

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>.

ADDRESSES: 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: lsrat@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]

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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

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
UCEDD Annual Report	67	1	1,412	94,604

Estimated Total Annual Burden Hours: 94,604.

Dated: May 25, 2016.

Kathy Greenlee,

Administrator & Assistant Secretary for Aging.

[FR Doc. 2016-13020 Filed 6-1-16; 8:45 am]

BILLING CODE 4154-01-P

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.

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. Grantees are required by Congress to provide information for use in program monitoring and for Government Performance and Results Act (GPRA) purposes. This information collection reports the number of active volunteers, issues and inquiries received, other SMP program outreach activities, and the number of Medicare dollars recovered, among other SMP performance outcomes. This information is used as the primary method for monitoring the SMP Projects. ACL estimates the burden of this collection of information as follows: Respondents: 54 SMP grantees at 23 hours per month (276 hours per year, per grantee). Total Estimated Burden Hours: 7,452 hours per year.

Dated: May 25, 2016.

Kathy Greenlee,
Administrator and Assistant Secretary for Aging.

[FR Doc. 2016-12868 Filed 6-1-16; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA-2016-N-0001]

Collaborating To Strengthen Food, Drug, and Medical Device Safety Systems; Notice of Conference

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of conference.

SUMMARY: The Food and Drug Administration (FDA) Philadelphia District Office, in co-sponsorship with the Association of Food and Drug Officials (AFDO), and the North Central Association of Food and Drug Officials, is announcing a conference entitled "Collaborating to Strengthen Food, Drug, and Medical Device Safety Systems." This conference is intended to provide information about FDA drug and device regulation to the regulated industry.

DATES: The conference will be held on June 25 to June 29, 2016. See

SUPPLEMENTARY INFORMATION for meeting times.

ADDRESSES: The Omni William Penn Hotel, 530 William Penn Pl., Pittsburgh, PA 15219. Attendees are responsible for their own accommodations.

FOR FURTHER INFORMATION CONTACT:

Randy Young, Association of Food and Drug Officials, 2550 Kingston Rd., Suite 311, York, PA 17402, 717-757-2888, FAX: 717-650-3650, ryoung@afdo.org.

SUPPLEMENTARY INFORMATION: FDA has made education of the food, feed, drug, and device manufacturing community a high priority to help ensure the quality of FDA-regulated products. The conference helps to achieve objectives set forth in section 406 of the Food and Drug Administration Modernization Act of 1997 (21 U.S.C. 393), which includes working closely with stakeholders and maximizing the availability and clarity of information for stakeholders and the public. The conference also is consistent with the Small Business Regulatory Enforcement Fairness Act of 1996 (Pub. L. 104-121), as outreach activities by government Agencies to small businesses.

The conference helps fulfill the U.S. Department of Health and Human Services' and FDA's important mission to protect the public health. The conference will provide FDA-regulated drug and device entities with information on a number of topics concerning FDA requirements related to the production and marketing of drugs and/or devices. Topics for discussion include, but are not limited to the following:

- FDA Program Alignment
- Recalls from the Perspective of the District
- Inspection of Licensed Producers under the Marijuana for Medical Purposes Regulations (Health Canada)
- Foreign inspections
- Regulatory Intelligence
- FDA Inspections: Challenges and Opportunities (Working Luncheon)
- Drug Shortages
- Drug Supply Chain Act: Wholesale Drug Distributor and 3rd Party Logistics Provider
- Medical Device Single Audit Program
- Compliance Questions Panel

The Conference Web site is: <http://afdo.org/conference>. The meeting times are as follows:

Date	Meeting time
June 25	8 a.m. to 5 p.m.
June 26	8 a.m. to 6 p.m.
June 27	8 a.m. to 5:30 p.m.
June 28	8 a.m. to 5 p.m.
June 29	8 a.m. to 11:30 a.m.

Registration: The AFDO registration fees cover the cost of facilities, materials, and breaks. Seats are limited and registration will close after the course is filled; therefore, please submit your registration as soon as possible.

Conference space will be filled in order of receipt of registration; those accepted will receive confirmation. Registration at the site is not guaranteed but may be possible on a space available basis on the day of the conference, beginning at 7:30 a.m. The cost of registration follows:

Category	Cost of registration
Member	\$475
Non-Member	575
Additional Fee for Registration Postmarked After June 1, 2016	100

To register, please complete and submit an AFDO conference registration form, available at <http://pitt.afdo.org/registration.html>, along with a check, money order payable to "AFDO"; the registrar will also accept Visa and MasterCard credit cards. Please mail your completed registration form and payment to: AFDO, 2550 Kingston Rd., Suite 311, York, PA 17402. To register online, please visit <http://pitt.afdo.org/registration.html> (FDA has verified the Web site address, but is not responsible for subsequent changes to the Web site after this document publishes in the **Federal Register**.) For more information on the conference, or for questions about registration, please contact Randy Young (see **FOR FURTHER INFORMATION CONTACT**), email inquiries will also be accepted at afdo@afdo.org, or visit <http://www.afdo.org>.

If you need special accommodations due to a disability, please contact Randy Young (see **FOR FURTHER INFORMATION CONTACT**) at least 7 days in advance of the conference.

Dated: May 26, 2016.

Leslie Kux,
Associate Commissioner for Policy.

[FR Doc. 2016-12942 Filed 6-1-16; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA-2016-P-0378]

Determination That TRIVARIS (Triamcinolone Acetonide) Injectable Suspension, 80 Milligrams/Milliliters, Was Not Withdrawn From Sale for Reasons of Safety or Effectiveness

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA or Agency) has