Regional Administrator's determination and a brief statement of the information that the requesting person intends to submit at such hearing; and 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. If EPA Region 5 does not receive a timely and appropriate request for a hearing and the Regional Administrator does not elect to hold a hearing on his own motion, this determination shall become final and effective on September 8, 2020 and no further public notice will be issued.

ADDRESSES: All documents relating to this determination are available for inspection at the following offices between the hours of 9 a.m. and 4 p.m., Monday through Friday, except for official holidays and unless the offices are inaccessible due to COVID 19: Illinois Environmental Protection Agency, 1021 North Grand Avenue East, P.O. Box 19276, Springfield, Illinois 62794–9276; and the U.S. Environmental Protection Agency Region 5, Ground Water and Drinking Water Branch (WG–15J), 77 W Jackson Blvd., Chicago, IL 60604. Requestors can email Cynthia Meyer, *meyer.cynthia@epa.gov,* to receive documents related to this determination if offices are inaccessible.

FOR FURTHER INFORMATION CONTACT: Cynthia Meyer, EPA Region 5, Ground Water and Drinking Water Branch, at the address given above, by telephone at 312–886–5868, or at *meyer.cynthia@epa.gov.*

Authority: Section 1413 of the Safe Drinking Water Act, 42 U.S.C. 300g–2, and the federal regulations implementing Section 1413 of the Act set forth at 40 CFR part 142.

Dated: July 30, 2020.

Kurt Thiede,

Regional Administrator, Region 5.

[FR Doc. 2020–17162 Filed 8–5–20; 8:45 am]

BILLING CODE 6560–50–P

ENVIRONMENTAL PROTECTION AGENCY

[EPA–HQ–OPP–2009–0308; FRL–10012–80]

Tetrachlorvinphos; Notice of Receipt of Requests to Voluntarily Cancel Certain Pesticide Registrations and Amend Registrations To Terminate Certain Uses

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 requests by the registrants to voluntarily cancel their registrations of certain products containing the pesticide tetrachlorvinphos (TCVP), or to amend their TCVP product registrations to terminate or delete one or more uses. The requests would terminate TCVP use of Chem-Tech, Ltd. (Chem-Tech) dust formulations on cats and dogs, voluntarily cancel TCVP dust formulations produced by The Hartz Mountain Corporation (Hartz) for domestic animals (cats and dogs) and cancel one of Hartz's pet collars for cats. The requests would not terminate the last TCVP products registered for use in the United States, or the last TCVP pesticide products registered in the United States for these uses. EPA intends to grant these requests 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 requests. If these requests are granted, any sale, distribution, or use of products listed in this notice will be permitted after the registrations have been canceled or uses terminated 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 September 8, 2020.

ADDRESSES: Submit your comments, identified by docket identification (ID) number EPA–HQ–OPP–2009–0308, 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.*

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.*

FOR FURTHER INFORMATION CONTACT: Patricia Biggio, Pesticide Re-Evaluation Division (7508P), Office of Pesticide Programs, Environmental Protection Agency, 1200 Pennsylvania Ave. NW, Washington, DC 20460–0001; telephone number: (703) 347–0547; email address: *biggio.patricia@epa.gov.*

SUPPLEMENTARY INFORMATION:

I. Executive Summary

There are two registrants for TCVP products registered for use on cats and dogs, Hartz and Chem-Tech. The registrants have submitted requests for voluntary cancelations or use terminations.

II. 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 and human health 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.*

III. Background on the Receipt of Requests To Cancel and/or Amend Registrations To Delete Uses

This notice announces receipt by EPA of requests from registrants Hartz and

Chem-Tech to cancel certain products or terminate certain uses of TCVP product registrations. TCVP is an organophosphate insecticide used to control fleas, ticks, various flies, lice, and insect larvae on livestock and domestic animals and their premises. In letters dated June 19, 2020 and July 10, 2020, Chem-Tech and Hartz, respectively, requested EPA to either cancel or amend registrations for certain pet use products containing TCVP. These pet products and their impending actions are identified in Tables 1 and 2 of Unit IV. These actions on the registrants' requests will not terminate the last TCVP products registered in the United States, or the last TCVP pesticide products registered in the United States for these uses.

IV. What action is the Agency taking?

This notice announces receipt by EPA of requests from registrants to cancel certain registrations or terminate certain uses of TCVP product registrations. The affected products and the registrants making the requests are identified in Tables 1 and 2 of this unit.

Unless the Agency determines that there are substantive comments that warrant further review of this request, EPA intends to issue an order canceling or terminating uses from the affected registrations.

TABLE 1—TCVP PRODUCT REGISTRATIONS WITH PENDING REQUESTS FOR TERMINATION OF USE

Registration No.	Product name	Company	Uses to be deleted
47000–123	Clean Crop Livestock 1% Rabon Dust ..	Chem-Tech, Ltd	Dogs, Cats.

TABLE 2—TCVP PRODUCT REGISTRATIONS WITH PENDING REQUESTS FOR CANCELLATION

Registration No.	Product name	Company
2596–63	Hartz 2 in 1 Plus Long Lasting Collar for Cats	The Hartz Mountain Corporation.
2596–78	Hartz 2 in 1 Flea and Tick Powder for Cats	The Hartz Mountain Corporation.
2596–79	Hartz 2 in 1 Flea and Tick Powder for Dogs	The Hartz Mountain Corporation.

Table 3 of this unit includes the names and addresses of record for the registrants of the products listed in Table 1 and Table 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 in Table 1 and Table 2 of this unit.

TABLE 3—REGISTRANTS REQUESTING VOLUNTARY CANCELLATION AND/OR TERMINATION OF USE

EPA company No.	Company name and address
2596	The Hartz Mountain Corporation, 400 Plaza Drive, Seacaucus, NJ 07094.
47000	Chem-Tech, Ltd., 620 Lesher Place, Lansing, MI 48912.

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**.

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)) provides for the possibility of a 180-day comment period where a voluntary cancellation involves a pesticide registered for at least one minor agricultural use.

Because the TCVP pet uses here do not involve any minor agricultural uses, the 180-day comment provision does not apply, and EPA is providing a 30-day comment period on the requests.

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 requests for voluntary cancellation and termination of 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 for termination of uses, EPA anticipates it will include the following provisions for the treatment of any existing stocks of the products listed in Tables 1 and 2 of Unit IV:

Hartz may not ''release for shipment,'' as that term is defined by 40 CFR 152.3, any product under EPA Reg. Nos. 2596–78 and 2596–79 (dust products) after July 31, 2020, or as soon as EPA issues an order on the request following the public comment period announced in this Notice, and may not sell or distribute existing stocks of its dust products after March 31, 2021, except for export consistent with FIFRA section 17 (7 U.S.C. 136o) or for proper disposal. Hartz may sell or distribute existing stocks of EPA Reg. No. 2596–63 (cat collar) until exhausted.

Once EPA has approved product labels reflecting the requested termination of uses pertaining to EPA Reg. No. 47000–123, registrants will be permitted to sell or distribute products under the previously approved labeling, if appropriate, according to the terms of the label approval. Thereafter, registrants will be prohibited from selling or distributing the products whose labels include the deleted uses identified in Table 1 of Unit IV., except for export consistent with FIFRA section 17 or for proper disposal.

Persons other than the registrants may sell, distribute, or use existing stocks of canceled 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 canceled products.

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

Dated: July 28, 2020.

Mary Reaves,

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

[FR Doc. 2020–17114 Filed 8–5–20; 8:45 am]

BILLING CODE 6560–50–P

FEDERAL DEPOSIT INSURANCE CORPORATION

Notice of Termination of Receiverships

The Federal Deposit Insurance Corporation (FDIC or Receiver), as Receiver for each of the following insured depository institutions, was charged with the duty of winding up the affairs of the former institutions and liquidating all related assets. The Receiver has fulfilled its obligations and made all dividend distributions required by law.

NOTICE OF TERMINATION OF RECEIVERSHIPS

Fund	Receivership name	City	State	Termination date
10145	United Security Bank	Sparta	GA	8/1/2020
10170	Town Community Bank & Trust	Antioch	IL	8/1/2020

The Receiver has further irrevocably authorized and appointed FDIC-Corporate as its attorney-in-fact to execute and file any and all documents that may be required to be executed by the Receiver which FDIC-Corporate, in its sole discretion, deems necessary, including but not limited to releases, discharges, satisfactions, endorsements, assignments, and deeds. Effective on the termination dates listed above, the Receiverships have been terminated, the Receiver has been discharged, and the Receiverships have ceased to exist as legal entities.

(Authority: 12 U.S.C. 1819)

Federal Deposit Insurance Corporation.

Dated at Washington, DC, on August 3, 2020.

James P. Sheesley,

Acting Assistant Executive Secretary.

[FR Doc. 2020–17208 Filed 8–5–20; 8:45 am]

BILLING CODE 6714–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Agency for Healthcare Research and Quality

Notice of Meeting

AGENCY: Agency for Healthcare Research and Quality, HHS.

ACTION: Notice.

SUMMARY: The Agency for Healthcare Research and Quality (AHRQ) announces a Special Emphasis Panel (SEP) meeting on ''HEALTHCARE INFORMATION TECHNOLOGY RESEARCH (HITR) 2020/10–ZHS1 HSR–F (01).'' This SEP meeting will be closed to the public.

DATES: August 26, 2020.

ADDRESSES: Agency for Healthcare Research and Quality (Video Assisted Review), 5600 Fishers Lane, Rockville, Maryland 20850.

FOR FURTHER INFORMATION CONTACT: Jenny Griffith, Committee Management Officer, Office of Extramural Research, Education and Priority Populations, Agency for Healthcare Research and Quality, (AHRQ), 5600 Fishers Lane, Rockville, Maryland 20850, Telephone: (301)427–1557.

SUPPLEMENTARY INFORMATION: A Special Emphasis Panel is a group of experts in fields related to health care research who are invited by the AHRQ, and agree to be available, to conduct on an as needed basis, scientific reviews of applications for AHRQ support. Individual members of the Panel do not attend regularly-scheduled meetings and do not serve for fixed terms or a long period of time. Rather, they are asked to participate in particular review meetings which require their type of expertise.

The SEP meeting referenced above will be closed to the public in accordance with the provisions set forth in 5 U.S.C. App. 2, section 10(d), 5 U.S.C. 552b(c)(4), and 5 U.S.C. 552b(c)(6). Grant applications for the ''HEALTHCARE INFORMATION TECHNOLOGY RESEARCH (HITR) 2020/10–ZHS1 HSR–F (01)'' is to be reviewed and discussed at this meeting. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Agenda items for this meeting are subject to change as priorities dictate.

Dated: August 3, 2020.

Virginia L. Mackay-Smith,

Associate Director.

[FR Doc. 2020–17224 Filed 8–5–20; 8:45 am]

BILLING CODE 4160–90–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2011–N–0424]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Temporary Marketing Permit Applications

AGENCY: Food and Drug Administration, Health and Human Services (HHS).

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a proposed collection of information has been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995.

DATES: Submit written comments (including recommendations) on the collection of information by September 8, 2020.

ADDRESSES: To ensure that comments on the information collection are received, OMB recommends that written comments be submitted to *https://*