



iPSC-Derived Human Neural Progenitor Cell Kit (African-American, Male Line)

Product Information

Catalog Number ASE-9740

Description Applied StemCell has developed an efficient integration-free method to differentiate high-quality neural progenitor cells (NPCs) from human iPSCs. The differentiated NPCs recapitulate the phenotype and functional parameters of primary and *in vivo* neural progenitor cells.

We provide NPCs differentiated from an integration-free, control human iPSC line (ASE-9211), reprogrammed from the fibroblasts of an African-American male donor. These high-purity ($\geq 95\%$) cells express high levels of the NPC biomarker Nestin (Figure 1).

To harness the full potential of our neural progenitor cells, we also provide optimized NPC Culture Media (ASE-9740MM) that supports robust maintenance and functionality of the NPCs in culture.

These iPSC-differentiated NPCs can be used as control lines to compare phenotype and functionality of patient-derived, genome edited iPSC-derived neural progenitor cells for drug screening applications. The NPCs can also be used for neurotoxicity assays.

Parental Tissue Control human iPSC (ASE-9211); p15
Age: Neonate
Gender: Male
Ethnicity: African-American
Tissue Source: Dermal Fibroblasts
Reprogramming Method: Episomal
Culture Conditions: Feeder-free

Clinical information Healthy (with no known disease phenotypes)

Shipping Dry ice

Storage and Stability Store the components of the kit at the appropriate storage conditions as indicated in the media and materials table, immediately upon arrival. Shelf-life of the product is contingent upon proper storage conditions

Quality Control Each lot of iPSC-derived human neural progenitor cells has been tested for growth, viability and purity ($\geq 95\%$) following recovery from cryopreservation. In addition, each lot has been tested for expression of NPC markers and for the absence of mycoplasma and pathogens.

Safety Precaution **PLEASE READ BEFORE HANDLING ANY FROZEN VIALS.** Please wear appropriate Personal Protection Equipment (lab coat, thermal gloves, safety goggles and a face shield) when handling frozen vials. Please be aware that the following scenario can occur: Liquid nitrogen can leak into the vials when the vials are submerged in liquid nitrogen. Upon thawing, the liquid nitrogen returns to the gas phase, resulting in a dangerous build-up of pressure within the vial. This can result in

Applied StemCell, Inc.

521 Cottonwood Dr. #111, Milpitas, CA 95035

Phone: 866-497-4180 (US Toll Free); 408-773-8007 Fax: 408-773-8238

info@appliedstemcell.com www.appliedstemcell.com

Copyright 2022, Applied StemCell, Inc. All rights reserved. This information is subject to change without notice.

the vial exploding and expelling not only the vial contents but also the vial cap and plastic fragments of the vial.

Warranty

The performance of Applied StemCell's iPSC-derived NPCs has been validated with the NPC Culture Media provided in the Neural Progenitor Cell Kit and the recommended additional reagents. Applied StemCell will not hold responsibility if components other than the components provided with the Neural Progenitor Cell Kit and those recommended are used to culture the Applied StemCell NPCs.

Restricted Use

This product is for research use only and not intended for human or animal diagnostic or therapeutic uses.

Characterization of the ASE-9740 Neural Progenitor Cells

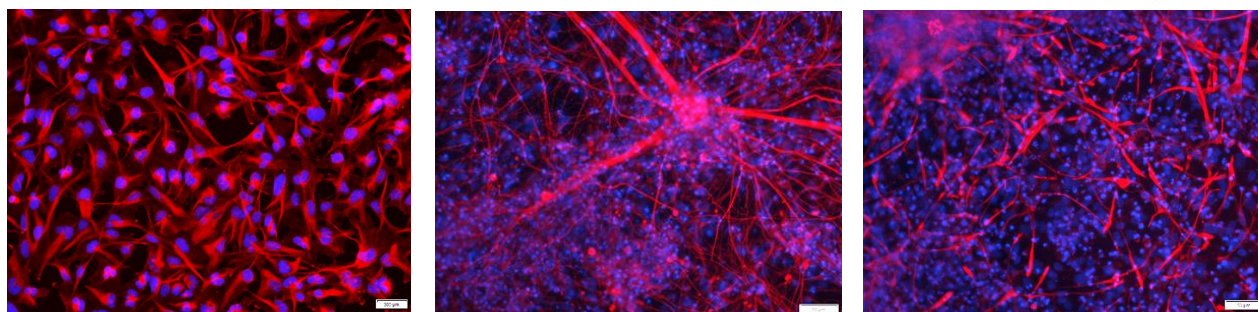


Figure 1. Immunostaining of the ASE-9740 Neural Progenitor Cells derived from Applied StemCell's control iPSC line ASE-9211. The neural progenitor cells (NPCs) (Left, Red: Nestin; Blue: DAPI) can be further differentiated into neurons (Middle, Red: Tuj1; Blue: DAPI) and astrocytes (Right, Red: GFAP; Blue: DAPI).

Media and Material

Neural Progenitor Cell Kit (ASE-9740)

Catalog #	Component	Amount	Storage	Shelf Life
ASE-9740-C	iPSC-Derived Neural Progenitor Cells (NPCs); African-American Male Line	≥1x10 ⁶ cells/ vial	Liq. N2	12 months
ASE-9740MM	NPC Culture Media	100 mL	-20°C	12 months

Additional Reagents Required

The below reagents are recommended for use with the neural progenitor cells. If you use reagents other than those recommended, we suggest that you do a batch-test to validate quality of the cells and culture protocol.

- Matrigel®, Corning, Cat# 354230
- Primary antibodies:
 - Nestin R&D systems MAB1259
 - Tuj1 R&D systems MAB1195
 - GFAP R&D systems AF2954
- Secondary antibodies: corresponding secondary antibodies were purchased from ThermoFisher

Protocol

1. Coating Cell Culture Vessels with Coating Matrix

- 1.1 Coat the plates with 80 µg/mL Matrigel®.
Note: Please follow manufacturer's instructions on coating plates using Matrigel®.
- 1.2 Incubate at room temperature for at least 1 hour before use.

2. Preparation of NPC Culture Media

- 2.1 Thaw the NPC Culture Media at room temperature before thawing the cryopreserved NPCs.
- 2.2 The Culture Media should be aliquoted and stored at -20°C if it will not be used immediately.
Note: The media can be stored at 4°C for up to 2 weeks or at -20°C for up to 12 months.

3. Thawing and Culturing Cryopreserved NPCs

- 3.1 To thaw the cryopreserved NPCs, remove one vial from the storage unit.
- 3.2 Immerse the vial in the water bath (up to 2/3rd of the vial) and thaw the cells rapidly until only a small piece of ice is still visible (approximately one minute).
Note: Do not shake the vial during thawing.
- 3.3 Bring the vial to the biological cabinet immediately and spray the outside of the vial thoroughly with 70% ethanol and wipe it with an autoclaved paper towel.
- 3.4 Remove the cells from the vial using a p1000 micropipette (or serological pipette) and transfer it slowly, drop-wise while swirling into a 15 mL conical tube containing 5 mL of pre-warmed NPC culture medium. Wash the vial with 1 mL medium from the 15 mL conical tube and transfer it back to the tube.
Note: Do not mix cells up and down and avoid generating bubbles.
- 3.5 Centrifuge cells at 250 x g for 5 minutes at room temperature.
- 3.6 Aspirate the medium very carefully using a vacuum (or pipette if preferred), leaving only a drop of liquid in the tube.
Note: Take extra care not to remove or disturb the cell pellet during aspiration of medium.
- 3.7 Using a p1000 micropipette, add 1 mL of the pre-warmed NPC culture medium into the tube and gently re-suspend cells by pipetting up and down 2-3 times.
- 3.8 Remove a 10 µL aliquot of the cell suspension and mix it with 10 µL of Trypan blue solution.
- 3.9 Count the cells.
- 3.10 Aspirate the coating matrix from the pre-warmed cell culture vessel.
- 3.11 Seed the NPCs at a density ranging from 100,000-150,000 live cells/cm² in NPC Culture Media.
- 3.12 Distribute the cells evenly.
- 3.13 Place the cell culture vessels in the incubator (37°C/ 5% CO₂/ humidity control) overnight.
- 3.14 We recommend you fully change the media daily and split the cells every 3-4 days.