ESTIMATED ANNUALIZED BURDEN HOURS—Continued

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total burden (in hours)
Pentagon/Shanksville Responder	World Trade Center Health Program, Pentagon/Shanksville Responder Application for Enrollment.	242	1	30/60	121
WTC Survivor	World Trade Center Health Program, Survivor Eligibility Application for Enrollment (all languages).	9,240	1	30/60	4,620
General responder	Clinic Selection Postcard for new general responders in NY/NJ to select a clinic.	3,830	1	15/60	958
Responder/Survivor/Advocate (physician).	Petition for the addition of health conditions.	35	1	1	35
Program Members	Designated Representative Appointment Form.	1,300	1	15/60	325
Program Members	HIPAA Release Form to allow the sharing of member information with a third party.	1,300	1	15/60	325
Program Members	Member Satisfaction Survey	6,600	1	30/60	3,300
General Public	WTCHP HIPAA Authorization for Deceased Individuals.	30	1	15/60	8
General Public	WTCHP General HIPAA Authorization to Third Parties.	30	1	15/60	8
Designated (DR) Representative Revocation Form.	DR form that removes the members current designated representative.	15	1	15/60	4
Total					12,882

Jeffrey M. Zirger,

Lead, Information Collection Review Office, Office of Scientific Integrity, Office of Science, Centers for Disease Control and Prevention.

[FR Doc. 2021–09095 Filed 4–29–21; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Assessing the Implementation and Cost of High-Quality Early Care and Education: Field Test, OMB 0970–0499

AGENCY: Office of Planning, Research, and Evaluation, Administration for Children and Families, HHS.

ACTION: Request for public comment.

SUMMARY: The Administration for Children and Families (ACF) at the U.S. Department of Health and Human Services (HHS) seeks approval to collect new information to use in testing measures of the implementation and costs of high-quality early care and education as part of the project, Assessing the Implementation and Cost of High-Quality Early Care and Education (ECE–ICHQ). The study received approval for a field test to validate and improve the psychometric properties of these measures in November 2019. This request is to add a measure to the approved field test to help further assess the associations between measures of implementation, cost, and quality.

DATES: *Comments due within 60 days of publication.* In compliance with the requirements of the Paperwork Reduction Act of 1995, ACF is soliciting public comment on the specific aspects of the information collection described above.

ADDRESSES: Copies of the proposed collection of information can be obtained, and comments may be forwarded by emailing *OPREinfocollection@acf.hhs.gov.* Alternatively, copies can also be obtained by writing to the Administration for Children and Families, Office of Planning, Research, and Evaluation, 330 C Street SW, Washington, DC 20201, Attn: OPRE Reports Clearance Officer. All requests, emailed or written, should be identified by the title of the information collection.

SUPPLEMENTARY INFORMATION:

Description: ACF seeks approval to collect new information to use in testing measures of the implementation and costs of high-quality early care and education as part of the ECE–ICHQ project. The project's goal is to create a technically sound and feasible instrument that will provide consistent, systematic measures of the implementation and costs of education and care in center-based settings that serve children from birth to age 5. The resulting measures will inform research, policy, and practice by improving understanding of variations in what centers do to support quality, their associated costs, and how resources for ECE may be better aligned with expectations for quality. The study received approval for a field test to validate and improve the psychometric properties of these measures in November 2019. For all previously approved materials for this study, see *https://www.reginfo.gov/public/do/PRAOMBHistory?ombControlNumber=0970-0499.* This request is to add a measure to the approved field test to help further assess the associations between measures of implementation, cost, and quality. The field test and this additional measure will include only remote data collection.

Respondents: Teachers and aids.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents (total over request period)	Number of responses per respondent (total over request period)	Avg. burden per response (in hours)	Total/annual burden (in hours)
Teaching staff survey	1,120	1	.50	560

Estimated Total Annual Burden Hours: 560.

Comments: The Department specifically requests comments on (a) whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology. Consideration will be given to comments and suggestions submitted within 60 days of this publication.

Authority: § 658O(a)(5) as amended by the CCDBG Act of 2014 § 9

Mary B. Jones,

ACF/OPRE Certifying Officer.

[FR Doc. 2021–09052 Filed 4–29–21; 8:45 am]

BILLING CODE 4184–73–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2014–N–0345]

Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Data to Support Drug Product Communications

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA or Agency) is announcing that a collection of information entitled ''Data to Support Drug Product Communications'' has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995 (PRA).

FOR FURTHER INFORMATION CONTACT: Ila S. Mizrachi, Office of Operations, Food and Drug Administration, Three White Flint North, 10A–12M, 11601 Landsdown St., North Bethesda, MD 20852, *301–796–7726, PRAStaff@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION: On February 2, 2020, the Agency submitted a proposed collection of information entitled ''Data to Support Drug Product Communications'' to OMB for review and 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 0910–0695. The approval expires on March 31, 2024. A copy of the supporting statement for this information collection is available on the internet at *https://www.reginfo.gov/public/do/PRAMain.*

Dated: April 26, 2021.

Lauren K. Roth,

Acting Principal Associate Commissioner for Policy.

[FR Doc. 2021–09043 Filed 4–29–21; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute of Dental & Craniofacial Research; 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 Dental and Craniofacial Research Special Emphasis Panel; End the HIV Epidemic.

Date: May 28, 2021.

Time: 3:00 p.m. to 5:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, 6701 Democracy Boulevard, Bethesda, MD 20892 (Virtual Meeting).

Contact Person: Yun Mei, MD, Scientific Review Officer, Scientific Review Branch, Natl Institute of Dental and Craniofacial Research, National Institutes of Health, 6701 Democracy Boulevard, Suite #670, Bethesda, MD 20892, (301) 827–4639, *yun.mei@nih.gov.*

(Catalogue of Federal Domestic Assistance Program No. 93.121, Oral Diseases and Disorders Research, National Institutes of Health, HHS)

Dated: April 26, 2021.

Melanie J. Pantoja,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2021–09041 Filed 4–29–21; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute on Aging; 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 Aging Initial Review Group Career Development Facilitating the Transition to Independence.

Date: June 10–11, 2021.

Time: 10:00 a.m. to 6:00 p.m.