CMS Manual System

Pub. 100-04 Medicare Claims Processing

Department of Health & Human Services (DHHS)
Centers for Medicare & Medicaid Services (CMS)

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CHANGE REQUEST 3153

I. SUMMARY OF CHANGES: This instruction revises Table 1 in the Medicare Claims Processing Manual, Chapter 17, Section 20, as published in Change Requests 3022 and 3025 on December 24, 2003.

NEW/REVISED MATERIAL - EFFECTIVE DATE: January 1, 2004 *IMPLEMENTATION DATE: March 26, 2004

Disclaimer for manual changes only: The revision date and transmittal number apply only to red italicized material. Any other material was previously published and remains unchanged.

II. CHANGES IN MANUAL INSTRUCTIONS:

(R = REVISED, N = NEW, D = DELETED)

R/N/D	CHAPTER/SECTION/SUBSECTION/TITLE	
R	17/20/Payment Allowance Limit for Drugs and Biologicals Not Paid on a Cost	
	or Prospective Payment Basis	

*III. FUNDING:

These instructions shall be implemented within your current operating budget.

IV. ATTACHMENTS:

X	Business Requirements
X	Manual Instruction
	Confidential Requirements
	One-Time Notification
	Recurring Update Notification

^{*}Medicare contractors only

Attachment - Business Requirements

SUBJECT: New Payment Allowance Percentages for DMERC Drugs

I. GENERAL INFORMATION

A. Background:

This Change Request (CR) revises Table 1 in the Medicare Claims Processing Manual, Chapter 17, Section 20, as published in CRs 3022 and 3025 dated December 24, 2003, by adding the payment limit percentage for the drug Capecitabine (Xeloda).

B. Policy:

Effective January 1, 2004, J8520 (Capecitabine, 150 mg) and J8521 (Capecitabine, 500 mg) will be paid at 90 percent of the April 1, 2003 average wholesale price.

C. Provider Education:

A provider education article related to this instruction will be available at www.cms.hhs.gov/medlearn/matters shortly after the CR is released. You will receive notification of the article release via the established "medlearn matters" listsery. Contractors shall post this article, or a direct link to this article, on their Web site and include information about it in a listsery message within one week of the availability of the provider education article. In addition, the provider education article must be included in your next regularly scheduled bulletin.

II. BUSINESS REQUIREMENTS

"Shall" denotes a mandatory requirement "Should" denotes an optional requirement

Requirement #	Requirements	Responsibility
3153.1	The payment limit allowance for J8520	FIs and DMERCs
	(Capecitabine, 150 mg) shall be 90 percent of	
	the April 1, 2003 average wholesale price.	
3153.2	The payment limit allowance for J8521	FIs and DMERCs
	(Capecitabine, 500 mg) shall be 90 percent of	
	the April 1, 2003 average wholesale price.	
3153.3	The effective date for the payment allowance	FIs and DMERCs
	percentages is January 1, 2004.	

3153.4	Contractors shall process claims using the following NDC codes: 00004-1100-20 150 mg 00004-1100-51 150 mg 00004-1101-16 500mg 00004-1101-50 500mg	FIs and DMERCs
3153.5	Contractors shall not search their files to either retract payment for claims already paid or to retroactively pay claims. However, contractors shall adjust claims brought to their attention.	FIs and DMERCs
3153.6	A provider education article related to this instruction will be available at www.cms.hhs.gov/medlearn/matters shortly after the CR is released. You will receive notification of the article release via the established "medlearn matters" listserv. Contractors shall post this article, or a direct link to this article, on their Web site and include information about it in a listserv message within one week of the availability of the provider education article. In addition, the provider education article must be included in your next regularly scheduled bulletin.	FIs and DMERCs

III. SUPPORTING INFORMATION AND POSSIBLE DESIGN CONSIDERATIONS

A. Other Instructions: N/A

X-Ref Requirement #	Instructions

B. Design Considerations: N/A

X-Ref Requirement #	Recommendation for Medicare System Requirements

C. Interfaces: N/A

D. Contractor Financial Reporting /Workload Impact: N/A

E. Dependencies: N/A

F. Testing Considerations: N/A

IV. SCHEDULE, CONTACTS, AND FUNDING

Effective Date: January 1, 2004	These instructions should be
Implementation Date: March 26, 2004	implemented within your current operating budget.
Pre-Implementation Contact(s): Appropriate Regional Office	
Post-Implementation Contact(s): Appropriate	
Regional Office	

20 - Payment Allowance Limit for Drugs and Biologicals Not Paid on a Cost or Prospective Payment Basis

(Rev. 131, 03-26-04)

AB-02-075, AB-02-174, PRM 2711.2 B.2, B3-5202, R1799B3

Prior to January 1, 2004, drugs and biologicals not paid on cost or prospective payment are paid based on the lower of the billed charge or 95 percent of the average wholesale price (AWP) as reflected in published sources (e.g., Red Book, Price Alert, etc.). Examples of drugs that are paid on this basis include, but are not limited to, drugs furnished incident to a physician's service, immunosuppressive drugs furnished by pharmacies, drugs furnished by pharmacies under the durable medical equipment benefit, covered oral anticancer drugs, and blood clotting factors.

The Medicare Prescription Drug, Improvement, and Modernization Act of 2003, changed the basis for payment of drugs and biologicals not paid on a cost or prospective payment basis. For January 1, 2004, through December 31, 2004, such drugs or biologicals are paid as described below:

- The payment limits for blood clotting factors will be 95 percent of the AWP.
- The payment limits for new drugs or biologicals will be 95 percent of the AWP. A new drug is defined as an unlisted drug (not currently covered by a HCPCS code) that was FDA approved subsequent to April 1,2003. A drug would not be considered new if: the brand or manufacturer of the drug changed; a new formulation of the vial size is developed; or the drug received a new indication.
- The payment limits for pneumococcal and hepatitis B drugs and biologicals will be 95 percent of the AWP.
- The payment limits for certain drugs studied by the OIG and GAO are based on the percentages of the April 1, 2003 AWPs specified on Table 1 below.
- The payment limits for infusion drugs furnished through an item of implanted durable medical equipment on or after January 1, 2004, will be 95 percent of the October 1, 2003 AWP.
- Drugs and biologicals not described above are paid at 85 percent of the April 1, 2003 AWP.

Payment limits determined under this instruction shall not be updated during 2004.

Table 1: Percentages of April 1, 2003 AWP for Selected Drugs

HCPCS	Applicable Percentage
J0640	80
J1100	86
J1260	80
J1440	81
J1441	81
J1561/J1563	80
J1626	80
J1642	80
J2405	87
J2430	85
J2820	80
J7320	82
J7517	86
J7608	80
J7619	80
J7631	80
J7644	80
J8520/J8521*	90
J9000	80
J9045	81
J9170	80
J9201	80

J9202	80
J9206	80
J9217	81
J9265	81
J9310	81
J9350	84
J9390	81
Q0136	87

* Use the following NDC numbers when processing claims:

00004-1100-20 150 mg

00004-1100-51 150 mg

00004-1101-16 500mg

00004-1101-50 500mg