

LAB_023 Tissue Collection - Tail Tip in Mice and Rats

Institutional author: **UQ Biological Resources** AEC Reviewed & Approved: April 2024

Version #7

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LAB_023 Tissue Collection - Tail Tip in Mice and Rats (Expires Dec 2024)

I. OBJECTIVE

To describe the tail tip technique used in mice and rats within UQBR facilities.

NB: The use of (*) indicates this statement is dependent on the facility procedures

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II. SAFETY

- 1. Facility and procedure appropriate PPE use is essential when handling laboratory rodents
- All accidents, injury or near misses are to be reported immediately to the Facility Manager and recorded on a UQ OHS Incident Report Form
 - This procedure has the risk of mouse bite injury take appropriate care.
 - This procedure has a risk of causing musculoskeletal injury when performed regularly consider suitable ergonomic design wherever possible

III. EQUIPMENT

PPE *

Minimum PPE is gloves and gown, additional PPE may be required based on facility or additional risk e.g. working with infectious material.

- Surgery scissors e.g. Iris Scissors
 - These should be in good working order and sharp to avoid discomfort for the rodent
- Disinfectant *
- Tissue collection tubes and tube holder
- Change station or Bio-safety cabinet *
- Snap lock bags/boxes
- Tissue Identification Slips *
- Forceps

IV. PREPARATION

- Turn on Change station or Biosafety Cabinet *
- 2. Clean all surfaces and equipment with disinfectant

This includes disinfecting tools – wiping thoroughly to remove any residue

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IV. PROCEDURE

Administration

Refer to individual facility workflows for administrative requirements

Refer to the animal management database workflows for database requirements

Restraint of Rodents

Refer to LAB 006 Handling and Restraint of Mice

Clean Technique

Use a clean technique when performing this procedure, this will minimise contamination from pathogens and subsequently infection in research animals.

Anaesthesia Training Consideration

UQ Biological Resources offers anaesthetic training courses to all staff and researchers. It is highly recommended that anaesthesia training is completed before anaesthetising rodents. For more information email uqbrtraincomp@uq.edu.au.

Tail Tip Procedure

- Ensure all instruments to be used have been disinfected and organic material removed between animals to prevent cross contamination.
- Note that this technique can only be performed in conscious mice and rats less than 21 days old.
- Rodents older than this must be anaesthetised for the procedure.
- These instructions assume tissue is being collected for genotyping purposes.
- The Code states 3.3.6 Methods used to identify animals must be (i) appropriate for the species and the circumstances (ii) be compatible with the purpose and aims for the project or activity (iii) involve non-invasive methods whenever possible. The use of invasive methods must conform with Clause 3.3.1 (iv) cause the least harm, including pain and distress, to the animals.
- In the unlikely event that genotype results need to be clarified, UQBR technicians may collect one additional tissue sample per rodent. Specific AEC approval is required where additional sampling is commonly requested. Where additional sampling is required, animals will be ear notched to avoid anaesthesia.

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1. Differentiate females and males. Refer to SOP Sexing Rodents

For identification purposes the numbers are consecutively grouped according to sex, the neonates are placed in a clean container to keep the sexes separated.

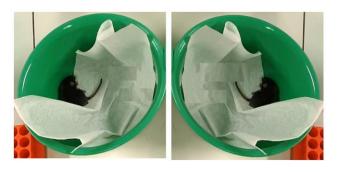


Figure 1. Females and males are separated into clean containers (UQBR 2020).

- 2. If identifying animals assign identification numbers to the litter starting with females

 The animal management database used at UQ is configured will create female ID's first. If tail tipping to
 collect tissue sometimes a separate procedure is performed to also identify the animal.
- 3. Label tissue collection tubes
- 4. Starting with females, restrain the rodent, holding the tail in one hand
- 5. Place the tail into the correct position within the scissors above the collection tube

 This will ensure the small tissue sample falls into the collection tube without contamination.



Figure 2. Placement of scissors and tissue into collection tube (UQBR 2020).

6. Swiftly incise the tip of the tail (maximum 1-2mm)

Ensure the scissors used are well maintained and sharp. Discard all blunted scissors and replace as necessary.

Adverse events should be referred to LAB_022 UQBR Veterinary Care Program.

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7. The tissue will land in the collection tube. Seal the collection tube.

If the tissue falls onto the work surface, use disinfected forceps to collect and place into the tissue collection tube. Note: the tissue sample may fall onto the work surface and be drawn into the Cabinet filter. To prevent losing the tissue sample, complete the work over a tray with raised edges ensuring airflow of the cabinet is not obstructed, or work well inside the work area.

- 8. Apply pressure to stem any blood flow when required
- 9. Disinfect scissors between animals by dipping into 70% ethanol, making sure to wipe dry *

 The scissors are dipped or sprayed with 70% ethanol. The scissors are then wiped dry with clean paper towel.

 This step is required because sensitive genotyping protocols could detect cells from the previous sample causing an inconclusive or incorrect genotype result.



Figure 3. Ethanol filled container to disinfect equipment (UQBR 2020).

- 10. Repeat Steps until all females are toed
- 11. Repeat Steps until all males are toed
- 12. Place collection tubes and tissue identification slip* into snap lock bag or tissue collection box *Follow the specific packing instructions for the genotyping provider that is used.*
- 13. Place the litter into the home cage and return to the rack
- 14. Disinfect the holding containers ready for next use
- 15. Disinfect instruments between litters, making sure to remove all residue
- 16. Continue from Step 1 for remaining litters to be toed
- 17. Place zip lock bag/box into designated fridge or collection point

 Generally tissues samples are stored in a fridge rather than a freezer to avoid degradation of the cells and to improve use of the tissue for genotyping. When tissue is placed into the freezer there is potential for the water molecules expand and burst the cells degrading the DNA.

V. REFERENCE INFORMATION

UQBR Training Consideration

For UQBR training purposes animals may remain for a number of days to monitor. Adverse effects may take time to develop and can assist with the assessment of competency.

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VI. REFERENCES

- National Health and Medical Research Council (NHMRC) 2008, Guidelines to promote the wellbeing of animals used for scientific purpose, viewed 11 April 2019, https://www.nhmrc.gov.au/aboutus/publications/guidelines-promote-wellbeing-animals-usedscientific-purposes
- 2. Office of the Gene Technology Regulator (OGTR) n.d., viewed 11 April 2019, http://www.ogtr.gov.au/
- 3. University of Queensland n.d., *Health, safety and wellbeing,* viewed 11 April 2019, https://staff.uq.edu.au/information-and-services/health-safety-wellbeing
- University of Queensland n.d., Incidents, injuries and hazard, viewed 11 April 2019, https://staff.uq.edu.au/information-and-services/health-safety-wellbeing/health-safetyworkplace/incidents-injuries-hazards
- 5. UQ Biological Resources n.d., *UQBR SOP's*, *viewed 11 April 2019*, https://biologicalresources.uq.edu.au/secure/reference-information#SOP's
- 6. UQ Biological Resources, 2020 UQBR Photo Library.

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