

data. You may submit comments on any topic related to this activity. Comments received, including attachments and other supporting materials, are part of the public record and subject to public disclosure. Do not include any information in your comment or supporting materials that you consider confidential or inappropriate for public disclosure. If you include your name, contact information, or other information that identifies you in the body of your comments, that information will be on public display. CDC will review all submissions and may choose to redact, or withhold, submissions containing private or proprietary information such as Social Security numbers, medical information, inappropriate language, or duplicate/near duplicate examples of a mass-mail campaign. CDC will carefully consider all comments submitted into the docket. CDC does not accept comment by email.

Oral Public Comment: The public is welcome to participate, via Zoom, during the public comment periods on September 28, 2021, from 11:30 a.m. to 12:00 p.m., EDT, and on September 29, 2021, from 12:10 p.m. to 12:40 p.m., EDT. Please note that the public comment periods end at the times indicated above. Each commenter will be provided up to five minutes for comment. A limited number of time slots are available and will be assigned on a first come-first served basis. Members of the public who wish to address the WTCHP STAC during the oral public comment sessions must sign up by providing their name and desired date for commenting to Mia Wallace, Committee Management Specialist, via email: MWallace@cdc.gov, or the addresses section provided in this notice by September 21, 2021.

Written Public Comment: Written comments will also be accepted from those unable to attend the public session per the instructions provided in the addresses section above. Written comments received in advance of the meeting will be included in the official record of the meeting. Written comments received by September 21, 2021 will be provided to the STAC prior to the meeting. The docket will close on September 29, 2021.

Policy on Redaction of Committee Meeting Transcripts (Public Comment): Transcripts will be prepared and posted to <http://www.regulations.gov> within 60 days after the meeting. If a person making a comment gives their name, no attempt will be made to redact that name. NIOSH will take reasonable steps to ensure that individuals making public comments are aware that their comments (including their name, if

provided) will appear in a transcript of the meeting posted on a public website. Such reasonable steps include a statement read at the start of the meeting stating that transcripts will be posted, and names of speakers will not be redacted. If individuals in making a statement reveal personal information (e.g., medical information) about themselves, that information will not usually be redacted. The CDC Freedom of Information Act coordinator will, however, review such revelations in accordance with the Freedom of Information Act and, if deemed appropriate, will redact such information. Disclosures of information concerning third party medical information will be redacted.

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.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Solicitation of Nominations for Membership on the Lead Exposure and Prevention Advisory Committee (LEPAC)

AGENCY: Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

ACTION: Notice.

SUMMARY: The Centers for Disease Control and Prevention (CDC) is soliciting nominations for membership on the LEPAC. The LEPAC is composed of 15 members that are Federal and non-Federal experts in fields associated with lead screening, the prevention of lead exposure, and services for individuals and communities affected by lead exposure. Nominations are being sought for individuals with expertise in the fields of epidemiology, toxicology, mental health, pediatrics, early childhood education, special education, diet and nutrition, and environmental

health. Members may be invited to serve for three-year terms. Selection of members is based on candidates' qualifications to contribute to the accomplishment of LEPAC objectives.

DATES: Nominations for membership on the LEPAC must be received no later than November 1, 2021. Packages received after this time will not be considered for the current membership cycle.

ADDRESSES: All nominations should be emailed to LEPAC@cdc.gov.

FOR FURTHER INFORMATION CONTACT: Paul Allwood, Ph.D., M.P.H., Designated Federal Officer, National Center for Environmental Health, CDC, 4770 Buford Highway NE, Atlanta, GA 30329-4018, Telephone: (770) 488-6774, PAllwood@cdc.gov.

SUPPLEMENTARY INFORMATION: The members of this committee are selected by the Secretary of the U.S. Department of Health and Human Services (HHS). The committee's objective is to advise the Secretary, HHS and the Director, Centers for Disease Control and Prevention/Administrator, Agency for Toxic Substances and Disease Registry on a range of activities to include: (1) Review of Federal programs and services available to individuals and communities exposed to lead; (2) review of the current research on lead exposure to identify additional research needs; (3) review of and identification of best practices, or the need for best practices regarding lead screening and the prevention of lead exposure; (4) identification of effective services, including services relating to healthcare, education, and nutrition for individuals and communities affected by lead exposure and lead poisoning, including in consultation with, as appropriate, the lead exposure registry as established in Public Law 114-322 Section 2203(b) (42 U.S.C. 300j-27); and (5) undertaking of any other review or activities that the Secretary determines to be appropriate.

Annually as determined necessary by the Secretary or as required by Congress, the committee shall submit a report to include: (1) An evaluation of the effectiveness of the Federal programs and services available to individuals and communities exposed to lead; (2) an evaluation of additional lead exposure research needs; (3) an assessment of any effective screening methods or best practices used or developed to prevent or screen for lead exposure; (4) input and recommendations for improved access to effective services relating to health care, education, or nutrition for individuals and communities impacted by lead exposure; and (5) any other recommendations for communities

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.

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Kalwant Smagh,

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

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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”).