clinical management of the cycle, which would include (but is not limited to) multiple aspects of the treatment such as patient selection, pre-treatment counseling and selection of the specific treatment protocol. The ART programs involved must have a method in place to ensure that these cycles can be prospectively reported by the ART program required to report them. In addition, all canceled cycles must be reported by the same ART program.

b. Cycles involving previously cryopreserved oocytes/embryos are to be reported by the ART program that accepts responsibility for thawing the oocytes/embryos.

Dated: June 7, 2022.

Angela K. Oliver,

Executive Secretary, Centers for Disease Control and Prevention.

[FR Doc. 2022–12528 Filed 6–9–22; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Behavioral Interventions To Advance Self-Sufficiency Next Generation (BIAS–NG) (OMB# 0970–0502)

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

ACTION: Request for public comment.

SUMMARY: The Office of Planning, Research, and Evaluation (OPRE) in the Administration for Children and Families (ACF), U.S. Department of Health and Human Services (HHS), requests Office of Management and Budget (OMB) approval to extend approval of the ACF Behavioral Interventions to Advance Self-Sufficiency Next Generation (BIAS–NG) Project Overarching Generic (OMB #: 0970–0502; Expiration date: 8/31/2022). Under this overarching generic, ACF collects data as part of rapid cycle testing and evaluation, in order to inform the design of interventions informed by behavioral science and to better understand the mechanisms and effects of such interventions. Interventions have been and will continue to be developed in the program area domains of Temporary Assistance for Needy Families (TANF), child welfare, and Early Head Start/Head Start (EHS/HS). These interventions are intended to improve outcomes for participants in these programs. No changes are proposed.

DATES: *Comments due within 30 days of publication.* OMB must make a decision about the collection of information between 30 and 60 days after publication of this document in the **Federal Register**. Therefore, a comment is best assured of having its full effect if OMB receives it within 30 days of publication.

ADDRESSES: Written comments and recommendations for the proposed information collection should be sent within 30 days of publication of this notice to *www.reginfo.gov/public/do/PRAMain.* Find this particular information collection by selecting ''Currently under 30-day Review—Open for Public Comments'' or by using the search function. You can also obtain copies of the proposed collection of information 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. This notice is a request for comments on ACF's proposal to extend approval of the overarching generic. Under the approved pilot generic clearance, OPRE has already completed work with five sites and has conducted five tests. The extended approval would allow OPRE to continue to work with at least three additional sites, conducting one or more tests of behavioral interventions. 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 generic clearance to conduct this research. Following standard OMB requirements for generic clearances, once instruments requiring burden 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/PRAViewICR?ref_nbr=201909-0970-003.*

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

Annual Burden Estimates (TANF, CW, EHS/HS): This request includes an extension to complete currently approved and ongoing phase 3 data collection in three sites (Matrix/Starfish and Hennepin County), and new data collection. Burden estimates for new requests are outlined below. Previously approved burden estimates can be found at the url above.

Instrument	Number of respondents (TANF, CW, EHS/HS) (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	0.25	100	33

Instrument	Number of respondents (TANF, CW, EHS/HS) (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)
Staff Survey	400	1	0.25	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	0.25	3,000	1,000
Staff Survey	1,200	1	0.25	300	100

Estimated Total Annual Burden Hours: 2,014.

Authority: 42 U.S.C. 1310.

Mary B. Jones,

ACF/OPRE Certifying Officer.

[FR Doc. 2022–12538 Filed 6–9–22; 8:45 am]

BILLING CODE 4184–07–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Center for Scientific Review; 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: Center for Scientific Review Special Emphasis Panel; Small Business: Radiation Therapy and Biology SBIR/STTR.

Date: July 7, 2022.

Time: 9:00 a.m. to 6:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Rockledge II, 6701 Rockledge Drive, Bethesda, MD 20892 (Virtual Meeting).

Contact Person: Bo Hong, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 6194, MSC 7804, Bethesda, MD 20892, 301–996–6208, *hongb@csr.nih.gov*.

Name of Committee: Infectious Diseases and Immunology B Integrated Review Group; HIV Comorbidities and Clinical Studies Study Section.

Date: July 12–13, 2022.

Time: 9:00 a.m. to 8:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Rockledge II, 6701 Rockledge Drive, Bethesda, MD 20892 (Virtual Meeting).

Contact Person: David C. Chang, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892, (301) 451–0290, *changdac@mail.nih.gov*.

Name of Committee: Center for Scientific Review Special Emphasis Panel; Small Business: Neuropathology, Developmental Disability, and Stem Education.

Date: July 13–14, 2022.

Time: 10:00 a.m. to 8:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Rockledge II, 6701 Rockledge Drive, Bethesda, MD 20892 (Virtual Meeting).

Contact Person: Heidi B. Friedman, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 1012A, MSC 7770, Bethesda, MD 20892, (301) 379–5632, *hfriedman@csr.nih.gov*.

Name of Committee: Center for Scientific Review Special Emphasis Panel; Fellowships: HIV/AIDS Biological.

Date: July 14, 2022.

Time: 10:00 a.m. to 8:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Rockledge II, 6701 Rockledge Drive, Bethesda, MD 20892 (Virtual Meeting).

Contact Person: Raj K. Krishnaraju, Ph.D., MS Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 6190, MSC 7804, Bethesda, MD 20892, (301) 435–1047, *kkrishna@csr.nih.gov*.

Name of Committee: Center for Scientific Review Special Emphasis Panel; Member Conflict: Motivated Behavior, Alcohol and Neurotoxicology.

Date: July 14, 2022.

Time: 11:00 a.m. to 6:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Rockledge II, 6701 Rockledge Drive, Bethesda, MD 20892 (Virtual Meeting).

Contact Person: Kirk Thompson, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 5184, MSC 7844, Bethesda, MD 20892, 301–435–1242, *kgt@mail.nih.gov*.

Name of Committee: Center for Scientific Review Special Emphasis Panel; Member Conflict: Metabolic Live Disease and Regeneration.

Date: July 15, 2022.

Time: 12:00 p.m. to 6:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Rockledge II, 6701 Rockledge Drive, Bethesda, MD 20892 (Virtual Meeting).

Contact Person: Ganesan Ramesh, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 2182, MSC 7818, Bethesda, MD 20892, (301) 827–5467, *ganesan.ramesh@nih.gov*.

Name of Committee: Center for Scientific Review Special Emphasis Panel; Small Business: Biomedical Sensing, Measurement and Instrumentation.

Date: July 19–20, 2022.

Time: 9:00 a.m. to 8:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Rockledge II, 6701 Rockledge Drive, Bethesda, MD 20892 (Virtual Meeting).

Contact Person: Michael L Bloom, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 6187, MSC 7804, Bethesda, MD 20892, 301–451–0132, *bloomm2@mail.nih.gov*.

Name of Committee: Center for Scientific Review Special Emphasis Panel; Small Business: Innovative Immunology.

Date: July 20, 2022.

Time: 10:00 a.m. to 8:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Rockledge II, 6701 Rockledge Drive, Bethesda, MD 20892 (Virtual Meeting).

Contact Person: Dayadevi Jirage, Ph.D. Scientific Review Officer, Center for Scientific Review, National Institutes of Health 6701 Rockledge Drive, Room 809–H,