

and CNF and these biomarkers of early effect, considering potential confounding factors such as smoking, age, gender, and workplace co-exposures, including non-engineered ultrafine particles.

The proposed project supports the NIOSH legislatively mandated industrywide studies program that conducts epidemiological and exposure

assessment research studies to identify the occupational causes of disease in the working population and their offspring and to effectively communicate study results to workers, scientists, industry, and the public.

The questionnaire will be administered one time only, at the worksite, to 100 workers involved in the production and use of CNT or CNT. The

study will be carried out during the participants' regular work shift. There is no cost to respondents or their employers other than their time. We estimate that the average burden per response to be 22 minutes, and that the total burden to all respondents will be 37 hours (see table below).

**ESTIMATED ANNUALIZED BURDEN HOURS**

Respondents	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total burden (in hours)
Nanomaterials Workers .....	100	1	22/60	37
<b>Total</b> .....				<b>37</b>

Dated: September 14, 2012.

**Ron A. Otten,**

Director, Office of Scientific Integrity (OSI), Office of the Associate Director for Science (OADS), Office of the Director, Centers for Disease Control and Prevention.

[FR Doc. 2012-23194 Filed 9-19-12; 8:45 am]

**BILLING CODE 4163-18-P**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Centers for Disease Control and Prevention**

**Healthcare Infection Control Practices Advisory Committee (HICPAC)**

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:

*Times and Dates:* 9:00 a.m.–5:00 p.m., October 11, 2012; 9:00 a.m.–12:00 p.m., October 12, 2012

*Place:* Renaissance Washington, DC Dupont Circle Hotel, City Center Ballroom, 1143 New Hampshire Avenue NW., Washington, District of Columbia 20037.

*Status:* Open to the public, limited only by the space available. Please register for the meeting at [www.cdc.gov/hicpac](http://www.cdc.gov/hicpac).

*Purpose:* The Committee is charged with providing advice and guidance to the Secretary, Department of Health and Human Services (HHS); the Director, Centers for Disease Control and Prevention (CDC); the Deputy Director, Office of Infectious Diseases (OID), CDC; and the Director, National Center for Emerging and Zoonotic Infectious Disease (NCEZID), CDC, regarding (1) the practice of infection control; (2) strategies for surveillance, prevention, and control of healthcare-associated infections (e.g., nosocomial infections) antimicrobial resistance and related events in settings where healthcare is provided, including

hospitals, ambulatory and long-term care facilities, and home health agencies; and (3) periodic updating of existing guidelines, development of new guidelines, guideline evaluation; and other policy statements regarding the prevention of healthcare-associated infections an healthcare-related conditions.

*Matters To Be Discussed:* The agenda will include updates on CDC's activities for healthcare associated infections (HAI), an update on the draft guideline for prevention of infections among patients in neonatal intensive care units (NICU), draft guideline for the prevention of surgical site infections, draft guidance for facility adjudication of infection data, and an update from the HICPAC surveillance working group.

Agenda items are subject to change as priorities dictate.

*Contact Person for More Information:* Erin Stone, M.S., HICPAC, Division of Healthcare Quality Promotion, NCEZID, CDC, 1600 Clifton Road, NE., Mailstop A-07, Atlanta, Georgia 30333 Telephone (404) 639-4045. Email: [hicpac@cdc.gov](mailto:hicpac@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.

Dated: September 12, 2012.

**Elaine L. Baker,**

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

[FR Doc. 2012-23193 Filed 9-19-12; 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—Ethics Subcommittee (ES)**

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:* 8:30 a.m.–2:30 p.m., EDT, Thursday, October 11, 2012.

*Place:* CDC, Thomas R. Harkin Global Communications Center, Distance Learning Auditorium, 1600 Clifton Road, NE., Atlanta, GA 30333. This meeting is also available by teleconference. Please dial (877) 928-1204 and enter code 4305992.

*Status:* Open to the public, limited only by the space available. The meeting room accommodates approximately 60 people. To accommodate public participation in the meeting, a conference telephone line will be available. The public is welcome to participate during the public comment period. The public comment period is tentatively scheduled for 2 p.m.–2:10 p.m.

*Purpose:* The ES will provide counsel to the ACD, CDC, regarding a broad range of public health ethics questions and issues arising from programs, scientists and practitioners.

*Matters To Be Discussed:* Agenda items will include the following topics: Ethical considerations relating to use of travel restrictions for the control of communicable diseases; addition of ethics standards to the accreditation process for public health departments; approaches for evaluating the impact of public health ethics activities; progress on developing practical tools to assist state, tribal, local, and territorial health departments in their efforts to address public health ethics challenges; and strategies for

increasing collaboration between public health ethics and public health law.

The agenda is subject to change as priorities dictate.

*Contact Person for More Information:* For security reasons, members of the public interested in attending the meeting should contact Drue Barrett, Ph.D., Designated Federal Officer, ACD, CDC—ES, 1600 Clifton Road NE., M/S D-50, Atlanta, Georgia 30333. Telephone: (404) 639-4690. Email: [d Barrett@cdc.gov](mailto:d Barrett@cdc.gov). The deadline for notification of attendance is October 1, 2012.

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.

Dated: September 12, 2012.

**Elaine L. Baker,**

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

[FR Doc. 2012-23192 Filed 9-19-12; 8:45 am]

**BILLING CODE 4163-18-P**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Administration for Children and Families**

**Submission for OMB Review; Comment Request**

*Title:* Title IV–E Plan for Foster Care, Adoption Assistance, and, optional, Guardianship Assistance Programs.

*OMB No.:* 0980-0141.

*Description:* A title IV–E plan is required by section 471, part IV–E of the Social Security Act (the Act) for each public child welfare agency requesting Federal funding for foster care, adoption assistance and guardianship assistance under the Act. Section 479B of the Act provides for an Indian tribe, tribal organization or tribal consortium (Tribe) to operate a title IV–E program in the same manner as a State with minimal exceptions. The Tribe must have an approved title IV–E Plan. The title IV–E plan provides assurances the

programs will be administered in conformity with the specific requirements stipulated in title IV–E. The plan must include all applicable State or Tribal statutory, regulatory, or policy references and citations for each requirement as well as supporting documentation. A title IV–E agency may use the pre-print format prepared by the Children’s Bureau of the Administration for Children and Families or a different format, on the condition that the format used includes all of the title IV–E plan requirements of the law.

*Respondents:* Title IV–E agencies administering or supervising the administration of the title IV–E programs.

**ANNUAL BURDEN ESTIMATES**

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Title IV–E Plan .....	17	1	16	272

*Estimated Total Annual Burden Hours: 272.*

**Additional Information**

Copies of the proposed collection may be obtained 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. All requests should be identified by the title of the information collection. Email address: [infocollection@acf.hhs.gov](mailto:infocollection@acf.hhs.gov).

**OMB Comment**

OMB is required to make a decision concerning the collection of information between 30 and 60 days after publication of this document in the **Federal Register**. Therefore, a comment is best assured of having its full effect if OMB receives it within 30 days of publication. Written comments and recommendations for the proposed information collection should be sent directly to the following: Office of Management and Budget, Paperwork Reduction Project, Email: [OIRA\\_SUBMISSION@OMB.EOP.GOV](mailto:OIRA_SUBMISSION@OMB.EOP.GOV), Attn: Desk Officer for the

Administration for Children and Families.

**Robert Sargis,**

*Reports Clearance Officer.*

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**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Administration for Children and Families**

[CFDA Number 93.631]

**Announcement of the Award of a Single-Source Program Expansion Supplement Grant to the University of Boston for the Institute for Community Inclusion (ICI) in Boston, MA**

**AGENCY:** Administration on Developmental and Intellectual Disabilities (AIDD), ACF, HHS.

**ACTION:** Announcing the award a single-source program expansion supplement to the University of Massachusetts for the Institute for Community Inclusion in Boston, MA, to support the additional employment grants that will be awarded.

**SUMMARY:** The Administration for Children and Families (ACF), Administration on Developmental and Intellectual Disabilities (AIDD) announces the award of a grant in the amount of \$300,000 to the University of Massachusetts for the Institute for Community Inclusion, Boston, MA.

**DATES:** The project period for the award is from September 30, 2012 to September 29, 2013.

**FOR FURTHER INFORMATION CONTACT:** Ophelia McLain, Supervisory Program Specialist, Administration on Intellectual and Developmental Disabilities, 370 L’Enfant Promenade SW., 2nd Floor East, Washington, DC 20447. Telephone: 202-690-7025; Email: [ophelia.mclain@acf.hhs.gov](mailto:ophelia.mclain@acf.hhs.gov).

**SUPPLEMENTARY INFORMATION:** In September 2011, the Administration on Developmental and Intellectual Disabilities (AIDD) awarded a grant to the ICI to serve as the training and technical assistance (T/TA) provider to recipients of Partnerships in Employment Systems Change grants, also awarded that same year. AIDD has expanding the Employment efforts by awarding two additional Partnerships in Employment Systems Change grants in