

should be mailed or delivered, with two complete copies, to the following address: Federal Trade Commission/ Office of the Secretary, Room 159-H (Annex B), 600 Pennsylvania Avenue, NW., Washington, DC 20580. If the comment contains any material for which confidential treatment is requested, it must be filed in paper (rather than electronic) form, and the first page of the document must be clearly labeled "Confidential."¹ The FTC is requesting that any comment filed in paper form be sent by courier or overnight service, if possible, because U.S. postal mail in the Washington area and at the Commission is subject to delay due to heightened security precautions. Comments filed in electronic form should be sent to the following e-mail box: spywareworkshop2004@ftc.gov.

The FTC Act and other laws the Commission administers permit the collection of public comments to consider and use in this proceeding as appropriate. All timely and responsive public comments, whether filed in paper or electronic form, will be considered by the Commission, and will be available to the public on the FTC Web site, to the extent practicable, at <http://www.ftc.gov/os/comments/spyware/index.html>. As a matter of discretion, the FTC makes every effort to remove home contact information for individuals from the public comments it receives before placing those comments on the FTC Web site. More information, including routine uses permitted by the Privacy Act, may be found in the FTC's privacy policy, at <http://www.ftc.gov/ftc/privacy.htm>.

Donald S. Clark,
Secretary.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Office of the Secretary

[Document Identifier: OS-0990-0208]

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Office of the Secretary, HHS.

In compliance with the requirement of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Office of the Secretary (OS), Department of Health and Human Services, is publishing the following summary of proposed collections for public comment. Interested persons are invited to send comments regarding this burden estimate or any other aspect of this collection of information, including any of the following subjects: (1) The necessity and utility of the proposed information collection for the proper performance of the agency's functions; (2) the accuracy of the estimated burden; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

1. Type of Information Collection Request: Extension of a currently approved collection;

Title of Information Collection: Applicant Background Survey;

Form/OMB No.: OS-0990-0208;

Use: This form will be used to ask applicants for employment how they learned about a vacancy to ensure that recruitment sources yield qualified women and minority applicants, as well as applicants with disabilities, in compliance with EEOC management directives.

Frequency: Reporting;

Affected Public: Individuals or Households;

Annual Number of Respondents: 30,000;

Total Annual Responses: 30,000;

Average Burden Per Response: 2 minutes;

Total Annual Hours: 1,000;

To obtain copies of the supporting statement and any related forms for the proposed paperwork collections referenced above, access the HHS Web site address at <http://www.hhs.gov/oirm/infocollect/pending/> or e-mail your request, including your address, phone number, OMB number, and OS document identifier, to naomi.cook@hhs.gov, or call the Reports Clearance Office on (202) 690-6162. Written comments and

recommendations for the proposed information collections must be mailed within 60 days of this notice directly to the OS Paperwork Clearance Officer designated at the following address: Department of Health and Human Services, Office of the Secretary, Assistant Secretary for Budget, Technology, and Finance, Office of Information and Resource Management, Attention: Naomi Cook (0990-0208), Room 531-H, 200 Independence Avenue, SW., Washington DC 20201.

Dated: March 23, 2004.

Robert E. Polson,

Office of the Secretary, Paperwork Reduction Act Reports Clearance Officer.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day-28-04]

Proposed Data Collections Submitted for Public Comment and Recommendations

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 or by fax to (202) 395-6974. Written comments should be received within 30 days of this notice.

Proposed Project

2004 American Indian Adult Tobacco Survey Pilot Test—New—National Center for Chronic Disease Prevention and Health Promotion (NCCDPHP), Centers for Disease Control and Prevention (CDC).

The purpose of this project is to test and pilot a culturally appropriate Adult Tobacco Survey questionnaire for American Indians and Alaska Natives. This questionnaire will expand data and existing knowledge of tobacco use among American Indians and Alaska Natives in order to benefit tobacco use and prevention surveillance at a tribal, state, and/or regional level. The questions will help to narrow existing gaps in knowledge of tobacco use among different tribes and inform development of tribal-specific interventions.

¹ Commission Rule 4.2(d), 16 CFR 4.2(d). The comment must also be accompanied by an explicit request for confidential treatment, including the factual and legal basis for the request, and must identify the specific portions of the comment to be withheld from the public record. The request will be granted or denied by the Commission's General Counsel, consistent with applicable law and the public interest. See Commission Rule 4.9(c), 16 CFR 4.9(c).

Current smoking prevalence among American Indians and Alaska Natives (36.0 percent) is highest compared to all other racial/ethnic groups (2000 NHIS). While national and regional data exist for American Indians and Alaska Natives, tribal level data is extremely limited. Currently, there are over 500 sovereign tribal nations in the U.S. In order to better understand tobacco use among American Indians and Alaska Natives, CDC is conducting a survey project that includes:

(1) Developing a culturally appropriate Adult Tobacco Survey questionnaire for tribes.

(2) Piloting the final instrument in approximately 24 tribes represented by six Tribal Support Centers (TSCs).

In an effort to better understand the effects of smoking in American Indian and Alaska Native populations, the Support Centers for Tobacco Programs (SCTP) will utilize a culturally appropriate questionnaire for pilot implementation in six different tribal centers. The centers are located in Alaska, California, Oklahoma, Michigan, along with two tribal centers located in the upper Midwest and upper Northwest. In total, the SCTPs will collect 2,691 completed surveys (the

number varying by Center respective to the size of each tribe, 18 years of age and older), which will be representative of distinct tribal communities conducting the survey. The SCTP will be responsible for obtaining the completed surveys. Trained individuals from each of the respective communities and/or support centers will conduct interviews. Most interviews will be conducted face-to-face, with a small proportion conducted by telephone. The total annualized burden is estimated to be 1,794 hours.

Location	Number of respondents	Number of responses per respondent	Avg. burden per response (in hrs.)
Alaska	450	1	40/60
California	466	1	40/60
Michigan	450	1	40/60
Oklahoma	600	1	40/60
Upper Midwest	350	1	40/60
Upper Northwest	375	1	40/60

Dated: March 25, 2004.

Joe E. Salter,

Acting Director, Management Analysis and Services Office, Centers for Disease Control and Prevention.

[FR Doc. 04-7308 Filed 3-31-04; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day-23-04]

Proposed Data Collections Submitted for Public Comment and Recommendations

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 or by fax to (202) 395-6974. Written comments should be received within 30 days of this notice.

Proposed Project: Descriptive Epidemiology of Missed or Delayed Diagnoses for Conditions Detected by Newborn Screening—New—National Center for Environmental Health

(NCEH), Centers for Disease Control and Prevention (CDC).

Every state in the United States and Washington, DC, has a public health program to test newborn babies for congenital metabolic and other disorders through laboratory testing of dried blood spots. These programs screen for between 4 and 30 different conditions including phenylketonuria (PKU) and congenital hypothyroidism, with testing performed in both state laboratories and private laboratories contracted by state health departments. The screening process or system is broader than the state public health newborn screening program, which is composed only of the laboratory and follow-up personnel. It involves the collection of blood from a newborn, analysis of the sample in a screening laboratory, follow-up of abnormal results, confirmatory testing and diagnostic work-up. Parents, hospitals, medical providers including primary care providers and specialists, state laboratory and follow-up personnel advocates, as well as other partners such as local health departments, police, child protection workers, and courts play important roles in this process.

Most children born with metabolic disease are identified in a timely manner and within the parameters defined by the newborn screening system of each state. These children are referred for diagnosis and treatment. However, some cases are not detected at all or the detection comes too late to prevent harm. These “missed cases”

often result in severe morbidity such as mental retardation or death.

In this project, we will update and expand a previous epidemiological study of missed cases of two disorders published in 1986. We will assess the number of cases of each disorder missed, and the reasons for the missed and legal outcomes, if any. The reasons for the missed will be tabulated according to which step or steps of the screening process it occurred. Data will be collected by asking state public health laboratory directors, newborn screening laboratory managers, follow-up coordinators, specialists at metabolic clinics and parent groups with an interest in newborn screening, for information regarding missed cases. An estimated 269 subjects (with an expected response rate of 80% from metabolic clinics, Lab Directors and Coordinators) will be requested to complete a short questionnaire that asks for information regarding the details of any missed cases of which they are aware.

The survey will highlight procedures and actions taken by states and other participants in newborn screening systems to identify causes of missed cases and to modify policies and procedures to prevent or minimize recurrences. The information gleaned from this study may be used to help craft changes in the screening protocols that will make the process more organized and efficient and less likely to fail an affected child. Furthermore, it is not clear that there is a systematic