www.grants.gov that contains a Grants.gov tracking number. The Administration on Aging will retrieve your application form from Grants.gov.

• We may request that you provide original signatures on forms at a later date.

2. Content and Form of Application Submission

a. DUNS Number

The Office of Management and Budget requires applicants to provide a Dun and Bradstreet (D&B) Data Universal Numbering System (DUNS) number when applying for Federal grants or cooperative agreements on or after October 1, 2003. It is entered on the SF 424. It is a unique, nine-digit identification number, which provides unique identifiers of single business entities. The DUNS number is free and easy to obtain.

Organizations can receive a DUNS number at no cost by calling the dedicated toll-free DUNS Number request line at 1–866–705–5711 or by using this link: *https://www.whitehouse.gov/omb/grants/duns_num_guide.pdf.*

Applicants unable to submit their application via *http://www.grants.gov* may request permission to submit a hard copy from the AoA Grants Management Officer: Stephen Daniels, *daniels.stephen@aoa.hhs.gov*, (202) 357–3464.

If you mail or hand deliver your application, you must submit one original application and two copies, plus a completed application checklist to AoA. The application deadline for applications sent by U.S. Postal Service must be post-marked by midnight July 14, 2006 or hand-delivered by 5 p.m. Eastern Time on July 14, 2006.

Submissions using the regular U.S. Postal Service must be addressed to: Department of Health and Human Services, Administration on Aging, Grants Management Division, Washington, DC 20201, Attention: Stephen Daniels.

Submissions by courier, overnight delivery, delivered in person, etc. should be addressed to: Department of Health and Human Services, Administration on Aging, Grants Management Division, One Massachusetts Avenue, NW., Room 4604, Washington, DC 20001, Attention: Stephen Daniels.

1. Submission Dates and Times

To receive consideration, applications must be received by the deadline listed in the ''Dates'' section of this Notice.

V. Responsiveness Criteria

Each application submitted will be screened to determine whether it was received by the closing date and time.

Applications received by the closing date and time will be screened for completeness and conformity with the requirements outlined in Sections III and IV of this Notice and the Program Announcement. Only complete applications that meet these requirements will be reviewed and evaluated competitively.

VI. Application Review Information

Eligible applications in response to this announcement will be reviewed according to the following evaluation criteria:

• Purpose and Need for Assistance—Weight: 20 points.

• Approach/Method—Work Plan and Activities—Weight: 30 points.

• Outcomes/Evaluation/Dissemination—Weight: 30 points.

• Level of Effort (Organization and Management; Budget and Resources)—Weight: 20 points.

VII. Agency Contacts

Direct inquiries regarding programmatic issues to the U.S. Department of Health and Human Services, Administration on Aging, Center for Wellness and Community-Based Care, Washington, DC 20201, telephone: (202) 357–3464.

Dated: May 30, 2006.

Josefina G. Carbonell,

Assistant Secretary for Aging.

[FR Doc. E6–8623 Filed 6–1–06; 8:45 am]

BILLING CODE 4154–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2006N–0187]

Agency Information Collection Activities: Proposed Collection; Comment Request; Survey of Health Care Professionals on the Food Safety and Nutrition Information That They Provide to Pregnant Women

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing an opportunity for public comment on the proposed collection of certain information by the agency. Under the Paperwork Reduction Act of 1995 (the PRA), Federal agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information and to allow 60 days for public comment in response to the notice. This notice solicits comments on a voluntary survey of health care professional on the food safety and nutrition information that they provide to pregnant women.

DATES: Submit written or electronic comments on the collection of information by August 1, 2006.

ADDRESSES: Submit electronic comments on the collection of information to *http://www.fda.gov/dockets/ecomments*. Submit written comments on the collection of information to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. All comments should be identified with the docket number found in brackets in the heading of this document.

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

SUPPLEMENTARY INFORMATION: Under the PRA (44 U.S.C. 3501–3520), Federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. ''Collection of information'' is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA (44 U.S.C. 3506(c)(2)(A)) requires Federal agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information before submitting the collection to OMB for approval. To comply with this requirement, FDA is publishing notice of the proposed collection of information set forth in this document.

With respect to the following collection of information, FDA invites comments on these topics: (1) Whether the proposed collection of information is necessary for the proper performance of FDA's functions, including whether the information will have practical utility; (2) the accuracy of FDA's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (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

respondents, including through the use of automated collection techniques, when appropriate, and other forms of information technology.

Survey of Health Care Professionals on the Food Safety and Nutrition Information that they Provide to Pregnant Women

Under section 903(b)(2) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 393(b)(2)), FDA is authorized to conduct research relating to foods and to conduct educational and public information programs relating to the safety of the nation's food supply. FDA is planning to conduct a survey of health care professionals to determine what information, advice, and recommendations they are offering to pregnant women about the following topics: (1) Methyl mercury and seafood consumption; (2) Listeriosis prevention; (3) weight control and nutrition; (4) dietary supplement usage; (5) food allergies; (6) Toxoplasmosis prevention; and (7) infant feeding practices. FDA is interested in obtaining this data since FDA has recently issued advice for pregnant women about food safety risks and diet risks such as mercury in seafood, Listeriosis, and Toxoplamosis. (''Food Safety for Moms-to-Be'', 2005 and ''What You Need to Know about Mercury in Fish and Shellfish'', 2004). Data from this survey will be used to evaluate whether health care professionals are aware of this advice and if they are educating their patients about information in the FDA advisories.

FDA will also use this survey to get a better understanding of what resources health care professionals use to stay abreast of current practices for caring for pregnant women. This will help FDA provide timely recommendations to health care professionals that will reach the largest audience.

A sample of 400 obstetrician/gynecologists, 200 nurse practitioners, 200 nurse midwives, 200 physician assistants, and 200 dietitians from the Special Supplemental Nutrition Program for Women, Infants, and Children (WIC) will be included in this survey. The sample of nurse practitioners, nurse midwives, and physician assistants will be drawn from those specializing in obstetrics. The samples will be randomly selected from lists obtained from national associations. The survey will be conducted using a mailed questionnaire. Cognitive interviews and a pretest will be conducted prior to fielding the survey.

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
1,200 - Survey	1	1,200	.167	200.4
75 - Pretest	1	75	.167	12.5
16 - Cognitive Interview	1	16	.75	12
Total	1	1,291		224.9

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

The burden estimate is based on FDA's experience with previous surveys.

Dated: May 25, 2006.

Jeffrey Shuren,

Assistant Commissioner for Policy.

[FR Doc. E6–8566 Filed 6–1–06; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2005N–0426]

Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Notice of Participation

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

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

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

SUPPLEMENTARY INFORMATION: In the **Federal Register** of March 16, 2006 (71 FR 13602), 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 0910–0191. The approval expires on May 31, 2009. A copy of the supporting statement for this information collection is available on the Internet at *http://www.fda.gov/ohrms/dockets*.

Dated: May 25, 2006.

Jeffrey Shuren,

Assistant Commissioner for Policy.

[FR Doc. E6–8567 Filed 6–1–06; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2005N–0393]

Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Investigational New Drug Regulations

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

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

FOR FURTHER INFORMATION CONTACT: Karen Nelson, Office of Management