Merit-Based Incentive Payment System (MIPS) Advancing Care Information Performance Category Measure

<u>Objective</u> :	Public Health and Clinical Data Registry Reporting
<u>Measure</u> :	Electronic Case Reporting The MIPS eligible clinician is in active engagement with a public health agency to electronically submit case reporting of reportable conditions.

Reporting Requirements

YES/NO

The MIPS eligible clinician must attest YES to being in active engagement with a public health agency to electronically submit case reporting of reportable conditions.

Scoring Information

BASE SCORE/PERFORMANCE SCORE/BONUS SCORE

- Required for Base Score (50%): No
- Eligible for bonus score: Yes, 5%

Note: Eligible clinicians must earn the full base score in order to earn any score in the Advancing Care Information performance category. In addition to the base score, eligible clinicians have the opportunity to earn additional credit through a performance score and the bonus score.

Regulatory References

- For further discussion, please see the Medicare Access and CHIP Reauthorization Act of 2015 (MACRA) final rule: <u>81 FR 77229</u>.
- In order to meet this objective and measure, an EP must use the capabilities and standards of CEHRT at 45 CFR 170.315 (f)(1), (f)(2), (f)(4), (f)(5), (f)(6) and (f)(7).

Certification and Standards Criteria

Below is the corresponding certification and standards criteria for electronic health record technology that supports achieving the meaningful use of this measure.

Certification Criteria (i) Consume and maintain a table of trigger codes to determine which encounters may be reportable. (ii) Match a patient visit or encounter to the trigger code based on the parameters of the trigger code table. (iii) Case report creation. Create a case report for electronic transmission: (A) Based on a matched trigger from paragraph (f)(5)(ii). (B) That includes, at a minimum: § 170.315(f)(5) Transmission to public health (1) The Common Clinical Data Set. agencies electronic case (2) Encounter diagnoses. Formatted according to at least one of the reporting following standards: (i) The standard specified in §170.207(i). (ii) At a minimum, the version of the standard specified in §170.207(a)(4). (3) The provider's name, office contact information, and reason for visit. (4) An identifier representing the row and version of the trigger table that triggered the case report.

§ 170.315(f)(6)	
Transmission to	
public health	
agencies—	
antimicrobial use	
and resistance	
reporting.	
8 170 215/f)/7)	

Create antimicrobial use and resistance reporting information for electronic transmission in accordance with the standard specified in §170.205(r)(1).

§ 170.315(f)(7)
Transmission to
public health
agencies—health
care surveys

Create health care survey information for electronic transmission in accordance with the standard specified in §170.205(s)(1).

Standards Criteria § 170.205(d)(2) HL7 2.5.1 (incorporated by reference in §170.299). **Electronic** submission to public health agencies for surveillance or reporting § 170.205(d)(3) Standard. HL7 2.5.1 (incorporated by reference in §170.299). **Electronic** Implementation specifications. PHIN Messaging Guide for Syndromic submission to Surveillance (incorporated by reference in §170.299) and Conformance public health Clarification for EHR Certification of Electronic Syndromic Surveillance, agencies for Addendum to PHIN Messaging Guide for Syndromic Surveillance surveillance or (incorporated by reference in §170.299). reporting Standard. HL7 2.5.1 (incorporated by reference in §170.299). § 170.205(d)(4) Implementation specifications. PHIN Messaging Guide for Syndromic **Electronic** Surveillance: Emergency Department, Urgent Care, Inpatient and submission to Ambulatory Care Settings, Release 2.0, April 21, 2015 (incorporated by public health reference in §170.299) and Erratum to the CDC PHIN 2.0 agencies for Implementation Guide, August 2015; Erratum to the CDC PHIN 2.0 surveillance or Messaging Guide, April 2015 Release for Syndromic Surveillance: reporting Emergency Department, Urgent Care, Inpatient and Ambulatory Care Settings (incorporated by reference in §170.299).

§ 170.207(a)(3)(4)	HTSDO SNOMED CT® International Release July 2012 (incorporated by reference in §170.299) and US Extension to SNOMED CT® March 2012 Release (incorporated by reference in §170.299).
	IHTSDO SNOMED CT®, U.S. Edition, September 2015 Release (incorporated by reference in §170.299).
§ 170.207(c)(2)(3)	Logical Observation Identifiers Names and Codes (LOINC®) Database version 2.40, a universal code system for identifying laboratory and clinical observations produced by the Regenstrief Institute, Inc. (incorporated by reference in §170.299).
	Logical Observation Identifiers Names and Codes (LOINC®) Database version 2.52, a universal code system for identifying laboratory and clinical observations produced by the Regenstrief Institute, Inc. (incorporated by reference in §170.299).

Additional standards criteria may apply. Review the <u>ONC 2015 Edition Final Rule</u> for more information.