## Brain Microvessel Isolation Kit – Mouse & All Species

Cat. #: P750-4 (4 reactions);

**Ship & Store:** Ship at RT. Upon receiving it, store Solution A & B at 4°C. Store the rest of the kit at RT.

Shelf Life: 3 months

**Description:** Efficient and reliable micro vessel isolation from mouse brain tissue. For research use only.

High yield

Easy-to-follow protocol

## **Product Components**

Components	Amount	Storage
Solution A	100 mL (24.75 mL / rxn)	4°C
Solution B	20 mL (5 mL / rxn)	4°C
Pestle	4	Room temperature
Tissue Strainer 1	4	Room temperature
Tissue Strainer 2	4	Room temperature

## Important note:

All procedures should be done in a clean station.

Always keep Solution A, B on ice. Always use ice cold Solution A, B for whole experiment.

## Protocol:

- 1. Sit Solution A, B on ice.
- 2. Euthanize one mouse and take its brain. This protocol is good for half or one whole mouse brain. Rinse the brain with 1x PBS to clean any debris. Dissect **cerebral cortex**, **hippocampus and stratum of mouse brain**, and put it in 8 mL ice cold Solution A in a 60mm Falcon Petri Dish (Corning #353004).
- 3. Put the dish under a dissection microscope and remove the meninges as soon as possible.
- 4. Make homogenate: Transfer the tissue only into a 1.5 mL microcentrifuge tube. Use a pestle (provided) to homogenize the tissue in the microcentrifuge tube on ice or in cold room:
  - 4.1 First, pestle the tissue without adding Solution A for 100 pestlings.
  - 4.2 Add 0.2 mL Solution A then do another 200 pestlings.
  - 4.3 Add another 0.2 mL Solution A and do another 100 pestlings.

A pestle motor can be used to grind the tissue. However, data shows that hand pestling can produce more consistent result than motor pestling.

- 5. Add 5 mL ice cold **Solution B** to a 50 mL centrifuge tube and put it on ice.
- 6. Pour the homogenate from the 1.5 mL microcentrifuge tube (step 4) into the 50 mL centrifuge tube, which containing 5 mL Solution B. Add 0.4 mL Solution A to the 1.5 mL microcentrifuge tube to rinse, then pour it into the 50 mL centrifuge tube. Repeat the rinse-pour for another 3 times. The total volume in the 50 mL centrifuge tube is 7 mL now (0.2 mL + 0.2 mL + 5 mL + 0.4 mL x 4).
- 7. Add another 3 mL Solution A into the same 50 mL centrifuge tube. The total volume is 10 mL now. Mix well by vortex, then centrifuge at 4,000 rpm for 10 minutes at 4°C.
- 8. Now you can see a fatty layer on the top in the 50 mL centrifuge tube. Remove the fatty layer carefully using 1 mL pipet tips (Cut the end of tip to make a wider opening. This makes the fat removing easier). Now, after removing the fatty layer, there should be 7–7.5 mL homogenate left in the 50 mL centrifuge tube.
- 9. Add an equal volume of Solution A (7–7.5 mL) into the 50 mL centrifuge tube, and vortex briefly to mix well. Then centrifuge at **4,000 rpm for 10 minutes at 4°C**.
- 10. Aspirate and discard the supernatant. Add 2mL Solution A to the pellet and Re-suspend it well by pipetting up and down.
- 11. Put a **Tissue Strainer 1** on a new 50 mL centrifuge tube, then pass the suspension from step 10 through the Tissue Strainer 1 to collect the filtrate.
- 12. Pass the collected filtrate through a **Tissue Strainer 2** on another container. Now the mouse brain microvessels are retained on the **Tissue Strainer 2**.
- 13. Wash the **Tissue Strainer 2** with 0.75 mL Solution A once. Be careful, wash evenly and gently to minimize the loss of some microvessels.
- 14. To collect the microvessels, flip over the **Tissue Strainer 2** and put it on a 35mm culture dish (Corning #430165). Use 1.5 mL Solution A to wash down the microvessels from the **Tissue Strainer 2** to the 35x10mm culture dish.

Now you have the isolated mouse brain microvessels suspended in 1.5 mL Solution A. Transfer all suspension to a **new** 1.5 mL microcentrifuge tube from the 35mm culture dish. Now the mouse brain microvessel isolation experiment is finished. Centrifuge the 1.5 mL tube at **5,000rpm for 10 minutes at 4°C** to collect pellet for downstream protein or/and RNA extraction or do it later by storing at **-80°C** for up to one month.

For downstream experiment:

Use our "Tissue RNA Storage Solution, Cat.# W0592" to store microvessels for up to 1 year.

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