ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)
Safety and Health Managers	Occupational Safety and Health Program Survey.	4,404	1	20/60
	Informed Consent Form	4,404	1	2/60
	Non Responder Interview	792	1	5/60

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. 2015–25194 Filed 10–2–15; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Disease, Disability, and Injury Prevention and Control Special Emphasis Panel: Initial review

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 a meeting for the initial review of applications in response to Funding Opportunity Announcement (FOA) PS16–001, Minority HIV/AIDS Research Initiative (MARI) to Build HIV Prevention Treatment and Research Capacity in Disproportionately Affected Black and Hispanic Communities and Among Historically Underrepresented Researchers.

DATES: 10:00 a.m.–5:00 p.m., December 9–10, 2015 (Closed).

ADDRESSES: Teleconference.

FOR FURTHER INFORMATION CONTACT: Gregory Anderson, M.S., M.P.H., Scientific Review Officer, National Center for HIV/AIDS, Viral Hepatitis, STD, and TB Prevention (NCHHSTP), CDC, 1600 Clifton Road NE., Mailstop E60, Atlanta, Georgia 30333, Telephone: (404) 718–8833.

SUPPLEMENTARY INFORMATION:

Status: The meeting 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, CDC, pursuant to Public Law 92–463.

Matters for Discussion: The meeting will include the initial review, discussion, and evaluation of applications received in response to ''Minority HIV/AIDS Research Initiative (MARI) to Build HIV Prevention Treatment and Research Capacity in Disproportionately Affected Black and Hispanic Communities and Among Historically Underrepresented Researchers'', FOA PS16–001.

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.

Catherine Ramadei,

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

[FR Doc. 2015–25274 Filed 10–2–15; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[Docket No. CDC–2015–0016]

Final Revised Vaccine Information Materials for Seasonal Influenza Vaccines

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

ACTION: Notice.

SUMMARY: Under the National Childhood Vaccine Injury Act (NCVIA) (42 U.S.C. 300aa–26), the HHS/CDC must develop vaccine information materials that all health care providers are required to give to patients/parents prior to administration of specific vaccines. On May 20, 2015, CDC published a notice in the **Federal Register** (80 FR 29009) seeking public comments on proposed new vaccine information materials for inactivated and live attenuated influenza vaccines. Following review of comments submitted and consultation as required under the law, CDC has finalized the materials. Copies of the final vaccine information materials for inactivated and live attenuated influenza vaccines are available to download from *http://www.cdc.gov/vaccines/hcp/vis/index.html* or *http://www.regulations.gov* (see Docket Number CDC–2015–0016).

DATES: Beginning no later than March 1, 2016, each health care provider who administers any seasonal influenza vaccine to any child or adult in the United States shall provide copies of the relevant revised vaccine information materials contained in this notice, in conformance with the August 7, 2015 CDC Instructions for the Use of Vaccine Information Statements prior to providing such vaccinations. These revised vaccine information materials may also be used earlier than that date. Prior to March 1, 2016, the previous edition of these two VISs can be used.

FOR FURTHER INFORMATION CONTACT: Suzanne Johnson-DeLeon (*msj1@cdc.gov*), National Center for Immunization and Respiratory Diseases, Centers for Disease Control and Prevention, Mailstop A–19, 1600 Clifton Road, NE., Atlanta, Georgia 30329.

SUPPLEMENTARY INFORMATION: The National Childhood Vaccine Injury Act of 1986 (Pub. L. 99–660), as amended by section 708 of Public Law 103–183, added section 2126 to the Public Health Service Act. Section 2126, codified at 42 U.S.C. 300aa–26, requires the Secretary of Health and Human Services to develop and disseminate vaccine information materials for distribution by all health care providers in the United States to any patient (or to the parent or legal representative in the case of a child) receiving vaccines covered under the National Vaccine Injury Compensation Program (VICP).

Development and revision of the vaccine information materials, also known as Vaccine Information Statements (VIS), have been delegated by the Secretary to the Centers for Disease Control and Prevention (CDC). Section 2126 requires that the materials be developed, or revised, after notice to the public, with a 60-day comment