§ 212.83 **Allocation of refiner's increased product costs.**

* * * * *

(c) *Allocation of increased costs.* * * *

(2) *General formulae.*

* * * * *

V^n=The total volume of all covered products (other than propane, which may be included only to the extent that it was refined by the refiner from crude petroleum) and all products refined from crude petroleum other than covered products sold in the period "n" (the consecutive three-month period of the preceding year such that the middle month of the period corresponds to the current month "u").

V_i^n=The total volume of a specific covered product or products of the type "i" (other than propane, which may be included only to the extent that it was refined by the refiner from crude petroleum) sold in the period "n" (the consecutive three-month period of the preceding year such that the middle month of the period corresponds to the current month "u").

[FR Doc.74-19652 Filed 8-23-74;8:45 am]

Title 12—Banks and Banking

CHAPTER II—FEDERAL RESERVE SYSTEM

SUBCHAPTER A—BOARD OF GOVERNORS OF THE FEDERAL RESERVE SYSTEM

PART 265—RULES REGARDING DELEGATION OF AUTHORITY

In order to further effectuate the reorganization of the Board's staff announced on July 16, 1974, the Board has amended its Rules Regarding Delegations of Authority pursuant to section 12(k) of the Federal Reserve Act (12 U.S.C. 248(k)) to transfer to the Director of the Office of Saver and Consumer Affairs certain delegated functions previously delegated to the Director of the Division of Supervision and Regulation. The transferred delegations concern the Board's Securities Credit Regulations, 12 CFR Parts 207, 220, 221 and 224. The transferred delegations include those previously designated as 12 CFR 265.2 (c)(13) and (18), and a previously unpublished delegation.

Section 265.2 is amended as follows:

§ 265.2 [Amended]

1. Paragraphs (c)(13) and (c)(18) are repealed.

2. Prior paragraph (c)(14) through (17) are redesignated as paragraph (c)(13) through (16).

3. A new paragraph (h) is added to § 265.2 as follows:

(h) The Director of the Office of Saver and Consumer Affairs (or, in his absence, the Acting Director) is authorized:

(1) Under the provisions of §§ 207.2 (f), 220.2(e), and 221.3(d) of this chapter (Regulations G, T, and U, respectively) to approve issuance of the list of OTC margin stocks and to add, omit, or remove any stock in circumstances indicating that such change is necessary or appropriate in the public interest.

(2) Under the provisions of § 207.4 (a)(2)(ii) of this chapter (Regulation G) to approve repayments of the "deficiency" with respect to stock option or employee stock purchase plan credit in lower amounts and over longer periods of time than those specified in the regulation.

(3) Pursuant to the provisions of Part 261 of this chapter, to make available reports and other information of the Board acquired pursuant to Parts 207, 220, 221, and 224 of this chapter (Regulations G, T, U and X) of the nature and in circumstances described in § 261.6 (a)(2) and (3) of Part 261 of this chapter.

Effective date: August 16, 1974.

By order of the Board of Governors, August 16, 1974.

[SEAL] CHESTER B. FELDBERG,
Secretary of the Board.

[FR Doc.74-19678 Filed 8-23-74;8:45 am]

Title 21—Food and Drugs

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

SUBCHAPTER A—GENERAL

PART 2—ADMINISTRATIVE FUNCTIONS, PRACTICES, AND PROCEDURES

Subpart M—Organization

Revised Field Structure in Regions I, IV, and V

The Commissioner of Food and Drugs is amending "Part 2—Administrative Functions, Practices, and Procedures" (21 CFR Part 2) to provide revised field structure in Region I, Boston; Region IV, Atlanta; and Region V, Chicago. Recent reorganizations provide for the establishment of the Winchester Engineering and Analytical Center in Region I; District Offices at Atlanta, GA, Orlando, FL, and Nashville, TN in Region IV; and the Minneapolis Center for Microbiological Investigations in Region V. A Houston Section, under a pilot project, is operationally conducting district office-type functions in Region VI. This amendment incorporates those additional units in the field structure.

Therefore, pursuant to provisions of the Federal Food, Drug, and Cosmetic Act (sec. 701(a), 52 Stat. 1055; 21 U.S.C. 371(a)) and under authority delegated to the Commissioner (21 CFR 2.120), Part 2 is amended by revising § 2.175 to read as follows:

§ 2.175 **Field structure.**

REGION I

Regional Field Office: 585 Commercial Street, Boston, MA 02109.
Winchester Engineering and Analytical Center: 109 Holton Street, Winchester, MA 01890.

REGION II

Regional Field Office: 850 Third Avenue, Brooklyn, NY 11232.
District Office: 850 Third Avenue, Brooklyn, NY 11232.
District Office: 599 Delaware Avenue, Buffalo, NY 14202.
District Office: Room 831, 970 Broad Street, Newark, NJ 07102.
District Office: Post Office Box S-4427, San Juan Station, San Juan, PR 00905.

REGION III

Regional Field Office: Room 1204, Second and Chestnut Streets, Philadelphia, PA 19106.
District Office: Room 1204, Second and Chestnut Streets, Philadelphia, PA 19106.
District Office: 900 Madison Avenue, Baltimore, MD 21201.

REGION IV

Regional Field Office: 880 West Peachtree Street, Atlanta, GA 30309.
District Office: 880 West Peachtree Street, Atlanta, GA 30309.
District Office: 297 Plus Park Boulevard, Nashville, TN 37217.
District Office: Post Office Box 118, Orlando, FL 32802.

REGION V

Regional Field Office: Room A-1945, 175 West Jackson Boulevard, Chicago, IL 60607.
District Office: Room 1222, 433 West Van Buren Street, Chicago, IL 60607.
District Office: 1141 Central Parkway, Cincinnati, OH 45202.
District Office: 1560 East Jefferson Avenue, Detroit, MI 48207.
District Office: 240 Hennepin Avenue, Minneapolis, MN 55401.
Minneapolis Center for Microbiological Investigations: 240 Hennepin Avenue, Minneapolis, MN 55401.

REGION VI

Regional Field Office: 3032 Bryan Street, Dallas, TX 75204.
District Office: 3032 Bryan Street, Dallas, TX 75204.
District Office: Room 222, 423 Canal Street, New Orleans, LA 70130.
Houston Section: Room 413, 201 Fannin Street, Houston, TX 77002.

REGION VII

Regional Field Office: 1009 Cherry Street, Kansas City, MO 64106.

REGION VIII

Regional Field Office: 721 19th Street, U.S. Customhouse, Denver, CO 80202.

REGION IX

Regional Field Office: Room 518, 50 Fulton Street, San Francisco, CA 94102.
District Office: Room 518, 50 Fulton Street, San Francisco, CA 94102.
District Office: 1521 West Pico Boulevard, Los Angeles, CA 90015.

REGION X

Regional Field Office: Room 5003, 909 First Avenue, Seattle, WA 98104.

Effective date. This order shall be effective August 26, 1974.

(Sec. 701(a), 52 Stat. 1055; 21 U.S.C. 371(a).)

Dated: August 19, 1974.

SAM D. FINE,
Associate Commissioner for Compliance.

[FR Doc.74-19656 Filed 8-23-74;8:45 am]

SUBCHAPTER D—DRUGS FOR HUMAN USE

PART 310—NEW DRUGS

Subpart E—Requirements for Specific New Drugs or Devices

SUBCHAPTER G—COSMETICS

PART 700—GENERAL

Subpart B—Requirements for Specific Cosmetic Products

Vinyl Chloride as an Ingredient of Drug and Cosmetic Aerosol Products

In the FEDERAL REGISTER of April 22, 1974 (39 FR 14215), the Commissioner of Food and Drugs issued a notice of proposed rule making regarding all drug and cosmetic aerosol products containing vinyl chloride as an ingredient, including propellant. The proposal was based on the Commissioner's determination that there are sufficient scientific data on which to base a decision that: (1) Vinyl chloride presents an unnecessary hazard to the public health when it is used as an ingredient in cosmetic aerosol products, and that such use should be banned; and (2) vinyl chloride, when used as an ingredient in drug aerosol products, is not generally recognized as safe and effective, is a new drug within the meaning of section 201(p) of the Federal Food, Drug, and Cosmetic Act, and requires an approved new drug application as a condition of marketing. Interested persons were invited to submit comments regarding the proposal on or before May 22, 1974.

Comments were received from the American Academy of Pediatrics, a municipal consumer affairs unit, and an individual. All the comments were in support of the Commissioner's proposal. Accordingly, the Commissioner concludes that the regulations should be adopted as proposed.

The notice of proposed rule making also requested data regarding the use of polyvinyl chloride in containers for food and cosmetics, and in devices. The Commissioner urgently requested that certain data be submitted on the extent of the usage of polyvinyl chloride containers, the rates of extraction of vinyl chloride monomer from these containers, and other matters that will pertain to the safety of these containers. The time limit for submission of data requested in the proposal was on or before June 21, 1974. However, to date, the Commissioner has received only a few responses and again requests that the pertinent information be sent to the Hearing Clerk, Food and Drug Administration, Rm. 4–65, 5600 Fishers Lane, Rockville, MD 20852, as soon as possible.

A companion notice in the same issue of the FEDERAL REGISTER (39 FR 14238) required each registrant under section 510 of the Federal Food, Drug, and Cosmetic Act to submit a list of all human drugs which are being manufactured, prepared, propagated, compounded, or processed for commercial distribution and which contain vinyl chloride as an ingredient or are packaged in polyvinyl chloride containers. The notice also requested registrants to furnish certain additional information relating to vinyl chloride and polyvinyl chloride.

Data being received by the Food and Drug Administration as a result of the FEDERAL REGISTER notices are being compiled and reviewed to determine whether any additional action by the Commissioner is needed to protect the public health.

Therefore, pursuant to provisions of the Federal Food, Drug, and Cosmetic Act (secs. 502, 505, 601(a), 701(a),, 52 Stat. 1050–1055, as amended; 21 U.S.C. 352, 355, 361(a), 371(a)) and under authority delegated to the Commissioner (21 CFR 2.120), Parts 310 and 700 are amended as follows:

1. By adding a new § 310.506 to Subpart E to read as follows:

§ 310.506 Use of vinyl chloride as an ingredient, including propellant, of aerosol drug products.

(a) Vinyl chloride has been used as a propellant in aerosol drug preparations. Evidence indicates that vinyl chloride inhalation can result in acute toxicity manifested by dizziness, headache, disorientation, and unconsciousness where inhaled at high concentrations. Cardiac effects, bone changes, and degenerative changes in the brain, liver, and kidneys have been reported in animals. Studies also demonstrate carcinogenic effects in animals as a result of inhalation exposure to vinyl chloride. Recently, vinyl chloride has been linked to liver disease, including liver cancer, in workers engaged in the polymerization of vinyl chloride.

(b) The Commissioner finds that there is a lack of general recognition by qualified experts of the safety or effectiveness of aerosol drug preparations containing vinyl chloride as an ingredient, including propellant. Therefore, any such product containing vinyl chloride is a new drug and a new drug application approved under section 505 of the Federal Food, Drug, and Cosmetic Act is required for marketing.

(c) A completed and signed "Notice of Claimed Investigational Exemption for a New Drug" (Form FD–1571), as set forth in § 312.1 of this chapter, is required to cover clinical investigations designed to obtain evidence that such preparations are safe and effective for the purposes intended.

(d) Any such drug within the jurisdiction of the act which is not in accord with this regulation is subject to regulatory action.

2. By adding a new § 700.14 to Subpart B to read as follows:

§ 700.14 Use of vinyl chloride as an ingredient, including propellant of cosmetic aerosol products.

(a) Vinyl chloride has been used as an ingredient in cosmetic aerosol products including hair sprays. Where such aerosol products are used in the confines of a small room, as is often the case, the level of vinyl chloride to which the individual may be exposed could be significantly in excess of the safe level established in connection with occupational exposure. Evidence indicates that vinyl chloride inhalation can result in acute toxicity manifested by dizziness, headache, disorientation, and unconsciousness where inhaled at high concentrations. Studies also demonstrate carcinogenic effects in animals as a result of inhalation exposure to vinyl chloride. Furthermore, vinyl chloride has recently been linked to liver disease, including liver cancer, in workers engaged in the polymerization of vinyl chloride. It is the view of the Commissioner that vinyl chloride is a deleterious substance which may render any cosmetic aerosol product that contains it as an ingredient injurious to users. Accordingly, any cosmetic aerosol product containing vinyl chloride as an ingredient is deemed to be adulterated under section 601(a) of the Federal Food, Drug, and Cosmetic Act.

(b) Any cosmetic aerosol product containing vinyl chloride as an ingredient shipped within the jurisdiction of the act is subject to regulatory action.

Effective date. This order shall be effective September 25, 1974.

(Secs. 502, 505, 601(a), 701(a), 52 Stat. 1050–1055, as amended; 21 U.S.C. 352, 355, 361(a), 371(a).)

Dated: August 20, 1974.

SAM D. FINE,
Associate Commissioner for Compliance.

[FR Doc.74–19657 Filed 8–23–74;8:45 am]

CHAPTER II—DRUG ENFORCEMENT ADMINISTRATION, DEPARTMENT OF JUSTICE

PART 1308—SCHEDULES OF CONTROLLED SUBSTANCES

Exempt Chemical Preparations

The Administrator of the Drug Enforcement Administration has received applications pursuant § 1308.23 of Title 21 of the Code of Federal Regulations requesting that several chemical preparations containing controlled substances be granted the exemptions provided for in § 1308.24 of Title 21 of the Code of Federal Regulations.

The Administrator hereby finds that each of the following chemical preparations and mixtures is intended for laboratory, industrial, educational, or special research purposes, is not intended for general administration to a human being or other animal, and either (a) contains no narcotic controlled substances and is packaged in such a form or concentration that the package quantity does not present any significant potential for abuse, (b) contains either a narcotic or nonnarcotic controlled substance and one or more adulterating or denaturing agents in such a manner, combination, quantity, proportion or concentration, that the preparation or mixture does not present any potential for abuse, or (c) the formulation of such preparation or mixture incorporates methods of denaturing or other means so that the controlled substance cannot in practice be removed, and therefore

the preparation or mixture does not present any significant potential for abuse. The Administrator further finds that exemption of the following chemical preparations and mixtures is consistent with the public health and safety as well as the needs of researchers, chemical analysts, and suppliers of these products.

Therefore, pursuant to section 202(d) of the Comprehensive Drug Abuse Prevention and Control Act of 1970 (21 U.S.C. 812(d)), and under the authority vested in the Attorney General by sections 301 and 501(b) of the Act (21 U.S.C. 821 and 871(b)) and delegated to the Administrator of the Drug Enforcement Administration by § 0.100 of Title 28 of the Code of Federal Regulations (see 38 FR 18380, July 2, 1973) the Administrator hereby orders that Part 1308 of Title 21 of the Code of Federal Regulations be amended as follows:

a. By amending § 1308.24(i) by adding the following chemical preparations:

§ 1308.24 Exempt chemical preparations.

* * * * *

(i) * * *

Manufacturer or supplier	Product name and supplier's catalog No.	Form of product	Date of application
* *	* *	* *	*
Abbott Laboratories	Digoxin I 125 Imusay ® diagnostic kit No. 7649.	Kit: 100 units	June 6, 1974
* *	* *	* *	*
Amersham/Searle Corp	Lysergic acid di[1-^{14}C] ethylamide, catalog No. C.F.A. 534.	Ampoule: 0.6 mg to 8.1 mg.	July 2, 1974
* *	* *	* *	*
Flow Laboratories and Oxoid Ltd.	Barbitone acetate buffer for electrophoresis code No. BR 11g.	Bottle: 100 gms.	June 21, 1974
Do	Complement fixation test, diluent tablets code No. BR 16.	Bottle: 100 tablets.	Do.
* *	* *	* *	*
Hoffman-La Roche Inc	TM Abuscreen radioimmunoassay for Methadone.	Kit containing vials of: 5 ml, 30 ml, and 100 ml, and bottle: 500 ml.	June 17, 1974
Do	TM Abuscreen radioimmunoassay for Methaqualone.	do	Do.
* *	* *	* *	*
Hyland Division, Travenol Labs., Inc.	T-21	Vial: 20 ml	July 11, 1974
Do	T-22	Vial: 20 ml	Do.
Do	T-23	Vial: 50 ml	Do.
Do	T-24	Vial: 50 ml	Do.
Do	T-25	Vial: 50 ml	Do.
Do	T-27	Vial: 20 ml	July 15, 1974
Do	T-28	Vial: 50 ml	Do.
Do	T-29	Vial: 50 ml	Do.
Do	T-30	Vial: 50 ml	Do.
* *	* *	* *	*
J.W.S. Delavau Co., Inc. and the Theta Corp.	Test mixture TM No. 1	Vial: 2 ml	June 19, 1974
Do	Test mixture TM No. 2	do	Do.
Do	Test mixture SM No. 1	do	Do.
Do	Test mixture SP No. 1	do	Do.
Do	Test mixture SM No. 2	do	Do.
Do	Test mixture SP No. 2	do	Do.
Do	Test mixture SM No. 3	do	Do.
Do	Test mixture SP No. 3	do	Do.
Do	Test mixture SM No. 4	do	Do.
Do	Test mixture SP No. 4	do	Do.
* *	* *	* *	*
Millipore Corp	Counterimmuno electrophoresis buffer 1, catalog No. ESCE 032 01.	Vial: 10.19 gm	June 17, 1974
Do	CIEP AgaroSlide, catalog No. ESCE 032 01.	Plate 3¼"x2¼"x$\frac{1}{16}$"	July 11, 1974
* *	* *	* *	*
Warner-Lambert Co. (General Diagnostics Division).	Russell's viper venom reagent	Vial: 7.3 ml containing 48 mg of powder.	July 8, 1974
* *	* *	* *	*

* * * * * * *

Effective date. This order is effective August 26, 1974. Any interested person may file written comments on or objections to the order on or before September 25, 1974. If any such comments or objections raise significant issues regarding any finding of fact or conclusion of law upon which the order is based, the Administrator shall immediately suspend the effectiveness of the order until he may reconsider the application in light of the comments and objections filed. Thereafter, the Administrator shall reinstate, revoke or amend his original order as he determines appropriate.

Dated: August 12, 1974.

JOHN R. BARTELS, JR.,
Administrator,
Drug Enforcement Administration.

[FR Doc.74-19572 Filed 8-23-74;8:45 am]

Title 29—Labor

CHAPTER XIV—EQUAL EMPLOYMENT OPPORTUNITY COMMISSION

PART 1601—PROCEDURAL REGULATIONS

New 706 Agencies; Formal Designations

Pursuant to § 1601.12(g), Title 29, Chapter XIV of the Code of Federal Regulations as revised and published in the FEDERAL REGISTER, (37 9214–9220), May 6, 1972, the Equal Employment Opportunity Commission (hereinafter referred to as the Commission) designates the agencies listed below as 706 Agencies (§ 1601.12 (c)) and Provisional Notice Agencies (as defined in § 1601.12(d)(2)), for purposes of receiving charges deferred by the Commission pursuant to section 706 (c) and (d) of title VII of the Civil Rights Act of 1964, as amended, and for purposes of according weight to the final findings and orders of those agencies pursuant to § 1601.12(g)(1), and ends the 15-day period within which any person or organization may file written comments as provided for under § 1601.12(g)(1).

§ 1601.12 [Amended]

The Commission through publication of the following designated agencies hereby amend § 1601.12(k). The designated 706 agencys are:

Iowa Commission on Civil Rights
Wisconsin Department of Industry, Labor and Human Relations

JOHN H. POWELL, Jr.,
Chairman.

[FR Doc.74–19742 Filed 8–23–74;8:45 am]

PART 1602—RECORDS AND REPORTS

Extension of Deadline for Filing Apprenticeship Information Report EEO–2

Notice is hereby given that the 1974 deadline for filing Apprenticeship Information Report EEO–2 as required by 29 CFR 1602.15 is extended from September 30, 1974, to December 31, 1974. The period during which statistics for Report EEO–2 must be obtained remains unchanged. Signed at Washington, D.C. this 14th day of August 1974.

JOHN H. POWELL, Jr.,
Chairman, Equal Employment Opportunity Commission.

[FR Doc.74–19717 Filed 8–23–74;8:45 am]

PART 1602—RECORDS AND REPORTS

Extension of Deadline for Filing Local Union Equal Employment Opportunity Report EEO–3

Notice is hereby given that the 1974 deadline for filing Local Union Equal Employment Opportunity Report EEO–3 required by 29 CFR 1602.22 is extended from November 30, 1974, to December 31, 1974. The period during which statistics for Report EEO–3 must be obtained remains unchanged. Signed at Washington, D.C., this 14th day of August 1974.

JOHN H. POWELL, Jr.,
Chairman, Equal Employment Opportunity Commission.

[FR Doc.74–19718 Filed 8–23–74;8:45 am]

PART 1602—RECORDS AND REPORTS

Filing Date Changes for State and Local Government Information Report EEO–4 and Elementary; Secondary Staff Information Report EEO–5

After gaining some experience concerning the difficulty jurisdictions are having in compiling reports EEO–4 and EEO–5, the Equal Employment Opportunity Commission has determined that the filing dates should be extended from July 30, 1974, and October 15, 1974, respectively, to September 30, 1974 and November 30, 1974, respectively, and annually thereafter. Corresponding changes are required under the appropriate recordkeeping requirements. The payroll reporting period for each form remains unchanged. Accordingly, §§ 1602.30, 1602.32(a), 1602.39, and 1602.41(a) of Part 1602, Chapter XVI, of Title 29 of the Code of Federal Regulations are hereby amended as follows below. These amendments are effective immediately.

§ 1602.30 [Amended]

1. Section 1602.30 is amended by striking the date "July 30, 1973," and inserting therein the date "September 30, 1974."

§ 1602.32 [Amended]

2. Section 1602.32(a) is amended by striking the date "July 30, 1974," and inserting therein the date "September 30, 1974."

§ 1602.39 [Amended]

3. Section 1602.39 is amended by striking the date "October 15, 1973," and inserting therein the date "November 30, 1974."

§ 1602.41(a) [Amended]

4. Section 1602.41(a) is amended by striking the date "October 15, 1974," and inserting therein the date "November 30, 1974."

Signed at Washington, D.C., this 15th day of August, 1974.

JOHN H. POWELL, Jr.,
Chairman, Equal Employment Opportunity Commission.

[FR Doc.74–19719 Filed 8–23–74;8:45 am]

Title 31—Money and Finance

CHAPTER II—FISCAL SERVICE; DEPARTMENT OF THE TREASURY

SUBCHAPTER B—BUREAU OF GOVERNMENT FINANCIAL OPERATIONS

PART 202—DEPOSITARIES AND FINANCIAL AGENTS OF THE GOVERNMENT

Collateral Security

Section 202.6(c) and (d) of Department of the Treasury Circular 176, Second Revision (31 CFR Part 202), is hereby amended to read as follows:

§ 202.6 **Collateral security.**

* * * * *

(c) *Deposits of securities.* Unless the Secretary of the Treasury provides otherwise, collateral security under this part must be deposited with the Federal Reserve Bank or Branch of the district in which the depository is located (depositaries located in Puerto Rico, the Virgin Islands, and the Panama Canal Zone will be considered as being located in the New York Federal Reserve district), or with a custodian or custodians within the United States designated by the Federal Reserve Bank, under terms and conditions prescribed by the Federal Reserve Bank. Securities deposited with a Federal Reserve Bank must be accompanied by a letter stating specifically the purpose for which the securities are being deposited.

(d) *Assignment.* A depository that pledges securities which are not negotiable without its endorsement or assignment may, in lieu of placing its unqualified endorsement on each security, furnish an appropriate resolution and irrevocable power of attorney authorizing the Federal Reserve Bank to assign the securities. The resolution and power of attorney shall conform to such terms and conditions as the Federal Reserve Banks shall prescribe.

(Sec. 10, 56 Stat. 356, as amended (12 U.S.C. 265))

Notice and public procedures thereon are unnecessary as the fiscal policy of the United States is involved.

Dated: August 14, 1974.

[SEAL] DAVID MOSSO,
Deputy Fiscal Assistant Secretary.

[FR Doc.74–19715 Filed 8–23–74;8:45 am]

Title 36—Parks, Forests and Memorials

CHAPTER I—NATIONAL PARK SERVICE, DEPARTMENT OF THE INTERIOR

PART 7—SPECIAL REGULATIONS, AREAS OF THE NATIONAL PARK SERVICE

Natchez Trace Parkway, Tennessee-Alabama-Mississippi; Trucks, Trailers, Buses and Towed Vehicles; Possession of Beer and Alcoholic Beverages

A proposal was published at pages 20489 and 20490 of the FEDERAL REGISTER of June 11, 1974, to revise paragraphs (c) (1) and (4) of § 7.43 and to add paragraphs (c) (5), (6), (7), and (d) to Title 36 of the Code of Federal Regulations. The effect on the revisions and additions is to clarify the use of the Natchez Trace Parkway motor road by trucks, trailers, buses and towed vehicles, and to establish restrictions on the possession of beer and alcoholic beverages on the parkway.

Interested persons were given 30 days for submitting written comments, suggestions, or objections with respect to the proposed revisions and additions. As a result of comments received, the pro-

posed regulation is adopted with the following changes: Deletion of ". . . in accordance with applicable State and local Regulations," from paragraph (d)(1), and deletion of ". . . in accordance with applicable State and local laws," from paragraph (d)(2).

These revisions and additions will become effective September 25, 1974 (5 U.S.C. 553; 16 U.S.C. 3; * * *)

Paragraph (c) (1) and (4) of § 7.43 is revised, and paragraph (c) (5), (6), (7) and (d) are added, to read as follows:

§ 7.43 Natchez Trace Parkway.

* * * * *

(c) Vehicles—(1) *Trucks.* Trucks over one ton rated capacity are not permitted on the parkway. Trucks, not exceeding one ton rated capacity, are permitted to travel on the Natchez Trace Parkway when used solely for transportation of persons, their baggage, camping equipment and related articles for recreational purposes only. Trucks used for the purpose of hauling non-recreational materials are not permitted.

* * * * *

(4) *Recreational vehicles.* Recreational vehicles, including but not limited to self-propelled mobile homes, campers, housetrailers, and vehicles up to 1½ ton rated capacity, when such recreational vehicles are used solely to carry persons for recreational purposes together with their baggage, camping equipment, and related articles for vacation or recreational purposes, are permitted on the parkway.

(5) *Trailers.* Trailers are permitted when used non-commercially to transport baggage, camping equipment, horses for recreational riding, small boats and other similar items used for vacation or recreational purposes, provided they meet the following criteria:

(i) Utility type trailers must be enclosed or covered and are not to exceed 5 feet by 8 feet.

(ii) Trailers must be equipped with red taillights, red stoplights and mechanical turn signals. Clearance lights are required on trailers over 6 feet high.

(iii) Trailers carrying over one ton must be properly equipped with brakes, adequate to control the movement thereof and to stop and to hold such vehicle, and designed so as to be applied by the driver of the towing motor vehicle from its cab; said brakes shall be so designed and connected that in case of an accidental breakaway of the towed vehicle the brakes shall be automatically applied.

(iv) All trailers shall be coupled to the towing vehicle by means of a safety chain or equivalent device in addition to the regular trailer hitch or coupling. No more slack shall be left in safety chains than shall be necessary to permit proper turning and the safety chains shall be so connected to the towed and towing vehicles and to the drawbar to prevent the drawbar from dropping to the ground if the drawbar fails and shall be of sufficient strength to control the trailer in event of failure of the regular trailer hitch or coupling.

(v) Only one trailer of any type may be towed by any one vehicle along the parkway. The towing vehicle and trailer must not exceed 55 feet in length.

(6) Buses. Commercial passenger carrying buses, when used for touring purposes, may travel the Natchez Trace Parkway by obtaining special written permission in advance from the Superintendent or his representative. School buses may travel on the parkway without such written permission when transporting people for special recreational or educational purposes.

(7) Towed vehicles other than trailers. Such vehicles must be towed with a rigid tow bar which does not require a driver for the towed vehicle. Tow bar must be equipped with safety chains that are so connected to the towed and towing vehicles and to the tow bar that, if the tow bar fails, it will not drop to the ground and the chains shall be of sufficient strength to prevent breakaway of the towed vehicle in the event of such tow bar failure. The towed vehicle must be equipped with brakelights, taillights, and signal lights in accordance with applicable State regulations. The towing vehicle and towed vehicle must not exceed 55 feet in length.

(d) *Beer and alcoholic beverages.* (1) The possession of beer or any alcoholic beverage in an open or unsealed container, is prohibited except in designated picnic, lodging, residence, and camping areas.

(2) The possession of beer or any alcoholic beverage in an open or unsealed container in any motor vehicle is prohibited.

C. W. OGLE,
Superintendent, Natchez Trace.

[FR Doc.74-19650 Filed 8-23-74;8:45 am]

Title 40—Protection of Environment

CHAPTER I—ENVIRONMENTAL PROTECTION AGENCY

SUBCHAPTER C—AIR PROGRAMS

[FRL 245-3]

PART 52—APPROVAL AND PROMULGATION OF IMPLEMENTATION PLANS

Puerto Rico; Approval of Compliance Schedules

On May 31, 1972 (37 FR 10842), pursuant to section 110 of the Clean Air Act and 40 CFR Part 51, the Administrator of the Environmental Protection Agency approved the implementation plan submitted by the Commonwealth of Puerto Rico for attainment and maintenance of national ambient air quality standards. On April 5, April 9, April 17 and September 10, 1973, proposed revisions to the approved plan were submitted by Puerto Rico in the form of compliance schedules covering sources located in the Commonwealth. Set forth below are regulations approving forty-seven of these compliance schedules.

These compliance schedules were adopted by Puerto Rico after notice and public hearings in conformity with the requirements of 40 CFR 51.4 and 51.6, as certified by supporting documents submitted by the Puerto Rico Environmental Quality Board. The Administrator determined that these schedules could not be approved as originally submitted by the Commonwealth, and requested that additional information be supplied relating to the procedures employed in adoption of these schedules by the Commonwealth, and to the enforceability of their terms. This information was received by EPA on February 1 and February 12, 1974. The Administrator thereupon found these schedules to be approvable as consistent with the requirements of the Clean Air Act, 40 CFR Part 51, and Puerto Rico's approved control strategies. The Administrator therefore proposed approval of these schedules in the FEDERAL REGISTER of May 13, 1974 (39 FR 17109). The comments of interested persons were invited during the thirty-day period following publication of this proposed rulemaking. No comments were received, and the thirty-day period established in the FEDERAL REGISTER of May 13, 1974 has since expired. The Administrator therefore approves these schedules in the form of the regulations set out below.

The date indicated for final compliance in forty-six of these schedules will have passed by the time of their final approval. This fact is attributable to delays occasioned by the necessity of ministering to infirmities existing at the time of their initial submission. The Administrator believes approval of these schedules to be nonetheless desirable inasmuch as each schedule reflects the consensus of EPA and the Puerto Rico Environmental Quality Board as to the abatement actions to be taken, and the date by which compliance should be achieved.

Each of these schedules establishes a new date by which the source involved must comply with the applicable emission limitation provisions of the federally-approved state implementation plan. This date is indicated in the table below, under the heading "Final Compliance Date". In no case does this compliance date fall beyond the April 30, 1975 date established in Puerto Rico's implementation plan for attainment of primary ambient air quality standards, nor is relaxation of any emission standard involved. Achievement of ambient air quality standards by the attainment date is therefore not jeopardized by approval of these schedules.

Copies of the forty-seven compliance schedules hereby approved are available for public inspection at the offices of the Environmental Protection Agency, Region II, 26 Federal Plaza, New York, New York 10007, and the Puerto Rico Environmental Quality Board, 1550 Ponce de Leon Avenue, Santurce, Puerto Rico 00910.

The Administrator finds that good cause exists for making these regulations effective immediately upon publication. This will not compromise any interest of any of the sources affected, as they were involved in the development of these schedules, and are therefore aware of their terms. In addition, under section 307 of the Clean Air Act, as amended, an affected source is able to challenge the

approval of a schedule immediately. The immediate effective date will serve EPA interests in allowing the Agency to enforce immediately the substantive terms of these regulations.

(42 U.S.C. 1857c-5)

Dated: August 16, 1974.

JOHN QUARLES,
Acting Administrator.

Part 52 of Chapter I, Title 40 of the Code of Federal Regulations is amended as follows:

Subpart BBB—Puerto Rico

1. Section 52.2720 is amended by adding a new paragraph (c) as follows:

§ 52.2720 Identification of plan.

* * * * *

(c) Supplemental information was submitted by the Commonwealth of Puerto Rico Environmental Quality Board on April 5, April 9, April 17, May 30, June 18 and September 10, 1973, and on February 1 and February 12, 1974.

2. A new § 52.2724 is added as follows:

§ 52.2724 Compliance schedules.

(a) The compliance schedules for the sources indentified below are approved as meeting the requirements of § 51.15 of this chapter. All regulations cited are air pollution control regulations of the Commonwealth, unless otherwise noted.

PUERTO RICO

Source	Location	Regulations involved	Date of adoption	Effective date	Final compliance date
Hormigonera Oriental, Inc	Fajardo	5.2, 5.5	Apr. 4, 1973	Immediately	Apr. 30, 1973
Hormigonera Metropolitana, Inc	Caguas	5.2, 5.5	do	do	Do.
Hormigonera, Inc	Caguas	5.1, 5.2, 5.5	Apr. 5, 1973	do	Do.
Industria Alimenticia	Barceloneta	5.5, 8	do	do	Nov. 30, 1973
Jocarirck	Vega Baja	5.2, 5.5	do	do	Apr. 28, 1973
Inabon Aggregates, Inc	Coto Laurel	5.2	do	do	Mar. 31, 1973
Ponce Asphalt, Inc.—Santa Isabela Plant.	Ponce	5.2, 5.5	do	do	Apr. 15, 1973
Cantera Diaz, Inc	Trujillo Alto	5.2	do	do	June 15, 1973
Inabon Asphalt, Inc	Coto Laurel	5.1, 5.2	do	do	June 1, 1973
Betteroads Asphalt Corp., plant 2	Aguada	5.1, 5.2	do	do	May 31, 1973
Hormigonera, Inc., plant 6	Bayamon	5.2	Apr. 4, 1973	do	Mar. 30, 1973
Hormigonera, Inc., plant 15	Canovanas	5.2	Apr. 5, 1973	do	Apr. 30, 1973
Hormigonera, Inc., plant 7	Fajardo	5.2	do	do	Mar. 30, 1973
Hormigonera, Inc., plant 5 (temporary).	Dorado	5.2	do	do	Mar. 31, 1973
Hormigonera Oriental, Inc	Humancao	5.2	Apr. 4, 1973	do	Apr. 30, 1973
Betteroads Asphalt Corp., plant 6	Arecibo	5.2, 5.4	Apr. 5, 1973	do	Dec. 31, 1973
Empress O'Neill	Manati	5.2, 5.4	do	do	July 15, 1973
General Electric Switchgear	Vieques	5.2, 4.1	Apr. 4, 1973	do	Mar. 16, 1973
Terrozos Pico, Inc	Ponce	5.1, 5.2	Apr. 5, 1973	do	June 30, 1973
South Caribe Aggregates, Inc	Ponce	5.1, 5.2	do	do	June 20, 1973
Rexach Block—Santurce	Santurce	5.1, 5.5	Apr. 9, 1973	do	Aug. 15, 1973
Integrated Industries	Mayaguez	5.5	Apr. 8, 1973	do	Dec. 22, 1972
Amrex Construction Co., Inc	Cayey	5.2	Apr. 9, 1973	do	Mar. 15, 1974
Bloques La Plata	Toa Baja	5.2, 5.5	do	do	Apr. 30, 1973
Rodfall Muebles	Saint Just	5.2, 5.5	do	do	Aug. 31, 1973
South Caribe Aggregates, Inc	Juana Diaz	5.2, 5.5	do	do	June 30, 1973
Rexach Asphalt Corp	Aguadilla	5.2	do	do	May 15, 1973
Betteroads Asphalt Corp	Sabana Grande	5.2	do	do	Dec. 31, 1973
Ponce Asphalt	Ponce	5.2	Apr. 8, 1973	do	Apr. 15, 1973
Puerto Rico Aggregates	Carolina	5.2	Apr. 5, 1973	do	July 31, 1973
Ibec Packing Co	Mayaguez	5.5, 8	Apr. 8, 1973	do	Aug. 15, 1973
Rexach Concrete Corp	Gurubo	5.2	Apr. 9, 1973	do	May 1, 1973
Omark Caribbean, Inc	Bayamon	5.1, 5.5	Apr. 8, 1973	do	Nov. 30, 1973
Best Contracting Corp	Rio Piedras	5.2	Apr. 5, 1973	do	Mar. 22, 1973
Betteroads Asphalt Corp	Rio Piedras	5.2	do	do	Nov. 30, 1973
Hormigonera, Inc	Rio Piedras	5.1, 5.2, 5.5	do	do	Dec. 31, 1973
Agregados Gurabo, Inc	Gurubo	5.1, 5.2, 5.5	Apr. 8, 1973	do	Mar. 12, 1973
Rexach Aggregates Corp	Caguas	5.1, 5.2, 5.5	Apr. 9, 1973	do	Feb. 28, 1973
Quality Concre-Mix, Inc. (Sabana Seca).	Bayamon	5.1, 5.2, 5.5	do	do	June 15, 1973
Rexach Concrete Corp	Aguirre	5.1, 5.2, 5.5	do	do	Apr. 1, 1973
Eli Lilly	Mayaguez	7.3	Apr. 8, 1973	do	Dec. 31, 1974
Tito Castro Ready Mix	Ponce	5.1, 5.2, 5.5	July 23, 1973	do	Aug. 22, 1973
Puerto Rico Concrete Products	Rio Piedras	5.2	July 5, 1973	do	Aug. 30, 1973
Ponce Concrete Products	Ponce	5.2	do	do	Aug. 31, 1973
Motors and Parts Distributors	Hato Rey	5.5	do	do	July 29, 1973
Hormigonera, Inc	Rio Piedras	5.1, 5.2, 5.5	June 28, 1973	do	Dec. 31, 1973
Inabon Ready Mix	Coto Laurel	5.1, 5.2	do	do	Apr. 27, 1973

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[FRL 225-1]

PART 52—APPROVAL AND PROMULGATION OF IMPLEMENTATION PLANS

Iowa and Missouri; Approval and Disapproval of Compliance Schedules

On May 31, 1972 (37 FR 10842), pursuant to section 110 of the Clean Air Act and 40 CFR Part 51, the Administrator approved portions of State plans for implementation of the national ambient air quality standards.

The States of Iowa and Missouri submitted to the Environmental Protection Agency compliance schedules to be considered as proposed revisions to the approved plans pursuant to 40 CFR 51.6. The approvable schedules were adopted by the States and submitted to the Environmental Protection Agency for review after notice and public hearings. The public hearings were held in accordance with the procedural requirements of 40 CFR 51.15 pertaining to compliance schedules. The compliance schedules have been determined to be consistent with the approved control strategies of Iowa and Missouri.

Accordingly, the Administrator proposed approval of these schedules on November 27, 1973, in the FEDERAL REGISTER (38 FR 32583). The proposed approval and disapproval of these schedules published in the November 27, 1973, FEDERAL REGISTER provided for a 30-day comment period. No comments concerning these schedules were received. Set forth below are specific compliance schedules which the Administrator finds approvable. Pursuant to 40 CFR 51.8, the Administrator approves the schedules listed below.

Certain schedules published as proposed disapproval notices for Missouri were sent back to Missouri for correction of deficiencies. Subsequently, the state notified the Environmental Protection Agency that some sources were now in compliance and the schedules would not be formally resubmitted for approval. The withdrawn schedules are for the following sources in Missouri: Harbinson Walker Refractories Company, Vandalia; Gooch Mill and Elevator Company, North Kansas City; St. Joe Minerals Corporation, Herculaneum; City of Joplin, Joplin; Homer Galbraith, Savannah; Kansas City Power and Light Company, Montrose. If the deficiencies in the remaining seven disapproved schedules are corrected they will be reproposed at a later date.

Each approved revision establishes a new date by which the individual source must comply with the applicable emission limitation in the Federally approved State implementation plan. This date is indicated in the table below, under the heading "Final Compliance Date." In all cases, the schedules include incremental steps toward compliance with the applicable emission limitations. While the tables below do not include these interim dates, the actual compliance schedules do.

Under Iowa law, the compliance schedule is not enforceable after the date on which the associated variance expires and variances cannot extend for more than one year. Therefore, to the extent that the Iowa schedules extend past the variance expiration date, they are not legally enforceable at this time. For this reason, the Environmental Protection Agency's approval of each compliance schedule is unconditional only as to that part of the schedule covered by the initial variance. Approval of the remainder of the schedule will be conditioned upon the State's renewal of the variance in identical form and substance to that included in the schedule submitted to the Environmental Protection Agency and approved herein. If the variance is renewed in this manner, the condition precedent will be satisfied and the approval of the next segment of the schedule will not require further action by the State or this Agency. If the variance is not renewed, or is modified from the version that is Federally approved herein, the condition will not be fulfilled, the approval of the remainder of the schedule would not be effective, and the State's