

training provided to Ryan White CARE Act grantees, resource allocation, and capacity expansion.

Trainees will be asked to complete the Participant Information Form for each activity they complete. The estimated

annual response burden to attendees of training programs is as follows:

Form	Number of respondents	Responses per respondent	Total responses	Hours per response	Total burden hours
Participant Information	75,000	2	150,000	0.2	30,000

The estimated annual burden to AETCs is as follows:

Form	Number of respondents	Responses per respondent	Total responses	Hours per response	Total burden hours
Program Record	12	500	6,000	0.1	600
Clinical Consultation	12	300	3,600	0.1	360
Group Clinical Consultation	12	75	900	0.1	90
Technical Assistance	12	250	3,000	0.1	300
Aggregate Data Set	12	3	36	32	1,152
Total	12	13,536	2,502

The total burden hours being requested are 32,502.

Written comments and recommendations concerning the proposed information collection should be sent within 30 days of this notice to: John Morrall, Human Resources and Housing Branch, Office of Management and Budget, New Executive Office Building, Room 10235, Washington, DC 20503.

Dated: September 30, 2003.

Jane M. Harrison,

Director, Division of Policy Review and Coordination.

[FR Doc. 03-25250 Filed 10-3-03; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Proposed Collection; Comment Request; National Cancer Institute Science Enrichment Program Surveys

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: NCI Science Enrichment Program (SEP) Surveys.

Type of Information Collection

Request: New.

Need and Use of Information

Collection: NCI SEP is a 5-week summer residential program on university campuses that serves under-represented minority and under-served students who have just completed ninth grade.

The program goals are to: (1) Encourage student participants to select careers in science, mathematics, and/or research, and (2) broaden and enrich students' science, research, and sociocultural backgrounds. The proposed data collection encompasses three surveys: (1) A follow-up survey of

SEP and control group students who participated in a five-year longitudinal evaluation of the program conducted between 1998 and 2003; (2) a post-program survey of parents of SEP 2004 participants; and (3) a follow-up survey of SEP 1990-1997 alumni. The information from the proposed data collection will supplement previous evaluation results, which have been and will continue to be used to judge program process and outcomes.

Frequency of Response: One time.

Affected Public: Individuals or households.

Type of Respondents: High school and college students, young adults, and parents of high school students participating in the program.

Cost to Respondents: \$4,070.

The annual reporting burden is as follows:

ESTIMATES OF HOUR BURDEN: BURDEN REQUESTED

Type of respondents	Average number of respondents/Yr.	Frequency of response	Average time per response	Average annual hour burden
SEP Participants	600	1	0.25	150
Control Group Students	300	1	0.25	75
Parents of SEP Participants	100	1	0.25	25
SEP 1990-1997 Alumni	627	1	0.25	157
Total	1,627	407

There are no Capitol Costs, Operating Costs, and/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 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 Mr. Frank Jackson, Center to Reduce Cancer Health Disparities, National Cancer Institute, National Institutes of Health, 6116 Executive Boulevard, Suite 602, Rockville, MD 20852, or call non-toll-free number (301) 496-8589, or E-mail your request, including your address to: fj12i@nih.gov.

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

Dated: September 29, 2003.
Reesa Nichols,
NCI Project Clearance Liaison.
 [FR Doc. 03-25294 Filed 10-6-03; 8:45 am]
BILLING CODE 4140-01-M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Submission for OMB Review; Comment Request: National Institute of Diabetes and Digestive and Kidney Diseases Information Clearinghouses Customer Satisfaction Survey

SUMMARY: Under the provision of section 3507(a)(1)(D) of the Paperwork Reduction Act of 1995, the National Institute of Diabetes and Digestive and Kidney Diseases (NIDDK), 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 April 30, 2003, pages 23150-23151 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: NIDDK Information Clearinghouses Customer Satisfaction Survey.

Type of Information Request: Extension. The OMB control number 0925-0480 expires July 31, 2003.

Need and Use of Information Collection: NIDDK is conducting a survey to evaluate the efficiency and effectiveness of services provided its three information clearinghouses: National Diabetes Information Clearinghouse, National Digestive Diseases Information Clearinghouse, National Kidney and Urologic Diseases Information Clearinghouse. The survey responds to Executive Order 12862, "Setting Customer Service Standards," which requires agencies and departments to identify and "survey their customers to determine the kind and quality of service they want and their level of satisfaction with existing service."

Frequency of Response: On occasion.
Affected Public: Individuals or households; clinics or doctor's offices.

Type of Respondents: Physicians, nurses, patients, family.

The annual reporting burden is as follows:

Estimated Number of Respondents: 12,000.

Estimated Number of Responses per Respondent: 1.

Estimated Average Burden Hours Per Response: 0.1671.

Estimated Total Annual Burden Hours Requested: 2,000. The annualized cost to respondents is estimated at \$39,000. There are no Capital Costs to report. There are no Operating or Maintenance Costs to report.

Type or respondents	Number of respondents	Frequency of response	Estimated average response time	Estimated annual burden hours.
Patients/Family	3,600	1.00	0.167	600
Phys. Asst.	7,200	1.00	0.167	1,200
Physicians	1,200	1.00	0.167	200
Totals	12,000	2,000

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 estimate 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