During any full calendar month you are in the medical care facility, you cannot receive more than the Federal benefit rate described in § 416.414(b)(1). We do not consider food or shelter provided during a medical confinement to be income.

(ii) If you enter a medical care facility and you are eligible for either benefit payable under § 416.212, we also consider this a temporary absence from your permanent living arrangement. We use the rules that apply to your permanent living arrangement to value any food, clothing, or shelter you receive during the month you enter the facility and throughout the period you are eligible for these benefits. We consider your absence to be temporary through the last month benefits under § 416.212 are paid unless you are discharged from the facility in the following month. In that case, we consider your absence to be temporary through the date of discharge.

* * * * *

15. Section 416.1167 is amended by revising paragraph (a) to read as follows:

§ 416.1167 Temporary absences and deeming rules.

(a) *General.* During a temporary absence, we continue to consider the absent person a member of the household. A temporary absence occurs when—

(1) You, your ineligible spouse, parent, or an ineligible child leaves the household but intends to and does return in the same month or the month immediately following; or

(2) You enter a medical care facility and are eligible for either benefit payable under § 416.212. We consider your absence to be temporary through the last month benefits under § 416.212 were paid unless you were discharged from the facility in the following month. In that case, we consider your absence to be temporary through the date of discharge.

* * * * *

Subpart T—State Supplementation Provisions; Agreement; Payments

16. The authority citation for subpart T of part 416 continues to read as follows:

Authority: Secs. 702(a)(5), 1616, 1618, and 1631 of the Social Security Act (42 U.S.C. 902(a)(5), 1382e, 1382g, and 1383); sec. 212, Pub. L. 93–66, 87 Stat. 155 (42 U.S.C. 1382 note); sec. 8 (a), (b)(1)–(b)(3), Pub. L. 93–233, 87 Stat. 956 (7 U.S.C. 612c note, 1431 note and 42 U.S.C. 1382e note); secs. 1 (a)–(c) and 2(a), 2(b)(1), 2(b)(2), Pub. L. 93–335, 88 Stat. 291 (42 U.S.C. 1382 note, 1382e note).

17. Section 416.2040 is amended by revising paragraph (a) and adding a new paragraph (c) to read as follows:

§ 416.2040 Limitations on eligibility.

* * * * *

(a) *Inmate of public institution.* A person who is a resident in a public institution for a month, is ineligible for a Federal benefit for that month under the provision of § 416.211(a), and does not meet the requirements for any of the exceptions in § 416.211 (b), (c), or (d), or § 416.212, also shall be ineligible for a federally administered State supplementary payment for that month.

* * * * *

(c) *Recipient eligible for benefits under § 416.212.* A recipient who is institutionalized and is eligible for either benefit payable under § 416.212 for a month or months may also receive federally administered State supplementation for that month. Additionally, a recipient who would be eligible for benefits under § 416.212 but for countable income which reduces his or her Federal SSI benefit to zero, may still be eligible to receive federally administered State supplementation.

[FR Doc. 96–5705 Filed 3–12–96; 8:45 am]

BILLING CODE 4190–29–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 101

[Docket No. 90N–0134]

RIN 0910–AA19

Food Labeling: Reference Daily Intakes; Correction

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule; correction.

SUMMARY: The Food and Drug Administration (FDA) is correcting a final rule that appeared in the Federal Register of December 28, 1995 (60 FR 67164). The final rule amended FDA regulations to establish Reference Daily Intakes (RDI's) for vitamin K, selenium, manganese, chromium, molybdenum, and chloride, but not for fluoride. The document was published with some typographical errors. This document corrects those errors.

EFFECTIVE DATE: January 1, 1997.

FOR FURTHER INFORMATION CONTACT: Camille E. Brewer, Center for Food Safety and Applied Nutrition (HFS–165), Food and Drug Administration, 200 C St. SW., Washington, DC 20204, 202–205–5483.

In FR Doc. 95–31197, appearing on page 67164 in the Federal Register of Thursday, December 28, 1995, the following corrections are made:

1. On page 67167, in the second column, in lines three, five, seven, and eight, ''mg'' is corrected to read ''μg.''

§ 101.36 Corrected

2. On page 67175, in the second column, in § 101.36(b)(3)(ii), in line fourteen, ''vitamin B6'' is corrected to read ''vitamin B_6'', and ''vitamin B12'' is corrected to read ''vitamin B_{12}''.

Dated: March 7, 1996.

William K. Hubbard,

Associate Commissioner for Policy Coordination.

[FR Doc. 96–6029 Filed 3–12–96; 8:45 am]

BILLING CODE 4160–01–F

ENVIRONMENTAL PROTECTION AGENCY

40 CFR Part 180

[PP 0E3889, 2E4113, and 5E4538/R2210; FRL–5352–8]

RIN 2070–AC78

Chlorothalonil; Pesticide Tolerances

AGENCY: Environmental Protection Agency (EPA).

ACTION: Final Rule.

SUMMARY: This document establishes tolerances for combined residues of the fungicide chlorothalonil and it metabolite in or on the raw agricultural commodities blueberries, filberts, and mushrooms. The Interregional Research Project No. 4 (IR-4) requested the regulation to establish a maximum permissible level for residues of the fungicide pursuant to the Federal Food, Drug and Cosmetic Act (FFDCA).

EFFECTIVE DATE: This regulation becomes effective March 13, 1996.

ADDRESSES: Written objections and hearing requests, identified by the document control number, [PP 0E3889, 2E4113, and 5E4538/R2210], may be submitted to: Hearing Clerk (1900), Environmental Protection Agency, Rm. M3708, 401 M St., SW., Washington, DC 20460. Fees accompanying objections and hearing requests shall be labeled ''Tolerance Petition Fees'' and forwarded to: EPA Headquarters Accounting Operations Branch, OPP (Tolerance Fees), P.O. Box 360277M, Pittsburgh, PA 15251. A copy of any objections and hearing requests filed with the Hearing Clerk should be

identified by the document control number and submitted to: Public Response and Program Resources Branch, Field Operations Division (7506C), Office of Pesticide Programs, Environmental Protection Agency, 401 M St., SW., Washington, DC 20460. In person, bring copy of objections and hearing requests to Rm. 1132, CM #2, 1921 Jefferson Davis Hwy., Arlington, VA 22202.

A copy of objections and hearing requests filed with the Hearing Clerk may also be submitted electronically by sending electronic mail (e-mail) to: opp-docket@epamail.epa.gov. Copies of objections and hearing requests must be submitted as an ASCII file avoiding the use of special characters and any form of encryption. Copies of objections and hearing requests will also be accepted on disks in WordPerfect 5.1 file format or ASCII file format. All copies of objections and hearing requests in electronic form must be identified by the docket number [PP 0E3889, 2E4113, and 5E4538/R2210]. No Confidential Business Information (CBI) should be submitted through e-mail. Electronic copies of objections and hearing requests on this rule may be filed online at many Federal Depository Libraries. Additional information on electronic submissions can be found below in this document.

FOR FURTHER INFORMATION CONTACT: By mail: Hoyt L. Jamerson, Registration Division (7505W), Environmental Protection Agency, 401 M St., SW., Washington, DC 20460. Office location and telephone number: Sixth Floor, Crystal Station #1, 2800 Jefferson Davis Highway, Arlington, VA 22202. (703) 308-8783, e-mail: jamerson.hoyt@epamail.epa.gov.

SUPPLEMENTARY INFORMATION: In the Federal Register of January 24, 1996 (61 FR 1884), EPA issued a proposed rule that gave notice that the Interregional Research Project No. 4 (IR-4), New Jersey Agricultural Experiment Station, P.O. Box 231, New Brunswick, NJ 08903, had submitted pesticide petitions (PP) 0E3889, 2E4113, and 5E4538 to EPA on behalf of the named Agricultural Experiment Stations. These petitions requested that the Administrator, pursuant to section 408(e) of the Federal Food, Drug and Cosmetic Act (FFDCA), 21 U.S.C. 346a(e) amend 40 CFR 180.275 by establishing tolerances for combined residues of the fungicide chlorothalonil (tetrachloroisophthalonitrile) and its metabolite 4-hydroxy-2,5,6-trichloroisophthalonitrile in or on certain raw agricultural commodities, as follows:

1. *PP 0E3889.* Petition submitted on behalf of the Agricultural Experiment Stations of Florida, Georgia, Kentucky, Louisiana, Michigan, North Carolina, Oklahoma, Pennsylvania, South Carolina, Tennessee, Texas, and Washington proposing a tolerance for blueberries at 1.0 part per million (ppm).

2. *PP 2E4113.* Petition submitted on behalf of the Oregon Agricultural Experiment Station proposing a tolerance for filberts at 0.1 ppm. The petitioner proposed that use of chlorothalonil on filberts be limited to Oregon based on the geographical representation of the residue data submitted. Additional residue data will be required to expand the area of usage. Persons seeking geographically broader registration should contact the Agency's Registration Division at the address provided above.

3. *PP 5E4538.* Petition submitted on behalf of the Pennsylvania Agricultural Experiment Station proposing a tolerance for mushrooms at 1.0 ppm.

There were no comments or requests for referral to an advisory committee received in response to the proposed rule.

The data submitted with the proposals and other relevant material have been evaluated and discussed in the proposed rule. Based on the data and information considered, the Agency concludes that the tolerances will protect the public health. Therefore, the tolerances are established as set forth below.

Any person adversely affected by this regulation may, within 30 days after publication of this document in the Federal Register, file written objections to the regulation and may also request a hearing on those objections. Objections and hearing requests must be filed with the Hearing Clerk, at the address given above (40 CFR 178.20). A copy of the objections and/or hearing requests filed with the Hearing Clerk should be submitted to the OPP docket for this rulemaking. The objections submitted must specify the provisions of the regulation deemed objectionable and the grounds for the objections (40 CFR 178.25). Each objection must be accompanied by the fee prescribed by 40 CFR 180.33(i). If a hearing is requested, the objections must include a statement of the factual issue(s) on which a hearing is requested, the requestor's contentions on such issues, and a summary of any evidence relied upon by the objector (40 CFR 178.27). A request for a hearing will be granted if the Administrator determines that the material submitted shows the following: There is genuine and substantial issue of fact; there is a reasonable possibility that available evidence identified by the requestor would, if established, resolve one or more of such issues in favor of the requestor, taking into account uncontested claims or facts to the contrary; and resolution of the factual issue(s) in the manner sought by the requestor would be adequate to justify the action requested (40 CFR 178.32).

A record has been established for this rulemaking under docket number [PP0E3889, 2E4113, and 5E4538/R2210] (including any objections and hearing requests submitted electronically as described below). A public version of this record, including printed, paper versions of electronic comments, which does not include any information claimed as CBI, is available for inspection from 8 a.m. to 4:30 p.m., Monday through Friday, excluding legal holidays. The public record is located in Room 1132 of the Public Response and Program Resources Branch, Field Operations Division (7506C), Office of Pesticide Programs, Environmental Protection Agency, Crystal Mall #2, 1921 Jefferson Davis Highway, Arlington, VA.

Written objections and hearing requests, identified by the document number [PP 0E3889, 2E4113, and 5E4538/R2210], may be submitted to the Hearing Clerk (1900), Environmental Protection Agency, Rm. 3708, 401 M St., SW., Washington, DC 20460.

A copy of electronic objections and hearing requests filed with the Hearing Clerk can be sent directly to EPA at:

opp-Docket@epamail.epa.gov

A copy of electronic objections and hearing requests filed with the Hearing Clerk must be submitted as an ASCII file avoiding the use of special characters and any form of encryption.

The official record for this rulemaking, as well as the public version, as described above will be kept in paper form. Accordingly, EPA will transfer any copies of objections and hearing requests received electronically into printed, paper form as they are received and will place the paper copies in the official rulemaking record which will also include all comments submitted directly in writing. The official rulemaking record is the paper record maintained at the address in ''ADDRESSES'' at the beginning of this document.

Under Executive Order 12866 (58 FR 51735, October 4, 1993), the Agency must determine whether the regulatory action is ''significant'' and therefore subject to review by the Office of Management and Budget (OMB) and the requirements of the Executive Order. Under section 3(f), the order defines ''a

significant regulatory action'' as an action that is likely to result in a rule (1) having an annual effect on the economy of $100 million or more, or adversely and materially affecting a sector of the economy, productivity, competition, jobs, the environment, public health or safety, or State, local or tribal governments or communities (also referred to as ''economically significant''); (2) creating serious inconsistency or otherwise interfering with an action taken or planned by another agency; (3) materially altering the budgetary impacts of entitlement, grants, user fees, or loan programs or the rights and obligations thereof; or (4) raising novel legal or policy issues arising out of legal mandates, the President's priorities, or the principles set forth in this Executive Order.

Pursuant to the terms of this Executive Order, EPA has determined that this rule is not ''significant'' and is therefore not subject to OMB review.

Pursuant to the requirements of the Regulatory Flexibility Act (Pub. L. 96-354, 94 Stat. 1164, 5 U.S.C. 601-612), the Administrator has determined that regulations establishing new tolerances or raising tolerance levels or establishing exemptions from tolerance requirements do not have a significant economic impact on a substantial number of small entities. A certification statement to this effect was published in the Federal Register of May 4, 1981 (46 FR 24950).

List of Subjects in 40 CFR Part 180

Environmental protection, Administrative practice and procedure, Agricultural commodities, Pesticides and pests, Reporting and recordkeeping requirements.

Dated: February 27, 1996.

Stephen L. Johnson,

Director, Registration Division, Office of Pesticide Programs.

Therefore, 40 CFR part 180 is amended as follows:

PART 180—[AMENDED]

1. The authority citation for Part 180 continues to read as follows:

Authority: 21 U.S.C. 346a and 371.

2. In § 180.275, by amending the table in paragraph (a) by adding alphabetically the raw agricultural commodities blueberries and mushrooms and by amending the table in paragraph (b), by adding alphabetically the raw agricultural commodity filberts to read as follows:

§ 180.275 Chlorothalonil; tolerances for residues.

(a) * * *

Commodities	Parts per million
* * * *	*
Blueberries	1.0
* * * *	*
Mushrooms	1.0
* * * *	*

(b) * * *

Commodities	Parts per million
* * * *	*
Filberts	0.1
* * * *	*

[FR Doc. 96–5536 Filed 3–12–96; 8:45 am]

BILLING CODE 6560–50–F

40 CFR Part 180

[OPP–300402A; FRL–4993–3]

RIN 2070–AB78

3,5-Dichloro-N-(1,1-Dimethyl-2-Propynyl)Benzamide; Pesticide Tolerances

AGENCY: Environmental Protection Agency (EPA).

ACTION: Final rule.

SUMMARY: EPA has completed the reregistration process and issued a Reregistration Eligibility Decision (RED) document for the pesticide 3,5-dichloro-*N*-(1,1-dimethyl-2-propynyl)benzamide, also known as pronamide. In the reregistration process, all information to support a pesticide's continued registration is reviewed for adequacy and, when needed, supplemented with new scientific studies. Based on the RED tolerance assessments for the pesticide chemical subject to this rule, EPA is issuing the following tolerance actions: to delete individual tolerances and establish crop-grouping tolerances, raise some tolerances and lower others, amend an incorrectly listed tolerance, and modify the statment under 40 CFR 180.317 for the pesticide pronamide.

EFFECTIVE DATE: This regulation becomes effective March 13, 1996.

ADDRESSES: Written objections and hearing requests, identified by the document control number, [OPP-300402A], may be submitted to: Hearing Clerk (1900), Environmental Protection Agency, Rm. M3708, 401 M St., SW., Washington, DC 20460. A copy of any objections and hearing requests filed with the Hearing Clerk should be identified by the document control number and submitted to: Public Response and Program Resources Branch, Field Operations Division (7506C), Office of Pesticide Programs, Environmental Protection Agency, 401 M St., SW., Washington, DC 20460. In person, bring copy of objections and hearing requests to Rm. 1132, CM #2, 1921 Jefferson Davis Hwy., Arlington, VA 22202. Fees accompanying objections shall be labeled ''Tolerance Petition Fees'' and forwarded to: EPA Headquarters Accounting Operations Branch, OPP (Tolerance Fees), P.O. Box 360277M, Pittsburgh, PA 15251. An electronic copy of objections and hearing requests filed with the Hearing Clerk may be submitted to OPP by sending electronic mail (e-mail) to:opp-docket@epamail.epa.gov.

Copies of electronic objections and hearing requests must be submitted as an ASCII file avoiding the use of special characters and any form of encryption. Copies of electronic objections and hearing requests will also be accepted on disks in WordPerfect 5.1 file format or ASCII file format. All copies of electronic objections and hearing requests must be identified by the docket number [OPP-300402A]. No Confidential Business Information (CBI) should be submitted through e-mail. Copies of electronic objections and hearing requests on this rule may be filed online at many Federal Depository Libraries. Additional information on electronic submissions can be found below in this document.

FOR FURTHER INFORMATION CONTACT: By mail: Philip Poli, (703)-308-8038; e-mail: poli.philip@epamail.epa.gov. By mail: Special Review and Reregistration Division (7508W), Environmental Protection Agency, 401 M St., SW., Washington, DC 20460. Office location: Special Review Branch, Crystal Station #1, 3rd Floor, 2800 Crystal Drive, Arlington, VA 22202.

SUPPLEMENTARY INFORMATION: EPA issued a proposed rule, published in the Federal Register of November 15, 1995 (60 FR 57379), which announced that based on a Reregistration Eligibility Decision (RED) for the pesticide 3,5-dichloro-*N*-(1,1-dimethyl-2-propynyl)benzamide, also known as pronamide, the Agency intended to revise 40 CFR 180.317 to delete individual tolerances and establish crop-grouping tolerances (as described in 40 CFR 180.34), raise some tolerances