confidential information. All the other completed model trust certificates and model trust documents (except for any trust provisions that relate to the testamentary disposition of trust assets) are retained and made publicly available based upon a proper request under EIGA (by filling out an OGE Form 201 access form) until the periods for retention of all other reports (usually the OGE Form 278 Public Financial Disclosure Reports) of the individual establishing the trust have lapsed (generally six years after the filing of the last other report). See 5 CFR 2634.600(g)(2) of OGE’s executive branch financial disclosure regulation.

The U.S. Office of Government Ethics administers the qualified trust program for the executive branch. At the present time, there are no active filers using the trust model certificates and documents. However, OGE intends to submit to OMB a request for extension of approval for two reasons. First, under OMB’s implementing regulations for the Paperwork Reduction Act, at 5 CFR 1320.3(c)(4)(I), any recordkeeping, reporting or disclosure requirement contained in a sponsoring agency rule of general applicability is deemed to meet the minimum threshold of ten or more persons. Second, OGE does anticipate possible limited use of these forms during the forthcoming three-year period 2016–2019. Therefore, the estimated burden figures, representing branchwide implementation of the forms, will remain the same as previously reported by OGE in its prior first and second round paperwork renewal notice for the trust forms (77 FR 76293–76294 (December 27, 2012) and 78 FR 40144–40146 (July 3, 2013)). The estimate is based on the amount of time imposed on a trust administrator or private representative.

<table>
<thead>
<tr>
<th>Type of Document</th>
<th>Total Annual Cost Burden</th>
<th>Avg. Annual Cost Burden per Model</th>
</tr>
</thead>
<tbody>
<tr>
<td>Trust Certificates:</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Certificate of Independence: Total filers (executive branch): 5; private citizen filers (100%): 5; private citizen burden hours (20 minutes/certificate): 2.</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>
| Certificate of Compliance: Total filers (executive branch): 10; private citizen filers (100%): 10; private citizen burden hours (20 minutes/certificate): 3; and ii. Model Qualified Trust Documents: A. Blind Trust Communications: Total users (executive branch): 5; private citizen users (100%): 5; communications documents (private citizens): 25 (based on an average of five communications per user, per year); private citizen burden hours (20 minutes/communication): 8. B. Model Qualified Blind Trust: Total users (executive branch): 2; private citizen users (100%): 2; private citizen burden hours (100 hours/model): 200. C. Model Qualified Diversified Trust: Total users (executive branch): 1; private citizen users (100%): 1; private citizen burden hours (100 hours/model): 100. D. –H. Of the five remaining model qualified trust documents: Total users (executive branch): 2; private citizen users (100%): 2; private citizen burden hours (100 hours/model): 200. I. –J. Of the two model confidentiality agreements: Total users (executive branch): 1; private citizen users (100%): 1; private citizen burden hours (50 hours/agreement): 50.

However, the total annual reporting hour burden on filers themselves is zero and not the 563 hours estimated above because OGE’s estimating methodology reflects the fact that all respondents hire private trust administrators or other private representatives to set up and maintain the qualified blind and diversified trusts. Respondents themselves, typically incoming private citizen Presidential nominees, therefore incur no hour burden. The estimated total annual cost burden to respondents resulting from the collection of information is $1,000,000. Those who use the model documents for guidance are private trust administrators or other private representatives hired to set up and maintain the qualified blind and diversified trusts of executive branch officials who seek to establish such qualified trusts. The cost burden figure is based primarily on OGE’s knowledge of the typical trust administrator fee structure (an average of 1 percent of total assets) and OGE’s experience with administration of the qualified trust program. The $1,000,000 annual cost figure is based on OGE’s estimate of an average of five possible active trusts anticipated to be under administration for each of the next three years with combined total assets of $100,000,000. However, OGE notes that the $1,000,000 figure is a cost estimate for the overall administration of the trusts, only a portion of which relates to information collection and reporting. For want of a precise way to break out the costs directly associated with information collection, OGE is continuing to report to OMB the full $1,000,000 estimate for paperwork clearance purposes.

On March 4, 2016, OGE published a first round notice of its intent to request paperwork clearance for the proposed unmodified qualified trust certificates and modified model trust documents. See 81 FR 11566–11567. OGE did not receive any responses to that notice. In this second notice, public comment is again invited on the model qualified trust certificates and model trust documents, and underlying regulatory provisions, as set forth in this notice, including specific views on the need for and practical utility of this set of collections of information, the accuracy of OGE’s burden estimate, the potential for enhancement of quality, utility and clarity of the information collected, and the minimization of burden (including the use of information technology).

Comments received in response to this notice will be summarized for, and may be included with, the OGE request for extension of the OMB paperwork approval for the set of the various existing qualified trust model certificates, the model communications package, and the model trust documents. The comments will also become a matter of public record.

Approved: May 27, 2016.

Walter M. Shaub, Jr.
Director, Office of Government Ethics.

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Informational Meeting: The Importation and Exportation of Infectious Biological Agents, Infectious Substances and Vectors; Public Webcast

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

ACTION: Notice of public webcast.

SUMMARY: The Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS) is hosting a public webcast to address import and export permit regulations for infectious biological agents, infectious substances, and vectors; and import and export permit exemptions. Presenters for this webcast will include representatives from the U.S. Department of Transportation (DOT), United States Department of Agriculture (USDA) Animal and Plant Health Inspection Service (APHIS), CDC Division of Global Migration and Quarantine, U.S. Customs and Border Protection, U.S. Department of Commerce, U.S. Food and Drug Administration, HHS/Office of the Assistant Secretary for Preparedness and Response/Biomedical Advanced Research and Development Authority (BARDA), U.S. Fish and Wildlife
Service, and the Public Health Agency of Canada.

DATES: The webcast will be held over two days, August 3, 2016 from 12 p.m. to 4 p.m. EDT and August 4, 2016 from 12:00 p.m. to 4:00 p.m. Registration instructions are found on the HHS/CDC Import Permit Program Web site, http://www.cdc.gov/od/eaipp/importApplication/agents.htm.

ADDITIONS: The webcast will be broadcast from the Centers for Disease Control and Prevention, 1600 Clifton Road NE., Atlanta, Georgia 30329.

FOR FURTHER INFORMATION CONTACT: Von McClee, Division of Select Agents and Toxins, Office of Public Health Preparedness and Response, Centers for Disease Control and Prevention, 1600 Clifton Road NE., MS A–46, Atlanta, GA 30333; phone: 404–718–2000; email: lrsa@cdc.gov.

SUPPLEMENTARY INFORMATION: This webcast is an opportunity for all interested parties (e.g., academic institutions and biomedical centers, commercial manufacturing facilities, federal, state, and local laboratories, including clinical and diagnostic laboratories, research facilities, exhibition facilities, and educational facilities) to obtain specific guidance and information regarding import and export permit regulations. The webcast will also provide assistance to those interested in applying for an import or export permit (or license) from federal agencies within the United States. Instructions for registration are found on the HHS/CDC Import Permit Program Web site, http://www.cdc.gov/od/eaipp/importApplication/agents.htm.

Participants must register by July 15, 2016. This is a webcast only event and there will be no on-site participation at the HHS/CDC broadcast facility.

Dated: May 27, 2016.

Veronica Kennedy,
Acting Executive Secretary, Centers for Disease Control and Prevention.

[FR Doc. 2016–13053 Filed 6–1–16; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Community Living

Agency Information Collection Activities; Submission for OMB Review; Comment Request; University Centers for Excellence in Developmental Disabilities Education, Research, and Service—Annual Report

AGENCY: Administration for Community Living, HHS.

ACTION: Notice.

SUMMARY: The Administration on Intellectual and Developmental Disabilities (AIDD), Administration for Community Living (ACL) is announcing an opportunity to comment on the proposed collection of information by the agency. Under the Paperwork Reduction Act of 1995 (the PRA), Federal agencies are required to publish notice in the Federal Register concerning each proposed collection of information, including each proposed extension of an existing collection of information, and to allow 60 days for public comment in response to the notice. This notice collects comments on the information collection requirements relating to the continuation of an existing collection for University Centers for Excellence in Developmental Disabilities Education, Research, and Service.

DATES: Submit written comments on the collection of information by August 1, 2016.

ADDRESSES: Submit written comments on the collection of information by email to Valerie.Bond@acl.hhs.gov.

FOR FURTHER INFORMATION CONTACT: Valerie Bond by email at Valerie.Bond@acl.hhs.gov or 202.795–7311.

SUPPLEMENTARY INFORMATION: Section 104 (42 U.S.C. 15004) of the Developmental Disabilities Assistance and Bill of Rights Act of 2000 (DD Act of 2000) directs the Secretary of Health and Human Services to develop and implement a system of program accountability to monitor the grantees funded under the DD Act of 2000. The program accountability system shall include the National Network of University Centers for Excellence in Developmental Disabilities Education, Research, and Service (UCEDDs) authorized under Part D of the DD Act of 2000. In addition to the accountability system, Section 154(e) (42 U.S.C. 15064) of the DD Act of 2000 includes requirements for a UCEDD Annual Report. ACL estimates the burden of this collection of information as follows:

Annual Burden Estimates

<table>
<thead>
<tr>
<th>Instrument</th>
<th>Number of respondents</th>
<th>Number of responses per respondent</th>
<th>Average burden hours per response</th>
<th>Total burden hours</th>
</tr>
</thead>
<tbody>
<tr>
<td>UCEDD Annual Report</td>
<td>67</td>
<td>1</td>
<td>1,412</td>
<td>94,604</td>
</tr>
</tbody>
</table>

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Community Living

Agency Information Collection Activities; Submission for OMB Review; Comment Request; Senior Medicare Patrol (SMP) Program Outcome Measurement

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 July 5, 2016.

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.

FOR FURTHER INFORMATION CONTACT: Phillip McKoy at 202.795.7397 or email: phillip.mckoy@acl.hhs.gov.