

organization or other affiliation, full address and phone, fax, and e-mail information or e-mail this information to *FindYouthInfo@air.org*. Additional identification documents may be required.

Dated: November 4, 2010.

**Sherry Glied,**

*Assistant Secretary for Planning and Evaluation.*

**Authority:** Division F, Pub. L. 111–8; E.O. 13459, 73 FR 8003, February 12, 2008.

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**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Centers for Disease Control and Prevention**

[30Day–10–10DE]

**Agency Forms Undergoing Paperwork Reduction Act Review**

The Centers for Disease Control and Prevention (CDC), Agency for Toxic Substances and Disease Registry (ATSDR) publishes a list of information collection requests under review by the Office of management and Budget (OMB) in compliance with the Paperwork Reduction Act (33 U.S.C. chapter 35). To request a copy of these requests, call the CDC Reports Clearance Officer, at (404) 639–5960 or send an e-mail to *omb@cdc.gov*. Send written comments to ATSDR 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**

Creation of State and Metropolitan Area-based Surveillance Projects for Amyotrophic Lateral Sclerosis (ALS)—New—Agency for Toxic Substances and Disease Registry (ATSDR).

**Background and Brief Description**

On October 10, 2008, President Bush signed S. 1382: ALS Registry Act which amended the Public Health Service Act to provide for the establishment of an Amyotrophic Lateral Sclerosis (ALS) Registry. The activities described are part of the effort to create the National ALS Registry. The purpose of the registry is to: (1) Better describe the incidence and prevalence of ALS in the United States; (2) examine appropriate factors, such as environmental and occupational, that might be associated with the disease; (3) better outline key demographic factors (such as age, race or ethnicity, gender, and family history of individuals diagnosed with the disease); and (4) better examine the connection between ALS and other motor neuron disorders that can be confused with ALS, misdiagnosed as ALS, and in some cases progress to ALS. The registry will collect personal health information that may provide a basis for further scientific studies of potential risks for developing ALS.

This project purposes to collect information-specific data related to

ALS. The objective of this project is to develop state-based and metropolitan area-based surveillance projects for ALS. The primary goal of the state-based and metropolitan area-based surveillance project is to use these data to evaluate the completeness of the National ALS Registry. The secondary goal of the surveillance project is to obtain reliable and timely information on the incidence and prevalence of ALS and to better describe the demographic characteristics (e.g., age, race, sex, and geographic location) of those with ALS.

Neurologists or their staff will complete an ALS Case Reporting Form on each of their ALS patients. This will be transmitted to the state or metropolitan health department. The contract surveillance staff assigned to the state and metropolitan area health departments will train medical personnel how to complete the ALS Case Reporting Form (Attachment 3) and assist with abstracting records as requested. An ALS Medical Record Verification Form will be collected on a subset of cases reported. Each medical provider reporting source should keep a line listing of individuals diagnosed with or thought to have ALS along with information on whether or not the case was reported and if not, the reason. Surveillance items to be collected include information to make sure that there are no duplicates. There are no costs to the respondents other than their time. The estimated annualized burden hours are 703.

**ESTIMATED ANNUALIZED BURDEN HOURS**

Type of data collection instrument	Type of respondent	Number of respondents	Number of responses per respondent	Average burden per response (in hours)
Training .....	Medical Personnel/Neurologist .....	243	1	30/60
ALS Case Reporting Form .....	Medical Personnel/Neurologist .....	2,250	1	5/60
ALS Medical Record Verification Form .....	Neurologist .....	450	1	20/60
Line Listing (record keeping) .....	Medical Personnel .....	243	1	1

Dated: November 4, 2010.

**Carol E. Walker,**

*Acting Reports Clearance Officer, Centers for Disease Control and Prevention/Agency for Toxic Substances and Disease Registry.*

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**BILLING CODE P**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**National Institutes of Health**

**Submission for OMB review; comment request; NCCAM Office of Communications and Public Liaison Communications Program Planning and Evaluation Research**

**SUMMARY:** Under the provisions of Section 3507(a)(1)(D) of the Paperwork Reduction Act of 1995, the National Center for Complementary and Alternative Medicine (NCCAM), the

National Institutes of Health (NIH), has submitted to the Office of Management and Budget (OMB) a request to review and approve the information collection listed below. This proposed information collection was previously published in the **Federal Register** on August 25, 2010 (Vol. 75, No. 164, p. 52349) and allowed 60-days for public comment. There was one public comments received during this time. The purpose of this notice is to allow an additional 30 days for public comment. The National Institutes of Health may not conduct or sponsor, and the respondent is not required to