

IV. Paperwork Reduction Act of 1995

This guidance refers to currently approved collections of information found in FDA regulations. These collections of information are subject to review by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995 (44 U.S.C. 3501–3520) (the PRA). The collections of information in 21 CFR part 801 are approved under OMB control number 0910–0485 and the collections of information in 21 CFR part 610, subpart G, are approved under OMB control number 0910–0338.

The labeling provisions recommended in this guidance are not subject to review by OMB because they do not constitute a “collection of information” under the PRA. Rather, the recommended labeling is a “public disclosure of information originally supplied by the Federal Government to the recipient for the purpose of disclosure to the public” (5 CFR 1320.3(c)(2)).

V. Comments

Interested persons may submit either electronic comments regarding this document to <http://www.regulations.gov> or written comments to the Division of Dockets Management (see **ADDRESSES**). It is only necessary to send one set of comments. Identify comments with the docket number found in brackets in the heading of this document. Received comments may be seen in the Division of Dockets Management between 9 a.m. and 4 p.m., Monday through Friday, and will be posted to the docket at <http://www.regulations.gov>.

Dated: November 25, 2014.

Leslie Kux,

Associate Commissioner for Policy.

[FR Doc. 2014–28265 Filed 12–1–14; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2014–N–1936]

Establishment of a Public Docket; Electronic Cigarettes and the Public Health Workshop

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; establishment of docket; request for data, information, and comments.

SUMMARY: The Food and Drug Administration (FDA), Center for Tobacco Products, is establishing a

public docket in conjunction with the first public workshop to gather scientific information about electronic cigarettes (e-cigarettes) as announced in Docket No. FDA–2014–N–0001–0079.

Regardless of attendance at the public workshop, interested parties are invited to submit comments, supported by research and data, regarding electronic cigarettes and the public health.

DATES: Submit written or electronic comments by April 15, 2015.

ADDRESSES: Submit electronic comments to <http://www.regulations.gov>. Submit written comments to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852. Identify comments with the docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Caryn Cohen, Office of Science, Center for Tobacco Products, Food and Drug Administration, Document Control Center, Bldg. 71, Rm. G335, 10903 New Hampshire Ave., Silver Spring, MD 20993–0002, 1–877–287–1373, email: workshop.CTPOS@fda.hhs.gov.

I. Background

On September 17, 2014, FDA announced a public workshop to gather information about e-cigarettes and the public health (Electronic Cigarettes and the Public Health; Public Workshop; 79 FR 55815, September 17, 2014, Docket No. FDA–2014–N–0001). The focus of the workshop is product science (specifically device designs and characteristics, and e-liquid and aerosol constituents), product packaging, constituent labeling, and environmental impact. FDA intends to follow the first workshop with two additional e-cigarette workshops; one on individual health effects and one on population health effects. As stated in the **Federal Register** notice of the public workshop, the workshops are not intended to inform the Agency’s deeming rulemaking. The workshops are intended to better inform FDA about these products. Should the Agency move forward as proposed to regulate e-cigarettes, additional information about the products would assist the Agency in carrying out its responsibilities under the law.

II. Submission of Comments

Regardless of attendance at the public workshop, interested parties are invited to submit comments, supported by research and data, regarding e-cigarettes and the public health. Information related to workshop presentations and

discussion topics, including specific questions to be addressed at the workshop, can be found at <http://www.fda.gov/TobaccoProducts/NewsEvents/ucm238308.htm>.

Interested persons may submit either electronic comments to this docket at <http://www.regulations.gov> or written comments to the Division of Dockets Management (see **ADDRESSES**). It is only necessary to send one set of comments. Identify comments with the docket number found in brackets in the heading of this document. Received comments may be seen in the Division of Dockets Management between 9 a.m. and 4 p.m., Monday through Friday, and will be posted to the docket at <http://www.regulations.gov>.

Dated: November 25, 2014.

Leslie Kux,

Associate Commissioner for Policy.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Agency Information Collection Activities: Submission to OMB for Review and Approval; Public Comment Request

AGENCY: Health Resources and Services Administration, HHS.

ACTION: Notice.

SUMMARY: In compliance with Section 3507(a)(1)(D) of the Paperwork Reduction Act of 1995, the Health Resources and Services Administration (HRSA) has submitted an Information Collection Request (ICR) to the Office of Management and Budget (OMB) for review and approval. Comments submitted during the first public review of this ICR will be provided to OMB. OMB will accept further comments from the public during the review and approval period.

DATES: Comments on this ICR should be received no later than January 2, 2015.

ADDRESSES: Submit your comments, including the Information Collection Request Title, to the desk officer for HRSA, either by email to OIRA_submission@omb.eop.gov or by fax to 202–395–5806.

FOR FURTHER INFORMATION CONTACT: To request a copy of the clearance requests submitted to OMB for review, email the HRSA Information Collection Clearance Officer at paperwork@hrsa.gov or call (301) 443–1984.