Certificate of analysis

SFC126-03-02

Signature: Dario Melguizo Sanchis

Date: 15.06.2015

Supervisor signature: Linda Lako

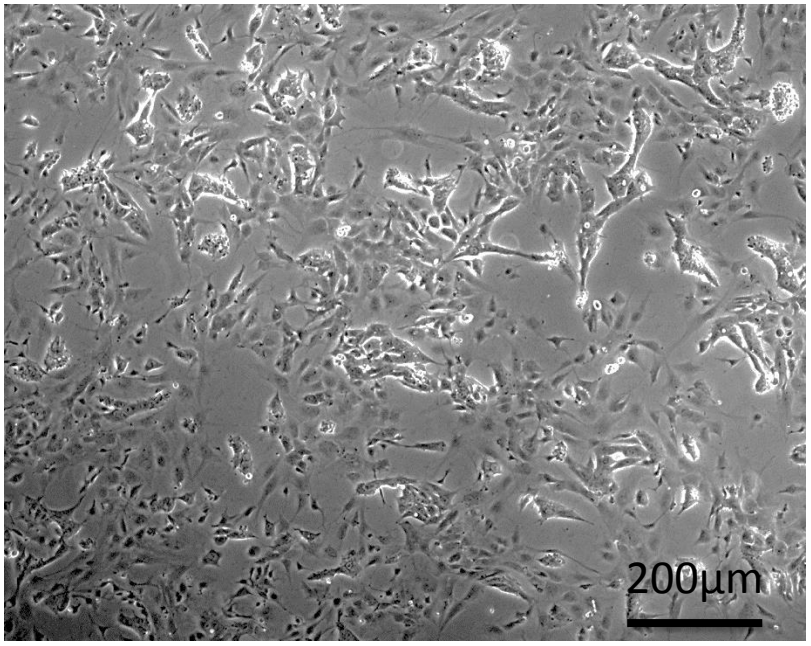
Date: 16.06.2015

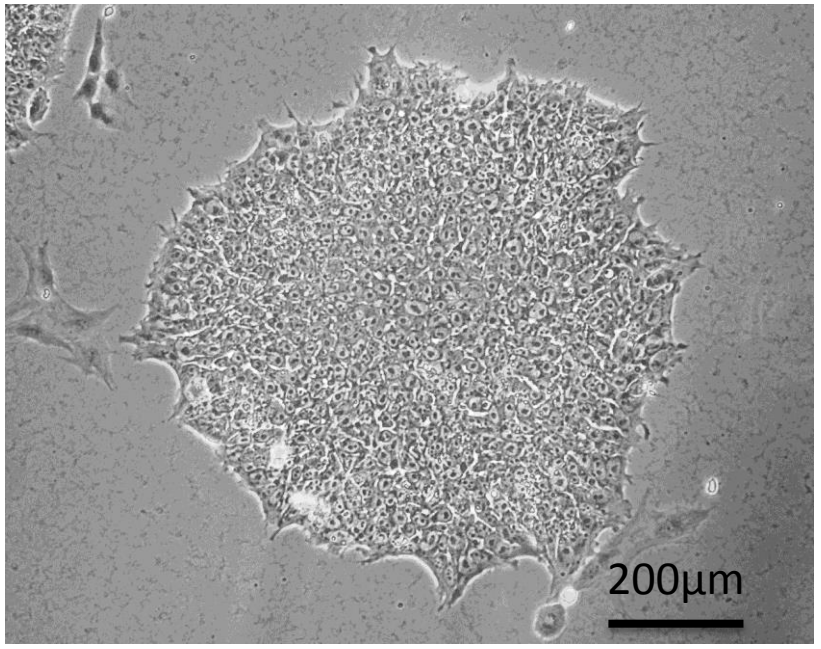
Source of fibroblasts and reprogramming information

- SF126 from UOXF 08/14
- Reprogrammed at UNEW ISV
- Reprogrammed on 30/01/2015 at passage 10
- Cytotune v2 WP3 SOP22

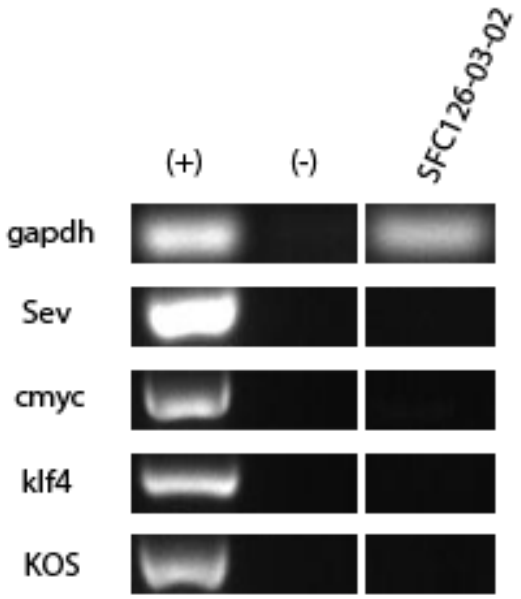
Viability post-thaw and Morphology according to SOP19 passage 14

- Cell count immediately post-thaw 2.8 x 10^6
- Viability immediately post-thaw 82%
- Photo at 24h post thaw (left) and 4d after ReLSR passaging (right):





Sendai clearance: according to WP3 SOP15 undetectable at passage 14

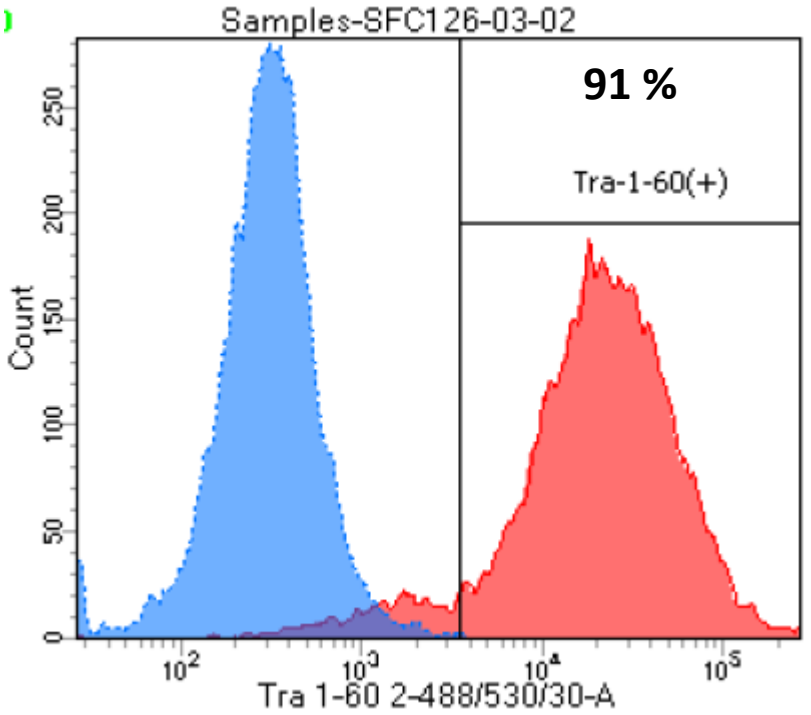


Mycoplasma test: Undetectable at passage 14

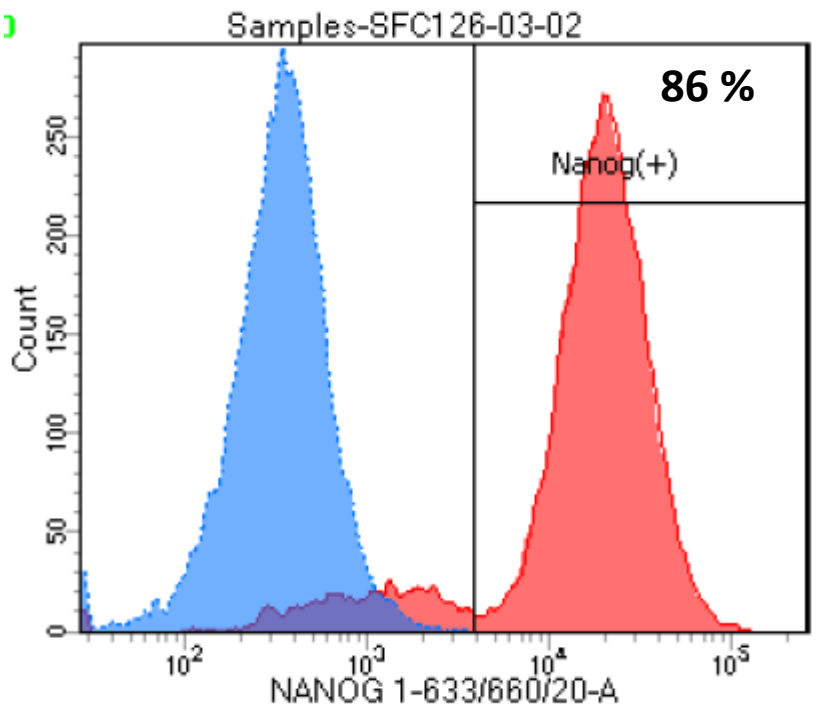
Owner	DM					
Date	08/06/2015					
Cell name	SFC126-03-02					
A	0.0284					
B	0.0161					
B/A	0.566901408					
> 1.2		Mycoplasma Contaminated			Positive Control	Negative Control
0.9-1.2		Status Unknown - Restest within 24 hours			0.0231	0.0862
0-0.9		Mycoplasma Free			1.769	0.0062
					76.58008658	0.071925754

Flow cytometric analysis according to WP3 SOP 20 and 21 passage 14

Tra-1-60:



NANOG:



SNP analysis
according to WP3 SOP Preparation of DNA and RNA samples for Illumina arrays

- Passage 14
- Identity to parent fibroblasts confirmed
- Karyotype abnormalities: none detected
- For details and raw data see StemDB

Comments

- Thawing in 2 wells of 6 well-plate recommended
- Clone prone to differentiation > ReLESR Passaging after thawing recommended
  - Add 1ml ReLESR, incubate for 3 min at 37˚C
  - Remove ReLESR
  - Add medium **dropwise** to collect undifferentiated cells