

the grantees may be experiencing in implementing their projects on a timely

manner, and, for writing Annual Report to Congress.

Home-Based Child Care Microenterprise Development Program 23

Respondents: Refugee Microenterprise Development Program 22. Refugee

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondents	Average burden hours per respondents	Total burden hours
Refugee Microenterprise Development Program .....	22	8	4	88
Refugee Home-Based Child Care Microenterprise Development Program ...	23	7	4	92
Total Burden .....				180

*Estimated Total Annual Burden Hours: 180.*

In compliance with the requirements of Section 506(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. 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, 370 L'Enfant Promenade SW., Washington, DC 20447, Attn: ACF Reports Clearance Officer. Email address: [infocollection@acf.hhs.gov](mailto:infocollection@acf.hhs.gov). All requests should be identified by the title of the information collection.

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.

**Robert Sargis,**

*Reports Clearance Officer.*

[FR Doc. 2015-01628 Filed 1-28-15; 8:45 am]

**BILLING CODE 4184-01-P**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Administration for Children and Families**

**Proposed Information Collection Activity; Comment Request**

**Proposed Projects**

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

*OMB No.:* New Collection.

*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.

In compliance with the requirements of Section 506(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. 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, 370 L'Enfant Promenade SW., Washington, DC 20447, Attn: ACF Reports Clearance Officer. Email address: [infocollection@acf.hhs.gov](mailto:infocollection@acf.hhs.gov). All requests should be identified by the title of the information collection.

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.

**Robert Sargis,**

*Reports Clearance Officer.*

[FR Doc. 2015-01643 Filed 1-28-15; 8:45 am]

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## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

[Docket No. FDA-2012-N-0115]

#### Agency Information Collection Activities; Proposed Collection; Comment Request; Guidance for Industry and Food and Drug Administration Staff—Class II Special Controls Guidance Document: Automated Blood Cell Separator Device Operating by Centrifugal or Filtration Principle

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Notice.

**SUMMARY:** The Food and Drug Administration (FDA) is announcing an opportunity for public comment on the proposed collection of certain information by the Agency. Under the Paperwork Reduction Act of 1995 (the PRA), Federal Agencies are required to publish 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 collection of information concerning class II special controls for an automated blood cell separator device operating by centrifugal or filtration separation principle.

**DATES:** Submit either electronic or written comments on the collection of information by March 30, 2015.

**ADDRESSES:** Submit electronic comments on the collection of information to <http://www.regulations.gov>. Submit written comments on the collection of information to the Division of Dockets Management (HFA-305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852. All comments should be identified with the docket number found in brackets in the heading of this document.

**FOR FURTHER INFORMATION CONTACT:** FDA PRA Staff, Office of Operations, Food and Drug Administration, 8455

Colesville Rd., COLE-14526, Silver Springs, MD 20993-0002, [PRASStaff@fda.hhs.gov](mailto:PRASStaff@fda.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. Section 3506(c)(2)(A) of the PRA (44 U.S.C. 3506(c)(2)(A)) 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, FDA is publishing notice of the proposed collection of information set forth in this document.

With respect to the following collection of information, FDA invites comments on these topics: (1) Whether the proposed collection of information is necessary for the proper performance of FDA's functions, including whether the information will have practical utility; (2) the accuracy of FDA's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (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.

#### Guidance for Industry and FDA Staff—Class II Special Controls Guidance Document: Automated Blood Cell Separator Device Operating by Centrifugal or Filtration Separation Principle (OMB Control Number 0910-0594)—Extension

Under the Safe Medical Devices Act of 1990 (Pub. L. 101-629), FDA may