Certificate of Analysis (CoA) for induced Pluripotent Stem Cells



This product is for research only

ECACC Catalogue No: 66540479

| Cell Line Name | UKKi018-B | Batch Number | P001 |
|------------------------------|--|-----------------------------------|--|
| Donor ID | NP0080 | | |
| Disease Association | Familial Long QT Syndrome | Phenotype of Donor | Affected |
| Tissue of Origin | PBMC | Sex | Female |
| Reprogramming Method | Non-integrating Sendai Virus (KLF4, MYC, POU5F1, SOX2) | | |
| Passage Number | Passage 37 | Cell number / vial | 1.32x10 ⁶ |
| Culture Matrix | Vitronectin | Culture Medium | Essential 8 TM /Essential 8 Flex TM |
| O ₂ Concentration | 20% | CO ₂ Concentration | 5% |
| Passaging Method | EDTA | Additional Culture Information | N/A |
| Cryopreservation Medium | 90% medium / 10% DMSO | | |
| Recommendation for thawing | Recommended thaw into 2 wells of a 6-well plate or per 10cm ² | | |
| Necommendation for thawing | Refer to cell line user protocols for further guidance at www.EBiSC.org | | |
| Additional Comments | Typical recovery after thaw, typical growth to confluency | | |
| Associated Publications | N/A | | |

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 |
|--------------------|--|--------------------------------|--|
| Sterility | Inoculation for microbiological growth | Not Detected | Pass |
| | qPCR for Mycoplasma | Not Detected | Pass |
| Cell Line Identity | Short Tandem Repeat analysis using PCR | N/A | Allele data recorded and available upon request. Gender 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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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 25-MAY-2017

| Test | Assay | Result |
|---------------------------------------|--|--|
| Dhanatana | Flow Cytometry | Positive Expression of CD90, SSEA-1, SSEA-4 and TRA-1-80 |
| Phenotype | Immunocyto-chemistry | Positive expression of TRA-1-80, POU5F1, Nanog and SSEA-4 |
| Karyotype | SNP Analysis (OmniExpress Exome Chip) | No larger chromosomal aberrations observed |
| Cell Line Identity | PowerPlex 16 STR Genotyping System | Match to donor profile |
| Clearance of Reprogramming Factors | PCR for Sendai virus | Not detected |
| Pluripotency | PCR | Pluripotency markers detected |
| Differentiation Potential | Trilineage differentiation | Differentiation to endoderm, ectoderm and mesoderm detected |
| Sterility | Virology (HBV, HCV, HIV1, HIV2) PCR | Not detected |
| Genetic Lesion | DNA sequencing hERG affected location | Mutation hERG: c.173A>G; Glu58Gly confirmed |

| 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 (and less

Date 23 feb 2018

