

madaly

madalynn.erb

Preparation of Free Floating Coronal Mouse Brain Sections 44



Protocol Info: madalynn.erb: Preparation of Free Floating Coronal Mouse Brain Sections. protocols.io https://protocols.io/view/preparati on-of-free-floating-coronalmouse-brain-s-c79uzr6w

Created: Jan 26, 2024

Last Modified: Jan 26, 2024

PROTOCOL integer ID: 94228

Keywords: ASAPCRN

Collect Mouse Brain Tissue

- 1 Deeply anesthetize mice via intraperitoneal injection of 2X Avertin solution
- 2 Perform transcardial perfusion using chilled saline solution
 - 2.1 0.9% NaCl kept on ice
 - 2.2 Use approximately 60mL saline solution per mouse
- **3** Switch from saline solution to chilled PFA

🐼 protocols.io

- **3.1** 4% paraformaldehyde in 0.1M phosphate buffer (PB) pH 7.4
- **3.2** Use approximately 60mL PFA per mouse
- 4 Remove brain immediately after PFA perfusion
- 5 Incubate brain in PFA for 24 hours at 4°C
- **6** Transfer brains to 30% Sucrose / 0.1M Phosphate Buffer (PB) keep at 4°C for \ge 24 hours
- 7 Tissue should be completely saturated with sucrose before sectioning

7.1 Brains will sink to the bottom of the vial when saturated

Section Tissue

- 8 Use a Leica SM2010R Microtome
 - 8.1 Blade: Leica 16cm, knife angle set at 0 degree
 - 8.2 Cut thickness: 35µm
- 9 Adjust the microtome platform so that it is level with the blade and lock it into place
- **10** Chill the microtome platform by covering it with crushed dry ice
- **11** Apply 5-10 drops of 30% Sucrose / 0.1M Phosphate Buffer (PB) solution to the chilled microtome platform and wait for it to solidify
- 12 Use the microtome to gently shave the solidified sucrose to make a flat surface
- **13** Use a razor blade to remove any spinal cord from the brain making a flat surface perpendicular to the rostral / caudal axis
- **14** Apply 2-3 drops of sucrose to the existing sucrose platform and quickly position the brain with the olfactory bulbs pointing upwards
 - **14.1** Apply 2-3 more drops of sucrose to the top of the brain to securely freeze it to the microtome platform
- **15** Gently cover the brain in crushed dry ice to freeze the tissue
- 16 Adjust the microtome platform so that the rostral / caudal axis is perpendicular to the blade and the dorsal / ventral axis is level with the blade
- 17 Collect sections in a 24-well plate prefilled with cryoprotectant solution (0.1M PB +30% Sucrose + 30% Ethylene Glycol)

🐼 protocols.io

Seal plate with parafilm and store tissue stored at -20°C