

## Dissociation of NSCs and Neurospheres with Accutase Dissociation of adherent human or rat NSCs

- 1. Aspirate the medium from culture dish
- 2. Add 2 mL of Accutase to culture dish.
- 3. Incubate for 2 to 5 minutes at 37 °C until individual single cells start to round up.
- 4. Gently rinse to remove cells off of the plate's surface.
- 5. Transfer cell suspension to 15 mL conical tube. Gently pipette up and down until cells are in a single cell suspension.
- 6. Add 8 mL of medium to rinse any remaining cells off of the dish's surface and transfer to the conical tube (from Step 5).
- 7. Take a 20uL sample of the cell suspension to determine viable cell density.
- 8. Centrifuge conical tube containing the cell suspension at 200g for 4 minutes.
- 9. Aspirate supernatant, resuspend in fresh medium and plate on coated dish(s). Incubate at 36 to 38°C in a humidified atmosphere of 4 to 6% CO<sub>2</sub> in air.

## Dissociation of human or rat neurosphere cultures

- 1. Remove neurosphere cell suspension from culture dish and transfer to a 15 mL conical tube.
- 2. Let neurospheres settle down in the tube (~2 to 5 minutes) before proceeding to Step 3. Alternatively, the cells can be centrifuged at 100g for 1 minute.
- 3. Gently aspirate medium leaving the neurospheres at the bottom of tube with approximately  $100 \,\mu\text{L}$  of media remaining.
- 4. Resuspend neurospheres in 5 mL DPBS.
- 5. Let neurospheres settle down in the tube (~2 to 5 minutes) before proceeding to Step 6. Alternatively, the cells can be centrifuged at 100g for 1 minute.
- 6. Gently aspirate DPBS leaving the neurospheres at the bottom of tube with approximately 100 μL of DPBS remaining.
- 7. Add 1mL of Accutase® to the neurospheres and incubate 10 minutes at room temperature.
- 8. Using the proper sized pipette tip (i.e.  $1000 \mu l$ ), pipette up and down until all the neurospheres are in a single cell suspension.
- 9. Add 4mL of fresh medium to the tube.
- 10. Centrifuge the cells at 200g for 4 minutes.
- 11. Gently aspirate the supernatant.
- 12. Resuspend cells in fresh medium, transfer to a new culture dish and incubate at 36 to 38°C in a humidified atmosphere of 4 to 6% CO<sub>2</sub> in air.

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