welfare cases; and (2) Training judges, attorneys, and other legal personnel in child welfare cases; and conducting cross-training with child welfare agency staff and contractors.

The statute requires separate applications for these two new grants. The annual burden estimates below describe the estimated burden for the two new grants. ACF collects information from the States about their work under these grants (applications, program reports) by way of a Program Instruction issued on June 15, 2006.

This Program Instruction describes the programmatic and fiscal provisions and reporting requirements for each of the grants, specifies the application submittal and approval procedures for the grants for fiscal years 2006 through 2010, and identifies technical resources for use by State courts during the course of the grants. The agency uses the information received to ensure compliance with the statute and provide training and technical assistance to the grantees.

Respondents: State Courts.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Application	52	2	40	4,160
Annual Program Report	52	2	36	3,744

Estimated Total Annual Burden Hours: 7,904.

Additional Information

Copies of the proposed collection may be obtained by writing to the Administration for Children and Families, Office of Administration, Office of Information Services, 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. E-mail 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: 202–395–6974, Attn: Desk Officer for the Administration for Children and Families.

Dated: January 18, 2008.

Janean Chambers,

Reports Clearance Officer.

[FR Doc. 08–324 Filed 1–25–08; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Awards to Eleven Office of Refugee Resettlement Shelter Care Providers

AGENCY: Office of Refugee Resettlement, Administration for Children and Families, HHS.

ACTION: Notice of Grant Awards.

CFDA #: 93.676

SUMMARY: This notice is hereby given that an award will be made to an unaccompanied alien shelter care provider, Southwest Regional Youth Village, Vincennes, Indiana, in the amount of $586,719. This funding will support services through September 30, 2008.

This funding will support the expansion of secure program bed capacity to meet the number of unaccompanied alien children referrals from the Department of Homeland Security (DHS).

The program is mandated by section 462 of the Homeland Security Act to ensure appropriate placement of all referrals from the DHS. ORR's ability to meet this mandate is often a challenge since the program is completely tied to DHS interior apprehension strategies and the sporadic number of border crossers.

The program has very specific requirements for the provision of services. This grantee is one of the only entities with the infrastructure, licensing, experience and appropriate level of trained staff to meet the service requirements for secure capacity. The program's ability to meet the number of secure referrals from DHS can only be accommodated through the expansion of this program through the supplemental award process.

FOR FURTHER INFORMATION CONTACT: Kenneth Tota, Office of Refugee Resettlement, Administration for Children and Families, 370 L'Enfant Promenade, SW., Washington, DC, 20447, telephone (202) 401–4858.

Dated: January 17, 2008.

Brent Orrell,

Acting Director, Office of Refugee Resettlement.

[FR Doc. E8–1360 Filed 1–25–08; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2008N–0019]

Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Survey of Food Safety and Nutrition Information Provided to Pregnant Women by Health Care Providers and the Special Supplemental Nutrition Program for Women, Infants, and Children Educators

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a collection of information entitled ''Survey of Food Safety and Nutrition Information Provided to Pregnant Women by Health Care Providers and WIC Educators'' has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

FOR FURTHER INFORMATION CONTACT: Jonna Capezzuto, Office of the Chief

Information Officer (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–4659.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of March 20, 2007 (72 FR 13117), the agency announced that the proposed information collection had been submitted to OMB for review and clearance under 44 U.S.C. 3507. An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. OMB has now approved the information collection and has assigned OMB control number 0910–0617. The approval expires on January 31, 2011. A copy of the supporting statement for this information collection is available on the Internet at *http://www.reginfo.gov/public/do/PRAMain*.

Dated: January 18, 2008.

Jeffrey Shuren,

Assistant Commissioner for Policy.

[FR Doc. E8–1353 Filed 1–25–08; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2008N–0018]

Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Food and Drug Administration Survey of Physicians' Perceptions of the Impact of Early Risk Communication About Medical Products

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a collection of information entitled ''FDA Survey of Physicians' Perceptions of the Impact of Early Risk Communication About Medical Products'' has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

FOR FURTHER INFORMATION CONTACT: Jonna Capezzuto, Office of the Chief Information Officer (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–4659.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of July 19, 2007 (72 FR 39628), the agency announced that the proposed information collection had been submitted to OMB for review and clearance under 44 U.S.C. 3507. An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. OMB has now approved the information collection and has assigned OMB control number 0910–0615. The approval expires on December 31, 2010. A copy of the supporting statement for this information collection is available on the Internet at *http://www.reginfo.gov/public/do/PRAMain*.

Dated: January 18, 2008.

Jeffrey Shuren,

Assistant Commissioner for Policy.

[FR Doc. E8–1355 Filed 1–25–08; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2008N–0017]

Agency Information Collection Activities; Proposed Collection; Comment Request; Exports: Notification and Recordkeeping Requirements

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 notification and recordkeeping requirements for persons exporting human drugs, biological products, devices, animal drugs, food, and cosmetics that may not be marketed or sold in the United States.

DATES: Submit written or electronic comments on the collection of information by March 28, 2008.

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: Jonna Capezzuto, Office of the Chief Information Officer (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–4659.

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.

Exports: Notification and Recordkeeping Requirements, 21 CFR Part 1 (OMB Control Number 0910–0482) — Extension

The respondents to this information collection are exporters who have notified FDA of their intent to export unapproved products that may not be sold or marketed in the United States as allowed under 801(e) of the Federal Food, Drug, and Cosmetic Act (the act) (21 U.S.C. 381). In general, the notification identifies the product being