(2) Criteria for pilot studies related to newborn screening;

(3) Center for Disease Control and Prevention's Enhancing Data Driven Disease Detection in Newborns (ED3N) Project; and

(4) ACHDNC Decision Matrix.

Agenda items are subject to change as priorities dictate. However, no votes will be held at this meeting to recommend including additional conditions for screening to the Recommended Uniform Screening Panel. Information about ACHDNC, including a roster of members and past meeting summaries, is also available on ACHDNC's website.

Members of the public also will have the opportunity to provide comments on any or all of the above agenda items. Public participants may request to provide general oral comments and may submit written statements in advance of the scheduled meeting. Oral comments will be honored in the order they are requested and may be limited as time allows. Members of the public registered to provide oral public comments on all other newborn screening related topics are tentatively scheduled to provide their statements on Friday, May 5, 2023. Requests to provide a written statement or make oral comments to the ACHDNC must be submitted via the registration website by 12:00 p.m. ET on Wednesday, April 19, 2023. Written comments will be shared with the Committee, so that they have an opportunity to consider them prior to the meeting.

Individuals who need special assistance or another reasonable accommodation should notify Alaina Harris at the address and phone number listed above at least 10 business days prior to the meeting.

Since this meeting occurs in a federal government building, attendees must go through a security check to enter the building. Non-U.S. citizen attendees must notify HRSA of their planned attendance at least 15 business days prior to the meeting in order to facilitate their entry into the building. All attendees are required to present government-issued identification prior to entry.

Maria G. Button,

Director, Executive Secretariat.

[FR Doc. 2023–07333 Filed 4–6–23; 8:45 am]

BILLING CODE 4165–15–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Agency Information Collection Activities: Proposed Collection: Public Comment Request; Information Collection Request Title: Rural Maternity and Obstetrics Management Strategies Program

AGENCY: Health Resources and Services Administration (HRSA), Department of Health and Human Services.

ACTION: Notice.

SUMMARY: In compliance with the requirement for opportunity for public comment on proposed data collection projects of the Paperwork Reduction Act of 1995, HRSA announces plans to submit an Information Collection Request (ICR), described below, to the Office of Management and Budget (OMB). Prior to submitting the ICR to OMB, HRSA seeks comments from the public regarding the burden estimate, below, or any other aspect of the ICR.

DATES: Comments on this ICR should be received no later than June 6, 2023.

ADDRESSES: Submit your comments to *paperwork@hrsa.gov* or mail the HRSA Information Collection Clearance Officer, Room 14N136B, 5600 Fishers Lane, Rockville, Maryland 20857.

FOR FURTHER INFORMATION CONTACT: 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 Samantha Miller, the HRSA Information Collection Clearance Officer, at (301) 594–4394.

SUPPLEMENTARY INFORMATION: When submitting comments or requesting information, please include the ICR title for reference.

Information Collection Request Title: Rural Maternity and Obstetrics Management Strategies Program, OMB No. 0906–xxxx–New.

Abstract: HRSA administers the Rural Maternity and Obstetrics Management Strategies (RMOMS) Program, which is authorized by sections 501(a)(2) and 711(b)(5) of the Social Security Act (42 U.S.C. 701(a)(2) and 912(b)(5), respectively), and sections 330A(e) and 330A–2 of the Public Health Service Act (42 U.S.C. 254c(e) and 254c–1b, respectively). These authorities allow HRSA to, among other things, award grants to promote rural health care services outreach by improving and expanding the delivery of health care services to include new and enhanced services in rural areas, through community engagement and evidence-based or innovative, evidence-informed models; as well as establish or continue collaborative improvement and innovation networks to improve access to, and delivery of, maternity and obstetrics care in rural areas.

The RMOMS program grants support networks that improve access to, and continuity of, maternal and obstetrics care in rural communities. The goals of the RMOMS program are to: (1) improve maternal and neonatal outcomes within a rural region; (2) develop a sustainable network approach to increase the delivery and access of preconception, prenatal, pregnancy, labor and delivery, and postpartum services; (3) develop a safe delivery environment with the support and access to specialty care for perinatal patients and infants; and (4) develop sustainable financing models for the provision of maternal and obstetrics care in rural hospitals and communities.

HRSA seeks OMB approval to collect information about RMOMS program grants using performance measures in HRSA's Electronic Handbooks via the Performance Improvement and Measurement System.

Need and Proposed Use of the Information: For this program, performance measures were drafted to provide data to the program and enable HRSA to provide aggregate program data required by Congress under the Government Performance and Results Act of 1993. These measures cover the principal topic areas of interest to the Federal Office of Rural Health Policy, including: (1) consortium/network; (2) sustainability; (3) population demographics; (4) project specific domains. The annual collection of this information helps further inform and substantiate the focus and objectives of the RMOMS program.

Likely Respondents: The respondents will be recipients of the Rural Maternity and Obstetrics Management Strategies Program awards.

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.

TOTAL ESTIMATED ANNUALIZED BURDEN HOURS

Form name	Number of respondents	Number of responses per respondent	Total responses	Average burden per response (in hours)	Total burden hours
Rural Maternity and Obstetrics Management Strategies Program Performance Improvement and Measurement System	10	1	10	9	90
	10	.....	10	.....	90

HRSA specifically 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.

Maria G. Button,

Director, Executive Secretariat.

[FR Doc. 2023–07275 Filed 4–6–23; 8:45 am]

BILLING CODE 4165–15–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute on Deafness and Other Communication Disorders; Notice of Meeting

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended, notice is hereby given of a meeting of the National Deafness and Other Communication Disorders Advisory Council.

The meeting will be open to the public as indicated below, with attendance limited to space available. Individuals who plan to attend as well as those who need special assistance, such as sign language interpretation or other reasonable accommodations, should notify the Contact Person listed below in advance of the meeting. The open session will be videocast and can be accessed from the NIH Videocasting website (*http://videocast.nih.gov/*).

The meeting 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 Deafness and Other Communication Disorders Advisory Council.

Date: May 18–19, 2023.

Closed: May 18, 2023, 11:00 a.m. to 1:00 p.m.

Agenda: To review and evaluate grant applications.

Place: Porter Neuroscience Research Center, Building 35A, Room 640/630, 35 Convent Drive, Bethesda, MD 20892.

Open: May 18, 2023, 2:00 p.m. to 4:40 p.m.

Agenda: Staff reports on divisional, programmatical, and special activities.

Place: Porter Neuroscience Research Center, Building 35A, Room 640/630, 35 Convent Drive, Bethesda, MD 20892.

Open: May 19, 2023, 9:00 a.m. to 11:45 a.m.

Agenda: Staff reports on divisional, programmatical, and special activities.

Place: Porter Neuroscience Research Center, Building 35A, Room 640/630, 35 Convent Drive, Bethesda, MD 20892.

Contact Person: Rebecca Wagenaar-Miller, Ph.D., Director, Division of Extramural Activities, NIDCD/NIH, 6001 Executive Boulevard, Bethesda, MD 20892, (301) 496–8693, *rebecca.wagenaar-miller@nih.gov.*

Any interested person may file written comments with the committee by forwarding the statement to the Contact Person listed on this notice. The statement should include the name, address, telephone number and when applicable, the business or professional affiliation of the interested person.

In the interest of security, NIH has procedures at *https://www.nih.gov/about-nih/visitor-information/campus-access-security* for entrance into on-campus and off-campus facilities. All visitor vehicles, including taxicabs, hotel, and airport shuttles will be inspected before being allowed on campus. Visitors attending a meeting on campus or at an off-campus federal facility will be asked to show one form of identification (for example, a government-issued photo ID, driver's license, or passport) and to state the purpose of their visit.

Information is also available on the Institute's/Center's home page: *https://www.nidcd.nih.gov/about/advisory-council,* where an agenda and any additional information for the meeting will be posted when available.

(Catalogue of Federal Domestic Assistance Program Nos. 93.173, Biological Research Related to Deafness and Communicative Disorders, National Institutes of Health, HHS)

Dated: April 3, 2023.

Tyeshia M. Roberson-Curtis,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2023–07319 Filed 4–6–23; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute of Allergy and Infectious Diseases; Notice of Closed Meeting

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended, notice is hereby given of the following meeting.

The meeting 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 Allergy and Infectious Diseases Special Emphasis Panel; NIH Support for Conferences and Scientific Meetings (Parent R13 Clinical Trial Not Allowed).

Date: May 2–4, 2023.

Time: 9:00 a.m. to 5:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institute of Allergy and Infectious Diseases, National Institutes of Health, 5601 Fishers Lane, Room 3G53, Rockville, MD 20892 (Virtual Meeting).

Contact Person: Konrad Krzewski, Ph.D., Scientific Review Officer, Scientific Review