statement is $2.3 million over the 3 year reporting period. See Table 5 and Section 6(b).

The estimated incremental burden cost provided in this ICR is more consistent with the dollar burden estimates provided in the 3 commenters than the previous estimates in the 1997 RIA.

What Is the Next Step in the Process for This ICR?

The EPA will consider the comments received under this notice 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. At that time, EPA will issue another **Federal Register** notice 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**.

Dated: November 6, 2006.

Scott L. Mathias,

Acting Director, Air Quality Policy Division, Office of Air Quality Planning and Standards, Office of Air and Radiation.

[FR Doc. E6–19376 Filed 11–14–06; 8:45 am]

BILLING CODE 6560–50–P

ENVIRONMENTAL PROTECTION AGENCY

[FRL–8242–2]

Clean Air Act Advisory Committee; Notice of Charter Renewal

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice of charter renewal.

The charter for the Environmental Protection Agency's Clean Air Act Advisory Committee (CAAAC) will be renewed for an additional two-year period, as a necessary committee which is in the public interest, in accordance with the provisions of the Federal Advisory Committee Act (FACA), 5 U.S.C. App. 2 section 9(c). The purpose of CAAAC is to provide advice and recommendations to the EPA Administrator on issues associated with policy and technical issues associated with implementation of the Clean Air Act.

It is determined that CAAAC is in the public interest in connection with the performance of duties imposed on the Agency by law.

Inquiries may be directed to Pat Childers, CAAAC Designated Federal Officer, U.S. EPA, Mail Code 6102A, 1200 Pennsylvania Ave., NW., Washington, DC 20460, or by e-mail *childers.pat@epa.gov*.

Dated: November 2, 2006.

William L. Wehrum,

Acting Assistant Administrator, Office of Air and Radiation.

[FR Doc. E6–19282 Filed 11–14–06; 8:45 am]

BILLING CODE 6560–50–P

ENVIRONMENTAL PROTECTION AGENCY

[FRL–8242–3]

Listening Session on Exploring Bottled Water as an Alternative Compliance Option in Limited Situations for Non-Transient, Non-Community Water Systems

AGENCY: Environmental Protection Agency.

ACTION: Notice.

SUMMARY: The U.S. Environmental Protection Agency (EPA) is announcing a listening session on the viability of bottled water as an alternative compliance option for chronic contaminants regulated under the Safe Drinking Water Act (SDWA). The purpose of this meeting is to identify information and data needed for EPA to evaluate the efficacy of bottled water as an alternative compliance option for non-transient, non-community water systems.

DATES: The listening session will be held in Washington, DC, on Tuesday, December 12, 2006, from 8:30 a.m. to 5 p.m. Registration will open at 8 a.m.

ADDRESSES: The listening session will take place at RESOLVE, Inc., 1255 23rd St., NW., Suite 275, Washington, DC 20037.

FOR FURTHER INFORMATION CONTACT: Interested participants from the public should contact Jennifer Moller, U.S. Environmental Protection Agency, Office of Ground Water and Drinking Water, Drinking Water Protection Division (Mail Code 4606M), 1200 Pennsylvania Avenue, NW., Washington, DC 20460. There is no charge for attending this workshop as an observer, but seats are limited, so register as soon as possible. Please contact Jennifer Moller at *Moller.Jennifer@epa.gov* or call 202–564–3891 to receive additional details.

SUPPLEMENTARY INFORMATION:

Background: At the request of the Association of State Drinking Water Administrators (ASDWA), EPA is convening a meeting to discuss information needed to explore whether and in what limited situations bottled water may be a safe and effective alternative compliance option to treatment technology and point-of-use devices. Under the Safe Drinking Water Act (SDWA) bottled water is allowed for use in very limited situations, such as in emergency situations or as a temporary measure under variances and exemptions. There is no statutory prohibition on the use of bottled water to achieve compliance. However, bottled water is prohibited by regulation (40 CFR 141.101) for use by a public water system to achieve compliance with a maximum contaminant level (MCL).

Public Comment: An opportunity for public comment will be provided during the listening session. Oral statements will be limited to five minutes; it is preferred that only one person present the statement on behalf of a group or organization. Written comments may be provided at the meeting or may be sent by mail to Jennifer Moller at the mail or e-mail address listed in the **FOR FURTHER INFORMATION CONTACT** section of this notice.

Special Accommodations: For information on access or services for individuals with disabilities, please contact Jennifer Moller at *Moller.Jennifer@epa.gov*. To request accommodation of a disability, please contact Jennifer Moller, preferably at least 10 days prior to the meeting, to give EPA as much time as possible to process your request.

Dated: November 8, 2006.

Cynthia C. Dougherty,

Director, Office of Ground Water and Drinking Water.

[FR Doc. E6–19266 Filed 11–14–06; 8:45 am]

BILLING CODE 6560–50–P

ENVIRONMENTAL PROTECTION AGENCY

[EPA–HQ–OPP–2005–0163; FRL–8099–7]

Aldicarb Revised Risk Assessments; Notice of Availability and Solicitation of Risk Reduction Options

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice.

SUMMARY: This notice announces the availability of EPA's revised risk assessments for the N-methyl carbamate pesticide aldicarb. In addition, this notice solicits public comment on risk reduction options for aldicarb, as well as an initial impacts and/or preliminary benefits assessmentfor a number of

aldicarb uses. The public is encouraged to suggest risk management ideas or proposals to address the risks identified. EPA is developing an Interim Reregistration Eligibility Decision (IRED) for aldicarb through the full, 6-Phase public participation process that the Agency uses to involve the public in developing pesticide reregistration and tolerance reassessment decisions. Subsequently, EPA will complete the cumulative assessment for N-methyl carbamate pesticides, including aldicarb. Additional risk mitigation for dietary concerns may be necessary for aldicarb at that time. Through these programs, EPA is ensuring that all pesticides meet current health and safety standards.

DATES: Comments must be received on or before January 16, 2007.

ADDRESSES: Submit your comments, identified by docket identification (ID) number EPA–HQ–OPP–2005–0163, by one of the following methods:

• *Federal e-Rulemaking Portal*: *http://www.regulations.gov*. Follow the on-line instructions for submitting comments.

• *Mail*: Office of Pesticide Programs (OPP) Regulatory Public Docket (7502P), Environmental Protection Agency, 1200 Pennsylvania Ave., NW., Washington, DC 20460-0001.

• *Delivery*: OPP Regulatory Public Docket (7502P), Environmental Protection Agency, Rm. S-4400, One Potomac Yard (South Building), 2777 S. Crystal Drive, Arlington, VA. Deliveries are only accepted during the Docket's normal hours of operation (8:30 a.m. to 4 p.m., Monday through Friday, excluding legal holidays). Special arrangements should be made for deliveries of boxed information. The Docket telephone number is (703) 305-5805.

Instructions: Direct your comments to docket ID number EPA–HQ–OPP–2005–0163. EPA's policy is that all comments received will be included in the docket without change and may be made available on-line at *http://www.regulations.gov*, including any personal information provided, unless the comment includes 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 regulations.gov or e-mail. The Federal regulations.gov website is an ''anonymous access'' system, which means EPA will not know your identity or contact information unless you provide it in the body of your comment. If you send an e-mail comment directly to EPA without going through regulations.gov, your e-mail address will be automatically captured and included as part of the comment that is placed in the docket and made available on the Internet. If you submit an electronic comment, EPA recommends that you include your name and other contact information in the body of your comment and with any disk or CD-ROM you submit. If EPA cannot read your comment due to technical difficulties and cannot contact you for clarification, EPA may not be able to consider your comment. Electronic files should avoid the use of special characters, any form of encryption, and be free of any defects or viruses.

Docket: All documents in the docket are listed in the docket 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, is not placed on the Internet and will be publicly available only in hard copy form. Publicly available docket materials are available either in the electronic docket at *http://www.regulations.gov*, or, if only available in hard copy, at the OPP Regulatory Public Docket in Rm. S-4400, One Potomac Yard (South Building), 2777 S. Crystal Drive, Arlington, VA. The hours of operation of this Docket Facility are from 8:30 a.m. to 4 p.m., Monday through Friday, excluding legal holidays. The Docket telephone number is (703) 305-5805.

FOR FURTHER INFORMATION CONTACT: Tracy Perry, Special Review and Reregistration Division (7508P), Office of Pesticide Programs, Environmental Protection Agency, 1200 Pennsylvania Ave., NW., Washington, DC 20460-0001; telephone number: (703) 308-0128; fax number: (703) 308-8005; e-mail address: *perry.tracy@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, 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 person listed under **FOR FURTHER INFORMATION CONTACT**.

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 e-mail. 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 submitting comments, remember to:

i. Identify the document by docket ID number and other identifying information (subject heading, **Federal Register** date and page number).

ii. Follow directions. The Agency may ask you to respond to specific questions or organize comments by referencing a Code of Federal Regulations (CFR) part or section number.

iii. Explain why you agree or disagree; suggest alternatives and substitute language for your requested changes.

iv. Describe any assumptions and provide any technical information and/or data that you used.

v. If you estimate potential costs or burdens, explain how you arrived at your estimate in sufficient detail to allow for it to be reproduced.

vi. Provide specific examples to illustrate your concerns and suggest alternatives.

vii. Explain your views as clearly as possible, avoiding the use of profanity or personal threats.

viii. Make sure to submit your comments by the comment period deadline identified.

II. Background

A. What Action is the Agency Taking?

EPA is making available the Agency's revised risk assessments, initially issued for comment through a **Federal Register** notice published on May 17, 2006 (71 FR 28693) (FRL-8064-8); a response to comments; and related documents for aldicarb. EPA is encouraging the public to submit information and suggestions for the risk management of aldicarb to

aid the Agency in its reregistration decision for this chemical (please refer to the Note to Reader in the aldicarb public docket). EPA is also soliciting public comment on preliminary benefits assessments for specific crops. EPA developed the risk assessments for aldicarb as part of its public process for making pesticide reregistration eligibility and tolerance reassessment decisions. Through these programs, EPA is ensuring that pesticides meet current standards under the Federal Insecticide, Fungicide, and Rodenticide Act (FIFRA) and the Federal Food, Drug, and Cosmetic Act (FFDCA), as amended by the Food Quality Protection Act of 1996 (FQPA).

Aldicarb is a restricted use systemic insecticide, acaricide and nematicide used to control leaf phylloxera; bud moth; citrus nematode (suppression); aphids; mites; white flies; thrips; fleahoppers, leafminers; leafhoppers; overwintering boll weevil; lygus; nematodes (suppression); cotton leaf perforator; seedcorn maggot; Mexican bean beetle; flea beetles; Colorado potato beetle; greenbug; chinch bug; three cornered alfalfa hopper (suppression); and sugar beet root maggot. Aldicarb is registered for use on agricultural crops including citrus, cotton, dry beans, peanuts, pecans, potatoes, sorghum, soybeans, sugar beets, sugarcane, sweet potatoes, and seed alfalfa (CA). In addition, aldicarb may be applied to field grown ornamentals (CA) and tobacco, and on coffee grown in Puerto Rico. There are no aldicarb products intended for sale to homeowners or for use in residential settings.

EPA is providing an opportunity, through this notice, for interested parties to provide risk management proposals or otherwise comment on risk management for aldicarb. Risks of concern associated with the use of aldicarb are: risks from rural drinking water wells in peanut/cotton growing regions in the southern coastal plain (Alabama, Georgia, South Carolina), ecological risks to mammals, birds, fish and aquatic invertebrates. In targeting these risks of concern, the Agency solicits information on effective and practical risk reduction measures.

EPA is applying the principles of public participation to all pesticides undergoing reregistration and tolerance reassessment. The Agency's Pesticide Tolerance Reassessment and Reregistration; Public Participation Process, published in the **Federal Register** on May 14, 2004 (69 FR 26819)(FRL–7357–9), explains that in conducting these programs, EPA is tailoring its public participation process to be commensurate with the level of risk, extent of use, complexity of issues, and degree of public concern associated with each pesticide. Due to its uses, risks, and other factors, aldicarb is being reviewed through the full 6-Phase public participation process.

All comments should be submitted using the methods in **ADDRESSES**, and must be received by EPA on or before the closing date. Comments and proposals will become part of the Agency Docket for aldicarb. Comments received after the close of the comment period will be marked ''late'' EPA is not required to consider these late comments.

After considering comments received, EPA will develop and issue for comment the aldicarb IRED. Further risk mitigation measures may be needed when EPA considers its cumulative assessment of the N-methyl carbamate pesticides.

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

Section 4(g)(2) of FIFRA as amended directs that, after submission of all data concerning a pesticide active ingredient, ''the Administrator shall determine whether pesticides containing such active ingredient are eligible for reregistration,'' before calling in product specific data on individual end-use products and either reregistering products or taking other ''appropriate regulatory action.''

Section 408(q) of the FFDCA, 21 U.S.C. 346a(q), requires EPA to review tolerances and exemptions for pesticide residues in effect as of August 2, 1996, to determine whether the tolerance or exemption meets the requirements of section 408(b)(2) or (c)(2) of FFDCA.

List of Subjects

Environmental protection, Pesticides and pests.

Dated: November 8, 2006.

Debra Edwards,

Director, Special Review and Reregistration Division, Office of Pesticide Programs.

[FR Doc. E6–19360 Filed 11–14–06; 8:45 am]

BILLING CODE 6560–50–S

ENVIRONMENTAL PROTECTION AGENCY

[EPA–HQ–OPP–2006–0875; FRL–8102–5]

Notice of Filing of Pesticide Petitions for Establishment or Amendment to Regulations for Residues of Pesticide Chemicals in or on Various Commodities

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice.

SUMMARY: This notice announces the initial filing of pesticide petitions proposing the establishment or amendment of regulations for residues of pesticide chemicals in or on various commodities.

DATES: Comments must be received on or before December 15, 2006

ADDRESSES: Submit your comments, identified by docket identification (ID) number EPA–HQ–OPP–2006–0875 and pesticide petition numbers (PP) 4E6867 and 6E7066, by one of the following methods:

- Federal eRulemaking Portal: *http://www.regulations.gov*. Follow the on-line instructions for submitting comments.
- *Mail*: Office of Pesticide Programs (OPP) Regulatory Public Docket (7502P), Environmental Protection Agency, 1200 Pennsylvania Ave., NW., Washington, DC 20460–0001.
- *Delivery*: OPP Regulatory Public Docket (7502P), Environmental Protection Agency, Rm. S–4400, One Potomac Yard (South Bldg.), 2777 S. Crystal Drive, Arlington, VA. Deliveries are only accepted during the Docket's normal hours of operation 8:30 a.m. to 4 p.m., Monday through Friday, excluding legal holidays. Special arrangements should be made for deliveries of boxed information. The Docket telephone number is (703) 305–5805.

Instructions: Direct your comments to docket ID number EPA–HQ–OPP–2006–0875. EPA's policy is that all comments received will be included in the docket without change and may be made available on-line at *http://www.regulations.gov*, including any personal information provided, unless the comment includes 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 regulations.gov or e-mail. The Federal regulations.gov website is an ''anonymous access'' system, which means EPA will not