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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

CONTROLLED SUBSTANCES IN SCHEDULES I AND II

1975 Final Aggregate Production Quotas

Section 306 of the Comprehensive Drug Abuse Prevention and Control Act of 1970 (21 U.S.C. 826) requires the Attorney General to establish aggregate production quotas for all controlled substances in Schedules I and II each year. This responsibility has been delegated to the Administrator of the Drug Enforcement Administration pursuant to § 0.100 of Title 28 of the Code of Federal Regulations.

On December 13, 1974, a notice of the proposed aggregate production quotas for these substances was published in the FEDERAL REGISTER (39 FR 241). All interested parties were invited to comment on or object to the proposed aggregate production quotas on or before January 13, 1975. Except with reference to Methylphenidate, no requests for hearings have been received by the Administration relative to the proposed quotas. Comments and objections relative to the proposed aggregate production quotas have been received from Western Fher Laboratories relative to Phenmetrazine, from Hoffman-La Roche Inc. relative to Alphaprodine, from Eli Lilly and Company relative to Amobarbital and Secobarbital, and from Richardson-Merrell Inc. relative to that portion of the Desoxyephedrine quota allocated for the production of Levo-Desoxyephedrine for use in the manufacture of a non-controlled substance. Specific details relative to these comments will be outlined in a future FEDERAL REGISTER notice.

Due to the fact that a request for a hearing has been received by the Administration with reference to the proposed aggregate production quota for Methylphenidate, the aggregate production quota for Methylphenidate does not appear in this order.

Based upon consideration of the factors set forth in 39 FR 241, the Administrator of the Drug Enforcement Administration, under the authority vested in the Attorney General by section 306 of the Comprehensive Drug Abuse Prevention and Control Act of 1970 (21 U.S.C. 826) and delegated to the Administrator by § 0.100 of Title 28 of the Code of Federal Regulations orders that the aggregate production quotas for 1975 for controlled substances, expressed in grams in terms of their respective anhydrous bases, be established as follows:

SCHEDULE I

Basic class:	Granted (1975)
1-alpha-acetylmethadole	300,000
Tetrahydrocannabinols	500

SCHEDULE II

Alphaprodine	34,500
Amobarbital	12,504,986
Amphetamine	3,291,300
Anileridine	88,133
Apomorphine	2,000
Cocaine	600,000
Codeine (for sale)	46,273,000
Codeine (for conversion)	1,165,000
Desoxyephedrine	[1] 1,215,374
Dihydrocodeine	731,000
Diphenoxylate	1,133,000
Feogine	200,000
Ethylmorphine	44,680
Fentanyl	2,000
Hydrocodone	800,000
Hydromorphone	70,200
Levorphanol	3,000
Methadone	3,245,000
Methadone Intermediate (4-cyano-2 dimethyl-amino-4, 4-diphenyl butane)	1,950,000
Methaqualone	19,628,335
Mixed Alkaloids of Opium	163,321
Morphine (for sale)	800,000
Morphine (for conversion)	62,182,000
Norpethidine	830,000
Opium (tinctures, extracts, etc. expressed in terms of opium)	1,904,000
Oxycodone (for sale)	1,769,400
Oxycodone (for conversion)	7,000
Oxymorphone	3,000
Pentobarbital	28,119,000
Pethidine	17,057,410
Phenazocine	225
Phenmetrazine	2,741,700
Secobarbital	18,430,000
Thebaine (for sale)	4,250,000
Thebaine (for conversion)	1,750,000

[1] 758,108 grams for the production of Levodesoxyephedrine for use in a non-controlled product, and 457,266 for production of Methamphetamine).

Pursuant to Title 21 Code of Federal Regulations, § 1303.23(c) the Administrator of the Drug Enforcement Administration will in early 1975 adjust individual manufacturing quotas allocated for 1975 based upon 1974 end of year inventory figures submitted by applicants and estimates of medical and scientific requirements to be provided by the Food and Drug Administration.

All persons who submitted an application for either an individual manufacturing quota or procurement quota for 1975 will be notified by mail as to their respective 1975 quota established by the Drug Enforcement Administration.

This order is effective on January 20, 1975.

Dated: January 15, 1975.

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

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