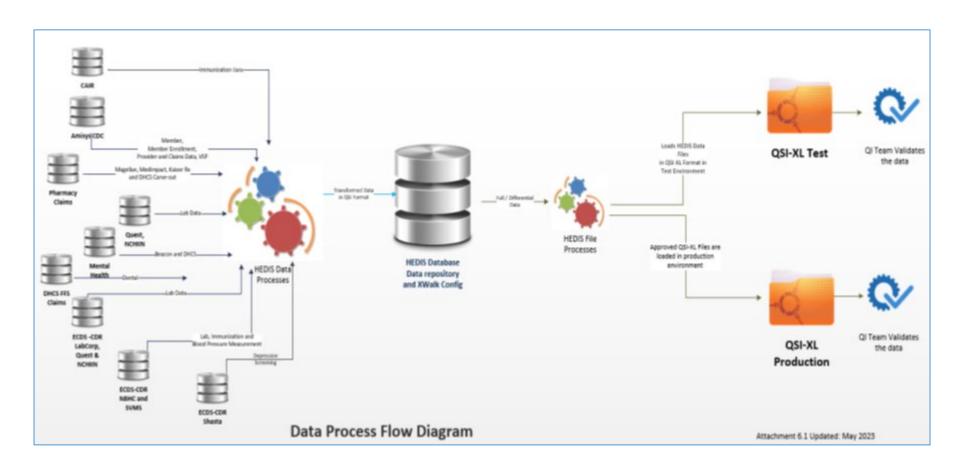
- Check your system for new items in structured data since last year:
 - ✓ New alcohol screening tools
 - ✓ New alcohol follow-up/counseling items
 - ✓ New depression screening tools

 Note that patient CIN can default to scientific notation format in Excel. Do NOT submit scientific notation





Data Flow Process







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ROADMAP Section 5 Purpose

- Record of Administration, Data Management and Processes (ROADMAP)
- The ROADMAP is a tool to help organizations of all types give auditors information about data used for HEDIS – where data comes from, and how the data is organized.
- The Roadmap also helps organizations send the right set of questions to the right people.
- If a third party entity is involved before the information reaches PHC, a description of the data flow is required (Section 5a).
- Section 5 and/or 5a is required in order to obtain auditor approval to integrate all supplemental data sources.





Section Description

- ROADMAP Section 5: Supplemental data and processes used during the measurement year.
- ROADMAP Section 5a: This section should be completed by the contracted data aggregator (e.g. HIE, data aggregator). Organizations validated through NCQA's Data Aggregator Validation (DAV) program do not need to complete this section.





ROADMAP Overview

 Let's take a deeper look at Section 5 and 5a of the ROADMAP

- A couple things to note:
 - On the document, Red indicates a response needed from the provider and Blue indicates PHC's response.
 - Providers can include an attachment such as a flow chart to describe their data flow process.
 - A list of organizations who are validated through NCQA's Data Aggregator Validation program can be found on the NCQA website.





Section 5: Supplemental Data

Section 5: Supplemental Data (IS 5)

Introduction

Supplemental data and processes used during the measurement year.

Complete a separate section for each supplemental data source being considered for use. Data received from an organization validated through NCQA's Data Aggregator Validation program must be identified with a Roadmap section.

Complete Section 5 (Tables 5.1, 5.5, 5.6) to address how data are received and used for HEDIS reporting. Indicate "NA" if questions are not applicable. You are not required to get responses from the validated entity.

Providers: please complete any section marked in red. PHC will provide responses in blue.

Organization
information

Organization name: Partnership HealthPlan of California

Date of completion: 01/23/2023

Name of supplemental data source: Provider

Table 5.1 General Information

5.1A The internal name of this data source (the name that will be used for supplemental data validation).

Provider

5.1B Summarize the intent or purpose of this data source (e.g., gap you are trying to fill, issue you are trying to solve by collecting these data).

To include depression screening results data for reported HEDIS measures.





Section 5a: Supplemental Data

Section 5a: Supplemental Data

Audit Roadmap section for data received from data aggregators

Introduction

This section should be completed by the contracted data aggregator (e.g., HIE, data aggregator). Organizations validated through NCQA's Data Aggregator Validation program do not need to complete this section.

If this section is completed by the data supplier, the organization must complete Section 5 (Tables 5.1, 5.4–5.6 and all applicable required documents) to address how it handles receipt of the data.

Providers: please complete any section marked in red. PHC will provide responses in blue.

Organization information

Organization name: Provider

Date of completion: 11/4/2022

Data aggregator type (e.g., HIE, data aggregator): Provider provides details. For example SQL Backend/Query Based

Name of entity that provides data: (i.e. NextGen, EPIC EHR)

Table 5A.1: System/Product Description

5A.1A Provide an overview of your product. ONC certified Ambulatory EHR Version 6.2021.1

Dravidor provides details. For example. The Heath plan posts undated member lists to their



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Audit Process

- To qualify for ECDS reporting, data must use standard layouts, meet the NCQA measure technical specifications and be accessible upon request.
- All supplemental data must be substantiated by proof-of-service documentation from the electronic health record.
- The ROADMAP document must be submitted and approved by the HEDIS auditor, prior to PSV.





Audit Process

- PHC must have clear policies and procedures describing how the data is collected and by whom.
- Documentation is required for only a sample, randomly selected by the auditor as part of the annual primary source verification process.





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Primary Source Verification Steps

- 1. PHC receives data from the provider
- 2. PHC sends the raw data file to the auditor
- Auditor randomly selects cases (approx. 16) for PSV
- 4. PHC sends Auditor selected cases to provider
- 5. Provider provides proof-of-service screen shots from their EHR, demonstrating they match the raw data file
- 6. PHC validates PSV POS, submits to auditor for approval





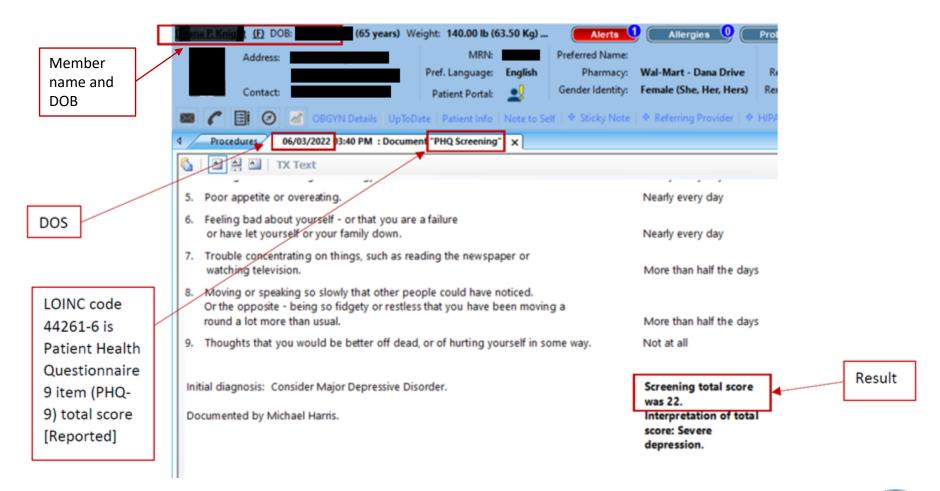
Proof-of-Service Documentation

- Documentation that is allowed includes:
 - A screen shot of the clinical summary from the visit for service, such as lab or radiology reports
 - A screen shot of the EHR or immunization registry records.
- Documentation that is not allowed includes:
 - Member surveys or documents completed by the member
 - Phone calls. Recorded phone calls to collect information about services rendered are not proof-of-service.





Sample Proof of Service







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QIP Spec Timeline

Step 1: Contact the PCP QIP team to let us know of your intent to implement ECDS and set-up an sFTP account: qip@partnershiphp.org

- Please provide the following details in an email to the PCP QIP inbox: o Name of Parent Organization
 - Contact name (one (1) contact per parent organization)
 - Contact phone number
 - Contact email address

Step 2: sFTP Account Set-Up (Must be completed by October 1, 2023)

- Once your contact information is received, an account will be generated in order to submit test files using the PHC ECDS Depression/Alcohol Screening & Counseling Templates.
 - An email will be sent to you from PHC's EDI team with log in credentials to access your new sFTP account

Step 3: Test File Submission (Data for January 1, 2023 – date the data is run)

- Acceptance of test file and HEDIS Roadmap 5 template via sFTP is due by October 15, 2023.
- The test file must include all data elements on the ECDS Depression/Alcohol Screening & Counseling Templates.
- The PHC HEDIS team will provide confirmation your test file has been received.
- If the test file data does not pass the HEDIS review, PCP sites will have until **November 1, 2023** to correct the data.
- In November and December, PHC's HEDIS team will reach out to PCPs to perform preliminary Primary Source Verification on a sample of data submitted.

Step 4: Final Data File Submission (Data for all of 2023)

- For incentive of the ECDS measure, acceptance of the final data file via sFTP is due between **January 7**, **2024 January 14**, **2024**.
- Final data files must include all data elements on the ECDS Depression/Alcohol Screening & Counseling Templates for all of 2023 (Jan 1 Dec 30).
- All PCP sites submitting final data are required to participate in the PHC HEDIS team Primary Source Verification Process.





Questions





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