

## Cell type: Mono-Allelic mEGFP-tagged TOMM20 WTC hiPSC line

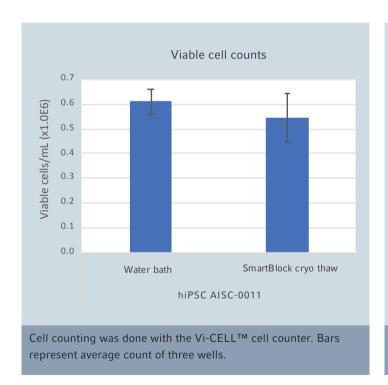
(AISC-0011, Coriell Institute)

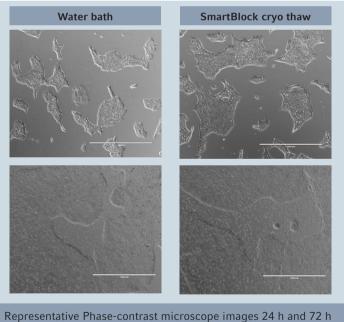
## Performance of the SmartBlock<sup>™</sup> cryo thaw for cell thawing

Freezing medium: mFreSR™ cryopreservation solution (Stem Cell Technologies, 05854)

**Procedure of freezing:** Cells were frozen when reaching a confluency of 60-70 % in 1 mL of cold (2-8 °C) mFreSR medium in Eppendorf CryoStorage Vials 2.0 mL (0030 079.485). Cells were frozen using a slow rate-controlled cooling freezing container (-1°C/minute) (Sigma Aldrich, CLS432000-1EA) at -80 °C for the first 24 hours and later transfer to a liquid nitrogen container in the vapor phase ( -135 °C) for long term storage.

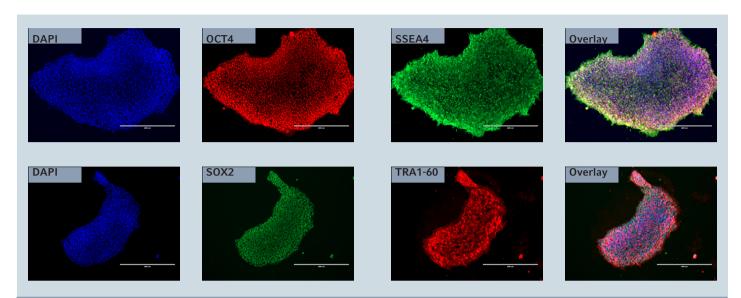
Procedure of thawing: The vials were thawed using the program 'Thawing cells' (3 minutes, 500 rpm). Parallel thawing with a water bath was done. After thawing, cells were suspended in 10 mL Essential 8<sup>™</sup> Flex Medium (Thermo Fisher Scientific, A2858501) and centrifuged at 200x g for 5 min. The cell pellet was resuspended in medium supplemented with RevitaCell™ (Thermo Fisher Scientific, A2644501) and cultivated on Matrigel® (Corning®, 354277) in a 6-well cell culture plate in 2 ml supplemented medium. Cell aggregates were incubated at 37 °C and 5 % CO<sub>2</sub>. Medium refreshment was performed each 24 or 48 hours. About 3-4 days were required between two successive passages.





post-thawing (40× and 100× magnification)





hiPSCs Immunostaining results of hiPSC thawed with the ThermoMixer C confirm the maintenance of pluripotency after 4 successive passages post-thawing (Magnification 100x). Staining was done with the Pluripotent Stem Cell 4-Marker Immunocytochemistry Kit (Thermo Fisher Scientific, A24881)

The program 'Thawing cells' of the Eppendorf SmartBlock cryo thaw allows optimal thawing and homogeneous growth of Mono-Allelic mEGFP-tagged TOMM20 WTC hiPSC line (compared to the existing method (here: water bath).

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