

3. On page 19816—

a. First and second columns, second paragraph from the bottom of the page through the sixth full paragraph; the paragraph beginning with the phrase “61. Section 423.100” through the paragraph ending with the phrase “on various individuals.” is corrected by deleting these paragraphs.

b. In the third column, second full paragraph, in the amendatory statement for § 423.120 (statement number 64), line 6 (immediately following amendatory statement C), the paragraph is corrected by adding the following amendatory statement “D. Adding a new paragraph (b)(2)(vi).”

4. On page 19817, third column, after the third full paragraph ((b)(1)(ix)) which ends with “* * * * *”, the paragraph is corrected by adding the following paragraphs:

“(2) * * *

(vi) Exceptions to paragraph (b)(2)(v) of this section are as follows:

(A) Drug products that are rated as therapeutically equivalent (under the Food and Drug Administration’s most recent publication of “Approved Drug Products with Therapeutic Equivalence Evaluations,” also known as the Orange Book).

(B) Utilization management processes that limit the quantity of drugs due to safety.

(C) Other drugs that CMS specifies through a process that is based upon scientific evidence and medical standards of practice (and, in the case of antiretroviral medications, is consistent with the Department of Health and Human Services Guidelines for the Use of Antiretroviral Agents in HIV-1–Infected Adults and Adolescents) and which permits public notice and comment.

5. On page 19818, second column, fifth paragraph from the bottom (regulations text for § 423.153(d)(1)(vii)(B)), lines 3 and 4, the term “comprehensive medical review” is corrected to read “comprehensive medication review”.

6. On page 19822, in the third column, third paragraph from the bottom of the page, in the amendatory statement for § 423.551 (statement number 84), line 2, the phrase “adding a new paragraph (g) is corrected to read “revising paragraph (g).”

Authority: Catalog of Federal Domestic Assistance Program No. 93.773, Medicare—Hospital Insurance; and Program No. 93.774, Medicare—Supplementary Medical Insurance Program.

Dated: June 4, 2010.

Dawn L. Smalls,

Executive Secretary to the Department.

[FR Doc. 2010–13923 Filed 6–7–10; 4:15 pm]

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GENERAL SERVICES ADMINISTRATION

48 CFR Part 505

[GSAR Amendment 2010–02; GSAR Case 2008–G503 (Change 45) Docket 2008–0007; Sequence 11]

RIN 3090–AI71

General Services Administration Acquisition Regulation; GSAR Case 2008–G503, Rewrite of GSAR Part 505, Publicizing Contract Actions

AGENCIES: Office of Acquisition Policy, General Services Administration (GSA).

ACTION: Final rule.

SUMMARY: The General Services Administration (GSA) is issuing a final rule amending GSA Acquisition Regulation (GSAR) which provides requirements for publicizing contract actions.

DATES: *Effective Date:* June 10, 2010.

FOR FURTHER INFORMATION CONTACT: For clarification of content, contact Beverly Cromer, Procurement Analyst, at (202) 501–1448. For information pertaining to status or publication schedules, contact the Regulatory Secretariat at (202) 501–4755. Please cite Amendment 2010–02, GSAR Case 2008–G503 (Change 45).

SUPPLEMENTARY INFORMATION:

A. Background

The GSA published a proposed rule, with request for comments, in the *Federal Register* at 73 FR 53404 on September 16, 2008. No comments were received in response to the proposed rule. This rule covers the GSAR portion of part 505. Currently, subparts 505.1, 505.2, and 505.5 are identified as “shaded” for regulatory coverage; however, the agency has deemed, these subparts as non-regulatory because the coverage addresses internal agency acquisition policy. These subparts have been revised and are moved to the non-regulatory portion of the GSA Acquisition Manual (GSAM).

This is not a significant regulatory action and, therefore, was not subject to review under Section 6(b) of Executive Order 12866, Regulatory Planning and Review, dated September 30, 1993. This rule is not a major rule under 5 U.S.C. 804.

B. Regulatory Flexibility Act

The General Services Administration certifies that this final rule will not have a significant economic impact on a substantial number of small entities within the meaning of the Regulatory Flexibility Act, 5 U.S.C. 601, *et seq.*, because the revisions are not considered substantive.

C. Paperwork Reduction Act

The Paperwork Reduction Act does not apply because the changes to the GSAR do not impose recordkeeping or information collection requirements, or otherwise collect information from offerors, contractors, or members of the public that require approval of the Office of Management and Budget under 44 U.S.C. chapter 35, *et seq.*

List of Subjects in 48 CFR Part 505

Government procurement.

Dated: May 17, 2010.

Rodney P. Lantier,

Acting Senior Procurement Executive, Office of Acquisition Policy, General Services Administration.

■ Therefore, under the authority of 40 U.S.C. 121(c), GSA removes and reserves 48 CFR part 505.

PART 505 [Removed and Reserved]

[FR Doc. 2010–13902 Filed 6–9–10; 8:45 am]

BILLING CODE 6820–61–S

DEPARTMENT OF TRANSPORTATION

Federal Motor Carrier Safety Administration

49 CFR Parts 390 and 395

Regulatory Guidance Concerning the Preparation of Drivers’ Record of Duty Status To Document Compliance With the Hours-of-Service Requirements

AGENCY: Federal Motor Carrier Safety Administration (FMCSA), DOT.

ACTION: Notice of regulatory guidance.

SUMMARY: The FMCSA announces regulatory guidance concerning the requirement for interstate commercial motor vehicle (CMV) drivers to prepare, in duplicate, a record of duty status for each 24-hour period. FMCSA has determined that the current requirement may be satisfied through the preparation of an original handwritten record, and subsequent electronic submission to the motor carrier of a scanned image of the original record; the driver would retain the original while the carrier maintains the electronic scanned electronic image along with any supporting documents.