comments and recommendations for the proposed information collection should be sent directly to the following: Office of Management and Budget, Paperwork Reduction Project, Fax: 202–395–7245, Email: *OIRA_SUBMISSION@OMB.EOP.GOV*, Attn: Desk Officer for the Administration for Children and Families.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 2013–02405 Filed 2–4–13; 8:45 am]

BILLING CODE P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request

Title: Personal Responsibility Education Program (PREP) Multi-Component Evaluation—Data Collection Related to the Performance Analysis Study and the Impact and the In-depth Implementation Study.

OMB No.: 0970–0398

Description: The Office of Data Analysis, Research, and Evaluation (HHS/ACF/ACYF/ODARE) in the Administration for Children, Youth and Families (ACYF) and the Office of Planning, Research, and Evaluation (HHS/ACF/OPRE) in the Administration for Children and Families (ACF) propose a data collection activity as part of the Personal Responsibility Education Program (PREP) Multi-Component Evaluation. The goals of the PREP Multi-Component Evaluation are to document how PREP programs are designed and implemented in the field, collect performance measure data for PREP programs, and assess the effectiveness of selected PREP-funded programs.

The PREP Multi-Component Evaluation contains three components: the ''Design and Implementation Study,'' the ''Performance Analysis Study,'' and the ''Impact and In-Depth Implementation Study.'' This notice is specific to data collection activities for both the Performance Analysis Study and the Impact and In-Depth Implementation Study.

The Performance Analysis Study component entails the development of performance measures and collection and analysis of performance measure data from PREP and Competitive PREP (CPREP) grantees. Data will be used to determine if PREP and CPREP grantees are meeting performance benchmarks related to the program's mission and priorities.

The Impact and In-depth Study component entails separate random assignment impact analyses accompanied by in-depth implementation analyses of four-to-five specific PREP-funded programs. The Impact and In-Depth Implementation Study will expand available evidence on whether the replication of evidence-based programs, or the substantial incorporation of elements of these programs, funded as part of PREP, are effective at delaying sexual activity, increasing condom or contraceptive use for sexually active youth, or reducing pregnancy among youth.

A description of the activities proposed follows:

Performance Analysis—CPREP Performance Measure Data Collection: Performance measurement data collection instruments will be administered to individuals representing CPREP grantees, CPREP program facilitators, other CPREP program staff, and CPREP program participants.

Impact Analysis—Follow-up Instruments: Two follow-up surveys (one at approximately 6 months post-program and one at approximately 12–24 months post-program) will be administered to study participants. Statistical analyses will be performed with these data to determine program effects.

Impact Analysis—Administrative Data Collection: Data will also be collected from administrative records (e.g. school or state agency data bases) and used to determine program effects.

In-depth Implementation Analysis—Implementation Instruments: Site visits will be conducted at two to three points during the program implementation period. During these site visits, in-person interviews, focus groups, and a web survey of program staff will be conducted. Data collected through these activities will provide an understanding of the selected programs' implementation and context and will allow for a description of each program and the treatment-control contrast to be evaluated in each site. Data will also help in interpreting impact findings.

Respondents: Program applicants (i.e., adolescents); Data managers (e.g., at schools or state agencies); Program administrators and staff; Participating youth; and Community members.

ANNUAL BURDEN ESTIMATES

Instrument	Total number of respondents	Number of responses per respondent	Average burden hours per response	Estimated total burden hours	Estimated annual burden hours
Performance Analysis: Competitive PREP Performance Measure Data Collection					
Participant entry survey	60,420	1	.08333	5,035	1,678
Participant exit survey	48,336	1	.16667	8,056	2,685
Performance reporting system data form	111	1	30	3,330	1,110
Implementation site data collection	900	1	14	12,600	4,200
Estimated Annual Burden—Subtotal for Performance Measure Data Collection					9,673
Impact Analysis: Follow-up Instruments					
First follow-up survey with program applicants	4,800	1	0.75	3,600	1,200
Second follow-up survey with program applicants	2,250	1	0.75	1,688	563
Estimated Annual Burden—Sub-total for Follow-up Instruments					1,763

ANNUAL BURDEN ESTIMATES—Continued

Instrument	Total number of respondents	Number of responses per respondent	Average burden hours per response	Estimated total burden hours	Estimated annual burden hours
Impact Analysis: Administrative Data Collection					
Administrative records	6	1	6	36	12
Estimated Annual Burden—Sub-total for Administrative Data Collection Instrument					12
In-depth Implementation Analysis: Implementation Instruments					
Focus group guide with participants	320	1	1.5	480	160
Semi-structured interview topic guide with program staff and other stakeholders	160	2	1	320	107
Web survey of program staff	100	2	0.5	100	33
Estimated Annual Burden—Sub-total for Implementation Instruments					300
TOTAL Estimated Annual Burden					11,748

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.

Steven M. Hanmer,

Reports Clearance Officer.

[FR Doc. 2013–02295 Filed 2–4–13; 8:45 am]

BILLING CODE 4184–37–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Submission for OMB Review; Comment Request (30-Day FRN): The Agricultural Health Study: A Prospective Cohort Study of Cancer and Other Disease Among Men and Women in Agriculture (NCI)

Summary: Under the provisions of Section 3507(a)(1)(D) of the Paperwork Reduction Act of 1995, the National Institutes of Health (NIH), has submitted to the Office of Management and Budget (OMB) a request to review and approve the information collection listed below. This proposed information collection was previously published in the **Federal Register** on December 6, 2012 (Vol. # 77, p. 72871) and allowed 60 days for public comment. No public comments have been received. The purpose of this notice is to allow an additional 30 days for public comment. The National Institutes of Health may not conduct or sponsor, and the respondent is not required to respond to, an information collection that has been extended, revised, or implemented on or after October 1, 1995, unless it displays a currently valid OMB control number.

Written comments and/or suggestions regarding the item(s) contained in this notice, especially regarding the estimated public burden and associated response time, should be directed to the Attention: NIH Desk Officer, Office of Management and Budget, at *OIRA_submission@omb.eop.gov* or by fax to 202–395–6974. To request more information on the proposed project or to obtain a copy of the data collection plans and instruments, contact Jane Hoppin, Sc.D., Epidemiology Branch, National Institute of Environmental Health Sciences, NIH, 111 T.W. Alexander Drive, PO Box 12233, MD A3–05, Research Triangle Park, NC 27709, or call non-toll-free number 919–541–7622, or email your request, including your address to: *hoppin1@niehs.nih.gov.*

Comments regarding this information collection are best assured of having their full effect if received within 30 days of the date of this publication.

Proposed Collection: The Agricultural Health Study: A Prospective Cohort Study of Cancer and Other Disease Among Men and Women in Agriculture, 0925–0406, Expiration Date 5/31/2013—REVISION—National Institute of Environmental Health Sciences (NIEHS), National Institutes of Health (NIH).

Need and Use of Information Collection: The purpose of this information collection is to continue and complete updating the occupational and environmental exposure information as well as medical history information for licensed pesticide applicators and their spouses enrolled in the Agricultural Health Study. This represents a request to complete phase IV (2013–2015) of the study and to continue and complete the buccal cell collection and the Study of Biomarkers of Exposures and Effects in Agriculture (BEEA). The primary objectives of the study are to determine the health effects resulting from occupational and environmental exposures in the agricultural environment. The phase IV follow up data will be collected by using one of three methods of the cohort member's choosing: Self-administered