

goals, such as reducing new HIV infections, increasing the use of condoms, and targeting high risk groups by describing and monitoring the HIV risk behaviors, HIV seroprevalence and incidence, and HIV prevention experiences of persons at highest risk for HIV infection.

The Centers for Disease Control and Prevention request two year approval for a new information collection. Data will be collected through anonymous, in-person interviews conducted with persons systematically selected from nine Metropolitan Statistical Areas (MSAs) throughout the United States; these nine MSAs were chosen based on having high HIV prevalence. A brief screening interview will be used to determine eligibility for participation in the behavioral assessment. Participants will be recruited through respondent-driven sampling, a scientifically proven recruitment strategy for reaching hidden, hard-to-reach, or stigmatized populations. Interview data will be recorded on secure portable computers, without internet connections. Data will

be transferred to secure, encrypted data servers. Data will be stored at CDC and shared with local health departments in accordance with existing data use agreements and the Assurance of Confidentiality for HIV/AIDS Surveillance Data. Data will be disseminated in aggregate through academic and agency publications, presentations, and reports. All data collection and activities will be anonymous.

Personally identifiable information (PII) is not included in the data collection. The CDC Privacy Officer has assessed this package for applicability of 5 U.S.C. 552a. The Privacy Act is not applicable because PII is not being collected under this CDC funded activity. The NHBS-Trans formative interview and optional HIV testing are anonymous (neither names nor Social Security numbers are collected). Data that will be collected through NHBS-Trans, while sensitive, are not personally identifying.

The data from the behavioral assessment will provide estimates of (1)

behavior related to the risk of HIV and other sexually transmitted diseases, (2) prior testing for HIV, (3) and use of HIV prevention services. All persons interviewed will also be offered an HIV test, and will participate in a pre-test counseling session. No other federal agency systematically collects this type of information from persons at risk for HIV infection. These data have substantial impact on prevention program development and monitoring at the local, state, and national levels.

The Burden Table below shows the estimated annualized burden hours for the participants' time. Annually, 990 participants will complete an eligibility screener (an average of 5 minutes to complete), 900 participants will complete the Behavioral Assessment (an average of 40 minutes to complete), and 900 will complete the Recruiter Debriefing Form (an average of two minutes to complete). The estimated total annualized burden would be 713 hours. Participation of respondents is voluntary. There are no costs to respondents other than their time.

#### ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total burden (in hours)
Persons Screened .....	Eligibility Screener .....	990	1	5/60	83
Eligible Participant .....	Behavioral Assessment .....	900	1	40/60	600
Peer Recruiters .....	Recruiter Debriefing .....	900	1	2/60	30
Total .....	.....	.....	.....	.....	713

Jeffrey M. Zirger,

Acting Chief, Information Collection Review Office, Office of Scientific Integrity, Office of the Associate Director for Science, Office of the Director, Centers for Disease Control and Prevention.

[FR Doc. 2018-11399 Filed 5-25-18; 8:45 am]

BILLING CODE 4163-18-P

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

[Docket No. FDA-2018-D-1468]

#### Registration of Food Facilities: What You Need To Know About the Food and Drug Administration Regulation; Small Entity Compliance Guide; Availability

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Notification of availability.

**SUMMARY:** The Food and Drug Administration (FDA or we) is announcing the availability of a final guidance for industry entitled "Registration of Food Facilities: What You Need To Know About the FDA Regulation—Small Entity Compliance Guide." The small entity compliance guide (SECG) is intended to help small entities comply with a final rule we issued in the **Federal Register** of July 14, 2016, entitled "Amendments to Registration of Food Facilities." The final rule amends the registration of food facilities regulations.

**DATES:** The announcement of the guidance is published in the **Federal Register** on May 29, 2018.

**ADDRESSES:** You may submit either electronic or written comments on Agency guidances at any time as follows:

#### 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

- *Mail/Hand delivery/Courier (for written/paper submissions):* Dockets Management Staff (HFA-305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

- For written/paper comments submitted to the Dockets Management Staff, FDA will post your comment, as well as any attachments, except for information submitted, marked and identified, as confidential, if submitted as detailed in “Instructions.”

*Instructions:* All submissions received must include the Docket No. FDA-2018-D-1468 for “Registration of Food Facilities: What You Need To Know About the FDA Regulation—Small Entity Compliance Guide.” Received comments will be placed in the docket and, except for those submitted as “Confidential Submissions,” publicly viewable at <https://www.regulations.gov> or at the Dockets Management Staff between 9 a.m. and 4 p.m., Monday through Friday.

- *Confidential Submissions*—To submit a comment with confidential information that you do not wish to be made publicly available, submit your comments only as a written/paper submission. You should submit two copies total. One copy will include the information you claim to be confidential with a heading or cover note that states “THIS DOCUMENT CONTAINS CONFIDENTIAL INFORMATION.” We will review this copy, including the claimed confidential information, in our consideration of comments. The second copy, which will have the claimed confidential information redacted/blacked out, will be available for public viewing and posted on <https://www.regulations.gov>. Submit both copies to the Dockets Management Staff. If you do not wish your name and contact information to be made publicly available, you can provide this information on the cover sheet and not in the body of your comments and you must identify this information as “confidential.” Any information marked as “confidential” will not be disclosed except in accordance with 21 CFR 10.20 and other applicable disclosure law. For more information about FDA’s posting of comments to public dockets, see 80 FR 56469, September 18, 2015, or access the information at: <https://www.gpo.gov/fdsys/pkg/FR-2015-09-18/pdf/2015-23389.pdf>.

*Docket:* For access to the docket to read background documents or the electronic and written/paper 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.

You may submit comments on any guidance at any time (see 21 CFR 10.115(g)(5)).

Submit written requests for single copies of the SECG to the Office of Compliance, Center for Food Safety and Applied Nutrition (HFS-800), Food and Drug Administration, 5001 Campus Dr., College Park, MD 20740. Send two self-addressed adhesive labels to assist that office in processing your request. See the **SUPPLEMENTARY INFORMATION** section for electronic access to the SECG.

**FOR FURTHER INFORMATION CONTACT:** Courtney Buchanan, Center for Food Safety and Applied Nutrition, Food and Drug Administration, 5001 Campus Dr., College Park, MD 20740, 240-402-2487.

#### **SUPPLEMENTARY INFORMATION:**

##### **I. Background**

The Public Health Security and Biodefense Act of 2002 (Biodefense Act) (Pub. L. 107-188) added section 415 to the Federal Food, Drug, and Cosmetic Act (FD&C Act). Section 415 of the FD&C Act (21 U.S.C. 350d) requires domestic and foreign facilities that manufacture, process, pack, or hold food for human or animal consumption in the United States to register with FDA.

On October 10, 2003, we issued an interim final rule (68 FR 58894) to implement section 415 of the FD&C Act. That rule established the food facility registration regulations in part 1, subpart H (21 CFR 1.225 through 1.243). Previously, this guidance restated FDA’s food facility registration regulations. This guidance also served as FDA’s SECG for part 1, subpart H.

The FDA Food Safety Modernization Act (FSMA), enacted on January 4, 2011, amended section 415 of the FD&C Act to require that facilities engaged in manufacturing, processing, packing, or holding food for consumption in the United States submit additional registration information to FDA, including an assurance that FDA will be permitted to inspect the facility at the times and in the manner permitted by the FD&C Act. Section 415 of the FD&C Act, as amended by FSMA, also requires food facilities required to register with

FDA to renew such registrations every other year, and provides FDA with authority to suspend the registration of a food facility in certain circumstances.

On July 14, 2016, we published a final rule (81 FR 45912) that amended our food facility registration regulations to reflect, among other things, the FSMA amendments to section 415 of the FD&C Act. Accordingly, FDA is revising this SECG to provide guidance intended to help small entities comply with the revised registration of food facilities requirements in part 1, subpart H.

We examined the economic implications of the final rule as required by the Regulatory Flexibility Act (5 U.S.C. 601–612). Although we stated that we did not believe that the final rule that amended our registration of food facilities regulations would have a significant economic impact on a substantial number of small entities, we analyzed various regulatory options to examine the impact on small entities. Consistent with section 212 of the Small Business Regulatory Enforcement Fairness Act (Pub. L. 104–121, as amended by Pub. L. 110–28), we are making available the SECG to explain the actions that a small entity must take to comply with the rule.

We are issuing the SECG consistent with our good guidance practices regulation (21 CFR 10.115(c)(2)). The SECG represents the current thinking of FDA on this topic. It does not establish any rights for any person and is not binding on FDA or the public. You can use an alternative approach if it satisfies the requirements of the applicable statutes and regulations. This guidance is not subject to Executive Order 12866.

##### **II. Paperwork Reduction Act of 1995**

The guidance refers to previously approved collections of information found in FDA regulations. The collections of information in part 1, subpart H have been approved under the Office of Management and Budget control number 0910-0502.

##### **III. Electronic Access**

Persons with access to the internet may obtain the SECG at either <https://www.fda.gov/FoodGuidances> or <https://www.regulations.gov>. Use the FDA website listed in the previous sentence to find the most current version of the guidance.

Dated: May 22, 2018.

**Leslie Kux,**

*Associate Commissioner for Policy.*

[FR Doc. 2018–11419 Filed 5–25–18; 8:45 am]

**BILLING CODE 4164-01-P**