

Dated: November 19, 2004.

Paul L. Johnson,

Project Clearance Liaison, NICHD, National Institutes of Health.

[FR Doc. 04-26539 Filed 12-1-04; 8:45 am]

BILLING CODE 4140-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

[OMB No. 0925-0454]

Submission for OMB Review; Comment Request; Case-Cohort Study of Cancer and Related Disorders Among Benzene-Exposed Workers in China

SUMMARY: Under the provisions of Section 3507(a)(1)(D) of the Paperwork Reduction Act of 1995, the National Cancer Institute (NCI), the National Institutes of Health (NIH) has submitted to the Office of Management and Budget (OMB) a request for review and approval of the information collection listed below. This proposed information collection was previously published in the **Federal Register** and allowed 60 days for public comment. No public comments were received. The purpose

of this notice is to allow an additional 30 days for public comment. The National Institutes of Health may not conduct or sponsor, and the respondent is not required to respond to, an information collection that has been extended, revised, or implemented on or after October 1, 1995, unless it displays a currently valid OMB control number.

Proposed Collection: Title: Case-Cohort Study of Cancer and Related Disorders Among Benzene-Exposed Workers in China. **Reinstatement With Change. Need and Use of Information Collection:** A case-cohort study will be performed to examine the risks of lymphohematopoietic cancers, other lymphohematopoietic disorders, benzene poisoning, and lung cancer among workers exposed to benzene. The study will attempt to determine with greater precision the risks of these disorders at low levels of benzene exposure, and to characterize the dose and time-specific relationship between benzene exposure and disease risk. Cases and controls will be selected from an existing cohort of 75,000 benzene-exposed workers and 36,000 comparison workers in 12 Chinese cities. There are 2 changes to the study from that previously approved by OMB in July 2001: (1) 386 more subjects

(including 155 more cases with benzene poisoning, 111 more cases with lung cancer, and 120 more controls) will be evaluated in the currently planned case-cohort study, which is now targeting a total of 2,156 subjects compared with 1,770 subjects as previously estimated; and (2) the questionnaire has been revised somewhat, although the average total time estimated for a subject to complete the questionnaire is unchanged from previously.

Frequency of Response: Single-time study. **Affected Public:** Individuals or households. **Type of Respondents:** Cases with lymphohematopoietic malignancies and related disorders, benzene poisoning and lung cancer among Chinese benzene-exposed and comparison workers; controls consist of a random sample of the Chinese worker cohort. The annual reporting burden is as follows: **Estimated Number of Respondents:** 862; **Estimated Number of Responses per Respondent:** 1; **Average Burden Hours per Response:** 0.3674; and **Estimated Total Annual Burden Hours Requested:** 396. The annualized cost to respondents is estimated at \$476. There are no Capital Costs to report. There are also no Operating or Maintenance Costs to report.

ESTIMATES OF HOUR BURDEN: BURDEN REQUESTED

Type of respondents	Estimated number of respondents	Estimated frequency of responses per respondent	Average burden time (in hours) per response	Estimated average annual hour burden
Workers in factories in China using or producing benzene and in comparison factories in which no benzene is used	1,078	1	0.37	396

Request for Comments: Written comments and/or suggestions from the public and affected agencies are invited on one or more of the following points: (1) Whether the proposed collection of information is necessary for the proper performance of the function of the agency, including whether the information will have practical utility; (2) the accuracy of the agency's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) ways to minimize the burden of the collection of information on those who are to respond, including the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology.

Direct Comments To OMB: Written comments and/or suggestions regarding

the item(s) contained in this notice, especially regarding the estimated public burden and associated response time, should be directed to the: Office of Management and Budget, Office of Regulatory Affairs, New Executive Office Building, Room 10235, Washington, DC 20503, Attention: Desk Officer for NIH. To request more information on the proposed project or to obtain a copy of the data collection plans and instruments, contact: Dr. Richard Hayes, OEB/EBP/DCEG/NCI 6120 Executive Boulevard, EPS Room 8114, Bethesda, MD 20892, or call non-toll-free number (301) 435-3973 or e-mail your request, including your address to: HayesR@mail.nih.gov.

Comments Due Date: Comments regarding this information collection are best assured of having their full effect if received within 30 days of the date of this publication.

Dated: November 19, 2004.

Rachelle Ragland-Greene,

NCI Project Clearance Liaison, National Institutes of Health.

[FR Doc. 04-26540 Filed 12-1-04; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Center for Scientific Review; Notice of Closed Meetings

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. Appendix 2), notice is hereby given of the following meetings.

The meetings will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C.,

as amended. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: Center for Scientific Review Special Emphasis Panel, Therapeutic Developments for Dermatologic and Rheumatologic Diseases Small Business.

Date: November 30, 2004.

Time: 8 a.m. to 5 p.m.

Agenda: To review and evaluate grant applications.

Place: Bethesda Marriott Suites, 6711 Democracy Boulevard, Bethesda, MD 20817.

Contact Person: Daniel F. McDonald, PhD, Chief, Renal and Urological Sciences IRG, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 4214, MSC 7814, Bethesda, MD 20892, (301) 435-1215, mcdonald@csr.nih.gov.

This notice is being published less than 15 days prior to the meeting due to the timing limitations imposed by the review and funding cycle.

Name of Committee: Center for Scientific Review Special Emphasis Panel, Visual Motion Processing.

Date: December 6, 2004.

Time: 11 a.m. to 12 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892, (Telephone conference call).

Contact Person: Michael A. Steinmetz, PhD, Scientific Review Administrator, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 5172, MSC 7844, Bethesda, MD 20892, (301) 435-1247, steinmem@csr.nih.gov.

This notice is being published less than 15 days prior to the meeting due to the timing limitations imposed by the review and funding cycle.

Name of Committee: Center for Scientific Review Special Emphasis Panel, Clinical Immunotherapy.

Date: December 10, 2004.

Time: 3 p.m. to 5 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892, (Telephone conference call).

Contact Person: Sharon K. Gubanich, PhD, Scientific Review Administrator, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 6204, MSC 7804, Bethesda, MD 20892, (301) 435-1767, guganics@csr.nih.gov.

This notice is being published less than 15 days prior to the meeting due to the timing limitations imposed by the review and funding cycle.

Name of Committee: Center for Scientific Review Special Emphasis Panel, Pharmacogenetics.

Date: December 15, 2004.

Time: 11:30 a.m. to 12:30 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892, (Telephone conference call).

Contact Person: Marcia Litwack, PhD, Scientific Review Administrator, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 6206, MSC 7804, Bethesda, MD 20892, (301) 435-1719, litwackm@csr.nih.gov. (Catalogue of Federal Domestic Assistance Program Nos. 93.306, Comparative Medicine; 93.333, Clinical Research, 93.306, 93.333, 93.337, 93.393-93.396, 93.837-93.844, 93.846-93.878, 93.892, 93.893, National Institutes of Health, HHS.)

Dated: November 23, 2004.

LaVerne Y. Stringfield,

Director, Office of Federal Advisory Committee Policy.

[FR Doc. 04-26538 Filed 12-1-04; 8:45 am]

BILLING CODE 4140-01-M

JUDICIAL CONFERENCE OF THE UNITED STATES

Hearings of The Judicial Conference Advisory Committees on Rules of Appellate, Bankruptcy, and Civil Procedure

AGENCY: Judicial Conference of the United States; Advisory Committees on Rules of Appellate, Bankruptcy, and Civil Procedure.

ACTION: Notice of proposed amendments and open hearings.

SUMMARY: The Advisory Committees on Rules of Appellate, Bankruptcy, and Civil Procedure have proposed amendments to the following rules:

Appellate Rule: 25.

Bankruptcy Rule: 5005.

Civil Rule: 5.

The text of the proposed rules amendments and the accompanying Committee Notes can be found at the United States Federal Courts' home page at <http://www.uscourts.gov/rules>.

The Judicial Conference Committee on Rules of Practice and Procedure submits these proposed rules amendments for public comment. All comments and suggestions with respect to them must be placed in the hands of the Secretary as soon as convenient and, in any event, not later than February 15, 2005. All written comments on the proposed rule amendments can be sent by one of the following three ways: by overnight mail to Peter G. McCabe, Secretary, Committee on Rules of Practice and Procedure of the Judicial Conference of the United States, Thurgood Marshall Federal Judiciary Building, Washington, DC 20544; by

electronic mail at <http://www.uscourts.gov/rules>; or by facsimile to Peter G. McCabe at (202) 502-1766. In accordance with established procedures all comments submitted on the proposed amendments are available to public inspection.

Public hearings are scheduled to be held on the amendments to:

- Appellate Rules in Washington, DC, on January 25, 2005;

- Bankruptcy Rules in Washington, DC, on February 3, 2005; and in San Francisco, California, on February 7, 2005; and

- Civil Rules in San Francisco, California, on January 12, 2005; in Dallas, Texas, on January 28, 2005; and in Washington, DC, on February 11, 2005.

Those wishing to testify should contact the Secretary at the address above in writing at least 30 days before the hearing.

FOR FURTHER INFORMATION CONTACT: John K. Rabiej, Chief, Rules Committee Support Office, Administrative Office of the United States Courts, Washington, DC 20544, telephone (202) 502-1820.

Dated: November 24, 2004.

John K. Rabiej,

Chief, Rules Committee Support Office.

[FR Doc. 04-26578 Filed 12-1-04; 8:45 am]

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DEPARTMENT OF LABOR

Employee Benefits Security Administration

Proposed Extension of Information Collection Request Submitted for Public Comment and Recommendations; Alternative Method of Compliance for Certain SEPs Pursuant to 29 CFR 2520.104-49

AGENCY: Employee Benefits Security Administration, Department of Labor.

ACTION: Notice.

SUMMARY: The Department of Labor, as part of its continuing effort to reduce paperwork and respondent burden, conducts a preclearance consultation program to provide the general public and Federal agencies with an opportunity to comment on proposed and continuing collections of information in accordance with the Paperwork Reduction Act of 1995 (PRA 95). This program helps to ensure that requested data can be provided in the desired format, reporting burden (time and financial resources) is minimized, collection instruments are clearly understood, and the impact of collection