The registration portion of the data collection will be limited to information that can be used to identify an individual to assure that there are not duplicate records for an individual. Avoiding duplication of registrants due to obtaining records from multiple sources is imperative to get accurate estimates of incidence and prevalence, as well as accurate information on demographic characteristics of the cases of ALS.

In addition to questions required for registration, there will be a series of short surveys to collect information on such things as military history, occupations, residential history, and family history that would not likely be available from other sources. This project proposes to add 10 additional risk factor surveys while continuing to collect information on individuals with ALS which can be combined with information obtained from existing sources of information. This combined data will become the National ALS Registry and will be used to provide more accurate estimates of the incidence and prevalence of disease as well as the demographic characteristics of the cases. Information obtained from the surveys will be used to better characterize potential risk factors for ALS which will lead to further in-depth studies.

The existence of the Web site (http://www.cdc.gov/als) is being advertised by ATSDR and advocacy groups such as the Amyotrophic Lateral Sclerosis Association (ALSA) and the Muscular Dystrophy Association (MDA). There are no costs to the respondents other than their time. The estimated annualized burden hours are 1,375.

### ESTIMATED ANNUALIZED BURDEN HOURS

<table>
<thead>
<tr>
<th>Type of respondent</th>
<th>Form name</th>
<th>Number of respondents</th>
<th>Number of responses per respondent</th>
<th>Average burden per response (in hours)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Person with ALS</td>
<td>Validation questions (Screener) for suspected ALS cases</td>
<td>1,670</td>
<td>1</td>
<td>2/60</td>
</tr>
<tr>
<td></td>
<td>Registration Form of ALS cases</td>
<td>1,500</td>
<td>1</td>
<td>7/60</td>
</tr>
<tr>
<td></td>
<td>Cases of ALS completing 1-time surveys</td>
<td>750</td>
<td>16</td>
<td>5/60</td>
</tr>
<tr>
<td></td>
<td>Cases of ALS completing twice yearly surveys</td>
<td>750</td>
<td>2.3</td>
<td>5/60</td>
</tr>
</tbody>
</table>

Ron A. Otten,
Director, Office of Scientific Integrity, Office of the Associate Director for Science, Office of the Director, Centers for Disease Control and Prevention.

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Administration for Native Americans Tribal Consultation; Notice of Meeting

AGENCY: Administration for Children and Families, Department of Health and Human Services.

ACTION: Notice of tribal consultation.

SUMMARY: The Department of Health and Human Services (HHS), Administration for Children and Families (ACF) will host a tribal consultation to consult on ACF programs and tribal priorities.

DATES: July 9–10, 2013.

ADDRESSES: 901 D Street SW., 7th Floor Multipurpose Room, Washington, DC 20024.

FOR FURTHER INFORMATION CONTACT: Lillian A. Sparks, Commissioner, Administration for Native Americans at 202–401–5590, by email at Lillian.sparks@acf.hhs.gov or by mail at 370 L’Enfant Promenade SW., 2 West, Washington, DC 20447.

SUPPLEMENTARY INFORMATION: On November 5, 2009, President Obama signed the “Memorandum for the Heads of Executive Departments and Agencies on Tribal Consultation.” The President stated that his Administration is committed to regular and meaningful consultation and collaboration with tribal officials in policy decisions that have tribal implications, including, as an initial step, through complete and consistent implementation of Executive Order 13175.

The United States has a unique legal and political relationship with Indian tribal governments, established through and confirmed by the Constitution of the United States, treaties, statutes, executive orders, and judicial decisions. In recognition of that special relationship, pursuant to Executive Order 13175 of November 6, 2000, executive departments and agencies are charged with engaging in regular and meaningful consultation and collaboration with tribal officials in the development of federal policies that have tribal implications, and are responsible for strengthening the government-to-government relationship between the United States and Indian tribes.

HHS has taken its responsibility to comply with Executive Order 13175 very seriously over the past decade, including the initial implementation of a department-wide policy on tribal consultation and coordination in 1997, and through multiple evaluations and revisions of that policy, most recently in 2010. ACF has also developed its own agency-specific consultation policy that complements the department-wide efforts.

ACF’s Administration for Native Americans (ANA) will hold a tribal consultation on the morning of July 9, 2013, to discuss the reauthorization of ANA’s authorizing legislation, the Native American Programs Act, and the development of data collection elements to collect information on the impact of ACF funding on the members of a tribal community.

A tribal resource day will begin the afternoon of July 9 and the ACF Tribal Consultation Session will begin the morning of July 10 and continue throughout the day until all discussions have been completed. Other ACF
DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Community Living

Agency Information Collection Activities; Submission for OMB Review; Comment Request; State Program Report

AGENCY: Administration for Community Living, HHS.

ACTION: Notice.

SUMMARY: The Administration for Community Living (ACL) is announcing that the proposed collection of information listed below has been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995.

DATES: Submit written comments on the collection of information by June 7, 2013.

ADDRESSES: Submit written comments on the collection of information by fax 202.395.5806 or by email to OIRA_submission@omb.eop.gov, Attn: OMB Desk Officer for ACL.

FURTHER INFORMATION CONTACT: Elena Fazio at 202–357–3583 or email: elena.fazio@acf.hhs.gov.

SUPPLEMENTARY INFORMATION: In compliance with 44 U.S.C. 3507, ACL has submitted the following proposed collection of information to OMB for review and clearance.

The Older Americans Act (OAA) requires annual program performance reports from States. In compliance with this OAA provision, ACL developed a State Program Report (SPR) in 1996 as part of its National Aging Program Information System (NAPIS).

The SPR collects information about how State Units on Aging expend their OAA funds as well as funding from other sources for OAA authorized supportive services. The SPR also collects information on the demographic and functional status of the recipients, and is a key source for AOA performance measurement. This collection includes minor revisions of the format from the 2010 approved version. The proposed revised version will be in effect for the FY 2014 reporting year and thereafter, while the current reporting, OMB Approval Number 0985–0008, will be extended to the end of the FY 2013 reporting cycle. The proposed FY 2014 version may be found on the ACL Web site link entitled Proposed SPR for Review available at http://www.aoa.gov/AoARoot/Program_Results/OAA_Performance.aspx#national ACL estimates the burden of this collection of information as follows: 2,600 hours.


Kathy Greenlee,
Administrator and Assistant Secretary for Aging.

[FR Doc. 2013–10921 Filed 5–7–13; 8:45 am]
BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2013–N–0093]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request: Evaluation of the Program for Enhanced Review Transparency and Communication for New Molecular Entity New Drug Applications and Original Biologics License Applications in Prescription Drug User Fee Act

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a proposed collection of information has been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995.

DATES: Fax written comments on the collection of information by June 7, 2013.

ADDRESSES: To ensure that comments on the information collection are received, OMB recommends that written comments be faxed to the Office of Information and Regulatory Affairs, OMB, Attn: FDA Desk Officer, FAX: 202–395–7285, or emailed to oira_submission@omb.eop.gov. All comments should be identified with the OMB control number 0910–New and title Evaluation of the Program for Enhanced Review Transparency and Communication for New Molecular Entity New Drug Applications and Original Biologics License Applications in Prescription Drug User Fee Act. Also include the FDA docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Ila S. Mizrachi, Office of Information Management, Food and Drug Administration, 1350 Piccard Dr., PI50–400B, Rockville, MD 20850, 301–796–7726, Ila.Mizrachi@fda.hhs.gov.

SUPPLEMENTARY INFORMATION: In compliance with 44 U.S.C. 3507, FDA has submitted the following proposed collection of information to OMB for review and clearance.