Modification Form for Permit BIO-RRI-0021

Permit Holder: Gregory Dekaban

Approved Personnel
(Please stroke out any personnel to be removed)

Bryan Au

Xizhong Zhang

Sonali deChickera

Christy Willert

John Barrett

Ryan Buensuceso

Additional Personnel
(Please list additional personnel here)

	Please stroke out any approved Biohazards to be removed below	**Write additional Biohazards for approval below. ***
Approved Microorganisms	E. coli (DH5 alpha), E. coli (Top 10)	
Approved Cells	Human (established), Rodent (established), Rodent (primary), HEK 293, HEK 293FT, CP1, CP2	
Approved Use of Human Source Material	Peripheral Blood Mononuclear cells (adult), Human cord blood cells, bone marrow	
Approved GMO	lentiviral vector backbone (Virapower), adenoirus-backed vectors (Ad 5), rAd5 GFP, proto-oncogene HER2/neu, wild type myxoma virus, recominant myxoma carry mRFP and or HER2/neu	wildtype raccoonpox virus, recombinant raccoonpox virus, vMyxgfp, vMyx135KO Ad My D88 CD40

** PLEASE ATTACH A MATERIAL SAFETY DATA SHEET OR EQUIVALENT FOR NEW BIOHAZARDS.*
*** PLEASE ATTACH A BRIEF DESCRIPTION OF THE WORK THAT EXPLAINS THE BIOHAZARDS USED AND HOW THEY WILL BE USED.*

Classification: 2+

Date of last Biohazardous Agents Registry Form: May 29, 2007

Signature of Permit Holder:

BioSafety Officer(s):

Chair,Biohazards Subcommittee:

Modification Form for Permit BIO-RRI-0021

Permit Holder: Gregory Dekaban

Approved use of Animals	rodent C57B1/6, NOD SCID, GFP	
Approved Toxin(s)		

* *PLEASE ATTACH A MATERIAL SAFETY DATA SHEET OR EQUIVALENT FOR NEW BIOHAZARDS.*
** *PLEASE ATTACH A BRIEF DESCRIPTION OF THE WORK THAT EXPLAINS THE BIOHAZARDS USED AND HOW THEY WILL BE USED.*

Classification: 2+

Date of last Biohazardous Agents Registry Form: May 29, 2007

Signature of Permit Holder: _____

BioSafety Officer(s): _____

Chair,Biohazards Subcommittee: _____

Raccoonpox virus is considered to be raccoon-specific. It is not harmful or pathogenic to humans. We are planning to use this virus in various mammalian cell lines to study host range and immune evasion strategies. As well, we are planning to sequence the genome. Towards this goal we will be growing the virus to high titres (10^8-10^9 pfu/ml) and also producing recombinant versions in which we will add fluorescent tags (EGFP and RFP) so that we can monitor infection. We know that this virus will grow in common monkey cells including Vero and BGMK cells and in rabbit kidney cells (RK13).

DeKaban

>> -----Original Message-----
>> From: Permit-Permis [mailto:permitpermis@phac-aspc.gc.ca]
>> Sent: August 14, 2009 1:31 PM
>> To: dekaban
>> Subject: Raccoonpox virus
>>
>> Dear Dr. Gregory A. Dekaban
>>
>> We do not regulate Raccoonpox virus as we consider this as a RG1, please contact your Canadian distributor and they will send you this pathogens without any documentation required form us.
>>
>> Regards
>>
>> Josee Davies
>> A/Regulatory Technologist/ technologiste en réglementation
>> Office of Laboratory Security/Bureau de la sécurité des laboratoires
>> Public Health Agency of Canada/ Agence de santé publique du Canada
>> 100 ch. Colonnade Rd. AL: 6201A Ottawa, Ontario, Canada K1A 0K9
>> Tel: (613) 957-1779
>> Fax: (613)941-0596
>>
>> **********Register Now**********
>>
>> A new law passed by Parliament requires all persons responsible for
>> human
>> pathogens of risk group 2, 3 or 4 or toxins on Schedule 1 of the Act to
>> register their laboratory or facility before midnight of September 21,
>> 2009. You must register, even if you already hold an Import Permit or
>> Compliance or Certification Letter. The registration website can be
>> found
>> at: http://www.phac-aspc.gc.ca/ols-bsl/pathogen/register-eng.php.
>> Please
>> submit both Form 1 and 2.
>>
>> *******Enregistrez-vous dès maintenant********
>>
>> Une nouvelle loi adoptée par le Parlement exige que toutes les personnes
>> qui sont responsables d'agents pathogènes humains de groupe de risque
>> 2, 3
>> ou 4 ou des toxines dans l'annexe 1 inscrivent leur laboratoire ou leur
>> établissement avant minuit le 21 septembre 2009. Vous devez vous
>> enregistrer, même si vous détenez un permis d'importation, une lettre de
>> conformité ou de certification. L'information concernant
>> l'inscription se
>> retrouve à l'adresse suivante :
>> http://www.phac-aspc.gc.ca/ols-bsl/pathogen/register-fra.php. S'il vous
>> plaît, assurez-vous de soumettre le Formulaire 1 et le Formulaire 2.
>>

ATCC
The Global Bioresource Center™

Search Catalog
Select a Category
Go

Login Search Options

About | Cultures and Products | Science | Standards | Deposit Services | Custom Services | Product Use Policy

ATCC Advanced Catalog Search » Product Details

Product Description

Before submitting an order you will be asked to read and accept the terms and conditions of ATCC's Material Transfer Agreement or, in certain cases, an MTA specified by the depositing institution.

Customers in Europe, Australia, Canada, China, Hong Kong, India, Japan, Korea, Macau, Mexico, New Zealand, Singapore, and Taiwan, R.O.C. must contact a local distributor for pricing information and to place an order for ATCC cultures and products.

Print this Page

Animal Viruses and Antisera

ATCC® Number: VR-838™ Order this Item **Price:** $325.00

Classification: Poxviridae, Orthopoxvirus

Agent: Raccoonpox virus deposited as Raccoonpox virus, Orthopoxvirus

Strain: Herman

Original Source: Isolated by Y.F. Herman from respiratory tract of raccoon with no clinical symptoms, Maryland, USA, 1964

Depositors: JH Nakano

Biosafety Level: 2 **Shipped:** frozen

Permits/Forms: In addition to the MTA mentioned above, other ATCC and/or regulatory permits may be required for the transfer of this ATCC material. Anyone purchasing ATCC material is ultimately responsible for obtaining the permits. Please click here for information regarding the specific requirements for shipment to your location.

Related Products

Host Organism : LLC-MK2 cells (ATCC CCL-7); Vero (ATCC CCL-81)
Vero cells (ATCC CCL-81); LLC-MK2 cells (ATCC CCL-7); CAM; suckling mouse

Incubation : **Temperature:** 35.5°C
Duration: 3-4 days

Effect : Yes, in vitro effects: Cytopathic effects (large plaques and giant cell syncytia) in Vero cells
Yes, in vivo effects: paralysis in suckling mouse
Yes, in vivo effects: pinpoint pocks on chorioallantoic membranes

Comments : The virus does not grow well on chorioallantoic membranes after 2 to 3 passages.
The virus cross-reacts serologically with the Connaught strain of vaccinia virus.
Raccoons inoculated with RPV show no apparent clinical symptoms; however, their sera demonstrate strong HAI reactions.
Does not grow well on CAM after 2-3 passages. Cross-reacts with Connaught strain on vaccinia virus. Raccoons inoculated with RPV show no apparent clinical symptoms; however, their sera demonstrate strong HAI reactions.

References : 33948: Thomas EK, et al. Further characterization of Raccoonpox virus. Arch. Virol. 49: 217-222, 1975. PubMed: 813616
33949: Herman YF. Bact. Proc. 64th Annual Meeting, ASM : 117, 1964.

Return to Top

Notices and Disclaimers

ATCC products are intended for laboratory research purposes only, unless noted otherwise. They are not intended for use in humans.

While ATCC uses reasonable efforts to include accurate and up-to-date information on this site, ATCC makes no warranties or representations as to its accuracy. Citations from scientific literature and patents are provided for informational purposes only. ATCC does not warrant that such information has been confirmed to be accurate.

All prices are listed in U.S. dollars and are subject to change without notice. A discount off the current list price will be applied to most cultures for nonprofit institutions in the United States. Cultures that are ordered as test tubes or flasks will carry an additional laboratory fee. Fees for permits, shipping, and handling may apply.

Back to my Search

Modification Form for Permit BIO-RRI-0021

Permit Holder: Gregory Dekaban

Approved Personnel
(Please stroke out any personnel to be removed)

~~Phillippe Alexander Gilbert~~

~~Carmen Simedrea~~

Additional Personnel
(Please list additional personnel here)

Christy Willert
Sonali dechickova
Xizhang Zhang
John Barrett
Bryan Au, Juno Jimenez

	Please stroke out any approved Biohazards to be removed below	Write additional Biohazards for approval below. *
Approved Microorganisms	E. coli (DH5 alpha), E. coli (Top 10)	
Approved Cells	Human (established), Rodent (established), Rodent (primary), HEK 293, HEK 293FT, CP1, CP2	
Approved Use of Human Source Material	Peripheral Blood Mononuclear cells (adult), Human cord blood cells, bone marrow	
Approved GMO	lentiviral vector backbone (Virapower), adenoirus-backed vectors (Ad 5), rAd5 GFP, proto-oncogene HER2/neu	wild type myxoma virus, recombinant myxoma carry an RFP and/or HER2/neu
Approved use of Animals	rodent C57B1/6, NOD SCID, GFP	

** PLEASE ATTACH A MATERIAL SAFETY DATA SHEET OR EQUIVALENT FOR NEW BIOHAZARDS.*
*** PLEASE ATTACH A BRIEF DESCRIPTION OF THE WORK THAT EXPLAINS THE BIOHAZARDS USED AND HOW THEY WILL BE USED.*

Date of last Biohazardous Agents Registry Form May 29, 2007

Signature of Permit Holder: Gregory A. Dekaban

BioSafety Officer(s): Terrill Woodside Stanley Roberts

Chair,Biohazards Subcommittee: G.M. Kidder

RECEIVED
16/3/14/07
KW

BIO-RRI-0021 (2+)
BIO-UWO-0031 (3)

robarts RESEARCH

BIOHAZARDOUS AGENTS REGISTRY FORM

Reviewed by Biosafety Subcommittee: February 2006

This form must be completed by each Principal Investigator when completing a grant application or grant renewal to be administered by the Robarts Research Institute, if the use of biohazardous and/or infectious agents is proposed. For any proposed animal work involving the use of biohazardous agents or animals carrying zoonotic agents infectious to humans, this form must also be completed.

COMPLETED FORMS ARE TO BE RETURNED TO BIOSAFETY SUBCOMMITTEE CHAIR, ROOM 3-34.1.

If there are any changes to the information on these forms (excluding grant title and funding agencies) a new form must be completed and sent to the Biosafety Subcommittee Chair BEFORE implementation of these changes can occur.

If multi-team grants are being applied for, each individual Investigator of the team must submit a Biohazardous Agents Registry Form to the Biosafety Subcommittee Chair.

Containment Levels will be required in accordance with Health Canada (HC), Laboratory Biosafety Guidelines, 3rd edition 2004, or Canadian Food Inspection Agency (CFIA), Containment Standards for Veterinary Facilities, 1st edition 1996.

For questions regarding this form, please contact Biosafety Subcommittee Chair at ext. 34125.

1.0 Contact Information

PRINCIPAL INVESTIGATOR: Gregory A. Dekaban

SIGNATURE: Gregory A Dekaban

DATE: March 19, 2007

DEPARTMENT: BTRG

ADDRESS: Rm 2-12 Robarts

TELEPHONE: x 34241

EMAIL: dekaban@robarts.ca

Location of experimental work to be carried out:

Building(s): Robarts

Room(s): Rm 2-12, external barrier, BTRT facility Rm 2-22

For work being performed at institutions affiliated with the Robarts Research Institute, the Safety Officer for the Institution where experiments will take place must sign the form prior to it being sent to Robarts Research Institute, Biosafety Subcommittee Chair. See Section 13.0, Approvals

/

GRANT TITLE(S):

ATTACH A BRIEF DESCRIPTION OF YOUR WORK, SUCH AS THE RESEARCH GRANT SUMMARY(S) EXPLAINING THE BIOHAZARD(S) USED.

FUNDING AGENCY/AGENCIES: Ontario Cancer Research Network

Anticipated Grant End Date: March 2010

Names of all personnel working under Principal Investigator's supervision in this location:

Greg DeKaban	Student to be named
Peter O'Connell	
Paula Porter	
Carmen Simedrea	
Tim Su	
Matilda Lean Parte	
Jonathan Snir	

Note : A list of human pathogens categorized according to Risk Group can be obtained by calling the Office of Laboratory Security directly at (613) 957-1779 or accessing their Web site : http://www.phac-aspc.gc.ca/pls-bsl/index.html

2.0 Microorganisms

2.1 Does your work involve the use of microorganisms? YES O NO O
If NO, please proceed to Section 3.0

2.2 Please complete the table below.

Name of Microorganism	Is microorganism a known human pathogen? YES/NO	Is microorganism a known animal pathogen? YES/NO	Is microorganism a known zoonotic agent? YES/NO	Maximum quantity to be cultured at one time?	Health Canada or CFIA Containment Level (select one)
E. coli DH5α Top10 stable 2	No	No	No	1-2L	1O 2O 3O
					1O 2O 3O
					1O 2O 3O

3.0 Cell Culture

3.1 Does your work involve the use of cell cultures? YES ✓ NO ○
If NO, please proceed to Section **4.0**.

3.2 Please indicate in the table below the type of cells that will be grown in culture.

Cell Type	Is this cell type used in your work? YES / NO	Established or Primary *	Supplier of Primary Cell Culture Tissue
Human	Yes	Both	Q. Jacque Galipeau;
Rodent	Yes (mouse)	Both	
Non-human primate			
Other (specify)			

* i.e. derived from fresh tissue

3.3 Complete the following table.

Specific Cell Line	Source / Supplier	HC or CFIA Containment Level (select one)
HEK 293, HEK293FT	Invitrogen or have already	1 ○ 2 ✓ 3 ○
CP.1* or CP.2*	D. Rod Nelson, U. B.C	1 ○ 2 ✓ 3 ○
		1 ○ 2 ○ 3 ○

* Express proto-oncogene Her2/neu and a dominant negative form of p53

4.0 Use of Human Source Materials

4.1 Does your work involve the use of human source materials? YES ✓ NO ○
If NO, please proceed to Section **5.0**

4.2 Indicate in the table below the Human Source Material to be used.

Human Source Material	Specify Source, or Not Applicable (NA)	Is Human Source Material known to be infected with an infectious agent? YES/NO	Name of Infectious Agent	HC or CFIA Containment Level (select one)
Human Blood (whole) or other Body Fluid				1 ○ 2 ○ 3 ○
Human Blood (fraction) or other Body Fluid	PBMC (adult) cord blood cells bone marrow *	NO		1 ○ 2 ✓ 3 ○
Human Organs (unpreserved)				1 ○ 2 ○ 3 ○
Human Tissues (unpreserved)				1 ○ 2 ○ 3 ○

* To be supplied by Dr. Jacque Galipeau (Montreal) or via David Hess as part of a collaboration.

5.0 Genetically Modified Organisms and Cell lines

5.1 Will genetic modifications be made to the organism, virus or cell line? YES ✓ NO ○
If NO, please proceed to Section 6.0

5.2 Will genetic sequences from any of the following be involved?

- HIV YES ✓ NO ○

If YES, specify: lentiviral vector backbone

- HTLV 1 or 2 YES ○ NO ✓

If YES, specify: _____

- Other human or animal pathogen and/or their toxins YES ○ NO ✓

If YES, specify: _____

5.2 Will intact genetic sequences be used from:

- SV 40 Large T antigen YES ○ NO ○
- Adeno E1A YES ✓ NO ○
- Known or suspected oncogenes YES ○ NO ○

If YES, specify: oncogenic dominant negative form of p53

5.4 Will a live vector(s) (viral or bacterial) be used for gene transduction? YES ✓ NO ○

If YES, name vector: Adenovirus-based vectors; lentiviral vector

5.5 List specific vector(s) to be used: Virapower lentiviral vector, Ad5

5.6 Will vector be replication defective? YES ✓ NO ○

5.7 Will vector be infectious to humans or animals? YES ✓ NO ○

5.8 Will this be expected to increase the Containment Level required? YES ✓ NO ○
2 of 3

6.0 Human Gene Therapy Trials

6.1 Will human clinical trials using the vector(s) in 5.5 be conducted? YES ○ NO ✓
If NO, please proceed to Section 7.0
If YES, attach a full description of the make-up of the virus.

6.2 Will vector be able to replicate in the host? YES ○ NO ○

6.3 How will the vector be administered? _____

6.4 Please give the Health Care Facility where the clinical trial will be conducted:

_____

6.5 Has human ethics approval been obtained? YES ○ NO ○

Approval # _____

7.0 Animal Experiments

7.1 Will any of the agents listed be used in live animals? YES ⊘ NO ○ under Peter O'Connell.
If NO, please proceed to section 8.0

7.2 Name of animal species to be used: C57Bl/6 ; NOD SCID

7.3 AUS protocol # pending and 2006-118-10 for expt's not involving virus vectors and NOD SCID mice.

7.4 If using murine cell lines, have they been tested for murine pathogens? YES ○ NO ⊘

8.0 Use of Animal species with Zoonotic Hazards

8.1 Will any of the following animals or their organs, tissues, lavages or other bodily fluids including blood be used?

- Pound source dogs YES ○ NO ⊘
- Pound source cats YES ○ NO ⊘
- Sheep or goats YES ○ NO ⊘
- Non- Human Primates YES ○ NO ⊘

If YES specify species _____

- Wild caught animals YES ○ NO ○

If YES specify species _____

9.0 Biological Toxins

9.1 Will toxins of biological origin be used? YES ○ NO ⊘
If NO, please proceed to Section 10.0
If YES, please name the toxin _____

9.2 What is the LD_{50} (specify species) of the toxin? _____

10.0 Import Requirements

10.1 Will the agent be imported? YES ○ NO ⊘
If NO, please proceed to Section 11.0
If YES, country of origin _____

10.2 Has an Import Permit been obtained from HC for human pathogens? YES ○ NO ○

10.3 Has an import permit been obtained from CFIA for animal pathogens? YES ○ NO ○

10.4 Has the import permit been sent to Biosafety Subcommittee Chair? YES ○ NO ○

If YES, Permit # _____

11.0 Training Requirements for Personnel Named on Form

All personnel named in section 1.0 of this form who will be using any of the above named agents are required to attend the following training courses given by OH&S.

- Biosafety
- Laboratory and Environmental/Waste Management Safety
- WHMIS

As the Principal Investigator, I have ensured that all of the personnel named on the form who will be using any of the biohazardous agents in Sections 2.0 to 10.0 have been trained as required.

SIGNATURE [signature]

12.0 Containment Levels

12.1 For the work described in sections **2.0** to **10.0**, select the highest HC or CFIA Containment Level required. 1○ 2● + 3●

12.2 Has the facility been certified by Biosafety Subcommittee Chair for this level of containment?

YES ● NO ○

If YES, give date: June 26, 2006 and permit number: 2006-06 (2-12.2)

13.0 Approvals

~~Robarts Research Institute~~ UWO Biohazards Subcommittee

Signature G.M. Cahill Date 29 May '07

Biosafety Officer for the Institution where experiments will take place

Signature [signature] Date May 28/07

Biosafety Officer of Robarts Research Institute (if different than above)

Signature _____ Date _____

Note: This permit will be in effect from _____ to _____,

subject to annual facility re-certification.