## **Certificate of Analysis (CoA) for induced Pluripotent Stem Cells**



This product is for research only

ECACC Catalogue No: 66540632

Cell Line Name	BIONi010-C-17	Batch / Lot Number	M001	
Reprogramming Method	Non-integrating Episomal vector (POU5F1, SOX2, MYC, Lin28, shP53 and KLF4)			
Genetic Modification	CRISPR associated (CRISPR/Cas) System			
Passage Number	40	Cell number / vial	3.98x10 <sup>5</sup>	
Culture Matrix	Matrigel <sup>™</sup> / Geltrex <sup>™</sup>	Culture Medium	Essential 8 <sup>™</sup>	
O <sub>2</sub> Concentration	21%	CO₂ Concentration	5%	
Passaging Method	EDTA	Additional Culture Information	Rho kinase inhibitor used at thaw	
Cryopreservation Medium	Cryostor®			
Recommendation for thawing	Recommended thaw into 2 well(s) of a 6-well plate or per 10cm <sup>2</sup> 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 <a href="https://cells.ebisc.org/">https://cells.ebisc.org/</a> 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 (HBV, HCV, HIV1, HIV2)	Not Detected	Confirmed pass by depositor
Cell Line Identity	STR / Fingerprinting	N/A	Allele data recorded and available upon request. Match to donor
Viability	Visual Assessment	Growth to confluence post-thaw	Acceptable
Phenotype	Continuous visual assessment of iPSC colony morphology	Recorded	Typical iPSC colonies with low differentiation levels



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	Flow Cytometry	SSEA-4 > 70% + TRA-1-60 > 70% + SSEA-1 < 10% + POU5F1 > 70% +	Pass
Differentiation Potential	Directed differentiation and qPCR for trilineage markers	Up-regulation of germ layer markers	Endoderm: Pass Mesoderm: Pass Ectoderm: Pass
Genomic Stability	G-Banding	Sex match to donor. 20 successful karyotypes recorded.	46, XY
Genetic Modification	Sanger sequencing at locus	Match to reported modification	TREM2: Gene knockout

Approved CoA	Signature	W. Philoots	Date	05/08/2022	

Additional guidance on storage, safety and usage can be found in the **EBISC Technical Information**.

