The information collection reflects an increase in 254,750 burden hours and 11,568 responses annually since the last OMB review and approval of the information collection. We attribute this to an increase in the number of submissions.

Dated: April 22, 2022.

Lauren K. Roth,

 $Associate\ Commissioner\ for\ Policy.$ [FR Doc. 2022–09070 Filed 4–27–22; 8:45 am]

BILLING CODE 4164-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Indian Health Service [OMB No. 0917–0041]

Request for Public Comment: 30-Day Information Collection: Indian Health Service Information Security Ticketing and Incident Reporting.

AGENCY: Indian Health Service, HHS. **ACTION:** Notice and request for comments. Request for extension of approval.

SUMMARY: In compliance with the Paperwork Reduction Act of 1995, the Indian Health Service (IHS) invites the general public to take this opportunity to comment on the information collection Office of Management and Budget (OMB) Control Number 0917–0041, titled, Information Security Ticketing and Incident Reporting. The purpose of this notice is to allow 30 days for public comment submitted

directly to OMB. A copy of the draft supporting statement is available at www.regulations.gov (see Docket ID IHS_FRDOC_001).

DATES: Comment Due Date: May 31, 2022. Your comments regarding this information collection are best assured of having full effect if received within 30 days of the date of this publication.

ADDRESSES: Direct Your Comments to OMB: Send your comments and suggestions regarding the proposed information collection contained in this notice, especially regarding the estimated public burden and associated response time to: Office of Management and Budget, Office of Regulatory Affairs, New Executive Office Building, Room 10235, Washington, DC 20503, Attention: Desk Officer for IHS.

FOR FURTHER INFORMATION CONTACT: To request additional information, please contact Evonne Bennett, Information Collection Clearance Officer at: *Evonne.Bennett@ihs.gov* or 301–443–4750.

SUPPLEMENTARY INFORMATION: This previously approved information collection project was last published in the Federal Register on February 17, 2022 (87 FR 9071), and allowed 60 days for public comment. No public comment was received in response to the notice. This notice announces our intent to submit this collection, which expires April 30, 2022, to OMB for approval of an extension, and to solicit comments on specific aspects for the proposed information collection.

Title: 0917–0041, "Information Security Ticketing and Incident Reporting."

Form(s) and Form number(s): Incident Reporting Form, Form F07–02b.

OMB Control Number: 0917-0041. Need and Use of Information Collection: This information collection activity provides a means for federal employees, Tribal employees, contractors, and other non-federal employees to report IHS information technology (IT) security and privacy incidents. This information collection has three purposes: To notify the CSIRT of an incident, provide updates about an open incident, and indicate resolution of an existing incident. The information collection furthers the IHS's ability to use secure IT, to enhance response time to IT incidents, and to maintain the agency's healthcare information security posture. This information collection also allows IHS to process privacy incidents and breaches within the IHS, in keeping with internal and external requirements.

Members of Affected Public: Federal employees, Tribal employees, contractors, and other non-federal employees accessing IHS IT systems.

Status of the Proposed Information Collection: Extension request.

Type of Respondents: Individuals. The table below provides: Types of data collection instruments, estimation of the number of respondents, number of responses per respondent, annual number of responses, average burden hour per response, and total annual burden hours.

Data collection instrument(s)	Estimated number of respondents	Responses per respondent	Annual number of responses	Average burden hour per response*	Total annual burden hours
IHS Federal and Non-Federal Staff	1700	1	1700	15/60	425
Total	1700	1	1700	15/60	425

^{*}For ease of understanding, the average burden per response is 15 minutes. There are no direct costs to respondents to report.

Requests for Comments: Your written comments and/or suggestions are invited on one or more of the following points:

- (a) whether the information collection activity is necessary to carry out an agency function;
- (b) whether the agency processes the information collected in a useful and timely fashion;
- (c) the accuracy of the public burden estimate (the estimated amount of time needed for individual respondents to provide the requested information);
- (d) whether the methodology and assumptions used to determine the estimates are logical;

- (e) ways to enhance the quality, utility, and clarity of the information being collected; and
- (f) ways to minimize the public burden through the use of automated, electronic, mechanical, or other technological collection techniques or other forms of information technology.

Elizabeth A. Fowler,

 $Acting\ Director, Indian\ Health\ Service.$ [FR Doc. 2022–09055 Filed 4–27–22; 8:45 am]

BILLING CODE 4165-16-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Government-Owned Inventions; Availability for Licensing

AGENCY: National Institutes of Health, HHS.

ACTION: Notice.

SUMMARY: The invention listed below is owned by an agency of the U.S. Government and is available for licensing to achieve expeditious commercialization of results of federally-funded research and

development. Foreign patent applications are filed on selected inventions to extend market coverage for companies and may also be available for licensing.

FOR FURTHER INFORMATION CONTACT:

Chris Kornak at 240–627–3705 or Chris.Kornak@nih.gov. Licensing information may be obtained by communicating with the Technology Transfer and Intellectual Property Office, National Institute of Allergy and Infectious Diseases, 5601 Fishers Lane, Rockville, MD 20852; tel. 301–496–2644. A signed Confidential Disclosure Agreement will be required to receive copies of unpublished information related to the invention.

SUPPLEMENTARY INFORMATION:

Technology description follows:

Replication-Competent Adenovirus Type-4 HIV Env Vaccines and Their

Description of Technology: National Institute of Allergy and Infectious Diseases (NIAID), International AIDS Vaccine Initiative (IAVI), Emergent, and Scripps have developed two recombinant adenovirus type 4 (Ad4) vector-based vaccine candidates. These replicating Ad4 vector-based candidates have shown improved activity against tier 2 HIV-1 isolates in experimental animals. Tier 2 isolates are among the most prevalent in infected populations. The two candidates, Ad4-Env150KN and Ad4-Env145NFL, incorporate novel design features based on Ad4-EnvC150 (1086c). Specifically, the truncation of the cytoplasmic tail of Env increases cell surface expression and has resulted in improved antigenicity from both candidates.

Additionally, the upper respiratory tract administration offers a way to bypass pre-existing Ad4 immunity in most people. Furthermore, unlike non-replicating vectors, these vaccines may evoke a durable immune response.

This technology is available for licensing for commercial development in accordance with 35 U.S.C. 209 and 37 CFR part 404, as well as for further development and evaluation under a research collaboration.

Potential Commercial Applications

• Prophylaxis against HIV-1.

Competitive Advantages

- Replicating vector may invoke durable immunity against HIV-1.
- Potential for prophylactic use in high-risk populations.
- Upper-respiratory (intranasal) administration will bypass pre-existing Ad4 immunity in most people.

Development Stage

• Phase 1 Clinical Trial (NCT03878121).

Inventors: Mark Connors (NIAID), Jeff Alexander (Emergent), Lo Vang (Emergent), Richard Wyatt (Scripps and IAVI), and Javier Guenaga (IAVI).

Publications: Alexander J., Mendy J., Vang L., Avanzini J.B., Garduno F., et al. (2013) Pre-Clinical Development of a Recombinant, Replication-Competent Adenovirus Serotype 4 Vector Vaccine Expressing HIV–1 Envelope 1086 Clade C. PLOS ONE 8(12): e82380. https://doi.org/10.1371/journal.pone.0082380.

Intellectual Property: HHS Reference No. E-105-2020-0-PCT-01—PCT Application No. PCT/US21/45389 filed on 10 August 2021.

Licensing Contact: To license this technology, please contact Chris Kornak at 240–627–3705 or Chris.Kornak@nih.gov, and reference E-105-2020.

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 this technology. In particular, NIAID would be very interested in a partnership with an entity that has a complementary HIV vaccine technology. For collaboration opportunities, please contact Chris Kornak at 240–627–3705 or Chris.Kornak@nih.gov.

Dated: April 25, 2022.

Surekha Vathyam,

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

[FR Doc. 2022–09158 Filed 4–27–22; 8:45 am]

BILLING CODE 4140-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Office of the Secretary; Notice of Meeting

Pursuant to section 10(a) of the Federal Advisory Committee Act, as amended, notice is hereby given of a meeting of the Interagency Pain Research Coordinating Committee.

The meeting will be open to the public via NIH Videocast https://videocast.nih.gov/. Individuals who plan to participate 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.

Name of Committee: Interagency Pain Research Coordinating Committee.

Date: June 7, 2022.

Time: 2:00 p.m. to 5:00 p.m. Eastern Daylight Time (EDT).

Agenda: The meeting will cover committee business items and IPRCC member updates. Items discussed will include updates on pain workforce enhancement, pain research, patient engagement, and diversity efforts.

Webcast Live: https://videocast.nih.gov/. Deadline: Submission of intent to submit written/electronic statement for comments: Tuesday, May 31st, by 5:00 p.m. EDT.

Place: National Institutes of Health, Building 31, 31 Center Drive, Bethesda, MD 20892 (Virtual Meeting).

Contact Person: Linda L. Porter, Ph.D., Director, Office of Pain Policy and Planning, Office of the Director, National Institute of Neurological Disorders and Stroke, NIH, 31 Center Drive, Room 8A31, Bethesda, MD 20892, Phone: (301) 451–4460, Email: Linda.Porter@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.

Visit the IPRCC website for more information: https://iprcc.nih.gov. Agenda and any additional information for the meeting will be posted when available.

Dated: April 25, 2022.

Tyeshia M. Roberson-Curtis,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2022–09140 Filed 4–27–22; $8:45~\mathrm{am}$]

BILLING CODE 4140-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Prospective Grant of Exclusive License, Inter-Institutional Agreement-Institution Lead: Engineered Influenza Neuraminidase Antigens

AGENCY: National Institutes of Health, HHS.

ACTION: Notice.

SUMMARY: The National Institute of Allergy and Infectious Diseases (NIAID), an institute of the National Institutes of Health (NIH), Department of Health and Human Services (HHS), is contemplating the grant of an exclusive, sublicensable patent license to the University of Washington, located in Seattle, State of Washington, U.S.A. in its rights to the inventions and the patent applications listed in the SUPPLEMENTARY INFORMATION section of this notice

DATES: Only written comments and/or applications for a license which are