

can also be submitted by e-mail to OCAS@CDC.GOV.

Dated: July 2, 2008.

John Howard,

Director, National Institute for Occupational Safety and Health.

[FR Doc. E8-16467 Filed 7-17-08; 8:45 am]

BILLING CODE 4163-19-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institute for Occupational Safety and Health; Final Effect of Designation of a Class of Employees for Addition to the Special Exposure Cohort

AGENCY: National Institute for Occupational Safety and Health (NIOSH), Department of Health and Human Services (HHS).

ACTION: Notice.

SUMMARY: The Department of Health and Human Services (HHS) gives notice concerning the final effect of the HHS decision to designate a class of employees at the Nuclear Materials and Equipment Corporation (NUMEC) facility in Parks Township, Pennsylvania, as an addition to the Special Exposure Cohort (SEC) under the Energy Employees Occupational Illness Compensation Program Act of 2000. On May 30, 2008, as provided for under 42 U.S.C. 7384q(b), the Secretary of HHS designated the following class of employees as an addition to the SEC:

All Atomic Weapons Employer (AWE) employees who worked at the Nuclear Materials and Equipment Corporation (NUMEC) facility in Parks Township, Pennsylvania, from June 1, 1960, through December 31, 1980, for a number of work days aggregating at least 250 work days occurring either solely under this employment or in combination with work days within the parameters established for one or more other classes of employees in the Special Exposure Cohort.

This designation became effective on June 29, 2008, as provided for under 42 U.S.C. 7384(14)(C). Hence, beginning on June 29, 2008, members of this class of employees, defined as reported in this notice, became members of the Special Exposure Cohort.

FOR FURTHER INFORMATION CONTACT:

Larry Elliott, Director, Office of Compensation Analysis and Support, National Institute for Occupational Safety and Health (NIOSH), 4676 Columbia Parkway, MS C-46, Cincinnati, OH 45226, telephone 1-800-CDC-INFO (1-800-232-4636) or directly at 1-513-533-6800 (this is not a toll-free number). Information requests

can also be submitted by e-mail to OCAS@CDC.GOV.

Dated: July 2, 2008.

John Howard,

Director, National Institute for Occupational Safety and Health.

[FR Doc. E8-16468 Filed 7-17-08; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Office of the Secretary

Findings of Scientific Misconduct

AGENCY: Office of the Secretary, HHS.

ACTION: Notice.

SUMMARY: Notice is hereby given that the Office of Research Integrity (ORI) and the Assistant Secretary for Health have taken final action in the following case:

J. Keith Hampton, St. Luke's Hospital: Based on the report of an investigation conducted by St. Luke's Hospital (SLH) in Chesterfield, MO, and additional analysis conducted by the Office of Research Integrity (ORI) during its oversight review, the U.S. Public Health Service (PHS) found that J. Keith Hampton, MSN, APRN, former Clinical Research Associate, SLH, engaged in scientific misconduct in research supported by National Cancer Institute (NCI), National Institutes of Health (NIH), awards U10 CA69651, U10 CA12027, and U10 CA33601.

PHS found that Mr. Hampton engaged in scientific misconduct by falsifying and fabricating data that were reported to the National Surgical Adjuvant Breast & Bowel Project (NSABP) and Cancer and Leukemia Group B (CALGB) cooperative research groups.

Specifically, PHS found that:

1. For protocol CALGB 90206,

Respondent:

(a) Falsified a patient's CT scan reports and registration forms and reported the falsified CT scan reports and registration worksheet to CALGB.

(b) Falsified a patient's performance status records (giving 80% performance status) and registration forms and reported the falsified performance status report and registration form to CALGB.

2. For protocol NSABP B-35,

Respondent:

(a) Falsified eligibility data related to hematology and chemistry assays and to the performance of a pelvic exam on one patient's registration form and reported the falsified registration forms to the National Cancer Institute Cancer Trial Support Unit (CTSU).

(b) Falsified pelvic exam eligibility on a second patient's registration form and

reported the falsified registration form to the CTSU.

(c) Falsified hematology and chemistry assay eligibility on a third patient's registration form and reported the falsified registration form to the CTSU.

3. For protocol NSABP B-36, Respondent falsified a patient's multigated acquisition test (MUGA—a test of heart function) records, cardiac function, and registration forms, certified the patient's eligibility, and reported the falsified MUGA test, cardiac function, and registration forms to the CTSU.

4. For protocol NSABP B-38, Respondent falsified hematology, chemistry, and MUGA eligibility for a patient on the registration form and reported the falsified registration form to the CTSU.

5. For protocol NSABP C-08, Respondent:

(a) Falsified urine protein/creatinine ratio eligibility for one patient on the registration form and reported the falsified registration form to the CTSU.

(b) Falsified urine protein/creatinine ratio eligibility for a second patient on the registration form and reported the falsified registration form to the CTSU.

(c) Falsified claims of the urine protein/creatinine ratio and PT(INR) eligibility for a third patient on the registration form and reported the falsified registration form to the CTSU.

6. For protocol NSABP R-04, Respondent falsified a patient's colonoscopy report and eligibility at registration and reported the falsified colonoscopy report and registration form to the CTSU.

Mr. Hampton has entered into a Voluntary Exclusion Agreement (Agreement) in which he has voluntarily agreed for a period of three (3) years, beginning on June 17, 2008:

(1) To exclude himself from any contracting or subcontracting with any agency of the United States Government and from eligibility or involvement in nonprocurement programs of the United States Government referred to as "covered transactions" pursuant to HHS' Implementation (2 CFR part 376 *et seq.*) of OMB Guidelines to Agencies on Governmentwide Debarment and Suspension (2 CFR part 180); and

(2) To exclude himself from serving in any advisory capacity to PHS, including but not limited to service on any PHS advisory committee, board, and/or peer review committee, or as a consultant or contractor to PHS.

FOR FURTHER INFORMATION CONTACT:

Director, Division of Investigative Oversight, Office of Research Integrity,

1101 Wootton Parkway, Suite 750,
Rockville, MD 20852, (240) 453-8800.

Chris B. Pascal,
Director, Office of Research Integrity.
[FR Doc. E8-16357 Filed 7-17-08; 8:45 am]
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**DEPARTMENT OF HEALTH AND
HUMAN SERVICES**

**Centers for Disease Control and
Prevention**

**Board of Scientific Counselors for the
National Center for Public Health
Informatics**

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 committee meeting:

Name: Board of Scientific Counselors for the National Center for Public Health Informatics.

Time and Date: 5 p.m.-9 p.m., August 27, 2008.

Place: The Westin Peachtree Plaza, 210 Peachtree Street, Atlanta, Georgia 30303.

Status: Open to the public, limited only by the space available.

Purpose: The committee shall advise the Secretary, HHS, and the Director, CDC, concerning strategies and goals for the programs and research within the national centers; shall conduct peer-review of scientific programs; and monitor the overall strategic direction and focus of the national centers. The board, after conducting its periodic reviews, shall submit a written description of the results of the review and its recommendations to the Director, CDC. The board shall perform second-level peer review of applications for grants-in-aid for research and research training activities, cooperative agreements, and research contract proposals relating to the broad areas within the national centers.

Matters To Be Discussed: The agenda will include an overview of the National Center for Public Health Informatics (NCPHI), including its mission, scope and goals. Detailed discussions will take place on the following issues: BioSense Strategic Planning, Open Source Models, and Organizational Issues for NCPHI.

Agenda items are subject to change as priorities dictate.

FOR FURTHER INFORMATION CONTACT:
Thomas G. Savel, M.D., Designated Federal Official, National Center for Public Health Informatics, CDC, 1600 Clifton Road, NE., MS E78, Atlanta, Georgia 30333. Telephone 404/498-2475.

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: July 8, 2008.

Diane Allen,
Acting Director, Management Analysis and Services Office, Centers for Disease Control and Prevention.
[FR Doc. E8-16449 Filed 7-17-08; 8:45 am]
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**DEPARTMENT OF HEALTH AND
HUMAN SERVICES**

**Centers for Disease Control and
Prevention**

**Safety and Occupational Health Study
Section: Notice of Charter Renewal**

This gives notice under the Federal Advisory Committee Act (Pub. L. 92-463) of October 6, 1972, that the Safety and Occupational Health Study Section, Centers for Disease Control and Prevention, Department of Health and Human Services, has been renewed for a 2-year period through June 30, 2010.

For More Information Contact: Price Connor, PhD, Executive Secretary, Safety and Occupational Health Study Section, Department of Health and Human Services, 1600 Clifton Road, NE., Mailstop E74, Atlanta, Georgia 30333, telephone 404/498-2511 or fax 404/498-2571.

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: July 14, 2008.

Diane Allen,
Acting Director, Management Analysis and Services Office, Centers for Disease Control and Prevention.
[FR Doc. E8-16450 Filed 7-17-08; 8:45 am]
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**DEPARTMENT OF HEALTH AND
HUMAN SERVICES**

**Administration for Children and
Families**

**Proposed Information Collection
Activity; Comment Request**

Proposed Projects

Title: Voluntary Surveys of Program Partners to Implement Executive Order 12862.

OMB No.: 0980-0266.

Description: Under the provisions of the Federal Paperwork Reduction Act of 1995 (Pub. L. 104-13), the Administration for Children and Families (ACF) is requesting clearance for instruments to implement Executive Order 12862 within ACF. The purpose of the data collection is to obtain customer satisfaction information from those entities who are funded to be our partners in the delivery of services to the American public. ACF partners are those entities that receive funding to deliver services or assistance from ACF programs. Examples of partners are state and local governments, territories, service providers, Indian Tribes and Tribal organizations, grantees, researchers, or other intermediaries serving target populations identified by and funded directly or indirectly by ACF. The surveys will obtain information about how well ACF is meeting the needs of our partners in operating the ACF programs.

Respondents: State, Local, & Tribal Govt. or not-for-profit Organizations

ANNUAL BURDEN ESTIMATES				
Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
State Governments, Territories and District of Columbia	54	10	1	540
Head Start Grantees and Delegates	200	1	0.50	100
Other Discretionary Grant Programs	200	10	0.50	1,000
Indian Tribes and Tribal Organizations	25	10	0.50	125

Estimated Total Annual Burden Hours: 1,765.

In compliance with the requirements of Section 506(c)(2)(A) of the Paperwork

Reduction Act of 1995, the Administration for Children and