Dated: August 1, 2002.

Nancy E. Cheal,

Acting Associate Director for Policy, Planning and Evaluation, Centers for Disease Control and Prevention.

[FR Doc. 02–20130 Filed 8–8–02; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[Program Announcement 00100]

Disease, Disability, and Injury Prevention and Control; Special Emphasis Panel: Community Based Strategies To Increase HIV Testing of Persons at High Risk in Communities of Color

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92–463), the Centers for Disease Control and Prevention (CDC) announces the following meeting:

Name: Disease, Disability, and Injury Prevention and Control Special Emphasis Panel (SEP): Community Based Strategies to Increase HIV Testing of Persons At High Risk in Communities of Color, PA# 00100.

Times and Dates: 9 a.m.–9:30 a.m., August, 26, 2002 (Open), 9:30 a.m.–4:30 p.m., August 26, 2002 (Closed), 9 a.m.–4:30 p.m., August 27–28, 2002 (Closed).

Place: Atlanta Marriott Century Center, 2000 Century Boulevard, NE., Atlanta, GA 30345.

Status: Portions of the meeting will be closed to the public in accordance with provisions set forth in section 552b(c) (4) and (6), Title 5 U.S.C., and the Determination of the Director, Management Analysis and Services Office, CDC, pursuant to Public Law 92–463.

Matters To Be Discussed: The meeting will include the review, discussion, and evaluation of applications received in response to PA# 00100.

Contact Person for More Information: Beth Wolfe, Prevention Support Office, National Center for HIV, STD, and TB Prevention, CDC, 1600 Clifton Road, NE., MS E–07, Atlanta, GA 30333, 404–639–8025.

The Director, Management Analysis and Services Office has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities, for both the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

Joseph E. Salter,

Acting Director, Management Analysis and Services Office, Centers for Disease Control and Prevention (CDC).

[FR Doc. 02–20296 Filed 8–7–02; 10:52 am]

BILLING CODE 4163–18–P

DEPARTMENT OF LABOR

Employment and Training Administration

[TA–W–38,307 and TA–W–38,307A]

Progress Lighting, Cowpens, SC; Progress Lighting, Philadelphia, PA; Amended Certification Regarding Eligibility To Apply for Worker Adjustment Assistance

In accordance with section 223 of the Trade Act of 1974 (19 U.S.C. 2273) the Department of Labor issued a Notice of Certification Regarding Eligibility to Apply for Worker Adjustment Assistance on November 28, 2000, applicable to workers of Progress Lighting, Cowpens, South Carolina. The notice was published in the **Federal Register** on December 21, 2000 (65 FR 80457).

At the request of the International Brotherhood of Electrical Workers, Local 2005, the Department reviewed the certification for workers of the subject firm. The company reports that worker separations occurred at the Philadelphia, Pennsylvania location of Progress Lighting. The Philadelphia, Pennsylvania location is a distribution center for the lighting fixtures produced in Cowpens, South Carolina.

Based on these findings, the Department is amending the certification to include workers of the Philadelphia, Pennsylvania location of Progress Lighting.

The intent of the Department's certification is to include all workers of Progress Lighting who were adversely affected by increased imports.

The amended notice applicable to TA–W–38,307 is hereby issued as follows:

All workers of Progress Lighting, Cowpens, South Carolina (TA–W–38,307) and Progress Lighting, Philadelphia, Pennsylvania (TA–W–38,307A) who became totally or partially separated from employment on or after October 28, 1999 through November 28, 2002, are eligible to apply for adjustment assistance under Section 223 of the Trade Act of 1974.

Signed at Washington, DC, this 25th day of July, 2002.

Edward A. Tomchick,

Director, Division of Trade Adjustment Assistance.

[FR Doc. 02–20192 Filed 8–8–02; 8:45 am]

BILLING CODE 4510–30–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 02N–0208]

Agency Information Collection Activities; Submission for OMB Review; Comment Request; State Enforcement Notifications

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that the proposed collection of information has been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995.

DATES: Submit written comments on the collection of information by September 9, 2002.

ADDRESSES: Submit written comments on the collection of information to the Office of Information and Regulatory Affairs, OMB, New Executive Office Bldg., 725 17th St. NW., rm. 10235, Washington, DC 20503, Attn: Stuart Shapiro, Desk Officer for FDA.

FOR FURTHER INFORMATION CONTACT: Peggy Schlosburg, Office of Information Resources Management (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–1223.

SUPPLEMENTARY INFORMATION: In compliance with 44 U.S.C. 3507, FDA has submitted the following proposed collection of information to OMB for review and clearance.

State Enforcement Notifications—21 CFR 100.2(d) (OMB Control Number 0910–0275)—Extension

Section 310(b) of the Federal Food, Drug, and Cosmetic Act (the act) (21 U.S.C. 337(b)) authorizes States to enforce certain sections of the act in their own names, but provides that States must notify FDA before doing so. Section 100.2(d) (21 CFR 100.2 (d)) sets forth the information that a State must provide to FDA in a letter of notification when it intends to take enforcement action under the act against a particular food located in the State. The information required under § 100.2(d) will enable FDA to identify the food against which the State intends to take action and advise the State whether Federal action has been taken against it. With certain narrow exceptions, Federal enforcement action precludes State action under the act.

FDA estimates the burden of this collection of information as follows: