

GM 23717*B

Certificate of Analysis

Product description	Human Fibroblast reprogrammed with seven		
	factors (Oct4, Sox2, Klf, c-Myc, Nanog, Lin-28, T-		
	antigen) using episomal vector		
Publication(s) describing iPSC establishment			
Parent cell line and cell type	Fibroblast	GM23311	
Diagnosis	Muscular dystrophy, congenital merosin-deficient,		
	1A; MDC1A		
Parent cell line freeze passage			
Passage of iPSC reported at submission	12		
Number of passages at Coriell	7		
Media	DMEM/F12 + 20% KOS	R + 100ng/ml bFGF	
Feeder	CF1 MEFs on 0.1% Gela	tin	
Passage method	Collagenase		
Split ratio	1:6; every 5-7 days		

The following testing specifications have been met for the specified product lot:

Test Description	Test Method	Test Specification	Result
Post-Thaw Viable Cell Recovery	Colony Doubling	Colony formation and diameter doubling within 5 days	Pass
Sterility	Growth on agar	Negative	Pass
Mycoplasma	PCR	Negative	Pass
Karyotype	G-banding	Normal Karyotype	Pass
Identity Match	STR (THO-1, D22S417, D10S526, vWA31, D5S592, and FES/FPS)	Match parent fibroblast line	Pass
Surface Antigen Expression of Stem Cell Markers	Immunostaining	> 80% expression of SSEA-4 < 10% expression of SSEA-1	Pass
Pluripotency	Pluripotency In vitro differentiation (cardiac, pancreatic and neuronal)		Pass
Teratoma Formation	eratoma Formation In Vivo Teratoma formation		Pass

Post-Thaw Viability

One vial of distribution lot was thawed. Cultures were observed daily. Colonies were photographed on the first day of appearance and then 5 days later. Colonies must double in diameter 5 days after first observation.

Days from Recovery to	Average Colony	Average Colony
First Colony Observation	Diameter (initial)	Diameter (post 7 days)
2 days	284	923



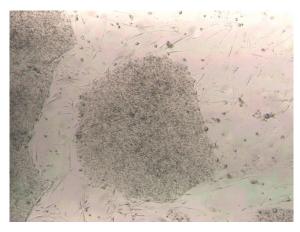


Figure 1A. Colony observed post thaw

Figure 1B. Colony 7 days after first observation

Karyotype Analysis

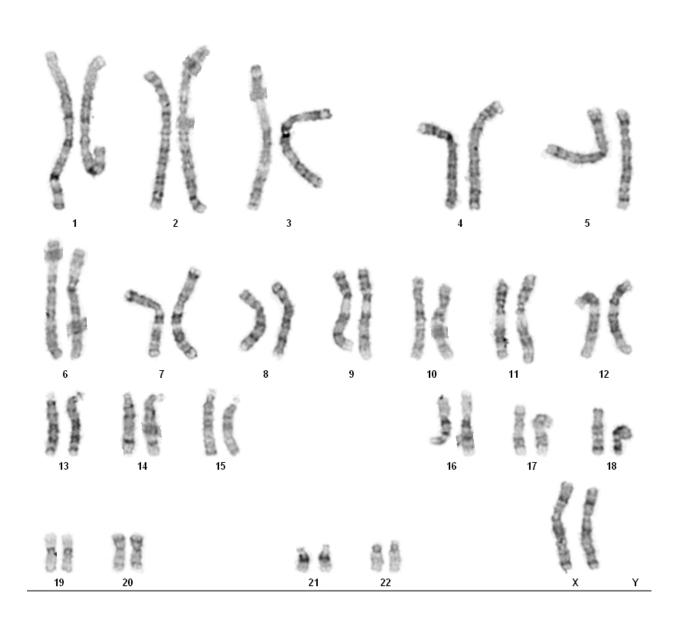


Figure 2: G-banded karyotype showing 46 XX

Surface Antigen Expression of Stem Cell Markers

Undifferentiated cells are stained for the surface antigens, SSEA4 and SSEA1. SSEA4 (stage specific embryonic antigen 4) is expressed on undifferentiated human stem cells. SSEA1 (stage specific embryonic antigen 1) is expressed on differentiated stem cells.

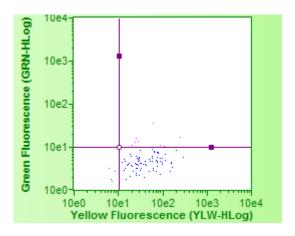


Figure 3A: Scatter plot of SSEA4 and SSEA1 double stained iPS cells.

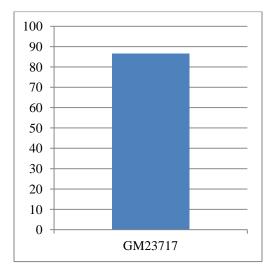


Figure 3B. Graph depicting percent SSEA4 positive cells in an undifferentiated cell culture.

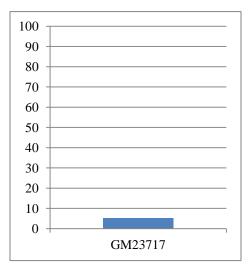


Figure 3C. Graph depicting percent SSEA1 positive cells in undifferentiated cell culture

Assessment of Pluripotency of a Cell Line

Cells are directed to differentiate to assess the pluripotency of the cell line. . RNA is harvested and gene expression is analyzed by real-time PCR. Ct values are adjusted for loading using a housekeeping gene. Gene expression is shown as fold difference to undifferentiated cells.

Embryoid Body (EB) Formation Assay

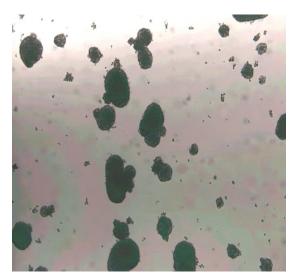


Figure 4A. Image of Embryoid Bodies, day 2

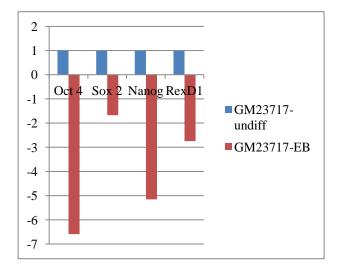


Figure 4B. Gene expression following EB differentiation. Fold difference is shown relative to undifferentiated iPS cell line.

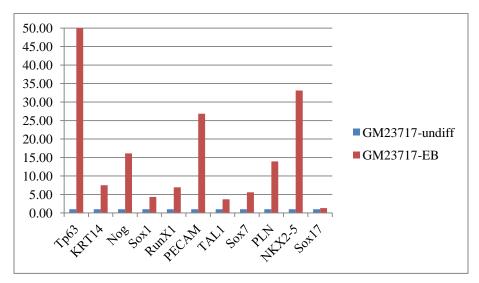


Figure 4C. Gene expression following EB differentiation. Fold difference is shown relative to undifferentiated iPS cell line.

	Tp63	KRT14	Nog	Sox1	RunX1	PECAM	TAL1	Sox7	PLN	NKX2-5	Sox17	AFP
GM23717- undiff	1.00	1.00	1.00	1.00	1.00	1.00	1.00	1.00	1.00	1.00	1.00	1.00
GM23717- EB	844	7.5	16.1	4.3	7.0	26.8	3.7	5.6	13.9	33.1	1.3	18823

Table 1. Fold difference values of gene expression of EB. Fold difference is shown as fold difference to undifferentiated cells.

Neural Differentiation

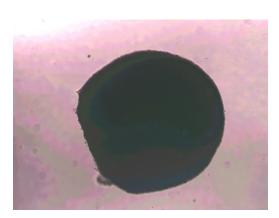


Figure 5A. Image of Neuronal Differentiation

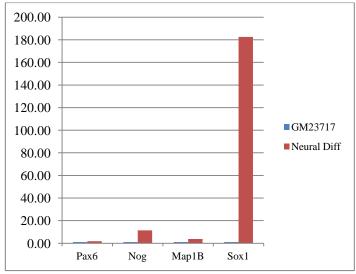


Figure 5B. Gene expression following neuronal differentiation. Fold difference is shown relative to undifferentiated iPS cell line.

Cardiac Differentiation

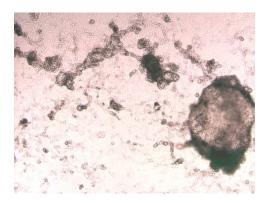


Figure 6A. Image of differentiated colony.

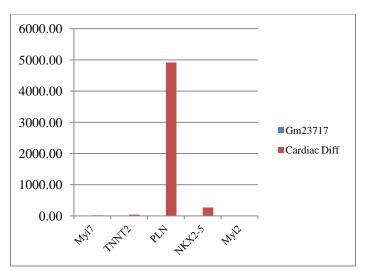


Figure 6B. Gene expression following cardiac differentiation. Fold difference is shown relative to undifferentiated iPS cell line.

Definitive Endoderm Differentiation



Figure 7A. Image of Definitive Endoderm Differentiation

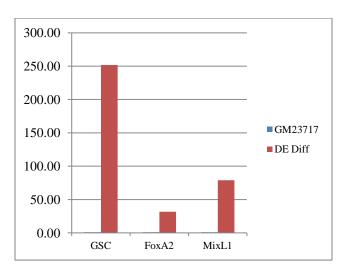


Figure 7B. Gene expression following pancreatic differentiation. Fold difference is shown relative to undifferentiated iPS cell line.

⊠ Pass	Steven Madre
☐ Fail	Steven Madore, PhD
Other:	Director, Stem Cell Biobank
	Date <u>June 29, 2012</u>



Teratoma Formation Analysis Report

Project Information

Service Title: Teratoma Formation Analysis

Customer: Coriell Institute

PI/Contact Person: Karen Fecenko-Tacka

Report date: May 31, 2012 Project manager: Qi Zheng

Contact person: Tianmin "Ivy" Zhang

Service Detail

Cell type: human iPS cells

Cell line & Passage: GM23717, P4

Feeder layer: CF1 MEF

Mouse type: Fox Chase SCID-beige, male, 6 week old, from Charles River

Cell concentration: 2-3 million/site, in 30% Matrigel

3 H&E slides

Injection date: March 21, 2012

	Mouse #1	Mouse #2	Mouse #3	Control
Lateration Office	kidney capsule	kidney capsule	kidney capsule	kidney capsule
Injection Sites	testis	testis	testis	testis
Tissue harvested	one kidney tumor and one testis tumor			
Days post-injection	47	47	47	47

H&E Histology Instruction

Histology: 10% Formalin fixed over night, embedded in paraffin, cut into 5-μm serial sections, H&E staining

Three embryonic germ cell layers: endoderm, mesoderm and ectoderm

Endoderm: digestive system (includes liver and pancreas), respiratory system, most glands

Mesoderm: muscle, blood vessels, much of the genital-urinary system, skeletal system

Ectoderm: skin, hair, nails, sweat and mammary glands, nervous system (including hypothalamus and both lobes of the pituitary gland), oral and nasal

Tel: 408-773-8007

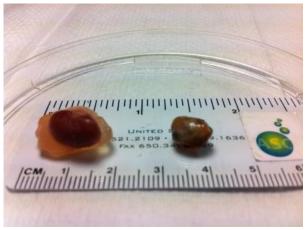
cavities, portions of the vagina, vestibule, penis and clitoris



Tumor and organ pictures



Mouse#1: one kidney tumor (left) and one testis tumor (right) harvested on day 47 after injection



Mouse#2: one kidney tumor (left) and one testis tumor (right) harvested on day 47 after injection



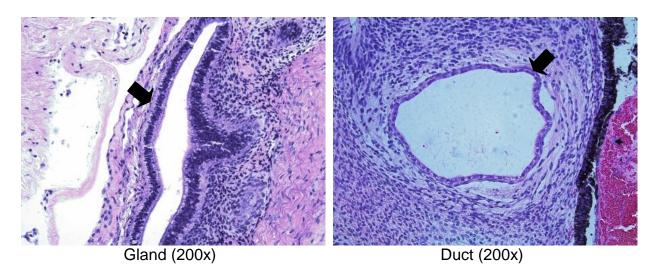
Mouse#3: one kidney tumor(left) and one testis tumor (right) harvested on day 47 after injection

Tel: 408-773-8007

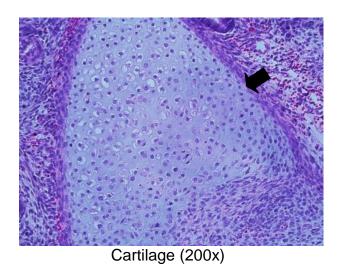


H&E staining results of kidney and testis tumors:

Endoderm

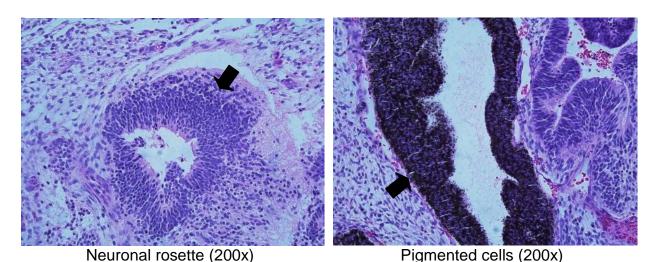


Mesoderm





Ectoderm



Summary

Three kidney tumors and three testis tumors are composed of scattered regions of differentiated cells and a large population of undifferentiated neoplastic cells. Three germ layers were clearly identified in histology analysis. The tissues listed above indicate that small areas of what might be these kinds of tissues were noted within the tumor. Overall, there is some degree of differentiation of these cells with organized structures, suggesting that some of these cells are pluripotent.

Tel: 408-773-8007

Project manager

Signature: Date: 5/31/2012

Qi Zheng, Ph.D. Senior Scientist

Reviewed and proved by

Signature: Date: 5/31/2012

Steve Yu, Ph.D.

Director of Service Department