contact John D. Hewes, Ph.D. at hewesj@mail.nih.gov.

Software for Evaluating Drug Induced Hepatotoxicity

Description of Technology: This invention pertains to a software tool for assisting differential medical diagnosis of drug-induced liver injury (hepatotoxicity) using clinical trial data. The software is capable of identifying a small subset of patients at risk for hepatotoxicity out of a pool of thousands of clinical trial participants. This software tool is the only one of its kind developed using SAS/IntrNet®.

Potential Commercial Applications:

• Hepatotoxicity detection
• Drug interactions

Competitive Advantages:

• Personalized predictions
• SAS/IntrNet® compatible

Development Stage: Prototype

Inventor: Ted J. Guo (FDA)

Publications:


• US Provisional Application No. 61/529,531 filed 31 August 2011
• PCT Application No. PCT/GB2012/052140 filed 31 August 2012

Licensing Contact: Jaime M. Greene; greenejaime@mail.nih.gov


Richard U. Rodriguez,
Director, Division of Technology Development and Transfer, Office of Technology Transfer, National Institutes of Health.

[FR Doc. 2013–30745 Filed 12–24–13; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Center For Scientific Review; Notice of Closed Meetings

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 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: Center for Scientific Review Special Emphasis Panel; Member Conflict: Biological Chemistry and Macromolecular Biophysics.

Date: January 6, 2014.
Time: 1:00 p.m. to 2:00 p.m.
Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892, (Telephone Conference Call).

Contact Person: John L. Bowers, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 4170, MSC 7806, Bethesda, MD 20892, (301) 435–1725, bowers@csr.nih.gov.

This notice is being published less than 15 days prior to the meeting due to the timing limitations imposed by the review and funding cycle.

Name of Committee: Center for Scientific Review Special Emphasis Panel; Myalgic Encephalomyelitis/Chronic Fatigue Syndrome.

Date: January 16, 2014.
Time: 2:30 p.m. to 4:30 p.m.
Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892, (Virtual Meeting).

Contact Person: Lynn E. Luethke, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 5166, MSC 7844, Bethesda, MD 20892, (301) 806–3323, luethke@csr.nih.gov.

Name of Committee: Center for Scientific Review Special Emphasis Panel; Program Project: Research Resources Reverse Site Visit.

Date: January 21–23, 2014.
Time: 7:30 p.m. to 12:30 p.m.
Agenda: To review and evaluate grant applications.

Place: Crowne Plaza Washington, DC—Rockville Hotel, 3 Research Ct., Rockville, MD 20850.

Contact Person: Lee Rosen, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 5116, MSC 7854, Bethesda, MD 20892, (301) 435–1171, rosen@csr.nih.gov.


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

[FR Doc. 2013–30747 Filed 12–24–13; 8:45 am]

BILLING CODE 4140–01–P