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.*

C. How can I get copies of this document and other related information?

A copy of the draft PR notice is available in the docket under docket identification (ID) number EPA–HQ–OPP–2016–0671.

II. What guidance does this PR Notice provide?

This draft PR Notice provides guidance to the registrant concerning the process for notifications, non-notifications and minor formulation amendments. This proposed notice updates and clarifies the scope of changes accepted by notification, non-notification and minor formulation amendments for all pesticide products. This notice supersedes both PR Notices 95–2 and 98–10 in their entirety. As per 40 CFR 152.46, EPA may determine that certain minor modifications to a registration having no potential to cause unreasonable adverse effects to the environment may be accomplished by notification or without notification to the Agency. Since the issuance of PR Notice 98–10, various regulatory and statutory changes have taken place. In particular, the Pesticide Registration Improvement Act (PRIA), the Pesticide Registration Improvement Renewal Act (PRIA 2), Pesticide Registration Improvement Extension Act (PRIA 3), and pending Pesticide Registration Enhancement Act of 2017 (PRIA 4) has resulted in a need for EPA to revise the notification procedures. Certain actions previously accepted under PR Notice 98–10 are now actions scheduled by the PRIA action tables. EPA is issuing this notice to align the notification program with the requirements of the Food Quality Protection Act (FQPA) and PRIA and to clarify the processes for accepting minor, low risk registration amendments to be accomplished through notification, non-notification or as accelerated amendments, previously established in PR Notice 98–10. EPA believes these changes will be useful to registrants as it presents a clarified and consolidated explanation for accomplishing these registration changes. No significant impacts or costs are expected as a result of this proposed PR Notice. However, the Agency is especially requesting impacted parties to provide through comments available information on projected cost implications of this draft updated guidance. The Paperwork Reduction Act (PRA) burdens associated with revisions to the PR Notice are accounted for in the current ICR entitled: Application for New and Amended Pesticide Registration, OMB ICR 2070–0060; EPA No. 0277.17. As noted above, no increase or decrease in the current PRA burden inventory is anticipated.

III. Do PR Notices contain binding requirements?

The PR Notice discussed in this notice is intended to provide guidance to EPA personnel and decision makers and to pesticide registrants. While the requirements in the statutes and Agency regulations are binding on EPA and the applicants, this PR Notice is not binding on either EPA or pesticide registrants, and EPA may depart from the guidance where circumstances warrant and without prior notice. Likewise, pesticide registrants may assert that the guidance is not appropriate generally or not applicable to a specific pesticide or situation.

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

Dated: August 2, 2017.

Richard P. Keigwin, Jr.,

Director, Office of Pesticide Programs.

[FR Doc. 2017–18765 Filed 9–5–17; 8:45 am]

BILLING CODE 6560–50–P

ENVIRONMENTAL PROTECTION AGENCY

[FRL–9967–00–OA]

Notification of a Public Meeting of the Science Advisory Board Chemical Assessment Advisory Committee

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice.

SUMMARY: The Environmental Protection Agency (EPA or Agency) Science Advisory Board (SAB) Staff Office announces a public meeting of the SAB Chemical Assessment Advisory Committee (CAAC) to receive a briefing from the EPA's National Center for Environmental Assessment (NCEA) on the content and presentation of assessment products to be released at early stages of development of draft assessments. These products represent an update to the materials released for the purposes of early stakeholder engagement, as outlined in the ''IRIS enhancements'' (2013). These materials are expected to add transparency to draft assessment development, while simultaneously increasing throughput and responsiveness to Agency needs.

DATES: The public face-to-face meeting will be held from Wednesday, September 27, 2017 through Thursday, September 28, 2017, from 9:00 a.m. to 5:00 p.m., (Eastern time) daily.

ADDRESSES: The public meeting will be held at Residence Inn Arlington Capital View, 2850 S. Potomac Ave., Arlington, VA 22202.

FOR FURTHER INFORMATION CONTACT: Any member of the public wishing to obtain further information concerning the meeting may contact Dr. Suhair Shallal, Designated Federal Officer (DFO), SAB Staff Office, by telephone at (202) 564–0257 or *shallal.suhair@epa.gov.* General information concerning the EPA Science Advisory Board, as well as any updates concerning the meeting announced in this notice, can be found at the EPA SAB Web site at *http://www.epa.gov/sab.*

SUPPLEMENTARY INFORMATION:

Background: The SAB was established pursuant to the Environmental Research, Development, and Demonstration Authorization Act (ERDAA) codified at 42 U.S.C. 4365, to provide independent scientific and technical advice to the Administrator on the technical basis for Agency positions and regulations. The SAB is a Federal Advisory Committee chartered under the Federal Advisory Committee Act (FACA), 5 U.S.C., App. 2. The SAB will comply with the provisions of FACA and all appropriate SAB Staff Office procedural policies. Pursuant to FACA and EPA policy, notice is hereby given that the SAB CAAC will hold a public meeting to receive a briefing from the EPA's National Center for Environmental Assessment (NCEA) on the content and presentation of assessment products to be released at early stages of draft development. These products represent an update to the materials released for the purposes of early stakeholder engagement, as outlined in the ''IRIS enhancements'' (2013). These materials are expected to add transparency to draft assessment development, while simultaneously increasing throughput and responsiveness to Agency needs. The CAAC will provide advice to the Administrator through the chartered SAB.

The NCEA continues to incorporate improvements in response to recommendations from the National Research Council and the SAB to (1) improve the scientific integrity of assessments; (2) improve the

productivity of the program; and (3) increase transparency so issues are identified early in the process. Information about this program is available at: *https://www.epa.gov/iris.*

Availability of Meeting Materials: Additional background on this SAB activity, the meeting agenda, and other materials for the meeting will be posted on the SAB Web site at *http://www.epa.gov/sab.*

Procedures for Providing Public Input: Public comment for consideration by EPA's federal advisory committees and panels has a different purpose from public comment provided to EPA program offices. Therefore, the process for submitting comments to a federal advisory committee is different from the process used to submit comments to an EPA program office. Federal advisory committees and panels, including scientific advisory committees, provide independent advice to the EPA. Members of the public can submit relevant comments pertaining to the meeting materials or the group conducting this SAB activity. Input from the public to the SAB will have the most impact if it consists of comments that provide specific scientific or technical information or analysis for SAB committees and panels to consider or if it relates to the clarity or accuracy of the technical information. Members of the public wishing to provide comment should contact the DFO directly.

Oral Statements: In general, individuals or groups requesting an oral presentation at a public meeting will be limited to five minutes per speaker. Interested parties wishing to provide comments should contact Dr. Suhair Shallal, DFO (preferably via email) at the contact information noted above by September 20, 2017, to be placed on the list of public speakers for the meeting.

Written Statements: Written statements will be accepted throughout the advisory process; however, for timely consideration by Committee members, statements should be supplied to the DFO (preferably via email) at the contact information noted above by September 20, 2017. It is the SAB Staff Office general policy to post written comments on the Web page for the advisory meeting. Submitters are requested to provide an unsigned version of each document because the SAB Staff Office does not publish documents with signatures on its Web sites. Members of the public should be aware that their personal contact information, if included in any written comments, may be posted to the SAB Web site. Copyrighted material will not be posted without explicit permission of the copyright holder.

Accessibility: For information on access or services for individuals with disabilities, please contact Dr. Suhair Shallal at (202) 564–0257 or at *shallal.suhair@epa.gov.* To request accommodation of a disability, please contact Dr. Shallal preferably at least ten days prior to the meeting, to give EPA as much time as possible to process your request.

Dated: August 21, 2017.

Christopher Zarba,

Director, EPA Science Advisory Board Staff Office.

[FR Doc. 2017–18764 Filed 9–5–17; 8:45 am]

BILLING CODE 6560–50–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60-Day–17–17AUQ; Docket No. CDC–2017–0064]

Proposed Data Collection Submitted for Public Comment and Recommendations

AGENCY: Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

ACTION: Notice with comment period.

SUMMARY: The Centers for Disease Control and Prevention (CDC), as part of its continuing efforts to reduce public burden and maximize the utility of government information, invites the general public and other Federal agencies to take this opportunity to comment on proposed and/or continuing information collections, as required by the Paperwork Reduction Act of 1995. This notice invites comment on Mobile Proximity Initial User Feedback information collection project.

DATES: Written comments must be received on or before November 6, 2017.

ADDRESSES: You may submit comments, identified by Docket No. CDC–2017–0064 by any of the following methods:

- *Federal eRulemaking Portal: Regulations.gov.* Follow the instructions for submitting comments.
- *Mail:* Leroy A. Richardson, Information Collection Review Office, Centers for Disease Control and Prevention, 1600 Clifton Road NE., MS–D74, Atlanta, Georgia 30329.

Instructions: All submissions received must include the agency name and Docket Number. All relevant comments received will be posted without change to *Regulations.gov*, including any personal information provided. For access to the docket to read background documents or comments received, go to *Regulations.gov*.

Please note: All public comment should be submitted through the Federal eRulemaking portal (*Regulations.gov*) or by U.S. mail to the address listed above.

FOR FURTHER INFORMATION CONTACT: To request more information on the proposed project or to obtain a copy of the information collection plan and instruments, contact Leroy A. Richardson, Information Collection Review Office, Centers for Disease Control and Prevention, 1600 Clifton Road NE., MS–D74, Atlanta, Georgia 30329; phone: 404–639–7570; Email: *omb@cdc.gov.*

SUPPLEMENTARY INFORMATION: Under the Paperwork Reduction Act of 1995 (PRA) (44 U.S.C. 3501–3520), Federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. In addition, the PRA also requires Federal agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each new proposed collection, each proposed extension of existing collection of information, and each reinstatement of previously approved information collection before submitting the collection to OMB for approval. To comply with this requirement, we are publishing this notice of a proposed data collection as described below.

Comments are invited on: (a) Whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) ways to enhance the quality, utility, and clarity of the information to be collected; (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology; and (e) estimates of capital or start-up costs and costs of operation, maintenance, and purchase of services to provide information. Burden means the total time, effort, or financial resources expended by persons to generate, maintain, retain, disclose or provide information to or for a Federal agency. This includes the time needed to review instructions; to develop, acquire, install and utilize technology and systems for the purpose of