based integrated approach to opioid prevention and treatment services in a single community?

**Health Economics:**
- What economic questions should be included as part of the study to inform systems and policy change?

**Implementation Research:**
- What implementation research questions should be included to develop best practices for replication in other communities impacted by the opioid crisis?
- What data should be collected to help develop metrics for determining the quality of an integrated approach to opioid prevention and treatment services, including policies and practices?
- Are there examples of prior implementation research studies that highlight implementation tools that can be used to replicate and scale up integrated approaches?

**Infrastructure, Partnerships, Collaboration:**
- What research, prevention, and treatment infrastructure and partnerships are needed to support a community-based pragmatic trial assessing the impact of an evidence-based integrated approach to opioid prevention and treatment services?
- What is the best approach to fostering collaboration and meaningful participation between state, county, and local governments; community stakeholders; medical/clinical service providers; and researchers?
- How do we construct a research initiative with the highest likelihood of having sustainable prevention and treatment services?
- What data would be of most interest to state and community partners?

Responses to this RFI are voluntary and may be submitted anonymously. Please do not include any personally identifiable or other information that you do not wish to make public.

Proprietary, classified, confidential, or sensitive information should not be included in responses. Comments submitted will be compiled for discussion and shared internally with NIDA, SAMHSA, NIH program staff, and participating leadership across the Department of Health and Human Services, as appropriate. Any personal identifiers (personal names, email addresses, etc.) will be removed when responses are compiled.

This RFI is for informational and planning purposes only and is not a solicitation for applications or an obligation on the part of the United States Government to provide support for any ideas identified in response to it. Please note that the United States Government will not pay for the preparation of any information submitted or for use of that information.

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**National Institutes of Health**

**National Institute of Diabetes and Digestive and Kidney Diseases; Notice of Closed Meetings**

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended, notice is hereby given of the following meetings.

The meetings 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 material, 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:** National Institute of Diabetes and Digestive and Kidney Diseases Special Emphasis Panel; T1D NIDDK Review.

**Date:** June 26, 2018.
**Time:** 11:00 a.m. to 1:00 p.m.
**Agenda:** To review and evaluate grant applications.

**Place:** National Institutes of Health, Two Democracy Plaza, 6707 Democracy Boulevard, Bethesda, MD 20892 (Telephone Conference Call).

**Contact Person:** Jason D. Hoffert, Ph.D., Scientific Review Officer, Review Branch, DEA, NIDDK, National Institutes of Health, Room 7343, 6707 Democracy Boulevard, Bethesda, MD 20817, 301–496–9010, jhoffert@niddk.nih.gov.

**Name of Committee:** National Institute of Diabetes and Digestive and Kidney Diseases Special Emphasis Panel; RFA–DK–17–020: Immune System Engineering for Targeted Tolerance in Type 1 Diabetes (R01).

**Date:** July 25, 2018.
**Time:** 11:00 a.m. to 4:30 p.m.
**Agenda:** To review and evaluate grant applications.

**Place:** National Institutes of Health, Two Democracy Plaza, 6707 Democracy Boulevard, Bethesda, MD 20892 (Telephone Conference Call).

**Contact Person:** Dianne Campl, Ph.D., Scientific Review Officer, Review Branch, DEA, NIDDK, National Institutes of Health, Room 7013, 6707 Democracy Boulevard, Bethesda, MD 20892–2542, 301–594–7682, campdextra.niddk.nih.gov.

**Name of Committee:** National Institute of Diabetes and Digestive and Kidney Diseases Special Emphasis Panel; SBR Phase II Clinical Trials.

**Date:** July 26, 2018.
**Time:** 11:00 a.m. to 1:00 p.m.
**Agenda:** To review and evaluate grant applications.

**Place:** National Institutes of Health, Two Democracy Plaza, 6707 Democracy Boulevard, Bethesda, MD 20892 (Telephone Conference Call).

**Contact Person:** Thomas A. Tatham, Ph.D., Scientific Review Officer, Review Branch, DEA, NIDDK, National Institutes of Health, Room 7021, 6707 Democracy Boulevard, Bethesda, MD 20892–5452, (301) 594–3993, ttatham@niddk.nih.gov.

**Name of Committee:** National Institute of Diabetes and Digestive and Kidney Diseases Special Emphasis Panel; PAR—18–042: NIDDK Ancillary Studies (R01).

**Date:** July 26, 2018.
**Time:** 11:30 a.m. to 1:30 p.m.
**Agenda:** To review and evaluate grant applications.

**Place:** National Institutes of Health, Two Democracy Plaza, 6707 Democracy Boulevard, Bethesda, MD 20892 (Telephone Conference Call).

**Contact Person:** Ann A. Jerkins, Ph.D., Scientific Review Officer, Review Branch, DEA, NIDDK, National Institutes of Health, Room 7349, 6707 Democracy Boulevard, Bethesda, MD 20892–5452, (301) 594–8894, begumn@niddk.nih.gov.
DEA, NIDDK. National Institutes of Health, Room 7119, 6707 Democracy Boulevard, Bethesda, MD 20892–5452, 301–594–2242, jerkinsa@niddk.nih.gov.


Date: July 30, 2018.

Time: 11:00 a.m. to 3:00 p.m.

Agenda: To review and evaluate cooperative agreement applications.

Place: National Institutes of Health, Two Democracy Plaza, 6707 Democracy Boulevard, Bethesda, MD 20892 (Telephone Conference Call).

Contact Person: Najma S. Begum, Ph.D., Scientific Review Officer Review Branch, DEA, NIDDK, National Institutes of Health, Room 7349, 6707 Democracy Boulevard, Bethesda, MD 20892–5452, (301) 594–8894, begumn@niddk.nih.gov.

(Catalogue of Federal Domestic Assistance Program Nos. 93.847, Diabetes, Endocrinology and Metabolic Research; 93.848, Digestive Diseases and Nutrition Research; 93.849, Kidney Diseases, Urology and Hematology Research, National Institutes of Health, HHS)


David D. Clary,
Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2018–13966 Filed 6–28–18; 8:45 am]
BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Center for Advancing Translational Sciences; Notice of Closed Meeting

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended, notice is hereby given of the following 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 contract proposals and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the contract proposals, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: National Center for Advancing Translational Sciences Special Emphasis Panel; SBIR Contract Review.

Date: July 25, 2018.

Time: 1:00 p.m. to 4:30 p.m.

Agenda: To review and evaluate contract proposals.

Place: National Institutes of Health, One Democracy Plaza, Room 1087, 6701 Democracy Boulevard, Bethesda, MD 20892 (Telephone Conference Call).

Contact Person: Rahat (Rani) Khan, Ph.D., Scientific Review Officer, Office of Scientific Review, National Center for Advancing Translational Sciences, 6701 Democracy Blvd., Rm 1078, Bethesda, MD 20892, 301–594–7319, khanzr2@csrc.nih.gov.

(Catalogue of Federal Domestic Assistance Program Nos. 93.859, Pharmacology, Physiology, and Biological Chemistry Research; 93.350, B—Cooperative Agreements; 93.859, Biomedical Research and Research Training, National Institutes of Health, HHS)


David D. Clary,
Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2018–13966 Filed 6–28–18; 8:45 am]
BILLING CODE 4140–01–P

DEPARTMENT OF HOMELAND SECURITY

Federal Emergency Management Agency

[Docket ID: FEMA–2018–0025; OMB No. 1660–0040]

Agency Information Collection Activities: Proposed Collection; Comment Request; Standard Flood Hazard Determination Form

AGENCY: Federal Emergency Management Agency, DHS.

ACTION: Notice and request for comments.

SUMMARY: The Federal Emergency Management Agency, as part of its continuing effort to reduce paperwork and respondent burden, invites the general public to take this opportunity to comment on an extension, without change, of a currently approved information collection. In accordance with the Paperwork Reduction Act of 1995, this notice seeks comments concerning this form which is used by regulated lending institutions, federal agency lenders, the Federal National Mortgage Association, the Federal Home Loan Mortgage Corporation, and the Government National Mortgage Association. Federally regulated lending institutions complete this form when making, increasing, extending, renewing or purchasing each loan for the purpose is of determining whether flood insurance is required and available. FEMA is responsible for maintaining the form and making it available.

DATES: Comments must be submitted on or before August 28, 2018.

ADDRESSES: To avoid duplicate submissions to the docket, please use only one of the following means to submit comments:

(1) Online. Submit comments at www.regulations.gov under Docket ID FEMA–2018–0025. Follow the instructions for submitting comments.

(2) Mail. Submit written comments to Docket Manager, Office of Chief Counsel, DHS/FEMA, 500 C Street SW, 8NE, Washington, DC 20472–3100.

All submissions received must include the agency name and Docket ID. Regardless of the method used for submitting comments or material, all submissions will be posted, without change, to the Federal eRulemaking Portal at http://www.regulations.gov, and will include any personal information you provide. Therefore, submitting this information makes it public. You may wish to read the Privacy Act notice that is available via the link in the footer of www.regulations.gov.

FOR FURTHER INFORMATION CONTACT:
Susan Bernstein, Insurance Specialist, FEMA, Marketing and Outreach Branch, 303–701–3595. You may contact the Information Management Division for copies of the proposed collection of information at email address: FEMA-Information-Collectons-Management@ fema.dhs.gov.

SUPPLEMENTARY INFORMATION: Section 1365 of the National Flood Insurance Act of 1968 (NFIA) (42 U.S.C. 4104b), as added by Section 528 of the National Flood Insurance Reform Act of 1994 (Pub. L. 103–325, title V), requires that FEMA develop a standard hazard determination form for recording the determination of whether a structure is located within an identified Special Flood Hazard Area and whether flood insurance is available. Regulated lending institutions, federal agency lenders, the Federal National Mortgage Association, the Federal Home Loan Mortgage Corporation, and the Government National Mortgage Association must complete this form for any loan made, increased, extended, renewed or purchased by these entities. The requirement for federally regulated lending institutions to determine whether a building or mobile home securing a loan is located in an area having special flood hazards and whether flood insurance is available has been in effect since the enactment of the Flood Disaster Protection Act of 1973, although the use of a standard form was not required until the enactment of the Section 1365 of the NFIA. The establishment of the Standard Flood Hazard Determination form has enabled