

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: May 31, 2001.

Bob Sargis,

Reports Clearance Officer.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request; Proposed Projects

Title: Low Income Home Energy Assistance Program (LIHEAP) Carryover and Reallotment Report.

OMB No.: 0970-0106.

Description: The LIHEAP statute and regulations require LIHEAP grantees to report certain information to HHS concerning funds forwarded and funds subject to reallotment. The 1994 reauthorization of the LIHEAP statute, the Human Service Amendments of 1994 (Pub. L. 103-252), requires that the carryover and reallotment report for one fiscal year be submitted to HHS by the

grantee before the Allotment for the next fiscal year may be awarded.

We are requesting changes in the collection of data by adding a form, the Carryover and Reallotment Report For FY 20___, for the collection of data previously requested by the Simplified Instructions for Timely Obligations of FY 20___ LIHEAP Funds and Reporting Funds for Carryover and Reallotment. The addition of the form will clarify the information being requested and ensure the submission of all the required information. Use of the form will be voluntary. It is being added in response to numerous queries each year concerning how to provide information. It will not add any additional burden on grantees. Grantees would have the option to use another format.

Respondents: State, Local or Tribal Govt.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Carryover and Reallotment	177	1	3	531
Estimated Total Annual Burden Hours:				531

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 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.

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: May 31, 2001.

Bob Sargis,

Reports Clearance Officer.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Blood Products Advisory Committee; Notice of Meeting

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

This notice announces a forthcoming meeting of a public advisory committee of the Food and Drug Administration (FDA). At least one portion of the meeting will be closed to the public.

Name of Committee: Blood Products Advisory Committee.

General Function of the Committee: To provide advice and recommendations to the agency on FDA's regulatory issues.

Date and Time: The meeting will be held on June 14, 2001, 8:30 a.m. to 5:30 p.m. and June 15, 2001, 8:30 a.m. to 3:45 p.m.

Location: Gaithersburg Hilton, 620 Perry Pkwy., Gaithersburg, MD 20877.

Contact: Linda A. Smallwood, Center for Biologics Evaluation and Research (HFM-302), Food and Drug Administration, 1401 Rockville Pike, Rockville, MD 20852-1448, 301-827-3514, or FDA Advisory Committee Information Line, 1-800-741-8138 (301-443-0572 in the Washington, DC area), code 19516. Please call the Information Line for up-to-date information on this meeting.

Agenda: On June 14, 2001, the following committee updates are tentatively scheduled: (1) Summary of the Public Health Service Advisory Committee on Blood Safety and Availability meeting, and (2) current thinking on clinical trial design and performance standards for approval of rapid human immunodeficiency virus (HIV) tests. In the morning, the committee will hear presentations, and discuss and make recommendations on re-entry for donors deferred because of HIV or hepatitis C virus (HCV) nucleic acid testing (NAT) or serological test results. In the afternoon, the committee will hear presentations, and discuss and make recommendations on the Clinical Laboratory Improvement Act (CLIA) criteria for invitro diagnostic tests: (1) Applicability of waivers to HIV rapid tests, and (2) revision of the uniform