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

<table>
<thead>
<tr>
<th>Type of respondents</th>
<th>Form name</th>
<th>Number of respondents</th>
<th>Number of responses per respondent</th>
<th>Average burden per response (in hrs)</th>
<th>Total burden (in hrs)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Tractor Parts Dealers</td>
<td>ROPS Questionnaire for Tractor Parts Dealers</td>
<td>468</td>
<td>1</td>
<td>5/60</td>
<td>39</td>
</tr>
<tr>
<td>Total</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>39</td>
</tr>
</tbody>
</table>

Leroy A. Richardson,  
Chief, Information Collection Review Office,  
Office of Scientific Integrity, Office of the Associate Director for Science, Office of the Director, Centers for Disease Control and Prevention.

[FR Doc. 2013–17740 Filed 7–23–13; 8:45 am]  
BILLING CODE 4163–18–P

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. Send written comments to CDC Desk Office, 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.
Estimated Annualized Burden Hours

<table>
<thead>
<tr>
<th>Type of respondent</th>
<th>Form name</th>
<th>Number of respondents</th>
<th>Number of responses per respondent</th>
<th>Average burden per response (in hours)</th>
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</thead>
<tbody>
<tr>
<td>Parents/Guardians</td>
<td>Web Survey</td>
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<tr>
<td></td>
<td>Follow-up web survey</td>
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</tr>
<tr>
<td></td>
<td>Focus Group Screener</td>
<td>60</td>
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<td>5/60</td>
</tr>
<tr>
<td></td>
<td>Focus Group Informed Consent</td>
<td>54</td>
<td>1</td>
<td>5/60</td>
</tr>
<tr>
<td></td>
<td>Focus Group Moderator</td>
<td>54</td>
<td>1</td>
<td>1</td>
</tr>
</tbody>
</table>

**DATES:** Comments on the collection(s) of information must be received by the OMB desk officer by August 23, 2013.

**ADDRESSES:** When commenting on the proposed information collections, please reference the document identifier or OMB control number. To be assured consideration, comments and recommendations must be received by the OMB desk officer via one of the following transmissions: OMB, Office of Information and Regulatory Affairs, Attention: CMS Desk Officer, Fax Number: (202) 395–6974, or Email: OIRA_submission@omb.eop.gov.

To obtain copies of a supporting statement and any related forms for the proposed collection(s) summarized in this notice, you may make your request using one of the following:
2. Email your request, including your address, phone number, OMB number, and CMS document identifier, to Paperwork@hhs.gov.
3. Call the Reports Clearance Office at (410) 786–1326.

**FURTHER INFORMATION CONTACT:**
Reports Clearance Office at (410) 786–1326.

**SUPPLEMENTARY INFORMATION:**
Under the Paperwork Reduction Act of 1995 (PRA) (44 U.S.C. 3501–3520), Federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. The term “collection of information” is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA (44 U.S.C. 3506(c)(2)(A)) requires Federal agencies to publish a 30-day notice in the Federal Register concerning each proposed collection of information, including each proposed extension or reinstatement of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, CMS is publishing this notice that summarizes the following proposed collection(s) of information for public comment:

1. **Type of Information Collection Request:** Reinstatement of a currently approved collection; **Title of Information Collection:** Consolidated Renal Operations in a Web Enabled Network (CROWNWeb) Third-party Submission Authorization Form; **Use:** The Consolidated Renal Operations in a Web Enabled Network (CROWNWeb) Third-Party Submission Authorization form (CWTPSA) is to be completed by “Facility Administrators” (administrators of CMS-certified dialysis facilities) if they intend to authorize a third party (a business with which the facility is associated, or an independent vendor) to submit data to us to comply with the recently-revised Conditions for Coverage of dialysis facilities. The CROWNWeb system is the system used as the collection point of data necessary for entitlement of ESRD patients to Medicare benefits and for Federal Government monitoring and assessing of the quality and types of care provided to renal patients. The information collected through the CWTPSA form will allow us along with our contractors to receive data from authorized parties acting on behalf of CMS-certified dialysis facilities. Since February 2009, we have received 4,160 CWTPSA forms and anticipates that they will continue to receive no more than 400 new CWTPSA forms annually to address the creation of new facilities under the current participating “third party submitters.” **Form Number:** CMS–10268 (OCN: 0938–1052); **Frequency:** Occasionally; **Affected Public:** Private Sector—Business or other for-profits and Not-for-profit institutions; **Number of Respondents:** 400; **Total Annual Responses:** 400; **Total Annual Hours:** 34. (For policy questions regarding this collection contact Michelle Tucker at 410–786–0736.)

2. **Type of Information Collection Request:** Extension of a currently approved collection; **Title of Information Collection:** Medicare Quality of Care Complaint Form; **Use:** In