

controlled substances listed in Schedules II:

Drug	Schedule
Methylphenidate (1724) .....	II
Phenylacetone (8501) .....	II
Methadone intermediate (9254) ...	II

The company plans to manufacture the listed controlled substance in bulk for sale to its customer.

Any other such applicant and any person who is presently registered with DEA to manufacture such a substance may file comments or objections to the issuance of the proposed registration pursuant to 21 CFR 1301.33(a).

Any such written comments or objections being sent via regular mail should be addressed, in quintuplicate, to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, Washington, DC. 20537, Attention: DEA **Federal Register** Representative/ODL; or any being sent via express mail should be sent to DEA Headquarters, Attention: DEA **Federal Register** Representative/ODL, 2401 Jefferson-Davis Highway, Alexandria, Virginia 22301; and must be filed no later than October 2, 2006.

Dated: July 26, 2006.

**Joseph T. Rannazzisi,**

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

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## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

#### Manufacturer of Controlled Substances; Notice of Application

Pursuant to Section 1301.33(a) Title 21 of the Code of Federal Regulations (CFR), this is notice that on June 27, 2006, Noramco Inc., 1440 Olympic Drive, Athens, Georgia 30601, made application by letter to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of basic drug code (1724) methylphenidate.

The company plans to bulk manufacture methylphenidate for a customer to use in the production of a controlled substance product.

Any other such applicant and any person who is presently registered with DEA to manufacture such a substance may file comments or objections to the issuance of the proposed registration pursuant to 21 CFR 1301.33(a).

Any such written comments or objections being sent via regular mail

should be addressed, in quintuplicate, to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, Washington, DC. 20537, Attention: DEA **Federal Register** Representative/ODL; or any being sent via express mail should be sent to DEA Headquarters, Attention: DEA Federal Representative/ODL, 2401 Jefferson-Davis Highway, Alexandria, Virginia 22301; and must be filed no later than October 2, 2006.

Dated: July 26, 2006.

**Joseph T. Rannazzisi,**

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

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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 February 19, 2006, Orasure Technologies, Inc., Lehigh University, Seeley G. Mudd-Building 6, Bethlehem, Pennsylvania 18015, made application by renewal, and by letter, to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the basic classes of controlled substances listed in Schedule I and II:

Drug	Schedule
Lysergic acid diethylamide (LSD) (7315) .....	I
4-Methoxyamphetamine (7411) ...	I
Normorphine (9313) .....	I
Tetrahydrocannabinols (THC) (7370) .....	I
Alphamethadol (9605) .....	I
Amphetamine (1100) .....	II
Methamphetamine (1105) .....	II
Cocaine (9041) .....	II
Hydromorphone (9150) .....	II
Benzoyllecgonine (9180) .....	II
Hydrocodone (9193) .....	II
Morphine (9300) .....	II
Oxycodone (9143) .....	II
Meperidine (9230) .....	II
Methadone (9250) .....	II
Oxymorphone (9652) .....	II

The company plans to manufacture the listed controlled substances in bulk to manufacture controlled substance derivatives. These derivatives will be used in diagnostic products created specifically for internal use only.

Any other such applicant and any person who is presently registered with DEA to manufacture such a substance

may file comments or objections to the issuance of the proposed registration pursuant to 21 CFR § 1301.33(a).

Any such written comments or objections being sent via regular mail may be addressed, in quintuplicate, to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, Washington, DC. 20537, Attention: DEA **Federal Register** Representative, Liaison and Policy Section (ODL); or any being sent via express mail should be sent to DEA Headquarters, Attention: DEA **Federal Register** Representative/ODL, 2401 Jefferson-Davis Highway, Alexandria, Virginia 22301; and must be filed no later than October 2, 2006.

Dated: July 26, 2006.

**Joseph T. Rannazzisi,**

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

[FR Doc. E6-12459 Filed 8-1-06; 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 January 10, 2006, Siegfried (USA), Inc., Industrial Park Road, Pennsville, New Jersey 08070, 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 in Schedule II:

Drug	Schedule
Amphetamine (1100) .....	II
Methylphenidate (1724) .....	II
Amobarbital (2125) .....	II
Pentobarbital (2270) .....	II
Secobarbital (2315) .....	II
Glutethimide (2550) .....	II
Codeine (9050) .....	II
Oxycodone (9143) .....	II
Hydromorphone (9150) .....	II
Hydrocodone (9193) .....	II
Methadone (9250) .....	II
Methadone intermediate (9254) ...	II
Dextropropoxyphene, bulk (non-dosage forms) (9273) .....	II
Morphine (9300) .....	II

The company plans to manufacture the listed controlled substances in bulk for distribution to its customers.

Any other such applicant and any person who is presently registered with DEA to manufacture such a substance may file comments or objections to the

issuance of the proposed registration pursuant to 21 CFR 1301.33(a).

Any such written comments or objections being sent via regular mail may be addressed, in quintuplicate, to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, Washington, DC 20537, Attention: DEA Federal Register Representative, Liaison and Policy Section (ODL); or any being sent via express mail should be sent to DEA Headquarters, Attention: DEA Federal Register Representative/ODL, 2401 Jefferson-Davis Highway, Alexandria, Virginia 22301; and must be filed no later than October 2, 2006.

July 26, 2006.

**Joseph T. Rannazzisi,**

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

[FR Doc. E6-12475 Filed 8-1-06; 8:45 am]

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## DEPARTMENT OF JUSTICE

### Office of Justice Programs

#### **Bureau of Justice Assistance; Agency Information Collection Activities: Extension of a Currently Approved Collection; Comments Requested**

**AGENCY:** Office of Justice Programs, Department of Justice.

**ACTION:** 30 Day Notice of Information Collection Under Review: Extension of a currently approved collection.

Bureau of Justice Assistance  
Application Form: *Southwest Border Prosecution Initiative* [OMB Number 1121-0270].

The Department of Justice (DOJ), Office of Justice Programs (OJP) 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 collection information is published to obtain comments from the public and affected agencies. This proposed information collection was previously published in the **Federal Register** [Volume 71, Number 104, pages 30962-30963 on May 31, 2006,] allowing for a 60 day comment period. The purpose of this notice is to allow for an additional 30 days for public comment until September 1, 2006. This process is conducted in accordance with 5 CFR 1320.10.

Written comments and/or suggestions regarding the item(s) contained in this notice, especially regarding 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 20530.

Additionally, comments may be submitted to OMB via facsimile to (202) 395-7285. Comments may also be submitted to the M. Pressley, Bureau of Justice Assistance, Office of Justice Programs, U.S. Department of Justice, 810 7th Street, NW., Washington, DC 20531 via facsimile to (202) 514-1590.

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 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:

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

(2) *The title of the form/collection:* Bureau of Justice Assistance Application Form for the Southwest Border Prosecution Initiative.

(3) *The agency form number, if any and the applicable component of the Department sponsoring the collection:* None.

(4) *Affected public who will be asked or required to respond, as well as a brief abstract:* Primary: State, Local or Tribal government.

*Other:* None.

*Abstract:* The Southwest Border Prosecutor Initiative was enacted in FY 2002 to reimburse state, county, parish, or municipal governments for the costs associated with the prosecution of criminal cases declined by local U.S. Attorneys. Each year, hundreds of criminal cases resulting from federal arrests are referred to local prosecutors to handle when the cases fall below certain monetary, quantity, or severity

thresholds. This places additional burdens on local government resources that are already stretched by the demands of prosecuting violations of local and state laws. This program provides funds to eligible jurisdictions in the four southwest border states, using a uniform payment-per-case basis for qualifying federally initiated and declined-referred criminal cases that were disposed of after October 1, 2001. Up to 220 eligible jurisdictions may apply. This includes county governments and the four state governments in Arizona, California, New Mexico, and Texas.

(5) *An estimate of the total number of respondents and the amount of time estimated for an average respondent to respond/reply:* It is estimated that no more than 220 respondents will apply. Each application takes approximately 60 minutes to complete and is submitted 4 times per year (quarterly).

(6) *An estimate of the total public burden (in hours) associated with the collection:* The total hour burden to complete the applications is 880 hours (880 applications (220 × 4 times a year) × 60 minutes per application = 52,800/60 minutes per hour = 880 burden hours).

*If additional information is required contact:* Lynn Bryant, Department Clearance Officer, United States Department of Justice, Policy and Planning Staff, Justice Management Division, 601 D Street, NW., Suite 1600, Washington, DC, 20530, or via phone at 202-514-4304.

Dated: July 28, 2006.

**Lynn Bryant,**

*Department Clearance Officer, Justice Management Division, PRA, United States Department of Justice.*

[FR Doc. E6-12454 Filed 8-1-06; 8:45 am]

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## NUCLEAR REGULATORY COMMISSION

### **Agency Information Collection Activities: Submission for the Office of Management and Budget (OMB) Review; Comment Request**

**AGENCY:** Nuclear Regulatory Commission (NRC).

**ACTION:** Notice of the OMB review of information collection and solicitation of public comment.

**SUMMARY:** The NRC has recently submitted to OMB for review the following proposal for the collection of information under the provisions of the Paperwork Reduction Act of 1995 (44 U.S.C. Chapter 35). The NRC hereby