Under section 307(b)(1) of the Clean Air Act, petitions for judicial review of this action must be filed in the United States Court of Appeals for the appropriate circuit by December 2, 2016. Filing a petition for reconsideration by the Administrator of this final rule does not affect the finality of this action for the purposes of judicial review nor does it extend the time within which a petition for judicial review may be filed, and shall not postpone the effectiveness of such rule or action. Parties with objections to this direct final rule are encouraged to file a comment in response to the parallel notice of proposed rulemaking for this action published in the proposed rules section of today's **Federal Register**, rather than file an immediate petition for judicial review of this direct final rule, so that the EPA can withdraw this direct final rule and address the comment in the proposed rulemaking. This action may not be challenged later in proceedings to enforce its requirements. (See section 307(b)(2).)

**List of Subjects in 40 CFR Part 62**

Environmental protection, Air pollution control, Solid waste incineration, Hospital/medical/infectious waste incineration.

**Authority:** 42 U.S.C. 7401 *et seq.*

Dated: May 17, 2016.

**Shaun L. McGrath,**

*Regional Administrator, Region 8.*

40 CFR part 62, subpart ZZ, is amended as follows:

## PART 62—APPROVAL AND PROMULGATION OF STATE PLANS FOR DESIGNATED FACILITIES AND POLLUTANTS

■ 1. The authority citation for part 62 continues to read as follows:

**Authority:** 42 U.S.C. 7401 *et seq.*

### Subpart ZZ—Wyoming

■ 2. Section 62.12610 is revised to read as follows:

**§ 62.12610 Identification of plan.**

Section 111(d)/129 Plan for Hospital/Medical/Infectious Waste Incinerators and the associated State regulation, Chapter 4, Section 5, and Chapter 5 of the Wyoming Air Quality Standards and Regulations, submitted by the State on September 7, 1999 and November 9, 1999, and as amended on May 13, 2015 and November 24, 2015.

■ 3. Section 62.12611 is revised to read as follows:

**§ 62.12611 Identification of sources.**

The plan applies to each individual hospital/medical/infectious waste incinerator:

(a) For which construction was commenced on or before June 20, 1996, or for which modification was commenced on or before March 16, 1998.

(b) For which construction was commenced after June 20, 1996 but no later than December 1, 2008, or for which modification is commenced after March 16, 1998 but no later than April 6, 2010.

■ 4. Section 62.12612 is revised to read as follows:

**§ 62.12612 Effective date.**

The effective date of the plan for hospital/medical/infectious waste incinerators is December 2, 2016.

**Editorial Note:** This document was received for publication by the Office of the Federal Register on September 26, 2016.

[FR Doc. 2016–23584 Filed 9–30–16; 8:45 am]

**BILLING CODE 6560–50–P**

## ENVIRONMENTAL PROTECTION AGENCY

### 40 CFR Part 180

**[EPA–HQ–OPP–2014–0920; FRL–9947–92]**

### Bacillus Mycoides Isolate J; 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 *Bacillus mycoides* isolate J in or on all agricultural commodities when used in accordance with label directions and good agricultural practices. Certis USA LLC submitted a petition to EPA under the Federal Food, Drug, and Cosmetic Act (FFDCA), requesting an exemption from the requirement of a tolerance. This regulation eliminates the need to establish a maximum permissible level for residues of *Bacillus mycoides* isolate J under FFDCA.

**DATES:** This regulation is effective October 3, 2016. Objections and requests for hearings must be received on or before December 2, 2016, 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–2014–0920, 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), West William Jefferson Clinton 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:** Robert McNally, Biopesticides and Pollution Prevention Division (7511P), Office of Pesticide Programs, Environmental Protection Agency, 1200 Pennsylvania Ave. NW., Washington, DC 20460–0001; main telephone number: (703) 305–7090; email address: *BPPDFRNotices@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(g), 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–2014–0920 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 December 2, 2016. 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–2014–0920, 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.html.* 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

In the **Federal Register** of January 28, 2015 (80 FR 4525) (FRL–9921–55), EPA issued a document pursuant to FFDCA section 408(d)(3), 21 U.S.C. 346a(d)(3), announcing the filing of a pesticide tolerance petition (PP 4F8252) by Certis USA LLC, 9145 Guilford Rd., Suite 175, Columbia, MD 21046. The petition requested that 40 CFR part 180 be amended by establishing an exemption from the requirement of a tolerance for residues of *Bacillus mycoides* isolate J, in or on all agricultural commodities. That document referenced a summary of the petition prepared by the petitioner Certis USA LLC, which is available in the docket via *http://www.regulations.gov.* There were no comments received in response to the notice of filing.

In addition, the Agency is removing the existing paragraph contained in section 180.1269 because that exemption from the requirement of a tolerance for *Bacillus mycoides* isolate J residues has expired.

## III. Final Rule

### *A. EPA's Safety Determination*

Section 408(c)(2)(A)(i) of FFDCA allows EPA to establish an exemption from the requirement for a tolerance (the legal limit for a pesticide chemical residue in or on a food) only if EPA determines that the exemption is ''safe.'' Section 408(c)(2)(A)(ii) of FFDCA defines ''safe'' to mean that ''there is a reasonable certainty that no harm will result from aggregate exposure to the pesticide chemical residue, including all anticipated dietary exposures and all other exposures for which there is reliable information.'' This includes exposure through drinking water and in residential settings but does not include occupational exposure. Pursuant to FFDCA section 408(c)(2)(B), in establishing or maintaining in effect an exemption from the requirement of a tolerance, EPA must take into account the factors set forth in FFDCA section 408(b)(2)(C), which require EPA to give special consideration to exposure of infants and children to the pesticide chemical residue in establishing a tolerance or tolerance exemption, and to ''ensure that there is a reasonable certainty that no harm will result to infants and children from aggregate exposure to the pesticide chemical residue . . . .'' Additionally, FFDCA section 408(b)(2)(D) requires that EPA consider ''available information concerning the cumulative effects of [a particular pesticide's] . . . residues and other substances that have a common mechanism of toxicity.''

EPA evaluated the available toxicity and exposure data on *Bacillus mycoides* isolate J and considered its validity, completeness, and reliability, as well as the relationship of this information to human risk. A full explanation of the data upon which EPA relied and its risk assessment based on that data can be found within the May 9, 2016, document entitled ''Federal Food, Drug, and Cosmetic Act (FFDCA) Considerations for *Bacillus mycoides* isolate J.'' This document, as well as other relevant information, is available in the docket for this action as described under **ADDRESSES**. Based upon its evaluation, EPA concludes that there is a reasonable certainty that no harm will result to the U.S. population, including infants and children, from aggregate exposure to residues of *Bacillus mycoides* isolate J. Therefore, an exemption from the requirement of a tolerance is established for residues of *Bacillus mycoides* isolate J in or on all agricultural commodities when used in accordance with label directions and good agricultural practices.

### *B. Analytical Enforcement Methodology*

An analytical method is not required for enforcement purposes for the reasons contained in the May 9, 2016, document entitled ''Federal Food, Drug, and Cosmetic Act (FFDCA) Considerations for *Bacillus mycoides* isolate J'' and because EPA is establishing an exemption from the requirement of a tolerance without any numerical limitation.

## IV. Statutory and Executive Order Reviews

This action establishes a tolerance exemption under FFDCA section 408(d) in response to a petition submitted to EPA. 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 action has been exempted from review under Executive Order 12866, this action 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 action 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).

Since tolerances and exemptions that are established on the basis of a petition under FFDCA section 408(d), such as the tolerance exemption in this action, do not require the issuance of a proposed rule, the requirements of the Regulatory Flexibility Act (RFA) (5 U.S.C. 601 *et seq.*) do not apply.

This action directly regulates growers, food processors, food handlers, and food retailers, not States or tribes. As a result, this action does not alter the relationships or distribution of power and responsibilities established by Congress in the preemption provisions of FFDCA section 408(n)(4). As such,

EPA 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, EPA 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 action. In addition, this action does not impose any enforceable duty or contain any unfunded mandate as described under Title II of the Unfunded Mandates Reform Act (UMRA) (2 U.S.C. 1501 *et seq.*).

This action does not involve any technical standards that would require EPA's consideration of voluntary consensus standards pursuant to section 12(d) of the National Technology Transfer and Advancement Act (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: September 15, 2016.

**Yu-Ting Guilaran,**

*Acting Director, 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. Revise § 180.1269 to read as follows:

**§ 180.1269 *Bacillus mycoides* isolate J; exemption from the requirement of a tolerance.**

An exemption from the requirement of a tolerance is established for residues of *Bacillus mycoides* isolate J in or on all agricultural commodities when used in accordance with label directions and good agricultural practices.

[FR Doc. 2016–23608 Filed 9–30–16; 8:45 am]

**BILLING CODE 6560–50–P**

## FEDERAL COMMUNICATIONS COMMISSION

### 47 CFR Part 54

**[WC Docket Nos. 11–42, 09–197 and 10–90; FCC 16–38]**

### Lifeline and Link Up Reform and Modernization, Telecommunications Carriers Eligible for Universal Service Support, Connect America Fund

**AGENCY:** Federal Communications Commission.

**ACTION:** Final rule; announcement of effective date.

**SUMMARY:** In this document, the Federal Communications Commission (Commission) announces that the Office of Management and Budget (OMB) has approved, for a period of three years, the information collection requirements associated with certain of the provision of the rules adopted as part of the Commission's Third Further Notice of Proposed Rulemaking, Order on Reconsideration, and Further Report and Order, (*Lifeline Third Reform Order*). This notice is consistent with the *Lifeline Third Reform Order,* which stated that the Commission would publish a document in the **Federal Register** announcing OMB approval and the effective date of those rules.

**DATES:** The rule amendments to 47 CFR 54.202(a)(6), (d), and (e), and 54.205(c) published at 81 FR 33025, May 24, 2016, will become effective October 3, 2016. The rule amendments to 47 CFR 54.101, 54.401(a)(2), (b), (c), (f), 54.403(a), 54.405(e)(1), (e)(3) through (e)(5), 54.407(a), (c)(2), (d), 54.408, 54.409(a)(2), 54.410(b) through (e), (g) through (h), 54.411, 54.416(a)(3), 54.420(b), and 54.422(b)(3) will become effective December 2, 2016 . The rule amendments to 47 CFR 54.410(f) will become effective January 1, 2017.

The rule amendments to 47 CFR 54.400(l) are applicable October 3, 2016. The rule amendments to 47 CFR 54.400(f), (j), and (m) through (o) are applicable December 2, 2016.

**FOR FURTHER INFORMATION CONTACT:** Christian Hoefly, Wireline Competition Bureau, Telecommunications Access Policy Division at (202) 418–3607 or at *christian.hoefly@fcc.gov.*

**SUPPLEMENTARY INFORMATION:** This document announces that, on September 20, 2016, OMB approved, for a period of three years, the information collection requirements contained in the Commission's *Order,* FCC 16–38, published at 81 FR 33025, May 24, 2016. The OMB Control Number is 3060–0819. The Commission publishes this notice as an announcement of the effective date rules requiring OMB approval. If you have any comments on the burden estimates listed below, or how the Commission can improve the collections and reduce any burdens caused thereby, please contact Nicole Ongele, Federal Communications Commission, Room 1–A620, 445 12th Street SW., Washington, DC 20554. Please include the OMB Control Number, 3060–0819, in your correspondence. The Commission will also accept your comments via email at *PRA@fcc.gov.*

To request material in accessible formation for people with disabilities (Braille, large print, electronic files, audio format), send an email to *fcc504@fcc.gov* or call the Consumer and Governmental Affairs Bureau at (202) 418–0530 (voice), (202) 418–0432 (TTY).

**Synopsis**

As required by the Paperwork Reduction Act of 1995 (44 U.S.C. 3507), the FCC is notifying the public that it received OMB approval on September 20, 2016, for the information collection requirements contained in the Commission's rules in 47 CFR part 54.

Under 5 CFR part 1320, an agency may not conduct or sponsor a collection of information unless it displays a current, 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 that does not display a current, valid OMB Control Number. The OMB Control Number is 3060–0819.

The foregoing notice is required by the Paperwork Reduction Act of 1995, Public Law 104–13, October 1, 1995, and 44 U.S.C. 3507.

The total annual reporting burdens and costs for the respondents are as follows:

*OMB Control Number:* 3060–0819.

*OMB Approval Date:* 9/20/2016.

*OMB Expiration Date:* 9/30/2019.

*Title:* Lifeline and Link Up Reform and Modernization, Telecommunications Carriers Eligible for Universal Service Support, Connect America Fund.

*Form Number:* FCC Forms 497, 481 & 555.

*Type of Review:* Revision of a currently approved collection.