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

<table>
<thead>
<tr>
<th>Instrument</th>
<th>Number of respondents</th>
<th>Number of responses per respondent</th>
<th>Average burden hours per response</th>
<th>Total annual burden hours</th>
</tr>
</thead>
<tbody>
<tr>
<td>Performance measure reporting form (for private sector affected public)</td>
<td>110</td>
<td>2</td>
<td>0.8</td>
<td>176</td>
</tr>
<tr>
<td>Performance measure reporting form (for State, local, and tribal government affected public)</td>
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<td>2</td>
<td>0.8</td>
<td>14</td>
</tr>
</tbody>
</table>

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:
infolinecollection@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 Sargsis,
Reports Clearance Officer.
[FR Doc. 2015–09189 Filed 4–20–15; 8:45 am]
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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
cumulation 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.

<table>
<thead>
<tr>
<th>Instrument</th>
<th>Number of respondents</th>
<th>Number of responses per respondent</th>
<th>Average burden hours per response</th>
<th>Total burden hours</th>
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<tbody>
<tr>
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<td>2</td>
<td>1</td>
<td>44</td>
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<tr>
<td>Community Impact Report</td>
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<td>2</td>
<td>1</td>
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</tr>
<tr>
<td>Demographic</td>
<td>22</td>
<td>2</td>
<td>1</td>
<td>44</td>
</tr>
</tbody>
</table>

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 Sargsj, Reports Clearance Officer.

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.