

Type of respondent	Estimated number of respondents	Frequency of response	Estimated number of responses	Average hours per response	Annual hour burden	Respondent cost**
Pilot RDD screener only	133	1	133	.0833	11	\$176
Pilot RDD screener and interview	200	2	400	*.2500	100	1,600
Additional RDD screeners for advance materials test	450	1	450	.0833	37	592
Pilot mail survey	640	1	640	.3333	213	3,408
RDD screener only	2,333	1	2,333	.0833	194	3,104
RDD screener and interview	3,500	2	7,000	*.2500	1,750	28,000
Mail survey	3,500	1	3,500	.3333	1,167	18,672
Telephone screener only for mail followup	457	1	457	.0833	38	608
Telephone screener and interview for mail followup	457	2	914	*.2500	229	3,664
Total	11,670		15,827		3,739	59,824

*(0.833 + 0.4167) / 2 = 0.2500.

** Hourly wage rate = \$16.00.

Request For Comments: Written comments and/or suggestions from the public and affected agencies are invited on one or more of the following points: (1) Whether the proposed collection of information is necessary for the proper performance of the function of the agency, including whether the information will have practical utility; (2) The accuracy of the agency's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (3) Ways to enhance the quality, utility, and clarity of the information to be collected; and (4) Ways to minimize the burden of the collection of information on those who are to respond, including the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology.

Direct Comments To OMB: Written comments and/or suggestions regarding the item(s) contained in this notice, especially regarding estimated public burden and associated response time, should be directed to the: Office of Management and Budget, Office of Regulatory Affairs, New Executive Office Building, Room 10235, Washington, DC 20503, *Attention:* Desk Office for NIH. To request more information on the proposed project or to obtain a copy of the data collection plans and instruments, contact Bradford W. Hesse, PhD, Project Officer, National Cancer Institute, NIH, EPN 4068, 6130 Executive Boulevard MSC 7365, Bethesda, Maryland 20892-7365, or call non-toll-free number 301-594-9904, or FAX your request to 301-480-2198, or E-mail your request, including your address, to heseb@mail.nih.gov.

Comments Due Date: Comments regarding this information collection are best assured of having their full effect if

received within 30-days of the date of this publication.

Dated: March 23, 2007.

Rachelle Ragland-Greene,

NCI Project Clearance Liaison, National Institutes of Health.

[FR Doc. E7-6064 Filed 3-30-07; 8:45 am]

BILLING CODE 4140-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Government-Owned Inventions; Availability for Licensing

AGENCY: National Institutes of Health, Public Health Service, 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. 207 to achieve expeditious commercialization of results of federally-funded research and development. Foreign patent applications are filed on selected inventions to extend market coverage for companies and may also be available for licensing.

ADDRESSES: 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.

High-Level Expression and Purification of Untagged and Histidine-Tagged Human Immunodeficiency Virus type-1 (HIV-1) Reverse Transcriptase

Description of Technology: This invention includes plasmids and protocols to express and purify large quantities of histidine-tagged and untagged HIV-1 reverse transcriptase (RT). Conditions have been optimized for overexpression and purification of p66 and p51 heterodimer RT in *E. coli*. High-level of expression was reached as RT represented approximately 30%–40% of total cell proteins. The subject invention enables the purification of large quantities of heterodimer RT necessary for structural and kinetic studies and facilitates subunit-specific amino acid alterations essential for structure/function investigations.

Applications: Research Tool.

Development Status: In vitro data available.

Inventors: Samuel H. Wilson, Rajendra Prasad, Esther W. Hou (NIEHS).

Related Publication: EW Hou, R Prasad, WA Beard, SH Wilson. High-level expression and purification of untagged and histidine-tagged HIV-1 reverse transcriptase. *Protein Expr Purif.* 2004 Mar;34(1):75–86.

Patent Status: HHS Reference No. E-141-2007/0—Research Tool.

Licensing Status: Available for non-exclusive licensing as biological material and research tool.

Licensing Contact: Sally Hu, PhD; 301/435-5606; HuS@mail.nih.gov.

Methods of Determining the Prognosis of an Adenocarcinoma

Description of Technology: Available for licensing and commercial development is a novel method for determining the prognosis of a subject with adenocarcinoma in an organ, such

as the lung, and to aid in the selection of a specific therapeutic regimen. Lung adenocarcinoma (AC) is the predominant histological subtype of lung cancer, which is the leading cause of cancer deaths worldwide. The risk of metastasis remains substantial in AC patients, even when a curative resection of early-stage AC is performed. The prognosis includes the determination of the likelihood of survival, the likelihood of metastasis, or both. The method includes quantization of the expression of a plurality of Th1 and Th2 cytokines of interest in the adenocarcinoma and in non-cancerous tissue in the organ. Altered expression of one or more of the Th1 and Th2 cytokines in the adenocarcinoma as compared to the non-cancerous tissue determines the prognosis for the subject. The method is capable of distinguishing patients with lymph node metastasis versus those with short term survival. Furthermore, methods are provided for evaluating the effectiveness of anti-cancer agents.

Applications: Prognosis of adenocarcinoma, aid in the selection of specific therapeutic regimens and evaluation of the effectiveness of anti-cancer agents.

Development Status: The technology is in early stage of development.

Inventors: Curtis C. Harris, Masahiro Seike, Xin Wei Wang (NCI).

Patent Status:

1. U.S. Provisional Application No. 60/830,936 filed 14 Jul 2006 (HHS Reference No. E-263-2006/0-US-01).

2. U.S. Provisional Application No. 60/885,101 filed 17 Jan 2007 (HHS Reference No. E-085-2007/0-US-01).

Licensing Status: Available for non-exclusive or exclusive licensing.

Licensing Contact: Cristina Thallhammer-Reyero, PhD, M.B.A.; 301/435-4507; thallhamc@mail.nih.gov.

Codon Optimized Genes for Subunit Vaccines

Description of Technology: Available for licensing from the NIH are gene constructs that express immunogenic proteins based on viral genes that have been optimized for expression in mammalian cells. Using vaccine vectors expressing respiratory syncytial virus (RSV) proteins from the optimized genes, this technology was shown to result in a potent RSV-specific cellular immune responses with favorable phenotypic patterns. Such optimized genes could be essential for development of an effective RSV subunit vaccine. Further, this optimization could have possible application to gene-based vectors for other viral vaccines.

Potential Applications of Technology: Vaccines; Improved protein expression.

Inventors: Barney S. Graham and Teresa R. Johnson (VRC/NIAID).

Patent Status: U.S. Provisional Application No. 60/872,071 filed 30 Nov 2006 (HHS Reference No. E-326-2006/0-US-01).

Licensing Status: Available for non-exclusive or exclusive licensing.

Licensing Contact: Susan Ano, PhD; 301/435-5515; anos@mail.nih.gov.

Dual Expression Vector for DNA Vaccines

Description of Technology: Available for licensing from the NIH is an expression vector for improved DNA vaccines. Activation of co-stimulatory molecules (e.g. signaling molecules, cytokines, chemokines) and the timing of the activation are important for adaptive immune response. This technology describes a new vector that expresses an antigen and a co-stimulatory molecule, the latter after a delay to allow accumulation of the antigen. It is known that in some circumstances an optimal immune response is achieved by administration of a co-stimulatory molecule after antigen delivery. The subject technology improves upon the existing concept by providing a vector for accomplishing this optimization in a single step. Exemplary animal studies have shown that the delayed expression of some co-stimulatory molecules in important signaling pathways resulted in enhancement of the cellular and/or humoral immune responses using HIV Env as a representative antigen.

Potential Applications: Improved DNA vaccines.

Inventors: Gary J. Nabel and Wataru Akahata (VRC/NIAID).

Patent Status: 1. U.S. Provisional Application No. 60/737,896 filed 18 Nov 2005 (HHS Reference No. E-043-2006/0-US-01).

2. PCT Application No. PCT/US2006/044552 filed 20 Nov 2006 (HHS Reference No. E-043-2006/2-PCT-01).

Licensing Status: Available for non-exclusive or exclusive licensing.

Licensing Contact: Susan Ano, PhD; 301/435-5515; anos@mail.nih.gov.

Methods and Systems for Efficient Analysis and Microdissection of Intact Biological Specimens

Description of Technology: Efficient and accurate analysis of intact biological specimens is needed to provide information related to a broad range of pathologies as well as normal physiological states. The available technology includes novel systems, methods, and platforms for selective

analysis of biological material such as whole cells, tissues and tumors. This platform may be used to identify and independently characterize specific components, such as cells, proteins, nucleic acids or other molecules that make-up the specimen.

The methods include placing the sample of interest on a surface such as a membrane, and activating the surface at selected sites adjacent to the section of interest. The activated sites become permeable and the cells or cell components adjacent to the permeable sites can then be selectively extracted and their content analyzed by standard biochemical procedures. In addition, the extract may be applied to microarray devices, such as cDNA arrays, for analysis of gene expression etc. The technique presents a convenient alternative to existing methods of tissue microdissection. For further convenience, the technique can be directly combined with a variety of analytical devices such as ELISAs, microarray biochips, or other devices which include multiple regions carrying multiple capture molecules.

Applications: Analysis of biological specimens such as whole cell tissues and tumors; High throughput analysis of individual components of intact biological specimens.

Inventors: Michael R. Emmert-Buck (NCI), Chad R. Englert-Haldeman (NCI), Robert F. Bonner (NICHD), and Lance A. Liotta (NCI).

Patent Status: 1. Patent Cooperation Treaty Application No. PCT/US01/08095 filed 14 Mar 2001, which published as WO 02/10751 on 07 Feb 2002; claiming priority to 26 Jul 2000 (HHS Reference No. E-197-2000/0-PCT-02).

2. National Phase Applications in:

a. U.S., Serial No. 10/333,374 filed 10 Jul 2003 (HHS Reference No. E-197-2000/0-US-03).

b. Canada, Serial No. 2415864 filed 14 Mar 2001 (HHS Reference No. E-197-2000/0-CA-04).

c. Europe, Serial No. 01918647.0 filed 14 Mar 2001 (HHS Reference No. E-197-2000/0-EP-05).

Licensing Status: Availability for non-exclusive or exclusive licensing.

Licensing Contact: Susan Ano, PhD; 301/435-5515; anos@mail.nih.gov.

Dated: March 26, 2007.

Steven M. Ferguson,

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

[FR Doc. E7-6066 Filed 3-30-07; 8:45 am]

BILLING CODE 4140-01-P