

Dated: March 2, 2006.

**Alvin Hall,**

Director, Management Analysis and Services  
Office, Centers for Disease Control and  
Prevention.

[FR Doc. E6-3261 Filed 3-7-06; 8:45 am]

BILLING CODE 4163-18-P

**DEPARTMENT OF HEALTH AND  
HUMAN SERVICES**

**Centers for Disease Control and  
Prevention**

**Government-Owned Inventions;  
Availability for Licensing and  
Cooperative Research and  
Development Agreements (CRADAs)**

**AGENCY:** Centers for Disease Control and  
Prevention Technology Transfer Office;  
Department of Health and Human  
Services.

**ACTION:** Notice.

**SUMMARY:** The invention named in this  
notice is owned by agencies of the  
United States Government and is  
available for licensing in the United  
States (U.S.) in accordance with 35  
U.S.C. 207, and is available for  
cooperative research and development  
agreements (CRADAs) in accordance  
with 15 U.S.C. 3710a, to achieve  
expeditious commercialization of  
results of federally funded research and  
development. A provisional patent  
application has been filed. A Patent  
Cooperation Treaty (PCT) application  
and national stage foreign patent  
applications claiming priority to the  
Patent Cooperation Treaty (PCT)  
application are expected to be filed  
within the appropriate deadlines to  
extend market coverage for U.S.  
companies and may also be available for  
licensing.

**ADDRESSES:** Licensing and CRADA  
information, and information related to  
the technology listed below, may be  
obtained by writing to Suzanne Seavello  
Shope, J.D., Technology Licensing and  
Marketing Scientist, Technology  
Transfer Office, Centers for Disease  
Control and Prevention (CDC), Mailstop  
K-79, 4770 Buford Highway, Atlanta,  
GA 30341, telephone (770)488-8613;  
facsimile (770)488-8615; or e-mail  
[sshope@cdc.gov](mailto:sshope@cdc.gov). A signed Confidential  
Disclosure Agreement (available under  
Forms at <http://www.cdc.gov/tto>) will be  
required to receive copies of  
unpublished patent applications and  
other information.

**Diagnostics**

*Immunoassay for Diagnosis of  
Orthopoxvirus Infection*

A CDC-developed immunoassay may  
be used for the diagnosis of infection  
with Orthopoxviruses (e.g. Monkeypox,  
Variola) by detection of acute phase  
immune responses that correlate to  
recent infection. With recent recognition  
of Orthopox viruses as emerging  
infectious agents with zoonotic  
transmission capabilities as well as  
select agents for bioterrorism, assays for  
the detection or diagnosis of infections  
are sought. This assay provides a rapid  
and simple method for detection of  
infection with these viruses related to  
zoonotic transmission or bioterrorism  
events involving such viruses.

Use of the assay produced high levels  
of sensitivity during the 2003  
Monkeypox outbreak in North America  
when compared to PCR.  
Commercialization of the ELISA test  
may provide a standard screening tool  
for diagnosis of Orthopoxvirus as well  
as a surveillance tool for exposure.

The immunoassay may also be useful  
at the state level for BT surveillance  
including an opportunity for use in  
reference labs. Reagents used in the  
assay are available through CDC  
laboratories and for commercial  
development of the assay. Further  
refinement of the assay may result in the  
development of additional reagents for  
incorporation into the assay.

*Inventors:* Kevin L. Karem, Inger K.  
Damon and Joanne L. Patton.  
*CDC Ref. #:* I-014-04.

**James D. Seligman,**

Chief Information Officer, Centers for Disease  
Control and Prevention.

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with 15 U.S.C. 3710a, to achieve  
expeditious commercialization of  
results of federally funded research and  
development. A provisional patent  
application has been filed. In addition,  
the invention is protected by copyright  
registration. A Patent Cooperation  
Treaty (PCT) application and national  
stage foreign patent applications  
claiming priority to the Patent  
Cooperation Treaty (PCT) application  
are expected to be filed within the  
appropriate deadlines to extend market  
coverage for U.S. companies and may  
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[sshope@cdc.gov](mailto:sshope@cdc.gov). A signed Confidential  
Disclosure Agreement (available under  
Forms at [www.cdc.gov/tto](http://www.cdc.gov/tto)) will be  
required to receive copies of  
unpublished patent applications and  
other information.

**Software**

*Computer Software for Automating  
Permeation Testing Data Analysis*

Data analysis for chemical protective  
clothing (CPC) permeation testing  
involves a number of equations and  
experimental factors. Experimenter bias  
and possible calculation errors are  
critical issues when determining  
permeation parameters. In order to  
compare results among different  
laboratories and manufacturers, the  
normalized breakthrough time is  
required since it is not dependent on the  
detection limits of the analytical system.  
However, calculating the normalized  
breakthrough time requires the use of  
polynomial curve fitting, polynomial  
derivatives, and quadratic equations.  
Solving these equations, without a  
computer program, would be very  
difficult. Therefore, a unique computer  
program using Microsoft Visual C++,  
referred to as "Permeation Calculator",  
has been developed at the National  
Institute for Occupational Safety and  
Health/National Personal Protective  
Technology Laboratory (NIOSH/NPPTL)  
to calculate the permeation parameters.  
The program imports data and then  
calculates the permeation parameters;