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
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 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: March 22, 2002.

Margaret M. Dotzel,

Associate Commissioner for Policy.

[FR Doc. 02–7580 Filed 3–28–02; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 01N–0301]

Agency Information Collection Activities; Announcement of OMB Approval; Customer/Partner Service Surveys

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

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

FOR FURTHER INFORMATION CONTACT: JonnaLynn P. Capezzuto, Office of Information Resources Management (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–4659.

SUPPLEMENTARY INFORMATION: Inthe **Federal Register** of December 20, 2001 (66 FR 65723), 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 0910–0360. The approval expires on March 31, 2005. A copy of the supporting statement for this information collection is available on the Internet at http://www.fda.gov/ohrms/dockets.

Dated: March 22, 2002.

Margaret M. Dotzel,

Associate Commissioner for Policy.

[FR Doc. 02–7524 Filed 3–28–02; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 01N–0402]

Agency Information Collection Activities; Announcement of OMB Approval; Medical Devices; Third-Party Premarket Submission Review and Quality System Inspections Under United States/European Community Mutual Recognition Agreement

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; Third-Party Premarket Submission Review and Quality System Inspections Under United States/European Community Mutual Recognition Agreement'' has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

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 the **Federal Register** of January 14, 2002 (67 FR 1770), 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 0910–0378. The approval expires on March 31, 2005. A copy of the supporting statement for this information collection is available on the Internet at http://www.fda.gov/ohrms/dockets.

Dated: March 22, 2002.

Margaret M. Dotzel,

Associate Commissioner for Policy.

[FR Doc. 02–7526 Filed 3–28–02; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Drug Manufacturing Inspections; Public Workshops

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of public workshops.

The Food and Drug Administration (FDA) is announcing a series of workshops to discuss the application of a systems-based approach to drug manufacturing inspections. The workshops, which will be held in collaboration with the Consumer Healthcare Products Association (CHPA), are intended to provide a regulatory perspective on the systems-based approach to inspections.

Date and Time: See table 1 following the ''Location'' section of this document.

Location: See table 1 below

TABLE1

Meeting Address	Date and Local Time	FDA Contact Person
NEW JERSEY: Sheraton Meadowlands Hotel, 2 Meadowlands Plaza, East Rutherford, NJ, 201–896–0500.	Monday, June 17, 2002, from 8:30 a.m. to 4:30 p.m.	Erik N. Henrikson
PUERTO RICO: San Juan Marriott Hotel, 1309 Ashford Ave., San Juan, PR, 800–981–8546.	Monday, July 15, 2002, from 8:30 a.m. to 4:30 p.m.	Do.
CALIFORNIA: Manhattan Beach Marriott Hotel, 1400 Parkview Dr., Manhattan Beach, CA, 310–546–7511.	Monday, August 5, 2002, from 8:30 a.m. to 4:30 p.m.	Do.

Contact:

For information regarding participation by FDA: Erik N. Henrikson, Center for Drug Evaluation and Research (HFD–320), Food and Drug Administration, 7520 Standish Pl., Rockville, MD 20855, 301–827–0072, FAX 301–594–2202.

For information regarding the program or registration: Bill Bradley, Consumer Healthcare Products Association (CHPA), 1150 Connecticut Ave. NW., Washington, DC 20036, 202–429–9260, FAX 202–223–6835.

Registration: Anyone interested in the workshops can obtain registration information from Bill Bradley, CHPA (address above), or a brochure with the program and registration form is available at http://www.chpa-info.org/meetings/pdfs/2002workshops_updated_22602.pdf. This material is also available from http://www.fda.gov.cder/calendar. Space is limited. Please preregister by the Friday prior to each of these meetings to confirm your participation. If you need special accommodations due to a disability, please contact Erik N. Henrikson (address above) at least 7 days in advance.

SUPPLEMENTARY INFORMATION:

Who Should Attend? This announcement is directed toward professionals involved in the manufacture, control, and regulation of prescription or over-the-counter drugs who will benefit from these workshops, including: Process/production engineers, quality assurance/quality control and regulatory affairs professionals, auditors, repackers and relabelers, consultants, regulatory investigators and good manufacturing practice compliance officials, and reviewing chemists. Other entities or individuals may also be interested in attending.

Is There a Registration Fee for This Workshop? Yes, a registration fee of $320.00 payable to CHPA is required for this workshop. This registration fee includes workshop reference materials and lunch on each day. Government employees qualify for a discounted rate of $75.00.

How Can I Get Additional Information, Including Copies of This Document or Other Related Documents? The notice of participation form, information about the workshops, and other related documents are available from the information contacts (addresses above) or on the Internet at http://www.fda.gov.cder/calender.

Dated: March 22, 2002.

Margaret M. Dotzel,

Associate Commissioner for Policy.

[FR Doc. 02–7579 Filed 3–28–02; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

FDA Food Labeling and Allergen Declaration; Public Workshop

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of public workshop.

SUMMARY: The Food and Drug Administration (FDA), Office of Regulatory Affairs Southwest Regional Small Business Program (Small Business Program), in collaboration with FDA's Center for Food Safety and Applied Nutrition and the Mid-Continental Association of Food and Drug Officials is announcing a public workshop entitled ''FDA Food Labeling and Allergen Declaration.'' This public workshop is intended to provide information about FDA food labeling regulations, allergen declaration and other related matters to the regulated industry, particularly small businesses and startups.

Date and Time: The public workshop will be held on August 14 and 15, 2002, from 8:30 a.m. to 5 p.m.

Location: The public workshop will be held at the Center for Community Cooperation, Oak Corner Room, 2900 Live Oak St., Dallas, TX 75204.

Contact: David Arvelo or Sue Thomason, Southwest Regional Office (HFR–SW16), Food and Drug Administration, 7920 Elmbrook Dr., suite 102, Dallas, TX 75247, 214–655–8100, ext. 130 or 128, FAX 214–655–8114, or e-mail: oraswrsbr@ora.fda.gov.

Registration: Pre-registration by July 31, 2002, is encouraged. The Mid-Continental Association of Food and Drug Officials has a $25 pre-registration fee to cover the cost of breaks. To pre-register, please complete the form below and send along with a check or money order for $25 payable to the Mid-Continental Association of Food and Drug Officials, 7920 Elmbrook Dr., suite 102, Dallas, TX 75247. As an alternative, the registration form can also be obtained on the Internet at http://www.fda.gov/ora/indust_assit/Default.htm. Directions to the facility are available at the Center for Community Cooperation Web site at http://www.cccdfw.org/pages/location.html. Seats are limited, please submit the registration form as soon as possible. Space will be filled in order of receipt of registration. Those accepted into the public workshop will receive written confirmation. Registration will close after the workshop is filled. Onsite registration will be done on a space-available basis on the day of the public workshop beginning at 8 a.m. The cost of onsite registration is $35 payable to the Mid-Continental Association of Food and Drug Officials. If you need special accommodations due to a disability, please contact David Arvelo or Sue Thomason at least 7 days in advance.

The following information is requested for registration:

Name: _____

Agency: _____

Mailing address: _____

_____

City: _____ State:_____

Zip code: _____

Phone: ( ) _____

FAX: ( ) _____

E-mail: _____

SUPPLEMENTARY INFORMATION: The workshop is being held in response to the large volume of food labeling inquiries from small food manufacturers