TABLE 3—REGISTRANTS REQUESTING VOLUNTARY CANCELLATION AND/OR AMENDMENTS

EPA company No.	Company name and address
100	Syngenta Crop Protection, LLC, 410 Swing Road, P.O. Box 18300, Greensboro, NC 27419–8300.
279	FMC Corporation, 2929 Walnut Street, Philadelphia, PA 19104.
2693	International Paint, LLC, 6001 Antoine Drive, Houston, TX 77091.
9688	Chemsico, A Division of United Industries Corp., One Rider Trail Plaza Drive, Suite 300, Earth City, MO 63045–1313.
19713	Drexel Chemical Company, P.O. Box 13327, Memphis, TN 38113–0327.
33270	Winfield Solutions, LLC, P.O. Box 64589, St. Paul, MN 55164–0589.
34704	Loveland Products, Inc., P.O. Box 1286, Greeley, CO 80632–1286.
42750	Albaugh, LLC, 1525 NE 36th Street, Ankeny, IA 50021.
55467	Tenkoz, Inc., 1725 Windward Concourse, Suite 410, Alpharetta, GA 30005.
62719	Corteva Agriscience, LLC, 9330 Zionsville Road, Indianapolis, IN 46268.
71173	Multi-Chem Group, LLC—Odessa, 6155 W Murphy St., Odessa, TX 79763–7511.

III. What is the Agency's authority for taking this action?

Section 6(f)(1) of FIFRA (7 U.S.C. 136d(f)(1)) provides that a registrant of a pesticide product may at any time request that any of its pesticide registrations be canceled or amended to terminate one or more uses. FIFRA further provides that, before acting on the request, EPA must publish a notice of receipt of any such request in the **Federal Register**.

Section 6(f)(1)(B) of FIFRA (7 U.S.C. 136d(f)(1)(B)) requires that before acting on a request for voluntary cancellation, EPA must provide a 30-day public comment period on the request for voluntary cancellation or use termination. In addition, FIFRA section 6(f)(1)(C) (7 U.S.C. 136d(f)(1)(C)) requires that EPA provide a 180-day comment period on a request for voluntary cancellation or termination of any minor agricultural use before granting the request, unless:

1. The registrants request a waiver of the comment period, or

2. The EPA Administrator determines that continued use of the pesticide would pose an unreasonable adverse effect on the environment.

The registrants have requested that EPA waive the 180-day comment period. Accordingly, EPA will provide a 30-day comment period on the proposed requests.

IV. Procedures for Withdrawal of Requests

Registrants who choose to withdraw a request for product cancellation or use termination should submit the withdrawal in writing to the person listed under **FOR FURTHER INFORMATION CONTACT**. If the products have been subject to a previous cancellation action, the effective date of cancellation and all other provisions of any earlier cancellation action are controlling.

V. Provisions for Disposition of Existing Stocks

Existing stocks are those stocks of registered pesticide products that are currently in the United States and that were packaged, labeled, and released for shipment prior to the effective date of the action. If the requests for voluntary cancellation and amendments to terminate uses are granted, the Agency intends to publish the cancellation order in the **Federal Register**.

In any order issued in response to these requests for cancellation of product registrations and amendments to terminate uses, EPA proposes to include the following provisions for the treatment of any existing stocks of the products listed in Table 1, Table 1A and Table 2 of Unit II.

For Product 100–1431

For product 100–1431 listed in Table 1A of Unit II, the registrant has requested March 30, 2022, as the effective date of cancellation, registrants will be permitted to sell and distribute existing stocks of voluntarily canceled products for 1 year after the effective date of the cancellation, which will be March 30, 2023. Thereafter, registrants will be prohibited from selling or distributing the product identified in Table 1A of Unit II, except for export consistent with FIFRA section 17 (7 U.S.C. 136o) or for proper disposal.

For all other voluntary product cancellations, listed in Table 1 of Unit II, registrants will be permitted to sell and distribute existing stocks of voluntarily canceled products for 1 year after the effective date of the cancellation, which will be the date of publication of the cancellation order in the **Federal Register**. Thereafter, registrants will be prohibited from selling or distributing the products identified in Table 1 of Unit II, except for export consistent with FIFRA section 17 (7 U.S.C. 136o) or for proper disposal.

Once EPA has approved product labels reflecting the requested amendments to terminate uses, registrants will be permitted to sell or distribute products under the previously approved labeling for a period of 18 months after the date of **Federal Register** publication of the cancellation order, unless other restrictions have been imposed. Thereafter, registrants will be prohibited from selling or distributing the products whose labels include the terminated uses identified in Table 2 of Unit II, except for export consistent with FIFRA section 17 or for proper disposal.

Persons other than the registrant may sell, distribute, or use existing stocks of canceled products & products whose labels include the terminated uses until supplies are exhausted, provided that such sale, distribution, or use is consistent with the terms of the previously approved labeling on, or that accompanied, the canceled products & terminated uses.

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

Dated: October 15, 2021.

Marietta Echeverria,

Acting Director, Registration Division, Office of Pesticide Programs.

[FR Doc. 2021–23178 Filed 10–22–21; 8:45 am]

BILLING CODE 6560–50–P

ENVIRONMENTAL PROTECTION AGENCY

[EPA–HQ–OPPT–2021–0146; FRL–8682–05–OCSPP]

Certain New Chemicals or Significant New Uses; Statements of Findings for August 2021

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice.

SUMMARY: The Toxic Substances Control Act (TSCA) requires EPA to publish in the **Federal Register** a statement of its findings after its review of certain TSCA

notices when EPA makes a finding that a new chemical substance or significant new use is not likely to present an unreasonable risk of injury to health or the environment. Such statements apply to premanufacture notices (PMNs), microbial commercial activity notices (MCANs), and significant new use notices (SNUNs) submitted to EPA under TSCA. This document presents statements of findings made by EPA on such submissions during the period from August 1, 2021 to August 31, 2021.

FOR FURTHER INFORMATION CONTACT: *For technical information contact:* Rebecca Edelstein, Chemical Control Division (7405M), Office of Pollution Prevention and Toxics, Environmental Protection Agency, 1200 Pennsylvania Ave. NW, Washington, DC 20460–0001; telephone number: 202–564–1667 email address: *edelstein.rebecca@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. General Information

A. Does this action apply to me?

This action is directed to the public in general. As such, the Agency has not attempted to describe the specific entities that this action may apply to. Although others may be affected, this action applies directly to the submitters of the PMNs addressed in this action.

B. How can I get copies of this document and other related information?

The docket for this action, identified by docket identification (ID) number EPA–HQ–OPPT–2021–0146, is available at *http://www.regulations.gov* or 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.

Due to the public health concerns related to COVID–19, the EPA Docket Center (EPA/DC) and Reading Room is closed to visitors with limited exceptions. The staff continues to provide remote customer service via email, phone, and webform. For the latest status information on EPA/DC services and docket access, visit *https://www.epa.gov/dockets.*

II. What action is the Agency taking?

This document lists the statements of findings made by EPA after review of notices submitted under TSCA section 5(a) that certain new chemical substances or significant new uses are not likely to present an unreasonable risk of injury to health or the environment. This document presents statements of findings made by EPA during the period from August 1, 2021 to August 31, 2021.

III. What is the Agency's authority for taking this action?

TSCA section 5(a)(3) requires EPA to review a TSCA section 5(a) notice and make one of the following specific findings:

- The chemical substance or significant new use presents an unreasonable risk of injury to health or the environment;
- The information available to EPA is insufficient to permit a reasoned evaluation of the health and environmental effects of the chemical substance or significant new use;
- The information available to EPA is insufficient to permit a reasoned evaluation of the health and environmental effects and the chemical substance or significant new use may present an unreasonable risk of injury to health or the environment;
- The chemical substance is or will be produced in substantial quantities, and such substance either enters or may reasonably be anticipated to enter the environment in substantial quantities or there is or may be significant or substantial human exposure to the substance; or
- The chemical substance or significant new use is not likely to present an unreasonable risk of injury to health or the environment.

Unreasonable risk findings must be made without consideration of costs or other non-risk factors, including an unreasonable risk to a potentially exposed or susceptible subpopulation identified as relevant under the conditions of use. The term ''conditions of use'' is defined in TSCA section 3 to mean ''the circumstances, as determined by the Administrator, under which a chemical substance is intended, known, or reasonably foreseen to be manufactured, processed, distributed in commerce, used, or disposed of.''

EPA is required under TSCA section 5(g) to publish in the **Federal Register** a statement of its findings after its review of a TSCA section 5(a) notice when EPA makes a finding that a new chemical substance or significant new use is not likely to present an unreasonable risk of injury to health or the environment. Such statements apply to PMNs, MCANs, and SNUNs submitted to EPA under TSCA section 5.

Anyone who plans to manufacture (which includes import) a new chemical substance for a non-exempt commercial purpose and any manufacturer or processor wishing to engage in a use of a chemical substance designated by EPA as a significant new use must submit a notice to EPA at least 90 days before commencing manufacture of the new chemical substance or before engaging in the significant new use.

The submitter of a notice to EPA for which EPA has made a finding of ''not likely to present an unreasonable risk of injury to health or the environment'' may commence manufacture of the chemical substance or manufacture or processing for the significant new use notwithstanding any remaining portion of the applicable review period.

IV. Statements of Administrator Findings Under TSCA Section 5(a)(3)(C)

In this unit, EPA provides the following information (to the extent that such information is not claimed as Confidential Business Information (CBI)) on the PMNs, MCANs and SNUNs for which, during this period, EPA has made findings under TSCA section 5(a)(3)(C) that the new chemical substances or significant new uses are not likely to present an unreasonable risk of injury to health or the environment:

- EPA case number assigned to the TSCA section 5(a) notice.
- Chemical identity (generic name if the specific name is claimed as CBI).
- website link to EPA's decision document describing the basis of the ''not likely to present an unreasonable risk'' finding made by EPA under TSCA section 5(a)(3)(C).

EPA Case No.	Chemical identity	Website link
P–20–0148, ... P–20–0149, ... P–20–0150, ... P–20–0151	Hydroxyalkanoic acid, salt, oxidized (Generic Name).	*https://www.epa.gov/system/files/documents/2021-09/p-20-0148-0151_determination_non-cbi_final_0.pdf.*

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

Dated: October 19, 2021.

Madison Le,

Director, New Chemicals Division, Office of Pollution Prevention and Toxics.

[FR Doc. 2021–23190 Filed 10–22–21; 8:45 am]

BILLING CODE 6560–50–P

ENVIRONMENTAL PROTECTION AGENCY

[EPA–HQ–OPP–2020–0692; 9179–01–OMS]

Information Collection Request Submitted to OMB for Review and Approval; Comment Request; Pesticide Environmental Stewardship Program Annual Measures Reporting (Renewal)

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice.

SUMMARY: The Environmental Protection Agency (EPA) has submitted an information collection request (ICR), Pesticide Environmental Stewardship Program Annual Measures Reporting, (EPA ICR Number 2415.04, OMB Control Number 2070–0188) 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 October 31, 2021. Public comments were previously requested via the **Federal Register** on March 31, 2021 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: Additional comments may be submitted on or before November 24, 2021.

ADDRESSES: Submit your comments to EPA, referencing Docket ID No. EPA–HQ–OPP–2020–0692, online using *www.regulations.gov* (our preferred method), by email to *siu.carolyn@epa.gov,* or by mail to: EPA Docket Center, Environmental Protection Agency, Mail Code 28221T, 1200 Pennsylvania Ave. 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: Carolyn Siu, Mission Support Division (7101M), Office of Program Support, Office of Chemical Safety and Pollution Prevention, Environmental Protection Agency, 1200 Pennsylvania Ave. NW, Washington, DC 20460; telephone number: (703) 347–0159; email address: *siu.carolyn@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, WJC 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: This ICR covers the information collection activities associated with voluntary participation in EPA's Pesticide Environmental Stewardship Program (PESP). The program uses the information collected to establish partner membership, develop stewardship strategies, measure progress towards stewardship goals, and award incentives. PESP is an EPA partnership program that encourages the use of the integrated pest management (IPM) strategies to reduce pests and pesticide risks. IPM is an approach that involves making the best choices from among a series of pest management practices that are both economical and pose the least possible hazard to people, property, and the environment.

PESP members include pesticide end-user and organizations which focus on training, educating, and/or influencing pesticide users. To become a PESP member, an organization submits an application and a five-year strategy outlining how environmental and human health risk reduction goals will be achieved through IPM implementation and/or education. The program encourages PESP members to track progress towards IPM goals such as: reductions in unnecessary use of pesticides, cost reductions, and knowledge shared about IPM methodologies. Entities participating in PESP also benefit from technical assistance, and through incentives for achievements at different levels.

Form Numbers: 9600–01, 9600–02, and 9600–03.

Respondents/affected entities: Pesticide user companies and organizations, or entities that practice IPM or promote the use of IPM through education and training.

Respondent's obligation to respond: Voluntary (5 CFR 1320.5(d)(2)).

Estimated number of respondents: 461 (total).

Frequency of response: Annual and on occasion.

Total estimated burden: 51,562 hours (per year). Burden is defined at 5 CFR 1320.03(b).

Total estimated cost: $3,605,562 (per year), includes $0 annualized capital or operation & maintenance costs.

Changes in the Estimates: There is a correction of the number of potential respondents from 419 to 461. There is an increase of 3,897 hours in the total estimated respondent burden compared with the ICR currently approved by OMB. This increase reflects EPA's updating of burden estimates for this collection based on historical information on the number of PESP members. Based on revised estimates, the number of IPM Promoters has decreased, while the number of IPM users has increased, and the number of National IPM users has decreased since the last ICR renewal. Although the estimated burden per response has not changed for any category, the shift in