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 7400.11B, Airspace Designations and Reporting Points, dated August 3, 2017, and effective September 15, 2017, is amended as follows:

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

* * * * *

AGL WI E5 Milwaukee, WI [Amended]

Milwaukee, General Mitchell International Airport, WI
(Lat. 42°56′49″ N., long. 87°53′49″ W.)
Racine, Batten International Airport, WI
(Lat. 42°45′40″ N., long. 87°48′50″ W.)
Waukesha, Waukesha County Airport, WI
(Lat. 43°02′28″ N., long. 88°14′13″ W.)
Milwaukee, Lawrence J. Timmerman Airport, WI
(Lat. 43°06′37″ N., long. 88°02′04″ W.)

That airspace extending upward from 700 feet above the surface within an 8.4-mile radius of General Mitchell International Airport, and within a 6.6-mile radius of Batten International Airport, and within a 7.5-mile radius of Waukesha County Airport, and within 2 miles each side of the 282° bearing from Waukesha County Airport extending from the 7.5-mile radius to 10.5 miles west of Waukesha County Airport, and within an 8.9-mile radius of Lawrence J. Timmerman Airport.

Issued in Fort Worth, Texas, on September 13, 2017.

Vonnie Royal,

Acting Manager, Operations Support Group, ATO Central Service Center.

[FR Doc. 2017–19948 Filed 9–21–17; 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.

SUMMARY: The Food and Drug Administration (FDA) is announcing that Zinpro Corp. has filed a 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.

DATES: Submit either electronic or written comments on the petitioner's environmental assessment by October 23, 2017.

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 October 23, 2017. The *https://www.regulations.gov* electronic filing system will accept comments until midnight Eastern Time at the end of October 23, 2017. 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 comment, 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 ''Food Additives Permitted in Feed and Drinking Water of Animals; Chromium DL-Methionine.'' 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 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 *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: Carissa Doody, Center for Veterinary Medicine, Food and Drug Administration, 7519 Standish Pl., Rockville, MD 20855, 240–402–6283, *carissa.doody@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 2300) has been filed by the Zinpro Corp., 10400 Viking Dr., Suite 240, Eden Prairie, MN 55344. The petition proposes 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 chromium DL-methionine as a nutritional source of chromium in cattle feed.

The potential environmental impact of this action is being reviewed. To encourage public participation consistent with regulations issued under the National Environmental Policy Act (40 CFR 1501.4(b)), the Agency is placing the environmental assessment (EA) submitted with the petition that is the subject of this notice on public display at the Dockets Management Staff for public review and comment (see **DATES** and **ADDRESSES**). FDA will also place on public display any amendments to, or comments on, the petitioner's EA without further announcement in the **Federal Register**.

If, based on its review, the Agency finds that an environmental impact statement is not required and this petition results in a regulation, the notice of availability of the Agency's finding of no significant impact and the evidence supporting that finding will be published with the regulation in the **Federal Register** in accordance with 21 CFR 25.51(b).

Dated: September 15, 2017.

Anna K. Abram,

Deputy Commissioner for Policy, Planning, Legislation, and Analysis.

[FR Doc. 2017–20195 Filed 9–21–17; 8:45 am]

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LIBRARY OF CONGRESS

Copyright Royalty Board

37 CFR Part 387

[Docket No. 15–CRB–0010–CA–S (Sports Rule Proceeding)]

Adjustment of Royalty Rates for Statutory Cable Retransmission License

AGENCY: Copyright Royalty Board (CRB), Library of Congress.

ACTION: Request for comments.

SUMMARY: The Copyright Royalty Judges solicit reply comments on the legal issue of the purported reach of the proposed rules relating to a cable system license royalty surcharge for retransmission of broadcasts of certain professional sports events.

DATES: Reply comments are due on or before October 23, 2017. Surreplies from original commenters are due on or before November 1, 2017.

ADDRESSES: You may make replies and surreplies, identified by docket number 15–CRB–0010–CA–S (Sports Rule Proceeding), by any of the following methods:

CRB's electronic filing application: Submit comments online in eCRB at *https://app.crb.gov/.*

U.S. mail: Copyright Royalty Board, P.O. Box 70977, Washington, DC 20024–0977; or

Overnight service (only USPS Express Mail is acceptable): Copyright Royalty Board, P.O. Box 70977, Washington, DC 20024–0977; or

Commercial courier: Address package to: Copyright Royalty Board, Library of Congress, James Madison Memorial Building, LM–403, 101 Independence Avenue SE., Washington, DC 20559–6000. Deliver to: Congressional Courier Acceptance Site, 2nd Street NE and D Street NE., Washington, DC; or

Hand delivery: Library of Congress, James Madison Memorial Building, LM–401, 101 Independence Avenue SE., Washington, DC 20559–6000.

Instructions: Unless submitting online, commenters must submit an original, five paper copies, and an electronic version on a CD. All submissions must include the CRB's name and docket number. All submissions received will be posted without change to eCRB on *https://app.crb.gov* including any personal information provided.

Docket: For access to the docket to read background documents or comments received, go to eCRB, the Copyright Royalty Board's electronic filing and case management system, at *https://app.crb.gov/* and search for docket number 15–CRB–0010–CA–S (Sports Rule Proceeding). For documents not yet uploaded to eCRB (because it is a new system), go to the agency Web site at *http://www.crb.gov/* or contact the CRB Program Specialist.

FOR FURTHER INFORMATION CONTACT: Anita Blaine, CRB Program Specialist, by telephone at (202) 707–7658 or email at *crb@loc.gov.*

SUPPLEMENTARY INFORMATION: In May 2017, the Copyright Royalty Judges (Judges) published notice of an agreed settlement and proposed rules to adjust royalties payable by certain cable system operators for a license to retransmit broadcast sports programming (the Sports Surcharge Rules). *See* 82 FR 24611 (May 30, 2017). Specifically, the rules as proposed would be applicable to ''Form 3'' cable systems [1] retransmitting ''eligible professional sports events.'' The proposed rules define ''eligible professional sports event'' as a game involving member teams of Major League Baseball, the National Basketball Association, the National Football League, the National Hockey League, and the Women's National Basketball Association.[2]

The Copyright Act (Act) directs that the Judges provide (1) an opportunity to comment to nonparticipants who would be bound and (2) an opportunity to comment *and object* to *participants* who would be bound. *See* 11 U.S.C. 801(b)(7)(A)(i). The Judges may decline to adopt an agreement as a basis for statutory terms and rates for ''participants that are not parties to the [settlement] agreement,'' if a *participant* objects to the agreement and the Judges conclude that the settlement ''does not provide a reasonable basis for setting'' rates or terms. *Id.* at § 801(b)(7)(A)(ii).

The statutory language does not prohibit the Judges from considering whether the proposed provisions are contrary to statutory law. *See [Register of Copyrights] Review of Copyright Royalty Judges Determination,* Docket no. 2009–1, 74 FR 4537, 4540 (Jan. 26, 2009) (Register's Opinion).[3] In the cited

[1] ''Form 3'' cable systems are those with semi-annual gross receipts, as defined by statute, greater than $527,600. *See* 17 U.S.C. 111(d)(1)(B), (E), & (F).

[2] The proposed sports programming surcharge would also apply to an ''eligible collegiate sports event'' as that term is defined in the proposed regulations. Eligible collegiate sports events are limited to games that involve certain Division I football or men's basketball teams. Proposed Rule 387.2(e)(5).

[3] The Act permits the Register of Copyrights (Register) to review for legal error the Judges' resolution of a material question of substantive law under the Act ''that underlies or is contained in a final determination'' by the Judges. *See* 17 U.S.C.