(FERC. No. 199), Relicensing for Existing 130-megawatt (MW), Santee and Cooper Rivers, Berkeley, Calhoun, Clarendon, Orangeburg and Sumter Counties, SC, Wait Period Ends: 12/03/2007, Contact: Monte Terhaar 202–502–6035.

EIS No. 20070460, Final EIS, FRC, WI, Guardian Expansion and Extension Project, Construction and Operation Natural Gas Pipeline Facilities, Jefferson, Dodge, Fond du Lac, Calument, Brown, Walworth, Outagamie Counties, WI, Wait Period Ends: 12/03/2007, Contact: Andy Black 1–866–208–3372.

EIS No. 20070461, Final EIS, IBR, 00, Colorado River Interim Guidelines for Lower Basin Shortages and Coordinated Operations for Lake Powell and Lake Mead, Implementation, Colorado River, CO and CA, Wait Period Ends: 12/03/2007, Contact: Nan Yoder 702–595–9851.

EIS No. 20070462, Final EIS, MMS, 00, Eastern Planning Area Outer Continental Shelf (OCS) Oil and Gas Lease Sale 224, Gulf of Mexico Offshore Marine Environment and Coastal Parshes/Counties of LA, MS, AL, and North Western Florida, Wait Period Ends: 12/03/2007, Contact: Dr. Sally Valdes 703–787–1707.

EIS No. 20070463, Draft EIS, CGD/MARAD, FL, Calypso Liquefied Natural Gas (LNG) Deepwater Port License Application, Proposes to Own, Construct, and Operate a Deepwater Port, Outer Continental Shelf (OCS) in the OCS NG 17–06 (Bahamas) Lease Area, 8 to 10 miles off the East Coast of Florida to the Northeast of Port Everglades, FL, Comment Period Ends: 12/17/2007, Contact: Mary Kate Jager 202–372–1454.

Amended Notices

EIS No. 20070385, Draft EIS, FHW, 00, Peace Bridge Expansion Project, Capacity Improvements to the Peace Bridge, Plazas and Connecting Roadways, U.S. Coast Guard Bridge Permit, U.S. Army COE Section 10 and 404 Permits. City of Buffalo, Erie County, NY and Town of Fort Erie, Ontario, Canada, Comment Period Ends: 11/13/2007, Contact: Amy Jackson-Grove 518–431–4131.

Revision of Fr Notice Published 09/14/2007: Extending Comment Period from 10/29/2007 to 11/13/2007.

Dated: October 30, 2007.

Ken Mittelholtz,

Environmental Protection Specialist, Office of Federal Activities.

[FR Doc. E7–21590 Filed 11–1–07; 8:45 am]

BILLING CODE 6560–50–P

ENVIRONMENTAL PROTECTION AGENCY

[EPA–HQ–OPP–2007–0588; FRL–8154–4]

Acrolein Human Health Risk Assessment; Notice of Availability

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice.

SUMMARY: This notice announces the availability of EPA's human health risk assessment, and related documents for the pesticide acrolein, and opens a public comment period on these documents (Phase 3 of 6–Phase Process). The public is encouraged to suggest risk management ideas or proposals to address the risks identified. EPA is developing a Reregistration Eligibility Decision (RED) for acrolein through a 6–Phase public participation process that the Agency uses to involve the public in developing pesticide reregistration decisions. Through this process, EPA is ensuring that all pesticides meet current health and safety standards.

DATES: Comments must be received on or before January 2, 2008.

ADDRESSES: Submit your comments, identified by docket identification (ID) number EPA–HQ–OPP–2007–0588, 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 Dr., 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 Facility telephone number is (703) 305–5805.

Instructions: Direct your comments to docket ID number EPA–HQ–OPP–2007–0588. 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 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 available in regulations.gov. To access the electronic docket, go to *http://www.regulations.gov*, select ''Advanced Search,'' then ''Docket Search.'' Insert the docket ID number where indicated and select the ''Submit'' button. Follow the instructions on the regulations.gov website to view the docket index or access available documents. 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 Bldg.), 2777 S. Crystal Dr., 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 Facility telephone number is (703) 305–5805.

FOR FURTHER INFORMATION CONTACT: Amaris Johnson, Special Review and

Reregistration Division (7508P), Office of Pesticide Programs, Environmental Protection Agency, 1200 Pennsylvania Ave., NW., Washington, DC 20460–0001; telephone number: (703) 305–9542; fax number: (703) 308–7070; e-mail address: *johnson.amaris@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 releasing for public comment its human health risk assessment and related documents for acrolein, an aldehyde, and soliciting public comment on risk management ideas or proposals. Acrolein is primarily used in irrigation canals and reservoirs to treat aquatic weeds, such as pondweed. Its secondary use is in oil fields, as a biocide to remove bacteria during petroleum production. EPA developed the human health risk assessment and risk characterization for acrolein as a part of its public process for making its pesticide reregistration eligibility decision. Through this program, EPA is ensuring that pesticides meet current standards under the Federal Insecticide, Fungicide, and Rodenticide Act (FIFRA). The ecological risk assessment for acrolein was posted in the docket for the Phase 3 public comment period on July 25, 2007. This comment period closed on September 23, 2007. The Agency intends to review the comments for both the ecological and human health risk assessments, revise the risk assessments if appropriate from comments received, and provide an additional public comment period on both the ecological and human health risk assessments for acrolein.

Acrolein is registered as a non-food use aquatic herbicide used primarily in irrigation canals. It is a non-specific biocide that treats aquatic weeds by breaking down their cell walls. Acrolein is formulated as a liquid, and is metered directly into irrigation canals or reservoirs.

EPA is providing an opportunity, through this notice, for interested parties to provide comments and input on the Agency's human health risk assessment for acrolein. Such comments and input could address, for example, the availability of additional data to further refine the risk assessments, such as residue data on root and tuber crops, air monitoring data, and residential bystander exposure and risk information, or could address the Agency's risk assessment methodologies and assumptions as applied to this specific pesticide.

Through this notice, EPA also is providing an opportunity for interested parties to provide risk management proposals or otherwise comment on risk management for acrolein. Risks of concern associated with the use of acrolein include:

1. Exposure to workers during application periods, and

2. Bystanders during post-application periods.

In targeting these risks of concern, the Agency solicits information on effective and practical risk reduction measures.

EPA seeks to achieve environmental justice, the fair treatment and meaningful involvement of all people, regardless of race, color, national origin, or income, 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, unusually high exposure to acrolein, compared to the general population.

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, the Agency is tailoring its public participation process to be commensurate with the level of risk, extent of use, complexity of the issues, and degree of public concern associated with each pesticide. For acrolein the full, 6–Phase process with 2 comment periods and ample opportunity for public consultation seems appropriate in view of its large number of users in the Pacific Northwest, multiple incidents, complex issues, acute toxicity, and numerous affected stakeholders.

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

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.''

List of Subjects

Environmental protection, Pesticides and pests.

Dated: October 23, 2007.

Steve Bradbury,

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

[FR Doc. E7–21438 Filed 11–1–07; 8:45 am]

BILLING CODE 6560–50–S

ENVIRONMENTAL PROTECTION AGENCY

[EPA–HQ–OPP–2007–1060; FRL–8155–4]

Pesticide Inert Ingredients: Support Status of Revoked Tolerance Exemptions

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice.

SUMMARY: EPA has received notice from various companies of their intention to submit data in order to support the reinstatement of a number of inert ingredient tolerance exemptions that were revoked because of insufficient data, revocations effective as of August 9, 2008, in a final rule published in the **Federal Register** on August 9, 2006 (71 FR 45415). This notice identifies the tolerance exemptions that the companies indicate they will be supporting with the submission of data. The information in this notice is for informational purposes only and does not affect the previous revocations. Based on the review of the submitted data, EPA will conduct rulemaking to establish new tolerance exemptions where appropriate.

FOR FURTHER INFORMATION CONTACT: Kerry Leifer, Registration Division (7505P), Office of Pesticide Programs, Environmental Protection Agency, 1200 Pennsylvania Ave., NW., Washington, DC 20460–0001; telephone number: (703) 308–8811; fax number: (703) 605–0781; e-mail address: *leifer.kerry@epa.gov*.

SUPPLEMENTARY INFORMATION:

I. General Information

A. Does this Action Apply to Me?

You may be potentially affected by this action if you are an agricultural producer, food manufacturer, or pesticide manufacturer. Potentially affected entities may include, but are not limited to:

- Crop production (NAICS code 111).
- Animal production (NAICS code 112).
- Food manufacturing (NAICS code 311).
- Pesticide manufacturing (NAICS code 32532).

This listing is not intended to be exhaustive, but rather provides a guide for readers regarding entities likely to be affected by this action. Other types of entities not listed in this unit could also be affected. The North American Industrial Classification System (NAICS) codes have been provided to assist you and others in determining whether this action might apply to certain entities. To determine whether you or your business may be affected by this action, you should carefully examine the applicability provisions in Unit II. 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. How Can I Get Copies of this Document and Other Related Information?

1. *Docket.* EPA has established a docket for this action under docket identification (ID) number EPA–HQ–OPP–2007–1060. 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 Office of Pesticide Programs (OPP) Regulatory Public Docket in Rm. S–4400, One Potomac Yard (South Bldg.), 2777 S. Crystal Dr., 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 Facility telephone number is (703) 305–5805.

2. *Electronic access.* You may access this **Federal Register** document electronically through the EPA Internet under the ''**Federal Register**'' listings at *http://www.epa.gov/fedrgstr*.

II. Background

A. Background on the Revocation of Pesticide Inert Ingredient Tolerance Exemptions on August 9, 2006

In a final rule published in the **Federal Register** on August 9, 2006 (71 FR 45415) (FRL–8084–1), EPA revoked inert ingredient tolerance exemptions because insufficient data were available to the Agency to make the safety determination required by FFDCA section 408(c)(2). In making the FFDCA reassessment safety determination, EPA considered the validity, completeness, and reliability of the data that are available to the Agency [FFDCA section 408 (b)(2)(D)] and the available information concerning the special susceptibility of infants and children (including developmental effects from *in utero* exposure) [FFDCA section 408 (b)(2)(C)]. Data gaps existed for these inert ingredients in areas that were critical to reassessment. Without these data, the assessment of possible effects to infants and children could not be made. EPA concluded it had insufficient data to make the safety finding of FFDCA section 408(c)(2) and revoked the inert ingredient tolerance exemptions identified in the final rule under 40 CFR 180.910, 180.920, 180.930, and 180.940, with the revocations effective two years after the date of publication. The tolerance exemptions will expire on August 9, 2008.

B. What Information Is Provided In This Notice?

EPA has received communications from pesticide registrants and inert ingredient manufacturers expressing interest in supporting certain inert ingredient tolerance exemptions that were revoked in the final rule of August 9, 2008. EPA developed voluntary guidance describing how interested parties could support these revoked tolerance exemptions, including consultations with the Agency and how they can demonstrate support, including identifying test materials and providing evidence that a laboratory has been hired to conduct the study. The voluntary guidance document, entitled ''Guidance for Supporting the Inert Ingredients Subject to the Revocation Notice of August 9, 2006'', is available on EPA's website at *http://www.epa.gov/opprd001/inerts/*.

In the interest of keeping the stakeholders informed about activities that may impact these revoked tolerance exemptions, EPA is publishing in this notice the support status of each of the revoked tolerance exemptions by indicating whether the Agency has received a demonstration of support (such as described in the guidance document discussed above).

Be advised that the information provided in this notice does not guarantee or in any way bind the Agency to reinstate tolerance