

Dated: February 24, 2006.

Thomas A. Readinger,

Associate Director for Offshore Minerals Management.

[FR Doc. E6-4303 Filed 3-23-06; 8:45 am]

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DEPARTMENT OF THE INTERIOR

Bureau of Reclamation

Upper Rio Grande Basin Water Operations Review, NM; Notice of Extension

AGENCY: Bureau of Reclamation, Interior.

ACTION: Notice of extension of public comment period for thirty days.

SUMMARY: Notice is hereby given that the comment period for the Draft Environmental Impact Statement (DEIS) for the Upper Rio Grande Water Operations Review, DES-05-80, is extended an additional 30 days to April 20, 2006.

DATES: The end of the public comment period, as noted in the **Federal Register** (71 FR 3323) on January 20, 2006, was March 21, 2006. The public comment period is now extended to April 20, 2006.

ADDRESSES: Written comments on the DEIS should be addressed to Valda Terauds, ALB-707, Bureau of Reclamation, Albuquerque Area Office, 555 Broadway, NW., Suite 100, Albuquerque, New Mexico 87102; faxogram (505) 462-3593; e-mail: vterauds@uc.usbr.gov. Our practice is to make comments, including names and home addresses of respondents, available for public review. Individual respondents may request that we withhold their home address from public disclosure, which we will honor to the extent allowable by law. If you wish us to withhold your name and/or address, you must state this prominently at the beginning of your comment. We will make all submissions from organizations or businesses, and from individuals identifying themselves as representatives or officials of organizations or businesses, available for public disclosure in their entirety.

FOR FURTHER INFORMATION CONTACT: Valda Terauds, Resource Management Planner, (505) 462-3584.

Dated: March 7, 2006.

Roger Slater,

Acting Regional Director—UC Region, Bureau of Reclamation.

[FR Doc. E6-4306 Filed 3-23-06; 8:45 am]

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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 June 13, 2005, Cerilliant API Services, LLC, 811 Paloma Drive, Suite A, Round Rock, Texas 78664, made application to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of the basic classes of controlled substances listed in Schedule I and II; and by letter dated September 2, 2005, to modify its name to Austin Pharma, LLC.

Drug	Schedule
Lysergic acid diethylamide (7315)	I
Marihuana (7360)	I
Tetrahydrocannabinols (7370)	I
3,4-Methylenedioxymphetamine (7400)	I
3,4-Methylenedioxy-N-ethylamphetamine (7404)	I
3,4-Methylenedioxymethamphetamine (7405)	I
Psilocyn (7438)	I
Acetyldihydrocodeine (9051)	I
Benzylmorphine (9052)	I
Codeine-N-oxide (9053)	I
Cyprenorphine (9054)	I
Desomorphine (9055)	I
Etorphine (9056)	I
Codeine methylbromide (9070)	I
Dihydromorphine (9145)	I
Heroin (9200)	I
Hydromorphanol (9301)	I
Methyldihydromorphine (9304)	I
Morphine methylbromide (9305)	I
Morphine-N-oxide (9307)	I
Alphamethadol (9605)	I
Normethadone (9635)	I
Amphetamine (1100)	II
Methamphetamine (1105)	II
Cocaine (9041)	II
Codeine (9050)	II
Dihydrocodeine (9120)	II
Oxycodone (9143)	II
Hydromorphone (9150)	II
Benzoylcegonine (9180)	II
Ecgonine (9180)	II
Hydrocodone (9193)	II
Levomethorphan (9210)	II
Methadone (9250)	II
Methadone intermediate (9254)	II
Morphine (9300)	II
Thebaine (9333)	II
Levo-alphaacetyl methadol (9648)	II
Oxymorphone (9652)	II
Poppy Straw Concentrate (9670)	II
Alfentanil (9737)	II
Remifentanil (9739)	II
Sufentanil (9740)	II
Carfentanil (9743)	II
Fentanyl (9801)	II

The company plans to manufacture the listed controlled substances in bulk 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.

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 May 23, 2006.

Dated: March 20, 2006.

Joseph T. Rannazzisi,

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

[FR Doc. E6-4302 Filed 3-23-06; 8:45 am]

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DEPARTMENT OF LABOR

Office of the Secretary

Submission for OMB Review: Comment Request

March 17, 2006.

The Department of Labor (DOL) has submitted the following public information collection requests (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 each ICR, with applicable supporting documentation, may be obtained by contacting Darrin King on 202-693-4129 (this is not a toll-free number) or e-mail: king.darrin@dol.gov.

Comments should be sent to Office of Information and Regulatory Affairs, Attn: OMB Desk Officer for the Occupational Safety and Health Administration (OSHA), 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: Occupational Safety and Health Administration.
Type of Review: Extension of currently approved collection.
Title: Lead in General Industry (29 CFR 1910.1025).
OMB Number: 1218-0092.
Frequency: On occasion; Quarterly; Semi-annually; and Annually.
Type of Response: Recordkeeping and Third party disclosure.
Affected Public: Business or other for-profit; Federal Government; and State, Local, or Tribal Government.
Number of Respondents: 62,357.
Number of Annual Responses: 4,068,503.
Estimated Time per Response: Ranges from 1 minute to notify an employee of their right to seek a second medical opinion to 2 hours for an employee to receive a medical examination.
Total Burden Hours: 1,242,562.
Total Annualized capital/startup costs: \$0.
Total Annual Costs (operating/maintaining systems or purchasing services): \$139,869,058.
Description: The purpose of 29 CFR 1910.1025 and its information collection requirements is to provide protection for employees from the adverse effects associated with occupational exposure to the carcinogen, lead. Employers must monitor employee exposure to lead, provide medical surveillance, train employees about the hazards of lead, and establish and maintain accurate records of employee exposure to lead. These records are used by employers, employees, physicians, and the

Government to ensure that employees are not being harmed by exposure to lead.

Agency: Occupational Safety and Health Administration.

Type of Review: Extension of currently approved collection.

Title: Lead in Construction Standard (29 CFR 1926.62).

OMB Number: 1218-0189.

Frequency: On occasion; Quarterly; Semi-annually; and Annually.

Type of Response: Recordkeeping and Third party disclosure.

Affected Public: Business or other for-profit; Federal Government; and State, Local, or Tribal Government.

Number of Respondents: 147,073.

Number of Annual Responses: 5,782,074.

Estimated Time per Response: Ranges from 1 minute to notify an employee of their right to seek a second medical opinion to 8 hours to develop a written compliance program.

Total Burden Hours: 1,560,717.

Total Annualized capital/startup costs: \$0.

Total Annual Costs (operating/maintaining systems or purchasing services): \$68,576,673.

Description: 29 CFR 1926.62 requires employers to train employees about the hazards of lead, monitor employee exposure, to provide medical surveillance, and maintain accurate records of employee exposure to lead. These records are used by employers, employees, physicians and the Government to ensure that employees are not harmed by exposure to lead in the workplace.

Ira L. Mills,

Departmental Clearance Officer.

[FR Doc. E6-4271 Filed 3-23-06; 8:45 am]

BILLING CODE 4510-26-P

DEPARTMENT OF LABOR

Office of the Secretary

Submission for OMB Review: Comment Request

March 17, 2006.

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 the Department of Labor. To obtain documentation contact Ira Mills on 202-693-4122 (this is not a toll-free number) or e-mail: Mills.Ira@dol.gov.

Comments should be sent to Office of Information and Regulatory Affairs, Attn: OMB Desk Officer for ETA, 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: Employment and Training Administration (ETA).
Type of Review: New Collection.
Title: Generic Solicitation for Grant Applications (SGAs).
OMB Number: 1205-0NEW.
Frequency: On occasion and annually.
Affected Public: State, Local or Tribal Government; Business or other for-profit; Not-for-profit institutions.
Type of Response: Reporting.
Number of Respondents: 50.
Annual Responses: 5,750.
Average Response time: 20.75 hours.
Total Annual Burden Hours: 119,312.
Total Annualized Capital/Startup Costs: \$2,836,058.
Total Annual Costs (operating/maintaining systems or purchasing services): 0.
Description: Requesting approval for a generic Solicitation for Grant Application form for information collection requirements for SGAs that extend beyond what is collected on currently approved standard forms. OMB approval of this generic SGA form will assist the Department to carry out its responsibilities under the Paperwork Reduction Act by accurately accounting for the public burden associated with grant applications through promoting a common structure for reporting the