have a significant economic impact on a substantial number of small entities.

DEA proposes placing the substances DOI and DOC (chemical names: 2,5-dimethoxy-4-iodoamphetamine [DOI] and 2,5-dimethoxy-4-chloroamphetamine [DOC]), including their salts, isomers, and salts of isomers whenever the existence of such salts, isomers, and salts of isomers is possible within the specific chemical designation, in schedule I of the CSA. This action is being taken, in part, to enable the United States to meet its obligations under the 1971 Convention for DOC. If finalized, this action would impose the regulatory controls and administrative, civil, and criminal sanctions applicable to schedule I controlled substances on persons who handle (manufacture, distribute, reverse distribute, import, export, engage in research, conduct instructional activities or chemical analysis with, or possess), or propose to handle DOI and DOC.

According to HHS, and also by DEA's findings in this proposed rule, DOI and DOC have a high potential for abuse, have no currently accepted medical use in treatment in the United States, and lack accepted safety for use under medical supervision. There appear to be no legitimate sources for DOI and DOC as marketed drugs in the United States, but DEA notes that these substances are available for purchase from legitimate suppliers for scientific research. There is no evidence of significant diversion of DOI and DOC from legitimate suppliers. As such, the proposed rule will not, if promulgated, result in a significant effect on a substantial number of small entities.

Unfunded Mandates Reform Act of 1995

On the basis of information contained in the ''Regulatory Flexibility Act'' section above, DEA has determined pursuant to the Unfunded Mandates Reform Act (UMRA) of 1995 (2 U.S.C. 1501 *et seq.*) that this proposed action would not result in any Federal mandate that may result ''in the expenditure by State, local, and tribal governments, in the aggregate, or by the private sector, of $100,000,000 or more (adjusted annually for inflation) in any 1 year . . . .'' Therefore, neither a Small Government Agency Plan nor any other action is required under UMRA of 1995.

List of Subjects in 21 CFR Part 1308

Administrative practice and procedure, Drug traffic control, Reporting and recordkeeping requirements.

For the reasons set out above, 21 CFR part 1308 is proposed to be amended to read as follows:

PART 1308—SCHEDULES OF CONTROLLED SUBSTANCES

■ 1. The authority citation for 21 CFR part 1308 continues to read as follows:

Authority: 21 U.S.C. 811, 812, 871(b), 956(b), unless otherwise noted.

■ 2. In § 1308.11, as proposed to be amended at 86 FR 16553 (March 30, 2021), 86 FR 37719 (July 16, 2021), 86 FR 69187 (December 7, 2021), and 87 FR 2383 (January 14, 2022), add paragraphs (d)(106) and (107) to read as follows:

§ 1308.11 Schedule I.

* * * * *

(d) * * *

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Anne Milgram,

Administrator.

[FR Doc. 2022–07648 Filed 4–8–22; 8:45 am]

BILLING CODE 4410–09–P

POSTAL REGULATORY COMMISSION

39 CFR Part 3010

[Docket No. RM2022–4; Order No. 6141]

RIN 3211–AA31

Rules of Practice and Procedure

AGENCY: Postal Regulatory Commission.

ACTION: Proposed rulemaking.

SUMMARY: The Commission is proposing to add rules which revise the Commission's rules of practice and procedure regarding notices, motions, and information requests. The Commission invites public comment on the proposed rules.

DATES: *Comments are due:* May 26, 2022.

ADDRESSES: For additional information, Order No. 6141 can be accessed electronically through the Commission's website at *https://www.prc.gov.* Submit comments electronically via the Commission's Filing Online system at *https://www.prc.gov.* Those who cannot submit comments electronically should contact the person identified in the **FOR FURTHER INFORMATION CONTACT** section by telephone for advice on filing alternatives.

FOR FURTHER INFORMATION CONTACT: David A. Trissell, General Counsel, at 202–789–6820.

SUPPLEMENTARY INFORMATION:

Table of Contents

I. Relevant Statutory Requirements

Pursuant to 39 U.S.C. 503, the Commission establishes this rulemaking docket to propose amendments to the Commission's rules of practice and procedure regarding notices, motions, and information requests.

II. Background

The proposed amendments provide rules relating to motions for reconsideration of final Commission orders.[1]

[1] Motions for review of other Commission determinations may be filed in accordance with 39 CFR 3010.160.

III. Basis and Purpose of Proposed Rules

The proposed amendments revise the Commission's rules on notices, motions, and information requests, within its rules of practice and procedure, to provide rules specific to motions for reconsideration. The proposed amendments reflect the Commission's current practice of hearing timely motions for reconsideration of its final orders.

Under the proposed rules, any party may file a motion for reconsideration of a Commission final order within 15 days of the issuance of the order subject to the motion. All motions for reconsideration must briefly and specifically allege material errors of fact or law, and the relief sought, and must be confined to new questions raised by the determination or action ordered and upon which the moving party had no prior opportunity to submit arguments. Finally, no motion for reconsideration shall stay the effect of an order of the Commission unless the Commission orders otherwise.

IV. Proposed Rules

Proposed § 3010.165(a). Proposed § 3010.165(a) is added to explain eligibility among parties for filing motions for reconsideration.

Proposed § 3010.165(b). Proposed § 3010.165(b) is added to clarify the timing and content requirements for motions for reconsideration.

Proposed § 3010.165(c). Proposed § 3010.165(c) is added to explain that motions for reconsideration do not, on their own, stay the effect of the underlying order.

List of Subjects in 39 CFR Part 3010

Administrative practice and procedure, Confidential business information, Freedom of information, Sunshine Act.

Erica A. Barker,

Secretary.

For the reasons stated in the preamble, the Commission proposes to amend chapter III of title 39 of the Code of Federal Regulations as follows:

PART 3010—RULES OF PRACTICE AND PROCEDURE

■ 1. The authority citation for part 3010 continues to read as follows:

Authority: 39 U.S.C. 404(d); 503; 504; 3661.

■ 2. Add § 3010.165 to read as follows:

§ 3010.165 Motions for reconsideration.

(a) Any person may file a motion requesting reconsideration of a final order by the Commission.

(b) The motion shall be filed within 15 days of the issuance of the final order that is the subject of the motion and must:

(1) Briefly and specifically allege material errors of fact or law and the relief sought; and

(2) Be confined to new questions raised by the determination or action ordered and upon which the moving party had no prior opportunity to submit arguments.

(c) Unless the Commission orders otherwise, the filing of a motion for reconsideration shall not stay the effect of an order of the Commission.

[FR Doc. 2022–07725 Filed 4–8–22; 8:45 am]

BILLING CODE 7710–FW–P

ENVIRONMENTAL PROTECTION AGENCY

40 CFR Part 52

[EPA–R08–OAR–2021–0807; FRL–9680–01–R8]

Approval and Promulgation of Implementation Plans; South Dakota; Revisions to South Dakota Codified Law and Administrative Rules of South Dakota

AGENCY: Environmental Protection Agency (EPA).

ACTION: Proposed rule.

SUMMARY: The Environmental Protection Agency (EPA or the ''Agency'') is proposing approval of South Dakota's submittal requesting that EPA recognize the merger of South Dakota's Department of Agriculture (DOA) with the Department of Environment and Natural Resources (DENR) to form the new Department of Agriculture and Natural Resources (DANR) and incorporate corresponding non-substantive revisions to the South Dakota Codified Law (SDCL) and the Administrative Rules of South Dakota (ARSD) into South Dakota's State Implementation Plan. Accordingly, EPA is proposing to approve South Dakota's submittal in accordance with the Clean Air Act (CAA).

DATES: Written comments must be received on or before May 11, 2022.

ADDRESSES: Submit your comments, identified by Docket ID No. EPA–R08–OAR–2021–0807, to the Federal Rulemaking Portal: *https://www.regulations.gov.* Follow the online instructions for submitting comments. Once submitted, comments cannot be edited or removed from *www.regulations.gov.* 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, the full EPA public comment policy, information about CBI or multimedia submissions, and general guidance on making effective comments, please visit *https://www.epa.gov/dockets/commenting-epa-dockets.*

Docket: All documents in the docket are listed in the *www.regulations.gov* index. Although listed in the index, some information is not publicly available, *e.g.,* CBI or other information whose disclosure is restricted by statute. Certain other material, such as copyrighted material, will be publicly available only in hard copy. Publicly available docket materials are available electronically in *www.regulations.gov.* To reduce the risk of COVID–19 transmission, for this action we do not plan to offer hard copy review of the docket. Please email or call the person listed in the **FOR FURTHER INFORMATION CONTACT** section if you need to make alternative arrangements for access to the docket.

FOR FURTHER INFORMATION CONTACT: Kate Gregory, Air and Radiation Division, U.S. Environmental Protection Agency (EPA), Region 8, Mail Code 8P–ARD–QP, 1595 Wynkoop Street, Denver, Colorado 80202–1129, telephone number: (303) 312–6175, email address: *gregory.kate@epa.gov.*

SUPPLEMENTARY INFORMATION: Throughout this document ''we,'' ''us,'' and ''our'' means EPA. In the Final Rules section of this **Federal Register**, EPA is approving the State's SIP submittal 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 relevant adverse comments are received in response to this rule, no further activity is contemplated. If EPA receives such 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 issue of the **Federal Register**.

List of Subjects in 40 CFR Part 52

Environmental protection, Air pollution control, Carbon monoxide, Greenhouse gases, Incorporation by reference, Intergovernmental relations, Lead, Nitrogen dioxide, Ozone,