An Adverse Event (AE) is any unfavorable and unintended sign, symptom, or disease associated with the research protocol. The AE could be experienced by a human or by an animal. An example of a human AE would be an investigator having an allergic reaction to a rat. An example of an animal AE would be the unanticipated death of a rat.

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| **PART I: PROTOCOL AND**  **EVENT TYPE** | |
| 1. IACUC Protocol Number: | |
| 2. IACUC Protocol Title: | |
| 3. Date this report completed: | |
| 4. Date of adverse event: | |
| **DEMOGRAPHICS** | |
| **PI Name** |  |
| PI Telephone Number |  |
| PI E-mail Address |  |
| **PART II: DETAILED ADVERSE EVENT INFORMATION** | |
| **Was this a human event? (use separate sheet to report if necessary)**  Severity/ Seriousness of the AE :  Minimal \_\_\_\_ Moderate \_\_\_\_ Life-threatening \_\_\_ Fatal \_\_\_\_\_\_  Required intervention to prevent permanent impairment/damage \_\_\_\_  Describe the adverse event and its relation to the protocol:  Were the procedures of the protocol deviated from in any way?  Describe treatment taken to address the AE.  If a student was injured, did the injured student report to Student Health Services?  What was the date(s) of treatment?  Was injury due to contact with animals in protocol?  Will the AE require follow-up treatment?  Will the protocol change as a result of the AE? If so state how protocol will change  What action will be taken to prevent an AE similar to this?    What was the outcome of the event? (resolved, unresolved, etc.)  Has CITI animal training certification been taken by person experiencing AE? Person completing this form?  Was the animal users risk assessment form completed? | |
| **Was this an animal event? (use separate sheet to report is necessary)**  Severity/ Seriousness of the AE :  Minimal \_\_\_\_ Moderate \_\_\_\_ Life-threatening \_\_\_ Fatal \_\_\_\_\_\_  Required intervention to prevent permanent impairment/damage \_\_\_\_  Describe the adverse event and its relation to the protocol:  Were the procedures of the protocol deviated from in any way?  Describe the treatment taken to address the AE.  Will the protocol change as a result of the AE? If so state how protocol will change.  What action will be taken to prevent an AE similar to this?  If fatal, was a necropsy performed?  What was the outcome of the event? (resolved, unresolved, etc.)  Has the CITI animal training been completed by person completing this form? | |
| Documentation accompanying the report  (e.g.Notes, Discharge Summary, Lab or Necropsy Reports, Other, etc.) | |

Name (person completing this report) \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

Relation of above to the protocol\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

Date \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

Phone \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

Email \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_