

Affected public: Individuals and households; businesses and other for profit, small businesses and organizations. **Types of respondents:** These surveys are designed to assess the satisfaction of the Clinical Center's major internal and external customers with the services provided. These

customers include, but are not limited to, the following groups of individuals: Clinical Center patients, family members of Clinical Center patients, visitors to the Clinical Center, National Institutes of Health investigators, NIH intramural collaborators, private physicians or organizations who refer

patients to the Clinical Center, volunteers, vendors and collaborating commercial enterprises, small businesses, regulators, and other organizations. The annual reporting burden is as follows:

TABLE 1.—BURDEN ESTIMATE

Customer	Type of survey	Estimated number to be surveyed	Expected response rate	Time to complete survey (minutes)	Estimated burden hours
Clinical Center Patients	Questionnaire/Telephone	11,100	66%	20	2436.6
Family Members of Patients	Questionnaire/Post-Card	8500	38%	10	533.3
Visitors to the Clinical Center	Questionnaire/Post-Card	3500	15%	10	87.5
Former physician employees and trainees.	Electronic	650	35%	10	38.2
Guest workers/Guest researchers	Electronic	950	60%	22	210
Extramural collaborators	Electronic	600	30%	15	45
Vendors and Collaborating Commercial Enterprises.	Questionnaire/Fax-Back	9500	17%	18	475
Professionals and Organizations Referring Patients.	Fax Back	9000	30%	28	1250
Regulators	Fax Back	85	82%	19	22
Volunteers	Questionnaire	850	58%	28	230
Total	n=16,812	5,327.6

Estimated costs to the respondents consists of their time; time is estimated using a rate of \$10.00 per hour for patients and the public; \$30.00 for vendors, regulators, organizations and \$55.00 for health care professionals. The estimated annual costs to respondents for each year for which the generic clearance is requested is \$72,894 for 1998, \$30,276 for 1999, and \$24,531 for 2000. There are no capital costs, operating costs and/or maintenance costs to report.

REQUESTS 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 functions of the Clinical Center and the agency, including whether the information shall 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 automated, electronic, mechanical, or other technological collection techniques or other forms of information technology.

FOR FURTHER INFORMATION: To request more information on the proposed

project, to obtain a copy of the data collection plans and instruments, or to submit comments, contact: Dr. David K. Henderson, Deputy Director for Clinical Care, Warren G. Magnuson Clinical Center, National Institutes of Health, Building 10, Room 2C 146, 9000 Rockville Pike, Bethesda, Maryland 20892, or call non-toll free: (301) 496-3515, or e-mail your request or comments, including your address to dhenderson@cc.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: April 17, 1997.

Michael Goldrich,

Deputy Director, CC.

[FR Doc. 97-10550 Filed 4-23-97; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Submission for OMB Review; Comment Request; Validation of a New Food Frequency Questionnaire

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 Institutes of Health (NIH), National Cancer Institute (NCI) will publish periodic summaries of proposed projects to be submitted to the Office of Management and Budget (OMB) for review and approval. This proposed information collection was previously published in the **Federal Register** on December 24, 1996, page 67839 and allowed 60-days for public comment. There were only two requests for additional information and no comments. 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:* Validation of a New Food Frequency Questionnaire. *Type of Information Collection Request:* New. *Need and use of Information Collection:* The agency conducts and funds studies examining the relationship between diet and chronic diseases. This information collection is needed to validate and further refine a new diet history questionnaire to be used in studies of diet and disease. The new questionnaire will be validated against reference data from four non-consecutive 24-hour dietary recalls among a national sample of persons 20-70 years of age. The

validity of the new questionnaire will be compared to two widely used food frequency questionnaires. As a further validation, biological nutrition measures

from blood specimens will be obtained from a 20% sub-sample of participants. *Frequency of response:* One-time study. *Affected public:* Individuals or

households. *Types of Respondents:* US adults 20–70 years of age. The annual reporting burden is as follows:

Data collection form	Estimated number of respondents	Estimated number of responses per respondent	Avg. burden hours per response	Estimated total hour burden	Estimated total annual burden hours requested
Screener	2700	1	0.167	450.9	300.6
Recalls interview #1	1620	1	0.75	1215.0	810.0
Recall interview #2	1563	1	0.5	781.5	521.0
Recall interview #3	1507	1	0.5	753.5	502.3
Recall interview #4	1451	1	0.5	725.5	483.7
New Questionnaire	1225	1	0.75	918.8	612.5
Food Questionnaire 1	612	1	0.5	306.0	204.0
Food Questionnaire 2	612	1	0.668	408.8	272.5
Opinion form	1225	1	0.167	204.6	136.4
Blood substudy	240	2	0.25	120.0	80.0
Total	2700	5884.6	3923.0

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 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, D.C. 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 Amy F. Subar, Ph.D., Project Officer, National Cancer Institute, EPN 313, 6130 EXECUTIVE BLVD MSC 7344, BETHESDA MD 20892–7344, or call non-toll-free number (301) 496–8500, or FAX your request to (301) 435–3710, or E-mail your request, including your address, to amy_subar@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: April 9, 1997.

Nancie L. Bliss,

OMB Project Clearance Liaison.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Substance Abuse and Mental Health Services Administration

Agency Information Collection Activities Under OMB Review

Periodically, the Substance Abuse and Mental Health Services Administration

(SAMHSA) will publish a list of information collection requests under OMB review, in compliance with the Paperwork Reduction Act (44 U.S.C. Chapter 35). To request a copy of these documents, call the SAMHSA Reports Clearance Officer on (301)443–8005.

Access to Community Care and Effective Services and Supports (ACCESS) Evaluation Study; Revision

The Center for Mental Health Services (CMHS) will continue an evaluation study that is assessing service systems integration (SI) approaches for homeless persons with serious mental illnesses. The evaluation study will collect data through interviews with homeless persons with serious mental illness and providers of services to homeless persons. SI sites will be contrasted with comparison sites to assess the impact of SI. The evaluation will describe approaches to SI, processes by which SI takes place, factors that influence SI, and the impact that SI has on homeless persons with serious mental illness. The estimated annualized burden is shown below.

	Number of respondents (5 years)	No. of responses per respondent	Average burden per response	Total burden hours (5 years)	Total annualized burden hours
Clients (Homeless Persons)	7,200	2.65	.98	18,702	3,740
Service Providers	1,426	78	.11	12,147	2,429