ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)
State Education Agency Contacts	State Recruitment Script	17	1	30/60
School District Contacts	District Recruitment Script	80	1	30/60
School Administrators	School Recruitment Script	133	1	30/60
Teachers	Data Collection Checklist and Make-up Form	400	1	15/60
Students	NYPANS Questionnaire	8,000	1	45/60
	Height and Weight Record Form	8,000	1	3/60
	Student Contact Form	1,200	1	2/60
	24–Hour Dietary Recall Interview Script	750	3	30/60

Dated: June 15, 2009.

Maryam I. Daneshvar,

Acting Reports Clearance Officer, Centers for Disease Control and Prevention.

[FR Doc. E9–14411 Filed 6–18–09; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request

Proposed Projects

Title: Head Start Eligibility Verification.

OMB No.: New Collection.

Description: The requirements for establishing proof of eligibility for the enrollment of children in Head Start programs are documented in 45 CFR 1305.4(e). Each child's record must include a signed document by an employee identifying those documents which were reviewed to determine eligibility. Presently there is no uniform document which the employee must sign. This form will be used to facilitate an efficient and accurate determination of childrens' eligibility for Head Start enrollment.

Respondents: Head Start grantees.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Head Start Eligibility Verification	1,600	750	0.08	96,000

Estimated Total Annual Burden Hours: 96,000.

In compliance with the requirements of Section 506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Administration for Children and Families is soliciting public comment on the specific aspects of the information collection described above. Copies of the proposed collection of information can be obtained and comments may be forwarded by writing to the Administration for Children and Families, Office of Administration, Office of Information Services, 370 L'Enfant Promenade, SW., Washington, DC 20447, Attn: ACF Reports Clearance Officer. E-mail address: *infocollection@acf.hhs.gov.* All requests should be identified by the title of the information collection.

The Department specifically requests comments on: (a) Whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology. Consideration will be given to comments and suggestions submitted within 60 days of this publication.

Dated: June 16, 2009.

Janean Chambers,

Reports Clearance Officer.

[FR Doc. E9–14482 Filed 6–18–09; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2009–N–0262]

Clinical Trials Transformation Initiative (U19)

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing the availability of grant funds to support the Clinical Trials Transformation Initiative (CTTI). The goal of CTTI is to support modernization of the clinical trial enterprise by identifying practices that will enhance human subject protection, boost the quality of information derived from clinical trials, and make the research process more efficient.

DATES: Important dates are as follows:

1. The application is due by: July 6, 2009.

2. The anticipated start date is in: September 2009.

FOR FURTHER INFORMATION CONTACT:
Programmatic/Review Contact: Melissa Robb, Office of the Commissioner, Food and Drug Administration, 5600 Fishers Lane, rm. 14B–45, Rockville, MD 20857, 301–827–1516, *Melissa.robb@fda.hhs.gov*
Grants Management Contact: Gladys M. Bohler, OAGS, Food and Drug Administration, 5630 Fishers Lane, rm. 2105, Rockville, MD 20857, 301–827–7168, *gmbohler@fda.hhs.gov*.

SUPPLEMENTARY INFORMATION:

I. Funding Opportunity Description

Funding Opportunity Number: RFA–FD–09–011

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.

FDA and Duke University's Department of Translational Medicine Institute (DTMI) co-founded CTTI. CTTI's goal is to systematically modernize the clinical trial process, a goal shared by FDA's Critical Path Initiative. CTTI is made up of a broad representation of member organizations including government, industry, patient advocacy groups, professional societies, and academia. The participants are working together to identify practices that through broad adoption will increase the quality and efficiency of clinical trials.

CTTI is generating evidence about how to improve the design and execution of clinical trials. Projects about design will address principles generally applicable to clinical trials to ensure that they are fit to accomplish their intended purpose.

B. Research Objectives

The goals of this program are to develop an administrative and scientific infrastructure to support the creation and execution of a series of projects under the auspices of CTTI, to complement the goals of FDA's Critical Path Initiative.

This funding opportunity will use a cooperative agreement award mechanism (U19). In the cooperative agreement mechanism, the Project Director/Principal Investigator (PD/PI) retains the primary responsibility and dominant role for planning, directing, and executing the proposed project, with FDA staff being substantially involved as a partner with the PD/PI. Substantive involvement includes, but is not limited to, the following: (1) FDA will work closely with the DTMI throughout the lifetime of this program and throughout all phases of planning, implementation, conduct and reporting of this program and all related projects; (2) FDA will appoint project officer (s) for the task(s) associated with this program and related projects; (3) FDA will identify appropriate staff to provide strategic and scientific input, as needed, throughout the life of this program and related projects.

C. Eligibility Information

This is a sole source award to DTMI located within Duke University to support the CTTI. Only one award will be made to the DTMI to support the CTTI.

II. Award Information/Funds Available

A. Award Amount

FDA anticipates providing up to $1.5 million (direct and indirect costs combined) during fiscal year 2009 to support research and related efforts of identified projects that are part of the Critical Path Initiative.

B. Length of Support

Subject to the availability of Federal funds and successful performance of the funding opportunity announcement (FOA) stated goals and objectives, 4 additional years of support may be available depending on annual appropriations. This award will be funded based on the quality of the application received and is subject to availability of Federal funds to support the program.

III. How to Submit a Paper Application

To submit a paper application in response to this FOA, applicants should first review the full announcement located at *http://www.fda.gov/ScienceResearch/SpecialTopics/CriticalPathInitiative/SpotlightonCPIProjects/ucm083241.htm*. Persons interested in applying for a grant may obtain application forms and instructions at *http://grants.nih.gov/grants/forms.htm*. For paper submissions, the following steps are required:

- Step 1: Obtain a Dun and Bradstreet Data Universal Numbering System (DUNS) Number
- Step 2: Register with Central Contractor Registration (CCR) Instructions on how to complete these steps can be found at *http://www07.grants.gov/applicants/organization_registration.jsp*

Submit paper applications to: Gladys M. Bohler, OAGS/GAAT, Food and Drug Administration, 5630 Fishers Lane (HFA–500), rm. 2105, Rockville, MD 20874.

Dated: June 15, 2009.

Jeffrey Shuren,

Associate Commissioner for Policy and Planning.

[FR Doc. E9–14436 Filed 6–18–09; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Office of Child Support Enforcement

AGENCY: Office of Child Support Enforcement (OCSE), Administration for Children and Families (ACF), Department of Health and Human Services (HHS).

ACTION: Notice to administratively impose a matching requirement.

SUMMARY: The Office of Child Support Enforcement (OCSE) in the Administration for Children and Families (ACF) hereby gives notice to the public that a matching requirement of five percent (5%) will be administratively imposed upon awards made under competitions governed by the following ''Section 1115'' funding opportunities in Fiscal Year 2009.

CFDA Number: 93.564.

Legislative Authority

Section 1115 of the Social Security Act [42 U.S.C. 1315] provides funds for experimental, pilot or demonstration projects that are likely to assist in promoting the objectives of Part D of the Title IV. The projects must be designed to improve the financial well-being of children or otherwise improve the operation of the child support program. Projects may not permit modifications in the child support program that would have the effect of disadvantaging children in need of support.