

Type of respondents	Estimated number of respondents	Estimated number of responses per respondent	Average burden hours per response	Estimated total annual burden hours requested
Adult Blood Donors	40,000	1	0.25	10,000

Request for Comments

Written comments and/or suggestions from the public and affected agencies are invited on one or more of the following points: (1) Whether the proposed collection of information is necessary for the proper performance of the function of the agency, including whether the information will have practical utility; (2) The accuracy of the agency'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 those who are to respond, including the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology.

Direct Comments to OMB

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: Office of Management and Budget, Office of Regulatory Affairs, New Executive Office Building, Room 10235, Washington, DC 20503, Attention: Desk Officer for NIH. To request more information on the proposed project or to obtain a copy of the data collection plans and instruments, contact: Dr. George J. Nemo, Group Leader, Transfusion Medicine, Scientific Research Group, Division of Blood Diseases and Resources, NHLBI, NIH, Two Rockledge Center, Suite 10042, 6701 Rockledge Drive, MSC 7950, Bethesda, MD 20892-7950, or call non-toll free number 301-435-0075, or e-mail your request, including your address to: nemog@nih.gov.

Comments Due Date

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

Dated: May 10, 2002.

Donald Christoferson,
Executive Officer, NHLBI.

[FR Doc. 02-13195 Filed 5-24-02; 8:45 am]

BILLING CODE 4140-01-M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Submission for OMB Review; Comment Request; Prostate, Lung, Colorectal and Ovarian Cancer Screening Trial

SUMMARY: Under the provisions of section 3507(a)(1)(D) of the Paperwork Reduction Act of 1995, the National Cancer Institute (NCI), the National Institutes of Health (NIH) has submitted to the Office of Management and Budget (OMB) a request for review and approval of the information collection listed below. This proposed information collection was previously published in the **Federal Register** on February 7, 2002, page 5834 and allowed 60 days for public comment. No public comments were 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.

Proposed Collection

Title: Prostate, Lung, Colorectal and Ovarian Cancer Screening Trial. **Type of Information Collection Request:** Revision, OMB control number 0925-0407, expiration date October 31, 2002. **Need and Use of Information Collection:** This trial is designed to determine if screening for prostate, lung, colorectal and ovarian cancer can reduce mortality from these cancers which currently cause an estimated 251,000 deaths annually in the U.S. The design is a two-armed randomized trial of men and women aged 55 to 74 at entry. The total sample size after more than 8 years of recruitment is 154,956. The primary endpoint of the trial is cancer-specific mortality for each of the four cancer sites (prostate, lung, colorectal, and ovary). In addition, cancer incidence, stage shift, and case survival are to be monitored to help understand and explain results. Biologic prognostic characteristics of the cancers will be measured and correlated with mortality to determine the mortality predictive value of these intermediate endpoints.

Basic demographic data, risk factor data for the four cancer sites and screening history data, as collected from all subjects at baseline, will be used to assure comparability between the screening and control groups and make appropriate adjustments in analysis. Further, demographic and risk factor information will be used to analyze the differential effectiveness of screening in high versus low risk individuals. **Frequency of Response:** On occasion. **Affected Public:** Individuals or households. **Type of Respondents:** Adult men and women. The annual reporting burden is as follows: **Estimated Number of Respondents:** 150,598; **Estimated Number of Responses per Respondent:** 1.38; **Average Burden Hours Per Response:** 0.19; and **Estimated Total Annual Burden Hours Requested:** 39,597. The annualized cost to respondents is estimated at: \$395,970. There are no Capital Costs to report. There are no Operating or Maintenance Costs to report.

Request for Comments

Written comments and/or suggestions from the public and affected agencies should address one or more of the following points: (1) Whether the proposed collection of information is necessary for the proper performance of the function of the agency, including whether the information will have practical utility; (2) The accuracy of the agency'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 those who are to respond, including the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology.

Direct Comments to OMB

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: Office of Management and Budget, Office of Regulatory Affairs, New Executive Office Building, Room 10235, Washington, DC 20503, Attention: Desk

Officer for NIH. To request more information on the proposed project or to obtain a copy of the data collection plans and instruments, contact: Dr. John Gohagan, Chief, Early Detection Research Group, National Cancer Institute, NIH, EPN Building, Suite 3100, 6130 Executive Boulevard, MSC7346, Bethesda, MD 20892-7346-, or call non-toll-free number (301) 496-3982 or E-mail your request, including your address to: JG72P@NIH.GOV.

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

Dated: May 21, 2002.

Reesa Nichols,

NCI Project Clearance Liaison.

[FR Doc. 02-13276 Filed 5-24-02; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Submission for OMB Review; Comment Request; Health Information National Trends Survey (HINTS)

SUMMARY: Under the provisions of Section 3507(a)(1)(D) of the Paperwork Reduction Act of 1995, the National Cancer Institute (NCI), the National Institutes of Health (NIH) has submitted to the Office of Management and Budget (OMB) a request for review and approval of the information collection listed below. This proposed information collection was previously published in

the **Federal Register** on December 21, 2000, pages 80444-80445, and allowed 60-days for public comment. No public comments were 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.

Proposed Collection

Title: Health Information National Trends Survey (HINTS). **Type of Information Collection Request:** NEW. **Need and Use of Information Collection:** As a result of the ongoing changes in the communication environment, there is an unprecedented opportunity to communicate information about cancer and other health topics to the general public. Developing appropriate messages for the public about cancer prevention, detection, diagnosis, treatment, and survivorship requires an understanding of individuals' sources and access to cancer-related information, their knowledge about cancer and other health information, or the factors that enhance or may hinder access, use, or knowledge of health information.

The HINTS is a new telephone survey designed to provide nationally representative, population-based standardized data on health knowledge and health information for the United States. The survey will establish important baseline data about cancer communication practices, preferences

for information, and cancer knowledge across the country. This survey will provide data on the public's perceived needs for cancer information, sources and access to health information (e.g., health care providers, Internet, mass media), current knowledge and understanding about cancer prevention and detection, and the barriers to more effectively understanding and utilizing cancer-related information. The HINTS is intended to be conducted every 2 years, and data will be used to measure progress in improving cancer knowledge and communication among the general public. The survey will be administered to one sample adult in 8,000 households, and is intended to have an adequate sample size to produce stable estimates for racial and ethnic minority populations.

Data from this survey are essential for NCI to develop improved cancer-related messages and materials and to tailor these messages for different audiences, especially for cancer prevention and detection. Data will be used to help selecting the best means of communicating cancer-related messages to different audiences (communication channels) to reach the diverse audiences in the United States. Finally, information obtained in this survey data will be used to identify research gaps and to guide the direction and decisions about NCI's research efforts in health promotion and health communication.

Frequency of response: One-time.

Affected public: Individuals or households. **Type of Respondents:** U.S. adults. The annual reporting burden is as follows:

Type of respondent	Estimated number of respondents	Frequency of response	Average hours per response	Annual hour burden
Adult respondents to Dress Rehearsal Interview	150	1	.4509	68
Adult respondents to Main Study Household Screener	11,200	1	.0334	374
Adult respondents to Main Study Sample Person Interview	8,000	1	.4509	3607
Total				4049

There are no Capital Costs to report. There are no Operating or Maintenance Costs to report.

Request for Comments

Written comments and/or suggestions from the public and affected agencies are invited on one or more of the following points: (1) Whether the proposed collection of information is necessary for the proposed performance of the functions of the agency, including whether the information shall have practical utility; (2) The accuracy of the

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 those who are to respond, including the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology.

Direct Comments to OMB

Written comments and/or suggestions regarding the item(s) contained in the notice, especially regarding the estimated public burden and associated response time, should be directed to the: Office of Management and Budget, Office of Regulatory Affairs, New Executive Office Building, Room 10235, Washington, DC 20503, Attention: Desk Officer for NIH. To request more information on the proposed project or to obtain a copy of the data collection plans and instruments, contact David E.