controlled substance listed in schedule

The company plans to import the listed controlled substance for the manufacture of a bulk controlled substance for distribution to its customer.

As noted in a previous notice published in the **Federal Register** on September 23, 1975, (40 FR 43745), all applicants for registration to import a basic class 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(b), (c), (d), (e), and (f) are satisfied.

Dated: March 27, 2008.

## Joseph T. Rannazzisi,

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

[FR Doc. E8-6774 Filed 4-1-08; 8:45 am]

#### **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 March 4, 2008, Lonza Riverside, 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 basic classes of controlled substances listed in schedules I and II:

Drug			Schedule
Gamma (2010).	hydroxybutyric	acid	I
Amphetamine (1100) Methylphenidate (1724)			II II

The company plans to manufacture bulk products for finished dosage units and distribution 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 being sent via regular mail should be addressed, in quintuplicate, to the Drug Enforcement Administration, Office of Diversion

Control, Federal Register Representative (ODL), Washington, DC 20537, or any being sent via express mail should be sent to Drug Enforcement Administration, Office of Diversion Control, Federal Register Representative (ODL), 8701 Morrissette Drive, Springfield, Virginia 22152; and must be filed no later than June 2, 2008.

Dated: March 27, 2008.

## Joseph T. Rannazzisi,

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

[FR Doc. E8–6775 Filed 4–1–08; 8:45 am]
BILLING CODE 4410–09–P

## **DEPARTMENT OF LABOR**

# **Employee Benefits Security Administration**

Proposed Extension of Information Collection Request Submitted for Public Comment; Consent To Receive Employee Benefit Plan Disclosure Electronically

**AGENCY:** Employee Benefits Security Administration, Department of Labor. **ACTION:** Notice.

**SUMMARY:** The Department of Labor, as part of its continuing effort to reduce paperwork and respondent burden, conducts a preclearance consultation program to provide the general public and Federal agencies with an opportunity to comment on proposed and continuing collections of information in accordance with the Paperwork Reduction Act of 1995 (PRA 95). This program helps to ensure that requested data can be provided in the desired format, reporting burden (time and financial resources) is minimized, collection instruments are clearly understood, and the impact of collection requirements on respondents can be properly assessed. Currently, the Employee Benefits Security Administration is soliciting comments on the proposed extension of an information collection request (ICR) incorporated in the Final Rules relating to the use of electronic communication and recordkeeping technologies by employee pension and welfare benefit plans (29 CFR 2520.104b-1).

A copy of the information collection request (ICR) can be obtained by contacting the individual shown in the **ADDRESSES** section of this notice or at <a href="http://www.RegInfo.gov">http://www.RegInfo.gov</a>.

**DATES:** Written comments must be submitted to the office shown in the **ADDRESSES** section on or before June 2, 2008.

ADDRESSES: G. Christopher Cosby, Department of Labor, Employee Benefits Security Administration, 200 Constitution Avenue NW., Washington, DC 20210, (202) 693–8410, FAX (202) 693–4745 (these are not toll-free numbers).

## SUPPLEMENTARY INFORMATION:

# I. Background

The Department established a safe harbor pursuant to which all pension and welfare benefit plans covered by Title I of ERISA may use electronic media to satisfy disclosure obligations under Title I of ERISA (29 CFR 2520.104b-1). Employee benefit plan administrators will be deemed to satisfy their disclosure obligations when furnishing documents electronically only if a participant who does not have access to the employer's electronic information system in the normal course of his duties, or a beneficiary or other person entitled to documents, has affirmatively consented to receive disclosure documents. Prior to consenting, the participant or beneficiary must also be provided with a clear and conspicuous statement indicating the types of documents to which the consent would apply, that consent may be withdrawn at any time, procedures for withdrawing consent and updating necessary information, the right to obtain a paper copy, and any hardware and software requirements. In the event of a hardware or software change that creates a material risk that the individual will be unable to access or retain documents that were the subject of the initial consent, the individual must be provided with information concerning the revised hardware or software, and an opportunity to withdraw a prior consent.

#### **II. Review Focus**

The Department of Labor (Department) is particularly interested in comments that:

- 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