

# Guidelines for Rodent Survival Surgery

## Scope

These guidelines apply to all surgical procedures performed on rodents at the NIH in which the animals are expected to recover from anesthesia. Prior to performing any survival surgery techniques on rodents, an approved Animal Study Proposal (ASP) must be in place with descriptions of the surgical procedures to be performed and the anesthesia/analgesia<sup>1,2</sup> that will be used pre-emptively, during and post-procedure. All personnel must also be appropriately trained. Specific procedures to accomplish these guidelines can be obtained from your veterinarian.<sup>3,4</sup>

## Definitions

- **Aseptic Technique:** is used to reduce microbial contamination to the lowest possible practical level.
- **Major Surgery:** Major survival surgery penetrates and exposes a body cavity, produces substantial impairment of physical or physiologic functions, or involves extensive tissue dissection or transection (e.g., laparotomy, thoracotomy, joint replacement, craniotomy, sciatic nerve cuff, and limb amputation).
- **Minor Surgery:** Minor survival surgery does not expose a body cavity and causes little or no physical impairment (e.g., wound suturing, peripheral vessel cannulation, percutaneous biopsy, and most procedures routinely done on an “outpatient” basis in veterinary clinical practice). Animals recovering from these minor procedures typically do not show significant signs of post-operative pain, have minimal complications, and return to normal function in a relatively short time.
- **Sterilization:** The process whereby all microorganisms including spores are eliminated or destroyed. The criterion of sterilization is the failure of organisms to grow if a growth supporting medium is supplied.
- **Disinfection:** The chemical or physical process that involves the destruction of pathogenic organisms. All disinfectants are effective against vegetative forms of organisms, but not necessarily spores.

## General

- Rodents do not vomit, so it is not necessary to fast them prior to surgery.<sup>5,6,7 6</sup>
- Appropriate pre-operative and post-operative care of animals in accordance with established veterinary medical and nursing practices are required.
- A designated animal procedure space for rodent surgeries is required; for example, a location within a procedure room or laboratory space free from clutter and easily disinfected prior to the surgical procedure. During the surgery period, the area should be dedicated to rodent surgery such that cleanliness is ensured, and contamination is minimized at the time of use.
- A “tips-only” technique restricts you to using only the sterile working ends of the surgical instruments to manipulate the surgical field. The gloved hand must never touch the working end of the instruments, the suture, suture needle, or any part of the surgical field.<sup>8</sup>

All survival surgery will be performed by using aseptic procedures, including masks, sterile gloves, sterile instruments, and aseptic techniques.

The *Guide* states that it is important for research personnel to be appropriately qualified and trained in all procedures to ensure that good surgical technique is practiced. Good technique includes:

- Aseptic technique
- Gentle tissue handling
- Minimal dissection of tissue
- Maintain tissue hydration
- Appropriate use of instruments
- Effective hemostasis
- Use of suture materials and patterns or other wound closure techniques that minimize trauma and remain intact.

In addition to clinically sound techniques, a surgical plan, as described in the ASP, should also consider the availability of personnel to provide anesthetic induction, aseptic preparation of the surgical site, and post-operative care appropriate to the surgical procedure.<sup>9</sup> Investigators must assure that the challenges of consecutive surgeries within one work session are adequately addressed. A surgical plan should include:

- Analgesia
- Preservation of corneal integrity/hydration
- Nutritional support
- Maintenance of body temperature
- Hydration

## **Procedures**

### **Personal Protective Equipment:**

- Clean covering such as a lab coat worn over work clothes
- Mask
- Gloves
  - Using sterile surgical gloves allows you to touch all areas of the sterile surgical field and surgical instruments with your gloved hand.
- Hair cover

### **Pre-Operative:**

- Surgery should be conducted in a disinfected, uncluttered area that promotes asepsis during surgery. Hard surfaces such as tabletops and non-surgical equipment should be disinfected prior to setting up surgical area (see Table 3).<sup>10</sup>
- Protect eyes from corneal desiccation by applying sterile ophthalmic ointment since anesthesia abolishes the blink reflex.<sup>11,12</sup>
- After anesthetizing the animal, remove the hair from the surgical site by either clipping, plucking, or using a depilatory. This procedure must be done at a separate area and not at the designated animal surgical space. If a depilatory is used, thoroughly rinse the chemical from the rodent's skin or apply a neutralizing agent.
- Administer analgesics (preemptive analgesia) as appropriate and approved in your ASP.
- Take measures to minimize hypothermia by providing heat.<sup>12</sup>
- Prepare the surgical site(s) with an appropriate skin disinfectant (see Table 1).<sup>13</sup> If using a stereotaxic frame, the rodent should be placed in the frame *before* the skin disinfectant is applied.
  - A surgical scrub agent can be alternated three times with 70% alcohol or sterile saline, followed by a final soaking with a disinfectant solution. Alcohol, by itself, is not an

adequate skin disinfectant.<sup>13-15</sup>

- Aseptic scrub skin preparation may contribute to hypothermia. Alternating with alcohol reduces body temperature but results in a rebound phase in which body temperature returns to baseline within a few minutes after application.<sup>14-16</sup>
- Surgeons should wash and dry their hands before aseptically donning sterile gloves.<sup>17</sup>
- Nitrile examination gloves can be either autoclaved or gas sterilized as an economical alternative to pre-packaged sterile surgical gloves.<sup>18</sup> Multiple pairs of nitrile gloves can be autoclaved in the same pack, but care must be used to avoid contamination of the gloves during donning. The same gloves can be worn between surgeries under the following circumstances:
  - The surgeon's gloves have not become contaminated during respective surgeries or
  - The "tips-only" technique is used. Examples of ways to prevent glove contamination are to have another person assist the surgeon by recovering and prepping subsequent animals for surgery, have the surgeon anesthetize and prep all animals having surgery before donning the gloves that s/he will wear during the procedure, etc.<sup>19</sup>
- When feasible, the surgical site should be draped aseptically with sterile material prior to making an incision to create a sterile surgical field. Draping is especially important when suture material will be used.<sup>20</sup>
- Instruments, suture material, suture needle, etc. must never touch outside of the sterile surgical field.
- When working alone and manipulation of non-sterile objects (e.g., anesthesia machines, microscopes, lighting, etc.) is required, it may be helpful to use sterile aluminum foil or sterile plastic covers to manipulate the objects.
- Consult with your IC's Animal Program Director or designee to ensure that your surgery practices meet the standards of aseptic surgery.

#### **Operative:**

- The animal must be maintained in a surgical plane of anesthesia throughout the procedure.<sup>21</sup>
  - If using the pedal withdrawal reflex to test depth of anesthesia, the rear paw has been shown to be more reliable than the forepaw.<sup>11,22</sup>
  - If neuromuscular blocking agents (e.g., pancuronium, succinylcholine) are administered then alternative indicators of anesthetic depth must be monitored.<sup>11</sup> Contact your veterinarian for equipment recommendations and information on how to interpret monitoring results. Animals on neuromuscular blockers must be mechanically ventilated.
- Provide an external heat source (preferably a feedback-controlled, infrared, warm water, or air-circulating heating device) throughout anesthesia and surgery. Hypothermia is a common cause of mortality in rodents undergoing a surgical procedure due to their high surface area to body mass ratio. Contact your veterinarian for information about alternative thermal support devices. Electric heating pads and heat lamps are not recommended because of their potential to cause burns.
- Begin surgery with sterile instruments and devices (e.g., implants and catheters). Handle instruments and devices aseptically (see Table 4).
- When using "tips-only" technique, the sterility of the instrument tips must be maintained throughout the procedure.
- Consider monitoring the animal's vital signs (e.g., respiratory rate, heart rate, body temperature) and tissue hydration.
- Ensure hemostasis and minimize blood loss.

- Close surgical wounds using appropriate techniques and materials (see Table 2).  
Suture gauge selection: Use the smallest gauge suture material that will perform adequately.  
Cutting and reverse cutting needles: Provide edges that will cut through dense, difficult to penetrate tissue, such as skin.  
Non-cutting, taper point or round needles: Have no edges to cut through tissue; used primarily for suturing easily torn tissues such as peritoneum or intestine.
- When surgical procedures are performed in series, utilized instruments, suture material and sterile gloves for multiple animals may be considered. Individual IC ACUC should base the number of animals undergoing a surgical procedure with the same sterile instrument pack, gloves, and suture package on performance standards to ensure animal welfare. In general, the recommendations are:
  - Instruments:
    - Begin with sterile instruments and utilize a “tips-only” technique to sterilize the tips of the instruments between each procedure using a hot bead sterilizer
    - Begin with sterile instruments and utilize new set or bead sterilize instruments between cages. <sup>8,23</sup>Clean instrument of blood and organic material prior to bead sterilizer or other sterilizing method.
    - Assure instruments are cooled after bead sterilizing, before touching tissue (sitting at room temperature for several minutes or dipping in sterile saline). Rinse alcohol after soaking in sterile saline or sterile water.
  - Gloves:
    - Begin with sterile gloves and utilize a tips only surgery technique to prevent cross-contamination
    - Begin with sterile gloves and dip the fingers of the gloves in alcohol for 30 seconds between surgeries to sanitize them<sup>23</sup>
    - Consider changing gloves between cages
  - Suture:
    - When using the same suture pack across serial surgeries to close muscle or skin the animal should be draped in and suture material must remain in the sterile field a between cages.
    - Use new suture pack between cages.
    - Suture selection may vary on surgical procedure, location of closure (deep vs superficial skin), and degree of tension. Recommend seeking veterinary guidance on suture selection.

#### **Post-Operative:**

- Move the animal to a warm, clean, dry area and continue to monitor during recovery. Return the animal to its routine housing only after it has exhibited the righting reflex.
- Continue to provide analgesics as appropriate and approved in your ASP.
- If appropriate, consider giving warm fluids and/or nutritional support.<sup>7</sup>
- Animals must be monitored and evaluated post-operatively. Some examples of monitoring parameters which may be employed include body weight, grimace scale, nesting behavior, or hydration status. Or refer to the [ARAC Guideline Pain and Distress in Laboratory Animals: Responsibilities, Recognition and Alleviation](#). Frequency and duration of post-operative evaluation are established in consultation with veterinary staff.
- Generally, remove skin closures 7 to 14 days post-operatively after verifying that the wound has healed.

**Surgical Record:**

- Creating and maintaining a surgical record with important operative and post-operative information (e.g., annotate cage card with procedure and date, body weight on the day of surgery, analgesic administration, wound closure removal, etc.) is required.
- Continue frequent monitoring of the animal post-surgery until stable (e.g., body weight, body condition, cage activities)

**Table 1. Skin Disinfectants**

AGENT	*EXAMPLES	COMMENTS
Iodophors	Betadine®, Prepodyne®, Wescodyne®	Reduced activity in presence of organic matter. Wide range of microbicidal action. Works best in pH 6-7.
Chlorhexidine	Nolvasan®, Hibiclens®	Presence of blood does not interfere with activity. Rapidly bactericidal and persistent. Effective against many viruses. Excellent for use on skin.
*The use of common brand names as examples does not indicate a product endorsement. ** alcohol is not a disinfectant by itself. It should be used with other skin disinfectants.		

**Table 2. Wound Closure Selection**

MATERIAL*	CHARACTERISTICS AND FREQUENT USES
Polyglactin 910 (Vicryl®), Polyglycolic acid (Dexon®)	Multifilament, Absorbable in 60-90 days; 25-50% loss of tensile strength in 14-21 days. Ligate or suture subcutaneous tissues where an absorbable suture is desirable. Not routinely recommended for skin closure due to high capillarity.
Polydioxanone (PDS®) or, Polyglyconate (Maxon®)	Monofilament, Absorbable in 6 months; 40% loss of tensile strength in 30-42 days. Ligate or suture tissues especially where an absorbable suture and extended wound support is desirable.
Polypropylene (Prolene®)	Monofilament, Non-absorbable. Inert.
Nylon (Ethilon®)	Monofilament, Non-absorbable. Inert. General skin closure.
Silk	Multifilament, Non-absorbable. (Caution: Tissue reactive and may wick microorganisms into the wound, so silk is not recommended for skin closure). Excellent handling. Preferred for cardiovascular procedures.
Stainless Steel Suture/Wound Clips/Wound Staples	Non-absorbable. General skin closure. Requires instrument for removal.
Cyanoacrylate (Vetbond®, Nexabond®, Tissue Mend®)	Tissue Adhesive, for non-tension bearing wounds.
*The use of common brand names as examples does not indicate a product endorsement.	

**Table 3. Recommended Non-Porous Disinfectants**

AGENT	EXAMPLES*	COMMENTS**
Alcohols	70% ethyl alcohol 85% isopropyl alcohol	Flammable*** When submerging instruments, recommended contact time is 15 minutes. Contaminated surfaces take longer to disinfect. Remove gross contamination before using.
Quaternary Ammonium	Roccal®, Quatricide®	Corrosive***. Rapidly inactivated by organic matter. Compounds may support growth of gram-negative bacteria.
Chlorine	Sodium hypochlorite (Clorox® 10% solution) Chlorine	Corrosive***. Presence of organic matter reduces activity. Chlorine dioxide must be fresh; kills vegetative organisms within 3 minutes of contact.
Phenolics	Lysol®, TBQ®	Less affected by organic material than other disinfectants.
Chlorhexidine	Nolvasan®, Hibiclens®	Presence of blood does not interfere with activity. Rapidly bactericidal and persistent. Effective against many viruses.
3 % or 6% Hydrogen peroxide, Accelerated Hydrogen Peroxide	3 % or 6% Hydrogen peroxide, Peroxigard®	This product is a one-step disinfectant cleaner and deodorant designed for general cleaning, disinfecting, and deodorizing of hard, non-porous surfaces.
Blend of peracetic acid, hydrogen peroxide, and acetic acid	Spor Klenz	Respiratory irritant***. Contact time 10 minutes. DOHS approved SOP is strongly recommended.
<p>*The use of common brand names as examples does not indicate a product endorsement.  ** Always follow manufacturer's instructions for dilution and expiration periods. All agents require personal protective equipment, and staff need to be aware of appropriate gloves and other items to ensure worker protection from chemical exposure.  *** Please read the SDS for hazardous compounds and follow DOHS recommendations</p>		

**Table 4. Recommended Sterilant for Surgical Instruments, Devices & Equipment**

AGENT	*EXAMPLES	COMMENTS
Steam Sterilization (moist heat)	Autoclave	Effectiveness dependent upon temperature, pressure and time, e.g., 121°C for 15 min vs 131°C for 3 min. Appropriate sterilization indicators should be used to ensure sterility.
Dry Heat	Hot Bead Sterilizer Dry Chamber	Fast Instruments must be cooled before contacting tissue. Only tips of instruments are sterilized with hot beads.

Gas sterilization	Ethylene Oxide, Vaporized hydrogen peroxide (VHP) Chlorine dioxide	Gas/ Vapors are irritating to tissues, levels required for sterilization are dangerous to personnel life and health; all materials require safe airing time and monitoring. Appropriate sterilization indicators should be used to ensure sterility.  DOHS clearance is required, approved SOP and trained staff when using gas sterilization***
Chlorine	Sterilant Levels of Chlorine dioxide (Clidox®, Alcide®) Sodium hypochlorite (Clorox® 10% solution)	Corrosive to instruments. Items must be clean and free of organic material. Instruments must be rinsed with sterile saline or sterile water before use. Use of room sprayers or decon units, please refer to requirements for room sterilization due to hazards to personnel. General use for room cleaning require standard lab PPE.
Glutaraldehydes	Glutaraldehyde (Cidex®, Cetylcode®, Metricide®)	Several hours required for sterilization. Corrosive and irritating. Instruments must be rinsed with sterile saline or sterile water before use. Product expiration dates must be adhered to as per manufacturer's instructions.  DOHS clearance, approved SOP and trained staff when using glutaraldehydes sterilization***
Blend of Peracetic acid, Hydrogen peroxide, and Acetic acid	Actril®	Respiratory irritant***. Several hours required for sterilization. Corrosive and irritating; only to be used in areas with strong ventilation. Instruments must be rinsed with sterile saline or sterile water before use. Use as room disinfectant or decontaminant requires DOHS review and approval, plus a risk assessment for PPE.
<p>* The use of common brand names as examples does not indicate a product endorsement. Note: Always follow manufacturer's instructions for dilution, exposure times and expiration periods. ** Alcohol is neither a sterilant nor high level disinfectant<sup>23,24</sup> *** Please read the SDS for hazardous compounds and follow DOHS recommendations</p>		

## References

1. Flecknell, P. Rodent analgesia: Assessment and therapeutics. *Vet J* **232**, 70-77, doi:10.1016/j.tvjl.2017.12.017 (2018).
2. Buitrago, S., Martin, T. E., Tetens-Woodring, J., Belicha-Villanueva, A. & Wilding, G. E. Safety and efficacy of various combinations of injectable anesthetics in BALB/c mice. *Journal of the American Association for Laboratory Animal Science : JAALAS* **47**, 11-17 (2008).
3. ACLAM Position Statement on Rodent Surgery. *Journal of the American Association for*

- Laboratory Animal Science : JAALAS **55**, 822-823 (2016).
4. Brown, M. J., Pearson, P. T. & Tomson, F. N. Guidelines for animal surgery in research and teaching. AVMA Panel on Animal Surgery in Research and Teaching, and the ASLAP (American Society of Laboratory Animal Practitioners). *Am J Vet Res* **54**, 1544-1559 (1993).
  5. Horn, C. C. et al. Why Can't Rodents Vomit? A Comparative Behavioral, Anatomical, and Physiological Study. *PLOS ONE* **8**, e60537, doi:10.1371/journal.pone.0060537 (2013).
  6. NRC. Guide for the Care and Use of Laboratory Animals: Eighth Edition. (The National Academies Press, 2011).
  7. Bennett K, Lewis K. Sedation and Anesthesia in Rodents. *Vet Clin North Am Exot Anim Pract.* 2022 Jan;**25**(1):211-255. doi: 10.1016/j.cvex.2021.08.013.
  8. Holdridge JA, Nichols MS, Dupont WD, Jones CP, Shuster KA. The Effectiveness of Hot Bead Sterilization in Maintaining Sterile Surgical Instrument Tips across Sequential Mouse Surgeries. *J Am Assoc Lab Anim Sci.* 2021 Nov 1;**60**(6):700-708. doi: 10.30802/AALAS-JAALAS-21-000047.
  9. Clevenger, R. R. et al. in *Management of Animal Care and Use Programs in Research, Education, and Testing* (eds Robert H Weichbrod, Gail A Thompson, & John N Norton) (CRC Press Taylor & Francis Group, 2018).
  10. Rutala, W. A. APIC guideline for selection and use of disinfectants. 1994, 1995, and 1996 APIC Guidelines Committee. Association for Professionals in Infection Control and Epidemiology, Inc. *Am J Infect Control* **24**, 313-342, doi:10.1016/s0196-6553(96)90066-8 (1996).
  11. Navarro KL, Huss M, Smith JC, Sharp P, Marx JO, Pacharinsak C. Mouse Anesthesia: The Art and Science. *ILAR journal.* 2021 Dec 31;**62**(1-2):238-273. doi: 10.1093/ilar/ilab016.
  12. Gargiulo, S. et al. Mice anesthesia, analgesia, and care, Part I: anesthetic considerations in preclinical research. *ILAR journal* **53**, E55-E69, doi:10.1093/ilar.53.1.55 (2012).
  13. Kroner, K. T., Budgeon, C. & Colopy, S. A. Update on Surgical Principles and Equipment. *Veterinary Clinics: Exotic Animal Practice* **19**, 13-32, doi:10.1016/j.cvex.2015.08.011 (2016).
  14. Del Valle, J. M. et al. Comparison of Aqueous and Alcohol-based Agents for Presurgical Skin Preparation Methods in Mice. *Journal of the American Association for Laboratory Animal Science : JAALAS* **57**, 401-414, doi:10.30802/AALAS-JAALAS-17-000128 (2018).
  15. Kick, B. L., Gumber, S., Wang, H., Moore, R. H. & Taylor, D. K. Evaluation of 4 Presurgical Skin Preparation Methods in Mice. *Journal of the American Association for Laboratory Animal Science : JAALAS* **58**, 71-77, doi:10.30802/AALAS-JAALAS-18-000047 (2019).
  16. Skorupski, A. M., Zhang, J., Ferguson, D., Lawrence, F. & Hankenson, F. C. Quantification of Induced Hypothermia from Aseptic Scrub Applications during Rodent Surgery Preparation. *Journal of the American Association for Laboratory Animal Science : JAALAS* **56**, 562-569 (2017).
  17. FOrcE, H. H. T. in *Morbidity and Mortality Weekly Report Vol. 51* (ed Epidemiology Program Office) (CDC, Atlanta, GA, October 25, 2002).
  18. LeMoine, D. M., Bergdall, V. K. & Freed, C. Performance analysis of exam gloves used for aseptic rodent surgery. *Journal of the American Association for Laboratory Animal Science : JAALAS* **54**, 311-316 (2015).
  19. Hoogstraten-Miller, S. L. & Brown, P. A. Techniques in aseptic rodent surgery. *Current protocols in immunology* **Chapter 1**, Unit-1.1.14, doi:10.1002/0471142735.im0112s82 (2008).
  20. Emmer, K. M., Celeste, N. A., Bidot, W. A., Perret-Gentil, M. I. & Malbrue, R. A. Evaluation of the Sterility of Press'n Seal Cling Film for Use in Rodent Surgery. *Journal of the American Association for Laboratory Animal Science : JAALAS* **58**, 235-239, doi:10.30802/AALAS- JAALAS-18-000096 (2019).
  21. Hein, M., Roehl, A. B. & Tolba, R. H. in *Small Animal Imaging: Basics and Practical Guide* (eds Fabian Kiessling, Bernd J. Pichler, & Peter Hauff) 117-126 (Springer International Publishing, 2017).



22. Samuel N, Taub AH, Paz R, Raz A. Implicit aversive memory under anaesthesia in animal models: a narrative review. *Br J Anaesth*. 2018 Jul;**121**(1):219-232. doi: 10.1016/j.bja.2018.05.046.
23. Keen, J. N., Austin, M., Huang, L.-S., Messing, S. & Wyatt, J. D. Efficacy of soaking in 70% isopropyl alcohol on aerobic bacterial decontamination of surgical instruments and gloves for serial mouse laparotomies. *Journal of the American Association for Laboratory Animal Science: JAALAS* **49**, 832-837 (2010).
24. Huerkamp, M. J. Alcohol as a disinfectant for aseptic surgery of rodents: crossing the thin blue line? *Contemp Top Lab Anim Sci* **41**, 10-12 (2002).

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