DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

[CFDA Number: 93.676]

Announcement of Intent To Issue an OPDIV-Initiated Supplement Under the Standing Announcement for Residential (Shelter) Services for Unaccompanied 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), U.S Department of Health and Human Services (HHS).

ACTION: Notice of intent to issue an OPDIV-Initiated Supplement.

SUMMARY: Administration for Children and Families, Office of Refugee Resettlement, announces the intent to issue an OPDIV-Initiated Supplement in multiple installments to BCFS Health and Human Services, San Antonio, TX. The aggregate total of the multiple installments will not exceed $367,860,381. The first two installments will be issued prior to September 30, 2018. The remaining installments will be issued after September 30, 2018 on to be determined dates. ORR has been identifying additional capacity to provide shelter for potential increases in apprehensions of Unaccompanied Children at the U.S. Southern 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 Unaccompanied Alien Children referred to its care by the Department of Homeland Security (DHS). To ensure sufficient capacity to provide shelter to unaccompanied children referred to HHS, BCFS proposed to provide ORR with 3,800 beds in an expedited manner.

DATES: Supplemental award funds will support activities through December 31, 2018.

FOR FURTHER INFORMATION CONTACT: Jallyn Sualog, Director, Division of Children's Services, Office of Refugee Resettlement, 330 C Street SW, Washington, DC 20447. Phone: 202–401–4997. Email: *DCSProgram@acf.hhs.gov.*

SUPPLEMENTARY INFORMATION: ORR is continuously monitoring its capacity to shelter the unaccompanied children 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 Unaccompanied Children referred to its care by DHS and so that the U.S. Border Patrol 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 Unaccompanied Alien Children from the Commissioner of the former Immigration and Naturalization Service (INS) to the Director of ORR of the Department of Health and Human Services (HHS).

(B) The Flores Settlement Agreement, Case No. CV85–4544RJK (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.

Karen Shields,

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

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity

AGENCY: Office of Planning, Research, and Evaluation; ACF; HHS.

ACTION: Request for public comment.

Title: Evaluation of Employment Coaching for TANF and Related Populations—Second Follow-Up Survey (OMB #0970–0506).

SUMMARY: The Administration for Children and Families (ACF) is proposing an additional data collection activity as part of the Evaluation of Employment Coaching for TANF and Related Populations. The Office of Management and Budget (OMB) Office of Information and Regulatory Affairs approved this information collection in March 2018 (0970–0506). ACF is proposing a second follow-up survey conducted as part of the evaluation.

This study will provide an opportunity to learn more about the potential of coaching to help clients achieve self-sufficiency and other desired employment-related outcomes. It will take place over five years in the following employment programs: MyGoals for Employment Success in Baltimore, MyGoals for Employment Success in Houston, Family Development and Self-Sufficiency program in Iowa, LIFT in New York City, Chicago, and Los Angeles; Work Success in Utah; and Goal4 It! in Jefferson County, Colorado. Together, these programs will include Temporary Assistance for Needy Families (TANF) agencies and other public or private employment programs that serve low-income individuals. Each site will have a robust coaching component and the capacity to conduct a rigorous impact evaluation. This study will provide information on whether coaching helps people obtain and retain jobs, advance in their careers, move toward self-sufficiency, and improve their overall well-being. To meet these objectives, this study includes an impact and implementation study, as approved by OMB.

This submission builds on the existing impact study, which randomly assigned participants to either a ''program group,'' who were paired with a coach, or to a ''control group,'' who were not paired with a coach. The effectiveness of the coaching will be determined by differences between members of the program and control groups in outcomes such as obtaining and retaining employment, earnings, measures of self-sufficiency, and measures of self-regulation.

The proposed information collection activity is a second follow-up survey, which will be available to participants approximately 21 months after random assignment. The second follow-up survey will provide rigorous evidence on whether the coaching interventions are effective, for whom, and under what circumstances.

Respondents: Individuals enrolled in the Evaluation of Employment Coaching for TANF and Related Populations. All participants will be able to opt out of participating in the data collection activities.

ANNUAL BURDEN ESTIMATES

Instrument	Total number of respondents	Annual number of respondents	Number of responses per respondent	Average burden hours per response	Annual burden hours
Second follow-up survey	4,800	1,600	1	1	1,600

Estimated Total Annual Burden Hours: 1,600.

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 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. Email address: *OPREinfocollection@acf.hhs.gov.* All requests should be identified by the title of the information collection.

SUPPLEMENTARY INFORMATION: 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: Section 413 of the Social Security Act, as amended by the FY 2017 Consolidated Appropriations Act, 2017 (Pub. L. 115–31).

Emily B. Jabbour,

ACF/OPRE Certifying Officer.

[FR Doc. 2018–20223 Filed 9–17–18; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2017–N–7022]

Post-Marketing Pediatric-Focused Product Safety Reviews; Establishment of a Public Docket; Request for Comments

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; establishment of a public docket; request for comments.

SUMMARY: The Food and Drug Administration (FDA) has established a public docket to collect comments related to the post-marketing, pediatric-focused safety reviews of products posted between April 2, 2018, and September 14, 2018, on FDA's website but not presented at the September 20, 2018, Pediatric Advisory Committee (PAC) meeting. These reviews are intended to be available for review and comment by members of the PAC, interested parties (such as academic researchers, regulated industries, consortia, and patient groups), and the general public.

DATES: Submit either electronic or written comments by September 28, 2018.

ADDRESSES: FDA has established a docket for public comment on this document. The docket number is FDA–2017–N–7022. The docket will close on September 28, 2018. Submit either electronic or written comments by that date. Please note that late, untimely filed comments will not be considered. Electronic comments must be submitted on or before September 28, 2018. The *https://www.regulations.gov* electronic filing system will accept comments until midnight Eastern Time at the end of September 28, 2018. Comments received by mail/hand delivery/courier (for written/paper submissions) will be considered timely if they are postmarked or the delivery service acceptance receipt is on or before that date.

You may submit comments 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 make available to the public, submit the comment as a written/paper submission and in the manner detailed (see ''Written/Paper Submissions'' and ''Instructions'').

Written/Paper Submissions

Submit written/paper submissions as follows:

- *Mail/Hand Delivery/Courier (for written/paper submissions):* Dockets Management Staff (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.
- For written/paper comments submitted to the Dockets Management Staff, FDA will post your comment, as well as any attachments, except for information submitted, marked and identified, as confidential, if submitted as detailed in ''Instructions.''

Instructions: All submissions received must include the Docket No. FDA–2017–N–7022 for ''Post-Marketing Pediatric-Focused Product Safety Reviews; Establishment of a Public Docket; Request for Comments.'' Received comments, those filed in a timely manner (see **ADDRESSES**), will be placed in the docket and, except for those submitted as ''Confidential Submissions,'' publicly viewable at *https://www.regulations.gov* or at the Dockets Management Staff between 9