the open public hearing by sending an email to OMPTfeedback@fda.hhs.gov by Tuesday, August 14, 2018. Requests for participation in the open public hearing are accepted until 9 a.m. on Tuesday September 4, 2018, and will be accepted as long as time allows. The email should contain complete contact information for each attendee (name, title, degree(s), affiliation, address, email address, and telephone number). For those wishing to present at the hearing, the email should also include a presentation title. Those without email access can register by contacting Allison Hoffman at 301–796–9203 by Tuesday, August 14, 2018 (see FOR FURTHER INFORMATION CONTACT). An email to omptfeedback@fda.hhs.gov no later than Tuesday, August 14, 2018 will be accepted for each attendee (name, affiliation, address, email address and telephone number). For those wishing to present at the hearing, the email should also include a presentation title. Those without email access can register by contacting Allison Hoffman at 301–796–9203 by Tuesday, August 14, 2018 (see FOR FURTHER INFORMATION CONTACT). An agenda for the hearing and any other background materials will be made available 5 days before the hearing at https://www.fda.gov/NewsEvents/MeetingsConferencesWorkshops/UCM610692.htm.

FDA will try to accommodate all persons who wish to make a presentation. Individuals wishing to present should identify the number of the specific question, or questions, they wish to address. This will help FDA organize the presentations. Individuals and organizations with common interests should consolidate or coordinate their presentations and request time for a joint presentation. FDA will notify registered presenters of their scheduled presentation times. The time allotted for each presenter will depend on the number of individuals who wish to speak. Presenters are encouraged to submit an electronic copy of their presentation (PowerPoint or PDF) to OMPTfeedback@fda.hhs.gov on or before Thursday, August 16, 2018. Those who are not giving electronic presentations are encouraged to submit a single slide (PowerPoint or PDF) with their name, affiliation, and topic. Persons registered to make either an oral presentation or participate as part of the open public hearing are encouraged to arrive at the hearing room early and check in at the onsite registration table to confirm their designated presentation time. An agenda for the hearing and any other background materials will be made available 5 days before the hearing at https://www.fda.gov/NewsEvents/MeetingsConferencesWorkshops/ucm580561.htm.

If you need special accommodations because of a disability, please contact OMPTfeedback@fda.hhs.gov (see FOR FURTHER INFORMATION CONTACT) no later than Friday, August 17, 2018, at 12 noon Eastern Time.

Streaming Webcast of the Public Hearing: For those unable to attend in person, FDA will provide a live webcast of the hearing. To join the hearing via the webcast, please go to https://collaboration.fda.gov/biosimilarspart15.

Transcripts: Please be advised that as soon as a transcript is available, it will be accessible at https://www.regulations.gov. It may be viewed at the Dockets Management Staff (see ADDRESSES).

IV. Notification of Hearing Under 21 CFR Part 15

The Commissioner of Food and Drugs is announcing that the public hearing will be held in accordance with part 15 (21 CFR part 15). The hearing will be conducted by a presiding officer, who will be accompanied by FDA senior management from the Office of the Commissioner, the Center for Drug Evaluation and Research, and the Center for Biologics Evaluation and Research. Under §15.30(f) (21 CFR 15.30(f)), the hearing is informal and the rules of evidence do not apply. No participant may interrupt the presentation of another participant. Only the presiding officer and panel members can pose questions; they can question any person during or at the conclusion of each presentation. Public hearings under part 15 are subject to FDA’s policy and procedures for electronic media coverage of FDA’s public administrative proceedings (21 CFR part 10, subpart C). Under 21 CFR 10.205, representatives of the media may be permitted, subject to certain limitations, to videotape, film, or otherwise record FDA’s public administrative proceedings, including presentations by participants. The hearing will be transcribed as stipulated in §15.30(b) (see Transcripts). To the extent that the conditions for the hearing, as described in this notification, conflict with any provisions set out in part 15, this notification acts as a waiver of those provisions as specified in §15.30(h). Dated: July 19, 2018.

Leslie Kux, Associate Commissioner for Policy.
by participating by webinar (See table below).

1. To attend the stakeholder meeting in Arlington, Virginia:
   • Address—201 12th Street South, Arlington, Virginia 22202.
   • When you enter the building, take the East elevators to your right, up to the 4th Floor reception area, 4E401, to check in. You will then be escorted to the conference room.
   • Nearest metro stations: Pentagon City, and Crystal City. Parking is available on the street and in the building.

2. To participate at the Webinar by Phone or WebEx:
   By Phone—
   • Dial the toll-free conference number (Verizon): 1–866–718–1874.
   • Attendee access code: 242 716 6.
   By WebEx—
   • To log into the Webinar, go to: https://dol.webex.com.
   • Enter Meeting number: 642 399 450.
   • Meeting password: MIne2018.

A. Stakeholder Meetings

SAFETY IMPROVEMENT TECHNOLOGIES FOR MOBILE EQUIPMENT AT SURFACE MINES, AND FOR BELT CONVEYORS AT SURFACE AND UNDERGROUND MINES STAKEHOLDER MEETINGS

<table>
<thead>
<tr>
<th>Date/time</th>
<th>Location</th>
<th>Contact No.</th>
</tr>
</thead>
<tbody>
<tr>
<td>August 9, 2018, 9 a.m. Central Time</td>
<td>DoubleTree by Hilton Hotel, Dallas-Market Center, 2015 Market Center Blvd., Dallas, Texas 75207.</td>
<td>214–741–7481</td>
</tr>
<tr>
<td>August 16, 2018, 11 a.m. Eastern Time</td>
<td>Renaissance Reno Downtown Hotel, One South Lake Street, Reno, Nevada 89501.</td>
<td>202–693–9440</td>
</tr>
<tr>
<td>August 21, 2018, 9 a.m. Pacific Time</td>
<td>Renaissance Reno Downtown Hotel, One South Lake Street, Reno, Nevada 89501.</td>
<td>775–682–3900</td>
</tr>
<tr>
<td>September 11, 2018, 9 a.m. Eastern Time</td>
<td>National Mine Health and Safety Academy, 1301 Airport Road, Beckley, West Virginia 25813 (Auditorium)</td>
<td>304–256–3100</td>
</tr>
<tr>
<td>September 20, 2018, 9 a.m. Eastern Time</td>
<td>Hilton Albany, 40 Lodge Street, Albany, New York 12207</td>
<td>518–462–6611</td>
</tr>
<tr>
<td>September 25, 2018, 9 a.m. Eastern Time</td>
<td>Mine Safety and Health Administration (Headquarters), 201 12th Street South, 4E401, Arlington, Virginia 22202.</td>
<td>202–693–9440</td>
</tr>
</tbody>
</table>

II. Background

On June 26, 2018, (83 FR 29716), MSHA published an RFI on Safety Improvement Technologies for Mobile Equipment at Surface Mines, and for Belt Conveyors at Surface and Underground Mines. MSHA is soliciting stakeholder comments, data and information on technologies that can reduce accidents involving mobile equipment at surface mines and belt conveyors at surface and underground mines. Specifically, the Agency is requesting information from the mining community regarding the types of engineering controls available, how to implement such engineering controls, and how these controls could be used in mobile equipment and belt conveyors to reduce accidents, fatalities and injuries. MSHA is also seeking suggestions from stakeholders on best practices, training materials, policies and procedures, innovative technologies, and any other information that stakeholders may have available to improve safety in and around mobile equipment, and working near and around belt conveyors. The meetings will provide the mining community an opportunity to discuss and share information about the issues raised in the RFI. Comments must be received or postmarked by midnight Eastern Standard Time on December 24, 2018.

David G. Zatezalo,
Assistant Secretary of Labor for Mine Safety and Health.

[FR Doc. 2018–15808 Filed 7–24–18; 8:45 am]

BILLING CODE 4520–43–P

FEDERAL COMMUNICATIONS COMMISSION

47 CFR Part 73
[MB Docket Nos. 18–202, 17–105; FCC 18–93]

Children’s Television Programming Rules; Modernization of Media Regulation Initiative

AGENCY: Federal Communications Commission.

ACTION: Proposed rule.

SUMMARY: In this document, the Commission proposes to revise the children’s television programming rules to modify outdated requirements and give broadcasters greater flexibility in serving the educational and informational needs of children. The proposed revisions reflect the dramatic changes in the video programming marketplace since the children’s television programming rules were first adopted more than 20 years ago.

DATES: Comments for this proceeding are due on or before September 24, 2018; reply comments are due on or before October 23, 2018.

ADDRESSES: You may submit comments, identified by MB Docket Nos. 18–202 and 17–105, by any of the following methods:

• Federal Communications Commission’s Website: http://www.fcc.gov/cgb/ecfs/. Follow the instructions for submitting comments.

• Mail: Filings can be sent by hand or messenger delivery, by commercial overnight courier, or by first-class or overnight U.S. Postal Service mail (although the Commission continues to experience delays in receiving U.S. Postal Service mail). All filings must be addressed to the Commission’s Secretary, Office of the Secretary, Federal Communications Commission.

• People with Disabilities: Contact the FCC to request reasonable accommodations (accessible format documents, sign language interpreters, CART, etc.) by email: FCC504@fcc.gov or phone: (202) 418–0530 or TTY: (202) 418–0432.

For detailed instructions for submitting comments and additional information on the rulemaking process, see the SUPPLEMENTARY INFORMATION section of this document.

FOR FURTHER INFORMATION CONTACT: For additional information, contact Kathy Berthot, Kathy.Berthot@fcc.gov; of the Media Bureau, Policy Division, (202) 418–7454.