

information should address one or more of the following four points:

(1) Evaluate whether the proposed collection of information is necessary for the proper performance of the function of the agency, including whether the information will have practical utility;

(2) Evaluate the accuracy of the agency's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;

(3) Enhance the quality, utility, and clarity of the information to be collected; and

(4) Minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, e.g., permitting electronic submission of responses.

Overview of This Information Collection

(1) *Type of information collection:* Extension of a currently approved collection.

(2) *The title of the form/collection:* ARCOS Transaction Reporting—DEA Form 333.

The agency form number, if any, and the applicable component of the Department sponsoring the collection: Form No.: DEA-333. Office of Diversion Control, Drug Enforcement Administration, U.S. Department of Justice.

(4) *Affected public who will be asked or required to respond, as well as a brief abstract:* Primary: Business or other for-profit. Other: None.

Necessary for U.S. to meet obligations under two international treaties: Single Convention on Narcotic Drugs and Psychotropic Substances. Treaties require information on the manufacture and consumption of certain substances. Information tracks substances from manufactured to sale to dispensing level.

(5) *An estimate of the total number of respondents and the amount of time estimated for an average respondent to respond/reply:* 1,126 respondents with an average 1 hour per paper response and 10 minutes per electronic response.

(6) *An estimate of the total public burden (in hours) associated with the collection:* 1,700 annual burden hours.

If additional information is required contact: Mr. Robert B. Briggs, Clearance Officer, United States Department of Justice, Information Management and Security Staff, Justice Management Division, United States Department of

Justice, Patrick Henry Building, Suite 1600, 601 D Street NW, Washington, DC 20004.

Dated: December 18, 2001.

Robert B. Briggs,

Department Clearance Officer, United States Department of Justice.

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whether the information will have practical utility;

(2) Evaluate the accuracy of the agency's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;

(3) Enhance the quality, utility, and clarity of the information to be collected; and

(4) Minimize the burden of the collection of information on those who are to respond, including through the use of appropriated automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, e.g., permitting electronic submission of responses.

Overview of This Information Collection

(1) *Type of information collection:* Extension of a currently approved collection.

(2) *The title of the form/collection:* U.S. Official Order Forms for Schedules I and II Controlled Substances (ACCOUNTABLE FORMS), Order Form Requisition.

(3) *The agency form number, if any, and the applicable component of the Department sponsoring the collection:* Form No.: DEA Form 222 and DEA Form 222a Office of Diversion Control, Drug Enforcement Administration, U.S. Department of Justice.

(4) *Affected public who will be asked or required to respond, as well as brief abstract:* Primary: Business or other for-profit. Other: Individuals or households, Federal Government, and State, Local or Tribal Government. DEA-222 is used to transfer or purchase Schedule I and II controlled substances and data is needed to provide an audit of transfer and purchase. DEA-222a Requisition Form is used to obtain the DEA-222 Order Form. Respondents are DEA registrants desiring to handle these controlled substances.

(5) *An estimate of the total number of respondents and the amount of time estimated for an average respondent to respond/reply:* 89,908 respondents with an average 17.5 minutes per response.

(6) *An estimate of the total public burden (in hours) associated with the collection:* 193,508 annual burden hours.

If additional information is required contact: Mr. Robert B. Briggs, Clearance Officer, United States Department of Justice, Information Management and Security Staff, Justice Management Division, United States Department of Justice, Patrick Henry Building, Suite 1600, 601 D Street NE, Washington, DC 20004.

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Agency Information Collection Activities: Proposed Collection; Comment Request

ACTION: 30-day notice of information collection under review: extension of a currently approved collection; U.S. official order forms for Schedules I and II controlled substances (ACCOUNTABLE FORMS), order form requisition.

The Department of Justice (DOJ), Drug Enforcement Administration (DEA), has submitted the following information collection request to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act of 1995. The proposed information collection is published to obtain comments from the public and affected agencies. This proposed information collection was previously published in the **Federal Register** Volume 66, Number 201, pages 52778–52779 on October 17, 2001, allowing for a 60-day comment period.

The purpose of this notice is to allow for an additional 30 days for public comment until January 28, 2002. This process is conducted in accordance with 5 CFR 1320.10.

Written comments and/or suggestions regarding the items contained in this notice, especially the estimated public burden and associated response time, should be directed to The Office of Management and Budget, Office of Information and Regulatory Affairs, Attention Department of Justice Desk Officer, Washington, DC 20503. Additionally, comments may be submitted to OMB via facsimile to (202)-395-7285.

Written comments and/or suggestions from the public and affected agencies concerning the proposed collection of information should address one or more of the following four points:

(1) Evaluate whether the proposed collection of information is necessary for the proper performance of the function of the agency, including

Dated: December 18, 2001.

Robert B. Briggs,

Department Clearance Officer, United States Department of Justice.

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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 September 27, 2001, Chattem Chemicals, Inc., 3801 St. Elmo Avenue, Building 18, Chattanooga, Tennessee 37409, made application by letter to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of dextropropoxyphene (9273), a basic class of controlled substance listed in Schedule II.

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

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, DC 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than February 25, 2002.

Dated: December 18, 2001.

Laura M. Nagel,

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

[FR Doc. 01-31821 Filed 12-26-01; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated August 24, 2001, and published in the **Federal Register** on September 6, 2001, (66 FR 46653), Chattem Chemicals, Inc., 3801 St. Elmo Avenue, Building 18, Chattanooga, Tennessee 37409, made application by renewal 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
4-Methoxyamphetamine (7411)	I
2,5-Dimethoxyamphetamine (7396)	I
Difenoxin	I
Amphetamine (1100)	II
Methamphetamine (1105)	II
Methylphenidate (1724)	II
Pentobarital (2270)	II
Secobarbital (2315)	II
Codeine (9050)	II
Oxycodone (9143)	II
Diphenoxylate (9170)	II
Hydrocodone (9193)	II
Meperidine (9230)	II
Morphine (9300)	II
Thebaine (9333)	II
Alfentanil (9737)	II
Sufentanil (9740)	II
Fentanyl (9801)	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: December 18, 2001.

Laura M. Nagel,

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

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DEPARTMENT OF LABOR

Office of the Secretary

Submission for OMB Review; Comment Request

December 13, 2001.

The Department of Labor (DOL) has submitted the following public information collection requests (ICRs) to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act of 1995 (Pub. L. 104-13, 44 U.S.C. Chapter 35). A copy of this ICR, with applicable supporting documentation, may be obtained by calling the Department of Labor. To obtain documentation contact Darrin King on (202) 693-4129 or E-Mail: *King-Darrin@dol.gov*.

Comments should be sent to Office of Information and Regulatory Affairs, Attn: OMB Desk Officer for OSHA, Office of Management and Budget, Room 10235, Washington, DC 20503 (202) 395-7316), within 30 days from the date of this publication in the **Federal Register**.

The OMB is particularly interested in comments which:

- Evaluate whether the proposed collection of information is unnecessary for the proper performance of the functions of the agency, including whether the information will have practical utility.
- Evaluate the accuracy of the agency's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;
- Enhance the quality, utility, and clarity of the information to be collected; and
- Minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, e.g., permitting electronic submission of responses.

Type of Review: Existing collection without an OMB control number.

Agency: Occupational Safety and Health Administration (OSHA).

Title: Occupational Safety and Health State Plan Information.

OMB Number: 1218-ONEW.

Affected Public: State, Local, or Tribal Government.

Frequency: On occasion, Quarterly, and Annually.

Type of Response: Reporting.

Number of Respondents: 26.

Number of Annual Responses: 1,552.