

Responses: 28,050; *Total Annual Hours:* 28,050. (For policy questions regarding this collection contact Bridgett Rider at 410–786–2602.)

5. *Type of Information Collection Request:* Extension of a currently approved collection; *Title of Information Collection:* Eligibility Error Rate Measurement in Medicaid and the Children’s Health Insurance Program; *Use:* The Improper Payments Information Act (IPIA) of 2002 requires CMS to produce national error rates for Medicaid and the Children’s Health Insurance Program (CHIP). To comply with the IPIA, CMS will use a national contracting strategy to produce error rates for Medicaid and CHIP fee-for-service and managed care improper payments. The federal contractor will review States on a rotational basis so that each State will be measured for improper payments, in each program, once and only once every three years. Subsequent to the first publication, we determined that we will measure Medicaid and CHIP in the same State. Therefore, States will measure Medicaid and CHIP eligibility in the same year measured for fee-for-service and managed care. We believe this approach will advantage States through economies of scale (e.g. administrative ease and shared staffing for both programs reviews). We also determined that interim case completion timeframes and reporting are critical to the integrity of the reviews and to keep the reviews on schedule to produce a timely error rate. Lastly, the sample sizes were increased slightly in order to produce an equal sample size per strata each month. Periodically, CMS will conduct Federal

re-reviews of States’ PERM files to ensure the accuracy of States’ review findings and the validity of the review process. CMS will select a random subsample of Medicaid and CHIP cases from the sample selection lists provided by each State. States will submit all pertinent information related to the review of each sampled case that is selected by CMS. *Form Number:* CMS–10184 (OMB control number: 0938–1012); *Frequency:* Annually, Quarterly *Affected Public:* State, Local, or Tribal Governments; *Number of Respondents:* 34; *Total Annual Responses:* 1,583; *Total Annual Hours:* 946,164. (For policy questions regarding this collection contact Bridgett Rider at 410–786–2602.)

Dated: April 5, 2016.
William N. Parham, III,
Director, Paperwork Reduction Staff, Office of Strategic Operations and Regulatory Affairs.
[FR Doc. 2016–08106 Filed 4–7–16; 8:45 am]
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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request

Title: National Survey of Child and Adolescent Well-Being-Third Cohort (NSCAW III): Agency Recruitment.
OMB No.: 0970–0202.
Description: The Administration for Children and Families (ACF) within the U.S. Department of Health and Human

Services (HHS) intends to collect data on a third cohort of children and families for the National Survey of Child and Adolescent Well-Being (NSCAW). NSCAW is the only source of nationally representative, longitudinal, firsthand information about the functioning and well-being, service needs, and service utilization of children and families who come to the attention of the child welfare system. The first two cohorts of NSCAW were collected beginning in 1999 and 2008 and studied children who had been the subject of investigation by Child Protective Services. Children were sampled from child welfare agencies nationwide. The proposed data collection plan for the third cohort of NSCAW includes two phases: Phase 1 includes child welfare agency recruitment and collection of files for sampling children, and Phase 2 includes baseline data collection and an 18-month follow-up data collection. The current data collection plan calls for selecting a new cohort of 4,565 children and families and repeating similar data collection procedures as the previous two cohorts. This Notice is specific to Phase 1. The overall goal is to recruit child welfare agencies in 83 primary sampling units nationwide. Child welfare agencies will be selected with probability proportional to size, based on the current distributions in the child welfare system. Child welfare agency recruitment will include: mail, email, phone calls, and site visits with child welfare agency administrators.

Respondents: Child welfare agency administrators and other personnel. Data collection will take place over a 2-year period.

ANNUAL BURDEN ESTIMATES					
Instrument	Total number of respondents	Annual number of respondents (rounded)	Number of responses per respondent	Average burden hours per response	Annual burden hours
Information package for agency administrators	83	42	1	.25	11
Initial visit or call with agency staff	83	42	1	1	42
Visit or call with agency staff explaining the sample file process	83	42	1	2	84
Agency staff monthly sample file generation and transmission	83	42	15	1	630

Estimated Total Annual Burden Hours: 767.
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 Planning, Research and Evaluation, 330 C Street SW., Washington, DC 20201, Attn: OPRE Reports Clearance Officer. Email address: OPREinfocollection@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,

ACF Certifying Officer.

[FR Doc. 2016-08018 Filed 4-7-16; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Title: Assets for Independence Program Performance Progress Report.
OMB No.: New.

Description: The Assets for Independence (AFI) Act (Title IV of the Community Opportunities, Accountability, and Training and Educational Services Act of 1998, Pub. L. 105-285, [42 U.S.C. 604 note]) requires that organizations operating AFI projects submit annual progress reports.

This request is to create an AFI program specific Performance Progress Report (PPR) to replace the semiannual standard form performance progress report (SF-PPR) and the annual data

report. The AFI PPR will collect data on project activities and attributes similar to the reports that it is replacing. The Office of Community Services (OCS) in the Administration for Children and Families (ACF) will use the data collected in the AFI PPR to prepare the annual AFI Report to Congress, to evaluate and monitor the performance of the AFI program overall and of individual projects, and to inform and support technical assistance efforts. The AFI PPR would fulfill AFI Act reporting requirements and program purposes.

The AFI PPR will be submitted quarterly: three times per year using an abbreviated short form and one time using a long form. Both draft data collection instruments are available for review online at <http://idaresources.acf.hhs.gov/AFIPPR>.

Respondents: Assets for Independence (AFI) program grantees.

Annual Burden Estimates:

Form name	Number of responses	Number of responses per respondent	Average burden hours per response	Total burden hours
AFI PPR Short Form	300	3	0.5	450
AFI PPR Long Form	300	1	3.8	1,140
Estimated Annual Burden Hours				1,590

Additional Information

Copies of the proposed collection may be obtained by writing to the Administration for Children and Families, Office of Planning, Research and Evaluation, 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. Email address: infocollection@acf.hhs.gov.

OMB Comment

OMB is required to make a decision concerning the collection of information between 30 and 60 days after publication of this document in the **Federal Register**. Therefore, a comment is best assured of having its full effect if OMB receives it within 30 days of publication. Written comments and recommendations for the proposed information collection should be sent directly to the following: Office of Management and Budget, Paperwork Reduction Project, Email: OIRA.SUBMISSION@OMB.EOP.GOV, Attn: Desk Officer for the Administration for Children and Families.

Robert Sargis,

Reports Clearance Officer.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA-2013-N-0519]

Agency Information Collection Activities; Proposed Collection; Comment Request; Guidance for Industry on How To Submit Information in Electronic Format to the Center for Veterinary Medicine Using the Food and Drug Administration Electronic Submission Gateway

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA, we) is announcing an opportunity for public comment on the proposed collection of certain information by the Agency. Under the Paperwork Reduction Act of 1995 (the PRA), Federal Agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, and to allow 60 days for public comment in response to the notice. This notice solicits comments on extending Office

of Management and Budget (OMB) approval on the existing reporting requirements relating to how one may submit information electronically to the Center for Veterinary Medicine (CVM) using the FDA Electronic Submissions Gateway (ESG).

DATES: Submit either electronic or written comments on the collection of information by June 7, 2016.

ADDRESSES: You may submit comments as follows:

Electronic Submissions

Submit electronic comments in the following way:

- Federal eRulemaking Portal: <http://www.regulations.gov>. Follow the instructions for submitting comments. Comments submitted electronically, including attachments, to <http://www.regulations.gov> will be posted to the docket unchanged. Because your comment will be made public, you are solely responsible for ensuring that your comment does not include any confidential information that you or a third party may not wish to be posted, such as medical information, your or anyone else's Social Security number, or confidential business information, such as a manufacturing process. Please note that if you include your name, contact information, or other information that