TABLE 1.—METHIDATHION PRODUCT CANCELLATIONS—Continued

Registration Number	Product Name	Chemical Name
10163-OR- 00-0010	Supracide 2E Insecti- cide	Methidathion
10163-OR- 02-0018	Supracide 2E Insecti- cide	Methidathion
10163-TX- 05-0003	Supracide 2E Insecti- cide	Methidathion
10163-UT- 00-0006	Supracide 2E Insecti- cide	Methidathion
10163-WA- 00-0006	Supracide 2E Insecti- cide	Methidathion
10163-WY- 05-0001	Supracide 2E Insecti- cide	Methidathion

Table 2 of this unit includes the names and addresses of record for all registrants of the products in Table 1 of this unit in sequence by EPA company number. This number corresponds to the first part of the EPA registration numbers of the products listed in Table 1 of this unit.

TABLE 2.—REGISTRANTS OF CANCELLED PRODUCTS

EPA Company Number	Company Name and Ad- dress
000100	Syngenta Crop Protec- tion, Inc., P. O. Box 18300, Greensboro, NC 27419–8300
010163	Gowan Company, P. O. Box 5569, Yuma, AZ 85366–5569

III. Summary of Public Comments Received and Agency Response to Comments

During the public comment period provided, EPA received no comments in response to the April 7, 2010 **Federal Register** notice (75 FR 17735; FRL– 8819–1) announcing the Agency's receipt of the requests for voluntary cancellations of products listed in Unit II., Table 1.

IV. Cancellation Order

Pursuant to FIFRA section 6(f), EPA hereby approves the requested cancellations of methidathion registrations identified in Unit II., Table 1. Accordingly, the Agency hereby orders that the product registrations identified in Unit II., Table 1 are canceled. The effective date of the cancellations that are the subject of this notice is December 30, 2012. Any distribution, sale, or use of existing stocks of the products identified in Unit II., Table 1, in a manner inconsistent with any of the provisions for disposition of existing stocks set forth in Unit VI. of this notice will be a violation of FIFRA.

V. What is the Agency's Authority for Taking this Action?

Section 6(f)(1) of FIFRA provides that a registrant of a pesticide product may, at any time, request that any of its pesticide registrations be canceled or amended to terminate one or more uses. FIFRA further provides that, before acting on the request, EPA must publish a notice of receipt of the request in the **Federal Register**. Thereafter, following the public comment period, the EPA Administrator may approve such a request. The notice of receipt for this action was published for comment on April 7, 2010. The comment period closed on May 7, 2010.

VI. Provisions for Disposition of Existing Stocks

Existing stocks are those stocks of registered pesticide products which are currently in the United States and were packaged, labeled, and released for shipment prior to the effective date of the action. The existing stock provisions for the products subject to this order are as follows:

After December 31, 2012, registrants are prohibited from selling or distributing existing stocks of products containing methidathion labeled for all uses.

After December 31, 2014, persons other than registrants are prohibited from selling or distributing existing stocks of products containing methidathion labeled for all uses.

After December 31, 2014, existing stocks of products containing methidathion labeled for all uses, already in the hands of users can be used legally until they are exhausted, provided that the use complies with the EPA approved label and labeling of the affected product.

List of Subjects

Environmental protection, Pesticides and pests.

Dated: May 21, 2010. Richard P. Keigwin, Jr., Director, Pesticide Re-evaluation Division, Office of Pesticide Programs.

[FR Doc. 2010–12925 Filed 6–1–10; 8:45 am] BILLING CODE 6560–50–S

ENVIRONMENTAL PROTECTION AGENCY

[FRL-9157-3; Docket ID No. EPA-HQ-ORD-2010-0396]

Draft Toxicological Review of Formaldehyde in Support of Summary Information on the Integrated Risk Information System (IRIS)

AGENCY: Environmental Protection Agency (EPA). **ACTION:** Notice of public comment period and listening session.

SUMMARY: EPA is announcing a 90-day public comment period and a public listening session for the external review draft human health assessment titled, "Toxicological Review of Formaldehyde Inhalation Assessment: In Support of Summary Information on the Integrated Risk Information System (IRIS)" (EPA/ 635/R-10/002C). The draft assessment was prepared by the National Center for Environmental Assessment (NCEA) within the EPA Office of Research and Development (ORD). EPA is releasing this draft assessment solely for the purpose of pre-dissemination peer review under applicable information quality guidelines. This draft assessment has not been formally disseminated by EPA. It does not represent and should not be construed to represent any Agency policy or determination. A committee of the National Research Council, acting under the auspices of National Academy of Sciences (NAS), will conduct an independent scientific peer review of the EPA draft human health assessment of formaldehyde. The peer review committee will hold meetings, some of which may involve public sessions. Public sessions will be announced before each meeting on the National Academies Web site (http:// www8.nationalacademies.org/cp/ projectview.aspx?kev=49207). The public comment period and NAS scientific peer review are separate processes that provide opportunities for all interested parties to comment on the assessment. Due to the timing of the NAS peer review meetings, one or more NAS meetings may take place before the close of EPA's public comment period (see DATES below). For NAS meetings that occur during the public comment period, EPA will provide all public

comments to the NAS at least 5 working days before the meeting date announced on the National Academies website. All comments provided to the EPA during the public comment period will inform the Agency's revision of the draft assessment.

An EPA listening session will be held on July 27, during the public comment period for this draft assessment. The purpose of the listening session is to allow all interested parties to present scientific and technical comments on draft IRIS health assessments to EPA and other interested parties attending the listening session. EPA welcomes the comments that will be provided to the Agency by the listening session participants. The comments will be considered by the Agency as it revises the draft assessment after the NAS external peer review. If listening session participants wish EPA to share their comments with the peer review committee before their meeting, they should also submit written comments at least 5 working days before the meeting date announced on the National Academies website, using the detailed and established procedures described in the SUPPLEMENTARY INFORMATION section of this notice.

DATES: The public comment period begins June 2, 2010, and ends August 31, 2010. Comments should be in writing and must be received by EPA by August 31, 2010.

The listening session on the draft assessment for formaldehyde will be held on July 27, beginning at 9 a.m. and ending at 4 p.m., Eastern Daylight Time. Interested parties who wish to attend the listening session should register no later than July 20. If you wish to present at the listening session, indicate in your registration that you would like to present oral comments and provide the length of your presentation. To register send an e-mail to

IRISListeningSession@epa.gov (subject line: Formaldehyde Listening Session); call Christine Ross at 703–347–8592; or fax a registration request to 703–347– 8689. Please reference the "Formeldehyde Listening Secsion" and

"Formaldehyde Listening Session" and include your name, title, affiliation, full address and contact information. Indicate if you will need audio-visual equipment (*e.g.*, laptop computer and slide projector). In general, each presentation should be no more than 30 minutes. If, however, there are more requests for presentations than the allotted time allows, then the time limit for each presentation will be adjusted. A copy of the agenda for the listening session will be available at the meeting. If no speakers have registered by July 20, the listening session will be cancelled, and EPA will notify those registered of the cancellation.

ADDRESSES: The draft "Toxicological Review of Formaldehyde Inhalation Assessment: In Support of Summary Information on the Integrated Risk Information System (IRIS)" is available primarily via the Internet on the NCEA home page under the Recent Additions and Publications menus at http:// www.epa.gov/ncea. A limited number of paper copies are available from the Information Management Team (Address: Information Management Team, National Center for Environmental Assessment (Mail Code: 8601P), U.S. Environmental Protection Agency, 1200 Pennsylvania Avenue, NW., Washington, DC 20460; telephone: 703-347-8561; facsimile: 703-347-8691). If you request a paper copy, please provide your name, mailing address, and the draft assessment title.

Comments may be submitted electronically via *http:// www.regulations.gov*, by e-mail, by mail, by facsimile, or by hand delivery/ courier. Please follow the detailed instructions provided in the **SUPPLEMENTARY INFORMATION** section of this notice.

The listening session on the draft assessment for formaldehyde will be held at the EPA offices at Potomac Yard North Building, N-4830, 2733 South Crystal Drive, Arlington, Virginia 22202. Please note that to gain entrance to this EPA building to attend the Formaldehyde Listening Session, you must have photo identification and must register at the guard's desk in the lobby. The guard will retain your photo identification and will provide you with a visitor's badge. At the guard's desk, you should provide the name Christine Ross and the telephone number 703-347–8592 to the guard on duty. The guard will contact Ms. Ross who will meet you in the reception area to escort you to the meeting room. When you leave the building, please return your visitor's badge to the guard and you will receive your photo identification.

A teleconference line will also be available for registered attendees/ speakers. The teleconference number is 866–299–3188, and the access code is 926–378–7897, followed by the pound sign (#). The teleconference line will be activated at 8:45 a.m., and you will be asked to identify yourself and your affiliation at the beginning of the call.

Information on Services for Individuals with Disabilities: EPA welcomes public attendance at the Formaldehyde Listening Session and will make every effort to accommodate persons with disabilities. For information on access or services for individuals with disabilities, please contact Christine Ross by phone at 703– 347–8592 or by e-mail at IRISListeningSession@epa.gov. To request accommodation for a disability, please contact Ms. Ross, preferably at least 10 days prior to the meeting, to give EPA as much time as possible to process your request.

Additional Information: For information on the docket, www.regulations.gov, or the public comment period, please contact the Office of Environmental Information (OEI) Docket (Mail Code: 2822T), U.S. Environmental Protection Agency, 1200 Pennsylvania Ave., NW., Washington, DC 20460; telephone: 202–566–1752; facsimile: 202–566–1753; or e-mail: *ORD.Docket@epa.gov.*

For information on the Formaldehyde Listening Session, please contact Christine Ross, IRIS Staff, National Center for Environmental Assessment (Mail Code: 8601P), U.S. Environmental Protection Agency, 1200 Pennsylvania Avenue, NW., Washington, DC 20460; telephone: 703–347–8592; facsimile: 703–347–8689; or e-mail: *IRISListeningSession@epa.gov.*

For information on the draft assessment, please contact John Whalan, National Center for Environmental Assessment (Mail Code: 8601P), U.S. Environmental Protection Agency, 1200 Pennsylvania Avenue, NW., Washington, DC 20460; telephone: 703-347–8639; facsimile: 703–347–8689; or e-mail: *FRN_Questions@epa.gov.* **SUPPLEMENTARY INFORMATION:**

I. Information About IRIS

EPA's IRIS is a human health assessment program that evaluates quantitative and qualitative risk information on effects that may result from exposure to chemical substances found in the environment. Through the IRIS Program, EPA provides the highest quality science-based human health assessments to support the Agency's regulatory activities. The IRIS database contains information for more than 540 chemical substances that can be used to support the first two steps (hazard identification and dose-response evaluation) of the risk assessment process. When supported by available data, IRIS provides oral reference doses (RfDs) and inhalation reference concentrations (RfCs) for chronic noncancer health effects and cancer assessments. Combined with specific exposure information, government and private entities use IRIS to help characterize public health risks of chemical substances in a site-specific

situation and thereby support risk management decisions designed to protect public health.

II. How to Submit Comments to the Docket at http://www.regulations.gov

Submit your comments, identified by Docket ID No. EPA–HQ–ORD–2010– 0396, by one of the following methods:

• http://www.regulations.gov: Follow the on-line instructions for submitting comments.

- E-mail: ORD.Docket@epa.gov.
- Facsimile: 202–566–1753.

• *Mail:* Office of Environmental Information (OEI) Docket (Mail Code: 2822T), U.S. Environmental Protection Agency, 1200 Pennsylvania Ave., NW., Washington, DC 20460. The telephone number is 202–566–1752. If you provide comments by mail, please submit one unbound original with pages numbered consecutively, and three copies of the comments. For attachments, provide an index, number pages consecutively with the comments, and submit an unbound original and three copies.

Hand Delivery: The OEI Docket is located in the EPA Headquarters Docket Center, EPA West Building, Room 3334, 1301 Constitution Ave., NW., Washington, DC. The EPA Docket Center 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. Deliveries are only accepted during the Public Reading Room's normal hours of operation, and special arrangements should be made for deliveries of boxed information. If you provide comments by hand delivery, please submit one unbound original with pages numbered consecutively, and three copies of the comments. For attachments, provide an index, number pages consecutively with the comments, and submit an unbound original and three copies.

Instructions: Direct your comments to Docket ID No. EPA-HQ-ORD-2010-0396. Please ensure that your comments are submitted within the specified comment period. Comments received after the closing date will be marked "late," and may only be considered if time permits. It is EPA's policy to include all comments it receives in the public docket without change and to make the comments available online at http://www.regulations.gov, including any personal information provided, unless comments include information claimed to be Confidential Business Information (CBI) or other information whose disclosure is restricted by statute. Do not submit information that you consider to be CBI or otherwise protected through http://

www.regulations.gov or e-mail. The *http://www.regulations.gov* website is an "anonymous access" system, which means that EPA will not know your identity or contact information unless you provide it in the body of your comments. If you send e-mail comments directly to EPA without going through http://www.regulations.gov, your e-mail address will be automatically captured and included as part of the comments that are placed in the public docket and made available on the Internet. If you submit electronic comments, EPA recommends that you include your name and other contact information in the body of your comments and with any disk or CD-ROM you submit. If EPA cannot read your comments due to technical difficulties and cannot contact you for clarification, EPA may not be able to consider your comments. Electronic files should avoid the use of special characters and any form of encryption and be free of any defects or viruses. For additional information about EPA's public docket, visit the EPA Docket Center homepage at http:// www.epa.gov/epahome/dockets.htm.

Docket: All documents in the docket are listed in the http:// www.regulations.gov index. Although listed in the index, some information is not publicly available, e.g., CBI or other information whose disclosure is restricted by statute. Certain other material, such as copyrighted material, will be publicly available only in hard copy. Publicly available docket materials are available either electronically at http:// www.regulations.gov or in hard copy at the OEI Docket in the EPA Headquarters Docket Center.

Dated: May 25, 2010.

Rebecca Clark,

Acting Director, National Center for Environmental Assessment. [FR Doc. 2010–13097 Filed 6–1–10; 8:45 am]

BILLING CODE 6560-50-P

ENVIRONMENTAL PROTECTION AGENCY

[FRL-9157-8]

Science Advisory Board Staff Office; Notification of Two Public Teleconferences of the SAB Trichloroethylene (TCE) Review Panel

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

SUMMARY: The EPA Science Advisory Board (SAB) Staff Office announces two public teleconferences of the SAB Trichloroethylene (TCE) Review Panel to conduct a follow-up discussion of its review of EPA's *Toxicological Review of Trichloroethylene in Support of Summary Information on the Integrated Risk Information System (IRIS),* External Review Draft.

DATES: There will be a public teleconference on June 24, 2010 from 12:30 to 4:30 p.m. (Eastern Daylight Time) and another public teleconference on August 5, 2010 from 9 to 11:30 a.m. (Eastern Daylight Time).

ADDRESSES: The teleconferences will be conducted by phone only.

FOR FURTHER INFORMATION CONTACT: Any member of the public wishing to obtain information concerning the public teleconferences may contact Dr. Holly Stallworth, Designated Federal Officer (DFO), EPA Science Advisory Board Staff Office (1400F), U.S. Environmental Protection Agency, 1200 Pennsylvania Avenue, NW., Washington, DC 20460; by telephone/voice mail at (202) 343-9867 or via e-mail at stallworth.holly@epa.gov. General information about the SAB, as well as any updates concerning the meeting announced in this notice, may be found on the SAB Web site at http:// www.epa.gov/sab.

SUPPLEMENTARY INFORMATION: Pursuant to the Federal Advisory Committee Act (FACA), 5 U.S.C., App. 2, notice is hereby given that the SAB Trichloroethylene (TCE) Review Panel will hold two public teleconferences to discuss its peer review report to EPA. The SAB was established pursuant to 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 FACA. The SAB will comply with the provisions of FACA and all appropriate SAB Staff Office procedural policies.

The TCE Review Panel met on May 10-12, 2010, to review EPA's Toxicological Review of Trichloroethylene in Support of Summary Information on the Integrated Risk Information System (IRIS), External **Review Draft** [Federal Register Notice dated March 31, 2010 (75 FR 16108-16109)]. Materials from the May meeting are posted on the SAB Web site at http://yosemite.epa.gov/sab/ sabproduct.nsf/MeetingCalBOARD/ BEEA3E70E29DE3A8852 576E3006B8F54?OpenDocument. The purpose of the June 24, 2010 teleconference is to further discuss the Panel's review of EPA's draft document. The purpose of the August 5, 2010