1311.42(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 1311.42(a), (b), (c), (d), (e), and (f) are satisfied.

Dated: July 5, 1995.

#### Gene R. Haislip,

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

 $[FR\ Doc.\ 95{-}17119\ Filed\ 7{-}12{-}95;\ 8{:}45\ am]$ 

BILLING CODE 4410-09-M

### Importer of Controlled Substances; Notice of Registration

By Notice dated May 18, 1995, and published in the **Federal Register** on May 25, 1995 (60 FR 27788), Calbiochem-Novabiochem Corporation, 10394 Pacific Center Court, Attn: Receiving Inspector, San Diego, California 92121–4340, made application to the Drug Enforcement Administration (DEA) to be registered as an importer of the basic classes of controlled substances listed below:

Drug	Schedule
Tetrahydrocannabinols (7370)  Mescaline (7381)  Amphetamine (1100)  Phencyclidine (7471)  Phenylacetone (8501)  Cocaine (9041)	    

A registered manufacturer did file a written request for a hearing with respect to Amphetamine. Therefore, pursuant to Section 1008(a) of the Controlled Substances Import and Export Act and in accordance with Title 21, Code of Federal Regulations, § 1311.42, the above firms is granted registration as an importer of the basic classes of controlled substances listed above with the exception of Amphetamine (1100).

Dated: July 5, 1995.

#### Gene R. Haislip,

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

[FR Doc. 95–17120 Filed 7–12–95; 8:45 am] BILLING CODE 4410–09–M

# Importer of Controlled Substances; Notice of Registration

By Notice dated May 17, 1995, and published in the **Federal Register** on May 25, 1995, (60 FR 27789), Research Biochemicals, Limited Partnership, One Strathmore Road, Natick, Massachusetts 01760, made application to the Drug Enforcement Administration (DEA) to be registered as an importer of the basic classes of controlled substances listed below:

Drug	Schedule
Methaqualone (2565)	

No comments or objections have been received. Therefore, pursuant to Section 1008(a) of the Controlled Substances Import and Export Act and in accordance with Title 21, Code of Federal Regulations, § 1311.42, the above firm is granted registration as an importer of the basic classes of controlled substances listed above.

Dated: July 5, 1995.

## Gene R. Haislip,

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

[FR Doc. 95–17121 Filed 7–12–95; 8:45 am] BILLING CODE 4410–09–M

#### **DEPARTMENT OF LABOR**

#### Office of the Secretary

# Commission on Family and Medical Leave; Notice of Public Hearing

**AGENCY:** Office of the Secretary, Labor. **ACTION:** Notice of public hearing.

SUMMARY: Pursuant to Title III of the Family Medical Leave Act (FMLA) of 1993 (Pub. L. 103–3) this is to announce a hearing on the experience of family and temporary medical leave policies for the Commission which is to take place on Friday, August 4, 1995. The purpose of the Commission is to, among other things, study the effects of existing and proposed policies relating to family and medical leave. The Commission has the practical task of conducting a comprehensive study of: (a) Existing and proposed mandatory and voluntary

policies relating to family and temporary medical leave, including policies provided by employers not covered under the act; (b) the potential costs, benefits, and impact on productivity, job creation and business growth of such policies on employers and employees; (c) possible differences in costs, benefits, impact on productivity, job creation and business growth of such policies on employers based on business type and size; (d) the impact of family and medical leave policies on the availability of employee benefits provided by employers, including employers not covered under this Act; (e) alternative and equivalent State enforcement of Title I with respect to employees described in section 108(a); (f) methods used by employers to reduce administrative costs of implementing family and medical leave policies; (g) the ability of the employers to recover, under section 104(c)(2), the premiums described in such section; and (h) the impact on employers and employees of policies that provided temporary wage replacement during periods of family and medical leave.

TIME AND PLACE: The hearing will be held on Friday, August 4, 1995, from 9 am until 12 pm, at the Department of Commerce, 14th Street and Constitution Avenue, NW, Washington, DC 20230, in the Departmental Auditorium.

**AGENDA:** The agenda for the hearing is as follows: Three panels of witnesses will give testimony on their experiences with family and temporary medical leave policies.

**STATEMENTS:** Interested persons may submit, in writing, data, information or views on employer or employee experiences with family and temporary medical leave policies prior to or at the hearing.

PUBLIC PARTICIPATION: The hearing will be open to the public. Seating will be available on a first-come, first-served basis. Seats will be reserved for the media. Persons with disabilities should contact the Commission no later than July 28, 1995, if special accommodations are needed.

#### FOR FURTHER INFORMATION CONTACT:

Susan King, Executive Director, Commission on Leave, U.S. Department of Labor, 200 Constitution Avenue, NW, Room S–3002, Washington, DC 20210, telephone: (202) 219–4526, Ext. 102.

Signed at Washington, DC, this 6th day of July, 1995.

#### Susan King,

Executive Director Commission on Leave. [FR Doc. 95–17206 Filed 7–12–95; 8:45 am] BILLING CODE 4510–23–M