2018 Individual Measures Applicable to Eligible OMS

Measure Number	Measure Description	Reporting Options/M ethods	Denominator Criteria (Eligible Cases)	Possible Denominator Codes Applicable to OMS *	Numerator	Possible Numerator Codes
Measure #21:	Perioperative Care: Selection of Prophylactic Antibiotic – First OR Second Generation Cephalosporin – National Quality Strategy Domain: Patient Safety 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.	Claims, Registry	All surgical patients aged 18 years and older undergoing procedures with the indications for a first OR second generation cephalosporin prophylactic antibiotic. Denominator Instructions: CPT Category I procedure codes billed by surgeons performing surgery on the same patient, submitted with modifier 62 (indicating two surgeons, i.e., dual procedures) will be included in the denominator population, therefore both surgeons will be fully accountable for the clinical action described in the measure.	Le Fort Fractures 21346, 21347, 21348, 21422, 21423, 21435, 21436 Mandibular Fractures 21454, 21461, 21462, 21465, 21470 Glossectomy 41130, 41135, 41140, 41145, 41150, 41153, 41155	Surgical patients who had an order for a first OR second generation cephalosporin for antimicrobial prophylaxis	Numerator Quality-Data Coding Options: Documentation of Order for First or Second Generation Cephalosporin for Antimicrobial Prophylaxis (written order, verbal order, or standing order/protocol) Performance Met: G9197: Documentation of order for first OR second generation cephalosporin for antimicrobial prophylaxis Note: G9197 is provided for antibiotic ordered or antibiotic given. Submit G9197 if a first or second generation cephalosporin was given for antimicrobial prophylaxis. OR Order for First or Second Generation Cephalosporin not ordered for Medical Reasons Denominator Exception: G9196: Documentation of medical reason(s) for not ordering a first OR second generation cephalosporin for antimicrobial prophylaxis (e.g., patients enrolled in clinical trials, patients with documented infection prior to surgical procedure of interest, patients who were receiving antibiotics more than 24 hours prior to surgery [except colon surgery patients taking oral prophylactic antibiotics], patients who were receiving antibiotics within 24 hours prior to arrival [except colon surgery patients taking oral prophylactic antibiotics], other medical reason(s)) OR

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			Denominator Criteria (Eligible Cases): Patients aged ≥ 18 years on date of encounter AND Patient procedure during the performance period (CPT): Listed below are surgical procedures with indications for first or second generation cephalosporin prophylactic antibiotic.			Order for First or Second Generation Cephalosporin not Ordered, Reason Not Given Performance Not Met: G9198: Order for first OR second generation cephalosporin for antimicrobial prophylaxis was not documented, reason not given
Measure #23:	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	Claims, Registry	Patients aged ≥ 18 years on date of encounter AND having one of the encounters listed in the next column to the right	Le Fort Fractures 21346, 21347, 21348, 21422, 21423, 21432, 21433, 21435, 21436 Mandibular Fractures 21454, 21461, 21462, 21465, 21470 Glossectomy 41130, 41135, 41140, 41145,	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	CPT II 4044F: Documentation that an order was given for venous thromboembolism (VTE) prophylaxis to be given within 24 hours prior to incision time or 24 hours after surgery end time OR 4044F with 1P: Documentation of medical reason(s) for patient not receiving any form of VTE prophylaxis (LMWH, LDUH, adjusted-dose warfarin, fondaparinux or mechanical prophylaxis) within 24 hours prior to incision time or 24 hours after surgery end time OR

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	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			41150, 41153, 41155		4044F with 8P: Order was not given for venous thromboembolism (VTE) prophylaxis to be given within 24 hours prior to incision time or 24 hours after surgery end time, reason not otherwise specified
Measure #46:	Medication Reconciliation Post- Discharge – National Quality Strategy Domain: Communication and Care Coordination DESCRIPTION: The percentage of discharges from any inpatient facility (e.g. hospital, skilled nursing facility, or rehabilitation facility) for patients 18 years of age and older seen within 30 days following discharge in the office by the physician, prescribing practitioner, registered nurse, or clinical pharmacist providing on-going care for whom the discharge medication list was reconciled	Claims, Registry	SUBMISSION CRITERIA 1: Patients 18-64 years of age on date of encounter SUBMISSION CRITERIA 2: Patients aged 65 years and older on date of encounter SUBMISSION CRITERIA 3: All Patients 18 years of age and older AND Patient encounter during the performance period (CPT or HCPCS) to the codes on the right column. AND Patient discharged from an inpatient facility (e.g., hospital, skilled	99201, 99202, 99203, 99204, 99205, 99212, 99213, 99214, 99215, 99324, 99325, 99326, 99327, 99328, 99334, 99335, 99341, 99342, 99343, 99344, 99345, 99347, 99348, 99349, 99350	Medication reconciliation conducted by a prescribing practitioner, clinical pharmacists or registered nurse on or within 30 days of discharge.	NUMERATOR NOTE: Medication reconciliation should be completed and documented on or within 30 days of discharge. If the patient has an eligible discharge but medication reconciliation is not performed and documented within 30 days, submit 1111F with 8P. Numerator Quality-Data Coding Options: Patient receiving Hospice Services, Patient Not Eligible: Denominator Exclusion: G9691: Patient had hospice services any time during the measurement period OR Documentation of Reconciliation of Discharge Medication with Current Medication List in the Medical Record Performance Met: CPT II 1111F: Discharge medications reconciled with the current medication list in outpatient medical record OR Discharge Medication not Reconciled with Current Medical Record, Reason Not Otherwise Specified

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	with the current medication list in the outpatient medical record. This measure is submitted as three rates stratified by age group: • Submission Criteria 1: 18-64 years of age • Submission Criteria 2: 65 years and older • Total Rate: All patients 18 years of age and older		nursing facility, or rehabilitation facility) within the last 30 days			Append a submission modifier (8P) to CPT Category II code 1111F to submit circumstances when the action described in the numerator is not performed and the reason is not otherwise specified. Performance Not Met: 1111F with 8P: Discharge medications not reconciled with the current medication list in outpatient medical record, reason not otherwise specified
Measure #47	Care Plan – National Quality Strategy Domain: Communication and Care Coordination	Claims, Registry	Patients aged ≥ 65 years on date of encounter AND Having one of the encounters (CPT or HCPCS codes) during the reporting period listed in the next column to the right	99201, 99202, 99203, 99204, 99205, 99212, 99213, 99214, 99215, 99218, 99219, 99220, 99221, 99222, 99223, 99231, 99232, 99234, 99235, 99236, 99291, 99304, 99305, 99306, 99307, 99308, 99309, 99310, 99324, 99325, 99326, 99327, 99328, 99334, 99335, 99341, 99342, 99343, 99344, 99345,	Patients who have an advance care plan or surrogate decision maker documented in the medical record or documentation in the medical record that an advance care plan was discussed but patient did not wish or was not able to name a surrogate decision maker or provide an	Advance Care Planning Discussed and Documented CPT II 1123F: Advance Care Planning discussed and documented; advance care plan or surrogate decision maker documented in the medical record OR CPT II 1124F: Advance Care Planning discussed and documented in the medical record; patient did not wish or was not able to name a surrogate decision maker or provide an advance care plan OR Advance Care Planning not Documented, Reason not Otherwise Specified Append a reporting modifier (8P) to CPT Category II code 1123F to report

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				99347, 99348, 99349, 99350, G0402	advance care plan	circumstances when the action described in the numerator is not performed and the reason is not otherwise specified. 1123F with 8P: Advance care planning not documented, reason not otherwise specified
Measure #130:	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/dietar y (nutritional) supplements AND must contain the medications' name, dosage, frequency and route of administration	Claims, Registry	Patients aged ≥ 18 years on date of encounter AND having one of the encounters (CPT or HCPCS codes) listed in the next column to the right	99201, 99202, 99203, 99204, 99205, 99212, 99213, 99214, 99215	Eligible professional or eligible clinician attests to documenting, updating or reviewing a patient's current medications using all immediate resources available on the date of encounter. This list must include ALL known prescriptions, over-the counters, herbals, and vitamin/mineral/di etary (nutritional) supplements AND must contain the medications' name, dosages, frequency and route of administration NUMERATOR NOTE: The eligible clinician	Numerator Quality-Data Coding Options: Current Medications Documented Performance Met: G8427: Eligible clinician attests to documenting in the medical record they obtained, updated, or reviewed the patient's current medications OR Current Medications not Documented, Patient not Eligible Denominator Exception: G8430: Eligible clinician attests to documenting in the medical record the patient is not eligible for a current list of medications being obtained, updated, or reviewed by the eligible clinician OR Current Medications with Name, Dosage, Frequency, or Route not Documented, Reason not Given Performance Not Met: G8428: Current list of medications not documented as obtained, updated, or reviewed by the eligible clinician, reason not given

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		etnods	Cases)	to OMS *	must document in the medical record they obtained, updated, or reviewed a medication list on the date of the encounter. Eligible clinicians submitting this measure may document medication information received from the patient, authorized representative(s), caregiver(s) or other available healthcare resources. By submitting the action described in this measure, the provider attests to having documented a list of current medications	
					utilizing all immediate resources available at the time of the encounter. G8427 should be submitted if the eligible clinician	

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					documented that the patient is not currently taking any medications	
Measure #131:	Pain Assessment and Follow-Up: Percentage of visits for patients aged 18 years and older with documentation of a pain assessment through discussion with the patient including the use of a standardized tool(s) on each visit AND documentation of a follow-up plan when pain is present	Claims, Registry	Patients aged ≥ 18 years on date of encounter AND having one of the encounters (CPT or HCPCS codes) listed in the next column to the right	92526, 96116, 96150, 99201, 99202, 99203, 99204, 99205, 99212, 99213, 99214, 99215	Patient visits with a documented pain assessment using a standardized tool(s) AND documentation of a follow-up plan when pain is present	G8730: Pain assessment documented as positive utilizing a standardized tool and a follow-up plan is documented OR G8731: Pain assessment documented as negative, no follow-up plan required OR G8442: Documentation that patient is not eligible for a pain assessment OR G8939: Pain assessment documented, follow-up plan not documented, patient not eligible/appropriate OR G8732: No documentation of pain assessment, reason not given OR G8509: Documentation of positive pain assessment; no documentation of a follow-up plan, reason not given

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Measure #137	Melanoma: Continuity of Care – Recall System – National Quality Strategy Domain: Communication and Care Coordination Percentage of patients, regardless of age, with a current diagnosis of melanoma or a history of melanoma whose information was entered, at least once within a 12 month period, into a recall system that includes: A target date for the next complete physical skin exam, AND A process to follow up with patients who either did not make an appointment within the specified timeframe or who missed a scheduled appointment	Registry	Diagnosis for melanoma or history of melanoma having one of diagnosis and patient encounter during the performance period. (CPT/(ICD-10-CM codes) listed to the right. WITHOUT Telehealth Modifier: GQ, GT, 95, POS 02	99201, 99202, 99203, 99204, 99205, 99212, 99213, 99214, 99215, 99241*, 99242*, 99243*, 99244*, 99245 C43.0, C43.10, C43.11, C43.12, C43.20, C43.21, C43.22, C43.30, C43.31, D03.30, D03.39, Z85.820	Patients whose information is entered, at least once within a 12 month period, into a recall system that includes: A target date for the next complete physical exam AND A process to follow up with patients who either did not make an appointment within the specified timeframe or who missed a scheduled appointment	Numerator Options: Performance Met: Patient information entered into a recall system that includes: target date for the next exam specified AND a process to follow up with patients regarding missed or unscheduled appointments (7010F) OR Denominator Exception: Documentation of system reason(s) for not entering patient's information into a recall system (e.g., melanoma being monitored by another physician provider) (7010F with 3P) OR Performance Not Met: Recall system not utilized, reason not otherwise specified (7010F with 8P)

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Measure #138	Melanoma: Coordination of Care – National Quality Strategy Domain: Communication and Care Coordination Percentage of patient visits, regardless of age, with a new occurrence of melanoma that have a treatment plan documented in the chart that was communicated to the physician(s) providing continuing care within one month of diagnosis THERE ARE TWO SUBMISSION CRITERIA FOR THIS MEASURE: 1) All visits for patients, regardless of age, diagnosed with a new occurrence of melanoma during excision of malignant lesion OR 2) All visits for patients, regardless of age, diagnosed with a new occurrence of melanoma during excision of malignant lesion OR 2) All visits for patients, regardless of age, diagnosed with a new occurrence of	Registry	Denominator Criteria (Eligible Cases) 1: Diagnosis for melanoma (ICD- 10-CM): C43.10, C43.11, C43.20, C43.30, C43.31, D03.30, D03.39 AND Patient encounter for excision of malignant melanoma (CPT): 11640, 11641, 11642, 11643, 11644, 11646, , 14040, 14041, 14060, 14061, 14301, 17311 All visits for patients, regardless of age, diagnosed with a new occurrence of melanoma Denominator Criteria (Eligible Cases) 2: Diagnosis for melanoma (ICD- 10-CM): C43.10, C43.11, C43.20, C43.30, C43.31, D03.30, D03.39 AND Patient encounter during the	(Eligible cases 1) C43.10, C43.11, C43.20, C43.30, C43.31, D03.30, D03.39 11640, 11641, 11642, 11643, 11644, 11646, 14040, 14041, 14060, 14061, 14301, 17311 (Eligible cases 2) C43.10, C43.11, C43.20, C43.30, C43.31, D03.30, D03.39 99201, 99202, 99203, 99204, 99205, 99212, 99213, 99214, 99215, 99214, 99215, 99241*, 99242*, 99243*, 99244*, 99245*	NUMERATOR (SUBMISSION CRITERIA): Patient visits with a treatment plan documented in the chart that was communicated to the physician(s) providing continuing care within one month of diagnosis.	NUMERATOR (SUBMISSION CRITERIA 1 & 2): Patient visits with a treatment plan documented in the chart that was communicated to the physician(s) providing continuing care within one month of diagnosis Performance Met: Treatment plan communicated to provider(s) managing continuing care within 1 month of diagnosis (5050F OR Denominator Exception: Documentation of patient reason(s) for not communicating treatment plan to the Primary Care Physician(s) (PCP) (s) (e.g., patient asks that treatment plan not be communicated to the physician(s) providing continuing care) (5050F with 2P) OR Denominator Exception: Documentation of system reason(s) for not communicating treatment plan to the PCP(s) (e.g., patient does not have a primary care physician or referring physician) (5050F with 3P) OR Performance Not Met: Treatment plan not communicated, reason not otherwise specified (5050F with 8P)

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	melanoma evaluated in an outpatient setting		performance period (CPT): 99201, 99202, 99203, 99204, 99205, 99212, 99213, 99214, 99215, 99241*, 99242*, 99243*, 99244*, 99245* WITHOUT Telehealth Modifier: GQ, GT, 95, POS 02 (CPT/(ICD-10- CM codes) listed to the right.			
Measure #224	Melanoma: Overutilization of Imaging Studies in Melanoma – National Quality Strategy Domain: Efficiency and Cost Reduction Percentage of patients, regardless of age, with a current diagnosis of Stage 0 through IIC melanoma or a history of melanoma of any stage, without signs or symptoms	Registry	THERE ARE TWO SUBMISSION CRITERIA FOR THIS MEASURE: 1) Patients with a diagnosis of Stage 0 through IIC melanoma without signs or symptoms suggesting systemic spread OR 2) Patients with a history of any stage melanoma without signs or symptoms	Diagnosis for melanoma (ICD-10-CM): C43.0, C43.11, C43.12, C43.20, C43.21, C43.22, C43.30, C43.31, C43.39, C43.4, C43.51, C43.52, C43.60, C43.61, C43.62, C43.70, C43.71, C43.72, C43.8, C43.9, D03.0, D03.10, D03.11, D03.12, D03.20, D03.21, D03.22, D03.30, D03.39, D03.4, D03.51, D03.52, D03.59, D03.60, D03.61,	NUMERATOR SUBMISSION CRITERIA: Patients for whom no diagnostic imaging studies were ordered	NUMERATOR SUBMISSION CRITERIA 1: Performance Met: None of the following diagnostic imaging studies ordered: chest x-ray, CT, Ultrasound, MRI, PET, or nuclear medicine scans (3320F) OR Denominator Exception: Documentation of medical reason(s) for ordering diagnostic imaging studies (e.g., patient has comorbid condition that warrants imaging, other medical reasons) (3319F with 1P) OR Denominator Exception: Documentation of system reason(s) for ordering diagnostic imaging studies (e.g., requirement for clinical trial enrollment, ordered by another provider, other system reasons) (3319F with 3P) OR Performance Not Met: One of the following diagnostic imaging studies ordered; chest x-ray, CT, Ultrasound, MRI, PET, or nuclear medicine scans (3319F)

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	suggesting systemic spread, seen for an office visit during the one-year measurement period, for whom no diagnostic imaging studies were ordered		suggesting systemic spread	D03.62, D03.70, D03.71, D03.72, D03.8, D03.9 AND Patient encounter during the performance period (CPT): 99201, 99202, 99203, 99204, 99205, 99212, 99213, 99214, 99215, 99241*, 99242*, 99243*, 99244*, 99245* WITHOUT Telehealth Modifier: GQ, GT, 95, POS 02 AND AJCC Melanoma Cancer Stage 0 through IIC Melanoma: G8944 AND Absence of signs of melanoma (cough, dyspnea, tenderness, localized neurologic signs such as weakness, jaundice, or any other sign suggesting systemic spread) or absence of symptoms of		
				melanoma (pain, paresthesia, or any other		

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				symptom suggesting the possibility of systemic spread of melanoma): G8749		
Measure # 226	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	Claims, Registry	Patients aged ≥ 18 years on date of encounter AND having one of the encounters (CPT or HCPCS codes) listed in the next column to the right	99201, 99202, 99203, 99204, 99205, 99212, 99213, 99214, 99215	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	Patient Screened for Tobacco Use CPT II 4004F: Patient screened for tobacco use AND received tobacco cessation counseling, if identified as a tobacco user OR Patient Screened for Tobacco Use and Identified as a Non-User of Tobacco CPT II 1036F: Current tobacco non-user OR Tobacco Screening not Performed for Medical Reasons Append a modifier (1P) to CPT Category II code 4004F to report documented circumstances that appropriately exclude patients from the denominator 4004F with 1P: Documentation of medical reason(s) for not screening for tobacco use (e.g., limited life expectancy) OR Tobacco Screening not Performed Reason Not Specified Append a reporting modifier (8P) to CPT Category II code 4004F to report circumstances when the action described in the numerator is not performed and the reason is not otherwise specified.

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Measure #238	Use of High-Risk Medications in the Elderly Percentage of patients 66 years of age and older who were ordered high- risk medications. Two rates are reported: a. Percentage of patients who were ordered at least one high-risk medication b. Percentage of patients who were ordered at least two different high-risk medications.	Registry	Patients aged ≥ 65 years on date of encounter AND having one of the encounters listed in the next column to the right	99201, 99202, 99203, 99204, 99205, 99212, 99213, 99214, 99215	Percentage of patients who were ordered at least one or two high-risk medication during the measurement period	4004F with 8P: Tobacco Screening not performed, reason not otherwise specified Drug Classifications applying to the OMS. Pain medication, skeletal relaxants: Carisoprodol Metaxalone Chlorzoxazone Methocarbamol Cyclobenzaprine Orphenadrine Pain medications, other: Indomethacin Ketorolac, include Meperidine parenteral Pentazocine
Measure #265	Biopsy Follow-Up: Percentage of new patients whose biopsy results have been reviewed and communicated to the primary care/referring physician and patient by the performing physician	Registry	All patients undergoing a biopsy	11100, 20200, 20205, 20206, 20220, 20225, 20240, 20245, 30100, 31050, 31051, 38500, 38505, 40490, 40808, 41100, 41108, 42100, 42400, 42405, 42800, 42802, 42804, 42806, 64795, 99201, 99202, 99203, 99204, 99205	Patients whose biopsy results have been reviewed and communicated to the primary care/referring physician and the patient by the physician performing the biopsy. The physician performing the biopsy must also acknowledge and/or document	G8883: Biopsy results reviewed, communicated, tracked and documented OR G88884: Clinician documented reason that patient's biopsy results were not reviewed, [e.g., patient asks that biopsy results not be communicated to the primary care/referring physician, patient does not have a primary care/referring physician or is a self-referred patient] OR G8885: Biopsy results NOT reviewed, communicated, tracked or documented

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					the communication in a biopsy tracking log and document in the patient's medical record.	
Measure #279	Sleep Apnea: Assessment of Adherence to Positive Airway Pressure Therapy – National Quality Strategy Domain: Effective Clinical Care	Registry	Patients aged ≥ 18 years on date of encounter AND Diagnosis for sleep apnea (ICD-10-CM): G47.30, G47.33 WITHOUT Telehealth Modifier: GQ, GT, 95, POS 02 AND Positive airway pressure therapy was prescribed: G8852 AND having one of the encounters (CPT or HCPCS codes) listed in the next column to the right	99201, 99202, 99203, 99204, 99205, 99212, 99213, 99214, 99215	Patient visits with documentation that adherence to positive airway pressure therapy was objectively measured	Patient visits with documentation that adherence to positive airway pressure therapy was objectively measured Numerator options: Performance Met: Objective measurement of adherence to positive airway pressure therapy, documented (G8851) OR Denominator Exception: Documentation of reason(s) for not objectively measuring adherence to positive airway pressure therapy (e.g., patient didn't bring data from continuous positive airway pressure [CPAP], therapy not yet initiated not available on machine) (G8854) OR Performance not met: Objective measurement of adherence to positive airway pressure therapy not performed, reason not given (G8855)
Measure #317	Preventive Care and Screening: Screening for High Blood Pressure and Follow- up Documented	Claims, Registry	Patients aged ≥ 18 years on date of encounter AND	99201, 99202, 99203, 99204, 99205, 99212, 99213, 99214, 99215, 99218,	Patients who were screened for high blood pressure AND have a	*Screening for High Blood Pressure not Documented, Patient not Eligible Denominator Exclusion: G9744: Patient not eligible due to active diagnosis of hypertension

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	Percentage of patients aged 18 years and older seen during the reporting period who were screened for high blood pressure AND a recommended follow-up plan is documented based on the current blood pressure (BP) reading as indicated		having one of the encounters (CPT or HCPCS codes) listed in the next column to the right WITHOUT Telehealth modifier: GQ, GT, 95, POS 02	99219, 99220, 99224, 99225, 99226, 99234, 99235, 99286, 99281, 99285, D7140, D7210, G0101, G0402, G0438, G0429	recommended follow-up plan documented, as indicated if the blood pressure is pre-hypertensive or hypertensive or hypertensing interval for a normal BP reading is every 2 years, to meet the intent of this measure, BP screening and follow-up must be performed once per performance period. For patients with Normal blood pressure, a follow-up plan is not required. If the blood pressure is pre-hypertensive (SBP > 120 and <139 OR DBP >80 and <89) at a Primary Care Provider (PCP) encounter follow up as directed by the PCP would meet the intent of	G8783: Normal blood pressure reading documented, follow-up not required OR G8950: Pre-Hypertensive or Hypertensive blood pressure reading documented, AND the indicated follow-up is documented OR **Screening or Follow-Up for High Blood Pressure not Completed, Documented Reason Denominator Exception: G9745: Documented reason for not screening or recommending a follow-up for high blood pressure OR G8785: Blood pressure reading not documented, reason not given OR G8952: Pre-Hypertensive or Hypertensive blood pressure reading documented, indicated follow-up not documented, reason not given

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Measure #331	Adult Sinusitis: Antibiotic Prescribed for Acute Sinusitis (Overuse) Percentage of patients, aged 18 years and older, with a diagnosis of acute sinusitis who were prescribed an antibiotic within 10 days after onset of symptoms	Registry	Denominator Criteria (Eligible Cases): Patients aged ≥ 18 years on date of encounter AND Diagnosis for acute sinusitis (ICD-10-CM): J01.00, J01.01, J01.10, J01.11, J01.20, J01.21, J01.30, J01.31, J01.40, J01.41, J01.80, J01.90 AND having one of the encounters (CPT code) listed in the next column to the right	99201, 99202, 99203, 99204, 99205, 99212, 99213, 99214, 99215, 99281, 99282, 99283, 99284, 99285 <u>WITHOUT</u> Telehealth Modifier: GQ, GT, 95, POS 02	the measure (G8783). Patients prescribed any antibiotic within 10 days after onset of symptoms	Numerator Options: Performance Met: Antibiotic regimen prescribed within 10 days after onset of symptoms (G9286) OR Denominator Exception: Antibiotic regimen prescribed within 10 days after onset of symptoms for documented medical reason (G9505) OR Performance Not Met: Antibiotic regimen not prescribed within 10 days after onset of symptoms (G9287)
Measure #332	Adult Sinusitis: Appropriate Choice of Antibiotic: Amoxicillin With or Without Clavulanate Prescribed for Patients with Acute Bacterial Sinusitis Percentage of patients aged 18 years and older with a diagnosis of acute	Registry	Denominator Criteria (Eligible Cases): Patients aged ≥ 18 years on date of encounter AND Diagnosis for acute sinusitis (ICD-10-CM): J01.00, J01.01, J01.10, J01.11, J01.20, J01.21, J01.30, J01.31,	99201, 99202, 99203, 99204, 99205, 99212, 99213, 99214, 99215, 99281, 99282, 99283, 99284, 99285 WITHOUT Telehealth Modifier: GQ, GT, 95, POS 02 AND	Patients who were prescribed amoxicillin, with or without clavulanate, as a first line antibiotic at the time of diagnosis	G9315: Amoxicillin, with or without clavulanate, prescribed as a first line antibiotic at the time of diagnosis OR G9313: Amoxicillin, with or without clavulanate, not prescribed as first line antibiotic at the time of diagnosis for documented reason (e.g., cystic fibrosis, immotile cilia disorders, ciliary dyskinesia, immune deficiency, prior history of sinus surgery within the past 12 months, and anatomic abnormalities, such as deviated

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	bacterial sinusitis that were prescribed amoxicillin, with or without clavulanate, as a first line antibiotic at the time of diagnosis		J01.40, J01.41, J01.80, J01.90 AND having one of the encounters (CPT code) listed in the next column to the right	Sinusitis caused by, or presumed to be caused by, bacterial infection: G9364 AND Antibiotic regimen prescribed: G9498		nasal septum, resistant organisms, allergy to medication, recurrent sinusitis, chronic sinusitis, or other reasons) OR G9314: Amoxicillin, with or without clavulanate, not prescribed as first line antibiotic at the time of diagnosis, reason not given
Measure #333	Adult Sinusitis: Computed Tomography (CT) for Acute Sinusitis (Overuse): Percentage of patients aged 18 years and older with a diagnosis of acute sinusitis who had a computerized tomography (CT) scan of the paranasal sinuses ordered at the time of diagnosis or received within 28 days after date of diagnosis	Registry	Patients aged ≥ 18 years on date of encounter AND Diagnosis for acute sinusitis Diagnosis for acute sinusitis (ICD-10-CM) for J01.00, J01.10, J01.20, J01.30, J01.40, J01.80, J01.90 AND having one of the encounters listed in the next column to the right	99201, 99202, 99204, 99205, 99212, 99213, 99214, 99215, 99281, 99282, 99283, 99284, 99285 WITHOUT Telehealth Modifier: GQ, GT, 95, POS 02	Patients who had a computerized tomography (CT) scan of the paranasal sinuses ordered at the time of diagnosis or received within 28 days after date of diagnosis	G9349: CT scan of the paranasal sinuses ordered at the time of diagnosis or received within 28 days after date of diagnosis OR G9348: CT scan of the paranasal sinuses ordered at the time of diagnosis for documented reasons (e.g., persons with sinusitis symptoms lasting at least 7 to 10 days, antibiotic resistance, immunocompromised, recurrent sinusitis, acute frontal sinusitis, acute sphenoid sinusitis, periorbital cellulitis, or other medical) OR G9350: CT scan of the paranasal sinuses not ordered at the time of diagnosis or received within 28 days after date of diagnosis
Measure #334	Adult Sinusitis: More than one Computerized	Registry	Patients aged ≥ 18 years on date of encounter	99201, 99202, 99204, 99205, 99212, 99213,	Patients who had more than one CT scan of the	G9352: More than one CT scan of the paranasal sinuses ordered or received

Measure Number	Measure Description	Reporting Options/M ethods	Denominator Criteria (Eligible Cases)	Possible Denominator Codes Applicable to OMS *	Numerator	Possible Numerator Codes
	Tomography (CT) Scan Within 90 Days for Chronic Sinusitis (Overuse): Percentage of patients aged 18 years and older with a diagnosis of chronic sinusitis who had more than one CT scan of the paranasal sinuses ordered or received within 90 days after date of diagnosis		AND Diagnosis for acute sinusitis Diagnosis for acute sinusitis (ICD-10-CM) J01.00, J01.10, J01.20, J01.30, J01.40, J01.80, J01.90 AND having one of the encounters listed in the next column to the right	99214, 99215, 99281, 99282, 99283, 99284, 99285 WITHOUT Telehealth Modifier: GQ, GT, 95, POS 02	paranasal sinuses ordered or received within 90 days after date of diagnosis	within 90 days after the date of diagnosis, reason not given OR G9353: More than one CT scan of the paranasal sinuses ordered or received within 90 days after the date of diagnosis for documented reasons (e.g., patients with complications, second CT obtained prior to surgery, other medical reasons) OR G9354: One CT scan or no CT scan of the paranasal sinuses ordered within 90 days after the date of diagnosis
Measure #342	Pain Brought Under Control Within 48 Hours: Patients aged 18 and older who report being uncomfortable because of pain at the initial assessment (after admission to palliative care services) who report pain was brought to a comfortable level within 48 hours	Registry	Patients aged ≥ 18 years AND having one of the encounters listed in the next column to the right AND Patient able to communicate and understand the language of	99324, 99325, 99326, 99327, 99328	Patients whose pain was brought to a comfortable level within 48 hours of initial assessment (after admission to palliative care services)	G9250: Documentation of patient pain brought to a comfortable level within 48 hours from initial assessment OR G9251: Documentation of patient with pain not brought to a comfortable level within 48 hours from initial assessment

Measure Number	Measure Description	Reporting Options/M ethods	Denominator Criteria (Eligible Cases)	Possible Denominator Codes Applicable to OMS *	Numerator	Possible Numerator Codes
			the person asking AND Patient self-reported uncomfortable due to pain at the initial assessment			
Measure #355	Unplanned Reoperation within the 30 day Postoperative Period Percentage of patients aged 18 years or older who had any unplanned reoperation within the 30 day postoperative period	Registry	All patients aged 18 years and older AND Patient procedure during the performance period having one of the encounters listed in the next column to the right	20200, 20205, 21552, 21554, 21555, 21556.	Unplanned return to the operating room for a surgical procedure, for any reason, within 30 days of the principal operative procedure	Unplanned return to the operating room for a surgical procedure, for complications of the principal operative procedure, within 30 days of the principal operative procedure (G9308) OR Unplanned return to the operating room for a surgical procedure, for complications of the principal operative procedure, within 30 days of the principal operative procedure (G9308)
Measure #358	Patient-Centered Surgical Risk Assessment and Communication – National Quality Strategy Domain: Person and Caregiver-Centered Experience and Outcomes Percentage of patients who	Registry	Patients aged ≥ 18 years on date of encounter AND Patient encounter during the performance period (Multiple CPT codes apply)	10121, 10140, 10160, 10180, 11000, 11001, 11010, 11011, 11042, 11043, 11044,11420, 11421, 14301, 15040, 15155, 15220, 15240, 15260, 15574, 15576, 21011, 21012, 21013, 21014, 21015,	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,	Numerator Options: Performance Met: Documentation of patient-specific risk assessment with a risk calculator based on multi-institutional clinical data, the specific risk calculator used, and communication of risk assessment from risk calculator with the patient or family (G9316) Performance Not Met: Documentation of patient-specific risk assessment with a risk calculator based on multi-institutional

Measure Number	Measure Description	Reporting Options/M ethods	Denominator Criteria (Eligible Cases)	Possible Denominator Codes Applicable to OMS *	Numerator	Possible Numerator Codes
	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		AND NOT DENOMINATOR EXCLUSION: Emergency surgery: G9752	21016, 21025, 21026, 21034, 21040, 21044, 21045, 21046, 21047, 21048, 21049, 21139, 21154, 21235, 21299, 21360, 40800, 40801, 40810, 40812, 40814, 40816, 41000, 41005, 41006, 41007, 41008, 41007, 41018, 41110, 41112, 41113, 41114, 41116, 41120, 41130, 41145, 41150, 41153, 41150,	and communication of risk assessment from risk calculator with the patient and/or family	clinical data, the specific risk calculator used, and communication of risk assessment from risk calculator with the patient or family not completed (G9317)
Measure #359	Optimizing Patient Exposure to Ionizing Radiation: Utilization of a Standardized Nomenclature for Computed Tomography (CT) Imaging Description - National Quality Strategy Domain: Communication and Care Coordination Percentage of computed tomography (CT) imaging reports for all	Registry	All patients regardless of age AND Patient procedure during the performance period (based on CPT codes to the right)	70486, 70487, 70488, 70490, 70491, 70492, 70496, 70498	CT imaging reports with the imaging study named according to a standardized nomenclature and the standardized nomenclature is used in institution's computer systems	Numerator Options: Performance Met: Imaging study named according to standardized nomenclature (G9318) OR Performance Not Met: Imaging study not named according to standardized nomenclature, reason not given (G9319)

Measure Number	Measure Description	Reporting Options/M ethods	Denominator Criteria (Eligible Cases)	Possible Denominator Codes Applicable to OMS *	Numerator	Possible Numerator Codes
	patients, regardless of age, with the imaging study named according to a standardized nomenclature and the standardized nomenclature is used in institution's computer systems					
Measure #361	Optimizing Patient Exposure to Ionizing Radiation: Reporting to a Radiation Dose Index Registry – National Quality Strategy Domain: Patient Safety Percentage of total computed tomography (CT) studies performed for all patients, regardless of age, that are submitted to a radiation dose index registry that is capable of collecting at a minimum selected data elements	Registry	All patients regardless of age AND Patient procedure during the performance period (based on CPT codes to the right)	70486, 70487, 70488, 70490, 70491, 70492, 70496, 70498	CT studies performed that are reported to a radiation dose index registry that is capable of collecting at a minimum all of the following data elements: Manufacturer Study description Manufacturer's model name Patient's weight Patient's size Patient's sex Patient's age Exposure time	Numerator Options: Performance Met: CT studies performed reported to a radiation dose index registry that is capable of collecting at a minimum all necessary data elements (G9327) OR Performance Not Met: CT studies performed not reported to a radiation dose index registry that is capable of collecting at a minimum all necessary data elements, reason not given (G9326)

Measure Number	Measure Description	Reporting Options/M ethods	Denominator Criteria (Eligible Cases)	Possible Denominator Codes Applicable to OMS *	Numerator	Possible Numerator Codes
					X-Ray tube current Kilovoltage (kV) Mean Volume Computed tomography dose index (CTDIvol) Dose-length product (DLP)	
Measure #362	Optimizing Patient Exposure to Ionizing Radiation: Computed Tomography (CT) Images Available for Patient Follow-up and Comparison Purposes – National Quality Strategy Domain: Communication and Care Coordination Percentage of final reports for computed tomography (CT) studies performed for all patients, regardless of age, which document that Digital Imaging and Communications in Medicine (DICOM) format image data are available to non-	Registry	All patients regardless of age AND Patient procedure during the performance period (based on CPT codes to the right)	70486, 70487, 70488, 70490, 70491, 70492, 70496, 70498	Final reports for CT studies which document that DICOM format image data are available to non-affiliated external healthcare facilities or entities on a secure, media-free, reciprocally searchable basis with patient authorization for at least a 12-month period after the study	Performance Met: Final report documented that DICOM format image data available to nonaffiliated external healthcare facilities or entities on a secure, media free, reciprocally searchable basis with patient authorization for at least a 12-month period after the study (G9340) OR Performance Not Met: DICOM format image data available to non-affiliated external healthcare facilities or entities on a secure, media free, reciprocally searchable basis with patient authorization for at least a 12-month period after the study not documented in final report, reason not given (G9329)

Measure Number	Measure Description	Reporting Options/M ethods	Denominator Criteria (Eligible Cases)	Possible Denominator Codes Applicable to OMS *	Numerator	Possible Numerator Codes
	affiliated external healthcare facilities or entities on a secure, media free, reciprocally searchable basis with patient authorization for at least a 12-month period after the study					
Measure #363	Optimizing Patient Exposure to Ionizing Radiation: Search for Prior Computed Tomography (CT) Studies Through a Secure, Authorized, Media-Free, Shared Archive – National Quality Strategy Domain: Communication and Care Coordination Percentage of final reports of computed tomography (CT) studies performed for all patients, regardless of age, which document that a search for Digital Imaging and Communications in Medicine (DICOM) format images was conducted for prior patient CT imaging	Registry	All patients regardless of age AND Patient procedure during the performance period (based on CPT codes to the right)	70486, 70487, 70488, 70490, 70491, 70492, 70496, 70498	Final reports of CT studies, which document that a search for DICOM format images was conducted for prior patient CT imaging studies completed at non-affiliated external healthcare facilities or entities within the past 12-months and are available through a secure, authorized, media-free, shared archive prior to an imaging study being performed	Numerator Options: Performance Met: Search conducted for prior patient CT studies completed at non-affiliated external healthcare facilities or entities within the past 12-months and are available through a secure, authorized, media-free, shared archive prior to an imaging study being performed (G9341) OR Denominator Exception: Due to system reasons search not conducted for DICOM format images for prior patient CT imaging studies completed at non-affiliated external healthcare facilities or entities within the past 12 months that are available through a secure, authorized, media-free, shared archive (e.g., non-affiliated external healthcare facilities or entities does not have archival abilities through a shared archival system) (G9344) OR Denominator Exception: Documentation of medical reason for not conducting a search for DICOM format images for prior patient CT imaging studies completed at non-affiliated external

Measure Number	Measure Description	Reporting Options/M ethods	Denominator Criteria (Eligible Cases)	Possible Denominator Codes Applicable to OMS *	Numerator	Possible Numerator Codes
	studies completed at non-affiliated external healthcare facilities or entities within the past 12-months and are available through a secure, authorized, media-free, shared archive prior to an imaging study being performed					healthcare facilities or entities within the past 12 months that are available through a secure, authorized, mediafree, shared archive (e.g., trauma, acute myocardial infarction, stroke, aortic aneurysm where time is of the essence) (G9753) OR Performance Not Met: Search not conducted prior to an imaging study being performed for prior patient CT studies completed at non-affiliated external healthcare facilities or entities within the past 12-months and are available through a secure, authorized, media-free, shared archive, reason not given (G9342)
Measure #374	Closing the Referral Loop: Receipt of Specialist Report – National Quality Strategy Domain: Effective Communication and Care Coordination Percentage of patients with referrals, regardless of age, for which the referring provider receives a report from the provider to whom the patient was referred	Registry	Patients regardless of age on the date of the encounter AND Patient encounter during the performance period (based on CPT codes to the right) WITHOUT Telehealth Modifier: GQ, GT, 95, POS 02 AND Patient was referred to another provider or	99201, 99202, 99203, 99204, 99205, 99212, 99213, 99214, 99215	Number of patients with a referral, for which the referring provider received a report from the provider to whom the patient was referred	Numerator Options: Performance Met: Provider who referred the patient to another provider received a report from the provider to whom the patient was referred G9969 OR Performance Not Met: Provider who referred the patient to another provider did not receive a report from the provider to whom the patient was referred G9970

Measure Number	Measure Description	Reporting Options/M ethods	Denominator Criteria (Eligible Cases)	Possible Denominator Codes Applicable to OMS *	Numerator	Possible Numerator Codes
			specialist during the performance period: G9968			

Measure Number	Measure Description	Reporting Options/M ethods	Denominator Criteria (Eligible Cases)	Possible Denominator Codes Applicable to OMS *	Numerator	Possible Numerator Codes
Measure #397	Melanoma Reporting - National Quality Strategy Domain: Communication and Care Coordination	Claims	Patients ≥ 18 years of age on date of encounter AND Diagnosis for malignant cutaneous melanoma (ICD- 10-CM): C43.0, C43.20, C43.21, C43.22, C43.30, C43.31, C43.39, C43.4, C43.51, C43.52, C43.59, C43.60, C43.61, C43.62, C43.70, C43.71, C43.72, C43.8, C43.9 AND Patient procedure during performance period (CPT): 88305		Pathology reports for primary malignant cutaneous melanoma that include the pT category and a statement on thickness and ulceration and for pT1, mitotic rate	Numerator Quality-Data Coding Options: If Patient is not Eligible for this Measure because the Specimen is not of Cutaneous Origin Denominator Exclusion: G9430: Specimen site other than anatomic cutaneous location OR Pathology reports that include the pT category and a statement on thickness and ulceration and for pT1, mitotic rate. Performance Met: G9428: Pathology report includes the pT Category and a statement on thickness and ulceration and for pT1, mitotic rate. OR Pathology Reports that does not Include the pT Category and a Statement on Thickness and Ulceration and for pT1, mitotic rate, not documented for Medical Reasons Denominator Exception: G9429: Documentation of medical reason(s) for not including pT Category and a statement on thickness and ulceration and for pT1, mitotic rate (e.g., negative skin biopsies in a patient with a history of melanoma or other documented medical reasons) OR

Measure Number	Measure Description	Reporting Options/M ethods	Denominator Criteria (Eligible Cases)	Possible Denominator Codes Applicable to OMS *	Numerator	Possible Numerator Codes
						Pathology Reports that does not include the pT category and a statement on thickness and ulceration and for pT1, mitotic rate, reason not given. Performance NOT met: G9431: Pathology report does not include the pT category and a statement on thickness and ulceration and for pT1, mitotic rate.
Measure #402	Tobacco Use and Help with Quitting Among Adolescents – National Quality Strategy Domain: Community / Population Health The percentage of adolescents 12 to 20 years of age with a primary care visit during the measurement year for whom tobacco use status was documented and received help with quitting if identified as a tobacco user	Registry	Patients aged 12 to 20 years on the date of encounter AND having one of the encounters (CPT or HCPCS codes) listed in the next column to the right	99201, 99202, 99203, 99204, 99205, 99212, 99213, 99214, 99215, 99406, 99407, G0438, G0439	Patients who were screened for tobacco use at least once within 18 months (during the measurement period or the six months prior to the measurement period) AND who received tobacco cessation counseling intervention if identified as a tobacco user	G9458: Patient documented as tobacco user AND received tobacco cessation intervention (must include at least one of the following: advice given to quit smoking or tobacco use, counseling on the benefits of quitting smoking or tobacco use, assistance with or referral to external smoking or tobacco cessation support programs, or current enrollment in smoking or tobacco use cessation program) if identified as a tobacco user OR G9459: Currently a tobacco non-user OR

Measure Number	Measure Description	Reporting Options/M ethods	Denominator Criteria (Eligible Cases)	Possible Denominator Codes Applicable to OMS *	Numerator	Possible Numerator Codes
Measure	Anesthesiology	Registry	Patients aged ≥	00100-00210	Current cigarette	G9460: Tobacco assessment OR tobacco cessation intervention not performed, reason not otherwise specified G9644: Patients who abstained from
#404	Smoking Abstinence: National Quality Strategy Domain: Effective Clinical Care The percentage of current smokers who abstain from cigarettes prior to anesthesia on the day of elective surgery or procedure	regiony	18 years AND having one of the encounters (CPT or HCPCS codes) listed in the next column to the right AND Current cigarette smokers (G9642) AND Elective Surgery (G9643) AND Seen preoperatively by anesthesiologist or proxy prior to day of surgery		smokers and who abstained from smoking prior to anesthesia on the day of surgery or procedure.	smoking prior to anesthesia on the day of surgery or procedure OR G9645: Patients who did not abstain from smoking prior to anesthesia on the day of surgery or procedure
Measure #408	Opioid Therapy Follow-up Evaluation – National Quality Strategy Domain:	Registry	(G9497) Patients aged ≥ 18 years on date of encounter AND Patient encounter	99201, 99202, 99203, 99204, 99205, 99212, 99213, 99214, 99215	Patients who had a follow-up evaluation conducted at least every three	Numerator Options: Performance Met: Patients who had a follow-up evaluation conducted at least every three months during opioid therapy (G9562) OR

Measure Number	Measure Description	Reporting Options/M ethods	Denominator Criteria (Eligible Cases)	Possible Denominator Codes Applicable to OMS *	Numerator	Possible Numerator Codes
	Effective Clinical Care All patients 18 and older prescribed opiates for longer than six weeks duration who had a follow-up evaluation conducted at least every three months during Opioid Therapy documented in the medical record		during the performance period (CPT codes in the next column): WITHOUT Telehealth Modifier: GQ, GT, 95, POS 02 AND Patients prescribed opiates for longer than six weeks: G9561		months during opioid therapy	Performance Not Met: Patients who did not have a follow-up evaluation conducted at least every three months during opioid therapy (G9563)
Measure #412	Documentation of Signed Opioid Treatment Agreement— National Quality Strategy Domain: Effective Clinical Care All patients 18 and older prescribed opiates for longer than six weeks duration who signed an opioid treatment agreement at least once during Opioid Therapy documented in the medical record	Registry	Patients aged ≥ 1 8 years on date of encounter AND Having one of the encounters (CPT) listed in the next column to the right WITHOUT Telehealth Modifier: GQ, GT, 95, POS 02 AND Patients prescribed opiates for longer than six weeks: G9577	99201-99205, 99212-99215, 99304-99310, 99324-99328, 99334-99337, 99341-99350	Patients who signed an opioid treatment agreement at least once during opioid therapy	G9578 (Performance Met): Documentation of signed opioid treatment agreement at least once during opioid therapy OR G9579 (Performance Not Met): No documentation of signed an opioid treatment agreement at least once during opioid therapy

Measure Number	Measure Description	Reporting Options/M ethods	Denominator Criteria (Eligible Cases)	Possible Denominator Codes Applicable to OMS *	Numerator	Possible Numerator Codes
Measure #414	Evaluation or Interview for Risk of Opioid Misuse — National Quality Strategy Domain: Effective Clinical Care All patients 18 and older prescribed opiates for longer than six weeks duration evaluated for risk of opioid misuse using a brief validated instrument (e.g. Opioid Risk Tool, SOAAP-R) or patient interview documented at least once during Opioid Therapy in the medical record	Registry	Patients aged ≥ 18 years on date of encounter AND Having one of the encounters (CPT) listed in the next column to the right WITHOUT Telehealth Modifier: GQ, GT, 95, POS 02 AND Patients prescribed opiates for longer than six weeks: G9583	99201-99205, 99212-99215, 99304-99310, 99324-99328, 99334-99337, 99341-99350 AND Patients prescribed opiates for longer than 6 weeks	Patients evaluated for risk of misuse of opiates by using a brief validated instrument (e.g., Opioid Risk Tool, SOAAP-R) or patient interview at least once during opioid therapy	G9584 (Performance Met): Patient evaluated for risk of misuse of opiates by using a brief validated instrument (e.g., Opioid Risk Tool, SOAAP-R) or patient interviewed at least once during opioid therapy OR G9585 (Performance Not Met): Patient not evaluated for risk of misuse of opiates by using a brief validated instrument (e.g., Opioid Risk Tool, SOAAP-R) or patient not interviewed at least once during opioid therapy
Measure #415	Emergency Medicine: Emergency Department Utilization of CT for Minor Blunt Head Trauma for Patients Aged 18 Years and Older - National Quality Strategy Domain: Efficiency and Cost Reduction Percentage of emergency department visits for patients aged 18 years and older who	Claims, Registry	Patients aged ≥ 18 years AND having one of the encounters (ICD-10-CM) listed in the next column to the right AND having one of the encounters (CPT codes) listed in	Diagnosis for minor blunt head trauma within the code range- S00-S09 Patient encounter during the performance period: 99281-99285	Emergency department visits for patients who have an indication for a head CT	Patient with Minor Blunt Head Trauma with a Valid Reason for a Head CT for Documented Reasons (Two G-codes [G9531 & G9530] are required on the claim form to submit this numerator option) Denominator Exclusion: G9531: Patient has documentation of ventricular shunt, brain tumor, multisystem trauma, pregnancy, or is currently taking an antiplatelet medication including: ASA/dipyridamole, clopidogrel, prasugrel, ticlopidine, ticagrelor, or cilostazol AND G9530: Patient presented within 24 hours of a minor blunt head trauma with a GCS

Measure Number	Measure Description	Reporting Options/M ethods	Denominator Criteria (Eligible Cases)	Possible Denominator Codes Applicable to OMS *	Numerator	Possible Numerator Codes
	presented within 24 hours of a minor blunt head trauma with a Glasgow Coma Scale (GCS) score of 15 and who had a head CT for trauma ordered by an emergency care provider who have an indication for a head CT		the next column to the right			score of 15 and had a head CT ordered for trauma by an emergency care provider OR If Patient is not Eligible for this Measure because of a Documented Reason as Indicated, Submit: (One G-code [G9532] is required on the claim form to submit this numerator option) Denominator Exclusion: G9532: Patient's head injury occurred greater than 24 hours before presentation to the emergency department, OR has a GCS score less than 15 or does not have a GCS score documented, OR had a head CT for trauma ordered by someone other than an emergency care provider, OR was ordered for a reason other than trauma OR Patient with Minor Blunt Head Trauma had an Appropriate Indication for a Head CT (Two G-codes [G9529 & G9530] are required on the claim form to submit this numerator option) Performance Met: G9529: Patient with minor blunt head trauma had an appropriate indication(s) for a head CT AND G9530: Patient presented within 24 hours of a minor blunt head trauma with a GCS score of 15 and had a head CT ordered for trauma by an emergency care provider OR Patient with Minor Blunt Head Trauma did not have an Appropriate Indication for a Head CT

Measure Number	Measure Description	Reporting Options/M ethods	Denominator Criteria (Eligible Cases)	Possible Denominator Codes Applicable to OMS *	Numerator	Possible Numerator Codes
						(Two G-codes [G9533 & G9530] are required on the claim form to submit this numerator option) Performance Not Met: G9533: Patient with minor blunt head trauma did not have an appropriate indication(s) for a head CT AND G9530: Patient presented within 24 hours of a minor blunt head trauma with a GCS score of 15 and had a head CT ordered for trauma by an emergency care provider
Measure #416	Emergency Medicine: Emergency Department Utilization of CT for Minor Blunt Head Trauma for Patients Aged 2 through 17 Years - National Quality Strategy Domain: Efficiency and Cost Reduction Percentage of emergency department visits for patients aged 2 through 17 years who presented within 24 hours of a minor blunt head trauma with a Glasgow Coma Scale (GCS) score of 15 and who had a head CT for trauma ordered by an emergency care	Claims, Registry	Patients aged 2 through 17 years on date of encounter AND having one of the encounters (ICD- 10-CM) listed in the next column to the right AND having one of the encounters (CPT codes) listed in the next column to the right	Diagnosis for minor blunt head trauma within the code range- S00-S09 Patient encounter during the performance period: 99281-99285	Emergency department visits for patients who are classified as low risk according to the Pediatric Emergency Care Applied Research Network (PECARN) prediction rules for traumatic brain injury	Pediatric Patient with Minor Blunt Head Trauma with a Valid Reason for Head CT for Documented Reasons (Two G-codes [G9595 & G9594] are required on the claim form to submit this numerator option) Denominator Exclusion: G9595: Patient has documentation of ventricular shunt, brain tumor, coagulopathy, including thrombocytopenia AND G9594: Patient presented within 24 hours of a minor blunt head trauma with a GCS score of 15 and had a head CT ordered for trauma by an emergency care provider OR If Patient is Not Eligible for this Measure because of a Documented Reason as Indicated, Submit: (One G-code [G9596] is required on the claim form to submit this numerator option) Denominator Exclusion: G9596: Pediatric patient's head injury occurred greater than 24 hours before presentation to the emergency department, OR has a GCS score less than 15 or does not have a GCS

Measure Number	Measure Description	Reporting Options/M ethods	Denominator Criteria (Eligible Cases)	Possible Denominator Codes Applicable to OMS *	Numerator	Possible Numerator Codes
	provider who are classified as low risk according to the PECARN prediction rules for traumatic brain injury					score documented, OR had a head CT for trauma ordered by someone other than an emergency care provider, OR was ordered for a reason other than trauma OR Pediatric Patient with Minor Blunt Head Trauma Classified as Low Risk According to the PECARN Prediction Rules with Order of Head CT (Two G-codes [G9593 & G9594] are required on the claim form to submit this numerator option) Performance Met: G9593: Pediatric patient with minor blunt head trauma classified as low risk according to the PECARN prediction rules AND G9594: Patient presented within 24 hours of a minor blunt head trauma with a GCS score of 15 and had a head CT ordered for trauma by an emergency care provider OR Pediatric Patient with Minor Blunt Head Trauma Not Classified as Low Risk According to the PECARN Prediction Rules with Order of Head CT (Two G-codes [G9597 & G9594] are required on the claim form to submit this numerator option) Performance Not Met: G9597: Pediatric patient with minor blunt head trauma not classified as low risk according to the PECARN prediction rules AND G9594: Patient presented within 24 hours of a minor blunt head trauma with a GCS score of 15 and had a head CT ordered for trauma by an emergency care provider

Measure Number	Measure Description	Reporting Options/M ethods	Denominator Criteria (Eligible Cases)	Possible Denominator Codes Applicable to OMS *	Numerator	Possible Numerator Codes
Measure #424	Perioperative Temperature Management — National Quality Strategy Domain: Patient Safety Percentage of patients, regardless of age, who undergo surgical or therapeutic procedures under general or neuraxial anesthesia of 60 minutes duration or longer for whom at least one body temperature greater than or equal to 35.5 degrees Celsius (or 95.9 degrees Fahrenheit) was recorded within the 30 minutes immediately before or the 15 minutes immediately after anesthesia end time	Registry	All patients regardless of age AND having one of the encounters (CPT codes) listed in the next column to the right AND Anesthesia of 60 minutes duration or longer: 4255F AND NOT DENOMINATOR EXCLUSIONS: Monitored Anesthesia Care (MAC): G9654 OR Peripheral Nerve Block (PNB): G9770	00100-00210	Patients for whom at least one body temperature greater than or equal to 35.5 degrees Celsius (or 95.9 degrees Fahrenheit) was recorded within the 30 minutes immediately before or the 15 minutes immediately after anesthesia end time	Performance Met: At least 1 body temperature measurement equal to or greater than 35.5 degrees Celsius (or 95.9 degrees Fahrenheit) achieved within the 30 minutes immediately before or the 15 minutes immediately after anesthesia end time (G9771) OR Denominator Exception: Documentation of one of the following medical reason(s) for not achieving at least 1 body temperature measurement equal to or greater than 35.5 degrees Celsius (or 95.9 degrees Fahrenheit) achieved within the 30 minutes immediately before or the 15 minutes immediately after anesthesia end time (e.g., Emergency cases, Intentional hypothermia, etc.) (G9772) OR Performance Not Met: At least 1 body temperature measurement equal to or greater than 35.5 degrees Celsius (or 95.9 degrees Fahrenheit) not achieved within the 30 minutes immediately before or the 15 minutes immediately after anesthesia end time (G9773)
Measure #426	Post-Anesthetic Transfer of Care Measure: Procedure Room to a Post Anesthesia Care Unit (PACU) – National Quality Strategy	Registry	All patients regardless of age AND having one of the encounters (CPT	00100-00210	Patients transferred directly from the procedure room to post- anesthesia care unit (PACU) for	G9655 (Performance Met): A transfer of care protocol or handoff tool/checklist that includes the required key handoff elements is used OR

Measure Number	Measure Description	Reporting Options/M ethods	Denominator Criteria (Eligible Cases)	Possible Denominator Codes Applicable to OMS *	Numerator	Possible Numerator Codes
	Domain: Communication and Care Coordination Percentage of patients, regardless of age, who are under the care of an anesthesia practitioner and are admitted to a PACU in which a post- anesthetic formal transfer of care protocol or checklist which includes the key transfer of care elements is utilized		codes) listed in the next column to the right AND Patient transferred directly from anesthetizing location to PACU: G9656 AND NOT Transfer of care during an anesthetic or to the intensive care unit: G9657		post-procedure care for whom a checklist or protocol which includes the key transfer of care elements is utilized.	G9658 (Performance Not Met): A transfer of care protocol or handoff tool/checklist that includes the required key handoff elements is not used
Measure #427	Post-Anesthetic Transfer of Care: Use of Checklist or Protocol for Direct Transfer of Care from Procedure Room to Intensive Care Unit (ICU) – National Quality Strategy Domain: Communication and Care Coordination Percentage of patients, regardless of age, who undergo a procedure under anesthesia and are admitted to an Intensive Care Unit	Registry	All patients regardless of age AND having one of the encounters (CPT codes) listed in the next column to the right AND Patient transferred directly from anesthetizing location to critical care unit: 0581F	00100-00210	Patients who have a documented use of a checklist or protocol for the transfer of care from the responsible anesthesia practitioner to the responsible ICU team or team member	OR OS83F with 8P: Transfer of care checklist used used

Measure Number	Measure Description	Reporting Options/M ethods	Denominator Criteria (Eligible Cases)	Possible Denominator Codes Applicable to OMS *	Numerator	Possible Numerator Codes
	(ICU) directly from the anesthetizing location, who have a documented use of a checklist or protocol for the transfer of care from the responsible anesthesia practitioner to the responsible ICU team or team member					
Measure #430	Prevention of Post- Operative Nausea and Vomiting (PONV) — Combination Therapy — National Quality Strategy Domain: Patient Safety Percentage of patients, aged 18 years and older, who undergo a procedure under an inhalational general anesthetic, AND who have three or more risk factors for post-operative nausea and vomiting (PONV), who receive combination therapy consisting of at least two prophylactic pharmacologic antiemetic agents of different classes preoperatively	Registry	Patients aged ≥ 18 years on date of encounter AND having one of the encounters (CPT codes) listed in the next column to the right AND Patient received inhalation anesthetic agent: 4554F AND Patient exhibits 3 or more risk factors for post- operative nausea and vomiting: 4556F	00100-00210	Patients who receive combination therapy consisting of at least two prophylactic pharmacologic anti-emetic agents of different classes preoperatively or intraoperatively	Performance Met: Patient received at least 2 prophylactic pharmacologic anti-emetic agents of different classes preoperatively and/or intraoperatively (G9775) OR Denominator Exception: Documentation of medical reason for not receiving at least 2 prophylactic pharmacologic anti-emetic agents of different classes preoperatively and/or intraoperatively (e.g., intolerance or other medical reason) (G9776) OR Performance Not Met: Patient did not receive at least 2 prophylactic pharmacologic anti-emetic agents of different classes preoperatively and/or intraoperatively (G9777)

Measure Number	Measure Description	Reporting Options/M ethods	Denominator Criteria (Eligible Cases)	Possible Denominator Codes Applicable to OMS *	Numerator	Possible Numerator Codes
Measure #431	Preventive Care and Screening: Unhealthy Alcohol Use: Screening & Brief Counseling- National Quality Strategy Domain: Community/ Population Health Percentage of patients aged 18 years and older who were screened at least once within the last 24 months for unhealthy alcohol use using a systematic screening method AND who received brief counseling if identified as an unhealthy alcohol user	Registry	Patients aged ≥ 18 years AND having one of the encounters (CPT or HCPCS codes) listed in the next column to the right WITHOUT Telehealth Modifier: GQ, GT, 95, POS 02 OR At Least One Preventive Visit during the performance period (CPT or HCPC): 96160, 96161, 99385*, 99386*, 99387*, 99395*, 99395*, 99395*, 99397*, 99401*, 99402*, 99403*, 99404*, 99411*, 99412*, 99429*, G0438, G0439 WITHOUT Telehealth Modifier: GQ, GT, 95, POS 02	99201-99205, 99212-99215, G0270, G0271	Patients who were screened at least once within the last 24 months for unhealthy alcohol use using a systematic screening method AND who received brief counseling if identified as an unhealthy alcohol user	identified as an unhealthy alcohol user when screened for unhealthy alcohol use using a systematic screening method and received brief counseling OR Performance Met: Patient identified as an unhealthy alcohol user when screened for unhealthy alcohol user when screened for unhealthy alcohol use using a systematic screening method and received brief counseling (G9621) G9622 (Performance Met): Patient not identified as an unhealthy alcohol user when screened for unhealthy alcohol user when screened for unhealthy alcohol use using a systematic screening method OR G9623 (Medical Performance Exclusion): Documentation of medical reason(s) for not screening for unhealthy alcohol use (e.g., limited life expectancy, other medical reasons) OR G9624 (Performance Not Met): Patient not screened for unhealthy alcohol screening using a systematic screening method OR patient did not receive brief counseling, reason not given
Measure #436	Radiation Consideration for Adult CT: Utilization of Dose Lowering Techniques	Claims, Registry	Patients aged ≥ 18 years AND	70450-70498, 0042T	Final reports with documentation that indicate an individualized dose optimization	G9637 Dose Reduction Techniques (Performance Met): Final reports with documentation of one or more dose reduction techniques (e.g., Automated exposure control, adjustment of the mA

Measure Number	Measure Description	Reporting Options/M ethods	Denominator Criteria (Eligible Cases)	Possible Denominator Codes Applicable to OMS *	Numerator	Possible Numerator Codes
	Percentage of final reports for patients aged 18 years and older undergoing CT with documentation that one or more of the following dose reduction techniques were used: • Automated exposure control • Adjustment of the mA and/or kV according to patient size • Use of iterative reconstruction technique		having one of the encounters (CPT or HCPCS codes) listed in the next column to the right		technique was used for the performed procedure, Dose optimization techniques include the following: • Automated expose control • Adjustment of the mA and/or kV according to patient size • Use of iterative reconstruction techniqe	and/or kV according to patient size, use of iterative reconstruction technique) OR G9638 Dose Reduction Techniques not Performed (Performance Not Met): Final reports without documentation of one or more dose reduction techniques (e.g., Automated exposure control, adjustment of the mA and/or kV according to patient size, use of iterative reconstruction technique)
Measure #463	Prevention of Post- Operative Vomiting (POV) – Combination Therapy (Pediatrics) – National Quality Strategy Domain: Effective Clinical Care Percentage of patients aged 3 through 17 years, who undergo a procedure under	Registry	Patients aged 3 through 17 years on date of encounter AND Patient procedure during the performance period (CPT codes on the next column) Patient received inhalational	00170, 00190	Patients who receive combination therapy consisting of at least two prophylactic pharmacologic anti-emetic agents of different classes preoperatively and/or intraoperatively	Numerator Options: Performance Met: Patient received combination therapy consisting of at least two prophylactic pharmacologic anti-emetic agents of different classes preoperatively and/or intraoperatively (G9956) OR Denominator Exception: Documentation of medical reason for not receiving combination therapy consisting of at least two prophylactic pharmacologic anti-emetic agents of different classes preoperatively and/or intraoperatively (e.g., intolerance or other medical reason) (G9957)

Measure Number	Measure Description	Reporting Options/M ethods	Denominator Criteria (Eligible Cases)	Possible Denominator Codes Applicable to OMS *	Numerator	Possible Numerator Codes
	general anesthesia in which an inhalational anesthetic is used for maintenance AND who have two or more risk factors for post-operative vomiting (POV), who receive combination therapy consisting of at least two prophylactic pharmacologic antiemetic agents of different classes preoperatively and/or intraoperatively		anesthetic agent: 4554F AND Patient exhibits 2 or more risk factors for post- operative vomiting: G9954 AND NOT DENOMINATOR EXCLUSION: Cases in which an inhalational anesthetic is used only for induction: G9955			OR Performance Not Met: Patient did not receive combination therapy consisting of at least two prophylactic pharmacologic anti-emetic agents of different classes preoperatively and/or intraoperatively (G9958)