Partial Withdrawal of Previous Notice

This document also withdraws a portion of a prior notice concerning the same CMS–10261–specific subject matter.

Specifically, on page 51276, in the second column, in the second paragraph, information collection CMS–10261 (OMB Control Number 0938–1054) that published in the Federal Register on August 24, 2015 (80 FR 51275) is hereby withdrawn.

Dated: September 15, 2015.

William N. Parham, III,
Director, Paperwork Reduction Staff, Office of Strategic Operations and Regulatory Affairs.

[FR Doc. 2015–23482 Filed 9–17–15; 8:45 am]

BILLING CODE 4120–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2015–N–3230]

Consumer Comments—Public Posting and Availability of Comments Submitted to Food and Drug Administration Dockets

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA or Agency) is changing the Agency’s long standing practice of not publicly posting on http://www.regulations.gov comments submitted by individuals in their individual capacity.

I. Background

Historically, FDA generally has not publicly posted comments submitted by individuals in their individual capacity (and not on behalf of an organization, corporation, or other entity). For comments submitted through http://www.regulations.gov, for example, such comments are identified as “Individual Consumer” under the field titled “Category (Required)” on the “Your Information” page. This non-posting practice has applied only to individual consumer comments which otherwise would be displayed on http://www.regulations.gov. These comments have been placed in the official FDA docket and are publicly available in FDA’s Reading Room or through Freedom of Information Act requests and have been considered by the Agency in finalizing its regulatory actions.

FDA is changing this practice and will post such consumer comments on http://www.regulations.gov, as it posts other comments. FDA has made this change so that its public dockets better serve their purpose of promoting transparency and the sharing of information.

In 1995, FDA explained that it routinely reviewed all comments for obvious confidential information before placing the comments in the docket (60 FR 66982), but this practice is no longer feasible given factors such as the volume of comments FDA receives and the adoption of a government-wide electronic portal system for submitting and posting comments at http://www.regulations.gov. FDA developed the practice of not posting individual consumer comments largely because of concerns about disclosing personal information of individuals who may not have realized, when submitting their comments, that their name, address, and other identifying information would be publicly viewable. This public viewability became more obvious as the Internet gained popularity and particularly when FDA dockets system was merged with the government-wide portal system for submission of all public comments on government regulatory actions at http://www.regulations.gov in 2007. This change so that its public dockets better serve their purpose of promoting transparency and the sharing of information.

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FDA selected “individual consumer” comments for non-posting because of previous concerns raised by individuals and the conclusion that such commenters may not be as familiar with the regulatory process and the public nature of dockets as are other entities, such as regulated industry.

In recent years, FDA has occasionally made exceptions to this non-posting practice, typically using the COMMENTS section in a particular Federal Register document to alert the public that all comments were subject to public posting. FDA Federal Register documents, requesting or providing for the submission of comments, published subsequent to this notice will contain new instructions and information concerning the posting of comments submitted to that particular docket.

This change fulfills a recommendation from the 2010 FDA Transparency Initiative and aligns with a 2013 recommendation from the Administrative Conference of the United States that “all agencies should manage their public rulemaking dockets to achieve maximum public disclosure” consistent with legal limitations and other claims of privilege. It also furthers an objective in Executive Order 13563, which directs Agencies to base their regulations on “public participation and an open exchange of ideas.”

II. Consumer Comments and Confidential Information

The commenter is solely responsible for ensuring that the submitted comment does not include any confidential information that the commenter or a third party may not wish to be posted, such as private medical information, the commenter’s or anyone else’s Social Security number, or confidential business information, such as a manufacturing process. If a name, contact information, or other information that identifies the commenter is included in the body of the submitted comment, that information will be posted on http://www.regulations.gov. FDA will post comments, as well as any attachments submitted electronically, on http://www.regulations.gov, along with the State/Province and country (if provided), the name of the commenter’s representative (if any), and the category selected to identify the commenter (e.g., individual, consumer, academic, industry).

The Agency expects that only in exceptional instances would a comment need to include private, personal, or confidential information. If a comment is submitted with confidential information that the commenter does not wish to be made available to the public, the comment would be submitted as a written/paper submission and in the manner detailed in the applicable Federal Register document. For written/paper comments submitted containing confidential information, FDA will post the redacted/blacked out version of the comment including any attachments submitted by the commenter. The unredacted copy will not be posted, assuming the commenter follows the instructions in the applicable Federal Register document. Any information marked as confidential will not be disclosed except in accordance with § 10.20 (21 CFR 10.20) and other applicable disclosure law.

FDA will include new information and standard instructions for submitting comments in all Federal Register documents requesting or providing for the submission of comments. The instructions will explain how to submit comments to the docket on that particular document via electronic means and also will explain the process for submission of comments, in written/paper format, that the commenter wishes to mark as confidential.

III. Date of Implementation

All comments submitted electronically through http://www.regulations.gov to any FDA docket, existing or new, after October 15, 2015, will be posted to the applicable docket and publicly viewable on http://www.regulations.gov. All comments submitted by mail or delivery to the Division of Dockets Management in written/paper format to any FDA docket, existing or new, after October 15, 2015, will be posted to the applicable docket and publicly viewable on http://www.regulations.gov unless submitted under the following conditions: (1) The written/paper submission is marked as confidential, and (2) the submitter provides an unredacted and a redacted version; the redacted version must have the information claimed as confidential redacted/blacked out. If submitted under these conditions, then the redacted/blacked out written/paper submission will be posted publicly on http://www.regulations.gov, except as otherwise provided by § 10.20 or other law.

Dated: September 14, 2015.

Leslie Kux,
Associate Commissioner for Policy.

[FR Doc. 2015–23389 Filed 9–17–15; 8:45 am]
BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES
Office of the Secretary


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

AGENCY: Office of the Secretary, HHS.

ACTION: Notice.

SUMMARY: In compliance with section 3507(a)(1)(D) of the Paperwork Reduction Act of 1995, the Office of the Secretary (OS), Department of Health and Human Services, has submitted an Information Collection Request (ICR), described below, to the Office of Management and Budget (OMB) for review and approval. The ICR is for renewal of the approved information collection assigned OMB control number 0990–0279, scheduled to expire on September 30, 2015. Comments submitted during the first public review of this ICR will be provided to OMB. OMB will accept further comments from the public on this ICR during the review and approval period.

DATES: Comments on the ICR must be received on or before October 19, 2015.

ADDRESSES: Submit your comments to OIRA_submission@omb.eop.gov or via facsimile to (202) 395–5806.

FOR FURTHER INFORMATION CONTACT: Information Collection Clearance staff, InformationCollectionClearance@hhs.gov or (202) 690–6162.

SUPPLEMENTARY INFORMATION: When submitting comments or requesting information, please include the OMB control number 0990–0279 and document identifier HHS–OS–30D for reference.

Information Collection Request Title: Institutional Review Board Form—OMB No. 0990–0279, Assistant Secretary for Health, Office for Human Research Protections.

Abstract: Section 491(a) of Public Law 99–158 states that the Secretary of HHS shall by regulation require that each entity applying for HHS support (e.g., a grant, contract, or cooperative agreement)