Centers for Medicare & Medicaid Services

Ambulatory Surgical Center Quality Reporting Program

Quality Measures Specifications Manual

Version 1.0

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# TABLE OF CONTENTS

<table>
<thead>
<tr>
<th>Section</th>
<th>Page</th>
</tr>
</thead>
<tbody>
<tr>
<td>BACKGROUND</td>
<td>1</td>
</tr>
<tr>
<td>THE SPECIFICATIONS MANUAL</td>
<td>3</td>
</tr>
<tr>
<td>Ambulatory Surgical Center (ASC) Quality Reporting Measures</td>
<td>5</td>
</tr>
<tr>
<td>Patient Burn</td>
<td>5</td>
</tr>
<tr>
<td>Patient Fall</td>
<td>8</td>
</tr>
<tr>
<td>Wrong Site, Wrong Side, Wrong Patient, Wrong Procedure, Wrong Implant</td>
<td>10</td>
</tr>
<tr>
<td>Hospital Transfer/Admission</td>
<td>12</td>
</tr>
<tr>
<td>Prophylactic Intravenous (IV) Antibiotic Timing</td>
<td>14</td>
</tr>
<tr>
<td>ASC Facility Volume Data on Selected ASC Surgical Procedures</td>
<td>17</td>
</tr>
<tr>
<td>Safe Surgery Checklist Use</td>
<td>19</td>
</tr>
<tr>
<td>APPENDIX A: DATA DEFINITIONS</td>
<td>20</td>
</tr>
</tbody>
</table>
BACKGROUND

Quality Reporting for Ambulatory Surgical Centers

Welcome to quality reporting for Ambulatory Surgical Centers (ASCs)! This manual provides specifications for quality measures finalized for reporting to meet requirements for this recently finalized program.

A quality reporting program for ASCs was finalized by the Centers for Medicare and Medicaid Services (CMS) in the Calendar Year (CY) 2012 OPPS/ASC Final Rule with Comment Period (CMS-1525-FC). Five claims-based measures (four outcome measures and one process of care measure) were adopted for the CY 2014 payment determination. For the CY 2015 payment determination, the same claims-based measures and two structural measures (surgical procedure volume and safe surgery checklist use) were adopted for a total of seven quality measures. For the CY 2016 payment determination, the same claims-based and structural measures as adopted for the CY 2015 payment determination and one process of care measure were adopted.

ASCs that do not meet program requirements for ASC Quality Reporting may receive a 2 percent reduction in their ASC annual payment update. Thus, only separately identifiable entities certified as ASC by Medicare are affected by program requirements and possible payment penalty under the ASC Quality Reporting Program. The definition of an ASC can be found in the Claims Processing Manual, Chapter 14, Section 10.1 located on the CMS website (www.cms.hhs.gov).

The below table summarizes the quality measures, reporting periods, and payment years affected.

Table 1: ASC Quality Measures, Reporting Periods, and Initial Payment Year Affected

<table>
<thead>
<tr>
<th>Measure</th>
<th>Reporting Period</th>
<th>Payments Affected</th>
</tr>
</thead>
</table>

The establishment of quality measure reporting procedures for ambulatory surgical centers was authorized under the Medicare Improvements and Extension Act of 2006 under Title I of the Tax Relief and Health Care Act of 2006 (Pub. L. 109-432).
Data Collection and Submission

Data for claims-based measures included in this specifications manual are to be reported for all Medicare fee-for-service (FFS) patients admitted to the ASC during required reporting periods (see Table 1).

For claims-based measures, the reporting period refers to dates of service, not to the claim submission date. For example, if a service was provided on September 30, 2012 with claim submission on October 1, 2012, this claim would not be included because the service date was prior to the reporting period.

Data for structural measures relates to all ASC patients.

Note that specifications for the Influenza Vaccination Coverage for Health Care Workers process of care measure are not included in this manual.

Claims-based Measures

ASCs are to submit information on the five claims-based measures using Quality Data Codes (QDCs) entered on their claims submitted using the CMS-1500 or associated electronic dataset. QDCs are specified CPT Category II codes or Level II G-codes that describe the clinical action required by a measure’s numerator. Clinical actions can apply to more than one condition and therefore, can also apply to more than one measure. Some measures require more than one clinical action and, therefore, have more than one associated QDC. Facilities should review numerator reporting instructions carefully.

The selected QDC(s) are to be reported in addition to any codes that would be standard for billing purposes (e.g., the ICD-9-CM diagnosis and Current Procedural Terminology (CPT) codes, Healthcare Common Procedure Coding System (HCPCS) Level II and CPT Category III codes for the services performed) on the ASC claim for the encounter.

Data completeness for the reporting of these measures has been proposed to be calculated by comparing the number of claims meeting measure specifications with the appropriate QDCs to the number of claims that would meet measure specifications without the appropriate QDCs on the submitted claim. Requirements for reporting completeness will be finalized prior to data collection beginning in October 1, 2012.

Structural Measures

Data for structural measures are to be submitted using a web-based tool that will be located on the QualityNet website located at www.QualityNet.org. Data collection for structural measures is required in 2013 and the tool will be available at this time for data entry.

Public Reporting

The Secretary of Health and Human Services must establish procedures to make data collected under the Quality Reporting Programs. Under the ASC Quality Reporting Program, facilities will be provided the opportunity to review their data prior to publication. Details on the publication of data, the ability to withdraw and not have data publicly reports, and reconsideration processes have been proposed and will be finalized prior to data collection beginning October 1, 2012.
THE SPECIFICATIONS MANUAL

This Specifications Manual provides measure specifications, associated QDCs with descriptions, and references for required ASC Quality Reporting Program quality measures.

The claims-based ASC quality measures adopted by CMS for the ASC Quality Reporting Program were developed by the ASC Quality Collaboration. These measures are the intellectual property of the ASC Quality Collaboration. Additional information about the ASC quality measures endorsed by the National Quality Forum (NQF) is available in the ASC Quality Collaboration Implementation Guide (www.ascquality.org).

Information for each of the ASC Quality Collaboration measures is displayed in the following format:

**Title of Measure** - Provides the title of the measure

**Quality Reporting Option** - States whether the measure is an outcome, structural, or a process of care measure.

**Description** - A brief description of what is being measured.

**Numerator** - The patient population experiencing the outcome or process of care being measured.

**Denominator** - The patient population evaluated.

**Numerator Inclusions** - Patients to be included in the patient population experiencing the outcome or process of care being measured.

**Numerator Exclusions** - Patients to be excluded from the patient population experiencing the outcome or process of care being measured.

**Denominator Inclusions** - Patients included in the population to be evaluated.

**Denominator Exclusions** - Patients to be excluded from the population to be evaluated.

**Coding options** - A list and description of the G-code(s) used to report the measure

**Data Sources** - The documents that typically contain the information needed to determine the numerator and denominator.

**Definitions** - Specific definitions for the terms included in the numerator and denominator statements.
IMPORTANT

A QDC has been established to report that the patient did **not** experience the events for four of the five claims-based outcome measures. If this code is used, none of the other QDCs should be used for these four measures.

**G8907:** Patient documented **not** to have experienced any of the following events: a burn prior to discharge; a fall within the facility; wrong site, wrong side, wrong patient, wrong procedure or wrong implant event; or a hospital transfer or hospital admission upon discharge from the facility.

**Note:** For surgical patients with an order for prophylactic antibiotics, information on the fifth measure, Prophylactic IV Antibiotic Timing, will be reported separately. If the patient received the prophylactic antibiotic on time and did not experience any of the events (a burn prior to discharge; a fall within the facility; wrong site, wrong side, wrong patient, wrong procedure or wrong implant event; or a hospital transfer or hospital admission upon discharge from the facility), the code listed above (G8907) would be used **in addition to** G8916. See each measure for the list of available codes.
Measure Title: Patient Burn

MEASURE ID #: ASC-1

REPORTING MECHANISM:
Medicare Fee-for-Service Claims

DESCRIPTION:
The number of admissions (patients) who experience a burn prior to discharge.

DENOMINATOR:
All ASC admissions
  Inclusions: All ASC admissions.
  Exclusions: None

NUMERATOR:
ASC admissions experiencing a burn prior to discharge.
  Inclusions: ASC admissions experiencing a burn prior to discharge.
  Exclusions: None

Numerator Quality-Data Coding Options for Reporting:
G8908: Patient documented to have received a burn prior to discharge.
G8909: Patient documented not to have received a burn prior to discharge.
G8907: Patient documented not to have experienced any of the following events: a burn prior to discharge; a fall within the facility; wrong site, wrong side, wrong patient, wrong procedure or wrong implant event; or a hospital transfer or hospital admission upon discharge from the facility.

Note: If using code G8908 or G8909, do not use code G8907.

DEFINITIONS:
Admission - completion of registration upon entry into the facility.
Burn - Unintended tissue injury caused by any of the six recognized mechanisms: scalds, contact, fire, chemical, electrical or radiation (e.g. warming devices, prep solutions, electrosurgical unit or laser).
Discharge - occurs when the patient leaves the confines of the ASC.

SELECTION BASIS:
There are numerous case reports in the literature regarding patient burns in the surgical and procedural setting. The diversity of the causative agents underscores the multitude of potential risks that must be properly mitigated to avoid patient burns.

The literature on burns suggests that electrosurgical burns are most common. A recent publication from the ECRI Institute (www.ecri.org) highlights the increased risk of burns with newer surgical devices that apply higher currents at longer activation times. Although electrical burns are most prevalent, other mechanisms of burn injury are frequently reported in case studies and case series. These include chemical and thermal burns.
Surgical fires are rare; however, their consequences can be grave, killing or seriously injuring patients and surgical staff. The risk of surgical fire is present whenever and wherever surgery is performed, whether in an operating room (OR), a physician’s office, or an outpatient clinic. Recognition of the diverse mechanisms by which a patient could sustain an unintentional burn in the ASC setting, scaling, contact, fire, chemical, electrical, or radiation, this will allow stakeholders to develop a better understanding of the incidence of these events and further refine preventive processes.

**CLINICAL RECOMMENDATION STATEMENTS:**
The risk of burns related to laser use can be reduced by adherence to the guidelines published by the American National Standards Institute (ANSI) for safe use of these devices in the health care setting. Similarly, the risk of burns related to the use of electrosurgical devices can be reduced by following the electrosurgery checklist published by ECRI Institute.

The risk of surgical fires can be reduced by minimizing ignition, oxidizer, and fuel risks (the “classic triangle”). The American Society of Anesthesiologist’s Practice Advisory for the Prevention and Management of Operating Room Fires seeks to prevent the occurrence of OR fires, reduce adverse outcomes associated with OR fires, and identify the elements of a fire response protocol.

These guidelines are available at: [http://www.asahq.org/For-Members/Practice-Management/Practice-Parameters.aspx](http://www.asahq.org/For-Members/Practice-Management/Practice-Parameters.aspx).

Guidance for the prevention of surgical fire has also been published by the Association of Perioperative Registered Nurses (AORN).

**REFERENCES**
Measure Title: Patient Fall

MEASURE ID #: ASC-2

REPORTING MECHANISMS:
Medicare Fee-for-Service Claims

DESCRIPTION:
The number of admissions (patients) who experience a fall within the ASC.

DENOMINATOR:
All ASC admissions
  Inclusions: All ASC admissions.
  Exclusions: None

NUMERATOR:
ASC admissions experiencing a fall within the confines of the ASC.
  Inclusions: ASC admissions experiencing a fall within the confines of the ASC.
  Exclusions: ASC admissions experiencing a fall outside the ASC.

Numerator Quality-Data Coding Options for Reporting:
G8910: Patient documented to have experienced a fall within the ASC.
G8911: Patient documented not to have experienced a fall within the ASC.
G8907: Patient documented not to have experienced any of the following events: a burn prior to discharge; a fall within the facility; wrong site, wrong side, wrong patient, wrong procedure or wrong implant event; or a hospital transfer or hospital admission upon discharge from the facility.

Note: If using code G8910 or G8911, do not use code G8907.

DEFINITIONS:
Admission - completion of registration upon entry into the facility.
Fall - a sudden, uncontrolled, unintentional, downward displacement of the body to the ground or other object, excluding falls resulting from violent blows or other purposeful actions (source: National Center for Patient Safety).

SELECTION BASIS:
“Falls per 100,000 patient days” has been endorsed as a serious reportable event by the NQF. While ASCs have a relatively low incidence of adverse events in general; information regarding the incidence of patient falls is not currently available. Stakeholders have expressed an interest in the public reporting of such adverse events. Due to the use of anxiolytics, sedatives, and anesthetic agents as adjuncts to procedures, patients undergoing outpatient surgery are at increased risk for falls.

CLINICAL RECOMMENDATION STATEMENTS:
The Agency for Healthcare Research and Quality’s (AHRQ) Prevention of Falls in Acute Care guidelines state that patient falls can be reduced by following a four-step approach: 1) evaluating and identifying risk factors for falls in the older patient; 2) developing an appropriate plan of care...
for prevention; 3) performing a comprehensive evaluation of falls that occur; and 4) performing a post-fall revision of plan of care as appropriate.

REFERENCES

- ECRI Institute. Falls Prevention Resources: https://www.ecri.org/Products/Pages/Fall_Prevention_Resources.aspx.
- American Medical Directors Association (AMDA). Falls and fall risk. Columbia, MD: American Medical Directors Association.
Measure Title: Wrong Site, Wrong Side, Wrong Patient, Wrong Procedure, Wrong Implant

MEASURE ID #: ASC-3

REPORTING MECHANISM:
Medicare Fee-for-Service Claims

DESCRIPTION:
Any ASC admissions (patients) who experience a wrong site, side, patient, procedure or implant.

DENOMINATOR:
All ASC admissions
  Inclusions: All ASC admissions.
  Exclusions: None

NUMERATOR:
All ASC admissions experiencing a wrong site, wrong side, wrong patient, wrong procedure or wrong implant.
  Inclusions: All ASC admissions experiencing a wrong site, wrong side, wrong patient, wrong procedure or wrong implant.
  Exclusions: None

Numerator Quality-Data Coding Options for Reporting:
G8912: Patient documented to have experienced a wrong site, wrong side, wrong patient, wrong procedure or wrong implant event.
G8913: Patient documented not to have experienced a wrong site, wrong side, wrong patient, wrong procedure or wrong implant event.
G8907: Patient documented not to have experienced any of the following events: a burn prior to discharge; a fall within the facility; wrong site, wrong side, wrong patient, wrong procedure or wrong implant event; or a hospital transfer or hospital admission upon discharge from the facility.

Note: If using code G8912 or G8913, do not use code G8907.

DEFINITIONS:
Admission - completion of registration upon entry into the facility.
Wrong - not in accordance with intended site, side, patient, procedure or implant.

SELECTION BASIS:
“Surgery performed on the wrong body part,” “surgery performed on the wrong patient,” and “wrong surgical procedure performed on a patient” have all been endorsed as serious reportable surgical events by NQF. This outcome measure serves as an indirect measure of providers’ adherence to The Joint Commission, an accreditation body, has developed a “Universal Protocol” guideline for eliminating wrong site, wrong procedure, wrong person surgery. The Universal Protocol is based on the consensus of experts and is endorsed by more than forty professional medical associations and organizations. To encompass the outcomes of all key identification verifications, the ASC
Quality Collaboration’s measure incorporates not only wrong site, wrong side, wrong patient and wrong procedure, but also wrong implant in its specifications.

**CLINICAL RECOMMENDATION STATEMENTS:**
The Joint Commission’s “Universal Protocol” is based on the consensus of experts from the relevant clinical specialties and professional disciplines and is endorsed by more than 40 professional medical associations and organizations.

**REFERENCES**
Measure Title: Hospital Transfer/Admission

MEASURE ID #: ASC-4

REPORTING MECHANISM:
Medicare-Fee-for-Service Claims

DESCRIPTION:
ASC admissions (patients) who are transferred or admitted to a hospital upon discharge from the ASC.

DENOMINATOR:
All ASC admissions
- Inclusions: All ASC admissions.
- Exclusions: None

NUMERATOR:
ASC admissions requiring a hospital transfer or hospital admission upon discharge from the ASC.
- Inclusions: ASC admissions requiring a hospital transfer or hospital admission upon discharge from the ASC.
- Exclusions: None

Numerator Quality-Data Coding Options for Reporting:
G8914: Patient documented to have experienced a hospital transfer or hospital admission upon discharge from ASC.

G8915: Patient documented not to have experienced a hospital transfer or hospital admission upon discharge from ASC.

G8907: Patient documented not to have experienced any of the following events: a burn prior to discharge; a fall within the facility; wrong site, wrong side, wrong patient, wrong procedure or wrong implant event; or a hospital transfer or hospital admission upon discharge from the facility.

Note: If using code G8914 or G8915, do not use code G8907.

DEFINITIONS:
- Admission - completion of registration upon entry into the facility.
- Hospital Transfer/Admission - any transfer/admission from an ASC directly to an acute care hospital including hospital emergency room.
- Discharge - occurs when the patient leaves the confines of the ASC.

SELECTION BASIS:
The need for transfer/admission is an unanticipated outcome and could be the result of insufficient rigor in patient or procedure selection. Hospital transfers/admissions can result in unplanned cost and time burdens that must be borne by patients and payers.

Selected states have expressed an interest in the public reporting of such events. While hospital transfers and admissions undoubtedly represent good patient care when necessary, high rates may be an indicator that practice patterns or patient selection guidelines are in need of review.
CLINICAL RECOMMENDATION STATEMENTS:
No clinical practice guidelines specifically addressing transfers or admissions from ASCs to acute care hospitals are available at this time.

REFERENCES

Measure Title: Prophylactic Intravenous (IV) Antibiotic Timing

MEASURE ID #: ASC-5

REPORTING MECHANISM: Medicare-Fee-for-Service Claims

DESCRIPTION: Intravenous (IV) antibiotics given for prevention of surgical site infection were administered on time.

DENOMINATOR: All ASC admissions with a preoperative order for a prophylactic IV antibiotic for prevention of surgical site infection.

Inclusions: All ASC admissions with a preoperative order for a prophylactic IV antibiotic for prevention of surgical site infection.

Exclusions: ASC admissions with a preoperative order for a prophylactic IV antibiotic for prevention of infections other than surgical site infections (e.g. bacterial endocarditis); ASC admissions with a preoperative order for a prophylactic antibiotic not administered by the intravenous route.

NUMERATOR: Number of ASC admissions with an order for a prophylactic IV antibiotic for prevention of surgical site infection who received the prophylactic antibiotic on time.

Inclusions: All ASC admissions with a preoperative order for a prophylactic IV antibiotic for prevention of surgical site infection.

Exclusions: None

Numerator Quality-Data Coding Options for Reporting:

G8916: Patient with preoperative order for IV antibiotic surgical site infection (SSI) prophylaxis, antibiotic initiated on time.

G8917: Patient with preoperative order for IV antibiotic surgical site infection (SSI) prophylaxis, antibiotic not initiated on time.

G8918: Patient without preoperative order for IV antibiotic surgical site infection (SSI) prophylaxis.

Note: The QDC of G8907 can be used if the patient did not experience any of the events for the four outcome measures (a burn prior to discharge; a fall within the facility; wrong site, wrong side, wrong patient, wrong procedure or wrong implant event; or a hospital transfer or hospital admission upon discharge from the facility); this code would be used plus one of the codes above for the prophylactic antibiotic timing measure for complete reporting of the 5 claims-based measures.

DEFINITIONS:

Admission - completion of registration upon entry into the facility.

Antibiotic administered on time - Antibiotic infusion is initiated within one hour prior to the time of the initial surgical incision or the beginning of the procedure (e.g., introduction of endoscope, insertion of needle, inflation of tourniquet) or two hours prior if vancomycin or fluoroquinolones are administered.
Intravenous - Administration of a drug within a vein, including bolus, infusion or IV piggyback.

Order - a written order, verbal order, standing order or standing protocol.

Prophylactic antibiotic - an antibiotic prescribed with the intent of reducing the probability of an infection related to an invasive procedure. For purposes of this measure, the following antibiotics are considered prophylaxis for surgical site infections: Ampicillin/sulbactam, Aztreonam, Cefazolin, Cefmetazole, Cefotetan, Cefoxitin, Cefuroxime, Ciprofloxacin, Clindamycin, Ertapenem, Erythromycin, Gatifloxacin, Gentamicin, Levofloxacin, Metronidazole, Moxifloxacin, Neomycin and Vancomycin.

SELECTION BASIS:
The CMS Surgical Infection Prevention performance measure states, “Surgical site infections occur in 2-5 percent of clean extra-abdominal surgeries and up to 20 percent of intra-abdominal surgeries. Each infection is estimated to increase a hospital stay by an average of 7 days and add over $3,000 in charges (1992 data). Patients who develop surgical site infections are 60 percent more likely to spend time in an ICU (intensive care unit), five times more likely to be readmitted to the hospital, and have twice the incidence of mortality. Despite advances in infection control practices, surgical site infections remain a substantial cause of morbidity and mortality among hospitalized patients. Studies indicate that appropriate preoperative administration of antibiotics is effective in preventing infection. Systemic and process changes that promote compliance with established guidelines and standards can decrease infectious morbidity.”

There is no literature available on variation in adherence to recommended prophylactic IV antibiotic timing among ASC providers. However, variability in the accuracy of timing of administration has been demonstrated in other clinical settings.

CLINICAL RECOMMENDATION STATEMENTS:
This performance measure is aligned with current surgical infection prevention guidelines recommending that prophylactic antibiotics be administered within one hour prior to surgical incision, or within two hours prior to incision when vancomycin or fluoroquinolones are used.

REFERENCES


Measure Title: ASC Facility Volume Data on Selected ASC Surgical Procedures

MEASURE ID#: ASC-6

REPORTING MECHANISM:
Web-based tool on QualityNet

DESCRIPTION: The aggregate count of selected surgical procedures. Most ASC procedures fall into one of eight categories: Cardiovascular, Eye, Gastrointestinal, Genitourinary, Musculoskeletal, Nervous System, Respiratory, and Skin. The eight categories and corresponding HCPCS are listed in the table below. The procedures and codes in Table 2 were selected based on recent ASC data and update the procedure codes listed in the Calendar Year (CY) 2012 OPPS/ASC Final Rule with Comment Period (CMS-1525-FC).

Measure ascertains response to the following question(s):
- What was the aggregate count of selected surgical procedures per category?


Table 2: Categories and HCPCS for ASC-6

<table>
<thead>
<tr>
<th>Organ System</th>
<th>CMS Procedure Category</th>
<th>Surgical Procedure Codes</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cardiovascular</td>
<td>Placement of long-term vascular access catheter</td>
<td>36561</td>
</tr>
<tr>
<td></td>
<td>Vascular procedures to improve blood flow to coronary (heart) vessels</td>
<td>92980, 92981, 92982, 92984</td>
</tr>
<tr>
<td>Eye</td>
<td>Organ transplant (eye)</td>
<td>65756, V2785</td>
</tr>
<tr>
<td></td>
<td>Laser procedure of eye</td>
<td>65855, 66761, 66821</td>
</tr>
<tr>
<td></td>
<td>Glaucoma procedures</td>
<td>66170, 66180</td>
</tr>
<tr>
<td></td>
<td>Cataract procedures</td>
<td>66982, 66984</td>
</tr>
<tr>
<td></td>
<td>Injection of eye</td>
<td>67028, J2778, J3300, J3396</td>
</tr>
<tr>
<td></td>
<td>Retina, macular and posterior segment procedures</td>
<td>67041, 67042, 67210, 67228</td>
</tr>
<tr>
<td></td>
<td>Repair of surrounding eye structures</td>
<td>67900, 67904, 67917, 67924</td>
</tr>
<tr>
<td>Gastrointestinal</td>
<td>GI endoscopy procedures</td>
<td>43239, 43235, 43248, 43249, 43251, 44361, 45330, 45331, 45378, 45380, 45381, 45383, 45384, 45385</td>
</tr>
<tr>
<td></td>
<td>Swallowing tube (esophagus)</td>
<td>43450</td>
</tr>
<tr>
<td></td>
<td>Hernia repair</td>
<td>49505</td>
</tr>
<tr>
<td></td>
<td>GI screening procedures</td>
<td>G0105, G0121</td>
</tr>
<tr>
<td>Genitourinary</td>
<td>Kidney stone fragmentation</td>
<td>50590</td>
</tr>
<tr>
<td></td>
<td>Bladder related procedures</td>
<td>52000, 52005, 52204, 52281, 52310, 52332,</td>
</tr>
<tr>
<td></td>
<td>Prostate biopsy</td>
<td>55700</td>
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<tr>
<td></td>
<td>Radiologic procedures (GU)</td>
<td>74420</td>
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<tr>
<td></td>
<td>Ultrasound procedures (GU)</td>
<td>76872</td>
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<tr>
<td>Category</td>
<td>Procedures</td>
<td>Codes</td>
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<tr>
<td>---------------</td>
<td>-----------------------------------------------------------------</td>
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</tr>
<tr>
<td>Musculoskeletal</td>
<td>Joint or muscle aspiration or injection</td>
<td>20610</td>
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<tr>
<td></td>
<td>Removal of musculoskeletal implants</td>
<td>20680</td>
</tr>
<tr>
<td></td>
<td>Repair of tendons and ligaments</td>
<td>23412</td>
</tr>
<tr>
<td></td>
<td>Repair of foot, toes, fingers, and wrist</td>
<td>26055, 28270, 28285, 28296, 29848</td>
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<tr>
<td></td>
<td>Removal of musculoskeletal lesion</td>
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<td>Joint arthroscopy</td>
<td>29824, 29826, 29827, 29880, 29881</td>
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<tr>
<td></td>
<td>Musculoskeletal drug injection</td>
<td>J0585</td>
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<tr>
<td>Nervous</td>
<td>Injection procedures in or around the spine</td>
<td>62310, 62311, 64479, 64483, 64484, 64490, 64491, 64492, 64493, 64494, 64495, 64622, 64623, 64626, 64627, G0260</td>
</tr>
<tr>
<td></td>
<td>Device implant</td>
<td>63650</td>
</tr>
<tr>
<td></td>
<td>Nerve decompression</td>
<td>64718</td>
</tr>
<tr>
<td></td>
<td>Repair of foot, toes, fingers, and wrist</td>
<td>64721</td>
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<tr>
<td>Respiratory</td>
<td>Sinus procedure</td>
<td>31255</td>
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<td>Skin</td>
<td>Skin procedures</td>
<td>11042, 13132, 14040, 14060, 15260, Q4101, Q4102, Q4106</td>
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<td>Repair of surrounding eye structures</td>
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<tr>
<td>Multi-system*</td>
<td>Brachytherapy</td>
<td>C2638, C2639, C2640, C2641</td>
</tr>
<tr>
<td></td>
<td>Cancer treatment</td>
<td>C9257</td>
</tr>
</tbody>
</table>

*Multi-System: procedures that can be performed in more than one organ system.
Measure Title: Safe Surgery Checklist Use

MEASURE ID #: ASC-7

REPORTING MECHANISM:
Web-based tool on QualityNet

Description: The use of a Safe Surgery Checklist for surgical procedures that includes safe surgery practices during each of the three critical perioperative periods: the period prior to the administration of anesthesia, the period prior to skin incision, and the period of closure of incision and prior to the patient leaving the operating room.

Measure ascertains response to the following question(s):
- Does/did your facility use a safe surgery checklist based on accepted standards of practice at any time during the designated period? Yes/No


Examples for Safe Surgery Practices*

<table>
<thead>
<tr>
<th>First critical point (period prior to administering anesthesia)</th>
<th>Second critical point (period prior to skin incision)</th>
<th>Third critical point (period of closure of incision and prior to patient leaving the operating room)</th>
</tr>
</thead>
<tbody>
<tr>
<td>- Verbal confirmation of patient identity</td>
<td>- Confirm surgical team members and roles</td>
<td>- Confirm the procedure</td>
</tr>
<tr>
<td>- Mark surgical site</td>
<td>- Confirm patient identity, procedure and surgical incision site</td>
<td>- Complete count of surgical instruments and accessories</td>
</tr>
<tr>
<td>- Check anesthesia machine/medication</td>
<td>- Administration of antibiotic prophylaxis within 60 minutes before incision</td>
<td>- Identify key patient concerns for recovery and management of the patient</td>
</tr>
<tr>
<td>- Assessment of allergies, airway and aspiration risk</td>
<td>- Communication among surgical team members of anticipated critical events</td>
<td></td>
</tr>
<tr>
<td></td>
<td>- Display of essential imaging as appropriate</td>
<td></td>
</tr>
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*Hospital safe surgery checklist items are not limited to the examples listed in this table.
APPENDIX A: DATA DEFINITIONS

**Admission:** Completion of registration upon entry into the facility.

**Antibiotic administered on time:** Antibiotic infusion is *initiated* within one hour prior to the time of the initial surgical incision or the beginning of the procedure (e.g., introduction of endoscope, insertion of needle, inflation of tourniquet) or two hours prior if vancomycin or fluoroquinolones are administered.

**Burn:** Unintended tissue injury caused by any of the six recognized mechanisms: scalds, contact, fire, chemical, electrical or radiation, (e.g. warming devices, prep solutions, electrosurgical unit or laser).

**Discharge:** Occurs when the patient leaves the confines of the ASC.

**Fall:** A sudden, uncontrolled, unintentional, downward displacement of the body to the ground or other object, excluding falls resulting from violent blows or other purposeful actions. (National Center for Patient Safety)

**Hospital transfer/admission:** Any transfer/admission from an ASC directly to an acute care hospital including hospital emergency room or emergency department.

**Intravenous:** Administration of a drug within a vein, including bolus, infusion or IV piggyback.

**Order:** A written order, verbal order, standing order or standing protocol.

**Prophylactic antibiotic:** An antibiotic prescribed with the intent of reducing the probability of an infection related to an invasive procedure. For purposes of the Prophylactic IV Antibiotic Timing measure, the following antibiotics are considered prophylaxis for surgical site infections: Ampicillin/sulbactam, Aztreonam, Cefazolin, Cefmetazole, Cefotetan, Cefoxitin, Cefuroxime, Ciprofloxacin, Clindamycin, Ertapenem, Erythromycin, Gatifloxacin, Gentamicin, Levofloxacin, Metronidazole, Moxifloxacin, Neomycin and Vancomycin.

**Quality Data Code (QDC):** Non-payable Healthcare Common Procedure Coding System (HCPCS) codes comprised of specified CPT Category II codes and/or G-codes that describe the clinical action required by a measure’s numerator.

**Wrong:** Not in accordance with intended site, side, patient, procedure or implant.

Additional information and resources, such as sample data collection sheets or logs and frequently asked questions (FAQs) about the measures, can be found on the ASC Quality Collaboration website at [www.ascquality.org](http://www.ascquality.org).