

FDA has drafted the set of questions below to help focus comments presented at the public meeting or otherwise communicated to the agency.

II. Questions

1. What concepts or underlying principles should guide the 2004 Produce Safety Action Plan? Are the seven objectives in the working draft appropriate for achieving the overarching goal to minimize foodborne illness associated with the consumption of fresh produce?

2. What major practices contribute to the contamination of fresh produce by harmful pathogens? What intervention strategies will prevent, reduce, or control this contamination?

3. The produce action plan covers fresh fruits and vegetables that have not been heat treated to reduce, control, or eliminate pathogens, or otherwise significantly processed. The draft action plan is not intended to cover frozen fruits and vegetables, fruit and vegetable juices, or other commodities such as tree nuts that are neither fruits nor vegetables and not typically regarded as produce. Should the produce action plan cover additional foods? If so, which foods?

4. What measurements should be used to measure progress toward the overarching goal (to minimize foodborne illness associated with fresh produce consumption)? What measures should be used to measure progress toward the individual objectives?

5. Does FDA's current GAPs/GMPs guidance (<http://www.foodsafety.gov/~dms/prodguid.html>) need to be expanded or otherwise revised? If yes, please describe generally the areas that need expansion or other revision.

6. In today's production and food preparation environments (farms, packing houses, retail establishments, and consumers), what conditions, practices, or other factors are the principal contributors to contamination of produce with a pathogen? What interventions would reduce, control, or eliminate this contamination?

7. There is broad variation within food operations including variations in size of establishments, the nature of the commodity produced, the practices used in production, and the vulnerability of a particular commodity to microbial hazards. How, if at all, should the produce action plan be structured to take into account such variation? For example, should there be different sets of interventions for identifiable segments of the fresh produce industry?

8. What roles can and should Federal, State, and local agencies and the food

industry play in developing and implementing action items to help achieve the objectives in this action plan?

9. Are there existing food safety systems or standards (such as international standards) that FDA should consider as part of the agency's development and implementation of a produce safety action plan? Please identify these systems or standards and explain what their consideration might contribute to this effort.

III. Registration

You may register through FDA's Web site <http://www.cfsan.fda.gov/> and choose "Public Meetings," by fax, or e-mail (see **FOR FURTHER INFORMATION CONTACT**). For security reasons and due to space limitations, we recommend that you register at least 5 days prior to the meeting. Registration will be accepted on a first-come-first-serve basis. You may register until close of business June 22, 2004. If you need special accommodations due to a disability, please inform the contact person at least 7 days in advance (see **FOR FURTHER INFORMATION CONTACT**). There is no registration fee for this public meeting, but early registration is encouraged because space is limited, and it will expedite entry into the building and its parking area. If you require parking, please include the vehicle make and tag number, if known, on your registration form. Because the meeting will be held in a Federal building, you should also bring a photo identification (ID) and plan for adequate time to pass through security screening systems. If you would like to make oral comments at the meeting, please specify your interest in speaking when you register. The amount of time for each oral presentation may be limited based upon the number of requests to speak. FDA encourages individuals or firms with relevant data or information to present such information at the meeting or in written comments to the record.

IV. Transcripts

A transcript will be made of the proceedings of the meeting. You may request a copy of the meeting transcript from FDA's Freedom of Information Office (HFI-35), Food and Drug Administration, 5600 Fishers Lane, rm. 12A-16, Rockville, MD 20857, approximately 30 working days after the meeting at a cost of 10 cents a page. The transcript of the public meeting and all comments submitted will be available for public examination at the Division of Dockets Management (see **ADDRESSES**) between 9 a.m. and 4 p.m., Monday through Friday.

V. Comments

In addition to presenting oral comments at the public meeting, interested persons may submit to the Division of Dockets Management (see **ADDRESSES**) written or electronic comments on the subject of this meeting. 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 the brackets in the heading of this document. Received comments may be seen in the Division of Dockets Management between 9 a.m. and 4 p.m., Monday through Friday.

Dated: June 9, 2004.

Jeffrey Shuren,

Assistant Commissioner for Policy.

[FR Doc. 04-13544 Filed 6-10-04; 1:35 pm]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Substance Abuse and Mental Health Services Administration

Agency Information Collection Activities: Proposed Collection; Comment Request

In compliance with section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995 concerning opportunity for public comment on proposed collections of information, the Substance Abuse and Mental Health Services Administration will publish periodic summaries of proposed projects. To request more information on the proposed projects or to obtain a copy of the information collection plans, call the SAMHSA Reports Clearance Officer on (301) 443-7978.

Comments are invited on: (a) Whether the proposed collections of information are necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) ways to enhance the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology.

Proposed Project: Reach Out Now National Teach-In Initiative Feedback Form

(OMB No. 0930-0258; Extension, no change)—Under section 515(b) of the Public Health Service Act (42 U.S.C. 290bb-21), the Center for Substance Abuse Prevention (CSAP) of the Substance Abuse and Mental Health Services Administration (SAMHSA) is directed to develop effective alcohol abuse prevention literature and, to assure the widespread dissemination of prevention materials among States, political subdivisions, and school

systems. Each April, SAMHSA collaborates with Scholastic Inc. in the April distribution of *Reach Out Now: Talk to Your Fifth Grader About Underage Alcohol Use*, a supplement created and distributed by Scholastic Inc.

Beginning in April 2004, SAMHSA sponsors an annual national Teach-In to foster a conversation with fifth graders on the dangers of early alcohol use. State substance abuse prevention directors nominate organizations to participate in this program. The Teach-In program builds upon the highly successful national initiative of the

Leadership to Keep Children Alcohol Free, which is focused on preventing alcohol use among children ages 9 to 15 and is spearheaded by more than 40 current and past Governors' spouses, who have held or supported Reach Out Now Teach-Ins in their States.

Organizations that agree to participate in this SAMHSA initiative are asked to provide feedback information about the implementation and results of the Teach-In event in their community school. The table that follows provides an estimate of the annual response burden for the feedback form.

Number of respondents	Responses/respondent	Burden/response (hrs.)	Total burden hours
200	1	.167	34

Send comments to Nancy Pearce, SAMHSA Reports Clearance Officer, Room 16-105, Parklawn Building, 5600 Fishers Lane, Rockville, MD 20857. Written comments should be received by August 16, 2004.

Dated: June 7, 2004.

Patricia S. Bransford,

Acting Executive Officer, SAMHSA.

[FR Doc. 04-13395 Filed 6-14-04; 8:45 am]

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DEPARTMENT OF HOMELAND SECURITY

Federal Emergency Management Agency

Agency Information Collection Activities: Submission for OMB Review; Comment Request

AGENCY: Federal Emergency Management Agency, Emergency Preparedness and Response Directorate, U.S. Department of Homeland Security.

ACTION: Notice and request for comments.

SUMMARY: The Federal Emergency Management Agency has submitted the following proposed information collection to the Office of Management and Budget (OMB) for review and clearance in accordance with the requirements of the Paperwork Reduction Act of 1995 (44 U.S.C. 3507).

The submission describes the nature of the information collection, the categories of respondents, the estimated burden (*i.e.*, the time, effort and resources used by respondents to respond) and cost, and includes the actual data collection instruments FEMA will use.

Title: EMI Independent Study Course Enrollment Application.

Type of Information Collection: Revision of a currently approved collection.

OMB Number: 1660-0046.

Form: FEMA Form 95-23.

Abstract: The purpose of this form is to collect information from individuals on what Independent Study courses they wish to enroll in. This form lists the courses available through FEMA's Independent Study Program and collects information from individuals so that these courses can be mailed to them.

Affected Public: Individuals or households.

Number of Respondents: 187,000.

Estimated Time per Respondent: 1 minute or .016666 hour.

Estimated Total Annual Burden Hours: 3,116 hours.

Frequency of Response: Other—As needed for participants to enroll in Independent Study courses.

Cost to Respondents: Annualized cost to all respondents is estimated at \$81,937. This figure is composed of two items: \$54,261 corresponding to the time spent completing the form at the national mean hourly rate of \$17.41 and \$27,676 comprising the cost of a regular postage stamp of \$0.37 for 74,800 respondents (40% of applications) who chose to apply via mail.

Comments: Interested persons are invited to submit written comments on the proposed information collection to the Office of Information and Regulatory Affairs at OMB, Attention: Desk Officer for the Emergency Preparedness and Response Directorate/Federal Emergency Management Agency, Department of Homeland Security, 725 17th Street, NW., Docket Library Room

10102, Washington, DC 20503. Comments must be submitted on or before July 15, 2004. In addition, interested persons may also send comments to FEMA (see contact information below).

FOR FURTHER INFORMATION CONTACT:

Requests for additional information or copies of the information collection should be made to Muriel B. Anderson, Chief, Records Management, FEMA at 500 C Street, SW., Room 316, Washington, DC 20472, facsimile number (202) 646-3347, or e-mail address FEMA-Information-Collections@dhs.gov.

Dated: June 2, 2004.

Edward W. Kernan,

Branch Chief, Information Resources Management Branch, Information Technology Services Division.

[FR Doc. 04-13456 Filed 6-14-04; 8:45 am]

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DEPARTMENT OF HOMELAND SECURITY

Federal Emergency Management Agency

Agency Information Collection Activities: Submission for OMB Review; Comment Request

AGENCY: Federal Emergency Management Agency, Emergency Preparedness and Response Directorate, U.S. Department of Homeland Security.

ACTION: Notice and request for comments.

SUMMARY: The Federal Emergency Management Agency (FEMA) has submitted the following information collection to the Office of Management