include a general description of the additional component(s) or additional scenario(s) and the basis for requiring the company to include the additional component(s) or additional scenario(s).

* * * * *

(iii) *Description of component.* The Board will respond in writing within 14 calendar days of receipt of the company's request. The Board will provide the covered company with a description of any additional component(s) or additional scenario(s) by March 1 of the calendar year in which the stress test is performed pursuant to this section.

■ 11. Section 252.55 is amended by revising paragraphs (a), (b)(4)(i), and (b)(4)(iii) to read as follows:

§ 252.55 Mid-cycle stress test.

(a) *Mid-cycle stress test requirement.* In addition to the stress test required under § 252.54, a covered company must conduct a mid-cycle stress test. The stress test must be conducted by September 30 of each calendar year based on data as of June 30 of that calendar year, unless the time or the as-of date is extended by the Board in writing.

(b) * * *

(4) *Notice and response*—(i) *Notification of additional component.* If the Board requires a covered company to include one or more additional components in its adverse and severely adverse scenarios under paragraph (b)(2) of this section or one or more additional scenarios under paragraph (b)(3) of this section, the Board will notify the company in writing. The Board will provide such notification no later than June 30. The notification will include a general description of the additional component(s) or additional scenario(s) and the basis for requiring the company to include the additional component(s) or additional scenario(s).

* * * * *

(iii) *Description of component.* The Board will provide the covered company with a description of any additional component(s) or additional scenario(s) by September 1 of the calendar year prior to the year in which the stress test is performed pursuant to this section.

■ 12. Section 252.57 is amended by revising paragraph (a) to read as follows:

§ 252.57 Reports of stress test results.

(a) *Reports to the Board of stress test results.* (1) A covered company must report the results of the stress test required under § 252.54 to the Board in the manner and form prescribed by the Board. Such results must be submitted by April 5 of the calendar year in which the stress test is performed pursuant to 12 CFR 252.54, unless that time is extended by the Board in writing.

(2) A covered company must report the results of the stress test required under § 252.55 to the Board in the manner and form prescribed by the Board. Such results must be submitted by October 5 of the calendar year in which the stress test is performed pursuant to 12 CFR 252.55, unless that time is extended by the Board in writing.

* * * * *

■ 13. Section 252.58 is amended by revising paragraph (a)(1)(ii) to read as follows:

§ 252.58 Disclosure of stress test results.

(a) * * *

(1) * * *

(ii) A covered company must publicly disclose a summary of the results of the stress test required under § 252.55. This disclosure must occur in the period beginning on October 5 and ending on November 4 of the calendar year in which the stress test is performed pursuant to 12 CFR 252.55, unless that time is extended by the Board in writing.

* * * * *

By order of the Board of Governors of the Federal Reserve System, September 26, 2016.

Robert deV. Frierson,

Secretary of the Board.

[FR Doc. 2016–23629 Filed 9–29–16; 8:45 am]

BILLING CODE 6210–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 573

[Docket No. FDA–2014–F–0988]

BASF Corp.; Filing of Food Additive Petition (Animal Use)

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of petition.

SUMMARY: The Food and Drug Administration (FDA) is announcing that BASF Corp., as a part of their petition (FAP 2286) proposing that the food additive regulations be amended to provide for the safe use of feed grade sodium formate as a feed acidifying agent in complete swine feeds, also proposed that FDA amend the animal food additive regulations for formic acid and ammonium formate to limit formic acid and formate salts from all added sources.

DATES: Submit either electronic or written comments on FDA's environmental assessment by October 31, 2016.

ADDRESSES: You may submit comments as follows:

Electronic Submissions

Submit electronic comments in the following way:

- *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 comment, 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–2014–F–0988 for ''Food Additives Permitted in Feed and Drinking Water of Animals; Feed Grade Sodium Formate.'' 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 comment 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: Chelsea Trull, Center for Veterinary Medicine, Food and Drug Administration, 7519 Standish Pl., Rockville, MD 20855, 240–402–6729, *chelsea.trull@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION: Under the Federal Food, Drug, and Cosmetic Act (section 409(b)(5) (21 U.S.C. 348(b)(5)), notice is given that the food additive petition (FAP 2286) filed by BASF Corp., 100 Park Ave., Florham Park, NJ 07932 proposing to amend Title 21 of the Code of Federal Regulations (CFR) in part 573 *Food Additives Permitted in Feed and Drinking Water of Animals* (21 CFR part 573) to provide for the safe use of feed grade sodium formate as a feed acidifying agent in complete swine feeds, also proposed that FDA amend the animal food additive regulations for formic acid (§ 573.480) and ammonium formate (§ 573.170) to limit formic acid and formate salts from all added sources to 1.2 percent of complete feed when multiple sources of formic acid and its salts are used in combination. This element of the petition was not described in the July 25, 2014, notice of petition (79 FR 43325).

Elsewhere in this issue of the **Federal Register**, FDA is publishing a regulation providing for the safe use of feed grade sodium formate as a feed acidifying agent in complete swine feeds.

The potential environmental impact of this action is being reviewed. The Agency will prepare a claim of categorical exclusion or an environmental assessment to evaluate the potential environmental impacts of these actions. Interested persons may submit to the Division of Dockets Management (see **ADDRESSES**) either electronic or written comments regarding this document. It is only necessary to send one set of comments. Identify comments with the docket number found in brackets in the heading of this document. Received comments may be seen in the Division of Dockets Management between 9 a.m. and 4 p.m., Monday through Friday. FDA will also place on public display any comments on potential environmental impact without further announcement in the **Federal Register**. If FDA determines a categorical exclusion applies, neither an environmental assessment nor an environmental impact statement is required. If FDA determines a categorical exclusion does not apply, FDA will prepare an environmental assessment and place it on public display at the Division of Dockets Management (see **DATES** and **ADDRESSES**) for public review and comment.

Dated: September 26, 2016.

Tracey H. Forfa,

Deputy Director, Center for Veterinary Medicine.

[FR Doc. 2016–23645 Filed 9–29–16; 8:45 am]

BILLING CODE 4164–01–P

ENVIRONMENTAL PROTECTION AGENCY

40 CFR Part 52

[EPA–R05–OAR–2016–0134; FRL–9953–51–Region 5]

Air Plan Approval; Wisconsin; NO_X as a Precursor to Ozone, $PM_{2.5}$ Increment Rules and PSD Infrastructure SIP Requirements

AGENCY: Environmental Protection Agency (EPA).

ACTION: Proposed rule.

SUMMARY: The Environmental Protection Agency (EPA) is proposing approval of a revision to Wisconsin's state implementation plan (SIP), revising portions of the State's Prevention of Significant Deterioration (PSD) and ambient air quality programs to address deficiencies identified in EPA's previous narrow infrastructure SIP disapprovals and Finding of Failure to Submit. This SIP revision request is consistent with the Federal PSD rules and addresses the required elements of the fine particulate matter ($PM_{2.5}$) PSD Increments, Significant Impact Levels (SILs) and Significant Monitoring Concentration (SMC) Rule. EPA is also proposing to approve elements of SIP submissions from Wisconsin regarding PSD infrastructure requirements of section 110 of the Clean Air Act (CAA) for the 1997 $PM_{2.5}$, 1997 ozone, 2006 $PM_{2.5}$, 2008 lead, 2008 ozone, 2010 nitrogen dioxide (NO_2), 2010 sulfur dioxide (SO_2), and 2012 $PM_{2.5}$ National Ambient Air Quality Standards (NAAQS). The infrastructure requirements are designed to ensure that the structural components of each state's air quality management program are adequate to meet the state's responsibilities under the CAA.

DATES: Comments must be received on or before October 31, 2016.

ADDRESSES: Submit your comments, identified by Docket ID No. EPA–R05–OAR–2016–0134 at *http://www.regulations.gov,* or via email to *damico.genevieve@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, 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.