leavened baked snack foods at levels not to exceed 400 IU of vitamin D_2 per 100 g in the finished food). The labeling requirement in § 172.381(d) ensures that when vitamin D_2 bakers yeast is used to make products, the manufacturer will have the information necessary to use the additive in conformance with the provisions of the regulation.

The second objection from AB Mauri asserts that if FDA is going to approve vitamin D_2 supplementation in baked products at higher levels than are currently permitted by the regulations, it should do so in a way that permits better control of vitamin D levels in finished products by considering the use of vitamin D_3 instead. AB Mauri questions whether vitamin D_2 is as effective for humans as vitamin D_3 at similar levels, and cites two peer-reviewed journal articles to support this claim.

Our evaluation of the petition was based solely on the safety of the proposed use of vitamin D_2 bakers yeast in yeast-containing baked goods. Therefore, expanding the scope of the final rule to provide for the safe use of vitamin D_3 is beyond the scope of the petition submitted by Lallemand. If AB Mauri is interested in obtaining approval for the expanded use of vitamin D_3 in food, they may do so by petitioning FDA for this use in accordance with section 409(b) of the FD&C Act.

IV. Summary and Conclusions

Section 409 of the FD&C Act requires that a food additive be shown to be safe prior to marketing. Under 21 CFR 170.3(i), a food additive is ''safe'' if there is a reasonable certainty in the minds of competent scientists that the substance is not harmful under the intended conditions of use. In the final rule authorizing the use of vitamin D_2 bakers yeast, we concluded that the data presented by the petitioner to establish safety of the additive demonstrate that vitamin D_2 bakers yeast is safe for its intended use in yeast-leavened baked products at levels not to exceed 400 IU of vitamin D_2 per 100 g in the finished food.

The petitioner has the burden to demonstrate the safety of the additive to gain FDA approval. Once we make a finding of safety, the burden shifts to an objector, who must come forward with evidence that calls into question our conclusion (see section 409(f)(1) of the FD&C Act). After evaluating the objections from AB Mauri, we have concluded that the objections do not provide any basis for us to reconsider our decision to issue the final rule authorizing the use of vitamin D_2 bakers yeast as a dual purpose nutrient supplement and leavening agent in yeast-containing baked products at levels not to exceed 400 IU of vitamin D_2 per 100 g in the finished food. Accordingly, we are not making any changes in response to the objections.

Dated: March 4, 2014.

Leslie Kux,

Assistant Commissioner for Policy.

[FR Doc. 2014–05060 Filed 3–10–14; 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]

New Animal Drugs; Change of Sponsor

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule; technical amendment.

SUMMARY: The Food and Drug Administration (FDA) is amending the animal drug regulations to reflect a change of sponsor for 110 approved new animal drug applications (NADAs) and 14 approved abbreviated new animal drug applications (ANADAs) for new animal drug for use in animal feed from Pfizer, Inc., including its several subsidiaries and divisions, to Zoetis, Inc.

DATES: This rule is effective March 11, 2014.

FOR FURTHER INFORMATION CONTACT: Steven D. Vaughn, Center for Veterinary Medicine (HFV–100), Food and Drug Administration, 7520 Standish Pl., Rockville, MD 20855; 240–276–8300, *steven.vaughn@fda.hhs.gov*.

SUPPLEMENTARY INFORMATION: Pfizer, Inc., 235 E. 42d St., New York, NY 10017, and its wholly owned subsidiaries Alpharma, LLC; Fort Dodge Animal Health, Division of Wyeth; Fort Dodge Animal Health, Division of Wyeth Holdings Corp.; and its division, Pharmacia & Upjohn Co., have informed FDA that they have transferred ownership of, and all rights and interest in, the 110 approved NADAs and 14 approved ANADAs in Table 1 to Zoetis, Inc., 333 Portage St., Kalamazoo, MI 4900.

TABLE 1—NADAS AND ANADAS TRANSFERRED FROM PFIZER, INC., TO ZOETIS, INC.

File No.	Product name
007–616	HISTOSTAT 50 (nitarsone) Type A Medicated Article.
011–116	ZOAMIX (zoalene) Type A Medicated Article.
012–375	ALBAMIX (novobiocin) Type A Medicated Article.
012–680	PHARMASTATIN 20 (nystatin) Type A Medicated Article.
013–747	Zoalene 90 Medicated Coccidiostat.
033–950	Sulfamerazine In Fish Grade.
034–085	LINCOMIX (lincomycin) Type A Medicated Article.
034–254	MGA (melengestrol acetate) Type A Medicated Article.
035–688	AUREOMIX Granular 500 (pen G, CTC, sulfamethazine) Type A Medicated Article.
035–805	AUREO S 700 Granular (CTC and sulfamethazine) Type A Medicated Article.
036–361	Amprolium and ethopabate/CTC (chlortetracycline)/sodium sulfate.
039–077	CSP (chlortetracycline, sulfathiazole, and penicillin G procaine) 250 and 500 Type A Medicated Articles.
039–402	MGA 500 (melengestrol acetate) Liquid Type A Medicated Article.
039–417	DECCOX (decoquinate) Type A Medicated Article.
040–209	ROFENAID 40 (sulfadimethoxine and ormetoprim) Type A Medicated Article.
041–647	AUREOMIX S 700–A (CTC and sulfamethazine) Type A Medicated Article.
041–648	AUREOMIX S 700–D (CTC and sulfamethazine) Type A Medicated Article.
041–649	AUREOMIX S 700–G (CTC and sulfamethazine) Type A Medicated Article.

TABLE 1—NADAS AND ANADAS TRANSFERRED FROM PFIZER, INC., TO ZOETIS, INC.—Continued

File No.	Product name
041–650	AUREOMIX S 700–E (CTC and sulfamethazine) Type A Medicated Article.
041–651	AUREOMIX S 700–F (CTC and sulfamethazine) Type A Medicated Article.
041–652	AUREOMIX S 700–C–2 (CTC and sulfamethazine) Type A Medicated Article.
041–653	AUREOMIX S 700–B (CTC and sulfamethazine) Type A Medicated Article.
041–654	AUREOMIX S 700–H (CTC and sulfamethazine) Type A Medicated Article.
044–820	AMPROL PLUS (amprolium and ethopabate)/LINCOMIX (lincomycin).
044–972	LINCOMIX (lincomycin)/COYDEN (clopidol).
045–348	ALBAC (bacitracin zinc)/DECCOX (decoquinate).
045–444	CHLORMAX (chlortetracycline)/DECCOX (decoquinate).
046–415	Tylosin Type A Medicated Articles.
046–592	BMD (bacitracin methylene disalicylate) Type A Medicated Articles.
046–666	Penicillin G Procaine Type A Medicated Articles.
046–699	CHLORMAX (chlortetracycline) Type A Medicated Articles.
046–718	MGA (liquid) (melengestrol acetate)/TERRAMYCIN (oxytetracycline).
046–719	MGA (dry) (melengestrol acetate)/TERRAMYCIN (oxytetracycline).
046–920	BACIFERM (bacitracin zinc) Type A Medicated Articles.
047–261	DECCOX (decoquinate)/LINCOMIX (lincomycin).
047–262	DECCOX (decoquinate)/LINCOMIX (lincomycin).
048–486	ROBENZ (robenidine) Type A Medicated Article.
048–761	AUREOMYCIN (chlortetracycline) Type A Medicated Article.
048–762	Chlortetracycline Type A Medicated Article.
048–763	Chlortetracycline Type A Medicated Article.
048–954	ZOAMIX (zoalene)/LINCOMIX (lincomycin).
049–287	PFICHLOR (chlortetracycline) Type A Medicated Article.
055–040	SF Mix 66 (chlortetracycline) Type A Medicated Article.
065–020	Micro CTC (chlortetracycline) 100 Type A Medicated Article.
091–668	CHLORMAX SP (chlortetracycline) 250 and 500 Type A Medicated Articles.
091–749	TYLAN 40 Plus Sulfa-G.
092–482	COBAN (monensin)/LINCOMIX (lincomycin).
092–507	ROBENZ/Aureomycin 500 Gm.
092–522	COBAN (monensin)/LINCOMIX (lincomycin).
093–106	ROBENZ (robenidine)/LINCOMIX (lincomycin).
096–298	AVATEC and BOVATEC (lasalocid) Type A Medicated Articles.
096–933	ROBENZ (robenidine)/Bacitracin Zn.
097–085	Bacitracin MD/ROBENZ (robenidine).
097–505	LINCOMIX (lincomycin) Type A Medicated Articles.
098–452	ALBAC 50 (bacitracin zinc) Type A Medicated Article.
100–901	PFICHLOR (chlortetracycline) 100S Milk Replacer Type A Medicated Article.
101–689	AVATEC (lasalocid)/LINCOMIX (lincomycin).
103–758	TERRAMYCIN (oxytetracycline) Type A Medicated Article.
107–347	CHEQUE (mibolerone) Medicated Dog Food.
107–996	AVATEC (lasalocid)/FORTRACIN (lasalocid).
114–794	AMPROL HI–E (amprolium and ethopabate)/BACIFERM (bacitracin zinc).
121–553	AUREOMYCIN (chlortetracycline)/COBAN (monensin).
124–309	MGA 100 and 200 (melengestrol acetate)/RUMENSIN (monensin).
125–476	MGA 500 (melengestrol acetate)/RUMENSIN (monensin).
128–686	BIO–COX (salinomycin) Type A Medicated Article.
133–334	Virginiamycin Type A Medicated Article.
134–284	BIO–COX (salinomycin)/FLAVOMYCIN (bambermycins).
134–830	ALBAC (bacitracin zinc)/COBAN (monensin).
135–746	BIO–COX (salinomycin)/BMD (bacitracin methylene disalicylate).
137–537	BIO–COX (salinomycin)/LINCOMIX (lincomycin).
138–456	COBAN (monensin)/BMD (bacitracin methylene disalicylate).
138–792	MGA 100 and 200 (melengestrol)/RUMENSIN (monensin)/TYLAN (tylosin).
138–870	MGA 500 (melengestrol acetate)/BOVATEC (lasalocid)/TYLAN (tylosin).
138–904	MGA (melengestrol acetate)/BOVATEC (lasalocid)/TYLAN (tylosin).
138–941	LINCOMIX (lincomycin)/BANMINTH (pyrantel).
138–992	MGA 200 (melengestrol acetate)/BOVATEC (lasalocid)/TYLAN (tylosin).
138–995	MGA (melengestrol acetate)/TYLAN (tylosin).
139–075	CYGRO (maduramicin) Type A Medicated Article.
139–192	MGA 500 (melengestrol acetate)/TYLAN (tylosin).
139–235	BACIFERM (bacitracin zinc)/BIO–COX (salinomycin).
139–876	MGA 200 (melengestrol acetate)/BOVATEC (lasalocid).
140–288	MGA 500 (melengestrol acetate)/BOVATEC (lasalocid).
140–443	HYGROMIX 1.6 (hygromycin B) Premix.
140–579	BOVATEC (lasalocid)/TERRAMYCIN (oxytetracycline).
140–853	BMD (bacitracin methylene disalicylate)/MONTEBAN (naracin).
140–859	AUREOMYCIN (chlortetracycline)/BIO–COX (salinomycin).
140–865	BACIFERM or ALBAC (bacitracin zinc)/MONTEBAN (naracin).
141–025	CATTLYST (laidlomycin) Type A Medicated Article.
141–059	BMD (bacitracin methylene disalicylate)/CTC (chlortetracycline).
141–083	AVATEC (lasalocid)/BACIFERM (bacitracin zinc).

TABLE 1—NADAS AND ANADAS TRANSFERRED FROM PFIZER, INC., TO ZOETIS, INC.—Continued

File No.	Product name
141–085	BMD (bacitracin methylene disalicylate)/ZOAMIX (zoalene).
141–088	HISTOSTAT (nitarsone)/BMD (bacitracin methylene disalicylate).
141–102	BMD (bacitracin methylene disalicylate)/DECCOX (decoquinate).
141–109	AVATEC (lasalocid)/BACIFERM (bacitracin zinc).
141–124	BMD (bacitracin methylene disalicylate)/MAXIBAN (naracin and nicarbazin).
141–132	HISTOSTAT (nitarsone)/ALBAC (bacitracin zinc).
141–136	BMD (bacitracin methylene disalicylate)/BIO–COX (salinomycin).
141–140	BMD (bacitracin methylene disalicylate)/COBAN (monensin).
141–144	BMD (bacitracin methylene disalicylate)/SAFE–GUARD (fenbendazole).
141–147	DECCOX (decoquinate)/CHLORMAX (chlortetracycline).
141–148	DECCOX (decoquinate)/RUMENSIN (monensin).
141–149	DECCOX (decoquinate)/RUMENSIN (monensin)/TYLAN (tylosin).
141–150	AVATEC (lasalocid)/STAFAC (virginiamycin).
141–154	BMD (bacitracin methylene disalicylate)/ROBENZ (robenidine).
141–156	BMD (bacitracin methylene disalicylate)/AMPROL (amprolium).
141–179	BMD (bacitracin methylene disalicylate)/AVATEC (lasalocid).
141–181	ALBAC (bacitracin zinc)/AVATEC (lasalocid).
141–185	DECCOX (decoquinate)/AUREOMYCIN (chlortetracycline).
141–201	CATTLYST (laidlomycin)/AUREOMYCIN (chlortetracycline).
141–250	BOVATEC (lasalocid)/AUREOMYCIN (chlortetracycline).
200–140	AUREOZOL (pen G, CTC, sulfathiazole) Type A Medicated Article.
200–167	AUREOZOL 500 Granular (pen G, CTC, sulfathiazole) Type A Medicated Article.
200–204	ALBAC (bacitracin zinc)/BIO–COX (salinomycin).
200–205	ALBAC (bacitracin zinc)/AMPROL HI–E (amprolium and ethopabate).
200–210	ALBAC (bacitracin zinc)/SACOX (salinomycin).
200–212	ALBAC (bacitracin zinc)/ROBENZ.
200–213	ALBAC (bacitracin zinc)/DECCOX (decoquinate).
200–218	ALBAC (bacitracin zinc)/COYDEN 25 (clopidol).
200–223	ALBAC 50 (bacitracin zinc) Type A Medicated Article.
200–242	BMD (bacitracin methylene disalicylate)/AUREOMYCIN (chlortetracycline).
200–261	CHLORMAX (chlortetracycline)/BIO–COX (salinomycin).
200–262	CHLORMAX (chlortetracycline)/SACOX (salinomycin).
200–263	CHLORMAX (chlortetracycline)/COBAN (monensin).
200–478	ALBAC 50 (bacitracin zinc)/NICARB (nicarbazin).

Accordingly, the Agency is amending the regulations in 21 CFR part 558 to reflect these transfers of ownership. In addition, the regulations are being amended to make minor corrections. This is being done to increase the accuracy and readability of the regulations.

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.

§ 558.76 [Amended]

■ 2. In § 558.76, in paragraph (a), remove ''046573'' and in its place add ''054771''; and in the table in paragraph (d)(1), in the ''Limitations'' column and in the ''Sponsor'' column, remove ''046573'' wherever it occurs and in its place add ''054771''.

§ 558.78 [Amended]

■ 3. In § 558.78, in paragraph (b), remove ''046573'' and in its place add ''054771''; and in the table in paragraph (d)(1), in the ''Sponsor'' column, remove ''046573'' wherever it occurs and in its place add ''054771''; and in paragraph (d)(2)(ii), remove ''046573'' and in its place add ''054771''.

§ 558.128 [Amended]

■ 4. Amend § 558.128 as follows:
■ a. In paragraph (b)(1), remove ''046573'' and in its place add ''054771'';
■ b. In the tables in paragraphs (e)(1), (e)(2), (e)(3), and (e)(5), in the ''Sponsor'' column, remove ''046573'' wherever it occurs and in its place add ''054771'';
■ c. In the table in paragraph (e)(4), in the ''Limitations'' column and in the ''Sponsor'' column, remove ''046573'' wherever it occurs and in its place add ''054771'';
■ d. In paragraph (e)(6)(iv), remove ''per head'' and in its place add ''per pound of body weight''; and
■ e. In paragraph (e)(6)(v), remove ''046573'' and in its place add ''054771''.

§ 558.140 [Amended]

■ 5. In § 558.140, in paragraph (a), remove ''046573'' and in its place add ''054771''; and revise paragraph (c)(3) to read as follows:

§ 558.140 Chlortetracycline and sulfamethazine.

* * * * *

(c) * * *

(3) *Limitations.* Feed for 28 days, withdraw 7 days prior to slaughter. A withdrawal period has not been established for this product in pre-ruminating calves. Do not use in calves to be processed for veal.

§ 558.145 [Amended]

■ 6. In § 558.145, in paragraph (a)(1), remove ''046573'' and in its place add ''054771''; and in paragraph (a)(2) remove ''046573 and 048164'' and in its place add ''Nos. 048164 and 054771''.

§ 558.155 [Amended]

■ 7. In § 558.155, in paragraphs (a)(1) and (2), remove ''046573'' and in its place add ''054771''.

§ 558.175 [Amended]

■ 8. In § 558.175, in the table in paragraph (d), in the ''Limitations'' column and in the ''Sponsor'' column, remove ''046573'' wherever it occurs and in its place add ''054771''; and in paragraph (d)(6), in the ''Sponsor'' column, remove ''000009'' and in its place add ''054771''.

§ 558.195 [Amended]

■ 9. In § 558.195, in paragraph (b), remove ''046573'' and in its place add ''054771''; in the tables in paragraphs (e)(1), (e)(2), and (e)(3), in the ''Limitations'' column and in the ''Sponsor'' column, remove ''046573'' wherever it occurs and in its place add ''054771''; and in the table in paragraph (e)(1), in the ''Limitations'' column and in the ''Sponsor'' column, remove ''000009'' wherever it occurs and in its place add ''054771''.

§ 558.198 [Amended]

■ 10. In § 558.198, in the tables in paragraphs (d)(1) and (2), in the ''Limitations'' column, remove ''046573'' wherever it occurs and in its place add ''054771''.

§ 558.258 [Amended]

■ 11. In § 558.258, in the tables in paragraphs (e)(2)(vi) and (vii), in the ''Limitations'' column and in the ''Sponsor'' column, remove ''046573'' and in its place add ''054771''; and in paragraphs (e)(2)(ii) through (v), in the ''Limitations'' column, remove ''000009'' and in its place add ''054771''.

§ 558.305 [Amended]

■ 12. In § 558.305, in paragraph (b), remove ''046573'' and in its place add ''054771''; and in the table in paragraph (e) in the ''Sponsor'' column, remove ''046573'' wherever it occurs and in its place add ''054771''.

§ 558.311 [Amended]

■ 13. In § 558.311, in paragraphs (b)(1) through (4) and (6) and (7), remove ''046573'' and in its place add ''054771''; in the table in paragraph (e)(1), in the ''Limitations'' column and in the ''Sponsor'' column, remove ''046573'' wherever it occurs and in its place add ''054771'', in the ''Limitations'' column, remove ''000009'' where it occurs and in its place add ''054771'', in the ''Limitations'' column, remove ''000004'' where it occurs and in its place add ''054771''; and in paragraphs (e)(2)(v), (e)(3)(v), and (e)(4)(v), remove ''046573'' and in its place add ''054771''.

§ 558.325 [Amended]

■ 14. In § 558.325, in paragraphs (a)(1) and (c)(3)(i), remove ''000009'' and in its place add ''054771''; and in the tables in paragraphs (d)(1) and (2), in the ''Sponsor'' column, remove ''000009'' wherever it occurs and in its place add ''054771''.

§ 558.340 [Amended]

■ 15. In § 558.340, in paragraph (a), remove ''046573'' and in its place add ''No. 054771''.

§ 558.342 [Amended]

■ 16. In § 558.342, in paragraph (b)(1), remove ''000009'' and in its place add ''054771''; in the table in paragraph (e)(1), in the ''Limitations'' column and in the ''Sponsor'' column, remove ''000009'' wherever it occurs and in its place add ''054771'', and in the ''Limitations'' column, remove ''046573'' wherever it occurs and in its place add ''054771''.

§ 558.348 [Amended]

■ 17. In § 558.348, in paragraph (a), remove ''000009'' and in its place add ''No. 054771''.

§ 558.355 [Amended]

■ 18. In § 558.355, in paragraphs (b)(8), (b)(9), and (b)(11), remove ''046573'' and in its place add ''054771''; in paragraphs (f)(1)(iii)(*b*), (f)(1)(iv)(*b*), (f)(1)(v)(*b*), (f)(1)(vii)(*b*), (f)(1)(xiv)(*b*), (f)(1)(xxv)(*b*), (f)(1)(xxix)(*b*), (f)(1)(xxx)(*b*), (f)(2)(ii)(*b*), (f)(2)(iii)(*b*), (f)(4)(ii)(*b*), (f)(4)(iii)(*b*), (f)(4)(iv)(*b*), (f)(4)(v)(*b*), remove ''046573'' and in its place add ''054771''; and revise paragraph (f)(1)(i)(*b*) to read as follows:

§ 558.355 Monensin.

* * * * *

(f) * * *

(1) * * *

(i) * * *

(*b*) *Limitations.* Feed continuously as the sole ration. In the absence of coccidiosis, the use of monensin with no withdrawal period may limit feed intake resulting in reduced weight gain. Do not feed to laying chickens.

* * * * *

§ 558.363 [Amended]

■ 19. In § 558.363, in paragraphs (a)(6) and (a)(7), remove ''046573'' and in its place add ''054771''; in paragraph (d)(1)(vi)(B) and (d)(1)(x)(B), remove ''046573'' and in its place add ''054771''.

§ 558.364 [Amended]

■ 20. In § 558.364, in paragraph (a), remove ''000009'' and in its place add ''054771''; and in the table in paragraph (d), in the ''Sponsor'' column, remove ''000009'' wherever it occurs and in its place add ''054771''.

§ 558.366 [Amended]

■ 21. In § 558.366, in the table in paragraph (d), in the ''Limitations'' column and in the ''Sponsor'' column, remove ''046573'' wherever it occurs and in its place add ''054771'' and in paragraph (d), in the ''Limitations'' column, remove ''000009'' wherever it occurs and in its place add ''054771''.

§ 558.369 [Amended]

■ 22. In § 558.369, in paragraph (a), remove ''046573'' and in its place add ''054771''.

§ 558.415 [Amended]

■ 23. In § 558.415, redesignate paragraphs (b) and (c) as paragraphs (c) and (d); revise paragraph (a); and add new paragraph (b) to read as follows:

§ 558.415 Novobiocin.

■ (a) *Specifications.* Type A medicated article containing 25 grams of novobiocin activity per pound.

(b) *Sponsor.* See No. 054771 in § 510.600(c) of this chapter.

* * * * *

■ 24. In § 558.430, redesignate paragraphs (b) and (c) as paragraphs (c) and (d); revise paragraph (a); and add new paragraph (b) to read as follows:

§ 558.430 Nystatin.

(a) *Specifications.* Type A medicated article containing 20 grams of nystatin activity per pound.

(b) *Sponsor.* See No. 054771 in § 510.600(c) of this chapter.

* * * * *

§ 558.460 [Amended]

■ 25. In § 558.460, in paragraph (b), remove ''046573'' and in its place add ''054771''; and in the table in paragraph (d)(1)(i), in the ''Sponsor'' column, remove ''000069, 046573'' and in its place add ''054771, 066104''; and in paragraphs (d)(1)(ii) and (iii), in the ''Sponsor'' column, remove ''Do.'' and in its place add ''054771, 066104''.

§ 558.464 [Amended]

■ 26. In § 558.464, in paragraphs (a)(1) and (2), remove ''000069'' and in its place add ''054771''.

§ 558.485 [Amended]

■ 27. In § 558.485, in paragraph (b)(7), remove ''000069 and 017135'' and in its place add ''017135 and 054771''; and in paragraph (e)(1)(xii)(C), remove ''000009'' and in its place add ''054771''.

§ 558.500 [Amended]

■ 28. In § 558.500, in paragraphs (e)(2)(viii) and (x), in the ''Limitations'' column, remove ''000009'' and in its place add ''054771''.

§ 558.515 [Amended]

■ 29. In § 558.515, in paragraph (a), remove ''046573'' and in its place add ''054771''; in the table in paragraph (d), in the ''Sponsor'' column, remove ''046573'' wherever it occurs and in its place add ''054771'' and in the ''Sponsor'' column, remove ''000009'' where it occurs and in its place add ''054771''.

§ 558.550 [Amended]

■ 30. In § 558.550, in paragraphs (b)(1), (d)(1)(iii)(*c*), (d)(1)(vi)(*c*), (d)(1)(vii)(*c*), (d)(1)(xvi)(*c*), (d)(1)(xx)(C), (d)(1)(xxi)(C), (d)(1)(xxii)(B), (d)(1)(xxiii)(*b*), (d)(3)(ii)(B), (d)(3)(iii)(B), (d)(3)(v)(B), (d)(4)(i)(*b*), remove ''046573'' and in its place add ''054771''; and in paragraph (d)(1)(xiii)(*c*), remove ''000009'' and in its place add ''054771''.

§ 558.555 [Amended]

■ 31. In § 558.555, in paragraphs (d)(2), (d)(3), (d)(4), and (d)(8), in the ''Limitations'' column, remove ''046573'' and in its place add ''054771''.

§ 558.575 [Amended]

■ 32. In § 558.575, in paragraph (a)(1), remove ''046573'' and in its place add ''054771''.

§ 558.582 [Amended]

■ 33. In § 558.582, in paragraph (a), remove ''046573'' and in its place add ''054771''.

§ 558.600 [Amended]

■ 34. In paragraph (e)(1)(iii) of § 558.600, in the ''Indications for use'' column, remove ''susceptible'' wherever it occurs and in its place add ''sensitive''; in the ''Limitations'' column, add ''Use as only source of tiamulin.''; and in the ''Limitations'' column, remove ''046573'' and in its place add ''054771''.

§ 558.625 [Amended]

■ 35. In § 558.625, remove and reserve paragraphs (b)(57) and (83); in paragraph (b)(54), remove ''046573'' and in its place add ''054771''; and add paragraph (b)(10) to read as follows:

§ 558.625 Tylosin.

* * * * *

(b) * * *

(10) To No. 012286: 0.4, 0.8, and 1.6 grams per pound, paragraph (f)(1)(vi)(*a*) of this section; 20, 40, and 100 grams per pound, paragraphs (f)(1)(i) through (vi) of this section.

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§ 558.630 [Amended]

■ 36. In § 558.630, in paragraph (b)(5), remove ''046573'' and in its place add ''054771''.

§ 558.635 [Amended]

■ 37. In § 558.635, in paragraph (a)(2), remove ''046573'' and in its place add ''054771''.

§ 558.680 [Amended]

■ 38. In § 558.680, in paragraph (b), remove ''046573'' and in its place add ''054771''.

Dated: February 28, 2014.

Steven D. Vaughn,

Director, Office of New Animal Drug Evaluation, Center for Veterinary Medicine.

[FR Doc. 2014–04937 Filed 3–10–14; 8:45 am]

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EQUAL EMPLOYMENT OPPORTUNITY COMMISSION

29 CFR Part 1625

RIN 3046–AA58

Waivers of Rights and Claims in Settlement of a Charge or Lawsuit Under the Age Discrimination in Employment Act; Corrections

AGENCY: Equal Employment Opportunity Commission.

ACTION: Correcting Amendments.

SUMMARY: The EEOC is correcting a cross-reference in its regulation concerning the requirements for a valid waiver of an individual's right to file a lawsuit under the Older Workers Benefit Protection Act (OWBPA) amendments to the Age Discrimination in Employment Act (ADEA). This is a technical correction.

DATES: *Effective Date:* March 11, 2014.

FOR FURTHER INFORMATION CONTACT: Carol R. Miaskoff, Assistant Legal Counsel, or Raymond L. Peeler, Senior Attorney-Advisor, at (202) 663–4640 (voice) or (202) 663–7026 (TTY). Requests for this document in an alternative format should be made to the Office of Communications and Legislative Affairs at (202) 663–4191 (voice) or (202) 663–4494 (TTY), or the Publications Information Center at 1–800–669–3362 (toll free).

SUPPLEMENTARY INFORMATION:

Background

In the Older Workers Benefit Protection Act of 1990 (OWBPA), Congress established requirements for the knowing and voluntary release of claims under the Age Discrimination in Employment Act (ADEA).[1] The OWBPA set basic requirements for all waivers of ADEA rights, and it imposed extra requirements when employers sought such waivers in connection with an exit incentive or group termination program. To implement the OWBPA, the EEOC issued a final negotiated rule at 29 CFR 1625.22 in 1998.[2]

Need for Correction

The EEOC now corrects a cross-reference in 29 CFR 1625.22(g)(3), the provision that states the basic requirements for waiving ADEA rights when settling an ADEA charge or lawsuit. Where subsection (g)(3) should cross reference the rule's ''knowing and voluntary'' requirements applicable to all ADEA waivers, it instead references the rule's additional requirements for *group termination programs.* Therefore, the EEOC now replaces the incorrect language in 29 CFR 1625.22(g)(3) (''set out in paragraph (f) of this section''), with language referencing the rule's general waiver requirements (''set out in paragraphs (b), (c), and (d) of this section.'').

Changes to Authority Citation

This rule also contains several changes to the existing authority citation for 29 CFR Part 1625. Some of these changes update existing citations to comply with **Federal Register** formatting conventions. Others streamline and consolidate several references to the Age Discrimination in Employment Act. The revisions also add Executive Order 12067 due to its discussion of the EEOC's leadership role in age discrimination in employment and the EEOC's responsibilities with respect to federal regulations concerning employment discrimination.

Retrospective Regulatory Review

Although the EEOC's rulemaking on waivers of rights and claims under the

[1] Public Law 101–433, 104 Stat. 978 (codified at 29 U.S.C. 626(f)).

[2] 63 FR 30624, 30631 (June 5, 1998) (EEOC Final Rule for Waiver of Rights and Claims under the ADEA).