## **Checklist for Adult Sponsor (1)**

This completed form is required for ALL projects.

ISEF forms are fillable, so you can type in responses within the PDF file and then print.

#### To be completed by the Adult Sponsor in collaboration with the student researcher(s):

Stude	ent	nt's Name(s):				
Proje	ct -	t Title:				
1. L		I have reviewed the ISEF Rules and Guidelines,	including the s	cience fair ethics statem	ent.	Fillable forms
2. <b>C</b>		I have reviewed the student's completed Stude	ent Checklist (1/	A) and Research Plan/Proj	ject Summary.	easier to fit student
3. <b>C</b>		I have worked with the student and we have di	scussed the po	ssible risks involved in th	e project.	names, long titles
4. E		<ul> <li>The project involves one or more of the following</li> <li>Humans</li> <li>Vertebrate Animals</li> </ul>	ing and require Pote □	s prior approval by an SR ntially Hazardous Biologi Microorganisms D r	C, IRB, IACUC or cal Agents DNA 🛛 Tis	ssues
5. <b>C</b>		Items to be completed for <b>ALL PROJECTS</b> <ul> <li>Adult Sponsor Checklist (1)</li> <li>Student Checklist (1A)</li> <li>Regulated Research Institutional/Indu</li> <li>Continuation/Research Progression Federation</li> </ul>	strial Setting Fo orm (7) (when a	Research Plan/Project Su Approval Form (1B) orm (1C) (when applicable pplicable)	mmary <	Only check the boxes that are appropriate to your research project.
Addit [	tio ]	<ul> <li>bonal forms required if the project includes the understanding student designed invention see full text of the rules.)</li> <li>Human Participants Form (4) or appropria</li> <li>Sample of Informed Consent Form (when Qualified Scientist Form (2) (when application)</li> </ul>	use of one or m ns/prototypes. te Institutional applicable and uble and/or requ	ore of the following (che (Requires prior approval b IRB documentation /or required by the IRB) ired by the IRB)	ck all that apply) by an Institution	): al Review Board (IRB);
[		<ul> <li>Vertebrate Animals (Requires prior approval, s</li> <li>Vertebrate Animal Form (5A) - for projects</li> <li>Vertebrate Animal Form (5B) - for projects</li> <li>Use Committee (IACUC) approval required</li> <li>Qualified Scientist Form (2) (Required for a</li> </ul>	see full text of t conducted in a conducted at a d prior experime all vertebrate a	he rules.) school/home/field resea Regulated Research Insti entation.) himal projects at a regulat	rch site (SRC pri tution. (Institutio ted research site	or approval required onal Animal Care and e or when applicable)
		<ul> <li>Potentially Hazardous Biological Agents (Req</li> <li>Potentially Hazardous Biological Agents Ri</li> <li>Human and Vertebrate Animal Tissue Form fresh or frozen tissue, primary cell cultures</li> <li>Qualified Scientist Form (2) (when applica</li> <li>The following are exempt from prior review similar microorganisms, for projects using projects using color change coliform water organisms.</li> </ul>	uires prior app isk Assessment n (6B)-to be co s, blood, blood ble) w but require a manure for co er test kits, mici	roval by SRC, IACUC or IB Form (6A) mpleted in addition to Fo products and body fluids Risk Assessment Form 3: mposting, fuel production obial fuel cells, and proje	C, see full text c rm 6A when pro c projects involvin n or other non-c ects involving de	of the rules.) ject involves the use of ng protists, archae and ulturing experiments, composing vertebrate
C	ב	Hazardous Chemicals, Activities and DevicesRisk Assessment Form (3)Qualified Scientist Form (2) (required for	(No SRC prior projects involvi	approval required, see fu ng DEA-controlled substa	ll text of the rule ances or when a	es.) oplicable)
[		Other Risk Assessment Form (3)	usually the ce/research er or parent, n	ot	This befo	must be dated re the "Actual Start " on Form 1A
Ľ		I attest to the information checked ab those a	are different).	nd agree to abide by	the science rair	etnics statement.
		E .				K
Adul	lt S	Sponsor's Printed Name Signature	Э		Date of Reviev	v (mm/dd/yy)
Phor	าค	Fmail				

# Student Checklist (1A)

This form is required for ALL projects.

This form is unique to each **project** not each student.

1.	a. Student/Team Leader:	Grade:		
	Email:	Phone:		
	b. Team Member: o	c. Team Member:		
2.	Title of Project:		This is usually the teacher or parent, not	
			the research mentor (if	
3.	School:	School Phone:		
Sc	hool Address:			
4.	Adult Sponsor: This is usually the teacher (or parent), not the reserved SPC/UPP (ACUC or other processor	Phone/Email:		
э. о	Does this project need SkC/IRB/IACUC of other pre-appr		This defaults to "No"	
6.	Is this a continuation/progression from a previous year? If Yes:		but change it for your	
	<ul> <li>a. Attach the previous year's □ Abstract and □ F</li> <li>b. Explain how this project is new and different from prev</li> <li>□ Continuation/Research Progression Form (7) Tr</li> </ul>	Research Plan/Project Sur vious years on his should be the date that	IF the student has continued their project, this year's poster	
7.	This year's experimentation/data collection:	illecting data.	should focus on the work from the current year.	
	Actual Start Date: (mm/dd/yy) End	Date: (mm/dd/yy)		
8.	Where will you conduct your experimentation? (check all	that apply)		
	□ Research Institution □ School □ Field □ F	Home 🛛 Other:		
9.	Source of Data:  Collected self/mentor  Other Describe/url:		Don't forget to say where you got your data (if other than yourself/mentor)	
10.	List the name and address of all non-home and non-sch virtually or on-site:	ool work site(s), whether y	you worked there	
Na	me			
Ad	dress:			
Pho	one/email			
11.	Complete a Research Plan/Project Summary following and attach to this form.	the Researc <mark>Every project</mark> r detailed Resea	nust have a structions rch Plan that is	
12.	An abstract is required for all projects after experiment	tation. <b>BEFORE</b> the pr	proved oject begins.	

### **Research Plan/Project Summary Instructions**

A complete Research Plan/Project Summary is required for ALL projects and must accompany Student Checklist (1A).

#### All projects must have a Research Plan/Project Summary

- a. The Research Plan is to be written prior to experimentation following the instructions below to detail the rationale, research question(s), methodology, and risk assessment of the proposed The Research Plan is the most important document of
- b. If changes are made during the research, such changes can be that some changes may require returning to the IRB or SRC for required, this addendum serves as a project summary to explain
- c. If no changes are made from the original research plan, no proj • Some studies, such as an engineering design or mathemat change through the course of research. If such changes or ossef.okstate.edu.
  - and can be appended to the original research plan.
  - The Research Plan/Project Summary should include the fol This detailed description of the research will guide the SRC
    - a. **RATIONALE:** Include a brief synopsis of the backgrour to be able to determine if the proper forms were completed research is important and if applicable, explain any sociand if they contain the correct information.
    - b. **RESEARCH QUESTION(S), HYPOTHESIS(ES), ENGINE** the rationale described above?
    - c. Describe the following in detail:
      - List of materials:

Research plan must be **VERY** detailed and clearly delineate the role of the student versus the role of the mentor.

- Procedures: Detail all procedures and experimental tense!!
- applicable, the source of data used. Describe only y
   Risk and Safety: Identify any potential risks and safe It must include proposed AND actual start and end dates.
- Data Analysis: Describe the procedures you will use
- d. **BIBLIOGRAPHY:** List major references (e.g. science jot Must include all details on the work site. If you plan to use vertebrate animals, one of these refe

# Items 1–4 below are subject-specific guidelines for additional items to Must identify the student and mentor role.

#### 1. Human participants research:

- **a. Participants:** Describe age range, gender, racial/ethnic composition of participants. Identify vulnerable populations (minors, pregnant women, prisoners, mentally disabled or economically disadvantaged).
- b. Recruitment: Where will you find your participants? How will they be invited to participate?
- c. Methods: What will participants be asked to do? Will you use any surveys, questionnaires or tests? If yes and not your own, how did you obtain? Did it require permissions? If so, explain. What is the frequency and length of time involved for each subject?
- **d. Risk Assessment:** What are the risks or potential discomforts (physical, psychological, time involved, social, legal, etc.) to participants? How will you minimize risks? List any benefits to society or participants.
- e. Protection of Privacy: Will identifiable information (e.g., names, telephone numbers, birth dates, email addresses) be collected? Will data be confidential/anonymous? If anonymous, describe how the data will be collected. If not anonymous, what procedures are in place for safeguarding confidentiality? Where will data be stored? Who will have access to the data? What will you do with the data after the study?
- f. Informed Consent Process: Describe how you will inform participants about the purpose of the study, what they will be asked to do, that their participation is voluntary and they have the right to stop at any time.

#### 2. Vertebrate animal research:

- a. Discuss potential ALTERNATIVES to vertebrate animal use and present justification for use of vertebrates.
- b. Explain potential impact or contribution of this research.
- c. Detail all procedures to be used, including methods used to minimize potential discomfort, distress, pain and injury to the animals and detailed chemical concentrations and drug dosages.
- d. Detail animal numbers, species, strain, sex, age, source, etc., include justification of the numbers planned.
- e. Describe housing and oversight of daily care.
- f. Discuss disposition of the animals at the end of the study.

#### Potentially hazardous biological agents research:

- a. Give source of the organism and describe BSL assessment process and BSL determination.
- b. Detail safety precautions and discuss methods of disposal.

#### 4. Hazardous chemicals, activities & devices:

- a. Describe Risk Assessment process, supervision, safety precautions and methods of disposal.
- b. Material Safety Data Sheets are not necessary to submit with paperwork.

### **Approval Form (1B)**

A completed form is required for each student, including all team members.



#### 2. To be completed by the local or affiliated Fair SRC (Required for projects requiring prior SRC/IRB APPROVAL. Sign 2a or 2b as appropriate.)

a. Required for projects that need prior SPC/IPB approval			b. Required for research conducted at all Regulated
BEFORE experit Do not write anything in this			Rese Do not write anything in
potentially haza box (2a) unless you are the			This proje this box (2b).
SRC/IRB has cai			(not home or high school, etc.), was reviewed and approved
Project Summary a designee.			by the proper institutional board before experimentation and
signature indicates approval of the Research Plan/Project			complies with the ISEF Rules. Attach (1C) and any required
Summary before the student begins experimentation.			institutional approvals (e.g. IACUC, IRB).
SRC/IRB Chair's Printed Name Signature Date of Approval (mm/dd/yy) (Must be prior to experimentation.)			SRC Chair's Printed Name Signature Date of Signature (mm/dd/yy) (May be after experimentation)

### 3. Final ISEF Affiliated Fair SRC Approval (Required for ALL Projects)

SRC Approval Af Do n	<b>( 3.</b> )			
Regional SRC Chair's Printed Name	Signature		Date of Approval (m	ım/dd/yy)
State/National SRC Chair's Printed Name (where applicable)	Signature		Date of Approval (m	nm/dd/yy)

### **Revised-Regulated Research Institutional/Industrial Setting Form (1C)**

This form must be completed AFTER experimentation by the adult supervising the student research either virtually or on site, conducted in a regulated research institution, industrial setting or any work site other than home, school or field.

Student's Name(s)

01							
Tit	itle of Proiect						
	If any of the research <b>was done</b> at a standard research facility						
To (Re	<ul> <li>(college or university, pharmaceutical company, environmental or State</li> <li>State</li> <li>biomedical testing facility, governmental facility, etc.) or a facility it's where advanced research is allowed (certain high schools or local</li> <li>class), then Form 1C is required.</li> </ul>	t <b>udent(</b> project	( <b>s)) a</b> boot	i <b>fter exp</b> h; please:	erime do not	n <b>tatio</b> print d	<b>n:</b> louble-sided.)
1	alabs), then Form TC is required.	apply).					
	If the science fair student's project will be data analysis only and the data are publicly available, then nothing other than this form is needed.			Yes Yes Yes Yes		No No No No	Some of these are default-clicked "no". Be sure to update these, if appropriate to your project.
2.	patient or student data) or from a private source (such as proprietary data), then the student must show documentation of how the data became available and how/why the student was allowed to view and study it.	e stude the pro	ent i ojec	n any pł t	nase of	the pr	oject, but
3.	The best thing to do for these kinds of private data is have the research mentor from the Regulated Research Institution include a short letter on their letterhead explaining that there was no HIPAA or other violations. This is even if the data are de- identified.	ection, did.	spe	ecific pro	ocedur	es perf	formed).
4.	<ul> <li>Did the student(s) work on the project as part of a group?</li> <li>Were there other high school students present? If yes, please list students names and describe how their work was related or differ</li> </ul>	the rent fro	□ m tł	Yes ne work (	□ of this	No projec	ct.
5.	. If this project is under a grant and needs to be acknolwedged, ple	ease list	t the	e grant s	tateme	ent her	e.
		hatany		ired rouid			

This must be the <b>mentor</b> from the Regulated Research Institution NOT the teacher.	been obtained. Copies are attached if applicated above and that any requested above above and that above abo	dired review This must be dated ble. I further AFTER the "End Date" any on Form 1A.
Direct Supervisor's Printed Name	Signature	Title
Institution		Date Signed (must be after experimenta- tion) (mm/dd/yy)
Address		Email/Phone

### **Qualified Scientist Form (2)**

May be requi biological ager	red for research involving human participants, vertebrate animals, potentially hazardous hts, and hazardous substances and devices. Must be completed and signed before the start of student experimentation.
Student's Name(s) _	
Title of Project	

#### To be completed by the Qualified Scientist:

Scientist Name: \_\_\_\_\_

Educational Background:

Experience/Training as relates to the student's area of research:

Position/Institution:

Email/Phone:

- 1. Have you reviewed the ISEF rules relevant to this project and the science □ Yes 🗆 No fair ethics statement relevant to this project?
- 2. Will any of the following be used?

	a. b.	Human participants Vertebrate animals	□ Yes □ Yes	
	с.	Potentially hazardous biological agents (microorganisms, rDNA and	☐ Yes	
	d.	Hazardous substances and devices	□ Yes	🗆 No
3.	Wi	II this study be a sub-set of a larger study?	□ Yes	🗆 No

4. Will you directly supervise the student?

To be completed by the	Qualified Scientist:	To be completed by the D when the Qualified Scient
I certify that I have reviewed Project Summary prior to the the student or Direct Supervi procedures, I will This must k advice and super <b>BEFORE</b> to knowledge of the Start Date" Research Plan/Pr 1A.	and approved the Research Plan/ start of the experimentation. If sor is not trained in the necessary be dated will provide ne "Actual n. I have a working on Form y the student in the	SU If this box is need, then this approval must be dated I c BEFORE the "Actual Start Su on Form 1A. Direct Supervisor's Printed Name
Qualified Scientist's Printed Nam	e	Experience/Training of Designated
olghatalo		Phone email

<b>su</b> If this box is need, then this approval must be dated I c <b>BEFORE</b> the "Actual Start Da Su by on Form 1A.	ate" <sup>ch</sup> Plan/Project techniques to be used t supervision.
Direct Supervisor's Printed Name	
Direct Supervisor's Printed Name Experience/Training of Designated St	upervisor

□ Yes

**No** 

Degree(s):

### **Risk Assessment Form (3)**

Must be completed before experimentation; <mark>recommended for all projects</mark>. May be required for projects involving Human Participants, Hazardous Chemicals, Materials or Devices or Potentially Hazardous Biological Agents.

Student's Name(s)\_\_\_\_\_

Title of Project \_\_\_\_\_

Some of the answers to questions on this form (3) will be part of your Research Plan too. It is a good idea to be detailed on both the form and Research Plan.

To be completed by the Student Researcher(s) in collaboration with Direct Supervisor/Qualified Scientist: (All questions must be answered; additional page(s) may be attached.)

- 1. Identify and assess the risks and hazards involved in this project.
- 2. a) List all hazardous chemicals, activities or devices to be used; b) identify and list all microorganisms to be used that are exempt from pre-approval (see Potentially Hazardous Biological Agent rules).
- 3. Describe the safety precautions and procedures that will be used to reduce the risks.
- 4. Describe the specific disposal procedures that will be used (when applicable).
- 5. List the source(s) of safety information.

To be completed and signed b I agree with the risk assessment and Research Plan/Project Summary and direct supervision.	y the Direct Superviso safety precautions and pro- the International Rules, inc	or (or Qualified Scientist, whe <sup>Thi</sup> cedures described above. I certify t luding the science fair ethics stater 1A.	s must be dated FORE the "Actual rt Date" on Form
Direct Supervisor's Printed Name	Signature	Date of	Review (mm/dd/yy)
Experience/Training as relates to the studer	it's area of research		
Position/Institution		Phone or email contact information	

### Human Participants Form (4)

Required for all research involving human participants not at a Regulated Research Institution. If at a Regulated Research Institution, use institutional approval forms for documentation of prior review and approval. (IRB approval required before recruitment or data collection.)

Student's Name(s) Tit	le of Project			
Adult Sponsor Pr	Phone/Email			
MUST BE COMPLETED BY STUDENT RESEARCHER(S) IN COLLABORATION	I WITH THE ADULT SPONSOR/DIRECT SUPERVISOR/QUALIFIED			
<ol> <li>I have submitted my Research Plan/Project Summary which address Research Plan/Project Summary Instructions</li> </ol>	eses ALL areas indicated in the Human Participants Section of the			
2. I have attached any surveys or questionnaires I will be using in my	project or other documents provided to human participants.			
Any published instrument(s) used was /were legally obtained.	Even though your			
4. ☐ Yes ☐ No Are you working with a Qualified Scientist? If yes,	attach the Qualified Scientist Form 2.			
	<u>given approval, the</u> study/project must still			
DELUW - IND	conform to all ISEF			
<b>MUST</b> be completed by Institutional Review Board (IRB) after review	v of the research plan. All questions mu <b>regulations.</b>			
Approval to be valid. (If not approved, fetulit paper work to the studies	a) and the following conditions (All 6 must be ensured)			
1.         Risk Level (check one) :         Image: Communication of the second s	al Risk			
2 Qualified Scientist (OS) Required (Form 2)	(a risk assessment form 3 is required).			
3. Risk Assessment Required (Form 3):				
4. Written Minor Assent required for minor participants.	is form is to be filled out by the SCHOOL IRB and not the			
5. Written Parental Permission required for minor particited	gional/state fair SRC. However, be sure that your school			
	B is aware of the rules and limitations of student research piects			
6. Written Informed Consent required for participants Pro				
Fo	r more information and a full list of rules, visit: https://			
IRB SIGNATURES (All 3 signatures required) None of these indiv	ww.societyforscience.org/isef/international-rules/			
scientist or related to (e.g., mother, father of) the student (conflict o	f interest).			
determination and that I agree with the decisions above the second	oxes above have been completed to indicate the IRB			
Must be s Medical or Mental Health Professional (a psychologist, medic: the "Actua	gned prior to I Start Date" rker, licensed clinical profe			
physician's assistant, doctor of pharmacy, or registered nurse on Form 1	A s project.			
Printed Name	Degree/Professional License			
Signature/Date (prior to experimentation)	Email			
This CANNOT be the same teacher that h	as signed the student's Adult Sponsor			
Educator				
Printed Name	Degree/Professional License			
Signature/Date (prior to experimentation) Must be sign	ned prior to the "Actual Start Date" on			
School Administrator				
Printed Name	Degree/Professional License			
Signature/Date (prior to experimentation)	ust be signed prior to the "Actual Start Date" on			

International Rules: Guidelines for Science and Engineering Fairs 2024–2025, societyforscience.org/ISEF

Human Inf	ormed Consent Form	
Instructions to the Student Researcher(s): A consultation with the Adult Sponsor, Direct Supervis This form is used to provide information to the resea informed consent, minor assent, and/or parental per • When written documentation is required, • Students may use this sample form or ma	This is just a sample consent form. You MUST submit a copy of the consent form that you used, if applicable. If the a survey was done online, submit a copy of the consent question used as part of that survey.	be developed in nt written
If the form is serving to document parental permission	on, a copy of any survey or questionnaire must be a	ttached.
Student Researcher(s): Title of Project:		
I am asking for your voluntary participation in my sci project. If you would like to participate, please sign i	ence fair project. Please read the following informa n the appropriate area below.	tion about the
Purpose of the project:		
If you participate, you will be asked to:		
Time required for participation:		
Potential Risks of Study:		
Benefits:		
How confidentiality will be maintained:		
If you have any questions about this study, feel free t	to contact:	
Adult Sponsor/QS/DS:	_ Phone/email:	
<b>Voluntary Participation:</b> Participation in this study is completely voluntary. If consequences. Please be aware that if you decide to decide not to answer any specific question.	you decide not to participate there will not be nega participate, you may stop participating at any time	tive and you may
By signing this form I am attesting that I have read a assent to participate or permission for my child to pa	nd understand the information above and I freely gi articipate.	ve my consent/
Adult Informed Consent or Minor Assent	Date Reviewed & Signed: (mm/dd/yy)	
Research Participant Printed Name:	Signature:	

Date Reviewed & Signed: (mm/dd/yy)

Signature:

International Rules: Guidelines for Science and Engineering Fairs 2024–2025, societyforscience.org/ISEF

Parental/Guardian Permission (if applicable)

Parent/Guardian Printed Name:

### Vertebrate Animal Form (5A)

Required for all research involving vertebrate animals that is conducted in a school/home/field research site. (SRC approval required before experimentation.)

Student's Name(s)		
Title of Project		

#### To be completed by Student Researcher:

- 1. Common name (or Genus, species) and number of animals used.
- 2. Describe completely the housing and husbandry to be provided. Include the cage/pen size, number of animals per cage, environment, bedding, type of food, frequency of food and water, how often animal is observed, etc. Add an additional page as necessary.
- 3. What will happen to the animals after experimentation?
- 4. Attach a copy of wildlife licenses or approval forms, as applicable
- 5. The ISEF Vertebrate Animal Rules require that any death, illness or unexpected weight loss be investigated and documented by a letter from the qualified scientist, direct supervisor or a veterinarian. If applicable, attach this letter with this form when submitting your paperwork to the SRC prior to competition.

To be completed by Local or Affiliate Fair Scientific Review C	committee (SRC) BEFORE experimentation.	
Level of Supervision Required for agricultural, behavio	ral or nutritional studies (select one):	
Direct Supervisor REQUIRED. Please have applicable person	sign below.	
Veterinarian and Direct Supervisor REQUIRED. Please have applicable persons sign below.		
Veterinarian, Direct Supervisor and Qualified Scientist REQUIRED. Please have applicable persons sign below and have the Qualified Scientist complete Form (2).		
The SRC has carefully reviewed this study and finds it is an appropria Local or Affiliate Fair SRC Pre-Approval Signature:	te study that may be conducted in a non-regulated research site.	
SRC Chair Printed Name Signature	Date of Approval (must be prior to experimentation) (mm/dd/yy)	
<ul> <li>To be completed by Veterinarian:</li> <li>I have reviewed this research and animal husbandry with the student before the start of existing This must be</li> <li>I have approved the use and dosa BEFORE the drugs and/or nutritional suppleme "Actual Start Date"</li> <li>I will provide veterinary medical a on Form 1A. of illness or emergency. (Fees may apply.)</li> </ul>	<ul> <li>To be completed by Direct Supervisor or Qualified Scientist when applicable:</li> <li>I have reviewed this research This must be the student before the start accept primary responsibilit of the animals in this project</li> <li>I will directly supervise the e on Form 1A.</li> </ul>	
Printed Name Email/Phone	Printed Name Email/Phone	
Signature Date of Approval (mm/dd/yy)	Signature Date of Approval (mm/dd/yy)	

# Vertebrate Animal Form (5B)

Required for all research involving vertebrate animals that is conducted in at a Regulated Research Institution. (IACUC approval required before experimentation. Form must be completed and signed after experimentation.)

Student's Name(s)	
Title of Project	
Title and Protocol Number of IACUC Approved Project	:
To be completed by Qualified Scientist or Principal In	vestigator:
1. Species of animals used:	Number of animals used:

- 2. Describe, in detail, the role of the student in this project: animal procedures and related equipment that were involved, oversight provided and safety precautions employed. (Attach extra pages if necessary.)
- 3. Was there any weight loss or death of any animal? If yes, attach a letter obtained from the qualified scientist, direct supervisor or a veterinarian documenting the situation and the results of the investigation.
- 4. Did the student's project also involve the use of tissues?
  - 🛛 No
  - □ Yes; complete Forms 6A and 6B
- 5. What laboratory training, including dates, was provided to the student?

#### 6. Attach a copy of the Regulated Research Institution IACUC Approval. A letter from the Qualified Scientist

or Principal Investigator is not sufficient. You MUST include a copy of

You **MUST** include a copy of **the actual IACUC form** with the approval protocol number.

Must be dated **after** the "Actual End Date" on Form 1A

Qualified Scientist/Principal Investigator		
Printed Name		V
Signature	Date (mm/dd/yy)	

### Potentially Hazardous Biological Agents Risk Assessment Form (6A)

Required for research involving microorganisms, rDNA, fresh/frozen tissue (including primary cell lines, human and other primate established cell lines and tissue cultures), blood, blood products and body fluids. SRC/IACUC/IBC approval required before experimentation.

Student's Name(s)\_\_\_\_\_

Title of Project

To be completed by the QUALIFIED SCIENTIST/DIRECT SUPERVISOR in collaboration with the student researcher(s). All questions are applicable and must be answered; additional page(s) may be attached.

#### SECTION 1: PROJECT ASSESSMENT

- 1. Identify potentially hazardous biological agents to be used in this experiment. Include the strain, source, quantity and the biosafety level risk group of each microorganism.
- 2. Describe the site of experimentation including the level of biological containment.
- 3. Describe the procedures that will be used to minimize risk (personal protective equipment, hood type, etc.).
- 4. What final biosafety level do you recommend for this project given the risk assessment you conducted?
- 5. Describe the method of disposal of all cultured materials and other potentially hazardous biological agents. If BSL-2 laboratory, include the BSL-2 checklist.

#### **SECTION 2: TRAINING**

this section (4).

- 1. What training will the student receive for this project?
- 2. Experience/training of Direct Supervisor as it relates to the student's area of research (if applicable).

SECTION 3: For ALL CELL LINES Direct Supervisor - Check the a Experimentation on the n Research Institution, but for BSL-2). [This study ha to experimentation.]	5, MICROORGANISMS AND TISSUES – To be ppropriate box(es) below: nicroorganisms/cell lines/tissues to be used in t will be conducted at a (check one)BSL-1 or _ s been reviewed by the local SRC and the proce	his study will NOT be conduct BSL-2 laboratory (include a c edures have been approved pr	ED SCIENTIST or ed at a Regulated copy of the checklist ior
Experimentation on the n Research Institution and forms are attached. Origin of cell lines:	nicroorganisms/cell lines/tissues to be used in t was approved by the appropriate institutional b Date of IACL	his study will be conducted at oard prior to experimentation; JC/IBC approval	a Regulated institutional approval
Experimentation on the microorganisms/cell lines/tissues to be used in this study will be conducted at a Regulated Research Institution, which does not require pre-approval for this type of study. The SRC has seen and approved the research plan and supporting documentation and acknowledges the accuracy of the responses above.			
CERTIFICATION – To be SIGNED	) by the QUALIFIED SCIENTIST or Direct St	his must be dated	
The QS/DS has seen this project's research plan and supporting documentation provided above. This study has been approved as a (check one) $\square$ BSL-1/ $\square$ BSI aboratory. BEFORE the BEFORE the mathematical start Date of the information ucted in an appropriate on Form 1A			
QS/DS Printed Name	Signature	Date of review (mn	n/dd/yy)
SECTION 4: CERTIFICATION completed by the LOCAL or AFFILIATEDR SRC			
The SRC has seen this project's research plan and the provided provided provided acknowledges the accuracy of the information provided.			
SRC Printed Name	Signatur	Date of review (mn	n/dd/yy)
Do <b>NOT</b> write anything in 🚄			

International Rules: Guidelines for Science and Engineering Fairs 2024–2025, societyforscience.org/ISEF

### Human and Vertebrate Animal Tissue Form (6B)

Required for research involving fresh/frozen tissue (including primary cell lines, human and other primate established cell lines and tissue cultures), blood, blood products and body fluids. If the research involves living organisms please ensure that the proper human or animal forms are completed. All projects using any tissue listed above must also complete Form 6A.

Student's Name(s)\_\_\_\_\_

Title of Project \_\_\_\_\_

#### To be completed by Student Researcher(s):

- 1. What vertebrate animal tissue will be used in this study? Check all that apply.
  - □ Fresh or frozen tissue sample
  - □ Fresh organ or other body part
  - □ Blood
  - □ Body fluids
  - Primary cell/tissue cultures
  - □ Human or other primate established cell lines
- 2. Where will the above tissue(s) be obtained? If using an established cell line include source and catalog number.
- 3. If the tissue will be obtained from a vertebrate animal study conducted at a research institution attach a copy of the IACUC certification with the name of the research institution, the title of the study, the IACUC approval number and a copy of IACUC approval. If human tissues were used, attach a copy of IRB approval.

			This should be	
<ul> <li>To be completed by the Qualifier</li> <li>I verify that the student will work him/her by myself or qualified per were euthanized for a purpose of AND/OR</li> <li>I certify that the blood, blood prostandards and guidance set forth Pathogens.</li> </ul>	ied Scientist or Direct Sup solely with de-identified organs rsonnel from the laboratory; an her than the student's research ducts, tissues or body fluids in in U.S. Occupational Safety an	<b>Dervisor:</b> s, tissues, cultures o d that if vertebrate a this project will be h d Health Act, 29CFR	dated <b>BEFORE</b> the "Actual Start Date" on Form 1A. handled in accordance wit s Subpart Z, 1910.1030 - <u>Bl</u>	to ley h the <u>bod Borne</u>
Printed Name	Signature		Date of Approval (mm/o (Must be prior to experiment	dd/yy) ation.)
Title		Phone/Email		
Institution				

### **Continuation/Research Progression Projects Form (7)**

Required for projects that are a continuation/progression in the same field of study as a previous project. This form must be accompanied by the previous year's abstract and Research Plan/Project Summary.

Student's Name(s)			
<b>To be completed by Student Researcher:</b> List all components of the current project <b>For any projects</b> from previous research.			
Components	<b>Current Research Project</b>	Previous Research Project: Year:	
1. Title			
2. Change in goal/ purpose/objective			
3. Changes in methodology	Continuation pr form. For the im student researce the Abstract an years further ba include the Abs year's work.	ojects <b>MUST</b> include this mediately prior year, the ther MUST include BOTH d Research Plan. For any ack, the researcher MUST tract for each additional	
4. Variable studied	For ALL project or began before	cts that were conducted re January 1, 2024.	
5. Additional changes			

Attached are:

Abstract and Research Plan/Project Summary, Year \_\_\_\_\_

I hereby certify that the above in board properly reflect work done	formation is correct and that the current year Abstract e only in the current year.	& Certification and project display
Student's Printed Name(s)	Signature	Date of Signature (mm/dd/yy)