GA 300° radials. From INT Rome, GA 060° and Volunteer, TN, 197° radials; Volunteer; London, KY; Lexington, KY; Cincinnati, KY; Shelbyville, IN; INT Shelbyville 313° and Boiler, IN, 136° radials; Boiler; Chicago Heights, IL; to INT Chicago Heights 358° and DuPage, IL, 101° radials. From Nodine, MN; to Gopher, MN. The airspace below 2,000 feet MSL outside the United States is excluded.

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V–157 [Amended]

From Key West, FL; INT Key West 038° and Dolphin, FL, 244° radials; to Dolphin. From Lakeland, FL; Ocala, FL; INT Ocala 346° and Taylor, FL, 170° radials; Taylor; to Waycross, GA. From Florence, SC; Fayetteville, NC; Kinston, NC; to Tar River, NC. From Robbinsville, NJ; INT Robbinsville 044° and LaGuardia, NY, 213° radials; LaGuardia; INT LaGuardia 032° and Deer Park, NY, 326° radials; INT Deer Park 326° and Kingston, NY, 191° radials; Kingston; to Albany, NY.

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V–159 [Amended]

From Melbourne, FL 269° T/276° M and Orlando, FL, 140° radials; Orlando; Ocala, FL; Cross City, FL; Greenville, FL; Pecan, GA; Eufaula, AL; INT Eufaula 320° and Vulcan, AL 139° radials to Vulcan. From Holly Springs, MS; Gilmore, AR; Walnut Ridge, AR; Dogwood, MO; Springfield, MO; Napoleon, MO; INT Napoleon 005° and St. Joseph, MO, 122° radials; St. Joseph; to Omaha, IA.

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V–225 [Amended]

From Key West, FL; to Lee County, FL.

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V–295 [Removed]

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V–437 [Amended]

From Melbourne, FL; INT Melbourne 322° and Ormond Beach, FL, 211° radials; Ormond Beach; INT Ormond Beach 360° and Savannah, GA, 177° radials; Savannah; INT Savannah 054° and Charleston, SC, 231° radials; Charleston; to Florence, SC. The airspace within R–2935 is excluded.

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V–492 [Amended]

From Palm Beach, FL; INT Palm Beach 356° and Melbourne, FL, 146° radials, to Melbourne.

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V–521 [Amended]

From Marianna, FL; Wiregrass, AL; INT Wiregrass 333° and Montgomery, AL, 129° radials; Montgomery; INT Montgomery 357° and Vulcan, AL, 139° radials; to Vulcan.

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V–529 [Removed]

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V–537 [Amended]

From INT Melbourne, FL, 269° T/276° M and Orlando, FL, 140° T/140° M radials; INT Orlando 140° and Melbourne 298° radials; INT Melbourne 298° and Ocala, FL 145° radials; Ocala; Gators, FL; to Greenville, FL.

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V–601 [Removed]

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Issued in Washington, DC, on July 11, 2024.

Brian Eric Konie,

Acting Manager, Rules and Regulations Group.

[FR Doc. 2024–15557 Filed 7–17–24; 8:45 am]

BILLING CODE 4910–13–P

CONSUMER PRODUCT SAFETY COMMISSION

16 CFR Part 1215

[Docket No. CPSC–2009–0064]

Notice of Availability and Request for Comment: Revision to the Voluntary Standard for Infant Bath Seats

AGENCY: Consumer Product Safety Commission.

ACTION: Notice of availability and request for comment.

SUMMARY: The U.S. Consumer Product Safety Commission's (Commission or CPSC) mandatory rule, Safety Standard for Infant Bath Seats, incorporates by reference ASTM F1967–19, Standard Consumer Safety Specification for Infant Bath Seats. ASTM notified the Commission that it has revised this incorporated voluntary standard. CPSC seeks comment on whether the revision improves the safety of infant bath seats.

DATES: Comments must be received by August 1, 2024.

ADDRESSES: You can submit comments, identified by Docket No. CPSC–2009–0064, by any of the following methods:

Electronic Submissions: Submit electronic comments to the Federal eRulemaking Portal at: *https://www.regulations.gov*. Follow the instructions for submitting comments. Do not submit through this website: confidential business information, trade secret information, or other sensitive or protected information that you do not want to be available to the public. CPSC typically does not accept comments submitted by email, except as described below.

Mail/Hand Delivery/Courier/Confidential Written Submissions: CPSC encourages you to submit electronic comments by using the Federal eRulemaking Portal. You may, however, submit comments by mail, hand delivery, or courier to: Office of the Secretary, Consumer Product Safety Commission, 4330 East West Highway, Bethesda, MD 20814; telephone: (301) 504–7479. If you wish to submit confidential business information, trade secret information, or other sensitive or protected information that you do not want to be available to the public, you may submit such comments by mail, hand delivery, or courier, or you may email them to: *cpsc-os@cpsc.gov*.

Instructions: All submissions must include the agency name and docket number. CPSC may post all comments without change, including any personal identifiers, contact information, or other personal information provided, to: *https://www.regulations.gov*. Do not submit to this website: confidential business information, trade secret information, or other sensitive or protected information that you do not want to be available to the public. If you wish to submit such information, please submit it according to the instructions for mail/hand delivery/courier/confidential written submissions.

Docket: For access to the docket to read background documents or comments received, go to: *https://www.regulations.gov,* and insert the docket number, CPSC–2009–0064, into the ''Search'' box, and follow the prompts.

FOR FURTHER INFORMATION CONTACT: Julia Kerns, Directorate for Engineering Sciences, U.S. Consumer Product Safety Commission, 5 Research Place, Rockville, MD 20850; telephone: (301) 987–2548; email: *JKerns@cpsc.gov*.

SUPPLEMENTARY INFORMATION: Section 104(b) of the Consumer Product Safety Improvement Act of 2008 (CPSIA) requires the Commission to adopt mandatory standards for durable infant or toddler products. 15 U.S.C. 2056a(b)(1). Mandatory standards must be ''substantially the same as'' voluntary standards, or they may be ''more stringent'' than the applicable voluntary standards, if the Commission determines that more stringent requirements would further reduce the risk of injury associated with the products. *Id.* Mandatory standards may be based, in whole or in part, on a voluntary standard.

Section 104(b)(4)(B) of the CPSIA specifies the process for when a voluntary standards organization revises a standard that the Commission incorporated by reference under section 104(b)(1). First, the voluntary standards organization must notify the Commission of the revision. Once the Commission receives this notification, the Commission may reject or accept the revised standard. To reject a revised standard, the Commission must notify the voluntary standards organization

within 90 days of receiving the notice that it has determined that the revised standard does not improve the safety of the consumer product and that it is retaining the existing standard. If the Commission does not take this action, the revised voluntary standard will be considered a consumer product safety standard issued under section 9 of the Consumer Product Safety Act (15 U.S.C. 2058), effective 180 days after the Commission received notification of the revision (or a later date specified by the Commission in the **Federal Register**). 15 U.S.C. 2056a(b)(4)(B).

In 2010, the Commission adopted a mandatory rule for infant bath seats under section 104(b)(1) of the CPSIA, which was codified in 16 CFR part 1215. The rule incorporated by reference ASTM F1967–08a, *Standard Consumer Safety Specification for Infant Bath Seats,* with modifications to make the standard more stringent. 75 FR 31691 (June 4, 2010). At the time the Commission published the final rule, ASTM F1967–08a was the current version of the voluntary standard. ASTM subsequently revised the voluntary standard five times. ASTM F1967 applies to infant bath seats, which it describes as products used in a bathtub, sink, or similar bathing enclosure and that provide support, at a minimum, to the front and back of a seated infant during bathing by a caregiver. The ASTM standard includes performance requirements, test methods, and labeling requirements to address hazards to infants associated with infant bath seats. After the Commission adopted the mandatory standard in 2010, the Commission updated the standard in 2012, 2013, and 2019 and the mandatory standard currently incorporates by reference ASTM F1967–19. 84 FR 49435 (September 20, 2019).

On July 8, 2024, ASTM notified CPSC that it had approved and published ASTM F1967–24. CPSC staff is assessing the revised voluntary standard to determine, consistent with section 104(b)(4)(B) of the CPSIA, its effect on the safety of consumer products covered by the standard. The Commission invites public comment on that question, to inform staff's assessment and any subsequent Commission consideration of the revisions in ASTM F1967–24.[1]

[1] The Commission voted (5–0) to approve this notice.

The currently incorporated voluntary standard (ASTM F1967–19) and the revised voluntary standard (ASTM F1967–24) are available for review in several ways. A read-only copy of the existing, incorporated standard (ASTM F1967–19) is available for viewing, at no cost, on the ASTM website at: *https://www.astm.org/READINGLIBRARY/*. A read-only copy of the revised standard (ASTM F1967–24), including red-lined versions that identify the changes from the 2019 version to the 2024 version, are available, at no cost, on ASTM's website at: *https://www.astm.org/CPSC.htm*. Interested parties can also download copies of the standards by purchasing them from ASTM International, 100 Barr Harbor Drive, P.O. Box C700, West Conshohocken, PA 19428–2959; phone: 610–832–9585; *https://www.astm.org*. Alternatively, interested parties can schedule an appointment to inspect copies of the standards at CPSC's Office of the Secretary, U.S. Consumer Product Safety Commission, 4330 East West Highway, Bethesda, MD 20814, telephone: 301–504–7479.

Comments must be received by August 1, 2024. Because of the short statutory time frame Congress established for the Commission to consider revised voluntary standards under section 104(b)(4) of the CPSIA, CPSC will not consider comments received after this date.

Alberta E. Mills,

Secretary, Consumer Product Safety Commission.

[FR Doc. 2024–15843 Filed 7–17–24; 8:45 am]

BILLING CODE 6355–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 73

[Docket No. FDA–2024–C–3229]

Phytolon Ltd.; Filing of Color Additive Petition

AGENCY: Food and Drug Administration, HHS.

ACTION: Notification of petition.

SUMMARY: The Food and Drug Administration (FDA or we) is announcing that we have filed a petition, submitted by Phytolon Ltd., proposing that the color additive regulations be amended to provide for the safe use of prickly pear yellow for the coloring of foods generally in amounts consistent with current good manufacturing practice.

DATES: The color additive petition was filed on July 3, 2024.

ADDRESSES: For access to the docket to read background documents or 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.

FOR FURTHER INFORMATION CONTACT: Kaiping Deng, Center for Food Safety and Applied Nutrition, Food and Drug Administration, 5001 Campus Dr., College Park, MD 20740, 708–924–0622.

SUPPLEMENTARY INFORMATION: Under section 721(d)(1) of the Federal Food, Drug, and Cosmetic Act ((21 U.S.C. 379e(d)(1)), we are giving notice that we have filed a color additive petition (CAP 4C0332), submitted by Phytolon Ltd., Ha-Tsmikha St, Yokne'am Illit, Israel. The petition proposes to amend the color additive regulations in 21 CFR part 73, ''Listing of Color Additives Exempt From Certification,'' to provide for the safe use of prickly pear yellow for the coloring of foods generally in amounts consistent with current good manufacturing practice.

The petitioner has claimed that this action is categorically excluded under 21 CFR 25.32(r), which applies to an action for substances which occur naturally in the environment, and for which the action does not significantly alter the concentration or distribution of the substance, its metabolites, or degradation products in the environment. In addition, the petitioner has stated that, to their knowledge, no extraordinary circumstances exist. If FDA determines a categorical exclusion applies, neither an environmental assessment nor an environmental impact statement is required. If FDA determines a categorical exclusion does not apply, we will request an environmental assessment and make it available for public inspection.

Dated: July 15, 2024.

Lauren K. Roth,

Associate Commissioner for Policy.

[FR Doc. 2024–15892 Filed 7–17–24; 8:45 am]

BILLING CODE 4164–01–P