

likely to begin and accelerates, often leading to substance abuse disorder. Notably, the peak period for the manifestation of cannabis-use disorder is age 18–19, and the past-year-prevalence for alcohol-use disorder is age 20–22. The TLI is designed to identify the propensity for these and other substance abuse prior to manifestation; as such, collecting data from the high school age group (14–18

years old) is critical to identifying at-risk youths for the purposes of early intervention. Thus, the TLI must be tested with data collected from youth populations, ages 14 to 18, comparable to those in existing studies. Moreover, the R&D team must provide psychometric external validation for the TLI through data collection from sets of identical and fraternal twins. Psychometric analyses are required to

show that the TLI performs according to expectations. Accordingly, studies will be performed on the collected information to demonstrate i) construct, ii) discriminative, iii) concurrent, and iv) predictive validity.

OMB approval is requested for 2 years. There are no costs to respondents other than their time. The total estimated annualized burden hours are 3,083.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent: individuals and households	Number of respondents	Responses per respondent	Average burden per response (in hours)	Annual hour burden
Parent of 14–17 year-old students: Consent Form	5,000	1	1/60	83
14–18 year-old students: School Survey (TLI)		1	30/60	2,500
14–18 year-old youths or their parents: Consent Form	600	1	1/60	10
14–18 year-old youths: Twins Survey (Demo/D&A)		1	10/60	100
14–18 year-old youths: Twins Survey (Dysregulation)		1	10/60	100
14–18 year-old youths: Twins Survey (TLI)		1	29/60	290

Dated: November 20, 2013.

Glenda J. Conroy,

Executive Officer (OM Director), NIDA, NIH.

[FR Doc. 2013–28985 Filed 12–2–13; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute of Neurological Disorders and Stroke; Notice of Closed Meeting

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. App.), 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 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: National Institute of Neurological Disorders and Stroke, Special Emphasis Panel, Stroke Trials Network-NDMC.

Date: December 18, 2013.

Time: 9:00 a.m. to 2:30 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Neuroscience Center, 6001 Executive

Boulevard, Rockville, MD 20852, (Telephone Conference Call).

Contact Person: Natalia Strunnikova, Ph.D., Scientific Review Officer, Scientific Review Branch, Division of Extramural Research, NINDS, NIH, NSC, 6001 Executive Blvd., Suite 3208, MSC 9529, Bethesda, MD 20892–9529, 301–402–0288, *Natalia.Strunnikova@nih.gov*.

(Catalogue of Federal Domestic Assistance Program Nos. 93.853, Clinical Research Related to Neurological Disorders; 93.854, Biological Basis Research in the Neurosciences, National Institutes of Health, HHS)

Dated: November 26, 2013.

Carolyn Baum,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2013–28854 Filed 12–2–13; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute of Diabetes and Digestive and Kidney Diseases; Notice of Closed Meeting

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. App.), 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 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, PAR13–228: Biomarkers for Diabetes, Digestive, Kidney and Urologic Diseases using Repository Biosamples.

Date: February 20, 2014.

Time: 2:00 p.m. to 5:00 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: Najma Begum, Ph.D., Scientific Review Officer, Review Branch, DEA, NIDDK, National Institutes of Health, Room 749, 6707 Democracy Boulevard, Bethesda, MD 20892–5452, (301) 594–8894, *begumn@nidk.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)

Dated: November 26, 2013.

David Clary,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2013–28851 Filed 12–2–13; 8:45 am]

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