Extensions to the period of performance may be requested.

Approved Proposals

NCHS/NCEH will provide a data file with the requested recoded variables and a randomly assigned unique identification number that is linked to the DNA specimen. No record connecting the new number with the original identification number will be kept after the samples have been sent. These samples can not be traced to any files maintained by NCHS.

Agency Agreement

A formal signed agreement in the form of a Materials Transfer Agreement (MTA) with individuals who have projects approved will be completed before the release of the samples. This agreement will contain the conditions for use of the DNA as stated in this document and as agreed upon by the investigators and CDC. A key component of this agreement is that no attempt will be made to link the results of the proposed research to any other data, including, but not limited to, the NHANES III public use data set. Also, the investigator agrees that the samples can not be used for commercial purposes.

Progress Reports

A progress report will be submitted annually. NHANES IRB continuation reports are also required annually.

Disposition of Results and Samples

No DNA samples provided can be used for any purpose other than those specifically requested in the proposal and approved by the Genetic Technical Panel and the NHANES IRB. No sample can be shared with others, including other investigators, unless specified in the proposal and so approved. Any unused samples must be returned to the Bank upon completion of the approved project. Researchers requesting DNA samples for age-race-gender studies and special studies will be required to provide NCHS with the results of all DNA tests performed for each anonymized sample. These results, once returned to NCHS, will be part of the public domain. Therefore, ample time will be given to the investigator to publish results prior to reporting the results to NCHS.

Proposed Cost Schedule For Providing NHANES III DNA Specimen Bank

A nominal processing fee of $38.00 is proposed for each sample received from the NHANES III DNA Specimen Bank. The costs are determined both for NCEH and NCHS and include the physical materials needed to process the samples at the NCEH laboratory as well as the materials to process the requests for samples at NCHS. These costs are inclusive of the staff needed for these activities at each Center. The fee is estimated to cover the costs of processing, handling and preparing the samples in accordance with the detailed requirements of the investigators. These costs were based on an assumption that NCEH and NCHS will receive and process 15 proposals in a year each requesting 1000 samples as shown in the table below.

The materials listed are for the recurring laboratory costs to dispense and prepare the samples for shipping; the computer software needed for the Web page and advertisements in scientific journals. Labor costs are based on the need for microbiologists, a proposal administrator and computer programmers for NCHS and NCEH to maintain the data bases and verify anonymity. Technical panel travel and expenses are based on the panel meeting twice a year. The space estimate is based on acquiring storage and aliquoting space.

Total Costs Per Sample if 15 Requests for 1,000 Samples

Materials	$1.90
Labor	22.00
Panel Travel/Expenses	2.69
Space	0.97
Subtotal	27.56
NCHS overhead (15%)	4.12
Subtotal	31.68
CDC/FMO overhead (20%)	6.32
Total	*38.00

*Shipping costs are not included in the $38.00 processing fee. These costs must also be paid by the investigator.

Comments are solicited on the proposed cost schedule. Comments are due by June 30, 1999.

Send Comments and for Information

Audrey L. Burwell, MS, Health Research Administrator, National Center for Health Statistics, Centers for Disease Control and Prevention, 6525 Belcrest Road, Room 1100, Hyattsville, MD 20782, Phone: 301–436–7062, 127, FAX: 301–436–4233, E-Mail: azb2@cdc.gov

References

1. Plan and Operation of the Third National Health and Nutrition Examination Survey, 1988–94. National Center for Health Statistics. Vital Health Stat (32) 1994.

2. Clayton EW, Steinberg KK, Khoury MJ, et al. Informed consent for genetic research on stored tissue samples. JAMA 1995;274:1786–1792.

Dated: May 25, 1999.

Joseph R. Carter,

Acting Associate Director for Management And Operation, Centers for Disease Control and Prevention.

[FR Doc. 99–13740 Filed 5–28–99; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Oncologic Drugs Advisory Committee; Amendment of Notice

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing an amendment to the notice of meeting of the Oncologic Drugs Advisory Committee. This meeting was announced in the **Federal Register** of May 20, 1999 (64 FR 27581). The meeting will be open to the public. This amendment is being made to change the meeting starting time and the procedures paragraph and to cancel the closed session.

FOR FURTHER INFORMATION CONTACT: Karen M. Templeton-Somers, Center for Drug Evaluation and Research (HFD–21), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–7001, or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area), code 12542. Please call the Information Line for up-to-date information on this meeting.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of May 20, 1999 (64 FR 27581), FDA announced that a meeting of Oncologic Drugs Advisory Committee would be held on June 7 and 8, 1999. On page 27581, in the third column, the ''Date and Time'' and ''Procedure'' portions are amended and on page 27582, in the first column, the ''Closed Committee Deliberations'' portion is removed to read as follows:

Date and Time: The meeting will be held on June 7, 1999, 9:30 a.m. to 5:30 p.m. and June 8, 1999, 8 a.m. to 5:30 p.m.

Procedure: Interested persons may present data, information, or views, orally or in writing, on issues pending before the committee. Written submissions may be made to the contact person by May 28, 1999. Oral presentations from the public will be scheduled between approximately 9:45 a.m. and 10:15 a.m. and 1:45 p.m. and 2 p.m. on June 7, 1999, and between approximately 8:15 a.m. and 8:45 a.m.

on June 8, 1999. Time allotted for each presentation may be limited. Those desiring to make formal oral presentations should notify the contact person before May 28, 1999, and submit a brief statement of the general nature of the evidence or arguments they wish to present, the names and addresses of proposed participants, and an indication of the approximate time requested to make their presentation. After the scientific presentations, a 15-minute open public session will be conducted for interested persons who have submitted their request to speak by May 28, 1999, to address issues specific to the submission or topic before the committee.

Dated: May 26, 1999.

Michael A. Friedman,

Deputy Commissioner for Operations.

[FR Doc. 99–13963 Filed 5–27–99; 4:50 pm]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Advisory Council; Meeting

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92–463), announcement is made of the following National Advisory body scheduled to meet during the month of June 1999.

Name: National Advisory Council (NAC) on the National Health Service Corps (NHSC).

Date and Time: Thursday, June 10; 7:30 a.m.–9 p.m., Friday, June 11; 8:30 a.m.–4:30 p.m., Saturday, June 12; 9 a.m.–5 p.m., Sunday, June 13; 8:30 a.m.–10 a.m.

Place: Oklahoma City Marriott, 3233 North West Expressway, Oklahoma City, OK 73112, (405) 842–6633.

The meeting is open to the public.

Agenda: Items will include updates on the NHSC and Scholarships and Loan Repayments program; HPSA designations; reports from the Dallas field office, State Primary Care Association, and State Department of Health. The NAC will also be discussing their draft position paper, ''The National Health Service Corps for the 21st Century'' in preparation for the year 2000 reauthorization. Site visits will be on Friday, June 11. Transportation for the public will not be available.

For further information, call Ms. Eve Morrow at (301) 594–4144.

Dated: May 26, 1999.

Jane Harrison,

Director, Division of Policy Review and Coordination.

[FR Doc. 99–13791 Filed 5–28–99; 8:45 am]

BILLING CODE 4160–15–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Submission for OMB Review; Comment Request; Evaluation of National Youth Anti-Drug Media Campaign

SUMMARY: Under the provision of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the National Institute on Drug Abuse of the National Institutes of Health submitted to the Office of Management and Budget (OMB) a request to review and approve the information collected listed below. This proposed information collection was previously published in the **Federal Register** on November 30, 1998 on page 65795 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 after October 1, 1995, unless it displays a currently valid OMB control number.

PROPOSED COLLECTION: *Title:* Evaluation of National Youth Anti-Drug Media Campaign. *Type of Information Collection Request:* New. *Need and Use of Information Collection:* The White House Office of National Drug Control Policy has transferred funds to NIDA to conduct an independent, scientifically designed and implemented evaluation of the National Youth Anti-Drug Media Campaign, the first prevention campaign to use paid advertising to discourage youth from drug use. The study will assess the outcomes and impact of the national campaign in reducing illegal drug use among children and adolescents.

For this study, two different surveys will be conducted: (1) The National Survey of Parents and Youth, a cross-sectional household survey; and (2) a Community Longitudinal Study of Parents and Youth in four communities with an ethnographic component. All data will be collected using a combination of computer-assisted personal interviews (CAPI) and audio computer-assisted self-interviews (ACASI). The findings will form the basis of semiannual and annual reports on campaign progress. These reports will provide assistance in improving the national campaign, and will help to establish a rich database of information about the process involved in changing attitudes and behaviors by the mass media.

Frequency of Response: The National Survey of Parents and Youth will be carried out in 6 waves over a three-year period. Each data collection wave will last 6 months. The Community Longitudinal Study will be carried out annually over three years. *Affected Public:* Individuals and households. *Type of Respondent:* Children and parents. The annual reporting burden is as follows:

TABLE 1.—RESPONDENT AND BURDEN ESTIMATE

Type of respondents	Estimated number of respondents	Number of responses per respondent	Average time in hours per response	Estimated total burden hours	Estimated annual hour burden (over 3 years)
National Survey of Youth and Parents (NSPY)					
Screener Respondent	164,754	1	.07	11,533	3,844
Youth 9–11	9,300	1	.58	5,394	1,798
Adolescents 12–18	19,200	1	.75	14,400	4,800
Parents	20,100	1	.92	18,492	6,164
Community Longitudinal Study of Parents and Youth (CLSPY)					
Screener Respondent	28,500	1	.07	1,995	[1] N/A
Youth 9–11	2,150	3	.65	4,193	1,398
Adolescents 12–14	2,150	3	.83	5,354	1,785