

## **Product Information**

Product Name	MIRJT6i-mND1-4					
Lot Number	WB0163					
Depositor	Morgridge Institute for Research – Laboratory of Dr. James Thomson					
Banked by	WiCell					
Thaw Recommendation	Thaw 1 vial into 1 well of a 6 well plate. WiCell recommends thawing using ROCK Inhibitor for best results.					
Culture Platform	Feeder Independent					
	Medium: E8 – WiCell recommends to passage using ROCK Inhibitor for best results.					
	Matrix: Matrigel					
Protocol	WiCell Feeder Independent E8 Medium Protocol modified to include ROCK Inhibitor at passage					
Passage Number	p32					
	These cells were cultured for 31 passages prior to freeze, 5 of them in E8/Matrigel. WiCell adds +1 to the passage number at freeze so that the number on the vial best represents the overall passage number of the cells at thaw.					
Date Vialed	21-May-2012					
Vial Label	WB0163 MIRJT6i-mND1-4 p32 21MAY12 DF					
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
Post-Thaw Viable Cell Recovery	WiCell	SOP-CH-305	≥ 15 Undifferentiated Colonies,	Pass
			≤ 30% Differentiation	
Identity by STR	UW Molecular Diagnostics	PowerPlex 16 HS	Consistent with known profile	Pass
	Laboratory	System by Promega		
Sterility	Apptec	30744	Negative	Pass
Mycoplasma	WiCell	SOP-QU-004	Negative	Pass
Karyotype by G-banding	WiCell	SOP-CH-003	Expected karyotype	Pass

Date of Lot Release	Quality Assurance Approval		
28- January-2013	1/25/2019 X RK		
	RK Quality Assurance Signed by: Kremers, Erik		



Histocompatibility/Molecular Diagnostics Laboratory

University of Wisconsin Hospital and Clinics

# Short Tandem Repeat Analysis\*

#### Sample Report: 10690-STR

Label on Tube: 10690-STR

Sample Date: 01/16/13 Lab Received 01/16/13

Requestor: WiCell Research Institute Test Date: 01/16/13

File Name: 130116 SLE

Report Date: 01/22/13

Sample Name: (label on tube) 10690-STR

**Description:** WI Cell Research Institute provided genomic DNA 254.5 ug/mL 260/280=1.79

Locus	Repeat #	STR Genotype
D16S539	5, 8-15	Identifying information
D7S820	6-14	has been redacted to
D13S317	7-15	protect donor
D5S818	7-15	more information is
CSF1PO	6-15	required, please,
TPOX	6-13	contact WiCell's
Amelogenin	NA	Technical Support.
TH01	5-11	
vWA	11, 13-21	

Comments: Based on the 10690-STR DNA submitted by WI Cell dated and received on 01/16/13, this sample (Label on Tube: 10690-STR) exactly matches the STR profile of the human stem cell line MIRJT6i-mND1-4 comprising 15 allelic polymorphisms across the 8 STR loci analyzed. No STR polymorphisms other than those corresponding to the human stem cell line MIRJT6i-mND1-4 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 suggest that the 10690-STR DNA sample submitted corresponds to the MIRJT6i-mND1-4 stem cell line and was not contaminated with any other human stem cells or a significant amount of mouse feeder layer cells. Sensitivity limits for detection of STR polymorphisms unique to either this or other human stem cell lines is ~5%.



Molecular Diagnostics Laboratory

\* Testing to assess engraftment following bone marrow transplantation 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.

This report is confidential. No part may be used for advertising or public announcement without written permission. Results apply only to the sample(s) tested.



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August 30, 2012 P.O. #:

WiCell Research Institute 505 S. Rosa Road Suite 120 Madison, WI 53719

Attn: Jessica Martin

#### STERILITY TEST REPORT

Sample Information:

1: DF19-9-7T-FTDL-01 10573 2: WA09-WB0156 10574 3: MIRJT6i-mND1-4-WB0163 10576 4: MIRJT6i-mND1-4-WB0162 10577 5: iPS(IMR90)-4-CB-01 10578 6: IISH6i-CML17-WB0170 10579 7: WA25-WB0169 10580

Date Received: Date in Test: Date Completed: August 09, 2012 August 15, 2012 August 29, 2012

**Test Information:** 

Test Codes: 30744, 30744A Immersion, USP / 21 CFR 610.12 Procedure #: BS210WCR.201

TEST PARAMETERS	PRODUCT			
Approximate Volume Tested	0.5 mL	0.5 mL		
Number Tested	14	14		
Type of Media	SCD	FTM		
Media Volume	400 mL	400 mL		
Incubation Period	14 Days	14 Days		
Incubation Temperature	20 °C to 25 °C	30 °C to 35 °C		
RESULTS	12 NEGATIVE 2 POSITIVE	12 NEGATIVE 2 POSITIVE		

Note: SCD and FTM Samples WA09-WB0156 10574 positive.

Testing conducted in accordance with current Good Manufacturing Practices.



## Mycoplasma Report

Testing Performed by WiCell RP WiCell /LRT 10690 1-17-13 FORM SOP-QU-004.01

Version B

Edition 01

Assay performed and reported by: MW Reviewed by: JB Equipment: 539 Berthold

		Readi	ng A	Α	Read	ing B	В	Ratio		
Sa	mple Number and ID	A1	A2	Average	B1	B2	Average	B/A	Mycoplasma Results	Comments/Suggestions
1	WB0163-Kp34 MW	469	477	473	217	223	220	0.47	Negative	
2	Positive (+) Control	153	153	153	13001	12986	12993.5	84.92	Positive	
3	Negative (-) Control	347	354	350.5	51	49	50	0.14	Negative	





Date Reported: Tuesday, January 15, 2013 Cell Line: MIRJT6i-mND1-4-WB0163 10690 Passage#: 34 Date of Sample: 1/9/2013 Specimen: iPSC Results: 46,XY





#### Interpretation:

This is a normal karyotype. No clonal abnormalities were detected at the stated band level of resolution.

Completed by: Reviewed and Interpreted by: A signed copy of this report is ava	CG(AS	SCP) , PhD, FACMG st.	
Date:	Sent By: Se	ent 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.