not impose additional requirements beyond those imposed by state law. For that reason, this proposed action:

- Is not a ''significant regulatory action'' subject to review by the Office of Management and Budget under Executive Orders 12866 (58 FR 51735, October 4, 1993) and 13563 (76 FR 3821, January 21, 2011);
- Is not an Executive Order 13771 (82 FR 9339, February 2, 2017) regulatory action because SIP approvals are exempted under Executive Order 12866;
- Does not impose an information collection burden under the provisions of the Paperwork Reduction Act (44 U.S.C. 3501 *et seq.*);
- Is certified as not having a significant economic impact on a substantial number of small entities under the Regulatory Flexibility Act (5 U.S.C. 601 *et seq.*);
- Does not contain any unfunded mandate or significantly or uniquely affect small governments, as described in the Unfunded Mandates Reform Act of 1995 (Pub. L. 104–4);
- Does not have Federalism implications as specified in Executive Order 13132 (64 FR 43255, August 10, 1999);
- Is not an economically significant regulatory action based on health or safety risks subject to Executive Order 13045 (62 FR 19885, April 23, 1997);
- Is not a significant regulatory action subject to Executive Order 13211 (66 FR 28355, May 22, 2001);
- Is not subject to requirements of section 12(d) of the National Technology Transfer and Advancement Act of 1995 (15 U.S.C. 272 note) because application of those requirements would be inconsistent with the Act; and
- Does not provide the EPA with the discretionary authority to address disproportionate human health or environmental effects with practical, appropriate, and legally permissible methods under Executive Order 12898 (59 FR 7629, February 16, 1994).

In addition, the SIP is not approved to apply on any Indian reservation land or in any other area where the EPA or an Indian tribe has demonstrated that a tribe has jurisdiction. In those areas of Indian country, the rule does not have tribal implications and will not impose substantial direct costs on tribal governments or preempt tribal law as specified by Executive Order 13175 (65 FR 67249, November 9, 2000).

List of Subjects in 40 CFR Part 52

Environmental protection, Air pollution control, Incorporation by reference, Infrastructure SIP, Interstate transport, Nitrogen oxides, Volatile organic compounds.

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

Dated: February 1, 2019.

Deborah Jordan,

Acting Regional Administrator, Region IX.

[FR Doc. 2019–03564 Filed 2–27–19; 8:45 am]

BILLING CODE 6560–50–P

ENVIRONMENTAL PROTECTION AGENCY

40 CFR Part 63

[EPA–HQ–OAR–2018–0794; FRL–9989–76–OAR]

RIN 2060–AT99

National Emission Standards for Hazardous Air Pollutants: Coal- and Oil-Fired Electric Utility Steam Generating Units—Reconsideration of Supplemental Finding and Residual Risk and Technology Review

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice of public hearing and extension of public comment period.

SUMMARY: On February 7, 2019, the Environmental Protection Agency (EPA) published a document in the **Federal Register** to announce its proposed National Emission Standards for Hazardous Air Pollutants: Coal- and Oil-Fired Electric Utility Steam Generating Units—Reconsideration of Supplemental Finding and Residual Risk and Technology Review. The document also requested public comment on the proposed action. The EPA is announcing that it will hold a public hearing to provide interested parties the opportunity to present data, views, or arguments concerning the proposed action. In addition, the EPA will extend the public comment period.

DATES: *Public Hearing:* The EPA will hold a public hearing on March 18, 2019, in Washington, DC. The deadline for accepting written comments is being extended by 9 days, to April 17, 2019. Please refer to the **SUPPLEMENTARY INFORMATION** section for additional information on the public hearing.

ADDRESSES: The hearing will be held at the EPA WJC East Building, 1201 Constitution Avenue NW, Room 1153, Washington, DC 20004. The hearing will convene at 8:00 a.m. (local time) and will conclude at 6:00 p.m. There will be a lunch break from noon to 1 p.m. The EPA's website for this rulemaking, which includes the proposal and information about the hearing, can be found at: *https://www.epa.gov/mats/proposed-revised-supplemental-finding-and-results-residual-risk-and-technology-review.* Written comments on the proposed rule may be submitted to the EPA electronically, by mail, facsimile, or through hand delivery/courier. Please refer to the proposal (84 FR 2670) for the addresses and detailed instructions.

Because this hearing is being held at a U.S. government facility, individuals planning to attend the hearing should be prepared to show valid picture identification to the security staff to gain access to the meeting room. Please note that the REAL ID Act, passed by Congress in 2005, established new requirements for entering federal facilities. For purposes of the REAL ID Act, the EPA will accept government-issued IDs, including driver's licenses from the District of Columbia and all states and territories. Acceptable alternative forms of identification include: Federal employee badges, passports, enhanced driver's licenses, and military identification cards. Additional information on the REAL ID Act is available at: *https://www.dhs.gov/real-id.*

Any objects brought into the building need to fit through the security screening system, such as a purse, laptop bag, or small backpack. Demonstrations will not be allowed on federal property for security reasons.

FOR FURTHER INFORMATION CONTACT: The EPA will begin pre-registering speakers for the hearing upon publication of this document in the **Federal Register**. To register to speak at the hearing, please use the online registration form available at *https://www.epa.gov/mats/proposed-revised-supplemental-finding-and-results-residual-risk-and-technology-review* or contact Adrian Gates at (919) 541–4860 or at *gates.adrian@epa.gov.* The last day to pre-register to speak at the hearing will be March 14, 2019. On March 15, 2019, the EPA will post at *https://www.epa.gov/mats/proposed-revised-supplemental-finding-and-results-residual-risk-and-technology-review* a general agenda for the hearing that will list pre-registered speakers in approximate order. The EPA will make every effort to follow the schedule as closely as possible on the day of the hearing; however, please plan for the hearing to run either ahead of schedule or behind schedule.

Additionally, requests to speak will be taken the day of the hearing at the hearing registration desk. The EPA will make every effort to accommodate all speakers who arrive and register, although preferences on speaking times may not be able to be fulfilled.

SUPPLEMENTARY INFORMATION: Each commenter will have 5 minutes to

provide oral testimony. The EPA encourages commenters to provide the EPA with a copy of their oral testimony electronically (via email) or in hard copy form.

The EPA may ask clarifying questions during the oral presentations, but will not respond to the presentations at that time. Written statements and supporting information submitted during the comment period will be considered with the same weight as oral comments and supporting information presented at the public hearing. Commenters should notify Adrian Gates if they will need specific equipment or if there are other special needs related to providing comments at the hearing. Verbatim transcripts of the hearing and written statements will be included in the docket for the rulemaking.

Please note that any updates made to any aspect of the hearing will be posted online at *https://www.epa.gov/mats/proposed-revised-supplemental-finding-and-results-residual-risk-and-technology-review.* While the EPA expects the hearing to go forward as set forth above, please monitor our website or contact Adrian Gates at (919) 541–4860 or *gates.adrian@epa.gov* to determine if there are any updates. The EPA does not intend to publish a document in the **Federal Register** announcing updates.

The EPA will not provide audiovisual equipment. Commenters should notify Adrian Gates when they pre-register to speak that they will require the service of a translator or special accommodations such as audio description. We may not be able to arrange accommodations without advanced notice.

Dated: February 25, 2019.

Panagiotis Tsirigotis,

Director, Office of Air Quality Planning and Standards.

[FR Doc. 2019–03518 Filed 2–27–19; 8:45 am]

BILLING CODE 6560–50–P

ENVIRONMENTAL PROTECTION AGENCY

40 CFR Part 180

[EPA–HQ–OPP–2017–0418; FRL–9970–24]

RIN 2070–ZA16

Fenoxaprop-ethyl, Flufenpyr-ethyl, Imazapyr, Maleic hydrazide, Pyrazon, Quinclorac, Triflumizole, et al.; Proposed Tolerance and Tolerance Exemption Actions

Correction

In proposed rule document 2019–00787, appearing on pages 1691 through 1697 in the issue of Tuesday, February 5, 2019, make the following correction:

On page 1691, in the first column, under the **DATES** heading, ''February 5, 2019'' should read ''April 8, 2019''.

[FR Doc. C1–2019–00787 Filed 2–27–19; 8:45 am]

BILLING CODE 1301–00–D

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

42 CFR Parts 405, 424, 455, and 457

[CMS–6058–RCN]

RIN 0938–AS84

Medicare, Medicaid, and Children's Health Insurance Programs; Program Integrity Enhancements to the Provider Enrollment Process; Extension of Timeline for Publication of the Final Rule

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

ACTION: Extension of timeline for publication of a final rule.

SUMMARY: This document announces the extension of the timeline for publication of the ''Medicare, Medicaid, and Children's Health Insurance Programs; Program Integrity Enhancements to the Provider Enrollment Process'' final rule. We are issuing this document in accordance with the Social Security Act (the Act), which requires notice to be provided in the **Federal Register** if there are exceptional circumstances that cause us to publish a final rule more than 3 years after the publication date of the proposed rule. In this case, the complexity of the rule and the scope of the comments received warrant the extension of the timeline for publication.

DATES: The timeline for publication of the final is extended for 1 year, until March 1, 2020.

FOR FURTHER INFORMATION CONTACT: Frank Whelan, (410) 786–1302.

SUPPLEMENTARY INFORMATION: In the March 1, 2016 **Federal Register** (81 FR 10720), we published a proposed rule titled ''Medicare, Medicaid, and Children's Health Insurance Programs; Program Integrity Enhancements to the Provider Enrollment Process'' that would implement sections of the Affordable Care Act that require Medicare, Medicaid, and Children's Health Insurance Program (CHIP) providers and suppliers to disclose certain current and previous affiliations with other providers and suppliers. This proposed rule would also provide us with additional authority to deny or revoke a provider's or supplier's Medicare enrollment. These and other important provisions in the proposed rule would: (1) Eliminate significant program integrity loopholes of long-standing concern to CMS and the Department; and (2) help halt and deter ongoing fraudulent and abusive behavior, including patient harm, in Medicare, Medicaid, and CHIP.

Section 1871(a)(3)(A) of the Act requires the Secretary of the Department of Health and Human Services, in consultation with the Director of the Office of Management and Budget (OMB), to establish a regular timeline for the publication of a final rule based on the previous publication of a proposed rule or an interim final rule. Section 1871(a)(3)(B) of the Act allows the timeline for publishing Medicare final regulations to vary based on the complexity of the regulation, the number and scope of comments received, and other related factors. The timeline for publishing the final rule, however, cannot exceed 3 years from the date of publishing the proposed regulation unless there are exceptional circumstances. The Secretary may extend the initial targeted publication date of the final rule if the Secretary provides public notice thereof, including a brief explanation of the justification for the variation, no later than the rule's previously established proposed publication date.

After consultation with the Director of OMB, the Department, through CMS, published a notice in the December 30, 2004 **Federal Register** (69 FR 78442) establishing a general 3-year timeline for publishing Medicare final rules after the publication of a proposed or interim final rule. Consistent with this, the final rule for the March 1, 2016 proposed rule was to be published by March 1, 2019.

This document announces an extension of the timeline for publication of the final rule due to exceptional circumstances. Based on both the public comments received and internal stakeholder feedback, we have determined that more time is needed to address and resolve certain complex policy and operational issues that the commenters and stakeholders raised. We stress that our decision in this matter to extend the timeline for issuing a final rule should not be viewed as a diminution of the Department's commitment to timely and effective rulemaking. Our goal remains to publish, as expeditiously as feasible, a final rule that strengthens our program integrity efforts while minimizing the