months of JCDH's approval and, thus, required a new construction permit; JCDH failed to issue an NSR permit for the EAF both initially and after the failure to timely commence construction; and the permit failed to assure compliance with a valid NSR permit that should have been issued by JCDH.

On June 16, 2022, the Administrator issued an Order denying the petition. The Order explains EPA's bases for denying the petition.

Dated: July 5, 2022.

Daniel Blackman,

Regional Administrator, Region 4.

[FR Doc. 2022–14902 Filed 7–12–22; 8:45 am]

BILLING CODE 6560–50–P

ENVIRONMENTAL PROTECTION AGENCY

[EPA–HQ–OPP–2022–0164; FRL–9998–01–OCSPP]

PB&ACSS JV, LLC.; Transfer of Data (June 2022)

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice.

SUMMARY: This notice announces that pesticide related information submitted to EPA's Office of Pesticide Programs (OPP) pursuant to the Federal Insecticide, Fungicide, and Rodenticide Act (FIFRA) and the Federal Food, Drug, and Cosmetic Act (FFDCA), including information that may have been claimed as Confidential Business Information (CBI) by the submitter, will be transferred to PB&ACSS JV, LLC. and its subcontractors and consultants, CDM Smith, Summitec LLC, ICF, Gibb Epidemiology, Jerrold Ward, DVM, and WinTech, LLC, in accordance with the CBI regulations. PB&ACSS JV, LLC. and its subcontractors and consultants have been awarded a contract to perform work for OPP, and access to this information will enable PB&ACSS JV, LLC. and its subcontractors and consultants to fulfill the obligations of the contract.

DATES: Data transfer and access to information will occur by July 18, 2022.

FOR FURTHER INFORMATION CONTACT: William Northern, Information Technology and Resources Management Division (7601T), Office of Pesticide Programs, Environmental Protection Agency, 1200 Pennsylvania Ave. NW, Washington, DC 20460–0001; telephone number: (703) 305–6478; email address: *northern.william@epa.gov.*

SUPPLEMENTARY INFORMATION:

I. General Information

A. Does this action apply to me?

This action applies to the public in general. As such, 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- EPA–HQ–OPP–2022–0164, is available at *http://www.regulations.gov.* Additional information about the docket, along with instructions for visiting the docket in-person, is available at *https://www.epa.gov/dockets.*

II. Contractor Requirements

Under Contract No. 68HERC22D0017, PB&ACSS JV, LLC. and its subcontractors and consultants, CDM Smith, Summitec, LLC, ICF, Gibb Epidemiology, Jerrold Ward, DVM, and WinTech, LLC, will perform critical reviews of EPA designated studies submitted by the registrants and/or from the open literature, and these reviews will be provided to the task order Contract Officer's Representative (COR) in DER or other similar study evaluation report form or system, as applicable. A template of the DER format provided to the contractor shall be followed in the preparation of Data Evaluation Records (DER)s. See DER Templates for Test Guidelines *https://www.epa.gov/pesticide-registration/study-profile-templates.* Specific guidance for completing each section is provided in the DER templates. Each review will encompass all items in the study that contribute to the overall knowledge of the pesticide, and will include the following:

1. An evaluation of the accuracy, credibility and scientific validity of that study;

2. Its suitability for meeting specific data requirements.

3. Any necessary graphic displays of data, and/or summary tables illustrating results of the study.

4. Sound scientific rationale for the conclusions reached on specific studies.

5. Clarity in data presentation and adherence to the template and overall guidance.

An evaluation for each study will include the following:

- Study identifying information.
- An in-depth examination of the materials and methods employed.
- An in-depth examination of the reported results.
- An in-depth discussion of the reviewer's scientific assessment of the study.
- A description of the reviewer's assessment which summarizes the overall conclusions and significance of the study.
- Establishment of a no observable adverse effects levels (NOAEL) and lowest observable adverse effects levels (LOAEL) based on significant toxicological effects, if applicable.
- Characterize Good Laboratory Practice (GLP) compliance in study protocols, reviews, and for data evaluations.
- Discussion of data's relevancy for supporting public health claims in review of efficacy data.

OPP has determined that access by PB&ACSS JV, LLC. and its subcontractors and consultants, CDM Smith, Summitec, LLC, ICF, Gibb Epidemiology, Jerrold Ward, DVM, and WinTech, LLC, to information on all pesticide chemicals is necessary for the performance of this contract.

Some of this information may be entitled to confidential treatment. The information has been submitted to EPA under FIFRA sections 3, 4, 6, and 7 and under FFDCA sections 408 and 409.

In accordance with the requirements of 40 CFR 2.307(h)(2), the contract with PB&ACSS JV, LLC. and its subcontractors and consultants, CDM Smith, Summitec, LLC, ICF, Gibb Epidemiology, Jerrold Ward, DVM, and WinTech, LLC, prohibits use of the information for any purpose not specified in the contract; prohibits disclosure of the information to a third party without prior written approval from the Agency; and requires that each official and employee of the contractor sign an agreement to protect the information from unauthorized release and to handle it in accordance with the *FIFRA Information Security Manual.* In addition, to the contractor and its subcontractors and consultants are required to submit for EPA approval a security plan under which any CBI will be secured and protected against unauthorized release or compromise. No information will be provided to the contractor and its subcontractors and consultants until the requirements in this document have been fully satisfied. Records of information provided to the contractor and its subcontractors and consultants will be maintained by EPA Project Officers for this contract. All information supplied to to the contractor and its subcontractors and consultants by EPA for use in connection with this contract will be returned to EPA when to the contractor

and its subcontractors and consultants have completed their work.

Authority: 7 U.S.C. 136 *et seq.;* 21 U.S.C. 301 *et seq.*

Dated: July 5, 2022.

Delores Barber,

Director, Information Technology and Resources Management Division, Office of Program Support.

[FR Doc. 2022–14937 Filed 7–12–22; 8:45 am]

BILLING CODE 6560–50–P

ENVIRONMENTAL PROTECTION AGENCY

[EPA–HQ–OPP–2017–0750; FRL–9968–01–OCSPP]

Pesticide Registration Review; Proposed Interim Decisions for Several Pesticides; Notice of Availability

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice.

SUMMARY: This notice announces the availability of EPA's proposed interim registration review decisions and opens a 60-day public comment period on the proposed interim decisions for the following pesticides: 2-methyl-1-butanol, Calcium Acetate, *Candida Oleophila,* Cedarwood Oil, Chlorflurenol Methyl Ester (CME), Citral, Heptyl butyrate, L-Carvone, Sedaxane, Tebuconazole, Triadimefon and Triadimenol.

DATES: Comments must be received on or before September 12, 2022.

ADDRESSES: Submit your comments, identified by the docket identification (ID) number for the specific pesticide of interest provided in Table 1 in Unit IV., through the *Federal eRulemaking Portal* at *https://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. Additional instructions on commenting and visiting the docket, along with more information about dockets generally, is available at *https://www.epa.gov/dockets.*

FOR FURTHER INFORMATION CONTACT:

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

For general information on the registration review program, contact: Melanie Biscoe, Pesticide Re-Evaluation Division (7508P), Office of Pesticide Programs, Environmental Protection Agency, 1200 Pennsylvania Ave. NW, Washington, DC 20460–0001; email address: *biscoe.melanie@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 for the pesticide of interest identified in Table 1 in Unit IV.

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 on 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: *https://www.epa.gov/dockets/commenting-epa-dockets.*

3. *Environmental justice.* EPA seeks to achieve environmental justice, the fair treatment and meaningful involvement of any group, including minority and/or low-income populations, in the development, implementation, and enforcement of environmental laws, regulations, and policies. To help address potential environmental justice issues, the Agency seeks information on any groups or segments of the population who, as a result of their location, cultural practices, or other factors, may have atypical or disproportionately high and adverse human health impacts or environmental effects from exposure to the pesticides discussed in this document, compared to the general population.

II. Background

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 proposed interim decisions for all pesticides listed in Table 1 in Unit IV. 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.

III. Authority

EPA is conducting its registration review of the chemicals listed in the Table 1 in Unit IV. pursuant to section 3(g) of the Federal Insecticide, Fungicide, and Rodenticide Act (FIFRA) and the Procedural Regulations for Registration Review at 40 CFR part 155, subpart C. FIFRA section 3(g) provides, among other things, that the registrations of pesticides are to be reviewed every 15 years. Under FIFRA, a pesticide product may be registered or remain registered only if it meets the statutory standard for registration given in FIFRA section 3(c)(5) (7 U.S.C. 136a(c)(5)). When used in accordance with widespread and commonly recognized practice, the pesticide product must perform its intended function without unreasonable adverse effects on the environment; that is, without any unreasonable risk to man or the environment, or a human dietary risk from residues that result from the use of a pesticide in or on food.

IV. What action is the Agency taking?

Pursuant to 40 CFR 155.58, this notice announces the availability of EPA's proposed interim registration review decisions for the pesticides shown in Table 1 and opens a 60-day public comment period on the proposed interim registration review decisions.