

List of Subjects in 38 CFR Part 17

Administrative practice and procedure, Alcohol abuse, Alcoholism, Claims, Day care, Dental health, Drug abuse, Foreign relations, Government contracts, Grant programs—health, Grant programs—veterans, Health care, Health facilities, Health professions, Health records, Homeless, Medical and dental schools, Medical devices, Medical research, Mental health programs, Nursing homes, Philippines, Reporting and recordkeeping requirements, Scholarships and fellowships, Travel and transportation expenses, Veterans.

Dated: May 20, 2016.

Michael Shores,

Acting Director, Office of Regulation Policy & Management, Office of Secretary, Department of Veterans Affairs.

For the reasons set forth in the preamble, we propose to amend 38 CFR part 17 as follows:

PART 17—MEDICAL

- 1. The authority citation for part 17 continues to read as follows:

Authority: 38 U.S.C. 501, and as noted in specific sections.

- 2. Amend part 17 by adding an undesignated center heading and § 17.415 immediately after § 17.410 to read as follows:

Nursing Services**§ 17.415 Full practice authority for advanced practice registered nurses.**

(a) *Advanced practice registered nurse (APRN).* For purposes of this section, an advanced practice registered nurse (APRN) is an individual who:

- (1) Has completed a nationally-accredited, graduate-level educational program that prepares them for one of the four APRN roles of Certified Nurse Practitioner (CNP), Certified Registered Nurse Anesthetist (CRNA), Clinical Nurse Specialist (CNS), or Certified Nurse-Midwife (CNM);
- (2) Has passed a national certification examination that measures knowledge in one of the APRN roles described in paragraph (a)(1) of this section;
- (3) Has obtained a license from a State licensing board in one of four recognized APRN roles described in paragraph (a)(1) of this section; and
- (4) Maintains certification and licensure as required by paragraphs (a)(2) and (3) of this section.

(b) *Full practice authority.* For purposes of this section, full practice authority means the authority of an APRN to provide services described in paragraph (d) of this section without the

clinical oversight of a physician, regardless of State or local law restrictions, when that APRN is working within the scope of their VA employment.

(c) *Granting of full practice authority.* VA may grant full practice authority to an APRN subject to the following:

(1) Verification that the APRN meets the requirements established in paragraph (a) of this section; and

(2) Determination that the APRN has demonstrated the knowledge and skills necessary to provide the services described in paragraph (d) of this section without the clinical oversight of a physician, and is thus qualified to be privileged for such scope of practice.

(d) *Services provided by an APRN with full practice authority.* (1) Subject to the limitations established in paragraph (d)(2) of this section, the full practice authority for each of the four APRN roles includes, but is not limited to, providing the following services:

(i) A CNP has full practice authority to:

(A) Take comprehensive histories, provide physical examinations and other health assessment and screening activities, diagnose, treat, and manage patients with acute and chronic illnesses and diseases;

(B) Order, perform, supervise, and interpret laboratory and imaging studies;

(C) Prescribe medication and durable medical equipment;

(D) Make appropriate referrals for patients and families, and request consultations;

(E) Aid in health promotion, disease prevention, health education, and counseling as well as the diagnosis and management of acute and chronic diseases.

(ii) A CRNA has full practice authority to:

(A) Plan and initiate anesthetic techniques (general, regional, local) and sedation;

(B) Provide post-anesthesia evaluation and discharge;

(C) Order and evaluate diagnostic tests;

(D) Request consultations;

(D) Perform point-of-care testing; and

(E) Respond to emergency situations for airway management.

(iii) A CNS has full practice authority to provide diagnosis and treatment of health or illness states, disease management, health promotion, and prevention of illness and risk behaviors among individuals, families, groups, and communities within their scope of practice.

(iv) A CNM has full practice authority to provide a range of primary health

care services to women, including gynecologic care, family planning services, preconception care (care that women veterans receive before becoming pregnant, including reducing the risk of birth defects and other problems such as the treatment of diabetes and high blood pressure), prenatal and postpartum care, childbirth, and care of a newborn, and treating the partner of their female patients for sexually transmitted disease and reproductive health, if the partner is also enrolled in the VA healthcare system or is not required to enroll.

(2) The full practice authority of an APRN is subject to the limitations imposed by the Controlled Substances Act, 21 U.S.C. 801 *et seq.*, and that APRN's State licensure on the authority to prescribe, or administer controlled substances, as well as any other limitations on the provision of VA care set forth in applicable Federal law and policy.

(e) *Preemption of State and local law.* To achieve important Federal interests, including but not limited to the ability to provide the same comprehensive care to veterans in all States under 38 U.S.C. 7301, this section preempts conflicting State and local laws relating to the practice of APRNs when such APRNs are working within the scope of their VA employment. Any State or local law, or regulation pursuant to such law, is without any force or effect on, and State or local governments have no legal authority to enforce them in relation to this section or decisions made by VA under this section.

(Authority: 38 U.S.C. 7301, 7304, 7402, and 7403)

[FR Doc. 2016-12338 Filed 5-24-16; 8:45 am]

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ENVIRONMENTAL PROTECTION AGENCY**40 CFR Part 52**

[EPA-R01-OAR-2014-0364; A-1-FRL-9936-62-Region 1]

Air Plan Approval; Connecticut; Sulfur Content of Fuel Oil Burned in Stationary Sources

AGENCY: Environmental Protection Agency (EPA).

ACTION: Proposed rule.

SUMMARY: The Environmental Protection Agency (EPA) is proposing to approve a State Implementation Plan (SIP) revision submitted by the State of Connecticut on April 22, 2014, with supplemental submittals on June 18,

2015 and September 25, 2015. This revision establishes sulfur in fuel oil content limits for use in stationary sources. In addition, the submittal includes minor clarifying revisions to the methods for sampling, emission testing, sample analysis, and reporting. The intended effect of this action is to propose approval of these requirements into the Connecticut SIP. This action is being taken under the Clean Air Act.

DATES: Written comments must be received on or before June 24, 2016.

ADDRESSES: Submit your comments, identified by Docket ID Number EPA–R01–OAR–2014–0364 by one of the following methods:

1. *www.regulations.gov*: Follow the on-line instructions for submitting comments.

2. *Email*: arnold.anne@epa.gov.

3. *Fax*: (617) 918–0047.

4. *Mail*: “Docket Identification Number EPA–R01–OAR–2014–0364, Anne Arnold, U.S. Environmental Protection Agency, EPA New England Regional Office, Office of Ecosystem Protection, Air Quality Planning Unit, 5 Post Office Square—Suite 100, (Mail code OEP05–2), Boston, MA 02109–3912.

5. *Hand Delivery or Courier*: Deliver your comments to: Anne Arnold, Manager, Air Quality Planning Unit, U.S. Environmental Protection Agency, EPA New England Regional Office, Office of Ecosystem Protection, Air Quality Unit, 5 Post Office Square—Suite 100, (mail code OEP05–2), Boston, MA 02109–3912. Such deliveries are only accepted during the Regional Office’s normal hours of operation. The Regional Office’s official hours of business are Monday through Friday, 8:30 a.m. to 4:30 p.m., excluding legal holidays.

Instructions: Direct your comments to Docket ID No. EPA–R01–OAR–2014–0364. EPA’s policy is that all comments received will be included in the public docket without change and may be made available online at *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 through *www.regulations.gov*, or email, information that you consider to be CBI or otherwise protected. The *www.regulations.gov* Web site is an “anonymous access” system, which means 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 EPA without going through *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, 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 EPA cannot read your comment due to technical difficulties and cannot contact you for clarification, 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.

Docket: All documents in the electronic docket are listed in the *www.regulations.gov* index. Although listed in the index, some information is not publicly available, *i.e.*, CBI or other information whose disclosure is restricted by statute. Certain other material, such as copyrighted material, is not placed on the Internet and will be publicly available only in hard copy form. Publicly available docket materials are available either electronically in *www.regulations.gov* or in hard copy at U.S. Environmental Protection Agency, EPA New England Regional Office, Office of Ecosystem Protection, Air Quality Planning Unit, 5 Post Office Square—Suite 100, Boston, MA. EPA requests that if at all possible, you contact the contact 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 a.m. to 4:30 p.m., excluding legal holidays.

In addition, copies of the state submittal are also available for public inspection during normal business hours, by appointment at the State Air Agency; Bureau of Air Management, Department of Energy and Environmental Protection, State Office Building, 79 Elm Street, Hartford, CT 06106–1630.

FOR FURTHER INFORMATION CONTACT:

Anne K. McWilliams, Air Quality Planning Unit, U.S. Environmental Protection Agency, New England Regional Office, 5 Post Office Square—Suite 100, (Mail code OEP05–2), Boston, MA 02109–3912, telephone (617) 918–1697, facsimile (617) 918–0697, email mcwilliams.anne@epa.gov.

SUPPLEMENTARY INFORMATION: In the Final Rules Section of this **Federal Register**, EPA is approving the State’s SIP submittals as a direct final rule without prior proposal because the Agency views this as a noncontroversial

submittal and anticipates no adverse comments. A detailed rationale for the approval is set forth in the direct final rule. If no adverse comments are received in response to this rule, no further activity is contemplated. If EPA receives 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 rule. EPA will not institute a second comment period. Any parties interested in commenting on this action should do so at this time. Please note that if EPA receives adverse comment on an amendment, paragraph, or section of this rule and if that provision may be severed from the remainder of the rule, EPA may adopt as final those provisions 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: November 5, 2015.

H. Curtis Spalding,

Regional Administrator, EPA New England.

[FR Doc. 2016–12118 Filed 5–24–16; 8:45 am]

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ENVIRONMENTAL PROTECTION AGENCY

40 CFR Parts 52 and 81

[EPA–R06–OAR–2015–0721; FRL–9946–85–Region 6]

Clean Air Act Redesignation Substitute for the Dallas-Fort Worth 1-Hour Ozone and 1997 8-Hour Ozone Nonattainment Areas; Texas

AGENCY: Environmental Protection Agency (EPA).

ACTION: Proposed rule.

SUMMARY: The Environmental Protection Agency (EPA) is proposing to approve a redesignation substitute and make a finding of attainment for both the 1-hour ozone and the 1997 8-hour ozone National Ambient Air Quality Standards (NAAQS) for the Dallas-Fort Worth nonattainment area (DFW area). The redesignation substitute demonstration states that the area has attained both the revoked 1-hour ozone and the revoked 1997 8-hour ozone NAAQS due to permanent and enforceable emission reductions, and that it will maintain those NAAQS for ten years from the date of the EPA’s approval of this demonstration. Final approval of the redesignation substitute will result in the area no longer being subject to any remaining applicable anti-backsliding requirements and the nonattainment