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

drug medicated feed. This correction is being made to improve the accuracy of the animal drug regulations.

List of Subjects in 21 CFR Part 558

Animal drugs, Animal feeds.

Accordingly, 21 CFR part 558 is corrected by making the following correcting amendment:

PART 558—NEW ANIMAL DRUGS FOR USE IN ANIMAL FEEDS

■ 1. The authority citation for 21 CFR part 558 continues to read as follows:

Authority: 21 U.S.C. 360b, 371.

§ 558.530 [Corrected]

■ 2. In § 558.530, remove and reserve paragraphs (d)(4)(i) and (d)(4)(xvii).

Dated: March 17, 2011.

Leslie Kux,

Acting Assistant Commissioner for Policy.

[FR Doc. 2011–6790 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]

New Animal Drugs for Use in Animal Feeds; Florfenicol; Correction

AGENCY: Food and Drug Administration, HHS.

ACTION: Correcting amendments.

SUMMARY: The Food and Drug Administration (FDA) published a document in the **Federal Register** of June 17, 2010 (75 FR 34361) revising the animal drug regulations to reflect approval of a supplemental new animal drug application (NADA). That document contained an incorrect table entry describing the maximum florfenicol concentration in Type B medicated swine feeds. 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 June 17, 2010 (75 FR 34361) revising the animal drug regulations to reflect approval of a supplemental new animal drug application (NADA). That document contained an incorrect table entry describing the maximum florfenicol concentration in Type B medicated swine feeds. This correction is being made to improve the accuracy of the animal drug regulations.

List of Subjects in 21 CFR Part 558

Animal drugs, Animal feeds.

Accordingly, 21 CFR part 558 is corrected by making the following correcting amendments:

PART 558—NEW ANIMAL DRUGS FOR USE IN ANIMAL FEEDS

■ 1. The authority citation for 21 CFR part 558 continues to read as follows:

Authority: 21 U.S.C. 360b, 371.

§ 558.4 [Corrected]

■ 2. In paragraph (d) of § 558.4, in the "Category II" table, in the "Type B maximum (100x)" column, in the entry for "Florfenicol", remove "Swine feed: n/a", "Catfish feed: n/a", and "Salmonid feed: n/a" and in their places add "9.1 g/lb (2.0%)".

Dated: March 17, 2011.

Leslie Kux,

Acting Assistant Commissioner for Policy.

[FR Doc. 2011–6789 Filed 3–23–11; 8:45 am]

BILLING CODE 4160–01–P

POSTAL SERVICE

39 CFR Parts 111 and 121

Combined Mailings of Standard Mail and Periodicals Flats

AGENCY: Postal Service.™

ACTION: Final rule; withdrawal.

SUMMARY: The Postal Service is withdrawing a final rule that would have provided a new option for mailers to combine mailings of Standard Mail ® flats and Periodicals flats within the same bundle, when placed on pallets, and to combine bundles of Standard Mail flats and bundles of Periodicals flats on the same pallet. The Postal Service also withdraws the *Code of Federal Regulations* revision to reflect that Standard Mail service standards apply to all Periodicals flats pieces entered in such combined mailings.

DATES: The final rule published on February 28, 2011 (76 FR 10757), is withdrawn effective March 24, 2011.

FOR FURTHER INFORMATION CONTACT: Jonathan Leon at 202–268–7443, or Kevin Gunther at 202–268–7208.

SUPPLEMENTARY INFORMATION: In a final rule published in the **Federal Register** on February 28, 2011, the Postal Service provided a new option for mailers to combine Standard Mail flats and Periodicals flats, when bundled and placed on pallets. Mailers using this option would have combined different-class mailpieces within the same bundle (comail), or combined separate same-class bundles (of different classes) on the same pallet (copalletize) to maximize presorting or to qualify for deeper destination entry discounts. All mailpieces prepared under this option were required to be bundled and placed on pallets.

In consideration of concerns expressed by members of the mailing community, the Postal Service has elected to withdraw this final rule and will publish these standards as a proposed rule concurrently.

The Postal Service also withdraws the revision to 39 CFR part 121.2 whereby we added a new item "c" to describe the USPS processing of Periodicals mailpieces included in combined mailings of Standard Mail flats and Periodicals flats, and specifying that Periodicals mailpieces included in these mailings will be assigned the service standards applicable to Standard Mail pieces.

Stanley F. Mires,

Chief Counsel, Legislative.

[FR Doc. 2011–6911 Filed 3–23–11; 8:45 am]

BILLING CODE 7710–12–P

ENVIRONMENTAL PROTECTION AGENCY

40 CFR Part 261

[EPA–R03–RCRA–2010–0132; FRL–9285–7]

Hazardous Waste Management System Identification and Listing of Hazardous Waste; Final Exclusion

AGENCY: Environmental Protection Agency.

ACTION: Final rule.

SUMMARY: The Environmental Protection Agency (EPA, also the Agency or we in this preamble) today is granting a petition submitted by Babcock & Wilcox Nuclear Operations Group, Inc., the current owner, and to BWX Technologies, Inc., as predecessor in interest to the current owner, identified collectively hereafter in this preamble as "B&W NOG," to exclude (or delist) on a one-time basis from the lists of hazardous waste, a certain solid waste generated at its Mt. Athos facility near Lynchburg, Virginia.

After careful analysis, we have concluded that the petitioned waste is