public hearing is received and accepted by the Agency.

FOR FURTHER INFORMATION CONTACT:

Karen Seeh, U.S. Environmental Protection Agency, Office of Environmental Information, Mail Stop 2823T, 1200 Pennsylvania Avenue NW., Washington, DC 20460, (202) 566–1175, seeh.karen@epa.gov.

SUPPLEMENTARY INFORMATION: On October 13, 2005, the final Cross-Media Electronic Reporting Rule (CROMERR) was published in the Federal Register (70 FR 59848) and codified as part 3 of title 40 of the CFR. CROMERR establishes electronic reporting as an acceptable regulatory alternative to paper reporting and establishes requirements to assure that electronic documents are as legally dependable as their paper counterparts. Subpart D of CROMERR requires that state, tribal or local government agencies that receive, or wish to begin receiving, electronic reports under their EPA-authorized programs must apply to EPA for a revision or modification of those programs and obtain EPA approval. Subpart D provides standards for such approvals based on consideration of the electronic document receiving systems that the state, tribe, or local government will use to implement the electronic reporting. Additionally, § 3.1000(b) through (e) of 40 CFR part 3, subpart D provides special procedures for program revisions and modifications to allow electronic reporting, to be used at the option of the state, tribe or local government in place of procedures available under existing programspecific authorization regulations. An application submitted under the subpart D procedures must show that the state, tribe or local government has sufficient legal authority to implement the electronic reporting components of the programs covered by the application and will use electronic document receiving systems that meet the applicable subpart D requirements.

Òn March 6, 2017, the South Carolina Department of Health and Environmental Control (SCDHEC) submitted an application titled Compliance Monitoring Data Portal (CMDP) for revision to its EPA-approved drinking water program under title 40 CFR to allow new electronic reporting. EPA reviewed SCDHEC's request to revise its EPA-authorized program and, based on this review, EPA determined that the application met the standards for approval of authorized program revision set out in 40 CFR part 3, subpart D. In accordance with 40 CFR 3.1000(d), this notice of EPA's decision to approve South Carolina's request to

revise its Part 142—National Primary Drinking Water Regulations Implementation program to allow electronic reporting under 40 CFR part 141 is being published in the **Federal Register**.

SCDHEC was notified of EPA's determination to approve its application with respect to the authorized program listed above.

Also, in today's notice, EPA is informing interested persons that they may request a public hearing on EPA's action to approve the State of South Carolina's request to revise its authorized public water system program under 40 CFR part 142, in accordance with 40 CFR 3.1000(f). Requests for a hearing must be submitted to EPA within 30 days of publication of today's Federal Register notice. Such requests should include the following information:

- (1) The name, address and telephone number of the individual, organization or other entity requesting a hearing;
- (2) A brief statement of the requesting person's interest in EPA's determination, a brief explanation as to why EPA should hold a hearing, and any other information that the requesting person wants EPA to consider when determining whether to grant the request;
- (3) The signature of the individual making the request, or, if the request is made on behalf of an organization or other entity, the signature of a responsible official of the organization or other entity.

In the event a hearing is requested and granted, EPA will provide notice of the hearing in the Federal Register not less than 15 days prior to the scheduled hearing date. Frivolous or insubstantial requests for hearing may be denied by EPA. Following such a public hearing, EPA will review the record of the hearing and issue an order either affirming today's determination or rescinding such determination. If no timely request for a hearing is received and granted, EPA's approval of the State of South Carolina's request to revise its part 142—National Primary Drinking Water Regulations Implementation program to allow electronic reporting will become effective 30 days after today's notice is published, pursuant to CROMERR section 3.1000(f)(4).

Matthew Leopard,

Director, Office of Information Management. [FR Doc. 2017–07140 Filed 4–7–17; 8:45 am]

BILLING CODE 6560-50-P

ENVIRONMENTAL PROTECTION AGENCY

[EPA-HQ-OPP-2016-0618; FRL-9959-38]

Cancellation Order for Certain Pesticide Registrations and/or Amendments To Terminate Uses

AGENCY: Environmental Protection

Agency (EPA). **ACTION:** Notice.

SUMMARY: This notice announces EPA's order for the cancellations and/or amendments to terminate uses, voluntarily requested by the registrants and accepted by the Agency, of products listed in Table 1 and 2 of Unit II. pursuant to the Federal Insecticide, Fungicide, and Rodenticide Act (FIFRA). This cancellation order follows a November 22, 2016 Federal Register Notice of Receipt of Requests from the registrants listed in Table 3 of Unit II. to voluntarily cancel and/or amend to terminate uses of these product registrations. In the November 22, 2016 notice, EPA indicated that it would issue an order implementing the cancellations and/or amendments to terminate uses, unless the Agency received substantive comments within the 30-day comment period that would merit its further review of these requests, or unless the registrants withdrew their requests. The Agency did not receive any comments on the notice. Further, the registrants did not withdraw their requests. Accordingly, EPA hereby issues in this notice a cancellation order granting the requested cancellations and/or amendments to terminate uses. Any distribution, sale, or use of the products subject to this cancellation order is permitted only in accordance with the terms of this order, including any existing stocks provisions.

DATES: The propoxur product cancellations are effective December 31, 2017. The remaining cancellations and/or amendments are effective April 10, 2017.

FOR FURTHER INFORMATION CONTACT:

Brittany Pruitt, Pesticide Re-Evaluation Division (7508P), Office of Pesticide Programs, Environmental Protection Agency, 1200 Pennsylvania Ave. NW., Washington, DC 20460–0001; telephone number: (703) 347–0289; email address: pruitt.brittany@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-2016-0618, is available at http://www.regulations.gov or at the Office of Pesticide Programs Regulatory Public Docket (OPP Docket) in the Environmental Protection Agency Docket Center (EPA/DC), West William Jefferson Clinton Bldg., Rm. 3334, 1301 Constitution Ave. NW., Washington, DC 20460-0001. 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 OPP Docket is (703) 305–5805. Please review the visitor instructions and additional information about the docket available at http://www.epa.gov/dockets.

II. What action is the agency taking?

This notice announces the cancellations and/or amendments to terminate uses, as requested by registrants, of products registered under FIFRA section 3 (7 U.S.C. 136a). These registrations are listed in sequence by registration number in Tables 1 and 2 of this unit.

TABLE 1—DIMETHOMORPH, METIRAM, PROFENOFOS, PROPOXUR AND SO-DIUM ACIFLUORFEN PRODUCT CAN-CELLATIONS

EPA registration No.	Product name
4–433	Bonide Kleen Up Grass and Weed Killer, Ready to Use.
100-598	Profenofos Technical.
100-699	Curacron 8E.
279-3395	CB Invader with Propoxur.
241–383	Acrobat MZ Fungicide.
241–395	Acrobat MZ WDG Fun- gicide.
241-410	Acrobat 50 WP Fungicide.
241–411	Stature MZ Fungicide.
241–419	Stature DM Fungicide.
3862-135	Drop Dead.
6218–24	Permacide Plus.
7969–105	Polyram 80 DF.
7969–321	Cabrio Plus Fungicide.
11556–33	Sendran Technical.
89459–39	Prentox Prenbay 1.5 BC.
89459–28	Prentox Prenbay 1% Oil.
FL980001	Polyram 80 DF.

TABLE 2—CAPTAN AND PROPOXUR PRODUCT REGISTRATION AMENDMENTS TO TERMINATE USES

EPA registration No.	Product name	Uses terminated
42750–230 42750–231	Captan Technical Captan 80 DF	Turf (golf courses and sod farms), seed beds and greenhouse bench treatment. Dichondra, turf grasses (golf courses, ornamental in non-pastured areas only), grasses (lawn seedbeds), turf (sod farms).
42750–235	Captan 50% WP	Dichondra, turf grasses (ornamentals in non-pastured areas only), grasses (lawn seed-beds).
42750–236	Captan 39.75% FL	Dichondra, turf grasses (ornamental in non-pastured areas only), grasses (lawn seed beds).
84396–12	Sungro Residual Spray	Indoor aerosol, spray, and liquid formulations; use in food handling establishments and indoor crack and crevice use.

Table 3 of this unit includes the names and addresses of record for all registrants of the products in Tables 1 and 2 of this unit, in sequence by EPA company number. This number corresponds to the first part of the EPA registration numbers of the products listed above.

TABLE 3—REGISTRANTS OF CAN-CELLED AND/OR AMENDED PROD-UCTS

EPA company No.	Company name and address
4	Bonide Products, Inc., 6301 Sutliff Road, Oriskany, NY 13424.
100	Syngenta Crop Protection, P.O. Box 18300, Greensboro, NC 27419.
241	BASF Corporation, 29 Davis Drive, Research Triangle Park, NC 27709.

TABLE 3—REGISTRANTS OF CANCELLED AND/OR AMENDED PROD-UCTS—Continued

EPA company No. 279		
Market Street, Philadel- phia, PA 19103. ABC Compounding Com- pany, Inc., P.O. Box 16247, Atlanta, GA 30321. 6218 Summit Chemical Com- pany, 8322 Sharon Drive, Frederick, MD 21704. BASF Corporation, P.O. Box 13528, Research Triangle Park, NC 27709. Bayer Healthcare, LLC, P.O. Box 390, Shawnee	company	
ABC Compounding Company, Inc., P.O. Box 16247, Atlanta, GA 30321. 6218 Summit Chemical Company, 8322 Sharon Drive, Frederick, MD 21704. 7969 BASF Corporation, P.O. Box 13528, Research Triangle Park, NC 27709. 11556 Bayer Healthcare, LLC, P.O. Box 390, Shawnee	279	Market Street, Philadel-
pany, 8322 Sharon Drive, Frederick, MD 21704. 7969	3862	ABC Compounding Company, Inc., P.O. Box 16247, Atlanta, GA
Box 13528, Research Triangle Park, NC 27709. 11556	6218	pany, 8322 Sharon Drive, Frederick, MD
P.O. Box 390, Shawnee	7969	Box 13528, Research Triangle Park, NC
	11556	P.O. Box 390, Shawnee

TABLE 3—REGISTRANTS OF CANCELLED AND/OR AMENDED PROD-UCTS—Continued

EPA company No.	Company name and address
42750	Albaugh LLC, P.O. Box 2127, Valdosta, GA 31604–2127.
84396	Sungro Products, LLC, 810 E. 18th Street, Los Angeles, CA 90021.
89459	Central Garden & Pet Company, 1501 E. Woodfield Road, Suite 200, West Schaumburg, IL 60173.

III. Summary of Public Comments Received and Agency Response to Comments

During the public comment period provided, EPA received no comments in response to the November 22, 2016 **Federal Register** notice announcing the Agency's receipt of the requests for voluntary cancellations and/or amendments to terminate uses of products listed in Tables 1 and 2 of Unit II.

IV. Cancellation Order

Pursuant to FIFRA section 6(f) (7 U.S.C. 136d(f)(1)), EPA hereby approves the requested cancellations and/or amendments to terminate uses of the registrations identified in Tables 1 and 2 of Unit II. Accordingly, the Agency hereby orders that the product registrations identified in Tables 1 and 2 of Unit II. are canceled and/or amended to terminate the affected uses. The effective date of the propoxur product cancellations that are subject to this notice is December 31, 2017. The effective date of the remaining cancellations that are subject to this notice is April 10, 2017. Any distribution, sale, or use of existing stocks of the products identified in Tables 1 and 2 of Unit II. in a manner inconsistent with any of the provisions for disposition of existing stocks set forth in Unit VI. will be a violation of

V. 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. Thereafter, following the public comment period, the EPA Administrator may approve such a request. The notice of receipt for this action was published for comment in the **Federal Register** of November 22, 2016 (81 FR 83833) (FRL-9954-80). The comment period closed on December 22, 2016.

VI. Provisions for Disposition of Existing Stocks

Existing stocks are those stocks of registered pesticide products which are currently in the United States and which were packaged, labeled, and released for shipment prior to the effective date of the action. The existing stocks provisions for the products subject to this order are as follows:

A. For Propoxur Products 279–3395, 3862–135, 6218–24, 11556–33, 89459– 28, 89459–39 Identified in Table 1 of Unit II.

At the request of the registrant FMC Corporation, the effective product

cancelation date for the propoxur products listed in Table 1 of Unit II. is December 31, 2017. The registrants may continue to sell and distribute existing stocks of the propoxur products listed in Table 1 of Unit II. until December 31, 2017. Thereafter, registrants will be prohibited from selling or distributing the propoxur products identified in Table 1 of Unit II., except for export consistent with FIFRA section 17 (7 U.S.C. 1360) or for proper disposal.

Persons other than the registrant may sell, distribute, or use existing stocks of the affected cancelled products 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 cancelled products.

B. For All Other Products Identified in Table 1 and 2 of Unit II.

For all other voluntary product cancellations noted, the registrants may continue to sell and distribute existing stocks of products listed in Table 1 of Unit II. until April 10, 2018, which is 1 year after publication of this cancellation order in the **Federal Register**. Thereafter, registrants are 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. 1360) or for proper disposal.

In the case of products for which there are requested amendments to terminate uses, once EPA has approved product labels reflecting the requested amendments to terminate uses, the registrant 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, the registrant will be prohibited from selling or distributing the products whose labels include the deleted 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 the affected cancelled products/ products under the previously approved labeling 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 cancelled products/ products under the previously approved labeling.

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

Dated: March 9, 2017.

Yu-Ting Guilaran,

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

[FR Doc. 2017-07133 Filed 4-7-17; 8:45 am]

BILLING CODE 6560-50-P

ENVIRONMENTAL PROTECTION AGENCY

[FRL 9957-03-OEI]

Cross-Media Electronic Reporting: Authorized Program Revision Approval, Commonwealth of Virginia

AGENCY: Environmental Protection

Agency (EPA). **ACTION:** Notice.

SUMMARY: This notice announces EPA's approval of the Commonwealth of Virginia's request to revise its National Primary Drinking Water Regulations Implementation EPA-authorized program to allow electronic reporting.

DATES: EPA's approval is effective May 10, 2017 for the Commonwealth of Virginia's National Primary Drinking Water Regulations Implementation program, if no timely request for a public hearing is received and accepted by the Agency.

FOR FURTHER INFORMATION CONTACT:

Karen Seeh, U.S. Environmental Protection Agency, Office of Environmental Information, Mail Stop 2823T, 1200 Pennsylvania Avenue NW., Washington, DC 20460, (202) 566–1175, seeh.karen@epa.gov.

SUPPLEMENTARY INFORMATION: On October 13, 2005, the final Cross-Media Electronic Reporting Rule (CROMERR) was published in the Federal Register (70 FR 59848) and codified as part 3 of title 40 of the CFR. CROMERR establishes electronic reporting as an acceptable regulatory alternative to paper reporting and establishes requirements to assure that electronic documents are as legally dependable as their paper counterparts. Subpart D of CROMERR requires that state, tribal or local government agencies that receive, or wish to begin receiving, electronic reports under their EPA-authorized programs must apply to EPA for a revision or modification of those programs and obtain EPA approval. Subpart D provides standards for such approvals based on consideration of the electronic document receiving systems that the state, tribe, or local government will use to implement the electronic reporting. Additionally, § 3.1000(b) through (e) of 40 CFR part 3, subpart D provides special procedures for program revisions and modifications to allow