

2023. An agency may neither conduct nor sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number.

Public comments were previously requested, via the **Federal Register** (87 FR 43843), on July 22, 2022 during a 60-day comment period. This notice allows for an additional 30 days for public comments. Supporting documents which explain in detail the information that the EPA will be collecting are available in the public docket for this ICR. The docket can be viewed online at <https://www.regulations.gov>, or in person, at the EPA Docket Center, WJC West Building, Room 3334, 1301 Constitution Ave. NW, Washington, DC. The telephone number for the Docket Center is 202-566-1744. For additional information about EPA's public docket, visit: <http://www.epa.gov/dockets>.

Abstract: The Emission Guidelines (EG) (40 CFR part 60, subpart Ce) for Hospital/Medical/Infectious Waste Incinerators were proposed on February 27, 1995; promulgated on September 15, 1997; and revised on both October 6, 2009 and April 4, 2011. The Federal Plan Requirements for these regulations (40 CFR part 62, subpart HHH) were proposed on July 6, 1999; promulgated on August 15, 2000; and revised on May 13, 2013. Subpart Ce requires either states or tribes to develop plans to implement the EG. If approvable state or tribal plans were not developed, the EPA was required to develop a Federal plan (Subpart HHH) to implement the Emission Guidelines for such states and tribes. The Federal plan is an interim measure to ensure that emissions standards are implemented until states assume their role as the preferred implementers of the EG. The 2013 rule finalized amendments to the HMIWI federal plan to implement the amended EG adopted on October 6, 2009, for those states that did not have an approved revised/new state plan in place within 2 years after promulgation of the EG. The regulations in 40 CFR part 60, subpart Ce and 40 CFR part 62, subpart HHH apply to each existing individual hospital/medical/infectious waste incinerator (HMIWI) that either commenced construction prior to December 2, 2008 or commenced modification prior to April 6, 2010. This information is being collected to assure compliance with 40 CFR part 60, subpart Ce and 40 CFR part 62, subpart HHH.

Form numbers: None.

Respondents/affected entities: Hospital/Medical/Infectious Waste Incinerators.

Respondent's obligation to respond: Mandatory (40 CFR part 60, subpart Ce and 40 CFR part 62, subpart HHH).

Estimated number of respondents: 28 existing respondents, consisting of 16 privately-owned, 1 Federally-owned, 1 State/locally owned HMIWI facilities, plus 10 States requiring State Plan Inventories (total).

Frequency of response: Semiannually and annually.

Total estimated burden: 19,200 hours (per year). Burden is defined at 5 CFR 1320.3(b).

Total estimated cost: \$2,430,000 (per year), which includes \$239,000 in annualized capital/startup and/or operation & maintenance costs.

Changes in the estimates: The decrease in burden from the most-recently approved ICR is due to a decrease in the number of sources due to a decline in the industry. This decrease is not due to any program changes. The number of sources has declined as HMIWI units have closed. Any new units that either commence construction after December 2, 2008 or commence modification after April 6, 2010 are subject to the new source performance standards at 40 CFR part 60, subpart Ec. The decrease in the operation and maintenance costs from the currently-approved ICR is also due to the decline in the number of HMIWI units. There are no changes to the capital costs because there are no new entities subject to 40 CFR part 60, subpart Ce or 40 CFR part 62, subpart HHH.

Courtney Kerwin,

Director, Regulatory Support Division.

[FR Doc. 2023-01760 Filed 1-27-23; 8:45 am]

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ENVIRONMENTAL PROTECTION AGENCY

[EPA-HQ-OLEM-2018-0317; FRL-10627-01-OMS]

Agency Information Collection Activities; Submission to the Office of Management and Budget for Review and Approval; Comment Request; Criteria for Classification of Solid Waste Disposal Facilities and Practices (Renewal)

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice.

SUMMARY: The Environmental Protection Agency (EPA) has submitted an information collection request (ICR), Criteria for Classification of Solid Waste Disposal Facilities and Practices,

Recordkeeping and Reporting Requirements (EPA ICR Number 1745.10, OMB Control Number 2050-0154) to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act. This is a proposed extension of the ICR, which is currently approved through February 28, 2023. Public comments were previously requested, via the **Federal Register**, on July 5, 2022 during a 60-day comment period. This notice allows for an additional 30 days for public comments.

DATES: Comments may be submitted on or before March 1, 2023.

ADDRESSES: Submit your comments, referencing Docket ID No. EPA-HQ-OLEM-2018-0317, to EPA online using www.regulations.gov (our preferred method), or by mail to: RCRA Docket (2822T), U.S. Environmental Protection Agency, 1200 Pennsylvania Avenue NW, Washington, DC 20460. EPA's policy is that all comments received will be included in the public docket without change including any personal information provided, unless the comment includes profanity, threats, information claimed to be Confidential Business Information (CBI) or other information whose disclosure is restricted by statute.

Submit written comments and recommendations to OMB for the proposed information collection 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.

FOR FURTHER INFORMATION CONTACT: Richard Huggins, Environmental Protection Agency, 1200 Pennsylvania Ave. NW, Washington, DC 20460; telephone number: 202-566-0543; email address: huggins.richard@epa.gov.

SUPPLEMENTARY INFORMATION: This is a proposed extension of the ICR, which is currently approved through February 28, 2023. An agency may not conduct or sponsor and a person is not required to respond to a collection of information unless it displays a currently valid OMB control number.

Public comments were previously requested via the **Federal Register** (87 FR 39830) on July 5, 2022 during a 60-day comment period. This notice allows for an additional 30 days for public comments. Supporting documents, which explain in detail the information that the EPA will be collecting, are available in the public docket for this ICR. The docket can be viewed online at www.regulations.gov. Materials can

also be viewed at the Reading Room located at the EPA Docket Center, WJC West Building, Room 3334, 1301 Constitution Avenue NW, Washington, DC 20004. The Docket Center's hours of operations are 8:30 a.m.–4:30 p.m., Monday–Friday (except Federal Holidays). The telephone number for the Docket Center is 202–566–1744.

Abstract: In order to effectively implement and enforce final changes to 40 CFR part 257—subpart B on a State level, owners/operators of construction and demolition waste landfills that receive CESQG hazardous wastes will have to comply with the final reporting and recordkeeping requirements. This continuing ICR documents the recordkeeping and reporting burdens associated with the location and ground-water monitoring provisions contained in 40 CFR part 257—subpart B.

Form numbers: None.

Respondents/affected entities: Entities potentially affected by this action are the private sector, as well as State, Local, or Tribal Governments.

Respondent's obligation to respond: Mandatory under Section 4010© and 3001(d)(4) of the Resource Conservation and Recovery Act (RCRA) of 1976.

Estimated number of respondents: 152.

Frequency of response: On occasion.

Total estimated burden: 11,219 hours (per year). Burden is defined at 5 CFR 1320.03(b).

Total estimated cost: \$1,951,843 per year, which includes \$1,577,659 annualized capital or operation & maintenance costs.

Changes in the estimates: There is no change in the number of hours in the total estimated respondent burden compared with the ICR currently approved by OMB.

Courtney Kerwin,

Director, Regulatory Support Division.

[FR Doc. 2023–01784 Filed 1–27–23; 8:45 am]

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ENVIRONMENTAL PROTECTION AGENCY

[EPA–HQ–OPP–2017–0751; FRL–10525–01–OCSPP]

Pesticide Registration Review; Decisions and Case Closures for Several Pesticides; Notice of Availability

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice.

SUMMARY: This notice announces the availability of EPA's interim or final registration review decisions for the following chemicals: 2-methyl-1-butanol, calcium acetate, *Candida oleophila*, cedarwood oil, citral, heptyl butyrate, *l*-carvone. In addition, this notice announces the closure of the registration review case for cetylpyridinium chloride (CPC) because the last U.S. registrations for this pesticide have been canceled.

DATES: Comments must be received on or before March 31, 2023.

ADDRESSES: Submit your comments, identified by docket identification (ID) number EPA–HQ–OPP–2017–0751, through the *Federal eRulemaking Portal* at <https://www.regulations.gov>. Follow the online instructions for submitting comments. Do not submit electronically any information you consider to be Confidential Business Information (CBI) or other information whose disclosure is restricted by statute. Additional instructions on commenting and visiting the docket, along with more information about dockets generally, is available at <https://www.epa.gov/dockets>.

FOR FURTHER INFORMATION CONTACT:

For pesticide specific information, contact: The Chemical Review Manager for the pesticide of interest identified in Table 1 in Unit IV.

For general information on the registration review program, contact: Melanie Biscoe, Pesticide Re-evaluation Division (7508P), Office of Pesticide Programs, Environmental Protection Agency, 1200 Pennsylvania Ave. NW, Washington, DC 20460–0001; telephone number: (202) 566–0701; email address: biscoe.melanie@epa.gov.

SUPPLEMENTARY INFORMATION:

I. General Information

Does this action apply to me?

This action is directed to the public in general and may be of interest to a wide range of stakeholders including environmental, human health, farm worker, and agricultural advocates; the chemical industry; pesticide users; and members of the public interested in the sale, distribution, or use of pesticides. Since others also may be interested, the Agency has not attempted to describe all the specific entities that may be affected by this action. If you have any questions regarding the applicability of this action to a particular entity, consult the pesticide specific contact person listed

under **FOR FURTHER INFORMATION CONTACT**.

II. Background

Registration review is EPA's periodic review of pesticide registrations to ensure that each pesticide continues to satisfy the statutory standard for registration, that is, the pesticide can perform its intended function without unreasonable adverse effects on human health or the environment. As part of the registration review process, the Agency has completed interim or final decisions for all pesticides listed in Table 1 in Unit IV. Through this program, EPA is ensuring that each pesticide's registration is based on current scientific and other knowledge, including its effects on human health and the environment.

III. Authority

EPA is conducting its registration review of the chemicals listed in Table 1 in Unit IV pursuant to section 3(g) of the Federal Insecticide, Fungicide, and Rodenticide Act (FIFRA) and the Procedural Regulations for Registration Review at 40 CFR part 155, subpart C. Section 3(g) of FIFRA provides, among other things, that the registrations of pesticides are to be reviewed every 15 years. Under FIFRA, a pesticide product may be registered or remain registered only if it meets the statutory standard for registration given in FIFRA section 3(c)(5) (7 U.S.C. 136a(c)(5)). When used in accordance with widespread and commonly recognized practice, the pesticide product must perform its intended function without unreasonable adverse effects on the environment; that is, without any unreasonable risk to man or the environment, or a human dietary risk from residues that result from the use of a pesticide in or on food.

IV. What action is the Agency taking?

Pursuant to 40 CFR 155.58, this notice announces the availability of EPA's interim or final registration review decisions for the pesticides shown in Table 1. The registration review decisions are supported by rationales included in the docket established for each chemical.