

GOVERNMENT PRINTING OFFICE

Depository Library Council to the Public Printer; Meeting

The Depository Library Council to the Public Printer (DLC) will meet on Sunday, April 6, 2003, through Wednesday, April 9, 2003, in Reno, Nevada. The sessions will take place from 1 p.m. until 4 p.m. and 7 p.m. to 10 p.m. on Sunday, 8:30 a.m. until 5 p.m. on Monday and Tuesday and from 8:30 a.m. until 3:30 p.m. on Wednesday. The meeting will be held at the Peppermill Hotel, 2707 South Virginia Street, Reno, Nevada. The purpose of this meeting is to discuss the Federal Depository Library Program. All sessions are open to the public.

A limited number of rooms are being held for Council attendees at the rate of \$55 (plus tax). The rate for a Friday and/or Saturday night stay is \$79 (plus tax). Reservations can be made by dialing toll free, 1-800-282-2444 or the hotel directly at (775) 826-2121. The rate is good for the meeting dates as well as the three (3) days prior to the meeting and the three (3) days after the meeting. To receive the Government rate, you must make your reservation no later than March 14, 2003 and mention the U.S. Government Printing Office or the Depository Library Council meeting. After that date, rooms will be subject to availability at the best obtainable rate.

Bruce R. James,
Public Printer.

[FR Doc. 03-5431 Filed 3-6-03; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30DAY-23-03]

Agency Forms Undergoing Paperwork Reduction Act Review

The Centers for Disease Control and Prevention (CDC) publishes a list of information collection requests under review by the Office of Management and Budget (OMB) in compliance with the Paperwork Reduction Act (44 U.S.C. Chapter 35). To request a copy of these requests, call the CDC Reports Clearance Officer at (404) 498-1210. Send written comments to CDC, Desk Officer, Human Resources and Housing Branch, New Executive Office Building, Room 10235, Washington, DC 20503. Written comments should be received within 30 days of this notice.

Proposed Project: Evaluating the Impact of Lymphedema and a Lymphedema Management Intervention for Women with Lymphatic Filariasis: Understanding Issues Related to Quality of Life—New—National Center for Infectious Diseases (NCID), Centers for Disease Control and Prevention (CDC). Lymphatic filariasis, a mosquito-transmitted parasitic disease affecting over 120 million people, is the second leading cause of permanent disability worldwide. Globally, lymphatic filariasis causes debilitating genital disease in an estimated 25 million men and lymphedema or elephantiasis of the leg in 15 million people, mostly women in poverty stricken countries. The World Health Organization (WHO) recently identified community management of chronic lymphedema as one of the top twenty lymphatic

filariasis research priorities. Recent advances in the management of chronic lymphedema include a prescribed hygiene and wound care intervention. This intervention has shown promising results in preventing bacterial infections thus reducing acute attacks, and anecdotally improving overall quality of life, alleviating pain and preventing further suffering.

This pilot study will provide a micro-level perspective of women's own experiences of living with lymphedema and others responses to it, illuminating the nature of the disease, the vulnerability of those disabled by the disease, and the impact of an intervention to influence the consequences of having the disease. This study will provide a better understanding, through a combination of qualitative and quantitative methods, the influence of lymphedema as well as the efficacy of a lymphedema management intervention in reducing episodes of bacterial infections and improving quality of life in women with lymphedema in two developing countries.

Women will be queried through in-depth interviews, focus groups, and questionnaire surveys as to the influence of lymphedema on their lives. Quality of life domains that will be explored include physical health, psychological health, social relationships, economic productivity, spiritual health, stigma, and environment. Recommendations will be derived from this study for the global community of lymphatic filariasis researchers in developing countries initiating national and local programs for the management of chronic lymphedema. There are no costs to respondents.

Forms	Number of respondents	Number of responses/respondent	Average burden/response (in hours)
In-depth Survey at Sites A and B	50	1	60/60
Cross-sectional Survey at Sites A and B	200	1	60/60

Dated: February 28, 2003.

Thomas Bartenfeld,

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

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30DAY-30-03]

Agency Forms Undergoing Paperwork Reduction Act Review

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review by the Office of Management and Budget (OMB) in compliance with the Paperwork Reduction Act (44 U.S.C. Chapter 35). To request a copy of these requests, call the CDC Reports Clearance Officer at (404) 498-1210. Send written comments to CDC, Desk Officer, Human Resources and Housing Branch, New Executive Office Building, Room 10235, Washington, DC 20503. Written comments should be received within 30 days of this notice.

Proposed Project: An Evaluation of Targeted Health Communication Messages: Folic Acid and Neural Tube Defects (OMB No. 0920-0461)—Revision—The National Center on Birth Defects and Developmental Disabilities (NCBDDD), Centers for Disease Control and Prevention (CDC).

Background

The Division of Birth Defects and Developmental Disabilities, within NCBDDD launched a national education campaign in January 1999 to increase women's knowledge about neural tube birth defects (NTDs) and the beneficial role folic acid, a B vitamin, plays in the prevention of NTDs. Studies show that a 50 to 70 percent reduction in the risk of neural tube birth defects is possible if all women capable of becoming pregnant consume 400 micrograms of

folic acid daily both prior to and during early pregnancy. Studies also indicate that Hispanic women have a greater risk for NTD-affected pregnancies than women in the general population. Specific, culturally sensitive, targeted media messages need to be directed at this population.

CDC and the March of Dimes Birth Defects Foundation developed health communication media messages and educational materials targeted to health care providers and English- and Spanish-speaking women. These media messages and educational materials consist of television and radio public service announcements (PSA), brochures and resource manuals. The Spanish-language folic acid communication evaluation survey examines the impact of Spanish-

language media messages on the levels of awareness, knowledge, and vitamin use among Hispanic women of childbearing age.

Hispanic women's exposure to Spanish-language media messages and educational materials on folic acid information will be collected and measured to determine whether these exposures influenced the women's knowledge and usage of folic acid. The number and frequency of women's exposures to the media messages such as television and radio PSAs will be collected from media channels and compared to information collected from survey data, National Council on Folic Acid organizations and the National Clearinghouse on Folic Acid activities. The estimated annualized burden is 250 hours.

Respondents	Number of respondents	Number of responses/respondent	Avg. burden/response (in hours)
Telephone Interview	1,000	1	15/60

Dated: February 28, 2003.

Thomas Bartenfeld,

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

[FR Doc. 03-5395 Filed 3-6-03; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30DAY-32-03]

Agency Forms Undergoing Paperwork Reduction Act Review

The Centers for Disease Control and Prevention (CDC) publishes a list of information collection requests under review by the Office of Management and Budget (OMB) in compliance with the Paperwork Reduction Act (44 U.S.C. Chapter 35). To request a copy of these requests, call the CDC Reports Clearance Officer at (404) 498-1210. Send written comments to CDC, Desk Officer, Human Resources and Housing Branch, New Executive Office Building, Room 10235, Washington, DC 20503. Written comments should be received within 30 days of this notice.

Proposed Project: Outcome Follow-up Survey for CDC's Youth Media Campaign—New—National Center for Chronic Disease Prevention and Health

Promotion (NCCDPPH), Centers for Disease Control and Prevention (CDC).

Background

In FY 2001, Congress established the Youth Media Campaign at the Centers for Disease Control and Prevention (CDC). Specifically, the House Appropriations Language said: The Committee believes that, if we are to have a positive impact on the future health of the American population, we must change the behaviors of our children and young adults by reaching them with important health messages. CDC, working in collaboration with federal partners, is coordinating an effort to plan, implement, and evaluate a campaign designed to clearly communicate messages that will help youth develop habits that foster good health over a lifetime. The Campaign is based on principles that have been shown to enhance success, including: designing messages based on research; testing messages with the intended audiences; involving young people in all aspects of Campaign planning and implementation; enlisting the involvement and support of parents and other influencers; refining the messages based on research; and measuring the effect of the campaign on the target audiences.

To measure the effect of the campaign on the target audiences, CDC designed a baseline survey for tween and parent dyads (Children's Youth Media Survey

and Parents' Youth Media Survey) that assessed aspects of the knowledge, attitudes, beliefs, and levels of involvement in positive activities of tweens and a parent or guardian. The baseline survey was conducted prior to the launch of the campaign from April 8, 2002 through June 21, 2002. The methodology was to use a panel design and to survey 3000 dyads (3000 parents and 3000 tweens) from a nationally representative sample and to survey 3000 dyads (again 3000 parents and 3000 tweens) from the six "high dose" communities for a total of 6000 dyads or 12,000 respondents. The survey was conducted using random digit dial.

The next steps in the measurement of effects of the campaign is to collect follow-up data one year post baseline survey and two years post baseline survey. The same panel members (minus attrition) of 6000 tween/parent dyads used in the baseline survey—nationally and in the six selected metropolitan areas—would be re-contacted to complete a survey that would be similar to that used at baseline. Items on campaign awareness would be added to the survey to enable segmentation of the respondents by awareness of the campaign. Thus, the data collection would be with approximately 4,200 tween/parent dyads in spring 2003 and 3,350 tween/parent dyads in 2004. The average annualized burden is 2,571 hours.