FDDA’s burden estimate is based on timed readings of each instrument, including the mail and field screeners, enrollment survey, baseline survey, and study A through D questionnaires. Of the total screening respondents, we expect 25 percent will respond only in the mail screening (household deemed ineligible), 65 percent will respond only in the field screening (mail screening nonrespondents), and the remaining 10 percent will respond in both the mail screening and the field screening. The latter includes eligible households from the mail screening that are subsequently field screened to sample the panel member, and the 10 percent quality control sample of households whose mail screening ineligibility is verified through in-person screening. The estimated burden published in the 60-day notice assumed an estimated 10,285 mail and field household screening respondents during yearly panel replenishment and 1,400 additional panel members will be recruited annually as part of the panel replenishment effort, as well as an additional 2,500 household screening respondents during replenishment and an additional 400 panel replenishment enrollment and baseline survey respondents should annual attrition rates be higher than expected.

The following references are on display at the Dockets Management Staff (see ADDRESSES) and are available for viewing by interested persons between 9 a.m. and 4 p.m., Monday through Friday; these are not available electronically at https://www.regulations.gov as these references are copyright protected. Some may be available at the website address, if listed. FDA has verified the website addresses, as of the date this document publishes in the Federal Register, but websites are subject to change over time.

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.

Title of the Collections: The SF–424 Mandatory Form.
Type of Collection: Reinstatement without change.
OMB No.: 4040–0002.
Abstract: The SF–424 Mandatory Form provides the Federal grant-making agencies an alternative to the Standard Form 424 data set and form. Agencies may use the SF–424 Mandatory Form for grant programs not required to collect all the data that is required on the SF–424 core data set and form. The IC expired on January 31, 2019. We are seeking reinstatement of this information collection and a three-year clearance.

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Terry Clark,
Paperwork Reduction Act Reports Clearance Officer, Office of the Secretary.

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute on Deafness and Other Communication Disorders; Notice of Closed Meetings

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended, 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 on Deafness and Other Communication Disorders Special Emphasis Panel; Voice, Speech, and Language Fellowship Review.
Date: June 14, 2019.
Time: 11:00 a.m. to 2:00 p.m.
Agenda: To review and evaluate grant applications.
Place: National Institutes of Health, Neuroscience Center Building (NSC), 6001 Executive Boulevard, Rockville, MD 20852, (Telephone Conference Call).
Contact Person: Eliane Lazar-Wesley, Ph.D., Scientific Review Officer, Scientific Review Branch, Division of Extramural Activities, 6001 Executive Boulevard, Room 8339, MSC 9670, Bethesda, MD 20892–8401, 301–496–8683, elaw@nih.gov.

Name of Committee: National Institute on Deafness and Other Communication Disorders Special Emphasis Panel; Hearing and Balance Fellowship Review.
Date: June 17, 2019.
Time: 8:00 a.m. to 5:00 p.m.
Agenda: To review and evaluate grant applications.
Place: Hyvatt Regency Bethesda, One Bethesda Metro Center, 7400 Wisconsin Avenue, Bethesda, MD 20814.
Contact Person: Shiguang Yang, DVM, Ph.D., Scientific Review Officer, Division of Extramural Activities, NIDCD, NIH, 6001 Executive Blvd., Room 8349, Bethesda, MD 20892, 301–496–8683, yangshi@niddc.nih.gov.

Name of Committee: National Institute on Deafness and Other Communication Disorders Special Emphasis Panel; NICHD Clinical Trial on Cochlear Implants Review.
Date: July 22, 2019.
Time: 12:00 p.m. to 1:30 p.m.
Agenda: To review and evaluate grant applications.
Place: National Institutes of Health, Neuroscience Center Building (NSC), 6001 Executive Boulevard, Rockville, MD 20852, (Telephone Conference Call).
Contact Person: Katherine Shim, Ph.D., Scientific Review Officer, Division of Extramural Activities, NIDCD, 6001 Executive Blvd., Room 8351, Bethesda, MD 20892, 301–496–8683, katherine.shim@nih.gov.

Name of Committee: National Institute on Deafness and Other Communication Disorders Special Emphasis Panel; NICHD.

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Eunice Kennedy Shriver National Institute of Child Health & Human Development; 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 Board of Scientific Counselors, NICHD.

The meeting will be open to the public as indicated below, with the attendance 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 Contact Person listed below in advance of the meeting.

The meeting will be closed to the public as indicated below in accordance with the provisions set forth in section 552b(c)(6), Title 5 U.S.C., as amended for the review, discussion, and evaluation of individual intramural programs and projects conducted by the EUNICE KENNEDY SHRIVER NATIONAL INSTITUTE OF CHILD HEALTH & HUMAN DEVELOPMENT, including consideration of personnel qualifications and performance, and the competence of individual investigators, the disclosure of which would