

Certificate of Analysis (CoA) for induced Pluripotent Stem Cells

This product is for research only

Cell Line Name	SIGi001-A-18	Batch / Lot Number	P002
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Reprogramming Method	Integrating Retrovirus (POU5F1, SOX2, MYC and KLF4)		
Genetic Modification	Isogenic modification MAPT (EX10 P301, homozygous; EX10 + 16 bp = C -> T, homozygous) Bi-Allelic MAPT HA-tag		
Passage Number	27	Cell number / vial	1-2 x10 ⁶
Culture Matrix	Matrigel™	Culture Medium	mTeSR™-1
O ₂ Concentration	21%	CO ₂ Concentration	5%
Passaging Method	Accutase	Additional Culture Information	Rho kinase inhibitor used at thaw
Cryopreservation Medium	40% FBS* / 50% mTeSR1 / 10% DMSO *Serum of Zone 1 origin		
Recommendation for thawing	Recommended thaw into 2X well(s) of a 6-well plate or per 10cm ² OR Recommended thaw into 2X 60mm plate(s) Refer to cell line user protocols for further guidance at www.EBiSC.org		
Additional Comments	Slow recovery after thaw, slow growth to confluency		

Please see <https://cells.ebisc.org/> for further information on Quality Control and characterisation applied to lines released by EBiSC. The following standard testing criteria have been determined within EBiSC, prior to release of this product:

Test	Assay	Acceptance Criteria	Result
Vector clearance	RT-PCR	Silenced (Retroviral)	Pass
Sterility	Inoculation for microbiological growth	Not Detected	Pass
	Mycoplasma	Not Detected	Pass
	Virology (HBV, HCV, HIV1, HIV2)	Not Detected	Absence of viral pathogens in other cell line clone from same donor

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Test	Assay	Acceptance Criteria	Result
Cell Line Identity	STR / Fingerprinting	85% match to donor Sex match to donor	Allele data recorded and available upon request. First profile recorded, match to donor.
Viability	Visual Assessment	Growth to confluence post-thaw	Low, slow recovery
Phenotype	Continuous visual assessment of iPSC colony morphology	Recorded	Typical PSC colonies with low differentiation levels
	Flow Cytometry	SSEA-4 > 70% + SSEA-1 < 10% + POU5F1 > 70% +	Pass by central facility
Differentiation Potential	Trilineage differentiation and qPCR for trilineage markers	Up-regulation of germ layer markers	Endoderm: Pass Mesoderm: Pass Ectoderm: Pass by central facility
Genomic Stability	G-Banding (10- 20 successful karyotypes recorded)	Sex match to donor.	No chromosomal abnormalities detected by central facility
Genetic Modification	Sanger sequencing at locus MAPT 17q21.31	Match to reported modification	Pass
	HA-tag on endogenous MAPT gene	Match to reported modification	Pass

Additional guidance on storage, safety and usage can be found in the [EBiSC Technical Information](#).

Approved CoA Signature *P. J. J. J.* Date 03.08.2026