

certification of those accredited Canadian laboratories will continue under DOT authority. The responsibility for conducting quarterly performance testing plus periodic on-site inspections of those LAPSA-accredited laboratories was transferred to the U.S. HHS, with the HHS' NLCP contractor continuing to have an active role in the performance testing and laboratory inspection processes. Other Canadian laboratories wishing to be considered for the NLCP may apply directly to the NLCP contractor just as U.S. laboratories do.

Upon finding a Canadian laboratory to be qualified, HHS will recommend that DOT certify the laboratory (**Federal Register**, July 16, 1996) as meeting the minimum standards of the Mandatory Guidelines published in the **Federal Register** on April 30, 2010 (75 FR 22809). After receiving DOT certification, the laboratory will be included in the monthly list of HHS-certified laboratories and participate in the NLCP certification maintenance program.

Janine Denis Cook,

*Chemist Division of Workplace Programs,
Center for Substance Abuse Prevention,
SAMHSA.*

[FR Doc. 2015-07423 Filed 3-31-15; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Agency for Toxic Substances and Disease Registry

[Docket No. ATSDR-2015-0001]

Proposed Substances To Be Evaluated for Set 29 Toxicological Profiles

AGENCY: Agency for Toxic Substances and Disease Registry (ATSDR), Department of Health and Human Services (HHS).

ACTION: Request for comments on the proposed substances to be evaluated for Set 29 toxicological profiles.

SUMMARY: ATSDR is initiating the development of its 29th set of toxicological profiles (CERCLA Set 29). This notice announces the list of proposed substances that will be evaluated for CERCLA Set 29 toxicological profile development. ATSDR's Division of Toxicology and Human Health Sciences is soliciting public nominations from the list of proposed substances to be evaluated for toxicological profile development. ATSDR also will consider the nomination of any additional, non-CERCLA substances that may have public health implications, on the basis

of ATSDR's authority to prepare toxicological profiles for substances not found at sites on the National Priorities List. The agency will do so in order to ". . . establish and maintain inventory of literature, research, and studies on the health effects of toxic substances" under CERCLA Section 104(i)(1)(B), to respond to requests for consultation under section 104(i)(4), and to support the site-specific response actions conducted by ATSDR, as otherwise necessary.

DATES: Nominations from the Substance Priority List and/or additional substances must be submitted within 30 days of the publication of this notice.

ADDRESSES: You may submit nominations, identified by Docket No. ATSDR-2015-0001, by any of the following methods:

**Internet:* Access the Federal eRulemaking portal at <http://www.regulations.gov>. Follow the instructions for submitting comments.

**Mail:* Division of Toxicology and Human Health Sciences, 1600 Clifton Rd. NE., MS F-57, Atlanta, Ga., 30333.

Instructions: All submissions must include the agency name and docket number for this notice. All relevant comments will be posted without change. This means that no confidential business information or other confidential information should be submitted in response to this notice. Refer to the section *Submission of Nominations* (below) for the specific information required.

FOR FURTHER INFORMATION CONTACT: For further information, please contact Commander Jessilyn B. Taylor, Division of Toxicology and Human Health Sciences, 1600 Clifton Rd. NE., MS F-57, Atlanta, Ga., 30333, Email: tpcandidatecomments@cdc.gov; phone: 1-800-232-4636.

SUPPLEMENTARY INFORMATION: The Superfund Amendments and Reauthorization Act of 1986 (SARA) [42 U.S.C. 9601 *et seq.*] amended the Comprehensive Environmental Response, Compensation, and Liability Act of 1980 (CERCLA or Superfund) [42 U.S.C. 9601 *et seq.*] by establishing certain requirements for ATSDR and the U.S. Environmental Protection Agency (EPA) with regard to hazardous substances most commonly found at facilities on the CERCLA National Priorities List (NPL). Among these statutory requirements is a mandate for the Administrator of ATSDR to prepare toxicological profiles for each substance included on the Priority List of Hazardous Substances. This list identifies 275 hazardous substances that ATSDR and EPA have determined pose

the most significant current potential threat to human health. The availability of the revised list of the 275 priority substances was announced in the **Federal Register** on May 28, 2014 (79 FR 30613). For prior versions of the list of substances, see **Federal Register** notices dated April 17, 1987 (52 FR 12866); October 20, 1988 (53 FR 41280); October 26, 1989 (54 FR 43619); October 17, 1990 (55 FR 42067); October 17, 1991 (56 FR 52166); October 28, 1992 (57 FR 48801); February 28, 1994 (59 FR 9486); April 29, 1996 (61 FR 18744); November 17, 1997 (62 FR 61332); October 21, 1999 (64 FR 56792); October 25, 2001 (66 FR 54014); November 7, 2003 (68 FR 63098); December 7, 2005 (70 FR 72840); and March 6, 2008 (73 FR 12178); November 3, 2011 (76 FR 68193).

Substances To Be Evaluated for Set 29 Toxicological Profiles

Each year, ATSDR develops a list of substances to be considered for toxicological profile development. The Set 29 nomination process includes consideration of all substances on ATSDR's Priority List of Hazardous Substances, also known as the Substance Priority List (SPL), as well as other substances nominated by the public. The 275 substances on the SPL will be considered for Set 29 Toxicological Profile development. This list may be found at the following Web site: www.atsdr.cdc.gov/SPL.

Submission of Nominations for the Evaluation of Set 29 Proposed Substances

Today's notice invites voluntary public nominations for substances included on the SPL and for substances not listed on the SPL. All nominations should include the full name of the nominator, affiliation, and email address. When nominating a non-SPL substance, please include the rationale for the nomination. Please note that email addresses will not be posted on www.atsdr.cdc.gov.

ATSDR will evaluate all data and information associated with nominated substances and will determine the final list of substances to be chosen for toxicological profile development. Substances will be chosen according to ATSDR's specific guidelines for selection. These guidelines can be found in the *Selection Criteria* announced in the **Federal Register** on May 7, 1993 (58 FR 27286-27287). A hard copy of the Selection Criteria is available upon request or may be accessed at: http://www.atsdr.cdc.gov/toxprofiles/guidance/criteria_for_selecting_tp_support.pdf.

Please ensure that your comments are submitted within the specified nomination period. Nominations received after the closing date will be marked as late and may be considered only if time and resources permit.

Sascha Chaney,

Acting Director, Office of Policy, Planning and Evaluation National Center for Environmental Health/Agency for Toxic Substances and Disease Registry.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Title: Generic Clearance for Grant Reviewer Recruitment Form.

OMB No.: New.

Description: This notice announces that the Administration for Children and Families intends to submit the proposed Information Collection Request (Generic ICR): Generic Clearance for Grant Reviewer Application Form under the Paperwork Reduction (PRA) (44 U.S.C. 3501 *et seq.*). Comments on specific aspects for

the proposed information collection are being solicited.

This request is for approval of a plan for conducting more than one information collection that is very similar, voluntary, low-burden and uncontroversial. Information collections under this generic clearance will be in compliance with U.S. Department of Health and Humans Services' Grants Policy Directive 2.04 "Awarding Grants", and the Awarding Agency Grants Administration Manual, Chapter 2.04C "Objective Review of Grant Applications. These forms will be used to select reviewers who will participate in the grant review process for the purpose of selecting successful applications.

Respondents: Grant Reviewer Candidates.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Recruitment Form	1,500	1	0.5	750

Estimated Total Annual Burden Hours: 750 Hours.

Additional Information: Copies of the proposed collection may be obtained by writing to the Administration for Children and Families, Office of Planning, Research and Evaluation, 370 L'Enfant Promenade SW., Washington, DC 20447, Attn: ACF Reports Clearance Officer. All requests should be identified by the title of the information collection. Email address: infocollection@acf.hhs.gov.

OMB Comment: OMB is required to make a decision concerning the collection of information between 30 and 60 days after publication of this document in the **Federal Register**. Therefore, a comment is best assured of having its full effect if OMB receives it within 30 days of publication. Written comments and recommendations for the proposed information collection should be sent directly to the following: Office of Management and Budget, Paperwork Reduction Project, Email: OIRA_SUBMISSION@OMB.EOP.GOV, Attn: Desk Officer for the Administration for Children and Families.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 2015-07352 Filed 3-31-15; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA-2009-N-0025]

Agency Information Collection Activities; Proposed Collection; Comment Request; Animal Food Labeling; Declaration of Certifiable Color Additives

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing an opportunity for public comment on the proposed collection of certain information by the 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 in response to the notice. This notice solicits comments on burden hours associated with the animal food industry declaring the presence of certified and noncertified color additives in their animal food products on the animal food label.

DATES: Submit either electronic or written comments on the collection of information by June 1, 2015.

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, 8455 Colesville Rd., COLE-14526, Silver Spring, MD 20993-0002, 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,