SUMMARY: Customs and Border Protection (CBP) published in the Federal Register of March 17, 2011, a document which adopted as a final rule, with some changes, interim amendments to the CBP regulations to revise, update, and consolidate the regulatory provisions relating to the country of origin of textile and apparel products. The final rule document contained two errors in the Background portion of the document. The first error concerns an inadvertent reference to imported “antique Persian carpets” in an example prepared by CBP. Because carpets of Iranian-origin are currently prohibited from importation into the United States, the example should not have referenced Persian antique carpets. The example is changed to reflect a non-prohibited article—a Turkish antique carpet. The second error consists of an outdated Internet address that was provided by CBP relating to certain instructions for the completion of CBP Form 7501. This document corrects these two errors.

DATES: Effective on March 24, 2011.

FOR FURTHER INFORMATION CONTACT: Cheryl Williams, Office of Medical Listings Improvement, Social Security Administration, 6401 Security Boulevard, Baltimore, Maryland 21235–6401, (410) 965–1020. For information on eligibility or filing for benefits, call our national toll-free number, 1–800–772–1213, or TTY 1–800–325–0778, or visit our Internet site, Social Security Online, at http://www.socialsecurity.gov.

SUPPLEMENTARY INFORMATION: On June 2, 2010, we published final rules that revised the portions of section 102.00 (the Special Senses and Speech body system for children) regarding the evaluation of hearing loss. The rules became effective on August 2, 2010. When we revised the listings for hearing loss in children, we inadvertently did not also revise listing 111.09C in the neurological disorders body system for children. That listing, which addresses communication impairment associated with a neurological disorder, cross-references to our prior listing for hearing loss in children: listing 102.08. We removed listing 102.08 when we published the 2010 final rules. Since we no longer have that listing, we must correct listing 111.09C.

Before the final rules we published in 2010 became effective, listing 102.08 was the only listing for hearing loss in children. We now have two such listings: listings 102.10 and 102.11. In the notice of proposed rulemaking for the hearing loss listing changes, we explained that we were revising listing 102.08 and changing it to listing 102.10, and that we would use it only for children who do not have cochlear implants. We also explained that we were adding a new listing 102.11 for children who have cochlear implants. Both listings include criteria for children with hearing loss who have communication impairments.

Therefore, we are changing the cross-reference in listing 111.09C to refer to both of the current listings.
Regulatory Procedures

We follow Administrative Procedure Act (APA) rulemaking procedures specified in 5 U.S.C. 553 when we develop regulations. Section 702(a)(5) of the Social Security Act, 42 U.S.C. 902(d)(5). The APA provides exceptions to its notice and public comment procedures when an agency finds there is good cause for dispensing with such procedures because they are impracticable, unnecessary, or contrary to the public interest. 5 U.S.C. 553(b)(B). The change we are making in this rule only corrects an error in a cross-reference. Because the change we are making is only a minor technical correction to a rule to correct an outdated cross-reference, we have determined that the opportunity for prior comment is unnecessary. Therefore, we are issuing this rule as a final rule.

In addition, we find that there is good cause for dispensing with the 30-day delay in the effective date of a substantive rule provided by 5 U.S.C. 553(d)(3). For the reasons already discussed, because this change is a minor technical correction to a rule, a delay in the effective date is unnecessary.

Executive Order 12866, as Supplemented by Executive Order 13563

We consulted with the Office of Management and Budget (OMB) and determined that this final rule does not meet the requirements for a significant regulatory action under Executive Order 12866, as supplemented by Executive Order 13563. Therefore, it was not subject to OMB review.

Regulatory Flexibility Act

We certify that this final rule will not have a significant economic impact on a substantial number of small entities because it affects only individuals. Therefore, a regulatory flexibility analysis was not required under the Regulatory Flexibility Act, as amended.

Paperwork Reduction Act

This rule does not create any new or affect any existing collections, and therefore, does not require Office of Management and Budget approval under the Paperwork Reduction Act. (Catalog of Federal Domestic Assistance Program Nos. 96.001, Social Security—Disability Insurance; 96.002, Social Security—Retirement Insurance; 96.004, Social Security—Survivors Insurance; and 96.006, Supplemental Security Income)

List of Subjects in 20 CFR Part 404

Administrative practice and procedure, Blind, Disability benefits, Old-Age, Survivors, and Disability Insurance, Reporting and recordkeeping requirements, Social Security.

Michael J. Astrue,
Commissioner of Social Security.

For the reasons set out in the preamble, we are amending subpart P of part 404 of chapter III of title 20 of the Code of Federal Regulations as set forth below:

PART 404—FEDERAL OLD-AGE, SURVIVORS AND DISABILITY INSURANCE (1950—

1. The authority citation for subpart P of part 404 continues to read as follows:

Authority: Secs. 202, 205(a)–(b), and (d)–(h), 216(f), 221(a), (i), and (j), 222(c), 223, 225, and 702(a)(5) of the Social Security Act (42 U.S.C. 402, 405(a)–(b), and (d)–(h), 416(i), 421(a), (i), and (j), 422(c), 423, 425, and 902(e)(5)); sec. 211(b), Pub. L. 104–193, 110 Stat. 2105, 2189; sec. 202, Pub. L. 108–203, 118 Stat. 509 (42 U.S.C. 902 note).

2. Amend appendix 1 to subpart P of part 404 by revising section 111.09C of part B to read as follows:

APPENDIX 1 TO SUBPART P OF PART 404—LISTING OF IMPAIRMENTS

* * * * *

Part B

* * * * *

111.00 [Neurological]

* * * * *

111.09 *

C. Impairment of hearing as described under the criteria in 102.10 or 102.11.

* * * * *

[FR Doc. 2011–6983 Filed 3–23–11; 8:45 am]

BILLING CODE 4191–02–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Parts 510 and 529

[Docket No. FDA–2010–N–0002]

New Animal Drugs; Change of Sponsor’s Name and Address; Corrections

AGENCY: Food and Drug Administration, HHS.

ACTION: Correcting amendments.

SUMMARY: The Food and Drug Administration (FDA) published a document in the Federal Register of April 20, 2010 (75 FR 20522) amending the animal drug regulations to reflect changes to a sponsor’s name and address. That document contained errors in the regulatory text. FDA is correcting the tables listing sponsors of approved animal drug applications (NADAs) by adding a change to the sponsor’s drug labeler code (DLC). Cross-references to the sponsor’s DLC are amended in two sections of the Code of Federal Regulations (CFR) containing the conditions of use approved animal drug products. These corrections are being made to improve the accuracy of the animal drug regulations.

DATES: This rule is effective March 24, 2011.

FOR FURTHER INFORMATION CONTACT: George K. Haibel, Center for Veterinary Medicine (HFV–6), Food and Drug Administration, 7519 Standish Pl., Rockville, MD 20855, 240–276–9019, e-mail: george.haibel@fda.hhs.gov.

SUPPLEMENTARY INFORMATION: The Food and Drug Administration (FDA) published a document in the Federal Register of April 20, 2010 (75 FR 20522) amending the animal drug regulations to reflect changes to a sponsor’s name and address. That document contained errors in the regulatory text. FDA is correcting the tables listing sponsors of approved animal drug applications (NADAs) by adding a change to the sponsor’s drug labeler code (DLC). Cross-references to the sponsor’s DLC are amended in two sections of the Code of Federal Regulations (CFR) containing the conditions of use approved animal drug products. These corrections are being made to improve the accuracy of the animal drug regulations.

List of Subjects

21 CFR Part 510

Administrative practice and procedure, Animal drugs, Labeling, Reporting and Recordkeeping requirements.

21 CFR Part 529

Animal drugs.

Accordingly, 21 CFR parts 510 and 529 are corrected by making the following correcting amendments:

PART 510—NEW ANIMAL DRUGS

1. The authority citation for 21 CFR part 510 continues to read as follows:


2. In § 510.600, in the table in paragraph (c)(1), revise the entry for “Piramal Critical Care, Inc.”, and in the table in paragraph (c)(2), remove the