exceeding $5,000 total direct costs only for the first year. An additional 4 years of support, up to $5,000 (direct costs only) each year may be available, depending upon fiscal year appropriations and successful performance of the conference.

D. Funding Plan

Continued funding for future year, noncompetitive segments, will be contingent upon satisfactory progress as determined annually by the FDA, the receipt of a PHS 2590 application, the approval of yearly task force reports, and the availability of Federal funds. An estimated amount of $160,000 is available in FY 2008. The number of grants funded will depend on the quality of the applications received, their relevance to FDA's mission, priorities, and the availability of funds.

VI. Review Procedure and Criteria

All applications submitted in response to this request for applications (RFA) will first be reviewed for responsiveness by grants management and program staff. Responsiveness is defined as submission of a complete on or before the required submission date as listed in the previous paragraphs. If applications are found to be nonresponsive, they will be returned to the applicant without further consideration.

Responsive applications will be reviewed and evaluated for scientific and technical merit by an ad hoc panel of experts. The following will be considered in making funding decisions: (1) Scientific merit of the proposed conference/scientific meeting as determined by the evaluation process; (2) availability of funds; and (3) relevance of program priorities. Final funding decisions will be made by the Commissioner of Food and Drugs or his or her designee.

Applicants are strongly encouraged to contact FDA Program staff to resolve any questions regarding criteria before the submission of their application.

VII. Submission Requirements

FDA is accepting new applications for this program electronically via *www.grants.gov*. To download the SF424 (5161) Application forms for this Funding Opportunity Announcement (FOA) link to *http://www.grants.gov/Apply* and follow the directions provided on that site. A one-time registration is required for institutions at: Grants.gov (*http://www.grants.gov/GetStarted*). The application receipt date is July 15, 2008.

Your organization will need to obtain a Data Universal Number System (DUNS) number as part of the Grants.gov registration process. The DUNS number is a 9-digit identification number, which uniquely identifies business entities. Obtaining a DUNS number is easy and there is no charge. The D&B number can be obtained by calling 866–705–5711 or through the Web site at *http://www.dnb.com/us.*

The applicant must also register in the Central Contractor Registration (CCR) database in order to be able to submit the application. Information about the CCR is available at *http://www.grants.gov/applicants/get_registered.jsp.*

VIII. Method of Application

A. Submission Instructions

The SF424 (5161) application has several components. Some components are required, others are optional. The forms package associated with this FOA in Grants.gov/APPLY includes all applicable components, required, and optional.

B. Format for Application

A completed application in response to this FOA includes the data in the following components:

The face page of the application should indicate ''Response to Food Protection Task Force Conference Grant Program RFA FD 08–006''.

Applications should include the following: (1) A title which has the term ''state food protection task force meetings'', ''conference'', ''council'', ''workshop'', ''alliance'' or other similar description to assist in the identification of the request; (2) location of the conference; (3) expected number of registrants and type of audience expected with their credentials; (4) dates of conference(s); (5) conference format and projected agenda(s), including list of principal areas or topics to be addressed; (6) physical facilities required for the conduct of the meeting; (7) justification of the conference(s), including the problems it intends to clarify and any developments it may stimulate; (8) brief biographical sketches of individuals responsible for planning the conference(s) and details concerning adequate support staff; (9) information about all related conferences held on this subject during the last 3 years (if available); (10) details of proposed per diem/subsistence rates, transportation, printing, supplies and facility rental costs; and (11) the necessary checklist and assurances pages provided in each application package.

IX. Freedom of Information

Data included in the application which have been specifically identified by the applicant as containing restricted and/or proprietary information may be entitled to confidential treatment as trade secret or confidential commercial information within the meaning of the Freedom of Information Act, (5 U.S.C. 552(b)(4), and FDA's implementing regulations (21 CFR 20.61).

Dated: June 3, 2008.

Jeffrey Shuren,

Associate Commissioner for Policy and Planning.

[FR Doc. E8–13015 Filed 6–9–08; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Proposed Collection; Comment Request; NIH-American Association for Retired Persons (AARP) Comprehensive Lifestyle Interview by Computer (CLIC) Study (NCI)

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 Cancer Institute (NCI), 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: NIH-American Association for Retired Persons (AARP) Comprehensive Lifestyle Interview by Computer (CLIC) Study. *Type of Information Collection Request:* New. *Need and Use of Information Collection:* The Nutritional Epidemiology Branch of the Division of Cancer Epidemiology and Genetics of the National Cancer Institute has planned this study to evaluate the feasibility of using these three new computerized questionnaires as well as the Diet and Health Questionnaire (DHQ), a well-established food frequency questionnaire in a population of early-to-late-middle-aged men and women. Participants will be asked to complete one of four different series (pathways) of computerized questionnaires over a 90 day period, with some questionnaires in a series being completed twice. This evaluation study comprises the necessary performance and feasibility tests for the new computerized questionnaires, which will provide an opportunity to assess the possibility of administering

computerized questionnaires in future large prospective cohort studies. The computerized questionnaires will support the ongoing examination between cancer and other health outcomes with nutritional, physical activity, and lifestyle exposures. The computerized questionnaires adhere to The Public Health Service Act, Section 412 (42 U.S.C. 285a–1) and Section 413 (42 U.S.C. 285a–2), which authorizes the Division of Cancer Epidemiology and Genetics of the National Cancer Institute (NCI) to establish and support programs for the detection, diagnosis, prevention and treatment of cancer; and to collect, identify, analyze and disseminate information on cancer research, diagnosis, prevention and treatment. *Frequency of Response:* Either 2 or 4 times, depending on the pathway. *Affected Public:* Individuals. *Type of Respondents:* U.S. adults (aged 50 and over). The annual reporting burden is displayed in the table below. The estimated total annualized burden hours being requested is 2616. The annualized cost to respondents is estimated at: $46,242. There are no Capital Costs, Operating Costs, and/or Maintenance Costs to report.

TABLE 1.—ESTIMATES OF ANNUAL BURDEN HOURS

Type of respondents	Instrument(s) tested	Frequency of response	Average time per response (minutes/hour)	Number of respondents/pathway	Annual hour burden
Senior Adults	Read Invitation	1	1/60	7500	125.000
	Pre-Enrollment	1	10/60	1046	174.333
	Enrollment Process	1	5/60	1035	86.250
	Assigned Pathway 1				
	ACT–24	2	15/60	156	78.000
	LHQ	1	20/60	156	52.000
	DHQ	1	30/60	156	78.000
	1 Web Re-entry	1	1/60	156	2.600
	Assigned Pathway 2				
	ASA24	2	30/60	156	156.000
	DHQ	1	30/60	156	78.000
	LHQ	1	20/60	156	52.000
	1 Web Re-Entry	1	1/60	156	2.600
	Assigned Pathway 3				
	ACT–24	2	15/60	362	181.000
	ASA24	2	30/60	362	362.000
	LHQ	1	20/60	362	120.667
	DHQ	1	30/60	362	181.000
	1 Web Re-Entry*	1	1/60	362	6.033
	Assigned Pathway 4				
	ACT–24	2	15/60	362	181.000
	ASA24	2	30/60	362	362.000
	LHQ	1	20/60	362	120.667
	DHQ	1	30/60	362	181.000
	3 Web Re-entries**	3	1/60	362	18.100
	Evaluation Survey	1	1/60	1035	17.250
Totals					2615.50

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) 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.

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 Arthur Schatzkin, M.D., Dr.P.H, Chief, Nutritional Epidemiology Branch, Division of Cancer Epidemiology and Genetics, National Cancer Institute, NIH, DHHS, Executive Plaza South, Room 3040, 6120 Executive Blvd., EPS–MSC 7242, Bethesda, MD 20892–7335 or call non-toll-free number 301–594–2931 or e-mail your request, including your address to: *schatzka@mail.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: June 2, 2008.

Vivian Horovitch-Kelley,

NCI Project Clearance Liaison Office, National Institutes of Health.

[FR Doc. E8–12920 Filed 6–9–08; 8:45 am]

BILLING CODE 4140–01–P