simulates an actual product evaluation. (See the Safer Choice Standard at http://www2.epa.gov/saferchoice/saferchoice-standard.)

3. Following the Procedures in the Safer Choice TPP Handbook. Use the Third-Party Profiler Handbook/ Guidance Manual to inform its profiling activities, including, for example, searching the approved set of data sources; digesting the key, distinguishing hazard data; and preparing a dossier on each chemical ingredient in a candidate product.

Beyond regular oversight, the TPP will conduct annual partner surveillance, including triennial on-site audits, to ensure compliance with program requirements. (See the Third-Party Profiler Handbook; contact Tony Thompson at thompson.tony@epa.gov or (202) 564–2296 for details.)

- 4. Submitting to Performance Review. In addition to regular feedback and recommendations from Safer Choice staff, engage in annual performance reviews, and once every 5 years, a full performance evaluation, which will culminate in a Safer Choice decision on whether the TPP retains its status as a qualified TPP.
- 5. Adopting and Using the Safer Choice Data System. Use the Safer Choice data system (known as the Safer Choice Community) to manage all Safer Choice product review and partnership activities, including purchase and maintenance of the necessary computer hardware and system licenses. (For more information, see the TPP Handbook.)
- 6. Performing Supplemental Duties. Fulfill other requests and functions related to its TPP duties that may arise from time to time, as needed.
- 7. Adhering to Good Business
  Practices. Demonstrate and follow good
  business practices in dealing with
  companies working with Safer Choice.
  Good business practices include:
  Offering open access to your services to
  legitimate potential clients; adopting
  and implementing customer service
  standards; and avoiding potential
  conflicts of interest.
- 8. Signing a Memorandum of Understanding. Qualified applicants will be asked to sign a MOU, formalizing their commitment to serve as a TPP and support the Safer Choice Program's human and environmental health protection mission. (See the boilerplate MOU in the docket ID number EPA-HQ-OPPT-2015-0841.)

# **II. Instructions for Applicants**

A. How to Submit an Application

Applicants for Safer Choice Third Party Profiler must complete an application form, available in the docket ID number EPA-HQ-OPPT-2015-0841. To be considered in the 2016 selection cycle, completed applications must be submitted to Safer Choice via one of the means listed under ADDRESSES and by the deadline listed under DATES indicated in this notice.

B. Requirements for Submitting Business Proprietary Information

Clearly mark the part or all of the information that you claim to be business proprietary. In addition to one complete version of the application that includes information claimed as proprietary, a copy of the application that does not contain the proprietary information must be submitted. Information marked as business proprietary will not be disclosed except in accordance with the procedures set forth in 40 CFR part 2.

## III. Selection of Safer Choice Third-Party Profilers

Safer Choice will review and evaluate all applications based on the TPP qualifications enumerated in this notice. From the applications submitted for the 2016 TPP cycle, the program will select up to two best-qualified candidates to participate in trial product reviews. After successful completion of these reviews—during which the program will assist the candidate in orienting to the process and gaining competencythe candidate will be a qualified Safer Choice TPP. The new TPP will be asked to sign the Safer Choice-TPP memorandum of understanding and, following signature, may accept clients and begin its service to the program.

The quality and integrity of the Safer Choice label rests on active program engagement with and oversight of our profilers. Safer Choice plans to add up to two new TPPs this year, doubling its pool of profilers, based on the program's ability to assimilate, train, and oversee the new TPPs while maintaining its other program functions. Safer Choice will continually evaluate its ability to bring on additional TPPs.

All qualified TPPs will be listed in and accessible via the Safer Choice Community, the data system product manufacturers use to apply for the Safer Choice label.

Authority: 15 U.S.C. 2601 et seq.

Dated: March 9, 2016.

### Wendy C. Hamnett,

Director, Office of Pollution Prevention and Toxics.

# ENVIRONMENTAL PROTECTION AGENCY

[EPA-HQ-OPPT-2013-0677; FRL-9943-57]

# Receipt of Test Data Under the Toxic Substances Control Act

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

**ACTION:** Notice.

SUMMARY: EPA is announcing its receipt of test data submitted pursuant to a test rule issued by EPA under the Toxic Substances Control Act (TSCA). As required by TSCA, this document identifies each chemical substance and/or mixture for which test data have been received; the uses or intended uses of such chemical substance and/or mixture; and describes the nature of the test data received. Each chemical substance and/or mixture related to this announcement is identified in Unit I. under SUPPLEMENTARY INFORMATION.

### FOR FURTHER INFORMATION CONTACT:

For technical information contact: Kathy Calvo, Chemical Control Division (7405M), Office of Pollution Prevention and Toxics, Environmental Protection Agency, 1200 Pennsylvania Ave. NW., Washington, DC 20460–0001; telephone number: (202) 564–8089; email address: calvo.kathy@epa.gov.

For general information contact: The TSCA-Hotline, ABVI-Goodwill, 422 South Clinton Ave., Rochester, NY 14620; telephone number: (202) 554–1404; email address: TSCA-Hotline@epa.gov.

### SUPPLEMENTARY INFORMATION:

## I. Chemical Substances and/or Mixtures

Information about the following chemical substance and/or mixture is provided in Unit IV.:

D-erythro-hex-2-enonic acid, gammalactone, monosodium salt (CAS RN 6381–77–7).

# II. Federal Register Publication Requirement

Section 4(d) of TSCA (15 U.S.C. 2603(d)) requires EPA to publish a notice in the **Federal Register** reporting the receipt of test data submitted pursuant to test rules promulgated under TSCA section 4 (15 U.S.C. 2603).

## III. Docket Information

A docket, identified by the docket identification (ID) number EPA-HQ-OPPT-2013-0677, has been established for this **Federal Register** document that announces the receipt of data. Upon EPA's completion of its quality assurance review, the test data received will be added to the docket for the

TSCA section 4 test rule that required the test data. Use the docket ID number provided in Unit IV. to access the test data in the docket for the related TSCA section 4 test rule.

The docket for this **Federal Register** document and the docket for each related TSCA section 4 test rule is available electronically at http:// www.regulations.gov or in person at the Office of Pollution Prevention and Toxics Docket (OPPT Docket), Environmental Protection Agency Docket Center (EPA/DC), West William Jefferson Clinton Bldg., Rm. 3334, 1301 Constitution Ave. NW., Washington, DC. The Public Reading Room is open from 8:30 a.m. to 4:30 p.m., Monday through Friday, excluding legal holidays. The telephone number for the Public Reading Room is (202) 566-1744, and the telephone number for the OPPT Docket is (202) 566-0280. Please review the visitor instructions and additional information about the docket available at http://www.epa.gov/dockets.

### IV. Test Data Received

This unit contains the information required by TSCA section 4(d) for the test data received by EPA.

D-erythro-hex-2-enonic acid, gammalactone, monosodium salt (CAS RN 6381–77–7).

1. Chemical Uses: Antioxidant in food applications for which the vitamin activity of ascorbic acid (Vitamin C) is not required. Specifically, the compound is most frequently used to develop and retain the coloring and taste in meat products. It is also used for seafood products, fruit, and vegetable preservation, in beverages, and as a developing agent in photographic applications.

2. Applicable Test Rule: Chemical testing requirements for second group of high production volume chemicals (HPV2), 40 CFR 799–5087.

3. Test Data Received: The following listing describes the nature of the test data received. The test data will be added to the docket for the applicable TSCA section 4 test rule and can be found by referencing the docket ID number provided. EPA reviews of test data will be added to the same docket upon completion.

Aquatic Toxicity Study (Algae) (C1). The docket ID number assigned to this data is EPA-HQ-OPPT-2007-0531.

### V. Correction

In the previous **Federal Register** notice published February 8, 2016 (81 FR 6511) (FRL–9942–65) in the heading, the Docket Number was listed incorrectly. The correct Docket Number is: EPA–HQ–OPPT–2013–0677.

Authority: 15 U.S.C. 2601 et seq.

Dated: March 10, 2016.

### Maria J. Doa,

Director, Chemical Control Division, Office of Pollution Prevention and Toxics.

[FR Doc. 2016-06053 Filed 3-16-16; 8:45 am]

BILLING CODE 6560-50-P

# ENVIRONMENTAL PROTECTION AGENCY

[EPA-HQ-OECA-2012-0502; FRL-9943-84-OEI]

Information Collection Request Submitted to OMB for Review and Approval; Comment Request; NSPS for Hospital/Medical/Infectious Waste Incinerators (Renewal)

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

**ACTION:** Notice.

**SUMMARY:** The Environmental Protection Agency has submitted an information collection request (ICR), "NSPS for Hospital/Medical/Infectious Waste Incinerators (40 CFR part 60, subpart Ec) (Renewal)" (EPA ICR No. 1730.10, OMB Control No. 2060-0363) to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act (44 U.S.C. 3501 et seq.). This is a proposed extension of the ICR, which is currently approved through March 31, 2016. Public comments were previously requested via the **Federal Register** (80 FR 32116) on June 5, 2015, during a 60day 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:** Additional comments may be submitted on or before April 18, 2016.

ADDRESSES: Submit your comments, referencing Docket ID Number EPA—HQ—OECA—2012—0502, to (1) EPA online using www.regulations.gov (our preferred method), by email to docket.oeca@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:

Patrick Yellin, Monitoring, Assistance, and Media Programs Division, Office of Compliance, Mail Code 2227A, Environmental Protection Agency, 1200 Pennsylvania Ave. NW., Washington, DC 20460; telephone number: (202) 564–2970; fax number: (202) 564–0050; email address: yellin.patrick@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 or in person at the EPA Docket Center, EPA West, 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 affected entities are subject to the General Provisions of the NSPS at 40 CFR part 60, subpart A and any changes, or additions to the Provisions specified at 40 CFR part 60, subpart Ec. Owners or operators of the affected facilities must make an initial notification, performance tests, periodic reports, and maintain records of the occurrence and duration of any startup, shutdown, or malfunction in the operation of an affected facility, or any period during which the monitoring system is inoperative. Reports are also required semiannually.

Form Numbers: None.

Respondents/affected entities: Hospital/medical/infectious waste incinerators.

Respondent's obligation to respond: Mandatory (40 CFR part 60, subpart Ec). Estimated number of respondents: 8 (total).

Frequency of response: Initially, occasionally, semiannually and annually.

Total estimated burden: 5,670 hours (per year). Burden is defined at 5 CFR 1320.3(b).

Total estimated cost: \$972,000 (per year), includes \$402,000 annualized capital or operation & maintenance costs.

Changes in the estimates: There is an overall increase in burden in this ICR from the most recently approved ICR. This is due to an increase in the estimated number of sources subject to the regulation, and is not caused by