ESTIMATED ANNUALIZED BURDEN HOURS—Continued

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)
Data manager at non-STD clinic health centers.	Non-STD clinic Data Elements	26	6	3
Public Health Laboratory Microbiologist	Laboratory Testing Data Elements	8	700	10/60
Public Health Laboratory Data Manager	Laboratory Data Elements	8	6	1
Gonorrhea Patients and Sexual Contacts	Field Investigation Data Elements	960	1	20/60

Jeffrey M. Zirger,

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

[FR Doc. 2021–14751 Filed 7–9–21; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

[CFDA Number: 93.676]

Announcement of Intent To Issue One Operating Division (OPDIV)-Initiated Supplement to BCFS Health and Human Services Under the Standing Announcement for Residential (Shelter) Services for Unaccompanied Alien Children, HHS–2017–ACF–ORR–ZU–1132

AGENCY: Unaccompanied Alien Children's (UAC) Program, Office of Refugee Resettlement (ORR), Administration for Children and Families (ACF), Department of Health and Human Services (HHS).

ACTION: Notice of intent to issue one OPDIV-Initiated Supplement to BCFS Health and Human Services (BCFS HHS), San Antonio, Texas under the UAC Program.

SUMMARY: ACF, ORR, announces the issuance of one OPDIV-Initiated Supplement to BCFS HHS, San Antonio, Texas in the amount of up to $475,868,102. ORR has been identifying additional capacity to provide shelter for potential increases in apprehensions of UAC at the Southwest Border. Planning for increased shelter capacity is a prudent step to ensure that ORR is able to meet its responsibility, by law, to provide shelter for UAC referred to its care by the Department of Homeland Security. To ensure sufficient capacity to provide shelter to UAC referred to HHS, ORR is requesting that BCFS HHS continue the use of up to 1008 hard-sided beds at Carrizo Springs, Texas.

DATES: Supplemental award funds will support activities until January 31, 2022.

FOR FURTHER INFORMATION CONTACT: Stephen Antkowiak, Office of Refugee Resettlement, Division of Unaccompanied Alien Children Operations, 330 Street SW, Washington, DC 20447. Phone: 202–260–6165. Email: *stephen.antkowiak@acf.hhs.gov.*

SUPPLEMENTARY INFORMATION: ORR is continuously monitoring its capacity to shelter the UAC referred to HHS, as well as the information received from interagency partners, to inform any future decisions or actions.

ORR has specific requirements for the provision of services. Award recipients must have the infrastructure, licensing, experience, and appropriate level of trained staff to meet those requirements. The expansion of the existing program and its services through this supplemental award is a key strategy for ORR to be prepared to meet its responsibility to provide shelter for UAC referred to its care by the Department of Homeland Security (DHS), and so the Customs and Border Protection can continue its vital national security mission to prevent illegal migration, trafficking, and protect the borders of the United States.

Statutory Authority: This program is authorized by—

(A) Section 462 of the Homeland Security Act of 2002, which in March 2003, transferred responsibility for the care and custody of UAC from the Commissioner of the former Immigration and Naturalization Service to the Director of ORR within HHS.

(B) The *Flores* Settlement Agreement, Case No. CV85–4544–RJK (C.D. Cal. 1996), as well as the William Wilberforce Trafficking Victims Protection Reauthorization Act of 2008 (Pub. L. 110–457), which authorizes post release services under certain conditions to eligible children. All programs must comply with the *Flores* Settlement Agreement, Case No. CV85–4544–RJK (C.D. Cal. 1996); pertinent regulations; and ORR policies and procedures.

Elizabeth Leo,

Senior Grants Policy Specialist, Office of Grants Policy, Office of Administration.

[FR Doc. 2021–14722 Filed 7–7–21; 4:15 pm]

BILLING CODE 4184–45–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Community Living

[OMB# 0985–New]

Agency Information Collection Activities; Proposed Collection; Comment Request; Evidence Based Program Fidelity Surveys

AGENCY: Administration for Community Living, HHS.

ACTION: Notice.

SUMMARY: The Administration for Community Living (ACL) is announcing an opportunity for the public to comment on the proposed collection of information listed above. Under the Paperwork Reduction Act of 1995 (the PRA), Federal agencies are required to publish a notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, and to allow 60 days for public comment in response to the notice. This notice solicits comments on the information collection requirements relating to a Grantee Survey and a Local Implementation Organization Survey that will be used by ACL to evaluate the fidelity with which ACL and its grantee organizations, under the Older Americans Act, implement the required evidence-based programs.

DATES: Comments on the collection of information must be submitted electronically by 11:59 p.m. (EST) or postmarked by September 10, 2021.

ADDRESSES: Submit electronic comments on the collection of information to: *Susan.Jenkins@acl.hhs.gov.* Submit written comments on the collection of information to

Administration for Community Living, Washington, DC 20201, Attention: Susan Jenkins.

FOR FURTHER INFORMATION CONTACT: Susan Jenkins, Administration for Community Living, Washington, DC 20201, 202–795–7369 or by email: *Susan.Jenkins@acl.hhs.gov.*

SUPPLEMENTARY INFORMATION: 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. ''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. The PRA requires Federal agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, ACL is publishing a notice of the proposed collection of information set forth in this document.

With respect to the following collection of information, ACL invites comments on our burden estimates or any other aspect of this collection of information, including:

(1) Whether the proposed collection of information is necessary for the proper performance of ACL's functions, including whether the information will have practical utility;

(2) the accuracy of ACL's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used to determine burden estimates; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and

(4) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques when appropriate, and other forms of information technology.

The Evidence Based Program Fidelity Surveys will be used by ACL to evaluate the fidelity with which ACL's grantee organizations, under the Older Americans Act, implement the required evidence-based programs. States that receive Older Americans Act funds under Title III–D are required to spend those funds on evidence-based programs to improve the health and well-being of their clients and to reduce disease and injury. Since 2003, the aging services network has been steadily moving towards wider implementation of disease prevention and health promotion programs that are based on scientific evidence and demonstrated to improve the health of older adults. The FY 2012 Congressional appropriations law included, for the first time, an evidence-based requirement related to Title III–D funds.

The results of this information collection will be used by ACL/AoA to:

- Effectively report its results to the President, to Congress, to the Department of Health and Human Services and to the public.
- Assess the effectiveness of ACL and its grantees in monitoring program fidelity.
- Aid in program refinement and continuous improvement.

To comment on this information collection please visit the ACL website: *https://www.acl.gov/about-acl/public-input.*

Estimated Program Burden

ACL estimates the burden associated with this collection of information as follows:

Respondent/data collection activity	Number of respondents	Responses per respondent	Hours per response	Annual burden hours
Grantee: Program selection process and survey	103	1	2.00	206
Local Implementation Organization Survey	412	1	0.67	275
Total	515	1	0.93	481

Dated: July 6, 2021.

Alison Barkoff,

Acting Administrator and Assistant Secretary for Aging.

[FR Doc. 2021–14700 Filed 7–9–21; 8:45 am]

BILLING CODE 4154–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2019–D–1997]

Food and Drug Administration Oversight of Food Products Covered by Systems Recognition Arrangements; Draft Guidance for Food and Drug Administration Staff; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of availability.

SUMMARY: The Food and Drug Administration (FDA or Agency) is announcing the availability of a draft guidance for FDA staff entitled ''FDA Oversight of Food Products Covered by Systems Recognition Arrangements.'' This draft guidance provides recommendations related to the FDA's regulatory oversight activities for food products imported from countries whose food safety systems the FDA has recognized in Systems Recognition Arrangements (SRAs).

DATES: Submit either electronic or written comments on the draft guidance by September 10, 2021 to ensure that the Agency considers your comment on the draft guidance before it begins work on the final version of the guidance.

ADDRESSES: You may submit comments on any guidance at any time as follows:

Electronic Submissions

Submit electronic comments in the following way:

- *Federal eRulemaking Portal: https://www.regulations.gov.* Follow the instructions for submitting comments. Comments submitted electronically, including attachments, to *https://www.regulations.gov* will be posted to the docket unchanged. Because your comment will be made public, you are solely responsible for ensuring that your comment does not include any confidential information that you or a third party may not wish to be posted, such as medical information, your or anyone else's Social Security number, or confidential business information, such as a manufacturing process. Please note that if you include your name, contact information, or other information that identifies you in the body of your comments, that information will be posted on *https://www.regulations.gov.*
- If you want to submit a comment with confidential information that you do not wish to be made available to the public, submit the comment as a