Drug	Schedule
Opium, raw (9600)	II
Opium extracts (9610)	II
Opium fluid extract (9620)	II
Opium tincture (9630)	II
Opium poppy/Poppy Straw (9650)	II
Poppy Straw Concentrate (9670)	II
Opium, granulated (9640)	II
Oxycodone (9143)	II
Oxymorphone (9652)	II
Pentobarbital (2270)	II
Phenazocine (9715)	II
Phencyclidine (7471)	II
Phenmetrazine (1631)	II
Phenylacetone (8501)	II
Piminodine (9730)	II
Powdered opium (9639)	II
Racemethorphan (9732)	II
Racemorphan (9733)	II
Remifentanil (9739)	II
Secobarbital (2315)	II
Sufentanil (9740)	II
Tapentadol (9780)	II
Thebaine (9333)	II

The company plans to import small quantities of the listed controlled substances for the National Institute on Drug Abuse (NIDA) for research activities.

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

In regard to the non-narcotic raw material, any bulk manufacturer who is presently, or is applying to be, registered with DEA to manufacture such basic classes of controlled substances listed in schedule 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 Morrissette Drive, Springfield, Virginia 22152; and must be filed no later than July 9, 2012.

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 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: May 31, 2012.

Joseph T. Rannazzisi,

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

[FR Doc. 2012–13920 Filed 6–7–12; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Registration; Meda Pharmaceuticals, Inc.

By Notice dated April 2, 2012, and published in the Federal Register on April 12, 2012, 77 FR 21998, Meda Pharmaceuticals, Inc., 705 Eldorado Street, Decatur, Illinois 62523, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as an importer of Nabilone (7379), a basic class of controlled substance listed in schedule II.

The company plans to import the listed controlled substance for distribution to its customers.

No comments or objections have been received. DEA has considered the factors in 21 U.S.C. 823(a) and 952(a), and determined that the registration of Meda Pharmaceuticals Inc. to import the basic class of controlled substance is consistent with the public interest, and with United States obligations under international treaties, conventions, or protocols in effect on May 1, 1971. DEA has investigated Meda Pharmaceuticals Inc. to ensure that the company's registration is consistent with the public interest. The investigation has 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. 952(a) and 958(a), and in accordance with 21 CFR 1301.34, the above named company is granted registration as an importer of the basic class of controlled substance listed.

Dated: May 31, 2012.

Joseph T. Rannazzisi,

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

[FR Doc. 2012–13916 Filed 6–7–12; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Application; Rhodes Technologies

Pursuant to § 1301.33(a), Title 21 of the Code of Federal Regulations (CFR), this is notice that on May 1, 2012, Rhodes Technologies, 498 Washington Street, Coventry, Rhode Island 02816, made application to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of

Morphine (9300), a basic class of controlled substance listed in schedule II.

The company plans to manufacture the listed controlled substance in bulk for conversion and sale to dosage form manufacturers.

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 pursuant to 21 CFR 1301.33(a).

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 Morrissette Drive, Springfield, Virginia 22152; and must be filed no later than August 7, 2012.

Dated: May 31, 2012.

Joseph T. Rannazzisi,

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

[FR Doc. 2012–13919 Filed 6–7–12; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Application; S&B Pharma, Inc.

Pursuant to § 1301.33(a), Title 21 of the Code of Federal Regulations (CFR), this is notice that on April 4, 2012, S&B Pharma Inc., 405 South Motor Avenue, Azusa, California 91702–3232, made application to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the following basic classes of controlled substances:

Drug	Schedule
Gamma Hydroxybutyric Acid (2010).	I
Tetrahydrocannabinols (7370)	I
Methamphetamine (1105)	II
Pentobarbital (2270)	II
Nabilone (7379)	II

The company plans to manufacture bulk controlled substances for use in product development and for distribution to its customers.

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 pursuant to 21 CFR 1301.33(a).

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 Morrissette Drive, Springfield, Virginia 22152; and must be filed no later than August 7, 2012.

Dated: May 31, 2012.

Joseph T. Rannazzisi,

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

[FR Doc. 2012–13918 Filed 6–7–12; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration; Pharmagra Labs, Inc.

By Notice dated January 30, 2012, and published in the **Federal Register** on February 6, 2012, 77 FR 5846, Pharmagra Labs Inc., 158 McLean Road, Brevard, North Carolina 28712, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of Pentobarbital (2270), a basic class of controlled substance in schedule II.

The company plans to manufacture the listed substance for analytical research and clinical trials.

No comments or objections have been received. DEA has considered the factors in 21 U.S.C. 823(a), and determined that the registration of Pharmagra Labs, Inc. to manufacture the listed basic class of controlled substance is consistent with the public interest at this time. DEA has investigated Pharmagra Labs, Inc. to ensure that the company's registration is consistent with the public interest. The investigation has 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(a), and in accordance with 21 CFR 1301.33, the above named company is granted registration as a bulk manufacturer of the basic class of controlled substance listed.

Dated: May 31, 2012.

Joseph T. Rannazzisi,

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

[FR Doc. 2012–13917 Filed 6–7–12; 8:45 am]

BILLING CODE 4410–09–P

OFFICE OF MANAGEMENT AND BUDGET

Office of Federal Procurement Policy

Value Engineering

AGENCY: Office of Federal Procurement Policy, Office of Management and Budget.

ACTION: Proposed revision to Office of Management and Budget Circular No. A–131, ''Value Engineering''.

SUMMARY: The Office of Federal Procurement Policy (OFPP) in the Office of Management and Budget (OMB) is proposing to revise OMB Circular A–131, Value Engineering, to update and reinforce policies associated with the consideration and use of Value Engineering (VE). VE is an effective technique for cutting waste and inefficiency—helping Federal agencies save billions of dollars in program and acquisition costs, improve performance, enhance quality, and foster the use of innovation. The proposed revisions are designed to ensure that the Federal Government has the capabilities and tools to consider and apply VE techniques to the maximum extent appropriate.

DATES: Interested parties should submit comments in writing to the address below on or before August 7, 2012.

ADDRESSES: Comments may be submitted by any of the following methods:

- *Online at: http://www.regulations.gov.*
- *Facsimile:* 202–395–5105.
- *Mail:* Office of Federal Procurement Policy, ATTN: Curtina Smith, New Executive Office Building, Room 9013, 725 17th Street NW., Washington, DC 20503.

Instructions: Please submit comments only and cite ''Proposed Revision to OMB Circular A–131'' in all correspondence. All comments received will be posted, without change or redaction, to *www.regulations.gov,* so commenters should not include information that they do not wish to be posted (for example because they consider it personal or business confidential).

FOR FURTHER INFORMATION CONTACT: Curtina Smith, OFPP, *csmith@omb.eop.gov.* Availability: Copies of the proposed revision to OMB Circular A–131 are available on OMB's Web site at *http://www.whitehouse.gov/omb/circulars_default/.*

SUPPLEMENTARY INFORMATION: