

review, institutions that might otherwise be subject to the RCR policy are under no obligation to implement the policy unless further public notice is issued in the **Federal Register**. Any future PHS action taken to implement the RCR policy would provide extended implementation time frames that take into consideration this suspension.

**FOR FURTHER INFORMATION CONTACT:** Barbara Bullman, J.D., Senior Program Analyst, Division of Education and Integrity, Office of Research Integrity, 5515 Security Lane, Suite 700, Rockville, MD 20852, (301) 443-5300.

**Chris B. Pascal, J.D.,**  
 Director, Office of Research Integrity.  
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**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Administration for Children and Families**

**Proposed Information Collection Activity; Comment Request**

**Proposed Projects**

*Title:* TANF High Performance Bonus Report, Assessment of Medicaid and SCHIP Enrollment.

*OMB No.:* New Collection.  
*Description:* Public Law 104-93 (PRWORA) established the Temporary Assistance for Needy Families (TANF) Program. It also included provisions for rewarding States that attain the highest levels of success in achieving the legislative goals of that program. The purpose of this collection is to obtain data upon which to base the

computation for measuring State performance in meeting those goals by providing Medicaid and SCHIP work supports. DHHS will use the information to allocate the Medicaid/SCHIP portion of the bonus grant funds appropriated under the law and implemented by 45 CFR part 270 published on August 30, 2000. States will not be required to submit this information unless they elect to compete in a Medicaid/SCHIP measure for the TANF High Performance Bonus awards in Federal fiscal years 2002 or 2003, or any subsequent Federal fiscal year for which Congress authorizes and appropriates bonus funds.

*Respondents:* Respondents may include any of the 50 States, the District of Columbia, and the U.S. Territories of Guam, Puerto Rico, and the Virgin Islands.

**ANNUAL BURDEN ESTIMATES**

Instrument	No. of respondents	No. of responses per respondent	Average burden hours per response	Total burden hours
TANF high performance bonus report, assessment of Medicaid and SCHIP enrollment among individuals after leaving TANF assistance .....	54	2	40	4,320
Estimated total annual burden hours .....				4,320

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: February 14, 2001.  
**Bob Sargis,**  
 Reports Clearance Officer.  
 [FR Doc. 01-4188 Filed 2-20-01; 8:45 am]  
**BILLING CODE 4184-01-M**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Food and Drug Administration**

**Allergenic 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:* Allergenic 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 March 5, 2001, 8:30 a.m. to 5 p.m.

*Location:* Holiday Inn, Versailles Ballrooms I and II, 8120 Wisconsin Ave., Bethesda, MD.  
*Contact:* William Freas or Pearline K. Muckelvene, Center for Biologics Evaluation and Research (HFM-71), Food and Drug Administration, 1401 Rockville Pike, Rockville, MD 20852, 301-827-0314, or FDA Advisory Committee Information Line, 1-800-741-8138 (301-443-0572 in the Washington, DC area), code 12388. Please call the Information Line for up-to-date information on this meeting.  
*Agenda:* On March 5, 2001, the committee will hear updates on: (1) The Laboratory of Immunobiochemistry personnel, (2) lot release statistics, (3) new guidance documents, (4) research and standardization programs, and (5) a compliance report. The committee will discuss whether master seed stocks of mold strains used for allergenic extracts should be rederived to reduce a theoretical risk of transmissible spongiform encephalopathy transmission. The committee will also discuss the statistical power of clinical studies used to assess bioequivalence as it applies to allergen extract studies. In the afternoon, the committee will discuss particulates that appear in allergen extracts and the effect of these particulates on the safety and efficacy on these products. In closed session, the committee will receive a report on the status of an investigational new drug application and product license application supplement.  
*Procedure:* On March 5, 2001, from 8:30 a.m. to 3:30 p.m., the meeting is open to the