CSI282Ra01

Primary Rat Dorsal Root Ganglion Neuron Cells (DRGN)

Organism Species: Rattus norvegicus (Rat)

Instruction manual

FOR RESEARCH USE ONLY
NOT FOR USE IN CLINICAL DIAGNOSTIC PROCEDURES

3rd Edition (Revised in Feb, 2025)

[DESCRIPTION]

Cell Type: Neuron cell

Synonyms: DGNC

Strain: Sprague Dawley Rat

Age: 1-3 days

Tissue Source: Dorsal root ganglion

Disease: Normal

Size: >5×10^5cell/vial

[PROPERTIES]

Cell activity: >85% (Viability by Trypan Blue Exclusion).

Formulation: Frozen 1 mL or T25 flask.

Biosafety: Negative for HIV-1, HBV, HCV, mycoplasma, bacteria, yeast and fungi.

Applications: For research use only. It is not approved for human or animal use, or for application in clinical diagnostic procedures.

Growth Properties: Adherent

[CONTENTS]

Form & Buffer: Supplied as solution form in frozen stock solution, containing 90% FBS+10% DMSO.

[USAGE]

Upon receiving the cells in a T-25 flask at room temperature, immediately transfer the cells to 37°C, 5% CO_2 incubator; the cells in vials, directly and immediately transfer the cells from dry ice to liquid nitrogen.

Culture conditions:

Special culture medium for neuronal cell:

Neurobasal-A Medium+B-27 Supplement (50X)+1%Penicillin-Streptomycin Solution

Temperature: 37°C

Condition: 95% air, 5% carbon dioxide

Cell recovery:

After receiving the cells, shake at 37°C in a water bath until completely dissolved, transfer to a 15 ml centrifuge tube, add 3-5 times complete culture solution, 1000 rpm for 5 min, discard the supernatant, and place in a T25 flask for culture.

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Cell passage:

Further culture of Primary Rat Dorsal Root Ganglion Neuron Cells are guaranteed under the conditions we provide; however, Primary Rat Dorsal Root Ganglion Neuron Cells are not recommended for expansion or long-term cultures because cells do not proliferate in culture.

[Shipping]

Dry ice.

[STORAGE]

Upon receiving, directly and immediately transfer the cells from dry ice to liquid nitrogen and keep the cells in liquid nitrogen until they are needed for experiments.

[IMPORTANTNOTE]

1. The cultured cycle of Primary Rat Dorsal Root Ganglion Neuron Cells is limited in *vitro*. It is suggested that after cell resuscitation, the special growth medium and correct operation method recommended by us should be used for culture, and it should be used for follow-up experiments as soon as possible.

2. It is recommended that culture bottles be coated with Collagen type I from rat tail, and the concentration of rat tail collagen coating is 0.1mg/ml.

3. The cell is for research use only, and we will not be responsible for any issue if the cell was used in clinical diagnostic or any other procedures.

[Figure]

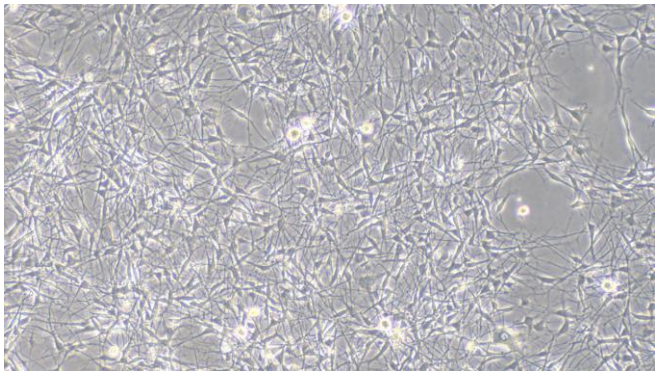

Figure 1

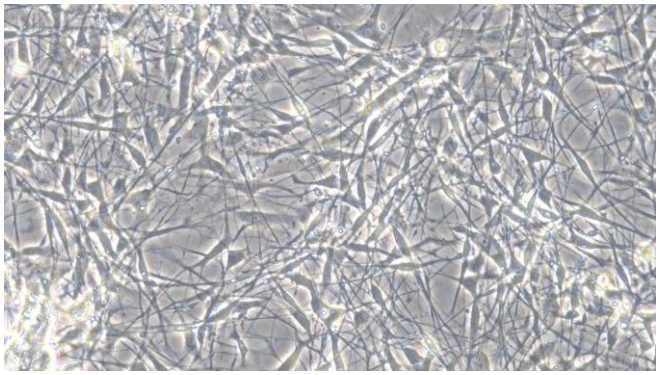

Figure 2

Figure 1 Morphology of Primary Rat Dorsal Root Ganglion Neuron Cells (Optical microscope,×100)

Figure 2 Morphology of Primary Rat Dorsal Root Ganglion Neuron Cells (Optical microscope,×200)

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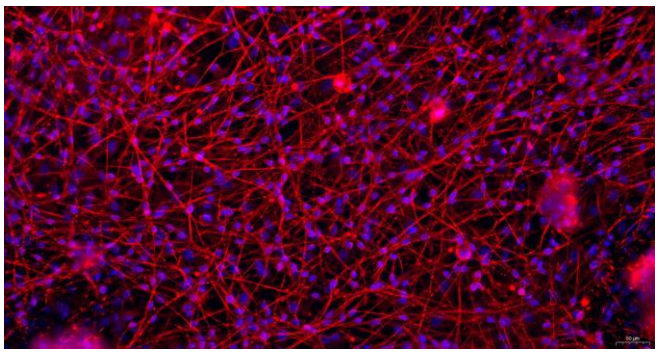

Figure 3

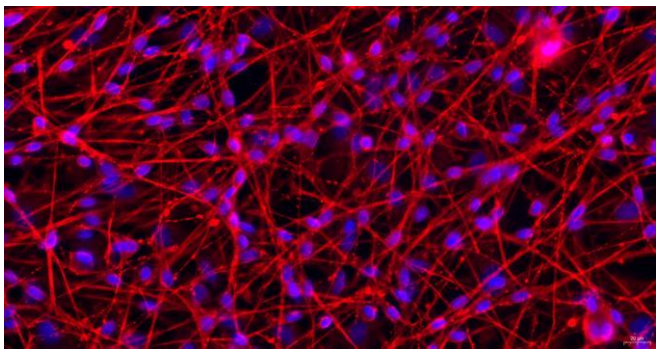

Figure 4

Figure 3 Immunofluorescence identification of Tubulin Beta specific antibody (×200)

Figure 4 Immunofluorescence identification of Tubulin Beta specific antibody (×400)

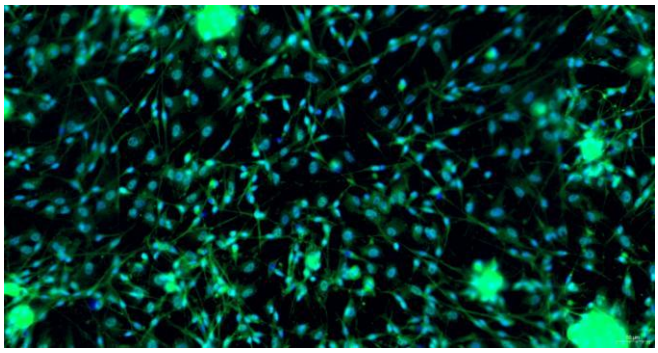

Figure 5

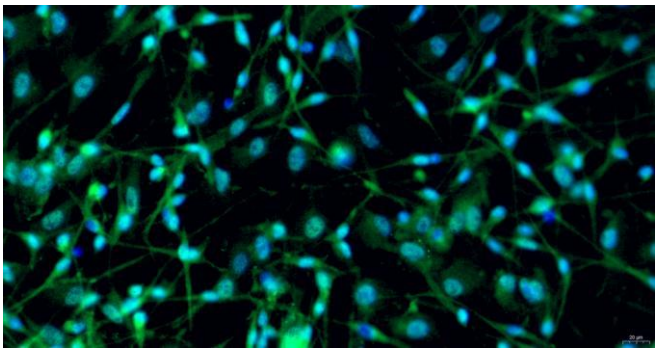

Figure 6

Figure 5 Immunofluorescence identification of MAP2 specific antibody (×200)

Figure 6 Immunofluorescence identification of MAP2 specific antibody (×400)

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