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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request

Title: Head Start Family and Child Experiences Survey (FACES).

OMB No.: 0970-0151.

Description: The Office of Planning, Research and Evaluation (OPRE), Administration for Children and Families (ACF), U.S. Department of Health and Human Services (HHS), is proposing to collect data for the Head Start Family and Child Experiences Survey (FACES). Featuring a new “Core Plus” study design, FACES will provide data on a set of key indicators, including information for performance measures. This design also allows for more rapid and frequent data reporting (Core study) and serves as a vehicle for studying more complex issues and topics in greater detail and with increased efficiency (Plus studies).

The FACES Core study will assess the school readiness skills of Head Start children, survey their parents, and ask their Head Start teachers to rate children’s social and emotional skills. In addition, FACES will include observations in Head Start classrooms, and program director, center director, and teacher surveys. FACES Plus studies include additional survey content of policy or programmatic interest, and may include additional programs or respondents beyond those participating in the Core FACES study.

Previous notices provided the opportunity for public comment on the proposed Head Start program recruitment and center selection process

(FR V.78, pg. 75569 12/12/2013; FR V.79, pg. 8461 02/12/2014). This notice describes the planned data collection activities for the FACES Core study and Plus studies. Direct child assessments, parent surveys, and teacher child reports for the Core study are included in this clearance package. Additionally, we describe instruments to support the Core study at the program and classroom levels and the Plus studies anticipated for future submission. Since these instruments will be informed by initial findings of FACES and emerging policy needs, they cannot be fully specified at this time. However, we describe the respondents and data collection methods, estimated respondent burden, and how the information will be used to the extent possible at this time. Subsequently, when fully developed in a manner consistent with the description provided in this 60-day notice and prior to use, we will submit these materials for a 30-day public comment period under the Paperwork Reduction Act.

Methods for Core data collection start with site visits in fall 2014 to 60 Head Start programs to directly assess the school readiness skills of 2,400 children sampled to participate in FACES. Parents of sampled children will complete surveys on the Web or by telephone about their children and family background. Head Start teachers will rate each sampled child (approximately 10 children per classroom) using the Web or paper-and-pencil forms. These activities will occur a second time in spring 2015. Additionally, the program sample size will increase to 180 programs in the spring to collect program- and classroom-level data. The methods of data collection for this phase will feature site visitors conducting observations of the types and quality of classroom activities. Head Start program

directors, center directors, and teachers will complete surveys about themselves and the services and instruction at Head Start. The program- and classroom-level data collection will occur a second time in spring 2017.

Plus study data collection will parallel the Core design in many ways, including recruitment and data collection procedures, to add new respondents, include new populations, or expand on the information gathered in the Core study. Additional early care and education administrators or providers (such as Education Coordinators or Family Service Staff) may be sampled. Plus studies may involve data collection in additional programs, such as programs serving different populations or programs implementing specific interventions. Data collection for these Plus studies may include child assessments, parent surveys, teacher child reports, and staff surveys. Plus studies may also feature topical modules to gather information in greater depth on particular topics (for example, parent engagement or program functioning). The methods of data collection will involve new methodologies such as qualitative interviews and supplemental surveys with expanded content.

The purpose of the Core data collection is to support the 2007 reauthorization of the Head Start program (P.L. 110-134), which calls for periodic assessments of Head Start’s quality and effectiveness. Plus data collection will further support understanding Head Start functioning for a broader set of programs or in more depth for particular topics.

Annual Burden Estimates

Respondents: Head Start children, parents of Head Start children, and Head Start teachers, directors, and other early care and education program staff.

Instrument	Total number of respondents	Annual number of respondents	Number of responses per respondent	Average burden hour per response	Estimated annual burden hours
Core Study					
Head Start core child assessment	2,400	800	2	0.75	1,200
Head Start core parent survey	2,400	800	2	0.25	400
Head Start fall parent supplemental survey	2,400	800	1	0.08	64
Head Start spring parent supplemental survey	2,400	800	1	0.08	64
Head Start core teacher child report	240	80	20	0.17	272
Head Start core teacher survey	720	240	2	0.50	240
Head Start core program director survey	180	60	2	0.25	30
Head Start core center director survey	360	120	2	0.25	60
Plus Studies					
Head Start parent qualitative interview	400	133	2	1.00	267
Head Start staff qualitative interview	300	100	2	1.00	200
Head Start child assessment for plus study	1,350	450	2	0.75	675

Instrument	Total number of respondents	Annual number of respondents	Number of responses per respondent	Average burden hour per response	Estimated annual burden hours
Head Start parent survey for plus study	1,350	450	2	0.25	225
Head Start parent supplemental survey for plus study	1,350	450	2	0.08	72
Head Start teacher child report for plus study	150	50	20	0.17	170
Head Start teacher survey for plus study	150	50	2	0.50	50
Head Start program director survey for plus study	50	17	2	0.25	8
Head Start center director survey for plus study	100	33	2	0.25	17
Early care and education administrators plus survey	600	200	2	0.50	200
Early care and education providers plus survey	900	300	2	0.50	300
Total					4,514

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, 370 L'Enfant Promenade, SW., Washington, DC 20447, 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.

Karl Koerper,

OPRE Reports Clearance Officer.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA-2013-N-1620]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Information From United States Firms and Processors That Export to the European Community

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 March 31, 2014.

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: 202-395-7285, or emailed to oira_submission@omb.eop.gov. All comments should be identified with the OMB control number 0910-0320. Also include the FDA 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: In compliance with 44 U.S.C. 3507, FDA has submitted the following proposed collection of information to OMB for review and clearance.

Information From United States Firms and Processors That Export to the European Community (OMB Control Number 0910-0320)—Extension

The European Community (EC) is a group of 27 European countries that have agreed to harmonize their commodity requirements to facilitate commerce among member States. EC legislation for intra-EC trade has been extended to trade with non-EC countries, including the United States. For certain food products, including those listed in this document, EC legislation requires assurances from the responsible authority of the country of origin that the processor of the food is in compliance with applicable regulatory requirements. The European Commission, the executive branch of the EC, requires countries trading with any of the EC member countries to provide lists of firms and processors approved to export certain animal-derived commodities to the EC. As stated in the notice published in the **Federal Register** of April 4, 1996 (61 FR 15077), we established a list of U.S. firms and processors that intended to export shell eggs, dairy products, and game meat and game meat products to the EC.

Although our 1996 **Federal Register** notice did not include on the list firms and processors exporting raw, bulk collagen, and gelatin intended for human consumption, EC directives require that shipments of raw, bulk collagen, and gelatin products be accompanied by certification stating that the product, derived from ruminant bones, bovine hides, and pigskins, has been produced in compliance with EC Council Directive 2003/863/EC. The directive contains the requirements for sourcing, manufacture, transport, and storage of raw materials and manufacture of finished products. Chapter III, Article 23, of the directive requires lists identifying non-EC firms and processors that meet EC requirements and have the appropriate animal and public health certificates.