

OFFICE OF THE VICE PRESIDENT FOR RESEARCH AND INNOVATION

INSTITUTIONAL ANIMAL CARE AND USE

Review Form

Date of application: Proposal number: Approved date: Expiration date:			
Initial submission	Renewa	1	Modification
	I. ADMINIS	TRATIVE DATA	
College / Department:			
Principal Investigator:			
E-mail address:			
Contact number:			
	Projec	t Title	
Funding source:			
Co-investigator(s)			
Name	College / Department	Contact number	E-mail address



Z/4 W W	II. ANIMAL REQUIR	REMENTS
Genus:		Species:
Strain/Subspecies/Breed:		Common name:
Approximate age: Approximate weight: Approximate size:		- - -
Sex:	Male(s)	Female(s) Not specific
Microbiological status:		
Source(s):		
Primary housing location(s):		
Location(s) where manipulation v	vill be conducted:	
Number of animals to be used:	Year 1: Y	ear 2: Year 3:
Total number of animals to be use	ed:	
III. TRA	ANSPORTATION	REMARKS / COMMENTS
Describe how animals will be transfacility, container(s) to be used, re	nsported from source(s) to housi	
	DY OBJECTIVES	REMARKS / COMMENTS
Discuss the main objective of the	study up to its specific objectives	S.



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V. RATIONALE FOR ANIMAL USE	REMARKS / COMMENTS
Explain the reasons why it is necessary to use animal models, justify the appropriateness of the selected species to the study and justify the number of animal(s) to be used in the study.	
VI. DESCRIPTION OF EXPERIMENTAL DESIGN AND ANIMAL PROCEDURES	REMARKS / COMMENTS
Briefly explain the experimental design and specifically describe all	
animal procedures to be employed in the study from the beginning up to	
the end of the experimentation. A flow chart with proper description or labels may be used to present the planned procedure.	
Include the following specific information if applicable:	
✓ Animal identification method(s)	
✓ Method(s) of restraint during sample collection	
✓ Experimental injections or inoculations (substance(s) to be injected or inoculated, dose, site, volume, route and schedule)	
✓ Sample collection (frequency, volume, site, technique)	
✓ Radiation (dosage, schedule)	
✓ Food or fluid restriction	
✓ SURGERY – discuss the whole process including the course of	
action to be performed on non-survival surgery, substance(s)	
to be used on animals such as anesthesia, etc. ✓ Other procedures that will be performed on animals	
oner procedures that will be performed on animals	
VII. PAIN OR DISTRESS CLASSIFICATION AND CONSIDERATION OF ALTERNATIVES	REMARKS / COMMENTS
Identify the pain or distress classification(s) for USDA covered species	
(Appendix I). Explain the possible effect(s) of the inflicted pain or distress to animal model.	
For procedures falling under USDA's classifications D or E, explain	
the possible alternatives that can be utilized or the non-availability of alternatives.	
For methods that can be produced in-vitro, explain the advantages and	
disadvantages of animal model utilization instead of in-vitro cell models or the likes.	



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VIII. ANESTHESIA, ANALGESIA, TRANQUILIZATION, REMARKS / COMMENTS **OTHER AGENTS** *Specify the anesthetic(s), analgesia(s), tranquilizer(s) and other agents* that will be used in the specific part of the experimentation and the necessity of applying such substances, possible side effect(s) if there would be any. Explain also the preparation, dose, volume, manner or route of administration (e.g. injections, topical application) and frequency of application. IX. METHOD OF EUTHANASIA OR DISPOSAL OF REMARKS / COMMENTS ANIMALS AT THE END OF THE STUDY Indicate the proposed method of euthanasia. If chemical agent is to be used, specify the dosage and route of administration. If the method of euthanasia is NOT consistent with the guidelines provided by BAI (Bureau of Animal Industry), provide specific justification as to why such method must be used. Also, indicate the method of carcass disposal. REMARKS / COMMENTS **HAZARDOUS AGENTS** Identify the hazardous agent(s) that will be used on animals (radionucleotides, biological agents, drugs, recombinant DNA, etc.). Explain the possible effects of these agents to the animal models, and the level of animal biosafety as the experimentation is performed. Also, describe the safety protocols in handling and disposal of hazardous agent identified. XI. BIOLOGICAL MATERIAL / ANIMAL PRODUCTS FOR REMARKS / COMMENTS

USE IN ANIMAL

Fill out the follo	wing fields if biological materials (e.g. cancer cell etc.) will be induced to model animal.	
Specify material: Source:		
	Sterile Attenuated	
Has the materia	l been tested for pathogen?	
	Yes (Attach copy of results) No	
passed through and/or the mate	fy that the tested materials to be used have not been a rodent species outside of animal facility in question erial is derived from the original tested sample. To the wledge the material remains uncontaminated with rodent	
	Signature of Principal Investigator	
XII.	GENETICALLY ENGINERRED ANIMALS	REMARKS / COMMENTS
manipulations t that the animal:		
XIII. ENI	EXCEPTIONS FROM ENVIRONMENTAL RICHMENT FOR NON-HUMAN PRIMATES OR	REMARKS / COMMENTS
	EXERCISE	
	g an exception for scientific reasons from the n for environment enrichment for non-human primates?	
	EXERCISE g an exception for scientific reasons from the	

institution s plan to provide dogs with opportunity for exercise? Yes (explain basis of request) No	
XIV. FIELD STUDY	REMARKS / COMMENTS
If animals in the wild will be used, describe how they will be observed, any interactions with the animals, whether the animals will be disturbed or affected, and any special procedures anticipated. Indicate if national and/or local permits are required and whether they have been obtained.	
XV. SPECIAL CONCERNS OR REQUIREMENTS OF THE STUDY	REMARKS / COMMENTS
List any special housing, equipment, animal care or departures from the Guide.	
XVI. PRINCIPAL INVESTIGATOR CERTIFICATIONS	REMARKS / COMMENTS
I certify that: • I have attended the institutionally required investigator training course.	
Year of attendance: Location:	
 I have determined that the research proposed herein is not unnecessarily duplicative of previously reported research. That the individuals listed in Section I are authorized to conduct procedures involving animals under this proposal, have attended institutionally required investigator training course, received training. I have reviewed the pertinent scientific literature and the sources 	
and/or databases as noted in Section VII and have found no valid alternative to any procedures described herein which may cause more than momentary pain or distress, whether it is relieved or not.	

significant stranges in this study. I will notify the IACUC regarding any unexpected study results that impact the animals. Any anticipated pain or distress, morbidity or
mortality will be reported to the attending veterinarian and the IACUC.
I am familiar with and will comply with all pertinent institutional and national rules and policies.
Signature over printed name of Principal Investigator / Date

XVII. CONSENT	REMARKS / COMMENTS
Adviser consent:	
Signature over printed name / Date	
Veterinarian certification:	
Signature over printed name / Date	
XVIII. FINAL APPROVAL	REMARKS / COMMENTS
Certification of review and approval by the Institutional Animal Care and Use Committee:	
Signature over printed name / Date	