Estimated total number of potential respondents: 4.

Frequency of response: On occasion.

Estimated total average number of responses for each respondent: 1.

Estimated total annual burden hours: 40 hours.

Estimated total annual costs: $2,788. This includes an estimated burden cost of $2,788 and an estimated cost of $0 for capital investment or maintenance and operational costs.

III. Are there changes in the estimates from the last approval?

There is a decrease of 100 hours in the total estimated respondent burden compared with that identified in the ICR currently approved by OMB. This decrease reflects EPA's expectation, based on past experience, that significantly fewer respondents will apply for recognition as Champions or Partners in the next three years. This change is an adjustment.

IV. What is the next step in the process for this ICR?

EPA will consider the comments received and amend the ICR as appropriate. The final ICR package will then be submitted to OMB for review and approval pursuant to 5 CFR 1320.12. EPA will issue another **Federal Register** document pursuant to 5 CFR 1320.5(a)(1)(iv) to announce the submission of the ICR to OMB and the opportunity to submit additional comments to OMB. If you have any questions about this ICR or the approval process, please contact the technical person listed under **FOR FURTHER INFORMATION CONTACT**.

Authority: 44 U.S.C. 3501 *et seq.*

Dated: September 22, 2017.

Louise P. Wise,

Deputy Assistant Administrator, Office of Chemical Safety and Pollution Prevention.

[FR Doc. 2017–21781 Filed 10–6–17; 8:45 am]

BILLING CODE 6560–50–P

ENVIRONMENTAL PROTECTION AGENCY

[EPA–HQ–OPP–2008–0316; FRL–9967–71]

Tetrachlorvinphos; Notice of Receipt of Request To Voluntarily Cancel Certain Pesticide Registrations

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice.

SUMMARY: In accordance with the Federal Insecticide, Fungicide, and Rodenticide Act (FIFRA), EPA is issuing a notice of receipt of a request by the registrant to voluntarily cancel their registrations of certain products containing the pesticide tetrachlorvinphos (TCVP). The request would not terminate the last TCVP products registered for use in the United States. EPA intends to grant this request at the close of the comment period for this announcement unless the Agency receives substantive comments within the comment period that would merit its further review of the request, or unless the registrant withdraws its request. If this request is granted, any sale, distribution, or use of products listed in this notice will be permitted after the registration has been cancelled only if such sale, distribution, or use is consistent with the terms as described in the final order.

DATES: Comments must be received on or before November 9, 2017.

ADDRESSES: Submit your comments, identified by docket identification (ID) number EPA–HQ–OPP–2008–0316, 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: Khue Nguyen, Pesticide Re-Evaluation Division (7508P), Office of Pesticide Programs, Environmental Protection Agency, 1200 Pennsylvania Ave. NW., Washington, DC 20460–0001; telephone number: 703–347–0248; email address: *nguyen.khue@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. 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. Background on the Receipt of Request To Cancel

This notice announces receipt by EPA of a request from the registrant Bayer Healthcare, LLC to cancel certain TCVP product registrations. TCVP is an organophosphate insecticide registered for use on livestock and livestock premises and as pet collars and pet dust/powders in residential settings. In a letter dated July 7, 2017, Bayer Healthcare, LLC requested EPA to cancel certain pesticide product registrations identified in Table 1 of Unit III. Specifically, Bayer stated that the pesticide product registrations identified in Table 1 were TCVP pet collar products that were never commercialized. Bayer noted that since the products identified in Table 1 were not in the channels of trade, no existing stocks provision is required for these products. The registrant's request will not terminate the last TCVP products registered in the United States.

III. What action is the agency taking?

This notice announces receipt by EPA of a request from a registrant to cancel certain TCVP product registrations. The affected products and the registrant making the request are identified in Tables 1 and 2 of this unit.

Unless a request is withdrawn by the registrant or if the Agency determines

that there are substantive comments that warrant further review of this request, EPA intends to issue an order canceling the affected registrations.

TABLE 1—TCVP PRODUCT REGISTRATIONS WITH PENDING REQUESTS FOR CANCELLATION

Registration No.	Product name	Company
11556–164	Americare Rabon Flea & Tick Collar for Dogs	Bayer Healthcare, LLC.
11556–165	Americare Rabon Flea & Tick Collar for Cats	Bayer Healthcare, LLC.

Table 2 of this unit includes the name and address of record for the registrant of the products listed in Table 1 of this unit. This number corresponds to the first part of the EPA registration numbers of the products listed in Table 1 of this unit.

TABLE 2—REGISTRANT REQUESTING VOLUNTARY CANCELLATION

EPA Company No.	Company name and address
11556	Bayer Healthcare, LLC., P.O. Box 390, Shawnee Mission, KS 66201–0390.

IV. 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 TCVP registrant has requested that EPA waive the 180-day comment period. Accordingly, EPA will provide a 30-day comment period on the proposed requests.

V. Procedures for Withdrawal of Requests

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

VI. 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 request for voluntary cancellation is granted, the Agency intends to publish the cancellation order in the **Federal Register**.

In any order issued in response to this request for cancellation of product registrations, EPA proposes to include the following provisions for the treatment of any existing stocks of the products listed in Table 1 of Unit III.

For this voluntary cancellation request, the registrant indicates that the products listed in Table 1 of Unit III are not in the channels of trade because they were never commercialized. Therefore, no existing stocks provision is needed. The cancellation will be effective on the date of publication of the cancellation order in the **Federal Register**. Thereafter, the registrant will be prohibited from selling or distributing the products identified in Table 1 of Unit III., except for export consistent with FIFRA section 17 (7 U.S.C. 136o) or for proper disposal.

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

Dated: September 19, 2017.

Yu-Ting Guilaran,

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

[FR Doc. 2017–21795 Filed 10–6–17; 8:45 am]

BILLING CODE 6560–50–P

ENVIRONMENTAL PROTECTION AGENCY

[EPA–HQ–OPP–2009–0879; FRL–9966–71]

Environmental Modeling Public Meeting; Notice of Public Meeting

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice.

SUMMARY: An Environmental Modeling Public Meeting (EMPM) will be held on Wednesday, October 18, 2017. This Notice announces the location and time for the meeting and provides tentative agenda topics. The EMPM provides a public forum for EPA and its stakeholders to discuss current issues related to modeling pesticide fate, transport, and exposure for pesticide risk assessments in a regulatory context.

DATES: The meeting will be held on October 18, 2017 from 9:00 a.m. to 4:30 p.m. Requests to participate in the meeting must be received on or before October 20, 2017.

To request accommodation of a disability, please contact the person listed under **FOR FURTHER INFORMATON CONTACT**, preferably at least 10 days prior to the meeting, to give EPA as much time as possible to process your request.

ADDRESSES: The meeting will be held at the Environmental Protection Agency, Office of Pesticide Programs (OPP), One Potomac Yard (South Building), First Floor Conference Center (S–1200), 2777 S. Crystal Drive, Arlington, VA 22202.

FOR FURTHER INFORMATION CONTACT: Stephen Wente or Jessica Joyce, Environmental Fate and Effects Division, Office of Pesticide Programs, Environmental Protection Agency, 1200 Pennsylvania Ave. NW., Washington, DC 20460–0001; telephone number: (703) 305–0001 and (703) 347–8191; fax number: (703) 305–0204; email address: *wente.stephen@epa.gov* and *joyce.jessica@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 required to