

IPS cell certification

Date: Sep 19, 2024

Cell Line: HEL352 clones HEL352.2 and HEL352.3

iPSC induction: Date of induction 19.02.2024. Fibroblasts were reprogrammed using the CRISPRa method (PMID: 35063129)

Cell line origin: 118 cd40

Investigator: Tapanainen Juha/Khatun Masuma

Culture Description: The iPSC lines were reprogrammed and cultured under feeder-free conditions. They were passaged using 0.5 mM EDTA on Matrigel-coated plates and maintained in E8 medium for over 10 passages.

Characterization:

- i) iPSC morphology assessment: Monitoring typical iPSC colony morphology.
- ii) Culture maintenance: Assessing the stability and survival of iPSC clones over ten passages.
- iii) RT-PCR: Used to confirm the removal of transgene vectors.
- iv) Mycoplasma test: Used to confirm the absence of mycoplasma.

Results:

- i) All iPSC lines maintained typical human pluripotent stem cell morphology during extended culture.
- ii) All clones survived and retained their morphological characteristics over multiple passages.
- iii) Both HEL352 clones (HEL352.2 and HEL352.3) tested negative for the episomal vector.
- iv) Mycoplasma test was negative for all lines.

Interpretation:

Based on the analysis, both HEL352 clones (HEL352.2 and HEL352.3) meet the common criteria for iPSCs.

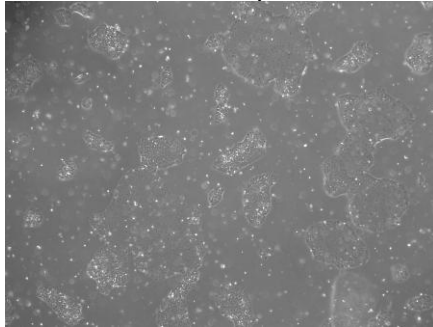
Recommendations:

- Use a Rho-kinase inhibitor (Y-27632) 24 hours after thawing to enhance cell survival.
- Fingerprinting has not been performed; it is important to match the produced iPSC lines to their corresponding donor cell lines.
- It is advisable to analyze the karyotype of the iPSC lines before using them in any experiments.
- Ensure compliance with the ISSCR Guidelines for Stem Cell Research and Clinical Translation ([ISSCR Guidelines](#)) for required characterizations, including morphology assessment, pluripotency verification, and genetic stability.

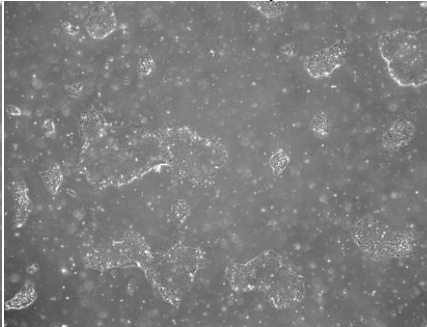
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A) Morphology assessment

HEL352.2 p10.



HEL352.3 p11



B) OriP RT-PCR

