requested by the Board, on information collection forms that have expired. The Board is removing references to the obsolete forms.

Regulatory Flexibility Act Analysis

Pursuant to section 605(b) of the Regulatory Flexibility Act (Pub. L. No. 96-354, 5 U.S.C. 601 *et seq.*), the Board certifies that the amendments would not have a significant economic impact on a substantial nubmer of small entities. The amendments simplify or reduce certain regulatory burdens for all depository institutions and have no particular effect on other small entities.

List of Subjects in 12 CFR Part 208

Membership, Banks, Accounting, Confidential business information, Federal Reserve System, Reporting and recordkeeping requirements, Securities, Disclosures of financial information.

Pursuant to the Board's authority under section 9 of the Federal Reserve Act, 12 U.S.C. 321 *et seq.*, the Board is amending 12 CFR Part 208 as follows:

PART 208—MEMBERSHIP OF STATE BANKING INSTITUTIONS IN THE FEDERAL RESERVE SYSTEM

1. The authority citation for 12 CFR Part 208 continues to read as follows:

Authority: Sections 9, 11(a), 11(c), 19, 21, 25, and 25(a) of the Federal Reserve Act, as amended (12 U.S.C. 321–338, 248(a), 248(c), 461, 481–486, 601, and 611, respectively); sections 4 and 13(j) of the Federal Deposit Insurance Act, as amended (12 U.S.C. 1814 and 1923(j), respectively); section 7(a) of the International Banking Act of 1978 (12 U.S.C. 3105); sections 907–910 of the International Lending Supervision Act of 1983 (12 U.S.C. 3906–3909); sections 2, 12(b), 12(g), 12(i), 15B(c)(5), 17, 17A, and 23 of the Securities Exchange Act of 1934 (15 U.S.C. 78b, 78*l*(b), 78*l*(g), 78*l*(i), 78o–4(c)(5), 78q, 78q–1, and 78w, respectively); and section 5155 of the Revised Statutes (12 U.S.C. 36) as amended by the McFadden Act of 1927.

2. Section 208.10(a)(3) is amended by changing the words "(Form FR 105a)" to read "(Forms FFIEC 031–034)" and by revising the last sentence to read as follows:

§ 208.10 [Amended]

(a) * * *

(3) * * * All signatures shall be the same in the published statement (although they may be typed or otherwise copied on the report for publication):

(i) As in the original report submitted to the Federal Reserve Bank if the bank does not submit its report of condition electronically, or

(ii) As retained in the bank's files in hard copy if the bank has filed its report of condition electronically. The hard copy retained in the bank's file must be made available to examiners upon request.

* * * * *

3. Section 208.10(a)(4) is revised to read as follows:

(a) * * *

(4) A copy of the printed report shall be retained in the bank's files and made available to examiners upon request.

* * * * *

4. Section 208.10(b) is amended by removing the first sentence in paragraph (2) and removing the words "attached to the certificate on Form FR 220a" at the end of paragraph (3).

By order of the Board of Governors of the Federal Reserve System, February 13, 1989.

William W. Wiles,

Secretary of the Board.

[FR Doc. 89-3713 Filed 2-16-89; 8:45 am]

BILLING CODE 6210-01-M

DEPARTMENT OF TRANSPORTATION

Research and Special Programs Administration

14 CFR Parts 217 and 241

[Docket No. 44999; Amendment No. 217-2; 241-57]

RIN 2137-AA97, 2137-AB01

Aviation Economic Regulations; Report of Traffic and Capacity Statistics; Collection of Service Segment and Charter Data; the "T-100 System"; Correction

AGENCY: Research and Special Programs Administration, DOT.

ACTION: Final rule; correction.

SUMMARY: This document corrects the final rule amending 14 CFR Parts 217 and 241 (Docket 44999) published in the **Federal Register** on November 16, 1988 (53 FR 46284). The rule established a new traffic reporting system known as the "T-100 System" for U.S. and foreign air carriers. This correction is not intended to address the pending petitions for reconsideration of the final rule (see notice published in the **Federal Register** on December 28, 1988 (53 FR 52404) suspending the effective date of the rule for all foreign air carriers.)

FOR FURTHER INFORMATION CONTACT: Donald Bright or Richard King, Office of Aviation Information Management, DAI-10, Research and Special Programs Administration, Department of Transportation, 400 Seventh Street SW., Washington, DC 20590, (202) 366-4384, or 366-4375, respectively.

SUPPLEMENTARY INFORMATION: In FR Doc. 88-26322 published in the **Federal Register** on Wednesday, November 16, 1988, corrections are made as follows:

1. On page 46291, the first column, the second full paragraph is corrected to read:

The Department has decided that it will not separately collect data behind a foreign carrier's homeland. Although not separately reported, these data will be included with the homeland data and reported as if the traffic enplaned or deplaned at the homeland. Except for the behind homeland data, the Department requires that traffic for any segment or market that includes a U.S. airport shall be duly reported. In its decision, the Department took into consideration the fact that other countries normally do not collect such information.

PART 217—[CORRECTED]

§ 217.5 [Corrected]

2. On page 46295, in the first column, § 217.5(b)(7), is corrected to read:

* * * * *

(b) * * *

(7) Revenue aircraft departures performed (Code 510). The number of revenue aircraft departures performed.

* * * * *

§ 217.10 [Corrected]

3. On page 46295, in the second column, in § 217.10 appendix, paragraph (a)(3), in the second line, "DAI-2" is corrected to "DAI-20."

4. On page 46299, in the second column, paragraph (g)(2)(ii) of § 217.10 appendix, is corrected by removing the last sentence.

PART 241—[CORRECTED]

Sec. 19–5—[Corrected]

5. On page 46307, section 19–5(c)(23), is correctly revised to read:

* * * * *

(c) * * *

(23) Revenue aircraft departures performed. The number of revenue aircraft departures performed.

* * * * *

§ 241.25 [Appendix corrected]

6. On page 46311, in the appendix to § 241.25, paragraph (j)(1), Field No. 18, positions 99–103, Mode 5N of the segement record layout in the description column is corrected to read: "Revenue aircraft departures scheduled (F, G520)"

7. Also in the appendix to § 241.25, the portion of Form 41 Schedule 100 appearing on page 46315 is corrected to read as follows:

BILLING CODE 4910-62-M

A-1. Air Carrier Name: _____ Code_____

A-2. Report Date: (Year) _____ (Month)_____

					C. ON-FLIGHT MARKET				
B-10 Revenue Freight Transp. (Pounds) F,L237 G,P237	B-11 Revenue Mail Transp. (Pounds) F,L239 G,P239	B-12 Revenue Acft. Dept. Sched. F520 6520	B-13 Revenue Aircraft Hours Ramp F,L630 G,P630	B-14 Revenue Aircraft Hours Airborne F,L610 G,P610	C-1 * Revenue Psgrs. Enplaned TOTAL/ First F,L110 F111	C-2 Revenue Psgrs. Enplaned Middle F113	C-3 Revenue Psgrs. Enplaned Coach F112	C-4 Revenue Freight Enplaned (Pounds) F,L217 G,P217	C-5 Revenue Mail Enplaned (Pounds) F,L219 G,P219
----------					-- Total for all aircraft types in market---				

BILLING CODE 4910-62-C

8. In the same appendix, Form 41 Schedule T–2 appearing on page 46317 is corrected as follows:

BILLING CODE 4910–62–M

(3) Form 41 Schedule T-2:

Provides data QUARTERLY to supplement detail T-100.

FORM 41 SCHEDULE T-2 U.S. AIR CARRIER TRAFFIC AND CAPACITY BY AIRCRAFT TYPE		Air Carrier Name:_____ Code:_____ Entity Code: _____ Report date: (Year)_____ (Month)_____			
		Aircraft Type Code:	Aircraft Type Code:	Aircraft Type Code:	Aircraft Type Code:
		_____	_____	_____	_____
SCHEDULED ALL-CARGO SERVICES:					
Revenue ton-miles	G240				
Available ton-miles	G280				
Revenue aircraft miles flown	G410				
Aircraft departures performed	G510				
NONSCHEDULED SERVICES:					
Aircraft departures performed	V510				
ALL SERVICES:					
Revenue passenger-miles (000)	Z140				
Available seat-miles (000)	Z320				
Revenue ton-miles	Z240				
Mail revenue ton-miles	Z249				
Freight revenue ton-miles	Z247				
Available ton-miles	Z280				
Revenue aircraft miles flown	Z410				
Aircraft departures performed	Z510				
Revenue aircraft hours (airborne)	Z610				
Revenue aircraft hours (ramp)	Z630				
Total aircraft hours (airborne)	Z650				
Aircraft days - equipment	Z810				
Aircraft days - routes	Z820				
Aircraft fuels issued	Z921				

RSPA Form 41 Schedule T-2

BILLING CODE 4910-62-C

Issued in Washington, DC, on February 7, 1989.

M. Cynthia Douglass,

Administrator, Research and Special Programs Administration, DOT.

[FR Doc. 89-3333 Filed 2-16-89; 8:45 am]

BILLING CODE 4910-62-M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 189

[Docket No. 87N-0055]

Substances Prohibited From Use in Human Food; Hydrogenated 4,4'-Isopropylidene-Diphenolphosphite Ester Resins

AGENCY: Food and Drug Administration.

ACTION: Final rule.

SUMMARY: The Food and Drug Administration (FDA) is amending the food additive regulations by adding hydrogenated 4,4'-isopropylidene-diphenolphosphite ester resins to the list of substances that are prohibited from use in human food. FDA is taking this action because there are no studies that establish safe conditions of use for this additive.

DATES: Effective March 20, 1989, except to any provisions that may be stayed by the filing of proper objections; written objections and requests for a hearing by March 20, 1989.

ADDRESS: Written objections to the Dockets Management Branch (HFA-305), Food and Drug Administration, Rm. 4-62, 5600 Fishers Lane, Rockville, MD 20857.

FOR FURTHER INFORMATION CONTACT: Marvin D. Mack, Center for Food Safety and Applied Nutrition (HFF-335), Food and Drug Administration, 200 C St. SW., Washington, DC 20204, 202-472-5690.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of September 9, 1987 (52 FR 33952), FDA issued a proposal to add hydrogenated 4,4'-isopropylidene-diphenolphosphite ester resins to the list of substances that are prohibited from use in human food (21 CFR Part 189). In the same issue of the **Federal Register** (52 FR 33929), the agency issued a final rule that removed the listing for hydrogenated 4,4'-isopropylidene-diphenolphosphite ester resins from § 178.2010 *Antioxidants and/or stabilizers for polymers* (21 CFR 178.2010). FDA took these actions because adverse neurological effects were observed in a study in which dogs were fed the additive, and because there are no studies that establish safe conditions of use for this additive.

FDA gave interested persons 60 days (until November 9, 1987) to file comments on the proposal to list hydrogenated 4,4'-isopropylidene-diphenolphosphite ester resins in 21 CFR Part 189 and 30 days (until October 9, 1987) to file objections to the revocation of its listing in the food additive regulations. No comments were received on the proposal (52 FR 33952), and no objections or requests for a hearing were received in response to the final rule (52 FR 33929) revoking the use of the additive. Therefore, FDA is listing hydrogenated 4,4' isopropylidene-diphenolphosphite ester resins as substances prohibited from use in food-contact surfaces, as proposed.

The agency has previously considered the environmental effects of this rule as announced in the proposed rule (52 FR 33952). No new information or comments have been received that would affect the agency's previous determination that there is no significant impact on the human environment and that an environmental impact statement is not required.

In accordance with the Regulatory Flexibility Act, the agency previously considered the potential effects that this rule would have on small entities, including small businesses. In accordance with section 605(b) of the Regulatory Flexibility Act, the agency has determined that no significant impact on a substantial number of small entities would derive from this action. FDA has not received any new information or comments that would alter its previous determination.

In accordance with Executive Order 12291, FDA has previously evaluated the economic impact of this final rule. The agency has determined that the final rule is not a major rule as defined by Executive Order 12291. The agency's findings of no major economic impact and no significant impact on a substantial number of small entities, and the evidence supporting these findings, are contained in a threshold assessment displayed in the Dockets Management Branch (address above).

Any person who will be adversely affected by this regulation may at any time on or before March 20, 1989, file with the Dockets Management Branch (address above) written objections thereto. Each objection shall be separately numbered, and each numbered objection shall specify with particularity the provisions of the regulation to which objection is made and the grounds for the objection. 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. Three copies of all documents shall be submitted and shall be identified with the docket number found in brackets in the heading of this document. Any objections received in response to the regulation 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 189

Food ingredients, Food packaging, Prohibited food ingredients.

Therefore, under the Federal Food, Drug, and Cosmetic Act and under authority delegated to the Commissioner of Food and Drugs, Part 189 is amended as follows:

PART 189—SUBSTANCES PROHIBITED FROM USE IN HUMAN FOOD

1. The authority citation for 21 CFR Part 189 continues to read as follows:

Authority: Secs. 201(s), 402, 409, 701, 52 Stat. 1046-1047 as amended, 1055-1056 as amended, 72 Stat. 1784-1788 as amended (21 U.S.C. 321(s), 342, 348, 371); 21 CFR 5.10.

2. New § 189.300 is added to Subpart C to read as follows:

§ 189.300 Hydrogenated 4,4'-isopropylidene-diphenolphosphite ester resins.

(a) Hydrogenated 4,4'-isopropylidene-diphenolphosphite ester resins are the condensation product of 1 mole of triphenyl phosphite and 1.5 moles of hydrogenated 4,4'-isopropylidene-diphenol such that the finished resins have a molecular weight in the range of 2,400 to 3,000. They are synthetic chemicals not found in natural products and have been used as antioxidants and as stabilizers in vinyl chloride polymer resins when such polymer resins are used in the manufacture of rigid vinyl chloride polymer bottles.

(b) Food containing any added or detectable levels of these substances is deemed to be adulterated and in violation of the Federal Food, Drug, and Cosmetic Act, based upon an order

published in the **Federal Register** of September 9, 1987 (52 FR 33929).

Dated: February 10, 1989.

Alan L. Hoeting,

Acting Associate Commissioner for Regulatory Affairs.

[FR Doc. 89-3730 Filed 2-16-89; 8:45 am]

BILLING CODE 4160-01-M

21 CFR Part 558

New Animal Drugs for Use in Animal Feeds; Bacitracin Methylene Disalicylate

AGENCY: Food and Drug Administration.

ACTION: Final rule.

SUMMARY: The Food and Drug Administration (FDA) is amending the animal drug regulations to reflect approval of a supplemental new animal drug application (NADA) filed by A. L. Laboratories, Inc., providing for use of Type A medicated articles containing 25, 40, and 50 grams of bacitracin methylene disalicylate per pound to make Type C medicated feeds for the prevention of ulcerative enteritis in quail.

EFFECTIVE DATE: February 17, 1989.

FOR FURTHER INFORMATION CONTACT: Dianne T. McRae, Center for Veterinary Medicine (HFV-135), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301-443-4913.

SUPPLEMENTARY INFORMATION: A. L. Laboratories, Inc., a subsidiary of A/S Apothekernes Laboratorium for Specialpraeparater, One Executive Dr., P.O. Box 1399, Fort Lee, NJ 07024, has filed a supplement to NADA 46-592 for bacitracin methylene disalicylate. The supplement provides for the use of Type A medicated articles containing 25, 40, and 50 grams of bacitracin methylene disalicylate per pound for making Type C medicated feeds for growing quail, for the prevention of ulcerative enteritis due to *Clostridium colinum* susceptible to bacitracin methylene disalicylate. The supplemental NADA is approved and 21 CFR 558.76(d)(1)(x) is amended to reflect 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 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 two environmental assessments, 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 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, 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: Sec. 512, 82 Stat. 343-351 (21 U.S.C. 360b); 21 CFR 5.10 and 5.83.

2. Section 558.76 is amended in the table in paragraph (d)(1)(x) by adding a new entry for "Quail" under the "Indications for use", "Limitations", and "Sponsor" columns to read as follows:

§ 558.76 Bacitracin methylene disalicylate.

* * * * *

(d) * * *

(1) * * *

Bacitracin methylene disalicylate in grams per ton	Combination in grams per ton	Indications for use	Limitations	Sponsor
(x) * * *		* * *		* * *
		Quail; for the prevention of ulcerative enteritis in growing quail due to *Clostridium colinum* susceptible to bacitracin methylene disalicylate	From Type A medicated articles containing 25, 40, or 50 grams of bacitracin methylene disalicylate. Feed continuously as the sole ration.	046573

* * * * *

Dated: February 9, 1989.

Richard H. Teske,

Deputy Director, Center for Veterinary Medicine.

[FR Doc. 89-3731 Filed 2-16-89; 8:45am]

BILLING CODE 4160-01-M

DEPARTMENT OF DEFENSE

Department of the Navy

32 CFR Part 706

Certifications and Exemptions Under the International Regulations for Preventing Collisions at Sea, 1972; Amendment

AGENCY: Department of the Navy, DOD.

ACTION: Final rule.

SUMMARY: The Department of the Navy is amending its certifications and exemptions under the International Regulations for Preventing Collisions at Sea, 1972 (72 COLREGS), to reflect that the Under Secretary of the Navy has (1) determined that USS ALBANY (SSN-753) is a vessel of the Navy which, due to its special construction and purpose, cannot comply fully with certain provisions of the 72 COLREGS without interfering with its special function as a naval submarine, and (2) has directed that certain corrections be made to the tables in the existing Part 706. The intended effect of this rule is to warn mariners in waters where 72 COLREGS apply.

EFFECTIVE DATE: February 7, 1989.

FOR FURTHER INFORMATION CONTACT: Captain P.C. Turner, JAGC, U.S. Navy, Admiralty Counsel, Office of the Judge Advocate General, Navy Department, 200 Stovall Street, Alexandria, VA 22332-2400. Telephone number: (202) 325-9744.

SUPPLEMENTARY INFORMATION: Pursuant to the authority granted in 33 U.S.C. 1605, the Department of the Navy amends 32 CFR Part 706. This amendment provides notice that the Under Secretary of the Navy, under authority delegated by the Secretary of the Navy, has certified that USS ALBANY (SSN-753) is a vessel of the Navy which, due to its special construction and purpose, cannot comply fully with 72 COLREGS: Rule