

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.

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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

computer assisted Web survey (CAWI); self-administered paper-and-pen (Paper/pen); or an interviewer administered computer assisted telephone interview (CATI). Proxy interviews for those cohort members unable to complete the follow up will be completed by using one of the three methods as well. Secondary objectives include evaluating biological markers that may be associated with agricultural exposures and risk of certain types of cancer.

Questionnaire data will be collected by using computer assisted telephone interview (CATI) and in-person interview (CAPI) systems for telephone screeners and home visit interviews, respectively. Some respondents will also be asked to participate in the collection of biospecimens including blood, urine, and buccal cells (loose cells from the respondent's mouth). The findings will provide valuable information concerning the potential

link between agricultural exposures and cancer and other chronic diseases among agricultural Health Study cohort members, and this information may be generalized to the entire agricultural community.

OMB approval is requested for 3 years. There are no costs to respondents other than their time. The total estimated annualized burden hours are 10,465.

Estimated Annualized Burden Hours

TABLE A.12-1—ESTIMATES ANNUALIZED BURDEN HOURS

Type of respondent	Form	Number of respondents	Number of responses per respondent	Average time per response (in hours)	Total annual burden hour
Private and Commercial Applicators and Spouses.	Reminder, Missing, and Damaged Scripts for Buccal Cell.	100	1	5/60	8
Private Applicators	BEEA CATI Screener	480	1	20/60	160
Private Applicators	BEEA Home Visit CAPI, Blood, & Urine x 1.	160	1	30/60	80
Private Applicators	BEEA Schedule Home Visit Script ..	20	3	5/60	5
Private Applicators	BEEA Home Visit CAPI, Blood, & Urine x 3.	20	3	30/60	30
Private Applicators	Paper/pen, CAWI or CATI	13,855	1	25/60	5,773
Spouses	Paper/pen, CAWI or CATI	10,201	1	25/60	4,250
Proxy	Paper/pen, CAWI or CATI	635	1	15/60	159
Total	10,465

Dated: January 30, 2013.

Vivian Horovitch-Kelley,

NCI Project Clearance Liaison, NCI, NIH.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Proposed Collection; Comment Request: Women's Health Initiative Observational Study

SUMMARY: In compliance with the requirement of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, for opportunity for public comment on proposed data collection projects, the

National Heart, Lung, and Blood Institute (NHLBI), the National Institutes of Health (NIH) will publish periodic summaries of proposed projects to be submitted to the Office of Management and Budget (OMB) for review and approval.

Proposed Collection: Title: The Women's Health Initiative (WHI) Observational Study. *Type of Information Collection Request:* Revision OMB #0925-0414. *Need and Use of Information Collection:* This study will be used by the NIH to evaluate risk factors for chronic disease among older women by developing and following a large cohort of postmenopausal women and relating subsequent disease development to baseline assessments of historical, physical, psychosocial, and physiologic

characteristics. In addition, the observational study will complement the clinical trial (which has received clinical exemption) and provide additional information on the common causes of frailty, disability and death for postmenopausal women, namely, coronary heart disease, breast and colorectal cancer, and osteoporotic fractures. Continuation of follow-up for ascertainment of medical history update forms will provide essential data for outcomes assessment for this population of aging women. *Frequency of Response:* Annually. *Affected Public:* Individuals or households and health care providers. *Type of Respondents:* Study participants, next-of-kin, and physician's office staff. The annual reporting burden is as follows:

ESTIMATE OF ANNUAL HOUR BURDEN

Type of respondent	Number of respondents	Frequency of response	Average time per response	Annual hour burden
OS Participants	41,495	1	20/60	13,929
Next of kin	936	1	6/60	92
Physician/Office Staff	17	1	5/60	1.4
Totals	42,448	14,023

¹ Annual burden is placed on health care providers and respondent relatives/informants through requests for information which will help in the compilation of the number and nature of new fatal and nonfatal events.