

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
<b>Total .....</b>		<b>9,000</b>	<b>27,000</b>		<b>18,000</b>

**Patricia M. Busche,**  
*Project Clearance Liaison, National Cancer Institute, National Institutes of Health.*  
 [FR Doc. 2019-11459 Filed 5-31-19; 8:45 am]  
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**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