ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
AFCARS	52	2	2,581	268,424
Estimated Total Annual Burden Hours				268,424

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 Administration, Office of Information Services, 370 L'Enfant Promenade, SW., Washington, DC 20447, Attn: ACF Reports Clearance Officer. E-mail 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 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: October 29, 2010.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 2010–27836 Filed 11–3–10; 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: Developmental Disabilities Program Independent Evaluation Project.

OMB No.: 0970–0372.

Description: The National Independent Study of the State Developmental Disabilities Programs (National Study) is an independent (non-biased) study to examine through rigorous and comprehensive research procedures the three programs funded under the Developmental Disabilities Assistance and Bill of Rights Act of 2000 (DD Act): (1) State Councils on Developmental Disabilities (SCDDs); (2) State Protection and Advocacy Systems for Individuals with developmental disabilities (P&As); and (3) University Centers for Excellence in Developmental Disabilities (UCEDDs). The purpose of the study is to assess program effectiveness and achievements, including collaborative efforts among these State developmental disabilities (DD) network programs. A component of the study will be an examination of the Administration on Developmental Disabilities' efficiency and effectiveness to support these DD Network programs. The results of this evaluation will provide a report to the Administration on Developmental Disabilities (ADD) (the agency that administers these programs) with information on the effectiveness of its programs and policies and serve as a way for ADD to promote accountability to the public.

The independent study is a response to accountability requirements for ADD as identified in the Developmental Disabilities Assistance and Bill of Rights Act of 2000 (DD Act), the Government Performance and Results Act (GPRA) of 1993, and the Program Assessment Rating Tool (PART), previously administered by the Office of Management and Budget (OMB).

ADD has OMB approval for all the evaluation tools (*e.g.,* data collection instruments) for this study, except a new one being proposed. The new evaluation tool would be an on-line survey tool designed to collect data for an assessment of ADD.

Respondents: For the ADD assessment survey being added, the respondents would be Staff of State Councils on Developmental Disabilities, State Protection and Advocacy Systems for Individuals with developmental disabilities, and University Centers for Excellence in Developmental Disabilities, Education, Research, and Service

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
ADD Assessment Survey	60	1	1	60
DD Council Estimate of Total Burden Hours for Activities to Support Administration of Proposed Information Collection Instruments	20	1	33.50	670
P&A Estimate of Total Burden Hours for Activities to Support Administration of Proposed Information Collection Instruments	20	1	33.50	670
UCEDD Estimate of Total Burden Hours for Activities to Support Administration of Proposed Information Collection Instruments	20	1	33.50	670
DD Council: Executive Director Interview	20	1	4	80
DD Council: Interview with Council Chair/Council Members	60	1	0.75	45
DD Council: Group Interview with Policymakers, Collaborators, and Grantees	160	1	2	320
UCEDD: Telephone Interview with Current and Graduated Students	100	1	0.75	75
UCEDD: Interview with the Consumer Advisory Committee	60	1	0.75	45

ANNUAL BURDEN ESTIMATES—Continued

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
UCEDD: Interview with Peer Researchers and Colleagues	100	1	0.75	75
UCEDD: Interview with Recipients of Community Services or Members of Organizations/Agencies that are Trained to Provide Community Services	100	1	0.75	75
UCEDD: Self-administered Form	20	1	8	160
P&A: Executive Director Interview	20	1	4	80
P&A: Staff Interview	60	1	0.75	45
P&A: Board of Directors (Commissioners)-Chair and Members	60	1	0.75	45
P&A: Group Interview with Policymakers and Collaborators	160	1	2	320
P&A: Interview with Recipient of Community Education	100	1	0.75	75
P&A: Interview with Clients	100	1	0.75	75
P&A: Self-administered Form	20	1	8	160
UCEDD: Interview with Director	20	1	4	80
DD Council: Group Interview with Recipients of Self-Advocacy and Leadership Education and Training	100	1	0.75	75
DD Council: Group Interview with Recipients of Education and Training to Improve Community Capacity	100	1	0.75	75
DD Council: Self-administered Form	20	1	8	160
Estimated Total Annual Burden Hours:				4,135

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.*

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:* 202–395–7285, *E-mail: OIRA_SUBMISSION @OMB.EOP.GOV., Attn:* Desk Officer for the Administration for Children and Families.

Dated: November 1, 2010.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 2010–27855 Filed 11–3–10; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2010–N–0564]

Agency Information Collection Activities; Proposed Collection; Comment Request; Restaurant Menu Labeling: Registration for Small Chains Under Section 4205 of the Patient Protection and Affordable Care Act of 2010

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, including each proposed extension of an existing 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 registration under the Patient Protection and Affordable Care Act of 2010 (Affordable Care Act).

DATES: Submit either electronic or written comments on the collection of information by January 3, 2011.

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: Denver Presley, Jr., Office of Information Management, Food and Drug Administration, 1350 Piccard Dr., PI50–400B, Rockville, MD 20850, 301–796–3793.

SUPPLEMENTARY INFORMATION:

I. Background

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, including each proposed extension of an existing 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