DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Application

Pursuant to § 1301.33(a) of Title 21 of the Code of Federal Regulations (CFR), this is notice that on January 5, 2007, Roche Diagnostics Operations, Inc., Attn: Regulatory Compliance, 9115 Hague Road, Indianapolis, Indiana 46250, 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 I and II:

Drug	Schedule
Lysergic Acid Diethylamide (7315)	I
Tetrahydrocannabinols (7370)	I
Alphamethadol (9605)	I
Phencyclidine (7471)	II
Ecgonine (9180)	II
Methadone (9250)	II
Morphine (9300)	II

The company plans to manufacture small quantities of the listed controlled substances for use in diagnostic products.

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, Attention: DEA Federal Register Representative/ODL, Washington, DC 20537; 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 May 29, 2007.

Dated: March 19, 2007.

Joseph T. Rannazzisi,

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

[FR Doc. E7–5510 Filed 3–26–07; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Application

Pursuant to § 1301.33(a) of Title 21 of the Code of Federal Regulations (CFR), this is notice that on January 26, 2007, Stepan Company, Natural Products Dept., 100 W. Hunter Avenue, Maywood, New Jersey 07607, 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
Cocaine (9041)	II
Ecgonine (9180)	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 should be addressed, in quintuplicate, to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, Attention: DEA Federal Register Representative/ODL, Washington, DC 20537; 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 May 29, 2007.

Dated: March 19, 2007.

Joseph T. Rannazzisi,

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

[FR Doc. E7–5509 Filed 3–26–07; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Application

Pursuant to 21 U.S.C. § 958(i), the Attorney General shall, prior to issuing a registration under this Section to a bulk manufacturer of a controlled substance in schedule I or II and prior to issuing a regulation under 21 U.S.C. 952(a) authorizing the importation of such a substance, provide manufacturers holding registrations for the bulk manufacture of the substance an opportunity for a hearing.

Therefore, in accordance with 21 CFR 1301.34(a), this is notice that on November 9, 2006, Tocris Cookson, Inc., 16144 Westwoods Business Park, Ellisville, Missouri 63021–4500, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as an importer of the basic classes of controlled substances listed in schedule I:

Drug	Schedule
Marihuana (7360)	I
Tetrahydrocannabinols (7370)	I

The company plans to import the above listed synthetic products for non-clinical laboratory based research only.

Any bulk manufacturer who is presently, or is applying to be, registered with DEA to manufacture such basic classes of controlled substances may file comments or objections to the issuance of the proposed registration and may, at the same time, file a written request for a hearing on such application pursuant to 21 CFR 1301.43 and in such form as prescribed by 21 CFR 1316.47.

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, Attention: DEA Federal Register Representative/ODL, Washington, DC 20537; 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 April 26, 2007.

This procedure is to be conducted simultaneously with and independent of the procedures described in 21 CFR 1301.34(b), (c), (d), (e) and (f). As noted in a previous notice published in the **Federal Register** on September 23, 1975, (40 FR 43745–46), all applicants for registration to import a basic class of any controlled substance listed in schedule I or II are, and will continue to be required to demonstrate to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, that the requirements for such registration pursuant to 21 U.S.C. 958(a), 21 U.S.C. 823(a), and 21 CFR 1301.34(b), (c), (d), (e) and (f) are satisfied.

Dated: March 19, 2007.

Joseph T. Rannazzisi,

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

[FR Doc. E7–5507 Filed 3–26–07; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Office of Justice Programs

[OJP (OJJDP) Docket No. 1468]

Meeting of the Federal Advisory Committee on Juvenile Justice

AGENCY: Office of Juvenile Justice and Delinquency Prevention, Office of Justice Programs, Justice.

ACTION: Notice of meeting.

SUMMARY: The Office of Juvenile Justice and Delinquency Prevention (OJJDP) is announcing the spring meeting of the Federal Advisory Committee on Juvenile Justice (FACJJ), which will be held in Washington, DC on April 23–24, 2007. The meeting times and location are noted below.

DATES: The schedule of events is as follows:

1. Sunday, April 22, 2007, 4 p.m. to 6 p.m.

2. Monday, April 23, 2007, 8:30 a.m. to 5 p.m.

3. Tuesday, April 24, 2007, 8:30 a.m. to 12:30 p.m.

ADDRESSES: All open meeting sessions will take place at the Office of Justice Programs, 810 Seventh Street, NW., main conference room, Washington, DC 20531.

FOR FURTHER INFORMATION CONTACT: Robin Delany-Shabazz, Designated Federal Official, OJJDP, *Robin.Delany-Shabazz@usdoj.gov*, or 202–307–9963. [Note: This is not a toll-free number.]

SUPPLEMENTARY INFORMATION: The Federal Advisory Committee on Juvenile Justice (FACJJ), established pursuant to Section 3(2)A of the Federal Advisory Committee Act (5 U.S.C. App.2) will meet to carry out its advisory functions under Section 223(f)(2)(C–E) of the Juvenile Justice and Delinquency Prevention Act of 2002. The FACJJ is composed of one representative from each state and territory. FACJJ duties include: Reviewing Federal policies regarding juvenile justice and delinquency prevention; advising the OJJDP Administrator with respect to particular functions and aspects of OJJDP; and advising the President and Congress with regard to State perspectives on the operation of OJJDP and Federal legislation pertaining to juvenile justice and delinquency prevention. More information, including a member list, may be found at *http://www.facjj.org.*

Meeting Agenda

1. Sunday, April 22, 2007

- 4 p.m.–6 p.m. New Member Orientation. (Closed Session).

2. Monday, April 23, 2007

- 8:30 a.m.–9:15 a.m. Call to Order by the Chair of the FACJJ and Remarks by the Administrator of OJJDP (Open Session).
- 9:15 a.m.–12 p.m. and 1:30 p.m.–5 p.m. Review, Discussion and Deliberation of the 2007 Draft Reports to the President, Congress, and the Administrator of OJJDP (Open Session).
- 12 p.m.–1:30 p.m. Subcommittee Meetings (Closed Sessions).

3. Tuesday, April 23, 2007

- 8:30 a.m.–12:30 p.m. Continuation of Review, Discussion and Deliberation of the 2007 Draft Reports; Presentations and Discussions concerning Effective Legal Counsel; and Other Business (Open Session).

For security purposes, members of the public who wish to attend open sessions of the meeting should register by sending an e-mail with their name, affiliation, address, phone number, and a list of sessions they plan to attend to *ddunston@edjassociates.com.* If e-mail is not available, fax this information to 240–221–4006, attention: Daryel Dunston. Because space is limited, notification of intent to attend should be sent by April 16, 2007.

Note: Photo identification will be required for admission to the meeting. Additional identification documents may be required. Space is limited.

Written Comments

Interested parties may submit written comments by Monday, April 16, 2007, to Robin Delany-Shabazz, Designated Federal Official for the Federal Advisory Committee on Juvenile Justice, OJJDP, at *Robin.Delany-Shabazz@usdoj.gov*. If e-mail is not available, fax your comments to 202–354–4063 and call Francesca Stern at 202–616–3551 to ensure fax was received. [Note: These are not toll-free numbers.] No oral presentations will be permitted at the meeting. However, written questions and comments from members of the public attending the meeting may be invited.

J. Robert Flores,

Administrator, Office of Juvenile Justice and Delinquency Prevention.

[FR Doc. E7–5544 Filed 3–26–07; 8:45 am]

BILLING CODE 4410–18–P

DEPARTMENT OF LABOR

Office of the Secretary

Submission for OMB Review: Comment Request

March 21, 2007.

The Department of Labor (DOL) has submitted the following public information collection request (ICR) 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 Ira Mills on 202–693–4122 (this is not a toll-free number) or E-Mail: *Mills.Ira@dol.gov*, or by accessing *http://www.reginfo.gov/public/do/PRAMain*.

Comments should be sent to Office of Information and Regulatory Affairs, Attn: OMB Desk Officer for U.S. Department of Labor/Bureau of Labor Statistic (BLS), Office of Management and Budget, Room 10235, Washington, DC 20503, 202–395–7316 (this is not a toll free number), 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 necessary 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.

Agency: Bureau of Labor Statistics.

Type of Review: Revision of a currently approved Collection.

Title: National Longitudinal Survey of Youth.

OMB Number: 1220–0109.

Frequency: Biennially.

Affected Public: Individuals or households.

Type of Response: Reporting.