Regional Haze Progress Report
Implementation Plans; Alaska:
Approval and Promulgation of State
71—Region 10

PART 52—APPROVAL AND
PROMULGATION OF
IMPLEMENTATION PLANS

Cathy Stepp,
Regional Administrator, Region 5.
40 CFR part 52 is amended as follows:

1. The authority citation for part 52 continues to read as follows:
   Authority: 42 U.S.C. 7401 et seq.

2. In § 52.720, the table in paragraph (e) is amended by adding the entry
   for section to

Regional Haze Progress Report.

**EPA-APPROVED ILLINOIS NONREGULATORY AND QUASI-REGULATORY PROVISIONS**

<table>
<thead>
<tr>
<th>Name of SIP provision</th>
<th>Applicable geographic or nonattainment area</th>
<th>State submittal date</th>
<th>EPA approval date</th>
<th>Comments</th>
</tr>
</thead>
<tbody>
<tr>
<td>Regional Haze Progress Report</td>
<td>Statewide</td>
<td>02/01/17</td>
<td>April 12, 2018, [insert Federal Register citation]</td>
<td></td>
</tr>
</tbody>
</table>

**SUMMARY:** The Environmental Protection Agency (EPA) is approving a revision to the Alaska regional haze State Implementation Plan (SIP), submitted by the State of Alaska on March 10, 2016. Alaska submitted its Regional Haze Progress Report ("progress report") and a negative declaration stating that further revision of the existing regional haze SIP is not needed at this time. Alaska submitted both the progress report and the negative declaration in the form of implementation plan revisions as required by federal regulations. The progress report addresses the federal Regional Haze Rule requirements under the Clean Air Act to submit a report describing progress in achieving reasonable progress goals established for regional haze and a determination of the adequacy of the state’s existing plan addressing regional haze. We are also approving minor updates to the Enhanced Smoke Management Plan, Long-Term Strategy, and Commitment to Future 308 Plan Revision sections of the regional haze SIP, submitted concurrently with the progress report.

**DATES:** This final rule is effective May 14, 2018.

**ADDRESSES:** The EPA has established a docket for this action under Docket ID No. EPA–R10–OAR–2016–0749. All documents in the docket are listed on the [https://www.regulations.gov](https://www.regulations.gov) website. Although listed in the index, some information may not be publicly available, i.e., Confidential Business Information or other information the disclosure of which is restricted by statute. Certain other material, such as copyrighted material, is not placed on the internet and is publicly available only in hard copy form. Publicly available docket materials are available at [https://www.regulations.gov](https://www.regulations.gov) and at EPA Region 10, Office of Air and Waste, 1200 Sixth Avenue, Seattle, Washington 98101. The EPA requests that you contact the person listed in the FOR FURTHER INFORMATION CONTACT section to schedule your inspection. The Regional Office’s official hours of business are Monday through Friday, 8:30 to 4:30, excluding federal holidays.

**FOR FURTHER INFORMATION CONTACT:** Jeff Hunt, Air Planning Unit, Office of Air and Waste (OAW–150), EPA Region 10, 1200 Sixth Ave Suite 900, Seattle, WA 98101; telephone number: (206) 553–0256; email address: hunt.jeff@epa.gov.

**SUPPLEMENTARY INFORMATION:**

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I. Background Information
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**I. Background Information**

On February 16, 2018, the EPA proposed to approve Alaska’s Regional Haze Progress Report, as well as minor updates to the Enhanced Smoke Management Plan, Long-Term Strategy, and Commitment to Future 308 Plan Revision sections of the regional haze SIP, submitted concurrently with the Alaska Regional Haze Progress Report.

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1 We received two comments in support of the proposed approval. We also received five comments that were not germane to the regional haze program or the Alaska submission. See “AK RH 5 year progress Memo to File reComment” included in the docket for this action.
III. Statutory and Executive Orders

Review

Under the Clean Air Act, the Administrator is required to approve a SIP submission that complies with the provisions of the Clean Air Act and applicable federal regulations. 42 U.S.C. 7410(k); 40 CFR 52.02(a). Thus, in reviewing SIP submissions, the EPA’s role is to approve state choices, provided that they meet the criteria of the CAA. Accordingly, this action merely approves 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 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 actions such as 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 this action does not involve technical standards; and  
- does not provide the 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).  

The SIP is not approved to apply on any Indian reservation land and is also not approved to apply 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 as specified by Executive Order 13175 (65 FR 67249, November 9, 2000).  

The Congressional Review Act, 5 U.S.C. 801 et seq., as added by the Small Business Regulatory Enforcement Fairness Act of 1996, generally provides that before a rule may take effect, the agency promulgating the rule must submit a rule report, which includes a copy of the rule, to each House of the Congress and to the Comptroller General of the United States. The EPA will submit a report containing this action 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. A major rule cannot take effect until 60 days after it is published in the Federal Register. This action is not a “major rule” as defined by 5 U.S.C. 804(2).  

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 June 11, 2018. 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. 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 52

Environmental protection, Air pollution control, Carbon monoxide, Incorporation by reference, Intergovernmental relations, Lead, Nitrogen dioxide, Ozone, Particulate matter, Reporting and recordkeeping requirements, Sulfur oxides, Volatile organic compounds.


Chris Hladick,
Regional Administrator, Region 10.

For the reasons set forth in the preamble, 40 CFR part 52 is amended as follows:

PART 52—APPROVAL AND PROMULGATION OF IMPLEMENTATION PLANS

§ 52.70 Identification of plan.

(1) The authority citation for part 52 continues to read as follows:  
Authority: 42 U.S.C. 7401 et seq.

Subpart C—Alaska

§ 52.70 [Amended]

1. In §52.70, the table in paragraph (e) is amended by revising the entries for “II.III.K. Area Wide Pollutant Control Program for Regional Haze” and “III.III.K. Area Wide Pollutant Control Program for Regional Haze” to read as follows:

<table>
<thead>
<tr>
<th>§ 52.70 Identification of plan.</th>
</tr>
</thead>
<tbody>
<tr>
<td>II.III.K. Area Wide Pollutant Control Program for Regional Haze.</td>
</tr>
</tbody>
</table>

2. In § 52.70, the table in paragraph (e) is amended by adding a new entry for “Statewide .......................... 4/12/2018, ..........................” as follows:

<table>
<thead>
<tr>
<th>Name of SIP provision</th>
<th>Applicable geographic or nonattainment area</th>
<th>State submittal date</th>
<th>EPA approval date</th>
<th>Explanations</th>
</tr>
</thead>
<tbody>
<tr>
<td>Statewide ..........................</td>
<td>Statewide ..........................</td>
<td>3/10/2016</td>
<td>4/12/2018, ..........................</td>
<td>[Insert Federal Register citation]</td>
</tr>
</tbody>
</table>

EPA–APPROVED ALASKA NONREGULATORY PROVISIONS AND QUASI–REGULATORY MEASURES
ENVIRONMENTAL PROTECTION AGENCY

40 CFR Part 180


Clethodim; Pesticide Tolerances

AGENCY: Environmental Protection Agency (EPA).

ACTION: Final rule.

SUMMARY: This regulation establishes tolerances for residues of clethodim in or on multiple commodities which are identified and discussed later in this document. In addition, this regulation removes several previously established tolerances that are superseded by this final rule. Interregional Research Project Number 4 (IR–4) requested these tolerances under the Federal Food, Drug, and Cosmetic Act (FFDCA).

DATES: This regulation is effective April 12, 2018. Objections and requests for hearings must be received on or before June 11, 2018, 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–2016–0651, 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:
Michael L. Goodis, Registration Division (7505P), Office of Pesticide Programs, Environmental Protection Agency, 1200 Pennsylvania Ave. NW, Washington, DC 20460–0001; main telephone number: (703) 305–7090; email address: RDFRNotices@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?


C. How can I file an objection or hearing request?

Under FFDCA section 408(g), 21 U.S.C. 346a, 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–2016–0651 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 June 11, 2018. 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–2016–0651, 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.

For more information, see Unit II.B. of this rule.

II. Legal Authority

To ensure uniformity of the Official Regulations for the United States, this final rule is issued under the authority of 40 CFR part 180 and 7 U.S.C. 346 and 21 U.S.C. 346a, as amended by the Food, Drug, and Cosmetic Act (FFDCA).

III. Notice of Proposed Rulemaking

A. Objectors

On June 11, 2018, the Environmental Protection Agency (EPA) issued a Notice of Proposed Rulemaking (NPRM) (FR Doc. 2018–07520, 83 FR 20234, April 12, 2018) for the establishment of tolerances for residues of clethodim in or on multiple commodities. Copies of the NPRM are available from the docket identified above.

This regulation is effective April 12, 2018. Objections and requests for hearings must be received on or before June 11, 2018. Objections and requests for hearings must be in writing, and must be filed 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–2016–0651 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 June 11, 2018. Addresses for mail and hand delivery of objections and hearing requests are provided in 40 CFR 178.25(b).