IV. Information Received

As specified by TSCA section 4(d), this unit identifies the information received by EPA: *Ethanedioic acid* (*CAS No. 144–62–7*).

1. *Chemical Use(s):* Ethanedioic acid is used as a rust remover; in antirust metal cleaners and coatings; as a flame-proofing and cross-linking agent in cellulose fabrics; as a reducing agent in mordent wool dying; as an acid dye stabilizing agent in nylon; as a scouring agent for cotton printing; and as a dye stripper for wool. Ethanedioic acid is also used for degumming silk; for the separation and recovery of rare earth elements from ore; for bleaching leather and masonry; for cleaning aluminum and wood decks; and as a synthetic intermediate for pharmaceuticals.

2. *Applicable Rule, Order, or Consent Agreement:* Chemical testing requirements for second group of high production volume chemicals (HPV2), 40 CFR 799.5087.

3. *Applicable docket ID number:* The information received will be added to docket ID number EPA–HQ–OPPT–2007–0531.

4. *Information Received:* EPA received the following information:

- Request for exemption from testing requirements.

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

Dated: January 4, 2017.

Maria J. Doa,

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

[FR Doc. 2017–02482 Filed 2–6–17; 8:45 am]

BILLING CODE 6560–50–P

ENVIRONMENTAL PROTECTION AGENCY

[EPA–HQ–OPP–2015–0022; FRL–9957–35]

Pesticide Product Registration; Receipt of Applications for New Uses

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice.

SUMMARY: EPA has received applications to register pesticide products containing active ingredients not included in any currently registered pesticide products. Pursuant to the Federal Insecticide, Fungicide, and Rodenticide Act (FIFRA), EPA is hereby providing notice of receipt and opportunity to comment on these applications.

DATES: Comments must be received on or before April 10, 2017.

ADDRESSES: Submit your comments, identified by docket identification (ID) numbers and the file symbols of interest as shown in the body of this notice, by one of the following methods:

- *Federal eRulemaking Portal: http://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.
- *Mail:* OPP Docket, Environmental Protection Agency Docket Center (EPA/DC), (28221T), 1200 Pennsylvania Ave. NW., Washington, DC 20460–0001.
- *Hand Delivery:* To make special arrangements for hand delivery or delivery of boxed information, please follow the instructions at *http://www.epa.gov/dockets/contacts.html.*

Additional instructions on commenting or visiting the docket, along with more information about dockets generally, is available at *http://www.epa.gov/dockets.*

FOR FURTHER INFORMATION CONTACT: Michael L. Goodis, Registration Division (7505P), Office of Pesticide Programs, Environmental Protection Agency, 1200 Pennsylvania Ave. NW., Washington, DC 20460–0001; main telephone number: (703) 305–7090; email address: *RDFRNotices@epa.gov.*

SUPPLEMENTARY INFORMATION:

I. General Information

A. Does this action apply to me?

You may be potentially affected by this action if you are an agricultural producer, food manufacturer, or pesticide manufacturer. The following list of North American Industrial Classification System (NAICS) codes is not intended to be exhaustive, but rather provides a guide to help readers determine whether this document applies to them. Potentially affected entities may include:

- Crop production (NAICS code 111).
- Animal production (NAICS code 112).
- Food manufacturing (NAICS code 311).
- Pesticide manufacturing (NAICS code 32532).

B. What should I consider as I prepare my comments for EPA?

1. *Submitting CBI.* Do not submit this information to EPA through *regulations.gov* or email. Clearly mark the part or all of the information that you claim to be CBI. For CBI information in a disk or CD–ROM that you mail to EPA, mark the outside of the disk or CD–ROM as CBI and then identify electronically within the disk or CD–ROM the specific information that is claimed as CBI. In addition to one complete version of the comment that includes information claimed as CBI, a copy of the comment that does not contain the information claimed as CBI must be submitted for inclusion in the public docket. Information so marked will not be disclosed except in accordance with procedures set forth in 40 CFR part 2.

2. *Tips for preparing your comments.* When preparing and submitting your comments, see the commenting tips at *http://www.epa.gov/dockets/comments.html.*

II. Registration Applications

EPA has received applications to register pesticide products containing active ingredients not included in any currently registered pesticide products. Pursuant to the provisions of FIFRA section 3(c)(4) (7 U.S.C. 136a(c)(4)), EPA is hereby providing notice of receipt and opportunity to comment on these applications. Notice of receipt of these applications does not imply a decision by the Agency on these applications.

1. *EPA Registration Numbers:* 100–542 and 100–620. *Docket ID Number:* EPA–HQ–OPP–2016–0495. *Applicant:* Syngenta Crop Protection, LLC, P.O. Box 18300, Greensboro, NC 27419. *Active ingredient:* Prometryn. *Product type:* Herbicide. *Proposed Use:* Sesame. *Contact:* RD.

2. *EPA Registration Numbers:* 8033–102 and 8033–103. *Docket ID Number:* EPA–HQ–OPP–2016–0649. *Applicant:* Nisso America Inc., on behalf of Nippon Soda Co., Ltd., 88 Pine Street, 14th Floor, New York, NY 10005. *Active Ingredient:* Cyflufenamid. *Product Type:* Fungicide. *Proposed Uses:* Cherry (Crop Sub-Group 12–12A), fruiting vegetables (Crop Group 8–10), and hops. *Contact:* RD.

3. *EPA Registration Number:* 11195–1. *Docket ID Number:* EPA–HQ–OPP–2016–0587. *Applicant:* Snowden Enterprises, Inc. c/o Pyxis Regulatory Consulting, Inc. 4110 136th St. Ct. NW., Gig Harbor, WA 98332. *Active Ingredient:* Sulfur dioxide. *Product Type:* Fungicide. *Proposed Use:* Fig. *Contact:* RD.

4. *EPA Registration Numbers:* 59639–2, 59639–3, 59639–83, 59639–132, and 59639–148. *Docket ID Number:* EPA–HQ–OPP–2016–0651. *Applicant:* Valent U.S.A. Corporation, 1600 Riviera Avenue, Suite #200, Walnut Creek, CA 94596. *Active Ingredient:* Clethodim. *Product Type:* Herbicide. *Proposed Uses:* Almond; brassica, leafy greens, subgroup 4–16B; leafy greens subgroup 4–16A; leaf petiole vegetable subgroup 22B; nut, tree, group 14–12; okra; onion, green, subgroup 3–07B; stalk and stem vegetable subgroup 22A; and vegetable,

fruiting, group 8–10, except okra. *Contact:* RD.

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

Dated: January 4, 2017.

Michael Goodis,

Acting Director, Registration Division, Office of Pesticide Programs.

[FR Doc. 2017–02483 Filed 2–6–17; 8:45 am]

BILLING CODE 6560–50–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

30-Day Submission Period for Requests for ONC-Approved Accreditor (ONC–AA) Status

AGENCY: Office of the National Coordinator for Health Information Technology, Department of Health and Human Services.

ACTION: Notice.

SUMMARY: This notice announces the 30-day period for submission of requests for ONC-Approved Accreditor (ONC–AA) status.

DATES: The 30-day submission period begins February 7, 2017 and will end on March 9, 2017.

FOR FURTHER INFORMATION CONTACT: Alicia Morton, Director, ONC Health IT Certification Program, Office of the National Coordinator for Health Information Technology (ONC), 202–690–7151.

SUPPLEMENTARY INFORMATION:

In accordance with 45 CFR 170.503(f)(2), an ONC-Approved Accreditor's (ONC–AA's) status will expire not later than 3 years from the date its status was granted by the National Coordinator for Health Information Technology (National Coordinator). The American National Standards Institute's (ANSI's) status as the ONC–AA for certifying bodies under the ONC Health IT Certification Program will expire on June 6, 2017. To ensure the continuity of the accreditation process and the ongoing responsibilities of the ONC–AA under the ONC Health IT Certification Program, we are seeking requests for ONC–AA status for the 3-year term that would follow the term of the current ONC–AA (ANSI). Accordingly, this notice is issued pursuant to § 170.503(b), which requires the National Coordinator to publish a notice in the **Federal Register** to announce the 30-day period during which requests for ONC–AA status may be submitted. In order to be considered for ONC–AA status, an accreditation organization must submit a written request to the National Coordinator that includes the information required by § 170.503(b) within the 30-day period specified in this notice. Section 170.503(b) requires an accreditation organization to submit the following information to demonstrate its ability to serve as an ONC–AA:

(1) A detailed description of the accreditation organization's conformance to ISO/IEC17011:2004 (incorporated by reference in § 170.599) and experience evaluating the conformance of certification bodies to ISO/IEC Guide 65:1996 (incorporated by reference in § 170.599);

(2) A detailed description of the accreditation organization's accreditation requirements, as well as how those requirements would complement the Principles of Proper Conduct for ONC-Authorized Certification Bodies (ONC–ACBs) and ensure the surveillance approaches used by ONC–ACBs include the use of consistent, objective, valid, and reliable methods;

(3) Detailed information on the accreditation organization's procedures that would be used to monitor ONC–ACBs;

(4) Detailed information, including education and experience, about the key personnel who review organizations for accreditation; and

(5) Procedures for responding to, and investigating, complaints against ONC–ACBs.

Requests for ONC–AA status may be submitted by email to *ONC-AA@hhs.gov* and should include ''Request for ONC–AA Status'' in the subject line. Alternatively, requests for ONC–AA status may be submitted by regular or express mail to: Office of the National Coordinator for Health Information Technology, Attention: ONC Health IT Certification Program—Request for ONC–AA Status, 330 C Street SW., Washington, DC 20201. In accordance with § 170.505, the official date of receipt of an email submission will be the date on which it was sent, and the official date of a submission by regular or express mail will be the date of the delivery confirmation. To clarify, email submissions may be sent up to and through 11:59 p.m. EST on the last day of the submission period. Additional information about requesting ONC–AA status and the ONC Health IT Certification Program can be found on the ONC Web site at: *http://healthit.gov/certification*.

Authority: 42 U.S.C. 300jj–11.

Jon White,

Acting National Coordinator for Health Information Technology.

[FR Doc. 2017–02458 Filed 2–6–17; 8:45 am]

BILLING CODE P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Office of the Secretary

[Document Identifier: 0990–New–30D]

Agency Information Collection Activities; Submission to OMB for Review and Approval; Public Comment Request

AGENCY: Office of the Secretary, HHS.

ACTION: Notice.

SUMMARY: In compliance with the Paperwork Reduction Act of 1995, the Office of the Secretary (OS), Department of Health and Human Services, has submitted an Information Collection Request (ICR), described below, to the Office of Management and Budget (OMB) for review and approval. The ICR is for a new collection. Comments submitted during the first public review of this ICR will be provided to OMB. OMB will accept further comments from the public on this ICR during the review and approval period.

DATES: Comments on the ICR must be received on or before March 9, 2017.

ADDRESSES: Submit your comments to *OIRA_submission@omb.eop.gov* or via facsimile to (202) 395–5806.

FOR FURTHER INFORMATION CONTACT: Information Collection Clearance staff, *Information.CollectionClearance@hhs.gov* or (202) 690–5683.

SUPPLEMENTARY INFORMATION: When submitting comments or requesting information, please include the Information Collection Request Title and document identifier 0990–New–30D for reference.

Information Collection Request Title: Evaluating Supporting Nursing Moms at Work.

Abstract: The HHS Office on Women's Health (OWH) is seeking approval by OMB on a new Information Collection Request. A Section of the Affordable Care Act (ACA) requires employers to provide basic breastfeeding accommodations for nursing mothers at work. These include a functional space, other than a bathroom, that is shielded from view and intrusion from coworkers and reasonable break time for women to express milk. OWH implemented outreach to businesses and industries across the nation to determine perceived barriers to compliance to this requirement, and became acutely aware of the sparse amount of information and resources that target worksite lactation needs and challenges of these employers.

Based upon these findings, in June, 2014, the HHS Office on Women's