analytes from animal matrices and LC/MS/MS detection. This method has been validated for the determination of penthiopyrad and its metabolites in chicken tissues, eggs, ruminant tissues and milk. The limit of quantification is 0.01 mg/kg for all animal matrix groups. Contact: Tawanda Maignan, (703) 308–8050, e-mail address:

maignan.tawanda@epa.gov. 5. *PP* 9G7677. (EPÁ–HQ–OPP–2010– 0346). State of Florida, Department of Citrus, 605 East Main Street, P.O. Box 9010, Bartow, FL 33831-9010, proposes to establish temporary tolerances in 40 CFR part 180 for residues of the fungicide 5-chloro-3-methyl-4-nitro-1Hpyrazole (CMNP) and its metabolite (5-chloro-4-nitro-1*H*-pyrazol-3-yl)methanol (CHNP), in or on orange at 0.80 ppm; and its processed commodities: Orange, juice at 0.025 ppm; orange, oil at 0.070 ppm; orange, dried pulp (also referred to as dried pomace) at 1.80 ppm. In all plant matrices, the residue of concern, parent CMNP and CHNP/CHNP glucoside, can be determined using HPLC/MS/MS following sample extraction, hydrolysis (to convert CHNP-glucoside to its aglycone, CHNP) and solid-phase cleanup. Contact: Tawanda Maignan, (703) 308-8050, e-mail address: maignan.tawanda@epa.gov.

Amended Tolerance

PP 0F7776. (EPA-HQ-OPP-2009-0012 Dow AgroSciences LLC, 9330 Zionsville Road, Indianapolis, IN 46268, proposes to reestablish the time-limited tolerances in 40 CFR 180.544 for indirect or inadvertent combined residues of the insecticide methoxyfenozide, (benzoic acid, 3methoxy-2-methyl-, 2-(3,5dimethylbenzoyl)-2-(1,1-dimethylethyl) hydrazide) and its metabolites RH-117,236 free phenol of methoxyfenozide; 3,5-dimethylbenzoic acid N-tert-butyl-N'-(3-hydroxy-2methylbenzoyl) hydrazide, RH-151,055 glucose conjugate of RH-117,236; 3,5dimethylbenzoic acid N-tert-butyl-N-[3 (β-D-glucopyranosyloxy)-2methylbenzoyl]-hydrazide) and RH-152,072 the malonylglycosyl conjugate of RH-117,236, in or on the raw agricultural commodities: Vegetable, root and tuber, group 1 at 0.1 ppm; vegetable, leaves of root and tuber, group 2 at 0.2 ppm; vegetable, bulb, group 3 at 0.2 ppm; vegetable, legume, group 6 at 0.1 ppm; vegetable, foliage of legume, group 7 at 10 ppm; grain, cereal, forage, fodder, and straw, group 16 at 10 ppm; grass, forage, fodder and hay, group 17 at 10 ppm; animal feed, non-grass, group 18 at 10 ppm; and herb and spice, group 19 at 10 ppm. Rohm

and Haas Company, requested these tolerances under the Federal Food, Drug and Cosmetic Act, as amended by the Food Quality Protection Act of 1996. A Notice of Filing was submitted and published in the Federal Register of March 19, 2001 (66 FR 15443) (FRL 6766–7). Based on the data submitted by Rohm and Haas Company, the Agency determined that only time-limited tolerances for these residues could be established. The Final Rule was published in the Federal Register of September 20, 2002 (67 FR 59193) (FRL-7198-5) with time-limited tolerances expiring on September 30, 2007. To enable establishment of permanent tolerances, 24 additional rotational crop trials were requested. The data were submitted to the Agency on March 3, and June 17, 2003. A Final Rule extending these time-limited tolerances to September 30, 2010, was subsequently published in the Federal Register of March 5, 2008 (73 FR 11820) (FRL-8352-2). A further extension of the tolerances set to expire September 30, 2013, is needed to allow for conclusion of the Agency review of the additional rotational crop data. Adequate enforcement methods are available for determination of methoxyfenozide residues in plant commodities, based on the Rohm and Haas Company Technical Report No. 34-98-87, "Tolerance Enforcement Method for Parent RH-2485 in Pome Fruit". The available Analytical Enforcement Methodology was previously reviewed in the Federal Register of September 20, 2002 (67 FR 59193) (FRL-7198-5). Contact: Clayton Myers, (703) 347-8874, e-mail address: myers.clayton@epa.gov.

List of Subjects

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

Dated: October 14, 2010.

G. Jeffrey Herndon,

Acting Director, Registration Division, Office of Pesticide Programs.

[FR Doc. 2010–26731 Filed 10–26–10; 8:45 am]

BILLING CODE 6560-50-P

ENVIRONMENTAL PROTECTION AGENCY

[EPA-HQ-OPP-2010-0008; FRL-8847-4]

Pesticide Products; Registration Applications

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 November 26, 2010.

ADDRESSES: Submit your comments, identified by the docket identification (ID) number for the pesticide of interest, specified within Unit II., 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 regulations.gov or email. The 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 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.

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 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. 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 vou 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). If you are commenting in a docket that addresses multiple products, please indicate to which registration number(s) your comment applies.
- 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. Registration Applications

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

- 1. Registration number: 7969–274. Docket number: EPA–HQ–OPP–2010–0616. Company name and address: BASF Corporation, 26 Davis Drive, Research Triangle Park, NC 27709–3528. Active ingredients: Boscalid and Pyraclostrobin. Proposed use(s): Seed treatment on rapeseed (cultivars, varieties, and/or hybrids, including canola and crambe). Contact: Heather Garvie, (703) 308–0034; garvie.heather@epa.gov.
- 2. Registration number: 7969–275.
 Docket number: EPA–HQ–OPP–2010–
 0755. Company name and address:
 BASF Corporation, 26 Davis Drive,
 Research Triangle Park, NC 27709–3528.
 Active ingredient: Saflufenacil.
 Proposed use(s): For the manufacture of herbicides for use as a harvest aid/desiccant in dry edible beans, dry peas, soybean, oilseeds canola/rapeseed subgroup 20A, oilseeds sunflower subgroup 20B, and oilseeds cottonseed subgroup 20C. Contact: Susan Stanton, (703) 305–5218; stanton.susan@epa.gov.
- 3. Registration number: 7969–278.

 Docket number: EPA–HQ–OPP–2010–
 0755. Company name and address:
 BASF Corporation, 26 Davis Drive,
 Research Triangle Park, NC 27709–3528.

 Active ingredient: Saflufenacil.

 Proposed use(s): As a harvest aid/
 desiccant in dry edible beans, dry peas,
 soybean, oilseeds canola/rapeseed
 subgroup 20A, oilseeds sunflower
 subgroup 20B, and oilseeds cottonseed
 subgroup 20C. Contact: Susan Stanton,
 (703) 305–5218, stanton.susan@epa.gov.
- 4. File symbol: 56799–U. Docket number: EPA–HQ–OPP–2010–0707. Company name and address: Productos Quimicos y Alimenticios OSKU S.A. El Guanaco 5212, Huechuraba, Santiago, Chile. Active ingredient: Sulfur Dioxide (from Sodium metabisulfite). Proposed use(s): Blueberries. Contact: Rosemary Kearns, (703) 305–5611, kearns.rosemary@epa.gov.
- 5. Registration numbers: 66330–64, 66330–65. Docket number: EPA–OPP–2010–0725. Company name and address: Arysta LifeScience North America Corporation, 15401 Weston Parkway, Suite 150, Cary, NC 27513. Active ingredient: Fluoxastrobin. Proposed use(s): Squash/cucumber subgroup 9B. Contact: Heather Garvie, (703) 308–0034; garvie.heather@epa.gov.

6. File symbol: 72500–EN. Docket number: EPA–HQ–OPP–2010–0769. Company name and address: Scimetrics, 9974 NE Frontage Rd., Wellington, CO 80549. Active ingredients: Warfarin and Imidacloprid. Proposed use(s): Rangeland and noncrop areas to control black-tailed and white-tailed prairie dogs and their fleas. Contact: Daniel Peacock, (703) 305–5407, peacock.dan@epa.gov.

7. File symbol: 82052–T. Docket number: EPA–HQ–OPP–2010–0767. Company name and address: Cutting Edge Formulation, Inc., 3057 Summer Oak Place, Buford, GA 30518. Active ingredient: D-limonene. Proposed use(s): Bacterial disease control by suppression of citrus canker. Contact: Rita Kumar, (703) 308–8291, kumar.rita@epa.gov.

List of Subjects

Environmental protection, Pesticides and pest.

Dated: October 15, 2010.

Daniel J. Rosenblatt,

Acting Director, Registration Division, Office of Pesticide Programs.

[FR Doc. 2010–26886 Filed 10–26–10; 8:45 am]

BILLING CODE 6560-50-P

FEDERAL COMMUNICATIONS COMMISSION

Notice of Public Information Collection(s) Being Submitted for Review and Approval to the Office of Management and Budget (OMB), Comments Requested

October 25, 2010.

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, 44 U.S.C. 3501-3520. 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; (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; and (e) ways to further reduce the information collection burdens on small

business concerns with fewer than 25 employees.

The FCC may not conduct or sponsor a collection of information unless it displays a currently valid OMB 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 OMB control number.

DATES: Submit written Paperwork Reduction Act (PRA) comments on or before November 26, 2010. 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 FCC contact listed below as soon as possible.

ADDRESSES: Submit all PRA comments to Nicholas A. Fraser, Office of Management and Budget, via fax at 202-395–5167 or the Internet at Nicholas A. Fraser@omb.eop.gov; and to Judith-B. Herman@fcc.gov, Federal Communications Commission. Send your PRA comments by e-mail to PRA@fcc.gov. To view a copy of this information collection request (ICR) submitted to OMB: (1) Go to the Web page http://reginfo.gov/public/do/ PRAMain, (2) look for the section of the Web page called "Currently Under Review", (3) click on the downwardpointing arrow in the "Select Agency" box below the "Currently Under Review" heading, (4) select "Federal Communications Commission" from the list of agencies presented in the "Select Agency" box, (5) click the "Submit" button to the right of the "Select Agency" box, and (6) when the list of FCC ICRs currently under review appears, look for the title of this ICR (or its OMB Control Number, if there is one) and then click on the ICR Reference Number to view detailed information about this ICR.

FOR FURTHER INFORMATION CONTACT: For additional information or copies of the information collection(s), contact Judith B. Herman at 202–418–0214 or via the Internet at Judith-B.Herman@fcc.gov.

SUPPLEMENTARY INFORMATION:

OMB Control Number: 3060–0999. Title: Hearing Aid Compatibility Status Report and Section 20.19, Hearing Aid-Compatible Mobile Handsets (Hearing Aid Compatibility Act).

Form No.: FCC Form 655—electronic only.

Type of Review: Revision of a currently approved collection.

Respondents: Business or other forprofit.

Number of Respondents: 925 respondents; 925 responses. Estimated Time per Response: 1

hour—2.5 hours.

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

Obligation to Respond: Required to obtain or retain benefits. Statutory authority for this information collection is contained in 47 U.S.C. 151, 154(i), 157, 160, 201, 202, 208, 214, 301, 303, 308, 309(j), 310 and 610.

Total Annual Burden: 12,063 hours. Total Annual Cost: N/A.

Privacy Act Impact Assessment: N/A. Nature and Extent of Confidentiality: Information in the reports may include confidential information. However, covered entities would be allowed to request that such materials submitted to the Commission be withheld from public inspection under 47 CFR 0.459 of the Commission's rules.

Needs and Uses: The Commission will submit this revised information collection (IC) to the Office of Management and Budget (OMB) during this comment period to obtain the full three-year clearance from them. The Commission is reporting no change in the burden estimates as previously

approved by the OMB.

On August 5, 2010, the Commission adopted final rules in a Second Report and Order, FCC 10-145, that, among other things, updated disclosure requirements for manufacturers and service providers. Manufacturers and service providers are now required to adequately inform consumers about the functionality and the limitations of their handsets in two specific situations. First, for handsets that meet hearing aid compatibility requirements over all air interfaces and frequency bands for which technical standards have been established, but that are also capable of supporting voice operations in new frequency bands and air interfaces for which standards do not exist, the following mandatory disclosure language must be clearly and effectively conveyed to consumers wherever the hearing aid compatibility rating for the handset is provided, including the point of sale and on company Web sites:

"This phone has been tested and rated for use with hearing aids for some of the wireless technologies that it uses. However, there may be some newer wireless technologies used in this phone that have not been tested yet for use with hearing aids. It is important to try the different features of this phone thoroughly and in different locations, using your hearing aid or cochlear implant, to determine if you hear any interfering noise. Consult with your