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. What Action is the Agency Taking?

EPA is printing a summary of a pesticide petition received under section 408 of the Federal Food, Drug, and Cosmetic Act (FFDCA), 21 U.S.C. 346a, proposing the establishment or amendment of regulations in 40 CFR part 180 for residues of pesticide chemicals in or on various food commodities. EPA has determined that this pesticide petition contains data or information regarding the elements set forth in FFDCA section 408(d)(2); however, EPA has not fully evaluated the sufficiency of the submitted data at this time or whether the data support granting of the pesticide petition. Additional data may be needed before EPA rules on this pesticide petition.

Pursuant to 40 CFR 180.7(f), a summary of the petition included in this notice, prepared by the petitioner along with a description of the analytical method available for the detection and measurement of the pesticide chemical residues is available on EPA's Electronic Docket at *http://www.regulations.gov*. To locate this information on the home page of EPA's Electronic Docket, select ''Quick Search'' and type the OPP docket ID number. Once the search has located the docket, clicking on the ''Docket ID'' will bring up a list of all documents in the docket for the pesticide including the petition summary.

New Tolerance

PP 0F6076. Nissan Chemical Industries, Ltd. (Nissan), 7-1, 3-Chome, Kanda-Nishiki-Cho Chiyoda-Ku, Tokyo, 101-0054 Japan, proposes to establish a tolerance for residues of the herbicide quizalofop-P-ethyl in or on barley, flax (seed), and wheat at 0.05 parts per million (ppm); and sunflower (seed) at 2.0 ppm. The analytical method used to collect sunflower and flax field and processing data involves refluxing samples with methanolic potassium hydroxide to convert quizalofop-P-ethyl and quizalofop-P residues to 2-methoxy-6-chloroquinoxaline (MeCHQ). The solution is then acidified and partitioned with hexane to extract the MeCHQ. The hexane fraction is cleaned up by gel permeation chromatography. The appropriate fraction is collected, concentrated and made up to final volume with hexane. Residues are quantified using normal phase high pressure liquid chromatography (HPLC) with fluorescence detection. The limit of quantitation of the method is 0.05 ppm. The analytical method used to collect wheat and barley field and processing data is similar to the method used for flax and sunflower, but has a few modifications. The modified method requires a silica solid phase extraction (SPE) purification for wheat and barley hay and straw matrices prior to gel permeation chromatography (GPC) cleanup. The determination and quantitation of the MeCHQ is conducted using reverse-phase HPLC with the fluorescence detection. The limit of quantitation of the method is still 0.05 ppm.

List of Subjects

Environmental protection, Agricultural commodities, Feed additives, Food additives, Pesticides and pests, Reporting and recordkeeping requirements.

Dated: July 26, 2006.

Lois Rossi,

Director, Registration Division, Office of Pesticide Programs.

[FR Doc. E6–12469 Filed 8–1–06; 8:45 am]

BILLING CODE 6560–50–S

ENVIRONMENTAL PROTECTION AGENCY

[EPA–HQ–OPPT–2003–0006; FRL–8078–7]

TSCA Chemical Testing; Receipt of Test Data

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice.

SUMMARY: This notice announces EPA's receipt of test data on *In Vitro* Dermal Absorption Rate Testing of certain chemicals of interest to the Occupational Safety and Health Administration (OSHA). EPA received data on the following chemicals: Dipropylene glycol methyl ether (DPGME) (CAS No. 34590–94–8); naphthalene (CAS No. 91–20–3); diphenylamine (DPA) (CAS No. 122–39–4); 1–nitropropane (CAS No. 108–03–2); 2–nitropropane (CAS No. 79–46–9); isophorone (CAS No. 78–59–1); p-nitrochlorobenzene (CAS No. 100–00–5); and benzyl chloride (CAS No. 100–44–7). These data were submitted pursuant to a test rule issued by EPA under section 4 of the Toxic Substances Control Act (TSCA).

FOR FURTHER 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*.

SUPPLEMENTARY INFORMATION:

I. General Information

A. Does this Action Apply to Me?

This action is directed to the public in general. This action may, however, be of interest to those persons who are concerned about data on health and/or environmental effects and other characteristics of this chemical. 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 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–OPPT–2003–0006. Publicly available docket materials are available electronically at *http://www.regulations.gov* or in hard copy at the OPPT Docket, EPA Docket Center (EPA/DC), EPA West, Room B102, 1301 Constitution Ave., NW., Washington, DC. The EPA Docket Center Public Reading Room is open from 8:30 a.m. to 4:30 p.m., Monday through Friday, excluding legal holidays. The telephone number for the Public Reading Room is (202) 566–1744, and the telephone number for the OPPT Docket 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/*.

II. Test Data Submissions

Section 4(d) of TSCA requires EPA to publish a notice in the **Federal Register** reporting the receipt of test data submitted pursuant to test rules promulgated under section 4(a) within 15 days after these data are received by EPA.

1. Test data for DPGME were submitted by the DPGME Dermal Absorption Task Group of the American Chemistry Council's Ethylene and Propylene Glycol Ethers Panel and received by EPA on February 22, 2006. The submission includes a final report titled ''Dipropylene Glycol Methyl Ether: *In Vitro* Dermal Absorption Rate Testing''. (See Document ID No. EPA–HQ–OPPT–2003–0006–0325).

2. Test data for naphthalene were submitted on behalf of the American Chemistry Council Naphthalene Panel's *In Vitro* Dermal Absorption Rate Testing Consortium and received by EPA on March 20, 2006. The submission includes a final report titled ''Naphthalene: *In Vitro* Dermal Absorption Rate Testing.'' (See Document ID No. EPA–HQ–OPPT–2003–0006–0328).

3. Test data for DPA were submitted by Chemtura Corporation and received by EPA on March 29, 2006. The submission includes a final report titled ''Determination of the *In Vitro* Absorption Rate of Diphenylamine.'' (See Document ID No. EPA–HQ–OPPT–2003–0006–0330).

4. Test data for 1–nitropropane were submitted by the Angus Chemical Company, a wholly owned subsidiary of the Dow Chemical Company, and received by EPA on October 13, 2005. The submission includes a final report titled ''1–Nitropropane: *In Vitro* Dermal Absorption Rate Testing.'' (See Document ID No. EPA–HQ–OPPT–2003–0006–0343).

5. Test data for 2–nitropropane were submitted by the Angus Chemical Company, a wholly owned subsidiary of the Dow Chemical Company, and received by EPA on October 13, 2005. The submission includes a final report titled ''2–Nitropropane: *In Vitro* Dermal Absorption Rate Testing.'' (See Document ID No. EPA–HQ–OPPT–2003–0006–0344).

6. Test data for isophorone were submitted by the Isophorone Dermal Absorption Task Group of the American Chemistry Council and received by EPA on February 14, 2006. An amended report was also received by EPA on April 17, 2006. The submissions include an original and amended final study report titled: ''Percutaneous Absorption and Cutaneous Disposition of [^{14}C]-Isophorone *In Vitro* in Human Skin.'' (See Document ID No. EPA–HQ–OPPT–2003–0006–0346).

7. Test data for p-nitrochlorobenzene were submitted by ATK Thiokol and received by EPA on May 25, 2006. The submission includes a final study report titled: ''p-Nitrochlorobenzene: *In Vitro* Dermal Absorption Rate Testing.'' (See Document ID No. EPA–HQ–OPPT–2003–0006–0351).

8. Test data for benzyl chloride were submitted by LANXESS Corporation and Ferro Corporation and received by EPA on May 30, 2006. The submission includes a final study report titled: ''Benzyl Chloride: *In Vitro* Dermal Absorption Rate Testing.'' (See Document ID No. EPA–HQ–OPPT–2003–0006–0352).

These chemical substances are used in a wide variety of applications as industrial solvents, which may result in exposures of a substantial number of workers as described in the support document for the proposed rule (64 FR 31074, June 9, 1999, Table 3–Exposure Information for Chemical Substances).

EPA has initiated its review and evaluation process for these submissions. At this time, the Agency is unable to provide any determination as to the completeness of these submissions.

Authority: 15 U.S.C. 2603.

List of Subjects

Environmental protection, Hazardous substances.

Dated: July 20, 2006.

James Willis,

Director, Chemical Control Division, Office of Pollution Prevention and Toxics.

[FR Doc. E6–12340 Filed 8–1–06; 8:45 am]

BILLING CODE 6560–50–S

FEDERAL COMMUNICATIONS COMMISSION

Notice of Public Information Collection(s) Being Submitted for Review to the Office of Management and Budget

July 19, 2006.

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 (PRA) 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 Paperwork Reduction Act (PRA) comments should be submitted on or before October 2, 2006. If you anticipate that you will be submitting PRA 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, Room 1-B441, 445 12th Street, SW., Washington, DC 20554 or via the Internet to *PRA@fcc.gov.* If you would like to obtain or view a copy of this information collection, you may do so by visiting the FCC PRA Web page at: *http://www.fcc.gov/omd/pra.*

FOR FURTHER INFORMATION CONTACT: For additional information or copies of the information collection(s), send an e-mail to *PRA@fcc.gov* or contact Judith B. Herman at 202–418–0214. If you would like to obtain or view a copy of this information collection, you may do so by visiting the FCC PRA Web page at: *http://www.fcc.gov/omd/pra.*

SUPPLEMENTARY INFORMATION:

OMB Control No.: 3060–XXXX.

Title: Prepaid Calling Card Service Provider Certification, WC Docket No. 05–68.

Form No.: N/A.

Type of Review: New collection.

Respondents: Business or other for-profit.

Number of Respondents: 787.

Estimated Time Per Response: 25 hours.

Frequency of Response: Quarterly reporting requirement, recordkeeping requirement and third party disclosure requirement.