DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request

Title: DHHS/ACF/OPRE Head Start Classroom-based Approaches and Resources for Emotion and Social skill promotion (CARES) project: Impact and Implementation Studies—Extension.

OMB No.: 0970–0364.

Description: The Head Start Classroom-based Approaches and Resources for Emotion and Social skill promotion (CARES) project is evaluating social emotional program enhancements within Head Start settings serving threeand four-year old children. This project focuses on identifying the central features of effective programs to provide the information federal policy makers and Head Start providers will need if they are to increase Head Start's capacity to improve the social and emotional skills and school readiness of preschool age children. The project is sponsored by the Office of Planning, Research, and Evaluation (OPRE) of the Administration for Children and Families (ACF). The Head Start CARES project uses a group-based randomized design to test the effects of three different evidence-based programs designed to improve the social and emotional development of children in Head Start classrooms.

Data to assess impacts of the program models in preschool was collected through surveys with teachers and parents, as well as direct child assessments. Data to assess implementation of the program models in preschool was collected through surveys and interviews with teachers, local coaches, trainers and center staff.

ANNUAL BURDEN ESTIMATES—EXTENSION

Data collection for both the impact and implementation studies occurred during the Head Start Year. The study sample involved 17 Head Start grantees/ delegate agencies, 104 centers, 307 classrooms, 1,042 selected 3-year old children and 2,885 selected 4-year old children.

The purpose of this request is to obtain an extension to finish impact data collection in the 2012 Follow-up Year (*e.g.*, Kindergarten for the 4-year olds). This data to assess impacts of the program models in the kindergarten year will be collected through teacher reports (surveys) and parent surveys.

Respondents: The respondents for the activities under the extension request for Follow-Up year data collection will be parents of children and kindergarten teachers of children in the study.

The annual burden estimates for both surveys covered by the extension are detailed below.

Instrument	Annual	Number of	Average	Estimated
	number of	responses per	burden hours	annual burden
	respondents	respondent	per response	hours
Teacher Report on Individual Children	962	1	0.33	317.5
Follow-up Parent Survey	962		0.33	317.5

Estimated Total Annual Burden Hours: 635.0.

In compliance with the requirements of Section 3506(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:

OPREinfocollection@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.

Dated: February 1, 2012.

Steven M. Hanmer,

OPRE Reports Clearance Officer. [FR Doc. 2012–2738 Filed 2–7–12; 8:45 am] **BILLING CODE 4184–22–P**

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request

Proposed Projects

Title: Tribal TANF Data Report, TANF Annual Report, and Reasonable Cause/ Corrective Action Documentation Process- Final.

OMB No.: 0970-0215.

Description

42 U.S.C. 612 (Section 412 of the Social Security Act as amended by Pub.

L. 104–193, the Personal Responsibility and Work Opportunity Reconciliation Act of 1996 (PRWORA), mandates that federally recognized Indian Tribes with an approved Tribal TANF program collect and submit to the Secretary of the Department of Health and Human Services data on the recipients served by the Tribes' programs. This information includes both aggregated and disaggregated data on case characteristics and individual characteristics. In addition, Tribes that are subject to a penalty are allowed to provide reasonable cause justifications as to why a penalty should not be imposed or may develop and implement corrective compliance procedures to eliminate the source of the penalty. Finally, there is an annual report, which requires the Tribes to describe program characteristics. All of the above requirements are currently approved by OMB and the Administration for Children and Families is simply proposing to extend them without any changes.

Respondents

Indian Tribes.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Final Tribal TANF Data Report Tribal TANF Annual Report Tribal TANF Reasonable Cause/Corrective	66 66 66	4 1 1	451 40 60	119,064 2,640 3,960
Estimated Total Annual Burden Hours				125,664

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. 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 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. 2012–2882 Filed 2–7–12; 8:45 am] BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Notice of Change in Application Requirements

AGENCY: Administration on Developmental Disabilities, ACF, HHS.

ACTION: Notification of change in allocation notification procedures to State Protection and Advocacy Systems (P&As) for mandatory awards under the Help America Vote Act (HAVA), Public Law 107–252.

CFDA Number: 93.617.

Statutory Authority: Title II, Subtitle D, Part 5, of HAVA 42 U.S.C. 15461–62; Section 102 of the Developmental Disabilities Assistance and Bill of Rights Act of 2000 (DD Act) (42 U.S.C. 15002); and Section 509 of the Rehabilitation Act of 1973 as amended (29 U.S.C. 794e)

SUMMARY: The Administration for Children and Families (ACF), Administration on Developmental Disabilities (ADD) has modified the application requirements for awards made to P&As under HAVA, Public Law 107–252. Under the program, formula grants are allotted to States based on population, financial need, and need for service. P&As provide services to individuals with developmental disabilities based on the identification of goals in the areas of emphasis listed in the DD Act and based on public input.

Section 291 of HAVA does not outline specific application requirements for P&As. Therefore, ADD has the discretion to alter the process by which P&As are notified of their annual allocations. Accordingly, P&As will no longer be required to submit an application; and, an annual Funding Opportunity Announcement (FOA) will no longer be published. Instead, ADD will now rely solely on the official notification provided to P&As by ACF's Division of Mandatory Grants. This notice informs P&As of the availability of their annual award allocations.

FOR FURTHER INFORMATION CONTACT:

Melvenia Wright, Program Specialist. Telephone: (202) 690–5557. Email: *Melvenia.Wright@acf.hhs.gov.*

Dated: February 2, 2012. Sharon Lewis,

Commissioner, Administration on Developmental Disabilities. [FR Doc. 2012–2920 Filed 2–7–12; 8:45 am] BILLING CODE 4184–38–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA-2011-N-0827]

Agency Information Collection Activities; Proposed Collection; Comment Request; Revisions to Labeling Requirements for Blood and Blood Components, Including Source Plasma; Correction

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; correction and extension of comment period.

SUMMARY: The Food and Drug Administration (FDA) is correcting a notice that appeared in the **Federal** Register of December 30, 2011. In the Federal Register of December 30, 2011, FDA published a notice entitled "Agency Information Collection Activities; Proposed Collection; Comment Request; Revisions to Labeling Requirements for Blood and **Blood Components**, Including Source Plasma," which provided incorrect publication information regarding the availability of the final rule. This document corrects this error and extends the comment period. Elsewhere in this issue of the Federal Register, FDA is publishing a companion final rule correction notice.

FOR FURTHER INFORMATION CONTACT: Joyce Strong, Office of Policy, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 32, Rm. 3208, Silver Spring, MD 20993–0002, 301– 796–9148.