

- A. Sexual functioning
- B. Ability to participate in an exercise program
- C. Ability to return to work
- D. Physical performance test pain (joint pain, joint aches)
- E. Regular daily activities
- F. Polypharmacy
- G. Admission to a skilled-nurse facility

XVII. Access to plastic surgery

XVIII. Readmissions/rehospitalizations

Timing:

No time limit

Setting:

Any

Sharon B. Arnold,

AHRQ Deputy.

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BILLING CODE 4160-90-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day-17-0770]

Agency Forms Undergoing Paperwork Reduction Act Review

The Centers for Disease Control and Prevention (CDC) has submitted the following information collection request to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act of 1995. The notice for the proposed information collection is published to obtain comments from the public and affected agencies.

Written comments and suggestions from the public and affected agencies concerning the proposed collection of information are encouraged. Your comments should address any of the following: (a) Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility; (b) Evaluate the accuracy of the agencies estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (c) Enhance the quality, utility, and clarity of the information to be collected; (d) Minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, e.g., permitting electronic submission of responses; and (e) Assess information collection costs.

To request additional information on the proposed project or to obtain a copy of the information collection plan and instruments, call (404) 639-7570 or send an email to omb@cdc.gov. Written comments and/or suggestions regarding the items contained in this notice should be directed to the Attention: CDC Desk Officer, Office of Management and Budget, Washington, DC 20503 or by fax to (202) 395-5806. Written comments should be received within 30 days of this notice.

Proposed Project

National HIV Behavioral Surveillance System ((NHBS), OMB Control No. 0920-0770, exp. 03/31/2017)—Revision—National Center for HIV, Hepatitis, STD, and TB Prevention (NCHHSTP), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

The CDC currently sponsors the National HIV Behavioral Surveillance (NHBS) System. The system is designed to describe and monitor the HIV risk behaviors, HIV seroprevalence and incidence, and HIV prevention experiences of persons at highest risk for HIV infection in the United States. NHBS awardees are state and local health departments that provide HIV-related services, conduct NHBS interviews, and submit non-identifiable information to CDC. To be eligible for NHBS funding, a health department must serve one of the 30 Metropolitan Statistical Areas (MSA) in the U.S. with high HIV prevalence. Twenty-two (22) programs receive NHBS funding and technical assistance from CDC at this time. Burden estimates are based on current availability of funds and recruitment targets for 22 CDC-funded NHBS awardees. If additional funding is received to support the participation of additional sites, CDC will submit a Change Request to make the appropriate adjustments to the total estimated annualized burden.

Information collection is based on rotating annual “cycles” of surveillance with three populations: Men who have sex with men (MSM), injecting drug users (IDUs), and heterosexuals at increased risk of HIV (HET). Screening interviews and specialized behavioral assessment interviews are conducted once every three years with each population: MSM in year 1, IDU in year 2, and HET in year 3. The target number of annual interviews for each NHBS-funded awardee is 500. Due to differences in the risk characteristics of the MSM, IDU and HET groups, the behavioral assessment is customized for each group. In addition, an HIV test and

pre-test counseling session are offered to all persons who participate in an NHBS interview.

The surveillance system is focused on behaviors directly related to HIV transmission and those that are amenable to intervention through prevention programs. Information collected through the NHBS System allows CDC to: (a) Describe the prevalence of and trends in risk behaviors; (b) describe the prevalence of and trends in HIV testing and HIV infection; (c) describe the prevalence of and trends in use of HIV prevention services; and (d) identify met and unmet needs for HIV prevention services in order to inform health departments, community-based organizations, community planning groups and other stakeholders. No other federal agency systematically collects this type of information from persons at risk for HIV infection.

Venue-based sampling methods are used to identify respondents for the MSM information collection cycle and respondent-driven sampling methods are used to identify respondents for the IDU cycle and the HET cycle. Consistent with these methods, persons who participate in the IDU and HET interviews may be trained to recruit additional respondents. Each person who serves as a peer recruiter will be asked to participate in a short debriefing interview.

CDC requests OMB approval to continue information collection for three years, with revisions. Selected questions in the eligibility screener and the behavioral assessment interview instruments will be updated to improve usability and data quality, and new questions will be added to provide measures of high priority emerging issues including pre-exposure prophylaxis, treatment as prevention, and opioid use and abuse. Lower priority questions and repetitive content will be deleted in order to manage project cost and respondent burden. There are no changes to the estimated burden per response for any information collection instrument. However, total burden will decrease due to a reduction in the number of health departments funded to participate in the NHBS System (from 25 to 22). Compared to the previous period of OMB approval, this will reduce the total estimated number of interviews for each cycle from 12,500 (4,167 annualized) to 11,000 (3,667 annualized).

Information collected through the NHBS has a substantial impact on the design and delivery of targeted prevention programs aimed at reducing new HIV infections and evaluating

progress towards national public health goals. Participation is voluntary and there is no cost to respondents other

than their time. The total estimated annualized burden hours are 8,735.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hr)
Persons Screened Eligible Participants	Eligibility Screener	13,142	1	5/60
	Behavioral Assessment for MSM	3,667	1	30/60
	Behavioral Assessment for IDU	3,667	1	54/60
	Behavioral Assessment for HET	3,667	1	39/60
Peer Recruiters	Recruiter Debriefing	3,667	1	2/60

Leroy A. Richardson,
Chief, Information Collection Review Office,
Office of Scientific Integrity, Office of the
Associate Director for Science, Office of the
Director, Centers for Disease Control and
Prevention.

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**DEPARTMENT OF HEALTH AND
HUMAN SERVICES**

**Centers for Disease Control and
Prevention**

[60Day-17-0904; Docket No. CDC-2016-0117]

**Proposed Data Collection Submitted
for Public Comment and
Recommendations**

AGENCY: Centers for Disease Control and
Prevention (CDC), Department of Health
and Human Services (HHS).

ACTION: Notice with comment period.

SUMMARY: The Centers for Disease
Control and Prevention (CDC), as part of
its continuing effort to reduce public
burden and maximize the utility of
government information, invites the
general public and other Federal
agencies to take this opportunity to
comment on proposed and/or
continuing information collections, as
required by the Paperwork Reduction
Act of 1995. This notice invites
comment on a proposed revision of the
“SEARCH for Diabetes in Youth Study,”
a national multi-center study aimed at
understanding more about diabetes
among children and young adults in the
United States.

DATES: Written comments must be
received on or before February 6, 2017.

ADDRESSES: You may submit comments,
identified by Docket No. CDC-2016-
0117 by any of the following methods:

- *Federal eRulemaking Portal:*
Regulations.gov. Follow the instructions
for submitting comments.

- *Mail:* Leroy A. Richardson,
Information Collection Review Office,
Centers for Disease Control and
Prevention, 1600 Clifton Road NE., MS-
D74, Atlanta, Georgia 30329.

Instructions: All submissions received
must include the agency name and
Docket Number. All relevant comments
received will be posted without change
to *Regulations.gov*, including any
personal information provided. For
access to the docket to read background
documents or comments received, go to
Regulations.gov.

Please note: All public comment
should be submitted through the
Federal eRulemaking portal
(*regulations.gov*) or by U.S. mail to the
address listed above.

FOR FURTHER INFORMATION CONTACT: To
request more information on the
proposed project or to obtain a copy of
the information collection plan and
instruments, contact the Information
Collection Review Office, Centers for
Disease Control and Prevention, 1600
Clifton Road NE., MS-D74, Atlanta,
Georgia 30329; phone: 404-639-7570;
Email: *omb@cdc.gov*.

SUPPLEMENTARY INFORMATION: Under the
Paperwork Reduction Act of 1995 (PRA)
(44 U.S.C. 3501-3520), Federal agencies
must obtain approval from the Office of
Management and Budget (OMB) for each
collection of information they conduct
or sponsor. In addition, the PRA also
requires Federal agencies to provide a
60-day notice in the **Federal Register**
concerning each proposed collection of
information, including each new
proposed collection, each proposed
extension of existing collection of
information, and each reinstatement of
previously approved information
collection before submitting the
collection to OMB for approval. To
comply with this requirement, we are
publishing this notice of a proposed
data collection as described below.

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; (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; and (e) estimates of capital
or start-up costs and costs of operation,
maintenance, and purchase of services
to provide information. Burden means
the total time, effort, or financial
resources expended by persons to
generate, maintain, retain, disclose or
provide information to or for a Federal
agency. This includes the time needed
to review instructions; to develop,
acquire, install and utilize technology
and systems for the purpose of
collecting, validating and verifying
information, processing and
maintaining information, and disclosing
and providing information; to train
personnel and to be able to respond to
a collection of information, to search
data sources, to complete and review
the collection of information; and to
transmit or otherwise disclose the
information.

Proposed Project

SEARCH for Diabetes in Youth Study
(OMB Control No. 0920-0904, Expires
8/31/2017)—Revision—National Center
for Chronic Disease Prevention and
Health Promotion (NCCDPHP), Centers
for Disease Control and Prevention
(CDC).

Background and Brief Description

Diabetes is one of the most common
chronic diseases among children in the
United States. When diabetes strikes
during childhood, it is routinely
assumed to be type 1, or juvenile-onset,
diabetes. Type 1 diabetes (T1D)