amended by the Protecting Access to Medicare Act of 2014, Pub. L. 113–93.

Christopher Beach,

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

[FR Doc. 2015–09074 Filed 4–20–15; 8:45 am]

BILLING CODE 4184–43–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

[OMB No.: 0970–0365]

Submission for OMB Review; Comment Request

Proposed Projects:

Title: Performance Measures for Community-Centered Healthy Marriage, Pathways to Responsible Fatherhood and Community-Centered Responsible Fatherhood Ex-Prisoner Reentry grant programs.

Description: The Office of Family Assistance (OFA), Administration for Children and Families (ACF), U.S. Department of Health and Human Services (HHS), intends to request approval from the Office of Management and Budget (OMB) to extend OMB Form 0970–0365 for the collection of performance measures from grantees for the Community-Centered Healthy Marriage, Pathways to Responsible Fatherhood and Community-Centered Responsible Fatherhood Ex-Prisoner Reentry discretionary grant programs. ACF offered a one year extension to all grants in an effort to increase the consistency and stability in program implementation, particularly in view of grantee progress toward achieving program goals. The performance measure data obtained from the grantees will be used by OFA to continue reporting on the overall performance of these grant programs.

Data will be collected from all 60 Community-Centered Healthy Marriage, 54 Pathways to Responsible Fatherhood and 5 Community-Centered Responsible Fatherhood Ex-Prisoner Reentry grantees in the OFA programs. Grantees will report on program and participant outcomes in such areas as participants' improvement in knowledge skills, attitudes, and behaviors related to healthy marriage and responsible fatherhood. Grantees will be asked to input data for selected outcomes for activities funded under the grants. Grantees will extract data from program records and will report the data twice yearly through an on-line data collection tool. Training and assistance will be provided to grantees to support this data collection process.

Respondents: Office of Family Assistance Funded Community-Centered Healthy Marriage, Pathways to Responsible Fatherhood and Community-Centered Responsible Fatherhood Ex-Prisoner Reentry Grantees.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total annual burden hours
Performance measure reporting form (for private sector affected public)	110	2	0.8	176
Performance measure reporting form (for State, local, and tribal government affected public)	9	2	0.8	14

Estimated Total Annual Burden Hours: 190.

Additional Information: Copies of the proposed collection may be obtained by writing to the Administration for Children and Families, Office of Planning, Research and Evaluation, 370 L'Enfant Promenade SW., Washington, DC 20447, Attn: ACF Reports Clearance Officer. All requests should be identified by the title of the information collection. Email address: *infocollection@acf.hhs.gov.*

OMB Comment: OMB is required to make a decision concerning 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. Written comments and recommendations for the proposed information collection should be sent directly to the following: Office of Management and Budget, Paperwork Reduction Project, Fax: 202–395–7285, Email: *OIRA_SUBMISSION@OMB.EOP.GOV,* Attn: Desk Officer for the Administration for Children and Families.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 2015–09189 Filed 4–20–15; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Title: Office of Refugee Resettlement Individual Development Accounts (ORR–IDA) Program.

OMB No.: New Collection.

Description: Description: The Office of Refugee Resettlement seeks OMB approval to develop three data collection tools for use in the ORR IDA Program.

The ORR IDA Program represents an anti-poverty strategy built on asset accumulation for low-income refugee individuals and families with the goal of promoting refugee economic independence.

IDAs are leveraged or matched, savings accounts. In the ORR Refugee IDA program, IDAs are matched with federal funds that have been allocated as ''match funds'' from at least 65 percent of the annual federal grant award. IDAs are established in insured accounts in qualified financial institutions. The funds are intended for the Asset Goals specified in this announcement. Although the refugee participant maintains control of all funds that the participant deposits in the IDA, including all interest that may accrue on the funds, the participant must sign a Savings Plan Agreement which specifies that the funds in the account will be used only for the participant's qualified Asset Goal(s) or for an emergency withdrawal.

The objectives of this program are to:

1. Establish IDAs for eligible participants;

2. Encourage regular saving habits among refugees;

3. Promote their participation in the financial institutions of this country;

4. Promote refugee acquisition of assets to build individual, family, and community resources;

5. Increase refugee knowledge of financial and monetary topics including developing a household budget;

6. Assist refugees in advancing their education;

7. Increase home ownership among refugees; and

8. Assist refugees in gaining access to capital.

The tools will collect information from grantees that will help ORR determine whether they are meeting the objectives of the program. Data to be collected will only include specialized, and relevant information to the program such as, number of people enrolled, amount in dollar allocated for matching IDA savings, number and value of assets purchased, confirmation of refugee status, and types and quantity of training provided. Tools will be used for semi-annual reports as well as for monitoring to ensure progress towards success, and appropriate use of federal funds.

Respondents: Office of Refugee Resettlement Individual Development Accounts Program grantees.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Program Status Report	22	2	1	44
Community Impact Report	22	2	1	44
Demographic	22	2	1	44

Estimated Total Annual Burden Hours: 132 hours.

Additional Information: Copies of the proposed collection may be obtained by writing to the Administration for Children and Families, Office of Planning, Research and Evaluation, 370 L'Enfant Promenade SW., Washington, DC 20447, Attn: ACF Reports Clearance Officer. All requests should be identified by the title of the information collection. Email address: *infocollection@acf.hhs.gov.*

OMB Comment: OMB is required to make a decision concerning 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. Written comments and recommendations for the proposed information collection should be sent directly to the following: Office of Management and Budget, Paperwork Reduction Project, Email: *OIRA_SUBMISSION@OMB.EOP.GOV, Attn:* Desk Officer for the Administration for Children and Families.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 2015–09192 Filed 4–20–15; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2015–N–0001]

Request for Nominations on the Allergenic Products Advisory Committee

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is requesting that any industry organizations interested in participating in the selection of a nonvoting industry representative to serve on the Allergenic Products Advisory Committee for the Center for Biologics Evaluation and Research notify FDA in writing. FDA is also requesting nominations for a nonvoting industry representative to serve on the Allergenic Products Advisory Committee. A nominee may either be self-nominated or nominated by an organization to serve as a nonvoting industry representative. Nominations will be accepted for current or upcoming vacancies effective with this notice.

DATES: Any industry organization interested in participating in the selection of an appropriate nonvoting member to represent industry interests must send a letter stating that interest to the FDA by May 21, 2015, (see sections I and II for further details). Concurrently, nomination materials for prospective candidates should be sent to FDA by May 21, 2015.

ADDRESSES: All statements of interest from interested industry organizations interested in participating in the selection process of nonvoting industry representative nomination should be sent to Janie Kim (see **FOR FURTHER INFORMATION CONTACT**). All nominations for nonvoting industry representatives may be submitted electronically by accessing the FDA Advisory Committee Membership Nomination Portal: *https://www.accessdata.fda.gov/scripts/FACTRSPortal/FACTRS/index.cfm* or by mail to Advisory Committee Oversight and Management Staff, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 32, Rm. 5103, Silver Spring, MD 20993–0002. Information about becoming a member of an FDA advisory committee can also be obtained by visiting FDA's Web site at *http://www.fda.gov/AdvisoryCommittees/default.htm.*

FOR FURTHER INFORMATION CONTACT: Janie Kim, Center for Biologics Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Silver Spring, MD 20993, 301–796–9016, FAX: 301–595–1307, email: *janie.kim@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION: The Agency intends to add a nonvoting industry representative to the following advisory committee:

I. Allergenic Products Advisory Committee

The Committee reviews and evaluates available data concerning the safety, effectiveness, and adequacy of labeling of marketed and investigational allergenic biological products or materials that are administered to humans for the diagnosis, prevention, or treatment of allergies and allergic disease, and makes appropriate recommendations to the Commissioner of Food and Drugs of its findings regarding the affirmation or revocation of biological product licenses, on the safety, effectiveness, and labeling of the products, on clinical and laboratory studies of such products, on amendments or revisions to regulations governing the manufacture, testing and licensing of allergenic biological products, and on the quality and relevance of FDA's research programs which provide the scientific support for regulating these agents.