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):

| Stud | aent | dent's Name(s): | | | | | |
|------|-------|--|---|--|--|--|--|
| Proj | ect | ect Title: | | | | | |
| 1. | | and the control of th | le forms | | | | |
| 2. | | I have reviewed the student's completed Student Checklist (1A) and Research Plan/Project Summary. should make it easier to fit studen | | | | | |
| 3. | | ☐ I have worked with the student and we have discussed the possible risks involved in the project. and n | s, long titles | | | | |
| 4. | | □ The project involves one or more of the following and requires prior approval by an SRC, IRB, IACUC or DC: □ Humans Potentially Hazardous Biological Agents □ Vertebrate Animals □ Microorganisms □ TDNA □ Tissues | iore. | | | | |
| 5. | | Student Checklist (1) Approval Form (1B) | check the boxe are appropriate to research projec | | | | |
| | litio | litional forms required if the project includes the use of one or more of the following (check all that apply): ☐ Humans, including student designed inventions/prototypes. (Requires prior approval by an Institutional Reviews ee full text of the rules.) ☐ Human Participants Form (4) or appropriate Institutional IRB documentation ☐ Sample of Informed Consent Form (when applicable and/or required by the IRB) ☐ Qualified Scientist Form (2) (when applicable and/or required by the IRB) | w Board (IRB); | | | | |
| | | □ Vertebrate Animals (Requires prior approval, see full text of the rules.) □ Vertebrate Animal Form (5A)-for projects conducted in a school/home/field research site (SRC prior approval vertebrate Animal Form (5B)-for projects conducted at a Regulated Research Institution. (Institutional An Use Committee (IACUC) approval required prior experimentation.) □ Qualified Scientist Form (2) (Required for all vertebrate animal projects at a regulated research site or whom | imal Care and | | | | |
| | | Potentially Hazardous Biological Agents (Requires prior approval by SRC, IACUC or IBC, see full text of the rule of Potentially Hazardous Biological Agents Risk Assessment Form (6A) ☐ Human and Vertebrate Animal Tissue Form (6B)-to be completed in addition to Form 6A when project inverses or frozen tissue, primary cell cultures, blood, blood products and body fluids. ☐ Qualified Scientist Form (2) (when applicable) ☐ The following are exempt from prior review but require a Risk Assessment Form 3: projects involving prot similar microorganisms, for projects using manure for composting, fuel production or other non-culturing projects using color change coliform water test kits, microbial fuel cells, and projects involving decompoorganisms. | rolves the use of ists, archae and gexperiments, | | | | |
| | | ☐ Hazardous Chemicals, Activities and Devices (No SRC prior approval required, see full text of the rules.) ☐ Risk Assessment Form (3) ☐ Qualified Scientist Form (2) (required for projects involving DEA-controlled substances or when applicable | le) | | | | |
| | | Risk Assessment Form (3) | 'Actual Start | | | | |
| | | ☐ I attest to the information checked ab the project mentor (if those are different). Indicate the information checked about the project mentor (if those are different). | statement. | | | | |
| Adı | ılt S | ult Sponsor's Printed Name Signature Date of Review (mm, | /dd/yy) | | | | |
| Pho | ne | nne 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: c. Te | |
| 2. | Title of Project: | |
| | | the research mento |
| 3. | School:(if multiple schools, list of the team leader or list all schools). | School Phone: |
| Sc | hool Address: | |
| 4. | Adult Sponsor: | Phone/Email: |
| 5. | This is usually the teacher (or parent), not the research Does this project need SRC/IRB/IACUC or other pre-approval | |
| | Is this a continuation/progression from a previous year? a. If yes, attach the previous year's Abstract and b. Explain how this project is new and different from previous | ☐ Yes ☐ No ☐ Research Plan/Proje This defaults to "No" but change it for your project, if appropriate. |
| | ☐ Continuation/Research Progression Form (7); include for | rms for all previous years IF the student has |
| 7. | This year's experimentation/data collection (include forms forms for the state of t | continued their project |
| | | : (mm/dd/yy) |
| 8. | Where will you conduct your experimentation? (check all that | apply) |
| | ☐ Research Institution ☐ School ☐ Field ☐ Hom | e 🗆 Other: |
| 9. | Source of Data: ☐ Collected self/mentor ☐ Other List all URL(s) in Rese | 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-school virtually or on-site: | vork site(s), whether you worked there |
| Naı | me | |
| Ado | dress: | |
| | | |
| | | |
| 11. | Complete a Research Plan/Project Summary following the land attach to this form. | Research Plan/Project Summary instructions Every project must have a |
| 12. | An abstract is required for all projects after experimentation | detailed Describ Dlan that is |

Research Plan/Project Summary Instructions

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

- The Research Plan is to be written prior to experimentation following the instructions below to detail the rationale, research guestion(s), methodology, and risk assessment of the proposed research.
- If changes are made during the research prior to competing in an affiliated fair, such changes can be added to the original research plan as an addendum, recognizing that son The Research Plan is the most important document of appropriate review and approvals. If no additional approvals all because it provides the regional and state SRC the to explain research that was conducted.
- - Some studies, such as an engineering design or mathemat change through the course of research. If such changes of ossef.okstate.edu. and can be appended to the original research plan.
 - - why this research is important and if applicable, and if they contain the correct information.
 - b. RESEARCH QUESTION(S), HYPOTHESIS(ES), EN this based on the rationale described above?
 - c. Describe the following in detail:
 - · List of materials:
 - and when applicable, the source of data used. Desc tense!! done by your mentor.

 - Data Analysis: Describe the procedures you will use
 - d. BIBLIOGRAPHY: List major references (e.g. scien Must include all details on the work site. literature review. If you plan to use vertebrate ani reference.

necessary details of your planned research.

If no changes are made from the original research plan, no p There is a helpful template in the student resources page of

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 back to be able to determine if the proper forms were completed

> 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 The entire Research Plan must be written in FUTURE

• Risk and Safety: Identify any potential risks and safe It must include proposed AND actual start and end dates.

Must identify the student and mentor role.

Items 1-4 below are subject-specific guidelines for additional items to applicable.

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.

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.

Hazardous chemicals, activities & devices:

- a. Describe Risk Assessment process, supervision, safety precautions and specific methods of disposal.
- b. 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.

| 1. To Be Completed by Studen a. Student Acknowledgment: | t and Parent | t | |
|--|---|---|--|
| I understand the risks and po | ssible dangers t | o me of the proposed re | esearch plan. |
| · · · · · · · · · · · · · · · · · · · | - | | national Pulse when conducting |
| this research. | | | DEEODE the WAsterel |
| I have read and agree to upher | old all aspects o | f the student researcher | BEFORE the "Actual rethi Start Date" on Form 14 |
| Student researchers are expected to main | | | |
| misconduct are not condoned at any level | | | |
| plagiarism, forgery, use or presentation of | | | fabrication of data. Fraudulent |
| projects will fail to qualify for competition | in affiliated fairs | and ISEF. | / |
| | | This must be dated | \bigvee |
| | | BEFORE the "Actual | |
| Student's Printed Name | Signature | Start Date" on Form | Date Acknowledged (mm/dd/yy) |
| b. Parent/Guardian Approval: I have | o road and und | 1A | (Must be prior to experimentation.) |
| Research Plan/Project Summar | | | |
| Research Flan/1 Toject Summar | y. i consent to n | iy ciliid participating in | uns research. |
| | | | |
| Parent/Guardian's Printed Name | Signature | _ | Date Acknowledged (mm/dd/yy) |
| | | | (Must be prior to experimentation.) |
| a. Required for projects that need prior SPG BEFORE experii Do not write anyth potentially haza box (2a) unless y SRC/IRB chairpe Project Summary at designee. signature indicates approval of the Research I | c/IPR approval hing in this ou are the rson or | b. Required for resonance Student This proje Write an (not hom (2b)). by the proper institution complies with the ISE | earch conducted at all Regulated S/teachers do not ything in this box institution approved onar poard perore experimentation and F Rules. Attach (1C) and any required |
| Summary before the student begins experime | entation. | institutional approval | ls (e.g. IA <mark>CUC, IRB)</mark> . |
| | | | |
| SRC/IRB Chair's Printed Name | | | |
| $oldsymbol{arV}$ | | SRC Chair's Printed N | ame |
| Signature Date of Appro | val (mm/dd/yy) | | V |
| | experimentation.) | | |
| | | Signature | Date of Signature (mm/dd/yy) (May be after experimentation) |
| | | | |
| 3. Final ISEF Affiliated Fair SRC | Approval(Re | guired for ALL Proi | ects) |
| | | te anything in this b | |
| SRC Approval After Experi | | , , | |
| I certify that this project adheres to the appro | ved research ria | e SRC Chairs com | nplies with all locit kules. |
| | | | |
| | | | |
| Regional SRC Chair's Printed Name | Signature | E | Date of Approval (mm/dd/yy) |
| | | | |
| State/National SRC Chair's Printed Name | Signature | | Date of Approval (mm/dd/yy) |
| (where applicable) | 2.3 | | _ 2.0 0 |

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.

| St | udent's Name(s) | | | | |
|-----------|---|---------------------------------|----------------------|--------------------------|---|
| Ti | tle of Project | | | | |
| To (Re | If any of the research was done at a standard research facility (college or university, pharmaceutical company, environmental or biomedical testing facility, governmental facility, etc.) or a facility where advanced research is allowed (certain high schools or local labs), then Form 1C is required. | nt's project boo | _ | | on: t double-sided.) |
| | 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 | □ No □ No □ No □ No | default-clicked "no". Be sure to update these, if appropriate to your |
| 2. | If the project data are covered by privacy rules/laws (such as 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. | y the student goals of the p | in any phas | □ No se of the ¡ | |
| 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 deidentified. | collection, sp lly did. | ecific proce | edures pe | erformed). |
| 4. | Provide details regarding data provided to the student: | | | | |
| 5. | Did the student(s) work on the project as part of a group? Were there other high school students present? If yes, please was related or different from the work of this projecct. | | I Yes nts names a | □ No ind descr | |
| 6. | If this project is under a grant and needs to be acknolwedged | d, please list th | ne grant stat | tement ho | ere. |
| | This must be the mentor from the Regulated Research Institution NOT the teacher. The must be the mentor from the Regulated Research Institution plicity in competition and I have plicity in competition and | ched if applical | ble. I further | i nis musi | t be dated ne "End Date" any |
| | Direct Supervisor's Printed Name Signature | | Title | V | _ |
| | Institution | | | Signed (mu (mm/dd/yy) | st be after experimenta- |
| | Education/Experience/Training | | Email | /Phone | |

Qualified Scientist Form (2)

May be required for research involving human participants, vertebrate animals, potentially hazardous biological agents, and hazardous substances and devices. Must be completed and signed before the start of student experimentation.

| St | Student's Name(s) | | | | | | | |
|----|---|--------------------------|-------------------------|----------------------|--|--|--|--|
| Ti | Title of Project | | | | | | | |
| _ | | | | | | | | |
| To | be completed by the Qualified Scientis | st: | | | | | | |
| Sc | ientist Name: | | | | | | | |
| | ucational Background: | | | | | | | |
| Ex | perience/Training as relates to the student's | area of research: | | | | | | |
| | | | | | | | | |
| | | | | | | | | |
| _ | | | | | | | | |
| Po | sition/Institution: | Email/Phone: | | | | | | |
| 1. | Have you reviewed the ISEF rules relevant fair ethics statement relevant to this project | | ☐ Yes | □No | | | | |
| 2. | Will any of the following be used? a. Human participants b. Animals c. Potentially hazardous biological agents tissues, including blood and blood pro- | | ☐ Yes ☐ Yes ☐ Yes | □ No □ No □ No | | | | |
| | d. Hazardous substances and devices | adotaj | ☐ Yes | □ No | | | | |
| 3. | Will this study be a sub-set of a larger stud | y? | ☐ Yes | □ No | | | | |
| 4. | Will you directly supervise the student? | | ☐ Yes | □ No | | | | |
| 5. | Did you provide any data; if yes, please pro | ovide source or describe | ☐ Yes | □ No | | | | |

| To be completed by the Qualified Scientist: | | | | | | |
|---|--|--|--|--|--|--|
| I certify that I have reviewed and Project Summary prior to the stathe student or Direct Supervisor procedures, I will This must be advice and super BEFORE the "knowledge of the Start Date" on Research Plan/Pr 1A. | rt of the experimentation. If is not trained in the necessary lated will provide Actual n. have a working | | | | | |
| Qualified Scientist's Printed Name | \downarrow | | | | | |
| Signature | Date of Approval (mm/dd/yy) | | | | | |

| To be completed by the Direct Supervisor when the Qualified Scientist cannot directly | | | | | | | |
|---|---|--|--|--|--|--|--|
| st If this box is need, then this approval must be dated I c BEFORE the "Actual Start Date by on Form 1A. | ted Plan/Project techniques to be used t supervision. | | | | | | |
| Direct Supervisor's Printed Name | | | | | | | |
| Experience/Training of Designated Supervisor | | | | | | | |
| Signature | Date of Approval (mm/dd/yy) | | | | | | |
| Phone email | | | | | | | |

Risk Assessment Form (3)

Must be completed before experimentation; recommended for all projects. 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) Plan too. It is a good idea to be detailed on both the | |
| To be completed by the Student Researcher(s) in co Scientist: (All questions must be answered; additional pa | ge(s) may be attached.) |
| 1. Identify and assess the risks and hazards involved in this pro | Use additional pages, if you need more room to sufficiently explain responses to these questions. |
| 2. a) List all hazardous chemicals, activities or devices to be are exempt from pre-approval (see Potentially Hazardous I | |
| 3. Describe the safety precautions and procedures that will be include permits received and safety plans, as applicable. | be used to reduce the risks. If you conducted field work, |
| 4. Describe the specific disposal procedures that will be use | d (when applicable). |
| 5. List the source(s) of safety information. | |
| To be completed and signed by the Direct Supervision I agree with the risk assessment and safety precautions and pages Research Plan/Project Summary and the International Rules, in direct supervision. | rocedures described above a certity i |
| Direct Supervisor's Printed Name Signature | Date of Review (mm/dd/yy) |
| Experience/Training as relates to the student'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) | Title of Project | | | | | | |
|--|---|--|--|--|--|--|--|
| Adult Sponsor | Phone/Email | | | | | | |
| MUST BE COMPLETED BY STUDENT RESEARCHER(S) IN COLLABO | PRATION WITH THE ADULT SPONSO | PR/DIRECT SUPERVISOR/QUALIFIED | | | | | |
| SCIENTIST: 1. | h addresses ALL areas indicated in th | ne Human Participants Section of the | | | | | |
| Research Plan/Project Summary Instructions. 2. I have attached any surveys or questionnaires I will be usin | a in my proiect or other documents : | provided to human participants. | | | | | |
| ☐ Any published instrument(s) used was /were legally ob | | Even though your | | | | | |
| 3. I have attached an informed consent that I would use if required to the state of the state | | school IRB may have | | | | | |
| 4. Yes No Are you working with a Qualified Scientist | ? If yes, attach the Qualified Scientis | given approvai, the | | | | | |
| BELOW | – IRB USE ONLY | study/project must still | | | | | |
| MUST be completed by Institutional Review Board (IRB) after review | | conform to all ISEF——nust be answer regulations. | | | | | |
| valid. (If not approved, return paperwork to the student with instruc | · | | | | | | |
| ☐ Approved with Full Committee Review (3 signatures required 1. Risk Level (check one): | Minimal Risk | l 6 must be answered) re than Minimal Risk risk assessment form 3 is required). | | | | | |
| 2. Qualified Scientist (QS) Required (Form 2): | Yes | risk assessment form 3 is required). | | | | | |
| Risk Assessment Required (Form 3): Written Minor Assent and written parental permission | Yes | | | | | | |
| ☐ Yes ☐ Not applicable (No r | ninors in this study) | | | | | | |
| 5. Written Informed Consent required for participants ≀8 ☐ Yes ☐ No ☐ | | | | | | | |
| 6. Facility for "protected groups" used, written approval | has I IRR is aware of the rules a | wever, be sure that your school | | | | | |
| L les L NO | projects | The infinite of the definition | | | | | |
| IRB SIGNATURES (All 3 signatures required) None of these scientist or related to (e.g., mother, father of) the student (e.g., | indiving the same of the same | | | | | | |
| scientist or related to (e.g., mother, father of) the student (co | For more information and a | a full list of rules, visit: https:// | | | | | |
| I attest that I have reviewed the student's project, that the determination and that I agree with the decisions above. | www.societyforscience.o | rg/iset/international-rules/ | | | | | |
| · · | | | | | | | |
| Medical or Mental Health Professional (a psychologist, medical do | ctor, licensed social worker, license | d clinical professional counselor, | | | | | |
| physician's assistant, doctor of pharmacy, or registered nurse Mu | | | | | | | |
| line and the second sec | "Actual Start Date" ^{cense} Form 1A | | | | | | |
| Signature | Date (prior to experimentation) | Email | | | | | |
| orginature | bute (priprite experimentation) | Email | | | | | |
| | V | | | | | | |
| | | | | | | | |
| Educator This CANNOT be the same teache | r/parent that has signed as the | student's Adult | | | | | |
| Print Name below Sponsor. | | | | | | | |
| | | | | | | | |
| Signature | Date (prior to experimentation) | Email | | | | | |
| | K | | | | | | |
| | Must be signed prior to the "Actual Start Date" on | | | | | | |
| | Form 1A | | | | | | |
| School Administrator | | | | | | | |
| Print Name below | Degree/Professional License | | | | | | |
| | Data (natan) | Final | | | | | |
| Signature | Date (prior to experimentation) | Email | | | | | |
| | Must be signed prior to the | Actual Staff Date Off | | | | | |

Human Informed Consent Form

Instructions to the Student Researcher(s): An MUST submit a copy of the consent form consultation with the Adult Sponsor, Direct Superviso that you used, if applicable. If the a survey This form is used to provide information to the researce was done online, submit a copy of the

Student Researcher(s):

Title of Project:

This is just a sample consent form. You informed consent, minor assent, and/or parental perm consent question used as part of that

When written documentation is required, to the consent question used as part of that

be developed in

nt written

• Students may use this sample form or may Survey.

| | $h \cap 1$ | torm ic | corving to c | locument paranta | Inarmicciar | $1 - 0 - 0 - 0 \cdot 1 - 0 \cdot$ | t anv curvav a | r alloctionnaire milet | no ottoo | $n \cap A$ |
|---|------------|---------|--------------|------------------|-------------|-----------------------------------|-----------------|---|----------|------------|
| ı | 115 | IOHH IS | SELVING TO C | iocument parenta | | 1. a (JUDV U | i aniv survev u | r questionnaire must | DE allac | HEU |
| | | | | | . p | ., | , | . 9000000000000000000000000000000000000 | | |

| am asking for your voluntary participation in my science fair project. Please read the following information about the project. If you would like to participate, please sign in the appropriate area below. | | | | | | |
|--|---|--|--|--|--|--|
| 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 | to contact: | | | | | |
| Adult Sponsor/QS/DS: | _ Phone/email: | | | | | |
| 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 negative participate, you may stop participating at any time and you may and understand the information above and I freely give my consent/ | | | | | |
| assent to participate or permission for my child to p | | | | | | |
| Adult Informed Consent or Minor Assent | Date Reviewed & Signed:(mm/dd/yy) | | | | | |
| Research Participant Printed Name: | Signature: | | | | | |
| Parental/Guardian Permission (if applicable) | Date Reviewed & Signed:(mm/dd/yy) | | | | | |
| Parent/Guardian Printed Name: | Signature: | | | | | |
| Page 38 Internat | ional Rules: Guidelines for Science and Engineering Fairs 2025–2026, societyforscience.org/ISEF | | | | | |

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 Stude | nt Researcher: | | | | | | |
| 1. Common name (or Genus | Common name (or Genus, species) and number of animals used. | | | | | | |
| per cage, environment, be | 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 ar | nimals after experimentation | ? | | | | | |
| 4. Attach a copy of wildlife li | censes or approval forms, as | applicable | | | | | |
| documented by a letter fro | | rect supervisor or a veterin | eight loss be investigated and arian. If applicable, attach this ition. | | | | |
| ☐ Veterinarian and Direct S | ed for agricultural, behavio ED. Please have applicable person upervisor REQUIRED. Please have ap rvisor and Qualified Scientist REQU mplete Form (2). Is study and finds it is an appropria | ral or nutritional studies (s sign below. plicable persons sign below. JIRED. Please have applicable per stee study that may be conducted Date of App | elect one): rsons sign below and have | | | | |
| the student before the sta | ch and animal husbandry with the of extended dosa BEFORE the ppleme "Actual Start Date" edical a on Form 1A. | Scientist when applica I have reviewed this rethe student before the accept primary respo | esearci This must be he start BEFORE the project "Actual Start Date" | | | | |

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 | | | | |
| | | | | |
| 1. Species of animals used: | Number of animals used: | | | |
| | roject: animal procedures and related equipment that autions employed. (Attach extra pages if necessary.) | | | |
| Was there any weight loss or death of any animal? scientist, direct supervisor or a veterinarian docur | If yes, attach a letter obtained from the qualified nenting the situation and the results of the investigation. | | | |
| 4. Did the student's project also involve the use of tis □ No □ Yes; complete Forms 6A and 6B | ssues? | | | |
| 5. What laboratory training, including dates, was pro | vided to the student? | | | |
| | | | | |
| or Principal Investigator is not sufficient. You MU the actu | on IACUC Approval. A letter from the Qualified Scientist ST include a copy of ual IACUC form with roval protocol number. Must be dated after the "Actual End Date" on Form 1A | | | |
| Qualified Scientist/Principal Investigator | | | | |
| Printed Name | | | | |
| 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 Identify potentially hazardous biological agents to be used in this experiment. In and the biosafety level risk group of each microorganism. | clude the strain, source, quantity | | | |
| 2. Describe the biosafety level of the experimentation site. | | | | |
| 3. Describe the procedures that will be used to minimize risk (personal protective e | equipment, safety cabinet type, etc.). | | | |
| Describe the method of disposal of all cultured materials and other potentially hazardous biological agents. If BSL-2 laboratory, not at an RRI, include the BSL-2 checklist | | | | |
| SECTION 2: TRAINING 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, MICROORGANISMS AND TISSUES – To be comp | leted by the QUALIFIED SCIENTIST | | | |
| or Direct Supervisor - Check the appropriate box(es) below: Experimentation on the microorganisms/cell lines/tissues to be used in this study will NOT be conducted at a Regulated Research Institution, but will be conducted at a (check one)BSL-1 orBSL-2 laboratory (include a copy of the checklist for BSL-2 . [This study has been reviewed by the local SRC and the procedures have been approved prior to experimentation.] | | | | |
| This project involves the culturing of Multi Drug Resistant Organisms (MDROs). or higher lab at a Regulated Research Institution and the required IBC pre-appropriate of IBC approval | | | | |
| Experimentation on the microorganisms/cell lines/tissues to be used in This makes arch Institution and was approved by the appropriate institutional approval forms are attached. Origin of cell lines: Date of IBC On For | RE the lation; institutional I Start Date" | | | |
| Experimentation on the microorganisms/cell lines/tissues to be used will be corwhich does not require IACUC or IBC approval for this type of study. | . \/ | | | |
| CERTIFICATION - To be SIGNED by the QUALIFIED SCIENTIST or Direct Superviso | r | | | |
| The QS/DS has seen this projects. The provided above. This study has been approved. The provided above above approved. The provided above approved above above approved above above approved above above approved above | knowledges the accuracy of the information , and will be conducted in an appropriate | | | |
| Do NOT write anything in | Date of review (mm/dd/yy) | | | |

this section (4).

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 R | Researcher(s): | | |
| 1. What vertebrate animal tissue w Fresh or frozen tissue sand result fresh organ or other bood Blood Body fluids Primary cell/tissue culture Human or other primate | mple dy part res | eck all that apply. | |
| 2. Where will the above tissue(s) | be obtained? If using an | established cell line ir | iclude source and catalog number. |
| | ne name of the research i | institution, the title of t | esearch institution attach a copy of the study, the IACUC approval num- of IRB approval. |
| To be completed by the Qua ☐ I verify that the student will wo him/her by myself or qualified were euthanized for a purpose AND/OR ☐ I certify that the blood, blood p standards and guidance set for Pathogens. | rk solely with de-identified of personnel from the laborate other than the student's respondents, tissues or body flu | organs, tissues, cultures bry; and that if vertebrate search. uids in this project will be | e ar <mark>on Form 1A. h</mark> ey |
| Printed Name | Signature | | Date of Approval (mm/dd/yy) (Must be prior to experimentation.) |
| 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) | | | | |
|--|---|--|--|--|
| To be completed by Studen previous research. | t Researcher: List all components of the curr | rent project. For any projects carried out (partially) before 2025. | | |
| Components | Current Research Project | Previous Research Project: Year: | | |
| 1. Title | | | | |
| 2. Change in goal/ purpose/objective | | | | |
| 2 Changes in | form. For the im | ojects MUST include this mediately prior year, the | | |
| 3. Changes in methodology | the Abstract and years further ba | student researcher MUST include BOTH the Abstract and Research Plan. For any years further back, the researcher MUST include the Abstract for each additional year's work. | | |
| 4. Variable studied | | ets that were conducted re January 1, 2025. | | |
| 5. Additional changes | | | | |
| | | | | |
| uttached are: Previous year's Abstra Previous Form 7s, if ap | ct and Research Plan/Project Summary, oplicable. | Year | | |
| | ove information is correct and that the currer done only in the current year. | nt year Abstract & Certification and project display | | |
| Student's Printed Name(s) | Signature | Date of Signature (mm/dd/w) | | |