

INSTITUTIONAL BIOSAFETY COMMITTEE SAN JOSÉ STATE UNIVERSITY BIOLOGICAL USE AUTHORIZATION APPLICATION

Attachment G. Research Organisms – Vertebrates, Invertebrates, or Plants

Chec	ck all that apply. Make sure to include details in the attached SOP.
\boxtimes	We will be introducing materials into research organisms (Complete Table I)
	☑ We will be introducing recombinant or synthetic nucleic acid molecules into research organisms. (Complete
	Tables II-III (animals – vertebrates or invertebrates) or Tables IV-V (plants) and Attachment A)
	\square We will be introducing infectious materials into research organisms. (Complete Tables II-III (animals –
	vertebrates or invertebrates) or Tables IV-V, and Attachment B)
	\square We will be introducing unfixed human or non-human primate organs, tissues, or cell cultures (OTCC) into
	research organisms (Complete Tables II-III (animals – vertebrates or invertebrates) or Tables IV-V (plants), and
	Attachment C)
\boxtimes	We will be using transgenic animals (vertebrates or invertebrates)
	☑ We will be creating transgenic animals (vertebrates or invertebrates)
	☑ We will be purchasing or obtaining transgenic animals (vertebrates or invertebrates) from another group
	☑ We will be breeding transgenic animals (vertebrates or invertebrates)
	☑ We will be transferring transgenic animals (vertebrates or invertebrates). Describe below and list prospective
_	recipients:
	Transgenic drosophila are shared upon request with other academic researchers
	We will be using transgenic plants
	☐ We will be creating transgenic plants
	\square We will be purchasing or obtaining transgenic plants from another group
	☐ We will be transferring transgenic plants. Describe below and list prospective recipients:
	Click or tap here to enter text.
	Work requires a USDA-APHIS Permit.
	We will be performing safety tests or screening for pathogenic or infectious agents before or after inoculation
	into animals (vertebrates or invertebrates).

A Standard Operating Procedure (SOP) must be attached that describes the work with research organisms. Describe the species, transporting, and decontamination methods, if applicable. For transgenic/genetically modified plants, describe any special growth requirements and whether recombinants are expected to be more pathogenic, as well as the transformation method used, the gene information, and containment requirements. A detailed step-by-step protocol is not necessary, but provide sufficient information on your procedures so that the committee can identify the steps that involve the greatest likelihood of worker or environmental exposure. Include the steps that will be conducted in a biological safety cabinet. Consult the SOP template for other required components.

Research Organism	(inclu	Recipie ude source na	Administration Route	Max. Conc. Administered	Max. Vol.		
(Genus, species, strain)	Recombinant DNA Construct (germ line /somatic?)	Other nucleic acid molecule (describe)	Microbe (only if microbe will be injected into research organism)	OTCC (unfixed human or non- human primate organs, tissues, or cell cultures)			
Aedes aegypti	NA	double stranded RNA	NS	NA	injected into mid thorax	1.2 ug	0.2 uL
Drosophila melanogaste r	Various insertions to alter signaling pathways using pUAST plasmid (germline)	NA	NA	NA	Outsourced		

⊠ N/A	I.B. Research Organism - Infectious Agent Details						
Species of Animal (vertebrates and invertebrates)/ Plant	# of Animals/ Plants	Infectious Agent	Duration of Infectivity	Route of Shedding/ Excretion & Interval			

⊠ N/A	II.A. Animal (Vertebrate and Invertebrate) Containment During Agent Introduction							
Species of Animal		Biological Agent or	Containment/Locations					
(vertebrate a	nd	recombinant DNA	Prior to procedures		During Procedures		Post Procedures	
invertebrate)/Strain		(include source details)	BSL/ABSL	Bldg/Room	BSL/ABSL	Bldg/Room	BSL/ABSL	Bldg/Room

□ N/A	II.B. Animal (Vertebrate and Invertebrate) Housing Post Administration						
After delivery o	After delivery of agents, animals (vertebrate and invertebrate) will be housed in (check all that apply):						
⊠ Separate cag	es from other animals						
☐ Well-labeled	cages indicating possible hazards to animal staff						
☐ Micro-isolate	☐ Micro-isolator cages						
☐ Conventional/ABSL-1 facility							
☐ ABSL-2 facility							
☐ ABSL-2 facility for 72 hours, then transferred to ABSL-1 facility							
☐ Live animals	☐ Live animals will not be returned to animal facilities after delivery						
☐ Other (specif	y): Click or tap here to enter text.						

□ N/A	III. General Animal (Vertebrate and Invertebrate) Housing or Handling							
Bldg/Room		Species	Procedures (i.e., breeding, handling, injections)	Shared room? (Y/N)	Proposed BSL			
DH 608A		Aedes aegypti, Drosophila melanogaster	breeding, handling, injections, dissection	N	1			

⊠ N/A	IV. Plant Research Organism					
Species of Plant	Transgenic (Y/N)		/Notification (Type, aber)	Noxious Weed (US or CA) (Y/N)	Method of Reproduction (self, wind pollinator, insect pollinator, human intervention required)	
		(Y/N)	(Type, Number)			

⊠ N/A	V. Plant Growth and Housing						
Bldg/Room	Bldg/Room Plant Stage of growth (i.e., seedling)		Shared room? (Y/N)	Proposed BSL			