b. The extent to which the applicant has demonstrated a community action component to improve and expand domestic violence intervention and prevention services throughout the state.

c. The extent to which the applicant has demonstrated an ability to plan and implement outreach and public education campaigns regarding domestic violence.

d. The extent to which the applicant has cooperated in or spearheaded the development of state protocols regarding domestic violence. Applicants should include examples of such protocols as Attachment A.

e. The extent to which the applicant has experience in funding and monitoring sub-awards.

f. The extent to which the applicant has demonstrated experience in providing training and technical assistance to local domestic violence programs whether through conferences or other training/technical assistance mechanisms.

g. The extent to which the applicant participated in the development of and/or referenced any state-level violence against women prevention plan.

6. Measures of Effectiveness (Not Scored)

The extent to which the applicant provided objective/quantifiable measures regarding the DELTA program's intended outcomes that will demonstrate the accomplishment of the various identified objectives of the cooperative agreement. Applicants are encouraged to consult the Program Guidance (See Attachment 3 in the application kit) for further clarification.

7. Budget (Not Scored)

The applicant should provide a detailed budget with complete line-item justification of all proposed costs consistent with the stated activities in this program announcement. Applicants should be precise about the purpose of each budget item and must provide itemized calculations of proposed costs. These funds should not be used to supplant existing efforts.

I. Other Requirements

Technical Reporting Requirements

Provide CDC with original plus two copies of:

1. Semiannual progress reports will be submitted as part of the grantee's continuation application. The progress report will include a data requirement that demonstrates measures of effectiveness. Specific guidance will be provided by NCIPC for the content of progress reports.

2. Financial status report, no more than 90 days after the end of the budget period.

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

Send all reports to the Grants Management Specialist identified in the ''Where to Obtain Additional Information'' section of this announcement.

The following additional requirements are applicable to this program. For a complete description of each, see Attachment I of the application kit.

AR–10 Smoke-Free Workplace Requirements

AR–11 Healthy People 2010

AR–12 Lobbying Restrictions

AR–13 Prohibition on Use of CDC Funds for Certain Gun Control Activities

AR–15 Proof of Non-Profit Status

J. Where To Obtain Additional Information

This and other CDC announcements, the necessary applications, and associated forms can be found on the CDC home page Internet address—*http://www.cdc.gov.* Click on ''Funding'' then ''Grants and Cooperative Agreements.''

For business management technical assistance, contact: Van A. King, Grants Management Specialist, Procurement and Grants Office, Centers for Disease Control and Prevention, 2920 Brandywine Road, Room 3000, Atlanta, GA 30341–4146, Telephone number (770) 488–2751, e-mail address: *VKing@cdc.gov.*

For program technical assistance, contact: Janet Saul, PhD, National Center for Injury Prevention and Control, Centers for Disease Control and Prevention, 4770 Buford Hwy, NW, Mailstop K–60, Atlanta, GA 30341–1125, Telephone number (770) 488–4733, e-mail address: *JSaul@cdc.gov.*

Dated: July 25, 2002.

Edward Schultz,

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

[FR Doc. 02–19284 Filed 7–30–02; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Disease, Disability, and Injury Prevention and Control Special Emphasis Panel: A Community-Based Intervention With Opinion Leaders to Achieve Syphilis Elimination, Program Announcement 02044

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): A Community-Based Intervention with Opinion Leaders to Achieve Syphilis Elimination, PA# 02044.

Times and Dates: 9 a.m.–9:30 a.m., August, 15, 2002 (Open), 9:30 a.m.–4:30 p.m., August 15, 2002 (Closed).

Place: Centers for Disease Control and Prevention 12 Corporate Square Boulevard—Room 1307 Atlanta, GA 30329

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# 02044.

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, Georgia 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.

Joe E. Salter,

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

[FR Doc. 02–19402 Filed 7–29–02; 12:22 pm]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Advisory Board on Radiation and Worker Health: Meeting

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 committee meeting.

Name: Advisory Board on Radiation and Worker Health (ABRWH).

Times and Dates: 12:30 p.m.–5:30 p.m., August 14, 2002. 8 a.m.–5 p.m., August 15, 2002.

Place: Hyatt Regency Cincinnati, 151 West Fifth Street, Cincinnati, Ohio 45202, telephone 513/579–1234, fax 513/352–0245.

Status: Open to the public, limited only by the space available. The meeting room accommodates approximately 65 people.

Background: The Advisory Board on Radiation and Worker Health (''the Board'') was established under the Energy Employees Occupational Illness Compensation Program Act of 2000 to advise the President, through the Secretary of Health and Human Services (HHS), on a variety of policy and technical functions required to implement and effectively manage the new compensation program. Key functions of the Board include providing advice on the development of probability of causation guidelines which have been promulgated by HHS, advice on methods of dose reconstruction which have been promulgated as an interim final rule, evaluation of the validity and quality of dose reconstructions conducted by the National Institute for Occupational Safety and Health (NIOSH) for qualified cancer claimants, and advice on the addition of classes of workers to the Special Exposure Cohort.

In December 2000 the President delegated responsibility for funding, staffing, and operating the Board to HHS, which subsequently delegated this authority to the CDC. NIOSH implements this responsibility for CDC. The charter was signed on August 3, 2001, and in November 2001 the President completed the appointment of an initial roster of 10 Board members. In April 2002 the President appointed an additional member to ensure more balanced representation on the Board. The initial tasks of the Board will be to review and provide advice on the proposed and interim rules of HHS.

Purpose: This board is charged with (a) providing advice to the Secretary, HHS, on the development of guidelines under Executive Order 13179; (b) providing advice to the Secretary, HHS, on the scientific validity and quality of dose reconstruction efforts performed for this Program; and (c) upon request by the Secretary, HHS, advise the Secretary on whether there is a class of employees at any Department of Energy facility who were exposed to radiation but for whom it is not feasible to estimate their radiation dose, and on whether there is reasonable likelihood that such radiation doses may have endangered the health of members of this class.

Matters To Be Discussed: Agenda for this meeting will include presentations for the Board's information on the adjudication of claims for atomic veterans, dose reconstruction for atomic veterans, and probability of causation determination for atomic veterans. The Board's agenda also includes development of comments on the Special Exposure Cohort Petitioning Process Guidelines (NPRM), dose reconstruction workgroup discussion and issues, and Board discussion of Board responsibilities.

Agenda items are subject to change as priorities dictate.

Contact Person for More Information: Larry Elliott, Executive Secretary, ABRWH, NIOSH, CDC, 4676 Columbia Parkway, Cincinnati, Ohio 45226, telephone 513/841–4498, fax 513/458–7125.

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.

Dated: July 25, 2002.

John C. Burckhardt,

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

[FR Doc. 02–19283 Filed 7–30–02; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 02N–0315]

Agency Information Collection Activities; Proposed Collection; Comment Request; Medical Devices: Humanitarian Use Devices

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing an opportunity for public comment on the proposed collection of certain information by the agency. Under the Paperwork Reduction Act of 1995 (the PRA), Federal agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing information collection, and to allow 60 days for public comment in response to the notice. This notice solicits comments on information collection requirements for humanitarian use devices.

DATES: Submit written and electronic comments on the collection of information by September 30, 2002.

ADDRESSES: Submit electronic comments on the collection of information to http://www.accessdata.fda.gov/scripts/oc/dockets/edockethome.cfm. Submit written comments on the collection of information to the Dockets Management Branch (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. All comments should be identified with the docket number found in brackets in the heading of this document.

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: Under the PRA (44 U.S.C. 3501–3520), Federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. ''Collection of information'' is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA (44 U.S.C. 3506(c)(2)(A)) requires Federal agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, FDA is publishing notice of the proposed collection of information set forth in this document.

With respect to the following collection of information, FDA invites comments on: (1) Whether the proposed collection of information is necessary for the proper performance of FDA's functions, including whether the information will have practical utility; (2) the accuracy of FDA's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) ways to minimize the