

Certificate of Analysis (CoA) for induced Pluripotent Stem Cells

This product is for research only



Cell Line Name	SIGi001-A-6	Batch / Lot Number	M001
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Reprogramming Method	Integrating Retrovirus (KLF4, MYC, POU5F1, SOX2)		
Genetic Modification	Isogenic modification MAPT (EX10 P301S, homozygous; EX10 + 16 bp = C, homozygous)		
Passage Number	31	Cell number / vial	2,0x10E6
Culture Matrix	Matrigel™ /Geltrex	Culture Medium	mTeSR™-1
O ₂ Concentration	21%	CO ₂ Concentration	5%
Passaging Method	EDTA	Additional Culture Information	Rho kinase inhibitor used at first passage
Cryopreservation Medium	Cryostor		
Recommendation for thawing	Recommended thaw into 60mm plates Refer to cell line user protocols for further guidance at www.EBiSC.org		
Additional Comments	Typical recovery after thaw, typical 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
Sterility	Inoculation for microbiological growth	Not Detected	Pass
	Mycoplasma	Not Detected	Pass
	Virology (HIV1, HIV2, HBV, HCV)	Not Detected	Confirmed pass by depositor
Cell Line Identity	STR / Fingerprinting	85% match to donor Sex match to donor	Allele data recorded and available upon request. Pass
Viability	Visual Assessment	Growth to confluence post-thaw	Acceptable
Phenotype	Continuous visual assessment of iPSC colony morphology	Recorded	Typical PSC colonies with low to medium differentiation levels



In case of queries, please get in touch via Contact@EBiSC.org

SIGi001-A-6.M001.CoA.v2

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Test	Assay	Acceptance Criteria	Result
	Flow Cytometry	SSEA-4 > 70% positive TRA-1-60 > 70% positive SSEA-1 < 10% positive POU5F1 > 70% positive	Pass
Karyotype	Karyolite BoBs	N/A	No autosomal or sex chromosome aneuploidies detected
Genetic Modification	Sanger sequencing at locus MAPT 17q21.31	Match to reported modification	Pass

Additional cell line characteristics have been determined by original reprogramming centres and have not been independently verified by EBiSC. Historical cell line data displayed here is accurate according to data provided by depositors on 02-SEP-2016.

Differentiation Potential	Spontaneous EB differentiation and qPCR for trilineage markers	Up-regulation of germ layer markers	Endoderm : Detected Mesoderm : Detected Ectoderm : Detected
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Additional guidance on storage, safety and usage can be found in the [EBiSC Technical Information](#).

Approved CoA Signature _____ Date _____



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