a Federal duty to which the information is relevant.

i. To the National Archives and Records Administration (NARA) for records management purposes.

j. To appropriate agencies, entities, and persons when (1) The Agency suspects or has confirmed that the security or confidentiality of information in the system of records has been compromised; (2) the Agency has determined that as a result of the suspected or confirmed compromise there is a risk of harm to economic or property interests, identity theft or fraud, or harm to the security or integrity of this system or other systems or programs (whether maintained by GSA or another agency or entity) that rely upon the compromised information; and (3) the disclosure made to such agencies, entities, and persons is reasonably necessary to assist in connection with GSA's efforts to respond to the suspected or confirmed compromise and prevent, minimize, or remedy such harm.

POLICIES AND PRACTICES FOR STORING, RETRIEVING, ACCESSING, RETAINING, AND DISPOSING OF RECORDS IN THE SYSTEM:

STORAGE:

All records are stored electronically in client-server computer format.

RETRIEVABILITY:

Records are retrievable with indexing values or other unique identifiers such as name or Social Security Number.

SAFEGUARDS:

System records are safeguarded in accordance with the requirements of the Privacy Act. Access is limited to authorized individuals with strengthened passwords, and the database is maintained behind a firewall that meets strict GSA OCIO security requirements.

RETENTION AND DISPOSAL:

System records are retained and disposed of according to GSA records maintenance and disposition schedules and the requirements of the National Archives and Records Administration.

SYSTEM MANAGER AND ADDRESS:

Director, Financial and Payroll Services Division, OCFO, GSA (BCE), 1500 E. Bannister Road, Kansas City, Missouri 66085.

NOTIFICATION PROCEDURE:

Individuals wishing to inquire if the system contains information about them should contact the program manager at the above address.

RECORD ACCESS PROCEDURE:

Individuals wishing to access their own records may do so by sending a request to the program manager listed above.

CONTESTING RECORD PROCEDURES:

GSA rules for access to records, and for contesting the contents and appealing initial determinations are provided in 41 CFR part 105–64.

RECORD SOURCE CATEGORIES:

The source for the image data in the system originates from the individuals and vendors who submit the documents on their own behalf. In addition, documents may come from Federal Government Agencies that may include Privacy Act information.

[FR Doc. E9–19102 Filed 8–7–09; 8:45 am]

BILLING CODE 6820–34–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Submission for OMB review; Comment Request; 24-hour Dietary Recall Method Comparison and the National Cancer Institute (NCI) Validation and Observational Feeding Studies

SUMMARY: Under the provisions of Section 3507(a)(1)(D) of the Paperwork Reduction Act of 1995, the National Cancer Institute (NCI), the National Institutes of Health (NIH), has submitted to the Office of Management and Budget (OMB) a request to review and approve the information collection listed below. This proposed information collection was previously published in the **Federal Register** on June 3, 2009 (74 FR 26702) and allowed 60-days for public comment. One public comment was received on June 5 requesting a copy of the data collection plans. The plans were sent to the responder on June 10. The purpose of this notice is to allow an additional 30 days for public comment. The National Institutes of Health may not conduct or sponsor, and the respondent is not required to respond to, an information collection that has been extended, revised, or implemented on or after October 1, 1995, unless it displays a currently valid OMB control number.

Proposed Collection: Title: 24-hour Dietary Recall Method Comparison and the National Cancer Institute (NCI) Validation and Observational Feeding Studies. *Type of Information Collection Request:* NEW. *Need and Use of Information Collection:* The objective of the two studies is to compare the performance of the newly developed computerized Automated Self-Administered 24-Hour Recall (ASA24) approach to collecting 24-hour recall (24HR) data with the current standard, the interviewer-administered Automated Multiple Pass Method (AMPM). The ultimate goal is to determine to what extent the new automated instrument can be used instead of the more expensive interviewer-administered instrument in the collection of dietary intake data. *Frequency of Response:* Twice. *Affected Public:* Individuals. *Type of Respondents:* For the 24-hour Dietary Recall Method Comparison study, approximately 1,200 adult members from three health maintenance organization plans (in Minnesota, California, and Michigan) between ages 20 and 70 years. For the NCI Observational Feeding Study, approximately 90 adult residents from the Washington, DC metropolitan area between ages 20 and 70 years. The annual reporting burden is estimated at 1052 hours (see table below). This amounts to an estimated 2105 burden hours over the 2-year data collection period with a total cost to the respondents $37,210. There are no Capital costs, Operating costs, and/or Maintenance Costs to report.

Study Questionnaire	Number of respondents	Frequency of response	Average time response (Minutes)	Annual hour burden
24HR recall comparison study:				
Information and Consent	650	1	15/60	162.50
Screener	600	1	3/60	30.00
Dietary Recall 1	540	1	30/60	270.00
Dietary Recall 2	486	1	30/60	243.00
Demographics questionnaire	540	1	8/60	72.00

Study Questionnaire	Number of respondents	Frequency of response	Average time response (Minutes)	Annual hour burden
Preference survey	243	1	3/60	12.15
Subtotal				789.65
NCI validation and observational feeding study:				
Screener	100	1	3/60	5.00
Reminder Telephone Call	90	1	3/60	4.50
Eating 3 meals	90	1	135/60	202.50
Dietary Recall	80	1	30/60	40.00
Demographics questionnaire	80	1	8/60	10.67
Subtotal				262.67
Total				1,052.32

Request for Comments: Written comments and/or suggestions from the public and affected agencies should address one or more of the following points: (1) Whether the proposed collection of information is necessary for the proposed performance of the functions of the agency, including whether the information may have practical utility; (2) The accuracy of the 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 those who are to respond, including the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology.

Direct Comments to OMB: Written comments and/or suggestions regarding the item(s) contained in this notice, especially regarding the estimated public burden and associated response time, should be directed to the Attention: NIH Desk Officer, Office of Management and Budget, at *OIRA_submission@omb.eop.gov* or by fax to 202–395–6974. To request more information on the proposed project or to obtain a copy of the data collection plans, contact Frances E. Thompson, PhD, Project Officer, National Cancer Institute, NIH, EPN 4095A, 6130 Executive Boulevard MSC 7335, Bethesda, Maryland 20892–7335, or call non-toll-free number 301–594–4410, or FAX your request to 301–435–3710, or e-mail your request, including your address, to *thompsof@mail.nih.gov*.

Comments Due Date: Comments regarding this information collection are best assured of having their full effect if received within 30 days of the date of this publication.

Dated: July 31, 2009.

Vivian Horovitch-Kelley,

NCI Project Clearance Liaison, National Institutes of Health.

[FR Doc. E9–19022 Filed 8–7–09; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2009–N–0366]

Office of Critical Path Programs—Critical Path Initiative

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing the availability of grant funds for the support of Office of Critical Path Programs (OCPP). The goal of OCPP is to develop an administrative and scientific infrastructure to support the creation and execution of a series of projects under the FDA's Critical Path Initiative.

DATES: Important dates are as follows:

1. The application due date is September 7, 2009.
2. The anticipated start date is in September 2009.
3. The opening date is August 10, 2009.
4. The expiration date is September 8, 2009.

FOR FURTHER INFORMATION AND ADDITIONAL REQUIREMENTS CONTACT:

Nancy Stanisic, Office of Critical Path Programs (HF–18), rm. 14B45, Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–1660.

Gladys M. Bohler, Grants Management Specialist, Office of Acquisitions and Grants Services (HFA–500), Food and Drug Administration, 5630 Fishers Lane, rm. 2105, Rockville, MD 20857, 301–827–7168.

For more information on this funding opportunity announcement (FOA) and to obtain detailed requirements, please refer to the full FOA located at *http://www.fda.gov/oc/initiatives/criticalpath/*.

SUPPLEMENTARY INFORMATION:

I. Funding Opportunity Description

Funding Opportunity Description Number: RFA FD–09–019
Catalog of Federal Domestic Assistance Number: 93.103

A. Background

The Critical Path Initiative, launched by FDA in 2004, has the objective of helping modernize the development, evaluation, manufacture, and use of FDA-regulated products. Through nationwide collaboration with other Federal, academic, scientific, and industry organizations, the initiative seeks to develop new tools to facilitate innovation in FDA-regulated product development. Examples of tools include novel biomarkers, laboratory assays, genetic tests, and state-of-the art information technologies, etc. In this initiative, FDA plays the role of a facilitator in the creation of partnerships and collaborations to support specific scientific projects.

B. Research Objectives

FDA's Office of the Commissioner is announcing its intent to accept and consider a single source application for the award of a Cooperative Agreement to the Critical Path Institute (C-Path).

FDA anticipates providing up to $1.5 million (direct and indirect costs combined) during fiscal year (FY) 2009 to support research and related efforts of