respond to requests for consultation, 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 received by August 30, 2021.

**ADDRESSES:** You may submit nominations, identified by Docket No. ATSDR–2021–0006 by any of the following methods:

- *Internet:* Access the Federal eRulemaking portal at *www.regulations.gov.* Follow the instructions for submitting comments.
- *Mail:* Office of Innovation and Analytics, Agency for Toxic Substances and Disease Registry, 1600 Clifton Rd. NE, Mail Stop S102–1, Atlanta, GA 30329–4027. Attn: Docket No. ATSDR–2021–0006.

*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 Kambria Haire, Agency for Toxic Substances and Disease Registry, Office of Innovation and Analytics, 1600 Clifton Rd. NE, Mail Stop S102–1, Atlanta, GA 30329–4027, Email: *ATSDRToxProfileFRNs@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) (for more information, visit *www.epa.gov/superfund/superfund-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 Substance Priority List. This list identifies 275 hazardous substances found at NPL sites that ATSDR and EPA have determined pose the most significant current potential threat to human health.

Substances to be evaluated for Toxicological Profile development: Each year, ATSDR develops a list of substances to be considered for Toxicological Profile development. The nomination process includes consideration of all substances on ATSDR's SPL, as well as other substances nominated by the public. For more information on ATSDR's SPL, visit *www.atsdr.cdc.gov/SPL/.*

Submission of nominations for Toxicological Profile development: This notice invites voluntary public nominations for substances included on the SPL and for substances not listed on the SPL. When nominating a non-SPL substance, please include the rationale for the nomination. ATSDR will evaluate 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,* which may be accessed at *www.atsdr.cdc.gov/toxprofiles/guidance/ATSDR_TP_Selection%20Criteria.pdf.*

**Donata Green,**

*Acting Director, Office of Policy, Partnerships and Planning, Agency for Toxic Substances and Disease Registry.*

[FR Doc. 2021–16195 Filed 7–28–21; 8:45 am]

**BILLING CODE 4163–70–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Agency for Toxic Substances and Disease Registry

**[Docket No. ATSDR–2021–0005]**

### Availability of Six Draft Toxicological Profiles

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

**ACTION:** Notice with comment period.

**SUMMARY:** The Agency for Toxic Substances and Disease Registry (ATSDR), within the Department of Health and Human Services (HHS), announces the opening of a docket to obtain comments on drafts of six updated Toxicological Profiles: Acetone, Aldrin/Dieldrin, Chlorophenols, 3,3-Dichlorobenzidine, Disulfoton, and Pentachlorophenol.

**DATES:** Written comments must be received on or before October 27, 2021.

**ADDRESSES:** You may submit comments, identified by docket number ATSDR–2021–0005, by any of the following methods:

- *Federal eRulemaking Portal: www.regulations.gov.* Follow the instructions for submitting comments.
- *Mail:* Office of Innovation and Analytics, Agency for Toxic Substances and Disease Registry, 1600 Clifton Rd. NE, Mail Stop S102–1, Atlanta, GA, 30329–4027. Attn: Docket No. ATSDR–2021–0005.

*Instructions:* All submissions must include the agency name and Docket Number. All relevant comments received will be posted without change to *http://regulations.gov,* including any personal information provided. For access to the docket to read background documents or comments received, go to *http://www.regulations.gov.*

**FOR FURTHER INFORMATION CONTACT:** Kambria Haire, Agency for Toxic Substances and Disease Registry, Office of Innovation and Analytics, 1600 Clifton Rd. NE, Mail Stop S102–1, Atlanta, GA, 30329–4027, Email: *ATSDRToxProfileFRNs@cdc.gov*; Phone: 1–800–232–4636.

**SUPPLEMENTARY INFORMATION:** ATSDR has prepared drafts of six updated toxicological profiles based on availability of new health effects and other information since their initial release. All toxicological profiles issued as ''Drafts for Public Comment'' represent the result of ATSDR's evidence-based evaluations to provide important toxicological information on priority hazardous substances. ATSDR is seeking public comments and additional information or reports on studies about the health effects of these six substances for review and potential inclusion in the profiles. ATSDR considers key studies for these substances during the profile development process. This notice solicits any relevant, additional studies. ATSDR will evaluate the quality and relevance of such data or studies for possible inclusion in the profile.

#### Public Participation

Interested persons or organizations are invited to participate by submitting written views, information, and data.

Please note that comments received, including attachments and other supporting materials, are part of the public record and are subject to public disclosure. Comments will be posted on *https://www.regulations.gov.* Therefore, do not include any information in your comment or supporting materials that you consider confidential or inappropriate for public disclosure. If

you include your name, contact information, or other information that identifies you in the body of your comments, that information will be on public display. ATSDR will review all submissions and may choose to redact, or withhold, submissions containing private or proprietary information such as Social Security numbers, medical information, inappropriate language, or duplicate/near duplicate examples of a mass-mail campaign. ATSDR will carefully consider all comments submitted in preparation of the final Toxicological Profiles and may revise the profiles as appropriate.

**Legislative Background**

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) regarding the 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 [also called the Substance Priority List (SPL)]. This list identifies 275 hazardous substances that ATSDR and EPA have determined pose the most significant potential threat to human health. The SPL is available online at *www.atsdr.cdc.gov/spl.* ATSDR is also mandated to revise and publish updated toxicological profiles, as necessary, to reflect updated health effects and other information.

In addition, CERCLA provides ATSDR with the authority to prepare toxicological profiles for substances not found on the SPL. CERCLA authorizes ATSDR to establish and maintain an inventory of literature, research, and studies on the health effects of toxic substances (CERCLA Section 104(i)(1)(B); 42 U.S.C. 9604(i)(1)(B)); to respond to requests for health consultations (CERCLA Section 104(i)(4); 42 U.S.C. 9604(i)(4)); and to support the site-specific response actions conducted by the agency. Public nominations for substances from the SPL (or other substances)for toxicological profile development were requested on April 18, 2018 (83FR17177–17178).

ATSDR has now prepared drafts of six updated toxicological profiles based on availability of new health effects and other information since their initial release.

**Availability**

The Draft Toxicological Profiles are available online at *http://www.atsdr.cdc.gov/ToxProfiles* and at *www.regulations.gov,* Docket No. ATSDR–2021–0005.

**Donata Green,**

*Acting Director, Office of Policy, Planning and Partnerships, Agency for Toxic Substances and Disease Registry.*

[FR Doc. 2021–16188 Filed 7–28–21; 8:45 am]

**BILLING CODE 4163–70–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Medicare & Medicaid Services

**[Document Identifier: CMS–10545 and CMS–R–185]**

### Agency Information Collection Activities: Submission for OMB Review; Comment Request

**AGENCY:** Centers for Medicare & Medicaid Services, Health and Human Services (HHS).

**ACTION:** Notice.

**SUMMARY:** The Centers for Medicare & Medicaid Services (CMS) is announcing an opportunity for the public to comment on CMS' intention to collect information from the public. Under the Paperwork Reduction Act of 1995 (PRA), federal agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension or reinstatement of an existing collection of information, and to allow a second opportunity for public comment on the notice. Interested persons are invited to send comments regarding the burden estimate or any other aspect of this collection of information, including the necessity and utility of the proposed information collection for the proper performance of the agency's functions, the accuracy of the estimated burden, ways to enhance the quality, utility, and clarity of the information to be collected, and the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

**DATES:** Comments on the collection(s) of information must be received by the OMB desk officer by August 30, 2021.

**ADDRESSES:** Written comments and recommendations for the proposed information collection should be sent within 30 days of publication of this notice to *www.reginfo.gov/public/do/PRAMain.*. Find this particular information collection by selecting ''Currently under 30-day Review—Open for Public Comments'' or by using the search function.

To obtain copies of a supporting statement and any related forms for the proposed collection(s) summarized in this notice, you may make your request using one of following:

1. Access CMS' website address at website address at: *https://www.cms.gov/Regulations-and-Guidance/Legislation/PaperworkReductionActof1995/PRA-Listing.html*

**FOR FURTHER INFORMATION CONTACT:** William Parham at (410) 786–4669.

**SUPPLEMENTARY INFORMATION:** Under the Paperwork Reduction Act of 1995 (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. The term ''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 publish a 30-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension or reinstatement of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, CMS is publishing this notice that summarizes the following proposed collection(s) of information for public comment:

1. *Type of Information Collection Request:* Revision of a currently approved collection; *Title of Information Collection:* Outcome and Assessment Information Set (OASIS) OASIS–D; *Use:* Due to the COVID–19 related Public Health Emergency, the next version of the Outcome and Assessment Information Set (OASIS), version E planned for implementation January 1, 2021, was delayed. This request is for the Office of Management and Budget (OMB) approval to extend the current OASIS–D expiration date in order for home health agencies to continue data collection required for participation in the Medicare program. The current version of the OASIS–D, data item set was approved by OMB on December 6, 2018 and implemented on January 1, 2019. This request includes updated calculations using 2020 data for