DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Proposed Collection; Comment Request; Request; OMB No. 0925–0177 "Special Volunteer and Guest Researcher Assignment," Form 590

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 information collection, the Office of Human Resources, 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: Special Volunteer and Guest Researcher Assignment for use in NIH facilities.

Type of Information Collection Request: Reinstatement, OMB 0925–0177, Expiration Date July 31, 2005.

Need and Use of Information Collection Request:

Form Number: NIH–590. A single Form NIH–590 is completed by an NIH official for each Guest Researcher or Special Volunteer prior to his/her arrival at NIH. The information on the form is necessary for the approving official to reach a decision on whether to allow a Guest Researcher to use NIH facilities, or whether to accept volunteer services offered by a Special Volunteer. If the original assignment is extended, another form notating the extension is completed to update the file.

Frequency of Response: Once.

Affected Public: Individuals.

Type of Respondents: Non-federal scientific professionals and/or individuals.

The annual Reporting burden is as follows:

Estimated Number of Respondents: 1660;

Estimated Number of Responses per Respondent: 1.0;

Average Burden Hours per Response: 0.1; and

Estimated Total Annual Burden Hours Requested: 166. The estimated annualized cost to respondents is $2,275.

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.

FOR FURTHER INFORMATION CONTACT: To request more information on the proposed project or to obtain a copy of the data collection plans and instruments, contact Mrs. Wanda Darwin, Office of Human Resources, Office of The Director, NIH, Building 31, Room 1C31E, One Center Drive, Bethesda, MD 20892–2269, or call non-toll-free number 301–402–2820, or E-mail your request, including your address to: [*darwinw@od.nih.gov*].

Comments Due Date: Comments regarding this information collection are best assured of having their full effect if received within 60 days of the date of this publication.

Dated: August 18, 2010.

Wanda R. Darwin,

Human Resources Specialist, Office of Human Resources, National Institutes of Health.

[FR Doc. 2010–21099 Filed 8–24–10; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Submission for OMB Review; Comment Request; Resource for the Collection and Evaluation of Human Tissues and Cells From Donors With an Epidemiology Profile (NCI)

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 to review and approve the information collection listed below. This proposed information collection was previously published in the **Federal Register** on June 16, 2010 (75 FR 34146) and allowed 60-days for public comment. One public comment was received on 7/16/2010 from a business informing us that they are able to provide a time-saving "batch processing service" to locate and verify "the most current addresses and phone numbers" of survey respondents. A response was sent on 7/26/2010 to the business which indicated the existence of similar devices and/or procedures in the current design of the project. 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: Resource for the Collection and Evaluation of Human Tissues and Cells from Donors with an Epidemiology Profile (NCI). Type of *Information Collection Request: New:* Need and Use of Information Collection: Under the auspices of three NCI IRB-approved protocols and instruments, the Laboratory of Human Carcinogenesis conducts case-control studies to investigate the relations between biomarkers, the environment, and human cancer. Human subjects recruited from the general population are needed as controls (Population Controls) for bio-specimens and personal histories (social, occupational and health) that serve as references for the significance of the frequency and prevalence of bio-markers found in cancer patients and thought to be important in the development, progression, and/or response to treatment of the malignant growths in cancer patients. The questionnaires will be used to obtain the personal histories to compare to the life styles and exposures and the biospecimens will serve as controls for the assay results obtained from cancer patients. The collection of information and specimens from the cancer cases received NIH Clinical Exemption (Request #2009–09–002) on October 28, 2009. *Frequency of Response:* Once. *Affected Public:* Adult and senior members of the licensed driver population in Baltimore, Maryland and eleven near-by counties, including the Eastern Shore. *Type of Respondents:* Responders will be English speaking, male and female, Caucasian, African-American and Asian. The total annual reporting burden is estimated to be 692 (see table below). There are no Capital Costs, Operating Costs, and/or Maintenance Costs to report.

TABLE 1—ESTIMATES OF ANNUAL BURDEN HOURS

Type of respondents	Survey instrument	Number of respondents	Frequency of response	Average time per response (minutes/hour)	Annual burden hours
Adults (40–79 years old)	Telephone Screener (Attachment 16)	1700	1	10/60 (0.17)	283
	Main Questionnaire (Attachment 6)	225	1	60/60 (1)	225
	Prostate Supplemental Questionnaire (Attachment 7).	125	1	30/60 (0.5)	63
	Liver Supplement (Attachment 8)	225	1	30/60 (0.5)	113
	Refusal Questionnaire Form (Attachment 21)	225	1	2/60 (0.03)	8
Totals		2500			692

Request for Comments: Written comments and/or suggestions from the public and affected agencies should address one or more of the following points: (1) Evaluate 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) Evaluate 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) Enhance the quality, utility, and clarity of the information to be collected; and (4) 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 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 Glenwood E. Trivers or Elise Bowman, Center for Cancer Research, NCI, NIH, 37 Convent Drive Room 3060–C or 3060–A, Building 37, Bethesda, Maryland 30893–4258 or call non-toll-free number 301–496–2094 or 301–496–2090 or e-mail your request, including your address to *triversg@mail.nih.gov* or *bowmane@mail.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: August 18, 2010.

Vivian Horovitch-Kelley

NCI Project Clearance Liaison, National Institutes of Health.

[FR Doc. 2010–21098 Filed 8–24–10; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2010–N–0079]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Experimental Study of Graphic Cigarette Warning Labels

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 September 24, 2010.

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–NEW and title "Experimental Study of Graphic Cigarette Warning Labels." Also include the FDA docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Jonna Capezzuto, Office of Information Management, Food and Drug Administration, 1350 Piccard Dr., PI50–400B, Rockville, MD 20850, 301–796–3794, *Jonnalynn.capezzuto@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.

Experimental Study of Graphic Cigarette Warning Labels—(OMB Control Number 0910–NEW)

Tobacco products are responsible for more than 440,000 deaths each year. The Centers for Disease Control and Prevention report that approximately 46 million U.S. adults smoke cigarettes in the United States, even though this behavior will result in death or disability for half of all regular users. Paralleling this enormous health burden is the economic burden of tobacco use, which is estimated to total $193 billion annually in medical expenditures and lost productivity. Curbing the significant adverse consequences of tobacco use is one of the most important public health goals of our time. One way to do this is through health warnings that describe and graphically depict the harm caused by cigarette use.

On June 22, 2009, the President signed the Family Smoking Prevention and Tobacco Control Act (Tobacco Control Act) (Public Law 111–31) into law. The Tobacco Control Act granted FDA important new authority to regulate the manufacture, marketing, and distribution of tobacco products to protect the public health generally and to reduce tobacco use by minors. Section 201 of the Tobacco Control Act, which amends section 4 of the Federal Cigarette Labeling and Advertising Act (15 U.S.C. 1333), requires FDA to issue "regulations that require color graphics depicting the negative health consequences of smoking to accompany the label statements specified in subsection (a)(1)." FDA conducts research relating to tobacco products under its statutory authority in section