

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA-2007-D-0372]

Agency Information Collection Activities; Proposed Collection; Comment Request; Adverse Event Reporting and Recordkeeping for Dietary Supplements as Required by the Dietary Supplement and Nonprescription Drug Consumer Protection Act

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA or we) is announcing an opportunity for public comment on the proposed collection of certain information by our 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. This notice solicits comments on the collection of information associated with adverse event reporting and recordkeeping for dietary supplements as required by the Dietary Supplement and Nonprescription Drug Consumer Protection Act (the DSNDCPA).

DATES: Submit either electronic or written comments on the collection of information by December 21, 2015.

ADDRESSES: You may submit comments as follows:

Electronic Submissions

Submit electronic comments in the following way:

- *Federal eRulemaking Portal:* <http://www.regulations.gov>. Follow the instructions for submitting comments. Comments submitted electronically, including attachments, to <http://www.regulations.gov> will be posted to the docket unchanged. Because your comment will be made public, you are solely responsible for ensuring that your comment does not include any confidential information that you or a third party may not wish to be posted, such as medical information, your or anyone else's Social Security number, or confidential business information, such as a manufacturing process. Please note that if you include your name, contact information, or other information that identifies you in the body of your comments, that information will be posted on <http://www.regulations.gov>.

- If you want to submit a comment with confidential information that you do not wish to be made available to the public, submit the comment as a written/paper submission and in the manner detailed (see "Written/Paper Submissions" and "Instructions").

Written/Paper Submissions

Submit written/paper submissions as follows:

- *Mail/Hand delivery/Courier (for written/paper submissions):* Division of Dockets Management (HFA-305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.
- For written/paper comments submitted to the Division of Dockets Management, FDA will post your comment, as well as any attachments, except for information submitted, marked and identified, as confidential, if submitted as detailed in "Instructions."

Instructions: All submissions received must include the Docket No. FDA-2007-D-0372 for the information collection request entitled, "Adverse Event Reporting and Recordkeeping for Dietary Supplements as Required by the DSNDCPA."

Received comments will be placed in the docket and, except for those submitted as "Confidential Submissions," publicly viewable at <http://www.regulations.gov> or at the Division of Dockets Management between 9 a.m. and 4 p.m., Monday through Friday.

- **Confidential Submissions**—To submit a comment with confidential information that you do not wish to be made publicly available, submit your comments only as a written/paper submission. You should submit two copies total. One copy will include the information you claim to be confidential with a heading or cover note that states "THIS DOCUMENT CONTAINS CONFIDENTIAL INFORMATION." The Agency will review this copy, including the claimed confidential information, in its consideration of comments. The second copy, which will have the claimed confidential information redacted/blacked out, will be available for public viewing and posted on <http://www.regulations.gov>. Submit both copies to the Division of Dockets Management. If you do not wish your name and contact information to be made publicly available, you can provide this information on the cover sheet and not in the body of your comments and you must identify this information as "confidential." Any information marked as "confidential" will not be disclosed except in accordance with 21 CFR 10.20 and other

applicable disclosure law. For more information about FDA's posting of comments to public dockets, see 80 FR 56469, September 18, 2015, or access the information at: <http://www.fda.gov/regulatoryinformation/dockets/default.htm>.

Docket: For access to the docket to read background documents or the electronic and written/paper comments received, go to <http://www.regulations.gov> and insert the docket number, found in brackets in the heading of this document, into the "Search" box and follow the prompts and/or go to the Division of Dockets Management, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

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

SUPPLEMENTARY INFORMATION: Under the PRA (44 U.S.C. 3501-3520), Federal Agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. "Collection of information" is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes Agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA (44 U.S.C. 3506(c)(2)(A)) requires Federal Agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, we are publishing this notice of the proposed collection of information set forth in this document.

With respect to the following collection of information, we invite comments on these topics: (1) Whether the proposed collection of information is necessary for the proper performance of FDA's functions, including whether the information will have practical utility; (2) the accuracy of FDA's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques, when appropriate, and other forms of information technology.

Adverse Event Reporting and Recordkeeping for Dietary Supplements as Required by the DSNDCPA, 21 U.S.C. 379aa–1(b)(1) OMB Control Number 0910–0635—Extension

The DSNDCPA (Pub. L. 109–462) amends the Federal Food, Drug, and Cosmetic Act (the FD&C Act) with respect to serious adverse event reporting and recordkeeping for dietary supplements and non-prescription drugs marketed without an approved application. Section 761(b)(1) of the FD&C Act (21 U.S.C. 379aa–1(b)(1)) requires the manufacturer, packer, or distributor whose name under section 403(e)(1) of the FD&C Act (21 U.S.C. 343(e)(1)) appears on the label of a dietary supplement marketed in the United States to submit to us all serious adverse event reports associated with the use of a dietary supplement, accompanied by a copy of the product label. The manufacturer, packer, or distributor of a dietary supplement is required by the DSNDCPA to use the MedWatch form (Form FDA 3500A) when submitting a serious adverse event report to FDA. In addition, under section 761(c)(2) of the FD&C Act, the

submitter of the serious adverse event report (referred to in the statute as the “responsible person”) is required to submit to FDA a follow up report of any related new medical information the responsible person receives within 1 year of the initial report. Section 761(e)(1) of the FD&C Act requires that responsible persons maintain records related to the dietary supplement adverse event reports they receive, whether or not the adverse event is serious. Under the statute, the records must be retained for a period of 6 years. As required by section 3(d)(3) of the DSNDCPA, we issued guidance to describe the minimum data elements for serious adverse event reports for dietary supplements. In the **Federal Register** of July 14, 2009 (74 FR 34024), we announced the availability of guidance entitled “Guidance for Industry: Questions and Answers Regarding Adverse Event Reporting and Recordkeeping for Dietary Supplements as Required by the Dietary Supplement and Nonprescription Drug Consumer Protection Act.” The guidance discusses how, when, and where to submit serious

adverse event reports for dietary supplements and followup reports. The guidance also provides our recommendation on records maintenance and access for serious and non-serious adverse event reports and related documents. The guidance recommends that the responsible person document their attempts to obtain the minimum data elements for a serious adverse event report. Along with these records, the guidance recommends that the responsible person keep the following other records: (1) Communications between the responsible person and the initial reporter of the adverse event and between the responsible person and any other person(s) who provided information about the adverse event; (2) the responsible person’s serious adverse event report to us with attachments; (3) any new information about the adverse event received by the responsible person; (4) any reports to us of new information related to the serious adverse event report. We estimate the annual reporting burden of this collection of information as follows:

TABLE 1—ESTIMATED ANNUAL REPORTING BURDEN ¹

21 U.S.C. section	Number of respondents	Number of responses per respondent	Total annual responses	Average burden per response	Total hours
21 U.S.C. 379aa–1(b)(1)—serious adverse event reports for dietary supplements	170	17	2,860	2	5,720
21 U.S.C. 379aa–1(c)(2)—followup reports of new medical information	42	17	715	1	715
Total					6,435

¹ There are no capital costs or operating and maintenance costs associated with this collection of information.

This estimate is based on our experience with similar adverse event reporting programs and the number of serious adverse event reports and followup reports received in the past 3 years. All dietary supplement manufacturers, packers, or distributors are subject to serious adverse event mandatory reporting. We received 2,435 initial serious adverse event reports in fiscal year (FY) 2012, 3,414 in FY2013, and 2,745 in FY2014. We averaged these figures (2,860 rounded to the nearest 10) as a basis for our estimated number of annual reports. We also used an average of the number of firms filing reports (170 rounded to the nearest 10). Finally, we estimate that it will take respondents

an average of 2 hours per report to collect information about a serious adverse event associated with a dietary supplement and report the information to us on Form FDA 3500A. Thus, the estimated burden associated with submitting initial dietary supplement serious adverse event reports is 5,720 hours (2,860 responses × 2 hours) as shown in row 1 of Table 1. If a respondent that has submitted a serious adverse event report receives new information related to the serious adverse event within 1 year of submitting the initial report, the respondent must provide the new information to us in a followup report. We estimate that 25 percent of serious adverse event reports related to dietary

supplements will have a followup report submitted, resulting in approximately 715 followup reports submitted annually (2,860 × 0.25 = 715). Dividing the annual number of reports among the 170 firms reporting results in approximately 17 reports for 42 respondents. We estimate that each followup report will require an hour to assemble and submit, including the time needed to copy and attach the initial serious adverse event report as recommended in the guidance. Thus the estimated burden for followup reports of new information is 715 hours (715 responses × 1 hour) as shown in row 2 of Table 1.

TABLE 2—ESTIMATED ANNUAL RECORDKEEPING BURDEN¹

21 U.S.C. section	Number of recordkeepers	Number of records per recordkeeper	Total annual records	Average burden per recordkeeping	Total hours
Dietary supplement adverse event records (21 U.S.C. 379aa–1(e)(1)).	1,700	74	125,800	0.5 (30 minutes)	62,900

¹ There are no capital costs or operating and maintenance costs associated with this collection of information.

All dietary supplement manufacturers, packers, or distributors are subject to serious adverse event recordkeeping. We estimate that there are 1,700 such respondents, based on the figure 1,460 as provided in our final rule of June 25, 2007 (72 FR 34751) on the “Current Good Manufacturing Practice in Manufacturing, Packaging, Labeling, or Holding Operations for Dietary Supplements,” and factoring a two percent annual growth rate. Estimating that each recordkeeper will keep approximately 74 records per year results in an annual burden of 125,800 records. Estimating that assembling and filing these records, including any necessary photocopying, will take approximately 30 minutes, or 0.5 hours, per record, results in an annual burden of 62,900 hours (125,800 records × 0.5 hours = 62,900 total hours).

Once the documents pertaining to an adverse event report have been assembled and filed in accordance with the safety reporting portal, we expect the records retention burden to be minimal, as we believe most establishments would normally keep this kind of record for at least several years after receiving the report, as a matter of usual and customary business practice.

Dated: October 15, 2015.

Leslie Kux,

Associate Commissioner for Policy.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2015–N–2126]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Evaluation of the Food and Drug Administration’s Campaign To Reduce Tobacco Use Among Lesbian, Gay, Bisexual, and Transgender Young Adults

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a proposed collection of information has been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995.

DATES: Fax written comments on the collection of information by November 20, 2015.

ADDRESSES: To ensure that comments on the information collection are received, OMB recommends that written comments be faxed to the Office of Information and Regulatory Affairs, OMB, Attn: FDA Desk Officer, FAX: 202–395–7285, or emailed to aira_submission@omb.eop.gov. All comments should be identified with the title “Evaluation of the Food and Drug Administration’s Campaign to Reduce Tobacco Use Among Lesbian, Gay, Bisexual, and Transgender Young Adults”. Also include the FDA docket number found in brackets in the heading of this document.

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: In compliance with 44 U.S.C. 3507, FDA has submitted the following proposed collection of information to OMB for review and clearance.

Evaluation of FDA’s Campaign To Reduce Tobacco Use Among Lesbian, Gay, Bisexual, and Transgender Young Adults—OMB Control Number 0910—New

The 2009 Family Smoking Prevention and Tobacco Control Act (Tobacco Control Act) (Pub. L. 111–31) amended the Federal Food, Drug, and Cosmetic Act (the FD&C Act) to grant FDA authority to regulate the manufacture, marketing, and distribution of tobacco products to protect public health and to reduce tobacco use by minors. Section 1003(d)(2)(D) of the FD&C Act (21 U.S.C. 393(d)(2)(D)) supports the development and implementation of FDA public education campaigns

related to tobacco use. Accordingly, FDA is currently developing and implementing public education campaigns to help prevent and reduce tobacco use among lesbian, gay, bisexual, and transgender (LGBT) young adults and thereby reduce the public health burden of tobacco. Overall the campaigns will feature events; advertisements on television and radio and in print; digital communications including social media; and other forms of media.

In support of the provisions of the Tobacco Control Act that require FDA to protect the public health and to reduce tobacco use, FDA requests OMB approval to collect information needed to evaluate FDA’s campaign to reduce tobacco use among LGBT young adults. Comprehensive evaluation of FDA’s public education campaigns is needed to ensure campaign messages are effectively received, understood, and accepted by those for whom they are intended. Evaluation is an essential organizational practice in public health and a systematic way to account for and improve public health actions.

FDA plans to conduct two studies to evaluate the effectiveness of its LGBT young adult tobacco prevention campaign: (1) An outcome evaluation study to evaluate the effectiveness of its LGBT young adult tobacco prevention campaign, and (2) a media tracking questionnaire to assess awareness of and receptivity to campaign messages. The timing of these studies will be designed to follow the multiple, discrete waves of media advertising planned for the campaigns.

I. Outcome Evaluation Study

Before the beginning of data collection for the outcome evaluation study, the 5-minute screening instrument will be tested in a small pilot study of LGBT young adults aged 18 to 24. The outcome evaluation study will then begin with a baseline survey of LGBT young adults aged 18 to 24 before the campaign launch. The baseline will be followed by three followup surveys of the target audience of young adults at approximately 6-month intervals after the campaign’s launch. Information will be collected