

Institutional Animal Care & Use Program - UTEP	
Title: Guidelines for Euthanasia	
Policy#: 008	Date in Effect: 21 November 2014
Version #: C	Rev Date: 30 June 2025
In Effect <input checked="" type="checkbox"/> Rescinded <input type="checkbox"/>	Date Rescinded:

A) RESPONSIBILITIES

It is the responsibility of all personnel using animals at The University of Texas at El Paso (UTEP) to abide by this policy. Exceptions to this policy must be approved by the Institutional Animal Care and Committee (IACUC) in the protocol to deviate from this policy.

B) APPLICATION

This policy applies to applicable animals used in research and teaching at UTEP.

C) BACKGROUND INFORMATION

To minimize animal suffering, laboratory animals must be euthanized either as described in the protocol at established endpoints, or expeditiously if criteria for humane endpoints have been reached. Animals must be continually observed and never be left unattended during the euthanasia procedure. All methods used must result in the confirmed death of the animal; for most methods this requires a secondary physical method after the primary chemical method to ensure death, such as cervical dislocation, removal of vital organ, decapitation, bilateral thoracotomy, or pithing, depending on the species. Secondary physical methods of euthanasia are also a means to confirm death in reptilian, amphibian, and aquatic species. Animal carcasses and tissues must be properly disposed of after euthanasia.

It is the PI's responsibility to determine that all personnel have been trained to perform the protocol-approved method of euthanasia. Personnel performing euthanasia must follow the approved procedures and apply them humanely and effectively. Training can be provided within the lab group if the existing staff has adequate expertise. Additional training in these techniques is available from LARC (Laboratory Animal Resource Center).

This policy is based on the [2020 AVMA Guidelines for the Euthanasia of Animals](#).

D) RODENT EUTHANASIA

Laboratory rodents should be euthanized in their home cage or a new cage and must not be placed or recombined in unfamiliar groups. Activities that contribute to distress in rodents include transport, handling (in animals not accustomed to it), disruption of compatible groups, and elimination of established scent marks. While eliminating all sources of distress may not be possible, the selected method of euthanizing rodents must minimize these sources of potential distress. Methods of euthanasia likely to elicit distress vocalizations or pheromones that other animals could hear or smell should be performed in another location, if transportation distress can be minimized. Rodent fetuses are in a state of unconsciousness during pregnancy, so euthanasia of the dam is considered sufficient to euthanize fetuses if they remain in the uterus. However, if fetuses must be removed from the uterus for tissue collection or experimentation, they should be treated as altricial neonates.

1. Chemical Methods

a. Carbon Dioxide Inhalation (≥ 10 days old)

- CO₂ exposure using a gradual fill method with a displacement rate from 30% to 70% of the chamber volume/min is recommended; see Appendix A.
- CO₂ must be supplied in a precisely regulated and purified form without contaminants or adulterants, typically from a commercially supplied cylinder or tank. An appropriate pressure-reducing regulator and flow meter is absolutely necessary.
- CO₂ flow should be maintained for at least 1 minute after respiratory arrest.
- Rodents should be kept in their home cage with familiar cagemates during CO₂ administration.
- The practice of immersion, where conscious animals are placed directly into a container prefilled with 100% CO₂, is unacceptable.

b. Injectable Anesthetic Overdose

- Intraperitoneal injection of at least 200 mg/kg sodium pentobarbital is recommended.

- Sodium pentobarbital containing solutions can be viscous and are best diluted to a concentration of no more than 60 mg/ml.
- Other injectable anesthetics may be approved and delivered at an overdose.
- Tribromoethanol is acceptable with conditions as a method for euthanasia of laboratory rodents. It should not be used as a sole agent and must accompany a physical method of euthanasia. Its use requires scientific justification. Recommended dose of greater than 500 mg/kg IP.

c. Inhalant Anesthetic Overdose

- Isoflurane inhalation during an overdose may be utilized as a method of euthanasia, preferably by precision vaporizer. Concentration more than or equal to 5% and duration: at least one minute after breathing stops. Open drop method is acceptable with a minimum of 5% (see table below)...
- TABLE:
(https://www.utep.edu/research/iacup/_files/docs/updated%20policies/iacuc%20policy%20020%20-%20isoflurane%20waste%20anesthetic%20gas%2005.26.20%20final.pdf)

d. Physical Means

- Decapitation or cervical dislocation of adult rodents must be scientifically justified in the protocol. Documented training for this procedure is required.

e. Neonates (<10 days old)

- Anesthetic overdose, as listed in the chemical methods above, can be used.
- Decapitation using scissors or sharp blades is acceptable as a sole means of euthanasia for most protocols.

E. ZEBRAFISH EUTHENASIA

Approved methods for zebrafish vary by age and agent.

1. Adults and Fry (>3 days post fertilization and older)

- Immerse fish in a solution of pharmaceutical grade MS-222 (e.g., [Syncaïne](#)). The solution must be buffered with sodium bicarbonate to a pH of 7.0-7.5.

- Keep fish in an immersion euthanasia solution for at least 30 minutes after cessation of opercular movements when using MS-222. If animals are too young to observe opercular movement, keep immersed for 30 minutes.
 - A secondary method of euthanasia/confirmatory method (exsanguination, decapitation, etc.) is required, even after cessation of opercular movement is noted.
- a. Other solutions inclusive of eugenol, isoeugenol and clove oil require fish to remain in the solution for at least 10 minutes after cessation of opercular movements. Other solutions require fish to remain in the solution for at least 30 minutes after cessation of opercular movements. If animals are too young to observe opercular movement, keep immersed for 30 minutes.
 - b. Rapid chilling
 - Submerge fish in 2-4°C water for 10 minutes for zebrafish >7 days post fertilization and older and 20 minutes for zebrafish 4-7 days post fertilization.
 - c. Sodium or calcium hypochlorite may be used as a single agent on zebrafish 4-7 days post fertilization. Animals should be immersed for at least 5 minutes
2. Embryos (<3 days post fertilization)
 - a. Tricaine, other agents, or rapid chilling may be used as above, but embryos <3 days post fertilization should be followed with an adjunctive method.
 - Adjunctive methods include immersion in sodium (10%) or calcium hypochlorite (10%).
 - b. Sodium or calcium hypochlorite may be used as a single agent on zebrafish embryos <3 days post fertilization. Animals should be immersed for at least 5 minutes.

F. REPTILE AND AMPHIBIAN EUTHENASIA

f. Chemical Methods

- Pharmaceutical grade MS 222 (e.g., [Syncline](#)) can be used either as an injectable agent (except intracoelomic injection in amphibians). Amphibians should be left in this solution for at least 1 hour following cessation of movement. Amphibians may also be fully anesthetized in a

properly buffered TMS 222 bath with a minimum 4-minute immersion that results in cessation of movement prior to application of euthanasia via a physical method. Alternatively, a minimum of 1 hour of immersion is recommended to ensure reliable euthanasia, especially for larger amphibians.


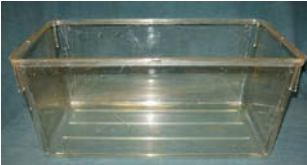

- For reptiles, an intracoelomic injection of 250-500mg/kg of neutral pH solution (0.7% - 1.0% MS 222) typically results in complete loss of consciousness by 4 minutes post-injection; unbuffered, 50% TMS 222 must be administered via intracoelomic injection to complete euthanasia.

g. Injectable Anesthetic Overdose

- Sodium pentobarbital (60 to 100 mg/kg of body weight) can be administered IV, intracoelomically, in the subcutaneous lymph spaces, or in the lymph sacs, although doses vary by species. Please consult with the Attending Veterinarian.

Appendix A

Rodent Euthanasia CO₂ (30 - 70% Concentration) Recommended Flow Rates per Cage Type and Size

Cage Type	Image of Cage	Concentration of CO ₂	Flow Rate Setting on CO ₂ Flow Meter
Rat Tecniplast		30-70%	6.96 -16.24 L/min-
Rat Standard		30-70%	5.64 – 13.16 L/min
Mouse Tecniplast		30-70%	2.58 – 6.02 L/min