identified above. For example, the Defense Contract Management Agency (DCMA) Contractor Purchasing System Review Group is a group dedicated to conducting CPSRs for the Department of Defense. As of June 2014 the group’s review workload included more than 700 contractors worldwide.

The cognizant ACO is responsible for granting, withholding, or withdrawing approval of a contractor’s purchasing system and for promptly notifying the contractor of same (FAR 44.305–1).

Related administrative requirements are as follows: FAR 44.305–2(c) requires that when recommendations are made for improvement of an approved system, the contractor shall be requested to reply within 15 days with a position regarding the recommendations. FAR 44.305–3(b) requires when approval of the contractor’s purchasing system is withheld or withdrawn, the ACO shall within 10 days after completing the in-plant review (1) inform the contractor in writing; (2) specify the deficiencies that must be corrected to qualify the system for approval; and (3) request the contractor to furnish within 15 days a plan for accomplishing the necessary actions. If the plan is accepted, the ACO must be corrected to qualify the system and for promptly notifying the ACO that the deficiencies have been corrected.

C. Annual Reporting Burden

1. Consent to Subcontract
   Respondents: 2,578.
   Responses per Respondent: 3.
   Total Annual Responses: 7,734.
   Hours per Response: 3.
   Total Burden Hours: 23,202.

2. Advance Notification
   Respondents: 1,861.
   Responses per Respondent: 3.
   Total Annual Responses: 5,583.
   Hours per Response: 0.25.
   Total Burden Hours: 1,396.

3. Contractors’ Purchasing System Review
   Respondents: 1,050.
   Responses per Respondent: 1.
   Total Annual Responses: 1,050.
   Hours per Response: 25.
   Total Burden Hours: 26,250.

4. Summary
   Respondents: 5,489.
   Total Annual Responses: 14,367.
   Total Burden Hours: 50,848.
   Affected Public: Businesses or other for-profit and not-for-profit institutions.

   Obtaining Copies: Requesters may obtain a copy of the information collection forms from the General Services Administration, Regulatory Secretariat Division (MVCB), 1800 F Street NW, Washington, DC 20405, telephone 202–501–4755. Please cite OMB Control No. 9000–0149, Subcontract Consent and Contractors’ Purchasing System Review, in all correspondence.

   Dated: August 20, 2018.

William Clark, Director, Office of Governmentwide Acquisition Policy, Office of Acquisition Policy, Office of Governmentwide Policy.

[FR Doc. 2018–18280 Filed 8–22–18; 8:45 am]
BILLING CODE 6820–EP–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES
Centers for Disease Control and Prevention
Mine Safety and Health Research Advisory Committee (MSHRAC), Metal Mining Automation and Advanced Technologies (MMAAT) Workgroup

AGENCY: Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

ACTION: Notice of meeting.

SUMMARY: In accordance with the Federal Advisory Committee Act, the CDC announces the following meeting for the Mine Safety and Health Research Advisory Committee (MSHRAC), Metal Mining Automation and Advanced Technologies (MMAAT) Workgroup. This meeting is open to the public, limited only by the space available. The public is welcome to submit written comments in advance of the meeting to the contact person below. Written comments received in advance of the meeting will be included in the official record of the meeting.

DATES: The meeting will be held on September 10, 2018, 8:00 a.m. to 4:00 p.m. MDT; and September 11, 2018, 8:00 a.m. to 12:00 noon MDT.

ADDRESS: University of Colorado, Anschutz Medical Campus, 13001 E 17th Place, Aurora, CO 80045. On September 10, the meeting will be held in the Krugman Conference Hall, and on September 11 in the Education 2 South Auditorium, both on that campus.

FOR FURTHER INFORMATION CONTACT: Todd Ruff, MMAAT Workgroup Designated Federal Officer, NIOSH, CDC, 315 E Montgomery Avenue, Spokane, Washington 99207, Telephone (509) 354–8003; Email ters@cdc.gov.

SUPPLEMENTARY INFORMATION: Purpose: This meeting will focus on emerging health & safety issues associated with the implementation of automation and advanced technology in the U.S. metal mining industry. The meeting is designed to identify to what extent automation and smart technologies will be implemented in metal mining and in what timeframe; to identify the related emerging health & safety concern; and to identify what gaps exist in occupational health & safety research related to automation and smart technologies.

Matters to Be Considered: The agenda will include updates on the state-of-the-art in mining automation and case studies on implementing automation at mine sites. The updates will be followed by panel discussions regarding: (1) Human factors considerations, (2) risk management, (3) automated haulage, (4) sensor technology, and (5) data analytics. Each panel will seek input and discuss the health and safety implications associated with these various topics, and identify gaps for further study. Agenda items are subject to change as priorities dictate.

The Director, Management Analysis and Services Office, has been delegated the authority to sign Federal Register notices pertaining to announcements of meetings and other committee management activities for both CDC and the Agency for Toxic Substances and Disease Registry.

Sherrill A. Berger, Chief Operating Officer, Centers for Disease Control and Prevention.

[FR Doc. 2018–18185 Filed 8–22–18; 8:45 am]
BILLING CODE 4163–19–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES
Centers for Disease Control and Prevention

Final National Occupational Research Agenda for Wholesale and Retail Trade

AGENCY: National Institute for Occupational Safety and Health (NIOSH) of the Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

ACTION: Notice of availability.

SUMMARY: NIOSH announces the availability of the final National Occupational Research Agenda for Wholesale and Retail Trade.

DATES: The final document was published August 17, 2018 on the CDC website.

ADDRESS: The document may be obtained at the following link: https://
DEPARTMENT OF HEALTH AND HUMAN SERVICES
Centers for Disease Control and Prevention
Advisory Committee on Breast Cancer in Young Women (ACBCYW);
Cancellation of Meeting

Notice is hereby given of a change in the meeting of the Advisory Committee on Breast Cancer in Young Women (ACBCYW); August 6, 2018, 1:00 p.m. to 5:00 p.m., Eastern.

The teleconference which was published in the Federal Register on June 18, 2018, Volume 83, Number 117, pages 28231–28232.

This meeting is being canceled in its entirety.

For Further Information Contact: Temeika L. Fairley, Ph.D., Designated Federal Officer, National Center for Chronic Disease Prevention and Health Promotion, CDC, 4770 Buford Hwy. NE, Mailstop K52, Atlanta, Georgia 30341, Telephone (770) 488–4518, Fax (770) 488–4760. Email: acbcyw@cdc.gov.

The Director, Management Analysis and Services Office, has been delegated the authority to sign Federal Register notices pertaining to announcements of meetings and other committee management activities, for both the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

Sherri Berger,
Chief Operating Officer, Centers for Disease Control and Prevention.

DEPARTMENT OF HEALTH AND HUMAN SERVICES
Centers for Disease Control and Prevention
[60Day–FY–2018; Docket No. CDC–2018–0063]

Proposed Data Collection Submitted for Public Comment and Recommendations

AGENCY: Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

ACTION: Notice with comment period.

SUMMARY: The Centers for Disease Control and Prevention (CDC), as part of its continuing effort to reduce public burden and maximize the utility of government information, invites the general public and other Federal agencies the opportunity to comment on a proposed and/or continuing information collection, as required by the Paperwork Reduction Act of 1995. This notice invites comment on a proposed information collection project titled “HIV prevention among Latina transgender women: Evaluation of a locally developed intervention”. The collection is part of a research study designed to evaluate the efficacy of a locally developed and culturally congruent two-session Spanish-language small-group intervention, ChiCAS (Chicas Creando Acceso a la Salud [Chicas: Girls Creating Access to Health]), which provides combination HIV prevention services to adult Hispanic/Latina transgender women at high risk for HIV infection.

DATES: CDC must receive written comments on or before October 22, 2018.

ADDRESSES: You may submit comments, identified by Docket No. CDC–2018–0063 by any of the following methods:

• Federal eRulemaking Portal: Regulations.gov. Follow the instructions for submitting comments.

• Mail: Jeffrey M. Zirger, Information Collection Review Office, Centers for Disease Control and Prevention, 1600 Clifton Road NE, MS–D74, Atlanta, Georgia 30329.

Instructions: All submissions received must include the agency name and Docket Number. CDC will post, without change, all relevant comments to Regulations.gov.

Please note: Submit all comments through the Federal eRulemaking portal (regulations.gov) or by U.S. mail to the address listed above.

FOR FURTHER INFORMATION CONTACT: To request more information on the proposed project or to obtain a copy of the information collection plan and instruments, contact Leroy A. Richardson, Information Collection Review Office, Centers for Disease Control and Prevention, 1600 Clifton Road NE, MS–D74, Atlanta, Georgia 30329; phone: 404–639–7570; Email: omb@cdc.gov.

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. In addition, the PRA also requires Federal agencies to provide a 60-day notice in the Federal Register concerning each proposed collection of information, including each new proposed collection, each proposed extension of existing collection of information, and each reinstatement of previously approved information collection before submitting the collection to the OMB for approval. To comply with this requirement, we are publishing this notice of a proposed data collection as described below.

The OMB is particularly interested in comments that will help:

1. Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility;
2. Evaluate the accuracy of the agency’s estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;
3. Enhance the quality, utility, and clarity of the information to be collected; and
4. Minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, e.g., permitting electronic submissions of responses.

5. Assess information collection costs.

Proposed Project

HIV prevention among Latina transgender women: Evaluation of a