

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 Clean Air 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, Intergovernmental relations, Ozone, Reporting and recordkeeping requirements, Volatile organic compounds.

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

Dated: December 21, 2017.

Alexis Strauss,

Acting Regional Administrator, Region IX.

[FR Doc. 2018–00022 Filed 1–8–18; 8:45 am]

BILLING CODE 6560–50–P

ENVIRONMENTAL PROTECTION AGENCY

40 CFR Part 52

[EPA–R05–OAR–2007–1092; FRL–9972–74–Region 5]

Air Plan Approval; Michigan Minor New Source Review

AGENCY: Environmental Protection Agency (EPA).

ACTION: Proposed rule; reopening of the comment period.

SUMMARY: The Environmental Protection Agency (EPA) is reopening the comment period for a proposed Clean Air Act rule published August 15, 2017. An appendix to one of the documents EPA proposed to approve was not available on *Regulations.gov* as required; therefore, EPA is reopening the comment period for 15 days.

DATES: Comments must be received on or before January 24, 2018.

ADDRESSES: Submit your comments, identified by Docket ID No. EPA–R05–OAR–2007–1092 at <https://www.regulations.gov>, or via email to damico.genvieve@epa.gov. For comments submitted at *Regulations.gov*, follow the online instructions for submitting comments. Once submitted, comments cannot be edited or removed from *Regulations.gov*. For either manner of submission, EPA may publish any comment received to its public docket. Do not submit electronically any information you consider to be Confidential Business Information (CBI) or other information whose disclosure is restricted by statute. Multimedia submissions (audio, video, etc.) must be accompanied by a written comment. The written comment is considered the official comment and should include discussion of all points you wish to make. EPA will generally not consider comments or comment contents located outside of the primary submission (*i.e.* on the web, cloud, or other file sharing system). For additional submission methods, please contact the person identified in the **FOR FURTHER INFORMATION CONTACT** section. For the full EPA public comment policy, information about CBI or multimedia submissions, and general guidance on making effective comments, please visit <https://www2.epa.gov/dockets/commenting-epa-dockets>.

FOR FURTHER INFORMATION CONTACT: Rachel Rineheart, Environmental Engineer, Air Permits Section, Air Programs Branch (AR–18J), Environmental Protection Agency, Region 5, 77 West Jackson Boulevard, Chicago, Illinois 60604, (312) 886–7017, Rineheart.rachel@epa.gov.

SUPPLEMENTARY INFORMATION:

Throughout this document whenever “we,” “us,” or “our” is used, we mean the EPA.

On August 15, 2017, EPA proposed to approve certain changes to Michigan’s minor new source review program which is contained in Part 2 of the Michigan Administrative Code. EPA had previously reopened the comment

period due to an incomplete docket from November 2, 2017 to December 4, 2017. The file containing the state’s September 2, 2003 submittal made available on *Regulations.gov* on September 12, 2017, was missing Attachment H which contained the state’s technical analysis of the rule changes. The missing attachment was made available on *regulations.gov* on December 6, 2017, and EPA is reopening the comment period for an additional 15 days. The comment period now closes on January 24, 2018.

Dated: December 20, 2017.

Robert Kaplan,

Acting Regional Administrator, Region 5.

[FR Doc. 2018–00023 Filed 1–8–18; 8:45 am]

BILLING CODE 6560–50–P

ENVIRONMENTAL PROTECTION AGENCY

40 CFR Part 282

[EPA–R06–UST–2017–0504; FRL–9968–28–Region 6]

Oklahoma: Final Approval of State Underground Storage Tank Program Revisions and Incorporation by Reference

AGENCY: Environmental Protection Agency (EPA).

ACTION: Proposed rule.

SUMMARY: Pursuant to the Resource Conservation and Recovery Act (RCRA or Act), the Environmental Protection Agency (EPA) is proposing to approve revisions to the State of Oklahoma’s Underground Storage Tank (UST) program submitted by the State. This action is based on EPA’s determination that these revisions satisfy all requirements needed for program approval. This action also proposes to codify EPA’s approval of Oklahoma’s state program and to incorporate by reference those provisions of the State regulations that we have determined meet the requirements for approval. The provisions will be subject to EPA’s inspection and enforcement authorities under sections 9005 and 9006 of RCRA subtitle I and other applicable statutory and regulatory provisions.

DATES: Send written comments by February 8, 2018.

ADDRESSES: Submit any comments, identified by EPA–R06–UST–2017–0504, by one of the following methods:

1. *Federal eRulemaking Portal:* <http://www.regulations.gov>. Follow the on-line instructions for submitting comments.
2. *Email:* lincoln.audray@epa.gov.
3. *Mail:* Audray Lincoln, Region 6, Project Officer, LUST Prevention/

Corrective Action Section (6MM-XU), Multimedia Division, EPA Region 6, 1445 Ross Avenue, Dallas, Texas 75202-2733.

4. *Hand Delivery or Courier:* Deliver your comments to Audray Lincoln, Region 6, Project Officer, LUST Prevention/Corrective Action Section (6MM-XU), Multimedia Division, EPA Region 6, 1445 Ross Avenue, Dallas, Texas 75202-2733.

Instructions: Direct your comments to Docket ID No. EPA-R06-UST-2017-0504. EPA's policy is that all comments received will be included in the public docket without change and may be available online at <http://www.regulations.gov>, including any personal information provided, unless the comment includes information claimed to be Confidential Business Information (CBI) or other information whose disclosure is restricted by statute. Do not submit information that you consider to be CBI or otherwise protected through <http://www.regulations.gov>, or email. The Federal <http://www.regulations.gov> website is an "anonymous access" system, which means the EPA will not know your identity or contact information unless you provide it in the body of your comment. If you send an email comment directly to the EPA without going through <http://www.regulations.gov>, your email address will be automatically captured and included as part of the comment that is placed in the public docket and made available on the internet. If you submit an electronic comment, the EPA recommends that you include your name and other contact information in the body of your comment and with any disk or CD-ROM you submit. If the EPA cannot read your comment due to technical difficulties, and cannot contact you for clarification, the EPA may not be able to consider your comment. Electronic files should avoid the use of special characters, any form of encryption, and be free of any defects or viruses.

You can view and copy the documents that form the basis for this codification and associated publicly available materials from 8:30 a.m. to 4:00 p.m. Monday through Friday at the following location: EPA Region 6, 1445 Ross Avenue, Dallas, Texas 75202-2733, phone number (214) 665-2239. Interested persons wanting to examine these documents should make an appointment with the office at least two weeks in advance.

FOR FURTHER INFORMATION CONTACT: Ms. Audray Lincoln, Region 6, Project Officer, LUST Prevention/Corrective Action Section (6MM-XU), Multimedia

Division, EPA Region 6, 1445 Ross Avenue, Dallas, Texas 75202-2733, phone number (214) 665-2239, email address lincoln.audray@epa.gov.

SUPPLEMENTARY INFORMATION: For additional information, see the direct final rule published in the "Rules and Regulations" section of this **Federal Register**.

Authority: This rule is issued under the authority of Sections 2002(a), 9004, and 7004(b) of the Solid Waste Disposal Act, as amended, 42 U.S.C. 6912, 6991c, 6991d, and 6991e.

Dated: November 3, 2017.

Samuel Coleman,

Acting Regional Administrator, Region 6.

[FR Doc. 2018-00038 Filed 1-8-18; 8:45 am]

BILLING CODE 6560-50-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

42 CFR Part 493

[CMS-3326-NC]

RIN 0938-ZB40

Request for Information: Revisions to Personnel Regulations, Proficiency Testing Referral, Histocompatibility Regulations and Fee Regulations Under the Clinical Laboratory Improvement Amendments of 1988 (CLIA)

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

ACTION: Request for information.

SUMMARY: This request for information seeks public comment regarding several items related to Clinical Laboratory Improvement Amendments of 1988 (CLIA) personnel requirements and histocompatibility requirements, which, with minor exception, have not been updated since 1992. We are also seeking public comment regarding the flexibility to impose alternative sanctions for laboratories issued a Certificate of Waiver (CoW) determined to have participated in proficiency testing (PT) referral. In addition, we are seeking public comment related to appropriate sanctions in situations where we determine that a laboratory has referred its PT samples to another laboratory and has reported the other laboratory's result as their own.

This request for information also seeks public comment regarding the updating of fees for determination of program compliance and additional fees

for laboratories established under the CLIA regulations. We are also seeking public comment regarding the collection of other fees we are authorized to collect such as fees for revised certificates, post survey follow-up visits, complaint investigations, and activities related to imposition of sanctions.

We intend to consider public comments (including information such as evidence, research, and trends) received in response to this request for information when we draft proposals, in consultation, as appropriate, with the Centers for Disease Control and Prevention (CDC), to update the existing CLIA regulations through future rulemaking. We are also soliciting public comment on other areas of CLIA which should be reviewed and potentially updated.

DATES: To be assured consideration, comments must be received at one of the addresses provided below, no later than 5 p.m. on March 12, 2018.

ADDRESSES: In commenting, refer to file code CMS-3326-NC. Because of staff and resource limitations, we cannot accept comments by facsimile (FAX) transmission.

You may submit comments in one of four ways (please choose only one of the ways listed):

1. *Electronically.* You may submit electronic comments on this regulation to <http://www.regulations.gov>. Follow the "Submit a comment" instructions.

2. *By regular mail.* You may mail written comments to the following address ONLY: Centers for Medicare & Medicaid Services, Department of Health and Human Services, Attention: CMS-3326-NC, P.O. Box 8016, Baltimore, MD 21244-8016.

Please allow sufficient time for mailed comments to be received before the close of the comment period.

3. *By express or overnight mail.* You may send written comments to the following address ONLY: Centers for Medicare & Medicaid Services, Department of Health and Human Services, Attention: CMS-3326-NC, Mail Stop C4-26-05, 7500 Security Boulevard, Baltimore, MD 21244-1850.

4. *By hand or courier.* Alternatively, you may deliver (by hand or courier) your written comments ONLY to the following addresses:

a. For delivery in Washington, DC—Centers for Medicare & Medicaid Services, Department of Health and Human Services, Room 445-G, Hubert H. Humphrey Building, 200 Independence Avenue SW, Washington, DC 20201.

(Because access to the interior of the Hubert H. Humphrey Building is not