additional three-year period. Responses will be submitted electronically using a web-based survey instrument. Minor changes to the instrument are proposed to address compliance with recommendations made in the updated PHS clinical practice guideline issued in May of 2008, such as coverage for combination therapies, smokeless tobacco use, and states’ familiarity with and use of the 2000 PHS guideline. The minor changes are not expected to affect the overall burden estimate. To minimize burden, each respondent will only be asked to record changes that occurred since the time of the previous submission. As in previous years, each respondent will also attach a copy of the state’s Medicaid coverage plan to their completed survey, in order to assist the research team with the interpretation of responses.

### ESTIMATED ANNUALIZED BURDEN HOURS

<table>
<thead>
<tr>
<th>Respondents</th>
<th>Number of respondents</th>
<th>Number of responses per respondent</th>
<th>Average burden per response (in hours)</th>
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</table>


Maryam I. Daneshvar,
Acting Reports Clearance Officer, Centers for Disease Control and Prevention.

[FR Doc. E0–1227 Filed 1–21–09; 8:45 am]
BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

The National Institute for Occupational Safety and Health (NIOSH) of the Centers for Disease Control and Prevention (CDC) Announces an Evaluation of Downdraft Vented Nail Salon Tables (VNTs)

Authority: 29 U.S.C. Sections 651 et seq.

AGENCY: National Institute for Occupational Safety and Health (NIOSH) of the Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

ACTION: Notice.

SUMMARY: The Division of Applied Research and Technology (DART), NIOSH, is conducting an evaluation of downdraft vented salon nail tables (VNTs). This notice invites developers, manufacturers, and vendors of VNTs to submit new, unused, downdraft VNTs for evaluation of operational characteristics and effectiveness in reducing levels of a source point tracer gas at standard distances from the vent. A 6-month supply of manufacturer recommended filters is to be submitted to NIOSH at the address below, together with the VNT. Evaluation parameters for the VNTs will include, but are not limited to:

- Airflow and capture characteristics
- Noise level, ergonomic features, and filter life
- Manufacturers, vendors, and developers who wish to submit VNTs with filters for evaluation are invited to respond to this announcement. A report on each VNT submitted for evaluation, including feedback on the evaluation parameters and staff recommendations, will be sent to the submitter. Results of the evaluation will potentially be used to develop educational materials for nail technicians and may also be disseminated through reports, publications, or presentations. NIOSH does not intend to identify manufacturers in its publications but testing information referencing particular manufacturers would be releasable if requested under the Freedom of Information Act (FOIA).

DATES: Written letter of interest must be received within 90 calendar days of publication in the Federal Register. The deadline for receipt of VNT and filter submissions is June 30, 2009. Evaluations will begin subject to the dates VNT and filter submissions are received. The VNTs will be retained for up to 10 months while being evaluated, after which they will be returned.

ADDRESSES: Manufacturers, vendors, and developers who wish to submit VNTs with filters for evaluation are invited to respond to this announcement by sending a written letter of interest to NIOSH/DART, Robert A. Taft Laboratories, 4676 Columbus Parkway, Mailstop C–23, Cincinnati, Ohio 45226, Attention: Susan Reutman, e-mail address: SReutman@cdc.gov.

SUPPLEMENTARY INFORMATION: Responses shall include: A description of the VNT including the manufacturer, schedule of availability of the VNT and filters for evaluation, and a statement of the terms under which the VNT will be made available for evaluation. Shipping and handling costs (including insurance) to ship the VNTs to NIOSH and for NIOSH to return the VNTs to the submitter will be the responsibility of the submitter. NIOSH reserves the right to decide which VNT submissions will be evaluated based on compliance with the specifications described above. NIOSH also reserves the right not to proceed in this manner.

Note: As a government entity, we cannot endorse any specific product directly, indirectly, or by implication. NIOSH will not be responsible for any costs related to usage, wear and tear or accidental damage to the VNT during transport or while the VNT is at NIOSH.

Contact Person for Technical Information: Susan Reutman, Ph.D., telephone (513) 533–8286, or e-mail SReutman@cdc.gov.


James D. Seligman,
Chief Information Officer, Centers for Disease Control and Prevention.

[FR Doc. E9–1193 Filed 1–21–09; 8:45 am]
BILLING CODE 4163–19–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2008–D–0264]

Compliance Policy Guide Sec. 540.370—Fish and Fishery Products—Decomposition; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing the
availability of Compliance Policy Guide Sec. 540.370—Fish and Fishery Products—Decomposition (the CPG). The CPG provides guidance for FDA staff on decomposition in fish and fishery products.

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

ADDRESS: Submit written comments on the CPG 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.regulations.gov. Submit written requests for single copies of the CPG to the Division of Compliance Policy (HFC–230), Office of Enforcement, Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857. Send two self-addressed adhesive labels to assist that office in processing your request, or fax your request to 240–632–6861. See the SUPPLEMENTARY INFORMATION section for electronic access to the CPG.

FOR FURTHER INFORMATION CONTACT: Robert D. Samuels, Center for Food Safety and Applied Nutrition (HFS–1061), Rockville, MD 20852. Submit written 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 CPG and received comments may be seen in the Division of Dockets Management between 9 a.m. and 4 p.m., Monday through Friday.

Please note that on January 15, 2008, the FDA Division of Dockets Management Web site transitioned to the Federal Dockets Management System (FDMS). FDMS is a Government-wide, electronic docket management system. Electronic comments or submissions will be accepted by FDA only through FDMS at http://www.regulations.gov.

III. Electronic Access

Persons with access to the Internet may obtain the CPG at http://www.fda.gov/ora/compliance_ref/cpg/default.htm.

Dated: January 12, 2009.

Michael A. Chappell,
Acting Associate Commissioner for Regulatory Affairs.

BILLING CODE 4160–01–S

I. Background

In the Federal Register of July 18, 2008 (73 FR 41361), FDA announced the availability of draft CPG Sec. 540.370—Fish and Fishery Products—Decomposition and gave interested parties an opportunity to submit comments. The agency received no comments on the draft CPG but on its own initiative made a few editorial changes for clarification purposes. The CPG provides guidance for FDA staff on decomposition in fish and fishery products. The CPG also contains information that may be useful to the regulated industry and to the public.

FDA is issuing this CPG as a level 1 guidance consistent with FDA’s good guidance practices regulation (21 CFR 10.115). The CPG represents the agency’s current thinking on FDA’s direct reference enforcement criteria related to decomposition in fish and fishery products. 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 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 CPG and received comments may be seen in the Division of Dockets Management between 9 a.m. and 4 p.m., Monday through Friday.

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SUPPLEMENTARY INFORMATION:

Intended Recipients of the Award

Seventeen programs in the LEND cohort are expected to compete in FY 2009. These programs presently have active LEND grants: University of Iowa, Johns Hopkins University, University of Missouri, University of Nebraska, Dartmouth Hitchcock Medical Center, Albert Einstein College of Medicine, Children’s Hospital of Pittsburgh, University of South Dakota, University of Vermont, Virginia Commonwealth University, West Virginia University, University of Massachusetts Medical School, Ohio State University, Vanderbilt University, Children’s Research Institute, Indiana University, University of Oklahoma.

Amount of Individual Supplemental Awards: Note: These funding levels are expected to continue in FYS 2009 and 2010.

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<thead>
<tr>
<th>Grantee</th>
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</table>

Current Project Periods: 7/1/04 through 6/30/09.

Period of Supplemental Funding: 7/1/09 through 6/30/11.


CFDA Number: 93.110.

FOR FURTHER INFORMATION CONTACT:

Laura Kavanagh; Branch Chief, MCH Training Program, Division of Research, Training and Education; Maternal and Child Health Bureau; (301) 443–2254.