

provisions of paragraph (b) (13) are amended to read as follows:

§ 905.559 Export Regulation 24.

(b) * * *

(13) Any seedless grapefruit, other than pink seedless grapefruit, grown in the production area, which do not grade at least Improved No. 2, or any pink seedless grapefruit which do not grade at least U.S. No. 2 Russet;

(Secs. 1-19, 48 Stat. 31, as amended; (7 U.S.C. 601-674).)

Dated: March 6, 1975, to become effective March 10, 1975.

CHARLES R. BRADER,
Deputy Director, Fruit and
Vegetable Division, Agricultural
Marketing Service.

[FR Doc.75-6306 Filed 3-10-75; 8:45 am]

[Grapefruit Reg. 15, Amdt. 3]

**PART 944—FRUITS; IMPORT
REGULATIONS**

**Minimum Grade Requirements for Imports
of Pink Seedless Grapefruit**

This amendment lowers the minimum grade requirement applicable to imported pink seedless grapefruit to U.S. No. 2 Russet on March 10, 1975. The requirement is the same as that applicable to grapefruit produced in Florida and regulated pursuant to Marketing Order No. 905.

This amendment is consistent with Section 8e of the Agricultural Marketing Agreement Act of 1937, as amended (7 U.S.C. 601-674). This section requires that whenever specified commodities, including grapefruit, are regulated under a federal marketing order, imports of that commodity must meet the same or comparable grade requirements as those in effect for the domestically produced commodity. This regulation fixes the same minimum grade requirement on imported pink seedless grapefruit as is effective under Marketing Order No. 905, as amended (7 CFR Part 905) regulating the handling of oranges, grapefruit, tangerines and tangelos grown in Florida.

Order. In § 944.111 (Grapefruit Regulation 15; 39 FR 33306, 37188; 40 FR 8322) paragraphs (a) (1), (2), and (3) are revised to read as follows:

§ 944.111 Grapefruit Regulation 15.

(a) * * *

(1) Seeded grapefruit shall grade at least U.S. No. 1 and be of a size not smaller than 3 1/16 inches in diameter except that a tolerance for seeded grapefruit smaller than such minimum size shall be permitted as specified in § 51.761 of the United States Standards for Florida Grapefruit;

(2) Seedless grapefruit, other than pink seedless grapefruit, shall grade at least Improved No. 2 ("Improved No. 2" shall mean grapefruit grading at least U.S. No. 2 and also meeting the requirements of the U.S. No. 1 grade as to shape (form) and color), and pink seedless

grapefruit shall grade at least U.S. No. 2 Russet; and

(3) Seedless grapefruit, other than pink seedless grapefruit, shall be of a size not smaller than 3 1/16 inches in diameter, and pink seedless grapefruit shall be of a size not smaller than 3 3/16 inches in diameter, except that a tolerance for seedless grapefruit smaller than such minimum sizes shall be permitted as specified in § 51.761 of the United States Standards for Florida Grapefruit.

It is hereby found that it is impracticable, unnecessary, and contrary to the public interest to give preliminary notice, engage in public rule-making procedure, and postpone the effective time of this amendment beyond that hereinafter specified (5 U.S.C. 553) in that (a) the requirements of this amended import regulation are imposed pursuant to section 8e of the Agricultural Marketing Agreement Act of 1937, as amended (7 U.S.C. 601-674), which makes such regulation mandatory; (b) this amendment fixes the same requirement for imports of pink seedless grapefruit as is applicable under amended Grapefruit Regulation 75 (§ 905.556 to the shipment of pink seedless grapefruit grown in Florida; and (c) this amendment lowers the minimum grade requirement applicable to imported pink seedless grapefruit.

(Secs. 1-19, 48 Stat. 31, as amended; (7 U.S.C. 601-674).)

Dated: March 6, 1975, to become effective March 10, 1975.

CHARLES R. BRADER,
Deputy Director, Fruit and Veg-
etable Division, Agricultural
Marketing Service.

[FR Doc.75-6305 Filed 3-10-75; 8:45 am]

**Title 9—Animal and Animal Products
CHAPTER I—ANIMAL AND PLANT HEALTH
INSPECTION SERVICE, DEPARTMENT
OF AGRICULTURE**

**SUBCHAPTER D—EXPORTATION AND IMPORTA-
TION OF ANIMALS (INCLUDING POULTRY)
AND ANIMAL PRODUCTS**

**PART 97—OVERTIME SERVICES
RELATING TO IMPORTS AND EXPORTS**

Commuted Traveltime Allowances

The purpose of this amendment is to establish commuted traveltime periods as nearly as may be practicable to cover the time necessarily spent in reporting to and returning from the place at which an employee of Veterinary Services performs overtime or holiday duty when such travel is performed solely on account of overtime or holiday duty. Such establishment depends upon facts within the knowledge of the Animal and Plant Health Inspection Service.

Therefore, pursuant to the authority conferred upon the Deputy Administrator, Veterinary Services, Animal and Plant Health Inspection Service by § 97.1 of the regulations concerning overtime services relating to imports and exports (9 CFR 97.1), administrative instructions 9 CFR 97.2 (1974 ed.), as amended

November 27, 1974 (39 FR 41356-41358), December 11, 1974 (39 FR 43294), January 3, 1975 (40 FR 757), and February 21, 1975 (40 FR 7620), prescribing the commuted traveltime that shall be included in each period of overtime or holiday duty, are hereby amended by adding to the respective lists therein as follows:

**§ 97.2 Administrative instructions pre-
scribing commuted travel time.**

WITHIN METROPOLITAN AREA

ONE HOUR

Helena, Montana.

(64 Stat. 561; (7 U.S.C. 2260).)

Effective date. The foregoing amendment shall become effective on March 11, 1975.

It is to the benefit of the public that this instruction be made effective at the earliest practicable date. It does not appear that public participation in this rulemaking proceeding would make additional relevant information available to the Department.

Accordingly, pursuant to 5 U.S.C. 553, it is found upon good cause that notice and public procedure on this instruction are impracticable, unnecessary, and contrary to the public interest and good cause is found for making it effective less than 30 days after publication in the FEDERAL REGISTER.

Done at Washington, D.C., this 6th day of March, 1975.

J. M. HEJL,
Deputy Administrator, Veteri-
nary Services, Animal and
Plant Health Inspection Serv-
ice.

[FR Doc.75-6307 Filed 3-10-75; 8:45 am]

**CHAPTER III—ANIMAL AND PLANT
HEALTH INSPECTION SERVICE, (MEAT
AND POULTRY PRODUCTS INSPEC-
TION), DEPARTMENT OF AGRICULTURE**

**SUBCHAPTER A—MANDATORY MEAT
INSPECTION**

**TENANTS AND SUBSIDIARIES IN
OFFICIAL ESTABLISHMENTS**

**Application for Grant of Inspection;
Correction**

The notice of final rulemaking published in the FEDERAL REGISTER of January 14, 1975 (40 FR 2575, FR Doc. 75-1043) concerning the requirements for application for grant of inspection by tenants and subsidiaries in official establishments contained an error. The second paragraph of the statement of considerations stated that the effective date of the change had been delayed one year from the date of publication. The effective date used, however, was July 14, 1976. The effective date used is correct. The words should read: "has accordingly been delayed for eighteen months from publication" rather than "has accordingly been delayed for one year from publication."

Done at Washington, D.C., on: March 5, 1975.

F. J. MULHERN,
Administrator, Animal and Plant
Health Inspection Service.

[FR Doc.75-6309 Filed 3-10-75;8:45 am]

**PART 317—LABELING, MARKING
DEVICES, AND CONTAINERS**

**PART 381—POULTRY PRODUCTS
INSPECTION REGULATIONS**

**Spanish Labeling of Products Distributed
Solely in Puerto Rico**

The Animal and Plant Health Inspection Service, pursuant to the authority conferred by the Federal Meat Inspection Act, as amended (21 U.S.C. 601 et seq.), and the Poultry Products Inspection Act, as amended (21 U.S.C. 451 et seq.), is amending § 317.2 of the meat inspection regulations (9 CFR Part 317) and § 381.116 of the poultry products inspection regulations (9 CFR Part 381) to provide for the labeling of products in the Spanish language when intended for distribution solely in Puerto Rico.

Statement of Considerations. On April 23, 1974, there were published in the Federal Register (39 FR 14351, 14352) proposed amendments to the meat inspection regulations and the poultry products inspection regulations. The proposal provided for labeling in Spanish of products distributed solely in Puerto Rico and gave interested parties until May 24, 1974, to comment.

It has been the policy of this Department's meat and poultry inspection labeling program for many years to require that mandatory labeling information be in English. However, regulations with respect to labeling of meat products for export (9 CFR 317.7) and for poultry products for foreign commerce (9 CFR 381.128) allow the presentation of mandatory labeling information, except for the inspection legend and establishment number, in a language other than English.

As processors and distributors attempt to provide more consumer information on the labeling and as new labeling requirements are imposed, the quantity of labeling information will expand. This expansion may result in severe space limitation for the display of legible information on the labels of small containers, such as baby foods.

The predominant language in Puerto Rico is Spanish. These amendments bring the Federal meat inspection regulations and the poultry products inspection regulations into accord with the regulations of the Food and Drug Administration, under which products distributed solely in Puerto Rico may be labeled in Spanish in lieu of English.

No comments were received concerning the proposed amendments. Therefore, it is concluded that there are no objections to the proposal and that the amendments as proposed do indeed provide desirable changes in the regulations. The Department has noted that

§ 317.7 of the meat inspection regulations and § 381.128 of the poultry products inspection regulations could be construed as in conflict with the addition as proposed. Therefore, a clarifying phrase has been added to the proposed regulations in cognizance of these sections.

With the exception of the clarification noted above, the amendments to the Federal meat inspection regulations and the poultry products inspection regulations are adopted as set forth in the proposal.

(Sec. 21, 34 Stat. 1260, as amended, 21 U.S.C. 621; sec. 14, 71 Stat. 441, as amended, 21 U.S.C. 463; 37 FR 23464, 23477)

The foregoing amendments shall become effective April 14, 1975.

Done at Washington, D.C., on: March 5, 1975.

F. J. MULHERN,
Administrator, Animal and Plant
Health Inspection Service.

1. Section 317.2 is amended by adding the following to the end of paragraph (b):

§ 317.2 Labels: definition; required features.

(b) * * * Except as provided in § 317.7, all words, statements, and other information required by or under authority of the Act to appear on the label or labeling shall appear thereon in the English language: *Provided, however,* That in the case of products distributed solely in Puerto Rico, Spanish may be substituted for English for all printed matter except the USDA inspection legend.

2. Section 381.116 is amended by adding the following to the end of paragraph (a):

§ 381.116 Wording on labels of immediate containers.

(a) * * * Except as provided in § 381.128, all words, statements and other information required by or under authority of the Act to appear on the label or labeling shall appear thereon in the English language: *Provided, however,* That in the case of products distributed solely in Puerto Rico, Spanish may be substituted for English for all printed matter except the USDA inspection legend.

[FR Doc.75-6310 Filed 3-10-75;8:45 am]

**Title 18—Conservation of Power and Water
Resources**

**CHAPTER I—FEDERAL POWER
COMMISSION**

**SUBCHAPTER D—APPROVED FORMS,
FEDERAL POWER ACT**

[Docket No. RM 75-21; Order No. 527]

**PART 141—STATEMENTS AND REPORTS
(SCHEDULES)**

**Order Prescribing Revision of FPC Form
No. 12F, Power Line and Generating
Plant Data**

Consonant with purposes of this Commission's Order No. 383-2 issued April 10,

1970, FPC Form No. 12F is hereby amended to accommodate the following changes resulting in simplification of the Form and causing it to conform with the established standards of Order No. 383-2.

Instruction No. 4 and its reporting Schedule D, identified "Scheduled Changes in System Generating Plants", of Form No. 12F are deleted to avoid needless duplication of data which is also presently submitted in response to FPC Form No. 12E1. This deletion necessitates revision of the Form title to "Power Line Construction Data", as well as revision of the title of § 141.57 to read "Form No. 12F, Power Line Construction Data".

The retitling of existing Schedule C of Form No. 12F to read "Lines Proposed for Installation in Next 6 Years But Not Under Construction" is made to conform with Order No. 383 data requirements and to eliminate confusion over the meaning of the present term "authorized".

Coding symbols are also changed for the purpose of simplification, and the addition of the name of the respondents to the back of the Form is made so that copying may be accomplished without loss of respondent identification.

The Form No. 12F adopted revisions do not require issuance of a notice of proposed rulemaking thereupon because no additional substantive information is to be collected and the overall effect of the revisions is to reduce respondent burden.

The Commission finds. (1) The revisions to FPC Form No. 12F as herein prescribed are necessary and appropriate for the administration of the Federal Power Act.

(2) Since the revisions to FPC Form No. 12F prescribed herein do not substantially alter the form and are primarily for the purpose of simplification or clarification, and since the adoption of such changes reduces the reporting burden and relates to matters of agency practice and procedure necessary and appropriate for the administration of the Federal Power Act, compliance with the notice, public procedure and effective date provisions of 5 U.S.C. 553 is unnecessary.

(3) Since the revisions to FPC Form No. 12F prescribed herein are for use for reports covering the calendar year ending June 30, 1975, good cause exists for making this order effective upon issuance.

The Commission, acting pursuant to the provisions of the Federal Power Act, as amended, particularly sections 309 and 311 (49 Stat. 858, 859 (16 U.S.C. 825h, 825j)) orders. (A) Deletion of Instruction No. 4 and its reporting Schedule D, identified "Scheduled Changes in System Generating Plants", from FPC Form No. 12F.

Revision of the title of Form No. 12F from "Power Line and Generating Sta-

¹ Form No. 12F is filed as part of the original document.

tion Construction Data" to "Power Line Construction Data";

Revision of the title of § 141.57 of the Commission's rules (18 CFR 141.57) from "Form No. 12F, Power Line and Generating Plant Data" to "Form No. 12F, Power Line Construction Data";

Retitling of Schedule C of Form No. 12F to "Lines Proposed for Installation in Next 6 Years But Not Under Construction";

Addition of the name of respondents to the back of Form No. 12F; and

Change of the coding symbols as indicated within the Form.

(B) The revisions adopted herein shall be effective upon issuance of this order.

(C) The Secretary shall cause prompt publication of this order to be made in the FEDERAL REGISTER.

By the Commission.

[SEAL] KENNETH F. PLUMB,
Secretary.

[FR Doc. 75-6219 Filed 3-19-75; 8:45 am]

Title 21—Food and Drugs

CHAPTER I—FOOD AND DRUG ADMINISTRATION, DEPARTMENT OF HEALTH, EDUCATION, AND WELFARE

SUBCHAPTER C—DRUGS

SUBCHAPTER D—DRUGS FOR HUMAN USE

Sterile Benzathine Cloxacillin for Intramammary Infusion

The Commissioner of Food and Drugs has evaluated a new animal drug application (55-068V) filed by E. R. Squibb & Sons, Inc., P.O. Box 4000, Princeton, NJ 08540, proposing the safe and effective use of sterile benzathine cloxacillin suspension for intramammary infusion in the treatment of bovine mastitis in dry cows. The application is approved.

The mailing address of the firm has been changed, and the address included in the regulations is changed accordingly.

Therefore, pursuant to provisions of the Federal Food, Drug, and Cosmetic Act (sec. 512 (i) and (n), 82 Stat. 347, 350-351; (21 U.S.C. 360b (i) and (n))) and under authority delegated to the Commissioner (21 CFR 2.120), Chapter I of Title 21 of the Code of Federal Regulations is amended as follows:

PART 135—NEW ANIMAL DRUGS

1. In § 135.501(c) by changing the address of code No. 035 as follows:

§ 135.501 [Amended]

Code No.: Firm name and address
035----- E. R. Squibb & Sons,
Inc., P.O. Box 4000,
Princeton, N.J. 08540.

PART 135d—NEW ANIMAL DRUGS FOR INTRAMAMMARY USE

2. By adding the following new section:

§ 135d.15 Sterile benzathine cloxacillin for intramammary infusion, veterinary.

(a) *Specifications.* The drug is sterile and contains benzathine cloxacillin equivalent to 500 milligrams cloxacillin in each 6 milliliters of peanut oil vehicle, and conforms to the certification requirements of § 149j.11 of this chapter.

(b) *Sponsor.* See code No. 035 in § 135.501(c) of this chapter.

(c) *Conditions of use.* (1) The drug is used for treatment of mastitis caused by *Staphylococcus aureus* and *Streptococcus agalactiae* in dairy cows at the time of drying-off of the cow.

(2) It is administered aseptically at the rate of 6 milliliters per infected quarter immediately after last milking at the time of drying-off of the cow.

(3) For use in dry cows only.

(4) Not to be used within 30 days of calving.

(5) Milk taken from treated cows prior to 72 hours (6 milkings) after calving must not be used for human food.

(6) Animals infused with this product the time of infusion until 72 hours after must not be slaughtered for food from calving.

(7) Federal law restricts this drug to use by or on the order of a licensed veterinarian.

3. By adding a new Part 149j consisting at this time of two sections to read as follows:

PART 149j—CLCXCACILLIN FOR VETERINARY USE

Sec. 149j.1 Sterile benzathine cloxacillin, veterinary.

149j.11 Sterile benzathine cloxacillin for intramammary infusion, veterinary.

AUTHORITY: Sec. 512 (i) and (n), 82 Stat. 347, 350-351; (21 U.S.C. 360b (i) and (n)).

§ 149j.1 Sterile benzathine cloxacillin, veterinary.

(a) *Requirements for certification.* (1) *Standards of identity, strength, quality, and purity.* Benzathine cloxacillin is the *N,N'*-dibenzylethylenediamine salt of 5-methyl-3-(*o*-chlorophenyl)-4-isoxazolyl penicillin. It is so purified and dried that:

(i) Its potency is not less than 704 nor more than 821 micrograms of cloxacillin per milligram on an anhydrous basis.

(ii) It is sterile.

(iii) It passes the safety test.

(iv) Its moisture content is not more than 5.0 percent.

(v) Its pH in an aqueous suspension containing 10 milligrams per milliliter is not less than 3.0 nor more than 6.5.

(vi) It passes the identity test.

(vii) It is crystalline.

(2) *Labeling.* It shall be labeled in accordance with the requirements of § 432.5 of this chapter.

(3) *Requests for certification; samples.* In addition to complying with the requirements of § 146.2 of this chapter, each such request shall contain:

(i) Results of tests and assays on the batch for potency, sterility, safety, moisture, pH, identity, and crystallinity.

(ii) Samples required:

(A) For all tests except sterility: 10 packages, each containing approximately 300 milligrams.

(B) For sterility testing: 20 packages, each containing approximately 600 milligrams.

(b) *Tests and methods of assay.* (1) *Potency.* Use the microbiological agar diffusion assay method: Proceed in accordance with § 436.105 of this chapter, using the cloxacillin working standard as the standard of comparison and prepare the sample for assay as follows: Dissolve an accurately weighed portion of the sample in sufficient methanol to give a convenient stock solution. Immediately dilute an aliquot of this stock solution with solution 1 to the reference concentration of 5.0 micrograms of cloxacillin per milliliter estimated.

(2) *Sterility.* Proceed as directed in § 436.20 of this chapter, using the method described in paragraph (e) (2) of that section, except use medium C in lieu of medium A, medium F in lieu of medium E, and during the period of incubation shake the tubes at least once daily.

(3) *Safety.* Proceed in accordance with § 436.33 of this chapter.

(4) *Moisture.* Proceed in accordance with § 436.201 of this chapter.

(5) *pH.* Proceed in accordance with § 436.202 of this chapter, using an aqueous suspension prepared by adding 10 milligrams per milliliter.

(6) *Identity.* Transfer approximately 20 milligrams of the sample to a 50-milliliter Erlenmeyer flask. Add 5.0 milliliters of 5N sodium hydroxide and heat in a steam bath 20 minutes. Cool. Transfer 1 milliliter to an extraction funnel; add approximately 10 milliliters of water and 1 milliliter of dilute sulfuric acid (1:2). Shake with 50 milliliters of ether. Discard the aqueous layer and wash the ether layer with approximately 30 milliliters of water. Discard the aqueous layer again and extract with approximately 50 milliliters of 0.1N sodium hydroxide. Obtain a spectrum of the 0.1N sodium hydroxide solution from 300 nanometers to 240 nanometers against a reagent blank. Treat about 15 milligrams of the cloxacillin working standard in the same manner. The sample is satisfactory if the spectrum obtained from the sample solution matches that of the standard solution with maximum at about 282 nanometers and minimum of about 257 nanometers.

(7) *Crystallinity.* Proceed as directed in § 436.203 of this chapter.