

GM 23230

Certificate of Analysis

Product description	Human fibroblast line reprogrammed with four				
	factors (Oct 4, Sox 2, c-Myc and Klf-4) using				
	retroviral vector				
Publication(s) describing iPSC establishment	Park et al., PMID <u>18691744</u>				
Parent cell line and cell type	GM04569 Fibroblast				
Diagnosis	Muscular Dystrophy, Becker Type; BMD				
Parent cell line freeze passage					
Passage of iPSC reported at submission	13				
Number of passages at Coriell	10				
Media	DMEM/F12 + 20% KOSR + 10 ng/ml bFGF				
Feeder	CF1 MEFs on 0.1% Gelatin				
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	Cell 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 In vitro differentiation (cardiac, pancreatic and neuronal)		Upregulation of genes appropriate to cell lineage	Pass	
Teratoma Formation	atoma 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	126	888

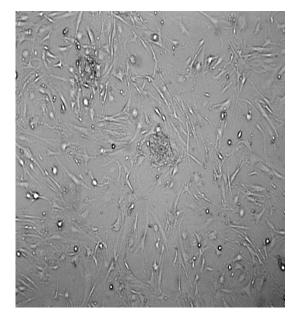


Figure 1A. Colony observed 2 days post thaw

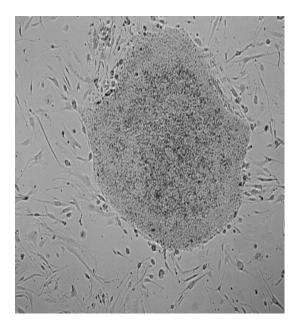


Figure 1B. Colony 5 days after first observation

Karyotype Analysis

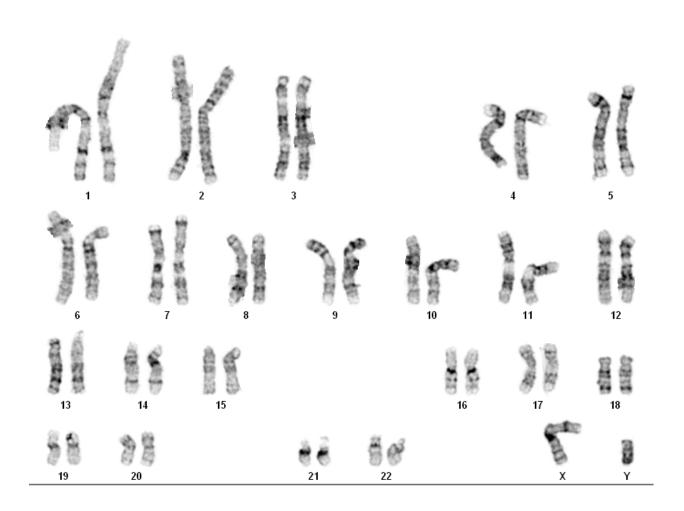


Figure 2: G-banded karyotype showing 46 XY

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.

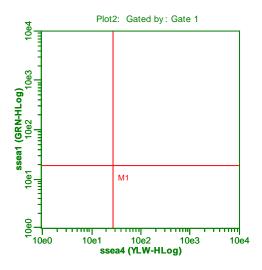


Figure 3A: Scatter plot of SSEA4 stained iPS cells.

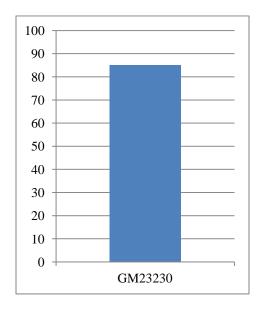


Figure 3B. Graph depicting percent SSEA4 positive cells in an undifferentiated cell culture.

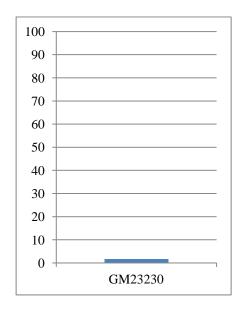


Figure 3C. Graph depicting percent SSEA1 positive cells in undifferentiated cell culture

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

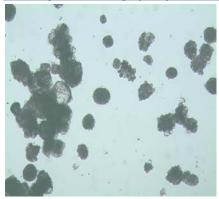


Figure 4A- Embryoid Body formation, day 11

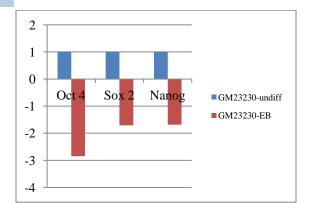


Figure 4B. Gene expression following EB differentiation. Fold difference is shown relative to undifferentiated iPS cell line.

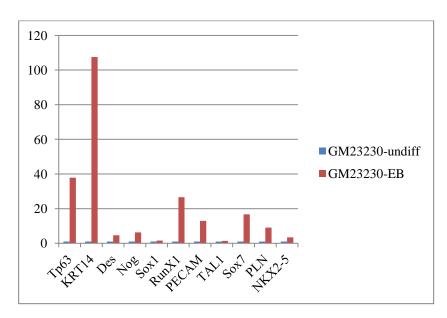


Figure 4C. Gene Expression following EB differentiation. Fold difference is shown relative to undifferentiated iPS cell line

	Tp63	KRT14	Des	Nog	Sox1	RunX1	PECAM	TAL1	Sox7	PLN	NKX2- 5	AF P
GM23 230- undiff	1	1	1	1	1	1	1	1	1	1	1	1
GM23 230- EB	38	107	5	6	2	27	13	1	17	9	3	87

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

Neural Differentiation

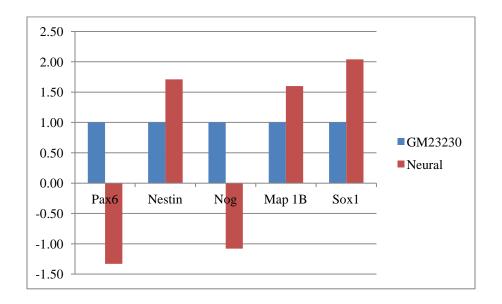


Figure 5B. Gene expression following neuronal differentiation. Fold difference is shown relative to undifferentiated iPS cell line.

Cardiac Differentiation



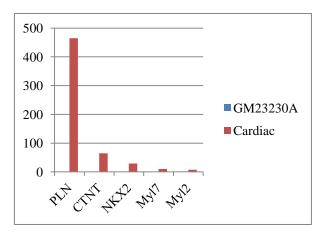


Figure 6A. Image of differentiated colony.

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

Pancreatic Differentiation

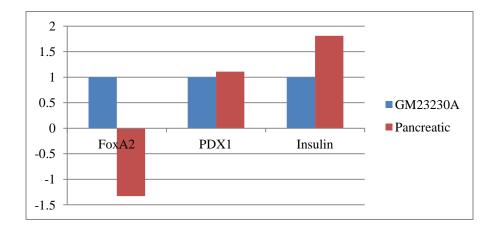


Figure 7B. Gene expression following pancreatic differentiation. Fold difference is shown relative to undifferentiated iPS cell line.



Teratoma Formation Analysis Report

Project Information

Service Title: Teratoma Formation Analysis

Customer: Coriell Institute

PI/Contact Person: Karen Fecenko-Tacka

Report date: October 7, 2011 Project manager: Qi Zheng

Contact person: Tianmin "Ivy" Zhang

Service Detail

Cell type: human iPS cells

Cell line & Passage: GM23230A, P3

Feeder layer: CF1 MEF

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

Cell concentration: 1.5 to 3 million/site, in 30% Matrigel

6 H&E slides

Injection date: August 17, 2011

	Mouse #1	Mouse #2	Mouse #3	Control
	kidney	kidney	kidney	kidney
Injection Sites	testis	testis	testis	testis
Tissue harvested	one kidney tumor and one testis tumor			
Days post-injection	41	41	41	41

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

Tel: 408-773-8007



Tumor pictures



Mouse#1: one kidney tumor (left) and one testis tumor (right) harvested on day 41 after injection



Mouse#2: one kidney tumor (left) and one testis tumor (right) harvested on day 41 after injection

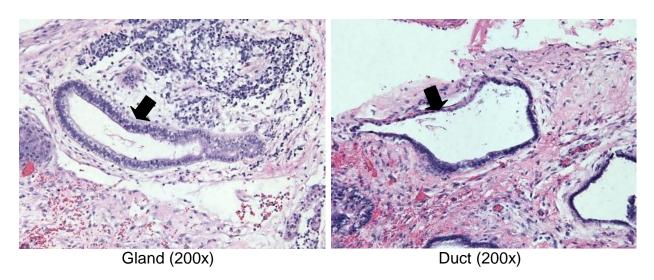


Mouse#3: one kidney tumor (left) and one testis tumor (right) harvested on day 41 after injection

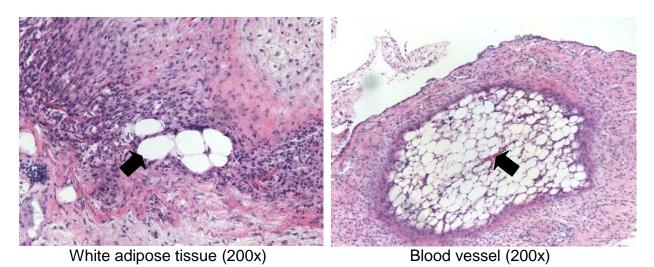


H&E staining results of kidney and testis tumors:

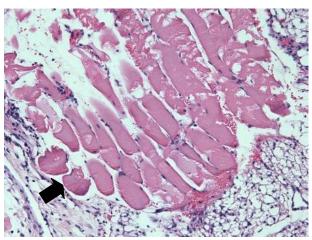
Endoderm

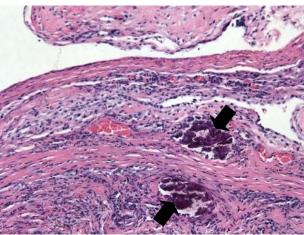


Mesoderm





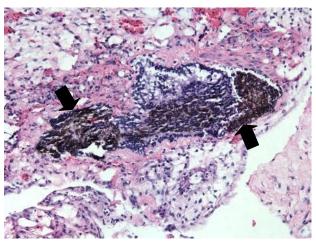




Skeletal muscle (200x)

Bone (200x)

Ectoderm



Pigmented cells (200x)

Summary

Three kidney tumors and three testis tumors harvested on day 41 after injection are composed of scattered regions of differentiated cells and a large population of undifferentiated neoplastic cells. In these tumors, 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.



Project manager

Signature: _____ Date: <u>10/7/2011</u>

Qi Zheng, Ph.D. Senior Scientist

Reviewed and proved by

Signature:_____ Date: <u>10/7/2011</u>

Steve Yu, Ph.D.

Director of Service Department

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