

strategic plan for Federal youth policy; and prepare recommendations to improve the coordination, effectiveness, and efficiency of programs affecting youth.

The Interagency Working Group on Youth Programs is comprised of staff from twelve Federal agencies that support programs and services that focus on youth: the U.S. Department of Agriculture; U.S. Department of Commerce; U.S. Department of Defense; U.S. Department of Education; U.S. Department of Health and Human Services (Chair); U.S. Department of Housing and Urban Development; U.S. Department of Justice (Vice-Chair); U.S. Department of Labor; U.S. Department of the Interior; U.S. Department of Transportation; Corporation for National and Community Service; and Office of National Drug Control Policy.

The Working Group seeks to promote achievement of positive results for at-risk youth through the following activities:

- Promoting enhanced collaboration at the Federal, State, and local levels, including with faith-based and other community organizations, as well as among families, schools and communities, in order to leverage existing resources and improve outcomes;
- Disseminating information about critical resources, including evidence-based programs, to assist interested citizens and decision-makers, particularly at the community level, to plan, implement, and participate in effective strategies for at-risk youth;
- Developing an overarching strategic plan for Federal youth policy, as well as recommendations for improving the coordination, effectiveness and efficiency of youth programs, using input from community stakeholders, including youth; and
- Producing a Federal Web site, *FindYouthInfo.gov*, to promote effective community-based efforts to reduce the factors that put youth at risk and to provide high-quality services to at-risk youth.

II. Registration, Security, Building, and Parking Guidelines

For security purposes, members of the public who wish to attend the meeting must pre-register online at <http://www.findyouthinfo.gov> no later than November 9, 2010. Should problems arise with Web registration, call the help desk at 1-877-231-7843 or send a request to register for the meeting to FindYouthInfo@air.org. To register, complete the online registration form, which will ask for your name, title,

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. [FR Doc. 2010-28392 Filed 11-9-10; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Office of the Secretary

Public Meeting To Solicit Input for a Strategic Plan for Federal Youth Policy

AGENCY: Office of the Assistant Secretary for Planning and Evaluation, DHHS.

ACTION: Notice of meeting.

SUMMARY: The U.S. Department of Health and Human Services, in its role as the Chair of the Interagency Working Group on Youth Programs, is announcing a meeting to solicit input from the public that will inform the development of a strategic plan for federal youth policy.

DATES: November 18, 2010, from 9 a.m. to 1 p.m.

ADDRESSES: The meeting will take place at the Houston Housing Authority Neighborhood Resource Center at 815 Crosby Street, Houston, TX 77019.

FOR FURTHER INFORMATION CONTACT: Visit the Web site for the Interagency Working Group on Youth Programs at <http://www.FindYouthInfo.gov> for information on how to register, or contact the Interagency Working Group on Youth Programs help desk, by telephone at 1-877-231-7843 [**Note:** this is a toll-free telephone number], or by e-mail at FindYouthInfo@air.org.

SUPPLEMENTARY INFORMATION:

I. Background

On March 11, 2009, the Congress passed the Omnibus Appropriations Act, 2009 (Pub. L. 111-8). The House Appropriations Committee Print, Division F—Departments of Labor, Health and Human Services, and Education, and Related Agencies Appropriations directed the Interagency Working Group on Youth Programs to solicit input from young people, State children's cabinet directors, and non-profit organizations on youth programs

and policies; develop an overarching strategic plan for Federal youth policy; and prepare recommendations to improve the coordination, effectiveness, and efficiency of programs affecting youth.

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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.
[FR Doc. 2010–28396 Filed 11–9–10; 8:45 am]
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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.
[FR Doc. 2010–28337 Filed 11–9–10; 8:45 am]
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