

AIRAC date	State	City	Airport	FDC No.	FDC date	Subject
1/9/14	CA	Los Angeles	Los Angeles Intl	3/9143	11/15/13	ILS or LOC Rwy 24R, ILS Rwy 24R (CAT II), ILS Rwy 24R (CAT III), Amdt 24A.
1/9/14	CA	Los Angeles	Los Angeles Intl	3/9144	11/22/13	ILS or LOC Rwy 24R, ILS Rwy 24R (CAT II), ILS Rwy 24R (CAT III), Amdt 24A.

[FR Doc. 2013-30482 Filed 12-26-13; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 520

[Docket No. FDA-2013-N-0002]

Withdrawal of Approval of New Animal Drug Applications; Roxarsone

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule.

SUMMARY: The Food and Drug Administration (FDA) is amending the animal drug regulations to reflect the withdrawal approval of five new animal drug applications (NADAs) for roxarsone oral dosage form products at the sponsor's request because the products are no longer manufactured or marketed.

DATES: Withdrawal of approval is effective January 6, 2014.

FOR FURTHER INFORMATION CONTACT: John Bartkowiak, Center for Veterinary Medicine (HFV-212), Food and Drug Administration, 7519 Standish Pl., Rockville, MD 20855, 240-276-9079, john.bartkowiak@fda.hhs.gov.

SUPPLEMENTARY INFORMATION: Zoetis Inc., 333 Portage St., Kalamazoo, MI 49007 has requested that FDA withdraw approval of the following five NADAs for roxarsone oral dosage form products, used to make medicated drinking water for chickens, turkeys, and swine, because the products are no longer manufactured or marketed:

NADA	Proprietary name
005-414	REN-O-SAL (roxarsone) Tablets.
006-019	Zuco Poultry Tablets.
006-081	Korum Improved Formula.
008-274	Pig Scour Tablets.
093-025	3-NITRO (roxarsone) Soluble.

Elsewhere in this issue of the **Federal Register**, FDA gave notice that approval of NADAs 005-414, 006-019, 006-081,

008-274, and 093-025, and all supplements and amendments thereto, is withdrawn, effective January 6, 2014. As provided in the regulatory text of this document, the animal drug regulations are amended to reflect these voluntary withdrawals of approval.

This rule does not meet the definition of "rule" in 5 U.S.C. 804(3)(A) because it is a rule of "particular applicability." Therefore, it is not subject to the congressional review requirements in 5 U.S.C. 801-808.

List of Subjects in 21 CFR Part 520

Animal drugs.

Therefore, under the Federal Food, Drug, and Cosmetic Act and under the authority delegated to the Commissioner of Food and Drugs and redelegated to the Center for Veterinary Medicine, 21 CFR part 520 is amended as follows:

PART 520—ORAL DOSAGE FORM NEW ANIMAL DRUGS

■ 1. The authority citation for 21 CFR part 520 continues to read as follows:

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

§§ 520.2087, 520.2088, and 520.2089 [Removed]

■ 2. Remove §§ 520.2087, 520.2088, and 520.2089.

Dated: December 20, 2013.

Bernadette Dunham,

Director, Center for Veterinary Medicine.

[FR Doc. 2013-30838 Filed 12-26-13; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 520

[Docket No. FDA-2013-N-0002]

Withdrawal of Approval of New Animal Drug Applications; Roxarsone

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of withdrawal.

SUMMARY: The Food and Drug Administration (FDA) is withdrawing approval of five new animal drug

applications (NADAs) for roxarsone oral dosage form products at the sponsor's request because the products are no longer manufactured or marketed.

DATES: Withdrawal of approval is effective January 6, 2014.

FOR FURTHER INFORMATION CONTACT: John Bartkowiak, Center for Veterinary Medicine (HFV-212), Food and Drug Administration, 7519 Standish Pl., Rockville, MD 20855, 240-276-9079, john.bartkowiak@fda.hhs.gov.

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005-414	REN-O-SAL Tablets.
006-019	Zuco Poultry Tablets.
006-081	Korum Improved Formula.
008-274	Pig Scour Tablets.
093-025	3-NITRO Soluble.

Therefore, under authority delegated to the Commissioner of Food and Drugs and redelegated to the Center for Veterinary Medicine, and in accordance with § 514.116 *Notice of withdrawal of approval of application* (21 CFR 514.116), notice is given that approval of NADAs 005-414, 006-019, 006-081, 008-274, and 093-025, and all supplements and amendments thereto, is hereby withdrawn, effective January 6, 2014.

Elsewhere in this issue of the **Federal Register**, FDA is amending the animal drug regulations to reflect the voluntary withdrawal of approval of these applications.

Dated: December 20, 2013.

Bernadette Dunham,

Director, Center for Veterinary Medicine.

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