

Information Summary

Organization: Neural Stem Cell Institute (NSCI) Address: One Discovery Drive, Rensselaer, NY 12144 Contact Person: Kathryn McIlroy Email: kathrynmcilroy@neuralsci.org Phone: 518-694-8188 x 226

SCRC Code: NSCI10

Sample Name: FTD-FF

Number of the Established iPSC Clones: 6

Characterization Performed:

- 1. Mycoplasma Test
- 2. Cell authentication Verification
- 3. Presence of the Pluripotency Marker
- 4. Spontaneous differentiation via embryoid body (EB) formation
- 5. Karyotyping

Qualified iPSC Clones: C1, C3, C4

Mycoplasma Test

Method: Determination of the mycoplasmal enzyme activity using Lonza MycoAlert Plus Detection kit.

iPSC Clone	NSCI10-C1	NSCI10-C3	NSCI10-C4
Cell passage	P6	P6	P6
B/A Ratio	0.52	0.84	0.44
Mycoplasma	NEG	NEG	NEG
Pass or Fail	Pass	Pass	Pass

Cell Authentication Verification

Markers	NSCI10 (FTD-FF)	NSCI10-C1	NSCI10-C3	NSCI10-C4
D3S1358	15, 16	15, 16	15, 16	15, 16
TH01	6	6	6	6
D21S11	30, 33.2	30, 33.2	30, 33.2	30, 33.2
D18S51	16, 19	16, 19	16, 19	16, 19
Penta E	13, 14	13, 14	13, 14	13, 14
D5S818	11, 13	11, 13	11, 13	11, 13
D13S317	11	11	11	11
D7S820	10, 11	10, 11	10, 11	10, 11
D16S539	11, 13	11, 13	11, 13	11, 13
CSF1PO	11	11	11	11
Penta D	11, 12	11, 12	11, 12	11, 12
vWA	14, 18	14, 18	14, 18	14, 18
D8S1179	13, 14	13, 14	13, 14	13, 14
ΤΡΟΧ	10, 11	10, 11	10, 11	10, 11
FGA	22, 23	22, 23	22, 23	22, 23
AMEL	х	х	х	х
Match with o	Match with original Fibroblast Yes Yes Y		Yes	



Expression of the Pluripotency Markers

Antibody staining for pluripotency markers: Oct4, Nanog, Tra-1-81, SSEA-4

NSCI10-C1	Oct4	Nanog	Tra-1-81	SSEA4
	DAPI	DAPI	DAPI	DAPI
NSCI10-C3	Oct4	Nanog	Tra-1-81	SSEA4
	DAPI	DAPI	DAPI	DAPI
	Oct4	Nanog	Tra-1-81	SSEA4
NSCI10-C4	DAPI	DAPI	DAPI	DAPI

Differentiation Potential via EB Formation

Method: The iPSC cells were disassociated into small clumps to form EB in the low attachment dish. After a 10day spontaneous differentiation, the EBs were harvested and RNAs were extracted. The expression levels of the selected 3-germ layer specific genes were analyzed by real-time PCR. Ct values are normalized for loading using a housekeeping gene.

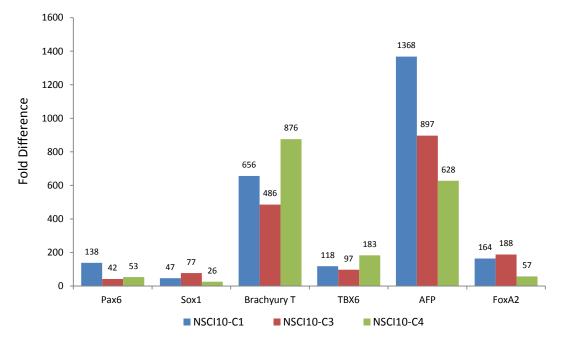


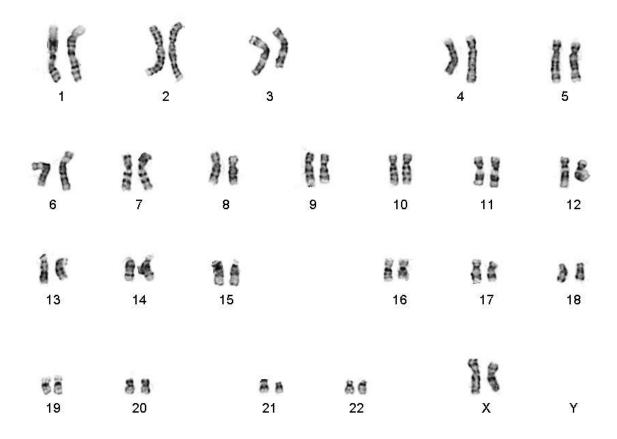
Figure: Lineage specific gene expression following EB differentiation. Fold difference is shown relative to undifferentiated iPS cell.



Karyotyping

Clone: NSCI10-C1 Passage: P7

- Tested:Number of metaphases counted: 20Banding and level: GTG/450Number of metaphases analyzed: 6Number of karyotypes: 2
- Result: 46,XX NORMAL FEMALE KARYOTYPE





Karyotyping

Clone: NSCI10-C3 Passage: Ρ7

- Number of metaphases counted: 20 Banding and level: GTG/550 Tested: Number of metaphases analyzed: 6 Number of karyotypes: 2
- **Result:** 46,XX NORMAL FEMALE KARYOTYPE

Channel of Channel	2	3			4	K 5
)(6	7	8)(9	10	요 전 11	12
13	14 14)) 15		16	置 17	18
) (19	音客 20	21	ð1 2:		X X	Y



Karyotyping

Clone: NSCI10-C4 Passage: P6

- Tested:Number of metaphases counted: 20Banding and level: GTG/450Number of metaphases analyzed: 6Number of karyotypes: 2
- **Result:** 46,XX NORMAL FEMALE KARYOTYPE

