Certificate of analysis

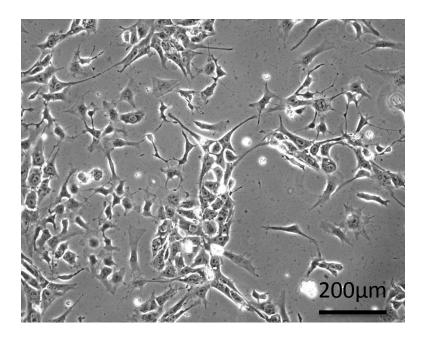
SFC888-07-03

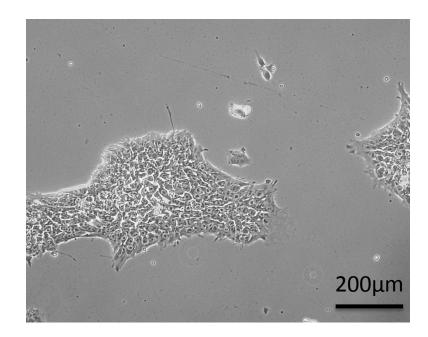
Signature: Dario Melguizo Sanchis Date: 12.06.2015 Supervisor signature: Linda Lako Date: 12.06.2015 Source of fibroblasts and reprogramming information

- SF888 from UOXF 01/15
- Reprogrammed at UNEW ISV
- Reprogrammed on 17/02/2015 at passage 8
- Cytotune v2 WP3 SOP22

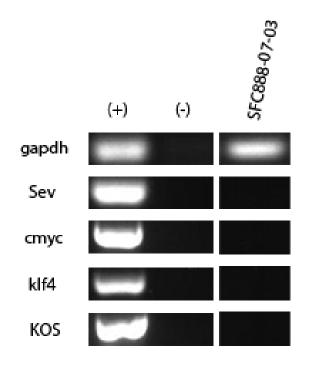
Viability post-thaw and Morphology according to SOP19 passage 11

- Cell count immediately post-thaw 1.6 x 10⁶
- Viability immediately post-thaw 92%
- Photo at 24h post thaw (left) and 3d after ReLSR passaging (right):





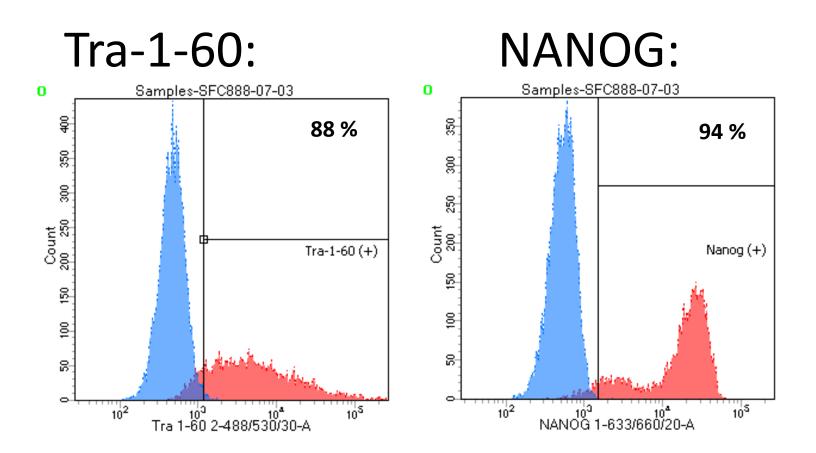
Sendai clearance: according to WP3 SOP15 undetectable at passage 11



Mycoplasma test: Undetectable at passage 11

Owner	DM				
Date	08/06/2015				
Cell name	SFC888-07-03				
Α	0.0249				
B	0.01				
B/A	0.401606426				
> 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 11



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

- Passage 11
- Identity to parent fibroblasts confirmed
- Karyotype abnormalities: Chr17 and Chr22 allelic imbalance which is not present in parental fibroblasts
- For details and raw data see StemDB

Comments

- Thawing in 2 wells of 6 well-plate recommended
- Clone prone to differentiation (see flow results weak positive population (2nd peak)) > ReLESR Passaging after thawing recommended
 - Add 1ml ReLESR, incubate for 3 min at 37°C
 - Remove ReLESR
 - Add medium **dropwise** to collect undifferentiated cells