

Requirements,” which referenced under the “Guideline Number” column the 91 series of test guidelines. EPA’s intention is to replace the 91 series test guideline designations with the appropriate 810 series test guideline designations. Therefore, at the time of the publication of the final rule, appropriate references to the 810 series test guideline numbers and names will be incorporated into the final rule.

#### IV. Are there any applicable voluntary consensus standards that EPA should consider?

This notice of availability does not involve a proposed regulatory action that would require the Agency to consider voluntary consensus standards pursuant to section 12(d) of the National Technology Transfer and Advancement Act of 1995 (NTTAA), Public Law 104–113, section 12(d) (15 U.S.C. 272 note). Section 12(d) of NTTAA directs EPA to use voluntary consensus standards in its regulatory activities unless to do so would be inconsistent with applicable law or otherwise impractical. Voluntary consensus standards are technical standards (e.g., materials specifications, test methods, sampling procedures, and business practices) that are developed or adopted by voluntary consensus standards bodies. NTTAA requires EPA to provide an explanation to Congress, through OMB, when the Agency decides not to use available and applicable voluntary consensus standards when NTTAA directs the Agency to do so.

These test guidelines represent an Agency effort to harmonize the test guidelines within OCSPP, as well as to harmonize the OCSPP test guidelines with those of OECD. The process for developing and amending these test guidelines, which began in 1991, includes public participation and the extensive involvement of the scientific community, including peer review by SAP and the SAB and other expert scientific organizations.

In the future, these test guidelines could be incorporated into regulatory actions taken by EPA under TSCA, *i.e.*, with regard to the TSCA section 4 testing program. Although, NTTAA requirements do not specifically apply to the issuance of these particular test guidelines, EPA invites your comment on whether or not there are any voluntary consensus standards that should be considered during the development of the final test guidelines or any future regulatory action that may be taken under TSCA. Future regulatory actions under TSCA section 4 may involve notice and comment rulemaking or negotiated voluntary testing enforcement consent agreements/orders/

decrees. Nevertheless, However, the Agency is interested in whether or not there are any voluntary consensus standards that EPA should consider either as part of the development of the final test guidelines themselves or in lieu of these final test guidelines when the Agency develops any future regulatory action that incorporates these test guidelines. Any comments provided will assist the Agency in complying with NTTAA by facilitating the Agency’s identification of voluntary consensus standards that should be addressed in the test guideline or considered during the development of a proposed regulatory action that incorporates any standards included in the final test guidelines. Please submit your comments as directed under **ADDRESSES**.

#### List of Subjects

Environmental protection, Antimicrobial agents, Chemicals, Harmonized test guidelines, Health and safety.

Dated: September 7, 2011.

**Stephen A. Owens,**

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

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**BILLING CODE 6560–50–P**

#### ENVIRONMENTAL PROTECTION AGENCY

[FRL–9465–2; Docket ID No. EPA–HQ–ORD–2011–0671]

#### Draft Toxicological Review of n-Butanol: 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; Extension.

**SUMMARY:** EPA announced a 60-day public comment period and a listening session on August 31, 2011 (76 FR 54227) for the external review draft human health assessment titled, “Toxicological Review of n-Butanol: In Support of Summary Information on the Integrated Risk Information System (IRIS)” (EPA/635/R–11/081A). EPA is extending the public comment period one week because of a one-week delay in the release of the Toxicological Review to the public. 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. After public review and comment, an EPA contractor will convene an expert panel for independent external peer review of this draft assessment. The public comment period and external peer review meeting are separate processes that provide opportunities for all interested parties to comment on the assessment. The external peer review meeting will be scheduled at a later date and announced in the **Federal Register**. Public comments submitted during the public comment period will be provided to the external peer reviewers before the panel meeting and considered by EPA in the disposition of public comments. Public comments received after the public comment period closes will not be submitted to the external peer reviewers and will only be considered by EPA if time permits.

The listening session will be held on October 26, 2011, 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. **DATES:** The public comment period will be extended to end November 7, 2011. Comments should be in writing and must be received by EPA by November 7, 2011.

The listening session on the draft assessment for n-Butanol will be held on October 26, 2011, beginning at 9 a.m. and ending at 4 p.m., Eastern Daylight Time or when the last presentation has been completed. To attend the listening session, interested parties should register no later than October 19, 2011, following the instructions in the August 31 **Federal Register** Notice (76 FR 54227). The location and instructions for entering the building can be found in the August 31, 2011, **Federal Register** Notice (76 FR 54227).

**ADDRESSES:** The draft “Toxicological Review of n-Butanol: 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 the August 31, 2011, **Federal Register** Notice (76 FR 54227).

*Additional Information:* For information on the docket, <http://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](mailto:ORD.Docket@epa.gov).

For information on the draft assessment, please contact Ambuja Bale, National Center for Environmental Assessment [Mail Code: (8601-P)], U.S. Environmental Protection Agency, National Center for Environmental Assessment, Office of Research and Development, U.S. Environmental Protection Agency, 1200 Pennsylvania Ave., NW., Washington, DC 20460; *telephone*: 703-347-8643; *facsimile*: 703-347-8689; or *e-mail*: [FRN\\_Questions@epa.gov](mailto:FRN_Questions@epa.gov).

Dated: September 9, 2011.

**Darrell A. Winner,**

*Acting Director, National Center for Environmental Assessment.*

[FR Doc. 2011-23664 Filed 9-14-11; 8:45 am]

**BILLING CODE 6560-50-P**

## ENVIRONMENTAL PROTECTION AGENCY

[EPA-HQ-OPP-2011-0005; FRL-8887-7]

### Pesticide Products; Receipt of Applications To Register New Uses

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

**ACTION:** Notice.

**SUMMARY:** This notice announces receipt of applications to register new uses for pesticide products containing currently registered active ingredients, pursuant to the provisions of section 3(c) of the Federal Insecticide, Fungicide, and Rodenticide Act (FIFRA), as amended.

EPA is publishing this Notice of such applications, pursuant to section 3(c)(4) of FIFRA.

**DATES:** Comments must be received on or before October 17, 2011.

**ADDRESSES:** Submit your comments, identified by the docket identification (ID) number specified within the table below, 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 Facility'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 the docket ID number specified for the pesticide of interest as shown in the registration application summaries. 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 [www.regulations.gov](http://www.regulations.gov) or e-mail. The [www.regulations.gov](http://www.regulations.gov) Web site 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 [www.regulations.gov](http://www.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 at <http://www.regulations.gov>. 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:** A contact person is listed at the end of each registration application summary and may be contacted by telephone or e-mail. The mailing address for each contact person listed is: Registration Division (7505P), Office of Pesticide Programs, Environmental Protection Agency, 1200 Pennsylvania Ave., NW., Washington, DC 20460-0001 or Biopesticides and Pollution Prevention Division (7511P), Office of Pesticide Programs, Environmental Protection Agency, 1200 Pennsylvania Ave., NW., Washington, DC 20460-0001.

#### 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