DEPARTMENT OF HEALTH AND HUMAN SERVICES
National Institutes of Health

Government-Owned Inventions; Availability for Licensing

AGENCY: National Institutes of Health, HHS.

ACTION: Notice.

SUMMARY: The inventions listed below are owned by an agency of the U.S. Government and are available for licensing in the U.S. in accordance with 35 U.S.C. 209 and 37 CFR part 404 to achieve expeditious commercialization of results of federally-funded research and development. Foreign patent applications listed below may be obtained by writing to the indicated licensing contact at the Office of Technology Transfer, National Institutes of Health, 6011 Executive Boulevard, Suite 325, Rockville, Maryland 20852–3804; telephone: 301–496–7057; fax: 301–402–0220. A signed Confidential Disclosure Agreement will be required to receive copies of the patent applications.

FOR FURTHER INFORMATION CONTACT: Licensing information and copies of the U.S. patent applications listed below may be obtained by writing to the indicated licensing contact at the Office of Technology Transfer, National Institutes of Health, 6011 Executive Boulevard, Suite 325, Rockville, Maryland 20852–3804; telephone: 301–496–7057; fax: 301–402–0220. A signed Confidential Disclosure Agreement will be required to receive copies of the patent applications.

SUPPLEMENTARY INFORMATION:

A Novel Rapid Point-of-Care Diagnostic Method for Infectious and Autoimmune Diseases

Description of Technology: A rapid assay for point-of-care diagnosis of infectious and autoimmune diseases. The assay takes 3.5 minutes, is highly efficient, and has low background. The National Institute of Neurological Disorders and Stroke, National Institutes of Health.

Potential Commercial Applications
• A rapid assay for point-of-care diagnosis of infectious and autoimmune diseases.
• Applications to different assay platforms, such as a portable, commercially available hand-held luminometer or an automated, high-throughput device.

Competitive Advantages
• Highly efficient, rapid, and easy to perform.
• Low background signals.

Development Stage
• Early-stage
• In vitro data available
• Prototype.

Inventor: Peter D. Burbelo (NIDCR)

Publications


Related Technologies

Licensing Contact: Sally Hu, Ph.D., M.B.A.; 301–435–5606; hus@mail.nih.gov.

Collaborative Research Opportunity: The National Institute of Dental and Craniofacial Research is seeking statements of capability or interest from parties interested in collaborative research to further develop, evaluate or commercialize using neodymium magnet for rapid diagnosis. For collaboration opportunities, please contact David Bradley, Ph.D. at bradleyd@niddcr.nih.gov.

A Mobile Health Platform

Description of Technology: The NIH inventors have developed a mobile health technology to monitor and predict a user’s psychological status and to deliver an automated intervention when needed. The technology uses smartphones to monitor the user’s location and ask questions about psychological status throughout the day. Continuous collected ambulatory psychological data are fused with data on location and responses to questions. The mobile data are combined with geospatial risk maps to quantify exposure to risk and predict a future psychological state. The future predictions are used to warn the user when he or she is at especially high risk of experiencing a negative event that might lead to an unwanted outcome (e.g., lapse to drug use in a recovering addict).

An internally developed mobile app is now being deployed to deliver an intervention in the context of drug addiction. The inventors are also seeking to test the technology for other health applications.

Potential Commercial Applications
• Real time behavior monitoring
• Therapeutic delivery of an intervention via a mobile device

Competitive Advantages
• Mobile device
• Real time
• Exposure to risk

Development Stage: Prototype

Inventors: Kenzie L. Preston, David H. Epstein, Matthew Tyburski, Massoud Vahabzadeh (all of NIDA)

Publications


Licensing Contact: Betty B. Tong, Ph.D.; 301–594–6565; tongb@mail.nih.gov

Collaborative Research Opportunity: The National Institute on Drug Abuse is seeking statements of capability or interest from parties interested in collaborative research to further develop, evaluate or commercialize
mhealth system to analyze and intervene. For collaboration opportunities, please contact Vio Conley at conleyv@mail.nih.gov.

**Detection and Discrimination of Classical and Atypical L-Type BSE Strains by RT-QuIC**

*Description of Technology:* Statutory surveillance of bovine spongiform encephalopathy (BSE) indicates that cattle are susceptible to both classical (C–BSE) and atypical forms of BSE. Atypical forms of BSE appear to be sporadic and thus may never be eradicated. A major challenge is the lack of sufficiently practical and sensitive tests for routine BSE detection and strain discrimination. The RT-QuIC test, which is based on prion-seeded fibrillization of recombinant prion protein (pPrPSc), is known to be highly specific and sensitive for detection of multiple human and animal prion diseases, but not BSE. This application claims methods for distinguishing whether a sheep, cow or goat has atypical L-bovine spongiform encephalopathy prion or classical bovine spongiform encephalopathy.

**Potential Commercial Applications**
- Detection and distinguishing of both BSE forms
- Rapid detection and discrimination of BSE forms

**Competitive Advantages**
- Orders of magnitude more sensitive than ELISA tests
- Eliminates need for multi-phase analyses of samples
- Can be applied to large scale testing of multiple samples

**Development Stage**
- *In vitro* data available
- *In vivo* data available (animal)
- Prototype

**Inventors:** Nigel H. Greig, Weiming Luo, David Tweedie, Harold W. Holloway, Qian-sheng Yu (all of NIA)

**Publication**

**Intelectual Property:** HHS Reference No. E–045–2012/0—
- US Patent No. 8,927,725 issued 06 Jan 2015
- US Patent No. 9,084,783 issued 21 Jul 2015

**Related Technologies:** HHS Reference No. E–189–2003/0—
- US Patent No. 8,546,430 issued 01 Oct 2013

**Inventors:** Byron W. Caughey (NIAID), Cristina D. Orrú (NIAID), Alessandra Favolez (EM), Cristina Casalone (EM), Maria Massa (EM), Cristiano Corona (EM)

**Related Patents/Intellectual Property:**
- and related international patents/patent applications

**License Contact:** Betty B. Tong, Ph.D.; 301–594–6565; tongb@mail.nih.gov

**Novel Regulatory B Cells for Treatment of Cancer and Autoimmune Disease**

*Description of Technology:* The manner by which cancers evade the immune response is not well-understood. What is known is that the manner is an active process that regulates immune responses employing at least two types of suppressive cells, myeloid-derived suppressive cells and regulatory T cells (Tregs), a key subset of CD4+ T cells that controls peripheral tolerance to self- and allo-antigens. Tregs are considered to play a key role in the escape of cancer cells from anti-tumor effector T cells. Cancer cells have been found to directly activate resting B cells to form suppressive regulatory B cells (tBregs) and utilize them to evade immune surveillance and mediate metastasis. tBregs directly inhibit CD4+ and CD8+ T cell activity in a cell contact-dependent manner, induce FoxP3+ T cell activity, and promote Treg-dependent metastasis. Researchers from the National Institute on Aging (NIA), NIH, have developed methods for the generation of tBregs, and for using tBregs to produce...
Tregs, and methods that inactivate or deplete Tregs. These methods have significant therapeutic value in the combat with cancer immune escape and metastasis, and in the control of harmful autoimmune diseases.

**Potential Commercial Applications:**
- Production of cellular cancer vaccines
- Treatments for immune-mediated disorders
- Treatments for cancer
- Treatments for chronic viral infections

**Development Stage:**
- Early-stage
- In vitro data available
- In vivo data available (animal)
- Ex vivo data available

**Inventors:** Bira Arya and Purevdorj Olkhanud (NIA)


**Licensing Contact:** Betty B. Tong, Ph.D.; 301–594–6565; tongb@mail.nih.gov

**Collaborative Research Opportunity:**
The National Institute on Aging, Laboratory of Molecular Biology and Immunology, is seeking statements of capability or interest from parties interested in collaborative research to further develop, evaluate, or commercialize the utilization of regulatory B cells to control autoimmune diseases and strategies that inactivate Tregs to control cancer immune escape. Please contact Nicole Darack, Ph.D. at 240–276–5493 or darackn@mail.nih.gov for more information.

**Immunogenic Tumor-associated Antigen SPANX–B for Selective Cancer Immunotherapy**

**Description of Technology:**
Researchers at the National Institute on Aging (NIA) have characterized a novel tumor-associated antigen, SPANX–B, which is naturally immunogenic and is expressed in a variety of human malignancies, including melanoma and lung, colon, renal, ovarian and breast carcinomas. In melanoma specifically, SPANX–B expression is associated with advanced and metastatic disease. Moreover, the researchers have found several agonist epitope peptides from SPANX–B which can be used to activate the immune system to eradicate tumors utilizing T cells. SPANX–B peptides have significant clinical and immunotherapeutic potential for the development of cancer diagnostic assays and potent protective and/or therapeutic vaccines to combat a wide-range of cancers.

**Potential Commercial Applications:**
- In vitro diagnostic assays for highly-metastatic melanomas or other cancers
- Therapeutic monoclonal antibodies
- Cancer vaccine development

**Competitive Advantages:**
- Immunogenic: SPANX–B peptides are naturally able to elicit immune response.
- Expressed in a wide-range of cancers.
- Use of epitope peptides facilitates the activation of cells of the more therapeutically effective branch of the immune system.
- Small epitope peptides: Can be more easily manufactured in contrast to recombinant proteins.

**Development Stage:**
- In vitro data available
- In vivo data available (animal)


**Inventors:** Bira Arya (NIA) and Vladimir Larionov (NCI)


**Licensing Contact:** Betty B. Tong, Ph.D.; 301–594–6565; tongb@mail.nih.gov

**Collaborative Research Opportunity:**
The National Institute on Aging, Laboratory of Molecular Biology and Immunology, is seeking statements of capability or interest from parties interested in collaborative research to further develop, evaluate, or commercialize the use of SPANX–B-based therapeutic approaches to combat cancers. Please contact Nicole Darack, Ph.D. at 240–276–5493 or darackn@mail.nih.gov for more information.

**Method for the Diagnosis and Prognosis of Age-Related Cardiovascular Disorders**

**Description of Technology:**
NIH investigators have discovered a method for the diagnosis and prognosis of cardiovascular aging. Current methodologies include the measurement of patient lipid profiles or expression of up to two proteins. In contrast, this technology utilizes the expression levels of a panel of proteins not previously known to be related to cardiovascular aging and may prove to be a more accurate diagnostic or prognostic of cardiovascular aging than currently available tests or it may improve the accuracy of currently available tests when used in concert.

The technology relates to methods for determining susceptibility to having an extremely common age-associated vascular disorder. It also describes the subsequent use of these proteins as markers for disease. While the underlying cellular and molecular mechanisms of age-related vascular disease remain largely undefined, the expression levels of the genes described in this technology have been empirically determined to differ between healthy and age-inflamed arterial tissue. Further, this technology includes a companion mass spectroscopic-based methodology for reproducible quantification of specific expression levels of interest.

**Potential Commercial Applications:**
Diagnosis of age-related vascular disorder.

**Inventors:** Mingyi Wang et al. (NIA)


**Licensing Contact:** Betty B. Tong, Ph.D.; 301–594–6565; tongb@mail.nih.gov

**Collaborative Research Opportunity:**
The National Institute on Aging, Laboratory of Cardiovascular Science, Cardiac Biology Section—Vascular Group, is seeking statements of capability or interest from parties interested in collaborative research to further develop, evaluate, or commercialize the use of these antisense oligos for the treatment of diseases. The antisense oligos target genes known to be associated with age-related vascular disease.

**Description of Technology:** The biological phenomenon of RNA interference (RNAi) has much promise for developing therapeutics to a variety of diseases. However, development of RNAi therapies remains mainly in preclinical stages largely because of difficulties in delivering small inhibitory RNAs (siRNA) and short hairpin RNAs (shRNA) into target cells. Although viral vector-based siRNA delivery systems are extensively used, their specificity and safety remain significant issues. Without a
solution to this delivery problem. RNAi cannot fulfill its therapeutic promise. Investigators at the National Institutes of Health have developed novel compositions and methods for delivering inhibitory oligonucleotides to cells in a targeted and efficient manner. The compositions and methods are based on utilizing a cell surface receptor targeting ligand, such as cytokine or chemokine, and a domain that binds an inhibitory oligonucleotide, to efficiently deliver the inhibitory oligonucleotide to the cell that expresses the cell surface receptor targeted ligand. Chemokine receptors are differentially expressed on various cells, including tumors; hence this technology allows targeting siRNA to aberrant cells. Gene silencing can also be achieved in variety of immune cells by targeting cytokine receptors. This technology has great potential for developing into a safe and effective means of delivering therapeutic siRNAs.

**Potential Commercial Applications**

- Treatment of cancers and autoimmune diseases by delivery of siRNA to tumor cells or various aberrantly functioning immune cells.
- This technology can be used to boost vaccine responses against cancers and chronic infectious diseases.
- Targeted delivery of fluorochrome-labeled RNA both in vitro and in vivo for diagnostic purposes, for example, to trace or localize various cells and to determine tumor metastasis and aberrant proliferation or homing of immune cells.

**Competitive Advantages**

- Simple method for linking siRNA to polypeptides to create non-covalent or covalent complexes
- In vitro targeted delivery of inhibitory RNAs into cells rather than systemically
- Delivery of multiple inhibitory RNAs to target multiple genes
- Long-term repression of target gene expression through RNAi phenomenon

**Development Stage**

- In vitro data available
- In vivo data available (animal)
- In situ data available

**Inventors:** Bira Arya, Purovedorj Olkhanud, Juan Espinoza (all of NIA)

**Publications**

- Collaborative Research Opportunity: The National Institute on Aging, Laboratory of Molecular Biology and Immunology, is seeking statements of capability or interest from parties interested in collaborative research to further develop, evaluate, or commercialize chemokine-based siRNA/shRNA technology for treatment of cancers and autoimmune diseases, i.e., to control expression of immunomodulatory cytokines and other factors that facilitate tumor escape, activity of regulatory T cells or Th2 type of cells. This technology can also be utilized to boost vaccine responses against cancers and chronic infectious diseases. Please contact John D. Hewes, Ph.D. at 240–276–5515 or john.hewes@nih.gov for more information.

**Dated:** September 17, 2015.

**Richard U. Rodriguez,**

**Acting Director, Office of Technology Transfer, National Institutes of Health.**

[FR Doc. 2015–24137 Filed 9–22–15; 8:45 am]

**BILLING CODE 4140–01–P**

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**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; Fellowships: Cell Biology, Developmental Biology, and Bioengineering.

**Date:** October 29–30, 2015.

**Time:** 8:00 a.m. to 6:00 p.m.

**Agenda:** To review and evaluate grant applications.

**Place:** Hyatt Regency Bethesda, One Bethesda Metro Center, 7400 Wisconsin Avenue, Bethesda, MD 20814.

**Contact Person:** Raj K. Krishnaraju, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 6190, Bethesda, MD 20892, 301–435–1047, kkrishna@csr.nih.gov.

**Name of Committee:** Molecular, Cellular and Developmental Neuroscience Integrated Review Group; Cellular and Molecular Biology of Glia Study Section.

**Date:** October 29–30, 2015.

**Time:** 8:00 a.m. to 2:00 p.m.

**Agenda:** To review and evaluate grant applications.

**Place:** Hilton McLean Tysons Corner, 7920 Jones Branch Drive, McLean, VA 22102.

**Contact Person:** Linda MacArthur, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 4187, Bethesda, MD 20892, 301–537–9986, macarthurb@csr.nih.gov.

**Name of Committee:** OncoG 1-Basic Translational Integrated Review Group; Tumor Progression and Metastasis Study Section.

**Date:** October 29–30, 2015.

**Time:** 8:00 a.m. to 6:00 p.m.

**Agenda:** To review and evaluate grant applications.

**Place:** Embassy Suites, DC Convention Center, 900 10 Street, Washington, DC 20001.

**Contact Person:** Rolf Jakobi, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 6187, MSC 7806, Bethesda, MD 20892, 301–495–1718, jakobi@cr.nih.gov.

**Name of Committee:** Center for Scientific Review Special Emphasis Panel; Topics in Bacterial Pathogenesis.

**Date:** October 29–30, 2015.

**Time:** 8:00 a.m. to 6:00 p.m.

**Agenda:** To review and evaluate grant applications.

**Place:** The Warwick Allerton Hotel, 701 North Michigan Avenue, Chicago, IL 60611.

**Contact Person:** Richard G. Kostriken, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 3192, MSC 7808, Bethesda, MD 20892, 240–519–7808, kostrikr@csr.nih.gov.

**Name of Committee:** Center for Scientific Review Special Emphasis Panel; Fellowships: Biophysical, Physiological, Pharmacological and Bioengineering Neuroscience.

**Date:** October 29–30, 2015.

**Time:** 8:00 a.m. to 5:00 p.m.

**Agenda:** To review and evaluate grant applications.

**Place:** The Westin Georgetown, 2350 M St. NW, Washington, DC 20037.