Quality ID #326 (NQF 1525): Atrial Fibrillation and Atrial Flutter: Chronic Anticoagulation Therapy

- National Quality Strategy Domain: Effective Clinical Care
- Meaningful Measure Area: Management of Chronic Conditions

2021 COLLECTION TYPE:

MEDICARE PART B CLAIMS

MEASURE TYPE:

Process

DESCRIPTION:

Percentage of patients aged 18 years and older with nonvalvular atrial fibrillation (AF) or atrial flutter who were prescribed warfarin OR another FDA-approved oral anticoagulant drug for the prevention of thromboembolism during the measurement period

INSTRUCTIONS:

This measure is to be submitted a minimum of <u>once per performance period</u> for patients with nonvalvular AF or atrial flutter seen during the performance period. 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.

NOTE: Patient encounters for this measure conducted via telehealth (e.g., encounters coded with GQ, GT, 95, or POS 02 modifiers) are allowable.

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 aged 18 years and older with a diagnosis of nonvalvular AF or atrial flutter who do not have a documented CHA₂ DS₂.VASc risk score of 0 or 1 for men; or 0, 1, or 2 for women.

DENOMINATOR NOTE: *Signifies that this CPT Category I code is a non-covered service under the Medicare Part B Physician Fee Schedule (PFS). These non-covered services will not be counted in the denominator population for Medicare Part B claims measures.

Denominator Criteria (Eligible Cases):

Patients aged ≥ 18 years on the date of encounter

AND

Diagnosis for nonvalvular atrial fibrillation or atrial flutter (ICD-10-CM): 148.0, 148.3, 148.4, 148.11, 148.19, 148.20, 148.21, 148.91, 148.92

AND

Patient encounter during the performance period (CPT): 99202, 99203, 99204, 99205, 99212, 99213, 99214, 99215, 99241*, 99242*, 99243*, 99244*, 99245*, 99304, 99305, 99306, 99307, 99308, 99309, 99310, 99324, 99325, 99326, 99327, 99328, 99334, 99335, 99336, 99337, 99341, 99342, 99343, 99344, 99345, 99347, 99348, 99349, 99350

NUMERATOR:

Patients with nonvalvular AF or atrial flutter for whom warfarin or another FDA-approved oral anticoagulant was prescribed

Version 5.0 November 2020

Definition:

Prescribed – also satisfied by documentation in current medication list.

Comfort Care Only - Refers to medical treatment of a dying person where the natural dying process is permitted to occur while assuring maximum comfort. It includes attention to the psychological and spiritual needs of the patient and support for both the dying patient and the patient's family. Comfort Measures Only is commonly referred to as "comfort care" by the general public. It may be completed in an inpatient, outpatient or home environment. Comfort Measures Only includes hospice, palliative and supportive treatment for patients who are suffering from a terminal illness—e.g., AIDS, cancer—or who have refused life-sustaining treatment. In order to use G9930, a patient must be on comfort care measures only and not be receiving any other types of care. It is not equivalent to a physician order to withhold emergency resuscitative measures such as Do Not Resuscitate (DNR).

CHA2DS2-VASc Stroke Risk Assessment - The assessment of patients with nonvalvular AF or atrial flutter, assessment of thromboembolic risk should include:

<u>Score</u>
1
1
2
1
2
1
1
1

NUMERATOR NOTE: Denominator Exclusions/Exception(s) are determined on the date of the denominator eliaible encounter.

The intent of the denominator exclusion **G9931** is to allow patients with a low risk for a thromboembolic event (i.e. a CHA2DS2-VASc score of 0 or 1 for men; or 0, 1, or 2 for women) to be excluded from the sample. This denominator exclusion serves as documentation that a patient's risk for a thromboembolic event was appropriately assessed using the CHA₂DS₂-VASc scoring tool and that the risk was low enough to not warrant anticoagulation treatment. In order to exclude low risk patients, eligible clinicians must use the CHA2 DS2-VASc assessment tool to determine a patient's risk score and must document either the numeric score (i.e. 0 or 1 for men; or 0, 1, or 2 for women) or all the individual risk factors assessed to support an assessment of the CHA₂DS₂-VASc score.

NUMERATOR QUALITY-DATA CODING OPTIONS:

If patient is not eligible for this measure because patient has transient or reversible cause of AF OR patient is receiving comfort care OR CHA₂DS₂-VASc of 0 or 1 for men; or 0, 1, or 2 for women (One of three quality-data codes [G9929 OR G9930 OR G9931] is required on the claim form to submit this numerator option)

Denominator Exclusion: G9929: Patient has transient or reversible cause of AF (e.g.,

pneumonia, hyperthyroidism, pregnancy, cardiac

surgery)

OR

Denominator Exclusion: G9930: Patients who are receiving comfort care only

Denominator Exclusion: G9931: Documentation of CHA₂DS₂-VASc risk score of 0 or 1 for

men; or 0, 1, or 2 for women

OR

Warfarin or Another FDA-Approved Oral Anticoagulant that is Prescribed

(One quality-data code [G8967] is required on the claim form to submit this numerator option)

Performance Met: G8967: Warfarin OR another FDA-approved oral anticoagulant

is prescribed

<u>OR</u>

Warfarin or Another FDA-Approved Anticoagulant that is Not Prescribed for Medical Reasons

(One quality-data code [G8968] is required on the claim form to submit this numerator option)

Denominator Exception: G8968: Documentation of medical reason(s) for not prescribing

warfarin OR another FDA-approved anticoagulant (e.g.,

atrial appendage device in place)

OR

Warfarin or Another FDA-Approved Anticoagulant Not Prescribed for PatientReasons

(One quality-data code [G8969] is required on the claim form to submit this numerator option)

Denominator Exception: G8969: Documentation of patient reason(s) for not prescribing

warfarin OR another oral anticoagulant that is FDA approved for the prevention of thromboembolism (e.g., patient choice of having atrial appendage device

placed)

<u>OR</u>

Warfarin or Another FDA-Approved Anticoagulant Not Prescribed for System Reasons

(One quality-data code [G9927] is required on the claim form to submit this numerator option)

Denominator Exception G9927: Documentation of system reason(s) for not prescribing

warfarin OR another FDA-approved anticoagulation due to patient being currently enrolled in a clinical trial

related to AF/atrial flutter treatment

<u>OR</u>

Warfarin OR Another FDA-Approved Anticoagulant Not Prescribed, Reason not Given

(One quality-data code [G9928] is required on the claim form to submit this numerator option)

Performance Not Met: G9928: Warfarin OR another FDA-approved anticoagulant not

prescribed, reason not given

RATIONALE:

AF, whether paroxysmal, persistent, or permanent and whether symptomatic or silent, significantly increases the risk of thromboembolic ischemic stroke. Nonvalvular atrial fibrillation increases the risk of stroke 5 times, and AF in the setting of mitral stenosis increases the risk of stroke 20 times over that of patients in sinus rhythm.

Thromboembolism occurring with AF is associated with a greater risk of recurrent stroke, more severe disability, and mortality. Silent AF is also associated with ischemic stroke. The appropriate use of antithrombotic therapy and the control of other risk factors, including hypertension and hypercholesterolemia, substantially reduce stroke risk.

One meta-analysis has stratified ischemic stroke risk among patients with nonvalvular AF using the following point scoring systems: AF Investigators; CHA₂DS₂ (congestive heart failure, hypertension, age 75 years, diabetes mellitus, prior stroke or TIA or thromboembolism [doubled]), or CHA₂DS₂-VASc (congestive heart failure, hypertension, age 75 years [doubled], diabetes mellitus, prior stroke or TIA or thromboembolism [doubled], vascular disease, age 65 to 74 years, sex category).

When compared with the CHA₂DS₂ score, the CHA₂DS₂-VASc score for nonvalvular AF has a broader score range (0 to 9) and includes a larger number of risk factors (female sex, 65 to 74 years of age, and vascular disease).

The selection of an antithrombotic agent should be based on shared decision making that takes into account risk factors, cost, tolerability, patient preference, potential for drug interactions, and other clinical characteristics, including time in the INR therapeutic range if the patient has been on warfarin, irrespective of whether the AF pattern is paroxysmal, persistent, or permanent.

CLINICAL RECOMMENDATION STATEMENTS:

- In patients with AF, anticoagulant therapy should be individualized based on shared decision-making after discussion of the absolute and RRs of stroke and bleeding, and the patient's values and preferences. (Class I, Level of Evidence: C)
- 2. Selection of anticoagulant therapy should be based on the risk of thromboembolism irrespective of whether the AF pattern is paroxysmal, persistent, or permanent. (Class I, Level of Evidence: B)
- 3. In patients with AF (except with moderate-to-severe mitral stenosis or a mechanical heart valve), the CHA2DS2-VASc score is recommended for assessment of stroke risk. (Class I, Level of Evidence: B)
- 4. For patients with AF who have mechanical heart valves, warfarin is recommended. (Class I, Level of Evidence: B)
- 5. For patients with AF and an elevated CHA2DS2-VASc score of 2 or greater in men or 3 or greater in women, oral anticoagulants are recommended.

Options include:

Warfarin (LOE: A) (S4.1.1-5–S4.1.1-7)

• Dabigatran (LOE: B) (S4.1.1-8) • Rivaroxaban (LOE: B) (S4.1.1-9)

 Apixaban (LOE: B) (S4.1.1-10), or • Edoxaban (LOE: B-R) (S4.1.1-11)

- 6. Among patients treated with warfarin, the INR should be determined at least weekly during initiation of anticoagulation therapy and at least monthly when anticoagulation (INR in range) is stable. (Class I, Level of Evidence: A)
- 7. For patients with nonvalvular AF unable to maintain a therapeutic INR level with warfarin, NOAC is recommended. (Class I, Level of Evidence: C)
- 8. Re-evaluation of the need for and choice of anticoagulation therapy at periodic intervals is recommended to reassess stroke and bleeding risks. (Class I, Level of Evidence: C)
- 9. For patients with atrial flutter, anticoagulation therapy is recommended according to the same risk profile used for AF. (Class I, Level of Evidence: C)

COPYRIGHT:

Physician performance measures and related data specifications were developed by the American College of Cardiology (ACC) and the American Heart Association (AHA) to facilitate quality improvement activities by physicians. These performance measures are not clinical quidelines and do not establish a standard of medical care, and have not been tested for all potential applications. While copyrighted, they can be reproduced and distributed, without modification, for noncommercial purposes, e.g., use by health care providers in connection with their practices. Commercial use is defined as the sale, license, or distribution of the performance measures for commercial gain, or incorporation of the performance measures into a product or service that is sold, licensed or distributed for commercial gain. Commercial uses of the measures require a license agreement between the user and the ACC or the AHA. Neither the ACC or the AHA, nor its members shall be responsible for any use of these measures.

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

© 2020 American College of Cardiology and American Heart Association. All Rights Reserved.

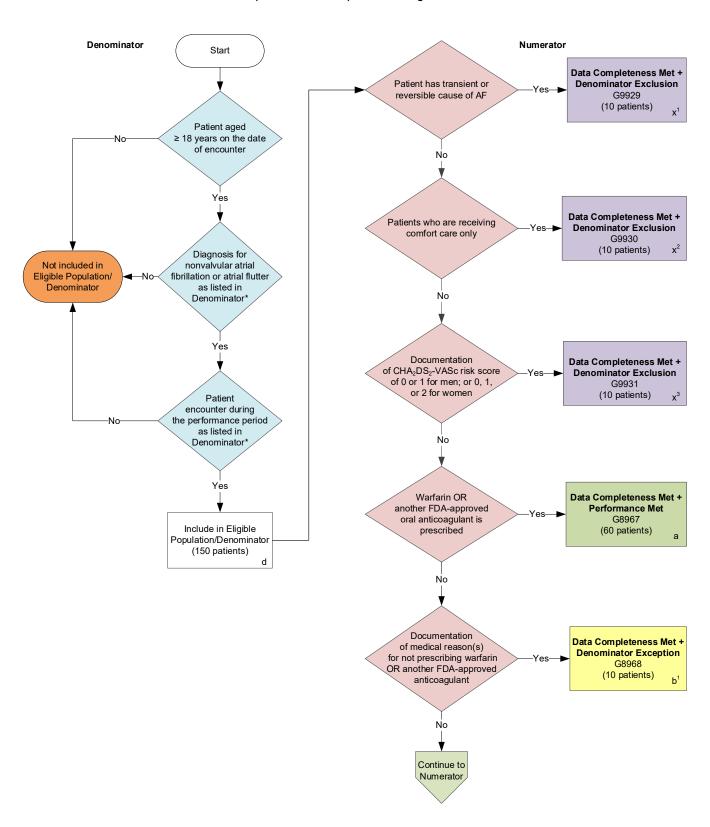
Limited proprietary coding is contained in the measure specifications for convenience. Users of the proprietary code sets should obtain all necessary licenses from the owners of these code sets. The ACC and the AHA, and its members disclaim all liability for use or accuracy of any Current Procedural Terminology (CPT®) or other coding contained in the specifications.

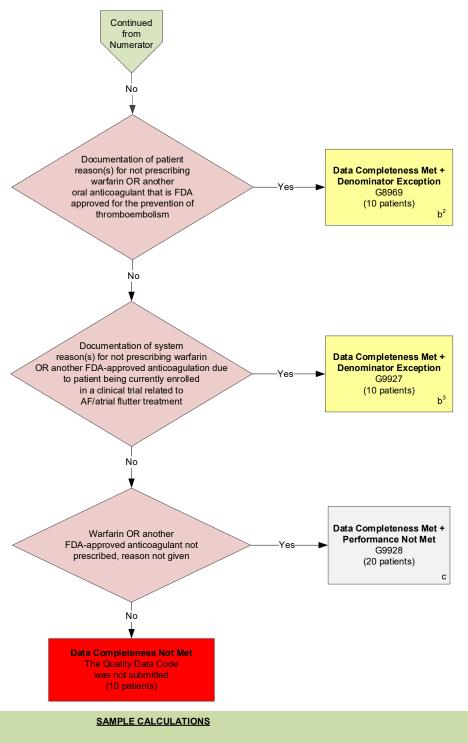
CPT® contained in the measures specifications is copyright 2004-2020 American Medical Association. LOINC® copyright 2004-2020 Regenstrief Institute, Inc. This material contains SNOMED CLINICAL TERMS (SNOMED CT®) copyright 2004-2020 International Health Terminology Standards Development Organisation. ICD-10 is copyright 2020 World Health Organization. All Rights Reserved. Use of SNOMED CT® is only authorized within the United States

The American Medical Association's and the PCPI® Foundation's significant past efforts and contributions to the performance measures are gratefully acknowledged.

2021 Medicare Part B Claims Flow for Quality ID #326 (NQF 1525): Atrial Fibrillation and Atrial Flutter: Chronic Anticoagulation Therapy

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





SAMPLE CALCULATIONS Data Completeness= Denominator Exclusion (x¹+x²+x³=30 pts) + Performance Met (a=60 pts) + Denominator Exception (b¹+b²+b³=30 pts) + Performance Not Met (c=20 pts) = 140 pts = 150 pts Eligible Population / Denominator (d=150 pts) = 150 pts = 150 pts Performance Rate= Performance Met (a=60 pts) = 60 pts = 75.00% Data Completeness Numerator (140 pts) - Denominator Exclusion (x¹+x²+x³=30 pts) - Denominator Exception (b¹+b²+b³=30 pts) = 80 pts

CPT only copyright 2020 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.

v5

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

NOTE: Submission Frequency: Patient-Process

CPT only copyright 2020 Amer
The measure diagrams were d

2021 Medicare Part B Claims Flow Narrative for Quality ID #326 (NQF 1525): Atrial Fibrillation and Atrial Flutter: Chronic Anticoagulation Therapy

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 encounter.
 - a. If Patients aged greater than or equal to 18 years on the date of encounter 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 encounter equals Yes, proceed to check Diagnosis for nonvalvular atrial fibrillation or atrial flutter as listed in Denominator*.
- 3. Check Diagnosis for nonvalvular atrial fibrillation or atrial flutter as listed in Denominator*:
 - a. If Diagnosis for nonvalvular atrial fibrillation or atrial flutter as listed in Denominator* equals No, do not include in Eligible Population/Denominator. Stop processing.
 - b. If Diagnosis for nonvalvular atrial fibrillation or atrial flutter as listed in Denominator* equals Yes, proceed to check Patient encounter during the performance period as listed in Denominator*.
- 4. Check Patient encounter during the performance period as listed in Denominator*:
 - a. If Patient encounter during the performance period as listed in Denominator* equals No, do not include in Eligible Population/Denominator. Stop processing.
 - b. If Patient encounter during the performance period as listed in Denominator* equals Yes, include in Eligible Population/Denominator.
- 5. Denominator Population:
 - Denominator Population is all Eligible Patients in the Denominator. Denominator is represented
 as Denominator in the Sample Calculation listed at the end of this document. Letter d equals
 150 patients in the Sample Calculation.
- 6. Start Numerator
- 7. Check Patient has transient or reversible cause of AF:
 - a. If Patient has transient or reversible cause of AF equals Yes, include in Data Completeness Met and Denominator Exclusion.
 - Data Completeness Met and Denominator Exclusion is represented in the Data Completeness and Performance Rate in the Sample Calculation listed at the end of this document. Letter x¹ equals 10 patients in the Sample Calculation.
 - b. If Patient has transient or reversible cause of AF equals No, proceed to check Patients who are receiving comfort care only.
- 8. Check Patients who are receiving comfort care only:
 - a. If Patients who are receiving comfort care only equals Yes, include in Data Completeness Met and Denominator Exclusion.

- Data Completeness Met and Denominator Exclusion is represented in the Data Completeness and Performance Rate in the Sample Calculation listed at the end of this document. Letter x² equals 10 patients in the Sample Calculation.
- b. If Patients who are receiving comfort care only equals No, proceed to check Documentation of CHA₂DS₂-VASc risk score of 0 or 1 for men; or 0, 1, or 2 for women.
- 9. Check Documentation of CHA₂DS₂-VASc risk score of 0 or 1 for men: or 0, 1, or 2 for women:
 - a. If Documentation of CHA₂DS₂-VASc risk score of 0 or 1 for men; or 0, 1, or 2 for women equals Yes, include in Data Completeness Met and Denominator Exclusion.
 - Data Completeness Met and Denominator Exclusion is represented in the Data Completeness and Performance Rate in the Sample Calculation listed at the end of this document. Letter x³ equals 10 patients in the Sample Calculation.
 - b. If Documentation of CHA₂DS₂-VASc risk score of 0 or 1 for men; or 0, 1, or 2 for women equals No, proceed to check Warfarin OR another FDA-approved oral anticoagulant is prescribed.
- 10. Check Warfarin OR another FDA-approved oral anticoagulant is prescribed:
 - a. If Warfarin OR another FDA-approved oral anticoagulant is prescribed equals Yes, include in Data Completeness Met and Performance Met.
 - Data Completeness Met and Performance Met is represented in the Data Completeness and Performance Rate in the Sample Calculation listed at the end of this document. Letter a equals 60 patients in the Sample Calculation.
 - b. If Warfarin OR another FDA-approved oral anticoagulant is prescribed equals No, proceed to check Documentation of medical reason(s) for not prescribing warfarin OR another FDA-approved anticoagulant.
- 11. Check Documentation of medical reason(s) for not prescribing warfarin OR another FDA-approved anticoagulant:
 - a. If Documentation of medical reason(s) for not prescribing warfarin OR another FDA-approved anticoagulant equals Yes, include in Data Completeness Met and Denominator Exception.
 - Data Completeness Met and Denominator Exception is represented in the Data Completeness and Performance Rate in the Sample Calculation listed at the end of this document. Letter b¹ equals 10 patients in the Sample Calculation.
 - b. If Documentation of medical reason(s) for not prescribing warfarin OR another FDA-approved anticoagulant equals No, proceed to check Documentation of patient reason(s) for not prescribing warfarin OR another oral anticoagulant that is FDA approved for the prevention of thromboembolism.
- 12. Check Documentation of patient reason(s) for not prescribing warfarin OR another oral anticoagulant that is FDA approved for the prevention of thromboembolism:
 - a. If Documentation of patient reason(s) for not prescribing warfarin OR another oral anticoagulant that is FDA approved for the prevention of thromboembolism equals Yes, include in the Data Completeness Met and Denominator Exception.
 - Data Completeness Met and Denominator Exception is represented in the Data Completeness

and Performance Rate in the Sample Calculation listed at the end of this document. Letter b² equals 10 patients in the Sample Calculation.

- b. If Documentation of patient reason(s) for not prescribing warfarin OR another oral anticoagulant that is FDA approved for the prevention of thromboembolism equals No, proceed to check Documentation of system reason(s) for not prescribing warfarin OR another FDA-approved anticoagulation due to patient being currently enrolled in a clinical trial related to AF/atrial flutter treatment.
- 13. Check Documentation of system reason(s) for not prescribing warfarin OR another FDA-approved anticoagulation due to patient being currently enrolled in a clinical trial related to AF/atrial flutter treatment:
 - a. If Documentation of system reason(s) for not prescribing warfarin OR another FDA-approved anticoagulation due to patient being currently enrolled in a clinical trial related to AF/atrial flutter treatment equals Yes, include in the Data CompletenessMet and Denominator Exception.
 - Data Completeness Met and Denominator Exception is represented in the Data Completeness and Performance Rate in the Sample Calculation listed at the end of this document. Letter b³ equals 10 patients in the Sample Calculation.
 - b. If Documentation of system reason(s) for not prescribing warfarin OR another FDA-approved anticoagulation due to patient being currently enrolled in a clinical trial related to AF/atrial flutter treatment equals No, proceed to check Warfarin OR another FDA-approved anticoagulant not prescribed, reason not given.
- 14. Check Warfarin OR another FDA-approved anticoagulant not prescribed, reason not given:
 - a. If Warfarin OR another FDA-approved anticoagulant not prescribed, reason not given equals Yes. include in Data Completeness Met and Performance Not Met.
 - Data Completeness Met and Performance Not Met is represented in the Data Completeness in the Sample Calculation listed at the end of this document. Letter c equals 20 patients in the Sample Calculation.
 - b. If Warfarin OR another FDA-approved anticoagulant not prescribed, reason not given, proceed to check Data Completeness Not Met.
- 15. Check Data Completeness Not Met:
 - a. If Data Completeness Not Met, the Quality Data Code was not submitted. 10 patients have been subtracted from the Data Completeness Numerator in the Sample Calculation.

Sample Calculations

Data Completeness equals Denominator Exclusion (x1 plus x2 plus x3 equals 30 patients) plus Performance Met (a equals 60 patients) plus Denominator Exception (b¹ plus b² plus b³ equals 30 patients) plus Performance Not Met (c equals 20 patients) divided by Eligible Population / Denominator (d equals 150 patients). All equals 140 patients divided by 150 patients). All equals 93.33 percent.

Performance Rate equals Performance Met (a equals 60 patients) divided by Data Completeness Numerator (140 patients) minus Denominator Exclusion (x1 plus x2 plus x3 equals 30 patients) minus Denominator Exception (b1 plus b² plus b³ equals 30 patients). All equals 60 patients divided by 80 patients. All equals 75.00 percent.

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

NOTE: Submission Frequency: Patient-Process

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.