ii. Paragraph (g)(3), each time it appears; and
iii. Paragraph (g)(4).

§ 92.104 [Amended]

14. Section 92.104 is amended by removing the word ''ornithosis'' and adding the word ''chlamydiosis'' in its place, in the following places:
(a) Paragraph (b)(2);
(b) Paragraph (b)(3);
(c) Paragraph (c)(3);
(d) Paragraph (c)(4);
(e) Paragraph (d)(3); and
(f) Paragraph (d)(4).

§ 92.106 [Amended]

15. In § 92.106, paragraph (c)(5)(iii), Cooperative and Trust Fund Agreement Between _____ (Name of Reporter) and the United States Department of Agriculture, Animal and Plant Health Inspection Service, is amended as follows:

a. In paragraph (A)(17), the words ''velogenic viscerotropic Newcastle disease'' are removed and the words ''exotic Newcastle disease'' are added in their place; and

b. The term ''VVND'' is removed and the term ''END'' is added in its place in the following places:
i. Paragraph (B)(4); and
ii. Paragraph (B)(5).

§ 92.209 [Amended]

16. In § 92.209, paragraph (a)(2) is redesignated as paragraph (b) and is amended by removing the words ''viscerotropic velogenic Newcastle disease'' and adding in their place the words ''exotic Newcastle disease'', and the paragraph designative (1) is removed in paragraph (a).

PART 94—RINDERPEST, FOOT-AND-MOUTH DISEASE, FOWL PEST (FOWL PLAGUE), EXOTIC NEWCASTLE DISEASE, AFRICAN SWINE FEVER, HOG CHOLERA, AND BOVINE SPONGIFORM ENCEPHALOPATHY: PROHIBITED AND RESTRICTED IMPORTATIONS.

17. The authority citation for part 94 continues to read as follows:

Authority: 7 U.S.C. 147a, 150ee, 161, 162, and 450; 19 U.S.C. 1306; 21 U.S.C. 111, 114a, 134a, 134b, 134c, 134f, 136, and 136a; 31 U.S.C. 9701; 42 U.S.C. 4331, 4332; 7 CFR 2.22, 2.80, and 371.2(d).

PART 94—AMENDED

18. The heading for part 94 is revised to read as set forth above.

19. In § 94.0, the definition of *Exotic Newcastle disease (VVND)* is removed and a definition of *Exotic Newcastle disease (END)* is added, in alphabetical order, to read as follows:

§ 94.0 Definitions.

* * * * *

Exotic Newcastle disease (END). Any velogenic Newcastle disease. Exotic Newcastle disease is an acute, rapidly spreading, and usually fatal viral disease of birds and poultry.

* * * * *

§ 94.6 [Amended]

20. Section 94.6 is amended as follows:

a. The term ''VVND'' is removed and the term ''END'' is added in its place in the following places:
i. The heading;
ii. Paragraph (a) introductory text;
iii. Paragraph (a)(1);
iv. Paragraph (a)(2);
v. Paragraph (c) introductory text, each time it appears;
vi. Paragraph (d) introductory text, each time it appears;
vii. Paragraph (d)(1)(ix) introductory text;
viii. Paragraph (d)(1)(ix)(A);
ix. Paragraph (d)(1)(ix)(B);
x. Paragraph (d)(1)(ix)(C) introductory text;
xi. Paragraph (d)(1)(ix)(C)(*1*);
xii. Paragraph (d)(1)(ix)(C)(*2*), each time it appears;
xiii. Paragraph (d)(2);
xiv. Paragraph (d)(3), both times it appears; and
xv. Paragraph (d)(4), both times it appears.

b. The term ''viscerotropic velogenic Newcastle disease'' is removed and the term ''END'' is added in its place in the following places:
i. Paragraph (c)(2); and
ii. Paragraph (c)(5).

PART 161—REQUIREMENTS AND STANDARDS FOR ACCREDITED VETERINARIANS AND SUSPENSION OR REVOCATION OF SUCH ACCREDITATION

21. The authority citation for part 161 continues to read as follows:

Authority: 15 U.S.C. 1828; 21 U.S.C. 105, 111–114, 114a, 114a–1, 115, 116, 120, 121, 125, 134b, 134f, 612, and 613; 7 CFR 2.22, 2.80, and 371.2(d).

§ 161.2 [Amended]

22. In § 161.2, paragraph (d)(6) is amended by removing the words ''psittacosis or ornithosis, and velogenic viscerotropic Newcastle disease'' and adding the words ''chlamydiosis and exotic Newcastle disease'' in their place.

Done in Washington, DC, this 29th day of October 1996.

A. Strating,

Acting Administrator, Animal and Plant Health Inspection Service.

[FR Doc. 96–28060 Filed 11–4–96; 8:45 am]

BILLING CODE 3410–34–P

SECURITIES AND EXCHANGE COMMISSION

17 CFR Part 200

[Release No. 34–37893]

Delegation of Authority to the General Counsel

AGENCY: Securities and Exchange Commission.

ACTION: Final rule.

SUMMARY: The Securities and Exchange Commission is amending its rules to delegate authority to the General Counsel to refer matters and information concerning possible professional misconduct to state bar associations and other state professional boards or societies.

EFFECTIVE DATE: November 5, 1996.

FOR FURTHER INFORMATION CONTACT: Barbara B. Hannigan, Ethics Counsel, at 942–0970.

SUPPLEMENTARY INFORMATION: The Securities and Exchange Commission (''Commission'') today announced amendments to its rules governing delegation of authority to the General Counsel.

The amendment to Rule 30–14 [1] authorizes the General Counsel to refer matters and information concerning possible professional misconduct to state bar associations and other state professional boards or societies.

Notwithstanding this delegation of authority, in instances where a referral of possible professional misconduct presents any unusual or noteworthy issues, the delegation would not be exercised and the matter would be submitted to the Commission.

The Commission finds, in accordance with Section 553(b)(3)(A) of the Administrative Procedure Act,[2] that this amendment relates solely to agency organization, procedure, or practice, and does not relate to a substantive rule. Accordingly, notice and opportunity for public comment are unnecessary, and publication of the amendment 30 days before its effective date is also unnecessary.

List of Subjects in 17 CFR Part 200

Administrative practice and procedure, Authority delegations (Government agencies).

Text of Amendment

For the reasons set out in the preamble, Title 17, Chapter II of the Code of Federal Regulations is amended as follows:

[1] 17 CFR 200.30–14.

[2] 5 U.S.C. 553(b)(3)(A).

PART 200—ORGANIZATION; CONDUCT AND ETHICS; AND INFORMATION AND REQUESTS

1. The authority citation for Part 200 continues to read in part as follows:

Authority: 15 U.S.C. 77s, 78d-1, 78d-2, 78w, 78ll(d), 79t, 77sss, 80a-37, 80b-11, unless otherwise noted.

* * * * *

2. Section 200.30–14 is amended by adding paragraph (k) to read as follows:

§ 200.30–14 Delegation of authority to the General Counsel.

* * * * *

(k) To refer matters and information concerning possible professional misconduct to state bar associations and other state professional boards or societies.

Dated: October 30, 1996.

By the Commission.

Margaret H. McFarland,

Deputy Secretary.

[FR Doc. 96–28386 Filed 11–4–96; 8:45 am]

BILLING CODE 8010–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 178

[Docket No. 93F–0101]

Indirect Food Additives: Adjuvants, Production Aids, and Sanitizers; Technical Amendment

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule; technical amendment.

SUMMARY: The Food and Drug Administration (FDA) is amending the food additive regulations to correct an error in the Chemical Abstracts Service (CAS) registry number for a component of a food additive. This document corrects that error.

EFFECTIVE DATE: November 5, 1996.

FOR FURTHER INFORMATION CONTACT: John R. Bryce, Center for Food Safety and Applied Nutrition (HFS–216), Food and Drug Administration, 200 C St. SW., Washington, DC 20204, 202–418–3023.

SUPPLEMENTARY INFORMATION: In the Federal Register of August 21, 1995 (60 FR 43370), the agency amended the food additive regulations to provide for the safe use of monomethyltin/dimethyltin isooctylmercaptoacetates as a stabilizer in rigid polyvinyl chloride and rigid vinyl chloride copolymers for use in contact with food. The CAS registry number for dimethyltin bis(2-ethylhexylmercaptoacetate was incorrectly published as ''(CAS Reg. No. 57583–35–43)'' instead of ''(CAS Reg. No. 57583–35–4)''. Accordingly, the agency is amending 21 CFR 178.2010 to correct the error.

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

List of Subjects in 21 CFR Part 178

Food additives, Food packaging.

Therefore, under the Federal Food, Drug, and Cosmetic Act and under authority delegated to the Commissioner of Food and Drugs and redelegated to the Director, Center for Food Safety and Applied Nutrition, 21 CFR part 178 is amended as follows:

PART 178—INDIRECT FOOD ADDITIVES: ADJUVANTS, PRODUCTION AIDS, AND SANITIZERS

1. The authority citation for 21 CFR part 178 continues to read as follows:

Authority: Secs. 201, 402, 409, 721 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 321, 342, 348, 379e).

§ 178.2010 [Amended]

2. Section 178.2010 *Antioxidants and/or stabilizers for polymers* is amended in the table in paragraph (b) under the heading ''Substances'' in the entry for ''Dimethyltin/monomethyltin isooctylmercaptoacetates'' by removing ''CAS Reg. No. 57583–35–43'' and adding in its place ''CAS Reg. No. 57583–35–4''.

Dated: October 16, 1996.

Fred R. Shank,

Director, Center for Food Safety and Applied Nutrition.

[FR Doc. 96–28290 Filed 11–4–96; 8:45 am]

BILLING CODE 4160–01–F

21 CFR Parts 520 and 556

Animal Drugs, Feeds, and Related Products; Enrofloxacin Oral Solution

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 a new animal drug application (NADA) filed by Bayer Corp. The NADA provides for the use of drinking water medicated with enrofloxacin for the control of mortality associated with certain bacteria in chickens and turkeys.

EFFECTIVE DATE: November 5, 1996.

FOR FURTHER INFORMATION CONTACT: George K. Haibel, Center for Veterinary Medicine (HFV–133), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–594–1644.

SUPPLEMENTARY INFORMATION: Bayer Corp., Agriculture Division, Animal Health, P.O. Box 390, Shawnee Mission, KS 66201, filed NADA 140–828 that covers Baytril® (enrofloxacin) 3.23% Concentrate Antimicrobial Solution. The concentrate is added to drinking water to produce a final concentration of 25 to 50 parts per million. The medicated drinking water is used in chickens for the control of mortality associated with *Escherichia coli* susceptible to enrofloxacin and in turkeys for the control of mortality associated with *E. coli* and *Pasteurella multocida* (fowl cholera) susceptible to enrofloxacin. The NADA is approved as of October 4, 1996, and the regulations are amended by adding new § 520.813 to reflect the approval. The regulations are also amended to provide for a tolerance for enrofloxacin residues in chickens and turkeys in new § 556.228. The drug product is available on a prescription basis. The basis of approval is discussed in the freedom of information summary.

In accordance with the freedom of information (FOI) provisions of part 20 (21 CFR part 20) and § 514.11(e)(2)(ii) (21 CFR 514.11(e)(2)(ii)), a summary of safety and effectiveness data and information submitted to support approval of this application (FOI summary) may be seen in the Dockets Management Branch (HFA–305), Food and Drug Administration, 12420 Parklawn Dr., rm. 1–23, Rockville, MD 20857, between 9 a.m. and 4 p.m., Monday through Friday. The FOI summary is also electronically available on the Center for Veterinary Medicine's home page on the World Wide Web (http://www.cvm.fda.gov/). The summaries are located in the section entitled, ''FDA CVM Documents and Databases—Information and Resources Library.''

Under section 512(c)(2)(F)(ii) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 360b(c)(2)(F)(ii)), this approval qualifies for 3 years of marketing exclusivity beginning October 4, 1996, because the NADA contains reports of new clinical or field investigations and new human food safety studies (other than bioequivalence or residue studies) essential to the approval of the