would also eliminate the Battle Mountain VORTAC as a reference point in the legal description as it is no longer required. This airspace would support IFR operations at Battle Mountain Airport, Battle Mountain, NV.

Class E airspace designations are published in paragraph 6002, 6004 and 6005 of FAA Order 7400.11D, dated August 8, 2019 and effective September 15, 2019, which is incorporated by reference in 14 CFR 71.1. The Class E airspace designation listed in this document will be published subsequently in the Order. FAA Order 7400.11, Airspace Designations and Reporting Points, is published yearly and effective on September 15.

Regulatory Notices and Analyses

The FAA has determined that this proposed regulation only involves an established body of technical regulations for which frequent and routine amendments are necessary to keep them operationally current, is non-controversial, and unlikely to result in adverse or negative comments. It, therefore: (1) Is not a ''significant regulatory action'' under Executive Order 12866; (2) is not a ''significant rule'' under DOT Regulatory Policies and Procedures (44 FR 11034; February 26, 1979); and (3) does not warrant preparation of a regulatory evaluation as the anticipated impact is so minimal. Given this is a routine matter that will only affect air traffic procedures and air navigation, it is certified that this proposed rule, when promulgated, would not have a significant economic impact on a substantial number of small entities under the criteria of the Regulatory Flexibility Act.

Environmental Review

This proposal will be subject to an environmental analysis in accordance with FAA Order 1050.1F, ''Environmental Impacts: Policies and Procedures'' prior to any FAA final regulatory action.

List of Subjects in 14 CFR Part 71

Airspace, Incorporation by reference, Navigation (air).

The Proposed Amendment

Accordingly, pursuant to the authority delegated to me, the Federal Aviation Administration proposes to amend 14 CFR part 71 as follows:

PART 71—DESIGNATION OF CLASS A, B, C, D, AND E AIRSPACE AREAS; AIR TRAFFIC SERVICE ROUTES; AND REPORTING POINTS

■ 1. The authority citation for 14 CFR part 71 continues to read as follows:

Authority: 49 U.S.C. 106(f), 106(g); 40103, 40113, 40120; E.O. 10854, 24 FR 9565, 3 CFR, 1959–1963 Comp., p. 389.

§ 71.1 [Amended]

■ 2. The incorporation by reference in 14 CFR 71.1 of FAA Order 7400.11D, Airspace Designations and Reporting Points, dated August 8, 2019, and effective September 15, 2019, is amended as follows:

Paragraph 6002 Class E Airspace Designated as Surface Areas.

* * * * *

AWP NV E2 Battle Mountain, NV [Amended]

Battle Mountain Airport, NV
(Lat. 40°35′57″ N, long. 116°52′28″ W)

That airspace extending upward from the surface to and including 2500 feet MSL within a 4.2-mile radius of Battle Mountain Airport, Battle Mountain, NV. This Class E airspace area is effective during the specific dates and times established in advance by a Notice to Airmen. The effective date and time will thereafter be continuously published in the Chart Supplement.

Paragraph 6004 Class E Airspace Areas Designated as an Extension to a Class D or Class E Surface Area.

* * * * *

AWP NV E4 Battle Mountain, NV [NEW]

Battle Mountain Airport, NV
(Lat. 40°35′57″ N, long. 116°52′28″ W)

That airspace extending upward from the surface within 1.3 miles each side of the 228° bearing from the Battle Mountain Airport extending from the 4.2 mile radius to 7 miles southwest of Battle Mountain Airport, Battle Mountain NV.

Paragraph 6005 Class E Airspace Areas Extending Upward From 700 Feet or More Above the Surface of the Earth.

* * * * *

AWP NV E5 Battle Mountain, NV [Amended]

Battle Mountain Airport, NV
(Lat. 40°35′57″ N, long. 116°52′28″ W)

That airspace extending upward from 700 feet above the surface within 16.5-mile radius of the Battle Mountain Airport beginning at the point where the 205° bearing intersects the 16.5-mile radius thence clockwise to the point where the 266° bearing intersects the 16.5-mile radius thence northeast along the 266° bearing to within 7 miles of the airport, thence clockwise along the 7-mile radius to the point where the 65° bearing intersects the 7-mile radius thence to the point where the 77° bearing intersects the 4.2-mile radius thence clockwise to the point where the 158° bearing intersects the 4.2 mile radius, thence to the point of beginning; and that airspace within 2 miles each side of the 49° bearing extending from the 4.2 mile radius to 10.5 miles from the airport.

Issued in Seattle, Washington, on November 19, 2019.

Byron Chew,

Manager, Operations Support Group, Western Service Center.

[FR Doc. 2019–25542 Filed 11–27–19; 8:45 am]

BILLING CODE 4910–13–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 573

[Docket No. FDA–2017–F–4399]

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

AGENCY: Food and Drug Administration, HHS.

ACTION: Notification; petition for rulemaking; amendment.

SUMMARY: The Food and Drug Administration (FDA or we) is announcing that the Zinpro Corp. (Zinpro) has amended their pending petition proposing that the food additive regulations be amended to provide for the safe use of chromium DL-methionine as a nutritional source of chromium in cattle feed. The amendment provides for a change in the feeding rate.

DATES: Submit either electronic or written comments on the petitioner's environmental assessment by December 30, 2019.

ADDRESSES: You may submit comments as follows. Please note that late, untimely filed comments will not be considered. Electronic comments must be submitted on or before December 30, 2019. The *https://www.regulations.gov* electronic filing system will accept comments until 11:59 p.m. Eastern Time at the end of December 30, 2019. Comments received by mail/hand delivery/courier (for written/paper submissions) will be considered timely if they are postmarked or the delivery service acceptance receipt is on or before that date.

Electronic Submissions

Submit electronic comments in the following way:

- *Federal eRulemaking Portal: https://www.regulations.gov*. Follow the instructions for submitting comments. Comments submitted electronically, including attachments, to *https://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 *https://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):* Dockets Management Staff (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

• For written/paper comments submitted to the Dockets Management Staff, 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–2017–F–4399 for ''Zinpro Corp.; Filing of Food Additive Petition (Animal Use).'' Received comments, those filed in a timely manner (see **ADDRESSES**), will be placed in the docket and, except for those submitted as ''Confidential Submissions,'' publicly viewable at *https://www.regulations.gov* or at the Dockets Management Staff 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 *https://www.regulations.gov*. Submit both copies to the Dockets Management Staff. 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: *https://www.gpo.gov/fdsys/pkg/FR-2015-09-18/pdf/2015-23389.pdf*.

Docket: For access to the docket to read background documents or the electronic and written/paper comments received, go to *https://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 Dockets Management Staff, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

FOR FURTHER INFORMATION CONTACT: Chelsea Cerrito, Center for Veterinary Medicine, Food and Drug Administration (HFV–224), 7519 Standish Pl., Rockville, MD 20855, 240–402–6729, *Chelsea.Cerrito@fda.hhs.gov*.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of September 22, 2017 (82 FR 44367), FDA announced that Zinpro Corp., 10400 Viking Dr., Suite 240, Eden Prairie, MN 55344 had filed a petition (FAP 2300) proposing to amend Title 21 of the Code of Federal Regulations (CFR) in part 573 (21 CFR part 573) *Food Additives Permitted in Feed and Drinking Water of Animals* to provide for the safe use of chromium DL-methionine as a nutritional source of chromium in cattle feed. Zinpro has amended the petition by changing the feeding rate.

Zinpro has submitted a revised environmental assessment which the Agency is placing on public display at the Dockets Management Staff for public review and comment (see **DATES** and **ADDRESSES**).

Dated: November 25, 2019.

Lowell J. Schiller,

Principal Associate Commissioner for Policy.

[FR Doc. 2019–25903 Filed 11–27–19; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 573

[Docket No. FDA–2019–F–5401]

Alzchem Trostberg GmbH; Filing of Food Additive Petition (Animal Use)

AGENCY: Food and Drug Administration, HHS.

ACTION: Notification; petition for rulemaking.

SUMMARY: The Food and Drug Administration (FDA or we) is announcing that Alzchem Trostberg GmbH has filed a petition proposing that the food additive regulations be amended to provide for the safe use of guanidinoacetic acid as a precursor of creatine in poultry feeds.

DATES: The food additive petition was filed on September 25, 2019.

ADDRESSES: For access to the docket, go to *https://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 Dockets Management Staff, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

FOR FURTHER INFORMATION CONTACT: Carissa Adams, Center for Veterinary Medicine, Food and Drug Administration,7519 Standish Pl., Rockville, MD 20855, 240–402–6283, *Carissa.Adams@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 a food additive petition (FAP 2309) has been filed by Alzchem Trostberg GmbH, CHEMIEPARK TROSTBERG, Dr.-Albert-Frank-Str. 32, 83308 Trostberg, Germany. The petition proposes to amend Title 21 of the Code of Federal Regulations (CFR) in part 573 (21 CFR part 573) *Food Additives Permitted in Feed and Drinking Water of Animals* to provide for the safe use of guanidinoacetic acid as a precursor of creatine in poultry feeds.

The petitioner has claimed that this action is categorically excluded under 21 CFR 25.32(r) because it is of a type that does not individually or cumulatively have a significant effect on the human environment. In addition, the petitioner has stated that, to their knowledge, no extraordinary circumstances exist. If FDA determines a categorical exclusion applies, neither an environmental assessment nor an environmental impact statement is