

Dated: January 6, 2012.

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 § 1301.33(a), of Title 21 of the Code of Federal Regulations (CFR), this is notice that on September 6, 2011 Cedarburg Pharmaceuticals, Inc., 870 Badger Circle, Grafton, Wisconsin 53024, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the following basic classes of controlled substances:

Drug	Schedule
4-Anilino-N-phenethyl-4-piperidine (8333).	II
Fentanyl (9801)	II

The company plans to manufacture the listed controlled substances in bulk for distribution to its customers. Regarding the drug code (8333), the company plans to use this controlled substance to manufacture another controlled substance.

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 Morrisette Drive, Springfield, Virginia 22152; and must be filed no later than March 19, 2012.

Dated: January 6, 2012.

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 § 1301.33(a), Title 21 of the Code of Federal Regulations (CFR), this is notice that on September 12, 2011, Johnson Matthey, Inc., Pharmaceuticals Materials, 900 River Road, Conshohocken, Pennsylvania 19428, made application by renewal 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
Amphetamine (1100)	II
Methylphenidate (1724)	II
Codeine (9050)	II
Oxycodone (9143)	II
Hydrocodone (9193)	II
Morphine (9300)	II
Thebaine (9333)	II

The company plans to manufacture the listed controlled substances in bulk for sale in bulk to its customers. The Thebaine (9333) will also be used to manufacture other controlled substances in bulk which will also be for sale in bulk 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 Morrisette Drive, Springfield, Virginia 22152; and must be filed no later than March 19, 2012.

Dated: January 6, 2012.

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 Registration

By Notice dated August 9, 2011, and published in the **Federal Register** on August 18, 2011, 76 FR 51401, Austin

Pharma LLC., 811 Paloma Drive, Suite C, Round Rock, Texas 78665-2402, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the following basic classes of controlled substances:

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

The company plans to manufacture bulk active pharmaceutical ingredients (APIs) for distribution to its customers. In reference to drug code 7360

(Marihuana), the company plans to bulk manufacture cannabidiol as a synthetic intermediate. This controlled substance will be further synthesized to bulk manufacture a synthetic THC (7370). No other activity for this drug code is authorized for this registration.

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 Austin Pharma LLC. to manufacture the listed basic classes of controlled substances is consistent with the public interest at this time. DEA has investigated Austin Pharma LLC. 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 classes of controlled substances listed.

Dated: January 6, 2012.

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 Registration

By Notice dated August 10, 2011, and published in the **Federal Register** on August 18, 2011, 76 FR 51401, AMRI Rensselaer, Inc., 33 Riverside Avenue, Rensselaer, New York 12144, made application by renewal to the Drug

Enforcement Administration (DEA) to be registered as a bulk manufacturer of the following basic classes of controlled substances:

Drug	Schedule
Marihuana (7360)	I
Tetrahydrocannabinols (7370)	I
Amphetamine (1100)	II
Lisdexamfetamine (1205)	II
Methylphenidate (1724)	II
Pentobarbital (2270)	II
Meperidine (9230)	II
4-Anilino-N-phenethyl-4-piperidine (8333)	II
Fentanyl (9801)	II

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

In reference to drug code 7360 (Marihuana), the company plans to bulk manufacture cannabidiol as a synthetic intermediate. This controlled substance will be further synthesized to bulk manufacture a synthetic THC (7370). No other activity for this drug code is authorized for this registration.

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 AMRI Rensselaer, Inc. to manufacture the listed basic classes of controlled substances is consistent with the public interest at this time. DEA has investigated AMRI Rensselaer, 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, and in accordance with 21 CFR 1301.33, the above named company is granted registration as a bulk manufacturer of the basic classes of controlled substances listed.

Dated: January 6, 2012.

Joseph T. Rannazzisi,

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

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DEPARTMENT OF JUSTICE

National Institute of Corrections

Solicitation for a Cooperative Agreement—Jail Resource Management: Review and Revision

AGENCY: National Institute of Corrections, U.S. Department of Justice.

ACTION: Solicitation for a Cooperative Agreement.

SUMMARY: The National Institute of Corrections (NIC) is seeking applications for the revision of its Jail Resource Management training program. The project will be for a 9-month period and will be carried out in conjunction with the NIC Jails Division. The awardee will work closely with NIC staff on all aspects of the project. To be considered, applicants must demonstrate, at a minimum, in-depth knowledge of (1) the purpose, functions, and operational complexities of local jails, (2) budget issues common in jails, (3) analysis of jail resource needs, (4) development and presentation of a budget request to appropriate governing bodies, (5) budget management, and (6) the resource constraints faced by many local governments and their jails. Also, the applicant must demonstrate expertise and experience in developing curricula based on adult learning principles, specifically the Instructional Theory into Practice (ITIP) model.

DATES: Applications must be received by 4 p.m. (EDT) on Thursday, February 9, 2012.

ADDRESSES: Mailed applications must be sent to: Director, National Institute of Corrections, 320 First Street NW., Room 5002, Washington, DC 20534. Applicants are encouraged to use Federal Express, UPS, or similar service to ensure delivery by the due date as mail at NIC is sometimes delayed due to security screening.

Applicants who wish to hand-deliver their applications should bring them to 500 First Street NW., Washington, DC 20534, and dial (202) 307-3106, ext. 0, at the front desk for pickup.

Faxed or emailed applications will not be accepted; however, electronic applications can be submitted via <http://www.grants.gov>.

FOR FURTHER INFORMATION CONTACT: A copy of this announcement and the required application forms can be downloaded from the NIC Web site at www.nic.gov/cooperativeagreements.

Questions about this project and the application procedures should be directed to Erika McDuffe, Correctional Program Specialist, National Institute of Corrections. Questions must be emailed

to Ms. McDuffe at emcduffe@bop.gov. Ms. McDuffe will respond by email to the individual. Also, all questions and responses will be posted on NIC's Web site at www.nic.gov for public review. (The names of those submitting the questions will not be posted.) The Web site will be updated regularly and postings will remain on the Web site until the closing date of this cooperative agreement solicitation.

SUPPLEMENTARY INFORMATION:

Background: NIC's Jail Resource Management course is a 3-day training program that focuses on the resource needs of the jail, development and presentation of a budget request, management of the budget, and identification of alternate funding options. This 3-day program is held in regions throughout the county.

Program participants are primarily sheriffs and administrators from smaller jails who have no specialized fiscal management staff. Most have no formally established mechanisms to identify, document, track, justify, or present resource needs. As a result, their jails often receive budget allocations that are inadequate to maintain a safe and secure jail.

NIC wishes to update the content of Jail Resource Management and ensure its design conforms to the ITIP model. The following reference materials are posted with this announcement on NIC's Web site: Jail Resource Management: Lesson Plans; Jail Resource Management: Participant Manual; Jail Resource Management: Presentation Slides; Jail Resource Management: Activities.

Scope of Work: The cooperative agreement awardee will revise the content of the current program to ensure it is current, accurate, and relevant. The awardee also will ensure that module sequencing is logical and enhances the flow of the program. Finally, the awardee will revise the program's design to conform to the ITIP model. The awardee will ensure that content, module sequencing, and instructional strategies effectively contribute to meeting the program's goal. To achieve this, the awardee will complete the following activities, at a minimum.

Initial Meeting: The cooperative agreement awardee, with subject matter expert and the curriculum specialist, will attend an initial meeting with the NIC staff for a project overview and preliminary planning. This will take place shortly after the cooperative agreement is awarded. The meeting will last up to one half day and will be conducted via Web conferencing.

Initial curriculum review: The awardee will review and become