Operations and Regulatory Affairs, Division of Regulations Development, Attention: Document Identifier/OMB Control Number _____, Room C4–26–05, 7500 Security Boulevard, Baltimore, Maryland 21244–1850.

Dated: May 8, 2012.

Martique Jones,

Director, Regulations Development Group, Division B Office of Strategic Operations and Regulatory Affairs.

[FR Doc. 2012–11441 Filed 5–10–12; 8:45 am]

BILLING CODE 4120–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[CMS–8050–N]

Medicare Program; Meeting of the Medicare Economic Index Technical Advisory Panel—May 21, 2012

Correction

In notice document 2012–10702 appearing on pages 26553–26554 in the issue of Friday, May 4, 2012 make the following corrections:

1. On page 26553, in the third column, in the last paragraph on the page ''III. Registration Instructions'', the third sentence should appear as set forth below:

''You may register online at *http://www.hcdi.com/mei/* or by phone by contacting Toya Via, HCD International, at (301) 552–8803, by 5:00 p.m. EDT, Monday, May 14, 2012.''

[FR Doc. C1–2012–10702 Filed 5–10–12; 8:45 am]

BILLING CODE 1501–01–D

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Title: Understanding Urban Indians' Interactions with ACF Programs and Services.

OMB No.: New Collection.

Description: As part of the ''Understanding Urban Indians' Interactions with ACF Programs and Services'' research study, telephone interviews will be conducted with up to 34 directors of Urban Indian Centers around the county, and site visits will be made to four urban areas with a high percentage of American Indians or Alaska Natives within the population: Phoenix, AZ; New York City, NY; Minneapolis, MN; and Anchorage, Alaska. Members of the research study team will utilize a telephone interview guide and field discussion guide to collect information from staff members at relevant programs and organizations (*e.g.,* American Indian Organizations, social service agencies serving urban Indians) in these areas.

The goal of this information collection is to assess the challenges and context for family self-sufficiency for urban Indians and their interaction with services and programs offered by ACF. The information gathered will help to generate recommendations and action items for ACF in seeking to better understand and meet the needs of the urban Indian population.

Respondents: Urban Indian Center directors, non-Native service providers, Native service providers, and AI/AN residents.

ANNUAL BURDEN ESTIMATES

Instrument	Annual number of respondents	Number of responses per respondent	Average burden hours per response	Total annual burden hours
Telephone interview guide for directors of Urban Indian Centers	34	1	1.25	42.5
Interview guide for Native service providers	24	1	1	24
Interview guide for non-Native service providers	24	1	1	24
Interview guide for AI/AN residents of CITY	64	1	1	64

Estimated Total Annual Burden Hours: 154.5.

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: OPRE Reports Clearance Officer. All requests should be identified by the title of the information collection. Email address: *OPREinfocollection@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.

Steven M. Hanmer,

OPRE Reports Clearance Officer.

[FR Doc. 2012–11273 Filed 5–10–12; 8:45 am]

BILLING CODE 4184–09–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Notice of the Award of a Single-Source Program Expansion Supplement to Pima County Community College District in Tucson, AZ

AGENCY: Office of Family Assistance, ACF, HHS.

ACTION: Award of a Single-Source Program Expansion Supplement to Pima County Community College District, a public/state controlled institution of higher education in Tucson, Arizona.

CFDA Number: 93.093.

Statutory Authority: Section 2008(a) of Title XX of the Social Security Act, as amended by Section 5507 of the Affordable Care Act (Pub. L. 111–148).

SUMMARY: This Administration for Children and Families (ACF), Office of

Family Assistance (OFA), Health Profession Opportunity Grants (HPOG) program announces the award of a single-source program expansion supplement to Pima County Community College District (PCC), a public/state controlled institution of higher education in Tucson, Arizona. Expansion supplement funds will support the acceleration of enrollment in Project Year Two, necessitated by a random assignment study entitled—Innovative Strategies for Increasing Self-Sufficiency (ISIS). This allows PCC to meet the sample size required by ISIS in the specific time period.

PCC was found to be an ideal fit as an ISIS evaluation site due to its unique program characteristics. The program has a clear articulated career pathway program, capacity to achieve a treatment sample of 500 or more over two project years, and a treatment sample that would be clearly distinct from the control sample because of the provision of intensive support and HPOG specific classes that are contextualized and compressed.

DATES: The project period for the award is September 30, 2011–September 29, 2012.

FOR FURTHER INFORMATION CONTACT: Stan Koutstaal, Program Manager, Office of Family Assistance, 370 L'Enfant Promenade SW., Washington, DC 20447. Telephone: 202–401–5457; Email: *stanley.koutstaal@acf.hhs.gov.*

Earl S. Johnson,

Director, Office of Family Assistance.

[FR Doc. 2012–11489 Filed 5–10–12; 8:45 am]

BILLING CODE 4184–48–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2012–N–0430]

Agency Information Collection Activities; Proposed Collection; Comment Request; Voluntary Submission of Food/Feed Facility Profile Information

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 and to allow 60 days for public comment in response to the notice. This notice solicits comments on the information collection provisions of FDA's program of voluntary submission of food facility profile information and new Form FDA 3797, which may be submitted electronically via the FDA Industry Systems Web site.

DATES: Submit either electronic or written comments on the collection of information by July 10, 2012.

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: Domini Bean, Office of Information Management, Food and Drug Administration, 1350 Piccard Dr., PI50–400T, Rockville, MD 20850, 301–796–5733, *domini.bean@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 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.

Voluntary Submission of Food/Feed Facility Profile Information (OMB Control Number 0910—New)

FDA has broad legal authority under the Federal Food, Drug, and Cosmetic Act (the FD&C Act) and the Public Health Service Act to protect the public health and the safety of the nation's food supply. In addition, under the Public Health Security and Bioterrorism Preparedness and Response Act of 2002 (Pub. L. 107–188) (the ''Bioterrorism Act'') FDA was further authorized to improve the ability of the United States to prevent, prepare for, and respond to bioterrorism and other public health emergencies. The Bioterrorism Act added section 415 of the FD&C Act (21 U.S.C. 350d), which requires domestic and foreign facilities that manufacture, process, pack, or hold food for human or animal consumption in the United States to register with FDA. FDA regulations at 21 CFR 1.230 through 1.235 set forth the procedures for registration of food (including animal food/feed) facilities. Information provided to FDA under these regulations helps us notify quickly the facilities that might be affected by a deliberate or accidental contamination of the food supply. Furthermore, the FDA Food Safety Modernization Act (Pub. L. 11–353) (FSMA) added section 421 of the FD&C Act (21 U.S.C. 350j), which directed FDA to allocate resources to inspect facilities according to the known safety risks of the facilities. We propose to collect additional food/feed facility profile information on a voluntary basis from firms that complete the FDA food facility registration process. Food facility profile information voluntarily provided to FDA will help us to determine whether a firm is high-risk or non-high-risk. We will use the profile information to assist us in determining the frequency at which we will inspect the firm. Facilities that voluntarily submit the food facility profile information would benefit from our advance preparation through interaction with better-informed investigators and potentially reduced inspection time. The need for this collection of information derives from our objective to obtain current, timely, and policy-relevant information to carry out our statutory functions. The FDA Commissioner is authorized to undertake this collection as specified in section 1003(d)(2) of the FD&C Act (21 U.S.C. 393(d)(2)).