

ways to enhance 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. Written comments should be received within 60 days of this notice.

### Proposed Project

Program Effectiveness Evaluation of Workplace Intervention for Intimate Partner Violence (IPV)—[OMB# 0920–0789] [expiration date 12/31/09]—Extension—National Center for Injury Prevention and Control (NCIPC), Centers for Disease Control and Prevention (CDC).

### Background and Brief Description

Intimate partner violence (IPV) affects a substantial number of Americans, and there has recently been increasing recognition of the impact it has on the workplace. In addition to direct impacts (batterers often stalk or even attack IPV victims at their place of work), IPV has indirect impacts on the workplace

environment through lost productivity due to medical leave, absenteeism, and fear and distraction on the part of victims and coworkers. The Centers for Disease Control and Prevention (CDC) contracted with RTI International (RTI) to evaluate an ongoing workplace IPV prevention program being implemented at a national corporation. The purpose of the proposed evaluation is to document in detail the workplace IPV prevention activities delivered by the company, to determine the impact of these activities on short-term and long-term outcomes, and to determine the cost-effectiveness of the program. All managers at the corporate office of the corporation have been screened to assess training experiences. More in-depth surveys were conducted with managers who had not completed the corporation's IPV training.

Approximately 200 managers have been surveyed at baseline, and 6 months later. Manager surveys focus on knowledge/awareness of IPV and company resources for IPV and number of referrals for IPV assistance. This extension is requested to cover the 12-month follow-up administration of this

survey. Due to unexpected delays at the evaluation site and an inability to field the 6-month follow up survey with managers when originally scheduled, the project will need to be continued an additional 3 months.

Employees (N = 400) of those managers who completed the baseline survey using an anonymous web-based survey at baseline have been surveyed. These employees will also be surveyed 12 months later (during the reinstatement period) to assess their self-evaluated productivity, absenteeism, and perceptions of manager behavior. Responses of managers (and their employees) who received the IPV training in the study period (*i.e.*, sometime between the baseline and 12 month surveys) with untrained managers will be compared. The study will provide CDC and employers information about the potential effectiveness and cost-effectiveness of workplace IPV intervention strategies.

There are no costs to respondents except their time to participate in the interview.

### ESTIMATED ANNUALIZED BURDEN HOURS

Respondents	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total burden (in hours)
Employee .....	400	1	30/60	200
Manager .....	200	2	30/60	200
Total .....				400

Dated: August 19, 2009.

**Maryam I. Daneshvar,**

*Acting Reports Clearance Officer, Centers for Disease Control and Prevention.*

[FR Doc. E9–20578 Filed 8–25–09; 8:45 am]

**BILLING CODE 4163–18–P**

### DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Administration for Children and Families

### Submission for OMB Review; Comment Request

*Title:* Grant Application Data Summary (GADS) Form.

**OMB No.:** 0970–0328.

**Description:** The Grant Application Data Summary (GADS) form collects information from applicants seeking grants from the Administration for Native Americans (ANA). Applicants complete the GADS form as part of their funding package. This standardized format allows ANA to evaluate applications for financial assistance and to determine the relative focus of the projects for which such assistance is requested. The data collected focuses on the specific ANA program area for which the applicant is applying. ANA awards annual grants in the following nine competitive areas: (1) Social & Economic Development Strategies (SEDS); (2)

Alaska SEDS; (3) Special Initiative: Family Preservation: Improving the Well-Being of Children Planning; (4) Special Initiative: Family Preservation: Improving the Well-Being of Children Implementation; (5) Native Language Preservation & Maintenance Assessment; (6) Native Language Preservation & Maintenance Planning; (7) Native Language Preservation & Maintenance Implementation; (8) Native Language Preservation & Maintenance Immersion; (9) Environmental Regulatory Enhancement.

**Respondents:** Federally Recognized Indian Tribes, Tribal Governments, Native American Non-profits, Tribal Colleges and Universities.

### ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Grant Application Data Summary (GADS) .....	500	1	0.50	250

*Estimated Total Annual Burden Hours:* 250

**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](mailto: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: 202-395-7245, Attn: Desk Officer for the Administration for Children and Families.

Dated: August 21, 2009.

**Robert Sargis,**

*Reports Clearance Officer.*

[FR Doc. E9-20547 Filed 8-25-09; 8:45 am]

**BILLING CODE 4184-01-P**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Administration for Children and Families**

**Submission for OMB Review; Comment Request**

*Title:* Summary Data Component, National Child Abuse and Neglect Data System (NCANDS).

*OMB No.:* 0980-0229.

*Description:* The Child Abuse and Neglect Treatment Act (42 U.S.C. 5101 et seq.) as amended requires States to annually work with the Secretary to

provide to the maximum extent practical, a report that includes 12 data items listed in the statute. The National Child Abuse and Neglect Data System (NCANDS), administered by the Children's Bureau, meets this reporting requirement. In addition, the amendments of 1988 require that the data system shall be universal and case specific and integrated with other case-based foster care and adoption data collected by the Secretary. There are two data components, the Detailed Case Data Component (DCDC), which includes the case-level data submitted through the Child File and some aggregated data submitted through the Agency File, and the Summary Data component (SC), which is used by States that cannot submit case-level data. No changes are being requested. The Summary Data Component will be phased out over the next few years as the number of States that can complete the Child File increases.

*Respondents:* State Child Welfare Agencies.

**ANNUAL BURDEN ESTIMATES**

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Child File .....	50	1	80	4,000
Agency File .....	50	1	24	1,200
Summary Data Component (SDC) .....	2	1	32	64

*Estimated Total Annual Burden Hours:* 5,264.

**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](mailto: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-

395-7245. Attn: Desk Officer for the Administration for Children and Families.

Dated: August 21, 2009.

**Robert Sargis,**

*Reports Clearance Officer.*

[FR Doc. E9-20546 Filed 8-25-09; 8:45 am]

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**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Food and Drug Administration**

[Docket No. FDA-2009-N-0383]

**Request for Notification From Industry Organizations Interested in Participating in the Selection Process for a Nonvoting Industry Representative on the Tobacco Products Scientific Advisory Committee and Request for Nominations for a Nonvoting Industry Representative on the Tobacco Products Scientific Advisory Committee**

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

**ACTION:** Notice.

**SUMMARY:** The Food and Drug Administration (FDA) is requesting that any industry organizations interested in participating in the selection of a nonvoting industry representative to serve on its Tobacco Products Scientific Advisory Committee notify FDA in writing. A nominee may either be self-nominated or nominated by an organization to serve as a nonvoting industry representative. Nominations will be accepted for current vacancies effective with this notice. Elsewhere in this issue of the **Federal Register**, FDA is publishing two separate documents announcing the establishment of the committee and the request for nomination of the Tobacco Products Scientific Advisory Committee.

**DATES:** Any industry organization interested in participating in the selection of an appropriate nonvoting member to represent industry interests must send a letter stating the interest to FDA by September 25, 2009, for vacancies listed in the notice. Concurrently, nomination material for