Certificate of Analysis (CoA) for induced Pluripotent Stem Cells



This product is for research only

ECACC Catalogue No: 66540005

Cell Line Name	UKBi002-A Alternative name: UKB002Ai	Batch Number	P001
Donor ID	LB-C2-36m		
Disease Association	No Disease Association	Phenotype of Donor	Unaffected Control
Tissue of Origin	Skin Fibroblasts	Sex	Male
Reprogramming Method	Retroviral vector (POU5F1, SOX2, KLF4 and MYC)		
Passage Number	Passage 15	Cell number / vial	1 - 2 x 10 ⁶
Culture Matrix	Matrigel/Geltrex	Culture Medium	mTeSR-1
O ₂ Concentration	20%	CO ₂ Concentration	5%
Passaging Method	EDTA	Additional Culture Information	N/A
Cryopreservation Medium	90% mTeSR-1 Medium +10% DMSO		
Recommendation for thawing	Recommended that	w into 1 well of a 6-well p	plate or per 10cm ²
	Refer to cell line user pr	otocols for further guida	nce at www.EBiSC.org
Additional Comments	A higher split ratio e. reaching confluency. To	g. 1:5 is recommended to o prevent spontaneous d o confluency to exceed 70	avoid cells rapidly ifferentiation, do not
Associated Publications	PubMed-ID 22113611		

Please see 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
	Inoculation for microbiological growth	Not Detected	Pass
Sterility	qPCR for Mycoplasma	Not Detected	Pass
	Virology (HBV, HCV, HIV1, HIV2)	Not Detected	Pass
Cell Line Identity	Short Tandem Repeat analysis using PCR	N/A	Allele data recorded and available upon request. Match to donor
Viability	Visual Assessment	Growth to confluence post-thaw	Acceptable



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Test	Assay	Acceptance Criteria	Result
Phenotype	Continuous visual assessment of iPSC colony morphology	Recorded	Morphology variable (ranging from iPSC emergence to Typical iPSC colonies) and dependent on TC practices. Differentiation Low to Medium
	Flow Cytometry	SSEA-4 > 70% + TRA-1-60 > 70% + SSEA-1 < 10% +	Pass
Differentiation Potential	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 20-APR-2015

Test	Assay	Result
Phenotype	Immunocytochemistry	Positive expression of POU5F1, SSEA-3, TRA-1-60 and TRA-1-81
Karyotype	SNP Karyotyping	Normal
Cell Line Identity	STR	Match to donor tissue.
Clearance of Reprogramming Factors	qPCR	No expression of reprogramming factors
Differentiation Potential	Teratoma Formation Tissues of all 3 germ la	Tissues of all 3 germ layers formed
- Total and Tota	Directed differentiation	Production of It-NES cells

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

Approved CoA

Signature Date 63 feb 2076



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