

clinical utility has been constrained, in part, by dose-limiting toxicity following systemic administration and the need for repeated dosing. The subject invention addresses these limitations through synthetic IL-15/21 sequences which incorporate flexible linker regions and cell membrane anchors. T cells engineered to express these constructs experience autocrine IL-15/21 signaling leading to enhanced anti-tumor function in vivo.

This Notice is made in accordance with 35 U.S.C. 209 and 37 CFR part 404. The prospective exclusive license will be royalty bearing, and the prospective exclusive license may be granted unless within fifteen (15) days from the date of this published Notice, the National Cancer Institute receives written evidence and argument establishing that the grant of the license would not be consistent with the requirements of 35 U.S.C. 209 and 37 CFR part 404.

In response to this Notice, the public may file comments or objections. Comments and objections, other than those in the form of a license application, will not be treated confidentially and may be made publicly available.

License applications submitted in response to this Notice will be presumed to contain business confidential information and any release of information from these license applications will be made only as required and upon a request under the Freedom of Information Act, 5 U.S.C. 552.

Dated: February 12, 2021.

Richard U. Rodriguez,
Associate Director, Technology Transfer
Center, National Cancer Institute.

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BILLING CODE 4140-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Submission for OMB Review; 30-Day Comment Request; CareerTrac

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 recommendations for the proposed information collection should be sent within 30 days of publication of this notice to www.reginfo.gov/public/do/PRAMain. Find this particular information collection by selecting "Currently under 30-day Review—Open for Public Comments" or by using the search function.

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: Dr. Kristi Pettibone, Health Scientist Administrator, Program Analysis Branch, Division of Extramural Research and Training, NIEHS, NIH, 560 Davis Dr., Morrisville, NC 27560, or call non-toll-free number (984) 287-3303 or Email your request, including your address to: pettibonekg@niehs.nih.gov.

SUPPLEMENTARY INFORMATION: This proposed information collection was previously published in the **Federal Register** on December 12, 2020, page 79493–79494 (64 FR 15367) 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 Fogarty International Center (FIC), National Cancer Institute (NCI), National Institute of Diabetes and Digestive and Kidney Diseases (NIDDK), and National Institute of Environmental Health Sciences (NIEHS), 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: CareerTrac-0925-0568—expiration date April 30, 2021, REVISION, Fogarty International Center (FIC), National Institute of Environmental Health Sciences (NIEHS), National Cancer Institute (NCI), National Institute of Diabetes and Digestive Kidney Diseases, (NIDDK), National Institutes of Health (NIH).

Need and Use of Information Collection: The purpose of this data collection system is to track, evaluate and report short and long-term outputs, outcomes and impacts of trainees involved in health research training programs—specifically tracking this for at least ten years following training by having Principal Investigators enter data after trainees have completed the program. The data collection system provides a streamlined, web-based application permitting principal investigators to record career achievement progress by trainee on a voluntary basis. FIC, NCI, NIDDK, and NIEHS management will use this data to monitor, evaluate and adjust grants to ensure desired outcomes are achieved, comply with OMB Part requirements, respond to congressional inquiries, and as a guide to inform future strategic and management decisions regarding the grant program.

OMB approval is requested for 3 years. There are no costs to respondents other than their time. The total estimated annualized burden hours are 12,705.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total annual burden hour
FIC Grantee	90	20	40/60	1,200
NIEHS Grantee	60	45	40/60	1,800
NCI CRCHD Grantee	244	22	40/60	3,579
NCI D43 Grantee	20	22	40/60	293
Superfund Grantee	30	105	40/60	2,100

ESTIMATED ANNUALIZED BURDEN HOURS—Continued

Type of respondent	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total annual burden hour
NIDDK Grantee	30	20	40/60	400
Trainees	5,000	1	40/60	3,333
Total	5,474	19,058	12,705

Jane M. Lambert,

Project Clearance Liaison, National Institute of Environmental Health Sciences, National Institutes of Health.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Proposed Collection; 60 Day Comment Request; The Impact of Clinical Research Training and Medical Education at the Clinical Center on Physician Careers in Academia and Clinical Research (Clinical Center)

AGENCY: National Institutes of Health, HHS.

ACTION: Notice.

SUMMARY: In compliance with the requirement of the Paperwork Reduction Act of 1995, for opportunity for public comment on proposed data collection projects, the Clinical Center, the National Institutes of Health (NIH) 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, contact: Robert M. Lembo, MD, Office of Clinical Research Training and Medical Education, NIH Clinical Center, National Institutes of Health, 10 Center Drive, Room 1N252C, Bethesda, MD 20892-1158, or call non-toll-free number (301) 496-2636, or Email your request, including your address to: robert.lembo@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 Impact of Clinical Research Training and

Medical Education at the Clinical Center on Physician Careers in Academia and Clinical Research, OMB #0925-0602 Expiration Date: 11/30/2022, REVISION, Clinical Center (CC), National Institutes of Health (NIH).

Need and Use of Information Collection: The information collected will allow continued assessment of the value of the training provided by the Office of Clinical Research Training and Medical Education (OCRTME) at the NIH Clinical Center and the extent to which this training promotes (a) patient safety; (b) research productivity and independence; and (c) future career development within clinical, translational, and academic research settings. The information received from respondents is presented to, evaluated by, and incorporated into the ongoing operational improvement efforts of the Director of the Office of Clinical Research Training and Education, and the Chief Executive Officer of the NIH Clinical Center. This information will enable the ongoing operational improvement efforts of the OCRTME and its commitment to providing clinical research training and medical education of the highest quality to each trainee.

OMB approval is requested for 3 years. There are no costs to respondents other than their time. The total estimated annualized burden hours 478.

ESTIMATED ANNUALIZED BURDEN HOURS

Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total annual burden hours requested
CRTP/MRSP Alumni Survey	704	1	20/60	235
Summer Internship Program Alumni Survey	280	1	20/60	93
Graduate Medical Education Graduate Survey	350	1	20/60	117
Clinical Electives Program 1 Year Alumni Surveys	100	1	20/60	33
Total	1,434	1,434	478