You may also provide the name, date, and **Federal Register** citation.

II. Registration Applications

EPA received applications as follows to register pesticide products containing active ingredients not included in any previously registered products pursuant to the provision of section 3(c)(4) of FIFRA. Notice of receipt of these applications does not imply a decision by the Agency on the applications.

Products Containing Active Ingredients not Included in any Previously Registered Products

1. *File Symbol*: 80224–R. *Applicant*: Innolytics, LLC, P.O. Box 675935, Rancho Santa Fe, CA 92067. *Product name*: LLC/Ovocontrol-P. *Active ingredient*: Nicarbazin. *Proposed classification/Use*: None. Control hatchability of feral pigeon eggs.

2. *File Symbol*: 80224–E. *Applicant*: Innolytics, LLC, P.O. Box 675935, Rancho Santa Fe, CA 92067. *Product name*: Nicarbazin 30% Granulated Premix. *Active ingredient*: Nicarbazin. *Proposed classification/Use*: None. Manufacturing-use product for formulation into end-use products to control the hatchability of resident Canada geese and feral pigeon eggs.

3. *File Symbol*: 80224–G. *Applicant*: Innolytics, LLC. *Product name*: LLC/Ovocontrol-G. *Active ingredient*: Nicarbazin. *Proposed classification/Use*: None. Control hatchability of resident Canada geese eggs.

List of Subjects

Environmental protection, Pesticides and pest.

Dated: March 21, 2005.

Betty Shackleford,

Acting Director, Registration Division, Office of Pesticide Programs.

[FR Doc. 05–6629 Filed 4–5–05; 8:45am]

BILLING CODE 6560–50–S

ENVIRONMENTAL PROTECTION AGENCY

[OPPT–2005–0017; FRL–7707–4]

Approval of Test Marketing Exemption for a Certain New Chemical

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice.

SUMMARY: This notice announces EPA's approval of an application for test marketing exemption (TME) under section 5(h)(1) of the Toxic Substances Control Act (TSCA) and 40 CFR 720.38. EPA has designated this application as TME–05–0001. The test marketing conditions are described in the TME application and in this notice.

DATES: Approval of this TME is effective March 24, 2005.

FOR FURTHER INFORMATION CONTACT: *For general information contact*: Colby Lintner, Regulatory Coordinator, Environmental Assistance Division (7408M), Office of Pollution Prevention and Toxics, Environmental Protection Agency, 1200 Pennsylvania Ave., NW., Washington, DC 20460–0001; telephone number: (202) 554–1404; e-mail address: *TSCA-Hotline@epa.gov*.

For technical information contact: Virginia Lee, Chemical Control Division (7405M), Office of Pollution Prevention and Toxics, Environmental Protection Agency, 1200 Pennsylvania Ave., NW., Washington, DC 20460–0001; telephone number: (202) 564–0883; e-mail address: *lee.virginia@epa.gov*.

SUPPLEMENTARY INFORMATION:

I. General Information

A. Does this Action Apply to Me?

This action is directed in particular to the chemical manufacturer and/or importer who submitted the TME to EPA. This action may, however, be of interest to the public in general. Since other entities may also 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 technical 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 OPPT–2005–0017. 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 EPA Docket Center, Rm. B102–Reading Room, EPA West, 1301 Constitution Ave., NW., Washington, DC. The EPA Docket Center is open from 8:30 a.m. to 4:30 p.m., Monday through Friday, excluding legal holidays. EPA Docket Center Reading Room telephone number is (202) 566–1744 and the telephone number for the OPPT Docket, which is located in EPA Docket Center, is (202) 566–0280.

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

Section 5(h)(1) of TSCA and 40 CFR 720.38 authorizes EPA to exempt persons from premanufacture notification (PMN) requirements and permit them to manufacture or import new chemical substances for test marketing purposes, if the Agency finds that the manufacture, processing, distribution in commerce, use, and disposal of the substances for test marketing purposes will not present an unreasonable risk of injury to health or the environment. EPA may impose restrictions on test marketing activities and may modify or revoke a test marketing exemption upon receipt of new information which casts significant doubt on its finding that the test marketing activity will not present an unreasonable risk of injury.

III. What Action is the Agency Taking?

EPA approves the above-referenced TME. EPA has determined that test marketing the new chemical substance, under the conditions set out in the TME application and in this notice, will not present any unreasonable risk of injury to health or the environment.

IV. What Restrictions Apply to this TME?

The test market time period, production volume, number of customers, and use must not exceed specifications in the application and this notice. All other conditions and restrictions described in the application and in this notice must also be met.

TME–05–0001

Date of receipt: February 7, 2005.

Notice of receipt: March 14, 2005, (70 FR 12478) (FRL–7704–9).

Applicant: PPG Industries, Inc..

Chemical: Alkanediocic acid, polymer with 1,3,5-tris(substituted alkyl)-1,3,5-triazine-2,4,6(*1H,3H,5H*)-trione, alkanotate (ester) 3-substituted-2-(substituted alkyl)-2-alkanoate (ester).

Use: Component of an automotive refinish direct-gloss topcoat.

Production volume: 5,000 kilogram/year (kg/yr).

Number of customers: 50.

Test marketing period: 365 days, commencing on first day of commercial manufacture.

The following additional restrictions apply to this TME. A bill of lading accompanying each shipment must state that the use of the substance is restricted to that approved in the TME. In addition, the applicant shall maintain the following records until 5 years after the date they are created, and shall make them available for inspection or copying in accordance with section 11 of TSCA:

1. Records of the quantity of the TME substance produced and the date of manufacture.

2. Records of dates of the shipments to each customer and the quantities supplied in each shipment.

3. Copies of the bill of lading that accompanies each shipment of the TME substance.

V. What was EPA's risk assessment for this TME?

EPA identified no significant health or environmental concerns for the test market substance. Therefore, the test market activities will not present any unreasonable risk of injury to human health or the environment.

VI. Can EPA Change Its Decision on this TME in the Future?

Yes. The Agency reserves the right to rescind approval or modify the conditions and restrictions of an exemption should any new information that comes to its attention cast significant doubt on its finding, that the test marketing activities will not present any unreasonable risk of injury to human health or the environment.

List of Subjects

Environmental protection, Test marketing exemptions.

Dated: March 24, 2005.

Miriam Wiggins-Lewis,

Acting Chief, New Chemicals Prenotice Management Branch, Office of Pollution Prevention and Toxics.

[FR Doc. 05–6628 Filed 4–5–05 8:45 am]

BILLING CODE 6560–50–S

FEDERAL COMMUNICATIONS COMMISSION

Notice of Public Information Collection(s) Being Reviewed by the Federal Communications Commission for Extension Under Delegated Authority

March 25, 2005.

SUMMARY: The Federal Communications Commission, 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(s), 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: Persons wishing to comment on this information collection should submit comments June 6, 2005. 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 Paperwork Reduction Act (PRA) comments to Judith B. Herman, Federal Communications Commission, 445 12th Street, SW., Room 1–C804, Washington, DC 20554 or via the internet to *Judith-B.Herman@fcc.gov*.

FOR FURTHER INFORMATION CONTACT: For additional information or copies of the information collections contact Judith B. Herman at 202–418–0214 or via the internet at *Judith-B.Herman@fcc.gov*.

SUPPLEMENTARY INFORMATION:

OMB Control No.: 3060–1005.

Title: Numbering Resource Optimization—Phase 3.

Form No.: N/A.

Type of Review: Extension of a currently approved collection.

Respondents: Business or other for profit and state, local or tribal government.

Number of Respondents: 53.

Estimated Time Per Response: 50–85 hours.

Frequency of Response: On occasion reporting requirement and third party disclosure requirement.

Total Annual Burden: 3,380 hours.

Annual Cost Burden: $12,000.

Privacy Act Impact Assessment: N/A.

Needs and Uses: In the Communications Act of 1934, as amended by the Telecommunications Act of 1996, the Federal Communications Commission (''Commission'') was given ''exclusive jurisdiction over those portions of the North American Numbering Plan (NANP) that pertains to the United States.'' In order for price cap local exchange carriers (LECs) to qualify for exogenous adjustment to access charges established under the federal cost recovery mechanism, they must demonstrate that pooling results in a net cost increase rather than a cost reduction. Applications to state commissions from carriers must demonstrate that certain requirements are met before states grant any use of the safety valve mechanism. State commissions seeking to implement service-specific and/or technology-specific area code overlays, must request delegated authority to do so.

OMB Control No.: 3060–1012.

Title: Schools and Libraries Universal Service Support Mechanism, CC Docket No. 02–6, Notice of Proposed Rulemaking (NPRM), Proposed Americans with Disabilities Act (ADA) Certification.

Form No.: N/A.

Type of Review: Extension of a currently approved collection.

Respondents: Business or other for profit and not-for-profit institutions.

Number of Respondents: 30,000.

Estimated Time Per Response: 2.5 minutes (0.4 hours).

Frequency of Response: On occasion reporting requirement and third party disclosure requirement.

Total Annual Burden: 1,200 hours.

Annual Cost Burden: N/A.

Privacy Act Impact Assessment: N/A.

Needs and Uses: The Commission is seeking an extension (no change) to this OMB-approved information collection. After the 60-day comment period, the Commission will submit this information collection to OMB in order to obtain the full three-year clearance from them. The NPRM solicited comment on whether the Commission