

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

*This product is for research only*

ECACC Catalogue No: 66540040

Cell Line Name	UNEWI002-A Alternative name: UNEW002Ai	Batch Number	P001
Donor ID	F150		
Disease Association	Retinitis Pigmentosa	Phenotype of Donor	Affected
Tissue of Origin	Dermal fibroblasts	Sex	Female
Reprogramming Method	Cytotune 2.0 Sendai Vectors ( SOX2, KLF4, MYC, POU5F1)		
Passage Number	Passage 24	Cell number / vial	1-2 x 10 <sup>6</sup>
Culture Matrix	Geltrex/Matrigel	Culture Medium	mTeSR-1
O <sub>2</sub> Concentration	20%	CO <sub>2</sub> Concentration	5%
Passaging Method	EDTA	Additional Culture Information	N/A
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 <a href="http://www.EBiSC.org">www.EBiSC.org</a>		
Additional Comments	Typical recovery after thaw, typical growth to confluency		
Associated Publications	N/A		

Please see [www.EBiSC.org](http://www.EBiSC.org) for further information on Quality Control 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
<b>Sterility</b>	Inoculation for microbiological growth	Not Detected	Pass
	qPCR for Mycoplasma	Not Detected	Pass
	Virology (HBV, HCV, HIV1, HIV2)	Not Detected	Pass
<b>Cell Line Identity</b>	Short Tandem Repeat analysis using PCR	N/A	Allele data recorded and available upon request. Gender match to donor
<b>Viability</b>	Visual Assessment	Growth to confluence post-thaw	Acceptable

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Test	Assay	Acceptance Criteria	Result
<b>Phenotype</b>	Continuous visual assessment of iPSC colony morphology	Recorded	Typical iPSC colonies with low differentiation levels
	Flow Cytometry	SSEA-4 > 70% + TRA-1-60 > 70% + SSEA-1 < 10% + POU5F1 > 70% +	Pass
<b>Differentiation Potential</b>	Spontaneous EB differentiation and qPCR for trilineage markers	Up-regulation of germ layer markers	Endoderm : Detected Mesoderm : Detected Ectoderm : Detected

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 29-JAN-2015

Test	Assay	Result
<b>Genetic Defect</b>	DNA sequencing	Mutation confirmed
<b>Phenotype</b>	Flow cytometry	Positive for markers TRA-1-60, NANOG, SSEA4, low expression of SSEA1
<b>Karyotype</b>	CytoSNP analysis	No clinically significant imbalance was detected
<b>Cell Line Identity</b>	CytoSNP analysis	Parental fibroblasts and clone are identical
<b>Clearance of Reprogramming Factors</b>	PCR for Sendai virus clearance	Reprogramming vector has been cleared
<b>Sterility</b>	qPCR for Mycoplasma	Negative
<b>Differentiation Potential</b>	EB spontaneous trilineage differentiation	Formed all germ layers

The following guidance can be found in the Instructions for Use	
<b>Intended use</b>	<b>Expiry Date</b>
<b>Product Format</b>	<b>Recommended storage conditions</b>
<b>Volume</b>	<b>Hazardous Information</b>

Approved CoA

Signature

Date

03 feb 2016



In case of queries, please contact [culturecollections.technical@phe.gov.uk](mailto:culturecollections.technical@phe.gov.uk). European Collection of Authenticated Cell Cultures (ECACC), Culture Collections, Public Health England, Tel: +44 (0) 1980 612684