

Name of Committee: National Institute on Drug Abuse Special Emphasis Panel; Mechanism for Time-Sensitive Drug Abuse Research.

Date: March 20, 2023.

Time: 12:00 p.m. to 1:30 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, National Institute on Drug Abuse, 301 North Stonestreet Avenue, Bethesda, MD 20892 (Virtual Meeting).

Contact Person: Sudhirkumar Udhavrao Yanpallewar, M.D., Scientific Review Officer, Scientific Review Branch, National Institute on Drug Abuse, NIH, 301 North Stonestreet Avenue, Bethesda, MD 20892, (301) 443-4577, sudhirkumar.yanpallewar@nih.gov.

Name of Committee: National Institute on Drug Abuse Special Emphasis Panel; HEAL Initiative: HEAL Data2Action—Innovation and Acceleration Projects, Phased Awards.

Date: March 22, 2023.

Time: 12:00 p.m. to 3:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, National Institute on Drug Abuse, 301 North Stonestreet Avenue, Bethesda, MD 20892 (Virtual Meeting).

Contact Person: Trinh T. Tran, Ph.D., Scientific Review Officer, Scientific Review Branch, Office of Extramural Policy, National Institute on Drug Abuse, NIH, 301 North Stonestreet Avenue, Bethesda, MD 20892, (301) 827-5843 trinh.tran@nih.gov.

Name of Committee: National Institute on Drug Abuse Special Emphasis Panel; Registry of Medical Cannabis Use and Health Outcomes.

Date: March 24, 2023.

Time: 11:00 a.m. to 5:00 p.m.

Agenda: To review and evaluate cooperative agreement applications.

Place: National Institutes of Health, National Institute on Drug Abuse, 301 North Stonestreet Avenue, Bethesda, MD 20892 (Virtual Meeting).

Contact Person: Gerald L. McLaughlin, Ph.D., Scientific Review Officer, Office of Extramural Policy and Review, National Institute on Drug Abuse, NIH, 301 North Stonestreet Avenue, Bethesda, MD 20892, (301) 827-5819, gm145a@nih.gov.

Name of Committee: National Institute on Drug Abuse Special Emphasis Panel; Omnibus Topic 167: Cause of Death Elucidated (CODE) in Drug Overdose: Research and Development of New Postmortem Toxicology Screening Devices That Are Portable, Rapid, Accurate, Affordable, and Accessible.

Date: March 30, 2023.

Time: 11:00 a.m. to 5:00 p.m.

Agenda: To review and evaluate contract proposals.

Place: National Institutes of Health, National Institute on Drug Abuse, 301 North Stonestreet Avenue, Bethesda, MD 20892 (Virtual Meeting).

Contact Person: Gerald L. McLaughlin, Ph.D., Scientific Review Officer, Office of Extramural Policy and Review, National Institute on Drug Abuse, NIH, 301 North Stonestreet Avenue, Bethesda, MD 20892, (301) 827-5819, gm145a@nih.gov.

(Catalogue of Federal Domestic Assistance Program Nos. 93.277, Drug Abuse Scientist Development Award for Clinicians, Scientist Development Awards, and Research Scientist Awards; 93.278, Drug Abuse National Research Service Awards for Research Training; 93.279, Drug Abuse and Addiction Research Programs, National Institutes of Health, HHS)

Dated: February 17, 2023.

Tyeshia M. Roberson-Curtis,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2023-03846 Filed 2-23-23; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Notice To Announce Updated Minimum Performance Standards for Experienced Firms That Receive Funding Through the Small Business Innovation Research (SBIR) and Small Business Technology Transfer (STTR) Programs

AGENCY: National Institutes of Health, HHS.

ACTION: Notice.

SUMMARY: The National Institutes of Health (NIH) announces the updated minimum performance standards for experienced firms funded through the Department of Health and Human Services (HHS) Small Business Innovation Research (SBIR) and Small Business Technology Transfer (STTR) Programs.

DATES: The relevant funding opportunity announcements have been updated to incorporate these changes. The updated performance standards will be required of any firms submitting SBIR or STTR grant or cooperative agreement applications on or after April 5, 2023.

ADDRESSES: Please visit our website to view the updated Minimum Performance Standards for Experienced Firms at <https://seed.nih.gov/small-business-funding/small-business-program-basics/eligibility-criteria>.

FOR FURTHER INFORMATION CONTACT: Stephanie Fertig, HHS Small Business Program Lead, Small business Education & Entrepreneurial Development (SEED) Office, Office of Extramural Research, NIH, Rockledge I, Suite 800, Bethesda, MD 20817. Email: seedinfo@nih.gov. Phone number (301) 435-2688.

SUPPLEMENTARY INFORMATION: The new minimum standards are aligned with Section 9 of the Small Business Act (15

U.S.C. 638), as amended by the SBIR and STTR Extension Act of 2022 (Pub. L. 117-183).

HHS is announcing the following changes:

Phase I to Phase II Transition Rate Benchmark: In accordance with guidance from the SBA, the HHS SBIR/STTR Program is implementing the Phase I to Phase II Transition Rate benchmark required by the SBIR/STTR Reauthorization Act of 2011 and the SBIR and STTR Extension Act of 2022. The benchmark establishes a minimum number of Phase II awards the company must have received for a given number of Phase I awards received during the 5-year time period. The Transition Rate is calculated as the total number of SBIR and STTR Phase II awards a company received during the past 5 fiscal years divided by the total number of SBIR and STTR Phase I awards it received during the past 5 fiscal years excluding the most recently-completed year.

Phase II to Commercialization Benchmark: In accordance with guidance from the SBA, HHS, including NIH, SBIR/STTR Programs are implementing the Phase II to Commercialization Rate benchmark for Phase I applicants, as required by the SBIR/STTR Reauthorization Act of 2011 and the SBIR and STTR Extension Act of 2022. The Commercialization Rate Benchmark was published in a **Federal Register** notice on August 8, 2013 (78 FR 48537).

This update is applicable to all HHS SBIR and STTR grants and cooperative agreements with application receipt dates on or after April 5, 2023. This update supersedes, in its entirety, previous Phase I to Phase II transition benchmarks established in May 2013 (78 FR 30951) and previous Commercialization Benchmarks established in September 2013 (78 FR 59410). Additional information can be found at: <https://grants.nih.gov/grants/guide/notice-files/not-od-23-092.htm>.

Dated: February 16, 2023.

Tara A. Schwetz,

Acting Principal Deputy Director, National Institutes of Health.

[FR Doc. 2023-03798 Filed 2-23-23; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Library of Medicine; Notice of Meeting

Pursuant to section 10(d) of the Federal Advisory Committee Act, as

amended, notice is hereby given of a meeting of the Board of Regents of the National Library of Medicine.

The meeting will be open to the public as indicated below. Individuals who plan to attend 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.

The meeting will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C., as amended. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable materials, and personal information concerning individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: Board of Regents of the National Library of Medicine.

Date: May 9, 2023.

Open: May 9, 2023, 10:00 a.m. to 4:00 p.m.

Agenda: Program Discussion.

Place: National Library of Medicine, 8600 Rockville Pike, Bethesda, MD 20892 (Virtual Meeting).

Closed: May 9, 2023, 4:00 p.m. to 4:30 p.m.

Agenda: To review and evaluate grant applications.

Contact Person: Christine Ireland, Committee Management Officer, Division of Extramural Programs, National Library of Medicine, Bethesda, MD 20892, 301-594-4929, irelanc@mail.nih.gov.

Any member of the public may submit written comments no later than 15 days in advance of the meeting. 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.

Information is also available on the Institute's/Center's home page: www.nlm.nih.gov/od/bor/bor.html where additional information for the meeting will be posted when available. This meeting will be broadcast to the public, and available for viewing at <https://videocast.nih.gov> on May 9, 2023.

(Catalogue of Federal Domestic Assistance Program No. 93.879, Medical Library Assistance, National Institutes of Health, HHS).

Dated: February 21, 2023.

Miguelina Perez,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2023-03845 Filed 2-23-23; 8:45 am]

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DEPARTMENT OF THE INTERIOR

Fish and Wildlife Service

[FWS-R4-ES-2023-N001;
FXES11140400000-223-FF04E00000]

Endangered Species; Recovery Permit Applications

AGENCY: Fish and Wildlife Service, Interior.

ACTION: Notice of receipt of permit applications; request for comments.

SUMMARY: We, the U.S. Fish and Wildlife Service, have received applications for permits to conduct activities intended to enhance the propagation or survival of endangered species under the Endangered Species Act. We invite the public and local, State, Tribal, and Federal agencies to comment on these applications. Before issuing any of the requested permits, we will take into consideration any information that we receive during the public comment period.

DATES: We must receive written data or comments on the applications by March 27, 2023.

ADDRESSES:

Reviewing Documents: Submit requests for copies of applications and other information submitted with the applications to Karen Marlowe (see **FOR FURTHER INFORMATION CONTACT**). All requests and comments should specify the applicant name and application number (e.g., Mary Smith, ESPER0001234).

Submitting Comments: If you wish to comment, you may submit comments by one of the following methods:

- *Email (preferred method):*

permitsR4ES@fws.gov. Please include your name and return address in your email message. If you do not receive a confirmation from the U.S. Fish and Wildlife Service that we have received your email message, contact us directly at the telephone number listed in **FOR FURTHER INFORMATION CONTACT**.

- *U.S. Mail:* U.S. Fish and Wildlife Service Regional Office, Ecological Services, 1875 Century Boulevard, Atlanta, GA 30345 (Attn: Karen Marlowe, Permit Coordinator).

FOR FURTHER INFORMATION CONTACT: Karen Marlowe, Permit Coordinator, 404-679-7097 (telephone) or karen_marlowe@fws.gov (email). Individuals in the United States who are deaf, deafblind, hard of hearing, or have a speech disability may dial 711 (TTY, TDD, or TeleBraille) to access telecommunications relay services. Individuals outside the United States should use the relay services offered

within their country to make international calls to the point-of-contact in the United States.

SUPPLEMENTARY INFORMATION: We, the U.S. Fish and Wildlife Service, invite review and comment from the public and local, State, Tribal, and Federal agencies on applications we have received for permits to conduct certain activities with endangered and threatened species under section 10(a)(1)(A) of the Endangered Species Act of 1973, as amended (ESA; 16 U.S.C. 1531 *et seq.*), and our regulations in the Code of Federal Regulations (CFR) at 50 CFR part 17. Documents and other information submitted with the applications are available for review, subject to the requirements of the Privacy Act of 1974, as amended (5 U.S.C. 552a) and the Freedom of Information Act (5 U.S.C. 552).

Background

With some exceptions, the ESA prohibits take of listed species unless a Federal permit is issued that authorizes such take. The ESA's definition of "take" includes hunting, shooting, harming, wounding, or killing, and also such activities as pursuing, harassing, trapping, capturing, or collecting.

A recovery permit issued by us under section 10(a)(1)(A) of the ESA authorizes the permittee to take endangered or threatened species while engaging in activities that are conducted for scientific purposes that promote recovery of species or for enhancement of propagation or survival of species. These activities often include the capture and collection of species, which would result in prohibited take if a permit were not issued. Our regulations implementing section 10(a)(1)(A) for these permits are found at 50 CFR 17.22 for endangered wildlife species, 50 CFR 17.32 for threatened wildlife species, 50 CFR 17.62 for endangered plant species, and 50 CFR 17.72 for threatened plant species.

Permit Applications Available for Review and Comment

The ESA requires that we invite public comment before issuing these permits. Accordingly, we invite local, State, Tribal, and Federal agencies, and the public to submit written data, views, or arguments with respect to these applications. The comments and recommendations that will be most useful and likely to influence agency decisions are those supported by quantitative information or studies. Proposed activities in the following permit requests are for the recovery and enhancement of propagation or survival of the species in the wild.