Dated: June 7, 2007.

James Scanlon,

Deputy Assistant Secretary for Planning and Evaluation (OSDP), Office of the Assistant Secretary for Planning and Evaluation.

[FR Doc. 07–2980 Filed 6–15–07; 8:45 am]

BILLING CODE 4151–05–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers of Disease Control and Prevention

[Docket Number NIOSH 105]

Notice of Request for Public To Submit Comments and Attend Public Meeting

AGENCY: National Institute for Occupational Safety and Health (NIOSH) of the Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

ACTION: Notice.

The National Institute for Occupational Safety and Health (NIOSH) of the Centers for Disease Control and Prevention (CDC) announces the following meeting and request for public comment on the following draft document: ''NIOSH Hazardous Drugs List Update.''

The documents can found at: *http://www.cdc.gov/niosh/topics/hazdrug/2007publicmeeting.*

Public Comment Period: Comments must be postmarked by September 20, 2007. Comments may be submitted electronically to *niocindocket@cdc.gov* or to Diane Miller, Robert A. Taft Laboratories, 4676 Columbia Parkway, MS C–34, Cincinnati, Ohio 45226. Comments should reference the NIOSH *Docket Number 105.*

Meeting Date and Time: August 28, 2007 9 a.m.–5 p.m.

Place: Marriott Metro Center Hotel, 775 12th Street, NW., Washington, DC 20005.

Purpose of Meeting: To discuss and obtain comments on the draft document, ''NIOSH Hazardous Drugs List Update.'' Special emphasis will be placed on discussion of the following:

(1) The appropriateness and relevancy of the NIOSH definition of Hazardous Drugs;

(2) the appropriateness and relevancy of the drugs that fit the NIOSH definition; and

(3) the appropriateness and relevancy of the drugs that do not fit the NIOSH definition.

Status: The forum will include scientists and representatives from various government agencies, industry, labor, and other stakeholders, and is open to the public, limited only by the space available. Persons interested in providing comments need to notify Diane Miller by July 20, 2007. Ms. Miller can be reached by telephone at (513) 533–8450 or by e-mail at *niocindocket@cdc.gov.* Persons wanting to provide comments will be permitted up to 10 minutes, subject to available time. If additional time becomes available, presenters will be notified. Oral comments given at the meeting will be recorded and included in the docket. Written comments will also be accepted at the meeting. NIOSH will use this information to help assess and revise the definition and/or list of Hazardous Drugs.

Contact for Technical Information: Barbara MacKenzie, NIOSH, Robert A. Taft Laboratories, 4676 Columbia Pkwy. MS C–26, Cincinnati, OH 45226, Telephone 513–533–8132, e-mail *hazardousdrugs@cdc.gov.*

Contact Person for Submitting Comments/Meeting Attendance: Diane Miller, Robert A. Taft Laboratories, 4676 Columbia Parkway, MS C–34, Cincinnati, Ohio 45226, telephone (513) 533–8450. All material submitted to the Agency should reference docket number NIOSH 105.

All information received in response to this notice will be available for public examination and copying at the NIOSH Docket Office, 4676 Columbia Parkway, Cincinnati, Ohio 45226.

Dated: June 11, 2007.

James D. Seligman,

Chief Information Officer, Centers for Disease Control and Prevention.

[FR Doc. E7–11680 Filed 6–15–07; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Statement of Organization, Functions and Delegation of Authority

Notice is hereby given that I have redelegated to the TANF Regional Program Managers the following authorities vested in me by the Director, Office of Family Assistance (OFA) in the memoranda dated April 17, 2007.

(a) Authorities Delegated

1. The authority to respond to general inquiries about established state, territorial, and tribal programs, to explain existing program policies, and to suggest referrals to other government agencies and private organizations.

2. The authority to request information from state, territorial, and tribal grantees relative to program policies and operations.

3. The authority to deem state TANF plan amendments as complete.

4. The authority to approve or disapprove state requests, as permitted under section 45 CFR 205.56(d), to utilize alternative data-matching sources for the Income Eligibility Verification System (IEVS).

5. The authority to approve or disapprove state IEVS targeting plans, pursuant to section 45 CFR 205.56(a)(1).

6. The authority to issue letters to state and tribal TANF grantees acknowledging their success in achieving work participation rates for a fiscal year.

7. The authority to advise and notify state TANF grantees concerning their applications for caseload reduction credits.

8. The authority to request caseload and expenditure data from state TANF grantees in order to establish a Tribal Family Assistance Grant (TFAG).

9. The authority to approve, request changes, or defer action on ''Letters of Intent'' submitted by tribal applicants who wish to operate a TANF program.

10. The authority to develop TANF technical assistance plans and proposals for the expenditure of technical assistance funds.

11. The authority to request additional information and consult with states and tribes on modifications to TANF corrective compliance plans submitted in response to a penalty determination.

12. The authority to serve as Approving Officials for audit resolution for state and tribal TANF programs, and for tribal NEW programs and Adult Assistance programs that do not involve cost disallowances.

13. The authority to approve Adult Assistance program plans and plan amendments.

14. The authority to approve Federal Financial Participation (FFP) in payments for repairs to homes owned by recipients of assistance under the Adult Assistance programs.

(b) Limitations

1. This delegation of authority shall be exercised under the Department's existing policies on delegations and regulations.

2. The authority to respond to general inquiries relative to established state, territorial, and tribal programs or to inquire about specific grantee policies and operations requires prior consultation with and concurrence of the Director, Division of State TANF

Policy for state TANF and Adult Assistance topics or the Director, Division of Tribal TANF Management for tribal TANF and NEW topics.

3. The authority to deem state TANF plan amendments complete requires prior consultation with and concurrence of the Director, Division of State TANF Policy.

4. The authority to approve/disapprove under 45 CFR 205.55(d), state applications to use alternate sources of information for income and eligibility (i.e., IEVS), requires prior consultation with and the concurrence of the Office of the Deputy Assistant Secretary for Administration, the Director, Division of State TANF Policy, and with the other programs affected by the request.

5. The authority to approve/disapprove under 45 CFR 205.56(a)(1), IEVS targeting plans, requires prior consultation with and the concurrence of the Office of the Deputy Assistance Secretary for Administration and the Director, Division of State TANF Policy.

6. The authority to issue letters to state and tribal TANF grantees acknowledging their success in achieving work participation rates for a fiscal year requires the concurrence of the Director, Division of State TANF Policy for state TANF grantees or the Director, Division of Tribal TANF Management for tribal TANF grantees.

7. The authority to advise and notify state TANF grantees on their caseload reduction credit applications requires prior consultation with the Director, Division of State TANF policy.

8. The authority to request caseload and expenditure data from state TANF grantees relative to establishing a TFAG requires consultation with and concurrence of the Director, Division of Tribal TANF Management.

9. The authority to approve, request changes, or defer action on ''Letters of Intent'' submitted by tribal applicants who wish to operate TANF programs requires prior consultation with and concurrence of the Director, Division of Tribal TANF Management.

10. The authority to implement TANF technical assistance plans and proposals for the expenditure of technical assistance funds requires approval of the Associate Director, TANF in consultation with the Director, Division of State & Territory Management.

11. The authority to request additional information and consult with states and tribes on modifications to TANF corrective compliance plans requires prior consultation with and concurrence of the Director, Division of State TANF Policy for state TANF plans or the Director, Division of Tribal TANF Management for tribal TANF plans.

12. The authority to approve audit resolutions letters requires prior consultation with and the concurrence of the Regional Office Grants Officer and the Director, Division of State TANF Policy for state TANF and Adult Assistance grantees, or the Director, Division of Tribal TANF Management for tribal TANF and NEW grantees.

13. This delegation of authority does not include the authority to make determinations on state appeals concerning audit questions or recommendations by the Department of Health and Human Services (HHS) Audit Agency which involve ACF program practices reviewed under titles I, X, XI, and XVI of the Social Security Act.

14. The authority to approve Adult Assistance Plans and amendments requires prior consultation with and concurrence of the Director, Division of State TANF Policy.

15. The authority to approve FFP in payments for repairs to homes owned by recipients of Adult Assistance requires prior consultation with and concurrence of the Regional Office Grants Officer and Director, Division of State TANF Policy.

(c) Effective Date

This delegation is effective upon the date of signature.

(d) Effect on Existing Delegations

As related to the authorities delegated herein, this delegation of authority supersedes all previous delegations of authority to the TANF Regional Program Managers.

I hereby affirm and ratify any actions taken by the TANF Regional Office Program Managers, Office of Family Assistance, which involved the exercise of the authorities delegated herein prior to the effective date of this delegation.

Dated: June 8, 2007.

Katherine Bradley,

Associate Director, TANF.

[FR Doc. E7–11707 Filed 6–15–07; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2007N–0225]

Anthrax Vaccines—Bridging Correlates of Protection in Animals to Immunogenicity in Humans; Public Workshop

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of public workshop.

The Food and Drug Administration (FDA) is announcing a public workshop entitled ''Anthrax Vaccines—Bridging Correlates of Protection in Animals to Immunogenicity in Humans.'' The purpose of the public workshop is to discuss possible strategies for bridging animal efficacy data to human immunogenicity data for investigational anthrax vaccines.

Date and Time: The public workshop will be held on November 8, 2007, from 8:30 a.m. to 5 p.m. and November 9, 2007, from 8:30 a.m. to 1 p.m.

Location: Hilton Washington DC North/Gaithersburg, 620 Perry Pkwy., Gaithersburg, MD, 20877, 1–301–977–8900.

Contact Person: Pier Minor, National Institutes of Health/Office of the Director, 6610 Rockledge Dr., Rm. 5WS6, Mail Stop Code 6604, Bethesda, MD 20892, 301–451–6809, FAX: 301–402–0659, e-mail: *minorp@mail.nih.gov*.

Registration: Mail or fax your registration information (including name, title, firm name, address, telephone, and fax number) to the contact person by October 19, 2007. There is no registration fee for the public workshop. Early registration is recommended because seating is limited. There will be no onsite registration.

If you need special accommodations due to a disability, please contact Wenda Minor (see *Contact Person*) at least 7 days in advance.

SUPPLEMENTARY INFORMATION: FDA's Center for Biologics Evaluation and Research, in cooperation with the National Institutes of Health and the Department of Health and Human Services' Office of Biomedical Advanced Research and Development Authority (BARDA) within the Office of the Assistant Secretary for Preparedness and Response, is holding this public workshop. The public workshop will include discussions on: (1) Background information on the ''Animal Rule'' (May 31, 2002, 67 FR 37988), immunological correlates of protection, and the toxin neutralizing assay that will be used to assess anthrax vaccine immunogenicity in animals and humans; (2) animal protection data for anthrax vaccines given both pre- and post-exposure; and (3) human immunogenicity data for anthrax vaccines. The goal of the public workshop is to discuss ways to expedite the development of new anthrax vaccines by providing additional information about bridging animal protection data to human immunogenicity.