Provide Medicare protection to individuals who are members of a Buy-in coverage group, as specified in the State’s Buy-in agreement. The Code of Federal Regulations at 42 CFR Section 407.40 provides for States to enroll in Medicare and pay the premiums for all eligible members covered under a Buy-in coverage group. Individuals enrolled in Medicare through the Buy-in program must be eligible for Medicare and be an eligible member of a Buy-in coverage group. The day to day operations of the State Buy-in program is accomplished through an automated data exchange process. The automated data exchange process is used to exchange Medicare and Buy-in entitlement information between the Social Security District Offices, State Medicaid Agencies and the Centers for Medicare & Medicaid Services (CMS). When problems arise that cannot be resolved through the normal data exchange process, clerical actions are required. The CMS–1957, “SSO Report of State Buy-In Problem” is used to report Buy-in problems cases. The CMS–1957 is the only standardized form available for communications between the aforementioned agencies for the resolution of beneficiary complaints and inquiries regarding State Buy-in eligibility.

**Bill: State, Local, or Tribal Governments; Number of Respondents:** 5,854; **Total Annual Responses:** 5,854; **Total Annual Hours:** 1,951. (For policy questions regarding this collection contact Keith Johnson at (410) 786–1148.)

**Dated:** May 21, 2019.

**William N. Parham, III,**

Director, Paperwork Reduction Staff, Office of Strategic Operations and Regulatory Affairs.

[FR Doc. 2019–10963 Filed 5–23–19; 8:45 am]

**BILLING CODE 4120–01–P**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Centers for Medicare & Medicaid Services**


**Agency Information Collection Activities: Submission for OMB Review; Comment Request**

**AGENCY:** Centers for Medicare & Medicaid Services, HHS.

**ACTION:** Notice.

**SUMMARY:** The Centers for Medicare & Medicaid Services (CMS) is announcing an opportunity for the public to comment on CMS’ intention to collect information from the public. Under the Paperwork Reduction Act of 1995 (PRA), federal agencies are required to publish notice in the Federal Register concerning each proposed collection of information, including each proposed extension or reinstatement of an existing collection of information, and to allow a second opportunity for public comment on the notice. Interested persons are invited to send comments regarding the burden estimate or any other aspect of this collection of information, including the necessity and utility of the proposed information collection for the proper performance of the agency’s functions, the accuracy of the estimated burden, ways to enhance the quality, utility, and clarity of the information to be collected, and the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

**DATES:** Comments on the collection(s) of information must be received by the OMB desk officer by June 24, 2019.

**ADDRESSES:** When commenting on the proposed information collections, please reference the document identifier or OMB control number. To be assured consideration, comments and recommendations must be received by the OMB desk officer via one of the following transmissions: OMB, Office of Information and Regulatory Affairs, Attention: CMS Desk Officer, Fax Number: (202) 395–5806 OR, Email: OIRA_submission@omb.eop.gov.

To obtain copies of a supporting statement and any related forms for the proposed collection(s) summarized in this notice, you may make your request using one of following:

2. Email your request, including your address, phone number, OMB number, and CMS document identifier, to Paperwork@cms.hhs.gov.
3. Call the Reports Clearance Office at (410) 786–1326.

**FOR FURTHER INFORMATION CONTACT:**

William N. Parham at (410) 786–4669.

**SUPPLEMENTARY INFORMATION:**

This notice sets out a summary of the use and burden associated with the following information collections. More detailed information can be found in each collection’s supporting statement and associated materials (see ADDRESSES).


Under the PRA (44 U.S.C. 3501–3520), federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. The term “collection of information” is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA requires federal agencies to publish a 60-day notice in the Federal Register concerning each proposed collection of information, including each proposed extension or reinstatement of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, CMS is publishing this notice.

**Information Collection**

1. **Type of Information Collection Request:** Extension without change of a currently approved collection; Title of Information Collection: Social Security Office Report of State Buy-in Problem; **Use:** The statutory authority for the State Buy-in program is Section 1843 of the Social Security Act, amended through 1989. Under Section 1843, a State can enter into an agreement to provide Medicare protection to individuals who are members of a Buy-in coverage group, as specified in the State’s Buy-in agreement. The Code of Federal Regulations at 42 CFR Section 407.40 provides for States to enroll in Medicare and pay the premiums for all eligible members covered under a Buy-in coverage group. Individuals enrolled in Medicare through the Buy-in program must be eligible for Medicare and be an eligible member of a Buy-in coverage group. The day to day operations of the State Buy-in program is accomplished through an automated data exchange process. The automated data exchange process is used to exchange Medicare and Buy-in entitlement information between the Social Security District Offices, State Medicaid Agencies and the Centers for Medicare & Medicaid Services (CMS). When problems arise that cannot be resolved through the normal data exchange process, clerical actions are required. The CMS–1957, “SSO Report of State Buy-In Problem” is used to report Buy-in problems cases. The CMS–1957 is the only standardized form available for communications between the aforementioned agencies for the resolution of beneficiary complaints and inquiries regarding State Buy-in eligibility. **Form Number:** CMS–1957 (OMB control number: 0938–0035); **Frequency:** Yearly; **Affected Public:** State, Local, or Tribal Governments; **Number of Respondents:** 5,854; **Total Annual Responses:** 5,854; **Total Annual Hours:** 1,951. (For policy questions regarding this collection contact Keith Johnson at (410) 786–1148.)

**Dated:** May 21, 2019.

**William N. Parham, III,**

Director, Paperwork Reduction Staff, Office of Strategic Operations and Regulatory Affairs.

[FR Doc. 2019–10963 Filed 5–23–19; 8:45 am]

**BILLING CODE 4120–01–P**
includes agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA (44 U.S.C. 3506(c)(2)(A)) requires federal agencies to publish a 30-day notice in the Federal Register concerning each proposed collection of information, including each proposed extension or reinstatement of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, CMS is publishing this notice that summarizes the following proposed collection(s) of information for public comment:

1. Type of Information Collection Request: Extension of a currently approved collection; Title of Information Collection: Medicaid Quality Assessment and Performance Improvement Programs, State Review of Accreditation Status, Medicaid Managed Care Quality Rating System, and Quality Strategy (QS) and Supporting Regulations; Use: Medicaid beneficiaries and stakeholders use the information collected and reported to understand the state’s quality improvement goals and objectives, and to understand how the state is measuring progress on its goals. States use this information to help monitor and assess the performance of their Medicaid managed care programs. This information may assist states in comparing the outcomes of quality improvement efforts and can assist them in identifying future performance improvement goals. CMS uses this information as a part of its oversight of Medicaid programs. Form Number: CMS–10553 (OMB control number: 0938–1281); Frequency: Yearly and occasionally; Affected Public: Private sector (business or other for-profits) and State, Local, or Tribal Governments; Number of Respondents: 603; Total Annual Responses: 6,441; Total Annual Hours: 52,343. (For policy questions regarding this collection contact Barbara Dailey at 410–786–9012.)

2. Type of Information Collection Request: Reinstatement; Title of Information Collection: End Stage Renal Disease Death Notification; Use: The ESRD Death Notification form (CMS–2746) is completed by all Medicare-approved ESRD facilities upon death of an ESRD patient. Its primary purpose is to collect fact of death and cause of death of ESRD patients. The ESRD Program Management and Medical Information System (PMMIS) has the responsibility of collecting, maintaining and disseminating, on a national basis, uniform data pertaining to ESRD patients and their treatment of care. All renal facilities approved to participate in the ESRD program are required by Public Law 95–292 to supply data to this system.

Federal regulations require that the ESRD Networks examine the mortality rates of every Medicare-approved facility within its area of responsibility. CMS–2746 provides the necessary data to assist the ESRD Networks in making decisions that result in improved patient care and in cost-effective distribution of ESRD resources. The data is used by the ESRD Networks to verify facility deaths and to monitor facility performance. The form is also used by health care planning agencies and researchers to determine survival rates by diagnoses. Form Number: CMS–2746 (OMB control number: 0938–0448); Frequency: Yearly; Affected Public: Private Sector (Business or other for-profits, Not-for-Profit Institutions); Number of Respondents: 7,311; Total Annual Responses: 92,023; Total Annual Hours: 46,011.50. (For policy questions regarding this collection contact Gequinica Polk at 410–786–2305.)

3. Type of Information Collection Request: Reinstatement; Title of Information Collection: End Stage Renal Disease Medical Evidence Report Medicare Entitlement and/or Patient Registration; Use: In accordance with section 226A of the law, the primary purpose of this form is to have a patient medically determined, by a physician, to have end stage renal disease for purposes of filing for Medicare benefits. The End Stage Renal Disease (ESRD) Medical Evidence (CMS–2728) is completed for all ESRD patients either by the first treatment facility or by a Medicare-approved ESRD facility when it is determined by a physician that the patient’s condition has reached that stage of renal impairment that a regular course of kidney dialysis or a kidney transplant is necessary to maintain life. The data reported on the CMS–2728 is used by the Federal Government, ESRD Networks, treatment facilities, researchers and others to monitor and assess the quality and type of care provided to end stage renal disease beneficiaries. Collection of these data are also necessary for entitlement of ESRD patients to Medicare benefits and also for the establishment and maintenance of a single, nationwide kidney disease registry for dialysis, transplant, and prospective transplant patients, and will store pertinent medical facts on each registrant. The data will enable individual practitioners and facility staff to manage care and improve ESRD patient treatment methods, which will permit local Medical Review Boards to more effectively monitor utilization and quality of medical care. Form Number: CMS–2728 (OMB control number: 0938–0046); Frequency: Yearly; Affected Public: Private Sector (Business or other for-profits, Not-for-Profit Institutions); Number of Respondents: 7,311; Total Annual Responses: 138,000; Total Annual Hours: 103,500. (For policy questions regarding this collection contact Gequinica Polk at 410–786–2305.)

4. Type of Information Collection Request: Revision of a currently approved collection; Title of Information Collection: The HIPAA Eligibility Transaction System (HETS); Use: HIPAA regulations require covered entities to verify the identity of the person requesting PHI and the person’s authority to have access to that information. Under the HIPAA Security rules, covered entities, regardless of their size, are required under 45 CFR Subtitle A, Subpart C 164.312(a)(2)(i) to “assign a unique name and/or number for identifying and tracking user identity.” A 'user' is defined in 164.304 as a “person or entity with authorized access”. Accordingly, the HIPAA Security rule requires covered entities to assign a unique name and/or number to each employee or workforce member who uses a system that receives, maintains or transmits electronic PHI so that system access and activity can be identified and tracked by user. This pertains to workforce members within small or large provider offices, health plans, group health plans, and clearinghouses. Federal law requires that CMS take precautions to minimize the security risk to the federal information system. Federal Information Processing Standards Publication (FIPS PUB) 1(1) 1–2 Paragraph 11.7—Security and Authentication states that: ‘‘Agencies shall employ risk management techniques to determine the appropriate mix of security controls needed to protect specific data and systems. The selection of controls shall take into account procedures required under applicable laws and regulations.’’ Accordingly, CMS requires that entities who wish to connect to the HETS application via the CMS Extranet and/or internet are uniquely identified. CMS is required to verify the identity of the person requesting the Protected Health Information (PHI) and the person’s authority to have access to Medicare eligibility information. Furthermore, CMS requires that trading partners who wish to conduct eligibility transactions on a real-time basis with CMS provide certain assurances as a condition of
receiving access to the Medicare eligibility information for the purpose of conducting real-time 270/271 inquiry/response transactions. Form Number: CMS–10157 (OMB control number: 0938–0960); Frequency: Yearly; Affected Public: Private Sector; Business or other for profits, Not-for-Profits Institutions; Number of Respondents: 1000; Total Annual Responses: 1000; Total Annual Hours: 250. (For policy questions regarding this collection contact Rupinder Singh at 410 786–7484.)

Dated: May 21, 2019.

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

[FR Doc. 2019–10964 Filed 5–23–19; 8:45 am]
BILLING CODE 4120–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

[CFDA Number: 93.647]

Announcement of an Unsolicited Single-Source Grant Award to the Woodson Center in Washington, DC

AGENCY: Office of Planning, Research and Evaluation, Administration for Children and Families, Department of Health and Human Services.

ACTION: Notice of issuance.

SUMMARY: The Administration for Children and Families (ACF), Office of Planning, Research and Evaluation announces the award of a grant in the amount of $150,000 to the Woodson Center plans to complete an environmental scan of privately managed foster care systems in multiple states. This environmental scan is expected to inform future networking efforts among similar organizations and identification and dissemination of best practices.

Statutory Authority: Social Security Act, Title XI, Section 1110, 42 U.S.C. 1310.

Elizabeth Leo,
Senior Grants Policy Specialist, Division of Grants Policy, Office of Administration.

[FR Doc. 2019–10897 Filed 5–23–19; 8:45 am]
BILLING CODE 4184–07–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families


AGENCY: Office of Community Services; Administration for Children and Families; HHS.

ACTION: Request for public comment.

SUMMARY: The Office of Community Services (OCS), Administration for Children and Families (ACF) is requesting approval of a three-year extension of the Assets for Independence (AFI) Performance Progress Report (PPR) Long Form and AFI PPR Short Form (OMB #0970–0483, expiration 8/31/2019). There are no changes requested to the forms. DATES: Comments due within 60 days of publication. In compliance with the requirements of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Administration for Children and Families 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 infocollection@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:


This request is for approval and extension of the current AFI PPR that expires August 31, 2019. OCS will continue to use the data collected in the AFI PPR to prepare the annual AFI Report to Congress, to evaluate and monitor the performance of the AFI program overall and of individual projects, and to inform and support technical assistance efforts. The AFI PPR will continue to fulfill AFI Act reporting requirements and program purposes.


Note: This request does not affect financial reporting requirements for AFI grantees. The SF–425 will still be required semiannually throughout the grant project period with a final report due 90 days after the grant project period ends.

Respondents: Assets for Independence (AFI) program grantees.

ANNUAL BURDEN ESTIMATES

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Estimated Total Annual Burden Hours: 623.5.

Comments: The Department specifically requests comments on (a) whether the proposed collection of information is necessary for the proper