Quality ID #429: Pelvic Organ Prolapse: Preoperative Screening for Uterine Malignancy

- National Quality Strategy Domain: Patient Safety
- Meaningful Measure Area: Preventable Healthcare Harm

2019 COLLECTION TYPE:

MEDICARE PART B CLAIMS

MEASURE TYPE:

Process - High Priority

DESCRIPTION:

Percentage of patients who are screened for uterine malignancy prior to vaginal closure or obliterative surgery for pelvic organ prolapse

INSTRUCTIONS:

This measure is to be submitted <u>each time</u> a prolapse organ repair surgery is performed during the performance period. There is no diagnosis associated with this measure. 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 using Medicare Part B claims. The listed denominator criteria are used to identify the intended patient population. The numerator quality-data codes included in this specification are used to submit the quality actions allowed by the measure on the claim form(s). All measure-specific coding should be submitted on the claim(s) representing the denominator eligible encounter and selected numerator option.

DENOMINATOR:

All patients undergoing surgery for pelvic organ prolapse involving vaginal closure/obliterative procedure

Denominator Criteria (Eligible Cases):

All patients, regardless of age

AND

Patient procedure during the performance period (CPT): 57106, 57110, 57120

NUMERATOR:

Number of patients screened for uterine malignancy or those that had an ultrasound and/or endometrial sampling of any kind

Numerator Quality-Data Coding Options:

Patient is not eligible for this measure because patient has had hysterectomy

Denominator Exclusion: G9774: Patients who have had a hysterectomy

OR

Documentation of Screening for Uterine Malignancy

Performance Met: G9618: Documentation of screening for uterine malignancy, or

those that had an ultrasound and/or endometrial

sampling of any kind

OR

Screening for Uterine Malignancy not Documented, Reason not Given

Performance Not Met: G9620: Patient not screened for uterine malignancy, or those that

have not had an ultrasound and/or endometrial sampling

of any kind, reason not given

RATIONALE:

This measure will promote screening of patients at risk for a uterine malignancy prior to obliterative vaginal surgery. The incidence of endometrial cancer found unsuspectingly in patients with POP ranges from 0.3-3.2%. In a review of all surgical pathology reports for patients undergoing a hysterectomy for pelvic organ prolapse, 644 women were evaluated and 2 were diagnosed with endometrial cancer (0.3%). Ensuring that providers ask about possible symptoms that may hint at the need for further evaluation would increase the quality of care provided to these patients.

CLINICAL RECOMMENDATION STATEMENTS:

This measure will help ensure that patients who do have a uterine malignancy are diagnosed prior to obliterative procedure and can be referred to a gynecologic oncologist for appropriate treatment for the malignancy. The incidence of endometrial cancer found unsuspectingly in patients with POP ranges from 0.3-3.2%. In a review of all surgical pathology reports for patients undergoing a hysterectomy for pelvic organ prolapse, 644 women were evaluated and 2 were diagnosed with endometrial cancer (0.3%). Ensuring that providers ask about possible symptoms that may hint at the need for further evaluation would increase the quality of care provided to these patients.

COPYRIGHT:

These performance measures were developed and are owned by the American Urogynecologic Society ("AUGS"). These performance measures are not clinical guidelines and do not establish a standard of medical care. AUGS makes no representations, warranties, or endorsement about the quality of any organization or physician that uses or reports performance measures and AUGS has no liability to anyone who relies on such measures. AUGS holds a copyright in this measure and can rescind or alter this measure at any time. Users of the measure shall not have the right to alter, enhance, or otherwise modify the measure and shall not disassemble, recompile, or reverse engineer the source code or object code relating to the measure. Anyone desiring to use or reproduce the measure without modification for a noncommercial purpose may do so without obtaining any approval from AUGS. All commercial uses must be approved by AUGS and are subject to a license at the discretion of AUGS. Use by health care providers in connection with their own practices is not commercial use. A "commercial use" refers to any sale, license, or distribution of a measure for commercial gain, or incorporation of a measure into any product or service that is sold, licensed, or distributed for commercial gain, even if there is no actual charge for inclusion of the measure.

Performance measures developed by AUGS for CMS may look different from the measures solely created and owned by AUGS.

THE MEASURES AND SPECIFICATIONS ARE PROVIDED "AS IS" WITHOUT WARRANTY OF ANY KIND.

Limited proprietary coding from Current Procedural Terminology (CPT®) is contained in the measure specifications. Users of this code set should obtain all necessary licenses. AUGS disclaims all liability for use or accuracy of any Current Procedural Terminology (CPT®) or other coding contained in the specifications.

Physician Performance Measures (Measures) and related data specifications developed by AUGS are intended to facilitate quality improvement activities by physicians. These Measures are intended to assist physicians in enhancing quality of care. They are designed for use by any physician who manages the care of a patient for a specific condition or for diagnosis or prevention. AUGS encourages use of this Measure by other health care professionals, where appropriate.

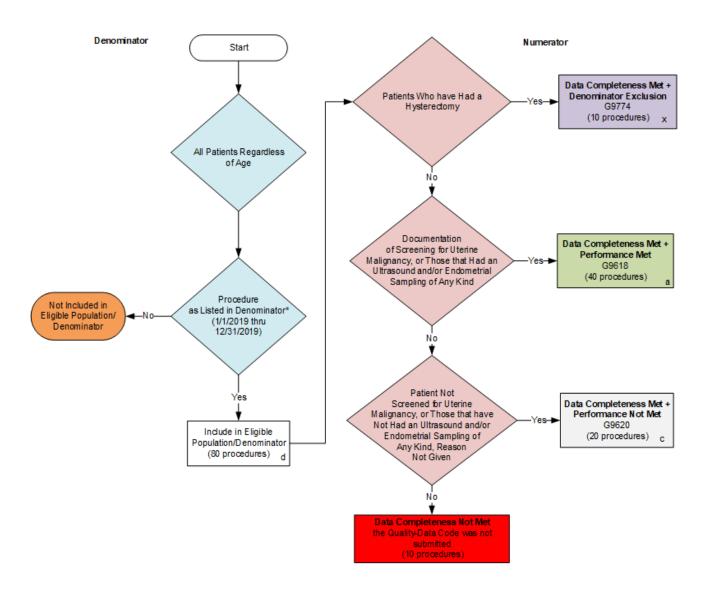
Measures are subject to review and may be revised or rescinded at any time by AUGS. They may not be altered without the prior written approval from AUGS. Measures developed by AUGS, while copyrighted, can be reproduced and distributed, without modification, for noncommercial purposes, e.g., use by health care providers in connection with their practices. Commercial use of the Measures is not permitted absent a license agreement between the user and AUGS. Commercial use is defined as the sale, license, or distribution of the Measures for commercial gain, or incorporation of the Measures into a product or service that is sold, licensed or distributed for commercial gain.

AUGS is not responsible for any harm to any party resulting from the use of these Measures.

Copyright © by the American Urogynecologic Society; 1100 Wayne Ave Suite 825 Silver Spring MD 20910. All Rights Reserved.

CPT® contained in the Measures specifications is copyright 2004-2018 American Medical Association. CPT® is a registered trademark of the American Medical Association.

2019 Medicare Part B Claims Flow for Quality ID #429: Pelvic Organ Prolapse: Preoperative Screening for Uterine Malignancy



SAMPLE CALCULATIONS: Data Completeness= Denominator Exclusion (x=10 procedures) + Performance Met (a=40 procedures) + Performance Not Met (c=20 procedures) = 70 procedures = 87.50% Eligible Population / Denominator (d=80 procedures) = 80 procedures Performance Rate= Performance Met (a=40 procedures) Data Completeness Numerator (70 procedures) - Denominator Exclusion (x=10 procedures) = 60.67% 60 procedures

*See the posted Measure Specification for specific coding and instructions to submit this measure. NOTE: Submission Frequency: Procedure

CPT only copyright 2018 American Medical Association. All rights reserved. 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.

2019 Medicare Part B Claims Flow Narrative for Quality ID #429: Pelvic Organ Prolapse: Preoperative Screening for Uterine Malignancy

Please refer to the specific section of the specification to identify the denominator and numerator information for use in submitting this Individual Specification.

- 1. Start with Denominator
- 2. All Patients Regardless of Age
- 3. Check Procedure Performed:
 - a. If Procedure as Listed in the Denominator equals No, do not include in Eligible Population. Stop Processing.
 - b. If Procedure as Listed in the Denominator equals Yes, include in Eligible Population.
- 4. Denominator Population:
 - a. 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.
- 5. Start Numerator
- 6. Check Patients Who have Had a Hysterectomy:
 - a. If Patients Who have Had a Hysterectomy equals Yes, include in Data Completeness Met and Denominator Exclusion.
 - b. Data Completeness Met and Denominator Exclusion letter is represented in the Data Completeness and Performance Rate in the Sample Calculation listed at the end of this document. Letter x equals 10 procedures in the Sample Calculation.
 - c. If Patients Who have Had a Hysterectomy equals No, proceed to check Documentation of Screening for Uterine Malignancy, or Those that Had an Ultrasound and/or Endometrial Sampling of Any Kind.
- 7. Check Documentation of Screening for Uterine Malignancy, or Those that Had an Ultrasound and/or Endometrial Sampling of Any Kind:
 - a. If Documentation of Screening for Uterine Malignancy, or Those that Had an Ultrasound and/or Endometrial Sampling of Any Kind equals Yes, include in Data Completeness Met and Performance Met.
 - b. 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 the Sample Calculation.
 - c. If Documentation of Screening for Uterine Malignancy, or Those that Had an Ultrasound and/or Endometrial Sampling of Any Kind equals No, proceed to check Patient Not Screened for Uterine Malignancy, or Those that have Not Had an Ultrasound and/or Endometrial Sampling of Any Kind, Reason Not Given.
- 8. Check Patient Not Screened for Uterine Malignancy, or Those that have Not Had an Ultrasound and/or Endometrial Sampling of Any Kind, Reason Not Given:

- a. If Patient Not Screened for Uterine Malignancy, or Those that have Not Had an Ultrasound and/or Endometrial Sampling of Any Kind, Reason Not Given equals Yes, include in Data Completeness Met and Performance Not Met.
- b. 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.
- c. If Patient not Screened for Uterine Malignancy, or those that have not had an Ultrasound and/or Endometrial Sampling of any kind, Reason not Given equals No, proceed to check Data Completeness Not Met.
- 9. Check Data Completeness Not Met:
 - a. If Data Completeness Not Met, the Quality-Data Code was not submitted. 10 procedures have been subtracted from the Data Completeness Numerator in the Sample Calculation.

SAMPLE CALCULATIONS: Data Completeness= Denominator Exclusion (x=10 procedures) + Performance Met (a=40 procedures) + Performance Not Met (o=20 procedures) = 70 procedures = 87.50% Eligible Population / Denominator (d=80 procedures) = 80 procedures Performance Rate= Performance Met (a=40 procedures) Data Completeness Numerator (70 procedures) - Denominator Exclusion (x=10 procedures) = 66.67%