Rodent Survival Surgery SOP

1. Purpose
This Standard Operating Procedure (SOP) describes methods to be followed for rodent survival surgery. This SOP follows the UBC Policy and CCAC guidelines for the survival surgery of rodents.

2. References:
Associated Guidelines and SOP’s can be found on the ACC SOP/Guidelines website: http://www.ors.ubc.ca/contents/animal-care-sops-guidelines
TECH 10 - IP injection in mice and rats
TECH 11 – Subcutaneous injection in mice and rats
SOP ACC-2012-01 Rodent Anesthesia
Rodent Anesthesia and Analgesia (Formulary)

Surgical Class and Analgesic Guidelines
NSAID Analgesia for Rodents: Ketoprofen (Anafen)
NSAID Analgesia for Rodents: Meloxicam
Opioid Analgesia for Rodents: Buprenorphine

Local Analgesia/Anesthesia for Rodents: Lidocaine, Bupivacaine, Ropivacaine
Policy # 16 Survival Surgery of Rodents (http://www.ors.ubc.ca/contents/acc-policies)
Policy # 17 Monitoring of Animals Used for Research, Teaching and Testing (http://www.ors.ubc.ca/contents/acc-policies)

Procedure Classifications for Mice and Rats

<table>
<thead>
<tr>
<th>Class</th>
<th>1</th>
<th>2</th>
<th>3</th>
<th>4</th>
</tr>
</thead>
<tbody>
<tr>
<td>Pain Level</td>
<td>Mild Pain</td>
<td>Moderate Pain</td>
<td>Moderate/Severe Pain</td>
<td>Severe Pain</td>
</tr>
<tr>
<td>Examples</td>
<td>Subcutaneous implant with trocar</td>
<td>Craniotomy with or without implant</td>
<td>Laparotomy with major organ manipulation or removal</td>
<td>Peritonitis/pancreatitis</td>
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<td></td>
<td>Ocular procedures</td>
<td>Simple laparotomy</td>
<td>Organ transplant</td>
<td>Thoracotomy</td>
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<td></td>
<td>Tracheal injections</td>
<td>Embryo transfer</td>
<td>Spinal surgery</td>
<td>Hind limb ischemia</td>
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<td></td>
<td>Skin biopsy/wound</td>
<td>Ovariectomy</td>
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<td>Spinal or nerve injury</td>
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<td></td>
<td>Vessel cut down or cannulation</td>
<td>Castration</td>
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<td>Cecal ligation and puncture</td>
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<td></td>
<td>Intramuscular injection</td>
<td>Intra-peritoneal osmotic pump</td>
<td></td>
<td>Orthopedics/Fractures</td>
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<tr>
<td></td>
<td>Subcutaneous osmotic pump</td>
<td>Dental extraction</td>
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<td>Bone cancer</td>
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Minimum Pre-operative Analgesic Requirements for Mice and Rats

<table>
<thead>
<tr>
<th>Class</th>
<th>1</th>
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<td>Moderate Pain</td>
<td>Moderate/Severe Pain</td>
<td>Severe Pain</td>
</tr>
<tr>
<td>Pre-operative Analgesics</td>
<td>• Local anesthetic at site of surgery and • Non-steroidal anti-inflammatory OR • Buprenorphine</td>
<td>• Local anesthetic at site of surgery and • Non-steroidal anti-inflammatory OR • Buprenorphine</td>
<td>• Local anesthetic at site of surgery and • Non-steroidal anti-inflammatory OR • Buprenorphine</td>
<td>• Local anesthetic at site of surgery and • Non-steroidal anti-inflammatory AND • Buprenorphine</td>
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</table>

Minimum Time Period and Frequency for Post-Procedure Monitoring for Signs of Pain

<table>
<thead>
<tr>
<th>Frequency (Times/Day)</th>
<th>1-2</th>
<th>2</th>
<th>2</th>
<th>3</th>
</tr>
</thead>
<tbody>
<tr>
<td>Duration (# of Days including Day 0)</td>
<td>2</td>
<td>3</td>
<td>4</td>
<td>4</td>
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</tbody>
</table>

Minimum Post-Procedure Analgesia – based on pain assessment and duration of analgesia based on drug. Treatment may need to be extended

| Analgesic Drugs and Frequency of Administration | Day 0: NSAID once a day or Buprenorphine twice a day | Day 0 and Day 1: NSAID once a day or Buprenorphine twice to three times a day | Day 0, Day 1 and Day 2: NSAID once a day AND Buprenorphine twice to three times a day | Day 0, Day 1 and Day 2: NSAID once a day AND Buprenorphine three times a day |

Recommended suture material and sizes for closing surgical incisions

<table>
<thead>
<tr>
<th>Type</th>
<th>Size</th>
<th>Closure Pattern</th>
</tr>
</thead>
<tbody>
<tr>
<td>Abdominal wall/muscle</td>
<td>Absorbable (Vicryl, PDS, Monocryl)</td>
<td>Rats (4-0 to 6-0) Mice (5-0 to 6-0)</td>
</tr>
<tr>
<td>Subcutaneous tissue</td>
<td>Absorbable (Vicryl, PDS, Monocryl)</td>
<td>Rats (4-0 to 6-0) Mice (5-0 to 6-0)</td>
</tr>
<tr>
<td>Skin</td>
<td>Non-absorbable, monofilament (Prolene, Nylon)</td>
<td>Rats (4-0 to 6-0) Mice (5-0 to 6-0)</td>
</tr>
</tbody>
</table>

*Absorbable suture and a simple continuous pattern can be used for skin suturing if a subcuticular (buried) suture pattern is used or if a monofilament absorbable suture such as Monocryl is used.

**A useful suture needle is a swaged on RB-1 taper needle

3. Responsibility
3.1. Principal Investigators (PIs) and their research staff, animal care staff and veterinary care staff.
3.2. All animal users performing surgery on rodents must have successfully completed the UBC Animal Care Centre (or equivalent) Rodent Biology and Husbandry, Rodent Anesthesia and Rodent Surgery courses.

4. Introduction
4.1. Aseptic technique is a set of specific practices and procedures performed under carefully controlled conditions with the goal of minimizing contamination by potential pathogens.
4.2. In order to comply with Policy # 16 Survival Surgery of Rodents, steps must be followed to prepare the surgical area, animal, surgeon, and surgical instruments and/or implants.
4.3. Though rodents may survive a surgical procedure that is not done aseptically, major physiological changes will occur as the animal’s immune system fights infection and these can significantly impact animal welfare and research results.
4.4. Though not required, aseptic techniques for non-recovery surgical procedures (especially those that are longer than 1 hour) are highly recommended in order to prevent physiological changes that confound research results.

4.5. The goals of a successful surgery are: maintenance of asepsis, minimizing surgery time, minimizing tissue trauma by gentle tissue handling, and maintenance of normal rodent physiological parameters for the given surgical model.

4.6. Never leave an anesthetized animal unattended. Monitor animal until it has recovered from anesthesia enough to be moving around the cage normally.

4.7. Monitoring of and record keeping for post surgical animals should occur at least twice a day for the initial post-op recovery period, more often if animals are showing signs of pain or clinical signs of illness, and must continue daily at least until the surgical incision is healed, sutures are removed and the animal has recovered to the point that is normal for the given surgical model (see Surgical Class and Analgesia Guidelines).

4.8. Suture material, especially braided suture material such as Vicryl, should not be reused from animal to animal.

5. Materials

5.1. Material or equipment to provide or conserve body heat (e.g. light cloth or gauze as "blanket", heating disc or pad, warm-water circulating pad, infrared heater, electrical heating pad set on "low").

5.2. Corneal protectant (i.e.: lacrilube, isoptotears, etc)

5.3. Accurate animal weigh scale

5.4. Calculator

5.5. 70% Isopropyl alcohol

5.6. Disinfectant for surfaces

5.7. Paper towels

5.8. Warmed SQ fluids (i.e.: sterile 0.9% saline or sterile lactated ringer’s solution) if anesthesia/surgery is expected to last more than 10 minutes.

5.9. Appropriately sized needles and syringes for size of animals (See TECH 11 SQ injection in mice and rats and TECH 10- IP injection in mice and rats).

5.10. Warmed recovery cage (either no bedding or paper towel placed over bedding to prevent recovering animals from inhaling bedding particles)

5.11. Easily accessible food and water or gel water replacement (e.g.: Transgel, Napanectar, Hydrogel, etc).

5.12. Appropriate analgesics (see Rodent Anesthesia and Analgesia, Surgical Class and Analgesia Guideline and Policy # 16 Survival Surgery of Rodents) and recommendations given above.

5.13. Appropriate anesthetics (see SOP ACC-2012-01 – Rodent Anesthesia)

5.14. Cage flags, surgical log and/or monitoring sheets (see Policy # 17 Monitoring of Animals Used for Research, Teaching and Testing).

5.15. Appropriately sized rectal thermometer if anesthesia is expected to last more than 15 minutes (typically, if the probe diameter is the smaller than the diameter of the animal’s normal fecal pellet diameter, it can be safely used)

5.16. Additional patient monitoring equipment if available – pulse oximeter, capnograph (CO2 monitor)

5.17. Electric fur clipper (with size 40-50 blade)

5.18. Gauze

5.19. Cotton tipped applicators

5.20. Chlorhexidine (Hibitane) 2% surgical scrub or equivalent

5.21. Water resistant surgical pack wraps (Cloth wrappers, disposable Steri-wraps or Steri-pouches)

5.22. Chemical sterilization indicators

5.23. Autoclave tape

5.24. Sterilized (via steam autoclave) surgical instruments appropriate for planned surgery

5.25. Sterile gauze

5.26. Sterile cotton tipped applicators

5.27. Sterile drape with appropriately sized fenestration (hole) for surgical incision

5.28. Hot bead sterilizer (if sterile tip surgery is planned).
5.29. Bouffants (hair covers)
5.30. Surgical masks or N95 masks
5.31. Non-sterile exam gloves
5.32. Clean lab coat, scrub top or gown
5.33. Sterile towels or sterile paper towels
5.34. Sterile surgical gloves of the appropriate size for surgeon (if full sterile technique is being used)
5.35. Sterile suture material of appropriate size and type for surgery (see recommendations above).
5.36. Animal surgical log listing type of surgery, anesthetics and analgesics given
5.37. Cage flag cards indicating type of surgery, anesthetics, analgesics given
5.38. Monitoring sheets with duration and frequency of monitoring for parameters listed on approved animal care protocol

6. Preparation of surgical instruments and sterile supplies (typically done a day or two before planned surgery)
6.1. Clean all surgical instruments needed for surgery and wrap in chosen surgical pack for autoclaving. Include a chemical sterilization indicator in each pack. Place autoclave tape on outer surface of each pack.
6.2. Label each pack with date of autoclaving (and contents if desired).
6.3. In packs, include and required additional sterile supplies such as drapes, gauze, cotton-tipped applicators, sterile tip fields, bowls, etc.
6.4. Prepare packs for autoclave and set autoclave parameters to the necessary time, pressure and temperature as required to achieve full sterilization of the packs
6.4.1. A biological indicator should be placed in the center of the largest pack in the center of the load once a month to ensure the autoclave parameters chosen are achieving sterility.
6.5. Remove dried packs from autoclave, check that autoclave tape indicates packs have been processed, and place in an area where the packs will not get wet or dirty.

7. Preparation of surgical and prep areas (performed right before planned surgery)
7.1. Choose location for surgery area – it should be away from busy areas (such as doorways), easily cleaned and disinfected, and not directly under an air supply duct.
7.2. Remove all unnecessary equipment and clutter from area and clean surfaces to remove dust, fur, bedding, etc.
7.3. Disinfect surfaces in surgical and prep areas with a disinfectant (i.e. Virkon, Percept, etc) and ensure proper contact time (refer to manufacturer's recommendations).
7.4. The surgical and prep areas should be cleaned and disinfected before and after each surgical procedure to remove blood, urine, feces, hair, etc.
7.5. Bring required support and monitoring equipment to surgical and prep areas (i.e. heating source, syringes, needles, clippers, etc).
7.6. Bring sterile surgical packs and supplies to surgery area but do not open yet.
7.7. Turn on hot bead sterilizer if using sterile tip technique

8. Preparation of Animal
8.1. Put on non-sterile exam gloves, surgical mask, hair bouffant and a clean lab coat or scrub top
8.2. Calculate and draw up required anesthetics, analgesics and fluids. Record all drugs and volumes on Surgical log.
8.3. Anesthetize the animal according to SOP ACC-2012-01– Rodent Anesthesia in the designated prep area.
8.4. Once immobile, place animal on heat support, apply corneal protectant in both eyes, and administer required analgesics (see above references) and SQ fluids.
8.5. Remove hair from planned surgical site with fur clippers being careful not to cut the skin with the clippers. Shave at least 1 cm perimeter around planned surgical site for mice and 2cm perimeter for rats.
8.6. Remove all loose hair (using gauze or a small handheld vacuum)
8.7. Clean the skin at the planned surgical site with 2% chlorhexidine soap (or equivalent) using gauze (rats) or cotton-tipped applicators (mice). Start cleaning directly over where the planned incision will be and extend outwards without touching the center again. Take care not to wet or soak the animal as this will lead to heat loss.
8.8. Remove soap with 70% isopropanol on a gauze or cotton tipped applicator in the same pattern as the soap was applied (centre to periphery). DO NOT spray animal with the alcohol or get it near the eyes.
8.9. Discard used gauze and cotton tipped applicators after each application of soap or alcohol.
8.10. Administer local anesthetic (see Local Analgesia/Anesthesia for Rodents) at the planned surgical incision site.
8.11. Repeat the soap application and alcohol removal procedure two more times.
8.12. Ensure all loose hair is removed from the animal and bring animal to the prepared surgical area and continue supportive care (heat, oxygen, inhalational anesthetic (if using).
8.13. Lubricate and place rectal thermometer and any other monitoring equipment (pulse oximeter, capnograph, etc).
8.14. Ensure animal is positioned correctly for the surgery and check breathing, color of extremities and surgical level of anesthesia (rear toe pinch withdrawal).
8.15. Adjust anesthetic level if required (adjust isoflurane % or inject an additional small dose of injectable anesthetic being careful not to exceed therapeutic index).
8.16. Change gloves to a new pair of exam gloves.

9. Initial preparation of Sterile Field
9.1. Carefully open the sterile instruments close to the anesthetized patient (within easy reach) without touching any inner surface of the pack.
9.2. Aseptically add all required sterile supplies to the sterile field by opening the outer package and letting them drop a short distance onto the sterile field (i.e.: scalpel blades, suture material, gauze, cotton tipped applicators, syringes, bowl to hold sterile saline or LRS, etc).
9.3. Draw up required sterile saline or LRS in a syringe and add fluids to sterile bowl (this is used to keep exposed tissues moist during surgery).
9.3.1. Ensure that they do not touch any non-sterile surfaces and the person does not touch any sterile surfaces.
9.4. Open outer package of sterile surgical gloves without contaminating inner surface.
9.5. Check animal’s breathing, temperature, color and anesthetic depth – make any required adjustments to support animal and maintain a surgical plane of anesthesia.

10. Preparation of the surgeon
10.1. Remove exam gloves and wash hands and arms well (up to elbows if exposed) with disinfectant soap (i.e.: Chlorhexidine).
10.1.1. Keep hands raised above elbows to prevent “dirty” water from running down arms and contaminating hands.
10.1.2. Pay particular attention to underneath fingernails, between fingers, palms and wrist areas.
10.1.3. A good hand wash should take about 90 seconds to complete.
10.1.4. Dry hands with a sterile towel or paper towels.
10.2. If wearing a sterile surgical gown, put this on.
10.3. Put on sterile surgical gloves without touching outer surface if using full sterile technique, otherwise put on a clean pair of exam gloves.
10.3.1. Do not touch any non-sterile surface with sterile gloves.

11. Continued preparation of sterile field
11.1. If performing full sterile technique, with sterile gloves, arrange all the sterile instruments on the sterile field. Use needle drivers to safely place scalpel blade onto scalpel handle (if using). If performing sterile tip technique, arrange instruments and sterile supplies without contaminating surfaces that must remain sterile (using a sterile instrument to arrange the supplies is helpful).
11.1.1. If sterile tip technique is planned for additional surgeries using the same instruments, arrange the sterile field so that the “sterile tip” zone is well demarcated from the “contaminated handle” zone on the sterile field.
11.1.2. Ensure the sterile gauze, cotton tipped applicators, suture, etc is in the “sterile tip” zone of the sterile field.
11.2. Check animal’s surgical plane of anesthesia (breathing rate, color, toe pinch withdrawal reflex (can use sterile forceps or gauze to protect sterile fingers – do not place contaminated objects back into sterile field)). Adjust anesthetic depth if required.

12. Surgery
12.1. Drape animal with sterile drape with fenestration (hole) over the planned surgical incision
12.1.1. Do not contaminate surfaces that should remain sterile.
12.2. Perform required surgical procedure
12.2.1. Keep blood loss to a minimum by gentle tissue handling and hemostasis
12.2.2. Keep all exposed tissues moist with sterile saline or LRS
12.2.3. Prevent contamination of surgical site and sterile field
12.2.4. Monitor surgical plane of anesthesia as describe above approximately every 5 minutes and adjust as required
12.2.5. Monitor and maintain physiological status (heart rate, respiratory rate and pattern, color of extremities and temperature and provide necessary supportive care as described in SOP ACC-2012-01 – Rodent Anesthesia throughout surgical procedure and during recovery from anesthetic
12.3. Once surgical procedure is completed, suture muscle layers and skin separately using appropriate suture material and suture pattern (see page 2 References).
12.4. Gently remove any blood from the surgery site or surrounding fur.
12.5. Ensure corneas are lubricated – apply additional lubrication if they appear dry.
12.6. Provide additional SQ fluids if surgery was longer than expected, urine production is high (such as when using xylazine or dexmedetomidine) or if more blood loss than expected.

13. Recovery of animal
13.1. Turn off gas anesthesia (if using) and continue to provide oxygen and heat support until animal begins to move.
13.2. Move animal to warmed recovery cage and continue to provide heat support.
13.2.1. Do not recover directly on bedding as the bedding can enter the eyes or mouth – remove bedding entirely or place paper towels on top of bedding to protect animal
13.2.2. Monitor closely until animal begins to move around on its own and begins to groom face.
13.3. Place food and supplemental water source (i.e. Transgel, hydrogel) on cage bottom so that animal can easily access it.
13.4. Provide “pre-made” nest for post-surgical mice so they can maintain their own body temperature.
13.5. Complete surgical log with description of surgical procedure, any complications or expected outcomes and all analgesics, anaesthetics, and drugs given.
13.6. Place cage flag card on cage to indicate what procedure animal had performed and all drugs administered.
13.7. Begin appropriate monitoring sheet and record when sutures are planned to be removed.
13.8. Return animal to housing room when able to move about cage normally and maintain its own body temperature
13.9. Monitor according to approved animal care protocol until animal healed and doing well and complete required monitoring sheets. Pay particular attention to surgical incision, sutures, weight and signs of pain.
13.9.1. Record all additional analgesics and supportive care on daily monitoring sheet.
13.10. External skin sutures should be removed once incision healed – typically 7-14 days.
13.11. If animal removes sutures and requires resuturing or if incision is infected, underlying tissues are exposed or the incision fails to heal normally, contact UBC Clinical Veterinarians for consultation.
13.12. Re-suturing an opened incision should only occur with Veterinary approval and only if incision is clean, shows no discharge and within 24 hours of initial suturing.

14. Additional surgeries using sterile tip technique
14.1. Blood should be cleaned off the instruments with water and the tips placed into a hot bead sterilizer for 15 seconds ((typically about ½ the length of the instrument can be inserted into the hot bead sterilizer)
14.1.1. Once removed from the hot bead sterilizer, the sterile “tips” that will be in contact with the tissues must be placed into the “sterile tip” area of the field while the contaminated handles must be in the “contaminated handle” area of the field.
14.1.2. Care must be taken to place and re-place the instruments properly so as to not contaminate the sterile area of the field.
14.1.3. The instruments will be very hot once removed from the hot bead sterilizer so should be
given at least 20 minutes to cool to room temperature.

14.1.3.1. It is helpful to let the instruments cool while prepping the next animal for surgery.

14.2. Anesthetize and prep the animal as previously described.

14.3. Add additional sterile supplies (suture, drapes, etc.) to the sterile area of the field.

14.4. Wash hands and arms as previously described but don non-sterile exam gloves

14.5. Drape animal while handling only the underside of the drape so the upper surface remains
sterile

14.6. Ensure non-sterile hands and contaminated handles of the instruments do not touch any
sterile surface including the top surface of the drape, suture material, etc.

14.7. Instruments can be used for up to 5 sterile tip surgeries before needing to be re-autoclaved

14.8. Recover the animal and complete surgical log and monitoring sheets as described above.