others by written request to terminate or modify the agreement.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration [Docket No. FDA-2009-D-0224]

Small Entity Compliance Guide: Bottled Water: Residual Disinfectants and Disinfection Byproducts; Availability

AGENCY: Food and Drug Administration,

HHS.

ACTION: Notice.

SUMMARY: The Food and Drug
Administration (FDA) is announcing the
availability of a guidance for industry
entitled "Bottled Water: Residual
Disinfectants and Disinfection
Byproducts—Small Entity Compliance
Guide" for a direct final rule published
in the Federal Register of March 28,
2001. This small entity compliance
guide (SECG) is intended to set forth in
plain language the requirements of the
regulation and to help small businesses
understand the regulation.

DATES: Submit written or electronic comments on the SECG at any time.

ADDRESSES: Submit written comments on the SECG to the Division of Dockets Management (HFA-305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. Submit electronic comments on the SECG to http://www.regulations.gov. Submit written requests for single copies of the SECG to the Division of Plant and Dairy Food Safety (HFS-317), Office of Food Safety, Center for Food Safety and Applied Nutrition, Food and Drug Administration, 5100 Paint Branch Pkwy., College Park, MD 20740, or fax your request to 301-436-2651. Send one self-addressed adhesive label to assist that office in processing your request. See the SUPPLEMENTARY INFORMATION section for electronic access to the SECG.

FOR FURTHER INFORMATION CONTACT:

Lauren Posnick Robin, Center for Food Safety and Applied Nutrition (HFS– 317), Food and Drug Administration, 5100 Paint Branch Pkwy., College Park, MD 20740, 301–436–1639.

SUPPLEMENTARY INFORMATION:

I. Background

In the **Federal Register** of March 28, 2001 (66 FR 16858), FDA issued a direct final rule amending its bottled water

quality standard regulations by establishing allowable levels for three residual disinfectants (chloramine, chlorine, and chlorine dioxide) and three types of disinfection byproducts (DBPs) (bromate, chlorite, and haloacetic acids (HAA5)), and by revising the existing allowable level for the DBP total trihalomethanes (TTHM). FDA also revised, for the three residual disinfectants and four types of DBPs only, the monitoring requirement for source water found in the current good manufacturing practice (CGMP) regulations for bottled water. On July 5, 2001 (66 FR 35373), FDA issued a technical amendment to correct an editorial error and confirmed the effective date of January 1, 2002.

FDA examined the economic implications of the direct final rule as required by the Regulatory Flexibility Act (5 U.S.C. 601-612) and determined that the rule would have a significant economic impact on a substantial number of small entities. In compliance with section 212 of the Small Business Regulatory Enforcement Fairness Act (Public Law 104-121), FDA is making available this SECG stating in plain language the legal requirements of the March 28, 2001, direct final rule set forth in 21 CFR parts 129 and 165 concerning the allowable levels and monitoring requirements for the three residual disinfectants (chloramine, chlorine, and chlorine dioxide) and four types of DBPs (bromate, chlorite, HAA5, and TTHM).

FDA is issuing this SECG as level 2 guidance consistent with FDA's good guidance practices regulation (21 CFR 10.115(c)(2)). The SECG represents the agency's current thinking on this topic. It does not create or confer any rights for or on any person and does not operate to bind FDA or the public. An alternative approach may be used if such approach satisfies the requirements of the applicable statutes and regulations.

II. Comments

Interested persons may submit to the Division of Dockets Management (see ADDRESSES) written or electronic comments regarding this SECG. Submit a single copy of electronic comments or two paper copies of any mailed comments, except that individuals may submit one paper copy. Comments are to be identified with the docket number found in brackets in the heading of this document. The SECG and received comments may be seen in the Division of Dockets Management between 9 a.m. and 4 p.m., Monday through Friday.

III. Electronic Access

Persons with access to the Internet may obtain the document at http://www.cfsan.fda.gov/guidance.html or http://www.regulations.gov.

Dated: May 22, 2009.

Jeffrey Shuren,

Associate Commissioner for Policy and Planning.

[FR Doc. E9–12671 Filed 5–29–09; 8:45 am] BILLING CODE 4160–01–S

DEPARTMENT OF HOMELAND SECURITY

[Docket No. DHS-2009-0079]

The National Infrastructure Advisory Council

AGENCY: Directorate for National Protection and Programs, Department of Homeland Security.

ACTION: Committee Management; Notice of Federal Advisory Council Meeting

SUMMARY: The National Infrastructure Advisory Council (NIAC) will meet on Tuesday July 14, 2009 at the J.W. Marriott's Salons E and F, 1331 Pennsylvania Avenue, Washington, DC 20004.

DATES: The National Infrastructure Advisory Council will meet Tuesday July 14, 2009 from 12:30 p.m. to 3:30 p.m. Please note that the meeting may close early if the committee has completed its business.

For additional information, please consult the NIAC Web site, http://www.dhs.gov/niac, or contact Matthew Sickbert by phone at 703–235–2888 or by e-mail at

Matthew.Sickbert@associates.dhs.gov.

ADDRESSES: The meeting will be held at the J.W. Marriott's Salons E and F, 1331 Pennsylvania Avenue, Washington, DC 20004. While we will be unable to accommodate oral comments from the public, written comments may be sent to Nancy J. Wong, Department of Homeland Security, Directorate for National Protection and Programs, Washington, DC 20528. Written comments should reach the contact person listed no later than July 7, 2009. Comments must be identified by DHS–2009–0079 and may be submitted by one of the following methods:

- Federal eRulemaking Portal: http://www.regulations.gov. Follow the instructions for submitting comments.
- *E-mail:* matthew.sickbert@associates.dhs.gov. Include the docket number in the subject line of the message.
 - Fax: 703–235–3055