

Drug	Schedule
Dimethyltryptamine (7435)	I
Psilocybin (7437)	I
Psilocyn (7438)	I
5-Methoxy-N,N-diisopropyltryptamine (7439)	I
N-Benzylpiperazine (7493)	I
MDPV (7535)	I
Methylone (7540)	I
Desomorphine (9055)	I
Etorphine (except HCl) (9056)	I
Heroin (9200)	I
Morphine-N-oxide (9307)	I
Normorphine (9313)	I
Pholcodine (9314)	I
Dextromoramide (9613)	I
Dipipanone (9622)	I
Racemoramide (9645)	I
Trimeperidine (9646)	I
1-Methyl-4-phenyl-4-propionoxypiperidine (9661)	I
Tilidine (9750)	I
Amphetamine (1100)	II
Methamphetamine (1105)	II
Methylphenidate (1724)	II
Amobarbital (2125)	II
Pentobarbital (2270)	II
Secobarbital (2315)	II
Phencyclidine (7471)	II
Phenylacetone (8501)	II
Cocaine (9041)	II
Codeine (9050)	II
Dihydrocodeine (9120)	II
Oxycodone (9143)	II
Hydromorphone (9150)	II
Ecgonine (9180)	II
Ethylmorphine (9190)	II
Meperidine (9230)	II
Methadone (9250)	II
Dextropropoxyphene, bulk (non-dosage forms) (9273)	II
Morphine (9300)	II
Oripavine (9330)	II
Thebaine (9333)	II
Levo-alphaacetylmethadol (9648) ..	II
Oxymorphone (9652)	II
Poppy Straw Concentrate (9670) ..	II
Alfentanil (9737)	II
Sufentanil (9740)	II
Fentanyl (9801)	II

The company plans to import small quantities of the listed controlled substances for the manufacture of analytical reference standards.

In reference to drug codes 7360 and 7370, the company plans to import a synthetic cannabidiol and a synthetic Tetrahydrocannabinol. No other activity for this drug code is authorized for this registration.

Comments and requests for hearing on applications to import narcotic raw material are not appropriate. 72 FR 3417 (2007).

Any bulk manufacturer who is presently, or is applying to be, registered with DEA to manufacture such basic classes of controlled substances listed in schedules I or II, which fall under the authority of section 1002(a)(2)(B) of the Act 21 U.S.C. 952 (a)(2)(B) may, in the circumstances set

forth in 21 U.S.C. 958(i), 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 should be addressed, in quintuplicate, to the Drug Enforcement Administration, Office of Diversion Control, Federal Register Representative (ODL), 8701 Morrisette Drive, Springfield, Virginia 22152; and must be filed no later than October 17, 2013.

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 substances in schedules 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: September 9, 2013.

Joseph T. Rannazzisi,

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

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DEPARTMENT OF JUSTICE

Office of Justice Programs

[OJP (OJP) Docket No. 1631]

Meeting of the Office of Justice Programs' Science Advisory Board

AGENCY: Office of Justice Programs (OJP), Justice.

ACTION: Notice of meeting.

SUMMARY: This notice announces a forthcoming meeting of OJP's Science Advisory Board ("Board"). The Board is chartered to provide OJP, a component of the Department of Justice, with valuable advice in the areas of science and statistics for the purpose of enhancing the overall impact and performance of its programs and activities in criminal and juvenile justice. To this end, the Board has designated six (6) subcommittees: National Institute of Justice (NIJ); Bureau of Justice Statistics (BJS); Office of Juvenile Justice and Delinquency

Prevention (OJJDP); Bureau of Justice Assistance; Quality and Protection of Science; and Evidence Translation/Integration.

DATES: The meeting will take place on Friday, October 4, 2013, from 8:30 a.m. to 4:00 p.m., ET, with a break for lunch at approximately noon.

ADDRESSES: The meeting will take place in the Main Conference Room on the third floor of the Office of Justice Programs, 810 7th Street, Northwest, Washington, DC 20531.

FOR FURTHER INFORMATION CONTACT: Phelan Wyrick, Designated Federal Officer ("DFO"), Office of the Assistant Attorney General, Office of Justice Programs, 810 7th Street Northwest, Washington, DC 20531; Phone: (202) 353–9254 [Note: this is not a toll-free number]; Email: phelan.wyrick@usdoj.gov.

SUPPLEMENTARY INFORMATION: This meeting is being convened to brief the OJP Assistant Attorney General and the Board members on the progress of the subcommittees, discuss any recommendations they may have for consideration by the full Board, and brief the Board on various OJP-related projects and activities. The final agenda is subject to adjustment, but it is anticipated that there will be a morning session and an afternoon session, with a break for lunch. These sessions will likely include briefings of the subcommittees' activities and discussion of future Board actions and priorities.

This meeting is open to the public. Members of the public who wish to attend this meeting must register with Phelan Wyrick at the above address at least seven (7) days in advance of the meeting. Registrations will be accepted on a space available basis. Access to the meeting will not be allowed without registration. Persons interested in communicating with the Board should submit their written comments to the DFO, as the time available will not allow the public to directly address the Board at the meeting. Anyone requiring special accommodations should notify Mr. Wyrick at least seven (7) days in advance of the meeting.

Phelan Wyrick,

Science Policy Advisor and Science Advisory Board DFO, Office of the Assistant Attorney General, Office of Justice Programs.

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