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



This product is for research only

ECACC Catalogue No: 66540617

| Cell Line Name               | UKBi011-A-1  | Batch / Lot Number                | M001                                 |  |
|------------------------------|--|-----------------------------------|--------------------------------------|--|
| Reprogramming Method         | Sendai CytoTune™ 2.0 (POU5F1, SOX2, cMYC, and KLF4)  |                                   |                                      |  |
| Genetic Modification         | CRISPR-Cas9  |                                   |                                      |  |
| Passage Number               | Passage 30   | Cell number / vial                | 2x10 <sup>6</sup>                    |  |
| Culture Matrix               | Matrigel <sup>™</sup> / Geltrex <sup>™</sup>   | Culture Medium                    | mTeSR <sup>™</sup> -1                |  |
| O <sub>2</sub> Concentration | 21%  | CO <sub>2</sub> Concentration     | 5%                                   |  |
| Passaging Method             | EDTA   | Additional Culture<br>Information | Rho kinase inhibitor<br>used at thaw |  |
| Cryopreservation Medium      | Cryostor®  |                                   |                                      |  |
| Recommendation for thawing   | Recommended thaw into a 50mm cell culture plate<br>Refer to cell line user protocols for further guidance at www.EBiSC.org |                                   |                                      |  |
| Additional Comments          | Slow recovery after thaw, slow 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,<br>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   | Low, slow recovery  |
|                              | Continuous visual assessment of iPSC colony morphology               | Recorded   | Typical PSC colonies with low differentiation levels            |
| Phenotype                    | Flow Cytometry   | SSEA-4 > 70% +<br>TRA-1-60 > 70% +<br>SSEA-1 < 10% +<br>POU5F1 > 70% + | Pass  |
| Differentiation<br>Potential | Spontaneous EB<br>differentiation and qPCR for<br>trilineage markers | Up-regulation of germ<br>layer markers                                 | Endoderm Pass<br>Mesoderm : Pass<br>Ectoderm : Pass             |



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|----------------------|----------------------------|--|---------------------------------------|
| Genomic Stability    | G-Banding                  | Sex match to donor. 20 successful karyotypes recorded. | No chromosomal abnormalities detected |
| Genetic Modification | Sanger sequencing at locus | Match to reported modification                         | ApoE: Gene knockout                   |

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

| Approved CoA | Signature | llus | Date | 28.05.2021 |  |
|--------------|-----------|------|------|------------|--|
|              |           |      |      |            |  |

