

**DEPARTMENT OF HEALTH AND HUMAN SERVICES****Centers for Disease Control and Prevention****Disease, Disability, and Injury Prevention and Control; Special Emphasis Panel (SEP): Initial Review**

The meeting announced below concerns Public Health Research on Modifiable Risk Factors for Spina Bifida, DD15–001, 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 the aforementioned meeting:

*Time and Date:* 10:00 a.m.–6:00 p.m., March 19, 2015 (Closed)

*Place:* Teleconference.

*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 Pub. L. 92–463.

*Matters for Discussion:* The meeting will include the initial review, discussion, and evaluation of applications received in response to “Public Health Research on Modifiable Risk Factors for Spina Bifida, DD15–001, initial review.”

*Contact Person for More Information:* M. Chris Langub, Ph.D., Scientific Review Officer, CDC, 4770 Buford Highway NE., Mailstop F–80, Atlanta, Georgia 30341, Telephone: (770) 488–3585, [EEO6@cdc.gov](mailto:EEO6@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.

**Elaine L. Baker,**

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

[FR Doc. 2015–03254 Filed 2–17–15; 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 (SEP): Initial Review**

The meeting announced below concerns Comparison and Validation of Screening Tools for Substance Use Among Pregnant Women, DP15–003, 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 the aforementioned meeting:

**Time and Date**

9:00 a.m.–6:30 p.m., March 18, 2015 (Closed)

*Place:* Teleconference.

*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 Pub. L. 92–463.

*Matters for Discussion:* The meeting will include the initial review, discussion, and evaluation of applications received in response to “Comparison and Validation of Screening Tools for Substance Use among Pregnant Women, DP15–003, initial review.”

*Contact Person for More Information:* M. Chris Langub, Ph.D., Scientific Review Officer, CDC, 4770 Buford Highway NE., Mailstop F–80, Atlanta, Georgia 30341, Telephone: (770) 488–3585, [EEO6@cdc.gov](mailto:EEO6@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.

**Elaine L. Baker,**

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

[FR Doc. 2015–03255 Filed 2–17–15; 8:45 am]

**BILLING CODE 4163–18–P**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES****Centers for Disease Control and Prevention****Advisory Committee to the Director (ACD), Centers for Disease Control and Prevention—State, Tribal, Local and Territorial (STLT) Subcommittee**

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 subcommittee:

*Time and Date:* 3:30 p.m.—5:00 p.m. EDT, March 13, 2015.

*Place:* This meeting will be held by teleconference.

*Status:* This meeting is open to the public, limited only by the availability of telephone ports. The public is welcome to participate during the public comment, which is tentatively scheduled from 4:45 p.m. to 4:55 p.m. To participate on the 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 and 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 Public Health Surveillance, Public Health Finance and Social Determinants of Health as they relate to STLT public health agencies.

The agenda is subject to change as priorities dictate.

*Contact Person for More Information:* Judith Monroe, M.D., Designated Federal Officer, State, Tribal, Local and Territorial Subcommittee, Advisory Committee to the Director, CDC, 1600 Clifton Road NE., M/S E–70, Atlanta, Georgia 30333, Telephone (404) 498–0300, Email: [OSTLTSDirector@cdc.gov](mailto:OSTLTSDirector@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.

**Elaine L. Baker,**

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

[FR Doc. 2015-03252 Filed 2-17-15; 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 (SEP): Initial Review**

The meeting announced below concerns Establishing a Vision and Eye Health Surveillance System for the Nation, DP15-004, 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 the aforementioned meeting:

*Time and Date:* 10:00 a.m.–5:00 p.m., March 10, 2015 (Closed).

*Place:* Teleconference.

*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 “Establishing a Vision and Eye Health Surveillance System for the Nation, DP15-004, initial review.”

*Contact Person for More Information:* M. Chris Langub, Ph.D., Scientific Review Officer, CDC, 4770 Buford Highway NE., Mailstop F-80, Atlanta, Georgia 30341, Telephone: (770) 488-3585, [EEO6@cdc.gov](mailto:EEO6@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.

**Elaine L. Baker,**

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

[FR Doc. 2015-03256 Filed 2-17-15; 8:45 am]

**BILLING CODE 4163-18-P**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Administration for Children and Families**

**Proposed Information Collection Activity: Comment Request**

*Title:* ANA Reviewer Profile for Panel Review Participation Form.

*OMB No.:* OMB No: 0970-265.

*Description:* The Department of Health and Human Services (HHS), Administration for Children and Families (ACF) proposes to revise the ANA Reviewer Profile for Panel Review Participation Form. The ANA Reviewer Profile for Panel Review Participation Form is used to collect information from prospective proposal reviewers in compliance with 42 U.S.C. 2991d 1. First time reviewers will be required to complete all sections of the form while returning reviewers will be required to complete the first section of the document and other necessary updates. The form allows the Commissioner of ANA to select qualified people to review grant applications submitted in response to funding opportunity announcements for ANA’s primary programs: Social and Economic Development Strategies (SEDS); Native Language Preservation and Maintenance; and Environmental Regulatory Enhancement. The panel review process is a legislative mandate in the ANA grant funding process.

*Respondents:* All US citizens including: Native Americans, Native Alaskans, Native Hawaiians and other Pacific Islanders.

**ANNUAL BURDEN ESTIMATES**

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
ANA Consultant and Evaluator Qualifications Form .....	300	1	.5	150

Estimated Total Annual Burden Hours: 150.

In compliance with the requirements of Section 506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Administration for Children and Families is soliciting public comment on the specific aspects of the information collection described above. Copies of the proposed collection of information can be obtained and comments may be forwarded by writing to the administration for Children and Families, Office of Planning, Research and Evaluation, 370 L’Enfant Promenade SW., Washington, DC 20447, Attn: ACF Reports Clearance Officer. Email address: [infocollection@acf.hhs.gov](mailto:infocollection@acf.hhs.gov). All requests should be

identified by the title of the information collection.

The Department specifically requests comments 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) the quality, utility, and clarity of the information to be collected; and (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. Consideration will be given to

comments and suggestions submitted within 60 days of this publication.

**Robert Sargis,**

*Report Clearance Officer.*

[FR Doc. 2015-03302 Filed 2-17-15; 8:45 am]

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