hours estimated for this ICR are summarized in the table below.

TOTAL ESTIMATED ANNUALIZED BURDEN—HOURS

Form name	Number of re- spondents	Number of re- sponses per respondent	Average burden per response (in hours)	Total burden hours
Adult Viral Hepatitis Prevention Coordinators	4 16 12 12 12 12	1 1 1 1 1	1.5 45/60 30/60 30/60 30/60 30/60	6 12 6 6 6 6
Total				42

Keith A. Tucker,

Information Collection Clearance Officer. [FR Doc. 2013–07541 Filed 4–1–13; 8:45 am] BILLING CODE 4150–47–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Meeting of the Presidential Advisory Council on HIV/AIDS

AGENCY: Department of Health and Human Services, Office of the Secretary, Office of the Assistant Secretary for Health.

ACTION: Notice.

SUMMARY: As stipulated by the Federal Advisory Committee Act, the U.S. Department of Health and Human Service (DHHS) is hereby giving notice that the Presidential Advisory Council on HIV/AIDS (PACHA) will hold a meeting to discuss implementation of the Patient Protection and Affordable Care Act. The meeting will be open to the public.

DATES: The meeting will be held April 22, 2013, from 9:00 a.m. to approximately 5:30 p.m. (EDT).

ADDRESSES: U.S. Department of Health and Human Services, 200 Independence Avenue SW., Washington, DC 20201 in the Auditorium.

FOR FURTHER INFORMATION CONTACT: Ms. Caroline Talev, Public Health Assistant, Presidential Advisory Council on HIV/ AIDS, Department of Health and Human Services, 200 Independence Avenue SW., Room 443H, Washington, DC 20201; (202) 205–1178. More detailed information about PACHA can be obtained by accessing the Council's Web site www.aids.gov/pacha.

SUPPLEMENTARY INFORMATION: PACHA was established by Executive Order 12963, dated June 14, 1995 as amended by Executive Order 13009, dated June 14, 1996. The Council was established to provide advice, information, and recommendations to the Secretary regarding programs and policies intended to promote effective prevention of HIV disease and AIDS. The functions of the Council are solely advisory in nature.

The Council consists of not more than 25 members. Council members are selected from prominent community leaders with particular expertise in, or knowledge of, matters concerning HIV and AIDS, public health, global health, philanthropy, marketing or business, as well as other national leaders held in high esteem from other sectors of society. Council members are appointed by the Secretary or designee, in consultation with the White House Office on National AIDS Policy. The agenda for the upcoming meeting will be posted on the Council's Web site at www.aids.gov/pacha.

Public attendance at the meeting is limited to space available. Individuals who plan to attend and need special assistance, such as sign language interpretation or other reasonable accommodations, should notify the designated contact person. Preregistration for public attendance is advisable and can be accomplished by contacting Caroline Talev at caroline.talev@hhs.gov. Members of the public will have the opportunity to provide comments at the meeting. Any individual who wishes to participate in the public comment session must register with Caroline Talev at *caroline.talev@hhs.gov;* registration for public comment will not be accepted by telephone. Public comment will be limited to two minutes per speaker. Any members of the public who wish to have printed material distributed to PACHA members at the meeting should submit, at a minimum, 1 copy of the materials to Caroline Taley, no later than close of business Monday, April 15, 2013.

Contact information for the PACHA contact person is listed above.

Dated: March 14, 2013.

B. Kaye Hayes,

Executive Director, Presidential Advisory Council on HIV/AIDS. [FR Doc. 2013–07614 Filed 4–1–13; 8:45 am] BILLING CODE 4150–43–P

BILLING CODE 4150-43-

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Call for Collaborating Partners for National Women's Health Week

AGENCY: Office on Women's Health, Office of the Assistant Secretary for Health, Office of the Secretary, Department of Health and Human Services.

ACTION: Notice.

SUMMARY: The Department of Health and Human Services (HHS), Office on Women's Health (OWH) invites public and private-sector health-related organizations to participate in National Women's Health Week (NWHW) as partners to help create awareness of women's health issues and educate women about improving their health and preventing disease.

DATES: Representatives of women's health organizations should submit expressions of interest no later than April 18, 2013.

ADDRESSES: Expressions of interest, comments, and questions may be submitted by electronic mail to *Henrietta.terry@hhs.gov* or by regular mail to Jill Wasserman, Office on Women's Health, Department of Health and Human Services, 200 Independence Avenue SW., Room 733E, Washington, DC 20201; or via fax to (202) 690–7172.

FOR FURTHER INFORMATION CONTACT: Henrietta Terry on (202) 205–1952. SUPPLEMENTARY INFORMATION: The OWH was established in 1991 to improve the health of American women by advancing and coordinating a comprehensive women's health agenda throughout HHS. The office fulfills its mission by advancing policy and issuing competitive contracts to an array of community, academic, and other organizations at the national and community levels. In addition, OWH's national educational campaigns provide information about the important steps women can take to improve and maintain their health, such as NWHW.

NWHW is a week-long health observance that kicks off on Mother's Day, Sunday, May 12 and ends Saturday, May 18, 2013. NWHW seeks to educate women about improving their physical and mental health and preventing disease. More than 2,200 events were held nationwide in 2012. Week-long, daily messages encourage women to make their health a top priority and take simple steps for a longer, healthier, and happier life. For more information about NWHW, please visit http://womenshealth.gov/nwhw/.

Dated: March 27, 2013.

Nancy C. Lee,

Deputy Assistant Secretary for Health— Women's Health.

[FR Doc. 2013–07617 Filed 4–1–13; 8:45 am] BILLING CODE 4150–33–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Statement of Organization, Functions, and Delegations of Authority

Part C (Centers for Disease Control and Prevention) of the Statement of Organization, Functions, and Delegations of Authority of the Department of Health and Human Services (45 FR 67772–76, dated October 14, 1980, and corrected at 45 FR 69296, October 20, 1980, as amended most recently at 78 FR 5812, dated January 28, 2013) is amended to reflect the reorganization of the Office for State, Tribal, Local, and Territorial Support.

Section C–B, Organization and Functions, is hereby amended as follows:

Delete in its entirety the title and function statements for the Knowledge Management Office (CQA5), Office of the Director (CQA).

Revise the functional statement for the Public Health Law Office (CQA2), Office of the Director (CQA) as follows:

After item (8), insert the following: (9) establish collaboration and coordination between clinical medicine and public health to better coordinate and partner for healthier communities.

Dated: March 22, 2013.

Sherri A. Berger,

Chief Operating Officer, Centers for Disease Control and Prevention. [FR Doc. 2013–07582 Filed 4–1–13; 8:45 am] BILLING CODE 4160–18–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Statement of Organization, Functions, and Delegations of Authority

Part C (Centers for Disease Control and Prevention) of the Statement of Organization, Functions, and Delegations of Authority of the Department of Health and Human Services (45 FR 67772–76, dated October 14, 1980, and corrected at 45 FR 69296, October 20, 1980, as amended most recently at 78 FR 5812, dated January 28, 2013) is amended to reflect the reorganization of the Office of the Associate Director for Science, Office of the Director, Centers for Disease Control and Prevention.

Section C–B, Organization and Functions, is hereby amended as follows:

Delete in its entirety the title and function statements for the Public Health Prevention Service Branch (CPLCC), Division of Leadership and Practice (CPLP).

Dated: March 22, 2013.

Sherri A. Berger,

Chief Operating Officer, Centers for Disease Control and Prevention.

[FR Doc. 2013–07545 Filed 4–1–13; 8:45 am] BILLING CODE 4160–18–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA-2013-N-0338]

Center for Devices and Radiological Health: Experiential Learning Program

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration's (FDA) Center for Devices and Radiological Health (CDRH or Center) is announcing an invitation for participation in its Experiential Learning Program (ELP). The ELP provides a formal training mechanism

for regulatory review staff to visit research, clinical, manufacturing, and health care facilities to observe firsthand how medical devices are designed, developed, and utilized. This training is intended to provide CDRH staff with an opportunity to observe the device development life cycle and provide a better understanding of the medical devices they review, and the challenges faced throughout development, testing, manufacturing, and clinical use. The purpose of this document is to invite medical device and health care facilities to participate in this formal training program for FDA's medical device review staff, or to contact CDRH for more information regarding the program.

DATES: Submit either an electronic or written request for participation in this program by May 2, 2013. The request should include a description of your facility relative to product areas CDRH regulates. Please include the Area of Interest/Medical Device or Technology (identified in table 10r 2) that the visit will demonstrate to CDRH staff.

ADDRESSES: Submit either electronic requests to *http://www.regulations.gov* or written requests to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville MD 20852. Identify comments with the docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT:

Latonya Powell, Center for Devices and Radiological Health, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 66, Rm. 4448, Silver Spring, MD 20993–0002, 301–796–6965, FAX: 301–827–3079,

Latonya.powell@fda.hhs.gov.

SUPPLEMENTARY INFORMATION:

I. Background

CDRH launched the ELP Pilot in 2012 and will fully implement the program in 2013. The Center is responsible for ensuring the safety and effectiveness of medical devices marketed in the United States. Furthermore, CDRH assures that patients and providers have timely and continued access to safe, effective, highquality medical devices and safe radiation-emitting products. In support of this mission, the Center launched various training and development initiatives to enhance performance of its regulatory review staff and other staff involved in the premarket review process. CDRH is driven to advance regulatory science; provide industry with predictable, consistent, transparent, and efficient regulatory pathways; and assure consumer