

The NADAs listed were identified as being affected by guidance for industry (GFI) #213, “New Animal Drugs and New Animal Drug Combination Products Administered in or on Medicated Feed or Drinking Water of Food-Producing Animals: Recommendations for Drug Sponsors for Voluntarily Aligning Product Use Conditions with GFI #209”, December 2013.

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that approval of NADA 039–077, ANADA 200–140, and ANADA 200–167, and all supplements and amendments thereto, is hereby withdrawn, effective March 31, 2014.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: March 12, 2014.

**Bernadette Dunham,**

*Director, Center for Veterinary Medicine.*

[FR Doc. 2014–05883 Filed 3–19–14; 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–2014–N–0002]

#### **Zoetis Inc., Withdrawal of Approval of New Animal Drug Applications; Chlortetracycline; Sulfathiazole; Penicillin**

**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 of approval of a new animal

drug application (NADA) and two abbreviated new animal drug applications (ANADAs) for three-way, fixed-ratio combination drug Type A medicated articles containing chlortetracycline, sulfathiazole, and penicillin. This action is being taken at the sponsor’s request because these products are no longer manufactured or marketed.

**DATES:** This rule is effective March 31, 2014.

#### **FOR FURTHER INFORMATION CONTACT:**

David Alterman, Center for Veterinary Medicine (HFV–212), Food and Drug Administration, 7519 Standish Pl., Rockville, MD 20855, 240–453–6843.

**SUPPLEMENTARY INFORMATION:** Zoetis Inc., 333 Portage St., Kalamazoo, MI 49007 has requested that FDA withdraw approval of the following NADA and two ANADAs because the products are no longer manufactured or marketed:

| NADA/ANADA    | Proprietary name   |
|---------------|--|
| 039–077 ..... | CSP 250 (chlortetracycline, sulfathiazole, and penicillin) Type A medicated article.               |
| 200–140 ..... | AUREOZOL (chlortetracycline, sulfathiazole, and penicillin) Type A medicated article.              |
| 200–167 ..... | AUREOZOL 500 Granular (chlortetracycline, sulfathiazole, and penicillin) Type A medicated article. |

The NADAs listed were identified as being affected by guidance for industry (GFI) #213, “New Animal Drugs and New Animal Drug Combination Products Administered in or on Medicated Feed or Drinking Water of Food-Producing Animals: Recommendations for Drug Sponsors for Voluntarily Aligning Product Use Conditions with GFI #209”, December 2013.

Elsewhere in this issue of the **Federal Register**, FDA gave notice that approval of NADA 039–077, ANADA 200–140, and ANADA 200–167, and all supplements and amendments thereto, is withdrawn, effective March 31, 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 558**

Animal drugs, Animal feeds.

Therefore, under the Federal Food, Drug, and Cosmetic Act and under authority delegated to the Commissioner

of Food and Drugs and redelegated to the Center for Veterinary Medicine, 21 CFR part 558 is amended as follows:

#### **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 [Amended]**

■ 2. In § 558.4(d), in the “Category II” table, remove the entry for “Sulfathiazole” and its respective following entries.

#### **§ 558.155 [Removed]**

■ 3. Remove § 558.155.

Dated: March 12, 2014.

**Bernadette Dunham,**

*Director, Center for Veterinary Medicine.*

[FR Doc. 2014–05882 Filed 3–19–14; 8:45 am]

**BILLING CODE 4160–01–P**

## DEPARTMENT OF VETERANS AFFAIRS

### 38 CFR Part 17

RIN 2900–AO85

#### **VA Dental Insurance Program—Federalism**

**AGENCY:** Department of Veterans Affairs.  
**ACTION:** Direct final rule; confirmation of effective date.

**SUMMARY:** The Department of Veterans Affairs (VA) published a direct final rule in the **Federal Register** on October 22, 2013, amending its regulations related to the VA Dental Insurance Program (VADIP), a pilot program to offer premium-based dental insurance to enrolled veterans and certain survivors and dependents of veterans. Specifically, this rule adds language to clarify the limited preemptive effect of certain criteria in the VADIP regulations. VA received no comments concerning this rule or its companion substantially identical proposed rule published in the **Federal Register** on October 23, 2013. This document confirms that the direct final rule became effective on December 23, 2013. In a companion document in this issue

of the **Federal Register**, we are withdrawing as unnecessary the proposed rule.

**DATES:** *Effective Date:* The effective date of December 23, 2013, for the final rule published October 22, 2013, 78 FR 62441, is confirmed.

**FOR FURTHER INFORMATION CONTACT:**

Kristin J. Cunningham, Director, Business Policy, Chief Business Office (10NB), Veterans Health Administration, Department of Veterans Affairs, 810 Vermont Ave. NW., Washington, DC 20420; (202) 461-1599. (This is not a toll-free number.)

**SUPPLEMENTARY INFORMATION:** In a direct final rule published in the **Federal Register** on October 22, 2013, 78 FR 62441, VA amended 38 CFR 17.169 to add language to clarify the limited preemptive effect of certain criteria in the VA Dental Insurance Program (VADIP), a pilot program to offer premium-based dental insurance to enrolled veterans and certain survivors and dependents of veterans. VA published a companion substantially identical proposed rule at 78 FR 63143 on October 23, 2013, to serve as a proposal for the provisions in the direct final rule in case adverse comments were received. The direct final rule and proposed rule each provided a 30-day comment period that ended on November 21 and November 22, 2013, respectively. No comments were received.

Under the direct final rule procedures that were described in 78 FR 62441 and 78 FR 63143, the direct final rule became effective on December 23, 2013, because no comments were received within the comment periods. In a companion document in this issue of the **Federal Register**, VA is withdrawing the proposed rulemaking, RIN 2900-AO86, published at 78 FR 63143, as unnecessary.

**Signing Authority**

The Secretary of Veterans Affairs, or designee, approved this document and authorized the undersigned to sign and submit the document to the Office of the Federal Register for publication electronically as an official document of the Department of Veterans Affairs. Jose D. Riojas, Chief of Staff, Department of Veterans Affairs, approved this document on March 11, 2014, for publication.

Dated: March 13, 2014.

**William F. Russo,**

*Deputy Director, Office of Regulation Policy and Management, Office of the General Counsel, Department of Veterans Affairs.*

[FR Doc. 2014-05911 Filed 3-19-14; 8:45 am]

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**DEPARTMENT OF HOMELAND SECURITY**

**Federal Emergency Management Agency**

**44 CFR Part 64**

[Docket ID FEMA-2013-0002; Internal Agency Docket No. FEMA-8325]

**Suspension of Community Eligibility**

**AGENCY:** Federal Emergency Management Agency, DHS.

**ACTION:** Final rule.

**SUMMARY:** This rule identifies communities where the sale of flood insurance has been authorized under the National Flood Insurance Program (NFIP) that are scheduled for suspension on the effective dates listed within this rule because of noncompliance with the floodplain management requirements of the program. If the Federal Emergency Management Agency (FEMA) receives documentation that the community has adopted the required floodplain management measures prior to the effective suspension date given in this rule, the suspension will not occur and a notice of this will be provided by publication in the **Federal Register** on a subsequent date. Also, information identifying the current participation status of a community can be obtained from FEMA's Community Status Book (CSB). The CSB is available at <http://www.fema.gov/fema/csb.shtm>.

**DATES:** The effective date of each community's scheduled suspension is the third date ("Susp.") listed in the third column of the following tables.

**FOR FURTHER INFORMATION CONTACT:** If you want to determine whether a particular community was suspended on the suspension date or for further information, contact David Stearrett, Federal Insurance and Mitigation Administration, Federal Emergency Management Agency, 500 C Street SW., Washington, DC 20472, (202) 646-2953.

**SUPPLEMENTARY INFORMATION:** The NFIP enables property owners to purchase Federal flood insurance that is not otherwise generally available from private insurers. In return, communities agree to adopt and administer local floodplain management measures aimed at protecting lives and new construction from future flooding. Section 1315 of the National Flood Insurance Act of 1968, as amended, 42 U.S.C. 4022, prohibits the sale of NFIP flood insurance unless an appropriate public body adopts adequate floodplain management measures with effective

enforcement measures. The communities listed in this document no longer meet that statutory requirement for compliance with program regulations, 44 CFR Part 59.

Accordingly, the communities will be suspended on the effective date in the third column. As of that date, flood insurance will no longer be available in the community. We recognize that some of these communities may adopt and submit the required documentation of legally enforceable floodplain management measures after this rule is published but prior to the actual suspension date. These communities will not be suspended and will continue to be eligible for the sale of NFIP flood insurance. A notice withdrawing the suspension of such communities will be published in the **Federal Register**.

In addition, FEMA publishes a Flood Insurance Rate Map (FIRM) that identifies the Special Flood Hazard Areas (SFHAs) in these communities. The date of the FIRM, if one has been published, is indicated in the fourth column of the table. No direct Federal financial assistance (except assistance pursuant to the Robert T. Stafford Disaster Relief and Emergency Assistance Act not in connection with a flood) may be provided for construction or acquisition of buildings in identified SFHAs for communities not participating in the NFIP and identified for more than a year on FEMA's initial FIRM for the community as having flood-prone areas (section 202(a) of the Flood Disaster Protection Act of 1973, 42 U.S.C. 4106(a), as amended). This prohibition against certain types of Federal assistance becomes effective for the communities listed on the date shown in the last column. The Administrator finds that notice and public comment procedures under 5 U.S.C. 553(b), are impracticable and unnecessary because communities listed in this final rule have been adequately notified.

Each community receives 6-month, 90-day, and 30-day notification letters addressed to the Chief Executive Officer stating that the community will be suspended unless the required floodplain management measures are met prior to the effective suspension date. Since these notifications were made, this final rule may take effect within less than 30 days.

**National Environmental Policy Act.** This rule is categorically excluded from the requirements of 44 CFR Part 10, Environmental Considerations. No environmental impact assessment has been prepared.

**Regulatory Flexibility Act.** The Administrator has determined that this