NE	V SUBMISSION PROCEDURE: PLEASE		DNS AT THE BOTTO		E Revision: 100601
R INSI	OBERT WOOD JOHNSON MEDICAL SCHOOL FUTIONAL ANIMAL CARE AND USE COMMITTEE	This application may be signed electronically. We suggest that PI's SAVE A COPY of the completed application BEFORE adding their electronic signature, as once the form is electronically signed, it will			FOR IACUC USE ONLY
		be locked from furthe	r changes. Saving the cor	npleted application	TRACKING NO.
		modifications are nee	ded in the future. The IAC	CUC application	CATEGORY
	Do not exceed length restrictions provided.	instructions may be for http://rwjms.umdnj.ed	ound on the Research Off u/research/orsp/ra/iacuc_	ice website below: forms.html	IACUC NO:
The	information in this application is confidential. Please	do not circulate outsid	e UMDNJ-Robert Wood J	ohnson Medical Schoo	DI.
	1.1 TITLE OF PROJECT				
	1.2 PRINCIPAL INVESTIGATOR INFORMATION		122 DEGREE(S)	123	ACADEMIC TITLE
			1.2.2 DEGREE(0)	1.2.0	
Е				CC (Dears Number Ct	reat City State Zin Cade)
INNC	1.2.4 DEPARIMENT		1.2.5 MAILING ADDRE	SS (Room Number, St	reet, City, State, Zip Code)
ERS			_		
÷.	1.2.6 CAMPUS				
NOI					
SECT	1.2.7 TELEPHONE AND FAX (Area Code, Number	and Extension)	1.2.8 EMAIL		
0,					
	1.3 GRANT TITLE (IF APPLICABLE)				
	1.4 FUNDING SOURCE				
	2.1 SPECIES Type:		Weight:		
	Strain:		Age:		
	Sex:				
	2.2 SUM TOTAL OF ANIMALS USED PER YEAR:		ESTIMATED NUMBER	OF ANIMALS USED F	PER 3 YEARS:
	2.3 REQUESTED SOURCE OF ANIMALS (Optional	/)			
ALS	2.4 SURGICALLY MODIFIED ANIMALS PURCHAS	ED Ves	If yes, list the source	and provide a protoco	approval letter from the source
ANIM			Source:		
N 2: /					
TIO	MEB Piscataway Camden		CHINJ Other	(Specify):	
SEC	2.6 List all Room Numbers (Laboratories, etc.) outsid	de the Vivarium where	animal procedures will be	performed and specify	y which procedures will be

1. Completed applications MUST be returned via SECURE e-mail VIA 'https://lift.umdnj.edu' ONLY to 'iacuc-rwjms@umdnj.edu' OR the application can be hand delivered on a CD or a flash drive to the ORSP.

Paper submissions will no longer be accepted.
 IACUC Policies may be found here: 'http://rwjms.umdnj.edu/research/orsp/ra/iacuc\_policies.html'

Institutional Animal Care and Use Committee Robert Wood Johnson Medical School Office of Research and Sponsored Programs 675 Hoes Lane, Room R-109 Piscataway, NJ 08854

	2.7 Are animals held outside of the Vivarium?	Yes No	
NUED)	If yes, please indicate the room locations(s) and the animals outside of the Vivarium or other authorized	he length of time that the animals will be housed outsided satellite facility for periods exceeding 12 hours, plea	de of the Vivarium. (If you need to house your se consult with the veterinarian.)
TIU	ROOM NUMBER	DURATION	FREQUENCY (Occurences per Week)
CON			
N 2 (			
стю			
SE	2.8 REQUEST FOR SPECIAL HOUSING (Option	al)	
	3.0 Briefly state the background for the project inc research. ( <i>Confine to the space provided</i> ):	cluding hypothesis to be tested, rationale, objectives, p	ublic health significance or importance of the
н			
ARC			
RESE			
OF F			
ACT			
<b>3</b> STR			
3: AE			
lion			
SECT			
•,			

4.0 Describe, in *lay language*, and sequence, in chronological order, *exactly* what will be done to the animals (*Detailed descriptions of in-vitro fertilization methodologies are unnecessary*):

If required, please use the continuation pages 13 & 14

	NOTE: Use of Hazardous Agents requires REHS. Links to some of the standard form (http://rwjms.umdnj.edu/research/orsp/ra/) approval will be given only after the use of	approval from the R s or respective webs All forms or letters <b>all</b> hazardous agen	WJMS Institutional sites are available u should be submitted ts has been approv	Biosafety C Inder the "R d simultaned ed.	ommittee, Labora equired Approvals ously to expedite t	tory Safety C s" section of t he process o	committe he ORS f approv	e, EOHS P website al. Final	S, and e IACUC	/or
	5.1 Does the use of chemicals or other agents pose potential danger to humans?									
	5.2 Can the natural or experimental disease or pathological condition under study be transmitted to humans?									
	If <b>yes</b> , what precautions are you taking to	protect persons who	will come into cont	act with the	se animals?					
	5.3 Indicate if your study involves any of the following. If any of these items are checked, clearance from the RWJMS Institutional Biosafety Committee, RWJMS Laboratory Safety Committee, EOHSS, and/or REHS is necessary. Please fill out the appropriate form and submit to the Office of Research and Sponsored Programs or the appropriate agency/committee.									
	Office of Research and Sponsored Progra 675 Hoes Lane, Room R-109 Piscataway, NJ 08854	ms Ph: 732-235-4 Fx: 732-235-5 E: iacuc-rwjm	Ph: 732-235-4338 Coriell Fx: 732-235-5534 Camd E: jacuc-rwims@umdni.edu		Institute for Medical Research Iddon Avenue In, NJ 08103		Ph: 856-757-2570 Fx: 856-968-9563 F: melerape@umdni.edu		du	
	Acute Toxins	Hu	man and/or Animal	Pathogens		Radioactive	Materia	ls		
	Chemical Carcinogens	- Hui	man Cells			Recombinar	nt DNA/F	RNA		
	Genetically Modified Cells					Other (Spec	;ify):			
	Identify agent(s) listed above:									
ENTS	AGENT:	DOSE:	ROUTE		FREQUENCY:		DURATION			
s age										
DOU										
AZAR										
5: H/										
TION										
SEC										
	5.4 Describe the potential health problems	to humans and/or a	nimals:							

	5.5 Describe special animal care required for biohazards and chemical hazards:
(CONTINUED)	
SECTION 5: HAZARDOUS AGENT	5.6 Describe special precautions for all personnel potentially exposed to this hazard:
	5.7 Will tumor cells, stem cells, tissue, sera, or other biological specimens from either animal or human sources be used as part of this protocol? Yes No List the supplier
	Human Animal ( <i>if animal</i> is selected, list the species. Reference Policy #6):
N	6.1 Breeding (Reference Policy #2)       Yes       No         If yes is genotyping required?       Yes       No         If yes answer questions 6.2 AND 6.3
3/IDENTIFICATIO	6.2 Genotyping (Reference Policy #8)       Tail Sample (specify length)         Other (specify tissue):       Other (specify tissue):         Age at tail clipping (specify):
6: GENOTYPING	6.3 Identification       Ear Tag:         Ear Punch       Tattoo         Oraclip (Provide justification, see Policy #9)
SECTION	
	This section intentionally left blank.

	7.1 \	What is the rationale for using animals in this study? (Check all that apply):				
		This research requires behavioral measurements from living animals.				
		This research requires biological measurements of tissue samples from living animals.				
		Animals are used to provide primary cell cultures.				
		Computer or other models cannot be used to replace animals in this research.				
		This research can not be done in-vitro.				
		Other ( <i>Explain</i> ):				
	7.2	Explain why the species used is appropriate for your research. ( <i>Check all that apply</i> ):				
		This research is a direct extension of previous work done on this species.				
		This research seeks to extend previous findings from other species specifically to this species.				
Į		Nothing is known about the physiological/behavioral phenomena of interest in this species.				
More is known about related aspects of the physiological/behavioral phenomena of interest in this species than any other.						
202		This species represents the best compromise between the simplest (lowest) organism that can be used and the most recent relevant model				
		system for human physiology/behavior.				
IVIAL		Other ( <i>Explain</i> ):				
NY.						
5						
วิ						
	73.	Justify the appropriateness of the number of animals to be used. (Check all that apply):				
		This number represents the lowest number needed for statistically significant tests of the hypothesis				
		A large number of physiological/behavioral parameters need to be measured in parametric fashion				
		A significant number of animals to be used are for breeding purposes				
		The physiological/behavioral parameters to be measured exhibit greater variablility, thus requiring a larger number of animals for statistically				
		significant analysis.				
		Meaningful data cannot be obtained from every animal used due to technical reasons, thus requiring a larger number of animals.				
		Other ( <i>Explain</i> ):				

	8.0 Define the number of animals to be used <b>p</b> provide committee members with an under	8.0 Define the number of animals to be used per year. (Outline experiments in as much detail as needed. Append a table or flow chart if necessary to provide committee members with an understanding of the experimental design.)							
AALS	In your description of your experimental de Number of Animals per Group and the Nun from the total number of animals per projec	In your description of your experimental design please indicate, where appropriate, the Experiment Title, the Number of Groups per Experiment, the Number of Animals per Group and the Number of Animals per Experiment. Please distinguish between the number of animals to be used per year from the total number of animals per project. Please use continuation pages 13 & 14 if necessary.							
- ANI	To help the commitee understand your inte protocol.	ntions, please provide an estimate	e of the total number of animals to be used over	the 3-year life of this					
SECTION 8: TOTAI									
	Sum total of animals used per year:		Estimated number of animals used per 3 yea	Irs:					
ITAL	9.0 Experimental use of animals: conditions that	at may create pain or distress (Ch	eck <b>every</b> appropriate category):						
MEN	Non-treated animals, euthanasia followed	by tissue harvest. (If only this iter	m is checked, proceed to SECTION 10.2).						
XPERI	Animals will suffer no pain or distress great analgesics, or tranquilizers.	ater than that induced by routine ir	njections or venipuncture and, therefore, will rec	eive no anesthetics,					
9: E	Animals will receive anesthetics, analgesi	cs, and/or tranquilizers to minimize	e or alleviate pain or distress during (check all the	nat apply):					
LION	Non-surgical procedures	Survival su	Irgery Non-surviv	al surgery					
SECI	Animals may experience pain or distress greater than induced by routine administrations or venipuncture, but will not receive anesthetics, analges or tranquilizers, since this will adversely affect the study. ( <b>MUST</b> answer SECTION 11.11 if this category is checked).								
	10.1 You must investigate alternatives to proce proposed research is not unnecessarily du more than slight or momentary pain and/o Complete Adjuvant use, and extensive irra available. The search or method should a cannot be eliminated.	dures likely to cause pain or distra uplicative [A painful procedure is or r distress in a human being to whi adiation.] List the sources or meth ilso include concepts of refinemer	ess. In addition this investigation should provide defined as any procedure that would reasonably ich the procedure is applied. Examples: termin hods that were used to determine that non-painf nt and reduction in order to minimize animal pair	evidence that the be expected to cause al surgery, Freund's ul alternatives are not and distress when it					
	Literature Search was conducted. Name	e of database searched: AL	TBIB AltWeb PubMed Other	Specify)					
S	Date	of search:							
TIVE	Year	s covered by search:							
ALTERNA	Keyw The v used	vords or search strategies used: word "Alternative" <b>MUST</b> be as one of the search terms.							
IVE /	The Animal Welfare Information Center of	the National Agricultural Library v	was consulted.						
IGAT	Recognized experts in the field were cons	ulted (give name and affiliation):							
/EST	NAME	AFFIL	IATION						
NI :C									
0N 10									
CTIC									
SE	Other ( <i>Explain</i> ):								
	10.2 I certify that this proposed research is not	unnecessarily duplicative and I ha	ave completed a database search						
	Name of database search:								
	Date of search: Key (do	Keywords or search strategies used (do not duplicate the keywords used in Section 10.1):							

11.2 Describe the non-surgical procedure, drug or cell injections, behavioral testing, blood sampling, etc.

11.3 Are you using an and	esthetic, ana	lgesic, or tranquilizer f	or a non-surgical	r a non-surgical procedure?			Yes No		
DRUG DOSE(S) r		S) mg/Kg ROUTE			FREQUENCY		PERSON ADMI	NISTERING	
-									
11.4 Are you collecting flu	uids?						Yes	No	
FLUID/SITE		VOLUME PER SAM	MPLE NUMBER OF		F SAMPLES FREQUEN		ICY OF SAMPLING SCHEDULE		
11.5 List the substances of	or drugs you	are administering (Oth	her than those list	ted in 11.3).	1		1		
DRUG/SUBSTANCE	DOSE(	S) mg/Kg	ROUTE		FREQUENCY		DURATION OF	TREATMENT	

	11.6 Will adjuvants be used for	or antibody production	? (Use of Freunds adjuvant	t <b>must</b> be jus	stified, please	e use continuation pa	ges).	Y	es	No
		INITIAL			BOOSTERS	3				
	ANTIGEN:									
	ADJUVANT:									
	INJECTION ROUTE:									
	INJECTION SITE:									
	VOLUME/SITE:									
	TOTAL VOLUME:									
	INJECTION SCHEDULE:									
	11.7 Will conscious animals v	will be restrained for o	oservation?					Yes		No
	NOTE: Short term (minutes) a procedures need not be desc	and skillful restrain of sribed.	animals for purposes of obs	ervation, phy	ysical examin	ation, or routine non	-stress	ful exper	rimental	
	FREQUENCY		DURATION			METHOD OF RES	FRAIN	Т		
ED)										
TINU										
CON										
URGICAL PROCEDUR	If <b>yes</b> , specify how long ani to onset of clinical signs). If 11.9 Could this condition ultin	imals will be maintaine death is the endpoint	ed in that state and the meth , justify and list the alternativ	nods to minin ves you have	nize or elimin e considered.	ate eventual pain or	distres	s (e.g. e Yes	uthanasia	a prior
N-SI	Will animals be euthanized	at the onset of clinica	I signs of pain or distress?					Yes		No
1: NC	If <b>yes</b> , describe the clinical	signs that will be mon	itored, the frequency of mor	nitoring, and	the criteria fo	or euthanization.				
ION 1	Tumors (See Policies #	±1, #3 and #5)	EAE (See Policy #7)	Ot	ther Diseases	S/Conditions (See Po	licies #	#3 and #	5) Specif	y Below
SECT										
	11.10 Will animals be subjec temperature extremes If <b>yes</b> , describe the stress,	eted to stressful physic s, or shock. the level, and the freq	al conditions such as high in uency:	ntensity nois	e, water imm	ersion,		Yes		No
	11.11 Will you be performing	a procedure which ma	ay result in unalleviated pair	n or distress?	?			Yes		No
	If <b>yes</b> , will you administer a	inalgesics or tranquiliz	ation?					Yes		No
	If <b>no</b> , please provide the sc	sientific justification for	omitting analgesia or tranq	uilization (Se	e Section 9.0	))				
	11.12 Will dietary manipulatio	ons be performed?						Yes		No
	If <b>yes</b> , describe the dietary	y manipulation.								

	12.1 Will a non-survival procedure be performed on an anesthetized animal which will not awaken from anesthesia? Yes No											
	If <b>no</b> , proceed to section 13											
AL)	12.2 List the person(s) performing the surgery:											
IRVIV												
N-SL												
ON) (	12.3 Describe the procedure(s). (Plea	se use continuation pages 13 & 14 if ne	cessary):									
URES												
CEDI												
PRO												
ICAL												
URG	12.4 Specify the following:		DOUTE									
12: S	ANESTHETIC	DOSE(S) mg/Kg	ROUTE	PERSON	ADMINISTER	RING						
NOI												
SECI												
	Asoptic tochnique includes wearing of		d facomasks: and roquires the use of s	torilo instru	monte and as	ontic						
	preparation of the surgical field. Survival surgery on rodents does not require a special facility, but should be performed using sterile instruments, surgical doves, and asentic procedures to prevent clinical infections. (For rodent asentic surgery reference Policy #10):											
	gioves, and aseptic procedures to prevent clinical infections. (For fodent aseptic surgery reference Policy #10):											
	13.2 Will the animals be subject to mo	]]	Yes		No							
	13.3 Person(s) performing the surgery:											
RES												
EDU	13.4 Describe the surgical procedure(	s). (Please use continuation pages 13 &	14 if necessary):									
ROC												
SAL P												
RGIC												
3: SU												
ON 1	13.5 Describe the anesthesia, analgesia, or sedatives used for the surgical procedure(s) Reference Policy #10 & #11:											
ECTI	ANESTHETIC	DOSE(S) mg/Kg	ROUTE	PERSON		RING						
S												
	13.6 If surgery will be performed on no	on-rodent animals, it must be done in an	approved surgical facility.									
	Name of Facility:											
	Location:											

	NOTE: Postoperative care must be in accordance with current established veterinary, medical, and nursing procedures (Animal Welfare Act). Postoperative care includes: Observing the animal to ensure uneventful recovery, providing adequate care for surgical incisions, and maintaining appropriate medical records. Administration of analgesics, or other medications and supporting fluids, is done as required.										
	(Public Health Service Policy).										
	14.1 Name the person(s) providing postoperative care:										
	NAME			NAME							
CARE	14.2 Provide a postoperative	e care plan by including detailed	answers to the fol	llowing:							
IVE (	FREQUENCY OF POSTOPE	ERATIVE EXAMS		ESTIMATED	DURATION OF MONITORING	3					
ERAT											
LOPE											
POS											
14:											
TION											
SEC	14.3 What complications could reasonably be expected?										
	·										
	If there are any complic	ations indicate that you will noti	fy Veterinary Staff	and follow the	eir recommendations.						
	14.4 List post-procedure ana	algesics / tranquilizers									
	MEDICATION	DOSE(S) mg/Kg	ROUTE		FREQUENCY	DURATION					
	15.0 A volatile anesthetic will	l be used (e.g. isofluorane)				Yes	No				
S	If <b>yes</b> , now will this be vented	d?									
ETIC											
STH											
ANE											
N 15:	What is the location of the ve	enting equipment?									
CTIO											
SEC											
SEC											

	16.0	Euthanasia, check or describe method(s) selected (Reference Polic	<b>y #</b> 4)	):		
		CO2 Inhalation (using compressed gas ONLY) followed by:		Pneumothorax		Cervical Dislocation Exsanguination
		Exsanguination under anesthesia (specify anesthesia and the physic	cal n	nethod:		
		Pentobarbital sodium 100 mg/Kg (for mice: 150 mg/Kg) specify:		I.V.		I.P. ( <i>Rodents ONLY</i> )
₹		Neonates (E16 to P9):		Decapitation		Injectible Anesthetic O.D. Cervical Dislocation
NAS		Other (specify and justify):				
SECTION 16: EUTH						
LIST OF IACUC POLICIES	This http: 1). T 2). C 3). N 4). E 5). E 6). C 7). E 8). N 9). N 10). 11).	section lists all of the applicable policies that pertain to IACUC applie //rwjms.umdnj.edu/research/orsp/ra/iacuc_policies.html Tumor Endpoints Deverorowding Aoribund Animals Euthanasia Body Weight Cell Line Usage EAE Mouse Tail Biopsy for Genetic Analysis Mouse Toeclip Guidelines for Rodent Survival Surgery Expired Drugs Policy	catio	ns. They may be	found	d at the UMDNJ-RWJMS web-site below:
		This section in	tentio	onally left blank.		

Please use this section for any additional information that could not be accommodated for in the application. (Text entry ONLY)

Please use this section for any additional information that could not be accommodated for in the application. (Text entry ONLY)

SECTION 17: RESEARCH PERSONNEL	17.0 Designate the Research personnel to be contacted in case of emergency. (Prior to starting this project off campus contact information <b>must</b> be provided to the Vivarium office.): Please contact the Vivarium to provide confidential emergency contact information.					
	NAME:				EMAIL:	OFFICE PHONE:
	PI:					
	SECONDARY:					
	ALT (1):					
	ALT (2):					
	ALT (3):					
	ALT (4):					
ION 18: TRAINING AND EXPERIENCE	18.1 Describe your training and experience with the procedures and techniques to be used on the animals you will be using in this protocol. If you are inexperienced in these procedures or with this species, describe how you will obtain the appropriate training.					
	18.2 List all individuals (including institution and title) who will be involved in the use of animals and indicate their experience with the specific experimental procedures and species employed in this application. If those individuals are inexperienced, indicate your plans for directly supervising them during training.					
	NAME E	EMAIL	TITLE/INSTITUTION	TRAIN	IING AND EXPERIENCE IN PROPOSED PROCED	URES
ECTI						
S						
SECTION 19: SIGNATURES	<ul> <li>Please indicate by check marks that you agree to all of the following statements:</li> <li>I declare that the information provided in the application is accurate to the best of my knowledge.</li> <li>I will notify the IACUC in writing of any changes to the animal care and use protocol. The request for these changes must be forwarded promptly in writing to the IACUC for approval. Changes include: species; drugs administered; method of anesthesia, analgesia, or euthanasia; surgical procedures; procedures that cause pain or distress in conscious animals; use of prolonged restraint; use of hazardous substances that involves possible exposure of personnel or animals in the animal facility; or change in number of animals used or change in personnel (deletions/additions). If adding personnel you, as the PI, must attest that new personnel either have experience in the procedures to be performed in this study or if not that you will personally train the new personnel</li> <li>I agree to abide by the provisions of the PHS/NIH Guide for the Care and Use of Laboratory Animals. As principal investigator, I assume responsibility for my co-investigators and other personnel involved in this project, with regard to their compliance with the above stated Policies.</li> <li>All animal studies proposed in the grant application cited are described and covered by this animal care and use protocol.</li> <li>All individuals listed in Section 18.2 have read and are familiar with the contents of this protocol.</li> </ul>					
					DATE.	