

<b>CMS Measure #</b>	<b>PQRS #</b>	<b>National Quality Forum (NQF) #</b>	<b>Measure Title</b>	<b>Measure Description</b>	<b>Numerator Statement</b>	<b>Denominator Statement</b>	<b>Domain</b>
122v2	1	0059	Diabetes: Hemoglobin A1c Poor Control	Percentage of patient 18 – 75 years of age with diabetes who had hemoglobin A1c >9.0% during the measurement period.	Patients whose most recent HbA1c level (performed during the measurement period) is >9.0%.	Patients 18-75 years of age with diabetes with a visit during the measurement period.	Effective Clinical Care
N/A	20	0270	Perioperative Care: Timing of Prophylactic Parenteral Antibiotic	Percentage of surgical patients aged 18 years and older undergoing procedures with the indications for prophylactic parenteral antibiotics, who have an order for prophylactic parenteral antibiotic to be given within one hour (if fluoroquinolone or vancomycin, two hours), prior to the surgical incision (or start of procedure when no incision is required).	Surgical patients who have an order for prophylactic parenteral antibiotic to be given within one hour (if fluoroquinolone or vancomycin, two hours) prior to the surgical incision (or start of procedure when no incision is required).	All surgical patients aged 18 years and older undergoing procedures with the indications for prophylactic parenteral Antibiotics.	Patient Safety

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N/A	21	0268	Perioperative Care: Selection of Prophylactic Antibiotic – First OR Second Generation Cephalosporin	Percentage of surgical patients aged 18 years and older undergoing procedures with the indications for a first OR second generation cephalosporin prophylactic antibiotic, who had an order for a first OR second generation cephalosporin for antimicrobial prophylaxis.	Surgical patients who had an order for cefazolin OR cefuroxime for antimicrobial prophylaxis.	All surgical patients aged 18 years and older undergoing procedures with the indications for a first OR second generation cephalosporin prophylactic antibiotic.	Patient Safety
	22	0271	Perioperative Care: Discontinuation of Prophylactic Parenteral Antibiotics	Percentage of non-cardiac surgical patients aged 18 years and older undergoing procedures with the indications for prophylactic parenteral antibiotics AND who received a prophylactic parenteral antibiotic, who have an order for discontinuation of prophylactic parenteral antibiotics within 24 hours of surgical end time.	Non-cardiac surgical patients who have an order for discontinuation of prophylactic parenteral antibiotics within 24 hours of surgical end time.	All non-cardiac surgical patients aged 18 years and older undergoing procedures with the indications for prophylactic parenteral antibiotics AND who received a prophylactic parenteral antibiotic.	Patient Safety

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N/A	23	0239	Perioperative Care: Venous Thromboembolism (VTE) Prophylaxis	Percentage of surgical patients aged 18 years and older undergoing procedures for which VTE prophylaxis is indicated in all patients, who had an order for Low Molecular Weight Heparin (LMWH), Low-Dose Unfractionated Heparin (LDUH), adjusted-dose warfarin, fondaparinux or mechanical prophylaxis to be given within 24 hours prior to incision time or within 24 hours after surgery end time.	Surgical patients who had an order for LMWH, LDUH, adjusted-dose warfarin, fondaparinux or mechanical prophylaxis to be given within 24 hours prior to incision time or within 24 hours after surgery end time.	All surgical patients aged 18 years and older undergoing procedures for which VTE prophylaxis is indicated in all patients.	Patient Safety
N/A	109	0050	Osteoarthritis (OA): Function and Pain Assessment	Percentage of patient visits for patients aged 21 years and older with a diagnosis of osteoarthritis (OA) with assessment for function and pain.	Patient visits with assessment for level of function and pain documented.	All patient visits for patients aged 21 years and older with a diagnosis of OA.	Person and Caregiver-Centered Experience and Outcomes

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68v3	130	419	Documentation of Current Medications in the Medical Record	Percentage of specified visits for patients aged 18 years and older for which the eligible professional attests to documenting a list of current medications to the best of his/her knowledge and ability. This list must include ALL prescriptions, over-the-counters, herbals, and vitamin/mineral/dietary (nutritional) supplements AND must contain the medications' name, dosage, frequency and route of administration.	Eligible professional attests to documenting a list of current medications to the best of his/her knowledge and ability. This list must include ALL prescriptions, over-the-counters, herbals and vitamin/mineral/dietary (nutritional) supplements AND must contain the medications' name, dosages, frequency and route of administration.	All visits occurring during the 12 month reporting period for patients aged 18 years and older before the start of the measurement period.	Patient Safety
N/A	131	0420	Pain Assessment and Follow-Up	Percentage of visits for patients aged 18 years and older with documentation of a pain assessment using a standardized tool(s) on each visit AND documentation of a follow-up plan when pain is present.	Patient visits with a documented pain assessment using a standardized tool(s) AND documentation of a follow-up plan when pain is present.	All visits for patients aged 18 years and older.	Community/Population Health

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N/A	142	0051	Osteoarthritis (OA): Assessment for Use of Anti-Inflammatory or Analgesic Over-the-Counter (OTC) Medications	Percentage of visits for patients aged 21 years and older with a diagnosis of osteoarthritis (OA) with an assessment for use of anti-inflammatory or analgesic over-the-counter (OTC) medications.	Patient visits with assessment for use of anti-inflammatory or analgesic OTC medications documented.	All patient visits for patients aged 21 years and older with a diagnosis of OA.	Effective Clinical Care
	217	0422	Functional Deficit: Change in Risk-Adjusted Functional Status for Patients with Knee Impairments	Percentage of patients aged 18 or older that receive treatment for a functional deficit secondary to a diagnosis that affects the knee in which the change in their Risk-Adjusted Functional Status is measured.	Patients presented Focus On Therapeutic Outcome (FOTO) Functional Intake Survey for the Knee at admission and FOTO Functional Status Survey at discharge for the purpose of calculating the patient's Risk-adjusted Functional Status Change Residual Score.	All patients aged 18 years and older who receive a treatment episode for a functional deficit related to the knee.	Communication and Care Coordination
	226	0028	Preventive Care and Screening: Tobacco Use: Screening and Cessation Intervention	Percentage of patients aged 18 years and older who were screened for tobacco use one or more times within 24 months AND who received cessation counseling intervention if identified as a tobacco user.	Patients who were screened for tobacco use at least once within 24 months AND who received tobacco cessation counseling intervention if identified as a tobacco user.	All patients aged 18 years and older.	Community/Population Health

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N/A	350	N/A	Total Knee Replacement: Shared Decision-Making: Trial of Conservative (Non-surgical) Therapy	Percentage of patients regardless of age or gender undergoing a total knee replacement with documented shared decision-making with discussion of conservative (non-surgical) therapy prior to the procedure	Surgical patients with documentation of shared decision-making with discussion of conservative (non-surgical) therapy prior to the procedure	All surgical patients undergoing procedures for which total knee replacement is indicated in all patients	Communication and Care Coordination
N/A	351	N/A	Total Knee Replacement: Venous Thromboembolic and Cardiovascular Risk Evaluation	Percentage of patients regardless of age or gender undergoing a total knee replacement who are evaluated for the presence or absence of venous thromboembolic and cardiovascular risk factors within 30 days prior to the procedure including history of Deep Vein Thrombosis (DVT), Pulmonary Embolism (PE), Myocardial Infarction (MI), Arrhythmia and Stroke	Surgical patients who are evaluated for the presence or absence of venous thromboembolic and cardiovascular risk factors within 30 days prior to the procedure including history of DVT, PE, MI, Arrhythmia and Stroke	All surgical patients undergoing procedures for which total knee replacement is indicated in all patients	Patient Safety
N/A	352	N/A	Total Knee Replacement: Preoperative Antibiotic Infusion with Proximal Tourniquet	Percentage of patients regardless of age undergoing a total knee replacement who had the prophylactic antibiotic completely infused prior to the inflation of the proximal tourniquet	Surgical patients who had the prophylactic antibiotic completely infused prior to the inflation of the proximal tourniquet	All surgical patients undergoing procedures for which total knee replacement is indicated in all patients	Patient Safety

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N/A	353	N/A	Total Knee Replacement: Identification of Implanted Prosthesis in Operative Report	Percentage of patients regardless of age or gender undergoing a total knee replacement whose operative report identifies the prosthetic implant specifications including the prosthetic implant manufacturer, the brand name of the prosthetic implant and the size of the prosthetic implant	Surgical patients whose operative report identifies the prosthetic implant specifications including the prosthetic implant manufacturer, the brand name of the prosthetic implant and the size of the prosthetic implant	All surgical patients undergoing procedures for which total knee replacement is indicated in all patients	Patient Safety
	358	N/A	Patient-centered Surgical Risk Assessment and Communication	Percentage of patients who underwent a non-emergency surgery who had their personalized risks of postoperative complications assessed by their surgical team prior to surgery using a clinical data-based, patient-specific risk calculator and who received personal discussion of those risks with the surgeon.	Documentation of empirical, personalized risk assessment based on the patient's risk factors with a validated risk calculator using multi-institutional clinical data, the specific risk calculator used, and communication of risk assessment from risk calculator with the patient and/or family.	The total number of adult patients (age 18 and over) having had non-emergency surgery.	Person and Caregiver-Centered Experience and Outcomes

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66v2	375	N/A	Functional Status Assessment for Knee Replacement: Knee injury and Osteoarthritis Outcome Score (KOOS)	Percentage of patients aged 18 years and older with primary total knee arthroplasty (TKA) who completed baseline and follow-up (patient-reported) functional status assessments.	Patients with patient reported functional status assessment results not more than 180 days prior to the primary TKA procedure, and at least 60 days and not more than 180 days after TKA procedure.	Adults, aged 18 and older, with a primary TKA and who had an outpatient encounter not more than 180 days prior to procedure, and at least 60 days and not more than 180 days after TKA procedure.	Person and Caregiver-Centered Experience and Outcomes
66v2	375	N/A	Functional Status Assessment for Knee Replacement: Oxford Knee Score	Percentage of patients aged 18 years and older with primary total knee arthroplasty (TKA) who completed baseline and follow-up (patient-reported) functional status assessments.	Patients with patient reported functional status assessment results not more than 180 days prior to the primary TKA procedure, and at least 60 days and not more than 180 days after TKA procedure.	Adults, aged 18 and older, with a primary TKA and who had an outpatient encounter not more than 180 days prior to procedure, and at least 60 days and not more than 180 days after TKA procedure.	Person and Caregiver-Centered Experience and Outcomes



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66v2	375	N/A	Functional Status Assessment for Knee Replacement: Knee Society Knee Scoring System	Percentage of patients aged 18 years and older with primary total knee arthroplasty (TKA) who completed baseline and follow-up (patient-reported) functional status assessments.	Patients with patient reported functional status assessment results not more than 180 days prior to the primary TKA procedure, and at least 60 days and not more than 180 days after TKA procedure.	Adults, aged 18 and older, with a primary TKA and who had an outpatient encounter not more than 180 days prior to procedure, and at least 60 days and not more than 180 days after TKA procedure.	Person and Caregiver-Centered Experience and Outcomes
66v2	375	N/A	Functional Status Assessment for Knee Replacement: SF-36	Percentage of patients aged 18 years and older with primary total knee arthroplasty (TKA) who completed baseline and follow-up (patient-reported) functional status assessments.	Patients with patient reported functional status assessment results not more than 180 days prior to the primary TKA procedure, and at least 60 days and not more than 180 days after TKA procedure.	Adults, aged 18 and older, with a primary TKA and who had an outpatient encounter not more than 180 days prior to procedure, and at least 60 days and not more than 180 days after TKA procedure.	Person and Caregiver-Centered Experience and Outcomes

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66v2	375	N/A	Functional Status Assessment for Knee Replacement: Veterans RAND 12 Item Health Survey (VR-12)	Percentage of patients aged 18 years and older with primary total knee arthroplasty (TKA) who completed baseline and follow-up (patient-reported) functional status assessments.	Patients with patient reported functional status assessment results not more than 180 days prior to the primary TKA procedure, and at least 60 days and not more than 180 days after TKA procedure.	Adults, aged 18 and older, with a primary TKA and who had an outpatient encounter not more than 180 days prior to procedure, and at least 60 days and not more than 180 days after TKA procedure.	Person and Caregiver-Centered Experience and Outcomes
66v2	375	N/A	Functional Status Assessment for Knee Replacement: Patient Reported Outcomes Measurement Information System (PROMIS)-10	Percentage of patients aged 18 years and older with primary total knee arthroplasty (TKA) who completed baseline and follow-up (patient-reported) functional status assessments.	Patients with patient reported functional status assessment results not more than 180 days prior to the primary TKA procedure, and at least 60 days and not more than 180 days after TKA procedure.	Adults, aged 18 and older, with a primary TKA and who had an outpatient encounter not more than 180 days prior to procedure, and at least 60 days and not more than 180 days after TKA procedure.	Person and Caregiver-Centered Experience and Outcomes