



Thaw and Culture Details

Cell Line Name	PENN068i-697-3
WiCell Lot Number	DB36611
Provider	University of Pennsylvania – Dr. Daniel Rader
Banked By	Penn Institute for Regenerative Medicine iPS Core Facility
Thaw and Culture Recommendations	WiCell recommends thawing 1 vial into 2 wells of a 6 well plate. WiCell recommends thawing using ROCK Inhibitor for best results.
Culture Platform	Feeder Dependent
	Medium: hESC Medium (KOSR)
	Matrix: MEF
Protocol	WiCell Feeder Dependent Protocol
Passage Number	p13 These cells were cultured for 13 passages prior to freeze and post colony picking. Therefore, plated cells at thaw should be labeled passage 14.
Date Vial	20-April-2015
Vial Label	iPS-697 SEV3 P13 4/20/2015 ZL
Biosafety and Use Information	Appropriate biosafety precautions should be followed when working with these cells. The end user is responsible for ensuring that the cells are handled and stored in an appropriate manner. WiCell is not responsible for damages or injuries that may result from the use of these cells. Cells distributed by WiCell are intended for research purposes only and are not intended for use in humans.

Testing Performed by WiCell

Test Description	Test Provider	Test Method	Test Specification	Result
Karyotype by G-banding	WiCell	SOP-CH-003	Expected karyotype	See Report
	<p>Results: 46,XY,dup(1)(q21q42)[15]/46,XY[3]/46,XX,t(1;13)(p22;q32)[2]/46,XX[1] Nonclonal Finding: 47,XY,+X,dup(1)(q21q42)</p> <p>Interpretation: There are both male and female cells in this culture; the most likely explanation for this is that two cultures have been mixed. Please resubmit the specimen. Both the male and female cell populations are karyotypically abnormal. In the male (XY) cells, there is a duplication of the long arm (q) of chromosome 1 in fifteen of 18 metaphases; this is a recurrent cytogenetic abnormality in human pluripotent stem cell cultures. In the female (XX) cells, there is a reciprocal translocation between the short arm (p) of chromosome 1 and the long arm (q) of chromosome 13 in 2 of three metaphases, not considered recurrently acquired in cultures of this cell type. No other clonal abnormalities were detected at the stated band level of resolution. There is also a nonclonal finding in the male cell population, listed above. Nonclonal findings likely result from technical artifact, but may be due to a developing clonal abnormality or to low-level mosaicism.</p>			
Post-Thaw Viable Cell Recovery	WiCell	SOP-CH-305	Recoverable attachment after passage	Pass
Identity by STR	UW Translational Research Initiatives in Pathology Laboratory	PowerPlex 16 HS System by Promega	Defines profile	Pass
Sterility	Steris	ST/07	Negative	Pass
Mycoplasma	WiCell	SOP-QU-004	Negative	Pass



Testing Reported by Provider

The Provider stated that some or all of the additional analyses listed below may have been performed for this cell line. For more information, publication and dbGaP links, where available, are provided on the cell line specific web page on the WiCell website.

- SNP microarray
- Flow Cytometry (Tra1-60 and SSEA-4)
- Differentiation into hepatocytes
- Infinium® Expanded Multi-Ethnic Genotyping Array (MEGA^{EX})

Approval Date	Quality Assurance Approval
23-June-2016	<p style="text-align: right;">6/5/2018</p> <p>X HEB</p> <p><small>HEB Quality Assurance Signed by Bruner, Haley</small></p>

Date Reported: Wednesday, May 02, 2018
Cell Line: PENN068i-697-3-DB36611-13636
Passage#: 15

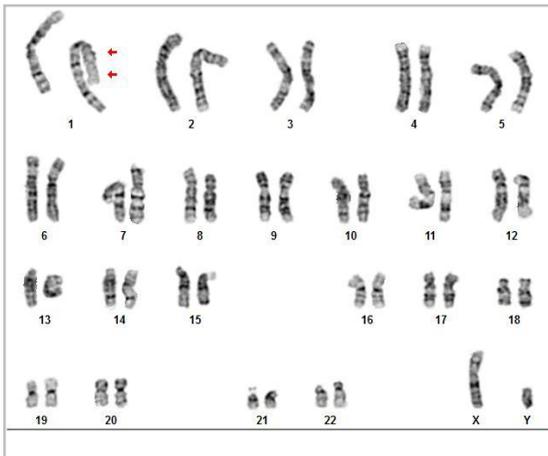
Cell Line Gender: Male
Reason for Testing: Lot Release Testing

Date of Sample: 4/24/2018
Specimen: Human IPS

Investigator: [REDACTED], WiCell

Results: 46,XY,dup(1)(q21q42)[15]/46,XY[3]/46,XX,t(1;13)(p22;q32)[2]/46,XX[1]

Nonclonal Finding: 47,XY,+X,dup(1)(q21q42)



Cell: 39
Slide: G02
Slide Type: Karyotype

Total Counted: 22
Total Analyzed: 10
Total Karyogrammed: 6
Band Resolution: 450-500

Interpretation:

There are both male and female cells in this culture; the most likely explanation for this is that two cultures have been mixed. Please resubmit the specimen.

Both the male and female cell populations are karyotypically abnormal. In the male (XY) cells, there is a duplication of the long arm (q) of chromosome 1 in fifteen of 18 metaphases; this is a recurrent cytogenetic abnormality in human pluripotent stem cell cultures. In the female (XX) cells, there is a reciprocal translocation between the short arm (p) of chromosome 1 and the long arm (q) of chromosome 13 in 2 of three metaphases, not considered recurrently acquired in cultures of this cell type. No other clonal abnormalities were detected at the stated band level of resolution.

There is also a nonclonal finding in the male cell population, listed above. Nonclonal findings likely result from technical artifact, but may be due to a developing clonal abnormality or to low-level mosaicism.

Completed by: [REDACTED] CG(ASCP)

Reviewed and Interpreted by: [REDACTED] Ph.D., FACMGG

A signed copy of this report is available upon request.

Date: _____ **Sent By:** _____ **Sent To:** _____ **QC Review By:** _____

Limitations: This assay allows for microscopic visualization of numerical and structural chromosome abnormalities. The size of structural abnormality that can be detected is >3-10Mb, dependent upon the G-band resolution obtained from this specimen. For the purposes of this report, band level is defined as the number of G-bands per haploid genome. It is documented here as "band level", i.e., the range of bands determined from the four karyograms in this assay. Detection of heterogeneity of clonal cell populations in this specimen (i.e., mosaicism) is limited by the number of metaphase cells examined, documented here as "# of cells counted".

This assay was conducted solely for listed investigator/institution. The results may not be relied upon by any other party without the prior written consent of the Director of the WiCell Cytogenetics Laboratory. The results of this assay are for research use only. If the results of this assay are to be used for any other purpose, contact the Director of the WiCell Cytogenetics Laboratory.

Unless otherwise mutually agreed in writing, the services provided to you hereunder by WiCell Research Institute, Inc. ("WiCell") are governed solely by WiCell's Terms and Conditions of Service, found at www.wicell.org/privacyandterms. Any terms you may attach to a purchase order or other document that are inconsistent, add to, or conflict with WiCell's Terms and Conditions of Service are null and void and of no legal force or effect.

Sample Report:

13636-STR

Sample Name on Tube: 13636-STR

154.1 ng/μL, (A260/280=1.93)

Sample Type: Cells

Cell Count: ~2 million cells

Requestor:

WiCell Research Institute

Quality Department

Sample Date: N/A

Receive Date: 04/30/18

Assay Date: 05/01/18

File Name: STR 180502 wmr

Report Date: 05/09/18

STR Locus	STR Genotype Repeat #	STR Genotype
FGA	16-18,18.2,19,19.2,20,20.2,21,21.2,22, 22.2, 23, 23.2, 24, 24.2, 25, 25.2, 26-30, 31.2, 43.2, 44.2,45.2, 46.2	Identifying information has been redacted to protect donor confidentiality. If more information is required, please, contact WiCell's Technical Support .
TPOX	6-13	
D8S1179	7-18	
vWA	10-22	
Amelogenin	X,Y	
Penta_D	2.2, 3.2, 5, 7-17	
CSF1PO	6-15	
D16S539	5, 8-15	
D7S820	6-14	
D13S317	7-15	
D5S818	7-16	
Penta_E	5-24	
D18S51	8-10, 10.2, 11-13, 13.2, 14-27	
D21S11	24,24.2,25,25.2,26-28,28.2,29,29.2, 30, 30.2,31, 31.2,32,32.2,33,33.2, 34,34.2,35,35.2,36-38	
TH01	4-9,9.3,10-11,13.3	
D3S1358	12-20	

Results: Based on the 13636-STR cells submitted by WiCell QA dated and received on 04/30/18, this sample (Label on Tube: 13636-STR) defines the STR profile of the human stem cell line PENN068i-697-3 comprising 26 allelic polymorphisms across the 15 STR loci analyzed.

Interpretation: No STR polymorphisms other than those corresponding to the human PENN068i-697-3 stem cell line were detected and the concentration of DNA required to achieve an acceptable STR genotype (signal/noise) was equivalent to that required for the standard procedure (~1 ng/amplification reaction) from human genomic DNA. This result suggests that the 13636-STR sample submitted corresponds to the PENN068i-697-3 stem cell line and was not contaminated with any other human stem cells or a significant amount of mouse feeder layer cells.

Sensitivity: Sensitivity limits for detection of STR polymorphisms unique to either this or other human stem cell lines is ~2-5%.

X *RMB* Digitally Signed on 05/10/18

X *WMR* Digitally Signed on 05/10/18

_____, BA
TRIP Laboratory, Molecular

_____, PhD, Director / Co-Director
UWHC Molecular Diagnostics Laboratory / UWSMPH TRIP Laboratory

Testing was accomplished by analysis of human genetic polymorphisms at STR loci. This methodology has not yet been approved by the FDA and is for investigational use only. Acknowledge TRIP in your publications, posters & presentations. For details, see: <http://www.pathology.wisc.edu/research/trip/acknowledging> TRIP agrees to maintain the confidentiality of any information provided to it in connection with its performance of this STR analysis on the same conditions as set forth in paragraph 2 of WiCell's Terms and Conditions of Service (<http://www.wicell.org/media.acux/1a429b84-2b54-44a4-8ad8-5c05db93dd8a>).



Mycoplasma Detection Assay Report

Testing Performed by WiCell

Lot Release Testing

April 16, 2018

FORM SOP-QU-004.01

Version G Edition 02

Reported by: AP

Reviewed by: DF

BD Monolight 180

#	Sample Name	Reading A		A Ave	Reading B		B Ave	Ratio B/A	Result	Comments/Suggestions
		RLU1	RLU2		RLU1	RLU2				
1	PENN068i-697-3-DB36611 13636	374	374	374	174	169	171.5	0.46	Negative	
2	Positive (+) Control	427	435	431	23689	23939	23814	55.25	Positive	
3	Negative (-) Control	669	659	664	80	83	81.5	0.12	Negative	

