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# Mouse stereotaxic surgery

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Protocol status: Working

We use this protocol and it's working

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## Disclaimer

The <u>protocols.io</u> team notes that research involving animals and humans must be conducted according to internationally-accepted standards and should always have prior approval from an Institutional Ethics Committee or Board.

### Abstract

This protocol details the mouse stereotaxic surgery.

### **Attachments**



1003-2591.docx

28KB



## **Materials**

#### **Materials:**

• Anesthesia : Inhalant Isoflurane

1. Induction: 3.0 - 3.5% 2. Maintenance: 2.0 - 2.5%

Analgesia:

1. Marcaine (Local)

2. Ethiqa XR / buprenorphine extended-release (Systemic)

■ Sterile 0.9% saline

Sterile ophthalmic ointment

Electric hair clippers

■ 70% ethanol

■ 3% Hydrogen peroxide

Motorized stereotaxic frame

Heating pad

Sterile surgical instruments

Sterile gauze and swabs

Surgical drill

Hamilton syringe

■ Steel needle

Surgical tubing

Pump

Surgical glue

# **Troubleshooting**



## **Procedure**

3d 0h 6m 30s

- 1 Turn on the heating pad to and set to \$\ 36.9 \circ\$.
- 2 Weigh mouse and record weight before starting surgery.
- 3 Place mouse in isoflurane chamber (3% isoflurane).
- 3.1 Record time of anesthesia.
- 4 After mouse is fully anesthetized, use hair clippers to shave head.
- 5 Place mouse back into isoflurane chamber (3% isoflurane) until fully anesthetized.
- 6 Transfer mouse onto stereotaxic frame and place the mouth / nose into nose cone with isoflurane.
- 6.1 Turn down isoflurane to 2%.
- 7 Apply sterile ophthalmic ointment to eyes using a sterile swab (this will prevent desiccation).
- 8 Use ear bars to secure mouse in stereotaxic frame ensuring head is level in all directions.
- 9 Using a sterile swab apply 70% EtOH to the scalp.
- 10 Using a sterile swab, apply povidone-iodine solution to the scalp.
- 10.1 Wait for povidone-iodine solution to dry before making surgical incision.



- Perform a subcutaneous injection of Ethiqa XR (buprenorphine extended-release) into the leg.
- 11.1 Use  $\underline{\underline{L}}$  0.05 mL per  $\underline{\underline{L}}$  20 g mouse.
- Inject  $4 30 \,\mu$  of Marcaine in 2-3 locations underneath the scalp near the incision site. This is a local analysesic.
- 12.1 Wait 30 seconds 00:01:00 for the Marcaine to diffuse before performing incision.

1m

- 13 Using a sterile scalpel, make a surgical incision to expose the skull.
- 13.1 Minimize the size of the incision as much as possible.

#### Note

You will need to see bregma and have access to the injection site. For substantia nigra injections this will be located at the caudal region of the skull.

- 14 Position your injection needle at bregma and save location in the AP and XY axis.
- 15 Enter and save injection coordinates into the motorized stereotaxic frame.
- 15.1 Coordinates for right substantia nigra
  - Anterior-posterior (AP): → ← -2.9 mm
  - Medial-lateral (ML): → ← -1.3 mm
  - Dorso-ventral (DV): -4.2
- Raise the needle away from the skull slightly and move to AP and ML injection coordinates.
- 17 Slowly lower the needle to touch the skull. Raise the needle and drill a small hole where the needle touched the skull.



- 17.1 Slowly drill through the skull keeping the drill shallow enough to not damage brain tissue.
  - 18 Once the hole is drilled lower the needle to the surface of the brain ensuring that the needle is not deflected by the skull.

#### Note

The needle should be completely straight.

- 19 Raise the needle → ← 30 mm - → ← 40 mm providing space to flush the needle and load with virus.
- 19.1 Flush the needle with sterile  $\blacksquare$  Room temperature  $\blacksquare$  H<sub>2</sub>O.
  - Cover the mouse's head with sterile gauze to absorb the H<sub>2</sub>O.
- 19.2 Draw up a  $\perp$  1  $\mu$ L air bubble.
- 19.3
  - Slowly draw up virus and watch the air / liquid interface to determine volume.
  - Needle / tubing should be marked with  $\perp \perp 1 \mu L$  intervals using a sharple to help with this step.
- 20 Move the prepared needle down to the surface of the brain. Move the needle to the desired DV coordinate at a slow speed (250µm / sec).
- 21 Wait 00:00:30 and then start the virus injections at 0.2 μL / min.

30s

22 After the injection is finished wait for 00:05:00, leaving the needle in place and allowing the virus to diffuse away from injection site.

5m

- 23 Remove the needle at slow speed (250  $\mu$ m / sec).
- 24 Move the needle up away from the skull and then release it and rotate it out of the way.



- 25 Close the incision site using sterile forceps and surgical glue.
- 26 Administer 4 0.5 mL sterile saline solution subcutaneously to avoid dehydration.

- 27 Place the mouse back into the home cage and monitor recovery.
- 27.1 Record time at recovery.
- 28 Monitor post operation recover for 5 72:00:00 and record any observations of pain or distress onto surgery cards.

3d