

evaluation. The ISIS project will test a range of promising strategies to promote employment, self-sufficiency, and reduce dependence on cash welfare. The ISIS project will evaluate multiple employment-focused strategies that build on previous approaches and are adapted to the current Federal, State, and local policy environment. The major goals of the project include increasing the empirical knowledge about the effectiveness of a variety of programs for low-income families to sustain employment and advance to positions that enable self-sufficiency, as well as producing useful findings for both policymakers and program administrators.

This proposed information collection activity focuses on collecting baseline data elements. Two data collection instruments will be completed by all participants prior to random assignment, and a third will be an interview guide to collect information from program staff. The first is a short baseline information form (BIF) that will collect basic identification, demographic, and contact information. The form will include relatively standard items from prior evaluations and national surveys. The second instrument will be a self-administered questionnaire (SAQ), covering information related to the project goals. The third instrument, baseline

implementation data collection interviews, will be used to collect information from knowledgeable informants about the service context for each evaluation site using a baseline implementation guide. The purpose of such interviews is to document and assess the service environment in which the evaluation is implemented and the opportunities for control group members to access the same or similar services as the treatment group members.

Respondents: Individuals enrolled in ISIS demonstration interventions, control group members, ISIS program operators (BIF and SAQ) and State and local informants (interviews).

ANNUAL BURDEN ESTIMATES

Instrument	Annual number of respondents	Number of responses per respondent	Average burden hours per response	Total annual burden hours
Baseline Information Form	4,800	1	0.75	3,600
Self-Administered Questionnaire	4,800	1	0.75	3,600
Baseline Implementation Data Collection Interviews	30	1	1	30

Estimated Total Annual Burden Hours: 7,230

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. E-mail address: OREinfocollection@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: June 10, 2011.
Steven M. Hanmer,
OPRE Reports Clearance Officer.
 [FR Doc. 2011-14869 Filed 6-15-11; 8:45 am]
BILLING CODE 4184-09-M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Title: Implementation, Systems and Outcome Evaluation of the Tribal and Low-Income Health Profession Opportunity Grants (HPOG).

OMB No.: New Collection.

Description: The Administration for Children and Families (ACF) is proposing information collection activities as part of the Implementation, Systems and Outcome Evaluation of the Health Profession Opportunity Grants (HPOG). Through this information collection, ACE seeks to develop comprehensive management and performance reports on the HPOG

initiative and design a feasible and reliable evaluation design to produce accurate evidence of the effect of HPOG on individuals and health job training programs systems.

The goals of the HPOG evaluation are to establish a performance management reporting process for HPOG, and design an evaluation of HPOG. Both goals require collecting information from HPOG grantees on a regular basis. The information collection proposed is an Internet-based collection of information from HPOG grantees on (1) program participants: Baseline characteristics, program participation and patterns, and participant outputs and outcomes; and (2) program designs and operating characteristics. The performance management system would collect information from grantees on their programs and participants on a bi-annual basis.

Respondents: Participant data to be collected by program staff in the 32 grantee organizations (higher education institutions, workforce investment boards, private training institutions, and tribal entities).

ANNUAL BURDEN ESTIMATES

Instrument	Annual number of respondents	Number of responses per respondent	Average burden hours per response	Total annual burden hours
Semi-annual HPOG Program Performance Report	32	2	31.25	2,000

Estimated Total Annual Burden Hours: 2,000.

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. *E-mail 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, *Fax:*
202-395-6974, *Attn:* Desk Officer for
the Administration, for Children and
Families.

Dated: June 9, 2011.

Steven M. Hanmer,
OPRE Reports Clearance Officer.

[FR Doc. 2011-14811 Filed 6-15-11; 8:45 am]

BILLING CODE 4184-09-M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Indian Health Service

Epidemiology Program for American Indian/Alaska Native Tribes and Urban Indian Communities; Correction

AGENCY: Indian Health Service, HHS.

ACTION: Notice; correction.

SUMMARY: The Indian Health Service published a document in the **Federal Register** on June 8, 2011, concerning competitive cooperative agreement applications to establish Tribal Epidemiology Centers serving American Indian/Alaska Native Tribes and urban Indian communities. The document contained one incorrect date.

FOR FURTHER INFORMATION CONTACT: Betty Gould, 301-443-1116.

Correction

In the **Federal Register** of June 8, 2011, in FR Doc. 2011-14131, on page 33318, in the first column, last complete sentence in the paragraph "Letters of Intent," correct the sentence to read:

"The LoI must be submitted to the Division of Grants Management to the

attention of Andrew Diggs by June 17, 2011."

Dated: June 9, 2011.

Betty Z. Gould,
Federal Register Liaison Officer, Indian Health Service.

[FR Doc. 2011-14794 Filed 6-15-11; 8:45 am]

BILLING CODE 4165-16-M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Proposed Collection; Comment Request; NINR End-of-Life and Palliative Care Science Needs Assessment: Funding Source Questionnaire (Survey of Authors)

SUMMARY: In compliance with the requirement of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, for opportunity for public comment on proposed data collection projects, the National Institute of Nursing Research (NINR), the National Institutes of Health (NIH) will publish periodic summaries of proposed projects to be submitted to the Office of Management and Budget (OMB).

Proposed Collection: Title: NINR End-of-Life and Palliative Care Science Needs Assessment: Funding Source Questionnaire (Survey of Authors). **Type of Information Collection Request:** NEW. **Need and Use of Information Collection:** The NINR End-of-Life Science Palliative Care (EOL PC) Needs Assessment: Funding Source Questionnaire will obtain information on funding sources of EOL PC research published studies for which a funding source is not cited or the information is unclear. Target participants are authors of publically-available EOL PC research studies published between 1997-2010 for whom a funding source is unknown or unclear. The questionnaire inquires about the funding source of the published study, type of funding received, year of funding, and duration of funded study. This is a 7-item questionnaire that takes approximately 5 minutes to complete. Data collected is part of a needs assessment to address the breadth and depth of EOL PC scientific issues for use in stimulating research capacity in the field. **Frequency of Response:** One time. **Affected Public:** Individual authors of publically available EOL PC research publications who do not list a funding source or the source is unclear within their publication. **Type of Respondents:** EOL PC researchers. The annual reporting burden is as follows: **Estimated Number of Respondents:** 1840; **Estimated**

Number of Responses per Respondent: 1; **Average Burden Hours per Response:** .08; and **Estimated Total Annual Burden Hours Requested:** 147. There are no Capital Costs, Operating or Maintenance Costs to report.

Request for Comments: Written comments and/or suggestions from the public and affected agencies are invited on one or more of the following points: (1) Whether the proposed collection of information is necessary for the proper performance of the function of the agency, including whether the information will have practical utility; (2) The accuracy of the agency'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 those who are to respond, including the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology.

FOR FURTHER INFORMATION CONTACT: To request more information on the proposed project or to obtain a copy of the data collection plans and instruments, contact Dr. Amanda Greene, Science Evaluation Officer, Office of Science Policy and Public Liaison, NINR, Democracy One, 6701 Democracy Blvd., Suite 710, Bethesda, MD 20892, or call non-toll-free number 301-496-9601, or e-mail your request to amanda.greene@nih.gov.

Comments Due Date: Comments regarding this information collection are best assured of having their full effect if received within 60 days of the date of this publication.

Dated: June 6, 2011.

Amanda Greene,
Science Evaluation Officer, NINR, National Institutes of Health.

[FR Doc. 2011-14878 Filed 6-15-11; 8:45 am]

BILLING CODE 4140-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Office of the Director; Notice of Charter Renewal for the Advisory Committee to the Director

In accordance with Title 41 of the U.S. Code of Federal Regulations, Section 102-3.65(a), notice is hereby given that the Charter for the Advisory Committee to the Director, National Institutes of Health was renewed for an