reached by a grant recipient or any partner organizations:

- Type of Trafficking (Labor, Sex, Labor and Sex, Unknown);
- Client Identifier (e.g., Initials, Date of Birth, and Country of Origin);
- Client information (Sex, Adult/Minor);
- Description of trafficking situation;
- Date that organization made contact with the victim began establishing trust and/or screened the person for victim status;
- Date that grantee positively identified person as a victim of a severe form of trafficking in persons;
- Documentation from the Department of Homeland Security (DHS) about the time of temporary status the victim is pursuing (e.g., Continued Presence, T Visa, U Visa, SIJS);
- Name of service agency assisting the victim;
- Date of HHS Certification or Eligibility; and
- Date the agency or victim terminated contact, with space for explanation.

Respondents: Rescue & Restore Victims of Human Trafficking Regional Program grantees.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Excel spreadsheet	20	4	4	16

Estimated Total Annual Burden Hours: 320.

In compliance with the requirements of Section 506(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, 370 L'Enfant Promenade, SW., Washington, DC 20447, Attn: ACF Reports Clearance Officer. Email 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.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 2014–07606 Filed 4–4–14; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Title: Permanency Innovations Initiative Evaluation: Phase 3.

OMB No.: 0970–0408.

Description: The Administration for Children and Families (ACF), U.S. Department of Health and Human Services (HHS) intends to collect data for an evaluation of the Permanency Innovations Initiative (PII). This 5-year initiative, funded by the Children's Bureau (CB) within ACF, is intended to build the evidence base for innovative interventions that enhance well-being and improve permanency outcomes for particular groups of children and youth who are at risk for long-term foster care and who experience the most serious barriers to timely permanency. The CB funded six grantees to identify local barriers to permanent placement and implement innovative strategies that mitigate or eliminate those barriers and reduce the likelihood that children will remain in foster care for 3 years or longer. In addition, evaluation plans were developed to support rigorous site-specific and cross-site studies to document the implementation and effectiveness of the grantees' projects and the initiative overall.

Data collection for the PII evaluation includes a number of components being launched at different points in time. Phase 1 included data collection for a cross-site implementation evaluation and site-specific evaluations of two PII grantees (approved August 2012; Washoe County, Nevada, and the State of Kansas). Phase 2 (approved August 2013) included data collection for site-specific evaluations of two PII grantees: Illinois Department of Children and Family Services (DCFS) and the Los Angeles Gay and Lesbian Center's Recognize Intervene Support Empower (RISE) project. Phase 3 includes data collection for a cross-site cost study, additional data collection components for the RISE project, and a cross-site administrative data study assessing outcomes. Phase 4 will include data collection for the California Department of Social Services' California Partnership for Permanency (CAPP) project. Data for the evaluations are collected through surveys of children, youth, foster parents, guardians, biological parents, permanency resources, and caseworkers, supervisors, administrators/managers, and other agency staff. The administrative data study does not impose any new data collection requirements but uses data already compiled and reported by the states.

Respondents: Children/youth and their parents, guardians, permanency resources, or foster caregivers; caseworkers, supervisors, administrators/managers, or other agency staff.

ANNUAL BURDEN ESTIMATES

Instrument	Annual number of respondents	Number of responses per respondent	Average burden hours per response	Total annual burden hours
RISE CCT Youth Interview (ages 11–19)	22	2	1.3	57
RISE CCT Qualitative Youth Interview (ages 11–19)	22	1	1.2	26
RISE CCT Facilitator Interview (Facilitator Burden)	2	4	0.2	2
RISE CCT Facilitator Interview (Child burden)	3	2	.5	3
RISE CCT Facilitator Survey	2	21	0.2	8
RISE CCT Facilitator submission of CAFAS data [1]	2	21	0.1	4
RISE CCT Permanency Resource Interview	11	2	1.0	22
RISE CCT Interview with Current Caregiver	11	2	0.6	13
RISE CCT burden	.....	.....	.....	135
RISE ORB Staff Follow-Up Survey	157	1	0.3	47
RISE ORB burden	.....	.....	.....	47
Cost Study Focus Group Preparation	9	1	1.5	14
Cost Study Focus Group	9	1	4.0	36
Trial Administration of Cost Study Activity Logs	9	1	1.5	14
Weekly Case Work Activity Log	123	52	0.4	2,558
Weekly Supervision Activity Log	39	52	0.4	811
Monthly Management/Administration Log	30	12	0.5	180
Cost study burden	.....	.....	.....	3,613
Administrative data submission, no added fields	1	12	0.3	2
Administrative data submission with added fields	1	12	0.8	10
Administrative data study burden	.....	.....	.....	12

[1] The CAFAS is administered as part of case planning, so the only burden is in submitting the CAFAS data to the evaluation team.

Estimated Total Annual Burden Hours: 3,807.

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: OPRE Reports Clearance Officer. All requests should be identified by the title of the information collection. Email address: *OPREinfocollection@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.

Karl Koerper,

OPRE Reports Clearance Officer.

[FR Doc. 2014–07614 Filed 4–4–14; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2004–N–0193]

Agency Information Collection Activities: Proposed Collection; Comment Request; Current Good Manufacturing Practice Regulations for Medicated Feeds

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) 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 the recordkeeping requirements for manufacturers of medicated animal feeds.

DATES: Submit either electronic or written comments on the collection of information by June 6, 2014.

ADDRESSES: Submit electronic comments on the collection of information to *http://www.regulations.gov.* Submit written comments on the collection of information to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852. All comments should be identified with the docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: FDA PRA Staff, Office of Operations, Food and Drug Administration, 1350 Piccard Dr., PI50–400B, Rockville, MD 20850, *PRAStaff@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION: Under the PRA (44 U.S.C. 3501–3520), Federal Agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. ''Collection of information'' is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes Agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA (44 U.S.C. 3506(c)(2)(A)) requires Federal