Quality ID \#249: Barrett's Esophagus

- National Quality Strategy Domain: Effective Clinical Care
- Meaningful Measure Area: Transfer of Health Information and Interoperability


## 2022 COLLECTION TYPE:

MIPS CLINICAL QUALITY MEASURES (CQMS)

## MEASURE TYPE:

Process

## DESCRIPTION:

Percentage of esophageal biopsy reports that document the presence of Barrett's mucosa that also include a statement about dysplasia

## INSTRUCTIONS:

This measure is to be submitted each time a patient's esophageal surgical pathology report demonstrates Barrett's Esophagus; however, only one quality data code (QDC) per date of service for a patient is required. This measure may be submitted by Merit-based Incentive Payment System (MIPS) eligible clinicians who perform the quality actions described in the measure based on the services provided and the measure-specific denominator coding.

## Measure Submission Type:

Measure data may be submitted by individual MIPS eligible clinicians, groups, or third party intermediaries. The listed denominatorcriteriaareusedtoidentifytheintended patientpopulation. Thenumeratoroptionsincludedinthis specification are used to submit the quality actions as allowed by the measure. The quality data codes listed do not need tobesubmitted byMIPSeligibleclinicians, groups, orthird party intermediariesthatutilizethismodalityfor submissions; however, these codes may be submitted for those third party intermediaries that utilize Medicare Part B claims data. For more information regarding Application Programming Interface (API), please refer to the Quality Payment Program (QPP) website.

## DENOMINATOR:

All surgical pathology esophageal biopsy reports for Barrett's Esophagus
Denominator Criteria (Eligible Cases):
Diagnosis for Barrett's Esophagus (ICD-10-CM): K22.70, K22.710, K22.711, K22.719
AND
Patient procedure during the performance period (CPT): 88305
WITHOUT
Telehealth Modifier: GQ, GT, 95, POS 02
AND NOT
DENOMINATOR EXCLUSION:
Specimen site other than anatomic location of esophagus: G8797

## NUMERATOR:

Esophageal biopsyreport documents the presence of Barrett's mucosa and includes astatementaboutdysplasia
Numerator Options:
Performance Met: Esophageal biopsy reports with the histological finding of Barrett's mucosa that contains a statement about dysplasia (present, absent, or indefinite and if present, contains appropriate grading)(3126F)

## OR

## DenominatorException:

Documentation of medical reason(s) for not submitting the histological finding of Barrett's mucosa (e.g., malignant neoplasm or absence of intestinal metaplasia) (3126F with

Performance Not Met:

Pathology report with the histological finding of Barrett's mucosa that does not contain a statement about dysplasia (present, absent, orindefinite, and if present, contains appropriate grading), reason not otherwise specified (3126F with 8P)

## RATIONALE:

Endoscopy is the technique of choice used to identify suspected Barrett's esophagus and to diagnose complications of GERD. Biopsy must beadded to confirm the presence of Barrett'sepithelium and to evaluatefor dysplasia(ACG, 2016; AGA, 2011).

There is a rapidly rising incidence of adenocarcinoma of the esophagus in the United States. A diagnosis of Barrett's esophagus increases a patient's risk for esophageal adenocarcinoma by 30 to 125 times that of people without Barrett's esophagus (although this risk is still small $0.4 \%$ to $0.5 \%$ per year)(Conteduca et al 2012, Intl J Onc). Esophageal adenocarcinoma is often not curable, partly because the disease is frequently discovered atalate stage and because treatments are not effective.A diagnosis of Barrett'sesophaguscould allow for appropriate screening of at risk patients as recommended by the American College of Gastroenterology.

Standard endoscopy with biopsy currently is the most reliable means of establishing a diagnosis of Barrett's esophagus. The definitive diagnosis of Barrett's esophagus requires a pathologist's review of an esophageal biopsy. Dysplasia is the first step in the neoplastic process, and information about dysplasia is crucial for clinical decision-making directing therapy. The presence and grade of dysplasia cannot be determined by routine endoscopy, and pathologist's review of a biopsy is essential for recognition of dysplasia, especially given that there are no recommended biomarkers for Barrett's esophagus. Endoscopic surveillance detects curable neoplasia in patients with Barrett's esophagus.

## CLINICAL RECOMMENDATION STATEMENTS:

The diagnosis of Barrett'sesophagus requires systematicbiopsy of the abnormal-appearing esophageal mucosato document intestinal metaplasia and to detect dysplasia (ACG, 2016).

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## 2022 Clinical Quality Measure Flow for Quality ID \#249: Barrett's Esophagus

Disclaimer: Refer to the measure specification for specific coding and instructions to submit this measure.

*See the posted measure specification for specific coding and instructions to submit this measure.

## 2022 Clinical Quality Measure Flow Narrative for Quality ID \#249: Barrett's Esophagus

Disclaimer: Refer to the measure specification for specific coding and instructions to submit this measure.

1. Start with Denominator
2. Check Diagnosis for Barrett's Esophagus as Listed in the Denominator*:
a. If Diagnosis for Barrett's Esophagus as Listed in the Denominator* equals No, do not include in Eligible Population/Denominator. Stop processing.
b. If Diagnosis for Barrett's Esophagus as Listed in the Denominator* equals Yes, proceed to check Patient procedure during the performance period as listed in Denominator*.
3. Check Patient procedure during the performance period as listed in Denominator*:
a. If Patient procedure during the performance period as listed in Denominator* equals No, do not include in Eligible Population/Denominator. Stop processing.
b. If Patient procedure during the performance period as listed in Denominator* equals Yes, proceed to check Telehealth Modifier.
4. Check Telehealth Modifier:
a. If Telehealth Modifier equals Yes, do not include in Eligible Population/Denominator. Stop processing.
b. If Telehealth Modifier equals No, proceed to check Specimen site other than anatomic location of esophagus.
5. Check Specimen site other than anatomic location of esophagus:
a. If Specimen site other than anatomic location of esophagus equals Yes, do not include in Eligible Population/Denominator. Stop processing.
b. If Specimen site other than anatomic location of esophagus equals No, include in Eligible Population/Denominator.
6. Denominator Population:

- Denominator Population is all Eligible Procedures in the Denominator. Denominator is represented as Denominator in the Sample Calculation listed at the end of this document. Letter d equals 80 procedures in the Sample Calculation.

7. Start Numerator
8. Check Esophageal biopsy reports with the histological finding of Barrett's mucosa that contains a statement about dysplasia:
a. If Esophageal biopsy reports with the histological finding of Barrett's mucosa that contains a statement about dysplasia equals Yes, Include in Data Completeness Met and Performance Met.

- Data Completeness Met and Performance Met letter is represented in the Data Completeness and Performance Rate in the Sample Calculation listed at the end of this document. Letter a equals 40 procedures in Sample Calculation.
b. If Esophageal biopsy reports with the histological finding of Barrett's mucosa that contains a statement
about dysplasia equals No, proceed to check Documentation of medical reason(s) for not submitting the histological finding of Barrett's mucosa.

9. Check Documentation of medical reason(s) for not submitting the histological finding of Barrett's mucosa:
a. If Documentation of medical reason(s) for not submitting the histological finding of Barrett's mucosa equals Yes, include in Data Completeness Met and Denominator Exception.

- Data Completeness Met and Denominator Exception letter is represented in the Data Completeness and Performance Rate in the Sample Calculation listed at the end of this document. Letter b equals 10 procedures in the Sample Calculation.
b. If Documentation of medical reason(s) for not submitting the histological finding of Barrett's mucosa equals No, proceed to check Pathology report with the histological finding of Barrett's mucosa that does not contain a statement about dysplasia, reason not otherwise specified.

10. Check Pathology report with the histological finding of Barrett's mucosa that does not contain a statement about dysplasia, reason not otherwise specified:
a. If Pathology report with the histological finding of Barrett's mucosa that does not contain a statement about dysplasia, reason not otherwise specified equals Yes, include in the Data Completeness Met and Performance Not Met.

- Data Completeness Met and Performance Not Met letter is represented in the Data Completeness in the Sample Calculation listed at the end of this document. Letter c equals 20 procedures in the Sample Calculation.
b. If Pathology report with the histological finding of Barrett's mucosa that does not contain a statement about dysplasia, reason not otherwise specified equals No, proceed to check Data Completeness Not Met.

11. Check Data Completeness Not Met:
a. If Data Completeness Not Met, the Quality Data Code or equivalent was not submitted. 10 procedures have been subtracted from Data Completeness Numerator in the Sample Calculation.

## Sample Calculations

Data Completeness equals Performance Met (a equals 40 procedures) plus Denominator Exception (b equals 10 procedures) plus Performance Not Met (c equals 20 procedures) divided by Eligible Population/Denominator (d equals 80 procedures). All equals 70 procedures divided by 80 procedures. All equals 87.50 percent.

Performance Rate equals Performance Met (a equals 40 procedures) divided by Data Completeness Numerator (70 procedures) minus Denominator Exception (b equals 10 procedures). All equals 40 procedures divided by 60 procedures. All equals 66.67 percent.
*See the posted measure specification for specific coding and instructions to submit this measure.
NOTE: Submission Frequency: Procedure
The measure diagrams were developed by CMS as a supplemental resource to be used in conjunction with the measure specifications. They should not be used alone or as a substitution for the measure specification.

