(8) The Program Review Panel's report that all written materials have been reviewed as required.

With the prior approval of CDC, copies of proceedings or publications resulting from the conference may be substituted for the performance report, provided they contain the information requested in items one through eight above.

b. Final financial and performance reports, no more than 90 days after the end of the project period.

The reports must be sent to the Grants Management Specialist listed in ''Agency Contacts'' section of this announcement.

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: Victoria E. Saho, Project Officer, Technical Information and Communications Branch, Division of HIV/AIDS Prevention—Intervention Research and Support, National Center for HIV, STD and TB Prevention, 1600 Clifton Road, NE, M/S E49, Atlanta, GA 30333, Telephone: (404) 639–5211, E-mail: *vsaho@cdc.gov*.

For business management and budget assistance, contact: Carlos Smiley, Grants Management Specialist, CDC Procurement and Grants Office, 2920 Brandywine Road, Room 3000, Atlanta, GA 30341–4146, Telephone: (770) 488–2722, E-mail: *csmiley1@cdc.gov*.

For business management and budget assistance in the territories contact: Cynthia Montgomery, Contract Specialist, CDC Procurement and Grants Office, 2920 Brandywine Rd., Atlanta, GA 30341–4146, Telephone: (770) 488–2632, E-mail: *caf5@cdc.gov*.

Dated: December 18, 2003.

Edward Schultz,

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

[FR Doc. 03–31830 Filed 12–22–03; 10:26 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[Program Announcement 04080]

Health Resources and Services Administration Rapid Expansion of Antiretroviral Therapy Programs for HIV-Infected Persons in Selected Countries in Africa and the Caribbean Under the President's Emergency Plan for AIDS Relief; Notice of Availability of Funds-Amendment

A notice announcing the availability of fiscal year (FY) 2004 funds for a cooperative agreement to rapidly expand ART for low-income HIV-infected persons in selected countries in Africa and the Caribbean under the President's Emergency Plan for AIDS Relief was published in the **Federal Register** on December 1, 2003, Volume 68, Number 230, pages 67186–67192. The notice is amended as follows: On page 67188, Column 1, Section ''III.1. Eligible Applicants,'' please insert the following between the first and second paragraphs:

The intent of this solicitation to support organizations that can rapidly implement ARV programs in three or more countries in which each applicant already has an operational presence. Although applications that consist of partnerships or consortia (of organizations that individually do not meet the eligibility criteria) that were formed specifically for the purpose of responding to this RFA would technically meet the eligibility requirements, the duration of the experience of partnerships or consortia (of organizations that individually do not meet the eligibility criteria) in working together will be considered in evaluating the strength of the applicants' proposal.

In addition, on page 67188, Column 2, Section ''IV.1. Address to Request Application Package,'' please disregard the first sentence and replace it with the following:

To apply for this funding opportunity use either application form CDC 5161–1 or CDC 0.1246(E), but we would prefer form CDC 5161–1.

Dated: December 17, 2003.

Sandra R. Manning,

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

[FR Doc. 03–31831 Filed 12–22–03; 10:26 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Citizens Advisory Committee on Public Health Service Activities and Research at Department of Energy (DOE) Sites: Idaho National Engineering and Environmental Laboratory Health Effects Subcommittee (INEELHES)

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92–463), the Agency for Toxic Substances and Disease Registry (ATSDR) and the Centers for Disease Control and Prevention (CDC) announce the following meeting.

Name: Citizens Advisory Committee on Public Health Service Activities and Research at Department of Energy (DOE) Sites: Idaho National Engineering and Environmental Laboratory Health Effects Subcommittee (INEELHES).

Times and Dates: 1 p.m.–4:45 p.m., January 21, 2004. 8 a.m.–4:30 p.m., January 22, 2004.

Place: The Hilton Garden Inn, 145 East Riverside Drive, Eagle, Idaho 83616, telephone 208–938–9600, fax 208–938–5200.

Status: Open to the public, limited only by the space available. The meeting room accommodates approximately 50 people.

Background: Under a Memorandum of Understanding (MOU) signed in December 1990 with DOE, and replaced by MOUs signed in 1996 and 2000, the Department of Health and Human Services (HHS) was given the responsibility and resources for conducting analytic epidemiologic investigations of residents of communities in the vicinity of DOE facilities, workers at DOE facilities, and other persons potentially exposed to radiation or to potential hazards from non-nuclear energy production use. HHS delegated program responsibility to CDC.

In addition, a memo was signed in October 1990 and renewed in November 1992, 1996, and in 2000, between ATSDR and DOE. The MOU delineates the responsibilities and procedures for ATSDR's public health activities at DOE sites required under sections 104, 105, 107, and 120 of the Comprehensive Environmental Response, Compensation, and Liability Act (CERCLA or ''Superfund''). These activities include health consultations and public health assessments at DOE sites listed on, or proposed for, the Superfund National Priorities List and at sites that are the subject of petitions from the public; and other health-related activities such as epidemiologic studies, health surveillance, exposure and disease registries, health education, substance-specific applied research, emergency response, and preparation of toxicological profiles.

Purpose: This subcommittee is charged with providing advice and recommendations to the Director, CDC, and the Administrator, ATSDR, regarding community concerns pertaining to CDC's and ATSDR's public health activities and research at this DOE

site. The purpose of this meeting is to provide a forum for community interaction and to serve as a vehicle for community concerns to be expressed as advice and recommendations to CDC and ATSDR.

Matters to be Discussed: Agenda items include Status of Stanford Cohen & Associates' Draft Report; INEEL Oversight Program of Food Products Grown Near the Aquifer; Presentation on Additional Food Products Grown Near the Aquifer; Overview of the Cancer Data Registry of Idaho and Minority Data; Progress Report on the National Institute for Occupational Safety and Health INEEL Cohort Data; Presentation of the Comprehensive Environmental Response, Compensation and Liability Act and the Relationship Between Maximum Contaminant Levels and Risk; Presentation on Fish as Bioconcentrators; and a Report on Other Activities at the Radiation Studies Branch. Agenda items are subject to change as priorities dictate.

For Further Information Contact: Ms. Natasha Friday, Executive Secretary, INEELHES, Radiation Studies Branch, Division of Environmental Hazards and Health Effects, National Center for Environmental Health, CDC, 1600 Clifton Road, NE (E–39), Atlanta, Georgia 30333, telephone (404) 498–1800, fax (404) 498–1811.

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 CDC and ATSDR.

Dated: December 18, 2003.

Alvin Hall,

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

[FR Doc. 03–31663 Filed 12–23–03; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[CMS–9019–N]

Medicare and Medicaid Programs; Quarterly Listing of Program Issuances—July 2003 Through September 2003

AGENCY: Centers for Medicare & Medicaid Services (CMS), HHS.

ACTION: Notice.

SUMMARY: This notice lists CMS manual instructions, substantive and interpretive regulations, and other **Federal Register** notices that were published from July 2003 through September 2003, relating to the Medicare and Medicaid programs. This notice provides information on national coverage determinations affecting specific medical and health care services under Medicare. Additionally, this notice identifies certain devices with investigational device exemption numbers approved by the Food and Drug Administration that potentially may be covered under Medicare. Finally, this notice also includes listings of all approval numbers from the Office of Management and Budget for collections of information in CMS regulations.

Section 1871(c) of the Social Security Act requires that we publish a list of Medicare issuances in the **Federal Register** at least every 3 months. Although we are not mandated to do so by statute, for the sake of completeness of the listing, and to foster more open and transparent collaboration efforts, we are also including all Medicaid issuances and Medicare and Medicaid substantive and interpretive regulations (proposed and final) published during this 3-month time frame.

FOR FURTHER INFORMATION CONTACT: It is possible that an interested party may have a specific information need and not be able to determine from the listed information whether the issuance or regulation would fulfill that need. Consequently, we are providing information contact persons to answer general questions concerning these items. Copies are not available through the contact persons. (*See* Section III of this notice for how to obtain listed material.)

Questions concerning items in Addendum III may be addressed to Karen Bowman, Office of Strategic Operations and Regulatory Affairs, Centers for Medicare & Medicaid Services, C5–16–03, 7500 Security Boulevard, Baltimore, MD 21244–1850, or you can call (410) 786–5252.

Questions concerning national coverage determinations in Addendum V may be addressed to Patricia Brocato-Simons, Office of Clinical Standards and Quality, Centers for Medicare & Medicaid Services, C1–09–06, 7500 Security Boulevard, Baltimore, MD 21244–1850, or you can call (410) 786–0261.

Questions concerning Investigational Device Exemptions items in Addendum VI may be addressed to Sharon Hippler, Office of Clinical Standards and Quality, Centers for Medicare & Medicaid Services, C5–13–27, 7500 Security Boulevard, Baltimore, MD 21244–1850, or you can call (410) 786–4633.

Questions concerning approval numbers for collections of information in Addendum VII may be addressed to Dawn Willinghan, Office of Strategic Operations and Regulatory Affairs, Regulations Development and Issuances Group, Centers for Medicare & Medicaid Services, C5–09–26, 7500 Security Boulevard, Baltimore, MD 21244–1850, or you can call (410) 786–6141.

Questions concerning all other information may be addressed to Gwendolyn Johnson, Office of Strategic Operations and Regulatory Affairs, Regulations Development and Issuances Group, Centers for Medicare & Medicaid Services, C5–12–26, 7500 Security Boulevard, Baltimore, MD 21244–1850, or you can call (410) 786–6954.

SUPPLEMENTARY INFORMATION:

I. Program Issuances

The Centers for Medicare & Medicaid Services (CMS) is responsible for administering the Medicare and Medicaid programs. These programs pay for health care and related services for 39 million Medicare beneficiaries and 35 million Medicaid recipients. Administration of the two programs involves (1) Furnishing information to Medicare beneficiaries and Medicaid recipients, health care providers, and the public and (2) maintaining effective communications with regional offices, State governments, State Medicaid agencies, State survey agencies, various providers of health care, all Medicare contractors that process claims and pay bills, and others. To implement the various statutes on which the programs are based, we issue regulations under the authority granted to the Secretary of the Department of Health and Human Services under sections 1102, 1871, 1902, and related provisions of the Social Security Act (the Act). We also issue various manuals, memoranda, and statements necessary to administer the programs efficiently.

Section 1871(c)(1) of the Act requires that we publish a list of all Medicare manual instructions, interpretive rules, statements of policy, and guidelines of general applicability not issued as regulations at least every 3 months in the **Federal Register**. We published our first notice June 9, 1988 (53 FR 21730). Although we are not mandated to do so by statute, for the sake of completeness of the listing of operational and policy statements, and to foster more open and transparent collaboration, we are continuing our practice of including Medicare substantive and interpretive regulations (proposed and final) published during the respective 3-month time frame.

II. How To Use the Addenda

This notice is organized so that a reader may review the subjects of manual issuances, memoranda,