

tool for monitoring of the P&As, including the public input requirement. Furthermore, it will provide an overview of program priorities, and permit AIDD to track accomplishments against goals, permitting the formulation of technical assistance and compliance with the Developmental Disabilities Assistance and Bill of Rights Act of 2000.

**DATES:** Submit written comments on the collection of information by August 10, 2015.

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

**FOR FURTHER INFORMATION CONTACT:** Valerie Bond, Administration on Community Living, Administration on Intellectual and Developmental Disabilities, Office of Program Support,

One Massachusetts Avenue NW., Room 4302, Washington, DC 20201, 202-690-5841.

**SUPPLEMENTARY INFORMATION:** In compliance with the requirements of section 506 (c)(2)(A) of the Paperwork Reduction Act of 1995, the Administration on Community Living is soliciting public comment on the specific aspects of the information collection described above. Copies of the proposed collection of information can be obtained and comments may be forwarded by writing to: Valerie Bond, Administration on Community Living, Administration on Intellectual and Developmental Disabilities, Office of Program, One Massachusetts Avenue, NW., Room 4302, Washington, DC 20201.

The Department specifically requests comments on: (a) Whether the proposed

Collection of information is necessary for the proper performance of the function of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) the quality, utility, and clarity of the information to be collected; (d) ways to minimize the burden information to be collected; and (e) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection technique comments and or other forms of information technology. Consideration will be given to comments and suggestions submitted within 60 days of this publication.

*Respondents:* 57 Protection and Advocacy Systems

**ANNUAL BURDEN ESTIMATES**

| Instrument   | Number of respondents | Number of responses per respondent | Average burden hours per response | Total burden hours |
|--|-----------------------|------------------------------------|-----------------------------------|--------------------|
| State Developmental Disabilities Council 5-Year State Plan ..... | 57                    | 1                                  | 44                                | 2,508              |

*Estimated Total Annual Burden Hours:* 2,508.

Dated: June 3, 2015.

**Kathy Greenlee,**  
*Administrator and Assistant Secretary for Aging.*

[FR Doc. 2015-14050 Filed 6-8-15; 8:45 am]

**BILLING CODE 4154-01-P**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Food and Drug Administration**

[Docket No. FDA-2014-N-0373]

**Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Risk and Benefit Perception Scale Development**

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Notice.

**SUMMARY:** The Food and Drug Administration (FDA) is announcing that a collection of information entitled "Risk and Benefit Perception Scale Development" has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

**FOR FURTHER INFORMATION CONTACT:** FDA PRA Staff, Office of Operations, Food and Drug Administration, 8455

Colesville Rd., COLE-14526, Silver Spring, MD 20993-0002, *PRAStaff@fda.hhs.gov*.

**SUPPLEMENTARY INFORMATION:** On November 28, 2015, the Agency submitted a proposed collection of information entitled "Risk and Benefit Perception Scale Development" to OMB for review and clearance under 44 U.S.C. 3507. An Agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. OMB has now approved the information collection and has assigned OMB control number 0910-0784. The approval expires on May 31, 2018. A copy of the supporting statement for this information collection is available on the Internet at <http://www.reginfo.gov/public/do/PRAMain>.

Dated: June 3, 2015.

**Leslie Kux,**  
*Associate Commissioner for Policy.*

[FR Doc. 2015-14027 Filed 6-8-15; 8:45 am]

**BILLING CODE 4164-01-P**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Food and Drug Administration**

[Docket No. FDA-2011-N-0672]

**Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Prominent and Conspicuous Mark of Manufacturers on Single-Use Devices**

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Notice.

**SUMMARY:** The Food and Drug Administration (FDA) is announcing that a collection of information entitled, "Prominent and Conspicuous Mark of Manufacturers on Single-Use Devices" has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

**FOR FURTHER INFORMATION CONTACT:** FDA PRA Staff, Office of Operations, Food and Drug Administration, 8455 Colesville Rd., COLE-14526, Silver Spring, MD 20993-0002, *PRAStaff@fda.hhs.gov*.

**SUPPLEMENTARY INFORMATION:** On April 23, 2015, the Agency submitted a proposed collection of information entitled, "Prominent and Conspicuous Mark of Manufacturers on Single-Use Devices" to OMB for review and clearance under 44 U.S.C. 3507. An