seeking nominations of qualified candidates to be considered for appointment as members of the Secretary's Advisory Committee on Human Research Protections (SACHRP). SACHRP provides advice and recommendations to the Secretary, HHS (Secretary), through the Assistant Secretary for Health, on matters pertaining to the continuance and improvement of functions within the authority of HHS directed toward protections for human subjects in research. SACHRP was established by the Secretary on October 1, 2002. OHRP is seeking nominations of qualified candidates to fill four positions on the Committee membership that will be vacated during the 2018 calendar year. **DATES:** Nominations for membership on

DATES: Nominations for membership on the Committee must be received no later than September 18, 2017.

ADDRESSES: Nominations should be mailed or delivered to Dr. Jerry Menikoff, Director, Office for Human Research Protections, Department of Health and Human Services, 1101 Wootton Parkway, Suite 200, Rockville, MD 20852. Nominations will not be accepted by email or by facsimile.

FOR FURTHER INFORMATION CONTACT: Julia Gorey, Executive Director, SACHRP, Office for Human Research Protections, 1101 Wootton Parkway, Suite 200, Rockville, MD 20852, telephone: 240–453–8141. A copy of the Committee charter and list of the current members can be obtained by contacting Ms. Gorey, accessing the SACHRP Web site at www.hhs.gov/ohrp/sachrp, or requesting via email at sachrp@osophs.dhhs.gov.

SUPPLEMENTARY INFORMATION: The Committee provides advice on matters pertaining to the continuance and improvement of functions within the authority of HHS directed toward protections for human subjects in research. Specifically, the Committee provides advice relating to the responsible conduct of research involving human subjects with particular emphasis on special populations such as neonates and children, prisoners, the decisionally impaired, pregnant women, embryos and fetuses, individuals and populations in international studies, investigator conflicts of interest and populations in which there are individually identifiable samples, data or information.

In addition, the Committee is responsible for reviewing selected ongoing work and planned activities of the OHRP and other offices/agencies within HHS responsible for human subjects protection. These evaluations

may include, but are not limited to, a review of assurance systems, the application of minimal research risk standards, the granting of waivers, education programs sponsored by OHRP, and the ongoing monitoring and oversight of institutional review boards and the institutions that sponsor research.

Nominations: The OHRP is requesting nominations to fill four positions for voting members of SACHRP. One position will become vacant in January 2018, with three others becoming vacant in October 2018. Nominations of potential candidates for consideration are being sought from a wide array of fields, including, but not limited to: public health and medicine, behavioral and social sciences, health administration, and biomedical ethics. To qualify for consideration of appointment to the Committee, an individual must possess demonstrated experience and expertise in any of the several disciplines and fields pertinent to human subjects protection and/or clinical research.

The individuals selected for appointment to the Committee can be invited to serve a term of up to four years. Committee members receive a stipend and reimbursement for per diem and any travel expenses incurred for attending Committee meetings and/or conducting other business in the interest of the Committee. Interested applicants may self-nominate.

Nominations should be typewritten. The following information should be included in the package of material submitted for each individual being nominated for consideration: (1) A letter of nomination that clearly states the name and affiliation of the nominee, the basis for the nomination (i.e., specific attributes which qualify the nominee for service in this capacity), and a statement that the nominee is willing to serve as a member of the Committee; (2) the nominator's name, address, daytime telephone number, and the home and/ or work address, telephone number, and email address of the individual being nominated; and (3) a current copy of the nominee's curriculum vitae. Federal employees should not be nominated for consideration of appointment to this Committee.

The Department makes every effort to ensure that the membership of HHS Federal advisory committees is fairly balanced in terms of points of view represented and the committee's function. Every effort is made to ensure that individuals from a broad representation of geographic areas, women and men, ethnic and minority groups, and the disabled are given

consideration for membership on HHS Federal advisory committees. Appointment to this Committee shall be made without discrimination on the basis of age, race, ethnicity, gender, sexual orientation, disability, and cultural, religious, or socioeconomic status.

Individuals who are selected to be considered for appointment will be required to provide detailed information regarding their financial holdings, consultancies, and research grants or contracts. Disclosure of this information is necessary in order to determine if the selected candidate is involved in any activity that may pose a potential conflict with the official duties to be performed as a member of SACHRP.

Authority: 42 U.S.C. 217a, Section 222 of the Public Health Service Act, as amended. The Committee is governed by the provisions of Public Law 92–463, as amended (5 U.S.C. Appendix 2), which sets forth standards for the formation and use of advisory committees.

Dated: July 26, 2017.

Julia Gorey,

Executive Director, Secretary's Advisory Committee on Human Research Protections, Office for Human Research Protections

[FR Doc. 2017-16156 Filed 8-1-17; 8:45 am]

BILLING CODE 4150-36-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute of Diabetes and Digestive and Kidney Diseases; Notice of Meetings

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. App.), notice is hereby given of meetings of the National Diabetes and Digestive and Kidney Diseases Advisory Council.

The meetings will be open to the public as indicated below, with attendance limited to space available. Individuals who plan to attend and need special assistance, such as sign language interpretation or other reasonable accommodations, should notify the Contact Person listed below in advance of the meeting.

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: National Diabetes and Digestive and Kidney Diseases Advisory Council.

Date: September 6, 2017. Open: 8:30 a.m. to 12:00 p.m.

Agenda: To present the Director's Report and other scientific presentations.

Place: National Institutes of Health, Natcher Building Forty-five, Conference Rooms E1/E2, 45 Center Drive, Bethesda, MD 20892.

Closed: 3:45 p.m. to 4:30 p.m. Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Natcher Building Forty-five, Conference Rooms E1/E2, 45 Center Drive, Bethesda, MD 20892

Contact Person: Brent B. Stanfield, Ph.D., Director, Division of Extramural Activities, National Institutes of Diabetes and Digestive and Kidney Diseases, 6707 Democracy Blvd. Room 7323, MSC 5452, Bethesda, MD 20892, (301) 594–8843, stanfibr@niddk.nih.gov.

Name of Committee: National Diabetes and Digestive and Kidney Diseases Advisory Council; Diabetes, Endocrinology and Metabolic Diseases.

Date: September 6, 2017. Closed: 1:00 p.m. to 2:00 p.m. Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Natcher Building Forty-five, Natcher Conference Center, Room E1, 45 Center Drive, Bethesda, MD 20892.

Open: 2:00 p.m. to 3:30 p.m.

Agenda: To review the Division's scientific and planning activities.

Place: National Institutes of Health, Natcher Building Forty-five, Natcher Conference Center, Room E1, 45 Center Drive, Bethesda, MD 20892.

Contact Person: Brent B. Stanfield, Ph.D., Director, Division of Extramural Activities, National Institutes of Diabetes and Digestive and Kidney Diseases, 6707 Democracy Blvd. Room 7323, MSC 5452, Bethesda, MD 20892, (301) 594–8843, stanfibr@niddk.nih.gov.

Name of Committee: National Diabetes and Digestive and Kidney Diseases Advisory Council; Kidney, Urologic and Hematologic Diseases.

Date: September 6, 2017. Open: 1:00 p.m. to 3:00 p.m.

Agenda: To review the Division's scientific and planning activities.

Place: National Institutes of Health, Natcher Building Forty-five, Natcher Conference Center, Room F2, 45 Center Drive, Bethesda, MD 20892.

Closed: 3:00 p.m. to 3:30 p.m. Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Natcher Building Forty-five, Natcher Conference Center, Room F2, 45 Center Drive, Bethesda, MD 20892.

Contact Person: Brent B. Stanfield, Ph.D., Director, Division of Extramural Activities, National Institutes of Diabetes and Digestive and Kidney Diseases, 6707 Democracy Blvd. Room 7323, MSC 5452, Bethesda, MD 20892, (301) 594–8843, stanfibr@niddk.nih.gov.

Name of Committee: National Diabetes and Digestive and Kidney Diseases Advisory Council; Digestive Diseases and Nutrition.

Date: September 6, 2017.

Open: 1:00 p.m. to 2:00 p.m.

Agenda: To review the Division's scientific and planning activities.

Place: National Institutes of Health, Natcher Building Forty-five, Natcher Conference Center, Room F1, 45 Center Drive, Bethesda, MD 20892.

Closed: 2:15 p.m. to 3:30 p.m. Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Natcher Building Forty-five, Natcher Conference Center, Room F1, 45 Center Drive, Bethesda, MD 20892.

Contact Person: Brent B. Stanfield, Ph.D., Director, Division of Extramural Activities, National Institutes of Diabetes and Digestive and Kidney Diseases, 6707 Democracy Blvd. Room 7323, MSC 5452, Bethesda, MD 20892, (301) 594–8843, stanfibr@niddk.nih.gov.

Any interested person may file written comments with the committee by forwarding the statement to the Contact Person listed on this notice. The statement should include the name, address, telephone number and when applicable, the business or professional affiliation of the interested person.

In the interest of security, NIH has instituted stringent procedures for entrance onto the NIH campus. All visitor vehicles, including taxicabs, hotel, and airport shuttles will be inspected before being allowed on campus. Visitors will be asked to show one form of identification (for example, a government-issued photo ID, driver's license, or passport) and to state the purpose of their visit.

Information is also available on the Institute's/Center's home page: www.niddk.nih.gov/fund/divisions/DEA/Council/coundesc.htm., where an agenda and any additional information for the meeting will be posted when available.

(Catalogue of Federal Domestic Assistance Program Nos. 93.847, Diabetes, Endocrinology and Metabolic Research; 93.848, Digestive Diseases and Nutrition Research; 93.849, Kidney Diseases, Urology and Hematology Research, National Institutes of Health, HHS)

Dated: July 27, 2017.

David Clary,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2017–16192 Filed 8–1–17; 8:45 am]

BILLING CODE 4140-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Request for Letters of Interest for NCI– MATCH Laboratories

SUMMARY: The National Cancer Institute (NCI) in collaboration with the NCI

Molecular Analysis for Therapy Choice (MATCH) trial leadership (NCT 02465060) invites applications for Clinical Laboratory Improvements Program (CLIA) certified/accredited laboratories that test tumor specimens from patients utilizing Next Generation Sequencing (NGS) assays to participate in the NCI MATCH trial. The NCI MATCH trial has implemented a new process for identifying patients for arms with rare variant eligibility criteria. Laboratories will contact any of the approximately 1100 sites that have activated NCI MATCH if a specimen sent from one of these sites has a rare variant that would potentially make the patient eligible for one of the treatment arms open in this initiative.

DATES: LOIs should be submitted to the National Cancer Institute (NCI), National Institutes of Health (NIH) on or before 5:00 p.m. EST on January 31, 2018.

ADDRESSES: Submit LOIs by email to NCIMATCHLabApps@nih.gov. 9609 Medical Center Drive, 3 West, Room 526, MSC 9728, Rockville, MD 20892.

FOR FURTHER INFORMATION CONTACT:

Questions about this request for LOIs should be directed to NCIMATCHLabApps@nih.gov. James V. Tricoli tricolij@mail.nih.gov can also

provide further information.

SUPPLEMENTARY INFORMATION: NCI—MATCH aims to establish whether patients with tumor mutations, amplifications or translocations in one of the genetic pathways of interest are likely to derive clinical benefit (primary objective: Objective response; secondary objective: Progression-free survival of at least 6 months) if treated with agents targeting that specific pathway in a single-arm design (see current arms below).

Patients with histologically documented solid tumors, lymphomas and multiple myeloma whose disease has progressed following at least one line of standard systemic therapy or for whom no standard therapy exists are eligible if they meet the eligibility criteria for the trial. Further information about the NCI–MATCH trial may be found at http://ecog-acrin.org/trials/ncimatch-eay131.

The selected collaborating laboratories may only act (*i.e.*, refer patients) on any of the rare variant arms for which their assay reports actionable mutations of interest (aMOIs). The assay must also report all exclusionary variants for the arm unless these occur at a frequency of >1% in cancer patients.

CLIA accredited/certified laboratories located in the United States may be