

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Centers for Disease Control and Prevention**

[30Day-16-16AOP]

**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](mailto: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**

*TRAUMATIC BRAIN INJURY (TBI) SURVEILLANCE SYSTEM—New—National Center for Injury Prevention and Control (NCIPC), Centers for Disease Control and Prevention (CDC)*

**Background and Brief Description**

The Centers for Disease Control and Prevention (CDC) requests Office of Management and Budget (OMB) approval for three years for a new data collection for a Traumatic Brain Injury (TBI) Surveillance System. TBI is a significant public health concern in the United States, contributing to an estimated 2.2 million Emergency Department (ED) visits, 280,000 hospitalizations, and 50,000 deaths in 2010. These numbers, however, underestimate the true public health and economic burden of TBI in the U.S. because they are based on healthcare

administrative data that only capture information on the number of ED visits, hospitalizations, and deaths identified as TBI-related. A surveillance system will accurately determine how many children and adults experience a TBI each year in the United States, and will collect information about the circumstances that identifies groups most at risk for TBI. By administering the surveillance system over time, the surveillance system can monitor trends and allow for an understanding of whether TBIs are increasing or decreasing, and whether prevention efforts are effective.

Data will be collected through household survey conducted as a random digit dial telephone survey utilizing both landline and cellphones; adult respondents will be asked about their own TBI history while adult respondents with children 5-17 years of age will serve as proxies and answer questions about their children's TBI history. Information collected will produce nationally representative incidence estimates of all TBI, with a particular focus on the incidence of sports- and recreation-related TBI (SRR-TBI) among youth 5-21 years of age. Another use of the data is to produce nationally-representative estimates of TBI-related disability. Participation in the information collection is voluntary. The survey will be conducted among English or Spanish speaking participants living in the United States.

The estimated annual burden hours are 4,612. There are no costs to respondents other than their time.

**ESTIMATED ANNUALIZED BURDEN HOURS**

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)
Adults 18 or older .....	Adult Eligibility Screener .....	2,611	1	2/60
	Adult Screener .....	14,167	1	14/60
	Adult Survey .....	2,500	1	21/60
Adolescent 12 to 17 years of age .....	Adolescent Screener .....	2,058	1	5/60
	Adolescent Survey .....	686	1	15/60

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 Prevention.

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