FOR FURTHER INFORMATION CONTACT: Dr. Sharon Holcombe, Office of Citizen Services and Communications, (202) 501–2719.

ADDRESSES: Submit comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden to Stephanie Morris, General Services Administration (MVA), Room 4035, 1800 F Street, NW., Washington, DC 20405.

SUPPLEMENTARY INFORMATION:

A. Purpose

The purpose of this information collection is to inform the General Services Administration (GSA) on how to best provide service and relevance to the American public via GSA's Web site, *http://www.GSA.gov.* The information collected from an online survey, focus groups, and Web site usability testing, will be used to refine the *http://www.GSA.gov* Web site. The questions to be asked are non-invasive and do not address or probe sensitive issues. It is important for the GSA to gain information from the many diffuse groups it serves; therefore, the GSA will be questioning individuals and households, and businesses and other-for-profit groups.

B. Annual Reporting Burden

Respondents: 190.

Responses Per Respondent: 1.

Total Responses: 190.

Hours Per Response: 72.6 minutes.

Total Burden Hours: 230.

Obtaining copies of proposals: Requesters may obtain a copy of the information collection documents from the General Services Administration, Regulatory and Federal Assistance Publications Division (MVA), 1800 F Street, NW., Room 4035, Washington, DC 20405, telephone (202) 208–7312, or by faxing your request to (202) 501–4067. Please cite Market Research Collection for the Office of Citizen Services and Communication in all correspondence.

Dated: October 2, 2002.

Michael Carleton,

Chief, Information Officer.

[FR Doc. 02–30868 Filed 12–5–02; 8:45 am]

BILLING CODE 6820–CX–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 02D–0463]

Guidance for Industry; Implementation of the Federal Food, Drug, and Cosmetic Act Regarding the Use of the Term ''Catfish;'' Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing the availability of a document entitled ''Guidance for Industry; Implementation of Section 403(t) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 343(t)) Regarding the Use of the Term 'Catfish.''' Section 10806 of the Farm Security and Rural Investment Act of 2002 amends the Federal Food, Drug, and Cosmetic Act (the act) to provide that a food shall be deemed to be misbranded ''[i]f it purports to be or is represented as catfish, unless it is fish classified within the family *Ictaluridae*.'' This guidance assists importers and domestic distributors of fish previously called ''catfish'' in selecting a new common or usual name that is consistent with the act.

DATES: Submit written or electronic comments at any time.

ADDRESSES: Submit written comments to the Dockets Management Branch (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. Submit electronic comments to http://www.fda.gov/dockets/ecomments.

Submit written requests for single copies of this guidance to the Office of Seafood (HFS–400), Center for Food Safety and Applied Nutrition, Food and Drug Administration, 5100 Paint Branch Pkwy., College Park, MD 20740. Include a self-adhesive address label to assist that office in processing your request, or include a fax number to which the guidance may be sent. See the **SUPPLEMENTARY INFORMATION** section for electronic access to this guidance document.

FOR FURTHER INFORMATION CONTACT: Spring C. Randolph, Center for Food Safety and Applied Nutrition (HFS–415), Food and Drug Administration, 5100 Paint Branch Pkwy., College Park, MD 20740, 301–436–2303, FAX 301–436–2599.

SUPPLEMENTARY INFORMATION:

I. Background

On May 13, 2002, Public Law 107–171, entitled the Farm Security and Rural Investment Act of 2002 (FSRIA), became law. Section 10806 of the FSRIA amends the food misbranding provision in section 403 of the act (21 U.S.C. 343) to provide that a food shall be deemed to be misbranded ''[i]f it purports to be or is represented as catfish, unless it is fish classified within the family *Ictaluridae*.'' This amendment overrides prior guidance that lists fish other than those from the family Ictaluridae as fish bearing the acceptable name ''catfish.''

The guidance document states that, consistent with the amendment to section 403 of the act, importers, domestic distributors, and sellers of fish in interstate commerce bearing the term ''catfish,'' that are not classified within the family *Ictaluridae*, may no longer use the term ''catfish'' on labeling, in whole or as part of their common or usual name. This guidance relates to all fish that are distributed in interstate commerce, including imports.

The document discusses how to apply FDA's common or usual name ''general principles'' regulation (21 CFR 102.5) in determining a name that can be used for the fish once known as ''catfish,'' but for which that name can no longer be used.

This guidance represents the agency's current thinking on acceptable common or usual names for fish bearing the name ''catfish'' that are not from the family *Ictaluridae*. 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.

This guidance is a level 1 guidance issued consistent with FDA's good guidance practices (GGPs) regulation (§ 10.115 (21 CFR 10.115)). Consistent with GGPs, the agency is soliciting public comment, but is implementing the guidance document immediately in accordance with § 10.115(g)(2) because the agency has determined that prior public participation is not feasible or appropriate. Section 403(t) of the act is now in effect and must be implemented immediately. Thus, FDA is making the guidance effective immediately.

II. Comments

Interested persons may submit to the Dockets Management Branch (see **ADDRESSES**) written or electronic comments on the guidance at any time. Two copies of any comments are to be submitted, except that individuals may submit one copy. Comments should be identified with the docket number

found in brackets in the heading of this document. A copy of the document and received comments may be seen in the Dockets Management Branch between 9 a.m. and 4 p.m., Monday through Friday.

III. Electronic Access

Persons with access to the Internet may obtain the document at http://www.cfsan.fda.gov/~dms/guidance.html.

Dated: November 15, 2002.

Margaret M. Dotzel,

Assistant Commissioner for Policy.

[FR Doc. 02–30901 Filed 12–5–02; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Proposed Collection; Comment Request; Request for Generic Clearance to Conduct Voluntary Customer/Partner Surveys

SUMMARY: In compliance with the requirement of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995 to provide opportunity for public comment on proposed data collection projects, the National Library of Medicine (NLM), the National Institutes of Health (NIH) will publish periodic summaries of proposed projects to be submitted to the Office of Management and Budget (OMB) for review and approval.

Proposed Collection

Title: Voluntary Customer Satisfaction Surveys.

Type of Information Collection Request: Extension. OMB Control No. 0925–0476, with an expiration date of March 31, 2003.

Need and Use of Information Collection: Executive Order 12962 directed agencies that provide significant services directly to the public to survey customers to determine the kind and quality of services they want and their level of satisfaction with existing services. Additionally, since 1994, the NLM has been a ''Federal Reinvention Laboratory '' with a goal of improving its methods of delivering information to the public. An essential strategy in accomplishing reinvention goals is the ability to periodically receive input and feedback from customers about the design and quality of the services they receive.

The NLM provides significant services directly to the public including health providers, researchers, universities, other federal agencies, state and local governments, and to others through a range of mechanisms, including publications, technical assistance, and Web sites. These services are primarily focused on health and medical information dissemination activities. The purpose of this submission is to obtain OMB's generic approval to continue to conduct satisfaction surveys of NLM's customers. The NLM will use the information provided by individuals and institutions to identify strengths and weaknesses in current services and to make improvements where feasible. The ability to periodically survey NLM's customers is essential to continually update and upgrade methods of providing high quality service.

Frequency of Response: Annually or biennially.

Affected Public: Individuals or households; businesses or other for profit; state or local governments; Federal agencies; non-profit institutions; small businesses or organizations.

Type of Respondents: Organizations, medical researchers, physicians and other health care providers, librarians, students, and the general public. Annual reporting burden is as follows:

Title of survey	Type of survey	Number of respondents	Estimated response time	Burden hours
Evaluation of Clinical Studies Database	Web-based	1,000	.167	167
Visible Human Project—Image Processing Tools	Electronic Mail	1,000	.25	250
PubMed	Web-based	5,000	.0835	418
Entrez	Web-based	2,000	.0835	167
GeneMap	Web-based	2,000	.0835	167
NCBI Web Site	Web-based	2,000	.0835	167
NLM Service Desk Survey	Interactive Voice Response telephone	400	.0835	33
NLM Onsite Reading Room Use	Exit Interview	500	.167	84
NLM Electronic Mail Customer Survey	Electronic Mail	1,000	.0835	84
MEDLINEplus User Survey	Web-based	500	.0835	59
Survey of Unified Medical Language System (UMLS) Use	Mail Survey	1,000	.5	500
NLM Services Satisfaction Survey	Web-based	2,000	.0835	167
Total		18,400		2,263

There are no capital costs to report. There are no operating or maintenance costs to report.

Request for Comments

Written comments and/or suggestions from the public and affected agencies are invited on one or more of the following points: (1) Whether the proposed collection of information is necessary for the proper performance of the function of the agency, including whether the information will have practical utility; (2) The accuracy of the agency's estimate of the burden of the proposed collection of information; (3) Ways to enhance the quality, utility, and clarity of the information to be collected; and (4) Ways to minimize the burden of the collection of information on those who are to respond, including the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology.

FOR FURTHER INFORMATION CONTACT: To request additional information on the proposed collection of information contact Ronald F. Stewart, National Library of Medicine, Building 38, Room 2N13, 8600 Rockville Pike, Bethesda, MD 20894, or call 301–496–6491 (not a toll-free number). You also may e-mail your request to: *ron_stewart@mail.nlm.nih.gov.*

Comments Due Date

Comments regarding this information collection are best assured of having their full effect if received within 60 days of the date of this publication.