clearance under 44 U.S.C. 3507. An Agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. OMB has now approved the information collection and has assigned OMB control number 0910–0625. The approval expires on March 31, 2016. A copy of the supporting statement for this information collection is available on the Internet at *http://www.reginfo.gov/public/do/PRAMain.*

Dated: April 15, 2013.

Leslie Kux,

Assistant Commissioner for Policy.

[FR Doc. 2013–09182 Filed 4–18–13; 8:45 am]

BILLING CODE 4160–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Agency Information Collection Activities; Proposed Collection; Comment Request

ACTION: Notice

SUMMARY: In compliance with the requirement for opportunity for public comment on proposed data collection projects (Section 3506(c)(2)(A) of Title 44, United States Code, as amended by the Paperwork Reduction Act of 1995, Pub. L. 104–13), the Health Resources and Services Administration (HRSA) publishes periodic summaries of proposed projects being developed for submission to the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995. To request more information on the proposed project or to obtain a copy of the data collection plans and draft instruments, email *paperwork@hrsa.gov* or call the HRSA Reports Clearance Officer at (301) 443–1984.

HRSA especially requests comments on: (1) The necessity and utility of the proposed information collection for the proper performance of the agency's functions, (2) the accuracy of the estimated burden, (3) ways to enhance the quality, utility, and clarity of the information to be collected, and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

Information Collection Request Title: Evaluating the Impact of 1115 Medicaid Waivers on Ryan White HIV/AIDS Program and Its Clients and Providers (OMB No. 0915–xxxx)—New

Abstract: Section 1115 of the Social Security Act allows states to develop, test, and implement new approaches to providing Medicaid coverage outside of federal program rules. Leading up to full implementation of the Affordable Care Act, states have begun to use Section 1115 Medicaid demonstration waivers as a ''bridge'' to 2014. This project will examine 1115 Medicaid waivers that have expanded eligibility to specifically include people living with HIV/AIDS (PLWH) who are not otherwise eligible for Medicaid services. Since 1990, the Ryan White HIV/AIDS Program (RWHAP) has provided funding for primary care, medications, and support services for PLWH, helping fill the health care and service gap for those who are uninsured or ineligible for Medicaid. Given the important role of the RWHAP and Medicaid in meeting the health care needs of PLWH, there is a need to better understand how Medicaid expansion and the 1115 Medicaid waivers will affect the RWHAP and how the waivers have prepared states for implementation of the Affordable Care Act.

As part of this project, case studies will be conducted in eight states that have implemented 1115 Medicaid waivers to expand Medicaid eligibility for PLWH. The case studies will include site visits and discussions with the state Medicaid programs and with RWHAP grantees and service providers to examine the waivers and their impact on PLWH. In addition, the studies will explore whether and how the 1115 Medicaid waivers have helped states and RWHAP grantees and providers prepare for implementation of the Affordable Care Act, including providing insights into Medicaid expansion. Data will be collected through qualitative interviews, guided by discussion tools with questions tailored for four specific groups of individuals from: (1) State Medicaid Agencies; (2) Ryan White HIV/AIDS Program Part B grantees and service providers; (3) Ryan White HIV/AIDS Program Part A grantees and service providers; and (4) Ryan White HIV/AIDS Program Part C grantees and clinical providers.

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 Information Collection Request are summarized in the table below.

The annual estimate of burden is as follows:

Form name	Number of respondents	Number of responses per respondent	Total responses	Average burden per response (in hours)	Total burden hours
Qualitative Interview Data Collection Tool—State Medicaid Agencies	40	1	40	2	80
Qualitative Interview Data Collection Tool—RWHAP Part A Administrators/Planning Councils	80	1	80	2	160
Qualitative Interview Data Collection Tool—RWHAP Part B/ADAP Directors and Coordinators	80	1	80	2	160
Qualitative Interview Data Collection Tool—RWHAP Part C Grantees/Clinical Providers	80	1	80	2	160
Total	280	.....	280	.....	560

ADDRESSES: Submit your comments to *paperwork@hrsa.gov* or mail the HRSA Reports Clearance Officer, Room 10–29, Parklawn Building, 5600 Fishers Lane, Rockville, MD 20857.

Deadline: Comments on this Information Collection Request must be received within 60 days of this notice.

Dated: April 12, 2013.

Bahar Niakan,

Director, Division of Policy and Information Coordination.

[FR Doc. 2013–09221 Filed 4–18–13; 8:45 am]

BILLING CODE 4165–15–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Advisory Council on Blood Stem Cell Transplantation; Notice of Meeting

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92–463), notice is hereby given of the following meeting:

Name: Advisory Council on Blood Stem Cell Transplantation.

Date and Time: May 16, 2013, 10:00 a.m. to 4:00 p.m. EST.

Place: The meeting will be via audio conference call and Adobe Connect Pro.

Status: The meeting will be open to the public.

Purpose: Pursuant to Public Law 109–129, 42 U.S.C. 274k (section 379 of the Public Health Service Act, as amended), the Advisory Council on Blood Stem Cell Transplantation (ACBSCT) advises the Secretary of the Department of Health and Human Services and the Administrator, Health Resources and Services Administration (HRSA), on matters related to the activities of the C.W. Bill Young Cell Transplantation Program (Program) and the National Cord Blood Inventory Program.

Agenda: The Council will hear reports from ACBSCT Work Groups including: Cord Blood Thawing and Washing; Access to Transplantation; and Advancing Hematopoietic Stem Cell Transplantation for Hemoglobinopathies. The Council will also hear presentations and discussions on topics including: Accreditation, Adverse Event Reporting, and Unmet Need. Agenda items are subject to change as priorities indicate.

After Council discussions, members of the public will have an opportunity to provide comments. Because of the Council's full agenda and the timeframe in which to cover the agenda topics, public comment may be limited. All public comments will be included in the record of the ACBSCT meeting. Meeting summary notes will be posted on HRSA's Program Web site at *http://bloodcell.transplant.hrsa.gov/ABOUT/Advisory_Council/index.html.*

The draft meeting agenda will be posted on *https://acbsctmeeting.com/public/sitePage.aspx?key=Home.* Those participating in this meeting should register by visiting *https://acbsctmeeting.com/public/sitePage.aspx?key=Home.* The deadline to register for this meeting is Tuesday, May 14, 2013. For all logistical questions and concerns, please contact Deborah Jones, Meeting Planner, by calling (301) 585–1261 or by sending an email to *registration@acbsctmeeting.com.*

The public can join the meeting by:

1. (Audio Portion) Calling the conference Phone Number 800–857–9638 and providing the Participant Code 75841; AND

2. (Visual Portion) Connecting to the ACBSCT Adobe Connect Pro Meeting using the following URL: *https://hrsa.connectsolutions.com/acbsctm/* (copy and paste the link into your browser if it does not work directly, and enter as a guest). Participants should call and connect 15 minutes prior to the meeting in order for logistics to be set up. If you have never attended an Adobe Connect meeting, please test your connection using the following URL: *https://hrsa.connectsolutions.com/common/help/en/support/meeting_test.htm* and get a quick overview by following URL: *http://www.adobe.com/go/connectpro_overview.* Call (301) 443–0437 or send an email to *ptongele@hrsa.gov* if you are having trouble connecting to the meeting site.

Public Comment: It is preferred that persons interested in providing an oral presentation submit a written request, along with a copy of their presentation to: Passy Tongele, MBA, Division of Transplantation, Healthcare Systems Bureau, HRSA, 5600 Fishers Lane, Room 12C–06, Rockville, Maryland 20857; or email at *ptongele@hrsa.gov.* Requests should contain the name, address, telephone number, email address, and any business or professional affiliation of the person desiring to make an oral presentation. Groups having similar interests are requested to combine their comments and present them through a single representative.

The allocation of time may be adjusted to accommodate the level of expressed interest. Persons who do not file an advance request for a presentation, but desire to make an oral statement, may request it at the time of the public comment period. Public participation and ability to comment will be limited to space and time as it permits.

For Further Information Contact: Patricia Stroup, MBA, MPA, Executive Secretary, HSB, Healthcare Systems Bureau, 5600 Fishers Lane, Room 12C–06, Rockville, Maryland 20857; telephone (301) 443–1127.

Dated: April 15, 2013.

Bahar Niakan,

Director, Division of Policy and Information Coordination.

[FR Doc. 2013–09222 Filed 4–18–13; 8:45 am]

BILLING CODE 4165–15–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute of Diabetes and Digestive and Kidney Diseases; Notice of Closed Meetings

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. App.), notice is hereby given of the following meetings.

The meetings will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C., as amended. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: National Institute of Diabetes and Digestive and Kidney Diseases Special Emphasis Panel; NIDDK Central Repositories Non-renewable Sample Access (X01): Hepatitis C and Type 1 Diabetes Biomarkers.

Date: May 9, 2013.

Time: 2:00 p.m. to 4:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Two Democracy Plaza, 6707 Democracy Boulevard, Bethesda, MD 20892 (Telephone Conference Call).

Contact Person: Najma Begum, Ph.D., Scientific Review Officer, Review Branch, DEA, NIDDK, National Institutes of Health, Room 749, 6707 Democracy Boulevard, Bethesda, MD 20892–5452, (301) 594–8894, *begumn@niddk.nih.gov.*

Name of Committee: National Institute of Diabetes and Digestive and Kidney Diseases Special Emphasis Panel; NIDDK–KUH Fellowship Review Committee.

Date: June 3, 2013.

Time: 8:00 a.m. to 5:00 p.m.

Agenda: To review and evaluate grant applications.