UK Dementia Research Institute





Certificate of analysis

DRICUi020-A

Operator: J Winston Date: 09/10/2024 Supervisor: H Hall-Roberts Date: 09/10/2024 Signature: *MRM*

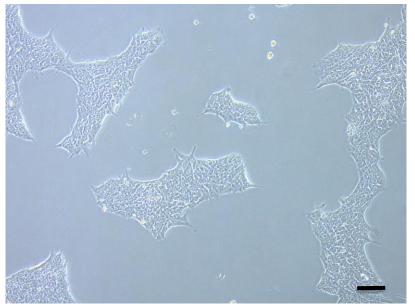
Source of cells and reprogramming information

- CF00C9047a T cells from Cardiff 16/12/2021
- Reprogrammed at UOXF AKA IPMAR22
- Reprogrammed on 04/2022 Sally Cowley (Oxford)
- Reprogramming system Cytotune v2
- Clone DRICUi020-A = IPMAR22A11
- Banked at p21 07/2023 Sarah Ellwood (Oxford)
- Cytotune performed at Oxford by Sarah Ellwood, other QC at Cardiff by Jincy Winston

Viability post-thaw and Morphology according to JMSCFSOP19 passage 22

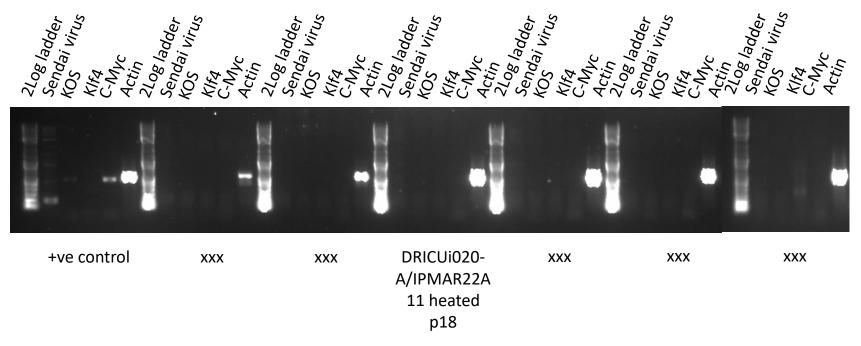
- Vial cell count immediately post-thaw 2.0 x 10⁶
- Viability immediately post-thaw 84%
- Photo at day 2 post-thaw (scale bar = 100μm):

Day 2 post-thaw, 20% plated to 1w.6wp



Sendai Cytotune 2 clearance: according to Cytotune manual

KOS and Klf4 undeterminable but Virus backbone and c-Myc undetectable at passage 18 so overall pass



Product sizes: SeV 181bp; KOS 528bp; SeV-Klf 410bp; SeV-Myc 532bp; Actin 623bp

Sterility:

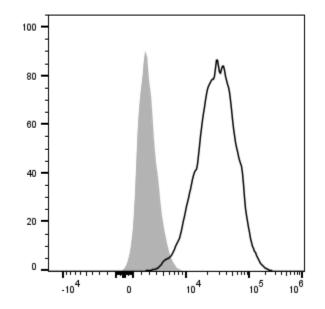
Mycoplasma test performed by Eurofins Genomics on 13/09/2024, undetectable at passage 23.

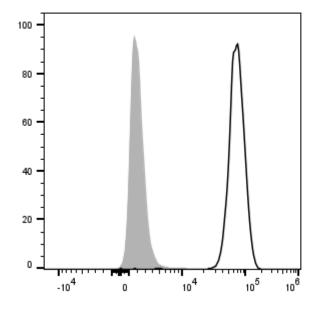
Visual inspection of thawed cells cultured without antibiotic/antimycotic for 4 days: no evidence of bacteria, yeast or fungus.

Flow cytometric analysis according to JMSCFSOP05 passage 22

DRICUI020-A TRA-1-60 98.1%

DRICUI020-A Nanog 98.9%



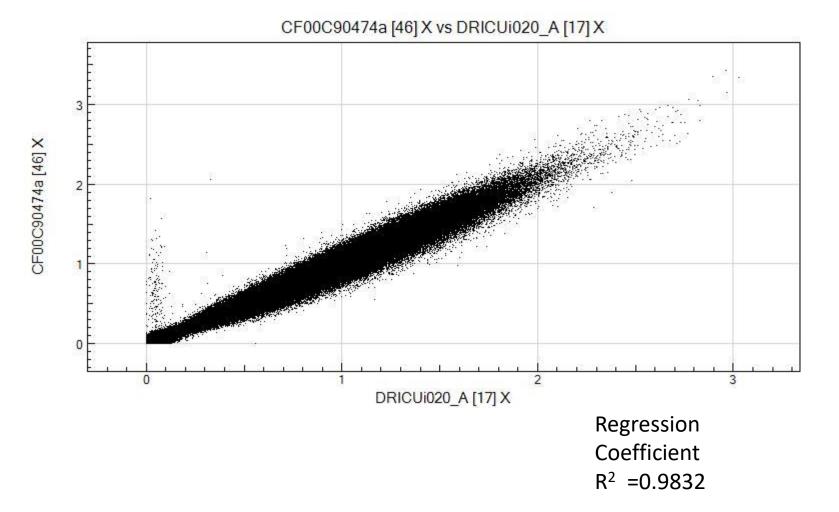


Illumina GSA SNP analysis according to JMSCFSOP16

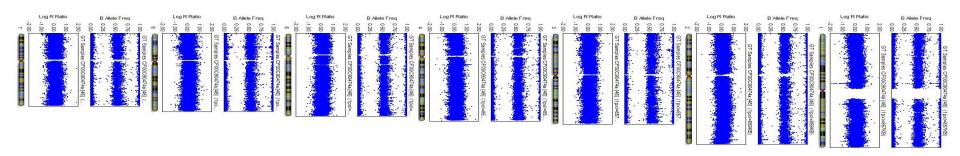
- Passage 21
- Identity to parent PBMC confirmed
- Karyotype abnormalities:

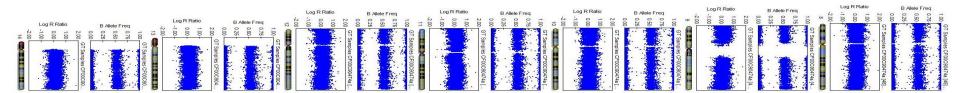
No gross abnormalities detected vs PBMC

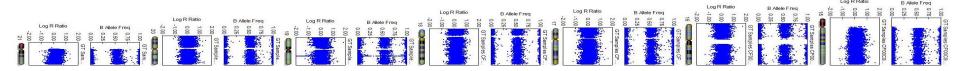
Alignment of CF00C90474A PBMC SNPs with DRICUi020-A

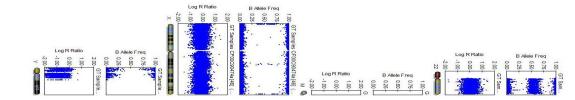


Karyogram CF00C90474A PBMC









Karyogram DRICUi020-A

