Customs district of Laredo, Tex. (Region VI), was published in the FEDERAL REGISTER on September 2, 1971 (36 F.R. 17579). The proposal was based upon the need to provide better Customs service in the Laredo, Tex., district. No objections to the proposal were received.

Accordingly, by virtue of the authority vested in the President by section 1 of the Act of August 1, 1914, 38 Stat. 623, as amended (19 U.S.C. 2), which was delegated to the Secretary of the Treasury by the President by Executive Order No. 10289, September 17, 1951 (3 CFR Ch. II), and pursuant to authority provided by Treasury Department Order No. 190, Rev. 7 (34 F.R. 15846), Progreso, Tex., is hereby designated as a port of entry in the Laredo, Tex., district (Region VI).

The geographical limits of the port of Progreso, Tex., include that part of the county of Hidalgo, Tex., encompassed by the following boundaries:

On the north by 26°6' North latitude, on the east by 97°54' West longitude, on the south by the United States-Mexico international boundary, and on the west by 98°00' West longitude.

To reflect this change, the table in § 1.2(c) of the Customs Regulations is amended by inserting in the column headed "Ports of Entry" in the Laredo, Tex., district (Region VI) "Progreso, Tex. (T.D. 71-278)," directly below Hidalgo.

(80 Stat. 379, sec. 1, 37 Stat. 434, sec. 1, 38 Stat. 623, as amended, R.S. 251, as amended, sec. 624, 46 Stat. 759; 5 U.S.C. 301, 19 U.S.C. 1, 2, 66, 1624)

This Treasury Decision shall become effective 30 days after publication in the FEDERAL REGISTER.

[SEAL] EUGENE T. ROSSIDES,
Assistant Secretary of the Treasury.

OCTOBER 26, 1971.

[FR Doc.71-16283 Filed 11-5-71;8:50 am]

[T.D. 71-276]

PART 16—LIQUIDATION OF DUTIES

Sugar Content of Certain Articles From Australia Subject to Countervailing Duties

Net amount of bounty declared for the period October 1970 through September 1971 for products of Australia subject to the countervailing duty order published in T.D. 54582. Section 16.24(f), Customs Regulations, amended.

The Treasury Department is in receipt of official information that the rates of bounties or grants paid or bestowed by the Australian Government within the meaning of section 303, Tariff Act of 1930 (19 U.S.C. 1303), on the exportation during the period October 1970 through September 1971 of approved fruit products and other approved products containing sugar are the amounts set forth in the following table:

MERCHANDISE—APPROVED FRUIT PRODUCTS AND OTHER APPROVED PRODUCTS

Month	*Net amount of bounty per 2,240 lbs. of sugar content (Australian dollars)*
October 1970	73.30
November 1970	72.40
December 1970	70.70
January 1971	70.40
February 1971	54.80
March 1971	56.40
April 1971	67.90
May 1971	65.20
June 1971	66.10
July 1971	67.40
August 1971	69.00
September 1971	65.70

The net amount of bounties or grants on the above-described commodities which are manufactured or produced in Australia is hereby ascertained, determined, and declared to be the rate stated in the above table. Additional duties on the above-described commodities, except those commodities covered by T.D. 55716 (27 F.R. 9595), whether imported directly or indirectly from that country, equal to the net amounts of the bounty shown above shall be assessed and collected.

The table in § 16.24(f) under "Australia—Sugar content of certain articles" is amended (1) by deleting therefrom the reference to T.D. 70-105, and (2) by adding a reference to this Treasury Decision. As amended the last three lines of the table under this commodity will read:

Country	Commodity	Treasury decision	Action
		70-197	New rate.
		70-225	New rate.
		71-273	New rate.

(R.S. 251, secs. 303, 624, 46 Stat. 687, 759; 19 U.S.C. 66, 1303, 1624)

[SEAL] LEONARD LEHMAN,
Acting Commissioner of Customs.

Approved: October 21, 1971.

EUGENE T. ROSSIDES,
Assistant Secretary of the Treasury.

[FR Doc.71-16284 Filed 11-5-71;8:50 am]

Title 21—FOOD AND DRUGS

Chapter II—Bureau of Narcotics and Dangerous Drugs, Department of Justice

PART 308—SCHEDULES OF CONTROLLED SUBSTANCES

Removal of Exceptions From Amphetamine and Methamphetamine Combination Products

A notice was published in the FEDERAL REGISTER of September 21, 1971 (36 F.R. 17849), proposing the removal of exceptions from amphetamine and methamphetamine combination products. These exceptions, from application of certain provisions of the Controlled Substances Act (21 U.S.C. 801 et seq.) and the Controlled Substances Import and Export Act (21 U.S.C. 951 et seq.), were granted under the Drug Abuse Control Amendments of 1965 and were continued under the Comprehensive Drug Abuse Prevention and Control Act of 1970 (Public Law 91-513) for administrative purposes.

All interested persons were given 30 days after publication to submit their objections to and comments on the proposal. No objections were received. The Narcotic and Drug Control Division of the State Board of Health of South Carolina, commented specifically on the product Edrisal, noting that the formula of the drug does not preclude a potential for abuse, that similar drugs (Edrisal with Codeine and Daprisal) are listed in Schedule II, and that as the stringent controls of Schedule II are applied to other stimulants, abuse of Edrisal is likely to increase.

After careful consideration of the comments submitted, in view of the fact that no objections were received, and based upon the investigations of the Bureau of Narcotics and Dangerous Drugs, the Director finds that no compound, mixture, or preparation containing any quantity of amphetamine (or its salts, optical isomers, or salts of its optical isomers) or methamphetamine (or its salts, isomers, or salts of its isomers) and one or more active medicinal ingredients not having a stimulant effect on the central nervous system, contains such ingredients in such combinations, quantity, proportion, or concentration as to vitiate the potential for abuse of the amphetamine or methamphetamine substances.

Therefore, under the authority vested in the Attorney General by section 202 (d) of the Comprehensive Drug Abuse Prevention and Control Act of 1970 (21 U.S.C. 812(d)), and redelegated to the Director, Bureau of Narcotics and Dangerous Drugs by § 0.100 of Title 28 of the Code of Federal Regulations, the Director hereby orders that:

1. Section 308.13(b)(1) of Title 21 of the Code of Federal Regulations be deleted and replaced with a new paragraph to read:

§ 308.13 Schedule III.

* * * * * *

(b) * * *

(1) Those compounds, mixtures, or preparations in dosage unit form containing any stimulant substances which compounds, mixtures, or preparations were listed on August 25, 1971, as excepted compounds under § 308.32, and any other drug of the quantitative composition shown in that list for those drugs or which is the same except that it contains a lesser quantity of controlled substances.

* * * * *

§ 308.32 [Amended]

2. Section 308.32(b) of Title 21 of the Code of Federal Regulations be amended by deleting the following drugs:

Trade name or other designation	Composition	Manufacturer or supplier
Edrisal	Tablet: Amphetamine sulfate 2.5 mg.; asprin, 162 mg.; phenacetin 162 mg.	Smith Kline & French Laboratories.
• • •		
Genegesic Capsules	Capsule: Methamphetamine hydrochloride, 1.2 mg.; chlorpheniramine maleate, 3.8 mg.; phenacetin, 120.0 mg.; salicylamide, 180.0 mg.; caffeine, 30.0 mg.; ascorbic acid, 50.0 mg.	General Pharmaceutical Products Inc.
• • •		
Hovizyme	Tablet: Methamphetamine hydrochloride, 0.5 mg.; conjugated estrogens-equine, 0.125 mg.; methyl testosterone, 1.25 mg.; amylase, 10.0 mg.; protease, 5.0 mg.; cellulase, 2.0 mg.; nicotinyl alcohol tartrate, 7.5 mg.; dehydrocholic acid, 50.0 mg.; ascorbic acid, 50.0 mg.; ferrous fumarate, 6.0 mg.	Ayerst Laboratories.
• • •		
Mediatric	Tablet of capsule: Methamphetamine hydrochloride, 1 mg.; conjugated estrogens-equine, 0.25 mg.; methyl testosterone, 2.5 mg.	Do.
Mediatric Liquid	Solution (15 cc.): Methamphetamine hydrochloride, 1 mg.; conjugated estrogens-equine, 0.25 mg.; methyl testosterone, 2.5 mg.	Do.
• • •		
Special Formula 711	Tablet: d-Amphetamine sulfate, 2.5 mg.; mephenesin, 500 mg.; salicylamine, 300 mg.	Detroit First Aid Co.
• • •		
Thora-Dex No. 1	Tablet: Dextroamphetamine sulfate, 2 mg., chlorpromazine hydrochloride, 10 mg.	Smith Kline & French Laboratories.
Thora-Dex No. 2	Tablet: Dextroamphetamine sulfate, 5 mg., chlorpromazine hydrochloride, 25 mg.	Do.
• • •		

In his order published July 7, 1971, in the FEDERAL REGISTER (36 F.R. 12736), the Director recognized that certain combination products containing amphetamine or methamphetamine exempted under the Drug Abuse Control Amendments of 1965 were not expressly excepted under § 308.32. The Director stated that, as a matter of policy, those substances would be deemed excepted under § 308.32 pending further action by the Bureau. This order applies to those substances as well.

The effect of this order is to subject all compounds, mixtures, or preparations containing amphetamine or methamphetamine, except those now listed in Schedule II, to all of the requirements of sections 305, 307, and 309 of the Controlled Substances Act (relating to labeling, recordkeeping, and prescription requirements for controlled substances), sections 1002, 1003, and 1004 of the Controlled Substances Import and Export Act (relating to importation, exportation, transshipment) and § 301.74(d) of Title 21 of the Code of Federal Regulations (relating to sampling of controlled substances).

This order does not transfer any formerly excepted compounds, mixtures, or preparations containing amphetamine or methamphetamine from Schedule III to Schedule II.

The requirements imposed upon the formerly excepted amphetamine and methamphetamine combination products by virtue of the removal of the excepted status shall become effective as follows:

1. *Labeling.* All labels on commercial containers of, and all labeling of, the above formerly excepted stimulant compounds, which are packaged on and after May 1, 1972, shall comply with the requirements of 21 CFR Part 302.

2. *Records and inventories.* Every registrant who is required to keep records under § 304.03 of Title 21 of the Code of Federal Regulations, and who is manufacturing, distributing or dispensing any of the above formerly excepted stimulant compounds, shall take an inventory of all stocks on hand on January 3, 1972, and thereafter shall keep all required records regarding these compounds.

3. *Prescriptions.* All prescriptions for the above formerly excepted stimulant compounds shall comply with 21 CFR Part 306, as applied to substances listed in Schedule III, on and after January 1, 1972.

4. *Importation and exportation.* All importation, exportation, transshipment, and in-transit shipment of the above formerly excepted stimulant compounds shall comply with the requirements of 21 CFR Part 312 on and after January 1, 1972.

5. *Security.* All of the above formerly excepted stimulant compounds shall be manufactured, stored, distributed and shipped in compliance with §§ 301.71–76 of Title 21 of the Code of Federal Regulations on and after January 1, 1972.

This order is effective on the date of its publication in the FEDERAL REGISTER (11–6–71).

Dated: November 1, 1971.

JOHN E. INGERSOLL,
Director, Bureau of Narcotics and Dangerous Drugs.

[FR Doc.71–16208 Filed 11–5–71;8:45 am]

Title 29—LABOR

Subtitle A—Office of the Secretary of Labor

PART 55—GRANTS UNDER THE EMERGENCY EMPLOYMENT ACT OF 1971

Subpart D—Grants Under the Secretary's Discretionary Authority

Subtitle A of Title 29, Code of Federal Regulations, is hereby amended by adding thereto a new subpart to Part 55, designated Subpart D, relating to grants under the Secretary's discretionary authority. The new Subpart D sets forth the regulations of the Secretary of Labor for making grants under section 9(a)(2) of the Emergency Employment Act of 1971 (Public Law 92–54).

The Emergency Employment Act of 1971 was designed to increase employment and was made effective by Congress on an emergency basis. The effective implementation of the discretionary grant program is not possible without regulations to enable the intended recipients of Federal financial assistance to know the requirements. Compliance with the notice and public procedure requirements of 5 U.S.C. 553 would involve a delay in making available the assistance provided by this Act; we find that under the circumstances such delay would be impracticable and contrary to the public interest. Accordingly, the new Subpart D shall be effective upon publication in the FEDERAL REGISTER (11–6–71).

It is the policy of this Department that rules relating to public property, loans, grants, benefits, or contracts shall be published for comment notwithstanding the provisions of 5 U.S.C. 553. See 29 CFR 2.7 published in the July 10, 1971 FEDERAL REGISTER, 36 F.R. 12976. In accordance with the spirit of the public policy set forth in the above mentioned section, interested parties may submit written comments, suggestions, data, or arguments to the Assistant Secretary for Manpower, U.S. Department of Labor, Washington, D.C. 20210, within 45 days of the publication of the regulations contained in this part. Material thus submitted will be evaluated and acted upon in the same manner as if this document were a proposal. Until it is revised, however, it shall remain effective, thus permitting the public business to proceed expeditiously.

Part 55 is hereby amended as follows:

1. A new Subpart D is hereby added as follows:

Subpart D—Grants Under the Secretary's Discretionary Authority