Board finds good cause for making this action effective immediately.

Lists of Subjects in 12 CFR Part 204.

Banks, banking, Currency, Federal Reserve System, Penalties, Reporting requirements.

PART 204—[AMENDED]

Pursuant to its authority under section 19(b) of the Federal Reserve Act (12 U.S.C. 461(b)), effective October 5, 1982, the Board amends Regulation D (12 CFR Part 204) by revising paragraph (e) of § 204.2 to read as follows:

§ 204.2 Definitions.

* * * * *

(e) * * *

(7) deposits or accounts maintained in connnection with an arrangement that permits the depositor to obtain credit directly or indirectly through the drawing of a negotiable or nonnegotiable check, draft, order or instruction or other similar device (including telephone or electronic order or instruction) on the issuing institution that can be used for the purpose of making payments or transfers to third persons or others, or to a deposit account of the depositor. Deposits that are subject to arrangements established before October 5, 1982, will not be regarded as transaction accounts (i) until the deposit issued in connection with the line of credit is extended, or matures and is renewed, or (ii) if the deposit issued in connection with the line of credit matures and is automatically renewed on or before December 31, 1982.

* * * * *

By order of the Board of Governors, November 16, 1982.

William W. Wiles,

Secretary of the Board.

[FR Doc. 82-31882 Filed 11-22-82; 8:45 am]

BILLING CODE 6210-01-M

CIVIL AERONAUTICS BOARD

14 CFR Part 382

[Reg. SPR-192; Special Regs. Amdt. No. 2 to Part 382]

Nondiscrimination on the Basis of Handicap; Notice of Approval by the Office of Management and Budget

AGENCY: Civil Aeronautics Board.

ACTION: Final rule.

SUMMARY: This final rule gives notice that the Office of Management and Budget (OMB) has approved the reporting requirements contained in new Part 382 of the Board's Special Regulations, "Nondiscrimination on the Basis of Handicap." This approval has been granted through June 30, 1984. OMB approval is required under the Paperwork Reduction Act of 1980.

DATES: Adopted: November 17, 1982. Effective: November 1, 1982.

FOR FURTHER INFORMATION CONTACT: Linda K. Koman, Data Requirements Section, Information Management Division, Office of Comptroller, Civil Aeronautics Board, 1825 Connecticut Avenue, N.W., Washington, D.C. 20428, (202) 673-6042.

Accordingly, the Civil Aeronautics Board amends Part 382 of its Special Regulations (14 CFR Part 382) by adding a note at the end of the table of contents to Part 382 to read:

Note.—The reporting requirements contained in §§ 382.21, 382.22 and 382.23 have been approved by the Office of Management and Budget under number 3024-0056.

This amendment is issued by the undersigned pursuant to delegation of authority from the Board to the Secretary in 14 CFR 385.24(b).

(Sec. 204 of the Federal Aviation Act of 1958, as amended, 72 Stat. 743; 49 U.S.C. 1324).

By the Civil Aeronautics Board.

Phyllis T. Kaylor,

Secretary.

[FR Doc. 82-32105 Filed 11-22-82; 8:45 am]

BILLING CODE 6320-01-M

SECURITIES AND EXCHANGE COMMISSION

17 CFR Part 270

[Release No. IC-12678; 57-907]

Adoption of Permanent Exemptions From Certain Provisions of the Investment Company Act of 1940 for Registered Separate Accounts and Other Persons

Correction

In FR Doc. 82-26630, beginning on page 42553, Tuesday, September 28, 1982 make the following corrections:

1. On page 42554, in the second column, in the first paragraph, in the twenty-third line, "a annunity" should read "an annuity".

2. On page 42555, in the third column, in the second line from the top, "or course," should read "of course,".

3. On page 42556, in the second column, in the ninth line from the bottom of the text, "company," should read "company;".

4. On page 42557, in the first column, the second line from the bottom of the text, "paragraph (1)" should read "paragraph (l)".

5. Also on page 42557, in the second column, in footnote 22, in the sixth line "[35" should read "[36".

6. On page 42559, in the second column, in the first paragraph, in the first line, "paragraph (1)" should read "paragraph (l)".

BILLING CODE 1505-01-M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Social Security Administration

20 CFR Parts 404 and 416

Federal Old Age, Survivors and Disability Insurance Benefits

AGENCY: Social Security Administration, HHS.

ACTION: Correction of a final regulation.

SUMMARY: This document corrects a citation to the Code of Federal Regulations that appears several times throughout final regulations implementing section 301 of Pub. L. 96-265 which were published July 21, 1982 (47 FR 31539). Section 301 provides that, in certain cases, a disabled person who has medically recovered will not have his or her disability benefits terminated if he or she is participating in an approved vocational rehabilitation program. In the preparation of these regulations several references were made to the Education Department regulations. With the creation of the Education Department, the regulations appearing in 45 CFR Part 1361 were transferred and redesignated to Title 34 of the Code of Federal Regulations, therefore making the citation to 45 CFR Part 1361 incorrect (45 FR 77368, November 21, 1980).

FOR FURTHER INFORMATION CONTACT: Mr. Harry J. Short, Office of Regulations, Social Security Administration, 6401 Security Boulevard, Baltimore, MD 21235, Telephone (301) 594-7337.

1. On page 31539, under Supplementary Information, the reference to 45 CFR 1361.2 and 1361.39 is corrected to 34 CFR Part 361.

2. On page 31542, in the middle column, in § 404.316(c)(1)(ii), the reference to 45 CFR 1361.39 is corrected to read 34 CFR Part 361.

3. On page 31543, in the third column, in § 404.1586(f)(1)(ii) the reference to 45 CFR 1361.39 is corrected to read 34 CFR Part 361.

4. On page 31544, in the first, middle, and third columns, in §§ 404.1596(c)(4), 416.1321(d), and 416.1338(a)(2) the references to 45 CFR 1361.39 are corrected to read 34 CFR Part 361.

Dated: November 14, 1982.

Robert F. Sermier,

Deputy Assistant Secretary for Management Analysis and Systems.

[FR Doc. 82-32010 Filed 11-22-82; 8:45 am]

BILLING CODE 4190-11-M

Food and Drug Administration

21 CFR Parts 74, 81 and 82

[Docket No. 82N-0268]

D&C Orange No. 5; Correction

AGENCY: Food and Drug Administration.

ACTION: Final rule; correction.

SUMMARY: The Food and Drug Administration (FDA) is correcting the effective date and the date for submittal of objections for a document that permanently listed D&C Orange No. 5 for use in lipsticks or other lip cosmetics and in drug and cosmetic mouthwashes and dentifrices.

DATES: The effective date for the rule published at 47 FR 49632 is December 3, 1982; objections by December 2, 1982.

FOR FURTHER INFORMATION CONTACT: Andrew D. Laumbach, Bureau of Foods (HFF-334), Food and Drug Administration, 200 C St. SW., Washington, DC 20204, 202-472-5690.

SUPPLEMENTARY INFORMATION: In FR Doc. 82-30082 appearing at page 49632 in the issue for Tuesday, November 2, 1982, the following corrections are made:

1. On page 49632, in the third column under "DATES", the words "Effective November 30, 1982; objections by November 29, 1982;" are corrected to read "Effective December 3, 1982; objections by December 2, 1982;".

2. On page 49636, in the second column, first line, the words "November 29" are corrected to read "December 2" and in the second line of the first full paragraph, the words "November 30" are corrected to read "December 3".

Dated: November 18, 1982.

William F. Randolph,

Acting Associate Commissioner for Regulatory Affairs.

[FR Doc. 82-32129 Filed 11-19-82; 12:55 pm]

BILLING CODE 4160-01-M

21 CFR Part 146

[Docket No. 76P-0163]

Identity and Quality Standard for Canned Pineapple Juice; Amendment and Ending of Stay After Commissioner's Final Decision

AGENCY: Food and Drug Administration.

ACTION: Final rule.

SUMMARY: The Food and Drug Administration (FDA) is ending the stay of effectiveness of the provision establishing the 13.5° Brix pineapple juice soluble solids requirement in § 146.185(b)(1)(i) (21 CFR 146.185(b)(1)(i)) and is setting a new Brix level, 12.8°. FDA is taking this action consistent with the Commissioner's Final Decision in the rulemaking proceeding that resulted in the stay. The Commissioner's Final Decision appears in the "Notices" section of this **Federal Register.**

EFFECTIVE DATE: May 23, 1983.

FOR FURTHER INFORMATION CONTACT: Ted Herman, Regulations Policy Staff (HFC-10), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301-443-3480.

SUPPLEMENTARY INFORMATION: FDA issued in the **Federal Register** of May 28, 1976 (41 FR 21768) an amendment to § 146.185(b)(1)(i) setting the minimum Brix level for pineapple juice from concentrate at 13.5°. FDA received an objection and a request for hearing on this provision. In the **Federal Register** of March 14, 1978 (43 FR 10552), FDA stayed the effectiveness of this Brix requirement, pending further notice. The Commissioner of Food and Drugs has issued his Final Decision on this matter—see the Final Decision in the "Notices" section of this **Federal Register.**

List of Subjects in 21 CFR Part 146

Canned fruit juice, Food standards, Fruit juices.

PART 145—CANNED FRUIT JUICES

§ 146.185 [Amended]

Therefore, under the Federal Food Drug, and Cosmetic Act (secs. 401, 701(e), 52 Stat. 1046 as amended, 70 Stat. 919 as amended (21 U.S.C. 341, 371(e))), under authority delegated to the Commissioner (21 CFR 5.10), and in accordance with the Commissioner's Final Decision, the stay of the provision establishing the 13.5° Brix requirement in § 146.185(b)(1)(i) is ended and § 146.185 *Canned pineapple juice* is amended in the second sentence of paragraph (b)(1)(i) by changing "13.5" to "12.8".

Effective date. Effective May 23, 1983, for affected products initially introduced or initially delivered for introduction into interstate commerce.

Dated: November 15, 1982.

Arthur Hull Hayes, Jr.,

Commissioner of Food and Drugs.

[FR Doc. 82-32044 Filed 11-22-82; 8:45 am]

BILLING CODE 4160-01-M

21 CFR Parts 510 and 520

Oral Dosage Form New Animal Drugs Not Subject to Certification; Dichlorophene and Toluene Capsules

AGENCY: Food and Drug Administration.

ACTION: Final rule.

SUMMARY: The Food and Drug Administration is amending the animal drug regulations to reflect approval of a new animal drug application (NADA) filed by Natchez Animal Supply Co. providing for safe and effective use of dichlorophene and toluene capsules for treating dogs and cats for certain helminth infections, and to add Natchez Animal Supply Co. to the list of sponsors of approved NADA's.

EFFECTIVE DATE: November 23, 1983.

FOR FURTHER INFORMATION CONTACT: Bob G. Griffith, Bureau of Veterinary Medicine (HFV-112), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301-443-3430.

SUPPLEMENTARY INFORMATION: Natchez Animal Supply Co., 201 John R. Junkin Dr., Natchez, MS 39120, filed an NADA (121-557) providing for use of dichlorophene and toluene capsules for treating dogs and cats for infections of certain ascarids, hookworms, and tapeworms. This product is the generic equivalent of a product that was the subject of a National Academy of Sciences/National Research Council (NAS/NRC) notice published in the **Federal Register** of February 1, 1969 (34 FR 1613) and reflected in 21 CFR 520.580. Approval of Natchez's product does not require submission of data to demonstrate bioequivalency because it is to be manufactured by the firms currently manufacturing the NAS/NRC-reviewed product. In addition, Natchez Animal Supply Co. was not included in the list of sponsors of approved applications found in 21 CFR 510.600(c). The NADA is approved and the regulations are amended to include this sponsor and the approval.

In accordance with the freedom of information 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 may be seen in the Dockets Management Branch (HFA-305), Food and Drug Administration, Rm. 4-62, 5600 Fishers Lane, Rockville, MD 20857, from 9 a.m. to 4 p.m., Monday through Friday.

The Bureau of Veterinary Medicine has determined pursuant to 21 CFR 25.24(d)(1)(i) (proposed December 11,

1979; 44 FR 71742) that this action is of a type that does not individually or cumulatively have a significant impact on the human environment. Therefore, neither an environmental assessment nor an environmental impact statement is required.

This action is governed by the provisions of 5 U.S.C. 556 and 557 and is therefore excluded for Executive Order 12291 by section 1(a)(1) of the Order.

List of Subjects

21 CFR Part 510

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

21 CFR Part 520

Animal drugs, Oral use.

PART 510—[AMENDED]

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 5.10) and redelegated to the Bureau of Veterinary Medicine (21 CFR 5.83), Parts 510 and 520 are amended as follows:

1. In Part 510, § 510.600 is amended by adding a new sponsor alphabetically to paragraph (c)(1) and numerically to paragraph (c)(2), to read as follows:

§ 510.600 Names, addresses, and drug labeler codes of sponsors of approved applications.

* * * * *

(c) ***

(1) ***

Firm name and address	Drug labeler code
* * * * *	
Natchez Animal Supply Co., 201 John R. Junkin Dr., Natchez, MS 39120	049968
* * * * *	

(2) * * *

Drug labeler code	Firm name and address
* * * * *	
049968	Natchez Animal Supply Co., 201 John R. Junkin Dr., Natchez, MS 39120.
* * * * *	

PART 520—[AMENDED]

2. In Part 520, § 520.580 is amended by revising paragraph (b)(1), to read as follows:

§ 520.580 Dichlorophene and toluene capsules.

* * * * *

(b) *Sponsor.* (1) For single dose only, see 000010, 000298, 000842, 000856, 010290, 010888, 011519, 011536, 011614, 015563, 017135, 023851, 049968, and 050906 in § 510.600(c) of this chapter.

* * * * *

Effective date. November 23, 1982.

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

Dated: November 17, 1982.

Lester M. Crawford,

Director, Bureau of Veterinary Medicine.

[FR Doc. 82-32034 Filed 11-22-82; 8:45 am]

BILLING CODE 4160-01-M

---

21 CFR Parts 510 and 558

New Animal Drugs for Use in Animal Feeds; Tylosin; Sponsors of Approved NADA's

AGENCY: Food and Drug Administration.

ACTION: Final rule.

SUMMARY: The Food and Drug Administration (FDA) is removing those portions of the regulations reflecting approval of a new animal drug application (NADA) providing for use of an 0.8-gram-per-pound tylosin (as tylosin phosphate) premix in making complete swine feeds used for increased rate of weight gain and improved feed efficiency. The sponsor, International Multifoods' Dixie Mills Co., requested the withdrawal of approval. Dixie Mills Co. is being removed from the list of sponsors of approved NADA's.

EFFECTIVE DATE: December 3, 1982.

FOR FURTHER INFORMATION CONTACT: Howard Meyers, Bureau of Veterinary Medicine (HFV-218), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301-443-4093.

SUPPLEMENTARY INFORMATION: In a notice published elsewhere in this issue of the **Federal Register**, approval of NADA 100-744 for Dixie Booster Pak-Tylan 1600 Medicated sponsored by International Multifoods' Dixie Mills Co. is withdrawn. This document amends the regulations to remove those portions of 21 CFR 510.600 and 558.625 which reflect approval of the NADA.

The Bureau of Veterinary Medicine has determined pursuant to 21 CFR 25.24(d)(2) (proposed December 11, 1979; 44 FR 71742) that this action is of a type that does not individually or cumulatively have a significant impact on the human environment. Therefore, neither an environmental assessment nor an environmental impact statement is required.

This action is governed by the provisions of 5 U.S.C. 556 and 557 and is therefore excluded from Executive Order 12291 by section 1(a)(1) of the Order.

List of Subjects

21 CFR Part 510

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

21 CFR Part 558

Animal drugs, Animal feeds.

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 5.10) and redelegated to the Bureau of Veterinary Medicine (21 CFR 5.84), Parts 510 and 558 are amended as follows:

PART 510—NEW ANIMAL DRUGS

§ 510.600 [Amended]

1. Part 510 is amended in § 510.600 *Names, addresses, and drug labeler codes of sponsors of approved applications* by by removing the entry for "Dixie Mills Co." from paragraph (c)(1) and removing the entry for "025066" from paragraph (c)(2).

PART 558—NEW ANIMAL DRUGS FOR USE IN ANIMAL FEEDS

§ 558.625 [Amended]

2. Part 558 is amended in § 558.625 *Tylosin* by removing the text of paragraph (b)(41) and marking it "[Reserved]."

Effective date. December 3, 1982.

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

Dated: November 17, 1982.

Lester M. Crawford,

Director, Bureau of Veterinary Medicine.

[FR Doc. 82-32043 Filed 11-22-82; 8:45 am]

BILLING CODE 4160-01-M

---

21 CFR Part 520

Oral Dosage Form New Animal Drugs not Subject to Certification; Dithiazanine Iodide and Piperazine Citrate Suspension

AGENCY: Food and Drug Administration.

ACTION: Final rule.

SUMMARY: The Food and Drug Administration (FDA) is amending the animal drug regulations to codify a previously approved new animal drug