# **IBC MAIN APPLICATION FORM**

IRBNet Number: Previous IRBNet Number (If applicable): Click here to enter text. Click here to enter text.

### **1: PRINCIPAL INVESTIGATOR INFORMATION**

#### **1.1. PRINCIPAL INVESTIGATOR INFORMATION**

Principal Investigator: Click here to enter text. Position/Title: Click here to enter text. Department/College: Click here to enter text. Office/Cell Phone #: CdTv&r2 (Iled[Pd[P)1 ()2 (e)1 (:)?)]J@J25 @376 rgT2 1 T@l[C2aCdil ad (Iled1 (6 (k h

Homan bood, tisse, or bdilyfid.

Transgnic and/or pthognic pants

Recombiant or systhetic nuleic acid molecles

Infected or ptentiallyinfected cell lines.

Animal bood, tisse, or bdilyfid

Radioactie materials.

Shipnopfoloigal materials

#### **1** | P a g e Institutional Biosafety Committee (IBC): Application Form Version 7.0, 2018-11-18

RESEARCH ACTIVITIES	BUILDING	ROOM	BIOSAFETY LEVEL	SHARED ROOM? (YES/NO)
			BSL 3 BSL 1 & BSL 2 BSL1, BSL2, BSL3	
Click here to enter text.	Click here to enter text.	Click here to enter text.	BSL 1 BSL 2 BSL 3 BSL 1 & BSL 2 BSL 1 & BSL 2 BSL1, BSL2, BSL3	Yes No
Click here to enter text.	Click here to enter text.	Click here to enter text.	BSL 1 BSL 2 BSL 3 BSL 1 & BSL 2 BSL 1 & BSL 2 BSL1, BSL2, BSL3	Yes No

## 2: NIH REVIEW CATEGORY & SUBCATEGORY

Check all the categories, subcategories and information that apply. NIH Office of Science Policy Website: <u>https://osp.od.nih.gov/</u>

Does your project include the use of recombinant or synthetic nucleic acid molecules?

Yes, complete the table below No, skip to section 3

CATEGORY	OVERSIGHT BY	INCLUDES/SUBCATEGORIES
	NIH Director, RAC & IBC	Studies that involve the deliberate transfer of drug resistance to microorgo()-1Cirtioct99 (t).9 h1 (c)-2 T@coorooii

CATEGORY OVERSIGHT BY

INCLUDES/SUBCATEGORIES

CATEGORY	OVERSIGHT BY	INCLUDES/SUBCATEGORIES		
	Form Submittal	more human research participants and meets the criteria of Section III-C, it is not exempt under this Section.		
		F-2: Those that are not in organisms, cells, or viruses and that have not been modified or manipulated (e.g., encapsulated into synthetic or natural vehicles) to render them capable of penetrating cellular membranes.		
		F-3: Those that consist solely of the exact recombinant or synthetic nucleic acid sequence from a single source that exists contemporaneously in nature.		
		F-4: Those that consist entirely of nucleic acids from a prokaryotic host, including its indigenous plasmids or viruses when propagated only in that host (or a closely related strain of the same species), or when transferred to another host by well-established physiological means.		
		F-5: Those that consist entirely of nucleic acids from a eukaryotic host including its chloroplasts, mitochondria, or plasmids (but excluding viruses) when propagated only in that host (or a closely related strain of the same species).		
		F-6: Those that consist entirely of DNA segments from different species that exchange DNA by known physiological processes, though one or more of the segments may be a synthetic equivalent.		
		F-7: Those genomic DNA molecules that have acquired a transposable element, provided the transposable element does not contain any recombinant and/or synthetic DNA.		
		F-8: Those that do not present a significant risk to health or the environment (see Section IV-C-1-b-(1)-(c))		

## 3: SPECIFIC AIMS OF PROJECT AND PROTOCOLS USED

#### **3.1. SPECIFIC AIMS**

3.1.1. Provide an overall summary of the project and briefly explain in **language understandable to the general public** the specific aim(s) of the study. Click here to enter text.

### **3.2.** BENEFITS

3.2.1. Explain in **language understandable to the general public** how the information gained in this study will benefit human or animal health, the advancement of knowledge, and/or server the good of society. Click here to enter text.

### **3.3. OUTLINE OF PROTOCOLS**

3.3.1. Outline the biohazard control plan for recombinant DNA work and other biohazardous work.

Briefly describe the general types of experimental procedures that will be performed. Address the potential sources of risk to personnel (aerosol generation, needle sticks, etc.) and/or the

environment, and how these risks will be managed.

Describe safety devices that will be used (e.g. biosafety cabinets, hand washing facilities, puncture resistant sharps containers, etc.)

Include decontamination/disinfection processes.

Include plans for disposing of materials.

Click here to enter text.

### 4: BIOLOGICAL MATERIALS IN PROJECT

4.1. List the recombinant DNA used in the proposed work.

Include cloned gene(s), vectors used; give both name and type of each vector.

Click here to enter text.

4.2. List the genes described above that will be expressed: Click here to enter text.

4.3. List organism(s) or cell lines are used in the proposed work: Click here to enter text.

4.4. Include information if any recombinant or synthetic DNA materials will be used in any vertebrate animal model (covers drosophila research, along with other animals): Click here to enter text.

4.5. List the infectious or pathogenic agents used in the proposed work.

### **5: PERSONNEL AND TRAINING**

5.1. Please list the PI and other personnel who will be handling biological agents. Include personnel who are graduate level or higher or who will have a role in training other lab members.

#### TABLE 5.1.A. PERSONNEL AND TRAINING

To add additional people, click on the + at the end of each box.

	PHONE #	EMAIL	CREDENTIALS	COMPLETED	ROLE IN			
		ADDRESS		TRAINING	PROJECT			

Sandals must not be worn with working in the laboratory. Other protective equipment, such as splash goggles, face shields, aprons, thermal or cut-resistant gloves, hearing protection or respirators, must be worn when conditions dictate.

In a class situation, student shall purchase or obtain the necessary and approved PPE designated by the

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