Dated: May 20, 2019.

Holli Richmond,

Executive Director, President's Council on Sports, Fitness, and Nutrition, Office of the Assistant Secretary for Health, U.S. Department of Health and Human Services.

[FR Doc. 2019–11476 Filed 5–31–19; 8:45 am]

BILLING CODE 4150–36–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Proposed Collection; 60-Day Comment Request; The Clinical Trials Reporting Program (CTRP) Database (NCI)

AGENCY: National Institutes of Health, HHS.

ACTION: Notice.

SUMMARY: In compliance with the requirement of the Paperwork Reduction Act of 1995 to provide opportunity for public comment on proposed data collection projects, the National Cancer Institute (NCI) will publish periodic summaries of proposed projects to be submitted to the Office of Management and Budget (OMB) for review and approval.

DATES: Comments regarding this information collection are best assured of having their full effect if received within 60 days of the date of this publication.

FOR FURTHER INFORMATION CONTACT: To obtain a copy of the data collection plans and instruments, submit comments in writing, or request more information on the proposed project, contact: Gisele Sarosy, MD, Coordinating Center for Clinical Trials (CCCT), National Cancer Institute, 9609 Medical Center Drive, 6W134, Rockville, MD 20852 or call non-toll-free number 240–276–6172 or Email your request, including your address to: *gisele.sarosy@nih.gov*. Formal requests for additional plans and instruments must be requested in writing.

SUPPLEMENTARY INFORMATION: Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995 requires: Written comments and/or suggestions from the public and affected agencies are invited to address 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.

Proposed Collection Title: The Clinical Trials Reporting Program (CTRP) Database, 0925–0600, Expiration Date 08/31/2019—EXTENSION, National Cancer Institute (NCI), National Institutes of Health (NIH).

Need and Use of Information Collection: The Clinical Trials Reporting Program (CTRP) Database is an electronic resource that serves as a single, definitive source of information about all NCI-supported clinical research. This resource allows the NCI to consolidate reporting, aggregate information and reduce redundant submissions. Information is submitted by clinical research administrators as designees of clinical investigators who conduct NCI-supported clinical research. The designees can electronically access the CTRP website to complete the initial trial registration. Subsequent to registration, four amendments and four study subject accrual updates occur per trial annually.

OMB approval is requested for 3 years. There are no costs to respondents other than their time. The estimated annualized burden hours are 18,000.

ESTIMATED ANNUALIZED BURDEN HOURS

Form name	Type of respondents	Number of respondents	Number of responses per respondent	Average time per response (in hours)	Total annual burden hours
Initial Registration	Clinical Trials	3,000	1	1	3,000
Amendment.		1,500	4	1	6,000
Update.		1,500	4	1	6,000
Accrual Updates.		3,000	4	15/60	3,000
Total		9,000	27,000		18,000

Patricia M. Busche,

Project Clearance Liaison, National Cancer Institute, National Institutes of Health.

[FR Doc. 2019–11459 Filed 5–31–19; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Submission for OMB Review; 30-Day Comment Request; Generic Clearance for Application Information for Fellowships, Internships, Training Programs, and Specialty Positions, National Cancer Institute (NCI)

AGENCY: National Institutes of Health, HHS.

ACTION: Notice.

SUMMARY: In compliance with the Paperwork Reduction Act of 1995, 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.

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

ADDRESSES: 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, *OIRA_submission@omb.eop.gov* or by

fax to 202–395–6974, Attention: Desk Officer for NIH.

FOR FURTHER INFORMATION CONTACT: To request more information on the proposed project or to obtain a copy of the data collection plans and instruments, contact: Vivian Horovitch-Kelley, Program Analyst, Office of Management Policy and Compliance, National Cancer Institute, 9609 Medical Center Drive, Room 2W444, Bethesda, Maryland 20892 or call non-toll-free number (240) 276–6850 or Email your request, including your address to: *vivian.horovitch-kelley@mail.nih.gov*.

SUPPLEMENTARY INFORMATION: This proposed information collection was previously published in the **Federal Register** on March 29 (Vol. 84, No. 61, Page 11987) 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 Cancer Institute (NCI), 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.

In compliance with Section 3507(a)(1)(D) of the Paperwork Reduction Act of 1995, 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.

Proposed Collection: Generic Clearance for Application Information from Fellows, Interns, and Trainees, 0925–XXXX, Exp., Date XX/XXXX, NEW, National Cancer Institute (NCI), National Institutes of Health (NIH).

Need and Use of Information Collection: This is a new generic information collection request to support the science and research in a multidisciplinary environment at the National Cancer Institute (NCI), a part of the National Institutes of Health. Applicants may possess a variety of degrees including, but not limited to, high school, post-baccalaureate, graduate, postdoctoral, Registered Nurse, and Doctor of Medicine (MD). Potential applicants may apply for cancer-related positions by submitting applications, resumes, curriculum vitae (CV), reference letters, letters of intent and interest, and other related documentation directly to the Divisions, Offices, and Centers. This information is necessary to evaluate the eligibility, merits, and quality of potential candidates and will also assist in matching potential candidates to various training and internship programs, and specialty positions. The information is for internal use to make decisions about candidates invited to visit and attend NCI fellowships, internships, training opportunities, and apply for specialized staff and faculty positions.

OMB approval is requested for 3 years. There are no costs to respondents other than their time. The total estimated annualized burden is 15,000 hours.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Number of respondents	Number of responses per respondent	Average time per response (in hours)	Total annual burden hours
Individuals (Applicants)	6,000	1	60/60	6,000
Individuals (Professional References)	18,000	1	30/60	9,000
Totals	24,000	24,000	.....	15,000

Patricia M. Busche,

Project Clearance Liaison, National Cancer Institute, National Institutes of Health.

[FR Doc. 2019–11461 Filed 5–31–19; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

All of Us Research Program, Tribal Consultation Meetings and Listening Sessions

AGENCY: National Institutes of Health, HHS.

ACTION: Notice.

SUMMARY: The National Institutes of Health (NIH) *All of Us* Research Program will host four Tribal Consultations to consult on the best practices for engaging with Tribal Nations to facilitate the inclusion of American Indian and Alaska Native (AI/AN) populations in this program. They will take place at three U.S. Department of Health and Human Services (HHS) Regional Consultations and at the National Congress of American Indians (NCAI) Mid-Year Conference and Marketplace.

DATES:

June 18, 2019: HHS Regional Consultation, Region 9 (Sacramento, CA).

June 24, 2019: NCAI Mid-Year Conference & Marketplace (Sparks, NV).

August 21, 2019: HHS Regional Consultation, Regions 1–4 (Washington, DC).

August 21, 2019: HHS Regional Consultation, Regions 6–8 (Denver, CO).

ADDRESSES: A full schedule of consultations with specific dates and times, as well as full location information, will be made available at *https://AllofUs.nih.gov/All-Us-Tribal-Engagement*.

FOR FURTHER INFORMATION CONTACT: The *All of Us* Tribal Engagement team by phone at 240–515–5317, by email at *AOUTribal@nih.gov,* or by mail at 6011 Executive Boulevard, Suite 214, Rockville, MD 20852.

SUPPLEMENTARY INFORMATION: In accordance with the U.S. Department of Health and Human Services' (HHS) Tribal Consultation Policy, *All of Us* announces that they will be holding four Tribal consultations during summer 2019: at the HHS Regional Consultation in Sacramento, California on June 18; at the NCAI Mid-Year Conference & Marketplace in Sparks, Nevada on June 24; at the HHS Regional Consultation in Washington, DC on July 16; and at the HHS Regional Consultation in Denver, Colorado on August 21.

The *All of Us* Research Program aims to accelerate health research and medical breakthroughs to enable an era of precision medicine for all. *All of Us* is committed to ensuring the program reflects the diversity of the United States. The goal for these events is to facilitate information exchange and provide an opportunity for Tribal and Urban Indian Organization leadership to have meaningful input as *All of Us* identifies priorities and opportunities around the inclusion of AI/AN