TABLE 11 TO SUBPART AAAAA OF PART 63—TOXICITY EQUIVALENCE FACTORS (TEFS) FOR HUMAN HEALTH RISK AS-SESSMENT OF POLYCHLORINATED DIBENZO-P-DIOXINS, DIBENZOFURANS, AND DIOXIN-LIKE POLYCHLORINATED BIPHENYLS—CONTINUED

Dioxin/Furan	2005 TEFs ¹
1,2,3,6,7,8-HxCDD	0.1
1,2,3,7,8,9-HxCDD	0.1
1,2,3,4,6,7,8-HpCDD	0.01
OCDD	0.0003
2,3,7,8-TCDF	0.1
1,2,3,7,8-PeCDF	0.03
2.3.4.7.8-PeCDF	0.3
1,2,3,4,7,8-HxCDF	0.1
1.2.3.6.7.8-HxCDF	0.1
1,2,3,7,8,9-HxCDF	0.1
2,3,4,6,7,8-HxCDF	0.1
1,2,3,4,6,7,8-HpCDF	0.01
1,2,3,4,7,8,9-HpCDF	0.01
OCDF	0.0003

¹EPA/100/R–10/005, "Recommended Toxicity Equivalence Factors (TEFs) for Human Health Risk Assessments of 2, 3, 7, 8-Tetrachlorodibenzo-p-dioxin and Dioxin-Like Compounds", December 2010. (See § 63.14 for availability.)

[FR Doc. 2024–14692 Filed 7–15–24; 8:45 am] BILLING CODE 6560–50–P

ENVIRONMENTAL PROTECTION AGENCY

40 CFR Part 170

[EPA-HQ-OPP-2022-0133; FRL-8528-04-OCSPP]

RIN 2070-AK92

Notification of Submission to the Secretary of Agriculture; Draft Final Rule; Pesticides; Agricultural Worker Protection Standard; Reconsideration of the Application Exclusion Zone Amendments

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notification of submission to the Secretary of Agriculture.

SUMMARY: This document notifies the public as required by the Federal Insecticide, Fungicide, and Rodenticide Act (FIFRA) that the EPA has forwarded a draft final rule to the U.S. Department of Agriculture (USDA) entitled "Pesticides; Agricultural Worker Protection Standard; Reconsideration of the Application Exclusion Zone Amendments." The draft regulatory document is not available to the public until after it has been signed and made available to the public by EPA.

DATES: See Unit I. under SUPPLEMENTARY INFORMATION.

ADDRESSES: The docket for this action, identified by docket identification (ID) number EPA–HQ–OPP–2022–0133, is available at *https://*

www.regulations.gov. That docket contains historical information and this

Federal Register document; it does not contain the draft final rule.

FOR FURTHER INFORMATION CONTACT:

Carolyn Schroeder, Pesticide Re-Evaluation Division (7508M), Office of Pesticide Programs, Environmental Protection Agency, 1200 Pennsylvania Ave. NW, Washington, DC 20460–0001; telephone number: (202) 566–2376; email address: *schroeder.carolyn@ epa.gov.*

SUPPLEMENTARY INFORMATION:

I. What action is EPA taking?

FIFRA section 25(a)(2)(B) requires the EPA to provide the USDA with a copy of any draft final rule at least 30 days before signing it in final form for publication in the Federal Register. The draft final rule is not available to the public until after it has been signed by EPA. If the Secretary of USDA comments in writing regarding the draft final rule within 15 days after receiving it, the EPA Administrator must include the comments of the USDA Secretary, if requested by the Secretary, and the EPA Administrator's response to those comments with the final rule that publishes in the Federal Register. If the Secretary of USDA does not comment in writing within 15 days after receiving the draft final rule, then the EPA Administrator may sign the final rule for publication in the Federal Register any time after the 15-day period.

II. Do any statutory and Executive Order reviews apply to this notification?

No. This document is merely a notification of submission to the Secretary of USDA. As such, none of the regulatory assessment requirements apply to this document.

List of Subjects in 40 CFR Part 170

Environmental protection, Pesticides, Agricultural worker, Pesticide handler, Employer, Farms, Forests, Nurseries, Greenhouses, Worker protection standard.

Dated: July 9, 2024.

Michal Freedhoff,

Assistant Administrator, Office of Chemical Safety and Pollution Prevention. [FR Doc. 2024–15447 Filed 7–15–24; 8:45 am]

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ENVIRONMENTAL PROTECTION AGENCY

40 CFR Part 180

[EPA-HQ-OPP-2020-0700; FRL-10420-01-OCSPP]

Trichoderma atroviride Strain K5 NRRL B–50520; Exemption From the Requirement of a Tolerance

AGENCY: Environmental Protection Agency (EPA). ACTION: Final rule.

SUMMARY: This regulation establishes an exemption from the requirement of a tolerance for residues of Trichoderma atroviride strain K5 NRRL B-50520 in or on all food commodities when used in accordance with label directions and good agricultural practices. Agrauxine Corp., submitted a petition to EPA under the Federal Food, Drug, and Cosmetic Act (FFDCA), requesting the exemption from a requirement of a tolerance. This regulation eliminates the need to establish a maximum permissible level for residues of Trichoderma atroviride strain K5 NRRL B-50520 under FFDCA when used in