Measure #44 (NQF 0236): Coronary Artery Bypass Graft (CABG): Preoperative Beta-Blocker in Patients with Isolated CABG Surgery – National Quality Strategy Domain: Effective Clinical Care

2016 PQRS OPTIONS FOR INDIVIDUAL MEASURES:
REGISTRY ONLY

DESCRIPTION:
Percentage of isolated Coronary Artery Bypass Graft (CABG) surgeries for patients aged 18 years and older who received a beta-blocker within 24 hours prior to surgical incision.

INSTRUCTIONS:
This measure is to be reported each time an isolated CABG procedure is performed during the reporting period. It is anticipated that eligible professionals who provide services for isolated CABG will submit this measure. The timeframe for this measure includes the entire 24 hour period prior to the surgical incision time.

Measure Reporting via Registry:
CPT codes and patient demographics are used to identify patients who are included in the measure’s denominator. The listed numerator options are used to report the numerator of the measure.

The quality-data codes listed do not need to be submitted for registry-based submissions; however, these codes may be submitted for those registries that utilize claims data.

DENOMINATOR:
Isolated CABG surgeries for patients aged 18 years and older

Definition:
Isolated CABG – Refers to CABG using arterial and/or venous grafts only.

Denominator Criteria (Eligible Cases):
Patients aged ≥ 18 years on date of encounter
AND
Patient encounter during the reporting period (CPT): 00566, 00567, 33510, 33511, 33512, 33513, 33514, 33516, 33517, 33518, 33519, 33521, 33522, 33523, 33533, 33534, 33535, 33536
OR
Patient encounter during the reporting period (CPT): 33510, 33511, 33512, 33513, 33514, 33516, 33517, 33518, 33519, 33521, 33522, 33523, 33530, 33533, 33534, 33535, 33536
AND
Patient encounter during the reporting period (CPT): 00562, 33530

NUMERATOR:
Patients who received a beta-blocker within 24 hours prior to surgical incision of isolated CABG surgeries

Definition:
Medical Reason – Eligible professional must document specific reason(s) for not administering beta-blockers.

Numerator Options:
Performance Met: Beta blocker administered within 24 hours prior to surgical incision (4115F)

OR
**Medical Performance Exclusion:** Documentation of medical reason(s) for not administering beta blocker within 24 hours prior to surgical incision (e.g., not indicated, contraindicated, other medical reason) (4115F with 1P)

**OR**

**Performance Not Met:** Beta blocker not administered within 24 hours prior to surgical incision, reason not otherwise specified (4115F with 8P)

**RATIONALE:**
Since its introduction in 1962, coronary artery bypass grafting (CABG) has continued to be the gold standard for revascularization of CAD, particularly in high-risk patients with multivessel disease. Evidence from multiple studies suggests that CABG prolongs survival especially in complex patients with diabetes, those aged 65 years or more, those with left main stem or triple-vessel disease, and those with impaired left ventricular function (El Bardissi et al., 2012)

The NHDS (NCHS) estimates that in 2010, in the United States, 219 000 patients underwent a total of 397,000 coronary artery bypass procedures (defined by procedure codes) (Go et al., 2014). Despite significant developments in PCI, CABG remains the most commonly used treatment option for patients with complex CAD and high-risk patients (El Bardissi et al., 2012).

Coronary revascularization, comprising coronary artery bypass graft (CABG) surgery and percutaneous coronary intervention (PCI), is among the most common major medical procedures provided by the US health care system, with more than 1 million procedures performed annually. It is also among the most costly procedure. Medicare inpatient payments to hospitals for coronary revascularizations exceeded $6.7 billion in fiscal year 2006 and are larger than the reimbursement for any other medical or surgical procedure (Epstein, 2011).

Postoperative atrial fibrillation (POAF) is a common complication following cardiac surgery, occurring in 25-40% of patients (Crystal, 2004, Burgess, 2006). POAF has been associated with increased rates of post-operative morbidity and mortality and consequently, increased costs (Mariscalco, 2008, Crystal, 2004, Bramer, 2010). Prophylactic administration of beta-blockers has been shown to reduce the risk of POAF and mortality following isolated coronary artery bypass graft surgery (Connolly, 2003, Mariscalco, 2008, Ferguson, 2002). Khan’s meta-analysis of RCTs found that “Preoperative BB initiation resulted in 52% reduction in the incidence of AF as compared to controls, however these results were not statistically significant”. El Bardissi (2012) showed a 19.5% increase in preoperative use of beta-blockers from 2000-2009.

Prophylaxis to prevent atrial fibrillation after cardiac surgery with any of the studied pharmacological or non-pharmacological interventions may be favored because of its reduction in the rate of atrial fibrillation, decrease in the length of stay and cost of hospital treatment and a possible decrease in the rate of stroke (Arsenault et al., 2013). "According to our findings, perioperative application of beta-blockers still plays a pivotal role in cardiac surgery, as they can substantially reduce the high burden of supraventricular and ventricular arrhythmias in the aftermath of surgery. Their influence on mortality, AMI, stroke, congestive heart failure, hypotension and bradycardia in this setting remains unclear (Blessberger et al., 2014)."

Postoperative AF after cardiac operations is associated with postoperative morbidities such as cerebrovascular accidents (CVA), infections (e.g., septicemia, pneumonia and mediastinitis), and renal failure. Previous studies have suggested that POAF after CABG is related to early and late mortality (Bramer et al., 2010). Development of AF immediately after coronary artery bypass surgery (CABG) results in a longer stay in the intensive care unit and in hospital, together with a significantly higher (two-to-three-fold) risk of post-operative stroke (Burgess et al., 2006).

AF complicates up to 40% of the 500,000 patients per year undergoing CABG and increase the cost of the procedure by 10,055 per case resulting in incremental cost of about & 2 billion annually. It also increases the length of stay to
additional 4-5 days and identifies a subset of patients at increased risk of morbidity, strokes and in-hospital and long-term mortality (Khan et al., 2013)

**CLINICAL RECOMMENDATION STATEMENTS:**

Preoperative Beta-blockers (ACCF/AHA, 2011):

Class I

1) Beta-blockers should be administered for at least 24 hours before CABG to all patients without contraindications to reduce the incidence or clinical sequelae of postoperative AF. (Level of Evidence: B), (ACCF/AHA, 2011)

Class IIa

1) Preoperative use of beta-blockers in patients without contraindications, particularly in those with an LV ejection fraction (LVEF) greater than 30%, can be effective in reducing the risk of in-hospital mortality. (Level of Evidence: B), (ACCF/AHA, 2011)

2) Beta-blockers can be effective in reducing the incidence of perioperative myocardial ischemia. (Level of Evidence: B), (ACCF/AHA, 2011)

Class IIb

1) The effectiveness of preoperative beta-blockers in reducing in-hospital mortality rate in patients with LVEF less than 30% is uncertain. (Level of Evidence: B), (ACCF/AHA, 2011)

Treatment of arrhythmias after revascularization (ESC/EACTS, 2014)

Class I

1) Beta-blockers are recommended to decrease the incidence of atrial fibrillation after CABG in the absence of contraindications. (Level of Evidence: A), (ESC/EACTS, 2014)

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2016 Registry Individual Measure Flow
Measure #44 NQF #0236: Coronary Artery Bypass Graft (CABG): Preoperative Beta-Blocker in Patients with Isolated CABG Surgery

Start

Denominator

Patient Age at Date of Service ≥18 Years

No

Not Included in Eligible Population/Denominator

Yes

Encounter as Listed in Denominator (1/1/2016 thru 12/31/2016)†

No

Encounter Code as Listed in Denominator (1/1/2016 thru 12/31/2016)†

Yes

Include in Eligible Population/Denominator (8 procedures) d

Numerator

Beta-Blocker Administered Within 24 Hours Prior to Surgical Incision

Yes

Reporting Met + Performance Met 4115F or equivalent (4 procedures) a

No

Beta-Blocker Not Administered Within 24 Hours Prior to Surgical Incision, Reason Not Specified

Yes

Reporting Met + Performance Not Met 4115F-SF or equivalent (2 procedures) c

No

Beta-Blocker Not Administered Within 24 Hours Prior to Surgical Incision for Medical Reasons

Yes

Reporting Met + Performance Exclusion 4115F-1P or equivalent (1 procedure) b

No

Reporting Not Met, Quality-Data Code or equivalent not reported (1 procedure)

SAMPLE CALCULATIONS:

Reporting Rate =
Performance Met (a=4 procedures) + Performance Exclusion (b=1 procedure) + Performance Not Met (c=2 procedures) = 7 procedures = 87.50%
Eligible Population / Denominator (d=8 procedures) =

Performance Rate =
Performance Met (a=4 procedures) = 4 procedures = 66.67%
Reporting Numerator (7 procedures) – Performance Exclusion (b=1 procedure) = 6 procedures

*See the posted Measure Specification for specific coding and instructions to report this measure.
NOTE: Reporting Frequency - Procedure

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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.
2016 Registry Individual Measure Flow
PQRS #44 NQF #0236: Coronary Artery Bypass Graft (CABG): Preoperative Beta-Blocker in Patients with Isolated CABG Surgery

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

1. Start with Denominator
2. Check Patient Age:
   a. If Age greater than or equal to 18 years of age on Date of Service equals No during the measurement period, do not include in Eligible Patient Population. Stop Processing.
   b. If Age greater than or equal to 18 years of age on Date of Service equals Yes during the measurement period, proceed to check Encounter for New CABG as Listed in the Denominator.
3. Check Encounter for Isolated CABG as Listed in the Denominator:
   a. If Encounter for Isolated CABG as Listed in the Denominator equals No, proceed to check Encounter for Re-Operation as Listed in the Denominator.
   b. If Encounter for Isolated CABG as Listed in the Denominator CABG equals Yes, proceed to include in Eligible population.
4. Check Encounter for Re-Operation as Listed in the Denominator:
   a. If Encounter for Re-Operation as Listed in the Denominator equals No, do not include in Eligible Patient Population. Stop Processing.
   b. If Encounter for Re-Operation as Listed in the Denominator equals Yes, proceed to check Encounter Code as Listed in the Denominator.
5. Check Encounter Code as Listed in the Denominator:
   a. If Encounter Code as Listed in the Denominator equals No, do not include in Eligible Patient Population. Stop Processing.
   b. If Encounter Code as Listed in the Denominator equals Yes, proceed to include in Eligible population.
6. Denominator Population:
   a. 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 8 procedures in the sample calculation.
7. Start Numerator
8. Check Beta-Blocker Administered Within 24 Hours Prior to Surgical Incision:
   a. If Beta-Blocker Administered Within 24 Hours Prior to Surgical Incision equals Yes, include in Reporting Met and Performance Met.
b. Reporting Met and Performance Met letter is represented in the Reporting Rate and Performance Rate in the Sample Calculation listed at the end of this document. Letter a equals 4 procedures in Sample Calculation.

c. If Beta-Blocker Administered Within 24 Hours Prior to Surgical Incision equals No, proceed to Beta-Blocker Not Administered Within 24 Hours Prior to Surgical Incision for Medical Reasons.

9. Check Beta-Blocker Not Administered Within 24 Hours Prior to Surgical Incision for Medical Reasons:
   a. If Beta-Blocker Not Administered Within 24 Hours Prior to Surgical Incision for Medical Reasons equals Yes, include in Reporting Met and Performance Exclusion.
   
   b. Reporting Met and Performance Exclusion is represented in the Reporting Rate in the Sample Calculation listed at the end of this document. Letter b equals 1 procedure in the Sample Calculation.
   
   c. If Beta-Blocker Not Administered Within 24 Hours Prior to Surgical Incision for Medical Reasons equals No, proceed to Beta-Blocker Not Administered Within 24 Hours Prior to Surgical Incision, Reason Not Specified.

10. Check Beta-Blocker Not Administered Within 24 Hours Prior to Surgical Incision, Reason Not Specified:
    a. If Beta-Blocker Not Administered Within 24 Hours Prior to Surgical Incision, Reason Not Specified equals Yes, include in the Reporting Met and Performance Not Met.
    
    b. Reporting Met and Performance Not Met letter is represented in the Reporting Rate in the Sample Calculation listed at the end of this document. Letter c equals 2 procedures in the Sample Calculation.
    
    c. If Beta-Blocker Not Administered Within 24 Hours Prior to Surgical Incision, Reason Not Specified equals No, proceed to Reporting Not Met.

11. Check Reporting Not Met:
    a. If Reporting Not Met equals No, Quality Data Code or equivalent not reported. 1 procedure has been subtracted from the reporting numerator in the sample calculation.

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<tr>
<th>SAMPLE CALCULATIONS:</th>
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<tr>
<td>Reporting Rate=</td>
</tr>
<tr>
<td>Performance Met (a=4 procedures) + Performance Exclusion (b=1 procedure) + Performance Not Met (c=2 procedures) = 7 procedures. = 87.50%</td>
</tr>
<tr>
<td>Eligible Population/Denominator (d=8 procedures) = 8 procedures</td>
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</tbody>
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| Performance Rate=     |
| Performance Met (a=4 procedures) = 4 procedures. = 66.67% |
| Reporting Numerator (7 procedures) – Performance Exclusion (b=1 procedure) = 6 procedures |