November 20, 2019

Chris Cramer, Ph.D. Vice President for Research and Institutional Official 419 Johnston Hall 101 Pleasant Street SE Minneapolis, MN 55455

Dear Dr. Cramer,

Attached is the official Fall 2020 Semiannual Report of the Institutional Animal Care and Use Committee. The IACUC concluded its semiannual evaluation of the institution's animal care and use program on October 22, 2019 using the Guide for the Care and Use of Laboratory Animals (Guide), the Guide for the Care and Use of Agricultural Animals in Agricultural Research and Teaching (Ag Guide), the refined OLAW checklist, and as applicable, the Animal Welfare Act Regulations as the basis for its evaluations. The IACUC met to discuss the results of semiannual inspections, conduct a Program Review, and evaluate other features of the animal care and use program.

The IACUC inspected all animal facilities and holding areas and many research laboratories; additionally, post-approval monitoring was conducted by IACUC compliance officers and committee members. Significant and minor deficiencies were documented. All of these deficiencies have been corrected at the time of this report. Please see the attached reports for details of each area inspected.

Please see the program review meeting notes for further detail on the issues discussed by the IACUC during this program review session.

Thank you for your continued support of IACUC activities.

# November 19, 2019 Program Review

Name Signature Dezhi Liao SCOTT MADILL Scott Opper Yeggy Norris Anna Robert Schumacher -ynn Impelluso Minie Vallary The Poltas Burnel Cooper BRUN A. CROOKER Ben Clark Keith Barker attended online Georgiy Aslanid! on

Please sign in below

# Fall 2019 Semiannual Program Review October 22, 2019

#### Voting Member Attendees:

Keith Barker, Dezhi Liao, Melanie Graham, Bob Schumacher, Sally Noll, Brian Crooker, George Aslanidi, Lynn Impelluso, Kristin Pilon, Nate Koewler, Don Martín, Marilyn Bennett, Henry Wong, Scott Madill,

# Alternate Member Attendees and Guests:

Frances Lawrenz, Laura Hocum-Stone, Geoff Ghose, Kakambi Nagaraja, Cynthia Lee, Ilana Cohen, Megan McCoy, Paul Lindstrom, Jennifer Borgert, Jodi Ogilvie, Margaret Luesse, Christine Sivula, Dan Montonye, Jen Hubbard, **Margaret Luesse**, Felicia Boynton, **Margaret Luesse**, Whitney McGee

# 1. Agenda:

- Inspection Summary (pg 2-4)
- IMHA Summary (pg 4)
- PAM Frequency Reduction Implementation Plan (pg 4-5)
- IACUC Office Administrative Summary (pg 6-7)
- Program Discussion Summary (from FCR Meetings) (pg 8-15)
- Cephalopod subcommittee update (pg 15)
- eProtocol Form Updates (pg 16-30)
- Pain classification (pg 31)
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- Health and Monitoring: does frequency of monitoring include RAR and lab monitoring frequency combined? (pg 32)
- Pre-worded answers available to PIs for common procedures or treatments (pg 32)
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- RAR training offered (pg 32-33)
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  - b. Inspection Dates (pg 49-51)
  - c. IMHA List and Justifications (pg 52-82)
  - d. Reduced PAM (pg 83-88)
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  - g. Administrative Summary and Graphs (pg 295-312)
  - h. Expired and Holding Protocol Summary (pg 313)
  - i. Repeat Significant Findings (pg 314-318)

#### 2. Inspection Summary

The Institutional Animal Care and Use Committee Senior Compliance Officer, Kristin Pilon, presented the Fall 2019 IACUC inspection report.

There were 286 inspections resulting in 201 findings for this six-month cycle. There were 166 Minor Findings and 35 Significant Findings. Twenty percent of the Significant Findings (7) were reported to OLAW.

- 7 reports sent to OLAW
- 192 laboratory areas that had no findings
- 5 repeat findings with three repeat significant findings (one of the same finding and two in the same laboratory but different types of significant findings noted)
- 13 notes to file
- 18 veterinary recommendations
- 39 laboratories that qualified for reduced post approval monitoring frequency

	Fall 2019 April 2019— September 2019	Spring 2019 October 2018— March 2019
Significant	35	29
Minor	166	165
Total	201	194



#### NOTE: Additional data and graphs located in appendices

There was a slight increase in percentage in total findings this six-month cycle from last six month cycle (0.04% increase—194 findings in Spring 2019 with 201 findings this six month cycle).

The increase can be attributed to the increase in significant findings, as the percent increase in minor findings was negligible.

At the time of Program Review, both the significant and minor findings had been corrected and the reports closed.

There was an increase (21%) in significant findings this past cycle (35 findings found during this sixmonth period and 29 findings noted during the Spring 2019 six-month reporting period). Of these 35 significant findings noted during this six-month cycle, five of them (14%) came from either self- reports or outside reports of non-compliance that came into the IACUC whereas 30 findings came from our standard inspection process during the last six months. Additionally, there were three investigators with repeat significant findings (one laboratory had the same finding and two laboratories had different significant findings) noted on their most recent inspections. An email was sent to each of these investigators offering guidance and help to better improve compliance in their laboratories.

There were three areas for which we saw the greatest increase in significant findings this six-month cycle. This includes a 400% increase for which animals were housed outside of RAR facilities without approval (4 found this six month cycle with no findings of this type noted in the Spring 2019 cycle), 500% increase for which anesthetic procedures were conducted on animals but not approved (5 findings of this type found this six month cycle with no finding of this type noted in Spring 2019) and a 700% increase for which analgesics were not given for the type or duration approved in the protocol (8 findings noted this six month cycle with only two of this type of finding noted in Spring 2019). The IACUC will be focusing efforts in order to try to minimize the number of analgesic deficiency type findings by sending out list serve announcements reminding investigators to pay closer attention to the timing of SR-buprenorphine or Carprofen administration.

Of these significant findings, fifteen of the thirty-five finding could be considered animal welfare issues (43%).

The increase in the Minor Findings category for this six-month reporting cycle was negligible with 165 findings noted in Spring 2019 and 166 findings noted this Fall 2019 cycle. The following categories that saw the increases were the following: Expired Items (4 findings of this type noted this six month cycle with 1 noted during the Spring 2019 cycle, Husbandry (14 noted this six month cycle with 6 noted in Spring 2019), and Occupational health program requirements not met (13 noted this six month cycle with 11 noted in Spring 2019).

During the six month cycle, Committee members have been alerted as to the uptick in aseptic technique findings. While we have seen an increase, it was still less than the last six month cycle (13 findings of this type noted this six month cycle with 18 noted in Spring 2019). Compliance staff will continue to distribute the RAR training that is offered so as to mitigate these types of findings. Additionally, the area which saw the greatest decrease in findings were the Ag areas with 6 findings noted this six month cycle and fifteen noted Spring 2019 (60% decrease).

Within the program review documents, findings have been defined by type (IACUC (178), DEHS (0), OHS (16) and Ag (7) as well as from what inspection/report they were identified or came from (PAM (100), Second Surgery (18), Initial Surgery (5), Initial surgery/semi-annual (0), Semi-annual (20), PAM/Semi-annual (27), Second Surgery/semi-annual (13), Ag (7), Self-Report (10), Outside Report (1)).

As stated previously, we had 286 inspections during this six-month period. These inspections have been categorized as to type performed with 117 Post Approval Monitoring (PAM) Inspections, 28 combined PAM/semi-annual, 36 Second Surgery inspections, 8 Second Surgery/Semi-annual inspections, 74 semi-

annual inspections and 16 Ag inspections conducted over this six-month period. Additionally, as part of our reduced PAM inspections we had 6 initial surgery inspections and 1 initial surgery/semi-annual inspections.

There was a slight increase in the number of laboratories that had no findings from the Spring 2019 sixmonth cycle versus this Fall 2019 cycle (190 to 192 respectively).

There was a slight decrease in the number of repeat findings found during inspections this last six month cycle (six noted in Spring 2019 and five noted this six month cycle). There were two repeat findings in the minor category with three significant repeat findings.

An overview of the Notes to File was given. These thirteen Notes to File were reported on a monthly basis during a full committee review meeting to confirm that these changes should not have required a submission from the PI. These Notes were obtained either through a submission to the IACUC office or through an inspection of the laboratory for which additional information was noted but did not change the procedures that were originally reviewed and approved by the IACUC.

There were eighteen veterinary recommendations this reporting period. These recommendations are changes to protocol for which the veterinary team approves while an amendment is submitted by the PI in cases where the change will be for the betterment of the animal. These veterinary recommendations were reviewed during the same time the monthly findings report and notes to file are discussed.

# 3. Current IMHA Summary

An overview of the IMHA spaces with justification for housing was presented to the IACUC at this program review. We have 120 PIs that have requested and approved housing of animals outside of Research Animal Resources in our tracking database. These PIs utilize 65 different areas.

# 4. PAM Frequency Reduction Implementation Plan (OVPR Risk Recalibration Initiative)

Reduced Post Approval Monitoring frequency was initiated in April 2012, after that Spring Program Review.

There were 145 PIs that did not qualify for reduced post approval monitoring inspections and had their post approval monitoring (PAM) or combined PAM/Semi-annual inspections completed due to not meeting our standard qualifications of reduced frequency. There were 39 PIs/laboratories that were due for post approval monitoring inspections but who did not receive their inspections as they qualified for reduced frequency (so either did not have an inspection or if surgery was conducted in the laboratory, only had a surgery portion of the visit and the PAM portion was not conducted). Additionally, there were 74 PIs that were due for PAM but either did not receive an inspection as they did not have any active studies, were put on hold (inactivation) due to no response at continuing review or had put their studies on hold due to limited funding or performing solely data analysis.

As background, post approval monitoring inspections are normally conducted yearly for all PI's who are currently performing experimental procedures on animals. The only PI's that do not currently receive yearly PAM inspections include: Fisheries and Wildlife studies except those that house animals on campus, Client Owned animal studies and Agricultural studies. The proposal includes reduction of PAM visit to every other year for those areas that meet the following criteria:

• No significant findings within the last two years

- No concerns or complaints made of the laboratory through RAR, RAR veterinarians, outside entities or through compliance staff concerns
- No repeat findings on most recent inspection
- No minor findings noted from most recent inspection other than ROHP non-compliance
- No additions of a different species to be used in the laboratory (either on original protocol or through a new submission)

If any of the above are noted, the laboratory would go back onto yearly PAM visits until all criteria are again met.

Laboratories that are considered "Euthanasia and Tissue Harvest" would be able to go onto this schedule if above criteria are met. Our office will continue to perform mandatory semi-annual inspections of laboratories that conduct either non-survival or survival surgical procedures for compliance. Those laboratories in good standing would receive twice-yearly inspections of surgical areas and procedures, records, aseptic technique, anesthetics/analgesics used, euthanasia methods, endpoints, ROHP compliance, etc...but would NOT receive a PAM inspection portion during these visits.

# 4. Administrative Statistics for Fall Program Review 2019:

- Total FCR submissions April 2019 September 2019: 65
- Total DMR submissions April 2019 September 2019: 531
- Review Outcomes:

# **FCR**

Number of new protocols: 51 Number of new protocols that received stipulations: 45 Number of new protocols that were approved as submitted: 5 Number of new protocols that were deferred: 1

Number of amendments: 14 Number of amendments that received stipulations: 11 Number of amendments that were approved as submitted: 2 Number of amendments that were deferred: 1

# <u>DMR</u>

Number of new protocols: 185 Number of new protocols that received stipulations: 149 Number of new protocols that were approved as submitted: 34 Number of new protocols that were sent to FCR: 1

Number of amendments: 346 Number of amendments that received stipulations: 205 Number of amendments that were approved as submitted: 138 Number of amendments that were sent to FCR: 0

• Median Approval Times for submission from October, 2018 – March 2018

# <u>FCR</u>

<u>New Protocols:</u> Days from receipt of submission to meeting: 8 Days from meeting date to initial letter sent: 2 Days from stips sent to responses received: 6 Days from submission to approval: 28

Amendments:

Days from receipt of submission to meeting: 10 Days from meeting date to initial letter sent: 2.5 Days from stips sent to responses received: 2 Days from submission to approval: 20

# DMR:

New Protocols:

Days from receipt of submission to agenda assignment: 2 Days from agenda assignment to all reviews received: 9 Days from reviews received to first letter sent: 2 Days from stips sent to responses received: 4 Days from submission to approval: 26

# Amendments:

Days from receipt of submission to agenda assignment: 3 Days from agenda assignment to all reviews received: 8 Days from reviews received to first letter sent: 3 Days from stips sent to responses received: 3 Days from submission to approval: 20

# 5. <u>Compilation of IACUC Discussion Notes April 2019 – September 2019</u>

# • INSTITUTIONAL PRACTICES, POLICIES, AND RESPONSIBILITIES

**APR.** The committee discussed a study that will use octopi. Although there are no current regulations in the U.S. mandating IACUC review of this work as they are invertebrates, due to the intelligence of the species, European institutions and AAALAC have begun to require a review of these studies. The committee discussed the issue further at program review and has established a subcommittee to research the topic and bring a recommendation back to the IACUC.

**AUG.** The committee discussed the previous approval of an exception for the use of unopened sterile saline bags for up to six months after opening. The exception had been granted in response to a shortage. Now that production and supply are back to normal, the temporary use of expired saline has been terminated. An announcement to PIs will be sent via the PI Google Groups.

# • PROTOCOL DISCUSSION AND REVIEW

**JUL.** The committee reviewed a proposal to update the text in the Breeding Procedure of the protocol form. The change would eliminate the question "What are the criteria for euthanasia of the Breeders?" and instead provide recommendations for euthanasia of rodent breeders (see below). Other links or recommendations may be provided in the future for other species.

"Recommended Mouse Breeder Retirement Criteria:

- They produce no litter within 60 days of mating (longer may be acceptable if delayed breeding is a strain characteristic).

- They produce no litter within 60 days of their last litter and are not visibly pregnant.

- They produce litters but do not wean pups for two to three litters."

The committee approved this change and we will strive to update the protocol form by 2019 Fall Program Review.

**JUL.** The committee discussed DEHS requirements for IACUC protocols. The committee was reminded that DEHS does not require inclusion of the following chemicals or SOPs for the following chemicals:

- CM
- 25 SO
- formaldehyde

- Isoflurane
- Luciferin
- MS-222
- LPS

# • SELF-REPORTS and OUTSIDE REPORTS

**APR.** The committee reviewed a self-report in which two mice were anesthetized with ketamine/xylazine in preparation for transurethral infection with E. coli. Following the procedure, cages were placed on heating pads to aid recovery, but the lab failed to return the animals to the housing rack. RAR returned them the following morning.

The following corrective action plan has been implemented to ensure that this does not happen again:

1. All Syntiron employees that are approved to handle animals will be informed about post-anesthesia monitoring requirements and directed to re-read the appropriate section of the IACUC protocol.

2. For any procedures that require anesthesia, one person from the Syntiron animal team will remain at the University to monitor the mice continuously until they recover (are mobile and able to access water).

3. If the monitor needs to leave the animal room, it will be for no more than 15 minutes, in accordance with current IACUC policy.

The committee had no additional concerns and considers the matter closed.

**APR.** The committee was updated on a self-report in which two client owned naïve goat kids were delivered but then returned to the client without being used on the study. They were replaced with two healthier animals. The PI transported the animals rather than RAR or the client as neither was available at the time. In the future the protocol will be amended to allow for transport by the investigator under special circumstances and with permission of RAR veterinarians. The committee had no additional concerns and considers the matter closed.

**MAY** The committee discussed a U Report that alleged an investigator had staff do procedures for outside biotech companies that were not approved on protocol, housed animals in his laboratory, took animals home, and pressured lab staff into not sharing information with IACUC. The IACUC chair and vice chair met with the PI and will attempt to contact lab staff regarding this allegations. The committee will be updated as additional information is collected.

**MAY** The committee discussed an RAR self-report regarding a rabbit kit that was found out of its cage. The kit appeared dehydrated and was administered SQ fluids and dextrose. It is believed the animal was out of the cage for less than 48 hours. There had been

additional equipment temporarily stored in the room due to flooding that had occurred in another area that may have prevented animal care staff from identifying the kit outside of the cage. Now that the animals are in a housing room without additional equipment and the PI is marking the card each time he takes a kit RAR feels that they can quickly recognize and address escaped animals moving forward and do not anticipate this happening again in the future. The committee suggested that RAR be reminded not to allow rooms to become cluttered and considers the matter closed.

**MAY** The committee was updated regarding a self-report in which a cranial window was applied to mice receiving cranial injections. While the surgical procedure was approved, the protocol listed closing with sutures and did not include the option for a cranial window. There were no adverse effects identified on these animals. The PI has submitted an amendment to account for this possibility in the future and the committee considers the matter closed.

**MAY** The committee was updated on a lab that contacted the IACUC office and area veterinarian to try perfusing a group of mice to improve immunolabeling. The lab perfused a group of animals under supervision of the area vet and personnel trained in perfusions. The lab has since updated the protocol to include perfusion for future animals. The committee had no additional concerns and consider the matter closed.

**MAY** The committee was updated on minor change that was approved by the area vet to allow an extension of single housing for mice for 2 additional days. The committee had no additional concerns and consider the matter closed.

**MAY** The committee discussed a self-report regarding the occurrence of frostbite after hypothermia induced anesthesia. After discussing this issue with the RAR veterinarian, the lab is planning to take the following steps to try and minimize the occurrence and extent of frostbite.

• "Slower rewarming. Rewarming is currently done by holding the pup under a heat lamp until they start breathing, and then placing them in a cage that is put on a heated pad. We discussed 3 strategies to slow down the rewarming process.

a. hold the pup further away from the heat lampb. warm the pup up by holding it in your hands until it starts to breathc. slow down rewarming by skipping heat lamp or hand held rewarming,and instead placing the pup in a heated cage (ideally 33°C)

- Protect the tail by applying vaseline before inducing anesthesia
- Change induction of anesthesia to a method where water is not added to the ice, and keep the pups on ice during the surgery. I mentioned how I would prefer to try this as a last resort strategy, because it doesn't lower the core temperature down to a complete cessation of heart beat, and this inherently has a higher risk of mortality when we resect the apex of the heart (due to bleeding out). "

The committee had no additional comments and considers the matter closed.

**MAY** The committee discussed an RAR self-report. On the weekend of May 4-5 RAR was responsible for providing post-operative care to an NHP that had undergone surgery on May 2 (craniotomy and viral tracer injection). Post-operative care included the administration of medications - Loxicam, Excede and diphphenhydrmine. The vet tech on duty misread the instructions for giving Loxicam on the treatment log and did not give that medication on Saturday, May 4 resulting in the NHP not receiving analgesia on that day. As this has occurred multiple times in recent months, the committee asks for the following additional information.

- Please clarify if this technician has previously provided the incorrect care for animals, in particular regarding to analgesic regimens. If so, provide the past occurrences.
- Please provide the training records for the veterinary technician and a summary of any applicable supplemental information regarding specific training on animal care that was provided to the technician
- Please provide a sample of the treatment log that was posted for the veterinary technician to review for weekend care
- Please clarify how cages are labeled to indicate the time of initial surgery for animals under post-op care
- Please address what changes in process may be leading to an increase in the number of incidents where veterinary technicians have not provide the prescribed treatments to be given to an animal
- Please provide potential secondary checks or other methods that could be implemented to ensure these events do not occur

**MAY** The committee discussed a self-report in which "on Feb 5, 2019 the lab initiated isolation stress treatment as stated in our protocol. Animals were placed in conical restrainer and placed back into the box. The care technician came in and tampered with the boxes and the restrainers. This extra handling led to the animals getting turned around in the restrainer, limiting their gas exchange. Three animals died of accidental asphyxiation." The committee asked for the following additional information regarding the incident:

- Please provide the records of the three animals in question
- Please clarify how long these animals were in the stress test
- Please provide additional information about the stress treatment including: whether the size of the conical in relation to the size of the mouse allows for animals to turn around; the size of the air holes in the conical and whether they are located at both ends of the conical to allow for air exchange in the event that one hole is blocked
- Please provide more detail regarding the experience the lab has with this particular stress test including the number of animals that have successfully

undergone this procedure versus the number that have died of asphyxiation or had other unexpected complications

• Please address whether additional provisions should be made for observation of the animal at more regular intervals or continuously

**MAY** The committee was updated on the death of a display snake in Morris. The animal was old and there were no signs of distress.

JUN. The committee was updated on the recent UReport that alleged an investigator had staff do procedures for outside biotech companies that were not approved on protocol, housed animals in his laboratory, took animals home, and pressured lab staff into not sharing information with IACUC. The IACUC chair talked with a former lab manager about these allegations and although the lab manager would not corroborate any of the items in the UReport, the manager did have concerns with the trusting the PI. The committee considers the matter closed as there was no further information confirming allegations, but will have some unannounced visits by the compliance staff.

**JUN.** The committee continued to discuss an RAR self-report in which weekend treatment was not provided as outlined. The committee reviewed treatment logs, the training records for the veterinary technician involved, and SOPs for weekend care. RAR staff also responded to questions from the committee regarding the communication between veterinarians and veterinary technicians to prepare for weekend care and during the weekend should questions or difficulties arise. In response to the information provided, the committee had two additional requests

- Following the RAR/ESS report discussed at the February 12, 2019 IACUC meeting, vet technicians were to receive retraining on the processes for weekend care. Please provide the committee with an update regarding this retraining session including whether it has been completed or when it will be scheduled
- The committee noted that there are several refinements to the communication process for weekend care between veterinarians, veterinary technicians, and other RAR staff that are not included in the SOPs provided to the IACUC. Please update the SOPs to reflect the current practices for weekend care including communication between area veterinarians, the weekend on call veterinarian, and veterinary technicians

**JUN**. The committee discussed lab responses to a self-report in which mice died of asphyxiation while under restraint. The lab outlined changes that have been made to prevent this from happening in the future. The committee has no additional concerns and considers the matter closed.

**JUN.** The committee discussed a self-report in which six mice were treated with diphtheria toxin without being transferred to the appropriate experimental protocol. The animals did not undergo any other experimental manipulations, but five of the six animals became ill and were euthanized as requested by the vet staff on 5/9/19. The sixth animal that was not showing signs of illness was also euthanized at the same time. Moving

forward, the lab will transfer animals from the breeding protocol to the experimental protocol at the time of weaning. The committee has no additional questions and considers the matter closed.

**JUN.** The committee discussed a protocol which had recently submitted an amendment to request additional animals after animals on study were not used for the second planned surgery due to scheduling conflicts with the surgeon. There were no welfare concerns. The committee had no additional questions and considers the matter closed.

**JUN.** The committee discussed a self-report in which an MD/PHD student was inadvertently not listed as personnel on a study that they were working on. The lab has since added the person to the protocol. The committee has no additional questions and considers the matter closed.

**JUL.** The committee discussed the responses received from RAR regarding recent issues with weekend care. The committee noted that the SOP should be updated to include the opportunity for weekend staff to visit areas and details on communication with PIs about weekend care. In addition, the current records system does not consistently maintain records in the same room that animals are housed and some PIs noted that a single folder or location that listed all RAR records, procedures and care for an animal would be helpful. IACUC leadership will continue to work with RAR discussing ways to improve communication with PIs and the availability of records.

**JUL.** The committee discussed a self-report regarding an animal that was burned by an electric heating pad. Insulation had been provided and the heating pad was set to the low setting. The lab will use a recirculating water heating device to prevent a recurrence of this event. The IACUC will also contact other labs conducting survival surgery on USDA regulated animals to inform them that recirculating water heating pads should be used rather than electric heating pads.

**JUL.** The committee discussed a self-report in which an animal on study with vestibular disease received treatments, but these treatments were not documented. The lab has retrained staff about the importance of documenting all treatments. The committee considers the matter closed.

**JUL.** The committee discussed a self-report in which a lab was not shaving the surgical site prior to surgery. The lab had used scissors to remove hair in this area as the incision was less than 2 mm. Moving forward the lab will use nair or shave the site to ensure that the hair is adequately removed. The committee considers the matter closed.

JUL. The committee discussed a self-report regarding animals that were housed in a lab for over 24 hours to acclimate the rats prior to undergoing behavioral tests. The lab did not have an approved IMHA and has been informed that they will need to get an IMHA approved before extended housing can be done again. The lab will work with the IACUC office to get an IMHA approved if it decides to continue using an extended acclimation that is longer than 24 hours. The committee considers the matter closed. **JUL.** The committee discussed an UReport regarding a pig on study. The report outlined an animal that was under dosed with SR Buprenorphine due to use of the wrong formulation. The animal did have multiple analgesics onboard and the endpoint was accelerated for a more humane endpoint. The IACUC leadership group is currently investigating the issue and will provide the committee with more information at the next IACUC meeting for further discussion.

**AUG.** The committee discussed a PI response to a request for an action plan to address ongoing non-compliance. The committee endorsed the plan outlined by the PI and had no additional comments.

**AUG.** The committee discussed a self-report in which a leg injury was identified in a client-owned duck that was on a UMN IACUC Protocol. Lab staff consulted with the area veterinarian and the animal was treated. The PI will update the protocol to outline a course of action for instances where non mortal injuries are identified in animals while on study.

**SEPT.** The committee discussed a UReport regarding a pig on study. The report outlined an animal that was under dosed with SR Buprenorphine due to use of the wrong formulation. The animal did have multiple analgesics onboard and the endpoint was accelerated for a more humane endpoint. The area veterinarian and other veterinary staff were consulted on the issue, and indicated that there was no change or increase in pain status due to the incorrect formulation. While notification of the PI did not occur, RAR has stated that moving forward, the PI will be notified in the event that issues like this occur. The committee considers the matter closed.

**SEPT.** The committee was notified of an anonymous report regarding primarily HR issues. The leadership group will collect additional information to bring back to the committee to determine if there are animal welfare concerns in addition to the HR component.

**SEPT.** The committee reviewed the updated RAR SOP outlining weekend care. The updates were made in response to a June RAR self-report. The committee had no additional comments or suggestions regarding the SOP. The committee considers the matter closed now.

**SEPT.** The committee discussed a self-report in which a naïve pig was prepared for a study on a separate protocol. As the pig had not undergone any previous procedures, the animal was transferred to the corresponding protocol. The committee had no additional comments and considers the matter closed.

**SEPT.** The committee discussed a self-report in which animals were not transferred appropriately between protocols. The animals are approved to be used for two labs occurring over the same afternoon, however, these procedures are outlined on two

separate protocols making it difficult to complete the transfer as the labs are concurrent. Procedures have been consolidated to one protocol to avoid this administrative oversight in the future. The committee had no additional comments and considers the matter closed.

**SEPT.** The committee discussed a report outlining a concern about an animal that was euthanized early due to complications with an implant. The animal had undergone emergency surgery to repair an implant that had come out. There was concern that the animal was too deeply anesthetized, but review of the surgical records and interviews with those involved with the surgery did not corroborate this concern. The committee considers the matter closed.

# • SUBCOMMITTEE UPDATES

#### Endpoints Subcommittee: No updates

**Subcommittee on Cephalopod Use:** The committee has reached out to the PI planning to use cephalopods. The PI was on sabbatical over the summer working with other investigators Marine Biological Laboratory to learn more about housing for cephalopods. The subcommittee will meet with the PI this semester now that PI has returned.

# 6. eProtocol Form Updates

# **IACUC Form Changes**

# September 2019 – Current PROD IACUC-R1S1 Form edited 2105 Effective Date 7/27/2018

Sections with Changes

- Page 17. Rationale Add New Question
- Page 18. Species Add Text
- Page 19. Procedures Update Link
- Page 20. Procedure Details Update Link
- Page 21. Procedure Antibody Production Remove Link
- Page 22. Procedure Behavior Update Text and Link
- Page 23. Procedure Breeding Remove Question, New Text, New Link
- Page 24. Procedure Dietary or Fluid Modification New Link
- Page 25. Procedure Physical Restraint New Link
- Page 26. DEHS Remove Checkboxes/Text, Update Text
- Page 27. Chemicals Update Text, Update Link
- Page 28. Radiations Update Link, Update Text
- Page 29. Continuing Review New Question
- Page 30. Amendment Update Text

# **Changes for the Rationale Page**

• Include under "study Objectives:

e. For 3 year renewals, please provide a brief summary of the research results from animals obtained during the prior approval period.

IACUC - IACUC-R1S1 Protocol Title: test	Protocol ID: 1908-37322A (Vegoe, Eric)		IACUC - IACUC-R1S1 Protocol Title: test	Protocol ID: 1908-373224 (Vegoe, Eric)
Trotoon Habries	Save   Spell Check   Help   Close		Trease and the second	Save   Spell Check   Help   Close
UNIVERSITY OF MINNESOTA Driven to Discover	Previous Next		UNIVERSITY OF MINNESOTA Driven to Discover Personal Information	Previous Next
Funding	test		Funding	test
Rationale	212		Rationale	
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Print View Event History	c. How will the results of this study be used? (1-2 sentences):		Print View Event History	c. How will the results of this study be used? (1-2 sentences):
	d. Summarize the specific aims (derived from the grant proposal):			d. Summarize the specific aims (derived from the grant proposal):
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# **Changes for the Species Page**

• Pain class information is no longer listed below "1. Pain Class Categorization (Check all that apply then add the number of animals by acquisition method. List all animals produced in conjunction with the study even those that are the wrong genotype and are euthanized.)" Please list the following below the species table:

"Class A: No pain, distress or use of pain-relieving drugs.

Class B: Potential pain/distress WITH appropriate analgesia/anesthesia/tranquilizers.

Class C: Pain/distress WITHOUT analgesia/anesthesia/tranquilizers."

Solid       Solid Case Likely       Control       Solid Case Likely       Case Likely       Case Likely       Cas			A (Vegoe, Eric)	locol ID: 1908-37322	Pro		ACUC - ACUC-R1S1				2A (Végbe, Eric)	ocol ID: 1998-3732	Proto		UC - IACUC-R1S1
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# **Changes for the Procedures Page**

Change link from <u>University of Minnesota Policy on</u>	To Information on Animal Transport
Animal Transport	

# https://drive.google.com/file/d/1w81ColPsLv\_f9oxcYq7RTchfmxhVA2bv/view

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# Changes for the "Anesthetic Regimen" and "Analgesic Regimen" fields listed on Procedure Pages

Change <u>recommended anesthetics</u> link to	recommended anesthetics			
https://research.umn.edu/units/iacuc/policies-gu	idelines/animal-use-guidelines-exceptions			

👲 eProtocol - University o	eProtocol - University of Minnesota Mozilla Firefax			-Xi -
🗇 🔒 https://eprotoco	ol.umn.ecu/applicationform/FORM_PROCEDURES.do	E	🖾 🕁	=
	Anesthetic Regimen	Save	Cancel	
	Will you be performing this procedure with anesthesia?	Yes	No	New Link:
	If yes, please add at least one Anesthetic Sedaive, Analgesic and/or Paralytic Agert. Click on "Add" to add Anesthetic Agent(s). For yet recommended anestetics and doses see link, <u>recommended anesthetics</u>	-		https://research.umn.edu/units/iacuc/policies-guidelines/animal-use-guidelines-exception
	Please click on "Add" to add Anesthetic Agent(s) and Sedatives(	5)		
	Click on "Add" to add Analgesic Agent(s). For vet recommended doses see link, <u>recommended analgesics</u> Analgesic agents (Prophylactic and Intra- procedural)		_	4
	Please click on "Add" to add Analgesic Agent(s). Click on "Add" to add Paralytic Agent(s).			

# Changes for the Antibody Production Procedures Page

Remove Guidelines for Immunization of Research Animals

Anti	body Production	Save Cancel	
Dies Gui	Se select Immunization and Do delines for Immunization of Research MUNIZATION AND POLYCLONAL	ANTIBODY PRODUCTION	Remove Text
1.	List antigens:		
2.	Adjuvant details:		
	Initial immunization:	Select One	

# Changes for the Behavior Procedures Page

Change these two links to one link IACUC Policy on Anesthesia, Surgery and Post- Procedural Records IACUC Sample Recordkeeping Forms	Sample Anesthesia, Surgery, and Post-Procedural Recordkeeping Forms
https://research.umn.edu/units/iacuc/policies-guidel exceptions#SampleAnesthesia	ines/animal-use-guidelines-



# Changes for the Breeding Procedures Page

Remove the question: "Anticipated number of offspring that will be used for experiments:"

Methods of Animal Identification broken	Remove link
<u>biopsy guidelines?</u> broken	biopsy guidelines?
Remove Question: "What are the criteria for euthanasia of the Breeders?"	<ul> <li>Include:</li> <li>Recommended criteria for euthanasia of mouse or rat breeders.</li> <li>They produce no litter within 60 days of mating (longer may be acceptable if delayed breeding is a strain characteristic).</li> <li>They produce no litter within 60 days of their last litter and are not visibly pregnant.</li> <li>They produce litters but do not wean pups for two to three litters.</li> </ul>
Remove Question: "Anticipated number of offspring that will be used for experiments:"	

Ereoding.	Save Cancel		Brocdiag	Save Ca	(pan)
You are required to keep accurate records of the number of animals produced and This information may be requested annually.	the r altimate dispositor.		You are required to keep accurate records of the number of animals produced and their up This information may be requested animally. Please explain the need for breeding on this protocol	ltimate dispo	isition.
Please explain the need for pressing on this protoce					
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Anticipated Number of Animals	hbΔ		Please cfck os "Add" to add Anticipated Number of Asimalia	NUU	-
Please click on "Add" to add Anticipated Number of Anima	15.				
Provide rationale/accentific justification for the number of offspring produced	-	Hatrove Sugation	Provide rationale/scientific justification for the number of offspring produced		
Anticipated number of offspring that will be used for experiments:			What will be done with the surplus animals?		
		New Last Nameva Last Has			
What will be done with the samplus animals?	~	Recommended distortel for optimunatio or more or real averages     They preduce no Hor willing 60 days of nationaling (longer may be acceptable if delayed benefing to a real-characteristic).	Recommended criteria for euthanasia of mouse or rat breeders. • They produce no litter within 60 days of mading (tonger may be acceptal breeding is a strain charactoristic), • They produce no litter within 60 days of their last litter and are not visibl	ble If delays	ed L
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NOTE If crossing pre-existing genetically modified strains, please contact the IBC to determine if BC approval is required.			Are there any interent problems for the animal associated with the genetically modified phenotype?	*25	No
Are there any inherent problems for the mirmal associated with the cesofically modified phenotype? If "Yes", dozcrito:	Tes 140		ll "Yes", describe:		
			Describe any special care or monitorine that may be required for these animals:		
Describe any special care or monitoring that may be required for these animals			Bethed of manifesting encourses of transcenary in the unionality		
Nethod of monitoring gressence of transaene in the animals:	-	Renove Link	in the ormanic my presence or managene in the dominate.		
			Methods of Animal Identification		
Mathods of Animal Identification		New Link	f a tail bionsy is performed, do you need an exception to the bioraxy guidelines?	Vac Bo	
The biopsy should not exceed Smm in size unless approval is obtained from the IACU	1	https://docs.aponia.com/document/d/182/Or/WCrM sEvation/TRF_siCKP/samil/Pages_0/acit	If yes, then provide details including the scientific justification:	100 110	1 L .
If a tail biopsy is performed, do you need an exception to the biopsy guidelines?	Yes Nov	entering of the second			

# Changes for the Dietary or Fluid Modifications Procedures Page

Under FOOD: "If Lab Staff/Other, do you need an exception to the <u>records keeping policy</u> ?" has a broken link. Update with link to the right	"If Lab Staff/Other, do you need an exception to the <u>records keeping policy</u> ?"		
https://docs.google.com/document/d/165zyyuZVEG1	TvWZyVHkm57ZJVcYImBxBdQf0npst4Mg/edit		
"If Lab Staff/Other, do you need an exception to the records keeping policy ?" has a broken link. Update	records keeping policy ?"		

Dietary or Fluid Modifications	Save Cancel	
Please select diet modification type (check all that apply):	🗌 Food 📋 Fluid	
FOOD		
ndicate whether diatapy manipulations area	Modified	
Describe Dietary Manipulations:	(Not restricted)	
Describe the expected impact of this manipulation on the animal's health and the p listress. Explain what will be done to alleviate or minimize these potential adverse	ootential for discomfort or effects:	
lescribe frequency of animal health monitoring during periods of food restrictions.		
OTE: The Guide for the Care and Use of Laboratory Animals states that "Body weig east weekly and more often for animals requiring greater restrictions. Written recore or each animal to document daily food and fluid consumption, hydration status, and hanges used as criteria for temporary or permanent removal of an animal from a r	ghts should be recorded at ords should be maintained d any behavioral and clinical protocol."	
Jo you need an exception to the weekly weighing policy?		
If yes, then provide details including the rationale:		
Describe length of time animals will be on experimental diet		
Will neonates be fasted beyond 8 hours, ruminants beyond 48 hours, or other mammals beyond 24 hours?		New Link: https://docs.google.com/document/d/165zyyuZVEG1TvWZyVHkm57ZJVcYImBxBdQl0npst4Mg/edit
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# **Changes for the Physical Restraint Procedures Page**

IACUC Guideline on Physical Restraint.

https://docs.google.com/document/d/1EO5YygfrGCnIBo1oR\_GhSgCKLKxeJ0dxzAgZJ5pis-g/edit

Physical Restraint	Save Cancel	
Complete if conscious animals will be physically restrained for r pain associated with restraint, the period of restraint should be the project. <u>IACUC Guideline on Physical Restraint</u> .	nore than one hour. To minimize distress or no longer than required to achieve the aims of	
Describe the method of restraint:		New Link:
Indicate the duration of restraint:		https://docs.google.com/document/d/1EO5YygfrGCnlBo1oR_GhSgCKLKxeJ0dxzAgZJ5pis-g/edit
Describe how the animal(s) will be acclimated to the restraint m	ethod:	

# **Changes for the DEHS Page**

# Checklist:

Remove the "Annual Safety Training" and "Hazardous Waste" checkboxes.

Change the text from "Radiations" to "Radioactive Material and/or Radiation Producing Equipment"



# Chemicals Section: Changes are highlighted Chemicals

# [Text Mark-Up]

In the table below list any toxic or poisonous chemicals, carcinogens, chemotherapeutic agents and other hazardous drugs, reproductive hazards or nanomaterials used in this protocol. Also list compounds where the hazards have not been fully studied. Contact DEHS, 612-626-6002, dehs@umn.edu labsafe@umn.edu, if you have questions regarding what to include.

You must attach a <del>Standard Operating Protocol (SOP) and an Animal Husbandry form for each chemical listed in this section. <mark>Chemicals with the same hazard class that are being used in similar ways can be listed on the same SOP and Animal Husbandry form.</mark> Toxic Hazard Class Standard Operating Procedure (SOP) that covers all the chemicals listed in this section.</mark> SOPs on file with DEHS may be used as long as they have been updated within the last 3 years.</del>

A collection of resources, including step-by-step instructions, form templates, sample SOPs, and lists of chemicals that require SOPs can be found at: <a href="http://www.z.umn.edu/iacucresources">www.z.umn.edu/iacucresources</a>

# [New Text]

In the table below list any toxic or poisonous chemicals, carcinogens, chemotherapeutic agents and other hazardous drugs, reproductive hazards or nanomaterials used in this protocol. Also list compounds where the hazards have not been fully studied. Contact DEHS, 612-626-6002, labsafe@umn.edu, if you have questions regarding what to include.

You must attach a Toxic Hazard Class Standard Operating Procedure (SOP) that covers all the chemicals listed in this section. SOPs on file with DEHS may be used as long as they have been updated within the last 3 years.

A collection of resources, including step-by-step instructions, form templates, sample SOPs, and lists of chemicals that require SOPs can be found at: www.z.umn.edu/iacucresources

# [Text Mark-Up]

Sample SOP: Toxic Hazard Class SOP Template

Tamoxifen, STZ, BrdU, BrdU in Drinking water, Progesterone

It is the responsibility of the PI to ensure that RAR staff (or other animal care staff if animals are not housed in RAR) are notified two weeks in advance of administering a hazardous agent to an animal.

Chemicals	therspectic scents and other		Chemicals	
<ul> <li>In the table below list any toxic or personous chemicals, carchogens, chemonazardous drugs, reproductive hazards or nanomaterials used in this protocol dehs@umn.edu, if you have questions regarding what to include.</li> <li>You must attach a Standard Operating Protocol (SOP) and an Animal Husband in this section. Chemicals with the same hazard class that are being used in same SOP and Animal Husbandry form. SOPs on file with DEHS may be used updated within the last 3 years.</li> <li>A collection of resources, including step-by-step instructions, form templates chemicals that require SOPs can be found at: www.z.umn.edu/iacucresource</li> </ul>	dry form for each chemical listed similar ways can be listed on the as long as they have been	Text Edit	In the table below list any toxic or poisonous chemicals, contact dous drugs, reproductive hazards or nanomaterials hazards have not been fully studied. Contact DEHS, 612-62 regarding what to include. You must attach a Toxic Hazard Class Standard Operating in this section. SOPs on file with DEHS may be used as lone. A collection of resources, including step-by-step instruction chemicals that require SOPs can be found at: www.z.umn	carcinogens, chemotherapeutic agents and other sused in this protocol. Also list compounds where the 26-6002, labsafe@umn.edu; if you have questions g Procedure (SOP) that covers all the chemicals listed ng as they have been updated within the last 3 years. ions, form templates, sample SOPs, and lists of n.edu/iacucresources
Chemicals	Add		Chemicals	bbA
Please click on Add to add Chemicals			Please click on Add to	to add Chemicals
Sample SOP's : <u>Tamoxifen</u> , <u>STZ</u> , <u>BrdU</u> , <u>BrdU in Drinking water</u> , <u>Progesterone</u> It is the responsibility of the PI to ensure that RAR staff (or other animal care staff if animals are not housed in RAR) are notified two weeks in advance of administering a hazardous agent to an animal.			Sample SOP: Toxic Hazard Class SOP Template	
		Text Edit New Link:	It is the responsibility of the PI to ensure that RAR staff (or RAR) are notified two weeks in advance of administering a	r other animal care staff if animals are not housed in a bazardous agent to an animal.



# Radiation Section: Changes are highlighted Radiations

Contact Radiation Protection (612-626-6764) for radiation protection forms or assistance, or visit the web at http://www.dehs.umn.edu/rad.htm https://radsafety.umn.edu/.

Where will the radiation be used (building and room no.)? Is this an RAR facility? Name of approved radioisotope permit holder or <mark>source owner machine owner</mark>: Duration of permit <mark>(if applicable)</mark>:

# **Controlled Substances Section: Changes are highlighted** Are you using Controlled Substances in this protocol? Yes No

Use of Controlled Substances requires DEHS Approval. <del>Please complete the controlled substances module for review and approval. Please submit a separate controlled substances protocol in eProtocol if you haven't already done so or if your current one needs updating (over three years old, safe moved, registrant changed, added controlled substances).</del>

Radiations ContactRadiation Protection (612-626-6764) for radiation protection forms or assistance, or visit the web at http://www.dehs.uom.edu/rad.htm	NewLink	Racitations Contact Rediation Protection (612-826-8764) for radiation protection forms or sesistance, or visit the web at https://radsalety.umm.edu
Where will the radiation be used (building and room no.)?		Where will the radiation be used (building and room no.)?
Is this an RAR facility?		Is this an RAR facility?
Name of approved radiosotope permit holder or source owners	Text Change	Name of approved radioisctose permit holder or machine owner:
	Text Change	
		Duration of permit (if applicable) :
Radiations Add		Radiations
Please click on Add to add Radiations		Please click on Add to add Radiations

# **Changes for the Continuing Review Page**

Update the continuing review page to include a question addressing breeding of animals beyond allotted amount

"Have you exceeded the allotted number of animals to be bred or used?

If so, provide an updated estimate to the number of animals needed here and update the species table and breeding procedures accordingly."



# **Changes for the Amendment Page**

Update to include text reminding investigators to update the Species section to account for additional animals as needed; the Health and Monitoring Section to account for additional adverse health concerns associated with new procedures or aims; and the experimental endpoints section to account for new endpoints.

To include after bullet points: 2. For changes in Protocol, complete the change request by clicking on that section in the left IMPORTANT: Update the following if new aims or navigation bar and completing the change in the procedures are included in the amendment: Protocol Form. Adding, removing, or Updating Personnel -"Species Section" to reflect new animals **Personnel Section** needed for additional aims or procedures Species - Species Section "Experimental Design" to reflect new aims or Animal Housing Location - Species Section procedures, how these are incorporated into Funding Source - Funding the previously approved design, and the Protocol rationale - Protocol Information impact on the sequential description of what Section animals will undergo Adding or modifying procedures - Protocol "Health and Monitoring Section" to reflect Information Section additional adverse health conditions related Other changes - see relevant section at left to new aims or procedures "Experimental Endpoints" to reflect new endpoints for additional aims



# 8. Pain classification

The committee discussed changing the pain classification to align with USDA categorization and voted to revise our current categorization. Previously the UMN IACUC implemented an A-C categorization rather than the USDA B-E categorization.

USDA Pain Categories	UMN Pain Categories
Class B- Animals being bred, conditioned,	Class A: No pain, distress or use of pain-
or held for use in teaching, testing,	relieving drugs.
experiments, research, or surgery but not	
yet used for such purposes	
Class C- Animals upon which teaching,	Class B: Potential pain/distress WITH
research, experiments, or tests were	appropriate analgesia/anesthesia/tranquilizers.
conducted involving no pain, distress, or	
use of pain-relieving drugs.	
Class D- Animals upon which	Class C: Pain/distress WITHOUT
experiments, teaching, research, surgery,	analgesia/anesthesia/tranquilizers.
or tests were conducted involving	
accompanying pain or distress to the	
animals and for which appropriate	
anesthetic, analgesic, or tranquilizing	
drugs were used.	
Class E- Animals upon which teaching,	
experiments, research, surgery, or tests	
were conducted involving accompanying	
pain or distress to the animals and for	
which the use of appropriate anesthetic,	
analgesic, or tranquilizing drugs would	
have adversely affected the procedures,	
results, or interpretation of the teaching,	
research, experiments, surgery, or tests.	

This change will require outreach to researchers in addition to updating eProtocol forms, so may not be completed in the next six month cycle.

The committee tabled discussion to standardize pain categories for the following common procedures that are often disputed regarding pain class.

- Infection models
- Food and water restriction
- Physical restraint
- Tumor studies
- Electro-fishing

# 9. DEHS SOPs and IACUC requirements for DEHS content

DEHS reminded the IACUC that their group is responsible for the SOPs of hazardous chemicals and works to maintain these documents. It is not necessary for PIs to include an attached SOP on all protocols as DEHS works directly with investigators on the content and maintenance of these documents. In the event that there is a chemical spill or questions about handling, DEHS should be contacted directly. DEHS would prefer that IACUC not make requests to the PI regarding these SOPs.

In the event that chemicals listed on a protocol are not included on the DEHS page, IACUC reviewers should provide a stipulation and the office will coordinate additional DEHS review.

# 10. Health and Monitoring section in the protocol: does frequency of monitoring include RAR and lab monitoring frequency combined?

RAR announced that they have refined their daily health check process for mice and rats with the goal of minimizing disruption of animals and potential impacts to your research. The daily health checks will now be comprised of a visual scan of each cage in a room. Cages will not be handled or opened unless concerns are identified during this visual scan. Animal health issues identified during health checks will still be managed according to our standard policies and procedures.

Propose updating the text in the Health and Monitoring section of the protocol to differentiate monitoring provided by RAR as a daily check and study based monitoring performed by the lab.

# 11. Pre-worded answers available to PIs for common procedures or treatments

Tabled until a future meeting

# 12. Veterinary pre-consult

Tabled until a future meeting

# 13. RAR training offered

Shannon May provided the committee with a presentation on RAR training courses provided and the registration process. RAR currently provides both required training and elective courses for researchers seeking hands-on experience or additional information and practice with common procedures. Required training is offered on a regular basis in multiple areas to facilitate timely completion of required modules.

# Required Training

- RAR Orientation (required for all RAR facility users)
- Facility Tour (required for all RAR facility users)

- Microisolator Training (required for access to RAR SPF facilities)
- NHP Training (required for access to RAR NHP facilities or to work with NHP tissues)

# Elective Training

- Mouse Basics
- Carbon Dioxide Euthanasia
- Mouse Injections (subcutaneous and intraperitoneal)
- Mouse Tail Vein Injections
- Mouse Blood Collection (saphenous and cardiac)
- Mouse Facial Vein Blood Collection
- Mouse Gavage
- Rat Handling and Injections
- Basics of Suturing
- Aseptic Technique in Rodents
- Anesthesia of Mice and Rats
- Rabbit Handling and Injections

In addition to these courses, specialized courses are available upon request.

See slides from the presentation located in the appendix for additional information.

# 14. Evaluation of the OLAW Checklist

The meeting attendees formed small groups to discuss different sections of the OLAW checklist which were then reviewed by the reconvened group. The committee determined that the Animal Use Program received a grade of "A" for all areas of the OLAW checklist except one deficiency that is currently being addressed. The committee also discussed potential ways that certain other areas could be improved to make the program stronger (see below).

# **Deficiency:**

- Institutional Policies and Responsibilities, Animal Care and Use Program *Resources necessary to manage program of veterinary care are provided.* 
  - It was noted that RAR does not currently have access to imaging equipment and an i-STAT blood analyzer. OVPR has identified funding and RAR has ordered an i-STAT. RAR is also looking into access for imaging equipment that could potentially be shared with investigators.

# **Opportunities for Improvement:**

- Institutional Policies and Responsibilities, Disaster Planning and Emergency Preparedness – Law enforcement and emergency personnel are provided a copy and integration with overall plan is in place.
  - The committee discussed whether outstate facilities had communicated the disaster plans to local security and law enforcement recently. Previously this was managed by the Attending Veterinarian through the institution's general emergency plans.
- Institutional Policies and Responsibilities, IACUC Protocol Review Special Considerations *Toe clipping only performed when no alternative performed aseptically and with pain relief.* 
  - Current posted guideline on the IACUC page specifies an age limit but does not list pain relief. Dr. Felicia Boynton has volunteered to look into this issue.
- Veterinary Care, Euthanasia Training is provided for appropriate methods for each species and considers psychological stress to personnel
  - We will look into providing more training opportunities for staff that are experiencing psychological stress. Potentially a program is already offered through the veterinary school and the IACUC could share resources for support materials. This information could also be shared during inspections.
#### 14. Sub-Groups for Evaluation of the OLAW Checklist

#### Group 1:

Institutional Policies and Responsibilities, Sections 1-4 ("Animal Care and Use Program", "Disaster Planning and Emergency Preparedness", "IACUC", and "IACUC Protocol Review - Special Considerations") (on pages 1-2)

Felicia Boynton Paul Lindstrom Megan McCoy Mimie Pollard Sam Baidoo Jennifer Hubbard Keith Barker Georgiy Aslanidi Jessica Sieber

#### Group 2:

Institutional Policies and Responsibilities Sections 5-8 ("IACUC Membership and Functions", "IACUC Training", "IACUC Records and Reporting Requirements", and "Veterinary Care") (on pages 2-3)

Ben Clark Ilana Cohen Scott Madill Carolyn Fairbanks Don Martin Geoff Ghose Laura Hocum-Stone

Jan Zimmerman

Group 3:

Institutional Policies and Responsibilities Sections 9-12 (, "Personnel Qualifications and Training", "Occupational Health and Safety of Personnel", "Personnel Security", and "Investigating & Reporting Animal Welfare Concerns") (on pages 3-5)

Brian Crooker Peggy Norris George Wilcox Sally Noll Henry Wong Dan Montonye Tricia Van Ee Molbert Cynthia Lee

#### Group 4:

Veterinary Care sections 1-3 ("Clinical Care and Management", "Animal Procurement and Transportation/Preventive Medicine", and "Surgery") (on pages 6-7)

Nathan Koewler Jennifer Borgert Jim Perry Dezhi Liao Kakambi Nagaraja Tim Goldsmith

Lynn Impelluso Christine Sivula

#### Group 5:

Veterinary Care sections 4-6, ("Pain, Distress, Anesthesia and Analgesia", "Euthanasia", and "Drug Storage and Control") (on page 7)

Melanie Graham Kristin Pilon Liz Pluhar Robert Schumacher Marilyn Bennett

Maggie Luesse Whitney McGee

# I. Semiannual Program Review Checklist

### **Institutional Policies and Responsibilities**

#### Date:

1.	An	imal Care and Use Program 📲	$\mathbf{A}^*$	м	S	С	NA
		Responsibility for animal well-being is assumed by all members of the program (Guide,					
		IO has authority to allocate needed resources (Guide, p.13)	Δ				
		Resources necessary to manage program of veterinary care are provided ( <i>Guide</i> , <u>p14</u> )			s		
	•	Sufficient resources are available to manage the program, including training of personnel in accord with regulations and the <i>Guide</i> ( <i>Guide</i> , pp 11, 15)	A				
		Program needs are regularly communicated to IO by AV and/or IACUC ( <i>Guide</i> , p13)	A				
	•	Responsibilities for daily animal care and facility management are assigned to specific individual(s) when a full-time veterinarian is not available on site ( <i>Guide</i> , p 14) [must]	A				
	•	Inter-institutional collaborations are described in formal written agreements ( <i>Guide</i> , $\underline{p}$ <u>15</u> )	A				
	•	Written agreements address responsibilities, animal ownership, and IACUC oversight ( <i>Guide</i> , <u>p 15</u> )	A				
2.	Di	saster Planning and Emergency Preparedness	<b>A</b> *	м	S	С	NA
	•	Disaster plans for each facility to include satellite locations are in place ( <i>Guide</i> , <u>p 35</u> , <u>p</u> 75) [must]	A				
		Plans include provisions for euthanasia ( <i>Guide</i> , p 35) [must]	A				
		Plans include triage plans to meet institutional and investigators' needs (Guide, p35)	A				
		Plans define actions to prevent animal injury or death due to HVAC or other failures ( <i>Guide</i> , p 35)	A		1		
		Plans describe preservation of critical or irreplaceable animals (Guide, p 35)	A				
		Plans include essential personnel and their training (Guide, p 35)	A				
	•	Animal facility plans are approved by the institution and incorporated into overall response plan ( <i>Guide</i> , <u>p 35</u> )	A				
-		Law enforcement and emergency personnel are provided a copy and integration with overall plan is in place ( <i>Guide</i> , <u>p 35</u> )	A				
3.	IA	CUC	Α*	м	S	C	NA
		Meets as necessary to fulfill responsibilities ( <i>Guide</i> , p.25) [must]	A				1
	•	IACUC Members named in protocols or with conflicts recuse themselves from protocol decisions ( <i>Guide</i> , p 26) [must]	A				
-	1 â II	Continuing IACUC oversight after initial protocol approval is in place ( <i>Guide</i> , p33)	A	-	1		
		IACUC evaluates the effectiveness of training programs (Guide, p15)	A	-		-	
л	τA	CLIC Protocol Review - Special Considerations	۸*	м	c	c	NA
	•	Humane endpoints are established for studies that involve tumor models, infectious diseases, vaccine challenge, pain modeling, trauma, production of monoclonal antibodies, assessment of toxicologic effects, organ or system failure, and models of cardiovascular shock ( <i>Guide</i> , p 27)	A				
		For pilot studies, a system to communicate with the IACUC is in place ( <i>Guide</i> , p 28)	A				
1	٠	For genetically modified animals, enhanced monitoring and reporting is in place ( <i>Guide</i> , p 28)	A		11		
	•	Restraint devices are justified in the animal use protocols (Guide, p 29) [must]	A				
	•	Alternatives to physical restraint are considered (Guide, p29)	Α	-			
		Period of restraint is the minimum to meet scientific objectives (Guide, p 29)	A				
		Training of animals to adapt to restraint is provided (Guide, p29)	Α				
		Animals that fail to adapt are removed from study (Guide, p29)	Α				
	•	Appropriate observation intervals of restrained animals are provided ( <i>Guide</i> , <u>p29</u> ) Veterinary care is provided if lesions or illness result from restraint ( <i>Guide</i> , <u>p30</u> )	Α				
		[must]	Α				

	•	Explanations of purpose and duration of restraint are provided to study personnel $(Guide, n, 30)$	Δ				
	•	Multiple surgical procedures on a single animal are justified and outcomes evaluated					
		(Guide, <u>p 30</u> )	Α				
	•	Major versus minor surgical procedures are evaluated on a case-by-case basis ( <i>Guide</i> , p. 20)					
-		<u>D-30</u> Multiple survival procedure justifications in non-regulated species conform to regulated	A				-
		species standards ( <i>Guide</i> , $\underline{p}$ <u>30</u> )	A				
	•	Animals on food/fluid restriction are monitored to ensure nutritional needs are met ( <i>Guide</i> , <u>p 31</u> )	A			11	
	•	Body weights for food/fluid restricted animals are recorded at least weekly ( <i>Guide</i> , $\underline{p}$ 31)	A				
		Daily written records are maintained for food/fluid restricted animals (Guide, p31)	A				
	•	Pharmaceutical grade chemicals are used , when available, for animal-related procedures ( <i>Guide</i> , <u>p 31</u> )	A				
	•	Non-pharmaceutical grade chemicals are described, justified, and approved by IACUC ( <i>Guide</i> , <u>p 31</u> ) Investigators conducting field studies know zoopotic diseases, safety issues, laws and	A				
		regulations applicable in study area ( $Guide, p32$ )	A				
_	•	Disposition plans are considered for species removed from the wild ( <i>Guide</i> , <u>p 32</u> )	A		-		-
	•	( <i>Guide</i> , $p$ 75)	A				1
5.	IA	CUC Membership and Functions	$\mathbf{A}^*$	м	S	С	NA
		IACUC is comprised of at least 5 members, appointed by CEO (PHS Policy, IV.A.3.)	Α				
	•	Members include a veterinarian, a scientist, a nonscientist, and a nonaffiliated non-lab					
-		IACUC authority and resources for oversight and evaluation of institution's program	A				-
	-	are provided ( <i>Guide</i> , p 14)	A				
	•	IACUC conducts semiannual evaluations of institutional animal care and use program (PHS Policy, IV.B.)	A				
		Conducts semiannual inspections of institutional animal facilities (PHS Policy, IV.B.)	A				1
	•	IACUC organizationally reports to the Institutional Official (PHS Policy, <u>IV.A.1.b.</u> )	Α				
	•	Methods for reporting and investigating animal welfare concerns are in place ( <i>Guide</i> , <u>p</u> 23) [must]	A				
		Reviews and investigates concerns about animal care and use at institution <sup>iii</sup> (PHS Policy, <u>IV.B.</u> )	A				
	*	Procedures are in place for review, approval, and suspension of animal activities <sup>iv</sup> (PHS Policy, <u>IV.B.</u> )	A				
	÷	Procedures are in place for review and approval of significant changes to approved activities (PHS Policy, <u>IV.B.</u> )	A				
	•	Policies are in place for special procedures (e.g., genetically modified animals, restraint, multiple survival surgery, food and fluid regulation, field investigations, agricultural animals) ( <i>Guide</i> , p 27-32)	A				
	•	Requests for exemptions from major survival surgical procedure restrictions are made to USDA/APHIS <sup><math>v</math></sup> ( <i>Guide</i> , <u>p</u> 30) [must]	A				
6.	IAC	CUC Training NEW	$\mathbf{A}^*$	м	s	C	NA
		All IACUC members should receive:					0.47
	-	<ul> <li>Formal orientation to institution's program (Guide, p17)</li> </ul>	-				
		<ul> <li>Training on legislation, regulations, guidelines, and policies (Guide, p17)</li> </ul>	A				
		<ul> <li>Training on how to inspect facilities and labs where animal use or housing occurs (<i>Guide</i>, <u>p 17</u>)</li> </ul>	A				
	_	• Training on how to review protocols as well as evaluate the program (Guide, p17)	A	-			
		<ul> <li>Ongoing training/education (Guide, p 17)</li> </ul>	A				
7.	IA	CUC Records and Reporting Requirements <sup>vi</sup>	$\mathbf{A}^{*}$	м	s	С	NA
		Semiannual report to the IO (PHS Policy, <u>IV.B.</u> )	1.	-		-	
		<ul> <li>Submitted to IO every 6 months</li> </ul>	A				12.5

0	Compiles program review a	and facility	inspection(s)	results (	includes all program
0	complies program review a	and racincy	mopection(5)	1 Courto (	moludes un program

A

		and facility deficiencies)	-				
		<ul> <li>Includes minority IACUC views</li> </ul>	A				
		<ul> <li>Describes IACUC-approved departures from the <i>Guide</i> or PHS Policy and the</li> </ul>					
		reasons for each departure <sup>vii</sup>	Δ				
-		Distinguishes significant from minor deficiencies	Δ				
_		<ul> <li>Includes a plan and schedule for correction for each deficiency identified<sup>viii</sup></li> </ul>	Δ				
-		Penorts to OLAW (PHS Policy IV F.)	in .				1
_	-	Annual report to OLAW documents program changes, dates of the semiannual	T	1	T	1	T
		annual report to OLAW documents program changes, dates of the semiannual program reviews and facility inspections and includes any minority views.	•				
		Dromptly advices OLAW of carious (angoing Cuide deviations or DHC Policy	A				
		o Prompting duvises OLAW of senous/ongoing Gaide deviations of PHS Policy					
-	_	Institute must premptly advice OLAW of any supportion of an animal activity by	A				
		• Institute must promptly duvise OLAW of any suspension of an animal activity by			-		-
	-	Departs to U.C. Department of Assimulture (UCDA) or Federal funding energy <sup>1</sup>	A		-		
-		Reports to U.S. Department of Agriculture (USDA) of Federal funding agency	-	-	-		1
		<ul> <li>Annual report to USDA contains required information including all</li> </ul>				1 1	
		exceptions/exemptions	A		-		-
		<ul> <li>Reporting mechanism to USDA is in place for IACUC-approved exceptions to the</li> </ul>				1	
		regulations and standards	A		_		-
		<ul> <li>Reports are filed within 15 days for failures to adhere to timetable for correction of</li> </ul>					
		significant deficiencies	Α				
		• Promptly reports suspensions of activities by the IACUC to USDA and any Federal	A				
		funding agency 🔒 🔜	1			12.1	
		Records (PHS Policy, <u>IV.E.</u> )					
		• IACUC meeting minutes and semiannual reports to the IO are maintained for 3		-	-	1	1
		years we have a second s	Α			$\sim 4$	
		• Records of IACUC reviews of animal activities include all required information <sup>x</sup>	Α				
		<ul> <li>Records of IACUC reviews are maintained for 3 years after the completion of the</li> </ul>					
		study	A				
	•	is in place including backup veterinary care <sup>xi</sup>	A				
		Veterinary access to all animals is provided ( <i>Guide</i> , <u>p 14</u> ) [must]	A		-		-
	•	Direct or delegated authority is given to the veterinarian to oversee all aspects of					
		animal care and use ( <i>Guide</i> , <u>p 14</u> ) [must]	A				-
	•	Veterinarian provides consultation when pain and distress exceeds anticipated level in	1				
	_	protocol ( <i>Guide</i> , <u>p.5</u> ) [must]	A				
	٠	Veterinarian provides consultation when interventional control is not possible (Guide, p					
_	_	<u>5) [must]</u>	Α			_	
		If part time /consulting veterinarian, visits meet programmatic needs (Guide, p14)	А				
	۲	Regular communication occurs between veterinarian and IACUC (Guide, p14)	А				
		Veterinarian(s) have experience and training in species used (Guide, p 15) [must]	A				
	•	Veterinarian(s) have experience in facility administration/management (Guide, p15)	А	-	_	1	
20				-	-		
9.	Pe	rsonnel Qualifications and Training	A	M	S	C	NA
		All personnel are adequately educated, trained, and/or qualified in basic principles of					
		laboratory animal science. Personnel included: [must]					
		<ul> <li>Veterinary/other professional staff (Guide, p15-16)</li> </ul>	Α				
		<ul> <li>IACUC members (Guide, p 17)</li> </ul>	A				
-		• Animal care personnel ( <i>Guide</i> , <u>p 16</u> )	Α				
		<ul> <li>Research investigators, instructors, technicians, trainees, and students (Guide, pp)</li> </ul>					
		<u>16-17</u> )	A				
		Continuing education for program and research staff provided to ensure high quality					
		care and reinforce training ( <i>Guide</i> , pp 16-17)	A				
			-				
	•	Training is available prior to starting animal activity ( <i>Guide</i> , p.17)	A				
	•	Training is available prior to starting animal activity ( <i>Guide</i> , <u>p17</u> ) Training is documented ( <i>Guide</i> , p15)	A				
		Training is available prior to starting animal activity ( <i>Guide</i> , <u>p17</u> ) Training is documented ( <i>Guide</i> , <u>p15</u> ) Training program content includes: ( <i>Guide</i> , p17)	A A				
	•	Training is available prior to starting animal activity ( <i>Guide</i> , <u>p17</u> ) Training is documented ( <i>Guide</i> , <u>p15</u> ) Training program content includes: ( <i>Guide</i> , <u>p17</u> ) Methods for reporting concerns ( <i>Guide</i> , <u>p17</u> )	A A				
	•	Training is available prior to starting animal activity ( <i>Guide</i> , <u>p17</u> ) Training is documented ( <i>Guide</i> , <u>p15</u> ) Training program content includes: ( <i>Guide</i> , <u>p17</u> ) • Methods for reporting concerns ( <i>Guide</i> , <u>p17</u> ) • Humane practices of animal care (e.g., housing, husbandry, handling) <sup>xii</sup>	A A A				
	•	Training is available prior to starting animal activity ( <i>Guide</i> , <u>p17</u> ) Training is documented ( <i>Guide</i> , <u>p15</u> ) Training program content includes: ( <i>Guide</i> , <u>p17</u> ) • Methods for reporting concerns ( <i>Guide</i> , <u>p17</u> ) • Humane practices of animal care (e.g., housing, husbandry, handling) <sup>xii</sup> • Humane practices of animal use (e.g., research procedures, use of anosthesia	A A A				

	pre- and post-operative care, aseptic surgical techniques and euthanasia (Guide, p	A				
	17) <sup>xiii</sup>					
	<ul> <li>Research/testing methods that minimize numbers necessary to obtain valid results (PHS Policy, <u>IV.A.1.g.</u>)</li> </ul>	A				
	<ul> <li>Research/testing methods that minimize animal pain or distress (PHS Policy, <u>IV.A.1.g.</u>)</li> </ul>	А				
	<ul> <li>Use of hazardous agents, including access to OSHA chemical hazard notices where applicable (<i>Guide</i>, <u>p 20</u>)</li> </ul>	A				
	• Animal care and use legislation (Guide, p17)	A				
	• IACUC function ( <i>Guide</i> , p 17)	Α	1			
	• Ethics of animal use and Three R's (Guide, p17)	A				
ιο.	Occupational Health and Safety of Personnel	$\mathbf{A}^*$	м	s	с	N
•	Program is in place and is consistent with federal, state, and local regulations ( <i>Guide</i> , $p \ 17$ ) [must]	A	-			
	Program covers all personnel who work in laboratory animal facilities (Guide, p18)	A	-		1	
1.	Changing, washing, and showering facilities are available as appropriate ( <i>Guide</i> , p19)	A				
•	Hazardous facilities are separated from other areas and identified as limited access $(Guide, p 19)$	A				
•	Personnel training is provided based on risk (e.g., zoonoses, hazards, personal hygiene, special precautions, animal allergies) ( <i>Guide</i> , <u>p 20</u> )	A				
- /• (	Personal hygiene procedures are in place (e.g., work clothing, eating/drinking/smoking policies) ( <i>Guide</i> , <u>p 20</u> )	A				
٠	Procedures for use, storage, and disposal of hazardous biologic, chemical, and physical agents are in place ( <i>Guide</i> , $p 21$ )	A				
•	Personal Protective Equipment for the work area is appropriate and available ( <i>Guide</i> , <u>p</u> 21)	A				
•	Program for medical evaluation and preventive medicine for personnel includes:					
	<ul> <li>Pre-employment evaluation including health history (Guide, p22)</li> </ul>	Α				
	<ul> <li>Immunizations as appropriate (e.g., rabies, tetanus) and tests as appropriate (Guide, p 22)</li> </ul>	A				
	<ul> <li>Zoonosis surveillance as appropriate (e.g., Q-fever, tularemia, Hantavirus, plague) (<i>Guide</i>, <u>p 23</u>)</li> </ul>	A				
	<ul> <li>Procedures for reporting and treating injuries, including accidents, bites, allergies, etc. (<i>Guide</i>, <u>p 23</u>)</li> </ul>	A				
	Promotes early diagnosis of allergies including preexisting conditions (Guide, p22)	A			_	
	<ul> <li>Considers confidentiality and other legal factors as required by federal, state and local regulations (<i>Guide</i>, p 22) [must]</li> </ul>	A				
	<ul> <li>If serum samples are collected, the purpose is consistent with federal and state laws (Guide, p 22) [must]</li> </ul>					NA
- e.	waste anestnetic gases are scavenged ( <i>Guide</i> , <u>p 21</u> )	A				
	Hearing protection is provided in high noise areas ( <i>Guide</i> , <u>p22</u> )	Α				
	Respiratory protection is available when performing airborne particulate work ( <i>Guide</i> , <u>p 22</u> )	A				
٠	Special precautions for personnel who work with nonhuman primates, their tissues or body fluids include:					
	<ul> <li>Tuberculosis screening provided for all exposed personnel (Guide, p23)</li> </ul>	А				
	<ul> <li>Training and implementation of procedures for bites, scratches, or injuries associated with macaques (<i>Guide</i>, <u>p 23</u>)</li> </ul>	A				
	<ul> <li>PPE is provided including gloves, arm protection, face masks, face shields, or goggles (<i>Guide</i>, <u>p 21</u>)</li> </ul>	A				
	<ul> <li>Injuries associated with macaques are carefully evaluated and treatment implemented (<i>Guide</i>, <u>p 23</u>)</li> </ul>	A				
•	Occupational safety and health of field studies is reviewed by OSH committee or office ( <i>Guide</i> , $p 32$ )	A				
	Descent of Computer MEN	.*		~	-	
11.	Personnel Security 1157	A	M	5	C	NA
	- proventive measures in place include pro employment ecreening and physical and LT			1		4

•	Preventive measures in place include pre-employment screening, and physical and IT		
	security ( <i>Guid</i> e, <u>p 23</u> )	A	i.

12.	Investigating & Reporting Animal Welfare Concerns	<b>A</b> *	м	s	С	NA
	Methods for investigating and reporting animal welfare concerns are established					
	(Guide, <u>p 23</u> ) [must]	A				
	Reported concerns and corrective actions are documented (Guide, p24)	A				
•	Mechanisms for reporting concerns are posted in facility and at applicable website with instructions ( <i>Guide</i> , $p 24$ )	A				
	<ul> <li>Includes multiple contacts (Guide, p 24)</li> </ul>	A				
	<ul> <li>Includes anonymity, whistle blower policy, nondiscrimination and reprisal protection (<i>Guide</i>, <u>p 24</u>)</li> </ul>	A				

\* A = acceptable

**M** = minor deficiency

S = significant deficiency (is or may be a threat to animal health or safety)
 C = change in program (PHS Policy <u>IV.A.1.a.-i.</u>) (include in semiannual report to IO and in annual report to OLAW)

NA = not applicable

NOTES:

## Veterinary Care

### Date:

1.	Cli	nical Care and Management	$\mathbf{A}^*$	м	s	С	NA
		Veterinary program offers high quality of care and ethical standards ( <i>Guide</i> , p105)					
		[must]	A				
		Veterinarian provides guidance to all personnel to ensure appropriate husbandry,					
		handling, treatment, anesthesia, analgesia, and euthanasia (Guide, p106)	Α				
		Veterinarian provides oversight to surgery and perioperative care (Guide, p106)	Α	<u> </u>			
		Veterinary care program is appropriate for program requirements (Guide, pp 113-114)	Α	-	-		
	•	Veterinarian(s) is familiar with species and use of animals and has access to medical					
	_	and experimental treatment records (Guide, <u>p 114</u> )	Α	1		1.1	-
-		Procedures to triage and prioritize incident reports are in place (Guide, p114)	Α		_	-	
_	•	Procedures are in place to address:		_			
		• Problems with experiments to determine course of treatment in consultation with				<b>.</b>	1.
		investigator(Guide, p114)	A				-
		<ul> <li>Recurrent or significant health problems with the IACUC and documentation of</li> </ul>					
		treatments and outcomes (Guide, p 114)	A				-
_		• Veterinary review and oversight of medical and animal use records ( <i>Guide</i> , <u>p115</u> )	A				
	•	Procedures established for timely reporting of animal injury, illness, or disease ( <i>Guide</i> , <u>p 114</u> ) [must]	A				
	•	Procedures established for veterinary assessment, treatment, or euthanasia ( <i>Guide</i> , <u>p</u> 114) [must]	A				
	•	Veterinarian is authorized to treat, relieve pain, and/or euthanize (Guide, p114)					
_	_		A				
2.	Ar	nimal Procurement and Transportation/Preventive Medicine	$\mathbf{A}^*$	M	S	С	NA
		Procedures for lawful animal procurement are in place ( <i>Guide</i> , p106) [must]	A				1
-		Sufficient facilities and expertise are confirmed prior to procurement ( <i>Guide</i> , p106)	A				1
-		Procurement is linked to IACUC review and approval (Guide, p 106)	A				
		Random source dogs and cats are inspected for identification (Guide, p106)					NA
		Population status of wildlife species is considered prior to procurement ( <i>Guide</i> , p106)	A				
		Appropriate records are maintained on animal acquisition ( <i>Guide</i> , p106)	A			1 - 1	
		Animal vendors are evaluated to meet program needs and guality ( <i>Guide</i> , p106)	A				1
		Breeding colonies are based on need and managed to minimize numbers (Guide, p					
		107)	A	_			
		Procedures for compliance with animal transportation regulations, including					
		international requirements, are in place ( <i>Guide</i> , p 107) [must]	A	_			
		Transportation is planned to ensure safety, security and minimize risk (Guide, p107)	A				
		Movement of animals is planned to minimize transit time and deliveries are planned to					
		ensure receiving personnel are available (Guide, pp 107-108)	A				
		Appropriate loading and unloading facilities are available (Guide, p109)	A				
		Environment at receiving site is appropriate (Guide, p109)	Α				
		Policies in place on separation by species, source, and health status (Guide, pp 109,		-			
	1	<u>111-112</u> )	A			_	
		Procedures in place for quarantine to include zoonoses prevention (Guide, p110)	Α				
	٠	Quarantined animals from different shipments are handled separately or physically					
-	~	Separateu (Galde, <u>p 110</u> ) Dreseduras in place for stabilization (acclimation (Galde, and 10, 111)	A				-
	•	Procedures in place for isolation of cick animation (Guide, pp 110-111)	A			-	-
	•	Protection place for isolation of SICK animals (Guide, p 112)	A			-	-
	•	program is in place for surveillance, diagnosis, treatment and control of disease to					
-		Diagnostic resources are available for proventive booth program (Quide a 112)	A	-			
		Diagnostic resources are available for preventive nearth program (Guide, p112)	A				
3.	Su	irgery	<b>A</b> *	М	S	С	NA
-		Surgical outcomes are assessed and corrective changes instituted (Guide, p115)	Α				
		Researchers have appropriate training to ensure good technique (Guide, p115)	1				
		[must]	A				

1	Da	in Distross Anasthasia and Analgosia	۸*	-		~	NA
	•	Post-operative monitoring and care are provided by trained personnel and documented (e.g., thermoregulation, physiologic function, analgesia, infection, removal of skin closures) ( <i>Guide</i> , pp 119-120)	A				
	•	For aquatic species, skin surfaces are kept moist during surgical procedures ( <i>Guide</i> , $p = 119$ )	A			- 1-	
	•	Procedures for monitoring surgical anesthesia and analgesia are in place ( <i>Guide</i> , $\underline{p}$ <u>119</u> )	A				-
2	•	Effective procedures for sterilizing instruments and monitoring expiration dates on sterile packs are in place ( <i>Guide</i> , <u>p 119</u> )	A				
	•	Aseptic technique is followed for survival surgical procedures ( <i>Guide</i> , <u>pp118-119</u> )	A		1.1	1.11	
-	•	For nonsurvival surgery, the site is clipped, gloves are worn and instruments and area are clean ( <i>Guide</i> , <u>p 118</u> )	A				
	•	Surgical procedures including laparoscopic procedures are categorized as major or minor ( <i>Guide</i> , <u>pp 117-118</u> )	A				
	•	Aseptic surgery is conducted in dedicated facilities or spaces, unless exception justified and IACUC approved ( <i>Guide</i> , <u>p 116</u> )	A				
	•	Pre-surgical plans are developed and include veterinary input (e.g., location, supplies, anesthetic and analgesic use, peri-operative care, recordkeeping) ( <i>Guide</i> , <u>p116</u> )	A				

ι.	Pain, Distress, Anesthesia and Analgesia	Α	M	S	С	NA
	<ul> <li>Guidelines for assessment and categorization of pain, distress and animal wellbeing are provided during training (<i>Guide</i>, <u>p 121</u>)</li> </ul>	A				
	<ul> <li>Selection of analgesics and anesthetics is based on professional veterinary judgment (Guide, p 121)</li> </ul>	A				
	<ul> <li>Painful procedures are monitored to ensure appropriate analgesic management (Guide, p 122)</li> </ul>	A				
	<ul> <li>Nonpharmacologic control of pain is considered as an element of postprocedural care (Guide, p 122)</li> </ul>	A				
	<ul> <li>Procedures are in place to assure antinoception before surgery begins (<i>Guide</i>, <u>p 122</u>)</li> <li>[must]</li> </ul>	A				
	<ul> <li>Guidelines for selection and use of analgesics and anesthetics are in place and regularly reviewed and updated (<i>Guide</i>, <u>p 122</u>)</li> </ul>	A				
	<ul> <li>Special precautions for the use of paralytics are in place to ensure anesthesia<sup>xiv</sup> (Guide, p 123)</li> </ul>	A				

## 5. Euthanasia

E	JINANASIA	A	M	3	C	NA
•	Methods are consistent with AVMA Guidelines on Euthanasia unless approved by the IACUC ( <i>Guide</i> , <u>p 123</u> )	A				
٠	Standardized methods are developed and approved by the veterinarian and IACUC that avoid distress and consider animal age and species ( <i>Guide</i> , <u>pp123-124</u> )	A				
•	Training is provided on appropriate methods for each species and considers psychological stress to personnel ( <i>Guide</i> , <u>p 124</u> )	A				
•	Procedures and training are in place to ensure death is confirmed ( <i>Guide</i> , <u>p124</u> ) [must]	A				
-	rug Storage and Control NEW	۸*	M		~	NA

## 6. Drug Storage and Control NEW

Drug Storage and Control des	m	IM	3	<b>U</b>	MA
<ul> <li>Program complies with federal regulations for human and veterinary drugs(Guide, p 115) [must]</li> </ul>	4		-1		
<ul> <li>Drug records and storage procedures are reviewed during facility inspections (<i>Guide</i>, <u>p</u> <u>115</u>)</li> </ul>	4				
<ul> <li>Procedures are in place to ensure analgesics and anesthetics are used within expiration date (<i>Guide</i>, <u>p 122</u>) [must]</li> </ul>	4				
<ul> <li>Anesthetics and analgesics are acquired, stored, and their use and disposal are recorded legally and safely (<i>Guide</i>, <u>p122</u>)</li> </ul>	4				

\* A = acceptable

**M** = minor deficiency

S = significant deficiency (is or may be a threat to animal health or safety)
 C = change in program (PHS Policy <u>IV.A.1.a.-i.</u>) (include in semiannual report to IO and in annual report to OLAW)

**NA** = not applicable

# Appendices

- a) RAR Training Presentation Slides (pg 45-48)
- b) Inspection Dates (pg 49-51)
- c) IMHA List and Justifications (pg 52-82)
- d) Reduced PAM (pg 83-88)
- e) Complete Inspection Report Summary (pg 89-165)
- f) Approved Protocol Exceptions (pg 166-294)
- g) Administrative Summary and Graphs (pg 295-312)
- h) Expired and Holding Protocol Summary (pg 313)
- i) Repeat Significant Findings (pg 314-318)









#### **Required Training**



- Facility Tour(s)
   Required for all RAR animal facility users
- Microisolator Training • Required for access to RAR SPF facilities
- NHP Training • Required for access to NHP facilities or for lab
  - personnel working with NHP tissues























Facility Name	Facility Code	Spring 2019	Fall 2019
	1	1/17/2019	7/18/2019
	1	1/9/2019	7/15/2019
	1	10/10/2018	4/19/2019
	1	1/17/2019	7/18/2019
	2	1/9/2019	7/15/2019
	2	1/9/2019	7/15/2019
	3	1/17/2019	7/18/2019
	4	1/22/2019	7/31/2019
	5	3/6/2019	9/17/2019
	6	12/14/2018	6/25/2019
	7	2/5/2019	7/31/2019
	8	12/17/2018	6/24/2019
	10	10/10/2018	4/23/2019
	10	10/30/2018	4/23/2019
	10	10/30/2018	4/10/2019
	10	10/30/2018	4/10/2019
	10	10/30/2018	4/10/2019
	10	11/29/2018	5/22/2019
	10	3/29/2019	9/27/2019
	12	11/19/2018	5/17/2019
	12	12/21/2018	6/21/2019
			5/17/2019
	12	3/18/2019	9/25/2019
	12	3/25/2019	9/23/2019
	12	3/13/2019	9/26/2019
	12	Not applicable	Not applicable
	13	10/15/2018	4/25/2019
	14	2/18/2019	8/6/2019
	15	3/28/2019	9/18/2019
	16	11/28/2018	5/15/2019
	17	1/8/2019	7/24/2019
	18	12/13/2018	6/6/2019
	19	10/29/2018	4/29/2019
	20	Not applicable	8/19/2019
	21	1/14/2019	7/26/2019
	22	3/26/2019	9/25/2019
	22	3/5/2019	9/30/2019
IMHA Areas:		Notapplicable	
		I Not applicable	

	2/21/2019	8/6/2019
	3/19/2019	9/30/2019
	1/22/2019	7/26/2019
	3/6/2019	9/10/2019
	Not applicable	9/26/2019
	Not applicable	Not applicable
	Not applicable	Not applicable
	Not applicable	Not applicable
	Not applicable	6/19/2019
	3/21/2019	9/20/2019
	2/19/2019	8/15/2019
	10/25/2018	Not applicable
	12/13/2018	6/6/2019
	2/15/2019	8/16/2019
	2/4/2019	8/16/2019
	Not applicable	Not applicable
	Not applicable	Not applicable
	2/18/2019	Not applicable
	Not applicable	
	Not applicable	Not applicable
	2/8/2019	8/7/2019
-	3/1/2019	8/7/2019
	3/7/2019	9/24/2019
-	Not applicable	Not applicable
	Not applicable	Not applicable
	11/26/2018	5/20/2019
	11/1/2018	5/17/2019
L	Not applicable	Not applicable
	Not applicable	Not applicable
	Not applicable	Not applicable
-	Not applicable	Not applicable
	3/26/2019	9/6/2019
	12/21/2018	6/21/2019
	1/2/2019	6/24/2019
	1/2/2019	6/24/2019
	1/25/2019	Not applicable
	Not applicable	Not applicable
	10/16/2018	4/30/2019
	2/25/2019	8/13/2019

	D=	11/2/2018	5/2/2019
		2/28/2019	8/20/2019
		2/19/2019	8/15/2019
		12/11/2018	6/20/2019
		11/27/2018	5/7/2019
		3/26/2019	9/27/2019
		3/27/2019	9/19/2019
		3/27/2019	9/19/2019
		Not applicable	Not applicable
		12/21/2018	6/17/2019
		2/15/2019	8/29/2019
		Not applicable	8/22/2019
	7	3/13/2019	9/17/2019
		3/18/2019	Not applicable
		Not applicable	8/22/2019
		2/21/2019	8/20/2019
		3/20/2019	9/11/2019
AG Sites:			
		3/20/2019	9/11/2019
	J	10/29/2018	4/17/2019
		10/29/2018	4/17/2019
		12/6/2018	5/31/2019
		12/6/2018	5/31/2019
	· · · · · · · · · · · · · · · · · · ·	11/15/2018	5/30/2019
		11/26/2018	5/10/2019
		11/26/2018	5/10/2019
		12/14/2018	6/21/2019
	0	12/14/2018	6/25/2019
		12/11/2018	6/25/2019
		N/A	6/21/2019
		12/11/2018	C/21/2010

Investigator	Species	Building	Room number	Protocol Number	Justification
Alejandro, Emilyn	mouse			1806-36072A	We need special housing for this mice that is only available in IBP Phenotyping core in CCRB
Aliota, Matthew	mouse	V		1804-35828A	The experiments to be performed are to be done at the second se
Bajer, Przemyslaw	fish		***	1906-37131A	Protocol has not yet been submitted
artolomucci, Alessandro	mouse			1701-34522A, 1706- 34930A, 1711-35305A	Our studies involve continuous monitoring of mice metabolic functions including daily food intake and body weight monitor as well as weekly body composition and indirect calorimetry analysis. Data acquisition systems are extremely expensive and therefore must be in a secured area. Also, it is absolutely crucial to our research that the animals are monitored in an experimentally controlled environment and that we are able to

Barrell, Emily	horse	***	1905-37027A	Horses are housed in the Barn presently as there is no RAR housing available; they will continue to be housed in this barn for the duration of the study.
Baughn, Anthony	mouse		1810-36444A	Mice will be infected with Mycobacterium tuberculosis, a bacterium that is classified as a Risk Group 3 pathogen due to its high infectivity and transmission via the respiratory route. According to the Biosafety in Microbiological and Biomedical Laboratories 5th ed (CDC), mice infected with M. tuberculosis do not pose an aerosol infection risk and can be maintained under services and processing of infected mice does present a significant aerosol exposure risk and must be conducted inside the services for work in the service and caging equipment used for housing the infected animals are designed to reduce the risk of personnel exposure to infectious organisms during husbandry.
Bee, Mark	frogs		1701-34456A	The main reasons for the long term housing of frogs in this room are the following: A major reason for housing these animals outside of RAR managed facilities concerns the time of day that the animals are used. It is likely that some tests will be conducted at night (e.g., between midnight and 4am) whe these frogs are naturally most active. It has been our experience that frogs are often more responsive when tested at night. Second, it is important for our experiments that we maintain some control over the acoustic environment in whic the frogs are kept. We may wish to broadcast recordings of natural frog choruses at night in the animal room to acoustically simulate the situation in which females are in natural breeding chorus.

Bee; Mark	frogs	1701-34456A	The main reasons for the long term housing of frogs in this room are the following: A major reason for housing these animals outside of RAR managed facilities concerns the time of day that the animals are used. It is likely that some tests will be conducted at night (e.g., between midnight and 4am) when these frogs are naturally most active. It has been our experience that frogs are often more responsive when tested at night. Second, it is important for our experiments that we maintain some control over the acoustic environment in which the frogs are kept. We may wish to broadcast recordings of
Belcher, John	mice	1712-35371A	natural frog choruses at night in the animal room to acoustically simulate the situation in which females are in natural breeding chorus. Mice are implanted with dorsal skin fold chambers in IMHA and then kept warm in humidified 32 degree C incubators in IMHA to minimize body heat loss via the implanted chamber.
Benneyworth, Michael	mice	1711-35294A	Our lab studies the effects of environmental experience on brain function and behavior. Our recent studies have shown that even mild stress (handling, injections and/or novelty exposure) can have significant impact on our outcome measures. The IMHA will enable us to minimize unwanted stress and control out animals' any for any for any for
Benneyworth, Michael	mice	1711-35294A	Our lab studies the effects of environmental experience on brain function and behavior. Our recent studies have shown that even mild stress (handling, injections and/or novelty exposure) can have significant impact on our outcome measures. The IMHA will enable us to minimize unwanted stress and control out animals' environment to reduce variability in our data

Bianco, Richard	sheep, pigs	pasture and barn	1610-34277A, 1703- 34665A, 1704-34726A, 1705-34789A, 1707- 34942A, 1710-35219A, 1711-35350A, 1802- 35610A, 1804-35772A, 1804-35780A, 1806- 36011A, 1807-36201A, 1808-36237A, 1810- 36420A, 1903-36852A, 1905-37042A, 1905- 37103A, 1907-37284A	Long term survival animals are provided a natural environment at This facility is capable of providing housing with pasture and appropriate shelter for large number of farm animals. This facility also has a veterinary technician living on the grounds. The technician is capable of blood draws, treatment, observing and assessing clinical health and is GLP completent and maintains the facility under AAALAC and GLP compliancy. Transportation is provided on short notice and during off hours. The facility does provide an excellent enriched environment for the test animals. In order to maintain the quality and scientific integrity of this study as well as to generate a reliable model for future studies, we are required to follow the regulations set by FDA under 21CFR part 58.
Bischof, John	mice		1701-34516A	We received IACUC approval for housing SPF mice in our animal surgery room located in the second structure in early 2002 and have continuously housed animals for this study there since then. In brief, the equipment needed for this project is required by other personnel in the lab at various times and cannot be moved to an RAR area.
Bischof, John	fish embryos		1804-35844A	Fish embryos will be considered vertebrates after they reach Day 3. They will be housed in the from Day 3 to Day 5 for observation. After which any surviving fish will be transferred to the zebrafish core and housed in tanks and grown out til three months. We will attempt the fish to spawn and evaluate the health of the embryos (non-vertebrates). After which the adult parent fish will be euthanized. The cryopreservation and observation procedures can be handled in NHH whereas the housing & care of fish post day 5 can be taken care in the zebrafish core. The zebrafish core has experts and an approved protocol for this procedure (#1506-32642A)

Bischof, John	zebrafish embryos		1804-35844A	Fish embryos will be considered vertebrates after they reach Day 3. They will be housed in from Day 3 to Day 5 for observation. After which any surviving fish will be transferred to the zebrafish core and housed in tanks and grown out til three months. We will attempt the fish to spawn and evaluate the health of the embryos (non-vertebrates). After which the adult parent fish will be euthanized. The cryopreservation and observation procedures can be handled in whereas the housing & care of fish post day 5 can be taken care in the zebrafish core. The zebrafish core has experts and an approved protocol for this procedure (#1506-32642A)
Bitterman, Peter	fish	-	1801-35483A	Zebrafish research facility, <b>Constant (IMHA)</b> is used only for egg/embryo production
Boe, Gail	fish, birds	***	1806-36052A	For display at UMM
Bold, Tyler	mouse		1812-36571A	Mice will be infected with Mycobacterium tuberculosis, a bacterium that is classified as a duration of the respiratory route. Mice infected with M. tuberculosis may release infectious bacteria to the air. Therefore, infected mice must be housed inside the duration of the air of the field of the duration of the field infectious organisms. Standard operating procedures for work in the duration and caging equipment used for housing the infected animals are designed to reduce the risk of personnel exposure to infectious organisms during husbandry.
Chen, Xiaoli	mouse		1801-35539A	Our studies involve continuous monitoring of mice metabolic functions including daily food intake and body weight monitor as well as weekly body composition and indirect calorimetry analysis. Data acquisition systems are extremely expensive and therefore must e in a secured area. Also, it is absolutely crucial to our research that the monitored in an experimentally controlled environment and that we are able to directly monitor the animals at all times.

Chen, Xiaoli	mouse	***	1801-35539A	Our studies involve continuous monitoring of mice metabolic functions including daily food intake and body weight monitor as well as weekly body composition and indirect calorimetry analysis. Data acquisition systems are extremely expensive and therefore must e in a secured area. Also, it is absolutely crucial to our research that the monitored in an experimentally controlled environment and that we are able to directly monitor the animals at all times.
Chen, Clark	mouse		1802-35597A	Also, for the second we request IMHA for up to 72 hours postop housing of animals undergoing survival surgeries. As described in the text, following a survival surgery the animal is placed in a clean cage in room the survival surgery the animal is placed in a the post-operative evaluation and special care of the mice with implants.
Collister, John	rats		1612-34405A, 1810- 36452A	The required equipment using in recording blood pressure and heart rate is located in room
Cvetanovic, Marija	mouse		1810-36435A	Behavioral test done by behavioral core. Recent studies have shown that even mild stress (handling, injections, and/or novelty exposure) can have significant impact on our outcome measures. The IMHA will enable us to minimize unwanted stress (e.g., transport from housing colony to lab) and control our animals' environment to reduce variability in our data
Cvetanovic, Marija	guinea pigs, frog, rat, mouse		1703-34631A	the proposed animals will be used for a teaching lab taught at the Itasca Biological Research Station at Lake Itasca State Park. This is sufficiently far from the Twin Cities that central RAR housing would not be possible
Denton, Robert	salamanders		1901-36686A	This secured cold room provides controlled conditions necessary to keep the salamanders at a cool temperature.

Desrosiers, Mark	turtle		1810-36465A	There are no turtle housing facilities in Research Animal Resources, and this is a display animal in the libraries
Dudley, Samuel	mouse		1703-34647A, 1704- 34710A	Use of telemetry system
Dougherty, Brendan	rat	Ĵ	1704-34724A	Rats receiving experimental spinal cord injuries receive specifi post-op care and monitoring to ensure appropriate recovery. We have found this to be best handled within the laboratory environment by trained staff with access to specific equipmen and drugs for the first 24-72 hours.
Ebner, Timothy	mouse		1803-35638A, 1808- 36330A	We request IMHA for up to 72 hours postop housing of animals undergoing survival surgeries. As described in the text following a survival surgery the animal is placed in a clean cage and the cage placed in the fume hood in room purpose of this is to facilitate the post-operative evaluation and special care of the mice with implants.
Engeland, William	mouse		1710-35244A	There are two reasons for housing mice in the <b>Second</b> First it enhances our ability to optimally perform chronic subordinate stress experiments, since the IMHA is the facility being used by our collaborators, Drs. A. Bartolomucci and M. Razzoli who have the experience required to effectively complete these experiments. Secondly the current housing space in the second stress and more importantly, my collaborators emphasize the importance of minimizing exposure of mice not involved in the experiment to mice undergoing daily bouts of dominant subordinate interactions. Since we house and breed mice for other experiments this space is not available fo the subordinate stress experiments

				Our studies involve continuous monitoring of mouse metabolic functions including body composition and indirect calorimetry
Ervasti, James	mouse		1806-36018A	analysis. Data acquisition systems are extremely expensive and therefore must be in a secured area. Also, it is absolutely crucial to our research that the animals are monitored in an experimentally controlled environment and that we are able to monitor the animals at all times.
Fallon, Ann	hamsters	2 <b></b> 2	1902-36743A	Blood-feeding arthropods cannot be transported to another building to be provided a blood meal.
Ferrington, Deborah	mice		1704-34752A	Rooms listed is where we have a hood set up to perform light stress experiments. These experiments require stringent control of the amount and timing of light so need to be performed outside the normal RAR facility where a 12/12 cycle of light and dark is maintained.
Firshman, Anna	Horse	***	1708-35084A	No Justification provided
Garry, Mary	pigs		1905-37039A	Young piglets require very close attention and specialized care. This level of care is provided by laboratory personnel.
Garry, Mary	pīgs		1806-36050A	Young piglets require very close attention and specialized care. This level of care is provided by laboratory personnel.
Gewirtz, Jonathan	mouse	-	1609-34140A	Animals are to be transported to a surgical procedures and behavioral experiments that are unavailable within the surger of the current housing is 'conventional', mice must be transported directly to the IMHA (a conventional space)

	NHP, mouse, rat	1706-34873A, 1706- 34898A, 1706-34903A, 1707-34995A, 1708- 35062A, 1805-35937A, 1806-35989A, 1806- 36065A, 1808-36291A, 1810-36463A, 1902- 36813A, 1902-36830A, 1903-36845A, 1904- 36948A, 1905-37026A	We have modified husbandry practices to be optimal for NHP and rodents used in complex disease models. This IMHA is capable of exceeding minimum expectations o the guide to provide our animals with varied enrichment, careful husbandry scheduling accommodating the highest level of care and complex environments/interactions that provide the best opportunity for expression of behaviors that represent the species typical repertoire.
Gregerson, Dale	mouse	1706-34882A	Maintenance of a specific, bright light environment continuously for up to 4 days can not be accommodated in RAR mouse rooms.
Griffith, Thomas	mouse	1906-37113A	We are conducting a study that requires mice to be housed in a personnel. Therefore, it is necessary to obtain clearance for the self management and maintenance of our mice housed in this
Hackel, Benjamin	mice	1710-35191A	Mice will be injected IV with 64Cu-labeled proteins. Housing in enables them to be kept in a remote, shielded location while radioactivity is present. This radioisotope has a 12.7 hour half life. Mice will be housed between imaging experiments (up to 48 hour post injection). Mice that are not euthanized for excised tissue biodistribution will be housed in until radioactivity decays to background (typically 10 half lives (127 hours))

Harmon, James	sheep	***	1908-37287A	Long term survival animals are provided a natural environme at This facility is capable of providing housing with pasture and appropriate shelter for a large number of farm animals. This facility also be a veterinary technician living on the grounds. The technician capable of blood draws, treatments, observing and assessin clinical health, and is GLP competent and maintains the facil under AAALAC and GLP compliancy. Transportation is provid on short notice and during off-hours. The facility does provid an excellent enriched environment for study animals. In ord to maintain the quality and scientific integrity of this study as well as to generate a reliable model for future studies, we a required to follow the regulations set by FDA under 21CFRp. 58. <sup>a</sup>
Haskell-Leuvano, Carrie	mice		1702-34552A, 1705- 34823A	To house and have access to the specialized TSE mouse metabolic cages described in the protocol. Depending on the experiment (exercise and feeding), the experimental mice need to be continuously housed in these cages for data collection purposes for up to 9 weeks of experimental and 1 weeks equilibration
Haskell-Leuvano, Carrie	mice	-	1702-34552A	To acclimate the mice to a different light schedule and to acclimate the mice to the new room before the behavior testing starts.

Hecht, Stephen	rats		1908-37306A	The AeroCore is a University ESO/ISO (external/internal service organization). AeroCore provides animal testing services, as such, the animals are housed in the IMHA and all services are performed in the IMHA.
Heîmpel, George	bīrds	***	1804-35830A	The zebra finches will be used to rear a quarantined insect. A certified quarantine facility is therefore needed to do the research
Henke, Craig	mice		1610-34253A, 1909- 37429A	Hypoxia chamber studies
Hove, Mark	fish		1902-36808A	To complete the life history and behavior studies mussels, animals (fish, amphibians, and aquatic invertebrates) need to be held in aquaria for experiments and observation. Recovery of microscopic mussel larvae from laboratory and naturally infested animals needs to be done in aquaria. Observations made during these studies will be used to improve natural resource management decisions.

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Hrabik, Thomas	fish		1612-34364A, 1803- 35684A	1612-34364A: This is a UROP provided by the University of Minnesota- Duluth. We need to house the fish here because it is the only place where there is enough space to hold all (6) of our holding tanks. We also have the rooms in which the tanks will be placed where we can control the amount of light and temperature.
lkramuddin, Sayeed	mouse		1612-34402A	Metabolic testing (Indirect Calorimetry, Meal pattern analysis, body composition) are only conducted at the se evaluations only 40 mice from this protocol will be operated or and housed at the second
	NHP		1901-36714A	Animals must be quarantined after MPTP injection (either IM or intracarotid) and are held in a room adjacent to the During this 72 hour period in which the animal is present, the room will be designated as an IMHA, otherwise it is managed by RAR.
Kawakami, Yasuhiko	zebra fish		1704-34728A, 1908- 37300A	No RAR housing is available for zebrafish. The zebra fish facility in <b>present of the setablished and utilized for years by</b> various research groups, therefore the facility is functionally able to house zebrafish.
Kim, Do-Hyung	moùse		1712-35414A	The protocol includes analysis of mouse physiology, such as the analysis of whole body fat content and food uptake. These assays are available at the Mouse Phenotyping Core in (Director: Dr. Alessandro Bartolomucci). The mice in the area will be kept 1-2 weeks before the assays are conducted. Once all the assays are completed, mice will be sacrified and tissues will be collected

Kotz, Catherine	mouse		1701-34527A, 1705- 34831A, 1706-34859A, 1902-36754A	In our study we will examine effects of optogenic stimulation/inhibition of orexin neurons in context of circadian rhythm. Our studies will include both calorimetry and SPA measurements as well as running wheel studies longer than 24h. Since those kind of observations can not be performed in we need to use IMHA
Kozak, Kenneth	salamanders		1704-34734A	Transporting salamanders between housing and testing facilities that are in different buildings has the potential to alter their basal metabolic rates by exposing them to sunlight and outdoor temperatures. Given that the animals are tested over a period of several months, this could adversely affect the results of the experiments as the outdoor climate fluctuates
Krook-Magnuson, Esther	mouse		1801-35497A	Our optogenetic experiments are done with 24-7 video EEG monitoring, and animals are tethered to allow light delivery. This requires special cages and investigator managed housing.
Kurtti, Timothy	hamsters		1809-36343A, 1904- 36955A	Because of difficulties and special facilities needed to contain ticks, and on occasion infected ticks, the animals will be held outside of the primary housing area. When animals are infested with ticks they need to be examined several times daily. Infected animals, cultures and ticks are handled by this team that has extensive training and experience in the handling of human and animal disease agents transmitted by arthropods.
Kyba, Michael	mouse		1708-35046A	Our studies involve continuous monitoring of mice metabolic functions including daily food intake and body weight monitor as well as weekly body composition and indirect calorimetry analysis. Data acquisition systems are extremely expensive and therefore must be in a secured area
Langlios, Ryan	mouse, rat		1811-36488A	We are conducting a study that requires rats to be housed in a ) that cannot be serviced by RAR personnel. Therefore, it is necessary to obtain clearance for the self-management and maintenance of our rats housed in this

Lee, Anna	mouse		1706-34906A, 1802- 35545A	An IMHA is required because the voluntary drug consumption studies must be conducted in home cages in a room with minimal noise and low foot traffic.
Lemos, Julia	mouse		1801-35436A	Many of the studies conducted in the laboratory entail prior stress or drug exposure and subsequent behavioral or electrophysiology assays. We are requesting that a subset of "in use" animals will be held in the IMHA (Mathematical Stressors) in the lectrophysiology apparation or der to minimize/prevent extraneous forms of environmental stressors introducing variability into our experimental procedures. By reducing variability, this measure will reduce of the number of animals needed to fulfill the experimental mission of our laboratory.
Liang, Jennifer	zebra fish		1703-34641A, 1708- 35044A, 1804-35821A	The provided housing is a state of the art aquatic system for housing zebrafish. This is not available anywhere else on campus that has a state of the art aquatic zebrafish facility that has been running since 2009. There are no other appropriate facilities for zebrafish on campus. Fish before 10- 14 dpf will be in petri dishes in incubators to the recirculating aquatic system in

Līn, Gufa	frogs, axolotis		1703-34677A	Tadpoles and axolotl juveniles need to be kept in the lab until feeding is well established and they are large enough to survive in the recirculating aquarium. Secondly, tadpoles or froglets that are under current experimentation, for example receiving daily warm pulse for transgene induction, or being kept in water with an added substance, or requiring regular observation under fluorescence microscope, will be kept in the lab. In the case of the added substance this cannot be done in the RAR area because the aquaria are recirculating type. For ongoing experiments requiring daily intervention the justification is that the tadpoles should not be subjected to frequent water changes and transport unnecessarily, as this may compromise their health and the success of the experiments. Some tadpoles or axolotls will be subjected to recording under light fluorescent microcopies which is not available in RAR facilities
Lin, Jizhen	mouse		1610-34251A	Once mice are injected with Radioactive Copper (Cu64) they are kept in the catch room adjacent to the machine behind lead for safety reasons.
Lowe, Dawn	mice		1907-37248A	Testing with sensitive physiology equipment that would be better suited in an investigator managed housing area rather than an RAR run facility because access will be limited to those familiar with the study
Lund, Troy	zebra fish	-	1906-37111A	
Madill, Scott	horses	***	1906-37132A, 1906- 37140A, 1906-37178A, 1907-37280A	No Justification provided

Martinson, Krishona	Horse		***	1808-36231A	No Justification provided
Martin, Cindy	pigs	_		1803-35699A	Animals will be undergoing a surgical procedure (arterial switch and atrial septectomy). It is anticipated that animals may need to stay anesthetized and supported by a ventilator for 12-48 hours to have time to recover from the surgical procedure.
Maragi, Frank	fish, amphibians, reptiles			1701-34515A	Fish and reptile species (i.e. turtles) are used in the Department of Biology 'Living Wall' to display animals discussed in Animal Physiology, Developmental Biology, and Genetics courses. It is necessary to house them in the classroom for students to observe during course instruction and discussion.
Mand, Sandy	fish			1611-34300A, 1702- 34545A, 1705-34846A, 1712-35413A	These are animals in laboratory classrooms where students car observe them. This IS their primary housing area, but it is not an RAR facility. RAR does not typically house fish.
Mand, Sandy	fish	-		1702-34545A	These are animals in laboratory classrooms where students car observe them. This is their primary housing area but it is not an RAR facility. RAR does not typically house fish. The advantage to housing in this area is that the tanks are fed by well water.
Mand, Sandy	fish, axolotis, reptiles			1611-34300A, 1702- 34545A, 1705-34846A, 1712-35413A, 1811- 36504A, 1907-37285A	These are animals in laboratory classrooms where students car observe them. This IS their primary housing area, but it is not an RAR facility. RAR does not typically house fish.
Malone, Erin	horses, sheep, cow, donkey, goat, camelids		¥*#¥	1805-35927A	Animals will be housed at the ast this is also the location of the student laboratories. By housing the animals near the students and at the lab site, we are able to minimize the risk and stress of transport across campus and to maximize animal use during the relevant time period.

Mashek, Doug	mouse	1704-34729A	Our studies involve continuous monitoring of mice metabolic functions including indirect calorimetry analysis and locomotion. Data acquisition systems are extremely expensive and therefore must be in a secured area. Also, it is absolutely crucial to our research that the animals are monitored in an experimentally controlled environment and that we are able to directly monitor the animals at all times.
Masino, Mark	zebra fish	1806-36051A, 1904- 36944A, 1905-37035A	Usage of zebrafish as a model organism requires the ability to breed fish to produce embryos for experimentation. As embryo production must be large enough to provide statistically meaningful results and embryos must be used within a few minutes of fertilization, in-house production of embryos is the only solution. This IMHA will also be housing and caring for additional animals (zebrafish) that are found on other protocols.
Masino, Mark	zebra fish embryos	1905-37035A	Our lab uses embryos/larvae from 1-7dpf for experiments, so we house them in the lab.
Masopust, David	mouse	1609-34184A, 1902- 36825A	We are conducting a study that requires mice to be housed in a that cannot be serviced by RAR personnel. Therefore, it is necessary to obtain clearance for the self-management and maintenance of our mice housed in this
McGaugh, Suzanne	fish	1705-34800A, 1906- 37158A, 1906-37186A	These animals need to be in a restricted and solitary space so we can easily set up and conduct behavior assays with them. RAR does not typically house fish.
McGaugh, Suzanne	fish	1705-34800A, 1906- 37158A, 1906-37186A	These animals need to be in a restricted and solitary space so we can easily set up and conduct behavior assays with them. RAR does not typically house fish.
Menken, Jennifer	snake, turtle, salamander, toad, gecko, fish	1611-34357A	The animals listed are part of the educational programming at the Bell Museum. They are used for display in the "Touch and See Room" as well as for demonstration in on-site and off-site classroom room programs. They are an important part of our programming with the public, they are used to demonstrate anatomy, ecology, evolutionary adaptations and biological diversity

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Mensinger, Allen	fish, frog			1807-36111A	The course tried to integrate physiology with behavior. We house the frogs in the lab so the students can observe their behavior and correlate with the experiments. The students ar also instructed in basic animal handling and care techniques and by having the frogs in the lab, we can teach the students this aspect of a science lab; we prefer the students be able to observe the behavior of the weakly electric fish
Mensinger, Allen	fish			1801-35509A, 1801- 35507A, 1903-36856A	Facility was built to specifically house aquatic animals
Mermelstein, Paul	rat			1809-36379A, 1811- 36486A	We plan to perform behavioral testing on rats in specially- constructed operant chambers as described in Experiment 10 of the approved protocol. This testing will be performed in modified to allow us to run our behavioral testing protocol. The only housing for rats in the second stress, and it will not be possible for us to maintain SPF within our operant chambers. Furthermore, SPF procedures would provide additional stress to our rats, and potentially interfere with the results of our behavioral testing.
Metzger, Joseph	mouse			1706-34883A	Our lab recently moved to and the and the second se
Michael, Kerry	rat			1605-33704A	The Morris campus of the University is physically isolated from the rest of the University systema three hour drive from the Twin Cities campus. It is therefore necessary for the sake of research productivity that we maintain our animals on site at the Morris campus.

More, Swati	mouse	1906-37128A	The telemetry system is only offered at the source offers this service. The mice must be housed in this area during the complete period of the study and they can not be taking in and out to avoid any stress to the mice. We will use the husbandary SOP for the source of the study and they can not be taking in and out to avoid any stress to the mice. We will use the husbandary SOP for the source of the study and they can not be taking in and out to avoid any stress to the mice. We will use the husbandary SOP for the source of the study and they can not be taking in and out to avoid any stress to the mice. We will use the husbandary SOP for the source of the study and they can not be taking in and out to avoid any stress to the mice. We will use the husbandary SOP	
Munderloh, Ulrike	mice, hamster	1607-33953A, 1804- 35774A, 1905-37105A	Because of difficulties and special facilities needed to contain ticks, and on occasion infected ticks, the animals will be held outside of the primary housing area. When animals are infested with ticks they need to be examined several times daily.	
Nakagawa, Yasushi	mouse	1712-35389A	Equipment for behavioral tests is housed in the and it will take up to 8 weeks to complete these tests. Therefore, mice housed in the second of the located in the second during behavioral testing. During sucrose preference test, we need to house the mice in IMHA.	
Netoff, Tay	rat	1704-34730A	We need to video record the rats 24/7 for 15 days to find whether or not they have behavioral seizures	
Niedernhofer, Laura	mouse	1808-36256A	We are conducting a study that requires mice to be housed in a better the serviced by RAR personnel. Therefore, it is necessary to obtain clearance for the self-management and maintenance of our mice housed in this	
Oliver, Jonathan	mouse		1609-34130A	Because of difficulties and special facilities needed to contain ticks, and on occasion infected ticks, the animals will be held outside of the primary housing area. When animals are infested with ticks they need to be examined several times daily. International was designed with the input and advice of the late Dr. Patrick Manning and brought to standard according to his recommendations. Access to rooms in and is limited to the principal investigators RAR personnel, technicians and students working on these projects. Infected animals, cultures and ticks are handled by this team that has extensive training and experience in the handling of human and animal disease agents transmitted by arthropods, The procedures for handling the animals are written out in detail in our laboratory SOP and all of our laboratory personnel are instructed in the procedures in case of emergency or absence of the principal investigators.
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Olson, Erin	fişh	1 <b>8</b> 48	1904-37007A	The fish under this protocol are housed for display in an office
Ondrey, Frank	mouse		1712-35415A, 1806- 36059A, 1902-36832A, 1905-37092A	We have had our own research facility (IMHA) for more than 40 years. This is our own research project, and specialized equipment is housed in the facility.

Ondrey, Frank	mouse		1806-36059A	Mice will be imaged five times each, when radioisotope copper 64 is injected, at Time Zero, and 12, 24, 36 and 48 hours, post injection. During this time the mice will be housed in the . This the second is intended to hold radioactive research animals, and is only accessible from . Once imaging is complete, these mice will be euthanized via cervical dislocation, under Isoflurane anesthesia. Carcasses will be disposed of in house, allowing the radioactivity to dissipate before disposal. 64Cu has a half life of 12.7 hours. Caging will be collected from radioactivity has dissipated.
Osborn, John	rat		1612-34416A, 1708- 35043A, 1805-35904A, 1812-36628A	Mice involved in this protocol will be instrumented with radio telemeter equipment which requires constant monitoring and measurement. In addition, animals waiting to be implanted with this equipment will also be housed in this room to reduce the stress of being moved prior to the study
Palmer, Melissa	fish	***	1711-35315A	The IMHA is needed in order to have the fish species on display for educational purposes. The location of the IMHA is crucial for the purpose of the display and need of the animals.
Patterson, Ned	dog		1901-36697A	For the at least the first 24 hours (and longer at the discretion of the study PI) the dogs will be in the that is staffed 24 hours a day by certified veterinary technicians; first technicians; first technicians; first technicians who check each patient at least once every hour. For any dog out of the RAR housing and in ICU or in the for more than 1 week, the PI will have another VMC clinician in the appropriate medical area co-manage the case, and the other clinician will have fina authority on all medical decisions that are not directly indicated in the protocol or by IACUC guidelines.

Paulsen, Megan	mouse		1708-35022A	Studies involve continuous monitoring of mice metabolic functions including daily food intake and body weight monitor as well as weekly body composition and indirect calorimetry analysis. Data acquisition systems are extremely expensive and therefore must be in a secured area. Also, it is absolutely crucial to our research that the animals are monitored in an experimentally controlled environment and that we are able to directly monitor the animals at all times.
Peterson, Lisa	mouse	-	1610-34220A	The equipment for the exposure of animals to the inhaled aldehyde vapors is in this area
Ponder, Julia	birds	****	1705-34793A, 1901- 36695A	The <b>Security Constant of</b> has cages, flight rooms and facilities specifically designed for the safe housing of raptors as well as technical staff with extensive experience in managing these birds. Specific needs include cage size, limited external visibility and perch designs
Portoghese, Phillip	mouse		1702-34546A, 1809- 36366A	In a number of our experiments we test the mice for acute tolerance. This requires the animals to be brought up the nigh- before LPS injection wait another 24 hours for testing of the compound and another 48 hours to test the ED80 dose of the drug to see if there is tolerance. for the sickle cell mice, these animals are quite fair and at this time we are unsure how they will respond to the treatment. There will not be a large number being used at one time and we would like to keep them upstairs for 72 hours for observation
Primus, Alexander	fish		1610-34232A	Housing for fish (larger than zebrafish) is not readily avlaiable in RAR facilities (that I am aware of). Also, this lab space is designed for housing aquatic organisms.

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Primus, Alexander	fish (various species)		1610-34203A	One of the goals of this course is to train students to manage or work in an aquaponics facility. Therefore students in this course need hands-on experience with live fish. The plant growth facility is also an ideal setting for the course because most aquaponics facilities in north-temperate climates rely on greenhouses for light and heat through winter.
Prīmus, Alexander	fish	***	1808-36276A	Fish will require high quality care to eliminate confounding variables of poor fish health for the experiment. Staff at MCL have the expertise and equipment required to maintain this level of care.
Robinson, Jerid	mouse		1906-37 <u>1</u> 64A	mice involved in this protocol will be instrumented with radio telemeter equipment which requires constant monitoring and measurement. This equipment is housed in the second of the sec
Rothwell, Patrick	mice		1810-36447A	Our lab studies the effects of behavioral experience on brain function and behavior. My previous studies have shown that even mild stress (handling, injections, and/or exposure to a novel environment) can have significant impact on our outcome measures. The IMHA will enable us to minimize unwanted stress (e.g., transport from housing colony to lab), and control our animals' environment to reduce variability in our data.

Rothwell, Patrick	mice	1810-36447A	Our lab studies the effects of behavioral experience on brain function and behavior. My previous studies have shown that even mild stress (handling, injections, and/or exposure to a novel environment) can have significant impact on our outcome measures. The IMHA will enable us to minimize unwanted stress (e.g., transport from housing colony to lab), and control our animals' environment to reduce variability in our data.
Ruan, Hai-Bin	mice	1702-34582A, 1811- 36529A	Our studies involve continuous monitoring of mice metabolic functions including daily food intake and body weight monitor as well as weekly body composition and indirect calorimetry analysis. Data acquisition systems are extremely expensive and therefore must be in a secured area. Also, it is crucial to our research that the animals are monitored in an experimentally controlled environment and that we are able to directly monitor the animals at all times.
Phelips, Nicholas	fish	1611-34299A	The <b>Example of the UMN</b> course Hort/FW4601 Aquaponics. It is an ideal setting for the course and our research because most aquaponics facilities in north-temperate climates rely on greenhouses for light and heat through winter.
Schwertfeger, Kaylee	mouse	1909-37381A	We are conducting a study that requires mice to be housed in a that cannot be serviced by RAR personnel. Therefore, it is necessary to obtain clearance for the self management and maintenance of our mice housed in this
Shimizu, Yoji	mouse	1802-35542A	We are conducting a study that requires mice to be housed in a that cannot be serviced by RAR personnel. Therefore, it is necessary to obtain clearance for the self management and maintenance of our mice housed in this

Sivaramakrishnan, Sivaraj	çichlids		1805-35886	RAR does not maintain Cichlid facilities
Smanski, Mîchael	fish	-	1904-36985A	There is no non-IMHA housing for zebrafish on campus
Smanski, Michael	fish		1711-35334A	These are animals in a laboratory classroom where students can work with and observe them. This is their primary housing area but it is not an RAR facility. RAR does not typically house fish
Sorensen, Peter	fish	***	1712-35381A, 1904- 36985A	RAR does not possess fish holding facilities that are either large enough or have well water for our carps
Sorensen, Peter	fish		1712-35381A	RAR does not possess fish holding facilities that are either large enough or have well water for our carps.
Spencer, Sade	mouse		1804-35790A, 1804- 35806A	Many of the studies conducted in the laboratory entail prior stress or drug exposure and subsequent behavioral or electrophysiology assays. We are requesting that a subset of "in use" animals will be held in the IMHA on ) that is in close proximity to the behavioral and electrophysiology apparati in order to minimize/prevent extraneous forms of environmental stressors introducing variability into our experimental procedures. By reducing variability, this measure will reduce of the number of animals needed to fulfill the experimental mission of our laboratory.

Stephens, David	bīrds	-	1707-34943A	We house blue jays and starlings in our facility in the <b>second starlings</b> in our facility in the <b>second starling</b> so that we can study their behavior as described in the accompanying protocol. The <b>second starling</b> allows us to maintain our blue jay colony in a large room adjacent to the procedure rooms; that is also readily accessible to our offices and data analysis facilities.
Stromnes, Ingunn	mouse		1703-34658A	We are conducting a study that requires mice to be housed in a personnel. Therefore, it is necessary to obtain clearance for the self-management and maintenance of our mice housed in this
Thayer, Stanley	rat		1612-34372A	These animals will be used for overnight, 24 hour sessions of ÉEG testing. Moving them back and forth from the will induce stress. Animals housed for three weeks in IMHA
Thayer, Stanley	rat		1612-34372A	These experiments require that the mice remain isolated to avoid stress that could disrupt behavioral testing.
Thomas, Mark	mice		1711-35337A	Our lab studies the effects of environmental experience on brain function. Our recent studies have shown that even mild stress (handling, injections and/or novelty exposure) can have significant impact on our outcome measures. The IMHA will enable us to minimize unwanted stress (e.g. transport from housing colony to lab) and control our animals' environment to reduce variability in our data
Thomas, Mark	rat		1711-35358A	Our lab studies the effects of environmental experience on brain function. Our recent studies have shown that even mild stress (handling, injections and/or novelty exposure) can have significant impact on our outcome measures. The IMHA will enable us to minimize unwanted stress (e.g. transport from housing colony to lab) and control our animals' environment to reduce variability in our data

Thomas, Mark	mice		1711-35337A	Our lab studies the effects of environmental experience on brain function. Our recent studies have shown that even mild stress (handling, injections and/or novelty exposure) can have significant impact on our outcome measures. The IMHA will enable us to minimize unwanted stress (e.g. transport from housing colony to lab) and control our animals' environment to reduce variability in our data
Tischler, Anna	mice		1707-34999A, 1709- 35107A, 1804-35785A	Mice will be infected with Mycobacterium tuberculosis, a bacterium that is classified as a due to its high infectivity and transmission via the respiratory route. Mice infected with M. tuberculosis may release infectious bacteria to the air. Therefore, infected mice must be housed inside the due to contain the highly infectious organisms. Standard operating procedures for work in the design equipment used for housing the infected animals are designed to reduce the risk of personnel exposure to infectious organisms during husbandry.
Todd, Jeffrey	cat, dogclient owned		1906-37145A	cats and dogs are only hospitalized within our hospital, the , until they are adopted or fostered as part of the blood donor program
Townsend, DeWayne	mice		1810-36460A	some of the studies proposed use specialized equipment that cannot be placed into the standard RAR managed rooms

Tranquillo, Robert	sheep		***	1701-34530A, 1709- 35109A	complete with outstanding methods of care, husbandry and research practices. It is capable of providing housing for a large number of animals with extended survival time-points. The farm is GLP compliant and is inspected biannually by University of MN IACUC and monthly by RAR veterinarians
Trent, Ava	horse, camelids	al	***	1712-35369A	Horses and camelids that are free of infectious disease and are not intended for advanced imaging will be housed at the disease and/or intended for practice with the advanced imaging modalities will be housed in the
Tretyakova, Natalia	mouse		•	1706-34933A	The equipment for the exposure of animals for the inhalations is in this location, which is the <b>second of</b> IMHA. The is an External Service Organization/Internal Service Organization for the University. This study is an ISO project.
Trumble, Troy	horse		***	1710-35272A, 1902- 36738A	1710-35272A: No Justification provided; 1902-36738A: RAR does not house horses
Trumble, Troy	horse.		***	1902-36738A	1902-36738A: No other housing option on Twin Cities/St. Paul campus for housing horses

	NHP		1904-36959A	Animals must be quarantined after MPTP injection (either IM or intracarotid) and are held in a room adjacent to the second second second second second second second second animal is present, the room will be designated as an IMHA, otherwise it is managed by RAR
Vossel, Ketih	mouse		1703-34627A	While animals are being recorded, they must be connected by insulated wire to data acquisition that feeds into AHC-secured devices. This system requires a significant amount of electrical power availability, as well as personnel who are familiar with the apparatus and data acquisition software to be able to identify and troubleshoot problems.
Wagner, Carston	moùse		1807-36095A	Currently we are approved for IMHA housing and have passed all required inspections for the animals we keep in our lab. The animals we keep in the lab have some type of neuropathy or turnor that requires constant supervision to make sure they are eating, clean bedding and to watch turnor growth. They also require testing more frequently and the stress of moving them back and forth from would provide an additional stress.
Ward, John	frogs		1902-36788A	Housing allows daily post operative monitoring by the PI to ensure that the frog does not have negative consequences to surgery
Weaver, Cyprian	newts		1612-34381A	We will be performing the resection surgery on the animals. After surgery they need to be continuously monitored for at least three days of surgery for any kind of discomfort or distress
Waye, Heather	snakes, amphibians		1701-34462A, 1710- 35172A, 1711-35353A, 1901-36655A, 1907- 37208A	These animals are housed at the where they are used for display purposes or experimental subjects in a variety of classroom situations/laboratory research
Wefel, Sara	horse	***	1708-35045A	No Justification provided

Wiedmann, Timothy	mouse	ă)	1610-34236A, 1611- 34288A	We have had our own research facility for more than 40 years. This lab is an External Service Organization/Internal Service Organization for the University. This project will be an ESO project.
Wong, Henry	mouse		1803-35719A	In brief, animals for screening are transported to the IMHA and allowed to acclimate for 24 hours prior to the start of any study. The mice are housed for all experimental manipulations for the duration of that particular study since they are tested or administered with compound daily. Animals would experience undue stress being transferred back and forth to the holding room everyday. In addition, the investigator will be more easily able to observe for any adverse effects. The longest duration study is the chronic tolerance test where mice will be housed in the IMHA for a maximum of 9 days. The animals are injected twice a day and are observed for any adverse effects due to the concentration of the compound being administered.
Wong, Henry	mouse		1803-35719A	In brief, animals for screening are transported to the IMHA and allowed to acclimate for 24 hours prior to the start of any study. The mice are housed for all experimental manipulations for the duration of that particular study since they are tested or administered with compound daily. Animals would experience undue stress being transferred back and forth to the holding room everyday. In addition, the investigator will be more easily able to observe for any adverse effects. The longest duration study is the chronic tolerance test where mice will be housed in the IMHA for a maximum of 9 days. The animals are injected twice a day and are observed for any adverse effects due to the concentration of the compound being administered.

. Zordoky, Beshay	mouse			1807-36187A	The mice for the stress studies will be housed in the stress studies will be cause our studies involve continuous monitoring of mice metabolic functions including daily food intake and body weight monitor. Also, it is absolutely crucial to our research that the animals are monitored in an experimentally controlled environment and that we are able to directly monitor the animals at all times for the behavioral assessment.
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David Stephens	
4/19	Quarterly Inspections
5/28/19	Veterinary Consult
6/19	Quarterly Inspections
7/19	Quarterly Inspections
8/16/19Semi-annual	No Deficiencies
9/19	Quarterly Inspections

Cyprian Weaver	
4/30/19	Veterinary Consult
5/19	No vet consults as all newts have been euthanized

Anthony Baughn	
4/22/2019	Veterinary Consult
5/20/19Semi-annual	No Deficiencies
6/17/19	Veterinary Consult
7/15/19	Veterinary Consult
8/26/2019	Veterinary Consult
9/16/2019	Veterinary Consult

Scott Madill	
4/19	Quarterly Inspections
5/19	Quarterly Inspections
6/6/19Semi-annual	No Deficiencies
7/19	Quarterly Inspections
8/19	Quarterly Inspections
9/19/19	No vet consult conducted

Deborah Ferrington	
4/19	Quarterly Inspections
5/19	Quarterly Inspections
6/11/19	Veterinary Consult
7/19	Quarterly Inspections
8/19	Quarterly Inspections
/24/19Semi-annual	No Deficiencies

David Masopust		
4/22/19	Veterinary Consult	
5/20/19Semi-annual	No Deficiencies	
6/17/19	Veterinary Consult	
7/15/19	Veterinary Consult	
8/26/19	Veterinary Consult	
9/16/19	Veterinary Consult	

Mark Masino	
4/19	Quarterly Inspections
5/16/19	Veterinary Consult
6/19	Quarterly Inspections
7/19	Quarterly Inspections
8/15/19Semi-annual	No Deficiencies
9/19	Quarterly Inspections

Mark Sanders	
4/19	No housing for month of April so no visit conducted
5/7/19Semi-annual	No Deficiencies
6/26/19	Veterinary Consult
7/12/19	Veterinary Consult
8/27/19	Veterinary Consult
9/5/19	Veterinary Consult

Anna Tischler	
4/22/19	Veterinary Consult
5/20/19Semi-annual	No Deficiencies
6/17/19	Veterinary Consult
7/15/19	Veterinary Consult
8/26/19	Veterinary Consult
9/16/19	Veterinary Consult

Philip Portoghese	
4/30/19Semi- annual/Second Surgery	No Deficiencies
5/19	Quarterly Inspections
6/19	Quarterly Inspections
7/31/19	Veterinary Consult
8/19	Quarterly Inspections
9/19	Quarterly Inspections

Frank Ondrey	and the second second
4/19	Quarterly inspections
5/19	Quarterly inspections
6/5/19	Veterinary Consult
7/19	Quarterly inspections
8/19	Quarterly inspections
9/10/19PAM/Semi-annual	No Deficiencies

Michael Benneyworth	
4/19	No animals housed for month of April so no visit conducted
5/19	No animals housed for month of May so no visit conducted
6/19	No animals housed for month of June so no inspection conducted
7/19	No animals housed for month of July so no visit conducted
8/7/19Semi-annual	No Deficiencies
9/26/19	Veterinary Consult

Melanie Graham	
4/3/2019	Veterinary Consult
6/21/19PAM/Semi- annual	No Deficiencies
7/3/2019	Veterinary Consult
8/6/2019	Veterinary Consult

11/18	PI has closed laboratory, no further housin
11/18	PI has closed laboratory, no further h

Patrick Rothwell	
4/24/2019	Veterinary Consult
5/23/2019	Veterinary Consult
6/26/2019	Veterinary Consult
7/31/2019	Veterinary Consult
8/29/2019	Veterinary Consult
	Minor: new housing chambers for home cage activity experiments that last 24 hours. Will need to do microbiological testing
9/30/19PAM/Semi-	Minor: container of food not labeled with mill/fil dates
annual	Minor: staff dumping cage bedding without proper PPE
	Minor: ROHP requirements not met by all staff listed on protocol

Mark Thomas	
4/3/19 and 4/17/19	Veterinary Consult
5/15/19 and 5/22/19	Veterinary Consult
6/19/2019	Veterinary Consult
7/17/2019	Veterinary Consult
8/21/2019	Veterinary Consult
9/17/19Second Surgery/Semi- annual	Minor: ROHP requirements not met by all staff listed on protocol
	Minor: mice undergoing targeting surgeries are euthanaized prior to recovery. Euthanasia method used not approved in protocol

sther Krook-Magnuson	
4/15/19	Veterinary Consultation
5/19	No InspectionReduced Frequency
6/26/19	Veterinary Consultation
7/19	No InspectionReduced Frequency
8/14/19 and 8/20/19 PAM/Semi-annual	No Deficiencies
9/19	No InspectionReduced Frequency

Richard Bianco	3
4/10/19	Veterinary Consult
5/19/19	No InspectionReduced Frequency
6/6/19	Veterinary Consult
7/24/19Semi-annual	No Deficiencies
8/19	No Inspection-Reduced Frequency
9/11/19	Veterinary Consult

Gufa Lin	
4/19	Quarterly Inspections
5/19	Quarterly Inspections
6/19/19Second Surgery/Semi- annual	Minor: autoclaved material not dated
	Minor: needs to have a barrier in between animal and cage bedding
	Minor: uses distilled water for injection
	Minor: needs to store working anesthetic/analgesic solutions in sterile container
	Minor: needs to add two extra days of post operative records for frog/axolotI surgical sheets
7/19	Quarterly Inspections
8/19	Quarterly Inspections
9/19	No animals housed for month of September so no vet consult conducted

Sivaraj Sivamakrishnan	
4/19	Quarterly inspections
5/2/19PAM/Semi-annual	No Deficiencies
6/19	Quarterly inspections
7/19	Quarterly inspections
9/16	Veterinary Consultation

John Bischof	
4/19	Quarterly Inspection
5/19	Quarterly Inspection
6/24/19PAM/Semi-annual	No Deficiencies
7/19	Quarterly Inspection
8/19	Quarterly Inspection
9/19	No Veterinary consult done as no fish housed in space

Tim Kurtti	
4/19	No InspectionReduced Frequency
5/28	Veterinary consult
6/19	No InspectionReduced Frequency
7/19	No Consult done
8/19	No Consult done
9/20/19PAM/Semi- annual	Minor: ROHP requirements not met by all staff listed on protocol

Tay Netoff		1
4/16/2019	Veterinary Consult	-
5/13/2019	Veterinary Consult	- 4
6/27/2019	Veterinary Consult	
7/3/19 and 7/29/19	Veterinary Consult	

Bartolomucci	
4/12/2019	Veterinary Consult
5/10/2019	Veterinary Consult
6/7/2019	Veterinary Consult
7/3/2019	Veterinary Consult
	Minor: tape used in to secure cage cards to cage card holders
8/6/19Semi-annual	Minor: cylindrical housing cages fro when rats are tethered for infusion experiments are hand washed without microbiological testing
	Minor: should find different mechanism of securing black bag to caging as tape is currently used
9/27/2019	Veterinary Consult

Brendan Dougherty	
4/19	No visit as PI has not housed animals for month of April
5/19	No visit as PI has not housed animals for month of May

	Minor: flurogold tracer surgically injected into rat and then animal euthanized but tracer injection not approved in protocol		
8/22/19PAM/Semi- annual	Minor: non-pharmaceutical grade glue used on skull of rats		
	Minor: no procedure details for special caging where animals are continuously tethered for seizure monitoring		
	Minor: when animals are not housed in the lab, the rodent trap should be removed from the floor		
9/25/2019	Veterinary Consult		

Keith Vossel	
6/19	No visit as PI has not housed animals for month of June
7/22/2019	Veterinary Consult
8/7/2019	Veterinary Consult
9/18/2019	Veterinary Consult

	Minor: needs one extra day of Carprofen administration
7/19	No animals housed for month of July so no veterinary consult done
8/19	No animals housed for month of August so no veterinary consult done
9/19/2019	Veterinary Consult

Matthew Aliota	
7/2/2019	Veterinary Consult
8/27/2019	Veterinary Consult
9/30/19Semi-annual	No Deficiencies

Timothy Ebner/Clark Chen	
7/26/2019	Veterinary Consult
8/19/2019	Veterinary Consult
9/26/19Second	Minor: ROHP requirements not met by all staff listed on protocol
Surgery/Semi-annual	Minor: vaporizer overdue for calibration

## Fall 2019 IACUC Inspection Report

Inspection Type	Number	Percentage
# of PAM Inspections	117	41%
# Second Surgery Inspections	36	13%
# of Initial Surgery Inspections	6	2%
#PAM/Semi-Annual	28	10%
#Second Surgery/Semi-annual	8	3%
#Initial Surgery/Semi-annual	1	0%
# of Semi-Annual Inspections	74	26%
# of Ag Inspections	16	6%
Total # of Inspections	286	100%
Total # of Findings	201	
terreture production of another	March True	
Inspection Finding Summary	Number	Percentage
Finding Categor	У	-
IACUC (% of total findings)	178	89%
DEHS (% of total findings)	0	0%
DEHS-CS (% of total findings)	0	0%
IBC (% of total findings)	0	0%
OHS (% of total findings)	16	8%
Ag (% of total findings)	7	3%
Type of Inspectio	n	
PAM (% of total findings)	100	50%
Second Surgery Inspection (% of total findings)	18	9%
Initial Surgery Inspection (% of total findings)	5	2%
Initial Surgery/Semi-annual (% of total findings)	٥	0%
Semi-annual (% of total findings)	20	10%
PAM/Semi-annual (% of total findings)	27	13%
Second Surgery/Semi-annual (% of total findings)	13	6%
Ag (% of total findings)	7	3%
Self Report (% total findings)	10	5%
Surgical Records Review (% of total findings)	Ó	0%
Committee Request (% of total findings)		0%
Unannounced visit (% of total findings)	0	0%
Outside reports of non-compliance (% of total	1	0%



\* Buildings where animals are housed or used as part of AAALAC Accredited Unit

Type of Finding			
Minor (% of total findings)Standard	160	80%	
Minor (% of total findings)Other	6	3%	
Significant ( % of total findings)Standard	30	15%	
Significant ( % of total findings)Other	5	2%	

Repeat Findings					
Fall 2019 Spring 201					
Minor> Minor:	2	5			
Significant> Significant:	3	1			
Total # of repeat findings	5	6			

## Fall 2019 IACUC Inspection Report

SIGNIFICANT Fall 2019			
Findings	Number	Percentage	
Euthanasia methods not	1	3%	
Analgesics not given after surgical or anesthetic procedures as outlined in protocol**	5	14%	
Analgesics not given (time/duration) as outlined in protocol**	8	23%	
Anesthetic not used when performing procedures but protocol indicates anesthetic will be used	o	0%	
Cats used as part of teaching protocol but species not approved	2	3%	
blood collection conducted but procedure not approved in protocol*	ĩ	3%	
Lidocaine/bupivacaine not given prior to or after surgery	O	0%	
Personnel working with animals but not listed as staff on study	1	3%	
Expired anesthetic/ analgesic/euthanasia solution in use	1	3%	
anesthetic procedures conducted but not approved in protocol**	5	14%	
uses left over teaching quail as live prey for raptors	i	3%	
Housing of animlas outside of RAR without approval	4	11%	

SIGNIFICANT Spring 2019				
Findings	Number	Percentage		
Euthanasia methods not	-1	3%		
Analgesics not given after surgical or anesthetic procedures as outlined in protocol	7	24%		
Analgesics not given (time/duration) as outlined in protocol**	2	7%		
Anesthetic not used when performing procedures but protocol indicates anesthetic will be used	1	3%		
Sheep that received sedation did not have vital monitoring performed every 15 minutes but only pre-sedation measurements were obtained	1	3%		
Animal work conducted without an approved protocol	2	7%		
Lidocaine/bupivacaine not given prior to or after surgery	2	7%		
Personnel working with animals but not listed as staff on study	1	3%		
Expired anesthetic/ analgesic/euthanasia solution in use	1	3%		
Body temperature and oxygen saturation levels low during surgical procedures without any intervention	1	3%		
Cage space requirements for pigs not followed overcrowding in nursery	1	3%		
three NHP underwent CT scans for anatomical imaging but CT scans not approved in protocol*	1	3%		

Survival surgery conducted but procedures not approved on protocol	4	11%	Survival surgery conducte on protocol that does no have survival procedures approved (animals ordere on wrong protocol)
Care not given to animals that were housed in IMHA for up to six days	1	3%	mice underwent up to thr cardiotoxin injury surgics procedures but protocol d not outline multiple injur surgeries in same anima
One NHP received less than the minimum water allotment due to transcription error	1	3%	Animals left unattended RAR while recovering fro anesthesia
controlled substances not properly stored	Q	0%	controlled substances no properly stored
CO2 tank unsecured	1.	3%	Staff euthanizing two cag of mice with CO2 left then euthanasia chamber unattended
Total Significant Findings	35	100%	due to miscommunication the weekend treatment sheet, two sheep did no receive antibiotics; last do of Carprofen or vitals. N pain seen on weekend b vets

\*Reported to OLAW

Survival surgery conducted on protocol that does not have survival procedures as approved (animals ordered on wrong protocol)	2	7%
mice underwent up to three cardiotoxin injury surgical procedures but protocol does not outline multiple injury surgeries in same animal	1	3%
Animals left unattended in RAR while recovering from anesthesia	1	3%
controlled substances not properly stored	1	3%
Staff euthanizing two cages of mice with CO2 left them in euthanasia chamber unattended	ì	3%
due to miscommunication on the weekend treatment sheet, two sheep did not receive antibiotics; last dose of Carprofen or vitals. No pain seen on weekend by vets	1	3%
Report details mouse found on floor of research area that was lethargic and underwent some type of surgical intervention as apparent in surgical repair of chest, animal not properly euthanized by cervical dislocation	1	3%
Total Significant Findings	29	100%

MINOR Fall 2019FindingsNumberPercentageROHP138%IPNFExpired items42%IPNFSurgical Records85%PNF4326%PNF-Other00%IPNF-Other00%IPNF-Other00%DEHS00%DEHS00%IPNF-Anesthetic Records85%AgOther00%IPNF-Personnel Training Records117%OHS32%14IPNF-Aseptic Technique140%					
Findings	Number	Percentage			
ROHP	13	8%			
IPNFExpired items	4	2%			
IPNF-Surgical Records	8	5%	IP		
PNF	43	26%			
PNF-Other	0	0%			
IPNFStandard	43	26%	-		
IPNF-Other	0	0%	11		
DEHS	Ó	0%	11		
DEHS-CS	0	0%	1		
IPNF-Anesthetic Records	8	5%	IPN		
AgStandard	6	4%	) (E		
AgOther	0	0%	1		
IPNF-Personnel Training Records	11	7%	IPN		
онѕ	3	2%	2.		
IPNF-Aseptic Technique	13	8%	IPN		
Husbandry	14	8%	1		
Total Minor Findings	166	100%	Т		

Ň	IINOR Spring 2019	
Findings	Number	Percentage
ROHP	11	7%
IPNFExpired items	1	1%
IPNF-Surgical Records	7	4%
PNF	49	30%
PNF-Other	1	1%
IPNFStandard	37	22%
IPNF-Other	0	0%
DEHS	1	1%
DEHS-CS	0	0%
PNF-Anesthetic Records	11	7%
AgStandard	15	9%
AgOther	D	0%
IPNF-Personnel Training Records	7	4%
OHS	1	1%
IPNF-Aseptic Technique	18	11%
Husbandry	6	4%
Total Minor Findings	165	100%

\*IPNF: IACUC Policy Not followed \*PNF: Protocol Not Followed

	Fall	Spring	Fall								
	2014	2015	2015	2016	2016	2017	2017	2018	2018	2019	2019
Minors	54	57	63	51	99	66	75	76	97	165	166
Significants	42	57	59	38	45	27	29	34	23	29	35





Obtained by Rise for Animals. Uploaded to Animal Research Laboratory Overview (ARLO) on 04/21/2021

	Fall 2014	Spring 2015	Fall 2015	Spring 2016	Fall 2016	Spring 2017	Fall 2017	Spring 2018	Fall 2018	Spring 2019	Fall 2019
Inspection Findings	353	634	548	454	474	341	387	292	120	184	192
Findings-Other	6	15	108	10	30	12	11	7	6	10	9



	Fall 2014	Spring 2015	Fall 2015	Spring 2016	Fall 2016	Spring 2017	Fall 2017	Spring 2018	Fall 2018	Spring 2019	Fall 2019
Total Findings-Standard	353	634	548	454	474	341	387	292	114	184	201
Total Inspections	317	322	331	331	273	266	275	278	293	287	286
No Findings	177	126	153	171	100	119	121	143	213	190	192
At least one finding	140	196	178	160	173	147	154	135	80	97	94



	Spring 2015	Fall 2015	Spring 2016	Fall 2016	Spring 2017	Fall 2017	Spring 2018	Fall 2018	Spring 2019	Fall 2019
Total Inspections	322	331	331	273	266	275	278	293	287	286
First Notices	528	445	372	353	260	294	189	0	0	0
Minors	57	61	50	94	65	75	76	95	164	160
Significants	49	42	32	27	16	29	27	19	21	30
	Spring 2015	Fall 2015	Spring 2016	Fall 2016	Spring 2017	Fall 2017	Spring 2018	Fall 2018	Spring 2019	Fall 2019
First Notices	1.6	1.3	1.1	1.3	1.0	1.1	0.7	0.0	0.0	0.0
Minors	0.2	0.2	0.2	0.3	0.2	0.3	0.3	0.3	0.6	0.6
Significants	0.2	0.1	0.1	0.1	0.1	0.1	0.1	0.1	0.1	0.1



Significants

0.2

0.1

0.1

0.1

0.1

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	Spring	5 Eall 2015	Spring	E-11 2016	Spring	Call 2017	Spring		Spring		
	2015	Fail 2015	2016		2017	Fail 2017	2018	Fall 2018	2019	Fall 2019	
First Notices	83.28%	81.20%	81.94%	74.47%	76.25%	73.87%	64.73%	0.00%	0.00%	0.00%	
Minors	8.99%	11.13%	11.01%	19.83%	19.06%	18.84%	26.03%	83.33%	88.65%	84.21%	
Significants	7.73%	7.66%	7.05%	5.70%	4.69%	7.29%	9.25%	16.67%	11.35%	15.79%	



Fall 2019 Significant Findings Report											
Type of Inspection	Building	Room #	Responsible Party	Date of Inspection	Deficiency Type-DEHS, OHS, etc.	Deficiency * = reported to OLAW ** = repeat finding	Corrective Action	Completion Date	Inspectors		
PAM			Christine Sivula	4/9/2019	IACUC	Last dose of Carprofen not given 12-24 hours as outlined in protocol	implementation or procedures so that when there are multiple early morning weekend treatments scheduled, the vet tech and on call veterinarian will confer on Friday to triage the cases for the weekend, ensuring critical treatments are completed in a timely	4/11/2019	Kristin Pilon		
Self Report		***	Yi-Mei Yang	4/23/2019	IACUC	surgery performed on mice that is not approved in the protocol	cranial window implantation done to check that brain was not damaged by the viral infusion over time. This surgery will not be done further until an amendment is submitted and approved	4/23/2019	Self Report		
Second Surgery Inspection			Demetri Yannopoulos	4/25/2019	IACUC	buprenorphine not given to seven pigs undergoing CPR procedures as outlined in protocol	buprenorphine will be given as outlined in study	4/26/2019	Kristin Pilon		
PAM			Stephanie Goldschmidt	5/27/2019	IACUC	cats used as part of teaching protocol but species not approved	shelter was asked to only bring dogs, but occassionally brought cats and PI mistakenly thought protocol had approval for both; a new shelter is being used for animals and they will only supply dogs	5/29/2019	Kristin Pilon		
Second Surgery Inspection			Samuel Dudley	5/29/2019	IACUC	SR buprenorphine not given two hours prior to surgery as outlined in protocol	PI will communicate with Pilar Guzman regarding the timing of SR-Bup prior to transmitter surgery so that analgesic will be given at	6/4/2019	Paul Lindstrom		
РАМ			Jizhen Lin	5/31/2019	IACUC	bioluminescence imaging conducted on mice but anesthetic event not approved in protocol	further imaging will not be conducted until protocol has been amended to include IVIS imaging	6/3/2019	Paul Lindstrom		

Fall 2019 Significant Findings Report										
Type of Inspection	Building	Room #	Responsible Party	Date of Inspection	Deficiency Type-DEHS, OHS, etc.	Deficiency * = reported to OLAW ** = repeat finding	Corrective Action	Completion Date	Inspectors	
Self Report			David Masopust	6/10/2019	IACUC	personnel working with animals not listed as staff on protocol	person has been added to protocol	6/10/2019	Self Report	
PAM			Michelle Willette	6/11/2019	IACUC	houses quail in raptor center but protocol not approved for housing	protocol will be renewed prior to next quail lab (September 2019) and IMHA information will be added at that time (no quail will be housed or used until protocol is renewed)	6/12/2019	Kristin Pilon	
PAM		- ( <b>1</b>	Michelle Willette	6/11/2019	IACUC	uses left over teaching quail as live prey for raptors	renewal protocol will be updated to include use of live quail as prey for raptors; no quail will be obtained or utilized until this change is made in renewal	6/12/2019	Kristin Pilon	
PAM			Michael Farrar	6/11/19 and 6/25/19	IACUC	blood collection done but not approved on protocol*	amendment submitted for blood collection	6/25/2019	llana Cohen	
PAM/Semi-annual			John Bishof	6/24/2019	IACUC	mice undergoing thermal therapty did not receive analgesics as outlined in the protocol (repeat finding)	amendment submitted making it clear that the thermal treatment is not a surgery and removing analgesics; RAR contacted for help in how to monitor animals after this procedure	7/9/2019	Paul Lindstrom and Georgiy Aslanidi	
PAM/Semi-annual			John Bishof	6/24/2019	IACUC	mice euthanized by cervical dislocation without anesthetics but protocol states anesthetics will be used	amendment submitted requesting cervical dislocation without the use of anesthesia; will not be performed until amendment approved	7/9/2019	Paul Lindstrom and Georgiy Aslanidi	
Ag		***	Kyle Rozeboom	6/25/2019	Ag	CO2 tank not secured	chained immediately by staff	6/25/2019	Paul Lindstrom and Scott Madill	
Second urgery/Semi-annual			Richard Bianco	6/27/2019	IACUC	eight sheep upderwent an additional balloon dilation surgical procedure that is not outlined in the protocol	amendment submitted fo additional dilatation procedure	6/28/2019	Kristin Pilon and Scott Madill	

Fall 2019 Significant Findings Report										
Type of Inspection	Building	Room #	Responsible Party	Date of Inspection	Deficiency Type-DEHS, OHS, etc.	Deficiency * = reported to OLAW ** = repeat finding	Corrective Action	Completion Date	Inspectors	
Outside Report of Non-Compliance			Walter Low	6/20/2019	IACUC	surgical procedures conducted on protocol that were not approved	staff will double check protocol prior to ordering animals to ensure proper protocol used	7/2/2019	Jennifer Borgert	
Second Surgery Inspection			Andrew Grande	6/20/2019	IACUC	surgical procedure conducted on protocol that is not outlined in study	amendment submitted updating protocol with second surgical procedure	7/15/2019	Jennifer Borgert	
PAM			Eric Newman	6/26/19 and 6/28/19	IACUC	anaglesics not given after electrolytic lesion surgery*	analgesics will be administered per protocol	7/1/2019	Megan McCoy	
PAM			Michael Smanski	7/18/2019	IACUC	anesthetic procedure not approved in protocol	amendment will be submitted adding anesthetic procedure; procedure will not be performed until approved	7/19/2019	Paul Lindstrom	
PAM			Clark Chen	7/11/19 and 7/16/19	IACUC	housing animals without approval	amendment submitted requesting IMHA housing; animals will not be housed until amendment approved	7/19/2019	Jennifer Borgert	
PAM			Nicola Grissom	7/24/2019	IACUC	Second dose of carprofen given upwards of 29-30 hours after initial dose	going forward, lab will ensure that mice have administration of carprofen within a span of 24 hours or less	7/26/2019	Kristin Pilon	
Self Report			Walter Low	7/22/2019	IACUC	animals housed in laboratory over 24 hours without approval	will amend the protocol for housing of animals over 24 hours in lab to acclimate rats to behavioral tests	7/22/2019	Self Report	
PAM			Aaron LeBeau	8/13/2019	IACUC	housing of animals without approval in protocol	protocol will be updated with IMHA information	8/21/2019	llana Cohen	
PAM			Aaron LeBeau	8/13/2019	IACUC	care not given to animals that were housed in IMHA for up to six days	going forward, lab will perform all animal care when IMHA is in use	8/21/2019	Ilana Cohen	
РАМ			Aaron LeBeau	8/13/2019	IACUC	bioluminescence imaging conducted but no longer approved on protocol as Pl removed this procedure through an amendment	imaging procedure was accidently removed from protocol; amendment will be submitted adding procedure back	8/21/2019	llana Cohen	

Fall 2019 Significant Findings Report										
Type of Inspection	Building	Room #	Responsible Party	Date of Inspection	Deficiency Type-DEHS, OHS, etc.	Deficiency * = reported to OLAW ** = repeat finding	Corrective Action	Completion Date	Inspectors	
Initial Surgery Inspection			John Osborn	8/13/2019	IACUC	mice that wereundergoing non-survival surgery did not receive the pre-operative dose of Meloxicam as stated in the protocol (repeat finding)	amendment submitted requesting removal of meloxicam from the acute procedure	8/15/2019	Paul Lindstrom	
ΡΑΜ			Phu Tran	8/20/2019	IACUC	SR buprenorphine not given for carotid artery ligation surgeries as outlined in protocol (repeat finding)	going forward, SR buprenorphine will be given as outlined in protocol	8/27/2019	Ilana Cohen	
PAM			Swati More	8/20/2019	IACUC	expired Xylazine in use	new xylazine purchased	8/22/2019	Paul Lindstrom	
РАМ			Kaylee Schwertfeger	8/23/2019	IACUC	animals underwent retro- orbital IV adminsitration of BM while under Isoflurane but anesthetic procedure not approved in protocol*	amendment submitted clarifying that bone marrow will be adminitered tail vein or retro-orbital vein, and retro-orbital administration will be done under isoflurane	8/30/2019	Megan McCoy	
РАМ			Sarah Greising	8/27/2019	IACUC	SR buprenorphine not given at time that as outlined in protocol*	going forward, SR- Buprenorphine will be given 2-4 hours prior to surgery as outlined in the protocol	9/3/2019	Kristin Pilon	
РАМ			Sarah Greising	8/27/2019	IACUC	animals undergo anesthetic procedure to remove tissue prior to euthanasia that is not outlined in protocol*	amendments submitted adding procedure to protocols	9/3/2019	Kristin Pilon	
Second Surgery Inspection			R. Scott McIvor	8/27/2019	IACUC	analgesic not given per timing that is approved in protocol*	going forward, all staff will administer analgesic according to timing approved in protocol	9/6/2019	Megan McCoy	

Fall 2019 Significant Findings Report									
Type of Inspection	PAM Building Room #		Responsible Party	Date of Inspection	Deficiency Type-DEHS, OHS, etc.	Deficiency * = reported to OLAW ** = repeat finding	Corrective Action	Completion Date	Inspectors
PAM			Yingjie Chen	9/4/2019	IACUC	SR buprenoprhine not given 2 3 hours prior to surgery	giving SR-Buprenorphine before the surgery negatively affected the survival rate; thus an amendment will be submitted amending the timing of analgesic administration; no surgery will be performed until amendment has been submitted and approved	9/5/2019	Jennifer Borgert
РАМ			Michael Benneyworth	9/16/2019	IACUC	anaglesic given upwards of 29 30 hours aftter initial dose, thus resulting in a time frame where pain was not managed	lab amended SOP such that analgesics will now be administered in the morning making sure the time is less than 24 hours between doses	9/19/2019	Kristin Pilon
PAM			Linda McLoon	9/24/2019	IACUC	SR buprenoprhine not given 2 4 hours prior to surgery as outlined in protocol	Going forward, SR- Buprenorphine will be given at least 2 hours prior to surgery	9/30/2019	Jennifer Borgert
Self Report		***		9/16/2019	IACUC	one NHP received less than the miniumum water allotment due to transcription error	water allotment corrected and vet contacted to assess animal	9/30/2019	Self Report

## Fall 2019 Minor Findings Report

Type of Inspection	Building	Room #	Responsible Party	Date of Inspection	Deficiency Type- DEHS, QHS, etc.	Minor Deficiency * = Repeat Finding	Corrective Action	Completion Date	inspectors	Category
Ag			Alfredo DiCostanzo	5/10/2019	Ag	Disaster plan overdue for review	disaster plan updated	5/13/2019	Paul Lindstrom	Ag
Ag			Alfredo DiCostanzo	5/10/2019	Ag	some entries for medical issues did not have resolution	animal care records updated	5/13/2019	Paul Lindstrom	Ag.
Ag		***	Dan Braaten	5/30/2019	Ag	hay bails used for feeding stored next to random equipment and gas can	hay bales had been moved during spring cleaning and have now been moved back to their normal storage location at the opposite end of the barn; the gas can has also been moved to the storage barn	6/4/2019	Kristin Pilon and Dan Montonye	Ag
Ag		*##	Wayne Martin	6/21/2019	Ag	feed storage area in St. Paul Poultry Facilities had bags of chicken feed stored on a pallet but pushed up against the wall	feed pallets moved away from the wall	7/16/2019	Jennifer Borgert and Jen Hubbard	Ág

## Fall 2019 Minor Findings Report

Type of Inspection	Building	Room #	Responsible Party	Date of Inspection	Deficiency Type- DEHS, OHS, etc.	Minor Deficiency * = Repeat Finding	Corrective Action	Completion Date	Inspectors	Category
Ag			Wayne Martin	6/21/2019	Ag	disaster plan overdue for review	disaster plan reviewed and cover sheet updated	7/26/2019	Jennifer Borgert and Jen Hubbard	Ag
Ag		***	Wayne Martin	6/21/2019	Ag	hutch doors are not locked during the day causing them to be slightly ajar; to prevent escapes, please secure the doors in the day too	chicken hutch doors will now be locked during the day	7/16/2019	Jennifer Borgert and Jen Hubbard	Ag
Semi-annual				4/10/2019	IACUC	baggies of soaked primate biscuits were observed that were not labeled with the contents or date of preparation	A system for labeling soaked biscuits, including contents and date prepared, has been put in place	4/29/2019	Melanie Graham and Brian Crooker	Husbandry
Semi-annual			Jennifer Erickson	6/25/2019	IACUC	mill/fill dates either missing or not clearly marked on food containers	mill/fill dates will now be recorded on a laminated card placed in a cage card holder to prevent writing from being smudged	7/3/2019	Kristin Pilon and Geoff Ghose	Husbandry
Type of Inspection	Building	Room #	Responsible Party	Date of Inspection	Deficiency Type- DEHS, OHS, etc.	Minor Deficiency * = Repeat Finding	Corrective Action	Completion Date	Inspectors	Category
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Semi-annual			Jennifer Erickson	6/25/2019	IACUC	needs to clean broom in 1-118	upon further inspection broom not dirty but worn from repeated use	7/14/2019	Kristin Pilon and Geoff Ghose	Husbandry
Semi-annual			Przemysław Bajer	7/26/2019	IACUC	containers of fish food in refrigerator had no expiration or fill dates	fish food containers have been labeled	8/15/2019	Paul Lindstrom and Mimie Pollard	Husbandry
Sem i-annual		ě.	Nicholas Phelps	7/26/2019	ACUC	containers of fish food in refrigerator had no expiration or fill dates	containers are now clearly labeled	8/14/2019	Paul Lindstrom and Mimie Pollard	Husbandry
Semi-annual			Emilyn Alejandro	8/6/2019	MOUC	tape used to secure cage cards to caging	staff will now use regular, pre-ordered RAR cage cards which should fit into the cage card holders	8/26/2019	Kristin Pilon and Laura Hocum Stone	Husbandry

Type of Inspection	Building	Room #	Responsible Party	Date of Inspection	Deficiency Type- DEHS, OHS, etc.	Minor Deficiency * = RepentFinding	Corrective Action	Completion Date	Inspectors	Category
Semi-annual			John Osborn	8/6/2019	IACUC	microbiological testing not done on caging equipment that is washed by staff	going forward either RAR will clean the cages or microbial testing will be done	8/8/2019	Kristin Pilon and Laura Hocum Stone	Husbandry
Semi-annual			John Osborn	8/6/2019	IACUC	tape used to secure black bag to outside of cage which leaves residue	a different method will be used to hold bags on in the future	8/8/2019	Kristin Pilon and Laura Hocum Stone	Husbandry
PAM/Semi-annual			Marija Cventanovic	8/19/2019	ACUC	temperature and humidity out of range in housing area so gave permission to house in temp/humidity controlled building. Needs to add this space as part of IMHA	am endment submitted adding alternative IMHA location	9/4/2019	Kristin Pilon and	Husbandry
PAM/Semi-annual		Buildíng	Marija Cventanovic	8/19/2019	IACUC:	rodent control not in place	live traps will be purchased, baited and placed in rooms during housing of any species	9/4/2019	Kristin Pilon and	Husbandry

Type of Inspection	Building	Room #	Responsible Party	Date of Inspection	Deficiency Type- DEHS, OHS, etc.	Minor Deficiency * = Repeat Finding	Corrective Action	Completion Date	Inspactors	Category
PAM/Semi-annual		Building <b>an an</b>	Marija Cventanovic	8/19/2019	ACUC	food not labeled with mill/fill dates	containers with animal food will be labeled with mill date of the food	9/4/1949	Kristin Pilon and	Husbandry
PAM/Semi-annual			Tay Netoff	8/22/2019	IACUC	rodent trap should be picked up when animals are not housed as it was not monitored	rodent traps are being placed in a bag in the cabinet when animals are not housed	9/4/2019	Kristin Pilon and Peggy Norris	Husbandry
Semi-annual			Jennifer Erickson	9/26/2019	IACUC	one of the Tence rails in paddock is loose and should be repaired	FM contacted and loose fence rail repaired on 9/27/19	9/27/2019	llana Cohen and Sally Noll	Husbandry
PAM/Semi-annual			Patrick Rothwell	9/30/2019	IACUC	microbiological testing needs to be done on new home cage chambers	Jessica Rugg will be contacted to obtain sample kits and IMHA section will be updated to include these chambers and disinfection process	10/8/2019	Kristin Pilon and Liz Pluhar	Husbandry

Type of Inspection	Building	Room #	Responsible Party	Date of Inspection	Deficiency Type- DEHS, OHS, etc.	Minor Deficiency * = Repeat Finding	Corrective Action	Completion Date	Inspectors	Category
PAM		_	Keith Vossel	4/15/2019	ACUC	sutures not removed from animals after surgery	going forward, absorbable sutures will be used and/or sutures will be removed in 7-14 days	4/22/2019	Megan McCoy	IPNF
Semi-annual			Zigang Dong	5/15/2019	IACUC	food and water had been pulled for stress induction but no time was noted on the caging or in the room as to when this food/water has been pulled so that staff could determine length of deprivation	going forward, time of food/water removal will be noted on the whiteboard in the housing room	5/20/2019	Kristin Pilon and Felicia Boynton	IPNF
Second Surgery Inspection			Sade Spencer	5/14/2019	ACUC	decapitation log not kept up to date	decapitation log has been updated using sacrifice records; machine shop will be used to sharpen guillotine blade as needed and log maintained	5/21/2019	Kristin Pilon	IPNF
PAM		1	lames Ervasti	5/22/2019	IACUC	SOP not available for the cleaning/disinfection of behavioral equipment	cleaning SOP drafted and staff trained on procedure	5/30/2019	Megan McCoy	IPNE
PAM		-	Ann Parr	5/29/2019	IACUC	anesthetic vaporizer overdue for inspection	vaporizer has been inspected	6/4/2019	Megan McCoy	IPNF

Type of Inspection	Building	Room #	Responsible Party	Date of Inspection	Deficiency Type- DEHS, OHS, etc.	Minor Deficiency * = Repent Finding	Corrective Action	Completion Date	Inspectors	Category
PAM/Semi- annual			Jennifer Menken	6/17/2019	IACUC	disaster plan overdue for review	disaster plan has been updated	7/13/2019	Megan McCoy and Marilyn Bennett	IPNF
Second Surgery/Semi- annual			Gufa Lin	6/19/2019	IACUC	barrier not used in between animal and cage bedding when recovering them from anesthesia	autoclaved paper towels will be placed between the animal and cage bedding	6/20/2019	Kristin Pilon and Sara Hashway	IPNF
Second Surgery/Semi- annual		5	Gufa Lin	6/19/2019	IACUC	uses distilled water for injection	injectable PBS will be used to dilute	6/20/2019	Kristin Pilon and Sara Hashway	IPNF
Second Surgery/Semi- annual			Gufa Lin	6/19/2019	IACUC	sterile container not used for storing diluted solutions used in animals	drugs will be stored in recommended containers	6/20/2019	Kristin Pilon and Sara Hashway	IPNF
PAM/Semi- annual		4	John Bishof	6/24/2019	IACUC	disaster plan overdue for review	disaster plan has been updated	6/26/2019	Paul Lindstrom and Georgiy Aslanidi	IPNF

Type of Inspection	Building	Room #	Responsible Party	Date of Inspection	Deficiency Type- DEHS, QHS, etc.	Minor Deficiency * = Repeat Finding	Corrective Action	Completion Date	Inspactors	Category
PAM/Semi- annual		I	Brendan Dougherty	6/20/2019	IACUC	disaster plan overdue for review	disaster plan reviewed and cover sheet updated	7/10/2019	Kristin Pilon and Henry Wong	IPNE
PAM/Semi- annual			Brendan Dougherty	6/20/2019	IACUC	Carprofen only given 2 days post-op instead of required 3	post-operative records updated to include another day of Carprofen administration	7/10/2019	Kristin Pilon and Henry Wong	IPNF
PAM			James Lokensgard	6/25/2019	IACUC	breeding colony records not kept	breeding spreadsheet created	6/28/2019	llana Cohen	IPNF
PAM			James Lokensgard	6/25/2019	IACUC	log not kept of maintenance and usage of decapitation equipment	decapitation log created	6/28/2019	llana Cohen	IPNF
PAM		-	James Lokensgard	6/25/2019	IACUC	SOP not available for cleaning/disinfection of behavioral equipment	behavioral equipment cleaning SOP created	6/28/2019	llana Cohen	IPNf

Type of Inspection	Building	Room #	Responsible Party	Date of Inspection	Deficiency Type- DEHS, OHS, etc.	Minor Deficiency * = Repeat Finding	Corrective Action	Completion Date	inspectors	Category
Semi-annual			Jennifer Erickson	6/25/2019	IACUC	treatment cards do not have monitoring plan listed so cannot tell how long or how often animals are monitored	According to RAR SOP rodents being monitored require rounds once per week; the inspection occurred on a Tuesday leaving the rest of the week for rounds to be completed; if more or less frequent checks are required (differing from standard SOP), that would be indicated on the vet plan card	7/14/2019	Kristin Pilon and Geoff Ghose	IPNF
Semi-annual		-	Dawn Lowe	6/25/2019	IACUC	single housing of female mice without exception in place	amendment submitted for both protocols adding justification for single housing	7/11/2019	Kristin Pilon and Geoff Ghose	IPNF
Self Report			Ned Patterson	6/28/2018	IACUC	lapse in monitoring documentation for a dog that developed vestibular disease	PI will ensure consistent and appropriate documentation; will inform Dr. Boynton of any medical issues that come up and plan will be documented in the book in the doo	6/28/2019	Self Report	IPNF

ype of inspection	Building	Room #	Responsible Party	Date of Inspection	Deficiency Type- DEHS, OHS, etc.	Minor Deficiency * = Repeat Finding	Corrective Action	Completion Date	Inspectors	Category
PAM		-	Xavier Revelo	7/8/2019	IACUC	non-pharmaceutical grade glucose used	pharmaceutical grade glucose will be used	7/16/2019	Kristin Pilon	IPNF
PAM			Xavier Revelo	7/8/2019	IACUC	SOP not in place for cleaning/disinfection of behavioral equipment	SOP is now in place of cleaning/disinfection of behavioral equipment; all staff have been trained	7/16/2019	Kristin Pilon	IPNF
рам			Jesse Williams	7/10/2019	IACUC	No flow meter on CO2 used for euthanasia	Ed Craig contacted to install and calibrate new flow meter; new custom lid designed for chamber and signage posted	7/25/2019	Kristin Pilon	IPNF
Semi-annual			Marc Schwabenlander	7/18/2019	IACUC	non-pharmaceutical grade xylazine used	pharmaceutical grade xylazine ordered from vendor and will be used going forward	8/24/2019	Jennifer Borgert and Scott Madill	IPNE

Type of Inspection	Building	Room #	Responsible Party	Date of Inspection	Deficiency Type- DEHS, OHS, etc.	Minor Deficiency * = Repent Finding	Corrective Action	Completion Date	Inspectors	Category
Semi-annual				7/23/2019	IACUC	some wooden ramps/pieces for the old animal chair/restraint systems were full of mold and smelled of urine and should be disposed	all wood items associated with the NHP chair immobilization equipment will be disposed; the plastic platforms may be used again, but the corroded feet will be removed and discarded	7/25/2019	Paul Lindstrom and Melanie Graham	IPNE
Second Surgery/Semi- annual			Ben Hayden	7/24/2019	IACUC	paint used in open field cage is peeling in areas of stress (door entry) and on items animals come into contact (barrels)	door repaired and will be monitoring for peeling paint; new barrels have been ordered to replace those with chipped paint	8/8/2019	Kristin Pilon and Melanie Graham	IPNF
PAM/Semi-annual			Mārija Cventanovic	8/19/2019	IACUC	Styrofoam container used as euthanasia chamber	non-porous plastic container will be purchased for use as euthanasia chamber	9/4/2019	Kristin Pilon and	IPNE
PAM/Semi-annual			Marija Cventanovic	8/19/2019	IACUC	housing space did not have disaster plan, IMHA information, or how to report animal concern signage	disaster plan, IMHA information and information on how to report animal welfare concerns will be posted	9/4/1949	Kristin Pilon and	IPNF

Type of inspection	Building	Room #	Responsible Party	Date of Inspection	Deficiency Type- DEHS, OHS, etc.	Minor Deficiency * = Repeat Finding	Corrective Action	Completion Date	Inspectors	Category
AM/Semi-annual			Suzanne McGaugh	8/16/2019	IACUC	disaster plan overdue for review	disaster plan has been reviewed	8/27/2019	Jennifer Borgert and Marilyn Bennett	IPNE
'AM/Semi-annual		-	Mark Desrosiers	8/20/2019	ACUC	staff could not access protocol and protocol not in lab	protocol has been printed and placed in binder	9/5/2019	Jennifer Borgert and Keith Barker	IPNF
AM/Semi-annual			Tay Netoff	8/22/2019	IACUC	non-pharmaceutical grade glue used to cement electrode to skull	pharmývet grade 3M Vetbond will be used going forward	9/4/2019	Kristin Pilon and Peggy Norris	IPNF
РАМ			Şarah Greising	8/27/2019	IACUC.	vaporizer overdue for calibration	appointment made; calibration will be done on 9/4/19	9/3/2019	Kristin Pilon	IPNF
PAM			Sarah Greising	8/27/2019	IACUC	needs cleaning/disinfection SOP for behavioral equipment	lab now has dedicated SOP for cleaning behavioral cages; SOP posted in animal housing room and in protocol binder	9/3/2019	Kristin Pilon	IPNF

Type of Inspection	Building	Room #	Responsible Party	Date of Inspection	Deficiency Type- DEHS, OHS, etc.	Minor Deficiency * = Repeat Finding	Corrective Action	Completion Date	Inspectors	Category
Semi-annual			Melissa Palmer	8/22/2019	IACUC	pieces of information missing from housing area	emergency plan and emergency contact information will be posted; animal reporting info will also be posted	9/12/2019	Kristin Pilon and Peggy Norris	IPNF
PAM			Erik Finger	8/29/2019	IACUC	vaporizer overdue for calibration	calibration service scheduled	9/17/2019	Kristin Pilon	IPNF
PAM		-	Atsushi Asakura	8/30/2019	ACUC	SOP not available for cleaning and disinfection of behavioral equipment	SOP has been created	9/5/2019	Jennifer Borgert	IPNF
PAM			Hongbo Pang	9/9/2019	ACUC	pH of Avertin not taken prior to injection	going forward, pH will be measured prior to injection	9/17/2019	Kristin Pilon	IPNF
PAM			Robert Cormier	9/18/2019	IACUC	non-pharmaceutical grade Carprofen used	pharmaceutical grade carprofen has been ordered and will be used going forward	9/19/2019	Paul Lindstrom	IPNF

ype of inspection	Building	Room #	Responsible Party	Date of Inspection	Deficiency Type- DEHS, OHS, etc.	Minor Deficiency * = Reposit Finding	Corrective Action	Completion Date	Inspectors	Category
Semi-annual			Thomas Hrabik	9/18/2019	IACUC	disäster plan overdue for review	disaster plan reviewed and new cover sheet attached	9/27/2019	Kristin Pilon and Felicia Boynton	IPNE
PAM		-	Linda McLoon	9/24/2019	IACUC	SOP not available for cleaning/disinfection of behavioral equipment	SOP will be posted in behavior area and staff trained	9/30/2019	Jennifer Borgert	IPNF
Second Surgery/Semi- annual			Tim othy Ebner	9/26/2019	IACUC	vaporizer overdue for inspection	vaporizer calibration scheduled; will not be used until calibration complete	9/27/2019	Paul Lindstrom and Christine Sivula	IPNE
AM/Semi-annual			Patrick Rothwell	9/30/2019	IACUC	food container in IMHA not labeled with mill/fill date	food container discarded; going forward all food containers will be Tabeled appropriately	10/8/2019	Kristin Pilon and Liz Pluhar	IPNE

Type of Inspection	Building:	Room #	Responsible Party	Date of Inspection	Deficiency Type- DEHS, QHS, etc.	Minor Deficiency * = Repeat Finding	Corrective Action	Completion Date	Inspectors	Category
PAM			Lucy Vulchanova	9/27/2019	IACUC	eye ointment not used for SNI surgeries	eye ointment will be used	10/13/2019	Ilana Cohen	IPNF
Initial Surgery Inspection			Harry Orr	6/12/2019	IACUC	sutures not removed following ICV injection surgery	amendment submitted to add anesthetic procedure to remove sutures	6/20/2019	Ilana Cohen	IPNF
РАМ			Clark Chen	7/11/19 and 7/16/19	IACUC	person performing surgery not listed as surgeon on protocol	amendment submitted adding surgeon	7/19/2019	Jennifer Borgert	IPNF
РАМ			Jizhen Lin	5/31/2019	IACUC	records not found for anesthetic procedures conducted on animals	anesthetic events are recorded, but records have not been kept; going forward, records will be kept and added to binder for storage	6/3/2019	Paul Lindstrom	IPNF-Anesthetic Records
РАМ		1	James Lokensgard	6/25/2019	IACUC	anesthetic record not kept for imaging procedures	anesthetic records kept	6/28/2019	llana Cohen	IPNF-Anesthetic Records

Type of Inspection	Building	Room #	Responsible Party	Date of Inspection	Deficiency Type- DEHS, OHS, etc.	Minor Deficiency * = RepentFinding	Corrective Action	Completion Date	(inspectors	Category
PAM		-	Clark Chen	7/11/19 and 7/16/19	IACUC	anesthetic records not kept for im aging procedures	going forward, anesthetic records will be kept	7/18/2019	Jennifer Borgert	IPNF-Anesthetic Records
PAM		:•0	Zohar Sachs	8/13/2019	ACUC	anesthetic records not kept	going forward, anesthetic records will be kept using the template provided	8/19/2019	Paul Lindstrom	IPNF-Anesthetic Records
PAM/Semi-annual			Marija Cventanovic	8/19/2019	ACUC	no anesthetic records in place	going forward, anesthetic records will be kept using the template provided	9/4/1949	Kristin Pilon and	IPNF-Anesthetic Records
PAM		-	Dale Gregerson	9/5/2019	IACUC	anesthetic records not kept	anesthetic records will be kept using the form provided	9/12/2019	Ilana Cohen	IPNF-Anesthetic Records
РАМ		-	Yingjie Chen	9/4/2019	IACUC	anesthetic records not kept	records will be kept in the lab going forward	9/5/2019	Jennifer Borgert	IPNF-Anesthetic Records

Type of inspection	Building	Room #	Responsible Party	Date of Inspection	Deficiency Type- DEHS, OHS, etc.	Minor Deficiency * = RepeatFinding	Corrective Action	Completion Date	Inspectors	Category
PAM			Linda McLoon	9/24/2019	IACUC	anesthetic records not kept	anesthetic records will be kept going forward	9/30/2019	Jennifer Borger t	IPNF-Anesthetic Records
PAM		***	Yibin Deng	5/15/2019	IACUC	animals not shaved prior to survival surgery procedure	in the future, hair will be removed from incision site by shaving or depilatory cream	5/28/2019	Paul Lindstrom	IPNF-Aseptic Technique
Second Surgery/Semi- annual			Gufa Lin	6/19/2019	IACUC	autociaved material not labeled as to sterilization date	date will be recorded on the autoclaved materials	6/20/2019	Kristin Pilon and Sara Hashway	IPNF-Aseptic Technique
Second Surgery/Semi- annual			Richard Bianco	6/27/2019	IACUC	crack in floor of OR that needs to be filled	floor to be repaired by 11/15/19	10/14/2019	Kristin Pilon and Scott Madill	IPNF-Aseptic Technique
РАМ			Paolo Provenzano	6/27/2019	IACUC	surgical mask and hair covering not worn during survival surgeries	mask and hair covering will be worn for survival surgery	7/1/2019	Jennifer Borgert	IPNF-Aseptic Technique

Type of inspection	Building	Room #	Responsible Party	Date of Inspection	Deficiency Type- DEHS, OHS, etc.	Minor Deficiency * = Repeat Finding	Corrective Action	Completion Date	inspectors	Category
Self Report			Yibin Deng	7/8/2019	IACUC	Tur not shaved from animals prior to Umplantsfor of tumor (repeat finding)	fur will be removed going forward	7/8/2019	Self Report	IPNF-Aseptic Technique
PAM			Nicola Grissom	7/24/2019	IACUC	surgical instruments not completely sterilized prior to use	all surgical equipment will be sterilized prior to surgery	7/26/2019	Kristin Pilon	IPNF-Aseptic Technique
PAM			Nicola Grissom	7/24/2019	IACUC	hair covering not worn during surgical procedures	hair covering will used during surgeries	7/26/2019	Kristin Pilon	IPNF-Aseptic Technique
PAM			Kaylee Schwertfeger	8/23/2019	IACUC	face mask not worn during surgical procedures	going forward, a face mask will be worn for surgical procedures	8/30/2019	Megan McCoy	IPNF-Aseptic Technique
PAM			Yingjie Chen	9/4/2019	IACUC	face mask not worn during surgical procedures	going forward, face mask will be used	9/5/2019	Jennifer Borgert	IPNF-Aseptic Technique

Type of Inspection	Building	Room #	Responsible Party	Date of Inspection	Deficiency Type- DEHS, OHS, etc.	Minor Deficiency * = Repeat Finding	Corrective Action	Completion Date	Inspectors	Category
PAM			Robert Cormier	9/18/2019	IACUC	hair covering not worn during surgical procedures	a hair net will be used going forward	9/19/2019	Paul Lindstrom	IPNF-Aseptic Technique
PAM		:•0	Linda McLoon	9/24/2019	IACUC	hair covering not worn during surgical procedures	going forward, hair covering will be worn	9/30/2019	Jennifer Borgert	IPNF:Aseptic Technique
РАМ			Linda McLoon	9/24/2019	IACUC	surgical instruments not completely sterilized prior to surgery	all înstruments will autoclave sterilized going forward	9/30/2019	Jennifer Borgert	IPNF-Aseptic Technique
РАМ			Lucy Vulchanova	9/27/2019	IACUC	sterile gloves expired	new sterile gloves have been purchased	10/13/2019	Ilana Cohen	IPNF-Aseptic Technique
Semi-annual		1	Jennifer Erickson	4/23/2019	IACUC	expired Accel cleaner in	expired solution discarded and replaced with solution prepared and dated on 4/23/19	5/1/2019	llana Cohen and Laura Hocum- Stone	IPNF-Expired Items

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Type of Inspection	Building	Room #	Responsible Party	Date of Inspection	Deficiency Type- DEHS, OHS, etc.	Minor Deficiency * = Repeat Finding	Corrective Action	Completion Date	Inspectors	Category
Semi-annual			Marc Schwabenlander	7/18/2019	IACUC	expired betadine scrub pad in exposure kit	new betadine scrub pads have been placed in the NHP kit	8/24/2019	Jennifer Borgert and Scott Madill	IPNF-Expired Items
Semi-annual			Michael Smanski	7/26/2019	IACUC	expired vials of Ovaprim	expired vials will be discarded and replaced with new	7/29/2019	Paul Lindstrom and Mimie Pollard	IPNF-Expired Items
РАМ		_	Masato Yamamoto	7/22/2019	IACUC	surgical gloves expired but in use	expired gloves disposed and will be replaced	8/1/2019	Jennifer Borgert	IPNF-Expired Items
PAM		***	Yibin Deng	5/15/2019	IACUC	no personnel training records for newest members of laboratory working with animals	training records will be completed for new lab staff members	5/28/2019	Paul Lindstrom	IPNF-Personnel Training Records
Second Surgery Inspection			Jakub Tolar	6/12/2019	IACUC	No training records for new employee	Training records for Courtney Popp added to binder	6/14/2019	Jennifer Borgert	IPNF-Personnel Training Records

Type of Inspection B	uilding	Room #	Responsible Party	Date of Inspection	Deficiency Type- DEHS, OHS, etc.	Minor Deficiency * = Repeat Finding	Corrective Action	Completion Date	Inspectors	Category
Second Surgery Inspection			Andrew Grande	6/20/2019	IACUC	training record for surgeon not available	training record submitted	6/27/2019	Jennifer Borgert	IPNF-Personnel Training Records
PAM			Michael Smanski	7/18/2019	IACUĆ	personnel training record detailing protocol specific procedures not available	records will be kept using template	7/19/2019	Paul Lindstrom	IPNF-Personnel Training Records
Second Surgery Inspection			David Bereiter	7/23/2019	IACUC	procedure training record missing for new employee	training documented and filed for new employee	8/13/2019	Ilana Cohen	IPNF-Personnel Training Records
PAM			Aaron LeBeau	8/13/2019	IACUC	no personnel training records kept for any of the new staff in the laboratory	training template filled out for each member of the lab and records now stored in the lab	8/21/2019	Ilana Cohen	IPNF-Personnel Training Records
PAM			Christopher Pennell	8/16/2019	IACUC	limited personnel training records	training records completed using template provided; records available in lab	9/12/2019	Megan McCoy	IPNF-Personnel Training Records

Type of inspection	Building.	Room #	Responsible Party	Date of Inspection	Deficiency Type- DEHS, OHS, etc.	Minor Deficiency * = Repeat Finding	Corrective Action	Completion Date	Inspectors	Category
PAM/Semi-annual		-	Mark Desrosiers	8/20/2019	IACUC	personnel training records not present	training documentation placed in binder	9/5/2019	Jenifer Borgert and Keith Barker	IPNF-Personnel Training Records
PAM			Kakambi Nagaraja	8/21/2019	IACUC	personnel training records not (nesent (recoar finding)	training records will be kept using template provided	9/11/2019	Kristin Pilon	IPNF-Personnel Training Records
PAM		-	Hongbo Pang	9/9/2019	IACUC	Personnel training records not kept	training records created and uploaded to protocol	9/17/2019	Kristin Pilon	IPNF-Personnel Training Records
PAM			Lucy Vulchanova	9/27/2019	IACUC	personnel training records missing for new staff member	training records for new staff have been created; going forward new staff training records will be created immediately	10/13/2019	Ilana Cohen	IPNF-Personnel Training Records
Second Surgery Inspection			Wei Chen	5/23/2019	IACUC	noted that p.o records kept on cage cards for rats undergoing survival surgery but cards not permanently kept	Post op records will now be kept	5/29/2019	Jennifer Borgert	IPNF-Surgical Records

Type of inspection	Building	Room #	Responsible Party	Date of Inspection	Deficiency Type- DEHS, QHS, etc.	Minor Deficiency * = Repent Finding	Corrective Action	Completion Date	Inspectors	Category
Second Surgery/Semi- annual		-	Gufa Lin	6/19/2019	IACUC	only keeps one day of post operative monitoring for frog/axolotl surgeries	three days of post- operative records will be kept	6/20/2019	Kristin Pilon and Sara Hashway	IPNF-Surgical Records
РАМ			Eric Newman	6/26/19 and 6/28/19	IACUC	post operative records not kept for last batch of surgeries	post operative records will be kept	7/1/2019	Megan McCoy	IPNF-Surgical Records
PAM		-	Clark Chen	7/11/19 and 7/16/19	IACUC	no post-operative records kept for surgical procedures	going forward, post- op records will be kept for all surgeries	7/18/2019	Jennifer Borgert	IPNF-Surgical Records
Initial Surgery Inspection		-	Jop Van Berlo	7/22/2019	ACUC	three days of post operative records not always kept	PI will retrain lab staff regarding proper record keeping procedures to ensure full compliance	8/2/2019	Kristin Pilon	JPNF-Surgical Records

Type of inspection	Building	Room #	Responsible Party	Date of Inspection	Deficiency Type- DEHS, OHS, etc.	Minor Deficiency * = Repeat Finding	Corrective Action	Completion Date	Inspectors	Category
Initial Surgery Inspection			Jop Van Berlo	7/22/2019	ACUC	myocardial infarction surgeries performed on 6/25 did not indicate analgesics were given as no documentation in surgical or controlled substance records	PI will retrain lab staff regarding proper record keeping procedures to ensure full compliance	8/2/2019	Kristin Pilon	IPNF-Surgical Records
Second Surgery Inspection			Emilyn Alejandro	9/6/2019	IACUC	needs to add the use of Lidocaine to surgical procedures	amendment submited adding lidocaine to protocol	10/3/2019	Megan McCoy	IPNF-Surgical Records
РАМ		-	Lucy Vulchanova	9/27/2019	IACUC	no surgical records for non-survival mouse spinal cord exposure surgeries	going forward, surgical records will be kept for all non- survival surgeries	10/13/2019	Ilana Cohen	IPNF-Surgical Records
PAM/Semi-annual		_	Marija Cventanovic	8/19/2019	OHS	personnel dumping cage bedding without proper PPE	personnel dumping cage bedding will contact Respiratory protection program	9/7/2019	Kristin Pilon and	OHS

Type of Inspection	Building	Room #	Responsible Party	Date of Inspection	Deficiency Type- DEHS, OHS, etc.	Minor Deficiency * = Repeat Finding	Corrective Action	Completion Date	Inspectors	Category
PAM		1	Sarah Greising	8/27/2019	OHS	staff dump bedding from physical/metabolic caging	staff performing this function are now enrolled in the Respiratory Protection Program	9/3/2019	Kristin Pilon	OHS
PAM/Semi-annual			Patrick Rothwell	9/30/2019	OHS	staff dumping bedding without proper PPE	Staff will dump bedding in a fume hood; if not deemed sufficient, RPP will be contacted for further evaluation	10/8/2019	Kristin Pilon and Liz Pluhar	OHS
Self Report			Jizhen Lin	4/18/2019	ACUC	endpoints of study not followed	lab is to consult with area veterinarian to view the animals and confirm that the tumor load is not beyond endpoint criteria	4/18/2019	Self Report	PNF
PAM/Semi-annual			Gail Boe	4/29/2019	IACUC	fish are sometimes adopted out but endpoint not approved	amendment submitted adding adoption as an endpoint for the fish	5/9/2019	Kristin Pilon and Christine Sivula	PNF
PAM			David Masopust	4/26/2019	IACUC	Avertin not approved for use for one procedure in protocol	amendment submitted adding avertin	5/7/2019	Megan McCoy	PNF

Type of inspection	Building	Room #	Responsible Party	Date of Inspection	Deficiency Type- DEHS, QHS, etc.	Minor Deficiency * = Repeat Finding	Corrective Action	Completion Date	(inspectors	Category
Second Surgery Inspection			Tim Starr	5/1/2019	IACUC	needs to give one more day of analgesic coverage; considered day 1 day of surgery instead of day 0	one additional dose of ketoprofen will be given to cover 3 full days	5/2/2019	llana Cohen	PNF
Second Surgery Inspection			Tim Starr	5/1/2019	ACUC	staff conducting surgeries but not listed as surgeon on protocol	amendment submitted adding surgeon	5/2/2019	llana Cohen	PNF
Self Report			Mark Herzberg	4/16/2019	ACUC	euthanasia method not followed as per approved protocol	amendment submitted to add perfusion as an approved method of euthanasia for future experiments	4/16/2019	Self Report	PNF
Second Surgery Inspection			Sade Spencer	5/14/2019	IACUC	xylocaine used as topical anesthetic for intracranial surgeries but not outlined for procedures	amendment submitted adding xylocaine	5/21/2019	Kristin Pilon	PNF
РАМ			Stephanie Goldschmidt	5/27/2019	ACUC	anesthetics used on dogs but not approved on protocol	protocol will be amended to include all anesthetics used	5/29/2019	Kristin Pilon	PNF

Type of inspection	Building	Room #	Responsible Party	Date of Inspection	Deficiency Type- DEHS, OHS, etc.	Minor Deficiency * = Repeat Finding	Corrective Action	Completion Date	Inspectors	Category
Second Surgery Inspection			Samuel Dudley	5/29/2019	IACUIC	person performing surgery not listed as surgeon in protocol	amendment submitted to add surgeon	6/27/2019	Paul Lindstrom	PNF
PAM		l	lizhen Lin	5/31/2019	MCUC	uses colon cancer /ovarian cells but only approved for melanoma and breast cancer cells	cell fines will be added to protocol; no work on cell fines will be done until am endment approved	6/3/2019	Paul Lindstrom	PNF
РАМ			Tyler Bold	6/17/2019	IACUC	endpoints of study not followed	amendment submitted extending endpoints	7/11/2019	Kristin Pilon	PNF
PAM		-	Walter Low	6/18/2019	IACUC	mice receive 1 million cells but only approved for 100,000; cells given 2-4 days later but protocol reads that they should be given 24-48 hours later	amendment submitted to increase number of cells and route of administration	7/1/2019	Paul Lindstrom	PNF
РАМ		—	Walter Low	6/18/2019	IACUC	behavioral tests conducted that are not listed in the protocol	amendment submitted to add additional behavioral tests	7/1/2019	Paul Lindstrom	PNF

Type of Inspection	Building	Room #	Responsible Party	Date of Inspection	Deficiency Type- DEHS, OHS, etc.	Minor Deficiency * = Repeat Finding	Corrective Action	Completion Date	Inspectors	Category
PAM		ļ	Alonso Guedes	6/21/2019	IACUC	monitoring description in protocol for animals undergoing surgery is not current practice	amendment submitted updating monitoring description	7/15/2019	Kristin Pilon	PNF
PAM		-	Steven Graves	6/25/2019	IACUC	person performing surgery not listed as a surgeon in the protocol	amendment submitted to add surgeon to protocol	7 <i>/2/2</i> 019	Paul Lindstrom	PNF
Second Surgery/Semi- annual			Richard Bianco	6/27/2019	IACUC	needs to update protocol to reflect that the two animals only undergoing Echo procedure every other week are not implanted	amendment submitted to clarify procedures for control animals	6/28/2019	Kristin Pilon and Scott Madill	PNF
PAM			Eric Newman	6/26/19 and 6/28/19	IACUC	person performing surgery not listed as a surgeon in the protocol	surgeon will be added to protocol	7/1/2019	Megan McCoy	PNF
РАМ		-	Xavier Revelo	7/8/2019	IACUC	fecal transplant study conducted but study not outlined in protocol	lab will no longer perform fecal transplant studies; if needed in the future, an amendment will be submitted to add procedure to protocol	7/16/2019	Kristin Pilon	PNF

Type of Inspection	Building	Room #	Responsible Party	Date of Inspection	Deficiency Type- DEHS, OHS, etc.	Minor Deficiency * = Repent Finding	Corrective Action	Completion Date	finspaceors	Category
PAM		-	Clark Chen	7/11/19 and 7/16/19	IACUC	Isoflurane anesthetic used for perfusions but Urethane/Acepromazin e approved in protocol for this procedure	amendment submitted requesting use of isoflurane for perfusions	7/19/2019	Jennifer Borgert	PNF
рам			Guisheng Song	7/22/2019	IACUC	adult mice receiving AAV via tail injection but protocol indicates that 2 day old mice will be injected via AAV via temporal vein	amendment submitted updating protocol to include adult mice	8/9/2019	Paul Lindstrom	PNF
РАМ			Nicola Grissom	7/24/2019	IACUC	dosage of ketamine used for experimental procedures prior to operant behavioral testing is over twice the dosage approved	amendment submitted updated ketamine dosage	7/26/2019	Kristin Pilon	PNF
РАМ			Nicola Grissom	7/24/2019	IACUC	needs to add lidocaine for use on ear bars prior to surgery	amendment submitted adding the use of lidocaine	7/26/2019	Kristin Pilon	PNF

Type of Inspection	Building	Room #	Responsible Party	Date of Inspection	Deficiency Type- DEHS, OHS, etc.	Minor Deficiency * = Repeat Finding	Corrective Action	Completion Date	Inspectors	Category
PAM			Michael Kyba	7/30/2019	NCUC	bilateral surgical procedure done to injure the muscle/cell transplant but protocol only reads that one hind limb muscle will be injured and transplanted	amendment submitted clarifying that the procedure is bilateral	8/2/2019	Megan McCoy	PNF
Second Surgery Inspection			Marija Cvetanovic	7/29/2019	IACUC	needs to reinstate giving SR buprenorphine 3-4 hours prior to surgery as per stipulations; Carprofen was on board at time of surgery (repeat finding)	Going forward, SR- Buprenorphine will be given 3-4 hours prior to surgery	8/15/2019	Jennifer Borgert	PNF
PAM		***	Sara Hamilton-Hart	8/6/2019	IACUC	endpoints of study not followed	amendment submitted extending study endpoints	8/15/2019	Kristin Pilon	PNF
PAM		_	Stephen Jameson	8/6/2019	IACUC	anesthetic used for intranasal infection not approved for this procedure	amendment submitted adding additional anesthetic	8/9/2019	Ilana Cohen	PNF

Type of Inspection	Building	Room #	Responsible Party	Date of Inspection	Deficiency Type- DEHS, OHS, etc.	Minor Deficiency * = Repent Finding	Corrective Action	Completion Date	Inspectors	Category
PAM			Stephen Jameson	8/6/2019	IACUC	euthanasia method for parabiosis study different than what is approved in protocol	amendment submitted adding alternative euthanasia method	8/9/2019	Ilana Cohen	PNF
PAM		-	Alexander Khoruts	8/15/2019	ACUC	monitoring of hamsters between 36-48 hours after infection not done as per approved protocol	amendment submitted updating and describing hamster monitoring	9/9/2019	Kristin Pilon	PNF
РАМ			PhuTran	8/20/2019	IACUC	euthanasia method used not outlined in protocol	amendment submitted adding euthanasia method to protocol	8/27/2019	llana Cohen	PNF
Self Réport			Felicia Boynton	8/20/2010	IACUC	ophthalmology procedures conducted on training protocol that were not outlined	amendment has been submitted to add them to training protocol	8/20/2019	Self Report	PNF
РАМ			Kakambi Nagaraja	8/21/2019	IACUC	blood collection method different than approved in protocol	protocol will be amended to include alternate route of blood collection	9/11/2019	Kristin Pilon	PNF

Type of inspection	Building	Room #	Responsible Party	Date of Inspection	Deficiency Type- DEHS, OHS, etc.	Minor Deficiency * = Repent Finding	Corrective Action	Completion Date	Inspectors	Category
PAM			Kakambi Nagaraja	8/21/2019	IACUC.	rabbits immunized weekly and with increasing volumes of antigen/adjuvant for a total of 8 weeks	since rabbits were no seroconverting with low antigen concentration, the antigen concentration was increased; this experiment is now complete but if the protocol is renewed, it will be amended with the increased concentrations	9/11/2019	Kristin Pilon	PNF
PAM/Semi-annual		-	Tay Netoff	8/22/2019	IACUC.	fluorogold tracer injection conducted but not approved in protocol	amendment submitted adding procedure to protocol; no fluorogold injections will be performed until amendment is approved	9/4/2019	Kristin Pilon and Peggy Norris	PNF
PAM/Semi-annual		÷	Tay Netoff	8/22/2019	IACUC	needs to update protocol to detail tethering of animals in special caging for one week	amendment submitted updating tethering procedures and length of time	9/4/2019	Kristin Pilon and Peggy Norris	PNF

Type of Inspection	Building	Room #	Responsible Party	Date of Inspection	Deficiency Type- DEHS, OHS, etc.	Minor Deficiency * = Repeat Finding	Corrective Action	Completion Date	Inspectors	Category
PAM			Sarah Greising	8/27/2019	IACUC	injections into muscle given as treatment do not require a surgery and thus do not require analgesics, needs to update protocols to make this distinction	amendment submitted clarifying that injections into muscles given as treatment is not a surgical procedure and therefore analgesics are not given	9/3/2019	Kristin Pilon	PNF
PAM			Sarah Greising	8/27/2019	IACUC.	protocol 1811-36513A should be updated to reflect that time 0 is a non-survival surgery not requiring analgesics	amendment submitted making clarification on time 0 not requiring analgesics	9/3/2019	Kristin Pilon	PNF
РАМ		-	Dale Gregerson	9/5/2019	IACUC	fundus imaging done under Isoflurane instead of ketämine/xylazine as approved in the protocol	amendment submitted adding alternate anesthetic for this procedure	9/12/2019	Ilana Cohen	PNF
рам			Hongbo Pang	9/9/2019	IACUC	five million cells injected SC but protocol only approved for one million tumor cells to be given	amendment submitted increasing range of number of cells to be injected	9/17/2019	Kristin Pilon	PNF

Type of Inspection	Building	Room #	Responsible Party	Date of Inspection	Deficiency Type- DEHS, OHS, etc.	Minor Deficiency * = Repeat Finding	Corrective Action	Completion Date	Inspectors	Category
РАМ		-	Hongbo Pang	9/9/2019	IACUC	Avertin used prior to perfusion but ketamine/xylazine listed as anesthetic to be used	amendment submitted adding avertin option	9/17/2019	Kristin Pilon	PNF
PAM			Robert Cormier	9/18/2019	IACUC	person performing surgery not listed as a surgeon	amendment submitted adding surgeon to protocol	9/19/2019	PaulLindstrom	PNF
Second Surgery/Semi- annual			Mark Thom as	9/17/2019	IACUC	euthanasia method not approved in protocol	amendment submitted updating euthanasia method	9/27/2019	Kristin Pilon and Jen Hubbard	PNF
Second Surgery Inspection			Jean Règal	9/18/2019	ACUC	needs to have an exception for single housing of animals that do not have catheters	amendment submitted updating social housing exception to include animals that do not have catheters	9/27/2019	Jennifer Borgert	PNF
Self Report			Vaiva Vezys	9/23/2019	IACUC	animal work done on wrong protocol as wrong animals transferred to for experiments	proper transfer paperwork put in and discrepancy resolved	9/24/2019	Self Report	PNF

Type of Inspection	Building	Room #	Responsible Party	Date of Inspection	Deficiency Type- DEHS, OHS, etc.	Minor Deficiency * = Repeat Finding	Corrective Action	Completion Date	fineplatero rs	Category
Second Surgery Inspection			Demetri Yannopoulos	4/25/2019	OHS	ROHP requirements not met by all staff listed on protocol	all staff now ROHP compliant	4/29/2019	Kristin Pilon	ROHP
Initial Surgery Inspection			Kevin Wickman	4/30/2019	оня	ROHP requirements not met by all staff listed on protocol	all staff now ROHP compliant	5/1/2019	Jennifer Borgert	ROHP
PAM			Alex Bianco	5/3/2019	OHS	ROHP requirements not met by all stafflisted on protocol	all staff now ROHP compliant	5/24/2019	Kristin Pilon	ROHP
PAM			Paolo Provenzano	6/27/2019	онз	ROHP requirements not met by all staff listed on protocol	ROHP requirements are being met	7/1/2019	Jennifer Borgert	ROHP
Second Surgery Inspection		-	Shalom Michaeli	8/6/2019	OHS.	ROHP requirements not met by all stafflisted on protocol	All staff now ROHP compliant	8/19/2019	Jennifer Borgert	ROHP

Type of Inspection	Building.	Room #	Responsible Party	Date of Inspection	Deficiency Type- DEHS, OHS, etc.	Minor Deficiency * = Repent Finding	Corrective Action	Completion Date	Inspectors	Category
PAM		-	George Wilcox	8/20/2019	онѕ	ROHP requirements not met by all staff listed on protocol	person removed from study	8/21/2019	Paul Lindstrom	ROHP
PAM			Robert Cormier	9/18/2019	онş	ROHP requirements not met by all staff listed on protocol	all staff now ROHP compliant	9/19/2019	Paul Lindstrom	ROHP
Second Surgery/Semi- annual		_	Mark Thomas	9/17/2019	OHS	ROHP requirements not met by all stafflisted on protocol	all staff now ROHP compliant	9/27/2019	Kristin Pilon and Jen Hubbard	ROHP
Semi-annual			Thomas Hrabik	9/18/2019	OHS	ROHP requirements not met by all stafflisted on protocol	all staffnow ROHP compliant	10/11/2019	Kristin Pilon and Felicia Boynton	ROHP
PAM/Semi-annual			Ulrike Munderloh	9/20/2019	OHS	ROHP requirements not met by all stafflisted on protocol	Tetanus scheduled for 10/16	9/26/2019	Jennifer Borgert and Keith Baker	ROHP

Type of inspection	Building	Room #	Responsible Party	Date of Inspection	Deficiency Type- DEHS, OHS, etc.	Minor Deficiency * = Repeat Finding	Corrective Action	Completion Date	Inspactors	Category
Šecond Surgery/Semi- annual			Tim othy Ebner	9/26/2019	онз	ROHP requirements not met by all stafflisted on protocol	ROHP requirement has been met	10/14/2019	Paul Lindstrom and Christine Sivula	ROHP
PAM/Semi-annual			Patrick Rothwell	9/30/2019	онѕ	ROHP requirements not met by all staff listed on protocol	staff member has an appointment to receive tetanus booster	10/8/2019	Kristin Pilon and Liz Pluhar	ROHP
PAM/Semi-annual		_	Jerrold Vitek and Matthew Johnson	9/27/2019	OHS	ROHP requirements not met by all staff listed on protocol	ROHP requirements are being met	10/1/2019	Kristin Pilon and Dan Montonye	ROHP

		Fall 2019 No Findings F	leport		
Type of Inspection	Building	Room #	Responsible Party	Date of Inspection	Inspectors
PAM	1		Ferenc Toth	4/9/2019	Kristin Pilon
PAM			Michael Garwood	4/10/2019	Paul Lindstrom
Semi-annual			Jennifer Erickson	4/10/2019	Melanie Graham and Brian Crooke
Semi-annual			Jennifer Erickson	4/10/2019	Melanie Graham and Brian Crooke
Semi-annual			Jeramy Kulesa	4/10/2019	Melanie Graham and Brian Crooke
PAM			A. David Redish	4/16/2019	Paul Lindstrom
PAM			Jayanth Panyam	4/16/2019	llana Cohen
Ag			Lee Johnston	4/17/2019	Paul Lindstrom an Felicia Boynton
Ag			Bradley Heins	4/17/2019	Paul Lindstrom an Felicia Boynton
Ag			David Israels-Swenson	4/17/2019	Paul Lindstrom an Felicia Boynton
PAM			Daniel Saltzman	4/18/2019	Paul Lindstrom
Semi-annual		Jennifer Erickson	4/19/2019	Megan McCoy and Don Martin	
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Semi-annual		Jennifer Erickson	4/23/2019	llana Cohen and Laura Hocum-Stone	
PAM		Manish Patel	4/24/2019	Paul Lindstrom	
Second Surgery Inspection		Robert Tranquillo	4/25/2019	Kristin Pilon	
Second Surgery Inspection		Ganesh Raveendran	4/25/2019	Kristin Pilon	
PAM		Andrew Grande	4/25/2019	Kristin Pilon	
PAM		Roy Cho	4/25/2019	Kristin Pilon	
Second Surgery Inspection		Derrick Green	4/25/2019	Kristin Pilon	
Semi-annual		Jennifer Erickson	4/25/2019	llana Cohen and Mimie Pollard	
Semi-annual	***	Tristan McNamara	4/25/2019	Ilana Cohen and Mimie Pollard	
PAM		Sunil David	4/26/2019	Jennifer Borgert	
Semi-annual		Jennifer Rees	4/25/2019	Jennifer Borgert and Christine Sivula	
Second Surgery Inspection	***	Sara Hashway	4/19/2019	Jennifer Borgert	

Semi-annual		Rob Denton	4/29/2019	Kristin Pilon and Christine Sivula
Semi-annual		Kerry Michael	4/29/2019	Kristin Pilon and Christine Sivula
Semi-annual		Heather Waye	4/29/2019	Kristin Pilon and Christine Sivula
Semi-annual		Heather Waye	4/29/2019	Kristin Pilon and Christine Sivula
Semi- annual/Second Surgery		Philip Portoghese	4/30/2019	Jennifer Borgert and Nate Koewler
PAM		Daniel Mueller	4/22/19 and 4/30/19	Jennifer Borgert
Initial Surgery Inspection		Kevin Wickman	4/30/2019	Jennifer Borgert
PAM		Prakash Kara	4/30/2019	Kristin Pilon
PAM		Eric Jensen	5/2/2019	Paul Lindstrom
PAM/Semi-annual	) <b></b> (	Sivaraj Sivaramakrishnan	5/2/2019	Jennifer Borgert and George Wilcox
Second Surgery Inspection		Greg Beilman	5/7/2019	Jennifer Borgert
PAM		Kirsten Nielsen	5/7/2019	Jennifer Borgert
PAM/Semi-annual		Gordon Smith	5/8/2019	Kristin Pilon and Peggy Norris

Semi-annual		Mark Sanders	5/7/2019	Paul Lindstrom and Liz Pluhar
Semi-annual		Mark Sanders	5/7/2019	Paul Lindstrom
Semi-annual	·	Alena Talkachova	5/8/2019	Paul Lindstrom and Don Martin
PAM		Tanya Freedman	5/8/2019	Ilana Cohen
Ag		Sally Noll	5/10/2019	Paul Lindstrom
PAM	***	Georgiy Aslanidi	5/15/2019	Paul Lindstrom
Second Surgery Inspection	***	Rebecca Morris	5/15/2019	Kristin Pilon
PAM		Beshay Zordoky	5/16/2019	llana Cohen
Second Surgery/Semi- annual Inspection		Carrie Haskell-Luevano	5/17/2019	Jennifer Borgert and Swayam Prabha
Semi-annual		Maxim Cheeran	5/20/2019	Paul Lindstrom and Sara Hashway
Semi-annual		Jennifer Erickson	5/22/2019	llana Cohen and George Wilcox
PAM		Mary Garry	5/21/2019	Paul Lindstrom
Second Surgery Inspection		Savita Rao	5/22/2019	Paul Lindstrom

Semi-annual	Jennifer Erickson	5/17/2019	Megan McCoy an Kakambi Nagaraj
Semi-annual	Jennifer Erickson	5/17/2019	Megan McCoy an Kakambi Nagaraj
Semi-annual	Ned Patterson	5/17/2019	Megan McCoy an Kakambi Nagaraj
Second Surgery Inspection	Sergio Gradilone	5/15/2019	Megan McCoy
PAM	Shujun Liu	5/15/2019	Megan McCoy
PAM	Andrew Nelson	5/28/2019	Kristin Pilon
PAM	Katie Satrom	5/28/2019	Megan McCoy
PAM	Ingunn Stromnes	5/24/2019	Jennifer Borgert
PAM	Geoffrey Hart	5/28/2019	Jennifer Borgert
PAM	Daniel Garry	5/31/2019	Paul Lindstrom
PAM	Gulin Oz	5/30/2019	Jennifer Borgert
Ag	Hugh Chester-Jones	5/31/2019	Jennifer Borgert
Ag	Samuel Baidoo	5/31/2019	Jennifer Borger

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Second Surgery Inspection		Mohammad Saleem Bhat	6/6/2019	Megan McCoy
Second Surgery Inspection		Timothy O'Connell	6/5/2019	Ilana Cohen
Second Surgery Inspection		Thomas Griffith	6/4/2019	llana Cohen
PAM		Dawn Lowe	6/4/2019	Paul Lindstrom
Semi-annual	***	Scott Madill	6/6/2019	Ben Clark and Sally Noll
PAM/Semi-annual		Christine Sivula	6/10/2019	Kristin Pilon and Peggy Norris
PAM	***	Paul Everson	6/13/2019	Jennifer Borgert
PAM		Daniel Vallera	6/13/2019	Paul Lindstrom
PAM	j 💻	David Largaespada	6/11/2019	Paul Lindstrom
РАМ		Yun You	6/17/2019	Jennifer Borgert
PAM		Anthony Baughn	6/20/2019	Jennifer Borgert
Semi-annual		Jennifer Erickson	6/21/2019	Paul Lindstrom and Brian Crooker
Semi-annual		Jennifer Erickson	6/24/2019	Kristin Pilon and George Aslanidi

PAM/Semi-annual		Melanie Graham	6/21/2019	Paul Lindstrom and Brian Crooker
PAM/Semi-annual		Melanie Graham	6/21/2019	Paul Lindstrom and Melanie Graham
PAM	- <b>m</b> 21	Lisa Peterson	6/19/2019	Megan McCoy
PAM		Kalpna Gupta	6/25/2019	Jennifer Borgert
Ag		Sally Noll	6/25/2019	Paul Lindstrom and Scott Madill
PAM/Semi-annual		Hubert Lim	6/25/2019	Paul Lindstrom and Nathan Koewler
PAM		Sergey Khasabov	6/26/2019	Paul Lindstrom
Second Surgery Inspection		Cindy Martin	6/27/2019	Kristin Pilon
PAM		Rosemary Kelly	6/27/2019	Kristin Pilon
PAM		Ratan Banik	6/26/2019	Paul Lindstrom
PAM		Rafael Andrade	6/27/2019	Kristin Pilon
Second Surgery Inspection		Julia Lemos	6/27/2019	Kristin Pilon
PAM		Peter Santi	6/26/2019	Megan McCoy

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Ag	***	Krishona Martinson	6/21/2019	Jennifer Borgert and Jen Hubbard
Ag	***	Brian Crooker	6/21/2019	Jennifer Borgert and Jen Hubbard
Semi-annual		Wei Chen	7/8/2019	Kristin Pilon and Jer Hubbard
Semi-annual		Christina Clarkson	7/8/2019	Kristin Pilon and Brian Crooker
Second Surgery Inspection		Joceyln Richard	7/11/2019	Kristin Pilon
PAM		Alfonso Araque	7/11/2019	Ilana Cohen
PAM		Stanley Thayer	7/17/2019	Paul Lindstrom
Semi-annual	***	Jennifer Erickson	7/18/2019	llana Cohen and Robert Schumache
Semi-annual	***	Jennifer Erickson	7/18/2019	llana Cohen and Robert Schumache
Semi-annual	***	Jennifer Erickson	7/18/2019	llana Cohen and Robert Schumache
PAM	-	Catherine Kotz	7/17/2019	Ilana Cohen
Second Surgery Inspection		Sven-Ulrik Gorr	7/12/2019	llana Cohen
Semi-annual		Jennifer Erickson	7/15/2019	Megan McCoy and Laura Hocum Stone

Semi-annual		Jennifer Erickson	7/15/2019	Megan McCoy and Laura Hocum Stone
Semi-annual		Dan Busian	7/19/2019	Paul Lindstrom and Sam Baidoo
PAM		Carston Wagner	7/22/2019	Megan McCoy
Second Surgery/Semi- annual		Paul laizzo	7/22/2019	Paul Lindstrom and Scott Madill
Initial Surgery Inspection		Yoji Shimizu	7/24/2019	llana Cohen
Semi-annual	***	Richard Bianco	7/24/2019	Paul Lindstrom and Brian Crooker
PAM	***	Scott Madill	7/19/2019	Jennifer Borgert
PAM	***	Alon Herschhorn	7/25/2019	Kristin Pilon
Second Surgery Inspection		Alik Widge	7/25/2019	Kristin Pilon
Initial Surgery/Semi- annual	- <b>-</b> 1	John Ward	7/26/2019	Paul Lindstrom and Mimie Pollard
PAM		Ivan Tkac	7/29/2019	Paul Lindstrom
PAM		Subree Subramanian	7/23/2019	Megan McCoy
PAM		Cheuk Leung	7/29/2019	Megan McCoy

PAM	Julia Davydova	7/22/2019	Jennifer Borgert
PAM	Julia Davydova	7/26/2019	Jennifer Borgert
Semi-annual	Jennifer Erickson	7/31/2019	Jennifer Borgert and Marilyn Bennett
Semi-annual	Jennifer Erickson	7/31/2019	Jennifer Borgert and Marilyn Bennett
PAM	Sunny Chan	8/1/2019	Ilana Cohen
Semi-annual	Alessandro Bartolomucci	8/7/2019	Kristin Pilon
Semi-annual	Michael Benneyworth	8/7/2019	Kristin Pilon
Semi-annual	Julia Ponder	8/6/2019	Ilana Cohen and Ji Perry
PAM	Dorraya El-Ashry	8/2/2019	Megan McCoy
PAM	Kristin Hogquist	8/6/2019	Ilana Cohen
PAM	Bryce Binstadt	8/8/2019	Ilana Cohen
РАМ	Christopher Staley	8/8/2019	Jennifer Borgert
Second Surgery Inspection	Benjamin Saunders	8/9/2019	Kristin Pilon

Second Surgery Inspection		***	Sarah Heilbronner	8/12/2019	Kristin Pilon
Semi-annual			Emily Taras	8/12/2019	Megan McCoy
PAM	-		Paulo Kofuji	8/14/2019	Ilana Cohen
Semi-annual			Anna Lee	8/13/2019	Paul Lindstrom and Dezhi Liao
Second Surgery Inspection			Suhasa Kodandaramaiah	8/15/2019	Paul Lindstrom
Semi-annual			Stephen Katz	8/15/2019	Paul Lindstrom and Sam Baidoo
Semi-annual			Mark Masino	8/15/2019	Jennifer Borgert and Christine Sivula
Semi-annual			Mark Masino	8/15/2019	Jennifer Borgert and Christine Sivula
PAM/Semi-annual			Esther Krook-Magnuson	8/14/19 and 8/20/19	Kristin Pilon and Christine Sivula
Semi-annual			David Stephens	8/16/2019	Jennifer Borgert and Marilyn Bennett
PAM		2	Alexandra Sobeck	8/20/2019	Megan McCoy
Initial Surgery Inspection			Carolyn Fairbanks	8/20/2019	Paul Lindstrom
PAM			Peter Crawford	8/21/2019	llana Cohen

Second Surgery Inspection	Karen Ashe	8/19/2019	Jennifer Borgert
PAM	DeWayne Townsend	8/22/2019	Paul Lindstrom
PAM	Jerid Robinson	8/28/2019	Kristin Pilon
Second Surgery Inspection	Rita Perlingeiro	8/30/2019	Megan McCoy
Semi-annual	Sandy Mand	8/29/2019	Jennifer Borgert and Felicia Boynto
Semi-annual	Sandy Mand	8/29/2019	Jennifer Borgert and Felicia Boynto
Second Surgery Inspection	Michael Olin	8/30/2019	Jennifer Borgert
PAM	John Belcher	8/26/2019	Jennifer Borgert
PAM	John Belcher	8/26/2019	Jennifer Borgert
PAM/Semi-annual	Mark Hove	9/6/2019	Paul Lindstrom an Jim Perry
PAM	Anna Lee	9/4/2019	Paul Lindstrom
PAM	York Marahrens	9/4/2019	Megan McCoy
PAM/Semi-annual	Frank Ondrey	9/10/2019	Jennifer Borgert and Robert Schumacher

PAM		Alexander Primus	9/6/2019	Jennifer Borgert
PAM		Massimo Costalonga	9/9/2019	Megan McCoy
PAM		Courtney Aldrich	9/6/2019	Megan McCoy
Ag		Nicky Overgaard	9/11/2019	Kristin Pilon and Felicia Boynton
Ag		Nicky Overgaard	9/11/2019	Kristin Pilon and Felicia Boynton
Ag		Terrill Giannonatti-Bradford	9/11/2019	Kristin Pilon and Felicia Boynton
Semi-annual		Rick Abrahamson	9/11/2019	Kristin Pilon and Felicia Boynton
Semi-annual		Jennifer Erickson	9/17/2019	Megan McCoy and Peggy Norris
Semi-annual		Mark Thomas and Julia Lemos	9/17/2019	Kristin Pilon and Je Hubbard
AM/Semi-annual		Ian Aldrich	9/18/2019	Kristin Pilon and Felicia Boynton
Semi-annual		Allen Mensinger	9/18/2019	Kristin Pilon and Felicia Boynton
'AM/Semi-annual	· * * * ·	Erin Olson	9/18/2019	Kristin Pilon and Felicia Boynton
Semi-annual		Jennifer Liang	9/18/2019	Kristin Pilon and Felicia Boynton

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PAM/Semi-annual		Frank Maragi	9/18/2019	Kristin Pilon and Felicia Boynton
PAM		Dezhi Liao	9/19/2019	Paul Lindstrom
Second Surgery Inspection		Amanda Klein	9/18/2019	Paul Lindstrom
PAM		Ruifeng Cao	9/18/2019	Jennifer Borgert
PAM		Yi-Mei Yang	9/18/2019	Jennifer Borgert
Semi-annual		Erin Malone.	9/23/2019	Kristin Pilon and Peggy Norris
Semi-annual		Sandra Allweiler and Kim Colvard	9/23/2019	Kristin Pilon and Peggy Norris
Second Surgery Inspection		Tate Gisslen	9/20/2019	Ilana Cohen
Semi-annual		Mark Schleiss	9/24/2019	Paul Lindstrom and Geoffrey Ghose
PAM/Semi-annual		Timothy Kurtti	9/20/2019	Jennifer Borgert and Keith Baker
PAM		Yasushi Nakagawa	9/20/2019	Jennifer Borgert
PAM/Semi-annual		Mark Bee	9/19/2019	Jennifer Borgert and Nathan Koewler
PAM/Semi-annual		Mark Bee	9/19/2019	Jennifer Borgert and Nathan Koewler

Semi-annual	Jennifer Erickson	9/25/2019	Paul Lindstrom and Mimie Pollard
Semi-annual	Brenda Mielke	9/25/2019	Kristin Pilon and Peggy Norris
Semi-annual	Sheryl Ferguson	9/25/2019	Kristin Pilon and Peggy Norris
Semi-annual	Flannery Miley	9/25/2019	Kristin Pilon and Peggy Norris
Semi-annual	Denise Obitz-Cooney	9/25/2019	Kristin Pilon and Peggy Norris
PAM	Yuying Liang	9/26/2019	Paul Lindstrom
Semi-annual	Jennifer Erickson	9/27/2019	Megan McCoy and Dezhi Liao
Semi-annual	Marc Jenkins	9/27/2019	Megan McCoy and Dezhi Liao
Semi-annual	Ann Fallon	9/27/2019	Jennifer Borgert and K. V. Nagaraja
Semi-annual	Deborah Ferrington	9/24/2019	Jennifer Borgert and Robert Schumacher
Semi-annual	Maxim Cheeran	9/30/2019	Paul Lindstrom and Felicia Boynton
Semi-annual	Michael Conzemius	9/30/2019	Paul Lindstrom and Felicía Boynton

## NOTES WRITTEN TO FILE

Investigator Name	Date of Inspection/submission	Protocol number (s)	Notes written to file
Jakub Tolar	5/13/2019	1808-36286A	PI request modification of a procedure regarding skin grafts to a less invasive method that will not require a chamber implant but uses a gelatin printed graft implant instead. Previously the chamber would be removed on day 5 requiring an additional anesthetic event. The gelatin printed graft negates this additional event. The area veterinary was in consult for this procedural improvement.
Gregory Beilman	5/14/2019	1805-35872A	PI submits to IACUC additional details of when supplemental fluid would be provided in pigs. The addition includes: after 4 hours of full resuscitation, if animal has a SBP>90 mmHG continue to observe; SBP < 90mmHG a lactate <2, and urine output >1cc/kg/hr, continue to observe urine output <1cc/kg/hour, give fluid bolus; Lactate >2, give fluid bolus
Daniel Gallaher	5/30/2019	1801-35506A	PI requests an additional six animals added to the protocol for use for training purposes.
Jizhen Lin	6/5/2019	1610-34251A	Noted on inspection that anti-PD antibodies are given by IP route but protocol indicates that they will be given IV
Brian Betts	6/5/2019	1807-36180A	PI requests clarification on timing of analgesics given (Carprofen every 24 hours as needed for breakthrough pain up to 72 hours post skin surgery)
Esther Krook-Magnuson	6/11/2019	1801-35497A	PI requests adding the distribution of sweet tasting food crumbs (e.g. chocolate sprinkles) to the open field behavioral test to encourage exploration
Jayanth Panyam	6/17/2019	1605-33821A	PI requests the use of a modified diet (purified AIN93G) for some mice that will be used in the imaging studies. The diet is nutritionally equivalent to normal feed but is better for immunofluorescence as it is supposed to decrease the background fluorescence when imaging.

## NOTES WRITTEN TO FILE

Valerie Brady	7/11/2019	1702-34612A	PI requests to increase the number of animals that are fin clipped than what was originally approved. This year they will be marking fishes in the marked lipping. Up to approximately 2000 fish will be marked (much likely less). The experimental design, procedures and attachment sections have all been updated to reflect the increase in the number of fishes receiving fin clips. Fin clipping is currently approved as a procedure in the protocol.
Yibin Deng	7/16/2019	1608-34064A	PI will use Nair to remove fur from mice prior to surgery. Currently, protocol indicates removal of top layer of fur using shearing scissors.
Courtney Johnson	7/19/2019	1806-36014A	Pl requests to include additional inclusion criteria for the dog study: to include dogs euthanized at the local animal control facility for reasons unrelated to the study and dogs euthanized at <b>the study</b> for reasons unrelated to the study
David Bereiter	7/30/2019	1802-3557A and 1802-35558A	Noted on inspection that only urethane is used for terminal procedures but protocol was updated last inspection to use both urethane and Isoflurane. Verbiage stating that either or both can be used during the approved terminal procedures
Natalia Tretyakova	9/25/2019	1706-34933A	PI requests 18 mice to be added to the protocol
Ulrike Munderloh	9/27/2019	1804-35774A	Noted on inspection that rickettsia inoculation and subsequent challenge injections done via IP injection but protocol is approved for IV tail vein injection.

Date of Recommendation	Investigator	Species	Protocol Number	Recommendation
4/8/2019	Dale Gregerson	mouse	1706-34882A	The lab is experiencing mortality of young (age 3-6 week) and parabiosed mice following anesthesia with 100mg/kg ketamine 10 mg/kg xylazine for fundus exams. This recommendation is to use an 8 mg/kg dose of ketamine for these more vulnerable mice. Lab staff will also provide 0.5-1mL warm saline SC at the time of anesthetic induction and cover the microscope stage and the mouse to retain heat. In addition, to prevent stress of parabiosesd mice associated with restraint for IP injections, paired mice will be anesthetized in an isoflurane induction chamber, then removed and injected with ketamine/xylazine IP
4/24/2019	Yibin Deng	mouse	1608-34064A	This veterinary recommendation is to modify both anesthesia and analgesia in the protocol to limit steps and stress to the mouse versus what is approved in the protocol. This recommendation is specifically for the allograft prostate cancer mouse models surgery but also pertains to the surgical castration surgery. For analgesia, use of SR buprenorphine at 2mg/kg, SC 2-4 hours prior to the surgical procedures. This is in place of starting with standard buprenorphine and then switching to SR buprenorphine. This will limit the number of injections the mouse receives. For anesthesia: use isoflurane at a range of 1-5% for both induction and maintenance (induction will be at the higher percentages and maintenance will be at the lower percentages. This is a change from starting with ketamine/xylazine for induction then switching to Isoflurane.

5/1/2019	Ferenc Toth	pig	1704-34762A, 1703-34645A; 1901-36656A	The drugs and doses listed below may be used for premedication, induction or maintenance of anesthesia (in addition to what is already listed on the protocol). The various options give lab staff tools to maintain physiologic homeostasis during surgical or imaging procedures, and minimize distress during premedication and induction: Telazol/xylazine; telazol/ketamine/xylazine; ketamine/acepromazine/atropine; Fentanyl; Also a urinary catheter may be placed in female pigs to aid in voiding during anesthesia. Male pigs may be manually expressed. If this is unsuccessful, cystocentesis may be performed to empty the bladder
5/7/2019	Dale Gregerson	mouse	1706-34882A	To improve post operative nutrition and reduce morbidity in parabiosed mice, lab members will administer 0.5-1MI SC fluids (LRB, LRB with 5% dextrose or 0.9% NaCl) to each mouse daily for 7 days after surgery. To encourage food consumption, moist food and diet get will be supplemented with autoclaved sunflower seeds and banana chips. These supplements will not replace the standard, fully-balance rodent diet but will be provided in small amounts to ensure that mice are in taking sufficient calories for appropriate recovery.
5/24/2019	Wei Chen	rat and cat	1903-36898, 1606-33914	Because the NeoPredEf powder is on backorder through the end of 2019, a veterinary recommendation was made to use topical supportive agents post operatively as needed for rats and cats undergoing craniotomies. Topical products include triple antibiotic ointment, Collasate, Rediheal and Wonderdust. The products can be used 1-2X daily either individually or in combination to aid in wound healing

6/4/2019	Brian Betts	mouse	1807-36180A	Recommended changes within the Skin Graft Surgery Procedure: make clear SR buprenorphine is the drug being used (versus standard buprenorphine) at a dose of 2mg/kg SC once pre-operatively 2-4 hours prior to surgery; Carprofen dosing should be changed to once a day dosing (for unrelieved pain not covered by SR buprenorphine on an as needed basis; Additionally please add range of carprofen from 5-10 mg/kg. Since it will be used synergistically with buprenophine , the lowest possible dose for effective pain control should be used; Starting with a lower dose would be the best in this case and then more can be added if pain is still noted
6/26/2019	Esther Krook-Magnuson	mouse	1801-35497A	The recommendation is to change from using specifically Neo- Predef to any topical antibiotic ointment/powder +/- steroid +/- topical anesthetic; Topical antibiotic ointments/powders are commonly used post surgery and may contain a variety of different combinations of antibiotics as well as being combined with a variety of different topical anti- inflammatories/anesthetics. Currently, NeoPredef is on backorder and unavailable. Any of the common alternative topical products would be an adequate replacement. Examples include but are not limited to TriTop, Neopolydex, Vetropolycin HC, Derma vet. Having the ability to use alternative treatments is critical to maintaining animal welfare when specific products are unavailable.

7/15/2019	Paul laizzo	sheep	1707-35001A	Premedication and induction relieve animal anxiety, reduce anesthetic total dose needed to maintain a surgical plan of anesthesia and allow for catheterization and intubation of animals. Currently, there are two sheep procedures approved on this protocol (Prototype Transcatheter Device Testing and Medial Sternotomy and general surgical procedures: Heart Explantation); Each of these procedures lists different premedication and induction agents. The recommendation is to allow the use of both premedication/induction regimens for either procedure as both could be appropriate based on the animals individual needs and different response to different medications. The PI or his designee will amend the protocol to reflect these changes by 10/15/19. Premedication: Ketamine of Ketmaine/Diazepam; Atropine may be given to help with oral secretions if deemed necessary for intubation; Induction: Propofol to effect (may not be needed if using Ketamine/diazepam combination)
8/19/2019	Alessandro Bartolomucci	mouse	1706-34930A	To reduce overall stress of animals associated with daily oral gavage, lab members may administer Tamoxifen in specially formulated and commercially available Tekland diet. The lab will be responsible for administering the diet, checking food levels daily to ensure mice have enough to eat and documenting when they have changed/filled the food

8/15/2019	Richard Bianco	sheep	1905-37042A	Based on previous history, it seems that the size of the device may result in some functional compromise of the heart during recovery. The drug digoxin can be used to increase cardiac contractility and potentially alleviate some of these effects. The use of the drug may be able to keep the animal healthy with very little risk of negative side effects. This will benefit both the animal and the study, and require less total animals if survival and well being can be improved. Following surgery if
8/19/2019	Marija Cventanovic	guinea pigs	1703-34631A	This vet recommendation is to allow housing of guinea pigs in building 75 of the Itasca Biological Station as an alternative housing are in the event that the environmental conditions fall outside the acceptable humidity range and the recommended temperature range in building 45 as dictated by the Guide. This recommendation is in effect for 8/19/19 through the end of the day on 8/23/19. Consideration should be given to the ventilation, illumination, noise level and other parameters that affect animal and handler health while housed in building 75
8/22/2019	Harry Orr	mouse	1708-35065A	To help mitigate weight loss secondary to daily IP injections of CCK receptor agonists lab members will provide animals with moist feed and sucrose water. Mice will continue to receive a standard, fully balanced rodent diet and may receive 0.5-1.0 mL SC fluids (LRS, LRS with 5% dextrose or 0.9% NaCl) if there is clinical evidence of dehydration. Lab members will euthanize mice if they lose 25% weight or if DVM determines it is appropriate
9/4/2019	Harry Orr	mouse	1708-35065A	To help mitigate weight loss secondary to daily IP injections of CCK receptor agonists, lab members will provide diet supplements including seeds, nuts, banana chips, diet get, etc. All supplements will be provided to mice in consultation with DVM. These supplements will NOT replace the standard, fully balanced rodent diet, but will be provided in small amounts to ensure mice are in taking sufficient nutrients.

9/12/2019	Robert Tranquillo	sheep	1808-36236A	Appropriate use of antibiotics is important for animal health and the prevention of unnecessary antibiotic resistance. The procedure "1-2 week, 4 week and 8 week angiogram to evaluate valve function" details in the post operative instructions to RAR to give ceftiofur/ceftriaxone for seven days after each of these angiograms. Assuming there is no gross contamination during the procedure, this use of antibiotics is likely unnecessary as the risk for post surgical infection is low. The recommendation is to change the instructions for post operative care from "Ceftiofur 3 mg/kg IM or Ceftrioxone 1 gIM every 24 hours for 7 days to "if there is a concern for an infection following procedure, a RAR veterinarian will be consulted for antibiotic recommendations"
9/13/2019	Mark Bee	frogs	1701-24456A	Tree frogs may be housed in either the modified flow through aquaria currently described in the IMHA section or in static aquaria. Animals may be housed with out without sphagnum moss. Newly captured animals have been temporarily housed in static aquaria without moss per veterinary instruction for approximately 8 weeks. Animals housed in static aquaria have exhibited a notably lower rate of attrition than animals housed in flow through aquaria in years past. The reason is unclear but the outcome suggests the animals benefit from this housing scheme. Animals will be provided fresh water daily in a cup/bowl on the floor of the enclosure in lieu of flow through water. Daily observations, cleaning schedules and methods, etc will remain unchanged. Long term static housing is a new endeavor for this group. Their PI and vet together may choose to make minor modifications based on experience and circumstances when deemed beneficial to animal health and well being. Minor changes will be included in the protocol renewal along with the description of static aquaria. Major changes will be submitted as amendments for IACUC review and approval

9/13/2019		NHP	1804-35859A	Side effects of harmaline administration used in the Tremor Induction procedure may include vomiting and vertigo. To kee the animal hydrated and comfortable, they may be administered 5-10 mL/kg of SC fluids (LRS or similar) and may be given one of the following anti-nausea medications: Maropitant 1-2 mg/kg SC, IV, PO or ondansetron 0.15 mg/kg Sc
9/23/2019	Vaiva Vezys	mouse	1611-34309A	A reversal agent may be administered to animals undergoing anesthesia for LINAC irradiation. Acceptable reversible agent are atipamizole (Antisedan, 1-2 mg/kg SC or IP) or yohibine (Yobine, 0.5-1 mg/kg IP). Reversal agents hasten recovery of consciousness and return to normal activity. When available animals may be anesthetized with inhaled Isoflurane instead of ketamine/xylazine. Isoflurane has a rapid onset and animals recover quickly once the gas is withdrawn. Animals do not need to be injected. The combined benefit is shorter duration of anesthesia and a marked decrease in handling time and number of injections.
9/30/2019	Patrick Rothwell	mouse	1810-36447	This veterinary recommendation is for using brief Isoflurane exposure to remove wound clips post operatively in mice. Th helps the animal remain still during clip removal and avoids ar additional stress to the surgical site related to tissue pulling. Clips should be removed 7-14 days after surgery

Protocol ID	Principal	Species	Guideline	Exception Request with
1909-37384A	Yang, Yi-Mei	Mice	EUTHANASIA	Mice are sacrificed by decapitation
	Yang, Yi-Mei		SOCIAL	
1909-37384A	(Amy)	Mice	HOUSING	see protocol for details
				Mice will develop glioma (20-70% of
				all cohorts) and the accompanying
				CNS symptoms - part paralysis
				(30%), seizures (1%), head tilting
				(20%), running/walking in circles
				(5%), cerebral edema (20%).
				Pronounced CNS symptoms will
				cause mice to be euthanized;
				however, we will not euthanized
				mice with moderate paralysis, head
				tilting or cerebral edema as long as
				they can access food and water and
				do not otherwise seam in distress.
				Mice in this state will be monitored
				for dresses progression. Although,
				the gliomas are not expected to
				cause the mice any direct pain mice
				with pronounced cranial swenning,
				debilitating paralysis or odoma
				maybe in pain and will be
				euthanized
			TUMOR	The paralysis of a single back leg
	Robinson			we consider nondebilitating, two leas
1908-37346A	lames	Mice		or a single front leg debilitating
1000 07 0 107	bameo	Miloc	ORTERN	
				Pentobarbitol will be purchased from
				Sigma, prepared in small filter-
				sterilized batches, and stored in
				injection vials. Because commercial
				pentobarbitol requires purchase in
				large quantities (10ml) we typically
				see variability in the potency of the
				arug over the lifetime of the vial that
				the mise shallonging. The shift to
			NON	me mice challenging. The ability to
				make smaller batches of drug, that
				us to more accurately does our miss
1008-373///	Nielsen Kirston	Mice Mice		and will result in fewer adverse
1000-07 044A	INCISCI, NISLEIL			and will result in rewer duverse

				For the parabiotic pairing/separation
				procedure, two surgeries will need to
				be performed on the mice. First is
				parabiotic surgery to join the two
				mice After 1-6 months the
				parabiosed mice will need to
				parabiosed mice will need to
				undergo second surgery to be
				separated for neuropenavioral
				assessments as described in the
				neurobehavioral testing procedure.
				Thus, the parabiotic
				pairing/separation surgeries are
				essential components of the same
				project. The separation surgery will
				not cause any more pain and
				distress than the first parabiotic
				surgery. Functionally, it will restore
				the functionality of the individual
				mice. 🗆
				П
				During the parabiosed period, some
				of the mice will be subjected to
				intracerebral injection of amyloid-
				beta (to induce Alzheimer's-type
				nathology including
				pathology mordaring
				neuronnammation). These
				procedures are necessary to study
				the mechanisms of regulation the
1000 070404			MULTIPLE	Immune cells in the brain. The
1908-37310A	Li, Ling	Місе	SURGERY	parabiosed mice are the best
				We use either under-gravel filters
				where the sediment waste on the
				gravel is siphoned out monthly or
				above-tank charcoal filter systems
				(especially for the marine tanks)
				where the charcoal cartridges are
				changed based on manufacturer's
				recommendations. We have protein
				skimmers for the marine tanks. We
			SANITATION	also have an Z-Hab system in Biosci
1907-37285A	Mand, Sandy	Fish (Other)	FREQUENCY	115 which includes a bio filter, filter,
			SOCIAL	
1907-37285A	Mand, Sandy	Fish (Other)	HOUSING	
				In order to assess how CNS
				leukemia evolves over time it may
				be necessary to sample CSF from
				the same mouse. However, similar
				to repeat blood draws, this is a
				relatively minor surgical procedure.
			MULTIPLE	CSF will not be obtained any more
1907-37283A	Gordon, Peter	Mice	SURGERY	than once per week for a up to a

				we are performing lancal apesthetic
				we are performing laboral anesthetic
				blocks for the purpose of diagnosis.
				We will record doses used and sites
				of administration butthe animal
				remains conscious and unsedated
				throughout - we will not be recording
				vital signs (Equine Lameness Exam -
				with local anesthetic blocks)
				sedation only; animals remain
				conscious and upright (Use of
				Sedation for Equine Procedures)
				a la mana antanan arang manan katang tang katang arang ar
			ANESTHESIA.	sedation only, animals remain
			SURGERY.	conscious throughout (Use of
			AND POST-	Sedation for Bovine Procedures)
			PROCEDURAL	
		Horse Cow	RECORDKEEP	local anesthetic bleb only for
1907-37280A	Madill Scott	(Biomedical)	ING	placement of jugular catheter (Blood
		()		Our lab has found in the preliminary
				studies from the expiring protocol
				that the frequent handling of the
				mice (weekly weighing and cage
				changes monthly
				echocardiography echo-MRI and
				blood pressure monitoring) has
				bindered the mice from gaining the
				weight we would expect on this
				HED In consultation with Dr. Cathy
				Ketz, an expert in chase mayor
				Roiz, all expert in obese mouse
				models, she suggested that we only
				weigh the mice bi-weekly. This bi-
				weekiy weigning, along with our
4007 07075 4	Time the	N 4:		decrease in data collection as
1907-37275A	i imotny	WICE		spelled out in this renewal (baseline,
1007 07000		o ·	SOCIAL	
1907-37262A	Schleiss, Mark	Guinea Pig	HOUSING	see protocol for details

1907-37261A	Dong, Zigang	Mice	MULTIPLE SURGERY	We will attempt to increase vascularity by first inducing a stromal reaction by implanting either a subcutaneous glass disc or Gelfoam dressing 2 weeks prior to the xenograft. Patel, Girish K et al. "A Humanized Stromal Bed Is Required for Engraftment of Isolated Human Primary Squamous Cell Carcinoma Cells in Immunocompromised Mice." The Journal of investigative dermatology 132.2 (2012): 284–290. PMC. Web. 30 May 2017. The implant will sit in the subcutaneous space and the incision will be small, therefore we
				only individuals with proper training
1907-372574	Eckfeldt Craig	Mice	METHOD	and a high degree of technical
1001 012017	London, ordig	inioo		pronoionoy win perform oerviour

				Simultaneous with ovariectomy a
				subset of mice will have herve cuffs
				implanted. The ovariectomy
				procedure takes less than 5 minutes
				beyond the never cuff and it is less
				stressful for the mouse than having
				U One coal of the preject is to
				One goal of the project is to
				determine the effect of estrogen on
				muscle regeneration. A second
				surgery we are requesting is freeze
				injury or cardiotoxin injury to the
				tibialis anterior muscle, which are
				minimally invasive surgeries. That
				is, each involves making a skin
				incision and then placing a freezing
				probe on the muscle or injecting the
				muscle with cardiotoxin to induce
				injury and subsequent muscle
				regeneration. Mice will be 2-8 weeks
				post-ovariectomy before either type
				injury is induced. A subset of
				cardiotoxin injured mice will be
				transplanted as well.□
				(Ovariectomy)
				The goal of one study is to
				determine the effect of estrogen on
				muscle regeneration. Thus, a
				second surgery following
			MULTIPLE	ovariectomy will be freeze injury to
1907-37248A	Lowe, Dawn	Mice, Mice	SURGERY	the muscle, which is minimally

				Antibiotics and anti-inflammatory
				reagents are acceptable if necessary
				for the first surgery nerve
				ouff/ovariactomy, but are not
				cultiovallectority, but are not
				acceptable after freeze injury or
				cardiotoxin injury because we are
				studying the inflammation. (Nerve
				cuff implantation)
				We will not give injections of Buprenorphine after the BaCl injury because the surgery is minimal (no body cavity is opened) and we have
				not noticed signs of post-surgical
				distress. Further, the BaCl injury is
				toone specific muscle in a group of muscles that cause dorsiflexion. We
				will continually work with the yet
				staff in order to be sure that there is
				adequate attention paid to the extent
				of post-surgical pain and function
				(ambulation).
				Parameters that will be continually
				monitored in these mice include impaired
				circulation in the foot, intense
				lethargy (as indicated by limited
				mobility, hunched posture.
			72 HOUR	limited aroomina) severe
			POST-OP	inflammation (as indicated by bright
				red coloration of the local tissue
1907-372484	Lowe Dawn	Mice Mice	POLICY	with
1001-012-01	Lowe, Duwn			To monitor physical activities (wheel
			SOCIAL	rupping capacity or cage activities)
1007 372484		Mico Mico		mice will be boused individually.
1307-07240A		wilce, wilce		To monitor physical activities (wheel
				rupping copposity or code activities
				mino will be boused individually. $\Box$
				The environment will influence the
				physical activity of the mice which in
				turn will affect skeletal muscle
				function. Because the magnitude of
				physical activities will likely vary
				depending on hormonal status an
				enriched environment would add
				another level of variables that at this
1907-37248A	Lowe, Dawn	Mice. Mice	ENRICHMENT	time we do not wish to explore
	, Dami		SOCIAL	
1907-37238A	Chen, Chi	Mice	HOUSING	see protocol for details

1907-37217A	Koewler, Nathan	Mice, Rat	EUTHANASIA METHOD	RAR veterinary staff are proficient in this procedure and in the case of health conditions where CO2 may not be readily available this procedure will be used to provide
		Dog, Frog (Xenopus), Gerbil, Goat, Guinea Pig, Hamster, Mice, Nonhuman Primate (Macaques), Pig (Biomedical), Rabbit, Rat, Sheep (Biomedical), Turkey, **ALL SPECIES DESIGNATION **, Cow (Biomedical), Chinchilla,	SOCIAL	
		Dog, Frog (Xenopus), Gerbil, Goat, Guinea Pig, Hamster, Mice, Nonhuman Primate (Macaques), Pig (Biomedical), Rabbit, Rat, Sheep (Biomedical), Turkey, **ALL SPECIES DESIGNATION **, Cow (Biomedical), Chinchilla,	EUTHANASIA	Euthasol® 0.22 ml/kg IV (~86 mg/kg

				We require a method of animal
				Identification that is unambiguous
				mice in developmental studies
				(postnatal pups, genotyping results
				required at P8) or in aging studies
				over 1 year. We cut toes after they
				are no longer webbed in mice P6-P8
				and use the toes for genotyping.
1907-37213A	Junge, Harald	Mice	TAIL BIOPSY	Aseptic practices are followed. In
				We observed in preliminary
				experiments that Mdm2 ECKO mice
				die about 1 week after tamoxifen
				Induced recombination using a Cdn5-
				death appears to be pleural effusion
				We suthanize the mice $5-6$ days
				after tamoxifen injection at which
				time animals are showing the first
				sign of being lethargic. We
				euthanize them as early as possible
			EUTHANASIA	to prevent distress but as late as
			DEATH/MORIB	necessary until the relevant
the second second second		a. 1999	UND	phenotype (blood-retina barrier
1907-37213A	Junge, Harald	Mice	ENDPOINT	defects) manifests. If the animal
				/
				sulfo-NHS biotin is not available
				sulfo-NHS biotin is not available USP grade, it will be ordered from
			NON-	sulfo-NHS biotin is not available USP grade, it will be ordered from Sigma. It will be made up fresh from
			NON- PHARMACAUT	sulfo-NHS biotin is not available USP grade, it will be ordered from Sigma. It will be made up fresh from powder in PBS (0.3 mg/ml) and
			NON- PHARMACAUT ICAL GRADE	sulfo-NHS biotin is not available USP grade, it will be ordered from Sigma. It will be made up fresh from powder in PBS (0.3 mg/ml) and sterile filtered. This agent is
1907-37213A	Junge, Harald	Mice	NON- PHARMACAUT ICAL GRADE COMPOUNDS	sulfo-NHS biotin is not available USP grade, it will be ordered from Sigma. It will be made up fresh from powder in PBS (0.3 mg/ml) and sterile filtered. This agent is frequently used in blood-brain
1907-37213A	Junge, Harald	Mice	NON- PHARMACAUT ICAL GRADE COMPOUNDS EUTHANASIA	sulfo-NHS biotin is not available USP grade, it will be ordered from Sigma. It will be made up fresh from powder in PBS (0.3 mg/ml) and sterile filtered. This agent is frequently used in blood-brain Sedation if not normally used in
1907-37213A 1907-37212A	Junge, Harald Dickerson, Erin	Mice Mice	NON- PHARMACAUT ICAL GRADE COMPOUNDS EUTHANASIA METHOD	sulfo-NHS biotin is not available USP grade, it will be ordered from Sigma. It will be made up fresh from powder in PBS (0.3 mg/ml) and sterile filtered. This agent is frequently used in blood-brain Sedation if not normally used in combination with barbiturate
1907-37213A 1907-37212A	Junge, Harald Dickerson, Erin	Mice Mice	NON- PHARMACAUT ICAL GRADE COMPOUNDS EUTHANASIA METHOD	sulfo-NHS biotin is not available USP grade, it will be ordered from Sigma. It will be made up fresh from powder in PBS (0.3 mg/ml) and sterile filtered. This agent is frequently used in blood-brain Sedation if not normally used in combination with barbiturate In order to assess the efficacy of the vaccine, the infection must be
1907-37213A 1907-37212A	Junge, Harald Dickerson, Erin	Mice Mice	NON- PHARMACAUT ICAL GRADE COMPOUNDS EUTHANASIA METHOD	sulfo-NHS biotin is not available USP grade, it will be ordered from Sigma. It will be made up fresh from powder in PBS (0.3 mg/ml) and sterile filtered. This agent is frequently used in blood-brain Sedation if not normally used in combination with barbiturate In order to assess the efficacy of the vaccine, the infection must be allowed to progress for 72 hours to
1907-37213A 1907-37212A	Junge, Harald Dickerson, Erin	Mice Mice	NON- PHARMACAUT ICAL GRADE COMPOUNDS EUTHANASIA METHOD	sulfo-NHS biotin is not available USP grade, it will be ordered from Sigma. It will be made up fresh from powder in PBS (0.3 mg/ml) and sterile filtered. This agent is frequently used in blood-brain Sedation if not normally used in combination with barbiturate In order to assess the efficacy of the vaccine, the infection must be allowed to progress for 72 hours to analyze the difference between
1907-37213A 1907-37212A	Junge, Harald Dickerson, Erin	Mice Mice	NON- PHARMACAUT ICAL GRADE COMPOUNDS EUTHANASIA METHOD EUTHANASIA DEATH/MORIB	sulfo-NHS biotin is not available USP grade, it will be ordered from Sigma. It will be made up fresh from powder in PBS (0.3 mg/ml) and sterile filtered. This agent is frequently used in blood-brain Sedation if not normally used in combination with barbiturate In order to assess the efficacy of the vaccine, the infection must be allowed to progress for 72 hours to analyze the difference between vaccinated and unvaccinated
1907-37213A 1907-37212A	Junge, Harald Dickerson, Erin Pravetoni.	Mice Mice	NON- PHARMACAUT ICAL GRADE COMPOUNDS EUTHANASIA METHOD EUTHANASIA DEATH/MORIB UND	sulfo-NHS biotin is not available USP grade, it will be ordered from Sigma. It will be made up fresh from powder in PBS (0.3 mg/ml) and sterile filtered. This agent is frequently used in blood-brain Sedation if not normally used in combination with barbiturate In order to assess the efficacy of the vaccine, the infection must be allowed to progress for 72 hours to analyze the difference between vaccinated and unvaccinated groups. While death will not be the
1907-37213A 1907-37212A 1906-37191A	Junge, Harald Dickerson, Erin Pravetoni, Marco	Mice Mice Mice	NON- PHARMACAUT ICAL GRADE COMPOUNDS EUTHANASIA METHOD EUTHANASIA DEATH/MORIB UND ENDPOINT	sulfo-NHS biotin is not available USP grade, it will be ordered from Sigma. It will be made up fresh from powder in PBS (0.3 mg/ml) and sterile filtered. This agent is frequently used in blood-brain Sedation if not normally used in combination with barbiturate In order to assess the efficacy of the vaccine, the infection must be allowed to progress for 72 hours to analyze the difference between vaccinated and unvaccinated groups. While death will not be the primary endpoint, it is known in this
1907-37213A 1907-37212A 1906-37191A	Junge, Harald Dickerson, Erin Pravetoni, Marco	Mice Mice Mice	NON- PHARMACAUT ICAL GRADE COMPOUNDS EUTHANASIA METHOD EUTHANASIA DEATH/MORIB UND ENDPOINT	sulfo-NHS biotin is not available USP grade, it will be ordered from Sigma. It will be made up fresh from powder in PBS (0.3 mg/ml) and sterile filtered. This agent is frequently used in blood-brain Sedation if not normally used in combination with barbiturate In order to assess the efficacy of the vaccine, the infection must be allowed to progress for 72 hours to analyze the difference between vaccinated and unvaccinated groups. While death will not be the primary endpoint, it is known in this Grouse are being captured in the
1907-37213A 1907-37212A 1906-37191A	Junge, Harald Dickerson, Erin Pravetoni, Marco	Mice Mice Mice	NON- PHARMACAUT ICAL GRADE COMPOUNDS EUTHANASIA METHOD EUTHANASIA DEATH/MORIB UND ENDPOINT	sulfo-NHS biotin is not available USP grade, it will be ordered from Sigma. It will be made up fresh from powder in PBS (0.3 mg/ml) and sterile filtered. This agent is frequently used in blood-brain Sedation if not normally used in combination with barbiturate In order to assess the efficacy of the vaccine, the infection must be allowed to progress for 72 hours to analyze the difference between vaccinated and unvaccinated groups. While death will not be the primary endpoint, it is known in this Grouse are being captured in the field with a noose pole - if a rare
1907-37213A 1907-37212A 1906-37191A	Junge, Harald Dickerson, Erin Pravetoni, Marco	Mice Mice Mice	NON- PHARMACAUT ICAL GRADE COMPOUNDS EUTHANASIA METHOD EUTHANASIA DEATH/MORIB UND ENDPOINT	sulfo-NHS biotin is not available USP grade, it will be ordered from Sigma. It will be made up fresh from powder in PBS (0.3 mg/ml) and sterile filtered. This agent is frequently used in blood-brain Sedation if not normally used in combination with barbiturate In order to assess the efficacy of the vaccine, the infection must be allowed to progress for 72 hours to analyze the difference between vaccinated and unvaccinated groups. While death will not be the primary endpoint, it is known in this Grouse are being captured in the field with a noose pole - if a rare injury occurs, there are limited
1907-37213A 1907-37212A 1906-37191A	Junge, Harald Dickerson, Erin Pravetoni, Marco	Mice Mice Mice	NON- PHARMACAUT ICAL GRADE COMPOUNDS EUTHANASIA METHOD EUTHANASIA UND ENDPOINT	sulfo-NHS biotin is not available USP grade, it will be ordered from Sigma. It will be made up fresh from powder in PBS (0.3 mg/ml) and sterile filtered. This agent is frequently used in blood-brain Sedation if not normally used in combination with barbiturate In order to assess the efficacy of the vaccine, the infection must be allowed to progress for 72 hours to analyze the difference between vaccinated and unvaccinated groups. While death will not be the primary endpoint, it is known in this Grouse are being captured in the field with a noose pole - if a rare injury occurs, there are limited options in the field. Rapid cervical
1907-37213A 1907-37212A 1906-37191A 1906-37189A	Junge, Harald Dickerson, Erin Pravetoni, Marco Ponder, Julia	Mice Mice Mice Bird (Other)	NON- PHARMACAUT ICAL GRADE COMPOUNDS EUTHANASIA METHOD ENTHANASIA DEATH/MORIB UND ENDPOINT EUTHANASIA METHOD	sulfo-NHS biotin is not available USP grade, it will be ordered from Sigma. It will be made up fresh from powder in PBS (0.3 mg/ml) and sterile filtered. This agent is frequently used in blood-brain Sedation if not normally used in combination with barbiturate In order to assess the efficacy of the vaccine, the infection must be allowed to progress for 72 hours to analyze the difference between vaccinated and unvaccinated groups. While death will not be the primary endpoint, it is known in this Grouse are being captured in the field with a noose pole - if a rare injury occurs, there are limited options in the field. Rapid cervical dislocation is the most expedient
1907-37213A 1907-37212A 1906-37191A 1906-37189A	Junge, Harald Dickerson, Erin Pravetoni, Marco Ponder, Julia	Mice Mice Mice Bird (Other)	NON- PHARMACAUT ICAL GRADE COMPOUNDS EUTHANASIA METHOD ENTHANASIA DEATH/MORIB UND ENDPOINT EUTHANASIA METHOD ENVIRONMEN	sulfo-NHS biotin is not available USP grade, it will be ordered from Sigma. It will be made up fresh from powder in PBS (0.3 mg/ml) and sterile filtered. This agent is frequently used in blood-brain Sedation if not normally used in combination with barbiturate In order to assess the efficacy of the vaccine, the infection must be allowed to progress for 72 hours to analyze the difference between vaccinated and unvaccinated groups. While death will not be the primary endpoint, it is known in this Grouse are being captured in the field with a noose pole - if a rare injury occurs, there are limited options in the field. Rapid cervical dislocation is the most expedient
1907-37213A 1907-37212A 1906-37191A 1906-37189A	Junge, Harald Dickerson, Erin Pravetoni, Marco Ponder, Julia	Mice Mice Mice Bird (Other)	NON- PHARMACAUT ICAL GRADE COMPOUNDS EUTHANASIA METHOD EUTHANASIA DEATH/MORIB UND ENDPOINT EUTHANASIA METHOD ENVIRONMEN TAL	sulfo-NHS biotin is not available USP grade, it will be ordered from Sigma. It will be made up fresh from powder in PBS (0.3 mg/ml) and sterile filtered. This agent is frequently used in blood-brain Sedation if not normally used in combination with barbiturate In order to assess the efficacy of the vaccine, the infection must be allowed to progress for 72 hours to analyze the difference between vaccinated and unvaccinated groups. While death will not be the primary endpoint, it is known in this Grouse are being captured in the field with a noose pole - if a rare injury occurs, there are limited options in the field. Rapid cervical dislocation is the most expedient

			SOCIAL	
1906-37182A	Hecht, Stephen	Rat	HOUSING	see protocol for details
	· · ·			I don't think we need a justification,
				but to clarify- lab staff will remove
				food from select cages 16 hours
			FOOD/FLUID	prior to testing and replace food at
			RESTRICTION	the appropriate time. These cages
			RECORDKEEP	will be marked by lab staff. During
1906-37180A	More, Swati	Mice	ING	all other times and for all un-marked
		Horse, Cow	SOCIAL	
1906-37178A	Madill, Scott	(Biomedical)	HOUSING	see protocol for details
			ANESTHESIA,	
			SURGERY,	
			AND POST-	Caudal epidural analgesia only,
			PROCEDURAL	animal is awake and standing
		Horse, Cow	RECORDKEEP	(Bovine superovulation and embryo
1906-37178A	Madill, Scott	(Biomedical)	ING	flush)
				We use either or both under-gravel
				filters where the sediment waste on
				the gravel is siphoned□
				out monthly or hang-on-tank
				charcoal filter systems where the
				charcoal cartridges are changed□
				~monthly. We will also have zeolite
	McGaugh,		SANITATION	on hand to add to filters as needed,
1906-37158A	Suzanne	Fish (Other)	FREQUENCY	to protect further against□
				Weighing fish is exceptionally
				stressful as the fish has to be netted
				and transferred to a weigh cup. $\square$
			WEEKLY	We will test sleep in these animals
			WEIGHING	at 0days, 15days, and 30days at
	McGaugh,		EXCEPTION	which point weight will be measured.
1906-37158A	Suzanne	Fish (Other)	(FOOD/FLUID)	Aside from those time points, we
			ANESTHESIA,	
			SURGERY,	This is a simple procedure taht is
			AND POST-	completed in less than a minute.
	1000 M		PROCEDURAL	anesthesia (if given) depth will only
the late by the second second of the	Cheeran,		RECORDKEEP	be measured at the beginning of the
1906-37154A	Maxim	Mice	ING	procedure. (Selection and breeding)

				I he two surgeries which are to be performed on the mice are a part of
				the same project. First, a brain
				tumor would be grown by
				intracranially implanting tumor cells.
				L The intrographic implementation of the
				electrodes (platinum wires) would be
				done at Day 5 after injecting the
				tumor cells in the mice brain. The
				electrodes would be inserted on the
				two sides of the region of tumor cell
				Implantation, such that this area lies
				electrodes. The procedure has been
				explained in the surgery section in
				detail. The electrodes would be fixed
				to the scalp and a dental cement
				would be used to fix the electrodes
				on the scalp, to make them
				he implanted both in the control
				group and the experimental group.
				However, the control group will not
				undergo electric field therapy.
				Since both the surgeries are
				Intracranial, there might be functional deficit during or after the
1906-37149A	Chen. Clark	Mice	SURGERY	procedure. In case there is
	,		SOCIAL	
1906-37149A	Chen, Clark	Mice	HOUSING	see protocol for details
				It is not standard protocol to sedate
1006 271 474	Dlubar Liz			pet dogs prior to euthanasia.
1900-37147A	Gallaher	בטק, בטק	SOCIAL	Eurranasia wiii not be performed
1906-37143A	Daniel	Rat	HOUSING	see protocol for details

				In veterinary practice, brood mares
				with a Caslick have it replaced each
				vear (it is opened prior to foaling so
				the tissue does not tear and
				replaced subsequently). While our
				mares are unlikely to get a Caslick
				each vear it is possible they could
				get another surgery related to their
				general care on the governing
				protocol. This would most likely be
				to sew up a laceration experienced
				through mishap or potentially
				placement of a subpalpebral lavage
				system to treat an ulcerated cornea.
				These are rare (generally $<1-2$ per
				vear) but might happen and since
				our horses stay in the herd a long
				time it is possible that over multiple
			MULTIPLE	vears a mare may have several
1906-37140A	Madill, Scott	Horse	SURGERY	minor surgeries to repair various
	,		72 HOUR	This is a simple skin incision and
			POST-OP	suturing, standard of care in
			ANALGESIA	veterinary practice for this surgery is
1906-37140A	Madill, Scott	Horse	POLICY	no analgesia beyond the local
				This is a simple skin-deep incision
				and suturing performed under local
				anesthesia (with sedation if
			ANESTHESIA,	required). Standard of care in
			SURGERY,	veterinary practice for this surgery is
			AND POST-	the animal is not specifically re-
			PROCEDURAL	examined until suture removal.
			RECORDKEEP	Should the incision break down
1906-37140A	Madill, Scott	Horse	ING	there is no danger to the animal and
			SOCIAL	
1906-37140A	Madill, Scott	Horse	HOUSING	see protocol for details
			EUTHANASIA	The goal is to determine if a hatch
			DEATH/MORIB	year raptor has the instinct to take
			UND	live prey by demonstrating the
1906-37139A	Ponder, Julia	Mice	ENDPOINT	ability to capture and kill said prey.
				We request not to give analgesics
				because we need for hyperalgesia to
				fully develop and analgesics may
				interfere with this process. Analgesic
				will be used if animals show
				excessive pain like behaviors (eg.
				vocalization, restlessness etc.). We
			70 110110	and others lab extensively used this
			72 HOUR	model. Based on our experience
				and available data, animals tolerate
			ANALGESIA	well this surgical procedure and do
1906-37138A	Banik, Ratan	Mice, Rat	POLICY	not show excessive pain behaviors.

				Animals can be subdued with a towel wrapped around them,
				allowing for injection of Euthasol.
				Mice will be euthanized with 0.2ml
				Euthasol (390 mg/ml sodium
				pentobarbital)
				Animala will be asvered with a sloth
4000 074004	Danile Datas	Mine Det		Animals will be covered with a cloth
1906-37138A	Banik, Ratan	Mice, Rat		to subdue them. Once they are
	Bradley,		SOCIAL	
1906-37137A	Elizabeth	Mice	HOUSING	see protocol for details
				The objective of this project is to
				study neuroinflammation after
				second injury and therefore it is
				essential to do second surgery on
				the same animal. We will perform
				craniotomy on both sides during the
				first surgery itself in order to
				minimize pain and distress during
				second surgery. The mice will be
				placed on a heating pad or under a
				heating lamp for recovery. Mice will
				be administered topical analgesic for
				pain control before surgery. Mice will
	Cheeran,		MULTIPLE	be assessed daily for hydration and
1906-37124A	Maxim	Mice	SURGERY	signs of distress. If animal cannot

				The animals will be monitored until
				they can independently maintain
				sternal recumbency or can stand
				and move about before leaving the
				and move about before leaving the
				surgery room. Fain post-surgery is
				scalp/skin incision. Animals will
				receive an application of lidocalne
				gel (2%) in and around the skinu
				Incision every 2 hours post-op as
				needed (evidence of discomfort,
				prolonged anorexia, etc.) to alleviate
				the irritation and pain associated
				with the surgery. From past
				experience animals□
				recover from the surgery and return
				to normal behaviors (feeding, etc)
				within 12-24 h. In addition to normal
				food and water, moist food will be
				provided in a petri dish on the floor
				of the cage during first 24 hours to
				facilitate easy access to water and
				food and prevent dehydration. Mice
				will be assessed daily for hydration
				and signs of distress. If required 1cc
				fluids will be given SQ in
				consultation with the veterinarian.
				NSAIDs or other antiinflammatory
				drugs will not be administered to
			72 HOUR	these animals at any point in the
			POST-OP	study as they will interfere with the
	Cheeran.		ANALGESIA	immunological parameters that are
1906-37124A	Maxim	Mice	POLICY	central to this study (Gomaa, S.
	Griffith.		EUTHANASIA	All staff have been trained in and
1906-37113A	Thomas	Mice	METHOD	are competent at cervical
				An IACUC exception is required for
				non-administration of analgesics
				after surgery, but is not required for
				tail snips in other situations where
			72 HOUR	bone is not cut. In this scenario
			POST-OP	zebrafish tails are comprised of
		Fish (Zebra		cartilage and soft tissues (like the tin
1906-371114	Lund Troy	fish)	POLICY	of a tail of a pre-weaping mouse
1000-07111A	Earla, Hoy	1907		or a tan or a pro-wearing mouse
				We request not to give analgesics
-------------	-----------------	----------------	------------	---------------------------------------
				because we need for hyperalgesia to
				fully develop and analgesics may
				interfere with this process.
				Methylene blue may provide
				analgesia in experimental animals.
				If animals are in excessive pain
				(vocalization, restlessness) and lose
			72 HOUR	10% of its weight, it will be
			POST-OP	euthanized. Based on our
			ANALGESIA	experience and available data,
1905-37106A	Banik, Ratan	Rat	POLICY	animals tolerate well this surgical
			EUTHANASIA	Rats are under minimal stress when
1905-37106A	Banik, Ratan	Rat	METHOD	injected for overdose.
	Munderloh,		SOCIAL	
1905-37105A	Ulrike	Hamster, Llama	HOUSING	see protocol for details
				As described in procedure, goal is to
				evaluated autologous stem cells in
				vascular graft after in vitro
				differentiation. In Vitro harvest and
				differentation take upto 2 weeks.
				The fat is harvested from each
				animal, isolated, and coated on
				graft's lumen surface prior to being
				implanted back in the same animal.
				Hence this require two procedures
				on each animal. (Adipose Fat
				Harvest)
				As described in study design,
				animals are implanted with
		Sheep	MULTIPLE	engineered graft coated with
1905-37103A	Bianco, Richard	(Biomedical)	SURGERY	autologous stem cells. To evaluate

	Nonhuman		Animals have previously been instrumented with a central vascular access port. The placement of a vascular access port is considered a minor surgical procedure (peripheral limb, not entering a major body cavity), essentially similar to routine catheter placement with the notable exception that the access site is placed underneath the skin. VAPs improve safety, and increase animal comfort when intermittent or recurrent venous access is required by avoiding painful 'catheter starts'. VAPs also allow NHPs to participate fully in enrichment programs and social group activities without the restrictions inherent in conventional approaches. This procedure is required to infuse the test chemical ablation under investigation in this study and is
1905-37094A	(Macaques)	SURGERY	mimic the approach intended in

<ul> <li>weight loss; a body condition score of less than or equal to 2 (based on Uliman-Culier 1999), primary flank solid tumor tissue measured by calipers to be of excessive size (&gt;2om length in any dimension) or impairing mobility, any health problems refractory to medical intervention such as labored breating; or on recommendation of the veterinary staff. □</li> <li>Swollen tissue, but palpably fluid-containing, either undermeath or adjoining solid tumor tissue will not be considered solid tissue in immunocompetent mice bearing tumors as this study is investigating an immunotherapy where enhanced infiltrates and edema concurrent with necrosis are features of enhanced T-cell activity described as pseudoprogression (Journal of Clinical Oncology, Vol 33, No 31 (November 1), 2015; pp 3541-3543). See http://jco.ascopubs.org/contentworland and provide and provide and the second trading and reports including impaired gait, labored breathing, and recommendation of veterinary</li> <li>1905-37092A Ondrey, Frank Mice</li> <li>TUMOR ENDPOINT CRITERIA</li> <li>Ondrey, Frank Mice</li> <li>TUMOR ENDPOINT Index and the pharmaceutical-grade breathing, and recommendation of veterinary</li> <li>We will use pharmaceutical-grade breathing, and recommendation of veterinary</li> <li>NON- Brown and the pharmaceutical-grade breathing and container. (General preparation)</li> <li>The pharmaceutical-grade pancuronium Bromide does not available, thus, non-pharmaceutical available, and container. (General preparation)</li> <li>The pharmaceutical-grade pancuronium Will be grade acuely using sterilized saline and container. (General preparation)</li> <li>The pharmaceutical-grade pancuronium Will be used.</li> </ul>					Endpoint criteria include >20%
1905-37092A       Ondrey, Frank       Mice       CRITERIA         1906-37092A       Ondrey, Frank       Mice       CRITERIA					weight loss a body condition score
<ul> <li>Uliman-Cultere 1999), primary flank solid tumor tissue measured by calipers to be of excessive size (&gt;2cm length in any dimension) or impairing mobility; any health problems refractory to medical intervention such as labored breathing; or on recommendation of the veterinary staff. U</li> <li>U</li> <li>Swollen tissue, but palpably fluid-containing, either undermeath or adjoining solid tumor tissue will not be considered solid tissue in immunocompetent mice bearing tumors as this study is investigating an immunotherapy where enhanced infiltrates and dedma concurrent with necrosis are features of enhanced T-cell activity described as pseudoprogression (Journal of Clinical Oncology, Vol 33, No 31 (November 1), 2015: pp 3541-3563). See http://jco.ascopubs.org/content/33/3 1/3541.1ul. Mice bearing swolen tissue will still be subject to all other endpoints including impaired gait, labored breathing, and recommendation of veterinary</li> <li>1905-37092A Ondrey, Frank Mice</li> <li>TUMOR</li> <li>ENDPOINT</li> <li>TUMOR tissue will sub be subject to all other endpoints including impaired gait, labored breathing, and recommendation of veterinary</li> <li>1905-arogize in of veterinary</li> <li>We will use pharmaceutical-grade pharucronium Will be prepared acutely using sterilized saline and container. (General preparation)</li> <li>The pharmaceutical-grade substance is not available. In that case, the non-pharmaceutical-grade pancuronium Bromide where or the pharmaceutical-grade substance is not available. In that case, the non-pharmaceutical-grade substance is not available. In the cose so ta available, thus, non-pharmaceutical compound will be used.</li> </ul>					of less than or equal to 2 (based on
1905-37092A       Ondrey, Frank       Mice       TUMOR CRITERIA       We will use pharmaceutical grade pharmaceutical grade south the veterinary staft.         1905-37092A       Ondrey, Frank       Mice       CRITERIA         1905-37092A       Ondrey, Frank       Mice       We will use pharmaceutical grade pharmaceutical grade south the pharmaceutical grade pharmaceutical grade south the pharmaceutical grade pharmaceutical grade pharmaceutical grade south the pharmaceutical grade pharmaceutical grade					Illiman-Cullere 1999); primary flank
1905-37092A       Ondrey, Frank       Mice       TUMOR         1905-37092A       Ondrey, Frank       Mice       We will use pharmaceutical-grade phancuronium Bromide whenever it is available, the non-pharmaceutical-grade phancuronium Bromide whenever it is not available, the non-pharmaceutical-grade acute/y using sterilized saline and container. (General prepared) acute/y using sterilized saline and container. (General prepared)					solid tumor tissue measured by
1905-37092A       Ondrey, Frank       Mice       TUMOR         1905-37092A       Ondrey, Frank       Mice       RITERIA         1905-37092A       Ondrey, Frank       Mice       We will use pharmaceutical grade paractronium will be used into available, in that case, the non-pharmaceutical grade paractronium will be repared acutely using sterilized saline and containing, and recommendation of veterinary					caliners to be of excessive size
1905-37092A       Ondrey, Frank       Mice       TUMOR ENDPOINT       Endpaintering of endersing and company and endpaintering of endersing and company and endpaintering of endp					(>2cm longth in any dimonsion) or
1905-37092A       Ondrey, Frank       Mice       CRITERIA       We will use pharmaceutical-grade paracuronium Bromide whenever it is available, the non-pharmaceutical-grade paracuronium will be used.         1905-37092A       Ondrey, Frank       Mice       CRITERIA					(22011 length in any dimension) of
1905-37092A       Ondrey, Frank       Mice       TUMOR       endpoints is lenacutical grade pancuronium Bromide whenever it is available, the non-pharmaceutical-grade pancuronium Will be prepared acutely using sterilized saline and containing in the sectional prepared acutely using sterilized saline and containing in the sectical grade pancuronium will be prepared acutely using sterilized saline and containing in the sectical grade pancuronium will be prepared acutely using sterilized saline and containing in the sectical grade pancuronium will be prepared acutely using sterilized saline and containing in the sectical grade pancuronium will be prepared acutely using sterilized saline and container. (General preparation)         1905-37092A       NON-       Prevention Such as labored prepared acutely using sterilized saline and container. (General preparation)         1905-37092A       NON-       Prevention Such as the sectical grade pancuronium will be prepared acutely using sterilized saline and container. (General preparation)					problems refractory to medical
1905-37092A       Ondrey, Frank       Mice       CRITERIA       We will use pharmaceutical-grade pancuronium Bromide does not available, thus, non-pharmaceutical-grade Pancuronium Bromide does not available, thus, non-pharmaceutical grade Pancuronium Bromide does not available, thus, non-pharmaceutical-grade Pancuronium Bromide does not availab					problems remaciony to medical
1905-37092A       Ondrey, Frank       Mice       TUMOR ENDPOINT       Bits explanable, the on-pharmaceutical-grade pancuronium Bits explanable, the on-pharmaceutical- grade         1905-37092A       Ondrey, Frank       Mice       Wice       We will use pharmaceutical-grade pancuronium With paradelical grade pancuronium With paramaceutical-grade pancuronium With paradelical grade pancuronium With paramaceutical-grade pancuronium With paradelical grade pancuronium With paradelical grade pancuronium With paramaceutical-grade pancuronium With paramaceutical-grade pancuronium With paradelical grade pancuronium With paramaceutical-grade pancuronium Bits parade pancuronium Bits parade pancur					hreathing: or on recommondation of
1905-37092A       Ondrey, Frank       Mice       Clinical Contention of veterinary statility of the program of th					
1905-37092A       Ondrey, Frank       Mice       CRITERIA       We will use pharmaceutical-grade pancuronium Bromide whenever it is available. In that case, the non-pharmaceutical grade pancuronium Bromide values the pharmaceutical grade container. (General preparation)         1905-37092A       Ondrey, Frank       Mice       RITERIA       We will use pharmaceutical-grade pancuronium Bromide whenever it is available. In that case, the non-pharmaceutical-grade pancuronium Bromide values and container. (General preparation)         1905-37092A       NON-PHARMACAUT       International preparation)       The pharmaceutical-grade pancuronium Bromide does not available. In that case, the non-pharmaceutical grade pancuronium Bromide solubatance is not available. In that case, the non-pharmaceutical grade pancuronium Bromide does not available. In that case, the non-pharmaceutical grade pancuronium Bromide does not available. In that case, the non-pharmaceutical grade pancuronium Bromide does not available. In that case, the non-pharmaceutical-grade pancuronium Bromide does not available. In that case, the non-pharmaceutical-grade pancuronium Bromide does not available. In that case, the non-pharmaceutical-grade pancuronium Bromide does not available. In that case, the non-pharmaceutical-grade pancuronium Bromide does not available. In that case, the non-pharmaceutical-grade pancuronium Bromide does not available. In that case, the non-pharmaceutical-grade pancuronium Bromide does not available. In that case, the non-pharmaceutical-grade pancuronium Bromide does not available. In that case, the non-pharmaceutical-grade pancuronium Bromide does not available. In that case, the non-pharmaceutical-grade pancuronium Bromide does not available. In that case the non-pharmaceutical-grade pancuronium Bromide does not available. In that case pancuro					
1905-37092A       Ondrey, Frank       Mice       CRITERIA       See Pharmaceutical-grade panceutical grade					L. Our all any time units and an also also be fluid.
1905-37092A       Ondrey, Frank       Mice       Crittering and impact of the pharmaceutical-grade pancuronium Bromide does not available, thus, non-pharmaceutical grade pancuronium Bromide does not available, thu					Swollen tissue, but parpapiy huid-
adjoining Solid tumor tissue will not be considered solid tissue in immunocompetent mice bearing tumors as this study is investigating an immunotherapy where enhanced infiltrates and edema concurrent with necrosis are features of enhanced T-cell activity described as pseudoprogression (Journal of Clinical Oncology, Vol 33, No 31 (November 1), 2015: pp 3541-3543). See          1905-37092A       Ondrey, Frank       Mice       CRITERIA       Roore beam including impaired gait, labored breathing, and recommendation of veterinary         1905-37092A       Ondrey, Frank       Mice       CRITERIA       We will use pharmaceutical-grade endpoints including impaired gait, labored breathing, and recommendation of veterinary         1905-37092A       Ondrey, Frank       Mice       CRITERIA       We will use pharmaceutical-grade endpoints including impaired gait, labored breathing, and recommendation of veterinary         1905-37092A       Ondrey, Frank       Mice       CRITERIA       We will use pharmaceutical-grade endpoints including impaired gait, labored breathing, and recommendation of veterinary         1905-37092A       Ondrey, Frank       Mice       CRITERIA       We will use pharmaceutical-grade endpoints including impaired gait, labored breathing, and recommendation of veterinary         1905-37092A       Ondrey, Frank       Mice       CRITERIA       Tomony and recommendation of veterinary         1905-37092A       Ondrey, Frank       Mice       CRITERIA       Tomony and will only be used when the pharmaceutical-grade pancuronium will be prepared acutely us					containing, either underneath or
1905-37092A       Ondrey, Frank       Mice       TUMOR       immunotherapy where enhanced infiltrates and edema concurrent with necrosis are features of enhanced T-cell activity described as pseudoprogression (Journal of Clinical Oncology, Vol 33, No 31 (November 1), 2015: pp 3541-3543). See         1905-37092A       Ondrey, Frank       Mice       TUMOR ENDPOINT Issue and the second					adjoining solid tumor tissue will not be considered solid tissue in
1905-37092AOndrey, FrankMiceCRITERIAWe will use pharmaceutical-grade pancuronium Bromide does not available, thus, non-pharmaceutical orantare. (General preparation)1905-37092AOndrey, FrankMiceRiftering Pancuronium Bromide does not available, thus, non-pharmaceutical orantare. (General preparation)1905-37092AOndrey, FrankMiceRiftering Pancuronium Bromide does not available, thus, non-pharmaceutical orantare.1905-37092AOndrey, FrankMiceRiftering Pancuronium Bromide does not available, thus, non-pharmaceutical oranged compound will be used.					immunocompetent mice bearing
1905-37092A       Ondrey, Frank       Mice       CRITERIA       an immunotherapy where enhanced infiltrates and edema concurrent with necrosis are features of enhanced T-cell activity described as pseudoprogression (Journal of Clinical Oncology, Vol 33, No 31 (November 1), 2015: pp 3541-3543). See         1905-37092A       Ondrey, Frank       Mice       CRITERIA       Nice acoupts.org/content/33/3 1/3541,full. Mice bearing swollen tissue will still be subject to all other endpoints including impaired gait, labored breathing, and recommendation of veterinary         1905-37092A       Ondrey, Frank       Mice       CRITERIA       We will use pharmaceutical-grade Pancuronium Bromide whenever it is available, the non-pharmaceutical grade pancuronium will be used when the pharmaceutical grade pancuronium will be prepared acutely using sterilized saline and container. (General preparation)         NON-       PHARMACAUT ICAL GRADE       NON-pharmaceutical-grade Pancuronium Bromide does not available, thus, non-pharmaceutical compound will be used.					tumors as this study is investigating
1905-37092A       Ondrey, Frank       Mice       CRITERIA       with necrosis are features of enhanced T-cell activity described as pseudoprogression (Journal of Clinical Oncology, Vol 33, No 31 (November 1), 2015: pp 3541-3543). See         1905-37092A       Ondrey, Frank       Mice       CRITERIA       recommendation of veterinary         1905-37092A       Ondrey, Frank       Mice       CRITERIA       we will use pharmaceutical-grade Pancuronium Bromide whenever it is available, the non-pharmaceutical grade pancuronium will only be used when the pharmaceutical grade pancuronium will be prepared acutely using sterilized saline and container. (General preparation)         NON-       PHARMACAUT       ICAL GRADE					an immunotherapy where enhanced
<ul> <li>with necrosis are features of enhanced T-cell activity described as pseudoprogression (Journal of Clinical Oncology, Vol 33, No 31 (November 1), 2015: pp 3541-3543). See http://jco.ascopubs.org/content/33/3 1/3541.full. Mice bearing swollen tissue will still be subject to all other endpoints including impaired gait, labored breathing, and recommendation of veterinary</li> <li>1905-37092A</li> <li>Ondrey, Frank</li> <li>Mice</li> <li>CRITERIA</li> <li>We will use pharmaceutical-grade Pancuronium Bromide whenever it is available, the non-pharmaceutical compound will only be used when the pharmaceutical-grade substance is not available. In that case, the non-pharmaceutical grade pancuronium will be prepared acutely using sterilized saline and container. (General preparation)</li> <li>The pharmaceutical-grade Pancuronium Bromide does not available, thus, non-pharmaceutical compound will be used.</li> </ul>					infiltrates and edema concurrent
1905-37092AOndrey, FrankMiceTUMOR ENDPOINT CRITERIAenhanced T-cell activity described as pseudoprogression (Journal of Clinical Oncology, Vol 33, No 31 (November 1), 2015: pp 3541-3543). See http://jco.ascopubs.org/content/33/3 1/3541.full. Mice bearing swollen tissue will still be subject to all other endpoints including impaired gait, labored breathing, and recommendation of veterinary1905-37092AOndrey, FrankMiceCRITERIAWe will use pharmaceutical-grade Pancuronium Bromide whenever it is available, the non-pharmaceutical compound will only be used when the pharmaceutical grade pancuronium will be prepared acutely using sterilized saline and container. (General preparation)NON- PHARMACAUT ICAL GRADENON- PHARMACAUTThe pharmaceutical-grade Pancuronium Bromide does not available, thus, non-pharmaceutical compound will be used.					with necrosis are features of
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Clinical Oncology, Vol 33, No 31 (November 1), 2015: pp 3541-3543). See http://jco.ascopubs.org/content/33/3 1/3541.full. Mice bearing swollen tissue will still be subject to all other endpoints including impaired gait, labored breathing, and recommendation of veterinary We will use pharmaceutical-grade Pancuronium Bromide whenever it is available, the non-pharmaceutical grade pancuronium will be used when the pharmaceutical-grade substance is not available. In that case, the non-pharmaceutical grade pancuronium will be prepared acutely using sterilized saline and container. (General prepared) acutely using sterilized saline and container.					as pseudoprogression (Journal of
1905-37092A       Ondrey, Frank       Mice       TUMOR       endpoints including impaired gait, labored breathing, and recommendation of veterinary         1905-37092A       Ondrey, Frank       Mice       CRITERIA       We will use pharmaceutical-grade Pancuronium Bromide whenever it is available, the non-pharmaceutical compound will only be used when the pharmaceutical grade pancuronium will be prepared acutely using sterilized saline and container. (General preparation)         NON-       PHARMACAUT ICAL GRADE       NON-pHARMACAUT ICAL GRADE					Clinical Oncology, Vol 33, No 31
1905-37092A       Ondrey, Frank       Mice       TUMOR ENDPOINT CRITERIA       Http://jco.ascopubs.org/content/33/3 1/3541.full.         1905-37092A       Ondrey, Frank       Mice       TUMOR ENDPOINT CRITERIA       endpoints including impaired gait, labored breathing, and recommendation of veterinary         1905-37092A       Ondrey, Frank       Mice       We will use pharmaceutical-grade Pancuronium Bromide whenever it is available, the non-pharmaceutical compound will only be used when the pharmaceutical-grade pancuronium will be prepared acutely using sterilized saline and container. (General preparation)         NON- PHARMACAUT ICAL GRADE       NON- PHARMACAUT ICAL GRADE       The pharmaceutical-grade pancuronium Bromide does not available, thus, non-pharmaceutical compound will be used.					(November 1), 2015: pp 3541-3543).
1905-37092AOndrey, FrankMiceTUMOR ENDPOINT CRITERIAhttp://jco.ascopubs.org/content/33/3 1/3541.full. Mice bearing swollen tissue will still be subject to all other endpoints including impaired gait, labored breathing, and recommendation of veterinary1905-37092AOndrey, FrankMiceCRITERIAWe will use pharmaceutical-grade Pancuronium Bromide whenever it is available, the non-pharmaceutical compound will only be used when the pharmaceutical grade pancuronium will be prepared acutely using sterilized saline and container. (General preparation)NON- PHARMACAUT ICAL GRADENON- PHARMACAUTThe pharmaceutical-grade Pancuronium Bromide does not available, thus, non-pharmaceutical compound will be used.					See
1905-37092A       Ondrey, Frank       Mice       TUMOR       tissue will still be subject to all other endpoints including impaired gait, labored breathing, and recommendation of veterinary         1905-37092A       Ondrey, Frank       Mice       CRITERIA       We will use pharmaceutical-grade Pancuronium Bromide whenever it is available, the non-pharmaceutical compound will only be used when the pharmaceutical-grade substance is not available. In that case, the non-pharmaceutical grade pancuronium will be prepared acutely using sterilized saline and container. (General preparation)         NON-       PHARMACAUT ICAL GRADE       The pharmaceutical-grade compound will be used.					http://ico.ascopubs.org/content/33/3
1905-37092A       Ondrey, Frank       Mice       TUMOR ENDPOINT CRITERIA       tissue will still be subject to all other endpoints including impaired gait, labored breathing, and recommendation of veterinary         1905-37092A       Ondrey, Frank       Mice       We will use pharmaceutical-grade Pancuronium Bromide whenever it is available, the non-pharmaceutical compound will only be used when the pharmaceutical-grade substance is not available. In that case, the non-pharmaceutical grade pancuronium will be prepared acutely using sterilized saline and container. (General preparation)         NON- PHARMACAUT ICAL GRADE       NON- PHARMACAUT ICAL GRADE       The pharmaceutical-grade Pancuronium Bromide does not available, thus, non-pharmaceutical compound will be used.					1/3541.full. Mice bearing swollen
1905-37092AOndrey, FrankMiceTUMOR ENDPOINT CRITERIAendpoints including impaired gait, labored breathing, and recommendation of veterinary1905-37092AOndrey, FrankMiceCRITERIAendpoints including impaired gait, labored breathing, and recommendation of veterinary1905-37092AOndrey, FrankMiceWe will use pharmaceutical-grade Pancuronium Bromide whenever it is available, the non-pharmaceutical compound will only be used when the pharmaceutical-grade substance is not available. In that case, the non-pharmaceutical grade pancuronium will be prepared acutely using sterilized saline and container. (General preparation)NON- PHARMACAUT ICAL GRADENON- PHARMACAUTThe pharmaceutical-grade Pancuronium Bromide does not available, thus, non-pharmaceutical compound will be used.					tissue will still be subject to all other
1905-37092A       Ondrey, Frank       Mice       ENDPOINT CRITERIA       Iabored breathing, and recommendation of veterinary         1905-37092A       Ondrey, Frank       Mice       CRITERIA       We will use pharmaceutical-grade Pancuronium Bromide whenever it is available, the non-pharmaceutical compound will only be used when the pharmaceutical-grade substance is not available. In that case, the non-pharmaceutical grade pancuronium will be prepared acutely using sterilized saline and container. (General preparation)         NON- PHARMACAUT ICAL GRADE       NON- PHARMACAUT       The pharmaceutical-grade Pancuronium Bromide does not available, thus, non-pharmaceutical				TUMOR	endpoints including impaired gait.
1905-37092A       Ondrey, Frank       Mice       CRITERIA       recommendation of veterinary         1905-37092A       Ondrey, Frank       Mice       CRITERIA       recommendation of veterinary         We will use pharmaceutical-grade       Pancuronium Bromide whenever it is available, the non-pharmaceutical compound will only be used when the pharmaceutical-grade substance is not available. In that case, the non-pharmaceutical grade pancuronium will be prepared acutely using sterilized saline and container. (General preparation)         NON-       PHARMACAUT ICAL GRADE       The pharmaceutical-grade compound will be used.				ENDPOINT	labored breathing, and
NON- PHARMACAUT ICAL GRADE	1905-37092A	Ondrev Frank	Mice	CRITERIA	recommendation of veterinary
NON- PHARMACAUT ICAL GRADENON- PHARMACAUT ICAL GRADEWe will use pharmaceutical-grade Pancuronium Bromide whenever it is available, the non-pharmaceutical compound will only be used when the pharmaceutical-grade substance is not available. In that case, the non-pharmaceutical grade pancuronium will be prepared acutely using sterilized saline and container. (General preparation)NON- PHARMACAUT ICAL GRADEThe pharmaceutical-grade Pancuronium Bromide does not available, thus, non-pharmaceutical compound will be used.	1000 01 002,1	onaroy, marite	Miloo	ORTERN	recommendation of votermary
We will use pharmaceutical-grade Pancuronium Bromide whenever it is available, the non-pharmaceutical compound will only be used when the pharmaceutical-grade substance is not available. In that case, the non-pharmaceutical grade pancuronium will be prepared acutely using sterilized saline and container. (General preparation)NON- PHARMACAUT ICAL GRADEThe pharmaceutical-grade Pancuronium Bromide does not available, thus, non-pharmaceutical compound will be used.					
NON-         NON-         NON-         NON-         NON-         NON-         PARCURATE         NON-         PARCURATE         NON-         PARCURATE         NON-         PHARMACAUT         ICAL GRADE					We will use pharmaceutical-grade
NON- NON- PHARMACAUT ICAL GRADE					Pancuropium Bromide whenever it
NON- NON- PHARMACAUT ICAL GRADE					is available, the pop-pharmaceutical
NON- PHARMACAUT ICAL GRADE					a a manual will only be used when
NON- PHARMACAUT ICAL GRADE					the pharmaceutical grade substance
NON- PHARMACAUT ICAL GRADE					is pot available. In that appende
NON- PHARMACAUT ICAL GRADE					non pharmagautical grade
NON- PHARMACAUT ICAL GRADE					
Acutely using sterilized saline and container. (General preparation)         The pharmaceutical-grade         Pancuronium Bromide does not         available, thus, non-pharmaceutical         PHARMACAUT         ICAL GRADE					pancuronium will be prepared
NON- PHARMACAUT ICAL GRADE					acutely using sterilized saline and
Image: Non- Pharmaceutical-grade Pancuronium Bromide does not available, thus, non-pharmaceutical compound will be used.ICAL GRADE					container. (General preparation)
NON- PAncuronium Bromide does not available, thus, non-pharmaceutical PHARMACAUT ICAL GRADE					The pharmaceutical-grade
NON- PHARMACAUT compound will be used. ICAL GRADE					Pancuronium Bromide does not
PHARMACAUT compound will be used. ICAL GRADE				NON-	available, thus, non-pharmaceutical
ICAL GRADE				PHARMACAUT	compound will be used.
				ICAL GRADE	
1905-37090A Chen, Wei Cat COMPOUNDS The non-pharmaceutical grade	1905-37090A	Chen, Wei	Cat	COMPOUNDS	The non-pharmaceutical grade

				In the past we have used
				nharmaceutical grade pancuronium
				Diomide from Sicol, Sicol
				Pharmaceuticals, Inc., Irvine, CA.
				However, this company was
				acquired by another company which
				no longer manufactures this product.
				We have been unable to find other
				sources for procuring
				pharmaceutical grade pancuronium
				bromide. Therefore, we use
				pancuronium bromide from Sigma-
				Aldrich, St. Louis, MO (Cat. No.
				P1918).
			NON-	Pancuronium bromide will be
			PHARMACAUT	dissolved in tissue culture grade
			ICAL GRADE	water at a concentration of 2 mg/ml.
1905-37065A	Rao, Savita	Mice	COMPOUNDS	The solution will be sterile filtered,
				We must perform multiple biopsies
				in order to track hair growth related
				marker protein expression between
				control animals and those receiving
			MULTIPLE	inhibitor over time. These samples
1905-37062A	Dong, Zigang	Mice	SURGERY	are superficial and should heal
				The intention of the spared nerve
				injury is to induce a state simulating
				the hyperalgesia experienced in
			72 HOUR	neuropathic pain. Administration of
			POST-OP	analgesics would be likely to alter
			ANALGESIA	the course of hyperalgesia
1905-37059A	Wilcox, George	Rat, Mice	POLICY	development, defeating the goal of
				Cervical dislocation may be
				indicated at times for emergency
				humane euthanasia where provision
			EUTHANASIA	of prior isoflurane anesthesia is
1905-37059A	Wilcox, George	Rat, Mice	METHOD	either not possible or would prolong
			SOCIAL	
1905-37059A	Wilcox, George	Rat, Mice	HOUSING	see protocol for details
			72 HOUR	Fish will be released in the wild
			POST-OP	making detailed monitoring and
	Sorensen,		ANALGESIA	administration of analgesics
1905-37046A	Peter	Fish (Other)	POLICY	impossible□
			ANESTHESIA,	
			SURGERY,	
			AND POST-	Fish will be released in the wild. The
			PROCEDURAL	movement of all fish will then be
	Sorensen,		RECORDKEEP	monitored as part of the experiment.
1905-37046A	Peter	Fish (Other)	ING	(Tagging)

				It is possible that single housing will
				be required if: 1) there is only one
		Pia		animal in the litter 2) littermates die
		(Biomedical).	ENVIRONMEN	or are euthanized 3) single housing
		Pia	TAI	is needed to minimize transmission
1905-37039A	Garry Mary	(Biomedical)		of illness among piglets or 4) if
1000 070007	Ourry, Mary	Pig		It is possible that we will need an
		(Biomedical)		exception to social bousing for the
		Pig	SOCIAL	reasons stated in 17B. If possible
1905-370394	Garny Many	(Biomedical)		however, we will house socially
1000 070007	Carry, Mary	(Biorriediodi)	1000110	NA: these are larval fish. (Non-
				survival surgery)
				NA: those are larval fish. (Coll
				Torgeted Lease Ablation
				rargeted Laser Abiation)
				NA: these are larged fich
				INA, these are larval lish.
				(Electrophysiology)
			ANESTHESIA,	
			SURGERY,	NA; these are larval fish. (In vivo
			AND POST-	neuronal labeling)
		Fish (Zebra	PROCEDURAL	
		fish), zebrafish	RECORDKEEP	NA; Tricaine used to euthanize fish.
1905-37035A	Masino, Mark	embryos/larvae	ING	(Fixation for IHC and ISH)
				Sedation is used for training the
				cervical dislocation technique.
				Isoflurane is used at 3-4X MAC and
				then dislocation performed under
			EUTHANASIA	anesthesia. In accordance with
1905-37029A	Finger, Erik	Mice, Rat	METHOD	changes in IACUC policy,
				In order to validate tolerance in long
				term graft survivors, a small subset
				of long term survivors will have a
				second skin grafting or islet
				transplant performed. This is to
				document donor specific tolerance
				and is an crucial immunologic
				outcome Additionally some islet
				transplant recipients will have a
				unilateral perfectomy in order to
				document that long term graft
1005 370284	Einger Erik	Mico		function is due to the graft and not
1303-37020A		WILLE	JUNGERT	No incisions are made. Inculin
				notificate naced by suboutanceus
				injection (with 14 a tracher). No
				Injection (with 14 g trochar). NO
1005 070004		Minn		pain medication has been required
1905-37028A	i⊢inger, Erik	INICE	POLICY	and pellets are well tolerated in

				Sedation is used for training the
		1		cervical dislocation technique.
		1		Isoflurane is used at 3-4X MAC and
		1		then dislocation performed under
		1	EUTHANASIA	anesthesia. In accordance with
1905-37028A	Finger, Erik	Mice	METHOD	changes in IACUC policy,
	Gallaher,		SOCIAL	
1905-37019A	Daniel	Rat	HOUSING	see protocol for details
			ENVIRONMEN	
	Faulk,	1	TAL	
1905-37011A	Christopher	Mice	ENRICHMENT	see protocol for details
	Faulk,		SOCIAL	· · · ·
1905-37011A	Christopher	Mice	HOUSING	see protocol for details
		Fish (Other),		
		Fish (Zebra	NON-	
		fish) Reptile	PHARMACAUT	
		(Other) Fish	ICAL GRADE	
1904-37007A	Olson Erin	(Other)	COMPOUNDS	see protocol for details
100101001		Amphibian		Furthanized animals will be
		(Other) Rodent		preserved as scientific museum
		(Other - Non-		specimen Additional methods to
		USDA) Bird		ensure euthanasia (e d
		(Other) Fish		decapitation cervical dislocation)
		(Other) Rentile		will destroy the future scientific
		(Other), Repuie		while of animals as museum
		(Uner), Uner		
		(NOTI-USDA),		specimens.
1004 270054	Kazak Kappoth	Ind-Sizeu anu		Descritation and/or convical
1904-37000A	Kozak, Kenneun	large manimais		Decapitation and/or cervical
		1		
		1		
4004.00000	Arrai Obiori			N1/A
1904-36992A	Arai, Shiori	Dog		N/A
		1		We are testing empryonic lethality of
				gene overexpression. Typically this
	Concernal di			Will be assessed before 72 hrs post
1001000054	Smanski,	tisn), ⊢isn		rertilization, but in rare cases we
1904-36985A	Michael	(Other)		might need to look for lethality in
			NON-	
	_ * ************	Fish (∠ebra		
	Smanski,	fish), Fish	ICAL GRADE	
1904-36985A	Michael	(Other)	COMPOUNDS	see protocol for details

				we sometimes add osmotic minipumps to other more invasive procedures, such as cardiac pressure overload or cardiac ischemic injury. The goal would be to add thymidine anologs such as BrdU or EdU to perform pulse chase experiments of proliferative cells to measure regeneration. Alternatively, we could add neurohormonal
1904-36978A	van Berlo, Jop	Mice, Rat	MULTIPLE SURGERY	stimulation in strains of mice that are especially resistant to cardiac injury. (Cardiac Ischemic Injury) This is a minimally invasive procedure that we sometimes add to other more invasive procedures, such as cardiac pressure overload or cardiac ischemic injury. The goal would be to add thymidine anologs
1904-36978A	van Berlo, Jop	Mice, Rat	EUTHANASIA METHOD	We will only perform decapitation using sharp scissors in neonatal pups younger than post-natal day 7 We will only use this method of euthanasia in newborn pups younger than post-natal day 7, where we will use sharp scissors to quickly
1904-36967A	Li, Yuzhi	Pig (Agricultural)	72 HOUR POST-OP ANALGESIA POLICY	Castration is a routine practice for piglet processing after birth. Currently, castration of piglets within the first week of birth without administering analgesia is the
1904-36960A	O'Connell, Timothy	Mice	SOCIAL HOUSING	see protocol for details

1904-36947A	Toth, Ferenc	Goat	EUTHANASIA METHOD	Barbiturate overdose will be performed by an experienced investigator with a single venipuncture. Administering a sedative before the barbiturate overdose would only prolong the
1904-36948A		Nonhuman Primate (Macaques)	MULTIPLE SURGERY	Animals have previously been instrumented with a central vascular access port. The placement of a vascular access port is considered a minor surgical procedure (peripheral limb, not entering a major body cavity), essentially similar to routine catheter placement with the notable exception that the access site is placed underneath the skin. VAPs improve safety, and increase animal comfort when intermittent or recurrent venous access is required by avoiding painful 'catheter starts'. VAPs also allow NHPs to participate fully in enrichment programs and social group activities without the restrictions inherent in conventional approaches. This procedure is required to implant the microcapsules and is designed

				The main objectives for all the studies need intracranial EEG monitoring and the device working. If a device fails, then the study would end for that dog and a new dog obtained, the supply of research dogs with spontaneous seizures is not large, and to also reduce numbers of dogs we plan to re- implant the device if needed, Under the previous protocol this was permitted and we did replace the device in 2 dogs (one now
				deceased, the other on the current protocol). Going forward, as some of these
1904-36933A	Patterson, Ned	Dog, Dog, Dog		dogs are and will be here for years, the PI will consult with the RAR vet about any concurrent health conditions, and the current seizure status of the dog before going ahead with a second intracranial surgery, and not proceeding if the concurrent conditions are deemed to be serious enough that the dog may not do well with the surgery and/or may not survive long enough post surgery to be valuable to the results. Specifically No second intracranial surgery will be performed without consultation with and approval by the RAR area vet, with their
1904-36928A	Pluhar, Liz	Dog, Doa	EUTHANASIA METHOD	It is not standard protocol to sedate pet dogs prior to euthanasia.
1903-36921A	Lim, Hubert	Guinea Pig	ANESTHESIA, SURGERY, AND POST- PROCEDURAL RECORDKEEP ING	We need to be outside of our sound chamber when performing the neural recordings to avoid electrical and acoustic noise contamination. Since our protocols usually requires up to 30 minutes of recordings for each session, we need to be able to
1903-36921A	Lim, Hubert	Guinea Pig	SOCIAL HOUSING	see protocol for details

T		1	
Willette, Michelle	Chicken, Bird (Other)	ANESTHESIA, SURGERY, AND POST- PROCEDURAL RECORDKEEP ING	This lab consists of 50 birds undergoing light anesthesia for non painful procedures. No individual records will be kept. (Exam and venipuncture) This lab consists of 50 birds under
Andrade			Implantation of the construct into the
Rafael	Rabbit	SURGERY	vascularization of the graft prior to
Andrade.	Rubbit	SOCIAL	associatization of the grant phon to
Rafael	Rabbit	HOUSING	see protocol for details
Asakura,		MULTIPLE	transplantation will be performed. I  After surgery is completed, animals will be cared and examined on a de bases by member of our  research staff, PI (Atsushi Asakura Dr. Shuichi Watanabe (Research Associate), Mayank Verma (Junior Scientist), or  (Undergraduate Student). They will also monitor those  animals everyday. Animal Facility staff is also monitoring regularly. The information will be transmitted by phone to PI's office and PI's laboratory. During weekend the information will be transmitted by  phone to PI's cell phone and by E- mail. Within 24 hours, the animals develop evidence(s) of distress will be performed for a terminal study. To alleviate pain during operation, we will frequently check the level of anesthesia after intraperitoneal injection of Avertin. We are going we will use a subq. injection of 5
	Willette, Michelle Andrade, Rafael Andrade, Rafael	Willette, Michelle       Chicken, Bird (Other)         Andrade, Rafael       Rabbit         Andrade, Rafael       Rabbit         Andrade, Rafael       Rabbit         Andrade, Rafael       Rabbit	Asakura,       Mice       Multiple         Michail       Mice       SURGERY, AND POST-PROCEDURAL RECORDKEEP ING         Andrade,       Rabbit       MULTIPLE SURGERY         Andrade,       Rabbit       SOCIAL HOUSING

1903-36906A	Asakura, Atsushi	Mice	NON- PHARMACAUT ICAL GRADE COMPOUNDS	Scientific justification for use of non- pharmaceutical grade Avertin: Avertin has been widely used as one of the anesthetic agents for mice. Accordingly, RAR site mentioned that "Avertin® is a non- pharmaceutical grade compound. Currently, there is no equivalent veterinary or human drug is available for experimental use of Avertin®. The highest grade equivalent chemical reagent will be
	Asakura,	a bee	SOCIAL	
1903-36906A	Atsushi	Mice	HOUSING	see protocol for details
1903-36904A	Farrar, Michael	Mice	WEEKLY WEIGHING EXCEPTION (FOOD/FLUID)	this should not affect their weight significantly. We certainly don't anticipate weightloss due to antibiotic administration. The
1903-36903A	Bereiter, David	Rat	MULTIPLE SURGERY	Exorbital gland removal and intra- CNS cannula placement will be performed concurrently. This may not qualify as multiple survival surgeries as there is no inter-
1903-36903A	Bereiter, David	Rat	72 HOUR POST-OP ANALGESIA POLICY	We have received previous permission (7.22.13) from IACUC that a single dose of ketoprofen is sufficient for the minor surgeries we propose. We will change the protocol to use carprofen instead. (Exorbital Gland removal) We had received permission (7.22.13) from IACUC to use

			NON- PHARMACAUT ICAL GRADE	Pharmaceutical grade formulations do not exist. Urethane is necessary because barbiturates often lead to unreliable OOemg activity. Please note that all animals are checked for response to toe pinch at 5 minute intervals to assess adequate anesthesia. (Thoracotomy) Pharmaceutical grade formulations do not exist. Barbiturates are not used here since they cause spontaneous muscle twitching that interferes with the evoked activity. (Dorsal brainstem surface exposure (electrophysiology)) Exorbitant cost increases effectively makes pharmaceutical grade formulations unavailable. (Ultraviolet light irradiation)
1903-36903A	Bereiter David	Rat	COMPOUNDS	
1903-36900A	Bangalore Kodandaramaia h, Suhasa	Mice, Rat	MULTIPLE SURGERY	We seek exception in surgeries when a virus injection (S2) is performed. We have previously found that in cases where virus injection is immediately followed by the device implantation (see procedures S3, S4 or S5), that the virus labeling is altered by the small brain displacements induced by device insertion. Since the viruses can take weeks to express, implanting the device later, after the viruses have fully expressed, minimizes the chance of a device- related deterioration. Thus, it may be very useful for experiments in which both Subprocedures S2 and S3 (or S4 or S5) are required, that an initial surgery with just procedure S2 can be performed (e.g., the viral infusion), and then the animal fully recovered, and then, 7-120 days
	Kodandaramaia		SOCIAL	
1903-36900A	h, Suhasa	Mice, Rat	HOUSING	see protocol for details

				Chloralose does not available, thus, non-pharmaceutical compound will be used. (Varying Brain States Using Different Anesthetic Drugs) The pharmaceutical-grade Pancuronium Bromide does not available, thus, non-pharmaceutical compound will be used. (Varying Brain States Using Different Anesthetic Drugs)
			NON-	Pharmaceutical-grade pancuronium
				promide is not available.
1903-36898A	Chen. Wei	Rat. Mice	COMPOUNDS	
				Pain level C.□
				This procedure in SA3f-1 and 3g-1 may lead to Pain Level C due to ulceration.□
				ulceration is observed on the treated tumor as a result of tumor necrosis. We will observe the small ulceration up to 7 days unless continuous
				bleeding or infection is observed or reaching other euthanization criteria. We will give analgesics as described in Health and Monitoring. □ Those which show large quantity bleeding infection or deterioration
			TUMOR	of general condition, will be
	Yamamoto,		ENDPOINT	euthanized.
1903-36889A	Masato	Mice	CRITERIA	Non-drug methods for supportive
				Bi-weekly body weights have been shown to be adequate to detect differences in□
			WEEKLY	dairy calf growth in previous SROC
			WEIGHING	calf studies. (Pre- and post weaning
1002 260074	Chester-Jones,	Cow		performance and health of nursery
1903-3088/A	nugn	(Agricultural)		uarry calves when ted a proprietary
		aoldfish round	PHARMACAUT	
	Mensinger.	goby, Fish	ICAL GRADE	
1903-36856A	Allen	(Other)	COMPOUNDS	see protocol for details

		Nonhuman Primate	MULTIPLE	The placement of a vascular access port is considered a minor surgical procedure (peripheral limb, not entering a major body cavity), essentially similar to routine catheter placement with the notable exception that the access site is placed underneath the skin. VAPs improve safety, and increase animal comfort when intermittent or recurrent venous access is required by avoiding painful 'catheter starts'. VAPs also allow NHPs to participate fully in enrichment programs and social group activities without the restrictions inherent in conventional approaches. Animals may subsequently be instrumented with hepatic portal access if they are intended for enrollment on an islet transplantation protocol that delivers islets using the standard intraportal route. In this situation, instrumentation allows animals to complete all surgical manipulation prior to induction of the disease state. (Central vascular access port
				In order to achieve successful engraftment of muscle stem and progenitor cells the muscle has to undergo an injury two days before transplantation to induce an essential injury response in the tissue. Muscle injection injury will be performed as described in the
1903-36840A	Chan, Sunny	Mice	MULTIPLE SURGERY	previous section, pain and distress in the animals will be monitored for 3 days post-procedure. Because the effects of analgesic compounds on muscle regeneration are unknown, and because we want to examine
1903-36840A	Chan, Sunny	Mice	72 HOUR POST-OP ANALGESIA POLICY	Because the effects of analgesic compounds on muscle regeneration are unknown, and because we want to examine the role of inflammatory cells in skeletal muscle

			EUTHANASIA	Embryos 15 days of gestation or
1903-36840A	Chan, Sunny	Mice	METHOD	greater will be decapitated.
				Staff are well-trained on the cervical
				dislocation procedure. Training
				to onsured staff are qualified to
				porform the procedure. Conviced
				dislocation in conscious condition
				provides a means to recover tissues
				and body fluids that are
				uncontaminated by anesthesia. It
				also provides a means of obtaining
				anatomically undamaged brain
				tissue for study. Handling and
				restraint required to perform this
				technique may be stressful to
				animals The equipment used to
				perform cervical dislocation will be
				maintained in good working
				condition.
				Staffs are well-trained on
				decapitation procedure.Training
				record is maintained and updated to
				ensured staff are qualified to
				perform the procedure. Decapitation
				provides a means to recover tissues
				and body fluids that are chemically
				uncontaminated when performed
				without anesthesia. It also provides
				a means of obtaining anatomically
				undamaged brain tissue for study.
				Handling and restraint required to
1000 26921 4		Mico		perform this technique may be
1902-30631A	newman, Enc	MICE	METHOD	distressi di to animais. The
				Post-operative analgesic therapy is
				contraindicated because the
				objective of the procedure is to
			72 HOUR	determine whether the experimental
	222		POST-OP	treatment causes hypersensitivity.
	Masopust,		ANALGESIA	Provision of analgesics would inhibit
1902-36825A	David	Mice	POLICY	the neurochemical changes required
				All personnel performing cervical
	Maconuct			extremely competent to provert
1902-368254	David	Mice	METHOD	inhumane euthanasia. Their training
1002-00020A	David	MICC.		minanane eunanasia. Their rialfillig

1902-36813A		Nonhuman Primate (Macaques)	MULTIPLE SURGERY	instrumented with a central vascular access port. The placement of a vascular access port is considered a minor surgical procedure (peripheral limb, not entering a major body cavity), essentially similar to routine catheter placement with the notable exception that the access site is placed underneath the skin. VAPs improve safety, and increase animal comfort when intermittent or recurrent venous access is required by avoiding painful 'catheter starts'. VAPs also allow NHPs to participate fully in enrichment programs and social group activities without the restrictions inherent in conventional approaches. This procedure is required to infuse the test therapy under investigation
1902-36795A	Pravetoni, Marco	Mice	BLOOD COLLECTION LIMIT	Animal will be euthanized by this method and will not survive the decapitation and subsequent blood collection (Trunk blood collection
1902-36788A	Ward, John	Frog (Xenopus)	MULTIPLE SURGERY	The main reason for using survival surgery in this protocol is to limit the number of frogs used. Some frogs (25%) do not have suitable oocytes, those that do can be used multiple times with good results. NIH guidelines permit 5 survival surgeries to remove oocytes and a sixth terminal surgery. If we did
1002-367884	Ward John	Frod (Xepopus)	SOCIAL	see protocol for details
1902-36781A	Lee, Michael	Mice	ANESTHESIA, SURGERY, AND POST- PROCEDURAL RECORDKEEP ING	Once the animal is under deep anesthesia, they are euthanized within 10 minutes. (Intracardial Purfusion)
1902-36781A	Lee, Michael	Mice	HOUSING	see protocol for details
1902-36774A	Lee, Michael	Mice	ANESTHESIA, SURGERY, AND POST- PROCEDURAL RECORDKEEP ING	Once the animal is under deep anesthesia, they are euthanized within 10 minutes. (Intracardial Purfusion)

				1 Decapitation of pregnant female
				mice is performed secondary to
				CO2 based euthanasia to ensure
				2. Direct descritation of mouse number
				S. Direct decapitation of mouse pups $(D0, 4)$ is assertable presedure.
				(P0-1) is acceptable procedure.
4000 007744		Marine and		""Based on the AVMA guidelines,
1902-36774A	Lee, Michael	wice		decapitation is an acceptable form
1000 0077 11			SOCIAL	
1902-36774A	Lee, Michael	Mice	HOUSING	see protocol for details
			ANESTHESIA,	
			SURGERY,	
			AND POST-	Once the animal is under deep
			PROCEDURAL	anesthesia, they are euthanized
		01 - MA	RECORDKEEP	within 10 minutes. (Intracardial
1902-36759A	Lee, Michael	Mice	ING	Purfusion)
			SOCIAL	
1902-36759A	Lee, Michael	Mice	HOUSING	see protocol for details
				Arthroscopy is required for baseline
				and post-study treatment. This does
				involve anesthesia, but is a
				minimally invasive procedure.
				Anesthesia is used to minimize pain
	Conzemius,		MULTIPLE	and distress. Post-operative pain will
1901-36726A	Michael	Dog	SURGERY	be treated with nonsteroidal anti-
				Simultaneous with (0 time point) or
				1,3,5, or 7 days after barium
				chloride injury we will measure
				physiological function of the muscle
				in the anesthetized mouse (in vivo
				contractile analysis). This is an
				extremely minimally invasive
				procedure in which only two small
				electrodes are place subcutaneously
				around the peroneal nerve of the
				leg. Immediately after this
			MULTIPLE	measurement, while the mouse is
1901-36722A	Lowe, Dawn	Mice	SURGERY	still anesthetized the skin will be

				Because we will be examining the
				role of macrophages in skeletal
				muscle
				autophagy/damage/regeneration,
				such as that which is induced by
				barium chloride injury, it is
				necessary to restrict the use of
				analgesia unless serious negative
				consequences result. We will
				continually work with the vet staff in
				order to be sure that there is
				adequate attention paid to the extent
				of post-surgical pain. 🛛
				If an analgesic is prescribed by
				RAR, analgesics that have fewer
				anti-inflammatory effects could be
				used in these cases, such as
				sustained release buprenorphine will
				be used first. Local anesthetics such
				as lidocaine or bupivicaine (sodium
				channel blockers) may also be used
				as regional analgesia if an analgesic
				is deemed necessary or it sustained
				release buprenorphine is not
			70 1 101 10	suitable/working. The most
			72 HOUR	commonly used local anesthetic
				agents are Lidocaine and
1001 267024		Mico		Bupivacaine. Lidocaine acts faster
1901-30722A	Lowe, Dawn	Mice		(within 2-5 minutes of injection) but
			SUBCEDV	
			AND DOST	Once the animal is under doon
				anesthesia, they are outbanized
				within 10 minutes (Intracardial
1001-367154	Lee Michael	Mico		
1901-00710A	Eee, Michael Faulk	MICE	SOCIAL	
1901-367084	Christopher	Mice		see protocol for details
1001 007 007	onnotopher	Miloe		Yes. The size of the ICU kennels are
				intended for animals which are in
				need of intensive care and are kept
			PRIMARY	in a more restricted space to keep
		Dog Dog Dog	ENCLOSURE	them quiet. These animals on this

				The main objectives for all the studies need intracranial EEG monitoring and the device working. If a device fails, then the study would end for that dog and a new dog obtained, the supply of research dogs with spontaneous seizures is not large, and to also reduce numbers of dogs we plan to re- implant the device if needed, Under the previous protocol this was permitted and we did replace the device in 2 dogs (one now deceased, the other on the current protocol).
1901-36697A	Patterson, Ned	Dog, Dog, Dog, Dog	MULTIPLE SURGERY	Going forward, as some of these dogs are and will be here for years, the PI will consult with the RAR vet about any concurrent health conditions, and the current seizure status of the dog before going ahead with a second intracranial surgery, and not proceeding if the concurrent conditions are deemed to be serious enough that the dog may not do well with the surgery and/or may not survive long enough post surgery to be valuable to the results. Specifically No second intracranial surgery will be performed without consultation with and approval by the RAR area vet, with their As the dogs are recovering from
1901-36697A	Patterson, Ned	Dog, Dog, Dog, Dog	ENVIRONMEN TAL ENRICHMENT	surgery, and need to be quiet and not have the EEG leads in the neck disturbed so for the 1-3 days of
1901-36697A	Patterson, Ned	Dog, Dog, Dog, Dog	SOCIAL HOUSING	As the dogs are recovering from surgery, and need to be quiet and not have the EEG leads in the neck
1901-36686A	Denton, Robert	Ampnibian (Other)	FREQUENCY	see protocol for details
1901-36686A	Denton, Robert	Amphibian (Other)	ENVIRONMEN TAL ENRICHMENT	see protocol for details

	Nonhuman		his eye position, a head post is implanted in an aseptic surgical procedure to permit monitoring of eye position (below). The animals is then trained to perform the task while maintaining fixation on a small spot on a video display under head restraint, and with some tasks to make specific eye movements to visual targets. This fixation control is necessary to insure that portion of the retina on which the stimuli fall is not an uncontrolled variable. □ □ Up to 12 months of training may be needed before an animal performs sufficiently well to allow us to collect neurophysiological data. When all training is complete, the animal undergoes another recovery surgery to implant an recording chamber. □ □ Implanting chambers in separate surgeries is scientifically justified because they have a limited useful lifetime. The multiple surgeries are needed for the scientific objectives of the experiments, and are related components of the research project. The stepwise surgical preparation of
1901-36681A	(Macaques)	SURGERY	to a small number of short surgeries,

				In the proposed research project, we
				intend to perform a primary surgical
				procedure during which an
				intravascular micro catheter inserted
				through the carotid artery is used to
				embolize the vascular supply of the
				femoral head. Monitoring the effects
				of the primary surgical procedure
				requires subsequent angiographic
				studies that also require access to
				the carotid artery. Thus, the second
				and potentially third and fourth
				surgical procedures (performed at
				least 7 days apart ) will entail only a
				minor cut-down procedure to the
				carotid artery (alternating the left
				and the right side) in anesthetized
				animals to allow insertion of the
				andiographic catheter and
				completion of the angiography
				These follow up surgeries are
				expected to be of very short
				duration (15-20 min) and expected
				to cause only minimum morbidity
				stress and discomfort These 2nd
		Pia	MULTIPLE	3rd and 4th surgical procedures that
1901-36656A	Toth Ferenc	(Biomedical)	SURGERY	are limited to surgical access to the
		Pia	SOCIAL	
1901-36656A	Toth, Ferenc	(Biomedical)	HOUSING	see protocol for details
	,	, ,	72 HOUR	
			POST-OP	
			ANALGESIA	
1901-36655A	Waye, Heather	salamander	POLICY	see protocol for details
			ENVIRONMEN	
			TAL	
1901-36655A	Waye, Heather	salamander	ENRICHMENT	see protocol for details
				Animals will be given a small
				incision in the femoral vein to
				administer cell treatment and/or
				Ferumoxytol injections. This will be
				done to ensure complete delivery of
				all therapeutic cells and contrast
				agent. Because the incision will be
				small and rats will be administered
				analgesics, the additional pain and
				distress experienced should be
				minimal.
	Grande,		MULTIPLE	
1901-36654A	Andrew	Rat, Mice	SURGERY	Prior to this procedure, rats undergo

				Skin biopsy will be performed up to
				3 times. There are two weeks
				between anv survival surgerv.
				Maximum of 3 survival surgeries per
			MULTIPLE	animal. These are minor survival
1812-36613A	Liu Liang	Mice	SURGERY	surgeries. The expected duration of
			NON-	
			PHARMACAUT	
			ICAL GRADE	We do not need an exception (Skin
1812-36613A	Liu Liang	Mice	COMPOUNDS	Biopsy)
			SOCIAL	
1812-36613A	Liu, Liang	Mice	HOUSING	see protocol for details
				One surgery is to perform AAV
				injections and the other is to perform
				the subsequent cranial window
				surgery. These are essential
				components of the same project.
				There will be no additional pain or
				distress due to having an additional
				survival surgery and only animals
				that are deemed healthy post the
1812-36610A	Lesne, Svlvain	Mice	SURGERY	initial surgery will move onto the
				Intratumoral injection is the only
				approach for NK cells effectively
				getting to GBM in orthotopic tumor
				model due to blood-brain barrier.
				Mice will be treated with
			MULTIPLE	Buprenorphine SR to cover the 72
1812-36595A	Chen, Clark	Mice	SURGERY	hour post-operative period. Mice will
	,			1. Cervical dislocation is used
				because organ congestion must be
				avoided to assess tumor
				dissemination and architecture. A
				common side effect of sedatives
				barbiturates and CO2 inhalation is
				congestion, which can obscures
				pathologic changes in vascular
				organs and tumor vasculature and
				cap thus render experiments most
				(unable to analyze vascular efforts
			FUTUANIAGIA	of genetic alteration or troatmont)
1812-365754	Modiano Jaime	Mice	METHOD	
1012 000707	Zhitnitskiv.	Pia	SOCIAL	
1811-36558A	Perle	(Agricultural)	HOUSING	see protocol for details
			EUTHANASIA	Field conditions without access to
1811-36497A	Barker, Fredrick	Bird (Other)	METHOD	CO2
	,	\	10000000 100000 100 000	19, 2040-001-00

				This study requires the implantation AND explantation of an osmotic mininump. Explantation is required
				for two reasons. 1) If the pump
				remained in the rat, drug would continue to be released and would
				affect assay results. 2) Removal of
				the pump allows us to record the
				accurate description of how much
				drug was administered within the
				time-frame studied (doing so following termination of the animal
				would add extra time for drug
				delivery).□
	Pravetoni,		MULTIPLE	□ Signs that will prompt additional
1811-36490A	Marco	Rat, Mice	SURGERY	analgesia□
			BLOOD	Animal will be euthanized by this
1911 264004	Pravetoni, Marco	Pat Miao		method and will not survive this
1811-30490A	Marco	Rat, Mice		In order to assess the relationship
				between metastatic burden as
				measured by IVIS (photon flux) and the visualization of
				macrometastases, we will be
				performing a pilot study which will
				allow the animals to live up to 10 weeks past injection of tumor cells
				This is necessary (Tail vein injection
				of breast cancer cells)
				In order to assess the full extent of
				which FAP-AT reduces tumor
			TUMOR	burden and metastases, we need to
1810-36480A	El-Ashry, Dorrava	Mice		animals up until the state at which
	,			In order to assess the full extent of
				which FAP-AT reduces tumor
				burden and metastases, we need to be able to determine a curve of
				metastatic burden as measured by
				IVIS versus macro-metastases that
				can be observed by eye. This will
				enable us to choose an endpoint
	El-Ashrv.			extent of drug efficacy by IVIS
1810-36480A	Dorraya	Mice	ENDPOINT	without having large numbers of

		1		The rat must first be injured and
	1	1	1	recover to model a chronic spinal
	1	1	ĺ	cord injury so that we can test our
	1	1	ĺ	scar ablation techniques and cell
	1	1	ĺ	transplantation. Pain and distress
	1	1		will be controlled through analgesics
	1	1	1	and antibiotics. 🗆
				(Spinal Cord Injury)
				The rat must first be injured and the
	1	1	ĺ	injury allowed to become chronic to
	1	1	1	test rose bengal scar clearance
	1	1		efficacy, then, the animal must be
	/	1	MULTIPLE	allowed to recover/secondary
1810-36461A	Parr, Ann	Rat	SURGERY	inflammatory response must
				Ovariectomy will be performed early
	1	1	ĺ	in life and is expected to be
	1	1	ĺ	completely healed in mice that will
	1	1	1	subsequently undergo additional
	Townsend,	1	MULTIPLE	surgical procedures, most
1810-36460A	DeWayne	Mice, Mice	SURGERY	commonly osmotic pump
				Several of our assay create a
	1	1	1	significant cardiac injury. This injury
	1	1	ĺ	can result in a moribund state. In
	1	1	1	some animals this period of
	1	1	ĺ	moribundity is temporary and the
	1	1		mice will eventually recover. In
	1	1	1	order to separate mice that will
	1	1	EUTHANASIA	ultimately survive from those with
			DEATH/MORIB	terminal dysfunction mice are
	Townsend,	1	UND	allowed to remain in a persistent
1810-36460A	DeWayne	Mice, Mice	ENDPOINT	moribund state. During these times,
	Townsend,	a and sources	SOCIAL	
1810-36460A	DeWayne	Mice, Mice	HOUSING	see protocol for details

	-			
				The multiple surgeries required in
				this study cannot be combined into
				one due to the severity and length of
				the individual procedures and
				recovery time required for the health
				of the animals. Specifically, the first
				surgery in the study requires a
				stereotaxic device, which would
				impede the success of the other
				surgeries and requires a different
				anesthetic regimen. The second
				procedure requires an extended
				recovery to allow for compensatory
				renal adaption. The third surgery
				requires that the first two surgeries,
				plus the collection of data over the
				control period, are already
				completed. (Electrolytic lesion of the
				OVLT)
				The multiple surgeries required in
1810-36452A	Collister, John	Rat	SURGERY	this study cannot be combined into
	,			Since food and water intake will be
				strictly monitored, any
				environmental enrichment (e.g.
				gnawing, chewing) could adversely
				affect the data, and due to the
			ENVIRONMEN	nature of the continual recording of
			TAL	blood pressure and heart rate in
1810-36452A	Collister, John	Rat	ENRICHMENT	each animal individually, social
				Food is not being restricted, and we
				strive to avoid any unnecessary
				handling of the rats. Food and fluid
				intake are measured and recorded
				daily, as well as urine output. 🏾
				However, upon noticing any unusual
			WEEKLY	hydration status or behavioral
			WEIGHING	changes, rats will be weighed, and if
1010 001501			EXCEPTION	the status persists or the weight of
1810-36452A	Collister, John	Kat		the animal is outside of what is
				Due to the nature of the continual
				heart rate in each animal
				individually, appial bauging will be
1810 364524	Collistor John	Pat		unaccontable due to cross talk of
1010-0040ZA		INCL	DUIDOUING	

-				
				two different series of experiments
				described separately below $\square$
				□ First Project 1 involves comparison
				of soveral patterns of epieid
				delivery. Our scientific chiestives
				require strict control over the total
				duration of obropic morphips
				avpagura (7 days), making it
				exposure (7 days), making it
				the pumpe at this time point. The
				numes must also be removed prior
				to tosting morphine conditioned
				place preference, to avoid any
				interference, to avoid any
				hitereferice with mobility during this
				L Second Projects 2 and 3 involve
				viral expression of light sensitive ion
				channels ("opsins"), to permit
				"aptegenetic" control of specific
				brain coll types and synaptic
				connections. One caveat to this
				connections. One caveat to this
				approach is that, even with viral
				transcriptional promotor, it takes
				time to accumulate sufficient ancin
				ume to accumulate sufficient opsin
				expression in brain cells to enable
				proposed experiments, it becomes
	Dothwoll			proposed experiments, it becomes
1910 264474	Rotriek	Mico		intracrapial virus injection before
1010-3044/A	Palnok	Mice	SURGERT	This will be used only for study of
				calcium signaling. Mice are
	Cyetanovic			decanitated with surgical sciesors
1810-364354	Marija	Mice	METHOD	without any anesthetic as anesthetic
	Cvetanovic	Miloo	SOCIAL	and any anconorio, as anconorio
1810-36435A	Marija	Mice	HOUSING	see protocol for details
	···· ··· ·j =·			

				The eximal will simply be reasing
				i ne animais will simply be receiving
				an injection of a euthanasia solution.
				This method will only be used in
				experiment 2 Here because we are
				measuring very labile molecular
				events (including phosphorylation)
				that require precise timing of
				sacrifice, sedatives or anesthesia
				would interfere with the scientific
				goals of the experiment.
				Only individuals trained in
			EUTHANASIA	decapitation will sacrifice the
1810-36429A	Meisel, Robert	Hamster	METHOD	animals. In addition the guillotine
			SOCIAL	
1810-36429A	Meisel, Robert	Hamster	HOUSING	see protocol for details
				All of the procedures performed on
				one animal are directly related to
				evaluation of the device. Clinically.
				this pulmonary valve would be
				implanted in a child, and when that
				abild group, they would underge a
				child grows, they would undergo a
				minimally invasive procedure to
				increase the diameter of the valve.
				The second procedure is a cut down
				to a vein and is minimally invasive
				This procedure will be performed
				under starile conditions and
				under sterne conditions and
				analgesia will be administered post
				operatively. (Pulmonary Valve
				Replacement)
				All of the procedures performed on
				one animal are directly related to
				one animal are uncerty related to
				evaluation of the device. Clinically,
				this pulmonary valve would be
				implanted in a child, and when that
				child grows, they would undergo a
				minimally invasive procedure to
				increase the diameter of the valve
				The subsequent expansions are to
				determine if the volve can be
				determine if the valve can be
				expanded multiple times, as this is
				the performance expectation for this
				device for clinical patients.
		Sheep	MULTIPI F	All of the procedures performed on
1810-364204	Rianco Richard	(Biomedical)	SURGERY	one animal are directly related to
1010-00420A	Dianco, Nichalu	(Diomedical)	SONOLINI	one animal are uncolly related to

		1	1	I
				The exception we request to the
				biopsy procedure is the use of
				isoflurane anesthesia for tail snips
				over 21 days. This would only be
	Costalonga,			done in the rare occasion that a
1810-36395A	Massimo	Mice	TAIL BIOPSY	second biopsy sample is needed
				AIM#2: As instructed by the
				inspector during the 2015 review,
				17d gestation fetuses must be
				decapitated before disposal.
				The oral candidiasis cortisone-
				induced mouse model we are
				studying may induce 25% weight
				loss at day 5 after inoculation of C.
				albicans. The experiment is only 5
	Costalonga,		EUTHANASIA	days long and in mice that on day 4
1810-36395A	Massimo	Mice	METHOD	are at 25% wight loss will be
			EUTHANASIA	If RAR veterinary staff require
			DEATH/MORIB	euthanasia of moribund mice, we
			UND	will follow through in the allowed
1810-36394A	Harris. Reuben	Mice	ENDPOINT	time.
	,		EUTHANASIA	We are collecting museum voucher
			DEATH/MORIB	samples and tissues from wild
			UND	rodents. These samples will allow
1809-36374A	Larsen. Peter	Wild Rodents	ENDPOINT	for metagenomic sequencing to
	,			Additionally, post-operative
				analgesic therapy is contraindicated
			72 HOUR	since provision of analgesics would
			POST-OP	inhibit the neurochemical changes
l I	Portoghese.		ANALGESIA	required to develop the
1809-36366A	Philip	Mice	POLICY	hypersensitive states to be studied
	Andrade.		SOCIAL	
1809-36347A	Rafael	Rabbit	HOUSING	see protocol for details
				We will harvest brain tissues from
				neonatal rodents vounder than 1
			EUTHANASIA	neonatal rodents younger than 1 week. According to IACUC

				For survival studies, and in
				particular preclinical drug trials,
				mice reaching the moribund state is
				considered an endpoint. This is
				applicable to mice with pancreas
				cancer. Animals are followed closely
				for signs and symptoms of
				advanced malignancy and are
				euthanized when they become
				moribund; it is our express aim not
				to let them progress to death for
				both humane and scientific reasons.
				Thus, we monitor the animals for
				general behavior and activity level;
				the development of severe
				cachexia, a cardinal manifestation
				of advanced pancreas cancer;
				and/or large palpable abdominal
				masses (> 2 cm). If these symptoms
				occur the mouse will be euthanized.
				It is further noteworthy that in our
				preclinical trials we carefully
				documents these symptoms in order
				to carefully identify the positive or
				negative effects of therapy on these
				hallmarks of pancreas cancer We
				note also that abdominal distension
				can develop and results from
				negative effects of therapy on these
			EUTHANASIA	hallmarks of pancreas cancer. We
			DEATH/MORIB	note also that abdominal distension
	Provenzano,		UND	can develop and results from the
1809-36341A	Paolo	Mice	ENDPOINT	accumulation of peritoneal ascites

				Avertin as opposed to an injection of
				Ketamine/Xylazine and an injection
				af Mahimahing marking Avarting own
				of Yonimpine makes Avertin our
				preferred choice. Additionally,
				ketamine/xylazine altered the course
				of pancreatic cancer as outlined in
				our protocol⊡
				Preparation:□
				Sterile filter with 0.2 micron filter. 🗆
				Store and use under sterile
				conditions. 🗆
				Store in the dark bottle or foil
				covered container. 🗆
				Store stock and working stock
				solutions at 4oC.□
			NON-	Do not use if the solution becomes
			PHARMACAUT	discolored or has a precipitate.□
	Provenzano,		ICAL GRADE	Check pH before each use and use
1809-36341A	Paolo	Mice	COMPOUNDS	only when greater than pH 5.□

				we are proposing to perform
				cervical dislocation without
				anesthesia due to the potential
				effects of anesthesia on the
				circulation and the induction of
				tissue injury. For example, we have
				consistently observed that lymph
				node, spleen, and bone marrow cell
				viability and function are adversely
				affected in situations in which cells
				are not rapidly obtained from the
				animal after elective sacrifice. We
				have noticed that new lab members
				who cannot rapidly harvest bone
				marrow and lymph nodes frequently
				have poor experimental results.
				Rapid acquisition of cells and
				tissues after circulation and
				oxygenation ceases is critical to
				preserve normal immune responses.
				The data are striking and illustrate
				that stem cells and lymphocytes are
				highly susceptible to apoptosis,
				corticosteriod-induced immune
				suppression as well as the well-
				known accumulation of nitric oxide
				and oxygen radicals that can occur
				with low circulation and oxygenation
				that will compromise our
				experiments, requiring more
				repetitions and hence more mice,
			EUTHANASIA	reagents, space in the colony and
1808-36332A	Tolar, Jakub	Mice	METHOD	resources. WHEN DONE

				Based on our initial trials of
				administering the AAV opsin
				construct followed by the immediate
				implantation of the fiber optic we
				noticed that we were not producing
				any expression of the opsin in the
				targeted region, but only along the
				shaft of the fiber optic. This was
				also confirmed in verbal
				communications with the Esther
				Krook-Magnuson lab. If the fiber
				optic is placed immediately after the
				delivery of the AAV, the AAV will
				concentrate around the fiber optic
				before it can be taken up by cells.
				Therefore, we are requesting to do
				an initial surgery to first inject the
				AAV opsin construct. This
				procedure is minimally invasive and
				typically only lasts an hour in length.
				The animal receives SR
				Buprenorphine for each surgical
				the mise handle each eurgen(
				constative national each surgery
				separately extremely well and do not
				any signs of pain or distress are
				seen following either procedure, we
				will euthanize the mouse If an
				animal fails to gain or maintain their
				weight, develops an infection, or
			MULTIPLE	starts showing signs of pain/distress
1808-36330A	Ebner, Timothy	Mice	SURGERY	following recovery of the first
			SOCIAL	,
1808-36330A	Ebner, Timothy	Mice	HOUSING	see protocol for details
			ENVIRONMEN	
			TAL	
1808-36330A	Ebner, Timothy	Mice	ENRICHMENT	see protocol for details

			-	Notes on use of non-pharmaceutical
				grade urethane:□
				To our knowledge, there is no
				pharmaceutical grade urethane
				available. However, we believe the
				use of urethane is still justified in our
				anesthetized, non-survival surgical
				procedures, as alternative
				anesthetics have considerable
				negative effects on physiology of the
				cerebral and cerebellar cortices. For
				example, ketamine is well known to
				block NMDA receptors in the brain,
				which are key receptors in the
				neuronal circuitry that we are
				studying (Bengtsson & Jorntell,
				2007). Barbiturates are also known
				to profoundly depress cerebellar
				circuitry (Sato, Y. et al., 1993).
				Additionally, isoflurane over time
				depresses cerebellar function (Loeb,
				A.L., et al., 1998), which is not ideal
				for our long, acute experiments.
				Therefore, urethane is the best
				available anesthetic agent for us to
				use to investigate cerebral and
				cerebellar function in our studies.□
				Bengtsson, F. & Jorntell, H.
			NON-	Ketamine and xylazine depress
				sensory-evoked parallel fiber and
1000 00000			ICAL GRADE	climbing fiber responses. J
1808-36330A	Ebner, Limothy	Mice		Neurophysiol 2007, 98(3):1697-
4000 0000 44	Gorr, Sven-	N4:	SOCIAL	
1808-36294A	UITIK	wice	HOUSING	see protocol for details

				Nake are proposible to parterm
				cervical dislocation without
				anesthesia due to the potential
				effects of anesthesia on the
				circulation and the induction of
				tissue injury. For example, we have
				consistently observed that lymph
				node spleen and hone marrow cell
				viability and function are adversely
				affected in situations in which cells
				are not rapidly obtained from the
				animal after elective sacrifice. We
				have noticed that new lab members
				who cannot rapidly harvest bone
				marrow and lymph nodes frequently
				have poor experimental results.
				Rapid acquisition of cells and
				tissues after circulation and
				oxygenation ceases is critical to
				preserve normal immune responses.
				The data are striking and illustrate
				that stem cells and lymphocytes are
				highly susceptible to
				apoptosis, corticosteriod-induced
				immune suppression as well as the
				well-known accumulation of nitric
				oxide and oxygen radicals that can
				occur with low circulation and
				oxygenation that will compromise
				our experiments, requiring more
				repetitions and hence more mice,
			EUTHANASIA	reagents, space in the colony and
1808-36286A	Tolar, Jakub	Mice	METHOD	resources. WHEN DONE
				Amputation is performed on mice
				that previously received intra-
				osseous tumor (by surgical
				procedure). The mice develop
				primary tumors relatively quickly,
				but our experimental aim is to
				achieve metastasis. We believe
				that the longer the tumor is present
	[]			the more likely metastasis will occur.
4000 000774	Largaespada,	haire -		we would perform the amputation
1808-36277A	David	IVIICE	ISUKGERY	when the mouse becomes

				The aggressive sampling scheme is
				necessary to capture the extent and
				duration of stress (measured by
				cortisol) and pain (measured by
				PGE2) associated with disbudding.
				We need 1ml of serum for both
				measures, so a volume of 3ml of
				whole blood is necessary. This
				aggressive sampling method has
				been described (Alvarez et al, 2009;
				Hempstead et al., 2018) with no
				apparent ill effects for the kids. $\square$
				Kids will be monitored for pallor
	N96		BLOOD	(pale oral mucous membranes) and
	Knauer,		COLLECTION	lethargy throughout the sampling
1808-36275A	Whitney	Goat	LIMIT	period. Kids will be fed a high plane
			NON-	
			PHARMACAUT	
Confliction and an all in the confliction and the state	2.91	10 - 10 - 1	ICAL GRADE	
1808-36261A	Pang, Hongbo	Mice	COMPOUNDS	see protocol for details
			72 HOUR	Subjects with peripheral neuropathy
			POST-OP	cannot receive post-operative
	Michaeli,	201 - 11	ANALGESIA	analgesics because these will inhibit
1808-36248A	Shalom	Rat	POLICY	the spinal neuroplasticity intended to

				Urethane is a widely used anesthetic
				in laboratory animal practice,
				especially in electrophysiologic
				studies (Maggi and Meli, 1986).
				Urethane has several advantages,
				including several possible
				administration routes, steady and
				long-lasting (6–12 h) surgical level
				of anesthesia, minimal effects on
				respiration and cardiovascular
				system, and muscle relaxation
				(Maggi and Meli, 1986, Hara and
				Harris, 2002). Although some
				thalamic and cortical suppression
				has been identified, several regions
				are only minimally modulated by
				urethane, and peripheral stimuli
				produce reflexes at the central
				nervous system level that modulate
				autonomic functions (Maggi and
				Meli, 1986). Nevertheless, urethane
				also has undesirable side effects. It
				causes hyperglycemia, and
				intraperitoneal injection induces
				necrosis in intra-abdominal organs
				(Maggi and Meli, 1986, Field and
				Lang, 1988). Urethane anesthesia is
				thus recommended to be terminal,
				Which precludes follow-up studies
				(Field and Lang, 1988).
	Miebooli			L
1000 262404	Nichaell,	Det		orethane has mild effects on
1000-30240A	эпают	Rai	COMPOUNDS	The neuropathic pain model, opinal
				ne neuropathic pain model, spinal
				cause some pain. Using analgesic
				drugs may antagonize the pain
				hypersensitivity and spinal neuronal
				(i.e. central) sensitization that is
			72 HOUR	developed after injury over time
			POST-OP	Administration of analgesic drugs
			ANALGESIA	post surgery however would
1808-36242A	Klein Amanda	Mice	POLICY	confound our behavioral
1000 002 12/ (			BLOOD	Blood collection will be performed
	Elmauist.		COLLECTION	post euthanasia. (Distributional
1807-36224A	William	Mice	LIMIT	Pharmacokinetics of anti-cancer
				In efficacy studies, the improvement
			EUTHANASIA	in survival upon treatment with
			DEATH/MORIB	novel anti-cancer agents will be
	Elmquist,		UND	evaluated. In these studies, tumor-
1807-36224A	William	Mice	ENDPOINT	bearing mice with deviations from
1807-36205A	Vannucci, Fabio	Pig (Agricultural)	EUTHANASIA METHOD	Per the AVMA guidelines, barbiturates have "a rapid onset of action, and loss of consciousness induced by barbiturates results in minimal or transient pain associated with venipuncture." In the event that an animal
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1807-36201A	Bianco, Richard	Sheep (Biomedical)	MULTIPLE SURGERY	This is a model development protocol. One of the primary difficulties in assessing transcatheter valves is the inability to anchor them in the annulus of a healthy animal. Other methods are used to mimic aortic stenosis e.g. external banding. This model most closely mimics the
1807-36197A		Mice, Rat, Dog, Cat, Rabbit, Guinea Pig, Chinchilla, Nonhuman Primate (Macaques), Pig (Biomedical), Sheep (Biomedical), Chicken, Turkey	SOCIAL HOUSING	

1807-36168A 1807-36152A 1807-36137A	Klein, Amanda Niedernhofer, Laura Georgieff, Michael	Mice Mice Mice	POLICY EUTHANASIA METHOD EUTHANASIA METHOD	Administration of analgesic drugs Culling purposes only. Preformed only on pups < 3 days. Decapitation without anesthesia or sedation will only be performed on postnatal day (P)0-P3 mice, for
			72 HOUR POST-OP ANALGESIA	The neuropathic pain model, spinal nerve ligation, proposed herein will cause some pain. Using analgesic drugs may antagonize the pain hypersensitivity and spinal neuronal (i.e. central) sensitization that is developed after injury over time.
1807-36193A		Nonhuman Primate (Macaques)	MULTIPLE SURGERY	Prior to training an animal to control his eye position, a head post is implanted in an aseptic surgical procedure to permit monitoring of eye position (see Headpost Implantation procedure). The animals is then trained to perform the task while maintaining fixation on a small spot on a video display under head restraint, and with some tasks to make specific eye movements to visual targets. This fixation control is necessary to insure that portion of the retina on which the stimuli fall is not an uncontrolled variable. (see Behavioral Training procedure). Up to 12 months of training may be needed before an animal performs sufficiently well to allow us to collect neurophysiological data. When all training is complete, the animal undergoes another recovery surgery to implant an recording chamber. Implanting chambers in separate surgeries is scientifically justified because they have a limited useful lifetime. The multiple surgeries are needed for the scientific objectives of the experiments, and are related components of the research project.

				To determine if mice in our model experience the same toxicities as
				patients, we request that we are
				criterion for morbidity and
				euthanasia. If mice can
				spontaneously recover after losing
				20% of their body weight then our
				model would not be that great. We
			EUTHANASIA	wouldn't know this if we had to
			DEATH/MORIB	euthanize them at the 20% weight
	Pennell,		UND	loss level. However, we think it
1807-36119A	Christopher	Mice	ENDPOINT	unlikely they could spontaneously
				Cervical dislocation is rapid and
				apparently painless. I have over 30
				years experience using this method
				We propose to develop a new
				model for clinical side effects of
				CAR immunotherapy. These side
				effects are CRS and neurologic
				adverse effects. Patients rapidly
				lose weight and experience systemic
				organ failure due to a sudden and
				systemic cytokine release. If left
				untreated, these toxicities are often
				fatal.□
				I o determine if mice in our model
				experience the same toxicities, we
				30% woight loss as one criterion for
				outhanasia (nlease note this
				exemption was granted in previously
				approved protocols). If mice can
				spontaneously recover after losing
				20% of their body weight, then our
				model would not be that great. We
				wouldn't know this if we had to
				euthanize them at the 20% weight
				loss level. However, we think it
				unlikely they could spontaneously
				recover after losing 30%. Since one
	Pennell,	5 cc	EUTHANASIA	of our goals is to reverse toxicity,
1807-36119A	Christopher	Mice	METHOD	we'd like to apply our proposed

				In order to test the efficiency of the
				virus to infect and express GFP on
				tumor tissue, we propose to use an
				orthotopic rat model injecting
				RDEney cells. For this approach we
				boothed cens. For this approach we
				need two surgenes, the first surgery
				will be the injection of the cells on
				the liver lobe and the second
				surgery will be the injection of the
				AAV virus on the portal vein or
				jugular vein. Anesthetics will be
	Gradilone,		MULTIPLE	titrated carefully and mice monitored
1807-36110A	Sergio	Rat	SURGERY	continuosly throughout monitoring
				For the jugular vein injection
			72 HOUR	procedure we spoke with the
			POST-OP	veterinarian and he told us that
	Gradilone		ANALGESIA	giving carprofen before the surgery
1807-36110A	Sergio	Rat	POLICY	and the day after would be enough
1007 001107	001910	i vat		The first surgery is for establishing
	Subramanian			the disease model. The secondary
1807 361034	Subree	Mice		surgeny is for treating the diseases
1007-30103A	Sublee	INICE	SUNGLINI	Bluet force will render the animals
				Biunt force will ferider the animals
				unconscious, after which the animal
				will be euthanized via cervical
				dislocation.
		2 101 1 102 0		Blunt force will render the animals
		Other* (Non-		unconscious, after which the animal
		USDA), Other*		will be euthanized via cervical
		(Non-USDA),		dislocation.
		Other* (Non-		
		USDA), Other*		Blunt force will render the animals
		(Non-USDA),		unconscious, after which the animal
		Other* (Non-		will be euthanized via cervical□
		USDA), Other*		dislocation.
		(Non-USDA)		
		Other* (Non-		Blunt force will render the animals
		USDA) Other*		unconscious, after which the animal
		(Non-LISDA)		will be euthanized via cervical
		Other* (Non		
				disioodion.
		(Other) Fish		Dlupt for an will reader the entire to
	Familia	(Other), FISN		Biunt force will render the animals
4000 000011	⊢orester,	(Other), Fish	LEUTHANASIA	unconscious, after which the animal
1806-36081A	James	(Other)		will be euthanized via cervical
	Alejandro,		SOCIAL	
1806-36072A	Emilyn	Mice, Mice	HOUSING	see protocol for details
	Alejandro,		EUTHANASIA	Only neonates (day 1) will be
1806-36072A	Emilyn	Mice, Mice	METHOD	euthanized via decapitation.

		Mias	NON- PHARMACAUT ICAL GRADE	There is no pharmaceutical grade avertin available. This animal line has high sensitivity to halogenated anesthetics (e.g. Isoflurane) to produce human malignant hyperthermia-like phenotype after exposure to these drugs (Chelu MG et al, FASEB J. 2006). In addition, in the prior studies, we have frequently experienced that this mouse lines has sensitivity to reduce cardiac function and hemodynamics during surgery (unpublished data). Only avertin among the drugs we tested did not change the basal cardiac function and hemodynamics compared to WT. Therefore, we will use avertin for all procedures we proposed in this animal protocol. □ 100% Tribromoethanol (avertin) stock solution will be prepared as follows. First, we will add non- pharmaceutical grade avertin (Sigma) to non-pharmaceutical grade tertiary amyl alcohol (Sigma) and completely dissolve it by heating and stirring. To use, we will dilute 100% stock to 2.5%, v/v, in diluent (0.8% NaCl, 1mM Tris (pH 7.4), 0.25mM EDTA, check the pH
1000-00040A		WICE		As indicated the birds are at a
				minimum of 2.5 Kg at the time of initial 24 hr fast. Once in the cages, we don't weigh them any more as there is an increased risk of injury taking them in and out of the cage repeatedly. Also with handling, feathers, scale, dander will contaminate the excreta collected leading to increased variability in the chemical composition. (Assay procedure-Turkey)
			VVEEKLY WEIGHING	As indicated the birds are at a minimum of 2.5 Kg at the time of
4000 000 45 4		Turkey,	EXCEPTION	initial 24 hr fast. Once in the cages,
1806-36045A	Noll, Sally	Chicken		we don't weigh them any more as
1806-36045A	Noll, Sally	Chicken	HOUSING	see protocol for details

				EAE is a paralytic disease that
				affects predominantly mobility of the
				experimental animals. Transient
				Debydration, fatigue and muscle
				waste are expected symptoms when
				mice reach a score of 3.0 (complete
				narelycic of bind limbe) and boyond
				These miss will reasive
				These fince will receive
4000 00000		Mine		supplemental nutrition, nutos and
1806-36038A	Lin, wensheng	wice		care on a twice daily basis. Animals
			ANESTHESIA,	Mice will be deeply anestnetized
			SURGERY,	with intraperitoneal injections of
			AND POST-	Avertin (425 mg/kg) prior to
			PROCEDURAL	transcardial perfusion. Depth of
			RECORDKEEP	anesthesia will be confirmed via lack
1806-36038A	Lin, Wensheng	Місе	ING	of toe pinch reflex. Euthanasia is
				When properly used by skilled
				personnel with well-maintained
				equipment, cervical dislocation may
				result in less fear and anxiety and
				be more rapid, painless, humane,
				and practical than other forms of
				euthanasia.
				Younger than 14-day-old pups will
				be will be euthanized by
				decapitation with scissors.
				Decapitation results in less fear and
				anxiety and be more rapid, painless,
				humane, and practical than other
				forms of euthanasia.
				EAE is a paralytic disease that
				EAE is a paralytic disease triat
				anects predominantly mobility of the
				experimental animals. Translent
				Denydration, fatigue and muscle
1000 00000			EUTHANASIA	waste are expected symptoms when
1806-36038A	Lin, Wensheng	Mice	METHOD	mice reach a score of 3.0 (complete

				Avertin has been the standard
				anesthetic in much mouse
				transgenic work. Advantages of
				Avertin are that it produces short-
				term (15-20 minutes) surgical
				anesthesia with good muscle
				relaxation and moderate respiratory
				depression, and that the mouse
				received it will recover within 30-60
				minutes. Usually, it takes less than 5
				minutes to perform EAE
				immunization. Moreover, we have
				used Avertin for EAE experiments
				for over 10 years (Avertin was
				approved for EAE experiments in
				our previous protocols 1209A21055
				and 1507-32810A). It is extremely
				important to use Avertin for our
				current and future EAE experiments,
				so that we make direct comparisons
				our previous, current, and future
				EAE work. Therefore, Avertin is
				selected due to its rapid induction of
				short-term anesthesia, quick
				recovery, low complication rate, and
				continuity of our EAE work. 🛛
				Avertin will be prepared according to
			NON-	the IACUC Guidelines. Briefly, ten
			PHARMACAUT	grams of 2,2,2-Tribromoethanol will
			ICAL GRADE	be suspended in 10 ml of tert-amyl
1806-36038A	Lin, Wensheng	Mice	COMPOUNDS	alcohol and serves as the stock
				DMD knockout model animals will
				have an end point of animal death
				without intervention in both the
				treated and untreated study groups.
			EUTHANASIA	The aim of the study is to determine
			DEATH/MORIB	the length of increase in disease
			UND	model animals with the treatment
1806-36033A	Garry, Daniel	Mice	ENDPOINT	and thus the treated and untreated

				Cervical dislocation will be
				performed by technically competent
				staff to ensure quick and complete
				euthanasia
				Decapitation will be used for
				euthanasia of mouse embryos and
				neonates up to day 10 $\square$
				Mouse peopates up to 10 days of
				age will be euthanized by methods
				according to NIH publication
				Guidelines for Euthanasia of Rodent
				Eetuses and Neonates (revisid
				6/22/16 Website
				https://oacu.oir.pib.gov/animal-
				research_advisory_committee_
				quidelines) Decapitation will be
				performed by now dispesable
				performed by new disposable
				of at the and of the precedure, or
				or at the end of the procedure, of
				⊔ Even event frame that NULL evide line v⊡
				Excerpt from the NIH guideline:
				Mouse, Rai, and Hamster Neonales
				up to 10 days of age. Acceptable
				methods for euthanasia include:
				injection of chemical anesthetics
				(e.g., pentobarbital), decapitation or
				cervical dislocation. Additionally,
1000 00000			EUTHANASIA	these animals are sensitive to
1806-36033A	Garry, Daniel	Місе		inhalant anesthetics; e.g., CO2, or
			72 HOUR	Our incisions are very small and
			POST-OP	seal right away with vetbond. Others
			ANALGESIA	who has performed this surgery did
1806-36019A	Siegfried, Jill	Mice	POLICY	not needed 3 days of analgenics

				we have read the Avertin guidelines
				in USDA Policy #3, Pharmaceutical-
				Grade Compounds in Research.
				However, we strongly feel that our
				planned use of Avertin is justified for
				the following reasons. First Avertin
				is the most widely used anesthetic in
				transgenic mouse research and is
				recommended by The Jackson
				Laboratory because it is easy to
				master administration, it is very fast
				acting and produces minimal
				obvious discomfort to the animal. In
				our experience with mouse
				anesthetics over the last 11 years at
			NON	the University of Missensin and
				University of Minnesota, we have
			ICAL GRADE	not experienced the significant post-
1806-36018A	Ervasti, James	Mice	COMPOUNDS	procedural mortality noted in the
				Currently our shelter cats (non-rerai)
				are receiving an iv dose of
				puprenorphine after induction then
				an oral dose of 0.1 mg/kg
				meloxicam after recovery and
				another 0.05 mg/kg dose of
				meloxicam to go home and
				administered 24 hours later.
				Unfortunately, there are no great
				take-home analgesics which are
				antiroly cofe to use in cote. NSAIDs
				entitlely sale to use in cats. NSAIDS
				are sale to use short-term but are
				not recommended to be used for
				multiple days in a row. Sublingual
				buprenorphine is a controlled
				substance and should not be
				dispensed to local rescue groups
				and shelters. Keeping these facts in
				mind, we consulted with a board-
				certified veterinary anesthesiologist
				and they recommended providing 2
				deeps of oral molovicom for pain in
				aste: they did not recommend
				providing a 3rd dose. This exact
				protocol has been approved by
				IACUC for this teaching lab for
				multiple years and we have not had
				any issues with post operative pain
				in our cats. This pain protocol is also
			72 HOUR	what is used at the Animal Humane
			POST-OP	Society (where most of our surgery
				natients are obtained from) and in
1906 360174	Horrlinger Sere	Cat. Doc		many votorinany benditale
1000-3001/A	rnenninger, sara	Cal, Dog		many veterinary nospitals

				L bo 2 day post operative menitoring
				of potients will be performed by the
				or patients will be performed by the
				rescue groups, toster parents, and
				local animal shelter staff. Each
				patient is discharged with detailed
				post-operative directions (see
				attachment labelled "detailed post
				operative care instructions") and
				caretakers are expected to review
				these documents and report any
				abnormalities to the U of MN faculty
				veterinarian via the emergency care
				phone line. Caretakers are expected
				to monitor each animal's activity
				level, mental attitude, elimination
				habits and food consumption and
				report any concerns. They are
				encouraged to check the incision
				daily and to report any redness,
				swelling, or discharge. They are also
				required to report any signs of pain
				or discomfort so additional pain
				medications can be prescribed. 🗆
				Last year, we added feral /
				community cat spay and neuter
				surgery to our teaching protocol.
			ANESTHESIA,	These cats are discharged to a
			SURGERY,	designated "feral cat coordinator"
			AND POST-	who is employed by the Animal
			PROCEDURAL	Humane Society or another rescue
			RECORDKEEP	group. After discharge the following
1806-36017A	Herrlinger, Sara	Cat, Dog	ING	steps are put into place to ensure
	<u>U</u>	· · · · ·	EUTHANASIA	Death is required to understand the
			DEATH/MORIB	effects of some of our experimental
			UND	therapies on GVHD. In order to
1806-36007A	Blazar, Bruce	Mice	ENDPOINT	understand their effect, death must

				Cervical dislocation. We are
				proposing to perform cervical
				dislocation without anesthesia due to
				the potential effects of anesthesia
				on the circulation and the induction
				of tissue injury. For example, we
				have consistently observed that
				lymph node, spleen, and bone
				marrow cell viability and function are
				adverselv affected in □
				situations in which cells are not
				rapidly obtained from the animal
				after elective sacrifice. We have
				noticed that new lab members who
				cannot rapidly harvest bone marrow
				and lymph nodes frequently have
				poor experimental results. Rapid
				acquisition of cells and tissues after
				circulation and oxygenation ceases
				is critical to preserve normal
				immune responses as measured in
				vitro by proliferation assays and in
				vivo. The data are striking and
				illustrate that stem cells and
				lymphocytes are highly susceptible
				to apoptosis, corticosteriod induced
				immune suppression as well as the
				well-known accumulation of nitric
				oxide and oxygen radicals that can
				occur with low circulation and
				oxvgenation that will compromise
			EUTHANASIA	our experiments, requiring more
1806-36007A	Blazar, Bruce	Mice	METHOD	repetitions and hence more mice.
	,		SOCIAL	· · · · · · · · · · · · · · · · · · ·
1806-36007A	Blazar, Bruce	Mice	HOUSING	see protocol for details
	,		MULTIPLE	
1806-35996A	Widge, Alik	Rat	SURGERY	see protocol for details
				The animal usually does not feel
				pain 24 hours post-surgery, but we
			72 HOUR	will monitor the rat everyday for at
			POST-OP	least 72 hours post-surgery.
			ANALGESIA	Analgesics will be given if the rat is
1806-35996A	Widge, Alik	Rat	POLICY	still feeling the pain either 48 or 72
	,			We need to weight animal everyday
			WEEKLY	to monitor their weight changes. The
			WEIGHING	weight will be an important indicator
			EXCEPTION	on whether we should increase or
1806-35996A	Widge, Alik	Rat	(FOOD/FLUID)	decrease the daily food distribution
			SOCIAL	· · · · · · · · · · · · · · · · · · ·
1806-35996A	Widge, Alik	Rat	HOUSING	see protocol for details

				This section is not applicable,
			FOOD/FLUID	poultry not housed in RAR facilities
			RESTRICTION	(Turkey Study Diets)
		Turkey,	RECORDKEEP	
1805-35968A	Noll, Sally	Chicken	ING	not applicable as the poutlry are not
				Due to the superficial nature of
				these melanoma tumors, the skin
				has a tendency to tighten and
				ulcerate at a very small size. In
				order to get sufficient data from our
				study and reduce the need to repeat,
				we would like to treat the ulceration
			TUMOR	with collasate ointment, instead of
			ENDPOINT	euthanizing the mouse before we
1805-35962A	Dong, Zigang	Mice	CRITERIA	can get sufficient data. We will treat
		Pig	EUTHANASIA	Personnel is trained to perform
1805-35959A	Pieters, Maria	(Agricultural)	METHOD	euthanasia (DVMs) directly in large
			NON-	
			PHARMACAUT	
		Pig	ICAL GRADE	
1805-35934A	laizzo, Paul	(Biomedical)	COMPOUNDS	see protocol for details
		Horse, Cow	ANESTHESIA,	
		(Biomedical),	SURGERY,	Standing sedation only. Clinical
		Goat, Sheep	AND POST-	cases are not monitored to this
		(Biomedical),	PROCEDURAL	level. Sedation is only used to the
		Camelid	RECORDKEEP	level needed to relax the animal, not
1805-35927A	Malone, Erin	(llamas &	ING	to perform surgery
			NON-	
			PHARMACAUT	
	10 54 Dec 65	Pig	ICAL GRADE	
1805-35922A	laizzo, Paul	(Biomedical)	COMPOUNDS	see protocol for details
	27500 1024 10 1000 10 10	N 10 10	SOCIAL	
1805-35921A	Meisel, Robert	Hamster	HOUSING	see protocol for details
				The animals will simply be receiving
				an injection of a euthanasia solution.
				I his method will only be used in
				Experiment 2. Here because we are
				measuring very labile molecular
			EUTHANASIA	events that require precise timing of
1805-35921A	Meisel, Robert	Hamster	METHOD	sacrifice, sedatives or anesthetics

			EUTHANASIA	C. difficile infection results in pseudomembranous enterocolitis in hamsters, which is typically a fatal condition. It is a rapid disease that generally peaks at 48 hours, and death/moribound state is the standard end-point in the literature. Of note, the human equivalent,
			DEATH/MORIB	which is responsible for $\sim 30,000$
1005 0501 14	Khoruts,	1.1	UND	deaths annually in the US, has 50%
1805-35914A	Alexander	Hamster		mortality with pest standard therapy.
1805-35914A	Alexander	Hamster	HOUSING	see protocol for details
				Uninephrectomy is required to study the effects of compounds on the left kidney, in isolation from any contralateral right kidney effects. Following uninephtrectomy the remaining left kidney undergoes hypertrophy. After 7-10 days hypertrophy is complete and a catheter can be placed into the outer medulla of the left kidney for compound administration. If the catheter was inserted during the same surgery as the uninephrectomy it's position would be displaced by the compensatory hypertrophy. (Uninephrectomy)
1805-35904A	Osborn Jr, John	Rat	MULTIPLE SURGERY	Prior uninephrectomy is required to study the effects of compounds on the left kidney, in isolation from any contralateral right kidney effects. Following uninephtrectomy the remaining left kidney undergoes hypertrophy. After 7-10 days
			NON	Inactin is non pharmaceutical and will be prepared in a manner that makes it compatible for animal use. As such we will take into account sterility, pH, purity and osmolarity when preparing the Inactin. New solutions will be made up daily. (Acute RBF/GFR)
1805-35904A	Osborn Jr, John	Rat	PHARMACAUT ICAL GRADE COMPOUNDS	will be prepared in a manner that makes it compatible for animal use. As such we will take into account

			ENVIRONMEN	
			TAL	
1805-35904A	Osborn Jr, John	Rat	ENRICHMENT	see protocol for details
				Anesthesia or sedation is generally
				not needed for iv administration of
the last sector and an and a sector and	2001 9 1004 1170	Cow	EUTHANASIA	barbituates to dairy cattle. If the
1805-35898A	Crooker, Brian	(Agricultural)	METHOD	animal is not calm, gentle restraint
				Sick mice cannot euthanized.We
				tried correlating histology to survival
				and surprisingly, did not find a
				correlation. It may be useful
				corollary data providing information
			EUTHANASIA	as to specific tissue site destruction
				but it does not correlate to survival.
4005 05004 4		N4:		Nor do data from in vitro assays
1805-35891A	Blazar, Bruce	MICe		(disparagingly referred to as 96-well
1805 35901 4	Blazar Pruco	Mice		see protocol for details
1000-30091A	Diazai, Diuce	wilce		Ve use cervical dislocation without
				anesthesia due to the potential
				depressive effects on the circulation
				and induction of tissue injury. We
				have consistently observed that
				lymph node, spleen and bone
				marrow cell viability and function are
				adversely affected in situations in
				which cells are not rapidly obtained
				from the animal after euthanasia.
				Rapid acquisition of cells and
				tissues after circulation and
				oxygenation ceases is critical to
				preserve normal immune responses
				as measured in vitro by proliferation
				assays and in vivo by GVHD effects.
				I he data are striking and illustrate
				that stem cells and lymphocytes are
				nigniy susceptible to apoptosis,
				curricosteroid-induced immune
				suppression as well as the well-
				and oxygen radicals that can occur
				with low circulation and poor
				oxygenation that will compromise
				our experiments. Carbon dioxide
				inhalation prior to cervical
				dislocation would adversely upset
				the acid-base balance resulting in
				acidosis which would be important
				on a cellular level for in vivo transfer
			EUTHANASIA	of cells and in vitro assays. When
1805-35891A	Blazar, Bruce	Mice	METHOD	done correctly, cervical dislocation

1805-35885A   Binstadt, Bryce   Mice   SURGERY   see protocol for details     1805-35885A   Binstadt, Bryce   Mice   SURGERY   see protocol for details     1805-35885A   Binstadt, Bryce   Mice   SURGERY   see protocol for details     1805-35883A   Low, Walter   Mice   SURGERY   see protocol for details     1805-35883A   Low, Walter   Mice   SURGERY   see protocol for details     1805-35883A   Low, Walter   Mice   SURGERY   see protocol for details     1805-35883A   Low, Walter   Mice   SURGERY   see protocol for details     1805-35883A   Low, Walter   Mice   SURGERY   see protocol for details     1805-35883A   Low, Walter   Mice   SURGERY   see protocol for details     1805-35883A   Low, Walter   Mice   SURGERY   see protocol for details     1805-35883A   Low, Walter   Mice   SURGERY   Surgery or surgeries to administer     1805-35883A   Low, Walter   Mice   SURGERY   This experiment requires additional surgery or surgeries to administer				MULTIPLE	
1 Init experiment requires additional surgery or surgeries to administer   2 ika virus or glioma cells infected with Zika virus. For the groups in Experiment 1B, these incoulations will be given on days 3, 7, and 14 following intracranial tumor implantation. Mice will be anesthetized during each surgical procedure. There are no sensory fibers within the brain to cause pain to the animal. Only the small incision will be noticed, and any minor pain or distress will be minimized with anesthetics and analgesics.   This experiment requires additional surgery or surgeries to administer Zika virus. Rats will be minimized with anesthetics and analgesics.   This experiment requires additional surgery or surgeries to administer Zika virus. Rats will be anesthetized during each surgical procedure. There are no sensory fibers within the brain to cause pain to the animal. Only the small incision will be noticed, and any minor pain or distress will be anesthetized during each surgical procedure. There are no sensory fibers within the brain to cause pain to the animal. Only the small incision will be noticed. and any minor pain or distress will be minimized with anesthetics and analgesics. (Induction of brain tumor cell line (mice))   1805-35683A Low, Walter Mice MULTIPLE This experiment requires additional surgery or surgeries to administer   1805-35683A Low, Walter Mice SURGERY Surgery or surgeries to administer   1805-35683A Low, Walter Mice EUTHANASIA Counterparts. Each animal will be count reduction ani/or ablation, and whether they live longer than their untreated counterparts. Each animal will be counterparts. Each animal w	1805-35885A	Binstadt, Bryce	Mice	SURGERY	see protocol for details
1805-35883A Low, Walter Mice Mul Mul Mul This experiment requires additional surgical procedure. There are no sensory fibers within the brain to cause pain to the animal. Only the small incision will be noticed, and any minor pain or distress will be minimized with anesthetics and analgesics.   This experiment requires additional surgical procedure. There are no sensory fibers within the brain to cause pain to the animal. Only the small incision will be noticed, and any minor pain or distress will be minimized with anesthetics and analgesics.   This experiment requires additional surgery or surgeries to administer Zika virus or glioma cells infected with Zika virus. Rats will be anesthetized during each surgical procedure. There are no sensory fibers within the brain to cause pain to the animal. Only the small incision will be noticed, and any minor pain or distress will be minimized with anesthetics and analgesics.   1805-35883A Low, Walter Mice MULTIPLE This experiment requires additional surgery or surgeries to administer animal. Only the small incision will be noticed, and any minor pain or distress will be minimized with anesthetics and analgesics. (Induction of brain tumor cell line (mice))   1805-35883A Low, Walter Mice SURGERY This experiment requires additional surgery or surgeries to administer and analgesics and analgesits achineved Animals with the path there untreated ana					i his experiment requires additional
Zika virus or glioma cells infected with Zika virus. For the groups in Experiment 18, these inoculations will be given on days 3, 7, and 14 following intracranial tumor implantation. Mice will be anesthetized during each surgical procedure. There are no sensory fibers within the brain to cause pain to the animal. Only the small incision will be noticed, and any minor pain or distress will be minimized with anesthetics and analgesics.   This experiment requires additional surgery or surgeries to administer Zika virus or glioma cells infected with Zika virus. Rats will be anesthetized during each surgical procedure. There are no sensory fibers within the brain to cause pain to the animal. Only the small incision will be noticed, and any minor pain or distress will be anesthetized during each surgical procedure. There are no sensory fibers within the brain to cause pain to the animal. Only the small incision will be noticed, and any minor pain or distress will be anesthetized during each surgical procedure. There are no sensory fibers within the brain to cause pain to the animal. Only the small incision will be noticed, and any minor pain or distress will be minimized with anesthetics and analgesics. (Induction of brain tumor cell line (mice))   1805-35883A Low, Watter Mice MULTIPLE This experiment requires additional surgery or surgeries to administer   1805-35883A Low, Watter Mice SURGERY Surgery or surgeries to administer   1805-35883A Low, Watter Mice EUTHANASIA polytical as quickly as possible once this state is achieved. Animals					surgery or surgeries to administer
1805-35883A Low, Walter Mice MULTIPLE This experiment requires additional surgery or surgerises additional analgesics. (Induction of brain tumor discusse pain to chase pain to the animal. Only the small incision will be noticed, and any minor pain or distress will be minimized with anesthetics and analgesics.   1805-35883A Low, Walter Mice MULTIPLE This experiment requires additional surgery or surgeries to administer SURGERY   1805-35883A Low, Walter Mice SURGERY This experiment requires additional surgery or surgeries to administer SURGERY   1805-35883A Low, Walter Mice SURGERY This experiment requires additional surgery or surgeries to administer SURGERY   1805-35883A Low, Walter Mice SURGERY This experiment requires additional surgery or surgeries to administer Surgery or surgery or surgeries to administery Surgery or surgery or surgeries to administery Surgery or surger					Zika virus or glioma cells infected
1805-35883A Low, Walter Mice MULTIPLE This experiment requires additional surgical procedure. There are no sensory fibers within the brain to cause pain to the animal. Only the small incision will be noticed, and any minor pain or distress will be minimized with anesthetics and analgesics.   1805-35883A Low, Walter Mice MULTIPLE This experiment requires additional surgical procedure. There are no sensory fibers within the brain to cause pain to the animal. Only the small incision will be noticed, and any minor pain or distress will be minimized with anesthetics and analgesics.   1805-35883A Low, Walter Mice MULTIPLE This experiment requires additional surgery or surgeries to administer and only the small incision will be noticed, and any minor pain or distress will be anesthetized during each surgical procedure. There are no sensory fibers within the brain to cause pain to the animal. Only the small incision will be noticed, and any minor pain or distress will be minimized with anesthetics and analgesics. (Induction of brain tumor cell line (mice))   1805-35883A Low, Walter Mice MULTIPLE This experiment requires additional surgery or surgeries to administer additional surgery or surgeries to additi					with Zika virus. For the groups in
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1805-35883A Low, Walter Mice MULTIPLE This experiment requires additional analgesics. (Induction of brain to cause pain to the animal. Only the small incision will be noticed, and any minor pain or distress will be minimized with anesthetics and analgesics.   1805-35883A Low, Walter Mice MULTIPLE This experiment requires additional incision will be noticed, and any minor pain or distress will be minimized with anesthetics and analgesics.   1805-35883A Low, Walter Mice MULTIPLE This experiment requires additional surgery or surgeries to administer Zika virus or glioma cells infected with Zika virus. Rats will be anesthetized during each surgical procedure. There are no sensory fibers within the brain to cause pain to the animal. Only the small incision will be noticed, and any minor pain or distress will be anesthetized during each surgical procedure. There are no sensory fibers within the brain to cause pain to the animal. Only the small incision will be noticed, and any minor pain or distress will be analgesics. (Induction of brain tumor cell line (mice))   1805-35883A Low, Walter Mice MULTIPLE This experiment requires additional analgesics. (Induction of brain tumor cell line (mice))   1805-35883A Low, Walter Mice SURGERY Surgery or surgeries to administer intered animals experience tumor reduction and/or ablation, and whether they live longer than their untreated counterparts. Each animal will be once this state is achieved. Animals WID once this state is achieved. Animals with the processible untanized as quickly as possible untanized as quickly as possible					will be given on days 3, 7, and 14
1805-35883ALow, WalterMiceMultTIPLEThis experiment requires additional analgesics. (Induction of brain tumor cell line (mice))1805-35883ALow, WalterMiceEUTHANASIA DEATH/MORIB UNDA state of morbundity may be reached to determine if treated animals experience tumor reduction and/or ablation, and whether they live longer than their untreated counterparts. Each animal will be counterparts. Each animal w					following intracranial tumor
1805-35883A Low, Walter Mice MULTIPLE This experiment requires additional surgery or surgerises to administer analyminor pain or distress will be an analgesics.   1805-35883A Low, Walter Mice EUTHANASIA A state of moribundity may be reached to determine if freated animals experience tumor reduction administer state is achieved. Animals will be minimized with anesthetics and analgesics.   1805-35883A Low, Walter Mice EUTHANASIA EUTHANASIA   1805-35883A Low, Walter Mice EUTHANASIA Councerthan their untreated curron their untreated curron their untreated curron their untreated curron to calked their the part to councerthin the curron their untreated to the their untreated to the their untreated to the treatment and/or ablation, and whether they live longer than their untreated curron their untreated to currengers. Each animal will be entremated to currengers. Each animal will be entremat					implantation. Mice will be
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1805-35883ALow, WalterMiceMULTIPLE SURGERYminor pain or distress will be minimized with anesthetics and analgesics.1805-35883ALow, WalterMiceEUTHANASIA DEATH/MORIB UNDA state of moribundity may be reached to determine if treated animals experience tumor reduction and/or ablation, and whether they live hore reached to determine if treated animals experience tumor reduction and/or ablation, and whether they live hore reached to determine if treated animals experience tumor reduction and/or ablation, and whether they live hore the state is achieved. Animals undot to the state is achieved. Animals undot to the state is achieved. Animals undot to the state is achieved. Animals1805-35883ALow, WalterMiceEUTHANASIA1805-35883ALow, Walter					incision will be noticed, and any
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1805-35883A Low, Walter Mice Feached to determine if treated animals experience tumor reduction and/or ablation, and whether they live longer than their untreated counterparts. Each animal will be euthanized as quickly as possible once this state is achieved. Animals   1805-35883A Low, Walter Mice ENDPOINT which respond to treatment and					A state of moribundity may be
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1805-35883A Low, Walter Mice EUTHANASIA live longer than their untreated counterparts. Each animal will be bEATH/MORIB   euthanized as quickly as possible UND once this state is achieved. Animals   this possesary to collect this many					and/or ablation, and whether they
EUTHANASIA counterparts. Each animal will be   DEATH/MORIB euthanized as quickly as possible   UND once this state is achieved. Animals   1805-35883A Low, Walter   Mice ENDPOINT   which respond to treatment and					live longer than their untreated
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1805-35883A Low, Walter Mice ENDPOINT which respond to treatment and					once this state is achieved. Animals
	1805-35883A	Low, Walter	Mice	ENDPOINT	which respond to treatment and
bloode to adoptiotative that					It is necessary to collect this many
biologs to adequately study the					production adequately study the
Animala will receive requesitation					Animals will receive recussitation
RI OOD If luids during the protocol as well as				BLOOD	fluids during the protocol as well as
Beilman Pig COLLECTION flushes after each blood draw		Beilman	Pig		flushes after each blood draw
1805-35872A Gregory (Biomedical) [1MIT Animals will not be allowed to waken	1805-35872A	Gregory	(Biomedical)		Animals will not be allowed to waken

1804-35863A	Yee, Douglas	Mice	NON- PHARMACAUT ICAL GRADE COMPOUNDS	Avertin does not come as a certified pharmaceutical grade compound. We have historically had problems with other anesthetics killing the mice and have found this one to be quite reliable. Avertin is stored as a powder until right before surgery. Then it is weighed and brought up in solution. As soon as it is no longer
				Re-catheterization in case of catheter failure. Intervals are based on the animals catheter patentcy.□ Typically recatheterization surgeries only occur within the 3 weeks of the original surgery if at all. Each rat would undergo a maximum of 1 re- catheterization using the alternate jugular vein. The initial surgery utilizes the rat's right jugular vein and the re-catheterization surgery (if necessary) uses the left side. Animals□ are anesthetized for the procedure. Breathing rate and animals sensitivity to touch will be monitored to□ determine the state of anesthesia and overall well being of the animal. We will use a heating pad from the□ time of anesthesia till the animal is awake and moving around normally. (Intravenous catheter surgery)
1804-35861A	Spencer, Sade	Rat	MULTIPLE SURGERY	usually performed in immediate succession (i.e. catheter surgery be
				Occasionally animals will have to be genotyped post-weaning if we require RAR staff to wean animals as opposed to laboratory staff or if we need to redo the biopsy due to accidental contamination. We will take every measure to minimize post-weaning tail biopsies as they can lead to additional stress and
1804-35861A	Spencer, Sade	Rat	TAIL BIOPSY	pain to the animals. In all cases we
180 <u>4</u> -3 <u>5</u> 861A	Spencer, Sade	Rat	HOUSING	see protocol for details

1804-35861A	Spencer, Sade	Rat	ENVIRONMEN TAL ENRICHMENT	see protocol for details
1804-35859A		Nonhuman Primate (Macagues)	MULTIPLE SURGERY	refinement we have implemented in which the electrodes are chronically implanted and sealed negating the need for chamber cleanings, and thus reducing the number of times that animal is required to leave its home cage. In all cases, revision surgeries may be necessary if one of the chambers is misaligned or becomes damaged during the course of the studies. These revision surgeries, however, are rare in our experience. (Chamber Implant)
1804-35856A	Li. Fagian	Mice	EUTHANASIA METHOD	Fetuses are neither sentient nor conscious prior to birth and thus incapable of actually perceiving pain.□ When fetuses (mouse>E15) are needed for study, euthanasia of individual fetuses induced by decapitation with surgical scissors is
1804-35852A	l in Gufa	Mice	MULTIPLE	The major objectives of this study is to investigate whether limb/digit regeneration can be enhanced by cell transplantations, with or without further application of growth factors. We have shown that the cell transplants did not survive well if cells were applied at the same time when the digit was amputated. The best time for cell transplantation in the mouse digit stumps is around 2 weeks after amputation. This has also been reported recently by Dr. Ken Muneoka's groups, published in

r	1			
				Avertin is used in this study, in
				addition to using isoflurane, as an
				anesthetic methods, to
				accommodate the imaging process,
				especially the fluorescence imaging
				under directing microscope, when
				proper isoflurane set up is difficult to
				set up to allow efficient imaging of
				the animals.
				Careful observation of the
				experimented mice in our previous
				work has indicated that there is no
				complications of ileitis, peritonitis or
			NON-	muscle necrosis. No signs of these
			PHARMACAUT	conditions have been indicated in
			ICAL GRADE	our post-mortem examination of the
1804-35852A	Lin, Gufa	Mice	COMPOUNDS	animal body and tissues collected.
				Petri dishes are replaced every day
1001050114	D: 1 ( 1 )	Fish (Zebra	SANITATION	and not used again. Tanks are
1804-35844A	Bischot, John	tisn)		replaced and cleaned at sigh of
			WEEKLY	
			WEIGHING	Net englischer (Diete fen Obiesting
4004.050004		<b>T</b>		Not applicable. (Diets for Objective
1804-35833A	INOII, Saliy	тигкеу		2 growth study)
1804 358334	Noll Sally	Turkey		see protocol for details
1004-000007	Non, Cany	Turkey		
			RESTRICTION	
			RECORDEEP	Not applicable (Diets for Objective 2
1804-358334	Noll Sally	Turkey	ING	arowth study)
1004-000007	I toll, Oally	Turkey		growth study)
			TAI	
1804-35833A	Noll Sally	Turkey	ENRICHMENT	see protocol for details
	i ton, ounj	, and y	SOCIAL	
1804-35829A	Noll, Sally	Turkev	HOUSING	see protocol for details
	· · · · · · · · · · · · · · · · · · ·			Producers do not have or use
			EUTHANASIA	sedation on their farms, we will only
1804-35829A	Noll, Sallv	Turkev	METHOD	use cervical dislocation for the
	, ,	,	SOCIAL	
1804-35828A	Aliota, Matthew	Mice	HOUSING	see protocol for details
			ENVIRONMEN	
			TAL	
1804-35825A	Noll, Sally	Turkey	ENRICHMENT	see protocol for details
			SOCIAL	
1804-35825A	Noll, Sally	Turkey	HOUSING	see protocol for details

				The exception for death as an
				endpoint will be pre-determined
				endpoint for the experiment
				tunically 100 days part transplant
				typically 100 days post transplant.
				As stated, control transplanted
				animals (BM only) are not expected
				to show signs of GvHD, and as such
				will typically survive for the term of
			EUTHANASIA	the experiment. Additionally, if all
			DEATH/MORIB	relevant experimental groups have
				succumbed to GvHD by for
1804 358154	Blazar Bruco	Mice		example, day 60 it would not be
1004-00010A		MICE		We use cervical dislocation without
				anesthesia due to the potential
				depressive effects on the circulation
				and induction of tissue injury. We
				have consistently observed that
				lymph pada, sploon and bono
				morrow coll vibility and function and
				manow cen viability and function are
				adversely affected in situations in
				which cells are not rapidly obtained
				from the animal after euthanasia.
				Rapid acquisition of cells and
				tissues after circulation and
				oxygenation ceases is critical to
				preserve normal immune responses
				as measured in vitro by proliferation
				assays and in vivo by GVHD offorts
				The data are striking and illustrate
				that stem cells and lymphocytes are
				highly susceptible to apoptosis,
				corticosteroid-induced immune
				suppression as well as the well-
				known accumulation of nitric oxide
				and oxygen radicals that can occur
				with low circulation and poor
				oxygenation that will compromise
				our experiments. Carbon dioxide
				inhalation prior to convical
				dialogotion would advargate uport
				dislocation would adversely upset
				the acid-base balance resulting in
				acidosis which would be important
				on a cellular level for in vivo transfer
			EUTHANASIA	of cells and in vitro assays. Other
1804-35815A	Blazar, Bruce	Mice	METHOD	inhaled anesthetics such as
				The neuropathic pain model (SNL)
				proposed herein will cause some
				pain. Administration of analgesic
			72 HOUR	drugs however, would confound our
			POST-OP	electrophysiological experiments in
				which we like to determine the
4004.050004		Mine		
1804-35808A	ikiein, Amanda	INICE	POLICY	enects of KATP channel modulators

				The neuropathic pain model, spinal
				nerve ligation, proposed herein will
				cause some pain. Using analgesic
				drugs may antagonize the pain
				hypersensitivity seen in primary
				afferent fibers and spinal neuronal
				(i.e. central) sensitization that is
				developed after injury over time
				Administration of analysis drugs
				Administration of analysis drugs
1004 250004	Klain Amanda	Mino		post surgery, nowever, would
1604-3060ZA	Klein, Amanua	MICE	POLICY	The neuropathic pain model, enjoy
				ne neuropathic pain model, spinal
				nerve ligation, proposed herein will
				cause some pain. Using analgesic
				drugs may antagonize the pain
				nypersensitivity and spinal neuronal
				(i.e. central) sensitization that is
			72 HOUR	developed after injury over time.
			POST-OP	Administration of analgesic drugs
			ANALGESIA	post surgery, however, would
1804-35801A	Klein, Amanda	Mice	POLICY	confound our behavioral
				We will perform experiments with
				small numbers of immune-deficient
				NOS2-/- and Irgm1-/- mice to
				measure virulence of M.
				tuberculosis mutant strains by
				monitoring survival of animals. We
				have also categorized all NOS2-/-
				and Irgm1-/- animals that will be
				used for the 6 week time points as
				Pain Class C, since this end point is
				close to the expected time of onset
			EUTHANASIA	of disease symptoms (7 weeks for
			DEATH/MORIB	Irgm1-/- mice, 9 weeks for NOS2-/-
			UND	mice), so these mice may
1804-35785A	Tischler, Anna	Mice	ENDPOINT	experience some overt signs of
				In project 4, for mouse sepsis
				protection model, we expect the
				control infected mice without
				treatment will die within 48 hours
				after infection. This protection model
				is a gold standard approach for
				validation of the efficacy of novel
				antibacterial agents (Ling et al. 2015
			EUTHANASIA	Nature 517:455). The mice will
			DEATH/MORIB	suffer from infections including pain
		Mice, Chicken		and distress: however we will
1803-35768A	Ji. Yinduo	Turkev	ENDPOINT	monitor the mice every two hours
	Gordon-Evans		EUTHANASIA	unlikely needed if owner has
1803-35743A	Wanda	Dog. Cat	METHOD	decided to euthanize
		. V		

		· · · · · · · · · · · · · · · · · · ·	· · · · · · · · · · · · · · · · · · ·	
				In order to "reduce and refine" these
				studies, we would like to keep
				animals on study that have non-
				cavitated ulcerations <1cm2. The
				animals will be treated with collasate
				ointment and monitored three times
				weekly by lab staff. Certain cell
				lines have a tendency to ulcerate
			TUMOR	the skin before the tumor is an
			ENDPOINT	adequate size, ergo if we treat the
1803-35739A	Dong, Zigang	Mice	CRITERIA	minor ulcerations we can keep the
				Mice will be euthanized on the same
				day they become moribund. Mice
				are allowed to reach moribund state
				because in order for our
			EUTHANASIA	experiments to produce good
			DEATH/MORIB	results, AML should be as prominent
			UND	in the mouse as possible. Often,
1803-35735A	Sachs, Zohar	Mice	ENDPOINT	this state co-occurs as moribundity.
	Richard,		SOCIAL	
1803-35725A	Jocelyn	Rat	HOUSING	see protocol for details
				Additionally, post-operative
				analgesic therapy is contraindicated
			72 HOUR	since provision of analgesics would
			POST-OP	inhibit the neurochemical changes
			ANALGESIA	required to develop the
1803-35719A	Wong, Henry	Mice	POLICY	hypersensitive states to be studied.

			1	IVVhen treatment works, some times
				ulceration is observed on the treated
				tumor as a result of tumor necrosis.
				We will observe the small ulceration
				up to 7 days unless continuous
				bleeding or infection is observed or
				reaching other euthanization criteria.
				We will give ketoprofen 5mg/kg SC
				q24h when necessary. Pain level C.
				EXP 1. intratumoral injection of
				adenovirus (head-neck sub-cu
				tumor, mice))
				When treatment works, some times
				ulceration is observed on the treated
				tumor as a result of tumor necrosis.
				We will observe the small ulceration
				up to 7 days unless continuous
				bleeding or infection is observed or
				reaching other euthanization criteria.
				We will give ketoproten smg/kg SC
				(24h when necessary. Pain level C.
				(EXP 3. combination therapy (sub-
				cu, it, panc, colori, neadneck, mice) j
				Oncolytic adenoviruses occationally
				induce tumor ulceration when anti-
				tumor effect is strong. Usually,
				ulceration is seen before the tumor
				disappears. This is a part of
			TUMOR	therapeutic effect and the ulceration
	Yamamoto,		ENDPOINT	is self limiting. We want to observe
1803-35712A	Masato	Mice, Hamster	CRITERIA	ulcerated tumor up to / days unless
				The piglet would undergo a
				septectomy, making the left and
				right atrial champers a single
				champer, which may create a
				mypoxic state for the pigret. We also
		Dia		(DA) to increase the pressure on the
		(Riomedical)		right ventricle (RV). Our rationale is
		Pig		that the piglet would have time to
1803-35699A	Martin Cindy	(Riomedical)	SURGERY	compensate for these smaller, but
1000 000000	With the group of the second s			This procedure is a moderate
				restrain. As such the animals are
			PHYSICAL	still able to move about the small
1803-35671A	Greising, Sarah	Mice	RESTRAINT	area, but is it merely restricted from

				As we stated above, along with the
				tumor lysis caused by hypoxia
				tumor ulceration might be observed
				in animals receiving subcutaneous
				tumor cells. Tumor pecrosis could
				be also an indicator of the
				successful therapeutic effect of the
				successful therapeditic effect of the
				suggested treatment. In some
				ulases, turnor necrosis reads to the
				is the phonomenon observed in the
				is the phenomenon observed in the
				L Ma would like to observe the
				we would like to observe the
				rodenis with uicers without
				antibiotics of analysia in order to
				avoid possible effect of those drugs
				to the tumor response. Thus, pain
				level C is requested in order to
				observe the ulcerated tumor without
				antibiotics of analgesia.
				Animais will be monitored as their
				tumor grows for assessment of
				tumor size and overall animal
				condition every second day and/or
				as frequently as is required by the
				area veterinarian after a treatment
				plan is discussed. This monitoring
				will include weekends and holidays.
			ENDPOINT	
1803-35667A	Davydova, Julia	Mice, Hamster	CRITERIA	
				The neuropathic pain model (SNL)
				proposed herein will cause some
				pain. Administration of analgesic
			72 HOUR	drugs, however, would contound our
			POST-OP	electrophysiological experiments in
			ANALGESIA	which we like to determine the
1802-35633A	Klein, Amanda	Місе	POLICY	effects of KAIP channel modulators
				Pumps will be implanted 3-5
				days post tumor inoculation.
				I his allows us the time to□
				Image mice to ensure tumor□
				growth. (Subcutaneous Alzet pump
				implantation)
			MULTIPLE	
1802-35613A	Olin, Michael	Mice	SURGERY	Tumors need to be established prior
			EUTHANASIA	
			DEATH/MORIB	
	565875898 64 18-10 5756		UND	Experiments need to determine
1802-35613A	Olin, Michael	Mice	ENDPOINT	moribundity to monitor treatments

				To determine if mice in our model
				experience the same toxicities as
				patients, we request that we are
				allowed to use 30% weight loss as a
				criterion for morbidity and
				euthanasia. If mice can
				spontaneously recover after losing
				20% of their body weight, then our
				model would not be that great. We
			EUTHANASIA	wouldn't know this if we had to
			DEATH/MORIB	euthanize them at the 20% weight
	Pennell,		UND	loss level. However, we think it
1802-35603A	Christopher	Mice	ENDPOINT	unlikely they could spontaneously
				Cervical dislocation is rapid and
				apparently painless. I have over 30
				years experience using this method
				of euthanasia.
				We propose to develop a new
				model for clinical side effects of
				CAR immunotherapy. These side
				effects are CRS and neurologic
				adverse effects. Patients rapidly
				lose weight and experience systemic
				organ failure due to a sudden and
				systemic cytokine release. If left
				untreated, these toxicities are often
				fatal. ⊓
				To determine if mice in our model
				experience the same toxicities we
				request that we are allowed to use
				30% weight loss as one criterion for
				euthanasia. If mice can
				spontaneously recover after losing
				20% of their body weight then our
				20% of their body weight, their our
				wouldn't know this if we had to
				wouldn't know this if we had to
				eumanize menn at the 20% weight
				IUSS IEVEI. HOWEVER, WE LITITIK IL
				uninkely they could spontaneously
				recover after losing 30%. Since one
				of our goals is to reverse toxicity,
				we d like to apply our proposed
	Pennell,		EUTHANASIA	treatment at a time when the mouse
1802-35603A	Christopher	Mice	METHOD	could not otherwise recover. 🗆

				For this experimental procedure, it is
				essential that tumor implantation be
				performed in the manner that has
				been established, that is with an
				intact cranium during the injection
				process. A subsequent period after
				tumor implantation with the intact
				cranium must be provided to allow
				for adequate establishment of the
				tumor cells to grow. Therefore, the
				cranioplasty procedure can not be
				performed at the time of tumor cell
				implantation. The tumor cell
				implantation surgery is minimally
				invasive and animals recover
				quickly and do quite well following
				this procedure. Furthermore, pain
				and distress post-operatively
				associated with multiple procedures
				will be mitigated appropriately as
				outlined in the individual procedure
				protocols. (Cranioplasty)
				For this experimental procedure, it
				is essential that tumor implantation
				be performed in the manner that has
				been established, that is with an
				intact cranium (or artificial bone
				implant) during the injection
				process. A subsequent period after
				tumor implantation with the intact
			MULTIPLE	cranium (or artificial bone implant)
1802-35597A	Chen, Clark	Місе	SURGERY	must be provided to allow for
4000 055074		N4:		
1802-3559/A	Chen, Clark	MICE	ENRICHMENT	see protocol for details

				TO OUL KNOWIEUGE, THELE IS NO
				pharmaceutical grade urethane
				available. However, we believe the
				use of urethane is still justified in our
				anesthetized, non-survival surgical
				procedures as alternative
				anesthetics have considerable
				negative effects on physiology of the
				cerebral and cerebellar cortices. For
				example, ketamine is well known to
				block NMDA recentors in the brain
				which are key receptors in the
				nouronal circuitry that we are
				atudving (Dangtagan & Jaratal)
				Studying (Bengtsson & Joniten,
				2007). Barbiturates are also known
				io protoundiy depress cerebenar
				Additionally (Salo, F. et al., 1995).
				Additionally, isonularie over time
				depresses cerebellar function (Loeb,
				A.L., et al., 1990), which is not ideal
				Therefore, wrothere is the best
				Therefore, urethane is the best
				available anesthetic agent for us to
				use to investigate cerebral and
				The procedure will also be carried
				out in 373 ME.
			19 103977978 M	Bengtsson, F. & Jorntell, H.
			NON-	Ketamine and xylazine depress
			PHARMACAUT	sensory-evoked parallel fiber and
	20092 0100 × 24	a	ICAL GRADE	climbing fiber responses. J
1802-35597A	Chen, Clark	Mice	COMPOUNDS	Neurophysiol 2007, 98(3):1697-705.
			SOCIAL	
1802-35597A	Chen, Clark	Mice	HOUSING	see protocol for details
				This procedure will be performed in
				mice with diabetes to measure
				effects of anti-diabetic therapy.
				Random checking will involve
				weekly measurements. For glucose
				tolerance test mice will be bled at 0,
				30, 60, 90 and 120 minutes after
			BLOOD	glucose intake. Each bleed is 5
			COLLECTION	mcroliters only. This is standard
1802-35591A	Czyzyk, Jan	Mice	LIMIT	GTT assay, which allows for
				A single microinjection of siRNA is
				made to block transcription in CFA-
				treated rats. The effectiveness of
			MULTIPLE	siRNA lasts for only a few days and
1802-35558A	Bereiter, David	Rat	SURGERY	cannot be given at the time CFA

				We have received previous
				permission (7.22.13) from IACUC
				that a single dose of ketoprofen is
				sufficient for the surgeries we
			72 HOUR	propose (Intra-cerebral drug
			POST-OP	administration)
				dammistrationy
1802 355584	Bereiter David	Pat		This is a single injection and not a
1002-00000A	Derenter, David	Ιλαί		Everbitant east increases offectively
				makes pharmacoutical grade
				formulations unavailable.
			NON	Dhamaaaautiaal grada farmaulatiana
			INON-	Pharmaceutical grade formulations
			PHARMACAUT	do not exist (Dorsal brainstem
			ICAL GRADE	surface exposure (electrophysiology,
1802-35558A	Bereiter, David	Rat	COMPOUNDS	microdialysis))
				Exorbital gland removal and chronic
				Intra-CNS cannula implantation can
				be performed concurrently.
				(Exorbital Gland removal)
				Exorbital gland removal and intra-
				CNS cannula placement will be
				performed concurrently. This may
				not qualify as multiple survival
				surgeries as there is no inter-
				procedure interval
			MULTIPLE	Exorbital glands are removed on
1802-35557A	Bereiter, David	Rat	SURGERY	day 0, but siRNA cannot be gi∨en
				We have received previous
				permission (7.22.13) from IACUC
				that a single dose of ketoprofen is
				sufficient for the minor surgeries we
				propose. We will change the
				protocol to use carprofen instead.
				We had received permission
				(7.22.13) from IACUC to use
				ketoprofen (2.5ma/ka. jp) but will
				change to carprofen (5 mg/kg, ip)
				(Chronic intra-cerebral quide
			POST-OP	cannula placement for drug
				administration)
1802 355574	Bereiter David	Pat		aanmiistration
1002-0000/A	Derener, Daviu	n val		

				Exorbitant cost increases effectively
				makes pharmaceutical grade
				formulations unavailable
				(Illtraviolet light irradiation)
				Recent exorbitant cost increases
				effectively makes pharmaceutical
				grade formulations unavailable
				(Chronic intra cerebral quide
				cannula placement for drug
				administration)
				administration
				Dharmaceutical grade formulations
1902 255574	Paraitar David	Pot		de pet eviet. Parbiturates are pet
1602-30007 A	Dereiter, Daviu	rai	SOCIAL	uo not exist. Barbiturates are not
1802 355/54		Mice		see protocol for details
1002-00040A	Lee, Anna	MICE		Decapitation will be used for P0 or
				E15 mice using sharp solesors. The
				addition of sodation to those animals
1902 255 45 4		Mico		addition of sedation to these animals
1602-30040A	Lee, Anna	MICE	METHOD	Carrinteriere with the success of
				Working on mice for more than 8
				years and has a high degree of
1001 255204	Chan Vinali	Mino	METHOD	mouse handling technical
1801-30039A	Chen, Xiaoli	MICE		proficiency. In addition, we will
1001 255204	Chan Vigali	Mico		and protocol for details
1601-30039A		MICE		See protocol for details
				(D1 D4) will be dependent of without
				(PT-P4) will be decapitated without
				anestnesia. Rapid decapitation is
				the preferred method to sacrifice
				pups of this age since it is the most
				rapid method, does not introduce
				adverse substances or gases that
	1227 Br. M. 1012	TRE MALE ANTINA ANT AVAILAN	EUTHANASIA	would affect sensitive neonatal
1801-35505A	Ashe, Karen	Mice, Rabbit	METHOD	neurons, and avoids any discomfort
				We are testing a novel
			EUTHANASIA	immunotherapy that may permit
			DEATH/MORIB	animals that appear sick (e.g. due to
	Pennell,		UND	immune-mediated inflammation) to
1801-35448A	Christopher	Mice	ENDPOINT	recover and become tumor-free.
			1111 11111 111 11 11 11 11 11 11 11 11	Cervical dislocation is rapid and
	Pennell,		EUTHANASIA	apparently painless. I have over 30
1801-35448A	Christopher	Mice	METHOD	years experience using this method

				Our research requires euthanization
				by convical dislocation without
				anestrisia. Anestriesia and carbon
				dioxide asphyxiation lead to an
				increase in catecholamine levels,
				which in turn stimulate lipolysis in
				adipose and glycogenolysis in liver.
				These alterations in lipolysis and
				blood glucose interfere with the
				analysis of insulin sensitivity.
				Immediately following euthanization
				mice are bled through the orbital
				plexus. Apesthetics are known to
				piexus. Anesthetics are known to
				which will interfere with our
				experiments. Blood collection will
Contraction and the second	Service discuss as all	18 - 19 x	EUTHANASIA	happen right after cervical
1712-35414A	Kim, Do-Hyung	Mice, Mice	METHOD	dislocation. 🗆
				non pharmaceutical-grade urethane
				is used as an anesthesia in our non-
				survival surgeries. Urethane comes
				in crystal form stored in secondary
				containment at room temperature.
				To prepare urethane from solid
				crystal, in a chemical fume hood.
				0.9g urethane is diluted in 5mL of
				saline and filtered using 0.22 micron
				Milley GP filter. Urethane in liquid
				form is stored at room tomporature
				I rethand is the best and ank antion
				orethane is the best and only option
				for this procedure because results
				will be comparable to previous
				research.
				(DOI:10.1523/JNEUROSCI.4801-
				06.2007 , DOI:
				10.1073/pnas.1520759113))
				(Terminal Epilepsy
				Analysis/electrophysiology)
				, , , ,
				non pharmaceutical-grade urethane
				is used as an anesthesia in our non-
				survival surgeries. Urethane comes
				in crystal form stored in secondary
				containment at reem temperature
				To propage unother from collid
				To prepare ureinane from solid
1710 05 11 10				crystal, in a chemical tume hood,
1/12-35414A	Kim, Do-Hyung	Mice, Mice	COMPOUNDS	0.9g urethane is diluted in 5mL of

				•
				We use either under-gravel filters
				where the sediment waste on the
				gravel is siphoned out monthly or
				above-tank charcoal filter systems
				where the charcoal cartridges are
				changed based on manufacturer's
				recommendations
		Fish (Other)		
		Amphibian		
		(Other) Rentile	SANITATION	We change the bedding
1712-35/130	Mand Sandy	(Other)		material/sand every month or two
17 12-00410A	Maria, Gariay	(Other)		We have a male leopard decko
		Amphibian		Male deckos fight when housed
		(Othor) Pontilo		together and we would prefer not to
1710 051104	Mand Candy	(Other), Repute		
1712-30413A	Mariu, Sariuy			Altered lipelytic activity and
				metapolites have been
				demonstrated in tissues and blood
				after anesthesia and sedation. This
		a	EUTHANASIA	euthanasia method will be used
1712-35406A	Bernlohr, David	Mice	METHOD	when determined to be necessary by
				Cervical dislocation of young poultry
				is necessary because of the
				extremely long time needed to
				euthanize poultry with CO2
			EUTHANASIA	especially up to one week of age
1712-35400A	Noll, Sally	Turkey, Turkey	METHOD	after hatch. They are resistant to
				This procedure only involves
			72 HOUR	puncture of one point of the skin and
			POST-OP	skull and is thus minimally invasive.
	Nakagawa,		ANALGESIA	I have consulted Dr. Nate Koewler
1712-35389A	Yasushi	Mice	POLICY	to confirm that we do not need
				There are no pharmaceutical-grade
			NON-	compound available. (In utero
			PHARMACAUT	electroporation)
	Nakagawa		ICAL GRADE	
1712-353894	Yasushi	Mice	COMPOLINIDS	No pharmaceutical-grade
L11 12 00000A	1 dodoni	miloo		rio pharmaocatoal-grade

				The major endpoint for our research on sickle cell disease (SCD) is microvascular stasis. We have published numerous papers demonstrating that pro-inflammatory compounds increase stasis and anti- inflammatory compounds inhibit stasis in our SCD mouse models. Thus the interventions that we evaluate for treating SCD are all anti-
1712-35371A	Belcher, John	Mice, Mice	72 HOUR POST-OP ANALGESIA POLICY	Use of anti-inflammatory drugs post- surgery will interfere with our measurement of microvascular stasis in our dorsal skin-fold chamber model. Buprenorphine has anti-inflammatory activity (Volker D, Bate M, Gentle R, Garg M. Oral buprenorphine is anti-inflammatory and modulates the pathogenesis of streptococcal cell wall polymer- induced arthritis in the Lew/SSN rat. Lab Anim. 2000 Oct;34(4):423-9). Unfortunately all of the analgesic
1710 252714	Polobor John	Mico Mico	NON- PHARMACAUT ICAL GRADE	The following is our scientific justification for the use of non-pharm grade chemicals. Urethane is widely used as an anesthetic for animal studies because of its minimal effects on cardiovascular and respiratory systems and maintenance of spinal reflexes. Alpha-chloralose is an anesthetic characterized by its ability to maintain animals in physiological conditions though immobilized and anesthetized. In addition, alpha- chloralose induces a loss of consciousness with little influence on either pain responses of
1712-30371A	Deloner, John			on enner pain response or

				Animals will receive virus infusion
				and optical fiber implantation in
				separate surgeries. This is
				experimentally necessary because it
				takes several weeks for AAV genetic
				nowloads to ovprose. For the
				payloads to express. For the
				animais well
				being and to maximize experimental
				success, we minimize the time that
				it carries fiber implants.
				Pain and distress will be monitored
				identically for each individual
				procedure, and animals will receive
				the same
				postoperative care, monitoring, and
				pain management. (Virus Vector
				Infusion (Rats, Survival))
				····· ································
				A subset of animals will receive
				virus infusion and optical fiber
				implantation in separate surgeries.
				This is experimentally necessary
				because it takes several weeks for
				AAV genetic payloads to express.
			MULTIPLE	For the animals' well being and to
1711-35358A	Thomas, Mark	Rat	SURGERY	maximize experimental success, we
	,		SOCIAL	
1711-35358A	Thomas, Mark	Rat	HOUSING	see protocol for details
		Reptile (Other),		
		Reptile (Other),		
		Amphibian	ENVIRONMEN	
		(Other), Frog	TAL	
17 <u>11-35353</u> A	Waye, Heather	(Other)	ENRICHMENT	see protocol for details
	Mereddy,		SOCIAL	
1711-35316A	Venkatram	Mice	HOUSING	see protocol for details
				Post-operative analgesic therapy is
				contraindicated because the
				objective of the procedure is to
			72 HOUR	determine whether the experimental
			POST-OP	treatment causes hypersensitivity.
	Vulchanova,		ANALGESIA	Provision of analgesics would inhibit
1711-35311A	Lucv	Mice	POLICY	the neurochemical changes required
				Cervical dislocation will be
				performed by technically competent
				staff to ensure quick and complete
				euthanasia
				cathanasia.
1711 352864	Garny Many	Mice		Decapitation will be used for
1711-00200A	Ourry, wary	WINCO		Decapitation will be used for

			-	
1710-35273A	Banik, Ratan	Rat	72 HOUR POST-OP ANALGESIA POLICY	We request not to give analgesics because we need for hyperalgesia to fully develop and analgesics may interfere with this process. Analgesic will be used if animals show excessive pain like behaviors (eg. vocalization, restlessness etc.). We and others lab extensively used this model. Based on our experience and available data, animals tolerate well this surgical procedure and do not show excessive pain behaviors. (Plantar skin+muscle incision) We request not to give analgesics because we need for hyperalgesia to fully develop and analgesics may interfere with this process. Capsaicin and methylene blue may provide analgesia in experimental animals. □ Analgesic will be used if animals show excessive pain like behaviors 1. An extensive history of this surgery in mice (since 1999) via the NIH Mouse Metabolic Phenotyping Consortium has shown that more that 24 hours of analgesics is not necessary for the well being of the mouse. The mice do not, except in rare cases, exhibit loss of weight, piloerection, unkempt coat, □ hunched posture. If this occurs additional analgesic will be provided. □ 2. The studies within this protocol are aimed at assessing metabolism. Keeping the use of □ analgesics to a minimum is to avoid motabolic of the patient pat
			72 HOUR POST-OP	metabolic alterations not intended by the experimental design is□
			ANALGESIA	imperative.□
1710-352184	Crawford Peter	Mice	POLICY	*Please note that we would like to
17 10-002 IOA				T lease hole that we would like to
4740.050404		N4:		
1710-35218A	Crawford, Peter	MICE	HOUSING	see protocol for details

	T		T	
				A subset of adult mice animals
				(approximately 40%) will be
				euthanized by cervical dislocation.
				This method is selected because
				does not chemically contaminate
				tissue (including hypoxia and
				acidosis), which is critical for
				metabolic studies involving high
				resolution chemical profiling of
				extracts derived from the tissues
				(e.g., LC/MS metabolomics and
				magnetic resonance spectroscopy).
				Moreover, cervical dislocation
				induces rapid loss of consciousness,
				and is rapidly accomplished.
				As stated on p. 49 of AVMA
				Guidelines for the Euthanasia of
				Animals: 2013 Edition, 'cervical
			EUTHANASIA	dislocation is acceptable with
1710-35218A	Crawford, Peter	Mice	METHOD	conditions for mice Personnel
			WEEKLY	
			WEIGHING	Diets are not restricted.
		Pig	EXCEPTION	(Supplementation of Camelina
1708-35078A	Li, Yuzhi	(Agricultural)	(FOOD/FLUID)	Press-cake in pig diets)
				A pharmaceutical grade is not
				available. As detailed in the SOP,
				urethane is prepared in a fume hood
				with PPE. After weighing the
			NON-	powder, e.g., 4 g, it is placed in a 50
			PHARMACAUT	mL Falcon centrifuge tube and
			ICAL GRADE	distilled water is added for a total
1708-35069A	Kara, Prakash	Місе		volume of 40 mL. The tube is clearly
4700.05000.4	One ithe Orenstern		MULTIPLE	
1708-35068A	Smith, Gordon	Other* (USDA)		see protocol for details
			DOST OD	
1708-350684	Smith Gordon		POLICY	see protocol for details
1700-0000A			NON-	
			PHARMACAUT	
1708-35068A	Smith, Gordon	Other* (USDA)	COMPOUNDS	see protocol for details
11.00.000001	ornan, ooraon		001100	

				8 Weeks in between□
				Bolus injection of ASO has been
				shown to result in more uniform
				delivery to all regions of the mouse
				brain than slower release pump
				delivery, however ASOs degrade
				over time so multiple bolus
				injections may be necessary to
				achieve full therapeutic potential.
				Our mice recover quickly from this
				short survival surgery and any
				additional surgeries will take place
				an absolute minimum of 4 weeks
				apart. In a currently oppoind clinical
				trial for ASO therapy in HD, patients
				receive intrathecal injection every 3
				months ASOs used in that trial are
				similar in chomical composition and
				mochanism of action to the ASOs
				L CCA 2 ASO injection and M D S
				SCA 3 ASO injection and M.R.S.
				12 Weeks in between⊔
				This mouse project has 4 total
				survival surgeries done to each
				mouse. One is conducted at the
				University of Michigan Ann Arbor,
				and then 3 more of the surgeries will
				be performed here at the University
			MULTIPLE	of MN. Due to the nature of the
1708-35065A	Orr, Harry	Mice	SURGERY	disease in these mice, the
			72 HOUR	
			POST-OP	This is a follow up procedure for the
			ANALGESIA	removal of stitches, and is
1708-35065A	Orr, Harry	Mice	POLICY	noninvasive. (Stitch Removal)
				Neonates up to P8 are euthanized
				by decapitation with surgical
				scissors as they do not have mature
			EUTHANASIA	nociceptors and are resistant to
1708-35065A	Orr, Harry	Mice	METHOD	hypoxia. P10 through P17 are
			ANESTHESIA.	Further monitoring of a given mouse
			SURGERY.	is not necessary as the procedure is
			AND POST-	terminal before 15 minutes pass.
			PROCEDURAL	(Non-survival perfusion)
			RECORDKEEP	· · · · · · · · · · · · · · · · · · ·
1708-35065A	Orr. Harry	Mice	ING	This is a follow up procedure for the
	Gallaher		SOCIAL	
1708-35056A	Daniel	Rat	HOUSING	see protocol for details
		10-5110-112-112		

			EUTHANASIA	It is highly possible that the
			DEATH/MORIB	treatments administered to the
			UND	animals could result in adverse
1708-35052A	LeBeau, Aaron	Mice	ENDPOINT	health effect such as drug-related
				In order to achieve successful
				engraftment of muscle stem and
				progenitor cells the muscle has to
				undergo an injury two days before
				transplantation to induce an
				essential injury response in the
				tissue. Cryo-injury or muscle
				injection injury will be performed as
				described in the previous section,
				pain and distress in the animals will
				be monitored for 3 days post-
				procedure. Because the effects of
			MULTIPLE	analgesic compounds on muscle
1708-35046A	Kyba, Michael	Mice, Mice	SURGERY	regeneration are unknown, and
				Because the effects of analgesic
				compounds on muscle regeneration
				are unknown, and because we want
				to examine the role of inflammatory
				cells in skeletal muscle
				damage/regeneration, we cannot
				us analgesics for this procedure.
				(Mascle mjury by injection)
				Because the entects of analyesic
				compounds on muscle regeneration
1700 250 464	Kuba Mishaal	Miss Miss		are unknown, and because we want
1708-30046A	Kyba, Michael	Mice, Mice	POLICY	
				our objective is not to have any
				due te unferseen effects of
				national treatments death is
1709 350 464	Kuba Michaol	Mico Mico		particular treatments, deatings
1700-33040A	rtyba, michael			Embryos 15 days of destation or
1708-350464	Kyha Michael	Mice Mice	METHOD	dreater will be decapitated
1700-00040A	rtyba, mionaei		SOCIAL	greater win be decapitated.
1708-35046A	Kyba Michael	Mice Mice	HOUSING	see protocol for details
1708-35046A 1708-35046A 1708-35046A 1708-35046A	Kyba, Michael Kyba, Michael Kyba, Michael Kyba, Michael	Mice, Mice Mice, Mice Mice, Mice Mice, Mice	SURGERY 72 HOUR POST-OP ANALGESIA POLICY EUTHANASIA DEATH/MORIB UND ENDPOINT EUTHANASIA METHOD SOCIAL HOUSING	regeneration are unknown, and Because the effects of analgesic compounds on muscle regeneration are unknown, and because we want to examine the role of inflammatory cells in skeletal muscle damage/regeneration, we cannot us analgesic for this procedure. (Muscle injury by injection) Because the effects of analgesic compounds on muscle regeneration are unknown, and because we want to examine the role of inflammatory Our objective is not to have any moribund animals, but on occasion, due to unforseen effects of particular treatments, death is possible, and we would euthanize Embryos 15 days of gestation or greater will be decapitated.
				For arenavirus (PICV and LCMV)- infected mice, we will use death as endpoints, which is the established
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				practice for LCMV-mouse model
				((von Herrath and Whitton, Animal
				choriomeningitis virus. Curr Protoc
				Immunol. 2001. Chapter 19: Unit
				19). After LCMV infection
				Intracranially, mice will begin to
				hunched posture, reduced mobility)
			EUTHANASIA	around day 6 to 7 post-challenge,
			DEATH/MORIB	and will succumb rapidly thereafter.
1708-35044A	Liang Yuving	Mice	UND ENDPOINT	Death may be delayed in certain denetically modified strains and/or
	Liang, raying			generically meaned channe analor
				vve nave used avertin routinely in
				2014). It is easier to store and use in
				the lab.
				Avertin will be prepared and stored
				using these guidelines:
				1.Sterlie filter with 0.2 micron filter
				conditions
				3.Store in the dark bottle or foil
				covered container
				4. Store stock and working stock
				5. Do not use if the solution
			NON-	becomes discolored or has a
			PHARMACAUT	precipitate
1708 350364	Dana Honaha	Mico		6. Check pH before each use and
1700-30030A	רמווץ, הטווטטט	INICE	SOCIAL	
1708-35036A	Pang, Hongbo	Mice	HOUSING	see protocol for details

				Plood chimorism acours at 10 to 14
				dove and we will be studying
				days and we will be studying
				mammary tumor progression at
				different time points (up to 13
				months). Keeping the mice joined
				together for the duration of the
				study, after blood chimerism has
				occurred, would put undue stress on
				the mice. Therefore, it is necessary
				for the mice to undergo a second
				survival surgery, 2 to 4 weeks after
				the initial surgery, to separate the
				joined mice. After the joined mice
				are separated, they will recover for 1
				to 2 weeks and cohabitate together
				for the duration of their different
				time points (up to 13 months). At
				which point, they will be euthanized
				for sample collection. (Parabiosis)
				Blood chimerism occurs at 10 to 14
				days and we will be studying
				mammary tumor progression at
				different time points (up to 13
				months) Keeping the mice joined
				together for the duration of the
	Morrie			study, after blood chimerism has
1708-350244	Rebecca	Mice	SURGERY	accurred would put undue stress on
11 00 0002 // (				No adverse affects are expected
				with the addition of antibiotics which
			WEEKLY	are known to be well tolerated
			WEIGHING	(Antibiotic Feeding)
			EXCEPTION	(, indefetto r courrig)
1708-35014A	Herzberg, Mark	Mice	(FOOD/FLUID)	No adverse impact on animal

				In our studies, we find that tumor
				bearing animals may develop
				involutions and ulcerations of the
				tumor as part of successful
				experimental treatment. Thus, it is
				important to follow these animals to
				document successful therapy.
				Animals with involuted/ulcerated
				tumors will be treated with topical
				collasate at least every other day
				until resolved. This protects the
				compromised tissue from infection
				or trauma, prevents the involutions
				from progressing, and in some
				instances allows the involutions to
				heal. Allowing these animals to
				remain on study prevents projects
				from having to be repeated on
				additional animals. Laboratory
				personnel will provide treatments in
				consultation with the veterinary staff.
				(Subcutaneous tumor induction)
				, ,
				In our studies, we find that tumor
				bearing animals may develop
				involutions and ulcerations of the
				tumor as part of successful
				experimental treatment. Animals
				with involuted/ulcerated tumors will
			TUMOR	be treated with topical collasate at
	Panvam		FNDPOINT	least every other day until resolved
1708-35013A	Javanth	Mice	CRITERIA	This protects the compromised
		Pia		
		(Biomedical).		
		Other* (USDA).	NON-	
		Sheep	PHARMACAUT	
		(Biomedical).	ICAL GRADE	
1707-35001A	laizzo, Paul	Dog	COMPOUNDS	see protocol for details

1707-349954		Nonhuman Primate (Macagues)	MULTIPLE	Animals will arrive on the protocol with an implanted vascular access port (VAP). The placement of a VAP is considered a minor surgical procedure (peripheral limb, not entering a major body cavity), essentially similar to routine catheter placement with the notable exception that the access site is placed underneath the skin. VAPs improve safety, and increase animal comfort when intermittent or recurrent venous access is required by avoiding painful catheter starts'
1707-34988A	Parr Ann	Rat	MULTIPLE	The rat must first be injured and recover to model a spinal cord injury so that we can test functional alterations after motor cortex stimulation. Pain and distress will be controlled through analgesics and antibiotics. (Motor Evoked Potential (MEP) Recording) The rat must first be injured and recover to model a spinal cord injury and viral vectors allowed time for transduction so that we can test optogenetic modulation of the motor cortex. Pain and distress will be controlled through analgesics and antibiotics.□ (AAV viral injection into the motor cortex and electrode implantation) The rat must first be injured and
1707-34943A	Stephens, David	Bird (Other), Bird (Other)	SOCIAL HOUSING	For our experiments we need to carefully control and monitor the amount of food each individual eats per day. This level of control would be impossible if animals were housed in the same cage. A second problem with social housing is territoriality. Though starlings are known for their spectacular grouping behavior, they become territorial when in breeding condition (typically

				The goal of one study is to
				determine plasmalemma excitability
				and the only way to do this is my
				activating muscle confraction via the
				implanted nerve cuff and
				simultaneously recording electrical
				activity of the muscle via the EMG
				electrodes. Thus, this second
			MULTIPLE	surgery to implant EMG recording
1707-34941A	Lowe, Dawn	Mice	SURGERY	electrodes, which is minimally
			ENVIRONMEN	
	74 Dect	55 . Sec.	TAL	
1707-34941A	Lowe, Dawn	Mice	ENRICHMENT	see protocol for details
				Cervical dislocation will be done on
				any poults to be euthanized before 3
				days of age per the unit SOP. Poults
				are resistent to carbon dioxide
				asphyxiation due to their hatching
				under a high CO2 environment and
				cervical dislocation is needed in
				order to have the poults euthanized
			EUTHANASIA	in a short time period. Cervical
1706-34934A	Johnson, Tim	Turkey, Turkey	METHOD	dislocation is allowed for poultry

				We request an exception to allow for
				our use of AAV viral infusion and
				surgical optic fiber implantation for
				optogenetic studies. Use of two
				separate surgical procedures $(\Delta\Delta V)$
				$\pm$ fiber implantation) is beneficial, as
				it will reduce the emount ourgen.
				it will reduce the amount surgery
				done at one time (reduces potential
				for tissue trauma), minimizes the
				need for anesthesia supplements
				during surgery, allow animals to fully
				recover between each class B
				surgery. It also maximizes the
				potential for animals to reach the
				study completion.
				Maximal viral-mediated gene
				transfer occurs approximately 3-8
				weeks after infection, depending on
				the brain region and neuronal
				process (cell body versus terminals)
				being targeted. In vitro and in vivo
				studies begin at this time, but
				require up to 1 weeks of additional
				time for completion. However
				antical implete are delicate, and
				optical implants are delicate, and
				generally only have reliable function
				for ~ 4 weeks. Therefore, is
				advantageous to implant them at a
				time when viral expression is
				maximal. This method will prevent
				subject loss due to optical implant
	Bartolomucci,		MULTIPLE	failure, and will substantially reduce
1706-34930A	Alessandro	Mice	SURGERY	the overall number of animals
	Bartolomucci,		SOCIAL	
1706-34930A	Alessandro	Mice	HOUSING	see protocol for details
				I have discontinued use of
			EUTHANASIA	Isofluorane, since it is not necessary
1706-34929A	Everson, Paul	Mice	METHOD	prior to euthanasia by cervical
		Horse, Cow	ANESTHESIA,	
		(Biomedical),	SURGERY,	
		Pig	AND POST-	
		(Biomedical),	PROCEDURAL	A sedative is administered prior to
	Clarkson.	Goat, Sheep	RECORDKEEP	IV injection of pentobarbitol for
1706-34914A	Christina	(Biomedical)	ING	euthanasia.
		(		These endpoints are highly
				susceptible to environmental
				changes: the use of anesthetics or
				CO2 has been demonstrated to alter
				the lovels of pourotransmitters as
				approved to convice dislocation
	Andorson			Dereappel in this group have wart
4700 0 40 40 4	Anderson,		LUTHANASIA	Personner in this group have years
1706-34913A	Grant	MICE	METHOD	of experience with cervical

The second second	Anderson,		EUTHANASIA	CO2 doesn't work well with pups so
1706-34912A	Grant	Rat, Mice	METHOD	we will use decapitation after rat
	1		SOCIAL	
1706-34906A	Lee, Anna	Mice	HOUSING	see protocol for details
1706-34906A	Lee, Anna	Mice	EUTHANASIA METHOD	Decapitation will be used for P0 or E15 mice using sharp scissors. The addition of sedation to these animals caninterfere with the success of
1706-34904A	Gupta, Kalpna	Mice	72 HOUR POST-OP ANALGESIA POLICY	In this study we are examining the effect of topical opioids on wound healing. Therefore, comparisons will be made between vehicle (without opioid or any analgesic) and opioid treated wounds on mice. The vehicle group therefore cannot receive any analgesics because that will interfere with the research question. We will make every effort
1706-34904A	Gupta, Kalpna	Mice	EUTHANASIA DEATH/MORIB UND ENDPOINT	Since to goal of Study #1 is to examine the effect of chronic treatment with morphine or cannabinoids on survival, it is necessary that the animals be
1706-34898A	Graham, Melanie	Mice	MULTIPLE SURGERY	Following transplant, after demonstration of a prolonged period of normoglycemia or insulin reduction greater than 50%, animals will undergo graft explant by removing the graft and following up the animals subsequent reversion to the diabetic state the functional state can be (or not) attributed to the graft. Animals that do not
1706-34897A		Nonhuman Primate (Macaques)	MULTIPLE SURGERY	see protocol for details

					Endpoint criteria that we will follow
					for this study are as follows:
					<ol> <li>Inability to move or perform</li> </ol>
					normal body functions
					2. Skin necrosis and/or ulceration
					over the tumor 🗆
					<ol><li>Clinical signs indicating tumor has</li></ol>
					metastasized (ie seizures, swollen
					abdomen, labored breathing and
					secondary masses)□
					Endpoint criteria that we cannot
					follow: 🗆
					1. At the end of the study, mice
					consume less food and are not as
					active as mice without tumors.
					<ol><li>Tumor burden that causes a</li></ol>
					significant loss of body mass -
					adipose and muscle mass. 🗆
					Scientific justification for not
					following these criteria:
					Cancer cachexia is a clinical
					syndrome characterized by
					weakness, fatigue, poor appetite,
					and muscle and adipose wasting
					(Fearon KC et al. Am J Clin Nutr
					2006;83:1345-50). The model we
				TUMOR	propose to use in our studies - male
	Metzger,			ENDPOINT	CD2F1 mice with colon-26
1706-34883A	Joseph	Mice, Rat	, Mice	CRITERIA	adenocarcinoma tumors is one of

 Metzger,	 <b>D</b> (	 EUTHANASIA	can be placed in a petri dish at the
			progression of cachexia in this model. Because mice are not housed individually, food intake for the cage can be monitored but not for individual mice. Moistened food
			Because the experiments specifically involve using a model of cancer cachexia, mice are expected to lose a significant amount of body weight, including both adipose and muscle mass. Therefore we cannot follow the first criteria for euthanasia of weight loss in the animals. Additionally, anorexia (decreased food intake) is a natural part of the

				Some mice will receive multiple
				procedures: the most complicated of
				these experiments are mice that are
				made to be radiation bone marrow
				chimorae, followed by the optic
				chimeras, followed by the optic
				I he bone marrow chimeras are
				done to introduce specific
				populations of immune cells (from
				Tg mice) that are sensitive to DTx,
				so that they can be eliminated upon
				toxin administration. The optic nerve
				crush is then done to stimulate
				recruitment of immune cells to the
				retina. In this protocol, we use mice
				that are > 6 wks of age to reduce
				radiation sensitivity. We use a large
				inoculum of BM cells (5-10 x 10^6)
				to promote rapid engraftment and
				recovery. Our previous experience
				with the diphtheria toxin allows us to
				use the minimum dose that is
				effective for either intraocular or iv
				administration, and the kinetics of its
				activity. We are now verv
				experienced in the optic nerve
				crush, and rarely experience
				problems []
				(3) Optic nerve crush)
	Gregerson		MULTIPI F	
1706-34882A	Dale	Mice, Mice	SURGERY	Many mice will receive multiple

				We originally requested an
				exception for use of post-op
				buprenorphine on $2/2/2011$ It was
				iustified as follows: "There is a
				growing literature on the beneficial
				effects of opioid receptor agonists
				on the survival of stressed or injured
				neurons. Since the goal of this
				research project is to discover the
				activities of endogenous factors that
				support neuron survival, these new
				finding raise the possibility that use
				of buprenorphine, or other opiates.
				will complicate our findings. Further,
				a recent paper suggests that opiates
				given at the time of injury, which
				would be post-op in our case, have
				the most significant effect. As a
				result, we request suspension of use
				of buprenorphine or similar agents
				post-op for the survival surgeries.
				Use of anti-inflammatories as post-
				op analgesics has been considered,
				including NSAIDS and
				glucocorticosteroids, but they have
				also been shown to affect neuron
				survival post-injury. For these
				reasons, we request exception to the
				common practice of using opiates
			72 HOUR	for this particular post-op
			POST-OP	procedure." 🗆
	Gregerson,		ANALGESIA	
1706-34882A	Dale	Mice, Mice	POLICY	In a series of letters, (on hand) from
1705 0 100 1 1			TAL	
1705-34831A	Kotz, Catherine	MICE		see protocol for details
1705 0 4004 4		N4:	SOCIAL	a a success and face alacta ital
1705-34831A	Kotz, Catherine	wice	HOUSING	see protocol for details

				In this model of cerebral malaria,
				there is some heterogeneity in the
				timing and incidence of severe
				disease (requiring outbanasia, since
				there are no defined treatments to
				reverse the lethal disease at that
				point). However, we need to
				distinguish between mice that
				become moribund and those with
				transient, mild illness, which may
				recover from the disease (for
				example, following the proposed
				cytokine treatments). We and
				others have explored use of other
				clinical features that would predict
				the inevitable onset of lethal disease
				prior to the criteria used (for
				example, we have collaborated with
				Dr. Aaron, Johnson at Maya Clinia
				Dr. Aaron Johnson at Mayo Cinic,
				Who has used WIRI scans of AINKA
1705 0 1000 1	Hamilton Hart,			infected mice as a potential way to
1705-34830A	Sara	Місе	ENDPOINT	anticipate the onset of disease
				The procedure proposed is not a
				restriction in access to water, but
				instead is provision of water
				containing antibiotics. We have
				never seen evidence that decrease
			WEEKLY	their water intake in the past when
			WEIGHING	using this procedure (when water
	Hamilton Hart		EXCEPTION	bottles were the norm - the
1705-34830A	Sara	Mice	(FOOD/FLUID)	consumption rate seemed similar to
			FUTHANASIA	For myositis studies, animal inability
				to ambulate to food or water will
				constitute an indication for
1705 249064	Determon Frik	Mico		immediate authonopia
170J-34620A	Feleison, Enk	MICE	ENDFOINT	
				we request use of non-
				pharmaceutical socium
				pentobarbital in these terminal
				animals, because the availability of
				pharmaceutical grade pentobarbital
				(Nembutal) is unreliable and
				inconsistent, rendering it essentially
			NON-	unavailable for use in animal
			PHARMACAUT	research. Solutions will be prepared
	Honda,		ICAL GRADE	using sterile procedures and sterile
1705-34824A	Christopher	Rat	COMPOUNDS	diluents (saline, ethanol, propylene
	Haskell-		SANITATION	· · · · · · · · · · · · · · · · · · ·
1705-34823A	Luevano Carrie	Mice	FREQUENCY	see protocol for details
11 00 0 1020/1				

			ENVIRONMEN	
	Haskell-		TAL	
1705-34823A	Luevano, Carrie	Mice	ENRICHMENT	see protocol for details
	Haskell-		SOCIAL	
1705-34823A	Luevano, Carrie	Mice	HOUSING	see protocol for details
				Using anesthesia interferes with
				gene expression for certain brain
				tissues collected and glucose
				measurements. For Decapitation of
				the mice, scissors are preferred to
				limit damage that can occur to the
				brain when using a guillotine and
	Haskell-		EUTHANASIA	scissors are easier to clean in
1705-34823A	Luevano, Carrie	Mice	METHOD	between animals to remove blood
				Requesting an exemption to social
				housing for education birds. Raptors
			ENVIRONMEN	are not social animals. They are
		Bird (Other),	TAL	predators and, as such, may present
1705-34793A	Ponder, Julia	Bird (Other)	ENRICHMENT	risk to each other in a group housing
				Requesting an exemption to social
				housing. Raptors are not social
		Bird (Other),	SOCIAL	animals. They are predators and, as
1705-34793A	Ponder, Julia	Bird (Other)	HOUSING	such, may present risk to each other

				Post-operative pain management
				has been addressed by our
				collaborator, Dr. Dale Gregerson. In
				their protocol, Dr. Gregerson
				explains the following. Regarding
				use of buprenorphine: An
				ophthalmologist (cornea surgeon).
				Dr. Steven Kaufman, at U of M. and
				also Dr. Roland Gunther, have been
				consulted about post-surgical
				analgesia for the intraocular
				injections we have proposed. Both
				confirm that only topical
				proparacaine is needed. We
				proposed to use topical
				proparacaine, and also ketamine &
				xylazine for restraint as the
				procedure requires precision. No
				post-operative analgesia is needed
				For reference, note that patients do
				not receive general anesthesia or
				post-op meds for such injections
				even when done repeatedly and for
				months to years. Only topical
				anesthetic drops, such as
				proparacaine is used for them. For
				these reasons we see no need for
				post-op meds in these mice
				Further there is a growing literature
			72 HOUR	on the beneficial effects of opioid
			POST-OP	receptor agonists on the survival of
	Ferrington.		ANALGESIA	stress or injured neurons. Since the
1704-34752A	Deborah	Mice	POLICY	goal of this research is to discover
				Any chemicals used for euthanasia
	Ferrington,		EUTHANASIA	and/or sedation may affect the cells
1704-34752A	Deborah	Mice	METHOD	and inhibit the success of cell growth

				In some situations, mice will have
				two surgeries performed intrarenal
				(i.r.) tumor implantation followed by
				i.r. immunotherany treatment. The
				I.r. Immunotherapy treatment. The
				second survival surgery is
				necessary to facilitate injection of
				the immunotherapeutic directly into
				the tumor-bearing kidney. It is
				important to emphasize that all
				injections will be through the
				peritoneum. The peritoneum will not
				be opened in any mice.
				(Intratumoral Therapy/Treatment)
				In some situations, mice will have
				two surgeries performed, intrarenal
				(i.r.) tumor implantation followed by
				i.r. immunotherapy treatment. The
	Griffith.		MULTIPLE	second survival surgery is
1704-34740A	Thomas	Mice	SURGERY	necessary to facilitate injection of
				The surgical procedure involves a
				simple skin incision to administer
				the intratumoral therapy/treatment
				into the kidney through the intact
				peritoneum. No incision is made in
				the peritoneum. The skin incision is
				then sealed with Vetbond. We will
				administer Bunivicaine (1.25 mg/kg)
				at the incision site on the day of
				at the inclusion site on the day of
				surgery. We are requesting only a
				Single aurillistration of Bupfvicame.
				I his decision was reached after
				consultation with CCRB area
				veterinarians Dr. C. Sivula and Dr.
				S. Hashway. (Intratumoral
				Therapy/Treatment)
				The surgical procedure involves a
				simple skin incision to administer
				the intratumoral therapy/treatment
				into the kidney through the intact
			72 HOUR	peritoneum. No incision is made in
			POST-OP	the peritoneum. The skin incision is
	Griffith			then sealed with Vetbond We will
1704-347404	Thomas	Mice	POLICY	administer Bunivicaine (1.25 mg/kg)
	Griffith		FUTHANASIA	All staff have been trained in and
1704-347404	Thomas	Mice	METHOD	are competent at cervical
1104-04/40A	rnomas	MICE		are competent at cervical

1704-347404	Griffith, Thomas	Mice	WEEKLY WEIGHING EXCEPTION	Since we are not restricting the amount of food/water, but rather changing the type of food they eat to one that most likely will increase their body weight, we will not be
1704-34740A	Thomas	MICE		their body weight, we will hot be
				Two surgeries are required as part of the same project: 1 epilepsy induction surgery and 2 implantation surgery. Epilepsy induction must be done separately from implantation because it is essential for rat to be quickly recovered for the kainic acid to work properly. (Chronic epilepsy induction)
			MULTIPLE	Two surgeries are required as part of the same project: 1 epilepsy induction surgery and 2 implantation surgery
1704-34730A	Netoff, Tay	Rat	SURGERY	Epilepsy induction must be done
1704-34729A	Mashek, Douglas	Mice, Mice	ENVIRONMEN TAL ENRICHMENT	see protocol for details
	Mashek.		SOCIAL	
1704-34729A	Douglas	Mice, Mice	HOUSING	see protocol for details
	Kawakami,	Fish (Zebra	72 HOUR POST-OP ANALGESIA	For zebrafish, the same treatment for other animals, such as mice, are not available, because zebrafish are aquatic animals. Thus, analgesia cannot be provided to zebrafish. If fish show distress or abnormal behavior after surgery (floating belly
1704-34728A	Y asuhiko	TISN)	POLICY	up or random swimming among the

	1		I our studios rolato to the ottoots of
			our studies relate to the effects of
			respiratory motor control and
			neuroplasticity. As most soy storoids
			are produced in the gapade
			are produced in the gonada,
			removal of the gonads is vital to
			answer rundamental experimental
			questions. Removing the gonads
			and allowing for a week of recovery
			creates a "new baseline" or reduced
			sex steroid production and
			circulation for which to conduct our
			studies. These procedures are have
			significant scientific ment and are
			considered standard procedures in
			the study of sex steroids. Rats
			receiving gonadectomies (or snam
			surgenes) receive pain medication
			at the time of surgery to minimize
			discomfort. The procedures are very
			rast (5-10 minutes per rat) allowing
			rais to recover from anestnesia
			anticipated. (Gonadectomy)
			As described with gapadastamy
			As described with gonadectority,
			our studies center on the effect of
			sex steroids in respiratory motor
			control and neuroplasticity. Tonowing
			removal gonads, our capacity to
	Dougharty		manner is critical to interpretation of
1704 247044	Dougherty,	Det	manner is childer to milerpretation of
17 04-347 248		rtal	Adoquato apalgosio offosto far thasa
			routing and yory brief, procedures
	Doughorty		are accomplished in a single pro-
1704 247244	Brondon	Det	are accomprished in a single pre-
1704-34724A	Dienuari	ral	operative auministration.

				We may utilize non-pharmaceutical- grade urethane as all acute neurophysiological experiments are terminal. Transition from inhaled isoflurane to Urethane is necessary for studies of respiratory neurophysiology because isoflurane is a profound respiratory depressant, while urethane maintains long- lasting anesthesia with minimal effect on cardio-respiratory function. (Acute neurophysiological measures)
			NON-	We may utilize non-pharmaceutical- grade Pancuronium Bromide if/when pharmaceutical grade is unavailable. The pharmaceutical grade version of this compound is currently available through only one
	David and a			vendor (Pfizer) and is frequently
1704-34724A	Brendan	Rat	COMPOUNDS	a time. Also, this compound is stable
	Dougherty,		SOCIAL	, ,
1704-34724A	Brendan	Rat	HOUSING	see protocol for details
				This procedure will be used in conjunction with the focused ultrasound procedure. (Ultrasound marker)
1704 247114	Ebbini Emad	Pot Mico		This surger ( will be used in
1704-34711A	Ebbini, Emaŭ	rai, Mille	JUNGERT	The procedure proposed is not a
			WEEKLY WEIGHING EXCEPTION	restriction in access to water, but instead is provision of water containing antibiotics. We have never seen evidence that decrease their water intake in the past when
1703-34693A	Hart, Geoffrey	Mice	(FOOD/FLUID)	using this procedure (when water

				The placement of the constrictor on
				the LAD is an essential component
				to the creation of hibernating
				myocardium. Pain and distress will
				be minimized by using a multi modal
				,balanced anesthetic protocol. The
				pig are socialized with with people to
				decrease the fear response before
				and after the procedure to ease in
				handling and transport. The pigs are
				monitored closely in recovery until
				they can stand on their own.
				(Thoracotomy - LAD Hibernation)
				,
				The revascularization procedure is
				an essential component of this study
				to evaluate the effects of bypass on
				hibernating myocardium and how
				stem cells or mitochondrial
				transplant may serve as an
				adjunctive therapy Pain and
	Kelly.	Pia	MULTIPLE	distress will be minimized by using a
1703-34685A	Rosemary	(Biomedical)	SURGERY	multi modal, balanced anesthetic
1100 0 1000/ (	rteeenary	Distriction	OUNCEIN	One prior surgery to implant
	Stromnes		MULTIPLE	orthotopic tumors into the pancreas
1703-34658A	Ingunn	Mice Mice	SURGERY	will be performed 80-120 days prior
1100 0 1000/	Ingaint		OUNCEIN	Frogs will be euthanized by double-
				pithing followed by decapitation. We
				have to resort to physical methods
				for euthanasia as chemical methods
				such as barbiturates tricaine or
		Frog (Other)		benzocaine produce long-lasting
	Cvetanovic	Rat Guinea	FUTHANASIA	blockade of voltage-dependent and
1703-346314	Marija	Pig Mice	METHOD	ligand-dependent ion channels
1703-34631A	Cvetanović, Marija	Rat, Guinea Pig, Mice	METHOD	ligand-dependent ion channels

				In the mice that are undergoing
				multiple survival procedures, we are
				interested in studying the effect of
				glial rectifying potassium channel
				(Kir4.1) overexpression on seizure
				activity. To ensure that expression
				of this protein is localized to specific
				brain regions, it is necessary to
				stereotaxically inject recombinant
				$\Delta \Delta V$ directly into the brain
				Stereotavic intracranial rAAV
				injection is a relatively cofe method
				or inducing gene expression with
				great spatiotemporal specificity and
				few, if any, side effects (Stoica et al.
				Curr Protoc Microbiol 2013. Ch 14,
				Unit 14D.5). We suspect that mice
				injected with rAAV containing Kir4.1
				will exhibit some seizure activity, but
				no other detrimental effects are
				anticipated from this procedure.□
				As part of this project, it is essential
				to determine not only if
				overexpression of Kir4 1 leads to
				seizures, but whether this channel
				may drive subclinical enilentic
				anikes and other other aborrant
				spikes and other other abenanic
4700 0 40074		N 4:		FEO Instructure of EEO
1703-34627A	vossel, Kelth	MICE	SURGERY	
				E18-21 mouse embryos and
				neonatal mouse pups P0-1 days will
				be euthanized by rapid decapitation
				using a large surgical-grade scissors
				without anesthesia. Rapid
				decapitation is the preferred method
				to sacrifice pups of this age since it
				is the most rapid method, does not
				introduce adverse substances or
				gases that would affect sensitive
				neonatal neurons, and avoids any
				discomfort associated with
			FUTHANASIA	injections. All individuals performing
1703-346274	Vossel Keith	Mice	METHOD	decapitation without anesthesia will
1100 040277			SOCIAL	decupitation without anostrosia will
1703-346274	Vossel Keith	Mice		see protocol for details
1700-04027A				
4702 040074		Minn		
1703-34627A	vossei, Keith	wice	ENRICHMENT	see protocol for details

				Avertin is not available as a
				pharmaceutical grade agent. We
				have used Avertin as an anesthetic
				for many years and find that it very
				quick and effective relative to the
				other anesthetics. It is a reliable
				agent for inducing anesthesia with
				low risk of respiratory suppression or
				occupational exposure in
				comparison with using inhaled
				a pestbetics for inducing apestbesia
				Avertin stock colution bas an
				assigned expiration date of 6
			NON	months and is stored at 4 degrees
				Coloius protected from light
				Censius protected from light.
1702 246074	Vacal Kaith	Mino		working solutions of Avenum will be
1703-34627A	vossei, Keitn	wiice	COMPOUNDS	made in a biosafety cabinet and will
				For the study of mecannisms
				analgesia we need to both induce a
				state of hypersensitivity (reflective
				of neuropathic pain) requiring
				peripheral nerve injury surgery and
				then later implant spinal
				microdialysis fibers in order to
				collect neurotransmitters during and
				immediately following application of
				electroacupuncture. These
				procedures will be separate by a
				week.
	Fairbanks,		MULTIPLE	
1703-34617A	Carolyn	Mice, Rat	SURGERY	For the study of the efficacy of
				Subjects with peripheral neuropathy
	T - i - i - i - i			cannot receive post-operative
1700 0 10 17 1	Hairbanks,		ANALGESIA	anaigesics because these will inhibit
1703-34617A	Carolyn	Mice, Rat	POLICY	the spinal neuroplasticity intended to
				vve are trained and experienced in
				the appropriate technique and
				anesthetization and sedation
				elevates the stress to the subject.
	Fairbanks,		EUTHANASIA	We only intend to use cervical
1703-34617A	Carolyn	Mice, Rat	METHOD	dislocation as an emergency method

				In some mice or rate, sciptic perve
				injection of AAV viral vectors or
				brain injections of viral vectors or
				neuronal tracers will precede (1-6
				weeks) or follow (2-4 weeks) Spared
				Nerve Injury. The addition of these
				injections are not expected to
				exacerbate the level of pain and
				distress associated with spared
				nerve injury. 🗆
				The rationale for administering the
				viral vector intraneurally is that
				Intraheural delivery will larget only
				delivery will be compared to
				intrathecal delivery which will result
				in transduction of both spinal cord
				and sensory neurons. The
				comparison between the two routes
				of delivery will provide information
				about the relative contribution of
				spinal and sensory neurons to the
				pain mechanisms under
				investigation. 🗆
				The rationale for injection in VPL or
				parabrachial nucleus prior to SNI is:
				1) to label projection neurons for
				Identification in subsequent ex vivo
	Vulabanava			physiological experiments, and 2) to
1703-346164	Vuichanova,	Mice Rat	SURGERY	projection neurons for subsequent
1700-040107	Lucy	Miloe, Rat	OUNCENT	projection neurons for subsequent
				Post-operative analgesic therapy is
				contraindicated because the
			72 HOUR	objective of the procedure is to
	1867 B. 1710.		POST-OP	determine whether the experimental
	Vulchanova,		ANALGESIA	treatment causes hypersensitivity.
1703-34616A		Mice, Rat		Provision of analgesics would inhibit
4700.040404	vuichanova,	Mine Det	SOCIAL	and much californitation
1703-34616A	LUCY	iviice, Rat	HOUSING	see protocol for details

1703-34616A	Vulchanova, Lucy	Mice, Rat	NON- PHARMACAUT ICAL GRADE COMPOUNDS	Exception is requested for the use of non-pharmaceutical grade urethane for non-survival surgeries. In our experience, this usually produces a very stable and long-lasting (8-16 hours) state of anesthesia, which is the reason for choosing this non- pharmaceutical grade anesthetic
				The rat must first be injured and recover to model a spinal cord injury so that we can test our cell transplants. Pain and distress will be controlled through analgesics and antibiotics. (Spinal Cord injury and Scaffold Transplantation)
				Three survival surgeries are necessary because a spinal cord injury is necessary to evaluate whether sNPC/scaffold transplantation is effective at resolving the CNS deficits associated with spinal cord injury. Then, anterograde axonal tracing is utilized to confirm that transplanted
1702-34614A	Parr, Ann	Mice, Rat	SURGERY	synaptic connections with the

				Dumke will follow INAD policy on
				anesthesia data recording (added as
				attachment). We are inducing minor
				sedation to make fish handleable, so
				their time in anesthesia until
				recovered is expected to be less
				than 15 minutes. INAD policy states
				that individual fish records do not
				need to be collected for fishes
				immersed in 40 mg/L solution,
				become handleable within 5 minutes
				and are therefore removed from
				immersion, and time to recovery
				does not exceed 20 minutes (pg. 11
				of INAD:AQUI-S 20E - Study
				Protocol). Deviations outside these
				parameters will be recorded to
				comply with INAD program. At
				minimum, we will be collecting and
				reporting information on dosage, fish
				species, numbers of each species
				treated, and whether the outcome
				was satisfactory (i.e. did fish
				become handleable within 5 minutes
				and did they recover within 20?).
				We will start with 40 mg/L solution
			ANESTHESIA,	strength but can adjust in the field.
			SURGERY,	For example, trout may enter
			AND POST-	unintended deep sedation in a 40
			PROCEDURAL	mg/L solution, and non-trout fishes
			RECORDKEEP	may take longer than 5 minutes to
1702-34612A	Brady, Valerie	Fish (Other)	ING	attain adequate sedation. If that

	are trung to
determine whether	and how the
genetic alterations of	n nometheren ( will
previde enviour ive	lemotrierapy will
mouse conorts. In o	rder to generate
a credible mouse su	irvival curve, we
need to observe the	mouse cohorts
up to "natural death	'. However,
these mice are valu	able and needed
for pathological and	biochemical
analyses, we canno	t allow them to
die spontaneously a	nd risk organ
deterioration. Rathe	r, we will harvest
mice when they are	determined to
be moribund-"close	to death
endpoint". One Crite	erion from UMN
IACUC guideline wi	I potentially
affect our studies to	observe
survival benefits. Ba	ased on "UMN
IACUC Euthanasia	Guideline'', we
have provided a ver	y strict criteria to
determine the morit	ound state, and
these must include	at least one of
the first two criteria	and at least one
of the three remaini	ng criteria: (1)
progressive weight	reductions up to
10% of body weight	measured on
two separate occasi	ons over a
EUTHANASIA period of one week:	(2) sudden.
	loss of at least
UND 10% of body weight	over a period of
1702-34610A Deng, Yibin Mice ENDPOINT one week; (3) failure	e to gain weight
Mice, Voles,	
Ground	
Squirrels, etc,	
Shrews, various	
species. Bats,	
various species.	
Small to	
Medium	
1702-34601A Whittaker, Joe Carnivores FREQUENCY see protocol for det	ails

1702-34601A	Whittaker Joe	Mice, Voles, Ground Squirrels, etc, Shrews, various species, Bats, various species, Small to Medium Carnivores	PHYSICAL	There will be no intermittent release. Once set traps will remain undisturbed. Any more frequent checks would impact other animals entering traps and lower capture rates. Our goal is to capture as many individuals and different species as possible. Extra time in the field would negatively impact this goal. (Live trapping - mice, voles, ground squirrels, etc.) Traps are routinely left overnight when trapping nocturnal or crepuscular animals. For the
1702-34001A		Carrivoles	RESTRAINT	We will only use cervical dislocation
				in the event of a severe injury and to bring about an immediate end to suffering. Attempts to provide any anesthesia would further prolong suffering.
1702-34601A	Whittaker, Joe	Mice, Voles, Ground Squirrels, etc, Shrews, various species, Bats, various species, Small to Medium Carnivores	EUTHANASIA METHOD	I have never had to do this procedure on any of the medium carnivores I have captured. If a medium-sized carnivore is in such distress to warrant euthanasia it will likely already be unconscious or debilitated and in such distress that attempts to sedate or use anesthesia would prolong suffering.
1702-346004	Starr Tim	Mice	EUTHANASIA DEATH/MORIB UND ENDPOINT	In cases where the tumor volume cannot be measured externally and mice do not display any of the specific tumor endpoint criteria, we will use moribundity as a terminal endpoint in the study. The nature of the cancers we work with (gastrointestinal tract, lung, and ovarian) do not generally manifest as measurable tumors and the other tumor endpoint criteria are not evident. For these animals we cannot easily determine morbidity or tumor mass as a percentage of body

				L bo oboloo of goporal aposthotio
				agents can be difficult and confusing
				since eveny agent has specific
				strengths and weaknesses. Avertin
				has been used as an anesthetic in
				mapy lobe for mapy years at the
				Inally labs for many years at the
				University of Minnesota and was
				routinely approved under IACOC
				protocols. Avertin has also been
				used at many other institutions,
				Including the University of
				vvisconsin and Jackson
				Laboratories, where it was the
				"anestnetic of choice" for mice
				undergoing snort surgical
				procedures. I have personally used
				Avertin under IACUC approved
				protocols in the past and it has
				worked well with no untoward or
				unexpected events.
				Avertin does not cause as much
				bradycardia including effects on
				loading conditions and ventricular
				function compared to ketamine
				(Hart, et al., Am J Physiol Heart Circ
				Physiol 2001) and Avertin has a
				lower mortality rate than Ketamine
				(Harrison, et al., Investigative
				Ophthalmology & Visual Science
			NON-	May 2008, and personal
			PHARMACAUT	experience).
			ICAL GRADE	Avertin not only acts as a general
1702-34600A	Starr, Tim	Mice	COMPOUNDS	anesthetic, it also provides good
				Dr. Tran and his staff are
				experienced in performing mouse
				decapitation proficiently.
				Rapid decapitation without sedation
				is performed with large sharp
				scissors only in juvenile rats (P0,
				P7, or P15 timepoints in our
				experimental design). All laboratory
				personnel who perform this
	L		EUTHANASIA	euthanasia method have
1702-34584A	Tran, Phu	Mice, Rat	METHOD	demonstrated technical skill with the

				In some few cases (Serial Injury) we
				will re-injure the muscle with
				cardiotoxin after 8-weeks, then 3-
				week, then 3 more weeks. This is to
				test if the cells are capable of
				regenerating under for stringent
				conditions after multiple injuries. As
				stated elsewhere, the the surgery is
			MULTIPLE	minor, involving a small incision with
1702-34580A	Perlingeiro, Rita	Mice	SURGERY	little noticeable pain, distress and
				Because the effect of analgesics on
				muscle regeneration is unknown,
				and because we want to examine
				the role of inflammatory cells in
				skeletal muscle
				damage/regeneration, we cannot
				use any agents that may interfere
				with inflammation. These mice can
				therefore not be treated with
				analgesics. The surgeries that
				required post-op analgesics are
				minimally invasive requiring only a
				small (~0.5cm) incision and needle
				injection. The veterinarian Dr.
				Hashway has commented that the
				surgery would cause very minor pain
				and therefore should not
				substantially negatively affect the
				welfare of the animals. If going
				forward, we find that we cannot
				perform these experiments due to
				excessive pain/distress we will then
				add an anaigesic. However, since
				opioids or local analgesics on
				the strong offect of addiction and
				dependence that the miss may
				ovporionoo, wo request a switch to
				Pain class C without the
1702-345804	Perlingeiro Rita	Mice	POLICY	administration of any analogsics
1702-34580A	Peningeiro, Rita	Mice	PULICY	administration of any analgesics.

				Our justification is that TBE is only
				used as anesthetic in non survival
				procedure in the context of
				physiological recording of muscle
				force using an organ bath, where our
				usual anesthetic ketamine/xylazine
				is not indicated due to its potential
				muscle relaxant effect as well as its
				sensory and motor uncoupling
				activity from the brain. The
				alternative of isoflurane pose special
				challenges with the scavenging of
				waste anesthetic gases and the
				requirement of an apparatus/system
				that is only located in our RAR
				facility that is too distant from our
				complex Organ Bath apparatus,
				neither of which can be moved.
				Additionally, TBE has been
				approved by our IACUC on our
				protocol.
				I BE is reconstituted in sterile
				conditions as 2,2,2 I ribromoethanol
				0.5 g in 1 mL of 2 methyl-2 butanol,
				mixed at 37 C and diluted with 40
				mL distilled sterile water. The final
			NON-	solution is then filtered through 0.22
			PHARMACAUT	micron filter and kept refrigerated
			ICAL GRADE	(4c) and protected from light. It can
1702-34580A	Perlingeiro, Rita	Mice	COMPOUNDS	be stored up to 2 weeks but it is
				We use either under-gravel filters
				where the sediment waste on the
				gravel is siphoned out monthly or
			SANITATION	above-tank charcoal filter systems
1702-34545A	Mand, Sandy	Fish (Other)	FREQUENCY	where the charcoal cartridges are

				NSAIDs or other anti-inflammatory
				drugs will not be administered to
				these animals at any point in the
				study as they will interfere with the
				immunological parameters that are
				central to this study. Recent articles
				that refer to the potential use of
				other drugs (e.g., Gabapentin,
				Memantin and Mexiletin) for
				analgesia were considered as
				alternate analgesic agents.
				However, these compounds have
				significant effects on neural/brain
				function and would interfere with our
				ability to study the oxidative stress
				response in the brain during viral
				encephalitis and hence will not be
				used. (Intracerebroventricular
			72 HOUR	stereotaxic injection of MCMV)
			POST-OP	009 0.0
	Lokensgard,		ANALGESIA	NSAIDs or other anti-inflammatory
1701-34539A	James	Mice	POLICY	drugs will not be administered to
			ENVIRONMEN	
	nina sa asar ana a		TAL	
1701-34527A	Kotz, Catherine	Mice	ENRICHMENT	see protocol for details
			SOCIAL	
1701-34527A	Kotz, Catherine	Mice	HOUSING	see protocol for details
				In the proposed animal studies, we
			EUTHANASIA	will determine genetic changes-
			DEATH/MORIB	caused tumor-related survival and
			UND	thus we need to euthanize the
1701-34523A	Deng, Yibin	Mice	ENDPOINT	mouse cohorts as possible as close

	1			
				we request an exception to allow for
				our use of AAV Viral Infusion and
				chronic icv peptide delivery in
				otherwise undisturbed animals. Use
				of two separate surgical procedures
				(AAV + minipump implantation) is
				beneficial, as it will reduce the
				amount surgery done at one time
				(reduces potential for tissue
				trauma), minimizes the need for
				anesthesia supplements during
				surgery, allow animals to fully
				recover between each class B
				surgery. It also maximizes the
				potential for animals to reach the
				study completion. □
				Maximal viral-mediated gene
				transfer occurs approximately 3-8
				weeks after infection, depending on
				the brain region and neuronal
				process (cell body versus terminals)
				being targeted. In vitro and in vivo
				studies begin at this time, but
				require up to 4 weeks of additional
				time for completion. This method
				will substantially reduce the overall
				number of animals required to
				complete experimental studies. This
				conservation of animal resources is
				particularly useful when studies
	Bartolomucci			occur in transgenic animals which
1701-345224	Alessandro	Mico		require considerable time and
1701-0-1022A	Alcooundio	MICC		The lab staffs are well trained and
1701-345164	Rischof John	Mice Rat	METHOD	proficient enough to perform the
1101-04010A		Wilde, Mat	METHOD	NSAIDs or other anti-inflammator/
				drugs will not be administered to
				these animals at any point in the
				study as they will interfere with the
				immunological parameters. The use
				of analgesics would interfere with
				some of the opdocints and immune
	Lokonsgard			assossments in this study and will
1701 245124	Lokensyaru,	Mico		assessments in this study and will
1701-34313A	James	WIICE		This is a model of hyperalgesia that
				is being used to investigate the role
				of CD38 in neuropathia pain. The
1610 244404		Miao		
1012-34440A	Guedes, Alonso	IVIICE		Use of analgesics will preclude
				Since animals wort have diet
				and due to the notestic land
4040 044404		N4:		and due to the potential long
1012-3444UA	Gueaes, Alonso	wice	I(FOOD/FLUID)	duration of the study, we would ask

				We request an exception for the use
				of non-pharmaceutical grade
				compounds for this procedure
				Avertin (tribrementhenel) is an
				Avertini (inbiomoethanoi) is an
				effective anestnetic agent for this
				procedure due to the fact that this
				procedure, is not a survival
				procedure and therefore there is no
				risk for ulcers or tissue necrosis at
				site of injection. Further Avertin is
				not harmful or a controlled
				substance and under proper
				preparation it is a potent anesthetic.
				Preparation of Avertin will be made
				following the university and IACUC
				auidelines:
				-Sterile fileter with 0.2 micron filter
				-Store and use under sterile
				conditions
				-Store in the dark bottle of foil
				covered container
				-Do not use if the solution becomes
				discolored or bas a precipitate
1612 344404	Guadas Alansa	Mico		Check pH before each use and use
1012-34440A	Gueues, Aloriso	MICE	COMPOUNDS	-Check pribelore each use and use
				In this procedure, animals undergo
				day 14 to acuse placental isohomia
				Then an approximation day 17 or 19 o
				Then on gestation day 17 of 16 a
				carolid aftery calleter is placed
				under isotiurane anestnesia for
				monitoring of blood pressure on
				gestation day 19 prior to
				exsanguination under anesthesia
				and necropsy. The carotid artery
				catheter is not placed at time of
				RUPP surgery on day 14 of
				gestation because maintaining
				patency of the catheters for
				prolonged periods of time is difficult
			MULTIPLE	in the rat and previous experiments
1612-34428A	Regal, Jean	Rat	SURGERY	by Gilbert have revealed that an
			SOCIAL	
1612-34428A	Regal, Jean	Rat	HOUSING	see protocol for details
				These experiements require a
				procedure to implant telemetry and
				a separate procedure for treatment
			MULTIPLE	groups, for baseline recordings two
1612-34416A	Osborn Jr. John	Mice	SURGERY	or more separate procedures are

				Sovere anomia is desired to mimic
				Severe allernia is desired to mimic
				the degree of anemia seen clinically
				in preterm neonates. This
				phiepotomy protocol has been
				validated and published (vvaliin DJ,
				I kac I, Stucker S, et al. Phiebotomy-
				induced anemia alters hippocampal
				neurochemistry in neonatal mice.
				Pediatric research. 2015;77(6):765-
				771.) and is used in our lab under
				IACUC protocol 1412-32111A in
				mice. As mice have tolerated this
				procedure, we expect the rats too as
				well, though they may be at
				increased risk of death given
				anemia + hyperoxia/ hypoxia
				chamber.
				Further, to justify the amendment to
				the protocol to increase volume
				phlebotomize/ 24 hour period, a
				study of iron deficiency anemia
				which fed dams an iron deficient diet
				and then after weaning, fed pups an
				iron deficient diet, found pups to
				have hematocrits of 18% at 6 weeks
				of life (Oh S, Shin P, Chung J.
				Effects of developmental iron
				deficiency and post-weaning iron
			BLOOD	repletion on the levels of iron
	Ingolfsland,		COLLECTION	transporter proteins in rats. Nutr Res
1612-34393A	Ellen	Rat	LIMIT	Pract. 2015; 9(6):613-618.) There
				tail clip: The tail tip will be immersed
				in ice cold isopropyl alcohol or for 10
				seconds (this will also serve to
				disinfect the tail). 3 mm of tail tip will
				be cleanly cut using clean gloves
				and a sterile sharp scalpel or razor
				blade.
				Bleeding to be controlled with direct
				pressure. Mouse to be observed in
1612-34391A	Lund, Iroy	Mice	TAIL BIOPSY	the cage after releasing to assure
				I his method of euthanasia will only
				pe used for narvesting tissue from
				neonatal mice. Neonates are
4640.040704	They are Oterstern	Det Mice	LEUTHANASIA	thus regist decentration with share
1012-34372A	r nayer, stanley	rtat, Milce		inus rapid decapitation with sharp
1612 3/265 4	Kofuii Paulo	Mico		see protocol for details
1012-34300A	Roruji, Paulo	MICE	DVIICOUTING	see protocorror details

				The food consumption of the mice
				should not change with this addition
				of 2W peptide and no weight loss is
				anticiptated. If a mouse is found to
				be in distress, an RAR vet or vet
				tech will be consulted and allowed to
				treat as needed or the mouse will be
				euthanized
				(2) ( $(2)$ (
				The food consumption of the mice
				should not change with this amino
				acid diet and no weight loss is
				anticiptated. If a mouse is found to
				be in distress, an RAR vet or vet
				tech will be consulted and allowed to
				treat as needed or the mouse will be
				euthanized. (amino acid diet)
				Bromodeoxyuridine is not acutely
				toxic to mice at the dose being
				given and no impact on the animals
				health is expected. If a mouse is
				found to be in distress, an RAR vet
				or vet tech will be consulted and
				allowed to treat as needed or the
				mouse will be euthanized.
				(Drinking water with
				bromodeoxyuridine)
			WEEKLY	
			WEIGHING	I he water consumption of the mice
1010 0 1000 1		N 41	EXCEPTION	should not change with the addition
1612-34360A	Jenkins, Marc	Mice	(FOOD/FLUID)	of OVA and no weight loss is
				Procedure is instant. Mice are
				dispatched immediately. Also, we
				do not want the empryos to be
				anesthetized upon harvest.
				We will only use decapitation for
				embryos, as described in the
		Mice, Pig	EUTHANASIA	procedures.
1611-34310A	Low, Walter	(Biomedical)	METHOD	
				In our model of intestinal pathology
				and autoimmunity, we use death as
				an endpoint, especially when we are
				testing any interventions to alleviate
				intestinal pathology. This is because
			EUTHANASIA	rescuing animals from death is a
			DEATH/MORIB	very high bar for efficacy. The
			UND	endpoints are death or recovery
1611-34309A	Vezys, Vaiva	Mice	ENDPOINT	from having malaise or being

				Avertin has been used in on-going
				and previous studies performed by
				our lab therefore in order to avoid
				introduction of a new variable and to
			NON-	allow for cross comparison with
				previous and current experiments
			ICAL GRADE	we request to continue the use of
1611-34309A	Vezvs Vaiva	Mice	COMPOUNDS	Avertin as an anesthetic vs other
	rozyc, ranta	inioo		We use either under-gravel filters
				where the sediment waste on the
				gravel is siphoned out monthly or
				above-tank charcoal filter systems
				where the charcoal cartridges are
			SANITATION	changed based on manufacturer's
1611 242004	Mand Sandy	Fich (Othor)		
1011-34300A	Mariu, Sariuy		FREQUENCT	We have a few variation of electric
				fich Ac montioned elsowhere in this
				nsh. As menuoned elsewhere in this
				protocol, when electric fish are
				of another fight and will change ite
				or another rish, one will change its
				signal in the jamming avoidance
				response." Since this is what we
				study in the class, it is important to
				allow the fish time alone before we
				begin our experiments.
				The ghost knives and elephant
				noses usually can be socially
				housed except just before and
				during the experimental period when
				we need their electrical signalling to
				settle into their innate pattern. So
				one request is to take these fish out
				of social bousing during the
				experimental period, returning them
				to social bousing afterward if they
				will settle in tegether (note that we
				will settle in together (note that we
				is picked on by all fich so it is power
				specially boused. We need to be oble
				to upp our boot judgement on sufficient
				individual fieb together.
				individual fish together.)
1611 242004	Mand Sandy	Fich (Other)		The third veriety of electric fich we
1011-34300A	ivianu, sanuy	Fish (Other)		The third variety of electric fish we
			DHARMACALIT	
	Whitley			
1610-342654	Chester	Mice		
1010-04200A	Chester	MICC		

				Based upon our previous studies.
				ulceration of xenograft tumors
				occurs in drug treated animals. If I
				kill the animals when skin ulceration
				occurs over tumors, I will achieve
				nothing in this project because
				immature termination of an effective
				treatment. I have observed that
				these early ulcerations on xenograft
				tumors formed crusts, fell off, and
				eventually healed. After that, only
				colored skin (the smooth surface
				with a little scar underneath) could
				be seen and xenograft tumors
				disappeared. Based upon this
				finding, we would like to make a
				modification on the original
				euthanization criteria regarding skin
				ulceration over xenograft tumors.
				This change allows me to continue
				the treatment till xenograft tumors
				disappear or treatment reaches its
				end point (1 month after injecting
				cancer cells or 10 weeks after
				starting the treatment). Early
				ulceration caused by drug treatment
				is not a sign to euthanize the
				animals when tumors remain small.
				Instead, it is a good sign to continue
				the protocol as defined in our
			TUMOR	original application when tumors
			ENDPOINT	remain small. It means that drug
1610-34251A	Lin, Jizhen	Mice	CRITERIA	treatment begins to kill some tumor
				A state of moribundity must be
				reached to determine if mice treated
			EUTHANASIA	with ultrasound experience tumor
			DEATH/MORIB	reduction and/or ablation, and
1010 0 1050 1				whether they live longer than their
1610-34250A	Low, Walter	Місе	ENDPOINT	untreated counterparts. Each
				Our animal studies are trying to
-------------	-------------	------	-------------	---
				determine whether and how the
				genetic alterations or
				chemoprevention/chemotherapy will
				provide any survival benefit for the
				mouse cohorts. In order to generate
				a credible mouse survival curve, we
				need to observe the mouse cohorts
				up to "natural death". However.
				these mice are valuable and needed
				for pathological and biochemical
				analyses. we cannot allow them to
				die spontaneously and risk organ
				deterioration. Rather, we will harvest
				mice when they are determined to
				be moribund-"close to death
				endpoint". One Criterion from UMN
				IACUC guideline will potentially
				affect our studies to observe
				survival benefits. Based on "UMN
				IACUC Euthanasia Guideline'', we
				have provided a very strict criteria to
				determine the moribund state, and
				these must include at least one of
				the first two criteria and at least one
				of the three remaining criteria: (1)
				progressive weight reductions up to
				10% of body weight measured on
				two separate occasions over a
			EUTHANASIA	period of one week; (2) sudden,
			DEATH/MORIB	unexplained weight loss of at least
			UND	10% of body weight over a period of
1610-34244A	Deng, Yibin	Mice	ENDPOINT	one week; (3) failure to gain weight

				Animals will receive an application
				of lidocaine gel (2%) in and around
				the skin incision to alleviate the
				irritation and pain associated with
				the surgery. From past experience
				animals recover from the surgery
				and return to normal behaviors
				(feeding, etc) within 12-24 h.
				Feeding and watering schedules will
				not be altered for the surgery or post-
				surgery. NSAIDs or other anti-
				inflammatory drugs will not be
				administered to these animals at
				any point in the study as they will
				interfere with the immunological
				parameters that are central to this
				study. In addition to monitoring
				animals post-anesthesia, daily
				monitoring for loose cannulas,
				wound infections, or mutilations will
				be performed up to 7 d post-surgery.
				Animals that lose their cannula
				before the treatment protocol is
				complete will be anesthetized and
				euthanized. (Canulation surgery)
				The animals will be monitored until
				they can independently maintain
				sternal recumbency or can stand
			72 HOUR	and move about before leaving the
			POST-OP	surgery room. Pain post-surgery is
	Cheeran,		ANALGESIA	expected due to injury to the
1610-34229A	Maxim	Mice	POLICY	scalp/skin incision. Animals will

				pased virus to transfect cells and the
				device. The trapefaction takes 2.4
				weeks to take full effect, and the
				by perdrive takes 1-2 weeks to reach
				its target with the best recordings
				occurring during the subsequent
				several weeks. This means that if
				we did both procedures in the same
				surgery, the optimal time for
				DREADD transfection and
				hyperdrive recording will be
				mismatched. Therefore, we will do
				two surgeries, one, first to transfect
				with DREADDs and the second to
				implant the hyperdrive. Rats will
				have at least 2 weeks between
				surgeries.
				This procedure will (by definition)
1610-342264	Redish David	Rat	SURGERY	he a second surgery This
1010-04220A		Γιαί	SOCIAL	
1610-34226A	Redish. David	Rat	HOUSING	see protocol for details

				The manufacturer of these osmotic pumps explains why removal is necessary (found here: http://www.alzet.com/products/guide _to_use/implantation_and_explantat ion html):
				Explanting ALZET Pumps
				Surgical removal of the ALZET pumps is accomplished in the anesthetized animal via a simple skin incision. If the pump has been in place longer than a couple of weeks, or the infusate is an irritant, it may be necessary to free the pump from surrounding connective tissue in order to remove it.
				The pump should be removed in the following circumstances:
				To verify delivery by measuring residual volume To verify stability & bioactivity of the test agent in solution No later than the recommended "explant by" date (see below) Note that an explanted pump cannot be reused.
1609-34180A	Geller, Melissa	Місе	SURGERY	

				Muce will be prepared for chronic
				awake cortical imaging in two steps
				In the first surgery, the skin over the
				and the first surgery, the skirt over the
				scalp will be removed and a metal
				nead bar will be permanently
				attached to the skull with
				cyanoacrylate glue and dental
				cement. The animal will then be
				allowed to recover and given
				adequate antibiotics Baytril (Bayer;
				5mg/kg) and pain medication
				Buprenex (2mg/kg) so that it fully
				recovers and is pain-free. The
				animal will then be acclimated to
				being fixed to a frame under a
				microscope for several days
				Following applimation, a second
				Following acclimation, a second
				surgery will be performed to create a
				tninned-skull cranial window for
				imaging the cortex.
				The reason to perform the surgery in
				two steps is that following the
				creation of the cranial window and
				receiver, from the second surgery
				which will take 2 to 3 days, we can
				which will take 2 to 3 days, we can
				Immediately begin our imaging
				sessions on the awake animal. The
				animal will have already been
				acclimated to being fixed under the
				microscope. However, this
			MULTIPLE	immediate imaging paradigm may
1609-34137A	Newman, Eric	Mice	SURGERY	not always be necessary and we will
				Because the primary tumors reach
				endpoint prior to establishment of
				robust metastasis, primary tumor
				resection will be necessary to
				assess metastatic growth and
				burden in these models. All
				procedures for tumor resection will
				include anesthesia and pain relief in
	Schwortfogor			the same manner as provided for
1600 244004	Sonwenneger,	Mino Mino		the initial ourgany Due to the
1609-34120A	naylee	wice, wice	SUKGEKY	the initial surgery. Due to the
				Decapitation is a standard approach
				for procurement of neural system
				tissue, sedatives may interfere with
				the cellular dynamics. Decapitation
				of guinea pigs will be provided by
		Mice, Guinea	EUTHANASIA	individuals with a demonstrated high
1608-34110A	Segura, Bradley	Pig	METHOD	degree of technical proficiency. We
	Yang, Yi-Mei		SOCIAL	· · · · · ·
1608-34101A	(Amy)	Mice	HOUSING	see protocol for details

				Mice are cogrificed by description
				wind a Personi Constant a starile
				using a Decapicone and a sterile
				snarp blade. The DecapiCone is a
				plastic conical rodent restraint
				device that is commonly used to
				restrain animals in order to reduce
				stress from handling, and to
	Yang, Yi-Mei		EUTHANASIA	minimize the chance of injury to
1608-34101A	(Amy)	Mice	METHOD	experimenters. This way, a rapid
				Our animal studies are trying to
				determine whether and how the
				genetic alterations or
				chemoprevention/chemotherapy will
				provide any survival benefit for the
				mouse cohorts. In order to generate
				a credible mouse survival curve, we
				need to observe the mouse cohorts
				up to "natural death". However
				these mice are valuable and needed
				for pathological and biochemical
				apply sos, we eappet allow them to
				dia opertaneously and risk organ
				die spontaneously and risk organ
				deterioration. Rather, we will harvest
				mice when they are determined to
				be moribund-"close to death
				endpoint". One Criterion from UMN
				IACUC guideline will potentially
				affect our studies to observe
				survival benefits. Based on "UMN
				IACUC Euthanasia Guideline'', we
				have provided a very strict criteria to
				determine the moribund state. and
				these must include at least one of
				the first two criteria and at least one
				of the three remaining criteria: (1)
				progressive weight reductions up to
				10% of body weight reactions up to
				two constants opposings over a
				two separate occasions over a
				perioa or one week; (2) sudden,
			DEATH/MORIB	unexplained weight loss of at least
			UND	10% of body weight over a period of
1608-34064A	Deng, Yibin	Mice	ENDPOINT	one week; (3) failure to gain weight

1607-33994A	Schleiss, Mark	Guinea Pig	HOUSING	see protocol for details
1608-34040A		(Xenopus), Gerbil, Goat, Guinea Pig, Hamster, Mice, Nonhuman Primate (Macaques), Pig (Biomedical), Rabbit, Rat, Sheep (Biomedical), Turkey, **ALL SPECIES DESIGNATION **, Cow (Biomedical), Chinchilla, Ferret, Cat	EUTHANASIA METHOD	Euthasol® 0.22 ml/kg IV (~86 mg/kg pentobarbital)
1608-34040A		Dog, Frog (Xenopus), Gerbil, Goat, Guinea Pig, Hamster, Mice, Nonhuman Primate (Macaques), Pig (Biomedical), Rabbit, Rat, Sheep (Biomedical), Turkey, **ALL SPECIES DESIGNATION **, Cow (Biomedical), Chinchilla, Ferret, Cat	SOCIAL HOUSING	see protocol for details

1607-33989A	Robinson, Jerid	Mice. Mice	Social Housing	The mice must be individually housed for the entire study while they have the telemetry transmitter implanted. The reason is because each transmi tter communicates to the receiver the mouse cage is in. This is to detect their individual arterial pressure and heart rate. The technology of the transmitters for now does not allow them to be housed in groups because the
1606-33864A	Cao, Ruifeng	Mice	EUTHANASIA	We need to look at protein kinase activation in the brain after acute stimulation, such as a light pulse. Due to the transient nature of the protein kinase activities, animals must be euthanized immediately and brain tissue must be harvested and processed within minutes after light exposure. The time it takes to sedate the animals are too long for
1605-33816A	Mand, Sandy	Fish (Other)	SANITATION	We use either under-gravel filters where the sediment waste on the gravel is siphoned out monthly or above-tank charcoal filter systems (especially for the marine tanks) where the charcoal cartridges are changed based on manufacturer's recommendations. We also have protein skimmers for the marine tanks. We also have an Z-Hab system in Biosci 115 which includes
1605 220164	Mand Sandy	Fish (Other)	SOCIAL	and protocol for dotaile
1000-330 IOA	wianu, Sandy	Nonhuman	PRIMARY	
1605-33678A		Primate (Macaques)	ENCLOSURE SIZE/SPACE	see protocol for details
1605-33678A		Nonhuman Primate (Macaques)	SANITATION FREQUENCY	see protocol for details
1605-33678A		Nonhuman Primate (Macaques)	MULTIPLE SURGERY	see protocol for details

# **IACUC RESEARCH SUBMISSIONS**

# APRIL 1, 2019 - SEPTEMBER 30, 2019

**TOTAL SUBMISSIONS: 596** 



New Protocols - 236

Changes in Protocol -360

# TOTAL SUBMISSIONS – 596 BY SUBMISSION TYPE APRIL 1 – SEPTEMBER 30, 2019



# TOTAL SUBMISSIONS – 596 BY SUBMISSION TYPE APRIL 1 – SEPTEMBER 30, 2019

NEW PROTOCOLS - 236

AMENDMENTS - 360



# TOTAL SUBMISSIONS – 596 BY SUBMISSION TYPE APRIL 1 – SEPTEMBER 30, 2019



# **REVIEW OUTCOMES- NEW STUDIES** APRIL 1 – SEPTEMBER 30, 2019

FCR AMENDMENTS - 14

DMR AMENDMENTS -346



# **REVIEW OUTCOMES - AMENDMENTS** APRIL 1 – SEPTEMBER 30, 2019



FCR SUBMISSIONS MEDIAN APPROVAL TIMES OCTOBER 1, 2018 – MARCH 31, 2019



# DMR SUBMISSIONS

MEDIAN APPROVAL TIMES OCTOBER 1, 2018 - MARCH 31, 2019



# SUBMISSION COMPARISON – TOTALS BY TYPE APRIL 2012 – SEPTEMBER 2019



SUBMISSION COMPARISON – TOTALS BY TYPE AND REVIEW PROCESS APRIL 2012 – SEPTEMBER 2019



# SUBMISSION COMPARISONS REVIEW OUTCOMES - NEW PROTOCOLS VIA FCR



# SUBMISSION COMPARISONS REVIEW OUTCOMES - NEW PROTOCOLS VIA DMR



# SUBMISSION COMPARISONS REVIEW OUTCOMES – AMENDMENTS VIA FCR



# SUBMISSION COMPARISONS REVIEW OUTCOMES - AMENDMENTS VIA DMR





TIME COMPARISON – FCR NEW PROTOCOLS APRIL 2014 – MARCH 2019





TIME COMPARISON – FCR AMENDMENTS APRIL 2014 – MARCH 2019





# TIME COMPARISON – DMR NEW PROTOCOLS APRIL 2014 – MARCH 2019



# TIME COMPARISON – DMR AMENDMENTS APRIL 2014 – MARCH 2019

### EXPIRED/SUSPENDED, EXTERNAL ANIMAL HOUSING, AND INCOMING ANIMAL TEMPORARY PROTOCOLS 4/1/2019 – 9/30/2019

#### Holding Protocol

Christine Sivula, 112600 1807-36197A 4-1-2019 - 9-30-2019

PI	Protocol ID	Species	Number of Animals	Expiration Date	New Protocol ID	Transfer Approval Date
Finger, Erik	1604-33640A	Mouse	22	5/12/2019	1905-37028A	5/31/2019
Asakura, Atsushi	1604-33660A	Mouse	132	5/23/2019	1903A36906	8/19/2019
Marahrens, York	1605-33812A	Mouse	115	9/12/2019	1809-36337A	9/17/2019 Amendment
*Lund, Troy	1605933717A	Mouse	2	6/16/2019	None	None

\*Dr. Troy Lund had one box of mice on protocol 1605A33717 (expired on 6/16/2019) that could not be moved to protocol 1809A36360 (where his other boxes of mice were transferred to on 6/16/2019) due to experiments that had been performed on these mice. That box of mice was outed on 9/18/2019.

### **External Animal Housing Protocol**

Angela Craig – transferred to Mark Suckow – transferred to Christine Sivula 1808-36233A 4-1-19 - 9-30`-19

PI	Company Name	Protocol ID	Specie	Number of Animals
*Sivula, Christine	Boston Sci	1808-36233	Dog	0

\* Formerly Mark Suckow.

#### **Incoming Animal Temporary Protocol**

Christine Sivula 1807-36151A 10-1-18 - 3-31-19

Incoming PI	Date of Arrival	Number of Animals Housed	Specie	New Protocol #
Junge, Harald	8/12, 8/13, 8/15	425	Mouse	1907A37213

### Andrew Grande:

### 6/20/19:

• It was noted during review of your surgical records for protocol 1807-36097A, that mice underwent a second stereotaxic surgery for intracranial virus injections 7 days after they had undergone the Controlled Cortical Impact surgery. Although the protocol states that mice will receive intracranial virus injections, it does not clearly state that this procedure is performed as a surgical intervention as there is no surgical procedure described in the protocol for this secondary surgery and the Controlled Cortical Impact Surgery states that there are no multiple surgeries. Please include this information in the protocol by first updating the Controlled Cortical Impact surgical procedure to indicate that these animals will undergo multiple survival surgeries (there is a checkbox under the "surgery" heading for this purpose) and also adding the second AAV administration surgical procedure to the protocol.

### 12/18/18 and 12/20/18:

- It was noted during the inspection that Andrew Crane is performing surgeries under protocol 1601-33310A; however, he is not listed on this protocol. Please add Andrew to this protocol. Also, please make sure he is listed as a surgeon on this study and provide his training and relative experience for review and approval by the IACUC.
- It was noted during the inspection that a femoral vein cutdown procedure is performed 48 hours after the controlled cortical impact surgery to facilitate the IV injection of cells, as approved; however, no analgesic was administered with this surgical procedure. The protocol specifies that Buprenorphine-SR will be administered at least 3 hours prior to the surgery if used alone; or, it will be administered upon sedation of the animal just prior to surgery if used in conjunction with standard formulation buprenorphine. Please confirm that analgesics will be administered for this procedure as stated in the protocol. Additionally, you may want to discuss with your area veterinarian, Dr. Misha Dunbar, this analgesic regimen to ensure proper dosing. (Please note that this is a repeat finding also noted on June 21, 2018 inspection report.)

### 6/20/18:

• It was noted during the inspection that a femoral vein cutdown procedure is performed 48 hours after the controlled cortical impact surgery to facilitate the IV injection of cells, as approved; however, no analgesic was administered with this surgical procedure. The

protocol specifies that Buprenorphine-SR will be administered at least 3 hours prior to the surgery if used alone; or, it will be administered upon sedation of the animal just prior to surgery if used in conjunction with standard formulation buprenorphine. Please confirm that analgesics will be administered for this procedure as stated in the protocol.

### 12/29/17:

- It was noted that two staff members working on your study are not listed as personnel on your protocol (Hui Xie and Wei Chen Lu). Please have them discontinue working on your study until such time they are added to the protocol and have completed their ROHP requirements, if applicable.
- In review of surgical records for this study, it was noted that on 10/16/17, rats (12) underwent traumatic brain injury ranging from mild to severe. The severe brain injury model is not outlined in your protocol. Please discontinue this model until you submit an amendment to your protocol. Additionally, staff indicated that this initial set of surgeries was to help set up TBI parameters as they are not well established for your study, hence no cell therapy was done on these animals only a range of injury models. Please also add a group of animals to be used for initial system set up if needed.

### 5/31/16:

Protocol 1505-32629A states that "animals may maximally undergo a weekly MRI procedure to evaluate the progression of aneurysm formation and then after treatment to assess treatment effect" but according to animal records, dogs GAM1 and GAM2 received two MRI's in the given week (GAM1 received an MRI on 1/13/16 and 1/14/16 and GAM2 received an MRI on 2/10/16 and 2/11/16 as well as on 2/22/16 and 2/23/16.. Please update your protocol to reflect that more than one study MRI may be conducted in a given week. Additionally, a provision in the protocol should be added that if adverse clinical signs are seen after aneurysm induction that additional MRIs may be warranted.

### 11/16/15:

 It was noted that some mice under protocol 1501-32253A were euthanized using Isoflurane anesthesia with subsequent perfusion. According to your study, only CO2, cervical dislocation or ketamine overdose and decapitation are approved methods. Please discontinue the Isoflurane anesthesia and perfusion euthanasia procedure until an amendment is submitted to your eProtocol submission and you are approved in writing by the IACUC.

### John Bischof:

### 6/24/19:

Mice undergoing thermal therapy procedures did not receive post procedure analgesics as indicated in the protocol. Appendix F for thermal therapy attached to your protocol states that buprenorphine will be given on the day of surgery, twice on the next day and once more the following day. The Health and Monitoring section of your protocol states that buprenorphine will be given at the time of surgery and ibuprofen in the drinking water for 7 days following thermal therapies. Please consult with your RAR veterinarian to assess appropriate post-procedure analgesia (or none) for the various thermal therapies on the protocol and submit an amendment with any changes to analgesia. This is a repeat finding also noted on the report sent 6/22/2018. In response to this finding last year it was stated that an amendment would be submitted and approved prior to resuming thermal therapy procedures.

### 6/22/18:

- It was noted that xylazine that expired in May 2018 was used for ketamine/xylazine anesthesia of mice for thermal therapy in June 2018. Expired anesthetics, analgesics and euthanasia agents may not be used in research animals. Please replace the xylazine prior to your next anesthetic procedure.
- Mice undergoing thermal therapy procedures did not receive post-surgical analgesics as indicated in the protocol. The appendix F for thermal therapy attached to your protocol states that buprenorphine will be given on the day of surgery, twice on the next day and once more the following day. The Health and Monitoring section of your protocol states that buprenorphine will be given at the time of surgery and ibuprofen in the drinking water for 7 days following thermal therapies. Please consult with your RAR veterinarian to assess post-procedure analgesia for the various thermal therapies on the protocol and submit an amendment with any changes to analgesia.

### **5/4/18** Self-Report:

This is in reference to pending protocol #1804-35844A submitted by PI Dr John Bischof. 8 We wanted to let the committee know that we had started housing two larval zebrafish prior to the approval of this protocol. The aim of our project is to develop a long term cryopreservation protocol for early stage zebrafish embryos. Most of the cryopreserved and rewarmed embryos don't make it past day 3 (vertebrate status). However, in past month we were able to get two embryos to develop to Day 5. Since this is a rare occurrence (and big breakthrough) for us, we immediately wanted to house these fish in the zebrafish core to see if their long-term development would be affected by our treatments. We were thinking that Zebrafish Core's fish room protocol would cover it. But after discussion with folks from IACUC and Dr Mark Masino, we were advised to submit a new protocol. Following his suggestion, we submitted an IACUC protocol last week and sincerely hope that we can avoid any breach of animal welfare policies in the future. We currently have one adult fish and one larval fish in separate tanks, which are considered to be IACUC approved housing. Please let us know if we can provide any other information regarding our protocols since we would like to completely transparent in this aspect.

### 6/16/17:

 It was noted during the inspection that tumor cell injections are sometimes performed by subcutaneous injection without the use of anesthesia; however, the approved protocol states that tumor cells will be injected under ketamine/xylazine anesthesia. Please reinstate the use of ketamine/xylazine injectable anesthesia for this procedure. If you would like to perform this procedure without anesthesia, please submit an amendment through eProtocol requesting this change.

### 1/12/17:

 It was noted that expired ketamine (exp. 6/2016) and expired xylazine (exp. 11/2016) were used to anesthetize mice for non-invasive procedures through December 2016. Expired anesthetics, analgesics and euthanasia agents may not be used in research animals. Please replace the ketamine and xylazine prior to your next anesthetic procedure.

### Clark Chen:

#### 7/11/19 and 7/16/19:

• It was noted during the inspection that mice under protocol 1802-35597A are housed in for 72 hours post-surgery; however, this protocol is not currently approved for housing in this space. In review of this protocol, it is stated that animals will be housed in this space overnight once surgery is complete and returned to the animal facility in the morning. If you would like to continue housing animals in this space for longer than 24 hours, please submit an amendment to the protocol to include this IMHA for the for housing mice under this study. Until such time that an amendment is approved, animals must not be housed in this space longer than 24 hours.

#### 1/18/19:

• In review of your surgical records, 5 mice on protocol 1707-34937A and 58 mice on protocol 1812-36595A underwent intracranial injection surgery from 10/26/18 to 1/16/19 and did not receive adequate post-operative analgesia. Although bupivacaine was given as indicated in the protocols, the first injectable dose of buprenorphine was not given until 24 hours after surgery. Analgesics must be given in accordance with the protocol and to cover the full 72 hour post-operative period. Please confirm that you will give buprenorphine on the day of surgery and for 72 hours following.

• Although diluted controlled substances are stored in a locked safe, this safe is not yet attached to the wall and therefore does not fully secure these drugs. Please move all controlled substances to the existing wall safe until the second safe has been secured by facilities management.