Application No. 63/180,534, filed April 27, 2021.

Licensing Contact: Peter Soukas, J.D., 301–496–2644; *peter.soukas@nih.gov.*

Collaborative Research Opportunity: The National Institute of Allergy and Infectious Diseases is seeking statements of capability or interest from parties interested in collaborative research to further develop, evaluate or commercialize for development of a vaccine for respiratory or other infections. For collaboration opportunities, please contact Peter Soukas, J.D., 301–496–2644; *peter.soukas@nih.gov.*

Dated: May 10, 2021.

Surekha Vathyam,

Deputy Director, Technology Transfer and Intellectual Property Office, National Institute of Allergy and Infectious Diseases.

[FR Doc. 2021–10818 Filed 5–21–21; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Proposed Collection; 60-Day Comment Request: Electronic Individual Development Plan (eIDP) (National Eye Institute)

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 Eye Institute of the National Institutes of Health will publish periodic summaries of propose 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: Dr. Cesar E. Perez-Gonzalez, Training Director, Office of the Scientific Director, National Eye Institute, NIH, Building 31, Room 6A22, MSC 0250, Bethesda, Maryland 20892 or call non-toll-free number (301) 451–6763 or Email your request, including your address to: *cesarp@nei.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: Electronic Individual Development Plans, 0925–NEW, expiration date XX/XX/XXXX, National Eye Institute (NEI), National Institutes of Health (NIH).

Need and Use of Information Collection: The National Eye Institute's (NEI) Office of the Scientific Director (OSD) goal is to train the next generation of vision researchers and ophthalmologists. Trainees who participate in NEI research come with different levels of education (student, postbaccalaureate, predoctoral including graduate and medical students, postdoctoral fellows) and for different amounts of time (6 months to 5 years). Training at the NEI focuses on scientific and professional skill development. To enhance their chances of obtaining their ideal career, completing an annual Individual Development Plan (IDP) is an important step in helping a trainee's career and professional development and is standard practice in graduate and postdoctoral education. An IDP is an effective tool for trainees to think about their career goals and skills needed to achieve them during their time at the NEI. Trainees work together with their research mentor to organize and summarize their research projects, consider career goals, and set training goals and expectations, both for the mentee and mentor.

This information collection request is to implement an electronic Individual Development Plan (eIDP). The data collected comes from a detailed questionnaire focused on responses to professional goals and expectations while the they are at the NEI. It is expected that the trainees will complete the eIDP annually and by doing so, it will help enhance the effectiveness of their training by setting clear goals that can be monitored not only by the trainee themselves but also by their mentor, the Training Director, and their Administrative Officer. In addition to this eIDP, the system will also implement an electronic exit survey. The data collected comes from a detailed questionnaire focused on responses to questions focused on trainee mentoring and professional experiences at the NEI as well as their plans after they depart. It is expected that the trainees will complete at the end of their tenure and that by doing so, the NEI Training Program can learn about ways to improve career development opportunities for future trainees as well as learn more about trainee job choices to better advise fellows. Additionally, we can use the survey to help determine mentor effectiveness and help identify problems in mentoring at the NEI.

OMB approval is requested for 3 years. There are no costs to respondents other than their time. The total estimated annualized burden hours are 450.

ESTIMATED ANNUALIZED BURDEN HOURS

	Type of respondent	Number of respondents	Number of responses per respondent	Average time per response (in hours)	Total annual burden hour
eIDP	Individuals	150	1	2	300
Exit Survey Part 1	Individuals	150	1	30/60	75
Exit Survey Part 2	Individuals	150	1	30/60	75

ESTIMATED ANNUALIZED BURDEN HOURS—Continued

	Type of respondent	Number of respondents	Number of responses per respondent	Average time per response (in hours)	Total annual burden hour
Total	150	150		450	

Dated: May 11, 2021.

Cesar E. Perez-Gonzalez,

Training Director, National Eye Institute, National Institutes of Health.

[FR Doc. 2021–10820 Filed 5–21–21; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Heart, Lung, and Blood Institute; Amended Notice of Meeting

Notice is hereby given of a change in the meeting of the National Heart, Lung, and Blood Advisory Council, June 08, 2021, 10:00 a.m. to June 08, 2021, 05:00 p.m., NIH, Rockledge 1, 6705 Rockledge Dr, Bethesda, MD 20892 which was published n the **Federal Register** on May 05, 2021, 294922.

The notice is amended to change the time of the meeting's public portion to 12:00 p.m. through 5:00 p.m. The meeting is partially closed to the public.

Dated: May 19, 2021.

David W. Freeman,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2021–10894 Filed 5–21–21; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Proposed Collection; 30 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 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 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: This proposed information collection was previously published in the **Federal Register** February 25, 2021 on pages 11550–11551 (86 FR 11550) 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 Clinical Center, 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: The Impact of Clinical Research Training and Medical Education at the Clinical Center on Physician Careers in Academia and Clinical Research, 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	Type of respondents	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total annual burden hours requested
CRTP/MRSP Alumni Survey	Post Doctoral Students	704	1	20/60	235