

TABLE 1—RODENTICIDES WITH PROPOSED INTERIM DECISIONS—Continued

Registration review case name and number	Docket ID No.	Chemical review manager and contact information
Bromadiolone, Case Number 2760	EPA-HQ-OPP-2015-0768	Steven R. Peterson, <i>OPPRodenticidesInquiries@epa.gov</i> , (202) 566-2218.
Bromethalin, Case Number 2765	EPA-HQ-OPP-2016-0077	Kent Fothergill, <i>OPPRodenticidesInquiries@epa.gov</i> , (202) 566-1943.
Chlorophacinone, Case Number 2100	EPA-HQ-OPP-2015-0778	Steven R. Peterson, <i>OPPRodenticidesInquiries@epa.gov</i> , (202) 566-2218.
Cholecalciferol, Case Number 7600	EPA-HQ-OPP-2016-0139	Kent Fothergill, <i>OPPRodenticidesInquiries@epa.gov</i> , (202) 566-1943.
Difenacoum, Case Number 7630	EPA-HQ-OPP-2015-0769	Steven R. Peterson, <i>OPPRodenticidesInquiries@epa.gov</i> , (202) 566-2218.
Difethialone, Case Number 7603	EPA-HQ-OPP-2015-0770	Steven R. Peterson, <i>OPPRodenticidesInquiries@epa.gov</i> , (202) 566-2218.
Diphacinone (and its sodium salt), Case Number 2205 ..	EPA-HQ-OPP-2015-0777	Steven R. Peterson, <i>OPPRodenticidesInquiries@epa.gov</i> , (202) 566-2218.
Strychnine, Case Number 3133	EPA-HQ-OPP-2015-0754	Shrestha, Srijana, <i>OPPRodenticidesInquiries@epa.gov</i> , (202) 566-2329.
Warfarin (and its sodium salt), Case Number 0011	EPA-HQ-OPP-2015-0481	Steven R. Peterson, <i>OPPRodenticidesInquiries@epa.gov</i> , (202) 566-2218.
Zinc Phosphide, Case Number 0026	EPA-HQ-OPP-2016-0140	Anna Senninger, <i>OPPRodenticidesInquiries@epa.gov</i> , (202) 566-2216.

The registration review docket for a pesticide includes earlier documents related to the registration review case. For example, the review opened with a Preliminary Work Plan, for public comment. A Final Work Plan was placed in the docket following public comment on the Preliminary Work Plan.

The documents in the dockets describe EPA's rationales for conducting additional risk assessments for the registration review of the rodenticides included in Table 1 in Unit IV, as well as the Agency's subsequent risk findings and consideration of possible risk mitigation measures. These proposed interim registration review decisions are supported by the rationales included in those documents. Following public comment, the Agency will issue interim or final registration review decisions for the rodenticides listed in Table 1 in Unit IV.

The registration review final rule at 40 CFR 155.58(a) provides for a minimum 60-day public comment period on all proposed interim registration review decisions. This comment period is intended to provide an opportunity for public input and a mechanism for initiating any necessary amendments to the proposed interim decision. All comments should be submitted using the methods in **ADDRESSES** and must be received by EPA on or before the closing date. These comments will become part of the docket for the pesticides included in Table 1 in Unit IV. Comments received after the close of the comment period will be marked "late." EPA is not required to consider these late comments. The Agency will carefully consider all comments received by the closing date and may provide a

"Response to Comments Memorandum" in the docket.

Background on the registration review program is provided at: <https://www.epa.gov/pesticide-reevaluation>.

Authority: 7 U.S.C. 136 *et seq.*

Dated: November 22, 2022.

Mary Reaves,

*Director, Pesticide Re-Evaluation Division,
Office of Pesticide Programs.*

[FR Doc. 2022-25978 Filed 11-28-22; 8:45 am]

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ENVIRONMENTAL PROTECTION AGENCY

[EPA-HQ-OAR-2004-0016; FRL-10442-01-OMS]

Information Collection Request Submitted to OMB for Review and Approval; Comment Request; Part 71 Federal Operating Permit Program (Renewal)

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice.

SUMMARY: The Environmental Protection Agency (EPA) has submitted an information collection request (ICR), part 71 Federal Operating Permit Program (Renewal) (EPA ICR Number 1713.13, OMB Control Number 2060-0336) to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act (PRA). This is a proposed extension of the ICR, which is currently approved through November 30, 2022. Public comments were previously requested via the **Federal Register** on January 7, 2022, during a 60-day

comment period. This notice allows for an additional 30 days for public comments. A fuller description of the ICR is given below, including its estimated burden and cost to the public. 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.

DATES: Comments must be submitted on or before December 29, 2022.

ADDRESSES: Submit your comments, referencing Docket ID No. EPA-HQ-OAR-2004-0016, online using <https://www.regulations.gov> (our preferred method), by email to a-and-r-docket@epa.gov, or by mail to: EPA Docket Center, Environmental Protection Agency, Mail Code 28221T, 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: Mayesha Choudhury, Air Quality Policy Division, Office of Air Quality Planning

and Standards, Environmental Protection Agency, Research Triangle Park, NC; telephone number: (919) 541-5297; fax number: (919) 541-5509; email address: choudhury.mayasha@epa.gov.

SUPPLEMENTARY INFORMATION:

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, Room 3334, 1301 Constitution Avenue NW, Washington, DC. The telephone number for the Docket Center is (202) 566-1744. The 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 or other information whose disclosure is restricted by statute. For additional information about EPA's public docket, visit <http://www.epa.gov/dockets>.

Abstract: Title V of the Clean Air Act (Act) requires the EPA to operate a federal operating permits program in areas not subject to an approved state program. The EPA regulations setting forth the requirements for the federal (EPA) operating permit program are at 40 CFR part 71. The part 71 program is designed to be implemented primarily by the EPA in all areas where state and local agencies do not have jurisdiction, such as Indian country and offshore, beyond states' seaward boundaries. The EPA may also delegate authority to implement the part 71 program on its behalf to a state, local or tribal agency, if the agency requests delegation and makes certain showings regarding its authority and ability to implement the program. One such delegate agency for the part 71 program exists at present.

In order to receive an operating permit for a major or other source subject to the permitting program, the applicant must conduct the necessary research, perform the appropriate analyses, and prepare the permit application with documentation to demonstrate that its facility meets all applicable statutory and regulatory requirements. Specific activities and requirements are listed and described in the Supporting Statement for the part 71 ICR.

Under part 71, the permitting authority (the EPA or a delegate agency) reviews permit applications, provides for public review of proposed permits, issues permits based on consideration of all technical factors and public input,

and reviews information submittals required of sources during the term of the permit. Under part 71, the EPA reviews certain actions and performs oversight of any delegate agency, consistent with the terms of a delegation agreement. Consequently, information prepared and submitted by sources is essential for sources to receive permits, and for federal and tribal permitting agencies to adequately review the permit applications and issue the permits, oversee implementation of the permits, and properly administer and manage the program.

Information that is collected is handled according to EPA's policies set forth in title 40, chapter 1, part 2, subpart B—Confidentiality of Business Information (see 40 CFR part 2). See also section 114(c) of the Act.

Form Numbers: The forms are 5900-01, 5900-02, 5900-03, 5900-04, 5900-05, 5900-06, 5900-79, 5900-80, 5900-81, 5900-82, 5900-83, 5900-84, 5900-85, and 5900-86.

Respondents/affected entities: Industrial plants (sources) and tribal permitting authorities.

Respondent's obligation to respond: mandatory (see 40 CFR part 71).

Estimated number of respondents: 89 (total); 88 industry sources and one tribal delegate permitting authority (the EPA serves as a permitting authority but is not a respondent).

Frequency of response: On occasion.
Total estimated burden: 23,845 hours (per year). Burden is defined at 5 CFR 1320.03(b).

Total estimated cost: \$1,858,914 (per year). There are no annualized capital or operation & maintenance costs.

Changes in estimates: There is an increase of 138 hours per year for the estimated respondent burden compared with the ICR currently approved by OMB. This increase is due to updated estimates of the number of sources and permits subject to the part 71 program, rather than any change in federal mandates.

Courtney Kerwin,

Director, Regulatory Support Division.

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NATIONAL SPACE COUNCIL

Notice of In-Space Authorization and Supervision Policy, Additional Listening Session

AGENCY: Executive Office of the President (EOP), National Space Council.

SUMMARY: On 9 September 2022, Vice President Kamala Harris, Chair of the

National Space Council, requested Council Members to provide “a proposal for the authorization and supervision of commercial novel space activities within 180 days[.]”

The White House National Space Council in the Executive Office of the President has held two virtual 2 hour listening sessions to engage with members of the public and learn about novel space capabilities and innovative missions, experiences with United States regulatory bodies, and approaches to mission authorization and supervision that can evolve over time. See **Federal Register** Notice ID NSPC-2022-0001-0001.

The National Space Council is now scheduling a third virtual listening session to accommodate additional speaker requests.

Perspectives gathered during the virtual listening sessions will inform the National Space Council as it develops a whole-of-government framework that provides a clear, predictable, and flexible process in furtherance of the *United States Space Priorities Framework (December 2021)* which states that “U.S regulations must provide clarity and certainty for the authorization and supervision of non-governmental space activities, including for novel activities such as on-orbit services, orbital debris removal, space-based manufacturing, commercial human spaceflight, and recovery and use of space resources.”

Dates

1. *Approaches for Authorization & Supervision continued:*
Thursday, 15 December 2022 1 p.m.–2 p.m. ET

Registration deadlines:
1. *Approaches for Authorization & Supervision continued:*
Thursday, 15 December 2022 1 p.m.

ADDRESSES: Register for a virtual listening session using the link below:
Approaches to Authorization & Supervision: <https://pitc.zoomgov.com/meeting/register/vJItc-6sqD8oHJZ0i2ezS2epxdLPUzub8eI>.

Please upload written comments to Regulations.gov.

FOR FURTHER INFORMATION CONTACT: Diane Howard at MBX.NSpC.IASP@ovp.eop.gov or by calling 202.456.7831.

SUPPLEMENTARY INFORMATION: Novel activities relate to those missions/activities that are not directly reviewed under existing regulatory regimes, including assembly and manufacturing, mining, and fueling stations. Participants are invited to share information about their missions—the different phases from cradle to grave as