DATES: Effective on April 1, 2005.

FOR FURTHER INFORMATION CONTACT: Patsy A. Bearden, 907–586–7008 or *patsy.bearden@noaa.gov*.

SUPPLEMENTARY INFORMATION:

Background

In January 2004, the U.S. Congress amended section 313(j) of the Magnuson-Stevens Fishery Conservation and Management Act (Magnuson-Stevens Act) through the Consolidated Appropriations Act of 2004 (Pub. L. 108–199, section 801). As amended, section 313(j)(1) requires the Secretary to approve and implement by regulation the so-called Crab Rationalization Program (Program), as it was approved by the North Pacific Fishery Management Council (Council) between June 2002 and April 2003, and all trailing amendments, including those reported to Congress on May 6, 2003. In June 2004, the Council consolidated its actions on the Program into the Council motion, which is contained in its entirety in Amendment 18 to the Fishery Management Plan for Bering Sea/Aleutian Islands King and Tanner Crabs (FMP). Additionally, in June 2004, the Council developed Amendment 19 to the FMP, which represents minor changes necessary to implement the Program. The Notice of Availability for these amendments was published in the **Federal Register** on September 1, 2004 (69 FR 53397). NMFS approved Amendments 18 and 19 on November 19, 2004. NMFS published a proposed rule to implement Amendments 18 and 19 in the **Federal Register** on October 29, 2004 (69 FR 63200). NMFS published a final rule to implement Amendments 18 and 19 on March 2, 2005 (70 FR 10174).

Need for Corrections

In the final rule, published on March 2, 2005 (70 FR 10174), five of the eight OMB control numbers listed in the crab final rule classification section were incorrect. This final rule replaces the incorrect numbers with correct OMB control numbers in the classification section of the final rule and in the regulatory text at 15 CFR part 902.1.

On page 10231, column 2, third heading, replace OMB No. 0648–0272 with OMB No. 0648–0517.

On page 10231, column 3, first heading, replace OMB No. 0648–0503 with OMB No. 0648–0516.

On page 10231, column 3, second heading, replace OMB No. 0648–0504 with OMB No. 0648–0514.

On page 10231, column 3, third heading, replace OMB No. 0648–0503 with OMB No. 0648–0515.

On page 10231, column 3, fourth heading, replace OMB No. 0648 0506 with OMB No. 0648–0518.

Classification

The Administrator, Alaska Region, NMFS (Regional Administrator), has determined that this final rule is necessary for the conservation and management of the BSAI crab fisheries. The Regional Administrator also has determined that this final rule is consistent with the Magnuson-Stevens Act and other applicable laws.

This final rule has been determined to be not significant for the purposes of Executive Order 12866.

Pursuant to 5 U.S.C. 553(b)(B), the Assistant Administrator of Fisheries, NOAA (AA) finds good cause to waive prior notice and opportunity for public comment otherwise required by the section. NOAA finds that prior notice and comment are unnecessary as this rule is purely technical in nature, having no substantive impact whatsoever. This action merely corrects OMB control numbers for approved collections-of-information, in no way altering those approved collections. NOAA finds that because of the non-substantive nature of the correction, no particular public interest exists in this final rule for which there is justification or need for prior notice and opportunity for comment.

Because this correcting amendment does not institute any substantive obligations for the public, the requirement for a 30-day delay in the effective date to this action pursuant to 5 U.S.C. 553(d) does not apply.

Because prior notice and opportunity for public comment are not required for this rule by 5 U.S.C., or any other law, the analytical requirements of the Regulatory Flexibility Act, 5 U.S.C. 601 *et seq.*, are inapplicable.

List of Subjects in 15 CFR Part 902

Reporting and recordkeeping requirements.

Dated: March 11, 2005.

William T. Hogarth

Assistant Administrator for Fisheries, National Marine Fisheries Service.

■ For the reasons set out in the preamble, 15 CFR part 902 is amended as follows:

PART 902—NOAA INFORMATION COLLECTION REQUIREMENTS UNDER THE PAPERWORK REDUCTION ACT; OMB CONTROL NUMBERS

■ 1. The authority citation for part 902 continues to read as follows:

Authority: 44 U.S.C. 3501 *et seq.*

■ 2. In § 902.1, the table in paragraph (b) under 50 CFR is amended by revising in numerical order entries for § 679.5(l)(3)(i), § 679.5(l)(4), § 680.4, § 680.5, § 680.6, § 680.20, § 680.21, § 680.40(f), (g), (h), (i), (j), (k), (l), and (m), § 680.41, § 680.43, and § 680.44 (a) through (f) to read as follows:

§ 902.1 OMB control numbers assigned pursuant to the Paperwork Reduction Act.

* * * * *

(b) * * *

CFR part or section where the information collection requirement is located	Current OMB control number (All numbers begin with 0648-)
* * * * *	
50 CFR	
* * * * *	
679.5(l)(3)(i), (l)(4)	-0272, -0517
* * * * *	
680.4	-0514
680.5	-0515
680.6	-0518
680.20	-0516
680.21	-0514
* * * * *	
680.40(f), (g), (h), (i), (j), (k), (l), and (m)	-0514
680.41	-0514
680.43	-0514
680.44(a), (b), (c), (d), (e)	-0515
680.44(f)	-0514
* * * * *	

[FR Doc. 05–5349 Filed 3–17–05; 8:45 am]

BILLING CODE 3510–22–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Parts 510 and 520

Oral Dosage Form New Animal Drugs; Tiamulin Soluble Powder

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule.

SUMMARY: The Food and Drug Administration (FDA) is amending the animal drug regulations to reflect approval of an abbreviated new animal drug application (ANADA) filed by Phoenix Scientific, Inc. The ANADA provides for use of tiamulin soluble powder to prepare medicated drinking water for the treatment of swine dysentery and swine pneumonia.

DATES: This rule is effective March 18, 2005.

FOR FURTHER INFORMATION CONTACT: Daniel A. Benz, Center for Veterinary Medicine (HFV–104), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–827–0223, e-mail: *daniel.benz@fda.gov*.

SUPPLEMENTARY INFORMATION: Phoenix Scientific, Inc., 3915 South 48th Street Ter., St. Joseph, MO 64503, filed a supplement to ANADA 200–344 that provides for use of Tiamulin Soluble Antibiotic to prepare medicated drinking water for the treatment of swine dysentery and swine pneumonia. Phoenix Scientific, Inc.'s Tiamulin Soluble Antibiotic is approved as a generic copy of Boehringer Ingelheim Vetmedica, Inc.'s DENAGARD (tiamulin) Soluble Antibiotic approved under NADA 134–644. The ANADA is approved as of February 16, 2005, and the regulations are amended in 21 CFR 520.2455 to reflect the approval. The basis of approval is discussed in the freedom of information summary.

FDA is also amending the regulations in 21 CFR 520.2455 to reflect a more recent genus name for the causative pathogen for swine dysentery and in the tables in 21 CFR 510.600(c) to reflect accepted style for the sponsor's street address. These actions are being taken to improve the accuracy of the regulations.

In accordance with the freedom of information provisions of 21 CFR part 20 and 21 CFR 514.11(e)(2)(ii), a summary of safety and effectiveness data and information submitted to support approval of this application may be seen in the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852, between 9 a.m. and 4 p.m., Monday through Friday.

The agency has determined under 21 CFR 25.33(a)(1) 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.

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

21 CFR Part 510

Administrative practice and procedure, Animal drugs, Labeling, Reporting and recordkeeping requirements.

21 CFR Part 520

Animal drugs.

■ 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 parts 510 and 520 are amended as follows:

PART 510—NEW ANIMAL DRUGS

■ 1. The authority citation for 21 CFR part 510 continues to read as follows:

Authority: 21 U.S.C. 321, 331, 351, 352, 353, 360b, 371, 379e.

§ 510.600 [Amended]

■ 2. Section 510.600 is amended in the table in paragraph (c)(1) in the entry for ''Phoenix Scientific, Inc.'' and in the table in paragraph (c)(2) in the entry for ''059130'' by removing ''St. Terrace'' and by adding in its place ''Street Ter.''.

PART 520—ORAL DOSAGE FORM NEW ANIMAL DRUGS

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

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

§ 520.2455 [Amended]

■ 4. Section 520.2455 is amended in paragraph (b) by removing ''*Sponsor.* See No. 000010'' and by adding in its place ''*Sponsors.* See Nos. 000010 and 059130''; and in paragraph (d)(1)(i) by removing ''*Treponema*'' and by adding in its place ''*Brachyspira*''.

Dated: March 9, 2005.

Stephen F. Sundlof,

Director, Center for Veterinary Medicine.

[FR Doc. 05–5380 Filed 3–17–05; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 573

Food Additives Permitted in Feed and Drinking Water of Animals; Poly(2–vinylpyridine-co-styrene); Salts of Volatile Fatty Acids

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule; technical amendment.

SUMMARY: The Food and Drug Administration (FDA) is amending the animal food additive regulations to correct the specifications for two food additives used in cattle feed. Incorrect symbols describing permitted levels of heavy metals such as lead and arsenic are being corrected with text to reflect the maximum permitted levels of these two impurities in these food additives. This action is being taken to improve the accuracy of the agency's regulations.

DATES: This rule is effective March 18, 2005.

FOR FURTHER INFORMATION CONTACT: Sharon Benz, Center for Veterinary Medicine (HFV–220), Food and Drug Administration, 7519 Standish Pl., Rockville, MD 20855, 240–453–6864, e-mail: *sbenz@cvm.fda.gov*.

SUPPLEMENTARY INFORMATION: FDA has found that part 573 (21 CFR part 573) of the Code of Federal Regulations does not accurately reflect the approved specifications for two food additives used in cattle feed, poly(2–vinylpyridine-co-styrene) and salts of volatile fatty acids. The greater than symbols in the tables describing the permitted levels of heavy metals such as lead and arsenic were incorrect. FDA is amending the regulations in §§ 573.870 and 573.914 to correctly reflect the maximum permitted levels of these two impurities in these food additives. This action is being taken to improve the accuracy of the agency's regulations.

Publication of this document constitutes final action on these changes under the Administrative Procedure Act (5 U.S.C. 553). Notice and public procedure are unnecessary because FDA is merely correcting nonsubstantive errors.

List of Subjects in 21 CFR 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