

HHS will publish a notice in the **Federal Register** reporting the addition of this class to the SEC or the result of any provision by Congress regarding the decision by HHS to add the class to the SEC.

FOR FURTHER INFORMATION CONTACT:

Larry Elliott, Director, Office of Compensation Analysis and Support, National Institute for Occupational Safety and Health, 4676 Columbia Parkway, MS C-46, Cincinnati, OH 45226, Telephone 513-533-6800 (this is not a toll-free number). Information requests can also be submitted by e-mail to OCAS@CDC.GOV.

Dated: September 20, 2005.

John Howard,

Director, National Institute for Occupational Safety and Health, Centers for Disease Control and Prevention.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request

Proposed Projects

Title: ACF-196 State Temporary Assistance for Needy Families Financial Report.

OMB No.: 0970-0247.

Description: This information collection is authorized under the Personal Responsibility and Work Opportunity Reconciliation Act of 1996 (PRWORA). The request is for renewal of approval to use the Administration for Children and Families (ACF) 196 form for periodic financial reporting under the Temporary Assistance for Needy Families (TANF) program.

Current approval expires on September 30, 2005.

States participating in the TANF program are required by statute to report financial data on a quarterly basis. This form meets the legal standard and provides essential data on the use of Federal funds. Failure to collect the data would seriously compromise ACF's ability to monitor program expenditures, estimate funding needs and prepare budget submissions required by Congress. Financial reporting under the TANF program is governed by 45 CFR Part 265.

Respondents: State TANF Agencies.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
ACF-196	54	4	8	1,728

Estimated Total Annual Burden Hours: 1,728.

Additional Information: 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, 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: September 20, 2005.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 05-19271 Filed 9-26-05; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

International Conference on Harmonisation Workshop on Oncolytic Viruses; Public Workshop

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of public workshop.

SUMMARY: The Food and Drug Administration (FDA) is announcing a public workshop entitled "ICH Workshop on Oncolytic Viruses." The workshop will be held in conjunction with the International Conference on Harmonisation (ICH) expert working group and steering committee meetings in Chicago, IL. The objective of the workshop is to identify and discuss issues relevant to clinical development

of oncolytic viruses including safety. The following viruses will be covered: Adenovirus, herpes simplex virus, reovirus, Newcastle disease virus, measles virus, and Sendai virus. Speakers will address selectivity, attenuation modes, shedding, clinical and viral safety, and proof of concept in support of the approach in animal and human setting.

Date and Time: The workshop will be held on November 7, 2005, from 8 a.m. to 5 p.m.

Location: The public workshop will be held at the Westin Michigan Avenue, 909 North Michigan Ave., Chicago, IL 60611.

Contact Person: Daniel Takefman, Center for Biologics Evaluation and Research (HFM-720), Food and Drug Administration, Rockville, MD 20852, 301-827-5102, e-mail: daniel.takefman@fda.hhs.gov.

Registration: Registrations are being collected by the Pharmaceutical Research and Manufacturers of America (PhRMA). Send registration information (including name, title, firm name, address, telephone, and fax number) to Liz Cross at PhRMA by FAX: 202-572-7797, or e-mail: lcross@phrma.org, no later than Friday, October 14, 2005. The registration fee for this workshop is \$450 for industry; \$175 for academia and government participants. To register via the Internet go to <http://>