

EARLY TERMINATIONS GRANTED—Continued

[November 1, 2014 thru November 28, 2014]

20150195	G	FFL/EM Holdings, LLC; Dr. H. Douglas Barnes; FFL/EM Holdings, ITC.
11/19/2014:		
20141588	G	Thoma Bravo Fund XI, L.P.; Compuware Corporation; Thoma Bravo Fund XI, L.P.
20150146	G	BAE Systems plc; Perimeter Internetworking Corp.; BAE Systems plc.
20150157	G	Green Equity Investors VI, L.P.; HP V AIV-4, L.P.; Green Equity Investors VI, L.P.
20150170	G	HGGC Fund II, L.P.; SSI/Opinionology Newco LLC; HGGC Fund II, L.P.
20150182	G	Platform Specialty Products Corporation; Permira IV L.P. 2; Platform Specialty Products Corporation.
20150185	G	DTE Energy Co.; LS Power Equity Partners II, L.P.; DTE Energy Co.
20150188	G	Littlejohn Fund V, L.P.; Lindsay Goldberg III L.P.; Littlejohn Fund V, L.P.
20150189	G	Hercules VB Holdings, Inc.; Herff Jones, Inc.; Hercules VB Holdings, Inc.
11/20/2014:		
20150131	G	Mr. Madhava Reddy; CareTech Solutions, Inc.; Mr. Madhava Reddy.
20150179	G	Aetna Inc.; bSwift LLC; Aetna Inc.
20150192	G	Schlumberger N.V.; Don Carruth; Schlumberger N.V.
11/21/2014:		
20150144	G	Saudi Arabian Oil Company; S-Oil Corporation; Saudi Arabian Oil Company.
20150158	G	Lockheed Martin Corporation; Albert Nardslivo; Lockheed Martin Corporation.
20150161	G	DI Parent, LP; Distribution International Holdings, LLC; DI Parent, LP.
20150181	G	Mitsubishi Estate Co., Ltd.; Michael A. Ruane; Mitsubishi Estate Co., Ltd.
20150187	G	Wind Point Partners VII-A, L.P.; KIPB Group Holdings, Inc.; Wind Point Partners VII-A, L.P.
20150194	G	EQT VI (No.1) Limited Partnership; Siemens AG; EQT VI (No.1) Limited Partnership.
20150210	G	Greenbriar Equity Fund III, L.P.; Robert W. Munch and Judith A. Munch; Greenbriar Equity Fund L.P.
20150212	G	Timothy P. Horne; 2003 Riverside Capital Appreciation Fund, L.P.; Timothy P. Horne.
20150215	G	OEC Holdings 4 L.P.; Platinum Equity Capital SPL Partners, L.P.; OEC Holdings 4 L.P.
20150217	G	Scott Rudolph; Natrol Holdings, Inc. (debtor-in-possession); Scott Rudolph.
11/24/2014:		
20150199	G	General Electric Company; The Resolute Fund II SIE, L.P.; General Electric Company.
20150206	G	ArcLight Energy Partners Fund V, L.P.; Sempra Energy; ArcLight Energy Partners Fund V, L.P.
11/25/2014:		
20150013	G	CVS Holdings I, LLC; Stephen Bolick; CVS Holdings I, LLC.
20150155	G	Permira V L.P. 2; Metalogix H&S Holdings Ltd.; Permira V L.P. 2.
20150223	G	American Capital Equity III, LP; Schulman IRB Holdings LP; American Capital Equity III, LP.
11/26/2014:		
20141011	G	GlaxoSmithKline plc; Leo Constellation Limited; GlaxoSmithKline plc.
20141013	G	GlaxoSmithKline plc; Novartis AG; GlaxoSmithKline plc.

FOR FURTHER INFORMATION CONTACT:

Renee Chapman, Contact Representative
or Theresa Kingsberry, Legal Assistant,
Federal Trade Commission, Premerger
Notification Office, Bureau of
Competition, Room CC-5301,
Washington, DC 20024, (202) 326-3100.

By Direction of the Commission.

Donald S. Clark,
Secretary.

[FR Doc. 2014-28940 Filed 12-10-14; 8:45 am]

BILLING CODE 6750-01-M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day-15-15GD]

Proposed Data Collections Submitted for Public Comment and Recommendations

The Centers for Disease Control and Prevention (CDC), as part of its continuing effort to reduce public burden, invites the general public and other Federal agencies to take this opportunity to comment on proposed

and/or continuing information collections, as required by the Paperwork Reduction Act of 1995. To request more information on the below proposed project or to obtain a copy of the information collection plan and instruments, call 404-639-7570 or send comments to Leroy A. Richardson, 1600 Clifton Road, MS-D74, Atlanta, GA 30333 or send an email to omb@cdc.gov.

Comments submitted in response to this notice will be summarized and/or included in the request for Office of Management and Budget (OMB) approval. 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; (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; and (e) estimates of capital or start-up costs and costs of operation,

maintenance, and purchase of services to provide information. Burden means the total time, effort, or financial resources expended by persons to generate, maintain, retain, disclose or provide information to or for a Federal agency. This includes the time needed to review instructions; to develop, acquire, install and utilize technology and systems for the purpose of collecting, validating and verifying information, processing and maintaining information, and disclosing and providing information; to train personnel and to be able to respond to a collection of information, to search data sources, to complete and review the collection of information; and to transmit or otherwise disclose the information. Written comments should be received within 60 days of this notice.

Proposed Project

Emergency Self Escape for Coal Miners—New—National Institute for Occupational Safety and Health (NIOSH), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

The Centers for Disease Control and Prevention’s (CDC) mission is to promote health and quality of life by preventing and controlling disease, injury, and disability. The National Institute for Occupational Safety and Health (NIOSH) provides national and world leadership to prevent work-related illness, injury, disability, and death by gathering information, conducting scientific research, and translating knowledge gained into products and services. NIOSH’s mission is critical to the health and safety of every American worker. The Office of Mine Safety and Health Research (OMSHR), one of the preeminent mining research laboratories in the world, is focused on occupational health and safety research for mine workers.

Recent research by the National Academy of Sciences (NAS) has called for a detailed, formal task analysis of mine self-escape (National Research Council, 2013). Such an analysis should identify the knowledge, skills, abilities, and other attributes (KSAOs) needed by mine personnel in the event of a mine disaster to successfully complete an emergency self-escape. This analysis will identify gaps between worker demands and capabilities, and propose recommendations to either minimize those gaps or enhance existing systems (e.g., communications, training, technology).

The purpose of the project is to enhance the ability of miners to escape from underground coal mines in the event of a fire, explosion, collapse of the mine structure, or flooding of the area by toxic gas or water. To escape, miners need to perform a set of tasks that apply specific knowledge and skills in moving through the mine, avoiding dangers, and using protective equipment. The project

will identify the tasks, knowledge and skills, procedures, equipment, communications, and physical requirements of self-escape. The results are expected to lead to recommendations for improvements to task requirements and procedures, equipment, training and communication processes.

NIOSH proposes this three-year study to better understand the requirements of emergency self-escape and to answer the following questions:

- What tasks (and critical tasks) do miners perform during self-escape?
- What knowledge beyond that needed to perform normal, routine mining tasks do miners require to facilitate successful self-escape?
- What are the cognitive requirements (such as reasoning, or weighing and deciding among alternatives, recognizing when a course of action is not producing the intended results) beyond that needed to perform normal, routine mining tasks?
- What other cognitive abilities or other cognitive competencies are needed?
- What gaps exist between what miners are required to do for self-escape and their capabilities?
- How can self-escape be improved by redesigning, eliminating, or modifying tasks or training, or by altering or introducing specific technologies/tools?

To answer these questions, we will use a task analysis study design that utilizes a multiple-method approach, to include (a) review of available research, (b) interviews and focus group meetings with participants, and (c) unobtrusive observation (e.g., of drills). During interviews and focus groups, targeted questions are asked to elicit the level and type of desired information. This

system of collecting information is “active” in that participants are presented stimuli (e.g., disaster scenarios, worker roles) and asked directly to provide their perceptions (e.g., of tasks or cognitive requirements needed to accomplish self-escape in that disaster). Observation checklists have been developed to capture relevant information during the unobtrusive naturalistic observations of self-escape drills. These data are then organized, collated, and re-presented to participants for confirmation of accuracy. Recommendations are generated based on study findings, related research and practices, and logical inference.

Participants will be mining personnel drawn from two operating coal mines, one large and one smaller mine, to represent the variety within the industry. The data collection schedule (e.g., timing and duration of interviews and focus groups) will be modified as needed to minimize disruption to mine operations. No more than 30 miner volunteers will participate in the study over three years. Minimal time (< 5 minutes each) will be spent in recruitment and obtaining informed consent. Semi-structured interviews with mine personnel will require 1.5–2 hours of their time depending on the interview. Focus group sessions will require approximately 12 hours of their time total but will be executed in smaller blocks of time. Observation of drills will occur as part of normal mine operations and will not result in any additional burden on the respondents. All participants will be between the ages of 18 and 75, currently employed, and living in the United States.

There is no cost to 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
Underground coal miners	Recruitment Script	30	1	5/60	3
Underground coal miners	Informed Consent	30	1	5/60	3
Underground coal miners	Initial Interviews	6	1	90/60	9
Underground coal miners	Cognitive Task Analysis Interviews	12	2	2	48
Underground coal miners	Initial focus group sessions	12	6	2	144
Underground coal miners	Hierarchical Task Analysis focus group sessions.	12	6	2	144
Total					351

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.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day-15-15GE]

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information, processing and maintaining information, and disclosing and providing information; to train personnel and to be able to respond to a collection of information, to search data sources, to complete and review the collection of information; and to transmit or otherwise disclose the information. Written comments should be received within 60 days of this notice.

Proposed Project

Improving the Impact of Laboratory Practice Guidelines: A New Paradigm for Metrics—Clinical and Laboratory Standards Institute—NEW—Center for Surveillance, Epidemiology and Laboratory Services (CSELS), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

The Centers for Disease Control and Prevention is funding three 5-year projects collectively entitled "Improving the Impact of Laboratory Practice Guidelines: A New Paradigm for Metrics". An "LPG" is defined as written recommendations for voluntary, standardized approaches for medical laboratory testing that takes into account processes for test selection, sample procurement and processing, analytical methods, and results reporting for effective diagnosis and management of disease and health conditions. LPGs may be disseminated to, and used by, laboratorians and clinicians to assist with test selection and test result interpretation. The overall purpose of these cooperative agreements is to increase the effectiveness of LPGs by defining measures and collecting information to inform better LPG creation, revision, dissemination, promotion, uptake, and impact on clinical testing and public health. The project will explore how these processes and their impediments and facilitators differ among various intended users of LPGs. Through this demonstration project, CDC seeks to understand how to customize LPG creation and promotion to better serve these intended users of LPGs. An important goal is to help organizations that sponsor the development of LPGs create a sustainable approach for continuous quality improvement to evaluate and improve an LPG's impact through better collection of information.

The CDC selected three organizations that currently create and disseminate LPGs to support activities under a cooperative agreement funding mechanism to improve the impact of their LPGs. The American Society for Microbiology, the Clinical and Laboratory Standards Institute (CLSI),

and the College of American Pathologists, will each use their LPGs as models to better understand how to improve uptake and impact of these and future LPGs. Only the CLSI submission will be described in this notice.

Specifically, the CLSI project will address two LPGs that are important to clinical testing and have a high public health impact: *POCT12, Point-of-Care Blood Glucose Testing in Acute and Chronic Care Facilities* and *POCT13, Glucose Monitoring in Settings without Laboratory Support*. These LPGs provide guidance and recommendations for personnel monitoring patient glucose levels at sites that have access to a hospital laboratory as well as locations, such as physician offices or nursing homes that do not have an on-site moderate or high complexity laboratory. It is expected that as a result of sustained improvements in the process of creating and updating these clinical LPGs, public health, which depends upon accurate and appropriate laboratory testing guided by the use of LPGs, will also generally benefit. The intended users of the CLSI's *POCT12* and *POCT13* LPGs will include point-of-care coordinators, clinical laboratory directors, medical technologists, nurses, and medical doctors.

The CLSI plans to collect information using the same survey instrument, "Fingerstick Glucose Survey" (FGS), on three separate occasions. During the first information collection (FGS1), all targeted respondents will be asked to complete the survey. Respondents who indicate that they are not familiar with either *POCT12* or *POCT13* will be asked to provide an email address and offered a free copy of the applicable LPG. This subset of respondents will be asked to complete the same survey (FGS2) 4-6 months after receiving the free LPG. After analysis of the information collected during the first two surveys, CLSI will make improvements to *POCT12* and *POCT13*, such as provision of educational materials or helpful products such as quality control logs, and may also alter their marketing campaigns to address issues related to awareness and use of CLSI documents.

The third survey (FGS3) will then be sent to all targeted respondents approximately 2.5 years after the first survey to obtain information that can be used to evaluate the impact of these improvements. Respondents that received a free copy of *POCT12* or *POCT13* following the first survey will also be contacted by email and asked to take the third survey.

A link to the survey will be distributed to all targeted respondents either by email or postcard. The CLSI