How do I submit confidential business information?

The Commission will provide confidential treatment for identified confidential information to the extent allowed by law. If your comments contain confidential information, you must submit the following by email to the address listed above under **ADDRESSES**:

- A transmittal letter requesting confidential treatment that identifies the specific information in the comments for which protection is sought and demonstrates that the information is a trade secret or other confidential research, development, or commercial information.
- A confidential copy of your comments, consisting of the complete filing with a cover page marked ''Confidential-Restricted,'' and the confidential material clearly marked on each page.
- A public version of your comments with the confidential information excluded. The public version must state ''Public Version—confidential materials excluded'' on the cover page and on each affected page and must clearly indicate any information withheld.

Will the Commission consider late comments?

The Commission will consider all comments received before the close of business on the comment closing date indicated above under **DATES**. To the extent possible, we will also consider comments received after that date.

How can I read comments submitted by other people?

You may read the comments received by the Commission at the Commission's Electronic Reading Room at *https://www2.fmc.gov/readingroom/.*

By the Commission.

William Cody,

Secretary.

[FR Doc. 2022–17582 Filed 8–12–22; 8:45 am]

BILLING CODE 6730–02–P

GOVERNMENT ACCOUNTABILITY OFFICE

Comptroller General's Advisory Council on Government Auditing Standards; Notice of Meeting

The Comptroller General's Advisory Council on Government Auditing Standards will hold a meeting on Wednesday, September 21, 2022, from 10:00 a.m. to 1:30 p.m. to discuss updates and revisions to the Government Auditing Standards. The meeting will be virtual and is open to the public.

The agenda, discussion materials and teleconference information for the virtual meeting will be available at *https://www.gao.gov/yellowbook* approximately one week before the meeting. Any interested person may attend the meeting as an observer. Members of the public will be provided an opportunity to present questions to the Council during a brief period in the afternoon on matters directly related to the proposed update and revision.

Questions concerning the meeting may be emailed to *YellowBook@gao.gov*. For further information, please contact Roger Bradley, Senior Auditor, at 202–512–7069. To request a reasonable accommodation (RA) for this event, email GAO's RA office at *ReasonableAccommodations@gao.gov*. Please request all accommodations at least 5 business days prior to the event (by September 14th).

Authority: Pub. L. 67–13, 42 Stat. 20 (June 10, 1921).

James R. Dalkin,

Director, Financial Management and Assurance, U.S. Government Accountability Office.

[FR Doc. 2022–17466 Filed 8–12–22; 8:45 am]

BILLING CODE 1610–02–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[Docket No. CDC–2022–0096; NIOSH 232]

Board of Scientific Counselors, National Institute for Occupational Safety and Health (BSC, NIOSH), National Firefighter Registry Subcommittee

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

ACTION: Notice of meeting and request for comment.

SUMMARY: In accordance with the Federal Advisory Committee Act, the CDC announces the following meeting of the Board of Scientific Counselors, National Institute for Occupational Safety and Health (BSC, NIOSH), National Firefighter Registry Subcommittee. This meeting is open to the public via virtual meeting, limited only by the number of web conference seats (500 web conference seats are available). Time will be available for public comment.

DATES: The meeting will be held on September 6, 2022, from 1:00 p.m. to 4:45 p.m., EDT.

Written comments must be received on or before August 30, 2022.

ADDRESSES: If you wish to attend the meeting, please register at the NIOSH website at *https://www.cdc.gov/niosh/bsc/nfrs/registration.html* or by telephone at (404) 498–2581 no later than August 30, 2022.

You may submit comments, identified by Docket No. CDC–2022–0096; NIOSH–232, by either of the methods listed below. Do not submit comments for the docket by email. CDC does not accept comments for the docket by email.

- *Federal eRulemaking Portal: https://www.regulations.gov.* Follow the instructions for submitting comments.
- *Mail:* Sherri Diana, NIOSH Docket Office, National Institute for Occupational Safety and Health, 1090 Tusculum Avenue, Mailstop C–34, Cincinnati, Ohio 45226. Attn: Docket No. CDC–2022–0096; NIOSH–232.

Instructions: All submissions received must include the Agency name and Docket Number. Docket No. CDC–2022–0096; NIOSH–232 will close August 30, 2022.

FOR FURTHER INFORMATION CONTACT: Emily J.K. Novicki, M.A., M.P.H., Designated Federal Officer, BSC, NIOSH, CDC, 1600 Clifton Road NE, Mailstop V24–4, Atlanta, Georgia, 30329–4027; Telephone: (404) 498–2581; Email: *ENovicki@cdc.gov.*

SUPPLEMENTARY INFORMATION:

Background: The Secretary of Health and Human Services, the Assistant Secretary for Health, and by delegation the Director, Centers for Disease Control and Prevention, are authorized under Sections 301 and 308 of the Public Health Service Act to conduct directly, or by grants or contracts, research, experiments, and demonstrations relating to occupational safety and health and to mine health.

The Board of Scientific Counselors, National Institute for Occupational Safety and Health provides advice to the Director, National Institute for Occupational Safety and Health, on NIOSH research and prevention programs. The Board also provides guidance on the Institute's research activities related to developing and evaluating hypotheses, systematically documenting findings, and disseminating results. In addition, the Board evaluates the degree to which the activities of NIOSH: (1) conform to those standards of scientific excellence appropriate for Federal scientific institutions in accomplishing objectives

in occupational safety and health; (2) address currently relevant needs in the fields of occupational safety and health either alone or in conjunction with other known activities inside and outside of NIOSH; and (3) produce their intended results in addressing important research questions in occupational safety and health, both in terms of applicability of the research findings and dissemination of the findings.

Purpose: The BSC, NIOSH National Firefighter Registry Subcommittee (the Subcommittee) provides scientific expertise to the Board of Scientific Counselors that will assist the BSC in advising the Director about NIOSH's efforts to establish and operate the National Firefighter Registry. Specifically, the Subcommittee advises the Board of Scientific Counselors on the following issues pertaining to the ''required strategy'' as mandated by the Firefighter Cancer Registry Act of 2018 (the Act): (1) Increase awareness of the National Firefighter Registry and encourage participation among all groups of firefighters; (2) consider data collection needs; (3) consider data storage and electronic access of health information; and (4) in consultation with subject matter experts, develop a method for estimating the number and type of fire incidents attended by a firefighter. Additional responsibilities of the Subcommittee are to provide guidance to the BSC regarding inclusion and the maintenance of data on firefighters as required by the Act.

Matters To Be Considered: The agenda for the meeting addresses issues related to: The National Firefighter Registry project overview and status, protocol updates, enrollment system demonstration, project launch, and future planning applicable to stakeholders. Agenda items are subject to change as priorities dictate.

The agenda is posted on the NIOSH website at *https://www.cdc.gov/niosh/bsc/nfrs/.*

Public Participation

Written Public Comment: Written comments will be accepted per the instructions provided in the **ADDRESSES** section above. Comments received in advance of the meeting are part of the public record and are subject to public disclosure. They will be included in the official record of the meeting. 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.

Written comments received by August 30, 2022, will be provided to the Subcommittee prior to the meeting.

Oral Public Comment: The public is welcome to participate during the public comment period, from 3:15 p.m. to 3:30 p.m., EDT, September 6, 2022. Each commenter will be provided up to 5 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 Subcommittee are requested to contact Designated Federal Officer for scheduling purposes (see **FOR FUTHER INFORMATION CONTACT** above).

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. 2022–17476 Filed 8–12–22; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2022–N–1633]

Soliciting Public Comment on Appendix A of the Food and Drug Administration's July 2018 Guidance Entitled ''Abbreviated New Drug Application Submissions—Amendments To Abbreviated New Drug Applications Under Generic Drug User Fee Amendments;'' Notice; Establishment of a Public Docket; Request for Comments

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; establishment of a public docket; request for comments.

SUMMARY: The Food and Drug Administration (FDA or the Agency) is announcing the establishment of a docket to solicit comments on the content of Appendix A in the July 2018 guidance for industry entitled ''ANDA Submissions—Amendments to Abbreviated New Drug Applications Under GDUFA'' (ANDA Amendments Guidance). We are soliciting comments on the content of Appendix A. The Agency is taking this action to fulfill the Agency's commitment described in section IX.B. of the GDUFA Reauthorization Performance Goals and Program Enhancements Fiscal Years 2023–2027 Commitment Letter (GDUFA III Commitment Letter).

DATES: Either electronic or written comments must be submitted by October 14, 2022.

ADDRESSES: You may submit comments as follows. Please note that late, untimely filed comments will not be considered. The *https://www.regulations.gov* electronic filing system will accept comments until 11:59 p.m. Eastern Time at the end of October 14, 2022. Comments received by mail/hand delivery/courier (for written/paper submissions) will be considered timely if they are received 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: