

for health care services rendered to Medicare beneficiaries. CMS requires that providers follow reasonable cost principles under 1861(v)(1)(A) of the Act when completing the Medicare cost report (MCR). Regulations at 42 CFR 413.20 and 413.24 require that providers submit acceptable cost reports on an annual basis and maintain sufficient financial records and statistical data, capable of verification by qualified auditors.

ESRD facilities participating in the Medicare program submit these cost reports annually to report cost and statistical data used by CMS to determine reasonable costs incurred for furnishing dialysis services to Medicare beneficiaries and to effect the year-end cost settlement for Medicare bad debts. *Form Number:* CMS–265–11 (OMB control number: 0938–0236); *Frequency:* Annually; *Affected Public:* Private Sector, Business or other for-profits, State, Local, or Tribal Governments); *Number of Respondents:* 7,492; *Total Annual Responses:* 7,492; *Total Annual Hours:* 494,472. (For questions regarding this collection contact Keplinger, Jill C at 410–786–4550.)

5. *Type of Information Collection Request:* Reinstatement without change; *Title of Information Collection:* Good Cause Processes; *Use:* Section 1851(g)(3)(B)(i) of the Act provides that MA organizations may terminate the enrollment of individuals who fail to pay basic and supplemental premiums after a grace period established by the plan. Section 1860D–1(b)(1)(B)(v) of the Act generally directs us to establish rules related to enrollment, disenrollment, and termination for Part D plan sponsors that are similar to those established for MA organizations under section 1851 of the Act. Consistent with these sections of the Act, subpart B in each of the Parts C and D regulations sets forth requirements with respect to involuntary disenrollment procedures at 42 CFR 422.74 and 423.44, respectively. In addition, section 1876(c)(3)(B) establishes that individuals may be disenrolled from coverage as specified in regulations. Thus, current regulations at 42 CFR 417.460 specify that a cost plan, specifically a Health Maintenance Organization (HMO) or competitive medical plan (CMP), may disenroll a member who fails to pay premiums or other charges imposed by the plan for deductible and coinsurance amounts.

These good cause provisions authorize CMS to reinstate a disenrolled individual's enrollment without interruption in coverage if the non-payment is due to circumstances that the individual could not reasonably foresee or could not control, such as an

unexpected hospitalization. At its inception, the process of accepting, reviewing, and processing beneficiary requests for reinstatement for good cause was carried out exclusively by CMS. *Form Number:* CMS–10544 (OMB control number: 0938–1271); *Frequency:* Annually; *Affected Public:* Business or other for-profits State, Local, or Tribal Governments); *Number of Respondents:* 312; *Total Annual Responses:* 41,289; *Total Annual Hours:* 27,499. (For questions regarding this collection contact Fabayo, Ronke at (410) 786–4460.)

Dated: February 16, 2022.

William N. Parham, III,
Director, Paperwork Reduction Staff, Office of Strategic Operations and Regulatory Affairs.

[FR Doc. 2022–03727 Filed 2–18–22; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

OMB No. 0970–0502

Proposed Information Collection Activity; Behavioral Interventions To Advance Self-Sufficiency Next Generation

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 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: 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. 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/PRAViewDocument?ref_nbr=201707-0970-005.

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

ANNUAL BURDEN ESTIMATES
[TANF, CW, EHS/HS]

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
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.

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 B. Jones,

ACF/OPRE Certifying Officer.

[FR Doc. 2022-03669 Filed 2-18-22; 8:45 am]

BILLING CODE 4184-07-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 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 of Neurological Disorders and Stroke Special Emphasis Panel; Detecting Cognitive Impairment, Including Dementia, in Primary Care and Other Everyday Clinical Settings for the General Public and Health Equity, Pragmatic Clinical Trials (U01 Clinical Trial Required).

Date: March 29, 2022.

Time: 10:00 a.m. to 5:00 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: Ana Olariu, Ph.D., Scientific Review Officer, Scientific Review Branch, Division of Extramural Activities, NINDS/NIH, NSC, 6001 Executive Boulevard, Room 3208, MSC 9529, Bethesda, MD 20892, 301-496-9223 Ana.Olariu@nih.gov.

Name of Committee: National Institute of Neurological Disorders and Stroke Special Emphasis Panel; BRAIN Initiative Advanced Postdoctoral Career Transition Award to

Promote Diversity (K99/R00 Independent Clinical Trial Not Allowed).

Date: March 30, 2022.

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

Agenda: To review and evaluate grant applications and/or proposals.

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

Contact Person: Lataisia Cherie Jones, Ph.D., Scientific Review Officer, Scientific Review Branch, Division of Extramural Activities, NINDS/NIH, NSC, 6001 Executive Boulevard, Suite 3208, Rockville, MD 20852, 301-496-9223, lataisia.jones@nih.gov.

Name of Committee: National Institute of Neurological Disorders and Stroke Special Emphasis Panel; Diversity K01 & MOSAIC Postdoctoral Career Transition Award to Promote Diversity (K99/R00 Independent Clinical Trial Not Allowed).

Date: March 31, 2022.

Time: 9:00 a.m. to 6:00 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: Lataisia Cherie Jones, Ph.D., Scientific Review Officer, Scientific Review Branch, Division of Extramural Activities, NINDS/NIH, NSC, 6001 Executive Boulevard, Suite 3208, Rockville, MD 20852, 301-496-9223, lataisia.jones@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)