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**DEPARTMENT OF HEALTH AND
 HUMAN SERVICES**

Food and Drug Administration

[Docket No. FDA-2013-N-0450]

**Agency Information Collection
 Activities; Submission for Office of
 Management and Budget Review;
 Comment Request; Abbreviated New
 Animal Drug Applications**

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 September
 25, 2013.

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 *oira_*
submission@omb.eop.gov. All
 comments should be identified with the
 OMB control number 0910-0669 and
 title "Abbreviated New Animal Drug
 Applications." 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, 1350 Piccard
 Dr., PI50-400B, Rockville, MD 20850,
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.

**Abbreviated New Animal Drug
 Applications—Section 512(b)(2) and
 (n)(1) of the Federal Food, Drug, and
 Cosmetic Act (21 U.S.C. 360b(b)(2) and
 (n)(1)) (OMB Control Number 0910-
 0669)—Extension**

On November 16, 1988, the President
 signed into law the Generic Animal
 Drug and Patent Restoration Act
 (GADPTRA) (Pub. L. 100-670). Under
 section 512(b)(2) of the Federal Food,
 Drug, and Cosmetic Act (the FD&C Act),
 as amended by GADPTRA, any person
 may file an abbreviated new animal
 drug application (ANADA) seeking

approval of a generic copy of an
 approved new animal drug. The
 information required to be submitted as
 part of an abbreviated application is
 described in section 512(n)(1) of the
 FD&C Act. Among other things, an
 abbreviated application is required to
 contain information to show that the
 proposed generic drug is bioequivalent
 to, and has the same labeling as, the
 approved drug referenced in the
 abbreviated application. FDA allows
 applicants to submit a complete
 ANADA or to submit information in
 support of an ANADA for phased
 review followed by the submission of an
 Administrative ANADA when FDA
 finds that all the applicable technical
 sections for an ANADA are complete.
 FDA requests that an applicant
 accompany ANADAs and requests for
 phased review of data to support
 ANADAs with the Form FDA 356v to
 ensure efficient and accurate processing
 of information to support approval of
 the generic new animal drug.

In the **Federal Register** of April 30,
 2013 (78 FR 25279), FDA published a
 60-day notice requesting public
 comment on the proposed collection of
 information. One comment was
 received; however the comment was not
 responsive to any of the four topics
 solicited by the notice. Therefore, FDA
 does not address the comment here.

FDA estimates the burden of this
 collection of information as follows:

TABLE 1—ANADAs: ESTIMATED ANNUAL REPORTING BURDEN

FD&C act section 512 (b)(2)	FDA form	Number of respondents	Number of responses per respondent	Total annual responses	Average burden per response	Total hours
ANADA	356v	18	1	18	159	2,862
Phased Review With Administrative ANADA	356v	3	5	15	31.8	477
Total						3,339

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

ANADA paperwork burden (section 512(b)(2) of the FD&C Act). Over the past 5 fiscal years, from October 2007 through September 2012, FDA has received an average of 21 ANADAs per year. FDA estimates that preparing the paperwork required under 21 U.S.C. 360b(n)(1) to be contained in an ANADA, whether all of the information is submitted with the ANADA or the applicant submits information for phased review followed by an Administrative ANADA that references that information, will take approximately 159 hours. (FDA is estimating that each ANADA that uses

the phased review process will have approximately five phased reviews per application. Therefore, assuming that three respondents will take advantage of the phased review option per year and an average of five phased reviews are submitted per application, times 31.8 hours per phased review, equals 477 total hours per year or 159 hours per application.)

Although over the last 5 fiscal years all sponsors chose to submit traditional ANADAs, some sponsors did indicate an interest in using the phased review option in the future. FDA believes that, with time, more and more sponsors will

take advantage of the phased review option as it provides greater flexibility and estimates that there will be three respondents for the phased review option. FDA also estimates that sponsors of ANADAs take approximately 25 percent less time to put together the information to support an ANADA than a new animal drug application (NADA) because they only need to provide evidence of bioequivalence and not the data required in a NADA to support a full demonstration of safety and effectiveness.

Form FDA 356v. FDA requests that an applicant fills out and sends in a Form FDA 356v with an ANADA, and with requests for phased review of data to support ANADAs, to ensure efficient and accurate processing of information to support the approval of a generic new animal drug.

Records and reports that are required post approval are described in 21 CFR 514.80, and that paperwork is already covered by that rule in OMB control number 0910-0284.

Dated: August 20, 2013.

Leslie Kux,

Assistant Commissioner for Policy.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA-2013-N-0878]

Agency Information Collection Activities; Proposed Collection; Comment Request; Premarket Notification for a New Dietary Ingredient

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 our proposed collection of certain information. 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 in response to the notice. This notice invites comments on the procedure by which a manufacturer or distributor of dietary supplements or of a new dietary ingredient is to submit information to us upon which it has based its conclusion that a dietary supplement containing a new dietary ingredient will reasonably be expected to be safe.

DATES: Submit either electronic or written comments on the collection of information by October 25, 2013.

ADDRESSES: Submit electronic comments on the collection of information to <http://www.regulations.gov>.

Submit written comments on the collection of information to the Division of Dockets Management (HFA-305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852. All comments should be identified with the 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, 1350 Piccard Dr., PI50-400B, Rockville, MD 20850, 301-796-3793, PRAStaff@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 notice of the proposed collection of information set forth in this document.

With respect to the following collection of information, FDA 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.

Premarket Notification for a New Dietary Ingredient—21 CFR 190.6 (OMB Control Number 0910-0330)—Extension

Section 413(a) of the Federal Food, Drug, and Cosmetic Act (FD&C Act) (21

U.S.C. 350b(a)) provides that at least 75 days before the introduction or delivery for introduction into interstate commerce of a dietary supplement that contains a new dietary ingredient, a manufacturer or distributor of dietary supplements or of a new dietary ingredient is to submit to us (as delegate for the Secretary of Health and Human Services) information upon which the manufacturer or distributor has based its conclusion that a dietary supplement containing a new dietary ingredient will reasonably be expected to be safe. Part 190 (21 CFR part 190) implements these statutory provisions. Section 190.6(a) requires each manufacturer or distributor of a dietary supplement containing a new dietary ingredient, or of a new dietary ingredient, to submit to the Office of Nutrition, Labeling, and Dietary Supplements (ONLDS) notification of the basis for their conclusion that said supplement or ingredient will reasonably be expected to be safe. Section 190.6(b) requires that the notification include the following: (1) The complete name and address of the manufacturer or distributor, (2) the name of the new dietary ingredient, (3) a description of the dietary supplements that contain the new dietary ingredient, and (4) the history of use or other evidence of safety establishing that the dietary ingredient will reasonably be expected to be safe.

The notification requirements described previously are designed to enable us to monitor the introduction into the food supply of new dietary ingredients and dietary supplements that contain new dietary ingredients, in order to protect consumers from the introduction of unsafe dietary supplements into interstate commerce. We use the information collected under these regulations to help ensure that a manufacturer or distributor of a dietary supplement containing a new dietary ingredient is in full compliance with the FD&C Act. We are currently developing an electronic means for submitting this information.

Description of Respondents: The respondents to this collection of information are firms in the dietary supplement industry, including dietary supplement and dietary ingredient manufacturers, packagers and re-packagers, holders, labelers and re-labelers, distributors, warehouses, exporters, and importers.

We estimate the burden of this collection of information as follows: