

Prevention (CDC), and the Director, National Centers for Injury Prevention and Control (NCIPC) regarding feasible goals for the prevention and control of injury. The committee makes recommendations regarding policies, strategies, objectives, and priorities, and reviews progress toward injury prevention and control.

Matters to be Discussed: The meeting will open to the public. The Advisory Committee for Injury Prevention and Control (ACIPC) will be discussing partnership activities and how the ACIPC can advance the field of injury prevention and control. Agenda items are subject to change as priorities dictate.

Contact Person for More Information: Ms. Amy Harris, Executive Secretary, ACIPC, NCIPC, CDC, 4770 Buford Highway, NE., M/S K61, Atlanta, Georgia 30341-3724, telephone (770) 488-4936.

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.

Dated: November 5, 2007.

Elaine L. Baker,

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

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Board of Scientific Counselors, National Institute for Occupational Safety and Health (BSC, NIOSH)

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 for the aforementioned committee:

Time and Date: 10 a.m.-2 p.m., December 13, 2007.

Place: Holiday Inn Capitol, 550 C Street, SW., Washington, DC 20024.

Status: Open to the public, limited only by the space available. The meeting room accommodates approximately 50 people. Teleconference available toll-free; please dial (888) 677-1819, Participant Pass Code 25404.

Purpose: The Secretary, the Assistant Secretary for Health, and by delegation the Director, Centers for Disease Control and Prevention, are authorized under sections 301 and 308 of the Public Health Service Act to conduct directly or by grants or contracts, research, experiments, and demonstrations relating to occupational safety and health and to mine health. The Board of Scientific Counselors shall provide guidance to the Director, National Institute for Occupational Safety and Health on research and prevention programs. Specifically, the Board shall

provide guidance on the Institute's research activities related to developing and evaluating hypotheses, systematically documenting findings and disseminating results. The Board shall evaluate the degree to which the activities of the National Institute for Occupational Safety and Health: (1) Conform to appropriate scientific standards, (2) address current, relevant needs, and (3) produce intended results.

Matters To Be Discussed: NIOSH Response to the National Academies of Science Program Reviews.

Agenda items are subject to change as priorities dictate.

For Further Information Contact: Roger Rosa, Executive Secretary, BSC, NIOSH, CDC, 395 E Street, SW., Suite 9200, Patriots Plaza Building, Washington, DC 20201, telephone (202) 245-0655, fax (202) 245-0664.

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 CDC and the Agency for Toxic Substances and Disease Registry.

Dated: November 5, 2007.

Elaine L. Baker,

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

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

Privacy Act of 1974; Report of a Modified System of Records

AGENCY: Centers for Medicare & Medicaid Services (CMS), Department of Health and Human Services (HHS).

ACTION: Notice of a Modified System of Records (SOR).

SUMMARY: In accordance with the requirements of the Privacy Act of 1974, we are proposing to modify an existing SOR titled, "Individuals Authorized Access to Centers for Medicare & Medicaid Services (CMS) Computer Services (IACS), System No. 09-70-0064," most recently modified at 67 FR 48911 (July 26, 2002). We propose to assign a new CMS identification number to this system to simplify the obsolete and confusing numbering system originally designed to identify the Bureau, Office, or Center that maintained information in the Health Care Financing Administration systems of records. The new identifying number for this system should read: System No. 09-70-0538.

We propose to broaden the scope of this system to include a CMS service

planned to provide a centralized user provisioning and administration service that supports the creation, deletion, and lifecycle management of enterprise identities. This service creates accounts, supports Role Based Access Control (RBAC), and provides business application integration points. RBAC is a form flow approval process and enterprise identity audit and recertification based on the role of the individual. The business application integration point allows business application owners to use the form flow process of the user provisioning service to approve or deny requests for access to business applications. This modification will permit CMS to implement a unified framework for managing user information and access rights, for those individuals who apply for and are granted access across multiple CMS systems and business contexts.

We propose to modify existing routine use number 1 that permits disclosure to agency contractors and consultants to include disclosure to CMS grantees who perform a task for the agency. CMS grantees, charged with completing projects or activities that require CMS data to carry out that activity, are classified separate from CMS contractors and/or consultants. The modified routine use will remain as routine use number 1. We will delete routine use number 2 authorizing disclosure to support constituent requests made to a congressional representative. If an authorization for the disclosure has been obtained from the data subject, then no routine use is needed. The Privacy Act allows for disclosures with the "prior written consent" of the data subject. Finally, we will delete the section titled "Additional Circumstances Affecting Routine Use Disclosures," that addresses "Protected Health Information (PHI)" and "small cell size." The requirement for compliance with HHS regulation "Standards for Privacy of Individually Identifiable Health Information" does not apply because this system does not collect or maintain PHI. In addition, our policy to prohibit release if there is a possibility that an individual can be identified through "small cell size" is not applicable to the data maintained in this system.

We are modifying the language in the remaining routine uses to provide a proper explanation as to the need for the routine use and to provide clarity to CMS's intention to disclose individual-specific information contained in this system. The routine uses will then be prioritized and reordered according to their usage. We will also take the