

Drug	Schedule
Diphenoxylate (9170)	II

The firm plans to manufacture the listed controlled substances to make finished dosage forms for distribution to its customers.

Any other such applicant and any person who is presently registered with DEA to manufacture such substances may file comments or objections to the issuance of the proposed registration.

Any such comments or objections may be addressed, in quintuplicate, to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, United States Department of Justice, Washington, D.C. 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than February 28, 2000.

Dated: December 16, 1999.

John H. King,

Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

[FR Doc. 99-33650 Filed 12-27-99; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Application

Pursuant to Section 1301.33(a) of Title 21 of the Code of Federal Regulations (CFR), this is notice that on October 15, 1999, Polaroid Corporation, 1265 Main Street, Building W6, Waltham, Massachusetts 02451, made application to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of 2, 5-dimethoxyamphetamine (7396), a basic class of controlled substance listed in Schedule I.

The firm plans to manufacture bulk 2, 5-dimethoxyamphetamine for conversion into a non-controlled substance.

Any other such applicant and any person who is presently registered with DEA to manufacture such substance may file comments or objections to the issuance of the proposed registration.

Any such comments or objections may be addressed, in quintuplicate, to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, United States Department of Justice, Washington, D.C. 20537, Attention: DEA Federal Register Representative (CCR),

and must be filed no later than February 28, 2000.

Dated: December 13, 1999.

John H. King,

Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

[FR Doc. 99-33647 Filed 12-27-99; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated December 14, 1998, and published in the **Federal Register** on December 23, 1998, (63 FR 71160), Pressure Chemical Company, 3419 Spellman Street, Pittsburgh, Pennsylvania 15201, made application to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of 2,5-dimethoxyamphetamine (7396), a basic class of controlled substance listed in Schedule I.

The firm plans to bulk manufacture 2,5-dimethoxyamphetamine for distribution to its customers.

DEA has considered the factors in Title 21, United States Code, Section 823(a) and determined that the registration of Pressure Chemical Company to manufacture 2,5-dimethoxyamphetamine is consistent with the public interest at this time. DEA has investigated the company to ensure that the company's continued registration is consistent with the public interest. The investigation included inspection and testing of the company's physical security systems, verification of the company's compliance with state and local laws, and a review of the company's background. Therefore, pursuant to 21 U.S.C. 823 and 28 C.F.R. §§ 0.100 and 0.104, the Deputy Assistant Administrator, Office of Diversion Control, hereby orders that the application submitted by the above firm for registration as a bulk manufacturer of the basic class of controlled substance listed above is granted.

Dated: December 17, 1999.

John H. King,

Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

[FR Doc. 99-33646 Filed 12-27-99; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[DEA No. 1861]

Controlled Substances: Established Initial Aggregate Production Quotas for 2000

AGENCY: Drug Enforcement Administration (DEA), Justice.

ACTION: Notice of aggregate production quotas for 2000.

SUMMARY: This notice establishes initial 2000 aggregate production quotas for controlled substances in Schedules I and II of the Controlled Substances Act (CSA).

EFFECTIVE DATE: December 28, 1999.

FOR FURTHER INFORMATION CONTACT:

Frank L. Sapienza, Chief, Drug and Chemical Evaluation Section, Drug Enforcement Administration, Washington, DC 20537, Telephone: (202) 307-7183.

SUPPLEMENTARY INFORMATION: Section 306 of the CSA (21 U.S.C. 826) requires that the Attorney General establish aggregate production quotas for each basic class of controlled substance listed in Schedules I and II. This responsibility has been delegated to the Administrator of the DEA by § 0.100 of Title 28 of the Code of Federal Regulations. The Administrator, in turn, has redelegated this function to the Deputy Administrator, pursuant to § 0.104 of Title 28 of the Code of Federal Regulations.

The 2000 aggregate production quotas represent those quantities of controlled substances that may be produced in the United States in 2000 to provide adequate supplies of each substance for: the estimated medical, scientific, research and industrial needs of the United States; lawful export requirements; and the establishment and maintenance of reserve stocks (21 U.S.C. 826(a) and 21 CFR 1303.11). These quotas do not include imports of controlled substances for use in industrial processes.

On October 21, 1999, a notice of the proposed initial 2000 aggregate production quotas for certain controlled substances in Schedules I and II was published in the **Federal Register** (64 FR 56809). All interested persons were invited to comment on or object to these proposed aggregate production quotas on or before November 22, 1999.

Six companies commented on a total of 16 Schedules I and II controlled substances within the published comment period. The companies commented that the proposed aggregate