

Participation in the Work@Health Program needs assessment and pilot training evaluation surveys is voluntary for employers. There are no costs to participants other than their time.

CDC will use the information collected in the needs assessment

survey to inform the development of the Work@Health training curriculum and delivery methods. The information collected in the pilot training surveys will be used to assess respondent satisfaction with and suggestions for the

procedures, methods, content and strategies employed in each Work@Health training model.

OMB approval is requested for one year. The total estimated annualized burden hours are 117.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hr)
Employers .....	Training Needs Assessment Survey .....	200	1	20/60
Employers Participating in the Work@Health Pilot Training Program.	Pilot Employer Application Form .....	400	1	5/60
	Pilot Training: Hands-on Model Evaluation Survey.	15	1	15/60
	Pilot Training: Online Model Evaluation Survey.	15	1	15/60
	Pilot Training: Blended Model Evaluation Survey.	15	1	20/60
	Pilot Training: Train-the-Trainer Model Evaluation Survey.	15	1	15/60

**Leroy A. Richardson,**

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**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Centers for Disease Control and Prevention**

[60Day-13-13[1]]

**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 and send comments to LeRoy Richardson, at CDC 1600 Clifton Road, MS-D74, Atlanta, GA 30333 or send an email to [omb@cdc.gov](mailto: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) 2 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**

ROPS Attributes Identified by Distribution Channel Intermediaries—New—National Institute for Occupational Safety and Health (NIOSH), Centers for Disease Control and Prevention (CDC).

*Background and Brief Description*

The mission of the National Institute for Occupational Safety and Health (NIOSH) is to promote safety and health at work for all people through research and prevention. The prevention of traumatic injury is within the purview of NIOSH, and elevated incidence and rates of traumatic injury are found in the farming community. High rates of traumatic injury are associated with the use of older tractors that are not equipped with rollover protective structures (ROPS), which have been proven to reduce tractor-rollovers, a leading cause of injury to agricultural workers. To reduce the incidence of traumatic injury among farm workers, NIOSH proposes to administer stated-preference questionnaires designed to assess preference among a group of

tractor-parts dealers in Pennsylvania, New York, New Hampshire and Vermont, who have membership in the Northeast Equipment Dealers' Association (NEDA). NEDA is a trade group for tractor parts dealers and is active in 12 States in the Northeast and Mid-Atlantic States. This information will be used to assess the impediments and barriers to adoption, as well as the incentives, for the distribution and sale of ROPS.

ROPS are generally provided to end users by tractor parts dealers, who constitute distribution channel intermediaries between the manufacturer and the consumer. However, little is known about the decision processes that tractor parts dealers follow in deciding whether or not to provide ROPS to end users. The current project will generate ranking scores for the importance given to various items of concern to tractor parts dealers; these most-important items were previously developed through review of relevant research studies.

CDC proposes to collect customized information, from 520 NEDA establishments, over a one-month period. This information will be of three kinds: 1. General screening information as to the appropriateness of administering a survey to the respondent organization; 2. Limited respondent perception of the demographic characteristics on the client base served by the NEDA establishment, and 3. Importance ranking of attributes of the process of providing ROPS, or the ROPS configuration itself.

This information will allow CDC to compile a systematic, quantifiable inventory of preference data for a group that is considered representative of tractor parts dealers nationwide. It will also allow CDC to develop

recommendations for overcoming the barriers that have compromised the effectiveness of occupational health and safety programs.

The total estimated burden for the one-time retrospective data collection is

39 hours which is based on a reduced response rate of 90% (468 respondents), as indicated in the table below. The average burden per response is 5 minutes. There are no costs to respondents other than their time.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hrs)	Total burden (in hrs)
Tractor Parts Dealers .....	ROPS Questionnaire for Tractor Parts Dealers.	468	1	5/60	39
Total .....	.....	.....	.....	.....	39

**Leroy A. Richardson,**

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**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Centers for Disease Control and Prevention**

[30Day-13-13PV]

**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 (404) 639-7570 or send an email to [omb@cdc.gov](mailto:omb@cdc.gov). Send written comments to CDC Desk Officer, Office of Management and Budget, Washington, DC 20503 or by fax to (202) 395-5806. Written comments should be received within 30 days of this notice.

**Proposed Project**

Study to Explore Educational Children’s Book in Pediatric Offices—NEW—National Center on Birth Defects and Developmental Disabilities (NCBDDD), Centers for Disease Control and Prevention (CDC).

*Background and Brief Description*

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-old children 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 low-income 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 clinic staff indicates that clinic staff are not only receptive to but supportive of the *Amazing Me* book. However, the web survey of RoR clinic staff does not provide information from the book’s target audience: Parents. If CDC wishes to expand book distribution beyond ROR clinic 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 three ROR pediatric practices and three 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 3 year 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 self-efficacy regarding monitoring developmental milestones. 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.

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. All survey responses (100%) will be submitted through a secure survey Web site established for this project.

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 (2) and greater Washington, DC (4).

This request is submitted to obtain OMB clearance for one year. The estimated annualized burden is 229 hours. There are no costs to the respondents other than their time.