



Medicaid EHR Incentive Program (Promoting Interoperability)

Eligible Professional Meaningful Use Attestation Manual

Program Year 2021



Table of Contents

1. Program Overview	1
1.1 Introduction	1
1.2 Background	2
2 Eligibility	2
2.1 Additional Requirements	3
2.2 Out-of-State Providers	4
2.3 Establishing Patient Volume.....	4
2.3.1 Patient Encounters Methodology	4
2.3.2 Eligible Professional Medicaid Encounter Definition	5
2.3.3 Definition of a Needy Individual Encounter	5
2.3.4 Group Practices	5
3 Payment Methodology	5
3.1 Payments	6
4 Provider Registration.....	6
5 Attestation Process & Validation	7
5.1 Attestation	7
5.2 Incentive Payments	8
6 Program Integrity	8
6.1 Attestation Appeal.....	9
6.2 Audit Appeal	9
7 Getting Started.....	14
7.1 Sign-in	14
7.2 Home Screen	15
7.3 Registration Data Screen	17
7.3.1 Provider CMS Registration Data	17
7.3.2 Provider Medicaid Attestation Data	18
7.4 Provider Eligibility Details Screen.....	19
7.4.1 Eligibility Details.....	19
7.4.2 Requesting KCHIP Report Data	21
7.4.3 Service Locations.....	24

7.5	Meaningful Use Questionnaire Screen	26
8	Requirements for Meaningful Use Measures	27
8.1	Meaningful Use Menu Screen	28
8.2	Meaningful Use Core Objectives – Stage 3	28
8.2.1	<i>MU Core Objective 1 – Protect Patient Health Information</i>	<i>28</i>
8.2.2	<i>MU Core Objective 2 – Electronic Prescribing (eRX)</i>	<i>29</i>
8.2.3	<i>MU Core Objective 3 – Clinical Decision Support.....</i>	<i>31</i>
8.2.4	<i>MU Core Objective 4 – Computerized Provider Order Entry.....</i>	<i>32</i>
8.2.5	<i>MU Core Objective 5 – Patient Electronic Access to Health Information</i>	<i>35</i>
8.2.6	<i>MU Core Objective 6 – Coordination of Care through Patient Engagement.....</i>	<i>37</i>
8.2.7	<i>MU Core Objective 7 – Health Information Exchange</i>	<i>40</i>
8.2.8	<i>MU Core Objective 8 – Public Health and Clinical Data Registry Reporting.....</i>	<i>43</i>
8.2.8.1	Measure 1: Immunization Registry Reporting	43
8.2.8.2	Measure 2: Syndromic Surveillance Reporting	45
8.2.8.3	Measure 3: Electronic Case Reporting	47
8.2.8.4	Measure 4: Public Health Registry Reporting	49
8.2.8.5	Measure 5: Clinical Data Registry Reporting.....	52
9	Electronic Clinical Quality Measures	55
9.1	Electronic Clinical Quality Measure Submission Selection Screen	56
9.2	Electronic Clinical Quality Measures Electronically Reported Selection Screen	57
9.3	Electronic Clinical Quality Measures Electronically Reported Summary	60
9.4	Electronic Clinical Quality Measures Manually Reported Selection Screen	61
9.5	Electronic Clinical Quality Measures Manually Reported	65
9.5.1	<i>CMS146</i>	<i>65</i>
9.5.2	<i>CMS137</i>	<i>66</i>
9.5.3	<i>CMS165</i>	<i>67</i>
9.5.4	<i>CMS156</i>	<i>69</i>
9.5.5	<i>CMS155</i>	<i>70</i>
9.5.6	<i>CMS138</i>	<i>72</i>
9.5.7	<i>CMS124</i>	<i>73</i>
9.5.8	<i>CMS153</i>	<i>75</i>
9.5.9	<i>CMS130</i>	<i>76</i>
9.5.10	<i>CMS117</i>	<i>77</i>
9.5.11	<i>CMS147</i>	<i>78</i>

9.5.12 CMS127 79

9.5.13 CMS131 80

9.5.14 CMS122 82

9.5.15 CMS134 83

9.5.16 CMS154 84

9.5.17 CMS145 85

9.5.18 CMS135 87

9.5.19 CMS144 88

9.5.20 CMS143 89

9.5.21 CMS142 90

9.5.22 CMS139 91

9.5.23 CMS161 92

9.5.24 CMS128 93

9.5.25 CMS136 94

9.5.26 CMS157 95

9.5.27 CMS129 96

9.5.28 CMS2 97

9.5.29 CMS68 99

9.5.30 CMS69 100

9.5.31 CMS133 102

9.5.32 CMS159 103

9.5.33 CMS177 104

9.5.34 CMS125 105

9.5.35 CMS149 107

9.5.36 CMS22 108

9.5.37 CMS50 109

9.5.38 CMS56 110

9.5.39 CMS66 111

9.5.40 CMS74 112

9.5.41 CMS75 113

9.5.42 CMS90 114

9.5.43 CMS249 115

9.5.44 CMS347 117

9.5.45 CMS349 119

9.5.46 CMS645 120

9.5.47 CMS771 121

10 Submitting Attestation 122

10.1 Pre-Attestation Summary Screen..... 122

 10.1.1 Objectives Summary..... 123

 10.1.2 Public Health Objectives Summary 128

 10.1.3 Electronic Clinical Quality Measures Summary (Electronically Reported)..... 129

 10.1.4 Electronic Clinical Quality Measures Summary (Manually Reported) 131

10.2 Incentive Payment Calculation Screen..... 133

10.3 Document Upload Screen 133

10.4 Attestation Statement Screen 135

10.5 Accepted Attestation Screen..... 138

10.6 Attestation Not Accepted Screen..... 138

10.7 Post Attestation Summary Screen 139

 10.7.1 Objectives Summary..... 139

 10.7.2 Public Health Objectives Summary 143

 10.7.3 Electronic Clinical Quality Measures Summary (Electronically Reported)..... 144

 10.7.4 Electronic Clinical Quality Measures Summary (Manually Reported) 145

10.8 Next Steps 150

1. Program Overview

1.1 Introduction

The Kentucky Medicaid Electronic Health Record (EHR) Incentive Program (also known as Promoting Interoperability) provides incentive payments to eligible professionals (EPs), eligible hospitals (EHs) and critical access hospitals (CAHs) as they demonstrate meaningful use (MU) of certified EHR technology (CEHRT). The purpose of this document is to provide instructions for providers to complete an attestation for the Kentucky Medicaid EHR Incentive Program (Promoting Interoperability) using the KYSLR system.

Resources:

- 42 CFR Parts 412, 413, 422 et al. Medicare and Medicaid Programs; Electronic Health Record Incentive Program Final Rule located at <https://www.federalregister.gov/documents/2010/07/28/2010-17207/medicare-and-medicaid-programs-electronic-health-record-incentive-program>
- 42 CFR Parts 412 and 495 et al. Medicare and Medicaid Programs; Electronic Health Record Incentive Program - Stage 3 and Modifications to Meaningful Use in 2015 Through 2017; Final Rule located at <https://www.federalregister.gov/documents/2015/10/16/2015-25595/medicare-and-medicaid-programs-electronic-health-record-incentive-program-stage-3-and-modifications>
- Kentucky State Medicaid HIT Plan (SMHP) located at <https://chfs.ky.gov/agencies/dms/ehr/Documents/SMHP%202019%20Annual%20Update%20v%201.0.pdf>
- Kentucky Medicaid EHR Application Portal located at <https://prd.webapps.chfs.ky.gov/KYSLR/login.aspx>
- Medicare and Medicaid Electronic Health records (EHR) Incentive Program (Promoting Interoperability) located at <https://www.cms.gov/Regulations-and-Guidance/Legislation/EHRIncentivePrograms/index.html?redirect=/EHRincentivePrograms/>
- Office of the National Coordinator for Health Information Technology located at <https://www.healthit.gov/>
- Kentucky Health Information Exchange located at <https://khie.ky.gov/PAGES/INDEX.ASPX>

Regional Extension Centers (RECs) have been designated to provide technical assistance to Kentucky providers. The RECs provide a full range of assistance and are listed below:

- Northeast Kentucky Area
Kentucky Rural Healthcare Information Organization (KRHIO)
Website: <https://krhio.org/>
Phone: 855-385-2089
E-mail: admin@nekyrhio.org

- Remaining Areas of Kentucky
Kentucky Regional Extension Center
Website: <http://www.kentuckyrec.com/>
Phone: 888-KY-REC-EHR or 859-323-3090
E-mail: kyrec@uky.edu

1.2 Background

The Centers for Medicare & Medicaid Services (CMS) has implemented, through provisions of the American Recovery and Reinvestment Act of 2009 (ARRA), incentive payments to EPs, EHS and CAHs, participating in Medicare and Medicaid programs that are meaningful users of CEHRT. The incentive payments are not a reimbursement, but are intended to encourage providers to adopt, implement, or upgrade CEHRT and use it in a meaningful manner.

Use of certified EHR systems is required to qualify for incentive payments. The Office of the National Coordinator for Health Information Technology (ONC) has issued rules defining certified EHR systems. More information about this process is available at <http://www.healthit.gov>. CMS continues its advancement of CEHRT utilization, focusing on burden reduction, and improving interoperability and patient access to health information.

The Kentucky Department for Medicaid Services (DMS) works closely with federal and state partners to ensure the Kentucky Medicaid EHR Incentive Program (Promoting Interoperability) fits into the overall strategic plan for the Kentucky Health Information Exchange (KHIE), thereby advancing national and Kentucky goals for HIE.

Providers can update their registration with the Medicare and Medicaid registration and attestation system (also referred to as the NLR). CMS' official Web site for the Medicare and Medicaid EHR Incentive Programs (Promoting Interoperability) can be found at <http://www.cms.gov/EHRIncentivePrograms/>. The site provides general and detailed information on the programs, including tabs to guide users on the path to payment, eligibility, MU, CEHRT, and frequently asked questions.

2 Eligibility

While providers could begin the program in Calendar Year (CY) 2011, they must have initiated participation in the program and received a payment no later than CY 2016. Program Year 2021 is the final year of participation. All attestations must be submitted no later than August 31st, 2021. Any attestation submitted after the deadline will not be eligible for the incentive payment.

The first tier of provider eligibility for the program is based on provider type and specialty. If the provider type and specialty for the submitting provider in the Kentucky Medicaid Partner Portal Application **does not** correspond to the provider types and specialties approved for participation in the Kentucky Medicaid EHR Incentive Program (Promoting Interoperability), the provider will receive an error message with a disqualification statement.

At this time, CHFS DMS has determined that the following providers are potentially eligible to enroll in the Kentucky Medicaid EHR Incentive Program (Promoting Interoperability):

- Physicians = Any provider who has a Provider Type 64 and Specialty other than 345 (Pediatrics)
- Physician Assistants (practicing in a FQHC [Provider Type 31 and Specialty 80] or RHC [Provider Type 35] led by a Physician Assistant) = Any provider with a Provider Type 95 and Specialty other than 959 (PA Group). A FQHC or RHC is considered to be PA led in the following instances:
 - The PA is the primary provider in a clinic (e.g., part time physician and full time PA in the clinic)
 - The PA is the clinical or medical director at a clinical site of the practice
 - The PA is the owner of the RHC
- Pediatricians = Any provider with a Provider Type 64 and Specialty 345
- Nurse Practitioners = Any provider with a Provider Type 78 and not Specialty 095 (CNM) or 789 (Nurse Practitioner Group)
- CNMs = Any provider with a Provider Type 78 and Specialty 095
- Dentists = Any provider with a Provider Type 60 (Individual)
- Optometrists = Any provider with a Provider Type 77
- Acute Care Hospital = Any provider with a Provider Type 01 and Specialty 010
- Children’s Hospital = Any provider with a Provider Type 01 and Specialty 015
- CAH = Any provider with a Provider Type 01 and Specialty 014

2.1 Additional Requirements

To qualify for an EHR incentive payment for each year the EP seeks the incentive payment, not be hospital-based and must:

1. Meet one of the following patient volume criteria:
 - a. Have a minimum of 30 percent patient volume attributable to individuals receiving TXIX and/or TXXI-CHIP (but not separate CHIPs) Medicaid services; **or**
 - b. Have a minimum 20 percent patient volume attributable to individuals receiving TXIX and/or TXXI-CHIP (but not separate CHIPs) Medicaid services, **and** be a pediatrician; **or**
 - c. Practice predominantly in a FQHC or RHC and have a minimum 30 percent patient volume attributable to needy individuals.
2. Have no sanctions and/or exclusions.

An individual EP may choose to receive the incentive directly or assign it to a Medicaid contracted clinic or group to which the provider is associated. The tax identification number (TIN) of the individual or entity receiving the incentive payment is required when registering with the National Level Registry (NLR) and must match a TIN linked to the individual provider in DMS’s system. If there is no contract on file with Kentucky Medicaid, the system will not be available to a provider for attestation until a contract has been approved by DMS. The following table is a summary of qualifying provider types and minimum patient encounter volumes.

Qualifying Providers by Type and Patient Volume

Program Entity	Percent Patient Volume Over Minimum 90-days	
Physicians	30%	Or the Medicaid EP practices predominantly in an FQHC or RHC -30% “needy individual” patient volume threshold
Pediatricians	20%	
Dentists	30%	
Optometrist	30%	
Physician Assistants when practicing at an FQHC/RHC led by a physician assistant	30%	
Nurse Practitioner	30%	

2.2 Out-of-State Providers

The Kentucky Medicaid EHR Incentive Program (Promoting Interoperability) welcomes out-of-state providers to participate in this program as long as they have at least one physical location in Kentucky. Kentucky must be the only state they are requesting an incentive payment from during that participation year. For audit purposes, out-of-state providers must make available any and all records, claims data, and other data pertinent to an audit by either the Kentucky DMS program or CMS. Records must be maintained as applicable by law in the state of practice or Kentucky, whichever is deemed longer.

2.3 Establishing Patient Volume

An eligible provider must annually meet patient volume requirements to participate in Kentucky’s Medicaid EHR Incentive Program (Promoting Interoperability) as established through the state’s CMS approved State Medicaid Health IT Plan (SMHP). The patient funding source identifies who can be counted in the patient volume: Title XIX (TXIX) – Medicaid and Title XXI (TXXI) – CHIP (but not separate CHIPs). All providers should calculate patient volume based on TXIX - Medicaid and/or TXXI-CHIP and out-of-state Medicaid patients.

2.3.1 Patient Encounters Methodology

- To calculate TXIX-Medicaid and/or TXXI-CHIP patient volume, an EP must divide:
 - The total TXIX and/or TXXI-CHIP Medicaid or out-of-state Medicaid patient encounters in any representative, continuous 90-day period in the prior calendar year or preceding 12 months from date of attestation; by
 - The total patient encounters in the same 90-day period.
- EPs Practicing Predominantly in an FQHC/RHC – to calculate needy individual patient volume, an EP must divide:
 - The total needy individual patient encounters in any representative, continuous 90-day period in the prior calendar year or preceding 12 months from date of attestation; by
 - The total patient encounters in the same 90-day period.

2.3.2 Eligible Professional Medicaid Encounter Definition

For purposes of calculating EP patient volume, a Medicaid encounter is defined as any service rendered on any one day to an individual enrolled in a Medicaid program whether or not Medicaid had a financial interest in the services that were rendered.

2.3.3 Definition of a Needy Individual Encounter

For purposes of calculating patient volume for an EP practicing predominantly in an FQHC/RHC, a needy individual encounter is defined as services rendered on any one day to an individual where medical services were:

- Furnished by the provider as uncompensated care; or
- Furnished at either no cost or reduced cost based on a sliding scale determined by the individual’s ability to pay.

2.3.4 Group Practices

Clinics or group practices will be permitted to calculate patient volume at the group practice/clinic level, but only in accordance with all of the following limitations:

- The clinic or group practice’s patient volume is appropriate as a patient volume methodology calculation for the EP.
- There is an auditable data source to support the clinic or group practice’s patient volume determination.
- **All** EPs in the group practice or clinic must use the same methodology for the payment year.
- The clinic or group practice uses the entire practice or clinic’s patient volume and does not limit patient volume in any way; and if an EP works inside and outside of the clinic or practice, then the patient volume calculation includes only those encounters associated with the clinic or group practice, and not the EP’s outside encounters.

3 Payment Methodology

The maximum incentive payment an EP could receive from Kentucky Medicaid is \$63,750, over a period of six years, or \$42,500 for pediatricians with a 20-29% Medicaid patient volume as shown below.

EP Patient Volume	EP (30%)	Pediatrician (20-29%)
Year 1	\$21,250	\$14,167
Year 2	\$8,500	\$5,666
Year 3	\$8,500	\$5,666
Year 4	\$8,500	\$5,666
Year 5	\$8,500	\$5,666
Year 6	\$8,500	\$5,666
Total Incentive Payment	\$63,750	\$42,500

Since pediatricians are qualified to participate as physicians, and therefore classified as EPs, they may qualify to receive the full incentive if the pediatrician can demonstrate that they meet the minimum 30% Medicaid patient volume requirements.

3.1 Payments

EP payments will be made in alignment with the calendar year and an EP must begin receiving incentive payments no later than CY 2016. EPs will assign the incentive payments to a tax ID (TIN) in the CMS EHR Registration and Attestation National Level Repository (NLR). The TIN must be associated in the Kentucky Medicaid Partner Portal Application system with either the EP him/herself or a group or clinic with whom the EP is affiliated. EPs who assign payment to himself or herself (and not a group or clinic) will be required to provide DMS with updated information. Each EP must have a current DMS contract and be contracted for at least 90 days.

For each year a provider wishes to receive a Medicaid incentive payment, determination must be made the provider was a meaningful user of EHR technology during that year. Medicaid EPs are not required to participate on a consecutive annual basis. However, the last year that an EP may begin receiving payments is 2016, and the last year the EP can receive payments is 2021. In the event that DMS determines monies have been paid inappropriately, incentive funds will be recouped and refunded to CMS.

The timeline for receiving incentive payments is illustrated below:

	CY 2011	CY 2012	CY 2013	CY 2014	CY 2015	CY 2016
CY 2011	\$21,250					
CY 2012	\$8,500	\$21,250				
CY 2013	\$8,500	\$8,500	\$21,250			
CY 2014	\$8,500	\$8,500	\$8,500	\$21,250		
CY 2015	\$8,500	\$8,500	\$8,500	\$8,500	\$21,250	
CY 2016	\$8,500	\$8,500	\$8,500	\$8,500	\$8,500	\$21,250
CY 2017		\$8,500	\$8,500	\$8,500	\$8,500	\$8,500
CY 2018			\$8,500	\$8,500	\$8,500	\$8,500
CY 2019				\$8,500	\$8,500	\$8,500
CY 2020					\$8,500	\$8,500
CY 2021						\$8,500
Total	\$63,750	\$63,750	\$63,750	\$63,750	\$63,750	\$63,750

4 Provider Registration

2016 was the last year a provider could initiate participation with the Kentucky Medicaid EHR Incentive Program (Promoting Interoperability). If changes to the registration need to be made, such as: address, phone number, taxpayer ID number (TIN) of the entity receiving the payment and the e-mail address; you may log into the NLR at <https://ehrincentives.cms.gov/hitech/login.action>.

The Quality Payment Program (QPP) is federal legislation altering the way clinicians are reimbursed for their Medicare Part B encounters. Clinicians have two tracks, Merit-based Incentive Payment System (MIPS) or Advanced Alternative Payment Models (APMs), to choose

from in the QPP based on their practice size, specialty, location or patient population. For more information, please visit <https://qpp.cms.gov/>.

5 Attestation Process & Validation

DMS uses the secure KYSLR system to house the attestation system. If an EP transfers from another state in which they participated in the program, updates their registration at the NLR and does not receive the link to the attestation system within two business days, assistance is available by contacting the Kentucky Medicaid EHR Incentive Program (Promoting Interoperability) at EHRIncentives@ky.gov or 502-564-0105 extension 2463.

5.1 Attestation

The following is a brief description of the information that a provider must report or attest to during the process:

1. The provider will log into the KYSLR at <https://prd.webapps.chfs.ky.gov/KYSLR/login.aspx> using their NPI and CMS assigned Registration Identifier.
2. The provider is asked to view the information displayed with the pre-populated data received from the NLR.
3. EPs will then enter two categories of data to complete the Eligibility Provider Details screen including: 1) patient volume characteristics, and 2) certification number for the ONC-ATCB certified EHR system (or numbers if obtained in modules).
4. EPs will submit MU data for objectives and electronic Clinical Quality Measures (eCQMs).
5. The EP will be asked to attest that:
 - The information submitted is accurate to the knowledge and belief of the EP.
 - The information submitted is accurate and complete for numerators, denominators and exclusions for functional measures applicable to the EP.
 - A zero was reported in the denominator of a measure when an EP did not care for any patients in the denominator population during the EHR reporting period.
 - The information submitted includes information on all patients to whom the measure applies.
 - As a meaningful EHR user, at least 50% of my patient encounters during the EHR reporting period occurred at the practice/location given in my attestation information and is equipped with CEHRT.
 - The information submitted for Objective 1, Protect Patient Health Information, requires a Security Risk Analysis to be completed within the calendar year of the EHR reporting period.
 - The information submitted for eCQM's was generated as output from an identified CEHRT.
 - The information submitted for eCQM's includes at least one outcome or high priority measure. If there are no outcome or high priority measures relevant to the EP's scope of practice, 6 relevant measures were reported.
 - Acknowledges the requirement to cooperate in good faith with ONC direct review of the EPs health information technology certified under the ONC Health IT Certification Program if a request to assist in ONC direct review

- is received.
- If requested, cooperated in good faith with ONC direct review of EPs health information technology certified under the ONC Health IT Certification Program as authorized by 45 CFR part 170, subpart E, to the extent that such technology meets (or can be used to meet) the definition of CEHRT, including by permitting timely access to such technology and demonstrating its capabilities as implemented and used by the EP in the field.
 - Acknowledges the option to cooperate in good faith with ONC-ACB surveillance of his or her health information technology certified under the ONC Health IT Certification Program if a request to assist in ONC-ACB surveillance is received
 - If requested, cooperated in good faith with ONC-ACB surveillance of the EPs health information technology certified under the ONC Health IT Certification Program as authorized by 45 CFR part 170, subpart E, to the extent that such technology meets (or can be used to meet) the definition of CEHRT, including by permitting timely access to such technology and demonstrating capabilities as implemented and used by the EP in the field.
6. The providers are asked to electronically sign the attestation.
- The provider or the agent/ staff member's initials are entered.
 - The providers NPI is entered.

The attestation itself is electronic and will require the provider to attest to meeting all requirements defined in the federal regulations. Some documentation will have to be provided to support specific elements of attestation. All providers are required to submit supporting documentation for patient volume claimed in the attestation. More information on documentation is provided in the attestation system. Once the electronic attestation is submitted by a qualifying provider and appropriate documentation is provided, DMS will conduct a pre-payment audit, which will include cross-checking for potential duplication payment requests, checking provider exclusion lists and verifying supporting documentation. All providers will be required to attest to MU to receive incentive payments.

5.2 Incentive Payments

Upon submission of the attestation and receipt of required documentation, verification is completed by DMS. Providers will be notified of approval for payment by email to the email address submitted with registration. Please be sure the email address provided is current.

6 Program Integrity

DMS has a contract with the Office of Inspector General (OIG) to perform audits and investigations of potential Medicaid fraud and/or abuse; therefore, OIG will conduct post payment incentive money audits. The audits conducted will investigate for all things attested; including, but not limited to the CEHRT component, percentage of Medicaid population

treated, Medicaid eligibility, etc. Any documentation to which an EP or EH attests, including future MU, will be audited. All reviews will ensure that no duplication of payment occurred. The OIG will submit reports on audit findings and recommendations to the DMS Division of Program Integrity. All documentation supporting the attestation is to be retained for six years.

6.1 Attestation Appeal

You may appeal the determination made by the Kentucky Department for Medicaid Services on your incentive payment application. In accordance with 907 KAR 6:005 Section 13, to appeal the provider must request a dispute resolution meeting. The request shall be in writing and mailed to and received by the department within 30 calendar days of the date the notice was received. The request must clearly identify each specific issue and dispute, and clearly state the basis on which the department's decision on each issue is believed to be erroneous. The provider shall also state the name, mailing address, and telephone number of individuals who are expected to attend the dispute resolution meeting on the provider's behalf. Any supporting documentation to the appeal should be included with the request. The address to send the request is below:

Division of Program Integrity
ATTN: EHR Appeal
Department for Medicaid Services
275 E. Main Street, 6E-A
Frankfort, KY 40621

6.2 Audit Appeal

You may appeal the determination made by DMS on your post payment audit. To request an appeal, the provider must log into the attestation website at

<https://prd.webapps.chfs.ky.gov/KYSLR/login.aspx>.

KY Medicaid EHR Incentive Program

In order to receive EHR incentive payments from Kentucky Medicaid, you first have to register at the [CMS Web Site](#) . After about 24 hours of successfully registering at the CMS level you should be able to complete your application on this site.

Please enter your NPI

Please enter the CMS assigned Registration Identifier

Step 1: After the provider has logged into the KYSLR, the provider will select 'view' on the Payment Year that is being appealed.

Provider Attestation Navigation			
ATTESTATION DETAILS AND NAVIGATION			
Payment Year	Status	AttestationID	Action
1	Paid	KY0001193	View
2	Attest_inProcess	-	Begin/Modify Attestation

Step 2: Click 'Appeals' on the navigation pane.

Send E-mail
Home
Reports
Document Upload
View All Payment Years
Issues/Concerns
Appeals
Additional Resources
KY Medicaid EHR Site
CMS EHR Site

Step 3: Select Audit Appeal.

Appeals (Year 1 Attestation)

What are you appealing:
--Select--
Attestation Appeal
Audit Appeal

Step 4: Select the Program Year you are appealing.

Appeals (Year 3 Attestation)

What are you appealing: Audit Appeal ▼

Provider Audit Appeal

Name	NPI	Payee NPI	CCN	Status	Audit Start date	Program Year	Select
Larry Crick	1821155532	1457784126		Audit Completed	2/19/2020	2011	Select

Step 5: Select the Appeal File Date, Appeal Type and enter Appeal Notes. The Appeal Reason is preselected.

(Year 2 Attestation)

Summary

Audit Case Number:	1982753695	Audit Status:	Audit Completed
Name:	Terri Watkins	NPI:	1982753695
Payee NPI:	1437691276	Address:	12 Mill creek park, bldg 12, Frankfort, KY, 40601
Audit Program Year:	2011	Audit Payment Year:	1

Appeal Status:

Appeal Setup
Findings
Appeal Document Upload
Appeal Outcome

Appeal Setup

Appeal File Date:

Appeal Reason:

Appeal Type:

Appeal Notes:

Step 6: To view the appeal findings, select the 'Findings' tab. This tab will display the appeal findings as well as any notes to and from the appeal reviewer.

(Year 3 Attestation)

Summary

Audit Case Number:	1821155532	Audit Status:	Audit Completed
Name:	Larry Crick	NPI:	1821155532
Payee NPI:	1457784126	Address:	12 Mill creek park, bldg 12, Frankfort, KY, 40601
Audit Program Year:	2011	Audit Payment Year:	1
Appeal Status:	Received filed appeal		

Appeal Findings

	Start Date	End Date	Notes	Provider Comments	Provider Action Required
<input type="button" value="Select"/>	7/3/2020	7/21/2020	Testing appeals. Plea...	Uploaded the s...	<input type="checkbox"/>
<input type="button" value="Select"/>	7/3/2020	7/21/2020	Testing appeals. Plea...	Uploaded the ...	<input type="checkbox"/>
<input type="button" value="Select"/>	6/29/2020	6/29/2020	test	1821155532 tes...	<input type="checkbox"/>

Findings Notes: Testing appeals. Please upload the supporting document for the audits.

Provider Comments: Uploaded the supporting documents
Testing for the appeal

Previous
Save
Save & Next

Step 7: Select the 'Appeal Document Upload' tab to upload supporting documentation to attach to your appeal. After you have uploaded your documentation, click on the 'Appeal Setup' tab to review your information. Once information is verified, click 'Next'.

Appeal Setup
Findings
Appeal Document Upload
Appeal Outcome

Document Upload

Upload Document: Browse...

Document Type: --Select-- ▼

Previous
Upload
Next

The appeal has been submitted. You may make changes to the appeal until 5pm. At 5pm, the appeal will be sent to CMS and locked for internal review. You can check the status of your appeal by viewing the summary.

Step 8: Select the 'Appeal Outcome' tab to view the outcome of the appeal.

(Year 5 Attestation)

Summary

Audit Case Number:	OIG00002	Audit Status:	Audit Completed
Name:	test002 test002	NPI:	0000000002
Payee NPI:	0000000002	Address:	12 Mill Creek Park, PO Box 1234, Frankfort, KY, 40621
Audit Program Year:	2018	Audit Payment Year:	4
Appeal Status:	Appeal dismissed with prejudice		

Appeal Outcome: Appeal Denied

7 Getting Started

EPs are required to provide details including patient volume characteristics, EHR details, upload requested documentation and electronically sign the attestation.

The provider begins the Kentucky Medicaid EHR Incentive Program (Promoting Interoperability) attestation process by accessing the KYSLR system at <https://prd.webapps.chfs.ky.gov/KYSLR/login.aspx>.

7.1 Sign-in

KY Medicaid EHR Incentive Program

In order to receive EHR incentive payments from Kentucky Medicaid, you first have to register at the [CMS Web Site](#) . After about 24 hours of successfully registering at the CMS level you should be able to complete your application on this site.

Please enter your NPI

Please enter the CMS assigned Registration Identifier

The provider enters the NPI and CMS assigned Registration Identifier that was returned by the NLR. Upon registration at the CMS registration site, you are assigned a CMS registration

identifier. The identifier is used for accessing the KYSLR and should be safeguarded as a password.

If the data submitted by the provider matches the data received from the NLR, the Home Screen will display. If the provider entry does not match, an error message with instructions will be returned. After five failed attempts, the provider will be locked out of the KYSLR for 15 minutes.

For assistance with registration, contact the NLR Production Support Help Desk toll free at (833) 238-0203 Monday through Friday, 8 a.m. to 5 p.m. Eastern time or email the help desk at NLRProdSupport@cms.hhs.gov.

7.2 Home Screen

The Home screen provides announcements, information about the provider’s current Kentucky attestation review as well as allows navigation for the provider to view a previous attestation or begin/modify a new attestation for their next EHR Incentive payment. This is also where the provider selects the Program Year they are attesting to. The status of their EHR is preselected to Meaningful User.

Home (Year 3 Attestation)

Announcements And Messages

Testing for PY 2021. Check all the screens and Reporting period.

Issues/Concerns

Clicking the below link will redirect you to the Issues/Concerns page, where you will be able to submit any issues and view the responses received from the DMS.

[Click Here](#)

Provider Information

You are currently enrolled in KY’s EHR Incentive Program.
Payment Year '3' is your current year attestation.

The current status of your application for the year 3 payment is 'AWAITING PROVIDER ATTESTATION'.

Stage of Meaningful Use

1st Year	2011	2019	2021	2022	2023	2024
2011	AIU 3	MU Stage 3 (90 Days)	MU Stage 3 (90 Days)			

Provider Status Flow

```

graph LR
    A[CMS Registration] --> B[Preliminary Verification]
    B --> C[Provider Attestation]
    style A fill:#add8e6,stroke:#333,stroke-width:1px
    style B fill:#add8e6,stroke:#333,stroke-width:1px
    style C fill:#add8e6,stroke:#333,stroke-width:1px
    
```

Completed
In Process

Provider Attestation Details

* For which program year are you applying? * Indicate the status of your EHR:

2021 Meaningful User

[Save Attestation Details](#)

Provider Attestation Navigation

ATTESTATION DETAILS AND NAVIGATION

Payment Year	Status	AttestationID	Action
1	Paid	KY0001042	View
2	Paid	KY0001559	View
3	Attest_inProcess	-	Begin/Modify Attestation

There are seven sections to the Home page listed below:

- Announcements and Messages – Displays messages or announcements for the provider.
- Issues/Concerns – Provides a link for the provider to submit a new issue or view a response to an issue.
- Provider Information – Provides a high-level status for the provider including the current payment year and the current status for the payment year.
- Stage of Meaningful Use – Supplies the stage of MU the provider will need to attest to according to the program year.
- Provider Status Flow – Displays a diagram showing the provider’s current year’s attestation. If the provider has been found not eligible for any reason, specific reasons for that finding is shown in this section.
- Provider Attestation Details – Provider selects the Program Year. EHR status is pre-selected as Meaningful User- currently meaningfully using CEHRT and are prepared to attest to MU and eCQMs.
- Provider Attestation Navigation – Lists the provider’s attestations by payment year and provides the navigation actions available for each year. These options may include:
 - View for a previously paid attestation;
 - View Attestation for a completed attestation;

- Begin/Modify for a new or not yet completed attestation.

7.3 Registration Data Screen

7.3.1 Provider CMS Registration Data

The data displayed in the Provider CMS Registration Data section is view only. If any of this data is incorrect, the data must be updated by logging in to the CMS Registration Module, making the updates and re-submission of the registration. Please allow 24 hours for the changes to be reflected.

Registration Data (Year 2 Attestation)

Provider CMS Registration Data

*** If any of this information is incorrect, please correct on the [CMS Registration Module](#).

Applicant NPI: 2111111111	Applicant TIN: 123456789	Name: Ubjk	Suffix:
Payee NPI: 9591196708	Address : Trzypfhp ,	Payee TIN: 73960002	City/State: Ubjk / GB
Program Option: MEDICAID	Zip Code: 40150 -	Medicaid State: KY	Phone Number: 5025640105
Provider Type: Physician	Email: EHRincentives@ky.gov	Participation Year: 2	Specialty: None
Federal Exclusions: None	State Rejection Reason: None		

Provider Medicaid Attestation Data

*** Please update the data below in reference to this attestation

Mailing Address:

Address 1: <input type="text" value="Trzypfhp"/>	Medicaid Provider Type: <input type="text" value="Nurse-Practitioner"/>
Address 2: <input type="text"/>	
City: <input type="text" value="testing"/>	State: <input type="text" value="GB"/>
ZipCode: <input type="text" value="40150"/>	<input type="text"/>

Were you assisted by a Regional Extension Center in Kentucky? Yes No

Previous
Next
Save
Cancel

The fields from the CMS registration are listed below:

- Applicant National Provider Identifier (NPI) – This is the eligible provider’s individual NPI. The NPI registered at CMS should be the same individual NPI that is enrolled in Kentucky Medicaid.
- Applicant TIN – This is the eligible providers Tax Identification Number. This TIN should be the same TIN that is listed for the provider in MMIS.
- Payee National Provider Identifier (NPI) – This is the eligible provider’s payee NPI given during the CMS registration. The Payee NPI should be enrolled in Kentucky Medicaid and listed as a payee with whom the individual provider is a member. **Note:** When a provider is linked to a Payee NPI that has multiple Medicaid ID’s enrolled in Kentucky Medicaid under that Payee NPI, the provider is required to select the appropriate Medicaid ID that the provider should be paid under.
- Payee TIN – The tax identification number associated with the payee NPI. This was the tax ID given during registration that will have the tax liability of the incentive payment. The Payee TIN should match the FEIN or SSN listed for the payee NPI within Kentucky Medicaid.
- Program Option – This program option was selected by the provider during their registration. It will be Medicaid if you are attesting with a State Agency and not Medicare.
- Medicaid State – This is the state that was selected during the provider’s registration.
- Provider Type – This is the provider type that was given during the registration at CMS. This type will be validated with your type of license.
- Participation year – This is the provider’s participation year with the program.
- Federal Exclusion – This will list any federal exclusion found on the provider if any during registration with CMS.
- Name – The Provider’s name listed on the CMS Registration.
- Address 1 – The provider’s street address listed on the CMS registration. Note: This is the address where all incentive monies will be mailed.
- Address 2 – The provider’s street address listed on the CMS registration.
- City/State – The provider’s city/state listed on the CMS registration.
- Zip Code – The provider’s zip code listed on the CMS registration.
- Phone Number – The provider’s phone number given on the CMS registration. This number is used for contact by EHR staff reviewing the attestations.
- Email – The provider’s email given during the CMS registration. This email address is used for system-generated emails on updates for the provider’s attestation and communication from the EHR review staff. **Note:** It is very important that this email address be accurate and up-to-date.
- Specialty – The provider’s specialty listed in the CMS registration.
- State Rejection Reason – This lists the state rejection reason if any are found. This will only list federal codes for rejection, for a more detailed state specific rejection see the home page.

7.3.2 Provider Medicaid Attestation Data

The data listed under the section Provider Medicaid Attestation Data is updatable by the

provider during attestation. If the Provider needs their paper check mailed to an address other than the one registered with CMS in the screen above, this is where it can be changed. Once the attestation is submitted by the provider, the data will become view only. These data fields are described below:


- Medicaid ID - This field only displays if you have multiple group Kentucky Medicaid Provider Numbers that are linked to the Payee NPI listed in your CMS registration. If so, you will need to select one of your Kentucky group Medicaid Numbers. **This Medicaid Number will be used for your incentive payments.**
- Mailing Address - The mailing address can be updated if the provider would like to give an alternate address from the one listed from CMS for correspondence. This change will only be used for mailing the provider's incentive payment. This will not change the address listed with CMS. If the mailing address is not current, this can delay receiving the incentive payment.
- Medicaid Provider Type - Please select the provider type from the list. This type should match the type of provider listed under your Kentucky Medicaid enrollment and your type of license.
- Were you assisted by a Regional Extension Center in Kentucky - Response to this question is required. If the response is yes, then please type the name of the person who assisted you during the attestation process.

7.4 Provider Eligibility Details Screen

EPs must enter two categories of information to complete the Eligibility Provider Details screen including Eligibility Details and Service Locations. Within the Eligibility Details section, the provider will enter data for Patient Volume and EHR Details.

7.4.1 Eligibility Details

Eligibility details section allows the provider to view or enter information depending on the source of the information and the status of the attestation. Information in this section includes patient volume and information about EHR use.

Provider Eligibility Details (Year 3 Attestation)	
Eligibility Details:	
All * fields are required fields.	
PATIENT VOLUME DETAILS	
Patient Volume:	1. Please indicate if your patient volume was calculated at a clinic or practice level for all Eligible professionals: * <input type="button" value="No"/>
	2. If yes, please enter the NPI of the clinic or group: <input type="text" value="0"/>
	3. For which program year are you applying? * <input type="button" value="2021"/>
	4. What is the time frame used for patient volume calculation? * <input type="button" value="Prior Calendar Year"/>
	5. Select the starting date of the 90-day period to calculate Medicaid encounter volume percentage: * <input type="text" value="10/1/2020"/> (mm/dd/yyyy)
	6. Medicaid patient encounters during this period (FQHCs/RHCs do NOT include uncompensated care volume in this count. Uncomp care volume needs to be included on the patient volume report.): * <input type="text" value="10"/>
	7. Total patient encounters during this period: * <input type="text" value="10"/>
	8. Medicaid patient volume percentage: 100.00%
EHR Details:	9. Enter the CMS EHR Certification ID of your EHR: * <input type="text" value="0015HTL9LXE85GB"/> 
	10. Indicate the status of your EHR: * <input checked="" type="radio"/> Meaningful User

Patient Volume

1. Indicate if patient volume was calculated at a clinic or practice level for all eligible professionals.
 - If submitting at the clinic or practice levels, **all** EPs from the clinic or practice must also submit their volume at the clinic or practice level for the same program year.
2. If submitting at the clinic or practice level, enter the NPI of the clinic or group.
3. The Program Year is display only from your selection made on the Home screen.
 - This should be the current year or the prior year, if the current date is on or before August 31.
4. Select the time frame used for patient volume calculation.
 - From the dropdown menu, select either the “Prior Calendar Year” or “Preceding 12 Months” of the date of attestation.
5. Select the starting date of the 90-day period to calculate the Medicaid encounter volume percentage. Enter as mm/dd/yyyy.
 - This date should be a continuous 90-day period.
6. Enter Medicaid patient encounters during this period.
7. Enter Total patient encounters during this period.

8. Medicaid patient volume percentage is auto-calculated based on the volume numbers entered and is displayed as a percentage with two decimal points.
 - o Volume thresholds are calculated using the EP’s total number of Medicaid member encounters for the 90-day period as the numerator and *all* patient encounters for the same EP over the same 90-day period as the denominator.

EHR Details

9. Enter the CMS EHR Certification ID
10. The status of your EHR is displayed only from your selection made on the Home screen.

7.4.2 Requesting KCHIP Report Data

To request a KCHIP Report, the provider will need to log into the attestation website at <https://prd.webapps.chfs.ky.gov/KYSLR/login.aspx>.

Click on the Reports link in the navigation menu and follow the instructions below to complete your request. Once the report is processed, an email will be sent to the email address provided at CMS registration.

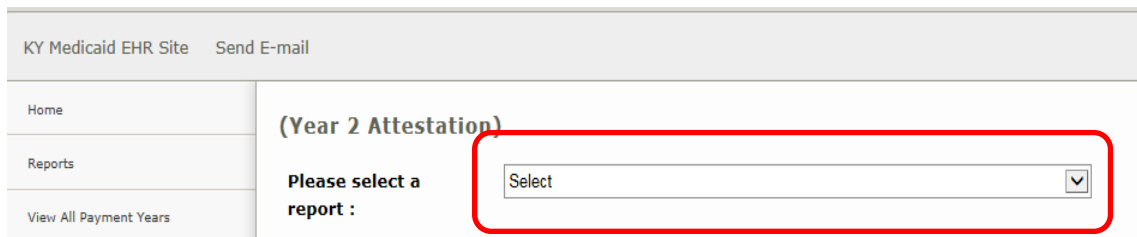
1st Year	2012	2013	2015	2017	2018	2019
2012	MU Stage 1 (90 Days)	MU Stage 1 (365 Days)	MU Mod Stage 2 (90 Days)	MU Mod Stage 2 or Stage 3 (90 Days)	MU Mod Stage 2 or Stage 3 (90 Days)	MU Stage 3 (365 Days)

The KCHIP data report will take approximately three hours to complete. Once the report is ready to be viewed, an email will be sent to the email address on file within the attestation. This email address can be verified on the ‘Registration Data’ screen of the attestation. If this email address is not correct, please go to the CMS Registration website to update this

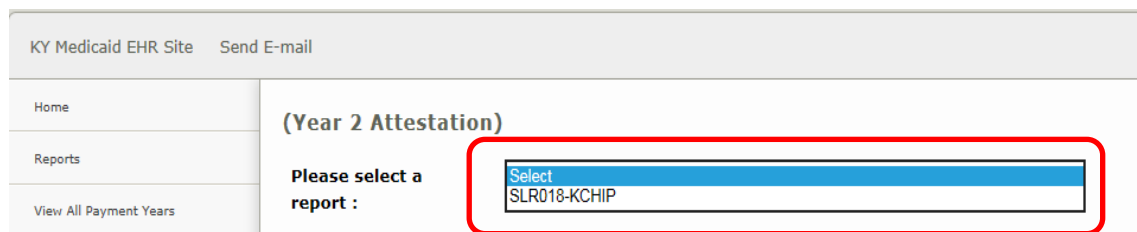
information. Email is our main form of communication with providers, so please take a moment to verify this information. Also, please be aware this update takes 24 hours to complete. Once you have received email notification that your KCHIP data is ready to be viewed, you will need to sign back into the attestation and click on the 'Reports' link located within the menu options located on the left hand side of the 'Home' screen and complete the following steps:



Step 1: Click the down arrow to select a report.



Step 2: Select 'SLR018-KCHIP'.



Step 3: Scroll down and locate the ‘Report Request Information’ heading. Click the ‘Select’ button next to the date you requested the report – also please confirm that the ‘Start Date and End Date’ are correct dates you will be attesting to for your 90 day patient volume.

REPORT REQUEST INFORMATION :						
	Date Requested	Report Name	NPI	Start Date	End Date	Status
Select	5/24/2018 9:26:04 AM	SLR018-KCHIP	2020202020	4/1/2017 12:00:00 AM	6/29/2017 12:00:00 AM	Completed - Successful

If KCHIP data is returned, subtract this total from the numerator value of your 90-day patient volume data, which is your total ‘Medicaid Encounters’. This adjusted total is what will be reported on line 6 on the ‘Eligibility Details’ page of the attestation. If ‘No Information Found’ is displayed, report your total Medicaid patients as you have calculated with no adjustments to line 6 on the ‘Eligibility Details’ page of the attestation and continue the completion of your attestation for review.

Provider Eligibility Details (Year 5 Attestation)

Eligibility Details:

All * fields are required fields.

PATIENT VOLUME DETAILS

Patient Volume: 1. Please indicate if your patient volume was calculated at a clinic or practice level for all Eligible professionals: *

2. If yes, please enter the NPI of the clinic or group:

3. For which program year are you applying? *

4. What is the time frame used for patient volume calculation? *

5. Select the starting date of the 90-day period to calculate Medicaid encounter volume percentage: *

6. Medicaid patient encounters during this period (FQHCs/RHCs do NOT include uncompensated care volume in this count. Uncomp care volume needs to be included on the patient volume report.): *

7. Total patient encounters during this period: *

8. Medicaid patient volume percentage: **100.00%**

EHR Details: 9. Enter the CMS EHR Certification ID of your EHR: *

10. Indicate the status of your EHR: * Meaningful User

7.4.3 Service Locations

In the Service location section, enter information about the service locations equipped with a certified EHR. Practice/Locations equipped with CEHRT can qualify for MU in the following ways:

- The CEHRT is permanently installed at the practice location.
- The CEHRT can be brought to the practice/location on a portable computing device.
- The CEHRT can be accessed remotely using computing devices at the practice/location.

Service Locations

The practice/location equipped with Certified EHR Technology (CEHRT) can be met in 3 ways:

1. CEHRT is permanently installed at the practice location
2. The CEHRT can be brought to the practice/location on a portable computing device
3. The CEHRT can be accessed remotely using computing devices at the practice/location

*Do you have multiple service locations? Yes No

*Enter the total number of locations:

*Enter the total number of locations with certified EHR Technology:

[Enter Service Location Address](#)

* Indicate below the service location(s) associated with this attestation that have Certified EHR Technology:

[Previous](#) [Next](#) [Save](#) [Cancel](#)

To complete this section, perform the following steps:

- Select Yes or No to indicate if there are multiple locations.
 - If Yes is selected, enter the total number of locations and the number of locations with a certified EHR.
 - A new section will open for entering an address. After entering the address, click on the Add button.
 - If No is selected, the total number of locations and locations with EHR technology will automatically populate with a 1.
- Enter the single service location address by clicking on the **Enter Service Location Address** button.

* Indicate below the service location(s) associated with this attestation that have Certified EHR Technology:

Address1:

Address 2:

City:

State:

Zip Code:

ZipCode Extension:

- Enter the Service location address information in the fields, then click the Add button.

* Indicate below the service location(s) associated with this attestation that have Certified EHR Technology:

Edit	Address Line 1	Address Line 2	City	State	Zip Code	Zip Code Extension	Delete
Modify	12 Millcreek Park		Frankfort	KY	40601		Delete
	<input type="text"/>	<input type="text"/>	<input type="text"/>	<input type="text"/>	<input type="text"/>	<input type="text"/>	<input type="button" value="ADD"/>

Once the address is added into the table, it can be modified or deleted, and more Service locations can be added.

- To edit or update a Service location, click the Modify link.
- To remove a Service location, click the Delete link.
- To add a new Service location, enter address information in to the fields and click the ADD button.

When final selections have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen.
- Click **Next** to move on to the next screen.
- Click **Save** to save selections and stay on the current screen.
- Click **Cancel** to remove selections and stay on the current screen.

7.5 Meaningful Use Questionnaire Screen

After entering the provider eligibility details, EPs will be directed to the MU Questionnaire screen. Here, the EP will enter the MU reporting period. The MU reporting period must be a 90-day consecutive period within the calendar year.

Meaningful Use Questionnaire (Year 3 Attestation)

Meaningful Use Questionnaire

*EHR Reporting Period Start Date: (mm/dd/yyyy)

*EHR Reporting Period End Date: (mm/dd/yyyy)

*Enter the percentage of unique patients who have structured data recorded on your certified EHR technology as of the reporting period above:

Previous **Next** **Save** **Cancel**

Enter responses for the following:

- Enter EHR Reporting Period Start Date
 - This is the starting date of the reporting period for the MU data.
- Enter EHR Reporting Period End Date
 - This is the end date of the reporting period for the MU data.
- Enter percentage of unique patients who have structured data recorded in the CEHRT as of the reporting period above.
 - This can be calculated by dividing the number of patients with structured data in your certified EHR by the total number of patients seen at service location(s) with CEHRT. Multiply by 100 to obtain the percentage. The amount of patients with structured data stored in your EHR should be at least 80%.

When final selections have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen.
- Click **Next** to move on to the next screen.
- Click **Save** to save selections and stay on the current screen.
- Click **Cancel** to remove selections and stay on the current screen.

8 Requirements for Meaningful Use Measures

Providers who are demonstrating MU for the Kentucky Medicaid EHR Program (Promoting Interoperability) will submit and attest to the following requirements:

- Medicaid provider eligibility requirements;
- Medicaid volume requirements;
- For Program Year 2021, Providers must select an EHR MU reporting period that is any continuous 90-day period within the current calendar year. Providers have until August 31, 2021 to attest to that EHR MU reporting period;
- For providers who work at multiple locations, 50% or more of patient encounters must occur at the location equipped with CEHRT;
- 80% of unique patients must have structured data recorded in the CEHRT;
- Must meet 8 MU Objectives for Stage 3;
- Must submit six eQMs. EPs must report on at least one outcome measure. If no outcome measure is relevant to his or her scope of practice, the EP must report on one high priority measure. If no high priority measures are relevant to their scope of practice, they may report on any six relevant measures.

Providers will be directed through the 8 MU Objectives listed below. The eQMs will not be available for attestation until the MU Objectives have been completed.

Meaningful Use Objectives

1. Protect Electronic Protected Health Information
2. Electronic Prescribing
3. Clinical Decision Support
4. Computerized Provider Order Entry
5. Patient Electronic Access to Health Information
6. Coordination of Care Through Patient Engagement
7. Health Information Exchange
8. Public Health and Clinical Data Registry Reporting
 - Immunization Registry Reporting
 - Syndromic Surveillance Reporting
 - Electronic Case Reporting
 - Public Health Registry Reporting
 - Clinical Data Registry Reporting

For additional information on MU Measures, please visit the CMS Web site <https://www.cms.gov/Regulations-and-Guidance/Legislation/EHRIncentivePrograms/index.html>.

8.1 Meaningful Use Menu Screen

The menu screen will only allow the user to select a group of measures, as they are available. For example, once the MU Core Objectives are completed, the Public Health Objectives will be active to select.

KY Medicaid EHR Incentive Program (Year 3 Attestation)

Please select a menu option below:

- [Meaningful Use Objectives](#)
- [Public Health Objective](#)
- [Electronic Clinical Quality Measures Submission](#)

Previous **Next**

Meaningful Use Core Objectives Link – Takes the EP to the first screen of the MU Core Objectives.

Public Health Objectives Link – Takes the EP to the first screen of the Public Health Objectives. This link is only active after the MU Core Objectives are completed.

Electronic Clinical Quality Measures Submission Link – Takes the EP to the first screen of the eQMs. This link is only active after the Public Health Objectives are completed.

If the EP does not wish to click the links for attestation, buttons at the bottom of the screen are available for navigation.

- Click **Previous** to go back to the previous screen.
- Click **Next** to move on to the next screen.

8.2 Meaningful Use Core Objectives – Stage 3

8.2.1 MU Core Objective 1 – Protect Patient Health Information

OBJECTIVE: Protect electronic protected health information (ePHI) created or maintained by the CEHRT through the implementation of appropriate technical, administrative, and physical safeguards.

Meaningful Use Objectives (Year 3 Attestation)

EP Objective 1 - Protect Patient Health Information

(* Red asterisk indicates a required field.)

Objective:	Protect electronic protected health information (ePHI) created or maintained by the CEHRT through the implementation of appropriate technical, administrative, and physical safeguards.
Measure:	Conduct or review a security risk analysis in accordance with the requirements under 45 CFR 164.308(a)(1), including addressing the security (including encryption) of data created or maintained by CEHRT in accordance with requirements under 45 CFR 164.312(a)(2)(iv) and 45 CFR 164.306(d)(3), implement security updates as necessary, and correct identified security deficiencies as part of the provider's risk management process.
<p>It is acceptable for the security risk analysis to be conducted outside the EHR reporting period. However, the analysis must be unique for each EHR reporting period, the scope must include the full EHR reporting period, and it must be conducted within the calendar year of the EHR reporting period</p> <p>Complete the following information:</p> <p>*Have you conducted or plan to conduct a security risk analysis in the calendar year 2021 in accordance to the requirements?</p> <p><input checked="" type="radio"/> Yes <input type="radio"/> No</p> <p>*Enter the date that the security risk analysis has been or will be conducted:</p> <p><input type="text" value="01/01/2021"/></p>	
<p style="text-align: right;"> <input type="button" value="Previous"/> <input type="button" value="Next"/> <input type="button" value="Save"/> <input type="button" value="Cancel"/> </p>	

In order for EPs to meet the objectives, they must be able to satisfy the measure.

To satisfy the Measure, select a response to the question and enter the date that the security risk analysis has been or will be conducted.

- If No is selected, upon navigation, a message will pop up stating that the entry for the measure does not meet the threshold to qualify for an incentive payment.

When final selections have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen. Selections will not be saved.
- Click **Next** to move on to the next screen. Selections will be saved.
- Click **Save** to save selections and stay on the current screen.
- Click **Cancel** to remove selections and stay on the current screen.

8.2.2 MU Core Objective 2 – Electronic Prescribing (eRX)

OBJECTIVE: Generate and transmit permissible prescriptions electronically.

Meaningful Use Objectives (Year 2 Attestation)

EP Objective 2 - Electronic Prescribing
 (*) Red asterisk indicates a required field.

Objective: Generate and transmit permissible prescriptions electronically (eRx).

Measure: More than 60% of all permissible prescriptions written by the EP are queried for a drug formulary and transmitted electronically using CEHRT.

***Patient Records:** Please select whether the data used to support the measure was extracted from all patient records or only from patient records maintained using certified EHR technology.

This data was extracted from ALL patient records not just those maintained using certified EHR technology.

This data was extracted only from patient records maintained using certified EHR technology.

EXCLUSION 1: Any EP who writes fewer than 100 permissible prescriptions during the EHR reporting period.

***Does this exclusion apply to you?**

Yes No

EXCLUSION 2: Any EP who does not have a pharmacy within his or her organization and there are no pharmacies that accept electronic prescriptions within 10 miles of the EPs practice location at the start of his or her EHR reporting period.

***Does this exclusion apply to you?**

Yes No

Complete the following information:

Numerator = The number of prescriptions in the denominator generated, queried for a drug formulary, and transmitted electronically using CEHRT.

Denominator = Number of prescriptions written for drugs requiring a prescription in order to be dispensed other than controlled substances during the EHR reporting period; or number of prescriptions written for drugs requiring a prescription in order to be dispensed during the EHR reporting period.

***Numerator :** ***Denominator :**

In order for EPs to meet the objective, they must satisfy the measure by claiming the exclusion or meeting the threshold.

To satisfy the Measure, make two selections.

- First, respond as to whether data used to support the measure was extracted from ALL patient records or only from patient records maintained using certified EHR technology.
- Second, respond to Exclusion 1.
 - If No is selected, respond to Exclusion 2.

- If No is selected, enter a whole number into both the Numerator and Denominator boxes. The Numerator/Denominator must be over 60% in order to successfully attest to the measure.

When final selections have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen. Selections will not be saved.
- Click **Next** to move on to the next screen. Selections will be saved.
- Click **Save** to save selections and stay on the current screen.
- Click **Cancel** to remove selections and stay on the current screen.

8.2.3 MU Core Objective 3 – Clinical Decision Support

OBJECTIVE: Implement clinical decision support (CDS) interventions focused on improving performance on high-priority health conditions.

Meaningful Use Objectives (Year 4 Attestation)

EP Objective 3 - Clinical Decision Support
 (*) Red asterisk indicates a required field.

In order to meet this objective, an EP must satisfy both measures for this objective through a combination of meeting the thresholds and exclusions.

Objective:	Implement clinical decision support (CDS) interventions focused on improving performance on high-priority health conditions.
Measure 1:	Implement five CDS interventions related to four or more clinical quality measures (CQMs) at a relevant point in patient care for the entire EHR reporting period. Absent four CQMs related to an EPs scope of practice or patient population, the CDS interventions must be related to high-priority health conditions.

Complete the following information:

*Have you implemented five CDS interventions related to four or more clinical quality measures at a relevant point in patient care for the entire EHR reporting period and absent four clinical quality measures related to your scope of practice or patient population, were the clinical decision support interventions related to high-priority health conditions?

Yes No

If you have implemented four or more CQMs related to five clinical decision support interventions for the entire EHR reporting period, please enter below.

Testing

Measure 2: Enable and implement the functionality for drug-drug and drug-allergy interaction checks for the entire EHR reporting period.

EXCLUSION: Any EP who writes fewer than 100 medication orders during the EHR reporting period may take an exclusion.

*Does this exclusion apply to you?

Yes No

Complete the following information:

*Has the EP enable and implement the functionality for drug-drug and drug-allergy interaction checks for the entire EHR reporting period?

Yes No

Previous Next Save Cancel

EP must satisfy both measures in order to meet the objective.

To satisfy Measure 1, respond to the question.

- If Yes is selected, enter four or more clinical quality measures related to the five clinical decision support interventions implemented.
- If No is selected, a pop up window stating the entry for the Measure does not qualify for an incentive payment.

To satisfy Measure 2, respond to the Exclusion.

- If No is selected, respond to the question for measure 2.
 - If No is selected in response to the question for measure 2, a pop up window stating the entry for Measure 2 does not qualify for an incentive payment.

When final selections have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen. Selections will not be saved.
- Click **Next** to move on to the next screen. Selections will be saved.
- Click **Save** to save selections and stay on the current screen.
- Click **Cancel** to remove selections and stay on the current screen.

8.2.4 MU Core Objective 4 – Computerized Provider Order Entry

OBJECTIVE: Use CPOE for medication, laboratory, and diagnostic imaging orders directly entered by any licensed healthcare professional, credentialed medical assistant, or a medical staff member credentialed to and performing the equivalent duties of a credentialed medical assistant, who can enter orders into the medical record per state, local, and professional guidelines.

Meaningful Use Objectives (Year 2 Attestation)

EP Objective 4 - Computerized Provider Order Entry

(*) Red asterisk indicates a required field.

In order to meet this objective, an EP must satisfy all three measures for this objective through a combination of meeting the thresholds and exclusions.

Objective: Use CPOE for medication, laboratory, and diagnostic imaging orders directly entered by any licensed healthcare professional, credentialed medical assistant, or a medical staff member credentialed to and performing the equivalent duties of a credentialed medical assistant, who can enter orders into the medical record per state, local, and professional guidelines.

Measure 1: More than 60% of medication orders created by the EP during the EHR reporting period are recorded using computerized provider order entry.

***Patient Records:** Please select whether the data used to support the measure was extracted from all patient records or only from patient records maintained using CEHRT.

- This data was extracted from ALL patient records not just those maintained using certified EHR technology.
- This data was extracted only from patient records maintained using certified EHR technology.

EXCLUSION: Any EP who writes fewer than 100 medication orders during the EHR reporting period.

* Does this exclusion apply to you?

- Yes No

Complete the following information:

Numerator = The number of orders in the denominator recorded using CPOE.

Denominator = Number of medication orders created by the EP during the EHR reporting period.

***Numerator:** ***Denominator:**

Measure 2: More than 60% of laboratory orders created by the EP during the EHR reporting period are recorded using computerized provider order entry.

***Patient Records:** Please select whether the data used to support the measure was extracted from all patient records or only from patient records maintained using CEHRT.

- This data was extracted from ALL patient records not just those maintained using certified EHR technology.
- This data was extracted only from patient records maintained using certified EHR technology.

EXCLUSION: Any EP who writes fewer than 100 laboratory orders during the EHR reporting period.

* Does this exclusion apply to you?

- Yes No

Complete the following information:

Numerator = The number of orders in the denominator recorded using CPOE.

Denominator = Number of laboratory orders created by the EP during the EHR reporting period.

***Numerator:** ***Denominator:**

Measure 3: More than 60% of diagnostic imaging orders created by the EP during the EHR reporting period are recorded using computerized provider order entry.

***Patient Records:** Please select whether the data used to support the measure was extracted from all patient records or only from patient records maintained using CEHRT.

This data was extracted from ALL patient records not just those maintained using certified EHR technology.

This data was extracted only from patient records maintained using certified EHR technology.

EXCLUSION: Any EP who writes fewer than 100 diagnostic imaging orders during the EHR reporting period.

*** Does this exclusion apply to you?**

Yes No

Complete the following information:

Numerator = The number of orders in the denominator recorded using CPOE.

Denominator = Number of diagnostic imaging orders created by the EP during the EHR reporting period.

***Numerator:** ***Denominator:**

An EP, through a combination of meeting the thresholds and exclusions (or both), must satisfy all three measures for this objective.

To satisfy Measure 1, make two selections.

- First, respond as to whether data used to support the measure was extracted from ALL patient records or only from patient records maintained using certified EHR technology.
- Second, respond to Exclusion.
 - If No is selected, enter a whole number into both the Numerator and Denominator boxes. The Numerator/Denominator must be over 60% in order to successfully attest to the measure.

To satisfy Measure 2, make two selections.

- First, respond as to whether data used to support the measure was extracted from ALL patient records or only from patient records maintained using certified EHR technology.

- Second, respond to the Exclusion.
 - If No is selected, enter a whole number into both the Numerator and Denominator boxes. The Numerator/Denominator must be over 60% in order to successfully attest to the measure.

To satisfy Measure 3, make two selections.

- First, respond as to whether data used to support the measure was extracted from ALL patient records or only from patient records maintained using certified EHR technology.
- Second, respond to the Exclusion.
 - If No is selected, enter a whole number into both the Numerator and Denominator boxes. The Numerator/Denominator must be over 60% in order to successfully attest to the measure.

When final selections have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen. Selections will not be saved.
- Click **Next** to move on to the next screen. Selections will be saved.
- Click **Save** to save selections and stay on the current screen.
- Click **Cancel** to remove selections and stay on the current screen.

8.2.5 MU Core Objective 5 – Patient Electronic Access to Health Information

OBJECTIVE: The EP provides patients (or patient-authorized representative) with timely electronic access to their health information and patient-specific education.

Meaningful Use Objectives (Year 2 Attestation)

EP Objective 5 - Patient Electronic Access to Health Information

(* Red asterisk indicates a required field.

In order to meet this objective, an EP must satisfy both measures for this objective through a combination of meeting the thresholds and exclusions.

Objective: The EP provides patients (or patient-authorized representatives) with timely electronic access to their health information and patient-specific education.

Measure 1: For more than 80% of all unique patients seen by the EP: (1) The patient (or the patient-authorized representative) is provided timely access to view online, download, and transmit his or her health information; and (2) The provider ensures the patient's health information is available for the patient (or patient-authorized representative) to access using any application of their choice that is configured to meet the technical specifications of the API in the provider's CEHRT.

EXCLUSION 1: Any EP with no office visits during the EHR reporting period.

*Does this exclusion apply to you?

Yes No

EXCLUSION 2: Any EP that conducts 50% or more of their patient encounters in a county that does not have 50% or more of its housing units with 4Mbps broadband availability according to the latest information available from the FCC on the first day of the EHR reporting period may exclude the measure.

*Does this exclusion apply to you?

Yes No

Complete the following information:

Numerator = The number of patients in the denominator (or patient-authorized representative) who are provided timely access to health information to view online, download, and transmit to a third party and to access using an application of their choice that is configured to meet the technical specifications of the API in the EP's CEHRT.

Denominator = The number of unique patients seen by the EP during the EHR reporting period.

*Numerator: *Denominator:

Measure 2: The EP must use clinically relevant information from CEHRT to identify patient-specific educational resources and provide electronic access to those materials to more than 35% of unique patients seen by the EP during the EHR reporting period.

EXCLUSION 1: Any EP with no office visits during the EHR reporting period.

*Does this exclusion apply to you?

Yes No

EXCLUSION 2: Any EP that conducts 50% or more of their patient encounters in a county that does not have 50% or more of its housing units with 4Mbps broadband availability according to the latest information available from the FCC on the first day of the EHR reporting period may exclude the measure.

*Does this exclusion apply to you?

Yes No

Complete the following information:

Numerator = The number of patients in the denominator who were provided electronic access to patient-specific educational resources using clinically relevant information identified from CEHRT during the EHR reporting period.

Denominator = The number of unique patients seen by the EP during the EHR reporting period.

*Numerator: *Denominator:

Previous
Next
Save
Cancel

EP must satisfy both measures in order to meet the objective.

To satisfy Measure 1, respond to Exclusion 1.

- If No is selected, respond to Exclusion 2.
 - If No is selected, enter a whole number into both the Numerator and Denominator boxes. The Numerator/Denominator must be over 80% in order to successfully attest to the measure.

To satisfy Measure 2, respond to Exclusion 1.

- If No is selected, respond to Exclusion 2.
 - If No is selected, enter a whole number into both the Numerator and Denominator boxes. The Numerator/Denominator must be over 35% in order to successfully attest to the measure.

When final selections have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen. Selections will not be saved.
- Click **Next** to move on to the next screen. Selections will be saved.
- Click **Save** to save selections and stay on the current screen.
- Click **Cancel** to remove selections and stay on the current screen.

8.2.6 MU Core Objective 6 – Coordination of Care through Patient Engagement

OBJECTIVE: Use CEHRT to engage with patients or their authorized representatives about the patients’ care.

Meaningful Use Objectives (Year 2 Attestation)

EP Objective 6 - Coordination of Care through Patient Engagement

(*) Red asterisk indicates a required field.

In order to meet this objective, EPs must attest to all three measures and meet the threshold for two measures. If the EP meets the criteria for exclusion from two measures, they must meet the threshold for the one remaining measure. If they meet the criteria for exclusion from all three measures, they may be excluded from meeting this objective.

Objective: Use CEHRT to engage with patients or their authorized representatives about the patient's care.

Measure 1: More than 5% of all unique patients (or their authorized representatives) seen by the EP actively engage with the EHR made accessible by the EP and either: (1) View, download, or transmit to a third party their health information; or (2) Access their health information through an API that can be used by applications chosen by the patient and configured to the API in the EP's CEHRT; or (3) A combination of (1) and (2).

<p>EXCLUSION 1: Any EP who has no office visits during the EHR reporting period.</p> <p>* Does this exclusion apply to you?</p> <p><input type="radio"/> Yes <input checked="" type="radio"/> No</p>
<p>EXCLUSION 2: Any EP that conducts 50% or more of their patient encounters in a county that does not have 50% or more of its housing units with 4Mbps broadband availability according to the latest information available from the FCC on the first day of the EHR reporting period may exclude the measure.</p> <p>* Does this exclusion apply to you?</p> <p><input type="radio"/> Yes <input checked="" type="radio"/> No</p>
<p>Complete the following information:</p> <p>Numerator = The number of unique patients (or their authorized representatives) in the denominator who have viewed online, downloaded, or transmitted to a third party the patient's health information during the EHR reporting period and the number of unique patients (or their authorized representatives) in the denominator who have accessed their health information through the use of an API during the EHR reporting period.</p> <p>Denominator = Number of unique patients seen by the EP during the EHR reporting period.</p> <p>*Numerator : <input style="width: 80px;" type="text" value="30"/> *Denominator : <input style="width: 80px;" type="text" value="30"/></p>
<p>Measure 2: For more than 5% of all unique patients seen by the EP during the EHR reporting period, a secure message was sent using the electronic messaging function of CEHRT to the patient (or the patient-authorized representative), or in response to a secure message sent by the patient or their authorized representative.</p>
<p>EXCLUSION 1: Any EP who has no office visits during the EHR reporting period.</p> <p>* Does this exclusion apply to you?</p> <p><input type="radio"/> Yes <input checked="" type="radio"/> No</p>
<p>EXCLUSION 2: Any EP that conducts 50% or more of their patient encounters in a county that does not have 50% or more of its housing units with 4Mbps broadband availability according to the latest information available from the FCC on the first day of the EHR reporting period may exclude the measure.</p> <p>* Does this exclusion apply to you?</p> <p><input type="radio"/> Yes <input checked="" type="radio"/> No</p>

Complete the following information:

Numerator = The number of patients in the denominator for whom a secure electronic message is sent to the patient (or patient-authorized representative) or in response to a secure message sent by the patient (or patient-authorized representative), during the EHR reporting period.

Denominator = Number of unique patients seen by the EP during the EHR reporting period.

***Numerator :** ***Denominator :**

Measure 3: Patient generated health data or data from a non-clinical setting is incorporated into the CEHRT for more than 5% of all unique patients seen by the EP during the EHR reporting period.

EXCLUSION 1: Any EP who has no office visits during the EHR reporting period.

*** Does this exclusion apply to you?**

Yes No

EXCLUSION 2: Any EP that conducts 50% or more of their patient encounters in a county that does not have 50% or more of its housing units with 4Mbps broadband availability according to the latest information available from the FCC on the first day of the EHR reporting period may exclude the measure.

*** Does this exclusion apply to you?**

Yes No

Complete the following information:

Numerator = The number of patients in the denominator for whom data from non-clinical settings, which may include patient generated health data, is captured through the CEHRT into the patient record during the EHR reporting period.

Denominator = Number of unique patients seen by the EP during the EHR reporting period.

***Numerator :** ***Denominator :**

An EP must attest to all three measures and meet the threshold for two measures for this objective. If the EP meets the criteria for exclusion from two measures, they must meet the threshold for the one remaining measure. If they meet the criteria for exclusion from all three measures, they may be excluded from meeting this objective.

To satisfy Measure 1, respond to Exclusion 1.

- If No is selected, respond to Exclusion 2.
 - If No is selected, enter a whole number into both the Numerator and Denominator boxes. The Numerator/Denominator must be over 5% in order to successfully attest to the measure.

To satisfy Measure 2, respond to Exclusion 1.

- If No is selected, respond to Exclusion 2.
 - If No is selected, enter a whole number into both the Numerator and Denominator boxes. The Numerator/Denominator must be over 5% in order to successfully attest to the measure.

To satisfy Measure 3, respond to Exclusion 1.

- If No is selected, respond to Exclusion 2.
 - If No is selected, enter a whole number into both the Numerator and Denominator boxes. The Numerator/Denominator must be over 5% in order to successfully attest to the measure.

When final selections have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen. Selections will not be saved.
- Click **Next** to move on to the next screen. Selections will be saved.
- Click **Save** to save selections and stay on the current screen.
- Click **Cancel** to remove selections and stay on the current screen.

8.2.7 MU Core Objective 7 – Health Information Exchange

OBJECTIVE: The EP provides a summary of care record when transitioning or referring their patient to another setting of care, receives or retrieves a summary of care record upon the receipt of a transition or referral or upon the first patient encounter with a new patient, and incorporates summary of care information from other providers into their EHR using the functions of CEHRT.

Meaningful Use Objectives (Year 2 Attestation)

EP Objective 7 - Health Information Exchange

(*) Red asterisk indicates a required field.

In order to meet this objective, EPs must attest to all three measures and meet the threshold for two measures. If the EP meets the criteria for exclusion from two measures, they must meet the threshold for the one remaining measure. If they meet the criteria for exclusion from all three measures, they may be excluded from meeting this objective.

Objective: The EP provides a summary of care record when transitioning or referring their patient to another setting of care, receives or retrieves a summary of care record upon the receipt of a transition or referral or upon the first patient encounter with a new patient, and incorporates summary of care information from other providers into their EHR using the functions of CEHRT.

Measure 1: For more than 50% of transitions of care and referrals, the EP that transitions or refers their patient to another setting of care or provider of care: (1) Creates a summary of care record using CEHRT; and (2) Electronically exchanges the summary of care record.

EXCLUSION 1: Any EP who transfers a patient to another setting or refers a patient to another provider fewer than 100 times during the EHR reporting period.

* Does this exclusion apply to you?

Yes No

EXCLUSION 2: Any EP that conducts 50% or more of their patient encounters in a county that does not have 50% or more of its housing units with 4Mbps broadband availability according to the latest information available from the FCC on the first day of the EHR reporting period.

* Does this exclusion apply to you?

Yes No

Complete the following information:

Numerator = The number of transitions of care and referrals in the denominator where a summary of care record was created using CEHRT and exchanged electronically.

Denominator = Number of transitions of care and referrals during the EHR reporting period for which the EP was the transferring or referring provider.

***Numerator** : ***Denominator** :

Measure 2: For more than 40% of transitions or referrals received and patient encounters in which the EP has never encountered the patient before, they incorporate into the patient's EHR an electronic summary of care document.

EXCLUSION 1: Any EP for whom the total transitions or referrals received and patient encounters in which they have never encountered the patient, is fewer than 100 during the EHR reporting period.

* Does this exclusion apply to you?

Yes No

EXCLUSION 2: Any EP that conducts 50% or more of their patient encounters in a county that does not have 50% or more of its housing units with 4Mbps broadband availability according to the latest information available from the FCC on the first day of the EHR reporting period.

* Does this exclusion apply to you?

Yes No

Complete the following information:

Numerator = Number of patient encounters in the denominator where an electronic summary of care record received is incorporated by the provider into the CEHRT.

Denominator = Number of patient encounters during the EHR reporting period for which an EP was the receiving party of a transition or referral or has never before encountered the patient and for which an electronic summary of care record is available.

*Numerator : *Denominator :

Measure 3: For more than 80% of transitions or referrals received and patient encounters in which the EP has never encountered the patient before, they perform a clinical information reconciliation. The EP must implement clinical information reconciliation for the following three clinical information sets: (1) Medication. Review of the patient's medication, including the name, dosage, frequency, and route of each medication. (2) Medication allergy. Review of the patient's known medication allergies. (3) Current Problem list. Review of the patient's current and active diagnoses.

EXCLUSION : Any EP for whom the total transitions or referrals received and patient encounters in which they have never before encountered the patient, is fewer than 100 during the EHR reporting period.

* Does this exclusion apply to you?

Yes No

Complete the following information:

Numerator = The number of transitions of care or referrals in the denominator where the following three clinical information reconciliations were performed: medication list, medication allergy list, and current problem list.

Denominator = Number of transitions of care or referrals during the EHR reporting period for which the EP was the recipient of the transition or referral or has never before encountered the patient.

*Numerator : *Denominator :

An EP must attest to all three measures and meet the threshold for two measures for this objective. If the EP meets the criteria for exclusion from two measures, they must meet the threshold for the one remaining measure. If they meet the criteria for exclusion from all three measures, they may be excluded from meeting this objective.

To satisfy Measure 1, the EP must respond to Exclusion 1.

- If No is selected, respond to Exclusion 2.
 - If No is selected, enter a whole number into both the Numerator and Denominator boxes. The Numerator/Denominator must be over 50% in order to successfully attest to the measure.

To satisfy Measure 2, respond to Exclusion 1.

- If No is selected, respond to Exclusion 2.
 - If No is selected, the EP must enter a whole number into both the Numerator and Denominator boxes. The Numerator/Denominator must be over 40% in order to successfully attest to the measure.

To satisfy Measure 3, respond to the Exclusion.

- If No is selected, respond to Exclusion 3.
 - If No is selected, enter a whole number into both the Numerator and Denominator boxes. The Numerator/Denominator must be over 80% in order to successfully attest to the measure.

When final selections have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen. Selections will not be saved.
- Click **Next** to move on to the next screen. Selections will be saved.
- Click **Save** to save selections and stay on the current screen.
- Click **Cancel** to remove selections and stay on the current screen.

8.2.8 MU Core Objective 8 – Public Health and Clinical Data Registry Reporting

OBJECTIVE: The EP is in active engagement with a public health agency (PHA) or clinical data registry (CDR) to submit electronic public health data in a meaningful way using certified electronic health record technology (CEHRT), except where prohibited, and in accordance with applicable law and practice.

In order to meet this objective, EPs need to meet two of the five measures. Exclusions do not count toward meeting the objective. If the EP qualifies for multiple exclusions and the remaining number of measures available is less than two, the EP can meet the objective by meeting all of the remaining measures available and claiming the applicable exclusions. If no measures remain available, you can meet the objective by claiming applicable exclusions for all measures. Available measures are ones for which the EP does not qualify for an exclusion.

8.2.8.1 Measure 1: Immunization Registry Reporting

MEASURE: The EP 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).

Public Health Objective Measures (Year 6 Attestation)

Immunization Registry Reporting

Objective

The EP is in active engagement with a public health agency (PHA) or clinical data registry (CDR) to submit electronic public health data in a meaningful way using CEHRT, except where prohibited, and in accordance with applicable law and practice.

Measure

The EP is 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).

*Would you like to attest to this measure?

Yes No

EXCLUSION 1: Does not administer immunizations to any of the populations for which data is collected by their jurisdiction's immunization registry or immunization information system (IIS) during the EHR reporting period.

*Does this exclusion apply to you?

Yes No

EXCLUSION 2: EP practices in a jurisdiction for which no immunization registry or immunization information system (IIS) is capable of accepting the specific standards required to meet the CEHRT definition at the start of the EHR reporting period.

*Does this exclusion apply to you?

Yes No

EXCLUSION 3: The EP practices in a jurisdiction where no immunization registry or immunization information system (IIS) has declared readiness to receive immunization data as of six months prior to the start of the EHR reporting period.

*Does this exclusion apply to you?

Yes No

Active Engagement Options:**Active Engagement Option 1 - Completed Registration to Submit Data:**

The EP 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 EHR reporting period; and the EP is awaiting an invitation from the PHA or CDR to begin testing and validation. This option allows EPs to meet the measure when the PHA or the CDR has limited resources to initiate the testing and validation process. EPs that have registered in previous years do not need to submit an additional registration to meet this requirement for each EHR reporting period.

Active Engagement Option 2 - Testing and Validation:

The EP is in the process of testing and validation of the electronic submission of data. EPs must respond to requests from the PHA or, where applicable, the CDR within 30 days; failure to respond twice within an EHR reporting period would result in that EP not meeting the measure.

Active Engagement Option 3 - Production:

The EP has completed testing and validation of the electronic submission and is electronically submitting production data to the PHA or CDR.

Please select the applicable active engagement option (may only select one).

- Option1
- Option2
- Option3

Previous

Next

Save

Cancel

To satisfy the Measure, respond to the question.

- If Yes is selected, respond to Exclusion 1.
- If No is selected, respond to Exclusion 2.
- If No is selected, respond to Exclusion 3.
 - If No is selected, select the applicable Active Engagement Option.

When final selections have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen. Selections will not be saved.
- Click **Next** to move on to the next screen. Selections will be saved.
- Click **Save** to save selections and stay on the current screen.
- Click **Cancel** to remove selections and stay on the current screen.

8.2.8.2 Measure 2: Syndromic Surveillance Reporting

MEASURE: The EP is in active engagement with a public health agency to submit syndromic surveillance data.

Public Health Objective Measures (Year 2 Attestation)

Syndromic Surveillance Reporting

Objective

The EP is in active engagement with a public health agency (PHA) or clinical data registry (CDR) to submit electronic public health data in a meaningful way using CEHRT, except where prohibited, and in accordance with applicable law and practice.

Measure

The EP is in active engagement with a PHA to submit syndromic surveillance data.

*Would you like to attest to this measure?

Yes No

EXCLUSION 1: An EP is not in a category of providers from which ambulatory syndromic surveillance data is collected by their jurisdiction's syndromic surveillance system.

*Does this exclusion apply to you?

Yes No

EXCLUSION 2: The EP practices in a jurisdiction for which no PHA is capable of receiving electronic syndromic surveillance data from EPs in the specific standards required to meet the CEHRT definition at the start of the EHR reporting period.

*Does this exclusion apply to you?

Yes No

EXCLUSION 3: The EP practices in a jurisdiction where no PHA has declared readiness to receive syndromic surveillance data from EPs as of six months prior to the start of the EHR reporting period.

*Does this exclusion apply to you?

Yes No

Active Engagement Options:**Active Engagement Option 1 - Completed Registration to Submit Data:**

The EP 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 EHR reporting period; and the EP is awaiting an invitation from the PHA or CDR to begin testing and validation. This option allows EPs to meet the measure when the PHA or the CDR has limited resources to initiate the testing and validation process. EPs that have registered in previous years do not need to submit an additional registration to meet this requirement for each EHR reporting period.

Active Engagement Option 2 - Testing and Validation:

The EP is in the process of testing and validation of the electronic submission of data. EPs must respond to requests from the PHA or, where applicable, the CDR within 30 days; failure to respond twice within an EHR reporting period would result in that EP not meeting the measure.

Active Engagement Option 3 - Production:

The EP has completed testing and validation of the electronic submission and is electronically submitting production data to the PHA or CDR.

Please select the applicable active engagement option (may only select one).

- Option1
- Option2
- Option3






To satisfy the Measure, respond to the question.

- If Yes is selected, respond to Exclusion 1.
- If No is selected, respond to Exclusion 2.
- If No is selected, respond to Exclusion 3.
 - If No is selected, select the applicable Active Engagement Option.

When final selections have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen. Selections will not be saved.
- Click **Next** to move on to the next screen. Selections will be saved.
- Click **Save** to save selections and stay on the current screen.
- Click **Cancel** to remove selections and stay on the current screen.

8.2.8.3 Measure 3: Electronic Case Reporting

MEASURE: The EP is in active engagement with a public health agency to submit case reporting of reportable conditions.

Public Health Objective Measures (Year 6 Attestation)

Electronic Case Reporting

Objective

The EP is in active engagement with a public health agency (PHA) or clinical data registry (CDR) to submit electronic public health data in a meaningful way using CEHRT, except where prohibited, and in accordance with applicable law and practice.

Measure

The EP is an active engagement with a PHA to submit case reporting of reportable conditions.

*Would you like to attest to this measure?

Yes No

EXCLUSION 1: Does not treat or diagnose any reportable diseases for which data is collected by their jurisdiction's reportable disease system during the EHR reporting period.

*Does this exclusion apply to you?

Yes No

EXCLUSION 2: The EP practices in a jurisdiction for which no PHA is capable of receiving electronic case reporting data in the specific standards required to meet the CEHRT definition at the start of the EHR reporting period.

*Does this exclusion apply to you?

Yes No

EXCLUSION 3: The EP practices in a jurisdiction where no PHA has declared readiness to receive electronic case reporting data as of six months prior to the start of the EHR reporting period.

*Does this exclusion apply to you?

Yes No

Active Engagement Options:**Active Engagement Option 1- Completed Registration to Submit Data:**

The EP 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 EHR reporting period; and the EP is awaiting an invitation from the PHA or CDR to begin testing and validation. This option allows EPs to meet the measure when the PHA or the CDR has limited resources to initiate the testing and validation process. EPs that have registered in previous years do not need to submit an additional registration to meet this requirement for each EHR reporting period.

Active Engagement Option 2 - Testing and Validation:

The EP is in the process of testing and validation of the electronic submission of data. EPs must respond to requests from the PHA or, where applicable, the CDR within 30 days; failure to respond twice within an EHR reporting period would result in that EP not meeting the measure.

Active Engagement Option 3 - Production:

The EP has completed testing and validation of the electronic submission and is electronically submitting production data to the PHA or CDR.

Please select the applicable active engagement option (may only select one).

- Option1
- Option2
- Option3

Previous

Next

Save

Cancel

To satisfy the Measure, respond to the question.

- If Yes is selected, respond to Exclusion 1.
- If No is selected, respond to Exclusion 2.
- If No is selected, respond to Exclusion 3.
 - If No is selected, select the applicable Active Engagement Option.

When final selections have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen. Selections will not be saved.
- Click **Next** to move on to the next screen. Selections will be saved.
- Click **Save** to save selections and stay on the current screen.
- Click **Cancel** to remove selections and stay on the current screen.

8.2.8.4 Measure 4: Public Health Registry Reporting

MEASURE: The EP is in active engagement with a public health agency to submit data to public health registries.

Public Health Objective Measures (Year 2 Attestation)

Public Health Registry Reporting

Objective

The EP is in active engagement with a public health agency (PHA) or clinical data registry (CDR) to submit electronic public health data in a meaningful way using CEHRT, except where prohibited, and in accordance with applicable law and practice.

Measure

The EP is in active engagement with a PHA to submit data to public health registries.

*Would you like to attest to this measure?

Yes No

EXCLUSION 1: Does not diagnose or directly treat any disease or condition associated with a public health registry in their jurisdiction during the EHR reporting period.

*Does this exclusion apply to you?

Yes No

EXCLUSION 2: The EP practices in a jurisdiction for which no PHA can accept electronic registry transactions in the specific standards required to meet the CEHRT definition at the start of the EHR reporting period.

*Does this exclusion apply to you?

Yes No

EXCLUSION 3: The EP practices in a jurisdiction where no PHA for which the EP is eligible to submit data has declared readiness to receive electronic registry transactions as of six months prior to the start of the EHR reporting period.

*Does this exclusion apply to you?

Yes No

Active Engagement Options:

Active Engagement Option 1 - Completed Registration to Submit Data:

The EP 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 EHR reporting period; and the EP is awaiting an invitation from the PHA or CDR to begin testing and validation. This option allows EPs to meet the measure when the PHA or the CDR has limited resources to initiate the testing and validation process. EPs that have registered in previous years do not need to submit an additional registration to meet this requirement for each EHR reporting period.

Active Engagement Option 2 - Testing and Validation:

The EP is in the process of testing and validation of the electronic submission of data. EPs must respond to requests from the PHA or, where applicable, the CDR within 30 days; failure to respond twice within an EHR reporting period would result in that EP not meeting the measure.

Active Engagement Option 3 - Production:

The EP has completed testing and validation of the electronic submission and is electronically submitting production data to the PHA or CDR.

*** Please select the applicable active engagement option (may only select one).**

- Option1
- Option2
- Option3

Instructions:

Provider may report to more than one public health registry and may count public health registry reporting more than one time to meet the required number of measures for the objective. You may enter as many registries as you wish but only two will be counted towards the objective.

A provider may count a public health registry if the provider achieved the phase of active engagement defined under Active Engagement Option 3: Production, including production data submission with the specialized registry in a prior year under the applicable requirements of the EHR Incentive Programs in 2015 through 2018.

To report the first Public Health Registry, enter the information in the text box, then click 'Add'. To report the additional Public Health Registries, select the active engagement option applicable for the next registry you are reporting, enter the information in the text box and click 'Add'. Public Health Registry information you are attesting to will be displayed in the Registry table below.

*** Please add the public health data registry below:**

- Other

LIST OF SPECIALIZED REGISTRIES YOU ADDED:

Type of Registry	Active Engagement Option	Description	Edit	Delete
Other	1	Brain	Edit	Delete

[Add](#)

[Previous](#)
[Next](#)
[Save](#)
[Cancel](#)

To satisfy the Measure, respond to the question.

- If Yes is selected, respond to Exclusion 1.
- If No is selected, respond to Exclusion 2.
- If No is selected, respond to Exclusion 3.
 - If No is selected, make two selections.
 - Select the applicable Active Engagement Option for each registry.
 - Add each public health registry to the table.
 - If Other is selected, type the name of the registry into the text box. Click **Add** to add it to the table.
 - To Edit the entries in the table, click the **Edit** link next to the registry to make changes. Click **Update** to accept changes or click **Cancel Edit Mode** to remove changes.
 - To Delete the entries in the table, click the **Delete** link next to the registry.

When final selections have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen. Selections will not be saved.
- Click **Next** to move on to the next screen. Selections will be saved.
- Click **Save** to save selections and stay on the current screen.
- Click **Cancel** to remove selections and stay on the current screen.

8.2.8.5 Measure 5: Clinical Data Registry Reporting

MEASURE: The EP is in active engagement to submit data to a clinical data registry.

Public Health Objective Measures (Year 2 Attestation)

Clinical Data Registry Reporting

Objective

The EP is in active engagement with a public health agency (PHA) or clinical data registry (CDR) to submit electronic public health data in a meaningful way using CEHRT, except where prohibited, and in accordance with applicable law and practice.

Measure

The EP is in active engagement to submit data to a CDR.

*Would you like to attest to this measure?

Yes No

EXCLUSION 1: Does not diagnose or directly treat any disease or condition associated with a CDR in their jurisdiction during the EHR reporting period.

*Does this exclusion apply to you?

Yes No

EXCLUSION 2: The EP practices in a jurisdiction for which no CDR is capable of accepting electronic registry transactions in the specific standards required to meet the CEHRT definition at the start of the EHR reporting period.

*Does this exclusion apply to you?

Yes No

EXCLUSION 3: The EP practices in a jurisdiction where no CDR for which the EP is eligible to submit data has declared readiness to receive electronic registry transactions as of six months prior to the start of the EHR reporting period.

*Does this exclusion apply to you?

Yes No

Active Engagement Options:

Active Engagement Option 1 - Completed Registration to Submit Data:

The EP 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 EHR reporting period; and

the EP is awaiting an invitation from the PHA or CDR to begin testing and validation. This option allows EPs to meet the measure when the PHA or the CDR has limited resources to initiate the testing and validation process. EPs that have registered in previous years do not need to submit an additional registration to meet this requirement for each EHR reporting period.

Active Engagement Option 2 - Testing and Validation:

The EP is in the process of testing and validation of the electronic submission of data. EPs must respond to requests from the PHA or, where applicable, the CDR within 30 days; failure to respond twice within an EHR reporting period would result in that EP not meeting the measure.

Active Engagement Option 3 - Production:

The EP has completed testing and validation of the electronic submission and is electronically submitting production data to the PHA or CDR.

*** Please select the applicable active engagement option (may only select one).**

- Option1
- Option2
- Option3

Instructions:

Provider may report to more than one Clinical Data Registry and may count Clinical Data Registry reporting more than one time to meet the required number of measures for the objective. You may enter as many registries as you wish but only two will be counted towards the objective.

To report the first Clinical Data Registry, enter the information in the text box, then click 'Add'. To report the additional Clinical Data Registries, select the active engagement option applicable for the next registry you are reporting, enter the information in the text box and click 'Add'. Clinical Data Registry information you are attesting to will be displayed in the Registry table below.

*** Please add the clinical data registry below:**

- Other

LIST OF SPECIALIZED REGISTRIES YOU ADDED:

Type of Registry	Active Engagement Option	Description	Edit	Delete
Other	1	Radiology	Edit	Delete
Other	3	PATHology	Edit	Delete
Other	2	ECG	Edit	Delete
Other	1	X-ray	Edit	Delete

Add

Previous

Next

Save

Cancel

To satisfy the Measure, respond to the question.

- If Yes is selected, respond to Exclusion 1.
- If No is selected, respond to Exclusion 2.
- If No is selected, respond to Exclusion 3.
 - If No is selected, make two selections.
 - Select the applicable Active Engagement Option for each registry.
 - Add each clinical data registry to the table.
 - If Other is selected, type the name of the registry into the text box. Click **Add** to add it to the table.
 - To Edit the entries in the table, click the **Edit** link next to the registry to make changes. Click **Update** to accept changes or click **Cancel Edit Mode** to remove changes.
 - To Delete the entries in the table, click the **Delete** link next to the registry.

When final selections have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen. Selections will not be saved.
- Click **Next** to move on to the next screen. Selections will be saved.
- Click **Save** to save selections and stay on the current screen.
- Click **Cancel** to remove selections and stay on the current screen.

9 Electronic Clinical Quality Measures

Electronic clinical quality measures (eCQMs) are tools that help measure and track the quality of health care services that EPs, EHs, and CAHs provide, as generated by a provider's EHR. Measuring and reporting eCQMs helps to ensure that our health care system is delivering effective, safe, efficient, patient-centered, equitable, and timely care.

The 2021 eCQM reporting period for EPs is any continuous **90-day period within the 2021 CY**. All participating EPs are required to report on any **6 eCQMs** relevant to their scope of practice from the set of 47 available. In addition, EPs must report on at least one outcome measure. If no outcome measure is relevant to his or her scope of practice, the EP must report on one high priority measure. If no high priority measures are relevant to their scope of practice, they may report on any six relevant measures.

9.1 Electronic Clinical Quality Measure Submission Selection Screen

Electronic Clinical Quality Measure (CQM) Submission Selection Screen (Year 2 Attestation)

Reporting Electronic Clinical Quality Measures

In order to report eCQMs, you will need to select the method of submission below. EPs must report on 6 of the 47 approved eCQMs related to their scope of practice. Medicaid EP's are required to report on at least one outcome or high priority measure. If there are no outcome or high priority measures relevant to an EP's scope of practice, they may report on any 6 relevant measures. For additional information on eCQM reporting, please [click here](#). In order to report eCQMs, you will need to select the method of submission below. Selecting electronically will satisfy uploading the QRDA file through this application or sending it through KHIE.

How would you like to submit eCQMs?

Manually Electronically

Providers who are attesting to Meaningful Use may use any continuous 90 days within the calendar year. Please provide the eCQM reporting period associated with this attestation:

*eCQM Reporting Period Start Date: (mm/dd/yy)

*eCQM Reporting Period End Date: (mm/dd/yy)

Previous
Next
Save
Cancel

Enter responses for the following:

- How would you like to submit eCQMs?
 - Select Manually or Electronically; selecting electronically will satisfy uploading the QRDA file through the KYSLR or sending it through KHIE.
- Enter the eCQM Reporting Period Start Date
 - This is the start date of the reporting period for selected eCQMs.
- Enter the eCQM Reporting Period End Date
 - This is the end date of the reporting period for selected eCQMs.

When final selections have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen. Selections will not be saved.
- Click **Next** to move on to the next screen. Selections will be saved.
- Click **Save** to save selections and stay on the current screen.
- Click **Cancel** to remove selections and stay on the current screen.

9.2 Electronic Clinical Quality Measures Electronically Reported Selection Screen

Electronic Clinical Quality Measures (Year 3 Attestation)

Electronic Clinical Quality Measures Details Report

QRDA File Upload

Please select the applicable button below to review and evaluate your eCQMs QRDA III files. QRDA III files are required for verification and must meet CMS defined thresholds in order to successfully attest. The most recent submission with an accepted status will be used for validation. If two files must be used for validation, then the two most recent submissions with an accepted status will be used. QRDA I files will be accepted however submission will not fulfil the requirements of electronic submission.

Upload QRDA files - This button will be used to upload the QRDA file(s) from your local computer into the attestation.

Upload KHIE QRDA files - This button will be used to upload files retrieved from KHIE (if available). You must select the file and then click on Upload KHIE QRDA button.

Upload QRDA files

KHIE QRDA Files

KHIE QRDA FILES

No Data Found!

Uploaded Invalid Files

INVALID FILES DETAILS

No errors to report.

To view the emeasures from your QRDA III file, click the select link in the corresponding row. Before proceeding, please review the emeasures and details.

The most recent submission with an accepted status will be used for validation. If two files must be used for validation, then the two most recent submissions with an accepted status will be used for validation.

Rows that do not include a select link are QRDA I documents that have been uploaded.

UPLOADED FILES				
	FileTransmissionID	Status	DateReceived	FileName
Select	587	Pending	8/17/2018 3:23:00 PM	QRDA_XML2_1013165786_2MSRS_new.xml
Select	586	Pending	8/17/2018 3:00:32 PM	QRDA_XML2_1013165786_4MSRS_1.xml
Select	585	Accepted	8/17/2018 2:46:26 PM	QRDA_XML2_1013165786__4Msrs.xml
Select	584	Accepted	8/17/2018 12:31:16 PM	MIPS_Sample_QRDA_III_2MSRS_2.xml
Select	583	Pending	8/17/2018 12:22:55 PM	MIPS_Sample_QRDA_III_2MSRS_LAst.xml
Select	582	Rejected	8/17/2018 12:22:22 PM	MIPS_Sample_QRDA_III_1MSRS.xml
Select	580	Rejected	8/17/2018 12:21:57 PM	MIPS_Sample_QRDA_III_2MSRS.xml

Selected File Electronic Clinical Quality Measures

ELECTRONIC CLINICAL QUALITY MEASURE DETAILS					
Measure Details				Domain	
eMeasure Title	Version Neutral ID	eMeasure Version Number	NQF Measure Number	Version Specific ID	Patient Safety
Use of High-Risk Medications in the Elderly	a3837ff8-1abc-4ba9-800e-fd4e7953adbd	2		40280381-3D61-56A7-013E-65C9C3043E54	
Member of Measure Set: NONE - eMeasure ID:5b3d9245-fb54-499c-b5c5-21c20564f0bd Initial Patient Population: 73 SexFemale: 48Male: 23Undifferentiated: 2 EthnicityHispanic or Latino: 7Not Hispanic or Latino: 5 PayerMEDICARE: 2BLUE CROSS/BLUE SHIELD: 1Unavailable / Unknown: 70 RaceAsian: 5Black or African American: 3White: 4Other Race: 2 Denominator: 73 SexFemale: 48Male: 23Undifferentiated: 2 EthnicityHispanic or Latino: 7Not Hispanic or Latino: 5 PayerMEDICARE: 2BLUE CROSS/BLUE SHIELD: 1Unavailable / Unknown: 70 RaceAsian: 5Black or African American: 3White: 4Other Race: 2 Numerator 1: 22 SexFemale: 14Male: 8 PayerUnavailable / Unknown: 22 Numerator 2: 10 SexFemale: 5Male: 5 PayerUnavailable / Unknown: 10					
eMeasure Title	Version Neutral ID	eMeasure Version Number	NQF Measure Number	Version Specific ID	Effective Clinical Care
CERVICAL CANCER SCREENING	42e7e489-790f-427a-a1a6-d6e807f65a6d	2		40280381-3D61-56A7-013E-669CBC034836	
Member of Measure Set: NONE - eMeasure ID:e1d695b0-acee-472f-bfaf-ddb6e1515933 Initial Patient Population: 107 SexFemale: 107 EthnicityNot Hispanic or Latino: 3 PayerMEDICARE: 20BLUE CROSS/BLUE SHIELD: 1Unavailable / Unknown: 86 RaceAmerican Indian or Alaska Native: 1Black or African American: 2 Denominator: 107 SexFemale: 107 EthnicityNot Hispanic or Latino: 3 PayerMEDICARE: 20BLUE CROSS/BLUE SHIELD: 1Unavailable / Unknown: 86 RaceAmerican Indian or Alaska Native: 1Black or African American: 2Numerator: 0Denominator Exclusions: 0					

eMeasure Title	Version Neutral ID	eMeasure Version Number	NQF Measure Number	Version Specific ID	Community / Population Health
Preventive Care and Screening: Influenza Immunization	a244aa29-7d11-4616-888a-86e376bfcc6f	2		40280381-3D61-56A7-013E-57F49972361A	

Member of Measure Set: NONE - eMeasure ID:40d83c98-aaa9-4436-9c12-54b0896b0b2c
 Initial Patient Population: 158 SexFemale: 51Male: 104Undifferentiated: 3
 EthnicityHispanic or Latino: 6Not Hispanic or Latino: 4 PayerMEDICARE: 6Unavailable / Unknown: 152 RaceAmerican Indian or Alaska Native: 3Asian: 4White: 1Other Race: 2
 Denominator: 83 SexFemale: 17Male: 66 EthnicityHispanic or Latino: 1Not Hispanic or Latino: 1 PayerMEDICARE: 1Unavailable / Unknown: 82 RaceAmerican Indian or Alaska Native: 1Asian: 1 Numerator: 15 SexFemale: 3Male: 12 PayerUnavailable / Unknown: 15
 Denominator Exceptions: 24 SexFemale: 6Male: 18 PayerUnavailable / Unknown: 24

Previous
Continue

To submit eCQMs electronically, click the Browse button to select the QRDA file you wish to upload. Once the file is selected click the Upload QRDA File button. If you wish to upload a file retrieved from KHIE ,you must select the file and then click on the Upload KHIE QRDA button.

- To view the emeasures from your QRDA III file, click the select link in the corresponding row. The most recent submission with an accepted status will be used for validation. If two files must be used for validation, the two most recent submissions with an accepted status will be used.

When final selections have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen. Selections will not be saved.
- Click **Continue** to move to the next screen.

9.3 Electronic Clinical Quality Measures Electronically Reported Summary

Electronic Clinical Quality Measure (Year 4 Attestation)

Electronic Clinical Quality Measure Summary Report

<u>Date Received</u>	<u>Report NPI</u>	<u>Reporting Period Start</u>	<u>Reporting Period End</u>
8/17/2018 2:46:26 PM	0505050505	1/1/2018 12:00:00 AM	12/31/2018 12:00:00 AM
<u>Document Type</u>	<u>eCQM Status:</u>	<u>Evaluated Date:</u>	<u>File Name:</u>
QRDA III	Accepted	8/17/2018 3:24:52 PM	QRDA_XML2_1013165786.xml, MIPS_Sample_QRDA_II_2.xml

QRDA_XML2_1013165786__4MSRS.XML

Domain	# of Measures
Communication and Care Coordination	0
Community/Population Health	2
Effective Clinical Care	1
Efficiency and Cost Reduction	0
Patient Safety	1
Person and Caregiver-Centered Experience and Outcomes	0

MIPS_SAMPLE_QRDA_III_2MSRS_2.XML

Domain	# of Measures
Communication and Care Coordination	0
Community/Population Health	1
Effective Clinical Care	1
Efficiency and Cost Reduction	0
Patient Safety	0
Person and Caregiver-Centered Experience and Outcomes	0

Total Required	6
Total Met	6
Final eCQM Result:	Submission Accepted, click Next to continue.

Previous
Evaluate eCQM Submissions
Next

To evaluate eCQMs submitted electronically, click Evaluate eCQM Submission button.

When final selections have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen. Selections will not be saved.
- Click **Cancel** to remove selections and stay on the current screen.

9.4 Electronic Clinical Quality Measures Manually Reported Selection Screen

Electronic Clinical Quality Measures (eCQMs) Selection Screen (Year 3 Attestation)

Instructions:

Select a minimum of 6 eCQMs from the list below. You will be prompted to enter numerator(s), denominator(s), performance rate(s), and exclusion(s) or exception(s), if applicable, for all selected eCQMs after you select the Save & Next button below.

Deselect All

PERSON AND CAREGIVER-CENTERED EXPERIENCE AND OUTCOMES

Selection	Measure #	Title
<input checked="" type="checkbox"/>	CMS157v9.2/NQF 0384e	Oncology: Medical and Radiation - Pain Intensity Quantified
<input checked="" type="checkbox"/>	CMS56v9.2/NQF XXXX	Functional Status Assessment for Total Hip Replacement
<input checked="" type="checkbox"/>	CMS66v9.3/NQF XXXX	Functional Status Assessment for Total Knee Replacement
<input checked="" type="checkbox"/>	CMS90v10.2/NQF XXXX	Functional Status Assessments for Congestive Heart Failure
<input checked="" type="checkbox"/>	CMS771v2.2/NQF XXXX	Urinary Symptom Score Change 6-12 Months After Diagnosis of Benign Prostatic Hyperplasia

PATIENT SAFETY

Selection	Measure #	Title
<input checked="" type="checkbox"/>	CMS156v9.3/NQF XXXX	Use of High-Risk Medications in the Older Adults
<input checked="" type="checkbox"/>	CMS139v9.2/NQF XXXX	Falls: Screening for Future Fall Risk
<input checked="" type="checkbox"/>	CMS68v10.3/NQF 0419e	Documentation of Current Medications in the Medical Record
<input checked="" type="checkbox"/>	CMS177v9.2/NQF 1365e	Child and Adolescent Major Depressive Disorder (MDD): Suicide Risk Assessment

COMMUNICATION AND CARE COORDINATION

Selection	Measure #	Title
<input checked="" type="checkbox"/>	CMS142v9.2/NQF XXXX	Diabetic Retinopathy: Communication with the Physician Managing Ongoing Diabetes Care
<input checked="" type="checkbox"/>	CMS50v9.2/NQF XXXX	Closing the Referral Loop: Receipt of Specialist Report

COMMUNITY/POPULATION HEALTH

Selection	Measure #	Title
<input checked="" type="checkbox"/>	CMS155v9.2/NQF XXXX	Weight Assessment and Counseling for Nutrition and Physical Activity for Children and Adolescents
<input checked="" type="checkbox"/>	CMS138v9.2/NQF 0028e	Preventive Care and Screening: Tobacco Use: Screening and Cessation Intervention
<input checked="" type="checkbox"/>	CMS153v9.2/NQF XXXX	Chlamydia Screening for Women
<input checked="" type="checkbox"/>	CMS117v9.2/NQF XXXX	Childhood Immunization Status
<input checked="" type="checkbox"/>	CMS147v10.2/NQF 0041e	Preventive Care and Screening: Influenza Immunization
<input checked="" type="checkbox"/>	CMS127v9.2/NQF XXXX	Pneumococcal Vaccination Status for Older Adults
<input checked="" type="checkbox"/>	CMS2v10.2/NQF 0418e	Preventive Care and Screening: Screening for Depression and Follow-Up Plan
<input checked="" type="checkbox"/>	CMS69v9.3/NQF XXXX	Preventive Care and Screening: Body Mass Index (BMI) Screening and Follow-Up Plan
<input checked="" type="checkbox"/>	CMS22v9.3/NQF XXXX	Preventive Care and Screening: Screening for High Blood Pressure and Follow-Up Documented
<input checked="" type="checkbox"/>	CMS75v9.2/NQF XXXX	Children Who Have Dental Decay or Cavities
<input checked="" type="checkbox"/>	CMS349v3.3/NQF XXXX	HIV Screening

EFFICIENCY AND COST REDUCTION

Selection	Measure #	Title
<input checked="" type="checkbox"/>	CMS146v9.2/NQF XXXX	Appropriate Testing for Pharyngitis
<input checked="" type="checkbox"/>	CMS154v9.2/NQF XXXX	Appropriate Treatment for Upper Respiratory Infection (URI)
<input checked="" type="checkbox"/>	CMS129v10.3/NQF 0389e	Prostate Cancer: Avoidance of Overuse of Bone Scan for Staging Low Risk Prostate Cancer Patients
<input checked="" type="checkbox"/>	CMS249v3.2/NQF 3475e	Appropriate Use of DXA Scans in Women Under 65 Years Who Do Not Meet the Risk Factor Profile for Osteoporotic Fracture

EFFECTIVE CLINICAL CARE

Selection	Measure #	Title
<input checked="" type="checkbox"/>	CMS137v9.3/NQF XXXX	Initiation and Engagement of Alcohol and Other Drug Dependence Treatment
<input checked="" type="checkbox"/>	CMS165v9.2/NQF XXXX	Controlling High Blood Pressure
<input checked="" type="checkbox"/>	CMS124v9.1/NQF XXXX	Cervical Cancer Screening
<input checked="" type="checkbox"/>	CMS130v9.2/NQF XXXX	Colorectal Cancer Screening
<input checked="" type="checkbox"/>	CMS131v9.2/NQF XXXX	Diabetes: Eye Exam
<input checked="" type="checkbox"/>	CMS122v9.3/NQF XXXX	Diabetes: Hemoglobin A1c (HbA1c) Poor Control (> 9%)
<input checked="" type="checkbox"/>	CMS134v9.3/NQF XXXX	Diabetes: Medical Attention for Nephropathy
<input checked="" type="checkbox"/>	CMS145v9.2/NQF 0070e	Coronary Artery Disease (CAD): Beta-Blocker Therapy-Prior Myocardial Infarction (MI) or Left Ventricular Systolic Dysfunction (LVEF < 40%)
<input checked="" type="checkbox"/>	CMS135v9.2/NQF 0081e	Heart Failure (HF): ACE Inhibitor or ARB or ARNI Therapy for Left Ventricular Systolic Dysfunction (LVSD)
<input checked="" type="checkbox"/>	CMS144v9.2/NQF 0083e	Heart Failure (HF): Beta-Blocker Therapy for Left Ventricular Systolic Dysfunction (LVSD)

<input checked="" type="checkbox"/>	CMS143v9.2/NQF 0086e	Primary Open-Angle Glaucoma (POAG): Optic Nerve Evaluation
<input checked="" type="checkbox"/>	CMS161v9.2/NQF 0104e	Adult Major Depressive Disorder (MDD): Suicide Risk Assessment
<input checked="" type="checkbox"/>	CMS128v9.2/NQF XXXX	Anti-depressant Medication Management
<input checked="" type="checkbox"/>	CMS136v10.2/NQF XXXX	Follow-Up Care for Children Prescribed ADHD Medication (ADD)
<input checked="" type="checkbox"/>	CMS133v9.2/NQF 0565e	Cataracts: 20/40 or Better Visual Acuity within 90 Days Following Cataract Surgery
<input checked="" type="checkbox"/>	CMS159v9.4/NQF 0710e	Depression Remission at Twelve Months
<input checked="" type="checkbox"/>	CMS125v9.2/NQF XXXX	Breast Cancer Screening
<input checked="" type="checkbox"/>	CMS149v9.2/NQF 2872e	Dementia: Cognitive Assessment
<input checked="" type="checkbox"/>	CMS74v10.2/NQF XXXX	Primary Caries Prevention Intervention as Offered by Primary Care Providers, including Dentists
<input checked="" type="checkbox"/>	CMS347v4.3/NQF XXXX	Statin Therapy for the Prevention and Treatment of Cardiovascular Disease
<input checked="" type="checkbox"/>	CMS645v4.1/NQF XXXX	Bone density evaluation for patients with prostate cancer and receiving androgen deprivation therapy

Previous
Save & Next

Select at least six of the eQMs out of the 47 eQMs available.

When final selections have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen. Selections will not be saved.
- Click **Save & Next** to save selections and to move on to the next screen.

9.5 Electronic Clinical Quality Measures Manually Reported

9.5.1 CMS146

Electronic Clinical Quality Measures (Year 3 Attestation)

Questionnaire 1 of 47

(* Red asterisk indicates a required field.)

Measure:	CMS146/NQF XXXX
Versions:	CMS146v9.2
Title:	Appropriate Testing for Pharyngitis
Description:	The percentage of episodes for patients 3 years and older with a diagnosis of pharyngitis that resulted in an antibiotic dispensing event and a group A streptococcus (strep) test.
Denominator:	Outpatient, telephone, online assessment, observation, or emergency department (ED) visits with a diagnosis of pharyngitis and an antibiotic dispensing event among patients 3 years or older.
Numerator:	A group A streptococcus test in the seven-day period from three days prior to the episode date through three days after the episode date.
Denominator Exclusions:	<p>Exclude episodes where the patients is taking antibiotics in the 30 days prior to the episode date.</p> <p>Exclude episodes where the patient had a competing comorbid condition during the 12 months prior to or on the episode date.</p> <p>Exclude episodes when the patient had hospice care overlapping with the measurement period.</p> <p>Exclude episodes where the patient had a competing diagnosis within three days after the episode date.</p>

Complete the following information:

Stratum 1: 3-17 years

* Denominator 1:	* Numerator 1:	* Performance Rate 1 (%):	* Exclusion 1:
4	3	0.00	0

Stratum 2: 18-64 years

* Denominator 1:	* Numerator 1:	* Performance Rate 1 (%):	* Exclusion 1:
1	1	1.00	1

Stratum 3: 65 years and older

* Denominator 1:	* Numerator 1:	* Performance Rate 1 (%):	* Exclusion 1:
3	1	6.00	6

Previous

Next

Save

Cancel

To satisfy this eCQM, enter a whole number into the Denominator, Numerator, Performance Rate, and Exclusion boxes.

When final selections have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen. Selections will not be saved.
- Click **Next** to move on to the next screen. Selections will be saved.
- Click **Save** to save selections and stay on the current screen.
- Click **Cancel** to remove selections and stay on the current screen.

9.5.2 CMS137

Electronic Clinical Quality Measures (Year 3 Attestation)

Questionnaire 2 of 47

(* Red asterisk indicates a required field.)

Measure:	CMS137/NQF XXXX
Versions:	CMS137v9.3
Title:	Initiation and Engagement of Alcohol and Other Drug Dependence Treatment
Description:	<p>Percentage of patients 13 years of age and older with a new episode of alcohol or other drug abuse or (AOD) dependence who received the following. Two rates are reported.</p> <p>a. Percentage of patients who initiated treatment including either an intervention or medication for the treatment of AOD abuse or dependence within 14 days of the diagnosis.</p> <p>b. Percentage of patients who engaged in ongoing treatment including two additional interventions or a medication for the treatment of AOD abuse or dependence within 34 days of the initiation visit. For patients who initiated treatment with a medication, at least one of the two engagement events must be a treatment intervention.</p>
Denominator:	Patients age 13 years of age and older who were diagnosed with a new episode of alcohol, opioid, or other drug abuse or dependency during a visit between January 1 and November 14 of the measurement period.
Numerator:	<p>Numerator 1: Initiation of treatment includes either an intervention or medication for the treatment of AOD abuse or dependence within 14 days of the diagnosis.</p> <p>Numerator 2: Engagement in ongoing treatment includes two additional interventions or a medication for the treatment of AOD abuse or dependence within 34 days of the initiation visit. For patients who initiated treatment with a medication, at least one of the two engagement events must be a treatment intervention (i.e., engagement for these members cannot be satisfied with medication treatment alone).</p>
Denominator Exclusions:	<p>Exclude patients with a previous active diagnosis of alcohol, opioid or other drug abuse or dependence in the 60 days prior to the first episode of alcohol or drug dependence.</p> <p>Exclude patients whose hospice care overlaps the measurement period.</p>
Complete the following information:	

Stratum 1: Patients age 13 - 17 at the end of the Measurement Period

* Denominator 1:	* Numerator 1:	* Performance Rate 1 (%):	* Exclusion 1:
0	0	0.00	0
* Denominator 2:	* Numerator 2:	* Performance Rate 2 (%):	* Exclusion 2:
7	7	7.00	8

Stratum 2: Patients age >=18 at the end of Measurement Period

* Denominator 1:	* Numerator 1:	* Performance Rate 1 (%):	* Exclusion 1:
9	9	9.00	9
* Denominator 2:	* Numerator 2:	* Performance Rate 2 (%):	* Exclusion 2:
9	9	9.00	9

Stratum 3: Total Score

* Denominator 1:	* Numerator 1:	* Performance Rate 1 (%):	* Exclusion 1:
5	5	5.00	5
* Denominator 2:	* Numerator 2:	* Performance Rate 2 (%):	* Exclusion 2:
5	5	5.00	5

Previous
Next
Save
Cancel

To satisfy this eCQM, enter a whole number into each of the Denominator, Numerator, Performance Rate, and Exclusion boxes.

When final selections have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen. Selections will not be saved.
- Click **Next** to move on to the next screen. Selections will be saved.
- Click **Save** to save selections and stay on the current screen.
- Click **Cancel** to remove selections and stay on the current screen.

9.5.3 CMS165

Electronic Clinical Quality Measures (Year 3 Attestation)

Questionnaire 3 of 47

(*) Red asterisk indicates a required field.

Measure:	CMS165/NQF XXXX
Versions:	CMS165v9.2
Title:	Controlling High Blood Pressure
Description:	Percentage of patients 18-85 years of age who had a diagnosis of hypertension overlapping the measurement period or the year prior to the measurement period, and whose most recent blood pressure was adequately controlled (< 140/90mmHg) during the measurement period.
Denominator:	Patients 18-85 years of age who had a visit and diagnosis of essential hypertension overlapping the measurement period or the year prior to the measurement period.
Numerator:	Patients whose most recent blood pressure is adequately controlled (systolic blood pressure < 140 mmHg and diastolic blood pressure < 90 mmHg) during the measurement period.

Denominator Exclusions: Patients with evidence of end stage renal disease (ESRD), dialysis or renal transplant before or during the measurement period. Also exclude patients with a diagnosis of pregnancy during the measurement period.
 Exclude patients whose hospice care overlaps the measurement period.
 Exclude patients 66 and older who are living long term in an institution for more than 90 consecutive days during the measurement period.
 Exclude patients 66 and older with advanced illness and frailty because it is unlikely that patients will benefit from the services being measured.

Complete the following information:

* Denominator 1:	* Numerator 1:	* Performance Rate 1 (%):	* Exclusion 1:
1	1	1.00	1

Previous
Next
Save
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To satisfy this eCQM, enter a whole number into the Denominator, Numerator, Performance Rate, and Exclusion boxes.

When final selections have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen. Selections will not be saved.
- Click **Next** to move on to the next screen. Selections will be saved.
- Click **Save** to save selections and stay on the current screen.
- Click **Cancel** to remove selections and stay on the current screen.

9.5.4 CMS156

Electronic Clinical Quality Measures (Year 3 Attestation)

Questionnaire 4 of 47

(*) Red asterisk indicates a required field.

Measure:	CMS156/NQF XXXX		
Versions:	CMS156v9.3		
Title:	Use of High-Risk Medications in the Older Adults		
Description:	Percentage of patients 65 years of age and older who were ordered at least two of the same high-risk medications.		
Denominator:	Patients 65 years and older who had a visit during the measurement period.		
Numerator:	Patients with at least two orders for the same high-risk medication on different days during the measurement period.		
Denominator Exclusions:	Exclude patients whose hospice care overlaps the measurement period.		
Complete the following information:			
* Denominator 1:	* Numerator 1:	* Performance Rate 1 (%):	* Exclusion 1:
<input type="text" value="4"/>	<input type="text" value="3"/>	<input type="text" value="3.00"/>	<input type="text" value="4"/>

Previous

Next

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Cancel

To satisfy this eQCM, enter a whole number into each of the Denominator, Numerator, Performance Rate, and Exclusion boxes.

When final selections have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen. Selections will not be saved.
- Click **Next** to move on to the next screen. Selections will be saved.
- Click **Save** to save selections and stay on the current screen.
- Click **Cancel** to remove selections and stay on the current screen.

9.5.5 CMS155

Electronic Clinical Quality Measures (Year 3 Attestation)

Questionnaire 5 of 47

(*) Red asterisk indicates a required field.

Measure:	CMS155/NQF XXXX
Versions:	CMS155v9.2
Title:	Weight Assessment and Counseling for Nutrition and Physical Activity for Children and Adolescents
Description:	<p>Percentage of patients 3-17 years of age who had an outpatient visit with a Primary Care Physician (PCP) or Obstetrician/Gynecologist (OB/GYN) and who had evidence of the following during the measurement period. Three rates are reported.</p> <ul style="list-style-type: none"> - Percentage of patients with height, weight, and body mass index (BMI) percentile documentation. - Percentage of patients with counseling for nutrition. - Percentage of patients with counseling for physical activity.

Denominator: Patients 3-17 years of age with at least one outpatient visit with a primary care physician (PCP) or an obstetrician/gynecologist (OB/GYN) during the measurement period.

Numerator:

Numerator 1: Patients who had a height, weight and body mass index (BMI) percentile recorded during the measurement period.

Numerator 2: Patients who had counseling for nutrition during the measurement period.

Numerator 3: Patients who had counseling for physical activity during the measurement period.

Denominator Exclusions: Patients who have a diagnosis of pregnancy during the measurement period. Exclude patients whose hospice care overlaps the measurement period.

Complete the following information:

Stratum 1: Patients age 3-11			
* Denominator 1:	* Numerator 1:	* Performance Rate 1 (%):	* Exclusion 1:
2	2	2.00	2
* Denominator 2:	* Numerator 2:	* Performance Rate 2 (%):	* Exclusion 2:
2	2	2.00	2
* Denominator 3:	* Numerator 3:	* Performance Rate 3 (%):	* Exclusion 3:
2	2	2.00	2
Stratum 2: Patients age 12-17			
* Denominator 1:	* Numerator 1:	* Performance Rate 1 (%):	* Exclusion 1:
1	1	1.00	1
* Denominator 2:	* Numerator 2:	* Performance Rate 2 (%):	* Exclusion 2:
1	1	1.00	1
* Denominator 3:	* Numerator 3:	* Performance Rate 3 (%):	* Exclusion 3:
1	1	1.00	1
Stratum 3: Total Score			
* Denominator 1:	* Numerator 1:	* Performance Rate 1 (%):	* Exclusion 1:
3	3	3.00	3
* Denominator 2:	* Numerator 2:	* Performance Rate 2 (%):	* Exclusion 2:
3	3	3.00	3
* Denominator 3:	* Numerator 3:	* Performance Rate 3 (%):	* Exclusion 3:
3	3	3.00	0

Previous
Next
Save
Cancel

To satisfy this eCQM, enter a whole number into each of the Denominator, Numerator, Performance Rate, and Exclusion boxes.

When final selections have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen. Selections will not be saved.
- Click **Next** to move on to the next screen. Selections will be saved.
- Click **Save** to save selections and stay on the current screen.
- Click **Cancel** to remove selections and stay on the current screen.

9.5.6 CMS138

Electronic Clinical Quality Measures (Year 3 Attestation)

Questionnaire 6 of 47

(* Red asterisk indicates a required field.)

Measure:	CMS138/NQF 0028e
Versions:	CMS138v9.2
Title:	Preventive Care and Screening: Tobacco Use: Screening and Cessation Intervention
Description:	<p>Percentage of patients aged 18 years and older who were screened for tobacco use one or more times within 12 months AND who received tobacco cessation intervention if identified as a tobacco user. Three rates are reported.</p> <p>a. Percentage of patients aged 18 years and older who were screened for tobacco use one or more times within 12 months.</p> <p>b. Percentage of patients aged 18 years and older who were identified as a tobacco user who received tobacco cessation intervention.</p> <p>c. Percentage of patients aged 18 years and older who were screened for tobacco use one or more times within 12 months AND who received tobacco cessation intervention if identified as a tobacco user.</p>
Denominator:	<p>Denominator 1: Population 1: All patients aged 18 years and older seen for at least two visits or at least one preventive visit during the measurement period.</p> <p>Denominator 2: Population 2: All patients aged 18 years and older seen for at least two visits or at least one preventive visit during the measurement period who were screened for tobacco use and identified as a tobacco user.</p> <p>Denominator 3: Population 3: All patients aged 18 years and older seen for at least two visits or at least one preventive visit during the measurement period.</p>
Numerator:	<p>Numerator 1: Population 1: Patients who were screened for tobacco use at least once within 12 months.</p> <p>Numerator 2: Population 2: Patients who received tobacco cessation intervention.</p> <p>Numerator 3: Population 3: Patients who were screened for tobacco use at least once within 12 months AND who received tobacco cessation intervention if identified as a tobacco user.</p>
Denominator Exceptions:	<p>Exception 1: Population 1: Documentation of medical reason(s) for not screening for tobacco use (e.g., limited life expectancy, other medical reason).</p> <p>Exception 2: Population 2: Documentation of medical reason(s) for not providing tobacco cessation intervention (e.g., limited life expectancy, other medical reason).</p> <p>Exception 3: Population 3: Documentation of medical reason(s) for not screening for tobacco use OR for not providing tobacco cessation intervention for patients identified as tobacco users (e.g., limited life expectancy, other medical reason).</p>
Complete the following information:	

Population 1:			
* Denominator 1:	* Numerator 1:	* Performance Rate 1 (%):	* Exception 1:
0	0	0.00	0
Population 2:			
* Denominator 2:	* Numerator 2:	* Performance Rate 2 (%):	* Exception 2:
0	0	0.00	0
Population 3:			
* Denominator 3:	* Numerator 3:	* Performance Rate 3 (%):	* Exception 3:
0	0	0.00	0

Previous
Next
Save
Cancel

To satisfy this eCQM, enter a whole number into the Denominator, Numerator, Performance Rate, and Exception boxes.

When final selections have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen. Selections will not be saved.
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- Click **Save** to save selections and stay on the current screen.
- Click **Cancel** to remove selections and stay on the current screen.

9.5.7 CMS124

Electronic Clinical Quality Measures (Year 3 Attestation)

Questionnaire 7 of 47

(*) Red asterisk indicates a required field.

Measure:	CMS124/NQF XXXX
Versions:	CMS124v9.1
Title:	Cervical Cancer Screening
Description:	<p>Percentage of women 21-64 years of age who were screened for cervical cancer using either of the following criteria:</p> <ul style="list-style-type: none"> * Women age 21-64 who had cervical cytology performed within the last 3 years. * Women age 30-64 who had cervical human papillomavirus (HPV) testing performed within the last 5 years.

Denominator: Women 23-64 years of age with a visit during the measurement period.

Numerator: Women with one or more screenings for cervical cancer. Appropriate screenings are defined by any one of the following criteria:
 * Cervical cytology performed during the measurement period or the two years prior to the measurement period for women who are at least 21 years old at the time of the test.
 * Cervical human papillomavirus (HPV) testing performed during the measurement period or the four years prior to the measurement period for women who are 30 years or older at the time of the test.

Denominator Exclusions: Women who had a hysterectomy with no residual cervix or a congenital absence of cervix.
 Exclude patients whose hospice care overlaps the measurement period.

Complete the following information:

* Denominator 1:	* Numerator 1:	* Performance Rate 1 (%):	* Exclusion 1:
32	12	23.00	32



To satisfy this eCQM, enter a whole number into the Denominator, Numerator, Performance Rate, and Exclusion boxes.

When final selections have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen. Selections will not be saved.
- Click **Next** to move on to the next screen. Selections will be saved.
- Click **Save** to save selections and stay on the current screen.
- Click **Cancel** to remove selections and stay on the current screen.

9.5.8 CMS153

Electronic Clinical Quality Measures (Year 3 Attestation)

Questionnaire 8 of 47

(*) Red asterisk indicates a required field.

Measure: CMS153/NQF XXXX

Versions: CMS153v9.2

Title: Chlamydia Screening for Women

Description: Percentage of women 16-24 years of age who were identified as sexually active and who had at least one test for chlamydia during the measurement period.

Denominator: Women 16 to 24 years of age who are sexually active and who had a visit in the measurement period.

Numerator: Women with at least one chlamydia test during the measurement period.

Denominator Exclusions: Women who are only eligible for the initial population due to a pregnancy test and who had an x-ray or an order for a specified medication within 7 days of the pregnancy test.
Exclude patients whose hospice care overlaps the measurement period.

Complete the following information:

Stratum 1: Patients age 16-20

* Denominator 1:	* Numerator 1:	* Performance Rate 1 (%):	* Exclusion 1:
<input type="text" value="1"/>	<input type="text" value="1"/>	<input type="text" value="1.00"/>	<input type="text" value="1"/>

Stratum 2: Patients age 21-24

* Denominator 1:	* Numerator 1:	* Performance Rate 1 (%):	* Exclusion 1:
<input type="text" value="1"/>	<input type="text" value="1"/>	<input type="text" value="11.00"/>	<input type="text" value="11"/>

Stratum 3: Total Score

* Denominator 1:	* Numerator 1:	* Performance Rate 1 (%):	* Exclusion 1:
<input type="text" value="3"/>	<input type="text" value="3"/>	<input type="text" value="33.00"/>	<input type="text" value="33"/>

To satisfy this eCQM, enter a whole number into the Denominator, Numerator, Performance Rate, and Exclusion boxes.

When final selections have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen. Selections will not be saved.
- Click **Next** to move on to the next screen. Selections will be saved.
- Click **Save** to save selections and stay on the current screen.
- Click **Cancel** to remove selections and stay on the current screen.

9.5.9 CMS130

Electronic Clinical Quality Measures (Year 3 Attestation)

Questionnaire 2 of 25

(*) Red asterisk indicates a required field.

Measure:	CMS130/NQF XXXX
Versions:	CMS130v9.2
Title:	Colorectal Cancer Screening
Description:	Percentage of adults 50-75 years of age who had appropriate screening for colorectal cancer.
Denominator:	Patients 50-75 years of age with a visit during the measurement period.
Numerator:	<p>Patients with one or more screenings for colorectal cancer. Appropriate screenings are defined by any one of the following criteria:</p> <ul style="list-style-type: none"> - Fecal occult blood test (FOBT) during the measurement period. - Flexible sigmoidoscopy during the measurement period or the four years prior to the measurement period. - Colonoscopy during the measurement period or the nine years prior to the measurement period. - FIT-DNA during the measurement period or the two years prior to the measurement period. - CT Colonography during the measurement period or the four years prior to the measurement period.
Denominator Exclusions:	<p>Exclude patients whose hospice care overlaps the measurement period.</p> <p>Exclude patients with a diagnosis or past history of total colectomy or colorectal cancer.</p> <p>Exclude patients 66 and older who are living long term in an institution for more than 90 consecutive days during the measurement period.</p> <p>Exclude patients 66 and older with advanced illness and frailty because it is unlikely that patients will benefit from the services being measured.</p>

Complete the following information:

* Denominator 1:	* Numerator 1:	* Performance Rate 1 (%):	* Exclusion 1:
<input type="text" value="436"/>	<input type="text" value="46"/>	<input type="text" value="346.00"/>	<input type="text" value="346"/>

To satisfy this eCQM, enter a whole number into each of the Denominator, Numerator, Performance Rate, and Exclusion boxes.

When final selections have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen. Selections will not be saved.
- Click **Next** to move on to the next screen. Selections will be saved.
- Click **Save** to save selections and stay on the current screen.
- Click **Cancel** to remove selections and stay on the current screen.

9.5.10 CMS117

Electronic Clinical Quality Measures (Year 3 Attestation)

Questionnaire 3 of 25

(*) Red asterisk indicates a required field.

Measure:	CMS117/NQF XXXX
Versions:	CMS117v9.2
Title:	Childhood Immunization Status
Description:	Percentage of children 2 years of age who had four diphtheria, tetanus and acellular pertussis (DTaP); three polio (IPV), one measles, mumps and rubella (MMR); three or four H influenza type B (Hib); three hepatitis B (Hep B); one chicken pox (VZV); four pneumococcal conjugate (PCV); one hepatitis A (Hep A); two or three rotavirus (RV); and two influenza (flu) vaccines by their second birthday.
Denominator:	Children who turn 2 years of age during the measurement period and who have a visit during the measurement period.
Numerator:	Children who have evidence showing they received recommended vaccines, had documented history of the illness, had a seropositive test result, or had an allergic reaction to the vaccine by their second birthday.

Denominator Exclusions: Exclude patients whose hospice care overlaps the measurement period.

Complete the following information:

* Denominator 1:	* Numerator 1:	* Performance Rate 1 (%):	* Exclusion 1:
346	4	46.00	43



To satisfy this eCQM, enter a whole number into the Denominator, Numerator, Performance Rate, and Exclusion boxes.

When final selections have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen. Selections will not be saved.
- Click **Next** to move on to the next screen. Selections will be saved.
- Click **Save** to save selections and stay on the current screen.
- Click **Cancel** to remove selections and stay on the current screen.

9.5.11 CMS147

Electronic Clinical Quality Measures (Year 3 Attestation)

Questionnaire 11 of 47

(*) Red asterisk indicates a required field.

Measure:	CMS147/NQF 0041e
Versions:	CMS147v10.2
Title:	Preventive Care and Screening: Influenza Immunization
Description:	Percentage of patients aged 6 months and older seen for a visit between October 1 and March 31 who received an influenza immunization OR who reported previous receipt of an influenza immunization.
Denominator:	All patients aged 6 months and older seen for a visit during the measurement period and seen for a visit between October 1 and March 31.
Numerator:	Patients who received an influenza immunization OR who reported previous receipt of an influenza immunization.

Denominator Exceptions: Documentation of medical reason(s) for not receiving influenza immunization (e.g., patient allergy, other medical reasons).
 Documentation of patient reason(s) for not receiving influenza immunization (e.g., patient declined, other patient reasons).
 Documentation of system reason(s) for not receiving influenza immunization (e.g., vaccine not available, other system reasons).

Complete the following information:

* Denominator 1:	* Numerator 1:	* Performance Rate 1 (%):	* Exception 1:
4	2	3.00	3



To satisfy this eCQM, enter a whole number into each of the Denominator, Numerator, Performance Rate, and Exception boxes.

When final selections have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen. Selections will not be saved.
- Click **Next** to move on to the next screen. Selections will be saved.
- Click **Save** to save selections and stay on the current screen.
- Click **Cancel** to remove selections and stay on the current screen.

9.5.12 CMS127

Electronic Clinical Quality Measures (Year 3 Attestation)

Questionnaire 12 of 47

(*) Red asterisk indicates a required field.

Measure:	CMS127/NQF XXXX
Versions:	CMS127v9.2
Title:	Pneumococcal Vaccination Status for Older Adults
Description:	Percentage of patients 65 years of age and older who have ever received a pneumococcal vaccine.

Denominator: Patients 65 years of age and older with a visit during the measurement period.

Numerator: Patients who have ever received a pneumococcal vaccination before the end of the measurement period.

Denominator Exclusions: Exclude patients whose hospice care overlaps the measurement period.

Complete the following information:

* **Denominator 1:** * **Numerator 1:** * **Performance Rate 1 (%):** * **Exclusion 1:**

To satisfy this eCQM, enter a whole number into the Denominator, Numerator, Performance Rate, and Exclusion boxes.

When final selections have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen. Selections will not be saved.
- Click **Next** to move on to the next screen. Selections will be saved.
- Click **Save** to save selections and stay on the current screen.
- Click **Cancel** to remove selections and stay on the current screen.

9.5.13 CMS131

Electronic Clinical Quality Measures (Year 3 Attestation)

Questionnaire 13 of 47

(*) Red asterisk indicates a required field.

Measure:	CMS131/NQF XXXX
Versions:	CMS131v9.2
Title:	Diabetes: Eye Exam

Description:	Percentage of patients 18-75 years of age with diabetes and an active diagnosis of retinopathy overlapping the measurement period who had a retinal or dilated eye exam by an eye care professional during the measurement period or diabetics with no diagnosis of retinopathy overlapping the measurement period who had a retinal or dilated eye exam by an eye care professional during the measurement period or in the 12 months prior to the measurement period.		
Denominator:	Patients 18-75 years of age with diabetes with a visit during the measurement period		
Numerator:	Patients with an eye screening for diabetic retinal disease. This includes diabetics who had one of the following: *Diabetic with a diagnosis of retinopathy that overlaps the measurement period and a retinal or dilated eye exam by an eye care professional in the measurement period. *Diabetic with no diagnosis of retinopathy overlapping the measurement period and a retinal or dilated eye exam by an eye care professional in the measurement period or the year prior to the measurement period.		
Denominator Exclusions:	Exclude patients whose hospice care overlaps the measurement period. Exclude patients 66 and older who are living long term in an institution for more than 90 consecutive days during the measurement period. Exclude patients 66 and older with advanced illness and frailty because it is unlikely that patients will benefit from the services being measured.		
Complete the following information:			
* Denominator 1:	* Numerator 1:	* Performance Rate 1 (%):	* Exclusion 1:
<input type="text" value="2"/>	<input type="text" value="2"/>	<input type="text" value="5.00"/>	<input type="text" value="5"/>



To satisfy this eCQM, enter a whole number into the Denominator, Numerator, Performance Rate, and Exclusion boxes.

When final selections have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen. Selections will not be saved.
- Click **Next** to move on to the next screen. Selections will be saved.
- Click **Save** to save selections and stay on the current screen.
- Click **Cancel** to remove selections and stay on the current screen.

9.5.14 CMS122

Electronic Clinical Quality Measures (Year 3 Attestation)

Questionnaire 14 of 47

(* Red asterisk indicates a required field.)

Measure:	CMS122/NQF XXXX		
Versions:	CMS122v9.3		
Title:	Diabetes: Hemoglobin A1c (HbA1c) Poor Control (> 9%)		
Description:	Percentage of patients 18-75 years of age with diabetes who had hemoglobin A1c > 9.0% during the measurement period.		
Denominator:	Patients 18-75 years of age with diabetes with a visit during the measurement period.		
Numerator:	Patients whose most recent HbA1c level (performed during the measurement period) is > 9.0%.		
Denominator Exclusions:	Exclude patients whose hospice care overlaps the measurement period. Exclude patients 66 and older who are living long term in an institution for more than 90 consecutive days during the measurement period. Exclude patients 66 and older with advanced illness and frailty because it is unlikely that patients will benefit from the services being measured.		
Complete the following information:			
* Denominator 1:	* Numerator 1:	* Performance Rate 1 (%):	* Exclusion 1:
<input type="text" value="0"/>	<input type="text" value="0"/>	<input type="text" value="0.00"/>	<input type="text" value="0"/>

Previous
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To satisfy this eCQM, enter a whole number into the Denominator, Numerator, Performance Rate and Exclusion boxes.

When final selections have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen. Selections will not be saved.
- Click **Next** to move on to the next screen. Selections will be saved.
- Click **Save** to save selections and stay on the current screen.
- Click **Cancel** to remove selections and stay on the current screen.

9.5.15 CMS134

Electronic Clinical Quality Measures (Year 3 Attestation)

Questionnaire 15 of 47

(*) Red asterisk indicates a required field.

Measure:	CMS134/NQF XXXX
Versions:	CMS134v9.3
Title:	Diabetes: Medical Attention for Nephropathy
Description:	The percentage of patients 18-75 years of age with diabetes who had a nephropathy screening test or evidence of nephropathy during the measurement period.
Denominator:	Patients 18-75 years of age with diabetes with a visit during the measurement period.
Numerator:	Patients with a screening for nephropathy or evidence of nephropathy during the measurement period.
Denominator Exclusions:	Exclude patients whose hospice care overlaps the measurement period. Exclude patients 66 and older who are living long term in an institution for more than 90 consecutive days during the measurement period. Exclude patients 66 and older with advanced illness and frailty because it is unlikely that patients will benefit from the services being measured.

Complete the following information:

* Denominator 1:	* Numerator 1:	* Performance Rate 1 (%):	* Exclusion 1:
<input type="text" value="8"/>	<input type="text" value="8"/>	<input type="text" value="0.00"/>	<input type="text" value="0"/>

Previous
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To satisfy this eCQM, enter a whole number into the Denominator, Numerator, Performance Rate, and Exclusion boxes.

When final selections have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen. Selections will not be saved.
- Click **Next** to move on to the next screen. Selections will be saved.
- Click **Save** to save selections and stay on the current screen.

- Click **Cancel** to remove selections and stay on the current screen.

9.5.16 CMS154

Electronic Clinical Quality Measures (Year 3 Attestation)

Questionnaire 16 of 47

(*) Red asterisk indicates a required field.

Measure:	CMS154/NQF XXXX
Versions:	CMS154v9.2
Title:	Appropriate Treatment for Upper Respiratory Infection (URI)
Description:	Percentage of episodes for patients 3 months of age and older with a diagnosis of upper respiratory infection (URI) that did not result in an antibiotic dispensing event.

Denominator:	Outpatient visits, telephone visits, online assessments, observation stays or emergency department visits with a diagnosis of URI during the measurement period among patients 3 months of age and older.
Numerator:	URI episodes without a prescription for antibiotic medication on or 3 days after the outpatient visit, telephone visit, online assessment, observation stay or emergency department visit for an upper respiratory infection.
Denominator Exclusions:	<p>Exclude URI episodes when the patient had a competing comorbid condition during the 12 months prior to or on the episode date.</p> <p>Exclude URI episodes when the patient had a new or refill prescription of antibiotics in the 30 days prior to or on the episode date.</p> <p>Exclude URI episodes when the patient had competing diagnosis on or three days after the episode date.</p> <p>Exclude URI episodes when the patient had hospice care overlapping with the measurement period.</p>

Complete the following information:

Stratum 1: 3 months - 17 years

* Denominator 1:	* Numerator 1:	* Performance Rate 1 (%):	* Exclusion 1:
1	1	1.00	1

Stratum 2: 18 - 64 years			
* Denominator 1:	* Numerator 1:	* Performance Rate 1 (%):	* Exclusion 1:
2	2	2.00	2
Stratum 3 : 65 years and older			
* Denominator 1:	* Numerator 1:	* Performance Rate 1 (%):	* Exclusion 1:
3	3	3.00	3

Previous
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To satisfy this eCQM, enter a whole number into the Denominator, Numerator, Performance Rate, and Exclusion boxes.

When final selections have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen. Selections will not be saved.
- Click **Next** to move on to the next screen. Selections will be saved.
- Click **Save** to save selections and stay on the current screen.
- Click **Cancel** to remove selections and stay on the current screen.

9.5.17 CMS145

Electronic Clinical Quality Measures (Year 3 Attestation)

Questionnaire 17 of 47

(*) Red asterisk indicates a required field.

Measure:	CMS145/NQF 0070e
Versions:	CMS145v9.2
Title:	Coronary Artery Disease (CAD): Beta-Blocker Therapy-Prior Myocardial Infarction (MI) or Left Ventricular Systolic Dysfunction (LVEF < 40%)
Description:	Percentage of patients aged 18 years and older with a diagnosis of coronary artery disease seen within a 12-month period who also have a prior MI or a current or prior LVEF < 40% who were prescribed beta-blocker therapy.

Denominator: All patients aged 18 years and older with a diagnosis of coronary artery disease seen within a 12-month period who also have prior (within the past 3 years) MI or a current or prior LVEF < 40%.

Numerator: Patients who were prescribed beta-blocker therapy.

Denominator Exceptions: Documentation of medical reason(s) for not prescribing beta-blocker therapy (e.g., allergy, intolerance, other medical reasons).
Documentation of patient reason(s) for not prescribing beta-blocker therapy (e.g., patient declined, other patient reasons).
Documentation of system reason(s) for not prescribing beta-blocker therapy (e.g., other reasons attributable to the health care system).

Complete the following information:

Population Criteria 1: Patients with left ventricular systolic dysfunction (LVEF < 40%).

* Denominator 1:	* Numerator 1:	* Performance Rate 1 (%):	* Exception 1:
8	8	8.00	8

Population Criteria 2: Patients with a prior (within the past 3 years) myocardial infarction.

* Denominator 1:	* Numerator 1:	* Performance Rate 1 (%):	* Exception 1:
2	2	2.00	2

Previous

Next

Save

Cancel

To satisfy this eCQM, enter a whole number into the Denominator, Numerator, Performance Rate, and Exception boxes.

When final selections have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen. Selections will not be saved.
- Click **Next** to move on to the next screen. Selections will be saved.
- Click **Save** to save selections and stay on the current screen.
- Click **Cancel** to remove selections and stay on the current screen.

9.5.18 CMS135

Electronic Clinical Quality Measures (Year 3 Attestation)

Questionnaire 18 of 47

(*) Red asterisk indicates a required field.

Measure:	CMS135/NQF 0081e		
Versions:	CMS135v9.2		
Title:	Heart Failure (HF): ACE Inhibitor or ARB or ARNI Therapy for Left Ventricular Systolic Dysfunction (LVSD)		
Description:	Percentage of patients aged 18 years and older with a diagnosis of heart failure (HF) with a current or prior left ventricular ejection fraction (LVEF) < 40% who were prescribed ACE inhibitor or ARB or ARNI therapy either within a 12-month period when seen in the outpatient setting OR at each hospital discharge.		
Denominator:	All patients aged 18 years and older with a diagnosis of heart failure with a current or prior LVEF < 40%.		
Numerator:	Patients who were prescribed ACE inhibitor or ARB or ARNI therapy either within a 12-month period when seen in the outpatient setting OR at each hospital discharge.		
Denominator Exceptions:	Documentation of medical reason(s) for not prescribing ACE inhibitor or ARB or ARNI therapy (e.g., hypotensive patients who are at immediate risk of cardiogenic shock, hospitalized patients who have experienced marked azotemia, allergy, intolerance, other medical reasons). Documentation of patient reason(s) for not prescribing ACE inhibitor or ARB or ARNI therapy (e.g., patient declined, other patient reasons). Documentation of system reason(s) for not prescribing ACE inhibitor or ARB or ARNI therapy (e.g., other system reasons).		
Complete the following information:			
* Denominator 1:	* Numerator 1:	* Performance Rate 1 (%):	* Exception 1:
<input type="text" value="22"/>	<input type="text" value="6"/>	<input type="text" value="44.00"/>	<input type="text" value="44"/>

To satisfy this eQCM, enter a whole number into the Denominator, Numerator, Performance Rate, and Exception boxes.

When final selections have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen. Selections will not be saved.
- Click **Next** to move on to the next screen. Selections will be saved.
- Click **Save** to save selections and stay on the current screen.
- Click **Cancel** to remove selections and stay on the current screen.

9.5.19 CMS144

Electronic Clinical Quality Measures (Year 3 Attestation)

Questionnaire 19 of 47

(*) Red asterisk indicates a required field.

Measure:	CMS144/NQF 0083e		
Versions:	CMS144v9.2		
Title:	Heart Failure (HF): Beta-Blocker Therapy for Left Ventricular Systolic Dysfunction (LVSD)		
Description:	Percentage of patients aged 18 years and older with a diagnosis of heart failure (HF) with a current or prior left ventricular ejection fraction (LVEF) < 40% who were prescribed beta-blocker therapy either within a 12-month period when seen in the outpatient setting OR at each hospital discharge.		
Denominator:	All patients aged 18 years and older with a diagnosis of heart failure with a current or prior LVEF < 40%.		
Numerator:	Patients who were prescribed beta-blocker therapy either within a 12-month period when seen in the outpatient setting OR at each hospital discharge.		
Denominator Exceptions:	Documentation of medical reason(s) for not prescribing beta-blocker therapy (e.g., low blood pressure, fluid overload, asthma, patients recently treated with an intravenous positive inotropic agent, allergy, intolerance, other medical reasons). Documentation of patient reason(s) for not prescribing beta-blocker therapy (e.g., patient declined, other patient reasons). Documentation of system reason(s) for not prescribing beta-blocker therapy (e.g., other reasons attributable to the healthcare system).		
Complete the following information:			
* Denominator 1:	* Numerator 1:	* Performance Rate 1 (%):	* Exception 1:
<input type="text" value="55"/>	<input type="text" value="55"/>	<input type="text" value="6.00"/>	<input type="text" value="77"/>

Previous
Next
Save
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To satisfy this eCQM, enter a whole number into the Denominator, Numerator, Performance Rate, and Exception boxes.

When final selections have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen. Selections will not be saved.
- Click **Next** to move on to the next screen. Selections will be saved.
- Click **Save** to save selections and stay on the current screen.
- Click **Cancel** to remove selections and stay on the current screen.

9.5.20 CMS143

Electronic Clinical Quality Measures (Year 3 Attestation)

Questionnaire 20 of 47

(*) Red asterisk indicates a required field.

Measure:	CMS143/NQF 0086e
Versions:	CMS143v9.2
Title:	Primary Open-Angle Glaucoma (POAG): Optic Nerve Evaluation
Description:	Percentage of patients aged 18 years and older with a diagnosis of primary open-angle glaucoma (POAG) who have an optic nerve head evaluation during one or more office visits within 12 months.

Denominator:	All patients aged 18 years and older with a diagnosis of primary open-angle glaucoma.
Numerator:	Patients who have an optic nerve head evaluation during one or more office visits within 12 months.
Denominator Exceptions:	Documentation of medical reason(s) for not performing an optic nerve head evaluation.

Complete the following information:

* Denominator 1:	* Numerator 1:	* Performance Rate 1 (%):	* Exception 1:
<input type="text" value="1"/>	<input type="text" value="1"/>	<input type="text" value="1.00"/>	<input type="text" value="1"/>

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To satisfy this eCQM, enter a whole number into the Denominator, Numerator, Performance Rate, and Exception boxes.

When final selections have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen. Selections will not be saved.
- Click **Next** to move on to the next screen. Selections will be saved.
- Click **Save** to save selections and stay on the current screen.
- Click **Cancel** to remove selections and stay on the current screen.

9.5.21 CMS142

Electronic Clinical Quality Measures (Year 3 Attestation)

Questionnaire 21 of 47

(* Red asterisk indicates a required field.)

Measure:	CMS142/NQF XXXX		
Versions:	CMS142v9.2		
Title:	Diabetic Retinopathy: Communication with the Physician Managing Ongoing Diabetes Care		
Description:	Percentage of patients aged 18 years and older with a diagnosis of diabetic retinopathy who had a dilated macular or fundus exam performed with documented communication to the physician who manages the ongoing care of the patient with diabetes mellitus regarding the findings of the macular or fundus exam at least once within 12 months.		
Denominator:	All patients aged 18 years and older with a diagnosis of diabetic retinopathy who had a dilated macular or fundus exam performed.		
Numerator:	Patients with documentation, at least once within 12 months, of the findings of the dilated macular or fundus exam via communication to the physician who manages the patient's diabetic care.		
Denominator Exceptions:	Documentation of medical reason(s) for not communicating the findings of the dilated macular or fundus exam to the physician who manages the ongoing care of the patient with diabetes. Documentation of patient reason(s) for not communicating the findings of the dilated macular or fundus exam to the physician who manages the ongoing care of the patient with diabetes.		
Complete the following information:			
* Denominator 1:	* Numerator 1:	* Performance Rate 1 (%):	* Exception 1:
<input type="text" value="1"/>	<input type="text" value="1"/>	<input type="text" value="22.00"/>	<input type="text" value="2"/>

To satisfy this eCQM, enter a whole number into each of the Denominator, Numerator, Performance Rate, and Exception boxes.

When final selections have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen. Selections will not be saved.
- Click **Next** to move on to the next screen. Selections will be saved.
- Click **Save** to save selections and stay on the current screen.
- Click **Cancel** to remove selections and stay on the current screen.

9.5.22 CMS139

Electronic Clinical Quality Measures (Year 3 Attestation)

Questionnaire 22 of 47

(*) Red asterisk indicates a required field.

Measure:	CMS139/NQF XXXX		
Versions:	CMS139v9.2		
Title:	Falls: Screening for Future Fall Risk		
Description:	Percentage of patients 65 years of age and older who were screened for future fall risk during the measurement period.		
Denominator:	Patients aged 65 years and older with a visit during the measurement period.		
Numerator:	Patients who were screened for future fall risk at least once within the measurement period.		
Denominator Exclusions:	Exclude patients whose hospice care overlaps the measurement period.		
Complete the following information:			
* Denominator 1:	* Numerator 1:	* Performance Rate 1 (%):	* Exclusion 1:
<input type="text" value="22"/>	<input type="text" value="22"/>	<input type="text" value="222.00"/>	<input type="text" value="5"/>

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To satisfy this eCQM, enter a whole number into the Denominator, Numerator, Performance Rate, and Exclusion boxes.

When final selections have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen. Selections will not be saved.
- Click **Next** to move on to the next screen. Selections will be saved.
- Click **Save** to save selections and stay on the current screen.
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9.5.23 CMS161

Electronic Clinical Quality Measures (Year 3 Attestation)

Questionnaire 23 of 47

(*) Red asterisk indicates a required field.

Measure:	CMS161/NQF 0104e	
Versions:	CMS161v9.2	
Title:	Adult Major Depressive Disorder (MDD): Suicide Risk Assessment	
Description:	All patient visits during which a new diagnosis of MDD or new diagnosis of recurrent MDD was identified for patients aged 18 years and older with a suicide risk assessment completed during the visit.	
Denominator:	Patient visits during which a new diagnosis of MDD, single or recurrent episode, was identified.	
Numerator:	Patient visits during which a new diagnosis of MDD, single or recurrent episode, was identified and a suicide risk assessment was completed during the visit.	
Complete the following information:		
* Denominator 1:	* Numerator 1:	* Performance Rate 1 (%):
<input type="text" value="35"/>	<input type="text" value="35"/>	<input type="text" value="3.00"/>



To satisfy this eCQM, enter a whole number into the Denominator, Numerator, and Performance Rate boxes.

When final selections have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen. Selections will not be saved.

- Click **Next** to move on to the next screen. Selections will be saved.
- Click **Save** to save selections and stay on the current screen.
- Click **Cancel** to remove selections and stay on the current screen.

9.5.24 CMS128

Electronic Clinical Quality Measures (Year 3 Attestation)

Questionnaire 24 of 47

(*) Red asterisk indicates a required field.

Measure:	CMS128/NQF XXXX		
Versions:	CMS128v9.2		
Title:	Anti-depressant Medication Management		
Description:	<p>Percentage of patients 18 years of age and older who were treated with antidepressant medication, had a diagnosis of major depression, and who remained on an antidepressant medication treatment. Two rates are reported.</p> <p>a. Percentage of patients who remained on an antidepressant medication for at least 84 days (12 weeks).</p> <p>b. Percentage of patients who remained on an antidepressant medication for at least 180 days (6 months).</p>		
Denominator:	Denominator 1: Patients 18 years of age and older who were dispensed antidepressant medications within 245 days (8 months) prior to the measurement period through the first 120 days (4 months) of the measurement period, and were diagnosed with major depression 60 days prior to, or 60 days after the dispensing event and had a visit 60 days prior to, or 60 days after the dispensing event.		
Numerator:	<p>Numerator 1: Patients who have received antidepressant medication for at least 84 days (12 weeks) of continuous treatment during the 114-day period following the Index Prescription Start Date.</p> <p>Numerator 2: Patients who have received antidepressant medications for at least 180 days (6 months) of continuous treatment during the 231-day period following the Index Prescription Start Date.</p>		
Denominator Exclusions:	<p>Patients who were actively on an antidepressant medication in the 105 days prior to the Index Prescription Start Date.</p> <p>Exclude patients whose hospice care overlaps the measurement period.</p>		
Complete the following information:			
* Denominator 1:	* Numerator 1:	* Performance Rate 1 (%):	* Exclusion 1:
<input type="text" value="1"/>	<input type="text" value="1"/>	<input type="text" value="1.00"/>	<input type="text" value="1"/>
* Denominator 2:	* Numerator 2:	* Performance Rate 2 (%):	* Exclusion 2:
<input type="text" value="2"/>	<input type="text" value="2"/>	<input type="text" value="2.00"/>	<input type="text" value="2"/>

To satisfy this eCQM, enter a whole number into the Denominator, Numerator, Performance Rate, and Exclusion boxes.

When final selections have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen. Selections will not be saved.
- Click **Next** to move on to the next screen. Selections will be saved.
- Click **Save** to save selections and stay on the current screen.
- Click **Cancel** to remove selections and stay on the current screen.

9.5.25 CMS136

Electronic Clinical Quality Measures (Year 3 Attestation)

Questionnaire 25 of 47

(* Red asterisk indicates a required field.)

Measure:	CMS136/NQF XXXX
Versions:	CMS136v10.2
Title:	Follow-Up Care for Children Prescribed ADHD Medication (ADD)
Description:	<p>Percentage of children 6-12 years of age and newly dispensed a medication for attention-deficit/hyperactivity disorder (ADHD) who had appropriate follow-up care. Two rates are reported.</p> <p>a. Percentage of children who had one follow-up visit with a practitioner with prescribing authority during the 30-Day Initiation Phase.</p> <p>b. Percentage of children who remained on ADHD medication for at least 210 days and who, in addition to the visit in the Initiation Phase, had at least two additional follow-up visits with a practitioner within 270 days (9 months) after the Initiation Phase ended.</p>
Denominator:	<p>Denominator 1: Initial Population 1: Children 6-12 years of age who were dispensed an ADHD medication during the Intake Period and who had a visit during the measurement period.</p> <p>Denominator 2: Initial Population 2: Children 6-12 years of age who were dispensed an ADHD medication during the Intake Period and who remained on the medication for at least 210 days out of the 300 days following the IPSP, and who had a visit during the measurement period.</p>
Numerator:	<p>Numerator 1: Patients who had at least one face-to-face visit with a practitioner with prescribing authority within 30 days after the IPSP.</p> <p>Numerator 2: Patients who had at least one face-to-face visit with a practitioner with prescribing authority during the Initiation Phase, and at least two follow-up visits during the Continuation and Maintenance Phase. One of the two visits during the Continuation and Maintenance Phase may be a telephone visit with a practitioner.</p>

Denominator Exclusions:

Exclusion 1: Exclude patients diagnosed with narcolepsy at any point in their history or during the measurement period.
 Exclude patients who had an acute inpatient stay with a principal diagnosis of mental health or substance abuse during the 30 days after the IPSD.
 Exclude patients who were actively on an ADHD medication in the 120 days prior to the Index Prescription Start Date.
 Exclude patients whose hospice care overlaps the measurement period.

Exclusion 2: Exclude patients diagnosed with narcolepsy at any point in their history or during the measurement period.
 Exclude patients who had an acute inpatient stay with a principal diagnosis of mental health or substance abuse during the 300 days after the IPSD.
 Exclude patients who were actively on an ADHD medication in the 120 days prior to the Index Prescription Start Date.
 Exclude patients whose hospice care overlaps the measurement period.

Complete the following information:

* Denominator 1:	* Numerator 1:	* Performance Rate 1 (%):	* Exclusion 1:
2	2	2.00	2
* Denominator 2:	* Numerator 2:	* Performance Rate 2 (%):	* Exclusion 2:
2	2	2.00	2

Previous
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To satisfy this eCQM, enter a whole number into the Denominator, Numerator, Performance Rate, and Exclusion boxes.

When final selections have been made, choose a navigation button at the bottom of the screen.

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9.5.26 CMS157

Electronic Clinical Quality Measures (Year 3 Attestation)

Questionnaire 26 of 47

(* Red asterisk indicates a required field.)

Measure:	CMS157/NQF 0384e
Versions:	CMS157v9.2

Title:	Oncology: Medical and Radiation - Pain Intensity Quantified	
Description:	Percentage of patient visits, regardless of patient age, with a diagnosis of cancer currently receiving chemotherapy or radiation therapy in which pain intensity is quantified.	
Denominator:	All patient visits, regardless of patient age, with a diagnosis of cancer currently receiving chemotherapy or radiation therapy.	
Numerator:	Patient visits in which pain intensity is quantified.	
Complete the following information:		
* Denominator 1:	* Numerator 1:	* Performance Rate 1 (%):
<input type="text" value="0"/>	<input type="text" value="0"/>	<input type="text" value="0.00"/>

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To satisfy this eCQM, enter a whole number into the Denominator, Numerator, and Performance Rate boxes.

When final selections have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen. Selections will not be saved.
- Click **Next** to move on to the next screen. Selections will be saved.
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9.5.27 CMS129

Electronic Clinical Quality Measures (Year 3 Attestation)

Questionnaire 27 of 47

(*) Red asterisk indicates a required field.

Measure:	CMS129/NQF 0389e
Versions:	CMS129v10.3
Title:	Prostate Cancer: Avoidance of Overuse of Bone Scan for Staging Low Risk Prostate Cancer Patients

Description:	Percentage of patients, regardless of age, with a diagnosis of prostate cancer at low (or very low) risk of recurrence receiving interstitial prostate brachytherapy, OR external beam radiotherapy to the prostate, OR radical prostatectomy who did not have a bone scan performed at any time since diagnosis of prostate cancer.		
Denominator:	All patients, regardless of age, with a diagnosis of prostate cancer at low (or very low) risk of recurrence receiving interstitial prostate brachytherapy, OR external beam radiotherapy to the prostate, OR radical prostatectomy.		
Numerator:	Patients who did not have a bone scan performed at any time since diagnosis of prostate cancer.		
Denominator Exceptions:	Documentation of reason(s) for performing a bone scan (including documented pain, salvage therapy, other medical reasons, bone scan ordered by someone other than reporting physician).		
Complete the following information:			
* Denominator 1:	* Numerator 1:	* Performance Rate 1 (%):	* Exception 1:
<input type="text" value="2"/>	<input type="text" value="2"/>	<input type="text" value="2.00"/>	<input type="text" value="2"/>

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To satisfy this eCQM, enter a whole number into the Denominator, Numerator, Performance Rate, and Exception boxes.

When final selections have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen. Selections will not be saved.
- Click **Next** to move on to the next screen. Selections will be saved.
- Click **Save** to save selections and stay on the current screen.
- Click **Cancel** to remove selections and stay on the current screen.

9.5.28 CMS2

Electronic Clinical Quality Measures (Year 3 Attestation)

Questionnaire 28 of 47

(*) Red asterisk indicates a required field.

Measure:	CMS2/NQF 0418e
Versions:	CMS2v10.2

Title:	Preventive Care and Screening: Screening for Depression and Follow-Up Plan		
Description:	Percentage of patients aged 12 years and older screened for depression on the date of the encounter or 14 days prior to the date of the encounter using an age-appropriate standardized depression screening tool AND if positive, a follow-up plan is documented on the date of the eligible encounter.		
Denominator:	All patients aged 12 years and older at the beginning of the measurement period with at least one eligible encounter during the measurement period.		
Numerator:	Patients screened for depression on the date of the encounter or up to 14 days prior to the date of the encounter using an age-appropriate standardized tool AND if positive, a follow-up plan is documented on the date of the eligible encounter.		
Denominator Exclusions:	Patients who have been diagnosed with depression or with bipolar disorder.		
Denominator Exceptions:	Patient Reason(s) Patient refuses to participate OR Medical Reason(s) Documentation of medical reason for not screening patient for depression (e.g., cognitive, functional, or motivational limitations that may impact accuracy of results; patient is an urgent or emergent situation where time is of the essence and to delay treatment would jeopardize the patient's health status).		
Complete the following information:			
Complete the following information:			
* Denominator 1:	* Numerator 1:	* Performance Rate 1 (%):	* Exclusion 1:
<input type="text" value="33"/>	<input type="text" value="33"/>	<input type="text" value="3.00"/>	<input type="text" value="3333"/>
* Exception 1:			
<input type="text" value="333"/>			

To satisfy this eCQM, enter a whole number into the Denominator, Numerator, Performance Rate, Exclusion, and Exception boxes.

When final selections have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen. Selections will not be saved.
- Click **Next** to move on to the next screen. Selections will be saved.
- Click **Save** to save selections and stay on the current screen.
- Click **Cancel** to remove selections and stay on the current screen.

9.5.29 CMS68

Electronic Clinical Quality Measures (Year 3 Attestation)

Questionnaire 29 of 47

(*) Red asterisk indicates a required field.

Measure:	CMS68/NQF 0419e
Versions:	CMS68v10.3
Title:	Documentation of Current Medications in the Medical Record
Description:	Percentage of visits for patients aged 18 years and older for which the eligible professional or eligible clinician attests to documenting a list of current medications using all immediate resources available on the date of the encounter.
Denominator:	All visits occurring during the 12 month measurement period for patients aged 18 years and older.
Numerator:	Eligible professional or eligible clinician attests to documenting, updating or reviewing the patient's current medications using all immediate resources available on the date of the encounter.
Denominator Exceptions:	Documentation of a medical reason(s) for not documenting, updating, or reviewing the patient's current medications list (e.g., patient is in an urgent or emergent medical situation where time is of the essence and to delay treatment would jeopardize the patient's health status).

Complete the following information:

* Denominator 1:	* Numerator 1:	* Performance Rate 1 (%):	* Exception 1:
<input type="text" value="22"/>	<input type="text" value="22"/>	<input type="text" value="0.00"/>	<input type="text" value="222"/>

[Previous](#)
[Next](#)
[Save](#)
[Cancel](#)

To satisfy this eCQM, enter a whole number into each of the Denominator, Numerator, Performance Rate, and Exception boxes.

When final selections have been made, choose a navigation button at the bottom of the screen.

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- Click **Save** to save selections and stay on the current screen.
- Click **Cancel** to remove selections and stay on the current screen.

9.5.30 CMS69

Electronic Clinical Quality Measures (Year 3 Attestation)

Questionnaire 30 of 47

(*) Red asterisk indicates a required field.

Measure:	CMS69/NQF XXXX
Versions:	CMS69v9.3
Title:	Preventive Care and Screening: Body Mass Index (BMI) Screening and Follow-Up Plan
Description:	Percentage of patients aged 18 years and older with a BMI documented during the current encounter or within the previous twelve months AND who had a follow-up plan documented if most recent BMI was outside of normal parameters.
Denominator:	All patients aged 18 and older on the date of the encounter with at least one eligible encounter during the measurement period.
Numerator:	Patients with a documented BMI during the encounter or during the previous twelve months, AND when the BMI is outside of normal parameters, a follow-up plan is documented during the encounter or during the previous twelve months of the current encounter.
Denominator Exclusions:	1. Patients who are pregnant. 2. Patients receiving palliative or hospice care.
Denominator Exceptions:	Patients with a documented medical reason for not documenting BMI or for not documenting a follow-up plan for a BMI outside normal parameters (e.g., elderly patients (65 or older) for whom weight reduction/weight gain would complicate other underlying health conditions such as illness or physical disability, mental illness, dementia, confusion or nutritional deficiency such as vitamin/mineral deficiency; patients in an urgent or emergent medical situation where time is of the essence and to delay treatment would jeopardize the patient's health status).

Patients who refuse measurement of height and/or weight.

Complete the following information:

* Denominator 1:	* Numerator 1:	* Performance Rate 1 (%):	* Exclusion 1:
<input type="text" value="22"/>	<input type="text" value="22"/>	<input type="text" value="222.00"/>	<input type="text" value="22"/>
* Exception 1:			
<input type="text" value="22"/>			

[Previous](#)[Next](#)[Save](#)[Cancel](#)

To satisfy this eCQM, enter a whole number into the Denominator, Numerator, Performance Rate, Exclusion, and Exception boxes.

When final selections have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen. Selections will not be saved.
- Click **Next** to move on to the next screen. Selections will be saved.
- Click **Save** to save selections and stay on the current screen.
- Click **Cancel** to remove selections and stay on the current screen.

9.5.31 CMS133

Electronic Clinical Quality Measures (Year 3 Attestation)

Questionnaire 31 of 47

(*) Red asterisk indicates a required field.

Measure:	CMS133/NQF 0565e		
Versions:	CMS133v9.2		
Title:	Cataracts: 20/40 or Better Visual Acuity within 90 Days Following Cataract Surgery		
Description:	Percentage of cataract surgeries for patients aged 18 and older with a diagnosis of uncomplicated cataract and no significant ocular conditions impacting the visual outcome of surgery and had best-corrected visual acuity of 20/40 or better (distance or near) achieved in the operative eye within 90 days following the cataract surgery.		
Denominator:	All cataract surgeries for patients aged 18 years and older who did not meet any exclusion criteria.		
Numerator:	Cataract surgeries with best-corrected visual acuity of 20/40 or better (distance or near) achieved in the operative eye within 90 days following cataract surgery.		
Denominator Exclusions:	Cataract surgeries in patients with significant ocular conditions impacting the visual outcome of surgery.		
Complete the following information:			
* Denominator 1:	* Numerator 1:	* Performance Rate 1 (%):	* Exclusion 1:
<input type="text" value="2"/>	<input type="text" value="0"/>	<input type="text" value="2.00"/>	<input type="text" value="5"/>

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To satisfy this eCQM, enter a whole number into the Denominator, Numerator, Performance Rate and Exclusion boxes.

When final selections have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen. Selections will not be saved.
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- Click **Save** to save selections and stay on the current screen.

- Click **Cancel** to remove selections and stay on the current screen.

9.5.32 CMS159

Electronic Clinical Quality Measures (Year 3 Attestation)

Questionnaire 32 of 47

(*) Red asterisk indicates a required field.

Measure:	CMS159/NQF 0710e
Versions:	CMS159v9.4
Title:	Depression Remission at Twelve Months
Description:	The percentage of adolescent patients 12 to 17 years of age and adult patients 18 years of age or older with major depression or dysthymia who reached remission 12 months (+/-60 days) after an index event.
Denominator:	Adolescent patients 12 to 17 years of age and adult patients 18 years of age and older with a diagnosis of major depression or dysthymia and an initial PHQ-9 or PHQ-9M score greater than nine during the index event. Patients may be screened using PHQ-9 and PHQ-9M up to 7 days prior to the office visit (including the day of the office visit).
Numerator:	Adolescent patients 12 to 17 years of age and adult patients 18 years of age and older who achieved remission at twelve months as demonstrated by a twelve month (+/- 60 days) PHQ-9 or PHQ-9M score of less than five.
Denominator Exclusions:	<ol style="list-style-type: none"> 1. Patients who died. 2. Patients who received hospice or palliative care services. 3. Patients who were permanent nursing home residents. 4. Patients with a diagnosis of bipolar disorder. 5. Patients with a diagnosis of personality disorder emotionally labile. 6. Patients with a diagnosis of schizophrenia or psychotic disorder. 7. Patients with a diagnosis of pervasive developmental disorder.

Complete the following information:

Stratum 1: Ages 12 to 17 at the time of the index assessment

* Denominator 1:	* Numerator 1:	* Performance Rate 1 (%):	* Exclusion 1:
11	11	2.00	11

Stratum 2: Ages 18 and older at the time of the index assessment

* Denominator 1:	* Numerator 1:	* Performance Rate 1 (%):	* Exclusion 1:
11	11	11.00	11



To satisfy this eCQM, enter a whole number into the Denominator, Numerator, Performance Rate, and Exclusion boxes.

When final selections have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen. Selections will not be saved.
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- Click **Cancel** to remove selections and stay on the current screen.

9.5.33 CMS177

Electronic Clinical Quality Measures (Year 3 Attestation)

Questionnaire 33 of 47

(*) Red asterisk indicates a required field.

Measure:	CMS177/NQF 1365e
Versions:	CMS177v9.2
Title:	Child and Adolescent Major Depressive Disorder (MDD): Suicide Risk Assessment
Description:	Percentage of patient visits for those patients aged 6 through 17 years with a diagnosis of major depressive disorder with an assessment for suicide risk.

Denominator: All patient visits for those patients aged 6 through 17 years with a diagnosis of major depressive disorder.

Numerator: Patient visits with an assessment for suicide risk.

Complete the following information:

* Denominator 1: * Numerator 1: * Performance Rate 1 (%):

To satisfy this eCQM, enter a whole number into the Denominator, Numerator, and Performance Rate boxes.

When final selections have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen. Selections will not be saved.
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- Click **Save** to save selections and stay on the current screen.
- Click **Cancel** to remove selections and stay on the current screen.

9.5.34 CMS125

Electronic Clinical Quality Measures (Year 3 Attestation)

Questionnaire 34 of 47

(*) Red asterisk indicates a required field.

Measure:	CMS125/NQF XXXX
Versions:	CMS125v9.2
Title:	Breast Cancer Screening
Description:	Percentage of women 50-74 years of age who had a mammogram to screen for breast cancer in the 27 months prior to the end of the Measurement Period.

Denominator: Women 51-74 years of age with a visit during the measurement period.

Numerator: Women with one or more mammograms during the 27 months prior to the end of the measurement period.

Denominator Exclusions: Women who had a bilateral mastectomy or who have a history of a bilateral mastectomy or for whom there is evidence of a right and a left unilateral mastectomy.
 Exclude patients whose hospice care overlaps the measurement period.
 Exclude patients 66 and older who are living long term in an institution for more than 90 consecutive days during the measurement period.
 Exclude patients 66 and older with advanced illness and frailty because it is unlikely that patients will benefit from the service being measured.

Complete the following information:

* Denominator 1:	* Numerator 1:	* Performance Rate 1 (%):	* Exclusion 1:
0	0	6.00	8



To satisfy this eCQM, enter a whole number into the Denominator, Numerator, Performance Rate, and Exclusion boxes.

When final selections have been made, choose a navigation button at the bottom of the screen.

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9.5.35 CMS149

Electronic Clinical Quality Measures (Year 3 Attestation)

Questionnaire 35 of 47

(*) Red asterisk indicates a required field.

Measure:	CMS149/NQF 2872e		
Versions:	CMS149v9.2		
Title:	Dementia: Cognitive Assessment		
Description:	Percentage of patients, regardless of age, with a diagnosis of dementia for whom an assessment of cognition is performed and the results reviewed at least once within a 12-month period.		
Denominator:	All patients, regardless of age, with a diagnosis of dementia.		
Numerator:	Patients for whom an assessment of cognition is performed and the results reviewed at least once within a 12-month period.		
Denominator Exceptions:	Documentation of patient reason(s) for not assessing cognition.		
	Complete the following information:		
* Denominator 1:	* Numerator 1:	* Performance Rate 1 (%):	* Exception 1:
<input type="text" value="4"/>	<input type="text" value="3"/>	<input type="text" value="6.00"/>	<input type="text" value="3"/>

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Next

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To satisfy this eCQM, enter a whole number into the Denominator, Numerator, Performance Rate, and Exception boxes.

When final selections have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen. Selections will not be saved.
- Click **Next** to move on to the next screen. Selections will be saved.
- Click **Save** to save selections and stay on the current screen.
- Click **Cancel** to remove selections and stay on the current screen.

9.5.36 CMS22

Electronic Clinical Quality Measures (Year 3 Attestation)

Questionnaire 36 of 47

(*) Red asterisk indicates a required field.

Measure:	CMS22/NQF XXXX		
Versions:	CMS22v9.3		
Title:	Preventive Care and Screening: Screening for High Blood Pressure and Follow-Up Documented		
Description:	Percentage of patient visits for patients aged 18 years and older seen during the measurement period who were screened for high blood pressure AND a recommended follow-up plan is documented, as indicated, if blood pressure is pre-hypertensive or hypertensive.		
Denominator:	All patient visits for patients aged 18 years and older at the beginning of the measurement period.		
Numerator:	Patient visits where patients were screened for high blood pressure AND have a recommended follow-up plan documented, as indicated, if the blood pressure is pre-hypertensive or hypertensive.		
Denominator Exclusions:	Patient has an active diagnosis of hypertension.		
Denominator Exceptions:	Documentation of medical reason(s) for not screening for high blood pressure (e.g., patient is in an urgent or emergent medical situation where time is of the essence and to delay treatment would jeopardize the patient's health status). Documentation of patient reason(s) for not screening for blood pressure measurements or for not ordering an appropriate follow-up intervention if patient is pre-hypertensive or hypertensive (e.g., patient refuses).		
Complete the following information:			
* Denominator 1:	* Numerator 1:	* Performance Rate 1 (%):	* Exclusion 1:
<input type="text" value="22"/>	<input type="text" value="22"/>	<input type="text" value="22.00"/>	<input type="text" value="22"/>
* Exception 1:			
<input type="text" value="22"/>			

To satisfy this eCQM, enter a whole number into the Denominator, Numerator, Performance Rate, Exclusion, and Exception boxes.

When final selections have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen. Selections will not be saved.
- Click **Next** to move on to the next screen. Selections will be saved.
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9.5.37 CMS50

Electronic Clinical Quality Measures (Year 3 Attestation)

Questionnaire 37 of 47

(*) Red asterisk indicates a required field.

Measure: CMS50/NQF XXXX

Versions: CMS50v9.2

Title: Closing the Referral Loop: Receipt of Specialist Report

Description: Percentage of patients with referrals, regardless of age, for which the referring provider receives a report from the provider to whom the patient was referred.

Denominator: Number of patients, regardless of age, who were referred by one provider to another provider, and who had a visit during the measurement period.

Numerator: Number of patients with a referral, for which the referring provider received a report from the provider to whom the patient was referred.

Complete the following information:

* Denominator 1:	* Numerator 1:	* Performance Rate 1 (%):
10	10	10.00

Previous
Next
Save
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To satisfy this eCQM, enter a whole number into the Denominator, Numerator, and Performance Rate boxes.

When final selections have been made, choose a navigation button at the bottom of the screen.

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9.5.38 CMS56

Electronic Clinical Quality Measures (Year 3 Attestation)

Questionnaire 38 of 47

(*) Red asterisk indicates a required field.

Measure:	CMS56/NQF XXXX		
Versions:	CMS56v9.2		
Title:	Functional Status Assessment for Total Hip Replacement		
Description:	Percentage of patients 18 years of age and older who received an elective primary total hip arthroplasty (THA) and completed a functional status assessment within 90 days prior to the surgery and in the 270-365 days after the surgery.		
Denominator:	Patients 19 years of age and older who had a primary total hip arthroplasty (THA) in the year prior to the measurement period and who had an outpatient encounter during the measurement period.		
Numerator:	Patients with patient-reported functional status assessment results (i.e., Veterans RAND 12-item health survey[VR-12], Patient-Reported Outcomes Measurement Information System [PROMIS]-10-Global Health, Hip Disability and Osteoarthritis Outcome Score [HOOS], HOOS Jr.) in the 90 days prior to or on the day of primary THA procedure, and in the 270-365 days after THA procedure.		
Denominator Exclusions:	Patients with two or more fractures indicating trauma at the time of the total hip arthroplasty or patients with severe cognitive impairment that overlaps the measurement period. Exclude patients whose hospice care overlaps the measurement period.		
Complete the following information:			
* Denominator 1:	* Numerator 1:	* Performance Rate 1 (%):	* Exclusion 1:
<input type="text" value="22"/>	<input type="text" value="2"/>	<input type="text" value="2.00"/>	<input type="text" value="22"/>

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To satisfy this eCQM, enter a whole number into the Denominator, Numerator, Performance Rate, and Exclusion boxes.

When final selections have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen. Selections will not be saved.
- Click **Next** to move on to the next screen. Selections will be saved.
- Click **Save** to save selections and stay on the current screen.
- Click **Cancel** to remove selections and stay on the current screen.

9.5.39 CMS66

Electronic Clinical Quality Measures (Year 3 Attestation)

Questionnaire 39 of 47

(*) Red asterisk indicates a required field.

Measure:	CMS66/NQF XXXX
Versions:	CMS66v9.3
Title:	Functional Status Assessment for Total Knee Replacement
Description:	Percentage of patients 18 years of age and older who received an elective primary total knee arthroplasty (TKA) and completed a functional status assessment within 90 days prior to the surgery and in the 270-365 days after the surgery.
Denominator:	Patients 19 years of age and older who had a primary total knee arthroplasty (TKA) in the year prior to the measurement period and who had an outpatient encounter during the measurement period.
Numerator:	Patients with patient-reported functional status assessment results (i.e., Veterans RAND 12-item health survey [VR-12], Patient-Reported Outcomes Measurement Information System [PROMIS]-10 Global Health, Knee Injury and Osteoarthritis Outcome Score [KOOS], KOOS Jr.) in the 90 days prior to or on the day of the primary TKA procedure, and in the 270-365 days after the TKA procedure.
Denominator Exclusions:	Patients with two or more fractures indicating trauma at the time of the total knee arthroplasty or patients with severe cognitive impairment that overlaps the measurement period. Exclude patients whose hospice care overlaps the measurement period.

Complete the following information:

* Denominator 1:	* Numerator 1:	* Performance Rate 1 (%):	* Exclusion 1:
<input type="text" value="0"/>	<input type="text" value="0"/>	<input type="text" value="0.00"/>	<input type="text" value="0"/>

To satisfy this eCQM, enter a whole number into the Denominator, Numerator Performance Rate and Exclusion boxes.

When final selections have been made, choose a navigation button at the bottom of the screen.

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9.5.40 CMS74

Electronic Clinical Quality Measures (Year 3 Attestation)

Questionnaire 40 of 47

(*) Red asterisk indicates a required field.

Measure:	CMS74/NQF XXXX
Versions:	CMS74v10.2
Title:	Primary Caries Prevention Intervention as Offered by Primary Care Providers, including Dentists
Description:	Percentage of children, 6 months - 20 years of age, who received a fluoride varnish application during the measurement period.
Denominator:	Children, 6 months - 20 years of age, with a visit during the measurement period.
Numerator:	Children who receive a fluoride varnish application during the measurement period.
Denominator Exclusions:	Exclude patients whose hospice care overlaps the measurement period.

Complete the following information:

Stratum 1: Population 1: age 6 months - 5 years

* Denominator 1:	* Numerator 1:	* Performance Rate 1 (%):	* Exclusion 1:
1	1	77.00	7

Stratum 2: Population 2: age 6-12

* Denominator 1:	* Numerator 1:	* Performance Rate 1 (%):	* Exclusion 1:
2	2	1.00	1

Stratum 3: Population 3: age 13-20

* Denominator 1:	* Numerator 1:	* Performance Rate 1 (%):	* Exclusion 1:
256	66	3.00	3

Previous
Next
Save
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To satisfy this eCQM, enter a whole number into the Denominator, Numerator, Performance Rate, and Exclusion boxes.

When final selections have been made, choose a navigation button at the bottom of the screen.

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- Click **Cancel** to remove selections and stay on the current screen.

9.5.41 CMS75

Electronic Clinical Quality Measures (Year 3 Attestation)

Questionnaire 41 of 47

(*) Red asterisk indicates a required field.

Measure:	CMS75/NQF XXXX
Versions:	CMS75v9.2
Title:	Children Who Have Dental Decay or Cavities
Description:	Percentage of children, 6 months - 20 years of age, who have had tooth decay or cavities during the measurement period.

Denominator: Children, 6 months - 20 years of age, with a clinical oral evaluation during the measurement period.

Numerator: Children who had a diagnosis of cavities or decayed teeth overlapping the measurement period.

Denominator Exclusions: Exclude patients whose hospice care overlaps the measurement period.

Complete the following information:

* Denominator 1: * Numerator 1: * Performance Rate 1 (%): * Exclusion 1:



To satisfy this eCQM, enter a whole number into the Denominator, Numerator, Performance Rate, and Exclusion boxes.

When final selections have been made, choose a navigation button at the bottom of the screen.

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- Click **Save** to save selections and stay on the current screen.
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9.5.42 CMS90

Electronic Clinical Quality Measures (Year 3 Attestation)

Questionnaire 42 of 47

(*) Red asterisk indicates a required field.

Measure:	CMS90/NQF XXXX
Versions:	CMS90v10.2
Title:	Functional Status Assessments for Congestive Heart Failure
Description:	Percentage of patients 18 years of age and older with congestive heart failure who completed initial and follow-up patient-reported functional status assessments.

Denominator: Patients 18 years of age and older who had two outpatient encounters during the measurement year and a diagnosis of congestive heart failure.

Numerator: Patients with patient-reported functional status assessment results (i.e., Veterans RAND 12-item health survey [VR-12]; VR-36; Kansas City Cardiomyopathy Questionnaire [KCCQ]; KCCQ-12; Minnesota Living with Heart Failure Questionnaire [MLHFQ]; Patient-Reported Outcomes Measurement Information System [PROMIS] -10 Global Health, PROMIS-29) present in the EHR two weeks before or during the initial FSA encounter and results for the follow-up FSA at least 30 days but no more than 180 days after the initial FSA.

Denominator Exclusions: Exclude patients with severe cognitive impairment that overlaps the measurement period.
Exclude patients whose hospice care overlaps the measurement period.

Complete the following information:

* Denominator 1:	* Numerator 1:	* Performance Rate 1 (%):	* Exclusion 1:
1	1	1.00	11



To satisfy this eCQM, enter a whole number into the Denominator, Numerator, Performance Rate, and Exclusion boxes.

When final selections have been made, choose a navigation button at the bottom of the screen.

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9.5.43 CMS249

Electronic Clinical Quality Measures (Year 3 Attestation)

Questionnaire 43 of 47

(*) Red asterisk indicates a required field.

Measure:	CMS249/NQF 3475e
Versions:	CMS249v3.2

Title:	Appropriate Use of DXA Scans in Women Under 65 Years Who Do Not Meet the Risk Factor Profile for Osteoporotic Fracture		
Description:	Percentage of female patients 50 to 64 years of age without select risk factors for osteoporotic fracture who received an order for a dual-energy x-ray absorptiometry (DXA) scan during the measurement period.		
Denominator:	Female patients ages 50 to 64 years with an encounter during the measurement period.		
Numerator:	Female patients who received an order for at least one DXA scan in the measurement period.		
Denominator Exclusions:	Exclude patients with a combination of risk factors (as determined by age) or one of the independent risk factors. Ages: 50-54 (>=4 combinations risk factors) or 1 independent risk factor. Ages: 55-59(>=3 combination risk factors) or 1 independent risk factor. Ages: 60-64 (>=2 combination risk factors) or 1 independent risk factor. For list of Risk Factors, please refer to the specifications sheet for the PY 2021.		
Complete the following information:			
* Denominator 1:	* Numerator 1:	* Performance Rate 1 (%):	* Exclusion 1:
<input type="text" value="3"/>	<input type="text" value="2"/>	<input type="text" value="10.00"/>	<input type="text" value="3"/>



To satisfy this eCQM, enter a whole number into the Denominator, Numerator, Performance Rate, and Exclusion boxes.

When final selections have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen. Selections will not be saved.
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9.5.44 CMS347

Electronic Clinical Quality Measures (Year 3 Attestation)

Questionnaire 44 of 47

(*) Red asterisk indicates a required field.

Measure:	CMS347/NQF XXXX
Versions:	CMS347v4.3
Title:	Statin Therapy for the Prevention and Treatment of Cardiovascular Disease
Description:	<p>Percentage of the following patients - all considered at high risk of cardiovascular events - who were prescribed or were on statin therapy during the measurement period:</p> <p>*Adults aged \geq 21 years who were previously diagnosed with or currently have an active diagnosis of clinical atherosclerotic cardiovascular disease (ASCVD); OR</p> <p>*Adults aged \geq 21 years who have ever had a fasting or direct low-density lipoprotein cholesterol (LDL-C) level \geq 190 mg/dL or were previously diagnosed with or currently have an active diagnosis of familial or pure hypercholesterolemia; OR</p> <p>*Adults aged 40-75 years with a diagnosis of diabetes with a fasting or direct LDL-C level of 70-189 mg/dL.</p>
Denominator:	<p>All patients who meet one or more of the following criteria (considered at "high risk" for cardiovascular events, under ACC/AHA guidelines): 1) Patients aged \geq 21 years at the beginning of the measurement period with clinical ASCVD diagnosis.</p> <p>Denominator 2: 2) Patients aged \geq 21 years at the beginning of the measurement period who have ever had a fasting or direct laboratory result of LDL-C \geq 190 mg/dL or were previously diagnosed with or currently have an active diagnosis of familial or pure hypercholesterolemia.</p> <p>Denominator 3: 3) Patients aged 40 to 75 years at the beginning of the measurement period with Type 1 or Type 2 diabetes and with an LDL-C result of 70-189 mg/dL recorded as the highest fasting or direct laboratory test result in the measurement year or during the two years prior to the beginning of the measurement period.</p>

Numerator: Patients who are actively using or who receive an order (prescription) for statin therapy at any point during the measurement period.

Denominator Exclusions:

1. Patients who have a diagnosis of pregnancy.
2. Patients who are breastfeeding.
3. Patients who have a diagnosis of rhabdomyolysis.

Denominator Exceptions:

1. Patients with adverse effect, allergy, or intolerance to statin medication.
2. Patients who are receiving palliative or hospice care.
3. Patients with active liver disease or hepatic disease or insufficiency.
4. Patients with end-stage renal disease (ESRD).
5. Patients with diabetes who have the most recent fasting or direct LDL-C laboratory test result < 70 mg/dL and are not taking statin therapy.

Complete the following information:

* Denominator 1:	* Numerator 1:	* Performance Rate 1 (%):	* Exclusion 1:
<input type="text" value="55"/>	<input type="text" value="11"/>	<input type="text" value="1.00"/>	<input type="text" value="1"/>
* Exception 1:	<input type="text" value="11"/>		
* Denominator 2:	* Numerator 2:	* Performance Rate 2 (%):	* Exclusion 2:
<input type="text" value="33"/>	<input type="text" value="33"/>	<input type="text" value="33.00"/>	<input type="text" value="33"/>
* Exception 2:	<input type="text" value="333"/>		
* Denominator 3:	* Numerator 3:	* Performance Rate 3 (%):	* Exclusion 3:
<input type="text" value="33"/>	<input type="text" value="5"/>	<input type="text" value="33.00"/>	<input type="text" value="33"/>
* Exception 3:	<input type="text" value="33"/>		

To satisfy this eCQM, enter a whole number into the Denominator, Numerator, Performance Rate, Exclusion, and Exception boxes.

When final selections have been made, choose a navigation button at the bottom of the screen.

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- Click **Save** to save selections and stay on the current screen.
- Click **Cancel** to remove selections and stay on the current screen.

9.5.45 CMS349

Electronic Clinical Quality Measures (Year 3 Attestation)

Questionnaire 45 of 47

(*) Red asterisk indicates a required field.

Measure:	CMS349/NQF XXXX		
Versions:	CMS349v3.3		
Title:	HIV Screening		
Description:	Percentage of patients aged 15-65 at the start of the measurement period who were between 15-65 years old when tested for HIV.		
Denominator:	Patients 15 to 65 years of age at the start of the measurement period AND who had at least one outpatient visit during the measurement period.		
Numerator:	Patients with documentation of an HIV test performed on or after their 15th birthday and before their 66th birthday.		
Denominator Exclusions:	Patients diagnosed with HIV prior to the start of the measurement period.		
Complete the following information:			
* Denominator 1:	* Numerator 1:	* Performance Rate 1 (%):	* Exclusion 1:
<input type="text" value="0"/>	<input type="text" value="0"/>	<input type="text" value="0.00"/>	<input type="text" value="55"/>

Previous

Next

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To satisfy this eCQM, enter a whole number into the Denominator, Numerator, Performance Rate, and Exclusion boxes.

When final selections have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen. Selections will not be saved.
- Click **Next** to move on to the next screen. Selections will be saved.
- Click **Save** to save selections and stay on the current screen.
- Click **Cancel** to remove selections and stay on the current screen.

9.5.46 CMS645

Electronic Clinical Quality Measures (Year 3 Attestation)

Questionnaire 46 of 47

(*) Red asterisk indicates a required field.

Measure:	CMS645/NQF XXXX		
Versions:	CMS645v4.1		
Title:	Bone density evaluation for patients with prostate cancer and receiving androgen deprivation therapy		
Description:	Patients determined as having prostate cancer who are currently starting or undergoing androgen deprivation therapy (ADT), for an anticipated period of 12 months or greater and who receive an initial bone density evaluation. The bone density evaluation must be prior to the start of ADT or within 3 months of the start of ADT.		
Denominator:	Male patients with a qualifying encounter in the measurement period AND with a diagnosis of prostate cancer AND with an order for ADT or an active medication of ADT with an intent for treatment greater than or equal to 12 months during the measurement period.		
Numerator:	Patients with a bone density evaluation within the two years prior to the start of or less than three months after the start of ADT treatment.		
Denominator Exceptions:	Patient refused recommendation for a bone density evaluation after the start of ADT therapy.		
Complete the following information:			
* Denominator 1:	* Numerator 1:	* Performance Rate 1 (%):	* Exception 1:
<input type="text" value="9"/>	<input type="text" value="9"/>	<input type="text" value="11.00"/>	<input type="text" value="9"/>

Previous
Next
Save
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To satisfy this eCQM, enter a whole number into the Denominator, Numerator, Performance Rate and Exception boxes.

When final selections have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen. Selections will not be saved.
- Click **Next** to move on to the next screen. Selections will be saved.

- Click **Save** to save selections and stay on the current screen.
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9.5.47 CMS771

Electronic Clinical Quality Measures (Year 3 Attestation)

Questionnaire 47 of 47

(*) Red asterisk indicates a required field.

Measure:	CMS771/NQF XXXX		
Versions:	CMS771v2.2		
Title:	Urinary Symptom Score Change 6-12 Months After Diagnosis of Benign Prostatic Hyperplasia		
Description:	Percentage of patients with an office visit within the measurement period and with a new diagnosis of clinically significant Benign Prostatic Hyperplasia who have International Prostate Symptoms Score (IPSS) or American Urological Association (AUA) Symptom Index (SI) documented at time of diagnosis and again 6-12 months later with an improvement of 3 points.		
Denominator:	Male patients with an initial diagnosis of benign prostatic hyperplasia 6 months prior to, or during the measurement period, and a urinary symptom score assessment within 1 month of initial diagnosis and a follow-up urinary symptom score assessment within 6-12 months, who had a qualifying visit during the measurement period.		
Numerator:	Patients with a documented improvement of at least 3 points in their urinary symptom score during the measurement period.		
Denominator Exclusions:	Patients with urinary retention that starts within 1 year of initial BPH diagnosis. Patients with an initial BPH diagnosis that starts during, or within 30 days of hospitalization. Patients with a diagnosis of morbid obesity, or with a BMI Exam >40 before the follow up urinary symptom score.		
Complete the following information:			
* Denominator 1:	* Numerator 1:	* Performance Rate 1 (%):	* Exclusion 1:
<input type="text" value="0"/>	<input type="text" value="0"/>	<input type="text" value="0.00"/>	<input type="text" value="0"/>

Previous
Next
Save
Cancel

To satisfy this eCQM, enter a whole number into the Denominator, Numerator, Performance Rate and Exclusion boxes.

When final selections have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen. Selections will not be saved.
- Click **Next** to move on to the next screen. Selections will be saved.
- Click **Save** to save selections and stay on the current screen.
- Click **Cancel** to remove selections and stay on the current screen.

10 Submitting Attestation

Prior to submitting the attestation for program staff review, EPs have the option to review and edit their responses with the Pre-Attestation Summary screens. EPs can navigate through screens and confirm their responses prior to submitting.

10.1 Pre-Attestation Summary Screen

Summary of Measures (Year 3 Attestation)

Please select the desired measure link below to review the details of your attestation. This is your last chance to view/edit the information you have entered before you attest. Please review your information as you will be unable to edit your information after you attest.

[Meaningful Use Objectives Summary](#)

[Public Health Objective Summary](#)

[Electronic Clinical Quality Measures Summary - Manual](#)

Previous
Next

Click on a link to review the summary.

When final reviews have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen. Selections will not be saved.
- Click **Next** to move on to the next screen. Selections will be saved.

10.1.1 Objectives Summary

Summary of Meaningful Use Core Measures (Year 2 Attestation)

Meaningful Use Core Measure List Table

Please select the edit link next to the measure you wish to update. If you do not wish to edit your measures you may select next to continue.

CORE OBJECTIVES SUMMARY

ObjectiveText	Description	Data Entered	Selection
Protect electronic protected health information (ePHI) created or maintained by the CEHRT through the implementation of appropriate technical, administrative, and physical safeguards.	Conduct or review a security risk analysis in accordance with the requirements under 45 CFR 164.308(a)(1), including addressing the security (including encryption) of data created or maintained by CEHRT in accordance with requirements under 45 CFR 164.312(a)(2)(iv) and 45 CFR 164.306(d)(3), implement security updates as necessary, and correct identified security deficiencies as part of the provider's risk management process.	Yes	Edit
Generate and transmit permissible prescriptions electronically.	More than 60% of all permissible prescriptions written by the EP are queried for a drug formulary and transmitted electronically using CEHRT.	Numerator = 50 Denominator = 77	Edit
Implement clinical decision support (CDS) interventions focused on improving performance on high-priority health conditions.	Implement five CDS interventions related to four or more clinical quality measures (CQMs) at a relevant point in patient care for the entire EHR reporting period. Absent four CQMs related to an EP's scope of practice or patient population, the CDS interventions must be related to high-priority health conditions.	Yes	Edit

<p>Implement clinical decision support (CDS) interventions focused on improving performance on high-priority health conditions.</p>	<p>Enable and implement the functionality for drug-drug and drug-allergy interaction checks for the entire EHR reporting period.</p>	<p>Yes</p>	<p>Edit</p>
<p>Use CPOE for medication, laboratory, and diagnostic imaging orders directly entered by any licensed healthcare professional, credentialed medical assistant, or a medical staff member credentialed to and performing the equivalent duties of a credentialed medical assistant, who can enter orders into the medical record per state, local, and professional guidelines.</p>	<p>More than 60% of medication orders created by the EP during the EHR reporting period are recorded using computerized provider order entry.</p>	<p>Numerator = 90 Denominator = 100</p>	<p>Edit</p>
<p>Use CPOE for medication, laboratory, and diagnostic imaging orders directly entered by any licensed healthcare professional, credentialed medical assistant, or a medical staff member credentialed to and performing the equivalent duties of a credentialed medical assistant, who can enter orders into the medical record per state, local, and professional guidelines.</p>	<p>More than 60% of laboratory orders created by the EP during the EHR reporting period are recorded using computerized provider order entry.</p>	<p>Numerator = 11110 Denominator = 11110</p>	<p>Edit</p>

<p>Use CPOE for medication, laboratory, and diagnostic imaging orders directly entered by any licensed healthcare professional, credentialed medical assistant, or a medical staff member credentialed to and performing the equivalent duties of a credentialed medical assistant, who can enter orders into the medical record per state, local, and professional guidelines.</p>	<p>More than 60% of diagnostic imaging orders created by the EP during the EHR reporting period are recorded using computerized provider order entry.</p>	<p>Numerator = 100 Denominator = 100</p>	<p>Edit</p>
<p>The EP provides patients (or patient-authorized representatives) with timely electronic access to their health information and patient-specific education.</p>	<p>For more than 80% of all unique patients seen by the EP: (1) The patient (or the patient-authorized representative) is provided timely access to view online, download, and transmit his or her health information; and (2) The provider ensures the patient's health information is available for the patient (or patient-authorized representative) to access using any application of their choice that is configured to meet the technical specifications of the API in the provider's CEHRT.</p>	<p>Numerator = 90 Denominator = 90</p>	<p>Edit</p>
<p>The EP provides patients (or patient-authorized representatives) with timely electronic access to their health information and patient-specific education.</p>	<p>The EP must use clinically relevant information from CEHRT to identify patient-specific educational resources and provide electronic access to those materials to more than 35% of unique patients seen by the EP during the EHR reporting period.</p>	<p>Numerator = 4 Denominator = 4</p>	<p>Edit</p>

<p>Use CEHRT to engage with patients or their authorized representatives about the patient's care.</p>	<p>More than 5% of all unique patients (or their authorized representatives) seen by the EP actively engage with the EHR made accessible by the EP and either: (1) View, download, or transmit to a third party their health information; or (2) Access their health information through an API that can be used by applications chosen by the patient and configured to the API in the EP's CEHRT; or (3) A combination of (1) and (2).</p>	<p>Numerator = 30 Denominator = 30</p>	<p>Edit</p>
<p>Use CEHRT to engage with patients or their authorized representatives about the patient's care.</p>	<p>For more than 5% of all unique patients seen by the EP during the EHR reporting period, a secure message was sent using the electronic messaging function of CEHRT to the patient (or the patient-authorized representative), or in response to a secure message sent by the patient or their authorized representative.</p>	<p>Numerator = 70 Denominator = 100</p>	<p>Edit</p>
<p>Use CEHRT to engage with patients or their authorized representatives about the patient's care.</p>	<p>Patient generated health data or data from a non-clinical setting is incorporated into the CEHRT for more than 5% of all unique patients seen by the EP during the EHR reporting period.</p>	<p>Numerator = 90 Denominator = 100</p>	<p>Edit</p>
<p>The EP provides a summary of care record when transitioning or referring their patient to another setting of care, receives or retrieves a summary of care record upon the receipt of a transition or referral or upon the first patient encounter with a new patient, and incorporates summary of care information from other providers into their EHR using the functions of CEHRT.</p>	<p>For more than 50% of transitions of care and referrals, the EP that transitions or refers their patient to another setting of care or provider of care: (1) Creates a summary of care record using CEHRT; and (2) Electronically exchanges the summary of care record.</p>	<p>Excluded</p>	<p>Edit</p>

<p>The EP provides a summary of care record when transitioning or referring their patient to another setting of care, receives or retrieves a summary of care record upon the receipt of a transition or referral or upon the first patient encounter with a new patient, and incorporates summary of care information from other providers into their EHR using the functions of CEHRT.</p>	<p>For more than 40% of transitions or referrals received and patient encounters in which the EP has never encountered the patient before, they incorporate into the patient's EHR an electronic summary of care document.</p>	<p>Excluded</p>	<p>Edit</p>
<p>The EP provides a summary of care record when transitioning or referring their patient to another setting of care, receives or retrieves a summary of care record upon the receipt of a transition or referral or upon the first patient encounter with a new patient, and incorporates summary of care information from other providers into their EHR using the functions of CEHRT.</p>	<p>For more than 80% of transitions or referrals received and patient encounters in which the EP has never encountered the patient before, they perform a clinical information reconciliation. The EP must implement clinical information reconciliation for the following three clinical information sets: (1) Medication. Review of the patient's medication, including the name, dosage, frequency, and route of each medication. (2) Medication allergy. Review of the patient's known medication allergies. (3) Current Problem list. Review of the patient's current and active diagnoses.</p>	<p>Excluded</p>	<p>Edit</p>

Previous
Next

The Objectives Summary lists each MU Objective attested to, with responses.

- If changes need to be made, click the Edit link for the MU Objective to update. This will redirect to the MU Objective details screen for changes to be made.
- It is important to be sure any changes are saved after edit is complete.

When final reviews/edits have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen. Selections will not be saved.
- Click **Next** to move on to the next screen. Selections will be saved.

10.1.2 Public Health Objectives Summary

Summary of Public Health Objective Measures (Year 5 Attestation)

Public Health Objective List Table

Please select the edit link next to the measure you wish to update. If you do not wish to edit your measures you may select next to continue.

PUBLIC HEALTH MEASURES SUMMARY			
ObjectiveText	Measure	Entered	Selection
The EP is in active engagement with a public health agency to submit electronic public health data from CEHRT except where prohibited and in accordance with applicable law and practice.	The EP is in active engagement with a public health agency to submit immunization data.	Option 2	Edit
The EP is in active engagement with a public health agency to submit electronic public health data from CEHRT except where prohibited and in accordance with applicable law and practice.	The EP is in active engagement with a public health agency to submit syndromic surveillance data.	Option 3	Edit
The EP is in active engagement with a public health agency to submit electronic public health data from CEHRT except where prohibited and in accordance with applicable law and practice.	The EP is in active engagement to submit data to a specialized registry.	Option 3 - KY Cancer Registry Option 1 - Skin	Edit

Previous
Next

The Public Health Objectives Summary lists each Public Health Measure attested to, with responses.

- If changes need to be made, click the Edit link for the PH Measure to update. This will redirect to the PH Measure details screen for changes to be made.
- It is important to be sure any changes are saved after edit is complete.

When final reviews/edits have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen. Selections will not be saved.
- Click **Next** to move on to the next screen. Selections will be saved.

10.1.3 Electronic Clinical Quality Measures Summary (Electronically Reported)

Electronic Clinical Quality Measures (Year 2 Attestation)

Electronic Clinical Quality Measures Summary Report

<u>Date Received</u>	<u>Report NPI</u>	<u>Reporting Period Start</u>	<u>Reporting Period End</u>
7/1/2020 1:01:24 PM	1851404065	3/1/2020 12:00:00 AM	6/22/2020 12:00:00 AM
<u>Document Type</u>	<u>eCQM Status:</u>	<u>Evaluated Date:</u>	
QRDA III	Accepted	7/1/2020 1:10:13 PM	
<u>File Name:</u>			
QRDA_XML2.xml, CPCSampleQRDA-III.xml			

QRDA_XML2_TESTFILE3.XML

Domain	# of Measures
Communication and Care Coordination	0
Community/Population Health	2
Effective Clinical Care	1
Efficiency and Cost Reduction	0
Patient Safety	1
Person and Caregiver-Centered Experience and Outcomes	0

CPCSAMPLEQRDA-III.XML

Domain	# of Measures
Communication and Care Coordination	1
Community/Population Health	4
Effective Clinical Care	9
Efficiency and Cost Reduction	0
Patient Safety	2
Person and Caregiver-Centered Experience and Outcomes	0

Total Required	<u>6</u>
Total Met	17
Final eCQM Result:	Submission Accepted, click Next to continue.
<div style="text-align: center;">Previous Evaluate eCQM Submissions Next</div>	

The Electronic Clinical Quality Measures Summary (Electronically Reported) lists each eCQM domain and number of measures attested to.

When final reviews/edits have been made, choose a navigation button at the bottom of the screen.

- Click Previous to go back to the previous screen. Selections will not be saved.
- Click Next to move on to the next screen. Selections will be saved.

10.1.4 Electronic Clinical Quality Measures Summary (Manually Reported)

Summary of Clinical Quality Measures (Year 4 Attestation)				
Clinical Quality Measures List Table				
<p>Please select the edit link next to the measure you wish to update. If you do not wish to edit your measures you may select next to continue.</p>				
PATIENT SAFETY				
Measure #	Title	Measure	Data Entered	Selection
CMS156v5.1/NQF 0022	Use of High-Risk Medications in the Elderly	Percentage of patients 66 years of age and older who were ordered high-risk medications. Two rates are reported. a. Percentage of patients who were ordered at least one high-risk medication. b. Percentage of patients who were ordered at least two different high-risk medications.	Denominator = 50 Numerator = 10 Performance Rate = 50.00 Denominator = 50 Numerator = 10 Performance Rate = 50.00	Edit
COMMUNITY/POPULATION HEALTH				
Measure #	Title	Measure	Data Entered	Selection
CMS117v5.1/NQF 0038	Childhood Immunization Status	Percentage of children 2 years of age who had four diphtheria, tetanus and acellular pertussis (DTaP); three polio (IPV), one measles, mumps and rubella (MMR); three H influenza type B (HiB); three hepatitis B (Hep B); one chicken pox (VZV); four pneumococcal conjugate (PCV); one hepatitis A (Hep A); two or three rotavirus (RV); and two influenza (flu) vaccines by their second birthday.	Denominator = 75 Numerator = 25 Performance Rate = 55.00	Edit

EFFECTIVE CLINICAL CARE				
Measure #	Title	Measure	Data Entered	Selection
CMS165v5.0/NQF 0018	Controlling High Blood Pressure	Percentage of patients 18-85 years of age who had a diagnosis of hypertension and whose blood pressure was adequately controlled (<140/90mmHg) during the measurement period.	Denominator = 100 Numerator = 50 Performance Rate = 50.00 Exclusion = 0	Edit
CMS135v5.2/NQF 0081	Heart Failure (HF): Angiotensin-Converting Enzyme (ACE) Inhibitor or Angiotensin Receptor Blocker (ARB) Therapy for Left Ventricular Systolic Dysfunction (LVSD)	Percentage of patients aged 18 years and older with a diagnosis of heart failure (HF) with a current or prior left ventricular ejection fraction (LVEF) < 40% who were prescribed ACE inhibitor or ARB therapy either within a 12 month period when seen in the outpatient setting OR at each hospital discharge.	Denominator = 50 Numerator = 30 Performance Rate = 35.00 Exception = 0 Exception = 0 Exception = 0	Edit
CMS142v5.2/NQF 0089	Diabetic Retinopathy: Communication with the Physician Managing Ongoing Diabetes Care	Percentage of patients aged 18 years and older with a diagnosis of diabetic retinopathy who had a dilated macular or fundus exam performed with documented communication to the physician who manages the ongoing care of the patient with diabetes mellitus regarding the findings of the macular or fundus exam at least once within 12 months.	Denominator = 50 Numerator = 10 Performance Rate = 50.00	Edit
CMS169v5.0/NQF XXXX	Bipolar Disorder and Major Depression: Appraisal for alcohol or chemical substance use	Percentage of patients with depression or bipolar disorder with evidence of an initial assessment that includes an appraisal for alcohol or chemical substance use.	Denominator = 75 Numerator = 35 Performance Rate = 50.00	Edit

Previous
Next

The Electronic Clinical Quality Measures Summary lists each eCQM attested to, with responses.

- If changes need to be made, click the Edit link for the eCQM to update. This will redirect to the eCQM details screen for changes to be made.
- It is important to be sure any changes are saved after edit is complete.

When final reviews/edits have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen. Selections will not be saved.
- Click **Next** to move on to the next screen. Selections will be saved.

10.2 Incentive Payment Calculation Screen

Incentive Payment Calculations (Year 2 Attestation)

Estimated Amount of Medicaid EHR Incentive Payment:	\$8,500.00
(This amount may also include adjustments)	

Previous
Next

The Incentive Payment Calculation screen is view only and provides the estimated amount of Medicaid EHR incentive payment.

When final reviews have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen. Selections will not be saved.
- Click **Next** to move on to the next screen. Selections will be saved.

10.3 Document Upload Screen

Document Upload (Year 3 Attestation)

Documentation needed to process your application may be attached below. If you cannot attach a PDF then use the Send E-mail link on the left to contact the EHR staff for assistance.

Documentation attached to the attestation does not alleviate the provider from being requested to produce additional documentation that may be requested during a pre-payment or post-payment audit. All documentation supporting the information attested should be kept for 6 years.

The following documents are required to submit with each Program Year of Participation:

1. Payment reassignment documentation if payment is assigned to any other NPI than the individual NPI.
2. Patient volume report. If you are using Medicaid patients from multiple states you could be requested to provide additional documentation.

The documents listed below are required to submit only if there has been a change from the previous Program Year:

3. Proof of CEHRT (Certified EHR Technology) being attested for your practice or facility. This can be: • a signed contract • a signed lease • a current invoice • a license agreement • a purchase order (PO) • or other legal documents showing that you have contracted with a certified EHR vendor.
4. KHIE on-boarding documentation. (Signed Participation Agreement's, MU Confirmation, Go-Live forms).

Please Note: Documentation loaded with the attestation does not alleviate the provider from being requested to produce additional documentation that may be requested during a pre payment or post payment audit. All documentation supporting the information attested by the Provider or Facility should be kept for 6 years.

UPLOADED DOCUMENTS				
	Payment Year	File Name	Description	
View	2	SLR Software Development Lifecycle Processes (002).pdf	Patient Volume	Delete
View	3	Sanchez_OIG 743891_EHR Report.pdf	Purchase Order	Delete
View	3	OutcomeBasedCertiication_CoP_06092020.pdf	Miscellaneous	Delete
View	3	Carol Martin_OIG 745850_EHR Report Review 05222020.pdf	Patient Volume	Delete

Upload a new PDF document: [Browse...](#)

Please select the documentation type:

[Upload](#)

[Previous](#) [Next](#)

The document upload screen allows EPs to submit PDF documents as part of the attestation. Documentation in support of the attestation includes but is not limited to; the Patient Volume Report, CEHRT ID documentation, MU report(s) from their CEHRT, and KHIE onboarding documentation. (Please note: the Patient Volume Report is a required upload in order to move to the next screen.)

Upload PDF documents by following the below steps:

- Select **Browse** to locate a document to upload.
- Select the documentation type from the dropdown.
- Click **Upload**.

When final selections have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen. Selections will not be saved.
- Click **Next** to move on to the next screen. Selections will be saved.

10.4 Attestation Statement Screen

Attestation Statement (Year 3 Attestation)

You are about to submit your attestation for participation in the Kentucky Medicaid EHR Incentive Program.

Please check the box next to each statement below to attest. Participation is required for ONC Direct Review and Participation is Optional for the ONC Surveillance. To complete your attestation, initial, enter your NPI and click the Submit button.

- The information submitted is accurate to the knowledge and belief of the EP.
- The information submitted is accurate and complete for numerators, denominators and exclusions for functional measures applicable to the EP.
- A zero was reported in the denominator of a measure when an EP did not care for any patients in the denominator population during the EHR reporting period.
- The information submitted includes information on all patients to whom the measure applies.
- As a meaningful EHR user, at least 50% of my patient encounters during the EHR reporting period occurred at the practice/location given in my attestation information and is equipped with certified EHR technology.
- The information submitted for Objective 1, Protect Patient Health Information, requires a Security Risk Analysis to be completed within the calendar year of the EHR reporting period.
- The information submitted for eCQM's was generated as output from an identified certified EHR technology.
- The information submitted for eCQM's includes at least one outcome or high priority measure. If there are no outcome or high priority measures relevant to the EP's scope of practice, 6 relevant measures were reported.

<p>1. Participation is Required for ONC Direct Review. The provider must answer question 1 (either 1a or 1a and 1b) -</p> <p>Supporting providers with the performance of CEHRT (SPPC). To engage in activities related to supporting providers with the performance of CEHRT the EP must attest that:</p>	
<input type="checkbox"/>	<p>1a. Acknowledges the requirement to cooperate in good faith with ONC direct review of the EPs health information technology certified under the ONC Health IT Certification Program if a request to assist in ONC direct review is received.</p>
<input type="checkbox"/>	<p>1b. If requested, cooperated in good faith with ONC direct review of EPs health information technology certified under the ONC Health IT Certification Program as authorized by 45 CFR part 170, subpart E, to the extent that such technology meets (or can be used to meet) the definition of CEHRT, including by permitting timely access to such technology and demonstrating its capabilities as implemented and used by the EP in the field.</p>
<p>2. Participation is Optional for ONC Surveillance. The provider may answer question 2 (either 2a or 2a and 2b) -</p>	
<input type="checkbox"/>	<p>2a. Acknowledges the option to cooperate in good faith with ONC-ACB surveillance of his or her health information technology certified under the ONC Health IT Certification Program if a request to assist in ONC-ACB surveillance is received; and</p>
<input type="checkbox"/>	<p>2b. If requested, cooperated in good faith with ONC-ACB surveillance of the EPs health information technology certified under the ONC Health IT Certification Program as authorized by 45 CFR part 170, subpart E, to the extent that such technology meets (or can be used to meet) the definition of CEHRT, including by permitting timely access to such technology and demonstrating capabilities as implemented and used by the EP in the field.</p>
<p>Support for health information exchange and the prevention of information blocking.</p>	
<input type="checkbox"/>	<p>Did not knowingly and willfully take action (such as disable functionality) to limit or restrict the compatibility or interoperability of certified EHR technology.</p>
<input type="checkbox"/>	<p>Implemented technologies, standards, policies, practices and agreements reasonably calculated to ensure, to the greatest extent practicable and permitted by law, that the certified EHR technology was at all relevant times –</p> <p>Connected in accordance with applicable law;</p> <p>Compliant with all standards applicable to the exchange of information, including the standards, implementation specifications and certification criteria adopted at 45 CFR part 170;</p> <p>Implemented in a manner that allowed for timely access by patients to their electronic health information; and</p>

Implemented in a manner that allowed for the timely, secure and trusted bi-directional exchange of structured electronic health information with other health care providers (as defined by 42 U.S.C. 300jj(3)), including unaffiliated providers, and with disparate certified EHR technology and vendors.

Responded in good faith and in a timely manner to requests to retrieve or exchange electronic health information, including from patients, health care providers (as defined by 42 U.S.C. 300jj (3)), and other persons, regardless of the requestor's affiliation or technology vendor.

I understand that I must have, and retain, documentation to support my eligibility for incentive payments and that the Department for Medicaid Services may ask for this documentation. I further understand that the Department for Medicaid Services will pursue repayment in all instances of improper or duplicate payment. I certify I am not receiving Medicaid EHR incentive funds from any other state or commonwealth and have not received a payment from the Kentucky Department for Medicaid Services for this year.

This is to certify that the foregoing information is true, accurate, and complete. I understand the Medicaid EHR incentive payments submitted under this provider number will be from Federal funds, and that any falsification, or concealment of a material fact may be prosecuted under Federal and State laws.

(*)Red asterik indicates a required field.

Initials: *

NPI: *

Note: Once you press the submit button below, you will not be able to change your information.

[Previous](#) [Submit](#)

All boxes must be checked appropriately in order to submit the attestation. Enter initials and NPI to submit the attestation.

When final selections have been made, choose a navigation button at the bottom of the screen.

- Click **Previous** to go back to the previous screen. Selections will not be saved.
- Click **Submit** to save and submit the attestation.

10.5 Accepted Attestation Screen

Attestation Summary Menu (Year 3 Attestation)

Success: Your attestation has been accepted.

All measures and their corresponding calculation have met compliance. Please select the desired measure link below to view the details of your submitted measures.

[Meaningful Use Core Objectives Summary](#)

[Public Health Objectives Summary](#)

[Electronic Clinical Quality Measures Summary - Manual](#)

Once the attestation is accepted, no updates can be made to any data from the attestation.

Click on the summary links to view the measure data that was submitted and accepted for attestation.

10.6 Attestation Not Accepted Screen

Attestation Summary Menu (Year 3 Attestation)

Alert: Your attestation cannot be accepted at this time.

One or more of the MU Core measure calculations did not meet MU minimum standards.
One or more of the Public health measures did not meet MU minimum standards.

Please select the summary of measures link below to view all measures and their corresponding calculation/compliance.

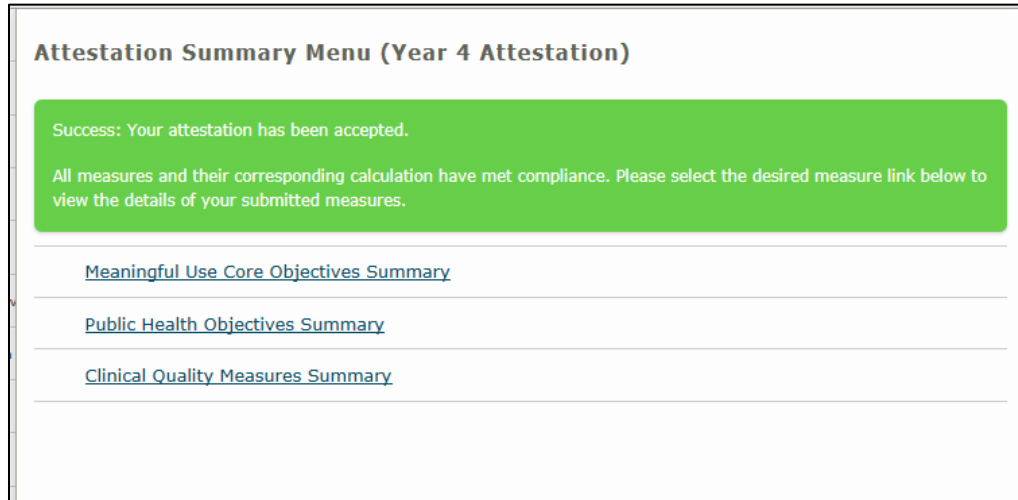
[Meaningful Use Core Objectives Summary](#)

[Public Health Objectives Summary](#)

[Electronic Clinical Quality Measures Summary - Manual](#)

Click on the summary links to view the measure data responses. The summary page will indicate which measures were accepted and which were rejected.

10.7 Post Attestation Summary Screen



After attestation is completed, a statement will appear that the attestation has been accepted.

Click on the summary links to view the measure data that was submitted. The summary page will indicate which measures were accepted.

10.7.1 Objectives Summary

Meaningful Use Core Measure Summary (Year 3 Attestation)			
CORE OBJECTIVES SUMMARY			
Objective	Measure	Entered	Status
Protect electronic protected health information (ePHI) created or maintained by the CEHRT through the implementation of appropriate technical, administrative, and physical safeguards.	Conduct or review a security risk analysis in accordance with the requirements under 45 CFR 164.308(a)(1), including addressing the security (including encryption) of data created or maintained by CEHRT in accordance with requirements under 45 CFR 164.312(a)(2)(iv) and 45 CFR 164.306(d)(3), implement security updates as necessary, and correct identified security deficiencies as part of the provider's risk management process.	Yes	Accepted
Generate and transmit permissible prescriptions electronically.	More than 60% of all permissible prescriptions written by the EP are queried for a drug formulary and transmitted electronically using CEHRT.	100%	Accepted

<p>Implement clinical decision support (CDS) interventions focused on improving performance on high-priority health conditions.</p>	<p>Implement five CDS interventions related to four or more clinical quality measures (CQMs) at a relevant point in patient care for the entire EHR reporting period. Absent four CQMs related to an EP's scope of practice or patient population, the CDS interventions must be related to high-priority health conditions.</p>	<p>Yes</p>	<p>Accepted</p>
<p>Implement clinical decision support (CDS) interventions focused on improving performance on high-priority health conditions.</p>	<p>Enable and implement the functionality for drug-drug and drug-allergy interaction checks for the entire EHR reporting period.</p>	<p>Yes</p>	<p>Accepted</p>
<p>Use CPOE for medication, laboratory, and diagnostic imaging orders directly entered by any licensed healthcare professional, credentialed medical assistant, or a medical staff member credentialed to and performing the equivalent duties of a credentialed medical assistant, who can enter orders into the medical record per state, local, and professional guidelines.</p>	<p>More than 60% of medication orders created by the EP during the EHR reporting period are recorded using computerized provider order entry.</p>	<p>60%</p>	<p>Accepted</p>
<p>Use CPOE for medication, laboratory, and diagnostic imaging orders directly entered by any licensed healthcare professional, credentialed medical assistant, or a medical staff member credentialed to and performing the equivalent duties of a credentialed medical assistant, who can enter orders into the medical record per state, local, and professional guidelines.</p>	<p>More than 60% of laboratory orders created by the EP during the EHR reporting period are recorded using computerized provider order entry.</p>	<p>60%</p>	<p>Accepted</p>

<p>Use CPOE for medication, laboratory, and diagnostic imaging orders directly entered by any licensed healthcare professional, credentialed medical assistant, or a medical staff member credentialed to and performing the equivalent duties of a credentialed medical assistant, who can enter orders into the medical record per state, local, and professional guidelines.</p>	<p>More than 60% of diagnostic imaging orders created by the EP during the EHR reporting period are recorded using computerized provider order entry.</p>	<p>60%</p>	<p>Accepted</p>
<p>The EP provides patients (or patient-authorized representatives) with timely electronic access to their health information and patient-specific education.</p>	<p>For more than 80% of all unique patients seen by the EP: (1) The patient (or the patient-authorized representative) is provided timely access to view online, download, and transmit his or her health information; and (2) The provider ensures the patient's health information is available for the patient (or patient-authorized representative) to access using any application of their choice that is configured to meet the technical specifications of the API in the provider's CEHRT.</p>	<p>Excluded 2</p>	<p>Accepted</p>
<p>The EP provides patients (or patient-authorized representatives) with timely electronic access to their health information and patient-specific education.</p>	<p>The EP must use clinically relevant information from CEHRT to identify patient-specific educational resources and provide electronic access to those materials to more than 35% of unique patients seen by the EP during the EHR reporting period.</p>	<p>Excluded 2</p>	<p>Accepted</p>
<p>Use CEHRT to engage with patients or their authorized representatives about the patient's care.</p>	<p>More than 5% of all unique patients (or their authorized representatives) seen by the EP actively engage with the EHR made accessible by the EP and either: (1) View, download, or transmit to a third party their health information; or (2) Access their health information through an API that can be used by applications chosen by the patient and configured to the API in the EP's CEHRT; or (3) A combination of (1) and (2).</p>	<p>5%</p>	<p>Accepted</p>

<p>Use CEHRT to engage with patients or their authorized representatives about the patient's care.</p>	<p>For more than 5% of all unique patients seen by the EP during the EHR reporting period, a secure message was sent using the electronic messaging function of CEHRT to the patient (or the patient-authorized representative), or in response to a secure message sent by the patient or their authorized representative.</p>	<p>5%</p>	<p>Accepted</p>
<p>Use CEHRT to engage with patients or their authorized representatives about the patient's care.</p>	<p>Patient generated health data or data from a non-clinical setting is incorporated into the CEHRT for more than 5% of all unique patients seen by the EP during the EHR reporting period.</p>	<p>4%</p>	<p>N/A</p>
<p>The EP provides a summary of care record when transitioning or referring their patient to another setting of care, receives or retrieves a summary of care record upon the receipt of a transition or referral or upon the first patient encounter with a new patient, and incorporates summary of care information from other providers into their EHR using the functions of CEHRT.</p>	<p>For more than 50% of transitions of care and referrals, the EP that transitions or refers their patient to another setting of care or provider of care: (1) Creates a summary of care record using CEHRT; and (2) Electronically exchanges the summary of care record.</p>	<p>100%</p>	<p>Accepted</p>
<p>The EP provides a summary of care record when transitioning or referring their patient to another setting of care, receives or retrieves a summary of care record upon the receipt of a transition or referral or upon the first patient encounter with a new patient, and incorporates summary of care information from other providers into their EHR using the functions of CEHRT.</p>	<p>For more than 40% of transitions or referrals received and patient encounters in which the EP has never encountered the patient before, they incorporate into the patient's EHR an electronic summary of care document.</p>	<p>100%</p>	<p>Accepted</p>

<p>The EP provides a summary of care record when transitioning or referring their patient to another setting of care, receives or retrieves a summary of care record upon the receipt of a transition or referral or upon the first patient encounter with a new patient, and incorporates summary of care information from other providers into their EHR using the functions of CEHRT.</p>	<p>For more than 80% of transitions or referrals received and patient encounters in which the EP has never encountered the patient before, they perform a clinical information reconciliation. The EP must implement clinical information reconciliation for the following three clinical information sets: (1) Medication. Review of the patient's medication, including the name, dosage, frequency, and route of each medication. (2) Medication allergy. Review of the patient's known medication allergies. (3) Current Problem list. Review of the patient's current and active diagnoses.</p>	<p>80%</p>	<p>Accepted</p>
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[Return to Menu](#)

10.7.2 Public Health Objectives Summary

Public Health Measures Summary (Year 4 Attestation)			
ObjectiveText	Measure	Entered	Status
<p>The EP is in active engagement with a public health agency to submit electronic public health data from CEHRT except where prohibited and in accordance with applicable law and practice.</p> <p>We further specify that providers must use the functions and standards as defined for CEHRT at § 495.4 where applicable; however, as noted for measure 3, providers may use functions beyond those established in CEHRT in accordance with state and local law.</p>	<p>The EP is in active engagement with a public health agency to submit immunization data.</p>	<p>Option 3</p>	<p>Accepted</p>
<p>The EP is in active engagement with a public health agency to submit electronic public health data from CEHRT except where prohibited and in accordance with applicable law and practice.</p> <p>We further specify that providers must use the functions and standards as defined for CEHRT at § 495.4 where applicable; however, as noted for measure 3, providers may use functions beyond those established in CEHRT in accordance with state and local law.</p>	<p>The EP is in active engagement to submit data to a specialized registry.</p>	<p>Option 2 - KY Cancer Registry</p>	<p>Accepted</p>

[Return to Menu](#)

10.7.3 Electronic Clinical Quality Measures Summary (Electronically Reported)

Electronic Clinical Quality Measures (Year 2 Attestation)

Electronic Clinical Quality Measures Summary Report

<u>Date Received</u>	<u>Report NPI</u>	<u>Reporting Period Start</u>	<u>Reporting Period End</u>
7/1/2020 1:01:24 PM	1851404065	3/1/2020 12:00:00 AM	6/22/2020 12:00:00 AM
<u>Document Type</u>	<u>eCQM Status:</u>	<u>Evaluated Date:</u>	
QRDA III	Accepted	7/1/2020 1:10:13 PM	
<u>File Name:</u>			
QRDA_XML2.xml, CPCSampleQRDA-III.xml			

QRDA_XML2_TESTFILE3.XML

Domain	# of Measures
Communication and Care Coordination	0
Community/Population Health	2
Effective Clinical Care	1
Efficiency and Cost Reduction	0
Patient Safety	1
Person and Caregiver-Centered Experience and Outcomes	0

CPCSAMPLEQRDA-III.XML

Domain	# of Measures
Communication and Care Coordination	1
Community/Population Health	4
Effective Clinical Care	9
Efficiency and Cost Reduction	0
Patient Safety	2
Person and Caregiver-Centered Experience and Outcomes	0

Total Required	6
Total Met	17
Final eCQM Result:	Submission Accepted.
<div style="background-color: #4CAF50; color: white; padding: 5px 15px; border-radius: 5px; display: inline-block;">Return To Menu</div>	

10.7.4 Electronic Clinical Quality Measures Summary (Manually Reported)

**Meaningful Use Electronic Clinical Quality Measures Summary
(Year 3 Attestation)**

Summary of Electronic Clinical Quality Measures		
PERSON AND CAREGIVER-CENTERED EXPERIENCE AND OUTCOMES		
Title	Description	Status
Oncology: Medical and Radiation - Pain Intensity Quantified	Percentage of patient visits, regardless of patient age, with a diagnosis of cancer currently receiving chemotherapy or radiation therapy in which pain intensity is quantified.	Accepted
Functional Status Assessment for Total Hip Replacement	Percentage of patients 18 years of age and older who received an elective primary total hip arthroplasty (THA) and completed a functional status assessment within 90 days prior to the surgery and in the 270-365 days after the surgery.	Accepted
Functional Status Assessment for Total Knee Replacement	Percentage of patients 18 years of age and older who received an elective primary total knee arthroplasty (TKA) and completed a functional status assessment within 90 days prior to the surgery and in the 270-365 days after the surgery.	Accepted
Functional Status Assessments for Congestive Heart Failure	Percentage of patients 18 years of age and older with congestive heart failure who completed initial and follow-up patient-reported functional status assessments.	Accepted
Urinary Symptom Score Change 6-12 Months After Diagnosis of Benign Prostatic Hyperplasia	Percentage of patients with an office visit within the measurement period and with a new diagnosis of clinically significant Benign Prostatic Hyperplasia who have International Prostate Symptoms Score (IPSS) or American Urological Association (AUA) Symptom Index (SI) documented at time of diagnosis and again 6-12 months later with an improvement of 3 points.	Accepted

PATIENT SAFETY

Title	Description	Status
Use of High-Risk Medications in the Older Adults	Percentage of patients 65 years of age and older who were ordered at least two of the same high-risk medications.	Accepted
Falls: Screening for Future Fall Risk	Percentage of patients 65 years of age and older who were screened for future fall risk during the measurement period.	Accepted
Documentation of Current Medications in the Medical Record	Percentage of visits for patients aged 18 years and older for which the eligible professional or eligible clinician attests to documenting a list of current medications using all immediate resources available on the date of the encounter.	Accepted
Child and Adolescent Major Depressive Disorder (MDD): Suicide Risk Assessment	Percentage of patient visits for those patients aged 6 through 17 years with a diagnosis of major depressive disorder with an assessment for suicide risk.	Accepted

COMMUNICATION AND CARE COORDINATION

Title	Description	Status
Diabetic Retinopathy: Communication with the Physician Managing Ongoing Diabetes Care	Percentage of patients aged 18 years and older with a diagnosis of diabetic retinopathy who had a dilated macular or fundus exam performed with documented communication to the physician who manages the ongoing care of the patient with diabetes mellitus regarding the findings of the macular or fundus exam at least once within 12 months.	Accepted
Closing the Referral Loop: Receipt of Specialist Report	Percentage of patients with referrals, regardless of age, for which the referring provider receives a report from the provider to whom the patient was referred.	Accepted

COMMUNITY/POPULATION HEALTH

Title	Description	Status
Weight Assessment and Counseling for Nutrition and Physical Activity for Children and Adolescents	Percentage of patients 3-17 years of age who had an outpatient visit with a Primary Care Physician (PCP) or Obstetrician/Gynecologist (OB/GYN) and who had evidence of the following during the measurement period. Three rates are reported. - Percentage of patients with height, weight, and body mass index (BMI) percentile documentation. - Percentage of patients with counseling for nutrition. - Percentage of patients with counseling for physical activity.	Accepted
Preventive Care and Screening: Tobacco Use: Screening and Cessation Intervention	Percentage of patients aged 18 years and older who were screened for tobacco use one or more times within 12 months AND who received tobacco cessation intervention if identified as a tobacco user. Three rates are reported. a. Percentage of patients aged 18 years and older who were screened for tobacco use one or more times within 12 months. b. Percentage of patients aged 18 years and older who were identified as a tobacco user who received tobacco cessation intervention. c. Percentage of patients aged 18 years and older who were screened for tobacco use one or more times within 12 months AND who received tobacco cessation intervention if identified as a tobacco user.	Accepted

Chlamydia Screening for Women	Percentage of women 16-24 years of age who were identified as sexually active and who had at least one test for chlamydia during the measurement period.	Accepted
Childhood Immunization Status	Percentage of children 2 years of age who had four diphtheria, tetanus and acellular pertussis (DTaP); three polio (IPV), one measles, mumps and rubella (MMR); three or four H influenza type B (Hib); three hepatitis B (Hep B); one chicken pox (VZV); four pneumococcal conjugate (PCV); one hepatitis A (Hep A); two or three rotavirus (RV); and two influenza (flu) vaccines by their second birthday.	Accepted
Preventive Care and Screening: Influenza Immunization	Percentage of patients aged 6 months and older seen for a visit between October 1 and March 31 who received an influenza immunization OR who reported previous receipt of an influenza immunization.	Accepted
Pneumococcal Vaccination Status for Older Adults	Percentage of patients 65 years of age and older who have ever received a pneumococcal vaccine.	Accepted
Preventive Care and Screening: Screening for Depression and Follow-Up Plan	Percentage of patients aged 12 years and older screened for depression on the date of the encounter or 14 days prior to the date of the encounter using an age-appropriate standardized depression screening tool AND if positive, a follow-up plan is documented on the date of the eligible encounter.	Accepted
Preventive Care and Screening: Body Mass Index (BMI) Screening and Follow-Up Plan	Percentage of patients aged 18 years and older with a BMI documented during the current encounter or within the previous twelve months AND who had a follow-up plan documented if most recent BMI was outside of normal parameters.	Accepted
Preventive Care and Screening: Screening for High Blood Pressure and Follow-Up Documented	Percentage of patient visits for patients aged 18 years and older seen during the measurement period who were screened for high blood pressure AND a recommended follow-up plan is documented, as indicated, if blood pressure is pre-hypertensive or hypertensive.	Accepted
Children Who Have Dental Decay or Cavities	Percentage of children, 6 months - 20 years of age, who have had tooth decay or cavities during the measurement period.	Accepted
HIV Screening	Percentage of patients aged 15-65 at the start of the measurement period who were between 15-65 years old when tested for HIV.	Accepted

EFFICIENCY AND COST REDUCTION

Title	Description	Status
Appropriate Testing for Pharyngitis	The percentage of episodes for patients 3 years and older with a diagnosis of pharyngitis that resulted in an antibiotic dispensing event and a group A streptococcus (strep) test.	Accepted
Appropriate Treatment for Upper Respiratory Infection (URI)	Percentage of episodes for patients 3 months of age and older with a diagnosis of upper respiratory infection (URI) that did not result in an antibiotic dispensing event.	Accepted

Prostate Cancer: Avoidance of Overuse of Bone Scan for Staging Low Risk Prostate Cancer Patients	Percentage of patients, regardless of age, with a diagnosis of prostate cancer at low (or very low) risk of recurrence receiving interstitial prostate brachytherapy, OR external beam radiotherapy to the prostate, OR radical prostatectomy who did not have a bone scan performed at any time since diagnosis of prostate cancer.	Accepted
Appropriate Use of DXA Scans in Women Under 65 Years Who Do Not Meet the Risk Factor Profile for Osteoporotic Fracture	Percentage of female patients 50 to 64 years of age without select risk factors for osteoporotic fracture who received an order for a dual-energy x-ray absorptiometry (DXA) scan during the measurement period.	Accepted

EFFECTIVE CLINICAL CARE

Title	Description	Status
Initiation and Engagement of Alcohol and Other Drug Dependence Treatment	Percentage of patients 13 years of age and older with a new episode of alcohol or other drug abuse or (AOD) dependence who received the following. Two rates are reported. a. Percentage of patients who initiated treatment including either an intervention or medication for the treatment of AOD abuse or dependence within 14 days of the diagnosis. b. Percentage of patients who engaged in ongoing treatment including two additional interventions or a medication for the treatment of AOD abuse or dependence within 34 days of the initiation visit. For patients who initiated treatment with a medication, at least one of the two engagement events must be a treatment intervention.	Accepted
Controlling High Blood Pressure	Percentage of patients 18-85 years of age who had a diagnosis of hypertension overlapping the measurement period or the year prior to the measurement period, and whose most recent blood pressure was adequately controlled (< 140/90mmHg) during the measurement period.	Accepted
Cervical Cancer Screening	Percentage of women 21-64 years of age who were screened for cervical cancer using either of the following criteria: * Women age 21-64 who had cervical cytology performed within the last 3 years. * Women age 30-64 who had cervical human papillomavirus (HPV) testing performed within the last 5 years.	Accepted
Colorectal Cancer Screening	Percentage of adults 50-75 years of age who had appropriate screening for colorectal cancer.	Accepted
Diabetes: Eye Exam	Percentage of patients 18-75 years of age with diabetes and an active diagnosis of retinopathy overlapping the measurement period who had a retinal or dilated eye exam by an eye care professional during the measurement period or diabetics with no diagnosis of retinopathy overlapping the measurement period who had a retinal or dilated eye exam by an eye care professional during the measurement period or in the 12 months prior to the measurement period.	Accepted

Diabetes: Hemoglobin A1c (HbA1c) Poor Control (> 9%)	Percentage of patients 18-75 years of age with diabetes who had hemoglobin A1c > 9.0% during the measurement period.	Accepted
Diabetes: Medical Attention for Nephropathy	The percentage of patients 18-75 years of age with diabetes who had a nephropathy screening test or evidence of nephropathy during the measurement period.	Accepted
Coronary Artery Disease (CAD): Beta-Blocker Therapy-Prior Myocardial Infarction (MI) or Left Ventricular Systolic Dysfunction (LVEF < 40%)	Percentage of patients aged 18 years and older with a diagnosis of coronary artery disease seen within a 12-month period who also have a prior MI or a current or prior LVEF < 40% who were prescribed beta-blocker therapy.	Accepted
Heart Failure (HF): ACE Inhibitor or ARB or ARNI Therapy for Left Ventricular Systolic Dysfunction (LVSD)	Percentage of patients aged 18 years and older with a diagnosis of heart failure (HF) with a current or prior left ventricular ejection fraction (LVEF) < 40% who were prescribed ACE inhibitor or ARB or ARNI therapy either within a 12-month period when seen in the outpatient setting OR at each hospital discharge.	Accepted
Heart Failure (HF): Beta-Blocker Therapy for Left Ventricular Systolic Dysfunction (LVSD)	Percentage of patients aged 18 years and older with a diagnosis of heart failure (HF) with a current or prior left ventricular ejection fraction (LVEF) < 40% who were prescribed beta-blocker therapy either within a 12-month period when seen in the outpatient setting OR at each hospital discharge.	Accepted
Primary Open-Angle Glaucoma (POAG): Optic Nerve Evaluation	Percentage of patients aged 18 years and older with a diagnosis of primary open-angle glaucoma (POAG) who have an optic nerve head evaluation during one or more office visits within 12 months.	Accepted
Adult Major Depressive Disorder (MDD): Suicide Risk Assessment	All patient visits during which a new diagnosis of MDD or new diagnosis of recurrent MDD was identified for patients aged 18 years and older with a suicide risk assessment completed during the visit.	Accepted
Anti-depressant Medication Management	Percentage of patients 18 years of age and older who were treated with antidepressant medication, had a diagnosis of major depression, and who remained on an antidepressant medication treatment. Two rates are reported. a. Percentage of patients who remained on an antidepressant medication for at least 84 days (12 weeks). b. Percentage of patients who remained on an antidepressant medication for at least 180 days (6 months).	Accepted
Follow-Up Care for Children Prescribed ADHD Medication (ADD)	Percentage of children 6-12 years of age and newly dispensed a medication for attention-deficit/hyperactivity disorder (ADHD) who had appropriate follow-up care. Two rates are reported. a. Percentage of children who had one follow-up visit with a practitioner with prescribing authority during the 30-Day Initiation Phase. b. Percentage of children who remained on ADHD medication for at least 210 days and who, in addition to the visit in the Initiation Phase, had at least two additional follow-up visits with a practitioner within 270 days (9 months) after the Initiation Phase ended.	Accepted

Cataracts: 20/40 or Better Visual Acuity within 90 Days Following Cataract Surgery	Percentage of cataract surgeries for patients aged 18 and older with a diagnosis of uncomplicated cataract and no significant ocular conditions impacting the visual outcome of surgery and had best-corrected visual acuity of 20/40 or better (distance or near) achieved in the operative eye within 90 days following the cataract surgery.	Accepted
Depression Remission at Twelve Months	The percentage of adolescent patients 12 to 17 years of age and adult patients 18 years of age or older with major depression or dysthymia who reached remission 12 months (+/-60 days) after an index event.	Accepted
Breast Cancer Screening	Percentage of women 50-74 years of age who had a mammogram to screen for breast cancer in the 27 months prior to the end of the Measurement Period.	Accepted
Dementia: Cognitive Assessment	Percentage of patients, regardless of age, with a diagnosis of dementia for whom an assessment of cognition is performed and the results reviewed at least once within a 12-month period.	Accepted
Primary Caries Prevention Intervention as Offered by Primary Care Providers, including Dentists	Percentage of children, 6 months - 20 years of age, who received a fluoride varnish application during the measurement period.	Accepted
Statin Therapy for the Prevention and Treatment of Cardiovascular Disease	Percentage of the following patients - all considered at high risk of cardiovascular events - who were prescribed or were on statin therapy during the measurement period: *Adults aged >= 21 years who were previously diagnosed with or currently have an active diagnosis of clinical atherosclerotic cardiovascular disease (ASCVD); OR *Adults aged >= 21 years who have ever had a fasting or direct low-density lipoprotein cholesterol (LDL-C) level >= 190 mg/dL or were previously diagnosed with or currently have an active diagnosis of familial or pure hypercholesterolemia; OR *Adults aged 40-75 years with a diagnosis of diabetes with a fasting or direct LDL-C level of 70-189 mg/dL.	Accepted
Bone density evaluation for patients with prostate cancer and receiving androgen deprivation therapy	Patients determined as having prostate cancer who are currently starting or undergoing androgen deprivation therapy (ADT), for an anticipated period of 12 months or greater and who receive an initial bone density evaluation. The bone density evaluation must be prior to the start of ADT or within 3 months of the start of ADT.	Accepted

[Return to Menu](#)

10.8 Next Steps

Thank you for participating in the Kentucky Medicaid EHR Incentive Program (Promoting Interoperability). The attestation will be reviewed as quickly as possible. Please be on the lookout for emails requesting additional information. A delayed response will delay the review process and thus will also delay receipt of your incentive payment.

A certificate of completion will be emailed once payment is processed for your sixth year of participation. You are no longer required to submit an attestation to the Kentucky Medicaid

EHR Incentive Program (Promoting Interoperability) however, providers are encouraged to participate in other programs available.

The Quality Payment Program (QPP) helps providers focus on care quality and making patients healthier. QPP also ends the Sustainable Growth Rate formula and gives the provider new tools, models, and resources to help give their patients the best possible care. Providers may select to participate in the Advanced Alternative Payment models (APMs) or the Merit-based Incentive Payment System (MIPS). If you participate in an Advanced APM, through Medicare Part B you may earn an incentive payment for participating in an innovative payment model. If you participate in MIPS, you will earn a performance-based payment adjustment. To check your participation status and for more information, providers can visit the [website](#).

CMS released the 2018 Quality Payment Program (QPP) Experience Report to provide insights into participation during the 2018 performance year. The report includes data regarding participation and performance in the MIPS and APMs tracks of QPP during the 2018 performance year. Key 2018 findings include:

- 98% of MIPS eligible clinicians participated in the program and avoided a negative payment adjustment, a one-year increase of 3 percentage points.
- 84% of clinicians earned an “exceptional performance” designation by earning 70 points or more.
- 356,353 MIPS eligible clinicians participated in MIPS through a MIPS APM, a 15,000-clinician increase from the 2017 performance year.
- The number of clinicians achieving Qualifying APM Participant (QP) status nearly doubled in one year, from 99,076 to 183,306 clinicians. This, along with the increase in MIPS APM participation, indicates a desire from clinicians and practices to transition toward value-based arrangements.
- 84% of small practices earned a positive payment adjustment, up 10 percentage points from the year prior.
- The percentage of rural practices earning a positive payment adjustment increased from 93% in 2017 to 97% in 2018.

For more information contact the Quality Payment Program at 1-866-288-8292 or by email at: gpp@cms.hhs.gov.