

programs. OMB reinstated the collection on August 7, 2002. This information is needed to facilitate nondiscrimination in GSA's Federal Financial Assistance Programs, consistent with Federal civil rights laws and regulations that apply to recipients of Federal financial assistance.

Public comments are particularly invited on: Whether this collection of information is necessary and whether it will have practical utility; whether our estimate of the public burden of this collection of information is accurate, and based on valid assumptions and methodology; ways to enhance the quality, utility, and clarity of the information to be collected.

DATES: *Comment Due Date:* February 24, 2003.

ADDRESSES: Submit comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden to Stephanie Morris, General Services Administration (MVA), Room 4035, 1800 F Street, NW., Washington, DC 20405.

FOR FURTHER INFORMATION CONTACT: K. Evelyn Britton, Office of Civil Rights, (202) 501-4347.

SUPPLEMENTARY INFORMATION:

A. Purpose

The General Services Administration (GSA) has mission responsibilities related to monitoring and enforcing compliance with Federal civil rights laws and regulations that apply to Federal Financial Assistance programs administered by GSA. Specifically, those laws provide that no person on the ground of race, color, national origin, disability, sex or age shall be excluded from participation in, be denied the benefits of, or be otherwise subjected to discrimination under any program in connection with which Federal financial assistance is extended under laws administered in whole or in part by GSA. These mission responsibilities generate the requirement to request and obtain certain data from recipients of Federal surplus property for the purpose of determining compliance, such as the number of individuals, based on race and ethnic origin, of the recipient's eligible and actual serviced population; race and national origin of those denied participation in the recipient's program(s); non-English languages encountered by the recipient's program(s) and how the recipient is addressing meaningful access for individuals that are Limited English Proficient; whether there has been complaints or lawsuits filed against the

recipient based on prohibited discrimination and whether there has been any findings; and whether the recipient's facilities are accessible to qualified individuals with disabilities.

B. Annual Reporting Burden

Respondents: 500.

Responses Per Respondent: 1.

Total Responses: 500.

Hours Per Response: 2.

Total Burden Hours: 1000.

Obtaining Copies of Proposals: A copy of this proposal may be obtained from the General Services Administration, Regulatory & Federal Assistance Publications Division, (MVA), Room 4035, 1800 F Street, NW., Washington, DC 20405, or by telephoning (202) 208-7312, or by faxing your request to (202) 501-4067. Please cite OMB Control No. 3090-0228, Nondiscrimination in Federal Financial Assistance Programs, in all correspondence.

Dated: November 26, 2002.

Michael Carleton,

Chief Information Officer.

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BILLING CODE 6820-34-M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day-03-24]

Proposed Data Collections Submitted for Public Comment and Recommendations

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 Centers for Disease Control and Prevention (CDC) will publish periodic summaries of proposed projects. To request more information on the proposed projects or to obtain a copy of the data collection plans and instruments, call the CDC Reports Clearance Officer on (404) 498-1210.

Comments are invited on: (a) Whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) ways to enhance the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information

on respondents, including through the use of automated collection techniques or other forms of information technology. Send comments to Anne O'Connor, CDC Assistant Reports Clearance Officer, 1600 Clifton Road, MS-D24, Atlanta, GA 30333. Written comments should be received within 60 days of this notice.

Proposed Project: Families, Communities, and Diabetes Management Project—New—National Center for Chronic Disease Prevention and Health Promotion (NCCDPHP), Centers for Disease Control and Prevention (CDC).

Diabetes Mellitus Type 2 is a chronic metabolic disease with a potential for serious health consequences that include both psychological and physical conditions. Effective management of this disease is important to prevent the development of these problems. No previous studies have systematically examined the ways in which psychological functioning, patient-provider relationships, family and social support, health insurance availability and utilization, lifestyle practices, and community support influence diabetes self-management among African American patients. Most diabetes management information is based on research conducted primarily with White patients. Accordingly, the Division of Diabetes Translation within the National Center for Chronic Disease Prevention and Health Promotion of the Centers for Disease Control and Prevention plans to conduct a longitudinal, family-centered research project to determine optimal ways of teaching African American patients and their families how to work together to manage diabetes successfully.

The research will involve samples of 40- to 64-year-old African American men and women with Type 2 diabetes and their close family members. Participating families will be divided into two groups, an intervention group that will receive the intervention at the beginning of the study, and a comparison group that will receive a modified version at the end. Measurements of self-care adherence and diabetes control will include both self-reports and objective measures such as blood glucose levels obtained through clinical lab work. Other data will include diabetes knowledge, community characteristics, social support, community support, psychological functioning, patient-provider relationships, and health care coverage. Participant involvement will occur over 13 month period. The estimated cost to respondents is \$213,236.

| respondents | No. of re-pondents | No. of re-sponses/re-pondent | Avg. burden/response (in hrs.) | Total burden (in hrs.) |
|--------------------------------------|--------------------|------------------------------|--------------------------------|------------------------|
| Adults with Diabetes: | | | | |
| Questionnaires | 400 | 3 | 60/60 | 1200 |
| Clinical Lab Work | 400 | 3 | 60/60 | 1200 |
| Glucose Test Meter Training | 400 | 1 | 60/60 | 400 |
| Family Members: Questionnaires | 400 | 3 | 45/60 | 900 |
| Teenagers: Questionnaires | 400 | 3 | 30/60 | 600 |
| Total | | | | 4300 |

Dated: December 19, 2002.

Nancy E. Cheal,

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

[FR Doc. 02-32516 Filed 12-24-02; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60 Day-03-25]

Proposed Data Collections Submitted for Public Comment and Recommendations

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Comments are invited on: (a) Whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including

whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) ways to enhance the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology. Send comments to Seleda Perryman, CDC Assistant Reports Clearance Officer, 1600 Clifton Road, MS-D24, Atlanta, GA 30333. Written comments should be received within 60 days of this notice.

Proposed Project

Reducing the Risk of Zoonotic Disease Transmission In Venues Where the Public Has Contact With Animals: A Survey of Current Practices—New—National Center for Infectious Diseases (NCID), Centers for Disease Control and Prevention. The purpose of this project is reduce transmission of zoonotic disease to those persons who interact with farm animals in a number of different settings. Though most of these interactions probably do not result in human illness, several recent outbreaks have highlighted the potential danger of infectious disease transmission in venues where the public comes into contact with animals and their environment. A large outbreak of *E. coli*

O157:H7 infections among visitors to a petting zoo in Pennsylvania in 2000 prompted CDC to develop recommendations to address this issue. Several large outbreaks of *E. coli* O157:H7 have also occurred at county fairs from persons being exposed to animals and their environment. No state or federal laws exist that deal specifically with public health issues relating to interactions between the public and farm animals.

The proposed study consists of a self-administered, written questionnaire mailed to petting zoos and fairs (state, regional, and county). The survey asks individuals to describe their zoo or fair's current practices regarding human interaction with animals, food and beverage consumption in relation to animal interaction areas, and handwashing facilities. The list of zoos comes from facilities licensed by the U.S. Department of Agriculture to show animals for commercial purposes. The list of fairs comes from the International Association of Fairs and Expositions, a private trade organization that volunteered to participate with CDC in having its members complete this survey. Study objectives are to describe current practices and to determine how CDC, other federal agencies, and non-governmental organizations can best educate zoos and fairs about safe animal-human interaction. There is no cost to respondents.

| Survey | No. of re-pondents | No. of responses/respondent | Average burden/response (in hours) | Total burden (in hours) |
|-----------------------------|--------------------|-----------------------------|------------------------------------|-------------------------|
| Written Questionnaire | 1400 | 1 | 10/60 | 233 |
| Total | | | | 233 |