

Lewis & Clark College IACUC

LABORATORY ANIMAL USE PROTOCOL APPLICATION

Purpose: Pls should complete this form prior to using live vertebrate animals in a research laboratory, for renewal (every three years at a minimum), and for modification. The completed form should be submitted to the Institutional Animal Care and Use Committee (IACUC).

Researcher and Laboratory	y Information							
Researcher First Last Name:	Title:		De	partment:		Email:		
Cell Phone:	Lab phone:		Lab	Building:		Lab room no:		
Does your lab have a current eme			t info	rmation? Yes N	10			
Does your lab keep training recor	ds for all personnel as requ	ired?		Yes N	10			
Project Information								
Protocol Application Date	Protocol Desired Star	t Date	Prop	oosal type:	If Re	newal or Mod, pro	vide curren	t/prior
Other Information			_	Initial submission	Proto	col no:		
Julei information				Renewal	Start I	Date:		
				Modification	Expira	ation Date:		
Funding: Please list active a	nd pendina fundina							
Sponsor 1:	Project Start Date:	Project End Da	ate:	Project Title 1:				
Current Pending	Acct no., if available:			I Drainat Title 2:				
Sponsor 2:	Project Start Date:	Project End Da	ate:	Project Title 2:				
Current Pending	Acct no., if available:							
Sponsor 3:	Project Start Date:	Project End Da	nto.	Project Title 3:				
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Current Pending	Acct no., if available:							
Senior and Key Personnel: additional pages as necessary).	List all individuals auth	orized to condu	ıct pı	rocedures involving anim	nals un	der this proposa	ıl (attach	
First Last Name:	Email:	Role (edit if other):		Level of Contact*	Ma eutha			
	Linaii.	(edit ii otilei).		2010. 0. 00	euma	IIIZE ! assessmen	t & docume	ntation:
Level of Contact dictates training minimal handling); Tier 3: Studen								
Collaborative Research							YES	NO
 Is any part of this research b If yes, please provide a desc 								
 Has collaborating institution's If yes, please provide outcon 								
Will an intra-institutional agree	eement or MOU be required	I for this work? If	yes, p	olease explain:				

Study Objectives Briefly explain the goals of the research; and the project's relevance to human or animal health, the advancement of knowledge, and/or the good of						
society. Us	e non-technical Ian	guage that IACUC me	mbers with var	ried backgrounds will understa	nd.	
mimal D	lucus onto					
	etotal number of an	nimals used for this pro	piect, including	breeders, experimental animal	ls. and animals not used exper	rimentally.
Senus		Species	<u>,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,</u>	Strain/Breed	Common Name	Total Number
'ear	Number	Age/Weight	Sex	Bacteriological status	Viral status	Source(s)
'ear	Number	Age/Weight	Sex	Bacteriological status	Viral status	Source(s)
'ear	Number	Age/Weight	Sex	Bacteriological status	Viral status	Source(s)
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ou.	Train 501	/ tgc/vvcigitt	OCA	Buoteriological status	Thu Status	334,33(3)
'ear	Number	Age/Weight	Sex	Bacteriological status	Viral status	Source(s)
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ear	Number	Age/Weight	Sex	Bacteriological status	VII al Status	Source(s)
ear	Number	Age/Weight	Sex	Bacteriological status	Viral status	Source(s)
4 ! 6 ! 4	tion for the Use	of Animala				
	tion for the Use		oggang it is not	populary to upo animala. Justify t	the appropriateness of the app	sice colocted Justify th
umber of a	animals to be used	I, including the method	s and sources	cessary to use animals. Justify tused to determine that alternati	ives (e.g. replacement, reduction	on, refinement) to the us
animals	and to the procedu	res have been conside	ered. Number n	nust be minimum required to o	btain scientifically valid results	

9 Attach Description of Experimental Design and Animal Procedures

7

Please attach a document clearly explaining the experimental design and specify all animal procedures. All procedures to be employed in the study must be described; the description should include all procedures performed on live animals, the number of procedures, the number of animals involved in each procedure, the number of procedures an individual animal will experience. This description should allow the IACUC to understand the experimental course of an animal from its entry into the experiment to the endpoint of the study. A flowchart may be an effective presentation of the planned procedure. Consider including relevant animal identification methods, methods of restraint, experimental injections or inoculations, blood withdrawals, radiation, food or fluid restriction, pharmaceutical-grade and non-pharmaceutical grade compounds, other procedures, resultant effects expected, other potential stressors, procedures to monitor and minimize distress, experimental endpoint criteria, veterinary care, and surgical procedures.

10	Housing, Environment, Transportation (if question is not applicable, please indicate N/A in appropriate space below)	
	1. How and where will the animals be housed?	
	2. Where will the manipulations be conducted?	
	Will the animals be exposed to non-standard housing? If yes, please explain.	
	4. Will any of the animals be singly housed? If yes, please provide scientific justification.	
	5. Will the animals be exposed to any non-standard environmental conditions? If yes, please explain.	
	6. Will the animals be exposed to any non-standard food or water, including withholding food? If yes, please explain.	
	7. Please describe any enrichment that will be provided to the animals.	
	8. Please describe how the health of the animals will be monitored.	
	9. Will animals be transported? If yes, please describe.	
	10. Will animals be physically restrained? If yes, please describe and provide scientific justification.	
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11	Method of Euthanasia or Disposition of Animals Please describe the proposed method of euthanasia and/or disposition. If a chemical agent is used, specify the dosage range and route of adrithe method of euthanasia is not consistent with the current (2020) AVMA Guidelines for the Euthanasia of Animals, provide scientific justification is used. Describe how you will determine that the euthanized animals are actually dead and what will happen to any animal euthanized. Indicate who will euthanize animals and indicate their training and experience with the method of euthanasia and the species involved.	on as to why als not

2	Will you use Anesthesia, Analgesia, Tranquilization, or Other Agents? Yes No or N/A
	If Yes, specify the anesthetics, analgesics, sedatives or tranquilizers that will be used. Include the name of the agent(s), the dosage range, route(s), possible side effects, and schedule of administration.
42	Described and the production of the production o
13	Does the Project Involve Surgery or Other Procedures? Yes No or N/A If yes, please respond to the questions below. 1. Identify and describe the surgical procedure(s) to be performed. Include preoperative procedures [e.g., fasting, analgesic loading], and monitoring and supportive care during surgery. Include the aseptic methods to be used.
	2. Identify the individual(s) that will perform surgery and their qualifications, training, and/or experience.
	Identify the location where surgery will be performed. [building(s) and room(s)]
	4. If survival surgery, describe postoperative care that will be provided and frequency of observation. Identify the responsible individual(s) and location(s) where care will be provided. [building(s) and room(s)] Include detection and management of postoperative complications during work hours, after hours, weekends and holidays.
	5. If non-survival surgery, describe how euthanasia will be provided and how death will be determined.
	6. Are paralytic agents used during surgery? If yes, please describe how ventilation will be maintained and how pain will be assessed.
	7. Has major or minor survival surgery been performed on any animal prior to being placed on this study? [Major survival surgery penetrates and exposes a body cavity or produces substantial impairment of physical or physiologic functions or involves extensive tissue dissection or transection (such as laparotomy, thoracotomy, craniotomy, joint replacement, or limb amputation)]. If yes, please explain.
	8. Will more than one survival surgery be performed on an animal while on this study? If yes, please justify.

1. What is the status of Institutional Biosafety Committee (IBC) review of this project? 2. If the IBC has approved this project, provide approval number and approval e 3. What is the Biosafety Level (BSL)? 4. Describe the practices and procedures required for the safe handling and disposal of contaminated a PPE, protective equipment such as fume hoods). 5. Describe methods for removal of radioactive waste and/or the monitoring of radioactivity, or indicate 6. Discuss any additional biosafety concerns:	
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Does the Project Involve Animal Breeding Colonies? Yes No or N/A If yes, provided the Project Involve Animal Breeding Colonies? Yes No or N/A If yes, provided the Project Involve Animal Breeding Colonies? Yes No or N/A If yes, provided the Project Involve Animal Breeding Colonies? Yes No or N/A If yes, provided the Project Involve Animal Breeding Colonies? Yes No or N/A If yes, provided the Project Involve Animals For use in approved experimental profequing that all animals or teaching must be counted again Number of breeders and number of young that cannot be used in experiments; and number manipulations 3. Describe breeding methods, including number of females per male or other 4. Indicate weaning age and describe how animals will be separated at weaning 5. Describe methods of identification of individual animals 6. Describe how you will dispose of founders, breeding pairs, or any non-transgenic animals	l/A.
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5. Describe methods of identification of individual animals 6. Describe how you will dispose of founders, breeding pairs, or any non-transgenic animals	
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Does the Project Involve Field Studies? Yes No or N/A If yes, please respond	
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Dood this i toject introller i leid otdales. I les introller il ves, please respond	
If animals in the wild will be used, describe how they will be observed, any interactions with the animals and any special procedures anticipated. Indicate if federal, state, and/or local permits are required and with the animals and any special procedures anticipated. Indicate if federal, state, and/or local permits are required and with the animals and any special procedures anticipated.	the question helow

17	Is this Application a Renewal or	Modification of a Prote	ocol Previously Appr	oved by the IACUC?	Yes No or N/A		
.,	If yes, provide a brief progress report, in the protocol.						
40	Special Concerns or Requireme	ents of the Study?					
18	List any special housing, equipment, an		e from the Guide				
19	Principal Investigator Certificati	ons					
	By signing below, I confirm that I will comply with all applicable regulations and guidelines for animal care and use, including but not limited to the U.S. Government Principles and relevant portions of the PHS Guide, PHS Policy, the Animal Welfare Act, and Lewis & Clark College's Animal Care and Use in Research and Teaching Policy. I will follow the guidelines and instructions of the Institutional Animal Care and Use Committee (IACUC), and notify the IACUC of any proposed changes prior to implementation. I am aware that deviations from an approved protocol, or violations of pertinent policies, guidelines, or laws could result in immediate suspension of the project. I certify that the number of animals proposed for use is the minimum number required to obtain scientifically valid results. I assume responsibility for the ethical conduct of this project and for protecting the welfare of the animal subjects and human participants. The research proposed herein is not unnecessarily duplicative of previously reported research. I will notify the IACUC regarding any unexpected study results that affect the animals. Any unanticipated pain or distress, morbidity or mortality will be reported to the attending veterinarian and the IACUC. I will take full responsibility for this protocol, make sure that all individuals involved have read and understand the procedures in the protocol, and confirm that all participants will be qualified and appropriately trained for the species used. A list of participants will be provided prior to their involvement.						
	PI Signature:	Date:	Co-PI Sig	nature:	Date:		
		1 / 11 14 011					
20	IACUC Review (this section to b			-1	New Doctor of New York (for the late)		
	Type of Review. If other explain below	Date Received by IACUC	Date Reviewed by IACU	C IACUC Decision	New Protocol Number (if applicable)		
					New Protocol Start Date:		
					New Protocol Expiration Date:		
	Notes for IACUC use only:						
	Chair First Last Name:	Chair	Signature:		Date:		

Researcher First Last Name	Title	Department	Protocol Number	Protocol Expiration Date

21 Laboratory Personnel Certifications

- A. I have successfully completed Lewis & Clark's Responsible Conduct of Research (RCR) online training modules.
- B. I have participated in Lewis & Clark's face-to-face RCR training.
- C. I have participated in Lewis & Clark's Biosafety Training.
- D. I have successfully completed the required online animal training modules, including both laboratory research and species-specific courses.
- E. I have completed the occupational safety and health training.
- F. I have read the procedures in this protocol and the approved endpoints described.
- G. I have successfully completed training on performing the procedures in which I will be participating.
- H. I agree to follow the procedures and approved endpoints.

First Last Name:	Signature:	Date:
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