

found in the brackets in the heading of this document. A copy of the document and received comments are available for public examination in the Dockets Management Branch 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 either <http://www.fda.gov/cber/guidelines.htm> or <http://www.fda.gov/ohrms/dockets/default.htm>.

Dated: August 20, 2002.

Margaret M. Dotzel

Associate Commissioner for Policy.

[FR Doc. 02-23105 Filed 9-11-02; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 02D-0333]

Draft Guidance for Industry: Juice HACCP Hazards and Controls Guidance, First Edition; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing the availability of a draft guidance document entitled "Guidance for Industry: Juice HACCP Hazards and Controls Guidance" (first edition) (the draft guidance). The draft guidance supports and complements the FDA regulation that requires a processor of juice to evaluate its operations using Hazard Analysis Critical Control Point (HACCP) principles and, if necessary, to develop and implement HACCP systems for its operations. The draft guidance represents FDA's views on potential hazards in juice products and how to control them, and it is designed to assist juice processors in the development of HACCP plans.

DATES: Submit written or electronic comments concerning the draft guidance and collection of information by November 12, 2002, to ensure adequate consideration in the preparation of the final guidance document. Comments on the draft guidance may be submitted at any time.

ADDRESSES: Submit written requests for single copies of the draft guidance to Michael E. Kashtock, Center for Food Safety and Applied Nutrition (see **FOR FURTHER INFORMATION CONTACT**). Send two self-addressed adhesive labels to assist that office in processing your

request. Submit written comments concerning the draft guidance to the Dockets Management Branch (HFA-305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. Submit electronic comments to <http://www.fda.gov/dockets.ecomments>. See the **SUPPLEMENTARY INFORMATION** section for electronic access to the draft guidance document.

FOR FURTHER INFORMATION CONTACT:

Michael E. Kashtock, Center for Food Safety and Applied Nutrition (HFS-305), Food and Drug Administration, 5100 Paint Branch Pkwy., College Park, MD 20740-3835, 301-436-2022, FAX 301-436-2651, e-mail: mkashtoc@cfstan.fda.gov.

SUPPLEMENTARY INFORMATION:

I. Background

FDA is announcing the availability of the first edition of the draft guidance entitled "Guidance for Industry: Juice HACCP Hazards and Controls Guidance."

Under the HACCP regulations in part 120 (21 CFR part 120), juice processors are required to evaluate their operations using HACCP principles and, if necessary, to develop and implement HACCP systems for their operations. Under § 120.9, juice products are adulterated under section 402(a)(4) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 342) if a processor or importer fails to have and implement a HACCP plan when one is necessary, or otherwise fails to meet any of the requirements of the regulations. The primary purpose of the draft guidance is: (1) To help processors and importers of juice products identify the likelihood that a food safety hazard may occur in their product, and (2) to guide them in the preparation of appropriate HACCP plans for those hazards that are reasonably likely to occur.

II. Significance of the Guidance

The draft guidance is a level 1 guidance issued consistent with FDA's good guidance practices regulations (21 CFR 10.115). The draft guidance represents the agency's current thinking on the potential hazards that are associated with various juice products and processing operations, and how the occurrence of these hazards can be avoided with HACCP controls when they are reasonably likely to occur, as required under part 120. 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 statute and regulations.

This draft guidance contains information collection provisions that are subject to review by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995 (44 U.S.C. 3501-3520). The collection of information in the draft guidance has been submitted to OMB for review and was approved under OMB control number 0910-0466.

III. Comments

Interested persons may submit to the Dockets Management Branch (see **ADDRESSES**) written or electronic comments on the draft guidance. Two copies of any comments are to be submitted, except that individuals may submit one copy. Comments are to be identified with the docket number found in brackets in the heading of this document. The draft guidance and received comments are available for public examination in the Dockets Management Branch between 9 a.m. and 4 p.m. Monday through Friday.

IV. Electronic Access

Persons with access to the Internet may obtain the draft guidance document at <http://www.cfsan.fda.gov/~dms/guidance.html>.

Dated: August 29, 2002.

Margaret M. Dotzel,

Associate Commissioner for Policy.

[FR Doc. 02-23106 Filed 9-11-02; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Substance Abuse and Mental Health Services Administration

The President's New Freedom Commission on Mental Health; Notice of Meeting

Pursuant to Executive Order 13263, notice is hereby given of a meeting of the President's New Freedom Commission on Mental Health in October 2002.

The meeting will be open and will consider how to accomplish the Commission's mandate to conduct a comprehensive study of the United States mental health service delivery system and to make recommendations on improving the delivery of public and private mental health services for adults and children. The Commission meeting will focus on issues relating to the Interim Report, which the Commission is required to send to the President by the end of October.

Attendance by the public will be limited to space available. Public comments are welcome. Please communicate with the individual listed as contact below to make arrangements to comment or to request special accommodations for persons with disabilities.

Additional information and a roster of Commission members may be obtained either by accessing the Commission Web site, www.mentalhealthcommission.gov, or by communicating with the contact whose name and telephone number is listed below.

Committee Name: The President's New Freedom Commission on Mental Health.

Meeting Date/Time:

Open: October 2, 2002, 4:30 p.m. to 6 p.m.

Open: October 3, 2002, 8:30 a.m. to 4:30 p.m.

Open: October 4, 2002, 8:30 a.m. to 1:30 p.m.

Place: Crystal Gateway Marriott, 1700 Jefferson Davis Highway, Arlington, Virginia 22202.

Contact: Claire Heffernan, Executive Secretary, 5600 Fishers Lane, Parklawn Building, Room 13C-26, Rockville, MD 20857, Telephone: (301) 443-1545; Fax: (301) 480-1554 and, e-mail: Cheffern@samhsa.gov, Web site: www.mentalhealthcommission.gov.

Dated: September 6, 2002.

Toian Vaughn,

Committee Management Officer, Substance Abuse and Mental Health Services Administration.

[FR Doc. 02-23107 Filed 9-11-02; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Substance Abuse and Mental Health Services Administration

Statement of Organization, Functions, and Delegations of Authority

Part M of the Substance Abuse and Mental Health Services Administration (SAMHSA) Statement of Organization, Functions, and Delegations of Authority for the Department of Health and Human Services as amended most recently at 67 FR 45136, July 8, 2002 is amended to: Abolish the Office of Evaluation, Scientific Analysis, and Synthesis (OESAS); replace the functional statement of two divisions of the Center for Substance Abuse Treatment (CSAT), the Division of State and Community Assistance and the Division of Services Improvement. The changes are to update and realign CSAT organizational structure to strengthen CSAT's programs and allow CSAT to

more effectively use its resources. The changes are as follows:

Section M.20, Functions is amended as follows:

Under the heading, *Office of Evaluation, Scientific Analysis, and Synthesis (MTC)*, delete the functional statement.

Under the heading, *Division of Services Improvement (MTB)*, delete the functional statement and substitute the following functional statement:

(1) Develops, plans, implements, and monitors national treatment capacity expansion and knowledge adoption program designed to improve treatment services throughout the United States, including services in other systems of care; (2) provides leadership and guidance to CSAT on the organization and financing of services for substance abuse treatment, HIPAA, and adoption of evidenced-based practices; (3) collaborates on the development of Guidelines for Applications (GFAs) and Requests for Contracts for the national treatment capacity expansion and services improvement agenda; (4) monitors grants, cooperative agreements, contracts, interagency agreements, and memoranda of understanding for treatment capacity expansion, knowledge adoption, and services improvement; (5) supports the development and testing of performance measures for public and private managed care plans and other systems of care; (6) collects, analyzes, and disseminates data and information pertaining to public and private financing and expenditures for treatment services; (7) identifies the need for, develops, and provides technical assistance to grantees, other service providers and systems of care, and others on adoption of evidence-based practices, capacity expansion, and organization and financing of services; (8) establishes and maintains collaborative relationships with other Federal, State, and local governmental agencies, national organizations, and constituency groups; (9) maintains internal expertise and collaborates with national experts on the science-to-services agenda; (10) develops funding levels for Division programs and activities; (11) provides guidance and oversight of training services for treatment professionals; and (12) provides leadership on workforce development activities.

Under the heading, *Division of State and Community Assistance (MTE)*, delete the functional statement and substitute the following functional statement:

(1) Administers the Substance Abuse Prevention and Treatment (SAPT) Block

Grant Program, including oversight and approval of Block Grant applications and maintenance of effort (MOE) issues; (2) administers the Substance Abuse Performance Partnership Grant (PPG), negotiating PPG agreements with States; (3) monitors and ensures State compliance with legislative and regulatory provisions which apply to PPG funds at State and provider levels; (4) provides guidance and technical assistance to States in preparation of State substance abuse plans; (5) conducts performance reviews of State agencies and treatment programs; (6) works closely with data and evaluation to assure proper reporting and data integrity; (7) administers the State Incentive Grant program for co-occurring disorders and the TCE grant program for co-occurring disorders; (8) works collaboratively with the Division of Services Improvement on performance measurement, GPRA, and HIPAA issues; and (9) serves as focus for State and local data infrastructure development issues.

Section M.40, Delegations of Authority: All delegations and redelegations of authority to officers and employees of SAMHSA which were in effect immediately prior to the effective date of this reorganization shall continue in them.

These organizational changes are effective August 12, 2002.

Dated: September 5, 2002.

Charles Curie,
Administrator.

[FR Doc. 02-23150 Filed 9-11-02; 8:45 am]

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DEPARTMENT OF THE INTERIOR

Fish and Wildlife Service

Notice of Availability of the Final Recovery Plan for the California Red-Legged Frog

AGENCY: Fish and Wildlife Service, Interior.

ACTION: Notice of document availability.

SUMMARY: We, the U.S. Fish and Wildlife Service (Service) announce the availability of a final recovery plan for the California red-legged frog (*Rana aurora draytonii*). The population of this subspecies of red-legged frog has been extirpated from 70 percent of its former range and is now found in coastal drainages of central California from Marin County, California, south to northern Baja California, Mexico. Actions needed for recovery include: (1) Protection of known populations and reestablishment of populations; (2)