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) * * *

Firm name and address	Drug labeler code
* * * * *	
Piramal Critical Care, Inc., 3850 Schelden Circle, Bethlehem, PA 18017	066794
* * * * *	

(2) * * *

Drug labeler code	Firm name and address
* * * * *	
066794	Piramal Critical Care, Inc., 3850 Schelden Circle, Bethlehem, PA 18017.
* * * * *	

PART 529—CERTAIN OTHER DOSAGE FORM NEW ANIMAL DRUGS

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

Authority: 21 U.S.C. 360b.

§ 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.

[FR Doc. 2011–6795 Filed 3–23–11; 8:45 am]

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.

List of Subjects in 21 CFR Part 529

Animal drugs.

Accordingly, 21 CFR part 529 is corrected by making the following correcting amendment:

PART 529—CERTAIN OTHER DOSAGE FORM NEW ANIMAL DRUGS

■ 1. The authority citation for part 529 continues to read as follows:

Authority: 21 U.S.C. 360b.

■ 2. In § 529.536, amend paragraph (c)(3) by adding a sentence after the first sentence to read as follows:

§ 529.536 Detomidine.

* * * * *

(c) * * *

(3) * * * Do not use in horses intended for human consumption.

Dated: March 17, 2011.

Leslie Kux,

Acting Assistant Commissioner for Policy.

[FR Doc. 2011–6791 Filed 3–23–11; 8:45 am]

BILLING CODE 4160–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 558

[Docket No. FDA–2010–N–0002]

Animal Drugs, Feeds, and Related Products; Withdrawal of Approval of New Animal Drug Applications; Aklomide; Levamisole Hydrochloride; Nitromide and Sulfanitran; Roxarsone; 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 October 26, 2010 (75 FR 65565) amending the animal drug regulations. The October 26, 2010, final rule amended the regulations by removing those portions that reflect approval of eight new animal drug applications. The final rule inadvertently failed to add conforming amendments in § 558.530. FDA is correcting the animal drug 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.*

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 those portions that reflect approval of eight new animal drug applications. The final rule inadvertently failed to add conforming amendments in § 558.530. FDA is correcting the animal drug regulations by removing cross references for use of the withdrawn drugs in combination