

(2024 MIPS) Immunization Registry Reporting

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If you are engaged in submitting data to a clinical data registry, you can attest to measure in your Healthmonix MIPSpro account.

Learn more about DrChrono's Immunization Registries partner, Iron Bridge.

- [Get Started: DrChrono Immunization Registries](#)
- [How to use the Iron Bridge Integration \(Immunization Registries Partner\)](#)

Description

The MIPS eligible clinician is in active engagement with a public health agency to submit immunization data and receive immunization forecasts and histories from the public health immunization registry/immunization information system (IIS).

Exclusions

Any MIPS eligible clinician meeting one or more of the following criteria may be excluded from the Immunization Registry Reporting measure if the MIPS eligible clinician: 1. Does not administer any immunizations to any of the populations for which data is collected by its jurisdiction's immunization registry or immunization information system during the performance period. OR 2. Operates in a jurisdiction for which no immunization registry or immunization information system is capable of accepting the specific standards required to meet the certified electronic health record technology (CEHRT) definition at the start of the performance period. OR 3. Operates in a jurisdiction where no immunization registry or immunization information system has declared readiness to receive immunization data as of 6 months prior to the start of the performance period.

Definitions

Active Engagement - The MIPS eligible clinician is in the process of moving towards sending "production data" to a public health agency (PHA) or clinical data registry (CDR), or is sending production data to a PHA or CDR. Active engagement may be demonstrated in one of the following ways:


- **Option 1 – Pre-Production and Validation:** The MIPS eligible clinician must first register to submit data with the PHA or, where applicable, the CDR to which the information is being submitted. Registration must be completed within 60 days after the start of the performance period, while awaiting an invitation from the PHA or CDR to begin testing and validation. MIPS eligible clinicians that have registered in previous years do not need to submit an additional registration for subsequent performance periods. Upon completion of the initial registration, the MIPS eligible clinician must begin the process of testing and validation of the electronic submission of data. The MIPS eligible clinician must respond to requests from the PHA or, where applicable, the CDR within 30 days; failure to respond twice within a performance period would result in the MIPS eligible clinician not meeting the measure.
- **Option 2 – Validated Data Production:** The MIPS eligible clinician has completed testing and validation of the electronic submission and is electronically submitting production data to the PHA or CDR.

Production data – Refers to data generated through clinical processes involving patient care, and it is used to distinguish between data and "test data" which may be submitted for the purposes of enrolling in and testing electronic data transfers.

Reporting Requirements

YES/NO - The MIPS eligible clinician must attest YES to being in active engagement with a PHA to submit immunization data and receive immunization forecasts and histories from the public health immunization registry/immunization information system (IIS).

In your Healthmonix MIPSpro PI dashboard, you can attest by selecting an option and clicking **Save**.

Help Dashboard Admin Home kareem.davies@drchrono.com

2022 / DrChrono Test / B Demo / PI checklist PI Score : 0/25

Immunization Registry Reporting (PI_PHCDRR_1)

Complete:

- This group is in active engagement with a public health agency to submit immunization data and receive immunization forecasts and histories from the public health immunization registry/immunization information system (IIS).
- This group did not administer any immunizations to any of the populations for which data is collected by its jurisdiction's immunization registry or immunization information system during the performance period.
- This group operates in a jurisdiction for which no immunization registry or immunization information system is capable of accepting the specific standards required to meet the CEHRT definition at the start of the performance period.
- This group operates in a jurisdiction where no immunization registry or immunization information system has declared readiness to receive immunization data as of 6 months prior to the start of the performance period.
- None of the above

Measure Details

Measure Title: Immunization Registry Reporting

Measure ID: PI_PHCDRR_1

Objective: Public Health and Clinical Data Exchange

Description

The MIPS eligible clinician is in active engagement with a public health agency to submit immunization data and receive immunization forecasts and histories from the public health immunization registry/immunization information system (IIS).

Definitions

Active engagement - The MIPS eligible clinician is in the process of moving towards sending "production data" to a public health agency or clinical data registry, or is sending production data to a public health agency (PHA) or clinical data registry (CDR).

Active engagement may be demonstrated in one of the following ways:

Option 1 - Completed Registration to Submit Data: The MIPS eligible clinician registered to submit data with the PHA or, where applicable, the CDR to which the information is being submitted; registration was completed within 60 days after the start of the MIPS performance period; and the MIPS eligible clinician is awaiting an invitation from the PHA or CDR to begin testing and validation. This option allows MIPS eligible clinicians to meet the measure when the PHA or the CDR has limited resources to initiate the testing and validation process. MIPS eligible clinicians that have registered in previous years do not need to submit an additional registration to meet this requirement for each MIPS performance period.

Option 2 - Testing and Validation: The MIPS eligible clinician is in the process of testing and validation of the electronic submission of data. MIPS eligible clinicians must respond to requests from the PHA or, where applicable, the CDR within 30 days; failure to respond twice within a MIPS performance period would result in that MIPS eligible clinician not meeting the measure.

Option 3 - Production: The MIPS eligible clinician has completed testing and validation of the electronic submission and is electronically submitting production data to the PHA or CDR.

Production data - Refers to data generated through clinical processes involving patient care, and it is used to distinguish between data and "test data" which may be submitted for the purposes of enrolling in and testing electronic data transfers.

Cancel Save