phenotypes are biological traits that may be influenced by variation at fewer genes and may mediate different aspects of the disease. The intermediate phenotype measurements that we will collect include resting EEG phenotypes (log voltae alpha (LVA) and beta spectral power), ERPs and heart rate variability (HRV). LVA has been found to be more abundant in alcoholics with co-morbid anxiety disorders. Increased beta power has been associated with increased risk of relapse. P300 ERP amplitude is reduced in alcoholics and their alcohol-naïve children. HRV is a potential intermediate phenotype for alcoholism and major depression. We also propose to administer the Temperament and Character Inventory, a standard, survey-based measure of harm avoidance, novelty seeking, reward dependence, and persistence. The use of such intermediate phenotypes and personality measures is likely to increase our ability to find vulnerability genes for alcoholism. We will use these EEG and EKG intermediate phenotypes and personality dimensions in (1) candidate gene analyses and (2) linkage analyses, utilizing the existing DNA, in order to determine the genes that increase an individuals's risk for alcoholism and anxiety disorders.

The re-recruitment of the original study participants will start in spring 2003. The study is expected to run for 6 months. Frequency of response: Once per respondent. Affected Public: Individuals. Type of Respondents: Adults members of the Southeastern American Indian tribe who were participants in the original study.

The annual reporting burden is as follows: Estimated Number of Respondents: It is estimated, after a survey by tribal members, that we will be able to re-recruit approximately 280 of the 294 original participants. Estimated Number of Responses per Respondent: One response per respondent. Average Burden Hours per Response: Three hours per individual, for a total respondent burden of 840 hours. Estimated Total Annual Burden Hours Requested: 840 hours. There are no Costs to Respondents to report. There are no Capital Costs to report. There are no Operating or Maintenance costs to

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: Mary-Anne Enoch M.D., NIH/NIAAA/DICBR/ LNG, 12420 Parklawn Drive, Park 5 Building, Room 451, MSC 8110, Bethesda, MD 20892-8110, or e-mail vour request to: maenoch@niaaa.nih.gov. Dr. Enoch can

2727.

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.

be contacted by telephone at 301–496–

Dated: February 14, 2003.

Stephen Long,

Executive Officer, NIAAA.

[FR Doc. 03-8862 Filed 4-10-03; 8:45 am]

BILLING CODE 4140-01-M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Cancer Institute; Notice of Closed Meeting

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

The meeting 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 contract proposals and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning

individuals associated with the contract proposals, the disclosure of which would constitute a clear unwarranted invasion of personal privacy.

Name of Committee: National Cancer Institute Special Emphasis Panel, Early Clinical Trials of Imaging Agents.

Date: May 2, 2003.

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

Agenda: To review and evaluate contract proposals.

Place: Bethesda Marriott, 5151 Pooks Hill Road, Bethesda, MD 20814.

Contact Person: Kenneth L. Bielat, PhD, Scientific Review Administrator, Division of Extramural Activities, National Cancer Institute, National Institutes of Health, 6116 Executive Boulevard, Room 7147, Bethesda, MD 20892. (301) 496–7576. bielatk@mail.nih.gov.

(Catalogue of Federal Domestic Assistance Program Nos. 93.392, Cancer Construction; 93.393, Cancer Cause and Prevention Research; 93.394, Cancer Detection and Diagnosis research; 93.395, Cancer Treatment Research; 93.396, Cancer Biology Research; 93.397, Cancer Centers Support; 93.398, Cancer research Manpower; 93.399, Cancer Control, National Institutes of Health, HHS)

Dated: April 4, 2003.

LaVerne Y. Stringfield,

Director, Office of Federal Advisory Committee Policy.

[FR Doc. 03–8858 Filed 4–10–03; 8:45 am] BILLING CODE 4140–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Cancer Institute; Notice of Closed Meeting

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

The meeting 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: National Cancer Institute Special Emphasis Panel, Spores in Skin Cancer.

Date: May 15-16, 2003.

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

Agenda: To review and evaluate grant applications.

Place: Holiday Inn Select, 8120 Wisconsin Avenue, Bethesda, MD 20892.