Information Office (A–101), 1200 Pennsylvania Ave., NW., Washington, DC 20460–0001. The request should: Identify the product name and registration number and specify the data or information desired.

A paper copy of the fact sheet, which provides more detail on this registration, may be obtained from the National Technical Information Service (NTIS), 5285 Port Royal Rd., Springfield, VA 22161.

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/*.

An electronic version of the public docket is available through EPA's electronic public docket and comment system, EPA Dockets. You may use EPA Dockets at *http://www.epa.gov/edocket/* to submit or view public comments, access the index listing of the contents of the official public docket, and to access those documents in the public docket that are available electronically. Although not all docket materials may be available electronically, you may still access any of the publicly available docket materials through the docket facility identified in Unit I.B.1. Once in the system, select ''search,'' then key in the appropriate docket ID number.

II. Did EPA Approve the Application?

The Agency approved the application after considering all required data on risks associated with the proposed use of Yeast Extract Hydrolysate from *Saccharomyces cerevisiae*, and information on social, economic, and environmental benefits to be derived from use. Specifically, the Agency has considered the nature of the chemical and its pattern of use, application methods and rates, and level and extent of potential exposure. Based on these reviews, the Agency was able to make basic health and safety determinations which show that use of Yeast Extract Hydrolysate from *Saccharomyces cerevisiae* when used in accordance with widespread and commonly recognized practice, will not generally cause unreasonable adverse effects to the environment.

III. Approved Application

EPA issued a notice, published in the **Federal Register** of August 6, 2003 (68 FR 46607) (FRL–7316–7), which announced that the Interregional Research Project No. 4 (IR-4), Rutgers University, Technology Center of New Jersey, 681 U.S. Highway #1 South, North Brunswick, NJ 08902–3390 on behalf of Morse Enterprises Limited, Inc., Brickell East, Floor Ten, 151 South East 15 Road, Miami, FL 33129, had submitted applications to register a manufacturing use product, Yeast Hydrolysate Liquid (EPA File Symbol 73512–E) and the end use product, KeyPlex 350 (EPA File Symbol 73512–R) containing the new active ingredient Yeast Extract Hydrolysate from *Saccharomyces cerevisiae*, which was not included in any previously registered product.

The following products were approved on February 19, 2004.

1. The manufacturing use product, Yeast Hydrolysate Liquid, containing 2.5% active ingredient (EPA Registration Number 73512–2) for use in the management of plant diseases.

2. The end use product, KeyPlex 350, containing 0.063% active ingredient (EPA Registration Number 73512–1 for use in the management of plant diseases.

List of Subjects

Environmental protection, Chemicals, Pesticides and pests.

Dated: June 29, 2004.

Janet L. Andersen,

Director, Biopesticides and Pollution Prevention Division, Office of Pesticide Programs.

[FR Doc. 04–15726 Filed 7–13–04; 8:45 am]

BILLING CODE 6560–50–S

ENVIRONMENTAL PROTECTION AGENCY

[OPP–2004–0192; FRL–7366–5]

Issuance of an Experimental Use Permit

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice.

SUMMARY: EPA has granted an experimental use permit (EUP) to the following pesticide applicant. An EUP permits use of a pesticide for experimental or research purposes only in accordance with the limitations in the permit.

FOR FURTHER INFORMATION CONTACT: Diana M. Horne, Biopesticides and Pollution Prevention Division (7511C), Office of Pesticide Programs, Environmental Protection Agency, 1200 Pennsylvania Ave., NW., Washington, DC 20460–0001; telephone number: (703) 308–8367; e-mail address: *horne.diana@epa.gov*.

SUPPLEMENTARY INFORMATION:

I. General Information

A. Does this Action Apply to Me?

This action is directed to the public in general. Although this action may be of particular interest to those persons who conduct or sponsor research on pesticides, 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 information in this action, 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 an official public docket for this action under docket identification (ID) number OPP–2004–0192. The official public docket consists of the documents specifically referenced in this action, any public comments received, and other information related to this action. Although a part of the official docket, the public docket does not include Confidential Business Information (CBI) or other information whose disclosure is restricted by statute. The official public docket is the collection of materials that is available for public viewing at the Public Information and Records Integrity Branch (PIRIB), Rm. 119, Crystal Mall #2, 1801 South Bell Street, Arlington, VA. This docket facility is open from 8:30 a.m. to 4 p.m., Monday through Friday, excluding legal holidays. The docket 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/*.

An electronic version of the public docket is available through EPA's electronic public docket and comment system, EPA Dockets. You may use EPA Dockets at *http://www.epa.gov/edocket/* to submit or view public comments, access the index listing of the contents of the official public docket, and to access those documents in the public docket that are available electronically. Although not all docket materials may be available electronically, you may still access any of the publicly available docket materials through the docket facility identified in Unit I.B.1. Once in the system, select ''search,'' then key in the appropriate docket ID number.

II. EUP

EPA has issued the following EUP:

69834–EUP–2. Issuance. EDEN Bioscience Corporation, 3830 Monte Villa Parkway, Suite 100, Bothell,

Washington 98021–7266. This EUP allows the use of 70.85 pounds of the biochemical pesticide Harpin αβ protein on 4,942 acres of citrus, cotton, field corn, ornamentals, peanut, rice, soybean, sugarcane, and wheat to evaluate the control of post harvest diseases, enhancing overall plant health, thus improving stand establishment and enhancing crop yield and quality. The program is authorized only in the States of Alabama, Arizona, Arkansas, Colorado, Florida, Georgia, Illinois, Indiana, Iowa, Kansas, Kentucky, Louisiana, Maryland, Minnesota, Mississippi, Missouri, Nebraska, New Jersey, New York, North Carolina, North Dakota, Ohio, Oklahoma, Pennsylvania, South Carolina, South Dakota, Tennessee, Texas, Virginia, and Washington. The EUP is effective from April 26, 2004 to April 26, 2006. A tolerance has been established for residues of the active ingredient in or on all food commodities.

Authority: 7 U.S.C. 136c.

List of Subjects

Environmental protection, Experimental use permits.

Dated: June 29, 2004.

Janet L. Andersen,

Director, Biopesticides and Pollution Prevention Division, Office of Pesticide Programs.

[FR Doc. 04–15725 Filed 7–13–04; 8:45 am]

BILLING CODE 6560–50–S

ENVIRONMENTAL PROTECTION AGENCY

[FRL–7787–1]

Notice of Review of Water Quality Indicators and/or Rapid Measurement Technology

AGENCY: U.S. Environmental Protection Agency, National Health and Environmental Effects Research Laboratory.

ACTION: Notice.

SUMMARY: This review will be conducted, and the information collected, by the Epidemiology and Biomarkers Branch, Human Studies Division, National Health and Environmental Effects Research Laboratory, Office of Research and Development, U.S. Environmental Protection Agency (EPA). Product information packages collected for review are strictly voluntary. This notice is a result of Section 3(a)(v)(1) of the Beaches Environmental Assessment and Coastal Health Act of 2000 and the strategic plan for EPA's Office of Research and Development (ORD) and the Office of Water entitled ''Action Plan for Beaches and Recreational Water.''

Current EPA guidelines (Section 303(c) Clean Water Act) recommend the use of cultural methods for E. coli and enterococci to measure recreational water quality. These methods produce results in 24 hours creating a delayed response for recreational water managers. This shortcoming in the current practices for measuring recreational water quality has led EPA to consider new technology and indicators that will provide rapid measurement of water quality (preferably 2 hours or less). This will give vendors the opportunity to showcase technology and determine needs for current industry.

Packages submitted should include the following information (if applicable):

1. Cost of the equipment/instrument?
2. Cost of supplies *per test* (List items needed with individual cost)?
3. Commercially available? How long has it been on the market? Are supplies commercially available?
4. Shipping time?
5. Is it portable? Laboratory or field use?
6. Skill level for operators? Do they need special training? Is it included in purchase?
7. Computer needed? Is it included? Specifications?
8. Specificity for target organisms? False-positive and false-negative rates?
9. Limit of detection?
10. Types of samples tested (primarily, but not limited to, types of water samples)?
11. Volume of sample required?
12. Detect live organisms?
13. Developed/validated method or experimental method?
14. Analysis time for one sample?
15. Preparation time for one sample?
16. How many samples can be analyzed per day?
17. Quality control procedures?
18. Is the system sterile? Cleaning and sanitization procedures?
19. References where equipment/instrument was used?
20. Data form (tables, graphs, etc.) and examples?
21. Can data be stored in the equipment/instrument, if portable?
22. Include brochures and pictures/photos/schematics.
23. Warranty and repair services?
24. Any distinctive elements that uniquely qualify this instrument or methods above others.

DATES: Review of information packages will begin on July 30, 2004. All interested parties should submit packages on or before March 31, 2005.

FOR FURTHER INFORMATION CONTACT: Elizabeth Sams, (919), 843–3161, FAX: (919) 966–0655, E-mail: *sams.elizabeth@epa.gov*, or by mailing a request Elizabeth Sams, U.S. EPA (MD 58–C), Research Triangle Park, NC 27711.

Dated: June 1, 2004.

Harold Zenick,

Associate Director for Health, Office of Research and Development.

[FR Doc. 04–15947 Filed 7–13–04; 8:45 am]

BILLING CODE 6560–50–P

FEDERAL COMMUNICATIONS COMMISSION

Notice of Public Information Collection(s) Being Submitted to OMB for Review and Approval

June 29, 2004.

SUMMARY: The Federal Communications Commissions, as part of its continuing effort to reduce paperwork burden invites the general public and other Federal agencies to take this opportunity to comment on the following information collection, as required by the Paperwork Reduction Act of 1995, Public Law 104–13. An agency may not conduct or sponsor a collection of information unless it displays a currently valid control number. No person shall be subject to any penalty for failing to comply with a collection of information subject to the Paperwork Reduction Act (PRA) that does not display a valid control number. Comments are requested concerning (a) whether the proposed collection of information is necessary for the proper performance of the functions of the Commission, including whether the information shall have practical utility; (b) the accuracy of the Commission's burden estimate; (c) ways to enhance the quality, utility, and clarity of the information collected; and (d) ways to minimize the burden of the collection of information on the respondents, including the use of automated collection techniques or other forms of information technology.

DATES: Written comments should be submitted on or before August 13, 2004. If you anticipate that you will be submitting comments, but find it difficult to do so within the period of time allowed by this notice, you should advise the contact listed below as soon as possible.

ADDRESSES: Direct all comments to Les Smith, Federal Communications Commission, Room 1–A804, 445 12th