Education, and Welfare, 200 C St. SW., Washington, DC 20204, 202–472–5690.

SUPPLEMENTARY INFORMATION: A notice published in the **Federal Register** of August 15, 1978 (43 FR 36143) announced that a food additive petition (FAP 8B3381) had been filed by Morton Chemical Co., 110 N. Wacker Dr., Chicago, IL 60606, proposing that the food additive regulations be amended to provide for the use of 2-sulfoethyl methacrylate, sodium salt as a component of food-contact coatings on metal and polyester film.

The Food and Drug Administration (FDA) has evaluated data in the petition and other relevant material and concludes that §§ 175.300 and 177.1630 (21 CFR 175.300 and 177.1630) should be amended as set forth below.

Therefore, under the Federal Food, Drug, and Cosmetic Act (sec. 409(c)(1), 72 Stat. 1786 (21 U.S.C. 348(c)(1))) and under authority delegated to the Commissioner of Food and Drugs (21 CFR 5.1), Parts 175 and 177 are amended as follows:

PART 175—INDIRECT FOOD ADDITIVES: ADHESIVE COATINGS AND COATINGS

1. Part 175 is amended in § 175.300 by alphabetically inserting a new item in the list of substances in paragraph (b)(3)(xxxiii) to read as follows:

§ 175.300 Resinous and polymeric coatings.

* * * * *

(b) * * *

(3) * * *

(xxxiii) * * *

2-Sulfoethyl methacrylate, sodium salt (CAS Reg. No. 10595–80–9). For use only in copolymer coatings on metal under conditions of use E, F, and G described in table 2 of paragraph (d) of this section, and limited to use at a level not to exceed 2.0 percent by weight of the dry copolymer coating.

* * * * *

PART 177—INDIRECT FOOD ADDITIVES: POLYMERS

2. Part 177 is amended in § 177.1630 by alphabetically inserting a new item in the list of substances in paragraph (e)(4)(iii) to read as follows:

§ 177.1630 Polyethylene phthalate polymers.

* * * * *

(e) * * *

(4) * * *

(iii) * * *

2-Sulfoethyl methacrylate, sodium salt (CAS Reg. No. 10595–80–9). For use only in copolymer coatings on polyethylene phthalate film under conditions of use E, F, and G described in table 2 of § 175.300(d) of this chapter, and limited to use at a level not to exceed 2.0 percent by weight of the dry copolymer coating.

* * * * *

Any person who will be adversely affected by the foregoing regulation may at any time on or before August 13, 1979, submit to the Hearing Clerk (HFA–305), Food and Drug Administration, Rm. 4–65, 5600 Fishers Lane, Rockville, MD 20857, written objections thereto and may make a written request for a public hearing on the stated 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. 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. Four copies of all documents shall be submitted and shall be identified with the Hearing Clerk docket number found in brackets in the heading of this regulation. Received objections may be seen in the above office between the hours of 9 a.m. and 4 p.m., Monday through Friday.

Effective date. This regulation shall become effective July 13, 1979.

(Sec. 409(c)(1), 72 Stat. 1786 (21 U.S.C. 348(c)(1)).)

Dated: July 3, 1979.

Sanford A. Miller,

Director, Bureau of Foods.

[FR Doc. 79–21335 Filed 7–12–79; 8:45 am]

BILLING CODE 4110–03–M

---

21 CFR Part 558

New Animal Drugs for use in Animal Feeds; Nicarbazin, Roxarsone and Lincomycin; Nicarbazin With Roxarsone; Nicarbazin With Lincomycin

AGENCY: Food and Drug Administration.

ACTION: Final rule.

SUMMARY: The animal drug regulations are amended to reflect approval of three new animal drug applications (NADA's) filed by Merck Sharp & Dohme Research Labs., providing for use of medicated broiler feeds containing nicarbazin in combination with roxarsone and lincomycin, or roxarsone, or lincomycin, for increased rate of weight gain and as an aid in prevention of coccidiosis.

EFFECTIVE DATE: July 13, 1979.

FOR FURTHER INFORMATION CONTACT: Lonnie W. Luther, Bureau of Veterinary Medicine (HFV–147), Food and Drug Administration, Department of Health, Education, and Welfare, 5600 Fishers Lane, Rockville, MD 20857, 301–443–4317.

SUPPLEMENTARY INFORMATION: Merck Sharp & Dohme Research Labs., Division of Merck & Co., Inc., P.O. Box 2000, Rahway, NJ 07065, filed three NADA's (107–997, 108–115, 108–116) providing for safe and effective use of a medicated broiler feed containing the above-mentioned drug combinations. Approval of these applications relies on certain portions of the efficacy and safety data contained in NADA's 9–476 for nicarbazin, 7–891 for roxarsone, and 34–085 for lincomycin. The Director of the Bureau of Veterinary Medicine concludes that the approval of these combination NADA's poses no increased risk from exposure to residues of the new animal drugs because the dosage levels and labeled indications for use are identical to those currently approved for each drug singly. Accordingly, under the agency's supplemental policy (42 FR 64367, December 23, 1977), this action did not require a reevaluation of the safety and effectiveness data in NADA's 9–476, 7–476, 7–891, and 34–085.

In amending the regulations to reflect this approval, § 558.366 *Nicarbazin* (21 CFR 558.366) is editorially revised to convert paragraph (e) *Conditions of use* from running text to table format.

In accordance with the regulations promulgated under the Freedom of Information Act (Part 20 (21 CFR Part 20)) and § 514.11(e)(2)(ii) of the animal drug regulations (21 CFR 514.11(e)(2)(ii)), summaries of safety and effectiveness data and information submitted to support approval of these applications are available for public examination at the office of the Hearing Clerk (HFA–305), Food and Drug Administration, Rm. 4–65, 5600 Fishers Lane, Rockville, MD 20857, between 9 a.m. and 4 p.m., Monday through Friday.

Therefore, under the Federal Food, Drug, and Cosmetic Act (sec. 512(i), 82 Stat. 347 (21 U.S.C. 360b(i))), under authority delegated to the Commissioner of Food and Drugs (21 CFR 5.1) and

redelegated to the Director of the Bureau of Veterinary Medicine (21 CFR 5.83), Part 558 is amended as follows:

1. In § 558.325, by adding new paragraphs (f)(3) (xi) and (xii) to read as follows:

§ 558.325 Lincomycin.

* * * * *

(f) * * *

(3) * * *

(xi) Nicarbazin and roxarsone as in § 558.366.

(xii) Nicarbazin as in § 558.366.

2. In § 558.366, by revising the text of paragraph (e) to table format, and adding to the table new combination uses for nicarbazin with roxarsone, roxarsone with lincomycin, and lincomycin, to read as follows:

§ 558.366 Nicarbazin.

* * * * *

(e) *Conditons of use.* It is used in chicken feed as follows:

Nicarbazin in grams per ton	Combination in grams per ton	Indications for use	Limitations	Sponsor
113.5 (0.0125 pct)[1]		Chickens; aid in preventing outbreaks of cecal (*Eimeria tenella*) and intestinal (*E. acervulina, E. maxima, E. necatrix,* and *E. brunetti*) coccidiosis [1].	Feed continuously as sole ration from time chicks are placed on litter until past the time when coccidiosis is ordinarily a hazard; do not use as a treatment for coccidiosis; do not use in flushing mashes; do not feed to laying hens; withdraw 4 days before slaughter.	000006
	Lincomycin 2 (0.00044 pct).	Broiler chickens; aid in preventing outbreaks of cecal (*Eimeria tenella*) and intestinal (*E. acervulina, E. maxima, E. necatrix* and *E. brunetti*) coccidiosis; for increased rate of weight gain.	do	000006
113.5 (0.0125 pct)	Roxarsone 22.7 (0.0025).	do	Feed continuously as sole ration from time chicks are placed on litter until past the time when coccidiosis is ordinarily a hazard; as sole source of organic arsenic; do not use a treatment for coccidiosis; do not use in flushing mashes; do not feed to laying hens; withdraw 5 days before slaughter.	000006
	Roxarsone 22.7 (0.0025) plus lincomycin 2 (0.0004).	do	do	000006

* * * * * * *

[1] These conditions are NAS/NRC reviewed and deemed effective. Applications for these uses need not include effectiveness data specified by § 514.111 of this chapter.

3. In § 558.530, by adding new paragraph (e)(4)(xix) to read as follows:

§ 558.530 Roxarsone.

* * * * *

(e) * * *

(4) * * *

(xix) Nicarbazin as in § 558.366.

Effective Date. This regulation is effective July 13, 1979.

(Sec. 512(i), 82 Stat. 347 (21 U.S.C. 360b(i)).)

Dated: July 5, 1979.

Terence Harvey,

Acting Director, Bureau of Veterinary Medicine.

[FR Doc. 79-21455 Filed 7-12-79; 8:45 am]

BILLING CODE 4110-03-M

21 CFR Part 558

New Animal Drugs for use in Animal Feeds; Tylosin

AGENCY: Food and Drug Administration.

ACTION: Final rule.

SUMMARY: The Food and Drug Administration (FDA) amends the regulations to reflect approval of a supplemental new animal drug application (NADA) filed by Zip Feed Mills providing for use of a 10-gram-per-pound tylosin premix for making complete swine feeds.

EFFECTIVE DATE: July 13, 1979.

FOR FURTHER INFORMATION CONTACT: Jack C. Taylor, Bureau of Veterinary Medicine (HFV-136), Food and Drug Administration, Department of Health, Education, and Welfare, 5600 Fishers Lane, Rockville, MD 20857, 301-443-5247.

SUPPLEMENTARY INFORMATION: Zip Feed Mills, 304 E. Eighth St., P.O. Box 500, Sioux Falls, SD 57101, filed a supplemental NADA (97-259) providing for safe and effective use of a premix containing 10 grams of tylosin (as tylosin phosphate) per pound for making complete swine feeds used to increase rate of weight gain and to improve feed efficiency. Approval of this application is based on safety and effectiveness data contained in Elanco Product Co.'s approved NADA 12-491. Use of these data has been authorized by Elanco.

The approval of this supplemental application poses no increased human risk from exposure to residues of the new animal drug because it does not alter usage of the drug. Accordingly, this approval did not require a reevaluation of the safety and effectiveness data in the parent application.

In accordance with the provisions of Part 20 (21 CFR Part 20) promulgated under the Freedom of Information Act (5 U.S.C. 552) and the freedom of information regulations in § 514.11(e)(2)(ii) of the animal drug regulations (21 CFR 514.11(e)(2)(ii)), a summary of safety and effectiveness data and information supporting approval of this application is available for public examination at the office of the Hearing Clerk (HFA-305), Food and Drug Administration, Rm. 4-65, 5600 Fishers Lane, Rockville, MD 20857, from 9 a.m. to 4 p.m., Monday through Friday.

Therefore, under the Federal Food, Drug, and Cosmetic Act (sec. 512(i), 82 Stat. 347 (21 U.S.C. 360b(i))) and under authority delegated to the Commissioner of Food and Drugs (21 CFR 5.1) and redelegated to the Director of the Bureau of Veterinary Medicine (21 CFR 5.83), § 558.625 is amended by revising paragraph (b)(18) to read as follows:

§ 558.625 Tylosin.

* * * * *

(b) * * *

(18) To 017434: 0.4, 4, and 10 grams per pound; paragraph (f)(1)(vi)(*a*) of this section.

* * * * *

Effective date July 13, 1979.

(Sec. 512(i), 82 Stat. 347 (21 U.S.C. 360b(i)).)

Dated: July 5, 1979.

Terence Harvey,

Director, Bureau of Veterinary Medicine.

[FR Doc. 79-21454 Filed 7-12-79; 8:45 am]

BILLING CODE 4110-03-M

21 CFR Part 558

[Docket No. 79N-0198]

New Animal Drugs for use in Animal Feeds; 2-Acetylamino-5-Nitrothiazole; Revocation of Applicable Portion of Regulations

AGENCY: Food and Drug Administration.

ACTION: Final rule.

SUMMARY: The Food and Drug Administration (FDA) is revoking that portion of the regulations reflecting approval of a new animal drug application (NADA) providing for use of a premix containing 2-acetylamino-5-nitrothiazole in turkey feed. The sponsor, American Cyanamid Co., has requested this action.

DATE: Effective July 23, 1979.

FOR FURTHER INFORMATION CONTACT: David N. Scarr, Bureau of Veterinary Medicine (HFV-214), Food and Drug Administration, Department of Health, Education, and Welfare, 5600 Fishers Lane, Rockville, MD 20857, 301-443-3183.

SUPPLEMENTARY INFORMATION: In a notice published elsewhere in this issue of the **Federal Register**, the agency announces that approval of NADA 9-424 is withdrawn. This document amends the animal drug regulations to delete the portion that reflects approval of the NADA. Approval is being withdrawn at the request of the sponsor, American Cyanamid Co.

§ 558.25 2-Acetylamino-5-nitrothiazole [Revoked]

Therefore, under the Federal Food, Drug, and Cosmetic Act (sec. 512(e), 82 Stat. 345-347 (21 U.S.C. 360b(e))) and under authority delegated to the Commissioner of Food and Drugs (21 CFR 5.1) and redelegated to the Director of the Bureau of Veterinary Medicine (21 CFR 5.84), Part 558 is amended by revoking § 558.25 *2-Acetylamino-5-nitrothiazole.*

EFFECTIVE DATE: This regulation shall become effective July 23, 1979.

(Sec. 512(e), 82 Stat. 345-347 (21 U.S.C. 360b(e)).)

Dated: July 6, 1979.

Terence Harvey,

Director, Bureau of Veterinary Medicine.

[FR Doc. 79-21597 Filed 7-12-79; 8:45 am]

BILLING CODE 4110-03-M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

21 CFR Part 1308

Schedules of Controlled Substances; Determination of Schedules for Preparations Containing Narcotic Drugs

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Final rule.

SUMMARY: This rule is issued to specify the method to be used in calculating the amount of a narcotic drug present in a Schedule III, IV or V preparation.

EFFECTIVE DATE: Effective July 26, 1979.

FOR FURTHER INFORMATION CONTACT: Howard McClain, Jr., Chief, Regulatory Control Division, Drug Enforcement Administration, Telephone 202-633-1366.

SUPPLEMENTARY INFORMATION: A notice was published in the **Federal Register** on April 26, 1979 (44 FR 24584), proposing rulemaking to specify the method to be used in calculating the amount of a narcotic drug present in a Schedule III, IV or V preparation and providing an opportunity for any interested party to submit comments or objections in writing regarding this proposal on or before May 29, 1979. No comments were received in response to this proposal.

Therefore, under the authority vested in him by the Act and by regulations of the Department of Justice, the Administrator of the Drug Enforcement Administration hereby orders that 21 CFR be modified as set forth below.

In 21 CFR Part 1308, the introductory paragraphs of § 1308.13(e), § 1308.14(b), and § 1308.15(b) are revised to read as follows:

§ 1308.13 Schedule III.

* * * * *

(e) *Narcotic Drugs.* Unless specifically excepted or unless listed in another schedule, any material, compound, mixture, or preparation containing any of the following narcotic drugs, or their salts calculated as the free anhydrous base or alkaloid, in limited quantities as set forth below:

* * * * *

§ 1308.14 Schedule IV.

* * * * *

(b) *Narcotic drugs.* Unless specifically excepted or unless listed in another schedule, any material, compound, mixture, or preparation containing any of the following narcotic drugs, or their salts calculated as the free anhydrous base or alkaloid, in limited quantities as set forth below:

* * * * *

§ 1308.15 Schedule V.

* * * * *

(b) Narcotic drugs containing non-narcotic active medicinal ingredients. Any compound, mixture, or preparation containing any of the following narcotic drugs, or their salts calculated as the free anhydrous base or alkaloid, in limited quantities as set forth below, which shall include one or more non-narcotic active medicinal ingredients in sufficient proportion to confer upon the compound, mixture, or preparation valuable medicinal qualities other than those possessed by narcotic drugs alone:

* * * * *

Peter B. Bensinger,

Administrator.

July 10, 1979.

[FR Doc. 79-21765 Filed 7-12-79; 8:45 am]

BILLING CODE 4110-09-M

DEPARTMENT OF HOUSING AND URBAN DEVELOPMENT

Office of Assistant Secretary for Housing—Federal Housing Commissioner

24 CFR Parts 203, 207, 220

[Docket No. R-79-683]

Debenture Interest Rates

AGENCY: Department of Housing and Urban Development.

ACTION: Final rule.

SUMMARY: This rule change provides for an increased debenture interest rate applicable to all home and project mortgages and loans under the National Housing Act (the Act), as amended, except for those loans or mortgages insured under the Act's section 221(g)(4) provision, committed or endorsed on or after July 1, 1979. The Secretary of the Treasury determines debenture interest rates in accordance with established procedure and the Act. The intended effect of this rule change is to increase debenture interest rates for appropriate mortgages.

EFFECTIVE DATE: This rule is effective August 13, 1979, retroactive to July 1, 1979.

FOR FURTHER INFORMATION CONTACT: T. J. O'Connor, Director, Office of Finance and Accounting, Department of Housing and Urban Development, 451 Seventh Street, SW., Room 2202, Washington, D.C. 20410, (202) 755-6310. (This is not a toll free number.)

SUPPLEMENTARY INFORMATION: The Secretary of the Treasury has determined in accordance with the provisions of section 224 of the National Housing Act, as amended, that the interest rate for the month of May 1979 is 8¼% and has approved the establishment of debenture interest rates at 8¼% to be effective as of July 1, 1979.

The Secretary of Housing and Urban Development has determined that advance publication and notice and public procedure are unnecessary since the debenture interest rate is set by the Secretary of the Treasury in accordance with a procedure established by statute.

A Finding of Inapplicability respecting the National Environmental Policy Act of 1969 has been made in accordance with HUD procedures. A copy of this Finding of Inapplicability will be available for public inspection during regular business hours at the office of the Rules Docket Clerk, Office of the General Counsel, Room 5218, Department of Housing and Urban Development, 451 7th Street, SW, Washington, D.C. 20410.

Accordingly, Chapter II is amended as follows:

PART 203—MUTUAL MORTGAGE INSURANCE AND INSURED HOME IMPROVEMENT LOANS

Subpart B—Contract Rights and Obligations

1. Section 203.405 is amended to read as follows:

§ 203.405 Debenture interest rate.

Debentures shall bear interest from the date of issue, payable semiannually on the first day of January and the first day of July of each year at the rate in effect as of the day the commitment was issued, or as of the date the mortgage was endorsed for insurance, whichever rate is higher.

The following interest rates are effective for the dates listed:

	On or after	Prior to
Effective rate (percent):		
6⅛	July 1, 1974	July 1, 1975
7	July 1, 1975	Jan. 1, 1976
7⅛	Jan. 1, 1976	July 1, 1976
7	July 1, 1976	Jan. 1, 1977
6⅝	Jan. 1, 1977	July 1, 1977
7¼	July 1, 1977	Jan. 1, 1978
7⅛	Jan. 1, 1978	July 1, 1978
7¾	July 1, 1978	Jan. 1, 1979
8	Jan. 1, 1979	July 1, 1979
8¼	July 1, 1979	

(Sec. 211, 52 Stat. 23; (12 U.S.C. 1715b). Interprets or applies sec. 203, 52 Stat. 10, as amended; (12 U.S.C. 1709).)

2. Section 203.479 is amended to read as follows:

§ 203.479 Debenture interest rate.

Debentures shall bear interest from the date of issue, payable semiannually on the first day of January and the first day of July of each year at the rate in effect as of the date the commitment was issued, or as of the date the loan was endorsed for insurance, whichever rate is the higher. The following interest rates are effective for the dates listed:

	On or after	Prior to
Effective rate (percent):		
6⅛	July 1, 1974	July 1, 1975
7	July 1, 1975	Jan. 1, 1976
7⅛	Jan. 1, 1976	July 1, 1976
7	July 1, 1976	Jan. 1, 1977
6⅝	Jan. 1, 1977	July 1, 1977
7¼	July 1, 1977	Jan. 1, 1978
7⅛	Jan. 1, 1978	July 1, 1978
7¾	July 1, 1978	Jan. 1, 1979
8	Jan. 1, 1979	July 1, 1979
8¼	July 1, 1979	

(Sec. 211, 52 Stat. 23; (12 U.S.C. 1715b). Interprets or applies sec. 203, 52 Stat. 10, as amended; (12 U.S.C. 1709).)

PART 207—MULTIFAMILY HOUSING MORTGAGE INSURANCE

Subpart B—Contract Rights and Obligations

3. In 207.259 paragraph (e)(6) is amended to read as follows:

§ 207.259 Insurance benefits.

* * * * *

(e) *Issuance of debentures.* * * *

(6) Bear interest from the date of issue, payable semiannually on the first day of January and the first day of July of each year at the rate in effect as of the date the commitment was issued, or as of the date of initial insurance endorsement of the mortgage, whichever rate is the higher. The following interest rates are effective for the dates listed:

	On or after	Prior to
Effective rate (percent):		
6⅛	July 1, 1974	July 1, 1975
7	July 1, 1975	Jan. 1, 1976
7⅛	Jan. 1, 1976	July 1, 1976
7	July 1, 1976	Jan. 1, 1977
6⅝	Jan. 1, 1977	July 1, 1977
7¼	July 1, 1977	Jan. 1, 1978
7⅛	Jan. 1, 1978	July 1, 1978
7¾	July 1, 1978	Jan. 1, 1979
8	Jan. 1, 1979	July 1, 1979
8¼	July 1, 1979	

(Sec. 211, 52 Stat. 23; (12 U.S.C. 1715b). Interprets or applies sec. 207, 52 Stat. 16, as amended; (12 U.S.C. 1713).)

PART 220—URBAN RENEWAL MORTGAGE INSURANCE AND INSURED IMPROVEMENT LOANS

Subpart D—Contract Rights and Obligations—Projects

4. Section 220.830 is amended to read as follows:

§ 220.830 Debenture interest rate.

Debentures shall bear interest from the date of issue, payable semianually on the first day of January and the first day of July of each year at the rate in effect as of the date the commitment was issued or as of the date the loan was endorsed for insurance, whichever rate is higher. The following interest rates are effective for the dates listed:

	On or after	Prior to
Effective rate (percent):		
6⅛	July 1, 1974	July 1, 1975
7	July 1, 1975	Jan. 1, 1976
7⅛	Jan. 1, 1976	July 1, 1976
7	July 1, 1976	Jan. 1, 1977
6⅝	Jan. 1, 1977	July 1, 1977
7¼	July 1, 1977	Jan. 1, 1978
7⅛	Jan. 1, 1978	July 1, 1978
7¾	July 1, 1978	Jan. 1, 1979
8	Jan. 1, 1979	July 1, 1979
8¼	July 1, 1979	

(Sec. 211, 52 Stat. 23; (12 U.S.C. 1715b). Interprets or applies sec. 220, 68 Stat. 596, as amended; (12 U.S.C. 1715k).)

Issued at Washington, D.C., July 10, 1979.

Lawrence B. Simons

Assistant Secretary for Housing, Federal Housing Commissioner.

[FR Doc. 79-21870 Filed 7-12-79; 8:45 am]

BILLING CODE 4210-01-M

24 CFR Part 221

[Docket No. R-79-558]

Mortgage and Loan Insurance Programs Under the National Housing Act; Multifamily Housing Projects Financed With Tax-Exempt Obligations

AGENCY: Department of Housing and Urban Development.

ACTION: Final rule.

SUMMARY: This rule clarifies that an amendment to 24 CFR 221.762, published on February 8, 1979, which provided that multifamily housing projects financed with tax-exempt obligations issued