SEARCH Phase 3 has completed visits for 1,839 cohort study participants.

CDC plans to continue information collection for two additional years, with minor changes. Participants in the registry study will continue to complete a Medication Inventory and an Initial Participant Survey; however, the in-person study examination will be discontinued. This change will result in a decrease in burden per respondent. CDC estimates that each clinical site will identify and register an average of 255 cases per year, for a total 1,275 cases across all sites.

No data collection changes are planned for the cohort study. CDC estimates that each clinical site will conduct follow-up on an average of 142 cases per year, for a total of 710 cases across all sites. The items collected for each case include a Health Questionnaire (Youth version), an additional Health Questionnaire (Parent version), Center for Epidemiologic Study-Depression, Quality of Care, Pediatric Quality of Life Survey (Peds QL), SEARCH Michigan Neuropathy Screening Instrument, Diabetes Eating Survey, Low Blood Sugar Survey, Supplemental Survey, Tanner Stage, Retinal Photo, Family Conflict Survey, Pediatric Diabetes Quality of Life Scale, Physical Exam, Specimen Collection, and Food Frequency Questionnaire.

Findings from the registry study will be used to estimate the incidence of diabetes in youth in the U.S. Findings from the cohort study will be used to estimate the prevalence and incidence of risk factors and complications associated with diabetes in youth, including chronic microvascular complications (retinopathy, nephropathy, and autonomic neuropathy) and selected markers of macrovascular complications (hypertension, arterial stiffness) of diabetes.

Participation is voluntary and 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 hr)	Total burden (in hr)
SEARCH Registry Study Participants	Medication Inventory	1,275	1	5/60	106
	Initial Participant Survey	1,275	1	10/60	213
SEARCH Cohort Study Participants	Health Questionnaire-Youth	710	1	15/60	178
	Health Questionnaire-Parent	710	1	15/60	178
	CES-Depression	710	1	4/60	47
	Quality of Care	710	1	13/60	154
	Peds QL	710	1	5/60	59
	SEARCH MNSI Neuropathy	710	1	10/60	118
	Diabetes Eating Survey	710	1	5/60	59
	Low Blood Sugar Survey	710	1	5/60	59
	Supplemental Survey	710	1	10/60	118
	Tanner Stage	710	1	5/60	59
	Retinal Photo	710	1	15/60	178
	Family Conflict Survey	710	1	5/60	59
	Pediatric Diabetes QOL Scale	710	1	5/60	59
	Physical Exam	710	1	3	2,130
	Specimen Collection	710	1	20/60	237
	Food Frequency Questionnaire	710	1	20/60	237
Total					4,248

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. 2014–04974 Filed 3–6–14; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day–14–0138]

Proposed Data Collections Submitted for Public Comment and Recommendations

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 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

Pulmonary Function Testing Course Approval Program, 29 CFR 1910.1043 (OMB No. 0920–0138, Expiration 8/31/2014)—Revision—The National Institute for Occupational Safety and Health (NIOSH), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

NIOSH has the responsibility under the Occupational Safety and Health Administration's Cotton Dust Standard, 29 CFR 1920.1043, for approving courses to train technicians to perform pulmonary function testing in the cotton industry. Successful completion of a NIOSH-approved course is mandatory under the standard.

To carry out its responsibility, NIOSH maintains a Pulmonary Function Testing Course Approval Program. The program consists of an application submitted by potential sponsors (universities, hospitals, and private consulting firms) who seek NIOSH approval to conduct courses, and if approved, notification to NIOSH of any course or faculty changes during the approval period, which is limited to five years. The application form and added materials, including an agenda,

curriculum vitae, and course materials are reviewed by NIOSH to determine if the applicant has developed a program which adheres to the criteria required in the standard. Following approval, any subsequent changes to the course are submitted by course sponsors via letter or email and reviewed by NIOSH staff to assure that the changes in faculty or course content continue to meet course requirements.

Course sponsors also voluntarily submit an annual report to inform NIOSH of their class activity level and any faculty changes. Sponsors who elect to have their approval renewed for an additional 5 year period submit a renewal application and supporting documentation for review by NIOSH staff to ensure the course curriculum meets all current standard requirements.

Approved course sponsors that elect to offer NIOSH-Approved Spirometry Refresher Courses must submit a separate application and supporting documents for review by NIOSH staff. Institutions and organizations throughout the country voluntarily submit applications and materials to become course sponsors and carry out training. Submissions are required for NIOSH to evaluate a course and determine whether it meets the criteria in the standard and whether technicians will be adequately trained as mandated under the standard. NIOSH will disseminate a one-time customer satisfaction survey to course directors and sponsor representatives to evaluate our service to courses, the effectiveness of the program changes implemented since 2005, and the usefulness of potential Program enhancements.

The annualized figures slightly over-estimate the actual burden, due to rounding of the number of respondents for even allocation over the three-year clearance period. The estimated annual burden to respondents is 201 hours. 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 hours)
Potential Sponsors	NIOSH-Approved Spirometry Testing Course Application.	3	1	3.5
	Annual Report	35	1	30/60
	NIOSH-Example of email request for course change.	12	1	45/60
	NIOSH-Approved Spirometry Course Sponsorship Renewal Application.	13	1	6
	NIOSH-Approved Spirometry Refresher Course Application.	10	1	8
	One-Time Customer Satisfaction Survey	23	1	12/60

Leroy 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. 2014–04972 Filed 3–6–14; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier: CMS–10518]

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Centers for Medicare & Medicaid Services, HHS.

ACTION: Notice.

SUMMARY: The Centers for Medicare & Medicaid Services (CMS) is announcing an opportunity for the public to comment on CMS' intention to collect information from the public. Under the Paperwork Reduction Act of 1995 (the PRA), federal agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information (including each proposed extension or reinstatement of an existing collection of information) and to allow 60 days for public comment on the proposed action. Interested persons are invited to send comments regarding our burden estimates or any other aspect of this collection of information, including any of the following subjects: (1) The necessity and utility of the proposed information collection for the proper performance of the agency's functions; (2) the accuracy of the estimated burden; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

DATES: Comments must be received by May 6, 2014.

ADDRESSES: When commenting, please reference the document identifier or OMB control number (OCN). To be assured consideration, comments and recommendations must be submitted in any one of the following ways:

1. *Electronically.* You may send your comments electronically to *http://www.regulations.gov.* Follow the instructions for ''Comment or Submission'' or ''More Search Options'' to find the information collection document(s) that are accepting comments.

2. *By regular mail.* You may mail written comments to the following address: CMS, Office of Strategic Operations and Regulatory Affairs, Division of Regulations Development, Attention: Document Identifier/OMB Control Number _____, Room C4–26–05, 7500 Security Boulevard, Baltimore, Maryland 21244–1850.

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 following:

1. Access CMS' Web site address at *http://www.cms.hhs.gov/PaperworkReductionActof1995.*

2. Email your request, including your address, phone number, OMB number, and CMS document identifier, to *Paperwork@cms.hhs.gov.*

3. Call the Reports Clearance Office at (410) 786–1326.

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

SUPPLEMENTARY INFORMATION: