Pursuant to 40 CFR 155.53(c), EPA is providing an opportunity, through this notice of availability, for interested parties to provide comments and input concerning the Agency's draft human health and ecological risk assessments for the registration review of chlorethoxyfos and the draft human health risk assessments for registration review of diazinon and phosmet. Such comments and input could address, among other things, the Agency's risk assessment methodologies and assumptions, as applied to a draft risk assessment. The Agency will consider all comments received during the public comment period and make changes, as appropriate, to a draft human health and/or ecological risk assessment. EPA may then issue a revised risk assessment, explain any changes to the draft risk assessment, and respond to comments. In the **Federal Register** notice announcing the availability of the revised risk assessment, if the revised risk assessment indicates risks of concern, the Agency may provide a comment period for the public to submit suggestions for mitigating the risk identified in the risk assessment before developing a proposed registration review decision on chlorethoxyfos, diazinon, and phosmet. The Agency is re-issuing phosmet draft risk assessment in this **Federal Register** notice, initiating a 60-day comment period for phosmet.

1. *Other related information.* Additional information on the registration review status of the chemicals listed in Table 1 of Unit III, as well as information on the Agency's registration review program and on its implementing regulation is available at *http://www.epa.gov/pesticide-reevaluation.*

2. *Information submission requirements.* Anyone may submit data or information in response to this document. To be considered during a pesticide's registration review, the submitted data or information must meet the following requirements:

- To ensure that EPA will consider data or information submitted, interested persons must submit the data or information during the comment period. The Agency may, at its discretion, consider data or information submitted at a later date.
- The data or information submitted must be presented in a legible and useable form. For example, an English translation must accompany any material that is not in English, and a written transcript must accompany any information submitted as an audiographic or videographic record. Written material may be submitted in paper or electronic form.
- Submitters must clearly identify the source of any submitted data or information.
- Submitters may request the Agency to reconsider data or information that the Agency rejected in a previous review. However, submitters must explain why they believe the Agency should reconsider the data or information in the pesticide's registration review.

As provided in 40 CFR 155.58, the registration review docket for each pesticide case will remain publicly accessible through the duration of the registration review process; that is, until all actions required in the final decision on the registration review case have been completed.

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

Dated: December 21, 2016.

Yu-Ting Guilaran,

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

EDITORIAL NOTE: The Office of the Federal Register received this document on May 22, 2017.

[FR Doc. 2017–10753 Filed 5–24–17; 8:45 am]

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ENVIRONMENTAL PROTECTION AGENCY

[EPA–HQ–OPP–2015–0393; FRL–9952–83]

Registration Review Interim Decisions; Notice of Availability

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice.

SUMMARY: This notice announces the availability of EPA's interim registration review decisions for 2-(Decylthio)ethanamine Hydrochloride, DTEA–HCl; Aliphatic Alcohols, C1–C5; Bentazon; Chlorfenapyr; Propoxur; Propoxycarbazone-sodium; Sodium Acifluorfen; and Thidiazuron. The Agency is also amending the interim registration review decision for Maleic Hydrazide. Registration review is EPA's periodic review of pesticide registrations to ensure that each pesticide continues to satisfy the statutory standard for registration, that is, that the pesticide can perform its intended function without causing unreasonable adverse effects on human health or the environment. Through this program, EPA is ensuring that each pesticide's registration is based on current scientific and other knowledge, including its effects on human health and the environment.

FOR FURTHER INFORMATION CONTACT:

For pesticide specific information, contact: The Chemical Review Manager for the pesticide of interest identified in Table 1 of Unit II.

For general information on the registration review program, contact: Richard Dumas, Pesticide Re-Evaluation Division (7508P), Office of Pesticide Programs, Environmental Protection Agency, 1200 Pennsylvania Ave. NW., Washington, DC 20460–0001; telephone number: (703) 308–8015; email address: *dumas.richard@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, farm worker, 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. If you have any questions regarding the applicability of this action to a particular entity, consult the Chemical Review Manager identified in Table 1 of Unit II.

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–2015–0393, 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?

Pursuant to 40 CFR 155.58(c), this notice announces the availability of EPA's interim registration review decisions for chemicals listed in Table 1.

Pursuant to 40 CFR 155.57, a registration review decision is the Agency's determination whether a pesticide meets, or does not meet, the

standard for registration in FIFRA. EPA has considered the chemicals listed in Table 1 in light of the FIFRA standard for registration. For the chemicals listed in Table 1, the Interim Decision documents in the docket describes the Agency's rationale for issuing a registration review interim decision for these pesticides.

In addition to the interim registration review decision documents, the registration review docket for the chemicals listed in Table 1 also includes other relevant documents related to the registration review of these cases. The proposed interim registration review decisions were posted to the docket and the public was invited to submit any comments or new information.

TABLE 1—DECISIONS BEING ISSUED OR AMENDED

Registration review case name and number	Docket ID number	Chemical review manager and contact information
2-(Decylthio)ethanamine Hydrochloride, DTEA–HCl, 5029.	EPA–HQ–OPP–2009–0336	SanYvette Williams, *Williams.sanyvette@epa.gov,* 703–305–7702.
Aliphatic Alcohols, C1–C5, 4003	EPA–HQ–OPP–2012–0340	SanYvette Williams, *Williams.sanyvette@epa.gov,* 703–305–7702.
Bentazon, 0182	EPA–HQ–OPP–2010–0117	Moana Appleyard, *Appleyard.moana@epa.gov,* 703–308–8175.
Chlorfenapyr, 7419	EPA–HQ–OPP–2010–0467	Margaret Hathaway, *hathaway.margaret@.epa.gov,* 703–305–5076.
Maleic Hydrazide, 0381	EPA–HQ–OPP–2009–0387	Ricardo Jones, *jones.ricardo@epa.gov,* 703–347–0493.
Propoxur, 2555	EPA–HQ–OPP–2009–0806	Brittany Pruitt, *pruitt.brittany@epa.gov,* 703–347–0289.
Propoxycarbazone-sodium, 7264	EPA–HQ–OPP–2015–0095	Marianne Mannix, *Mannix.marianne@epa.gov,* 703–347–0275.
Sodium Acifluorfen, 2605	EPA–HQ–OPP–2010–0135	Nathan Sell, *sell.nathan@epa.gov,* (703) 347–8020.
Thidiazuron, 4092	EPA–HQ–OPP–2015–0381	Christina Motilall, *motilall.christina@epa.gov,* 703–603–0522.

EPA addresses the comments or information received during the 60-day comment period in the discussion for each pesticide listed in Table 1. From the 60-day comment period, public comments received may or may not affect the Agency's interim decision.

Pursuant to 40 CFR 155.58(c), the registration review case docket for the chemicals listed in Table 1 will remain open until all actions required in the interim decision have been completed.

Background on the registration review program is provided at: *http://www.epa.gov/pesticide-reevaluation.* Earlier documents related to the registration review of a pesticide are provided in the chemical specific dockets listed in Table 1.

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

Dated: January 11, 2017.

Yu-Ting Guilaran,

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

[FR Doc. 2017–10671 Filed 5–24–17; 8:45 am]

BILLING CODE 6560–50–P

ENVIRONMENTAL PROTECTION AGENCY

[EPA–HQ–OPP–2015–0794; FRL–9957–98]

Registration Review; Draft Human Health and/or Ecological Risk Assessment(s), and Final Tetrachlorvinphos Occupational and Residential Exposure Risk Assessment, and the Agency's Decision To Rely on Data From Human Health Research; Notice of Availability

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice.

SUMMARY: This notice announces the availability of EPA's draft human health and ecological risk assessments for the registration review of bromacil, cyprodinil, and propamocarb; the draft human health risk assessment for the registration review of cyphenothrin; and the draft ecological risk assessment for the registration review of 2, 4–D, and opens a public comment period on these documents. This notice also announces the availability of EPA's final occupational and residential exposure assessment for the registration review of tetrachlorvinphos (TCVP) and EPA's explanation for relying on TCVP data from human research on TCVP exposure from pet collars. The TCVP draft risk assessments were published for a 60-day public comment period in the **Federal Register** of January 20, 2016 (81 FR 3128) (FRL–9940–81). Registration review is EPA's periodic review of pesticide registrations to ensure that each pesticide continues to satisfy the statutory standard for registration; that is, the pesticide can perform its intended function without unreasonable adverse effects on human health or the environment. As part of the registration review process, the Agency has completed comprehensive draft human health and draft ecological risk assessments for the registration review of bromacil, cyprodinil, and propamocarb; the draft human health risk assessment for the registration review of cyphenothrin; the final occupational and residential exposure assessment for TCVP; and the draft ecological risk assessment for the registration review of 2, 4–D. After reviewing comments received during the public comment period for all pesticide cases named above (excluding TCVP), EPA may issue a revised risk assessment, explain any changes to the draft risk assessment, and respond to comments and may request public input on risk mitigation before completing a proposed registration review decision for the pesticides identified above. Regarding TCVP, the EPA has published a revised human health and final occupational and residential exposure assessment in addition to response to comments and other support documents, which explain changes to the preliminary risk assessments and responds to substantive comments. Through the registration review program, the EPA is ensuring that each pesticide's registrations are based on current scientific and other knowledge, including its effects on human health and the environment.

DATES: Comments must be received on or before July 24, 2017.

ADDRESSES: Submit your comments, identified by docket identification (ID) number EPA–HQ–OPP–2015–0794, 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.*