The HHS Appropriations Act requires that when issuing statements, press releases, requests for proposals, bid solicitations, and other documents describing projects or programs funded in whole or in part with Federal money, grantees shall clearly state the percentage and dollar amount of the total costs of the program or project which will be financed with Federal money, and the percentage and dollar amount of the total costs of the project or program that will be financed by non-governmental sources.

3. *Reporting:* Each year of the project period, the cooperative agreement recipient is required to submit a non-competing application which includes an annual progress report for the current year, and a project work plan, budget, and budget justification for the upcoming year. The progress report should contain, at a minimum, a report on both process and outcome objectives, including evaluation of the activities as a whole, as well as for each component of NTC activities.

For component requirements, the progress report should contain, at a minimum, information related to the following:

Component A., *Compile, Coordinate, and Disseminate Training Information:* Obtaining and incorporating stakeholder input; gathering and cataloging training materials; design, implementation, and maintenance of NTC Web site; mechanisms for making information available; and, use of electronic technologies in addressing the requirements of this component.

Component B., *Conduct Training Meetings:* (a) Title of training event; (b) location; (c) topic(s) covered; (d) presenter(s) (as applicable); (e) number of participants; (f) agencies sponsoring participants; and (g) evaluation summary (including whether meeting objectives were met); and, (h) credit hours or CEUs available.

Component C., *Develop Training Resources and/or Materials:* The progress report should contain process and outcome information related to any materials developed as a result of this cooperative agreement.

All information should be provided in adequate detail for the reviewer to assess the planning, implementation, evaluation, and status of project activities compared to the approved work plan. If deviations from the approved work plan were necessary, the progress report should fully explain and justify modifications.

The work plan for the upcoming year should clearly reflect proposed NTC activities, including timeline; justification for any modifications from the previous year; S.M.A.R.T. objectives; and evaluation plan. The budget should reflect proposed costs to carry out the project plan. Sufficient detail should be provided so that the reviewer is able to determine the adequacy and appropriateness of budgeted items related to the proposed activities.

The cooperative agreement recipient is required to submit an annual Financial Status Report (FSR) within 90 days after the end of each budget period. Agencies that receive a total of $500,000 or greater of Federal funds must undergo an independent audit in accordance with OMB Circular A–133.

VII. Agency Contacts

Administrative and Budgetary Requirements

For application kits, submission of hard copy applications, and information on budget and business aspects of the application, please contact: WilDon Solutions, Office of Grants Management Operations Center, 1515 Wilson Blvd., Third Floor Suite 310, Arlington, VA 22209 at 1–888–203–6161, e-mail *OPHSgrantinfo@teamwildon.com*, or fax 703–351–1138.

Program Requirements

For information related to family planning program requirements, contact: WilDon Solutions, Office of Grants Management Operations Center, 1515 Wilson Blvd., Third Floor Suite 310, Arlington, VA 22209 at 1–888–203–6161, e-mail *OPHSgrantinfo@teamwildon.com*, or fax 703–351–1138. Identify that your inquiry is related to the Office of Population Affairs/Office of Family Planning program announcement for the National Family Planning Center Cooperative Agreement.

VIII. Other Information

Definitions: For the purposes of this announcement, the following definitions apply:

Family planning training—job-specific skill development, the purpose of which is to promote and improve the delivery of family planning services. Further description of family planning services may be found in the authorizing legislation, implementing regulations, and program guidelines.

Application—a request for financial support of a project submitted to OPA on specified forms and in accordance with instructions provided.

Cooperative Agreement—An award instrument of financial assistance where ''substantial involvement'' is anticipated between the HHS awarding agency and the recipient during performance of the contemplated project or activity. ''Substantial involvement'' means that the recipient can expect Federal programmatic collaboration or participation in managing the award. The entity that receives a Federal cooperative agreement assumes the legal and financial responsibility and accountability for the awarded funds and performance of activities approved for funding, and is held to all requirements for Federal grants.

Evidence-based—relevant scientific evidence that has undergone comprehensive review and rigorous analysis.

Project—those activities described in the application and supported under the approved budget.

Technical Assistance Conference Call: There will be an opportunity for prospective applicants to participate in a technical assistance conference call to be held within one month after publication of this Notice in the **Federal Register**. For more information regarding this opportunity, including date, registration information, and how to join the call, please consult the OPA Web site at *http://opa.osophs.dhhs.gov*.

Dated: May 1, 2007.

Evelyn M. Kappeler,

Acting Director, Office of Population Affairs.

[FR Doc. E7–8668 Filed 5–4–07; 8:45 am]

BILLING CODE 4150–34–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Agency for Healthcare Research and Quality

National Advisory Council for Healthcare Research and Quality: Request for Nominations for Public Members

AGENCY: Agency for Healthcare Research and Quality (AHRQ), HHS.

ACTION: Request for nominations for public members.

SUMMARY: 42 U.S.C. 299c, section 931 of the Public Health Service (PHS Act), established a National Advisory Council for Healthcare Research and Quality (the Council). The Council is to advise the Secretary of HHS and the Director of the Agency for Healthcare Research and Quality (AHRQ) on matters related to actions of the Agency to improve the quality, safety, efficiency, and effectiveness of health care for all Americans.

Seven current members' terms will expire in November 2007. To fill these positions in accordance with the legislative mandate establishing the

Council, we are seeking individuals who are distinguished: (1) In the conduct of research, demonstration projects, and evaluations with respect to health care; (2) in the fields of health care quality research or health care improvement; (3) in the practice of medicine; (4) in other health professions; (5) in representing the private health care sector (including health plans, providers, purchasers) or administrators of health care delivery systems; (6) in the fields of health care economics, information systems, law, ethics, business, or public policy; and, (7) in representing the interests of patients and consumers of health care. Individuals are particularly sought with experience and success in activities specified in the summary above.

DATES: Nominations should be received on or before June 15, 2007.

ADDRESSES: Nominations should be sent to Ms. Deborah Queenan, AHRQ, 540 Gaither Road, Room 3238, Rockville, Maryland 20850. Nominations also may be faxed to (301) 427–1341.

FOR FURTHER INFORMATION CONTACT: Ms. Deborah Queenan, AHRQ, at (301) 427–1330.

SUPPLEMENTARY INFORMATION: 42 U.S.C. 299c, section 931, of the PHS Act, provides that the National Advisory Council for Healthcare Research and Quality shall consist of 21 appropriately qualified representatives of the public appointed by the Secretary of Health and Human Services and, in addition, ex officio representatives from other Federal agencies specified in the authorizing legislation, principally agencies that conduct or support health care research, as well as Federal officials the Secretary may consider appropriate. The Council meets in the Washington, DC., metropolitan area, generally in Rockville, Maryland, approximately three times a year to provide broad guidance to the Secretary and AHRQ's Director on the direction of and programs undertaken by AHRQ.

Seven individuals will be selected presently by the Secretary to serve on the Council beginning with the meeting in the spring of 2008. Members generally serve 3-year terms. Appointments are staggered to permit an orderly rotation of membership.

Interested persons may nominate one or more qualified persons for membership on the Council. Self-nominations are accepted. Nominations shall include: (1) A copy of the nominee's resume or curriculum vitae; and (2) a statement that the nominee is willing to serve as a member of the Council. Selected candidates will be asked to provide detailed information concerning their financial interests, consultant positions and research grants and contracts, to permit evaluation of possible sources of conflict of interest.

The Department seeks a broad geographic representation and has special interest in assuring that women, minority groups, and the physically handicapped are adequately represented on advisory bodies, and therefore, particularly encourages nominations for appropriately qualified female, minority, and/or physically handicapped candidates.

Dated: April 26, 2007.

Carolyn M. Clancy,

Director.

[FR Doc. 07–2239 Filed 5–4–07; 8:45 am]

BILLING CODE 4160–90–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Agency for Healthcare Research and Quality Notice of Meeting

In accordance with section 10(d) of the Federal Advisory Committee Act (5 U.S.C., Appendix 2), announcement is made of a Health Care Policy and Research Special Emphasis Panel (SEP) meeting.

A Special Emphasis Panel is group of experts in fields related to health care research who are invited by the Agency for Healthcare Research and Quality (AHRQ), and agree to be available, to conduct on an as needed basis, scientific reviews of applications for AHRQ support. Individual members of the Panel do not attend regularly-scheduled meetings and do not serve for fixed terms or a long period of time. Rather, they are asked to participate in particular review meetings which require their type of expertise.

Substantial segments of the upcoming SEP meeting listed below will be closed to the public in accordance with the Federal Advisory Committee Act, section 10(d) of 5 U.S.C., Appendix 2 and 5 U.S.C. 552b(c)(6). Grant applications for ''Ambulatory Care Patient Safety Proactive Risk Assessment (P20),'' are to be reviewed and discussed at this meeting. These discussions are likely to reveal personal information concerning individuals associated with the applications. This information is exempt from mandatory disclosure under the above-cited statutes.

SEP Meeting on: Ambulatory Care Patient Safety Proactive Risk Assessment (P20).

Date: May 21–22, 2007 (Open on May 21 from 8 a.m. to 8:15 a.m. and closed for the remainder of the meeting).

Place: John M. Eisenberg Building, AHRQ Conference Center, 540 Gaither Road, Rockville, Maryland 20850.

Contact Person: Anyone wishing to obtain a roster of members, agenda or minutes of the non-confidential portions of this meeting should contact Mrs. Bonnie Campbell, Committee Management Officer, Office of Extramural Research, Education and Priority Populations, AHRQ, 540 Gaither Road, Room 2038, Rockville, Maryland 20850, Telephone (301) 427–1554.

Agenda items for this meeting are subject to change as priorities dictate.

Dated: April 23, 2007.

Carolyn M. Clancy,

Director.

[FR Doc. 07–2238 Filed 5–2–07; 8:45 am]

BILLING CODE 4160–90–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Pediatric Oncology Subcommittee of the Oncologic Drugs Advisory Committee; Notice of Meeting

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

This notice announces a forthcoming meeting of a public advisory committee of the Food and Drug Administration (FDA). The meeting is open to the public.

Name of Committee: Pediatric Oncology Subcommittee of the Oncologic Drugs Advisory Committee.

General Function of the Committee: To provide advice and recommendations to the agency on FDA's regulatory issues.

Date and Time: The meeting will be held on June 27, 2007, from 8 a.m. to 4 p.m.

Location: Food and Drug Administration, Center for Drug Evaluation and Research Advisory Committee Conference Room, rm. 1066, 5630 Fishers Lane, Rockville, MD.

Contact Person: Mimi Phan, Center for Drug Evaluation and Research (HFD–21), Food and Drug Administration, 5600 Fishers Lane, (for express delivery, 5630 Fishers Lane, rm. 1093), Rockville, MD 20857, 301–827–7001, FAX: 301–827–6776, e-mail: *Mimi.Phan@fda.hhs.gov*, or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area), code 3014512542. Please call the Information Line for up-to-date information on this meeting.

Agenda: The subcommittee will do the following: (1) Discuss review of