Controlled substance	Drug code	Schedule
Phenazocine	9715	II
Thiafentanil	9729	II
Piminodine	9730	II
Racemethorphan	9732	II
Racemorphan	9733	II
Alfentanil	9737	II
Remifentanil	9739	II
Sufentanil	9740	II
Carfentanil	9743	II
Tapentadol	9780	II
Bezitramide	9800	II
Moramide-intermediate	9802	II

The company plans to import small quantities of the listed controlled substances to support research activities funded by the National Institute on Drug Abuse. No other activities for these drug codes are authorized for this registration.

Approval of permit applications will occur only when the registrant's business activity is consistent with what is authorized under 21 U.S.C. 952(a)(2). Authorization will not extend to the import of Food and Drug Administration-approved or non-approved finished dosage forms for commercial sale.

Matthew Strait,

Deputy Assistant Administrator.

[FR Doc. 2025–08414 Filed 5–13–25; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–1543]

Importer of Controlled Substances Application: Pall Life Sciences PR, LLC

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Pall Life Sciences PR, LLC has applied to be registered as an importer of basic class(es) of controlled substance(s). Refer to **SUPPLEMENTARY INFORMATION** listed below for further drug information.

DATES: Registered bulk manufacturers of the affected basic class(es), and applicants, therefore, may submit electronic comments on, or objections to the issuance of the proposed registration on or before June 13, 2025. Such persons may also file a written request for a hearing on the application on or before June 13, 2025.

ADDRESSES: The Drug Enforcement Administration requires that all comments be submitted electronically through the Federal eRulemaking Portal, which provides the ability to type short comments directly into the comment field on the webpage or attach a file for lengthier comments. Please go to *https://www.regulations.gov* and follow the online instructions at that site for submitting comments. Upon submission of your comment, you will receive a Comment Tracking Number. Please be aware that submitted comments are not instantaneously available for public view on *https://www.regulations.gov.* If you have received a Comment Tracking Number, your comment has been successfully submitted and there is no need to resubmit the same comment. All requests for a hearing must be sent to: (1) Drug Enforcement Administration, Attn: Hearing Clerk/OALJ, 8701 Morrissette Drive, Springfield, Virginia 22152; and (2) Drug Enforcement Administration, Attn: DEA Federal Register Representative/DPW, 8701 Morrissette Drive, Springfield, Virginia 22152. All requests for a hearing should also be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrissette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.34(a), this is notice that on February 25, 2025, Pall Life Sciences PR, LLC, Road 194, Kilometer 0.4, Fajardo, Puerto Rico 00738–0000, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Oxycodone	9143	II
Hydromorphone	9150	II
Methadone	9250	II
Morphine	9300	II
Fentanyl	9801	II

The company plans to import listed controlled substances for research purposes, drug testing, and analysis to support foreign regulatory compliance of finished dosage forms to foreign markets. No other activities for these drug codes are authorized for this registration.

Approval of permit applications will occur only when the registrant's business activity is consistent with what is authorized under 21 U.S.C. 952(a)(2). Authorization will not extend to the import of Food and Drug Administration-approved or non-approved finished dosage forms for commercial sale.

Matthew Strait,

Deputy Assistant Administrator.

[FR Doc. 2025–08413 Filed 5–13–25; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF LABOR

Agency Information Collection Activities; Submission for OMB Review; Comment Request; Pilot Study and Prospective Analysis of the Draft Revised Form 33, Safety and Health Program Assessment Worksheet

ACTION: Notice of availability; request for comments.

SUMMARY: The Department of Labor (DOL) is submitting this Occupational Safety & Health Administration (OSHA)-sponsored information collection request (ICR) to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act of 1995 (PRA). Public comments on the ICR are invited.

DATES: The OMB will consider all written comments that the agency receives on or before June 13, 2025.

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: Nicole Bouchet by telephone at 202–693–0213, or by email at *DOL_PRA_PUBLIC@dol.gov*.

SUPPLEMENTARY INFORMATION: OSHA conducts validity and reliability analyses of a safety and health program assessment worksheet, the Draft Revised Form 33. The studies will enable OSHA to ensure that a valid, reliable, and efficient tool is provided to On-Site Consultation programs in the fifty states, the District of Columbia, and several United States territories to replace the current OSHA Form 33, thereby, enhancing the quality of consultative services. The studies for which OSHA has requested approval comprises of a pre-test, a follow-up study, and a Prospective Analysis and OSHA is requesting additional time to complete the remaining studies. For additional substantive information about this ICR, see the related notice published in the **Federal Register** on January 13, 2025 (90 FR 2756).

Comments are invited on: (1) whether the collection of information is necessary for the proper performance of the functions of the Department, including whether the information will have practical utility; (2) the accuracy of the agency's estimates of the burden and cost of the collection of information, including the validity of the methodology and assumptions used; (3) ways to enhance the quality, utility and clarity of the information collection; and (4) ways to minimize the burden of the collection of information on those who are to respond, including the use of automated collection techniques or other forms of information technology.

This information collection is subject to the PRA. A Federal agency generally cannot conduct or sponsor a collection of information, and the public is generally not required to respond to an information collection, unless the OMB approves it and displays a currently valid OMB Control Number. In addition, notwithstanding any other provisions of law, no person shall generally be subject to penalty for failing to comply with a collection of information that does not display a valid OMB Control Number. *See* 5 CFR 1320.5(a) and 1320.6.

DOL seeks PRA authorization for this information collection for three (3) years. OMB authorization for an ICR cannot be for more than three (3) years without renewal. The DOL notes that information collection requirements submitted to the OMB for existing ICRs receive a month-to-month extension while they undergo review.

Agency: DOL–OSHA.

Title of Collection: Pilot Study and Prospective Analysis of the Draft Revised Form 33, Safety and Health Program Assessment Worksheet.

OMB Control Number: 1218–0280.

Affected Public: State, Local, and Tribal Governments.

Total Estimated Number of Respondents: 503.

Total Estimated Number of Responses: 810.

Total Estimated Annual Time Burden: 888 hours.

Total Estimated Annual Other Costs Burden: $0.

(Authority: 44 U.S.C. 3507(a)(1)(D))

Nicole Bouchet,

Senior Paperwork Reduction Act Analyst.

[FR Doc. 2025–08489 Filed 5–13–25; 8:45 am]

BILLING CODE 4510–26–P

NATIONAL FOUNDATION ON THE ARTS AND THE HUMANITIES

National Endowment for the Humanities

Meeting of Humanities Panel

AGENCY: National Endowment for the Humanities; National Foundation on the Arts and the Humanities.

ACTION: Notice of meeting.

SUMMARY: The National Endowment for the Humanities (NEH) will hold three meetings, by video conference, of the Humanities Panel, a federal advisory committee, during May 2025. The purpose of the meetings is for panel review, discussion, evaluation, and recommendation of applications for financial assistance under the National Foundation on the Arts and the Humanities Act of 1965.

DATES: See **SUPPLEMENTARY INFORMATION** for meeting dates. The meetings will open at 8:30 a.m. and will adjourn by 5 p.m. on the dates specified below.

FOR FURTHER INFORMATION CONTACT: Elizabeth Voyatzis, Committee Management Officer, 400 7th Street SW, Room 4060, Washington, DC 20506; (202) 606–8322; *evoyatzis@neh.gov*.

SUPPLEMENTARY INFORMATION: Pursuant to section 10(a)(2) of the Federal Advisory Committee Act (5 U.S.C. 10), notice is hereby given of the following meetings:

1. Date: May 8, 2025

This video meeting will discuss applications on the topics of Senior Research Grants, for the Documenting Endangered Languages—Preservation grant program, submitted to the Division of Preservation and Access.

2. Date: May 9, 2025

This video meeting will discuss applications on the topics of Senior Research Grants, for the Documenting Endangered Languages—Preservation grant program, submitted to the Division of Preservation and Access.

3. Date: May 12, 2025

This video meeting will discuss applications on the topics of Senior Research Grants, for the Documenting Endangered Languages—Preservation grant program, submitted to the Division of Preservation and Access.

Because these meetings will include review of personal and/or proprietary financial and commercial information given in confidence to the agency by grant applicants, the meetings will be closed to the public pursuant to sections 552b(c)(4) and 552b(c)(6) of title 5, U.S.C., as amended. I have made this determination pursuant to the authority granted me by the Chair's Delegation of Authority to Close Advisory Committee Meetings dated April 15, 2016.

Dated: May 8, 2025.

Jessica Graves,

Paralegal Specialist, National Endowment for the Humanities.

[FR Doc. 2025–08460 Filed 5–13–25; 8:45 am]

BILLING CODE 7536–01–P

NATIONAL SCIENCE FOUNDATION

Proposal Review Panel for Computing & Communication Foundations; Notice of Meeting

In accordance with the Federal Advisory Committee Act (Pub. L. 92–463, as amended), the National Science Foundation (NSF) announces the following meeting:

Name and Committee Code: Proposal Review Panel for Computing & Communication Foundations (#1192) (Virtual Site Visit).

Date and Time: June 2, 2025; 1 p.m.–2 p.m. (PDT); June 3, 2025; 7 a.m.–3:30 p.m. (PDT); June 4, 2025; 7 a.m.–3:45 p.m. (PDT).

Place: NSF, 2415 Eisenhower Avenue, Alexandria, VA 22314 (Virtual).

Type of Meeting: Part-Open.

Contact Persons: Basu, Sankar, National Science Foundation, 2415 Eisenhower Avenue, Alexandria, VA 22314; Telephone: 703–292–7843.

Purpose of Meeting: NSF virtual site visit to provide advice and recommendations concerning the progress of the TILOS AI Institute and to conduct a renewal review during year 4 of the award period as stipulated in the cooperative agreement.