this action merely proposes to approve state law as meeting federal requirements and does not impose additional requirements beyond those imposed by state law. For that reason, this action:

- Is not a ''significant regulatory action'' subject to review by the Office of Management and Budget under Executive Order 12866 (58 FR 51735, October 4, 1993);
- 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 CAA; and
- does not provide EPA with the discretionary authority to address, as appropriate, disproportionate human health or environmental effects, using practicable and legally permissible methods, under Executive Order 12898 (59 FR 7629, February 16, 1994).

In addition, this proposed rule does not have tribal implications as specified by Executive Order 13175 (65 FR 67249, November 9, 2000), because the SIP is not approved to apply in Indian country located in the state, and EPA notes that it will not impose substantial direct costs on tribal governments or preempt tribal law.

List of Subjects in 40 CFR Part 52

Environmental protection, Air pollution control, Incorporation by reference, Intergovernmental relations, Nitrogen dioxides, Ozone, Particulate matter, Reporting and recordkeeping requirements, Volatile organic compounds.

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

Dated: March 27, 2014.

Ron Curry,

Regional Administrator, Region 6.

[FR Doc. 2014–08342 Filed 4–14–14; 8:45 am]

BILLING CODE 6560–50–P

ENVIRONMENTAL PROTECTION AGENCY

40 CFR Part 62

[EPA–R07–OAR–2013–0692; FRL 9909–44–Region 7]

Approval and Promulgation of State Plans for Designated Facilities and Pollutants; Air Emissions From Existing Municipal Solid Waste Landfills; State of Missouri

AGENCY: Environmental Protection Agency (EPA).

ACTION: Proposed rule.

SUMMARY: The Environmental Protection Agency (EPA) is proposing to approve the revision to the state section 111 plan submitted by the State of Missouri for controlling emissions from existing municipal solid waste (MSW) landfills. The revised State Plan incorporates revisions to the Emissions Guidelines (EG) for MSW landfills promulgated by EPA in 2000 and 2006. The plan also corrects typographical and administrative changes in the Missouri rules. The plan was submitted to fulfill the requirements of section 111 of the Clean Air Act (CAA).

DATES: Comments on this proposed action must be received in writing by May 15, 2014.

ADDRESSES: Submit your comments, identified by Docket ID No. EPA–R07–OAR–2013–0692, by mail to Craig Bernstein, Environmental Protection Agency, Air Planning and Development Branch, 11201 Renner Boulevard, Lenexa, Kansas 66219. Comments may also be submitted electronically or through hand delivery/courier by following the detailed instructions in the **ADDRESSES** section of the direct final rule located in the rules section of this **Federal Register**.

FOR FURTHER INFORMATION CONTACT: Craig Bernstein, Environmental Protection Agency, Air Planning and Development Branch, 11201 Renner Boulevard, Lenexa, Kansas 66219; at 913–551–7688; or by email at *Bernstein.craig@epa.gov.*

SUPPLEMENTARY INFORMATION: In the final rules section of today's **Federal Register**, EPA is approving the state's 111(d) plan revision as a direct final rule without prior proposal because the Agency views this as a noncontroversial action and anticipates no relevant adverse comments because the revisions are administrative and consistent with Federal regulations. A detailed rationale for the approval is set forth in the direct final rule. If no relevant adverse comments are received in response to this action, no further activity is contemplated in relation to this action. If EPA receives relevant adverse comments, the direct final rule will be withdrawn and all public comments received will be addressed in a subsequent final rule based on this proposed action. EPA will not institute a second comment period on this action. Any parties interested in commenting on this action should do so at this time. Please note that if EPA receives adverse comment on part of this rule and if that part can be severed from the remainder of the rule, EPA may adopt as final those parts of the rule that are not the subject of an adverse comment. For additional information, see the direct final rule which is located in the rules section of this **Federal Register**.

Dated: April 3, 2014.

Karl Brooks,

Regional Administrator, Region 7.

[FR Doc. 2014–08337 Filed 4–14–14; 8:45 am]

BILLING CODE 6560–50–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

42 CFR Part 100

RIN 0906–AB00

National Vaccine Injury Compensation Program: Revisions to the Vaccine Injury Table

AGENCY: Office of the Secretary, HHS.

ACTION: Notice of public hearing.

SUMMARY: This document announces a public hearing to receive information and views on the Notice of Proposed Rulemaking (NPRM) entitled ''National Vaccine Injury Compensation Program: Revisions to the Vaccine Injury Table.''

DATES: The public hearing will be held on April 28, 2014, from 10:00 a.m.–11:30 a.m. (EDT).

ADDRESSES: The public hearing will be held in Conference Room 10–65 in the Parklawn Building, 5600 Fishers Lane, Rockville, Maryland 20857.

FOR FURTHER INFORMATION CONTACT: Dr. Avril Melissa Houston, Acting Director, Division of Vaccine Injury Compensation, at 855–266–2427 or by email at *ahouston@hrsa.gov.*

SUPPLEMENTARY INFORMATION: The National Childhood Vaccine Injury Act