specific aspects of the information collection described above.

ADDRESSES: You can obtain copies of the proposed collection of information and submit comments by emailing *OPREinfocollection@acf.hhs.gov.* Identify all requests by the title of the information collection.

SUPPLEMENTARY INFORMATION:

Description: OPRE is conducting the BIAS–NG project, which uses behavioral insights to design and test interventions intended to improve the efficiency, operations, and efficacy of human services programs. The BIAS–NG project is applying and testing behavioral insights to ACF programs including TANF, child welfare, and EHS/HS, and intends to expand these efforts to child care. This notice is a request for comments on ACF's proposal to revise and extend a previously approved collection, which included data collection to design and test interventions in the TANF, child welfare, and EHS/HS domains. Under the approved pilot generic clearance, OPRE has already conducted work with seven sites to conduct seven tests, and is planning to continue to work with at least one additional site, conducting one or more tests of behavioral interventions for a total of nine tests of behavioral interventions. All approved information collection activities can be found here: *https://www.reginfo.gov/public/do/PRAICList?ref_nbr=202206-0970-002.*

In addition to extending approval, this approval would also allow OPRE to conduct tests in the newly added program area of child care. The design and testing of BIAS–NG interventions is rapid and, to the extent possible, iterative. Each specific intervention is designed in consultation with agency leaders and launched as quickly as possible. To maximize the likelihood that the intervention produces measurable, significant, and positive effects on outcomes of interest, rapid cycle evaluation techniques will be employed in which proximate outcomes will be measured to allow the research team to more quickly iterate and adjust the intervention design, informing subsequent tests. Due to the rapid and iterative nature of this work, OPRE sought and received approval for an overarching generic clearance to conduct this research. Following standard OMB requirements for generic clearances, once instruments subject to PRA are tailored to a specific site and the site's intervention, OPRE submits an individual generic information collection request under this umbrella clearance. Each request includes the individual instrument(s), a justification specific to the individual information collection, a description of the proposed intervention, and any supplementary documents. Each specific information collection includes up to two submissions—one submission for the formative stage research and another submission for any further data collection requiring burden during the testing phase. The type of information to be collected and the uses of the information is described in the supporting statements, found here: *https://www.reginfo.gov/public/do/PRAViewDocument?ref_nbr=202206-0970-002.*

Respondents: (1) Program Administrators, (2) Program Staff, and (3) Program Clients.

ANNUAL BURDEN ESTIMATES (TANF, CHILD WELFARE, EHS/HS, CHILD CARE)

Instrument	Number of respondents (TANF, CW, EHS/HS, CC) (total over request period)	Number of responses per respondent (total over request period)	Average burden hours per response (in hours)	Total burden (in hours)	Annual burden (in hours)
Phase 3: Diagnosis and Design					
Administrator interviews/focus groups	48	1	1	48	16
Staff interviews/focus groups	400	1	1	400	133
Client interviews/focus groups	400	1	1	400	133
Client survey	400	1	.25	100	33
Staff Survey	400	1	.25	100	33
Phase 4: Evaluation					
Administrator interviews/focus groups	96	1	1	96	32
Staff interviews/focus groups	800	1	1	800	267
Client interviews/focus groups	800	1	1	800	267
Client survey	12,000	1	.25	3,000	1,000
Staff Survey	1,200	1	.25	300	100

Estimated Total Annual Burden Hours: 2,014.

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: 42 U.S.C. 1310.

Mary C. Jones,

ACF/OPRE Certifying Officer.

[FR Doc. 2024–11077 Filed 5–20–24; 8:45 am]

BILLING CODE 4184–07–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Statement of Organization, Functions, and Delegations of Authority

AGENCY: Office of Population Affairs, Office of the Assistant Secretary for Health, Office of the Secretary, Department of Health and Human Services (HHS).

ACTION: Notice.

SUMMARY: The Office of the Assistant Secretary for Health, Office of Population Affairs (OPA) has modified its organizational structure.

DATES: This new organizational structure was approved by the Secretary of Health and Human Services and takes effect on May 14, 2024.

FOR FURTHER INFORMATION CONTACT: Jessica Swafford Marcella, Deputy Assistant Secretary for Population Affairs, Office of Population Affairs, Office of the Assistant Secretary for Health, Department of Health and Human Services at *Jessica.marcella@hhs.gov* and 240–453–2800.

SUPPLEMENTARY INFORMATION: Part A (Office of the Secretary, U.S. Department of Health and Human Services) of the Statement of Organization, Functions, and Delegations of Authority of the Department of Health and Human Services (60 FR 56605, dated November 9, 1995, and corrected at 75 FR 53304, August 31, 2010, amended at 82 FR 3005, dated January 10, 2017, and amended, most recently at 84 FR 14951, dated April 12, 2019) is amended to reflect the reorganization of the Office of Population Affairs (OPA), Office of the Assistant Secretary for Health (OASH). This reorganization will streamline operations and improve the efficiency and effectiveness of OPA. OPA is responsible for implementing and administering the Title X family planning program, managing the Office of Adolescent Health, which is headed by a director, and implementing and administering the Teen Pregnancy Prevention program and other adolescent health activities. The changes are as follows:

I. *Under Part C, section C–G, Organization, revise to organize OPA staff across six Divisions:* Administration and Operations, Adolescent Health Programs, Clinical and Scientific Affairs, Policy and External Affairs, Research and Evaluation, and Title X Service Delivery. The Division of Administration and Operations will manage overall operations for OPA, including staff recruitment and retention, budgeting, training, and travel. The Division of Adolescent Health Programs will lead implementation of the Teen Pregnancy Prevention Program and other adolescent health programmatic activities. The Division of Clinical and Scientific Affairs will provide clinical and scientific expertise, consultation, and oversight for all OPA staff and activities. The Division of Policy and External Affairs will lead all policy and communication activities for the office. The Division of Research and Evaluation will oversee data collection for OPA programs and will lead OPA's research and evaluation activities. The Division of Title X Service Delivery will lead implementation of the Title X service delivery program.

II. *Delegations of Authority:* All delegations and redelegations of authority made to officials and employees of affected organizational components will continue in them or their successors pending further redelegations, if allowed, provided they are consistent with this reorganization.

(Authority: 44 U.S.C. 3101)

Xavier Becerra,

Secretary, U.S. Department of Health and Human Services.

[FR Doc. 2024–11058 Filed 5–20–24; 8:45 am]

BILLING CODE P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute of Neurological Disorders and Stroke; Notice of Closed Meetings

Pursuant to section 1009 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: Neurological Sciences Training Initial Review Group; NST–4 Study Section.

Date: June 14, 2024.

Time: 9:30 a.m. to 6:30 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Neuroscience Center, 6001 Executive Boulevard, Rockville, MD 20852 (Virtual Meeting).

Contact Person: Steven G Britt, MD, Scientific Review Officer, Scientific Review Branch, Division of Extramural Activities, NINDS/NIH/HHS NSC, 6001 Executive Boulevard, Rockville, MD 20852, 301–480–1953, *steve.britt@nih.gov*.

Name of Committee: National Institute of Neurological Disorders and Stroke Special Emphasis Panel; Review of UE5 Training Applications.

Date: July 12, 2024.

Time: 2:30 p.m. to 6:30 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institute of Health, Neuroscience Center, 6001 Executive Boulevard, Rockville, MD 20852 (Virtual Meeting).

Contact Person: William C. Benzing, Ph.D., Scientific Review Officer, Scientific Review Branch, Division of Extramural Activities, NINDS/NIH/HHS NSC, 6001 Executive Boulevard, Rockville, MD 20852, 301–496–0660, *benzingw@mail.nih.gov*.

(Catalogue of Federal Domestic Assistance Program Nos. 93.853, Clinical Research Related to Neurological Disorders; 93.854, Biological Basis Research in the Neurosciences, National Institutes of Health, HHS).

Dated: May 15, 2024.

Lauren A. Fleck,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2024–11067 Filed 5–20–24; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute on Drug Abuse; Notice of Closed Meetings

Pursuant to section 1009 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 Drug Abuse Special Emphasis Panel; HEAL Initiative: Translating Research To Practice to End the Overdose Crisis.

Date: June 27, 2024.

Time: 10:00 a.m. to 6:30 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institute of Health, National Institute on Drug Abuse, 301 North Stonestreet Avenue, Bethesda, MD 20892 (Virtual Meeting).

Contact Person: Sheila Pirooznia, Ph.D., Scientific Review Officer, Division of Extramural Review, Scientific Review Branch, National Institute on Drug Abuse, NIH, 301 North Stonestreet Avenue, MSC,