requirements under the titles. However, on July 23, 1997, representatives of the State of Florida notified FSIS that, because of a lack of funding, Florida will no longer continue to administer its State meat and poultry inspection programs after November 30, 1997. The representatives have requested that the Department assume responsibility for the meat and poultry inspection programs.

In view of the termination date, it is determined that the State of Florida would not effectively enforce requirements at least equal to those imposed under the titles. Therefore, the Secretary of Agriculture must designate the State of Florida under section 301(c)(3) of the FMIA and section 5(c)(3) of the PPIA. Therefore, on and after December 2, 1997, the provisions of the titles will apply to operations and transactions within the State of Florida, unless exempt under sections 23 or 301(c)(2) of the FMIA or sections 5(c)(2) or 15 of the PPIA.

Owners or operators of Florida's meat and poultry establishments wishing to continue operations after November 30, 1997, must contact the FSIS District Office in order to receive Federal inspection. This office will provide information concerning requirements and exemptions under the FMIA and the PPIA, applications for inspection, and requests for surveys of establishments. Address correspondence to USDA/FSIS District Office, 100 Alabama Street, SW, Suite 3R90, Atlanta, GA 30303.

The Administrator, FSIS, has determined that there is good cause for issuing this final rule without prior notice and opportunity for public comment. Because the State of Florida has advised FSIS that its State-operated meat and poultry inspection programs will be discontinued, the Agency is mandated by law to assume the responsibilities for administering the meat and poultry inspection programs. It is necessary, therefore, to designate the State of Florida immediately, in accordance with section 301(c)(3) of the FMIA and section 5(c)(3) of the PPIA, in order to carry out the Secretary's responsibilities under the FMIA and PPIA.

In addition, it does not appear that additional relevant information would be made available to the Secretary by public participation in this rulemaking proceeding. Accordingly, under the administrative procedures in 5 U.S.C. 553, it is found upon good cause that notice and other public procedures are impracticable and contrary to the public interest.

Executive Order 12866 and Regulatory Flexibility Act

This final rule has been determined to be not significant under Executive Order 12866 and, therefore, has not been reviewed by the Office of Management and Budget.

The Administrator, FSIS, has determined that this final rule will not have a significant economic impact on a substantial number of small entities, as defined by the Regulatory Flexibility Act (5 U.S.C. 601). The U.S. Department of Agriculture, pursuant to law, is assuming the responsibility, previously held by the State of Florida, of administering the meat and poultry inspection programs with respect to operations and transactions within the State of Florida. This action will affect approximately 122 State, 26 custom exempt, and 0 Talmadge Aiken meat and poultry establishments in Florida, and most, if not all, of which may be presumed to be small businesses. However, this is not a substantial number of establishments given the approximately 6,800 small meat and small poultry establishments nationwide, which are either federally or State inspected. In addition, the application of certain Federal facility and other requirements will be flexible, and each facility will be reviewed with regard to the circumstances peculiar to that establishment. Further, it is not anticipated that significant costs will be incurred by these Florida establishments as a result of this action.

Executive Order 12988

This final rule has been reviewed under Executive Order 12988, Civil Justice Reform. This rule: (1) Preempts State and local laws and regulations that are inconsistent with this rule; (2) has no retroactive effect; and (3) does not require administrative proceedings before parties may file suit in court challenging this rule. However, the administrative procedures specified in 9 CFR 306.5 and 381.35 must be exhausted prior to any judicial challenge of the application of the provisions of this rule, if the challenge involves any decision of an FSIS employee relating to inspection services provided under the FMIA or the PPIA.

Paperwork Requirements

This rule has been reviewed under the Paperwork Reduction Act and imposes no new paperwork or recordkeeping requirements.

List of Subjects

9 CFR Part 331

Meat inspection.

9 CFR Part 381

Poultry and poultry products.

Accordingly, 9 CFR parts 331 and 381 are amended as follows:

PART 331—SPECIAL PROVISIONS FOR DESIGNATED STATES AND TERRITORIES; AND FOR DESIGNATION OF ESTABLISHMENTS WHICH ENDANGER PUBLIC HEALTH AND FOR SUCH DESIGNATED ESTABLISHMENTS

1. The authority citation for part 331 continues to read as follows:

Authority: 21 U.S.C. 601–695; 7 CFR 2.18, 2.53.

§ 331.2 [Amended]

2. The table in section 331.2 is amended in the ''State'' column by adding ''Florida'' immediately below ''Connecticut'' and in the ''Effective date of application of Federal provisions'' column, by adding ''Dec. 2, 1997'' on the line with ''Florida.''

PART 381—POULTRY PRODUCTS INSPECTION

3. The authority citation for Part 381 continues to read as follows:

Authority: 7 U.S.C. 138f, 450; 21 U.S.C. 451–470; 7 CFR 2.18, 2.53.

§ 381.221 [Amended]

4. The table in section 381.221 is amended in the ''States'' column by adding ''Florida'' immediately below ''Connecticut'' and in the ''Effective date of application of Federal provisions'' column, by adding ''Dec. 2, 1997,'' on the line with ''Florida.''

Done at Washington, DC, on: November 4, 1997.

Thomas J. Billy,

Administrator.

[FR Doc. 97–29928 Filed 11–13–97; 8:45 am]

BILLING CODE 3410–DM–P

DEPARTMENT OF TRANSPORTATION

Federal Aviation Administration

14 CFR Part 39

[Docket No. 96–ANE–35; Amendment 39–10134; AD 97–19–13]

RIN 2120–AA64

Airworthiness Directives; Pratt & Whitney JT8D–200 Series Turbofan Engines

AGENCY: Federal Aviation Administration, DOT.

ACTION: Final rule; correction.

SUMMARY: This document makes a correction to Airworthiness Directive (AD) 97–19–13 applicable to Pratt & Whitney (PW) JT8D–200 series turbofan engines that was published in the **Federal Register** on September 19, 1997 (62 FR 49135). The paragraph references to the Accomplishment Instructions of PW Alert Service Bulletin (ASB) No. 5944, Revision 3, dated December 16, 1994, in paragraph (a)(3) of the compliance section are incorrect. This document corrects the paragraph references. In all other respects, the original document remains the same.

EFFECTIVE DATE: November 14, 1997.

FOR FURTHER INFORMATION CONTACT: Christopher Spinney, Aerospace Engineer, Engine Certification Office, FAA, Engine and Propeller Directorate, 12 New England Executive Park, Burlington, MA 01803–5299; telephone (781) 238–7175, fax (781) 238–7199.

SUPPLEMENTARY INFORMATION: A final rule airworthiness directive applicable to Pratt & Whitney (PW) JT8D–200 series turbofan engines, was published in the **Federal Register** on September 19, 1997 (62 FR 49135). The following correction is needed:

§ 39.13 [Corrected]

On page 49136, in the third column, in the Compliance Section, in paragraph (a)(3), in the sixth line, ''2.A.(2) (c) and (d) or (f) and (g)'' is corrected to read ''2.A.(2) (a) and (b) or (d) and (e)''.

Issued in Burlington, MA, on November 6, 1997.

Jay J. Pardee,

Manager, Engine and Propeller Directorate, Aircraft Certification Service.

[FR Doc. 97–29968 Filed 11–13–97; 8:45 am]

BILLING CODE 4910–13–U

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 558

New Animal Drugs For Use In Animal Feeds; Carbarsone and Bacitracin Zinc

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 an abbreviated new animal drug application (ANADA) filed by Alpharma Inc. The ANADA provides for using approved carbarsone and bacitracin zinc Type A medicated articles to make Type C medicated turkey feeds used for prevention of blackhead, increased rate of weight gain, and improved feed efficiency.

EFFECTIVE DATE: November 14, 1997.

FOR FURTHER INFORMATION CONTACT: Jeffrey M. Gilbert, Center for Veterinary Medicine (HFV–128), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–594–1602.

SUPPLEMENTARY INFORMATION: Alpharma Inc., One Executive Dr., P.O. Box 1399, Fort Lee, NJ 07024, is sponsor of ANADA 200–203 that provides for combining approved carbarsone and bacitracin zinc Type A medicated articles to make Type C medicated feeds for turkeys containing carbarsone 227 to 340.5 grams per ton (g/t) and bacitracin zinc 4 to 45 g/t. The Type C medicated feed is used as an aid in the prevention of blackhead, for increased rate of weight gain, and improved feed efficiency.

Alpharma Inc.'s, ANADA 200–203 is approved as a generic copy of Hoffmann-LaRoche's NADA 136–484. The ANADA is approved as of November 14, 1997, and the regulations are amended in § 558.120 (21 CFR 558.120) to reflect the approval. The basis for approval is discussed in the freedom of information summary.

In addition, § 558.120 is revised by redesignating paragraph (c) as (d), by reserving paragraph (c), and newly redesignated paragraph (d)(1)(iii)(*b*) is amended to reflect the approval.

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

The agency has determined under 21 CFR 25.33(a)(1) 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.

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.

§ 558.120 [Amended]

2. Section 558.120 *Carbarsone (not U.S.P.)* is amended by redesignating paragraph (c) as paragraph (d), by reserving paragraph (c), and in newly redesignated paragraph (d)(1)(iii)(*b*) by removing ''No. 000004'' and adding in its place ''Nos. 000004 and 046573''.

Dated: October 22, 1997.

Stephen F. Sundlof,

Director, Center for Veterinary Medicine.

[FR Doc. 97–30033 Filed 11–13–97; 8:45 am]

BILLING CODE 4160–01–F

OCCUPATIONAL SAFETY AND HEALTH REVIEW COMMISSION

29 CFR Part 2200

Rules of Procedure for E–Z Trials

AGENCY: Occupational Safety and Health Review Commission.

ACTION: Final rule; correcting amendment.

SUMMARY: This document restores the selection provision for commencing E–Z Trial, 29 CFR 2200.203(a), which was inadvertently removed.

DATES: November 14, 1997.

FOR FURTHER INFORMATION CONTACT: Earl R. Ohman, Jr., General Counsel, (202) 606–5410, Occupational Safety and Health Review Commission, 1120 20th Street NW., 9th Floor, Washington, DC 20036–3419.

SUPPLEMENTARY INFORMATION: On October 30, 1997, (62 FR 58650), paragraph (a) of § 2200.203 was inadvertently removed. In order for the Rules of Procedures for E–Z trial to operate effectively, paragraph (a) must be restored.

List of Subjects in 29 CFR Part 2200

Administrative practice and procedure, Hearing and appeal procedures.

For the reasons set forth in the preamble, the Occupational Safety and Health Review Commission amends Title 29, Chapter XX, Part 2200, Subpart M of the Code of Federal Regulations as follows:

PART 2200—RULES OF PROCEDURE

1. The authority citation continues to read as follows:

Authority: 29 U.S.C. 661(g).

2. Section 2200.203 is amended by adding paragraph (a), to read as follow: