Comments Due: 5 p.m. ET 3/23/21.

Docket Numbers: ER21–1251–000.

Applicants: Bighorn Solar 1, LLC.

Description: Baseline eTariff Filing: MBR Application with Request for Waivers & Expediting to be effective 4/1/2021.

Filed Date: 3/2/21.

Accession Number: 20210302–5130.

Comments Due: 5 p.m. ET 3/23/21.

The filings are accessible in the Commission's eLibrary system (*https://elibrary.ferc.gov/idmws/search/fercgensearch.asp*) by querying the docket number.

Any person desiring to intervene or protest in any of the above proceedings must file in accordance with Rules 211 and 214 of the Commission's Regulations (18 CFR 385.211 and 385.214) on or before 5:00 p.m. Eastern time on the specified comment date. Protests may be considered, but intervention is necessary to become a party to the proceeding.

eFiling is encouraged. More detailed information relating to filing requirements, interventions, protests, service, and qualifying facilities filings can be found at: *http://www.ferc.gov/docs-filing/efiling/filing-req.pdf.* For other information, call (866) 208–3676 (toll free). For TTY, call (202) 502–8659.

Dated: March 2, 2021.

Nathaniel J. Davis, Sr.,

Deputy Secretary.

[FR Doc. 2021–04752 Filed 3–5–21; 8:45 am]

BILLING CODE 6717–01–P

DEPARTMENT OF ENERGY

Federal Energy Regulatory Commission

[Docket No. EL21–48–000]

Tucson Electric Power Company; Notice of Institution of Section 206 Proceeding and Refund Effective Date

On March 1, 2021, the Commission issued an order in Docket No. EL21–48–000, pursuant to section 206 of the Federal Power Act (FPA), 16 U.S.C. 824e, instituting an investigation to determine whether Tucson Electric Power Company's (Tucson Electric) proposed market-based rates in the Tucson Electric balancing authority area are unjust and unreasonable. *Tucson Electric Power Company,* 174 FERC ¶ 61,165 (2021).

The refund effective date in Docket No. Docket No. EL21–48–000, established pursuant to section 206(b) of the FPA, will be the date of publication of this notice in the **Federal Register**.

Any interested person desiring to be heard in Docket No. EL21–48–000 must file a notice of intervention or motion to intervene, as appropriate, with the Federal Energy Regulatory Commission, in accordance with Rule 214 of the Commission's Rules of Practice and Procedure, 18 CFR 385.214 (2020), within 21 days of the date of issuance of the order.

In addition to publishing the full text of this document in the **Federal Register**, the Commission provides all interested persons an opportunity to view and/or print the contents of this document via the internet through the Commission's Home Page (*http://www.ferc.gov*) using the ''eLibrary'' link. Enter the docket number excluding the last three digits in the docket number field to access the document. At this time, the Commission has suspended access to the Commission's Public Reference Room, due to the proclamation declaring a National Emergency concerning the Novel Coronavirus Disease (COVID–19), issued by the President on March 13, 2020. For assistance, contact FERC at *FERCOnlineSupport@ferc.gov* or call toll-free, (886) 208–3676 or TYY, (202) 502–8659.

The Commission strongly encourages electronic filings of comments, protests and interventions in lieu of paper using the ''eFile'' link at *http://www.ferc.gov.* In lieu of electronic filing, you may submit a paper copy. Submissions sent via the U.S. Postal Service must be addressed to: Kimberly D. Bose, Secretary, Federal Energy Regulatory Commission, 888 First Street NE, Room 1A, Washington, DC 20426. Submissions sent via any other carrier must be addressed to: Kimberly D. Bose, Secretary, Federal Energy Regulatory Commission, 12225 Wilkins Avenue, Rockville, Maryland 20852.

Dated: March 2, 2021.

Nathaniel J. Davis, Sr.,

Deputy Secretary.

[FR Doc. 2021–04751 Filed 3–5–21; 8:45 am]

BILLING CODE 6717–01–P

ENVIRONMENTAL PROTECTION AGENCY

[EPA–HQ–OPP–2017–0720; FRL–10019–47]

Pesticide Registration Review; Draft Human Health and/or Ecological Risk Assessments for Several Pesticides; Notice of Availability

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice.

SUMMARY: This notice announces the availability of EPA's draft human health and/or ecological risk assessments for the registration review of diuron, famoxadone, fluometuron, indaziflam, inorganic chlorates, mancozeb, napropamide, nicarbazin, peroxy compounds, propiconazole, tetraconazole, and zinc pyrithione. In addition, the preliminary work plan for indaziflam is also being published for public comment at this time.

DATES: Comments must be received on or before May 7, 2021.

ADDRESSES: Submit your comments, to the docket identification (ID) number for the specific pesticide of interest provided in the Table in Unit IV, 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, are available at *http://www.epa.gov/dockets.*

Due to the public health concerns related to COVID–19, the EPA Docket Center (EPA/DC) and Reading Room are closed to public 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: For pesticide specific information contact: The Chemical Review Manager for the pesticide of interest identified in the Table in Unit IV.

For general questions 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; telephone number: (703) 305–7106; 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 identified in the Table 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 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.*

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 comprehensive draft human health and/or ecological risk assessments for all pesticides listed in the Table in Unit IV. After reviewing comments received during the public comment period, 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 listed in the Table 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 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. Section 3(g) of FIFRA 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 human health and/or ecological risk assessments for the pesticides shown in the following table and opens a 60-day public comment period on the risk assessments.

TABLE—DRAFT RISK ASSESSMENTS BEING MADE AVAILABLE FOR PUBLIC COMMENT

Registration review case name and No.	Docket ID No.	Chemical review manager and contact information
Diuron, Case 0046	EPA–HQ–OPP–2015–0077	Theodore Varns, *varns.theodore@epa.gov*, (703) 347–8589.
Famoxadone, Case 7038	EPA–HQ–OPP–2015–0094	Christina Scheltema, *scheltema.christina@epa.gov*, (703) 308–2201.
Fluometuron, Case 0049	EPA–HQ–OPP–2015–0746	Ana Pinto, *pinto.ana@epa.gov*, (703) 347–8421.
Indaziflam, Case 7288	EPA–HQ–OPP–2020–0587	Kent Fothergill, *fothergill.kent@epa.gov*, (703) 347–8299.
Inorganic Chlorates, Case 4049	EPA–HQ–OPP–2016–0080	Christian Bongard, *bongard.christian@epa.gov*, (703) 347–0337.
Mancozeb, Case 0643	EPA–HQ–OPP–2015–0291	Alex Hazlehurst, *hazlehurst.alexander@epa.gov*, (703) 347–0221.
Napropamide, Case 2450	EPA–HQ–OPP–2016–0019	Carolyn Smith, *smith.carolyn@epa.gov*, (703) 347–8325.
Nicarbazin, Case 7628	EPA–HQ–OPP–2015–0101	Samantha Thomas, *thomas.samantha@epa.gov*, (703) 347–0514.
Peroxy Compounds, Case 6059	EPA–HQ–OPP–2009–0546	Joseph Mabon, *mabon.joseph@epa.gov*, (703) 347–0177.
Propiconazole, Case 3125	EPA–HQ–OPP–2015–0459	Anna Romanovsky, *romanovsky.anna@epa.gov*, (703) 347–0203.
Tetraconazole, Case 7043	EPA–HQ–OPP–2015–0061	Veronica Dutch, *dutch.veronica@epa.gov*, (703) 308–8585.
Zinc Pyrithione, Case 2480	EPA–HQ–OPP–2014–0158	Michael McCarroll, *mccarroll.michael@epa.gov*, (703) 347–0147.

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/or ecological risk assessments for the pesticides listed in the Table in Unit IV. 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.

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: March 2, 2021.

Mary Reaves,

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

[FR Doc. 2021–04706 Filed 3–5–21; 8:45 am]

BILLING CODE 6560–50–P

ENVIRONMENTAL PROTECTION AGENCY

[FRL–10020–80–Region 5]

Public Meeting for Great Lakes Advisory Board

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice of public meeting for Great Lakes Advisory Board.

SUMMARY: Pursuant to the Federal Advisory Committee Act (FACA), the Environmental Protection Agency (EPA) is announcing a public meeting of the Great Lakes Advisory Board on March 30th, 2021 from 1:00 p.m. to 4:00 p.m. Central Standard Time and March 31st from 9:00 a.m. to 12:00 p.m. Central Standard Time with remote participation only.

DATES: This virtual public meeting will be held on March 30th, 2021 from 1:00 p.m. to 4:00 p.m. Central Standard Time and March 31st from 9:00 a.m. to 12:00 p.m. Central Standard Time. Members of the public seeking to view the meeting (but not provide oral comments) must register by 3:00 p.m. Central Standard Time on March 26th, 2021. Members of the public seeking to make oral comments during the virtual meeting must register and indicate their request to make public comments by contacting the Designated Federal Officer (DFO) directly by 3:00 p.m. Central Standard Time on March 21st, 2021 to be placed on a list of registered commenters and receive special instructions for participation. For information on how to register, please see [How do I participate in the meeting] below.

FOR FURTHER INFORMATION CONTACT: Edlynzia Barnes, Designated Federal Officer (DFO), at *barnes.Edlynzia@epa.gov* or 312–886–6249.

SUPPLEMENTARY INFORMATION:

I. General Information

The GLAB is chartered in accordance with the Federal Advisory Committee Act (FACA) of 1972 (5 U.S.C., Appendix 2, as amended) and 41 CFR 102–3.50(d). The Advisory Board provides advice and recommendations on matters related to the Great Lakes Restoration Initiative. The Advisory Board also advises on domestic matters related to implementation of the Great Lakes Water Quality Agreement between the U.S. and Canada. The major objectives are to provide advice and recommendations on: Great Lakes protection and restoration activities; long-term goals, objectives, and priorities for Great Lakes protection and restoration; and other issues identified by the Great Lakes Interagency Task Force/Regional Working Group.

II. How do I participate in the remote public meeting?

A. Remote Meeting

This meeting will be conducted as a virtual meeting on March 30th, 2021 from 1:00 p.m. to 4:00 p.m. Central Standard Time and March 31st from 9:00 a.m. to 12:00 p.m. Central Standard Time. You must register by 3:00 p.m. Central Standard Time on March 26th, 2021 to receive information on how to participate. You may also submit written or oral comments for the committee by contacting the DFO directly per the processes outlined below.

B. Registration

To register and receive information on how to attend this virtual meeting, please send an email to the DFO at *barnes.edlynzia@epa.gov* with the SUBJECT line of ''Request to Register for March 2021 GLAB Meeting'' and include the following information: Name, Title, Organization, Email, and Phone Number. Attendees must register by 3:00 p.m. Central Standard Time on March 26th, 2021 to receive instructions for participation.

C. Procedures for Providing Public Comments

Oral Statements: In general, oral comments at this virtual conference will be limited to the Public Comments portions of the meeting agenda. Members of the public may provide oral comments limited to up to three minutes per individual or group and submit further information in written comments. Persons interested in providing oral statements should contact the DFO directly at *barnes.edlyzia@epa.gov* by 3:00 p.m. Central Standard Time on March 21st, 2021 with the SUBJECT line of ''Request to Register for March 2021 GLAB Meeting—Provide Oral Statement'' to be placed on the list of registered speakers and receive special instructions for participation. The following information should be included in the email: Name, Title, Organization, Email, and Phone Number. Oral commenters will be provided an opportunity to speak in the order in which their request was received by the DFO and to the extent permitted by the number of comments and the scheduled length of the meeting. Persons not able to provide oral comments during the meeting, will be given an opportunity to provide written comments after the meeting.

Written Statements: Persons interested in providing written statements pertaining to this committee meeting may email them to the DFO prior to 3:00 p.m. Central Standard Time on March 21st, 2021 with the SUBJECT line of ''Request to Register for March 2021 GLAB Meeting—Provide A Written Statement''. The following information should be included in the email: Name, Title, Organization, Email, and Phone Number.

D. Availability of Meeting Materials

The meeting agenda and other materials for the virtual conference will be posted on the GLAB website at *www.glri.us/glab.*