

(4) *Before January 1, 2001, if you worked in a sheltered workshop.* Before January 1, 2001, if you worked in a sheltered workshop or a comparable facility especially set up for severely impaired persons, we will ordinarily consider that your earnings from this work show that you have engaged in substantial gainful activity if your earnings averaged more than the amounts in the table in paragraph (b)(2) of this section. Average monthly earnings from a sheltered workshop or a comparable facility that are equal to or less than those amounts indicated in table 1 of paragraph (b)(2) of this section will ordinarily show that you have not engaged in substantial gainful activity without the need to consider other information, as described in paragraph (b)(6) of this section, regardless of whether they are more or less than those indicated in paragraph (b)(3) of this section. When your earnings from a sheltered workshop or comparable facility are equal to or less than those amounts indicated in table 1 of paragraph (b)(2), we will consider the provisions of paragraph (b)(6) of this section only if there is evidence showing that you may have engaged in substantial gainful activity. For work performed in a sheltered workshop in months beginning January 2001, the rules of paragraphs (b)(2), (3), and (6) apply the same as they do to any other work done by an employee.

* * * * *

(6) *Earnings that are not high enough to ordinarily show that you engaged in substantial gainful activity.*

(i) *Before January 1, 2001,* if your average monthly earnings were between the amounts shown in paragraphs (b)(2) and (3) of this section, we will generally consider other information in addition to your earnings (see paragraph (b)(6)(iii) of this section). This rule generally applies to employees who did not work in a sheltered workshop or a comparable facility, although we may apply it to some people who work in sheltered workshops or comparable facilities (see paragraph (b)(4) of this section).

(ii) *Beginning January 1, 2001,* if your average monthly earnings are equal to or less than the amounts determined under paragraph (b)(2) of this section, we will generally not consider other information in addition to your earnings unless there is evidence indicating that you may be engaging in substantial gainful activity or that you are in a position to defer or suppress your earnings.

(iii) *Examples of other information we may consider include, whether—*

(A) Your work is comparable to that of unimpaired people in your

community who are doing the same or similar occupations as their means of livelihood, taking into account the time, energy, skill, and responsibility involved in the work, and

(B) Your work, although significantly less than that done by unimpaired people, is clearly worth the amounts shown in paragraph (b)(2) of this section, according to pay scales in your community.

* * * * *

3. The authority citation for Subpart K of Part 416 continues to read as follows:

Authority: Secs. 702(a)(5), 1602, 1611, 1612, 1613, 1614(f), 1621, and 1631 of the Social Security Act (42 U.S.C. 902(a)(5), 1381a, 1382, 1382a, 1382b, 1382c(f), 1382j, and 1383); sec. 211, Pub. L. 93–66, 87 Stat. 154 (42 U.S.C. 1382 note).

4. Section 416.1112 is amended by revising paragraph (c)(3) to read as follows:

§ 416.1112 Earned income we do not count.

* * * * *

(c) * * *

(3) If you are a blind or disabled child who is a student regularly attending school as described in § 416.1861:

(i) *For earned income beginning January 1, 2002,* monthly and yearly maximum amounts that are the larger of:

(A) The monthly and yearly amounts for the previous year, or

(B) Monthly and yearly maximum amounts increased for changes in the cost-of-living, calculated in the same manner as the Federal benefit rates described in § 416.405, except that we will use the calendar year 2001 amounts as the base amounts and will round the resulting amount to the next higher multiple of \$10 where such amount is a multiple of \$5 but not of \$10 and to the nearest multiple of \$10 in any other case.

(ii) *For earned income before January 1, 2002,* the amounts indicated in Table 1 of this section.

TABLE 1

For months	Up to per month	But not more than in a calendar year
In calendar years before 2001	\$400	\$1,620
In calendar year 2001	1,290	5,200

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 558

New Animal Drugs for Use in Animal Feeds; Decoquinatate and Monensin

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 Alpharma, Inc. The NADA provides for use of approved, single-ingredient decoquinatate and monensin Type A medicated articles to make two-way combination drug Type B and Type C medicated feeds used for prevention of coccidiosis and improved feed efficiency in cattle fed in confinement for slaughter.

DATES: This rule is effective December 29, 2000.

FOR FURTHER INFORMATION CONTACT:

Janis R. Messenheimer, Center for Veterinary Medicine (HFV–135), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–827–7578.

SUPPLEMENTARY INFORMATION: Alpharma Inc., One Executive Dr., P.O. Box 1399, Fort Lee, NJ 07024, filed NADA 141–148 that provides for use of DECCOX® (27.2 gram per pound (g/lb) decoquinatate) and Rumensin® (20, 30, 45, 60, 80, or 90.7 g/lb monensin activity as monensin sodium) Type A medicated articles to make two-way combination Type B and Type C medicated feeds. The Type C medicated feeds contain 13.6 to 27.2 g/ton decoquinatate and 5 to 30 g/ton monensin, and are used for prevention of coccidiosis caused by *Eimeria bovis* and *E. zuernii*, and improved feed efficiency in cattle fed in confinement for slaughter. The NADA is approved as of November 16, 2000, and the regulations in 21 CFR 558.195 and 558.355 are amended to reflect the approval. The basis for approval is discussed in the freedom of information summary.

In accordance with the freedom of information provisions of 21 CFR part 20 and 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 Dockets Management Branch (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)(2) 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 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.

2. Section 558.195 is amended in the table in paragraph (d) by adding an entry following "13.6 to 27.2 (0.0015 to 0.003 pct)" and before "Chlortetracycline approximately 400" to read as follows:

§ 558.195 Decoquinatate.

* * * * *

(d) * * *

Decoquinatate in grams per ton	Combination in grams per ton	Indications for use	Limitations	Sponsor
*	*	*	*	*
	Monensin 5 to 30	Cattle fed in confinement for slaughter; for prevention of coccidiosis caused by <i>Eimeria bovis</i> and <i>E. zuernii</i> , and improved feed efficiency.	Feed only to cattle fed in confinement for slaughter. Feed continuously as the sole ration to provide 22.7 mg of decoquinatate per 100 lb body weight per day and 50 to 360 mg of monensin per head per day. Feed at least 28 days during period of exposure to coccidiosis or when it is likely to be a hazard. Do not feed to animals producing milk for food. Also see (c)(1) of this paragraph and § 558.355(d)(8). Monensin as monensin sodium provided by 000986 in § 510.600(c) of this chapter.	046573
*	*	*	*	*

3. Section 558.355 is amended by adding paragraph (f)(7) to read as follows:

§ 558.355 Monensin.

* * * * *

(f) * * *

(7) Monensin may also be used in combination with decoquinatate as in § 558.195.

Dated: December 20, 2000.

Stephen F. Sundlof,

Director, Center for Veterinary Medicine.

[FR Doc. 00–33217 Filed 12–28–00; 8:45 am]

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DEPARTMENT OF TRANSPORTATION

Federal Highway Administration

23 CFR Part 777

[FHWA Docket No. FHWA–97–2514; 96–8]

RIN 2125–AD78

Mitigation of Impacts to Wetlands and Natural Habitat

AGENCY: Federal Highway Administration (FHWA), DOT.

ACTION: Final rule.

SUMMARY: This document revises the rule concerning the eligibility for Federal-aid transportation funding of activities to mitigate impacts to wetlands and natural habitats due to highway projects funded pursuant to provisions of title 23, U.S. Code. It updates the FHWA's wetlands regulation to conform with wetland and natural habitat mitigation provisions contained in the Intermodal Surface Transportation Efficiency Act of 1991 (ISTEA) and the Transportation Equity Act for the 21st Century (TEA–21),

which allow increased flexibility for Federal funding participation under title 23, U.S. Code, in mitigation measures for impacts of federally funded highway projects to wetlands and natural habitats

EFFECTIVE DATE: January 29, 2001.

FOR FURTHER INFORMATION CONTACT: Mr. Paul Garrett, Office of Natural Environment, (303) 969–5772, ext. 332, email address:

paul.garrett@fhwa.dot.gov; FHWA, 555 Zang Street, Lakewood, CO 80228, office hours are from 8 a.m. to 5 p.m., m.t., Monday through Friday, except Federal holidays; or Mr. Robert J. Black, Office of the Chief Counsel, HCC–30, (202) 366–1359, email address:

robert.black@fhwa.dot.gov, 400 Seventh Street, SW., Washington, D.C. 20590–0001. Office hours are from 7:45 a.m. to 4:15 p.m., e.t., Monday through Friday, except Federal holidays.

SUPPLEMENTARY INFORMATION:

Electronic Access

Internet users may access all comments received by the U.S. DOT Dockets, Room PL–401, by using the universal resource locator (URL): <http://>