5. LIHEAP Program Integrity Assessment Supplement—OMB Control No. 0970–0075; and
6. LIHEAP Model Plan (Detailed and Abbreviated)—OMB Control No. 0970–0075.

The content and annual burden estimates for the above existing data collections will remain unchanged. The only modification is the instrument of the data collections, which will now be through OLDC.

The information is being collected for the Department's annual LIHEAP Report to Congress. The data also provides information about the need for LIHEAP funds. Finally, the data are used in the calculation of LIHEAP performance measures under the Government Performance and Results Act of 1993. The data elements will improve the accuracy of measuring LIHEAP targeting performance and LIHEAP cost efficiency.

Respondents: State Governments and the District of Columbia

ANNUAL BURDEN ESTIMATES FOR PERFORMANCE MEASURES

Performance measure	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Percentage of Reduction in Household Energy Burden	51	1	60	3,060
Number of Utility Service Restorations	51	1	20	1,020
Number of Crises Averted	51	1	20	1,020

Estimated Total Annual Burden Hours: 5,100.

As LIHEAP is a block grant, there is a wide spectrum of capacity to collect and report data among grantees. The estimated burden hours displayed above are for the average LIHEAP grantee, assuming data collection systems and agreements already in place. For those grantees that would need to establish such agreements and systems, estimated burden for the initial year of reporting would more closely resemble 400 hours for each performance measure. However, after the systems are in place, estimated burden for the collection of these data will more closely reflect the figures in the table above.

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 Administration, Office of Information Services, 370 L'Enfant Prmenade 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 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. 2013–13384 Filed 6–5–13; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Community Living

Agency Information Collection Activities; Proposed Collection; Comment Request; University Centers for Excellence in Developmental Disabilities Education, Research, and Service—Annual Report

AGENCY: The Administration on Intellectual and Developmental Disabilities (AIDD), Administration for Community Living (ACL), HHS.

ACTION: Notice.

SUMMARY: The Administration on Intellectual and Developmental Disabilities (AIDD), now part of the Administration for Community Living, is announcing that the proposed collection of information listed below has been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995.

DATES: Submit written comments on the collection of information by July 8, 2013.

ADDRESSES: *OIRA_submission@omb.eop.gov* or by fax to 202.395.5806. Attn: OMB Desk Officer for ACL, Office of Information and Regulatory Affairs, OMB.

FOR FURTHER INFORMATION CONTACT: Jennifer Johnson, at 202–690–5982 or *jennifer.johnson@acl.hhs.gov.*

SUPPLEMENTARY INFORMATION: In compliance with 44 U.S.C. 3507, the Administration on Intellectual and Developmental Disabilities (now part of the Administration for Community Living) has submitted the following proposed collection of information to OMB for review and clearance.

Section 104 (42 U.S.C. 15004) of the Developmental Disabilities Assistance and Bill of Rights Act of 2000 (DD Act of 2000) directs the Secretary of Health and Human Services to develop and implement a system of program accountability to monitor the grantees funded under the DD Act of 2000. The program accountability system shall include the National Network of University Centers for Excellence in Developmental Disabilities Education, Research, and Service (UCEDDs) authorized under Part D of the DD Act of 2000. In addition to the accountability system, Section 154 (e) (42 U.S.C. 15064) of the DD Act of 2000 includes requirements for a UCEDD Annual Report. In response to the 60-day **Federal Register** notice related to this proposed data collection and published on January 15, 2013 in Volume 78, ten sets of comments were received. Most of the comments provided recommendations for enhancing the quality and clarity of the information to be collected. The comments resulted in some revisions to the proposed data collection tools. The originally proposed data collection tools, the comments with responses and a revised set of data collection tools may be obtained by contacting Jennifer Johnson at *jennifer.johnson@acl.hhs.gov* or 202–690–5982. AIDD estimates the

burden of this collection of information as 1,412 average burden hours per responses, for 67 UCEDDs—Total burden is 94,604 hours per year.

Dated: June 3, 2013.

Kathy Greenlee,

Administrator and Assistant Secretary for Aging.

[FR Doc. 2013–13421 Filed 6–5–13; 8:45 am]

BILLING CODE 4154–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2013–N–0190]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Requirements Under the Comprehensive Smokeless Tobacco Health Education Act of 1986, as Amended by the Family Smoking Prevention and Tobacco Control Act

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a proposed collection of information has been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995.

DATES: Fax written comments on the collection of information by July 8, 2013.

ADDRESSES: To ensure that comments on the information collection are received, OMB recommends that written comments be faxed to the Office of Information and Regulatory Affairs, OMB, Attn: FDA Desk Officer, FAX: 202–395–7285, or emailed to *oira_submission@omb.eop.gov*. All comments should be identified with the OMB control number 0910–0671. Also include the FDA docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Daniel Gittleson, Office of Information Management, Food and Drug Administration, 1350 Piccard Dr., PI50–400B, Rockville, MD 20850, 301–796–5156, *daniel.gittleson@fda.hhs.gov*.

SUPPLEMENTARY INFORMATION: In compliance with 44 U.S.C. 3507, FDA has submitted the following proposed collection of information to OMB for review and clearance.

Requirements Under the Comprehensive Smokeless Tobacco Health Education Act of 1986, as Amended by the Family Smoking Prevention and Tobacco Control Act—(OMB Control Number 0910–0671)—Extension

On June 22, 2009, the President signed the Family Smoking Prevention and Tobacco Control Act (the Tobacco Control Act) (Pub. L. 111–31) into law. Section 3 of the Comprehensive Smokeless Tobacco Health Education Act of 1986 (the Smokeless Tobacco Act) (15 U.S.C. 4402), as amended by section 204 of the Tobacco Control Act, requires, among other things, that all smokeless tobacco product packages and advertisements bear one of four required warning statements. Section 3(b)(3)(A) of the Smokeless Tobacco Act requires that the warnings be displayed on packaging and advertising for each brand of smokeless tobacco ''in accordance with a plan submitted by the tobacco product manufacturer, importer, distributor, or retailer'' to, and approved by, FDA.

This information collection—the submission to FDA of warning plans for smokeless tobacco products—is statutorily mandated. The warning plans will be reviewed by FDA, as required by the Smokeless Tobacco Act, to determine whether the companies' plans for the equal distribution and display of warning statements on packaging and the quarterly rotation of warning statements in advertising for each brand of smokeless tobacco products comply with section 3 of the Smokeless Tobacco Act, as amended.

Based on the Federal Trade Commission's (FTC's) previous experience with the submission of warning plans and FDA's experience with smokeless tobacco companies (e.g., correspondence associated with user fees under section 919 of the Federal Food, Drug, and Cosmetic Act, as amended by the Tobacco Control Act (21 U.S.C. 387s)), FDA estimates that there are 36 companies affected by this information collection. To account for the entry of new smokeless tobacco companies that may be affected by this information collection, FDA is estimating the total number of respondents to be 100.

When the FTC requested an extension of their approved information collection in 2007, based on over 20 years implementing the warning plan requirements and taking into account increased computerization and improvements in electronic communication, the FTC estimated submitting an initial plan would take 60 hours. Based on FDA's experience over the past several years, FDA believes the estimate of 60 hours to complete an initial rotational plan continues to be reasonable.

In the **Federal Register** of March 18, 2013 (78 FR 16678), FDA published a 60-day notice requesting public comment on the proposed collection of information. No comments were received.

FDA estimates the burden of this collection of information as follows:

TABLE 1—ESTIMATED ANNUAL REPORTING BURDEN [1]

Activity	Number of respondents	Number of responses per respondent	Total annual responses	Average burden per response	Total hours	Total capital costs
Submission of rotational plans for health warning label statements	100	1	100	60	6,000	$1,200

[1] There are no operating and maintenance costs associated with this collection of information.

FDA estimates a total of 100 respondents at 1 response each and 60 burden hours per response for a total of 6,000 burden hours (100 respondents × 1 response × 60 burden hours = 6,000 total burden hours). In addition, capital costs are based on all 100 respondents mailing in their submission at a postage rate of $12 for a 5-pound parcel (business parcel post mail delivered from the farthest delivery zone). Therefore, FDA estimates that the total postage cost for mailing the rotational warning plans to be $1,200.

Dated: June 3, 2013.

Leslie Kux,

Assistant Commissioner for Policy.

[FR Doc. 2013–13448 Filed 6–5–13; 8:45 am]

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