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(c)(5)); sec. 211(b), Pub. L. 104–193, 110 Stat. 2105, 2189; secs. 202, Pub. L. 108–205, 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

111.09

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
entry for “060307” and in numerical sequence, add an entry for “066794” to read as follows:

§ 510.600 Names, addresses, and drug labeler codes of sponsors of approved applications.

- (c) * * *
- (1) * * *


<table>
<thead>
<tr>
<th>Firm name and address</th>
<th>Drug labeler code</th>
</tr>
</thead>
<tbody>
<tr>
<td>Piramal Critical Care, Inc., 3850 Schelden Circle, Bethlehem, PA 18017</td>
<td>066794</td>
</tr>
<tr>
<td>* * * * * * *</td>
<td>* * * * * *</td>
</tr>
</tbody>
</table>

PART 529—CERTAIN OTHER DOSAGE FORM NEW ANIMAL DRUGS

3. The authority citation for 21 CFR part 529 continues to read as follows:


§ 529.1186 [Corrected]

4. In paragraph (b) of § 529.1186, remove “060307, and 065085” and in its place add “065085, and 066794”.

§ 529.2150 [Corrected]

5. In paragraph (b) of § 529.2150, remove “060307” and in its place add “066794”.

Dated: March 17, 2011.

Leslie Kux,
Acting Assistant Commissioner for Policy.

BILLING CODE 4160–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 529

[Docket No. FDA–2010–N–0002]

Certain Other Dosage Form New Animal Drugs; Detomidine; Correction

AGENCY: Food and Drug Administration, HHS.

ACTION: Correcting amendment.

SUMMARY: The Food and Drug Administration (FDA) published a document in the Federal Register of April 23, 2010 (75 FR 21162), that amended the animal drug regulations to reflect approval of an original new animal drug application (NADA). FDA is correcting a paragraph describing limitations to the approved conditions of use for detomidine hydrochloride oromucosal gel in horses. This correction is 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: FDA published a document in the Federal Register of April 23, 2010 (75 FR 21162), that amended the animal drug regulations to reflect approval of an original NADA. FDA is correcting a paragraph describing limitations to the approved conditions of use for detomidine hydrochloride oromucosal gel in horses. This correction is 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 October 26, 2010 (75 FR 65565) amending the animal drug regulations. The October 26, 2010, final rule amended the regulations by removing cross references for use of the withdrawn drugs in combination drug medicated feed. This correction is 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.

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