finances. This information is also included in a published annual statistical and financial report, available to the general public.

Pub. L. 109–171, the Deficit Reduction Act of 2005, contains a number of provisions that will impact the States' completion and submission of these quarterly financial reports. These changes become effective in fiscal years 2006, 2007 and 2008. These changes require revisions to some of the data entry lines and reporting instructions currently contained on these forms. In addition, a periodic review of the data currently requested on these forms will assure that OCSE collections the information needed in the most efficient format feasible.

Respondents: State agencies administering the Child Support Enforcement Program.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
OCSE–396A	54	4	8	1,728
OCSE–34A	54	4	8	1,728
Estimate Total Annual Burden Hours:				3,456

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. E-mail address: *infocollection@acf.hhs.gov.*

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: November 28, 2006.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 06–9503 Filed 12–1–06; 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: Third Grade Follow-up to the Heat Start Impact Study.

OMB No.: 0970–0229.

Description: The Administration for Children and Families (ACF) within the Department of Health and Human Services (HHS) is requesting comments on plans to implement a third grade follow-up to the Head Start Impact Study. This study will collect information for determining, on a national basis, how Head Start affects outcomes in the third grade for children who participated in the program as compared to children not enrolled in Head Start and to determine under which conditions Head Start Works best and for which children.

The Head Start Impact Study was longitudinal study that involved approximately 5,000 first-time-enrolled, three- and four-year-old preschool children across 84 nationally representative grantee/delegate agencies (in communities where there were more eligible children and families than can be served by the program). The participating children were randomly assigned to either a Head Start group (that could enroll in Head Start services) or a control group (that could not enroll in Head Start services but could enroll in other available services selected by their parents). Data collection for the study began in the fall of 2002 and extended through spring 2006.

It is the intention of the Administration for Children and Families to examine outcomes for this sample of children and families during the spring of the children's third grade year. Data will be collected in the spring of 2007 (for the four-year-old cohort) and the spring of 2008 (for the three-year-old cohort). The domains of development to be assessed include demographic characteristics of the children and families, as well as children's cognitive development, school achievement and adjustment, socio-emotional functioning, heath and access to health care, and relationships with peers. Information will also be collected on parents' involvement in educational activities, mental health and well-being, and monitoring and other parenting practices, and information related to the characteristics and quality of the schools and classrooms that children attend.

Respondents: Individuals or households and school districts (principals and teachers).

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Year 1 (spring 2007):				
Parent Tracking Interview	2,559	1	0.166	426
Parent Interview	1,720	1	1.00	1,720

ANNUAL BURDEN ESTIMATES—Continued

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Child Assessment	1,720	1	1.00	1,720
Teacher Survey/TCR	2,580	1	0.50	1,290
Principal Survey	1,462	1	0.33	487
Year 2 (fall 2007):				
Parent Tracking Interview	4,667	1	0.166	778
Year 2 (spring 2008):				
Parent Tracking Interview	2,108	1	0.166	351
Parent Interview	2,042	1	1.00	2,042
Child Assessment	2,042	1	1.00	2,042
Teacher Survey/TCR	3,063	1	0.50	1,532
Principal	1,736	1	0.33	579

Estimated Total Annual Burden Hours: 6,484

Additional Information: Copies of the proposed collection may be obtained 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. E-mail address: *infocollection@ach.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, 725 17th Street, NW., Washington, DC 20503, E-mail address: *Karen_T._Matsuoka@omb.eop.gov*, Attn: Desk Officer for ACF.

Dated: November 28, 2006.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 06–9504 Filed 12–1–06; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2006N–0472]

Agency Information Collection Activities; Proposed Collection; Comment Request; Substances Prohibited from Use in Animal Food or Feed; Animal Proteins Prohibited in Ruminant Feed

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 extending OMB approval on the existing recordkeeping requirements for this information collection, regarding animal proteins prohibited in ruminant feed.

DATES: Submit written or electronic comments on the collection of information by February 2, 2007.

ADDRESSES: Submit electronic comments on the collection of information to: *http://www.fda.gov/dockets/ecomments.* 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: Denver Presley, Jr., Office of the Chief Information Officer (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–1472.

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 agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, FDA is publishing notice of the proposed collection of information set forth in this document.

With respect to the following collection of information, FDA invites comments on these topics: (1) Whether the proposed collection of information is necessary for the proper performance of FDA's functions, including whether the information will have practical utility; (2) the accuracy of FDA's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques, when appropriate, and other forms of information technology.

Substances Prohibited From Use in Animal Food or Feed; Animal Proteins Prohibited in Ruminant Feed—21 CFR 589.2000(e)(1)(iv) (OMB Control Number 0910–0339)—Extension

This information collection was established because epidemiological evidence gathered in the United Kingdom suggested that bovine spongiform encephalopathy (BSE), a progressively degenerative central nervous system disease, is spread to ruminant animals by feeding protein derived from ruminants infected with BSE. This regulation places general requirements on persons that manufacture, blend, process, and