Community Development Activities Program (RF Program).

Amount of Award: $500,000.

SUMMARY: Notice is hereby given that a noncompetitive grant award is being made to the Rural Community Assistance Program, Inc. to provide training and technical assistance to small communities struggling to deal with the safety and security of small and very small community water and wastewater treatment systems. This award addresses Congressional concern that many small and very small community water and wastewater treatment systems might be most vulnerable to terrorist attack, yet the least prepared to deal with the issues.

The application is not within the scope of any existing or expected to be issued program announcement for the Fiscal Year 2005—Rural Community Development Activities Program (RF) as authorized under the Community Services Block Grant Act of 1998, as amended; Sections 680(a)(3)(B) of the Community Opportunities Accountability, and Training and Educational Services (COATES) Act (Pub. L. 105–285). This application is expected to provide valuable on-site training and technical assistance to small and very small communities struggling to deal with the safety and security of small community water and wastewater treatment systems. This announcement is inviting application for a 12-month budget period and a 12 month project period.

The funds are not being competed due to the Senate appropriation language in FY 2005 that directs the Office of Community Services to support a Rural Community Assistance Program Small Community Infrastructure Safety and Training and Technical project. Congress intends the funds to go to an organization that is capable of conducting a project that is national in scope that provides State, regional and national infrastructure safety training workshops and on-site technical assistance targeted to small and very small community water and wastewater treatment systems.

Contact for Further Information: Administration for Children and Families, Office of Community Services, 370 L'Enfant Promenade, SW., Washington, DC 20447, Veronica Terrell—(202) 401–5295, *vterrell@acf.hhs.gov.*

Dated: May 5, 2005.

Josephine B. Robinson,

Director, Office of Community Services.

[FR Doc. 05–9912 Filed 5–17–05; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2004N–0564]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Temporary Marketing Permit Applications; Correction

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; correction.

SUMMARY: The Food and Drug Administration is correcting a notice that appeared in the **Federal Register** of April 8, 2005 (70 FR 18029). The document announced a submission for the Office of Management and Budget review and request for comments on temporary marketing permit applications. The document was published with an incorrect docket number. This document corrects that error.

FOR FURTHER INFORMATION CONTACT: Joyce Strong, Office of Policy and Planning (HF–27), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–7010.

SUPPLEMENTARY INFORMATION: In FR Doc. 05–7021, appearing on page 18029 in the **Federal Register** of Friday, April 8, 2005, the following correction is made:

1. On page 18029, in the first column, in the heading of the document, ''[Docket No. 2005N–0564]'' is corrected to read ''[Docket No. 2004N–0564]''.

Dated: May 11, 2005.

Jeffrey Shuren,

Assistant Commissioner for Policy.

[FR Doc. 05–9808 Filed 5–17–05; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2004N–0498]

Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Medical Devices; Medical Device Tracking

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a collection of information entitled ''Medical Devices; Medical Device Tracking'' has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

FOR FURTHER INFORMATION CONTACT: Peggy Robbins, Office of Management Programs (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–1223.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of March 4, 2005 (70 FR 10648), the agency announced that the proposed information collection had been submitted to OMB for review and clearance under 44 U.S.C. 3507. An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. OMB has now approved the information collection and has assigned OMB control number 0910–0442. The approval expires on April 30, 2008. A copy of the supporting statement for this information collection is available on the Internet at *http://www.fda.gov/ohrms/dockets.*

Dated: May 11, 2005.

Jeffrey Shuren,

Assistant Commissioner for Policy.

[FR Doc. 05–9809 Filed 5–17–05; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2004N–0525]

Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Reports of Corrections and Removals

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a collection of information entitled ''Reports of Corrections and Removals'' has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

FOR FURTHER INFORMATION CONTACT: Peggy Robbins, Office of Management Programs (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–1223.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of March 4, 2005 (70 FR 10647), the agency announced that the proposed information collection had been submitted to OMB for review and clearance under 44 U.S.C. 3507. An

agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. OMB has now approved the information collection and has assigned OMB control number 0910–0359. The approval expires on April 30, 2008. A copy of the supporting statement for this information collection is available on the Internet at *http://www.fda.gov/ohrms/dockets*.

Dated: May 11, 2005.

Jeffrey Shuren,

Assistant Commissioner for Policy.

[FR Doc. 05–9810 Filed 5–17–05; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2003D–0263] (formerly Docket No. 03D–0263)

Guidance for Industry on Channels of Trade Policy for Commodities With Residues of Pesticide Chemicals, for Which Tolerances Have Been Revoked, Suspended, or Modified by the Environmental Protection Agency Pursuant to Dietary Risk Considerations; 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 ''Channels of Trade Policy for Commodities With Residues of Pesticide Chemicals, for Which Tolerances Have Been Revoked, Suspended, or Modified by the Environmental Protection Agency Pursuant to Dietary Risk Considerations.'' This guidance presents FDA's general policy for implementing the channels of trade provision in the Federal Food, Drug, and Cosmetic Act (the act), as amended by the Food Quality Protection Act of 1996 (the FQPA), for food containing residues of pesticide chemicals, for which tolerances have been revoked, suspended, or modified pursuant to dietary risk considerations.

DATES: You may submit written or electronic comments on agency guidances at any time.

ADDRESSES: Submit written requests for single copies of the guidance to Michael E. Kashtock, Office of Plant and Dairy Foods, Center for Food Safety and Applied Nutrition (HFS–305), Food and Drug Administration, 5100 Paint Branch Pkwy., College Park, MD 20740. Include a self-addressed adhesive label to assist that office in processing your request. Submit written comments on the guidance to the Division of Dockets Management (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 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, 301–436–2022, e-mail: *mkashtoc@cfsan.fda.gov*.

SUPPLEMENTARY INFORMATION:

I. Background

In the **Federal Register** of July 23, 2003 (68 FR 43535), FDA announced the availability of a draft guidance document entitled ''Guidance for Industry: Channels of Trade Policy for Commodities With Residues of Pesticide Chemicals, for Which Tolerances Have Been Revoked, Suspended, or Modified by the Environmental Protection Agency.'' This guidance presents FDA's general policy for implementing the channels of trade provision in the act, as amended by the FQPA. Interested persons were given until September 22, 2003, to comment on the draft guidance.

FDA received five written comments on the draft guidance document. The agency reviewed and evaluated these comments and has modified the guidance where appropriate. In particular, FDA has modified the guidance document, including its title, to make it clear that it applies solely to food commodities that contain residues of pesticide chemicals for which the applicable tolerance was revoked, suspended, or modified by the Environmental Protection Agency (EPA) pursuant to dietary risk considerations as addressed under section 408(l)(2) of the FQPA. A comment pointed out that this condition was implied in the draft guidance document, but that it should be explicit in the final guidance.

FDA is issuing this guidance as a level 1 guidance consistent with FDA's good guidance practices regulation (21 CFR 10.115). The guidance represents the agency's current thinking on its planned enforcement approach to the channels of trade provision of the act and how that provision relates to FDA-regulated products with residues of pesticide chemicals for which tolerances have been revoked, suspended, or modified by EPA pursuant to dietary risk considerations. 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 it satisfies the requirements of the applicable statutes and regulations.

II. Paperwork Reduction Act of 1995

This 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 this guidance was approved under OMB Control No. 0910–0562. The approval expires on May 31, 2008. An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number.

III. Comments

Interested persons may submit to the Division of Dockets Management (see **ADDRESSES**), written or electronic comments regarding this document. 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 guidance and received comments may be seen in the Division of Dockets Management between 9 a.m. and 4 p.m., Monday through Friday.

IV. Electronic Access

Interested persons also may access the guidance document at *http://www.cfsan.fda.gov/guidance.html.*

Dated: March 10, 2005.

Jeffrey Shuren,

Assistant Commissioner for Policy.

[FR Doc. 05–9811 Filed 5–17–05; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Proposed Collection; Comment Request; 5 A Day Customized Survey

Summary: In compliance with the requirement of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995 for opportunity for public comment on proposed data collection projects, the National Cancer Institute (NCI), National Institutes of Health (NIH) will publish periodic summaries of proposed projects to be submitted to the Office of