considered to have been waived, and the contract will then be awarded to the party that has submitted the best responsive offer.

The Secretary will consider and evaluate all offers received as a result of this notice. Any offer, including that of the previous concessioner, must be received by the Superintendent, Glacier National Park, West Glacier, Montana 59936, no later than sixty (60) days following release of the prospectus to be considered and evaluated.

Dated: October 2, 1998.

Michael D. Snyder,

Deputy Regional Director, Rocky Mountain Intermountain Region.

[FR Doc. 98–27226 Filed 10–8–98; 8:45 am]

BILLING CODE 4310–70–P

DEPARTMENT OF THE INTERIOR

National Park Service

National Preservation Technology and Training Board: Meeting

AGENCY: National Park Service, Department of the Interior.

ACTION: Notice of meeting of the National Preservation Technology and Training Board.

Notice is hereby given in accordance with the Federal Advisory Committee Act, 5 U.S.C. Appendix (1988), that the National Preservation Technology and Training Board will meet on November 2, 3, and 4, 1998, in Natchitoches, Louisiana.

The Board was established by Congress to provide leadership, policy advice, and professional oversight to the National Center for Preservation Technology and Training, as required under the National Historic Preservation Act of 1966, as amended (16 U.S.C. 470).

The Board will meet on the campus of Northwestern State University of Louisiana in the Board Room of the Louisiana School for Math, Science and the Arts at 715 College Street, Natchitoches, Louisiana. Matters to be discussed will include, officer and committee reports; Northwestern University report; staff program updates; the establishment of non-Federal support for the Center's programs; budget review; grant program, cooperating organizations, task force reports on NCPTT development and systems, and Millenium projects.

Monday, November 3 the meeting will start at 10 a.m. and end at 5 p.m. On Tuesday, November 4 the meeting will start at 8:30 a.m. and end at 5 p.m. On Wednesday, November 5, the meeting will be begin at 8:30 a.m. and end at 11:30 a.m. Meetings will be open to the public. However, facilities and space for accommodating members of the public are limited and persons will be accommodated on a first-come, first-served basis. Any member of the public may file a written statement concerning the matters to be discussed with Dr. Elizabeth A. Lyon, Chair, National Preservation Technology and Training Board, PO Box 1269, Flowery Branch, Georgia 30542.

Persons wishing more information concerning this meeting, or who wish to submit written statements, may do so by contacting Mr. E. Blaine Cliver, Chief, HABS/HAER, National Park Service, 1849 C Street NW, Washington, DC 20240, telephone: (202) 343–9573. Draft summary minutes of the meeting will be available for public inspection about eight weeks after the meeting at the office of the Preservation Assistance Division, Suite 200, 800 North Capitol Street, Washington, DC.

Dated: October 5, 1998.

E. Blaine Cliver,

Chief, HABS/HAER, Designated Federal Official, National Park Service.

[FR Doc. 98–27275 Filed 10–8–98; 8:45 am]

BILLING CODE 4310–70–P

DEPARTMENT OF THE INTERIOR

Bureau of Reclamation

Meeting of the Conservation Advisory Group, Yakima River Basin Water Enhancement Project, Yakima, WA

AGENCY: Bureau of Reclamation, Department of the Interior.

ACTION: Notice of meeting.

SUMMARY: As required by the Federal Advisory Committee Act, notice is hereby given that the Conservation Advisory Group, Yakima River Basin Water Enhancement Project, Yakima, Washington established by the Secretary of the Interior, will hold a public meeting. The purpose of the Conservation Advisory Group is to provide technical advice and counsel to the Secretary and the State on the structure, implementation, and oversight of the Yakima River Basin Water Conservation Program.

DATES: Thursday, October 15, 1998, 9 a.m.–4 p.m.

ADDRESSES: Bureau of Reclamation Office, 1917 Marsh Road, Yakima, Washington.

FOR FURTHER INFORMATION CONTACT: James Esget, Manager, Yakima River Basin Water Enhancement Project, P.O. Box 1749, Yakima, Washington 98907; (509) 575–5848, extension 267.

SUPPLEMENTARY INFORMATION: The purpose of the meeting will be to review the Bureau of Reclamation's water acquisition process and procedures and develop recommendations on the process to facilitate voluntary sale or lease of water. Progress Reports will be provided on the Basin Conservation Plan and the Yakima River Basin Wetlands and Floodplain Habitat Plan.

Dated: October 1, 1998.

Loren Kjeldgaard,

Acting Area Manager.

[FR Doc. 98–27123 Filed 10–8–98; 8:45 am]

BILLING CODE 4310–94–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Application

Pursuant to Section 1301.33(a) of Title 21 of the Code of Federal Regulations (CFR), this is notice that on August 5, 1998, Ansys Diagnostics, Inc., 25200 Commercentre Drive, Lake Forest, California 92630, made application by renewal to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of the basic classes of controlled substances listed below:

Drug	Schedule
Phencyclidine (7471)	II
1-Piperidinocyclohexanecarbonitrile (PCC) (8603).	II
Benzoylecgonine (9180)	II

The firm plans to manufacture the listed controlled substances to produce standards and controls for in-vitro diagnostic drug testing systems.

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.

Any such comments or objections may be addressed, in quintuplicate, to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, United States Department of Justice, Washington, DC 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than December 8, 1998.

Dated: October 1, 1998.

John H. King,

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

[FR Doc. 98–27100 Filed 10–8–98; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importation of Controlled Substances; Notice of Application

Pursuant to Section 1008 of the Controlled Substances Import and Export Act (21 U.S.C. 958(i)), 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 Section 1002(a) authorizing the importation of such a substance, provide manufacturers holding registrations for the bulk manufacture of the substance an opportunity for a hearing.

Therefore, in accordance with Section 1301.34 of Title 21, Code of Federal Regulations (CFR), notice is hereby given that on June 3, 1998, Calbiochem-Novabiochem Corporation, 10394 Pacific Center Court, Attn: Receiving Inspector, San Diego, California 92121–4340, made application to the Drug Enforcement Administration to be registered as an importer of the basic classes of controlled substances listed below:

Drug	Schedule
Tetrahydrocannabinols (7370)	I
Mescaline (7381)	I
Phencyclidine (7471)	II
Phenylacetone (8501)	II
Cocaine (9041)	II

The firm plans to import small quantities of the listed controlled substances to make reagents for distribution to the biomedical research community.

Any manufacturer holding, or applying for, registration as a bulk manufacturer of this basic classes of controlled substances may file written comments on or objections to the application described above and may, at the same time, file a written request for a hearing on such application in accordance with 21 CFR 1301.43 in such form as prescribed by 21 CFR 1316.47.

Any such comments, objections or requests for a hearing may be addressed, in quintuplicate, to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, United States Department of Justice, Washington, D.C. 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than November 9, 1998.

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 at 40 FR 43745–46 (September 23, 1975), all applicants for registration to import basic classes 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 (a), (b), (c), (d), (e), and (f) are satisfied.

Dated: October 1, 1998.

John H. King,

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

[FR Doc. 98–27101 Filed 10–8–98; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importation of Controlled Substances; Notice of Application

Pursuant to Section 1008 of the Controlled Substances Import and Export Act (21 U.S.C. 958(i)), 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 Section 1002(a) authorizing the importation of such a substance, provide manufacturers holding registrations for the bulk manufacture of the substance an opportunity for a hearing.

Therefore, in accordance with Section 1301.34 of Title 21, Code of Federal Regulations (CFR), notice is hereby given that on August 11, 1998, Chiragene, Inc., 7 Powder Horn Drive, Warren, New Jersey 07059, made application to the Drug Enforcement Administration to be registered as an importer of the basic classes of controlled substances listed below:

Drug	Schedule
2,5-Dimethoxyamphetamine (7396).	I
Phenylacetone (8501)	II

The firm plans to import the phenylacetone to manufacture amphetamine and the 2,5-dimethoxyamphetamine for distribution.

Any manufacturer holding, or applying for, registration as a bulk manufacturer of this basic classes of controlled substances may file written comments on or objections to the application described above and may, at the same time, file a written request for a hearing on such application in accordance with 21 CFR 1301.43 in such form as prescribed by 21 CFR 1316.47.

Any such comments, objections or requests for a hearing may be addressed, in quintuplicate, to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, United States Department of Justice, Washington, D.C. 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than November 9, 1998.

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 at 40 FR 43745–46 (September 23, 1975), all applicants for registration to import basic classes 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(a), (b), (c), (d), (e), and (f) are satisfied.

Dated: October 1, 1998.

John H. King,

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

[FR Doc. 98–27102 Filed 10–8–98; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Application

Pursuant to Section 1301.33(a) of Title 21 of the Code of Federal Regulations (CFR), this is notice that on July 20, 1998, Hoffmann-LaRoche, Inc., 240 Kingsland Street, Nutley, New Jersey 07110, made application by renewal to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of levorphanol (9220), a