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	66	4	451	119,064
Tribal TANF Annual Report	66	1	40	2,640
Tribal TANF Reasonable Cause/Corrective	66	1	60	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]

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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.