## ENVIRONMENTAL PROTECTION AGENCY

**[EPA–HQ–OLEM–2018–0102, FRL–8935–01–OLEM]**

### Agency Information Collection Activities; Proposed Collection; Comment Request; RCRA Expanded Public Participation, EPA ICR No. 1688.09, OMB Control No. 2050–0149

**AGENCY:** Environmental Protection Agency (EPA).

**ACTION:** Notice.

**SUMMARY:** The Environmental Protection Agency (EPA) is planning to submit the information collection request (ICR), RCRA Expanded Public Participation (EPA ICR No. 1688.09, OMB Control No. 2050–0149) to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act (PRA). Before doing so, the EPA is soliciting public comments on specific aspects of the proposed information collection as described in **SUPPLEMENTARY INFORMATION**. This is a proposed extension of the ICR, which is currently approved through May 31, 2021. 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 13, 2021.

**ADDRESSES:** Submit your comments, referencing by Docket ID No. EPA–HQ–OLEM–2018–0102, to: (1) EPA online using *www.regulations.gov* (our preferred method), by email to *rcra-docket@epa.gov,* or by mail to: EPA Docket Center, Environmental Protection Agency, Mail Code 28221T, 1200 Pennsylvania Ave. NW, Washington, DC 20460, and (2) OMB via email to *oira_submission@omb.eop.gov.* Address comments to OMB Desk Officer for EPA.

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.

**FOR FURTHER INFORMATION CONTACT:** Peggy Vyas, Environmental Protection Agency, 1200 Pennsylvania Ave. NW, Washington, DC 20460; telephone number: 202–566–0453; fax number: email address: *vyas.peggy@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 *www.regulations.gov.* Out of an abundance of caution for members of the public and our staff, the EPA Docket Center and Reading Room is closed to the public, with limited exceptions, to reduce the risk of transmitting COVID–19. Our Docket Center staff will continue to provide remote customer service via email, phone and webform. For further information about the EPA's public docket, Docket Center services and the current status, please visit us online at *https://www.epa.gov/dockets.* The telephone number for the Docket Center is 202–566–1744.

Pursuant to section 3506(c)(2)(A) of the PRA, the EPA is soliciting comments and information to enable it to: (i) Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the Agency, including whether the information will have practical utility; (ii) evaluate the accuracy of the Agency's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (iii) enhance the quality, utility, and clarity of the information to be collected; and (iv) minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated electronic, mechanical, or other technological collection techniques or other forms of information technology, *e.g.,* permitting electronic submission of responses. The EPA will consider the comments received and amend the ICR as appropriate. The final ICR package will then be submitted to OMB for review and approval. At that time, the EPA will issue another **Federal Register** notice to announce the submission of the ICR to OMB and the opportunity to submit additional comments to OMB.

*Abstract:* Section 7004(b) of RCRA gives EPA broad authority to provide for, encourage, and assist public participation in the development, revision, implementation, and enforcement of any regulation, guideline, information, or program under RCRA. In addition, the statute specifies certain public notices (*i.e.,* radio, newspaper, and a letter to relevant agencies) that EPA must provide before issuing any RCRA permit. The statute also establishes a process by which the public can dispute a permit and request a public hearing to discuss it. EPA carries out much of its RCRA public involvement at 40 *CFR* Parts 124 and 270.

*Form Numbers:* None.

*Respondents/affected entities:* Entities potentially affected by this action are Businesses and other for-profit.

*Respondent's obligation to respond:* Mandatory (RCRA 7004(b)).

*Estimated number of respondents:* 46.

*Frequency of response:* On occasion.

*Total estimated burden:* 4,375 Burden is defined at 5 CFR 1320.03(b).

*Total estimated cost:* $326,263 (per year), which includes $321,833 annualized labor and $4,430 annualized capital and operation & maintenance costs.

*Changes in Estimates:* The burden hours are likely to stay substantially the same.

Dated: October 5, 2021.

**Carolyn Hoskinson,**

*Director, Office of Resource Conservation and Recovery.*

[FR Doc. 2021–22098 Filed 10–8–21; 8:45 am]

**BILLING CODE 6560–50–P**

## ENVIRONMENTAL PROTECTION AGENCY

**[EPA–HQ–RCRA–2007–0932, FRL–8936–01–OLEM]**

### Agency Information Collection Activities; Proposed Collection; Comment Request; Management Standards for Hazardous Waste Pharmaceuticals Title of ICR, EPA ICR No. 2486.03, OMB Control No. 2050–0212

**AGENCY:** Environmental Protection Agency (EPA).

**ACTION:** Notice.

**SUMMARY:** The Environmental Protection Agency (EPA) is planning to submit the information collection request (ICR), Management Standards for Hazardous Waste Pharmaceuticals (EPA ICR No. 2486.03, OMB Control No. 2050–0212) to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act (PRA). Before doing so, the EPA is soliciting public comments on specific aspects of the proposed information collection as described in **SUPPLEMENTARY INFORMATION**. This is a proposed extension of the ICR, which is currently approved through May 31, 2022. 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 13, 2021.

**ADDRESSES:** Submit your comments, referencing by Docket ID No. EPA–HQ–

RCRA–2007–0932, to: (1) EPA online using *www.regulations.gov* (our preferred method), by email to *rcra-docket@epa.gov,* or by mail to: EPA Docket Center, Environmental Protection Agency, Mail Code 28221T, 1200 Pennsylvania Ave. NW, Washington, DC 20460, and (2) OMB via email to *oira_submission@omb.eop.gov*. Address comments to OMB Desk Officer for EPA.

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.

**FOR FURTHER INFORMATION CONTACT:** Kristin Fitzgerald, Environmental Protection Agency, 1200 Pennsylvania Ave. NW, Washington, DC 20460; telephone number: 202–566–0512; email address: *fitzgerald.kristin@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 *www.regulations.gov*. Out of an abundance of caution for members of the public and our staff, the EPA Docket Center and Reading Room is closed to the public, with limited exceptions, to reduce the risk of transmitting COVID–19. Our Docket Center staff will continue to provide remote customer service via email, phone and webform. For further information about the EPA's public docket, Docket Center services and the current status, please visit us online at *https://www.epa.gov/dockets*. The telephone number for the Docket Center is 202–566–1744.

Pursuant to section 3506(c)(2)(A) of the PRA, the EPA is soliciting comments and information to enable it to: (i) Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the Agency, including whether the information will have practical utility; (ii) evaluate the accuracy of the Agency's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (iii) enhance the quality, utility, and clarity of the information to be collected; and (iv) minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated electronic, mechanical, or other technological collection techniques or other forms of information technology, *e.g.,* permitting electronic submission of responses. The EPA will consider the comments received and amend the ICR as appropriate. The final ICR package will then be submitted to OMB for review and approval. At that time, the EPA will issue another **Federal Register** notice to announce the submission of the ICR to OMB and the opportunity to submit additional comments to OMB.

*Abstract:* Some pharmaceuticals are regulated as hazardous waste under the Resource Conservation and Recovery Act (RCRA) when discarded. This final rule added regulations for the management of hazardous waste pharmaceuticals by healthcare facilities and reverse distributors. Healthcare facilities (for both humans and animals) and reverse distributors now manage their hazardous waste pharmaceuticals under a new set of sector-specific standards in lieu of the existing hazardous waste generator regulations. These regulations are found in 40 CFR 266, Subpart P, and are mandatory. The new requirements include labeling containers holding non-creditable hazardous waste pharmaceuticals and evaluated hazardous waste pharmaceuticals with the words ''Hazardous Waste Pharmaceuticals''. Healthcare facilities and reverse distributors must also track or manage rejected shipments by sending a copy of the manifest to the designated facility that returned or rejected the shipment. Additionally, healthcare facilities and reverse distributors must submit exception reports for a missing copy of a manifest. Reverse distributors are required to amend their contingency plan under 40 CFR 262 Subpart M. A reverse distributor must submit an unauthorized hazardous waste report if it receives waste it is not authorized to receive.

*Form Numbers:* None.

*Respondents/affected entities:* Entities potentially affected by this action are the private sector.

*Respondent's obligation to respond:* Mandatory (RCRA Section 3001).

*Estimated number of respondents:* 13,373.

*Frequency of response:* Annual.

*Total estimated burden:* 43,577 hours. Burden is defined at 5 CFR 1320.03(b).

*Total estimated cost:* $2,543,409, which includes $2,543,409 annualized labor costs and $0 annualized capital or O&M costs.

*Changes in Estimates:* The burden hours are expected to decrease as some of the burden associated with the rule have been incorporated into other existing ICRs.

Dated: October 5, 2021.

**Carolyn Hoskinson,**

*Director, Office of Resource Conservation and Recovery.*

[FR Doc. 2021–22097 Filed 10–8–21; 8:45 am]

**BILLING CODE 6560–50–P**

## FEDERAL COMMUNICATIONS COMMISSION

**[WC Docket No. 17–97; DA 21–1103; FR ID 50347]**

### Call Authentication Trust Anchor

**AGENCY:** Federal Communications Commission.

**ACTION:** Notice and request for comments.

**SUMMARY:** In this document, the Wireline Competition Bureau (Bureau) addresses a statutory obligation under the Pallone-Thune Telephone Robocall Abuse Criminal Enforcement and Deterrence (TRACED) Act relating to the Commission's caller ID authentication rules. Specifically, the Bureau seeks comment on STIR/SHAKEN implementation extensions granted by the Commission and associated burdens and barriers to the implementation of STIR/SHAKEN.

**DATES:** Comments are due on or before November 12, 2021; reply comments are due on or before November 26, 2021.

**ADDRESSES:** Pursuant to sections 1.415 and 1.419 of the Commission's rules, 47 CFR 1.415, 1.419, interested parties may file comments and reply comments on or before the dates indicated in this document. Comments and reply comments may be filed using the Commission's Electronic Comment Filing System (ECFS). See *Electronic Filing of Documents in Rulemaking Proceedings,* 63 FR 24121 (1998). Interested parties may file comments or reply comments, identified by WC Docket No. 17–97 by any of the following methods:

- *Electronic Filers:* Comments may be filed electronically using the internet by accessing ECFS: *https://www.fcc.gov/ecfs/*.
- *Paper Filers:* Parties who choose to file by paper must file an original and one copy of each filing.
- Filings can be sent by commercial overnight courier, or by first-class or overnight U.S. Postal Service mail. All filings must be addressed to the Commission's Secretary, Office of the Secretary, Federal Communications Commission.
- Commercial overnight mail (other than U.S. Postal Service Express Mail and Priority Mail) must be sent to 9050