

Quality ID #384: Adult Primary Rhegmatogenous Retinal Detachment Surgery: No Return to the Operating Room Within 90 Days of Surgery

2026 COLLECTION TYPE:

MERIT-BASED INCENTIVE PAYMENT SYSTEM (MIPS) CLINICAL QUALITY MEASURE (CQM)

MEASURE TYPE:

Outcome – High Priority

DESCRIPTION:

Patients aged 18 years and older who had surgery for primary rhegmatogenous retinal detachment who did not require a return to the operating room within 90 days of surgery.

INSTRUCTIONS:

Reporting Frequency:

This measure is to be submitted each time for denominator eligible cases as defined in the denominator criteria.

Intent and Clinician Applicability:

This measure is intended to reflect the quality of services provided for patients receiving primary rhegmatogenous retinal detachment surgery. This measure may be submitted by Merit-based Incentive Payment System (MIPS) eligible clinicians who perform the quality actions as defined by the numerator based on the services provided and the measure-specific denominator coding.

Measure Strata and Performance Rates:

This measure contains one strata defined by a single submission criteria.

This measure produces a single performance rate.

Implementation Considerations:

For the purposes of MIPS implementation, this procedure measure is submitted each time a procedure is performed.

This is an outcome measure and will be calculated solely using Merit-based Incentive Payment System (MIPS) eligible clinician, group, or third-party intermediary submitted data.

- For patients who receive the surgical procedures specified in the denominator coding, it should be submitted whether or not the patient had to return to the operating room within 90 days of surgery.
- Include only procedures performed between **January 1st and September 30th** of the performance period. This will allow the post-operative period to occur before third-party intermediaries must submit data to CMS.

Telehealth:

NOT TELEHEALTH ELIGIBLE: This measure is not appropriate for nor applicable to the telehealth setting. This measure is procedure based and therefore doesn't allow for the denominator criteria to be conducted via telehealth. It would be appropriate to remove these patients from the denominator eligible patient population. Telehealth eligibility is at the measure level for inclusion within the denominator eligible patient population and based on the measure specification definitions which are independent of changes to coding and/or billing practices.

Measure Submission:

The quality data codes listed do not need to be submitted by MIPS eligible clinicians, groups, or third party intermediaries that utilize this collection type for submissions; however, these codes may be submitted for those third party intermediaries that utilize Medicare Part B claims data. The coding provided to identify the measure criteria: Denominator or Numerator, may be an example of coding that could be used to identify patients that meet the intent of this clinical topic. When

implementing this measure, please refer to the 'Reference Coding' section to determine if other codes or code languages that meet the intent of the criteria may also be used within the medical record to identify and/or assess patients. For more information regarding Application Programming Interface (API), please refer to the Quality Payment Program (QPP) website.

DENOMINATOR:

Patients aged 18 years or older who had surgery for primary rhegmatogenous retinal detachment.

Denominator Criteria (Eligible Cases):

Patients aged \geq 18 years on the date of the procedure

AND

Patient procedure during the performance period (CPT): 67107, 67108, 67110

AND NOT

DENOMINATOR EXCLUSION:

Surgical procedures that included the use of silicone oil: G9756

NUMERATOR:

Patients who did not return to the operating room within 90 days for complications within the operative eye.

NUMERATOR NOTE:

For the purposes of meeting the numerator, complications are only those related to the following procedures: 67107, 67108, 67110

Numerator Options:

Performance Met:

Patient did not require a return to the operating room within 90 days of surgery (G9515)

OR

Performance Not Met:

Patient required a return to the operating room within 90 days of surgery (G9514)

RATIONALE:

The goal of treatment for retinal breaks is to create a firm chorioretinal adhesion in the attached retina immediately adjacent to and surrounding the retinal tear using cryotherapy or laser photocoagulation to halt the progression of subretinal fluid from detaching the neurosensory retina. Treatment of peripheral horseshoe tears should be extended to the ora serrata if the tear cannot be surrounded using laser or cryotherapy. The most common cause of failure in treating horseshoe tears is failure to adequately treat the tear, particularly the anterior border. Continued vitreous traction may extend the tear beyond the treated area and allow fluid to dissect through the subretinal space to cause a clinical retinal detachment. Treatment of dialyses must extend over the entire length of the dialysis, reaching the ora serrata beyond each horn or end of the dialysis.

Sufficient evidence exists for treating acute, symptomatic horseshoe tears. There is insufficient evidence for management of other vitreoretinal abnormalities. A Cochrane systematic review found that in making the decision to treat other vitreoretinal abnormalities, including lattice degeneration and asymptomatic retinal breaks, the risks that treatment will be unnecessary, ineffective, or harmful must be weighed against the possible benefit of reducing the rate of subsequent retinal detachment.

In a study published in 2011, Schall and colleagues studied the success rate with 4 surgical techniques. Initial success rate for retinal reattachment was 86% for scleral buckling only, 90% for vitrectomy only, 94% for the combination of scleral buckling and vitrectomy, and 63% for pneumatic retinopexy surgery. Patients undergoing pneumatic retinopexy had a lower initial success rate, however there was no statistically significant difference in initial reattachment rates between the other three groups. In a 2002 study, Ling and colleagues reported an 85% success rate with a single procedure. Of the 15% that initially failed, 97% were successful with one additional surgery.

CLINICAL RECOMMENDATION STATEMENTS:

This is an outcome measure. As such, no clinical recommendations are included.

REFERENCES:

Day S, Grossman DS, Mruthyunjaya P, Sloan FA, Lee PP. One year outcomes after retinal detachment surgery among Medicare beneficiaries. *Am J Ophthalmol* 2010; 150(3):338-45, Massachusetts Eye and Ear Infirmary, Harvard Medical School. *Ophthalmology Quality & Outcomes Report* 2012.

Flaxel CJ, Adelman RA, Bailey ST et al. Posterior Vitreous Detachment, Retinal Breaks, and Lattice Degeneration Preferred Practice Pattern®. *Ophthalmology*. 2020;127:P146-P181.

Ling, et al, Retinal detachment surgery in district general hospitals: An Audit of Changing Practice, *Br J Ophthalmology* 2002; 86:827-833.

Schall S, Sherman MP, Barr CC, Kaplan HJ, Primary retinal detachment repair: comparison of 1-year outcomes of four surgical techniques. *Retina* 2011 Sep; 31(8):1500-4.

Sullivan PM, Luff AJ, Aylward GW. Results of primary retinal reattachment surgery: a prospective audit. *Eye* 1997; 11:869-71.

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The American Association of Eye and Ear Centers of Excellence's (AAECE) significant past efforts and contributions to the development and updating of the measure is acknowledged. The Academy is solely responsible for the review and enhancement ("Maintenance") of the measure as of June 5, 2015.

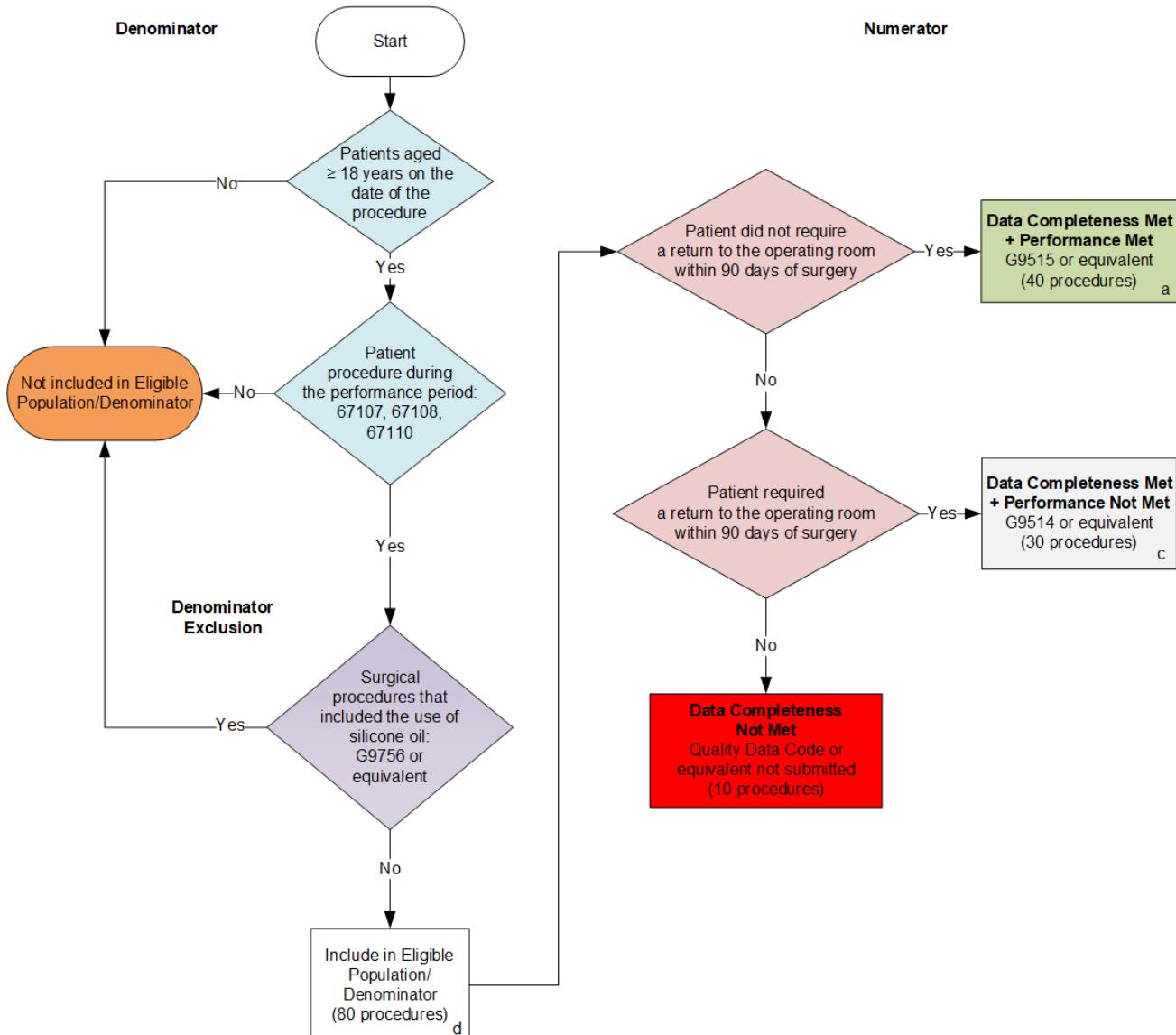
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**2026 Clinical Quality Measure Flow for Quality ID #384:
Adult Primary Rhegmatogenous Retinal Detachment Surgery:
No Return to the Operating Room Within 90 Days of Surgery**

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



SAMPLE CALCULATIONS

Data Completeness=

$$\frac{\text{Performance Met (a=40 procedures)} + \text{Performance Not Met (c=30 procedures)}}{\text{Eligible Population / Denominator (d=80 procedures)}} = \frac{70 \text{ procedures}}{80 \text{ procedures}} = 87.50\%$$

Performance Rate=

$$\frac{\text{Performance Met (a=40 procedures)}}{\text{Data Completeness Numerator (70 procedures)}} = \frac{40 \text{ procedures}}{70 \text{ procedures}} = 57.14\%$$

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

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**2026 Clinical Quality Measure Flow Narrative for Quality ID #384:
Adult Primary Rhegmatogenous Retinal Detachment Surgery:
No Return to the Operating Room Within 90 Days of Surgery**

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

1. Start with Denominator
2. Check *Patients aged greater than or equal to 18 years on the date of the procedure*:
 - a. If *Patients aged greater than or equal to 18 years on the date of the procedure* equals No, do not include in *Eligible Population/Denominator*. Stop processing.
 - b. If *Patients aged greater than or equal to 18 years on the date of the procedure* equals Yes, proceed to check *Patient procedure during the performance period*.
3. Check *Patient procedure during the performance period*:
 - a. If *Patient procedure during the performance period*: equals No, do not include in *Eligible Population/Denominator*. Stop processing.
 - b. If *Patient procedure during the performance period* equals Yes, proceed to check *Surgical procedures that included the use of silicone oil*.
4. Check *Surgical procedures that included the use of silicone oil*:
 - a. If *Surgical procedures that included the use of silicone oil* equals Yes, do not include in *Eligible Population/Denominator*. Stop processing.
 - b. If *Surgical procedures that included the use of silicone oil* equals No, include in *Eligible Population/Denominator*.
5. 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.
6. Start Numerator
7. Check *Patient did not require a return to the operating room within 90 days of surgery*:
 - a. If *Patient did not require a return to the operating room within 90 days of surgery* 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 the Sample Calculation.

- b. If *Patient did not require a return to the operating room within 90 days of surgery* equals No, proceed to check *Patient required a return to the operating room within 90 days of surgery*.
8. Check *Patient required a return to the operating room within 90 days of surgery*:
 - a. If *Patient required a return to the operating room within 90 days of surgery* equals Yes, include in *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 30 procedures in the Sample Calculation.
 - b. If *Patient required a return to the operating room within 90 days of surgery* equals No, proceed to check *Data Completeness Not Met*.
9. Check *Data Completeness Not Met*:
 - If *Data Completeness Not Met*, the Quality Data Code or equivalent was not submitted. 10 procedures have been subtracted from the Data Completeness Numerator in the Sample Calculation.

Sample Calculations

Data Completeness equals Performance Met (a equals 40 procedures) plus Performance Not Met (c equals 30 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). All equals 40 procedures divided by 70 procedures. All equals 57.14 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.