& Walk events. This deviation allows the bridge to remain in the closed position to allow for the safe movement of event participants.

DATES: This deviation is effective from 7:30 a.m. on April 7, 2013, to 8:30 a.m. on April 14, 2013.

ADDRESSES: The docket for this deviation, [USCG–2013–0123] is available at *http://www.regulations.gov*. Type the docket number in the ''SEARCH'' box and click ''SEARCH.'' Click on Open Docket Folder on the line associated with this deviation. You may also visit the Docket Management Facility in Room W12–140 on the ground floor of the Department of Transportation West Building, 1200 New Jersey Avenue SE., Washington, DC 20590, between 9 a.m. and 5 p.m., Monday through Friday, except Federal holidays.

FOR FURTHER INFORMATION CONTACT: If you have questions on this temporary deviation, call or email Randall Overton, Bridge Administrator, Coast Guard Thirteenth District; telephone 206–220–7282, email *Randall.D.Overton@uscg.mil*. If you have questions on viewing the docket, call Barbara Hairston, Program Manager, Docket Operations, telephone 202–366–9826.

SUPPLEMENTARY INFORMATION: Multnomah County has requested that the Broadway Bascule Bridge remain closed to vessel traffic to facilitate safe, uninterrupted roadway passage of participants in the Race for the Roses and the Bridge to Brews Run & Walk events. The Broadway Bridge crosses the Willamette River at mile 11.7 and provides 90 feet of vertical clearance above Columbia River Datum 0.0 while in the closed position. Vessels that do not require a bridge opening may continue to transit beneath the bridge during the closure periods. Under normal conditions this bridge operates in accordance with 33 CFR 117.897, which allows for the bridge to remain closed between 7 a.m. and 9 a.m. and 4 p.m. and 6 p.m. Monday through Friday, and also requires advance notification when a bridge opening is needed. This deviation period is effective from 7:30 a.m. on April 7, 2013, to 8:30 a.m. on April 14, 2013. The deviation allows the bascule span of the Broadway Bridge across the Willamette River, mile 11.7, to remain in the closed position and need not open for maritime traffic from 7:30 a.m. until 12:01 p.m. on April 7, 2013, and from 5:30 a.m. until 8:30 a.m. on April 14, 2013. The bridge shall operate in accordance to 33 CFR 117.897 at all other times. Waterway usage on this stretch of the Willamette River includes vessels ranging from commercial tug and barge to small pleasure craft. Mariners will be notified and kept informed of the bridge's operational status via the Coast Guard Notice to Mariners publication and Broadcast Notice to Mariners as appropriate. The draw span will be required to open, if needed, for vessels engaged in emergency response operations during this closure period.

In accordance with 33 CFR 117.35(e), the drawbridge must return to its regular operating schedule immediately at the end of the designated time period. This deviation from the operating regulations is authorized under 33 CFR 117.35.

Dated: February 27, 2013.

Randall D. Overton,

Bridge Administrator.

[FR Doc. 2013–05711 Filed 3–12–13; 8:45 am]

BILLING CODE 9110–04–P

ENVIRONMENTAL PROTECTION AGENCY

40 CFR Part 180

[EPA–HQ–OPP–2011–0360; FRL–9380–9]

RIN 2070–ZA16

Tetrachlorvinphos; Extension of Time-Limited Interim Pesticide Tolerances

AGENCY: Environmental Protection Agency (EPA).

ACTION: Final rule.

SUMMARY: This regulation extends the time-limited interim tolerances for the combined residues of the insecticide tetrachlorvinphos, in or on multiple commodities which are identified in Unit III. of this document, under the Federal Food, Drug, and Cosmetic Act (FFDCA).

DATES: This regulation is effective March 13, 2013. Objections and requests for hearings must be received on or before May 13, 2013, and must be filed in accordance with the instructions provided in 40 CFR part 178 (see also Unit I.C. of the **SUPPLEMENTARY INFORMATION**).

ADDRESSES: The docket for this action, identified by docket identification (ID) number EPA–HQ–OPP–2011–0360, is available at *http://www.regulations.gov* or at the Office of Pesticide Programs Regulatory Public Docket (OPP Docket) in the Environmental Protection Agency Docket Center (EPA/DC), EPA West Bldg., Rm. 3334, 1301 Constitution Ave. NW., Washington, DC 20460–0001. The 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 OPP Docket is (703) 305–5805. Please review the visitor instructions and additional information about the docket available at *http://www.epa.gov/dockets.*

FOR FURTHER INFORMATION CONTACT: Carmen Rodia, Registration Division (7504P), Office of Pesticide Programs, Environmental Protection Agency, 1200 Pennsylvania Avenue NW., Washington, DC 20460–0001; telephone number: (703) 306–0327; email address: *rodia.carmen@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 following list of North American Industrial Classification System (NAICS) codes is not intended to be exhaustive, but rather provides a guide to help readers determine whether this document applies to them. Potentially affected entities may include:

- Crop production (NAICS code 111).
- Animal production (NAICS code 112).
- Food manufacturing (NAICS code 311).
- Pesticide manufacturing (NAICS code 32532).

B. How can I get electronic access to other related information?

You may access a frequently updated electronic version of 40 CFR part 180 through the Government Printing Office's e-CFR site at *http://www.ecfr.gov/cgi-bin/text-idx?&c=ecfr&tpl=/ecfrbrowse/Title40/40tab_02.tpl.*

C. How can I file an objection or hearing request?

Under FFDCA section 408(g), 21 U.S.C. 346a, any person may file an objection to any aspect of this regulation and may also request a hearing on those objections. You must file your objection or request a hearing on this regulation in accordance with the instructions provided in 40 CFR part 178. To ensure proper receipt by EPA, you must identify docket ID number EPA–HQ–OPP–2011–0360 in the subject line on the first page of your submission. All objections and requests for a hearing must be in writing, and must be received by the Hearing Clerk on or before May 13, 2013. Addresses for mail and hand delivery of objections and

hearing requests are provided in 40 CFR 178.25(b).

In addition to filing an objection or hearing request with the Hearing Clerk as described in 40 CFR part 178, please submit a copy of the filing (excluding any Confidential Business Information (CBI)) for inclusion in the public docket. Information not marked confidential pursuant to 40 CFR part 2 may be disclosed publicly by EPA without prior notice. Submit the non-CBI copy of your objection or hearing request, identified by docket ID number EPA–HQ–OPP–2011–0360, by one of the following methods:

- *Federal eRulemaking Portal: http://www.regulations.gov.* Follow the online instructions for submitting comments. Do not submit electronically any information you consider to be CBI or other information whose disclosure is restricted by statute.
- *Mail:* OPP Docket, Environmental Protection Agency Docket Center (EPA/DC), (28221T), 1200 Pennsylvania Ave. NW., Washington, DC 20460–0001.
- *Hand Delivery:* To make special arrangements for hand delivery or delivery of boxed information, please follow the instructions at *http://www.epa.gov/dockets/contacts.htm.* Additional instructions on commenting or visiting the docket, along with more information about dockets generally, is available at *http://www.epa.gov/dockets.*

II. Background

A detailed summary of the background related to EPA's extension of the time-limited interim tolerances for the combined residues of the insecticide tetrachlorvinphos, including its metabolites, in or on multiple commodities can be found in the **Federal Register** notices of June 8, 2011 (76 FR 33184) (FRL–8874–7); September 16, 2011 (76 FR 57657) (FRL–8887–5); and March 6, 2013 (78 FR 14487) (FRL–9380–8). The referenced documents are available in the docket established by this action, which is described under **ADDRESSES**. Locate and click on the hyperlink for docket ID number EPA–HQ–OPP–2011–0360. Double-click on the documents to view the referenced background summary information. There were no substantive comments received in response to the proposed rule of March 6, 2013.

III. Conclusion

For the reasons stated in the proposed rule of March 6, 2013, EPA is finalizing its proposal to extend to August 18, 2013, the expiration dates for the following tolerances listed in 40 CFR 180.252 for tetrachlorvinphos, (Z)-2-chloro-1-(2,4,5-trichlorophenyl) vinyl dimethyl phosphate, including its metabolites, 1-(2,4,5-trichlorophenyl)-ethanol (free and conjugated forms), 2,4,5-trichloroacetophenone, and 1-(2,4,5-trichlorophenyl)-ethanediol in or on: Cattle, fat (of which no more than 0.1 part per million (ppm) is tetrachlorvinphos *per se*) at 0.2 (ppm); cattle, kidney (of which no more than 0.05 ppm is tetrachlorvinphos *per se*) at 1.0 ppm; cattle, liver (of which no more than 0.05 ppm is tetrachlorvinphos *per se*) at 0.5 ppm; cattle, meat (of which no more than 2.0 ppm is tetrachlorvinphos *per se*) at 2.0 ppm; cattle, meat byproducts, except kidney and liver at 1.0 ppm; egg (of which no more than 0.05 ppm is tetrachlorvinphos *per se*) at 0.2 ppm; hog, fat (of which no more than 0.1 ppm is tetrachlorvinphos *per se*) at 0.2 ppm; hog, kidney (of which no more than 0.05 ppm is tetrachlorvinphos *per se*) at 1.0 ppm; hog, liver (of which no more than 0.05 ppm is tetrachlorvinphos *per se*) at 0.5 ppm; hog, meat (of which no more than 2.0 ppm is tetrachlorvinphos *per se*) at 2.0 ppm; hog, meat byproducts, except kidney and liver at 1.0 ppm; milk, fat (reflecting negligible residues in whole milk and of which no more than 0.05 ppm is tetrachlorvinphos *per se*) at 0.05 ppm; poultry, fat (of which no more than 7.0 ppm is tetrachlorvinphos *per se*) at 7.0 ppm; poultry, liver (of which no more than 0.05 ppm is tetrachlorvinphos *per se*) at 2.0 ppm; poultry, meat (of which no more than 3.0 ppm is tetrachlorvinphos *per se*) at 3.0 ppm; and poultry, meat byproducts, except liver at 2.0 ppm.

IV. Statutory and Executive Order Reviews

This final rule establishes tolerances under FFDCA section 408(e) of FFDCA. The Office of Management and Budget (OMB) has exempted these types of actions from review under Executive Order 12866, entitled ''Regulatory Planning and Review'' (58 FR 51735, October 4, 1993). Because this final rule has been exempted from review under Executive Order 12866, this final rule is not subject to Executive Order 13211, entitled ''Actions Concerning Regulations That Significantly Affect Energy Supply, Distribution, or Use'' (66 FR 28355, May 22, 2001) or Executive Order 13045, entitled ''Protection of Children from Environmental Health Risks and Safety Risks'' (62 FR 19885, April 23, 1997). This final rule does not contain any information collections subject to OMB approval under the Paperwork Reduction Act (PRA) (44 U.S.C. 3501 *et seq.*), nor does it require any special considerations under Executive Order 12898, entitled ''Federal Actions to Address Environmental Justice in Minority Populations and Low-Income Populations'' (59 FR 7629, February 16, 1994).

In addition, under the Regulatory Flexibility Act (RFA) (5 U.S.C. 601 et seq.), the Agency previously assessed whether establishing tolerances, exemptions from tolerances, raising tolerance levels, or expanding exemptions might adversely impact small entities and concluded, as a generic matter, that there is no adverse economic impact. The factual basis for the Agency's generic certification for tolerance actions was published on May 4, 1981 (46 FR 24950), and was provided to the Chief Counsel for Advocacy of the Small Business Administration.

This final rule directly regulates growers, food processors, food handlers, and food retailers, not States or tribes, nor does this action alter the relationships or distribution of power and responsibilities established by Congress in the preemption provisions of FFDCA section 408(n)(4). As such, the Agency has determined that this action will not have a substantial direct effect on States or tribal governments, on the relationship between the national government and the States or tribal governments, or on the distribution of power and responsibilities among the various levels of government or between the Federal Government and Indian tribes. Thus, the Agency has determined that Executive Order 13132, entitled ''Federalism'' (64 FR 43255, August 10, 1999) and Executive Order 13175, entitled ''Consultation and Coordination with Indian Tribal Governments'' (65 FR 67249, November 9, 2000) do not apply to this final rule. In addition, this final rule does not impose any enforceable duty or contain any unfunded mandate as described under Title II of the Unfunded Mandates Reform Act of 1995 (UMRA) (2 U.S.C. 1501 *et seq.*).

This action does not involve any technical standards that would require Agency consideration of voluntary consensus standards pursuant to section 12(d) of the National Technology Transfer and Advancement Act of 1995 (NTTAA) (15 U.S.C. 272 note).

V. Congressional Review Act

Pursuant to the Congressional Review Act (5 U.S.C. 801 *et seq.*), EPA will submit a report containing this rule and other required information to the U.S. Senate, the U.S. House of Representatives, and the Comptroller General of the United States prior to publication of the rule in the **Federal**

Register. This action is not a ''major rule'' as defined by 5 U.S.C. 804(2).

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 26, 2013.

G. Jeffrey Herndon,

Acting Director, Registration Division, Office of Pesticide Programs.

Therefore, 40 CFR chapter I is amended as follows:

PART 180—[AMENDED]

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

Authority: 21 U.S.C. 321(q), 346a and 371.

■ 2. In § 180.252 revise the table in paragraph (a) to read as follows:

§ 180.252 Tetrachlorvinphos; tolerances for residues.

(a) * * *

Commodity	Parts per million	Expiration/revocation date
Cattle, fat (of which no more than 0.1 ppm is tetrachlorvinphos *per se*)	0.2	8/18/13
Cattle, kidney (of which no more than 0.05 ppm is tetrachlorvinphos *per se*)	1.0	8/18/13
Cattle, liver (of which no more than 0.05 ppm is tetrachlorvinphos *per se*)	0.5	8/18/13
Cattle, meat (of which no more than 2.0 ppm is tetrachlorvinphos *per se*)	2.0	8/18/13
Cattle, meat byproducts, except kidney and liver	1.0	8/18/13
Egg (of which no more than 0.05 ppm is tetrachlorvinphos *per se*)	0.2	8/18/13
Hog, fat (of which no more than 0.1 ppm is tetrachlorvinphos *per se*)	0.2	8/18/13
Hog, kidney (of which no more than 0.05 ppm is tetrachlorvinphos *per se*)	1.0	8/18/13
Hog, liver (of which no more than 0.05 ppm is tetrachlorvinphos *per se*)	0.5	8/18/13
Hog, meat (of which no more than 2.0 ppm is tetrachlorvinphos *per se*)	2.0	8/18/13
Hog, meat byproducts, except kidney and liver	1.0	8/18/13
Milk, fat (reflecting negligible residues in whole milk and of which no more than 0.05 ppm is tetrachlorvinphos *per se*)	0.05	8/18/13
Poultry, fat (of which no more than 7.0 ppm is tetrachlorvinphos *per se*)	7.0	8/18/13
Poultry, liver (of which no more than 0.05 ppm is tetrachlorvinphos *per se*)	2.0	8/18/13
Poultry, meat (of which no more than 3.0 ppm is tetrachlorvinphos *per se*)	3.0	8/18/13
Poultry, meat byproducts, except liver	2.0	8/18/13

* * * * *

[FR Doc. 2013–05814 Filed 3–12–13; 8:45 am]

BILLING CODE 6560–50–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

42 CFR Parts 412, 413, 424, and 476

[CMS–1588–CN4]

RIN 0938–AR12

Medicare Program; Hospital Inpatient Prospective Payment Systems for Acute Care Hospitals and the Long Term Care Hospital Prospective Payment System and Fiscal Year 2013 Rates; Hospitals' Resident Caps for Graduate Medical Education Payment Purposes; Quality Reporting Requirements for Specific Providers and for Ambulatory Surgical Centers; Corrections

AGENCY: Centers for Medicare & Medicaid Services (CMS), HHS.

ACTION: Final rule; correction.

SUMMARY: This document corrects technical errors in the correcting document that appeared in the October 3, 2012 **Federal Register** entitled ''Medicare Program; Hospital Inpatient Prospective Payment Systems for Acute Care Hospitals and the Long Term Care Hospital Prospective Payment System and Fiscal Year 2013 Rates; Hospitals' Resident Caps for Graduate Medical Education Payment Purposes; Quality Reporting Requirements for Specific Providers and for Ambulatory Surgical Centers; Correction.''

DATES: *Effective date:* This correcting document is effective March 12, 2013.

Applicability Date: This correcting document is applicable to discharges on or after October 1, 2012.

FOR FURTHER INFORMATION CONTACT: Tzvi Hefter, (410) 786–4487.

SUPPLEMENTARY INFORMATION:

I. Background

In the August 31, 2012 **Federal Register** (77 FR 53258), we published a final rule entitled ''Medicare Program; Hospital Inpatient Prospective Payment Systems for Acute Care Hospitals and the Long Term Care Hospital Prospective Payment System and Fiscal Year 2013 Rates; Hospitals' Resident Caps for Graduate Medical Education Payment Purposes; Quality Reporting Requirements for Specific Providers and for Ambulatory Surgical Centers'' (hereinafter referred to as the FY 2013 IPPS/LTCH PPS final rule). To correct typographical and technical errors in the FY 2013 IPPS/LTCH PPS final rule, we published correcting documents that appeared in the October 3, 2012 **Federal Register** (77 FR 60315); October 17, 2012 **Federal Register** (77 FR 63751); and the October 29, 2012 **Federal Register** (77 FR 65495).

The October 3, 2012 correcting document (77 FR 60315) included several corrections to figures and data for the Hospital Readmissions Reduction program. Since that time, we have determined that these corrections still contained errors. Therefore, in this correcting document, we will identify and correct the errors related to the Hospital Readmissions Reduction Program included in October 3, 2012 correcting document (FR Doc. 2012–24307).

II. Summary of Errors and Corrections to Tables Posted on the CMS Web Site

A. Errors in the October 3, 2012 Correcting Document

On page 60317, in corrections to figures regarding the Hospital Readmissions Reduction Program, we made an error in the: (1) Amount by which payments to hospitals would be reduced; and (2) number of hospitals that will have their base operating DRG payments reduced by the readmissions adjustment.