people. The public is welcome to participate during the public comment, which is tentatively scheduled from 3:15 to 3:35 p.m. This meeting is also available by teleconference. Please dial (888) 233–0592 and enter code 33288611.

Purpose: The Subcommittee will provide advice to the CDC Director through the ACD on strategies, future needs, and challenges faced by State, Tribal, Local and Territorial health agencies, and will provide guidance on opportunities for CDC.

Matters for Discussion: The STLT subcommittee members will discuss progress on implementation of ACD-adopted recommendations related to the health department of the future, other emerging challenges and how CDC can best support STLT health departments in the transforming health system.

The agenda is subject to change as priorities dictate.

Contact Person for More Information

John Auerbach, MBA, Designated Federal Officer, STLT Subcommittee, ACD, CDC, 4770 Buford Hwy, MS E70, Atlanta, GA 30341, Telephone (404) 498–0300, Email: *OSTLTSDirector@cdc.gov*. Please submit comments to *OSTLTSDirector@cdc.gov* no later than August 4, 2016.

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.

Elaine L. Baker,

Director, Management Analysis and Services Office, Centers for Disease Control and Prevention.

[FR Doc. 2016–15932 Filed 7–5–16; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Board of Scientific Counselors, National Center for Injury Prevention and Control (BSC, NCIPC)

In accordance with Section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92–463), the Centers for Disease Control and Prevention (CDC) announces, the following meeting of the aforementioned committee:

Time and Date: 1:00 p.m.–4:00 p.m., August 1, 2016 (CLOSED).

Place: Teleconference.

Status: The meeting as designated above will be closed to the public in accordance with provisions set forth in Section 552b(c)(4) and (6), Title 5, U.S.C., and the Determination of the Director, Management Analysis and Services Office.

Purpose: The Board of Scientific Counselors makes recommendations regarding policies, strategies, objectives, and priorities; and reviews progress toward injury prevention goals and provides evidence in injury prevention-related research and programs. The Board also provides advice on the appropriate balance of intramural and extramural research, the structure, progress, and performance of intramural programs. The Board is designed to provide guidance on extramural scientific program matters, including the: (1) Review of extramural research concepts for funding opportunity announcements; (2) conduct of Secondary Peer Review of extramural research grants, cooperative agreements, and contracts applications received in response to the funding opportunity announcements as it relates to the Center's programmatic balance and mission; (3) submission of secondary review recommendations to the Center Director of applications to be considered for funding support; (4) review of research portfolios, and (5) review of program proposals. The board shall provide guidance on the National Center for Injury Prevention and Control's programs and research activities by conducting scientific peer review of intramural research and programs within the National Center for Injury Prevention and Control; by ensuring adherence to Office of Management and Budget requirements for intramural peer review; and by monitoring the overall direction, focus, and success of the National Center for Injury Prevention and Control.

Matters for Discussion: The BSC, NCIPC will meet to conduct a Secondary Peer Review of extramural research grant applications received in response to two (2) Funding Opportunity Announcements (FOAs): Evaluation of Practice-based Strategies from CDC's Rape Prevention and Education (RPE) Program to Build Evidence for Primary Prevention of Sexual Violence, FOA RFA–CE–16–005; PHS 2014–02 Omnibus Solicitation of the NIH, CDC, FDA and ACF for Small Business Innovation Research Grant Applications (Parent SBIR [R43/R44]), FOA PA–14–071. Applications will be assessed for applicability to the Center's mission and programmatic balance. Recommendations from the secondary review will be voted upon and the applications will be forwarded to the Center Director for consideration for funding support.

Agenda items are subject to change as priorities dictate.

Contact Person for More Information: Gwendolyn H. Cattledge, Ph.D., M.S.E.H., Deputy Associate Director for Science, NCIPC, CDC, 4770 Buford Highway NE., Mailstop F–63, Atlanta, GA 30341, Telephone (770) 488–1430.

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.

Elaine L. Baker,

Director, Management Analysis and Services Office, Centers for Disease Control and Prevention.

[FR Doc. 2016–15931 Filed 7–5–16; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[Docket Number CDC–2016–0020, NIOSH–289]

Issuance of Final Publication: National Institute for Occupational Safety and Health (NIOSH) Quality Assurance Review of B Readers' Classifications Submitted in the Department of Labor (DOL) Black Lung Benefits Program

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 issuance of final publication.

SUMMARY: The National Institute for Occupational Safety and Health (NIOSH) of the Centers for Disease Control and Prevention (CDC) announces the availability of the following publication: *National Institute for Occupational Safety and Health (NIOSH) Quality Assurance Review of B Readers' Classifications Submitted in the Department of Labor (DOL) Black Lung Benefits Program.*

ADDRESSES: The document may be obtained at the following link: *http://www.cdc.gov/niosh/topics/chestradiography/breader-blacklung-benefits-qa-program.html.*

FOR FURTHER INFORMATION CONTACT: Dr. Eileen Storey, NIOSH, Respiratory Health Division, Surveillance Branch, 1095 Willowdale Road, Morgantown, WV 26505. Telephone (304) 285–5754 (this is not a toll-free number).

Dated: June 30, 2016.

Frank Hearl,

Chief of Staff, National Institute for Occupational Safety and Health, Centers for Disease Control and Prevention.

[FR Doc. 2016–15978 Filed 7–5–16; 8:45 am]

BILLING CODE 4163–19–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Opportunity To Co-Sponsor Office for Human Research Protections Educational Workshops

AGENCY: Office for Human Research Protections, Office of the Assistant Secretary for Health, Office of the Secretary, Department of Health and Human Services.

ACTION: Notice.

SUMMARY: The Office for Human Research Protections (OHRP) announces the opportunity for non-federal public and private sector entities to co-sponsor OHRP Educational Workshops. Potential co-sponsors must have an approved Federal-wide Assurance with OHRP, be recipients of HHS grants for human subject research, and have a demonstrated interest and experience in the protection of human subjects in research. Potential co-sponsors must be willing to participate substantively in the co-sponsored activity.

DATES: Requests for co-sponsorships of OHRP Educational Workshops are received throughout the year at the email address below. OHRP co-sponsors a limited number of Educational Workshops with institutions each year. Requests are being received for Educational Workshops that will take place in the fall of 2016 or beyond.

ADDRESSES: Requests for co-sponsorships should be sent to *OHRP-EDU@HHS.GOV* with ''Co-sponsorship for OHRP Educational Workshops'' in the subject field or by mail to OHRP at 1101 Wootton Parkway, Suite 200, Rockville MD 20852.

FOR FURTHER INFORMATION CONTACT: *OHRP-EDU@HHS.GOV* or call OHRP's Division of Education and Development (DED) at 240–453–6900.

SUPPLEMENTARY INFORMATION:

Description

The Office for Human Research Protections (OHRP) provides leadership in the protection of the rights, welfare, and well-being of subjects involved in research conducted or supported by the U.S. Department of Health and Human Services (HHS). The OHRP is a program office within the Office of the Assistant Secretary for Health, Office of the Secretary, HHS.

OHRP provides clarification and guidance, develops educational programs and materials, maintains regulatory oversight, and provides advice on ethical and regulatory issues in biomedical and behavioral research. OHRP also supports the Secretary's Advisory Committee on Human Research Protections (SACHRP), which advises the Secretary of Health and Human Services on issues of human subject protections.

Consistent with OHRP's mission and the applicable statutory authority, 42 U.S.C. 289, OHRP Educational Workshops aim to provide clarification and guidance to the public on how to interpret, implement, and comply with the HHS-regulations on the protection of human subjects in research. Workshops are moderate size half-day or one-day events that typically accept between 120 and 140 attendees.

Co-sponsors will assist with workshop and agenda development, coordination, financial management, and meeting logistics in conjunction with OHRP staff.

Co-sponsors can charge registration fees to recover costs associated with the events; however, co-sponsors may not set registration fees at an amount higher than necessary to recover related conference expenses. Further, we expect co-sponsors to be solely responsible for collecting and handling any registration fees collected.

Eligibility for Co-Sponsorship: The co-sponsoring institution must have an approved Federal-wide Assurance with OHRP and be a recipient of HHS grants for human subject research. The selected co-sponsoring organization(s) shall furnish the necessary personnel, materials, services, and facilities to administer its responsibility for the workshop. These duties will be outlined in a co-sponsorship agreement with OHRP that will set forth the details of the co-sponsored activity, including the requirements that any fees collected by the co-sponsor shall be limited to the amount necessary to cover the co-sponsor's related conference expenses.

Co-sponsoring institutions will be asked to sign a Co-Sponsorship Agreement with HHS. This Co-Sponsorship Agreement does not represent an endorsement by OHRP of the co-sponsors' policies, positions, or activities. Additionally, this agreement will not affect any determination concerning activities by the co-sponsors that are regulated by OHRP.

The following Model Co-Sponsorship Agreement is presented only as an example. The assignment of duty and responsibilities in the Agreement will be discussed and agreed upon with each co-sponsor on a case by case basis and as applicable.

Model Co-Sponsorship Agreement

The Office for Human Research Protections (OHRP) and [co-sponsor] (if more than one co-sponsor, include all names followed by ''jointly referred to as co-sponsoring institutions'') agree to co-sponsor an Educational Workshop according to the understanding expressed below:

1. Background

The event is an OHRP Educational Workshop/Event tentatively titled, [title].

The Workshop/Event will be held on [Date] at [Location].

The Workshop/Event is a [half/1-day] educational outreach initiative that provides education and training focusing on the HHS policies and regulations on human research protections and their applicability. The Workshop/Event is designed for professionals engaged in human subject research, including, but not limited to, institutional review board (IRB) chairs, members and staff, investigators and research staff, and institutional officials.

The co-sponsoring institution for this educational activity, [co-sponsor], has an approved Federal-wide Assurance with OHRP and is a recipient of HHS grants for human subject research. OHRP has collaborated with [co-sponsor] (if more than one co-sponsor, include, [co-sponsor], and others) to develop a comprehensive agenda that addresses the provisions of the HHS Protection of Human Subjects Regulations, 45 CFR part 46, and the ethical principles of The Belmont Report.

OHRP fulfills its mission pursuant to 42 U.S.C. 289 by providing an education program where clarification and guidance with respect to ethical issues raised in connection with biomedical or behavioral research involving human subjects can be addressed. This workshop/event co-sponsored with [co-sponsor] is an important component of the OHRP educational program for fiscal year [year].

2. Responsibilities for Developing the Event

OHRP and [co-sponsor] have collaborated, and will continue to