Lakeview Interagency Office, 1301 South G Street, Lakeview, OR 97630, (541) 947–6107, or *ptalbott@or.blm.gov* and/or from the following Web site *http://www.or.blm.gov/SEOR-RAC.*

Dated: March 30, 2005.

Shirley Gammon,

Manager, Lakeview District.

[FR Doc. 05–6677 Filed 4–4–05; 8:45 am]

BILLING CODE 4310–33–P

DEPARTMENT OF THE INTERIOR

Bureau of Land Management

[CO–922–05–1310–FI; COC63406]

Notice of Proposed Reinstatement of Terminated Oil and Gas Lease COC63406

AGENCY: Bureau of Land Management; Interior.

ACTION: Notice of proposed reinstatement of terminated oil and gas lease.

SUMMARY: Pursuant to the provisions of 30 U.S.C. 188(d) and (e), and 43 CFR 3108.2–3(a) and (b)(1), a petition for reinstatement of oil and gas lease COC63406 for lands in Delta County, Colorado, was timely filed and was accompanied by all the required rentals accruing from the date of termination.

FOR FURTHER INFORMATION CONTACT: Bureau of Land Management, Beverly A. Derringer, Chief, Fluid Minerals Adjudication, at 303–239–3765.

SUPPLEMENTARY INFORMATION: The lessee has agreed to the amended lease terms for rentals and royalties at rates of $5.00 per acre, or fraction thereof, per year and 16⅔ percent, respectively. The lessee has paid the required $500 administrative fee and $155 to reimburse the Department for the cost of this **Federal Register** notice. The lessee has met all the requirements for reinstatement of the lease as set out in Section 31 (d) and (e) of the Mineral Lands Leasing Act of 1920 (30 U.S.C. 188), and the Bureau of Land Management is proposing to reinstate lease COC63406 effective August 1, 2004, subject to the original terms and conditions of the lease and the increased rental and royalty rates cited above.

Dated: February 8, 2005.

Beverly A. Derringer,

Chief, Fluid Minerals Adjudication.

[FR Doc. 05–6691 Filed 4–4–05; 8:45 am]

BILLING CODE 4310–JB–M≤

DEPARTMENT OF THE INTERIOR

Bureau of Reclamation

Prospective Grant of Exclusive Patent License

AGENCY: Bureau of Reclamation, Interior.

ACTION: Notice.

SUMMARY: This notice is issued in accordance with 35 U.S.C. 209(c)(1) and 37 CFR 404.7(a)(1)(I). The Bureau of Reclamation (Reclamation) is contemplating the granting of an exclusive license in the United States to practice the invention embodied in U.S. Patent No. 6,466,009 B1, entitled, ''Flexible Printed Circuit Magnetic Flux Probe.'' The exclusive license is to be granted to Iris Power Engineering having a place of business in Toronto, Canada. Iris Power Engineering is owned by Koch Industries, Inc. of Wichita, Kansas. The patent rights in this invention have been assigned to the United States of America.

The prospective exclusive license will be royalty-bearing and will comply with the terms and conditions of 35 U.S.C. 209 and 37 CFR 404.7. While the primary purpose of this notice is to announce Reclamation's intent to grant an exclusive license to practice the above listed patent, it also serves to publish the availability of this patent for licensing in accordance with law. The prospective license may be granted unless Reclamation receives written evidence and argument which establish that the grant of the license would not be consistent with the requirements of 35 U.S.C. 209 and 37 CFR 404.7.

DATES: Written evidence and arguments against granting the prospective license must be received by fifteen (15) days from the date of this notice.

ADDRESSES: Inquiries, comments, and other materials relating to the contemplated license may be submitted to Chuck Hennig, Research Coordinator, Bureau of Reclamation, Office of the Research Director, D–9000, P.O. Box 25007, Denver, CO 80225–0007.

A copy of the above identified patent may be purchased from the U.S. Patent and Trademark Office, by calling (703) 308–9726 or (800) 972–6382 or downloaded free of charge from the U.S. Patent and Trademark Office Web site at *http://www.uspto.gov.*

FOR FURTHER INFORMATION CONTACT: Chuck Hennig (*chennig@do.usbr.gov*), Research Coordinator, at 303–445–2134 or Siegie Potthoff (*spotthoff@do.usbr.gov*), Program Administrator, at 303–445–2136.

SUPPLEMENTARY INFORMATION: The invention relates to a flexible magnetic flux probe useful in the field of rotary generator and motor condition monitoring. Additional R&D is required to develop a commercially viable version of the device. In addition, a significant investment must be made to develop a market niche for the device.

Competing applications completed and received by Reclamation in response to this notice within fifteen (15) days of this posting will be considered as objections to the grant of the contemplated license. Application forms are available from the Office of the Research Director, Bureau of Reclamation at the address above.

Reclamation's 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. There may be circumstances in which we would withhold a respondent's identity from public disclosure, as 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.

Dated: March 2, 2005.

Charles Hennig,

Research Coordinator, Denver Office.

[FR Doc. 05–6711 Filed 4–4–05; 8:45 am]

BILLING CODE 4310–MN–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated December 21, 2004, and published in the **Federal Register** on January 4, 2005, (70 FR 388–389), Cambrex Charles City, Inc., 1205 11th Street, Charles City, Iowa 50616, made application by renewal and on October 13, 2004 by letter 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
Amphetamine (1100)	II
Methylphenidate (1724)	II
Dextropropoxyphene (9273)	II

Drug	Schedule
Fentanyl (9801)	II

The company plans to manufacture the listed controlled substances in bulk 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 determined that the registration of Cambrex Charles City, Inc. to manufacture the listed basic classes of controlled substances is consistent with the public interest at this time. DEA has investigated Cambrex Charles City, 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(a), the above named company is granted registration as a bulk manufacturer of the basic classes of controlled substances listed.

Dated: March 25, 2005.

William J. Walker,

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

[FR Doc. 05–6701 Filed 4–4–05; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated December 21, 2004, and published in the **Federal Register** on January 4, 2005, (70 FR 391), Eli-Elsohly Laboratories, Inc., Mahmoud A. Elsohly PhD., 5 Industrial Park Drive, Oxford, Mississippi 38655, made application by letter to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of Thebaine (9333), a basic class of controlled substance listed in Schedule II.

The company plans to manufacture the listed controlled substance in bulk for use in analysis and drug test standards.

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 Eli-Elsohly Laboratories, Inc. to manufacture the listed basic class of controlled substance is consistent with the public interest at this time. DEA has investigated Eli-Elsohly Laboratories, 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(a), the above named company is granted registration as a bulk manufacturer of the basic class of controlled substance listed.

Dated: March 25, 2005.

William J. Walker,

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

[FR Doc. 05–6699 Filed 4–4–05; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importation of Controlled Substances; Notice of Application

Pursuant to 21 U.S.C. 958(1)), 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)(2)(b) authorizing the importation of such substances, provide manufacturers holding registrations for the bulk manufacture of the substances an opportunity for a hearing.

Therefore, in accordance with 21 CFR 1301.34(a), this is notice that on May 20, 1998, Ethical Nutritional, LLC, 176 University Parkway, Pomona, California 91768–4300, made application to the Drug Enforcement Administration (DEA) for registration as an importer of Coca Leaves (9040), a basic class of controlled substances listed in Schedule II.

The company plans to import small quantities of the listed controlled substance to manufacture homeopathic medications for human consumption.

Any manufacturer who is presently, or is applying to be, registered with DEA to manufacture such basic classes of controlled substances may file written 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 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 5, 2005.

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 the basic classes of controlled substances listed 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: March 25, 2005.

William J. Walker,

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

[FR Doc. 05–6700 Filed 4–4–05; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated December 21, 2004, and published in the **Federal Register** on January 4, 2005, (70 FR 391), Houba, Inc., P.O. Box 190, 16235 State Road 17, Culver, Indiana 46511, 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; and by letter dated October 1, 2004, to modify its name to Acura Pharmaceutical Technologies, Inc., and change the address by removing the P.O. Box 190.

Drug	Schedule
Oxycodone (9143)	II
Hydrocodone (9193)	II

The company plans to manufacture the listed controlled substances in bulk for distribution to its customers.