ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average hours per response	Total burden hours
IAATP: Trainee Survey Pre-Test Administration	1,200	1	.15	180
IAATP: Trainee Survey Follow-Up Administration	1,200	1	.10	120

Estimated Total Annual Burden Hours: 300

In compliance with the requirement of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Administration for Children and Families is publishing the following summary of a proposed collection for public comment. Copies of the proposed collection of information can be requested from 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: January 8, 2008.

Janean Chambers,

Reports Clearance Officer.

[FR Doc. 08–95 Filed 1–14–08; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request

Proposed Projects

Title: Statewide Automated Child Welfare Information System (SACWIS) Assessment Review Guide (SARGE).

OMB No.: 0970–0159.

Description: The Department of Health and Human Services (HHS) cannot fulfill its obligation to effectively serve the nation's adoption and foster care populations, nor report meaningful and reliable information to Congress about the extent of problems facing these children or the effectiveness of assistance provided to this population, without access to timely and accurate information. Currently, SACWIS support State efforts to meet the following Federal reporting requirements: The Adoption and Foster Care Analysis and Reporting System (AFCARS) required by section 479(b)(2) of the Social Security Act; the National Child Abuse and Neglect Data System (NCANDS); Child Abuse Prevention and Treatment Act (CAPTA); and the Chafee Independent Living Program. These systems also support State efforts to provide the information to conduct the Child and Family Service Reviews. Currently, 42 States and the District of Columbia have developed, or are developing, a SACWIS with Federal financial participation. The purpose of these reviews is to ensure that all aspects of the project, as described in the approved Advance Planning Document, have been adequately completed, and conform to applicable regulations and policies.

To initiate a review, States will submit the completed SACWIS Assessment Review Guide (SARGE) and other documentation at the point that they have completed system development and the system is operational statewide. The additional documents submitted as part of this process should all be readily available to the State as a result of good project management practices.

The information collected in the SACWIS Assessment Review Guide will allow State and Federal officials to determine if the State's SACWIS meets the requirements of title IV–E Federal Financial Participation (FFP) defined at 45 CFR 1355.50. Additionally, other States will be able to use the documentation provided as part of this review process in their own system development efforts.

Respondents: State Title IV–E Agencies.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
SACWIS Assessment Review Guide (SARGE)	3	1	250	750

Estimated Total Annual Burden Hours: 750.

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: January 8, 2008.

Janean Chambers,

Reports Clearance Officer.

[FR Doc. 08–96 Filed 1–14–08: 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2006D–0419]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Voluntary National Retail Food Regulatory Program Standards

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 February 14, 2008.

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: 202–395–6974, or e-mailed to *baguilar@omb.eop.gov*. All comments should be identified with the OMB control number 0910–NEW and title, Voluntary National Retail Food Regulatory Program Standards. Also include the FDA docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Jonna Capezzuto, Office of the Chief Information Officer (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–4659.

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.

Voluntary National Retail Food Regulatory Program Standards

The Program Standards define nine essential elements of an effective regulatory program for retail food establishments, establish basic quality control criteria for each element, and provide a means of recognition for those state, local, and tribal regulatory programs that meet the Program Standards. The program elements addressed by the Program Standards are as follows: (1) Regulatory foundation, (2) trained regulatory staff, (3) inspection program based on Hazard Analysis and Critical Control Point (HACCP) principles, (4) uniform inspection program, (5) foodborne illness and food defense preparedness and response, (6) compliance and enforcement, (7) industry and community relations, (8) program support and resources, and (9) program assessment. Each standard includes a list of records needed to document compliance with the standard (referred to in the Program Standards document as ''quality records'') and has one or more corresponding appendices that contain forms and worksheets to facilitate the collection of information needed to assess the retail food regulatory program against that standard. The respondents are state, local and tribal government agencies. Regulatory agencies may use existing, available records or may choose to develop and use alternate forms and worksheets that capture the same information.

In the course of their normal activities, state, local, and tribal regulatory agencies already collect and keep on file many of the records needed as quality records to document compliance with each of the Program Standards. Although the detail and format in which this information is collected and recorded may vary by jurisdiction, records that are kept as a usual and customary part of normal agency activities include inspection records, written quality assurance procedures and records of quality assurance checks, staff training certificates and other training records, a log or database of food-related illness or injury complaints, records of investigations resulting from such complaints, an inventory of inspection equipment, records of outside audits, and records of outreach efforts (e.g., meeting agendas and minutes, documentation of food safety education activities). No new recordkeeping burden is associated with these existing records, which are already a part of usual and customary program recordkeeping activities by state, local, and tribal regulatory agencies, and which can serve as quality records under the Program Standards.

State, local, and tribal regulatory agencies that enroll in the Program Standards and seek listing in the FDA National Registry are required to report to FDA on the completion of the following three management tasks outlined in the Program Standards: (1) Conducting a program self assessment; (2) conducting a baseline survey of the regulated industry; and (3) obtaining an independent outside audit (verification audit). All three tasks must initially be completed within a 3-year time span. The results are reported to FDA on Form FDA 3519, ''FDA National Registry Report'' and Form FDA 3520, ''Permission to Publish in National Registry.'' These forms are located in Appendix I of the Program Standards document. If a regulatory agency follows all the recordkeeping recommendations in the individual standards and their appendices, it will have all the information needed to complete the forms. The time required to complete the forms is minimal.

In the **Federal Register** of November 14, 2006 (71 FR 66337), FDA published a 60-day notice requesting public comment on the information collection provisions. FDA received two letters in response to the notice, but the letters contained comments that were not responsive to the four PRA comment requests. These comments will not be addressed in this document.

In April 2006, the Conference for Food Protection approved changes to the Program Standards that have been incorporated into a draft 2007 revision, which is available at *http://www.cfsan.fda.gov/~dms/ret4toc.html*. FDA analyzed whether incorporation of the changes alters its estimate of the recordkeeping and reporting burdens as set forth in the 60-day notice. FDA concluded that the changes cause a minor increase and decrease in the recordkeeping burden, resulting in no net change in the recordkeeping burden estimate. FDA further concluded that the reporting burden estimate should be increased by adding a line to table 2 to reflect the addition of 150 hours. This is because the revision to Standard 2 establishes an Assessment of Training Needs (ATN) process and forms that can be used by regulatory retail food program managers/training officers to prepare Food Safety Inspection officers (FSIOs) to conduct retail food and foodservice inspections (new Appendix B–2 and its accompanying Attachments