Ryan (*Caroline.Ryan@acl.hhs.gov*) to register. Registrants will receive confirmation after they have been accepted.

Webcast of the Public Workshop: This public workshop will also be available by webcast. Those interested in joining the webcast must register no later than October 14, 2014. Early registration is recommended; conference lines are limited. To register for the webcast, please visit: *https://attendee.gotowebinar.com/register/2329630204976253953.* Call details will be sent to all registered participants after October 14, 2014.

Requests for Public Comment: This public workshop will include topic-focused sessions with opportunities for public comment in-person or via webcast. The agencies will do their best to accommodate requests to make public comments. Individuals and organizations with common interests are urged to consolidate or coordinate their comments.

Comments: The agencies seek broad input from stakeholders and experts on the use of health information technology (health IT) for the purposes of developing an integrated, person-centered plan, including how to improve communication and collaboration among community-based organizations and health care partners. To obtain broad public comment, the Agencies are soliciting comments on all aspects of the public workshop topics.

The deadline for submitting comments related to this public workshop is October 31, 2014.

Please submit comments to: *personcenteredhealthIT@hhs.gov*

Transcript: The meeting will be transcribed. A transcript and meeting materials will be posted on the healthit.gov Web site at: *http://healthitgov-stage.ahrqstg.org/person-centered-care.*

Dated: October 1, 2014.

Jacob Reider,

Deputy National Coordinator for Health Information Technology.

[FR Doc. 2014–23931 Filed 10–7–14; 8:45 am]

BILLING CODE 4150–45–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2011–N–0627]

Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; General Administrative Procedures: Citizen Petitions; Petition for Reconsideration or Stay of Action; Advisory Opinions

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a collection of information entitled ''General Administrative Procedures: Citizen Petitions; Petition for Reconsideration or Stay of Action; Advisory Opinions'' 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 June 12, 2014, the Agency submitted a proposed collection of information entitled ''General Administrative Procedures: Citizen Petitions; Petition for Reconsideration or Stay of Action; Advisory Opinions'' 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 0910–0183. The approval expires on August 31, 2017. A copy of the supporting statement for this information collection is available on the Internet at *http://www.reginfo.gov/public/do/PRAMain.*

Dated: October 2, 2014.

Leslie Kux,

Assistant Commissioner for Policy.

[FR Doc. 2014–24051 Filed 10–7–14; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2010–N–0493]

Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Additional Criteria and Procedures for Classifying Over-the-Counter Drugs as Generally Recognized as Safe and Effective and Not Misbranded

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a collection of information entitled ''Additional Criteria and Procedures for Classifying Over-the-Counter Drugs as Generally Recognized as Safe and Effective and Not Misbranded'' 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 June 25, 2014, the Agency submitted a proposed collection of information entitled ''Additional Criteria and Procedures for Classifying Over-the-Counter Drugs as Generally Recognized as Safe and Effective and Not Misbranded'' 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 0910–0688. The approval expires on August 31, 2017. A copy of the supporting statement for this information collection is available on the Internet at *http://www.reginfo.gov/public/do/PRAMain.*

Dated: October 2, 2014.

Leslie Kux,

Assistant Commissioner for Policy.

[FR Doc. 2014–23956 Filed 10–7–14; 8:45 am]

BILLING CODE 4164–01–P