Consistent with the statute and regulations, ACF requests extension of the ACF–800.

Respondents: States, the District of Columbia, and Territories including Puerto Rico, Guam, the Virgin Islands, American Samoa, and the Northern Marianna Islands.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
ACF–800	56	1	40	2,240

Estimated Total Annual Burden Hours: 2,240.

Additional Information

Copies of the proposed collection may be obtained by writing to the Administration for Children and Families, Office of Planning, Research and Evaluation, 370 L'Enfant Promenade SW., Washington, DC 20447, Attn: ACF Reports Clearance Officer. All requests should be identified by the title of the information collection. Email address: *infocollection@acf.hhs.gov*.

OMB Comment

OMB is required to make a decision concerning the collection of information between 30 and 60 days after publication of this document in the **Federal Register**. Therefore, a comment is best assured of having its full effect if OMB receives it within 30 days of publication. Written comments and recommendations for the proposed information collection should be sent directly to the following: Office of Management and Budget, Paperwork Reduction Project, Email: *OIRA_SUBMISSION@OMB.EOP.GOV,* Attn: Desk Officer for the Administration for Children and Families.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 2012–12302 Filed 5–21–12; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Award of a Single Source Cooperative Agreement Grant to the Congressional Hunger Center in Washington, DC

AGENCY: Office of Policy, Research and Evaluation, ACF, HHS.

ACTION: The Administration for Children and Families (ACF) announces the award of a single source cooperative agreement to the Congressional Hunger Center in Washington, DC to support a Bill Emerson National Hunger Fellow.

C.F.D.A. Number: 93.647.

Statutory Authority: The award is authorized under Section 1110 of the Social Security Act, 42 U.S.C. § 613.

SUMMARY: The Administration for Children and Families (ACF), Office of the Assistant Secretary (OAS), announces the award of a single source cooperative agreement for $3,000 with the Congressional Hunger Center (CHC) in Washington, DC, to support a Bill Emerson National Hunger Fellow who will work on hunger and obesity issues for young children. The Fellow will work closely with the ACF health team on strengthening its strategic vision to improve health and nutrition in children's programs. The Fellow will work with the team to examine programs in the Office of Child Care (OCC), OCC Tribal Maternal, Infant, and Early Childhood Home Visiting (TMIECHV) Grant Program under the Affordable Care Act (ACA), and the Office of Head Start, and will communicate with other agencies on child-focused nutrition programs. There is currently no individual in ACF designated to work specifically on these nutrition- and hunger-related issues. Additionally, the Fellow will work with the ACF health team to synthesize ideas emergent from this investigative work to further develop strategies for integrating hunger- and obesity-prevention strategies into ACF's childhood programming.

DATES: March 1, 2012—February 28, 2013.

FOR FURTHER INFORMATION CONTACT: George Askew, MD, FAAP, Senior Policy Advisor, Office of the Assistant Secretary, 901 D Street SW., Washington, DC 20447. Telephone: 202–401–1399; Email: *george.askew@acf.hhs.gov*.

George Askew,

Senior Policy Advisor, Office of the Assistant Secretary.

[FR Doc. 2012–12297 Filed 5–21–12; 8:45 am]

BILLING CODE 4184–07–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute of Child Health and Human Development; Submission for OMB Review; Comment Request; Stress and Cortisol Measurement Substudy for the National Children's Study

SUMMARY: Under the provisions of Section 3507(a)(1)(D) of the Paperwork Reduction Act of 1995, the National Institute of Child Health and Human Development (NICHD), the National Institutes of Health (NIH) has submitted to the Office of Management and Budget (OMB) a request for review and approval of the information collection listed below. This proposed information collection was previously published in the **Federal Register** on February 17, 2012, pages 9666–9668 (Volume 77, Number 33) of the **Federal Register** and allowed 60 days for public comment. One comment was received. The commenter questioned the value of the National Children's Study overall and suggested that the NCS be eliminated. The NCS is implemented to meet the requirements of the Children's Health Act of 2000 (Pub. L. 106–310). 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: Stress and Cortisol Measurement Substudy for the National Children's Study (NCS). *Type of Information Collection Request:* New. Need and Use of Information Collection: The Children's Health Act of 2000 (Pub. L. 106–310) states:

(a) PURPOSE.—It is the purpose of this section to authorize the National Institute of Child Health and Human Development* to conduct a national longitudinal study of environmental influences (including

physical, chemical, biological, and psychosocial) on children's health and development.

(b) IN GENERAL.—The Director of the National Institute of Child Health and Human Development* shall establish a consortium of representatives from appropriate Federal agencies (including the Centers for Disease Control and Prevention, the Environmental Protection Agency) to—

(1) Plan, develop, and implement a prospective cohort study, from birth to adulthood, to evaluate the effects of both chronic and intermittent exposures on child health and human development; and

(2) Investigate basic mechanisms of developmental disorders and environmental factors, both risk and protective, that influence health and developmental processes.

(c) REQUIREMENT.—The study under subsection (b) shall—

(1) Incorporate behavioral, emotional, educational, and contextual consequences to enable a complete assessment of the physical, chemical, biological, and psychosocial environmental influences on children's well-being;

(2) Gather data on environmental influences and outcomes on diverse populations of children, which may include the consideration of prenatal exposures; and

(3) Consider health disparities among children, which may include the consideration of prenatal exposures.

To fulfill the requirements of the Children's Health Act, the Stress and Cortisol Measurement Substudy will develop an optimized, item-reduced measure of self-reported stress that is supported empirically through convergent validity analysis of stress biomarkers. Specifically, key moderators of stress biomarkers will be evaluated to inform the efficiency and quality of measurements during pregnancy. Development of a scientifically robust maternal stress measure would measure chronic stress more efficiently, would not require biospecimen collection and biomarker analyses, and would thereby reduce participant burden and NCS Vanguard (Pilot) and NCS Main Study costs. With this information collection request, the NCS seeks to obtain OMB's clearance to conduct a substudy aimed at developing a validated questionnaire that will reflect specific biological and physiological measures of maternal stress.

Background: The National Children's Study is a prospective, national longitudinal study of the interaction between environment, genetics on child health and development. The Study defines ''environment'' broadly, taking a number of natural and man-made environmental, biological, genetic, and psychosocial factors into account. By studying children through their different phases of growth and development, researchers will be better able to understand the role these factors have on health and disease. Findings from the Study will be made available as the research progresses, making potential benefits known to the public as soon as possible. The National Children's Study is led by a consortium of federal partners: The U.S. Department of Health and Human Services (including the *Eunice Kennedy Shriver* National Institute of Child Health and Human Development and the National Institute of Environmental Health Sciences of the National Institutes of Health and the Centers for Disease Control and Prevention), and the U.S. Environmental Protection Agency.

To conduct the detailed preparation needed for a study of this size and complexity, the NCS was designed to include a preliminary pilot study known as the Vanguard Study. The purpose of the Vanguard Study is to assess the feasibility, acceptability, and cost of the recruitment strategy, study procedures, and outcome assessments that are to be used in the NCS Main Study. The Vanguard Study begins prior to the NCS Main Study and will run in parallel with the Main Study. At every phase of the NCS, the multiple methodological studies conducted during the Vanguard phase will inform the implementation and analysis plan for the Main Study.

In this information collection request, the NCS requests approval from OMB to perform a multi-center substudy, called the Stress and Cortisol Measurement Substudy. This substudy aims to determine the most reliable, acceptable, and cost-efficient approach for assessing maternal stress. Maternal stress is of particular interest to the NCS due to studies that have shown an association between maternal stress and negative health outcomes, including preterm birth which is one of the most important problems in maternal-child health in the U.S. Stress factors are also more prevalent in the population of socio-demographically disadvantaged women who are at an increased risk for preterm birth. Maternal stress is associated with additional health outcomes, such as still-birth, low birth weight, problems in offspring brain function and behavior (including lower IQ and impaired executive function), immune-related problems such as allergies and asthma, congenital malformations, infections, and numerous disorders of organ systems.

Development of a scientifically robust and validated questionnaire to reflect specific physiological measures of stress would allow us to measure chronic stress more efficiently, would not require biospecimen collection and biomarker analyses, and would thereby reduce participant burden and Study costs. To develop this instrument, the NCS will collect several types of information from substudy participants through medical record abstraction, questionnaires (a series of validated stress measures), physiological measures (heart rate and self-reported stress), and several types of biospecimens.

Frequency of Response: Annual [As needed].

Affected Public: Pregnant women and their children.

Type of Respondents: Pregnant women who are not geographically eligible to enroll in the NCS Vanguard Study.

Annual reporting burden: See Table 1. The annualized cost to respondents is estimated at: $73,500 (based on $10 per hour). There are no Capital Costs to report. There are no Operating or Maintenance Costs to report.

TABLE 1—ESTIMATED ANNUAL REPORTING BURDEN AND COST SUMMARY, STRESS AND CORTISOL MEASUREMENTS

Data collection activity	Type of respondent	Estimated number of respondents	Estimated number of responses per respondent	Average burden hours per response (in hours)	Estimated total annual burden hours	Estimated total annual respondent cost
Screening	Members of NCS target population (not NCS participants).	2,100	1	5/60	175	$1,750
Saliva Self-Collection Demonstration.	Members of NCS target population (not NCS participants).	700	1	15/60	175	1,750

TABLE 1—ESTIMATED ANNUAL REPORTING BURDEN AND COST SUMMARY, STRESS AND CORTISOL MEASUREMENTS—Continued

Data collection activity	Type of respondent	Estimated number of respondents	Estimated number of responses per respondent	Average burden hours per response (in hours)	Estimated total annual burden hours	Estimated total annual respondent cost
Urine Self-Collection Instructions.	Members of NCS target population (not NCS participants).	700	1	5/60	58	583
Ecological Momentary Assessment Training.	Members of NCS target population (not NCS participants).	700	1	30/60	350	3,500
Visit 1 Stress Questionnaire	Members of NCS target population (not NCS participants).	700	1	60/60	700	7,000
Adult Blood	Members of NCS target population (not NCS participants).	700	2	30/60	700	7,000
Adult Urine	Members of NCS target population (not NCS participants).	700	1	15/60	175	1,750
Adult Hair	Members of NCS target population (not NCS participants).	700	2	15/60	350	3,500
Adult Saliva	Members of NCS target population (not NCS participants).	700	28	3/60	980	9,800
Demographic and Health Interview.	Members of NCS target population (not NCS participants).	700	1	60/60	700	7,000
Participant Contact Information Sheet.	Members of NCS target population (not NCS participants).	700	1	5/60	58	583
Take-Home Questionnaire	Members of NCS target population (not NCS participants).	700	1	30/60	350	3,500
Time Diary	Members of NCS target population (not NCS participants).	700	72	2/60	1,680	16,800
Heart Monitoring	Members of NCS target population (not NCS participants).	700	1	2/60	23	233
Visit 2 Stress Questionnaire	Members of NCS target population (not NCS participants).	700	1	45/60	525	5,250
Stressful Life Events Schedule Checklist.	Members of NCS target population (not NCS participants).	700	1	30/60	350	3,500
Total		2,100			7,350	73,500

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, including the validity of the methodology and assumptions used; (3) 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 Office of Management and Budget, Office of Information and Regulatory Affairs, Attn: NIH Desk Officer, by Email to *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 and instruments, contact Ms. Jamelle E. Banks, Public Health Analyst, Office of Science Policy, Analysis and Communication, National Institute of Child Health and Human Development, 31 Center Drive Room 2A18, Bethesda, Maryland 20892, or call a non-toll free number (301) 496–1877 or Email your request, including your address to *banksj@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: May 16, 2012.

Jamelle E. Banks,

Project Clearance Liaison, Office of Science Policy, Analysis and Communications, National Institute of Child Health and Human Development.

[FR Doc. 2012–12367 Filed 5–21–12; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Cancer Institute; Closed Meetings

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. App.), notice is hereby given of the following meetings.

The meetings will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C., as amended. The grant applications/contract proposals and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant applications/contract proposals the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: National Cancer Institute Special Emphasis Panel; Tools To Probe DNA Repair and Damage Signaling Networks.

Date: June 6, 2012.

Time: 11:00 a.m. to 12:00 p.m.

Agenda: To review and evaluate contract proposals.

Place: National Institutes of Health, 6116 Executive Boulevard, Rockville, MD 20852, (Telephone Conference Call).

Contact Person: David G. Ransom, Ph.D., Scientific Review Officer, Research Programs Review Branch, Division of Extramural Activities, National Cancer Institute, NIH, 6116 Executive Blvd., Room 8133, Bethesda, MD 20892–8328, 301–451–4757, *david.ransom@nih.gov.*

This notice is being published less than 15 days prior to the meeting due to scheduling conflicts.

Name of Committee: National Cancer Institute Special Emphasis Panel; Cancer Prevention Research Small Grant Program (R03).

Date: June 28, 2012.

Time: 8:00 a.m. to 6:00 p.m.

Agenda: To review and evaluate grant applications.

Place: Bethesda North Marriott Hotel Conference & Center, 5701 Marinelli Road, North Bethesda, MD.

Contact Person: Clifford W Schweinfest, Ph.D., Scientific Review Officer, Special Review and Logistics Branch, Division of Extramural Activities, National Cancer Institute, NIH, 6116 Executive Blvd., Room 8050a, Bethesda, MD 20892–8329, 301–402–9415, *schweinfestcw@mail.nih.gov.*

Information is also available on the Institute's/Center's home page: *http://deainfo.nci.nih.gov/advisory/sep/sep.htm,* where an agenda and any additional information for the meeting will be posted when available.

(Catalogue of Federal Domestic Assistance Program Nos. 93.392, Cancer Construction; 93.393, Cancer Cause and Prevention Research; 93.394, Cancer Detection and Diagnosis Research; 93.395, Cancer Treatment Research; 93.396, Cancer Biology Research; 93.397, Cancer Centers Support; 93.398, Cancer Research Manpower; 93.399, Cancer Control, National Institutes of Health, HHS)

Dated: May 16, 2012.

Jennifer S. Spaeth,

Director, Office of Federal Advisory Committee Policy.

[FR Doc. 2012–12387 Filed 5–21–12; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Center for Scientific Review Amended; Notice of Meeting

Notice is hereby given of a change in the meeting of the Neurotransporters, Receptors, and Calcium Signaling Study Section, June 7, 2012, 8:00 a.m. to June 7, 2012, 5:00 p.m., Embassy Suites at the Chevy Chase Pavilion, 4300 Military Road NW., Washington, DC, 20015 which was published in the **Federal Register** on May 8, 2012, 77FR 27073–27075.

The meeting will be held on June 7, 2012. The meeting location has been changed to the Mayflower Renaissance 1127 Connecticut Ave. NW., Washington, DC 20036. The meeting is closed to the public.

Dated: May 16, 2012.

Jennifer S. Spaeth,

Director, Office of Federal Advisory Committee Policy.

[FR Doc. 2012–12363 Filed 5–21–12; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HOMELAND SECURITY

Office of the Secretary

[Docket No. DHS–2012–0019]

Privacy Act of 1974; U.S. Customs and Border Protection, DHS/CBP–006—Automated Targeting System, System of Records

AGENCY: Privacy Office, DHS.

ACTION: Notice of Privacy Act system of records.

SUMMARY: In accordance with the Privacy Act of 1974, the Department of Homeland Security proposes to update and expand an existing Department of Homeland Security system of records notice titled, U.S. Customs and Border Protection, DHS/CBP–006—Automated Targeting System (ATS) 72 FR 43650, August 6, 2007. The Department of Homeland Security (DHS) and U.S. Customs and Border Protection (CBP) have designed ATS to efficiently perform risk assessments on information pertaining to international travelers and import and export shipments attempting to enter or leave the United States. ATS uses a rule-managed technology that facilitates the targeting of high-risk travelers and cargo.

DHS/CBP is publishing this System of Records Notice (SORN) to update ATS and to update and expand the categories of individuals, categories of records, routine uses, access provisions, and sources of data stored in ATS. Elsewhere in the **Federal Register**, the Department of Homeland Security is concurrently issuing a Notice of Proposed Rulemaking exempting this system of records from certain provisions of the Privacy Act. This updated and expanded system will be included in the Department of Homeland Security's inventory of record systems.

DATES: Submit comments on or before June 21, 2012. This system will be effective June 21, 2012.

ADDRESSES: You may submit comments, identified by docket number DHS–2012–0019 by one of the following methods:

- *Federal e-Rulemaking Portal: http://www.regulations.gov.* Follow the instructions for submitting comments.
- *Fax:* 703–483–2999.
- *Mail:* Mary Ellen Callahan, Chief Privacy Officer, Privacy Office, Department of Homeland Security, Washington, DC 20528.

Instructions: All submissions received must include the agency name and docket number for this rulemaking. All comments received will be posted without change to *http://www.regulations.gov,* including any personal information provided.

Docket: For access to the docket to read background documents or comments received go to *http://www.regulations.gov.*

FOR FURTHER INFORMATION CONTACT: For general questions please contact: Laurence E. Castelli (202–325–0280), CBP Privacy Officer, Office of