TABLE 1.—ESTIMATED ANNUAL REPORTING BURDEN[1]

21 CFR Section	No. of Respondents	Annual Frequency per Response	Total Annual Responses	Hours per Response	Total Hours
1002(d)	1	1	1	10	10

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

The reporting burden for § 100.2(d) is insignificant because enforcement notifications are seldom used by States. During the last 3 years, FDA has not received any enforcement notifications. Since the enactment of section 403A(b) of the act (21 U.S.C. 343–1(b)) as part of the Nutrition Labeling and Education Act of 1990, FDA has received only a few enforcement notifications. Although FDA believes that the burden will be insignificant, it believes these information collection provisions should be extended to provide for the potential future need of a State government to submit enforcement notifications informing FDA when it intends to take enforcement action under the act against a particular food located in the State.

Dated: August 2, 2002.

Margaret M. Dotzel,

Associate Commissioner for Policy.

[FR Doc. 02–20121 Filed 8–8–02; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 02N–0104]

Agency Information Collection Activities; Submission for OMB Review; Comment Request; Consumer Handling of Ready-to-Eat Foods

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

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

DATES: Submit written comments on the collection of information by September 9, 2002.

ADDRESSES: Submit written comments on the collection of information to the Office of Information and Regulatory Affairs, OMB, New Executive Office Bldg., 725 17th St. NW., rm. 10235, Washington, DC 20503, Attn: Stuart Shapiro, Desk Officer for FDA.

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

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.

Consumer Handling of Ready-to-Eat Foods

Section 402 of the Federal Food, Drug, and Cosmetic Act (the act) (21 U.S.C. 342) authorizes FDA to regulate foods so that they are not adulterated. FDA's research in food safety seeks to reduce the incidence of foodborne illness by improving the ability to find new ways to detect, enumerate, and control pathogens in the food supply. FDA's Center for Food Safety and Applied Nutrition awarded two grants of research funds in September 2001 to support research into consumer refrigeration practices and shelf life for ready-to-eat (RTE) foods entitled ''Consumer Storage Length Practices for Ready-to-Eat Foods'' and ''Consumer Handling of Ready-to-Eat Foods After Purchase.''

The information that will be collected concerns consumer handling of RTE food products. The research will provide data on the storage of RTE foods in unopened and opened packages in home refrigerators; consumer understanding of expiration dates; and consumer use of this information in making decisions regarding purchases, consumption, and home storage conditions. The data from these surveys will be used to refine the Department of Health and Human Services and U.S. Department of Agriculture *Listeria monocytogenes* (LM) risk assessment, issued in draft for public comment on January 19, 2001 (66 FR 5515). The values used for home storage of foods in the draft LM risk assessment were largely based on expert opinion, not statistically supportable data. Thus, the consumer storage data from these two grants will improve FDA's confidence in the predicted risks by reducing the uncertainty in consumer practices.

For the ''Consumer Storage Length Practices for Ready-to-Eat Foods,'' approximately 2,400 respondents will be selected from an already existing nationally representative Web-enabled panel. For ''Consumer Handling of Ready-to-Eat Foods After Purchase,'' a more traditional survey approach will be used and will be conducted in three parts. In part 1, approximately 400 in-person interviews will be conducted in Tennessee, Illinois, Kansas, Missouri, Florida, and New York. Participants will be selected to represent both sexes, different income groups, and education levels, and a wide range of adults from different ethnic groups. In part 2, 100 respondents from part 1 will complete food diaries of specific foods from the day the food dairy is initiated until those foods are consumed or discarded. In part 3, two mass mailings of questionnaires will be conducted one in fall-winter and the second in spring-summer for a total of 2,000 respondents.

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

TABLE 1.—ESTIMATED ANNUAL REPORTING BURDEN[1]

	No. of Respondents	Annual Frequency per Response	Total Annual Responses	Hours per Response	Total Hours
Web-enabled panel survey	2,400	1	2,400	0.25	600
Interview survey	400	1	400	0.5	200
Food diary	100	1	100	0.5	50
Mail survey	2,000	1	2,000	0.3	600

TABLE 1.—ESTIMATED ANNUAL REPORTING BURDEN[1]—Continued

	No. of Respondents	Annual Frequency per Response	Total Annual Responses	Hours per Response	Total Hours
Total					1,450

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

The number of respondents given in table 1 of this document is based on the study design in the two grant applications. The hours per response was estimated based on experience of the grantees for similar surveys and also on the number of questions to be included in each survey instrument.

Dated: August 2, 2002.

Margaret M. Dotzel,

Associate Commissioner for Policy.

[FR Doc. 02–20122 Filed 8–8–02; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Nonclinical Studies Subcommittee of the Advisory Committee for Pharmaceutical Science; Notice of Meeting

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

This notice announces a forthcoming meeting of a public advisory committee of the Food and Drug Administration (FDA). The meeting will be open to the public.

Name of Committee: Nonclinical Studies Subcommittee of the Advisory Committee for Pharmaceutical Science.

General Function of the Committee: To provide advice and recommendations to the agency on FDA's regulatory issues.

Date and Time: The meeting will be held on September 9, 2002, from 1 p.m. to 4:30 p.m. and September 10, 2002, from 8 a.m. to 12 noon.

Location: Center for Drug Evaluation and Research, Advisory Committee conference room 1066, 5630 Fishers Lane, Rockville, MD.

Contact Person: Kathleen Reedy, Center for Drug Evaluation and Research (HFD–21), Food and Drug Administration, 5600 Fishers Lane (for express delivery, 5630 Fishers Lane, rm. 1093), Rockville, MD 20857, 301–827–7001, e-mail: *reedyk@cder.fda.gov*, or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area), code 12539. Please call the Information Line for up-to-date information on this meeting.

Agenda: On September 9, 2002, the Chair of the expert working group on cardiotoxicity will report on the progress and activities of the group and the subcommittee will be allotted time to discuss the report. The subcommittee will then discuss a plan to transfer oversight of the subcommittee from the Advisory Committee for Pharmaceutical Science to the Science Advisory Board of the National Center for Toxicological Research. On September 10, 2002, the Chair of the expert working group on vascular damage will report on the progress and activities of the group and the subcommittee members will be allotted time to discuss the report.

Procedure: Interested persons may present data, information, or views, orally or in writing, on issues pending before the subcommittee. Written submissions may be made to the contact person by August 23, 2002. Oral presentations from the public will be scheduled between approximately 10 a.m. and 11 a.m. on September 10, 2002. Time allotted for each presentation may be limited. Those desiring to make formal oral presentations should notify the contact person before August 23, 2002, and submit a brief statement of the general nature of the evidence or arguments they wish to present, the names and addresses of proposed participants, and an indication of the approximate time requested to make their presentation.

Persons attending FDA's advisory committee meetings are advised that the agency is not responsible for providing access to electrical outlets.

FDA welcomes the attendance of the public at its advisory committee meetings and will make every effort to accommodate persons with physical disabilities or special needs. If you require special accommodations due to a disability, please contact Carolyn Jones, 301–827–7001, at least 7 days in advance of the meeting.

Notice of this meeting is given under the Federal Advisory Committee Act (5 U.S.C. app. 2).

Dated: August 2, 2002.

Linda Arey Skladany,

Senior Associate Commissioner for External Relations.

[FR Doc. 02–20162 Filed 8–8–02; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HOUSING AND URBAN DEVELOPMENT

[Docket No. FR–4737–N–06]

Notice of Proposed Information Collection for Public Comment: Infrastructure Study

AGENCY: Office of Policy Development and Research, HUD.

ACTION: Notice.

SUMMARY: The proposed information collection requirement concerning a project to obtain information on the infrastructure needs of tribally designated housing entities and tribal housing agencies will be submitted to the Office of Management and Budget (OMB) for review, as required by the Paperwork Reduction Act. The Department is soliciting public comments on the subject proposal.

DATES: *Comments Due Date:* October 8, 2002.

ADDRESSES: Interested persons are invited to submit comments regarding this proposal. Comments should refer to the proposal by name and/or OMB number and should be sent to: Reports Liaison Officer, Office of Policy Development and Research, Department of Housing and Urban Development, 451 7th Street, SE, 8228, Washington, DC 20410.

FOR FURTHER INFORMATION CONTACT: Ndeye Jackson, telephone number (202) 708–5537 X5737; *Ndeye J. Jackson@HUD.gov.* Copies of the proposed forms and other available documents may be obtained from Ndeye Jackson.

SUPPLEMENTARY INFORMATION: The Department will submit the proposed information collection to OMB for review, as required by the Paperwork Reduction Act of 1995 (44 U.S.C. Chapter 35, as amended).

This Notice is soliciting comments from members of the public and affected agencies concerning the proposed