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

ANM MT E5 Kalispell, MT [Amended]

Glacier Park International Airport, MT (Lat. 48°18'38" N, long. 114°15'22" W)

That airspace extending upward from 700 feet above the surface within a 7.5-mile radius of the airport, and within 2.3 miles each side of the 138° bearing from the airport, extending from the 7.5-mile radius to 13.4 miles southeast of the airport, and with 2 miles each side of the 215° bearing from the airport, extending from the 7.5-mile radius to 19.5 miles southwest of the airport; and that airspace extending upward from 1,200 feet above the surface within a 25-mile radius of the airport beginning at the 270° bearing from the airport, clockwise to the 090° bearing from the airport, thence along the 090° bearing to 45 miles east of the airport, thence within a 45-mile radius of the airport clockwise to the 270° bearing from the airport, thence along the 270° bearing to the point of beginning, 25 miles west of Glacier Park International Airport.

Issued in Seattle, Washington, on September 18, 2020.

B.G. Chew,
Acting Group Manager, Operations Support Group, Western Service Center.

September 18, 2020.

FOR FURTHER INFORMATION CONTACT:


SUPPLEMENTARY INFORMATION: On August 19, 2020, the Commission issued a Petition for rulemaking informing that Bloom Energy Corporation submitted a petition for rulemaking requesting that the Commission clarify that the thermal energy output produced by a topping-cycle facility’s solid oxide fuel cell system when used to reform methane and produce hydrogen for fuel for electricity generation by that facility is useful thermal energy output that would enable the facility powered by such fuel cells to be certified as a qualifying cogeneration facility, all as more fully explained in the petition.


Nathaniel J. Davis, Sr.,
Deputy Secretary.

[FR Doc. 2020–21037 Filed 10–2–20; 8:45 am]
BILLING CODE 6717–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 1
[Docket No. FDA–2014–N–0053]

Requirements for Additional Traceability Records for Certain Foods; Proposed Rule; Public Meetings; Request for Comments

AGENCY: Food and Drug Administration, HHS.

ACTION: Notification of public meeting; request for comments.

SUMMARY: The Food and Drug Administration (FDA, the Agency, or we) is announcing three virtual public meetings entitled “Requirements for Additional Traceability Records for Certain Foods; Proposed Rule.” The purpose of the public meetings is to discuss the proposed rule entitled “Requirements for Additional Traceability Records for Certain Foods,” which was issued under the FDA Food Safety Modernization Act (FSMA). These public meetings are intended to facilitate and support the public’s evaluation and commenting process on the proposed rule.

DATES: The public meetings will be held on November 6, 2020, from 8:30 a.m. Eastern Time to 3:30 p.m. Eastern Time; November 18, 2020, from 9:30 a.m. Eastern Time to 4:30 p.m. Eastern Time; and December 2, 2020, from 11:30 a.m. Eastern Time to 6:30 p.m. Eastern Time. Submit either electronic or written comments on the notice by January 21, 2021. See “How to Participate in the Public Meetings” in the SUPPLEMENTARY INFORMATION section of this document for closing dates for advanced registration and other information regarding meeting participation.

ADDRESSES: Due to the impact of the COVID–19 pandemic, these meetings will be held virtually to help protect the public and limit the spread of the virus. 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 January 21, 2021. The https://www.regulations.gov electronic filing system will accept comments until 11:59 p.m. Eastern Time at the end of January 21, 2021. 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, Rockville, MD 20852.

For written/paper comments submitted to the Dockets Management...