to pay, it would lead to ability-to-pay studies for project use power contractors.

Comment: Since Reclamation intends to maintain the effective project use power rate at 2.5 mils/kWh, through application of the ability-to-pay test, what is the purpose of Reclamation's proposed rate adjustment?

Response: While present economic conditions create depressed agriculture and the majority of the irrigators will pay 2.5 mils/kWh, the adjusted rate will allow Reclamation to capture additional revenues if and when the economics of agriculture improve. It will also provide current rate structure for use in decisions and legislation related to proposed new projects. Reclamation is required to have accurate numbers for operations, maintenance, and replacement costs.

NEPA

In compliance with the National Environmental Policy Act of 1969 (NEPA), 43 U.S.C. 4321 *et seq.* Council on Environmental Quality Regulations (40 CFR parts 1500–1508); and the Department of Energy's NEPA Implementing Procedures (10 CFR part 1021), Reclamation has determined that this action is categorically excluded from the preparation of an Environmental Assessment or Environmental Impact Statement.

Power Rate Schedules

The existing rate schedule MRB–P10 placed into effect on November 1, 1986, will be replaced by rate schedule MRB–P11. Rate Schedule MRB–P11 is as follows: *Effective:* 30 days after being published in the **Federal Register**. *Location:* In the areas generally described as central and eastern Montana, North and South Dakota, Nebraska, eastern Colorado, Wyoming, Kansas, western Iowa, and western Minnesota.

Applicable

For use in the operation of congressionally authorized irrigation and drainage pumping plants on irrigation projects for power service supplied through metering at specified points of delivery.

Character and Conditions of Service

Alternating current, 60 hertz, three phase, delivered and metered at the point identified in the contract upon demand during the summer irrigation season.

Availability

Available at 60 hertz at the pumping plant upon demand during the summer irrigation season.

Monthly Rate

Demand Charge: None.

Energy Charge: 10.76 mils/kWh for all energy use; subject to ability-to-pay but not less than 2.5 mils/kWh.

Seasonal Minimum Bill: $2.75 per kilowatt of the maximum 30-minute integrated demand established during service months of each year specified in the contract.

Adjustments

For Power Factor: The customer will normally be required to maintain a power factor at a point of delivery of not less than 95 percent lagging or leading.

Approval of Project Use Power Rate by the Commissioner, Bureau of Reclamation

The Commissioner approved the rate of 10.76 mils/kWh by memorandum dated January 15, 2002.

Dated: January 23, 2002.

Gerald W. Kelso,

Assistant Regional Director.

[FR Doc. 02–4025 Filed 2–19–02; 8:45 am]

BILLING CODE 4310–MN–P

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 March 7, 2001, Cedarburg Pharmaceutical LLC, 870 Badger Circle, Grafton, Wisconsin 53024, made application to the Drug Enforcement Administration to be registered as an importer of the basic classes of controlled substances listed below:

Drug	Schedule
Codeine (9050)	II
Morphine (9300)	II
Thebaine (9333)	II

The firm plans to import the listed controlled substances to repackage and sell as bulk controlled substances, and also use as starting materials to manufacture bulk active pharmaceuticals ingredients.

Any manufacturer holding, or applying for, registration as a bulk manufacturer of these 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, DC 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than (30 Days from publication).

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: February 6, 2002.

Laura M. Nagel,

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

[FR Doc. 02–4057 Filed 2–19–02; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated August 9, 2001, and published in the **Federal Register** on August 10, 2001, (66 FR 42240), Roche Diagnostics Corporation, Attn:

Regulatory Compliance, 9115 Hague Road, Indianapolis, Indiana 46250, 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 below:

Drug	Schedule
Lysergic acid diethylamide (7315)	I
Tetrahydrocannabinols (7370)	I
Phencyclidine (7471)	II
Benzoylecgonine (9180)	II
Methadone (9250)	II
Morphine (9300)	II

Roche Diagnostics Corporation plans to manufacture small quantities of the above listed controlled substances for incorporation in drug of abuse detection kits.

No comments or objections have been received. DEA has considered the factors in Title 21, United States Code, section 823(a) and determined that the registration of Roche Diagnostics Corporation to manufacture the listed controlled substances is consistent with the public interest at this time. DEA has investigated Roche Diagnostics Corporation to ensure that the company's continued registration is consistent with the public interest. These investigations have included inspection and testing of the company's physical security systems, audits of the company's records, 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. section 823 and 28 CFR secs. 0.100 and 0.104, the Deputy Assistant Administrator, Office of Diversion Control, hereby orders that the application submitted by the above firm for registration as a bulk manufacturer of the basic classes of controlled substances listed above is granted.

Dated: February 1, 2002.

Laura M. Nagel,

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

[FR Doc. 02–4056 Filed 2–19–02; 8:45 am]

BILLING CODE 4410–09–M

OVERSEAS PRIVATE INVESTMENT CORPORATION

Submission for OMB Review; Comment Request

AGENCY: Overseas Private Investment Corporation

ACTION: Request for comments.

SUMMARY: Under the provisions of the Paperwork Reduction Act (44 U.S.C. chapter 35), agencies are required to publish a Notice in the **Federal Register** notifying the public that the Agency is preparing an information collection request for OMB review and approval and to request public review and comment on the submission. OPIC published its first Federal Register Notice on this information collection request on December 12, 2002, in 66 FR 239, p. 64312, at which time a 60-calendar day comment period was announced. This comment period ended February 11, 2002. No comments were received in response to this notice.

This information collection submission has now been submitted to OMB for review. Comments are again being solicited on the need for the information, its practical utility, the accuracy of the Agency's burden estimate, and on ways to minimize the reporting burden, including automated collection techniques and uses of other forms of technology. The proposed form under review is summarized below.

DATES: Comments must be received on or before March 22, 2002.

ADDRESSES: Copies of the subject form and the request for review submitted to OMB may be obtained from the Agency Submitting Officer. Comments on the form should be submitted to the OMB Reviewer.

FOR FURTHER INFORMATION CONTACT:

OPIC Agency Submitting Officer

Carol Brock, Records Manager, Overseas Private Investment Corporation, 1100 New York Avenue, NW., Washington, DC 20527; 202/336–8563.

OMB Reviewer

David Rostker, Office of Information and Regulatory Affairs, Office of Management and Budget, New Executive Office Building, Docket Library, Room 10102, 725 17th Street, NW., Washington, DC 20503, 202/395–3897.

Summary of Form Under Review

Type of Request: Form Amendment.

Title: Application for Political Risk Investment Insurance.

Form Number: OPIC–52.

Frequency of Use: Once per investor, per project.

Type of Respondents: Business or other institutions.

Standard Industrial Classification Codes: All.

Description of Affected Public: U.S. companies investing overseas.

Reporting Hours: 6½ hours per project.

Number of Responses: 150 per year.

Federal Cost: $24,300 per year.

Authority for Information Collection: Sections 231 and 234(a) of the Foreign Assistance Act of 1961, as amended.

Abstract (Needs and Uses): The OPIC 52 form is the principal document used by OPIC to determine the investor's and the project's eligibility, assess the environmental impact and developmental effects of the project, measure the economic effects for the United States and the host country economy, and collect information for underwriting analysis.

Dated: February 13, 2002.

Rumu Sarkar,

Assistant General Counsel, Administrative Affairs, Department of Legal Affairs.

[FR Doc. 02–4018 Filed 2–19–02; 8:45 am]

BILLING CODE 3210–01–M

RAILROAD RETIREMENT BOARD

Proposed Data Collection; Comment Request

SUMMARY: In accordance with the requirement of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995 which provides opportunity for public comment on new or revised data collections, the Railroad Retirement Board will publish periodic summaries of proposed data collections.

Comments are invited on: (a) Whether the proposed information collection is necessary for the proper performance of the functions of the agency, including whether the information has practical utility; (b) the accuracy of the RRB's estimate of the burden of the collection of the information: (c) ways to enhance the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden related to the collection of information on respondents, including the use of automated collection techniques or other forms of information technology.

Title and Purpose of information collection: Statement Regarding Contributions and Support: OMB 3220–0099 Under section 2 of the Railroad Retirement Act, dependency on an employee for one-half support at the time of an employee's death can be a condition affecting entitlement to a survivor annuity and can affect the amount of both spouse and survivor annuities. One-half support is also a condition which may negate the public service pension offset in Tier I for a spouse or widow(er). The Railroad Retirement Board (RRB) utilizes Form G–134, Statement Regarding Contributions and Support, to secure