CTAF/UNICOM frequency and therefore, a control zone is not needed. The airspace study for this action indicated the establishment of a control zone would not require aircraft to have two-way radio capability because this airport does not have an operating control tower.

The airport is served by an ILS to runway 18, with a minimum Decision Height of 300 feet AGL and a minimum visibility of 1 mile. The last 400 feet authorized on this approach lie in uncontrolled airspace, in which ATC cannot guarantee separation from all other aircraft. Additionally, there are two VOR approaches. However their Minimum Descent Altitudes are above the floor of the Transition area.

At present, the airport CTAF/UNICOM frequency is 122.8 Mhz., on which it is not mandatory for either aircraft to communicate or monitor, although highly prudent.

The Wyoming State Aeronautics Division petitioned the FAA to establish a control zone for this airport, and the principle air carrier user of the airport endorsed the proposal.

Based on all the information available, the FAA has decided to adopt the rule as proposed, with a clarification that the miles referenced in the rule are nautical miles.

Control zones are published in § 71.171 of Handbook 7400.7 effective November 1, 1991, which is incorporated by reference in 14 CFR 71.1. The control zone listed in this document will be published subsequently in the Handbook.

The Rule

This amendment to part 71 of the Federal Aviation Regulations establishes a control zone for the Jackson Hole Airport, Wyoming, to provide additional controlled airspace for aircraft executing instrument approach procedures to the Jackson Hole Airport. This action will segregate aircraft operating in VFR from those aircraft operating in IFR. The airspace will be depicted on aeronautical charts for pilot reference. The FAA has determined that this regulation only involves an established body of technical regulations for which frequent and routine amendments are necessary to keep them operationally current. It, therefore—(1) is not a "major rule" under Executive Order 12291; (2) is not a "significant rule" under DOT Regulatory Policies and Procedures (44 FR 11034; February 26, 1979); and (3) does not warrant preparation of a regulatory evaluation as the anticipated impact is so minimal. Since this is a routine matter that will only affect air traffic procedures and air navigation, it is certified that this rule will not have a significant economic impact on a substantial number of small entities under the criteria of the Regulatory Flexibility Act.

List of Subjects in 14 CFR Part 71

Aviation safety, Control zones, Incorporation by reference

Adoption of the Amendment

Accordingly, pursuant to the authority delegated to me, part 71 of the Federal Aviation Regulations (14 CFR part 71) is amended, as follows:

PART 71—[AMENDED]

1. The authority citation for 14 CFR part 71 continues to read as follows:

Authority: 49 U.S.C. app. 1348(a), 1354(a), 1510; E.O. 10854, 24 FR 9565, 3 CFR, 1959–1963 Comp., p. 389; 49 U.S.C. 106(g); 14 CFR 11.69.

§ 71.1 [Amended]

2. The incorporation by reference in 14 CFR 71.1 of the Federal Aviation Administration Order 7400.7, Compilation of Regulations, published April 30, 1991, and effective November 1, 1991, is amended as follows:

Section 71.171 Designation

* * * * *

ANM WY CZ Jackson, WY [New]
Jackson Hole Airport, WY
(lat. 43°36'24" N, long. 110°44'15" W.)

Within a 4.3 nautical mile radius of Jackson Hole Airport, Wyoming.

* * * * *

Issued in Seattle, Washington on September 4, 1992.

Temple H. Johnson, Jr.,

Manager, Air Traffic Division.

[FR Doc. 92–22196 Filed 9–14–92; 8:45 am]

BILLING CODE 4910-13-M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 529

Certain Other Dosage Form New Animal Drugs; Formalin 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 Western Chemical, Inc. The NADA provides for the use of formalin solution (aqueous solution of 37 percent formaldehyde) to control certain external protozoa and monogenetic trematodes on salmon, trout, catfish, largemouth bass, and bluegill and to control certain fungi on salmon, trout, and esocid eggs.

EFFECTIVE DATE: September 15, 1992.

FOR FURTHER INFORMATION CONTACT: George K. Haibel, Center for Veterinary Medicine (HFV–133), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–295–8644.

SUPPLEMENTARY INFORMATION: Western Chemical, Inc., 1269 Lattimore Rd., Ferndale, WA 98248, has filed NADA 140–989. The NADA provides for the use of formalin solution (aqueous solution of 37 percent formaldehyde) in tanks, raceways, and ponds to control: (1) the external protozoa: *Ichthyophthirius* spp., *Chilodonella* spp., *Costia* spp., *Epistylis* spp., *Scyphidia* spp., and *Trichodina* spp.; (2) the monogenetic trematodes: *Cleidodiscus* spp., *Gyrodactylus* spp., and *Dactylogyrus* spp. on salmon, trout, catfish, largemouth bass, and bluegill; and (3) in incubation tanks to control fungi of the family Saprolegniaceae on salmon, trout, and esocid eggs. The NADA is approved as of July 31, 1992, and the regulations are amended in 21 CFR 529.1030(b) to reflect this approval. The basis for approval is discussed in the freedom of information summary.

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. 1–23, 12420 Parklawn Dr., Rockville, MD 20857, between 9 a.m. and 4 p.m., Monday through Friday.

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 does not qualify for marketing exclusivity because no new clinical or field investigations (other than bioequivalence or residue studies) and no new human food safety studies (other than bioequivalence or residue studies) were essential to the approval and conducted or sponsored by the applicant.

The agency has carefully considered the potential environmental effects of this action. FDA has concluded that the action will not have a significant impact on the human environment, and that an environmental impact statement is not required. The agency's finding of no significant impact and the evidence supporting that finding, contained in an

environmental assessment, may be seen in the Dockets Management Branch (address above) between 9 a.m. and 4 p.m., Monday through Friday.

List of Subjects in 21 CFR Part 529

Animal drugs.

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 529 is amended as follows:

PART 529—CERTAIN OTHER DOSAGE FORM NEW ANIMAL DRUGS

1. The authority citation for 21 CFR part 529 continues to read as follows:

Authority: Sec. 512 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 360b).

§ 529.1030 [Amended]

2. Section 529.1030 *Formalin solution* is amended in paragraph (b) by revising the phrase "See Nos. 049968 and 051212" to read "See Nos. 049968, 050378, and 051212".

Dated: September 9, 1992.

Gerald B. Guest,

Director, Center for Veterinary Medicine.

[FR Doc. 92–22237 Filed 9–14–92; 8:45 a.m.]

BILLING CODE 4160–01–F

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21 CFR Part 520

Animal Drugs, Feeds, and Related Products; Oxytetracycline Hydrochloride Soluble Powder

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule.

SUMMARY: The Food and Drug Administration (FDA) is amending the animal drug regulations to codify an approved new animal drug application (NADA) held by Fermenta Animal Health Co. The NADA provides for the use of an oxytetracycline hydrochloride soluble powder as an antibacterial in the drinking water of growing turkeys.

EFFECTIVE DATE: September 15, 1992.

FOR FURTHER INFORMATION CONTACT: Dianne T. McRae, Center for Veterinary Medicine (HFV–102), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–295–8623.

SUPPLEMENTARY INFORMATION: Fermenta Animal Health Co., 10150 North Executive Hills Blvd., Kansas City, MO 64153, is the sponsor of NADA 38–200, which provides for the use of oxytetracycline hydrochloride soluble powder as an antibacterial in the drinking water of growing turkeys for control of hexamitiasis caused by *Hexamita meleagridis*. The NADA was originally approved as published in the **Federal Register** of October 13, 1970 (35 FR 16041). That approval should have been but was not codified in part 520 (21 CFR part 520). Accordingly, the regulations at § 520.1660d(b) (21 CFR 520.1660d(b)) are now amended to provide for that approval.

List of Subjects in 21 CFR Part 520

Animal drugs.

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 520 is amended as follows:

PART 520—ORAL DOSAGE FORM NEW ANIMAL DRUGS NOT SUBJECT TO CERTIFICATION

1. The authority citation for 21 CFR part 520 continues to read as follows:

Authority: Sec. 512 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 360b).

2. Section 520.1660d is amended by revising paragraph (b) to read as follows:

§ 520.1660d Oxytetracycline hydrochloride soluble powder.

* * * * *

(b) *Sponsors.* See No. 017144 in § 510.600(c) of this chapter for use as in paragraph (e)(1) of this section. See No. 054273 for use as in paragraph (e)(1)(ii) of this section.

* * * * *

Dated: September 9, 1992.

Gerald B. Guest,

Director, Center for Veterinary Medicine.

[FR Doc. 92–22239 Filed 9–14–92; 8:45 a.m.]

BILLING CODE 4160–01–F

DEPARTMENT OF THE TREASURY

Internal Revenue Service

26 CFR Part 1

[T.D. 8423]

RIN 1545–AP93

Taxation of Tax-Exempt Organizations' Income From Ordinary and Routine Investments; Correction

AGENCY: Internal Revenue Service, Treasury.

ACTION: Correction to final regulations.

SUMMARY: This document contains a correction to Treasury Decision 8423, which was published in the **Federal Register** Wednesday, July 29, 1992 (57 FR 33442), relating to the taxation of tax-exempt organizations' unrelated business taxable income.

EFFECTIVE DATE: July 29, 1992.

FOR FURTHER INFORMATION CONTACT: Regina L. Oldak, (202) 622–6080 (not a toll-free number).

SUPPLEMENTARY INFORMATION:

Background

The final regulations that are the subject of this correction amend the regulations under section 512(b)(1) to conform to and incorporate the 1978 statutory amendment regarding securities loans.

Need for Correction

As published, T.D. 8423 contains an error which may prove to be misleading and is in need of clarification.

Correction of Publication

Accordingly, the publication of the final regulation (T.D. 8423), which was the subject of FR Doc. 92–17677, is corrected as follows:

1. On page 33443, column 3, § 1.512(b)–1(a)(3), line 4, the language "prior to this Treasury decision remain" is corrected to read "prior to August 30, 1991, remain".

Dale D. Goode,

Federal Register Liaison Officer, Assistant Chief Counsel (Corporate).

[FR Doc. 92–22230 Filed 9–14–92; 8:45 am]

BILLING CODE 4830–01–M

PENSION BENEFIT GUARANTY CORPORATION

29 CFR Part 2676

Valuation of Plan Benefits and Plan Assets Following Mass Withdrawal—Interest Rates

AGENCY: Pension Benefit Guaranty Corporation.

ACTION: Final rule.

SUMMARY: This is an amendment to the Pension Benefit Guaranty Corporation's regulation on Valuation of Plan Benefits and Plan Assets Following Mass Withdrawal (29 CFR part 2676). The regulation prescribes rules for valuing benefits and certain assets of multiemployer plans under sections 4219(c)(1)(D) and 4281(b) of the Employee Retirement Income Security Act of 1974. Section 2676.15(c) of the regulation contains a table setting forth, for each calendar month, a series of interest rates to be used in any valuation performed as of a valuation date within that calendar month. On or about the fifteenth of each month, the

PBGC publishes a new entry in the table for the following month, whether or not the rates are changing. This amendment adds to the table the rate series for the month of October 1992.

EFFECTIVE DATE: October 1, 1992.

FOR FURTHER INFORMATION CONTACT: Deborah C. Murphy, Attorney, Office of the General Counsel (22500), Pension Benefit Guaranty Corporation, 2020 K Street NW., Washington, DC 20006; 202–778–8820 (202–778–1958) for TTY and TDD). (These are not toll-free numbers.)

SUPPLEMENTARY INFORMATION: The PBGC finds that notice of and public comment on this amendment would be impracticable and contrary to the public interest, and that there is good cause for making this amendment effective immediately. These findings are based on the need to have the interest rates in this amendment reflect market conditions that are as nearly current as possible and the need to issue the interest rates promptly so that they are available to the public before the beginning of the period to which they apply. (See 5 U.S.C. 553 (b) and (d).) Because no general notice of proposed rulemaking is required for this amendment, the Regulatory Flexibility Act of 1980 does not apply (5 U.S.C. 601(2)).

The PBGC has also determined that this amendment is not a "major rule" within the meaning of Executive Order 12291 because it will not have an annual effect on the economy of $100 million or more; or create a major increase in costs or prices for consumers, individual industries, or geographic regions; or have significant adverse effects on competition, employment, investment, or innovation, or on the ability of United States-based enterprises to compete with foreign-based enterprises in domestic or export markets.

List of Subjects in 29 CFR Part 2676

Employee benefit plans and pensions.

In consideration of the foregoing, part 2676 of subchapter H of chapter XXVI of title 29, Code of Federal Regulations, is amended as follows:

PART 2676—VALUATION OF PLAN BENEFITS AND PLAN ASSETS FOLLOWING MASS WITHDRAWAL

1. The authority citation for part 2676 continues to read as follows:

Authority: 29 U.S.C. 1302(b)(3), 1399(c)(1)(D), and 1441(b)(1).

2. In § 2676.15, paragraph (c) is amended by adding to the end of the table of interest rates therein the following new entry:

§ 2676.15 Interest.

* * * * *

(c) *Interest Rates.*

For valuation dates occurring in the month:	The values for i_k are:															
	i_1	i_2	i_3	i_4	i_5	i_6	i_7	i_8	i_9	i_{10}	i_{11}	i_{12}	i_{13}	i_{14}	i_{15}	i_u
*		*			*		*		*			*			*	
October 1992...	.06375	.0625	.06125	.06	.05875	.0575	.0575	.0575	.0575	.0575	.05625	.05625	.05625	.05625	.05625	.055

Issued at Washington, DC, on this 8th day of September 1992.

James B. Lockhart III,

Executive Director, Pension Benefit Guaranty Corporation.

[FR Doc. 92–22258 Filed 9–14–92; 8:45 am]

BILLING CODE 7708–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Care Financing Administration

42 CFR Parts 405, 413, 414, and 415

[BPD–712–CN]

RIN 0938–AE91

Medicare Program; Fee Schedule for Physicians' Services

AGENCY: Health Care Financing Administration (HCFA), HHS.

ACTION: Correction of final rule.

SUMMARY: In the November 25, 1991 final rule (56 FR 59624) on the Medicare fee schedule for physician services, we inadvertently set forth regulations on the fee schedule at 42 CFR, part 415. However, our plan for the recodification of HCFA regulations calls for general regulations on payment for Part B medical and other health services to be codified in part 414, with part 415 reserved for regulations on payment to teaching physicians, teaching hospitals, and provider-based physicians. Therefore, in this correction notice, we are redesignating in their entirety the physician fee schedule regulations contained in part 415, subpart A to part 414, subpart A, and reserving part 415 for future use.

Also, this document corrects technical errors that appeared in the final rule published in the **Federal Register** on November 25, 1991 (56 FR 59502) entitled "Medicare Program; Fee Schedule for Physicians' Services".

EFFECTIVE DATE: January 1, 1992.

FOR FURTHER INFORMATION CONTACT: Nancy Miller, (410) 966–4494.

SUPPLEMENTARY INFORMATION: In **Federal Register** Document 91–27785, beginning on page 59502, published November 25, 1991, make the following corrections:

Preamble

A. Page 59507

In column 1, paragraph g., the parenthetical reference "(§ 415.60)" is removed from the first sentence. Immediately following the parenthetical statement at the end of the first sentence, two new sentences are added to read: "We will address in a future FR document the coverage of psychological testing services furnished by CPs and CSWs. Until that document is published, we will continue our present policy of covering psychological testing if the service is furnished by a physician (including incident-to services) or by a qualified psychologist (whether or not a clinical psychologist)."

B. Page 59508

In column 2, section C.1., in the second paragraph the first sentence is corrected to read: "The work RVUs, along with the RVUs for practice expense, malpractice, and technical components, are considered to be 'initial' RVUs."

C. Page 59515

In column 2, paragraph (2), the third through fifth sentences are corrected to read: "We have added 0.01804 work RVUs, 0.00531 practice expense RVUs, and 0.00032 malpractice RVUs or approximately $0.73 per service to the office visits, office consultations, and emergency visit codes to reflect the work in EKG interpretations. We added 0.02706 work RVUs, 0.00807 practice