

2. *EPA Registration Numbers:* 264–824 and 264–825. *Docket ID number:* EPA–HQ–OPP–2015–0327. *Applicant:* Bayer CropScience LP, 2 T.W. Alexander Drive, P.O. Box 12014, RTP, NC 27709. *Active ingredient:* Prothioconazole. *Product type:* Fungicide. *Proposed Use:* Sorghum seed treatment. *Contact:* RD.

3. *EPA File Symbol:* 89825–R. *Docket ID number:* EPA–HQ–OPP–2015–0328. *Applicant:* Barnacle-Blocker, LLC., 12907 Yacht Club Place, Cortez, FL 34215. *Active ingredient:* Capsaicin. *Product type:* Antifoulant. *Proposed Use:* Boats. *Contact:* AD.

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

Dated: June 3, 2015.

**Jennifer McInain,**

*Acting, Director, Antimicrobials Division, Office of Pesticide Programs.*

[FR Doc. 2015–15180 Filed 6–18–15; 08:45 am]

**BILLING CODE 6560–50–P**

## ENVIRONMENTAL PROTECTION AGENCY

[EPA–HQ–OPPT–2015–0236; FRL–9928–94]

### Agency Information Collection Activities; Proposed Renewal and Comment Request; TSCA Section 8(a) Preliminary Assessment Information Rule (PAIR)

**AGENCY:** Environmental Protection Agency (EPA).

**ACTION:** Notice.

**SUMMARY:** In compliance with the Paperwork Reduction Act (PRA), this document announces that EPA is planning to submit an Information Collection Request (ICR) to the Office of Management and Budget (OMB). The ICR, entitled: “TSCA Section 8(a) Preliminary Assessment Information Rule (PAIR)” and identified by EPA ICR No. 0586.13 and OMB Control No. 2070–0054, represents the renewal of an existing ICR that is scheduled to expire on March 31, 2016. Before submitting the ICR to OMB for review and approval, EPA is soliciting comments on specific aspects of the proposed information collection that is summarized in this document. The ICR and accompanying material are available in the docket for public review and comment.

**DATES:** Comments must be received on or before August 18, 2015.

**ADDRESSES:** Submit your comments, identified by docket identification (ID) number EPA–HQ–OPPT–2015–0236, 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:* Document Control Office (7407M), Office of Pollution Prevention and Toxics (OPPT), Environmental Protection Agency, 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:** *For technical information contact:* Mike Mattheisen, Chemical Control Division (7405 M), Office of Pollution Prevention and Toxics, Environmental Protection Agency, 1200 Pennsylvania Ave. NW., Washington, DC 20460–0001; telephone number: (202) 564–3077; email address: [mattheisen.mike@epa.gov](mailto:mattheisen.mike@epa.gov).

*For general information contact:* The TSCA–Hotline, ABVI–Goodwill, 422 South Clinton Ave., Rochester, NY 14620; telephone number: (202) 554–1404; email address: [TSCA-Hotline@epa.gov](mailto:TSCA-Hotline@epa.gov).

#### SUPPLEMENTARY INFORMATION:

##### I. What Information is EPA Particularly Interested in?

Pursuant to PRA section 3506(c)(2)(A) (44 U.S.C. 3506(c)(2)(A)), EPA specifically solicits comments and information to enable it to:

1. Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the Agency, including whether the information will have practical utility.
2. Evaluate the accuracy of the Agency’s estimates of the burden of the proposed collection of information, including the validity of the methodology and assumptions used.
3. Enhance the quality, utility, and clarity of the information to be collected.
4. Minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated electronic, mechanical, or other technological collection techniques or other forms of information technology, *e.g.*, permitting electronic submission of responses. In particular, EPA is requesting comments from very small businesses (those that employ less than 25) on examples of specific additional efforts that EPA

could make to reduce the paperwork burden for very small businesses affected by this collection.

##### II. What Information Collection Activity or ICR Does this Action Apply to?

*Title:* TSCA Section 8(a) Preliminary Assessment Information Rule (PAIR).

*ICR number:* EPA ICR No. 0586.13.

*OMB control number:* OMB Control No. 2070–0054.

*ICR status:* This ICR is currently scheduled to expire on March 31, 2016. An Agency may not conduct or sponsor, and a person is not required to respond to, a collection of information, unless it displays a currently valid OMB control number. The OMB control numbers for EPA’s regulations in title 40 of the Code of Federal Regulations (CFR), after appearing in the **Federal Register** when approved, are listed in 40 CFR part 9, are displayed either by publication in the **Federal Register** or by other appropriate means, such as on the related collection instrument or form, if applicable. The display of OMB control numbers for certain EPA regulations is consolidated in 40 CFR part 9.

*Abstract:* Section 8(a) of the Toxic Substances Control Act (TSCA) authorizes EPA to promulgate rules under which manufacturers, importers and processors of chemical substances and mixtures must maintain records and submit reports to EPA. EPA has promulgated the Preliminary Assessment Information Rule (PAIR) under TSCA section 8(a). EPA uses PAIR to collect information to identify, assess and manage human health and environmental risks from chemical substances, mixtures and categories. PAIR requires chemical manufacturers and importers to complete a standardized reporting form to help evaluate the potential for adverse human health and environmental effects caused by the manufacture or importation of identified chemical substances, mixtures or categories. Chemicals identified by EPA or any other federal agency, for which a justifiable information need for production, use or exposure-related data can be satisfied by the use of the PAIR are proper subjects for TSCA section 8(a) PAIR rulemaking. In most instances the information that EPA receives from a PAIR report is sufficient to satisfy the information need in question. This information collection addresses the reporting and recordkeeping requirements associated with TSCA section 8(a).

Responses to the collection of information are mandatory (see 40 CFR parts 712, 766, and 792). Respondents

may claim all or part of a response confidential. EPA will disclose information that is covered by a claim of confidentiality only to the extent permitted by, and in accordance with, the procedures in TSCA section 14 and 40 CFR part 2.

**Burden statement:** The annual public reporting and recordkeeping burden for this collection of information is estimated to average 31.5 hours per response. Burden is defined in 5 CFR 1320.3(b).

The ICR, which is available in the docket along with other related materials, provides a detailed explanation of the collection activities and the burden estimate that is only briefly summarized here:

**Respondents/Affected Entities:**

Entities potentially affected by this ICR are companies that manufacture, process or import chemical substances, mixtures or categories.

**Estimated total number of potential respondents:** 1.

**Frequency of response:** On occasion.

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

**Estimated total annual burden hours:** 31.5 hours.

**Estimated total annual costs:** \$2,388. This includes an estimated burden cost of \$2,388 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 916 hours in the total estimated respondent burden compared with that identified in the ICR currently approved by OMB. This decrease reflects additional both adjustment changes from a reduction in the assumed number of PAIR reports filed annually, and program changes resulting from mandatory electronic submissions of PAIR reports. In recent years (FY 2011–FY 2014), EPA has received no PAIR submissions and, for the purposes of this analysis, EPA assumes an annual rate of one submission per year. At the time OMB last renewed this ICR, EPA estimated an average of 33 reports from 14.8 submitters based on fiscal year 2006–2010 data. The ICR supporting statement provides a detailed analysis of the change in burden estimate. This change is both an adjustment and a program change.

### 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: June 10, 2015.

**James Jones,**

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

[FR Doc. 2015–14946 Filed 6–18–15; 8:45 am]

**BILLING CODE 6560–50–P**

## ENVIRONMENTAL PROTECTION AGENCY

[EPA–HQ–OPPT–2015–0305; FRL–9928–69]

### Use of High Throughput Assays and Computational Tools; Endocrine Disruptor Screening Program; Notice of Availability and Opportunity for Comment

**AGENCY:** Environmental Protection Agency (EPA).

**ACTION:** Notice.

**SUMMARY:** This document describes how EPA is planning to incorporate an alternative scientific approach to screen chemicals for their ability to interact with the endocrine system. This will improve the Agency's ability to fulfill its statutory mandate to screen pesticide chemicals and other substances for their ability to cause adverse effects by their interaction with the endocrine system. The approach incorporates validated high throughput assays and a computational model and, based on current research, can serve as an alternative for some of the current assays in the Endocrine Disruptor Screening Program (EDSP) Tier 1 battery. EPA has partial screening results for over 1800 chemicals that have been evaluated using high throughput assays and a computational model for the estrogen receptor pathway. In the future, EPA anticipates that additional alternative methods will be available for EDSP chemical screening based on further advancements of high throughput assays and computational models for other endocrine pathways. Use of these alternative methods will accelerate the pace of screening, decrease costs, and reduce animal testing. In addition, this approach advances the goal of providing sensitive, specific, quantitative, and

efficient screening using alternative test methods to some assays in the Tier 1 battery to protect human health and the environment.

**DATES:** Comments must be received on or before August 18, 2015.

**ADDRESSES:** Submit your comments, identified by docket identification (ID) number EPA–HQ–OPPT–2015–0305, 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:** Document Control Office (7407M), Office of Pollution Prevention and Toxics (OPPT), Environmental Protection Agency, 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:** For technical information contact: Jane Robbins, Office of Science Coordination and Policy (OSCP), Office of Chemical Safety and Pollution Prevention, Environmental Protection Agency, 1200 Pennsylvania Ave. NW., Washington, DC 20460–0001; telephone number: (202) 564–6625; email address: [robbins.jane@epa.gov](mailto:robbins.jane@epa.gov).

For general information contact: The TSCA–Hotline, ABVI–Goodwill, 422 South Clinton Ave., Rochester, NY 14620; telephone number: (202) 554–1404; email address: [TSCA-Hotline@epa.gov](mailto:TSCA-Hotline@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 those interested in endocrine testing of chemicals (including pesticides), and the EDSP in general. 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 is the agency authority for taking this action?

The EDSP is established under section 408(p) of the Federal Food, Drug and