June 30, 2011. We are requesting a three-year clearance of this collection. The 4040–0001 proposed collection encompasses 14 forms.

There are four requested changes to the SF 424 (R&R) Application for Federal Assistance (Cover) and, there are four requested changes to the R&R Other Project Information form.

These changes to the instructions will increase data quality and clarity for the collection. Agencies will not be required to collect all of the information in the proposed data set. The agency will identify the data that must be provided by applicants through instructions that will accompany the application forms.

ESTIMATED ANNUALIZED BURDEN TABLE FOR SF–424 R&R

Agency	Type of respondent	Number of annual respondents	Number of responses per respondent	Average burden on respondent per response in hours	Total burden hours
DHS	Grant Applicant	173	1	60	10,380
DOC	Grant Applicant	165	1	60	9,900
DOD	Grant Applicant	17,943	1	60	1,076,580
DOE	Grant Applicant	7,292	1	60	437,520
DOI	Grant Applicant	41	1	60	2,460
DOT	Grant Applicant	370	1	60	22,200
ED	Grant Applicant	2,000	1	60	120,000
HHS	Grant Applicant	62,133	1	60	3,727,980
NARA	Grant Applicant	1	1	60	60
NASA	Grant Applicant	102	1	60	6,120
NRC	Grant Applicant	2	1	60	120
NSF	Grant Applicant	1,001	1	60	60,060
USAID	Grant Applicant	9	1	60	540
USDA	Grant Applicant	6,349	1	60	380,940
Total		97,581			5,854,860

Mary Forbes,

Paperwork Reduction Act Reports Clearance Officer, Office of the Secretary.

[FR Doc. 2011–10250 Filed 4–27–11; 8:45 am]

BILLING CODE 4151–AE–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Statement of Organization, Functions, and Delegations of Authority; National Institutes of Health

Part N, National Institutes of Health, of the Statement of Organization, Functions, and Delegations of Authority for the Department of Health and Human Services (40 FR 22859, May 27, 1975, as amended most recently at 66 FR 6617, January 22, 2001, and redesignated from Part HN as Part N at 60 FR 56605, November 9, 1995), is amended as set forth below to establish the Division of the National Toxicology Program (NTP) within the National Institute of Environmental Health Sciences (NIEHS).

Section N–V, Organization and Functions, is amended as follows:

Immediately after the paragraph headed "Office of Translational Research" (N V4, formerly HN V4), insert the following:

Division of the National Toxicology Program (N V5, formerly HN V5). (1) Provides toxicological evaluations on substances of public health concern; (2) develops and validates improved toxicology methods (more sensitive, specific, and rapid); (3) develops approaches and generates data to strengthen the science base for risk assessments; and (4) communicates results with all stakeholders. Program goals are achieved through a highly integrated, cooperative research and testing program carried out through in-house research, research and development contracts, cooperative agreements, and other support mechanisms.

Biomolecular Screening Branch (N V52, formerly HN V52). (1) Develops research and testing activities in high and medium throughput screening assays for rapid detection of biological activities of significance to toxicology and carcinogenesis, (2) carries out the NTP automated screening assays with C. elegans, (3) develops analysis tools and approaches to allow an integrated assessment of high throughput screening endpoints and associations with findings from traditional toxicology and cancer models, and (4) develops assays and approaches to understand the genetic and epigenetic bases for differences in susceptibility.

Cellular and Molecular Pathology Branch (N V53, formerly HN V53). Responsible for (1) managing, evaluating, reviewing, and reporting all pathology data generated through conduct of NTP toxicity and carcinogenicity studies; (2) establishing standards, terminology, and diagnostic criteria for rodent pathology; (3) providing laboratory animal medicine support for the NTP and Division of Intramural Research (DIR); (4) maintaining the NTP Archives; and (5) managing pathology, toxicology, and other contracts to support NTP and DIR investigators. Staff veterinary scientists provide collaborative pathology diagnostic support for DIR investigators and mentoring/training in toxicologic pathology and laboratory animal medicine.

Program Operations Branch (N V54, formerly HN V54). (1) Provides recommendations to the NTP for scientific, administrative, and fiscal procedures and requirements by which NTP goals may be accomplished through in-house and contract activities; (2) provides resources for analytical chemistry, toxicokinetics, and evaluations of bioavailability and biotransformation; (3) initiates the contract award process and participates with the NIEHS contracts office in the review and award of the contract; (5) manages toxicity and carcinogenicity studies performed under contract and monitors them for technical and fiscal performance; (6) manages the receipt, maintenance, tracking, and dissemination of NTP documents and data.

Toxicology Branch (N V55, formerly HN V55). (1) Responsible for the design, interpretation, review, and reporting of general toxicology and carcinogenicity studies, usually in rodent models, as well as studies to evaluate targeted effects on the immune system, reproduction, development, and interference with chromosomes and

DNA for substances studied by the NTP; (2) integrates information derived from studies of absorption, metabolism, distribution, and excretion of test substances within the body and the development of mathematical models that utilize this information in the extrapolation and prediction of findings across different species and exposure conditions; (3) oversees analysis and development of models using information derived from studies of gene expression in different tissues; (4) incorporates systems biology approaches; (5) reports results from all these specialized toxicology studies; (6) develops new methodologies for toxicological assessments; and (7) provides guidance on the proper utilization of new types of toxicology information in hazard identification, hazard characterization, and regulatory decision-making.

NTP Laboratory (NTPL) (N V56, formerly, HN V56). Responsible for providing laboratory capabilities and support for the performance of agent-specific, targeted research directly related to specific substances nominated to the NTP, issues of central importance to programs of the NTP, or the development of new methods to advance the scientific programs of the NTP.

Delegations of Authority Statement: All delegations and redelegations of authority to officers and employees of NIH that were in effect immediately prior to the effective date of this reorganization and are consistent with this reorganization shall continue in effect, pending further redelegation.

Dated: April 20, 2011.

Francis S. Collins,

Director.

[FR Doc. 2011–10318 Filed 4–27–11; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30-Day–11–10GI]

Agency Forms Undergoing Paperwork Reduction Act Review

The Centers for Disease Control and Prevention (CDC) publishes a list of information collection requests under review by the Office of Management and Budget (OMB) in compliance with the Paperwork Reduction Act (44 U.S.C. chapter 35). To request a copy of these requests, call the CDC Reports Clearance Officer at (404) 639–5960 or send an e-mail to *omb@cdc.gov.* Send written comments to CDC Desk Officer, Office of Management and Budget, Washington, DC or by fax to (202) 395–5806. Written comments should be received within 30 days of this notice.

Proposed Project

Evaluating Act Against AIDS Social Marketing Campaign Phases Targeting Consumers—New—National Center for HIV/AIDS, Viral Hepatitis, STD and TB Prevention (NCHHSTP), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

In response to the continued HIV epidemic in our country, CDC has launched Act Against AIDS (AAA), a 5-year, multifaceted communication campaign to reduce HIV incidence in the United States. CDC plans to release the campaign in phases, with some of the phases running concurrently. Each phase of the campaign will use mass media and direct-to-consumer channels to deliver HIV prevention and testing messages. Some components of the campaign will be designed to provide basic education and increase awareness of HIV/AIDS among the general public, and others will be targeted to specific subgroups or communities at greatest risk of infection. The current study addresses the need to assess the effectiveness of these social marketing messages aimed at increasing HIV awareness and delivering HIV prevention and testing messages among at-risk populations.

This study will evaluate the *AAA* social marketing campaign aimed at increasing HIV/AIDS awareness, increasing prevention behaviors, and improving HIV testing rates among consumers. The study will consist of a quarterly tracking survey of *AAA* target audiences to measure exposure to each phase of the campaign and interventions implemented under *AAA.* Each extended survey will have a core set of items asked in all rounds, as well as a module of questions relating to specific *AAA* activities and communication initiatives that are occurring during a given quarter. Each extended survey sample will consist of 1,000 respondents selected from a combination of sources, including a national opt-in e-mail list sample and respondent lists generated by partnership organizations (*e.g.*, the National Urban League, the National Medical Association). Participants will self-administer the extended survey at home on personal computers. The research will include 12 data collections over a 3-year period: four self-administered quarterly extended surveys per year over 3 years, with a total of 12,000 respondents. There is no cost to the respondents other than their time. The total estimated annual burden hours are 2667.

ESTIMATED ANNUALIZED BURDEN HOURS

Respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)
Individuals (male and female) aged 18 years and older/Study Screener.	Study Screener	20,000	1	2/60
Individuals (male and female) aged 18 years and older.	Extended survey	4,000	1	30/60