



IPMAR

IPSC Platform to Model Alzheimer's Disease Risk

Certificate of analysis

DRICUi008-A

Operators: J Winston/C Bridge/R O'Donoghue

Date: 03/06/2026

Supervisor: H Hall-Roberts

Date: 19/6/2026

Signature: *HCRoberts*

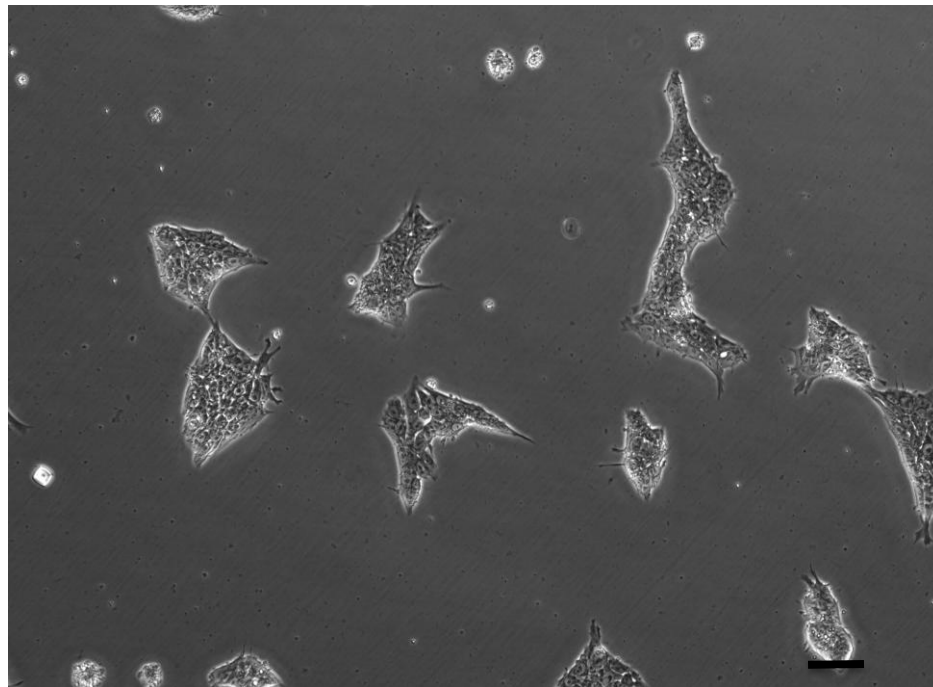
Source of cells and reprogramming information

- ADANG14293UC T cells from Cardiff 27/07/2021
- Reprogrammed at UOXF AKA IPMAR09
- Reprogrammed on 09/2021 Sally Cowley/ Sarah Ellwood (Oxford)
- Reprogramming system Cytotune v2
- Clone DRICUi008-A =IPMAR09E8a4(subclone A7)
- Banked at p24, 12/2023 Emily Maguire (Cardiff)

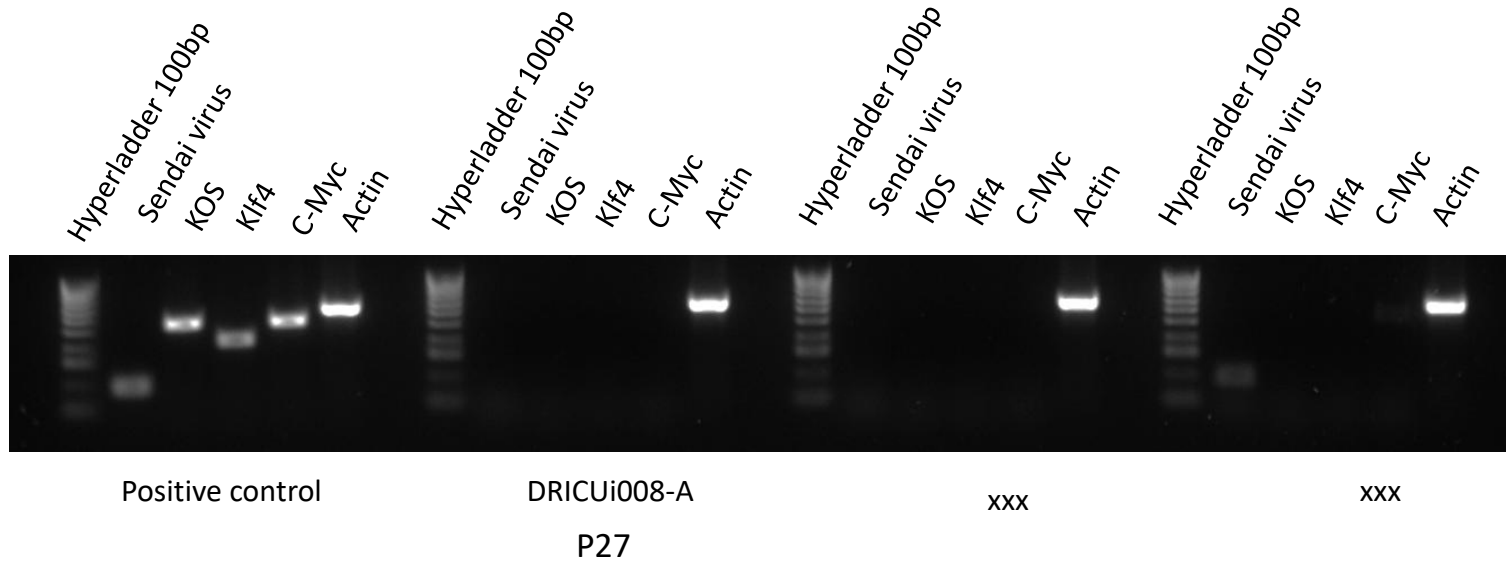
Viability post-thaw and Morphology according to JMSCFSOP19 passage 26

- Vial cell count immediately post-thaw 1.76×10^6
- Viability immediately post-thaw 93.9%
- Photo at day 2 post-thaw (scale bar = $100\mu\text{m}$):

Day 2 post-thaw, 25% plated to 1w.6wp



Sendai Cytotune 2 clearance: according to Cytotune manual Virus undetectable at Passage 27



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

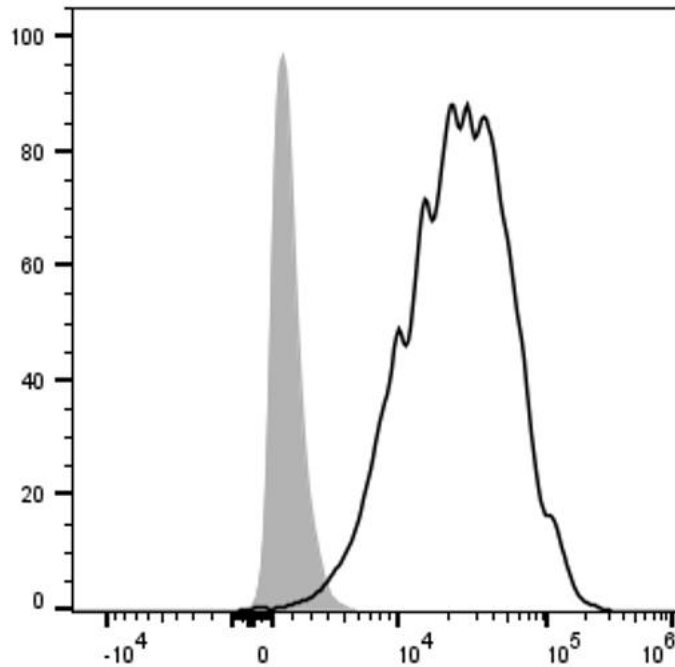
Sterility:

Mycoplasma test performed by Eurofins Genomics on 27/04/26, undetectable at passage 28.

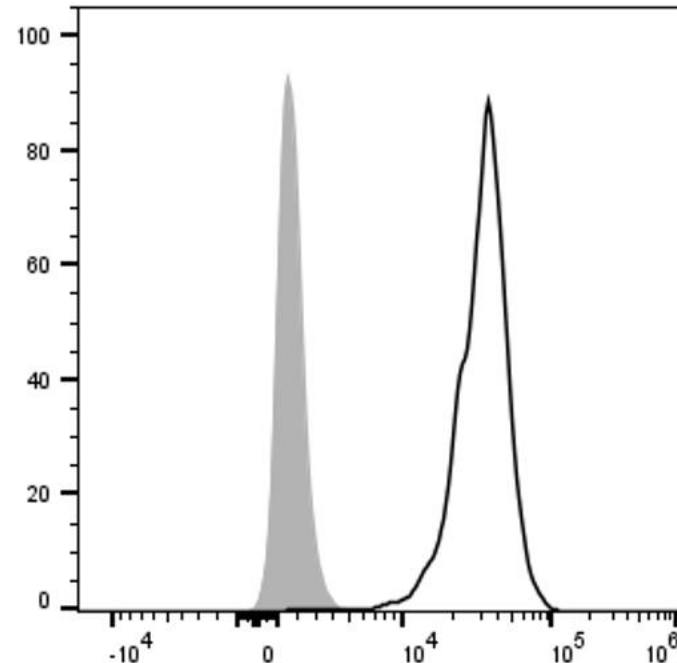
Visual inspection of thawed cells cultured without antibiotic/antimycotic for 4 days:
no evidence of bacteria, yeast or fungus.

Flow cytometric analysis according to JMSCFSOP05 passage 21

DRICUi008-A TRA-1-60 98.7%



DRICUi008-A Nanog 99.1%

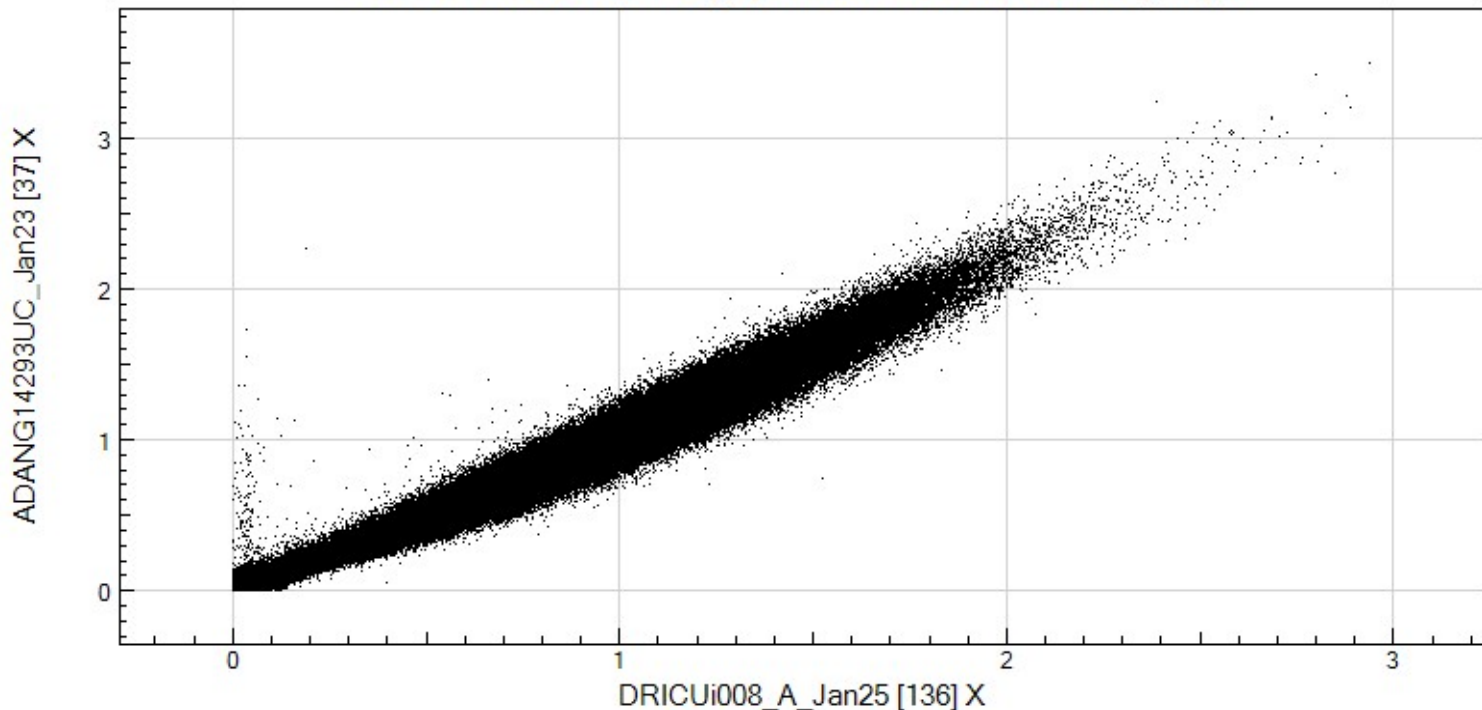


Illumina GSA SNP analysis according to JMSCFSOP16

- Passage 26
- Identity to parent PBMC confirmed
- Karyotype abnormalities:
 - No gross abnormalities detected vs PBMC

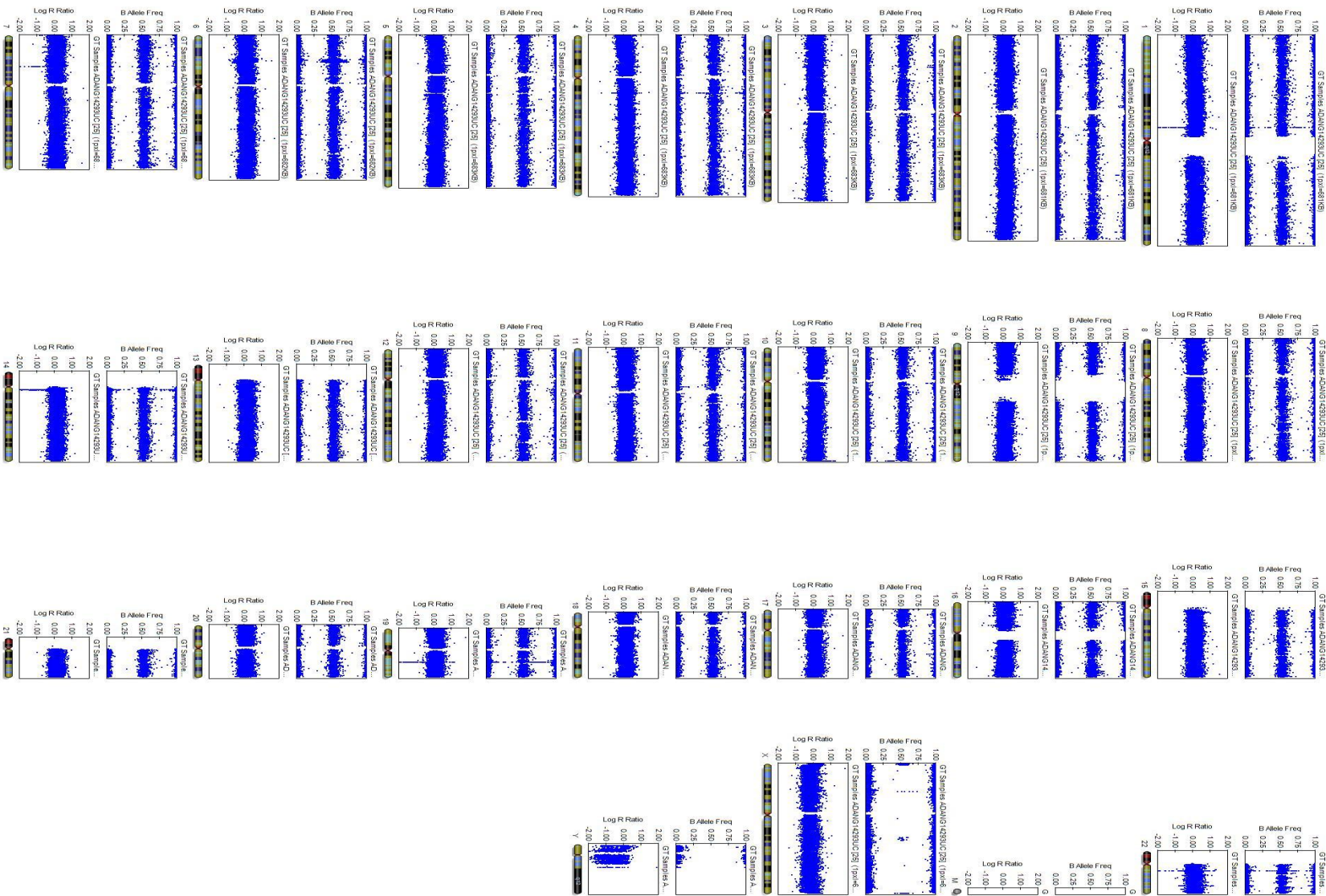
Alignment of ADANG14293UC PBMC SNPs with DRICUi008-A

ADANG14293UC_Jan23 [37] X vs DRICUi008_A_Jan25 [136] X



Regression Coefficient R^2 : 0.9830

Karyogram ADANG14293UC PBMC



Karyogram DRICUi008-A

