

3. *By hand delivery or courier.* Deliver your comments to: Public Information and Records Integrity Branch (PIRIB), Office of Pesticide Programs (OPP), Environmental Protection Agency, Rm. 119, Crystal Mall #2, 1921 Jefferson Davis Hwy., Arlington, VA, Attention: Docket ID Number OPP-2002-0146. Such deliveries are only accepted during the docket's normal hours of operation as identified in Unit I.B.1.

D. How Should I Handle CBI That I Want to Submit to the Agency?

Do not submit information that you consider to be CBI electronically through EPA's electronic public docket or by e-mail. You may claim information that you submit to EPA as CBI by marking any part or all of that information as CBI (if you submit CBI on disk or CD ROM, 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 CBI). Information so marked will not be disclosed except in accordance with procedures set forth in 40 CFR part 2.

In addition to one complete version of the comment that includes any 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 and EPA's electronic public docket. If you submit the copy that does not contain CBI on disk or CD ROM, mark the outside of the disk or CD ROM clearly that it does not contain CBI. Information not marked as CBI will be included in the public docket and EPA's electronic public docket without prior notice. If you have any questions about CBI or the procedures for claiming CBI, please consult the person listed under **FOR FURTHER INFORMATION CONTACT**.

E. What Should I Consider as I Prepare My Comments for EPA?

You may find the following suggestions helpful for preparing your comments:

1. Explain your views as clearly as possible.
2. Describe any assumptions that you used.
3. Provide copies of any technical information and/or data you used that support your views.
4. If you estimate potential burden or costs, explain how you arrived at the estimate that you provide.
5. Provide specific examples to illustrate your concerns.
6. Offer alternative ways to improve the notice or collection activity.
7. Make sure to submit your comments by the deadline in this document.

8. To ensure proper receipt by EPA, be sure to identify the docket control number assigned to this action in the subject line on the first page of your response. You may also provide the name, date, and **Federal Register** citation.

II. Background

A. What Action is the Agency Taking?

This notice constitutes and announces the availability of the TRED for tebutiuron. This decision has been developed as part of the public participation process that EPA and the U.S. Department of Agriculture (USDA) are using to involve the public in the reassessment of pesticide tolerances under FFDCA. EPA must review tolerances and tolerance exemptions that were in effect when FQPA was enacted in August 1996, to ensure these existing pesticide residues limits for food and feed commodities meet the safety standard of the new law.

FFDCA requires EPA to review all the tolerances for registered chemicals in effect on or before the date of the enactment. In reviewing these tolerances, the Agency must consider, among other things, aggregate risks from non-occupational sources of pesticide exposure, whether there is increased susceptibility to infants and children, and the cumulative effects of pesticides with a common mechanism of toxicity. The tolerances are considered reassessed once the safety finding has been made or a revocation occurs.

FFDCA requires that the Agency, when considering whether to establish, modify, or revoke a tolerance, consider "available information" concerning the cumulative effects of a particular pesticide's residues and "other substances that have a common mechanism of toxicity."

As indicated above, the Agency will also evaluate the cumulative risk, if necessary, posed by the entire group of chemicals with which a common mechanism of toxicity is shared, and issues a final tolerance reassessment decision once the cumulative assessment for that group is completed. At this time, tebutiuron has not been identified as sharing a common mechanism of toxicity and is not scheduled for a cumulative risk assessment.

The tolerance reassessment program is being conducted under Congressionally mandated time frames, and EPA recognizes both the need to make timely tolerance decisions and to involve the public. Therefore, EPA is issuing this TRED as a final document with a 30-day comment period. All

comments will be considered by the Agency. If any comment significantly affects a TRED, EPA will amend the TRED by publishing the amendment in the **Federal Register**.

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

The authority for this TRED is found in section 408(q) of the FFDCA, 21 U.S.C. 346a(q). Section 408(q) requires EPA to review tolerances and exemptions for pesticide chemical residues in effect on August 2, 1996, to determine whether the tolerance or exemption meets the requirements of 408(b)(2) or (c)(2). This review is to be completed by August 3, 2006.

List of Subjects

Environmental protection, Chemicals, Pesticides and tolerances.

Dated: March 19, 2003.

Lois A. Rossi,

Director, Special Review and Reregistration Division, Office of Pesticide Programs.

[FR Doc. 03-7981 Filed 4-1-03; 8:45 am]

BILLING CODE 6560-50-S

ENVIRONMENTAL PROTECTION AGENCY

[OPP-2003-0097; FRL-7298-7]

Thiamethoxam; Notice of Filing a Pesticide Petition to Establish a Tolerance for a Certain Pesticide Chemical in or on Food

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice.

SUMMARY: This notice announces the initial filing of a pesticide petition proposing the establishment of regulations for residues of a certain pesticide chemical in or on various food commodities.

DATES: Comments, identified by docket ID number OPP-2003-0097, must be received on or before May 2, 2003.

ADDRESSES: Comments may be submitted electronically, by mail, or through hand delivery/courier. Follow the detailed instructions as provided in Unit I. of the **SUPPLEMENTARY INFORMATION**.

FOR FURTHER INFORMATION CONTACT: Dani Daniel, Registration Division (7505C), Office of Pesticide Programs, Environmental Protection Agency, 1200 Pennsylvania Ave., NW., Washington, DC 20460-0001; telephone number: (703) 305-5409; e-mail address: daniel.dani@epa.gov.

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 111)
- Animal production (NAICS 112)
- Food manufacturing (NAICS 311)
- Pesticide manufacturing (NAICS

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. How Can I Get Copies of this Document and Other Related Information?

1. *EPA Docket.* EPA has established an official public docket for this action under docket ID number OPP-2003-0097. 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, 1921 Jefferson Davis Hwy., 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.

Certain types of information will not be placed in the EPA dockets. Information claimed as CBI and other information whose disclosure is restricted by statute, which is not included in the official public docket, will not be available for public viewing in EPA's electronic public docket. EPA's policy is that copyrighted material will not be placed in EPA's electronic public docket but will be available only in printed, paper form in the official public docket. To the extent feasible, publicly available docket materials will be made available in EPA's electronic public docket. When a document is selected from the index list in EPA dockets, the system will identify whether the document is available for viewing in EPA's electronic public docket. 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. EPA intends to work towards providing electronic access to all of the publicly available docket materials through EPA's electronic public docket.

For public commenters, it is important to note that EPA's policy is that public comments, whether submitted electronically or on paper, will be made available for public viewing in EPA's electronic public docket as EPA receives them and without change, unless the comment contains copyrighted material, CBI, or other information whose disclosure is restricted by statute. When EPA identifies a comment containing copyrighted material, EPA will provide a reference to that material in the version of the comment that is placed in EPA's electronic public docket. The entire printed comment, including the copyrighted material, will be available in the public docket.

Public comments submitted on computer disks that are mailed or delivered to the docket will be transferred to EPA's electronic public docket. Public comments that are mailed or delivered to the docket will be scanned and placed in EPA's electronic public docket. Where practical, physical objects will be photographed, and the photograph will be placed in EPA's electronic public docket along with a

brief description written by the docket staff.

C. How and to Whom Do I Submit Comments?

You may submit comments electronically, by mail, or through hand delivery/courier. To ensure proper receipt by EPA, identify the appropriate docket ID number in the subject line on the first page of your comment. Please ensure that your comments are submitted within the specified comment period. Comments received after the close of the comment period will be marked "late." EPA is not required to consider these late comments. If you wish to submit CBI or information that is otherwise protected by statute, please follow the instructions in Unit I.D. Do not use EPA dockets or e-mail to submit CBI or information protected by statute.

1. *Electronically.* If you submit an electronic comment as prescribed in this unit, EPA recommends that you include your name, mailing address, and an e-mail address or other contact information in the body of your comment. Also, include this contact information on the outside of any disk or CD ROM you submit, and in any cover letter accompanying the disk or CD ROM. This ensures that you can be identified as the submitter of the comment and allows EPA to contact you in case EPA cannot read your comment due to technical difficulties or needs further information on the substance of your comment. EPA's policy is that EPA will not edit your comment, and any identifying or contact information provided in the body of a comment will be included as part of the comment that is placed in the official public docket, and made available in EPA's electronic public docket. 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.

i. *EPA Dockets.* Your use of EPA's electronic public docket to submit comments to EPA electronically is EPA's preferred method for receiving comments. Go directly to EPA dockets at <http://www.epa.gov/edocket/>, and follow the online instructions for submitting comments. Once in the system, select "search," and then key in docket ID number OPP-2003-0097. The system is an "anonymous access" system, which means EPA will not know your identity, e-mail address, or other contact information unless you provide it in the body of your comment.

ii. *E-mail.* Comments may be sent by e-mail to opp-docket@epa.gov, Attention: Docket ID Number OPP-2003-0097. In contrast to EPA's

electronic public docket, EPA's e-mail system is not an "anonymous access" system. If you send an e-mail comment directly to the docket without going through EPA's electronic public docket, EPA's e-mail system automatically captures your e-mail address. E-mail addresses that are automatically captured by EPA's e-mail system are included as part of the comment that is placed in the official public docket, and made available in EPA's electronic public docket.

iii. *Disk or CD ROM.* You may submit comments on a disk or CD ROM that you mail to the mailing address identified in Unit I.C.2. These electronic submissions will be accepted in WordPerfect or ASCII file format. Avoid the use of special characters and any form of encryption.

2. *By mail.* Send your comments to: Public Information and Records Integrity Branch (PIRIB) (7502C), Office of Pesticide Programs (OPP), Environmental Protection Agency, 1200 Pennsylvania Ave., NW., Washington, DC 20460-0001, Attention: Docket ID number OPP-2003-0097.

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4. If you estimate potential burden or costs, explain how you arrived at the estimate that you provide.
5. Provide specific examples to illustrate your concerns.
6. Make sure to submit your comments by the deadline in this notice.
7. To ensure proper receipt by EPA, be sure to identify the docket ID number assigned to this action in the subject line on the first page of your response. You may also provide the name, date, and **Federal Register** citation.

II. What Action is the Agency Taking?

EPA has received a pesticide petition as follows proposing the establishment and/or amendment of regulations for residues of a certain pesticide chemical in or on various food commodities under section 408 of the Federal Food, Drug, and Cosmetic Act (FFDCA), 21 U.S.C. 346a. EPA has determined that this 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 petition. Additional data may be needed before EPA rules on the petition.

List of Subjects

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

Dated: March 24, 2003.

Debra Edwards,

Acting Director, Registration Division, Office of Pesticide Programs.

Summary of Petition

The petitioner's summary of the pesticide petitions is printed below as required by FFDCA section 408(d)(3). The summary of the petitions was

prepared by Interregional Research Project 4 (IR-4) and represents the view of the petitioners. The petition summary announces the availability of a description of the analytical methods available to EPA for the detection and measurement of the pesticide chemical residues or an explanation of why no such method is needed.

Interregional Research Project 4 (IR-4) Syngenta Crop Protection Inc.

PP 2E6505, 2E6363, 2E6508, 3E6524, 1E6349, 9F5051 and 0F6142

This notice is a summary of pesticide petitions proposing the establishment/ amendment of a regulation for residues of thiamethoxam and its metabolite in or on coffee (imported), pecans, leafy vegetable crop group, head and stem brassica subgroup, leafy brassica subgroup, succulent beans, stone fruit crop group, sunflower seed, peppermint tops and spearmint tops. This summary was prepared by the petitioners.

EPA has received seven pesticide petitions; four from Interregional Research Project 4 (IR-4), 681 U.S. Highway #1 South, North Brunswick, NJ 08902-3390, PP 2E6505, 2E6363, 2E6508 and 3E6524 and three from Syngenta Crop Protection Inc., P.O. Box 18300, Greensboro, NC 27419-8300, PP 0F6142, 1E6349, and 9F5051 proposing, pursuant to section 408(d) of the Federal Food, Drug, and Cosmetic Act (FFDCA), 21 U.S.C. 346a(d), to amend 40 CFR 180.565 by establishing a tolerance for residues of the insecticide thiamethoxam [3-(chloro-5-thiazolyl)methyl]tetrahydro-5-methyl-N-nitro-4H-1,3,5-oxadiazin-4-imine] (CAS Reg. No. 153719-23-4) and its metabolite N-(2-chloro-thiazol-5-ylmethyl)-N'-methyl-N'-nitro-guanidine) in or on the agricultural commodities:

A. IR-4 Petitions

1. PP 2E6505 proposes to establish tolerances for stone fruits, group 12 at 0.5 ppm.

2. PP 2E6363 proposes to establish tolerances for peppermint and spearmint, tops at 4.0 ppm.

3. PP 2E6508 proposes to establish tolerances for beans, succulent at 0.02 ppm.

4. PP 3E6524 proposes to establish tolerances for sunflower, seed at 0.02 ppm.

B. Syngenta Petitions

5. PP 0F6142 proposes to establish tolerances for pecans at 0.02 ppm.

6. PP 9F5051 proposes to establish tolerances for:

- Leafy vegetables, group 4 at 2.0 ppm
- Head and stem brassica vegetables, subgroup 5A at 1.0 ppm.
- Leafy brassica vegetables, subgroup 5B at 2.0 ppm.

7. PP 1E6349 proposes to establish tolerances for imported green and roasted coffee beans and instant coffee at 0.05 ppm.

EPA has determined that the petitions contain data or information regarding the elements set forth in section 408(d)(2) of the FFDCa; however, EPA has not fully evaluated the sufficiency of the submitted data at this time or whether the data support granting of the petitions. Additional data may be needed before EPA rules on the petitions.

A. Residue Chemistry

1. *Plant metabolism.* The primary metabolic pathways of thiamethoxam in plants (corn, rice, pears, and cucumbers) were similar to those described for animals, with certain extensions of the pathway in plants. Parent compound and CGA-322704 were the major residues in all crops. The metabolism of thiamethoxam in plants and animals is understood for the purposes of the proposed tolerances. Parent thiamethoxam and the metabolite, CGA-322704, are the residues of concern for tolerance setting purposes.

2. *Analytical method.* Syngenta Crop Protection Inc. has submitted practical analytical methodology for detecting and measuring levels of thiamethoxam in or on raw agricultural commodities. The method is based on crop specific cleanup procedures and determination by liquid chromatography with either ultraviolet (UV) or mass spectroscopy (MS) detection. The limit of detection (LOD) for each analyte of this method is 1.25 nanogram (ng) injected for samples analyzed by UV and 0.25 ng injected for samples analyzed by MS, and the limit of quantitation (LOQ) is 0.005 ppm for milk and juices and 0.01 ppm for all other substrates.

3. *Magnitude of residues.* IR-4 has submitted complete residue data for thiamethoxam on succulent beans, sunflower seed, peppermint and spearmint tops and stone fruits. Syngenta has submitted complete residue data for the proposed imported coffee, pecan, leafy vegetable, head and stem brassica vegetables, leafy brassica vegetables. Details of the Syngenta residue data on these crops were provided in previously published Notices of Filing.

B. Toxicological Profile

1. *Acute toxicity.* The acute oral LD₅₀ for thiamethoxam in the rat is 1,563 milligrams/kilogram body weight (mg/kg bwt). The acute dermal LD₅₀ of thiamethoxam is >2000 mg/kg bwt. Thiamethoxam is non-toxic at atmospheric concentrations of 3.72 mg/L. Thiamethoxam is minimally irritating to the eye, non-irritating to skin and is not a dermal sensitizer.

In an acute neurotoxicity screening study in rats (OPPTS Harmonized Guideline 870.6200), the no observed adverse affect level (NOAEL) was 100 mg/kg/day with a NOAEL of 500 mg/kg/day based on drooped palpebral closure, decrease in rectal temperature and locomotor activity and increase in forelimb grip strength (males only). At higher dose levels, mortality, abnormal body tone, ptosis, impaired respiration, tremors, longer latency to first step in the open field, crouched over posture, gait impairment, hypo-arousal, decreased number of rears, uncoordinated landing during the righting reflex test, slight lacrimation (females only) and higher mean average input stimulus value in the auditory startle response test (males only).

2. *Genotoxicity.* In gene mutation studies with *S. typhimurium* and *E. coli* (OPPTS Harmonized Guidelines, 870.5100 and 870.5265), there was no evidence of gene mutation when tested up to 5,000 µg/plate and there was no evidence of cytotoxicity.

In a gene mutation study with chinese hamster V79 cells at hypoxanthine guanine phosphoribosyl transferase (HGPRT) focus (OPPTS Harmonized Guideline 870.5300) there was no evidence of a gene mutation when tested up to the solubility limit.

In a CHO cell cytogenetics study (OPPTS Harmonized Guideline 870.5375) there was no evidence of chromosomal aberrations when tested up to cytotoxic or solubility limit concentrations.

An *in vivo* mouse bone marrow micronucleus study (OPPTS Harmonized Guideline 870.5395) was negative when tested up to levels of toxicity in whole animals; however, there was no evidence of target cell cytotoxicity. An unscheduled DNA synthesis (UDS) assay (OPPTS Harmonized Guideline 870.5550) was negative when tested up to precipitating concentrations.

3. *Reproductive and developmental toxicity.* A prenatal developmental study in the rat (OPPTS Harmonized Guideline 870.3700) resulted in maternal and developmental NOAELs of 30 mg/kg/day and 200 mg/kg/day,

respectively. The maternal lowest observed adverse effect level (LOAEL) is 200 mg/kg/day based on decreased body weight, body weight gain and food consumption. The developmental LOAEL was 750 mg/kg/day based on decreased fetal body weight and an increased incidence of skeletal anomalies.

A prenatal developmental study in the rabbit (OPPTS Harmonized Guideline 870.3700) resulted in maternal and developmental NOAELs of 50 mg/kg/day. The maternal and developmental LOAEL is 150 mg/kg/day. The maternal LOAEL is based on maternal deaths, hemorrhagic discharge, decreased body weight and food intake during the dosing period. The developmental LOAEL is based on decreased fetal body weights, increased incidence of post-implantation loss and a slight increase in the incidence of a few skeletal anomalies/variations.

In a reproduction and fertility effects study in rats (OPPTS Harmonized Guideline 870.3800), the parental/systemic NOAEL is 1.84 (males), 202.06 (females) mg/kg/day; the reproductive NOAEL is 0.61 (males), 202.06 (females) mg/kg/day and the offspring NOAEL is 61.25 (males), 79.20 (females) mg/kg/day. The parental/systemic LOAEL is 61.25 (males), not determined (females) mg/kg/day based on increased incidence of hyaline change in renal tubules in F0 and F1 males. The reproductive LOAEL is 1.84 (males), not determined (females) mg/kg/day based on increased incidence and severity of tubular atrophy observed in testes of the F1 generation males. The offspring LOAEL is 158.32 (males), 202.06 (females) mg/kg/day based on reduced body weight gain during the lactation period in all litters.

4. *Subchronic toxicity.* A 90-day oral toxicity study in rats (OPPTS Harmonized Guideline 870.3100) resulted in a NOAEL of 1.74 males and 92.5 (females) mg/kg/day. The LOAEL is 17.64 (male) and 182.1 (female) mg/kg/day based on increased incidence of hyaline change of renal tubules epithelium (males), fatty change in adrenal gland of females, liver changes in females, all at the LOAEL.

A 90-day oral toxicity study in mice (OPPTS Harmonized Guideline 870.3100) resulted in an NOAEL of 1.41 (males) and 19.2 (females) mg/kg/day. The LOAEL was 14.3 (male) and 231 (female) mg/kg/day based on increased incidence of hepatocellular hypertrophy. At higher dose levels: Decrease in body weight and body weight gain, necrosis of individual hepatocytes, pigmentation of Kupffer cells, and lymphocytic infiltration of the

liver in both sexes; slight hematologic effects and decreased absolute and relative kidney weights in males; and ovarian atrophy, decreased ovary and spleen weights and increased liver weights in females.

In a 90-day oral toxicity study in dogs (OPPTS Harmonized Guideline 870.3150), the NOAEL is 8.23 (males) and 9.27 (females) mg/kg/day. The LOAEL is 32.0 (male) and 33.9 (female) mg/kg/day based on slightly prolonged prothrombin times and decreased plasma albumin and A/G ration (both sexes); decreased calcium levels and ovary weights and delayed maturation in the ovaries (female); decreased cholesterol and phospholipid levels, testis weights, spermatogenesis, and spermatid giant cells in testes (male).

In a 28-day dermal study in rats (OPPTS Harmonized Guideline 870.3200), the NOAEL was 250 (male) and 60 (female) mg/kg/day. The LOAEL was 1,000 (male) and 250 (female) mg/kg/day based on increased plasma glucose, triglyceride levels, and alkaline phosphatase activity and inflammatory cell infiltration in the liver and necrosis of single hepatocytes in females and hyaline change in renal tubules and a very slight reduction in body weight in males. At higher dose levels in females, chronic tubular lesions in the kidneys and inflammatory cell infiltration in the adrenal cortex were observed.

In a subchronic neurotoxicity screening study in rats (OPPTS Harmonized Guideline 870.6200) the NOAEL was 95.4 (male) and 216.4 (female) mg/kg/day, both at highest dose tested. The LOAEL was not determined. No treatment related observations at any dose level. LOAEL was not achieved. May not have been tested at sufficiently high dose levels; however, a new study is not required because the weight of the evidence from other toxicity studies indicates no evidence of concern.

5. Chronic toxicity. In a chronic toxicity study in dogs (OPPTS Harmonized Guideline 870.4100) the NOAEL was 4.05 (male) and 4.49 (female) mg/kg/day. The LOAEL was 21.0 (male) and 24.6 (female) mg/kg/day based on increase of creatinine in both sexes, transient decrease in food consumption in females, and occasional increase in urea levels, decrease in ALT, and atrophy of seminiferous tubules in males.

In a mouse carcinogenicity study (OPPTS Harmonized Guideline 870.4200), the NOAEL was 2.63 (male) and 3.68 (female) mg/kg/day. The LOAEL was 63.8 (male) and 87.6 (female) mg/kg/day based on hepatocyte hypertrophy, single cell necrosis, inflammatory cell infiltration, pigment

deposition, foci of cellular alteration, hyperplasia of kupffer cells and increased mitotic activity, also an increase in the incidence of hepatocellular adenoma (both sexes). At higher doses, there was an increase in the incidence of hepatocellular adenocarcinoma (both sexes) and the number of animals with multiple tumors, evidence of carcinogenicity.

In a combined chronic carcinogenicity study in rats (OPPTS Harmonized Guideline 870.4300) the NOAEL was 21.0 (male) and 50.3 (female) mg/kg/day. The LOAEL was 63.0 (male) and 255 (female) mg/kg/day based on increased incidence of lymphocytic infiltration of the renal pelvis and chronic nephropathy in males and decreased body weight gain, slight increase in the severity of hemosiderosis of the spleen, foci of cellular alteration in liver and chronic tubular lesions in kidney in females. No evidence of carcinogenicity.

In a hepatic cell proliferation study in mice, the NOAEL was 16 (male) and 20 (female) mg/kg/day. The LOAEL was 72 (male) and 87 (female) mg/kg/day based on proliferative activity of hepatocytes. At higher dose levels, increases in absolute and relative liver weights, speckled liver, hepatocellular glycogenesis/fatty change, hepatocellular necrosis, apoptosis and pigmentation were observed.

In a 28-day feeding study to assess replicative DNA synthesis in the male rat, the NOAEL was 711 mg/kg/day. The LOAEL was not established. Immunohistochemical staining of liver sections from control and high dose animals for proliferating cell nuclear antigen gave no indication for a treatment related increase in the fraction of DNA synthesizing hepatocytes in S-phase. CGA-293343 did not stimulate hepatocyte cell proliferation in male rats.

In a special study to assess liver biochemistry in the mouse, the NOAEL was 17 (male) and 92 (female) mg/kg/day. The LOAEL was 74 (male), 92 (female) mg/kg/day based on marginal to slight increases in absolute and relative liver weights, a slight increase in the microsomal protein content of the livers, moderate increases in the cytochrome P450 content, slight to moderate increases in the activity of several microsomal enzymes, slight to moderate induction of cytosolic glutathione S-transferase activity. Treatment did not affect peroxisomal fatty acid B-oxidation.

6. Animal metabolism. The metabolism of thiamethoxam in rats and livestock animals is adequately understood. The residues of concern

have been determined to be parent thiamethoxam and its metabolite N-(2-chloro-thiazol-5-ylmethyl)-N'-methyl-N-nitro-guanidine.

7. Metabolite toxicology. For risk assessment purposes, residues of the metabolite corrected for molecular weight are considered to be toxicologically equivalent to parent thiamethoxam.

C. Aggregate Exposure

1. Dietary exposure. Permanent tolerances have been established (40 CFR 180.565) for the combined residues of the insecticide thiamethoxam, 3-[(2-chloro-5-thiazolyl)methyl]tetrahydro-5-methyl-N-nitro-4H-1,3,5-oxadiazin-4-imine and its metabolite N-(2-chloro-thiazol-5-ylmethyl)-N'-methyl-N-nitro-guanidine, in or on a variety of raw agricultural commodities levels ranging from 0.02 parts per million (ppm) to 1.5 ppm (including barley, canola, corn, cotton, sorghum, wheat, cucurbit vegetables, fruiting vegetables, pome fruits, tuberous and corm vegetables and livestock commodities).

Pending tolerances include coffee (imported), grapes, raisins, grape juice, pecans, sunflower seed, stone fruits, succulent beans, peppermint and spearmint tops, head and stem brassica, leafy brassica greens and leafy vegetables.

Tier III chronic and Tier I acute dietary exposure evaluations were made using the Dietary Exposure Evaluation Model (DEEM™), version 7.76 from Exponent. All processing factors were taken from the EPA assessment of August 28, 2000 (DP Barcode D268606, PC Code 060109). These assessment results include all current tolerances and the proposed tolerances on stone fruit, mint, succulent beans, sunflower seed, pecans, leafy vegetables, leafy brassica vegetables, brassica head and stem vegetables and imported coffee.

For the Tier I acute assessment, the proposed tolerance residues for these commodities (stone fruit, 0.5 ppm; mint, 4.0 ppm; succulent beans, 0.02 ppm; sunflower seed, 0.02 ppm; pecans, 0.02 ppm; leafy vegetables, 2.0 ppm; leafy brassica vegetables, 2.0 ppm; brassica head and stem vegetables, 1.0 ppm; and imported coffee, 0.05 ppm) were used along with the published tolerances for all other commodities. One hundred percent of crop treated was assumed for all commodities in the acute assessment.

In the chronic assessments, residue values for secondary animal commodities, pome fruits, ginger, turmeric, peppers, potatoes, wheat and barley were taken from the EPA assessment of August 28, 2000 which

uses average field trial residue data with $\frac{1}{2}$ LOQ substitutions for all non-detectable residues. In addition, a residue value of 0.011 ppm ($\frac{1}{2}$ LOQ) and a percent crop treated value of 6.6% were used for all corn commodities. For the remaining registered and the proposed commodities listed above, the following residue data was used in the DEEM™: Cucurbit, leafy and brassica vegetables and tomatoes - average field trial residues from soil-only application residue studies; stone fruits, mint, succulent beans, sunflower seed, and coffee - average field trial residue data with $\frac{1}{2}$ LOQ substitutions for non-detectable residues; and pecans - the proposed tolerance.

In regard to the cucurbit vegetables and tomatoes, the current tolerances are based upon soil and foliar uses; however, Syngenta is currently limiting the use of thiamethoxam in these crops to soil only applications - thus, the refinement in DEEM™ inputs described above. Likewise, the proposed tolerances on leafy vegetables, leafy brassica vegetables and head and stem brassica vegetables are based upon soil and foliar applications of thiamethoxam; however, Syngenta is currently pursuing only the soil application use - thus, the refinement of DEEM inputs described above. Syngenta will pursue foliar applications for these crops at a later date; therefore, the proposed and current tolerances on these crops remain based upon soil and foliar applications.

All consumption data for these assessments was taken from the USDA's Continuing Survey of Food Intake by Individuals (CSFII) with the 1994-96 consumption data base and the supplemental CSFII children's survey (1998) consumption data base. For the chronic assessments, the following percent of crop treated values were used for the proposed uses: Coffee, 16.7%; sunflower, 15%; mint, 10%; leaf lettuce, 24.6%; head lettuce, 32%; spinach and cress, 15.6%; all other leafy vegetables, 19.4%; broccoli, 26.2%, cabbage, 15.3%; all other brassica vegetables 17.2%; beans, 20%; stone fruits, 15%; and pecans, 100%. A percent crop treated value of 5% was used for apples and a value of 6.6% was used for all corn commodities. All other percent of crop treated values were taken from the August 28, 2000 EPA assessment.

i. *Food*. For the purposes of assessing the potential dietary exposure under the proposed tolerances, Syngenta Crop Protection has estimated aggregate exposure from all crops for which tolerances are established or proposed. The Tier I acute assessment utilized tolerance values and 100% of crop

treated values. The Tier III chronic assessments utilized the residue and percent of crop treated values described above.

a. *Acute exposure*. An acute reference dose of 0.10 mg/kg bwt/day for all population subgroups was based on a NOAEL of 100 mg/kg bwt/day from an acute neurotoxicity study in rats and an uncertainty factor of 100X (100X for combined interspecies and intraspecies variability). An additional Food Quality Protection Act (FQPA) safety factor of 10X was applied to all population subgroups due to the absence of a developmental neurotoxicity study. For the purpose of aggregate risk assessment, the exposure value was expressed in terms of margin of exposure (MOE). The MOE was calculated by dividing the NOAEL by the exposure for each population subgroup. In addition, exposure was expressed as a percent of the acute reference dose (%aRfD). Acute exposure to the most exposed subpopulation (children 1-6 years old) resulted in a MOE of 6,452 (15.5% of the aRfD of 0.10 mg/kg/ bwt/day) at the 95th percentile of exposure. Since the benchmark MOE for this assessment was 1,000, and since EPA generally has no concern for exposures below 100% of the aRfD, Syngenta believes that there is a reasonable certainty that no harm will result from acute dietary (food) exposure to residues arising from the current and proposed uses for thiamethoxam chronic exposure.

b. *Chronic exposure*. The chronic reference dose (cRfD) for thiamethoxam is 0.0006 mg/kg/ bwt/day for all population subgroups and is based on a NOAEL of 0.6 mg/kg/bwt/day from a two generation rat reproduction study. An uncertainty factor of 100X (for combined interspecies and intraspecies variability) and an additional FQPA safety factor of 10X was applied due to evidence of increased susceptibility to young rats following prenatal/postnatal exposure. Exposure was expressed as MOE and percent of the reference dose (%RfD). Chronic exposure to the most exposed subpopulation (non-nursing infants) resulted in a MOE of 11,538 (8.6% of the cRfD of 0.0006 mg/kg/bw/day). Since the benchmark MOE for this assessment was 1,000 and since EPA generally has no concern for exposures below 100% of the RfD, Syngenta believes that there is a reasonable certainty that no harm will result from chronic dietary (food) exposure to residues arising from the current and proposed uses for thiamethoxam.

c. *Lifetime exposure*. The Q* value for thiamethoxam is 0.0377 (mg/kg/day)⁻¹ and is based on benign and malignant

heptocellular tumors in mice in an 18-month carcinogenicity study. Lifetime exposure to the U.S. population resulted in a lifetime risk of 8.17×10^{-7} which represents 81.7% of EPA's lifetime risk limit of 1.0×10^{-6} .

ii. *Drinking water*. EPA used the Pesticide Root Zone/Exposure Analysis Modeling System (PRZM/EXAMS) to estimate pesticide concentrations in surface water and SCI-GROW, which predicts pesticide concentrations in ground water. None of these models include consideration of the impact processing (mixing, dilution, or treatment) of raw water for distribution as drinking water would likely have on the removal of pesticides from the source water. The primary use of these models by the Agency at this stage is to provide a coarse screen for sorting out pesticides for which it is highly unlikely that drinking water concentrations would ever exceed human health levels of concern. Based on the SCI-GROW and PRZM/EXAMS models, EPA calculated that estimated environmental concentrations of thiamethoxam at the highest use rate (0.125 lb a.i./acre) are 1.9 parts per billion (ppb) for acute and chronic exposure to ground water and 11.4 ppb and 0.77 ppb for acute and chronic exposure, respectively, to surface water. Based on field and laboratory data as well as on going prospective ground water monitoring studies, Syngenta believes that the potential exposure to ground water is much lower than that predicted by the conservative SCI-GROW model.

Preliminary results from the prospective ground water monitoring studies have indicated no detections of thiamethoxam in ground water. EPA determined estimated environmental concentrations (EECs) are used for comparison to Drinking Water Levels of Comparison (DWLOC).

a. *Acute drinking water risk*. Acute DWLOCs were calculated based on an acute populated adjusted dose (aPAD) of 0.1 mg/kg/day. For the acute assessment, the children (1-6 yrs) subpopulation generated the lowest acute DWLOC of approximately 845 ppb. EPA has determined that the surface water acute EEC is 11.4 ppb and the ground water EEC is 1.9 ppb. Since the surface water value is greater than the ground water value, the surface water value will be used for comparison purposes and will protect for any concerns for ground water concentrations. Since the acute DWLOC of 845 ppb is considerably higher than the acute EEC of 11.4 ppb, Syngenta believes that EPA should not have a concern for acute risk to either surface or ground water.

b. *Chronic drinking water risk.* Chronic DWLOCs were calculated based on a cPAD of 0.0006 mg/kg/day. For the chronic assessment, the non-nursing infants subpopulation generated the lowest chronic DWLOC of approximately 5.5 ppb. EPA has determined that the surface water chronic EEC is 0.77 ppb and the ground water EEC is 1.9 ppb. Since the ground water value is greater than the surface water value, the ground water value will be used for comparison purposes and will protect for any concerns for surface water concentrations. Since the chronic DWLOC of 5.5 ppb is higher than the chronic EEC of 1.9 ppb, Syngenta believes that EPA should not have a concern for chronic risk to either surface or ground water.

c. *Lifetime drinking water risk.* Based on currently registered and proposed uses for thiamethoxam, Syngenta has determined a DWLOC of 2.0 ppb. At the currently registered maximum use rate of 0.125 lbs. a.i. per acre per growing season, EPA has used the SCI-GROW model to predict a ground water EEC of 1.9 ppb. Thus, the ground water EEC is below the lifetime DWLOC for the general population. The Agency used a screening level model designed to estimate pesticide concentrations in shallow ground water. A number of factors demonstrate that the actual lifetime exposure through drinking water will be less than the lifetime DWLOC. These reasons are as follows:

- Thiamethoxam is a systemic pesticide. EPA's Tier I ground water model assumes that all of the product that is applied to the crop is available for run off. Syngenta has submitted data to show that a percentage (15–25%) of the product is absorbed by the plant, resulting in that much less product available to leach into ground water. Although, data submitted is on only two crops (beans and cucumbers), it is likely that the total amount of thiamethoxam available for ground water leaching is less than the amount EPA uses as a model input.

- Although, the Agency model is based on aerobic soil half lives, EPA's lifetime risk assessment is for lifetime exposure. Data indicate the anaerobic aquatic half-life for thiamethoxam is shorter than the aerobic soil half-life and longer than the aerobic aquatic half-life. Although, EPA is unable to predict, with a high degree of certainty, what happens to thiamethoxam ground water over time, this does provide some support for the expectation that concentrations in ground water will decline between annual applications.

- Shallow ground water modeling is not the perfect model for representing

all drinking water from ground water sources. It is likely to be an over estimate of most drinking water concentrations, which tend to originate from deeper sources. EPA's experience is that the model is reasonably accurate for shallow drinking water, but it is less accurate for estimating concentrations in drinking water from deeper sources.

- The Agency has established conditions of registration for the previous uses that include two prospective ground water studies and a retrospective monitoring study, so that the reasonable certainty of no harm finding will be sustained. Preliminary results have indicated no detections of thiamethoxam in ground water.

- The dietary food risk is based on residue data derived from the average of field trials, which were performed at a higher application rate than what was accepted by EPA. It is not unusual in the Agency's experience for field trial data to be an order of magnitude above actual monitoring. Since thiamethoxam has only recently been registered, actual monitoring data are not yet available. It is likely that the actual risk contribution from food will be much lower than current data indicate, which would result in a larger lifetime DWLOC. Syngenta expects that this refined lifetime DWLOC would be larger than the EECs for the proposed uses. Based on the previous points, Syngenta does not expect that the general population would be exposed to levels exceeding the lifetime DWLOC.

2. *Non-dietary exposure.*

Thiamethoxam is not currently registered for use on any sites that would result in residential exposure.

D. *Cumulative Effects*

The potential for cumulative effects of thiamethoxam and other substances that have a common mechanism of toxicity has also been considered.

Thiamethoxam belongs to a new pesticide chemical class known as the neonicotinoids. There is no reliable information to indicate that toxic effects produced by thiamethoxam would be cumulative with those of any other chemical including another pesticide. Therefore, Syngenta believes it is appropriate to consider only the potential risks of thiamethoxam in an aggregate risk assessment.

E. *Safety Determination*

Syngenta concludes, as described above, that there is reasonable certainty that no harm to the U.S. population will result from aggregate acute or chronic dietary exposure to thiamethoxam residues including the proposed commodities.

F. *International Tolerances*

There are no codex MRLs established for residues of thiamethoxam.

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ENVIRONMENTAL PROTECTION AGENCY

[OPP–2003–0102; FRL–7299–6]

Fludioxonil; Notice of Filing Pesticide Petitions to Establish a Tolerance for a Certain Pesticide Chemical in or on Food

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice.

SUMMARY: This notice announces the initial filing of a pesticide petition proposing the establishment of regulations for residues of a certain pesticide chemical in or on various food commodities.

DATES: Comments, identified by docket ID number OPP–2003–0102, must be received on or before May 2, 2003.

ADDRESSES: Comments may be submitted electronically, by mail, or through hand delivery/courier. Follow the detailed instructions as provided in Unit I.C. of the **SUPPLEMENTARY INFORMATION**.

FOR FURTHER INFORMATION CONTACT: Sidney Jackson, Registration Division (7505C), Office of Pesticide Programs, Environmental Protection Agency, 1200 Pennsylvania Ave., NW., Washington, DC 20460–0001; telephone number: (703) 305–7610; e-mail address: jackson.sidney@epa.gov.

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. 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. Potentially affected entities may include, but are not limited to:

- Crop production (NAICS 111)
- Animal production (NAICS 112)
- Food manufacturing (NAICS 311)
- Pesticide manufacturing (NAICS 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