News Flash – The Centers for Medicare & Medicaid Services (CMS) recently issued a final rule that will change how Medicare pays for dialysis services for Medicare beneficiaries who have End-Stage Renal Disease (ESRD). CMS also issued a proposed rule that would establish a new Quality Incentive Program (QIP) to promote high quality services in dialysis facilities by linking a facility’s payments to performance standards. The QIP is the first pay-for-performance program in a Medicare Fee-For-Service (FFS) payment system. For additional information please see the CMS Fact sheet (7/26) at http://www.cms.gov/apps/media/fact_sheets.asp on the CMS website.

MLN Matters® Number: SE1028 Revised Related Change Request (CR) #: N/A
Related CR Release Date: N/A Effective Date: N/A
Related CR Transmittal #: N/A Implementation Date: N/A

Recovery Audit Contractor (RAC) Demonstration High-Risk Diagnosis Related Group (DRG) Coding Vulnerabilities for Inpatient Hospitals

This is the third in a series of articles that will disseminate information on RAC Demonstration high dollar improper payment vulnerabilities. The purpose of this article is to provide inpatient hospital education regarding four RAC demonstration-identified inpatient hospital coding vulnerabilities, in an effort to prevent these same problems from occurring in the future. With the expansion of the RAC Program and the initiation of complex coding review in all four RAC regions, it is essential that providers understand the lessons learned from the demonstration and implement appropriate corrective actions.

Note: This article was updated on August 21, 2012, to reflect current Web addresses. Previously, this article was revised on October 29, 2010, to clarify requirements for coding diagnosis codes by attending physicians.

Provider Types Affected

This article is for all Inpatient Hospital providers that submit Fee-For-Service claims to Medicare Fiscal Intermediaries (FIs) or Part A/B Medicare Administrative Contractors (MACs).

Disclaimer

This article was prepared as a service to the public and is not intended to grant rights or impose obligations. This article may contain references or links to statutes, regulations, or other policy materials. The information provided is only intended to be a general summary. It is not intended to take the place of either the written law or regulations. We encourage readers to review the specific statutes, regulations and other interpretive materials for a full and accurate statement of their contents.
Provider Action Needed

Review the article and take steps, if necessary, to meet Medicare’s documentation requirements to avoid unnecessary denial of your claims.

Background

Effective March 2005, the RAC Demonstration began in California, Florida, and New York. In 2007, the program expanded to include Massachusetts, Arizona, and South Carolina before ending on March 27, 2008. The primary goal of the RAC demonstration was to determine if recovery auditing could be effective in Medicare. The Centers for Medicare & Medicaid Services (CMS) directed the RAC staff to organize their efforts primarily to attain that goal. Supplemental goals, such as correcting identified vulnerabilities, were identified after the fact and were not required tasks. CMS did collect improper hospital payment information from the RACs. However, it was on a voluntary basis, at the claim level and focused on the collection and not the principal and secondary diagnoses on a claim. Four of the high risk inpatient hospital coding vulnerabilities identified are listed in Table 1 below. These claims were denied because the demonstration RACs determined that the medical record documentation submitted did not support the codes billed.

Table 1

<table>
<thead>
<tr>
<th>Provider Type</th>
<th>Improper Payment Amount (pre-appeal)</th>
<th>RAC Demonstration Findings</th>
</tr>
</thead>
<tbody>
<tr>
<td>1 Inpatient Hospital</td>
<td>$15,999,757</td>
<td>Respiratory System Diagnosis with Vent support – (CMS DRG 475) – Principal diagnosis on the claim did not match the principal diagnosis in the medical record.</td>
</tr>
<tr>
<td>2 Inpatient Hospital</td>
<td>$11,769,645</td>
<td>Closed Biopsy of Lung (CMS DRG 076, 077,120) - A transbronchial lung biopsy was billed but the medical record showed a transbronchial biopsy was performed.</td>
</tr>
<tr>
<td>3 Inpatient Hospital</td>
<td>$10,014,530</td>
<td>OR Procedure for Infections, Parasitic Diseases (CMS DRG 415) – The codes on the claim did not match information in the medical record.</td>
</tr>
</tbody>
</table>
For example, one of the coding vulnerabilities the RACs identified was that hospitals were inappropriately reporting a surgical code 33.27, closed endoscopic biopsy of the lung. The medical record documentation indicated that the site of the biopsy was the bronchus, not the lung, and therefore the correct code to bill is the non-surgical code 33.24, closed endoscopic biopsy of the bronchus.

**Inpatient Hospital Medical Documentation Reminders**


The general rules for reporting secondary diagnoses (Section III, “Reporting Additional Diagnoses” of the Official Coding Guidelines) are that they affect patient care in terms of requiring:

1. Clinical evaluation or
2. Diagnostic treatment or
3. Therapeutic treatment or
4. Causes an increase in the Length of Stay (LOS) or
5. Increased nursing care and/or monitoring.

CMS encourages providers to ensure that all fields on documentation tools (such as assessments, flow sheets, checklists, etc.) are completed. If a field is not applicable, CMS recommends that providers use an entry such as “N/A” to show that the questions were reviewed and answered. Fields that are left blank often lead the reviewer to make an inaccurate determination.

Documentation that is not legible has a direct effect on the RAC reviewer’s ability to support that the services billed were coded correctly, medically necessary and were provided in an appropriate setting.

During the RAC demonstration reviewers noted that entries in the medical records were not consistent. CMS encourages providers to ensure all entries are consistent with other parts of the medical record (assessments, treatment plans, and physician orders, nursing notes, medication and treatment records, etc. and other facility documents such as admission and discharge data, pharmacy records, etc.). If an entry is made that contradicts documentation found elsewhere in the record, clarification should be obtained and documented by the attending physician.

Lastly, CMS reminds providers to ensure that any information that affects the billed services and is acquired after physician documentation is complete, must be added to the existing documentation in accordance with accepted standards for amending medical record documentation.

**Additional Information**

Providers are also encouraged to visit the CMS RAC website at [http://www.cms.gov/Research-Statistics-Data-and-Systems/Monitoring-Programs/recovery-audit-program/index.htm](http://www.cms.gov/Research-Statistics-Data-and-Systems/Monitoring-Programs/recovery-audit-program/index.htm) for updates on the National RAC Program. On that website, you can register to receive email updates and view current RAC activities nationwide.

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