Women respondents	Number of respondents	Number of responses per respondents	Average burden per response (in hours)	Total burden (in hours)
Quantitative Survey in site A and site B	200	1	1	200
Total	250			225

Dated: July 30, 2001.

Nancy Cheal,

Acting Associate Director for Policy, Planning and Evaluation, Centers for Disease Control and Prevention.

[FR Doc. 01–19656 Filed 8–6–01; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[Program Announcement 01196]

Evaluation of Breast Cancer Incidence; Notice of Availability of Funds

A. Purpose

The Centers for Disease Control and Prevention (CDC) announces the availability of fiscal year (FY) 2001 funds for a grant program for an Evaluation of Breast Cancer Incidence in DuPage County, Illinois. This program addresses the ''Healthy People 2010'' focus areas of Cancer and Environmental Health.

The purpose of the program is to conduct an analysis of data routinely collected by health service organizations on breast cancer morbidity and mortality in DuPage County, Illinois. Through this program, the DuPage County Health Department will be able to determine the incidence of breast cancer in the county and to outline a plan to address the programmatic and health issues identified.

No human subjects research will be supported under this program announcement.

B. Eligible Applicant

Assistance will be provided only to the DuPage County Health Department in Wheaton, Illinois. No other applications are solicited. Eligibility is limited to the DuPage County Health Department because Fiscal Year 2001 federal appropriations specifically direct the Centers for Disease Control and Prevention to award funds to evaluate the high incidence of breast cancer in DuPage County, Illinois.

Note: Title 2 of the United States Code, Chapter 26, Section 1611 states that an organization described in section 501(c)(4) of the Internal Revenue Code of 1986 that engages in lobbying activities is not eligible to receive Federal funds constituting an award, grant, cooperative agreement, contract, loan, or any other form.

C. Availability of Funds

Approximately $92,000 is available in FY 2001 to fund this award. It is expected that the award will begin on or about September 30, 2001, and will be made for a one year project period. Funding estimates may change.

D. Where To Obtain Additional Information

This and other CDC announcements can be found on the CDC home page Internet address—http://www.cdc.gov Click on ''Funding'' then ''Grants and Cooperative Agreements.''

To obtain business management technical assistance, contact: Sharron Orum, Grants Management Specialist, Grants Management Branch, Procurement and Grants Office, Centers for Disease Control and Prevention, 2920 Brandywine Road, Room 3000, Atlanta, GA 30341–4146, Telephone number: (770) 488–2716, Email address: spo2@cdc.gov

For program technical assistance, contact: Ronney Lindsey, Deputy Director, Division of Environmental Hazards and Health Effects, National Center for Environmental Health, Centers for Disease Control and Prevention, 4770 Buford Highway, Mailstop E19, Atlanta, GA 30341, Telephone number: (404) 498–1308, Email address: rll3@cdc.gov

Dated: August 1, 2001.

John L. Williams,

Director, Procurement and Grants Office Centers for Disease Control and Prevention (CDC).

[FR Doc. 01–19693 Filed 8–6–01; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 01N–0319]

Agency Information Collection Activities; Proposed Collection; Comment Request; Health and Diet Survey

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing an opportunity for public comment on the proposed collection of certain information by the agency. Under the Paperwork Reduction Act of 1995 (the PRA), Federal agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information and to allow 60 days for public comment in response to the notice. This notice solicits comments on a voluntary consumer survey about knowledge, perceptions, attitudes, and practices related to dietary supplements and food.

DATES: Submit written or electronic comments on the collection of information by October 9, 2001.

ADDRESSES: Submit electronic comments on the collection of information to http://www.accessdata.fda.gov/scripts/oc/dockets/edockethome.cfm. Submit written comments on the collection of information to the Dockets Management Branch (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. All comments should be identified with the docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Peggy Schlosburg, Office of Information Resources Management (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–1223.

SUPPLEMENTARY INFORMATION: Under the PRA (44 U.S.C. 3501–3520), Federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor.

''Collection of information'' is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA (44 U.S.C. 3506(c)(2)(A)) requires Federal agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information before submitting the collection to OMB for approval. To comply with this requirement, FDA is publishing notice of the proposed collection of information set forth in this document.

With respect to the following collection of information, FDA invites comments on: (1) Whether the proposed collection of information is necessary for the proper performance of FDA's functions, including whether the information will have practical utility; (2) the accuracy of FDA'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 respondents, including through the use of automated collection techniques, when appropriate, and other forms of information technology.

Health and Diet Survey

The authority for FDA to collect the information derives from the authority of the Commissioner of Food and Drugs, as specified in section 903(d)(2) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 393(d)(2)). The Health and Diet Survey will provide FDA information about consumers' knowledge, perceptions, attitudes, and practices related to dietary supplements and food. A nationally representative sample of 2,000 adults in the 48 contiguous States and the District of Columbia will be selected at random and interviewed by telephone. Participation will be voluntary. The survey will collect information about: (1) Prevalence, experience, and purposes of use of dietary supplements; (2) knowledge of health benefits, health risks, and regulation of dietary supplements; (3) sources of dietary supplement information; (4) perceptions of dietary supplement labels; (5) replacement and combination use of supplements and drugs; (6) adverse experience with dietary supplements; (7) children's and teenagers' use of dietary supplements; (8) knowledge of diet-health relationships; (9) dietary management practices; and (10) use of food labels.

Some of the questions to be asked (items 8 through 10 listed in the previous paragraph) replicate the ones asked in the 1995 Health and Diet Survey. Responses to these questions will help FDA identify and measure any changes in consumer knowledge, perceptions, attitudes, and practices with regard to diet, health, and use of food labels. The information will also help the agency evaluate the effectiveness of the Nutrition Labeling and Education Act of 1990 in promoting the public health.

The agency will use the other questions in the proposed survey to enhance its understanding of consumer knowledge, perceptions, attitudes, and practices regarding dietary supplements. Subsequent to the enactment of the Dietary Supplement Health and Education Act of 1994, the consumption of dietary supplements in the United States has been increasing. FDA needs current, timely, and policy-relevant consumer information to help it identify needs for and develop consumer education programs and regulatory policies to ensure safe and appropriately labeled supplement products. The survey will help the agency measure prevalence and distribution of consumer knowledge, perceptions, attitudes, and practices. This information can be used to understand and describe the consumer environment that is the intended target of labeling and education initiatives.

FDA estimates the burden of this collection of information as follows:

TABLE1.— ESTIMATED ANNUAL REPORTING BURDEN[1]

Activity	No. of Respondents	Annual Frequency per Response	Total Annual Responses	Hours per Response	Total Hours
Cognitive interview	9	1	9	1.5	13.5
Pretest	9	1	9	0.5	4.5
Screener	4,200	1	4,200	0.02	84
Survey	2,000	1	2,000	0.5	1,000
Total					1,102

[1] There are no capital costs or operating and maintenance costs associated with this collection of information.

These estimates are based on FDA's experience with previous consumer surveys. Prior to the administration of the survey, the agency plans to conduct a series of nine cognitive interviews and a series of nine pretests to ensure the quality of the survey. Cognitive interviews will help the agency understand respondent comprehension of the meanings of questions and words, and how respondents answer questions. Pretests will help the agency examine and reduce problems in the administration of the final questionnaire. The agency will use a screener to select an eligible adult respondent in each household to participate in the survey.

Dated: August 1, 2001.

Margaret M. Dotzel,

Associate Commissioner for Policy.

[FR Doc. 01–19626 Filed 8–6–01; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Oncologic Drugs Advisory Committee; Notice of Meeting

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

This notice announces a forthcoming meeting of a public advisory committee of the Food and Drug Administration (FDA). The meeting will be open to the public.