

Guidelines for Survival Bleeding of Mice and Rats

These guidelines have been developed to assist investigators and NIH Animal Care and Use Committees (ACUC) in their choice and application of survival rodent bleeding techniques. The guidelines are based on peer-reviewed publications^{1,2,3,4,5,6} as well as on data and experience accumulated at NIH. The Investigator and veterinary staff should decide which method of blood withdrawal to use. As with any procedure, training is critically important. The comfort and level of skill of an investigator with a procedure as well as the sample volume and frequency of sampling should be considered when choosing a method. For example, some institutions recommend the use of retro-orbital bleeding only for specific applications and encourage investigators to use other techniques, whereas other institutions place no restrictions on retro-orbital bleeding performed by trained investigators. It is the responsibility of both the investigator and ACUC to ensure use of techniques and procedures which result in the least pain and distress to the animal, while adequately addressing the needs of the experimental design. Any exceptions to these guidelines, e.g., increase in blood volume to be collected or retro-orbital bleeding without use of anesthesia as outlined below, must be described in IC ACUC guidelines or must be scientifically justified in the ASP and approved by the IC ACUC.

Training and experience of the phlebotomist in the chosen procedure are of paramount importance. Training opportunities and resources, including access to experienced investigators and veterinarians, must be made available to new personnel. Each ACUC should establish lines of accountability to oversee the training of its personnel. The procedures utilized must be reviewed and approved by the ACUC prior to their implementation.

Factors to consider in choosing the blood withdrawal technique appropriate for the purpose at hand include, but are not limited to:

- The species to be bled.
- The size of the animal to be bled.
- The type of the sample required (e.g. serum, whole cells, etc.).
- The quality of the sample required (sterility, tissue fluid contamination, etc.)
- The quantity of blood required.
- The frequency of sampling.
- Health status of the animal being bled.
- The training and experience of the phlebotomist.
- The effect of restraint or anesthesia on the blood parameter measured.

The acceptable quantity and frequency of blood sampling is dependent on the circulating blood volume of the animal and the red blood cell (RBC) turnover rate[‡]. The approximate circulating blood volume of rodents is 55 to 70 ml/kg of body weight. Of the circulating blood volume, approximately 10% of the total volume can be safely removed every 2 to 4 weeks, 7.5% every 7 days, and 1% every 24 hours.^{7,8} Volumes greater than recommended should be justified in the ASP and appropriate fluid and/or cellular replacement provided. Blood sample ranges, based on body weight, are provided in Table 1.

[‡] RBC life span of the mouse: 38-47 days. RBC life span of the rat: 42-65 days.^{9,10,11}

Body weight (g)	*CBV(ml)	1% CBV (ml) every 24 hrs†	7.5% CBV (ml) every 7 days†	10% CBV (ml) every 2 - 4 wks†
20	1.10 - 1.40	.011 - .014	.082 - .105	.11 - .14
25	1.37 - 1.75	.014 - .018	.10 - .13	.14 - .18
30	1.65 - 2.10	.017 - .021	.12 - .16	.17 - .21
35	1.93 - 2.45	.019 - .025	.14 - .18	.19 - .25
40	2.20 - 2.80	.022 - .028	.16 - .21	.22 - .28
125	6.88 - 8.75	.069 - .088	.52 - .66	.69 - .88
150	8.25 - 10.50	.082 - .105	.62 - .79	.82 - 1.0
200	11.00 - 14.00	.11 - .14	.82 - 1.05	1.1 - 1.4
250	13.75 - 17.50	.14 - .18	1.0 - 1.3	1.4 - 1.8
300	16.50 - 21.00	.17 - .21	1.2 - 1.6	1.7 - 2.1
350	19.25 - 24.50	.19 - .25	1.4 - 1.8	1.9 - 2.5
*Circulating blood volume		†Maximum sample volume for that sampling frequency		

The following guidelines refer to the most frequently used survival sampling sites: a) retro-orbital; b) mandibular; c) saphenous; d) tail; and e) jugular. Blood withdrawal by cardiac puncture is considered a euthanasia procedure and should be performed only after ensuring that the animal is under deep anesthesia, as evidenced by lack of response to a painful stimulus (e.g., toe or tail pinch). A list of the issues that should guide the choice of survival blood collection route(s) is noted below, and an abbreviated summary is provided in Table 2.

Retro-orbital Sinus/Plexus Sampling: ¹²

- Retro-orbital sampling can be used in both mice and rats (though not a preferred method in the rat) by penetrating the retro-orbital sinus in mice or plexus in rats with a capillary tube or Pasteur pipette.
- Rapid – large number of animals can be bled within a short period of time.
- Obtainable volume: medium to large.
- Good sample quality. Potential contamination with topical anesthetic, if used, should be taken into account.
- A minimum of 10 days should be allowed for tissue repair before repeat sampling from the same orbit. Otherwise the healing process may interfere with blood flow.
- Alternating orbits should not be attempted until the phlebotomist is proficient in obtaining samples from the orbit accessed most readily by the dominant hand i.e. a right handed individual should gain proficiency withdrawing samples from the right orbit before attempting to obtain samples from the left orbit.
- In the hands of an unskilled phlebotomist, retro-orbital sampling has a greater potential than other blood collection routes to result in complications.
- In mice, general anesthesia is recommended if compatible with experimental design. If retro-orbital bleeding is conducted without general anesthesia a topical ophthalmic anesthetic e.g. proparacaine or tetracaine drops, must be applied prior to the procedure. ARAC believes that

retro-orbital bleeding performed in mice by a trained practitioner represents “minimal or transient pain and distress” and therefore should be considered a USDA Column “C” procedure.

- In rats, the presence of a venous plexus rather than a sinus can lead to greater orbital tissue damage than in the mouse. General anesthesia must be used unless scientific justification is provided and approved by the IACUC. In addition, a topical ophthalmic anesthetic, e.g. proparacaine or tetracaine drops, is recommended prior to the procedure. ARAC believes that retro-orbital bleeding performed in rats by a trained practitioner represents more than “minimal or transient pain and distress” and therefore should be considered a USDA Column “D” procedure.
- In both mice and rats, care must be taken to ensure adequate hemostasis following the procedure.
- Use of sterile capillary tubes and pipettes are recommended for use to help avoid periorbital infection and potential long-term damage to the eye. The edges of the tubes should be checked for smoothness to also decrease likelihood of eye damage.

Mandibular (Facial vein/artery) Sampling (limited to adult mice):¹³

- Obtainable blood volumes: medium to large.
- Repeated sampling is possible by alternating sides of the face.
- Sample may be a mixture of venous and arterial blood.
- Requires less hands-on training than tail or retro-orbital sampling to reliably withdraw a reasonable quantity of blood.
- Perform on awake animals to achieve proper restraint, which in turn results in proper site alignment and venous compression for good blood flow.
- Can be performed rapidly and with a minimal amount of equipment, allowing for rapid completion.
- Sample volume can be partially controlled with the size of needle (20 gauge or smaller) or lancet (4 mm) used to puncture the site.

Saphenous Sampling (medial or lateral approach):^{14, 15}

- Can be used in both rats and mice by piercing the saphenous vein with a needle.
- Obtainable blood volumes: small to medium.
- Variable sample quality.
- The procedure is customarily done on an awake animal but effective restraint is required.¹⁶
- Requires more hands-on training than tail or retro-orbital sampling to reliably withdraw more than a minimal amount of blood.
- Although more esthetically acceptable than retro-orbital sampling, prolonged restraint and site preparation time can result in increased animal distress when handling an awake animal.
- Temporary favoring of limb may be noted following the procedure.
- Application of sterile petroleum jelly to the site may assist the blood to bead and in turn enhance total blood volumes captured.
- The clot/scab can be gently removed for repeated small samples if serial collection is required.

Lateral Tail Vein or Ventral/Dorsal Artery Sampling:

- Can be used in both rats and mice by cannulating the blood vessel or by superficially nicking the vessel perpendicular to the tail.
- Sample collection by nicking the vessel is easily performed in both species, but produces a sample of variable quality that may be contaminated with tissue and skin products. Sample quality decreases with prolonged bleeding times and “milking” of the tail.
- Sample collection using a needle (cannulation) minimizes contamination of the sample, but is more difficult to perform in the mouse.
- Obtainable volumes for cannulation or nicking: artery – medium to large; vein – small; In general, arterial sampling produces larger volumes and is faster, but special care must be taken to ensure adequate hemostasis. For this reason, the artery should only be used if large volumes are needed.
- Repeated collections possible. With tail nicking, the clot/scab can be gently removed for repeated small samples if serial testing is required (e.g., glucose measures, etc.)
- In most cases warming the tail with the aid of a heat lamp or warm compresses will increase obtainable blood volume.
- Cannulation and tail nicking are routinely done without anesthesia, although effective restraint is required.

Tail Clip Sampling¹⁷

- Can be used in both rats and mice by clipping (e.g. amputating) no more than 1mm of the distal tail in mice or 2 mm in rats
- Produces a sample of variable quality that may be contaminated with tissue and skin products. Sample quality decreases with prolonged bleeding times and “milking” of the tail.
- Obtainable volume: small.
- Repeated collections possible. The clot/scab can be gently removed for repeated small samples if serial testing is required (e.g., glucose measures, etc.)
- In most cases warming the tail with the aid of a heat lamp or warm compresses will increase obtainable blood volume.
- When performing tail clipping, consideration should be given to anesthesia/analgesia, particularly if the tail has been previously clipped for genotyping. If a topical hypothermic anesthetic is used, blood will flow as the tail re-warms. If a local anesthetic is applied, adequate contact time should be allowed for it to take effect.

Jugular Sampling (limited to the rat):

- Obtainable blood volumes: medium to large.
- Results in high quality sample.
- Jugular sampling can be conducted without anesthesia, although the use of anesthesia greatly facilitates the procedure.
- Does not easily lend itself to repeated serial sampling.

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Approved - 2/14/01

Revised - 1/12/05, 12/18/07, 9/03/08, 9/08/10

Table 2: Summary of Blood Sampling Techniques

Route	General anesthesia required	Speed and efficiency		Sample quality		Repeated sampling	Relative volumes obtainable	Species	Comments
		Mouse	Rat	Mouse	Rat				
Retro-orbital	Mouse – Recommended ¹ Rat – yes ²	+++	++	+++	++	Should alternate eyes	Medium to large	Rat, Mouse	Rapid, potential for complications
Mandibular	No	+++	N/A	+++	N/A	Yes	Medium to large	Mouse	Rapid, easy and repeated samples possible
Saphenous	No	++	++	++	++	Yes	Small to medium	Rat, Mouse	Not as rapid as other techniques, low potential for tissue damage
Tail Vein or Artery	No	++ Vein +++ Artery	+++ Vein +++ Artery	± to ++ ³	++ to +++	Yes	Small to medium (vein) Medium to large (artery)	Rat, Mouse	Repeatable, simple, variable sample quality
Tail clip	No	+++	+++	+/-	+/-	Yes	1-2 drops	Rat, Mouse	Repeatable if gently pull scab
Jugular	Recommended	N/A	+ /++	N/A	+++	Difficult	Large	Rat	Limited application, poor for repeated sampling

¹ Topical anesthesia must be used if general anesthesia is not used

² Topical anesthesia is recommended in addition to general anesthesia

³ Depending on method and amount of manipulation