

to the lack of potential for adverse effects. Because there are no threshold levels of concern with the toxicity, pathogenicity, or infectivity of *Kosakonia cowanii* strain SYM00028, EPA determined that no additional margin of safety is necessary to protect infants and children as part of the qualitative assessment conducted. Based upon its evaluation in the *Kosakonia cowanii* strain SYM00028 Human Health Assessment, which concludes that there are no risks of concern from aggregate exposure to *Kosakonia cowanii* strain SYM00028, 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 *Kosakonia cowanii* strain SYM00028.

B. Analytical Enforcement Methodology

An analytical method is not required for *Kosakonia cowanii* strain SYM00028 because EPA is establishing an exemption from the requirement of a tolerance without any numerical limitation.

C. Conclusion

Therefore, an exemption from the requirement of a tolerance is established for residues of *Kosakonia cowanii* strain SYM00028 in or on all food commodities when used in accordance with label directions and good agricultural practices.

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, 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 (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 (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 (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: December 6, 2021.

Edward Messina,

Director, Office of Pesticide Programs.

Therefore, for the reasons stated in the preamble, EPA is amending 40 CFR chapter I as follows:

PART 180—TOLERANCES AND EXEMPTIONS FOR PESTICIDE CHEMICAL RESIDUES IN FOOD

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

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

■ 2. Add § 180.1387 to subpart D to read as follows:

§ 180.1387 *Kosakonia cowanii* strain SYM00028; exemption from the requirement of a tolerance.

An exemption from the requirement of a tolerance is established for residues of *Kosakonia cowanii* strain SYM00028 in or on all food commodities when used in accordance with label directions and good agricultural practices.

[FR Doc. 2021–26846 Filed 12–13–21; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

42 CFR Parts 413 and 512

[CMS–1749–CN]

RIN 0938–AU39

Medicare Program; End-Stage Renal Disease Prospective Payment System, Payment for Renal Dialysis Services Furnished to Individuals With Acute Kidney Injury, End-Stage Renal Disease Quality Incentive Program, and End-Stage Renal Disease Treatment Choices Model; Correction

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

ACTION: Final rule; correction.

SUMMARY: This document corrects a typographic error that appeared in the final rule published in the **Federal Register** on November 8, 2021 entitled “Medicare Program; End-Stage Renal Disease Prospective Payment System, Payment for Renal Dialysis Services Furnished to Individuals With Acute Kidney Injury, End-Stage Renal Disease Quality Incentive Program, and End-Stage Renal Disease Treatment Choices Model.”

DATES: This correction is effective January 1, 2022.

FOR FURTHER INFORMATION CONTACT:

ESRDPayment@cms.hhs.gov, for issues related to the ESRD PPS and coverage and payment for renal dialysis services furnished to individuals with AKI.

ESRDApplications@cms.hhs.gov, for issues related to the Transitional Add-On Payment Adjustment for New and Innovative Equipment and Supplies (TPNIES).

Delia Houseal, (410) 786-2724, for issues related to the ESRD QIP.

ETC-CMMI@cms.hhs.gov, for issues related to the ESRD Treatment Choices (ETC) Model.

SUPPLEMENTARY INFORMATION:**I. Background**

In FR Doc. 2021-23907 of November 8, 2021 (86 FR 61874), there was a typographic error that is identified and corrected by the Correction of Errors section below. The correction in this document is effective as if it had been included in the document published November 8, 2021. Accordingly, the correction is effective January 1, 2022.

II. Summary of Error

On page 61874, in the third sentence of the first column, we inadvertently left the number “412” in the CFR citation at the top of the document. Therefore, the number “412” should be deleted.

III. Waiver of Proposed Rulemaking

We ordinarily publish a notice of proposed rulemaking in the **Federal Register** to provide a period for public comment before the provisions of a rule take effect in accordance with section 553(b) of the Administrative Procedure Act (APA) (5 U.S.C. 553(b)). However, we can waive this notice and comment procedure if the Secretary finds, for good cause, that the notice and comment process is impracticable, unnecessary, or contrary to the public interest, and incorporates a statement of the finding and the reasons therefore in the notice.

Section 553(d) of the APA ordinarily requires a 30-day delay in effective date of final rules after the date of their publication in the **Federal Register**. This 30-day delay in effective date can be waived, however, if an agency finds for good cause that the delay is impracticable, unnecessary, or contrary to the public interest, and the agency incorporates a statement of the findings and its reasons in the rule issued.

We believe that this correcting document does not constitute a rule that would be subject to the notice and comment or delayed effective date requirements. This document corrects a typographic error and does not make

substantive changes to the policies or payment methodologies that were adopted in the final rule. Thus, this correcting document is intended to ensure that the information is accurately reflected in the final rule.

Even if this were a rulemaking to which the notice and comment and delayed effective date requirements applied, we find that there is good cause to waive such requirements. Undertaking further notice and comment procedures to incorporate the correction in this document into the calendar year (CY) 2022 End-Stage Renal Disease (ESRD) Prospective Payment System (PPS) final rule or delaying the effective date of the correction would be contrary to the public interest because it is in the public interest to ensure that the rule accurately reflects our policies as of the date they take effect. Further, such procedures would be unnecessary because we are not making any substantive revisions to the final rule, but rather, we are simply correcting the **Federal Register** document to reflect the policies that we previously proposed, received public comment on, and subsequently finalized in the CY 2022 ESRD PPS final rule. For these reasons, we believe there is good cause to waive the requirements for notice and comment and delay in effective date.

IV. Correction of Errors

In FR Doc. 2021-23907 of November 8, 2021 (86 FR 61874), make the following correction:

On page 61874, in the first column; in the third sentence, remove the number “412” from the CFR citation.

Karuna Seshasai,

*Executive Secretary to the Department,
Department of Health and Human Services.*

[FR Doc. 2021-26914 Filed 12-13-21; 8:45 am]

BILLING CODE 4120-01-P

**FEDERAL COMMUNICATIONS
COMMISSION****47 CFR Part 54**

[WC Docket No. 21-93; DA 21-1499; FR ID 61508]

**Establishing Emergency Connectivity
Fund To Close the Homework Gap**

AGENCY: Federal Communications Commission.

ACTION: Final rule.

SUMMARY: In this document, the Wireline Competition Bureau (the Bureau) grants a petition for an expedited waiver of the Emergency

Connectivity Fund (ECF) Program's invoice filing deadline submitted by the State E-rate Coordinators' Alliance (SECA) and clarifies the service delivery date for certain funding requests.

DATES: Effective December 14, 2021.

FOR FURTHER INFORMATION CONTACT: For further information, please contact Gabriela Gross, Telecommunications Access Policy Division, Wireline Competition Bureau, at *gabriela.gross@fcc.gov*.

SUPPLEMENTARY INFORMATION: This is a summary of the Commission's Order in WC Docket No. 21-93; DA 21-1499, adopted and released on December 2, 2021. The full text of this document is available for public inspection on the Commission's website at <https://www.fcc.gov/document/wcb-waives-ecf-invoice-deadline-and-clarifies-service-delivery-date>.

Synopsis**I. Introduction**

1. In the Order, the Bureau grants a petition for an expedited waiver of the ECF Program's invoice filing deadline submitted by SECA. Specifically, and subject to the limitations stated in the Order, the Bureau waives §§ 54.1711(d) and (e) of the Commission's rules to provide relief to applicants that: (a) Applied for ECF funding during the first or second application filing windows; (b) incorrectly used June 30, 2022 as the service delivery date on their ECF FCC Form 471 applications for equipment and/or other non-recurring services, rather than the actual service delivery date; and (c) received a funding commitment decision letter (FCDL) or revised funding commitment decision letter (RFCDL) noting August 29, 2022 as the invoice filing deadline based on the incorrect service delivery date (Affected Program Participants).

2. Accordingly, the Bureau directs the Universal Service Administrative Company (USAC), the Administrator of the ECF Program, to continue to use the August 29, 2022 invoice filing deadline noted on the Affected Program Participants' FCDLs and RFCDLs and allow them to submit their requests for reimbursement on or before this date. To the extent other applicants incorrectly used June 30, 2022 as the service delivery date for equipment and/or non-recurring services, rather than the actual delivery date, but have not yet received an FCDL or RFCDL with an invoice filing deadline, the Bureau directs USAC to use June 30, 2022 as the service delivery date for these requests. The Bureau also extends this relief to service providers that agreed to file requests for reimbursement on behalf of