

Drug	Schedule
Alpha-Methylthiofentanyl (9832)	I
3-Methylthiofentanyl (9833)	I
Thiofentanyl (9835)	I
Amphetamine (1100)	II
Methamphetamine (1105)	II
Phenmetrazine (1631)	II
Methylphenidate (1724)	II
Amobarbital (2125)	II
Pentobarbital (2270)	II
Secobarbital (2315)	II
Glutethimide (2550)	II
Nabilone (7379)	II
1-Phenylcyclohexylamine (7460)	II
Phencyclidine (7471)	II
1-Piperidinocyclohexanecarbonitrile (8603)	II
Alphaprodine (9010)	II
Cocaine (9041)	II
Codeine (9050)	II
Dihydrocodeine (9120)	II
Oxycodone (9143)	II
Hydromorphone (9150)	II
Diphenoxylate (9170)	II
Benzoyllecgonine (9180)	II
Ethylmorphine (9190)	II
Hydrocodone (9193)	II
Levomethorphan (9210)	II
Levorphanol (9220)	II
Isomethadone (9226)	II
Meperidine (9230)	II
Methadone (9250)	II
Methadone-intermediate (9254)	II
Morphine (9300)	II
Thebaine (9333)	II
Levo-alphaacetylmethadol (9648)	II
Oxymorphone (9652)	II
Noroxymorphone (9668)	II
Alfentanil (9737)	II
Sufentanil (9740)	II
Fentanyl (9801)	II

The firm plans to manufacture small quantities of the listed controlled substances to make deuterated and non-deuterated drug reference standards which will be distributed to analytical and forensic laboratories for drug testing programs.

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, DC 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than April 16, 2001.

Dated: January 25, 2001.

Laura M. Nagel,

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

[FR Doc. 01-3752 Filed 2-13-01; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated September 25, 2000, and published in the **Federal Register** on October 3, 2000, (65 FR 59018), Chattem Chemicals, Inc., 3801 St. Elmo Avenue, Building 18, Chattanooga, Tennessee 37409, made application by letter to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the basic classes of controlled substances listed below:

Drug	Schedule
N-Ethylamphetamine (1475)	I

Drug	Schedule
4-Methoxyamphetamine (7411)	I
2,5-Dimethoxyamphetamine (7396)	I
Difenoxin (9168)	I
Methylphenidate (1724)	II
Pentobarbital (2270)	II
Secobarbital (2315)	II
Diphenoxylate (9170)	II
Hydrocodone (9193)	II

The firm plans to bulk manufacture the listed controlled substances to produce products for distribution to its customers.

No comments or objections have been received. DEA has considered the factors in Title 21, United States Code, Section 823(a) and determined that the registration of Chattem Chemicals, Inc. to manufacture the listed controlled substances is consistent with the public interest at this time. DEA has investigated Chattem Chemicals, Inc. to ensure that the company's registration is consistent with the public interest. This 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 and history. Therefore, pursuant to 21 U.S.C. 823 and 28 CFR 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 classes of controlled substances listed above is granted.

Dated: January 25, 2001.

Laura M. Nagel,

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

[FR Doc. 01-3750 Filed 2-13-01; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Registration

By Notice dated October 31, 2000, and published in the **Federal Register** on November 14, 2000, (65 FR 68158), Stepan Company, Natural Products Department, 100 W. Hunter Avenue, Maywood, New Jersey 07605, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as an importer of coca leaves (9040), a basic class of controlled substance listed in Schedule II.

The firm plans to import the coca leaves to manufacture bulk controlled substance.

No comments or objections have been received. DEA has considered the factors in Title 21, United States Code, Section 823(a) and determined that the registration of Stepan Company, Natural Products Department to import coca leaves is consistent with the public interest and with United States obligations under international treaties, conventions, or protocols in effect on May 1, 1971, at this time. DEA has investigated Stepan Company, Natural Products Department on a regular basis to ensure that the company's continued registration is consistent with the public interest. This investigation included inspection and testing of the company's physical security systems, audits of the company's records, verification of the company's compliance with state and local laws, and a review of the company's background and history. Therefore, pursuant to section 1008(a) of the Controlled Substances Import and Export Act and in accordance with Title 21, Code of Federal Regulations, section 1301.34, the above firm is granted

registration as an importer of the basic class of controlled substance listed above.

Dated: January 25, 2001.

Laura M. Nagel,

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

[FR Doc. 01-3751 Filed 2-13-01; 8:45 am]

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DEPARTMENT OF LABOR

Pension and Welfare Benefits Administration

Extension of Information Collection; Comment Request; Prohibited Transaction Exemption 91-55

ACTION: Notice.

SUMMARY: The Department of Labor, as part of its continuing effort to reduce paperwork and respondent burden, conducts a preclearance consultation program to provide the general public and Federal agencies with an opportunity to comment on proposed and continuing collections of information in accordance with the Paperwork Reduction Act of 1995 (PRA 95) 44 U.S.C. 3506(c)(2)(A). This helps to ensure that requested data can be provided in the desired format, reporting burden (time and financial resources) is minimized, collection instruments are clearly understood, and the impact of collection requirements on respondents can be properly assessed.

Currently, the Pension and Welfare Benefits Administration is soliciting comments concerning the information collection request (ICR) incorporated in Prohibited Transaction Exemption 91-55, Transactions Between Individual Retirement Accounts and Authorized Purchasers of American Eagle Coins. A copy of the ICR may be obtained by contacting the office listed in the Addresses section of this notice.

DATES: Written comments must be submitted to the office shown in the Addresses section below on or before April 16, 2001.

ADDRESSES: Gerald B. Lindrew, Office of Policy and Research, U.S. Department of Labor, Pension and Welfare Benefits Administration, 200 Constitution Avenue, NW, Washington, D.C. 20210. Telephone: (202) 219-4782 (this is not a toll-free number); FAX: (202) 219-4745. These are not toll-free numbers.

SUPPLEMENTARY INFORMATION:

I. Background

Prohibited Transaction Exemption 91-55 permits purchases and sales by

certain "individual retirement accounts," as defined in Internal Revenue Code section 408 (IRAs) of American Eagle bullion coins ("Coins") in principal transactions from or to broker-dealers in Coins that are "authorized purchasers" of Coins in bulk quantities from the United States Mint and which are also "disqualified persons," within the meaning of Code section 4975(e)(2), with respect to IRAs. The exemption also describes the circumstances under which an interest-free extension of credit in connection with such sales and purchases is permitted. In the absence of an exemption, such purchases and sales and extensions of credit would be impermissible under the Employee Retirement Income Security Act of 1974 (ERISA).

The information collection request for this exemption includes three requirements. First, certain information related to covered transactions in Coins must be disclosed by the authorized purchaser to persons who direct the transaction for the IRA. Currently, it is standard industry practice that most of this information is provided to persons directing investments in an IRA when transactions in Coins occur. The exemption also requires that the disqualified person maintain for a period of at least six years such records as are necessary to allow accredited persons, as defined in the exemption, to determine whether the conditions of the transaction have been met. Finally, an authorized purchaser must provide a confirmation statement with respect to each covered transaction to the person who directs the transaction for the IRA.

The recordkeeping requirement facilitates the Department's ability to make findings under section 408 of ERISA and section 4975(c) of the Code. The confirmation and disclosure requirements protect a participant or beneficiary investing in IRAs transacting in Coins with authorized purchasers by providing the investor or the person directing his or her investments with timely information about the market in Coins and about the individual's account in particular.

II. Desired Focus of Comments

The Department of Labor is particularly interested in comments that:

- Evaluate whether the proposed collection of information is necessary for the proper performance of the