Darius Taylor,

Deputy, Information Collection Clearance Officer.

[FR Doc. 2013–19137 Filed 8–7–13; 8:45 am]

BILLING CODE 4150–38–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Board of Scientific Counselors, National Center for Health Statistics

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), National Center for Health Statistics (NCHS) announces the following meeting of the aforementioned committee:

Times and Dates:

11:00 a.m.–5:30 p.m., September 19, 2013

8:30 a.m.–1:00 p.m., September 20, 2013

Place: NCHS Headquarters, 3311 Toledo Road, Hyattsville, Maryland 20782.

Status: This meeting is open to the public; however, visitors must be processed in accordance with established federal policies and procedures. For foreign nationals or non-US citizens, pre-approval is required (please contact Gwen Mustaf, 301–458–4500, *glm4@cdc.gov* or Virginia Cain, *vcain@cdc.gov* at least 10 days in advance for requirements). All visitors are required to present a valid form of picture identification issued by a state, federal or international government. As required by the Federal Property Management Regulations, Title 41, Code of Federal Regulation, Subpart 101–20.301, all persons entering in or on Federal controlled property and their packages, briefcases, and other containers in their immediate possession are subject to being x-rayed and inspected. Federal law prohibits the knowing possession or the causing to be present of firearms, explosives and other dangerous weapons and illegal substances. The meeting room accommodates approximately 100 people.

Purpose: This committee is charged with providing advice and making recommendations to the Secretary, Department of Health and Human Services; the Director, CDC; and the Director, NCHS, regarding the scientific and technical program goals and objectives, strategies, and priorities of NCHS.

Matters to be Discussed: The agenda will include welcome remarks by the Acting Director, NCHS; Demo of the NHIS Online Analytic Real-time System (OARS); initiation of Office of Analysis and Epidemiology review.

Requests to make oral presentations should be submitted in writing to the contact person listed below. All requests must contain the name, address, telephone number, and organizational affiliation of the presenter.

Written comments should not exceed five single-spaced typed pages in length and must be received by September 4, 2013.

The agenda items are subject to change as priorities dictate.

Contact Person for more Information: Virginia S. Cain, Ph.D., Director of Extramural Research, NCHS/CDC, 3311 Toledo Road, Room 7208, Hyattsville, Maryland 20782, telephone (301) 458–4500, fax (301) 458–4020.

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 Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

Elaine L. Baker,

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

[FR Doc. 2013–19156 Filed 8–7–13; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2013–N–0001]

Pediatric Ethics Subcommittee of the Pediatric Advisory Committee; Notice of Meeting

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

This notice announces a forthcoming meeting of a subcommittee of a public advisory committee of the Food and Drug Administration (FDA). The meeting will be open to the public.

Name of Subcommittee: Pediatric Ethics Subcommittee of the Pediatric Advisory Committee.

General Function of the Subcommittee: To advise and make recommendations to the Pediatric Advisory Committee on pediatric ethical issues.

Date and Time: The meeting will be held on September 9, 2013, from 8 a.m. to 5:30 p.m. and September 10, 2013, from 8 a.m. to 3 p.m.

Location: Doubletree Hilton Hotel, 8727 Colesville Rd., Silver Spring, MD 20910, 301–589–5200 or visit the hotel's Web site at *http://doubletree3.hilton.com/en/hotels/maryland/doubletree-by-hilton-hotel-washington-dc-silver-spring-DCASSDT/index.html.*

Contact Person: Walter Ellenberg, Office of the Commissioner, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 32, rm. 5154, Silver Spring, MD 20993, 301–796–0885, email *walter.ellenberg@fda.hhs.gov* or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area). A notice in the **Federal Register** about last minute modifications that impact a previously announced subcommittee meeting cannot always be published quickly enough to provide timely notice. Therefore, you should always check the Agency's Web site at *http://www.fda.gov/AdvisoryCommittees/default.htm* and scroll down to the appropriate advisory committee meeting link, or call the advisory committee information line to learn about possible modifications before coming to the meeting.

Agenda: On September 9 and 10, 2013, the Pediatric Ethics Subcommittee of the Pediatric Advisory Committee will meet to discuss ethical issues in pediatric product development, including medical counter measures, focusing on the concepts of minimal risk, disorder or condition, and exposure of pediatric subjects to risks under 21 CFR 50.54.

FDA intends to make background material available to the public no later than 2 business days before the meeting. If FDA is unable to post the background material on its Web site prior to the meeting, the background material will be made publicly available at the location of the subcommittee meeting, and the background material will be posted on FDA's Web site after the meeting. Background material is available at *http://www.fda.gov/AdvisoryCommittees/Calendar/default.htm.* Scroll down to the appropriate advisory committee meeting link.

Procedure: Interested persons may present data, information, or views, orally or in writing, on issues pending before the committee. Written submissions may be made to the contact person on or before September 9, 2013. Oral presentations from the public will be scheduled between approximately 2 p.m. and 3 p.m. Those individuals interested in making formal oral presentations should notify the contact

person and submit a brief statement of the general nature of the evidence or arguments they wish to present, the names and addresses of proposed participants, and an indication of the approximate time requested to make their presentation on or before August 30, 2013. Time allotted for each presentation may be limited. If the number of registrants requesting to speak is greater than can be reasonably accommodated during the scheduled open public hearing session, FDA may conduct a lottery to determine the speakers for the scheduled open public hearing session. The contact person will notify interested persons regarding their request to speak by September 3, 2013.

Persons attending FDA's subcommittee meetings are advised that the Agency is not responsible for providing access to electrical outlets.

FDA welcomes the attendance of the public at this meeting and will make every effort to accommodate persons with physical disabilities or special needs. If you require special accommodations due to a disability, please contact Walter Ellenberg at 301–796–0885, email *walter.ellenberg@fda.hhs.gov,* at least 7 days in advance of the meeting.

FDA is committed to the orderly conduct of its advisory committee meetings. Please visit our Web site at *http://www.fda.gov/AdvisoryCommittees/AboutAdvisoryCommittees/ucm111462.htm* for procedures on public conduct during advisory committee meetings.

Notice of this meeting is given under the Federal Advisory Committee Act (5 U.S.C. app. 2).

Dated: August 2, 2013.

Leslie Kux,

Assistant Commissioner for Policy.

[FR Doc. 2013–19138 Filed 8–7–13; 8:45 am]

BILLING CODE 4160–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Agency Information Collection Activities: Submission to OMB for Review and Approval; Public Comment Request

AGENCY: Health Resources and Services Administration, HHS.

ACTION: Notice.

SUMMARY: In compliance with Section 3507(a)(1)(D) of the Paperwork Reduction Act of 1995, the Health Resources and Services Administration (HRSA) has submitted an Information Collection Request (ICR) to the Office of Management and Budget (OMB) for review and approval. Comments submitted during the first public review of this ICR will be provided to OMB. OMB will accept further comments from the public during the review and approval period.

DATES: Comments on this ICR should be received within 30 days of this notice.

ADDRESSES: Submit your comments, including the Information Collection Request Title, to the desk officer for HRSA, either by email to *OIRA_submission@omb.eop.gov* or by fax to 202–395–5806.

FOR FURTHER INFORMATION CONTACT: To request a copy of the clearance requests submitted to OMB for review, email the HRSA Information Collection Clearance Officer at *paperwork@hrsa.gov* or call (301) 443–1984.

SUPPLEMENTARY INFORMATION:

Information Collection Request Title: Analyzing Title V Programs in the Context of the Affordable Care Act

OMB No.: 0915–xxxx—New

Abstract: The Affordable Care Act (ACA) will make affordable health coverage available to all legal U.S. residents, as well as guide transformation in the delivery of medicine and public health services. For children, expanded coverage has come about gradually over the past two decades and implementation of major coverage provisions of the ACA in 2014 will result in some shifts in child health coverage.

The Title V Maternal and Child Health (MCH) Block Grant, administered by the Health Resources and Services Administration's Maternal and Child Health Bureau, provides the foundation for ensuring the health of the nation's mothers, women, children, and youth, including children and youth with special health care needs and their families. Many ACA provisions, like state Medicaid expansions and mandatory health insurance, will change the face of health insurance demand and services provided. In response, State Title V programs will focus on increasing access, equality, and health equity.

A proposed data collection form has been developed to collect health care services budget information from Title V MCH Block Grant recipients to better understand the types of direct services currently provided by Title V MCH programs. This form will request information on expenditures for medical services in addition to data on the individuals served.

Need and Proposed Use of the Information: As children shift between coverage categories as a result of implementation of the ACA, HRSA would like to quantify the impact of these shifts on the federal investment in Title V funding specifically through the federal funds provided via the Title V MCH Block Grant. To do this, HRSA will need to survey states to collect information on whether they use federal Title V MCH Block Grant funds to reimburse health care practitioners who provide services to children and pregnant women.

Likely Respondents: The respondents to the survey will be the Title V Program Directors in the states, the District of Columbia, and Puerto Rico.

Burden Statement: Burden in this context means the time expended by persons to generate, maintain, retain, disclose or provide the information requested. This includes the time needed to review instructions; to develop, acquire, install and utilize technology and systems for the purpose of collecting, validating and verifying information, processing and maintaining information, and disclosing and providing information; to train personnel and to be able to respond to a collection of information; to search data sources; to complete and review the collection of information; and to transmit or otherwise disclose the information. The total annual burden hours estimated for this ICR are summarized in the table below.