

140–340, 140–443, 140–947, 141–090, 200–171, 200–569, 200–570, and 200–580, and all supplements and amendments thereto, is hereby withdrawn, effective December 31, 2016.

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, 2016.

Tracey H. Forfa,

Deputy Director, Center for Veterinary Medicine.

[FR Doc. 2016–31082 Filed 12–23–16; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 573

[Docket No. FDA–2015–F–4282]

Food Additives Permitted in Feed and Drinking Water of Animals; Feed Grade Sodium Formate

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule.

SUMMARY: The Food and Drug Administration (FDA, we, the Agency) is amending the regulations for food additives permitted in feed and drinking water of animals to provide for the safe use of feed grade sodium formate as a feed acidifying agent in complete poultry feeds. This action is in response to a food additive petition filed by BASF Corp.

DATES: This rule is effective December 27, 2016. Submit either written or electronic objections and requests for a hearing by January 26, 2017. See section V of this document for information on the filing of objections.

ADDRESSES: You may submit objections and requests for a hearing as follows:

Electronic Submissions

Submit electronic objections in the following way:

- *Federal eRulemaking Portal:* <https://www.regulations.gov>. Follow the instructions for submitting comments. Objections submitted electronically, including attachments, to <https://www.regulations.gov> will be posted to the docket unchanged. Because your objection will be made public, you are solely responsible for ensuring that your objection does not include any confidential information that you or a

third party may not wish to be posted, such as medical information, your or anyone else's Social Security number, or confidential business information, such as a manufacturing process. Please note that if you include your name, contact information, or other information that identifies you in the body of your objection, that information will be posted on <https://www.regulations.gov>.

- If you want to submit an objection with confidential information that you do not wish to be made available to the public, submit the objection as a written/paper submission and in the manner detailed (see "Written/Paper Submissions" and "Instructions").

Written/Paper Submissions

Submit written/paper submissions as follows:

- *Mail/Hand delivery/Courier (for written/paper submissions):* Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

- For written/paper objections submitted to the Division of Dockets Management, FDA will post your objection, as well as any attachments, except for information submitted, marked and identified, as confidential, if submitted as detailed in "Instructions."

Instructions: All submissions received must include the Docket No. FDA–2015–F–4282 for "Food Additives Permitted in Feed and Drinking Water of Animals; Feed Grade Sodium Formate." Received objections will be placed in the docket and, except for those submitted as "Confidential Submissions," publicly viewable at <https://www.regulations.gov> or at the Division of Dockets Management between 9 a.m. and 4 p.m., Monday through Friday.

- **Confidential Submissions**—To submit an objection with confidential information that you do not wish to be made publicly available, submit your objections only as a written/paper submission. You should submit two copies total. One copy will include the information you claim to be confidential with a heading or cover note that states "THIS DOCUMENT CONTAINS CONFIDENTIAL INFORMATION." The Agency will review this copy, including the claimed confidential information, in its consideration of objections. The second copy, which will have the claimed confidential information redacted/blacked out, will be available for public viewing and posted on <https://www.regulations.gov>. Submit both copies to the Division of Dockets Management. If you do not wish your name and contact information to be

made publicly available, you can provide this information on the cover sheet and not in the body of your objections and you must identify this information as "confidential." Any information marked as "confidential" will not be disclosed except in accordance with 21 CFR 10.20 and other applicable disclosure law. For more information about FDA's posting of comments to public dockets, see 80 FR 56469, September 18, 2015, or access the information at: <http://www.fda.gov/regulatoryinformation/dockets/default.htm>.

Docket: For access to the docket to read background documents or the electronic and written/paper objections received, go to <https://www.regulations.gov> and insert the docket number, found in brackets in the heading of this document, into the "Search" box and follow the prompts and/or go to the Division of Dockets Management, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

FOR FURTHER INFORMATION CONTACT: Chelsea Trull, Center for Veterinary Medicine, Food and Drug Administration, 7519 Standish Pl., Rockville, MD 20855, 240–402–6729, chelsea.trull@fda.hhs.gov.

SUPPLEMENTARY INFORMATION:

I. Background

In a notice published in the **Federal Register** of November 24, 2015 (80 FR 73153), FDA announced that we had filed a food additive petition (animal use) (FAP 2293) submitted by BASF Corp., 100 Park Ave., Florham Park, NJ 07932. The petition proposed that the regulations for food additives permitted in feed and drinking water of animals be amended to provide for the safe use of feed grade sodium formate as a feed acidifying agent in complete poultry feeds.

II. Conclusion

FDA concludes that the data establish the safety and utility of feed grade sodium formate for use as a feed acidifying agent in complete poultry feeds and that the food additive regulations should be amended as set forth in this document.

III. Public Disclosure

In accordance with § 571.1(h) (21 CFR 571.1(h)), the petition and the documents that we considered and relied upon in reaching our decision to approve the petition will be made available for public disclosure (see **FOR FURTHER INFORMATION CONTACT**). As provided in § 571.1(h), we will delete from the documents any materials that are not available for public disclosure.

IV. Environmental Impact

The Agency has determined under 21 CFR 25.32(r) that this action is of a type that does not individually or cumulatively have a significant effect on the human environment. Therefore, neither an environmental assessment, nor an environmental impact statement is required.

V. Objections and Hearing Requests

Any person who will be adversely affected by this regulation may file with the Division of Dockets Management (see ADDRESSES) either electronic or written objections. Each objection shall be separately numbered, and each numbered objection shall specify with particularity the provision of the regulation to which objection is made and the grounds for the objection. Each numbered objection on which a hearing is requested shall specifically so state. Failure to request a hearing for any particular objection shall constitute a waiver of the right to a hearing on that objection. Each numbered objection for which a hearing is requested shall include a detailed description and analysis of the specific factual information intended to be presented in support of the objection in the event that a hearing is held. Failure to include such a description and analysis for any particular objection shall constitute a waiver of the right to a hearing on the objection.

Any objections received in response to the regulation may be seen in the Division of Dockets Management between 9 a.m. and 4 p.m., Monday through Friday, and will be posted to the docket at <https://www.regulations.gov>.

List of Subjects in 21 CFR Part 573

Animal feeds, Food additives.

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 573 is amended as follows:

PART 573—FOOD ADDITIVES PERMITTED IN FEED AND DRINKING WATER OF ANIMALS

■ 1. The authority citation for part 573 continues to read as follows:

Authority: 21 U.S.C. 321, 342, 348.

■ 2. In § 573.696, revise the introductory text and paragraph (b) to read as follows:

§ 573.696 Feed grade sodium formate.

The food additive, feed grade sodium formate, may be safely used in the

manufacture of complete swine and poultry feeds in accordance with the following prescribed conditions:

* * * * *

(b) The additive is used or intended for use as a feed acidifying agent, to lower the pH, in complete swine and poultry feeds at levels not to exceed 1.2 percent of the complete feed.

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Dated: December 20, 2016.

Tracey H. Forfa,

Deputy Director, Center for Veterinary Medicine.

[FR Doc. 2016–31079 Filed 12–23–16; 8:45 am]

BILLING CODE 4164–01–P

AFRICAN DEVELOPMENT FOUNDATION

22 CFR Part 1506

RIN 3005–AA00

Collection of Claims

AGENCY: U.S. African Development Foundation.

ACTION: Final rule.

SUMMARY: The U.S. African Development Foundation (USADF) is revising its regulations on collection of claims in accordance with the Debt Collection Improvement Act of 1996 (DCIA), as implemented by the Department of Justice (Justice) and the Department of the Treasury (Treasury) in the revised Federal Claims Collection Standards (FCCS). The FCCS prescribes the standards that Federal agencies must use in the administrative collection, offset, compromise, and suspension or termination of collection activity for civil claims of money, funds, or property as defined by law.

DATES: This final rule is effective February 27, 2017.

FOR FURTHER INFORMATION CONTACT: June B. Brown, 202–233–8882.

SUPPLEMENTARY INFORMATION:

I. Background

In accordance with the requirements of the DCIA and the implementing regulations promulgated by Justice and Treasury at 31 CFR parts 900–904, USADF is revising its regulations to establish procedures for the administrative collection, offset, compromise, suspension and termination of collection activity for civil claims for money, funds, or property, as defined by 31 U.S.C. 3701(b), and the process by which USADF can refer civil claims to Treasury, Treasury-designated debt

collection centers, or Justice for collection by further administrative action or litigation, as applicable. The regulations do not apply to claims between federal agencies. The rules affect USADF's debtors. The regulations clarify and prescribe the steps USADF must take before initiating debt collection to ensure that individuals' rights are protected. These steps include notifying the debtor of the debt and the consequences of failing to resolve the debt.

II. Section-by-Section Analysis

Subpart A announces the purpose and scope of the regulations, defines terms used in Part 1506, and addresses whether USADF can impose sanctions or remedies other than those prescribed in Part 1506, whether USADF will subdivide a claim exceeding \$100,000, and how claims involving fraud are processed.

Subpart B describes the steps involved in a collection action, including the information USADF includes in a written demand for payment, a debtor's request for review of a claim, the determination of interest, penalty and administrative costs, and the reporting and consequences of delinquent debts.

Subpart C provides for salary offset collection procedures, notice and hearing requirements prior to offset, and USADF's use of offset for claims of another Federal agency.

Subpart D addresses the compromise of debts through reduction or negotiation of the claim amount, joint and several liability on a claim, and releasing the debtor after full payment of a compromised amount.

Subpart E prescribes the circumstances and criteria for USADF to suspend or terminate a collection action.

Subpart F describes the circumstances for USADF to discharge a delinquent debt and reporting a discharge of debt to the Internal Revenue Service.

Subpart G addresses when USADF refers claims to the Department of Justice for litigation.

Subpart H addresses when USADF is required to transfer debts to the Financial Management Service of the Department of the Treasury.

III. Matters of Regulatory Procedure

Executive Order 12866

The proposed regulations have been determined to be non-significant within the meaning of Executive Order 12866.

Regulatory Flexibility Act

The USADF President, in accordance with the Regulatory Flexibility Act, 5