enrollments, finances, faculty salaries, technology transfer activities, and institutional rankings over a 40-year period, 1970–2011. IDA also includes census information concerning neighborhoods surrounding colleges and universities.

160 Institutes of Higher Education (IHE) will be sampled from the IDA in order to collect information from key informants and key leaders from the surrounding community. Information gathered from these respondents will be

used to: (1) Develop and revise customized marketing and program materials targeting potential campus and community stakeholders; and (2) inform strategies for the marketing plan, which aims to facilitate adoption of the Safer Campuses and Communities intervention by IHEs.

The online survey will be completed by: College Administrators and staff, campus and municipal police; as well as selected community leaders. The IHEs will be contacted via email, with a maximum of 12 participants per IHE for a total of 1800 respondents. All respondent information will be maintained in a secure, electronic format accessible to a limited number of project staff. The amount of time required for a respondent to complete the survey is estimated to be 1 hour.

There are no costs to respondents other than their time. The total estimated annual burden hours are 1.800.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Form name	Number of respondents	Number of responses per respondent	Average burden per respondent (in hours)
College Administrator	CDC Questionnaire (Attachment C)	600 600 600	1 1 1	1 1 1

Dated: March 28, 2013.

Ron A. Otten.

Director, Office of Scientific Integrity, Office of the Associate Director for Science, Office of the Director, Centers for Disease Control and Prevention.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60-Day-13-13PV]

Proposed Data Collections Submitted for Public Comment and Recommendations

In compliance with the requirement of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995 for opportunity for public comment on proposed data collection projects, the Centers for Disease Control and Prevention (CDC) will publish periodic summaries of proposed projects. To request more information on the proposed projects or to obtain a copy of the data collection plans and instruments, call 404–639–7570 or send comments to Kimberly S. Lane, 1600 Clifton Road, MS–D74, Atlanta, GA 30333 or send an email to omb@cdc.gov.

Comments are invited 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) ways to enhance 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. Written comments should be received within 60 days of this notice.

Proposed Project

"Study to Explore Distribution, Reach, and Influence of Educational Children's Book *Amazing Me. It's Busy Being 3!* in Pediatric Office Settings"— NEW—National Center on Birth Defects and Developmental Disabilities (NCBDDD), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

Developmental disabilities have reached epidemic proportions in the U.S., with approximately 17 percent of children experiencing developmental delays. Impairment in physical, learning, language, or behavior areas can have a lifetime impact on everyday activities of life for a child and into adulthood. Research has shown that parents can be reliable sources of information about their children's development. Several studies have found that parents' concerns about their children's development are generally valid and predictive of developmental delays. These studies suggest that efforts to raise parental awareness of developmental milestones can increase the likelihood that children with developmental disabilities are identified

early and connected with appropriate services and support.

Using a children's picture book format, CDC developed Amazing Me: It's Busy Being 3! to increase awareness of developmental milestones among parents of 3-year-olds and actively engage them in the monitoring of their child's development. CDC partnered with Lysol and Reach Out and Read (ROR), a non-profit organization that promotes early literacy among lowincome families by distributing books in pediatric exam rooms, to disseminate copies of *Amazing Me* to parents. In spring 2012, 250 of ROR's largest pediatric clinics each received 300 copies of *Amazing Me* for distribution to parents of 3-year-old children during well-child visits. Distribution of Amazing Me through ROR practices was used as a vehicle to reach those at higher risk for developmental delays and disabilities: children insured by Medicaid and children from families with low incomes.

Preliminary data gathered from a web survey of ROR clinical staff indicates that clinical staff are not only receptive to but supportive of the *Amazing Me* book. However, the web survey of ROR clinical staff does not provide information from the book's target audience—parents. If CDC wishes to expand book distribution beyond ROR clinical settings, it will be important to gather data on parents' experiences receiving the Amazing Me book as part of a pediatric visit, and what kind of influence, if any, the book has had on their knowledge, attitudes, and beliefs about developmental milestones.

To this end, CDC will identify and recruit 3 ROR pediatric practices and 3 non-ROR practices in the greater Atlanta, Georgia and greater Washington, DC areas to distribute copies of Amazing Me to parents/ guardians of 3-year-olds, soon to be 3year-olds, or recently turned 4-year-olds attending the selected practices. The study will gather feedback from parents/ guardians about (1) their experiences receiving the book as part of a pediatric visit, and (2) the influence of the book on their awareness, attitudes, and selfefficacy regarding monitoring developmental milestones. Data will be gathered through a web survey of 900 parents/guardians who have received a copy of the Amazing Me book from

participating ROR and non-ROR practices. Parents/guardians will access the web survey by logging onto a URL address provided on a sticker affixed to the inside cover of each *Amazing Me* book. We estimate that we will screen 900 parents/guardians in order to recruit 900 respondents for the web survey.

CDC will also conduct six follow-up focus groups with survey respondents to gather more in-depth information from parents about their experiences reading the *Amazing Me* book at home with their children and assessing their child's development using the book. We estimate that we will screen 60 parents/guardians to recruit 54 participants for the focus groups. These six focus groups

will be conducted in greater Atlanta, Georgia and greater Washington, DC.

Findings from the parent web survey and focus groups will help CDC to determine if a children's book is an effective channel for reaching parents, whether more books like *Amazing Me* for other age groups should be developed, and if the ROR book distribution model is an effective means to reach low-income and at-risk families.

This request is submitted to obtain Office of Management and Budget (OMB) clearance for two years. The estimated annualized burden hours for this data collection activity are 139. There are no costs to the respondents other than their time.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total burden hours			
Web Survey								
Parents/Guardians	Web Screener and SurveyFollow-up Contact Survey	900 900	1 1	4/60 1/60	60 15			
Focus Groups								
Parents/Guardians	Screener	60 54 54	1 1 1	5/60 5/60 1	5 5 54			
Total					139			

Dated: March 28, 2013.

Ron A. Otten,

Director, Office of Scientific Integrity, Office of the Associate Director for Science, Office of the Director, Centers for Disease Control and Prevention.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day-13-0924]

Proposed Data Collections Submitted for Public Comment and Recommendations

In compliance with the requirement of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995 for opportunity for public comment on proposed data collection projects, the Centers for Disease Control and Prevention (CDC) will publish periodic summaries of proposed projects. To request more information on the

proposed projects or to obtain a copy of the data collection plans and instruments, call 404–639–5960 or send comments to Ron Otten, 1600 Clifton Road, MS–D74, Atlanta, GA 30333 or send an email to *omb@cdc.gov*.

Comments are invited 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) ways to enhance 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. Written comments should be received within 60 days of this notice.

Proposed Project

Survey of Rapid Influenza Diagnostic Test (RIDT) Practices in Clinical Laboratories and Evaluation of Laboratory Course—Reinstatement (OMB Control No. 0920–0924) with change—the Office of Surveillance, Epidemiology, and Laboratory Services (OSELS), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

The purpose of this request is to obtain Office of Budget and Management (OMB) approval to reinstate with change, the data collection for the Survey of Rapid Influenza Diagnostic Test (RIDT) Practices in Clinical Laboratories (OMB Control No. 0920-0924). OMB approval for the 2012 RIDT project expired February 28, 2012. CDC seeks a threeyear approval to conduct the RIDT project. Changes incorporated into this reinstatement request include changing the name of the collection to "Survey of Rapid Influenza Diagnostic Test (RIDT) Practices in Clinical Laboratories and Evaluation of Laboratory Course" and adding a question about whether or not the participants have taken the free CDC rapid influenza testing course, Strategies for Improving Rapid Influenza Testing