TABLE 1.—ESTIMATED ANNUAL REPORTING BURDEN[1]

FDA Form No.	No. of Respondents	Annual Frequency per Respondent	Total Annual Responses	Hours per Response	Total Hours
3487	12	27	324	0.81	262

[1] There are no capital costs or operating and maintenance costs associated with this collection of information.

Dated: December 31, 2003.

Jeffrey Shuren,

Assistant Commissioner for Policy.

[FR Doc. 04–347 Filed 1–7–04; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2003N–0509]

Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Experimental Study of Health Claim Disclaimers on Foods

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a collection of information entitled ''Experimental Study of Health Claim Disclaimers on Foods'' 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 November 10, 2003 (68 FR 63802), 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 0910–0531. The approval expires on June 30, 2004. A copy of the supporting statement for this information collection is available on the Internet at *http://www.fda.gov/ohrms/dockets*.

Dated: December 31, 2003.

Jeffrey Shuren,

Assistant Commissioner for Policy.

[FR Doc. 04–348 Filed 1–7–04; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2000D–1314]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Guidance for Industry on How to Use E-Mail to Submit a Notice of Intent to Slaughter for Human Food Purposes

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that the proposed collection of information listed below has been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995.

DATES: Fax written comments on the collection of information by February 9, 2004.

ADDRESSES: OMB is still experiencing significant delays in the regular mail, including first class and express mail, and messenger deliveries are not being accepted. To ensure that comments on the information collection are received, OMB recommends that written comments be faxed to the Office of Information and Regulatory Affairs, OMB, Attn: Fumie Yokota, Desk Officer for FDA, FAX: (202) 395–6974.

FOR FURTHER INFORMATION CONTACT: Denver Presley, Office of Management Programs (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, (301) 827–1472.

SUPPLEMENTARY INFORMATION: In compliance with 44 U.S.C. 3507, FDA has submitted the following proposed collection of information to OMB for review and clearance.

Guidance for Industry on How to Use E-Mail to Submit a Notice of Intent to Slaughter for Human Food Purposes—(OMB Control Number 0910–0450)—Extension

Section 512(j) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 360b(j)) gives FDA the authority to issue regulations setting out the conditions for marketing animals treated with investigational new animal drugs for food use. Under this authority, FDA's regulations at § 511.1(b)(4) (21 CFR 511.1(b)(4)), provide that sponsors must obtain authorization to slaughter these animals for food. The Center for Veterinary Medicine (CVM) may grant such authorization to a sponsor under § 511.1(b)(5). If CVM authorizes the slaughter of investigational animals for food use, CVM issues a slaughter authorization letter to new animal drug sponsors which sets the terms under which such animals treated with investigational new animal drugs may be slaughtered. The authorization letter states that sponsors must submit slaughter notices each time such animals are to be slaughtered unless CVM waives this notice in the authorization letter. Currently, slaughter notices are submitted to CVM on paper. This guidance will give sponsors the option to submit a slaughter notice electronically as an e-mail attachment. The electronic submission of slaughter notices is part of CVM's ongoing initiative to provide a method for paperless submissions. The likely respondents to this collection of information are new animal drug sponsors who have conducted clinical studies under § 511.1(b).

In the **Federal Register** of August 7, 2003 (68 FR 47076), FDA published a 60-day notice requesting public comment on the information collection provisions. No comments were received.

FDA estimates the burden for this collection of information as follows:

TABLE 1.—ESTIMATED ANNUAL REPORTING BURDEN[1]

Form No.	No. of Respondents	Annual Frequency per Respondent	Total Annual Responses	Hours per Response	Total Hours
FDA Form 3488	12	7	84	0.40	33.6

[1] There are no capital costs or operating and maintenance costs associated with this collection of information.

Submitting a slaughter notice electronically represents a new medium for submission of information currently submitted on paper. The reporting burden for compilation and submission of this information on paper is included in OMB clearance of the information collection provisions of § 511.1 (OMB control number 0910–0117). The estimates in table 1 of this document reflect the burden associated with putting the same information on FDA Form No. 3488 and resulted from discussions with sponsors about the time necessary to complete this form.

Dated: December 31, 2003.

Jeffrey Shuren,

Assistant Commissioner for Policy.

[FR Doc. 04–349 Filed 1–7–04; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF THE INTERIOR

National Park Service

Chesapeake and Ohio Canal National Historical Park Advisory Commission; Notice of Public Meeting

Notice is hereby given in accordance with the Federal Advisory Committee Act that a meeting of the Chesapeake and Ohio Canal National Historical Park Federal Advisory Commission is scheduled for Friday, January 16, 2004, at park headquarters, 1850 Dual Highway, Suite 100, Hagerstown, Maryland. The meeting will begin at 10 a.m.

The Commission was established by *Pub. L. 91–664* to meet and consult with the Secretary of the Interior on general policies and specific matters related to the administration and development of the Chesapeake and Ohio Canal National Historical Park.

The members of the Commission are as follows: Mrs. Sheila Rabb Weidenfeld, Chairman, Mr. Charles J. Weir, Mr. Barry A. Passett, Mr. Terry W. Hepburn, Ms. Elise B. Heinz, Ms. JoAnn M. Spevacek, Mrs. Mary E. Woodward, Mrs. Donna Printz, Mrs. Ferial S. Bishop, Ms. Nancy C. Long, Mrs. Jo Reynolds, Dr. James H. Gilford, Brother James Kirkpatrick.

Topics that will be presented during the meeting include:

1. Major planning initiatives, construction and development projects.
2. Park operational issues.

The meeting will be open to the public. Any member of the public may file with the Commission a written statement concerning the matters to be discussed. Persons wishing further information concerning this meeting, or who wish to submit written statements, may contact Kevin D. Brandt, Acting Superintendent, C&O Canal National Historical Park, 1850 Dual Highway, Suite 100, Hagerstown, Maryland 21740.

Minutes of the meeting will be available for public inspection six (6) weeks after the meeting at park headquarters, Hagerstown, Maryland.

Dated: December 2, 2003.

Kevin Brandt,

Acting Superintendent, C&O Canal National Historical Park.

[FR Doc. 04–133 Filed 1–7–04; 8:45 am]

BILLING CODE 4310–SV–M

INTERNATIONAL TRADE COMMISSION

[Investigations Nos. 731–TA–1063–1068 (Preliminary)]

Certain Frozen and Canned Warmwater Shrimp and Prawns From Brazil, China, Ecuador, India, Thailand, and Vietnam

AGENCY: United States International Trade Commission.

ACTION: Institution of antidumping investigations and scheduling of preliminary phase investigations.

SUMMARY: The Commission hereby gives notice of the institution of investigations and commencement of preliminary phase antidumping investigations Nos. 731–TA–1063–1068 (Preliminary) under section 733(a) of the Tariff Act of 1930 (19 U.S.C. 1673b(a)) (the Act) to determine whether there is a reasonable indication that an industry in the United States is materially injured or threatened with material injury, or the establishment of an industry in the United States is materially retarded, by reason of imports from Brazil, China, Ecuador, India, Thailand, and Vietnam of certain frozen or canned warmwater shrimp and prawns,[1] provided for in subheadings 0306.13.00 and 1605.20.10 of the Harmonized Tariff Schedule of the United States, that are alleged to be sold in the United States at less than fair value. Unless the Department of Commerce extends the time for initiation pursuant to section 732(c)(1)(B) of the Act (19 U.S.C. 1673a(c)(1)(B)), the Commission must reach a preliminary determination in antidumping investigations in 45 days, or in this case by February 17, 2004. The Commission's views are due at Commerce within five business days thereafter, or by February 24, 2004.

For further information concerning the conduct of these investigations and rules of general application, consult the Commission's Rules of Practice and Procedure, part 201, subparts A through E (19 CFR part 201), and part 207, subparts A and B (19 CFR part 207).

EFFECTIVE DATE: December 31, 2003.

FOR FURTHER INFORMATION CONTACT: Jim McClure (202–205–3191) or Elizabeth Haines (202–205–3200), Office of Investigations, U.S. International Trade Commission, 500 E Street SW., Washington, DC 20436. Hearing-impaired persons can obtain information on this matter by contacting the Commission's TDD terminal on 202–205–1810. Persons with mobility impairments who will need special assistance in gaining access to the Commission should contact the Office of the Secretary at 202–205–2000. General information concerning the Commission may also be obtained by accessing its Internet server (*http://www.usitc.gov*). The public record for these investigations may be viewed on the Commission's electronic docket (EDIS) at *http://edis.usitc.gov.*

SUPPLEMENTARY INFORMATION:

Background—These investigations are being instituted in response to petitions

[1] For purposes of these investigations, the products covered are defined as certain warmwater shrimp and prawns, whether frozen or canned, wild-caught (ocean harvested) or farm-raised (produced by aquaculture), head-on or head-off, shell-on or peeled, tail-on or tail-off, deveined or not deveined, cooked or raw, or otherwise processed in frozen or canned form. Excluded from this definition are fresh shrimp and prawns, whether shell-on or peeled; coldwater shrimp and prawns, in any state of processing; shrimp and prawns in prepared meals; breaded shrimp and prawns; and dried shrimp and prawns.