

(v) Veterans will be disenrolled, and reenrolled, in the order of the priority categories listed with veterans in priority category 1 being the last to be disenrolled and the first to be reenrolled. Similarly, within priority categories 7 and 8, veterans will be disenrolled, and reenrolled, in the order of the priority subcategories listed with veterans in subcategory (i) being the last to be disenrolled and first to be reenrolled.

* * * * *

(5) *Disenrollment.* A veteran enrolled in the VA health care system under paragraph (d)(2) or (d)(4) of this section will be disenrolled only if:

(i) The veteran submits to a VA medical center or the VA Health Eligibility Center, 1644 Tullie Circle, Atlanta, Georgia 30329, a signed document stating that the veteran no longer wishes to be enrolled; or

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(Authority: 38 U.S.C 101, 501, 1521, 1701, 1705, 1710, 1721, 1722).

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DEPARTMENT OF VETERANS AFFAIRS

38 CFR Part 36

RIN 2900-AF00

Schedule for Rating Disabilities; The Skin

AGENCY: Department of Veterans Affairs.

ACTION: Final rule; correction.

SUMMARY: In a document published in the *Federal Register* on July 31, 2002, (67 FR 49590), we amended that portion of the Department of Veterans Affairs (VA) Schedule for Rating Disabilities that addresses the skin. The document contains an error in the Supplementary Information portion of the preamble. That error consists of an incorrect restatement of regulatory text. This document corrects that error.

DATES: *Effective Date:* This correction is effective July 31, 2002.

FOR FURTHER INFORMATION CONTACT: Carroll McBrine, M.D., Consultant, Policy and Regulations Staff (211B), Compensation and Pension Service, Veterans Benefits Administration, Department of Veterans Affairs, 810 Vermont Avenue NW., Washington, DC 20420, (202) 273-7230.

SUPPLEMENTARY INFORMATION: In rule FR Doc. 02-19331, published on July 31, 2002 (67 FR 49590), on page 49595, in column 1, the first paragraph, the phrase "a 30-percent evaluation calls for

recurrent debilitating episodes at least four times during the past 12-month period despite ongoing immunosuppressive therapy" is corrected to read "a 30-percent evaluation calls for recurrent debilitating episodes at least four times during the past 12-month period, and requiring intermittent systemic immunosuppressive therapy."

Approved: October 1, 2002.

Roland Halstead,

Acting Director, Office of Regulatory Law.

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ENVIRONMENTAL PROTECTION AGENCY

40 CFR Part 52

[IA 154-1154a; FRL-7392-6]

Approval and Promulgation of Implementation Plans; State of Iowa

AGENCY: Environmental Protection Agency (EPA).

ACTION: Direct final rule.

SUMMARY: EPA is announcing it is approving revisions to the Iowa State Implementation Plan (SIP). The SIP revisions, regarding the state's construction permitting rules as they pertain to industrial anaerobic lagoons and anaerobic lagoons for animal feeding operations in Iowa, will help ensure Federal enforceability of the state's air program.

DATES: This direct final rule will be effective December 9, 2002, unless EPA receives adverse comments by November 8, 2002. If adverse comments are received, EPA will publish a timely withdrawal of the direct final rule in the *Federal Register* informing the public that the rule will not take effect.

ADDRESSES: Comments may be mailed to Lynn M. Slugantz, Environmental Protection Agency, Air Planning and Development Branch, 901 North 5th Street, Kansas City, Kansas 66101.

Copies of documents relative to this action are available for public inspection during normal business hours at the above-listed Region 7 location. The interested persons wanting to examine these documents should make an appointment with the office at least 24 hours in advance.

FOR FURTHER INFORMATION CONTACT: Lynn M. Slugantz at (913) 551-7883.

SUPPLEMENTARY INFORMATION: Throughout this document whenever "we," "us," or "our" is used, we mean EPA. This section provides additional

information by addressing the following questions:

What is an SIP?

What is the Federal approval process for an SIP?

What does Federal approval of a state regulation mean to me?

What is being addressed in this action?

Have the requirements for approval of an SIP revision been met?

What action is EPA taking?

What Is a SIP?

Section 110 of the Clean Air Act (CAA) requires states to develop air pollution regulations and control strategies to ensure that state air quality meets the national ambient air quality standards established by EPA. These ambient standards are established under section 109 of the CAA, and they currently address six criteria pollutants. These pollutants are: carbon monoxide, nitrogen dioxide, ozone, lead, particulate matter, and sulfur dioxide.

Each state must submit these regulations and control strategies to us for approval and incorporation into the Federally-enforceable SIP.

Each Federally-approved SIP protects air quality primarily by addressing air pollution at its point of origin. These SIPs can be extensive, containing state regulations or other enforceable documents and supporting information such as emission inventories, monitoring networks, and modeling demonstrations.

What Is the Federal Approval Process for a SIP?

In order for state regulations to be incorporated into the Federally-enforceable SIP, states must formally adopt the regulations and control strategies consistent with state and Federal requirements. This process generally includes a public notice, public hearing, public comment period, and a formal adoption by a state-authorized rulemaking body.

Once a state rule, regulation, or control strategy is adopted, the state submits it to us for inclusion into the SIP. We must provide public notice and seek additional public comment regarding the proposed Federal action on the state submission. If adverse comments are received, they must be addressed prior to any final Federal action by us.

All state regulations and supporting information approved by EPA under section 110 of the CAA are incorporated into the Federally-approved SIP. Records of such SIP actions are maintained in the Code of Federal Regulations (CFR) at Title 40, part 52, entitled "Approval and Promulgation of