• *Federal eRulemaking Portal: http://www.regulations.gov*. Follow the instructions for submitting comments. Comments submitted electronically, including attachments, to *http://www.regulations.gov* will be posted to the docket unchanged. Because your comment will be made public, you are solely responsible for ensuring that your comment does not include any confidential information that you or a third party may not wish to be posted, such as medical information, your or anyone else's Social Security number, or confidential business information, such as a manufacturing process. Please note that if you include your name, contact information, or other information that identifies you in the body of your comments, that information will be posted on *http://www.regulations.gov*.

• If you want to submit a comment with confidential information that you do not wish to be made available to the public, submit the comment as a written/paper submission and in the manner detailed (see ''Written/Paper Submissions'' and ''Instructions'').

Written/Paper Submissions

Submit written/paper submissions as follows:

• *Mail/Hand delivery/Courier (for written/paper submissions):* Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

• For written/paper comments submitted to the Division of Dockets Management, FDA will post your comment, as well as any attachments, except for information submitted, marked and identified, as confidential, if submitted as detailed in ''Instructions.''

Instructions: All submissions received must include the Docket No. FDA–2016–N–1111 for ''Proposal to Ban Electrical Stimulation Devices Used to Treat Self-Injurious or Aggressive Behavior.'' Received comments will be placed in the docket and, except for those submitted as ''Confidential Submissions,'' publicly viewable at *http://www.regulations.gov* or at the Division of Dockets Management between 9 a.m. and 4 p.m., Monday through Friday.

• *Confidential Submissions*—To submit a comment with confidential information that you do not wish to be made publicly available, submit your comments only as a written/paper submission. You should submit two copies total. One copy will include the information you claim to be confidential with a heading or cover note that states ''THIS DOCUMENT CONTAINS CONFIDENTIAL INFORMATION.'' The Agency will review this copy, including the claimed confidential information, in its consideration of comments. The second copy, which will have the claimed confidential information redacted/blacked out, will be available for public viewing and posted on *http://www.regulations.gov*. Submit both copies to the Division of Dockets Management. If you do not wish your name and contact information to be made publicly available, you can provide this information on the cover sheet and not in the body of your comments and you must identify this information as ''confidential.'' Any information marked as ''confidential'' will not be disclosed except in accordance with 21 CFR 10.20 and other applicable disclosure law. For more information about FDA's posting of comments to public dockets, see 80 FR 56469, September 18, 2015, or access the information at: *http://www.fda.gov/regulatoryinformation/dockets/default.htm*.

Docket: For access to the docket to read background documents or the electronic and written/paper comments received, go to *http://www.regulations.gov* and insert the docket number, found in brackets in the heading of this document, into the ''Search'' box and follow the prompts and/or go to the Division of Dockets Management, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

FOR FURTHER INFORMATION CONTACT: Rebecca Nipper, Center for Devices and Radiological Health, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 66, Rm. 1540, Silver Spring, MD 20993–0002, 301–796–6527.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of April 25, 2016, FDA published a proposed rule with a 30-day comment period to request comments on a proposal to ban ESDs used for SIB or AB. Comments on the proposed ban will inform FDA's rulemaking.

The Agency has received requests for a 60-day extension of the comment period for the proposed rule. Each request conveyed concern that the current 30-day comment period does not allow sufficient time to develop a meaningful or thoughtful response to the proposed rule.

FDA has considered the requests and is extending the comment period for the proposed rule for 60 days, until July 25, 2016. The Agency believes that a 60-day extension allows adequate time for interested persons to submit comments without significantly delaying rulemaking on this important issue.

Dated: May 17, 2016.

Leslie Kux,

Associate Commissioner for Policy.

[FR Doc. 2016–12026 Filed 5–20–16; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF THE INTERIOR

Bureau of Ocean Energy Management

30 CFR Part 550

[Docket ID: BOEM–2013–0081]

RIN 1010–AD82

Air Quality Control, Reporting, and Compliance

AGENCY: Bureau of Ocean Energy Management (BOEM), Interior.

ACTION: Proposed rule; notice of extension of public comment period.

SUMMARY: BOEM is extending the public comment period to submit comments on the proposed rule entitled ''Air Quality Control, Reporting, and Compliance,'' which was published in the **Federal Register** on April 5, 2016. The original public comment period to submit comments on this rulemaking would have ended on June 6, 2016. However, BOEM has received public comments requesting an extension of the comment period. BOEM has reviewed the extension requests and has determined that a 14-day comment period extension to June 20, 2016, is appropriate. The proposed rule specified a separate, shorter period to submit comments to the Office of Management and Budget on the information collection (IC) burden in this rulemaking. That comment period ended on May 5, 2016, and will not be extended.

DATES: The comment period for comments on the substance of the proposed rule published on April 5, 2016 (81 FR 19717), has been extended. Written comments must be received by the extended due date of June 20, 2016. BOEM may not fully consider comments received after this date.

ADDRESSES: You may submit comments identified by the number 1010–AD82, by any of the following methods:

• *Federal rulemaking portal: http://www.regulations.gov.* Follow the instructions for submitting comments.

• *Mail:* Department of the Interior, Bureau of Ocean Energy Management, Office of Policy, Regulation and Analysis, Attention: Peter Meffert, 45600 Woodland Road, Sterling, Virginia 20166.

• *Hand delivery:* Front Desk, Department of the Interior, Bureau of

Ocean Energy Management, Office of Policy, Regulation and Analysis, Attention: Peter Meffert, 45600 Woodland Road, Sterling, Virginia 20166.

• *Public Availability of Comments:* BOEM does not consider anonymous comments; please include your name and address as part of your submittal. Before including your name, address, phone number, email address, or other personally identifiable information in your comment, you should be aware that your entire comment—including your personally identifiable information—may be made publicly available at any time. While you can ask us in your comment to withhold your personally identifiable information from public review, we cannot guarantee that we will be able to do so.

FOR FURTHER INFORMATION CONTACT: Peter Meffert, Bureau of Ocean Energy Management, Office of Policy, Regulation and Analysis, at *peter.meffert@boem.gov* or mail to 45600 Woodland Road, Sterling, Virginia 20166; or call (703) 787–1610.

SUPPLEMENTARY INFORMATION: BOEM published a proposed rule on Air Quality Control, Reporting, and Compliance on April 5, 2016. The proposed rule is intended to revise and replace BOEM's air quality regulations with a new set of regulations that reflect a number of policy changes with respect to the existing air quality regulatory program. The key policy changes in the proposed rule relate to: (1) Fulfilling BOEM's statutory responsibility under section 5(a)(8) of the Outer Continental Shelf Lands Act by addressing all relevant criteria and major precursor air pollutants and by cross-referencing the ambient air quality standards and benchmarks (AAQSB) for those pollutants to those of the U.S. Environmental Protection Agency; (2) formalizing the concept and application of the term ''attributed emissions''; (3) changing the methods for determining the locations from which air emissions will be measured and evaluated; (4) modifying the process by which emission exemption threshold (EETs) are established and updated; (5) changing the circumstances when emission reduction measures (ERM), including Best Available Control Technology (BACT), are required, and establishing new criteria for the application of ERM; (6) revising the boundary at which BOEM determines air quality compliance to the State seaward boundary (SSB), rather than the coastline; (7) formalizing requirements for the consolidation of emissions from multiple facilities; (8) consistent with BOEM's existing regulatory authority, articulating a schedule for ensuring that plans, including previously approved plans, will be compliant with these updated regulations; (9) adding an air quality component to the submission of right-of-use and easement, right-of-way, and lease term pipeline applications; (10) expanding use of the offsets as an alternative in circumstances where BACT was previously required; and (11) adding a new requirement for all plans to be reviewed at least every 10 years, to ensure ongoing compliance with the National Ambient Air Quality Standards (NAAQS), as amended from time to time.

After publication of the proposed rule, BOEM received public comments requesting an extension.

On March 17, 2016, BOEM issued a press release to notify the public that the proposed rule would be issued and provided an internet link to an advance copy of the proposed rule. On April 5, 2016, BOEM issued the proposed rule with a requirement to submit comments on the substance of this rulemaking by June 6, 2016. BOEM is extending the comment period to June 20, 2016. With more than 90 days of public inspection, BOEM has concluded that interested parties will have sufficient opportunity to analyze the proposed rule and provide comment. Accordingly, written comments on the substance of this rulemaking must be submitted by the extended due date of June 20, 2016.

The proposed rule specified a separate, shorter period to submit comments to the Office of Management and Budget on the information collection (IC) burden in this rulemaking. That comment period ended on May 5, 2016, and will not be extended.

Dated: May 17, 2016.

Amanda C. Leiter,

Acting Assistant Secretary—Land and Minerals Management.

[FR Doc. 2016–12099 Filed 5–20–16; 8:45 am]

BILLING CODE 4310–MR–P

ENVIRONMENTAL PROTECTION AGENCY

40 CFR Part 52

[EPA–R01–OAR–2012–0289; FRL–9946–68–Region 1]

Air Plan Approval; New Hampshire; Ozone Maintenance Plan

AGENCY: Environmental Protection Agency.

ACTION: Proposed rule.

SUMMARY: The Environmental Protection Agency (EPA) is proposing to approve a State Implementation Plan (SIP) revision submitted by the State of New Hampshire that contains an ozone maintenance plan for New Hampshire's former 1-hour ozone nonattainment areas. The Clean Air Act requires that areas that are designated attainment for the 1997 8-hour ozone standard, and also had been previously designated either nonattainment or maintenance for the 1-hour ozone standard, develop a plan showing how the state will maintain the ozone standard for the area. The intended effect of this action is to propose approval of New Hampshire's maintenance plan. This action is being taken in accordance with the Clean Air Act.

DATES: Written comments must be received on or before June 22, 2016.

ADDRESSES: Submit your comments, identified by Docket ID No. EPA–R01–OAR–2012–0289 at *http://www.regulations.gov,* or via email to *arnold.anne@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, the 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. The 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 *http://www2.epa.gov/dockets/commenting-epa-dockets.*

FOR FURTHER INFORMATION CONTACT: Anne Arnold, Air Quality Planning Unit, U.S. Environmental Protection Agency, Suite 100, Mail Code OEP05–02, Boston, MA 02109–3912, telephone number (617) 918–1047, fax number (617) 918–0047, email *arnold.anne@epa.gov*.

SUPPLEMENTARY INFORMATION: In the Final Rules Section of this **Federal**