medical care services in the United States. Ambulatory services are rendered in a wide variety of settings. including physicians' offices and hospital outpatient and emergency departments. The NAMCS target population consists of all office visits made by ambulatory patients to non-Federal office-based physicians (excluding those in the specialties of anesthesiology, radiology, and pathology) who are engaged in direct patient care. To complement these data, NCHS initiated the National Hospital Ambulatory Medical Care Survey (NHAMCS, OMB No. 0920-0278) to provide data concerning patient visits to hospital outpatient and emergency departments.

The NAMCS provides a range of baseline data on the characteristics of the users and providers of ambulatory medical care. Data collected include the patients' demographic characteristics, reason(s) for visit, physicians' diagnosis, diagnostic services, medications and visit disposition. In addition to the annual statistics normally collected, a key focus of the 2005–2006 survey will be on the prevention and treatment of selected chronic conditions. These data, together with trend data, may be used to monitor the effects of change in the health care system, provide new insights into ambulatory medical care,

and stimulate further research on the use, organization, and delivery of ambulatory care.

Users of NAMCS data include, but are not limited to, congressional and other federal government agencies, state and local governments, medical schools, schools of public health, researchers, administrators, and health planners. NAMCS plans to extend its data collection into 2005 and 2006. To calculate the burden hours the number of respondents for NAMCS is based on a sample of 3,000 physicians with a 50 percent participation rate (this includes physicians who are out-of-scope as well as those who refuse). There is no cost to respondents.

Respondents	Number of re- spondents	Number of re- sponses/re- spondent	Average bur- den/response (in hrs)	Total burden hours
Office-based physicians: Induction Form Patient Record Form	1,500 1,500	1 30	25/60 5/60	625 3,750
Total				4,375

Dated: December 19, 2003.

# Ron Ergle,

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

[FR Doc. 03–32164 Filed 12–30–03; 8:45 am] BILLING CODE 4163–18–P

#### DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Disease Control and Prevention

## [30 Day-13-04]

### 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: Housing and Health Study—New—National Center for HIV, STD, and TB Prevention (NCHSTP), Centers for Disease Control and Prevention (CDC). CDC is requesting OMB approval to conduct a study to examine the impact of providing housing for homeless or unstably housed people (people who are in temporary housing programs or doubled up with others) while living with HIV.

This project includes a unique collaboration with the Department of Housing and Urban Development (HUD). HUD is providing funding for housing vouchers for study participants. CDC will use the results of the data collection to inform policy makers about the types of housing and other affiliated services most likely to reduce HIV transmission and disease progression in the homeless population.

The population to be studied will be drawn from persons living with HIV/ AIDS who are seeking housing services from three communities with unmet housing needs. These needs are evidenced by a waiting list for services, or other evidence of unmet housing need through the Housing Opportunities for Persons with AIDS (HOPWA) program. The project will be a longitudinal cohort study, following participants for 18 months. Participants will be randomized into two groups. One group will receive vouchers for housing subsidies plus a 2-session behavioral intervention; the other group will receive referral to housing resources through participating agencies and other agencies plus the 2-session behavioral intervention. No study participants will be denied access to other housing services that are available through participating agencies or other community resources.

Since, all participants receive the behavioral intervention, the study technically assesses the effects of housing over and above the behavioral intervention. A cost study will also be conducted to determine the resources needed for this approach and the cost benefits of providing housing for homeless and unstably housed people living with HIV. The purpose of the cost study is to evaluate the effects of housing affordability and the costeffectiveness (i.e. cost-utility ratio) of this strategy relative to other interventions in other public health and other HIV prevention interventions.

Study participants will be surveyed at the beginning of the project (baseline) and at 6, 12, and 18 months after baseline. HUD site service providers will also be surveyed. Blood samples for CD4 and viral load counts will also be collected for all participants. The annualized burden for this data collection is 6,030 hours.

Respondents	Number of respondents	Responses per respondent	Average bur- den per re- sponse (in hours)
HOPWA Program Participants	1,000	4	1.5
HUD Site Service Providers	15	1	2

Dated: December 22, 2003.

Ron Ergle,

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

[FR Doc. 03–32165 Filed 12–30–03; 8:45 am] BILLING CODE 4163–18–P

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

#### Centers for Disease Control and Prevention

[60Day-04-19]

#### 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–E11, Atlanta, GA 30333. Written comments should be received within 60 days of this notice.

*Proposed Project:* HIV/AIDS Prevention and Surveillance Project Reports, OMB No. 0920–0208— Extension—National Center for HIV, STD and TB Prevention (NCHSTP), Centers for Disease Control and Prevention (CDC).

CDC is requesting to extend the use of the currently approved form, OMB No. 0920-0208, for collecting HIV counseling, testing, and referral (CTR) program data. This current form expires March 30, 2004. This request is for an 18-month clearance past this date. Extension of the current form will allow grantees to continue to collect CTR data as they transition to the new set of CTR variables and the new program evaluation and monitoring system (PEMS). Over the next year, grantees will either transition to the new variables once they have reprogrammed their existing computer systems, or as the CDC-provided PEMS is made available. CDC funds cooperative agreements for 65 HIV prevention projects (50 states, 6 cities, 7 territories, Washington, DC, and Puerto Rico) and approximately 50 community based organizations to support HIV counseling, testing, and referral programs.

HIV counseling, testing, and referral services in STD clinics, women's health centers, drug treatment centers, and other health facilities have been described as a primary prevention strategy of the national HIV prevention program. The funded public health departments and community based organizations have increased the provision of HIV counseling, testing, and referral activities to those at increased risk for acquiring or transmitting HIV, as well as minority communities and women of child bearing age.

CDC is responsible for monitoring and evaluating HIV prevention programs conducted under HIV prevention cooperative agreements. HIV counseling, testing, and referral services are a vital component of HIV prevention programs. Without data to monitor and evaluate the impact of HIV counseling, testing, and referral programs, HIV prevention program priorities cannot be assessed and improved to prevent further spread of the epidemic. CDC needs minimal core data from all grantees describing CTR services provided for at-risk persons. Until grantees are prepared for collecting the new CTR variables and reporting data electronically through PEMS, it is essential that they be allowed to continue to collect the current CTR data using the existing forms.

Completing the initial data submission will take approximately 5 minutes per form. Approximately two (2) million records annually are expected from over 11,000 directly and indirectly funded grantee facilities. The total estimated burden is 167,000 hours annually. This is the estimated burden if no one transitions to the new system during the year, but it is expected that many of the grantees will transition to PEMS in phases throughout the year. Following this notice, a separate data collection for PEMS will be submitted for public comment and will include the revised CTR data variables and associated burden estimate. There is no cost to respondents.

Respondents	Number of re- spondents	Number of re- sponses/re- spondent	Average bur- den/response (in hours)	Total burden (in hours)
Directly or Indirectly Funded Facilities	11,000	182	5/60	167,000
Total				167,000