VI.2. Administrative and National Policy Requirements

45 CFR Part 74 and Part 92

For more information on the Code of Federal Regulations, see the National Archives and Records Administration at the following Internet address: *http://www.access.gpo.gov/nara/cfr/cfr-table-search.html.*

The following additional requirements apply to this project:

- AR–4 HIV/AIDS Confidentiality Provisions.
- AR–7 Executive Order 12372.
- AR–8 Public Health System Reporting Requirements.
- AR–9 Paperwork Reduction Act Requirements.
- AR–11 Healthy People 2010.
- AR–14 Accounting System Requirements.
- AR–16 Security Clearance Requirement.

Additional information on these requirements can be found on the CDC Web site at the following Internet address: *http://www.cdc.gov/od/pgo/funding/ARs.htm.*

VII. Agency Contacts

For general questions about this announcement, contact: Technical Information Management Section, CDC Procurement and Grants Office, 2920 Brandywine Road, Atlanta, GA 30341, Telephone: 770–488–2700.

For program technical assistance, contact: Ron Sanders, Program Consultant, National Center for HIV, STD and TB Prevention, Division of HIV AIDS Prevention, 1600 Clifton Road, NE Mail stop E–47, Atlanta, GA 30333, Telephone: 404–639–4678, E-mail: *RLS5@cdc.gov.*

For financial, grants management, or budget assistance, contact: Kang Lee, Grants Management Specialist, CDC Procurement and Grants Office, 2920 Brandywine Road, Atlanta, GA 30341, Telephone: 770–488–2733, E-mail: *kil8@cdc.gov.*

Dated: June 28, 2004.

Alan Kotch,

Acting Director, Procurement and Grants Office, Centers for Disease Control and Prevention.

[FR Doc. 04–15070 Filed 7–1–04; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[Program Announcement 05003]

Tuberculosis Elimination and Laboratory; Notice of Availability of Funds—Amendment

A notice announcing the availability of fiscal year (FY) 2004 funds for Tuberculosis Elimination and Laboratory was published in the **Federal Register** on May 27, 2004, Volume 69, Number 103, pages 30300–30312. The notice is amended as follows: On page 30300, Column 1, ''Application Deadline'', change deadline date to July 29, 2004. On page 30308, Column 3, ''Application Deadline'', change deadline date to July 29, 2004.

Dated: June 28, 2004.

Alan Kotch,

Acting Director, Procurement and Grants Office, Centers for Disease Control and Prevention.

[FR Doc. 04–15066 Filed 7–1–04; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Request for Nominations for Nonvoting Members Representing Industry Interests on Public Advisory Panels or Committees

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is requesting nominations for nonvoting industry representatives to serve on certain device panels of the Medical Devices Advisory Committee in the Center for Devices and Radiological Health.

FDA has a special interest in ensuring that women, minority groups, individuals with physical disabilities, and small businesses are adequately represented on its advisory committees. Therefore, the agency encourages nominations for appropriately qualified candidates from these groups.

DATES: Industry organizations interested in participating in the selection of a nonvoting member to represent industry for vacancies listed in this notice must send a letter to FDA by August 2, 2004, stating their interest in one or more panels. Concurrently, nomination materials for prospective candidates should be sent to FDA by August 2, 2004. A nominee may either be self-nominated or nominated by an organization to serve as a nonvoting industry representative.

ADDRESSES: All letters of interest and nominations should be sent to Kathleen L. Walker, Center for Devices and Radiological Health (HFZ–17), Food and Drug Administration, 2098 Gaither Rd., Rockville, MD 20850, 301–594–1283, ext. 114, e-mail: *KLW@CDRH.FDA.GOV*.

FOR FURTHER INFORMATION CONTACT: Kathleen L. Walker, Center for Devices and Radiological Health (HFZ–17), Food and Drug Administration, 2098 Gaither Rd., Rockville, MD 20850, 301–594–1283, ext. 114, e-mail: *KLW@CDRH.FDA.GOV*.

SUPPLEMENTARY INFORMATION: Section 520(f)(3) of the Federal Food, Drug and Cosmetic Act (21 U.S.C. 360j(f)(3)), as amended by the Medical Device Amendments of 1976, provides that each medical device panel include one nonvoting member to represent the interests of the medical device manufacturing industry.

FDA is requesting nominations for nonvoting members representing industry interests for the vacancies listed below:

Medical Device Panels of the Medical Device Advisory Committee	Approximate Date Representative is Needed
Circulatory System Devices Panel	July 1, 2005
Ear, Nose, and Throat Devices Panel	Nov. 1, 2004
Immunology Devices Panel	Mar. 1, 2005
Medical Devices Dispute Resolution Panel	Oct. 1, 2004
Neurological Devices Panel	Dec. 1, 2004
Obstetrics and Gynecology Devices Panel	Feb 1, 2005
Orthopaedic and Rehabilitation Devices Panel	Sept. 1, 2004

I. Functions

The functions of the medical device panels are listed as follows: (1) Review and evaluate data on the safety and effectiveness of marketed and investigational devices and make recommendations for their regulation; (2) advise the Commissioner of Food