difficulties or needs further information on the substance of your comment. Any identifying or contact information provided in the body of a comment will be included as part of the comment that is placed in the official public docket and made available in EPA's electronic public docket. If EPA cannot read your comment due to technical difficulties and cannot contact you for clarification, EPA may not be able to consider your comment.

Use of the *https://www.regulations.gov* website to submit comments to EPA electronically is EPA's preferred method for receiving comments. The electronic public docket system is an ''anonymous access'' system, which means EPA will not know your identity, email address, or other contact information unless you provide it in the body of your comment.

Please ensure that your comments are submitted within the specified comment period. Comments received after the close of the comment period will be marked ''late.'' EPA is not required to consider these late comments.

Gautam Srinivasan,

Associate General Counsel.

[FR Doc. 2022–13295 Filed 6–21–22; 8:45 am]

BILLING CODE 6560–50–P

ENVIRONMENTAL PROTECTION AGENCY

[EPA–HQ–OPP–2017–0466; FRL–9928–01–OCSPP]

Spirodiclofen; Rescinding Cancellation Order for Certain Pesticide Registrations

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice.

SUMMARY: In the **Federal Register** of November 19, 2021, EPA amended the effective date of the cancellation for the two spirodiclofen registrations, EPA Registration Nos. 10163–382 and 10163–383, as requested by the registrant (Gowan), for the two registrations from December 31, 2021, until June 30, 2022. This notice announces that EPA is rescinding the cancellation order based on the submission of outstanding data that facilitated the risk assessments and registration review decision for spirodiclofen, as well as the registrant's commitment to implement label changes that adequately address the Agency's risk concerns. The registrant has submitted an amended label for the sole end-use product (EPA Registration No. 10163–383) that reflects the risk mitigation proposed by the Agency.

DATES: This recission of the cancellation order for the two spirodiclofen registrations, EPA Registration Nos. 10163–382 and 10163–383 is effective June 22, 2022.

FOR FURTHER INFORMATION CONTACT: Veronica Dutch, Pesticide Re-evaluation Division (7508M), Office of Pesticide Programs, Environmental Protection Agency, 1200 Pennsylvania Ave. NW, Washington, DC 20460–0001; telephone number: (202) 566–2352; email address: *dutch.veronica@epa.gov.*

SUPPLEMENTARY INFORMATION:

I. General Information

A. Does this action apply to me?

This action is directed to the public in general and may be of interest to a wide range of stakeholders including environmental, human health, and agricultural advocates; the chemical industry; pesticide users; and members of the public interested in the sale, distribution, or use of pesticides. Since others also may be interested, the Agency has not attempted to describe all the specific entities that may be affected by 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–OPP–2017–0466, is available at *https://www.regulations.gov.* Additional information about the docket and visiting EPA for docket access, visit *https://www.epa.gov/dockets.*

II. What action is the Agency taking?

This notice is being issued to rescind the cancellation order for the two spirodiclofen registrations (EPA Registration Nos. 10163–382 and 10163–383). Bayer CropScience had requested the voluntary cancellation of these registrations in August 2017, as an alternative to developing data required by the registration review data call-in for spirodiclofen, GDCI–124871–1883. The notice of receipt of the request to cancel was published in the **Federal Register** on October 3, 2017 (82 FR 46052) (FRL–9966–85). The Agency solicited public comment on the notice and received no comments.

The history of subsequent related FRN actions is summarized in the table below.

TABLE—CANCELLATION ACTIONS FOR EPA REGISTRATION NOS. 10163–382 AND 10163–383

FR citation	Title	Effective date	Notes
FRL–9971–10, 80 FR 60985, 12/26/2017.	Product Cancellation Order for Certain Pesticide Registrations and Amendments to Terminate Uses.	6/30/2019	
FRL–9975–97, 83 FR 16076, 4/13/2018.	Product Cancellation Orders: Certain Pesticide Registrations and Amendments to Terminate Uses; Correction.	12/31/2020	This notice amended the previous order to correct an error in the effective date.
FRL–10017–47, 85 FR 83078, 12/21/2020.	Product Cancellation Order for Certain Pesticide Registrations.	12/31/2021	This notice extended the cancellation effective date for the two spirodiclofen registrations to allow time for generating the outstanding data from GDCI–124871–1883.
FRL–9272–01, 86 FR 64929, 11/19/2021.	Product Cancellation Order for Certain Pesticide; Amendment.	6/30/2022	This notice extended the cancellation effective date for the two spirodiclofen product registrations to allow time for developing the spirodiclofen registration review risk assessments.
FRL–9928–01 6/22/2022 ...	Spirodiclofen; Rescinding Cancellation Order for Certain Pesticide Registrations.	6/22/2022	This notice rescinds the cancellation order for the two spirodiclofen product registrations.

A history of subsequent related actions is summarized here. The Agency issued the spirodiclofen registration review generic data call-in (GDCI–124871–1883) on May 11, 2016. Bayer CropScience requested the voluntary cancellation of spirodiclofen as an alternative to developing the required data. The two spirodiclofen registrations were transferred from Bayer CropScience to Gowan Company effective March 18, 2021, and Gowan requested that the cancellation order be amended to facilitate the submission of outstanding data identified in the GDCI before the cancellations would become effective. Based on data submitted by both Bayer and Gowan to fulfill the requirements of the DCI, EPA subsequently completed draft ecological and human health risk assessments (DRAs) for the registration review of spirodiclofen. The DRAs were published for public comment on October 29, 2021 and identified potential risks of concern associated with the use of spirodiclofen. EPA subsequently extended the effective date of cancellation to allow time for developing the proposed interim registration review decision (PID) for spirodiclofen. On April 4, 2022, EPA issued a PID addressing the human health and ecological risks of concern identified in the spirodiclofen DRAs and proposing measures to mitigate those risks. Gowan subsequently submitted an amended end-use product label that is responsive to the risk mitigation measures proposed by EPA. The comment period on the PID closed on June 6, 2022. The Agency intends to issue an Interim Decision (ID) for spirodiclofen after considering comments received on the PID. Because the cancellation order for the spirodiclofen product registrations was set to take effect on June 30, 2022, and because Gowan has acted in good faith to address the Agency's risk concerns and Gowan has submitted an amended label for the sole end-use product (EPA Registration No. 10163–383) that reflects the risk mitigation proposed by the Agency, EPA is now rescinding the cancellation order. The ID will account for any comments received on the PID as well as the amended label submitted by Gowan.

The cancellation order for EPA Registration Nos. 10163–382 and 10163–383 is hereby rescinded.

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

Dated: June 15, 2022.

Mary Elissa Reaves,

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

[FR Doc. 2022–13340 Filed 6–21–22; 8:45 am]

BILLING CODE 6560–50–P

ENVIRONMENTAL PROTECTION AGENCY

[EPA–HQ–OGC–2022–0510; FRL–9949–01–OGC]

Proposed Consent Decree, Clean Air Act Citizen Suit

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice of proposed consent decree; request for public comment.

SUMMARY: In accordance with the Clean Air Act, as amended (CAA or the Act), notice is given of a proposed consent decree in *Our Children's Earth Foundation* v. *Regan,* No. 3:22–cv–00695–WHA (N.D. CA). On February 2, 2022, Plaintiff Our Children's Earth Foundation filed a complaint in the United States District Court for the Northern District of California alleging that the Environmental Protection Agency (EPA or the Agency) failed to perform certain non-discretionary duties in accordance with the Act to take action on several Nevada SIP submittals by the required deadlines. The proposed consent decree would establish deadlines for EPA to take specified actions.

DATES: Written comments on the proposed consent decree must be received by July 22, 2022.

ADDRESSES: Submit your comments, identified by Docket ID No. EPA–HQ–OGC–2022–0510, online at *https://www.regulations.gov* (EPA's preferred method). Follow the online instructions for submitting comments.

Instructions: All submissions received must include the Docket ID number for this action. Comments received may be posted without change to *https://www.regulations.gov,* including any personal information provided. For detailed instructions on sending comments and additional information on the rulemaking process, see the ''Additional Information about Commenting on the Proposed Consent Decree'' heading under the **SUPPLEMENTARY INFORMATION** section of this document.

FOR FURTHER INFORMATION CONTACT: Emily Seidman, Air and Radiation Law Office, Office of General Counsel, U.S. Environmental Protection Agency; telephone (202) 564–0906; email address *seidman.emily@epa.gov.*

SUPPLEMENTARY INFORMATION:

I. Obtaining a Copy of the Proposed Consent Decree

The official public docket for this action (identified by Docket ID No. EPA–HQ–OGC–2022–0510) contains a copy of the proposed consent decree. The official public docket is available for public viewing at the Office of Environmental Information (OEI) Docket in the EPA Docket Center, EPA West, Room 3334, 1301 Constitution Ave. NW, Washington, DC. The EPA Docket Center 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 OEI Docket is (202) 566–1752.

The electronic version of the public docket for this action contains a copy of the proposed consent decree and is available through *https://www.regulations.gov.* You may use *https://www.regulations.gov* to submit or view public comments, access the index listing of the contents of the official public docket, and access those documents in the public docket that are available electronically. Once in the system, key in the appropriate docket identification number then select ''search.''

II. Additional Information About the Proposed Consent Decree

The proposed consent decree would resolve a lawsuit filed by Our Children's Earth Foundation seeking to compel the Agency to approve, disapprove, or conditionally approve, in whole or in part, several Nevada SIP submittals by the required deadlines. Specifically, the proposed consent decree would require that the appropriate EPA official or officials sign a notice or notices of final rule for publication in the **Federal Register** to approve, disapprove, conditionally approve, or approve in part and conditionally approve in part: by February 28, 2023, the Nevada Infrastructure SIP for 2012 p.m. 2.5 submittal and the PM Revised Air Quality Standards and Definitions submittal; and by April 1, 2023, seven submittals revising the Clark County Air Quality Regulations portion of the Nevada SIP and the non-transport provisions of the Nevada 2015 Ozone i-SIP submittal.

On April 21, 2022 and May 19, 2022, final rules were published in the **Federal Register** approving in full, eight Nevada submittals revising the Clark County Air Quality Regulations portion