

Institute of Experimental Immunology / University of Zürich Viral Immunobiology / Prof. Dr. Christian Münz		
Date 01062015	Standard Operating Procedure <u>Anesthesia</u>	Pages 1/3

Isoflurane anesthesia

1. Assemble the induction chamber with the tube for discharging used isoflurane/oxygen mixture and the tube for delivering fresh isoflurane/oxygen mixture.
2. Clean the induction chamber well with 70% EtOH.
3. Place animals in the chamber.
4. Turn on the oxygen supply.
5. Set vaporizer value to 5% for the induction of anesthesia. Reduce to 3% after induction.
6. Set the flow to 2 liters per minute.
7. Wait until respiration rate is 50-60 per minute.
8. Take animal out and turn of isoflurane and oxygen flow.
9. Do not put animals in chambers already filled with gas. Empty chamber and let the gases flow out before starting another anesthesia.

Isoflurane anesthesia (IVIS)

Turn on "Evacuation Pump" and the oxygen supply.

Turn on gas flow in IVIS Imaging Chamber

Turn on the gas flow to the XGI-8

Set Vaporizer Value to 0% agent

Turn on "IVIS FLOW"

Turn on Flow to Induction chamber

Set the Vaporizer to appropriate setting for animal induction, typically at 2.5%.

Place animal in induction chamber

Turn off "chamber on/off" prior to removing animals from induction Chamber

Place animals in the nose cones

At the end of the procedure: turn the vaporizer to the off position.

Turn on the "Chamber on/off" to allow the pure oxygen to flow through the induction Chamber for 5 minutes.

Turn off the supply oxygen/the XGI-8/the evacuation pump

Injection anesthesia

1. Mice are weighted.
2. Mice are anesthetized by i.p. injection of 0.05 mg/kg Fentanyl, 5 mg/kg Midazolam, 0.5 mg/kg Medetomidin.
3. After onset of narcosis mice are placed on a heating pad to prevent hypothermia.
4. After 5 to 10 min, when surgical tolerance is reached (lack of retraction reflex when pinching the foot between two fingers), animals are allowed to aspirate 25-50µl of influenza A virus.
5. Observe animals for another 5-10 min for complete aspiration of fluid and abnormal breathing rate.
6. Abnormal breathing can be counteracted by placing the animal in an upright position and/or immediate antagonisation of anesthesia.
7. Animals are antagonized by i.p. injection of 2.5 mg/kg Atipamezolin, 1.2 mg/kg Naloxon, 0.5 mg/kg Flumazenil.

Injection anesthesia (alternative)

1. Mice are weighted.
2. Mice are anesthetized by i.p. injection of 100 mg/kg Ketamin, 14 mg/kg Xylazin.
3. After onset of narcosis mice are placed on a heating pad to prevent hypothermia.
4. After 5 to 10 min, when surgical tolerance is reached, animals are allowed to aspirate 25-50µl of influenza A virus.
5. Observe animals for another 5-10 min for complete aspiration of fluid and abnormal breathing rate.
6. Abnormal breathing can be counteracted by placing the animal in an upright position and/or immediate antagonisation of anesthesia.
7. Animals are antagonized by i.p. injection of 5 mg/kg Atipamezolin.

Comment VetA: Fentanyl und Midazolam müssen zuerst antagonisiert werden (mit Naloxon und Flumazenil). Atipamezol darf erst 10 Minuten danach verabreicht werden, falls die Tiere noch nicht wach sind. Ketamin-Xylazin-Anästhesie: Atipamezol darf nicht früher als 60 Minuten nach initialer

Verabreichung von Ketamin gegeben werden, um eine durch einen Ketamin-Hangover verursachte Katalepsie zu verhindern