affected by lead exposure, as appropriate.

At least half of the committee will consist of Federal representatives from a range of agencies that may include the Department of Housing and Urban Development; the Environmental Protection Agency; the Consumer Product Safety Commission; the Centers for Medicare and Medicaid Services; the Health Resources and Services Administration; the Food and Drug Administration; the U.S. Department of Agriculture; the Occupational Safety and Health Administration; the National Institute of Environmental Health Sciences; the U.S. Geological Survey; and such additional federal, state, tribal, and local public and private officials as the Secretary deems necessary for the committee to carry out its function. The rest of the committee will consist of non-Federal members. Only non-Federal members are being solicited with this announcement.

The U.S. Department of Health and Human Services policy stipulates that committee membership be balanced in terms of points of view represented, and the committee's function. Appointments shall be made without discrimination on the basis of age, race, ethnicity, gender, sexual orientation, gender identity, HIV status, disability, and cultural, religious, or socioeconomic status. Nominees must be U.S. citizens and cannot be full-time employees of the U.S. Government. Current participation on federal workgroups or prior experience serving on a federal advisory committee does not disqualify a candidate; however, HHS policy is to avoid excessive individual service on advisory committees and multiple committee memberships.

Committee members are Special Government Employees, requiring the filing of financial disclosure reports at the beginning and annually during their terms. CDC reviews potential candidates for LEPAC membership each year and provides a slate of nominees for consideration to the Secretary of HHS for final selection. HHS notifies selected candidates of their appointment as soon as the HHS selection process is completed. Note that the need for different expertise varies from year to year and a candidate who is not selected in one year may be reconsidered in a subsequent year. Candidates should submit the following items:

- Current curriculum vitae, including complete contact information (telephone numbers, mailing address, email address).
- At least one letter of recommendation from person(s) not employed by the U.S. Department of Health and Human Services. (Candidates may submit letter(s) from current HHS employees if they wish, but at least one letter must be submitted by a person not employed by an HHS agency (*e.g.,* CDC, NIH, FDA, etc.)

Nominations may be submitted by the candidate him- or herself or by the person/organization recommending the candidate.

The Director, Strategic Business Initiatives Unit, Office of the Chief Operating Officer, Centers for Disease Control and Prevention, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities, for both the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

Kalwant Smagh,

Director, Strategic Business Initiatives Unit, Office of the Chief Operating Officer, Centers for Disease Control and Prevention.

[FR Doc. 2021–19222 Filed 9–3–21; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2021–N–0929]

Food and Drug Administration New Era of Smarter Food Safety Summit on E-Commerce; Public Meeting; Request for Comments

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of public meeting; request for comments.

SUMMARY: The Food and Drug Administration (FDA or we) is announcing a virtual public meeting entitled ''FDA New Era of Smarter Food Safety Summit on E-Commerce: Ensuring the Safety of Foods Ordered Online and Delivered Directly to Consumers.'' The purpose of the public meeting is to engage with stakeholders and invite input on various topics pertaining to the implementation of Core Element 3.1 of the New Era of Smarter Food Safety Blueprint. We intend to use information resulting from the public meeting to determine what action(s), if any, should be taken to help ensure the safe production and delivery of human and animal foods sold through new e-commerce business models.

DATES: The public meeting will be held over 3 days on October 19, 2021, from 11:30 a.m. to 5:30 p.m. Eastern Time; October 20, 2021, from 11:30 a.m. to 5:15 p.m. Eastern Time, and October 21, 2021, from 11:30 a.m. to 3:45 p.m. Eastern Time. Submit either electronic or written comments on this public meeting by November 20, 2021. See ''Participating in the Public Meeting'' in the **SUPPLEMENTARY INFORMATION** section of this document for registration and other information regarding meeting participation.

ADDRESSES: The public meeting will be held virtually. For more information on the public meeting, see: *https://www.fda.gov/food/workshops-meetings-webinars-food-and-dietary-supplements/new-era-smarter-food-safety-summit-e-commerce-ensuring-safety-foods-ordered-online-and-delivered.*

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 November 20, 2021. The *https://www.regulations.gov* electronic filing system will accept comments until 11:59 p.m. Eastern Time at the end of November 20, 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, 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–2021–N–0929 for ''FDA New Era of Smarter Food Safety Summit on E-Commerce.'' 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, 240–402–7500.

- 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.'' We will review this copy, including the claimed confidential information, in our 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.govinfo.gov/content/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, 240–402–7500.

FOR FURTHER INFORMATION CONTACT: Juanita Yates, Center for Food Safety and Applied Nutrition, (HFS–009), Food and Drug Administration, 5001 Campus Dr., College Park, MD 20740, 240–402–1731, *juanita.yates@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION:

I. Background

In April 2019, we announced the New Era of Smarter Food Safety initiative and in July 2020, we released the New Era of Smarter Food Safety Blueprint (see *https://www.fda.gov/food/new-era-smarter-food-safety/new-era-smarter-food-safety-blueprint*) (Blueprint). The Blueprint outlines the effort to modernize approaches to food safety as we respond to unique demands on our food system and work to ensure the safety and security of our food supply. This virtual public meeting (Summit) will focus on Core Element 3.1: Ensure Safety of Food Produced or Delivered Using New Business Models. Specifically, we invite discussion and input on human and animal foods sold through Business to Consumer (B2C) e-commerce. B2C e-commerce is the manufacturing, packaging, labeling, storage, and delivery of human and animal foods sold directly to consumers, through commercial transactions conducted electronically on the internet.

The Summit is an opportunity for us to share our current understanding of human and animal foods sold through new business models and hear from the public. The Summit will enhance our knowledge of possible food safety risks related to these new business models and help us identify what additional courses of action, if any, are needed to address potential food safety vulnerabilities.

We invite industry, consumers, consumer and public health organizations, academia, Federal, State, local, and tribal governments, foreign governments, and other interested parties to join the discussion and provide their perspectives on these issues.

II. Topics for Discussion at the Public Meeting

The Summit will address a variety of topics related to human and animal foods sold through B2C e-commerce, including:

- Types of B2C e-commerce models (*e.g.,* produce and meal kit subscription services, ghost kitchens, dark stores);
- Safety risks associated with foods sold through B2C e-commerce;
- Standards of care used by industry to control these safety risks;
- Types of delivery models (*e.g.,* third-party delivery, autonomous delivery models);
- Regulatory approaches to food sold through B2C e-commerce, including challenges and gaps that need to be addressed; and
- Labeling of foods sold through B2C e-commerce.

During the Summit, experts from FDA, industry, academia, consumer and public health organizations, domestic and foreign governments will be asked to address these topics. Each day, there will also be an opportunity for registered participants to ask questions and engage with these experts, as well as to offer open public comment for those who select this option when registering (see Part III, ''Participating in the Public Meeting'').

Before the meeting date, we will post the agenda and additional background materials on the internet at: *https://www.fda.gov/food/workshops-meetings-webinars-food-and-dietary-supplements/new-era-smarter-food-safety-summit-e-commerce-ensuring-safety-foods-ordered-online-and-delivered.* Registered participants will be notified when these materials are posted. There will be an opportunity for interested stakeholders to submit written comments following the meeting.

III. Participating in the Public Meeting

Registration: To register for the public meeting, please visit the following website to register: *https://www.fda.gov/food/workshops-meetings-webinars-food-and-dietary-supplements/new-era-smarter-food-safety-summit-e-commerce-ensuring-safety-foods-ordered-online-and-delivered.* Please provide complete contact information for each attendee, including name, title, affiliation, address, email, and telephone.

This is a virtual public meeting. Registration is free and will be open through the day of the meeting. Persons interested in attending this public meeting must register at: *https://www.fda.gov/food/workshops-meetings-webinars-food-and-dietary-supplements/new-era-smarter-food-safety-summit-e-commerce-ensuring-safety-foods-ordered-online-and-delivered.* Upon registering, they will receive a confirmation email with a link for the public meeting. Approximately

24 to 48 hours before the event, registrants will receive an email with the meeting link and a formal calendar invitation.

Request to Provide Open Public Comment: During online registration, you may indicate if you wish to make open public comments during the public meeting and which topic(s) you would like to address. All requests to make public comments must be received by October 8, 2021 at 11:59 p.m. Eastern Time. We will do our best to accommodate requests to make public comments. We are seeking to have a broad representation of ideas and issues presented at the meeting. Individuals and organizations with common interests are urged to consolidate or coordinate their comments. FDA will determine the amount of time allotted to each commenter, the meeting day, the approximate time open public comments are to be provided and notify all registrants who requested to make public comments.

Streaming Webcast of the Public Meeting: This public meeting will be broadcast via YouTube.

Transcripts: Please be advised that as soon as a transcript of the public meeting is available, it will be accessible at *https://www.regulations.gov.* It may be viewed at the Dockets Management Staff (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20850. A link to the transcript will also be available on the internet at: *https://www.fda.gov/food/workshops-meetings-webinars-food-and-dietary-supplements/new-era-smarter-food-safety-summit-e-commerce-ensuring-safety-foods-ordered-online-and-delivered.*

Dated: August 31, 2021.

Lauren K. Roth,

Acting Principal Associate Commissioner for Policy.

[FR Doc. 2021–19219 Filed 9–3–21; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2013–N–1529]

Agency Information Collection Activities; Proposed Collection; Comment Request; Reclassification Petitions for Medical Devices

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA, Agency, or we) is announcing an opportunity for public comment on the proposed collection of certain information by the Agency. Under the Paperwork Reduction Act of 1995 (PRA), Federal Agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, and to allow 60 days for public comment in response to the notice. This notice solicits comments on information collection associated with reclassification of medical devices.

DATES: Submit either electronic or written comments on the collection of information by November 8, 2021.

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 November 8, 2021. The *https://www.regulations.gov* electronic filing system will accept comments until 11:59 p.m. Eastern Time at the end of November 8, 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, 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–2013–N–1529 for ''Agency Information Collection Activities; Proposed Collection; Comment Request; Reclassification Petitions for Medical Devices.'' 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, 240–402–7500.

- 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.govinfo.gov/content/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